

APPROVED DRUG PRODUCTS

WITH

**THERAPEUTIC
EQUIVALENCE
EVALUATIONS**

42nd 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**

2022

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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, 2021.

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Therapeutic Equivalence Evaluations**

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

**APPROVED DRUG PRODUCTS
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Therapeutic Equivalence Evaluations**

PREFACE TO FORTY SECOND 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the inclusion of any product is that the product is the subject of an application with an approval that has not been withdrawn for safety or effectiveness reasons.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 revisions. 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 amended the FD&C Act to establish, among other things, the 505(b)(2) and 505(j) approval pathways. 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 have qualified 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 under section 505(b) of the FD&C Act or an ANDA under section 505(j) of the FD&C Act 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 Central Document Room, Attn: Director, Division of Orange Book Publication and Regulatory Assessment (DOBPR), Office of Generic Drug Policy, Center for Drug Evaluation and Research, Food and Drug Administration, 5901-B Ammendale Rd., Beltsville, MD 20705-1266. 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 effectiveness reasons subsequent to being discontinued from marketing.¹ This publication also includes indices of prescription and OTC drug products by proprietary name (brand name or trade name) or, if no proprietary name exists, established name of the active ingredient and by applicant name, which have been abbreviated for this publication. Established names for active ingredients generally conform to 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 effectiveness and for which NDAs 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 Division of Orange Book Publication and Regulatory Assessment 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 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 equivalent to the applicant's drug product even if the applicant's drug

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).

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, 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

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

⁶ 21 CFR 314.3(b).

⁷ 21 CFR 314.3(b).

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

⁸ 21 CFR 314.3(b).

⁹ 21 CFR 320.24.

¹⁰ We note that prior to the 36th edition of the Orange Book, 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 <https://www.fda.gov/drugs/guidance-compliance-regulatory-information/guidances-drugs> and FDA Drugs guidance (Product-Specific Guidances for Generic Drug Development) Web page at <https://www.fda.gov/drugs/guidances-drugs/product-specific-guidances-generic-drug-development>.

1.4 Reference Listed Drug and Reference Standard

A reference listed drug is the listed drug¹¹ identified by FDA as the drug product upon which an applicant relies in seeking approval of its ANDA.¹² FDA's general practice is to designate as RLDs drug products that have been approved for safety and effectiveness under section 505(c) of the FD&C Act. 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.¹³

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.¹⁴ 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 for reasons other than safety or effectiveness, FDA may select an ANDA that is therapeutically equivalent to this reference listed drug 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.

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 prescription 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.

¹¹ A "listed drug" is a new drug product that has been approved under section 505(c) of the FD&C Act for safety and effectiveness or under section 505(j) of the FD&C Act, which has not been withdrawn or suspended under section 505(e) (1) through (5) or section 505(j) (6) of the FD&C Act, and which has not been withdrawn from sale for what FDA has determined are reasons of safety or effectiveness. Listed drug status is evidenced by the drug product's identification in the current edition of FDA's "Approved Drug Products With Therapeutic Equivalence Evaluations" (the list) as an approved drug. A drug product is deemed to be a listed drug on the date of approval for the NDA or ANDA for that drug product (21 CFR 314.3(b)).

¹² 21 CFR 314.3(b).

¹³ 21 CFR 314.94(a) (3) (i).

¹⁴ 21 CFR 314.3(b).

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

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 (i.e., not on the Discontinued Drug Product list) from more than one manufacturer. For such products, a therapeutic equivalence code generally 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. Products listed in the Orange Book are identified by the applicant's name (firm name on the Form FDA 356h in the application). Where the applicant's name does not appear on the label, a person wishing to relate a specific product to the applicant name in the Orange Book may refer to FDA's NDC Directory¹⁵ and match its search terms to information on the label, such as the NDC Code if available.

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 NDA or ANDA that has been approved and that has not been withdrawn for safety or effectiveness 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, prescription 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

¹⁵ <https://www.fda.gov/drugs/drug-approvals-and-databases/national-drug-code-directory>.

prescription drug product (e.g., a particular strength of an approved drug that is not on the Discontinued Drug Product list) as therapeutically equivalent to other pharmaceutically equivalent prescription drug 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. 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.

For example, 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 ingredient(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.¹⁶ 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 x mg/mL. Generally, the amount of dry powder or lyophilized powder in a container is identified as the strength, expressed as x mg/vial.

However, FDA subsequently realized that the format of the Orange Book with respect to parenteral solutions should be changed to reflect that each strength of a drug is considered to be a separate listed drug. The Orange Book displays the strength of all new approvals of parenteral solutions. Previously (i.e., prior to 2003), we would have displayed only the concentration of an approved parenteral solution, e.g. 50 mg/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. 1 gm/20 mL (50 mg/mL) and 3 gm/60 mL (50 mg/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, suppositories, and inserts. 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.

¹⁶ The strengths of certain parenteral drug products, including contrast agents, may be expressed as a percentage.

"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, sprays, suppositories, and inserts 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.¹⁷ 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 therapeutic equivalence codes may be potentially confusing and inadequate for these drug products. Looking at the Orange Book listing alone for a product identified as a reference listed drug or reference standard, it may be difficult to determine to which therapeutic equivalence code the reference listed drugs and/or reference standard designation corresponds. For example, Unithroid 0.3 mg strength has been assigned the therapeutic equivalence codes AB1, AB2, and AB3 and it is identified as the reference listed drug and reference standard, but it is unclear that the reference listed drug and reference standard designations are associated with the AB1 therapeutic equivalence code.

Accordingly, FDA provides the following chart, which identifies (1) a reference listed drug for each therapeutic equivalence code in the Orange Book and (2) and the reference standard products in the Active Section of the Orange Book.¹⁸

- Therapeutic equivalence has been established between products that have the same AB+number therapeutic equivalence code (i.e. AB1, AB2, AB3 or AB4).
- More than one therapeutic equivalence code may apply to some products. One common therapeutic equivalence code indicates therapeutic equivalence between products. For example, Unithroid has been assigned therapeutic equivalence codes AB1, AB2, and AB3, and therefore, Unithroid tablets are considered therapeutically equivalent to other levothyroxine sodium products of the same strength with these therapeutic equivalence codes.

TE Code	Proprietary Name	Applicant	Strength	Appl No	RLD	RS
AB1	UNITHROID	STEVENS J	0.3MG	N021210	RLD	RS
AB2	SYNTHROID	ABBVIE	0.3MG	N021402	RLD	RS

¹⁷ In previous editions of the Orange Book, FDA provided a chart outlining therapeutic equivalence codes for all .025 mg levothyroxine sodium drug products in the Active Section of the Orange Book. FDA has decided, for ease of review, to revise the chart to identify the NDAs for the reference listed drugs for each therapeutic equivalence code (i.e., AB1, AB2, AB3, and AB4), and their corresponding reference standards, which are identified in 0.2 and 0.3 mg strengths.

¹⁸ The chart is current as of the date of publication of the annual edition. See the most current monthly cumulative supplement for updates to this information available at <https://www.fda.gov/media/72973/download>. Please consult the Active Section for information on other strengths.

AB3	LEVOXYL	KING PHARMS	0.2MG	N021301	RLD	RS
AB4	THYRO-TABS	ALVOGEN INC	0.3MG	N021116	RLD	-
AB4	LEVOTHYROXINE SODIUM ¹⁹	MYLAN	0.3MG	A076187	-	RS

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.²⁰ (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, as applicable, other applications blocked by the relevant exclusivity. If the listed drug is also protected by one or more patents, the approval date for an ANDA or 505(b)(2) application that relies on the listed drug 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

¹⁹ Alvogen INC tablets (NDA 021116) (previously known as Levothroid) previously was listed in the Discontinued Drug Product List section of the Orange Book. It is the RLD for therapeutic equivalents identified with the AB4 code. During this time, Mylan's levothyroxine product (ANDA 076187) was selected as the reference standard for ANDA applicants to use to establish bioequivalence to Thyro-Tabs. It remains 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).

²⁰ 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).

holder of the NDA may waive its exclusivity as to any or all applications that might otherwise be blocked by such exclusivity. If an NDA sponsor waives its exclusivity, qualified applications may be accepted for review and/or approved, as applicable. An NDA for which the holder has waived its exclusivity as to all 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 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, Office of Generic Drugs, Center for Drug Evaluation and Research, Food and Drug Administration, 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 generally 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 effectiveness 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 Division of Orange Book Publication and Regulatory Assessment (DOBPR) 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.²¹ 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 DOBPR 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 or 2 of the Orange Book (as discussed in Section 1.1), must be submitted to DOBPR by

²¹ See, e.g., Section 506I(d) of the FD&C Act.

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, DOBPPRA 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 listed in the Orange Book, but rather intends to apply the change prospectively to drug products as they are added to the Orange Book.

You can contact DOBPPRA by email at orangebook@fda.hhs.gov.

1.13 Availability of the 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. 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 effectiveness 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)	→	<u>AP</u>	! 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
	→	<u>AP</u>	! KENDRA PHARM	150MG/ML	A079444	001	OCT 31, 1999
APPLICANT	→						
AVAILABLE STRENGTH(S) OF A PRODUCT	→						
APPLICATION 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	→	<table border="0"> <tr> <td>TABLET; ORAL</td> <td></td> </tr> <tr> <td>HYDROCHLOROTHIAZIDE, RESERPINE AND HYDRALAZINE HCL</td> <td></td> </tr> <tr> <td>REINWALD LABS</td> <td>25MG; 15MG; 0.1MG A069808 001 JAN 18, 1982</td> </tr> </table>	TABLET; ORAL		HYDROCHLOROTHIAZIDE, RESERPINE AND HYDRALAZINE HCL		REINWALD LABS	25MG; 15MG; 0.1MG A069808 001 JAN 18, 1982
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*.

SULFASALAZINE

TABLET; ORAL

FAZINE

AB PARKLAND **500MG** **A042999** **001**

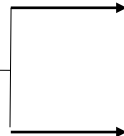
SULAZINE

AB URSA **500MG** **A042222** **001**

SULFASALAZINE

BP BROWN 500MG A041297 001

PRODUCTS CONSIDERED THERAPEUTICALLY EQUIVALENT TO EACH OTHER



PRODUCTS CONSIDERED **NOT** THERAPEUTICALLY EQUIVALENT TO ANY OTHER PRODUCTS LISTED

SULFASALAZINE

TABLET; ORAL

FAZINE

AB PARKLAND **500MG** **A042999** **001**

SULFASALAZINE

BP BROWN 500MG A041297 001

SOUTH 500MG A067627 001

PRODUCTS CONSIDERED **NOT** THERAPEUTICALLY EQUIVALENT TO EACH OTHER



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 HLTHCARE	EQ 20MG BASE/ML	N020978 001	Dec 17, 1998
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TABLET;ORAL

ABACAVIR SULFATE

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 HLTHCARE	EQ 300MG BASE	N020977 001	Dec 17, 1998
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ABACAVIR SULFATE; DOLUTEGRAVIR SODIUM; LAMIVUDINE

TABLET;ORAL

TRIUMEQ

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

EPZICOM

AB	+! VIIV HLTHCARE	EQ 600MG BASE;300MG	N021652 001	Aug 02, 2004
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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 HLTHCARE	EQ 300MG BASE;150MG;300MG	N021205 001	Nov 14, 2000

ABALOPARATIDE

SOLUTION;SUBCUTANEOUS

TYMLOS

+!	RADIUS HEALTH INC	3.12MG/1.56ML (2MG/ML)	N208743 001	Apr 28, 2017
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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		500MG	A208327 002	Dec 23, 2020
AB	APOTEX	250MG	A208453 001	Oct 31, 2018
AB	DR REDDYS LABS LTD	250MG	A208416 001	May 18, 2020
AB	GLENMARK PHARMS	250MG	A209227 001	Oct 16, 2019
AB	HIKMA	250MG	A208339 001	Oct 31, 2018
AB	MSN	250MG	A210686 001	Jul 10, 2019
AB	MYLAN	250MG	A208446 001	Oct 31, 2018
AB		500MG	A208446 002	Dec 14, 2020
AB	ONCOGEN PHARMA	250MG	A215947 001	Jan 05, 2022
AB		500MG	A215947 002	Jan 05, 2022
AB	QILU	250MG	A212462 001	Sep 27, 2019
AB		500MG	A212462 002	Jun 25, 2021
AB	RISING	250MG	A208371 001	Feb 25, 2019
AB	TEVA PHARMS USA	250MG	A208432 001	Oct 31, 2018
AB	WOCKHARDT BIO AG	250MG	A208380 001	Feb 27, 2019

PRESCRIPTION DRUG PRODUCT LIST

ABIRATERONE ACETATE

TABLET; ORAL

ZYTIGA

<u>AB</u>	+	JANSSEN BIOTECH	<u>250MG</u>	<u>N202379</u>	<u>001</u>	Apr 28, 2011
<u>AB</u>	+	YONSA	<u>500MG</u>	<u>N202379</u>	<u>002</u>	Apr 14, 2017
	+	SUN PHARMA GLOBAL	125MG	N210308	001	May 22, 2018

ACALABRUTINIB

CAPSULE; ORAL

CALQUENCE

	+	ASTRAZENECA	100MG	N210259	001	Oct 31, 2017
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ACAMPROSATE CALCIUM

TABLET, DELAYED RELEASE; ORAL

ACAMPROSATE CALCIUM

<u>AB</u>	!	GLENMARK GENERICS	<u>333MG</u>	<u>A202229</u>	<u>001</u>	Jul 16, 2013
<u>AB</u>		MYLAN	<u>333MG</u>	<u>A200142</u>	<u>001</u>	Mar 11, 2014
<u>AB</u>		ZYDUS PHARMS	<u>333MG</u>	<u>A205995</u>	<u>001</u>	May 26, 2017

ACARBOSE

TABLET; ORAL

ACARBOSE

<u>AB</u>		EMCURE PHARMS LTD	<u>25MG</u>	<u>A202271</u>	<u>001</u>	Feb 07, 2012
<u>AB</u>			<u>50MG</u>	<u>A202271</u>	<u>002</u>	Feb 07, 2012
<u>AB</u>			<u>100MG</u>	<u>A202271</u>	<u>003</u>	Feb 07, 2012
<u>AB</u>		HIKMA	<u>25MG</u>	<u>A078470</u>	<u>001</u>	May 07, 2008
<u>AB</u>			<u>50MG</u>	<u>A078470</u>	<u>002</u>	May 07, 2008
<u>AB</u>			<u>100MG</u>	<u>A078470</u>	<u>003</u>	May 07, 2008
<u>AB</u>		IMPAX LABS	<u>25MG</u>	<u>A078441</u>	<u>001</u>	May 14, 2009
<u>AB</u>			<u>50MG</u>	<u>A078441</u>	<u>002</u>	May 14, 2009
<u>AB</u>			<u>100MG</u>	<u>A078441</u>	<u>003</u>	May 14, 2009
<u>AB</u>	!	STRIDES PHARMA	<u>25MG</u>	<u>A090912</u>	<u>001</u>	Jul 27, 2011
<u>AB</u>			<u>50MG</u>	<u>A090912</u>	<u>002</u>	Jul 27, 2011
<u>AB</u>			<u>100MG</u>	<u>A090912</u>	<u>003</u>	Jul 27, 2011
<u>AB</u>		VIRTUS PHARM	<u>25MG</u>	<u>A091343</u>	<u>001</u>	Oct 17, 2013
<u>AB</u>			<u>50MG</u>	<u>A091343</u>	<u>002</u>	Oct 17, 2013
<u>AB</u>			<u>100MG</u>	<u>A091343</u>	<u>003</u>	Oct 17, 2013
<u>AB</u>		WATSON LABS	<u>25MG</u>	<u>A077532</u>	<u>001</u>	May 07, 2008
<u>AB</u>			<u>50MG</u>	<u>A077532</u>	<u>002</u>	May 07, 2008
<u>AB</u>			<u>100MG</u>	<u>A077532</u>	<u>003</u>	May 07, 2008

ACEBUTOLOL HYDROCHLORIDE

CAPSULE; ORAL

ACEBUTOLOL HYDROCHLORIDE

<u>AB</u>	!	AMNEAL PHARM	<u>EQ 200MG BASE</u>	<u>A075047</u>	<u>001</u>	Dec 30, 1999
<u>AB</u>	!		<u>EQ 400MG BASE</u>	<u>A075047</u>	<u>002</u>	Dec 30, 1999
<u>AB</u>		MYLAN	<u>EQ 200MG BASE</u>	<u>A074288</u>	<u>001</u>	Apr 24, 1995
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A074288</u>	<u>002</u>	Apr 24, 1995

ACETAMINOPHEN

SOLUTION; INTRAVENOUS

ACETAMINOPHEN

<u>AP</u>		BAXTER HLTHCARE CORP	<u>1GM/100ML (10MG/ML)</u>	<u>A214331</u>	<u>001</u>	Sep 17, 2021
<u>AP</u>		CUSTOPHARM INC	<u>1GM/100ML (10MG/ML)</u>	<u>A202605</u>	<u>001</u>	Jun 13, 2016
<u>AP</u>		EUGIA PHARMA	<u>1GM/100ML (10MG/ML)</u>	<u>A210969</u>	<u>001</u>	Oct 21, 2020
<u>AP</u>		MYLAN	<u>1GM/100ML (10MG/ML)</u>	<u>A213255</u>	<u>001</u>	Aug 07, 2020
<u>AP</u>		SANDOZ INC	<u>1GM/100ML (10MG/ML)</u>	<u>A204052</u>	<u>001</u>	Mar 22, 2016
		<u>OFIRMEV</u>				
<u>AP</u>	+	MALLINCKRODT HOSP	<u>1GM/100ML (10MG/ML)</u>	<u>N022450</u>	<u>001</u>	Nov 02, 2010
		ACETAMINOPHEN				
	+	B BRAUN MEDICAL INC	500MG/50ML (10MG/ML)	N204957	001	Feb 18, 2021
	+		1GM/100ML (10MG/ML)	N204957	002	Feb 18, 2021
		FRESENIUS KABI USA	1GM/100ML (10MG/ML)	N204767	001	Oct 28, 2015

ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE

TABLET; ORAL

APADAZ

	+	KVK TECH INC	325MG;EQ 4.08MG BASE	N208653	002	Jan 04, 2019
	+		325MG;EQ 6.12MG BASE	N208653	001	Feb 23, 2018
	+		325MG;EQ 8.16MG BASE	N208653	003	Jan 04, 2019

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; BUTALBITAL

CAPSULE;ORAL

BUTALBITAL AND ACETAMINOPHEN

AA	GRANULES	<u>300MG;50MG</u>	<u>A213115</u>	<u>001</u>	Nov 22, 2019
AA	! MAYNE PHARMA INC	<u>300MG;50MG</u>	<u>A207313</u>	<u>001</u>	Dec 27, 2017

TABLET;ORAL

BUTALBITAL AND ACETAMINOPHEN

AA	ALVOGEN	<u>300MG;50MG</u>	<u>A207635</u>	<u>001</u>	Jun 05, 2017
AA		<u>325MG;50MG</u>	<u>A205120</u>	<u>001</u>	Oct 30, 2015
AA	LARKEN LABS INC	<u>325MG;50MG</u>	<u>A203484</u>	<u>002</u>	Dec 04, 2015
AA	! LGM PHARMA	<u>300MG;50MG</u>	<u>A090956</u>	<u>001</u>	Aug 23, 2011
AA	MIKART	<u>300MG;50MG</u>	<u>A207386</u>	<u>001</u>	Nov 15, 2016
BUTAPAP					
AA	! MIKART	<u>325MG;50MG</u>	<u>A089987</u>	<u>001</u>	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	<u>325MG;50MG;40MG</u>	<u>A204733</u>	<u>001</u>	Sep 26, 2018
AA	GRANULES	<u>300MG;50MG;40MG</u>	<u>A213321</u>	<u>001</u>	Apr 08, 2020
AA	LANNETT CO INC	<u>300MG;50MG;40MG</u>	<u>A212082</u>	<u>001</u>	Dec 17, 2019
AA		<u>325MG;50MG;40MG</u>	<u>A212083</u>	<u>001</u>	Dec 17, 2019
AA	! LGM PHARMA	<u>300MG;50MG;40MG</u>	<u>A040885</u>	<u>001</u>	Nov 16, 2009
AA	MAYNE PHARMA INC	<u>300MG;50MG;40MG</u>	<u>A210817</u>	<u>001</u>	Dec 17, 2019
AA	!	<u>325MG;50MG;40MG</u>	<u>A089007</u>	<u>001</u>	Mar 17, 1986
AA	NUVO PHARMS INC	<u>300MG;50MG;40MG</u>	<u>A207118</u>	<u>001</u>	Oct 28, 2016
AA	RICONPHARMA LLC	<u>300MG;50MG;40MG</u>	<u>A215047</u>	<u>001</u>	Nov 17, 2021
AA	SENORES PHARMS	<u>300MG;50MG;40MG</u>	<u>A214087</u>	<u>001</u>	Aug 13, 2021
AA		<u>325MG;50MG;40MG</u>	<u>A214087</u>	<u>002</u>	Aug 13, 2021
AA	TARO PHARM INDS LTD	<u>300MG;50MG;40MG</u>	<u>A213046</u>	<u>001</u>	Jul 01, 2020
AA	XSPIRE PHARMA	<u>300MG;50MG;40MG</u>	<u>A206615</u>	<u>001</u>	Aug 04, 2017

SOLUTION;ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

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

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

AA	ABHAI LLC	<u>325MG;50MG;40MG</u>	<u>A211106</u>	<u>001</u>	Sep 26, 2018
AA	+ ACTAVIS LABS UT INC	<u>325MG;50MG;40MG</u>	<u>A088616</u>	<u>001</u>	Nov 09, 1984
AA	ALVOGEN	<u>325MG;50MG;40MG</u>	<u>A204984</u>	<u>001</u>	Jan 10, 2017
AA	LANNETT CO INC	<u>325MG;50MG;40MG</u>	<u>A200243</u>	<u>001</u>	Sep 13, 2012
AA	LGM PHARMA	<u>325MG;50MG;40MG</u>	<u>A209587</u>	<u>001</u>	Oct 31, 2018
AA	MIKART	<u>325MG;50MG;40MG</u>	<u>A089175</u>	<u>001</u>	Jan 21, 1987
AA	NESHER PHARMS	<u>325MG;50MG;40MG</u>	<u>A211543</u>	<u>001</u>	Jul 17, 2020
AA	! STRIDES PHARMA	<u>325MG;50MG;40MG</u>	<u>A040511</u>	<u>001</u>	Aug 27, 2003
AA		<u>325MG;50MG;40MG</u>	<u>A203647</u>	<u>001</u>	Sep 21, 2020

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE;ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE

AB	LGM PHARMA	<u>325MG;50MG;40MG;30MG</u>	<u>A076560</u>	<u>001</u>	Jun 10, 2004
AB	NOSTRUM LABS INC	<u>325MG;50MG;40MG;30MG</u>	<u>A075929</u>	<u>001</u>	Apr 22, 2002
AB	STRIDES PHARMA	<u>325MG;50MG;40MG;30MG</u>	<u>A204649</u>	<u>001</u>	Jul 08, 2020

FIORICET W/ CODEINE

AB	+! ACTAVIS LABS UT INC	<u>325MG;50MG;40MG;30MG</u>	<u>N020232</u>	<u>001</u>	Jul 30, 1992
BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE					
	LGM PHARMA	300MG;50MG;40MG;30MG	A076560	002	Jul 19, 2012

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE;ORAL

TREZIX

	XSPIRE PHARMA	320.5MG;30MG;16MG	A204785	001	Nov 26, 2014
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TABLET;ORAL

ACETAMINOPHEN, CAFFEINE AND DIHYDROCODEINE BITARTRATE

	LARKEN LABS INC	325MG;30MG;16MG	A204209	001	Sep 30, 2016
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ACETAMINOPHEN; CODEINE PHOSPHATE

SOLUTION;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

AA	AKORN	<u>120MG/5ML;12MG/5ML</u>	<u>A040119</u>	<u>001</u>	Apr 26, 1996
AA	ANDA REPOSITORY	<u>120MG/5ML;12MG/5ML</u>	<u>A089450</u>	<u>001</u>	Oct 27, 1992
AA	! PHARM ASSOC	<u>120MG/5ML;12MG/5ML</u>	<u>A087508</u>	<u>001</u>	

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

<u>AA</u>	AMNEAL PHARMS NY	<u>300MG;30MG</u>	<u>A040779 001</u>	May 29, 2008
<u>AA</u>	AUROLIFE PHARMA LLC	<u>300MG;15MG</u>	<u>A202800 001</u>	Apr 15, 2013
<u>AA</u>		<u>300MG;30MG</u>	<u>A202800 002</u>	Apr 15, 2013
<u>AA</u>		<u>300MG;60MG</u>	<u>A202800 003</u>	Apr 15, 2013
<u>AA</u>	ELITE LABS INC	<u>300MG;15MG</u>	<u>A212418 001</u>	Sep 10, 2019
<u>AA</u>		<u>300MG;30MG</u>	<u>A212418 002</u>	Sep 10, 2019
<u>AA</u>		<u>300MG;60MG</u>	<u>A212418 003</u>	Sep 10, 2019
<u>AA</u>	EYWA	<u>300MG;15MG</u>	<u>A211610 001</u>	Jun 27, 2019
<u>AA</u>		<u>300MG;30MG</u>	<u>A211610 002</u>	Jun 27, 2019
<u>AA</u>		<u>300MG;60MG</u>	<u>A211610 003</u>	Jun 27, 2019
<u>AA</u>	NOSTRUM LABS INC	<u>300MG;15MG</u>	<u>A088629 002</u>	Mar 06, 1985
<u>AA</u>		<u>300MG;30MG</u>	<u>A088629 003</u>	Mar 06, 1985
<u>AA</u>	+!	<u>300MG;60MG</u>	<u>A088629 001</u>	Mar 06, 1985
<u>AA</u>	! SPECGX LLC	<u>300MG;15MG</u>	<u>A040419 001</u>	May 31, 2001
<u>AA</u>	!	<u>300MG;30MG</u>	<u>A040419 002</u>	May 31, 2001
<u>AA</u>		<u>300MG;60MG</u>	<u>A040419 003</u>	May 31, 2001
<u>AA</u>	SUN PHARM INDS LTD	<u>300MG;30MG</u>	<u>A085868 001</u>	
<u>AA</u>		<u>300MG;60MG</u>	<u>A087083 001</u>	

ACETAMINOPHEN; HYDROCODONE BITARTRATE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<u>AA</u>	! ANDA REPOSITORY	<u>325MG/15ML;7.5MG/15ML</u>	<u>A040482 001</u>	Sep 25, 2003
<u>AA</u>	GENUS	<u>325MG/15ML;7.5MG/15ML</u>	<u>A040894 001</u>	Jul 19, 2011
<u>AA</u>	! MIKART	<u>300MG/15ML;10MG/15ML</u>	<u>A040881 001</u>	Feb 25, 2010
<u>AA</u>	PHARM ASSOC	<u>325MG/15ML;7.5MG/15ML</u>	<u>A040838 001</u>	May 10, 2013
<u>AA</u>	TRIS PHARMA INC	<u>300MG/15ML;10MG/15ML</u>	<u>A201295 002</u>	Dec 30, 2021
<u>AA</u>		<u>325MG/15ML;7.5MG/15ML</u>	<u>A201295 001</u>	Dec 30, 2021
<u>AA</u>	VISTAPHARM	<u>325MG/15ML;7.5MG/15ML</u>	<u>A200343 001</u>	Jan 25, 2012
<u>AA</u>	WES PHARMA INC	<u>325MG/15ML;7.5MG/15ML</u>	<u>A211023 001</u>	Mar 08, 2019
<u>AA</u>	! PHARM ASSOC	<u>325MG/15ML;10MG/15ML</u>	<u>A040834 001</u>	Apr 18, 2008

TABLET; ORAL

ANEXSIA 5/325

<u>AA</u>	! SPECGX LLC	<u>325MG;5MG</u>	<u>A040409 001</u>	Oct 20, 2000
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ANEXSIA 7.5/325

<u>AA</u>	! SPECGX LLC	<u>325MG;7.5MG</u>	<u>A040405 001</u>	Sep 08, 2000
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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>	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>	! ANDA REPOSITORY	<u>325MG;2.5MG</u>	<u>A040846 001</u>	Jun 09, 2010
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A040432 001</u>	Jan 22, 2003
<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>	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>	! 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>	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

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<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>	PRINSTON INC	<u>325MG; 5MG</u>	<u>A214928 001</u>	Dec 30, 2021
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A214928 002</u>	Dec 30, 2021
<u>AA</u>		<u>325MG; 10MG</u>	<u>A214928 003</u>	Dec 30, 2021
<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>	STRIDES PHARMA	<u>325MG; 5MG</u>	<u>A040655 001</u>	Jan 19, 2006
<u>AA</u>		<u>325MG; 5MG</u>	<u>A202935 002</u>	Jun 15, 2016
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A040656 001</u>	Jan 19, 2006
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A202935 003</u>	Jun 15, 2016
<u>AA</u>		<u>325MG; 10MG</u>	<u>A040355 001</u>	May 31, 2000
<u>AA</u>		<u>325MG; 10MG</u>	<u>A202935 004</u>	Jun 15, 2016
<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>	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

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

SOLUTION; ORAL

OXYCODONE AND ACETAMINOPHEN

<u>AA</u>	!	ABHAI LLC	<u>325MG/5ML; 5MG/5ML</u>	<u>A211499 001</u>	Dec 31, 2018
<u>AA</u>		NOSTRUM LABS INC	<u>325MG/5ML; 5MG/5ML</u>	<u>A201448 001</u>	Aug 26, 2021
		MIKART INC	<u>300MG/5ML; 10MG/5ML</u>	<u>A202142 001</u>	Nov 27, 2018

TABLET; ORAL

OXYCET

<u>AA</u>		SPECGX LLC	<u>325MG; 5MG</u>	<u>A087463 001</u>	Dec 07, 1983
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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
<u>AA</u>			<u>325MG; 10MG</u>	<u>A201447 004</u>	Apr 12, 2013
<u>AA</u>		ALVOGEN	<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>		CHARTWELL	<u>325MG; 5MG</u>	<u>A207834 001</u>	Aug 15, 2019

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE AND ACETAMINOPHEN

<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A207834 002</u>	Aug 15, 2019
<u>AA</u>		<u>325MG; 10MG</u>	<u>A207834 003</u>	Aug 15, 2019
<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>	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>	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

PERCO CET

<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

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

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

<u>AB</u>	ALKEM LABS LTD	<u>325MG; 37.5MG</u>	<u>A202076 001</u>	Mar 30, 2012
<u>AB</u>	AMNEAL PHARMS	<u>325MG; 37.5MG</u>	<u>A090485 001</u>	Dec 09, 2009
<u>AB</u>	AUROBINDO PHARMA LTD	<u>325MG; 37.5MG</u>	<u>A207152 001</u>	Mar 22, 2017
<u>AB</u>	MICRO LABS LTD INDIA	<u>325MG; 37.5MG</u>	<u>A201952 001</u>	Dec 14, 2012
<u>AB</u>	RISING PHARMA	<u>325MG; 37.5MG</u>	<u>A077858 001</u>	Sep 26, 2008
<u>AB</u>	SUN PHARM INDS INC	<u>325MG; 37.5MG</u>	<u>A077184 001</u>	Dec 16, 2005
<u>AB</u>	ZYDUS PHARMS USA INC	<u>325MG; 37.5MG</u>	<u>A090460 001</u>	Sep 06, 2012

ULTRACET

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

CAPSULE, EXTENDED RELEASE; ORAL

ACETAZOLAMIDE

<u>AB</u>	ACCORD HLTHCARE	<u>500MG</u>	<u>A207659 001</u>	Oct 18, 2018	
<u>AB</u>	ALEMBIC PHARMS LTD	<u>500MG</u>	<u>A210423 001</u>	Feb 19, 2019	
<u>AB</u>	CADILA	<u>500MG</u>	<u>A205301 001</u>	Jan 16, 2019	
<u>AB</u>	HERITAGE PHARMA	<u>500MG</u>	<u>A040904 001</u>	Dec 10, 2008	
<u>AB</u>	!	INDICUS PHARMA	<u>500MG</u>	<u>A090779 001</u>	Jul 14, 2011
<u>AB</u>	MICRO LABS LTD INDIA	<u>500MG</u>	<u>A207401 001</u>	Oct 01, 2020	
<u>AB</u>	NOSTRUM LABS INC	<u>500MG</u>	<u>A204691 001</u>	Mar 29, 2016	
<u>AB</u>	NOVAST LABS	<u>500MG</u>	<u>A203434 001</u>	Sep 30, 2016	

TABLET; ORAL

ACETAZOLAMIDE

<u>AB</u>	APPCO	<u>125MG</u>	<u>A211372 001</u>	Feb 22, 2021
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PRESCRIPTION DRUG PRODUCT LIST

ACETAZOLAMIDE

TABLET; ORAL

ACETAZOLAMIDE

<u>AB</u>		<u>250MG</u>	<u>A211372 002</u>	Feb 22, 2021
<u>AB</u>	BRECKENRIDGE	<u>125MG</u>	<u>A207503 001</u>	Apr 30, 2020
<u>AB</u>		<u>250MG</u>	<u>A207503 002</u>	Apr 30, 2020
<u>AB</u>	EYWA PHARMA	<u>125MG</u>	<u>A211556 001</u>	Oct 18, 2019
<u>AB</u>		<u>250MG</u>	<u>A211556 002</u>	Oct 18, 2019
<u>AB</u>	HERITAGE PHARMA	<u>125MG</u>	<u>A205530 001</u>	Oct 27, 2016
<u>AB</u>		<u>250MG</u>	<u>A205530 002</u>	Oct 27, 2016
<u>AB</u>	LANNETT	<u>250MG</u>	<u>A084840 001</u>	
<u>AB</u>	MANKIND PHARMA	<u>125MG</u>	<u>A214282 001</u>	Oct 07, 2020
<u>AB</u>		<u>250MG</u>	<u>A214282 002</u>	Oct 07, 2020
<u>AB</u>	NOVITIUM PHARMA	<u>125MG</u>	<u>A210588 001</u>	Oct 17, 2019
<u>AB</u>		<u>250MG</u>	<u>A210588 002</u>	Oct 17, 2019
<u>AB</u>	RUBICON	<u>125MG</u>	<u>A215101 001</u>	Aug 19, 2021
<u>AB</u>		<u>250MG</u>	<u>A215101 002</u>	Aug 19, 2021
<u>AB</u>	STRIDES PHARMA	<u>125MG</u>	<u>A209734 001</u>	Nov 20, 2017
<u>AB</u>		<u>250MG</u>	<u>A209734 002</u>	Nov 20, 2017
<u>AB</u>	TARO	<u>125MG</u>	<u>A040195 001</u>	May 28, 1997
<u>AB</u>	!	<u>250MG</u>	<u>A040195 002</u>	May 28, 1997

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

<u>AP</u>	EMCURE PHARMS LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A202693 001</u>	Dec 19, 2014
<u>AP</u>	HIKMA	<u>EQ 500MG BASE/VIAL</u>	<u>A040089 001</u>	Feb 28, 1995
<u>AP</u>	MYLAN ASI	<u>EQ 500MG BASE/VIAL</u>	<u>A200880 001</u>	May 09, 2012
<u>AP</u>	! XGEN PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A040784 001</u>	Dec 10, 2008
<u>AP</u>	ZYDUS PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A206533 001</u>	Apr 15, 2019

ACETIC ACID, GLACIAL

SOLUTION; IRRIGATION, URETHRAL

ACETIC ACID 0.25% IN PLASTIC CONTAINER

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

SOLUTION/DROPS; OTIC

ACETIC ACID

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

VOSOL

<u>AT</u>	AKORN	<u>2%</u>	<u>N012179 001</u>	
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ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS; OTIC

HYDROCORTISONE AND ACETIC ACID

<u>AT</u>	COSETTE	<u>2%;1%</u>	<u>A040609 001</u>	Feb 06, 2006
<u>AT</u>	TARO PHARM INDS LTD	<u>2%;1%</u>	<u>A088759 001</u>	Mar 04, 1985

VOSOL HC

<u>AT</u>	+! AKORN	<u>2%;1%</u>	<u>N012770 001</u>	
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ACETOHYDROXAMIC ACID

TABLET; ORAL

LITHOSTAT

+!	MISSION PHARMA	250MG	N018749 001	May 31, 1983
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ACETYLCHOLINE CHLORIDE

FOR SOLUTION; OPHTHALMIC

MIOCHOL-E

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

INJECTABLE; INTRAVENOUS

ACETADOTE

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

<u>AP</u>	AKORN	<u>6GM/30ML (200MG/ML)</u>	<u>A203173 001</u>	Mar 24, 2015
<u>AP</u>	EUGIA PHARMA	<u>6GM/30ML (200MG/ML)</u>	<u>A207358 001</u>	Feb 29, 2016
<u>AP</u>	EXELA PHARMA	<u>6GM/30ML (200MG/ML)</u>	<u>A204797 001</u>	Apr 15, 2021
<u>AP</u>	FRESENIUS KABI USA	<u>6GM/30ML (200MG/ML)</u>	<u>A200644 001</u>	Nov 07, 2012
<u>AP</u>	SAGENT PHARMS INC	<u>6GM/30ML (200MG/ML)</u>	<u>A091684 001</u>	Oct 31, 2017
<u>AP</u>	ZYDUS PHARMS	<u>6GM/30ML (200MG/ML)</u>	<u>A208166 001</u>	Jul 20, 2018

PRESCRIPTION DRUG PRODUCT LIST

ACETYLCYSTEINE

SOLUTION;INHALATION, ORAL

ACETYLCYSTEINE

<u>AN</u>	ALVOGEN	<u>10%</u>	<u>A204674</u>	<u>001</u>	Feb 11, 2014
<u>AN</u>		<u>20%</u>	<u>A203853</u>	<u>001</u>	Jun 21, 2012
<u>AN</u>	! AM REGENT	<u>10%</u>	<u>A072489</u>	<u>001</u>	Jul 28, 1995
<u>AN</u>	!	<u>20%</u>	<u>A072547</u>	<u>001</u>	Jul 28, 1995
<u>AN</u>	HOSPIRA	<u>10%</u>	<u>A073664</u>	<u>001</u>	Aug 30, 1994
<u>AN</u>		<u>20%</u>	<u>A074037</u>	<u>001</u>	Aug 30, 1994

ACITRETIN

CAPSULE;ORAL

ACITRETIN

<u>AB</u>	BARR LABS INC	<u>10MG</u>	<u>A091455</u>	<u>001</u>	Apr 04, 2013
<u>AB</u>		<u>25MG</u>	<u>A091455</u>	<u>002</u>	Apr 04, 2013
<u>AB</u>	IMPAX LABS INC	<u>10MG</u>	<u>A202552</u>	<u>001</u>	Dec 23, 2015
<u>AB</u>		<u>17.5MG</u>	<u>A202552</u>	<u>002</u>	Dec 23, 2015
<u>AB</u>		<u>22.5MG</u>	<u>A202552</u>	<u>003</u>	Dec 23, 2015
<u>AB</u>		<u>25MG</u>	<u>A202552</u>	<u>004</u>	Dec 23, 2015
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A202148</u>	<u>001</u>	Sep 10, 2015
<u>AB</u>		<u>25MG</u>	<u>A202148</u>	<u>002</u>	Sep 10, 2015
<u>AB</u>	SIGMAPHARM LABS LLC	<u>10MG</u>	<u>A204633</u>	<u>001</u>	May 22, 2015
<u>AB</u>		<u>17.5MG</u>	<u>A204633</u>	<u>002</u>	May 22, 2015
<u>AB</u>		<u>22.5MG</u>	<u>A204633</u>	<u>003</u>	May 22, 2015
<u>AB</u>	!	<u>25MG</u>	<u>A204633</u>	<u>004</u>	May 22, 2015
<u>AB</u>	TEVA PHARMS USA	<u>17.5MG</u>	<u>A202897</u>	<u>001</u>	Apr 04, 2013
<u>AB</u>		<u>22.5MG</u>	<u>A202897</u>	<u>002</u>	Apr 04, 2013

ACOLIDINIUM BROMIDE

POWDER, METERED;INHALATION

TUDORZA PRESSAIR

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

ACOLIDINIUM BROMIDE; FORMOTEROL FUMARATE

POWDER, METERED;INHALATION

DUAKLIR PRESSAIR

+! ASTRAZENECA 0.4MG/INH;0.012MG/INH N210595 001 Mar 29, 2019

ACYCLOVIR

CAPSULE;ORAL

ACYCLOVIR

<u>AB</u>	! APOTEX	<u>200MG</u>	<u>A075677</u>	<u>001</u>	Sep 28, 2005
<u>AB</u>	CADILA	<u>200MG</u>	<u>A204313</u>	<u>001</u>	Mar 25, 2016
<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>	HERITAGE PHARMS INC	<u>200MG</u>	<u>A074889</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>	KENTON	<u>200MG</u>	<u>A075090</u>	<u>001</u>	Jan 26, 1999
<u>AB</u>	TEVA	<u>200MG</u>	<u>A074578</u>	<u>001</u>	Apr 22, 1997
<u>AB</u>	YILING	<u>200MG</u>	<u>A212173</u>	<u>001</u>	Sep 14, 2020

CREAM;TOPICAL

ACYCLOVIR

<u>AB</u>	AMNEAL	<u>5%</u>	<u>A208766</u>	<u>001</u>	Nov 09, 2020
<u>AB</u>	! PADAGIS ISRAEL	<u>5%</u>	<u>A208702</u>	<u>001</u>	Feb 04, 2019

ZOVIRAX

<u>AB</u>	+ BAUSCH	<u>5%</u>	<u>N021478</u>	<u>001</u>	Dec 30, 2002
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OINTMENT;TOPICAL

ACYCLOVIR

<u>AB</u>	ALEMBIC 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>	APOTEX	<u>5%</u>	<u>A210774</u>	<u>001</u>	Sep 06, 2019
<u>AB</u>	CADILA	<u>5%</u>	<u>A205974</u>	<u>001</u>	Mar 15, 2019
<u>AB</u>	CIPLA	<u>5%</u>	<u>A211794</u>	<u>001</u>	Jan 18, 2019
<u>AB</u>	COSETTE	<u>5%</u>	<u>A205591</u>	<u>001</u>	Nov 13, 2017
<u>AB</u>	FOUGERA PHARMS INC	<u>5%</u>	<u>A206633</u>	<u>001</u>	May 11, 2016
<u>AB</u>	GLENMARK PHARMS SA	<u>5%</u>	<u>A205510</u>	<u>001</u>	Jul 31, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>5%</u>	<u>A212444</u>	<u>001</u>	May 19, 2021
<u>AB</u>	MYLAN PHARMS INC	<u>5%</u>	<u>A202459</u>	<u>001</u>	Apr 03, 2013
<u>AB</u>	PRINSTON INC	<u>5%</u>	<u>A212202</u>	<u>001</u>	Nov 15, 2021
<u>AB</u>	SOLARIS PHARMA CORP	<u>5%</u>	<u>A212495</u>	<u>001</u>	Apr 07, 2020
<u>AB</u>	TARO	<u>5%</u>	<u>A205469</u>	<u>001</u>	Dec 21, 2016
<u>AB</u>	TORRENT	<u>5%</u>	<u>A209971</u>	<u>001</u>	Jan 11, 2019
<u>AB</u>	XIROMED	<u>5%</u>	<u>A201501</u>	<u>001</u>	Jan 29, 2020

ZOVIRAX

<u>AB</u>	+! BAUSCH	<u>5%</u>	<u>N018604</u>	<u>001</u>	Mar 29, 1982
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PRESCRIPTION DRUG PRODUCT LIST

ACYCLOVIR

SUSPENSION; ORAL

ACYCLOVIR

AB	ACTAVIS MID ATLANTIC	200MG/5ML	A074738 001	Apr 28, 1997
AB	AKORN	200MG/5ML	A077026 001	Jun 07, 2005
AB	HERITAGE PHARMS INC	200MG/5ML	A212718 001	Apr 23, 2020
AB	NOVITIUM PHARMA	200MG/5ML	A212252 001	Jul 10, 2020
AB	VISTAPHARM	200MG/5ML	A213951 001	Jan 11, 2021

ZOVIRAX

AB	+ ! MYLAN	200MG/5ML	N019909 001	Dec 22, 1989
	TABLET; BUCCAL			
	SITAVIG			
	+ ! EPI HLTH	50MG	N203791 001	Apr 12, 2013
	TABLET; ORAL			

ACYCLOVIR

AB	APOTEX INC	400MG	A077309 001	Sep 29, 2005
AB		800MG	A077309 002	Sep 29, 2005
AB	CADILA PHARMS LTD	400MG	A202168 001	Nov 15, 2013
AB		800MG	A202168 002	Nov 15, 2013
AB	CARLSBAD	400MG	A075382 001	Apr 30, 1999
AB		800MG	A075382 002	Apr 30, 1999
AB	HERITAGE PHARMS INC	400MG	A074891 001	Oct 31, 1997
AB		800MG	A074891 002	Oct 31, 1997
AB	HETERO LABS LTD V	400MG	A203834 001	Oct 29, 2013
AB	!	800MG	A203834 002	Oct 29, 2013
AB	SQUARE PHARMS	400MG	A209366 001	Oct 07, 2019
AB		800MG	A209366 002	Oct 07, 2019
AB	STRIDES PHARMA	400MG	A074946 001	Nov 19, 1997
AB		800MG	A074946 002	Nov 19, 1997
AB	TEVA	400MG	A074556 002	Apr 22, 1997
AB		800MG	A074556 003	Apr 22, 1997
AB	YILING	400MG	A210401 001	Mar 07, 2018
AB		800MG	A210401 002	Mar 07, 2018
AB	ZYDUS PHARMS	400MG	A204314 001	Aug 19, 2014
AB		800MG	A204314 002	Aug 19, 2014

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR SODIUM

AP	EUGIA PHARMA	EQ 50MG BASE/ML	A203701 001	Oct 11, 2013
AP	! FRESENIUS KABI USA	EQ 50MG BASE/ML	A074930 001	May 13, 1998
AP	NAMIGEN LLC	EQ 50MG BASE/ML	A207919 001	Jun 17, 2020
AP	ZYDUS PHARMS	EQ 50MG BASE/ML	A206535 001	Aug 31, 2018

ACYCLOVIR; HYDROCORTISONE

CREAM; TOPICAL

XERESE

+ !	BAUSCH	5%;1%	N022436 001	Jul 31, 2009
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ADAPALENE

CREAM; TOPICAL

ADAPALENE

AB	FOUGERA PHARMS	0.1%	A090824 001	Jun 30, 2010
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DIFFERIN

AB	+ ! GALDERMA LABS LP	0.1%	N020748 001	May 26, 2000
	GEL; TOPICAL			

ADAPALENE

AB	ACTAVIS MID ATLANTIC	0.3%	A201000 001	Oct 27, 2014
AB	ALEOR	0.3%	A213508 001	Jun 18, 2020
AB	DERMACEUTICALS			
AB	TARO	0.3%	A208322 001	Jun 23, 2016

DIFFERIN

AB	+ ! GALDERMA LABS LP	0.3%	N021753 001	Jun 19, 2007
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LOTION; TOPICAL

DIFFERIN

+ !	GALDERMA LABS LP	0.1%	N022502 001	Mar 17, 2010
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SOLUTION; TOPICAL

ADAPALENE

AB	CALL INC	0.1%	A203981 001	Sep 23, 2016
AB		0.1%	A204593 001	Jan 05, 2016

PRESCRIPTION DRUG PRODUCT LIST

ADAPALENE; BENZOYL PEROXIDE

GEL; TOPICAL

ADAPALENE AND BENZOYL PEROXIDE

AB	ACTAVIS MID ATLANTIC	0.1%;2.5%	A203790 001	Sep 30, 2015
AB	GLENMARK PHARMS LTD	0.1%;2.5%	A208108 001	Nov 08, 2019
AB	PADAGIS ISRAEL	0.1%;2.5%	A205033 001	Jan 23, 2018
AB	TARO	0.1%;2.5%	A206959 001	Jan 24, 2018
AB		0.3%;2.5%	A209148 001	Oct 17, 2018

EPIDUO

AB	+ ! GALDERMA LABS LP	0.1%;2.5%	N022320 001	Dec 08, 2008
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EPIDUO FORTE

AB	+ ! GALDERMA LABS	0.3%;2.5%	N207917 001	Jul 15, 2015
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ADEFOVIR DIPIVOXIL

TABLET; ORAL

ADEFOVIR DIPIVOXIL

AB	APOTEX	10MG	A205459 001	Jul 06, 2018
AB	SIGMAPHARM LABS LLC	10MG	A202051 001	Aug 29, 2013

HEPSERA

AB	+ ! GILEAD	10MG	N021449 001	Sep 20, 2002
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ADENOSINE

INJECTABLE; INJECTION

ADENOSINE

AP	! AKORN	3MG/ML	A078076 001	Oct 31, 2008
AP	FRESENIUS KABI USA	3MG/ML	A077133 001	Apr 27, 2005
AP		3MG/ML	A205568 001	Apr 16, 2018
AP	GLAND PHARMA LTD	3MG/ML	A077283 001	Jun 14, 2007
AP		3MG/ML	A206778 001	Feb 16, 2018
AP	HIKMA	3MG/ML	A076404 001	Jun 16, 2004
AP		3MG/ML	A076500 001	Jun 16, 2004
AP	MYLAN LABS LTD	3MG/ML	A078686 001	May 13, 2009

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	EMCURE PHARMS LTD	60MG/20ML (3MG/ML)	A202313 001	Sep 15, 2014
AP		90MG/30ML (3MG/ML)	A202313 002	Sep 15, 2014
AP	EUGIA PHARMA	60MG/20ML (3MG/ML)	A205331 001	Nov 02, 2017
AP		90MG/30ML (3MG/ML)	A205331 002	Nov 02, 2017
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

AFAMELANOTIDE

IMPLANT; SUBCUTANEOUS

SCENESSE

+ !	CLIVUNEL INC	16MG	N210797 001	Oct 08, 2019
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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

AIR POLYMER-TYPE A

FOAM; INTRAUTERINE

EXEM FOAM KIT

+ !	GISKIT	10ML	N212279 001	Nov 07, 2019
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ALBENDAZOLE

TABLET; ORAL

ALBENDAZOLE

AB	ACTAVIS ELIZABETH	200MG	A208094 001	May 20, 2019
AB	CIPLA LTD	200MG	A210434 001	Sep 21, 2018
AB	DR REDDYS	200MG	A211034 001	Jan 26, 2021
AB	EDENBRIDGE PHARMS	200MG	A211117 001	May 14, 2019
AB	MSN	200MG	A213435 001	Jan 21, 2021

PRESCRIPTION DRUG PRODUCT LIST

ALBENDAZOLE

TABLET; ORAL

ALBENDAZOLE

AB	STRIDES PHARMA	200MG	A210011 001	Dec 07, 2018
AB	ZYDUS PHARMS	200MG	A208979 001	Dec 14, 2018

ALBENZA

AB	+! IMPAX LABS INC	200MG	N020666 001	Jun 11, 1996
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ALBUMIN HUMAN

INJECTABLE; INJECTION

OPTISON

+!	GE HEALTHCARE	10MG/ML	N020899 001	Dec 31, 1997
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ALBUTEROL SULFATE

AEROSOL, METERED; INHALATION

ALBUTEROL SULFATE

AB1	CIPLA	EQ 0.09MG BASE/INH	A209959 001	Apr 08, 2020
AB1	SANDOZ INC	EQ 0.09MG BASE/INH	A207085 001	Jun 01, 2021

PROVENTIL-HFA

AB1	+! KINDEVA	EQ 0.09MG BASE/INH	N020503 001	Aug 15, 1996
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ALBUTEROL SULFATE

AB2	LUPIN	EQ 0.09MG BASE/INH	A209954 001	Aug 24, 2020
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PROAIR HFA

AB2	+! TEVA BRANDED PHARM	EQ 0.09MG BASE/INH	N021457 001	Oct 29, 2004
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VENTOLIN HFA

BX	+! GLAXOSMITHKLINE	EQ 0.09MG BASE/INH	N020983 001	Apr 19, 2001
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POWDER, METERED; INHALATION

PROAIR DIGIHALER

+	TEVA BRANDED PHARM	EQ 0.09MG BASE/INH	N205636 002	Dec 21, 2018
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PROAIR RESPICLICK

+!	TEVA BRANDED PHARM	EQ 0.09MG BASE/INH	N205636 001	Mar 31, 2015
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SOLUTION; INHALATION

ACCUNEB

AN	+! MYLAN SPECIALITY LP	EQ 0.021% BASE	N020949 002	Apr 30, 2001
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AN	+!	EQ 0.042% BASE	N020949 001	Apr 30, 2001
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ALBUTEROL SULFATE

AN	AKORN	EQ 0.5% BASE	A074543 001	Jan 15, 1998
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AN	LUOXIN AUROVITAS	EQ 0.083% BASE	A206224 001	Oct 17, 2017
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AN	NEPHRON	EQ 0.021% BASE	A076355 002	Mar 31, 2010
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AN		EQ 0.042% BASE	A076355 001	Jun 28, 2004
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AN	!	EQ 0.083% BASE	A074880 001	Sep 17, 1997
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AN	!	EQ 0.5% BASE	A075664 001	Jun 26, 2001
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AN	RITEDOSE CORP	EQ 0.021% BASE	A214531 001	Dec 28, 2021
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AN		EQ 0.042% BASE	A214531 002	Dec 28, 2021
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AN		EQ 0.083% BASE	A077839 001	Dec 16, 2008
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AN	SUN PHARM	EQ 0.083% BASE	A207857 001	Jul 21, 2017
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AN	WATSON LABS	EQ 0.021% BASE	A077772 001	Sep 25, 2007
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AN		EQ 0.042% BASE	A077772 002	Sep 25, 2007
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SYRUP; ORAL

ALBUTEROL SULFATE

AA	AKORN	EQ 2MG BASE/5ML	A074749 001	Jan 30, 1998
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AA	AMNEAL PHARMS	EQ 2MG BASE/5ML	A079241 001	May 12, 2010
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AA	COSETTE	EQ 2MG BASE/5ML	A074454 001	Sep 25, 1995
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AA	LANNETT CO INC	EQ 2MG BASE/5ML	A078105 001	Dec 27, 2006
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AA	QUAGEN	EQ 2MG BASE/5ML	A212197 001	Sep 06, 2019
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AA	! TEVA	EQ 2MG BASE/5ML	A073419 001	Mar 30, 1992
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TABLET; ORAL

ALBUTEROL SULFATE

AB	AMNEAL PHARMS CO	EQ 2MG BASE	A208804 001	May 21, 2018
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AB		EQ 4MG BASE	A208804 002	May 21, 2018
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AB	APPCO	EQ 2MG BASE	A210948 001	Mar 15, 2019
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AB		EQ 4MG BASE	A210948 002	Mar 15, 2019
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AB	MYLAN	EQ 2MG BASE	A072894 002	Jan 17, 1991
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AB	!	EQ 4MG BASE	A072894 001	Jan 17, 1991
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AB	RISING	EQ 2MG BASE	A207046 001	Jun 29, 2018
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AB		EQ 4MG BASE	A207046 002	Jun 29, 2018
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AB	SUN PHARM	EQ 2MG BASE	A072637 002	Dec 05, 1989
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INDUSTRIES

AB		EQ 4MG BASE	A072637 001	Dec 05, 1989
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AB	VIRTUS PHARM	EQ 2MG BASE	A211397 001	Oct 26, 2018
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AB		EQ 4MG BASE	A211397 002	Oct 26, 2018
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AB	ZYDUS PHARMS	EQ 2MG BASE	A208884 001	Oct 22, 2020
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AB		EQ 4MG BASE	A208884 002	Oct 22, 2020
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PRESCRIPTION DRUG PRODUCT LIST

ALBUTEROL SULFATE

TABLET, EXTENDED RELEASE;ORAL

VOSPIRE ER

STRIDES PHARMA

EQ 4MG BASE

A076130 002 Sep 26, 2002

ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

SOLUTION; INHALATION

ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE**AN** CIPLA**EQ 0.083% BASE;0.017%****A077559 001** Dec 31, 2007**AN** ! NEPHRON**EQ 0.083% BASE;0.017%****A076749 001** Dec 31, 2007**AN** RITEDOSE CORP**EQ 0.083% BASE;0.017%****A202496 001** Oct 01, 2012**AN** SUN PHARM**EQ 0.083% BASE;0.017%****A207875 001** Aug 07, 2017

SPRAY, METERED; INHALATION

COMBIVENT RESPIMAT

+! BOEHRINGER

EQ 0.1MG BASE/INH;0.02MG/INH

N021747 001 Oct 07, 2011

INGELHEIM

ALCLOMETASONE DIPROPIONATE

CREAM; TOPICAL

ALCLOMETASONE DIPROPIONATE**AB** ! FOUGERA PHARMS**0.05%****A076973 001** Jul 12, 2005**AB** GLENMARK GENERICS**0.05%****A079061 001** Jun 23, 2009**AB** TARO**0.05%****A076587 001** Sep 15, 2005

OINTMENT; TOPICAL

ALCLOMETASONE DIPROPIONATE**AB** ! FOUGERA PHARMS**0.05%****A076884 001** Jul 18, 2005**AB** GLENMARK GENERICS**0.05%****A079227 001** Jul 30, 2009**AB** TARO**0.05%****A076730 001** Jul 29, 2004ALCOHOL

SOLUTION; INTRA-ARTERIAL

ABLYSINOL

+ BELCHER

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

! HIKMA

EQ 70MG BASE/75ML

A090520 001 Feb 25, 2013

TABLET; ORAL

ALENDRONATE SODIUM**AB** APOTEX**EQ 5MG BASE****A077982 001** Aug 04, 2008**AB****EQ 10MG BASE****A077982 002** Aug 04, 2008**AB****EQ 35MG BASE****A077982 003** Aug 04, 2008**AB****EQ 70MG BASE****A077982 004** Aug 04, 2008**AB**

AUROBINDO PHARMA

EQ 10MG BASE**A090124 001** Aug 04, 2008**AB****EQ 35MG BASE****A090124 002** Aug 04, 2008**AB****EQ 70MG BASE****A090124 003** Aug 04, 2008**AB**

CIPLA

EQ 5MG BASE**A076768 001** Aug 04, 2008**AB****EQ 10MG BASE****A076768 002** Aug 04, 2008**AB****EQ 35MG BASE****A076768 003** Aug 04, 2008**AB****EQ 40MG BASE****A076768 004** Aug 04, 2008**AB****EQ 70MG BASE****A076768 005** Aug 04, 2008**AB**

HANGZHOU BINJIANG

EQ 5MG BASE**A090258 001** Sep 24, 2009**AB****EQ 10MG BASE****A090258 002** Sep 24, 2009**AB****EQ 35MG BASE****A090258 003** Sep 24, 2009**AB****EQ 70MG BASE****A090258 004** Sep 24, 2009**AB**

SUN PHARM

EQ 5MG BASE**A090022 001** Sep 10, 2008**AB****EQ 10MG BASE****A090022 002** Sep 10, 2008**AB****EQ 35MG BASE****A090022 003** Sep 10, 2008**AB****EQ 70MG BASE****A090022 004** Sep 10, 2008**AB**

WATSON LABS

EQ 35MG BASE**A076984 001** Aug 04, 2008**AB****EQ 40MG BASE****A076984 002** Aug 04, 2008**AB****EQ 70MG BASE****A076984 003** Aug 04, 2008FOSAMAX**AB** +! MERCK AND CO INC**EQ 70MG BASE****N020560 005** Oct 20, 2000

TABLET, EFFERVESCENT; ORAL

BINOSTO

+! ASCEND THERAPS US

EQ 70MG BASE

N202344 001 Mar 12, 2012

PRESCRIPTION DRUG PRODUCT LIST

ALENDRONATE SODIUM; CHOLECALCIFEROL

TABLET; ORAL

FOSAMAX PLUS D

+ ORGANON

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**AP** +! AKORN**EQ 0.5MG BASE/ML****N019353 001** Dec 29, 1986ALFENTANIL**AP** HOSPIRA**EQ 0.5MG BASE/ML****A075221 001** Oct 28, 1999ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ALFUZOSIN HYDROCHLORIDE**AB** APOTEX INC**10MG****A079013 001** Jul 18, 2011**AB** AUROBINDO PHARMA**10MG****A079060 001** Aug 30, 2012

LTD

AB INVAGEN PHARMS**10MG****A090284 001** Jan 17, 2012**AB** SUN PHARM**10MG****A079057 001** Jul 18, 2011**AB** UNICHEM**10MG****A203192 001** Jan 28, 2016UROXATRAL**AB** +! CONCORDIA**10MG****N021287 001** Jun 12, 2003ALISKIREN HEMIFUMARATE

TABLET; ORAL

ALISKIREN HEMIFUMARATE**AB** ANCHEN PHARMS**EQ 150MG BASE****A206665 001** Mar 22, 2019**AB****EQ 300MG BASE****A206665 002** Mar 22, 2019TEKTURNA**AB** + NODEN PHARMA**EQ 150MG BASE****N021985 001** Mar 05, 2007**AB** +!**EQ 300MG BASE****N021985 002** Mar 05, 2007ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

TEKTURNA HCT

+! NODEN PHARMA

EQ 300MG BASE;12.5MG

N022107 003 Jan 18, 2008

+!

EQ 300MG BASE;25MG

N022107 004 Jan 18, 2008

ALITRETINOIN

GEL; TOPICAL

PANRETIN

+! CONCORDIA

EQ 0.1% BASE

N020886 001 Feb 02, 1999

ALLOPURINOL

TABLET; ORAL

ALLOPURINOL**AB** ACCORD HLTHCARE**100MG****A203154 001** May 06, 2013**AB****300MG****A203154 002** May 06, 2013**AB** CHARTWELL**100MG****A077353 001** Sep 08, 2005**AB****300MG****A077353 002** Sep 08, 2005**AB** INDOCO**100MG****A204467 001** Jul 28, 2016**AB****300MG****A204467 002** Jul 28, 2016**AB** IPCA LABS LTD**100MG****A090637 001** Mar 16, 2011**AB****300MG****A090637 002** Mar 16, 2011**AB** MYLAN**100MG****A018659 001** Oct 24, 1986**AB****300MG****A018659 002** Oct 24, 1986**AB** NORTHSTAR HLTHCARE**100MG****A078253 001** Sep 11, 2007**AB****300MG****A078253 002** Sep 11, 2007**AB** SUN PHARM**100MG****A071450 001** Jan 09, 1987

INDUSTRIES

AB**300MG****A071450 001** Jan 09, 1987**AB** UNICHEM**100MG****A211820 001** Mar 12, 2019**AB****300MG****A211820 002** Mar 12, 2019**AB** VINTAGE PHARMS**100MG****A075798 001** Jun 27, 2003**AB****300MG****A075798 002** Jun 27, 2003**AB** WATSON LABS**100MG****N018832 002** Sep 28, 1984**AB****300MG****N018877 001** Sep 28, 1984**AB** ZYDUS PHARMS**100MG****A210117 001** Oct 12, 2017**AB****300MG****A210117 002** Oct 12, 2017LOPURIN**AB** DR REDDYS LA**100MG****A071586 001** Apr 02, 1987**AB****300MG****A071587 001** Apr 02, 1987

PRESCRIPTION DRUG PRODUCT LIST

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 CANADA INC	2 IU/ML; 40MG/ML; 12MCG/ML; 40 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	N021163 001	May 18, 2000
+	!		2 IU/ML; 40MG/ML; 12MCG/ML; 40 IU/ML; 1MCG/ML; 3MG/ML; 120MCG/ML; 8MG/ML; 1 .2MG/ML; 0.72MG/ML; 1.2MG/ML; 660 IU/ML; 30MCG/ML	N021163 002	Jun 16, 2003

ALPRAZOLAM

CONCENTRATE; ORAL

ALPRAZOLAM

! HIKMA

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	<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>		NATCO	<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>		NOVITIUM PHARMA	<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>		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>		STRIDES PHARMA	<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>AB</u>		SUN PHARM	<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

XANAX

<u>AB</u>	+	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

ALPRAZOLAM

<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

PRESCRIPTION DRUG PRODUCT LIST

ALPRAZOLAM

TABLET, EXTENDED RELEASE;ORAL

ALPRAZOLAM

<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>	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>XANAX XR</u>				
<u>AB</u>	+	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>	HIKMA	<u>0.5MG/ML</u>	<u>A074815 001</u>	Jan 20, 1998
<u>AP</u>	TEVA PHARMS USA	<u>0.5MG/ML</u>	<u>A075196 001</u>	Apr 30, 1999

CAVERJECT

<u>AP</u>	+	PFIZER	<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>	+	PFIZER	<u>0.5MG/ML</u>	<u>N018484 001</u>
CAVERJECT IMPULSE				
		PFIZER	0.01MG/VIAL	N021212 001 Jun 11, 2002
			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

ALVIMOPAN

CAPSULE;ORAL

ALVIMOPAN

<u>AB</u>	WATSON LABS TEVA	<u>12MG</u>	<u>A208295 001</u>	Dec 19, 2019
<u>ENTEREG</u>				
<u>AB</u>	+	CUBIST PHARMS	<u>12MG</u>	<u>N021775 001</u> May 20, 2008

AMANTADINE HYDROCHLORIDE

CAPSULE;ORAL

AMANTADINE HYDROCHLORIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>100MG</u>	<u>A208966 001</u>	Jun 21, 2017
<u>AB</u>	!	BIONPHARMA INC	<u>100MG</u>	<u>A078720 001</u> May 29, 2008
<u>AB</u>	HERITAGE PHARMA	<u>100MG</u>	<u>A209171 001</u>	Jun 12, 2017
<u>AB</u>	NOVELGENIX THERAPS	<u>100MG</u>	<u>A210129 001</u>	Mar 02, 2020
<u>AB</u>	RUBICON	<u>100MG</u>	<u>A212044 001</u>	May 21, 2020

PRESCRIPTION DRUG PRODUCT LIST

AMANTADINE HYDROCHLORIDE

CAPSULE;ORAL

AMANTADINE HYDROCHLORIDE

AB	SANDOZ	100MG	A071293 001	Feb 18, 1987
AB	UPSHER SMITH LABS	100MG	A070589 001	Aug 05, 1986
AB	WATSON LABS INC	100MG	A208107 001	Dec 06, 2016
AB	ZYDUS PHARMS	100MG	A208278 001	May 31, 2016
BX	STRIDES PHARMA	100MG	A209047 001	Jun 07, 2017

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	!	AKORN	50MG/5ML	A074170 001	Oct 28, 1994
AA	!	ANDA REPOSITORY	50MG/5ML	A074028 001	Jun 28, 1993
AA	!	CMP PHARMA INC	50MG/5ML	A075819 001	Sep 11, 2002
AA		NOVELGENIX THERAPS	50MG/5ML	A214178 001	Aug 20, 2021
AA	!	PHARM ASSOC	50MG/5ML	A074509 001	Jul 17, 1995

TABLET;ORAL

AMANTADINE HYDROCHLORIDE

AB		ALEMBIC PHARMS LTD	100MG	A214284 001	Oct 15, 2020
AB		JUBILANT GENERICS	100MG	A210403 001	Feb 07, 2018
AB		NOVELGENIX THERAPS	100MG	A210215 001	Mar 10, 2020
AB	!	UPSHER SMITH LABS	100MG	A076186 001	Dec 16, 2002
AB		WATSON LABS INC	100MG	A208096 001	Dec 15, 2016
BX		STRIDES PHARMA	100MG	A209035 001	Jun 09, 2017

TABLET, EXTENDED RELEASE;ORAL

OSMOLEX ER

+	ADAMAS PHARMA	EQ 129MG BASE	N209410 001	Feb 16, 2018
+		EQ 193MG BASE	N209410 002	Feb 16, 2018

AMBRISENTAN

TABLET;ORAL

AMBRISENTAN

AB		CIPLA	5MG	A210715 001	Apr 26, 2019
AB			10MG	A210715 002	Apr 26, 2019
AB		MYLAN	5MG	A208441 001	Mar 28, 2019
AB			10MG	A208441 002	Mar 28, 2019
AB		PAR PHARM INC	5MG	A209509 001	Apr 10, 2019
AB			10MG	A209509 002	Apr 10, 2019
AB		SIGMAPHARM LABS LLC	5MG	A208354 001	Apr 10, 2019
AB			10MG	A208354 002	Apr 10, 2019
AB		SUN PHARM	5MG	A210784 001	Mar 28, 2019
AB			10MG	A210784 002	Mar 28, 2019
AB		WATSON LABS INC	5MG	A208252 001	Mar 28, 2019
AB			10MG	A208252 002	Mar 28, 2019
AB		ZYDUS PHARMS	5MG	A210058 001	Mar 28, 2019
AB			10MG	A210058 002	Mar 28, 2019

LETAIRIS

AB	+	GILEAD	5MG	N022081 001	Jun 15, 2007
AB	+	!	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
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OINTMENT;TOPICAL

AMCINONIDE

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

AMIFAMPRIDINE

TABLET;ORAL

RUZURGI

+	!	JACOBUS PHARM CO INC	10MG	N209321 001	May 06, 2019
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PRESCRIPTION DRUG PRODUCT LIST

AMIFAMPRIDINE PHOSPHATE

TABLET; ORAL

FIRDAPSE

+! CATALYST PHARMS EQ 10MG BASE N208078 001 Nov 28, 2018

AMIFOSTINE

INJECTABLE; INJECTION

AMIFOSTINEAP SUN PHARM 500MG/VIAL A077126 001 Mar 14, 2008ETHYOLAP +! CLINIGEN 500MG/VIAL N020221 001 Dec 08, 1995AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATEAP ! EMCURE PHARMS LTD EQ 250MG BASE/ML A204040 001 Dec 12, 2013AP FRESENIUS KABI USA EQ 50MG BASE/ML A205605 001 Dec 09, 2015AP EQ 250MG BASE/ML A205604 001 Dec 09, 2015AP ! HIKMA EQ 50MG BASE/ML A063313 001 Apr 11, 1994AP EQ 250MG BASE/ML A063315 001 Apr 11, 1994AP SAGENT PHARMS INC EQ 250MG BASE/ML A203323 001 May 12, 2016AP TEVA PHARMS USA EQ 250MG BASE/ML A064045 002 Sep 28, 1993

SUSPENSION, LIPOSOMAL; INHALATION

ARIKAYCE KIT

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

AMILORIDE HYDROCHLORIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDEAB ! PAR PHARM 5MG A070346 001 Jan 22, 1986AB SIGMAPHARM LABS LLC 5MG A079133 001 Jan 30, 2009AB USPHARMA WINDLAS 5MG A204180 001 Aug 07, 2015MIDAMORAB + PADAGIS US 5MG N018200 001AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDEAB BARR EQ 5MG ANHYDROUS; 50MG A071111 001 May 10, 1988AB ! RISING PHARMA EQ 5MG ANHYDROUS; 50MG A073209 001 Oct 31, 1991AMINO ACIDS

INJECTABLE; INJECTION

AMINO ACIDS

B BRAUN 15% (150GM/1000ML) A091112 001 Apr 13, 2012

15% (300GM/2000ML) A091112 002 Apr 13, 2012

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

TRAVASOL 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE 10% (10GM/100ML) N018931 003 Aug 23, 1984

PRESCRIPTION DRUG PRODUCT LIST

AMINO ACIDS

INJECTABLE; INJECTION

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

PRESCRIPTION DRUG PRODUCT LIST

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

EMULSION; INTRAVENOUS

PERIKABIVEN IN PLASTIC CONTAINER

+	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; 3.5GM/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
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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	
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AMINOCAPROIC ACID

INJECTABLE; INJECTION

AP	AMINOCAPROIC ACID	250MG/ML	A071192 001	Dec 01, 1987
	LUITPOLD			
AP	AMINOCAPROIC ACID IN PLASTIC CONTAINER	250MG/ML	A070010 001	Mar 09, 1987
	HOSPIRA			
	SOLUTION; ORAL			
AA	AMICAR	0.25GM/ML	N015230 002	
	AKORN			
AA	AMNEAL	0.25GM/ML	A212780 001	Aug 23, 2019
AA	BELCHER	0.25GM/ML	A213825 001	Apr 08, 2021
AA	LEADING PHARMA LLC	0.25GM/ML	A214140 001	Jan 26, 2021
AA	TULEX PHARMS INC	0.25GM/ML	A212494 001	Aug 11, 2020
AA	VISTAPHARM	0.25GM/ML	A212814 001	Feb 26, 2020
	TABLET; ORAL			
AB	AMICAR	500MG	N015197 001	
AB	AMICAR	1GM	N015197 002	Jun 24, 2004
AB	AMNEAL	500MG	A212492 001	Nov 26, 2019
AB	ANI PHARMS	500MG	A211629 001	Dec 14, 2020
AB	LEADING PHARMA LLC	500MG	A213928 001	Feb 12, 2021
AB	LEADING PHARMA LLC	1GM	A213928 002	Feb 12, 2021

PRESCRIPTION DRUG PRODUCT LIST

AMINOCAPROIC ACID

TABLET; ORAL

AMINOCAPROIC ACID

<u>AB</u>	MSN	<u>500MG</u>	<u>A212938</u>	<u>001</u>	Nov 06, 2020
<u>AB</u>	SUNNY	<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
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GEL; TOPICAL

AMELUZ

+	!	BIOFRONTERA	10%	N208081	001	May 10, 2016
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SOLUTION; TOPICAL

LEVULAN

+	!	DUSA	20%	N020965	001	Dec 03, 1999
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AMINOPHYLLINE

INJECTABLE; INJECTION

AMINOPHYLLINE

!		HOSPIRA	25MG/ML	A087242	001	Oct 26, 1983
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AMINOSALICYLIC ACID

GRANULE, DELAYED RELEASE; ORAL

PASER

!		JACOBUS	4GM/PACKET	A074346	001	Jun 30, 1994
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AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

<u>AP</u>	ACELLA	<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	
<u>AP</u>	EUGIA PHARMA	<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 INC	<u>50MG/ML</u>	<u>A203884</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
	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>	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>	RUBICON	<u>100MG</u>	<u>A078578</u>	<u>002</u>	Feb 26, 2021	
<u>AB</u>		<u>200MG</u>	<u>A078578</u>	<u>001</u>	Nov 06, 2008	
<u>AB</u>		<u>400MG</u>	<u>A078578</u>	<u>003</u>	Feb 26, 2021	
<u>AB</u>	TARO	<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>	UNICHEM	<u>200MG</u>	<u>A213446</u>	<u>001</u>	Jul 21, 2020	
<u>AB</u>	!	UPSHER SMITH LABS	<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>	ZYDUS PHARMS USA INC	<u>200MG</u>	<u>A079029</u>	<u>001</u>	Sep 16, 2008	
	<u>PACERONE</u>					
<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	
<u>AB</u>		<u>400MG</u>	<u>A075135</u>	<u>003</u>	Jul 02, 2020	
	AMIODARONE HYDROCHLORIDE					
	TARO	300MG	A076362	002	Dec 02, 2003	

PRESCRIPTION DRUG PRODUCT LIST

AMISULPRIDE

SOLUTION; INTRAVENOUS

BARHEMSYS

+! ACACIA

5MG/2ML (2.5MG/ML)

N209510 001 Feb 26, 2020

+!

10MG/4ML (2.5MG/ML)

N209510 002 Sep 01, 2020

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>	MANKIND PHARMA	<u>10MG</u>	<u>A213999 001</u>	Feb 19, 2021
<u>AB</u>		<u>25MG</u>	<u>A213999 002</u>	Feb 19, 2021
<u>AB</u>		<u>50MG</u>	<u>A213999 003</u>	Feb 19, 2021
<u>AB</u>		<u>75MG</u>	<u>A213999 004</u>	Feb 19, 2021
<u>AB</u>		<u>100MG</u>	<u>A213999 005</u>	Feb 19, 2021
<u>AB</u>		<u>150MG</u>	<u>A213999 006</u>	Feb 19, 2021
<u>AB</u>	+ SANDOZ	<u>10MG</u>	<u>A085968 004</u>	
<u>AB</u>	+!	<u>25MG</u>	<u>A085968 002</u>	
<u>AB</u>	+	<u>50MG</u>	<u>A085968 001</u>	
<u>AB</u>	+	<u>75MG</u>	<u>A085968 006</u>	
<u>AB</u>	+	<u>100MG</u>	<u>A085968 003</u>	
<u>AB</u>	+	<u>150MG</u>	<u>A085968 005</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>	UNICHEM	<u>10MG</u>	<u>A214548 001</u>	May 19, 2021
<u>AB</u>		<u>25MG</u>	<u>A214548 002</u>	May 19, 2021
<u>AB</u>		<u>50MG</u>	<u>A214548 003</u>	May 19, 2021
<u>AB</u>		<u>75MG</u>	<u>A214548 004</u>	May 19, 2021
<u>AB</u>		<u>100MG</u>	<u>A214548 005</u>	May 19, 2021
<u>AB</u>		<u>150MG</u>	<u>A214548 006</u>	May 19, 2021
<u>AB</u>	UPSHER SMITH LABS	<u>10MG</u>	<u>A212654 002</u>	Sep 29, 2021
<u>AB</u>		<u>25MG</u>	<u>A212654 001</u>	Apr 07, 2020
<u>AB</u>		<u>50MG</u>	<u>A212654 003</u>	Sep 29, 2021
<u>AB</u>		<u>75MG</u>	<u>A212654 004</u>	Sep 29, 2021
<u>AB</u>		<u>100MG</u>	<u>A212654 005</u>	Sep 29, 2021
<u>AB</u>		<u>150MG</u>	<u>A212654 006</u>	Sep 29, 2021
<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	<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

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BENZOATE

SUSPENSION; ORAL

KATERZIA

+! AZURITY

EQ 1MG BASE/ML

N211340 001 Jul 08, 2019

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>	CHARTWELL RX	<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>	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
<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>	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>	ORBION PHARMS	<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 PHARMA	<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 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>	UNICHEM	<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>	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
<u>NORVASC</u>				
<u>AB</u>	+ UPJOHN	<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

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM

TABLET; ORAL

AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM

AB	APOTEX	<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A205199 001</u>	Nov 18, 2019
AB		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A205199 002</u>	Nov 18, 2019
AB		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A205199 003</u>	Nov 18, 2019
AB		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A205199 004</u>	Nov 18, 2019
AB		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A205199 005</u>	Nov 18, 2019
AB		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A205199 006</u>	Nov 18, 2019
AB		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A205199 007</u>	Nov 18, 2019
AB		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A205199 008</u>	Nov 18, 2019
AB	DR REDDYS	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<u>A203874 001</u>	Mar 07, 2014
AB		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<u>A203874 002</u>	Mar 07, 2014
AB		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<u>A203874 003</u>	Mar 07, 2014
AB		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A203874 004</u>	Mar 07, 2014
AB		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A203874 005</u>	Mar 07, 2014
AB		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A203874 006</u>	Mar 07, 2014
AB		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A203874 007</u>	Mar 07, 2014
AB		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A203874 008</u>	Mar 07, 2014
AB		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A203874 009</u>	Mar 07, 2014
AB		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A203874 010</u>	Mar 07, 2014
AB		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A203874 011</u>	Mar 07, 2014
AB	MYLAN	<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A200465 004</u>	Nov 29, 2013
AB		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A200465 005</u>	Nov 29, 2013
AB		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A200465 006</u>	Nov 29, 2013
AB		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A200465 007</u>	Nov 29, 2013
AB		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A200465 008</u>	Nov 29, 2013
AB		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A200465 009</u>	Nov 29, 2013
AB		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A200465 010</u>	Nov 29, 2013
AB		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A200465 011</u>	Nov 29, 2013
AB	ZYDUS PHARMS	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<u>A207762 001</u>	Jan 11, 2019
AB		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<u>A207762 002</u>	Jan 11, 2019
AB		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<u>A207762 003</u>	Jan 11, 2019
AB		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A207762 004</u>	Jan 11, 2019
AB		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A207762 005</u>	Jan 11, 2019
AB		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A207762 006</u>	Jan 11, 2019
AB		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A207762 007</u>	Jan 11, 2019
AB		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A207762 008</u>	Jan 11, 2019
AB		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A207762 009</u>	Jan 11, 2019
AB		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A207762 010</u>	Jan 11, 2019
AB		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A207762 011</u>	Jan 11, 2019
CADUET				
AB	+ PHARMACIA	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<u>N021540 009</u>	Jul 29, 2004
AB	+	<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<u>N021540 010</u>	Jul 29, 2004
AB	+	<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<u>N021540 011</u>	Jul 29, 2004
AB	+	<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>N021540 001</u>	Jan 30, 2004
AB	+	<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>N021540 002</u>	Jan 30, 2004
AB	+	<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>N021540 003</u>	Jan 30, 2004
AB	+	<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>N021540 004</u>	Jan 30, 2004
AB	+	<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>N021540 005</u>	Jan 30, 2004
AB	+	<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>N021540 006</u>	Jan 30, 2004
AB	+	<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>N021540 007</u>	Jan 30, 2004
AB	+	<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

AB	APOTEX	<u>EQ 2.5MG BASE;10MG</u>	<u>A091431 001</u>	Dec 30, 2013
AB		<u>EQ 5MG BASE;10MG</u>	<u>A091431 002</u>	Dec 30, 2013
AB		<u>EQ 5MG BASE;20MG</u>	<u>A091431 003</u>	Dec 30, 2013
AB		<u>EQ 5MG BASE;40MG</u>	<u>A091431 004</u>	Dec 30, 2013
AB		<u>EQ 10MG BASE;20MG</u>	<u>A091431 005</u>	Dec 30, 2013
AB		<u>EQ 10MG BASE;40MG</u>	<u>A091431 006</u>	Dec 30, 2013
AB	AUROBINDO PHARMA LTD	<u>EQ 2.5MG BASE;10MG</u>	<u>A202239 001</u>	Sep 05, 2012
AB		<u>EQ 5MG BASE;10MG</u>	<u>A202239 002</u>	Sep 05, 2012
AB		<u>EQ 5MG BASE;20MG</u>	<u>A202239 003</u>	Sep 05, 2012
AB		<u>EQ 5MG BASE;40MG</u>	<u>A202239 004</u>	Sep 05, 2012
AB		<u>EQ 10MG BASE;20MG</u>	<u>A202239 005</u>	Sep 05, 2012
AB		<u>EQ 10MG BASE;40MG</u>	<u>A202239 006</u>	Sep 05, 2012
AB	DR REDDYS LABS INC	<u>EQ 2.5MG BASE;10MG</u>	<u>A077183 001</u>	Apr 15, 2010
AB		<u>EQ 5MG BASE;10MG</u>	<u>A077183 002</u>	Apr 15, 2010
AB		<u>EQ 5MG BASE;20MG</u>	<u>A077183 003</u>	Apr 15, 2010

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE;ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

<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>	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

LOTREL

<u>AB</u>	+	NOVARTIS	<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; 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	<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
<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

TRIBENZOR

<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>EQ 5MG BASE;12.5MG;160MG</u>	<u>A206180 001</u>	Dec 19, 2017
<u>AB</u>			<u>EQ 5MG BASE;25MG;160MG</u>	<u>A206180 002</u>	Dec 19, 2017
<u>AB</u>			<u>EQ 10MG BASE;12.5MG;160MG</u>	<u>A206180 003</u>	Dec 19, 2017
<u>AB</u>			<u>EQ 10MG BASE;25MG;160MG</u>	<u>A206180 004</u>	Dec 19, 2017
<u>AB</u>			<u>EQ 10MG BASE;25MG;320MG</u>	<u>A206180 005</u>	Dec 19, 2017
<u>AB</u>		LUPIN LTD	<u>EQ 5MG BASE;12.5MG;160MG</u>	<u>A200797 001</u>	Jun 03, 2015
<u>AB</u>			<u>EQ 5MG BASE;25MG;160MG</u>	<u>A200797 002</u>	Jun 03, 2015
<u>AB</u>			<u>EQ 10MG BASE;12.5MG;160MG</u>	<u>A200797 003</u>	Jun 03, 2015
<u>AB</u>			<u>EQ 10MG BASE;25MG;160MG</u>	<u>A200797 004</u>	Jun 03, 2015
<u>AB</u>			<u>EQ 10MG BASE;25MG;320MG</u>	<u>A200797 005</u>	Jun 03, 2015

EXFORGE HCT

<u>AB</u>	+	NOVARTIS	<u>EQ 5MG BASE;12.5MG;160MG</u>	<u>N022314 001</u>	Apr 30, 2009
<u>AB</u>	+		<u>EQ 5MG BASE;25MG;160MG</u>	<u>N022314 002</u>	Apr 30, 2009
<u>AB</u>	+		<u>EQ 10MG BASE;12.5MG;160MG</u>	<u>N022314 003</u>	Apr 30, 2009
<u>AB</u>	+		<u>EQ 10MG BASE;25MG;160MG</u>	<u>N022314 004</u>	Apr 30, 2009
<u>AB</u>	+		<u>EQ 10MG BASE;25MG;320MG</u>	<u>N022314 005</u>	Apr 30, 2009

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

AMLODIPINE AND OLMESARTAN MEDOXOMIL

<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
<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>	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>	TORRENT	<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	<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

+	ADHERA	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>	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	<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	<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

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; VALSARTAN

TABLET; ORAL

AMLODIPINE BESYLATE AND VALSARTAN

<u>AB</u>	ALEMBIC 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	<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>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 HLTH 414	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203700 001</u>	Feb 25, 2013
<u>AP</u>	DECATUR	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204465 001</u>	Oct 23, 2014
<u>AP</u>	ESSENTIAL ISOTOPES	<u>3.75mCi-260mCi/ML</u>	<u>A205687 001</u>	Dec 17, 2015
<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>	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>	METHODIST	<u>3.75mCi-260mCi/ML</u>	<u>A215083 001</u>	Jul 09, 2021
<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>	NCM USA BRONX LLC	<u>3.75mCi-260mCi/ML</u>	<u>A204515 001</u>	Feb 04, 2015
<u>AP</u>	NUKEMED	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204455 001</u>	Apr 23, 2015
<u>AP</u>	PETNET	<u>30mCi-300mCi (3.75-37.5mCi/ML)</u>	<u>A204510 001</u>	Nov 02, 2015
<u>AP</u>	PRECISION NUCLEAR	<u>3.75mCi-260mCi/ML</u>	<u>A204547 001</u>	Aug 14, 2015
<u>AP</u>	SHERTECH LABS LLC	<u>3.75mCi-260mCi/ML</u>	<u>A204366 001</u>	Sep 19, 2014
<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>		<u>3.75mCi-260mCi/ML</u>	<u>A203543 001</u>	Dec 14, 2012
<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>	UNIV TX SW MEDCTR	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A209507 001</u>	Nov 01, 2019
<u>AP</u>	UNIV WISCONSIN	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A211740 001</u>	Sep 09, 2020
<u>AP</u>	WA UNIV SCH MED	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204506 001</u>	Feb 07, 2014
<u>AP</u>	WISCONSIN	<u>3.75mCi-260mCi/ML</u>	<u>A204356 001</u>	Dec 18, 2014

AMMONIUM CHLORIDE

INJECTABLE; INJECTION

AMMONIUM CHLORIDE IN PLASTIC CONTAINER

!	HOSPIRA	5MEQ/ML	A088366 001	Jun 13, 1984
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PRESCRIPTION DRUG PRODUCT LIST

AMMONIUM LACTATE

CREAM; TOPICAL

AMMONIUM LACTATE

<u>AB</u>	!	PADAGIS ISRAEL	<u>EQ 12% BASE</u>	<u>A075774 001</u>	May 01, 2002
<u>AB</u>		TARO	<u>EQ 12% BASE</u>	<u>A075883 001</u>	Apr 10, 2003

LOTION; TOPICAL

AMMONIUM LACTATE

<u>AB</u>	!	PADAGIS ISRAEL	<u>EQ 12% BASE</u>	<u>A075570 001</u>	Jun 23, 2004
<u>AB</u>		TARO	<u>EQ 12% BASE</u>	<u>A076216 001</u>	May 28, 2004

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>		AUROBINDO	<u>250MG</u>	<u>A065271 001</u>	Nov 09, 2005
<u>AB</u>			<u>500MG</u>	<u>A065271 002</u>	Nov 09, 2005
<u>AB</u>		CHARTWELL	<u>250MG</u>	<u>A062058 001</u>	
<u>AB</u>			<u>500MG</u>	<u>A062058 002</u>	
<u>AB</u>		HIKMA PHARMS	<u>250MG</u>	<u>A065291 001</u>	Feb 05, 2007
<u>AB</u>			<u>500MG</u>	<u>A065291 002</u>	Feb 05, 2007
<u>AB</u>		SANDOZ	<u>250MG</u>	<u>A064076 001</u>	Sep 30, 1994
<u>AB</u>			<u>500MG</u>	<u>A064076 002</u>	Sep 30, 1994
<u>AB</u>		TEVA	<u>250MG</u>	<u>A061926 001</u>	
<u>AB</u>	!		<u>500MG</u>	<u>A061926 003</u>	

AMOXIL

<u>AB</u>		US ANTIBIOTICS	<u>250MG</u>	<u>A062216 001</u>	
<u>AB</u>			<u>500MG</u>	<u>A062216 004</u>	

FOR SUSPENSION; ORAL

AMOXICILLIN

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

AMOXICILLIN PEDIATRIC

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

<u>AB</u>		US ANTIBIOTICS	<u>50MG/ML</u>	<u>A062226 005</u>	
<u>AB</u>			<u>125MG/5ML</u>	<u>A062226 001</u>	
<u>AB</u>			<u>250MG/5ML</u>	<u>A062226 002</u>	

LAROTID

<u>AB</u>		US ANTIBIOTICS	<u>125MG/5ML</u>	<u>A062226 003</u>	
<u>AB</u>			<u>250MG/5ML</u>	<u>A062226 004</u>	

TABLET; ORAL

AMOXICILLIN

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

PRESCRIPTION DRUG PRODUCT LIST

AMOXICILLIN

TABLET, CHEWABLE;ORAL

AMOXICILLIN

TEVA

125MG

A064013 002 Sep 11, 1995

!

250MG

A064013 001 Dec 22, 1992

AMOXICILLIN; CLARITHROMYCIN; LANSOPRAZOLE

CAPSULE, TABLET, CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED)AB ! RISING500MG;500MG;30MGA206006 001 Oct 07, 2016AB SANDOZ INC500MG;500MG;30MGA202588 001 Mar 04, 2014AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUMAB AUROBINDO PHARMA LTD125MG/5ML;EQ 31.25MG BASE/5MLA209371 001 Apr 19, 2019AB 200MG/5ML;EQ 28.5MG BASE/5MLA201090 001 Dec 20, 2011AB 250MG/5ML;EQ 62.5MG BASE/5MLA209371 002 Apr 19, 2019AB 400MG/5ML;EQ 57MG BASE/5MLA201090 002 Dec 20, 2011AB 600MG/5ML;EQ 42.9MG BASE/5MLA201091 001 Dec 20, 2011AB HIKMA PHARMS 200MG/5ML;EQ 28.5MG BASE/5MLA065191 002 Jan 25, 2005AB 400MG/5ML;EQ 57MG BASE/5MLA065191 001 Jan 25, 2005AB 600MG/5ML;EQ 42.9MG BASE/5MLA065373 001 Nov 09, 2007AB MICRO LABS LTD 200MG/5ML;EQ 28.5MG BASE/5MLA205187 001 May 21, 2021

INDIA

AB 400MG/5ML;EQ 57MG BASE/5MLA205187 002 May 21, 2021AB SANDOZ 200MG/5ML;EQ 28.5MG BASE/5MLA065066 001 Jun 05, 2002AB 400MG/5ML;EQ 57MG BASE/5MLA065066 002 Jun 05, 2002AB SANDOZ INC 200MG/5ML;EQ 28.5MG BASE/5MLA065098 001 Dec 16, 2002AB 400MG/5ML;EQ 57MG BASE/5MLA065098 002 Dec 16, 2002AB 600MG/5ML;EQ 42.9MG BASE/5MLA065358 001 Aug 13, 2007AB TEVA 200MG/5ML;EQ 28.5MG BASE/5MLA065089 001 May 25, 2004AB ! 400MG/5ML;EQ 57MG BASE/5MLA065089 002 May 25, 2004AB ! 600MG/5ML;EQ 42.9MG BASE/5MLA065162 001 Mar 12, 2004AB WOCKHARDT BIO AG 250MG/5ML;EQ 62.5MG BASE/5MLA065431 001 Nov 25, 2008AB 600MG/5ML;EQ 42.9MG BASE/5MLA065420 001 Dec 02, 2013AUGMENTIN '125'AB + US ANTIBIOTICS125MG/5ML;EQ 31.25MG BASE/5MLN050575 001 Aug 06, 1984AUGMENTIN '250'AB +! US ANTIBIOTICS250MG/5ML;EQ 62.5MG BASE/5MLN050575 002 Aug 06, 1984AUGMENTIN ES-600AB + US ANTIBIOTICS600MG/5ML;EQ 42.9MG BASE/5MLN050755 001 Jun 22, 2001

TABLET;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUMAB AUROBINDO PHARMA LTD250MG;EQ 125MG BASEA091569 001 Jan 20, 2012AB 500MG;EQ 125MG BASEA091569 002 Jan 20, 2012AB 875MG;EQ 125MG BASEA091568 001 Jan 20, 2012AB HIKMA PHARMS 875MG;EQ 125MG BASEA203824 001 Aug 23, 2016AB MICRO LABS LTD 250MG;EQ 125MG BASEA205707 001 Dec 30, 2016

INDIA

AB 500MG;EQ 125MG BASEA205707 002 Dec 30, 2016AB 875MG;EQ 125MG BASEA204755 003 Dec 30, 2016AB ! SANDOZ 250MG;EQ 125MG BASEA065189 001 Aug 23, 2005AB 500MG;EQ 125MG BASEA065064 001 Mar 15, 2002AB ! 875MG;EQ 125MG BASEA065063 001 Mar 14, 2002AB ! SANDOZ INC 500MG;EQ 125MG BASEA065117 001 Nov 27, 2002AB 875MG;EQ 125MG BASEA065093 001 Nov 21, 2002AB TEVA 500MG;EQ 125MG BASEA065101 001 Oct 30, 2002AB TEVA PHARMS USA 875MG;EQ 125MG BASEA065096 001 Oct 29, 2002AUGMENTIN '875'AB + US ANTIBIOTICS875MG;EQ 125MG BASEN050720 001 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

!

SANDOZ

1GM;EQ 62.5MG BASE

A090227 001 Apr 21, 2010

PRESCRIPTION DRUG PRODUCT LIST

AMOXICILLIN; OMEPRAZOLE MAGNESIUM; RIFABUTIN

CAPSULE, DELAYED RELEASE;ORAL

TALICIA

+! REDHILL 250MG;EQ 10MG BASE;12.5MG N213004 001 Nov 01, 2019

AMPHETAMINE

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

TABLET;ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

<u>AB</u>	RHODES PHARMS	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A213111</u>	<u>003</u>	Jan 13, 2021
<u>AB</u>		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A213111</u>	<u>005</u>	Jan 13, 2021
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>A213111</u>	<u>006</u>	Jan 13, 2021

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL

ADDERALL XR 10AB + TAKEDA PHARMS USA 2.5MG;2.5MG;2.5MG;2.5MG N021303 001 Oct 11, 2001ADDERALL XR 15AB + TAKEDA PHARMS USA 3.75MG;3.75MG;3.75MG;3.75MG N021303 006 May 22, 2002ADDERALL XR 20AB + TAKEDA PHARMS USA 5MG;5MG;5MG;5MG N021303 002 Oct 11, 2001ADDERALL XR 25AB + TAKEDA PHARMS USA 6.25MG;6.25MG;6.25MG;6.25MG N021303 004 May 22, 2002ADDERALL XR 30AB +! TAKEDA PHARMS USA 7.5MG;7.5MG;7.5MG;7.5MG N021303 003 Oct 11, 2001ADDERALL XR 5AB + TAKEDA PHARMS USA 1.25MG;1.25MG;1.25MG;1.25MG N021303 005 May 22, 2002DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATEAB ASCENT PHARMS INC 1.25MG;1.25MG;1.25MG;1.25MG A214959 001 Sep 29, 2021AB 2.5MG;2.5MG;2.5MG;2.5MG A214959 002 Sep 29, 2021AB 3.75MG;3.75MG;3.75MG;3.75MG A214959 003 Sep 29, 2021AB 5MG;5MG;5MG;5MG A214959 004 Sep 29, 2021AB 6.25MG;6.25MG;6.25MG;6.25MG A214959 005 Sep 29, 2021AB 7.5MG;7.5MG;7.5MG;7.5MG A214959 006 Sep 29, 2021AB LANNETT CO INC 1.25MG;1.25MG;1.25MG;1.25MG A214403 001 Nov 26, 2021AB 2.5MG;2.5MG;2.5MG;2.5MG A214403 002 Nov 26, 2021AB 3.75MG;3.75MG;3.75MG;3.75MG A214403 003 Nov 26, 2021AB 5MG;5MG;5MG;5MG A214403 004 Nov 26, 2021AB 6.25MG;6.25MG;6.25MG;6.25MG A214403 005 Nov 26, 2021AB 7.5MG;7.5MG;7.5MG;7.5MG A214403 006 Nov 26, 2021DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATEAB ACTAVIS ELIZABETH 1.25MG;1.25MG;1.25MG;1.25MG A077302 001 Jun 22, 2012AB 2.5MG;2.5MG;2.5MG;2.5MG A077302 002 Jun 22, 2012AB 3.75MG;3.75MG;3.75MG;3.75MG A077302 003 Jun 22, 2012AB 5MG;5MG;5MG;5MG A077302 004 Jun 22, 2012AB 6.25MG;6.25MG;6.25MG;6.25MG A077302 005 Jun 22, 2012AB 7.5MG;7.5MG;7.5MG;7.5MG A077302 006 Jun 22, 2012AB ANI PHARMS 1.25MG;1.25MG;1.25MG;1.25MG A205401 001 Jan 22, 2019AB 2.5MG;2.5MG;2.5MG;2.5MG A205401 002 Jan 22, 2019AB 3.75MG;3.75MG;3.75MG;3.75MG A205401 003 Jan 22, 2019AB 5MG;5MG;5MG;5MG A205401 004 Jan 22, 2019AB 6.25MG;6.25MG;6.25MG;6.25MG A205401 005 Jan 22, 2019AB 7.5MG;7.5MG;7.5MG;7.5MG A205401 006 Jan 22, 2019AB ELITE LABS INC 1.25MG;1.25MG;1.25MG;1.25MG A212037 001 Dec 11, 2019AB 2.5MG;2.5MG;2.5MG;2.5MG A212037 002 Dec 11, 2019AB 3.75MG;3.75MG;3.75MG;3.75MG A212037 003 Dec 11, 2019AB 5MG;5MG;5MG;5MG A212037 004 Dec 11, 2019AB 6.25MG;6.25MG;6.25MG;6.25MG A212037 005 Dec 11, 2019AB 7.5MG;7.5MG;7.5MG;7.5MG A212037 006 Dec 11, 2019AB IMPAX LABS 1.25MG;1.25MG;1.25MG;1.25MG A076852 001 Feb 16, 2016AB 2.5MG;2.5MG;2.5MG;2.5MG A076852 002 Feb 16, 2016AB 3.75MG;3.75MG;3.75MG;3.75MG A076852 003 Feb 16, 2016AB 5MG;5MG;5MG;5MG A076852 004 Feb 16, 2016

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		<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	PAR PHARM INC	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A206159 001</u>	May 31, 2019
AB		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A206159 002</u>	May 31, 2019
AB		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A206159 003</u>	May 31, 2019
AB		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A206159 004</u>	May 31, 2019
AB		<u>6.25MG; 6.25MG; 6.25MG; 6.25MG</u>	<u>A206159 005</u>	May 31, 2019
AB		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A206159 006</u>	May 31, 2019
AB	RHODES PHARMS	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A210651 001</u>	May 17, 2019
AB		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A210651 002</u>	May 17, 2019
AB		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A210651 003</u>	May 17, 2019
AB		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A210651 004</u>	May 17, 2019
AB		<u>6.25MG; 6.25MG; 6.25MG; 6.25MG</u>	<u>A210651 005</u>	May 17, 2019
AB		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A210651 006</u>	May 17, 2019
AB	SPECGX LLC	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A211547 001</u>	Apr 22, 2019
AB		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A211547 002</u>	Apr 22, 2019
AB		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A211547 003</u>	Apr 22, 2019
AB		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A211547 004</u>	Apr 22, 2019
AB		<u>6.25MG; 6.25MG; 6.25MG; 6.25MG</u>	<u>A211547 005</u>	Apr 22, 2019
AB		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A211547 006</u>	Apr 22, 2019
AB	SUN PHARM INDUSTRIES	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A211715 001</u>	May 17, 2019
AB		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A211715 002</u>	May 17, 2019
AB		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A211715 003</u>	May 17, 2019
AB		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A211715 004</u>	May 17, 2019
AB		<u>6.25MG; 6.25MG; 6.25MG; 6.25MG</u>	<u>A211715 005</u>	May 17, 2019
AB		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A211715 006</u>	May 17, 2019
	MYDAYIS			
	+ TAKEDA PHARMS USA	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	ACCORD HLTHCARE	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A214347 001</u>	Nov 22, 2021
AB		<u>1.875MG; 1.875MG; 1.875MG; 1.875MG</u>	<u>A214347 002</u>	Nov 22, 2021
AB		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A214347 003</u>	Nov 22, 2021
AB		<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>A214347 004</u>	Nov 22, 2021
AB		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A214347 005</u>	Nov 22, 2021
AB		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A214347 006</u>	Nov 22, 2021
AB		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A214347 007</u>	Nov 22, 2021
AB	ACTAVIS ELIZABETH	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A206340 001</u>	Feb 05, 2016
AB		<u>1.875MG; 1.875MG; 1.875MG; 1.875MG</u>	<u>A206340 002</u>	Feb 05, 2016
AB		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A206340 003</u>	Feb 05, 2016
AB		<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>A206340 004</u>	Feb 05, 2016
AB		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A206340 005</u>	Feb 05, 2016
AB		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A206340 006</u>	Feb 05, 2016
AB		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A206340 007</u>	Feb 05, 2016
AB	ALVOGEN	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A207388 001</u>	Jul 28, 2017
AB		<u>1.875MG; 1.875MG; 1.875MG; 1.875MG</u>	<u>A207388 002</u>	Jul 28, 2017
AB		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A207388 003</u>	Jul 28, 2017
AB		<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>A207388 004</u>	Jul 28, 2017
AB		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A207388 005</u>	Jul 28, 2017
AB		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A207388 006</u>	Jul 28, 2017
AB		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A207388 007</u>	Jul 28, 2017
AB	ASCENT PHARMS INC	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A213709 001</u>	Apr 22, 2021
AB		<u>1.875MG; 1.875MG; 1.875MG; 1.875MG</u>	<u>A213709 002</u>	Apr 22, 2021
AB		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A213709 003</u>	Apr 22, 2021
AB		<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>A213709 004</u>	Apr 22, 2021
AB		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A213709 005</u>	Apr 22, 2021
AB		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A213709 006</u>	Apr 22, 2021
AB		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A213709 007</u>	Apr 22, 2021
AB	AUROLIFE PHARMA LLC	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A202424 001</u>	Nov 27, 2013
AB		<u>1.875MG; 1.875MG; 1.875MG; 1.875MG</u>	<u>A202424 002</u>	Nov 27, 2013
AB		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A202424 003</u>	Nov 27, 2013
AB		<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>A202424 004</u>	Nov 27, 2013
AB		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A202424 005</u>	Nov 27, 2013
AB		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A202424 006</u>	Nov 27, 2013
AB		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A202424 007</u>	Nov 27, 2013

PRESCRIPTION DRUG PRODUCT LIST

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET;ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

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	ELITE LABS INC	1.25MG;1.25MG;1.25MG;1.25MG	A211352	001	Dec 07, 2018
AB		1.875MG;1.875MG;1.875MG;1.875MG	A211352	002	Dec 07, 2018
AB		2.5MG;2.5MG;2.5MG;2.5MG	A211352	003	Dec 07, 2018
AB		3.125MG;3.125MG;3.125MG;3.125MG	A211352	004	Dec 07, 2018
AB		3.75MG;3.75MG;3.75MG;3.75MG	A211352	005	Dec 07, 2018
AB		5MG;5MG;5MG;5MG	A211352	006	Dec 07, 2018
AB		7.5MG;7.5MG;7.5MG;7.5MG	A211352	007	Dec 07, 2018
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	GRANULES	1.25MG;1.25MG;1.25MG;1.25MG	A215771	001	Dec 28, 2021
AB		1.875MG;1.875MG;1.875MG;1.875MG	A215771	002	Dec 28, 2021
AB		2.5MG;2.5MG;2.5MG;2.5MG	A215771	003	Dec 28, 2021
AB		3.125MG;3.125MG;3.125MG;3.125MG	A215771	004	Dec 28, 2021
AB		3.75MG;3.75MG;3.75MG;3.75MG	A215771	005	Dec 28, 2021
AB		5MG;5MG;5MG;5MG	A215771	006	Dec 28, 2021
AB		7.5MG;7.5MG;7.5MG;7.5MG	A215771	007	Dec 28, 2021
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	RHODES PHARMS	1.25MG;1.25MG;1.25MG;1.25MG	A213111	001	Jan 13, 2021
AB		1.875MG;1.875MG;1.875MG;1.875MG	A213111	002	Jan 13, 2021
AB		3.125MG;3.125MG;3.125MG;3.125MG	A213111	004	Jan 13, 2021
AB		7.5MG;7.5MG;7.5MG;7.5MG	A213111	007	Jan 13, 2021
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
AB		1.875MG;1.875MG;1.875MG;1.875MG	A040440	002	Oct 07, 2003
AB		2.5MG;2.5MG;2.5MG;2.5MG	A040440	003	Oct 07, 2003
AB		3.125MG;3.125MG;3.125MG;3.125MG	A040440	004	Oct 07, 2003
AB		3.75MG;3.75MG;3.75MG;3.75MG	A040440	005	Oct 07, 2003
AB		5MG;5MG;5MG;5MG	A040440	006	Oct 07, 2003
AB		7.5MG;7.5MG;7.5MG;7.5MG	A040440	007	Oct 07, 2003
AB	SUN PHARM INDUSTRIES	1.25MG;1.25MG;1.25MG;1.25MG	A040480	001	Sep 09, 2003
AB		1.875MG;1.875MG;1.875MG;1.875MG	A040480	002	Sep 09, 2003
AB		2.5MG;2.5MG;2.5MG;2.5MG	A040480	003	Sep 09, 2003
AB		3.125MG;3.125MG;3.125MG;3.125MG	A040480	004	Sep 09, 2003
AB		3.75MG;3.75MG;3.75MG;3.75MG	A040480	005	Sep 09, 2003
AB		5MG;5MG;5MG;5MG	A040480	006	Sep 09, 2003
AB		7.5MG;7.5MG;7.5MG;7.5MG	A040480	007	Sep 09, 2003
AB	USPHARMA WINDLAS	1.25MG;1.25MG;1.25MG;1.25MG	A210293	001	Apr 03, 2020
AB		1.875MG;1.875MG;1.875MG;1.875MG	A210293	002	Apr 03, 2020
AB		2.5MG;2.5MG;2.5MG;2.5MG	A210293	003	Apr 03, 2020
AB		3.75MG;3.75MG;3.75MG;3.75MG	A210293	004	Apr 03, 2020

PRESCRIPTION DRUG PRODUCT LIST

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB		5MG; 5MG; 5MG; 5MG	A210293 005	Apr 03, 2020
AB		7.5MG; 7.5MG; 7.5MG; 7.5MG	A210293 006	Apr 03, 2020

AMPHETAMINE SULFATE

TABLET; ORAL

AMPHETAMINE SULFATE

AA	ALKEM LABS LTD	5MG	A213720 001	Oct 27, 2020
AA		10MG	A213720 002	Oct 27, 2020
AA	AMNEAL PHARMS	5MG	A211139 001	Sep 26, 2018
AA		10MG	A211139 002	Sep 26, 2018
AA	AUROLIFE PHARMA LLC	5MG	A211639 001	Apr 17, 2019
AA		10MG	A211639 002	Apr 17, 2019
AA	BIONPHARMA INC	5MG	A212919 001	Nov 22, 2019
AA		10MG	A212919 002	Nov 22, 2019
AA	CEROVENE INC	5MG	A212582 001	Feb 04, 2020
AA		10MG	A212582 002	Feb 04, 2020
AA	EPIC PHARMA LLC	5MG	A213980 001	Oct 27, 2020
AA		10MG	A213980 002	Oct 27, 2020
AA	GRANULES	5MG	A212619 001	Aug 05, 2019
AA		10MG	A212619 002	Aug 05, 2019
AA	PRINSTON INC	5MG	A211861 001	Mar 11, 2020
AA		10MG	A211861 002	Mar 11, 2020
AA	RHODES PHARMS	5MG	A213852 001	Sep 07, 2021
AA		10MG	A213852 002	Sep 07, 2021
AA	SENORES PHARMS	5MG	A212901 001	May 22, 2020
AA		10MG	A212901 002	May 22, 2020
AA	SPECGX LLC	5MG	A213583 001	Jan 22, 2021
AA		10MG	A213583 002	Jan 22, 2021
AA	SUN PHARM INDS INC	5MG	A214574 001	Jan 27, 2021
AA		10MG	A214574 002	Jan 27, 2021

EVEKEO

AA	ARBOR PHARMS LLC	5MG	A200166 001	Aug 09, 2012
AA	!	10MG	A200166 002	Aug 09, 2012

TABLET, ORALLY DISINTEGRATING; ORAL

EVEKEO ODT

+	ARBOR PHARMS LLC	5MG	N209905 001	Jan 30, 2019
+		10MG	N209905 002	Jan 30, 2019
+		15MG	N209905 003	Jan 30, 2019
+	!	20MG	N209905 004	Jan 30, 2019

AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE

SUSPENSION, EXTENDED RELEASE; ORAL

DYANAVEL XR

+	TRIS PHARMA INC	2MG/ML; EQ 0.5MG BASE/ML	N208147 001	Oct 19, 2015
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TABLET, EXTENDED RELEASE; ORAL

DYANAVEL XR 10

+	TRIS PHARMA INC	8MG; EQ 2MG BASE	N210526 002	Nov 04, 2021
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DYANAVEL XR 15

+	TRIS PHARMA INC	12MG; EQ 3MG BASE	N210526 003	Nov 04, 2021
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DYANAVEL XR 20

+	TRIS PHARMA INC	16MG; EQ 4MG BASE	N210526 004	Nov 04, 2021
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DYANAVEL XR 5

+	TRIS PHARMA INC	4MG; EQ 1MG BASE	N210526 001	Nov 04, 2021
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AMPHOTERICIN B

INJECTABLE; INJECTION

AMPHOTERICIN B

!	XGEN PHARMS	50MG/VIAL	A063206 001	Apr 29, 1992
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INJECTABLE, LIPID COMPLEX; INJECTION

ABELCET

+	LEADIANT BIOSCI INC	5MG/ML	N050724 001	Nov 20, 1995
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INJECTABLE, LIPOSOMAL; INJECTION

AMBISOME

AB	+	ASTELLAS	50MG/VIAL	N050740 001	Aug 11, 1997
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AMPHOTERICIN B

AB		SPIL	50MG/VIAL	A212514 001	Dec 14, 2021
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PRESCRIPTION DRUG PRODUCT LIST

AMPICILLIN SODIUM

INJECTABLE; INJECTION

AMPICILLIN SODIUM

AP	ACS DOBFAR SPA	<u>EQ 10GM BASE/VIAL</u>	<u>A090889 001</u>	Apr 03, 2013
AP	ANTIBIOTICE	<u>EQ 250MG BASE/VIAL</u>	<u>A090354 001</u>	Dec 28, 2009
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A090354 002</u>	Dec 28, 2009
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A090354 003</u>	Dec 28, 2009
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A090354 004</u>	Dec 28, 2009
AP	AUROBINDO PHARMA	<u>EQ 250MG BASE/VIAL</u>	<u>A065499 002</u>	Aug 17, 2010
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A065499 003</u>	Aug 17, 2010
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A065499 004</u>	Aug 17, 2010
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A065499 005</u>	Aug 17, 2010
AP		<u>EQ 10GM BASE/VIAL</u>	<u>A065493 001</u>	Aug 17, 2010
AP	HQ SPECTL PHARMA	<u>EQ 250MG BASE/VIAL</u>	<u>A062772 006</u>	Apr 15, 1993
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A062772 007</u>	Apr 15, 1993
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A062772 001</u>	Apr 15, 1993
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A062772 003</u>	Apr 15, 1993
AP		<u>EQ 10GM BASE/VIAL</u>	<u>A063142 001</u>	Apr 15, 1993
AP	ISTITUTO BIO ITA SPA	<u>EQ 10GM BASE/VIAL</u>	<u>A201404 001</u>	Dec 20, 2013
AP		<u>EQ 250MG BASE/VIAL</u>	<u>A062719 001</u>	May 12, 1987
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A062719 003</u>	May 12, 1987
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A062719 002</u>	May 12, 1987
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A062797 002</u>	Jul 12, 1993
AP	MYLAN LABS LTD	<u>EQ 250MG BASE/VIAL</u>	<u>A201025 001</u>	Apr 09, 2014
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A201025 002</u>	Apr 09, 2014
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A201025 003</u>	Apr 09, 2014
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A201025 004</u>	Apr 09, 2014
AP		<u>EQ 10GM BASE/VIAL</u>	<u>A202198 001</u>	Apr 07, 2014
AP	SAGENT PHARMS INC	<u>EQ 125MG BASE/VIAL</u>	<u>A090583 001</u>	Nov 27, 2015
AP		<u>EQ 250MG BASE/VIAL</u>	<u>A090583 002</u>	Nov 27, 2015
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A090583 003</u>	Nov 27, 2015
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A090583 004</u>	Nov 27, 2015
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A090583 005</u>	Nov 27, 2015
AP		<u>EQ 10GM BASE/VIAL</u>	<u>A090581 001</u>	Oct 20, 2015
AP	! SANDOZ	<u>EQ 125MG BASE/VIAL</u>	<u>A061395 001</u>	
AP	!	<u>EQ 250MG BASE/VIAL</u>	<u>A061395 002</u>	
AP	!	<u>EQ 500MG BASE/VIAL</u>	<u>A061395 003</u>	
AP	!	<u>EQ 1GM BASE/VIAL</u>	<u>A061395 004</u>	
AP	!	<u>EQ 2GM BASE/VIAL</u>	<u>A061395 005</u>	
AP	!	<u>EQ 10GM BASE/VIAL</u>	<u>A061395 006</u>	

POWDER; INTRAVENOUS

AMPICILLIN SODIUM

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

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

AP	ACS DOBFAR	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065406 001</u>	Dec 22, 2009
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065406 002</u>	Dec 22, 2009
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065403 001</u>	Dec 23, 2009
AP	ANTIBIOTICE	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A201406 001</u>	Dec 07, 2015
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A201406 002</u>	Dec 07, 2015
AP	ASTRAL	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090579 001</u>	Jan 08, 2016
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090579 002</u>	Jan 08, 2016
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A090578 001</u>	Jan 11, 2016
AP	AUROBINDO PHARMA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090340 001</u>	Sep 20, 2010
AP		<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090349 001</u>	Sep 20, 2010
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090340 002</u>	Sep 20, 2010
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090349 002</u>	Sep 20, 2010
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A090339 001</u>	Sep 20, 2010
AP	HIKMA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065074 001</u>	Mar 19, 2002
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065074 002</u>	Mar 19, 2002
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065076 001</u>	Mar 19, 2002
AP	HQ SPECTL PHARMA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065176 001</u>	Nov 30, 2005
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065176 002</u>	Nov 30, 2005
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065188 001</u>	Nov 25, 2005
AP	ISTITUTO BIO ITA SPA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065222 001</u>	Nov 29, 2005
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065222 002</u>	Nov 29, 2005
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065314 001</u>	Nov 27, 2006
AP	MYLAN LABS LTD	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A201024 001</u>	Apr 07, 2014

PRESCRIPTION DRUG PRODUCT LIST

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

<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>UNASYN</u>				
<u>AP</u>	! PFIZER	<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
<u>AP</u>	+	<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>N050608 005</u>	Dec 10, 1993

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMPICILLIN TRIHYDRATE

! SANDOZ

EQ 500MG BASE

A064082 002 Aug 29, 1995

ANAGRELIDE HYDROCHLORIDE

CAPSULE; ORAL

AGRYLIN

<u>AB</u>	TAKEDA PHARMS USA	<u>EQ 0.5MG BASE</u>	<u>N020333 001</u>	Mar 14, 1997
<u>ANAGRELIDE HYDROCHLORIDE</u>				
<u>AB</u>	IMPAX LABS	<u>EQ 0.5MG BASE</u>	<u>A076910 001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A076910 002</u>	Apr 18, 2005
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 0.5MG BASE</u>	<u>A076468 001</u>	Apr 18, 2005
<u>AB</u>	!	<u>EQ 1MG BASE</u>	<u>A076468 002</u>	Apr 18, 2005
<u>AB</u>	TORRENT	<u>EQ 0.5MG BASE</u>	<u>A209151 001</u>	Jun 30, 2017
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A209151 002</u>	Jun 30, 2017

ANASTROZOLE

TABLET; ORAL

ANASTROZOLE

<u>AB</u>	ACCORD HLTHCARE	<u>1MG</u>	<u>A090568 001</u>	Jun 28, 2010
<u>AB</u>	BEIJING YILING	<u>1MG</u>	<u>A206037 001</u>	Nov 09, 2018
<u>AB</u>	CIPLA	<u>1MG</u>	<u>A091164 001</u>	Jun 28, 2010
<u>AB</u>	EUGIA PHARMA	<u>1MG</u>	<u>A212434 001</u>	Jul 24, 2020
<u>AB</u>	KENTON	<u>1MG</u>	<u>A078944 001</u>	Jun 28, 2010
<u>AB</u>	NATCO PHARMA LTD	<u>1MG</u>	<u>A079220 001</u>	Jun 28, 2010
<u>AB</u>	TEVA PHARMS	<u>1MG</u>	<u>A078058 001</u>	Jun 28, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>1MG</u>	<u>A078921 001</u>	Jun 28, 2010
<u>ARIMIDEX</u>				
<u>AB</u>	+	ANI PHARMS	<u>1MG</u>	<u>N020541 001</u> Dec 27, 1995

ANGIOTENSIN II ACETATE

SOLUTION; INTRAVENOUS

GIAPREZA

+! LA JOLLA PHARMA

EQ 0.5MG BASE/ML (EQ 0.5MG BASE/ML)

N209360 003 Dec 23, 2021

+!

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

N209360 001 Dec 21, 2017

ANIDULAFUNGIN

POWDER; INTRAVENOUS

ERAXIS

+! VICURON HOLDINGS

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

APIXABAN

<u>AB</u>	ACCORD HLTHCARE	<u>2.5MG</u>	<u>A210180 001</u>	Jul 28, 2020
<u>AB</u>		<u>5MG</u>	<u>A210180 002</u>	Jul 28, 2020
<u>AB</u>	INDOCO	<u>2.5MG</u>	<u>A209898 001</u>	Sep 11, 2020
<u>AB</u>		<u>5MG</u>	<u>A209898 002</u>	Sep 11, 2020
<u>ELIQUIS</u>				
<u>AB</u>	+	BRISTOL MYERS	<u>2.5MG</u>	<u>N202155 001</u> Dec 28, 2012

PRESCRIPTION DRUG PRODUCT LIST

APIXABAN

TABLET; ORAL

ELIQUIS

SQUIBB

AB	+		5MG	N202155	002	Dec 28, 2012
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APOMORPHINE HYDROCHLORIDE

FILM; SUBLINGUAL

KYNMOBI

	+	SUNOVION PHARMS INC	10MG	N210875	001	May 21, 2020
	+		15MG	N210875	002	May 21, 2020
	+		20MG	N210875	003	May 21, 2020
	+		25MG	N210875	004	May 21, 2020
	+		30MG	N210875	005	May 21, 2020

INJECTABLE; SUBCUTANEOUS

APOKYN

	+	MDD US	30MG/3ML (10MG/ML)	N021264	002	Apr 20, 2004
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APRACLONIDINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

APRACLONIDINE HYDROCHLORIDE

AT		AKORN	EQ 0.5% BASE	A077764	001	Mar 12, 2009
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IOPIDINE

AT	+	NOVARTIS	EQ 0.5% BASE	N020258	001	Jul 30, 1993
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	+		EQ 1% BASE	N019779	001	Dec 31, 1987
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APREMILAST

TABLET; ORAL

APREMILAST

AB		ALKEM LABS LTD	10MG	A211761	001	Sep 21, 2021
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AB			20MG	A211761	002	Sep 21, 2021
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AB			30MG	A211761	003	Sep 21, 2021
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AB		AMNEAL	10MG	A211782	001	Jun 30, 2021
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AB			20MG	A211782	002	Jun 30, 2021
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AB			30MG	A211782	003	Jun 30, 2021
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AB		ZYDUS PHARMS	10MG	A211859	001	Sep 21, 2021
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AB			20MG	A211859	002	Sep 21, 2021
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AB			30MG	A211859	003	Sep 21, 2021
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OTEZLA

AB	+	AMGEN INC	10MG	N205437	001	Mar 21, 2014
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AB	+		20MG	N205437	002	Mar 21, 2014
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AB	+		30MG	N205437	003	Mar 21, 2014
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APREPITANT

CAPSULE; ORAL

APREPITANT

AB		GLENMARK PHARMS SA	40MG	A207777	001	Oct 12, 2017
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AB			80MG	A207777	002	Oct 12, 2017
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AB			125MG	A207777	003	Oct 12, 2017
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AB		SANDOZ	40MG	A090999	001	Sep 24, 2012
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AB			80MG	A090999	002	Sep 24, 2012
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AB			125MG	A090999	003	Sep 24, 2012
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AB		TORRENT	40MG	A211835	001	Oct 21, 2020
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AB			80MG	A211835	002	Oct 21, 2020
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AB			125MG	A211835	003	Oct 21, 2020
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EMEND

AB	+	MERCK	80MG	N021549	001	Mar 26, 2003
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AB	+		125MG	N021549	002	Mar 26, 2003
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EMULSION; INTRAVENOUS

CINVANTI

	+	HERON THERAPS INC	130MG/18ML (7.2MG/ML)	N209296	001	Nov 09, 2017
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FOR SUSPENSION; ORAL

EMEND

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

SOLUTION; INHALATION

ARFORMOTEROL TARTRATE

AN		AXAR PHARMS INC	EQ 0.015MG BASE/2ML	A213762	001	Jun 22, 2021
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AN		CIPLA	EQ 0.015MG BASE/2ML	A207306	001	Jun 22, 2021
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AN		GLENMARK PHARMS LTD	EQ 0.015MG BASE/2ML	A213132	001	Jun 22, 2021
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AN		TEVA PHARMS USA	EQ 0.015MG BASE/2ML	A200293	001	Nov 09, 2021
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BROVANA

AN	+	SUNOVION	EQ 0.015MG BASE/2ML	N021912	001	Oct 06, 2006
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PRESCRIPTION DRUG PRODUCT LIST

ARGATROBAN

INJECTABLE; INJECTION

ARGATROBAN

<u>AP</u>	AMNEAL PHARMS CO	<u>250MG/2.5ML (100MG/ML)</u>	<u>A206698 001</u>	Jan 26, 2018
<u>AP</u>	CAPLIN	<u>50MG/50ML (1MG/ML)</u>	<u>A214235 001</u>	Jan 21, 2021
<u>AP</u>	FRESENIUS KABI USA	<u>250MG/2.5ML (100MG/ML)</u>	<u>N201811 001</u>	Mar 23, 2015
<u>AP</u>	+! HIKMA PHARM CO LTD	<u>50MG/50ML (1MG/ML)</u>	<u>N203049 002</u>	Sep 30, 2016
<u>AP</u>	+!	<u>250MG/2.5ML (100MG/ML)</u>	<u>N203049 001</u>	Jan 05, 2012
<u>AP</u>	HOSPIRA INC	<u>250MG/2.5ML (100MG/ML)</u>	<u>A204120 001</u>	Sep 21, 2016
<u>AP</u>	MYLAN INSTITUTIONAL	<u>250MG/2.5ML (100MG/ML)</u>	<u>A202626 001</u>	Jun 30, 2014
<u>AP</u>	+! NOVARTIS	<u>250MG/2.5ML (100MG/ML)</u>	<u>N020883 001</u>	Jun 30, 2000
<u>AP</u>	PAR STERILE PRODUCTS	<u>250MG/2.5ML (100MG/ML)</u>	<u>A091665 001</u>	Jun 30, 2014

INJECTABLE; INTRAVENOUS

ARGATROBAN IN SODIUM CHLORIDE

<u>AP</u>	GLAND PHARMA LTD	<u>125MG/125ML (1MG/ML)</u>	<u>A205570 001</u>	May 22, 2017
<u>AP</u>	+! SANDOZ	<u>125MG/125ML (1MG/ML)</u>	<u>N022485 001</u>	May 09, 2011

SOLUTION; INTRAVENOUS

ARGATROBAN IN SODIUM CHLORIDE

+!	ACCORD HLTHCARE	50MG/50ML (1MG/ML)	N212035 001	Jun 07, 2021
	EUGIA PHARMA	50MG/50ML (1MG/ML)	N209552 001	Nov 27, 2018
	SPECLTS			

ARGININE HYDROCHLORIDE

INJECTABLE; INJECTION

R-GENE 10

+!	PHARMACIA AND UPJOHN	10GM/100ML	N016931 001	
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ARIPIPIRAZOLE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

ABILIFY MAINTENA KIT

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

SOLUTION; ORAL

ARIPIPIRAZOLE

<u>AA</u>	! AMNEAL PHARMS	<u>1MG/ML</u>	<u>A203906 001</u>	Aug 14, 2015
<u>AA</u>	APOTEX INC	<u>1MG/ML</u>	<u>A204094 001</u>	Sep 30, 2015
<u>AA</u>	AUROBINDO PHARMA LTD	<u>1MG/ML</u>	<u>A210479 001</u>	Jan 29, 2019
<u>AA</u>	LANNETT CO INC	<u>1MG/ML</u>	<u>A204171 001</u>	Aug 14, 2015
<u>AA</u>	VISTAPHARM	<u>1MG/ML</u>	<u>A212870 001</u>	Dec 26, 2019

TABLET; ORAL

ABILIFY

<u>AB</u>	+ OTSUKA	<u>2MG</u>	<u>N021436 006</u>	Nov 15, 2002
<u>AB</u>	+!	<u>5MG</u>	<u>N021436 005</u>	Nov 15, 2002
<u>AB</u>	+!	<u>10MG</u>	<u>N021436 001</u>	Nov 15, 2002
<u>AB</u>	+	<u>15MG</u>	<u>N021436 002</u>	Nov 15, 2002
<u>AB</u>	+	<u>20MG</u>	<u>N021436 003</u>	Nov 15, 2002
<u>AB</u>	+	<u>30MG</u>	<u>N021436 004</u>	Nov 15, 2002

ARIPIPIRAZOLE

<u>AB</u>	ACCORD HLTHCARE	<u>2MG</u>	<u>A206251 001</u>	Dec 07, 2016
<u>AB</u>		<u>5MG</u>	<u>A206251 002</u>	Dec 07, 2016
<u>AB</u>		<u>10MG</u>	<u>A206251 003</u>	Dec 07, 2016
<u>AB</u>		<u>15MG</u>	<u>A206251 004</u>	Dec 07, 2016
<u>AB</u>		<u>20MG</u>	<u>A206251 005</u>	Dec 07, 2016
<u>AB</u>		<u>30MG</u>	<u>A206251 006</u>	Dec 07, 2016
<u>AB</u>	AJANTA PHARMA LTD	<u>2MG</u>	<u>A206174 001</u>	Sep 12, 2016
<u>AB</u>		<u>5MG</u>	<u>A206174 002</u>	Sep 12, 2016
<u>AB</u>		<u>10MG</u>	<u>A206174 003</u>	Sep 12, 2016
<u>AB</u>		<u>15MG</u>	<u>A206174 004</u>	Sep 12, 2016
<u>AB</u>		<u>20MG</u>	<u>A206174 005</u>	Sep 12, 2016
<u>AB</u>		<u>30MG</u>	<u>A206174 006</u>	Sep 12, 2016
<u>AB</u>	ALEMBIC PHARMS LTD	<u>2MG</u>	<u>A202101 001</u>	Apr 28, 2015
<u>AB</u>		<u>5MG</u>	<u>A202101 002</u>	Apr 28, 2015
<u>AB</u>		<u>10MG</u>	<u>A202101 003</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A202101 004</u>	Apr 28, 2015
<u>AB</u>		<u>20MG</u>	<u>A202101 005</u>	Apr 28, 2015
<u>AB</u>		<u>30MG</u>	<u>A202101 006</u>	Apr 28, 2015
<u>AB</u>	ALKEM LABS LTD	<u>2MG</u>	<u>A207105 001</u>	Feb 21, 2019
<u>AB</u>		<u>5MG</u>	<u>A207105 002</u>	Feb 21, 2019
<u>AB</u>		<u>10MG</u>	<u>A207105 003</u>	Feb 21, 2019

PRESCRIPTION DRUG PRODUCT LIST

ARIPIPRAZOLE

TABLET; ORAL

ARIPIPRAZOLE

<u>AB</u>		<u>15MG</u>	<u>A207105 004</u>	Feb 21, 2019
<u>AB</u>		<u>20MG</u>	<u>A207105 005</u>	Feb 21, 2019
<u>AB</u>		<u>30MG</u>	<u>A207105 006</u>	Feb 21, 2019
<u>AB</u>	AMNEAL PHARMS	<u>2MG</u>	<u>A204838 001</u>	Jun 17, 2016
<u>AB</u>		<u>5MG</u>	<u>A204838 002</u>	Jun 17, 2016
<u>AB</u>		<u>10MG</u>	<u>A204838 003</u>	Jun 17, 2016
<u>AB</u>		<u>15MG</u>	<u>A204838 004</u>	Jun 17, 2016
<u>AB</u>		<u>20MG</u>	<u>A204838 005</u>	Jun 17, 2016
<u>AB</u>		<u>30MG</u>	<u>A204838 006</u>	Jun 17, 2016
<u>AB</u>	APOTEX INC	<u>2MG</u>	<u>A078583 001</u>	Jul 24, 2015
<u>AB</u>		<u>5MG</u>	<u>A078583 002</u>	Jul 24, 2015
<u>AB</u>		<u>10MG</u>	<u>A078583 003</u>	Jul 24, 2015
<u>AB</u>		<u>15MG</u>	<u>A078583 004</u>	Jul 24, 2015
<u>AB</u>		<u>20MG</u>	<u>A078583 005</u>	Jul 24, 2015
<u>AB</u>		<u>30MG</u>	<u>A078583 006</u>	Jul 24, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>2MG</u>	<u>A203908 001</u>	Oct 08, 2015
<u>AB</u>		<u>5MG</u>	<u>A203908 002</u>	Oct 08, 2015
<u>AB</u>		<u>10MG</u>	<u>A203908 003</u>	Oct 08, 2015
<u>AB</u>		<u>15MG</u>	<u>A203908 004</u>	Oct 08, 2015
<u>AB</u>		<u>20MG</u>	<u>A203908 005</u>	Oct 08, 2015
<u>AB</u>		<u>30MG</u>	<u>A203908 006</u>	Oct 08, 2015
<u>AB</u>	BOSCOGEN	<u>2MG</u>	<u>A091279 001</u>	Jan 09, 2017
<u>AB</u>		<u>5MG</u>	<u>A091279 002</u>	Jan 09, 2017
<u>AB</u>		<u>10MG</u>	<u>A091279 003</u>	Jan 09, 2017
<u>AB</u>		<u>15MG</u>	<u>A091279 004</u>	Jan 09, 2017
<u>AB</u>		<u>20MG</u>	<u>A091279 005</u>	Jan 09, 2017
<u>AB</u>		<u>30MG</u>	<u>A091279 006</u>	Jan 09, 2017
<u>AB</u>	HETERO LABS LTD V	<u>2MG</u>	<u>A205064 001</u>	Apr 28, 2015
<u>AB</u>		<u>5MG</u>	<u>A205064 002</u>	Apr 28, 2015
<u>AB</u>		<u>10MG</u>	<u>A205064 003</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A205064 004</u>	Apr 28, 2015
<u>AB</u>		<u>20MG</u>	<u>A205064 005</u>	Apr 28, 2015
<u>AB</u>		<u>30MG</u>	<u>A205064 006</u>	Apr 28, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>2MG</u>	<u>A204111 001</u>	Oct 07, 2016
<u>AB</u>		<u>5MG</u>	<u>A204111 002</u>	Oct 07, 2016
<u>AB</u>		<u>10MG</u>	<u>A204111 003</u>	Oct 07, 2016
<u>AB</u>		<u>15MG</u>	<u>A204111 004</u>	Oct 07, 2016
<u>AB</u>		<u>20MG</u>	<u>A204111 005</u>	Oct 07, 2016
<u>AB</u>		<u>30MG</u>	<u>A204111 006</u>	Oct 07, 2016
<u>AB</u>	ORBION PHARMS	<u>2MG</u>	<u>A202683 001</u>	May 23, 2017
<u>AB</u>		<u>5MG</u>	<u>A202683 002</u>	May 23, 2017
<u>AB</u>		<u>10MG</u>	<u>A202683 003</u>	May 23, 2017
<u>AB</u>		<u>15MG</u>	<u>A202683 004</u>	May 23, 2017
<u>AB</u>		<u>20MG</u>	<u>A202683 005</u>	May 23, 2017
<u>AB</u>		<u>30MG</u>	<u>A202683 006</u>	May 23, 2017
<u>AB</u>	PRINSTON INC	<u>2MG</u>	<u>A205363 001</u>	Dec 04, 2017
<u>AB</u>		<u>5MG</u>	<u>A205363 002</u>	Dec 04, 2017
<u>AB</u>		<u>10MG</u>	<u>A205363 003</u>	Dec 04, 2017
<u>AB</u>		<u>15MG</u>	<u>A205363 004</u>	Dec 04, 2017
<u>AB</u>		<u>20MG</u>	<u>A205363 005</u>	Dec 04, 2017
<u>AB</u>		<u>30MG</u>	<u>A205363 006</u>	Dec 04, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>2MG</u>	<u>A206383 001</u>	Sep 29, 2016
<u>AB</u>		<u>5MG</u>	<u>A206383 002</u>	Sep 29, 2016
<u>AB</u>		<u>10MG</u>	<u>A206383 003</u>	Sep 29, 2016
<u>AB</u>		<u>15MG</u>	<u>A206383 004</u>	Sep 29, 2016
<u>AB</u>		<u>20MG</u>	<u>A206383 005</u>	Sep 29, 2016
<u>AB</u>		<u>30MG</u>	<u>A206383 006</u>	Sep 29, 2016
<u>AB</u>	TORRENT	<u>2MG</u>	<u>A201519 001</u>	Apr 28, 2015
<u>AB</u>		<u>10MG</u>	<u>A201519 003</u>	Apr 28, 2015
<u>AB</u>		<u>5MG</u>	<u>A201519 002</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A201519 004</u>	Apr 28, 2015
<u>AB</u>		<u>20MG</u>	<u>A201519 005</u>	Apr 28, 2015
<u>AB</u>		<u>30MG</u>	<u>A201519 006</u>	Apr 28, 2015
<u>AB</u>	UNICHEM	<u>2MG</u>	<u>A203025 001</u>	Dec 01, 2021
<u>AB</u>		<u>5MG</u>	<u>A203025 002</u>	Dec 01, 2021
<u>AB</u>		<u>10MG</u>	<u>A203025 003</u>	Dec 01, 2021
<u>AB</u>		<u>15MG</u>	<u>A203025 004</u>	Dec 01, 2021
<u>AB</u>		<u>20MG</u>	<u>A203025 005</u>	Dec 01, 2021
<u>AB</u>		<u>30MG</u>	<u>A203025 006</u>	Dec 01, 2021

PRESCRIPTION DRUG PRODUCT LIST

ARIPIPIRAZOLE

TABLET;ORAL

ABILIFY MYCITE KIT

+	OTSUKA	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

AB	!	ALEMBIC PHARMS LTD	10MG	A202102	001	Apr 28, 2015
AB			15MG	A202102	002	Apr 28, 2015
AB		ORBION PHARMS	10MG	A202547	001	Dec 11, 2017
AB			15MG	A202547	002	Dec 11, 2017
AB		SCIEGEN PHARMS INC	10MG	A207240	001	Apr 18, 2018
AB			15MG	A207240	002	Apr 18, 2018
AB		SQUARE PHARMS	10MG	A090165	001	Aug 28, 2018
AB			15MG	A090165	002	Aug 28, 2018
			20MG	A090165	003	Aug 28, 2018
			30MG	A090165	004	Aug 28, 2018

ARIPIPIRAZOLE 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
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ARMODAFINIL

TABLET;ORAL

ARMODAFINIL

AB		AUROBINDO PHARMA LTD	50MG	A206069	001	Mar 06, 2018
AB			150MG	A206069	002	Mar 06, 2018
AB			200MG	A206069	004	Dec 07, 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			100MG	A200043	004	May 09, 2019
AB			150MG	A200043	002	Jun 01, 2012
AB			200MG	A200043	005	May 09, 2019
AB			250MG	A200043	003	Jun 01, 2012
AB		NATCO PHARMA LTD	50MG	A202768	001	Nov 28, 2016
AB			100MG	A202768	004	Sep 28, 2017
AB			150MG	A202768	002	Nov 28, 2016
AB			200MG	A202768	005	Sep 28, 2017
AB			250MG	A202768	003	Nov 28, 2016
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

ARSENIC TRIOXIDE

INJECTABLE; INJECTION

ARSENIC TRIOXIDE

AP		AMNEAL	1MG/ML	A210739	001	Jan 25, 2021
AP			2MG/ML	A210739	002	Aug 19, 2021
AP		AMRING PHARMS	1MG/ML	A210802	001	Nov 13, 2018
AP		EUGIA PHARMA	2MG/ML	A214011	001	Oct 15, 2021
AP		FRESENIUS KABI USA	1MG/ML	A208231	001	Aug 31, 2018
AP		GLAND PHARMA LTD	2MG/ML	A215059	001	Oct 07, 2021
AP		INGENUS PHARMS LLC	1MG/ML	A209315	001	Nov 15, 2018
AP		NEXUS PHARMS	1MG/ML	A209780	001	Nov 15, 2018
AP		SANDOZ INC	2MG/ML	A215359	001	Dec 02, 2021
AP		STI PHARMA LLC	1MG/ML	A209873	001	May 06, 2019
AP		ZYDUS PHARMS	1MG/ML	A206228	001	Nov 13, 2018

PRESCRIPTION DRUG PRODUCT LIST

ARSENIC TRIOXIDE

INJECTABLE; INJECTION

ARSENIC TRIOXIDE

AP		2MG/ML	A206228 002	Aug 30, 2019
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TRISENOX

AP	+ !	CEPHALON	2MG/ML	N021248 002	Oct 13, 2017
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ARTEMETHER; LUMEFANTRINE

TABLET; ORAL

COARTEM

+ !	NOVARTIS	20MG;120MG	N022268 001	Apr 07, 2009
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ARTESUNATE

POWDER; INTRAVENOUS

ARTESUNATE

+ !	AMIVAS	110MG/VIAL	N213036 001	May 26, 2020
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ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

ORABLOC

+	PIERREL	4%;EQ 0.009MG BASE/1.8ML (EQ 0.005MG BASE/ML)	N022466 001	Feb 26, 2010
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+ !		4%;EQ 0.018MG BASE/1.8ML (EQ 0.01MG BASE/ML)	N022466 002	Feb 26, 2010
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SEPTOCAINE

+ !	DEPROCO	4%; EQ 0.0085MG BASE/1.7ML (4%; EQ 0.005MG BASE/ML)	N020971 002	Mar 30, 2006
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+ !		4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML)	N020971 001	Apr 03, 2000
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ASCIMINIB HYDROCHLORIDE

TABLET; ORAL

SCEMBLIX

+	NOVARTIS	EQ 20MG BASE	N215358 001	Oct 29, 2021
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+ !		EQ 40MG BASE	N215358 002	Oct 29, 2021
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ASCORBIC ACID

SOLUTION; INTRAVENOUS

ASCOR

+ !	MCGUFF	25,000MG/50ML (500MG/ML)	N209112 001	Oct 02, 2017
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ASCORBIC ACID; BIOTIN; CHOLECALCIFEROL; CYANOCOBALAMIN; DEXPANTHENOL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE; RIBOFLAVIN; THIAMINE; TOCOPHEROL ACETATE; VITAMIN A; VITAMIN K

INJECTABLE; INTRAVENOUS

INFUVITE PEDIATRIC

+ !	SANDOZ CANADA 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
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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
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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
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PRESCRIPTION DRUG PRODUCT LIST

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

FOR SOLUTION; ORAL

MOVIPREP

AA	+ !	SALIX PHARMS	4.7GM;100GM;1.015GM;5.9GM;2.691GM;7.5GM	N021881 001	Aug 02, 2006
AA		NOVEL LABS INC	4.7GM;100GM;1.015GM;5.9GM;2.691GM;7.5GM	A090145 001	Jan 25, 2012
		PLENVU			
	+ !	SALIX	7.54GM;140GM;2.2GM;48.11GM;5.2GM;9GM	N209381 001	May 04, 2018

ASENAPINE

SYSTEM; TRANSDERMAL

SECUADO

	+	HISAMITSU	3.8MG/24HR	N212268 001	Oct 11, 2019
	+		5.7MG/24HR	N212268 002	Oct 11, 2019
	+ !		7.6MG/24HR	N212268 003	Oct 11, 2019

ASENAPINE MALEATE

TABLET; SUBLINGUAL

ASENAPINE MALEATE

AB		ALEMBIC PHARMS LTD	EQ 2.5MG BASE	A206098 003	Jul 19, 2021
AB			EQ 5MG BASE	A206098 001	Dec 10, 2020
AB			EQ 10MG BASE	A206098 002	Dec 10, 2020
AB		BRECKENRIDGE	EQ 2.5MG BASE	A205960 001	Dec 10, 2020
AB			EQ 5MG BASE	A205960 003	Mar 08, 2021
AB			EQ 10MG BASE	A205960 002	Dec 10, 2020
AB		SIGMAPHARM LABS LLC	EQ 5MG BASE	A206107 001	Dec 10, 2020
AB			EQ 10MG BASE	A206107 002	Dec 10, 2020

SAPHRIS

AB	+	ALLERGAN	EQ 2.5MG BASE	N022117 003	Mar 12, 2015
AB	+		EQ 5MG BASE	N022117 001	Aug 13, 2009
AB	+ !		EQ 10MG BASE	N022117 002	Aug 13, 2009

ASPIRIN

CAPSULE, EXTENDED RELEASE; ORAL

DURLAZA

	+ !	ESPERO	162.5MG	N200671 001	Sep 04, 2015
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ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

FIORINAL

AA	+ !	ALLERGAN	325MG;50MG;40MG	N017534 005	Apr 16, 1986
AA		LANNETT	325MG;50MG;40MG	A086996 002	Oct 11, 1985
		BUTALBITAL, ASPIRIN AND CAFFEINE			
		LGM PHARMA	500MG;50MG;40MG	A205230 001	Oct 18, 2021
		TABLET; ORAL			
		BUTALBITAL, ASPIRIN AND CAFFEINE			
	!	STRIDES PHARMA	325MG;50MG;40MG	A204195 001	Sep 22, 2016

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

AB		LGM PHARMA	325MG;50MG;40MG;30MG	A075231 001	Nov 30, 2001
AB		MAYNE PHARMA INC	325MG;50MG;40MG;30MG	A203335 001	Oct 30, 2015
AB		STEVENS J	325MG;50MG;40MG;30MG	A074951 001	Aug 31, 1998
AB	+ !	ALLERGAN	325MG;50MG;40MG;30MG	N019429 003	Oct 26, 1990

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL

ORPHENGESIC FORTE

	!	GALT PHARMS	770MG;60MG;50MG	A075141 002	May 29, 1998
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ASPIRIN; CARISOPRODOL; CODEINE PHOSPHATE

TABLET; ORAL

CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE

	!	INGENUS PHARMS NJ	325MG;200MG;16MG	A040860 001	Jan 07, 2010
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ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE; ORAL

AGGRENOX

AB	+ !	BOEHRINGER INGELHEIM	25MG;200MG	N020884 001	Nov 22, 1999
AB		AMNEAL PHARMS	25MG;200MG	A206392 001	Mar 08, 2016
AB		BARR	25MG;200MG	A078804 001	Aug 14, 2009

PRESCRIPTION DRUG PRODUCT LIST

ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE; ORAL

ASPIRIN AND DIPYRIDAMOLE

<u>AB</u>	DR REDDYS	<u>25MG;200MG</u>	<u>A209048 001</u>	Oct 10, 2018
<u>AB</u>	GLENMARK PHARMS SA	<u>25MG;200MG</u>	<u>A210318 001</u>	May 24, 2019
<u>AB</u>	MICRO LABS	<u>25MG;200MG</u>	<u>A209929 001</u>	Aug 11, 2021
<u>AB</u>	PAR PHARM INC	<u>25MG;200MG</u>	<u>A207944 001</u>	Jan 18, 2017
<u>AB</u>	SANDOZ INC	<u>25MG;200MG</u>	<u>A206739 001</u>	Jan 18, 2017
<u>AB</u>	ZYDUS PHARMS	<u>25MG;200MG</u>	<u>A206753 001</u>	Aug 29, 2017

ASPIRIN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE AND ASPIRIN

<u>AA</u>	EPIC PHARMA LLC	<u>325MG;4.8355MG</u>	<u>A040910 001</u>	Jul 16, 2020
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PERCODAN

<u>AA</u>	+! ENDO PHARMS	<u>325MG;4.8355MG</u>	<u>N007337 007</u>	Aug 05, 2005
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ATAZANAVIR SULFATE

CAPSULE; ORAL

ATAZANAVIR SULFATE

<u>AB</u>	AMNEAL	<u>EQ 150MG BASE</u>	<u>A209717 002</u>	Jun 01, 2020
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A209717 003</u>	Jun 01, 2020
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A209717 004</u>	Jun 01, 2020
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 100MG BASE</u>	<u>A204806 001</u>	Jun 25, 2018
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A204806 002</u>	Jun 25, 2018
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A204806 003</u>	Jun 25, 2018
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A204806 004</u>	Jun 25, 2018
<u>AB</u>	CIPLA	<u>EQ 100MG BASE</u>	<u>A200626 001</u>	Aug 09, 2018
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A200626 002</u>	Aug 09, 2018
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A200626 003</u>	Aug 09, 2018
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A200626 004</u>	Aug 09, 2018
<u>AB</u>	LAURUS	<u>EQ 150MG BASE</u>	<u>A212579 001</u>	Apr 30, 2021
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A212579 002</u>	Apr 30, 2021
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A212579 003</u>	Apr 30, 2021
<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	<u>EQ 50MG BASE/PACKET</u>	<u>N206352 001</u>	Jun 02, 2014
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ATAZANAVIR SULFATE; COBICISTAT

TABLET; ORAL

EVOTAZ

+!	BRISTOL-MYERS SQUIBB	<u>EQ 300MG BASE;150MG</u>	<u>N206353 001</u>	Jan 29, 2015
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ATENOLOL

TABLET; ORAL

ATENOLOL

<u>AB</u>	ALVOGEN	<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>	HLTHCARE	<u>25MG</u>	<u>A073026 002</u>	May 01, 1992
<u>AB</u>		<u>50MG</u>	<u>A073026 003</u>	Sep 17, 1991
<u>AB</u>		<u>100MG</u>	<u>A073026 001</u>	Sep 17, 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>	TEVA	<u>25MG</u>	<u>A074056 003</u>	Jul 19, 2004
<u>AB</u>		<u>50MG</u>	<u>A074056 001</u>	Jan 18, 1995

PRESCRIPTION DRUG PRODUCT LIST

ATENOLOL

TABLET; ORAL

ATENOLOL

<u>AB</u>		<u>100MG</u>	<u>A074056</u>	<u>002</u>	Jan 18, 1995
<u>AB</u>	UNICHEM	<u>25MG</u>	<u>A213136</u>	<u>001</u>	Nov 21, 2019
<u>AB</u>		<u>50MG</u>	<u>A213136</u>	<u>002</u>	Nov 21, 2019
<u>AB</u>		<u>100MG</u>	<u>A213136</u>	<u>003</u>	Nov 21, 2019
<u>AB</u>	UNIQUE PHARM LABS	<u>25MG</u>	<u>A077443</u>	<u>001</u>	Sep 13, 2006
<u>AB</u>		<u>50MG</u>	<u>A077443</u>	<u>002</u>	Sep 13, 2006
<u>AB</u>		<u>100MG</u>	<u>A077443</u>	<u>003</u>	Sep 13, 2006
<u>AB</u>	ZYDUS PHARMS USA	<u>25MG</u>	<u>A076900</u>	<u>001</u>	Jan 28, 2005
<u>AB</u>		<u>50MG</u>	<u>A076900</u>	<u>002</u>	Jan 28, 2005
<u>AB</u>		<u>100MG</u>	<u>A076900</u>	<u>003</u>	Jan 28, 2005
<u>TENORMIN</u>					
<u>AB</u>	+ ALMATICA	<u>25MG</u>	<u>N018240</u>	<u>004</u>	Apr 09, 1990
<u>AB</u>	+	<u>50MG</u>	<u>N018240</u>	<u>001</u>	
<u>AB</u>	+!	<u>100MG</u>	<u>N018240</u>	<u>002</u>	

ATENOLOL; CHLORTHALIDONE

TABLET; ORAL

ATENOLOL AND CHLORTHALIDONE

<u>AB</u>	ALVOGEN	<u>50MG;25MG</u>	<u>A072302</u>	<u>002</u>	May 31, 1990
<u>AB</u>		<u>100MG;25MG</u>	<u>A072302</u>	<u>001</u>	May 31, 1990
<u>AB</u>	NOVITIUM PHARMA	<u>50MG;25MG</u>	<u>A215560</u>	<u>001</u>	Oct 25, 2021
<u>AB</u>		<u>100MG;25MG</u>	<u>A215560</u>	<u>002</u>	Oct 25, 2021
<u>AB</u>	UNICHEM	<u>50MG;25MG</u>	<u>A213302</u>	<u>001</u>	Nov 25, 2020
<u>AB</u>		<u>100MG;25MG</u>	<u>A213302</u>	<u>002</u>	Nov 25, 2020
<u>AB</u>	WATSON LABS	<u>50MG;25MG</u>	<u>A073665</u>	<u>001</u>	Jul 02, 1992
<u>AB</u>		<u>100MG;25MG</u>	<u>A073665</u>	<u>002</u>	Jul 02, 1992
<u>AB</u>	ZYDUS PHARMS	<u>50MG;25MG</u>	<u>A210028</u>	<u>001</u>	Mar 08, 2019
<u>AB</u>		<u>100MG;25MG</u>	<u>A210028</u>	<u>002</u>	Mar 08, 2019
<u>TENORETIC 100</u>					
<u>AB</u>	+! ALMATICA	<u>100MG;25MG</u>	<u>N018760</u>	<u>001</u>	Jun 08, 1984
<u>TENORETIC 50</u>					
<u>AB</u>	+ ALMATICA	<u>50MG;25MG</u>	<u>N018760</u>	<u>002</u>	Jun 08, 1984

ATOGEPAANT

TABLET; ORAL

QULIPTA

+	ABEVIE INC	10MG	N215206	001	Sep 28, 2021
+		30MG	N215206	002	Sep 28, 2021
+		60MG	N215206	003	Sep 28, 2021

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

ATOMOXETINE HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>10MG</u>	<u>A078983</u>	<u>001</u>	May 30, 2017
<u>AB</u>		<u>18MG</u>	<u>A078983</u>	<u>002</u>	May 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A078983</u>	<u>003</u>	May 30, 2017
<u>AB</u>		<u>40MG</u>	<u>A078983</u>	<u>004</u>	May 30, 2017
<u>AB</u>		<u>60MG</u>	<u>A078983</u>	<u>005</u>	May 30, 2017
<u>AB</u>		<u>80MG</u>	<u>A078983</u>	<u>006</u>	May 30, 2017
<u>AB</u>		<u>100MG</u>	<u>A078983</u>	<u>007</u>	May 30, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A079016</u>	<u>001</u>	May 30, 2017
<u>AB</u>		<u>18MG</u>	<u>A079016</u>	<u>002</u>	May 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A079016</u>	<u>003</u>	May 30, 2017
<u>AB</u>		<u>40MG</u>	<u>A079016</u>	<u>004</u>	May 30, 2017
<u>AB</u>		<u>60MG</u>	<u>A079016</u>	<u>005</u>	May 30, 2017
<u>AB</u>		<u>80MG</u>	<u>A079016</u>	<u>006</u>	May 30, 2017
<u>AB</u>		<u>100MG</u>	<u>A079016</u>	<u>007</u>	May 30, 2017
<u>AB</u>	DR REDDYS LABS LTD	<u>10MG</u>	<u>A090609</u>	<u>001</u>	Feb 23, 2018
<u>AB</u>		<u>18MG</u>	<u>A090609</u>	<u>002</u>	Feb 23, 2018
<u>AB</u>		<u>25MG</u>	<u>A090609</u>	<u>003</u>	Feb 23, 2018
<u>AB</u>		<u>40MG</u>	<u>A090609</u>	<u>004</u>	Feb 23, 2018
<u>AB</u>		<u>60MG</u>	<u>A090609</u>	<u>005</u>	Feb 23, 2018
<u>AB</u>		<u>80MG</u>	<u>A090609</u>	<u>006</u>	Feb 23, 2018
<u>AB</u>		<u>100MG</u>	<u>A090609</u>	<u>007</u>	Feb 23, 2018
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MG</u>	<u>A079019</u>	<u>001</u>	May 30, 2017
<u>AB</u>		<u>18MG</u>	<u>A079019</u>	<u>002</u>	May 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A079019</u>	<u>003</u>	May 30, 2017
<u>AB</u>		<u>40MG</u>	<u>A079019</u>	<u>004</u>	May 30, 2017
<u>AB</u>		<u>60MG</u>	<u>A079019</u>	<u>005</u>	May 30, 2017
<u>AB</u>		<u>80MG</u>	<u>A079019</u>	<u>006</u>	May 30, 2017

PRESCRIPTION DRUG PRODUCT LIST

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

ATOMOXETINE HYDROCHLORIDE

<u>AB</u>		<u>100MG</u>	<u>A079019 007</u>	May 30, 2017
<u>AB</u>	HETERO LABS LTD V	<u>10MG</u>	<u>A202682 001</u>	Mar 11, 2021
<u>AB</u>		<u>18MG</u>	<u>A202682 002</u>	Mar 11, 2021
<u>AB</u>		<u>25MG</u>	<u>A202682 003</u>	Mar 11, 2021
<u>AB</u>		<u>40MG</u>	<u>A202682 004</u>	Mar 11, 2021
<u>AB</u>		<u>60MG</u>	<u>A202682 005</u>	Mar 11, 2021
<u>AB</u>		<u>80MG</u>	<u>A202682 006</u>	Mar 11, 2021
<u>AB</u>		<u>100MG</u>	<u>A202682 007</u>	Mar 11, 2021
<u>AB</u>	TEVA PHARMS USA	<u>10MG</u>	<u>A079022 001</u>	May 30, 2017
<u>AB</u>		<u>18MG</u>	<u>A079022 002</u>	May 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A079022 003</u>	May 30, 2017
<u>AB</u>		<u>40MG</u>	<u>A079022 004</u>	May 30, 2017
<u>AB</u>		<u>60MG</u>	<u>A079022 005</u>	May 30, 2017
<u>AB</u>		<u>80MG</u>	<u>A079022 006</u>	May 30, 2017
<u>AB</u>		<u>100MG</u>	<u>A079022 007</u>	May 30, 2017
<u>AB</u>	ZYDUS PHARMS USA INC	<u>18MG</u>	<u>A079017 001</u>	Sep 17, 2010
<u>AB</u>		<u>25MG</u>	<u>A079017 002</u>	Sep 17, 2010
<u>AB</u>		<u>40MG</u>	<u>A079017 003</u>	Sep 17, 2010
<u>AB</u>		<u>60MG</u>	<u>A079017 004</u>	Sep 17, 2010
<u>AB</u>		<u>80MG</u>	<u>A079017 005</u>	Sep 17, 2010
<u>AB</u>		<u>100MG</u>	<u>A079017 006</u>	Sep 17, 2010
<u>STRATTERA</u>				
<u>AB</u>	+ LILLY	<u>10MG</u>	<u>N021411 002</u>	Nov 26, 2002
<u>AB</u>	+	<u>18MG</u>	<u>N021411 003</u>	Nov 26, 2002
<u>AB</u>	+	<u>25MG</u>	<u>N021411 004</u>	Nov 26, 2002
<u>AB</u>	+	<u>40MG</u>	<u>N021411 005</u>	Nov 26, 2002
<u>AB</u>	+!	<u>60MG</u>	<u>N021411 006</u>	Nov 26, 2002
<u>AB</u>	+	<u>80MG</u>	<u>N021411 007</u>	Feb 14, 2005
<u>AB</u>	+	<u>100MG</u>	<u>N021411 008</u>	Feb 14, 2005

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 10MG BASE</u>	<u>A207687 001</u>	Mar 30, 2018
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207687 002</u>	Mar 30, 2018
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207687 003</u>	Mar 30, 2018
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A207687 004</u>	Mar 30, 2018
<u>AB</u>	ALKEM LABS LTD	<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>	APOTEX INC	<u>EQ 10MG BASE</u>	<u>A090548 001</u>	May 29, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A090548 002</u>	May 29, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090548 003</u>	May 29, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A090548 004</u>	May 29, 2012
<u>AB</u>	AUGUST	<u>EQ 10MG BASE</u>	<u>A214969 001</u>	Sep 02, 2021
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A214969 002</u>	Sep 02, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A214969 003</u>	Sep 02, 2021
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A214969 004</u>	Sep 02, 2021
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 10MG BASE</u>	<u>A091650 001</u>	Jul 17, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A091650 002</u>	Jul 17, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A091650 003</u>	Jul 17, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A202357 001</u>	Jul 17, 2012
<u>AB</u>	GRAVITI PHARMS	<u>EQ 10MG BASE</u>	<u>A209912 001</u>	Jun 18, 2018
<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>	LUPIN LTD	<u>EQ 10MG BASE</u>	<u>A204991 001</u>	Mar 06, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A204991 002</u>	Mar 06, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A204991 003</u>	Mar 06, 2019
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A204991 004</u>	Mar 06, 2019

PRESCRIPTION DRUG PRODUCT LIST

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

<u>AB</u>	MICRO LABS LTD INDIA	<u>EQ 10MG BASE</u>	<u>A205945 001</u>	Nov 07, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205945 002</u>	Nov 07, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205945 003</u>	Nov 07, 2019
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A205945 004</u>	Nov 07, 2019
<u>AB</u>	MSN	<u>EQ 10MG BASE</u>	<u>A211933 001</u>	Feb 08, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A211933 002</u>	Feb 08, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A211933 003</u>	Feb 08, 2019
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A211933 004</u>	Feb 08, 2019
<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>	UMEDICA LABS PVT LTD	<u>EQ 10MG BASE</u>	<u>A213853 001</u>	Aug 19, 2020
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A213853 002</u>	Aug 19, 2020
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A213853 003</u>	Aug 19, 2020
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A213853 004</u>	Aug 19, 2020
<u>AB</u>	ZYDUS PHARMS	<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

LIPITOR

<u>AB</u>	+ UPJOHN	<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

ATOVAQUONE

SUSPENSION; ORAL

ATOVAQUONE

<u>AB</u>	ABHAI LLC	<u>750MG/5ML</u>	<u>A210510 001</u>	May 31, 2019
<u>AB</u>	ABON PHARMS LLC	<u>750MG/5ML</u>	<u>A214272 001</u>	Oct 25, 2021
<u>AB</u>	AMNEAL PHARMS	<u>750MG/5ML</u>	<u>A202960 001</u>	Mar 18, 2014
<u>AB</u>	APOTEX	<u>750MG/5ML</u>	<u>A209750 001</u>	Oct 11, 2017
<u>AB</u>	BIONPHARMA INC	<u>750MG/5ML</u>	<u>A212918 001</u>	Mar 30, 2021
<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>	PADAGIS US	<u>750MG/5ML</u>	<u>A207833 001</u>	Apr 28, 2017

MEPRON

<u>AB</u>	+! GLAXOSMITHKLINE LLC	<u>750MG/5ML</u>	<u>N020500 001</u>	Feb 08, 1995
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ATOVAQUONE; PROGUANIL HYDROCHLORIDE

TABLET; ORAL

ATOVAQUONE AND PROGUANIL HYDROCHLORIDE

<u>AB</u>	GLENMARK GENERICS	<u>62.5MG;25MG</u>	<u>A091211 002</u>	Apr 06, 2015
<u>AB</u>		<u>250MG;100MG</u>	<u>A091211 001</u>	Jan 12, 2011
<u>AB</u>	MYLAN	<u>62.5MG;25MG</u>	<u>A202362 001</u>	May 27, 2014
<u>AB</u>		<u>250MG;100MG</u>	<u>A202362 002</u>	May 27, 2014

MALARONE

<u>AB</u>	+! GLAXOSMITHKLINE	<u>250MG;100MG</u>	<u>N021078 001</u>	Jul 14, 2000
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MALARONE PEDIATRIC

<u>AB</u>	+	<u>GLAXOSMITHKLINE</u>	<u>62.5MG;25MG</u>	<u>N021078 002</u>	Jul 14, 2000
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PRESCRIPTION DRUG PRODUCT LIST

ATRACURIUM BESYLATE

INJECTABLE; INJECTION

ATRACURIUM BESYLATE

<u>AP</u>	EUGIA PHARMA	<u>10MG/ML</u>	<u>A206011 001</u>	Apr 08, 2015
<u>AP</u>	! HIKMA	<u>10MG/ML</u>	<u>A074901 001</u>	Jul 18, 1997
<u>AP</u>	HOSPIRA INC	<u>10MG/ML</u>	<u>A090761 001</u>	Oct 18, 2012
<u>AP</u>	MEITHEAL	<u>10MG/ML</u>	<u>A091489 001</u>	Feb 17, 2012

ATRACURIUM BESYLATE PRESERVATIVE FREE

<u>AP</u>	EUGIA PHARMA	<u>10MG/ML</u>	<u>A206010 001</u>	Apr 08, 2015
<u>AP</u>	! HIKMA	<u>10MG/ML</u>	<u>A074900 001</u>	Jul 18, 1997
<u>AP</u>	HOSPIRA INC	<u>10MG/ML</u>	<u>A090782 001</u>	Oct 18, 2012
<u>AP</u>	MEITHEAL	<u>10MG/ML</u>	<u>A091488 001</u>	Feb 17, 2012

ATROPINE

SOLUTION; INTRAMUSCULAR

ATROPEN

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

ATROPINE SULFATE

SOLUTION; INTRAVENOUS

ATROPINE SULFATE

<u>AP</u>	ACCORD HLTHCARE	<u>0.25MG/5ML (0.05MG/ML)</u>	<u>A212868 001</u>	Jul 26, 2021		
<u>AP</u>		<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A212868 002</u>	Jul 26, 2021		
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A212868 003</u>	Jul 26, 2021		
<u>AP</u>	+	!	HOSPIRA	<u>0.25MG/5ML (0.05MG/ML)</u>	<u>N021146 002</u>	Jul 09, 2001
<u>AP</u>	+	!		<u>0.5MG/5ML (0.1MG/ML)</u>	<u>N021146 004</u>	Aug 17, 2017
<u>AP</u>	+	!		<u>1MG/10ML (0.1MG/ML)</u>	<u>N021146 003</u>	Jul 09, 2001
<u>AP</u>	INTL MEDICATION SYS	<u>1MG/10ML (0.1MG/ML)</u>	<u>A212461 001</u>	Oct 05, 2020		
+	!	ACCORD HLTHCARE	0.4MG/ML (0.4MG/ML)	N214652 001	Sep 29, 2020	
+	!		1MG/ML (1MG/ML)	N214652 002	Sep 29, 2020	

SOLUTION; INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS, INTRAOSSEOUS, ENDOTRACHEAL

ATROPINE SULFATE

<u>AP</u>	ACCORD HLTHCARE	<u>8MG/20ML (0.4MG/ML)</u>	<u>A213424 001</u>	Mar 19, 2021		
<u>AP</u>	+	!	FRESENIUS KABI USA	<u>8MG/20ML (0.4MG/ML)</u>	<u>N209260 001</u>	Jan 26, 2018
<u>AP</u>	HIKMA	<u>8MG/20ML (0.4MG/ML)</u>	<u>A213561 001</u>	Dec 01, 2021		

SOLUTION/DROPS; OPHTHALMIC

ATROPINE SULFATE

<u>AT</u>	+	!	AKORN	<u>1%</u>	<u>N206289 001</u>	Jul 18, 2014
<u>AT</u>			APOTEX INC	<u>1%</u>	<u>A215624 001</u>	Nov 26, 2021
			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
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ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

SOLUTION; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

!	!	HIKMA	0.025MG/5ML; 2.5MG/5ML	A087708 001	May 03, 1982
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TABLET; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

<u>AA</u>	ANI PHARMS	<u>0.025MG; 2.5MG</u>	<u>A086727 001</u>	
<u>AA</u>	BAYSHORE PHARMS LLC	<u>0.025MG; 2.5MG</u>	<u>A210819 001</u>	Nov 13, 2018
<u>AA</u>	LANNETT	<u>0.025MG; 2.5MG</u>	<u>A085372 001</u>	
<u>AA</u>	LEADING PHARMA LLC	<u>0.025MG; 2.5MG</u>	<u>A213413 001</u>	Feb 20, 2020
<u>AA</u>	MAYNE PHARMA INC	<u>0.025MG; 2.5MG</u>	<u>A210789 001</u>	Jun 03, 2020
<u>AA</u>	MYLAN	<u>0.025MG; 2.5MG</u>	<u>A085762 001</u>	
<u>AA</u>	PHARMA LIFE	<u>0.025MG; 2.5MG</u>	<u>A207128 001</u>	Oct 21, 2020
<u>AA</u>	SPECGX LLC	<u>0.025MG; 2.5MG</u>	<u>A213335 001</u>	Oct 06, 2020
<u>AA</u>	UPSHER SMITH LABS	<u>0.025MG; 2.5MG</u>	<u>A210571 001</u>	Aug 31, 2018
<u>AA</u>	WINDER LABS LLC	<u>0.025MG; 2.5MG</u>	<u>A211362 001</u>	Jan 27, 2021

LOMOTIL

<u>AA</u>	+	!	PFIZER	<u>0.025MG; 2.5MG</u>	<u>N012462 001</u>
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PRESCRIPTION DRUG PRODUCT LIST

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

AVACOPAN

CAPSULE; ORAL

TAVNEOS

+! CHEMOCENTRYX 10MG N214487 001 Oct 07, 2021

AVANAFIL

TABLET; ORAL

STENDRA

+ METUCHEN PHARMS 50MG N202276 001 Apr 27, 2012

+ 100MG N202276 002 Apr 27, 2012

+! 200MG N202276 003 Apr 27, 2012

AVAPRITINIB

TABLET; ORAL

AYVAKIT

+ BLUEPRINT MEDICINES 25MG N212608 004 Jun 16, 2021

+ 50MG N212608 005 Jun 16, 2021

+ 100MG N212608 001 Jan 09, 2020

+ 200MG N212608 002 Jan 09, 2020

+! 300MG N212608 003 Jan 09, 2020

AVATROMBOPAG MALEATE

TABLET; ORAL

DOPTELET

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

AVIBACTAM SODIUM; CEFTAZIDIME

POWDER; INTRAVENOUS

AVYCAZ

+! ALLERGAN 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	100MG/VIAL	A201537 001	Sep 16, 2013
AP	EUROHLTH INTL SARL	100MG/VIAL	A209337 001	Jun 08, 2020
AP	MEITHEAL	100MG/VIAL	A212128 001	Nov 02, 2020
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
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TABLET; ORAL

ONUREG

+ CELGENE CORP 200MG N214120 001 Sep 01, 2020

+! 300MG N214120 002 Sep 01, 2020

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	ALKEM LABS LTD	25MG	A208687 001	Mar 27, 2020
AB		50MG	A208687 002	Mar 27, 2020
AB		75MG	A208687 003	Mar 27, 2020

PRESCRIPTION DRUG PRODUCT LIST

AZATHIOPRINE

TABLET; ORAL

AZATHIOPRINE

<u>AB</u>		<u>100MG</u>	<u>A208687</u>	<u>004</u>	Mar 27, 2020
<u>AB</u>	AMNEAL	<u>50MG</u>	<u>A074069</u>	<u>001</u>	Feb 16, 1996
<u>AB</u>		<u>75MG</u>	<u>A074069</u>	<u>002</u>	Nov 02, 2021
<u>AB</u>		<u>100MG</u>	<u>A074069</u>	<u>003</u>	Nov 02, 2021
<u>AB</u>	RISING	<u>50MG</u>	<u>A075568</u>	<u>001</u>	Dec 13, 1999
<u>AB</u>	ZYDUS PHARMS USA	<u>25MG</u>	<u>A077621</u>	<u>002</u>	Sep 05, 2008
<u>AB</u>		<u>50MG</u>	<u>A077621</u>	<u>001</u>	Mar 15, 2007
<u>AB</u>		<u>75MG</u>	<u>A077621</u>	<u>003</u>	Sep 05, 2008
<u>AB</u>		<u>100MG</u>	<u>A077621</u>	<u>004</u>	Sep 05, 2008

IMURAN

<u>AB</u>	+!	SEBELA IRELAND LTD	<u>50MG</u>	<u>N016324</u>	<u>001</u>
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AZATHIOPRINE SODIUM

INJECTABLE; INJECTION

AZATHIOPRINE SODIUM

!

HIKMA

EQ 100MG BASE/VIAL

A074419 001 Mar 31, 1995

AZELAIC ACID

AEROSOL, FOAM; TOPICAL

FINACEA

<u>AB</u>	+!	LEO PHARMA AS	<u>15%</u>	<u>N207071</u>	<u>001</u>	Jul 29, 2015
		CREAM; TOPICAL				
		AZELEX				
	+!	ALMIRALL	20%	N020428	001	Sep 13, 1995
		GEL; TOPICAL				

AZELAIC ACID

<u>AB</u>		ACTAVIS LABS UT INC	<u>15%</u>	<u>A208011</u>	<u>001</u>	Nov 19, 2018
<u>AB</u>		GLENMARK PHARMS	<u>15%</u>	<u>A204637</u>	<u>001</u>	Nov 19, 2018
<u>AB</u>		TARO	<u>15%</u>	<u>A210549</u>	<u>001</u>	Aug 23, 2019
<u>AB</u>		TOLMAR	<u>15%</u>	<u>A208724</u>	<u>001</u>	Nov 19, 2018

FINACEA

<u>AB</u>	+!	LEO PHARMA AS	<u>15%</u>	<u>N021470</u>	<u>001</u>	Dec 24, 2002
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AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AZELASTINE HYDROCHLORIDE

<u>AT</u>		AKORN	<u>0.05%</u>	<u>A203660</u>	<u>001</u>	Nov 08, 2016
<u>AT</u>		ALEMBIC PHARMS LTD	<u>0.05%</u>	<u>A209620</u>	<u>001</u>	Mar 20, 2019
<u>AT</u>		APOTEX INC	<u>0.05%</u>	<u>A078621</u>	<u>001</u>	Aug 03, 2009
<u>AT</u>		GLAND PHARMA LTD	<u>0.05%</u>	<u>A210092</u>	<u>001</u>	Feb 25, 2020
<u>AT</u>	!	SANDOZ INC	<u>0.05%</u>	<u>A202305</u>	<u>001</u>	May 31, 2012
<u>AT</u>		SOMERSET THERAPS LLC	<u>0.05%</u>	<u>A207411</u>	<u>001</u>	Mar 29, 2019
<u>AT</u>		SUN PHARM	<u>0.05%</u>	<u>A078738</u>	<u>001</u>	Jun 21, 2010

SPRAY, METERED; NASAL

AZELASTINE HYDROCHLORIDE

<u>AB</u>		AKORN	<u>0.137MG/SPRAY</u>	<u>A207610</u>	<u>001</u>	May 17, 2019
<u>AB</u>			<u>0.2055MG/SPRAY</u>	<u>A210032</u>	<u>001</u>	Aug 23, 2019
<u>AB</u>		ALKEM LABS LTD	<u>0.137MG/SPRAY</u>	<u>A208156</u>	<u>001</u>	Aug 18, 2017
<u>AB</u>		AMNEAL	<u>0.137MG/SPRAY</u>	<u>A204660</u>	<u>001</u>	Aug 28, 2017
<u>AB</u>			<u>0.2055MG/SPRAY</u>	<u>A208199</u>	<u>001</u>	Dec 15, 2017
<u>AB</u>	!	APOTEX INC	<u>0.137MG/SPRAY</u>	<u>A077954</u>	<u>001</u>	Apr 30, 2009
<u>AB</u>			<u>0.2055MG/SPRAY</u>	<u>A201846</u>	<u>001</u>	Aug 31, 2012
<u>AB</u>		AUROBINDO PHARMA LTD	<u>0.137MG/SPRAY</u>	<u>A212289</u>	<u>001</u>	May 08, 2020
<u>AB</u>			<u>0.2055MG/SPRAY</u>	<u>A212775</u>	<u>001</u>	Nov 12, 2020
<u>AB</u>		BRECKENRIDGE	<u>0.137MG/SPRAY</u>	<u>A090176</u>	<u>001</u>	Jul 28, 2015
<u>AB</u>		HIKMA	<u>0.137MG/SPRAY</u>	<u>A091444</u>	<u>001</u>	Oct 24, 2014
<u>AB</u>			<u>0.2055MG/SPRAY</u>	<u>A207243</u>	<u>001</u>	Sep 22, 2017
<u>AB</u>		PADAGIS ISRAEL	<u>0.2055MG/SPRAY</u>	<u>A202743</u>	<u>001</u>	May 08, 2014
<u>AB</u>		SUN PHARM	<u>0.137MG/SPRAY</u>	<u>A090423</u>	<u>001</u>	May 23, 2012
<u>AB</u>		UPSHER SMITH LABS	<u>0.137MG/SPRAY</u>	<u>A202609</u>	<u>001</u>	Mar 17, 2017
<u>AB</u>		ZYDUS PHARMS	<u>0.137MG/SPRAY</u>	<u>A091409</u>	<u>001</u>	Aug 14, 2017

AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE

SPRAY, METERED; NASAL

AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE

<u>AB</u>		APOTEX	<u>0.137MG/SPRAY; 0.05MG/SPRAY</u>	<u>A207712</u>	<u>001</u>	Apr 28, 2017
<u>AB</u>		PADAGIS ISRAEL	<u>0.137MG/SPRAY; 0.05MG/SPRAY</u>	<u>A208111</u>	<u>001</u>	Feb 18, 2021
<u>AB</u>	+!	MYLAN SPECIALITY LP	<u>0.137MG/SPRAY; 0.05MG/SPRAY</u>	<u>N202236</u>	<u>001</u>	May 01, 2012

PRESCRIPTION DRUG PRODUCT LIST

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

<u>AB</u>	AMNEAL	<u>EQ 100MG BASE/5ML</u>	<u>A205666</u>	<u>001</u>	Jul 19, 2018
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A205666</u>	<u>002</u>	Jul 19, 2018
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 100MG BASE/5ML</u>	<u>A209201</u>	<u>001</u>	Oct 09, 2018
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A209201</u>	<u>002</u>	Oct 09, 2018
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 100MG BASE/5ML</u>	<u>A207531</u>	<u>001</u>	Apr 09, 2018
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A207531</u>	<u>002</u>	Apr 09, 2018
<u>AB</u>	PLIVA	<u>EQ 100MG BASE/5ML</u>	<u>A065246</u>	<u>002</u>	Jul 05, 2006
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A065246</u>	<u>001</u>	Jul 05, 2006
<u>AB</u>	TARO	<u>EQ 200MG BASE/5ML</u>	<u>A211521</u>	<u>001</u>	Dec 11, 2019
<u>AB</u>	ZYDUS	<u>EQ 100MG BASE/5ML</u>	<u>A211147</u>	<u>001</u>	Jul 31, 2018
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A211147</u>	<u>002</u>	Jul 31, 2018

ZITHROMAX

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

INJECTABLE;INJECTION

AZITHROMYCIN

<u>AP</u>	EUGIA PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A203294</u>	<u>001</u>	Jun 19, 2015
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 500MG BASE/VIAL</u>	<u>A065179</u>	<u>001</u>	Dec 13, 2005
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A065501</u>	<u>001</u>	Nov 09, 2009
<u>AP</u>	HAINAN POLY PHARM	<u>EQ 500MG BASE/VIAL</u>	<u>A203412</u>	<u>001</u>	Oct 09, 2018
<u>AP</u>	HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>A065500</u>	<u>001</u>	Jun 26, 2009
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A065511</u>	<u>001</u>	Jun 26, 2009
<u>AP</u>	SUN PHARM INDS LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A090923</u>	<u>001</u>	Apr 02, 2013

ZITHROMAX

<u>AP</u>	+	!	PFIZER	<u>EQ 500MG BASE/VIAL</u>	<u>N050733</u>	<u>001</u>	Jan 30, 1997
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SOLUTION/DROPS;OPHTHALMIC

AZASITE

+	!	AKORN	1%	N050810	001	Apr 27, 2007
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TABLET;ORAL

AZITHROMYCIN

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 250MG BASE</u>	<u>A211791</u>	<u>001</u>	Jan 28, 2020
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A211792</u>	<u>001</u>	Jan 28, 2020
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A211793</u>	<u>001</u>	Jan 27, 2020
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 250MG BASE</u>	<u>A207370</u>	<u>001</u>	Jul 05, 2018
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A207398</u>	<u>001</u>	Jul 05, 2018
<u>AB</u>	BIONPHARMA INC	<u>EQ 250MG BASE</u>	<u>A210000</u>	<u>001</u>	Feb 26, 2019
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A210001</u>	<u>001</u>	Feb 26, 2019
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A209999</u>	<u>001</u>	Dec 26, 2018
<u>AB</u>	CSPC OUYI	<u>EQ 250MG BASE</u>	<u>A208250</u>	<u>001</u>	Apr 17, 2019
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A208249</u>	<u>001</u>	Oct 25, 2018
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A207566</u>	<u>001</u>	Sep 24, 2018
<u>AB</u>	LUPIN LTD	<u>EQ 250MG BASE</u>	<u>A065398</u>	<u>001</u>	May 15, 2015
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065399</u>	<u>001</u>	May 15, 2015
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A065400</u>	<u>001</u>	May 15, 2015
<u>AB</u>	PLIVA	<u>EQ 250MG BASE</u>	<u>A065225</u>	<u>001</u>	Nov 14, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065223</u>	<u>001</u>	Nov 14, 2005
<u>AB</u>	!	<u>EQ 600MG BASE</u>	<u>A065218</u>	<u>001</u>	Nov 14, 2005
<u>AB</u>	SANDOZ	<u>EQ 250MG BASE</u>	<u>A065211</u>	<u>001</u>	Nov 14, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065212</u>	<u>001</u>	Nov 14, 2005
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A065209</u>	<u>001</u>	Nov 14, 2005
<u>AB</u>	SUNSHINE	<u>EQ 250MG BASE</u>	<u>A209045</u>	<u>001</u>	Dec 07, 2018
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A209044</u>	<u>001</u>	Dec 07, 2018
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A209043</u>	<u>001</u>	Dec 06, 2018
<u>AB</u>	TEVA	<u>EQ 500MG BASE</u>	<u>A065193</u>	<u>001</u>	Nov 14, 2005
<u>AB</u>	WOCKHARDT	<u>EQ 250MG BASE</u>	<u>A065404</u>	<u>001</u>	Feb 11, 2008
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065405</u>	<u>001</u>	Feb 11, 2008

PRESCRIPTION DRUG PRODUCT LIST

AZITHROMYCIN

TABLET; ORAL

AZITHROMYCIN

<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A065302 003</u>	Feb 11, 2008
<u>AB</u>	YUNG SHIN PHARM	<u>EQ 600MG BASE</u>	<u>A211068 001</u>	May 08, 2020

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

AZTREONAM

FOR SOLUTION; INHALATION

CAYSTON

+	!	GILEAD	75MG/VIAL	N050814 001	Feb 22, 2010
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INJECTABLE; INJECTION

AZACTAM

<u>AP</u>	+	!	BRISTOL MYERS SQUIBB	<u>1GM/VIAL</u>	<u>N050580 002</u>	Dec 31, 1986
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<u>AP</u>	+	!		<u>2GM/VIAL</u>	<u>N050580 003</u>	Dec 31, 1986
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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
<u>AP</u>		HOSPIRA INC	<u>1GM/VIAL</u>	<u>A206517 001</u>	Nov 08, 2021
<u>AP</u>			<u>2GM/VIAL</u>	<u>A206517 002</u>	Nov 08, 2021
		FRESENIUS KABI USA	500MG/VIAL	A065439 001	Jun 18, 2010

BACITRACIN

OINTMENT; OPHTHALMIC

BACITRACIN

!		PADAGIS US	500 UNITS/GM	A061212 001	
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BACITRACIN ZINC; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE

!		PADAGIS US	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A062166 002	
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BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE

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

OINTMENT; OPHTHALMIC

LUMI-SPORYN

<u>AT</u>	+	CASPER PHARMA LLC	<u>EQ 400 UNITS/GM;EQ 3.5MG BASE/GM;EQ 10,000 UNITS/GM</u>	<u>N050417 001</u>	
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NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC

<u>AT</u>		AKORN	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A065088 001</u>	Feb 06, 2004
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<u>AT</u>	!	BAUSCH AND LOMB	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A064064 001</u>	Oct 30, 1995
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<u>AT</u>		PADAGIS US	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A060764 002</u>	
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BACITRACIN ZINC; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

BACITRACIN ZINC AND POLYMYXIN B SULFATE

<u>AT</u>		AKORN	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>A064028 001</u>	Jan 30, 1995
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<u>AT</u>	!	BAUSCH AND LOMB	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>A064046 001</u>	Jan 26, 1995
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<u>AT</u>		PADAGIS US	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>A065022 001</u>	Feb 27, 2002
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BACLOFEN

GRANULES; ORAL

LYVISPAH

+		SAOL THERAPS RES LTD	5MG/PACKET	N215422 001	Nov 22, 2021
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+			10MG/PACKET	N215422 002	Nov 22, 2021
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+	!		20MG/PACKET	N215422 003	Nov 22, 2021
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INJECTABLE; INTRATHECAL

BACLOFEN

<u>AP</u>		EMERALD INTL LTD	<u>0.05MG/ML</u>	<u>A091193 001</u>	May 03, 2016
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<u>AP</u>			<u>0.5MG/ML</u>	<u>A091193 002</u>	May 03, 2016
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<u>AP</u>			<u>2MG/ML</u>	<u>A091193 003</u>	May 03, 2016
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<u>AP</u>		MAIA PHARMS INC	<u>0.05MG/ML</u>	<u>A210777 001</u>	Jan 15, 2021
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<u>AP</u>			<u>0.5MG/ML</u>	<u>A210048 001</u>	Sep 11, 2019
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PRESCRIPTION DRUG PRODUCT LIST

BACLOFEN

INJECTABLE; INTRATHECAL

BACLOFEN

<u>AP</u>		<u>1MG/ML</u>	<u>A210315 001</u>	Jul 30, 2019
<u>AP</u>		<u>2MG/ML</u>	<u>A210048 002</u>	Sep 11, 2019
<u>AP</u>	MYLAN LABS LTD	<u>0.5MG/ML</u>	<u>A209592 001</u>	Mar 21, 2018
<u>AP</u>		<u>1MG/ML</u>	<u>A209594 001</u>	Mar 06, 2018
<u>AP</u>		<u>2MG/ML</u>	<u>A209592 002</u>	Mar 21, 2018

GABLOFEN

<u>AP</u>	+	PIRAMAL CRITICAL	<u>0.05MG/ML</u>	<u>N022462 001</u>	Nov 19, 2010
<u>AP</u>	+		<u>0.5MG/ML</u>	<u>N022462 002</u>	Nov 19, 2010
<u>AP</u>	+		<u>1MG/ML</u>	<u>N022462 004</u>	Jun 22, 2012
<u>AP</u>	+		<u>2MG/ML</u>	<u>N022462 003</u>	Nov 19, 2010

LIORESAL

<u>AP</u>	+	SAOL THERAPS RES LTD	<u>0.05MG/ML</u>	<u>N020075 003</u>	Nov 07, 1996
<u>AP</u>	+		<u>0.5MG/ML</u>	<u>N020075 001</u>	Jun 17, 1992
<u>AP</u>	+		<u>2MG/ML</u>	<u>N020075 002</u>	Jun 17, 1992

SOLUTION; ORAL

OZOBAX

	+	METACEL PHARMS LLC	5MG/5ML	N208193 001	Sep 18, 2019
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TABLET; ORAL

BACLOFEN

<u>AB</u>		AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A214099 001</u>	Jul 13, 2021
<u>AB</u>			<u>20MG</u>	<u>A214099 002</u>	Jul 13, 2021
<u>AB</u>		BEXIMCO PHARMS USA	<u>10MG</u>	<u>A214114 001</u>	Jul 16, 2021
<u>AB</u>			<u>20MG</u>	<u>A214114 002</u>	Jul 16, 2021
<u>AB</u>		EYWA PHARMA	<u>5MG</u>	<u>A211555 003</u>	Nov 30, 2021
<u>AB</u>			<u>10MG</u>	<u>A211555 001</u>	Feb 01, 2019
<u>AB</u>			<u>20MG</u>	<u>A211555 002</u>	Feb 01, 2019
<u>AB</u>		IMPAX	<u>5MG</u>	<u>A077971 003</u>	Jul 07, 2021
<u>AB</u>			<u>10MG</u>	<u>A077971 001</u>	Oct 26, 2007
<u>AB</u>			<u>20MG</u>	<u>A077971 002</u>	Oct 26, 2007
<u>AB</u>		INNOGENIX	<u>5MG</u>	<u>A212378 003</u>	Apr 30, 2021
<u>AB</u>			<u>10MG</u>	<u>A212378 001</u>	Oct 09, 2020
<u>AB</u>			<u>20MG</u>	<u>A212378 002</u>	Oct 09, 2020
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>10MG</u>	<u>A072234 001</u>	Jul 21, 1988
<u>AB</u>	!		<u>20MG</u>	<u>A072235 001</u>	Jul 21, 1988
<u>AB</u>		KARTHA	<u>5MG</u>	<u>A214374 001</u>	Mar 05, 2021
<u>AB</u>			<u>10MG</u>	<u>A214374 002</u>	Mar 05, 2021
<u>AB</u>			<u>20MG</u>	<u>A214374 003</u>	Mar 05, 2021
<u>AB</u>		LANNETT CO INC	<u>5MG</u>	<u>A077241 003</u>	Sep 22, 2021
<u>AB</u>			<u>10MG</u>	<u>A077241 002</u>	Jul 06, 2007
<u>AB</u>			<u>20MG</u>	<u>A077241 001</u>	Dec 20, 2005
<u>AB</u>		NORTHSTAR HLTHCARE	<u>10MG</u>	<u>A078401 002</u>	Sep 18, 2009
<u>AB</u>			<u>20MG</u>	<u>A078401 001</u>	Sep 18, 2009
<u>AB</u>		OXFORD PHARMS	<u>10MG</u>	<u>A077088 002</u>	Oct 31, 2007
<u>AB</u>			<u>20MG</u>	<u>A077088 001</u>	Oct 31, 2007
<u>AB</u>		RUBICON	<u>5MG</u>	<u>A209102 001</u>	Nov 28, 2017
<u>AB</u>			<u>10MG</u>	<u>A209102 002</u>	Nov 28, 2017
<u>AB</u>			<u>20MG</u>	<u>A209102 003</u>	Nov 28, 2017
<u>AB</u>		UNICHEM	<u>10MG</u>	<u>A212067 001</u>	Jul 09, 2020
<u>AB</u>			<u>20MG</u>	<u>A212067 002</u>	Jul 09, 2020
<u>AB</u>		UPSHER SMITH LABS	<u>10MG</u>	<u>A074584 001</u>	Aug 19, 1996
<u>AB</u>			<u>20MG</u>	<u>A074584 002</u>	Aug 19, 1996
<u>AB</u>		VINTAGE PHARMS	<u>10MG</u>	<u>A077068 002</u>	Aug 30, 2005
<u>AB</u>			<u>20MG</u>	<u>A077068 001</u>	Aug 30, 2005
<u>AB</u>		ZYDUS	<u>5MG</u>	<u>A211659 003</u>	Apr 17, 2020
<u>AB</u>			<u>10MG</u>	<u>A211659 001</u>	Nov 23, 2018
<u>AB</u>			<u>20MG</u>	<u>A211659 002</u>	Nov 23, 2018

BALOXAVIR MARBOXIL

FOR SUSPENSION; ORAL

XOFLUZA

	+	GENENTECH INC	2MG/ML	N214410 001	Nov 23, 2020
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TABLET; ORAL

XOFLUZA

	+	GENENTECH INC	20MG	N210854 001	Oct 24, 2018
	+		40MG	N210854 002	Oct 24, 2018
	+		80MG	N210854 003	Mar 18, 2021

PRESCRIPTION DRUG PRODUCT LIST

BALSALAZIDE DISODIUM

CAPSULE;ORAL

BALSALAZIDE DISODIUM

AB	APOTEX INC	750MG	A077883 001	Dec 28, 2007
AB	HIKMA	750MG	A077806 001	Dec 28, 2007

COLAZAL

AB	+ !	VALEANT PHARMS INTL	750MG	N020610 001	Jul 18, 2000
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BARICITINIB

TABLET;ORAL

OLUMIANT

+	ELI LILLY AND CO	1MG	N207924 002	Oct 08, 2019
+ !		2MG	N207924 001	May 31, 2018

BARIUM SULFATE

FOR SUSPENSION;ORAL

E-Z-HD

+ !	BRACCO	98% (334GM/BOT)	N208036 001	Jan 11, 2016
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E-Z-PAQUE

+ !	BRACCO	96% (169GM/BOT)	N208036 002	Apr 07, 2017
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VARIBAR THIN LIQUID

+ !	BRACCO	81% (120GM/BOT)	N208036 004	Apr 30, 2019
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PASTE;ORAL

VARIBAR PUDDING

	BRACCO	40%	N208844 001	Oct 14, 2016
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SUSPENSION;ORAL

ENTERO VU 24%

+ !	BRACCO	24% (144GM/600ML)	N208143 008	May 29, 2020
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LIQUID E-Z-PAQUE

+ !	BRACCO	60% (213GM/BOT)	N208143 003	Mar 01, 2017
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READI-CAT 2

+ !	BRACCO	2% (9GM/BOT)	N208143 001	Jan 15, 2016
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READI-CAT 2 SMOOTHIE

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

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

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

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

+ !	BRACCO	40% (100GM/250ML)	N208143 006	Jan 23, 2018
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BAZEDOXIFENE ACETATE; ESTROGENS, CONJUGATED

TABLET;ORAL

DUAVEE

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

AEROSOL, METERED;INHALATION

QVAR REDIHALER

+	NORTON WATERFORD	0.04MG/INH	N207921 001	Aug 03, 2017
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+		0.08MG/INH	N207921 002	Aug 03, 2017
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AEROSOL, METERED;NASAL

QNASL

+	TEVA BRANDED PHARM	0.04MG/ACTUATION	N202813 002	Dec 17, 2014
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+ !		0.08MG/ACTUATION	N202813 001	Mar 23, 2012
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BECLOMETHASONE DIPROPIONATE MONOHYDRATE

SPRAY, METERED;NASAL

BECONASE AQ

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

TABLET;ORAL

SIRTURO

+	JANSSEN THERAP	EQ 20MG BASE	N204384 002	May 27, 2020
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+ !		EQ 100MG BASE	N204384 001	Dec 28, 2012
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BELINOSTAT

POWDER;INTRAVENOUS

BELEODAQ

+ !	ACROTECH	500MG/VIAL	N206256 001	Jul 03, 2014
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PRESCRIPTION DRUG PRODUCT LIST

BELUMOSUDIL MESYLATE

TABLET; ORAL

REZUROCK

+! KADMON PHARMS LLC EQ 200MG BASE N214783 001 Jul 16, 2021

BELZUTIFAN

TABLET; ORAL

WELIREG

+! MERCK SHARP DOHME 40MG N215383 001 Aug 13, 2021

BEMPEDOIC ACID

TABLET; ORAL

NEXLETOL

+! ESPERION THERAPS 180MG N211616 001 Feb 21, 2020
INCBEMPEDOIC ACID; EZETIMIBE

TABLET; ORAL

NEXLIZET

+! ESPERION THERAPS 180MG;10MG N211617 001 Feb 26, 2020
INCBENAZEPRIL 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>	ANNORA PHARMA	<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>	CADILA	<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
<u>AB</u>	CHARTWELL RX	<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>	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
<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>LOTENSIN</u>				
<u>AB</u>	+ VALIDUS PHARMS	<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>	SANDOZ	<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
<u>LOTENSIN HCT</u>				
<u>AB</u>	+ VALIDUS PHARMS	<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
BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE				
	SANDOZ	5MG;6.25MG	A076631 001	Feb 11, 2004

PRESCRIPTION DRUG PRODUCT LIST

BENDAMUSTINE HYDROCHLORIDE

POWDER; IV (INFUSION)

TRENDA

+	!	CEPHALON	25MG/VIAL	N022249	002	May 01, 2009
+	!		100MG/VIAL	N022249	001	Mar 20, 2008

SOLUTION; IV (INFUSION)

BELRAPZO

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

+	!	EAGLE PHARMS	100MG/4ML (25MG/ML)	N208194	001	Dec 07, 2015
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BENOXINATE HYDROCHLORIDE; FLUORESCEIN SODIUM

SOLUTION/DROPS; OPHTHALMIC

ALTAFLUOR BENOX

+	!	ALTAIRE PHARMS INC	0.4%;0.25%	N208582	001	Dec 14, 2017
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FLUORESCEIN SODIUM AND BENOXINATE HYDROCHLORIDE

+	!	BAUSCH LOMB IRELAND	0.4%;0.3%	N211039	001	Mar 09, 2020
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BENZNIDAZOLE

TABLET; ORAL

BENZNIDAZOLE

+		CHEMO RESEARCH SL	12.5MG	N209570	001	Aug 29, 2017
+	!		100MG	N209570	002	Aug 29, 2017

BENZONATATE

CAPSULE; ORAL

BENZONATATE

<u>AA</u>		ALEMBIC LABS	<u>100MG</u>	<u>A040682</u>	<u>001</u>	Jul 30, 2007
<u>AA</u>			<u>200MG</u>	<u>A040682</u>	<u>002</u>	Jul 30, 2007
<u>AA</u>		APOTEX INC	<u>100MG</u>	<u>A091310</u>	<u>001</u>	Jan 16, 2015
<u>AA</u>			<u>200MG</u>	<u>A091310</u>	<u>002</u>	Jan 16, 2015
<u>AA</u>		ASCENT PHARMS INC	<u>100MG</u>	<u>A211518</u>	<u>001</u>	Feb 22, 2019
<u>AA</u>			<u>150MG</u>	<u>A211518</u>	<u>002</u>	Feb 22, 2019
<u>AA</u>			<u>200MG</u>	<u>A211518</u>	<u>003</u>	Feb 22, 2019
<u>AA</u>		BIONPHARMA INC	<u>100MG</u>	<u>A081297</u>	<u>001</u>	Jan 29, 1993
<u>AA</u>			<u>200MG</u>	<u>A081297</u>	<u>002</u>	Oct 30, 2007
<u>AA</u>		CSPC-NBP PHARM	<u>100MG</u>	<u>A202765</u>	<u>002</u>	Aug 25, 2017
<u>AA</u>			<u>200MG</u>	<u>A202765</u>	<u>001</u>	Jul 31, 2015
<u>AA</u>		MIKART	<u>100MG</u>	<u>A040851</u>	<u>001</u>	Nov 09, 2009
<u>AA</u>			<u>150MG</u>	<u>A040851</u>	<u>002</u>	Nov 09, 2009
<u>AA</u>			<u>200MG</u>	<u>A040851</u>	<u>003</u>	Nov 09, 2009
<u>AA</u>		PURACAP PHARM LLC	<u>100MG</u>	<u>A206948</u>	<u>001</u>	Dec 19, 2018
<u>AA</u>			<u>200MG</u>	<u>A206948</u>	<u>002</u>	Dec 19, 2018
<u>AA</u>		QINGDAO BAHEAL PHARM	<u>100MG</u>	<u>A210562</u>	<u>001</u>	Nov 09, 2018
<u>AA</u>			<u>150MG</u>	<u>A210562</u>	<u>002</u>	Nov 09, 2018
<u>AA</u>			<u>200MG</u>	<u>A210562</u>	<u>003</u>	Nov 09, 2018
<u>AA</u>		STRIDES PHARMA	<u>100MG</u>	<u>A091133</u>	<u>001</u>	Jul 30, 2015
<u>AA</u>			<u>200MG</u>	<u>A091133</u>	<u>002</u>	Jul 30, 2015
<u>AA</u>	!	THEPHARMANETWORK LLC	<u>100MG</u>	<u>A040627</u>	<u>001</u>	Mar 30, 2007
<u>AA</u>	!		<u>150MG</u>	<u>A201209</u>	<u>001</u>	Sep 24, 2014
<u>AA</u>	!		<u>200MG</u>	<u>A040749</u>	<u>001</u>	Jul 25, 2007
<u>AA</u>		ZYDUS PHARMS USA	<u>100MG</u>	<u>A040597</u>	<u>001</u>	Jun 08, 2007
<u>AA</u>			<u>200MG</u>	<u>A040597</u>	<u>002</u>	Jun 08, 2007

TESSALON

<u>AA</u>	+	PFIZER	<u>100MG</u>	<u>N011210</u>	<u>001</u>	
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BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

ACANYA

<u>AB</u>	+	!	BAUSCH	<u>2.5%;EQ 1.2% BASE</u>	<u>N050819</u>	<u>001</u>	Oct 23, 2008
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BENZACLIN

<u>AB</u>	+	!	BAUSCH	<u>5%;EQ 1% BASE</u>	<u>N050756</u>	<u>001</u>	Dec 21, 2000
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CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE

<u>AB</u>		ACTAVIS LABS UT INC	<u>2.5%;EQ 1.2% BASE</u>	<u>A205128</u>	<u>001</u>	Jun 19, 2015
<u>AB</u>		ENCUBE	<u>5%;1.2%</u>	<u>A212433</u>	<u>001</u>	Apr 28, 2021
<u>AB</u>		GLENMARK PHARMS	<u>5%;EQ 1% BASE</u>	<u>A209252</u>	<u>001</u>	Mar 14, 2019
<u>AB</u>		MYLAN PHARMS INC	<u>5%;EQ 1% BASE</u>	<u>A065443</u>	<u>001</u>	Aug 11, 2009
<u>AB</u>		PADAGIS ISRAEL	<u>2.5%;EQ 1.2% BASE</u>	<u>A205397</u>	<u>001</u>	Sep 09, 2019
<u>AB</u>			<u>5%;EQ 1% BASE</u>	<u>A202440</u>	<u>001</u>	Sep 21, 2015
<u>AB</u>	!		<u>5%;1.2%</u>	<u>A090979</u>	<u>001</u>	Jun 26, 2012
<u>AB</u>		TARO	<u>2.5%;EQ 1.2% BASE</u>	<u>A206575</u>	<u>001</u>	Aug 19, 2019
<u>AB</u>			<u>5%;EQ 1% BASE</u>	<u>A208776</u>	<u>001</u>	May 25, 2018
<u>AB</u>		TARO PHARMS	<u>5%;1.2%</u>	<u>A206218</u>	<u>001</u>	Dec 15, 2017

PRESCRIPTION DRUG PRODUCT LIST

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE

<u>AB</u>	ZYDUS PHARMS	<u>5%;1.2%</u>	<u>A210794 001</u>	Dec 28, 2018
	<u>DUAC</u>			
<u>AB</u>	+ STIEFEL	<u>5%;1.2%</u>	<u>N050741 001</u>	Aug 26, 2002
	ONEXTON			
	+! BAUSCH	3.75%;EQ 1.2% BASE	N050819 002	Nov 24, 2014

BENZOYL PEROXIDE; ERYTHROMYCIN

GEL; TOPICAL

BENZAMYCIN

<u>AB</u>	+! VALEANT INTL	<u>5%;3%</u>	<u>N050557 001</u>	Oct 26, 1984
	<u>ERYTHROMYCIN AND BENZOYL PEROXIDE</u>			
<u>AB</u>	LYNE	<u>5%;3%</u>	<u>A065385 001</u>	Sep 18, 2015

BENZOYL PEROXIDE; TRETINOIN

CREAM; TOPICAL

TWYNEO

	+! SOL-GEL	3%;0.1%	N214902 001	Jul 26, 2021
	TECHNOLOGIES			

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL

BENZPHETAMINE HYDROCHLORIDE

<u>AA</u>	ANDA REPOSITORY	<u>50MG</u>	<u>A090473 002</u>	Sep 15, 2010
<u>AA</u>	EPIC PHARMA LLC	<u>50MG</u>	<u>A090346 001</u>	Dec 15, 2015
<u>AA</u>	! KVK TECH	<u>50MG</u>	<u>A090968 001</u>	Jul 20, 2010
	ANDA REPOSITORY	25MG	A090473 001	Sep 15, 2010

BENZTROPINE MESYLATE

INJECTABLE; INJECTION

BENZTROPINE MESYLATE

<u>AP</u>	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A090233 001</u>	Jul 28, 2009
<u>AP</u>	HIKMA	<u>1MG/ML</u>	<u>A209442 001</u>	Oct 14, 2021
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A090287 001</u>	Aug 31, 2009
<u>AP</u>	NAVINTA LLC	<u>1MG/ML</u>	<u>A091525 001</u>	Feb 05, 2013
	<u>COGENTIN</u>			
<u>AP</u>	+! AKORN	<u>1MG/ML</u>	<u>N012015 001</u>	

TABLET; ORAL

BENZTROPINE MESYLATE

<u>AA</u>	! ASPEN GLOBAL INC	<u>0.5MG</u>	<u>A204713 001</u>	Apr 14, 2015
<u>AA</u>	!	<u>1MG</u>	<u>A204713 002</u>	Apr 14, 2015
<u>AA</u>	!	<u>2MG</u>	<u>A204713 003</u>	Apr 14, 2015
<u>AA</u>	CHARTWELL RX	<u>1MG</u>	<u>A081265 002</u>	Jan 23, 1992
<u>AA</u>		<u>2MG</u>	<u>A081265 001</u>	Jan 23, 1992
<u>AA</u>	EPIC PHARMA LLC	<u>0.5MG</u>	<u>A072264 001</u>	Feb 27, 1989
<u>AA</u>		<u>1MG</u>	<u>A072265 001</u>	Feb 27, 1989
<u>AA</u>		<u>2MG</u>	<u>A072266 001</u>	Feb 27, 1989
<u>AA</u>	INVAGEN PHARMS	<u>0.5MG</u>	<u>A090294 001</u>	Mar 29, 2010
<u>AA</u>		<u>1MG</u>	<u>A090294 002</u>	Mar 29, 2010
<u>AA</u>		<u>2MG</u>	<u>A090294 003</u>	Mar 29, 2010
<u>AA</u>	LEADING PHARMA LLC	<u>0.5MG</u>	<u>A090168 001</u>	Nov 28, 2012
<u>AA</u>		<u>1MG</u>	<u>A090168 002</u>	Nov 28, 2012
<u>AA</u>		<u>2MG</u>	<u>A090168 003</u>	Nov 28, 2012
<u>AA</u>	PLIVA	<u>0.5MG</u>	<u>A089058 001</u>	Aug 10, 1988
<u>AA</u>		<u>1MG</u>	<u>A089059 001</u>	Aug 10, 1988
<u>AA</u>		<u>2MG</u>	<u>A089060 001</u>	Aug 10, 1988
<u>AA</u>	VINTAGE	<u>0.5MG</u>	<u>A040715 001</u>	Aug 27, 2007
<u>AA</u>		<u>1MG</u>	<u>A040715 002</u>	Aug 27, 2007
<u>AA</u>		<u>2MG</u>	<u>A040715 003</u>	Aug 27, 2007

BEPOTASTINE BESILATE

SOLUTION/DROPS; OPHTHALMIC

BEPOTASTINE BESILATE

<u>AT</u>	APOTEX	<u>1.5%</u>	<u>A206066 001</u>	Mar 05, 2019
<u>AT</u>	MYLAN	<u>1.5%</u>	<u>A206220 001</u>	Mar 18, 2019
	<u>BEPREVE</u>			
<u>AT</u>	+! BAUSCH AND LOMB INC	<u>1.5%</u>	<u>N022288 001</u>	Sep 08, 2009

PRESCRIPTION DRUG PRODUCT LIST

BEROTRALSTAT HYDROCHLORIDE

CAPSULE;ORAL

ORLADEYO

+ BIOCRYST

EQ 110MG BASE

N214094 001 Dec 03, 2020

+!

EQ 150MG BASE

N214094 002 Dec 03, 2020

BESIFLOXACIN HYDROCHLORIDE

SUSPENSION/DROPS;OPHTHALMIC

BESIVANCE

+! BAUSCH AND LOMB

EQ 0.6% BASE

N022308 001 May 28, 2009

BETAINE

FOR SOLUTION;ORAL

BETAINEAB NOVITIUM PHARMA1GM/SCOOPFULA214864 001 Nov 23, 2021CYSTADANEAB +! RECORDATI RARE1GM/SCOOPFULN020576 001 Oct 25, 1996BETAMETHASONE ACETATE; BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE;INJECTION

BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATEAB AM REGENT3MG/ML;EQ 3MG BASE/MLA090747 001 Jul 31, 2009CELESTONE SOLUSPANAB +! ORGANON3MG/ML;EQ 3MG BASE/MLN014602 001BETAMETHASONE DIPROPIONATE

CREAM;TOPICAL

BETAMETHASONE DIPROPIONATEAB ACTAVIS MIDEQ 0.05% BASEA070885 001 Feb 03, 1987

ATLANTIC

AB COSETTEEQ 0.05% BASEA210217 001 Oct 12, 2018AB +! FOUGERA PHARMSEQ 0.05% BASEN019137 001 Jun 26, 1984AB TAROEQ 0.05% BASEA073552 001 Apr 30, 1992AB ZYDUS PHARMSEQ 0.05% BASEA208885 001 Jan 11, 2019

CREAM, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATEAB FOUGERA PHARMSEQ 0.05% BASEA076215 001 Dec 09, 2003AB ! GLENMARK GENERICSEQ 0.05% BASEA078930 001 Sep 23, 2008AB PADAGIS ISRAELEQ 0.05% BASEA076592 001 Dec 09, 2003AB TAROEQ 0.05% BASEA076543 001 Dec 09, 2003

GEL, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATEAB ! FOUGERA PHARMSEQ 0.05% BASEA075276 001 May 13, 2003AB TAROEQ 0.05% BASEA076508 001 Dec 02, 2003

LOTION;TOPICAL

BETAMETHASONE DIPROPIONATEAB AKORNEQ 0.05% BASEA209896 001 Feb 06, 2018AB COSETTEEQ 0.05% BASEA071467 001 Aug 10, 1987AB ! FOUGERA PHARMS INCEQ 0.05% BASEA070275 001 Aug 12, 1985AB PADAGIS USEQ 0.05% BASEA072538 001 Jan 31, 1990

LOTION, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATEAB AKORNEQ 0.05% BASEA208849 001 Oct 11, 2019AB FOUGERA PHARMSEQ 0.05% BASEA077111 001 May 21, 2007AB ! TAROEQ 0.05% BASEA077477 001 May 21, 2007AB TELIGENTEQ 0.05% BASEA206389 001 Feb 13, 2018

OINTMENT;TOPICAL

BETAMETHASONE DIPROPIONATEAB ACTAVIS MIDEQ 0.05% BASEA071012 001 Feb 03, 1987

ATLANTIC

AB CADILAEQ 0.05% BASEA214048 001 Jul 14, 2020AB +! FOUGERA PHARMS INCEQ 0.05% BASEN019141 001 Sep 04, 1984AB TAROEQ 0.05% BASEA074271 001 Sep 15, 1994

OINTMENT, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATEAB ACTAVIS MIDEQ 0.05% BASEA074304 001 Aug 31, 1995

ATLANTIC

AB FOUGERA PHARMSEQ 0.05% BASEA075373 001 Jun 22, 1999AB LUPIN LTDEQ 0.05% BASEA209106 001 Dec 18, 2019AB TAROEQ 0.05% BASEA076753 001 Oct 12, 2004AB TELIGENTEQ 0.05% BASEA206118 001 Nov 09, 2017DIPROLENEAB +! MERCK SHARP DOHMEEQ 0.05% BASEN018741 001 Jul 27, 1983

PRESCRIPTION DRUG PRODUCT LIST

BETAMETHASONE DIPROPIONATE

SPRAY;TOPICAL

SERNIVO

+! PRIMUS PHARMS EQ 0.05% BASE/SPRAY N208079 001 Feb 05, 2016

BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE

AEROSOL, FOAM;TOPICAL

ENSTILAR

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

CREAM;TOPICAL

WYNZORA

+! MC2 0.064%;0.005% N213422 001 Jul 20, 2020

OINTMENT;TOPICAL

CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE**AB** PADAGIS 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

CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE**AB** TARO PHARMS **0.064%;0.005%** **A213269 001** Sep 02, 2020**AB** TOLMAR **0.064%;0.005%** **A210765 001** May 11, 2020CALCIPOTRIENE AND BETHAMETHASONE DIPROPIONATE**AB** PADAGIS ISRAEL **0.064%;0.005%** **A212367 001** Sep 11, 2020TACLONEX**AB** +! LEO PHARMA AS **0.064%;0.005%** **N022185 001** May 09, 2008BETAMETHASONE 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, 2004BETAMETHASONE VALERATE

AEROSOL, FOAM;TOPICAL

BETAMETHASONE VALERATE**AB** PADAGIS ISRAEL **0.12%** **A078337 001** Nov 26, 2012**AB** TARO **0.12%** **A208204 001** May 24, 2017LUXIQ**AB** +! MYLAN **0.12%** **N020934 001** Feb 28, 1999

CREAM;TOPICAL

BETA-VAL**AB** COSETTE **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** ANIMA **EQ 0.1% BASE** **A070052 001** Jul 31, 1985**AB** +! FOUGERA PHARMS INC **EQ 0.1% BASE** **N018866 001** Aug 31, 1983

OINTMENT;TOPICAL

BETA-VAL**AB** COSETTE **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, 1983

PRESCRIPTION DRUG PRODUCT LIST

BETAXOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

BETAXOLOL HYDROCHLORIDE

<u>AT</u>	ACELLA	<u>EQ 0.5% BASE</u>	<u>A078694 001</u>	Nov 16, 2009
<u>AT</u>	AKORN	<u>EQ 0.5% BASE</u>	<u>A075386 001</u>	Jun 30, 2000
<u>AT</u>	MEDIMETRIKS PHARMS	<u>EQ 0.5% BASE</u>	<u>A075630 001</u>	Apr 12, 2001

BETOPTIC

<u>AT</u>	+! SANDOZ INC	<u>EQ 0.5% BASE</u>	<u>N019270 001</u>	Aug 30, 1985
	SUSPENSION/DROPS;OPHTHALMIC			
	BETOPTIC S			
	+! NOVARTIS	EQ 0.25% BASE	N019845 001	Dec 29, 1989
	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>		BIOPHARM	<u>5MG</u>	<u>A040728 002</u>	Oct 26, 2007
<u>AA</u>			<u>10MG</u>	<u>A040728 003</u>	Oct 26, 2007
<u>AA</u>			<u>25MG</u>	<u>A040728 004</u>	Oct 26, 2007
<u>AA</u>			<u>50MG</u>	<u>A040728 001</u>	Oct 26, 2007
<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>DUVOID</u>				
<u>AA</u>		CHARTWELL RX	<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>	

BEXAROTENE

CAPSULE;ORAL

BEXAROTENE

<u>AB</u>		AMNEAL PHARMS NY	<u>75MG</u>	<u>A210105 001</u>	Sep 04, 2018
<u>AB</u>		ANI PHARMS	<u>75MG</u>	<u>A209861 001</u>	May 08, 2018
<u>AB</u>		BIONPHARMA INC	<u>75MG</u>	<u>A203174 001</u>	Aug 12, 2014
<u>AB</u>		HIKMA	<u>75MG</u>	<u>A203663 001</u>	Jun 16, 2020
<u>AB</u>		TEVA PHARMS USA	<u>75MG</u>	<u>A209931 001</u>	Jan 14, 2021
<u>AB</u>		UPSHER SMITH LABS	<u>75MG</u>	<u>A209886 001</u>	Jul 25, 2018

TARGRETIN

<u>AB</u>	+!	VALEANT LUXEMBOURG	<u>75MG</u>	<u>N021055 001</u>	Dec 29, 1999
		GEL;TOPICAL			
		TARGRETIN			
		+! BAUSCH	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>		KENTON	<u>50MG</u>	<u>A091011 001</u>	Jun 10, 2015
<u>AB</u>		SANDOZ	<u>50MG</u>	<u>A078575 001</u>	Jul 06, 2009
<u>AB</u>		SUN PHARM	<u>50MG</u>	<u>A079110 001</u>	Jul 06, 2009
<u>AB</u>		WATSON LABS TEVA	<u>50MG</u>	<u>A078634 001</u>	Aug 28, 2009
<u>AB</u>		ZYDUS PHARMS USA INC	<u>50MG</u>	<u>A079089 001</u>	Jul 06, 2009

CASODEX

<u>AB</u>	+!	ANI PHARMS	<u>50MG</u>	<u>N020498 001</u>	Oct 04, 1995
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PRESCRIPTION DRUG PRODUCT LIST

BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

BIKTARVY

+	GILEAD SCIENCES INC	EQ 30MG BASE;120MG;EQ 15MG BASE	N210251	002	Oct 07, 2021
+	!	EQ 50MG BASE;200MG;EQ 25MG BASE	N210251	001	Feb 07, 2018

BIMATOPROST

IMPLANT;OPHTHALMIC

DURYSTA

+	ALLERGAN INC	10MCG	N211911	001	Mar 04, 2020
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SOLUTION/DROPS;OPHTHALMIC

BIMATOPROST

AT	!	ALEMBIC PHARMS LTD	<u>0.03%</u>	<u>A210263</u>	<u>001</u>	Apr 12, 2019
AT		APOTEX INC	<u>0.03%</u>	<u>A090449</u>	<u>001</u>	Jul 20, 2015
AT		GLAND PHARMA LTD	<u>0.03%</u>	<u>A210126</u>	<u>001</u>	Mar 22, 2019
AT		LUPIN LTD	<u>0.03%</u>	<u>A203991</u>	<u>001</u>	Feb 20, 2015
AT		MICRO LABS	<u>0.03%</u>	<u>A202505</u>	<u>001</u>	Sep 08, 2020
AT		SANDOZ INC	<u>0.03%</u>	<u>A202565</u>	<u>001</u>	May 05, 2015
AT		SOMERSET THERAPS LLC	<u>0.03%</u>	<u>A207601</u>	<u>001</u>	Jun 19, 2019

LUMIGAN

+	ALLERGAN	0.01%	N022184	001	Aug 31, 2010
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SOLUTION/DROPS;TOPICAL

BIMATOPROST

AT		AKORN	<u>0.03%</u>	<u>A203051</u>	<u>001</u>	Oct 09, 2018
AT		ALEMBIC PHARMS LTD	<u>0.03%</u>	<u>A210515</u>	<u>001</u>	Jan 21, 2020
AT		APOTEX INC	<u>0.03%</u>	<u>A201894</u>	<u>001</u>	Dec 01, 2014
AT		SANDOZ INC	<u>0.03%</u>	<u>A202719</u>	<u>001</u>	Apr 19, 2016

LATISSE

AT	+	ALLERGAN	<u>0.03%</u>	<u>N022369</u>	<u>001</u>	Dec 24, 2008
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BINIMETINIB

TABLET;ORAL

MEKTOVI

+	ARRAY BIOPHARMA INC	15MG	N210498	001	Jun 27, 2018
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BISMUTH SUBCITRATE POTASSIUM; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

CAPSULE;ORAL

PYLERA

+	ALLERGAN	140MG;125MG;125MG	N050786	001	Sep 28, 2006
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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
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BISOPROLOL FUMARATE

TABLET;ORAL

BISOPROLOL FUMARATE

AB		ALEMBIC LABS	<u>5MG</u>	<u>A204891</u>	<u>001</u>	Jan 11, 2017
AB			<u>10MG</u>	<u>A204891</u>	<u>002</u>	Jan 11, 2017
AB		AUROBINDO PHARMA	<u>5MG</u>	<u>A077910</u>	<u>001</u>	Dec 27, 2006
AB			<u>10MG</u>	<u>A077910</u>	<u>002</u>	Dec 27, 2006
AB		FRONTIDA BIOPHARM	<u>5MG</u>	<u>A075474</u>	<u>001</u>	Oct 25, 2002
AB			<u>10MG</u>	<u>A075474</u>	<u>002</u>	Oct 25, 2002
AB		NOVITIUM PHARMA	<u>5MG</u>	<u>A215563</u>	<u>001</u>	Oct 29, 2021
AB			<u>10MG</u>	<u>A215563</u>	<u>002</u>	Oct 29, 2021
AB		RUBICON	<u>5MG</u>	<u>A075643</u>	<u>001</u>	Nov 16, 2000
AB			<u>10MG</u>	<u>A075643</u>	<u>002</u>	Nov 16, 2000
AB		UNICHEM	<u>5MG</u>	<u>A078635</u>	<u>001</u>	Aug 18, 2009
AB	!		<u>10MG</u>	<u>A078635</u>	<u>002</u>	Aug 18, 2009

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

AB		EDENBRIDGE PHARMS	<u>2.5MG;6.25MG</u>	<u>A212678</u>	<u>001</u>	Jul 09, 2020
AB			<u>5MG;6.25MG</u>	<u>A212678</u>	<u>002</u>	Jul 09, 2020
AB			<u>10MG;6.25MG</u>	<u>A212678</u>	<u>003</u>	Jul 09, 2020
AB		MYLAN	<u>2.5MG;6.25MG</u>	<u>A075768</u>	<u>001</u>	Sep 25, 2000
AB			<u>5MG;6.25MG</u>	<u>A075768</u>	<u>002</u>	Sep 25, 2000
AB			<u>10MG;6.25MG</u>	<u>A075768</u>	<u>003</u>	Sep 25, 2000
AB		NOVITIUM PHARMA	<u>2.5MG;6.25MG</u>	<u>A215562</u>	<u>001</u>	Nov 04, 2021
AB			<u>5MG;6.25MG</u>	<u>A215562</u>	<u>002</u>	Nov 04, 2021
AB			<u>10MG;6.25MG</u>	<u>A215562</u>	<u>003</u>	Nov 04, 2021
AB		SANDOZ	<u>2.5MG;6.25MG</u>	<u>A075579</u>	<u>001</u>	Sep 25, 2000

PRESCRIPTION DRUG PRODUCT LIST

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

<u>AB</u>		<u>5MG; 6.25MG</u>	<u>A075579 002</u>	Sep 25, 2000
<u>AB</u>		<u>10MG; 6.25MG</u>	<u>A075579 003</u>	Sep 25, 2000
<u>AB</u>	UNICHEM	<u>2.5MG; 6.25MG</u>	<u>A079106 001</u>	Jul 28, 2010
<u>AB</u>		<u>5MG; 6.25MG</u>	<u>A079106 002</u>	Jul 28, 2010
<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
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BIVALIRUDIN

<u>AP</u>		ACCORD HLTHCARE	<u>250MG/VIAL</u>	<u>A206551 001</u> Nov 22, 2017
<u>AP</u>		DR REDDYS LABS LTD	<u>250MG/VIAL</u>	<u>A201577 001</u> May 26, 2017
<u>AP</u>		EUGIA PHARMA	<u>250MG/VIAL</u>	<u>A205962 001</u> Jul 27, 2018
<u>AP</u>		FRESENIUS KABI USA	<u>250MG/VIAL</u>	<u>A090189 001</u> Oct 28, 2016
<u>AP</u>		HAINAN POLY PHARM	<u>250MG/VIAL</u>	<u>A213078 001</u> May 28, 2021
<u>AP</u>		HONG KONG	<u>250MG/VIAL</u>	<u>A091602 001</u> Jul 16, 2018
<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
<u>AP</u>		SHUANGCHENG	<u>250MG/VIAL</u>	<u>A210031 001</u> Oct 23, 2019

SOLUTION; INTRAVENOUS

ANGIOMAX RTU

	+	MAIA PHARMS INC	250MG/50ML (5MG/ML)	N211215 001 Jul 25, 2019
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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>		HIKMA	<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
<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>		MEITHEAL	<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>		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

BORTEZOMIB

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

VELCADE

	+	TAKEDA PHARMS USA	3.5MG/VIAL	N021602 001 May 13, 2003
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POWDER; INTRAVENOUS

BORTEZOMIB

		DR REDDYS LABS LTD	3.5MG/VIAL	N206927 001 Oct 04, 2019
		FRESENIUS KABI USA	3.5MG/VIAL	N205004 001 Nov 06, 2017

BOSENTAN

TABLET; ORAL

BOSENTAN

<u>AB</u>		HIKMA	<u>62.5MG</u>	<u>A208695 001</u> Apr 26, 2019
<u>AB</u>			<u>125MG</u>	<u>A208695 002</u> Apr 26, 2019
<u>AB</u>		PAR PHARM INC	<u>62.5MG</u>	<u>A205699 001</u> Apr 26, 2019
<u>AB</u>			<u>125MG</u>	<u>A205699 002</u> Apr 26, 2019
<u>AB</u>		SUN PHARM	<u>62.5MG</u>	<u>A209324 001</u> Apr 26, 2019
<u>AB</u>			<u>125MG</u>	<u>A209324 002</u> Apr 26, 2019
<u>AB</u>		WATSON LABS INC	<u>62.5MG</u>	<u>A207110 001</u> Apr 26, 2019
<u>AB</u>			<u>125MG</u>	<u>A207110 002</u> Apr 26, 2019
<u>AB</u>		ZYDUS PHARMS	<u>62.5MG</u>	<u>A207760 001</u> Apr 26, 2019
<u>AB</u>			<u>125MG</u>	<u>A207760 002</u> Apr 26, 2019

TRACLEER

<u>AB</u>	+	ACTELION	<u>62.5MG</u>	<u>N021290 001</u> Nov 20, 2001
<u>AB</u>	+		<u>125MG</u>	<u>N021290 002</u> Nov 20, 2001

PRESCRIPTION DRUG PRODUCT LIST

BOSENTAN

TABLET, FOR SUSPENSION;ORAL

TRACLEER

+! ACTELION 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

BREMELANOTIDE ACETATE

SOLUTION;SUBCUTANEOUS

VYLEESI (AUTOINJECTOR)

+! PALATIN EQ 1.75MG BASE/0.3ML (EQ 1.75MG N210557 001 Jun 21, 2019
TECHNOLOGIES BASE/0.3 ML)BRETYLIUM TOSYLATE

INJECTABLE;INJECTION

BRETYLIUM TOSYLATE

! BRECKENRIDGE 50MG/ML A204386 001 Dec 21, 2018

BREXANOLONE

SOLUTION;INTRAVENOUS

ZULRESSO

+! SAGE THERAP 100MG/20ML (5MG/ML) N211371 001 Jun 17, 2019

BREXPIPIRAZOLE

TABLET;ORAL

REXULTI

+ OTSUKA 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

+ TAKEDA PHARMS USA 30MG N208772 001 Apr 28, 2017

+ 90MG N208772 002 Apr 28, 2017

+! 180MG N208772 003 Oct 02, 2017

BRILLIANT BLUE G

SOLUTION;OPHTHALMIC

TISSUEBLUE

+! DUTCH OPHTHALMIC 0.025% N209569 001 Dec 20, 2019

BRIMONIDINE TARTRATE

GEL;TOPICAL

BRIMONIDINE TARTRATEAB PADAGIS ISRAEL EQ 0.33% BASE A209158 001 Sep 23, 2021MIRVASOAB +! GALDERMA LABS LP EQ 0.33% BASE N204708 001 Aug 23, 2013

SOLUTION/DROPS;OPHTHALMIC

ALPHAGAN PAT +! ALLERGAN 0.15% N021262 001 Mar 16, 2001BRIMONIDINE TARTRATEAT AKORN 0.2% A076439 001 Mar 14, 2006AT ! BAUSCH AND LOMB 0.2% A076260 001 May 28, 2003AT INDOCO 0.2% A091691 001 Nov 18, 2014AT SANDOZ INC 0.2% A076254 001 Sep 16, 2003AT 0.2% A078075 001 Jan 30, 2008AT SOMERSET THERAPS 0.2% A208992 001 Mar 11, 2019
LLCOOLIANAAT +! SANDOZ INC 0.15% N021764 001 May 22, 2006

ALPHAGAN P

+! ALLERGAN 0.1% N021770 001 Aug 19, 2005

PRESCRIPTION DRUG PRODUCT LIST

BRIMONIDINE TARTRATE; BRINZOLAMIDE

SUSPENSION/DROPS;OPHTHALMIC

SIMBRINZA

+! ALCON LABS INC 0.2%;1% N204251 001 Apr 19, 2013

BRIMONIDINE TARTRATE; TIMOLOL MALEATE

SOLUTION/DROPS;OPHTHALMIC

COMBIGAN

+! ALLERGAN 0.2%;EQ 0.5% BASE N021398 001 Oct 30, 2007

BRINCIDOFIVIR

SUSPENSION;ORAL

TEMBEXA

+! CHIMERIX 10MG/ML N214460 001 Jun 04, 2021

TABLET;ORAL

TEMBEXA

+! CHIMERIX 100MG N214461 001 Jun 04, 2021

BRINZOLAMIDE

SUSPENSION/DROPS;OPHTHALMIC

AZOPT**AB** +! NOVARTIS **1%** **N020816 001** Apr 01, 1998BRINZOLAMIDE**AB** BAUSCH **1%** **A204884 001** Aug 18, 2021**AB** WATSON LABS INC **1%** **A209406 001** Nov 27, 2020BRIVARACETAM

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

BROMFENAC SODIUM

SOLUTION/DROPS;OPHTHALMIC

BROMFENAC SODIUM**AT2** ! AKORN **EQ 0.09% ACID** **A203395 001** Jan 22, 2014**AT2** ALEMBIC PHARMS LTD **EQ 0.09% ACID** **A210560 001** Jun 21, 2019**AT2** GLAND PHARMA LTD **EQ 0.09% ACID** **A211029 001** Mar 17, 2020

BROMSITE

+! SUN PHARM 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, 2008PARLODEL**AB** + VALIDUS PHARMS **EQ 5MG BASE** **N017962 002** Mar 01, 1982

TABLET;ORAL

BROMOCRIPTINE MESYLATE**AB** ! PADAGIS US **EQ 2.5MG BASE** **A077646 001** Oct 01, 2008**AB** SANDOZ INC **EQ 2.5MG BASE** **A074631 001** Jan 13, 1998PARLODEL**AB** + VALIDUS PHARMS **EQ 2.5MG BASE** **N017962 001**

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 **2MG/5ML;10MG/5ML;30MG/5ML** **A203375 001** Sep 20, 2016**AA** ALKEM LABS LTD **2MG/5ML;10MG/5ML;30MG/5ML** **A210647 001** Jul 14, 2020

PRESCRIPTION DRUG PRODUCT LIST

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP; ORAL

BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

<u>AA</u>	BRECKENRIDGE	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A203997 001</u>	Sep 30, 2020
<u>AA</u>	EYWA	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A211170 001</u>	Jun 16, 2020
<u>AA</u>	LANNETT CO INC	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A213125 001</u>	Apr 17, 2020
<u>AA</u>	MAYNE PHARMA INC	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A207676 001</u>	Dec 04, 2018
<u>AA</u>	PADAGIS US	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A205292 001</u>	Jul 15, 2014
<u>AA</u>	RHODES PHARMS	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A202955 001</u>	Nov 05, 2020
<u>AA</u>	TARO	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A205112 001</u>	Feb 27, 2017

BUDESONIDE

AEROSOL, FOAM; RECTAL

UCERIS

+! SALIX

2MG/ACTUATION

N205613 001 Oct 07, 2014

CAPSULE, DELAYED RELEASE; ORAL

BUDESONIDE

<u>AB</u>	AMNEAL PHARMS	<u>3MG</u>	<u>A206200 001</u>	Jul 31, 2017
<u>AB</u>	AUROBINDO PHARMA USA	<u>3MG</u>	<u>A090410 001</u>	May 16, 2011
<u>AB</u>	MAYNE PHARMA	<u>3MG</u>	<u>A206623 001</u>	Apr 08, 2016
<u>AB</u>	RISING	<u>3MG</u>	<u>A207367 001</u>	Apr 07, 2017
<u>AB</u>	SCIECURE PHARMA INC	<u>3MG</u>	<u>A209041 001</u>	Sep 28, 2017
<u>AB</u>	ZYDUS PHARMS	<u>3MG</u>	<u>A206134 001</u>	May 04, 2017

ENTOCORT EC

<u>AB</u>	+! PADAGIS US	<u>3MG</u>	<u>N021324 001</u>	Oct 02, 2001
	TARPEYO			
	+! CALLIDITAS	4MG	N215935 001	Dec 15, 2021
	CAPSULE, EXTENDED RELEASE; ORAL			
	ORTIKOS			
	+ SUN PHARM INDS INC	6MG	N211929 001	Jun 13, 2019
	+!	9MG	N211929 002	Jun 13, 2019

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

<u>AN</u>	CIPLA	<u>0.25MG/2ML</u>	<u>A205710 001</u>	Nov 16, 2017
<u>AN</u>		<u>0.5MG/2ML</u>	<u>A205710 002</u>	Nov 16, 2017
<u>AN</u>		<u>1MG/2ML</u>	<u>A205710 003</u>	Nov 16, 2017
<u>AN</u>	IMPAX LABS INC	<u>0.25MG/2ML</u>	<u>A078404 001</u>	Jul 31, 2012
<u>AN</u>		<u>0.5MG/2ML</u>	<u>A078404 002</u>	Jul 31, 2012
<u>AN</u>	LUPIN	<u>0.5MG/2ML</u>	<u>A210897 001</u>	Nov 09, 2018
<u>AN</u>	NEPHRON	<u>0.25MG/2ML</u>	<u>A078202 001</u>	Mar 30, 2009
<u>AN</u>		<u>0.5MG/2ML</u>	<u>A078202 002</u>	Mar 30, 2009
<u>AN</u>	SANDOZ INC	<u>0.25MG/2ML</u>	<u>A201966 003</u>	Sep 27, 2013
<u>AN</u>		<u>0.5MG/2ML</u>	<u>A201966 002</u>	Sep 27, 2013
<u>AN</u>		<u>1MG/2ML</u>	<u>A201966 001</u>	Sep 27, 2013
<u>AN</u>	SUN PHARM	<u>0.25MG/2ML</u>	<u>A211922 001</u>	Apr 14, 2021
<u>AN</u>		<u>0.5MG/2ML</u>	<u>A211922 002</u>	Apr 14, 2021
<u>AN</u>		<u>1MG/2ML</u>	<u>A211922 003</u>	Apr 14, 2021
<u>AN</u>	TEVA PHARMS	<u>0.25MG/2ML</u>	<u>A077519 001</u>	Nov 18, 2008
<u>AN</u>		<u>0.5MG/2ML</u>	<u>A077519 002</u>	Nov 18, 2008
<u>AN</u>	TEVA PHARMS USA	<u>1MG/2ML</u>	<u>A204548 001</u>	Mar 08, 2016

PULMICORT RESPULES

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

TABLET, EXTENDED RELEASE; ORAL

BUDESONIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>9MG</u>	<u>A205457 001</u>	Jul 03, 2018
<u>AB</u>	MYLAN	<u>9MG</u>	<u>A208851 001</u>	Sep 17, 2020

UCERIS

<u>AB</u>	+! SALIX	<u>9MG</u>	<u>N203634 001</u>	Jan 14, 2013
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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

PRESCRIPTION DRUG PRODUCT LIST

BUDESONIDE; FORMOTEROL FUMARATE; GLYCOPYRROLATE

AEROSOL, METERED; INHALATION

BREZTRI AEROSPHERE

+! ASTRAZENECA AB 0.16MG/INH;0.0048MG/INH;0.009MG/INH N212122 001 Jul 23, 2020

BUMETANIDE

INJECTABLE; INJECTION

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

TABLET; ORAL

BUMETANIDEAB AMNEAL PHARMS CO 0.5MG A209724 001 Oct 18, 2017AB 1MG A209724 002 Oct 18, 2017AB 2MG A209724 003 Oct 18, 2017AB HERITAGE PHARMA 0.5MG A074225 001 Apr 24, 1995AB 1MG A074225 002 Apr 24, 1995AB 2MG A074225 003 Apr 24, 1995AB SANDOZ 0.5MG A074700 001 Nov 21, 1996AB 1MG A074700 002 Nov 21, 1996AB ! 2MG A074700 003 Nov 21, 1996AB UPSHER SMITH LABS 0.5MG A209916 001 Jan 23, 2018AB 1MG A209916 002 Jan 23, 2018AB 2MG A209916 003 Jan 23, 2018AB ZYDUS PHARMS 0.5MG A202900 001 Apr 30, 2018AB 1MG A202900 002 Apr 30, 2018AB 2MG A202900 003 Apr 30, 2018BUMEXAB + VALIDUS PHARMS 0.5MG N018225 002 Feb 28, 1983AB + 1MG N018225 001 Feb 28, 1983AB + 2MG N018225 003 Jun 14, 1985

BUMETANIDE

BX RISING 0.5MG A212019 001 Dec 12, 2019

BX 1MG A212019 002 Dec 12, 2019

BX 2MG A212019 003 Dec 12, 2019

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

IMPLANT; IMPLANTATION

XARACOLL

+! INNOCOLL PHARMS 100MG N209511 001 Aug 28, 2020

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDEAP EUGIA PHARMA 0.25% A207183 001 May 13, 2016AP 0.5% A207183 002 May 13, 2016AP HIKMA PHARMS 0.25% A205141 001 Feb 11, 2021AP 0.5% A205141 002 Feb 11, 2021AP HOSPIRA 0.25% A070583 001 Feb 17, 1987AP 0.25% A070590 001 Feb 17, 1987AP 0.5% A070584 001 Feb 17, 1986AP 0.5% A070597 001 Mar 03, 1987AP 0.5% A070609 001 Mar 03, 1987AP 0.75% A070585 001 Mar 03, 1987BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREEAP EUGIA PHARMA 0.25% A203895 001 Nov 05, 2013AP 0.5% A203895 002 Nov 05, 2013AP 0.75% A203895 003 Nov 05, 2013AP HIKMA PHARMS 0.25% A204842 001 Feb 11, 2021AP 0.5% A204842 002 Feb 11, 2021AP 0.75% A204842 003 Feb 11, 2021MARCAINE HYDROCHLORIDEAP +! HOSPIRA 0.25% N016964 001AP +! 0.5% N016964 006MARCAINE HYDROCHLORIDE PRESERVATIVE FREEAP +! HOSPIRA 0.25% N016964 012AP +! 0.5% N016964 005AP +! 0.75% N016964 009

PRESCRIPTION DRUG PRODUCT LIST

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

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>	B BRAUN MEDICAL INC	<u>0.75%</u>	<u>A209087 001</u>	Apr 02, 2019
<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
<u>AP</u>	HUONS	<u>0.75%</u>	<u>A212822 001</u>	Dec 30, 2019

MARCAINE

<u>AP</u>	+! HOSPIRA	<u>0.75%</u>	<u>N018692 001</u>	May 04, 1984
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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
	!	0.25%;0.005MG/ML	A071165 001	Jun 16, 1988
		0.25%;0.005MG/ML	A071167 001	Jun 16, 1988

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

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
	BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE			
	! SEPTODONT	0.5%;0.0091MG/ML	A077250 001	Sep 27, 2006

BUPIVACAINE; MELOXICAM

SOLUTION, EXTENDED RELEASE; PERIARTICULAR

ZYNRELEF KIT

	+! HERON THERAPS INC	60MG/2.3ML (29.25MG/ML); 1.8MG/2.3ML (0.88MG/ML)	N211988 001	May 12, 2021
	+!	200MG/7ML (29.25MG/ML); 6MG/7ML (0.88MG/ML)	N211988 002	May 12, 2021
	+!	300MG/10.5ML (29.25MG/ML); 9MG/10.5ML (0.88MG/ML)	N211988 003	May 12, 2021
	+!	400MG/14ML (29.25MG/ML); 12MG/14ML (0.88MG/ML)	N211988 004	May 12, 2021

BUPRENORPHINE

FILM, EXTENDED RELEASE; TRANSDERMAL

BUPRENORPHINE

<u>AB</u>	AMNEAL	<u>5MCG/HR</u>	<u>A211586 001</u>	Apr 14, 2020
<u>AB</u>		<u>7.5MCG/HR</u>	<u>A211586 002</u>	Apr 14, 2020
<u>AB</u>		<u>10MCG/HR</u>	<u>A211586 003</u>	Apr 14, 2020
<u>AB</u>		<u>15MCG/HR</u>	<u>A211586 004</u>	Apr 14, 2020
<u>AB</u>		<u>20MCG/HR</u>	<u>A211586 005</u>	Apr 14, 2020
<u>AB</u>	AVEVA	<u>5MCG/HR</u>	<u>A210272 001</u>	Sep 23, 2021
<u>AB</u>		<u>7.5MCG/HR</u>	<u>A210272 002</u>	Sep 23, 2021
<u>AB</u>		<u>10MCG/HR</u>	<u>A210272 003</u>	Sep 23, 2021
<u>AB</u>		<u>15MCG/HR</u>	<u>A210272 004</u>	Sep 23, 2021
<u>AB</u>		<u>20MCG/HR</u>	<u>A210272 005</u>	Sep 23, 2021
<u>AB</u>	WATSON LABS TEVA	<u>5MCG/HR</u>	<u>A204937 001</u>	Nov 20, 2018
<u>AB</u>		<u>7.5MCG/HR</u>	<u>A204937 005</u>	Jun 29, 2021
<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

PRESCRIPTION DRUG PRODUCT LIST

BUPRENORPHINE

FILM, EXTENDED RELEASE;TRANSDERMAL

BUTRANS

<u>AB</u>	+	PURDUE PHARMA LP	<u>5MCG/HR</u>	<u>N021306 001</u>	Jun 30, 2010
<u>AB</u>	+		<u>7.5MCG/HR</u>	<u>N021306 005</u>	Jun 30, 2014
<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

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

<u>AB</u>	+	BDSI	<u>EQ 0.075MG BASE</u>	<u>N207932 001</u>	Oct 23, 2015
<u>AB</u>	+		<u>EQ 0.15MG BASE</u>	<u>N207932 002</u>	Oct 23, 2015
<u>AB</u>	+		<u>EQ 0.3MG BASE</u>	<u>N207932 003</u>	Oct 23, 2015
<u>AB</u>	+		<u>EQ 0.45MG BASE</u>	<u>N207932 004</u>	Oct 23, 2015
<u>AB</u>	+		<u>EQ 0.6MG BASE</u>	<u>N207932 005</u>	Oct 23, 2015
<u>AB</u>	+		<u>EQ 0.75MG BASE</u>	<u>N207932 006</u>	Oct 23, 2015
<u>AB</u>	+		<u>EQ 0.9MG BASE</u>	<u>N207932 007</u>	Oct 23, 2015

BUPRENORPHINE HYDROCHLORIDE

<u>AB</u>		ALVOGEN	<u>EQ 0.075MG BASE</u>	<u>A211594 001</u>	Aug 03, 2021
<u>AB</u>			<u>EQ 0.15MG BASE</u>	<u>A211594 002</u>	Aug 03, 2021
<u>AB</u>			<u>EQ 0.3MG BASE</u>	<u>A211594 003</u>	Aug 03, 2021
<u>AB</u>			<u>EQ 0.45MG BASE</u>	<u>A211594 004</u>	Aug 03, 2021
<u>AB</u>			<u>EQ 0.6MG BASE</u>	<u>A211594 005</u>	Aug 03, 2021
<u>AB</u>			<u>EQ 0.75MG BASE</u>	<u>A211594 006</u>	Aug 03, 2021
<u>AB</u>			<u>EQ 0.9MG BASE</u>	<u>A211594 007</u>	Aug 03, 2021

INJECTABLE;INJECTION

BUPRENEX

<u>AP</u>	+	INDIVIOR INC	<u>EQ 0.3MG BASE/ML</u>	<u>N018401 001</u>	
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BUPRENORPHINE HYDROCHLORIDE

<u>AP</u>		AM REGENT	<u>EQ 0.3MG BASE/ML</u>	<u>A078331 001</u>	Mar 27, 2007
<u>AP</u>		HIKMA	<u>EQ 0.3MG BASE/ML</u>	<u>A076931 001</u>	Mar 02, 2005
<u>AP</u>		HOSPIRA	<u>EQ 0.7MG BASE/ML</u>	<u>A074137 001</u>	Jun 03, 1996
<u>AP</u>		PAR STERILE PRODUCTS	<u>EQ 0.3MG BASE/ML</u>	<u>A206586 001</u>	Jul 28, 2015

TABLET;SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE

<u>AB</u>		ACTAVIS ELIZABETH	<u>EQ 2MG BASE</u>	<u>A090819 001</u>	Feb 19, 2015
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A090819 002</u>	Feb 19, 2015
<u>AB</u>		ETHYPHARM	<u>EQ 2MG BASE</u>	<u>A090622 001</u>	Sep 24, 2010
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A090622 002</u>	Sep 24, 2010
<u>AB</u>		HIKMA	<u>EQ 2MG BASE</u>	<u>A078633 001</u>	Oct 08, 2009
<u>AB</u>	!		<u>EQ 8MG BASE</u>	<u>A078633 002</u>	Oct 08, 2009
<u>AB</u>		RHODES PHARMS	<u>EQ 2MG BASE</u>	<u>A207276 001</u>	Mar 27, 2017
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A207276 002</u>	Mar 27, 2017
<u>AB</u>		RUBICON	<u>EQ 2MG BASE</u>	<u>A090279 001</u>	Jun 10, 2015
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A090279 002</u>	Jun 10, 2015
<u>AB</u>		SUN PHARM	<u>EQ 2MG BASE</u>	<u>A201760 001</u>	Jan 29, 2016
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A201760 002</u>	Jan 29, 2016

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

FILM;BUCCAL, SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

<u>AB</u>		ALVOGEN	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A205954 001</u>	Jan 24, 2019
<u>AB</u>			<u>EQ 4MG BASE;EQ 1MG BASE</u>	<u>A205954 002</u>	Jan 24, 2019
<u>AB</u>			<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A205954 003</u>	Jan 24, 2019
<u>AB</u>			<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>A205954 004</u>	Jan 24, 2019
<u>AB</u>		DR REDDYS LABS SA	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A205299 001</u>	Jun 14, 2018
<u>AB</u>			<u>EQ 4MG BASE;EQ 1MG BASE</u>	<u>A205806 001</u>	Jun 14, 2018
<u>AB</u>			<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A205299 002</u>	Jun 14, 2018
<u>AB</u>			<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>A205806 002</u>	Jun 14, 2018
<u>AB</u>		MYLAN TECHNOLOGIES	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A211785 001</u>	Apr 17, 2020
<u>AB</u>			<u>EQ 4MG BASE;EQ 1MG BASE</u>	<u>A211785 002</u>	Apr 17, 2020
<u>AB</u>			<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A207607 001</u>	Jun 14, 2018
<u>AB</u>			<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>A207607 002</u>	Jun 14, 2018

SUBOXONE

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

PRESCRIPTION DRUG PRODUCT LIST

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

FILM;BUCCAL, SUBLINGUAL

SUBOXONE

<u>AB</u>	+	!	<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>N022410 004</u>	Aug 10, 2012
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>		ALKEM LABS LTD	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A214930 001</u>	Jun 15, 2021
<u>AB</u>			<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A214930 002</u>	Jun 15, 2021
<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>		HIKMA	<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
<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>		RHODES PHARMS	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A205601 001</u>	Mar 30, 2020
<u>AB</u>			<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A205601 002</u>	Mar 30, 2020
<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	<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>		WES PHARMA INC	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A209069 001</u>	Jul 17, 2020
<u>AB</u>			<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A209069 002</u>	Jul 17, 2020
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

	+	BAUSCH	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>		APNAR PHARMA LP	<u>75MG</u>	<u>A075584 001</u>	Feb 07, 2000
<u>AB</u>			<u>100MG</u>	<u>A075584 002</u>	Feb 07, 2000
<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>		CADILA PHARMS LTD	<u>75MG</u>	<u>A208606 001</u>	Jan 16, 2020
<u>AB</u>			<u>100MG</u>	<u>A208606 002</u>	Jan 16, 2020
<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>		MICRO LABS	<u>75MG</u>	<u>A207403 001</u>	Apr 17, 2020
<u>AB</u>			<u>100MG</u>	<u>A207403 002</u>	Apr 17, 2020

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>		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>		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

PRESCRIPTION DRUG PRODUCT LIST

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

BUPROPION HYDROCHLORIDE

<u>AB1</u>		<u>200MG</u>	<u>A202304</u>	<u>003</u>	May 26, 2015
<u>AB1</u>	SANDOZ	<u>100MG</u>	<u>A075932</u>	<u>001</u>	Nov 25, 2003
<u>AB1</u>		<u>150MG</u>	<u>A075932</u>	<u>002</u>	Mar 22, 2004
<u>AB1</u>		<u>200MG</u>	<u>A075932</u>	<u>003</u>	Jun 22, 2005
<u>AB1</u>	SCIEGEN PHARMS INC	<u>100MG</u>	<u>A205794</u>	<u>001</u>	Mar 01, 2016
<u>AB1</u>		<u>150MG</u>	<u>A205794</u>	<u>002</u>	Mar 01, 2016
<u>AB1</u>		<u>200MG</u>	<u>A205794</u>	<u>003</u>	Mar 01, 2016
<u>AB1</u>	SUN PHARM	<u>100MG</u>	<u>A078866</u>	<u>001</u>	Apr 06, 2010
<u>AB1</u>		<u>150MG</u>	<u>A078866</u>	<u>002</u>	Apr 06, 2010
<u>AB1</u>		<u>200MG</u>	<u>A078866</u>	<u>003</u>	Apr 06, 2010
<u>AB1</u>	YICHANG HUMANWELL	<u>100MG</u>	<u>A211347</u>	<u>001</u>	Oct 16, 2018
<u>AB1</u>		<u>150MG</u>	<u>A211347</u>	<u>002</u>	Oct 16, 2018
<u>AB1</u>		<u>200MG</u>	<u>A211347</u>	<u>003</u>	Oct 16, 2018

WELLBUTRIN SR

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

BUPROPION HYDROCHLORIDE

<u>AB2</u>	!	ACTAVIS LABS FL INC	<u>150MG</u>	<u>A079094</u>	<u>001</u>	Mar 24, 2009
<u>AB2</u>		ANCHEN PHARMS	<u>150MG</u>	<u>A091520</u>	<u>001</u>	Jun 09, 2011
<u>AB2</u>		IMPAX LABS	<u>150MG</u>	<u>A075914</u>	<u>001</u>	May 27, 2004
<u>AB2</u>		SANDOZ INC	<u>150MG</u>	<u>A077475</u>	<u>001</u>	Mar 12, 2008
<u>AB2</u>		SCIEGEN PHARMS INC	<u>150MG</u>	<u>A206122</u>	<u>001</u>	Aug 17, 2016
<u>AB3</u>		ACCORD HLTHCARE	<u>150MG</u>	<u>A210497</u>	<u>001</u>	Oct 31, 2018
<u>AB3</u>			<u>300MG</u>	<u>A210497</u>	<u>002</u>	Oct 31, 2018
<u>AB3</u>		ACTAVIS LABS FL INC	<u>150MG</u>	<u>A077715</u>	<u>001</u>	Nov 26, 2008
<u>AB3</u>		ADAPTIS	<u>150MG</u>	<u>A211020</u>	<u>001</u>	Jan 28, 2019
<u>AB3</u>			<u>300MG</u>	<u>A211020</u>	<u>002</u>	Jan 28, 2019
<u>AB3</u>		ANBISON LAB	<u>150MG</u>	<u>A207224</u>	<u>001</u>	Jun 30, 2017
<u>AB3</u>			<u>300MG</u>	<u>A207224</u>	<u>002</u>	Jun 30, 2017
<u>AB3</u>		ANCHEN PHARMS	<u>150MG</u>	<u>A077284</u>	<u>001</u>	Dec 14, 2006
<u>AB3</u>			<u>300MG</u>	<u>A077284</u>	<u>002</u>	Dec 14, 2006
<u>AB3</u>		INVAGEN PHARMS	<u>150MG</u>	<u>A206556</u>	<u>001</u>	Aug 26, 2016
<u>AB3</u>			<u>300MG</u>	<u>A206556</u>	<u>002</u>	Aug 26, 2016
<u>AB3</u>		LUPIN LTD	<u>150MG</u>	<u>A090693</u>	<u>001</u>	Apr 06, 2017
<u>AB3</u>			<u>300MG</u>	<u>A090693</u>	<u>002</u>	Apr 06, 2017
<u>AB3</u>		SCIEGEN PHARMS INC	<u>150MG</u>	<u>A207479</u>	<u>001</u>	Apr 12, 2017
<u>AB3</u>			<u>300MG</u>	<u>A207479</u>	<u>002</u>	Apr 12, 2017
<u>AB3</u>		SINOTHERAPEUTICS INC	<u>150MG</u>	<u>A208652</u>	<u>001</u>	Aug 21, 2017
<u>AB3</u>			<u>300MG</u>	<u>A208652</u>	<u>002</u>	Aug 21, 2017
<u>AB3</u>		SUN PHARM	<u>150MG</u>	<u>A200216</u>	<u>001</u>	Nov 30, 2020
<u>AB3</u>			<u>300MG</u>	<u>A203650</u>	<u>001</u>	Dec 31, 2020
<u>AB3</u>		TWI PHARMS	<u>150MG</u>	<u>A210081</u>	<u>001</u>	Nov 03, 2017
<u>AB3</u>			<u>300MG</u>	<u>A210081</u>	<u>002</u>	Nov 03, 2017
<u>AB3</u>		WATSON LABS INC	<u>150MG</u>	<u>A077285</u>	<u>001</u>	Nov 26, 2008
<u>AB3</u>			<u>300MG</u>	<u>A077285</u>	<u>002</u>	Aug 15, 2008
<u>AB3</u>		WOCKHARDT LTD	<u>150MG</u>	<u>A202189</u>	<u>001</u>	Nov 21, 2012
<u>AB3</u>		YICHANG HUMANWELL	<u>150MG</u>	<u>A210015</u>	<u>001</u>	Jun 14, 2018
<u>AB3</u>			<u>300MG</u>	<u>A210015</u>	<u>002</u>	Jun 14, 2018
<u>AB3</u>		ZHEJIANG JUTAI PHARM	<u>150MG</u>	<u>A211200</u>	<u>002</u>	Apr 29, 2020
<u>AB3</u>			<u>300MG</u>	<u>A211200</u>	<u>001</u>	Sep 05, 2019
<u>AB3</u>		ZYDUS PHARMS	<u>150MG</u>	<u>A201567</u>	<u>002</u>	Jul 23, 2018
<u>AB3</u>			<u>300MG</u>	<u>A201567</u>	<u>001</u>	Jan 17, 2014

WELLBUTRIN XL

<u>AB3</u>	+	BAUSCH	<u>150MG</u>	<u>N021515</u>	<u>001</u>	Aug 28, 2003
<u>AB3</u>	+	!	<u>300MG</u>	<u>N021515</u>	<u>002</u>	Aug 28, 2003
		FORFIVO XL				
		+	ALMATICA	450MG	N022497	001
						Nov 10, 2011

BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CONTRAIVE

+	!	NALPROPION	90MG; 8MG	N200063	001	Sep 10, 2014
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PRESCRIPTION DRUG PRODUCT LIST

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>7.5MG</u>	<u>A078246 005</u>	Feb 21, 2020
<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>	EPIC PHARMA LLC	<u>5MG</u>	<u>A208972 001</u>	Apr 16, 2019
<u>AB</u>		<u>7.5MG</u>	<u>A208972 002</u>	Apr 16, 2019
<u>AB</u>		<u>10MG</u>	<u>A208972 003</u>	Apr 16, 2019
<u>AB</u>		<u>15MG</u>	<u>A208972 004</u>	Apr 16, 2019
<u>AB</u>		<u>30MG</u>	<u>A208972 005</u>	Apr 16, 2019
<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>5MG</u>	<u>A075388 001</u>	May 09, 2002
<u>AB</u>		<u>10MG</u>	<u>A075388 002</u>	May 09, 2002
<u>AB</u>		<u>15MG</u>	<u>A075388 003</u>	May 09, 2002
<u>AB</u>		<u>30MG</u>	<u>A078302 001</u>	Dec 17, 2007
<u>AB</u>	RUBICON	<u>5MG</u>	<u>A075521 001</u>	Apr 05, 2002
<u>AB</u>		<u>7.5MG</u>	<u>A075521 004</u>	Mar 16, 2021
<u>AB</u>		<u>10MG</u>	<u>A075521 002</u>	Apr 05, 2002
<u>AB</u>		<u>15MG</u>	<u>A075521 003</u>	Apr 05, 2002
<u>AB</u>		<u>30MG</u>	<u>A075521 005</u>	Mar 16, 2021
<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>	UNICHEM	<u>5MG</u>	<u>A210907 001</u>	Nov 14, 2019
<u>AB</u>		<u>10MG</u>	<u>A210907 002</u>	Nov 14, 2019
<u>AB</u>		<u>15MG</u>	<u>A210907 003</u>	Nov 14, 2019
<u>AB</u>		<u>30MG</u>	<u>A210907 004</u>	Nov 14, 2019
<u>AB</u>	YILING	<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	<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

PRESCRIPTION DRUG PRODUCT LIST

BUSULFAN

INJECTABLE; INJECTION

BUSULFAN

AP	ACCORD HLTHCARE INC	6MG/ML	A210148 001	Feb 22, 2019
AP	AM REGENT	6MG/ML	A202259 001	Dec 22, 2015
AP	AMNEAL	6MG/ML	A209580 001	Dec 18, 2017
AP	APOTEX	6MG/ML	A210448 001	May 07, 2019
AP	ARTHUR GRP	6MG/ML	A205106 001	Sep 21, 2018
AP	HOSPIRA INC	6MG/ML	A205672 001	Jul 31, 2018
AP	MEITHEAL	6MG/ML	A212127 001	Oct 23, 2020
AP	MYLAN INSTITUTIONAL	6MG/ML	A208536 001	Nov 20, 2017
AP	NEXUS PHARMS	6MG/ML	A207794 001	Jan 14, 2019
AP	PHARMASCIENCE INC	6MG/ML	A207050 001	Mar 24, 2017
AP	SHILPA	6MG/ML	A210931 001	Apr 18, 2019

BUSULFEX

AP	+ ! OTSUKA PHARM	6MG/ML	N020954 001	Feb 04, 1999
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TABLET; ORAL

MYLERAN

+ !	ASPEN GLOBAL	2MG	N009386 001	
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BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL

MENTAX

+ !	MYLAN	1%	N020524 001	Oct 18, 1996
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BUTOCONAZOLE NITRATE

CREAM; VAGINAL

GYNAZOLE-1

!	PADAGIS ISRAEL	2%	A200923 001	May 18, 2012
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BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATE

AP	HIKMA	2MG/ML	A075046 001	Aug 12, 1998	
AP	HIKMA FARMACEUTICA	1MG/ML	A078400 001	May 01, 2009	
AP		2MG/ML	A078400 002	May 01, 2009	
<u>BUTORPHANOL TARTRATE PRESERVATIVE FREE</u>					
AP	HIKMA	1MG/ML	A075045 001	Aug 12, 1998	
AP		2MG/ML	A075045 002	Aug 12, 1998	
AP	!	HOSPIRA	1MG/ML	A074626 001	Jan 23, 1997
AP	!		2MG/ML	A074626 002	Jan 23, 1997

SPRAY, METERED; NASAL

BUTORPHANOL TARTRATE

AB	APOTEX INC	1MG/SPRAY	A075499 001	Dec 04, 2002	
AB	HIKMA	1MG/SPRAY	A075824 001	Mar 12, 2002	
AB	!	RISING PHARMA	1MG/SPRAY	A075759 001	Aug 08, 2001

CABAZITAXEL

SOLUTION; INTRAVENOUS

CABAZITAXEL

+ !	ACCORD HLTHCARE	60MG/3ML (20MG/ML)	N207949 001	Dec 29, 2021
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JEVTANA KIT

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

TABLET; ORAL

CABERGOLINE

AB	!	INGENUS PHARMS LLC	0.5MG	A204735 001	Aug 01, 2018
AB		IVAX SUB TEVA PHARMS	0.5MG	A077750 001	Mar 07, 2007

CABOTEGRAVIR

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

APRETUDE

+ !	VIIV HLTHCARE	600MG/3ML (200MG/ML)	N215499 001	Dec 20, 2021
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CABOTEGRAVIR SODIUM

TABLET; ORAL

VOCABRIA

+ !	VIIV HLTHCARE	EQ 30MG BASE	N212887 001	Jan 21, 2021
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PRESCRIPTION DRUG PRODUCT LIST

CABOTEGRAVIR; RILPIVIRINE

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

CABENUVA KIT

+!	VIIV HLTHCARE	400MG/2ML (200MG/ML); 600MG/2ML (300MG/ML)	N212888 001	Jan 21, 2021
+!		600MG/3ML (200MG/ML); 900MG/3ML (300MG/ML)	N212888 002	Jan 21, 2021

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

CAFFEINE CITRATE

SOLUTION; INTRAVENOUS

CAFSCI

AP +! HIKMA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **N020793 001** Sep 21, 1999

CAFFEINE CITRATE

AP AM REGENT EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **A077906 001** May 15, 2007
AP EUGIA PHARMA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **A205013 001** Sep 22, 2015
AP EXELA PHARMA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **A077233 001** Sep 21, 2006
 SCIENCE
AP FRESENIUS KABI USA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **A077997 001** Jul 20, 2007
AP MICRO LABS EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **A207400 001** Dec 14, 2017
AP SAGENT PHARMS EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **A090827 001** Aug 29, 2012

SOLUTION; ORAL

CAFSCI

AA +! HIKMA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **N020793 002** Apr 12, 2000

CAFFEINE CITRATE

AA EXELA PHARMA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **A077304 001** Sep 21, 2006
AA FRESENIUS KABI USA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **A078002 001** Jan 31, 2008
AA MICRO LABS EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **A213202 001** Dec 16, 2019
AA SAGENT PHARMS EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **A091102 001** Aug 29, 2012
AA SUN PHARM EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **A090357 001** Sep 30, 2009

CAFFEINE; ERGOTAMINE TARTRATE

SUPPOSITORY; RECTAL

MIGERGOT

! COSETTE 100MG; 2MG A086557 001 Oct 04, 1983

TABLET; ORAL

ERGOTAMINE TARTRATE AND CAFFEINE

! MIKART 100MG; 1MG A040590 001 Sep 16, 2005

CALCIFEDIOL

CAPSULE, EXTENDED RELEASE; ORAL

RAYALDEE

+! EIRGEN 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 0.005% **A205772 001** Jun 09, 2015
AB TOLMAR 0.005% **A200935 001** May 30, 2012

DOVONEX

AB +! LEO PHARMA AS 0.005% **N020554 001** Jul 22, 1996

OINTMENT; TOPICAL

CALCIPOTRIENE

! GLENMARK PHARMS INC 0.005% A090633 001 Mar 24, 2010

SOLUTION; TOPICAL

CALCIPOTRIENE

AT AKORN 0.005% **A077579 001** Nov 19, 2009
AT COSETTE 0.005% **A078468 001** Mar 24, 2011
AT FOUGERA PHARMS 0.005% **A078305 001** May 06, 2008
AT NOVEL LABS INC 0.005% **A207163 001** Dec 26, 2017
AT ! TOLMAR 0.005% **A077029 001** Nov 20, 2009

PRESCRIPTION DRUG PRODUCT LIST

CALCITONIN SALMON

INJECTABLE; INJECTION

CALCITONIN-SALMON

AP	CUSTOPHARM INC	200 IU/ML	A212416 001	May 14, 2021
AP	PAR STERILE PRODUCTS	200 IU/ML	A209358 001	Nov 10, 2021

MIACALCIN

AP	+! MYLAN IRELAND LTD	200 IU/ML	N017808 002	Mar 29, 1991
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CALCITONIN-SALMON

AB	! APOTEX INC	200 IU/SPRAY	A076396 001	Nov 17, 2008
AB	PAR PHARM	200 IU/SPRAY	A076979 001	Jun 08, 2009

CALCITRIOL

CAPSULE; ORAL

CALCITRIOL

AB	AMNEAL PHARMS	0.25MCG	A203289 002	Jun 14, 2017
AB		0.5MCG	A203289 001	Jun 14, 2017
AB	BIONPHARMA INC	0.25MCG	A091174 001	May 24, 2013
AB		0.5MCG	A091174 002	May 24, 2013
AB	HIKMA	0.25MCG	A076917 001	Mar 27, 2006
AB	STRIDES PHARMA	0.25MCG	A091356 001	Dec 12, 2014
AB		0.5MCG	A091356 002	Dec 12, 2014
AB	SUN PHARM	0.25MCG	A204556 001	Feb 21, 2019
AB		0.5MCG	A204556 002	Feb 21, 2019
AB	TEVA	0.25MCG	A075765 001	Oct 12, 2001
AB		0.5MCG	A075765 002	Oct 12, 2001

ROCALTROL

AB	+ VALIDUS PHARMS	0.25MCG	N018044 001	
AB	+!	0.5MCG	N018044 002	

INJECTABLE; INJECTION

CALCITRIOL

AP	! AKORN	0.001MG/ML	A078066 001	Jan 29, 2008
AP	GLAND PHARMA LTD	0.001MG/ML	A211030 001	Feb 03, 2020

OINTMENT; TOPICAL

VECTICAL

	+! GALDERMA LABS LP	3MCG/GM	N022087 001	Jan 23, 2009
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SOLUTION; ORAL

CALCITRIOL

AA	ELYSIUM	1MCG/ML	A209798 001	Nov 21, 2018
AA	HIKMA	1MCG/ML	A076242 001	Jul 18, 2003

ROCALTROL

AA	+! VALIDUS PHARMS	1MCG/ML	N021068 001	Nov 20, 1998
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CALCIUM ACETATE

CAPSULE; ORAL

CALCIUM ACETATE

AB	CHARTWELL RX	667MG	A091312 001	Jun 01, 2012
AB	HERITAGE PHARMS INC	667MG	A202315 001	Jun 29, 2015
AB	HIKMA	667MG	A077728 001	Feb 26, 2008
AB	INVAGEN PHARMS	667MG	A203135 001	Feb 07, 2013
AB	LUPIN LTD	667MG	A202127 001	Jul 09, 2015
AB	NOSTRUM LABS INC	667MG	A203179 001	Oct 26, 2015
AB	SUVEN PHARMS	667MG	A211038 001	Feb 21, 2020

PHOSLO GELCAPS

AB	+! FRESENIUS MEDCL	667MG	N021160 003	Apr 02, 2001
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SOLUTION; ORAL

PHOSLYRA

	+! FRESENIUS MEDCL	667MG/5ML	N022581 001	Apr 18, 2011
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TABLET; ORAL

CALCIUM ACETATE

AB	! CHARTWELL MOLECULAR	667MG	A202420 001	Feb 05, 2013
AB	HERITAGE PHARMS INC	667MG	A202885 001	Jan 22, 2015
AB	PADAGIS US	667MG	A091561 001	Apr 13, 2011

CALCIUM CHLORIDE

INJECTABLE; INJECTION

CALCIUM CHLORIDE 10%

AP	AM REGENT	100MG/ML	A209088 001	Jul 27, 2017
AP	INTL MEDICATION SYS	100MG/ML	A203477 001	May 09, 2018
AP	MEDEFIL INC	100MG/ML	A211553 001	May 01, 2019

CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER

AP	+! HOSPIRA	100MG/ML	N021117 001	Jan 28, 2000
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PRESCRIPTION DRUG PRODUCT LIST

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/M L; 0.38MG/ML; 2.1MG/ML; 7.14MG/ML; 0.42MG/M L	N018469 001
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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; 3.0 5GM/1000ML; 0.314GM/1000ML; 2.21GM/1000ML ; 7.07GM/1000ML (5000ML)	N021703 011	Oct 10, 2008
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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
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PRISMASOL BGK 2/0 IN PLASTIC CONTAINER

+!	BAXTER HLTHCARE CORP	N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.0 3GM/1000ML; 0.157GM/1000ML; 3.09GM/1000ML ; 6.46GM/1000ML (5000ML)	N021703 002	Oct 25, 2006
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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/100 0ML; 6.46GM/1000ML (5000ML)	N021703 003	Oct 25, 2006
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PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER

+!	BAXTER HLTHCARE CORP	N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.4 4GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML ; 6.46GM/1000ML (5000ML)	N021703 015	Oct 10, 2008
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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/100 0ML; 6.46GM/1000ML (5000ML)	N021703 004	Oct 25, 2006
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PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER

+!	BAXTER HLTHCARE CORP	N/A/1000ML; N/A/1000ML; 5.4GM/1000ML; 2.44 GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6.46 GM/1000ML (5000ML)	N021703 014	Oct 10, 2008
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CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DELFLX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT	+!	FRESENIUS MEDCL	25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 5 67MG/100ML; 392MG/100ML	N018883 001	Nov 30, 1984
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DELFLX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	+!	FRESENIUS MEDCL	25.7MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N018883 004	Nov 30, 1984
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DELFLX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT	+!	FRESENIUS MEDCL	18.4MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N020171 001	Aug 19, 1992
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DELFLX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT	+!	FRESENIUS MEDCL	25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 5 67MG/100ML; 392MG/100ML	N018883 002	Nov 30, 1984
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DELFLX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	+!	FRESENIUS MEDCL	25.7MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N018883 005	Nov 30, 1984
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DELFLX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT	+!	FRESENIUS MEDCL	18.4MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N020171 002	Aug 19, 1992
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DELFLX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT	+!	FRESENIUS MEDCL	25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 5 567MG/100ML; 392MG/100ML	N018883 003	Nov 30, 1984
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DELFLX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	+!	FRESENIUS MEDCL	25.7MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5 538MG/100ML; 448MG/100ML	N018883 006	Nov 30, 1984
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DELFLX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT	+!	FRESENIUS MEDCL	18.4MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5 538MG/100ML; 448MG/100ML	N020171 003	Aug 19, 1992
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DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	18.3MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N020183 001	Dec 04, 1992
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DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	18.3MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N020183 002	Dec 04, 1992
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DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	18.3MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5 538MG/100ML; 448MG/100ML	N020183 004	Dec 04, 1992
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PRESCRIPTION DRUG PRODUCT LIST

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DIANEAL LOW CALCIUM W/DEXTROSE 1.5% IN PLASTIC CONTAINER

<u>AT</u>	+	BAXTER HLTHCARE	<u>18.3MG/100ML;1.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML</u>	<u>N017512 012</u>	Jan 10, 1989
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DIANEAL LOW CALCIUM W/DEXTROSE 2.5% IN PLASTIC CONTAINER

<u>AT</u>	+	BAXTER HLTHCARE	<u>18.3MG/100ML;2.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML</u>	<u>N017512 013</u>	Jul 11, 1990
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DIANEAL LOW CALCIUM W/DEXTROSE 3.5% IN PLASTIC CONTAINER

<u>AT</u>	+	BAXTER HLTHCARE	<u>18.3MG/100ML;3.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML</u>	<u>N017512 014</u>	Jul 11, 1990
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DIANEAL LOW CALCIUM W/DEXTROSE 4.25% IN PLASTIC CONTAINER

<u>AT</u>	+	BAXTER HLTHCARE	<u>18.3MG/100ML;4.25GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML</u>	<u>N017512 015</u>	Jul 11, 1990
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DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

<u>AT</u>	+	BAXTER HLTHCARE	<u>25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML</u>	<u>N017512 004</u>	
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<u>AT</u>	+		<u>25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML</u>	<u>N020163 001</u>	Dec 04, 1992
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DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

<u>AT</u>	+	BAXTER HLTHCARE	<u>25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML</u>	<u>N017512 005</u>	
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<u>AT</u>	+		<u>25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML</u>	<u>N020163 002</u>	Dec 04, 1992
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DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

<u>AT</u>	+	BAXTER HLTHCARE	<u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML</u>	<u>N017512 006</u>	
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<u>AT</u>	+		<u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML</u>	<u>N020163 003</u>	Dec 04, 1992
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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
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CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

<u>AP</u>	+	ICU MEDICAL INC	<u>20MG/100ML;5GM/100ML;30MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N017608 001</u>	
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DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER

<u>AP</u>		B BRAUN	<u>20MG/100ML;5GM/100ML;30MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N019634 003</u>	Feb 24, 1988
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LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>		BAXTER HLTHCARE	<u>20MG/100ML;5GM/100ML;30MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N016679 001</u>	
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POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

<u>AP</u>		BAXTER HLTHCARE	<u>20MG/100ML;5GM/100ML;254MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N019367 006</u>	Apr 05, 1985
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POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

<u>AP</u>		BAXTER HLTHCARE	<u>20MG/100ML;5GM/100ML;179MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N019367 004</u>	Apr 05, 1985
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<u>AP</u>			<u>20MG/100ML;5GM/100ML;328MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N019367 005</u>	Apr 05, 1985
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<u>AP</u>		FRESENIUS KABI USA	<u>20MG/100ML;5GM/100ML;179MG/100ML;600MG/100ML;310MG/100ML</u>	<u>A211428 001</u>	Nov 09, 2021
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<u>AP</u>	+	ICU MEDICAL INC	<u>20MG/100ML;5GM/100ML;179MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N019685 002</u>	Oct 17, 1988
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POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

<u>AP</u>		BAXTER HLTHCARE	<u>20MG/100ML;5GM/100ML;254MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N019367 007</u>	Apr 05, 1985
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POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

<u>AP</u>		BAXTER HLTHCARE	<u>20MG/100ML;5GM/100ML;328MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N019367 008</u>	Apr 05, 1985
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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
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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
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			20MG/100ML;5GM/100ML;179MG/100ML;600MG/100ML;310MG/100ML	N019367 003	Apr 05, 1985
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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
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PRESCRIPTION DRUG PRODUCT LIST

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
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CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

SOLUTION; IRRIGATION

BALANCED SALT

AT		AKORN	<u>0.48MG/ML; 0.3MG/ML; 0.75MG/ML; 3.9MG/ML; 6.4MG/ML; 1.7MG/ML</u>	A075503 001	Sep 27, 2006
AT		B BRAUN	<u>0.48MG/ML; 0.3MG/ML; 0.75MG/ML; 3.9MG/ML; 6.4MG/ML; 1.7MG/ML</u>	A091387 001	Feb 03, 2010

BSS

AT	+	ALCON	<u>0.48MG/ML; 0.3MG/ML; 0.75MG/ML; 3.9MG/ML; 6.4MG/ML; 1.7MG/ML</u>	N020742 001	Dec 10, 1997
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CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

INJECTABLE; INJECTION

PHOXILLUM B22K 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
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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
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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>	A075323 001	Apr 21, 2000
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PLEGISOL IN PLASTIC CONTAINER

AT	+	HOSPIRA	<u>17.6MG/100ML; 325.3MG/100ML; 119.3MG/100ML; 643MG/100ML</u>	N018608 001	Feb 26, 1982
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CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

RINGER'S IN PLASTIC CONTAINER

AP		B BRAUN	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	N020002 001	Apr 17, 1992
AP		BAXTER HLTHCARE	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	N016693 001	
AP		ICU MEDICAL INC	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	N018251 001	

SOLUTION; IRRIGATION

RINGER'S IN PLASTIC CONTAINER

AT		B BRAUN	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	N018156 001	
AT		BAXTER HLTHCARE	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	N018495 001	Feb 19, 1982
AT		ICU MEDICAL INC	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	N017635 001	

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>	N019632 001	Feb 29, 1988
AP	+	BAXTER HLTHCARE	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	N016682 001	
AP		FRESENIUS KABI USA	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	A209338 001	Jan 28, 2019
AP		ICU MEDICAL INC	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	N017641 001	

SOLUTION; IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER

AT	+	B BRAUN	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	N018681 001	Dec 27, 1982
AT		BAXTER HLTHCARE	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	N018494 001	Feb 19, 1982
AT	+		<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	N018921 001	Apr 03, 1984
AT	+	ICU MEDICAL INC	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	N019416 001	Jan 17, 1986

PRESCRIPTION DRUG PRODUCT LIST

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

+!	FRESENIUS KABI USA	1GM/50ML (20MG/ML)	N208418	004	Jun 17, 2021
+!		2GM/100ML (20MG/ML)	N208418	005	Jun 17, 2021
+!	HQ SPCLT PHARMA	1GM/50ML (20MG/ML)	N210906	001	Oct 29, 2018
+!		1GM/100ML (10MG/ML)	N210906	003	Jun 04, 2021
+!		2GM/100ML (20MG/ML)	N210906	002	Oct 29, 2018

CALCIUM OXYBATE; MAGNESIUM OXYBATE; POTASSIUM OXYBATE; SODIUM OXYBATE

SOLUTION; ORAL

XYWAV

+!	JAZZ	0.234GM/ML; 0.096GM/ML; 0.13GM/ML; 0.04GM/ML	N212690	001	Jul 21, 2020
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CANAGLIFLOZIN

TABLET; ORAL

INVOKANA

+	JANSSEN PHARMS	100MG	N204042	001	Mar 29, 2013
+!		300MG	N204042	002	Mar 29, 2013

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	<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	<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	<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	<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

PRESCRIPTION DRUG PRODUCT LIST

CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>		<u>32MG;12.5MG</u>	<u>A204100 002</u>	Feb 27, 2015
<u>AB</u>		<u>32MG;25MG</u>	<u>A204100 003</u>	Feb 27, 2015
<u>AB</u>	MYLAN	<u>16MG;12.5MG</u>	<u>A090704 001</u>	Dec 04, 2012
<u>AB</u>		<u>32MG;12.5MG</u>	<u>A090704 002</u>	Dec 04, 2012
<u>AB</u>		<u>32MG;25MG</u>	<u>A090704 003</u>	Dec 04, 2012
<u>AB</u>	PRINSTON INC	<u>16MG;12.5MG</u>	<u>A207455 001</u>	Apr 11, 2018
<u>AB</u>		<u>32MG;12.5MG</u>	<u>A207455 002</u>	Apr 11, 2018
<u>AB</u>		<u>32MG;25MG</u>	<u>A207455 003</u>	Apr 11, 2018
<u>AB</u>	ZYDUS PHARMS	<u>16MG;12.5MG</u>	<u>A203466 001</u>	Nov 27, 2017
<u>AB</u>		<u>32MG;12.5MG</u>	<u>A203466 002</u>	Nov 27, 2017
<u>AB</u>		<u>32MG;25MG</u>	<u>A203466 003</u>	Nov 27, 2017

CANGRELOR

POWDER; INTRAVENOUS

KENGREAL

+! CHIESI 50MG/VIAL N204958 001 Jun 22, 2015

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>	DR REDDYS LABS LTD	<u>150MG</u>	<u>A204345 001</u>	Dec 04, 2020
<u>AB</u>		<u>500MG</u>	<u>A204345 002</u>	Dec 04, 2020
<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>	HIKMA	<u>150MG</u>	<u>A200483 001</u>	Jul 14, 2016
<u>AB</u>		<u>500MG</u>	<u>A200483 002</u>	Jul 14, 2016
<u>AB</u>	MSN	<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>	RELIANCE LIFE	<u>150MG</u>	<u>A211724 001</u>	Apr 27, 2020
<u>AB</u>		<u>500MG</u>	<u>A211724 002</u>	Apr 27, 2020
<u>AB</u>	RISING PHARMA	<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	<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>	SUN PHARM	<u>150MG</u>	<u>A204668 001</u>	Jun 21, 2019
<u>AB</u>		<u>500MG</u>	<u>A204668 002</u>	Jun 21, 2019
<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

XELODA

<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

CAPMATINIB HYDROCHLORIDE

TABLET; ORAL

TABRECTA

+ NOVARTIS PHARM EQ 150MG BASE N213591 001 May 06, 2020

+! EQ 200MG BASE N213591 002 May 06, 2020

CAPSAICIN

PATCH; TOPICAL

QUTENZA

+! AVERITAS 8% N022395 001 Nov 16, 2009

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

<u>AB</u>	AJANTA PHARMA LTD	<u>12.5MG</u>	<u>A212809 001</u>	Dec 13, 2019
<u>AB</u>		<u>25MG</u>	<u>A212809 002</u>	Dec 13, 2019
<u>AB</u>		<u>50MG</u>	<u>A212809 003</u>	Dec 13, 2019
<u>AB</u>		<u>100MG</u>	<u>A212809 004</u>	Dec 13, 2019
<u>AB</u>	ANNORA PHARMA	<u>12.5MG</u>	<u>A074737 001</u>	Oct 28, 1998

PRESCRIPTION DRUG PRODUCT LIST

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

<u>AB</u>		<u>25MG</u>	<u>A074737 002</u>	Oct 28, 1998
<u>AB</u>		<u>50MG</u>	<u>A074737 003</u>	Oct 28, 1998
<u>AB</u>		<u>100MG</u>	<u>A074737 004</u>	Oct 28, 1998
<u>AB</u>	BOSCOGEN	<u>12.5MG</u>	<u>A074677 004</u>	May 30, 1997
<u>AB</u>		<u>25MG</u>	<u>A074677 002</u>	May 30, 1997
<u>AB</u>		<u>50MG</u>	<u>A074677 001</u>	May 30, 1997
<u>AB</u>		<u>100MG</u>	<u>A074677 003</u>	May 30, 1997
<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>	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>	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

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 001</u>	Nov 25, 2011
<u>AB</u>		<u>200MG</u>	<u>A078986 002</u>	Nov 25, 2011
<u>AB</u>		<u>300MG</u>	<u>A078986 003</u>	Nov 25, 2011
<u>AB</u>	NOSTRUM LABS INC	<u>100MG</u>	<u>A076697 001</u>	May 20, 2011
<u>AB</u>		<u>200MG</u>	<u>A076697 002</u>	May 20, 2011
<u>AB</u>		<u>300MG</u>	<u>A076697 003</u>	May 20, 2011
<u>AB</u>	TARO	<u>100MG</u>	<u>A201106 001</u>	Jun 21, 2013
<u>AB</u>		<u>200MG</u>	<u>A201106 002</u>	Jun 21, 2013
<u>AB</u>		<u>300MG</u>	<u>A201106 003</u>	Jun 21, 2013
<u>AB</u>	TEVA PHARMS	<u>100MG</u>	<u>A078592 001</u>	Sep 20, 2012
<u>AB</u>		<u>200MG</u>	<u>A078592 002</u>	Sep 20, 2012
<u>AB</u>		<u>300MG</u>	<u>A078592 003</u>	Sep 20, 2012

CARBATROL

<u>AB</u>	+ TAKEDA PHARMS USA	<u>100MG</u>	<u>N020712 003</u>	Sep 30, 1997
<u>AB</u>	+	<u>200MG</u>	<u>N020712 001</u>	Sep 30, 1997
<u>AB</u>	+!	<u>300MG</u>	<u>N020712 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 001</u>	Jun 05, 2002
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TEGRETOL

<u>AB</u>	+! NOVARTIS	<u>100MG/5ML</u>	<u>N018927 001</u>	Dec 18, 1987
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TERIL

<u>AB</u>	TARO	<u>100MG/5ML</u>	<u>A076729 001</u>	Sep 20, 2004
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TABLET; ORAL

CARBAMAZEPINE

<u>AB</u>	APOTEX INC	<u>200MG</u>	<u>A075948 001</u>	Feb 27, 2002
<u>AB</u>	TARO	<u>200MG</u>	<u>A074649 001</u>	Oct 03, 1996
<u>AB</u>	TORRENT PHARMS	<u>100MG</u>	<u>A077272 001</u>	Dec 07, 2005
<u>AB</u>		<u>200MG</u>	<u>A077272 002</u>	Dec 07, 2005
<u>AB</u>	UMEDICA LABS PVT LTD	<u>100MG</u>	<u>A207798 001</u>	Apr 15, 2020
<u>AB</u>		<u>200MG</u>	<u>A207798 002</u>	Apr 15, 2020
<u>AB</u>	VGYAAN	<u>200MG</u>	<u>A214328 001</u>	Aug 16, 2021

EPITOL

<u>AB</u>	TEVA	<u>200MG</u>	<u>A070541 001</u>	Sep 17, 1986
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TEGRETOL

<u>AB</u>	+! NOVARTIS	<u>200MG</u>	<u>N016608 001</u>	
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PRESCRIPTION DRUG PRODUCT LIST

CARBAMAZEPINE

TABLET; ORAL

CARBAMAZEPINE

TORRENT PHARMS	300MG	A077272 003	Dec 07, 2005
	400MG	A077272 004	Dec 07, 2005

TABLET, CHEWABLE; ORAL

CARBAMAZEPINE

AB	!	TARO PHARM INDS	100MG	A075687 001	Oct 24, 2000
AB		TORRENT PHARMS	100MG	A075712 001	Jul 05, 2001

EPITOL

AB		TEVA	100MG	A073524 001	Jul 29, 1992
		CARBAMAZEPINE			
	!	TARO PHARM INDS	200MG	A075687 002	Jul 29, 2002

TABLET, EXTENDED RELEASE; ORAL

CARBAMAZEPINE

AB		ANBISON LAB	100MG	A212948 001	Sep 30, 2021
AB			200MG	A212948 002	Sep 30, 2021
AB			400MG	A212948 003	Sep 30, 2021
AB		CSPC OUYI	100MG	A213311 001	Apr 13, 2021
AB			200MG	A213311 002	Apr 13, 2021
AB			400MG	A213311 003	Apr 13, 2021
AB		TARO	100MG	A078115 001	Mar 31, 2009
AB			200MG	A078115 002	Mar 31, 2009
AB			400MG	A078115 003	Mar 31, 2009
AB		UNIQUE PHARM	100MG	A211623 001	Apr 24, 2020
AB			200MG	A211623 002	Apr 24, 2020
AB			400MG	A211623 003	Apr 24, 2020
AB		ZYDUS PHARMS	100MG	A205571 001	Feb 07, 2019
AB			200MG	A205571 002	Feb 07, 2019
AB			400MG	A205571 003	Feb 07, 2019
		<u>TEGRETOL-XR</u>			
AB	+	NOVARTIS	100MG	N020234 001	Mar 25, 1996
AB	+		200MG	N020234 002	Mar 25, 1996
AB	+		400MG	N020234 003	Mar 25, 1996

CARBIDOPA

TABLET; ORAL

CARBIDOPA

AB		ALVOGEN	25MG	A204291 001	Jan 08, 2016
AB		ANI PHARMS	25MG	A203261 001	Mar 10, 2014
AB		AUROBINDO PHARMA LTD	25MG	A211055 001	Oct 21, 2019
AB		EDENBRIDGE PHARMS	25MG	A205304 001	Feb 17, 2016
AB		NOVEL LABS INC	25MG	A204763 001	Oct 20, 2017
AB		ZYDUS PHARMS	25MG	A209910 001	May 07, 2018

LODOSYN

AB	+	ATON	25MG	N017830 001	
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CARBIDOPA; ENTACAPONE; LEVODOPA

TABLET; ORAL

CARBIDOPA, LEVODOPA AND ENTACAPONE

AB		SUN PHARM	25MG; 200MG; 100MG	A079085 001	May 10, 2012
AB			37.5MG; 200MG; 150MG	A079085 002	May 10, 2012
		<u>STALEVO 100</u>			
AB	+	ORION PHARMA	25MG; 200MG; 100MG	N021485 002	Jun 11, 2003
		<u>STALEVO 150</u>			
AB	+	ORION PHARMA	37.5MG; 200MG; 150MG	N021485 003	Jun 11, 2003
		STALEVO 125			
	+	ORION PHARMA	31.25MG; 200MG; 125MG	N021485 006	Aug 29, 2008
		STALEVO 200			
	+	ORION PHARMA	50MG; 200MG; 200MG	N021485 004	Aug 02, 2007
		STALEVO 50			
	+	ORION PHARMA	12.5MG; 200MG; 50MG	N021485 001	Jun 11, 2003
		STALEVO 75			
	+	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

PRESCRIPTION DRUG PRODUCT LIST

CARBIDOPA; LEVODOPA

SUSPENSION; ENTERAL

DUOPA

+! ABBVIE INC

4.63MG/ML; 20MG/ML

N203952 001 Jan 09, 2015

TABLET; ORAL

CARBIDOPA AND LEVODOPAAB ACTAVIS ELIZABETH10MG; 100MGA074260 001 Sep 03, 1993AB 25MG; 100MG A074260 002 Sep 03, 1993AB ! 25MG; 250MG A074260 003 Sep 03, 1993AB APOTEX INC 10MG; 100MG A077120 001 Jun 02, 2008AB 25MG; 100MG A077120 002 Jun 02, 2008AB 25MG; 250MG A077120 003 Jun 02, 2008AB MAYNE PHARMA 10MG; 100MG A073618 001 Aug 28, 1992AB 25MG; 100MG A073589 001 Aug 28, 1992AB 25MG; 250MG A073607 001 Aug 28, 1992AB MYLAN 10MG; 100MG A090324 001 Sep 28, 2009AB 25MG; 100MG A090324 002 Sep 28, 2009AB 25MG; 250MG A090324 003 Sep 28, 2009AB SCIEGEN PHARMS INC 10MG; 100MG A214092 001 May 07, 2021AB 25MG; 100MG A214092 002 May 07, 2021AB 25MG; 250MG A214092 003 May 07, 2021AB SUN PHARM INDS 10MG; 100MG A078536 001 Oct 28, 2008AB 25MG; 100MG A078536 002 Oct 28, 2008AB 25MG; 250MG A078536 003 Oct 28, 2008SINEMETAB + ORGANON 10MG; 100MG N017555 001AB + 25MG; 100MG N017555 003AB + 25MG; 250MG N017555 002

DHIVY

+! AVION PHARMS

25MG; 100MG

N214869 001 Nov 12, 2021

TABLET, EXTENDED RELEASE; ORAL

CARBIDOPA AND LEVODOPAAB ACCORD HLTHCARE 25MG; 100MG A202323 001 Feb 08, 2013AB 50MG; 200MG A202323 002 Feb 08, 2013AB ALEMBIC PHARMS LTD 25MG; 100MG A210341 001 Jun 05, 2019AB 50MG; 200MG A210341 002 Jun 05, 2019AB APOTEX 25MG; 100MG A076212 001 Jun 16, 2004AB 50MG; 200MG A076212 002 Jun 16, 2004AB IMPAX LABS 25MG; 100MG A076521 001 May 14, 2004AB 50MG; 200MG A076521 002 May 14, 2004AB MYLAN 25MG; 100MG A075091 002 Apr 21, 2000AB 50MG; 200MG A075091 001 Sep 30, 1999AB SCIEGEN PHARMS INC 25MG; 100MG A214091 001 Oct 05, 2021AB 50MG; 200MG A214091 002 Oct 05, 2021AB SUN PHARM INDS 25MG; 100MG A077828 001 Aug 23, 2007AB ! 50MG; 200MG A077828 002 Aug 23, 2007

TABLET, ORALLY DISINTEGRATING; ORAL

CARBIDOPA AND LEVODOPAAB SUN PHARM 10MG; 100MG A078690 001 Jul 31, 2009AB 25MG; 100MG A078690 002 Jul 31, 2009

! 25MG; 250MG A078690 003 Jul 31, 2009

CARBINOXAMINE MALEATE

SOLUTION; ORAL

CARBINOXAMINE MALEATE

! MIKART

4MG/5ML

A040458 001 Apr 25, 2003

SUSPENSION, EXTENDED RELEASE; ORAL

KARBINAL ER

+! AYTU

4MG/5ML

N022556 001 Mar 28, 2013

TABLET; ORAL

CARBINOXAMINE MALEATEAA INVAGEN PHARMS 4MG A090435 001 Apr 15, 2010AA ! MIKART 4MG A040442 001 Mar 19, 2003AA MISSION PHARMACAL 4MG A090756 001 May 27, 2011

! MIKART 6MG A207484 001 May 31, 2016

CARBOPLATIN

INJECTABLE; INTRAVENOUS

CARBOPLATINAP ACCORD HLTHCARE 50MG/5ML (10MG/ML) A206775 001 Feb 09, 2017AP 150MG/15ML (10MG/ML) A206775 002 Feb 09, 2017AP 450MG/45ML (10MG/ML) A206775 003 Feb 09, 2017AP 600MG/60ML (10MG/ML) A206775 004 Feb 09, 2017

PRESCRIPTION DRUG PRODUCT LIST

CARBOPLATIN

INJECTABLE; INTRAVENOUS

CARBOPLATIN

<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>		<u>600MG/60ML (10MG/ML)</u>	<u>A205487 004</u>	Aug 03, 2020
<u>AP</u>	FRESENIUS KABI USA	<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>	HIKMA	<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
<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
<u>AP</u>	NOVAST LABS	<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>	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>	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 PHARM	<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
	! ACCORD HLTHCARE	<u>1GM/100ML (10MG/ML)</u>	<u>A206775 005</u>	Apr 06, 2020

CARBOPROST TROMETHAMINE

INJECTABLE; INJECTION

CARBOPROST TROMETHAMINE

<u>AP</u>	DR REDDYS LABS LTD	<u>EQ 0.25MG BASE/ML</u>	<u>A211941 001</u>	Jul 02, 2019
<u>AP</u>	SUNNY	<u>EQ 0.25MG BASE/ML</u>	<u>A213118 001</u>	Mar 25, 2021
	<u>HEMABATE</u>			
<u>AP</u>	+! PFIZER	<u>EQ 0.25MG BASE/ML</u>	<u>N017989 001</u>	

CARFILZOMIB

POWDER; INTRAVENOUS

CARFILZOMIB

<u>AP</u>	DR REDDYS	<u>60MG/VIAL</u>	<u>A209422 001</u>	Sep 09, 2019
	<u>KYPROLIS</u>			
<u>AP</u>	+! ONYX THERAP	<u>60MG/VIAL</u>	<u>N202714 001</u>	Jul 20, 2012
	+	10MG/VIAL	N202714 003	Jun 07, 2018
	+	30MG/VIAL	N202714 002	Jun 03, 2016

CARGLUMIC ACID

TABLET, FOR SUSPENSION; ORAL

CARBAGLU

<u>AB</u>	+! RECORDATI RARE	<u>200MG</u>	<u>N022562 001</u>	Mar 18, 2010
	<u>CARGLUMIC ACID</u>			
<u>AB</u>	NOVITIUM PHARMA	<u>200MG</u>	<u>A213729 001</u>	Oct 13, 2021

PRESCRIPTION DRUG PRODUCT LIST

CARIPRAZINE HYDROCHLORIDE

CAPSULE; ORAL

VRAYLAR

+	ALLERGAN	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>	FOSUN PHARMA	<u>350MG</u>	<u>A081025 001</u>	Apr 13, 1989
<u>AA</u>	MLV	<u>350MG</u>	<u>A040245 001</u>	Sep 08, 1997
<u>AA</u>		<u>350MG</u>	<u>A211789 001</u>	Oct 20, 2021
<u>AA</u>	NATCO	<u>350MG</u>	<u>A090988 001</u>	Oct 28, 2014
<u>AA</u>	NOSTRUM LABS INC	<u>350MG</u>	<u>A207237 002</u>	Sep 21, 2020
<u>AA</u>	NOVAST LABS	<u>350MG</u>	<u>A040823 001</u>	Oct 22, 2008
<u>AA</u>	ORIENT PHARMA CO LTD	<u>350MG</u>	<u>A205085 001</u>	Oct 28, 2014
<u>AA</u>	OXFORD PHARMS	<u>350MG</u>	<u>A040188 001</u>	Mar 07, 1997
<u>AA</u>	SCIEGEN PHARMS INC	<u>350MG</u>	<u>A203374 001</u>	Jan 27, 2014
<u>AA</u>	WATSON LABS	<u>350MG</u>	<u>A087499 001</u>	Apr 20, 1982
<u>AA</u>	WILSHIRE PHARMS INC	<u>350MG</u>	<u>A205126 002</u>	Jul 08, 2015

SOMA

<u>AA</u>	+	MYLAN SPECIALITY LP	<u>350MG</u>	<u>N011792 001</u>
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CARISOPRODOL

<u>AB</u>	AUROBINDO PHARMA	<u>250MG</u>	<u>A040792 002</u>	Nov 08, 2016
<u>AB</u>	NOSTRUM LABS INC	<u>250MG</u>	<u>A207237 001</u>	May 11, 2017
<u>AB</u>	WILSHIRE PHARMS INC	<u>250MG</u>	<u>A205126 001</u>	Jul 08, 2015

SOMA

<u>AB</u>	+	!	MYLAN SPECIALITY LP	<u>250MG</u>	<u>N011792 004</u>	Sep 13, 2007
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CARMUSTINE

IMPLANT; INTRACRANIAL

GLIADEL

+	ARBOR PHARMS LLC	7.7MG	N020637 001	Sep 23, 1996
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INJECTABLE; INJECTION

BICNU

<u>AP</u>	+	!	AVET LIFESCIENCES	<u>100MG/VIAL</u>	<u>N017422 001</u>
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CARMUSTINE

<u>AP</u>	AMNEAL	<u>100MG/VIAL</u>	<u>A211229 001</u>	Oct 16, 2018
<u>AP</u>	DR REDDYS	<u>100MG/VIAL</u>	<u>A213207 001</u>	Oct 22, 2020
<u>AP</u>	JIANGSU PHARMS	<u>100MG/VIAL</u>	<u>A211202 001</u>	Mar 12, 2021
<u>AP</u>	MEITHEAL	<u>100MG/VIAL</u>	<u>A213460 001</u>	Aug 02, 2021
<u>AP</u>	NAVINTA LLC	<u>100MG/VIAL</u>	<u>A210179 001</u>	Sep 11, 2018
<u>AP</u>	STI PHARMA LLC	<u>100MG/VIAL</u>	<u>A209278 001</u>	Apr 02, 2019

CARTEOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CARTEOLOL HYDROCHLORIDE

!	SANDOZ INC	1%	A075476 001	Jan 03, 2000
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CARVEDILOL

TABLET; ORAL

CARVEDILOL

<u>AB</u>	AUROBINDO PHARMA	<u>3.125MG</u>	<u>A078332 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078332 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078332 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078332 004</u>	Sep 05, 2007
<u>AB</u>	BEXIMCO USA	<u>3.125MG</u>	<u>A078384 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078384 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078384 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078384 004</u>	Sep 05, 2007
<u>AB</u>	CHARTWELL MOLECULAR	<u>3.125MG</u>	<u>A077474 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077474 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077474 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077474 004</u>	Sep 05, 2007
<u>AB</u>	DR REDDYS LABS LTD	<u>3.125MG</u>	<u>A076649 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A076649 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A076649 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A076649 004</u>	Sep 05, 2007
<u>AB</u>	GLENMARK GENERICS	<u>3.125MG</u>	<u>A078251 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078251 002</u>	Sep 05, 2007

PRESCRIPTION DRUG PRODUCT LIST

CARVEDILOL

TABLET; ORAL

CARVEDILOL

<u>AB</u>		<u>12.5MG</u>	<u>A078251 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078251 004</u>	Sep 05, 2007
<u>AB</u>	LUPIN	<u>3.125MG</u>	<u>A078217 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078217 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078217 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078217 004</u>	Sep 05, 2007
<u>AB</u>	MYLAN	<u>3.125MG</u>	<u>A077316 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077316 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077316 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077316 004</u>	Sep 05, 2007
<u>AB</u>	RUBICON	<u>3.125MG</u>	<u>A078165 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078165 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078165 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078165 004</u>	Sep 05, 2007
<u>AB</u>	SANDOZ	<u>3.125MG</u>	<u>A078227 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078227 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078227 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078227 004</u>	Sep 05, 2007
<u>AB</u>	SUN PHARM INDS LTD	<u>3.125MG</u>	<u>A076989 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A076989 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A076989 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A076989 004</u>	Sep 05, 2007
<u>AB</u>	TARO	<u>3.125MG</u>	<u>A077780 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077780 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077780 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077780 004</u>	Sep 05, 2007
<u>AB</u>	TEVA	<u>3.125MG</u>	<u>A076373 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A076373 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A076373 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A076373 004</u>	Sep 05, 2007
<u>AB</u>	ZYDUS PHARMS USA INC	<u>3.125MG</u>	<u>A077614 004</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077614 001</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077614 002</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077614 003</u>	Sep 05, 2007
<u>COREG</u>				
<u>AB</u>	+ WOODWARD	<u>3.125MG</u>	<u>N020297 004</u>	May 29, 1997
<u>AB</u>	+	<u>6.25MG</u>	<u>N020297 003</u>	Sep 14, 1995
<u>AB</u>	+!	<u>12.5MG</u>	<u>N020297 002</u>	Sep 14, 1995
<u>AB</u>	+	<u>25MG</u>	<u>N020297 001</u>	Sep 14, 1995

CARVEDILOL PHOSPHATE

CAPSULE, EXTENDED RELEASE; ORAL

CARVEDILOL PHOSPHATE

<u>AB</u>	IMPAX LABS INC	<u>10MG</u>	<u>A204717 001</u>	May 07, 2018
<u>AB</u>		<u>20MG</u>	<u>A204717 002</u>	May 07, 2018
<u>AB</u>		<u>40MG</u>	<u>A204717 003</u>	May 07, 2018
<u>AB</u>		<u>80MG</u>	<u>A204717 004</u>	May 07, 2018
<u>AB</u>	SUN PHARM INDUSTRIES	<u>10MG</u>	<u>A090132 001</u>	Oct 25, 2017
<u>AB</u>		<u>20MG</u>	<u>A090132 002</u>	Oct 25, 2017
<u>AB</u>		<u>40MG</u>	<u>A090132 003</u>	Oct 25, 2017
<u>AB</u>		<u>80MG</u>	<u>A090132 004</u>	Oct 25, 2017
<u>COREG CR</u>				
<u>AB</u>	+ WOODWARD	<u>10MG</u>	<u>N022012 001</u>	Oct 20, 2006
<u>AB</u>	+	<u>20MG</u>	<u>N022012 002</u>	Oct 20, 2006
<u>AB</u>	+!	<u>40MG</u>	<u>N022012 003</u>	Oct 20, 2006
<u>AB</u>	+	<u>80MG</u>	<u>N022012 004</u>	Oct 20, 2006

CASIMERSEN

SOLUTION; INTRAVENOUS

AMONDYS 45

+! SAREPTA THERAPS INC 100MG/2ML (50MG/ML)

N213026 001 Feb 25, 2021

CASPOFUNGIN ACETATE

POWDER; INTRAVENOUS

CANCIDAS

<u>AP</u>	+! MERCK	<u>50MG/VIAL</u>	<u>N021227 001</u>	Jan 26, 2001
<u>AP</u>	+!	<u>70MG/VIAL</u>	<u>N021227 002</u>	Jan 26, 2001
<u>CASPOFUNGIN ACETATE</u>				
<u>AP</u>	FRESENIUS KABI USA	<u>50MG/VIAL</u>	<u>N206110 001</u>	Dec 30, 2016

PRESCRIPTION DRUG PRODUCT LIST

CASPOFUNGIN ACETATE

POWDER; INTRAVENOUS

CASPOFUNGIN ACETATE

<u>AP</u>		<u>70MG/VIAL</u>	<u>N206110 002</u>	Dec 30, 2016
<u>AP</u>	GLAND	<u>50MG/VIAL</u>	<u>A207092 001</u>	Sep 29, 2017
<u>AP</u>		<u>70MG/VIAL</u>	<u>A207092 002</u>	Sep 29, 2017
<u>AP</u>	JIANGSU PHARMS	<u>50MG/VIAL</u>	<u>A200833 001</u>	Jun 28, 2018
<u>AP</u>		<u>70MG/VIAL</u>	<u>A200833 002</u>	Jun 28, 2018
<u>AP</u>	UBI	<u>50MG/VIAL</u>	<u>A211263 001</u>	Oct 01, 2021
<u>AP</u>		<u>70MG/VIAL</u>	<u>A211263 002</u>	Oct 01, 2021
<u>AP</u>	XELLIA PHARMS APS	<u>50MG/VIAL</u>	<u>A205923 001</u>	Jul 02, 2018
<u>AP</u>		<u>70MG/VIAL</u>	<u>A205923 002</u>	Jul 02, 2018

CEDAZURIDINE; DECITABINE

TABLET; ORAL

INQOVI

+! OTSUKA

100MG; 35MG

N212576 001 Jul 07, 2020

CEFACLOR

CAPSULE; ORAL

CEFACLOR

<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

CEFACLOR

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

CEFACLOR

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>	! TEVA PHARMS	<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
	! TEVA PHARMS	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>	QILU	<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

PRESCRIPTION DRUG PRODUCT LIST

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

CEFAZOLIN SODIUM

AP		EQ 10GM BASE/VIAL	A062831 003	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
	SOLUTION; INTRAVENOUS			
	CEFAZOLIN IN PLASTIC CONTAINER			
	+! BAXTER HLTHCARE	EQ 1GM BASE/50ML (EQ 20MG BASE/ML)	N207131 002	Feb 01, 2021
	CORP			
	+!	EQ 2GM BASE/100ML (EQ 20MG BASE/ML)	N207131 001	Aug 07, 2015

CEFDINIR

CAPSULE; ORAL

CEFDINIR

AB	ALKEM LABS LTD	300MG	A210220 001	Feb 19, 2021
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	ALKEM LABS LTD	125MG/5ML	A210534 001	Feb 19, 2021
AB		250MG/5ML	A210534 002	Feb 19, 2021
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	QILU	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 INC	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

CEFIDEROCOL SULFATE TOSYLATE

POWDER; INTRAVENOUS

FETROJA

	+! SHIONOGI INC	EQ 1GM BASE/VIAL	N209445 001	Nov 14, 2019
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PRESCRIPTION DRUG PRODUCT LIST

CEFIXIME

CAPSULE; ORAL

CEFIXIME

! ALKEM LABS LTD

400MG

A210574 001 Oct 09, 2018

FOR SUSPENSION; ORAL

CEFIXIMEAB ALKEM LABS LTD100MG/5MLA211775 001 Feb 19, 2021AB200MG/5MLA211775 002 Feb 19, 2021AB AUROBINDO PHARMA LTD100MG/5MLA204835 001 Apr 14, 2015AB200MG/5MLA204835 002 Apr 14, 2015AB BELCHER100MG/5MLA206938 001 Feb 06, 2017AB200MG/5MLA206938 002 Feb 06, 2017AB500MG/5MLA206939 001 Feb 06, 2017SUPRAXAB +! LUPIN LTD500MG/5MLN202091 001 Feb 20, 2013AB ! LUPIN PHARMS200MG/5MLA065355 001 Apr 10, 2007

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

! HIKMA

EQ 500MG BASE/VIAL

A065072 001 Nov 20, 2002

!

EQ 1GM BASE/VIAL

A065072 002 Nov 20, 2002

!

EQ 2GM BASE/VIAL

A065072 003 Nov 20, 2002

!

EQ 10GM BASE/VIAL

A065071 001 Nov 20, 2002

CEFOTETAN DISODIUM

INJECTABLE; INJECTION

CEFOTANAP + TELIGENTEQ 1GM BASE/VIALN050588 001 Dec 27, 1985AP +EQ 2GM BASE/VIALN050588 002 Dec 27, 1985CEFOTETANAP ! FRESENIUS KABI USAEQ 1GM BASE/VIALA065374 001 Aug 09, 2007AP !EQ 2GM BASE/VIALA065374 002 Aug 09, 2007

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

CEFOXITINAP ! ACS DOBFAREQ 1GM BASE/VIALA065414 001 Jun 12, 2009AP !EQ 2GM BASE/VIALA065414 002 Jun 12, 2009AP !EQ 10GM BASE/VIALA065415 001 May 19, 2010AP HIKMAEQ 1GM BASE/VIALA065051 001 Sep 11, 2000APEQ 2GM BASE/VIALA065051 002 Sep 11, 2000APEQ 10GM BASE/VIALA065050 001 Sep 11, 2000AP HIKMA FARMACEUTICAEQ 1GM BASE/VIALA065238 001 Mar 12, 2010APEQ 2GM BASE/VIALA065238 002 Mar 12, 2010APEQ 10GM BASE/VIALA065239 001 Mar 02, 2010CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINERAP +! B BRAUNEQ 1GM BASE/VIALN065214 001 Mar 10, 2006AP +!EQ 2GM BASE/VIALN065214 002 Mar 10, 2006

POWDER; INTRAVENOUS

CEFOXITIN IN PLASTIC CONTAINER

SAMSON MEDCL

EQ 100GM BASE

A200938 001 Nov 16, 2015

CEFPODOXIME PROXETIL

FOR SUSPENSION; ORAL

CEFPODOXIME PROXETIL

AUROBINDO PHARMA LTD

EQ 50MG BASE/5ML

A065409 001 Jun 08, 2007

!

EQ 100MG BASE/5ML

A065409 002 Jun 08, 2007

TABLET; ORAL

CEFPODOXIME PROXETILAB AUROBINDO PHARMAEQ 100MG BASEA065370 001 Jun 11, 2007ABEQ 200MG BASEA065370 002 Jun 11, 2007AB ORCHID HLTHCAREEQ 100MG BASEA065388 001 Nov 14, 2007ABEQ 200MG BASEA065388 002 Nov 14, 2007AB SANDOZEQ 100MG BASEA065462 001 May 28, 2008

PRESCRIPTION DRUG PRODUCT LIST

CEFPODOXIME PROXETIL

TABLET; ORAL

CEFPODOXIME PROXETIL

AB	!		<u>EQ 200MG BASE</u>	<u>A065462 002</u>	May 28, 2008
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CEFPROZIL

FOR SUSPENSION; ORAL

CEFPROZIL

AB		APOTEX INC	<u>125MG/5ML</u>	<u>A065351 001</u>	Feb 29, 2012
AB			<u>250MG/5ML</u>	<u>A065351 002</u>	Feb 29, 2012
AB		AUROBINDO PHARMA	<u>125MG/5ML</u>	<u>A065381 001</u>	Jan 30, 2007
AB			<u>250MG/5ML</u>	<u>A065381 002</u>	Jan 30, 2007
AB		LUPIN	<u>125MG/5ML</u>	<u>A065261 001</u>	Dec 19, 2005
AB	!		<u>250MG/5ML</u>	<u>A065261 002</u>	Dec 19, 2005
AB		ORCHID HLTHCARE	<u>125MG/5ML</u>	<u>A065284 002</u>	Dec 30, 2005
AB			<u>250MG/5ML</u>	<u>A065284 001</u>	Dec 30, 2005
AB		SANDOZ	<u>125MG/5ML</u>	<u>A065257 001</u>	Dec 08, 2005
AB			<u>250MG/5ML</u>	<u>A065257 002</u>	Dec 08, 2005

TABLET; ORAL

CEFPROZIL

AB		APOTEX INC	<u>250MG</u>	<u>A065327 001</u>	Mar 26, 2008
AB			<u>500MG</u>	<u>A065327 002</u>	Mar 26, 2008
AB		AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A065340 001</u>	May 24, 2007
AB			<u>500MG</u>	<u>A065340 002</u>	May 24, 2007
AB		CHARTWELL RX	<u>250MG</u>	<u>A065235 001</u>	Nov 14, 2005
AB			<u>500MG</u>	<u>A065235 002</u>	Nov 14, 2005
AB		LUPIN	<u>250MG</u>	<u>A065276 001</u>	Dec 08, 2005
AB	!		<u>500MG</u>	<u>A065276 002</u>	Dec 08, 2005
AB		ORCHID HLTHCARE	<u>250MG</u>	<u>A065267 001</u>	Dec 19, 2005
AB			<u>500MG</u>	<u>A065267 002</u>	Dec 19, 2005
AB		TEVA	<u>250MG</u>	<u>A065208 001</u>	Dec 06, 2005
AB			<u>500MG</u>	<u>A065208 002</u>	Dec 06, 2005

CEFTAROLINE FOSAMIL

POWDER; INTRAVENOUS

CEFTAROLINE FOSAMIL

AP		APOTEX	<u>400MG/VIAL</u>	<u>A208075 001</u>	Sep 21, 2021
AP			<u>600MG/VIAL</u>	<u>A208075 002</u>	Sep 21, 2021
		<u>TEFLARO</u>			
AP	+	ALLERGAN	<u>400MG/VIAL</u>	<u>N200327 001</u>	Oct 29, 2010
AP	+		<u>600MG/VIAL</u>	<u>N200327 002</u>	Oct 29, 2010

CEFTAZIDIME

INJECTABLE; INJECTION

CEFTAZIDIME

AP		ACS DOBFAR	<u>1GM/VIAL</u>	<u>A062640 002</u>	Nov 20, 1985
AP			<u>2GM/VIAL</u>	<u>A062640 003</u>	Nov 20, 1985
AP			<u>6GM/VIAL</u>	<u>A062640 004</u>	Feb 03, 1992
		<u>FORTAZ</u>			
AP	+	TELIGENT	<u>500MG/VIAL</u>	<u>N050578 001</u>	Jul 19, 1985
AP	+		<u>1GM/VIAL</u>	<u>N050578 002</u>	Jul 19, 1985
AP	+		<u>2GM/VIAL</u>	<u>N050578 003</u>	Jul 19, 1985
AP	+		<u>6GM/VIAL</u>	<u>N050578 004</u>	Jul 19, 1985
		<u>TAZICEF</u>			
AP		HOSPIRA	<u>500MG/VIAL</u>	<u>A062662 001</u>	Mar 06, 1986
AP			<u>1GM/VIAL</u>	<u>A062662 002</u>	Mar 06, 1986
AP			<u>1GM/VIAL</u>	<u>A064032 001</u>	Oct 31, 1993
AP			<u>2GM/VIAL</u>	<u>A062662 003</u>	Mar 06, 1986
AP			<u>2GM/VIAL</u>	<u>A064032 002</u>	Oct 31, 1993
AP			<u>6GM/VIAL</u>	<u>A062662 004</u>	Mar 06, 1986

CEFTAZIDIME IN DEXTROSE CONTAINER

+	B BRAUN	<u>EQ 1GM BASE</u>	<u>N050823 001</u>	Jun 13, 2011
+		<u>EQ 2GM BASE</u>	<u>N050823 002</u>	Jun 13, 2011

CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM

POWDER; INTRAVENOUS

ZERBAXA

+	CUBIST PHARMS LLC	<u>EQ 1GM BASE/VIAL;EQ 0.5GM BASE/VIAL</u>	<u>N206829 001</u>	Dec 19, 2014
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PRESCRIPTION DRUG PRODUCT LIST

CEFTRIAXONE SODIUM

INJECTABLE; INJECTION

CEFTRIAXONE

AP	ACS DOBFAR	<u>EQ 500MG BASE/VIAL</u>	<u>A065329 001</u>	Jul 24, 2008
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A065329 002</u>	Jul 24, 2008
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A065329 003</u>	Jul 24, 2008
AP		<u>EQ 10GM BASE/VIAL</u>	<u>A065328 001</u>	Jul 24, 2008
AP	QILU	<u>EQ 10GM BASE/VIAL</u>	<u>A209218 001</u>	Oct 17, 2018
AP	! SANDOZ	<u>EQ 10GM BASE/VIAL</u>	<u>A065168 001</u>	May 17, 2005
AP	! SANDOZ INC	<u>EQ 1GM BASE/VIAL</u>	<u>A065204 001</u>	May 03, 2005
AP	!	<u>EQ 2GM BASE/VIAL</u>	<u>A065204 002</u>	May 03, 2005
AP	WOCKHARDT	<u>EQ 1GM BASE/VIAL</u>	<u>A065180 001</u>	May 12, 2006

CEFTRIAXONE AND DEXTROSE IN DUPLIX CONTAINER

AP	+! B BRAUN	<u>EQ 1GM BASE/VIAL</u>	<u>N050796 001</u>	Apr 20, 2005
AP	+!	<u>EQ 2GM BASE/VIAL</u>	<u>N050796 002</u>	Apr 20, 2005

CEFTRIAXONE SODIUM

AP	ASTRAL	<u>EQ 10GM BASE/VIAL</u>	<u>A091117 001</u>	Jan 20, 2017
AP	HIKMA	<u>EQ 10GM BASE/VIAL</u>	<u>A090701 001</u>	Oct 04, 2017

CEFTRIAXONE

SAMSON MEDCL EQ 100GM BASE/VIAL A090057 001 Apr 25, 2014

CEFTRIAXONE 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

CEFTRIAXONE

AP	AKORN	<u>EQ 250MG BASE/VIAL</u>	<u>A065305 001</u>	Jan 11, 2008
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A065305 002</u>	Jan 11, 2008
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A065305 003</u>	Jan 11, 2008
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A065305 004</u>	Jan 11, 2008
AP	ASTRAL	<u>EQ 250MG BASE/VIAL</u>	<u>A091049 001</u>	Jun 11, 2018
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A091049 002</u>	Jun 11, 2018
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A091049 003</u>	Jun 11, 2018
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A091049 004</u>	Jun 11, 2018
AP	HIKMA FARMACEUTICA	<u>EQ 250MG BASE/VIAL</u>	<u>A065342 001</u>	Jan 10, 2008
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A065342 002</u>	Jan 10, 2008
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A065342 003</u>	Jan 10, 2008
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A065342 004</u>	Jan 10, 2008
AP	LUPIN	<u>EQ 250MG BASE/VIAL</u>	<u>A065125 001</u>	Sep 30, 2003
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A065125 002</u>	Sep 30, 2003
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A065125 003</u>	Sep 30, 2003
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A065125 004</u>	Sep 30, 2003
AP	QILU	<u>EQ 250MG BASE/VIAL</u>	<u>A203702 001</u>	Jun 29, 2016
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A203702 002</u>	Jun 29, 2016
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A203702 003</u>	Jun 29, 2016
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A203702 004</u>	Jun 29, 2016
AP	! SANDOZ	<u>EQ 250MG BASE/VIAL</u>	<u>A065169 001</u>	May 09, 2005
AP	!	<u>EQ 500MG BASE/VIAL</u>	<u>A065169 002</u>	May 09, 2005
AP	!	<u>EQ 1GM BASE/VIAL</u>	<u>A065169 003</u>	May 09, 2005
AP	!	<u>EQ 2GM BASE/VIAL</u>	<u>A065169 004</u>	May 09, 2005
AP	WOCKHARDT	<u>EQ 250MG BASE/VIAL</u>	<u>A065391 001</u>	Apr 12, 2007
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A065391 002</u>	Apr 12, 2007
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A065391 003</u>	Apr 12, 2007

CEFUROXIME AXETIL

TABLET; ORAL

CEFUROXIME AXETIL

AB	ALKEM LABS LTD	<u>EQ 250MG BASE</u>	<u>A065496 001</u>	Jun 07, 2010
AB		<u>EQ 500MG BASE</u>	<u>A065496 002</u>	Jun 07, 2010
AB	APOTEX	<u>EQ 250MG BASE</u>	<u>A065069 001</u>	Oct 02, 2002
AB		<u>EQ 500MG BASE</u>	<u>A065069 002</u>	Oct 02, 2002
AB	AUROBINDO PHARMA LTD	<u>EQ 125MG BASE</u>	<u>A065308 001</u>	Mar 29, 2006
AB		<u>EQ 250MG BASE</u>	<u>A065308 002</u>	Mar 29, 2006
AB		<u>EQ 500MG BASE</u>	<u>A065308 003</u>	Mar 29, 2006
AB	LUPIN	<u>EQ 250MG BASE</u>	<u>A065135 001</u>	Jul 25, 2003
AB	!	<u>EQ 500MG BASE</u>	<u>A065135 002</u>	Jul 25, 2003
AB	ORCHID HLTHCARE	<u>EQ 125MG BASE</u>	<u>A065359 001</u>	Feb 15, 2008
AB		<u>EQ 250MG BASE</u>	<u>A065359 002</u>	Feb 15, 2008
AB		<u>EQ 500MG BASE</u>	<u>A065359 003</u>	Feb 15, 2008
AB	WOCKHARDT	<u>EQ 125MG BASE</u>	<u>A065166 001</u>	Jul 29, 2005
AB		<u>EQ 250MG BASE</u>	<u>A065166 002</u>	Jul 29, 2005
AB		<u>EQ 500MG BASE</u>	<u>A065166 003</u>	Jul 29, 2005

PRESCRIPTION DRUG PRODUCT LIST

CEFUROXIME SODIUM

INJECTABLE; INJECTION

CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER

<u>AP</u>	<u>+!</u>	<u>B BRAUN</u>	<u>EQ 750MG BASE/VIAL</u>	<u>N050780 001</u>	Feb 21, 2001
<u>AP</u>	<u>+!</u>		<u>EQ 1.5GM BASE/VIAL</u>	<u>N050780 002</u>	Feb 21, 2001

CEFUROXIME SODIUM

<u>AP</u>		<u>ACS DOBFAR SPA</u>	<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>		<u>HIKMA FARMACEUTICA</u>	<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

ZINACEF

<u>AP</u>	<u>+!</u>	<u>TELIGENT</u>	<u>EQ 1.5GM BASE/VIAL</u>	<u>N050558 003</u>	Oct 19, 1983
<u>AP</u>	<u>+!</u>		<u>EQ 7.5GM BASE/VIAL</u>	<u>N050558 004</u>	Oct 23, 1986

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFUROXIME SODIUM

<u>AB</u>		<u>ACS DOBFAR SPA</u>	<u>EQ 750MG BASE/VIAL</u>	<u>A064125 001</u>	May 30, 1997
<u>AB</u>		<u>HIKMA FARMACEUTICA</u>	<u>EQ 750MG BASE/VIAL</u>	<u>A065048 001</u>	Jan 09, 2004

ZINACEF

<u>AB</u>	<u>+!</u>	<u>TELIGENT</u>	<u>EQ 750MG BASE/VIAL</u>	<u>N050558 002</u>	Oct 19, 1983
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CELECOXIB

CAPSULE; ORAL

CELEBREX

<u>AB</u>	<u>+</u>	<u>UPJOHN</u>	<u>50MG</u>	<u>N020998 004</u>	Dec 15, 2006
<u>AB</u>	<u>+</u>		<u>100MG</u>	<u>N020998 001</u>	Dec 31, 1998
<u>AB</u>	<u>+</u>		<u>200MG</u>	<u>N020998 002</u>	Dec 31, 1998
<u>AB</u>	<u>+!</u>		<u>400MG</u>	<u>N020998 003</u>	Aug 29, 2002

CELECOXIB

<u>AB</u>		<u>ALEMBIC PHARMS LTD</u>	<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>		<u>APOTEX INC</u>	<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>		<u>AUROBINDO PHARMA LTD</u>	<u>50MG</u>	<u>A206827 001</u>	Feb 01, 2016
<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>		<u>CADILA PHARMS LTD</u>	<u>50MG</u>	<u>A208701 001</u>	Nov 14, 2019
<u>AB</u>			<u>100MG</u>	<u>A208701 002</u>	Nov 14, 2019
<u>AB</u>			<u>200MG</u>	<u>A208701 003</u>	Nov 14, 2019
<u>AB</u>			<u>400MG</u>	<u>A208701 004</u>	Nov 14, 2019
<u>AB</u>		<u>CIPLA</u>	<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>		<u>CSPC OUYI</u>	<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>		<u>LUPIN LTD</u>	<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>		<u>MACLEODS PHARMS LTD</u>	<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>		<u>MICRO LABS</u>	<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>		<u>MYLAN</u>	<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>		<u>NANJING</u>	<u>200MG</u>	<u>A213598 001</u>	May 13, 2020
<u>AB</u>		<u>QINGDAO BAHEAL PHARM</u>	<u>50MG</u>	<u>A208856 001</u>	Aug 07, 2019
<u>AB</u>			<u>100MG</u>	<u>A208856 002</u>	Aug 07, 2019
<u>AB</u>			<u>200MG</u>	<u>A208856 003</u>	Aug 07, 2019
<u>AB</u>			<u>400MG</u>	<u>A208856 004</u>	Aug 07, 2019

PRESCRIPTION DRUG PRODUCT LIST

CELECOXIB

CAPSULE; ORAL

CELECOXIB

AB	SCIEGEN PHARMS INC	50MG	<u>A205129 001</u>	Dec 03, 2020
AB		100MG	<u>A205129 002</u>	Dec 03, 2020
AB		200MG	<u>A205129 003</u>	Dec 03, 2020
AB		400MG	<u>A205129 004</u>	Dec 03, 2020
AB	TEVA	50MG	<u>A076898 001</u>	May 30, 2014
AB		100MG	<u>A076898 002</u>	May 30, 2014
AB		200MG	<u>A076898 003</u>	May 30, 2014
AB		400MG	<u>A076898 004</u>	May 30, 2014
AB	TIANJIN TIANYAO	50MG	<u>A207872 001</u>	Feb 25, 2020
AB		100MG	<u>A207872 002</u>	Feb 25, 2020
AB		200MG	<u>A207872 003</u>	Feb 25, 2020
AB		400MG	<u>A207872 004</u>	Feb 25, 2020
AB	TORRENT	50MG	<u>A207677 001</u>	Dec 23, 2015
AB		100MG	<u>A207677 002</u>	Dec 23, 2015
AB		200MG	<u>A207677 003</u>	Dec 23, 2015
AB		400MG	<u>A207677 004</u>	Dec 23, 2015
AB	UMEDICA LABS PVT LTD	50MG	<u>A210628 001</u>	Nov 27, 2019
AB		100MG	<u>A210628 002</u>	Nov 27, 2019
AB		200MG	<u>A210628 003</u>	Nov 27, 2019
AB		400MG	<u>A210628 004</u>	Nov 27, 2019
AB	UNICHEM	50MG	<u>A213301 001</u>	Jan 12, 2021
AB		100MG	<u>A213301 002</u>	Jan 12, 2021
AB		200MG	<u>A213301 003</u>	Jan 12, 2021
AB		400MG	<u>A213301 004</u>	Jan 12, 2021
AB	WATSON LABS INC	50MG	<u>A200562 001</u>	Feb 11, 2015
AB		100MG	<u>A200562 002</u>	Feb 11, 2015
AB		200MG	<u>A200562 003</u>	Feb 11, 2015
AB		400MG	<u>A200562 004</u>	Feb 11, 2015
AB	YILING	50MG	<u>A211412 001</u>	Mar 06, 2020
AB		100MG	<u>A211412 002</u>	Mar 06, 2020
AB		200MG	<u>A211412 003</u>	Mar 06, 2020
AB		400MG	<u>A211412 004</u>	Mar 06, 2020
BX	AMNEAL PHARMS	50MG	A208833 001	May 31, 2018
BX		100MG	A208833 002	May 31, 2018
BX		200MG	A208833 003	May 31, 2018
BX		400MG	A208833 004	May 31, 2018

SOLUTION; ORAL

ELYXYB

+! BDSI

25MG/ML

N212157 001 May 05, 2020

CELECOXIB; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

SEGLENTIS

+! KOWA PHARMS

56MG; 44MG

N213426 001 Oct 15, 2021

CENOBAAMATE

TABLET; ORAL

XCOPRI

+! SK LIFE

12.5MG

N212839 001 Mar 10, 2020

+

25MG

N212839 002 Mar 10, 2020

+

50MG

N212839 003 Mar 10, 2020

+

100MG

N212839 004 Mar 10, 2020

+

150MG

N212839 005 Mar 10, 2020

+

200MG

N212839 006 Mar 10, 2020

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

AB	ALKEM LABS LTD	EQ 250MG BASE	<u>A090836 001</u>	Dec 20, 2010
AB		EQ 500MG BASE	<u>A090836 002</u>	Dec 20, 2010
AB	AUROBINDO PHARMA LTD	EQ 250MG BASE	<u>A065253 001</u>	Nov 16, 2005
AB		EQ 500MG BASE	<u>A065253 002</u>	Nov 16, 2005
AB	BELCHER PHARMS	EQ 250MG BASE	<u>A062713 001</u>	Jul 15, 1988
AB		EQ 500MG BASE	<u>A062713 002</u>	Jul 15, 1988
AB	CHARTWELL RX	EQ 250MG BASE	<u>A065152 001</u>	Feb 24, 2005
AB		EQ 500MG BASE	<u>A065152 002</u>	Feb 24, 2005
AB	HIKMA	EQ 250MG BASE	<u>A065215 001</u>	Jan 24, 2006
AB		EQ 500MG BASE	<u>A065215 002</u>	Jan 24, 2006
AB	LUPIN	EQ 250MG BASE	<u>A065229 001</u>	Nov 25, 2005

PRESCRIPTION DRUG PRODUCT LIST

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

<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>	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
	ALKEM LABS LTD	EQ 333MG BASE	A090836 003	Mar 29, 2013
	!	EQ 750MG BASE	A090836 004	Mar 29, 2013

FOR SUSPENSION; ORAL

CEPHALEXIN

<u>AB</u>	ALKEM LABS LTD	<u>EQ 125MG BASE/5ML</u>	<u>A210221 001</u>	Mar 26, 2019
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A210221 002</u>	Mar 26, 2019
<u>AB</u>	LUPIN	<u>EQ 125MG BASE/5ML</u>	<u>A065234 001</u>	Aug 17, 2005
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065234 002</u>	Aug 17, 2005
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 125MG BASE/5ML</u>	<u>A065326 001</u>	Jul 10, 2006
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065326 002</u>	Jul 10, 2006
<u>AB</u>	TEVA	<u>EQ 125MG BASE/5ML</u>	<u>A062703 001</u>	Feb 13, 1987
<u>AB</u>	!	<u>EQ 250MG BASE/5ML</u>	<u>A062703 002</u>	Feb 13, 1987
<u>AB</u>	YUNG SHIN PHARM	<u>EQ 125MG BASE/5ML</u>	<u>A065336 001</u>	Jul 25, 2007
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065336 002</u>	Jul 25, 2007

TABLET; ORAL

CEPHALEXIN

TEVA

!

EQ 250MG BASE	A063023 001	Jan 12, 1989
EQ 500MG BASE	A063024 001	Jan 12, 1989

CERITINIB

TABLET; ORAL

ZYKADIA

+! NOVARTIS

150MG	N211225 001	Mar 18, 2019
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CETIRIZINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

QUZYTIR

+! JDP

10MG/ML (10MG/ML)	N211415 001	Oct 04, 2019
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SOLUTION/DROPS; OPHTHALMIC

ZERVIAE

+! EYEVANCE

EQ 0.24% BASE	N208694 001	May 30, 2017
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SYRUP; ORAL

CETIRIZINE HYDROCHLORIDE

<u>AA</u>	AMNEAL PHARMS	<u>5MG/5ML</u>	<u>A090766 001</u>	Oct 07, 2009
<u>AA</u>	BRECKENRIDGE	<u>5MG/5ML</u>	<u>A078488 001</u>	Oct 06, 2008
<u>AA</u>	LANNETT CO INC	<u>5MG/5ML</u>	<u>A078876 001</u>	May 11, 2012
<u>AA</u>	MICRO LABS	<u>5MG/5ML</u>	<u>A090191 001</u>	Nov 12, 2009
<u>AA</u>	! PADAGIS US	<u>5MG/5ML</u>	<u>A078398 001</u>	Jun 17, 2008
<u>AA</u>	TARO	<u>5MG/5ML</u>	<u>A076601 001</u>	Jun 20, 2008
<u>AA</u>	TEVA PHARMS	<u>5MG/5ML</u>	<u>A077279 001</u>	May 27, 2008

CETRORELIX

INJECTABLE; INJECTION

CETROTIDE

+! EMD SERONO INC

EQ 0.25MG BASE/ML	N021197 001	Aug 11, 2000
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CEVIMELINE HYDROCHLORIDE

CAPSULE; ORAL

CEVIMELINE HYDROCHLORIDE

<u>AB</u>	HIKMA	<u>30MG</u>	<u>A091591 001</u>	Jul 08, 2013
<u>AB</u>	NOVEL LABS INC	<u>30MG</u>	<u>A204746 001</u>	Dec 30, 2016
<u>AB</u>	RISING	<u>30MG</u>	<u>A203775 001</u>	Jun 04, 2014

EVOXAC

<u>AB</u>	+! DAIICHI SANKYO INC	<u>30MG</u>	<u>N020989 002</u>	Jan 11, 2000
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CHENODIOL

TABLET; ORAL

CHENODIOL

! LGM PHARMA

250MG	A091019 001	Oct 22, 2009
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CHLORAMBUCIL

TABLET; ORAL

LEUKERAN

+! ASPEN GLOBAL INC

2MG	N010669 002	
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PRESCRIPTION DRUG PRODUCT LIST

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION

CHLORAMPHENICOL SODIUM SUCCINATE

! FRESENIUS KABI USA EQ 1GM BASE/VIAL

A062365 001 Aug 25, 1982

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE

AB	BARR	5MG	A084768	001	
AB		10MG	A083116	001	
AB		25MG	A084769	001	

LIBRIUM

AB	VALEANT PHARM INTL	5MG	A085461	001	
AB		10MG	A085472	001	
AB	+	25MG	A085475	001	

CHLORDIAZEPOXIDE HYDROCHLORIDE; CLIDINIUM BROMIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE

AB	ALKEM LABS LTD	5MG; 2.5MG	A214065	001	Apr 26, 2021
AB	AMNEAL	5MG; 2.5MG	A215555	001	Oct 25, 2021
AB	DR REDDYS	5MG; 2.5MG	A214698	001	May 10, 2021
AB	MISEMER	5MG; 2.5MG	A210579	001	Jul 29, 2020
AB	NUVO PHARMS INC	5MG; 2.5MG	A211421	001	Jul 07, 2020
AB	PHARMA LIFE	5MG; 2.5MG	A213530	001	Oct 20, 2020
AB	TEVA PHARMS USA	5MG; 2.5MG	A211476	001	Nov 02, 2021
LIBRAX					
AB	+	BAUSCH	5MG; 2.5MG	N012750	001

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL

CHLORHEXIDINE GLUCONATE

AT	AKORN	0.12%	A074356	001	May 07, 1996
AT	BAJAJ	0.12%	A075561	001	Nov 14, 2000
AT	LYNE	0.12%	A074291	001	Dec 28, 1995
AT	PHARM ASSOC	0.12%	A074522	001	Dec 15, 1995
AT	WOCKHARDT BIO AG	0.12%	A075006	001	Mar 03, 2004
AT	XTTRIUM	0.12%	A077789	001	Jun 18, 2009
PAROEX					
AT	SUNSTAR AMERICAS	0.12%	A076434	001	Nov 29, 2005
PERIDEX					
AT	+	3M	0.12%	N019028	001 Aug 13, 1986
PERIOGARD					
AT	COLGATE PALMOLIVE CO	0.12%	A073695	001	Jan 14, 1994
AT	COLGATE-PALMOLIVE CO	0.12%	A203212	001	Jan 28, 2016
TABLET; DENTAL					
PERIOCHIP					
	+	DEXCEL PHARMA	2.5MG	N020774	001 May 15, 1998

CHLOROPROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHLOROPROCAINE HYDROCHLORIDE

AP	HIKMA	2%	A040273	001	Sep 09, 1998
AP		3%	A040273	002	Sep 09, 1998
NESACAINE					
AP	+	FRESENIUS KABI USA	2%	N009435	002
NESACAINE-MPF					
AP	+	FRESENIUS KABI USA	2%	N009435	006 May 02, 1996
AP	+		3%	N009435	007 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

AA	IPCA LABS LTD	250MG	A090610	001	Dec 03, 2009
AA		500MG	A090249	001	Dec 03, 2009
AA	!	NATCO PHARMA LTD	250MG	A091621	001 Jan 21, 2011
AA	!		500MG	A090612	001 Jan 21, 2011
AA	SUVEN PHARMS	500MG	A214756	001	Sep 03, 2021

PRESCRIPTION DRUG PRODUCT LIST

CHLOROTHIAZIDE

SUSPENSION; ORAL

DIURIL

+! SALIX PHARMS 250MG/5ML N011870 001

CHLOROTHIAZIDE SODIUM

INJECTABLE; INJECTION

CHLOROTHIAZIDE SODIUM

AP	AM REGENT	EQ 500MG BASE/VIAL	A202561 001	Apr 22, 2013
AP	FRESENIUS KABI USA	EQ 500MG BASE/VIAL	A090896 001	Oct 16, 2009
AP	RK PHARMA	EQ 500MG BASE/VIAL	A202493 001	Jun 18, 2014
AP	SAGENT PHARMS INC	EQ 500MG BASE/VIAL	A202462 001	May 29, 2015
AP	SUN PHARM	EQ 500MG BASE/VIAL	A091546 001	Jul 26, 2011

DIURIL**AP** +! AKORN **EQ 500MG BASE/VIAL** **N011145 005**CHLORPHENIRAMINE MALEATE; CODEINE PHOSPHATE

TABLET, EXTENDED RELEASE; ORAL

TUXARIN ER

MAINPOINTE 8MG; 54.3MG N206323 001 Jun 22, 2015

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

! PADAGIS US 4MG/5ML; 5MG/5ML; 60MG/5ML A204627 001 Apr 29, 2014

CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

TUZISTRA XR

+! AYTU 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**AB** TRIS PHARMA INC **EQ 8MG MALEATE/5ML; EQ 10MG BITARTRATE/5ML** **A091632 001** Oct 01, 2010HYDROCODONE POLISTIREX AND CHLORPHENIRAMNE POLISTIREX**AB** ! NEOS THERAP INC **EQ 8MG MALEATE/5ML; EQ 10MG BITARTRATE/5ML** **A091671 001** Jun 29, 2012CHLORPROMAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

CHLORPROMAZINE HYDROCHLORIDE

! GENUS 30MG/ML A214542 001 Jun 02, 2021

! 100MG/ML A214542 002 Jun 02, 2021

INJECTABLE; INJECTION

CHLORPROMAZINE HYDROCHLORIDE**AP** EUGIA PHARMA **25MG/ML** **A211816 001** Jul 07, 2020**AP** ! WEST-WARD PHARMS **25MG/ML** **A083329 001**

INT

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE**AB** ENALTEC **10MG** **A212630 001** Nov 29, 2021**AB** **25MG** **A212630 002** Nov 29, 2021**AB** **50MG** **A212630 003** Nov 29, 2021**AB** **100MG** **A212630 004** Nov 29, 2021**AB** **200MG** **A212630 005** Nov 29, 2021**AB** GLENMARK PHARMS LTD **10MG** **A212144 001** Mar 23, 2021**AB** **25MG** **A212144 002** Mar 23, 2021**AB** **50MG** **A212144 003** Mar 23, 2021**AB** **100MG** **A212144 004** Mar 23, 2021**AB** **200MG** **A212144 005** Mar 23, 2021**AB** LANNETT CO INC **10MG** **A212996 001** Jan 22, 2021**AB** **25MG** **A212996 002** Jan 22, 2021**AB** **50MG** **A212996 003** Jan 22, 2021**AB** **100MG** **A212996 004** Jan 22, 2021**AB** **200MG** **A212996 005** Jan 22, 2021**AB** SUN PHARM **10MG** **A214256 001** Oct 26, 2020**AB** **25MG** **A214256 002** Oct 26, 2020**AB** **50MG** **A214256 003** Oct 26, 2020**AB** **100MG** **A214256 004** Oct 26, 2020**AB** **200MG** **A214256 005** Oct 26, 2020

PRESCRIPTION DRUG PRODUCT LIST

CHLORPROMAZINE HYDROCHLORIDE

TABLET;ORAL

CHLORPROMAZINE HYDROCHLORIDE

<u>AB</u>	TEVA PHARMS	<u>10MG</u>	<u>A215659 001</u>	Oct 25, 2021
<u>AB</u>		<u>25MG</u>	<u>A215659 002</u>	Oct 25, 2021
<u>AB</u>		<u>50MG</u>	<u>A215659 003</u>	Oct 25, 2021
<u>AB</u>		<u>100MG</u>	<u>A215659 004</u>	Oct 25, 2021
<u>AB</u>		<u>200MG</u>	<u>A215659 005</u>	Oct 25, 2021
<u>AB</u>	+	UPSHER SMITH LABS	<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>
<u>AB</u>	ZYDUS	<u>10MG</u>	<u>A213368 001</u>	Jan 17, 2020
<u>AB</u>		<u>25MG</u>	<u>A213368 002</u>	Jan 17, 2020
<u>AB</u>		<u>50MG</u>	<u>A213368 003</u>	Jan 17, 2020
<u>AB</u>		<u>100MG</u>	<u>A213368 004</u>	Jan 17, 2020
<u>AB</u>		<u>200MG</u>	<u>A213368 005</u>	Jan 17, 2020
BX	AMNEAL PHARMS CO	10MG	A209755 001	Sep 10, 2018
BX		25MG	A209755 002	Sep 10, 2018
BX		50MG	A209755 003	Sep 10, 2018
BX		100MG	A209755 004	Sep 10, 2018
BX		200MG	A209755 005	Sep 10, 2018

CHLOROTHALIDONE

TABLET;ORAL

CHLOROTHALIDONE

<u>AB</u>	AJANTA PHARMA LTD	<u>25MG</u>	<u>A214129 001</u>	Nov 27, 2020
<u>AB</u>		<u>50MG</u>	<u>A214129 002</u>	Nov 27, 2020
<u>AB</u>	ALKEM LABS LTD	<u>25MG</u>	<u>A213412 001</u>	Feb 11, 2020
<u>AB</u>		<u>50MG</u>	<u>A213412 002</u>	Feb 11, 2020
<u>AB</u>	AMNEAL PHARMS CO	<u>25MG</u>	<u>A207204 001</u>	Jul 01, 2019
<u>AB</u>		<u>50MG</u>	<u>A207204 002</u>	Jul 01, 2019
<u>AB</u>	APPCO	<u>25MG</u>	<u>A210742 001</u>	Oct 12, 2018
<u>AB</u>		<u>50MG</u>	<u>A210742 002</u>	Oct 12, 2018
<u>AB</u>	+	MYLAN	<u>25MG</u>	<u>A086831 002</u>
<u>AB</u>	+	!	<u>50MG</u>	<u>A086831 001</u>
<u>AB</u>	NOVAST LABS	<u>25MG</u>	<u>A206904 001</u>	Mar 30, 2017
<u>AB</u>		<u>50MG</u>	<u>A206904 002</u>	Mar 30, 2017
<u>AB</u>	SUN PHARM INDUSTRIES	<u>25MG</u>	<u>A089286 002</u>	Jul 21, 1986
<u>AB</u>		<u>50MG</u>	<u>A089286 001</u>	Jul 21, 1986
<u>AB</u>	UNICHEM	<u>25MG</u>	<u>A211627 001</u>	Aug 06, 2019
<u>AB</u>		<u>50MG</u>	<u>A211627 002</u>	Aug 06, 2019
<u>AB</u>	VISTAPHARM	<u>25MG</u>	<u>A211063 001</u>	Feb 26, 2019
<u>AB</u>		<u>50MG</u>	<u>A211063 002</u>	Feb 26, 2019
<u>AB</u>	ZYDUS PHARMS	<u>25MG</u>	<u>A207813 001</u>	May 10, 2019
<u>AB</u>		<u>50MG</u>	<u>A207813 002</u>	May 10, 2019
BX	UMEDICA LABS PVT LTD	25MG	A207222 001	May 24, 2018
BX		50MG	A207222 002	May 24, 2018
	THALITONE			
BX	+	CASPER PHARMA LLC	N019574 002	Feb 12, 1992
	+		N019574 001	Dec 20, 1988

CHLORZOXAZONE

TABLET;ORAL

CHLORZOXAZONE

<u>AA</u>	AUROBINDO PHARMA LTD	<u>500MG</u>	<u>A089853 001</u>	May 04, 1988
<u>AA</u>	MAYNE PHARMA	<u>500MG</u>	<u>A211849 002</u>	Jul 21, 2020
<u>AA</u>	!	MIKART	<u>250MG</u>	<u>A207483 001</u>
<u>AA</u>	NOVITIUM PHARMA	<u>500MG</u>	<u>A212254 001</u>	Sep 12, 2019
<u>AA</u>	SENORES PHARMS	<u>250MG</u>	<u>A215158 001</u>	Jul 29, 2021
<u>AA</u>	!	WATSON LABS	<u>500MG</u>	<u>A089859 001</u>
<u>AB</u>	GLENMARK PHARMS LTD	<u>375MG</u>	<u>A212185 001</u>	May 26, 2020
<u>AB</u>		<u>750MG</u>	<u>A212185 002</u>	May 26, 2020
<u>AB</u>	I3 PHARMS	<u>375MG</u>	<u>A212053 001</u>	Sep 14, 2020
<u>AB</u>		<u>750MG</u>	<u>A212053 002</u>	Sep 14, 2020
<u>AB</u>	MAYNE PHARMA	<u>375MG</u>	<u>A211849 001</u>	Jul 21, 2020
<u>AB</u>		<u>750MG</u>	<u>A211849 003</u>	Jul 21, 2020
<u>AB</u>	!	MIKART	<u>375MG</u>	<u>A040861 001</u>
<u>AB</u>	!		<u>750MG</u>	<u>A040861 002</u>
<u>AB</u>	NOVITIUM PHARMA	<u>375MG</u>	<u>A212253 001</u>	Nov 27, 2019

PRESCRIPTION DRUG PRODUCT LIST

CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

<u>AB</u>		<u>750MG</u>	<u>A212253</u>	<u>002</u>	Nov 27, 2019
<u>AB</u>	PAR PHARM INC	<u>375MG</u>	<u>A212743</u>	<u>002</u>	Apr 29, 2021
<u>AB</u>		<u>750MG</u>	<u>A212743</u>	<u>001</u>	Nov 02, 2020
<u>AB</u>	TEVA PHARMS USA INC	<u>375MG</u>	<u>A212898</u>	<u>001</u>	Jun 17, 2020
<u>AB</u>		<u>750MG</u>	<u>A212898</u>	<u>002</u>	Jun 17, 2020

CHOLESTYRAMINE

POWDER; ORAL

CHOLESTYRAMINE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 4GM RESIN/PACKET</u>	<u>A211119</u>	<u>001</u>	Apr 06, 2020
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A211119</u>	<u>002</u>	Apr 06, 2020
<u>AB</u>	ALKEM LABS LTD	<u>EQ 4GM RESIN/PACKET</u>	<u>A211856</u>	<u>001</u>	Oct 19, 2021
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A211856</u>	<u>002</u>	Oct 19, 2021
<u>AB</u>	ANI PHARMS	<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>	! SANDOZ	<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>	TAGI	<u>EQ 4GM RESIN/PACKET</u>	<u>A209597</u>	<u>001</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A209597</u>	<u>002</u>	Mar 09, 2021
<u>AB</u>	ZYDUS PHARMS	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A202901</u>	<u>001</u>	Jul 02, 2018

CHOLESTYRAMINE LIGHT

<u>AB</u>	ALKEM LABS LTD	<u>EQ 4GM RESIN/PACKET</u>	<u>A211799</u>	<u>001</u>	Oct 19, 2021
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A211799</u>	<u>002</u>	Oct 19, 2021
<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>	! SANDOZ	<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>	TAGI	<u>EQ 4GM RESIN/PACKET</u>	<u>A209599</u>	<u>001</u>	Nov 12, 2020
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A209599</u>	<u>002</u>	Nov 12, 2020
<u>AB</u>	ZYDUS PHARMS	<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

+	TRAVERE	50MG	N205750	001	Mar 17, 2015
+	!	250MG	N205750	002	Mar 17, 2015

CHOLINE C-11

INJECTABLE; INTRAVENOUS

CHOLINE C-11

<u>AP</u>	DECATUR	<u>4-33.1mCi/ML</u>	<u>A206319</u>	<u>001</u>	Nov 13, 2015
<u>AP</u>	+! MCPRF	<u>4-33.1mCi/ML</u>	<u>N203155</u>	<u>001</u>	Sep 12, 2012
<u>AP</u>	WA UNIV SCH MED	<u>4-33.1mCi/ML</u>	<u>A208413</u>	<u>001</u>	Jan 10, 2017
	UNIV TX MD ANDERSON	4-100mCi/ML	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>	ADAPTIS	<u>EQ 45MG FENOFIBRIC ACID</u>	<u>A211626</u>	<u>001</u>	Jul 18, 2019
<u>AB</u>		<u>EQ 135MG FENOFIBRIC ACID</u>	<u>A211626</u>	<u>002</u>	Jul 18, 2019
<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>	AUROBINDO PHARMA LTD	<u>EQ 45MG FENOFIBRIC ACID</u>	<u>A212598</u>	<u>001</u>	Jul 25, 2019
<u>AB</u>		<u>EQ 135MG FENOFIBRIC ACID</u>	<u>A212598</u>	<u>002</u>	Jul 25, 2019
<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>	MICRO LABS	<u>EQ 45MG FENOFIBRIC ACID</u>	<u>A213450</u>	<u>001</u>	Jun 16, 2020
<u>AB</u>		<u>EQ 135MG FENOFIBRIC ACID</u>	<u>A213450</u>	<u>002</u>	Jun 16, 2020
<u>AB</u>	TWI PHARMS	<u>EQ 45MG FENOFIBRIC ACID</u>	<u>A210469</u>	<u>001</u>	Jul 05, 2019
<u>AB</u>		<u>EQ 135MG FENOFIBRIC ACID</u>	<u>A210469</u>	<u>002</u>	Jul 05, 2019

PRESCRIPTION DRUG PRODUCT LIST

CHOLINE FENOFIBRATE

CAPSULE, DELAYED RELEASE;ORAL

FENOFIBRIC ACID

AB	YICHANG HUMANWELL	EQ 45MG FENOFIBRIC ACID	A212562 001	Dec 23, 2020
AB		EQ 135MG FENOFIBRIC ACID	A212562 002	Dec 23, 2020

TRILIPIX

AB	+ ABBVIE	EQ 45MG FENOFIBRIC ACID	N022224 001	Dec 15, 2008
AB	+!	EQ 135MG FENOFIBRIC ACID	N022224 002	Dec 15, 2008

CHROMIC CHLORIDE

INJECTABLE;INJECTION

CHROMIC CHLORIDE IN PLASTIC CONTAINER

+!	HOSPIRA	EQ 0.004MG CHROMIUM/ML	N018961 001	Jun 26, 1986
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CICLESONIDE

AEROSOL, METERED;INHALATION

ALVESCO

+!	COVIS	0.08MG/INH	N021658 002	Jan 10, 2008
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+!		0.16MG/INH	N021658 003	Jan 10, 2008
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AEROSOL, METERED;NASAL

ZETONNA

+!	COVIS	0.037MG/INH	N202129 001	Jan 20, 2012
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SPRAY, METERED;NASAL

OMNARIS

+!	COVIS	0.05MG/SPRAY	N022004 001	Oct 20, 2006
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CICLOPIROX

CREAM;TOPICAL

CICLOPIROX

AB	COSETTE	0.77%	A078463 001	Dec 20, 2010
AB	FOUGERA PHARMS	0.77%	A076435 001	Dec 29, 2004
AB	GLENMARK PHARMS	0.77%	A090273 001	Nov 10, 2009
AB	PADAGIS ISRAEL	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
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GEL;TOPICAL

CICLOPIROX

AB	+! ALVOGEN	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	PADAGIS US	0.77%	A078266 001	Jan 07, 2009

SHAMPOO;TOPICAL

CICLOPIROX

AT	ACTAVIS MID	1%	A090490 001	Nov 24, 2009
	ATLANTIC			
AT	FOUGERA PHARMS	1%	A090146 001	May 25, 2010
AT	PADAGIS US	1%	A078594 001	Feb 16, 2010
AT	TARO	1%	A090269 001	Feb 23, 2011
AT	TELLIGENT	1%	A209975 001	Apr 05, 2018

LOPROX

AT	+! BAUSCH	1%	N021159 001	Feb 28, 2003
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SOLUTION;TOPICAL

CICLOPIROX

AT	ACELLA	8%	A078172 001	Sep 18, 2007
AT	ACTAVIS MID	8%	A078046 001	Sep 18, 2007
	ATLANTIC			
AT	AKORN	8%	A078270 001	Sep 18, 2007
AT	COSETTE	8%	A078233 001	Sep 18, 2007
AT	PADAGIS US	8%	A077623 001	Sep 18, 2007
AT	RISING	8%	A078124 001	Sep 18, 2007
AT	TARO PHARM INDS	8%	A078144 001	Sep 18, 2007

PENLAC

AT	+! VALEANT BERMUDA	8%	N021022 001	Dec 17, 1999
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SUSPENSION;TOPICAL

CICLOPIROX

AB	FOUGERA PHARMS	0.77%	A076422 001	Aug 06, 2004
AB	PADAGIS ISRAEL	0.77%	A077676 001	Dec 15, 2006
AB	TARO	0.77%	A077092 001	Aug 10, 2005

LOPROX

AB	+! MEDIMETRIKS PHARMS	0.77%	N019824 001	Dec 30, 1988
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PRESCRIPTION DRUG PRODUCT LIST

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	HQ SPCLT PHARMA	EQ 500MG BASE/VIAL; 500MG/VIAL	A207594 001	Dec 12, 2019
AP	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

CILASTATIN SODIUM; IMPENEM; RELEBACTAM

POWDER; INTRAVENOUS

RECARBRIO

+! MSD MERCK CO

EQ 500MG

N212819 001 Jul 16, 2019

BASE/VIAL; 500MG/VIAL; 250MG/VIAL

CILOSTAZOL

TABLET; ORAL

CILOSTAZOL

AB	APOTEX INC	50MG	A077030 001	Dec 10, 2004
AB		100MG	A077030 002	Dec 10, 2004
AB	CASI PHARMS INC	50MG	A077310 001	Nov 08, 2005
AB		100MG	A077021 001	Nov 23, 2004
AB	CHARTWELL RX	50MG	A077831 002	Sep 24, 2012
AB		100MG	A077831 001	Sep 24, 2012
AB	HIKMA	50MG	A077024 001	May 17, 2005
AB		100MG	A077024 002	May 17, 2005
AB	SLATE	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

CIMETIDINE

TABLET; ORAL

CIMETIDINE

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

SOLUTION; ORAL

CIMETIDINE HYDROCHLORIDE

AA	! AKORN	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	ACCORD HLTHCARE	EQ 30MG BASE	A211892 001	May 15, 2020
AB		EQ 60MG BASE	A211892 002	May 15, 2020
AB		EQ 90MG BASE	A211892 003	May 15, 2020
AB	ACME LABS	EQ 30MG BASE	A213325 001	May 18, 2020
AB		EQ 60MG BASE	A213325 002	May 18, 2020
AB		EQ 90MG BASE	A213325 003	May 18, 2020
AB	ALKEM LABS LTD	EQ 30MG BASE	A210570 001	May 17, 2019
AB		EQ 60MG BASE	A210570 002	May 17, 2019
AB		EQ 90MG BASE	A210570 003	May 17, 2019
AB	AMNEAL PHARMS	30MG	A204364 001	Dec 02, 2021
AB		60MG	A204364 002	Dec 02, 2021
AB		90MG	A204364 003	Dec 02, 2021
AB	AUROBINDO PHARMA LTD	EQ 30MG BASE	A206125 001	Mar 08, 2018
AB		EQ 60MG BASE	A206125 002	Mar 08, 2018

PRESCRIPTION DRUG PRODUCT LIST

CINACALCET HYDROCHLORIDE

TABLET; ORAL

CINACALCET HYDROCHLORIDE

<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A206125 003</u>	Mar 08, 2018
<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>	DR REDDYS	<u>EQ 30MG BASE</u>	<u>A208368 001</u>	Sep 18, 2020
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A208368 002</u>	Sep 18, 2020
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A208368 003</u>	Sep 18, 2020
<u>AB</u>	HETERO LABS LTD V	<u>EQ 30MG BASE</u>	<u>A209403 001</u>	Oct 07, 2020
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A209403 002</u>	Oct 07, 2020
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A209403 003</u>	Oct 07, 2020
<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 PHARM	<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

+	BAYER HLTHCARE	250MG/5ML	N020780 001	Sep 26, 1997
+	!	500MG/5ML	N020780 002	Sep 26, 1997

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>CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
<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

+	!	ALK ABELLO	6% (60MG/ML)	N207986 001	Dec 10, 2015
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CIPROFLOXACIN HYDROCHLORIDE

OINTMENT; OPHTHALMIC

CILOXAN

+	!	NOVARTIS	EQ 0.3% BASE	N020369 001	Mar 30, 1998
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SOLUTION/DROPS; OPHTHALMIC

CILOXAN

<u>AT</u>	+	!	NOVARTIS	<u>EQ 0.3% BASE</u>	<u>N019992 001</u>	Dec 31, 1990
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CIPROFLOXACIN HYDROCHLORIDE

<u>AT</u>		AKORN	<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	<u>EQ 0.3% BASE</u>	<u>A077689 001</u>	Dec 13, 2006
<u>AT</u>		TELIGENT	<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

SOLUTION/DROPS; OTIC

CETRALAX

+	!	WRASER PHARMS	EQ 0.2% BASE	N021918 001	May 01, 2009
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PRESCRIPTION DRUG PRODUCT LIST

CIPROFLOXACIN HYDROCHLORIDE

TABLET; ORAL

CIPRO

<u>AB</u>	+	BAYER HLTHCARE	<u>EQ 250MG BASE</u>	<u>N019537 002</u>	Oct 22, 1987
<u>AB</u>	+	!	<u>EQ 500MG BASE</u>	<u>N019537 003</u>	Oct 22, 1987

CIPROFLOXACIN HYDROCHLORIDE

<u>AB</u>		AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A077859 001</u>	Apr 26, 2007
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A077859 002</u>	Apr 26, 2007
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A077859 003</u>	Apr 26, 2007
<u>AB</u>		CARLSBAD	<u>EQ 250MG BASE</u>	<u>A076126 002</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076126 003</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076126 004</u>	Jun 09, 2004
<u>AB</u>		CHARTWELL	<u>EQ 250MG BASE</u>	<u>A076896 001</u>	Nov 04, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076896 002</u>	Nov 04, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076896 003</u>	Nov 04, 2004
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 100MG BASE</u>	<u>A075593 002</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 250MG BASE</u>	<u>A075593 003</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A075593 004</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A075593 001</u>	Jun 09, 2004
<u>AB</u>		HIKMA	<u>EQ 250MG BASE</u>	<u>A076558 002</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076558 003</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076558 004</u>	Jun 09, 2004
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>EQ 250MG BASE</u>	<u>A076089 002</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076089 003</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076089 004</u>	Jun 09, 2004
<u>AB</u>		RISING PHARMA	<u>EQ 500MG BASE</u>	<u>A075817 003</u>	Jun 09, 2004
<u>AB</u>		TARO	<u>EQ 100MG BASE</u>	<u>A076912 001</u>	Feb 18, 2005
<u>AB</u>			<u>EQ 250MG BASE</u>	<u>A076912 002</u>	Oct 06, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076912 003</u>	Oct 06, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076912 004</u>	Oct 06, 2004
<u>AB</u>		UNIQUE PHARM LABS	<u>EQ 250MG BASE</u>	<u>A076639 001</u>	Sep 10, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076639 002</u>	Sep 10, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076639 003</u>	Sep 10, 2004
<u>AB</u>		WATSON LABS	<u>EQ 100MG BASE</u>	<u>A076794 001</u>	Feb 10, 2005
<u>AB</u>			<u>EQ 250MG BASE</u>	<u>A076794 002</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076794 003</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076794 004</u>	Jun 09, 2004
<u>AB</u>		YILING	<u>EQ 250MG BASE</u>	<u>A208921 001</u>	Jun 22, 2018
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A208921 002</u>	Jun 22, 2018

CIPROFLOXACIN HYDROCHLORIDE; FLUOCINOLONE ACETONIDE

SOLUTION/DROPS; OTIC

OTOVEL

+	!	LABORATORIOS SALVAT	<u>EQ 0.3% BASE;0.025%</u>	<u>N208251 001</u>	Apr 29, 2016
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CIPROFLOXACIN HYDROCHLORIDE; HYDROCORTISONE

SUSPENSION/DROPS; OTIC

CIPRO HC

+	!	NOVARTIS	<u>EQ 0.2% BASE;1%</u>	<u>N020805 001</u>	Feb 10, 1998
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CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CIPROFLOXACIN EXTENDED RELEASE

<u>AB</u>	!	ANCHEN PHARMS	<u>425.2MG;EQ 574.9MG BASE</u>	<u>A078166 001</u>	Nov 27, 2007
<u>AB</u>		DR REDDYS LABS LTD	<u>425.2MG;EQ 574.9MG BASE</u>	<u>A077701 001</u>	Mar 26, 2007
	!	ANCHEN PHARMS	<u>212.6MG;EQ 287.5MG BASE</u>	<u>A078166 002</u>	Nov 27, 2007

CIPROFLOXACIN; DEXAMETHASONE

SUSPENSION/DROPS; OTIC

CIPRODEX

<u>AB</u>	+	!	NOVARTIS	<u>0.3%;0.1%</u>	<u>N021537 001</u>	Jul 18, 2003
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CIPROFLOXACIN AND DEXAMETHASONE

<u>AB</u>		DR REDDYS LABS LTD	<u>0.3%;0.1%</u>	<u>A205548 001</u>	Aug 10, 2020
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CISATRACURIUM BESYLATE

INJECTABLE; INJECTION

CISATRACURIUM BESYLATE

<u>AP</u>		EUGIA PHARMA	<u>EQ 2MG BASE/ML</u>	<u>A209144 001</u>	May 08, 2020
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A203183 001</u>	Feb 26, 2015
<u>AP</u>		HOSPIRA INC	<u>EQ 2MG BASE/ML</u>	<u>A203236 001</u>	Mar 30, 2018
<u>AP</u>			<u>EQ 2MG BASE/ML</u>	<u>A203238 001</u>	Mar 30, 2018
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A203236 002</u>	Mar 30, 2018
<u>AP</u>		JIANGSU PHARMS	<u>EQ 2MG BASE/ML</u>	<u>A209334 001</u>	Aug 30, 2017
<u>AP</u>		MEITHEAL	<u>EQ 2MG BASE/ML</u>	<u>A211668 001</u>	Apr 25, 2019

PRESCRIPTION DRUG PRODUCT LIST

CISATRACURIUM BESYLATE

INJECTABLE; INJECTION

CISATRACURIUM BESYLATE

<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A211669 001</u>	Apr 25, 2019
<u>AP</u>		<u>EQ 10MG BASE/ML</u>	<u>A211668 002</u>	Apr 25, 2019
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 2MG BASE/ML</u>	<u>A201836 001</u>	Nov 06, 2020
<u>AP</u>		<u>EQ 10MG BASE/ML</u>	<u>A201836 002</u>	Nov 06, 2020
<u>AP</u>	SANDOZ INC	<u>EQ 2MG BASE/ML</u>	<u>A200159 001</u>	Feb 03, 2012
<u>AP</u>	SOMERSET	<u>EQ 2MG BASE/ML</u>	<u>A209132 001</u>	Apr 24, 2019
<u>AP</u>	ZYDUS PHARMS	<u>EQ 2MG BASE/ML</u>	<u>A212171 001</u>	Nov 04, 2019
<u>AP</u>		<u>EQ 10MG BASE/ML</u>	<u>A212171 002</u>	Nov 04, 2019

CISATRACURIUM BESYLATE PRESERVATIVE FREE

<u>AP</u>	EUGIA PHARMA	<u>EQ 2MG BASE/ML</u>	<u>A209665 001</u>	Oct 27, 2020
<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 PHARMS	<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
<u>AP</u>	SOMERSET THERAPS LLC	<u>EQ 2MG BASE/ML</u>	<u>A206791 001</u>	Feb 20, 2019
<u>AP</u>		<u>EQ 10MG BASE/ML</u>	<u>A206791 002</u>	Feb 20, 2019

NIMBEX

<u>AP</u>	+! ABBVIE	<u>EQ 2MG BASE/ML</u>	<u>N020551 001</u>	Dec 15, 1995
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NIMBEX PRESERVATIVE FREE

<u>AP</u>	+! ABBVIE	<u>EQ 2MG BASE/ML</u>	<u>N020551 003</u>	Dec 15, 1995
<u>AP</u>	+!	<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>	HIKMA	<u>1MG/ML</u>	<u>A075036 001</u>	Nov 07, 2000
<u>AP</u>	+ HQ SPCLT PHARMA	<u>1MG/ML</u>	<u>N018057 004</u>	Nov 08, 1988
<u>AP</u>	PHARMACHEMIE BV	<u>1MG/ML</u>	<u>A074656 001</u>	May 16, 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>	! HIKMA	<u>EQ 10MG BASE/5ML</u>	<u>A077043 001</u>	Dec 13, 2004
<u>AA</u>	LANNETT CO INC	<u>EQ 10MG BASE/5ML</u>	<u>A077629 001</u>	Jun 15, 2006

TABLET; ORAL

CELEXA

<u>AB</u>	+ ALLERGAN	<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>	COSETTE	<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>	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>	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

PRESCRIPTION DRUG PRODUCT LIST

CITALOPRAM HYDROBROMIDE

TABLET; ORAL

CITALOPRAM HYDROBROMIDE

<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
<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>	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; LACTIC ACID; POTASSIUM BITARTRATE

GEL; VAGINAL

PHEXXI

+! EVOFEM INC 1%; 1.8%; 0.4% N208352 001 May 22, 2020

CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE

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 4GM; 75MG 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>	HIKMA	<u>1MG/ML</u>	<u>A075405 001</u>	Feb 28, 2000
<u>AP</u>	HISUN PHARM HANGZHOU	<u>1MG/ML</u>	<u>A210856 001</u>	Nov 25, 2019
<u>AP</u>	MYLAN LABS LTD	<u>1MG/ML</u>	<u>A200510 001</u>	Oct 06, 2011
TABLET; ORAL				
MAVENCLAD				
	+! EMD SERONO INC	10MG	N022561 001	Mar 29, 2019

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>	! 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>	CHARTWELL	<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>	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>	STRIDES PHARMA	<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>	TEVA	<u>250MG</u>	<u>A065155 001</u>	May 31, 2005
<u>AB</u>		<u>500MG</u>	<u>A065155 002</u>	May 31, 2005

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>	! MAYNE PHARMA	<u>500MG</u>	<u>A065154 001</u>	May 18, 2005
<u>AB</u>	SUNSHINE	<u>500MG</u>	<u>A208987 001</u>	Jul 09, 2018

PRESCRIPTION DRUG PRODUCT LIST

CLASCOTERONE

CREAM; TOPICAL

WINLEVI

+! SUN PHARM

1%

N213433 001 Aug 26, 2020

CLEMASTINE FUMARATE

SYRUP; ORAL

CLEMASTINE FUMARATE

! GENUS

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

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** + PFIZER**EQ 75MG BASE****N050162 001****AB** +**EQ 150MG BASE****N050162 002****AB** +!**EQ 300MG BASE****N050162 003** Apr 14, 1988CLINDAMYCIN HYDROCHLORIDE**AB** AUROBINDO PHARMA**EQ 150MG BASE****A065442 001** Aug 26, 2009**AB****EQ 300MG BASE****A065442 002** Aug 26, 2009**AB** COSETTE**EQ 150MG BASE****A063029 001** Sep 20, 1989**AB****EQ 300MG BASE****A063029 002** Aug 05, 2005**AB** EPIC PHARMA LLC**EQ 150MG BASE****A065194 001** Mar 22, 2004**AB****EQ 300MG BASE****A065194 002** Mar 22, 2004**AB** LANNETT CO INC**EQ 75MG BASE****A065243 002** 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, 2005CLINDAMYCIN PALMITATE HYDROCHLORIDE

FOR SOLUTION; ORAL

CLEOCIN**AA** ! PFIZER**EQ 75MG BASE/5ML****A062644 001** Apr 07, 1986CLINDAMYCIN PALMITATE HYDROCHLORIDE**AA** ALEMBIC LABS**EQ 75MG BASE/5ML****A206958 001** May 05, 2017**AA** AMNEAL PHARMS**EQ 75MG BASE/5ML****A203513 001** Mar 13, 2014**AA** AUROBINDO PHARMA**EQ 75MG BASE/5ML****A202409 001** Apr 30, 2013**AA** LTD**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** PADAGIS US**EQ 75MG BASE/5ML****A090902 001** Jul 07, 2010CLINDAMYCIN PHOSPHATE

AEROSOL, FOAM; TOPICAL

CLINDAMYCIN PHOSPHATE**AT** GLENMARK PHARMS LTD**1%****A210778 001** Sep 20, 2021**AT** PADAGIS ISRAEL**1%****A090785 001** Mar 31, 2010EVOCLIN**AT** +! MYLAN**1%****N050801 001** Oct 22, 2004

CREAM; VAGINAL

CLEOCIN**AB** +! PFIZER**EQ 2% BASE****N050680 002** Mar 02, 1998CLINDAMYCIN PHOSPHATE**AB** FOUGERA PHARMS**EQ 2% BASE****A065139 001** Dec 27, 2004

CLINDESSE

+! PADAGIS US

EQ 2% BASE

N050793 001 Nov 30, 2004

GEL; TOPICAL

CLEOCIN T**AB1** +! PFIZER**EQ 1% BASE****N050615 001** Jan 07, 1987

PRESCRIPTION DRUG PRODUCT LIST

CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

CLINDAMYCIN PHOSPHATE

<u>AB1</u>	ENCUBE	<u>EQ 1% BASE</u>	<u>A212438 001</u>	Mar 11, 2021
<u>AB1</u>	FOUGERA PHARMS	<u>EQ 1% BASE</u>	<u>A064160 001</u>	Jan 28, 2000
<u>AB1</u>	GLENMARK PHARMS LTD	<u>EQ 1% BASE</u>	<u>A214251 001</u>	Feb 10, 2021
<u>AB1</u>	PADAGIS ISRAEL	<u>EQ 1% BASE</u>	<u>A212104 001</u>	Dec 31, 2020
<u>AB1</u>	SOLARIS PHARMA CORP	<u>EQ 1% BASE</u>	<u>A211872 001</u>	Jul 29, 2020
<u>AB1</u>	TARO PHARMS	<u>EQ 1% BASE</u>	<u>A214052 001</u>	Nov 10, 2020

CLINDAGEL

<u>AB2</u>	+! BAUSCH	<u>EQ 1% BASE</u>	<u>N050782 001</u>	Nov 27, 2000
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CLINDAMYCIN PHOSPHATE

<u>AB2</u>	SOLARIS PHARMA CORP	<u>EQ 1% BASE</u>	<u>A212842 001</u>	Aug 13, 2021
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GEL; VAGINAL

XACIATO

+!	DARE	EQ 2% BASE	N215650 001	Dec 07, 2021
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INJECTABLE; INJECTION

CLEOCIN PHOSPHATE

<u>AP</u>	PFIZER	<u>EQ 150MG BASE/ML</u>	<u>A062803 001</u>	Oct 16, 1987
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<u>AP</u>	+!	<u>EQ 150MG BASE/ML</u>	<u>N050441 001</u>	
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CLINDAMYCIN PHOSPHATE

<u>AP</u>	ALMAJECT	<u>EQ 150MG BASE/ML</u>	<u>A062800 001</u>	Jul 24, 1987
<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>A062943 001</u>	Sep 29, 1988
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 150MG BASE/ML</u>	<u>A065346 001</u>	Mar 29, 2007
<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>A065347 001</u>	May 09, 2007
<u>AP</u>	HIKMA	<u>EQ 150MG BASE/ML</u>	<u>A062889 001</u>	Apr 25, 1988
<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>A065206 001</u>	Sep 24, 2004
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 150MG BASE/ML</u>	<u>A090108 001</u>	Sep 30, 2011
<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>A090109 001</u>	Sep 30, 2011

CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER

<u>AP</u>	AKORN	<u>EQ 6MG BASE/ML</u>	<u>A203048 001</u>	Apr 04, 2013
<u>AP</u>		<u>EQ 12MG BASE/ML</u>	<u>A203048 002</u>	Apr 04, 2013
<u>AP</u>		<u>EQ 18MG BASE/ML</u>	<u>A203048 003</u>	Apr 04, 2013
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 6MG BASE/ML</u>	<u>A208084 001</u>	Jun 28, 2017
<u>AP</u>		<u>EQ 12MG BASE/ML</u>	<u>A208084 002</u>	Jun 28, 2017
<u>AP</u>		<u>EQ 18MG BASE/ML</u>	<u>A208084 003</u>	Jun 28, 2017
<u>AP</u>	! SANDOZ INC	<u>EQ 6MG BASE/ML</u>	<u>A201692 001</u>	May 31, 2012
<u>AP</u>	!	<u>EQ 12MG BASE/ML</u>	<u>A201692 002</u>	May 31, 2012
<u>AP</u>	!	<u>EQ 18MG BASE/ML</u>	<u>A201692 003</u>	May 31, 2012

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%

+!	ABRAXIS PHARM	EQ 900MG BASE/100ML	N050635 001	Dec 22, 1989
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LOTION; TOPICAL

CLEOCIN T

<u>AB</u>	+! PFIZER	<u>EQ 1% BASE</u>	<u>N050600 001</u>	May 31, 1989
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CLINDAMYCIN PHOSPHATE

<u>AB</u>	FOUGERA PHARMS	<u>EQ 1% BASE</u>	<u>A065067 001</u>	Jan 31, 2002
<u>AB</u>	PADAGIS ISRAEL	<u>EQ 1% BASE</u>	<u>A214604 001</u>	Mar 08, 2021
<u>AB</u>	TARO PHARMS	<u>EQ 1% BASE</u>	<u>A214526 001</u>	Jun 28, 2021

SOLUTION; INTRAVENOUS

CLINDAMYCIN PHOSPHATE IN 0.9% SODIUM CHLORIDE

+!	BAXTER HLTHCARE CORP	EQ 300MG BASE/50ML (EQ 6MG BASE/ML)	N208083 001	Apr 20, 2017
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+!		EQ 600MG BASE/50ML (EQ 12MG BASE/ML)	N208083 002	Apr 20, 2017
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+!		EQ 900MG BASE/50ML (EQ 18MG BASE/ML)	N208083 003	Apr 20, 2017
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SOLUTION; TOPICAL

CLINDA-DERM

<u>AT</u>	PADAGIS US	<u>EQ 1% BASE</u>	<u>A063329 001</u>	Sep 30, 1992
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CLINDAMYCIN PHOSPHATE

<u>AT</u>	CADILA	<u>EQ 1% BASE</u>	<u>A208767 001</u>	Jul 16, 2018
<u>AT</u>	ENCUBE ETHICALS	<u>EQ 1% BASE</u>	<u>A209914 001</u>	Jan 28, 2019
<u>AT</u>	FOUGERA PHARMS INC	<u>EQ 1% BASE</u>	<u>A064159 001</u>	Jun 05, 1997
<u>AT</u>	GLASSHOUSE PHARMS	<u>EQ 1% BASE</u>	<u>A209846 001</u>	Feb 08, 2018
<u>AT</u>	! PADAGIS US	<u>EQ 1% BASE</u>	<u>A064050 001</u>	Nov 30, 1995
<u>AT</u>	TARO PHARM INDS	<u>EQ 1% BASE</u>	<u>A065184 001</u>	Mar 31, 2004
<u>AT</u>	TELIGENT	<u>EQ 1% BASE</u>	<u>A206945 001</u>	Dec 30, 2016

SUPPOSITORY; VAGINAL

CLEOCIN

+!	PFIZER	100MG	N050767 001	Aug 13, 1999
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SWAB; TOPICAL

CLINDAMYCIN PHOSPHATE

<u>AT</u>	AKORN	<u>EQ 1% BASE</u>	<u>A065513 001</u>	Jun 17, 2010
<u>AT</u>	! PADAGIS US	<u>EQ 1% BASE</u>	<u>A065049 001</u>	May 25, 2000

PRESCRIPTION DRUG PRODUCT LIST

CLINDAMYCIN PHOSPHATE

SWAB; TOPICAL

CLINDETS

AT	PADAGIS US	EQ 1% BASE	A064136 001	Sep 30, 1996
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CLINDAMYCIN PHOSPHATE; TRETINOIN

GEL; TOPICAL

CLINDAMYCIN PHOSPHATE AND TRETINOIN

AB	ACTAVIS MID ATLANTIC	1.2%;0.025%	A202564 001	Jun 12, 2015
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ZIANA

AB	+! BAUSCH	1.2%;0.025%	N050802 001	Nov 07, 2006
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VELTIN

BX	+! ALMIRALL	1.2%;0.025%	N050803 001	Jul 16, 2010
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CLOBAZAM

FILM; ORAL

SYMPAZAN

+	AQUESTIVE	5MG	N210833 001	Nov 01, 2018
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+		10MG	N210833 002	Nov 01, 2018
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+		20MG	N210833 003	Nov 01, 2018
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SUSPENSION; ORAL

CLOBAZAM

AB	ALKEM LABS LTD	2.5MG/ML	A213039 001	May 06, 2021
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AB	AMNEAL	2.5MG/ML	A210039 001	Oct 22, 2018
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AB	BIONPHARMA INC	2.5MG/ML	A208819 001	Oct 22, 2018
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AB	HETERO LABS LTD III	2.5MG/ML	A209796 001	Feb 24, 2020
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AB	HIKMA	2.5MG/ML	A209715 001	Oct 22, 2018
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AB	LANNETT CO INC	2.5MG/ML	A213110 001	Apr 24, 2020
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AB	LUPIN LTD	2.5MG/ML	A210546 001	Dec 28, 2018
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AB	MYLAN	2.5MG/ML	A211259 001	Oct 22, 2018
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AB	TARO	2.5MG/ML	A210978 001	Apr 15, 2019
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AB	UPSHER SMITH LABS	2.5MG/ML	A210569 001	Oct 22, 2018
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AB	VISTAPHARM	2.5MG/ML	A210746 001	Jul 10, 2019
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ONFI

AB	+! LUNDBECK PHARMS LLC	2.5MG/ML	N203993 001	Dec 14, 2012
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TABLET; ORAL

CLOBAZAM

AB	ALKEM LABS LTD	10MG	A212714 001	Sep 06, 2019
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AB		20MG	A212714 002	Sep 06, 2019
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AB	AMNEAL PHARMS CO	10MG	A209718 001	Oct 22, 2018
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AB		20MG	A209718 002	Oct 22, 2018
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AB	BIONPHARMA INC	10MG	A208825 001	Oct 22, 2018
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AB		20MG	A208825 002	Oct 22, 2018
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AB	BRECKENRIDGE	10MG	A209308 001	Oct 22, 2018
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AB		20MG	A209308 002	Oct 22, 2018
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AB	HETERO LABS LTD III	10MG	A209795 001	Oct 22, 2018
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AB		20MG	A209795 002	Oct 22, 2018
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AB	HIKMA	10MG	A208785 001	Oct 22, 2018
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AB		20MG	A208785 002	Oct 22, 2018
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AB	LUPIN LTD	10MG	A210545 001	Dec 14, 2018
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AB		20MG	A210545 002	Dec 14, 2018
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AB	MICRO LABS	10MG	A211711 001	Jan 30, 2019
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AB		20MG	A211711 002	Jan 30, 2019
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AB	PIRAMAL HLTHCARE UK	10MG	A209808 001	Oct 22, 2018
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AB		20MG	A209808 002	Oct 22, 2018
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AB	UPSHER SMITH LABS	10MG	A209687 001	Oct 22, 2018
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AB		20MG	A209687 002	Oct 22, 2018
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AB	ZYDUS PHARMS	10MG	A211449 001	Oct 22, 2018
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AB		20MG	A211449 002	Oct 22, 2018
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ONFI

AB	+ LUNDBECK PHARMS LLC	10MG	N202067 002	Oct 21, 2011
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AB	+!	20MG	N202067 003	Oct 21, 2011
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CLOBAZAM

	ALKEM LABS LTD	5MG	A212714 003	Dec 21, 2020
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CLOBETASOL PROPIONATE

AEROSOL, FOAM; TOPICAL

CLOBETASOL PROPIONATE

AB1	GLENMARK PHARMS LTD	0.05%	A210809 001	Feb 15, 2019
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AB1	PADAGIS ISRAEL	0.05%	A077763 001	Mar 10, 2008
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AB1	TARO	0.05%	A208779 001	Oct 04, 2018
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OLUX

AB1	+! MYLAN	0.05%	N021142 001	May 26, 2000
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PRESCRIPTION DRUG PRODUCT LIST

CLOBETASOL PROPIONATE

AEROSOL, FOAM;TOPICAL

CLOBETASOL PROPIONATE

AB2	GLENMARK PHARMS LTD	0.05%	A211450 001	Sep 09, 2019
AB2	PADAGIS ISRAEL	0.05%	A201402 001	Aug 14, 2012

OLUX E

AB2	+! MYLAN	0.05%	N022013 001	Jan 12, 2007
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CREAM;TOPICAL

CLOBETASOL PROPIONATE

AB1	ALEOR DERMACEUTICALS	0.05%	A213291 001	Jan 27, 2020
AB1	AMNEAL	0.05%	A211256 001	Dec 26, 2018
AB1	AUROBINDO PHARMA USA	0.05%	A075338 001	Feb 09, 2001
AB1	COSETTE	0.05%	A074139 001	Aug 03, 1994
AB1	ENCUBE	0.05%	A212982 001	Aug 28, 2020
AB1	FOUGERA PHARMS INC	0.05%	A074392 001	Sep 30, 1996
AB1	GLENMARK PHARMS	0.05%	A209095 001	May 10, 2018
AB1	LUPIN LTD	0.05%	A210208 001	Jan 30, 2018
AB1	RISING	0.05%	A211401 001	Jan 11, 2019
AB1	TARO	0.05%	A074249 001	Jul 08, 1996
AB1	TELIGENT	0.05%	A209974 001	Apr 17, 2018
AB1	TORRENT	0.05%	A211836 001	Dec 30, 2019
AB1	XIROMED	0.05%	A210034 001	Jun 15, 2018
AB1	ZYDUS PHARMS	0.05%	A211074 001	Oct 15, 2018

CORMAX

AB1	! AKORN	0.05%	A074220 001	May 16, 1997
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CLOBETASOL PROPIONATE (EMOLLIENT)

AB2	ANI PHARMS	0.05%	A075733 001	Aug 22, 2001
AB2	! FOUGERA PHARMS	0.05%	A075430 001	May 26, 1999
AB2	TARO	0.05%	A075633 001	May 17, 2000
AB2	TELIGENT	0.05%	A209411 001	Aug 21, 2017

EMBELINE E

AB2	AKORN	0.05%	A075325 001	Dec 24, 1998
	IMPOYZ			
	+! PRIMUS PHARMS	0.025%	N209483 001	Nov 28, 2017

GEL;TOPICAL

CLOBETASOL PROPIONATE

AB	! FOUGERA PHARMS	0.05%	A075368 001	Feb 15, 2000
AB	PADAGIS US	0.05%	A075027 001	Oct 31, 1997
AB	TARO	0.05%	A075279 001	May 28, 1999
AB	TELIGENT	0.05%	A208881 001	Mar 06, 2017

EMBELINE

AB	AKORN	0.05%	A076141 001	Apr 12, 2002
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LOTION;TOPICAL

CLOBETASOL PROPIONATE

AB	ACTAVIS MID ATLANTIC	0.05%	A078223 001	Dec 04, 2008
AB	CADILA	0.05%	A205249 001	Sep 24, 2019
AB	LUPIN LTD	0.05%	A209147 001	Sep 22, 2017
AB	TARO	0.05%	A200302 001	Jul 02, 2012
AB	TELIGENT	0.05%	A208667 001	Nov 29, 2016

CLOBEX

AB	+! GALDERMA LABS LP	0.05%	N021535 001	Jul 24, 2003
	IMPEKLO			
	+! MYLAN	0.05%	N213691 001	May 19, 2020

OINTMENT;TOPICAL

CLOBETASOL PROPIONATE

AB	ALEOR DERMACEUTICALS	0.05%	A211800 001	Mar 04, 2019
AB	AUROBINDO PHARMA USA	0.05%	A075057 001	Aug 12, 1998
AB	COSETTE	0.05%	A074089 001	Feb 16, 1994
AB	ENCUBE	0.05%	A211295 001	Nov 15, 2019
AB	! FOUGERA PHARMS	0.05%	A074407 001	Feb 23, 1996
AB	GLENMARK PHARMS	0.05%	A208933 001	Mar 20, 2017
AB	NOVEL LABS INC	0.05%	A208841 001	May 04, 2018
AB	TARO	0.05%	A074248 001	Jul 12, 1996
AB	TELIGENT	0.05%	A208589 001	Jan 23, 2019
AB	XIROMED	0.05%	A209701 001	Apr 17, 2018
AB	ZYDUS PHARMS	0.05%	A210199 001	Oct 27, 2017

EMBELINE

AB	AKORN	0.05%	A074221 001	Mar 31, 1995
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PRESCRIPTION DRUG PRODUCT LIST

CLOBETASOL PROPIONATE

SHAMPOO; TOPICAL

CLOBETASOL PROPIONATE

AB	ACTAVIS MID ATLANTIC	0.05%	A078854 001	Jun 07, 2011
AB	AKORN	0.05%	A209871 001	Oct 27, 2017
AB	ALEOR DERMACEUTICALS	0.05%	A213290 001	May 18, 2020
AB	AMNEAL	0.05%	A214895 001	Jun 14, 2021
AB	PADAGIS ISRAEL	0.05%	A090974 001	Aug 09, 2012
AB	TARO PHARMS	0.05%	A214867 001	Mar 25, 2021

CLOBEX

AB	+! GALDERMA LABS SOLUTION; TOPICAL	0.05%	N021644 001	Feb 05, 2004
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CLOBETASOL PROPIONATE

AT	ALEOR DERMACEUTICALS	0.05%	A212881 001	Oct 21, 2019
AT	COSETTE	0.05%	A074331 001	Dec 15, 1995
AT	FOUGERA PHARMS	0.05%	A075391 001	Feb 08, 1999
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	PRINSTON INC	0.05%	A213139 001	Feb 08, 2021
AT	SAPTALIS PHARMS	0.05%	A211494 001	Oct 02, 2019
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	WOCKHARDT BIO AG	0.05%	A075205 001	Nov 13, 1998

EMBELINE

AT	! AKORN SPRAY; TOPICAL	0.05%	A074222 001	Dec 06, 1995
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CLOBETASOL PROPIONATE

AT	ALEOR DERMACEUTICALS	0.05%	A211191 001	Oct 02, 2019
AT	GLENMARK PHARMS	0.05%	A209004 001	Mar 26, 2018
AT	LUPIN LTD	0.05%	A208125 001	Mar 26, 2018
AT	PADAGIS US	0.05%	A090898 001	Jun 16, 2011
AT	TARO	0.05%	A208842 001	Mar 26, 2018
AT	ZYDUS PHARMS	0.05%	A206378 001	Feb 16, 2017

CLOBEX

AT	+! GALDERMA LABS LP	0.05%	N021835 001	Oct 27, 2005
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CLOCORTOLONE PIVALATE

CREAM; TOPICAL

CLOCORTOLONE PIVALATE

AB	TARO	0.1%	A206370 001	Apr 21, 2020
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CLODERM

AB	+! EPI HLTH	0.1%	N017765 001	
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CLOFARABINE

SOLUTION; INTRAVENOUS

CLOFARABINE

AP	ABON PHARMS LLC	20MG/20ML (1MG/ML)	A204029 001	May 09, 2017
AP	ACCORD HLTHCARE	20MG/20ML (1MG/ML)	A212034 001	Feb 22, 2019
AP	AMNEAL	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	MEITHEAL	20MG/20ML (1MG/ML)	A213461 001	Oct 23, 2020
AP	MSN	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
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CLOMIPHENE CITRATE

TABLET; ORAL

CLOMIPHENE CITRATE

!	PAR PHARM	50MG	A075528 001	Aug 30, 1999
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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

PRESCRIPTION DRUG PRODUCT LIST

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

CLOMIPRAMINE HYDROCHLORIDE

<u>AB</u>	AJANTA PHARMA LTD	<u>25MG</u>	<u>A213897 001</u>	Oct 02, 2020
<u>AB</u>		<u>50MG</u>	<u>A213897 002</u>	Oct 02, 2020
<u>AB</u>		<u>75MG</u>	<u>A213897 003</u>	Oct 02, 2020
<u>AB</u>	ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A211822 001</u>	Aug 04, 2021
<u>AB</u>		<u>50MG</u>	<u>A211822 002</u>	Aug 04, 2021
<u>AB</u>		<u>75MG</u>	<u>A211822 003</u>	Aug 04, 2021
<u>AB</u>	AMNEAL PHARMS CO	<u>25MG</u>	<u>A208632 001</u>	Oct 31, 2018
<u>AB</u>		<u>50MG</u>	<u>A208632 002</u>	Oct 31, 2018
<u>AB</u>		<u>75MG</u>	<u>A208632 003</u>	Oct 31, 2018
<u>AB</u>	JUBILANT CADISTA	<u>25MG</u>	<u>A212218 001</u>	Oct 21, 2019
<u>AB</u>		<u>50MG</u>	<u>A212218 002</u>	Oct 21, 2019
<u>AB</u>		<u>75MG</u>	<u>A212218 003</u>	Oct 21, 2019
<u>AB</u>	LEADING PHARMA LLC	<u>25MG</u>	<u>A211364 001</u>	Feb 07, 2020
<u>AB</u>		<u>50MG</u>	<u>A211364 002</u>	Feb 07, 2020
<u>AB</u>		<u>75MG</u>	<u>A211364 003</u>	Feb 07, 2020
<u>AB</u>	LUPIN LTD	<u>25MG</u>	<u>A209294 001</u>	Nov 21, 2018
<u>AB</u>		<u>50MG</u>	<u>A209294 002</u>	Nov 21, 2018
<u>AB</u>		<u>75MG</u>	<u>A209294 003</u>	Nov 21, 2018
<u>AB</u>	MANKIND PHARMA	<u>25MG</u>	<u>A211767 001</u>	Apr 08, 2019
<u>AB</u>		<u>50MG</u>	<u>A211767 002</u>	Apr 08, 2019
<u>AB</u>		<u>75MG</u>	<u>A211767 003</u>	Apr 08, 2019
<u>AB</u>	MICRO LABS	<u>25MG</u>	<u>A213219 001</u>	Jun 22, 2020
<u>AB</u>		<u>50MG</u>	<u>A213219 002</u>	Jun 22, 2020
<u>AB</u>		<u>75MG</u>	<u>A213219 003</u>	Jun 22, 2020
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A074947 001</u>	Apr 30, 1998
<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>	TULEX PHARMS INC	<u>25MG</u>	<u>A210653 001</u>	Apr 03, 2020
<u>AB</u>		<u>50MG</u>	<u>A210653 002</u>	Apr 03, 2020
<u>AB</u>		<u>75MG</u>	<u>A210653 003</u>	Apr 03, 2020
<u>AB</u>	UNIQUE PHARM LABS	<u>25MG</u>	<u>A212285 001</u>	Aug 07, 2020
<u>AB</u>		<u>50MG</u>	<u>A212285 002</u>	Aug 07, 2020
<u>AB</u>		<u>75MG</u>	<u>A212285 003</u>	Aug 07, 2020
<u>AB</u>	VGYAAN	<u>25MG</u>	<u>A213221 001</u>	Jun 22, 2020
<u>AB</u>		<u>50MG</u>	<u>A213221 002</u>	Jun 22, 2020
<u>AB</u>		<u>75MG</u>	<u>A213221 003</u>	Jun 22, 2020
<u>AB</u>	ZYDUS PHARMS	<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>	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>	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>KLONOPIN</u>				
<u>AB</u>	+ CHEPLAPHARM	<u>0.5MG</u>	<u>N017533 001</u>	
<u>AB</u>	+!	<u>1MG</u>	<u>N017533 002</u>	

PRESCRIPTION DRUG PRODUCT LIST

CLONAZEPAM

TABLET; ORAL

KLONOPIN

<u>AB</u>	+	<u>2MG</u>	<u>N017533</u>	<u>003</u>	
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TABLET, ORALLY DISINTEGRATING; ORAL

CLONAZEPAM

<u>AB</u>	ALEMBIC PHARMS LTD	<u>0.125MG</u>	<u>A211033</u>	<u>001</u>	Jun 28, 2019
<u>AB</u>		<u>0.25MG</u>	<u>A211033</u>	<u>002</u>	Jun 28, 2019
<u>AB</u>		<u>0.5MG</u>	<u>A211033</u>	<u>003</u>	Jun 28, 2019
<u>AB</u>		<u>1MG</u>	<u>A211033</u>	<u>004</u>	Jun 28, 2019
<u>AB</u>		<u>2MG</u>	<u>A211033</u>	<u>005</u>	Jun 28, 2019
<u>AB</u>	BARR	<u>0.125MG</u>	<u>A077194</u>	<u>001</u>	Aug 10, 2005
<u>AB</u>		<u>0.25MG</u>	<u>A077194</u>	<u>002</u>	Aug 10, 2005
<u>AB</u>		<u>0.5MG</u>	<u>A077194</u>	<u>003</u>	Aug 10, 2005
<u>AB</u>		<u>1MG</u>	<u>A077194</u>	<u>004</u>	Aug 10, 2005
<u>AB</u>		<u>2MG</u>	<u>A077194</u>	<u>005</u>	Aug 10, 2005
<u>AB</u>	PAR PHARM	<u>0.125MG</u>	<u>A077171</u>	<u>001</u>	Aug 03, 2005
<u>AB</u>		<u>0.25MG</u>	<u>A077171</u>	<u>002</u>	Aug 03, 2005
<u>AB</u>		<u>0.5MG</u>	<u>A077171</u>	<u>003</u>	Aug 03, 2005
<u>AB</u>	!	<u>1MG</u>	<u>A077171</u>	<u>004</u>	Aug 03, 2005
<u>AB</u>		<u>2MG</u>	<u>A077171</u>	<u>005</u>	Aug 03, 2005
<u>AB</u>	SUN PHARM INDS INC	<u>0.125MG</u>	<u>A078654</u>	<u>001</u>	Aug 27, 2014
<u>AB</u>		<u>0.25MG</u>	<u>A078654</u>	<u>002</u>	Aug 27, 2014
<u>AB</u>		<u>0.5MG</u>	<u>A078654</u>	<u>003</u>	Aug 27, 2014
<u>AB</u>		<u>1MG</u>	<u>A078654</u>	<u>004</u>	Aug 27, 2014
<u>AB</u>		<u>2MG</u>	<u>A078654</u>	<u>005</u>	Aug 27, 2014

CLONIDINE

SYSTEM; TRANSDERMAL

CATAPRES-TTS-1

<u>AB</u>	+	LAVIPHARM	<u>0.1MG/24HR</u>	<u>N018891</u>	<u>001</u>	Oct 10, 1984
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CATAPRES-TTS-2

<u>AB</u>	+	LAVIPHARM	<u>0.2MG/24HR</u>	<u>N018891</u>	<u>002</u>	Oct 10, 1984
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CATAPRES-TTS-3

<u>AB</u>	+	!	LAVIPHARM	<u>0.3MG/24HR</u>	<u>N018891</u>	<u>003</u>	Oct 10, 1984
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CLONIDINE

<u>AB</u>	ACTAVIS LABS UT INC	<u>0.1MG/24HR</u>	<u>A090873</u>	<u>001</u>	May 06, 2014
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A090873</u>	<u>002</u>	May 06, 2014
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A090873</u>	<u>003</u>	May 06, 2014
<u>AB</u>	AVEVA	<u>0.1MG/24HR</u>	<u>A076157</u>	<u>001</u>	Aug 18, 2009
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A076157</u>	<u>002</u>	Aug 18, 2009
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A076157</u>	<u>003</u>	Aug 18, 2009
<u>AB</u>	MAYNE PHARMA	<u>0.1MG/24HR</u>	<u>A079090</u>	<u>001</u>	Aug 20, 2010
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A079090</u>	<u>002</u>	Aug 20, 2010
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A079090</u>	<u>003</u>	Aug 20, 2010
<u>AB</u>	MYLAN TECHNOLOGIES	<u>0.1MG/24HR</u>	<u>A076166</u>	<u>001</u>	Jul 16, 2010
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A076166</u>	<u>002</u>	Jul 16, 2010
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A076166</u>	<u>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</u>	<u>001</u>	Jul 08, 2011
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A200673</u>	<u>002</u>	Jul 08, 2011
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/10ML (0.1MG/ML)</u>	<u>A200300</u>	<u>001</u>	Jan 26, 2011
<u>AP</u>	!	<u>5MG/10ML (0.5MG/ML)</u>	<u>A200300</u>	<u>002</u>	Jan 26, 2011
<u>AP</u>	XGEN PHARMS	<u>1MG/10ML (0.1MG/ML)</u>	<u>A203167</u>	<u>001</u>	Oct 29, 2013
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A203167</u>	<u>002</u>	Oct 29, 2013
<u>AP</u>	ZYDUS PHARMS	<u>1MG/10ML (0.1MG/ML)</u>	<u>A202601</u>	<u>001</u>	Feb 20, 2014
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A202601</u>	<u>002</u>	Feb 20, 2014

DURACLON

<u>AP</u>	+	MYLAN INSTITUTIONAL	<u>1MG/10ML (0.1MG/ML)</u>	<u>N020615</u>	<u>001</u>	Oct 02, 1996
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TABLET; ORAL

CLONIDINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>0.1MG</u>	<u>A070974</u>	<u>001</u>	Dec 16, 1986
<u>AB</u>		<u>0.2MG</u>	<u>A070975</u>	<u>001</u>	Dec 16, 1986
<u>AB</u>		<u>0.3MG</u>	<u>A070976</u>	<u>001</u>	Dec 16, 1986
<u>AB</u>	ALEMBIC PHARMS LTD	<u>0.1MG</u>	<u>A091368</u>	<u>001</u>	Dec 06, 2011
<u>AB</u>		<u>0.2MG</u>	<u>A091368</u>	<u>002</u>	Dec 06, 2011
<u>AB</u>		<u>0.3MG</u>	<u>A091368</u>	<u>003</u>	Dec 06, 2011
<u>AB</u>	FRONTIDA BIOPHARM	<u>0.1MG</u>	<u>A070923</u>	<u>003</u>	Sep 04, 1987
<u>AB</u>		<u>0.2MG</u>	<u>A070923</u>	<u>002</u>	Sep 04, 1987
<u>AB</u>		<u>0.3MG</u>	<u>A070923</u>	<u>001</u>	Sep 04, 1987

PRESCRIPTION DRUG PRODUCT LIST

CLONIDINE HYDROCHLORIDE

TABLET;ORAL

CLONIDINE HYDROCHLORIDE

<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>	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>	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
<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>	NOVAST LABS	<u>0.1MG</u>	<u>A209675 001</u>	Mar 05, 2019
<u>AB1</u>	XIAMEN LP PHARM CO	<u>0.1MG</u>	<u>A209757 001</u>	Nov 20, 2017

KAPVAY

<u>AB1</u>	+!	CONCORDIA PHARMS INC	<u>0.1MG</u>	<u>N022331 003</u>	Sep 28, 2010
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CLONIDINE HYDROCHLORIDE

<u>AB2</u>	ACTAVIS ELIZABETH	<u>0.1MG</u>	<u>A202792 001</u>	May 15, 2015
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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>	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>	GRAVITI PHARMS	<u>EQ 75MG BASE</u>	<u>A204359 001</u>	Feb 02, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A202928 001</u>	Feb 10, 2014
<u>AB</u>	POLYGEN PHARMS	<u>EQ 75MG BASE</u>	<u>A213351 001</u>	Jul 17, 2020
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A213351 002</u>	Jul 17, 2020
<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	<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>	TORRENT PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A090844 001</u>	May 17, 2012

PLAVIX

<u>AB</u>	+	SANOFI AVENTIS US	<u>EQ 75MG BASE</u>	<u>N020839 001</u>	Nov 17, 1997
<u>AB</u>	+		<u>EQ 300MG BASE</u>	<u>N020839 002</u>	Sep 20, 2007

CLORAZEPATE DIPOTASSIUM

TABLET;ORAL

CLORAZEPATE DIPOTASSIUM

<u>AB</u>	AUROLIFE PHARMA LLC	<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>15MG</u>	<u>A071858 001</u>	Jul 17, 1987
<u>AB</u>	TARO	<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

TRANXENE

<u>AB</u>	+	RECORDATI RARE	<u>7.5MG</u>	<u>N017105 007</u>
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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>	NOVITIUM PHARMA	<u>1%</u>	<u>A209815</u>	<u>001</u>	Feb 14, 2019
<u>AT</u>	! TARO	<u>1%</u>	<u>A074580</u>	<u>001</u>	Jul 29, 1996
<u>AT</u>	TASMAN PHARMA	<u>1%</u>	<u>A212281</u>	<u>001</u>	Jul 25, 2019
<u>AT</u>	TEVA	<u>1%</u>	<u>A073306</u>	<u>001</u>	Feb 28, 1995

TROCHE/LOZENGE; ORAL

CLOTRIMAZOLE

<u>AB</u>	! HIKMA	<u>10MG</u>	<u>A076387</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>	PADAGIS US	<u>10MG</u>	<u>A076763</u>	<u>001</u>	Oct 28, 2005

CLOZAPINE

SUSPENSION; ORAL

VERSACLOZ

+	! TASMAN PHARMA	50MG/ML	N203479	001	Feb 06, 2013
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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>50MG</u>	<u>N019758</u>	<u>003</u>	May 20, 2019
<u>AB</u>	+!	<u>100MG</u>	<u>N019758</u>	<u>002</u>	Sep 26, 1989
<u>AB</u>	+	<u>200MG</u>	<u>N019758</u>	<u>004</u>	May 20, 2019

CLOZAPINE

	IVAX SUB TEVA PHARMS	12.5MG	A074949	003	Jul 31, 2003
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TABLET, ORALLY DISINTEGRATING; ORAL

CLOZAPINE

<u>AB</u>	BARR LABS INC	<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	<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
	BARR LABS INC	12.5MG	A090308	003	Apr 09, 2018
	TEVA PHARMS USA	150MG	A203039	001	Nov 25, 2015
		200MG	A203039	002	Nov 25, 2015

COBICISTAT

TABLET; ORAL

TYBOST

+	! GILEAD SCIENCES INC	150MG	N203094	001	Sep 24, 2014
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PRESCRIPTION DRUG PRODUCT LIST

COBICISTAT; DARUNAVIR

TABLET;ORAL

PREZCOBIX

+! JANSSEN PRODS 150MG;800MG N205395 001 Jan 29, 2015

COBICISTAT; DARUNAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

SYM TUZA

+! JANSSEN PRODS 150MG;800MG;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

NUMBRINO

+! CODY LABS INC 4% N209575 001 Jan 10, 2020

CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE**AA** ! PHARM ASSOC 10MG/5ML;5MG/5ML;6.25MG/5ML **A040660 001** Dec 07, 2006PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE**AA** AKORN 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 AKORN 10MG/5ML;6.25MG/5ML **A040151 001** Aug 26, 1997**AA** AMNEAL PHARMS 10MG/5ML;6.25MG/5ML **A200894 001** Apr 24, 2013**AA** NOSTRUM LABS INC 10MG/5ML;6.25MG/5ML **A090180 001** Mar 17, 2010**AA** QUAGEN 10MG/5ML;6.25MG/5ML **A214238 001** Oct 08, 2020**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** PHARM ASSOC 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** + HIKMA 15MG **N022402 001** Jul 16, 2009**AB** + 30MG **N022402 002** Jul 16, 2009**AB** +! 60MG **N022402 003** Jul 16, 2009**AB** LANNETT CO INC 15MG **A203046 001** Jun 13, 2014**AB** 30MG **A203046 002** Jun 13, 2014**AB** 60MG **A203046 003** Jun 13, 2014COLCHICINE

CAPSULE;ORAL

MITIGARE

+! HIKMA INTL PHARMS 0.6MG N204820 001 Sep 26, 2014

SOLUTION;ORAL

GLOPERBA

+! ROMEG 0.6MG/5ML N210942 001 Jan 30, 2019

PRESCRIPTION DRUG PRODUCT LIST

COLCHICINE

TABLET; ORAL

COLCHICINE

AB	ALKEM LABS LTD	<u>0.6MG</u>	<u>A211250 001</u>	Feb 08, 2019
AB	AMNEAL PHARMS	<u>0.6MG</u>	<u>A204711 001</u>	Sep 28, 2016
AB	AUROBINDO PHARMA LTD	<u>0.6MG</u>	<u>A215444 001</u>	Jan 06, 2022
AB	DR REDDYS	<u>0.6MG</u>	<u>A209876 001</u>	Sep 06, 2019
AB	GRANULES	<u>0.6MG</u>	<u>A210425 001</u>	Feb 05, 2020
AB	HETERO LABS LTD V	<u>0.6MG</u>	<u>A208993 001</u>	Aug 13, 2021
AB	MYLAN	<u>0.6MG</u>	<u>A209470 001</u>	Sep 16, 2019
AB	PAR PHARM INC	<u>0.6MG</u>	<u>A203976 001</u>	Aug 12, 2021
AB	WATSON LABS INC	<u>0.6MG</u>	<u>A204461 001</u>	Jul 31, 2019
AB	ZYDUS PHARMS	<u>0.6MG</u>	<u>A211519 001</u>	Feb 19, 2019

COLCRYS

AB	+ ! TAKEDA PHARMS USA	<u>0.6MG</u>	<u>N022352 001</u>	Jul 29, 2009
	COLCHICINE			
	ZYDUS PHARMS	0.3MG	A211519 002	Nov 14, 2019

COLCHICINE; PROBENECID

TABLET; ORAL

COL-PROBENECID

AB	+ ! WATSON LABS	<u>0.5MG;500MG</u>	<u>A084279 001</u>	
	<u>PROBENECID AND COLCHICINE</u>			
AB	NOVAST LABS	<u>0.5MG;500MG</u>	<u>A040618 001</u>	May 13, 2008

COLESEVELAM HYDROCHLORIDE

FOR SUSPENSION; ORAL

COLESEVELAM HYDROCHLORIDE

AB	ALKEM LABS LTD	<u>3.75GM/PACKET</u>	<u>A210316 001</u>	May 06, 2019
AB	GLENMARK PHARMS LTD	<u>3.75GM/PACKET</u>	<u>A202190 002</u>	Jul 16, 2018

WELCHOL

AB	+ ! DAIICHI SANKYO	<u>3.75GM/PACKET</u>	<u>N022362 002</u>	Oct 02, 2009
	COLESEVELAM HYDROCHLORIDE			
	GLENMARK PHARMS LTD	1.875GM/PACKET	A202190 001	Jul 16, 2018

TABLET; ORAL

COLESEVELAM HYDROCHLORIDE

AB	ALKEM LABS LTD	<u>625MG</u>	<u>A209038 001</u>	Oct 05, 2018
AB	BEIJING TIDE PHARM	<u>625MG</u>	<u>A206036 001</u>	Oct 14, 2021
AB	BIONPHARMA INC	<u>625MG</u>	<u>A208670 001</u>	Sep 13, 2019
AB	DR REDDYS	<u>625MG</u>	<u>A210889 001</u>	Oct 05, 2018
AB	GLENMARK PHARMS LTD	<u>625MG</u>	<u>A203480 001</u>	May 18, 2018
AB	IMPAX LABS INC	<u>625MG</u>	<u>A091600 001</u>	May 16, 2018
AB	INVENTIA	<u>625MG</u>	<u>A212050 001</u>	Dec 04, 2020
AB	LUPIN LTD	<u>625MG</u>	<u>A201354 001</u>	Dec 17, 2020
AB	ZHEJIANG JINGXIN	<u>625MG</u>	<u>A209946 001</u>	Jul 15, 2020
AB	ZYDUS PHARMS	<u>625MG</u>	<u>A207765 001</u>	Oct 07, 2019

WELCHOL

AB	+ ! DAIICHI SANKYO	<u>625MG</u>	<u>N021176 001</u>	May 26, 2000
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COLESTIPOL HYDROCHLORIDE

GRANULE; ORAL

COLESTID

AB	+ PFIZER	<u>5GM/SCOOPFUL</u>	<u>N017563 003</u>	Sep 22, 1995
AB	+ ! PFIZER	<u>5GM/PACKET</u>	<u>N017563 004</u>	Sep 22, 1995

COLESTIPOL HYDROCHLORIDE

AB	IMPAX LABS	<u>5GM/SCOOPFUL</u>	<u>A077277 001</u>	May 02, 2006
AB		<u>5GM/PACKET</u>	<u>A077277 002</u>	May 02, 2006

FLAVORED COLESTID

+ PFIZER

5GM/PACKET

N017563 001

+

5GM/SCOOPFUL

N017563 002

TABLET; ORAL

COLESTID

AB	+ ! PFIZER	<u>1GM</u>	<u>N020222 001</u>	Jul 19, 1994
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COLESTIPOL HYDROCHLORIDE

AB	IMPAX LABS	<u>1GM</u>	<u>A077510 001</u>	Oct 24, 2006
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COLISTIMETHATE SODIUM

INJECTABLE; INJECTION

COLISTIMETHATE SODIUM

AP	EMCURE PHARMS LTD	<u>EQ 150MG BASE/VIAL</u>	<u>A202359 001</u>	Sep 28, 2012
AP	FRESENIUS KABI USA	<u>EQ 150MG BASE/VIAL</u>	<u>A065364 001</u>	Apr 17, 2008
AP	NEXUS PHARMS	<u>EQ 150MG BASE/VIAL</u>	<u>A065177 001</u>	Mar 19, 2004
AP	SAGENT PHARMS INC	<u>EQ 150MG BASE/VIAL</u>	<u>A201365 001</u>	Feb 19, 2014
AP	XELLIA PHARMS APS	<u>EQ 150MG BASE/VIAL</u>	<u>A205356 001</u>	May 29, 2015

PRESCRIPTION DRUG PRODUCT LIST

COLISTIMETHATE SODIUM

INJECTABLE; INJECTION

COLISTIMETHATE SODIUM

AP	XGEN PHARMS	<u>EQ 150MG BASE/VIAL</u>	<u>A064216 001</u>	Feb 26, 1999
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COLY-MYCIN M

AP	+! PAR STERILE PRODUCTS	<u>EQ 150MG BASE/VIAL</u>	<u>N050108 002</u>	
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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	
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CONIVAPTAN HYDROCHLORIDE

INJECTABLE; INTRAVENOUS

VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER

+!	CUMBERLAND PHARMS	20MG/100ML (0.2MG/ML)	N021697 002	Oct 08, 2008
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COPANLISIB DIHYDROCHLORIDE

POWDER; INTRAVENOUS

ALIQOPA

+!	BAYER HEALTHCARE	60MG/VIAL	N209936 001	Sep 14, 2017
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COPPER

INTRAUTERINE DEVICE; INTRAUTERINE

PARAGARD T 380A

+!	COOPERSURGICAL	309MG/COPPER	N018680 001	Nov 15, 1984
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COPPER DOTATATE CU-64

SOLUTION; INTRAVENOUS

DETECTNET

+!	RADIOMEDIX	4mL (1mCi/ML)	N213227 001	Sep 03, 2020
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CORTICOTROPIN

INJECTABLE; INJECTION

ACTHAR GEL

+!	MALLINCKRODT ARD	80 UNITS/ML	N008372 008	
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PURIFIED CORTROPHIN GEL

+!	ANI PHARMS	80 UNITS/ML	N008975 002	
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CORTISONE ACETATE

TABLET; ORAL

CORTISONE ACETATE

!	HIKMA INTL PHARMS	25MG	A080776 002	
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COSYNTROPIN

INJECTABLE; INJECTION

CORTROSYN

AP	+! AMPHASTAR PHARMS INC	<u>0.25MG/VIAL</u>	<u>N016750 001</u>	
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COSYNTROPIN

AP	MYLAN INSTITUTIONAL	<u>0.25MG/VIAL</u>	<u>A090574 001</u>	Dec 17, 2009
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AP	SANDOZ	<u>0.25MG/VIAL</u>	<u>A202147 001</u>	Jun 29, 2012
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CRISABOROLE

OINTMENT; TOPICAL

EUCRISA

+!	ANACOR PHARMS INC	2%	N207695 001	Dec 14, 2016
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CRIZOTINIB

CAPSULE; ORAL

XALKORI

+	PF PRISM CV	200MG	N202570 001	Aug 26, 2011
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+!		250MG	N202570 002	Aug 26, 2011
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CROFELEMER

TABLET, DELAYED RELEASE; ORAL

MYTESI

+!	NAPO PHARMS INC	125MG	N202292 001	Dec 31, 2012
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CROMOLYN SODIUM

CONCENTRATE; ORAL

CROMOLYN SODIUM

AA	AILEX PHARMS LLC	<u>100MG/5ML</u>	<u>A209264 001</u>	Oct 16, 2017
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AA	MICRO LABS LTD	<u>100MG/5ML</u>	<u>A202745 001</u>	Apr 04, 2013
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INDIA

AA	RISING	<u>100MG/5ML</u>	<u>A202583 001</u>	Oct 27, 2011
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PRESCRIPTION DRUG PRODUCT LIST

CROMOLYN SODIUM

CONCENTRATE; ORAL

GASTROCROM

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 ! 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

CROTAMITON

LOTION; TOPICAL

CROTAN

AT MARNEL PHARMS **10%** **A087204 001**

EURAX

AT +! JOURNEY **10%** **N009112 003**

CUPRIC CHLORIDE

INJECTABLE; INJECTION

CUPRIC CHLORIDE IN PLASTIC CONTAINER

+! HOSPIRA EQ 0.4MG COPPER/ML **N018960 001** Jun 26, 1986

CUPRIC SULFATE; MANGANESE SULFATE; SELENIOUS ACID; ZINC SULFATE

SOLUTION; INTRAVENOUS

MULTRY'S

+! AM REGENT EQ 60MCG COPPER/ML; EQ 3MCG BASE/ML; EQ 6MCG SELENIUM/ML; EQ 1000MCG BASE/ML (1ML) **N209376 003** Jun 30, 2021

TRALEMENT

+! AM REGENT EQ 0.3MG COPPER/ML; EQ 55MCG BASE/ML; EQ 60MCG SELENIUM/ML; EQ 3MG BASE/ML (1ML) **N209376 001** Jul 02, 2020

+! EQ 0.3MG COPPER/ML; EQ 55MCG BASE/ML; EQ 60MCG SELENIUM/ML; EQ 3MG BASE/ML (5ML) **N209376 002** Dec 02, 2020

CYANOCOBALAMIN

INJECTABLE; INJECTION

CYANOCOBALAMIN

AP +! AM REGENT **1MG/ML** **A080737 001**

AP EUGIA PHARMA **1MG/ML** **A213874 001** Dec 08, 2020

AP GLAND PHARMA LTD **1MG/ML** **A214390 001** Sep 24, 2020

AP MYLAN LABS LTD **1MG/ML** **A204829 001** Jun 05, 2017

AP SANDOZ INC **1MG/ML** **A212915 001** Jan 04, 2021

AP SOMERSET THERAPS **1MG/ML** **A206503 001** Dec 11, 2015

LLC

AP **1MG/ML** **A209429 001** Dec 18, 2018

AP VITRUVIAS THERAP **1MG/ML** **A209255 001** Dec 18, 2018

AP WEST-WARD PHARMS **1MG/ML** **A080515 002**

INT

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, 2007

CYCLOBENZAPRINE HYDROCHLORIDE

AB APOTEX **15MG** **A206703 001** Jul 24, 2018

AB **30MG** **A206703 002** Jul 24, 2018

AB TWI PHARMS INC **15MG** **A091281 001** Jan 31, 2013

AB **30MG** **A091281 002** Jan 31, 2013

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 ALEMBIC LABS **5MG** **A078218 002** Jun 19, 2015

AB **7.5MG** **A078218 003** Nov 03, 2020

AB **10MG** **A078218 001** Apr 18, 2008

PRESCRIPTION DRUG PRODUCT LIST

CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

<u>AB</u>	ANDA REPOSITORY	<u>5MG</u>	<u>A073541 002</u>	Apr 06, 2006
<u>AB</u>		<u>10MG</u>	<u>A073541 001</u>	May 23, 1995
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A078643 001</u>	Sep 26, 2008
<u>AB</u>		<u>10MG</u>	<u>A078643 002</u>	Sep 26, 2008
<u>AB</u>	INVAGEN PHARMS	<u>5MG</u>	<u>A090478 001</u>	Jul 23, 2010
<u>AB</u>		<u>10MG</u>	<u>A090478 002</u>	Jul 23, 2010
<u>AB</u>	JUBILANT CADISTA	<u>5MG</u>	<u>A077563 001</u>	Apr 19, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A077563 003</u>	Aug 25, 2017
<u>AB</u>		<u>10MG</u>	<u>A077563 002</u>	Apr 19, 2006
<u>AB</u>	KVK TECH	<u>5MG</u>	<u>A078048 001</u>	Feb 28, 2011
<u>AB</u>		<u>10MG</u>	<u>A078048 002</u>	Feb 28, 2011
<u>AB</u>	OXFORD PHARMS	<u>5MG</u>	<u>A077209 002</u>	Feb 03, 2006
<u>AB</u>		<u>10MG</u>	<u>A077209 001</u>	Oct 04, 2005
<u>AB</u>	PRINSTON INC	<u>5MG</u>	<u>A077797 001</u>	Feb 28, 2007
<u>AB</u>		<u>10MG</u>	<u>A077797 002</u>	Feb 28, 2007
<u>AB</u>	RUBICON	<u>5MG</u>	<u>A208170 001</u>	May 31, 2017
<u>AB</u>		<u>7.5MG</u>	<u>A208170 002</u>	May 31, 2017
<u>AB</u>	!	<u>10MG</u>	<u>A208170 003</u>	May 31, 2017
<u>AB</u>	SUN PHARM INDS LTD	<u>5MG</u>	<u>A078722 001</u>	May 12, 2008
<u>AB</u>		<u>7.5MG</u>	<u>A078722 002</u>	May 12, 2008
<u>AB</u>		<u>10MG</u>	<u>A078722 003</u>	May 12, 2008
<u>AB</u>	UNICHEM	<u>5MG</u>	<u>A213324 001</u>	Jul 06, 2020
<u>AB</u>		<u>7.5MG</u>	<u>A213324 002</u>	Jul 06, 2020
<u>AB</u>		<u>10MG</u>	<u>A213324 003</u>	Jul 05, 2020

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AKPENTOLATE

<u>AT</u>	AKORN	<u>1%</u>	<u>A040164 001</u>	Jan 13, 1997
<u>AT</u>		<u>2%</u>	<u>A040165 001</u>	Jan 13, 1997

CYCLOGYL

<u>AT</u>	+!	ALCON LABS INC	<u>0.5%</u>	<u>A084109 001</u>
<u>AT</u>	+!		<u>1%</u>	<u>A084110 001</u>
<u>AT</u>	+!		<u>2%</u>	<u>A084108 001</u>

CYCLOPENTOLATE HYDROCHLORIDE

<u>AT</u>	AKORN	<u>0.5%</u>	<u>A205937 001</u>	Dec 09, 2015
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PENTOLAIR

<u>AT</u>	BAUSCH AND LOMB	<u>1%</u>	<u>A040075 001</u>	Apr 29, 1994
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CYCLOPENTOLATE HYDROCHLORIDE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CYCLOMYDRIL

!	ALCON LABS INC	0.2%;1%	A084300 001	
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CYCLOPHOSPHAMIDE

CAPSULE; ORAL

CYCLOPHOSPHAMIDE

<u>AB</u>	ANI PHARMS	<u>25MG</u>	<u>A207014 001</u>	Mar 19, 2018	
<u>AB</u>		<u>50MG</u>	<u>A207014 002</u>	Mar 19, 2018	
<u>AB</u>	CIPLA	<u>25MG</u>	<u>A211608 001</u>	Jan 18, 2019	
<u>AB</u>		<u>50MG</u>	<u>A211608 002</u>	Jan 18, 2019	
<u>AB</u>	+	HIKMA	<u>25MG</u>	<u>N203856 001</u>	Sep 16, 2013
<u>AB</u>	+!		<u>50MG</u>	<u>N203856 002</u>	Sep 16, 2013
<u>AB</u>	STI PHARMA LLC	<u>25MG</u>	<u>A209872 001</u>	May 07, 2018	
<u>AB</u>		<u>50MG</u>	<u>A209872 002</u>	May 07, 2018	

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

<u>AP</u>	AMNEAL	<u>500MG/VIAL</u>	<u>A210046 001</u>	May 25, 2018	
<u>AP</u>		<u>1GM/VIAL</u>	<u>A210046 002</u>	May 25, 2018	
<u>AP</u>		<u>2GM/VIAL</u>	<u>A210046 003</u>	May 25, 2018	
<u>AP</u>	!	BAXTER HLTHCARE	<u>500MG/VIAL</u>	<u>A040745 001</u>	May 21, 2008
<u>AP</u>	!		<u>1GM/VIAL</u>	<u>A040745 002</u>	May 21, 2008
<u>AP</u>	!		<u>2GM/VIAL</u>	<u>A040745 003</u>	May 21, 2008
<u>AP</u>	JIANGSU PHARMS	<u>500MG/VIAL</u>	<u>A204555 001</u>	Oct 31, 2014	
<u>AP</u>		<u>1GM/VIAL</u>	<u>A204555 002</u>	Oct 31, 2014	
<u>AP</u>		<u>2GM/VIAL</u>	<u>A204555 003</u>	Oct 31, 2014	

SOLUTION; INTRAVENOUS

CYCLOPHOSPHAMIDE

+!	EUGIA PHARMA SPECLTS	500MG/2.5ML (200MG/ML)	N210735 001	Aug 25, 2021
+!		1GM/5ML (200MG/ML)	N210735 002	Aug 25, 2021

PRESCRIPTION DRUG PRODUCT LIST

CYCLOPHOSPHAMIDE

SOLUTION; INTRAVENOUS

CYCLOPHOSPHAMIDE

+!	INGENUS PHARMS LLC	500MG/2.5ML (200MG/ML)	N212501 001	Jul 30, 2020
+!		1GM/5ML (200MG/ML)	N212501 002	Jul 30, 2020
+!		2GM/10ML (200MG/ML)	N212501 003	Nov 19, 2021

TABLET; ORAL

CYTOXAN

+	BAXTER HLTHCARE	25MG	N012141 002	
+!		50MG	N012141 001	

CYCLOSERINE

CAPSULE; ORAL

SEROMYCIN

!	PURDUE	250MG	A060593 001	
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CYCLOSPORINE

CAPSULE; ORAL

CYCLOSPORINE

AB1	APOTEX	25MG	A210721 001	Jul 10, 2019
AB1		50MG	A210721 002	Jul 10, 2019
AB1		100MG	A210721 003	Jul 10, 2019
AB1	IVAX SUB TEVA PHARMS	25MG	A065110 003	Mar 29, 2005
AB1		50MG	A065110 001	Mar 29, 2005
AB1		100MG	A065110 002	Mar 29, 2005
AB1	MAYNE PHARMA	25MG	A065044 002	Dec 20, 2000
AB1		100MG	A065044 001	Dec 20, 2000
AB1	SANDOZ	25MG	A065017 002	Jan 13, 2000
AB1		100MG	A065017 001	Jan 13, 2000
	GENGRAF			
AB1	ABBEVIE	25MG	A065003 001	May 12, 2000
AB1		100MG	A065003 003	May 12, 2000
	NEORAL			
AB1	+ NOVARTIS	25MG	N050715 001	Jul 14, 1995
AB1	+!	100MG	N050715 002	Jul 14, 1995
	CYCLOSPORINE			
AB2	APOTEX	25MG	A065040 001	May 09, 2002
AB2		100MG	A065040 002	May 09, 2002
	SANDIMMUNE			
AB2	+ NOVARTIS	25MG	N050625 001	Mar 02, 1990
AB2	+!	100MG	N050625 002	Mar 02, 1990
BX	+	50MG	N050625 003	Nov 23, 1992
	EMULSION; OPHTHALMIC			
	RESTASIS			
	+! ALLERGAN	0.05%	N050790 001	Dec 23, 2002
	RESTASIS MULTIDOSE			
	+! ALLERGAN	0.05%	N050790 002	Oct 27, 2016
	VERKAZIA			
	+! SANTEN	0.1%	N214965 001	Jun 23, 2021
	INJECTABLE; INJECTION			
	CYCLOSPORINE			
AP	HIKMA	50MG/ML	A065004 001	Oct 29, 1999
AP	PADAGIS US	50MG/ML	A065151 001	Oct 07, 2003
	SANDIMMUNE			
AP	+! NOVARTIS	50MG/ML	N050573 001	Nov 14, 1983
	SOLUTION; OPHTHALMIC			
	CEQUA			
	+! SUN PHARM	0.09%	N210913 001	Aug 14, 2018
	SOLUTION; ORAL			
	CYCLOSPORINE			
AB1	ABBEVIE	100MG/ML	A065025 001	Mar 03, 2000
AB1	IVAX SUB TEVA PHARMS	100MG/ML	A065078 001	Mar 25, 2005
AB1	MAYNE PHARMA	100MG/ML	A065054 001	Dec 18, 2001
	NEORAL			
AB1	+! NOVARTIS	100MG/ML	N050716 001	Jul 14, 1995
	CYCLOSPORINE			
AB2	WOCKHARDT BIO AG	100MG/ML	A065133 001	Sep 17, 2004
	SANDIMMUNE			
AB2	+! NOVARTIS	100MG/ML	N050574 001	Nov 14, 1983

PRESCRIPTION DRUG PRODUCT LISTCYPROHEPTADINE HYDROCHLORIDE

SYRUP; ORAL

CYPROHEPTADINE HYDROCHLORIDE

<u>AA</u>	ANDA REPOSITORY	<u>2MG/5ML</u>	<u>A204823</u>	<u>001</u>	Dec 27, 2016
<u>AA</u>	ELYSIUM	<u>2MG/5ML</u>	<u>A209108</u>	<u>001</u>	Oct 16, 2018
<u>AA</u>	LANNETT CO INC	<u>2MG/5ML</u>	<u>A203191</u>	<u>001</u>	Jul 13, 2017
<u>AA</u>	! LYNE	<u>2MG/5ML</u>	<u>A040668</u>	<u>001</u>	Jun 28, 2006
<u>AA</u>	PHARM ASSOC	<u>2MG/5ML</u>	<u>A091295</u>	<u>001</u>	Mar 28, 2013
<u>AA</u>	QUAGEN	<u>2MG/5ML</u>	<u>A212423</u>	<u>001</u>	May 22, 2019
<u>AA</u>	TRIS PHARMA INC	<u>2MG/5ML</u>	<u>A205431</u>	<u>001</u>	Dec 21, 2021

TABLET; ORAL

CYPROHEPTADINE HYDROCHLORIDE

<u>AA</u>	APPCO	<u>4MG</u>	<u>A206553</u>	<u>001</u>	Nov 29, 2016
<u>AA</u>	BEXIMCO PHARMS USA	<u>4MG</u>	<u>A206676</u>	<u>001</u>	Apr 12, 2019
<u>AA</u>	BOSCOGEN	<u>4MG</u>	<u>A040644</u>	<u>001</u>	May 30, 2006
<u>AA</u>	ELYSIUM	<u>4MG</u>	<u>A207555</u>	<u>001</u>	Jan 31, 2017
<u>AA</u>	+! HERITAGE PHARMA	<u>4MG</u>	<u>A087056</u>	<u>001</u>	
<u>AA</u>	MOUNTAIN	<u>4MG</u>	<u>A040537</u>	<u>001</u>	Sep 30, 2003
<u>AA</u>	NOVAST LABS	<u>4MG</u>	<u>A205087</u>	<u>001</u>	Sep 23, 2015
<u>AA</u>	QUAGEN	<u>4MG</u>	<u>A212491</u>	<u>001</u>	Feb 24, 2021
<u>AA</u>	RISING	<u>4MG</u>	<u>A207783</u>	<u>001</u>	Dec 29, 2016
<u>AA</u>	STRIDES PHARMA	<u>4MG</u>	<u>A209172</u>	<u>001</u>	Apr 11, 2018
<u>AA</u>	ZYDUS PHARMS	<u>4MG</u>	<u>A208938</u>	<u>001</u>	May 19, 2017

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

PROCYSBI

+	HORIZON	EQ 25MG BASE	N203389	001	Apr 30, 2013
+	!	EQ 75MG BASE	N203389	002	Apr 30, 2013

GRANULE, DELAYED RELEASE; ORAL

PROCYSBI

+	HORIZON PHARMA USA	EQ 75MG BASE/PACKET	N213491	001	Feb 14, 2020
+	!	EQ 300MG BASE/PACKET	N213491	002	Feb 14, 2020

CYSTEAMINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CYSTADROPS

+	!	RECORDATI RARE	EQ 0.37% BASE	N211302	001	Aug 19, 2020
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CYSTARAN

+	!	LEADIANT BIOSCI INC	EQ 0.44% BASE	N200740	001	Oct 02, 2012
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CYSTEINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

ELCYS

+	!	EXELA PHARMA	500MG/10ML (50MG/ML)	N210660	001	Apr 16, 2019
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CYTARABINE

INJECTABLE; INJECTION

CYTARABINE

<u>AP</u>	!	FRESENIUS KABI USA	<u>100MG/ML</u>	<u>A076512</u>	<u>001</u>	Jan 15, 2004
<u>AP</u>		GLAND PHARMA LTD	<u>100MG/VIAL</u>	<u>A211937</u>	<u>001</u>	Dec 23, 2019
<u>AP</u>			<u>2GM/VIAL</u>	<u>A211938</u>	<u>001</u>	Dec 23, 2019
<u>AP</u>		HIKMA	<u>100MG/VIAL</u>	<u>A071471</u>	<u>001</u>	Aug 02, 1989
<u>AP</u>	!	HOSPIRA	<u>20MG/ML</u>	<u>A071868</u>	<u>001</u>	Jun 04, 1990
<u>AP</u>	!		<u>20MG/ML</u>	<u>A072168</u>	<u>001</u>	Aug 31, 1990
<u>AP</u>	!		<u>20MG/ML</u>	<u>A072945</u>	<u>001</u>	Feb 28, 1994
<u>AP</u>			<u>100MG/ML</u>	<u>A075383</u>	<u>001</u>	Nov 22, 1999
<u>AP</u>		MEITHEAL	<u>100MG/ML</u>	<u>A205696</u>	<u>001</u>	Jul 17, 2018
<u>AP</u>		RISING PHARMA	<u>20MG/ML</u>	<u>A200915</u>	<u>001</u>	Dec 13, 2011
<u>AP</u>			<u>100MG/ML</u>	<u>A201784</u>	<u>001</u>	Jan 30, 2012
<u>AP</u>	!	WEST-WARD PHARMS	<u>2GM/VIAL</u>	<u>A074245</u>	<u>002</u>	Aug 31, 1994
		INT				
	!	HIKMA	500MG/VIAL	A071472	001	Aug 02, 1989
	!	WEST-WARD PHARMS	1GM/VIAL	A074245	001	Aug 31, 1994
		INT				

PRESCRIPTION DRUG PRODUCT LIST

CYTARABINE; DAUNORUBICIN

POWDER; INTRAVENOUS

VYXEOS

+! CELATOR PHARMS 100MG;44MG N209401 001 Aug 03, 2017

DABIGATRAN ETEXILATE MESYLATE

CAPSULE; ORAL

DABIGATRAN ETEXILATE MESYLATE**AB** ALKEM LABS LTD **EQ 75MG BASE** **A208040 001** Mar 11, 2020**AB** **EQ 150MG BASE** **A208040 002** Mar 11, 2020**AB** HETERO LABS LTD III **EQ 75MG BASE** **A207961 001** May 06, 2020**AB** **EQ 150MG BASE** **A207961 002** May 06, 2020PRADAXA**AB** + BOEHRINGER **EQ 75MG BASE** **N022512 001** Oct 19, 2010
INGELHEIM**AB** +! **EQ 150MG BASE** **N022512 002** Oct 19, 2010+ **EQ 110MG BASE** N022512 003 Nov 20, 2015

PELLETS; ORAL

PRADAXA+ BOEHRINGER **EQ 20MG BASE/PACKET** N214358 001 Jun 21, 2021

+ INGELHEIM

+ **EQ 30MG BASE/PACKET** N214358 002 Jun 21, 2021+ **EQ 40MG BASE/PACKET** N214358 003 Jun 21, 2021+ **EQ 50MG BASE/PACKET** N214358 004 Jun 21, 2021+ **EQ 110MG BASE/PACKET** N214358 005 Jun 21, 2021+! **EQ 150MG BASE/PACKET** N214358 006 Jun 21, 2021DABRAFENIB MESYLATE

CAPSULE; ORAL

TAFINLAR

+ NOVARTIS **EQ 50MG BASE** N202806 001 May 29, 2013+! **EQ 75MG BASE** N202806 002 May 29, 2013DACARBAZINE

INJECTABLE; INJECTION

DACARBAZINE**AP** ! FRESENIUS KABI USA **200MG/VIAL** **A075371 002** Aug 27, 1999**AP** HIKMA **200MG/VIAL** **A075812 001** Jun 15, 2001**AP** **500MG/VIAL** **A075812 002** Oct 31, 2002**AP** HOSPIRA **200MG/VIAL** **A075940 001** Oct 18, 2001**AP** TEVA PHARMS USA **200MG/VIAL** **A075259 002** Aug 27, 1998**AP** ! **500MG/VIAL** **A075259 001** Sep 22, 2000

! FRESENIUS KABI USA 100MG/VIAL A075371 001 Aug 27, 1999

DACOMITINIB

TABLET; ORAL

VIZIMPRO

+ PFIZER 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 **0.5MG/VIAL** **N050682 001**DACTINOMYCIN**AP** EUGIA PHARMA **0.5MG/VIAL** **A203385 001** Nov 09, 2017**AP** HISUN PHARM **0.5MG/VIAL** **A207232 001** Jul 16, 2019

HANGZHOU

AP MEITHEAL **0.5MG/VIAL** **A213463 001** Nov 13, 2020**AP** XGEN PHARMS **0.5MG/VIAL** **A203999 001** May 20, 2019DALBAVANCIN HYDROCHLORIDE

POWDER; INTRAVENOUS

DALVANCE

+! ALLERGAN **EQ 500MG BASE/VIAL** N021883 001 May 23, 2014DALFAMPRIDINE

TABLET, EXTENDED RELEASE; ORAL

AMPYRA**AB** +! ACORDA **10MG** **N022250 001** Jan 22, 2010DALFAMPRIDINE**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 **10MG** **A206811 001** Jan 23, 2017

LTD

PRESCRIPTION DRUG PRODUCT LIST

DALFAMPRIDINE

TABLET, EXTENDED RELEASE;ORAL

DALFAMPRIDINE

<u>AB</u>	HIKMA	<u>10MG</u>	<u>A206646</u>	<u>001</u>	Oct 24, 2018
<u>AB</u>	MICRO LABS	<u>10MG</u>	<u>A210158</u>	<u>001</u>	Mar 11, 2019
<u>AB</u>	SUN PHARM	<u>10MG</u>	<u>A208292</u>	<u>001</u>	May 21, 2019

DALFOPRISTIN; QUINUPRISTIN

INJECTABLE; INTRAVENOUS

SYNERCID

+	KING PHARMS	350MG/VIAL;150MG/VIAL	N050748	001	Sep 21, 1999
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DALTEPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

FRAGMIN

+	PFIZER	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

<u>AB</u>	BARR	<u>50MG</u>	<u>A074582</u>	<u>003</u>	May 29, 1998
<u>AB</u>		<u>100MG</u>	<u>A074582</u>	<u>002</u>	May 29, 1998
<u>AB</u>	!	<u>200MG</u>	<u>A074582</u>	<u>001</u>	Aug 09, 1996
<u>AB</u>	LANNETT CO INC	<u>50MG</u>	<u>A077246</u>	<u>002</u>	Apr 19, 2007
<u>AB</u>		<u>100MG</u>	<u>A077246</u>	<u>003</u>	Apr 19, 2007
<u>AB</u>		<u>200MG</u>	<u>A077246</u>	<u>001</u>	Sep 28, 2005

DANTROLENE SODIUM

CAPSULE; ORAL

DANTRIUM

<u>AB</u>	+	PAR STERILE PRODUCTS	<u>25MG</u>	<u>N017443</u>	<u>001</u>
<u>AB</u>	+		<u>50MG</u>	<u>N017443</u>	<u>003</u>
<u>AB</u>	+	!	<u>100MG</u>	<u>N017443</u>	<u>002</u>

DANTROLENE SODIUM

<u>AB</u>	ELITE LABS INC	<u>25MG</u>	<u>A076686</u>	<u>001</u>	Oct 24, 2005
<u>AB</u>		<u>50MG</u>	<u>A076686</u>	<u>002</u>	Oct 24, 2005
<u>AB</u>		<u>100MG</u>	<u>A076686</u>	<u>003</u>	Oct 24, 2005
<u>AB</u>	IMPAX LABS	<u>25MG</u>	<u>A076856</u>	<u>001</u>	Mar 01, 2005
<u>AB</u>		<u>50MG</u>	<u>A076856</u>	<u>002</u>	Mar 01, 2005
<u>AB</u>		<u>100MG</u>	<u>A076856</u>	<u>003</u>	Mar 01, 2005

FOR SUSPENSION; INTRAVENOUS

RYANODEX

+	EAGLE PHARMS	250MG/VIAL	N205579	001	Jul 22, 2014
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INJECTABLE; INJECTION

DANTRIUM

<u>AP</u>	+	PAR STERILE PRODUCTS	<u>20MG/VIAL</u>	<u>N018264</u>	<u>001</u>
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DANTROLENE SODIUM

<u>AP</u>	HIKMA	<u>20MG/VIAL</u>	<u>A204762</u>	<u>001</u>	Jun 19, 2017
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REVONTO

<u>AP</u>	USWM	<u>20MG/VIAL</u>	<u>A078378</u>	<u>001</u>	Jul 24, 2007
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DAPAGLIFLOZIN

TABLET; ORAL

FARXIGA

+	ASTRAZENECA AB	5MG	N202293	001	Jan 08, 2014
+	!	10MG	N202293	002	Jan 08, 2014

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

PRESCRIPTION DRUG PRODUCT LIST

DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE

TABLET; ORAL

QTERN

+	ASTRAZENECA AB	5MG;EQ 5MG BASE	N209091	002	May 02, 2019
+	!	10MG;EQ 5MG BASE	N209091	001	Feb 27, 2017

DAPSONE

GEL; TOPICAL

ACZONE

<u>AB</u>	+	ALLERGAN	<u>5%</u>	<u>N021794</u>	<u>001</u>	Jul 07, 2005
<u>AB</u>	+	ALMIRALL	<u>7.5%</u>	<u>N207154</u>	<u>001</u>	Feb 24, 2016

DAPSONE

<u>AB</u>		TARO	<u>5%</u>	<u>A209506</u>	<u>001</u>	Oct 16, 2017
<u>AB</u>		TARO PHARMS	<u>7.5%</u>	<u>A210191</u>	<u>001</u>	Jun 26, 2019

TABLET; ORAL

DAPSONE

<u>AB</u>		ACTAVIS LLC	<u>25MG</u>	<u>A204380</u>	<u>001</u>	Mar 23, 2017
<u>AB</u>			<u>100MG</u>	<u>A204380</u>	<u>002</u>	Mar 23, 2017
<u>AB</u>		ALVOGEN	<u>25MG</u>	<u>A205429</u>	<u>001</u>	Jan 07, 2016
<u>AB</u>			<u>100MG</u>	<u>A205429</u>	<u>002</u>	Jan 07, 2016
<u>AB</u>	+	JACOBUS	<u>25MG</u>	<u>A086841</u>	<u>001</u>	
<u>AB</u>	+	!	<u>100MG</u>	<u>A086842</u>	<u>001</u>	
<u>AB</u>		NOSTRUM LABS INC	<u>25MG</u>	<u>A203887</u>	<u>001</u>	May 06, 2016
<u>AB</u>			<u>100MG</u>	<u>A203887</u>	<u>002</u>	May 06, 2016
<u>AB</u>		NOVITIUM PHARMA	<u>25MG</u>	<u>A206505</u>	<u>001</u>	Dec 01, 2016
<u>AB</u>			<u>100MG</u>	<u>A206505</u>	<u>002</u>	Dec 01, 2016
<u>AB</u>		RISING	<u>100MG</u>	<u>A207165</u>	<u>001</u>	May 08, 2019
<u>AB</u>		VIRTUS PHARMS	<u>25MG</u>	<u>A204074</u>	<u>001</u>	May 10, 2016
<u>AB</u>			<u>100MG</u>	<u>A204074</u>	<u>002</u>	May 10, 2016

DAPTOMYCIN

POWDER; INTRAVENOUS

CUBICIN

<u>AP</u>	+	CUBIST PHARMS LLC	<u>500MG/VIAL</u>	<u>N021572</u>	<u>002</u>	Sep 12, 2003
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DAPTOMYCIN

<u>AP</u>		ACCORD HLTHCARE	<u>350MG/VIAL</u>	<u>A212667</u>	<u>001</u>	Jul 12, 2019
<u>AP</u>			<u>500MG/VIAL</u>	<u>A211961</u>	<u>001</u>	Jun 24, 2019
<u>AP</u>		BE PHARMS	<u>350MG/VIAL</u>	<u>A213425</u>	<u>001</u>	Aug 20, 2020
<u>AP</u>			<u>500MG/VIAL</u>	<u>A212513</u>	<u>001</u>	Jun 26, 2019
<u>AP</u>		DR REDDYS	<u>350MG/VIAL</u>	<u>A211403</u>	<u>001</u>	Aug 31, 2020
<u>AP</u>			<u>500MG/VIAL</u>	<u>A208375</u>	<u>001</u>	May 01, 2019
<u>AP</u>		EUGIA PHARMA	<u>500MG/VIAL</u>	<u>A213171</u>	<u>001</u>	Sep 02, 2021
<u>AP</u>		FRESENIUS KABI USA	<u>500MG/VIAL</u>	<u>A206077</u>	<u>001</u>	Apr 11, 2018
<u>AP</u>		HANGZHOU ZHONGMEI	<u>500MG/VIAL</u>	<u>A215215</u>	<u>001</u>	Nov 26, 2021
<u>AP</u>		JIANGSU PHARMS	<u>500MG/VIAL</u>	<u>A212022</u>	<u>001</u>	Aug 22, 2019
<u>AP</u>		MEITHEAL	<u>350MG/VIAL</u>	<u>A213786</u>	<u>001</u>	Jun 29, 2021
<u>AP</u>			<u>500MG/VIAL</u>	<u>A213623</u>	<u>001</u>	Jun 29, 2021
<u>AP</u>		MYLAN LABS LTD	<u>500MG/VIAL</u>	<u>A205037</u>	<u>001</u>	Jun 05, 2018
<u>AP</u>		QILU PHARM HAINAN	<u>500MG/VIAL</u>	<u>A215316</u>	<u>001</u>	Aug 24, 2021
<u>AP</u>	+	SAGENT PHARMS INC	<u>350MG/VIAL</u>	<u>N208385</u>	<u>001</u>	Sep 12, 2017
<u>AP</u>			<u>500MG/VIAL</u>	<u>A207104</u>	<u>001</u>	Nov 15, 2019
<u>AP</u>		TEVA PHARMS USA	<u>500MG/VIAL</u>	<u>A091039</u>	<u>001</u>	Mar 25, 2016
<u>AP</u>		XELLIA PHARMS APS	<u>500MG/VIAL</u>	<u>A206005</u>	<u>001</u>	Jun 15, 2016

CUBICIN RF

+	CUBIST PHARMS LLC	500MG/VIAL	N021572	003	Jul 06, 2016
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DAPTOMYCIN

+	HOSPIRA INC	350MG/VIAL	N210282	001	Jun 21, 2021
+	!	500MG/VIAL	N210282	002	Jun 21, 2021
+	XELLIA PHARMS APS	350MG/VIAL	N209949	001	Oct 20, 2017

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE; ORAL

DARIFENACIN

<u>AB</u>		MACLEODS PHARMS LTD	<u>EQ 7.5MG BASE</u>	<u>A207302</u>	<u>001</u>	Jul 28, 2017
<u>AB</u>	!	!	<u>EQ 15MG BASE</u>	<u>A207302</u>	<u>002</u>	Jul 28, 2017

DARIFENACIN HYDROBROMIDE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>EQ 7.5MG BASE</u>	<u>A207681</u>	<u>001</u>	Dec 08, 2017
<u>AB</u>			<u>EQ 15MG BASE</u>	<u>A207681</u>	<u>002</u>	Dec 08, 2017
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 7.5MG BASE</u>	<u>A206743</u>	<u>001</u>	Sep 19, 2016
<u>AB</u>			<u>EQ 15MG BASE</u>	<u>A206743</u>	<u>002</u>	Sep 19, 2016
<u>AB</u>		CIPLA	<u>EQ 7.5MG BASE</u>	<u>A207664</u>	<u>001</u>	Sep 01, 2016
<u>AB</u>			<u>EQ 15MG BASE</u>	<u>A207664</u>	<u>002</u>	Sep 01, 2016
<u>AB</u>		POLYGEN PHARMS	<u>EQ 7.5MG BASE</u>	<u>A211045</u>	<u>001</u>	Jan 06, 2020

PRESCRIPTION DRUG PRODUCT LIST

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE;ORAL

DARIFENACIN HYDROBROMIDE

<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A211045 002</u>	Jan 06, 2020
<u>AB</u>	TORRENT	<u>EQ 7.5MG BASE</u>	<u>A205209 001</u>	Nov 17, 2016
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A205209 002</u>	Nov 17, 2016
BX	XIROMED	EQ 7.5MG BASE	A209571 002	Oct 22, 2019
BX		EQ 15MG BASE	A209571 001	Oct 22, 2019

DAROLUTAMIDE

TABLET;ORAL

NUBEQA

+	!	BAYER HEALTHCARE	300MG	N212099 001	Jul 30, 2019
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DARUNAVIR

SUSPENSION;ORAL

PREZISTA

+	!	JANSSEN PRODS	100MG/ML	N202895 001	Dec 16, 2011
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TABLET;ORAL

DARUNAVIR

<u>AB</u>		TEVA PHARMS USA	<u>600MG</u>	<u>A202118 001</u>	Nov 21, 2017
<u>AB</u>	+	JANSSEN PRODS	<u>600MG</u>	<u>N021976 002</u>	Feb 25, 2008
	+		75MG	N021976 004	Dec 18, 2008
	+		150MG	N021976 005	Dec 18, 2008
	+	!	800MG	N021976 006	Nov 09, 2012

DASATINIB

TABLET;ORAL

DASATINIB

<u>AB</u>		APOTEX	<u>80MG</u>	<u>A203180 001</u>	Nov 23, 2021
<u>AB</u>			<u>140MG</u>	<u>A203180 002</u>	Nov 23, 2021
<u>AB</u>	+	BRISTOL MYERS SQUIBB	<u>80MG</u>	<u>N021986 005</u>	Oct 28, 2010
<u>AB</u>	+		<u>140MG</u>	<u>N021986 006</u>	Oct 28, 2010
	+		20MG	N021986 001	Jun 28, 2006
	+		50MG	N021986 002	Jun 28, 2006
	+		70MG	N021986 003	Jun 28, 2006
	+	!	100MG	N021986 004	May 30, 2008

DASIGLUCAGON HYDROCHLORIDE

SOLUTION;SUBCUTANEOUS

ZEGALOGUE

+	!	ZEALAND PHARMA	EQ 0.6MG BASE/0.6ML (EQ 0.6MG BASE/0.6ML)	N214231 001	Mar 22, 2021
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ZEGALOGUE (AUTOINJECTOR)

+	!	ZEALAND PHARMA	EQ 0.6MG BASE/0.6ML (EQ 0.6MG BASE/0.6ML)	N214231 002	Mar 22, 2021
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DAUNORUBICIN HYDROCHLORIDE

INJECTABLE;INJECTION

CERUBIDINE

<u>AP</u>	!	HIKMA	<u>EQ 20MG BASE/VIAL</u>	<u>A064103 001</u>	Feb 03, 1995
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DAUNORUBICIN HYDROCHLORIDE

<u>AP</u>		FRESENIUS KABI USA	<u>EQ 20MG BASE/VIAL</u>	<u>A065000 001</u>	May 25, 1999
<u>AP</u>	+	HIKMA	<u>EQ 5MG BASE/ML</u>	<u>N050731 001</u>	Jan 30, 1998
<u>AP</u>		HISUN PHARM HANGZHOU	<u>EQ 5MG BASE/ML</u>	<u>A208759 001</u>	Apr 12, 2019
<u>AP</u>		TEVA PHARMS USA	<u>EQ 5MG BASE/ML</u>	<u>A065035 001</u>	Jan 24, 2000
		FRESENIUS KABI USA	EQ 5MG BASE/VIAL	A065034 001	Nov 20, 2001

DECITABINE

INJECTABLE;INTRAVENOUS

DACOGEN

<u>AP</u>	+	!	OTSUKA	<u>50MG/VIAL</u>	<u>N021790 001</u>	May 02, 2006
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DECITABINE

<u>AP</u>		ACCORD HLTHCARE	<u>50MG/VIAL</u>	<u>A203475 001</u>	Feb 27, 2017
<u>AP</u>		CHEMI SPA	<u>50MG/VIAL</u>	<u>A206033 001</u>	Sep 22, 2017
<u>AP</u>		DR REDDYS	<u>50MG/VIAL</u>	<u>A203131 001</u>	Jul 11, 2013
<u>AP</u>		EUGIA PHARMA	<u>50MG/VIAL</u>	<u>A214569 001</u>	Sep 20, 2021
<u>AP</u>		GLAND	<u>50MG/VIAL</u>	<u>A205539 001</u>	Nov 23, 2020
<u>AP</u>		INGENUS PHARMS LLC	<u>50MG/VIAL</u>	<u>A210984 001</u>	Sep 16, 2019
<u>AP</u>		LUPIN LTD	<u>50MG/VIAL</u>	<u>A210756 001</u>	Nov 09, 2018
<u>AP</u>		MEITHEAL	<u>50MG/VIAL</u>	<u>A212959 001</u>	Jul 02, 2021
<u>AP</u>		MSN	<u>50MG/VIAL</u>	<u>A212265 001</u>	Aug 28, 2019

PRESCRIPTION DRUG PRODUCT LIST

DECITABINE

INJECTABLE; INTRAVENOUS

DECITABINE

<u>AP</u>	NIVAGEN PHARMS INC	<u>50MG/VIAL</u>	<u>A212117</u>	<u>001</u>	Dec 07, 2020
<u>AP</u>	PHARMASCIENCE INC	<u>50MG/VIAL</u>	<u>A204607</u>	<u>001</u>	May 31, 2017
<u>AP</u>	QILU PHARM HAINAN	<u>50MG/VIAL</u>	<u>A212826</u>	<u>001</u>	Apr 12, 2021
<u>AP</u>	SAGENT PHARMS INC	<u>50MG/VIAL</u>	<u>A207100</u>	<u>001</u>	Mar 16, 2018
<u>AP</u>	SANDOZ INC	<u>50MG/VIAL</u>	<u>A202969</u>	<u>001</u>	Aug 28, 2014
<u>AP</u>	WOCKHARDT BIO AG	<u>50MG/VIAL</u>	<u>A209056</u>	<u>001</u>	Apr 09, 2019
<u>AP</u>	ZYDUS PHARMS	<u>50MG/VIAL</u>	<u>A214486</u>	<u>001</u>	Nov 19, 2021

POWDER; INTRAVENOUS

DECITABINE

+	SUN PHARM	50MG/VIAL	N205582	001	Jan 28, 2014
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DEFERASIROX

GRANULE; ORAL

DEFERASIROX

<u>AB</u>	ALKEM LABS LTD	<u>90MG</u>	<u>A213374</u>	<u>001</u>	Jul 14, 2020
<u>AB</u>		<u>180MG</u>	<u>A213374</u>	<u>002</u>	Jul 14, 2020
<u>AB</u>		<u>360MG</u>	<u>A213374</u>	<u>003</u>	Jul 14, 2020
<u>AB</u>	AMNEAL	<u>90MG</u>	<u>A214194</u>	<u>003</u>	Aug 02, 2021
<u>AB</u>		<u>180MG</u>	<u>A214194</u>	<u>001</u>	Feb 09, 2021
<u>AB</u>		<u>360MG</u>	<u>A214194</u>	<u>002</u>	Feb 09, 2021
<u>AB</u>	AUCTA	<u>90MG</u>	<u>A214559</u>	<u>001</u>	Mar 09, 2021
<u>AB</u>		<u>180MG</u>	<u>A214559</u>	<u>002</u>	Mar 09, 2021
<u>AB</u>		<u>360MG</u>	<u>A214559</u>	<u>003</u>	Mar 09, 2021
<u>AB</u>	MSN	<u>180MG</u>	<u>A214650</u>	<u>001</u>	Mar 17, 2021
<u>AB</u>		<u>360MG</u>	<u>A214650</u>	<u>002</u>	Mar 17, 2021
<u>AB</u>	TEVA PHARMS USA	<u>90MG</u>	<u>A214180</u>	<u>001</u>	Nov 19, 2021
<u>AB</u>		<u>180MG</u>	<u>A214180</u>	<u>002</u>	Nov 19, 2021
<u>AB</u>		<u>360MG</u>	<u>A214180</u>	<u>003</u>	Nov 19, 2021

JADENU SPRINKLE

<u>AB</u>	+	NOVARTIS	<u>90MG</u>	<u>N207968</u>	<u>001</u>	May 18, 2017
<u>AB</u>	+		<u>180MG</u>	<u>N207968</u>	<u>002</u>	May 18, 2017
<u>AB</u>	+	!	<u>360MG</u>	<u>N207968</u>	<u>003</u>	May 18, 2017

TABLET; ORAL

DEFERASIROX

<u>AB</u>	ACTAVIS ELIZABETH	<u>90MG</u>	<u>A208697</u>	<u>001</u>	Dec 13, 2019
<u>AB</u>		<u>180MG</u>	<u>A208697</u>	<u>002</u>	Dec 13, 2019
<u>AB</u>		<u>360MG</u>	<u>A208697</u>	<u>003</u>	Dec 13, 2019
<u>AB</u>	ALEMBIC PHARMS LTD	<u>90MG</u>	<u>A211824</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>		<u>180MG</u>	<u>A211824</u>	<u>003</u>	Jun 15, 2020
<u>AB</u>		<u>360MG</u>	<u>A211824</u>	<u>002</u>	Nov 20, 2019
<u>AB</u>	ALKEM LABS LTD	<u>90MG</u>	<u>A210555</u>	<u>001</u>	Mar 30, 2020
<u>AB</u>		<u>180MG</u>	<u>A210555</u>	<u>003</u>	Jul 02, 2020
<u>AB</u>		<u>360MG</u>	<u>A210555</u>	<u>002</u>	Mar 30, 2020
<u>AB</u>	AMNEAL	<u>180MG</u>	<u>A210727</u>	<u>003</u>	Jun 15, 2020
<u>AB</u>	ANNORA PHARMA	<u>90MG</u>	<u>A214341</u>	<u>001</u>	May 14, 2021
<u>AB</u>		<u>180MG</u>	<u>A214341</u>	<u>002</u>	May 14, 2021
<u>AB</u>		<u>360MG</u>	<u>A214341</u>	<u>003</u>	May 14, 2021
<u>AB</u>	CELLTRION	<u>90MG</u>	<u>A212669</u>	<u>001</u>	May 27, 2021
<u>AB</u>		<u>180MG</u>	<u>A212669</u>	<u>002</u>	May 27, 2021
<u>AB</u>		<u>360MG</u>	<u>A212669</u>	<u>003</u>	May 27, 2021
<u>AB</u>	CIPLA	<u>90MG</u>	<u>A211852</u>	<u>001</u>	Feb 11, 2020
<u>AB</u>		<u>180MG</u>	<u>A211852</u>	<u>003</u>	Jun 15, 2020
<u>AB</u>		<u>360MG</u>	<u>A211852</u>	<u>002</u>	Feb 11, 2020
<u>AB</u>	MSN	<u>90MG</u>	<u>A210945</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>		<u>180MG</u>	<u>A210945</u>	<u>003</u>	Jun 16, 2020
<u>AB</u>		<u>360MG</u>	<u>A210945</u>	<u>002</u>	Nov 20, 2019
<u>AB</u>	PIRAMAL HLTHCARE UK	<u>90MG</u>	<u>A212995</u>	<u>001</u>	Dec 30, 2019
<u>AB</u>		<u>180MG</u>	<u>A212995</u>	<u>003</u>	Jun 15, 2020
<u>AB</u>		<u>360MG</u>	<u>A212995</u>	<u>002</u>	Dec 30, 2019
<u>AB</u>	SUN PHARM	<u>90MG</u>	<u>A211641</u>	<u>001</u>	Jan 02, 2020
<u>AB</u>		<u>180MG</u>	<u>A211641</u>	<u>003</u>	Jun 15, 2020
<u>AB</u>		<u>360MG</u>	<u>A211641</u>	<u>002</u>	Jan 02, 2020
<u>AB</u>	TEVA PHARMS USA	<u>90MG</u>	<u>A209223</u>	<u>001</u>	Nov 25, 2019
<u>AB</u>		<u>180MG</u>	<u>A209223</u>	<u>003</u>	Apr 24, 2020
<u>AB</u>		<u>360MG</u>	<u>A209223</u>	<u>002</u>	Nov 25, 2019
<u>AB</u>	ZYDUS PHARMS	<u>90MG</u>	<u>A211383</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>		<u>180MG</u>	<u>A211383</u>	<u>003</u>	Jun 15, 2020
<u>AB</u>		<u>360MG</u>	<u>A211383</u>	<u>002</u>	Nov 20, 2019

PRESCRIPTION DRUG PRODUCT LIST

DEFERASIROX

TABLET; ORAL

JADENU

<u>AB</u>	+	NOVARTIS PHARMS CORP	<u>90MG</u>	<u>N206910</u>	<u>001</u>	Mar 30, 2015
<u>AB</u>	+		<u>180MG</u>	<u>N206910</u>	<u>002</u>	Mar 30, 2015
<u>AB</u>	+	!	<u>360MG</u>	<u>N206910</u>	<u>003</u>	Mar 30, 2015

TABLET, FOR SUSPENSION; ORAL

DEFERASIROX

<u>AB</u>		ACTAVIS ELIZABETH	<u>125MG</u>	<u>A203560</u>	<u>001</u>	Jan 26, 2016
<u>AB</u>			<u>250MG</u>	<u>A203560</u>	<u>002</u>	Jan 26, 2016
<u>AB</u>			<u>500MG</u>	<u>A203560</u>	<u>003</u>	Jan 26, 2016
<u>AB</u>		ALEMBIC PHARMS LTD	<u>125MG</u>	<u>A210060</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>			<u>250MG</u>	<u>A210060</u>	<u>002</u>	Nov 20, 2019
<u>AB</u>			<u>500MG</u>	<u>A210060</u>	<u>003</u>	Nov 20, 2019
<u>AB</u>		ALKEM LABS LTD	<u>125MG</u>	<u>A210519</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>			<u>250MG</u>	<u>A210519</u>	<u>002</u>	Nov 20, 2019
<u>AB</u>			<u>500MG</u>	<u>A210519</u>	<u>003</u>	Nov 20, 2019
<u>AB</u>		BIONPHARMA INC	<u>125MG</u>	<u>A210920</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>			<u>250MG</u>	<u>A210920</u>	<u>002</u>	Nov 20, 2019
<u>AB</u>			<u>500MG</u>	<u>A210920</u>	<u>003</u>	Nov 20, 2019
<u>AB</u>		ICHNOS	<u>125MG</u>	<u>A209433</u>	<u>001</u>	Jan 06, 2020
<u>AB</u>			<u>250MG</u>	<u>A209433</u>	<u>002</u>	Jan 06, 2020
<u>AB</u>			<u>500MG</u>	<u>A209433</u>	<u>003</u>	Jan 06, 2020
<u>AB</u>		MSN	<u>125MG</u>	<u>A209878</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>			<u>250MG</u>	<u>A209878</u>	<u>002</u>	Nov 20, 2019
<u>AB</u>			<u>500MG</u>	<u>A209878</u>	<u>003</u>	Nov 20, 2019
<u>AB</u>		SUN PHARM	<u>125MG</u>	<u>A209782</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>			<u>250MG</u>	<u>A209782</u>	<u>002</u>	Nov 20, 2019
<u>AB</u>			<u>500MG</u>	<u>A209782</u>	<u>003</u>	Nov 20, 2019
<u>EXJADE</u>						
<u>AB</u>	+	NOVARTIS	<u>125MG</u>	<u>N021882</u>	<u>001</u>	Nov 02, 2005
<u>AB</u>	+		<u>250MG</u>	<u>N021882</u>	<u>002</u>	Nov 02, 2005
<u>AB</u>	+	!	<u>500MG</u>	<u>N021882</u>	<u>003</u>	Nov 02, 2005

DEFERIPRONE

SOLUTION; ORAL

FERRIPROX

+! CHIESI

100MG/ML

N208030 001 Sep 09, 2015

TABLET; ORAL

DEFERIPRONE

<u>AB</u>		HIKMA	<u>500MG</u>	<u>A213239</u>	<u>001</u>	Mar 29, 2021
<u>AB</u>		TARO PHARM INDS LTD	<u>500MG</u>	<u>A208800</u>	<u>001</u>	Feb 08, 2019
<u>FERRIPROX</u>						
<u>AB</u>	+	CHIESI	<u>500MG</u>	<u>N021825</u>	<u>001</u>	Oct 14, 2011
	+		1GM	N021825	002	Jul 25, 2019
	+	!	1GM	N212269	001	May 19, 2020

DEFEROXAMINE MESYLATE

INJECTABLE; INJECTION

DEFEROXAMINE MESYLATE

<u>AP</u>		FRESENIUS KABI USA	<u>500MG/VIAL</u>	<u>A078718</u>	<u>001</u>	Sep 15, 2009
<u>AP</u>			<u>2GM/VIAL</u>	<u>A078718</u>	<u>002</u>	Sep 15, 2009
<u>AP</u>		GLAND PHARMA LTD	<u>500MG/VIAL</u>	<u>A207384</u>	<u>001</u>	Sep 29, 2017
<u>AP</u>			<u>2GM/VIAL</u>	<u>A207384</u>	<u>002</u>	Sep 29, 2017
<u>AP</u>		HOSPIRA	<u>500MG/VIAL</u>	<u>A076019</u>	<u>001</u>	Mar 17, 2004
<u>AP</u>	!		<u>2GM/VIAL</u>	<u>A076019</u>	<u>002</u>	Mar 17, 2004
<u>AP</u>		WEST-WARD PHARMS INT	<u>500MG/VIAL</u>	<u>A078086</u>	<u>001</u>	May 30, 2007
<u>AP</u>			<u>2GM/VIAL</u>	<u>A078086</u>	<u>002</u>	May 30, 2007
<u>DESFERAL</u>						
<u>AP</u>	+	NOVARTIS	<u>500MG/VIAL</u>	<u>N016267</u>	<u>001</u>	

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

PRESCRIPTION DRUG PRODUCT LISTDEFLAZACORT

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
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TABLET; ORAL

BAXDELA

+	!	MELINTA	EQ 450MG BASE	N208610	001	Jun 19, 2017
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DEMECLOCYCLINE HYDROCHLORIDE

TABLET; ORAL

DEMECLOCYCLINE HYDROCHLORIDE

AB	AKORN	150MG	A065389	001	Dec 01, 2008
AB		300MG	A065389	002	Dec 01, 2008
AB	AMNEAL PHARM	150MG	A065425	001	Feb 27, 2008
AB	!	300MG	A065425	002	Feb 27, 2008
AB	EPIC PHARMA LLC	150MG	A065447	001	Aug 18, 2015
AB		300MG	A065447	002	Aug 18, 2015

DEOXYCHOLIC ACID

SOLUTION; SUBCUTANEOUS

DEOXYCHOLIC ACID

AP	SLAYBACK PHARMA LLC	20MG/2ML (10MG/ML)	A212296	001	Apr 02, 2021		
	KYBELLA						
AP	+	!	KYTHERA BIOPHARMS	20MG/2ML (10MG/ML)	N206333	001	Apr 29, 2015

DESFLURANE

LIQUID; INHALATION

DESFLURANE

AN	SHANGHAI HENGRUI	100%	A208234	001	Feb 26, 2018		
	SUPRANE						
AN	+	!	BAXTER HLTHCARE	100%	N020118	001	Sep 18, 1992

DESIPRAMINE HYDROCHLORIDE

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

AB	ACTAVIS TOTOWA	10MG	A074430	001	Feb 09, 1996
AB		25MG	A071601	001	Jun 05, 1987
AB		50MG	A071588	001	Jun 05, 1987
AB		75MG	A071602	001	Oct 05, 1987
AB		100MG	A071766	001	Oct 05, 1987
AB		150MG	A074430	002	Feb 09, 1996
AB	ALEMBIC PHARMS LTD	10MG	A209785	001	Jul 07, 2021
AB		25MG	A209785	002	Jul 07, 2021
AB		50MG	A209785	003	Jul 07, 2021
AB		75MG	A209785	004	Jul 07, 2021
AB		100MG	A209785	005	Jul 07, 2021
AB		150MG	A209785	006	Jul 07, 2021
AB	AMNEAL PHARMS CO	10MG	A208105	001	Mar 17, 2016
AB		25MG	A208105	002	Mar 17, 2016
AB		50MG	A208105	003	Mar 17, 2016
AB		75MG	A208105	004	Mar 17, 2016
AB		100MG	A208105	005	Mar 17, 2016
AB		150MG	A208105	006	Mar 17, 2016
AB	HERITAGE PHARMS INC	10MG	A207433	001	May 05, 2016
AB		25MG	A207433	002	May 05, 2016
AB		50MG	A207433	003	May 05, 2016
AB		75MG	A207433	004	May 05, 2016
AB		100MG	A207433	005	May 05, 2016
AB		150MG	A207433	006	May 05, 2016
AB	NOVAST LABS	10MG	A204963	001	Dec 26, 2017
AB		25MG	A204963	002	Dec 26, 2017

PRESCRIPTION DRUG PRODUCT LIST

DESIPRAMINE HYDROCHLORIDE

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

<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

NORPRAMIN

<u>AB</u>	+	VALIDUS PHARMS	<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>	
<u>AB</u>	+		<u>100MG</u>	<u>N014399</u>	<u>005</u>	
<u>AB</u>	+		<u>150MG</u>	<u>N014399</u>	<u>006</u>	

DESLORATADINE

TABLET; ORAL

CLARINEX

<u>AB</u>	+	ORGANON	<u>5MG</u>	<u>N021165</u>	<u>001</u>	Dec 21, 2001
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DESLORATADINE

<u>AB</u>		BELCHER PHARMS	<u>5MG</u>	<u>A078355</u>	<u>001</u>	Apr 19, 2012
<u>AB</u>		DR REDDYS LABS LTD	<u>5MG</u>	<u>A078365</u>	<u>001</u>	Mar 08, 2011
<u>AB</u>		LUPIN PHARMS	<u>5MG</u>	<u>A078352</u>	<u>001</u>	Oct 25, 2010
<u>AB</u>		ORBION PHARMS	<u>5MG</u>	<u>A078357</u>	<u>001</u>	Feb 19, 2010
<u>AB</u>		PERRIGO	<u>5MG</u>	<u>A078361</u>	<u>001</u>	Dec 22, 2011
<u>AB</u>		SANDOZ	<u>5MG</u>	<u>A078364</u>	<u>001</u>	Dec 03, 2010

TABLET, ORALLY DISINTEGRATING; ORAL

DESLORATADINE

		REDDYS	2.5MG	A078367	001	Jul 12, 2010
!			5MG	A078367	002	Jul 12, 2010

DESLORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

CLARINEX-D 12 HOUR

	+	ORGANON	2.5MG;120MG	N021313	001	Feb 01, 2006
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DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR

!		DR REDDYS LABS LTD	5MG;240MG	A078366	001	Apr 26, 2011
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DESMOPRESSIN ACETATE

INJECTABLE; INJECTION

DDAVP

<u>AP</u>	+	FERRING PHARMS INC	<u>0.004MG/ML</u>	<u>N018938</u>	<u>001</u>	Mar 30, 1984
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DESMOPRESSIN ACETATE

<u>AP</u>		AM REGENT	<u>0.004MG/ML</u>	<u>A091374</u>	<u>001</u>	Feb 14, 2019
<u>AP</u>		SAGENT PHARMS INC	<u>0.004MG/ML</u>	<u>A204695</u>	<u>001</u>	Aug 22, 2017
<u>AP</u>			<u>0.004MG/ML</u>	<u>A204751</u>	<u>001</u>	Aug 22, 2017
<u>AP</u>		SUN PHARM INDS LTD	<u>0.004MG/ML</u>	<u>A091280</u>	<u>001</u>	Jan 25, 2013
<u>AP</u>		TEVA PHARMS USA	<u>0.004MG/ML</u>	<u>A074888</u>	<u>001</u>	Oct 15, 1997
<u>AP</u>		UBI	<u>0.004MG/ML</u>	<u>A210218</u>	<u>001</u>	Feb 14, 2020
<u>AP</u>			<u>0.004MG/ML</u>	<u>A210223</u>	<u>001</u>	Sep 17, 2020

SPRAY, METERED; NASAL

DESMOPRESSIN ACETATE

<u>AB</u>	!	BAUSCH AND LOMB	<u>0.01MG/SPRAY</u>	<u>A074830</u>	<u>001</u>	Jan 25, 1999
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DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION)

<u>AB</u>		APOTEX INC	<u>0.01MG/SPRAY</u>	<u>A076703</u>	<u>001</u>	Jan 27, 2005
<u>AB</u>	!	SUN PHARM	<u>0.01MG/SPRAY</u>	<u>A078271</u>	<u>001</u>	Dec 23, 2013
<u>AB</u>		ZYDUS PHARMS	<u>0.01MG/SPRAY</u>	<u>A091345</u>	<u>001</u>	Oct 03, 2017

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>		ABHAI LLC	<u>0.1MG</u>	<u>A210371</u>	<u>001</u>	Jan 28, 2019
<u>AB</u>			<u>0.2MG</u>	<u>A210371</u>	<u>002</u>	Jan 28, 2019
<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

PRESCRIPTION DRUG PRODUCT LIST

DESMOPRESSIN ACETATE

TABLET; ORAL

DESMOPRESSIN ACETATE

AB	GLENMARK PHARMS LTD	<u>0.1MG</u>	<u>A201831 001</u>	May 28, 2015
AB		<u>0.2MG</u>	<u>A201831 002</u>	May 28, 2015
AB	HERITAGE PHARMA	<u>0.1MG</u>	<u>A207880 001</u>	May 26, 2017
AB		<u>0.2MG</u>	<u>A207880 002</u>	May 26, 2017
AB	NOVAST LABS	<u>0.1MG</u>	<u>A208357 001</u>	Jun 06, 2019
AB		<u>0.2MG</u>	<u>A208357 002</u>	Jun 06, 2019

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

AB	LUPIN LTD	<u>0.15MG, N/A; 0.02MG, 0.01MG</u>	<u>A202226 001</u>	Aug 12, 2015
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CYCLESSA

AB	+	!	ASPEN GLOBAL INC	<u>0.1MG, 0.125MG, 0.15MG; 0.025MG, 0.025MG, 0.025MG</u>	<u>N021090 001</u>	Dec 20, 2000
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DESOGESTREL AND ETHINYL ESTRADIOL

AB	!	DURAMED PHARMS BARR	<u>0.15MG; 0.03MG</u>	<u>A075256 002</u>	Aug 12, 1999
AB		MAYNE PHARMA	<u>0.15MG, N/A; 0.02MG, 0.01MG</u>	<u>A076916 001</u>	Dec 29, 2008
AB			<u>0.1MG, 0.125MG, 0.15MG; 0.025MG, 0.025MG, 0.025MG</u>	<u>A077182 001</u>	Jan 24, 2006
AB		MYLAN LABS LTD	<u>0.15MG, N/A; 0.02MG, 0.01MG</u>	<u>A202296 001</u>	Aug 30, 2013
AB		NAARI PTE LTD	<u>0.15MG, N/A; 0.02MG, 0.01MG</u>	<u>A209170 001</u>	Jun 05, 2017
AB		NOVAST LABS	<u>0.15MG; 0.03MG</u>	<u>A091234 001</u>	Jul 12, 2013
AB		WATSON LABS	<u>0.15MG; 0.03MG</u>	<u>A076915 001</u>	Jul 29, 2005

ENSKYCE

AB	LUPIN LTD	<u>0.15MG; 0.03MG</u>	<u>A201887 001</u>	Mar 07, 2013
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ISIBLOOM

AB	XIROMED	<u>0.15MG; 0.03MG</u>	<u>A202789 001</u>	Aug 12, 2015
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KALLIGA

AB	AUROBINDO PHARMA LTD	<u>0.15MG; 0.03MG</u>	<u>A207081 001</u>	May 17, 2017
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KARIVA

AB	!	BARR	<u>0.15MG, N/A; 0.02MG, 0.01MG</u>	<u>A075863 001</u>	Apr 05, 2002
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KIMIDESS

AB	VINTAGE PHARMS	<u>0.15MG, N/A; 0.02MG, 0.01MG</u>	<u>A076681 001</u>	Apr 30, 2015
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PIMTREA

AB	NOVAST LABS	<u>0.15MG, N/A; 0.02MG, 0.01MG</u>	<u>A091247 001</u>	Aug 01, 2013
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SIMLIYA

AB	AUROBINDO PHARMA LTD	<u>0.15MG, N/A; 0.02MG, 0.01MG</u>	<u>A206853 001</u>	Mar 22, 2017
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VELIVET

AB	DURAMED PHARMS BARR	<u>0.1MG, 0.125MG, 0.15MG; 0.025MG, 0.025MG, 0.025MG</u>	<u>A076455 001</u>	Feb 24, 2004
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VIORELE

AB	GLENMARK GENERICS	<u>0.15MG, N/A; 0.02MG, 0.01MG</u>	<u>A091346 001</u>	Apr 02, 2012
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VOLNEA

AB	XIROMED	<u>0.15MG, N/A; 0.02MG, 0.01MG</u>	<u>A202689 001</u>	Sep 09, 2016
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DESONIDE

AEROSOL, FOAM; TOPICAL

VERDESO

+	!	ALMIRALL	0.05%	N021978 001	Sep 19, 2006
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CREAM; TOPICAL

DESONIDE

AB	CADILA	<u>0.05%</u>	<u>A210198 001</u>	Nov 20, 2019	
AB	COSETTE	<u>0.05%</u>	<u>A074027 001</u>	Sep 28, 1992	
AB	GLENMARK PHARMS	<u>0.05%</u>	<u>A209729 001</u>	Jul 24, 2017	
AB	+	!	PADAGIS US	<u>0.05%</u>	<u>N017010 001</u>
AB	TARO	<u>0.05%</u>	<u>A073548 001</u>	Jun 30, 1992	

DESOWEN

AB	GALDERMA LABS LP	<u>0.05%</u>	<u>N019048 001</u>	Dec 14, 1984
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GEL; TOPICAL

DESONIDE

!	CINTEX SVCS	0.05%	A202470 001	May 11, 2020
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LOTION; TOPICAL

DESONIDE

AB	ALEOR	<u>0.05%</u>	<u>A213632 001</u>	Aug 24, 2020
AB	DERMACEUTICALS	<u>0.05%</u>	<u>A075860 001</u>	Mar 19, 2002

PRESCRIPTION DRUG PRODUCT LIST

DESONIDE

LOTION; TOPICAL

DESONIDE

<u>AB</u>	GLENMARK PHARMS	<u>0.05%</u>	<u>A209494</u>	<u>001</u>	Sep 26, 2017
<u>AB</u>	TARO	<u>0.05%</u>	<u>A202161</u>	<u>001</u>	Oct 31, 2014
<u>AB</u>	TELIGENT	<u>0.05%</u>	<u>A207855</u>	<u>001</u>	Sep 28, 2017

DESOWEN

<u>AB</u>	! GALDERMA LABS LP	<u>0.05%</u>	<u>A072354</u>	<u>001</u>	Jan 24, 1992
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OINTMENT; TOPICAL

DESONIDE

<u>AB</u>	AKORN	<u>0.05%</u>	<u>A208836</u>	<u>001</u>	Mar 27, 2017
<u>AB</u>	ALEOR	<u>0.05%</u>	<u>A212473</u>	<u>001</u>	Oct 23, 2019
<u>AB</u>	DERMACEUTICALS				
<u>AB</u>	ENCUBE ETHICALS	<u>0.05%</u>	<u>A210998</u>	<u>001</u>	Jan 30, 2019
<u>AB</u>	FOUGERA PHARMS	<u>0.05%</u>	<u>A075751</u>	<u>001</u>	Mar 12, 2001
<u>AB</u>	GLENMARK PHARMS LTD	<u>0.05%</u>	<u>A209996</u>	<u>001</u>	Sep 15, 2017
<u>AB</u>	+! PADAGIS US	<u>0.05%</u>	<u>N017426</u>	<u>001</u>	
<u>AB</u>	TARO	<u>0.05%</u>	<u>A074254</u>	<u>001</u>	Aug 03, 1994
<u>AB</u>	TELIGENT	<u>0.05%</u>	<u>A212002</u>	<u>001</u>	Mar 12, 2019

DESOWEN

<u>AB</u>	GALDERMA LABS LP	<u>0.05%</u>	<u>A071425</u>	<u>001</u>	Jun 15, 1988
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DESOXIMETASONE

CREAM; TOPICAL

DESOXIMETASONE

<u>AB</u>	ACTAVIS MID	<u>0.25%</u>	<u>A205082</u>	<u>001</u>	Sep 04, 2015
<u>AB</u>	ATLANTIC				
<u>AB</u>	CADILA	<u>0.25%</u>	<u>A205620</u>	<u>001</u>	Sep 28, 2018
<u>AB</u>	COSETTE	<u>0.25%</u>	<u>A209595</u>	<u>001</u>	Mar 04, 2020
<u>AB</u>	FOUGERA PHARMS	<u>0.25%</u>	<u>A078369</u>	<u>001</u>	Jun 29, 2010
<u>AB</u>	LUPIN	<u>0.05%</u>	<u>A208163</u>	<u>001</u>	Jan 10, 2017
<u>AB</u>		<u>0.25%</u>	<u>A208164</u>	<u>001</u>	Jan 09, 2017
<u>AB</u>	PADAGIS ISRAEL	<u>0.25%</u>	<u>A076510</u>	<u>001</u>	Jul 01, 2003
<u>AB</u>	RISING	<u>0.05%</u>	<u>A210980</u>	<u>001</u>	Dec 21, 2018
<u>AB</u>		<u>0.25%</u>	<u>A205594</u>	<u>001</u>	Jul 02, 2018

TOPICORT

<u>AB</u>	! TARO PHARM INDS LTD	<u>0.05%</u>	<u>A073210</u>	<u>001</u>	Nov 30, 1990
<u>AB</u>	!	<u>0.25%</u>	<u>A073193</u>	<u>001</u>	Nov 30, 1990

GEL; TOPICAL

DESOXIMETASONE

<u>AB</u>	AKORN	<u>0.05%</u>	<u>A090727</u>	<u>001</u>	Mar 10, 2011
<u>AB</u>	PADAGIS US	<u>0.05%</u>	<u>A077552</u>	<u>001</u>	Jan 09, 2006
<u>AB</u>	RISING	<u>0.05%</u>	<u>A204675</u>	<u>001</u>	Aug 12, 2016

TOPICORT

<u>AB</u>	! TARO PHARM INDS LTD	<u>0.05%</u>	<u>A074904</u>	<u>001</u>	Jul 14, 1998
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OINTMENT; TOPICAL

DESOXIMETASONE

<u>AB</u>	ACTAVIS MID	<u>0.25%</u>	<u>A204965</u>	<u>001</u>	Nov 07, 2016
<u>AB</u>	ATLANTIC				
<u>AB</u>	AKORN	<u>0.25%</u>	<u>A201005</u>	<u>001</u>	Apr 24, 2014
<u>AB</u>	CADILA	<u>0.25%</u>	<u>A205206</u>	<u>001</u>	Sep 19, 2017
<u>AB</u>	COSETTE	<u>0.25%</u>	<u>A206740</u>	<u>001</u>	Dec 23, 2016
<u>AB</u>	FOUGERA PHARMS	<u>0.25%</u>	<u>A078657</u>	<u>001</u>	Sep 28, 2012
<u>AB</u>	! GLENMARK GENERICS	<u>0.25%</u>	<u>A202838</u>	<u>001</u>	Sep 20, 2013
<u>AB</u>	LUPIN	<u>0.05%</u>	<u>A208044</u>	<u>001</u>	Dec 12, 2016
<u>AB</u>		<u>0.25%</u>	<u>A208104</u>	<u>001</u>	Dec 01, 2016
<u>AB</u>	NOVEL LABS INC	<u>0.25%</u>	<u>A206792</u>	<u>001</u>	May 10, 2016
<u>AB</u>	PADAGIS ISRAEL	<u>0.25%</u>	<u>A077770</u>	<u>001</u>	Apr 20, 2015
<u>AB</u>	RISING	<u>0.25%</u>	<u>A204272</u>	<u>001</u>	Nov 30, 2016
<u>AB</u>	TELIGENT	<u>0.05%</u>	<u>A209973</u>	<u>001</u>	Oct 23, 2018
<u>AB</u>		<u>0.25%</u>	<u>A208101</u>	<u>001</u>	Feb 25, 2016

TOPICORT

<u>AB</u>	+! TARO PHARM INDS LTD	<u>0.05%</u>	<u>N018594</u>	<u>001</u>	Jan 17, 1985
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SPRAY; TOPICAL

DESOXIMETASONE

<u>AT</u>	LUPIN	<u>0.25%</u>	<u>A208124</u>	<u>001</u>	Mar 16, 2018
<u>AT</u>	PADAGIS ISRAEL	<u>0.25%</u>	<u>A206441</u>	<u>001</u>	Jan 20, 2017

TOPICORT

<u>AT</u>	+! TARO PHARMS	<u>0.25%</u>	<u>N204141</u>	<u>001</u>	Apr 11, 2013
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PRESCRIPTION DRUG PRODUCT LIST

DESVENLAFAXINE

TABLET, EXTENDED RELEASE;ORAL

DESVENLAFAXINE

+	ALEMBIC PHARMS LTD	50MG	N204150	001	Mar 04, 2013
+	!	100MG	N204150	002	Mar 04, 2013

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE;ORAL

DESVENLAFAXINE SUCCINATE

AB	ACTAVIS LABS FL	<u>EQ 25MG BASE</u>	<u>A204065</u>	<u>001</u>	Jul 29, 2016
AB		<u>EQ 50MG BASE</u>	<u>A204065</u>	<u>002</u>	Jul 29, 2016
AB		<u>EQ 100MG BASE</u>	<u>A204065</u>	<u>003</u>	Jul 29, 2016
AB	ALEMBIC PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A204003</u>	<u>003</u>	Sep 14, 2018
AB		<u>EQ 50MG BASE</u>	<u>A204003</u>	<u>001</u>	Jun 29, 2015
AB		<u>EQ 100MG BASE</u>	<u>A204003</u>	<u>002</u>	Jun 29, 2015
AB	HIKMA	<u>EQ 25MG BASE</u>	<u>A204082</u>	<u>002</u>	Aug 28, 2017
AB		<u>EQ 50MG BASE</u>	<u>A204082</u>	<u>001</u>	Feb 16, 2016
AB		<u>EQ 100MG BASE</u>	<u>A204083</u>	<u>001</u>	Feb 16, 2016
AB	INTELLIPHARMACEUTIC S	<u>EQ 50MG BASE</u>	<u>A204805</u>	<u>001</u>	May 07, 2019
AB		<u>EQ 100MG BASE</u>	<u>A204805</u>	<u>002</u>	May 07, 2019
AB	LUPIN LTD	<u>EQ 50MG BASE</u>	<u>A204172</u>	<u>001</u>	Jun 29, 2015
AB		<u>EQ 100MG BASE</u>	<u>A204172</u>	<u>002</u>	Jun 29, 2015
AB	RUBICON	<u>EQ 50MG BASE</u>	<u>A204028</u>	<u>001</u>	Jun 29, 2015
AB		<u>EQ 100MG BASE</u>	<u>A204028</u>	<u>002</u>	Jun 29, 2015
AB	YICHANG HUMANWELL	<u>EQ 25MG BASE</u>	<u>A210014</u>	<u>003</u>	Oct 13, 2020
AB		<u>EQ 50MG BASE</u>	<u>A210014</u>	<u>001</u>	Oct 01, 2018
AB		<u>EQ 100MG BASE</u>	<u>A210014</u>	<u>002</u>	Oct 01, 2018
AB	ZYDUS PHARMS	<u>EQ 50MG BASE</u>	<u>A204020</u>	<u>001</u>	Oct 11, 2017
AB		<u>EQ 100MG BASE</u>	<u>A204020</u>	<u>002</u>	Oct 11, 2017

PRISTIQ

AB	+	PF PRISM CV	<u>EQ 25MG BASE</u>	<u>N021992</u>	<u>003</u>	Aug 20, 2014
AB	+		<u>EQ 50MG BASE</u>	<u>N021992</u>	<u>001</u>	Feb 29, 2008
AB	+	!	<u>EQ 100MG BASE</u>	<u>N021992</u>	<u>002</u>	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

!	HIKMA	1MG/ML	A088252	001	Sep 01, 1983
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ELIXIR;ORAL

DEXAMETHASONE

AA	!	ANIMA	<u>0.5MG/5ML</u>	<u>A084754</u>	<u>001</u>	
AA		LANNETT CO INC	<u>0.5MG/5ML</u>	<u>A091188</u>	<u>001</u>	May 11, 2011
AA		LYNE	<u>0.5MG/5ML</u>	<u>A090891</u>	<u>001</u>	Jul 12, 2011

IMPLANT;INTRAVITREAL

OZURDEX

+	!	ALLERGAN	0.7MG	N022315	001	Jun 17, 2009
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INSERT;OPHTHALMIC

DEXTENZA

+	!	OCULAR THERAPEUTIX	0.4MG	N208742	001	Nov 30, 2018
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SOLUTION;ORAL

DEXAMETHASONE

!	HIKMA	0.5MG/5ML	A088248	001	Sep 01, 1983
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SUSPENSION;INTRAOCULAR

DEXYCU KIT

+	!	EYEPOINT PHARMS	9%	N208912	001	Feb 09, 2018
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SUSPENSION/DROPS;OPHTHALMIC

MAXIDEX

+	!	NOVARTIS	0.1%	N013422	001	
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TABLET;ORAL

DEXAMETHASONE

AB		AMNEAL	<u>4MG</u>	<u>A215106</u>	<u>001</u>	Oct 14, 2021
AB			<u>6MG</u>	<u>A215106</u>	<u>002</u>	Oct 14, 2021
AB	+	HIKMA	<u>1.5MG</u>	<u>A084610</u>	<u>001</u>	
AB	+		<u>4MG</u>	<u>A084612</u>	<u>001</u>	
AB	+	!	<u>6MG</u>	<u>A088316</u>	<u>001</u>	Sep 15, 1983
AB		LARKEN LABS INC	<u>1.5MG</u>	<u>A201270</u>	<u>001</u>	Jul 17, 2017

PRESCRIPTION DRUG PRODUCT LIST

DEXAMETHASONE

TABLET; ORAL

DEXAMETHASONE

BP	ALVOGEN	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	HIKMA	0.5MG	A084611 001	
BP	+	0.75MG	A084613 001	
BP	+	1MG	A088306 001	Sep 15, 1983
BP	+	2MG	A087916 001	Aug 26, 1982
BP	XSPIRE PHARMA	1.5MG	A088237 001	Apr 28, 1983
	HEMADY			
	+	DEXCEL PHARMA	20MG	N211379 001 Oct 03, 2019

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

DEXAMETHASONE SODIUM PHOSPHATE

AP	AMNEAL	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A208689 001</u>	Aug 22, 2018	
AP	EUGIA PHARMA	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A206781 001</u>	Dec 01, 2015	
AP		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A210966 001</u>	Jun 05, 2020	
AP		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A210967 001</u>	Jun 07, 2019	
AP	!	FRESENIUS KABI USA	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A084916 001</u>	
AP			<u>EQ 4MG PHOSPHATE/ML</u>	<u>A203129 001</u>	Sep 30, 2015
AP	!		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A040572 001</u>	Apr 22, 2005
AP			<u>EQ 10MG PHOSPHATE/ML</u>	<u>A209192 001</u>	Jul 06, 2018
AP	GLAND PHARMA LTD	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A215654 001</u>	Aug 04, 2021	
AP	MYLAN LABS LTD	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A040803 001</u>	Aug 29, 2008	
AP		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A040802 001</u>	Aug 29, 2008	
AP	SOMERSET	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A207521 001</u>	Jun 08, 2018	
AP	SOMERSET THERAPS	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A211036 001</u>	May 10, 2019	
	LLC				
AP	WEST-WARD PHARMS	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A084282 001</u>		
	INT				
AP	!	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A087702 001</u>	Sep 07, 1982	

DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE

AP	AMNEAL	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A208690 001</u>	Aug 22, 2018	
AP	!	FRESENIUS KABI USA	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A040491 001</u>	Apr 11, 2003
AP	SOMERSET THERAPS	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A207442 001</u>	Apr 19, 2018	
	LLC				

SOLUTION/DROPS; OPHTHALMIC, OTIC

DEXAMETHASONE SODIUM PHOSPHATE

!	BAUSCH AND LOMB	EQ 0.1% PHOSPHATE	A040069 001	Jul 26, 1996
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DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

MAXITROL

AT	+	NOVARTIS	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>N050065 002</u>	
			<u>NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE</u>		
AT		BAUSCH AND LOMB	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A064063 001</u>	Jul 25, 1994
AT		PADAGIS US	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A062938 001</u>	Jul 31, 1989

SUSPENSION/DROPS; OPHTHALMIC

DEXASPORIN

AT		BAUSCH AND LOMB	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A064135 001</u>	Sep 13, 1995
			<u>MAXITROL</u>		
AT	+	NOVARTIS	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N050023 002</u>	
AT		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

+	NOVARTIS	0.1%;0.3%	N050616 001	Sep 28, 1988
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SUSPENSION/DROPS; OPHTHALMIC

TOBRADEX

AB	+	NOVARTIS	<u>0.1%;0.3%</u>	<u>N050592 001</u>	Aug 18, 1988
			<u>TOBRAMYCIN AND DEXAMETHASONE</u>		
AB		AMNEAL	<u>0.1%;0.3%</u>	<u>A212991 001</u>	Jul 15, 2021
AB		BAUSCH AND LOMB	<u>0.1%;0.3%</u>	<u>A064134 001</u>	Oct 27, 1999
		TOBRADEX ST			
	+	EYEVANCE	0.05%;0.3%	N050818 001	Feb 13, 2009

PRESCRIPTION DRUG PRODUCT LIST

DEXCHLORPHENIRAMINE MALEATE

SYRUP; ORAL

DEXCHLORPHENIRAMINE MALEATE

<u>AA</u>	<u>+!</u>	WOCKHARDT BIO AG	<u>2MG/5ML</u>	<u>A088251</u>	<u>001</u>	Mar 23, 1984
<u>AA</u>		CAPPELLON PHARMS LLC	<u>2MG/5ML</u>	<u>A202520</u>	<u>001</u>	Jul 16, 2018

DEXLANSOPRAZOLE

CAPSULE, DELAYED RELEASE; ORAL

DEXILANT

<u>AB</u>	<u>+!</u>	TAKEDA PHARMS USA	<u>60MG</u>	<u>N022287</u>	<u>002</u>	Jan 30, 2009
<u>AB</u>		PAR PHARM INC	<u>60MG</u>	<u>A202294</u>	<u>001</u>	Apr 19, 2017
		DEXILANT				
	<u>+</u>	TAKEDA PHARMS USA	<u>30MG</u>	<u>N022287</u>	<u>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</u>	<u>001</u>	Feb 09, 2016
<u>AP</u>		ACTAVIS INC	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A204686</u>	<u>001</u>	Oct 17, 2016
<u>AP</u>		AKORN	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A202585</u>	<u>001</u>	Nov 24, 2014
<u>AP</u>		AMNEAL PHARMS CO	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A207551</u>	<u>001</u>	May 20, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A207551</u>	<u>002</u>	May 20, 2020
<u>AP</u>		BAXTER HLTHCARE CORP	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A208532</u>	<u>001</u>	Aug 21, 2018
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A208532</u>	<u>002</u>	Aug 21, 2018
<u>AP</u>		EUGIA PHARMA	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A205867</u>	<u>001</u>	Mar 17, 2016
<u>AP</u>			<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A210321</u>	<u>001</u>	Dec 07, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A210321</u>	<u>002</u>	Dec 07, 2020
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u>	<u>A208129</u>	<u>001</u>	Nov 29, 2018
<u>AP</u>			<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A201072</u>	<u>001</u>	Sep 18, 2015
<u>AP</u>			<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A208129</u>	<u>002</u>	Nov 29, 2018
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A208129</u>	<u>003</u>	Nov 29, 2018
<u>AP</u>		GLAND	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A202126</u>	<u>001</u>	Aug 20, 2015
<u>AP</u>		GLAND PHARMA LTD	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A209307</u>	<u>001</u>	Jun 03, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A209307</u>	<u>002</u>	Jun 03, 2020
<u>AP</u>		HIKMA	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A205046</u>	<u>001</u>	Apr 26, 2017
<u>AP</u>			<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A206407</u>	<u>001</u>	Jan 30, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A206407</u>	<u>002</u>	Jan 30, 2020
<u>AP</u>		HONG KONG	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A204843</u>	<u>001</u>	Jan 18, 2019
<u>AP</u>		JIANGSU PHARMS	<u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u>	<u>A209065</u>	<u>002</u>	Jun 12, 2020
<u>AP</u>			<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A209065</u>	<u>001</u>	Sep 19, 2017
<u>AP</u>			<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A209065</u>	<u>003</u>	Jun 12, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A209065</u>	<u>004</u>	Jun 12, 2020
<u>AP</u>		MYLAN INSTITUTIONAL	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A202881</u>	<u>001</u>	Aug 18, 2014
<u>AP</u>		MYLAN LABS LTD	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A212571</u>	<u>001</u>	Aug 27, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A212571</u>	<u>002</u>	Aug 27, 2020
<u>AP</u>		PAR STERILE PRODUCTS	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A203972</u>	<u>001</u>	Aug 18, 2014
<u>AP</u>			<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A208266</u>	<u>001</u>	Sep 15, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A208266</u>	<u>002</u>	Sep 15, 2020
<u>AP</u>		PIRAMAL CRITICAL	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A214794</u>	<u>001</u>	Sep 03, 2021
<u>AP</u>		SANDOZ INC	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A091465</u>	<u>001</u>	Jun 14, 2016
<u>AP</u>		SLAYBACK PHARMA LLC	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A212791</u>	<u>001</u>	Dec 04, 2019
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A212791</u>	<u>002</u>	Dec 04, 2019
<u>AP</u>		TAGI	<u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u>	<u>A212857</u>	<u>001</u>	Nov 23, 2020
<u>AP</u>			<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A212857</u>	<u>002</u>	Nov 23, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A212857</u>	<u>003</u>	Nov 23, 2020
<u>AP</u>		TEVA PHARMS USA	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A205272</u>	<u>001</u>	Nov 28, 2017
<u>AP</u>		ZYDUS PHARMS	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A206798</u>	<u>001</u>	Feb 27, 2018
		<u>PRECEDEX</u>				
<u>AP</u>	<u>+!</u>	HOSPIRA	<u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u>	<u>N021038</u>	<u>004</u>	Nov 14, 2014
<u>AP</u>	<u>+!</u>		<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>N021038</u>	<u>001</u>	Dec 17, 1999
<u>AP</u>	<u>+!</u>		<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>N021038</u>	<u>002</u>	Mar 13, 2013
<u>AP</u>	<u>+!</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>N021038</u>	<u>003</u>	Mar 13, 2013
	<u>+!</u>		<u>EQ 1MG BASE/250ML (EQ 4MCG BASE/ML)</u>	<u>N021038</u>	<u>005</u>	Jan 31, 2020

SOLUTION; INTRAVENOUS

DEXMEDETOMIDINE HYDROCHLORIDE

	<u>+!</u>	HQ SPCLT PHARMA	<u>EQ 1MG BASE/10ML (EQ 100MCG BASE/ML)</u>	<u>N206628</u>	<u>002</u>	Oct 21, 2015
	<u>+</u>		<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>N206628</u>	<u>003</u>	Jun 22, 2018
	<u>+</u>		<u>EQ 400MCG BASE/4ML (EQ 100MCG BASE/ML)</u>	<u>N206628</u>	<u>001</u>	Oct 21, 2015
	<u>+</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>N206628</u>	<u>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>	ASCENT PHARMS INC	<u>5MG</u>	<u>A215523 001</u>	Dec 08, 2021
<u>AB</u>		<u>10MG</u>	<u>A215523 002</u>	Dec 08, 2021
<u>AB</u>		<u>15MG</u>	<u>A215523 003</u>	Dec 08, 2021
<u>AB</u>		<u>20MG</u>	<u>A215523 004</u>	Dec 08, 2021
<u>AB</u>		<u>25MG</u>	<u>A215523 005</u>	Dec 08, 2021
<u>AB</u>		<u>30MG</u>	<u>A215523 006</u>	Dec 08, 2021
<u>AB</u>		<u>35MG</u>	<u>A215523 007</u>	Dec 08, 2021
<u>AB</u>		<u>40MG</u>	<u>A215523 008</u>	Dec 08, 2021
<u>AB</u>	GRANULES	<u>5MG</u>	<u>A213813 001</u>	Sep 09, 2020
<u>AB</u>		<u>10MG</u>	<u>A213813 002</u>	Sep 09, 2020
<u>AB</u>		<u>15MG</u>	<u>A213813 003</u>	Sep 09, 2020
<u>AB</u>		<u>20MG</u>	<u>A213813 004</u>	Sep 09, 2020
<u>AB</u>		<u>25MG</u>	<u>A213813 005</u>	Sep 09, 2020
<u>AB</u>		<u>30MG</u>	<u>A213813 006</u>	Sep 09, 2020
<u>AB</u>		<u>35MG</u>	<u>A213813 007</u>	Sep 09, 2020
<u>AB</u>		<u>40MG</u>	<u>A213813 008</u>	Sep 09, 2020
<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>5MG</u>	<u>A078992 001</u>	Nov 23, 2021
<u>AB</u>		<u>10MG</u>	<u>A078992 002</u>	Nov 23, 2021
<u>AB</u>		<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>		<u>20MG</u>	<u>A078992 005</u>	Nov 23, 2021
<u>AB</u>		<u>40MG</u>	<u>A078992 006</u>	Nov 23, 2021
<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>	SUN PHARM INDS INC	<u>5MG</u>	<u>A206734 001</u>	Nov 05, 2021
<u>AB</u>		<u>10MG</u>	<u>A206734 002</u>	Nov 05, 2021
<u>AB</u>		<u>15MG</u>	<u>A206734 003</u>	Nov 05, 2021
<u>AB</u>		<u>20MG</u>	<u>A206734 004</u>	Nov 05, 2021
<u>AB</u>		<u>25MG</u>	<u>A206734 005</u>	Nov 05, 2021
<u>AB</u>		<u>30MG</u>	<u>A206734 006</u>	Nov 05, 2021
<u>AB</u>		<u>35MG</u>	<u>A206734 007</u>	Nov 05, 2021
<u>AB</u>		<u>40MG</u>	<u>A206734 008</u>	Nov 05, 2021
<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

PRESCRIPTION DRUG PRODUCT LIST

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

FOCALIN XR

<u>AB</u>	<u>+!</u>	<u>40MG</u>	<u>N021802</u>	<u>006</u>	Aug 11, 2010
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TABLET;ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	ABHAI INC	<u>2.5MG</u>	<u>A206931</u>	<u>001</u>	Dec 04, 2015
<u>AB</u>		<u>5MG</u>	<u>A206931</u>	<u>002</u>	Dec 04, 2015
<u>AB</u>		<u>10MG</u>	<u>A206931</u>	<u>003</u>	Dec 04, 2015
<u>AB</u>	ALKEM LABS LTD	<u>2.5MG</u>	<u>A212631</u>	<u>001</u>	Jul 19, 2019
<u>AB</u>		<u>5MG</u>	<u>A212631</u>	<u>002</u>	Jul 19, 2019
<u>AB</u>		<u>10MG</u>	<u>A212631</u>	<u>003</u>	Jul 19, 2019
<u>AB</u>	CEDIPROF INC	<u>5MG</u>	<u>A209211</u>	<u>001</u>	Sep 19, 2018
<u>AB</u>		<u>10MG</u>	<u>A209211</u>	<u>002</u>	Sep 19, 2018
<u>AB</u>	NOVEL LABS INC	<u>2.5MG</u>	<u>A204534</u>	<u>001</u>	Dec 04, 2015
<u>AB</u>		<u>5MG</u>	<u>A204534</u>	<u>002</u>	Dec 04, 2015
<u>AB</u>		<u>10MG</u>	<u>A204534</u>	<u>003</u>	Dec 04, 2015
<u>AB</u>	RHODES PHARMS	<u>2.5MG</u>	<u>A208756</u>	<u>001</u>	Nov 20, 2017
<u>AB</u>		<u>5MG</u>	<u>A208756</u>	<u>002</u>	Nov 20, 2017
<u>AB</u>		<u>10MG</u>	<u>A208756</u>	<u>003</u>	Nov 20, 2017
<u>AB</u>	SUN PHARM INDUSTRIES	<u>2.5MG</u>	<u>A201231</u>	<u>001</u>	Sep 24, 2015
<u>AB</u>		<u>5MG</u>	<u>A201231</u>	<u>002</u>	Sep 24, 2015
<u>AB</u>		<u>10MG</u>	<u>A201231</u>	<u>003</u>	Sep 24, 2015
<u>AB</u>	TRIS PHARMA INC	<u>2.5MG</u>	<u>A207901</u>	<u>001</u>	Aug 26, 2016
<u>AB</u>		<u>5MG</u>	<u>A207901</u>	<u>002</u>	Aug 26, 2016
<u>AB</u>		<u>10MG</u>	<u>A207901</u>	<u>003</u>	Aug 26, 2016

FOCALIN

<u>AB</u>	<u>+</u>	NOVARTIS	<u>2.5MG</u>	<u>N021278</u>	<u>001</u>	Nov 13, 2001
<u>AB</u>	<u>+</u>		<u>5MG</u>	<u>N021278</u>	<u>002</u>	Nov 13, 2001
<u>AB</u>	<u>+!</u>		<u>10MG</u>	<u>N021278</u>	<u>003</u>	Nov 13, 2001

DEXMETHYLPHENIDATE HYDROCHLORIDE; SERDEXMETHYLPHENIDATE CHLORIDE

CAPSULE;ORAL

AZSTARYS

<u>+</u>	COMMAVE THERAP	EQ 5.2MG BASE;EQ 26.1MG BASE	N212994	001	May 07, 2021
<u>+</u>		EQ 7.8MG BASE;EQ 39.2MG BASE	N212994	002	May 07, 2021
<u>+!</u>		EQ 10.4MG BASE;EQ 52.3MG BASE	N212994	003	May 07, 2021

DEXRAZOXANE HYDROCHLORIDE

INJECTABLE;INJECTION

DEXRAZOXANE HYDROCHLORIDE

<u>AP</u>	<u>!</u>	EUGIA PHARMA	<u>EQ 250MG BASE/VIAL</u>	<u>A200752</u>	<u>001</u>	Oct 19, 2011
<u>AP</u>	<u>!</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A200752</u>	<u>002</u>	Oct 19, 2011
<u>AP</u>		GLAND	<u>EQ 250MG BASE/VIAL</u>	<u>A207321</u>	<u>002</u>	Dec 16, 2019
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A207321</u>	<u>001</u>	Nov 28, 2016
<u>AP</u>		HIKMA	<u>EQ 250MG BASE/VIAL</u>	<u>A076068</u>	<u>001</u>	Sep 28, 2004
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A076068</u>	<u>002</u>	Sep 28, 2004

DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL

DEXEDRINE

<u>AB</u>	<u>+</u>	IMPAX LABS INC	<u>5MG</u>	<u>N017078</u>	<u>001</u>
<u>AB</u>	<u>+</u>		<u>10MG</u>	<u>N017078</u>	<u>002</u>
<u>AB</u>	<u>+!</u>		<u>15MG</u>	<u>N017078</u>	<u>003</u>

DEXTROAMPHETAMINE SULFATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>5MG</u>	<u>A203901</u>	<u>001</u>	Nov 30, 2012
<u>AB</u>		<u>10MG</u>	<u>A203901</u>	<u>002</u>	Nov 30, 2012
<u>AB</u>		<u>15MG</u>	<u>A203901</u>	<u>003</u>	Nov 30, 2012
<u>AB</u>	MAYNE PHARMA	<u>5MG</u>	<u>A076137</u>	<u>001</u>	Jan 18, 2002
<u>AB</u>		<u>10MG</u>	<u>A076137</u>	<u>002</u>	Jan 18, 2002
<u>AB</u>		<u>15MG</u>	<u>A076137</u>	<u>003</u>	Jan 18, 2002
<u>AB</u>	NESHER PHARMS	<u>5MG</u>	<u>A209111</u>	<u>001</u>	Jun 27, 2017
<u>AB</u>		<u>10MG</u>	<u>A209111</u>	<u>002</u>	Jun 27, 2017
<u>AB</u>		<u>15MG</u>	<u>A209111</u>	<u>003</u>	Jun 27, 2017
<u>AB</u>	SPECGX LLC	<u>5MG</u>	<u>A076353</u>	<u>001</u>	May 06, 2003
<u>AB</u>		<u>10MG</u>	<u>A076353</u>	<u>002</u>	May 06, 2003
<u>AB</u>		<u>15MG</u>	<u>A076353</u>	<u>003</u>	May 06, 2003
<u>AB</u>	STRIDES PHARMA	<u>5MG</u>	<u>A205673</u>	<u>001</u>	Oct 31, 2017
<u>AB</u>		<u>10MG</u>	<u>A205673</u>	<u>002</u>	Oct 31, 2017
<u>AB</u>		<u>15MG</u>	<u>A205673</u>	<u>003</u>	Oct 31, 2017

SOLUTION;ORAL

DEXTROAMPHETAMINE SULFATE

<u>AA</u>	<u>!</u>	OUTLOOK PHARMS	<u>5MG/5ML</u>	<u>A040776</u>	<u>001</u>	Jan 29, 2008
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PRESCRIPTION DRUG PRODUCT LIST

DEXTROAMPHETAMINE SULFATE

SOLUTION;ORAL

DEXTROAMPHETAMINE SULFATE

AA	TRIS PHARMA INC	5MG/5ML	A203644	001	May 29, 2013
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TABLET;ORAL

DEXTROAMPHETAMINE SULFATE

AA	ARBOR PHARMS LLC	2.5MG	A090533	001	Oct 25, 2011
AA		5MG	A090533	002	Oct 25, 2011
AA		7.5MG	A090533	003	Oct 25, 2011
AA		10MG	A090533	004	Oct 25, 2011
AA		15MG	A090533	005	Oct 25, 2011
AA		20MG	A090533	006	Oct 25, 2011
AA		30MG	A090533	007	Oct 25, 2011
AA	AUROLIFE PHARMA LLC	5MG	A202893	001	Jul 31, 2013
AA		10MG	A202893	002	Jul 31, 2013
AA	AVANTHI INC	5MG	A203548	001	Nov 23, 2015
AA		10MG	A203548	002	Nov 23, 2015
AA	BARR	5MG	A040361	001	Jan 31, 2001
AA	!	10MG	A040361	002	Jan 31, 2001
AA	NESHER PHARMS	5MG	A206588	001	Mar 28, 2016
AA		10MG	A206588	002	Mar 28, 2016
AA	NOVEL LABS INC	5MG	A204330	001	Mar 16, 2016
AA		10MG	A204330	002	Mar 16, 2016
AA	NUVO PHARM	5MG	A210059	001	Oct 18, 2017
AA		10MG	A210059	002	Oct 18, 2017
AA	SPECGX LLC	5MG	A040436	001	Jan 29, 2002
AA		10MG	A040436	002	Jan 29, 2002
AA	WINDER LABS LLC	2.5MG	A212160	001	Jun 07, 2021
AA		5MG	A212160	002	Jun 07, 2021
AA		7.5MG	A212160	003	Jun 07, 2021
AA		10MG	A212160	004	Jun 07, 2021
AA		15MG	A212160	005	Jun 07, 2021
AA		20MG	A212160	006	Jun 07, 2021
AA		30MG	A212160	007	Jun 07, 2021

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PROMETHAZINE DM

AA	!	SLATE	15MG/5ML; 6.25MG/5ML	A040649	001	Feb 14, 2006
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PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

AA		AMNEAL PHARMS	15MG/5ML; 6.25MG/5ML	A090575	001	Feb 08, 2011
AA		TRIS PHARMA INC	15MG/5ML; 6.25MG/5ML	A091687	001	Jun 28, 2012

PROMETHAZINE W/ DEXTROMETHORPHAN

AA		WOCKHARDT BIO AG	15MG/5ML; 6.25MG/5ML	A088864	001	Jan 04, 1985
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DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE

CAPSULE;ORAL

DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE

AB		ACTAVIS ELIZABETH	20MG;10MG	A202934	001	Oct 10, 2017
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NUDEXTA

AB	+	AVANIR PHARMS	20MG;10MG	N021879	001	Oct 29, 2010
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DEXTROSE

INJECTABLE;INJECTION

DEXTROSE 10% IN PLASTIC CONTAINER

AP	+	B BRAUN	10GM/100ML	N019626	004	Feb 02, 1988
AP	+	BAXTER HLTHCARE	10GM/100ML	N016694	001	
AP		FRESENIUS KABI USA	10GM/100ML	A209448	001	Jul 16, 2018
AP	+	ICU MEDICAL INC	10GM/100ML	N018080	001	

DEXTROSE 5% IN PLASTIC CONTAINER

AP	+	B BRAUN	50MG/ML	N016730	002	
AP	+		5GM/100ML	N016730	001	
AP	+		5GM/100ML	N019626	002	Feb 02, 1988
AP	+	BAXTER HLTHCARE	50MG/ML	N016673	003	Oct 30, 1985
AP	+		50MG/ML	N020179	002	Dec 07, 1992
AP	+		5GM/100ML	N016673	001	
AP	+		5GM/100ML	N020179	001	Dec 07, 1992
AP		FRESENIUS KABI USA	50MG/ML	A207449	001	Oct 21, 2016
AP	+	HOSPIRA	5GM/100ML	N019466	001	Jul 15, 1985
AP	+		5GM/100ML	N019479	001	Sep 17, 1985
AP	+	ICU MEDICAL INC	50MG/ML	N016367	002	

DEXTROSE 50%

AP	+	HOSPIRA	500MG/ML	N019445	001	Jun 03, 1986
AP		INTL MEDICATION SYS	500MG/ML	A203451	001	Mar 26, 2021

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 50% IN PLASTIC CONTAINER

AP	+	BAXTER HLTHCARE	50GM/100ML	N020047 001	Jul 02, 1991
AP	+	ICU MEDICAL INC	50GM/100ML	N018563 001	Mar 23, 1982

DEXTROSE 70% IN PLASTIC CONTAINER

AP	+	B BRAUN	70GM/100ML	N019626 005	Feb 18, 2015
AP	+	BAXTER HLTHCARE	70GM/100ML	N020047 003	Jul 02, 1991
AP	+	ICU MEDICAL INC	70GM/100ML	N018561 001	Mar 23, 1982
AP	+		70GM/100ML	N019893 001	Dec 26, 1989

DEXTROSE 20% IN PLASTIC CONTAINER

+	ICU MEDICAL INC	20GM/100ML	N018564 001	Mar 23, 1982
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DEXTROSE 25%

+	HOSPIRA	250MG/ML	N019445 002	Nov 23, 1998
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DEXTROSE 30% IN PLASTIC CONTAINER

+	ICU MEDICAL INC	30GM/100ML	N019345 001	Jan 26, 1985
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DEXTROSE 40% IN PLASTIC CONTAINER

+	ICU MEDICAL INC	40GM/100ML	N018562 001	Mar 23, 1982
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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
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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
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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
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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
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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
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DEXTROSE; POTASSIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER

AP	+	BAXTER HLTHCARE	5GM/100ML; 75MG/100ML	N017634 004
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DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER

AP	+	BAXTER HLTHCARE	5GM/100ML; 150MG/100ML	N017634 001
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DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

AP	+	BAXTER HLTHCARE	5GM/100ML; 300MG/100ML	N017634 002
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POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER

AP		B BRAUN	5GM/100ML; 150MG/100ML	N019699 004	Sep 29, 1989
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POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER

AP		B BRAUN	5GM/100ML; 300MG/100ML	N019699 006	Sep 29, 1989
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DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER

+	BAXTER HLTHCARE	5GM/100ML; 224MG/100ML	N017634 003
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POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER

ICU MEDICAL INC	5GM/100ML; 149MG/100ML	N018371 001
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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
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AP		BAXTER HLTHCARE	5GM/100ML; 150MG/100ML; 200MG/100ML	N018037 007	Apr 13, 1982
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DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K)

AP		BAXTER HLTHCARE	5GM/100ML; 224MG/100ML; 200MG/100ML	N018037 004
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PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDEINJECTABLE; INJECTION

	<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ</u>			
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;200MG/100ML	N018037 008	Apr 13, 1982
	<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K)</u>			
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;200MG/100ML	N018037 001	
	<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ</u>			
AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;200MG/100ML	N018037 005	Apr 13, 1982
	<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ</u>			
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;200MG/100ML	N018037 009	Apr 13, 1982
	<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ</u>			
AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;200MG/100ML	N018037 002	
	<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K)</u>			
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;200MG/100ML	N018037 003	
	<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER</u>			
AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;330MG/100ML	N018629 005	Mar 23, 1982
AP		5GM/100ML;150MG/100ML;330MG/100ML	N018629 002	Mar 23, 1982
	<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER</u>			
AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;330MG/100ML	N018629 003	Mar 23, 1982
	<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER</u>			
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;330MG/100ML	N018629 004	Mar 23, 1982
AP		5GM/100ML;300MG/100ML;330MG/100ML	N018629 006	Mar 23, 1982
	<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER</u>			
AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;330MG/100ML	N018629 007	Mar 23, 1982
	<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER</u>			
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;330MG/100ML	N018629 008	Mar 23, 1982
	<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER</u>			
AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;330MG/100ML	N018629 001	Mar 23, 1982
	<u>DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER</u>			
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;450MG/100ML	N018008 010	
	<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML;75MG/100ML;200MG/100ML	N019630 008	Feb 17, 1988
	<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML;75MG/100ML;330MG/100ML	N019630 014	Feb 17, 1988
	<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML;75MG/100ML;450MG/100ML	N019630 020	Feb 17, 1988
	<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML;75MG/100ML;900MG/100ML	N019630 026	Feb 17, 1988
	<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML;150MG/100ML;200MG/100ML	N019630 010	Feb 17, 1988
	<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML;150MG/100ML;330MG/100ML	N019630 016	Feb 17, 1988
	<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML;150MG/100ML;450MG/100ML	N019630 022	Feb 17, 1988
	<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML;150MG/100ML;900MG/100ML	N019630 028	Feb 17, 1988
	<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML;300MG/100ML;200MG/100ML	N019630 012	Feb 17, 1988
	<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML;300MG/100ML;330MG/100ML	N019630 018	Feb 17, 1988
	<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML;300MG/100ML;450MG/100ML	N019630 024	Feb 17, 1988
	<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML;300MG/100ML;900MG/100ML	N019630 030	Feb 17, 1988
	<u>POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;450MG/100ML	N018008 005	Apr 28, 1982
AP		5GM/100ML;150MG/100ML;450MG/100ML	N018008 006	Apr 28, 1982
AP	+! ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;450MG/100ML	N018362 009	Jul 05, 1983
AP	+!	5GM/100ML;149MG/100ML;450MG/100ML	N018362 005	Mar 28, 1988
	<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;450MG/100ML	N018008 007	Apr 28, 1982
AP	+! ICU MEDICAL INC	5GM/100ML;149MG/100ML;450MG/100ML	N018362 010	Jul 05, 1983
	<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;900MG/100ML	N019308 005	Apr 05, 1985
AP	+! ICU MEDICAL INC	5GM/100ML;149MG/100ML;900MG/100ML	N019691 005	Mar 24, 1988
	<u>POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;450MG/100ML	N018008 008	Apr 28, 1982
AP	+! ICU MEDICAL INC	5GM/100ML;224MG/100ML;450MG/100ML	N018362 002	
	<u>POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;450MG/100ML	N018008 009	Apr 28, 1982
AP	+! ICU MEDICAL INC	5GM/100ML;298MG/100ML;450MG/100ML	N018362 003	

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

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**

MD-GASTROVIEW

AA LIEBEL-FLARSHEIM **66%;10%** **A087388 001**

DIAZEPAM

CONCENTRATE; ORAL

DIAZEPAM INTENSOL
! HIKMA 5MG/ML A071415 001 Apr 03, 1987

GEL; RECTAL

DIASTAT
+ BAUSCH 2.5MG/0.5ML (5MG/ML) N020648 001 Jul 29, 1997
DIASTAT ACUDIAL
+ BAUSCH 10MG/2ML (5MG/ML) N020648 007 Sep 15, 2005
+! 20MG/4ML (5MG/ML) N020648 006 Sep 15, 2005

INJECTABLE; INJECTION

DIAZEPAM

AP ! BELOTECA INC **10MG/2ML (5MG/ML)** **A210363 001** Mar 18, 2019
AP **50MG/10ML (5MG/ML)** **A211998 001** Dec 26, 2019
AP HIKMA **10MG/2ML (5MG/ML)** **A070313 001** Dec 16, 1985
AP HOSPIRA **10MG/2ML (5MG/ML)** **A072079 001** Dec 20, 1988
AP ! **50MG/10ML (5MG/ML)** **A071583 001** Oct 13, 1987

SOLUTION; ORAL

DIAZEPAM

AA ! HIKMA **5MG/5ML** **A070928 001** Apr 03, 1987
AA LANNETT CO INC **5MG/5ML** **A206477 001** Jun 24, 2016

SPRAY; NASAL

VALTOCO
+ NEURELIS INC 5MG/SPRAY N211635 001 Jan 10, 2020
+ 7.5MG/SPRAY N211635 002 Jan 10, 2020
+! 10MG/SPRAY N211635 003 Jan 10, 2020

TABLET; ORAL

DIAZEPAM

AB IVAX SUB TEVA **2MG** **A071307 001** Dec 10, 1986
PHARMS
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
AB **5MG** **A070325 003** Sep 04, 1985
AB **10MG** **A070325 001** Sep 04, 1985
AB STRIDES PHARMA **2MG** **A077749 001** Mar 31, 2006

PRESCRIPTION DRUG PRODUCT LIST

DIAZEPAM

TABLET; ORAL

DIAZEPAM

<u>AB</u>		<u>5MG</u>	<u>A077749</u>	<u>002</u>	Mar 31, 2006
<u>AB</u>		<u>10MG</u>	<u>A077749</u>	<u>003</u>	Mar 31, 2006

VALIUM

<u>AB</u>	+	ROCHE	<u>2MG</u>	<u>N013263</u>	<u>002</u>
<u>AB</u>	+		<u>5MG</u>	<u>N013263</u>	<u>004</u>
<u>AB</u>	+	!	<u>10MG</u>	<u>N013263</u>	<u>006</u>

DIAZOXIDE

SUSPENSION; ORAL

DIAZOXIDE

<u>AB</u>		E5 PHARMA INC	<u>50MG/ML</u>	<u>A211050</u>	<u>001</u>	Dec 20, 2019
<u>AB</u>		NOVITIUM PHARMA	<u>50MG/ML</u>	<u>A210799</u>	<u>001</u>	Jul 08, 2020

PROGLYCEM

<u>AB</u>	+	!	TEVA BRANDED PHARM	<u>50MG/ML</u>	<u>N017453</u>	<u>001</u>
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DICHLORPHENAMIDE

TABLET; ORAL

KEVEYIS

+	!	XERIS	50MG	N011366	002	Aug 07, 2015
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DICLOFENAC

CAPSULE; ORAL

ZORVOLEX

+	!	ZYLA	35MG	N204592	002	Oct 18, 2013
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DICLOFENAC EPOLAMINE

SYSTEM; TOPICAL

FLECTOR

+	!	INST BIOCHEM	1.3%	N021234	001	Jan 31, 2007
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LICART

+	!	IBSA INST BIO	1.3%	N206976	001	Dec 19, 2018
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DICLOFENAC POTASSIUM

CAPSULE; ORAL

DICLOFENAC POTASSIUM

<u>AB</u>		AUROBINDO PHARMA LTD	<u>25MG</u>	<u>A213875</u>	<u>001</u>	Oct 19, 2021
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<u>AB</u>		BIONPHARMA INC	<u>25MG</u>	<u>A204648</u>	<u>001</u>	Feb 23, 2016
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ZIPSOR

<u>AB</u>	+	!	ASSERTIO	<u>25MG</u>	<u>N022202</u>	<u>001</u>	Jun 16, 2009
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FOR SOLUTION; ORAL

CAMBIA

<u>AB</u>	+	!	ASSERTIO	<u>50MG</u>	<u>N022165</u>	<u>001</u>	Jun 17, 2009
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DICLOFENAC POTASSIUM

<u>AB</u>		PAR FORM	<u>50MG</u>	<u>A202964</u>	<u>001</u>	May 02, 2016
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TABLET; ORAL

CATAFLAM

<u>AB</u>		AMICI	<u>25MG</u>	<u>A076561</u>	<u>002</u>	Jul 21, 2021
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<u>AB</u>			<u>50MG</u>	<u>A076561</u>	<u>001</u>	Mar 18, 2004
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DICLOFENAC POTASSIUM

<u>AB</u>		RICONPHARMA LLC	<u>50MG</u>	<u>A215585</u>	<u>001</u>	Oct 08, 2021
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<u>AB</u>	!	RK PHARMA	<u>50MG</u>	<u>A075463</u>	<u>001</u>	Jul 26, 1999
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<u>AB</u>		RUBICON	<u>25MG</u>	<u>A075229</u>	<u>002</u>	Sep 16, 2021
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<u>AB</u>			<u>50MG</u>	<u>A075229</u>	<u>001</u>	Nov 20, 1998
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<u>AB</u>		TEVA	<u>50MG</u>	<u>A075219</u>	<u>001</u>	Aug 06, 1998
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DICLOFENAC SODIUM

GEL; TOPICAL

DICLOFENAC SODIUM

<u>AB</u>		ACTAVIS MID ATLANTIC	<u>3%</u>	<u>A206493</u>	<u>001</u>	Dec 02, 2015
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<u>AB</u>		AKORN	<u>1%</u>	<u>A209484</u>	<u>001</u>	Nov 21, 2018
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<u>AB</u>		CIPLA	<u>1%</u>	<u>A209903</u>	<u>001</u>	Aug 03, 2018
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<u>AB</u>		GLENMARK PHARMS LTD	<u>3%</u>	<u>A208301</u>	<u>001</u>	Sep 13, 2016
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<u>AB</u>		PADAGIS ISRAEL	<u>3%</u>	<u>A210893</u>	<u>001</u>	Jul 27, 2018
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<u>AB</u>	!	TARO	<u>3%</u>	<u>A206298</u>	<u>001</u>	Apr 28, 2016
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<u>AB</u>		TOLMAR	<u>3%</u>	<u>A200936</u>	<u>001</u>	Oct 28, 2013
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SOLUTION; TOPICAL

DICLOFENAC SODIUM

<u>AT</u>		AKORN	<u>1.5%</u>	<u>A206655</u>	<u>001</u>	Jan 28, 2021
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<u>AT</u>		AMNEAL PHARMS	<u>1.5%</u>	<u>A206116</u>	<u>001</u>	Sep 02, 2016
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<u>AT</u>		CADILA	<u>1.5%</u>	<u>A206411</u>	<u>001</u>	Apr 17, 2018
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<u>AT</u>		LUPIN LTD	<u>1.5%</u>	<u>A204132</u>	<u>001</u>	Aug 20, 2015
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PRESCRIPTION DRUG PRODUCT LIST

DICLOFENAC SODIUM

SOLUTION;TOPICAL

DICLOFENAC SODIUM

AT	NOVEL LABS INC	<u>1.5%</u>	<u>A205878</u>	<u>001</u>	Dec 09, 2015
AT	! TARO	<u>1.5%</u>	<u>A203818</u>	<u>001</u>	Nov 26, 2014
AT	TELIGENT	<u>1.5%</u>	<u>A202769</u>	<u>001</u>	Jul 08, 2015
AT	TWI PHARMS	<u>1.5%</u>	<u>A202393</u>	<u>001</u>	Nov 24, 2014
AT	WATSON LABS INC	<u>1.5%</u>	<u>A202852</u>	<u>001</u>	Nov 24, 2014

PENNSAID

+! HORIZON

2%

N204623 001 Jan 16, 2014

SOLUTION/DROPS;OPHTHALMIC

DICLOFENAC SODIUM

AT	AKORN	<u>0.1%</u>	<u>A077845</u>	<u>001</u>	Apr 17, 2008
AT	ALTAIRE PHARMS INC	<u>0.1%</u>	<u>A203383</u>	<u>001</u>	Nov 16, 2015
AT	! BAUSCH AND LOMB	<u>0.1%</u>	<u>A078792</u>	<u>001</u>	Dec 28, 2007
AT	RISING	<u>0.1%</u>	<u>A078553</u>	<u>001</u>	Dec 28, 2007
AT	SANDOZ INC	<u>0.1%</u>	<u>A078031</u>	<u>001</u>	Feb 06, 2008

TABLET, DELAYED RELEASE;ORAL

DICLOFENAC SODIUM

AB	ACTAVIS ELIZABETH	<u>50MG</u>	<u>A074514</u>	<u>001</u>	Mar 26, 1996
AB		<u>75MG</u>	<u>A074514</u>	<u>002</u>	Mar 26, 1996
AB	CARLSBAD	<u>25MG</u>	<u>A075185</u>	<u>002</u>	Nov 13, 1998
AB		<u>50MG</u>	<u>A075185</u>	<u>003</u>	Nov 13, 1998
AB		<u>75MG</u>	<u>A075185</u>	<u>001</u>	Nov 13, 1998
AB	! UNIQUE PHARM LABS	<u>25MG</u>	<u>A090066</u>	<u>001</u>	Dec 01, 2010
AB	!	<u>50MG</u>	<u>A090066</u>	<u>002</u>	Dec 01, 2010
AB	!	<u>75MG</u>	<u>A077863</u>	<u>003</u>	Jun 08, 2007

TABLET, EXTENDED RELEASE;ORAL

DICLOFENAC SODIUM

AB	! DEXCEL LTD	<u>100MG</u>	<u>A076201</u>	<u>001</u>	Nov 06, 2002
AB	VPNA	<u>100MG</u>	<u>A075492</u>	<u>001</u>	Feb 11, 2000

DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE;ORAL

ARTHROTEC

AB	+ PFIZER	<u>50MG;0.2MG</u>	<u>N020607</u>	<u>001</u>	Dec 24, 1997
AB	+!	<u>75MG;0.2MG</u>	<u>N020607</u>	<u>002</u>	Dec 24, 1997

DICLOFENAC SODIUM AND MISOPROSTOL

AB	ACTAVIS LABS FL INC	<u>50MG;0.2MG</u>	<u>A201089</u>	<u>001</u>	Jul 09, 2012
AB		<u>75MG;0.2MG</u>	<u>A201089</u>	<u>002</u>	Jul 09, 2012
AB	AMNEAL PHARMS	<u>50MG;0.2MG</u>	<u>A203995</u>	<u>001</u>	Nov 25, 2016
AB		<u>75MG;0.2MG</u>	<u>A203995</u>	<u>002</u>	Nov 25, 2016
AB	MICRO LABS	<u>50MG;0.2MG</u>	<u>A204355</u>	<u>001</u>	Jul 15, 2021
AB		<u>75MG;0.2MG</u>	<u>A204355</u>	<u>002</u>	Jul 15, 2021
AB	SANDOZ	<u>50MG;0.2MG</u>	<u>A200158</u>	<u>001</u>	May 09, 2013
AB		<u>75MG;0.2MG</u>	<u>A200158</u>	<u>002</u>	May 09, 2013
AB	YUNG SHIN PHARM	<u>50MG;0.2MG</u>	<u>A205143</u>	<u>001</u>	Feb 19, 2020
AB		<u>75MG;0.2MG</u>	<u>A205143</u>	<u>002</u>	Feb 19, 2020

DICLOXACILLIN SODIUM

CAPSULE;ORAL

DICLOXACILLIN SODIUM

TEVA

EQ 250MG BASE

A062286 001 Jun 03, 1982

!

EQ 500MG BASE

A062286 002 Jun 03, 1982

DICYCLOMINE HYDROCHLORIDE

CAPSULE;ORAL

DICYCLOMINE HYDROCHLORIDE

AB	! LANNETT	<u>10MG</u>	<u>A084285</u>	<u>001</u>	
AB	MYLAN	<u>10MG</u>	<u>A040319</u>	<u>001</u>	Sep 07, 1999
AB	WATSON LABS	<u>10MG</u>	<u>A085082</u>	<u>001</u>	Jun 19, 1986
AB	WEST WARD	<u>10MG</u>	<u>A040204</u>	<u>001</u>	Feb 28, 1997

INJECTABLE; INJECTION

BENTYL

AP	+! ALLERGAN	<u>10MG/ML</u>	<u>N008370</u>	<u>001</u>	Oct 15, 1984
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BENTYL PRESERVATIVE FREE

AP	+! ALLERGAN	<u>10MG/ML</u>	<u>N008370</u>	<u>002</u>	Oct 15, 1984
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DICYCLOMINE HYDROCHLORIDE

AP	AKORN	<u>10MG/ML</u>	<u>A207084</u>	<u>001</u>	May 04, 2018
AP	AM REGENT	<u>10MG/ML</u>	<u>A208353</u>	<u>001</u>	Feb 17, 2017
AP	CUSTOPHARM INC	<u>10MG/ML</u>	<u>A210788</u>	<u>001</u>	Feb 11, 2019
AP	FOSUN PHARMA	<u>10MG/ML</u>	<u>A210979</u>	<u>001</u>	Jul 02, 2018
AP	FRESENIUS KABI USA	<u>10MG/ML</u>	<u>A210257</u>	<u>001</u>	Jan 25, 2019
AP	GENE YORK PHARMS	<u>10MG/ML</u>	<u>A212058</u>	<u>001</u>	Apr 26, 2019

PRESCRIPTION DRUG PRODUCT LIST

DICYCLOMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DICYCLOMINE HYDROCHLORIDE

AP	NEXUS PHARMS	<u>10MG/ML</u>	<u>A206468</u>	<u>001</u>	Feb 01, 2019
AP	SLATE	<u>10MG/ML</u>	<u>A207076</u>	<u>001</u>	Nov 02, 2018

DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE)

AP	HIKMA	<u>10MG/ML</u>	<u>A040465</u>	<u>001</u>	Jun 30, 2003
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SYRUP; ORAL

DICYCLOMINE HYDROCHLORIDE

AA	! GENERICS	<u>10MG/5ML</u>	<u>A040169</u>	<u>001</u>	Mar 24, 2005
AA	HIKMA	<u>10MG/5ML</u>	<u>A212286</u>	<u>001</u>	May 22, 2020
AA	NOVITIUM PHARMA	<u>10MG/5ML</u>	<u>A214721</u>	<u>001</u>	Apr 23, 2021

TABLET; ORAL

DICYCLOMINE HYDROCHLORIDE

AB	HIKMA PHARMS	<u>20MG</u>	<u>A040161</u>	<u>001</u>	Oct 01, 1996
AB	LANNETT	<u>20MG</u>	<u>A040230</u>	<u>001</u>	Feb 26, 1999
AB	! WATSON LABS	<u>20MG</u>	<u>A085223</u>	<u>001</u>	Jul 30, 1986

DIENOGEST; ESTRADIOL VALERATE

TABLET; ORAL

NATAZIA

+	!	BAYER HLTHCARE	N/A, 2MG, 3MG, N/A, N/A; 3MG, 2MG, 2MG, 1MG, N/A	<u>N022252</u>	<u>001</u>	May 06, 2010
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DIETHYLPROPION HYDROCHLORIDE

TABLET; ORAL

DIETHYLPROPION HYDROCHLORIDE

AA	AVANTHI INC	<u>25MG</u>	<u>A201212</u>	<u>001</u>	Dec 22, 2010
AA	! LANNETT CO INC	<u>25MG</u>	<u>A200177</u>	<u>001</u>	Jul 18, 2011

TABLET, EXTENDED RELEASE; ORAL

DIETHYLPROPION HYDROCHLORIDE

AB	! LANNETT CO INC	<u>75MG</u>	<u>A091680</u>	<u>001</u>	Oct 24, 2011
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DIFELIKEFALIN ACETATE

SOLUTION; INTRAVENOUS

KORSUVA

+	!	CARA THERAP	EQ 0.065MG BASE/1.3ML (EQ 0.05MG BASE/ML)	<u>N214916</u>	<u>001</u>	Aug 23, 2021
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DIFLORASONE DIACETATE

CREAM; TOPICAL

DIFLORASONE DIACETATE

BX	!	ANI PHARMS	0.05%	<u>A076263</u>	<u>001</u>	Dec 20, 2002
BX	!	TARO	0.05%	<u>A075508</u>	<u>001</u>	Apr 24, 2000

OINTMENT; TOPICAL

DIFLORASONE DIACETATE

AB	FOUGERA PHARMS	<u>0.05%</u>	<u>A075374</u>	<u>001</u>	Apr 27, 1999
AB	RISING	<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	TELGENT	<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	<u>500MG</u>	<u>A203547</u>	<u>001</u>	Jun 16, 2017

DIFLUPREDNATE

EMULSION; OPHTHALMIC

DIFLUPREDNATE

AB	AMNEAL	<u>0.05%</u>	<u>A211526</u>	<u>001</u>	Nov 17, 2021
AB	CIPLA	<u>0.05%</u>	<u>A211776</u>	<u>001</u>	Aug 09, 2021

DUREZOL

AB	! NOVARTIS	<u>0.05%</u>	<u>N022212</u>	<u>001</u>	Jun 23, 2008
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DIGOXIN

ELIXIR; ORAL

DIGOXIN

AA	! HIKMA	<u>0.05MG/ML</u>	<u>N021648</u>	<u>001</u>	Aug 26, 2004
AA	NOVITIUM PHARMA	<u>0.05MG/ML</u>	<u>A213000</u>	<u>001</u>	Oct 04, 2019

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>	

PRESCRIPTION DRUG PRODUCT LIST

DIGOXIN

INJECTABLE; INJECTION

LANOXIN

<u>AP</u>	<u>+</u> !	COVIS	<u>0.25MG/ML</u>	<u>N009330</u>	<u>002</u>	
		LANOXIN PEDIATRIC				
	<u>+</u> !	COVIS	0.1MG/ML	N009330	004	

TABLET; ORAL

DIGOXIN

<u>AB</u>		HIKMA INTL PHARMS	<u>0.125MG</u>	<u>A077002</u>	<u>002</u>	Oct 30, 2007
<u>AB</u>			<u>0.25MG</u>	<u>A077002</u>	<u>001</u>	Oct 30, 2007
<u>AB</u>		IMPAX LABS	<u>0.125MG</u>	<u>A078556</u>	<u>001</u>	Jul 20, 2009
<u>AB</u>			<u>0.25MG</u>	<u>A078556</u>	<u>002</u>	Jul 20, 2009
<u>AB</u>		NOVITIUM PHARMA	<u>0.125MG</u>	<u>A215307</u>	<u>001</u>	Nov 22, 2021
<u>AB</u>			<u>0.25MG</u>	<u>A215307</u>	<u>002</u>	Nov 22, 2021
<u>AB</u>		RISING PHARMA	<u>0.125MG</u>	<u>A040282</u>	<u>001</u>	Dec 23, 1999
<u>AB</u>			<u>0.25MG</u>	<u>A040282</u>	<u>002</u>	Dec 23, 1999
<u>AB</u>		STEVENS J	<u>0.125MG</u>	<u>A076268</u>	<u>001</u>	Jul 26, 2002
<u>AB</u>			<u>0.25MG</u>	<u>A076268</u>	<u>002</u>	Jul 26, 2002
<u>AB</u>		SUN PHARM INDS INC	<u>0.125MG</u>	<u>A076363</u>	<u>001</u>	Jan 31, 2003
<u>AB</u>			<u>0.25MG</u>	<u>A076363</u>	<u>002</u>	Jan 31, 2003

LANOXIN

<u>AB</u>	<u>+</u>	CONCORDIA	<u>0.125MG</u>	<u>N020405</u>	<u>002</u>	Sep 30, 1997
<u>AB</u>	<u>+</u> !		<u>0.25MG</u>	<u>N020405</u>	<u>004</u>	Sep 30, 1997
	<u>+</u>		0.0625MG	N020405	001	Sep 30, 1997

DIHYDROERGOTAMINE MESYLATE

INJECTABLE; INJECTION

D.H.E. 45

<u>AP</u>	<u>+</u> !	BAUSCH	<u>1MG/ML</u>	<u>N005929</u>	<u>001</u>	
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DIHYDROERGOTAMINE MESYLATE

<u>AP</u>		HIKMA	<u>1MG/ML</u>	<u>A040453</u>	<u>001</u>	Jun 09, 2003
<u>AP</u>		HIKMA PHARMS	<u>1MG/ML</u>	<u>A206621</u>	<u>001</u>	Sep 15, 2017
<u>AP</u>		PADAGIS US	<u>1MG/ML</u>	<u>A040475</u>	<u>001</u>	Apr 28, 2003
<u>AP</u>		PROVEPHARM SAS	<u>1MG/ML</u>	<u>A212046</u>	<u>001</u>	Jan 07, 2020
<u>AP</u>		SAGENT PHARMS INC	<u>1MG/ML</u>	<u>A207264</u>	<u>001</u>	Jul 11, 2018

SPRAY, METERED; NASAL

DIHYDROERGOTAMINE MESYLATE

<u>AB</u>		AMNEAL	<u>0.5MG/SPRAY</u>	<u>A214105</u>	<u>001</u>	Jan 04, 2022
<u>AB</u>		CIPLA	<u>0.5MG/SPRAY</u>	<u>A212907</u>	<u>001</u>	May 20, 2020
<u>AB</u>		CUSTOPHARM INC	<u>0.5MG/SPRAY</u>	<u>A211393</u>	<u>001</u>	Feb 28, 2020

MIGRANAL

<u>AB</u>	<u>+</u> !	BAUSCH	<u>0.5MG/SPRAY</u>	<u>N020148</u>	<u>001</u>	Dec 08, 1997
		TRUDHESA				
	<u>+</u> !	IMPEL NEUROPHARMA	0.725MG/SPRAY	N213436	001	Sep 02, 2021

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

DILTIAZEM HYDROCHLORIDE

<u>AB1</u>		GLENMARK PHARMS LTD	<u>60MG</u>	<u>A212317</u>	<u>001</u>	Mar 22, 2021
<u>AB1</u>			<u>90MG</u>	<u>A212317</u>	<u>002</u>	Mar 22, 2021
<u>AB1</u>			<u>120MG</u>	<u>A212317</u>	<u>003</u>	Mar 22, 2021
<u>AB1</u>		MYLAN	<u>60MG</u>	<u>A074910</u>	<u>001</u>	May 02, 1997
<u>AB1</u>			<u>90MG</u>	<u>A074910</u>	<u>002</u>	May 02, 1997
<u>AB1</u>	<u>!</u>		<u>120MG</u>	<u>A074910</u>	<u>003</u>	May 02, 1997
<u>AB2</u>		ACCORD HLTHCARE	<u>120MG</u>	<u>A206997</u>	<u>001</u>	Apr 28, 2020
<u>AB2</u>			<u>180MG</u>	<u>A206997</u>	<u>002</u>	Apr 28, 2020
<u>AB2</u>			<u>240MG</u>	<u>A206997</u>	<u>003</u>	Apr 28, 2020
<u>AB2</u>		APOTEX	<u>120MG</u>	<u>A074943</u>	<u>003</u>	Dec 19, 2000
<u>AB2</u>			<u>180MG</u>	<u>A074943</u>	<u>002</u>	Dec 19, 2000
<u>AB2</u>	<u>!</u>		<u>240MG</u>	<u>A074943</u>	<u>001</u>	Aug 06, 1998

CARDIZEM CD

<u>AB3</u>	<u>+</u>	BAUSCH	<u>120MG</u>	<u>N020062</u>	<u>001</u>	Aug 10, 1992
<u>AB3</u>	<u>+</u>		<u>180MG</u>	<u>N020062</u>	<u>002</u>	Dec 27, 1991
<u>AB3</u>	<u>+</u>		<u>240MG</u>	<u>N020062</u>	<u>003</u>	Dec 27, 1991
<u>AB3</u>	<u>+</u>		<u>300MG</u>	<u>N020062</u>	<u>004</u>	Dec 27, 1991
<u>AB3</u>	<u>+</u> !		<u>360MG</u>	<u>N020062</u>	<u>005</u>	Aug 24, 1999

CARTIA XT

<u>AB3</u>		ACTAVIS LABS FL INC	<u>120MG</u>	<u>A074752</u>	<u>002</u>	Jul 09, 1998
<u>AB3</u>			<u>180MG</u>	<u>A074752</u>	<u>001</u>	Jul 09, 1998
<u>AB3</u>			<u>240MG</u>	<u>A074752</u>	<u>003</u>	Jul 09, 1998
<u>AB3</u>			<u>300MG</u>	<u>A074752</u>	<u>004</u>	Jul 09, 1998

DILTIAZEM HYDROCHLORIDE

<u>AB3</u>		ACTAVIS ELIZABETH	<u>360MG</u>	<u>A202463</u>	<u>001</u>	Dec 07, 2012
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PRESCRIPTION DRUG PRODUCT LIST

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DILTIAZEM HYDROCHLORIDE

<u>AB3</u>	NOVAST LABS	<u>120MG</u>	<u>A208783 001</u>	Jun 14, 2019
<u>AB3</u>		<u>180MG</u>	<u>A208783 002</u>	Jun 14, 2019
<u>AB3</u>		<u>240MG</u>	<u>A208783 003</u>	Jun 14, 2019
<u>AB3</u>		<u>300MG</u>	<u>A208783 004</u>	Jun 14, 2019
<u>AB3</u>		<u>360MG</u>	<u>A208783 005</u>	Jun 14, 2019
<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>	SUN PHARM	<u>120MG</u>	<u>A090492 001</u>	Oct 28, 2011
<u>AB3</u>		<u>120MG</u>	<u>A203023 001</u>	Jun 08, 2017
<u>AB3</u>		<u>180MG</u>	<u>A090492 002</u>	Oct 28, 2011
<u>AB3</u>		<u>180MG</u>	<u>A203023 002</u>	Jun 08, 2017
<u>AB3</u>		<u>240MG</u>	<u>A090492 003</u>	Oct 28, 2011
<u>AB3</u>		<u>240MG</u>	<u>A203023 003</u>	Jun 08, 2017
<u>AB3</u>		<u>300MG</u>	<u>A090492 004</u>	Oct 28, 2011
<u>AB3</u>		<u>300MG</u>	<u>A203023 004</u>	Jun 08, 2017
<u>AB3</u>		<u>360MG</u>	<u>A090492 005</u>	Oct 28, 2011
<u>AB3</u>		<u>360MG</u>	<u>A203023 005</u>	Jun 08, 2017
<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	<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 PHARM	<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	<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
<u>TAZTIA XT</u>				
<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
<u>TIAZAC</u>				
<u>AB4</u>	+ BAUSCH	<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
INJECTABLE; INJECTION				
<u>DILTIAZEM HYDROCHLORIDE</u>				
<u>AP</u>	AKORN	<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

PRESCRIPTION DRUG PRODUCT LIST

DILTIAZEM HYDROCHLORIDE

INJECTABLE; INJECTION

DILTIAZEM HYDROCHLORIDE

<u>AP</u>	WEST-WARD PHARMS	<u>5MG/ML</u>	<u>A078538</u>	<u>001</u>	Dec 17, 2008
	INT				
	! HOSPIRA	100MG/VIAL	A075853	001	Dec 17, 2002
	SOLUTION; INTRAVENOUS				
	DILTIAZEM HYDROCHLORIDE IN DEXTROSE 5%				
	+! EXELA PHARMA	125MG/125ML (1MG/ML)	N215252	001	Oct 28, 2021
	+!	250MG/250ML (1MG/ML)	N215252	002	Oct 28, 2021

TABLET; ORAL

CARDIZEM

<u>AB</u>	+ BAUSCH	<u>30MG</u>	<u>N018602</u>	<u>001</u>	Nov 05, 1982
<u>AB</u>	+	<u>60MG</u>	<u>N018602</u>	<u>002</u>	Nov 05, 1982
<u>AB</u>	+	<u>90MG</u>	<u>N018602</u>	<u>003</u>	Dec 08, 1986
<u>AB</u>	+!	<u>120MG</u>	<u>N018602</u>	<u>004</u>	Dec 08, 1986

DILTIAZEM HYDROCHLORIDE

<u>AB</u>	EDENBRIDGE PHARMS	<u>30MG</u>	<u>A211596</u>	<u>001</u>	Nov 18, 2019
<u>AB</u>		<u>60MG</u>	<u>A211596</u>	<u>002</u>	Nov 18, 2019
<u>AB</u>		<u>90MG</u>	<u>A211596</u>	<u>003</u>	Nov 18, 2019
<u>AB</u>		<u>120MG</u>	<u>A211596</u>	<u>004</u>	Nov 18, 2019
<u>AB</u>	TEVA	<u>30MG</u>	<u>A074185</u>	<u>001</u>	May 31, 1995
<u>AB</u>		<u>60MG</u>	<u>A074185</u>	<u>002</u>	May 31, 1995
<u>AB</u>		<u>90MG</u>	<u>A074185</u>	<u>003</u>	May 31, 1995
<u>AB</u>		<u>120MG</u>	<u>A074185</u>	<u>004</u>	May 31, 1995

TABLET, EXTENDED RELEASE; ORAL

CARDIZEM LA

<u>AB</u>	+ BAUSCH	<u>120MG</u>	<u>N021392</u>	<u>001</u>	Feb 06, 2003
<u>AB</u>	+	<u>180MG</u>	<u>N021392</u>	<u>002</u>	Feb 06, 2003
<u>AB</u>	+	<u>240MG</u>	<u>N021392</u>	<u>003</u>	Feb 06, 2003
<u>AB</u>	+	<u>300MG</u>	<u>N021392</u>	<u>004</u>	Feb 06, 2003
<u>AB</u>	+	<u>360MG</u>	<u>N021392</u>	<u>005</u>	Feb 06, 2003
<u>AB</u>	+!	<u>420MG</u>	<u>N021392</u>	<u>006</u>	Feb 06, 2003

DILTIAZEM HYDROCHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>120MG</u>	<u>A077686</u>	<u>006</u>	Mar 15, 2010
<u>AB</u>		<u>180MG</u>	<u>A077686</u>	<u>005</u>	Mar 15, 2010
<u>AB</u>		<u>240MG</u>	<u>A077686</u>	<u>004</u>	Mar 15, 2010
<u>AB</u>		<u>300MG</u>	<u>A077686</u>	<u>003</u>	Mar 15, 2010
<u>AB</u>		<u>360MG</u>	<u>A077686</u>	<u>002</u>	Mar 15, 2010
<u>AB</u>		<u>420MG</u>	<u>A077686</u>	<u>001</u>	Mar 15, 2010

DIMENHYDRINATE

INJECTABLE; INJECTION

DIMENHYDRINATE

!	FRESENIUS KABI USA	50MG/ML	A040519	001	Jun 23, 2004
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DIMERCAPROL

INJECTABLE; INJECTION

BAL

+!	AKORN	10%	N005939	001	
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DIMETHYL FUMARATE

CAPSULE, DELAYED RELEASE; ORAL

DIMETHYL FUMARATE

<u>AB</u>	ACCORD HLTHCARE	<u>120MG</u>	<u>A210499</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>		<u>240MG</u>	<u>A210499</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>	ALKEM LABS LTD	<u>120MG</u>	<u>A210440</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>		<u>240MG</u>	<u>A210440</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>	AMNEAL	<u>120MG</u>	<u>A210402</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>		<u>240MG</u>	<u>A210402</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>	CIPLA	<u>120MG</u>	<u>A210305</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>		<u>240MG</u>	<u>A210305</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>	GLENMARK PHARMS LTD	<u>120MG</u>	<u>A210309</u>	<u>001</u>	Oct 06, 2020
<u>AB</u>		<u>240MG</u>	<u>A210309</u>	<u>002</u>	Oct 06, 2020
<u>AB</u>	HETERO LABS LTD III	<u>120MG</u>	<u>A210500</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>		<u>240MG</u>	<u>A210500</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>	LUPIN	<u>120MG</u>	<u>A210226</u>	<u>001</u>	Oct 05, 2020
<u>AB</u>		<u>240MG</u>	<u>A210226</u>	<u>002</u>	Oct 05, 2020
<u>AB</u>	MSN	<u>120MG</u>	<u>A210460</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>		<u>240MG</u>	<u>A210460</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>	MYLAN	<u>120MG</u>	<u>A210531</u>	<u>001</u>	Aug 17, 2020
<u>AB</u>		<u>240MG</u>	<u>A210531</u>	<u>002</u>	Aug 17, 2020
<u>AB</u>	PHARMATHEN	<u>120MG</u>	<u>A210436</u>	<u>001</u>	Mar 26, 2021
<u>AB</u>		<u>240MG</u>	<u>A210436</u>	<u>002</u>	Mar 26, 2021

PRESCRIPTION DRUG PRODUCT LIST

DIROXIMEL FUMARATE

CAPSULE, DELAYED RELEASE;ORAL

VUMERITY

+! BIOGEN INC

231MG

N211855 001 Oct 29, 2019

DISOPYRAMIDE PHOSPHATE

CAPSULE;ORAL

DISOPYRAMIDE PHOSPHATEAB TEVAEQ 100MG BASEA070101 001 Feb 22, 1985ABEQ 150MG BASEA070102 001 Feb 22, 1985NORPACEAB + PFIZEREQ 100MG BASEN017447 001AB +!EQ 150MG BASEN017447 002

CAPSULE, EXTENDED RELEASE;ORAL

NORPACE CR

+ PFIZER

EQ 100MG BASE

N018655 001 Jul 20, 1982

+!

EQ 150MG BASE

N018655 002 Jul 20, 1982

DISULFIRAM

TABLET;ORAL

DISULFIRAMAB ALVOGEN250MGA091681 001 Aug 08, 2013AB CHARTWELL MOLECULES250MGA091563 001 Dec 31, 2012AB500MGA091563 002 Dec 31, 2012AB HIKMA250MGA202652 001 Feb 05, 2014AB !500MGA202652 002 Feb 05, 2014AB SIGMAPHARM LABS LLC250MGA091619 001 Mar 28, 2011AB500MGA091619 002 Mar 28, 2011DIVALPROEX SODIUM

CAPSULE, DELAYED REL PELLETS;ORAL

DEPAKOTEAB +! ABBVIEEQ 125MG VALPROIC ACIDN019680 001 Sep 12, 1989DIVALPROEX SODIUMAB AJANTA PHARMA LTDEQ 125MG VALPROIC ACIDA213181 001 Mar 02, 2020AB DR REDDYS LABS LTDEQ 125MG VALPROIC ACIDA078979 001 Jan 23, 2009AB ZYDUS PHARMS USAEQ 125MG VALPROIC ACIDA078919 001 Jan 27, 2009

INC

TABLET, DELAYED RELEASE;ORAL

DEPAKOTEAB + ABBVIEEQ 125MG VALPROIC ACIDN018723 003 Oct 26, 1984AB +EQ 250MG VALPROIC ACIDN018723 001 Mar 10, 1983AB +!EQ 500MG VALPROIC ACIDN018723 002 Mar 10, 1983DIVALPROEX SODIUMAB ACTAVIS LABS FL INCEQ 500MG VALPROIC ACIDA079080 001 Feb 25, 2011AB APOTEXEQ 125MG VALPROIC ACIDA077615 003 Jul 29, 2008ABEQ 250MG VALPROIC ACIDA077615 002 Jul 29, 2008ABEQ 500MG VALPROIC ACIDA077615 001 Jul 29, 2008AB AUROBINDO PHARMAEQ 125MG VALPROIC ACIDA090554 001 Apr 21, 2011

LTD

ABEQ 250MG VALPROIC ACIDA090554 002 Apr 21, 2011ABEQ 500MG VALPROIC ACIDA090554 003 Apr 21, 2011AB CELLTRIONEQ 125MG VALPROIC ACIDA077296 001 Jul 31, 2008ABEQ 250MG VALPROIC ACIDA077296 002 Jul 31, 2008ABEQ 500MG VALPROIC ACIDA077296 003 Jul 31, 2008AB DR REDDYS LABS LTDEQ 125MG VALPROIC ACIDA078755 001 Jul 29, 2008ABEQ 250MG VALPROIC ACIDA078755 002 Jul 29, 2008ABEQ 500MG VALPROIC ACIDA078755 003 Jul 29, 2008AB INVATECHEQ 125MG VALPROIC ACIDA078290 003 Jul 29, 2008ABEQ 250MG VALPROIC ACIDA078290 002 Jul 29, 2008ABEQ 500MG VALPROIC ACIDA078290 001 Jul 29, 2008AB LUPINEQ 125MG VALPROIC ACIDA078790 001 Jul 29, 2008ABEQ 250MG VALPROIC ACIDA078790 002 Jul 29, 2008ABEQ 500MG VALPROIC ACIDA078790 003 Jul 29, 2008AB ORBION PHARMSEQ 125MG VALPROIC ACIDA078853 001 Nov 25, 2008ABEQ 250MG VALPROIC ACIDA078853 002 Nov 25, 2008ABEQ 500MG VALPROIC ACIDA078853 003 Nov 25, 2008AB PRINSTON INCEQ 125MG VALPROIC ACIDA090210 001 Nov 30, 2009ABEQ 250MG VALPROIC ACIDA090210 002 Nov 30, 2009ABEQ 500MG VALPROIC ACIDA090210 003 Nov 30, 2009AB SUN PHARM INDSEQ 125MG VALPROIC ACIDA078597 001 Jul 29, 2008ABEQ 250MG VALPROIC ACIDA078597 002 Jul 29, 2008ABEQ 500MG VALPROIC ACIDA078597 003 Jul 29, 2008AB UNICHEM LABS LTDEQ 125MG VALPROIC ACIDA079163 001 Apr 05, 2011

PRESCRIPTION DRUG PRODUCT LIST

DIVALPROEX SODIUM

TABLET, DELAYED RELEASE;ORAL

DIVALPROEX SODIUM

<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>	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>	+	ABBVIE	<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>		AMNEAL PHARMS	<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>		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>		LUPIN LTD	<u>EQ 250MG VALPROIC ACID</u>	<u>A209286 001</u>	Oct 18, 2019
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A209286 002</u>	Oct 18, 2019
<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>		HIKMA	<u>EQ 12.5MG BASE/ML</u>	<u>A074277 001</u>	Oct 31, 1994
<u>AP</u>	!	HOSPIRA	<u>EQ 12.5MG BASE/ML</u>	<u>A074086 001</u>	Nov 29, 1993

DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+	!	BAXTER HLTHCARE	<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

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>				<u>160MG/8ML (20MG/ML)</u>	<u>N203551 004</u>	Sep 21, 2015
<u>AP</u>			AMNEAL	<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>			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>			EUGIA PHARMA	<u>20MG/2ML (10MG/ML)</u>	<u>A214575 001</u>	Jun 25, 2021
<u>AP</u>				<u>80MG/8ML (10MG/ML)</u>	<u>A214575 002</u>	Jun 25, 2021
<u>AP</u>				<u>160MG/16ML (10MG/ML)</u>	<u>A214575 003</u>	Jun 25, 2021
<u>AP</u>			GLAND PHARMA LTD	<u>20MG/2ML (10MG/ML)</u>	<u>A213510 001</u>	Jul 01, 2021
<u>AP</u>				<u>80MG/8ML (10MG/ML)</u>	<u>A213510 002</u>	Jul 01, 2021
<u>AP</u>				<u>160MG/16ML (10MG/ML)</u>	<u>A213510 003</u>	Jul 01, 2021
<u>AP</u>			HIKMA	<u>20MG/ML (20MG/ML)</u>	<u>A204490 001</u>	Jan 14, 2021
<u>AP</u>				<u>80MG/4ML (20MG/ML)</u>	<u>A204490 002</u>	Jan 14, 2021
<u>AP</u>			HONG KONG	<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

PRESCRIPTION DRUG PRODUCT LIST

DOCETAXEL

INJECTABLE; INJECTION

DOCETAXEL

<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>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 PHARMS	<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>			<u>160MG/8ML (20MG/ML)</u>	<u>A208137 001</u>	Apr 01, 2019
<u>AP</u>			<u>160MG/16ML (10MG/ML)</u>	<u>A208859 001</u>	Apr 30, 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>		SHILPA	<u>20MG/ML (20MG/ML)</u>	<u>A210327 001</u>	May 16, 2019
<u>AP</u>	+		<u>20MG/ML (20MG/ML)</u>	<u>N205934 001</u>	Dec 22, 2015
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A210327 002</u>	May 16, 2019
<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>A210327 003</u>	May 16, 2019
<u>AP</u>	+		<u>160MG/8ML (20MG/ML)</u>	<u>N205934 003</u>	Dec 22, 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
	+	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

SOLUTION; INTRAVENOUS

DOCETAXEL

<u>AP</u>		SUN PHARM	<u>20MG/ML (20MG/ML)</u>	<u>N022534 003</u>	Jan 08, 2019
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>N022534 004</u>	Jan 08, 2019
<u>AP</u>			<u>160MG/8ML (20MG/ML)</u>	<u>N022534 005</u>	Jan 08, 2019

DOFETILIDE

CAPSULE; ORAL

DOFETILIDE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>0.125MG</u>	<u>A210740 001</u>	Jan 22, 2019
<u>AB</u>			<u>0.25MG</u>	<u>A210740 002</u>	Jan 22, 2019
<u>AB</u>			<u>0.5MG</u>	<u>A210740 003</u>	Jan 22, 2019

DOFETILIDE

<u>AB</u>		ACCORD HLTHCARE	<u>0.125MG</u>	<u>A213338 001</u>	Jun 19, 2020
<u>AB</u>			<u>0.25MG</u>	<u>A213338 002</u>	Jun 19, 2020
<u>AB</u>			<u>0.5MG</u>	<u>A213338 003</u>	Jun 19, 2020
<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>		GRANULES	<u>0.125MG</u>	<u>A212750 001</u>	Oct 14, 2021
<u>AB</u>			<u>0.25MG</u>	<u>A212750 002</u>	Oct 14, 2021
<u>AB</u>			<u>0.5MG</u>	<u>A212750 003</u>	Oct 14, 2021
<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>		MSN	<u>0.125MG</u>	<u>A213220 001</u>	Jan 29, 2020
<u>AB</u>			<u>0.25MG</u>	<u>A213220 002</u>	Jan 29, 2020
<u>AB</u>			<u>0.5MG</u>	<u>A213220 003</u>	Jan 29, 2020
<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>		STRIDES PHARMA	<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>		SUN PHARM	<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

TIKOSYN

<u>AB</u>	+	PFIZER	<u>0.125MG</u>	<u>N020931 001</u>	Oct 01, 1999
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PRESCRIPTION DRUG PRODUCT LIST

DOFETILIDE

CAPSULE;ORAL

TIKOSYN

AB	+		0.25MG	N020931 002	Oct 01, 1999
AB	+	!	0.5MG	N020931 003	Oct 01, 1999

DOLASETRON MESYLATE

TABLET;ORAL

ANZEMET

+	!	VALIDUS PHARMS	50MG	N020623 001	Sep 11, 1997
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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

TABLET, FOR SUSPENSION;ORAL

TIVICAY PD

+	!	VIIV HLTHCARE	EQ 5MG BASE	N213983 001	Jun 12, 2020
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DOLUTEGRAVIR SODIUM; LAMIVUDINE

TABLET;ORAL

DOVATO

+	!	VIIV HLTHCARE	EQ 50MG BASE;300MG	N211994 001	Apr 08, 2019
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DOLUTEGRAVIR SODIUM; RILPIVIRINE HYDROCHLORIDE

TABLET;ORAL

JULUCA

+	!	VIIV HLTHCARE	EQ 50MG BASE;EQ 25MG BASE	N210192 001	Nov 21, 2017
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DONEPEZIL HYDROCHLORIDE

TABLET;ORAL

ARICEPT

AB	+	EISAI INC	5MG	N020690 002	Nov 25, 1996
AB	+	!	10MG	N020690 001	Nov 25, 1996
AB	+	!	23MG	N022568 001	Jul 23, 2010

DONEPEZIL HYDROCHLORIDE

AB		ACI	5MG	A078662 001	May 31, 2011
AB			10MG	A078662 002	May 31, 2011
AB		ALEMBIC PHARMS LTD	5MG	A201724 001	Feb 25, 2013
AB			10MG	A201724 002	Feb 25, 2013
AB		AUROBINDO	5MG	A090056 001	May 31, 2011
AB			10MG	A090056 002	May 31, 2011
AB		CADILA	5MG	A090100 001	Oct 24, 2012
AB			10MG	A090100 002	Oct 24, 2012
AB		CADILA PHARMS LTD	5MG	A204609 001	Sep 19, 2017
AB			10MG	A204609 002	Sep 19, 2017
AB		CIPLA LTD	5MG	A077518 001	May 31, 2011
AB			10MG	A077518 002	May 31, 2011
AB		DEXCEL PHARMA	23MG	A203713 001	Feb 19, 2016
AB		DR REDDYS	23MG	A202723 001	Jul 24, 2013
AB		DR REDDYS LABS LTD	5MG	A201001 001	May 31, 2011
AB			10MG	A201001 002	May 31, 2011
AB		GRAVITI PHARMS	5MG	A202114 001	Jul 05, 2013
AB			10MG	A202114 002	Jul 05, 2013
AB		HETERO LABS LTD V	5MG	A203034 001	Jan 30, 2015
AB			10MG	A203034 002	Jan 30, 2015
AB		INDICUS PHARMA	5MG	A201634 001	Jun 13, 2012
AB			10MG	A201634 002	Jun 13, 2012
AB			23MG	A203419 001	Apr 12, 2016
AB		JUBILANT GENERICS	5MG	A090768 001	May 31, 2011
AB			10MG	A090768 002	May 31, 2011
AB		LUPIN LTD	23MG	A202782 001	Oct 30, 2015
AB		MACLEODS PHARMS LTD	5MG	A201146 001	Aug 17, 2012
AB			10MG	A201146 002	Aug 17, 2012
AB			23MG	A202631 001	Jan 22, 2014
AB		PLIVA HRVATSKA DOO	5MG	A090425 001	May 31, 2011
AB			10MG	A090425 002	May 31, 2011
AB		PRINSTON INC	5MG	A200292 001	May 31, 2011
AB			10MG	A200292 002	May 31, 2011
AB		SANDOZ	5MG	A090290 001	May 31, 2011
AB			10MG	A090290 002	May 31, 2011
AB		SCIEGEN PHARMS INC	5MG	A203907 001	Oct 29, 2014
AB			10MG	A203907 002	Oct 29, 2014

PRESCRIPTION DRUG PRODUCT LIST

DONEPEZIL HYDROCHLORIDE

TABLET;ORAL

DONEPEZIL HYDROCHLORIDE

<u>AB</u>	STRIDES PHARMA	<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>	TORRENT PHARMS	<u>5MG</u>	<u>A090686</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090686</u>	<u>002</u>	May 31, 2011
<u>AB</u>	TWI PHARMS	<u>23MG</u>	<u>A203104</u>	<u>001</u>	Oct 29, 2014
<u>AB</u>	ZYDUS PHARMS	<u>23MG</u>	<u>A203162</u>	<u>001</u>	Aug 31, 2017

TABLET, ORALLY DISINTEGRATING;ORAL

DONEPEZIL HYDROCHLORIDE

<u>AB</u>	HISUN PHARM HANGZHOU	<u>5MG</u>	<u>A205269</u>	<u>001</u>	Jul 27, 2018
<u>AB</u>		<u>10MG</u>	<u>A205269</u>	<u>002</u>	Jul 27, 2018
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A201787</u>	<u>001</u>	Dec 14, 2012
<u>AB</u>		<u>10MG</u>	<u>A201787</u>	<u>002</u>	Dec 14, 2012
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>A091198</u>	<u>001</u>	May 10, 2011
<u>AB</u>	!	<u>10MG</u>	<u>A091198</u>	<u>002</u>	May 10, 2011

DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

NAMZARIC

+	ALLERGAN	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</u>	<u>001</u>	Apr 11, 2018
<u>AP</u>		<u>80MG/ML</u>	<u>A207707</u>	<u>002</u>	Apr 11, 2018
<u>AP</u>	+!	<u>40MG/ML</u>	<u>N018132</u>	<u>001</u>	
<u>AP</u>	+!	<u>80MG/100ML</u>	<u>N018132</u>	<u>002</u>	Feb 04, 1982
<u>AP</u>	+!	<u>80MG/ML</u>	<u>N018132</u>	<u>004</u>	Jul 09, 1982
<u>AP</u>	+!	<u>160MG/100ML</u>	<u>N018132</u>	<u>003</u>	Feb 04, 1982

DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%

<u>AP</u>	+!	B BRAUN	<u>80MG/100ML</u>	<u>N019099</u>	<u>002</u>	Oct 15, 1986
<u>AP</u>	+!		<u>320MG/100ML</u>	<u>N019099</u>	<u>004</u>	Oct 15, 1986

DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+!	B BRAUN	<u>160MG/100ML</u>	<u>N019099</u>	<u>003</u>	Oct 15, 1986
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DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+!	BAXTER HLTHCARE	<u>80MG/100ML</u>	<u>N019615</u>	<u>001</u>	Mar 27, 1987
<u>AP</u>	+!		<u>160MG/100ML</u>	<u>N019615</u>	<u>002</u>	Mar 27, 1987
<u>AP</u>	+!		<u>320MG/100ML</u>	<u>N019615</u>	<u>003</u>	Mar 27, 1987
<u>AP</u>	+!	HOSPIRA	<u>80MG/100ML</u>	<u>N018826</u>	<u>001</u>	Sep 30, 1983
<u>AP</u>	+!		<u>160MG/100ML</u>	<u>N018826</u>	<u>002</u>	Sep 30, 1983
<u>AP</u>	+!		<u>320MG/100ML</u>	<u>N018826</u>	<u>003</u>	Sep 30, 1983

DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER

+	B BRAUN	40MG/100ML	N019099	001	Oct 15, 1986
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DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

+	BAXTER HLTHCARE	640MG/100ML	N019615	004	Mar 27, 1987
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DORAVIRINE

TABLET;ORAL

PIFELTRO

+	MSD MERCK CO	100MG	N210806	001	Aug 30, 2018
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DORAVIRINE; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

DELSTRIGO

+	MSD MERCK CO	100MG;300MG;300MG	N210807	001	Aug 30, 2018
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DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE

<u>AT</u>	AKORN	<u>EQ 2% BASE</u>	<u>A077846</u>	<u>001</u>	Oct 28, 2008
<u>AT</u>	ALEMBIC PHARMS LTD	<u>EQ 2% BASE</u>	<u>A212639</u>	<u>001</u>	Aug 09, 2019
<u>AT</u>	BAUSCH AND LOMB	<u>EQ 2% BASE</u>	<u>A090143</u>	<u>001</u>	Jun 25, 2009
<u>AT</u>	FDC LTD	<u>EQ 2% BASE</u>	<u>A205294</u>	<u>001</u>	Jan 24, 2019
<u>AT</u>	INDOCO	<u>EQ 2% BASE</u>	<u>A202053</u>	<u>001</u>	Sep 11, 2014
<u>AT</u>	MICRO LABS	<u>EQ 2% BASE</u>	<u>A204778</u>	<u>001</u>	Nov 08, 2019
<u>AT</u>	SANDOZ INC	<u>EQ 2% BASE</u>	<u>A078748</u>	<u>001</u>	Nov 06, 2008

PRESCRIPTION DRUG PRODUCT LIST

DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE

<u>AT</u>		<u>EQ 2% BASE</u>	<u>A078981 001</u>	Apr 13, 2009
	<u>TRUSOPT</u>			
<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>	+! AKORN	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>N020869 001</u>	Apr 07, 1998
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COSOPT PF

<u>AT</u>	+! AKORN	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>N202667 001</u>	Feb 01, 2012
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DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE

<u>AT</u>	AKORN	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A077847 001</u>	Oct 28, 2008
<u>AT</u>		<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A203058 001</u>	Sep 22, 2014
<u>AT</u>	ALEMBIC PHARMS LTD	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A213099 001</u>	May 04, 2021
<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>	EUGIA PHARMA	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A207629 001</u>	May 14, 2021
<u>AT</u>		<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A207630 001</u>	Jul 24, 2018
<u>AT</u>	FDC LTD	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A205295 001</u>	Jun 13, 2019
<u>AT</u>	INDOCO	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A202054 001</u>	Sep 03, 2014
<u>AT</u>	MICRO LABS	<u>EQ 2.0% BASE;EQ 0.5% BASE</u>	<u>A204777 001</u>	May 28, 2020
<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>	SOMERSET	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A207523 001</u>	Jun 25, 2019
<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>	+! HIKMA	<u>20MG/ML</u>	<u>N014879 001</u>	
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DOXAPRAM HYDROCHLORIDE

<u>AP</u>	ATHENEX INC	<u>20MG/ML</u>	<u>A076266 001</u>	Jan 10, 2003
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DOXAZOSIN MESYLATE

TABLET; ORAL

CARDURA

<u>AB</u>	+! UPJOHN	<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>	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>	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
<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	<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
BX	HERITAGE PHARMA	EQ 1MG BASE	A205210 001	Feb 13, 2018
BX		EQ 2MG BASE	A205210 002	Feb 13, 2018
BX		EQ 4MG BASE	A205210 003	Feb 13, 2018
BX		EQ 8MG BASE	A205210 004	Feb 13, 2018

PRESCRIPTION DRUG PRODUCT LIST

DOXAZOSIN MESYLATE

TABLET, EXTENDED RELEASE;ORAL

CARDURA XL

+ UPJOHN

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>	AJANTA PHARMA LTD	<u>EQ 10MG BASE</u>	<u>A212624 001</u>	Sep 13, 2019
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A212624 002</u>	Sep 13, 2019
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A212624 003</u>	Sep 13, 2019
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A212624 004</u>	Sep 13, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A212624 005</u>	Sep 13, 2019
<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A215076 001</u>	Apr 21, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A215076 002</u>	Apr 21, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A215076 003</u>	Apr 21, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A215076 004</u>	Apr 21, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A215076 005</u>	Apr 21, 2021
<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>	APPCO	<u>EQ 10MG BASE</u>	<u>A214908 001</u>	Apr 01, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A214908 002</u>	Apr 01, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A214908 003</u>	Apr 01, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214908 004</u>	Apr 01, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A214908 005</u>	Apr 01, 2021
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 10MG BASE</u>	<u>A211603 001</u>	Mar 27, 2019
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A211603 002</u>	Mar 27, 2019
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A211603 003</u>	Mar 27, 2019
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A211603 004</u>	Mar 27, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A211603 005</u>	Mar 27, 2019
<u>AB</u>	CONTRACT PHARMACAL	<u>EQ 10MG BASE</u>	<u>A213474 001</u>	Jul 28, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213474 002</u>	Jul 28, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A213474 003</u>	Jul 28, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A213474 004</u>	Jul 28, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A213474 005</u>	Jul 28, 2020
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 10MG BASE</u>	<u>A210675 001</u>	Oct 16, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A210675 002</u>	Oct 16, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A210675 003</u>	Oct 16, 2020
<u>AB</u>	LEADING PHARMA LLC	<u>EQ 10MG BASE</u>	<u>A211619 001</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A211619 002</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A211619 003</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A211619 004</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A211619 005</u>	Mar 09, 2021
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 150MG BASE</u>	<u>A211618 001</u>	Mar 01, 2021
<u>AB</u>	!	<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 10MG BASE</u>	<u>A071422 002</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A071422 003</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A071422 004</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A071422 005</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A071422 001</u>	Nov 09, 1987
<u>AB</u>	TARO PHARM INDS LTD	<u>EQ 10MG BASE</u>	<u>A213063 001</u>	Jul 01, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213063 002</u>	Jul 01, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A213063 003</u>	Jul 01, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A213063 004</u>	Jul 01, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A213063 005</u>	Jul 01, 2020
!	PAR PHARM	EQ 150MG BASE	A071422 006	Nov 09, 1987
CONCENTRATE;ORAL				
<u>DOXEPIN HYDROCHLORIDE</u>				
<u>AA</u>	!	LANNETT CO INC	<u>EQ 10MG BASE/ML</u>	<u>A074721 001</u> Dec 29, 1998
<u>AA</u>		WOCKHARDT BIO AG	<u>EQ 10MG BASE/ML</u>	<u>A071918 001</u> Jul 20, 1988
CREAM;TOPICAL				
ZONALON				
	+	!	MYLAN	5% N020126 001 Apr 01, 1994

PRESCRIPTION DRUG PRODUCT LIST

DOXEPIN HYDROCHLORIDE

TABLET; ORAL

DOXEPIN HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 3MG BASE</u>	<u>A201951 001</u>	Jul 26, 2013
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A201951 002</u>	Jul 26, 2013
<u>AB</u>	RK PHARMA	<u>EQ 3MG BASE</u>	<u>A202337 001</u>	Jan 20, 2016
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A202337 002</u>	Jan 20, 2016
<u>SILENOR</u>				
<u>AB</u>	+ CURRAX	<u>EQ 3MG BASE</u>	<u>N022036 001</u>	Mar 17, 2010
<u>AB</u>	+!	<u>EQ 6MG BASE</u>	<u>N022036 002</u>	Mar 17, 2010

DOXERCALCIFEROL

CAPSULE; ORAL

DOXERCALCIFEROL

<u>AB</u>	AVET	<u>0.5MCG</u>	<u>A205360 001</u>	Sep 15, 2020
<u>AB</u>		<u>1MCG</u>	<u>A205360 002</u>	Sep 15, 2020
<u>AB</u>		<u>2.5MCG</u>	<u>A205360 003</u>	Sep 15, 2020
<u>AB</u>	RISING	<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>HECTOROL</u>				
<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	<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	<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>	GLAND PHARMA LTD	<u>4MCG/2ML (2MCG/ML)</u>	<u>A210452 001</u>	Sep 26, 2019
<u>AP</u>	HIKMA	<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
<u>AP</u>	LUPIN LTD	<u>4MCG/2ML (2MCG/ML)</u>	<u>A210801 001</u>	Nov 01, 2018
<u>AP</u>	MEITHEAL	<u>4MCG/2ML (2MCG/ML)</u>	<u>A211670 001</u>	Feb 07, 2020
<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

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	<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>	HIKMA	<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
<u>AP</u>	MYLAN LABS LTD	<u>50MG/VIAL</u>	<u>A200170 002</u>	Oct 28, 2011
<u>AP</u>	+! PFIZER	<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>	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>	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
	+ PFIZER	150MG/75ML	N050629 003	Mar 28, 2011

INJECTABLE, LIPOSOMAL; INJECTION

DOXIL (LIPOSOMAL)

<u>AB</u>	+ BAXTER HLTHCARE CORP	<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

PRESCRIPTION DRUG PRODUCT LIST

DOXORUBICIN HYDROCHLORIDE

INJECTABLE, LIPOSOMAL; INJECTION

DOXORUBICIN HYDROCHLORIDE

<u>AB</u>	AYANA PHARMA LTD	<u>20MG/10ML (2MG/ML)</u>	<u>A207228 001</u>	Oct 12, 2021
<u>AB</u>		<u>50MG/25ML (2MG/ML)</u>	<u>A207228 002</u>	Oct 12, 2021
<u>AB</u>	ZYDUS	<u>20MG/10ML (2MG/ML)</u>	<u>A212299 001</u>	Sep 10, 2020
<u>AB</u>		<u>50MG/25ML (2MG/ML)</u>	<u>A212299 002</u>	Sep 10, 2020

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>	!	SUN PHARM	<u>A203263 001</u>	Feb 04, 2013
<u>AB</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>	COSETTE	<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>	STRIDES PHARMA	<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
<u>AB</u>	!	<u>EQ 100MG BASE</u>	<u>A065053 002</u>	Nov 22, 2000
<u>AB</u>	ZYDUS PHARMS	<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>	+	CHARTWELL RX	<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	<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>
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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>	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	<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

PRESCRIPTION DRUG PRODUCT LIST

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>	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>	CHANGZHOU PHARM	<u>EQ 100MG BASE</u>	<u>A209402 001</u>	Oct 07, 2019
<u>AB</u>	CHARTWELL	<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>	NESHER PHARMS	<u>EQ 50MG BASE</u>	<u>A208263 001</u>	Nov 22, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A208263 002</u>	Nov 22, 2021
<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	<u>EQ 50MG BASE</u>	<u>A207774 001</u>	May 31, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A207774 002</u>	May 31, 2018

VIBRAMYCINAB +! PFIZEREQ 100MG BASEN050007 002

INJECTABLE; INJECTION

DOXY 100AP ! FRESENIUS KABI USA EQ 100MG BASE/VIAL A062475 001 Dec 09, 1983DOXY 200AP ! FRESENIUS KABI USA EQ 200MG BASE/VIAL A062475 002 Dec 09, 1983DOXYCYCLINEAP ! HIKMA EQ 100MG BASE/VIAL A062569 001 Mar 09, 1988AP MYLAN LABS LTD EQ 100MG BASE/VIAL A091406 001 Aug 21, 2012AP ZYDUS PHARMS EQ 100MG BASE/VIAL A207757 001 Sep 28, 2017AP EQ 200MG BASE/VIAL A207757 002 Sep 28, 2017DOXYCYCLINE HYCLATEAP WEST-WARD PHARMS EQ 100MG BASE/VIAL A062992 001 Feb 16, 1989

INT

AP EQ 200MG BASE/VIAL A062992 002 Feb 16, 1989

TABLET; ORAL

ACTICLATEAB + ALMIRALL EQ 75MG BASE N205931 001 Jul 25, 2014AB +! EQ 150MG BASE N205931 002 Jul 25, 2014DOXYCYCLINE HYCLATEAB ACELLA EQ 100MG BASE A210664 001 Mar 16, 2020AB ACTAVIS LABS FL INC EQ 100MG BASE A062421 001 Feb 02, 1983AB AJANTA PHARMA LTD EQ 75MG BASE A211584 001 Jun 01, 2020AB EQ 150MG BASE A211584 002 Jun 01, 2020AB ALEMBIC PHARMS LTD EQ 20MG BASE A210537 001 Mar 03, 2020AB EQ 50MG BASE A210536 002 Sep 14, 2021AB EQ 75MG BASE A211744 001 Jun 30, 2020AB EQ 150MG BASE A211744 002 Jun 30, 2020AB APOTEX EQ 75MG BASE A209243 001 Apr 15, 2019AB EQ 150MG BASE A209243 002 Apr 15, 2019AB CADILA EQ 100MG BASE A207773 001 Oct 30, 2017AB CARIBE HOLDINGS EQ 50MG BASE A062269 003 Oct 05, 1983AB ! EQ 100MG BASE A062269 002 Nov 08, 1982AB CHANGZHOU PHARM EQ 100MG BASE A211343 001 Oct 09, 2019AB CHARTWELL EQ 50MG BASE A062505 002 Jul 27, 2021AB EQ 100MG BASE A062505 001 Sep 11, 1984AB EMCURE PHARMS LTD EQ 100MG BASE A209969 001 Nov 09, 2018AB EPIC PHARMA LLC EQ 20MG BASE A065182 001 May 13, 2005AB EQ 75MG BASE A214207 001 Dec 16, 2020AB EQ 150MG BASE A214207 002 Dec 16, 2020AB HIKMA INTL PHARMS EQ 100MG BASE A065095 001 Jul 02, 2003AB ! LANNETT CO INC EQ 20MG BASE A065277 001 Nov 10, 2005AB LARKEN LABS EQ 20MG BASE A065287 001 Feb 28, 2006AB LUPIN LTD EQ 75MG BASE A208818 001 Sep 27, 2017AB EQ 150MG BASE A208818 002 Sep 27, 2017AB MAYNE PHARMA INC EQ 75MG BASE A208765 001 Jun 14, 2017AB EQ 150MG BASE A208765 002 Jun 14, 2017

PRESCRIPTION DRUG PRODUCT LIST

DOXYCYCLINE HYCLATE

TABLET;ORAL

DOXYCYCLINE HYCLATE

<u>AB</u>	NOVEL LABS INC	<u>EQ 100MG BASE</u>	<u>A207558 001</u>	Sep 06, 2017
<u>AB</u>	SUN PHARM INDUSTRIES	<u>EQ 20MG BASE</u>	<u>A065134 001</u>	May 13, 2005
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062677 001</u>	Jul 10, 1986

TABLET, DELAYED RELEASE;ORAL

DORYX

<u>AB</u>	+	MAYNE PHARMA	<u>EQ 50MG BASE</u>	<u>N050795 006</u>	Dec 19, 2014
<u>AB</u>	+		<u>EQ 75MG BASE</u>	<u>N050795 001</u>	May 06, 2005
<u>AB</u>	+		<u>EQ 100MG BASE</u>	<u>N050795 002</u>	May 06, 2005
<u>AB</u>	+		<u>EQ 150MG BASE</u>	<u>N050795 003</u>	Jun 20, 2008
<u>AB</u>	+	!	<u>EQ 200MG BASE</u>	<u>N050795 005</u>	Apr 11, 2013

DOXYCYCLINE HYCLATE

<u>AB</u>		ACTAVIS ELIZABETH	<u>EQ 50MG BASE</u>	<u>A090134 003</u>	May 22, 2018
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A090134 001</u>	Dec 14, 2011
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A090134 002</u>	Dec 14, 2011
<u>AB</u>			<u>EQ 200MG BASE</u>	<u>A090134 004</u>	May 22, 2018
<u>AB</u>		ALEMBIC PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A213075 001</u>	Jan 03, 2022
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A213075 002</u>	Jan 03, 2022
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A213075 003</u>	Jan 03, 2022
<u>AB</u>			<u>EQ 200MG BASE</u>	<u>A213075 004</u>	Jan 03, 2022
<u>AB</u>		HERITAGE PHARMS INC	<u>EQ 75MG BASE</u>	<u>A200856 001</u>	Apr 30, 2013
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A200856 002</u>	Apr 30, 2013
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A200856 003</u>	Apr 30, 2013
<u>AB</u>			<u>EQ 200MG BASE</u>	<u>A200856 004</u>	Nov 13, 2018
<u>AB</u>		PRINSTON INC	<u>EQ 50MG BASE</u>	<u>A207494 003</u>	Feb 19, 2019
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A207494 001</u>	Nov 15, 2016
<u>AB</u>			<u>EQ 200MG BASE</u>	<u>A207494 002</u>	Nov 15, 2016

DORYX

	+	MAYNE PHARMA	EQ 80MG BASE	N050795 004	Apr 11, 2013
		DORYX MPC			
	+	!	MAYNE PHARMA	EQ 120MG BASE	N050795 008

DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE

TABLET, DELAYED RELEASE;ORAL

DICLEGIS

<u>AB</u>	+	!	DUCHESNAY	<u>10MG;10MG</u>	<u>N021876 001</u>	Apr 08, 2013
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DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE

<u>AB</u>		ACTAVIS LABS FL INC	<u>10MG;10MG</u>	<u>A205811 001</u>	Aug 19, 2016
<u>AB</u>		MYLAN PHARMS INC	<u>10MG;10MG</u>	<u>A207825 001</u>	Jul 06, 2020
<u>AB</u>		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
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DRONABINOL

CAPSULE;ORAL

DRONABINOL

<u>AB</u>		AKORN	<u>2.5MG</u>	<u>A079217 001</u>	Jun 20, 2014
<u>AB</u>			<u>5MG</u>	<u>A079217 002</u>	Jun 20, 2014
<u>AB</u>			<u>10MG</u>	<u>A079217 003</u>	Jun 20, 2014
<u>AB</u>		ASCENT PHARMS INC	<u>2.5MG</u>	<u>A207421 001</u>	Feb 07, 2020
<u>AB</u>			<u>5MG</u>	<u>A207421 002</u>	Feb 07, 2020
<u>AB</u>			<u>10MG</u>	<u>A207421 003</u>	Feb 07, 2020
<u>AB</u>		LANNETT CO INC	<u>2.5MG</u>	<u>A201463 001</u>	May 18, 2018
<u>AB</u>			<u>5MG</u>	<u>A201463 002</u>	May 18, 2018
<u>AB</u>			<u>10MG</u>	<u>A201463 003</u>	May 18, 2018
<u>AB</u>		SVC PHARMA	<u>2.5MG</u>	<u>A078292 001</u>	Jun 27, 2008
<u>AB</u>			<u>5MG</u>	<u>A078292 002</u>	Jun 27, 2008
<u>AB</u>			<u>10MG</u>	<u>A078292 003</u>	Jun 27, 2008

MARINOL

<u>AB</u>	+	ALKEM LABS LTD	<u>2.5MG</u>	<u>N018651 001</u>	May 31, 1985
<u>AB</u>	+	!	<u>5MG</u>	<u>N018651 002</u>	May 31, 1985
<u>AB</u>	+		<u>10MG</u>	<u>N018651 003</u>	May 31, 1985

SOLUTION;ORAL

SYNDROS

	+	!	BENUVIA	5MG/ML	N205525 001	Mar 23, 2017
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PRESCRIPTION DRUG PRODUCT LIST

DRONEDARONE HYDROCHLORIDE

TABLET; ORAL

MULTAQ

+! SANOFI AVENTIS US EQ 400MG BASE N022425 001 Jul 01, 2009

DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL**AP** AM REGENT 2.5MG/ML A072123 001 Oct 24, 1988**AP** EUROHLTH INTL SARL 2.5MG/ML A208197 001 Dec 14, 2017**AP** HOSPIRA 2.5MG/ML A071981 001 Feb 29, 1988INAPSINE**AP** +! AKORN 2.5MG/ML N016796 001DROSPIRENONE

TABLET; ORAL

SLYND

+! EXELTIS USA INC 4MG N211367 001 May 23, 2019

DROSPIRENONE; ESTETROL

TABLET; ORAL

NEXTSTELLIS

+! MAYNE PHARMA 3MG;14.2MG N214154 001 Apr 15, 2021

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 3MG;0.02MG A078515 001 Mar 30, 2009**AB** GLENMARK PHARMS LTD 3MG;0.02MG A204296 001 Aug 17, 2015**AB** HETERO LABS LTD 3MG;0.02MG A211944 001 Mar 22, 2019**AB** HLTHCARE 3MG;0.02MG A203291 001 Jul 18, 2017**AB** MYLAN LABS LTD 3MG;0.02MG A202594 001 Oct 22, 2015**AB** WATSON LABS 3MG;0.02MG A078833 001 Nov 28, 2011LO-ZUMANDIMINE**AB** AUROBINDO PHARMA LTD 3MG;0.02MG A209632 001 Feb 27, 2018LORYNA**AB** XIROMED 3MG;0.02MG A079221 001 Mar 28, 2011MELAMISA**AB** NOVAST LABS 3MG;0.02MG A202016 001 Jan 26, 2016NIKKI**AB** LUPIN LTD 3MG;0.02MG A201661 001 May 27, 2014YAZ**AB** +! BAYER HLTHCARE 3MG;0.02MG N021676 001 Mar 16, 2006

TABLET; ORAL-28

DROSPIRENONE AND ETHINYL ESTRADIOL**AB** BARR 3MG;0.03MG A077527 001 May 09, 2008**AB** GLENMARK PHARMS LTD 3MG;0.03MG A204848 001 Mar 25, 2016**AB** HETERO LABS LTD 3MG;0.03MG A213034 001 Jan 24, 2020**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**AB** NAARI PTE LTD 3MG;0.03MG A207245 001 Nov 22, 2016SYEDA**AB** XIROMED 3MG;0.03MG A090114 001 Mar 28, 2011YAELA**AB** NOVAST LABS 3MG;0.03MG A202015 001 Nov 19, 2014YASMIN**AB** +! BAYER HLTHCARE 3MG;0.03MG N021098 001 May 11, 2001ZUMANDIMINE**AB** AUROBINDO PHARMA LTD 3MG;0.03MG A209407 001 Mar 26, 2018DROSPIRENONE; 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, 2010DROSPIRENONE, 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

PRESCRIPTION DRUG PRODUCT LIST

DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM

TABLET;ORAL

DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM

AB		<u>3MG,N/A;0.03MG,N/A;0.451MG,0.451MG</u>	<u>A203594 001</u>	Oct 11, 2016
	<u>SAFYRAL</u>			
AB	+! BAYER HLTHCARE	<u>3MG,N/A;0.03MG,N/A;0.451MG,0.451MG</u>	<u>N022574 001</u>	Dec 16, 2010
	<u>TYDEMY</u>			
AB	LUPIN LTD	<u>3MG,N/A;0.03MG,N/A;0.451MG,0.451MG</u>	<u>A205948 001</u>	Dec 12, 2017

DROXIDOPA

CAPSULE;ORAL

DROXIDOPA

AB	AJANTA PHARMA LTD	<u>100MG</u>	<u>A214391 001</u>	Feb 18, 2021
AB		<u>200MG</u>	<u>A214391 002</u>	Feb 18, 2021
AB		<u>300MG</u>	<u>A214391 003</u>	Feb 18, 2021
AB	ALKEM LABS LTD	<u>100MG</u>	<u>A213911 001</u>	Feb 18, 2021
AB		<u>200MG</u>	<u>A213911 002</u>	Feb 18, 2021
AB		<u>300MG</u>	<u>A213911 003</u>	Feb 18, 2021
AB	ANNORA	<u>100MG</u>	<u>A211726 002</u>	Aug 09, 2021
AB		<u>200MG</u>	<u>A211726 003</u>	Aug 09, 2021
AB		<u>300MG</u>	<u>A211726 001</u>	Feb 18, 2021
AB	AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A214387 001</u>	Feb 18, 2021
AB		<u>200MG</u>	<u>A214387 002</u>	Feb 18, 2021
AB		<u>300MG</u>	<u>A214387 003</u>	Feb 18, 2021
AB	BIONPHARMA INC	<u>100MG</u>	<u>A213033 001</u>	Apr 28, 2021
AB		<u>200MG</u>	<u>A213033 002</u>	Apr 28, 2021
AB		<u>300MG</u>	<u>A213033 003</u>	Apr 28, 2021
AB	HIKMA	<u>100MG</u>	<u>A212835 001</u>	Feb 18, 2021
AB		<u>200MG</u>	<u>A212835 002</u>	Feb 18, 2021
AB		<u>300MG</u>	<u>A212835 003</u>	Feb 18, 2021
AB	LUPIN PHARMS	<u>100MG</u>	<u>A211652 001</u>	Feb 18, 2021
AB		<u>200MG</u>	<u>A211652 002</u>	Feb 18, 2021
AB		<u>300MG</u>	<u>A211652 003</u>	Feb 18, 2021
AB	MSN PHARMS INC	<u>100MG</u>	<u>A211741 001</u>	Feb 18, 2021
AB		<u>200MG</u>	<u>A211741 002</u>	Feb 18, 2021
AB		<u>300MG</u>	<u>A211741 003</u>	Feb 18, 2021
AB	SCIEGEN PHARMS INC	<u>100MG</u>	<u>A214017 001</u>	Feb 18, 2021
AB		<u>200MG</u>	<u>A214017 002</u>	Feb 18, 2021
AB		<u>300MG</u>	<u>A214017 003</u>	Feb 18, 2021
AB	SLATE	<u>100MG</u>	<u>A215265 001</u>	Nov 01, 2021
AB		<u>200MG</u>	<u>A215265 002</u>	Nov 01, 2021
AB		<u>300MG</u>	<u>A215265 003</u>	Nov 01, 2021
AB	SUN PHARM	<u>100MG</u>	<u>A214384 001</u>	Feb 18, 2021
AB		<u>200MG</u>	<u>A214384 002</u>	Feb 18, 2021
AB		<u>300MG</u>	<u>A214384 003</u>	Feb 18, 2021
AB	TASMAN PHARMA	<u>100MG</u>	<u>A213661 001</u>	Feb 18, 2021
AB		<u>200MG</u>	<u>A213661 002</u>	Feb 18, 2021
AB		<u>300MG</u>	<u>A213661 003</u>	Feb 18, 2021
AB	TRIS PHARMA INC	<u>100MG</u>	<u>A214543 001</u>	May 05, 2021
AB		<u>200MG</u>	<u>A214543 002</u>	May 05, 2021
AB		<u>300MG</u>	<u>A214543 003</u>	May 05, 2021
AB	ZYDUS PHARMS	<u>100MG</u>	<u>A211818 001</u>	Feb 18, 2021
AB		<u>200MG</u>	<u>A211818 002</u>	Feb 18, 2021
AB		<u>300MG</u>	<u>A211818 003</u>	Feb 18, 2021
	<u>NORTHERA</u>			
AB	+ LUNDBECK NA LTD	<u>100MG</u>	<u>N203202 001</u>	Feb 18, 2014
AB	+	<u>200MG</u>	<u>N203202 002</u>	Feb 18, 2014
AB	+!	<u>300MG</u>	<u>N203202 003</u>	Feb 18, 2014

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS;ORAL

CYMBALTA

AB	+ LILLY	<u>EQ 20MG BASE</u>	<u>N021427 001</u>	Aug 03, 2004
AB	+	<u>EQ 30MG BASE</u>	<u>N021427 002</u>	Aug 03, 2004
AB	+!	<u>EQ 60MG BASE</u>	<u>N021427 004</u>	Aug 03, 2004

DULOXETINE HYDROCHLORIDE

AB	ACTAVIS ELIZABETH	<u>EQ 20MG BASE</u>	<u>A090776 001</u>	Dec 17, 2013
AB		<u>EQ 30MG BASE</u>	<u>A090776 002</u>	Dec 17, 2013
AB		<u>EQ 60MG BASE</u>	<u>A090776 003</u>	Dec 17, 2013
AB	AJANTA PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A208706 001</u>	Jan 06, 2017
AB		<u>EQ 30MG BASE</u>	<u>A208706 002</u>	Jan 06, 2017
AB		<u>EQ 40MG BASE</u>	<u>A208706 004</u>	Mar 11, 2019
AB		<u>EQ 60MG BASE</u>	<u>A208706 003</u>	Jan 06, 2017

PRESCRIPTION DRUG PRODUCT LIST

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS;ORAL

DULOXETINE HYDROCHLORIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A202949 001</u>	Jun 09, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A202949 002</u>	Jun 09, 2014
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A202949 003</u>	Jun 09, 2014
<u>AB</u>	ALKEM LABS LTD	<u>EQ 20MG BASE</u>	<u>A203197 001</u>	Aug 26, 2015
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A203197 002</u>	Aug 26, 2015
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A203197 003</u>	Aug 26, 2015
<u>AB</u>	ANCHEN PHARMS	<u>EQ 20MG BASE</u>	<u>A090780 001</u>	Oct 28, 2015
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090780 002</u>	Oct 28, 2015
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090780 003</u>	Oct 28, 2015
<u>AB</u>	APOTEX	<u>EQ 20MG BASE</u>	<u>A202045 001</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A202045 002</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A202045 003</u>	Jun 11, 2014
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A090778 001</u>	Dec 11, 2013
<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	<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	<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	<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>	QINGDAO BAHEAL PHARM	<u>EQ 20MG BASE</u>	<u>A210599 001</u>	Apr 17, 2019
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A210599 002</u>	Apr 17, 2019
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A210599 003</u>	Apr 17, 2019
<u>AB</u>	SUN PHARM	<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>	SUNSHINE	<u>EQ 20MG BASE</u>	<u>A212328 001</u>	Feb 11, 2021
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A212328 002</u>	Feb 11, 2021
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A212328 003</u>	Feb 11, 2021
<u>AB</u>	TORRENT	<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	<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
	DRIZALMA SPRINKLE			
+	SUN PHARMA GLOBAL	EQ 20MG BASE	N212516 001	Jul 19, 2019
+		EQ 30MG BASE	N212516 002	Jul 19, 2019
+		EQ 40MG BASE	N212516 003	Jul 19, 2019
+		EQ 60MG BASE	N212516 004	Jul 19, 2019

PRESCRIPTION DRUG PRODUCT LIST

DUTASTERIDE

CAPSULE; ORAL

AVODART

AB	+ !	WOODWARD	0.5MG	N021319	001	Nov 20, 2001
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DUTASTERIDE

AB		ACELLA	0.5MG	A206373	001	Mar 17, 2016
AB		AMNEAL PHARMS	0.5MG	A203118	001	Nov 20, 2015
AB		ASCENT PHARMS INC	0.5MG	A206574	001	Oct 21, 2016
AB		AUROBINDO PHARMA LTD	0.5MG	A202660	001	Nov 20, 2015
AB		BARR	0.5MG	A090095	001	Dec 21, 2010
AB		BIONPHARMA INC	0.5MG	A200899	001	Nov 20, 2015
AB		CADILA	0.5MG	A204373	001	Oct 04, 2017
AB		HUMANWELL PURACAP	0.5MG	A209909	001	Nov 21, 2017
AB		MARKSANS PHARMA	0.5MG	A204376	001	Apr 07, 2017
AB		STRIDES PHARMA	0.5MG	A204262	001	Nov 20, 2015
AB		VINTAGE	0.5MG	A202421	001	Nov 20, 2015

DUTASTERIDE; TAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL

DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE

AB		ANCHEN PHARMS	0.5MG; 0.4MG	A202509	001	Feb 26, 2014
AB		ZYDUS PHARMS	0.5MG; 0.4MG	A207769	001	May 24, 2018
AB	+ !	WOODWARD	0.5MG; 0.4MG	N022460	001	Jun 14, 2010

DUVELISIB

CAPSULE; ORAL

COPIKTRA

	+	SECURA	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

	+ !	FERA PHARMS LLC	0.125%	N011963	001	
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ECONAZOLE NITRATE

AEROSOL, FOAM; TOPICAL

ECOZA

	+ !	RESILIA PHARMS	1%	N205175	001	Oct 24, 2013
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CREAM; TOPICAL

ECONAZOLE NITRATE

AB	!	PADAGIS ISRAEL	1%	A076479	001	Jun 23, 2004
AB		TARO	1%	A076005	001	Nov 26, 2002
AB		TELGENT	1%	A076574	001	Dec 17, 2004

SPECTAZOLE

AB	+	ALVOGEN	1%	N018751	001	Dec 23, 1982
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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

	+ !	BAUSCH	200MG/ML	N008922	001	
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EDOXABAN 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

PRESCRIPTION DRUG PRODUCT LIST

EFAVIRENZ

CAPSULE; ORAL

EFAVIRENZ

AB	AUROBINDO PHARMA LTD	50MG	A078064 001	Dec 15, 2017
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AB		200MG	A078064 003	Dec 15, 2017
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SUSTIVA

AB	+ BRISTOL MYERS SQUIBB	50MG	N020972 001	Sep 17, 1998
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AB	+!	200MG	N020972 003	Sep 17, 1998
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EFAVIRENZ

	AUROBINDO PHARMA LTD	100MG	A078064 002	Dec 15, 2017
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TABLET; ORAL

EFAVIRENZ

AB	AUROBINDO PHARMA LTD	600MG	A077673 001	Sep 21, 2018
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AB		600MG	A205322 001	Aug 30, 2018
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AB	CIPLA	600MG	A204766 001	Jun 15, 2018
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AB	HETERO LABS LTD III	600MG	A078886 001	Apr 27, 2018
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AB	STRIDES PHARMA	600MG	A204869 001	Mar 12, 2018
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SUSTIVA

AB	+! BRISTOL MYERS SQUIBB	600MG	N021360 002	Feb 01, 2002
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EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

ATRIPLA

AB	+! GILEAD SCIENCES	600MG; 200MG; 300MG	N021937 001	Jul 12, 2006
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EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE

AB	AUROBINDO PHARMA LTD	600MG; 200MG; 300MG	A203041 001	Sep 04, 2018
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AB	CIPLA	600MG; 200MG; 300MG	A206894 001	Jun 03, 2019
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AB	LAURUS	600MG; 200MG; 300MG	A213541 001	Dec 22, 2021
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AB	MACLEODS PHARMS LTD	600MG; 200MG; 300MG	A204287 001	Sep 13, 2021
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AB	TEVA PHARMS USA	600MG; 200MG; 300MG	A091215 001	Nov 09, 2018
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EFAVIRENZ; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

EFAVIRENZ, LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE

AB	LAURUS	400MG; 300MG; 300MG	A213038 001	May 14, 2020
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AB		600MG; 300MG; 300MG	A212786 001	May 14, 2020
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SYMFI

AB	+! MYLAN LABS LTD	600MG; 300MG; 300MG	N022142 001	Mar 22, 2018
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SYMFI LO

AB	+! MYLAN	400MG; 300MG; 300MG	N208255 001	Feb 05, 2018
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EFINACONAZOLE

SOLUTION; TOPICAL

EFINACONAZOLE

AB	ACRUX DDS	10%	A211969 001	Jun 21, 2021
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AB	TEVA PHARMS USA	10%	A211827 001	Dec 16, 2020
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JUBLIA

AB	+! BAUSCH	10%	N203567 001	Jun 06, 2014
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EFLORNITHINE HYDROCHLORIDE

CREAM; TOPICAL

VANIQA

	+! SKINMEDICA	13.9%	N021145 001	Jul 27, 2000
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ELAGOLIX SODIUM

TABLET; ORAL

ORILISSA

	+ ABBVIE INC	EQ 150MG BASE	N210450 001	Jul 23, 2018
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	+!	EQ 200MG BASE	N210450 002	Jul 23, 2018
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ELAGOLIX SODIUM, ESTRADIOL, NORETHINDRONE ACETATE; ELAGOLIX SODIUM

CAPSULE; ORAL

ORIAHNN (COPACKAGED)

	+! ABBVIE INC	EQ 300MG BASE, 1MG, 0.5MG; EQ 300MG BASE	N213388 001	May 29, 2020
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PRESCRIPTION DRUG PRODUCT LIST

ELBASVIR; GRAZOPREVIR

TABLET; ORAL

ZEPATIER

+! MERCK SHARP DOHME 50MG; 100MG N208261 001 Jan 28, 2016

ELETRIPTAN HYDROBROMIDE

TABLET; ORAL

ELETRIPTAN HYDROBROMIDEAB AJANTA PHARMA LTD EQ 20MG BASE A205186 001 Aug 29, 2017AB EQ 40MG BASE A205186 002 Aug 29, 2017AB AUROBINDO PHARMA LTD EQ 20MG BASE A210708 001 Jan 15, 2019AB EQ 40MG BASE A210708 002 Jan 15, 2019AB MYLAN EQ 20MG BASE A205152 001 Aug 11, 2017AB EQ 40MG BASE A205152 002 Aug 11, 2017AB TEVA PHARMS USA EQ 20MG BASE A202040 001 Jun 27, 2017AB EQ 40MG BASE A202040 002 Jun 27, 2017AB ZYDUS PHARMS EQ 20MG BASE A206409 001 Jun 16, 2017AB EQ 40MG BASE A206409 002 Jun 16, 2017RELPAKAB + UPJOHN EQ 20MG BASE N021016 001 Dec 26, 2002AB +! EQ 40MG BASE N021016 002 Dec 26, 2002ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR

TABLET; ORAL

TRIKAFTA (COPACKAGED)

+ VERTEX PHARMS INC 50MG, 37.5MG, 25MG; 75MG N212273 002 Jun 08, 2021

+! 100MG, 75MG, 50MG; 150MG N212273 001 Oct 21, 2019

ELIGLUSTAT TARTRATE

CAPSULE; ORAL

CERDELGAAB +! GENZYME CORP EQ 84MG BASE N205494 001 Aug 19, 2014ELIGLUSTAT TARTRATEAB APPCO EQ 84MG BASE A212463 001 Sep 08, 2021AB TEVA PHARMS USA INC EQ 84MG BASE A212474 001 Dec 27, 2021ELTROMBOPAG OLAMINE

FOR SUSPENSION; ORAL

PROMACTA KIT

+ NOVARTIS EQ 12.5MG ACID/PACKET N207027 002 Sep 27, 2018

+! EQ 25MG ACID/PACKET N207027 001 Aug 24, 2015

TABLET; ORAL

PROMACTA

+ NOVARTIS EQ 12.5MG ACID N022291 004 Oct 20, 2011

+ EQ 25MG ACID N022291 001 Nov 20, 2008

+ EQ 50MG ACID N022291 002 Nov 20, 2008

+! EQ 75MG ACID N022291 003 Sep 08, 2009

ELUXADOLINE

TABLET; ORAL

VIBERZI

+ ALLERGAN HOLDINGS 75MG N206940 001 May 27, 2015

+! 100MG N206940 002 May 27, 2015

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; LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TRIJARDY XR

+ BOEHRINGER 5MG; 2.5MG; 1GM N212614 001 Jan 27, 2020

+ INGELHEIM 10MG; 5MG; 1GM N212614 002 Jan 27, 2020

+ 12.5MG; 2.5MG; 1GM N212614 003 Jan 27, 2020

+! 25MG; 5MG; 1GM N212614 004 Jan 27, 2020

PRESCRIPTION DRUG PRODUCT LIST

EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET;ORAL

SYNJARDY

+	BOEHRINGER INGELHEIM	5MG;500MG	N206111	001	Aug 26, 2015
+		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 INGELHEIM	5MG;1GM	N208658	001	Dec 09, 2016
+		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
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EMTRIVA

AB	+	GILEAD	200MG	N021500	001	Jul 02, 2003
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SOLUTION;ORAL

EMTRIVA

+	!	GILEAD	10MG/ML	N021896	001	Sep 28, 2005
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EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

ODEFSEY

+	!	GILEAD SCIENCES INC	200MG;EQ 25MG BASE;EQ 25MG BASE	N208351	001	Mar 01, 2016
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EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

COMPLERA

+	!	GILEAD SCIENCES INC	200MG;EQ 25MG BASE;300MG	N202123	001	Aug 10, 2011
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EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

DESCOVY

+	!	GILEAD SCIENCES INC	200MG;EQ 25MG BASE	N208215	001	Apr 04, 2016
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EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE

AB	HETERO LABS LTD III	200MG;300MG	A201806	001	Oct 07, 2021
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EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE

AB	AMNEAL PHARMS CO	100MG;150MG	A209721	001	Aug 22, 2018
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AB		133MG;200MG	A209721	002	Aug 22, 2018
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AB		167MG;250MG	A209721	003	Aug 22, 2018
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AB		200MG;300MG	A209721	004	Aug 22, 2018
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AB	APOTEX	200MG;300MG	A208740	001	Jun 16, 2021
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AB	AUROBINDO PHARMA LTD	200MG;300MG	A090513	001	Jan 26, 2018
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AB	CIPLA	200MG;300MG	A090958	001	Apr 02, 2021
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AB	LAURUS	200MG;300MG	A212114	001	Jul 26, 2019
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AB	LUPIN LTD	200MG;300MG	A204131	001	Jun 04, 2021
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AB	MACLEODS PHARMS LTD	200MG;300MG	A203442	001	May 15, 2020
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AB	MYLAN	200MG;300MG	A206436	001	Apr 09, 2018
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AB	STRIDES PHARMA	200MG;300MG	A091055	001	Jan 13, 2021
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AB	TEVA PHARMS USA	200MG;300MG	A090894	001	Jun 08, 2017
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AB	ZYDUS PHARMS	100MG;150MG	A212689	002	Jul 01, 2021
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AB		133MG;200MG	A212689	003	Jul 01, 2021
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AB		167MG;250MG	A212689	004	Jul 01, 2021
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AB		200MG;300MG	A212689	001	Feb 28, 2020
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TRUVADA

AB	+	GILEAD	100MG;150MG	N021752	002	Mar 10, 2016
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AB	+		133MG;200MG	N021752	003	Mar 10, 2016
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AB	+		167MG;250MG	N021752	004	Mar 10, 2016
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AB	+	!	200MG;300MG	N021752	001	Aug 02, 2004
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PRESCRIPTION DRUG PRODUCT LIST

ENALAPRIL MALEATE

SOLUTION;ORAL

ENALAPRIL MALEATE

AB	BIONPHARMA INC	1MG/ML	A212408 001	Aug 10, 2021
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EPANED

AB	+!	AZURITY	1MG/ML	N208686 001	Sep 20, 2016
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TABLET;ORAL

ENALAPRIL MALEATE

AB	HERITAGE PHARMA	2.5MG	A075479 001	Aug 22, 2000
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AB		5MG	A075479 002	Aug 22, 2000
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AB		10MG	A075479 003	Aug 22, 2000
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AB		20MG	A075479 004	Aug 22, 2000
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AB	SANDOZ INC	2.5MG	A075496 001	Aug 22, 2000
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AB		5MG	A075496 002	Aug 22, 2000
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AB		10MG	A075459 001	Aug 22, 2000
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AB		20MG	A075459 002	Aug 22, 2000
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AB	TARO	2.5MG	A075657 001	Jan 23, 2001
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AB		5MG	A075657 002	Jan 23, 2001
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AB		10MG	A075657 003	Jan 23, 2001
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AB		20MG	A075657 004	Jan 23, 2001
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AB	WOCKHARDT LTD	2.5MG	A075483 001	Aug 22, 2000
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AB		5MG	A075483 002	Aug 22, 2000
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AB		10MG	A075483 003	Aug 22, 2000
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AB		20MG	A075483 004	Aug 22, 2000
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VASOTEC

AB	+	BAUSCH	2.5MG	N018998 005	Jul 26, 1988
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AB	+		5MG	N018998 001	Dec 24, 1985
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AB	+		10MG	N018998 002	Dec 24, 1985
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AB	+!		20MG	N018998 003	Dec 24, 1985
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ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE

AB	COSETTE	5MG;12.5MG	A075727 001	Sep 18, 2001
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AB		10MG;25MG	A075727 002	Sep 18, 2001
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AB	DR REDDYS LABS LTD	5MG;12.5MG	A075909 001	Oct 15, 2001
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AB		10MG;25MG	A075909 002	Oct 15, 2001
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AB	TARO PHARM INDS	5MG;12.5MG	A075788 001	Sep 18, 2001
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AB		10MG;25MG	A075788 002	Sep 18, 2001
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VASERETIC

AB	+	BAUSCH	5MG;12.5MG	N019221 003	Jul 12, 1995
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AB	+!		10MG;25MG	N019221 001	Oct 31, 1986
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ENALAPRILAT

INJECTABLE;INJECTION

ENALAPRILAT

AP	!	ATHENEX INC	1.25MG/ML	A075634 001	Aug 22, 2000
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AP		HIKMA FARMACEUTICA	1.25MG/ML	A078687 001	Dec 23, 2008
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AP	!	HOSPIRA	1.25MG/ML	A075458 001	Aug 22, 2000
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ENASIDENIB MESYLATE

TABLET;ORAL

IDHIFA

+	CELGENE CORP	EQ 50MG BASE	N209606 001	Aug 01, 2017
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+		EQ 100MG BASE	N209606 002	Aug 01, 2017
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ENCORAFENIB

CAPSULE;ORAL

BRAFTOVI

+	ARRAY BIOPHARMA INC	75MG	N210496 002	Jun 27, 2018
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ENFUVRTIDE

INJECTABLE;SUBCUTANEOUS

FUZEON

+	ROCHE	90MG/VIAL	N021481 001	Mar 13, 2003
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ENOXAPARIN SODIUM

INJECTABLE;INTRAVENOUS, SUBCUTANEOUS

ENOXAPARIN SODIUM

AB	AMPHASTAR PHARMS	300MG/3ML (100MG/ML)	A208600 001	Mar 14, 2019
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AB	INC			
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AB	SANDOZ INC	300MG/3ML (100MG/ML)	A078660 001	Nov 28, 2011
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LOVENOX

AB	+	SANOFI AVENTIS US	300MG/3ML (100MG/ML)	N020164 009	Jan 23, 2003
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PRESCRIPTION DRUG PRODUCT LIST

ENOXAPARIN SODIUM

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>	NANJING KING-FRIEND	<u>30MG/0.3ML (100MG/ML)</u>	<u>A206834 001</u>	Nov 29, 2019
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A206834 002</u>	Nov 29, 2019
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A206834 003</u>	Nov 29, 2019
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A206834 004</u>	Nov 29, 2019
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A206834 005</u>	Nov 29, 2019
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A206834 006</u>	Nov 29, 2019
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A206834 007</u>	Nov 29, 2019
<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>	ZYDUS PHARMS	<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
<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
<u>ENTACAPONE</u>				
<u>AB</u>	AJANTA PHARMA LTD	<u>200MG</u>	<u>A205792 001</u>	Aug 31, 2017
<u>AB</u>	ALEMBIC PHARMS LTD	<u>200MG</u>	<u>A212601 001</u>	Jan 04, 2022
<u>AB</u>	ATLANTIDE	<u>200MG</u>	<u>A078941 001</u>	Aug 16, 2012
<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 PHARM	<u>200MG</u>	<u>A090690 001</u>	Jul 16, 2012
<u>AB</u>	SUNSHINE	<u>200MG</u>	<u>A206669 001</u>	Oct 03, 2018

ENTECAVIR

SOLUTION; ORAL

BARACLUDE

+	BRISTOL MYERS SQUIBB	0.05MG/ML	N021798 001	Mar 29, 2005
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TABLET; ORAL

BARACLUDE

<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
<u>ENTECAVIR</u>				
<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

PRESCRIPTION DRUG PRODUCT LIST

ENTECAVIR

TABLET; ORAL

ENTECAVIR

<u>AB</u>	AMNEAL PHARMS	<u>0.5MG</u>	<u>A206652</u>	<u>001</u>	Nov 12, 2015
<u>AB</u>		<u>1MG</u>	<u>A206652</u>	<u>002</u>	Nov 12, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.5MG</u>	<u>A206217</u>	<u>001</u>	Aug 26, 2015
<u>AB</u>		<u>1MG</u>	<u>A206217</u>	<u>002</u>	Aug 26, 2015
<u>AB</u>	BRECKENRIDGE	<u>0.5MG</u>	<u>A208721</u>	<u>001</u>	Mar 15, 2018
<u>AB</u>		<u>1MG</u>	<u>A208721</u>	<u>002</u>	Mar 15, 2018
<u>AB</u>	BRIGHTGENE	<u>0.5MG</u>	<u>A212126</u>	<u>001</u>	Sep 25, 2019
<u>AB</u>		<u>1MG</u>	<u>A212126</u>	<u>002</u>	Sep 25, 2019
<u>AB</u>	CASI PHARMS INC	<u>0.5MG</u>	<u>A206672</u>	<u>001</u>	May 11, 2017
<u>AB</u>		<u>1MG</u>	<u>A206672</u>	<u>002</u>	May 11, 2017
<u>AB</u>	CIPLA	<u>0.5MG</u>	<u>A206872</u>	<u>001</u>	Dec 06, 2016
<u>AB</u>		<u>1MG</u>	<u>A206872</u>	<u>002</u>	Dec 06, 2016
<u>AB</u>	HETERO LABS LTD V	<u>0.5MG</u>	<u>A205740</u>	<u>001</u>	Aug 21, 2015
<u>AB</u>		<u>1MG</u>	<u>A205740</u>	<u>002</u>	Aug 21, 2015
<u>AB</u>	PHARMADAX INC	<u>0.5MG</u>	<u>A212106</u>	<u>001</u>	Aug 10, 2020
<u>AB</u>		<u>1MG</u>	<u>A212106</u>	<u>002</u>	Aug 10, 2020
<u>AB</u>	PRINSTON INC	<u>0.5MG</u>	<u>A208782</u>	<u>001</u>	Oct 10, 2017
<u>AB</u>		<u>1MG</u>	<u>A208782</u>	<u>002</u>	Oct 10, 2017
<u>AB</u>	YUNG SHIN PHARM	<u>0.5MG</u>	<u>A208195</u>	<u>001</u>	Nov 10, 2021
<u>AB</u>		<u>1MG</u>	<u>A208195</u>	<u>002</u>	Nov 10, 2021
<u>AB</u>	ZYDUS PHARMS	<u>0.5MG</u>	<u>A206745</u>	<u>001</u>	Jun 23, 2017
<u>AB</u>		<u>1MG</u>	<u>A206745</u>	<u>002</u>	Jun 23, 2017

ENTRECTINIB

CAPSULE; ORAL

ROZLYTREK

+ GENENTECH INC

+!

100MG

200MG

N212725 001 Aug 15, 2019

N212725 002 Aug 15, 2019

ENZALUTAMIDE

CAPSULE; ORAL

ENZALUTAMIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>40MG</u>	<u>A209614</u>	<u>001</u>	May 14, 2021
<u>AB</u>	<u>XTANDI</u>				
<u>AB</u>	+! ASTELLAS	<u>40MG</u>	<u>N203415</u>	<u>001</u>	Aug 31, 2012
	TABLET; ORAL				
	XTANDI				
	+ ASTELLAS	40MG	N213674	001	Aug 04, 2020
	+!	80MG	N213674	002	Aug 04, 2020

EPHEDRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

REZIPRES

+! ETON

23.5MG/5ML (4.7MG/ML)

N213536 001 Jun 14, 2021

EPHEDRINE SULFATE

SOLUTION; INTRAVENOUS

AKOVAZ

<u>AP</u>	+! EXELA PHARMA	<u>50MG/ML (50MG/ML)</u>	<u>N208289</u>	<u>001</u>	Apr 29, 2016
	<u>CORPHEDRA</u>				
<u>AP</u>	PAR STERILE PRODUCTS	<u>50MG/ML (50MG/ML)</u>	<u>N208943</u>	<u>001</u>	Jan 27, 2017
	<u>EPHEDRINE SULFATE</u>				
<u>AP</u>	AKORN	<u>50MG/ML (50MG/ML)</u>	<u>N208609</u>	<u>001</u>	Mar 01, 2017
<u>AP</u>	AMNEAL	<u>50MG/ML (50MG/ML)</u>	<u>A212932</u>	<u>001</u>	Oct 23, 2019
<u>AP</u>	DR REDDYS LABS LTD	<u>50MG/ML (50MG/ML)</u>	<u>A212649</u>	<u>001</u>	Oct 03, 2020
<u>AP</u>	EUGIA PHARMA	<u>50MG/ML (50MG/ML)</u>	<u>A214579</u>	<u>001</u>	Jun 14, 2021
<u>AP</u>	FRESENIUS KABI USA	<u>50MG/ML (50MG/ML)</u>	<u>A209646</u>	<u>001</u>	Aug 04, 2020
<u>AP</u>	HIKMA	<u>50MG/ML (50MG/ML)</u>	<u>A214334</u>	<u>001</u>	Dec 15, 2020
<u>AP</u>	SANDOZ INC	<u>50MG/ML (50MG/ML)</u>	<u>A209784</u>	<u>001</u>	Aug 23, 2017
	AKOVAZ				
	+! EXELA PHARMA	25MG/5ML (5MG/ML)	N208289	002	Aug 02, 2021
	EMERPHED				
	+! NEXUS PHARMS	50MG/10ML (5MG/ML)	N213407	001	Apr 17, 2020

PRESCRIPTION DRUG PRODUCT LIST

EPINASTINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

ELESTAT

AT	+ !	ALLERGAN	0.05%	N021565	001	Oct 16, 2003
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EPINASTINE HYDROCHLORIDE

AT		AKORN	0.05%	A204055	001	May 05, 2017
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AT		APOTEX	0.05%	A090919	001	Oct 31, 2011
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AT		BRECKENRIDGE	0.05%	A090870	001	Mar 14, 2011
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AT		SOMERSET THERAPS LLC	0.05%	A090951	001	Oct 31, 2011
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EPINEPHRINE

INJECTABLE;INTRAMUSCULAR, SUBCUTANEOUS

EPINEPHRINE (AUTOINJECTOR)

AB		TEVA PHARMS USA	0.15MG/DELIVERY	A090589	002	Aug 16, 2018
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AB			0.3MG/DELIVERY	A090589	001	Aug 16, 2018
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EPIPEN

AB	+ !	MYLAN SPECIALITY LP	0.3MG/DELIVERY	N019430	001	Dec 22, 1987
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EPIPEN JR.

AB	+ !	MYLAN SPECIALITY LP	0.15MG/DELIVERY	N019430	002	Dec 22, 1987
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ADRENACLICK

BX	+ !	IMPAX	EQ 0.15MG/DELIVERY	N020800	003	Nov 25, 2009
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BX	+ !		EQ 0.3MG/DELIVERY	N020800	004	Nov 25, 2009
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SOLUTION;INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

ADRENALIN

AP	+ !	PAR STERILE PRODUCTS	EQ 30MG BASE/30ML (EQ 1MG BASE/ML)	N204640	001	Dec 18, 2013
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EPINEPHRINE

AP		INTL MEDICATION SYS	EQ 30MG BASE/30ML (EQ 1MG BASE/ML)	A211880	001	Apr 24, 2020
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ADRENALIN

+ !	PAR STERILE PRODUCTS	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	N204200	001	Dec 07, 2012
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SOLUTION;INTRAMUSCULAR, SUBCUTANEOUS

AUVI-Q

BX	+ !	KALEO INC	EQ 0.15MG/DELIVERY	N201739	002	Aug 10, 2012
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BX	+		EQ 0.3MG/DELIVERY	N201739	001	Aug 10, 2012
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+			EQ 0.1MG/DELIVERY	N201739	003	Nov 17, 2017
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SYMJEPI

+ !	ADAMIS PHARMS CORP	0.15MG/0.3ML (0.15MG/0.3ML)	N207534	002	Sep 27, 2018
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+ !		0.3MG/0.3ML (0.3MG/0.3ML)	N207534	001	Jun 15, 2017
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SOLUTION;INTRAVENOUS

EPINEPHRINE

+ !	HOSPIRA INC	1MG/10ML (0.1MG/ML)	N209359	001	Nov 05, 2019
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SOLUTION;INTRAVENOUS, INTRAOCULAR, INTRAMUSCULAR, SUBCUTANEOUS

EPINEPHRINE

+ !	BELCHER	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	N205029	001	Jul 29, 2014
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EPINEPHRINE BITARTRATE; LIDOCAINE HYDROCHLORIDE

INJECTABLE;INJECTION

LIGNOSPAN FORTE

!	DEPROCO	EQ 0.02MG BASE/ML;2%	A088389	001	Jan 22, 1985
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LIGNOSPAN STANDARD

!	DEPROCO	EQ 0.01MG BASE/ML;2%	A088390	001	Jan 22, 1985
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EPINEPHRINE BITARTRATE; PRILUCAINE HYDROCHLORIDE

INJECTABLE;INJECTION

CITANEST FORTE DENTAL

AP	+ !	DENTSPLY PHARM	0.005MG/ML;4%	N021383	001	
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PRILUCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE

AP		SEPTODONT INC	0.005MG/ML;4%	A078959	001	Aug 30, 2011
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EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE;INJECTION

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

AP		B BRAUN MEDICAL INC	0.005MG/ML;1.5%	A208475	001	Sep 08, 2021
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AP		HOSPIRA	0.005MG/ML;0.5%	A089635	001	Jun 21, 1988
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AP			0.005MG/ML;1.5%	A088571	001	Sep 13, 1985
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AP			0.005MG/ML;1.5%	A089645	001	Jun 21, 1988
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AP			0.005MG/ML;2%	A089651	001	Jun 21, 1988
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AP			0.01MG/ML;1%	A089644	001	Jun 21, 1988
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AP	!		0.01MG/ML;2%	A089646	001	Jun 21, 1988
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XYLOCAINE W/ EPINEPHRINE

AP	+ !	FRESENIUS KABI USA	0.005MG/ML;0.5%	N006488	012	
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AP	+ !		0.005MG/ML;1.5%	N006488	017	
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AP	+ !		0.005MG/ML;2%	N006488	019	Nov 13, 1986
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PRESCRIPTION DRUG PRODUCT LIST

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

XYLOCAINE W/ EPINEPHRINE

<u>AP</u>	<u>+</u> !		<u>0.01MG/ML;1%</u>	<u>N006488</u>	<u>004</u>	
<u>AP</u>	<u>+</u> !		<u>0.02MG/ML;2%</u>	<u>N006488</u>	<u>005</u>	
		<u>+</u> !	<u>0.005MG/ML;1%</u>	<u>N006488</u>	<u>018</u>	<u>Nov 13, 1986</u>

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ELLEENCE

<u>AP</u>	<u>+</u> !	<u>PFIZER INC</u>	<u>200MG/100ML (2MG/ML)</u>	<u>N050778</u>	<u>001</u>	<u>Sep 15, 1999</u>
<u>AP</u>	<u>+</u>		<u>50MG/25ML (2MG/ML)</u>	<u>N050778</u>	<u>002</u>	<u>Sep 15, 1999</u>

EPIRUBICIN HYDROCHLORIDE

<u>AP</u>		<u>ACTAVIS TOTOWA</u>	<u>50MG/25ML (2MG/ML)</u>	<u>A065445</u>	<u>002</u>	<u>Sep 18, 2008</u>
<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A065445</u>	<u>003</u>	<u>Sep 18, 2008</u>
<u>AP</u>		<u>AKORN</u>	<u>50MG/25ML (2MG/ML)</u>	<u>A090163</u>	<u>001</u>	<u>Jun 24, 2009</u>
<u>AP</u>		<u>CIPLA LTD</u>	<u>50MG/25ML (2MG/ML)</u>	<u>A065361</u>	<u>001</u>	<u>Oct 22, 2007</u>
<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A065361</u>	<u>002</u>	<u>Oct 22, 2007</u>
<u>AP</u>		<u>HIKMA</u>	<u>50MG/25ML (2MG/ML)</u>	<u>A065289</u>	<u>001</u>	<u>Jun 27, 2007</u>
<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A065289</u>	<u>002</u>	<u>Jun 27, 2007</u>
<u>AP</u>		<u>HISUN PHARM</u>	<u>50MG/25ML (2MG/ML)</u>	<u>A090075</u>	<u>001</u>	<u>Mar 25, 2010</u>
		<u>HANGZHOU</u>				
<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A090075</u>	<u>002</u>	<u>Mar 25, 2010</u>
<u>AP</u>		<u>IMPAX LABS INC</u>	<u>50MG/25ML (2MG/ML)</u>	<u>A065331</u>	<u>001</u>	<u>Aug 09, 2007</u>
<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A065331</u>	<u>002</u>	<u>Aug 09, 2007</u>

EPLERENONE

TABLET; ORAL

EPLERENONE

<u>AB</u>		<u>ACCORD HLTHCARE</u>	<u>25MG</u>	<u>A206922</u>	<u>001</u>	<u>Jul 13, 2017</u>
<u>AB</u>			<u>50MG</u>	<u>A206922</u>	<u>002</u>	<u>Jul 13, 2017</u>
<u>AB</u>		<u>BRECKENRIDGE</u>	<u>25MG</u>	<u>A208283</u>	<u>001</u>	<u>Sep 14, 2018</u>
<u>AB</u>			<u>50MG</u>	<u>A208283</u>	<u>002</u>	<u>Sep 14, 2018</u>
<u>AB</u>		<u>CHARTWELL RX</u>	<u>25MG</u>	<u>A078482</u>	<u>001</u>	<u>Jul 30, 2008</u>
<u>AB</u>			<u>50MG</u>	<u>A078482</u>	<u>002</u>	<u>Jul 30, 2008</u>
<u>AB</u>		<u>PRASCO</u>	<u>25MG</u>	<u>A203896</u>	<u>001</u>	<u>Feb 02, 2017</u>
<u>AB</u>			<u>50MG</u>	<u>A203896</u>	<u>002</u>	<u>Feb 02, 2017</u>
<u>AB</u>		<u>SANDOZ</u>	<u>25MG</u>	<u>A078510</u>	<u>001</u>	<u>Aug 01, 2008</u>
<u>AB</u>			<u>50MG</u>	<u>A078510</u>	<u>002</u>	<u>Aug 01, 2008</u>
<u>AB</u>		<u>WESTMINSTER PHARMS</u>	<u>25MG</u>	<u>A207842</u>	<u>001</u>	<u>Oct 25, 2021</u>
<u>AB</u>			<u>50MG</u>	<u>A207842</u>	<u>002</u>	<u>Oct 25, 2021</u>

INSPIRA

<u>AB</u>	<u>+</u>	<u>UPJOHN</u>	<u>25MG</u>	<u>N021437</u>	<u>001</u>	<u>Sep 27, 2002</u>
<u>AB</u>	<u>+</u> !		<u>50MG</u>	<u>N021437</u>	<u>002</u>	<u>Sep 27, 2002</u>

EPOPROSTENOL SODIUM

INJECTABLE; INJECTION

EPOPROSTENOL SODIUM

<u>AP1</u>		<u>TEVA PHARMS USA</u>	<u>EQ 0.5MG BASE/VIAL</u>	<u>A078396</u>	<u>001</u>	<u>Apr 23, 2008</u>
<u>AP1</u>			<u>EQ 1.5MG BASE/VIAL</u>	<u>A078396</u>	<u>002</u>	<u>Apr 23, 2008</u>

FLOLAN

<u>AP1</u>	<u>+</u> !	<u>GLAXOSMITHKLINE LLC</u>	<u>EQ 0.5MG BASE/VIAL</u>	<u>N020444</u>	<u>001</u>	<u>Sep 20, 1995</u>
<u>AP1</u>	<u>+</u> !		<u>EQ 1.5MG BASE/VIAL</u>	<u>N020444</u>	<u>002</u>	<u>Sep 20, 1995</u>

EPOPROSTENOL SODIUM

<u>AP2</u>		<u>SUN PHARM</u>	<u>EQ 0.5MG BASE/VIAL</u>	<u>A210473</u>	<u>001</u>	<u>Jan 15, 2021</u>
<u>AP2</u>			<u>EQ 1.5MG BASE/VIAL</u>	<u>A210473</u>	<u>002</u>	<u>Jan 15, 2021</u>

VELETRI

<u>AP2</u>	<u>+</u> !	<u>ACTELION</u>	<u>EQ 0.5MG BASE/VIAL</u>	<u>N022260</u>	<u>002</u>	<u>Jun 28, 2012</u>
<u>AP2</u>	<u>+</u> !		<u>EQ 1.5MG BASE/VIAL</u>	<u>N022260</u>	<u>001</u>	<u>Jun 27, 2008</u>

EPTIFIBATIDE

INJECTABLE; INJECTION

EPTIFIBATIDE

<u>AP</u>		<u>ACCORD HLTHCARE</u>	<u>2MG/ML</u>	<u>A205557</u>	<u>001</u>	<u>Nov 06, 2017</u>
<u>AP</u>			<u>75MG/100ML</u>	<u>A205557</u>	<u>002</u>	<u>Nov 06, 2017</u>
<u>AP</u>		<u>AKORN</u>	<u>2MG/ML</u>	<u>A204589</u>	<u>001</u>	<u>Apr 18, 2017</u>
<u>AP</u>			<u>75MG/100ML</u>	<u>A204589</u>	<u>002</u>	<u>Apr 18, 2017</u>
<u>AP</u>		<u>BAXTER HLTHCARE</u>	<u>2MG/ML</u>	<u>A208554</u>	<u>001</u>	<u>Nov 23, 2018</u>
		<u>CORP</u>				
<u>AP</u>			<u>75MG/100ML</u>	<u>A208554</u>	<u>002</u>	<u>Nov 23, 2018</u>
<u>AP</u>		<u>EUGIA PHARMA</u>	<u>2MG/ML</u>	<u>A206127</u>	<u>001</u>	<u>Dec 08, 2015</u>
<u>AP</u>			<u>75MG/100ML</u>	<u>A206127</u>	<u>002</u>	<u>Dec 08, 2015</u>
<u>AP</u>		<u>HAINAN POLY PHARM</u>	<u>2MG/ML</u>	<u>A209864</u>	<u>001</u>	<u>Jan 25, 2019</u>
<u>AP</u>			<u>75MG/100ML</u>	<u>A209864</u>	<u>002</u>	<u>Jan 25, 2019</u>
<u>AP</u>	<u>!</u>	<u>MYLAN LABS LTD</u>	<u>2MG/ML</u>	<u>A203258</u>	<u>001</u>	<u>Jul 20, 2018</u>

PRESCRIPTION DRUG PRODUCT LIST

EPTIFIBATIDE

INJECTABLE; INJECTION

EPTIFIBATIDE

<u>AP</u>	!		<u>75MG/100ML</u>	<u>A203258</u>	<u>002</u>	Jul 20, 2018
<u>AP</u>		SAGENT PHARMS INC	<u>2MG/ML</u>	<u>A204693</u>	<u>001</u>	Mar 07, 2018
<u>AP</u>			<u>75MG/100ML</u>	<u>A204693</u>	<u>002</u>	Mar 07, 2018
<u>AP</u>		SHUANGCHENG	<u>2MG/ML</u>	<u>A213081</u>	<u>001</u>	Sep 07, 2021
<u>AP</u>		TEVA PHARMS USA	<u>2MG/ML</u>	<u>A090854</u>	<u>001</u>	Jun 12, 2015

ERAVACYCLINE DIHYDROCHLORIDE

POWDER; INTRAVENOUS

XERAVA

+	!	TETRAPHASE PHARMS	EQ 50MG BASE/VIAL	N211109	001	Aug 27, 2018
+	!		EQ 100MG BASE/VIAL	N211109	002	Jun 03, 2020

ERDAFITINIB

TABLET; ORAL

BALVERSA

+		JANSSEN BIOTECH	3MG	N212018	001	Apr 12, 2019
+			4MG	N212018	002	Apr 12, 2019
+	!		5MG	N212018	003	Apr 12, 2019

ERGOALCIFEROL

CAPSULE; ORAL

DRISDOL

<u>AA</u>	+	!	VALIDUS PHARMS	<u>50,000 IU</u>	<u>N003444</u>	<u>001</u>
<u>ERGOALCIFEROL</u>						
<u>AA</u>			ALEMBIC LABS	<u>50,000 IU</u>	<u>A040833</u>	<u>001</u>
<u>AA</u>			PURACAP PHARM LLC	<u>50,000 IU</u>	<u>A204276</u>	<u>001</u>
<u>AA</u>			STRIDES PHARMA	<u>50,000 IU</u>	<u>A090455</u>	<u>001</u>
<u>AA</u>			SUN PHARM INDS INC	<u>50,000 IU</u>	<u>A040865</u>	<u>001</u>

VITAMIN D

<u>AA</u>			BIONPHARMA INC	<u>50,000 IU</u>	<u>A080704</u>	<u>001</u>
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ERGOLOID MESYLATES

TABLET; ORAL

ERGOLOID MESYLATES

!		SUN PHARM INDUSTRIES	1MG	A081113	001	Oct 31, 1991
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ERGOTAMINE TARTRATE

TABLET; SUBLINGUAL

ERGOMAR

!		TERSERA	2MG	A087693	001	Feb 24, 1983
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ERIBULIN MESYLATE

SOLUTION; INTRAVENOUS

HALAVEN

+	!	EISAI INC	1MG/2ML (0.5MG/ML)	N201532	001	Nov 15, 2010
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ERLOTINIB HYDROCHLORIDE

TABLET; ORAL

ERLOTINIB HYDROCHLORIDE

<u>AB</u>			ALEMBIC PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A214719</u>	<u>001</u>	Jul 08, 2021
<u>AB</u>				<u>EQ 100MG BASE</u>	<u>A214719</u>	<u>002</u>	Jul 08, 2021
<u>AB</u>				<u>EQ 150MG BASE</u>	<u>A214719</u>	<u>003</u>	Jul 08, 2021
<u>AB</u>			APOTEX	<u>EQ 25MG BASE</u>	<u>A208396</u>	<u>001</u>	Nov 05, 2019
<u>AB</u>				<u>EQ 100MG BASE</u>	<u>A208396</u>	<u>002</u>	Nov 05, 2019
<u>AB</u>				<u>EQ 150MG BASE</u>	<u>A208396</u>	<u>003</u>	Nov 05, 2019
<u>AB</u>			MSN	<u>EQ 25MG BASE</u>	<u>A214366</u>	<u>001</u>	May 10, 2021
<u>AB</u>				<u>EQ 100MG BASE</u>	<u>A214366</u>	<u>002</u>	May 10, 2021
<u>AB</u>				<u>EQ 150MG BASE</u>	<u>A214366</u>	<u>003</u>	May 10, 2021
<u>AB</u>			NATCO PHARMA LTD	<u>EQ 25MG BASE</u>	<u>A208488</u>	<u>001</u>	Nov 05, 2019
<u>AB</u>				<u>EQ 100MG BASE</u>	<u>A208488</u>	<u>002</u>	Nov 05, 2019
<u>AB</u>				<u>EQ 150MG BASE</u>	<u>A208488</u>	<u>003</u>	Nov 05, 2019
<u>AB</u>			RISING	<u>EQ 25MG BASE</u>	<u>A091002</u>	<u>001</u>	Jun 11, 2014
<u>AB</u>				<u>EQ 100MG BASE</u>	<u>A091002</u>	<u>002</u>	Jun 11, 2014
<u>AB</u>				<u>EQ 150MG BASE</u>	<u>A091002</u>	<u>003</u>	Jun 11, 2014
<u>AB</u>			SHILPA	<u>EQ 25MG BASE</u>	<u>A211960</u>	<u>001</u>	Nov 05, 2019
<u>AB</u>				<u>EQ 100MG BASE</u>	<u>A211960</u>	<u>002</u>	Nov 05, 2019
<u>AB</u>				<u>EQ 150MG BASE</u>	<u>A211960</u>	<u>003</u>	Nov 05, 2019
<u>AB</u>			SUN PHARM	<u>EQ 25MG BASE</u>	<u>A210300</u>	<u>001</u>	Nov 05, 2019
<u>AB</u>				<u>EQ 100MG BASE</u>	<u>A210300</u>	<u>002</u>	Nov 05, 2019
<u>AB</u>				<u>EQ 150MG BASE</u>	<u>A210300</u>	<u>003</u>	Nov 05, 2019
<u>AB</u>			TEVA PHARMS USA INC	<u>EQ 25MG BASE</u>	<u>A091059</u>	<u>001</u>	Nov 09, 2020
<u>AB</u>				<u>EQ 100MG BASE</u>	<u>A091059</u>	<u>002</u>	Aug 28, 2015

PRESCRIPTION DRUG PRODUCT LIST

ERLOTINIB HYDROCHLORIDE

TABLET; ORAL

ERLOTINIB HYDROCHLORIDE

AB		<u>EQ 150MG BASE</u>	<u>A091059 003</u>	Aug 28, 2015
AB	ZYDUS PHARMS	<u>EQ 25MG BASE</u>	<u>A213065 001</u>	Apr 16, 2020
AB		<u>EQ 100MG BASE</u>	<u>A213065 002</u>	Apr 16, 2020
AB		<u>EQ 150MG BASE</u>	<u>A213065 003</u>	Apr 16, 2020
<u>TARCEVA</u>				
AB	+ OSI PHARMS	<u>EQ 25MG BASE</u>	<u>N021743 001</u>	Nov 18, 2004
AB	+	<u>EQ 100MG BASE</u>	<u>N021743 002</u>	Nov 18, 2004
AB	+!	<u>EQ 150MG BASE</u>	<u>N021743 003</u>	Nov 18, 2004

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	EUGIA PHARMA	<u>EQ 1GM BASE/VIAL</u>	<u>A209133 001</u>	Jun 25, 2018
AP	GLAND PHARMA LTD	<u>EQ 1GM BASE/VIAL</u>	<u>A212040 001</u>	Mar 26, 2021
AP	SAVIOR LIFETEC CORP	<u>EQ 1GM BASE/VIAL</u>	<u>A207647 001</u>	Mar 19, 2019
<u>INVANZ</u>				
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>	
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ERYTHROMYCIN

AB	ARBOR PHARMS LLC	<u>250MG</u>	<u>A062746 001</u>	Dec 22, 1986
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GEL; TOPICAL

ERYGEL

AT	+! MYLAN	<u>2%</u>	<u>N050617 001</u>	Oct 21, 1987
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ERYTHROMYCIN

AT	FOUGERA PHARMS	<u>2%</u>	<u>A064184 001</u>	Sep 30, 1997
AT	PADAGIS US	<u>2%</u>	<u>A063211 001</u>	Jan 29, 1993
AT	TELIGENT	<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	! PADAGIS US	<u>0.5%</u>	<u>A062447 001</u>	Sep 26, 1983

SOLUTION; TOPICAL

ERYTHRA-DERM

AT	SAPTALIS PHARMS	<u>2%</u>	<u>A062687 001</u>	Feb 05, 1988
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ERYTHROMYCIN

AT	! PADAGIS US	<u>2%</u>	<u>A063038 001</u>	Jan 11, 1991
AT	TELIGENT	<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	! PADAGIS US	<u>2%</u>	<u>A064126 001</u>	Jul 03, 1996

TABLET; ORAL

ERYTHROMYCIN

AB	ARBOR PHARMS LLC	<u>250MG</u>	<u>A061621 001</u>	
AB	!	<u>500MG</u>	<u>A061621 002</u>	

PRESCRIPTION DRUG PRODUCT LIST

ERYTHROMYCIN

TABLET;ORAL

ERYTHROMYCIN

<u>AB</u>	CADILA PHARMS LTD	<u>250MG</u>	<u>A213628 001</u>	Jun 28, 2021
<u>AB</u>		<u>500MG</u>	<u>A213628 002</u>	Jun 28, 2021
<u>AB</u>	TEVA PHARMS USA INC	<u>250MG</u>	<u>A214549 001</u>	Feb 11, 2021
<u>AB</u>		<u>500MG</u>	<u>A214549 002</u>	Feb 11, 2021
<u>AB</u>	TORRENT	<u>250MG</u>	<u>A212015 001</u>	Jul 06, 2020
<u>AB</u>		<u>500MG</u>	<u>A212015 002</u>	Jul 06, 2020
BX	AMNEAL PHARMS CO	250MG	A209720 001	Mar 09, 2018
BX		500MG	A209720 002	Mar 09, 2018

TABLET, DELAYED RELEASE;ORAL

ERY-TAB

<u>AB</u>	ARBOR PHARMS LLC	<u>250MG</u>	<u>A062298 001</u>	
<u>AB</u>		<u>333MG</u>	<u>A062298 003</u>	Mar 29, 1982
<u>AB</u>	!	<u>500MG</u>	<u>A062298 002</u>	

ERYTHROMYCIN

<u>AB</u>	AMNEAL PHARMS CO	<u>250MG</u>	<u>A210954 001</u>	Jul 02, 2019
<u>AB</u>		<u>333MG</u>	<u>A210954 002</u>	Jul 02, 2019
<u>AB</u>		<u>500MG</u>	<u>A210954 003</u>	Jul 02, 2019
<u>AB</u>	TORRENT	<u>250MG</u>	<u>A211975 001</u>	Jul 26, 2021
<u>AB</u>		<u>500MG</u>	<u>A211975 002</u>	Jul 26, 2021

ERYTHROMYCIN ETHYLSUCCINATE

GRANULE;ORAL

E.E.S.

<u>AB</u>	+	ARBOR PHARMS LLC	<u>EQ 200MG BASE/5ML</u>	<u>N050207 001</u>
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ERYPED

<u>AB</u>	+	ARBOR PHARMS LLC	<u>EQ 200MG BASE/5ML</u>	<u>N050207 003</u>	Mar 30, 1987
<u>AB</u>	+	!	<u>EQ 400MG BASE/5ML</u>	<u>N050207 002</u>	

ERYTHROMYCIN ETHYLSUCCINATE

<u>AB</u>	AMNEAL PHARMS	<u>EQ 200MG BASE/5ML</u>	<u>A211204 001</u>	Nov 01, 2019
<u>AB</u>		<u>EQ 400MG BASE/5ML</u>	<u>A211204 002</u>	Nov 01, 2019
<u>AB</u>	ANI PHARMS	<u>EQ 200MG BASE/5ML</u>	<u>A062055 001</u>	
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A062055 003</u>	Nov 02, 2018
<u>AB</u>		<u>EQ 400MG BASE/5ML</u>	<u>A062055 002</u>	Nov 02, 2018

TABLET;ORAL

E.E.S. 400

BX	!	ARBOR PHARMS LLC	EQ 400MG BASE	A061905 002	Aug 12, 1982
BX	!	ARBOR PHARMS LLC	EQ 400MG BASE	A061904 001	

ERYTHROMYCIN LACTOBIONATE

INJECTABLE;INJECTION

ERYTHROCIN

<u>AP</u>		HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>A062638 001</u>	Oct 31, 1986
<u>AP</u>		+	<u>EQ 500MG BASE/VIAL</u>	<u>N050609 001</u>	Sep 24, 1986

ERYTHROMYCIN LACTOBIONATE

<u>AP</u>		EXELA PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A211086 001</u>	Sep 17, 2020
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ERYTHROMYCIN STEARATE

TABLET;ORAL

ERYTHROCIN STEARATE

	!	ARBOR PHARMS LLC	EQ 250MG BASE	A060359 001	
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ESCITALOPRAM OXALATE

SOLUTION;ORAL

ESCITALOPRAM OXALATE

<u>AA</u>		AMNEAL PHARMS	<u>EQ 5MG BASE/5ML</u>	<u>A202227 001</u>	Mar 14, 2012
<u>AA</u>		AUROBINDO PHARMA LTD	<u>EQ 5MG BASE/5ML</u>	<u>A079062 001</u>	Apr 02, 2012
<u>AA</u>	!	HETERO LABS LTD III	<u>EQ 5MG BASE/5ML</u>	<u>A202221 001</u>	Jun 12, 2012
<u>AA</u>		LANNETT CO INC	<u>EQ 5MG BASE/5ML</u>	<u>A090477 001</u>	Jun 12, 2013
<u>AA</u>		MACLEODS PHARMS LTD	<u>EQ 5MG BASE/5ML</u>	<u>A202754 001</u>	Mar 31, 2016
<u>AA</u>		TARO	<u>EQ 5MG BASE/5ML</u>	<u>A079121 001</u>	May 03, 2012

TABLET;ORAL

ESCITALOPRAM OXALATE

<u>AB</u>		ACCORD HLTHCARE	<u>EQ 5MG BASE</u>	<u>A202389 001</u>	Sep 11, 2012
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A202389 002</u>	Sep 11, 2012
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A202389 003</u>	Sep 11, 2012
<u>AB</u>		AMNEAL PHARMS	<u>EQ 5MG BASE</u>	<u>A205619 001</u>	May 17, 2017
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A205619 002</u>	May 17, 2017
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A205619 003</u>	May 17, 2017
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A090432 001</u>	Sep 11, 2012

PRESCRIPTION DRUG PRODUCT LIST

ESCITALOPRAM OXALATE

TABLET; ORAL

ESCITALOPRAM OXALATE

<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090432 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A090432 003</u>	Sep 11, 2012
<u>AB</u>	CADILA	<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>AB</u>	GRAVITI PHARMS	<u>EQ 5MG BASE</u>	<u>A078777 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078777 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078777 003</u>	Sep 11, 2012
<u>AB</u>	INVAGEN PHARMS	<u>EQ 5MG BASE</u>	<u>A078604 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078604 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078604 003</u>	Sep 11, 2012
<u>AB</u>	JUBILANT GENERICS	<u>EQ 5MG BASE</u>	<u>A202280 001</u>	Sep 12, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202280 002</u>	Sep 12, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A202280 003</u>	Sep 12, 2012
<u>AB</u>	LUPIN LTD	<u>EQ 5MG BASE</u>	<u>A078169 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078169 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078169 003</u>	Sep 11, 2012
<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

LEXAPRO

<u>AB</u>	+	ALLERGAN	<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

ESKETAMINE HYDROCHLORIDE

SPRAY; NASAL

SPRAVATO

+	JANSSEN PHARMS	EQ 28MG BASE	N211243 001	Mar 05, 2019
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ESLICARBAZEPINE ACETATE

TABLET; ORAL

APTIOM

<u>AB</u>	+	SUNOVION PHARMS INC	<u>200MG</u>	<u>N022416 001</u>	Nov 08, 2013
<u>AB</u>	+		<u>400MG</u>	<u>N022416 002</u>	Nov 08, 2013
<u>AB</u>	+		<u>600MG</u>	<u>N022416 003</u>	Nov 08, 2013
<u>AB</u>	+	!	<u>800MG</u>	<u>N022416 004</u>	Nov 08, 2013

ESLICARBAZEPINE ACETATE

<u>AB</u>	DR REDDYS LABS LTD	<u>200MG</u>	<u>A211238 001</u>	Jun 29, 2021
<u>AB</u>		<u>400MG</u>	<u>A211238 002</u>	Jun 29, 2021
<u>AB</u>		<u>600MG</u>	<u>A211238 003</u>	Jun 29, 2021
<u>AB</u>		<u>800MG</u>	<u>A211238 004</u>	Jun 29, 2021

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

BREVIBLOC

<u>AP</u>	+	BAXTER HLTHCARE	<u>10MG/ML</u>	<u>N019386 006</u>	Feb 25, 2003
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ESMOLOL HYDROCHLORIDE

<u>AP</u>	EUGIA PHARMA	<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>	GLAND PHARMA LTD	<u>10MG/ML</u>	<u>A208538 001</u>	Aug 14, 2019
<u>AP</u>	HIKMA	<u>10MG/ML</u>	<u>A076323 001</u>	Aug 10, 2004
<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 INC	<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

BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER

+	BAXTER HLTHCARE	2GM/100ML	N019386 005	Jan 27, 2003
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BREVIBLOC IN PLASTIC CONTAINER

+	BAXTER HLTHCARE	1GM/100ML	N019386 004	Feb 16, 2001
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PRESCRIPTION DRUG PRODUCT LIST

ESMOLOL HYDROCHLORIDE

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>	CISEN	<u>EQ 20MG BASE</u>	<u>A213158 001</u>	Sep 22, 2020
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A213158 002</u>	Sep 22, 2020
<u>AB</u>	CSPC OUYI	<u>EQ 20MG BASE</u>	<u>A212949 001</u>	Oct 02, 2020
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A212949 002</u>	Oct 02, 2020
<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>	ETHYPHARM	<u>EQ 20MG BASE</u>	<u>A090841 001</u>	Mar 31, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090841 002</u>	Mar 31, 2021
<u>AB</u>	GLENMARK PHARMS	<u>EQ 20MG BASE</u>	<u>A209495 001</u>	May 10, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A209495 002</u>	May 10, 2019
<u>AB</u>	GRAVITI PHARMS	<u>EQ 20MG BASE</u>	<u>A213486 001</u>	Mar 19, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A213486 002</u>	Mar 19, 2021
<u>AB</u>	HETERO LABS LTD III	<u>EQ 20MG BASE</u>	<u>A202784 001</u>	Sep 21, 2015
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A211977 001</u>	Jun 02, 2020
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A202784 002</u>	Sep 21, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A211977 002</u>	Jun 02, 2020
<u>AB</u>	INDCHEMIE HEALTH	<u>EQ 20MG BASE</u>	<u>A210559 001</u>	Feb 26, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A210559 002</u>	Feb 26, 2021
<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
<u>AB</u>	MYLAN	<u>EQ 20MG BASE</u>	<u>A078936 001</u>	Aug 02, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078936 002</u>	Aug 03, 2015
<u>AB</u>	SUN PHARM	<u>EQ 20MG BASE</u>	<u>A209735 001</u>	Apr 30, 2018
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A209735 002</u>	Apr 30, 2018
<u>AB</u>	TORRENT	<u>EQ 20MG BASE</u>	<u>A203636 001</u>	Oct 19, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A203636 002</u>	Oct 19, 2015
<u>AB</u>	ZYDUS PHARMS	<u>EQ 20MG BASE</u>	<u>A206296 001</u>	May 22, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A206296 002</u>	May 22, 2019

NEXIUM

<u>AB</u>	+ ASTRAZENECA	<u>EQ 20MG BASE</u>	<u>N021153 001</u>	Feb 20, 2001
<u>AB</u>	+!	<u>EQ 40MG BASE</u>	<u>N021153 002</u>	Feb 20, 2001

FOR SUSPENSION, DELAYED RELEASE;ORAL

ESOMEPRAZOLE MAGNESIUM

<u>AB</u>	CIPLA	<u>EQ 10MG BASE/PACKET</u>	<u>A211752 001</u>	Mar 23, 2020
<u>AB</u>		<u>EQ 20MG BASE/PACKET</u>	<u>A211751 001</u>	Mar 23, 2020
<u>AB</u>		<u>EQ 40MG BASE/PACKET</u>	<u>A211751 002</u>	Mar 23, 2020

NEXIUM

<u>AB</u>	+ ASTRAZENECA	<u>EQ 10MG BASE/PACKET</u>	<u>N022101 001</u>	Feb 27, 2008
<u>AB</u>	+	<u>EQ 20MG BASE/PACKET</u>	<u>N021957 001</u>	Oct 20, 2006
<u>AB</u>	+!	<u>EQ 40MG BASE/PACKET</u>	<u>N021957 002</u>	Oct 20, 2006
	+	EQ 2.5MG BASE/PACKET	N021957 003	Dec 15, 2011
	+	EQ 5MG BASE/PACKET	N021957 004	Dec 15, 2011

ESOMEPRAZOLE MAGNESIUM; NAPROXEN

TABLET, DELAYED RELEASE;ORAL

NAPROXEN AND ESOMEPRAZOLE MAGNESIUM

<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 20MG BASE;375MG</u>	<u>A204206 001</u>	Feb 18, 2020
<u>AB</u>		<u>EQ 20MG BASE;500MG</u>	<u>A204206 002</u>	Feb 18, 2020
<u>AB</u>	MYLAN	<u>EQ 20MG BASE;375MG</u>	<u>A204920 001</u>	Jul 20, 2021
<u>AB</u>		<u>EQ 20MG BASE;500MG</u>	<u>A204920 002</u>	Jul 20, 2021

VIMOVO

<u>AB</u>	+ HORIZON	<u>EQ 20MG BASE;375MG</u>	<u>N022511 002</u>	Apr 30, 2010
<u>AB</u>	+!	<u>EQ 20MG BASE;500MG</u>	<u>N022511 001</u>	Apr 30, 2010

PRESCRIPTION DRUG PRODUCT LIST

ESOMEPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS

ESOMEPRAZOLE SODIUM

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 40MG BASE/VIAL</u>	<u>A205379 001</u>	Sep 25, 2015
<u>AP</u>	DEVA HOLDING AS	<u>EQ 40MG BASE/VIAL</u>	<u>A207181 001</u>	Mar 06, 2017
<u>AP</u>	! EUGIA PHARMA	<u>EQ 40MG BASE/VIAL</u>	<u>A204657 002</u>	Aug 10, 2016
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 20MG BASE/VIAL</u>	<u>A203349 001</u>	Apr 01, 2020
<u>AP</u>		<u>EQ 40MG BASE/VIAL</u>	<u>A203349 002</u>	Apr 01, 2020
<u>AP</u>	MYLAN LABS LTD	<u>EQ 40MG BASE/VIAL</u>	<u>A202686 002</u>	May 17, 2017
<u>AP</u>	SUN PHARMA GLOBAL	<u>EQ 40MG BASE/VIAL</u>	<u>A200882 002</u>	Mar 18, 2013

NEXIUM IV

<u>AP</u>	+! ASTRAZENECA	<u>EQ 40MG BASE/VIAL</u>	<u>N021689 002</u>	Mar 31, 2005
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ESTAZOLAM

TABLET; ORAL

ESTAZOLAM

<u>AB</u>	MAYNE PHARMA	<u>1MG</u>	<u>A074921 001</u>	Jul 10, 1997
<u>AB</u>	!	<u>2MG</u>	<u>A074921 002</u>	Jul 10, 1997
<u>AB</u>	NOVITIUM PHARMA	<u>1MG</u>	<u>A074826 001</u>	Jul 03, 1997
<u>AB</u>		<u>2MG</u>	<u>A074826 002</u>	Jul 03, 1997
<u>AB</u>	WATSON LABS	<u>1MG</u>	<u>A074818 001</u>	Aug 19, 1997
<u>AB</u>		<u>2MG</u>	<u>A074818 002</u>	Aug 19, 1997

ESTRADIOL

CREAM; VAGINAL

ESTRACE

<u>AB</u>	+! ALLERGAN	<u>0.01%</u>	<u>A086069 001</u>	Jan 31, 1984
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ESTRADIOL

<u>AB</u>	ALVOGEN	<u>0.01%</u>	<u>A209767 001</u>	Mar 05, 2018
<u>AB</u>	MYLAN	<u>0.01%</u>	<u>A208788 001</u>	Dec 29, 2017
<u>AB</u>	PADAGIS ISRAEL	<u>0.01%</u>	<u>A210194 001</u>	Jan 22, 2018
<u>AB</u>	PRASCO LABS LLC	<u>0.01%</u>	<u>A212313 001</u>	Jul 15, 2021
<u>AB</u>	TEVA PHARMS USA	<u>0.01%</u>	<u>A210488 001</u>	Mar 30, 2018

FILM, EXTENDED RELEASE; TRANSDERMAL

CLIMARA

<u>AB</u>	+ BAYER HLTHCARE	<u>0.06MG/24HR</u>	<u>N020375 006</u>	May 27, 2003
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ESTRADIOL

<u>AB</u>	MYLAN TECHNOLOGIES	<u>0.06MG/24HR</u>	<u>A075182 005</u>	Jul 20, 2006
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CLIMARA

<u>AB2</u>	+ BAYER HLTHCARE	<u>0.025MG/24HR</u>	<u>N020375 004</u>	Mar 05, 1999
<u>AB2</u>	+	<u>0.0375MG/24HR</u>	<u>N020375 005</u>	May 27, 2003
<u>AB2</u>	+	<u>0.05MG/24HR</u>	<u>N020375 001</u>	Dec 22, 1994
<u>AB2</u>	+	<u>0.075MG/24HR</u>	<u>N020375 003</u>	Mar 23, 1998
<u>AB2</u>	+!	<u>0.1MG/24HR</u>	<u>N020375 002</u>	Dec 22, 1994

ESTRADIOL

<u>AB2</u>	MYLAN TECHNOLOGIES	<u>0.025MG/24HR</u>	<u>A075182 003</u>	Jan 26, 2005
<u>AB2</u>		<u>0.0375MG/24HR</u>	<u>A075182 004</u>	Jul 20, 2006
<u>AB2</u>		<u>0.05MG/24HR</u>	<u>A075182 006</u>	Feb 24, 2000
<u>AB2</u>		<u>0.075MG/24HR</u>	<u>A075182 002</u>	Jan 26, 2005
<u>AB2</u>		<u>0.1MG/24HR</u>	<u>A075182 001</u>	Feb 24, 2000
<u>AB3</u>	AMNEAL	<u>0.025MG/24HR</u>	<u>A211396 001</u>	Sep 28, 2020
<u>AB3</u>		<u>0.0375MG/24HR</u>	<u>A211396 002</u>	Sep 28, 2020
<u>AB3</u>		<u>0.05MG/24HR</u>	<u>A211396 003</u>	Sep 28, 2020
<u>AB3</u>		<u>0.075MG/24HR</u>	<u>A211396 004</u>	Sep 28, 2020
<u>AB3</u>		<u>0.1MG/24HR</u>	<u>A211396 005</u>	Sep 28, 2020
<u>AB3</u>	MYLAN TECHNOLOGIES	<u>0.025MG/24HR</u>	<u>A206685 001</u>	Aug 15, 2018
<u>AB3</u>		<u>0.0375MG/24HR</u>	<u>A206685 002</u>	Aug 15, 2018
<u>AB3</u>		<u>0.05MG/24HR</u>	<u>A206685 003</u>	Aug 15, 2018
<u>AB3</u>		<u>0.075MG/24HR</u>	<u>A206685 004</u>	Aug 15, 2018
<u>AB3</u>		<u>0.1MG/24HR</u>	<u>A206685 005</u>	Aug 15, 2018

MINIVELLE

<u>AB3</u>	+ NOVEN	<u>0.025MG/24HR</u>	<u>N203752 005</u>	Sep 23, 2014
<u>AB3</u>	+	<u>0.0375MG/24HR</u>	<u>N203752 001</u>	Oct 29, 2012
<u>AB3</u>	+	<u>0.05MG/24HR</u>	<u>N203752 003</u>	Oct 29, 2012
<u>AB3</u>	+	<u>0.075MG/24HR</u>	<u>N203752 002</u>	Oct 29, 2012
<u>AB3</u>	+!	<u>0.1MG/24HR</u>	<u>N203752 004</u>	Oct 29, 2012

GEL; TRANSDERMAL

DIVIGEL

+	! VERTICAL PHARMS LLC	<u>0.1%</u>	<u>N022038 001</u>	Jun 04, 2007
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GEL, METERED; TRANSDERMAL

ELESTRIN

+	! MYLAN SPECIALITY LP	<u>0.06% (0.87GM/ACTIVATION)</u>	<u>N021813 001</u>	Dec 15, 2006
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PRESCRIPTION DRUG PRODUCT LIST

ESTRADIOL

GEL, METERED; TRANSDERMAL

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

+! PFIZER 0.0075MG/24HR N020472 001 Apr 26, 1996

SPRAY; TRANSDERMAL

EVAMIST

+! PADAGIS US 1.53MG/SPRAY N022014 001 Jul 27, 2007

SYSTEM; TRANSDERMAL

ESTRADIOLAB1 MYLAN TECHNOLOGIES 0.025MG/24HR A201675 001 Dec 19, 2014AB1 0.0375MG/24HR A201675 002 Dec 19, 2014AB1 0.05MG/24HR A201675 003 Dec 19, 2014AB1 0.075MG/24HR A201675 004 Dec 19, 2014AB1 0.1MG/24HR A201675 005 Dec 19, 2014VIVELLE-DOTAB1 + NOVARTIS 0.025MG/24HR N020538 009 May 03, 2002AB1 + 0.0375MG/24HR N020538 005 Jan 08, 1999AB1 + 0.05MG/24HR N020538 006 Jan 08, 1999AB1 + 0.075MG/24HR N020538 007 Jan 08, 1999AB1 +! 0.1MG/24HR N020538 008 Jan 08, 1999

ESTRADIOL

BX AMNEAL 0.025MG/24HR A211293 001 Feb 04, 2019

BX 0.0375MG/24HR A211293 002 Feb 04, 2019

BX 0.05MG/24HR A211293 003 Feb 04, 2019

BX 0.075MG/24HR A211293 004 Feb 04, 2019

BX 0.1MG/24HR A211293 005 Feb 04, 2019

MENOSTAR

+! BAYER HLTHCARE 0.014MG/24HR N021674 001 Jun 08, 2004

TABLET; ORAL

ESTRADIOLAB BARR LABS INC 0.5MG A040197 001 Oct 22, 1997AB 1MG A040197 002 Oct 22, 1997AB ! 2MG A040197 003 Oct 22, 1997AB EPIC PHARMA INC 0.5MG A040275 001 Dec 29, 1998AB 1MG A040275 002 Dec 29, 1998AB 2MG A040275 003 Dec 29, 1998AB MAYNE PHARMA 0.5MG A040114 003 Mar 14, 1996AB 1MG A040114 001 Mar 14, 1996AB 2MG A040114 002 Mar 14, 1996

TABLET; VAGINAL

ESTRADIOLAB AMNEAL PHARMS 10MCG A205256 001 May 29, 2015AB GLENMARK PHARMS LTD 10MCG A210264 001 Sep 14, 2018AB TEVA PHARMS USA 10MCG A206388 001 Jul 21, 2017VAGIFEMAB +! NOVO NORDISK INC 10MCG N020908 002 Nov 25, 2009ESTRADIOL ACETATE

INSERT, 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

! PFIZER 5MG/ML A085470 003

ESTRADIOL VALERATE

INJECTABLE; INJECTION

DELESTROGENAO +! PAR STERILE 10MG/ML N009402 002
PRODUCTSAO +! 20MG/ML N009402 004AO +! 40MG/ML N009402 003ESTRADIOL VALERATEAO AM REGENT 20MG/ML A090920 001 Jan 19, 2010

PRESCRIPTION DRUG PRODUCT LIST

ESTRADIOL VALERATE

INJECTABLE; INJECTION

ESTRADIOL VALERATE

<u>AO</u>		<u>40MG/ML</u>	<u>A090920 002</u>	Jan 19, 2010
<u>AO</u>	HIKMA	<u>10MG/ML</u>	<u>A203723 001</u>	Apr 21, 2020
<u>AO</u>		<u>20MG/ML</u>	<u>A203723 002</u>	Apr 21, 2020
<u>AO</u>		<u>40MG/ML</u>	<u>A203723 003</u>	Apr 21, 2020

ESTRADIOL; LEVONORGESTREL

FILM, EXTENDED RELEASE; TRANSDERMAL

CLIMARA PRO

+	!	BAYER HLTHCARE	0.045MG/24HR; 0.015MG/24HR	N021258 001	Nov 21, 2003
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ESTRADIOL; NORETHINDRONE ACETATE

FILM, 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

<u>AB</u>	+	!	AMNEAL	<u>1MG; 0.5MG</u>	<u>N020907 001</u>	Nov 18, 1998
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ESTRADIOL AND NORETHINDRONE ACETATE

<u>AB</u>			BARR	<u>1MG; 0.5MG</u>	<u>A079193 001</u>	May 11, 2010
<u>AB</u>			BRECKENRIDGE PHARM	<u>0.5MG; 0.1MG</u>	<u>A078324 002</u>	Jun 09, 2011
<u>AB</u>				<u>1MG; 0.5MG</u>	<u>A078324 001</u>	Apr 17, 2008
<u>AB</u>			MYLAN LABS LTD	<u>0.5MG; 0.1MG</u>	<u>A207261 001</u>	Feb 10, 2017
<u>AB</u>				<u>1MG; 0.5MG</u>	<u>A207261 002</u>	Feb 10, 2017
<u>AB</u>			NOVAST LABS	<u>0.5MG; 0.1MG</u>	<u>A210612 001</u>	Apr 03, 2019
<u>AB</u>				<u>1MG; 0.5MG</u>	<u>A210612 002</u>	Apr 03, 2019
			AMABELZ			
BX			LUPIN LTD	0.5MG; 0.1MG	A203339 001	Jun 20, 2016
BX				1MG; 0.5MG	A203339 002	Jun 20, 2016

ESTRADIOL; NORETHINDRONE ACETATE; RELUGOLIX

TABLET; ORAL

MYFEMBREE

+	!	MYOVANT SCIENCES	1MG; 0.5MG; 40MG	N214846 001	May 26, 2021
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ESTRADIOL; NORGESTIMATE

TABLET; ORAL

ESTRADIOL AND NORGESTIMATE

!		BARR	1MG, 1MG; N/A, 0.09MG	A076812 001	Apr 29, 2005
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ESTRADIOL; PROGESTERONE

CAPSULE; ORAL

BIJUVA

+		THERAPEUTICSMD INC	0.5MG; 100MG	N210132 002	Dec 28, 2021
+	!		1MG; 100MG	N210132 001	Oct 28, 2018

ESTRAMUSTINE PHOSPHATE SODIUM

CAPSULE; ORAL

EMCYT

+	!	PHARMACIA AND UPJOHN	EQ 140MG PHOSPHATE	N018045 001	
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ESTROGENS, CONJUGATED

CREAM; TOPICAL, VAGINAL

PREMARIN

+	!	WYETH PHARMS	0.625MG/GM	N020216 001	
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INJECTABLE; INJECTION

PREMARIN

+	!	WYETH PHARMS	25MG/VIAL	N010402 001	
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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
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PREMPRO

+	!	WYETH PHARMS	0.3MG; 1.5MG	N020527 005	Jun 04, 2003
+	!		0.45MG; 1.5MG	N020527 004	Mar 12, 2003

PRESCRIPTION DRUG PRODUCT LIST

ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE

TABLET; ORAL-28

PREMPRO

+	!	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

OGEN 5

+		PFIZER	6MG	A083220	004
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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	ORBION PHARMS	1MG	A091113	001	Jun 10, 2014
AB		2MG	A091113	002	Jun 10, 2014
AB		3MG	A091113	003	Jun 10, 2014
AB	SUN PHARM	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

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

EDECRIIN

AP	+	!	BAUSCH	EQ 50MG BASE/VIAL	N016093	001
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ETHACRYNATE SODIUM

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

PRESCRIPTION DRUG PRODUCT LIST

ETHACRYNATE SODIUM

INJECTABLE; INJECTION

ETHACRYNATE SODIUM

AP	VPI PHARMS INC	EO 50MG BASE/VIAL	A208663 001	Jun 09, 2020
AP	ZYDUS PHARMS	EO 50MG BASE/VIAL	A207758 001	Nov 17, 2017

ETHACRYNIC ACID

TABLET; ORAL

EDECRIN

AB	+ BAUSCH	25MG	N016092 001	
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ETHACRYNIC ACID

AB	AGNITIO	25MG	A211809 001	Jul 12, 2019
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	HIKMA	25MG	A207262 001	Feb 23, 2017
AB	LUPIN LTD	25MG	A211719 001	Sep 06, 2019
AB	PAR PHARM INC	25MG	A208501 001	Jul 21, 2017
AB	SCIEGEN PHARMS INC	25MG	A211232 001	Aug 27, 2019
AB	STRIDES PHARMA	25MG	A213240 001	Oct 19, 2020
AB	UPSHER SMITH LABS	25MG	A212417 001	Feb 19, 2020

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

MYAMBUTOL

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
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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
AB	KELNOR			
AB	BARR	0.035MG;1MG	A076785 001	May 23, 2005
AB	LO-MALMOREDE			
AB	NOVAST LABS	0.035MG;1MG	A209548 001	Feb 11, 2019
AB	MALMOREDE			
AB	NOVAST LABS	0.05MG;1MG	A209547 001	Jul 25, 2018
AB	ZOVIA 1/35E-28			
AB	MAYNE PHARMA	0.035MG;1MG	A072721 001	Dec 30, 1991
AB	ZOVIA 1/50E-28			
AB	! WATSON LABS	0.05MG;1MG	A072723 001	Dec 30, 1991

ETHINYL ESTRADIOL; ETONOGESTREL

RING; VAGINAL

ELURYNG

AB	AMNEAL	0.015MG/24HR;0.12MG/24HR	A210830 001	Dec 11, 2019
AB	ETHINYL ESTRADIOL; ETONOGESTREL			
AB	TEVA PHARMS USA INC	0.015MG/24HR;0.12MG/24HR	A204305 001	Jan 13, 2021
AB	NUVARING			
AB	+ ORGANON SUB MERCK	0.015MG/24HR;0.12MG/24HR	N021187 001	Oct 03, 2001
AB	VERARING			
AB	DR REDDYS LABS SA	0.015MG/24HR;0.12MG/24HR	A207577 001	Dec 09, 2021

ETHINYL ESTRADIOL; LEVONORGESTREL

SYSTEM; TRANSDERMAL

TWIRLA

+	AGILE	0.03MG/24HR;0.12MG/24HR	N204017 001	Feb 14, 2020
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TABLET; ORAL

ASHLYNA

AB	GLENMARK GENERICS	0.03MG,0.01MG;0.15MG,N/A	A203163 001	Feb 23, 2015
AB	DAYSEE			
AB	LUPIN LTD	0.03MG,0.01MG;0.15MG,N/A	A091467 001	Apr 10, 2013

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL

<u>DOLISHALE</u>					
AB	NOVAST LABS	<u>0.02MG;0.09MG</u>	<u>A091692</u>	<u>001</u>	Oct 22, 2020
<u>ICLEVIA</u>					
AB	AUROBINDO PHARMA LTD	<u>0.03MG;0.15MG</u>	<u>A206850</u>	<u>001</u>	Jun 29, 2018
<u>INTROVALE</u>					
AB	XIROMED	<u>0.03MG;0.15MG</u>	<u>A079064</u>	<u>001</u>	Sep 27, 2010
<u>JAIMIESS</u>					
AB	XIROMED	<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>A203770</u>	<u>001</u>	Dec 27, 2017
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL</u>					
AB	AMNEAL PHARMS	<u>0.03MG;0.15MG</u>	<u>A203871</u>	<u>001</u>	Nov 13, 2015
AB		<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>A203872</u>	<u>001</u>	Dec 22, 2015
AB	GLENMARK GENERICS	<u>0.02MG;0.09MG</u>	<u>A202791</u>	<u>001</u>	Apr 09, 2015
AB	GLENMARK PHARMS LTD	<u>0.03MG;0.15MG</u>	<u>A203164</u>	<u>001</u>	Jun 12, 2015
AB	LUPIN LTD	<u>0.03MG;0.15MG</u>	<u>A091440</u>	<u>001</u>	Oct 23, 2012
AB	MYLAN LABS LTD	<u>0.03MG;0.15MG</u>	<u>A200490</u>	<u>001</u>	Apr 21, 2015
AB	! WATSON LABS	<u>0.02MG;0.09MG</u>	<u>A079218</u>	<u>001</u>	Jun 06, 2011
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL</u>					
AB	LUPIN LTD	<u>0.02MG,0.1MG;0.01MG,N/A</u>	<u>A091674</u>	<u>001</u>	Oct 26, 2011
AB	MAYNE PHARMA	<u>0.02MG,0.1MG;0.01MG,N/A</u>	<u>A200407</u>	<u>001</u>	Oct 25, 2011
AB		<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>A078834</u>	<u>001</u>	May 31, 2011
AB	MYLAN LABS LTD	<u>0.02MG,0.15MG;</u>	<u>A206053</u>	<u>001</u>	Oct 02, 2017
		<u>0.025MG,0.15MG;0.03MG,0.15MG;</u>			
AB		<u>0.02MG,0.1MG;0.01MG,N/A</u>	<u>A200493</u>	<u>001</u>	Jun 17, 2015
AB		<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>A200492</u>	<u>001</u>	May 27, 2015
AB	XIROMED	<u>0.02MG,0.1MG;0.01MG,N/A</u>	<u>A205131</u>	<u>001</u>	Dec 14, 2017
<u>LO SIMPESSÉ</u>					
AB	AUROBINDO PHARMA LTD	<u>0.02MG,0.1MG;0.01MG,N/A</u>	<u>A206852</u>	<u>001</u>	Apr 28, 2017
<u>LOSEASONIQUE</u>					
AB	TEVA BRANDED PHARM	<u>0.02MG,0.1MG;0.01MG,N/A</u>	<u>N022262</u>	<u>001</u>	Oct 24, 2008
<u>QUARTETTE</u>					
AB	+! TEVA BRANDED PHARM	<u>0.02MG,0.15MG;0.025MG,0.15MG;0.03MG,0.15MG;0.01MG,N/A</u>	<u>N204061</u>	<u>001</u>	Mar 28, 2013
<u>QUASENSE</u>					
AB	WATSON LABS	<u>0.03MG;0.15MG</u>	<u>A077101</u>	<u>001</u>	Sep 06, 2006
<u>SEASONALE</u>					
AB	+! TEVA BRANDED PHARM	<u>0.03MG;0.15MG</u>	<u>N021544</u>	<u>001</u>	Sep 05, 2003
<u>SEASONIQUE</u>					
AB	+! TEVA BRANDED PHARM	<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>N021840</u>	<u>001</u>	May 25, 2006
<u>SETLAKIN</u>					
AB	NOVAST LABS	<u>0.03MG;0.15MG</u>	<u>A090716</u>	<u>001</u>	Sep 15, 2014
<u>SIMPESSÉ</u>					
AB	AUROBINDO PHARMA LTD	<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>A206851</u>	<u>001</u>	Apr 07, 2017
	BALCOLTRA				
	+! AVION PHARMS	0.02MG;0.1MG	N208612	001	Jan 09, 2018
	TYBLUME				
	+! EXELTIS USA INC	0.02MG;0.1MG	N209405	001	Mar 30, 2020
TABLET; ORAL-28					
<u>ALTAVERA</u>					
AB	XIROMED	<u>0.03MG;0.15MG</u>	<u>A079102</u>	<u>001</u>	Aug 03, 2010
<u>AYUNA</u>					
AB	AUROBINDO PHARMA LTD	<u>0.03MG;0.15MG</u>	<u>A206866</u>	<u>001</u>	Sep 23, 2016
<u>ENPRESSE-28</u>					
AB	DURAMED PHARMS BARR	<u>0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG</u>	<u>A075809</u>	<u>002</u>	Jul 16, 2001
<u>KURVELO</u>					
AB	LUPIN LTD	<u>0.03MG;0.15MG</u>	<u>A091408</u>	<u>001</u>	Oct 17, 2012
<u>LEVONEST</u>					
AB	NOVAST LABS LTD	<u>0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG</u>	<u>A090719</u>	<u>001</u>	Dec 29, 2010
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL</u>					
AB	AMNEAL PHARMS	<u>0.03MG;0.15MG</u>	<u>A201095</u>	<u>001</u>	Dec 08, 2014
AB	MYLAN LABS LTD	<u>0.03MG;0.15MG</u>	<u>A091663</u>	<u>001</u>	Dec 21, 2012
AB	NAARI PTE LTD	<u>0.02MG;0.1MG</u>	<u>A207065</u>	<u>001</u>	Aug 17, 2020
AB		<u>0.03MG;0.15MG</u>	<u>A207033</u>	<u>001</u>	Oct 09, 2020
<u>LEVORA 0.15/30-28</u>					
AB	! MAYNE PHARMA	<u>0.03MG;0.15MG</u>	<u>A073594</u>	<u>001</u>	Dec 13, 1993

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL-28

<u>MARLISSA</u>					
AB	GLENMARK GENERICS	0.03MG;0.15MG	A091452 001	Feb 29, 2012	
<u>MYZILRA</u>					
AB	VINTAGE PHARMS LLC	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A077502 001	Nov 23, 2011	
<u>PORTIA-28</u>					
AB	BARR	0.03MG;0.15MG	A075866 002	May 23, 2002	
<u>TRIVORA-28</u>					
AB	MAYNE PHARMA	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A074538 002	Dec 18, 1997	
<u>AFIRMELLE</u>					
AB1	AUROBINDO PHARMA LTD	0.02MG;0.1MG	A206886 001	Nov 14, 2016	
<u>AVIANE-28</u>					
AB1	DURAMED PHARMS BARR	0.02MG;0.1MG	A075796 001	Apr 30, 2001	
<u>FALMINA</u>					
AB1	NOVAST LABS LTD	0.02MG;0.1MG	A090721 001	Mar 28, 2012	
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL</u>					
AB1	AMNEAL PHARMS	0.02MG;0.1MG	A201108 001	Feb 05, 2014	
AB1	MAYNE PHARMA	0.02MG;0.1MG	A076625 001	Nov 18, 2004	
AB1	MYLAN LABS LTD	0.02MG;0.1MG	A200245 001	Oct 09, 2013	
<u>VIENVA</u>					
AB1	XIROMED	0.02MG;0.1MG	A201088 001	May 21, 2015	
<u>LESSINA-28</u>					
AB2	BARR	0.02MG;0.1MG	A075803 002	Mar 20, 2002	
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL</u>					
AB2	MAYNE PHARMA	0.02MG;0.1MG	A077681 001	May 31, 2006	
BX	LUPIN LTD	0.02MG;0.1MG	A091425 001	Jan 18, 2013	
BX		0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A200248 001	Nov 19, 2015	

ETHINYL ESTRADIOL; NORELGESTROMIN

FILM, EXTENDED RELEASE; TRANSDERMAL

<u>ONSURA</u>					
AB	TEVA PHARMS USA	0.035MG/24HR;0.15MG/24HR	A213977 001	Aug 25, 2021	
<u>XULANE</u>					
AB	MYLAN TECHNOLOGIES	0.035MG/24HR;0.15MG/24HR	A200910 001	Apr 16, 2014	
<u>ETHINYL ESTRADIOL AND NORELGESTROMIN</u>					
BX	AMNEAL	0.035MG/24HR;0.15MG/24HR	A213950 001	Feb 25, 2021	

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21

<u>NORTREL 1/35-21</u>					
AB	BARR	0.035MG;1MG	A072693 001	Feb 28, 1992	
<u>NORTREL 7/7/7</u>					
	BARR	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A075478 001	Aug 30, 2002	

TABLET; ORAL-28

<u>ALYACEN 1/35</u>					
AB	GLENMARK GENERICS	0.035MG;1MG	A091634 001	Jan 19, 2012	
<u>ALYACEN 7/7/7</u>					
AB	GLENMARK GENERICS	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A091636 001	Jan 19, 2012	
<u>ARANELLE</u>					
AB	BARR	0.035MG,0.035MG,0.035MG;0.5MG,1MG,0.5MG	A076783 001	Sep 29, 2004	
<u>BALZIVA-28</u>					
AB	BARR	0.035MG;0.4MG	A076238 001	Apr 22, 2004	
<u>BRIELLYN</u>					
AB	GLENMARK GENERICS	0.035MG;0.4MG	A090538 001	Mar 22, 2011	
<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	MAYNE PHARMA	0.035MG;0.5MG	A070686 001	Jan 29, 1987	
AB	NAARI PTE LTD	0.035MG;1MG	A206864 001	Apr 28, 2017	
AB	WATSON LABS	0.035MG;0.4MG	A078323 001	Feb 04, 2010	

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-28

NORETHINDRONE AND ETHINYL ESTRADIOL

AB	WATSON LABS TEVA	0.035MG;1MG	A070687	001	Jan 29, 1987
	<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>PHILITH</u>				
AB	NOVAST LABS LTD	0.035MG;0.4MG	A090947	001	Dec 22, 2011
	<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
	PIRMELLA 1/35				
BX	LUPIN LTD	0.035MG;1MG	A201512	001	Apr 24, 2013
	PIRMELLA 7/7/7				
BX	LUPIN LTD	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A201510	001	Apr 24, 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

KAITLIB FE

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	AMNEAL PHARMS	0.035MG;0.4MG	A078892	001	Sep 26, 2011
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
AB	NAARI PTE LTD	0.035MG;0.4MG	A207066	001	Mar 29, 2017

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

CAPSULE; ORAL

GEMMILY

AB	XIROMED	0.02MG;1MG	A213317	001	Nov 09, 2020
	<u>MERZEE</u>				
AB	SLAYBACK PHARMA LLC	0.02MG;1MG	A212706	001	Dec 18, 2020
	<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>				
AB	AMNEAL PHARMS	0.02MG;1MG	A214292	001	Jul 20, 2021
	<u>TAYTULLA</u>				
AB	+! 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
	<u>BLISOVI 24 FE</u>				
AB	LUPIN LTD	0.02MG;1MG	A091398	001	Oct 28, 2015
	<u>FYAVOLV</u>				
AB	LUPIN LTD	0.005MG;1MG	A204213	002	Dec 10, 2015
AB		0.0025MG;0.5MG	A204213	001	Dec 10, 2015
	<u>GILDESS 24 FE</u>				
AB	VINTAGE PHARMS	0.02MG;1MG	A090293	001	Dec 01, 2014
	<u>LARIN 24 FE</u>				
AB	NOVAST LABS	0.02MG;1MG	A202994	001	Feb 18, 2015
	<u>LERIBANE</u>				
AB	NOVAST LABS	0.0025MG;0.5MG	A203435	002	Jun 03, 2016
AB		0.005MG;1MG	A203435	001	Jun 03, 2016

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

AB	!	BARR LABS INC	<u>0.005MG;1MG</u>	<u>A076221</u>	<u>001</u>	Nov 06, 2009
AB		GLENMARK GENERICS	<u>0.0025MG;0.5MG</u>	<u>A203038</u>	<u>001</u>	Apr 02, 2015
AB			<u>0.005MG;1MG</u>	<u>A203038</u>	<u>002</u>	Apr 02, 2015

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

AB	!	AMNEAL PHARMS	<u>0.02MG;1MG</u>	<u>A078267</u>	<u>001</u>	Sep 01, 2009
AB		BARR LABS INC	<u>0.02MG;1MG</u>	<u>A090938</u>	<u>001</u>	Dec 01, 2014
AB		GLENMARK PHARMS LTD	<u>0.02MG;1MG</u>	<u>A204847</u>	<u>001</u>	Nov 17, 2017

LO LOESTRIN FE

+!

TABLET; ORAL-21

AUROVELA 1.5/30

AB		AUROBINDO PHARMA LTD	<u>0.03MG;1.5MG</u>	<u>A207581</u>	<u>001</u>	Jun 26, 2017
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AUROVELA 1/20

AB		AUROBINDO PHARMA LTD	<u>0.02MG;1MG</u>	<u>A207506</u>	<u>001</u>	Jun 16, 2017
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HAILEY 1.5/30

AB		GLENMARK PHARMS	<u>0.03MG;1.5MG</u>	<u>A209297</u>	<u>001</u>	Jun 05, 2018
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JUNEL 1.5/30

AB		BARR	<u>0.03MG;1.5MG</u>	<u>A076381</u>	<u>001</u>	May 30, 2003
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JUNEL 1/20

AB		BARR	<u>0.02MG;1MG</u>	<u>A076380</u>	<u>001</u>	May 30, 2003
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LARIN 1.5/30

AB		NOVAST LABS	<u>0.03MG;1.5MG</u>	<u>A202996</u>	<u>001</u>	Mar 20, 2014
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LARIN 1/20

AB		NOVAST LABS	<u>0.02MG;1MG</u>	<u>A202995</u>	<u>001</u>	Dec 04, 2013
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LOESTRIN 21 1.5/30

AB	+	TEVA BRANDED PHARM	<u>0.03MG;1.5MG</u>	<u>N017875</u>	<u>001</u>	
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LOESTRIN 21 1/20

AB	+	TEVA BRANDED PHARM	<u>0.02MG;1MG</u>	<u>N017876</u>	<u>001</u>	
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LOESTRIN FE 1.5/30

AB	+!	TEVA BRANDED PHARM	<u>0.03MG;1.5MG</u>	<u>N017355</u>	<u>001</u>	
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MICROGESTIN 1.5/30

AB		MAYNE PHARMA	<u>0.03MG;1.5MG</u>	<u>A075548</u>	<u>002</u>	Jul 30, 2003
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MICROGESTIN 1/20

AB		MAYNE PHARMA	<u>0.02MG;1MG</u>	<u>A075647</u>	<u>002</u>	Jul 30, 2003
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NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

AB		GLENMARK PHARMS LTD	<u>0.02MG;1MG</u>	<u>A206969</u>	<u>001</u>	Jan 20, 2016
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AB		MYLAN LABS LTD	<u>0.02MG;1MG</u>	<u>A202771</u>	<u>001</u>	Nov 06, 2013
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AB			<u>0.03MG;1.5MG</u>	<u>A202770</u>	<u>001</u>	Feb 19, 2015
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TRI-LEGEST 21

BARR

TABLET; ORAL-28

AUROVELA FE 1.5/30

AB		AUROBINDO PHARMA LTD	<u>0.03MG;1.5MG</u>	<u>A207580</u>	<u>001</u>	Jun 15, 2017
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AUROVELA FE 1/20

AB		AUROBINDO PHARMA LTD	<u>0.02MG;1MG</u>	<u>A207505</u>	<u>001</u>	Jun 16, 2017
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BLISOVI FE 1/20

AB		LUPIN LTD	<u>0.02MG;1MG</u>	<u>A201584</u>	<u>001</u>	Nov 18, 2015
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CHABELINA FE

AB	!	NOVAST LABS	<u>0.02MG,0.03MG,0.035MG;1MG,1MG,1MG</u>	<u>A202962</u>	<u>001</u>	Apr 15, 2020
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HAILEY FE 1.5/30

AB		GLENMARK PHARMS	<u>0.03MG;1.5MG</u>	<u>A209031</u>	<u>001</u>	Jun 05, 2018
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HAILEY FE 1/20

AB		GLENMARK PHARMS LTD	<u>0.02MG;1MG</u>	<u>A206597</u>	<u>001</u>	Nov 21, 2017
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JUNEL FE 1.5/30

AB		BARR	<u>0.03MG;1.5MG</u>	<u>A076064</u>	<u>001</u>	Sep 18, 2003
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JUNEL FE 1/20

AB		BARR	<u>0.02MG;1MG</u>	<u>A076081</u>	<u>001</u>	Sep 18, 2003
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LARIN FE 1.5/30

AB		NOVAST LABS	<u>0.03MG;1.5MG</u>	<u>A091453</u>	<u>001</u>	Aug 23, 2013
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LARIN FE 1/20

AB		NOVAST LABS	<u>0.02MG;1MG</u>	<u>A091454</u>	<u>001</u>	Aug 26, 2013
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LOESTRIN FE 1/20

AB	+	TEVA BRANDED PHARM	<u>0.02MG;1MG</u>	<u>N017354</u>	<u>001</u>	
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MICROGESTIN FE 1.5/30

AB		MAYNE PHARMA	<u>0.03MG;1.5MG</u>	<u>A075548</u>	<u>001</u>	Feb 05, 2001
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PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL-28

MICROGESTIN FE 1/20

AB	MAYNE PHARMA	<u>0.02MG;1MG</u>	<u>A075647</u>	<u>001</u>	Feb 05, 2001
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NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

AB	MAYNE PHARMA	<u>0.02MG,0.03MG,0.035MG;1MG,1MG,1MG</u>	<u>A076629</u>	<u>001</u>	Mar 18, 2010
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AB	MYLAN LABS LTD	<u>0.02MG;1MG</u>	<u>A202772</u>	<u>001</u>	Nov 14, 2013
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AB		<u>0.03MG;1.5MG</u>	<u>A202741</u>	<u>001</u>	Feb 20, 2015
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TRI-LEGEST FE

AB	BARR	<u>0.02MG,0.03MG,0.035MG;1MG,1MG,1MG</u>	<u>A076105</u>	<u>001</u>	Oct 26, 2007
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BLISOVI FE 1.5/30

EX	LUPIN LTD	0.03MG;1.5MG	A201585	001	Nov 18, 2015
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TABLET, CHEWABLE; ORAL

MIBELAS 24 FE

AB	LUPIN	<u>0.02MG;1MG</u>	<u>A206287</u>	<u>001</u>	May 24, 2016
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MINASTRIN 24 FE

AB	+! APIL	<u>0.02MG;1MG</u>	<u>N203667</u>	<u>001</u>	May 08, 2013
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NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

AB	AMNEAL PHARMS	<u>0.02MG;1MG</u>	<u>A207514</u>	<u>001</u>	Sep 11, 2017
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AB	GLENMARK PHARMS LTD	<u>0.02MG;1MG</u>	<u>A210369</u>	<u>001</u>	Dec 26, 2017
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AB	XIROMED	<u>0.02MG;1MG</u>	<u>A209609</u>	<u>001</u>	Jul 16, 2018
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ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

ESTARYLLA

AB	XIROMED	<u>0.035MG;0.25MG</u>	<u>A090794</u>	<u>001</u>	Jan 30, 2013
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MILI

AB	AUROBINDO PHARMA LTD	<u>0.035MG;0.25MG</u>	<u>A205449</u>	<u>001</u>	Jul 07, 2016
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MONO-LINYAH

AB	NOVAST LABS LTD	<u>0.035MG;0.25MG</u>	<u>A090523</u>	<u>001</u>	May 23, 2012
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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
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AB		<u>0.035MG;0.25MG</u>	<u>A203865</u>	<u>001</u>	Oct 27, 2015
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AB		<u>0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG</u>	<u>A203873</u>	<u>001</u>	May 12, 2016
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AB	!	<u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u>	<u>A200494</u>	<u>001</u>	Jun 17, 2011
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AB	GLENMARK GENERICS	<u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u>	<u>A200538</u>	<u>001</u>	Apr 05, 2012
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AB	!	<u>0.035MG;0.25MG</u>	<u>A204057</u>	<u>001</u>	Feb 23, 2016
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AB	GLENMARK PHARMS	<u>0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG</u>	<u>A205588</u>	<u>001</u>	Apr 26, 2016
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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
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AB		<u>0.035MG;0.25MG</u>	<u>A205630</u>	<u>001</u>	Oct 27, 2016
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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
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AB		<u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u>	<u>A200383</u>	<u>001</u>	Apr 07, 2015
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AB	NAARI PTE LTD	<u>0.035MG;0.035MG;0.035MG;0.18MG;0.215MG;0.25MG</u>	<u>A200383</u>	<u>001</u>	Apr 07, 2015
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AB		<u>0.035MG;0.25MG</u>	<u>A200384</u>	<u>001</u>	Apr 07, 2015
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PREVIFEM

AB	VINTAGE PHARMS LLC	<u>0.035MG;0.25MG</u>	<u>A076334</u>	<u>001</u>	Jan 09, 2004
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SPRINTEC

AB	!	<u>0.035MG;0.25MG</u>	<u>A075804</u>	<u>001</u>	Sep 25, 2002
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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
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TRI-ESTARYLLA

AB	XIROMED	<u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u>	<u>A090793</u>	<u>001</u>	Jan 30, 2013
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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
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TRI-LO-ESTARYLLA

AB	XIROMED	<u>0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG</u>	<u>A091232</u>	<u>001</u>	Jun 29, 2015
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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
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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
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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
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PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-21

CRYSELLE**AB** DURAMED PHARMS BARR **0.03MG;0.3MG** **A075840 001** Nov 30, 2001

TABLET; ORAL-28

CRYSELLE**AB** DURAMED PHARMS BARR **0.03MG;0.3MG** **A075840 002** Nov 30, 2001ELINEST**AB** NOVAST LABS LTD **0.03MG;0.3MG** **A091105 001** Mar 28, 2012LOW-OGESTREL-28**AB** MAYNE PHARMA **0.03MG;0.3MG** **A075288 002** Jul 28, 1999ETHINYL ESTRADIOL; SEGESTERONE ACETATE

RING; VAGINAL

ANNOVERA

+! THERAPEUTICSMID INC 0.013MG/24HR;0.15MG/24HR N209627 001 Aug 10, 2018ETHIODIZED OIL

OIL; INTRALYMPHATIC, INTRAUTERINE

LIPIODOL

+! GUERBET EQ 4.8GM IODINE/10ML (EQ 480MG IODINE/ML) N009190 001ETHIONAMIDE

TABLET; ORAL

TRECATOR

+! WYETH PHARMS 250MG N013026 002ETHOSUXIMIDE

CAPSULE; ORAL

ETHOSUXIMIDE**AB** AKORN **250MG** **A040686 001** May 28, 2008**AB** BIONPHARMA INC **250MG** **A040430 001** Oct 28, 2002**AB** STRIDES PHARMA **250MG** **A211928 001** Feb 19, 2019ZARONTIN**AB** **+**! PARKE DAVIS **250MG** **N012380 001**

ETHOSUXIMIDE

BX HERITAGE PHARMS INC 250MG A200892 001 Sep 25, 2012

BX PURACAP PHARM LLC 250MG A210654 001 Mar 16, 2020

SYRUP; ORAL

ETHOSUXIMIDE**AA** MIKART **250MG/5ML** **A040506 001** Dec 22, 2003**AA** PHARM ASSOC **250MG/5ML** **A040253 001** Nov 22, 2000ZARONTIN**AA** **+**! PARKE-DAVIS **250MG/5ML** **A080258 001**ETODOLAC

CAPSULE; ORAL

ETODOLAC**AB** ANI PHARMS **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** BAYSHORE PHARMS LLC **400MG** **A210704 001** Dec 16, 2020**AB** **500MG** **A210704 002** Dec 16, 2020**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

TABLET, EXTENDED RELEASE; ORAL

ETODOLAC**AB** BAYSHORE PHARMS LLC **400MG** **A212263 001** Nov 24, 2020**AB** **500MG** **A212263 002** Nov 24, 2020**AB** **600MG** **A212263 003** Nov 24, 2020**AB** TARO **400MG** **A076174 001** Mar 13, 2003

PRESCRIPTION DRUG PRODUCT LIST

ETODOLAC

TABLET, EXTENDED RELEASE;ORAL

ETODOLAC

<u>AB</u>		<u>500MG</u>	<u>A076174 002</u>	Mar 13, 2003
<u>AB</u>		<u>600MG</u>	<u>A076174 003</u>	Mar 13, 2003
<u>AB</u>	TEVA	<u>400MG</u>	<u>A075665 003</u>	Feb 05, 2001
<u>AB</u>		<u>500MG</u>	<u>A075665 002</u>	Jul 31, 2000
<u>AB</u>	!	<u>600MG</u>	<u>A075665 001</u>	Jul 31, 2000
<u>AB</u>	ZYDUS PHARMS	<u>400MG</u>	<u>A091134 001</u>	Jan 23, 2014
<u>AB</u>		<u>500MG</u>	<u>A091134 002</u>	Jan 23, 2014
<u>AB</u>		<u>600MG</u>	<u>A091134 003</u>	Jan 23, 2014

ETOMIDATE

INJECTABLE;INJECTION

AMIDATE

<u>AP</u>	+!	HOSPIRA	<u>2MG/ML</u>	<u>N018227 001</u>	Sep 07, 1982
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ETOMIDATE

<u>AP</u>		CAPLIN	<u>2MG/ML</u>	<u>A215028 001</u>	Dec 18, 2020
<u>AP</u>		EMCURE PHARMS LTD	<u>2MG/ML</u>	<u>A204618 001</u>	Aug 13, 2014
<u>AP</u>		EUGIA PHARMA	<u>2MG/ML</u>	<u>A206126 001</u>	Feb 24, 2017
<u>AP</u>		GLAND PHARMA LTD	<u>2MG/ML</u>	<u>A209058 001</u>	Apr 18, 2017
<u>AP</u>		HIKMA	<u>2MG/ML</u>	<u>A074593 001</u>	Nov 04, 1996
<u>AP</u>			<u>2MG/ML</u>	<u>A202354 001</u>	Feb 25, 2016
<u>AP</u>		MYLAN LABS LTD	<u>2MG/ML</u>	<u>A201044 001</u>	Feb 07, 2017
<u>AP</u>		ZYDUS PHARMS	<u>2MG/ML</u>	<u>A202360 001</u>	Jul 18, 2014

ETONOGESTREL

IMPLANT;IMPLANTATION

NEXPLANON

+!	ORGANON USA INC	68MG/IMPLANT	<u>N021529 002</u>	May 31, 2011
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ETOPOSID

CAPSULE;ORAL

ETOPOSID

!	MYLAN	50MG	<u>A075635 001</u>	Sep 19, 2001
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INJECTABLE;INJECTION

ETOPOSID

<u>AP</u>		ACCORD HLTHCARE	<u>20MG/ML</u>	<u>A074513 001</u>	Mar 14, 1996
<u>AP</u>	!	FRESENIUS KABI USA	<u>20MG/ML</u>	<u>A074983 001</u>	Sep 30, 1998
<u>AP</u>		HIKMA	<u>20MG/ML</u>	<u>A074290 001</u>	Jul 17, 1995
<u>AP</u>		TEVA PHARMS USA	<u>20MG/ML</u>	<u>A074529 001</u>	Jul 24, 1996

ETOPOSID PHOSPHATE

INJECTABLE;INJECTION

ETOPOPHOS PRESERVATIVE FREE

+!	CHEPLAPHARM	EQ 100MG BASE/VIAL	<u>N020457 001</u>	May 17, 1996
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ETRAVIRINE

TABLET;ORAL

ETRAVIRINE

<u>AB</u>		AMNEAL	<u>100MG</u>	<u>A214196 002</u>	Jun 14, 2021
<u>AB</u>			<u>200MG</u>	<u>A214196 003</u>	Jun 14, 2021

INTELENCE

<u>AB</u>	+	JANSSEN R AND D	<u>100MG</u>	<u>N022187 001</u>	Jan 18, 2008
<u>AB</u>	+!		<u>200MG</u>	<u>N022187 002</u>	Dec 22, 2010
	+		25MG	<u>N022187 003</u>	Mar 26, 2012

EVEROLIMUS

TABLET;ORAL

AFINITOR

<u>AB</u>	+	NOVARTIS	<u>2.5MG</u>	<u>N022334 003</u>	Jul 09, 2010
<u>AB</u>	+!		<u>5MG</u>	<u>N022334 001</u>	Mar 30, 2009
<u>AB</u>	+		<u>7.5MG</u>	<u>N022334 004</u>	Mar 30, 2012
<u>AB</u>	+		<u>10MG</u>	<u>N022334 002</u>	Mar 30, 2009

EVEROLIMUS

<u>AB</u>		ALKEM LABS LTD	<u>0.25MG</u>	<u>A214138 001</u>	Nov 26, 2021
<u>AB</u>			<u>0.5MG</u>	<u>A214138 002</u>	Nov 26, 2021
<u>AB</u>			<u>0.75MG</u>	<u>A214138 003</u>	Nov 26, 2021
<u>AB</u>			<u>1MG</u>	<u>A214138 004</u>	Nov 26, 2021
<u>AB</u>		BIOCON PHARMA	<u>2.5MG</u>	<u>A214182 001</u>	Feb 11, 2021
<u>AB</u>			<u>5MG</u>	<u>A214182 002</u>	Feb 11, 2021
<u>AB</u>			<u>7.5MG</u>	<u>A214182 003</u>	Feb 11, 2021
<u>AB</u>			<u>10MG</u>	<u>A214182 004</u>	Feb 11, 2021
<u>AB</u>		BRECKENRIDGE	<u>0.25MG</u>	<u>A205432 001</u>	May 20, 2021
<u>AB</u>			<u>0.5MG</u>	<u>A205432 002</u>	May 20, 2021

PRESCRIPTION DRUG PRODUCT LIST

EVEROLIMUS

TABLET;ORAL

EVEROLIMUS

<u>AB</u>		<u>0.75MG</u>	<u>A205432</u>	<u>003</u>	May 20, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A205426</u>	<u>001</u>	Mar 05, 2021
<u>AB</u>		<u>5MG</u>	<u>A205426</u>	<u>002</u>	Mar 05, 2021
<u>AB</u>		<u>7.5MG</u>	<u>A205426</u>	<u>003</u>	Mar 05, 2021
<u>AB</u>		<u>10MG</u>	<u>A205426</u>	<u>004</u>	Mar 05, 2021
<u>AB</u>	HIKMA	<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
<u>AB</u>		<u>1MG</u>	<u>A206133</u>	<u>004</u>	Nov 18, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A207486</u>	<u>001</u>	Jun 08, 2020
<u>AB</u>		<u>5MG</u>	<u>A207486</u>	<u>002</u>	Jun 08, 2020
<u>AB</u>		<u>7.5MG</u>	<u>A207486</u>	<u>003</u>	Jun 08, 2020
<u>AB</u>		<u>10MG</u>	<u>A207486</u>	<u>004</u>	Nov 23, 2021
<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>A212936</u>	<u>001</u>	Jun 08, 2020
<u>AB</u>		<u>5MG</u>	<u>A212936</u>	<u>002</u>	Jun 08, 2020
<u>AB</u>		<u>7.5MG</u>	<u>A212936</u>	<u>003</u>	Jun 08, 2020
<u>AB</u>		<u>10MG</u>	<u>A212936</u>	<u>004</u>	Jun 08, 2020
<u>AB</u>	PAR PHARM	<u>0.25MG</u>	<u>A205775</u>	<u>001</u>	Oct 18, 2021
<u>AB</u>		<u>0.5MG</u>	<u>A205775</u>	<u>002</u>	Oct 18, 2021
<u>AB</u>		<u>0.75MG</u>	<u>A205775</u>	<u>003</u>	Oct 18, 2021
<u>AB</u>		<u>1MG</u>	<u>A205775</u>	<u>004</u>	Oct 18, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A207934</u>	<u>001</u>	Dec 09, 2019
<u>AB</u>		<u>5MG</u>	<u>A207934</u>	<u>002</u>	Dec 09, 2019
<u>AB</u>		<u>7.5MG</u>	<u>A207934</u>	<u>003</u>	Dec 09, 2019
<u>AB</u>		<u>10MG</u>	<u>A207934</u>	<u>004</u>	Dec 09, 2020
<u>AB</u>	TEVA PHARMS USA	<u>2.5MG</u>	<u>A210050</u>	<u>001</u>	Dec 09, 2019
<u>AB</u>		<u>5MG</u>	<u>A210050</u>	<u>002</u>	Dec 09, 2019
<u>AB</u>		<u>7.5MG</u>	<u>A210050</u>	<u>003</u>	Dec 09, 2019

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
<u>AB</u>	+	!	<u>1MG</u>	<u>N021560</u>	<u>004</u>	Aug 10, 2018

TABLET, FOR SUSPENSION;ORAL

AFINITOR DISPERZ

<u>AB</u>	+	NOVARTIS PHARM	<u>2MG</u>	<u>N203985</u>	<u>001</u>	Aug 29, 2012
<u>AB</u>	+		<u>3MG</u>	<u>N203985</u>	<u>002</u>	Aug 29, 2012
<u>AB</u>	+	!	<u>5MG</u>	<u>N203985</u>	<u>003</u>	Aug 29, 2012

EVEROLIMUS

<u>AB</u>		MYLAN	<u>2MG</u>	<u>A210130</u>	<u>001</u>	Apr 19, 2019
<u>AB</u>			<u>3MG</u>	<u>A210130</u>	<u>002</u>	Apr 19, 2019
<u>AB</u>			<u>5MG</u>	<u>A210130</u>	<u>003</u>	Apr 19, 2019

EXEMESTANE

TABLET;ORAL

AROMASIN

<u>AB</u>	+	!	PFIZER	<u>25MG</u>	<u>N020753</u>	<u>001</u>	Oct 21, 1999
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EXEMESTANE

<u>AB</u>		ALVOGEN	<u>25MG</u>	<u>A200898</u>	<u>001</u>	Jul 28, 2014
<u>AB</u>		BRECKENRIDGE	<u>25MG</u>	<u>A211031</u>	<u>001</u>	Feb 21, 2019
<u>AB</u>		CIPLA	<u>25MG</u>	<u>A210323</u>	<u>001</u>	Apr 27, 2018
<u>AB</u>		HIKMA	<u>25MG</u>	<u>A077431</u>	<u>001</u>	Apr 01, 2011
<u>AB</u>		QILU	<u>25MG</u>	<u>A213547</u>	<u>001</u>	Apr 13, 2020
<u>AB</u>		RISING	<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>		ZYDUS PHARMS	<u>25MG</u>	<u>A202602</u>	<u>001</u>	Oct 03, 2018

EXENATIDE SYNTHETIC

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

SUSPENSION, EXTENDED RELEASE;SUBCUTANEOUS

BYDUREON BCISE

+	!	ASTRAZENECA AB	2MG/0.85ML (2MG/0.85ML)	N209210	001	Oct 20, 2017
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PRESCRIPTION DRUG PRODUCT LIST

EZETIMIBE

TABLET;ORAL

EZETIMIBE

<u>AB</u>	ACCORD HLTHCARE	<u>10MG</u>	<u>A211550 001</u>	Oct 26, 2018
<u>AB</u>	ALKEM LABS LTD	<u>10MG</u>	<u>A209234 001</u>	Dec 21, 2017
<u>AB</u>	AMNEAL PHARMS CO	<u>10MG</u>	<u>A208803 001</u>	Jun 12, 2017
<u>AB</u>	APOTEX	<u>10MG</u>	<u>A208332 001</u>	Jun 12, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A209838 001</u>	Aug 25, 2017
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MG</u>	<u>A078560 001</u>	Jun 26, 2015
<u>AB</u>	OHM LABS INC	<u>10MG</u>	<u>A207311 001</u>	Jun 12, 2017
<u>AB</u>	SANDOZ INC	<u>10MG</u>	<u>A203931 001</u>	Jun 12, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>10MG</u>	<u>A210673 001</u>	Oct 23, 2020
<u>AB</u>	WATSON LABS INC	<u>10MG</u>	<u>A200831 001</u>	Jun 12, 2017
<u>AB</u>	ZYDUS PHARMS	<u>10MG</u>	<u>A204331 001</u>	Jun 12, 2017

ZETIA

<u>AB</u>	+!	ORGANON	<u>10MG</u>	<u>N021445 001</u>	Oct 25, 2002
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EZETIMIBE; ROSUVASTATIN CALCIUM

TABLET;ORAL

ROSZET

+	ALThERA PHARMS	10MG;EQ 5MG BASE	N213072 001	Mar 23, 2021
+		10MG;EQ 10MG BASE	N213072 002	Mar 23, 2021
+		10MG;EQ 20MG BASE	N213072 003	Mar 23, 2021
+!		10MG;EQ 40MG BASE	N213072 004	Mar 23, 2021

EZETIMIBE; SIMVASTATIN

TABLET;ORAL

EZETIMIBE AND SIMVASTATIN

<u>AB</u>	ALKEM LABS LTD	<u>10MG;10MG</u>	<u>A209222 001</u>	Dec 22, 2017
<u>AB</u>		<u>10MG;20MG</u>	<u>A209222 002</u>	Dec 22, 2017
<u>AB</u>		<u>10MG;40MG</u>	<u>A209222 003</u>	Dec 22, 2017
<u>AB</u>		<u>10MG;80MG</u>	<u>A209222 004</u>	Dec 22, 2017
<u>AB</u>	AMNEAL PHARMS CO	<u>10MG;10MG</u>	<u>A208831 001</u>	Nov 21, 2017
<u>AB</u>		<u>10MG;20MG</u>	<u>A208831 002</u>	Nov 21, 2017
<u>AB</u>		<u>10MG;40MG</u>	<u>A208831 003</u>	Nov 21, 2017
<u>AB</u>		<u>10MG;80MG</u>	<u>A208831 004</u>	Nov 21, 2017
<u>AB</u>	DR REDDYS LABS SA	<u>10MG;10MG</u>	<u>A200909 001</u>	Apr 26, 2017
<u>AB</u>		<u>10MG;20MG</u>	<u>A200909 002</u>	Apr 26, 2017
<u>AB</u>		<u>10MG;40MG</u>	<u>A200909 003</u>	Apr 26, 2017
<u>AB</u>		<u>10MG;80MG</u>	<u>A200909 004</u>	Apr 26, 2017
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MG;10MG</u>	<u>A208699 001</u>	Jun 27, 2019
<u>AB</u>		<u>10MG;20MG</u>	<u>A208699 002</u>	Jun 27, 2019
<u>AB</u>		<u>10MG;40MG</u>	<u>A208699 003</u>	Jun 27, 2019
<u>AB</u>		<u>10MG;80MG</u>	<u>A208699 004</u>	Jun 27, 2019
<u>AB</u>	WATSON LABS INC	<u>10MG;10MG</u>	<u>A202968 001</u>	Apr 26, 2017
<u>AB</u>		<u>10MG;20MG</u>	<u>A202968 002</u>	Apr 26, 2017
<u>AB</u>		<u>10MG;40MG</u>	<u>A202968 003</u>	Apr 26, 2017
<u>AB</u>		<u>10MG;80MG</u>	<u>A202968 004</u>	Apr 26, 2017

VYTORIN

<u>AB</u>	+	ORGANON	<u>10MG;10MG</u>	<u>N021687 001</u>	Jul 23, 2004
<u>AB</u>	+		<u>10MG;20MG</u>	<u>N021687 002</u>	Jul 23, 2004
<u>AB</u>	+		<u>10MG;40MG</u>	<u>N021687 003</u>	Jul 23, 2004
<u>AB</u>	+!		<u>10MG;80MG</u>	<u>N021687 004</u>	Jul 23, 2004

FAMCICLOVIR

TABLET;ORAL

FAMCICLOVIR

<u>AB</u>	APOTEX	<u>125MG</u>	<u>A091480 001</u>	Jul 22, 2011
<u>AB</u>		<u>250MG</u>	<u>A091480 002</u>	Jul 22, 2011
<u>AB</u>		<u>500MG</u>	<u>A091480 003</u>	Jul 22, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>125MG</u>	<u>A091114 001</u>	Mar 21, 2011
<u>AB</u>		<u>250MG</u>	<u>A091114 002</u>	Mar 21, 2011
<u>AB</u>		<u>500MG</u>	<u>A091114 003</u>	Mar 21, 2011
<u>AB</u>	CIPLA	<u>125MG</u>	<u>A078278 001</u>	Mar 21, 2011
<u>AB</u>		<u>250MG</u>	<u>A078278 002</u>	Mar 21, 2011
<u>AB</u>		<u>500MG</u>	<u>A078278 003</u>	Mar 21, 2011
<u>AB</u>	HETERO LABS LTD V	<u>125MG</u>	<u>A202438 001</u>	Sep 10, 2014
<u>AB</u>		<u>250MG</u>	<u>A202438 002</u>	Sep 10, 2014
<u>AB</u>		<u>500MG</u>	<u>A202438 003</u>	Sep 10, 2014
<u>AB</u>	MACLEODS PHARMS LTD	<u>125MG</u>	<u>A201022 001</u>	Jan 12, 2012
<u>AB</u>		<u>250MG</u>	<u>A201022 002</u>	Jan 12, 2012
<u>AB</u>		<u>500MG</u>	<u>A201022 003</u>	Jan 12, 2012

PRESCRIPTION DRUG PRODUCT LIST

FAMCICLOVIR

TABLET; ORAL

FAMCICLOVIR

<u>AB</u>	TEVA PHARMS	<u>125MG</u>	<u>A077487</u>	<u>001</u>	Aug 24, 2007
<u>AB</u>		<u>250MG</u>	<u>A077487</u>	<u>002</u>	Aug 24, 2007
<u>AB</u>	!	<u>500MG</u>	<u>A077487</u>	<u>003</u>	Aug 24, 2007

FAMOTIDINE

FOR SUSPENSION; ORAL

FAMOTIDINE

<u>AB</u>	AKORN	<u>40MG/5ML</u>	<u>A201995</u>	<u>001</u>	May 30, 2014
<u>AB</u>	!	<u>40MG/5ML</u>	<u>A090440</u>	<u>001</u>	Jun 29, 2010
<u>AB</u>	NAVINTA LLC	<u>40MG/5ML</u>	<u>A091020</u>	<u>001</u>	May 27, 2010
<u>AB</u>	NOVEL LABS INC	<u>40MG/5ML</u>	<u>A201695</u>	<u>001</u>	Dec 17, 2012
<u>AB</u>	NOVITIUM PHARMA	<u>40MG/5ML</u>	<u>A215043</u>	<u>001</u>	Apr 20, 2021

INJECTABLE; INJECTION

FAMOTIDINE

<u>AP</u>	ATHENEX INC	<u>10MG/ML</u>	<u>A075651</u>	<u>001</u>	Apr 16, 2001
<u>AP</u>		<u>10MG/ML</u>	<u>A075684</u>	<u>001</u>	Apr 16, 2001
<u>AP</u>	FRESENIUS KABI USA	<u>10MG/ML</u>	<u>A075709</u>	<u>001</u>	Apr 16, 2001
<u>AP</u>	!	<u>10MG/ML</u>	<u>A075488</u>	<u>001</u>	Apr 16, 2001
<u>AP</u>	MYLAN LABS LTD	<u>10MG/ML</u>	<u>A078641</u>	<u>001</u>	Jun 25, 2008

FAMOTIDINE PRESERVATIVE FREE

<u>AP</u>	ATHENEX INC	<u>10MG/ML</u>	<u>A075622</u>	<u>001</u>	Apr 16, 2001
<u>AP</u>		<u>10MG/ML</u>	<u>A075825</u>	<u>001</u>	Apr 17, 2001
<u>AP</u>	FRESENIUS KABI USA	<u>10MG/ML</u>	<u>A075813</u>	<u>001</u>	Apr 16, 2001
<u>AP</u>	!	<u>10MG/ML</u>	<u>A075486</u>	<u>001</u>	Apr 16, 2001
<u>AP</u>	MYLAN LABS LTD	<u>10MG/ML</u>	<u>A078642</u>	<u>001</u>	Jun 25, 2008

FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER

!	BAXTER HLTHCARE	0.4MG/ML	A075591	001	May 10, 2001
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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>	ANNORA PHARMA	<u>20MG</u>	<u>A215767</u>	<u>001</u>	Nov 04, 2021
<u>AB</u>		<u>40MG</u>	<u>A215767</u>	<u>002</u>	Nov 04, 2021
<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>	ASCENT PHARMS INC	<u>20MG</u>	<u>A215689</u>	<u>001</u>	Oct 15, 2021
<u>AB</u>		<u>40MG</u>	<u>A215689</u>	<u>002</u>	Oct 15, 2021
<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>	CELLTRION	<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>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>	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>	VKT PHARMA	<u>20MG</u>	<u>A215630</u>	<u>001</u>	Jan 07, 2022
<u>AB</u>		<u>40MG</u>	<u>A215630</u>	<u>002</u>	Jan 07, 2022

FAMOTIDINE; IBUPROFEN

TABLET; ORAL

DUEXIS

<u>AB</u>	+!	HORIZON	<u>26.6MG; 800MG</u>	<u>N022519</u>	<u>001</u>	Apr 23, 2011
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IBUPROFEN AND FAMOTIDINE

<u>AB</u>	ALKEM LABS LTD	<u>26.6MG; 800MG</u>	<u>A211890</u>	<u>001</u>	Aug 03, 2021
<u>AB</u>	TEVA PHARMS USA	<u>26.6MG; 800MG</u>	<u>A211278</u>	<u>001</u>	Oct 29, 2021

FEBUXOSTAT

TABLET; ORAL

FEBUXOSTAT

<u>AB</u>	ALEMBIC PHARMS LTD	<u>40MG</u>	<u>A205421</u>	<u>001</u>	Jul 01, 2019
<u>AB</u>		<u>80MG</u>	<u>A205421</u>	<u>002</u>	Jul 01, 2019
<u>AB</u>	ALKEM LABS LTD	<u>40MG</u>	<u>A212924</u>	<u>001</u>	Dec 07, 2021
<u>AB</u>		<u>80MG</u>	<u>A212924</u>	<u>002</u>	Dec 07, 2021
<u>AB</u>	DR REDDYS LABS LTD	<u>40MG</u>	<u>A205374</u>	<u>001</u>	Oct 22, 2020
<u>AB</u>		<u>80MG</u>	<u>A205374</u>	<u>002</u>	Oct 22, 2020

PRESCRIPTION DRUG PRODUCT LIST

FEBUXOSTAT

TABLET; ORAL

FEBUXOSTAT

<u>AB</u>	HIKMA	<u>40MG</u>	<u>A205414</u>	<u>001</u>	Oct 15, 2019
<u>AB</u>		<u>80MG</u>	<u>A205414</u>	<u>002</u>	Oct 15, 2019
<u>AB</u>	INDOCO	<u>40MG</u>	<u>A210292</u>	<u>001</u>	Dec 30, 2019
<u>AB</u>		<u>80MG</u>	<u>A210292</u>	<u>002</u>	Dec 30, 2019
<u>AB</u>	MSN	<u>40MG</u>	<u>A210461</u>	<u>001</u>	Dec 30, 2019
<u>AB</u>		<u>80MG</u>	<u>A210461</u>	<u>002</u>	Dec 30, 2019
<u>AB</u>	MYLAN	<u>40MG</u>	<u>A205385</u>	<u>001</u>	Jul 01, 2019
<u>AB</u>		<u>80MG</u>	<u>A205385</u>	<u>002</u>	Jul 01, 2019
<u>AB</u>	SUN PHARM	<u>40MG</u>	<u>A205467</u>	<u>001</u>	Jul 01, 2019
<u>AB</u>		<u>80MG</u>	<u>A205467</u>	<u>002</u>	Jul 01, 2019
<u>AB</u>	SUNSHINE	<u>40MG</u>	<u>A213069</u>	<u>001</u>	Jun 02, 2020
<u>AB</u>		<u>80MG</u>	<u>A213069</u>	<u>002</u>	Jun 02, 2020

ULORIC

<u>AB</u>	+	TAKEDA PHARMS USA	<u>40MG</u>	<u>N021856</u>	<u>001</u>	Feb 13, 2009
<u>AB</u>	+	!	<u>80MG</u>	<u>N021856</u>	<u>002</u>	Feb 13, 2009

FEDRATINIB HYDROCHLORIDE

CAPSULE; ORAL

INREBIC

+	!	IMPACT	EQ 100MG BASE	N212327	001	Aug 16, 2019
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FELBAMATE

SUSPENSION; ORAL

FELBAMATE

<u>AB</u>		AMNEAL PHARMS	<u>600MG/5ML</u>	<u>A202385</u>	<u>001</u>	Dec 16, 2011
<u>AB</u>		NOVITIUM PHARMA	<u>600MG/5ML</u>	<u>A211333</u>	<u>001</u>	May 31, 2019
<u>AB</u>		TARO	<u>600MG/5ML</u>	<u>A206314</u>	<u>001</u>	Jun 16, 2017

FELBATOL

<u>AB</u>	+	!	MYLAN SPECIALITY LP	<u>600MG/5ML</u>	<u>N020189</u>	<u>003</u>	Jul 29, 1993
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TABLET; ORAL

FELBAMATE

<u>AB</u>		ALVOGEN	<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	<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>		CADILA	<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>AB</u>		TARO	<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

FELBATOL

<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
<u>AB</u>		HERITAGE PHARMS INC	<u>2.5MG</u>	<u>A201964</u>	<u>001</u>	Nov 08, 2013
<u>AB</u>			<u>5MG</u>	<u>A201964</u>	<u>002</u>	Nov 08, 2013
<u>AB</u>			<u>10MG</u>	<u>A201964</u>	<u>003</u>	Nov 08, 2013
<u>AB</u>		ORBION PHARMS	<u>2.5MG</u>	<u>A203032</u>	<u>001</u>	May 21, 2015
<u>AB</u>			<u>5MG</u>	<u>A203032</u>	<u>002</u>	May 21, 2015
<u>AB</u>			<u>10MG</u>	<u>A203032</u>	<u>003</u>	May 21, 2015
<u>AB</u>		SUN PHARM INDS LTD	<u>2.5MG</u>	<u>A091200</u>	<u>001</u>	Dec 13, 2013
<u>AB</u>			<u>5MG</u>	<u>A091200</u>	<u>002</u>	Dec 13, 2013
<u>AB</u>			<u>10MG</u>	<u>A091200</u>	<u>003</u>	Dec 13, 2013
<u>AB</u>		SUN PHARM INDUSTRIES	<u>2.5MG</u>	<u>A075896</u>	<u>001</u>	Nov 02, 2004
<u>AB</u>			<u>5MG</u>	<u>A075896</u>	<u>002</u>	Nov 02, 2004
<u>AB</u>			<u>10MG</u>	<u>A075896</u>	<u>003</u>	Nov 02, 2004
<u>AB</u>		TORRENT PHARMS LTD	<u>2.5MG</u>	<u>A202170</u>	<u>001</u>	Nov 28, 2011
<u>AB</u>			<u>5MG</u>	<u>A202170</u>	<u>002</u>	Nov 28, 2011
<u>AB</u>		!	<u>10MG</u>	<u>A202170</u>	<u>003</u>	Nov 28, 2011

PRESCRIPTION DRUG PRODUCT LIST

FELODIPINE

TABLET, EXTENDED RELEASE;ORAL

FELODIPINE

<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	<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
<u>AB</u>	YUNG SHIN PHARM	<u>2.5MG</u>	<u>A204800 001</u>	Apr 29, 2019
<u>AB</u>		<u>5MG</u>	<u>A204800 002</u>	Apr 29, 2019
<u>AB</u>		<u>10MG</u>	<u>A204800 003</u>	Apr 29, 2019

FENFLURAMINE HYDROCHLORIDE

SOLUTION;ORAL

FINTEPLA

+! ZOGENIX INC

EQ 2.2MG BASE/ML

N212102 001 Jun 25, 2020

FENOFIBRATE

CAPSULE;ORAL

ANTARA (MICRONIZED)

<u>AB</u>	+ LUPIN	<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>	ALEMBIC PHARMS LTD	<u>67MG</u>	<u>A213842 001</u>	Oct 19, 2020
<u>AB</u>		<u>134MG</u>	<u>A213842 002</u>	Oct 19, 2020
<u>AB</u>		<u>200MG</u>	<u>A213842 003</u>	Oct 19, 2020
<u>AB</u>	ANI PHARMS	<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	<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>	AUROBINDO PHARMA LTD	<u>67MG</u>	<u>A212232 001</u>	Sep 20, 2021
<u>AB</u>		<u>134MG</u>	<u>A212232 002</u>	Sep 20, 2021
<u>AB</u>		<u>200MG</u>	<u>A212232 003</u>	Sep 20, 2021
<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>	REYOUNG	<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>	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	<u>67MG</u>	<u>A210782 001</u>	Jun 26, 2018
<u>AB</u>		<u>134MG</u>	<u>A210782 002</u>	Jun 26, 2018
<u>AB</u>		<u>200MG</u>	<u>A210782 003</u>	Jun 26, 2018

ANTARA (MICRONIZED)

+ LUPIN

30MG

N021695 004 Oct 18, 2013

+!

90MG

N021695 005 Oct 18, 2013

LIPOFEN

+ CIPHER PHARMS INC

50MG

N021612 001 Jan 11, 2006

+!

150MG

N021612 003 Jan 11, 2006

TABLET;ORAL

FENOFIBRATE

<u>AB</u>	ADAPTIS	<u>54MG</u>	<u>A210606 001</u>	Aug 17, 2018
<u>AB</u>		<u>160MG</u>	<u>A210606 002</u>	Aug 17, 2018
<u>AB</u>	AJANTA PHARMA LTD	<u>54MG</u>	<u>A210138 001</u>	Jul 23, 2018
<u>AB</u>		<u>160MG</u>	<u>A210138 002</u>	Jul 23, 2018

PRESCRIPTION DRUG PRODUCT LIST

FENOFIBRATE

TABLET; ORAL

FENOFIBRATE

<u>AB</u>	ALEMBIC LABS	<u>54MG</u>	<u>A209660</u>	<u>001</u>	Feb 11, 2019
<u>AB</u>		<u>160MG</u>	<u>A209660</u>	<u>002</u>	Feb 11, 2019
<u>AB</u>	ALEMBIC PHARMS LTD	<u>48MG</u>	<u>A210476</u>	<u>001</u>	Aug 09, 2019
<u>AB</u>		<u>145MG</u>	<u>A210476</u>	<u>002</u>	Aug 09, 2019
<u>AB</u>	AMNEAL	<u>48MG</u>	<u>A209951</u>	<u>001</u>	Feb 09, 2018
<u>AB</u>		<u>54MG</u>	<u>A209950</u>	<u>001</u>	Mar 19, 2018
<u>AB</u>		<u>145MG</u>	<u>A209951</u>	<u>002</u>	Feb 09, 2018
<u>AB</u>		<u>160MG</u>	<u>A209950</u>	<u>002</u>	Mar 19, 2018
<u>AB</u>	AUROBINDO PHARMA LTD	<u>48MG</u>	<u>A205118</u>	<u>001</u>	May 05, 2016
<u>AB</u>		<u>145MG</u>	<u>A205118</u>	<u>002</u>	May 05, 2016
<u>AB</u>	AUSTARPHARMA	<u>48MG</u>	<u>A208476</u>	<u>001</u>	Feb 10, 2021
<u>AB</u>		<u>54MG</u>	<u>A207803</u>	<u>001</u>	Dec 19, 2017
<u>AB</u>		<u>145MG</u>	<u>A208476</u>	<u>002</u>	Feb 10, 2021
<u>AB</u>		<u>160MG</u>	<u>A207803</u>	<u>002</u>	Dec 19, 2017
<u>AB</u>	CIPLA	<u>48MG</u>	<u>A208709</u>	<u>001</u>	Dec 15, 2016
<u>AB</u>		<u>145MG</u>	<u>A208709</u>	<u>002</u>	Dec 15, 2016
<u>AB</u>	DR REDDYS	<u>54MG</u>	<u>A210670</u>	<u>001</u>	Sep 06, 2019
<u>AB</u>		<u>160MG</u>	<u>A210670</u>	<u>002</u>	Sep 06, 2019
<u>AB</u>	GRAVITI PHARMS	<u>48MG</u>	<u>A211122</u>	<u>001</u>	Mar 18, 2020
<u>AB</u>		<u>145MG</u>	<u>A211122</u>	<u>002</u>	Mar 18, 2020
<u>AB</u>	HETERO LABS LTD III	<u>48MG</u>	<u>A204598</u>	<u>001</u>	Jul 12, 2016
<u>AB</u>		<u>145MG</u>	<u>A204598</u>	<u>002</u>	Jul 12, 2016
<u>AB</u>	IMPAX LABS	<u>54MG</u>	<u>A076509</u>	<u>001</u>	Mar 26, 2008
<u>AB</u>	!	<u>160MG</u>	<u>A076509</u>	<u>002</u>	Mar 26, 2008
<u>AB</u>	LUPIN LTD	<u>48MG</u>	<u>A090856</u>	<u>001</u>	Dec 23, 2011
<u>AB</u>		<u>54MG</u>	<u>A204019</u>	<u>001</u>	Aug 17, 2015
<u>AB</u>		<u>145MG</u>	<u>A090856</u>	<u>002</u>	Dec 23, 2011
<u>AB</u>		<u>160MG</u>	<u>A204019</u>	<u>002</u>	Aug 17, 2015
<u>AB</u>	MANKIND PHARMA	<u>54MG</u>	<u>A213864</u>	<u>001</u>	Jun 12, 2020
<u>AB</u>		<u>160MG</u>	<u>A213864</u>	<u>002</u>	Jun 12, 2020
<u>AB</u>	MYLAN	<u>54MG</u>	<u>A076520</u>	<u>001</u>	Oct 25, 2007
<u>AB</u>		<u>160MG</u>	<u>A076520</u>	<u>003</u>	Oct 25, 2007
<u>AB</u>	MYLAN PHARMS INC	<u>40MG</u>	<u>A204475</u>	<u>001</u>	Jun 23, 2016
<u>AB</u>		<u>48MG</u>	<u>A202856</u>	<u>001</u>	Dec 07, 2012
<u>AB</u>		<u>120MG</u>	<u>A204475</u>	<u>002</u>	Jun 23, 2016
<u>AB</u>		<u>145MG</u>	<u>A202856</u>	<u>002</u>	Dec 07, 2012
<u>AB</u>	PRINSTON INC	<u>48MG</u>	<u>A211080</u>	<u>001</u>	Aug 28, 2018
<u>AB</u>		<u>145MG</u>	<u>A211080</u>	<u>002</u>	Aug 28, 2018
<u>AB</u>	RHODES PHARMS	<u>54MG</u>	<u>A076433</u>	<u>001</u>	May 13, 2005
<u>AB</u>		<u>160MG</u>	<u>A076433</u>	<u>002</u>	May 13, 2005
<u>AB</u>	SUN PHARM	<u>48MG</u>	<u>A200884</u>	<u>001</u>	Sep 07, 2017
<u>AB</u>		<u>145MG</u>	<u>A200884</u>	<u>002</u>	Sep 07, 2017
<u>AB</u>	SUN PHARM INDS LTD	<u>54MG</u>	<u>A076635</u>	<u>001</u>	Oct 31, 2005
<u>AB</u>		<u>160MG</u>	<u>A076635</u>	<u>003</u>	Oct 31, 2005
<u>AB</u>	VALEANT PHARMS NORTH	<u>48MG</u>	<u>A090715</u>	<u>001</u>	Apr 05, 2012
<u>AB</u>		<u>145MG</u>	<u>A090715</u>	<u>002</u>	Apr 05, 2012

FENOGLIDE

<u>AB</u>	+ SALIX	<u>40MG</u>	<u>N022118</u>	<u>001</u>	Aug 10, 2007
<u>AB</u>	+	<u>120MG</u>	<u>N022118</u>	<u>002</u>	Aug 10, 2007

TRICOR

<u>AB</u>	+ ABBVIE	<u>48MG</u>	<u>N021656</u>	<u>001</u>	Nov 05, 2004
<u>AB</u>	+	<u>145MG</u>	<u>N021656</u>	<u>002</u>	Nov 05, 2004

TRIGLIDE

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

	+	ATHENA	105MG	N022418	002	Aug 14, 2009
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FENOLDOPAM MESYLATE

INJECTABLE; INJECTION

CORLOPAM

<u>AP</u>	+	HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>N019922</u>	<u>001</u>	Sep 23, 1997
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FENOLDOPAM MESYLATE

<u>AP</u>		HIKMA	<u>EQ 10MG BASE/ML</u>	<u>A076582</u>	<u>001</u>	Oct 12, 2004
<u>AP</u>		SANDOZ INC	<u>EQ 10MG BASE/ML</u>	<u>A077155</u>	<u>001</u>	Feb 15, 2005

PRESCRIPTION DRUG PRODUCT LIST

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

FENTANYL-100AB AVEVA100MCG/HRA077449 004 Oct 20, 2008AB KINDEVA100MCG/HRA202097 005 Nov 04, 2016AB MYLAN TECHNOLOGIES100MCG/HRA076258 004 Jan 28, 2005AB SPECGX LLC100MCG/HRA077154 004 Feb 09, 2011FENTANYL-12AB AVEVA12.5MCG/HRA077449 005 Sep 11, 2015AB KINDEVA12.5MCG/HRA202097 001 Nov 04, 2016AB MYLAN TECHNOLOGIES12.5MCG/HRA076258 005 Jan 23, 2007AB SPECGX LLC12.5MCG/HRA077154 005 Jun 11, 2015FENTANYL-25AB AVEVA25MCG/HRA077449 001 Oct 20, 2008AB KINDEVA25MCG/HRA202097 002 Nov 04, 2016AB MYLAN TECHNOLOGIES25MCG/HRA076258 001 Jan 28, 2005AB ! SPECGX LLC25MCG/HRA077154 001 Feb 09, 2011FENTANYL-37AB AVEVA37.5MCG/HRA077449 006 Dec 06, 2017AB MYLAN TECHNOLOGIES37.5MCG/HRA076258 006 Dec 29, 2014AB SPECGX LLC37.5MCG/HRA077154 006 Jan 14, 2020FENTANYL-50AB AVEVA50MCG/HRA077449 002 Oct 20, 2008AB KINDEVA50MCG/HRA202097 003 Nov 04, 2016AB MYLAN TECHNOLOGIES50MCG/HRA076258 002 Jan 28, 2005AB SPECGX LLC50MCG/HRA077154 002 Feb 09, 2011FENTANYL-62AB AVEVA62.5MCG/HRA077449 007 Dec 06, 2017AB MYLAN TECHNOLOGIES62.5MCG/HRA076258 007 Dec 29, 2014AB SPECGX LLC62.5MCG/HRA077154 007 Jan 14, 2020FENTANYL-75AB AVEVA75MCG/HRA077449 003 Oct 20, 2008AB KINDEVA75MCG/HRA202097 004 Nov 04, 2016AB MYLAN TECHNOLOGIES75MCG/HRA076258 003 Jan 28, 2005AB SPECGX LLC75MCG/HRA077154 003 Feb 09, 2011FENTANYL-87AB AVEVA87.5MCG/HRA077449 008 Dec 06, 2017AB MYLAN TECHNOLOGIES87.5MCG/HRA076258 008 Dec 29, 2014

SPRAY; SUBLINGUAL

SUBSYS

+ BTCP PHARMA

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 FRESENIUS KABI USAEQ 0.05MG BASE/MLA212086 001 Sep 01, 2020AP HOSPIRAEQ 0.05MG BASE/MLN019115 001 Jan 12, 1985FENTANYL CITRATE PRESERVATIVE FREEAP FRESENIUS KABI USAEQ 0.05MG BASE/MLA210762 001 May 03, 2019AP +! HIKMAEQ 0.05MG BASE/MLN019101 001 Jul 11, 1984AP HOSPIRAEQ 0.05MG BASE/MLA072786 001 Sep 24, 1991SUBLIMAZE PRESERVATIVE FREEAP +! AKORNEQ 0.05MG BASE/MLN016619 001

SPRAY, METERED; NASAL

LAZANDA

+ BTCP PHARMA

EQ 0.1MG BASE

N022569 001 Jun 30, 2011

+!

EQ 0.4MG BASE

N022569 002 Jun 30, 2011

PRESCRIPTION DRUG PRODUCT LIST

FENTANYL CITRATE

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

TROCHE/LOZENGE;TRANSMUCOSAL

ACTIQ

AB	+	CEPHALON	<u>EQ 0.2MG BASE</u>	<u>N020747 001</u>	Nov 04, 1998
AB	+	!	<u>EQ 0.4MG BASE</u>	<u>N020747 002</u>	Nov 04, 1998
AB	+		<u>EQ 0.6MG BASE</u>	<u>N020747 003</u>	Nov 04, 1998
AB	+		<u>EQ 0.8MG BASE</u>	<u>N020747 004</u>	Nov 04, 1998
AB	+		<u>EQ 1.2MG BASE</u>	<u>N020747 005</u>	Nov 04, 1998
AB	+		<u>EQ 1.6MG BASE</u>	<u>N020747 006</u>	Nov 04, 1998

FENTANYL CITRATE

AB		SPECGX LLC	<u>EQ 0.2MG BASE</u>	<u>A078907 001</u>	Oct 30, 2009
AB			<u>EQ 0.4MG BASE</u>	<u>A078907 002</u>	Oct 30, 2009
AB			<u>EQ 0.6MG BASE</u>	<u>A078907 003</u>	Oct 30, 2009
AB			<u>EQ 0.8MG BASE</u>	<u>A078907 004</u>	Oct 30, 2009
AB			<u>EQ 1.2MG BASE</u>	<u>A078907 005</u>	Oct 30, 2009
AB			<u>EQ 1.6MG BASE</u>	<u>A078907 006</u>	Oct 30, 2009

FERRIC CARBOXYMALTOSE

SOLUTION;INTRAVENOUS

INJECTAFER

+	AM REGENT	750MG IRON/15ML (50MG IRON/ML)	N203565 001	Jul 25, 2013
+	!	500MG IRON/10ML (50MG IRON/ML)	N203565 002	Oct 08, 2020
+	!	1GM IRON/20ML (50MG IRON/ML)	N203565 003	Apr 28, 2021

FERRIC CITRATE

TABLET;ORAL

AURYXIA

+	KERYX BIOPHARMS	EQ 210MG IRON	N205874 001	Sep 05, 2014
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FERRIC DERISOMALTOSE

SOLUTION;INTRAVENOUS

MONOFERRIC

+	PHARMACOSMOS AS	1GM/10ML (100MG/ML)	N208171 003	Jan 16, 2020
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FERRIC HEXACYANOFERRATE(II)

CAPSULE;ORAL

RADIOGARDASE (PRUSSIAN BLUE)

+	HEYL CHEMISCH	500MG	N021626 001	Oct 02, 2003
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FERRIC MALTOL

CAPSULE;ORAL

ACCRUFER

+	SHIELD TX	30MG IRON	N212320 001	Jul 25, 2019
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FERRIC OXYHYDROXIDE

INJECTABLE;INJECTION

FERLECIT

AB	+	SANOFI AVENTIS US	<u>EQ 62.5MG IRON/5ML (EQ 12.5MG IRON/ML)</u>	<u>N020955 001</u>	Feb 18, 1999
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SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE

AB		WEST-WARD PHARMS INT	<u>EQ 62.5MG IRON/5ML (EQ 12.5MG IRON/ML)</u>	<u>A078215 001</u>	Mar 31, 2011
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INFED

BP	+	ALLERGAN	EQ 100MG IRON/2ML (EQ 50MG IRON/ML)	N017441 001	
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INJECTABLE;INTRAVENOUS

VENOFER

+	AM REGENT	EQ 50MG IRON/2.5ML (EQ 20MG IRON/ML)	N021135 002	Mar 20, 2005
+	!	EQ 100MG IRON/5ML (EQ 20MG IRON/ML)	N021135 001	Nov 06, 2000
+		EQ 200MG IRON/10ML (EQ 20MG IRON/ML)	N021135 004	Feb 09, 2007

TABLET, CHEWABLE;ORAL

VELPHORO

+	VIFOR FRESENIUS	EQ 500MG IRON	N205109 001	Nov 27, 2013
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FERRIC PYROPHOSPHATE CITRATE

FOR SOLUTION;INTRAVENOUS

TRIFERIC

+	ROCKWELL MEDICAL INC	272MG IRON/PACKET	N208551 001	Apr 25, 2016
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PRESCRIPTION DRUG PRODUCT LIST

FERRIC PYROPHOSPHATE CITRATE

SOLUTION;INTRAVENOUS

TRIFERIC

+	!	ROCKWELL MEDICAL INC	27.2MG IRON/5ML (5.44MG IRON/ML)	N206317	001	Jan 23, 2015
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TRIFERIC AVNU

+	!	ROCKWELL MEDICAL INC	6.75MG IRON/4.5ML (1.5MG IRON/ML)	N212860	001	Mar 27, 2020
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FERUMOXYTOL

SOLUTION;INTRAVENOUS

FERAHEME

<u>AB</u>	+	!	COVIS	<u>EQ 510MG IRON/17ML (EQ 30MG IRON/ML)</u>	<u>N022180</u>	<u>001</u>	Jun 30, 2009
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FERUMOXYTOL

<u>AB</u>			SANDOZ INC	<u>EQ 510MG IRON/17ML (EQ 30MG IRON/ML)</u>	<u>A206604</u>	<u>001</u>	Jan 15, 2021
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FESOTERODINE FUMARATE

TABLET, EXTENDED RELEASE;ORAL

FESOTERODINE FUMARATE

<u>AB</u>			AUROBINDO PHARMA LTD	<u>4MG</u>	<u>A205007</u>	<u>001</u>	Feb 17, 2017
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<u>AB</u>				<u>8MG</u>	<u>A205007</u>	<u>002</u>	Feb 17, 2017
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<u>AB</u>			ZYDUS PHARMS	<u>4MG</u>	<u>A204946</u>	<u>001</u>	Oct 03, 2017
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<u>AB</u>				<u>8MG</u>	<u>A204946</u>	<u>002</u>	Oct 03, 2017
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TOVIAZ

<u>AB</u>	+		PFIZER	<u>4MG</u>	<u>N022030</u>	<u>001</u>	Oct 31, 2008
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<u>AB</u>	+	!		<u>8MG</u>	<u>N022030</u>	<u>002</u>	Oct 31, 2008
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FEXINIDAZOLE

TABLET;ORAL

FEXINIDAZOLE

+	!	SANOFI	600MG	N214429	001	Jul 16, 2021
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FEXOFENADINE HYDROCHLORIDE

TABLET;ORAL

FEXOFENADINE HYDROCHLORIDE

<u>AB</u>			DR REDDYS LABS LTD	<u>30MG</u>	<u>A076502</u>	<u>001</u>	Apr 11, 2006
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<u>AB</u>				<u>60MG</u>	<u>A076502</u>	<u>002</u>	Apr 11, 2006
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<u>AB</u>				<u>180MG</u>	<u>A076502</u>	<u>003</u>	Apr 11, 2006
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<u>AB</u>			RISING	<u>60MG</u>	<u>A077081</u>	<u>003</u>	Apr 11, 2008
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<u>AB</u>				<u>180MG</u>	<u>A077081</u>	<u>001</u>	Apr 16, 2007
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<u>AB</u>			TEVA	<u>30MG</u>	<u>A076447</u>	<u>001</u>	Sep 01, 2005
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<u>AB</u>				<u>60MG</u>	<u>A076447</u>	<u>002</u>	Sep 01, 2005
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<u>AB</u>				<u>180MG</u>	<u>A076447</u>	<u>003</u>	Sep 01, 2005
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FIDAXOMICIN

FOR SUSPENSION;ORAL

DIFICID

+	!	CUBIST PHARMS LLC	40MG/ML	N213138	001	Jan 24, 2020
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TABLET;ORAL

DIFICID

+	!	CUBIST PHARMS LLC	200MG	N201699	001	May 27, 2011
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FINASTERIDE

TABLET;ORAL

FINASTERIDE

<u>AB</u>			ACCORD HLTHCARE	<u>1MG</u>	<u>A091643</u>	<u>001</u>	Nov 05, 2013
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<u>AB</u>				<u>5MG</u>	<u>A090121</u>	<u>001</u>	Feb 23, 2010
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<u>AB</u>			ALKEM LABS LTD	<u>1MG</u>	<u>A207750</u>	<u>001</u>	Jan 06, 2017
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<u>AB</u>				<u>5MG</u>	<u>A204304</u>	<u>001</u>	Jan 05, 2017
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<u>AB</u>			AUROBINDO PHARMA	<u>5MG</u>	<u>A078341</u>	<u>001</u>	Oct 30, 2007
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<u>AB</u>			AUROBINDO PHARMA LTD	<u>1MG</u>	<u>A203687</u>	<u>001</u>	Nov 05, 2013
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<u>AB</u>			CIPLA	<u>1MG</u>	<u>A077335</u>	<u>001</u>	Nov 20, 2014
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<u>AB</u>			DR REDDYS LABS INC	<u>1MG</u>	<u>A076436</u>	<u>001</u>	Jul 28, 2006
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<u>AB</u>			DR REDDYS LABS LTD	<u>5MG</u>	<u>A076437</u>	<u>001</u>	Feb 28, 2007
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<u>AB</u>			HETERO LABS LTD III	<u>1MG</u>	<u>A090060</u>	<u>001</u>	Jul 01, 2013
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<u>AB</u>				<u>5MG</u>	<u>A090061</u>	<u>001</u>	Jun 07, 2010
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<u>AB</u>			SUN PHARM	<u>1MG</u>	<u>A090508</u>	<u>001</u>	Jul 01, 2013
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<u>AB</u>				<u>5MG</u>	<u>A090507</u>	<u>001</u>	Aug 16, 2011
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<u>AB</u>			TEVA	<u>5MG</u>	<u>A076511</u>	<u>001</u>	Dec 15, 2006
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<u>AB</u>			ZYDUS PHARMS USA	<u>5MG</u>	<u>A078900</u>	<u>001</u>	Dec 28, 2009
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PROPECIA

<u>AB</u>	+	!	MERCK	<u>1MG</u>	<u>N020788</u>	<u>001</u>	Dec 19, 1997
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PRESCRIPTION DRUG PRODUCT LIST

FINASTERIDE

TABLET; ORAL

PROSCAR

AB	+ !	MERCK	5MG	N020180	001	Jun 19, 1992
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FINASTERIDE; TADALAFIL

CAPSULE; ORAL

ENTADFI

+ !	VERU	5MG; 5MG	N215423	001	Dec 09, 2021
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FINERENONE

TABLET; ORAL

KERENDIA

+	BAYER HLTHCARE	10MG	N215341	001	Jul 09, 2021
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+ !		20MG	N215341	002	Jul 09, 2021
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FINGOLIMOD HYDROCHLORIDE

CAPSULE; ORAL

FINGOLIMOD HYDROCHLORIDE

AB		ACCORD HLTHCARE	EQ 0.5MG BASE	A207991	001	Oct 28, 2020
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AB		HEC PHARM CO LTD	EQ 0.5MG BASE	A207939	001	Nov 10, 2021
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AB		MYLAN	EQ 0.5MG BASE	A208005	001	Jan 19, 2021
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AB		TEVA PHARMS USA	EQ 0.25MG BASE	A212152	001	Nov 12, 2021
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GILENYA

AB	+	NOVARTIS	EQ 0.25MG BASE	N022527	002	May 11, 2018
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AB	+ !		EQ 0.5MG BASE	N022527	001	Sep 21, 2010
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FINGOLIMOD LAURYL SULFATE

TABLET, ORALLY DISINTEGRATING; ORAL

TASCENSO ODT

+ !	HANDA	EQ 0.25MG BASE	N214962	001	Dec 23, 2021
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FISH OIL TRIGLYCERIDES

EMULSION; INTRAVENOUS

OMEGAVEN

+ !	FRESENIUS KABI USA	5GM/50ML (0.1GM/ML)	N210589	001	Jul 27, 2018
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+ !		10GM/100ML (0.1GM/ML)	N210589	002	Jul 27, 2018
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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
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+ !		3GM/100ML; 6GM/100ML; 5GM/100ML; 6GM/100ML (250ML)	N207648	002	Jul 13, 2016
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+ !		3GM/100ML; 6GM/100ML; 5GM/100ML; 6GM/100ML (500ML)	N207648	003	Jul 13, 2016
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+ !		3GM/100ML; 6GM/100ML; 5GM/100ML; 6GM/100ML (1000ML)	N207648	004	Aug 10, 2018
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FLAVOXATE HYDROCHLORIDE

TABLET; ORAL

FLAVOXATE HYDROCHLORIDE

AB		EPIC PHARMA	100MG	A076835	001	Nov 30, 2005
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AB	!	PADAGIS US	100MG	A076831	001	Dec 16, 2004
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FLECAINIDE ACETATE

TABLET; ORAL

FLECAINIDE ACETATE

AB		AMNEAL PHARM	50MG	A075442	001	Jul 31, 2001
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AB			100MG	A075442	002	Jul 31, 2001
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AB			150MG	A075442	003	Jul 31, 2001
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AB		ANI PHARMS	50MG	A075882	001	Oct 28, 2002
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AB			100MG	A075882	002	Oct 28, 2002
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AB			150MG	A075882	003	Oct 28, 2002
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AB		AUROBINDO PHARMA LTD	50MG	A202821	001	Nov 03, 2017
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AB			100MG	A202821	002	Nov 03, 2017
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AB			150MG	A202821	003	Nov 03, 2017
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AB		BEXIMCO PHARMS USA	50MG	A210683	001	Sep 16, 2020
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AB			100MG	A210683	002	Sep 16, 2020
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AB			150MG	A210683	003	Sep 16, 2020
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AB		HIKMA	50MG	A076278	001	Jan 14, 2003
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AB			100MG	A076278	002	Jan 14, 2003
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AB	!		150MG	A076278	003	Jan 14, 2003
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AB		SUN PHARM INDS LTD	50MG	A076421	001	Mar 28, 2003
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AB			100MG	A076421	002	Mar 28, 2003
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PRESCRIPTION DRUG PRODUCT LIST

FLECAINIDE ACETATE

TABLET; ORAL

FLECAINIDE ACETATE

AB		150MG	A076421 003	Mar 28, 2003
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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
INC

10-30ML (13.5-51mCi/ML)

N202008 002 Apr 06, 2012

+! 10-50ML (13.5-51mCi/ML)

N202008 003 Apr 06, 2012

FLORTAUCIPIR F-18

SOLUTION; INTRAVENOUS

TAUVID

+! AVID RADIOPHARMS
INC

30ML (8.1-51mCi/ML)

N212123 001 May 28, 2020

+! 50ML (8.1-51mCi/ML)

N212123 002 May 28, 2020

FLOXURIDINE

INJECTABLE; INJECTION

FLOXURIDINE

AP	FRESENIUS KABI USA	500MG/VIAL	A075837 001	Feb 22, 2001
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AP	! HIKMA	500MG/VIAL	A075387 001	Apr 16, 2000
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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
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AB	+!	200MG/5ML	N020090 002	Dec 23, 1993
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FLUCONAZOLE

AB	AUROBINDO PHARMA LTD	50MG/5ML	A079150 001	Sep 18, 2009
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AB		200MG/5ML	A079150 002	Sep 18, 2009
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INJECTABLE; INJECTION

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

AP	BAXTER HLTHCARE CORP	200MG/100ML (2MG/ML)	A077947 001	May 26, 2010
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AP		400MG/200ML (2MG/ML)	A077947 002	May 26, 2010
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AP	FRESENIUS KABI USA	200MG/100ML (2MG/ML)	A076145 001	Jul 29, 2004
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AP		400MG/200ML (2MG/ML)	A076145 002	Jul 29, 2004
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AP	HIKMA	200MG/100ML (2MG/ML)	A076087 001	Jul 29, 2004
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AP		400MG/200ML (2MG/ML)	A076087 003	Jul 29, 2004
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AP	HIKMA FARMACEUTICA	200MG/100ML (2MG/ML)	A076736 001	Aug 23, 2005
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FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	200MG/100ML (2MG/ML)	A076766 001	Jul 29, 2004
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AP		400MG/200ML (2MG/ML)	A076766 002	Jul 29, 2004
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AP	HIKMA FARMACEUTICA	200MG/100ML (2MG/ML)	A078698 001	Jan 30, 2012
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AP		400MG/200ML (2MG/ML)	A078698 002	Jan 30, 2012
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AP	HOSPIRA	200MG/100ML (2MG/ML)	A076303 001	Jul 29, 2004
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AP		400MG/200ML (2MG/ML)	A076303 002	Jul 29, 2004
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AP	! INFORLIFE	200MG/100ML (2MG/ML)	A079104 001	Jul 30, 2009
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AP	!	400MG/200ML (2MG/ML)	A079104 002	Jul 30, 2009
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AP	WEST-WARD PHARMS INT	200MG/100ML (2MG/ML)	A078107 001	Jul 30, 2008
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AP		400MG/200ML (2MG/ML)	A078107 002	Jul 30, 2008
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AP	WOODWARD	200MG/100ML (2MG/ML)	A077909 001	May 26, 2010
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AP		400MG/200ML (2MG/ML)	A077909 002	May 26, 2010
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FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

	HIKMA	100MG/50ML (2MG/ML)	A076087 002	Sep 26, 2008
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PRESCRIPTION DRUG PRODUCT LIST

FLUCONAZOLE

INJECTABLE; INJECTION

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
WOODWARD 100MG/50ML (2MG/ML)

A077909 003 Apr 20, 2015

TABLET; ORAL

DIFLUCAN

<u>AB</u>	+	PFIZER	<u>50MG</u>	<u>N019949</u>	<u>001</u>	Jan 29, 1990
<u>AB</u>	+		<u>100MG</u>	<u>N019949</u>	<u>002</u>	Jan 29, 1990
<u>AB</u>	+		<u>150MG</u>	<u>N019949</u>	<u>004</u>	Jun 30, 1994
<u>AB</u>	+!		<u>200MG</u>	<u>N019949</u>	<u>003</u>	Jan 29, 1990

FLUCONAZOLE

<u>AB</u>		ANI PHARMS	<u>50MG</u>	<u>A078423</u>	<u>001</u>	Mar 07, 2011
<u>AB</u>			<u>100MG</u>	<u>A078423</u>	<u>002</u>	Mar 07, 2011
<u>AB</u>			<u>150MG</u>	<u>A078423</u>	<u>003</u>	Mar 07, 2011
<u>AB</u>			<u>200MG</u>	<u>A078423</u>	<u>004</u>	Mar 07, 2011
<u>AB</u>		AUROBINDO PHARMA	<u>50MG</u>	<u>A077731</u>	<u>001</u>	Oct 07, 2008
<u>AB</u>			<u>100MG</u>	<u>A077731</u>	<u>002</u>	Oct 07, 2008
<u>AB</u>			<u>150MG</u>	<u>A077731</u>	<u>003</u>	Oct 07, 2008
<u>AB</u>			<u>200MG</u>	<u>A077731</u>	<u>004</u>	Oct 07, 2008
<u>AB</u>		CHARTWELL	<u>50MG</u>	<u>A076665</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>			<u>100MG</u>	<u>A076665</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>			<u>150MG</u>	<u>A076665</u>	<u>003</u>	Jul 29, 2004
<u>AB</u>			<u>200MG</u>	<u>A076665</u>	<u>004</u>	Jul 29, 2004
<u>AB</u>		DR REDDYS LABS INC	<u>50MG</u>	<u>A076658</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>			<u>100MG</u>	<u>A076658</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>			<u>150MG</u>	<u>A076658</u>	<u>003</u>	Jul 29, 2004
<u>AB</u>			<u>200MG</u>	<u>A076658</u>	<u>004</u>	Jul 29, 2004
<u>AB</u>		GLENMARK GENERICS	<u>50MG</u>	<u>A077253</u>	<u>001</u>	Jan 25, 2006
<u>AB</u>			<u>100MG</u>	<u>A077253</u>	<u>002</u>	Jan 25, 2006
<u>AB</u>			<u>150MG</u>	<u>A077253</u>	<u>003</u>	Jan 25, 2006
<u>AB</u>			<u>200MG</u>	<u>A077253</u>	<u>004</u>	Jan 25, 2006
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>50MG</u>	<u>A076077</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>			<u>100MG</u>	<u>A076077</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>			<u>150MG</u>	<u>A076077</u>	<u>003</u>	Jul 29, 2004
<u>AB</u>			<u>200MG</u>	<u>A076077</u>	<u>004</u>	Jul 29, 2004
<u>AB</u>		TARO	<u>50MG</u>	<u>A076507</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>			<u>100MG</u>	<u>A076507</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>			<u>150MG</u>	<u>A076507</u>	<u>003</u>	Jul 29, 2004
<u>AB</u>			<u>200MG</u>	<u>A076507</u>	<u>004</u>	Jul 29, 2004
<u>AB</u>		UNIQUE PHARM LABS	<u>50MG</u>	<u>A076957</u>	<u>001</u>	Sep 28, 2005
<u>AB</u>			<u>100MG</u>	<u>A076957</u>	<u>002</u>	Sep 28, 2005
<u>AB</u>			<u>150MG</u>	<u>A076957</u>	<u>004</u>	Feb 27, 2017
<u>AB</u>			<u>200MG</u>	<u>A076957</u>	<u>003</u>	Sep 28, 2005
<u>AB</u>		ZYDUS PHARMS	<u>50MG</u>	<u>A208963</u>	<u>001</u>	Feb 16, 2017
<u>AB</u>			<u>100MG</u>	<u>A208963</u>	<u>002</u>	Feb 16, 2017
<u>AB</u>			<u>150MG</u>	<u>A208963</u>	<u>003</u>	Feb 16, 2017
<u>AB</u>			<u>200MG</u>	<u>A208963</u>	<u>004</u>	Feb 16, 2017

FLUCYTOSINE

CAPSULE; ORAL

ANCOBON

<u>AB</u>	+	BAUSCH	<u>250MG</u>	<u>N017001</u>	<u>001</u>	
<u>AB</u>	+!		<u>500MG</u>	<u>N017001</u>	<u>002</u>	

FLUCYTOSINE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A213665</u>	<u>001</u>	May 01, 2020
<u>AB</u>			<u>500MG</u>	<u>A213665</u>	<u>002</u>	May 01, 2020
<u>AB</u>		HIKMA	<u>250MG</u>	<u>A206550</u>	<u>001</u>	Oct 17, 2017
<u>AB</u>			<u>500MG</u>	<u>A206550</u>	<u>002</u>	Oct 17, 2017
<u>AB</u>		NOVEL LABS INC	<u>250MG</u>	<u>A204652</u>	<u>001</u>	Jul 07, 2017
<u>AB</u>			<u>500MG</u>	<u>A204652</u>	<u>002</u>	Jul 07, 2017
<u>AB</u>		SIGMAPHARM LABS LLC	<u>250MG</u>	<u>A201566</u>	<u>001</u>	Jun 28, 2011
<u>AB</u>			<u>500MG</u>	<u>A201566</u>	<u>002</u>	Jun 28, 2011

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARABINE PHOSPHATE

<u>AP</u>		ACTAVIS LLC	<u>50MG/2ML (25MG/ML)</u>	<u>A203738</u>	<u>001</u>	Feb 28, 2017
<u>AP</u>	!	ACTAVIS TOTOWA	<u>50MG/VIAL</u>	<u>A078610</u>	<u>001</u>	Feb 11, 2009
<u>AP</u>		AREVA PHARMS	<u>50MG/2ML (25MG/ML)</u>	<u>A090724</u>	<u>001</u>	Sep 27, 2010
<u>AP</u>		CUSTOPHARM INC	<u>50MG/VIAL</u>	<u>A076349</u>	<u>001</u>	Aug 28, 2003
<u>AP</u>	!	FRESENIUS KABI USA	<u>50MG/2ML (25MG/ML)</u>	<u>A078393</u>	<u>001</u>	Oct 15, 2007
<u>AP</u>			<u>50MG/VIAL</u>	<u>A078544</u>	<u>001</u>	Oct 15, 2007

PRESCRIPTION DRUG PRODUCT LIST

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARABINE PHOSPHATE

<u>AP</u>	SAGENT PHARMS INC	<u>50MG/2ML (25MG/ML)</u>	<u>A076661 001</u>	Apr 28, 2004
<u>AP</u>	+! SANDOZ	<u>50MG/2ML (25MG/ML)</u>	<u>N022137 001</u>	Sep 21, 2007

FLUDEOXYGLUCOSE F-18

INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F18

<u>AP</u>	3D IMAGING DRUG	<u>20-300mCi/ML</u>	<u>A203778 001</u>	Oct 30, 2015
<u>AP</u>	BIOMEDCL RES FDN	<u>20-300mCi/ML</u>	<u>A203710 001</u>	May 01, 2015
<u>AP</u>		<u>20-300mCi/ML</u>	<u>A203837 001</u>	May 01, 2015
<u>AP</u>	BRIGHAM WOMENS	<u>20-300mCi/ML</u>	<u>A203816 001</u>	Oct 30, 2014
<u>AP</u>	CARDINAL HEALTH 414	<u>20-300mCi/ML</u>	<u>A203603 001</u>	Nov 13, 2015
<u>AP</u>		<u>20-500mCi/ML</u>	<u>A203603 002</u>	Sep 27, 2018
<u>AP</u>	CHILDRENS HOSP MI	<u>20-300mCi/ML</u>	<u>A204385 001</u>	Oct 29, 2014
<u>AP</u>	DECATUR	<u>20-300mCi/ML</u>	<u>A204463 001</u>	Oct 21, 2014
<u>AP</u>	ESSENTIAL ISOTOPES	<u>20-300mCi/ML</u>	<u>A203946 001</u>	Feb 05, 2014
<u>AP</u>	+! FEINSTEIN	<u>20-400mCi/ML</u>	<u>N021870 002</u>	Nov 21, 2008
<u>AP</u>	ISOLOGIC INNOVATIVE	<u>20-300mCi/ML</u>	<u>A204525 001</u>	Oct 29, 2014
<u>AP</u>	JUBILANT DRAXIMAGE	<u>20-300mCi/ML</u>	<u>A203920 001</u>	Jun 23, 2015
<u>AP</u>	KETTERING MEDCTR	<u>4-40mCi/ML</u>	<u>A204759 001</u>	Oct 27, 2015
<u>AP</u>	KREITCHMAN PET CTR	<u>10-100mCi/ML</u>	<u>A203942 001</u>	Apr 11, 2016
<u>AP</u>	MA GENERAL HOSP	<u>20-300mCi/ML</u>	<u>A204333 001</u>	Sep 25, 2014
<u>AP</u>	MCPRF	<u>20-240mCi/ML</u>	<u>A203612 001</u>	Aug 05, 2013
<u>AP</u>	MEM SLOAN-KETTERING	<u>20-300mCi/ML</u>	<u>A208679 001</u>	Dec 08, 2016
<u>AP</u>	METHODIST HOSP RES	<u>20-300mCi/ML</u>	<u>A203904 001</u>	Apr 23, 2015
<u>AP</u>	MIPS CRF	<u>20-300mCi/ML</u>	<u>A204472 001</u>	Sep 11, 2015
<u>AP</u>	NCM USA BRONX LLC	<u>20-300mCi/ML</u>	<u>A204512 001</u>	Jan 07, 2015
<u>AP</u>	! PETNET	<u>20-200mCi/ML</u>	<u>A079086 001</u>	Feb 25, 2011
<u>AP</u>	PHARMALOGIC HLDGS	<u>20-200mCi/ML</u>	<u>A203664 001</u>	Feb 04, 2014
<u>AP</u>	PRECISION NUCLEAR	<u>20-500mCi/ML</u>	<u>A204546 001</u>	Apr 07, 2015
<u>AP</u>	! QUEEN HAMAMATSU PET	<u>10-100mCi/ML</u>	<u>A203771 001</u>	Aug 31, 2015
<u>AP</u>	SHERTECH LABS LLC	<u>20-300mCi/ML</u>	<u>A204264 001</u>	Dec 18, 2014
<u>AP</u>	SOFIE	<u>20-300mCi/ML</u>	<u>A203591 001</u>	Aug 31, 2015
<u>AP</u>	! TRUSTEES UNIV PA	<u>20-200mCi/ML</u>	<u>A203665 001</u>	Feb 14, 2013
<u>AP</u>	! UCLA BIOMEDICAL	<u>4-40mCi/ML</u>	<u>A203801 001</u>	Oct 29, 2014
<u>AP</u>	UCSF RODIOPHARM	<u>20-300mCi/ML</u>	<u>A203811 001</u>	Jun 27, 2013
<u>AP</u>	UCSF RODIOPHARM	<u>20-300mCi/ML</u>	<u>A203902 001</u>	May 09, 2014
<u>AP</u>	UIHC PET IMAGING	<u>20-300mCi/ML</u>	<u>A203990 001</u>	Aug 06, 2014
<u>AP</u>	UNIV MICHIGAN	<u>20-300mCi/ML</u>	<u>A204531 001</u>	Jul 17, 2015
<u>AP</u>	UNIV SOUTHERN CA	<u>20-300mCi/ML</u>	<u>A209341 001</u>	Dec 16, 2020
<u>AP</u>	UNIV TX MD ANDERSON	<u>20-300mCi/ML</u>	<u>A203246 002</u>	Jan 13, 2014
<u>AP</u>	UNIV UTAH CYCLOTRON	<u>20-300mCi/ML</u>	<u>A204498 001</u>	Jun 23, 2015
<u>AP</u>	WISCONSIN	<u>20-500mCi/ML</u>	<u>A203709 001</u>	Oct 23, 2013
<u>AP</u>	WUSM CYCLOTRON	<u>20-300mCi/ML</u>	<u>A203935 001</u>	Feb 05, 2014
	HOT SHOTS NM LLC	4-500mCi/ML	A203937 001	Oct 30, 2014
	NORTHLAND	4-500mCi/ML	A203994 001	Feb 04, 2015
	NUKEMED	4-500mCi/ML	A203911 001	Apr 22, 2015
	UNIV TX MD ANDERSON	20-150mCi/ML	A203246 001	Jan 13, 2014

FLUDROCORTISONE ACETATE

TABLET; ORAL

FLUDROCORTISONE ACETATE

<u>AB</u>	BARR	<u>0.1MG</u>	<u>A040425 001</u>	Jan 21, 2003
<u>AB</u>	! IMPAX LABS	<u>0.1MG</u>	<u>A040431 001</u>	Mar 18, 2002

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

<u>AP</u>	BAXTER HLTHCARE CORP	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076755 002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076755 001</u>	Oct 12, 2004
<u>AP</u>	FRESENIUS KABI USA	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076955 002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076955 001</u>	Oct 12, 2004
<u>AP</u>	HIKMA	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076256 002</u>	Oct 12, 2004
<u>AP</u>		<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076787 002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076256 001</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076787 001</u>	Oct 12, 2004
<u>AP</u>	HIKMA FARMACEUTICA	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A078527 001</u>	Mar 23, 2009
<u>AP</u>	! RISING PHARMA	<u>1MG/10ML (0.1MG/ML)</u>	<u>A078527 002</u>	Mar 23, 2009
<u>AP</u>	SAGENT PHARMS	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A078595 001</u>	May 13, 2008
<u>AP</u>		<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A090584 001</u>	Aug 28, 2012
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A090584 002</u>	Aug 28, 2012

PRESCRIPTION DRUG PRODUCT LIST

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

<u>AP</u>	SANDOZ INC	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A077071</u>	<u>001</u>	May 03, 2005
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A077071</u>	<u>002</u>	May 03, 2005

FLUNISOLIDE

SPRAY, METERED; NASAL

FLUNISOLIDE

<u>AB</u>	!	BAUSCH AND LOMB	<u>0.025MG/SPRAY</u>	<u>A074805</u>	<u>001</u>	Feb 20, 2002
<u>AB</u>		RISING PHARMA	<u>0.025MG/SPRAY</u>	<u>A077704</u>	<u>001</u>	Aug 03, 2006
	!	APOTEX	0.029MG/SPRAY	A077436	001	Aug 09, 2007

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>		COSETTE	<u>0.01%</u>	<u>A089526</u>	<u>001</u>	Jul 26, 1988
<u>AT</u>			<u>0.025%</u>	<u>A210747</u>	<u>001</u>	Nov 05, 2018
<u>AT</u>		FOUGERA PHARMS INC	<u>0.01%</u>	<u>A088170</u>	<u>001</u>	Dec 16, 1982
<u>AT</u>			<u>0.025%</u>	<u>A088169</u>	<u>001</u>	Dec 16, 1982
<u>AT</u>		TARO	<u>0.025%</u>	<u>A087104</u>	<u>001</u>	Apr 27, 1982

SYNALAR

<u>AT</u>	+	!	MEDIMETRIKS PHARMS	<u>0.01%</u>	<u>N012787</u>	<u>004</u>
<u>AT</u>	+	!		<u>0.025%</u>	<u>N012787</u>	<u>002</u>
<u>AT</u>	+	!		<u>0.025%</u>	<u>N012787</u>	<u>005</u>

IMPLANT; INTRAVITREAL

ILUVIEN

	+	!	ALIMERA SCIENCES INC	0.19MG	N201923	001	Sep 26, 2014
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RETISERT

	+	!	BAUSCH AND LOMB	0.59MG	N021737	001	Apr 08, 2005
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YUTIQ

	+	!	EYEPOINT PHARMS	0.18MG	N210331	001	Oct 12, 2018
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OIL; TOPICAL

DERMA-SMOOTH/FS

<u>AT</u>	+	!	HILL DERMAC	<u>0.01%</u>	<u>N019452</u>	<u>001</u>	Feb 03, 1988
<u>AT</u>	+	!		<u>0.01%</u>	<u>N019452</u>	<u>002</u>	Nov 09, 2005

FLUCINOLONE ACETONIDE

<u>AT</u>		GLENMARK PHARMS LTD	<u>0.01%</u>	<u>A210556</u>	<u>001</u>	Oct 25, 2018
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FLUOCINOLONE ACETONIDE

<u>AT</u>		AMNEAL	<u>0.01%</u>	<u>A201759</u>	<u>001</u>	Oct 17, 2011
<u>AT</u>			<u>0.01%</u>	<u>A201764</u>	<u>001</u>	Oct 17, 2011
<u>AT</u>		LYNE	<u>0.01%</u>	<u>A090982</u>	<u>001</u>	Apr 25, 2016
<u>AT</u>			<u>0.01%</u>	<u>A203377</u>	<u>001</u>	Apr 25, 2016
<u>AT</u>		PADAGIS ISRAEL	<u>0.01%</u>	<u>A202847</u>	<u>001</u>	Aug 09, 2013
<u>AT</u>			<u>0.01%</u>	<u>A202848</u>	<u>001</u>	Aug 09, 2013
<u>AT</u>		QUAGEN	<u>0.01%</u>	<u>A212760</u>	<u>001</u>	Apr 02, 2021
<u>AT</u>			<u>0.01%</u>	<u>A212761</u>	<u>001</u>	Apr 02, 2021
<u>AT</u>		TARO	<u>0.01%</u>	<u>A202368</u>	<u>001</u>	May 19, 2016
<u>AT</u>			<u>0.01%</u>	<u>A209336</u>	<u>001</u>	May 19, 2016

FLUOCINONIDE ACETONIDE

<u>AT</u>		GLENMARK PHARMS LTD	<u>0.01%</u>	<u>A210539</u>	<u>001</u>	Oct 26, 2018
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OIL/DROPS; OTIC

DERMOTIC

<u>AT</u>	+	!	HILL DERMAC	<u>0.01%</u>	<u>N019452</u>	<u>003</u>	Nov 09, 2005
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FLAC

<u>AT</u>		ANDA REPOSITORY	<u>0.01%</u>	<u>A210736</u>	<u>001</u>	Apr 11, 2018
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FLUOCINOLONE ACETONIDE

<u>AT</u>		AMNEAL	<u>0.01%</u>	<u>A091306</u>	<u>001</u>	Oct 17, 2011
<u>AT</u>		LYNE	<u>0.01%</u>	<u>A203378</u>	<u>001</u>	Apr 25, 2016
<u>AT</u>		PADAGIS ISRAEL	<u>0.01%</u>	<u>A202849</u>	<u>001</u>	Jul 17, 2017
<u>AT</u>		QUAGEN	<u>0.01%</u>	<u>A212762</u>	<u>001</u>	Apr 02, 2021
<u>AT</u>		TASMAN PHARMA	<u>0.01%</u>	<u>A213264</u>	<u>001</u>	Feb 05, 2021

FLUOCINONIDE ACETONIDE

<u>AT</u>		GLENMARK PHARMS LTD	<u>0.01%</u>	<u>A211815</u>	<u>001</u>	Dec 14, 2018
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OINTMENT; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>		COSETTE	<u>0.025%</u>	<u>A089524</u>	<u>001</u>	Jul 26, 1988
<u>AT</u>		FOUGERA PHARMS INC	<u>0.025%</u>	<u>A088168</u>	<u>001</u>	Dec 16, 1982
<u>AT</u>		TARO	<u>0.025%</u>	<u>A040041</u>	<u>001</u>	Sep 15, 1994

SYNALAR

<u>AT</u>	+	!	MEDIMETRIKS PHARMS	<u>0.025%</u>	<u>N013960</u>	<u>001</u>
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PRESCRIPTION DRUG PRODUCT LIST

FLUOCINOLONE ACETONIDE

SHAMPOO; TOPICAL

CAPEX

+! GALDERMA LABS LP

0.01%

N020001 001 Aug 27, 1990

SOLUTION; TOPICAL

FLUOCINOLONE ACETONIDE**AT** ENCUBE ETHICALS0.01%A209913 001 Feb 13, 2019**AT** FOUGERA PHARMS INC0.01%A088167 001 Dec 16, 1982**AT** GLASSHOUSE PHARMS0.01%A209596 001 Dec 26, 2017**AT** LUPIN0.01%A206422 001 Sep 02, 2015**AT** TARO0.01%A089124 001 Sep 11, 1985SYNALAR**AT** +! MEDIMETRIKS PHARMS0.01%N015296 001FLUOCINOLONE 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** AMNEAL0.1%A211111 001 Jun 04, 2018**AB** CADILA0.1%A208989 001 Feb 10, 2020**AB** FOUGERA PHARMS INC0.1%A200735 001 Jul 14, 2014**AB** GLENMARK GENERICS0.1%A091282 001 Jul 14, 2014**AB** PADAGIS ISRAEL0.1%A090256 001 Jan 14, 2014**AB** TARO0.1%A200734 001 Jul 14, 2014**AB** TELIGENT0.1%A211758 001 Apr 03, 2019VANOS**AB** +! BAUSCH0.1%N021758 001 Feb 11, 2005FLUOCINONIDE**AB1** AMNEAL0.05%A210554 001 Aug 21, 2018**AB1** COSETTE0.05%A073085 001 Feb 14, 1992**AB1** FOUGERA PHARMS INC0.05%A073030 001 Oct 17, 1994**AB1** TARO0.05%A071500 001 Jun 10, 1987**AB1** +!0.05%N019117 001 Jun 26, 1984**AB1** TEVA0.05%A072488 001 Feb 06, 1989FLUOCINONIDE EMULSIFIED BASE**AB2** COSETTE0.05%A074204 001 Jun 13, 1995**AB2** FOUGERA PHARMS0.05%A076586 001 Jun 23, 2004**AB2** ! TARO PHARM INDS LTD0.05%A072494 001 Jan 19, 1989**AB2** TEVA0.05%A072490 001 Feb 07, 1989LIDEX-E**AB2** + ALVOGEN0.05%N016908 003

GEL; TOPICAL

FLUOCINONIDE**AB** + ALVOGEN0.05%N017373 001**AB** COSETTE0.05%A072537 001 Feb 07, 1989**AB** FOUGERA PHARMS INC0.05%A072933 001 Dec 30, 1994**AB** ! TARO0.05%A074935 001 Jul 29, 1997**AB** TELIGENT0.05%A209030 001 Jun 19, 2018

OINTMENT; TOPICAL

FLUOCINONIDE**AB** FOUGERA PHARMS0.05%A074905 001 Aug 26, 1997**AB** NOVEL LABS INC0.05%A207538 001 Jul 31, 2017**AB** ! TARO0.05%A075008 001 Jun 30, 1999**AB** TELIGENT0.05%A207680 001 Sep 28, 2018**AB** TEVA0.05%A073481 001 Dec 27, 1991**AB** XIROMED0.05%A212976 001 Nov 26, 2019LIDEX**AB** + ALVOGEN0.05%N016909 002

SOLUTION; TOPICAL

FLUOCINONIDE**AT** COSETTE0.05%A071535 001 Dec 02, 1988**AT** ENCUBE ETHICALS0.05%A209699 001 Nov 29, 2018**AT** FOUGERA PHARMS INC0.05%A072934 001 Feb 27, 1995**AT** GLASSHOUSE PHARMS0.05%A209118 001 Apr 23, 2018**AT** MACLEODS PHARMS LTD0.05%A209283 001 Apr 23, 2018

PRESCRIPTION DRUG PRODUCT LIST

FLUOCINONIDE

SOLUTION; TOPICAL

FLUOCINONIDE

<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>	TELLIGENT	<u>0.05%</u>	<u>A207554 001</u>	Mar 18, 2019
<u>AT</u>	ZYDUS PHARMS	<u>0.05%</u>	<u>A208948 001</u>	Jul 17, 2018

LIDEX

<u>AT</u>	+ ALVOGEN	<u>0.05%</u>	<u>N018849 001</u>	Apr 06, 1984
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FLUORESCEIN 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
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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	<u>EQ 500MG BASE/2ML (EQ 250MG BASE/ML)</u>	<u>N022186 002</u>	Aug 08, 2008

FLUORODOPA F-18

SOLUTION; INTRAVENOUS

FLUORODOPA F18

+!	FEINSTEIN	0.42-8.33mCi/ML	<u>N200655 001</u>	Oct 10, 2019
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FLUOROESTRADIOL F-18

SOLUTION; INTRAVENOUS

CERIANNA

+!	ZIONEXA	50ML (4-100mCi/ML)	<u>N212155 001</u>	May 20, 2020
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FLUOROMETHOLONE

OINTMENT; OPHTHALMIC

FML

+!	ALLERGAN	0.1%	<u>N017760 001</u>	Sep 04, 1985
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SUSPENSION/DROPS; OPHTHALMIC

FML

+!	ALLERGAN	0.1%	<u>N016851 002</u>	Jul 28, 1982
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FML FORTE

+!	ALLERGAN	0.25%	<u>N019216 001</u>	Apr 23, 1986
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FLUOROMETHOLONE ACETATE

SUSPENSION/DROPS; OPHTHALMIC

FLAREX

+!	EYEVANCE	0.1%	<u>N019079 001</u>	Feb 11, 1986
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FLUOROURACIL

CREAM; TOPICAL

EFUDEX

<u>AB</u>	+! BAUSCH	<u>5%</u>	<u>N016831 003</u>	
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FLUOROURACIL

<u>AB</u>	ACCORD HLTHCARE	<u>5%</u>	<u>A214845 001</u>	Oct 07, 2021
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<u>AB</u>	MAYNE PHARMA	<u>5%</u>	<u>A077524 001</u>	Apr 11, 2008
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<u>AB</u>	TARO	<u>5%</u>	<u>A090368 001</u>	Mar 05, 2010
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CARAC

+!	VALEANT PHARMS	0.5%	<u>N020985 001</u>	Oct 27, 2000
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NORTH

TOLAK

+!	HILL DERMACEUTICALS	4%	<u>N022259 001</u>	Sep 18, 2015
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INJECTABLE; INJECTION

FLUOROURACIL

<u>AP</u>	! ACCORD HLTHCARE	<u>500MG/10ML (50MG/ML)</u>	<u>A040743 002</u>	Apr 26, 2007
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<u>AP</u>	!	<u>1GM/20ML (50MG/ML)</u>	<u>A040743 001</u>	Apr 26, 2007
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<u>AP</u>	!	<u>2.5GM/50ML (50MG/ML)</u>	<u>A040798 002</u>	Apr 26, 2007
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<u>AP</u>	!	<u>5GM/100ML (50MG/ML)</u>	<u>A040798 001</u>	Apr 26, 2007
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<u>AP</u>	FRESENIUS KABI USA	<u>500MG/10ML (50MG/ML)</u>	<u>A040279 002</u>	Sep 30, 1998
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<u>AP</u>		<u>1GM/20ML (50MG/ML)</u>	<u>A040279 001</u>	Sep 30, 1998
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<u>AP</u>		<u>2.5GM/50ML (50MG/ML)</u>	<u>A040278 001</u>	Sep 30, 1998
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<u>AP</u>		<u>5GM/100ML (50MG/ML)</u>	<u>A040278 002</u>	Sep 30, 1998
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<u>AP</u>	GLAND PHARMA LTD	<u>500MG/10ML (50MG/ML)</u>	<u>A210123 001</u>	Oct 27, 2017
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<u>AP</u>		<u>1GM/20ML (50MG/ML)</u>	<u>A210123 002</u>	Oct 27, 2017
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<u>AP</u>		<u>2.5GM/50ML (50MG/ML)</u>	<u>A210124 001</u>	Dec 26, 2017
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<u>AP</u>		<u>5GM/100ML (50MG/ML)</u>	<u>A210124 002</u>	Dec 26, 2017
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<u>AP</u>	SAGENT PHARMS INC	<u>500MG/10ML (50MG/ML)</u>	<u>A203608 001</u>	May 11, 2017
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<u>AP</u>		<u>1GM/20ML (50MG/ML)</u>	<u>A203608 002</u>	May 11, 2017
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<u>AP</u>		<u>2.5GM/50ML (50MG/ML)</u>	<u>A203609 001</u>	Feb 17, 2016
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<u>AP</u>		<u>5GM/100ML (50MG/ML)</u>	<u>A203609 002</u>	Feb 17, 2016
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<u>AP</u>	TEVA PHARMS USA	<u>500MG/10ML (50MG/ML)</u>	<u>A040333 001</u>	Jan 27, 2000
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PRESCRIPTION DRUG PRODUCT LIST

FLUOROURACIL

SOLUTION;TOPICAL

EFUDEX

<u>AT</u>	<u>+!</u>	BAUSCH	<u>2%</u>	<u>N016831</u>	<u>001</u>	
<u>FLUOROURACIL</u>						
<u>AT</u>		TARO	<u>2%</u>	<u>A076526</u>	<u>001</u>	Nov 05, 2003
		!	<u>5%</u>	<u>A076526</u>	<u>002</u>	Nov 05, 2003

FLUOXETINE HYDROCHLORIDE

CAPSULE;ORAL

FLUOXETINE HYDROCHLORIDE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>EQ 40MG BASE</u>	<u>A090223</u>	<u>003</u>	Mar 19, 2009
<u>AB</u>		APNAR PHARMA LP	<u>EQ 40MG BASE</u>	<u>A075049</u>	<u>003</u>	Jan 29, 2002
<u>AB</u>		AUROBINDO PHARMA	<u>EQ 40MG BASE</u>	<u>A078619</u>	<u>003</u>	Jan 31, 2008
<u>AB</u>		CADILA PHARMS LTD	<u>EQ 40MG BASE</u>	<u>A206993</u>	<u>003</u>	May 23, 2019
<u>AB</u>		HERITAGE PHARMS INC	<u>EQ 40MG BASE</u>	<u>A201336</u>	<u>003</u>	Oct 01, 2012
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>EQ 40MG BASE</u>	<u>A075245</u>	<u>003</u>	Sep 28, 2004
<u>AB</u>		MARKSANS PHARMA	<u>EQ 40MG BASE</u>	<u>A075465</u>	<u>003</u>	Aug 02, 2001
<u>AB</u>		SCIEGEN PHARMS INC	<u>EQ 40MG BASE</u>	<u>A204597</u>	<u>003</u>	Mar 16, 2015
<u>AB</u>		SUN PHARM INDS LTD	<u>EQ 40MG BASE</u>	<u>A076990</u>	<u>001</u>	Dec 13, 2004
<u>AB</u>		TEVA	<u>EQ 40MG BASE</u>	<u>A075452</u>	<u>003</u>	Jan 29, 2002

PROZAC

<u>AB</u>	<u>+!</u>	ELI LILLY AND CO	<u>EQ 40MG BASE</u>	<u>N018936</u>	<u>003</u>	Jun 15, 1999
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FLUOXETINE HYDROCHLORIDE

<u>AB1</u>		ALEMBIC PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A090223</u>	<u>001</u>	Mar 19, 2009
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A090223</u>	<u>002</u>	Mar 19, 2009
<u>AB1</u>		APNAR PHARMA LP	<u>EQ 10MG BASE</u>	<u>A075049</u>	<u>001</u>	Aug 02, 2001
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A075049</u>	<u>002</u>	Jan 29, 2002
<u>AB1</u>		AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	<u>A078619</u>	<u>001</u>	Jan 31, 2008
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A078619</u>	<u>002</u>	Jan 31, 2008
<u>AB1</u>		CADILA PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A206993</u>	<u>001</u>	May 23, 2019
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A206993</u>	<u>002</u>	May 23, 2019
<u>AB1</u>		HERITAGE PHARMS INC	<u>EQ 10MG BASE</u>	<u>A201336</u>	<u>001</u>	Oct 01, 2012
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A201336</u>	<u>002</u>	Oct 01, 2012
<u>AB1</u>		IVAX SUB TEVA PHARMS	<u>EQ 10MG BASE</u>	<u>A075245</u>	<u>002</u>	Jan 31, 2002
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A075245</u>	<u>001</u>	Jan 31, 2002
<u>AB1</u>		LANDELA PHARM	<u>EQ 10MG BASE</u>	<u>A075464</u>	<u>001</u>	Jan 30, 2002
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A075464</u>	<u>002</u>	Jan 30, 2002
<u>AB1</u>		MARKSANS PHARMA	<u>EQ 10MG BASE</u>	<u>A075465</u>	<u>001</u>	Jan 29, 2002
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A075465</u>	<u>002</u>	Jan 29, 2002
<u>AB1</u>		SCIEGEN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A204597</u>	<u>001</u>	Mar 16, 2015
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A204597</u>	<u>002</u>	Mar 16, 2015
<u>AB1</u>		TEVA	<u>EQ 10MG BASE</u>	<u>A075452</u>	<u>001</u>	Jan 29, 2002
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A075452</u>	<u>002</u>	Jan 29, 2002
<u>AB1</u>		TEVA PHARMS USA	<u>EQ 10MG BASE</u>	<u>A076001</u>	<u>001</u>	Jan 29, 2002
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A076001</u>	<u>002</u>	Jan 29, 2002

PROZAC

<u>AB1</u>	<u>+</u>	ELI LILLY AND CO	<u>EQ 10MG BASE</u>	<u>N018936</u>	<u>006</u>	Dec 23, 1992
<u>AB1</u>	<u>+</u>		<u>EQ 20MG BASE</u>	<u>N018936</u>	<u>001</u>	Dec 29, 1987

CAPSULE, DELAYED REL PELLETS;ORAL

FLUOXETINE HYDROCHLORIDE

!		DR REDDYS LABS LTD	<u>EQ 90MG BASE</u>	<u>A078572</u>	<u>001</u>	Mar 22, 2010
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SOLUTION;ORAL

FLUOXETINE HYDROCHLORIDE

<u>AA</u>		AUROBINDO PHARMA LTD	<u>EQ 20MG BASE/5ML</u>	<u>A079209</u>	<u>001</u>	Mar 20, 2009
<u>AA</u>		LANNETT CO INC	<u>EQ 20MG BASE/5ML</u>	<u>A077849</u>	<u>001</u>	Feb 09, 2007
<u>AA</u>	<u>!</u>	PHARM ASSOC	<u>EQ 20MG BASE/5ML</u>	<u>A076015</u>	<u>001</u>	Jan 30, 2002
<u>AA</u>		TEVA	<u>EQ 20MG BASE/5ML</u>	<u>A075506</u>	<u>001</u>	Aug 02, 2001
<u>AA</u>		WOCKHARDT BIO AG	<u>EQ 20MG BASE/5ML</u>	<u>A075514</u>	<u>001</u>	Aug 29, 2002

TABLET;ORAL

FLUOXETINE HYDROCHLORIDE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A208698</u>	<u>001</u>	Apr 05, 2017
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A208698</u>	<u>002</u>	Apr 05, 2017
<u>AB</u>	<u>+!</u>	ALMATICA	<u>EQ 60MG BASE</u>	<u>N202133</u>	<u>001</u>	Oct 06, 2011
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 10MG BASE</u>	<u>A213286</u>	<u>001</u>	Apr 08, 2020
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A213286</u>	<u>002</u>	Apr 08, 2020
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A213265</u>	<u>001</u>	Jun 10, 2020
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 10MG BASE</u>	<u>A076006</u>	<u>001</u>	Jan 30, 2002
<u>AB</u>	<u>!</u>		<u>EQ 20MG BASE</u>	<u>A076006</u>	<u>002</u>	Apr 23, 2018
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A211721</u>	<u>001</u>	Jan 25, 2019

PRESCRIPTION DRUG PRODUCT LIST

FLUOXETINE HYDROCHLORIDE

TABLET; ORAL

FLUOXETINE HYDROCHLORIDE

<u>AB</u>	INVENTIA HLTHCARE	<u>EQ 60MG BASE</u>	<u>A209695 001</u>	Nov 20, 2017
<u>AB</u>	LUPIN LTD	<u>EQ 10MG BASE</u>	<u>A211653 001</u>	Apr 15, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A211653 002</u>	Apr 15, 2019
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A211632 001</u>	Feb 08, 2019
<u>AB</u>	PAR FORM	<u>EQ 10MG BASE</u>	<u>A203836 001</u>	Aug 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A203836 002</u>	Aug 19, 2016
<u>AB</u>	PAR PHARM INC	<u>EQ 60MG BASE</u>	<u>A209419 001</u>	Nov 16, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A210935 001</u>	Mar 20, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A210935 002</u>	Mar 20, 2019
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A211282 001</u>	Jan 10, 2019
<u>AB</u>	TARO	<u>EQ 60MG BASE</u>	<u>A211477 001</u>	Nov 21, 2018
<u>AB</u>	TEVA	<u>EQ 10MG BASE</u>	<u>A075872 001</u>	Jan 29, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A075872 002</u>	Jan 04, 2019
<u>AB</u>	TEVA PHARMS USA	<u>EQ 60MG BASE</u>	<u>A211051 001</u>	Dec 03, 2018
<u>AB</u>	UPSHER SMITH LABS	<u>EQ 10MG BASE</u>	<u>A211696 001</u>	Jan 30, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A211696 002</u>	Jan 30, 2019
	TORRENT	EQ 10MG BASE	A206937 001	Oct 21, 2016
!		EQ 20MG BASE	A206937 002	Oct 21, 2016

FLUOXETINE HYDROCHLORIDE; OLANZAPINE

CAPSULE; ORAL

OLANZAPINE AND FLUOXETINE HYDROCHLORIDE

<u>AB</u>	PAR PHARM	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>A077742 001</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>A077742 002</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>A077742 003</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>A077742 004</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>A077742 005</u>	Nov 02, 2012
<u>AB</u>	SANDOZ	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>A078901 005</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>A078901 001</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>A078901 003</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>A078901 002</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>A078901 004</u>	Nov 16, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>A202074 001</u>	Mar 25, 2013
<u>AB</u>		<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>A077528 001</u>	Jun 19, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>A077528 002</u>	Jun 19, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>A077528 003</u>	Jun 19, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>A077528 004</u>	Jun 19, 2012
<u>SYMBYAX</u>				
<u>AB</u>	+ LILLY	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>N021520 001</u>	Apr 09, 2007
<u>AB</u>	+	<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>N021520 002</u>	Dec 24, 2003
<u>AB</u>	+	<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>N021520 004</u>	Dec 24, 2003
<u>AB</u>	+!	<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>N021520 003</u>	Dec 24, 2003
<u>AB</u>	+	<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>N021520 005</u>	Dec 24, 2003

FLUPHENAZINE DECANOATE

INJECTABLE; INJECTION

FLUPHENAZINE DECANOATE

<u>AO</u>	EUGIA PHARMA	<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>	HIKMA	<u>25MG/ML</u>	<u>A074531 001</u>	Aug 30, 1996
<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

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>	AMNEAL	<u>1MG</u>	<u>A213647 001</u>	Jul 09, 2020
<u>AB</u>		<u>2.5MG</u>	<u>A213647 002</u>	Jul 09, 2020
<u>AB</u>		<u>5MG</u>	<u>A213647 003</u>	Jul 09, 2020
<u>AB</u>		<u>10MG</u>	<u>A213647 004</u>	Jul 09, 2020
<u>AB</u>	CEROVENE INC	<u>1MG</u>	<u>A214534 001</u>	Jan 07, 2021

PRESCRIPTION DRUG PRODUCT LIST

FLUPHENAZINE HYDROCHLORIDE

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

<u>AB</u>		<u>2.5MG</u>	<u>A214534 002</u>	Jan 07, 2021
<u>AB</u>		<u>5MG</u>	<u>A214534 003</u>	Jan 07, 2021
<u>AB</u>		<u>10MG</u>	<u>A214534 004</u>	Jan 07, 2021
<u>AB</u>	ENALTEC	<u>1MG</u>	<u>A215141 001</u>	Oct 20, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A215141 002</u>	Oct 20, 2021
<u>AB</u>		<u>5MG</u>	<u>A215141 003</u>	Oct 20, 2021
<u>AB</u>		<u>10MG</u>	<u>A215141 004</u>	Oct 20, 2021
<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>	NOVITIUM PHARMA	<u>1MG</u>	<u>A214674 001</u>	Mar 01, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A214674 002</u>	Mar 01, 2021
<u>AB</u>		<u>5MG</u>	<u>A214674 003</u>	Mar 01, 2021
<u>AB</u>		<u>10MG</u>	<u>A214674 004</u>	Mar 01, 2021
<u>AB</u>	ZYDUS	<u>1MG</u>	<u>A214552 001</u>	May 27, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A214552 002</u>	May 27, 2021
<u>AB</u>		<u>5MG</u>	<u>A214552 003</u>	May 27, 2021
<u>AB</u>		<u>10MG</u>	<u>A214552 004</u>	May 27, 2021

FLURANDRENOLIDE

CREAM; TOPICAL

CORDRAN SP

<u>AT</u>	+!	ALMIRALL	<u>0.05%</u>	<u>N012806 002</u>
<u>AT</u>		CINTEX SVCS	<u>0.05%</u>	<u>A205342 001</u> Apr 13, 2016

LOTION; TOPICAL

CORDRAN

<u>AT</u>	+!	ALMIRALL	<u>0.05%</u>	<u>N013790 001</u>
<u>AT</u>		CINTEX SVCS	<u>0.05%</u>	<u>A205343 001</u> Dec 22, 2016
<u>AT</u>		PADAGIS ISRAEL	<u>0.05%</u>	<u>A207133 001</u> Aug 30, 2016

OINTMENT; TOPICAL

CORDRAN

<u>AT</u>	+!	ALMIRALL	<u>0.05%</u>	<u>N012806 001</u>
<u>AT</u>		TELGENT	<u>0.05%</u>	<u>A207851 001</u> Dec 30, 2016

TAPE; TOPICAL

CORDRAN

	+!	ALMIRALL	0.004MG/SQ CM	N016455 001
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FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL

FLURAZEPAM HYDROCHLORIDE

		MYLAN PHARMS INC	15MG	A070345 002 Nov 27, 1985
	!		30MG	A070345 001 Nov 27, 1985

FLURBIPROFEN

TABLET; ORAL

FLURBIPROFEN

	!	TEVA	100MG	A074431 001 May 31, 1995
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FLURBIPROFEN SODIUM

SOLUTION/DROPS; OPHTHALMIC

FLURBIPROFEN SODIUM

	!	BAUSCH AND LOMB	0.03%	A074447 001 Jan 04, 1995
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FLUTAMIDE

CAPSULE; ORAL

FLUTAMIDE

	!	WAYLIS THERAP	125MG	A075298 001 Sep 18, 2001
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FLUTEMETAMOL F-18

INJECTABLE; INTRAVENOUS

VIZAMYL

	+!	GE HEALTHCARE	121.5mCi/30ML (4.05mCi/ML)	N203137 002 Oct 25, 2013
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PRESCRIPTION DRUG PRODUCT LIST

FLUTICASONE FUROATE

POWDER; INHALATION

ARNUTTY 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
+	!		0.2MG/INH;EQ 0.0625MG BASE/INH;EQ 0.025MG BASE/INH	N209482	002	Sep 09, 2020

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

AB		COSETTE	0.05%	A077055	001	Jun 30, 2006
AB		FOUGERA PHARMS	0.05%	A076451	001	May 14, 2004
AB	!	PADAGIS ISRAEL	0.05%	A076793	001	May 14, 2004

LOTION; TOPICAL

CUTIVATE

AB	+	!	FOUGERA PHARMS	0.05%	N021152	001	Mar 31, 2005
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FLUTICASONE PROPIONATE

AB		GLENMARK GENERICS	0.05%	A090759	001	May 02, 2011
AB		PADAGIS ISRAEL	0.05%	A091553	001	Jul 30, 2013

OINTMENT; TOPICAL

FLUTICASONE PROPIONATE

AB		COSETTE	0.005%	A077168	001	Mar 03, 2006
AB	!	PADAGIS ISRAEL	0.005%	A076668	001	May 14, 2004

POWDER; INHALATION

ARMONAIR DIGIHALER

+		TEVA PHARM	0.055MG/INH	N208798	004	Feb 20, 2020
+			0.113MG/INH	N208798	005	Feb 20, 2020
+	!		0.232MG/INH	N208798	006	Feb 20, 2020

FLOVENT DISKUS 100

+	!	GLAXO GRP LTD	0.1MG/INH	N020833	002	Sep 29, 2000
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FLOVENT DISKUS 250

+	!	GLAXO GRP LTD	0.25MG/INH	N020833	003	Sep 29, 2000
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FLOVENT DISKUS 50

+	!	GLAXO GRP LTD	0.05MG/INH	N020833	001	Sep 29, 2000
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SPRAY, METERED; NASAL

FLUTICASONE PROPIONATE

AB		AKORN	0.05MG/SPRAY	A077570	001	Jan 16, 2008
AB		APOTEX INC	0.05MG/SPRAY	A077538	001	Sep 12, 2007
AB	!	HIKMA	0.05MG/SPRAY	A076504	001	Feb 22, 2006
AB		WOCKHARDT BIO AG	0.05MG/SPRAY	A078492	001	Jan 09, 2012

XHANCE

+	!	OPTINOSE US INC	0.093MG	N209022	001	Sep 18, 2017
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FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE

AEROSOL, METERED; INHALATION

ADVAIR 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

ADVAIR DISKUS 100/50

AB	+	!	GLAXO GRP LTD	0.1MG/INH;EQ 0.05MG BASE/INH	N021077	001	Aug 24, 2000
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ADVAIR DISKUS 250/50

AB	+	!	GLAXO GRP LTD	0.25MG/INH;EQ 0.05MG BASE/INH	N021077	002	Aug 24, 2000
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PRESCRIPTION DRUG PRODUCT LIST

FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE

POWDER; INHALATION

ADVAIR DISKUS 500/50

AB	+!	GLAXO GRP LTD	<u>0.5MG/INH;EQ 0.05MG BASE/INH</u>	<u>N021077 003</u>	Aug 24, 2000
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FLUTICASONE PROPIONATE AND SALMETEROL XINAFOATE

AB		HIKMA	<u>0.1MG/INH;EQ 0.05MG BASE/INH</u>	<u>A203433 001</u>	Dec 17, 2020
AB			<u>0.25MG/INH;EQ 0.05MG BASE/INH</u>	<u>A203433 002</u>	Dec 17, 2020
AB		TEVA PHARMS USA	<u>0.1MG/INH;EQ 0.05MG BASE/INH</u>	<u>A213948 001</u>	Dec 13, 2021
AB			<u>0.25MG/INH;EQ 0.05MG BASE/INH</u>	<u>A213948 002</u>	Dec 13, 2021
AB			<u>0.5MG/INH;EQ 0.05MG BASE/INH</u>	<u>A213948 003</u>	Dec 13, 2021

WIXELA INHUB

AB		MYLAN	<u>0.1MG/INH;EQ 0.05MG BASE/INH</u>	<u>A208891 001</u>	Jan 30, 2019
AB			<u>0.25MG/INH;EQ 0.05MG BASE/INH</u>	<u>A208891 002</u>	Jan 30, 2019
AB			<u>0.5MG/INH;EQ 0.05MG BASE/INH</u>	<u>A208891 003</u>	Jan 30, 2019
		AIRDUO DIGIHALER			
	+	TEVA PHARM	0.055MG/INH;EQ 0.014MG BASE/INH	N208799 004	Jul 12, 2019
	+		0.113MG/INH;EQ 0.014MG BASE/INH	N208799 005	Jul 12, 2019
	+		0.232MG/INH;EQ 0.014MG BASE/INH	N208799 006	Jul 12, 2019
		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/INH;EQ 0.014MG BASE/INH	N208799 003	Jan 27, 2017

FLUVASTATIN SODIUM

CAPSULE; ORAL

FLUVASTATIN SODIUM

AB		MYLAN PHARMS INC	<u>EQ 20MG BASE</u>	<u>A090595 001</u>	Apr 11, 2012
AB	!		<u>EQ 40MG BASE</u>	<u>A090595 002</u>	Apr 11, 2012
AB		TEVA PHARMS	<u>EQ 20MG BASE</u>	<u>A078407 001</u>	Jun 12, 2012
AB			<u>EQ 40MG BASE</u>	<u>A078407 002</u>	Jun 12, 2012

TABLET, EXTENDED RELEASE; ORAL

FLUVASTATIN SODIUM

AB		BEIJING	<u>EQ 80MG BASE</u>	<u>A209397 001</u>	Apr 26, 2021
AB		TEVA PHARMS USA	<u>EQ 80MG BASE</u>	<u>A079011 001</u>	Jan 27, 2016

LESCOL XL

AB	+!	NOVARTIS	<u>EQ 80MG BASE</u>	<u>N021192 001</u>	Oct 06, 2000
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FLUVOXAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

FLUVOXAMINE MALEATE

AB		ACTAVIS ELIZABETH	<u>100MG</u>	<u>A091482 001</u>	Apr 23, 2013
AB	!		<u>150MG</u>	<u>A091482 002</u>	Nov 18, 2013
AB		ANCHEN PHARMS	<u>100MG</u>	<u>A091476 001</u>	Mar 13, 2013
AB			<u>150MG</u>	<u>A091476 002</u>	Mar 13, 2013
AB		BIONPHARMA INC	<u>150MG</u>	<u>A212182 001</u>	May 11, 2020

TABLET; ORAL

FLUVOXAMINE MALEATE

AB		APOTEX	<u>25MG</u>	<u>A075902 001</u>	May 07, 2001
AB			<u>50MG</u>	<u>A075902 002</u>	May 07, 2001
AB			<u>100MG</u>	<u>A075902 003</u>	May 07, 2001
AB		UPSHER SMITH LABS	<u>25MG</u>	<u>A075888 001</u>	Nov 29, 2000
AB			<u>50MG</u>	<u>A075888 002</u>	Nov 29, 2000
AB	!		<u>100MG</u>	<u>A075888 003</u>	Nov 29, 2000

LUVOX

AB		ANI PHARMS	<u>25MG</u>	<u>N021519 001</u>	Dec 20, 2007
AB			<u>50MG</u>	<u>N021519 002</u>	Dec 20, 2007
AB			<u>100MG</u>	<u>N021519 003</u>	Dec 20, 2007

FOLIC ACID

INJECTABLE; INJECTION

FOLIC ACID

AP	!	FRESENIUS KABI USA	<u>5MG/ML</u>	<u>A089202 001</u>	Feb 18, 1986
AP		XGEN PHARMS	<u>5MG/ML</u>	<u>A202522 001</u>	Nov 06, 2019

TABLET; ORAL

FOLIC ACID

AA	!	AMNEAL PHARM	<u>1MG</u>	<u>A040625 001</u>	Jul 21, 2005
AA		ATHEM	<u>1MG</u>	<u>A211064 001</u>	Mar 08, 2019
AA		CADILA PHARMS LTD	<u>1MG</u>	<u>A202437 001</u>	Jan 27, 2014
AA		CHARTWELL MOLECULAR	<u>1MG</u>	<u>A090035 001</u>	Jun 09, 2009
AA		LEADING PHARMA LLC	<u>1MG</u>	<u>A040796 001</u>	Jan 12, 2009
AA		NUVO PHARMS INC	<u>1MG</u>	<u>A204418 001</u>	Jul 28, 2015
AA		QINGDAO BAHEAL PHARM	<u>1MG</u>	<u>A091145 001</u>	Jul 12, 2013
AA	+!	WATSON LABS	<u>1MG</u>	<u>A080680 001</u>	

PRESCRIPTION DRUG PRODUCT LIST

FOMEPIZOLE

INJECTABLE; INJECTION

FOMEPIZOLE

<u>AP</u>	AM REGENT	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078368 001</u>	Dec 14, 2007
<u>AP</u>	! MYLAN INSTITUTIONAL	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078639 001</u>	Mar 03, 2008
<u>AP</u>	NAVINTA LLC	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078537 001</u>	Mar 06, 2008

FONDAPARINUX SODIUM

INJECTABLE; SUBCUTANEOUS

ARIXTRA

<u>AP</u>	+! MYLAN IRELAND LTD	<u>2.5MG/0.5ML</u>	<u>N021345 001</u>	Dec 07, 2001
<u>AP</u>	+!	<u>5MG/0.4ML</u>	<u>N021345 002</u>	May 28, 2004
<u>AP</u>	+!	<u>7.5MG/0.6ML</u>	<u>N021345 003</u>	May 28, 2004
<u>AP</u>	+!	<u>10MG/0.8ML</u>	<u>N021345 004</u>	May 28, 2004

FONDAPARINUX SODIUM

<u>AP</u>	DR REDDYS LABS LTD	<u>2.5MG/0.5ML</u>	<u>A091316 001</u>	Jul 11, 2011
<u>AP</u>		<u>5MG/0.4ML</u>	<u>A091316 002</u>	Jul 11, 2011
<u>AP</u>		<u>7.5MG/0.6ML</u>	<u>A091316 003</u>	Jul 11, 2011
<u>AP</u>		<u>10MG/0.8ML</u>	<u>A091316 004</u>	Jul 11, 2011
<u>AP</u>	EUGIA PHARMA	<u>2.5MG/0.5ML</u>	<u>A206918 001</u>	Dec 26, 2017
<u>AP</u>		<u>5MG/0.4ML</u>	<u>A206918 002</u>	Dec 26, 2017
<u>AP</u>		<u>7.5MG/0.6ML</u>	<u>A206918 003</u>	Dec 26, 2017
<u>AP</u>		<u>10MG/0.8ML</u>	<u>A206918 004</u>	Dec 26, 2017
<u>AP</u>	JIANGSU PHARMS	<u>2.5MG/0.5ML</u>	<u>A206812 001</u>	May 15, 2018
<u>AP</u>		<u>5MG/0.4ML</u>	<u>A206812 002</u>	May 15, 2018
<u>AP</u>		<u>7.5MG/0.6ML</u>	<u>A206812 003</u>	May 15, 2018
<u>AP</u>		<u>10MG/0.8ML</u>	<u>A206812 004</u>	May 15, 2018
<u>AP</u>	SCINOPHARM TAIWAN	<u>2.5MG/0.5ML</u>	<u>A208615 001</u>	Nov 14, 2018
<u>AP</u>		<u>5MG/0.4ML</u>	<u>A208615 002</u>	Nov 14, 2018
<u>AP</u>		<u>7.5MG/0.6ML</u>	<u>A208615 003</u>	Nov 14, 2018
<u>AP</u>		<u>10MG/0.8ML</u>	<u>A208615 004</u>	Nov 14, 2018

FORMOTEROL FUMARATE

SOLUTION; INHALATION

FORMOTEROL FUMARATE

<u>AN</u>	ALEMBIC PHARMS LTD	<u>0.02MG/2ML</u>	<u>A215078 001</u>	Nov 22, 2021
<u>AN</u>	TEVA PHARMS USA INC	<u>0.02MG/2ML</u>	<u>A091141 001</u>	Jun 22, 2021
<u>AN</u>	+! MYLAN SPECLT	<u>0.02MG/2ML</u>	<u>N022007 001</u>	May 11, 2007

FORMOTEROL FUMARATE; GLYCOPYRROLATE

AEROSOL, METERED; INHALATION

BEVESPI AEROSPHERE

+!	ASTRAZENECA	0.0048MG/INH;0.0090MG/INH	N208294 001	Apr 25, 2016
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FORMOTEROL FUMARATE; MOMETASONE FUROATE

AEROSOL, METERED; INHALATION

DULERA

+	ORGANON	0.005MG/INH;0.05MG/INH	N022518 003	Aug 12, 2019
+!		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 HLTHCARE	EQ 50MG BASE/ML	N022116 001	Jun 14, 2007
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TABLET; ORAL

FOSAMPRENAVIR CALCIUM

<u>AB</u>	MYLAN	<u>EQ 700MG BASE</u>	<u>A204060 001</u>	Apr 15, 2016
<u>AB</u>	SUN PHARM	<u>EQ 700MG BASE</u>	<u>A204024 001</u>	Nov 20, 2019
<u>AB</u>	+! VIIV HLTHCARE	<u>EQ 700MG BASE</u>	<u>N021548 001</u>	Oct 20, 2003

FOSAPREPITANT DIMEGLUMINE

POWDER; INTRAVENOUS

EMEND

<u>AP</u>	+! MERCK AND CO INC	<u>EQ 150MG BASE/VIAL</u>	<u>N022023 002</u>	Nov 12, 2010
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FOSAPREPITANT DIMEGLUMINE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 150MG BASE/VIAL</u>	<u>A204025 001</u>	Aug 26, 2020
<u>AP</u>	APOTEX	<u>EQ 150MG BASE/VIAL</u>	<u>A205020 001</u>	Sep 05, 2019
<u>AP</u>	ASPIRO	<u>EQ 150MG BASE/VIAL</u>	<u>A214616 001</u>	Jul 29, 2021
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 150MG BASE/VIAL</u>	<u>A211860 001</u>	Sep 05, 2019
<u>AP</u>	BE PHARMS	<u>EQ 150MG BASE/VIAL</u>	<u>A212309 001</u>	Sep 05, 2019
<u>AP</u>	CHIA TAI TIANQING	<u>EQ 150MG BASE/VIAL</u>	<u>A212143 001</u>	Mar 03, 2021
<u>AP</u>	DR REDDYS	<u>EQ 150MG BASE/VIAL</u>	<u>A211160 001</u>	Dec 09, 2020

PRESCRIPTION DRUG PRODUCT LIST

FOSAPREPITANT DIMEGLUMINE

POWDER; INTRAVENOUS

FOSAPREPITANT DIMEGLUMINE

<u>AP</u>	EUGIA PHARMA	<u>EQ 150MG BASE/VIAL</u>	<u>A210625 001</u>	Jan 12, 2021
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 150MG BASE/VIAL</u>	<u>A206197 001</u>	Jun 09, 2016
<u>AP</u>	LUPIN LTD	<u>EQ 150MG BASE/VIAL</u>	<u>A210689 001</u>	Sep 05, 2019
<u>AP</u>	MSN	<u>EQ 150MG BASE/VIAL</u>	<u>A209965 001</u>	Sep 05, 2019
<u>AP</u>	MYLAN LABS LTD	<u>EQ 150MG BASE/VIAL</u>	<u>A204015 002</u>	Sep 05, 2019
<u>AP</u>	NAVINTA LLC	<u>EQ 150MG BASE/VIAL</u>	<u>A212957 002</u>	Aug 20, 2020
<u>AP</u>	PRAXGEN	<u>EQ 150MG BASE/VIAL</u>	<u>A211624 001</u>	Sep 05, 2019
<u>AP</u>		<u>EQ 150MG BASE/VIAL</u>	<u>A213199 001</u>	Oct 04, 2021
<u>AP</u>	QILU PHARM HAINAN	<u>EQ 150MG BASE/VIAL</u>	<u>A213106 001</u>	Sep 08, 2020
<u>AP</u>	SANDOZ INC	<u>EQ 150MG BASE/VIAL</u>	<u>A203939 001</u>	Dec 08, 2020
	TEVA PHARMS USA	EQ 150MG BASE/VIAL	N210064 001	Sep 05, 2019

FOSCARNET SODIUM

INJECTABLE; INJECTION

FOSCARNET SODIUM

<u>AP</u>	FRESENIUS KABI USA	<u>2.4GM/100ML</u>	<u>A212483 001</u>	Jan 29, 2021
<u>AP</u>	GLAND PHARMA LTD	<u>2.4GM/100ML</u>	<u>A213001 001</u>	Apr 21, 2021

FOSCAVIR

<u>AP</u>	+!	CLINIGEN HLTHCARE	<u>2.4GM/100ML</u>	<u>N020068 001</u>	Sep 27, 1991
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FOSDENOPTERIN HYDROBROMIDE

POWDER; INTRAVENOUS

NULIBRY

	+!	ORIGIN	EQ 9.5MG BASE/VIAL	N214018 001	Feb 26, 2021
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FOSFOMYCIN TROMETHAMINE

FOR SOLUTION; ORAL

FOSFOMYCIN TROMETHAMINE

<u>AA</u>	ALKEM LABS LTD	<u>EQ 3GM BASE/PACKET</u>	<u>A214554 001</u>	Oct 21, 2021
<u>AA</u>	XIROMED	<u>EQ 3GM BASE/PACKET</u>	<u>A212548 001</u>	Oct 06, 2020

MONUROL

<u>AA</u>	+!	ZAMBON SPA	<u>EQ 3GM BASE/PACKET</u>	<u>N050717 001</u>	Dec 19, 1996
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FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

<u>AB</u>	APOTEX INC	<u>10MG</u>	<u>A076906 001</u>	May 17, 2005
<u>AB</u>		<u>20MG</u>	<u>A076906 002</u>	May 17, 2005
<u>AB</u>		<u>40MG</u>	<u>A076906 003</u>	May 17, 2005
<u>AB</u>	AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A091163 001</u>	Mar 30, 2011
<u>AB</u>		<u>20MG</u>	<u>A091163 002</u>	Mar 30, 2011
<u>AB</u>		<u>40MG</u>	<u>A091163 003</u>	Mar 30, 2011
<u>AB</u>	CHARTWELL RX	<u>10MG</u>	<u>A076483 001</u>	Apr 23, 2004
<u>AB</u>		<u>20MG</u>	<u>A076483 002</u>	Apr 23, 2004
<u>AB</u>		<u>40MG</u>	<u>A076483 003</u>	Apr 23, 2004
<u>AB</u>	INVAGEN PHARMS	<u>10MG</u>	<u>A077222 001</u>	Apr 20, 2005
<u>AB</u>		<u>20MG</u>	<u>A077222 002</u>	Apr 20, 2005
<u>AB</u>		<u>40MG</u>	<u>A077222 003</u>	Apr 20, 2005
<u>AB</u>	PRINSTON INC	<u>10MG</u>	<u>A205670 001</u>	Aug 29, 2016
<u>AB</u>		<u>20MG</u>	<u>A205670 002</u>	Aug 29, 2016
<u>AB</u>		<u>40MG</u>	<u>A205670 003</u>	Aug 29, 2016
<u>AB</u>	TEVA	<u>10MG</u>	<u>A076139 001</u>	Nov 25, 2003
<u>AB</u>		<u>20MG</u>	<u>A076139 002</u>	Nov 25, 2003
<u>AB</u>	!	<u>40MG</u>	<u>A076139 003</u>	Nov 25, 2003

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

<u>AB</u>	AUROBINDO PHARMA	<u>10MG;12.5MG</u>	<u>A079245 001</u>	Jul 09, 2009
<u>AB</u>	!	<u>20MG;12.5MG</u>	<u>A079245 002</u>	Jul 09, 2009
<u>AB</u>	INVAGEN PHARMS	<u>10MG;12.5MG</u>	<u>A090228 001</u>	Jul 09, 2009
<u>AB</u>		<u>20MG;12.5MG</u>	<u>A090228 002</u>	Jul 09, 2009
<u>AB</u>	SANDOZ	<u>10MG;12.5MG</u>	<u>A076961 001</u>	Sep 28, 2005
<u>AB</u>		<u>20MG;12.5MG</u>	<u>A076961 002</u>	Sep 28, 2005

FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE

POWDER; INTRAVENOUS

AKYNZEO

	+!	HELSINN HLTHCARE	EQ 235MG BASE/VIAL;EQ 0.25MG BASE/VIAL	N210493 001	Apr 19, 2018
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PRESCRIPTION DRUG PRODUCT LIST

FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE

SOLUTION; INTRAVENOUS

AKYNZEO

+	!	HELINN HLTHCARE	EQ 235MG BASE/20ML (EQ 11.75MG BASE/ML); EQ 0.25MG BASE/20ML (EQ 0.0125MG BASE/ML)	N210493	002	May 27, 2020
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FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

CEREBYX

AP	+	!	PARKE DAVIS	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>N020450</u>	<u>001</u>	Aug 05, 1996
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FOSPHENYTOIN SODIUM

AP			FRESENIUS KABI USA	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078052</u>	<u>001</u>	Aug 06, 2007
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AP			HIKMA	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A077481</u>	<u>001</u>	Aug 06, 2007
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AP				<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A077989</u>	<u>001</u>	Aug 06, 2007
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AP			HIKMA FARMACEUTICA	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078765</u>	<u>001</u>	Dec 02, 2009
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AP			MYLAN LABS LTD	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078736</u>	<u>001</u>	Jun 08, 2010
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AP			SUN PHARM	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078417</u>	<u>001</u>	Mar 18, 2008
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AP			WOCKHARDT	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078137</u>	<u>001</u>	Aug 06, 2007
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FOSTAMATINIB DISODIUM

TABLET; ORAL

TAVALISSE

+			RIGEL PHARMS INC	EQ 100MG BASE	N209299	001	Apr 17, 2018
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+	!			EQ 150MG BASE	N209299	002	Apr 17, 2018
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FOSTEMSAVIR TROMETHAMINE

TABLET, EXTENDED RELEASE; ORAL

RUKOBIA

+	!		VIIV HLTHCARE	EQ 600MG BASE	N212950	001	Jul 02, 2020
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FROVATRIPTAN SUCCINATE

TABLET; ORAL

FROVA

AB	+	!	ENDO PHARMS	<u>EQ 2.5MG BASE</u>	<u>N021006</u>	<u>001</u>	Nov 08, 2001
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FROVATRIPTAN SUCCINATE

AB			AMNEAL PHARMS CO	<u>EQ 2.5MG BASE</u>	<u>A211292</u>	<u>001</u>	Nov 06, 2018
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AB			GLENMARK PHARMS LTD	<u>EQ 2.5MG BASE</u>	<u>A204730</u>	<u>001</u>	Mar 11, 2016
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AB			MYLAN	<u>EQ 2.5MG BASE</u>	<u>A202931</u>	<u>001</u>	Aug 28, 2014
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FULVESTRANT

INJECTABLE; INTRAMUSCULAR

FASLODEX

AO	+	!	ASTRAZENECA	<u>50MG/ML</u>	<u>N021344</u>	<u>001</u>	Apr 25, 2002
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FULVESTRANT

AO			ACCORD HLTHCARE	<u>50MG/ML</u>	<u>A211689</u>	<u>001</u>	Nov 17, 2020
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AO			AMNEAL	<u>50MG/ML</u>	<u>A210044</u>	<u>001</u>	Mar 04, 2019
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AO			APOTEX	<u>50MG/ML</u>	<u>A211730</u>	<u>001</u>	Jun 11, 2021
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AO			CHIA TAI TIANQING	<u>50MG/ML</u>	<u>A211422</u>	<u>001</u>	Feb 07, 2020
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AO			DR REDDYS	<u>50MG/ML</u>	<u>A209246</u>	<u>001</u>	Aug 07, 2020
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AO			EUGIA PHARMA	<u>50MG/ML</u>	<u>A208811</u>	<u>001</u>	Jul 23, 2019
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AO			GLENMARK PHARMS INC	<u>50MG/ML</u>	<u>A207754</u>	<u>001</u>	Aug 22, 2019
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AO			HBT LABS INC	<u>50MG/ML</u>	<u>A209714</u>	<u>001</u>	Nov 21, 2019
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AO			SAGENT PHARMS INC	<u>50MG/ML</u>	<u>A205871</u>	<u>001</u>	Aug 22, 2019
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AO			SANDOZ INC	<u>50MG/ML</u>	<u>A205935</u>	<u>001</u>	May 14, 2019
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AO			XIROMED	<u>50MG/ML</u>	<u>A213553</u>	<u>001</u>	Aug 13, 2021
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SOLUTION; INTRAMUSCULAR

FULVESTRANT

			FRESENIUS KABI USA	250MG/5ML (50MG/ML)	N210326	001	May 20, 2019
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			TEVA PHARMS USA INC	250MG/5ML (50MG/ML)	N210063	001	Aug 19, 2019
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FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

AP			ACCORD HLTHCARE	<u>10MG/ML</u>	<u>A070017</u>	<u>001</u>	Dec 15, 1986
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AP			AREVA PHARMS	<u>10MG/ML</u>	<u>A208435</u>	<u>001</u>	Dec 18, 2020
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AP			ATHENEX INC	<u>10MG/ML</u>	<u>A214766</u>	<u>001</u>	Jan 27, 2021
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AP	!		BAXTER HLTHCARE	<u>10MG/ML</u>	<u>A202747</u>	<u>001</u>	Jan 27, 2014
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CORP

AP			EMCURE PHARMS LTD	<u>10MG/ML</u>	<u>A203428</u>	<u>001</u>	Aug 26, 2014
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AP			EUGIA PHARMA	<u>10MG/ML</u>	<u>A212174</u>	<u>001</u>	May 03, 2019
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AP			FRESENIUS KABI USA	<u>10MG/ML</u>	<u>N018902</u>	<u>001</u>	May 22, 1984
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AP			GLAND PHARMA LTD	<u>10MG/ML</u>	<u>A213902</u>	<u>001</u>	Jul 01, 2020
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AP			HOSPIRA	<u>10MG/ML</u>	<u>A075241</u>	<u>001</u>	May 28, 1999
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AP				<u>10MG/ML</u>	<u>N018667</u>	<u>001</u>	May 28, 1982
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AP			WOCKHARDT	<u>10MG/ML</u>	<u>A077941</u>	<u>001</u>	Mar 22, 2007
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PRESCRIPTION DRUG PRODUCT LIST

FUROSEMIDE

SOLUTION;ORAL

FUROSEMIDE

<u>AA</u>	!	HIKMA	<u>10MG/ML</u>	<u>A070434</u>	<u>001</u>	Apr 22, 1987
<u>AA</u>		WOCKHARDT BIO AG	<u>10MG/ML</u>	<u>A070655</u>	<u>001</u>	Oct 02, 1987
		HIKMA	40MG/5ML	A070433	001	Apr 22, 1987

TABLET;ORAL

FUROSEMIDE

<u>AB</u>		EPIC PHARMA LLC	<u>20MG</u>	<u>N018569</u>	<u>002</u>	
<u>AB</u>			<u>40MG</u>	<u>N018569</u>	<u>001</u>	
<u>AB</u>			<u>80MG</u>	<u>N018569</u>	<u>005</u>	Aug 14, 1984
<u>AB</u>		HIKMA	<u>20MG</u>	<u>N018823</u>	<u>001</u>	Nov 10, 1983
<u>AB</u>			<u>40MG</u>	<u>N018823</u>	<u>002</u>	Nov 10, 1983
<u>AB</u>			<u>80MG</u>	<u>A070086</u>	<u>001</u>	Jan 24, 1986
<u>AB</u>		IPCA LABS LTD	<u>20MG</u>	<u>A078010</u>	<u>001</u>	Sep 18, 2006
<u>AB</u>			<u>40MG</u>	<u>A078010</u>	<u>002</u>	Sep 18, 2006
<u>AB</u>			<u>80MG</u>	<u>A078010</u>	<u>003</u>	Sep 18, 2006
<u>AB</u>		LEADING PHARMA LLC	<u>20MG</u>	<u>A077293</u>	<u>001</u>	Nov 09, 2005
<u>AB</u>			<u>40MG</u>	<u>A077293</u>	<u>002</u>	Nov 09, 2005
<u>AB</u>			<u>80MG</u>	<u>A077293</u>	<u>003</u>	Nov 09, 2005
<u>AB</u>		MYLAN	<u>20MG</u>	<u>N018487</u>	<u>001</u>	
<u>AB</u>			<u>40MG</u>	<u>N018487</u>	<u>002</u>	
<u>AB</u>			<u>80MG</u>	<u>A070082</u>	<u>001</u>	Oct 29, 1986
<u>AB</u>		PRINSTON INC	<u>20MG</u>	<u>A076796</u>	<u>001</u>	Mar 26, 2004
<u>AB</u>			<u>40MG</u>	<u>A076796</u>	<u>002</u>	Mar 26, 2004
<u>AB</u>			<u>80MG</u>	<u>A076796</u>	<u>003</u>	Mar 26, 2004

LASIX

<u>AB</u>	+	VALIDUS PHARMS	<u>20MG</u>	<u>N016273</u>	<u>002</u>	
<u>AB</u>	+		<u>40MG</u>	<u>N016273</u>	<u>001</u>	
<u>AB</u>	+		<u>80MG</u>	<u>N016273</u>	<u>003</u>	

GABAPENTIN

CAPSULE;ORAL

GABAPENTIN

<u>AB</u>		ACI	<u>100MG</u>	<u>A206943</u>	<u>001</u>	May 14, 2018
<u>AB</u>			<u>300MG</u>	<u>A206943</u>	<u>002</u>	May 14, 2018
<u>AB</u>			<u>400MG</u>	<u>A206943</u>	<u>003</u>	May 14, 2018
<u>AB</u>		ACTAVIS ELIZABETH	<u>100MG</u>	<u>A075350</u>	<u>001</u>	Sep 12, 2003
<u>AB</u>			<u>300MG</u>	<u>A075350</u>	<u>002</u>	Sep 12, 2003
<u>AB</u>			<u>400MG</u>	<u>A075350</u>	<u>003</u>	Sep 12, 2003
<u>AB</u>		ALKEM	<u>100MG</u>	<u>A090858</u>	<u>001</u>	Dec 17, 2010
<u>AB</u>			<u>300MG</u>	<u>A090858</u>	<u>002</u>	Dec 17, 2010
<u>AB</u>			<u>400MG</u>	<u>A090858</u>	<u>003</u>	Dec 17, 2010
<u>AB</u>		AMNEAL PHARMS NY	<u>100MG</u>	<u>A078428</u>	<u>001</u>	Jul 25, 2007
<u>AB</u>			<u>300MG</u>	<u>A078428</u>	<u>002</u>	Jul 25, 2007
<u>AB</u>			<u>400MG</u>	<u>A078428</u>	<u>003</u>	Jul 25, 2007
<u>AB</u>		ASCENT PHARMS INC	<u>100MG</u>	<u>A214956</u>	<u>001</u>	May 10, 2021
<u>AB</u>			<u>300MG</u>	<u>A214956</u>	<u>002</u>	May 10, 2021
<u>AB</u>			<u>400MG</u>	<u>A214956</u>	<u>003</u>	May 10, 2021
<u>AB</u>		AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A078787</u>	<u>001</u>	Jan 31, 2008
<u>AB</u>			<u>300MG</u>	<u>A078787</u>	<u>002</u>	Jan 31, 2008
<u>AB</u>			<u>400MG</u>	<u>A078787</u>	<u>003</u>	Jan 31, 2008
<u>AB</u>		CSPC OUYI	<u>100MG</u>	<u>A075477</u>	<u>001</u>	Mar 23, 2005
<u>AB</u>			<u>300MG</u>	<u>A075477</u>	<u>002</u>	Mar 23, 2005
<u>AB</u>			<u>400MG</u>	<u>A075477</u>	<u>003</u>	Mar 23, 2005
<u>AB</u>		GRANULES	<u>100MG</u>	<u>A075360</u>	<u>001</u>	Apr 06, 2005
<u>AB</u>			<u>300MG</u>	<u>A075360</u>	<u>002</u>	Apr 06, 2005
<u>AB</u>			<u>400MG</u>	<u>A075360</u>	<u>003</u>	Apr 06, 2005
<u>AB</u>		GRAVITI PHARMS	<u>100MG</u>	<u>A207099</u>	<u>001</u>	Mar 24, 2017
<u>AB</u>			<u>300MG</u>	<u>A207099</u>	<u>002</u>	Mar 24, 2017
<u>AB</u>			<u>400MG</u>	<u>A207099</u>	<u>003</u>	Mar 24, 2017
<u>AB</u>		INVAGEN PHARMS	<u>100MG</u>	<u>A090705</u>	<u>001</u>	Dec 30, 2009
<u>AB</u>			<u>300MG</u>	<u>A090705</u>	<u>002</u>	Dec 30, 2009
<u>AB</u>			<u>400MG</u>	<u>A090705</u>	<u>003</u>	Dec 30, 2009
<u>AB</u>		MARKSANS PHARMA	<u>100MG</u>	<u>A090007</u>	<u>001</u>	Jul 21, 2011
<u>AB</u>			<u>300MG</u>	<u>A090007</u>	<u>002</u>	Jul 21, 2011
<u>AB</u>			<u>400MG</u>	<u>A090007</u>	<u>003</u>	Jul 21, 2011
<u>AB</u>		SCIEGEN PHARMS INC	<u>100MG</u>	<u>A204989</u>	<u>001</u>	Feb 18, 2016
<u>AB</u>			<u>300MG</u>	<u>A204989</u>	<u>002</u>	Feb 18, 2016
<u>AB</u>			<u>400MG</u>	<u>A204989</u>	<u>003</u>	Feb 18, 2016
<u>AB</u>		STRIDES PHARMA	<u>100MG</u>	<u>A211314</u>	<u>001</u>	Oct 16, 2018
<u>AB</u>			<u>300MG</u>	<u>A211314</u>	<u>002</u>	Oct 16, 2018

PRESCRIPTION DRUG PRODUCT LIST

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

<u>AB</u>		<u>400MG</u>	<u>A211314</u>	<u>003</u>	Oct 16, 2018
<u>AB</u>	SUN PHARM INDS LTD	<u>100MG</u>	<u>A077242</u>	<u>001</u>	Aug 24, 2006
<u>AB</u>		<u>300MG</u>	<u>A077242</u>	<u>002</u>	Aug 24, 2006
<u>AB</u>		<u>400MG</u>	<u>A077242</u>	<u>003</u>	Aug 24, 2006
<u>AB</u>	TARO	<u>100MG</u>	<u>A077261</u>	<u>001</u>	Aug 02, 2013
<u>AB</u>		<u>300MG</u>	<u>A077261</u>	<u>002</u>	Aug 02, 2013
<u>AB</u>		<u>400MG</u>	<u>A077261</u>	<u>003</u>	Aug 02, 2013
<u>AB</u>	ZHEJIANG YONGTAI	<u>100MG</u>	<u>A213603</u>	<u>001</u>	Aug 17, 2020
<u>AB</u>		<u>300MG</u>	<u>A213603</u>	<u>002</u>	Aug 17, 2020
<u>AB</u>		<u>400MG</u>	<u>A213603</u>	<u>003</u>	Aug 17, 2020

NEURONTIN

<u>AB</u>	+ UPJOHN	<u>100MG</u>	<u>N020235</u>	<u>001</u>	Dec 30, 1993
<u>AB</u>	+ UPJOHN	<u>300MG</u>	<u>N020235</u>	<u>002</u>	Dec 30, 1993
<u>AB</u>	+!	<u>400MG</u>	<u>N020235</u>	<u>003</u>	Dec 30, 1993

SOLUTION; ORAL

GABAPENTIN

<u>AA</u>	ACELLA PHARMS LLC	<u>250MG/5ML</u>	<u>A076403</u>	<u>001</u>	May 01, 2012
<u>AA</u>	AKORN	<u>250MG/5ML</u>	<u>A078974</u>	<u>001</u>	Feb 18, 2011
<u>AA</u>	AMNEAL PHARMS	<u>250MG/5ML</u>	<u>A202024</u>	<u>001</u>	Mar 23, 2012
<u>AA</u>	TARO	<u>250MG/5ML</u>	<u>A076672</u>	<u>001</u>	Jul 03, 2013
<u>AA</u>	TRIS PHARMA INC	<u>250MG/5ML</u>	<u>A091286</u>	<u>001</u>	Mar 14, 2016

NEURONTIN

<u>AA</u>	+! UPJOHN	<u>250MG/5ML</u>	<u>N021129</u>	<u>001</u>	Mar 02, 2000
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TABLET; ORAL

GABAPENTIN

<u>AB</u>	ACI	<u>600MG</u>	<u>A203244</u>	<u>002</u>	Jul 12, 2013
<u>AB</u>		<u>800MG</u>	<u>A203244</u>	<u>001</u>	Jul 12, 2013
<u>AB</u>	ACTAVIS ELIZABETH	<u>600MG</u>	<u>A075694</u>	<u>001</u>	Oct 21, 2004
<u>AB</u>		<u>800MG</u>	<u>A075694</u>	<u>002</u>	Oct 21, 2004
<u>AB</u>	ALKEM LABS LTD	<u>600MG</u>	<u>A206402</u>	<u>001</u>	Dec 23, 2015
<u>AB</u>		<u>800MG</u>	<u>A206402</u>	<u>002</u>	Dec 23, 2015
<u>AB</u>	APOTEX INC	<u>100MG</u>	<u>A077894</u>	<u>001</u>	Oct 10, 2006
<u>AB</u>		<u>300MG</u>	<u>A077894</u>	<u>002</u>	Oct 10, 2006
<u>AB</u>		<u>400MG</u>	<u>A077894</u>	<u>003</u>	Oct 10, 2006
<u>AB</u>	ASCENT PHARMS INC	<u>600MG</u>	<u>A214957</u>	<u>001</u>	Oct 01, 2021
<u>AB</u>		<u>800MG</u>	<u>A214957</u>	<u>002</u>	Oct 01, 2021
<u>AB</u>	AUROBINDO PHARMA LTD	<u>600MG</u>	<u>A200651</u>	<u>001</u>	Oct 06, 2011
<u>AB</u>		<u>800MG</u>	<u>A200651</u>	<u>002</u>	Oct 06, 2011
<u>AB</u>	CSPC OUYI	<u>600MG</u>	<u>A207057</u>	<u>001</u>	Oct 26, 2017
<u>AB</u>		<u>800MG</u>	<u>A207057</u>	<u>002</u>	Oct 26, 2017
<u>AB</u>	GLENMARK PHARMS LTD	<u>600MG</u>	<u>A077662</u>	<u>001</u>	Aug 18, 2006
<u>AB</u>		<u>800MG</u>	<u>A077662</u>	<u>002</u>	Aug 18, 2006
<u>AB</u>	INVAGEN PHARMS	<u>600MG</u>	<u>A202764</u>	<u>001</u>	Oct 16, 2012
<u>AB</u>		<u>800MG</u>	<u>A202764</u>	<u>002</u>	Oct 16, 2012
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>100MG</u>	<u>A076017</u>	<u>001</u>	Apr 28, 2004
<u>AB</u>		<u>300MG</u>	<u>A076017</u>	<u>002</u>	Apr 28, 2004
<u>AB</u>		<u>400MG</u>	<u>A076017</u>	<u>003</u>	Apr 28, 2004
<u>AB</u>	RUBICON	<u>600MG</u>	<u>A077661</u>	<u>004</u>	Sep 13, 2006
<u>AB</u>		<u>800MG</u>	<u>A077661</u>	<u>005</u>	Sep 13, 2006
<u>AB</u>	SCIEGEN PHARMS INC	<u>600MG</u>	<u>A205101</u>	<u>001</u>	Feb 04, 2016
<u>AB</u>		<u>800MG</u>	<u>A205101</u>	<u>002</u>	Feb 04, 2016
<u>AB</u>	SUN PHARM INDS LTD	<u>600MG</u>	<u>A077525</u>	<u>001</u>	Aug 24, 2006
<u>AB</u>		<u>800MG</u>	<u>A077525</u>	<u>002</u>	Aug 24, 2006
<u>AB</u>	ZYDUS PHARMS USA INC	<u>600MG</u>	<u>A078926</u>	<u>001</u>	Feb 11, 2011
<u>AB</u>		<u>800MG</u>	<u>A078926</u>	<u>002</u>	Feb 11, 2011

NEURONTIN

<u>AB</u>	+ UPJOHN	<u>600MG</u>	<u>N020882</u>	<u>001</u>	Oct 09, 1998
<u>AB</u>	+!	<u>800MG</u>	<u>N020882</u>	<u>002</u>	Oct 09, 1998

GRALISE

BX	+! ALMATICA	300MG	N022544	001	Jan 28, 2011
BX	+!	600MG	N022544	002	Jan 28, 2011

PRESCRIPTION DRUG PRODUCT LIST

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

GADOBUTROL

SOLUTION; INTRAVENOUS

GDAVIST

+	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

GADOTERATE MEGLUMINE

SOLUTION; INTRAVENOUS

CLARISCAN

<u>AP</u>	GE HEALTHCARE	<u>7.538GM/20ML (376.9MG/ML)</u>	<u>A210016</u>	<u>003</u>	Nov 01, 2019
<u>AP</u>		<u>1.8845GM/5ML (376.9MG/ML)</u>	<u>A210016</u>	<u>005</u>	Nov 24, 2020
<u>AP</u>		<u>3.769GM/10ML (376.9MG/ML)</u>	<u>A210016</u>	<u>001</u>	Nov 01, 2019
<u>AP</u>		<u>5.6535GM/15ML (376.9MG/ML)</u>	<u>A210016</u>	<u>002</u>	Nov 01, 2019
<u>AP</u>		<u>37.69GM/100ML (376.9MG/ML)</u>	<u>A210016</u>	<u>004</u>	Aug 04, 2020

DOTAREM

<u>AP</u>	+	GUERBET	<u>37.69GM/100ML (376.9MG/ML)</u>	<u>N204781</u>	<u>001</u>	Mar 20, 2013
<u>AP</u>	+	!	<u>1.8845GM/5ML (376.9MG/ML)</u>	<u>N204781</u>	<u>005</u>	Mar 31, 2017
<u>AP</u>	+	!	<u>3.769GM/10ML (376.9MG/ML)</u>	<u>N204781</u>	<u>002</u>	Mar 20, 2013
<u>AP</u>	+	!	<u>5.6535GM/15ML (376.9MG/ML)</u>	<u>N204781</u>	<u>003</u>	Mar 20, 2013
<u>AP</u>	+	!	<u>7.538GM/20ML (376.9MG/ML)</u>	<u>N204781</u>	<u>004</u>	Mar 20, 2013

GADOTERIDOL

INJECTABLE; INJECTION

PROHANCE

+	BRACCO	279.3MG/ML	N020131	001	Nov 16, 1992
+	!		N021489	001	Oct 09, 2003

PROHANCE MULTIPACK

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

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 8MG BASE</u>	<u>A204895</u>	<u>001</u>	Aug 05, 2016
<u>AB</u>		<u>EQ 16MG BASE</u>	<u>A204895</u>	<u>002</u>	Aug 05, 2016
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A204895</u>	<u>003</u>	Aug 05, 2016
<u>AB</u>	BARR	<u>EQ 8MG BASE</u>	<u>A078189</u>	<u>001</u>	Sep 15, 2008
<u>AB</u>		<u>EQ 16MG BASE</u>	<u>A078189</u>	<u>002</u>	Sep 15, 2008
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A078189</u>	<u>003</u>	Sep 15, 2008
<u>AB</u>	SUN PHARM	<u>EQ 8MG BASE</u>	<u>A090178</u>	<u>001</u>	Feb 02, 2011
<u>AB</u>		<u>EQ 16MG BASE</u>	<u>A090178</u>	<u>002</u>	Feb 02, 2011
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A090178</u>	<u>003</u>	Feb 02, 2011
<u>AB</u>	WATSON LABS	<u>EQ 8MG BASE</u>	<u>A079028</u>	<u>001</u>	Dec 15, 2008
<u>AB</u>		<u>EQ 16MG BASE</u>	<u>A079028</u>	<u>002</u>	Dec 15, 2008

PRESCRIPTION DRUG PRODUCT LIST

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE;ORAL

GALANTAMINE HYDROBROMIDE

AB		<u>EQ 24MG BASE</u>	<u>A079028 003</u>	Dec 15, 2008
	<u>RAZADYNE ER</u>			
AB	+ ! JANSSEN PHARMS	<u>EQ 8MG BASE</u>	<u>N021615 001</u>	Apr 01, 2005
AB	+	<u>EQ 16MG BASE</u>	<u>N021615 002</u>	Apr 01, 2005
AB	+	<u>EQ 24MG BASE</u>	<u>N021615 003</u>	Apr 01, 2005

SOLUTION;ORAL

GALANTAMINE HYDROBROMIDE

! HIKMA

4MG/ML

A078185 001 Jan 30, 2009

TABLET;ORAL

GALANTAMINE HYDROBROMIDE

AB	AUROBINDO PHARMA LTD	<u>EQ 4MG BASE</u>	<u>A090957 001</u>	Mar 29, 2011
AB		<u>EQ 8MG BASE</u>	<u>A090957 002</u>	Mar 29, 2011
AB		<u>EQ 12MG BASE</u>	<u>A090957 003</u>	Mar 29, 2011
AB	BARR	<u>EQ 4MG BASE</u>	<u>A077605 001</u>	Aug 28, 2008
AB		<u>EQ 8MG BASE</u>	<u>A077605 002</u>	Aug 28, 2008
AB		<u>EQ 12MG BASE</u>	<u>A077605 003</u>	Aug 28, 2008
AB	DR REDDYS LABS LTD	<u>EQ 4MG BASE</u>	<u>A077593 001</u>	Sep 11, 2008
AB		<u>EQ 8MG BASE</u>	<u>A077593 002</u>	Sep 11, 2008
AB		<u>EQ 12MG BASE</u>	<u>A077593 003</u>	Sep 11, 2008
AB	SANDOZ	<u>EQ 4MG BASE</u>	<u>A077589 001</u>	Jun 22, 2009
AB		<u>EQ 8MG BASE</u>	<u>A077589 002</u>	Jun 22, 2009
AB		<u>EQ 12MG BASE</u>	<u>A077589 003</u>	Jun 22, 2009
AB	! YABAO PHARM	<u>EQ 4MG BASE</u>	<u>A077604 001</u>	Feb 06, 2009
AB		<u>EQ 8MG BASE</u>	<u>A077604 002</u>	Feb 06, 2009
AB		<u>EQ 12MG BASE</u>	<u>A077604 003</u>	Feb 06, 2009
AB	ZYDUS PHARMS USA INC	<u>EQ 4MG BASE</u>	<u>A078898 001</u>	Feb 17, 2011
AB		<u>EQ 8MG BASE</u>	<u>A078898 002</u>	Feb 17, 2011
AB		<u>EQ 12MG BASE</u>	<u>A078898 003</u>	Feb 17, 2011

GALLIUM CITRATE GA-67

INJECTABLE;INJECTION

GALLIUM CITRATE GA 67

BS	CURIUM	2mCi/ML	N018058 001	
BS	LANTHEUS MEDCL	2mCi/ML	N017478 001	

GALLIUM DOTATATE GA-68

POWDER;INTRAVENOUS

NETSPOT

+! AAA USA INC

2.1-5.5mCi/ML

N208547 001 Jun 01, 2016

GALLIUM DOTATOC GA-68

SOLUTION;INTRAVENOUS

GALLIUM DOTATOC GA 68

+! UIHC PET IMAGING

0.5-4mCi/ML

N210828 001 Aug 21, 2019

GALLIUM GA-68 GOZETOTIDE

POWDER;INTRAVENOUS

ILLUCCIX

+! TELIX

6.7mCi/ML

N214032 001 Dec 17, 2021

GALLIUM GA-68 PSMA-11

SOLUTION;INTRAVENOUS

GALLIUM GA 68 PSMA-11

+! UNIV CA LOS ANGELES 30ML (0.5-5mCi/mL)

N212642 001 Dec 01, 2020

+! UNIV OF CA SAN FRAN 20ML (0.5-5mCi/mL)

N212643 001 Dec 01, 2020

GANCICLOVIR

GEL;OPHTHALMIC

ZIRGAN

+! BAUSCH AND LOMB

0.15%

N022211 001 Sep 15, 2009

SOLUTION;INTRAVENOUS

GANZYK-RTU

+! EXELA PHARMA

500MG/250ML (2MG/ML)

N209347 001 Feb 17, 2017

GANCICLOVIR SODIUM

INJECTABLE;INJECTION

GANCICLOVIR SODIUM

AP	FRESENIUS KABI USA	<u>EQ 500MG BASE/VIAL</u>	<u>A090658 001</u>	Jun 21, 2010
AP	HAINAN POLY PHARM	<u>EQ 500MG BASE/VIAL</u>	<u>A204204 001</u>	Nov 08, 2018
AP	HIKMA	<u>EQ 500MG BASE/VIAL</u>	<u>A076222 001</u>	Jul 16, 2003
AP	PAR STERILE	<u>EQ 500MG BASE/VIAL</u>	<u>A204950 001</u>	Dec 06, 2016

PRESCRIPTION DRUG PRODUCT LIST

GANCICLOVIR SODIUM

INJECTABLE; INJECTION

GANCICLOVIR SODIUM

PRODUCTS

<u>AP</u>	!	PHARMASCIENCE INC	<u>EQ 500MG BASE/VIAL</u>	<u>A207645 001</u>	Dec 08, 2017
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GANIRELIX ACETATE

INJECTABLE; INJECTION

GANIRELIX ACETATE

<u>AP</u>	+	!	ORGANON USA INC	<u>250MCG/0.5ML</u>	<u>N021057 001</u>	Jul 29, 1999
<u>AP</u>			SUN PHARM	<u>250MCG/0.5ML</u>	<u>A204246 001</u>	Nov 30, 2018

GATIFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

GATIFLOXACIN

<u>AT</u>			AKORN	<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>			SANDOZ INC	<u>0.5%</u>	<u>A204227 001</u>	Jul 11, 2016
<u>AT</u>			TORRENT	<u>0.5%</u>	<u>A213542 001</u>	Nov 03, 2021

ZYMAXID

<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	250MG	N206995 001	Jul 13, 2015
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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>			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>			FRESENIUS KABI USA	<u>EQ 200MG BASE/VIAL</u>	<u>A090799 001</u>	Jul 25, 2011
<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>A090242 003</u>	May 16, 2011
<u>AP</u>				<u>EQ 2GM BASE/VIAL</u>	<u>A090799 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>			HIKMA INTL PHARMS	<u>EQ 200MG BASE/VIAL</u>	<u>A206617 001</u>	Jun 25, 2021
<u>AP</u>				<u>EQ 1GM BASE/VIAL</u>	<u>A206617 002</u>	Jun 25, 2021
<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>			MEITHEAL	<u>200MG/5.26ML (38MG/ML)</u>	<u>A212129 001</u>	Dec 11, 2020
<u>AP</u>				<u>1GM/26.3ML (38MG/ML)</u>	<u>A212129 002</u>	Dec 11, 2020
<u>AP</u>				<u>2GM/52.6ML (38MG/ML)</u>	<u>A212129 003</u>	Dec 11, 2020
<u>AP</u>			MYLAN LABS LTD	<u>200MG/5.26ML (38MG/ML)</u>	<u>A205242 001</u>	Dec 06, 2017
<u>AP</u>				<u>1GM/26.3ML (38MG/ML)</u>	<u>A205242 002</u>	Dec 06, 2017
<u>AP</u>				<u>2GM/52.6ML (38MG/ML)</u>	<u>A205242 003</u>	Dec 06, 2017
<u>AP</u>			SAGENT PHARMS INC	<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>			SHILPA	<u>EQ 200MG BASE/VIAL</u>	<u>A207575 001</u>	Feb 22, 2019
<u>AP</u>				<u>200MG/5.26ML (38MG/ML)</u>	<u>A210991 001</u>	Oct 04, 2019
<u>AP</u>				<u>EQ 1GM BASE/VIAL</u>	<u>A207575 002</u>	Feb 22, 2019
<u>AP</u>				<u>1GM/26.3ML (38MG/ML)</u>	<u>A210991 002</u>	Oct 04, 2019
<u>AP</u>				<u>2GM/52.6ML (38MG/ML)</u>	<u>A210991 003</u>	Oct 04, 2019
<u>AP</u>			SUN PHARM	<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

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	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

AB		APOTEX	600MG	A075034	001	Jul 20, 1998
AB		ASCENT PHARMS INC	600MG	A214603	001	Jan 13, 2021
AB		AUROBINDO PHARMA LTD	600MG	A202726	001	Sep 16, 2015
AB		CADILA	600MG	A204189	001	Aug 28, 2018
AB		CADILA PHARMS LTD	600MG	A203266	001	Jun 17, 2016
AB		CARIBE HOLDINGS	600MG	A078012	001	Mar 26, 2007
AB		CHARTWELL MOLECULES	600MG	A074270	001	Sep 27, 1993
AB		IMPAX PHARMS	600MG	A078207	001	Jun 01, 2007
AB		INVAGEN PHARMS	600MG	A077836	001	Jul 27, 2006
AB		NORTHSTAR HLTHCARE	600MG	A079072	001	Sep 13, 2010

LOPID

AB	+	!	PFIZER PHARMS	600MG	N018422	003	Nov 20, 1986
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GEMIFLOXACIN MESYLATE

TABLET; ORAL

FACTIVE

AB	+	!	LG CHEM LTD	EQ 320MG BASE	N021158	001	Apr 04, 2003
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GEMIFLOXACIN MESYLATE

AB		ORBION PHARMS	EQ 320MG BASE	A090466	001	Jun 15, 2015
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GENTAMICIN SULFATE

CREAM; TOPICAL

GENTAMICIN SULFATE

AT		COSETTE	EQ 0.1% BASE	A064056	001	Apr 29, 1994
AT	!	PADAGIS US	EQ 0.1% BASE	A062307	001	

INJECTABLE; INJECTION

GENTAMICIN SULFATE

AP	!	FRESENIUS KABI USA	EQ 10MG BASE/ML	A062366	002	Feb 06, 1986
AP	!		EQ 40MG BASE/ML	A062366	001	Aug 04, 1983
AP		HOSPIRA	EQ 10MG BASE/ML	A062420	001	Aug 15, 1983
AP			EQ 40MG BASE/ML	A062420	002	Aug 15, 1983

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP		BAXTER HLTHCARE	EQ 1.2MG BASE/ML	A062373	007	Sep 07, 1982
AP			EQ 1.6MG BASE/ML	A062373	008	Sep 07, 1982
AP			EQ 80MG BASE/100ML	A062373	002	Sep 07, 1982
AP			EQ 100MG BASE/100ML	A062373	005	Sep 07, 1982
AP		HOSPIRA	EQ 1.2MG BASE/ML	A062414	001	Aug 15, 1983
AP			EQ 1.6MG BASE/ML	A062414	003	Aug 15, 1983
AP			EQ 80MG BASE/100ML	A062414	008	Aug 15, 1983
AP			EQ 100MG BASE/100ML	A062414	010	Aug 15, 1983
	!	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	EQ 0.3% BASE	A064093	001	Aug 31, 1995
AT		FERA PHARMS LLC	EQ 0.3% BASE	A065024	001	Jul 30, 2004

OINTMENT; TOPICAL

GENTAMICIN SULFATE

AT		COSETTE	EQ 0.1% BASE	A064054	001	Apr 29, 1994
AT		FOUGERA PHARMS INC	EQ 0.1% BASE	A062533	001	Oct 05, 1984
AT	!	PADAGIS US	EQ 0.1% BASE	A062351	001	Feb 18, 1982

PRESCRIPTION DRUG PRODUCT LIST

GENTAMICIN SULFATE

OINTMENT; TOPICAL

GENTAMICIN SULFATE

AT	TARO	EQ 0.1% BASE	A062477 001	Dec 23, 1983
AT	TELLIGENT	EQ 0.1% BASE	A209233 001	Dec 31, 2018

SOLUTION/DROPS; OPHTHALMIC

GENOPTIC

AT	! ALLERGAN	EQ 0.3% BASE	A062452 001	Oct 10, 1984
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GENTAK

AT	AKORN	EQ 0.3% BASE	A064163 001	Oct 12, 2001
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GENTAMICIN SULFATE

AT	AKORN	EQ 0.3% BASE	A062635 001	Jan 08, 1987
AT	BAUSCH AND LOMB	EQ 0.3% BASE	A064048 001	May 11, 1994
AT	PADAGIS US	EQ 0.3% BASE	A065121 001	Jan 30, 2004
AT	SANDOZ INC	EQ 0.3% BASE	A062196 001	

GENTAMICIN SULFATE; PREDNISOLONE ACETATE

OINTMENT; OPHTHALMIC

PRED-G

+	! ALLERGAN	EQ 0.3% BASE; 0.6%	N050612 001	Dec 01, 1989
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SUSPENSION/DROPS; OPHTHALMIC

PRED-G

+	! ALLERGAN	EQ 0.3% BASE; 1%	N050586 001	Jun 10, 1988
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GILTERITINIB FUMARATE

TABLET; ORAL

XOSPATA

+	! ASTELLAS	EQ 40MG BASE	N211349 001	Nov 28, 2018
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GIVOSIRAN SODIUM

SOLUTION; SUBCUTANEOUS

GIVLAARI

+	! ALNYLAM PHARMS INC	EQ 189MG BASE/ML (EQ 189MG BASE/ML)	N212194 001	Nov 20, 2019
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GLASDEGIB MALEATE

TABLET; ORAL

DAURISMO

+	PFIZER	EQ 25MG BASE	N210656 001	Nov 21, 2018
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+	!	EQ 100MG BASE	N210656 002	Nov 21, 2018
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GLATIRAMER ACETATE

INJECTABLE; SUBCUTANEOUS

COPAXONE

AP	+	! TEVA PHARMS USA	20MG/ML	N020622 002	Feb 12, 2002
AP	+	!	40MG/ML	N020622 003	Jan 28, 2014

GLATIRAMER ACETATE

AP	MYLAN	20MG/ML	A091646 001	Oct 03, 2017
AP		40MG/ML	A206936 001	Oct 03, 2017

GLATOPIA

AP	SANDOZ INC	20MG/ML	A090218 001	Apr 16, 2015
AP		40MG/ML	A206921 001	Feb 12, 2018

GLECAPREVIR; PIBRENTASVIR

PELLETS; ORAL

MAVYRET

+	! ABBVIE INC	50MG; 20MG/PACKET	N215110 001	Jun 10, 2021
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TABLET; ORAL

MAVYRET

+	! ABBVIE INC	100MG; 40MG	N209394 001	Aug 03, 2017
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GLIMEPIRIDE

TABLET; ORAL

AMARYL

AB	+	! SANOFI AVENTIS US	1MG	N020496 001	Nov 30, 1995
AB	+		2MG	N020496 002	Nov 30, 1995
AB	+		4MG	N020496 003	Nov 30, 1995

GLIMEPIRIDE

AB	ACCORD HLTHCARE	1MG	A078181 001	Aug 23, 2007
AB		2MG	A078181 002	Aug 23, 2007
AB		4MG	A078181 003	Aug 23, 2007
AB	AUROBINDO PHARMA LTD	1MG	A202759 001	Jun 29, 2012
AB		2MG	A202759 002	Jun 29, 2012
AB		4MG	A202759 003	Jun 29, 2012
AB	CARLSBAD	1MG	A077911 001	Sep 22, 2009
AB		2MG	A077911 002	Sep 22, 2009

PRESCRIPTION DRUG PRODUCT LIST

GLIMEPIRIDE

TABLET; ORAL

GLIMEPIRIDE

<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>	MICRO LABS	<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
<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
	MICRO LABS	3MG	A091220	003	Jun 29, 2012
		6MG	A091220	005	Jun 29, 2012

GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

DUETACT

<u>AB</u>	<u>+</u> !	TAKEDA PHARMS USA	<u>2MG;30MG</u>	<u>N021925</u>	<u>001</u>	Jul 28, 2006
<u>AB</u>	<u>+</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	<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>	<u>+</u>	PFIZER	<u>5MG</u>	<u>N017783</u>	<u>001</u>	May 08, 1984
<u>AB</u>	<u>+</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>		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	<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>	<u>+</u>	PFIZER	<u>2.5MG</u>	<u>N020329</u>	<u>003</u>	Aug 10, 1999
<u>AB</u>	<u>+</u>		<u>5MG</u>	<u>N020329</u>	<u>001</u>	Apr 26, 1994
<u>AB</u>	<u>+</u> !		<u>10MG</u>	<u>N020329</u>	<u>002</u>	Apr 26, 1994

PRESCRIPTION DRUG PRODUCT LIST

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

AB	EPIC PHARMA LLC	2.5MG;250MG	A077507 001	Oct 27, 2005
AB		2.5MG;500MG	A077507 002	Oct 27, 2005
AB		5MG;500MG	A077507 003	Oct 27, 2005
AB	HERITAGE PHARMS INC	2.5MG;250MG	A078728 001	Jun 23, 2010
AB		2.5MG;500MG	A078728 002	Jun 23, 2010
AB		5MG;500MG	A078728 003	Jun 23, 2010
AB	TEVA PHARMS	2.5MG;250MG	A077270 001	Oct 28, 2005
AB		2.5MG;500MG	A077270 002	Oct 28, 2005
AB	!	5MG;500MG	A077270 003	Oct 28, 2005
AB	ZYDUS PHARMS USA INC	2.5MG;250MG	A078905 001	Jan 31, 2011
AB		2.5MG;500MG	A078905 002	Jan 31, 2011
AB		5MG;500MG	A078905 003	Jan 31, 2011

GLUCAGON

INJECTABLE; INJECTION

GLUCAGON

AP	AMPHASTAR PHARMS INC	1MG/VIAL	A208086 001	Dec 28, 2020
AP	+! LILLY	1MG/VIAL	N020928 001	Sep 11, 1998
	POWDER; NASAL BAQSIMI			
	+! ELI LILLY AND CO	3MG	N210134 001	Jul 24, 2019
	SOLUTION; SUBCUTANEOUS GVOKE HYOPEN			
	+! XERIS	0.5MG/0.1ML (0.5MG/0.1ML)	N212097 003	Sep 10, 2019
	+!	1MG/0.2ML (1MG/0.2ML)	N212097 004	Sep 10, 2019
	GVOKE PFS			
	+! XERIS	0.5MG/0.1ML (0.5MG/0.1ML)	N212097 001	Sep 10, 2019
	+!	1MG/0.2ML (1MG/0.2ML)	N212097 002	Sep 10, 2019

GLUCAGON HYDROCHLORIDE

INJECTABLE; INJECTION

GLUCAGON

	+! 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)

AB	HIKMA	1.5MG	A075890 001	Jul 31, 2003
AB		3MG	A075890 002	Jul 31, 2003
AB		6MG	A075890 003	Jul 31, 2003
AB	TEVA	1.5MG	A074686 001	Apr 20, 1999
AB		3MG	A074686 002	Apr 20, 1999
AB		6MG	A074686 004	Apr 20, 1999

GLYNASE

AB	+ PFIZER	1.5MG	N020051 001	Mar 04, 1992
AB	+	3MG	N020051 002	Mar 04, 1992
AB	+!	6MG	N020051 004	Sep 24, 1993

GLYBURIDE

AB1	CADILA PHARMS LTD	1.25MG	A203379 001	Jan 04, 2019
AB1		2.5MG	A203379 002	Jan 04, 2019
AB1		5MG	A203379 003	Jan 04, 2019
AB1	EPIC PHARMA LLC	1.25MG	A076257 001	Jun 27, 2002
AB1		2.5MG	A076257 002	Jun 27, 2002
AB1		5MG	A076257 003	Jun 27, 2002
AB1	HERITAGE PHARMS INC	1.25MG	A090937 001	Feb 28, 2011
AB1		2.5MG	A090937 002	Feb 28, 2011
AB1		5MG	A090937 003	Feb 28, 2011
AB1	ORIENT PHARMA CO LTD	1.25MG	A206483 001	Feb 22, 2019
AB1		2.5MG	A206483 002	Feb 22, 2019
AB1		5MG	A206483 003	Feb 22, 2019
AB1	SUNNY	1.25MG	A203581 001	Apr 14, 2016
AB1		2.5MG	A203581 002	Apr 14, 2016
AB1		5MG	A203581 003	Apr 14, 2016
AB1	TEVA	1.25MG	A074388 001	Aug 29, 1995
AB1		2.5MG	A074388 002	Aug 29, 1995

PRESCRIPTION DRUG PRODUCT LIST

GLYBURIDE

TABLET; ORAL

GLYBURIDE

<u>AB1</u>	!		<u>5MG</u>	<u>A074388</u>	<u>003</u>	Aug 29, 1995
<u>AB1</u>		ZYDUS PHARMS	<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 (MICRONIZED)				
		TEVA	4.5MG	A074686	003	Apr 20, 1999

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>		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	<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 THERAP	1.1GM/ML	N203284	001	Feb 01, 2013
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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>	
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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>	

GLYCOPYRROLATE

INJECTABLE; INJECTION

GLYCOPYRROLATE

<u>AP</u>		ACCORD HLTHCARE	<u>0.2MG/ML</u>	<u>A213238</u>	<u>001</u>	Jul 08, 2020
<u>AP</u>		AM REGENT	<u>0.2MG/ML</u>	<u>A089335</u>	<u>001</u>	Jul 23, 1986
<u>AP</u>		AMNEAL	<u>0.2MG/ML</u>	<u>A208973</u>	<u>001</u>	Jun 15, 2017
<u>AP</u>		APOTEX	<u>0.2MG/ML</u>	<u>A210246</u>	<u>001</u>	Oct 29, 2019
<u>AP</u>		ATHENEX INC	<u>0.2MG/ML</u>	<u>A210083</u>	<u>001</u>	Feb 21, 2020
<u>AP</u>		CAPLIN	<u>0.2MG/ML</u>	<u>A211705</u>	<u>001</u>	Mar 20, 2019
<u>AP</u>		EUGIA PHARMA	<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>		GLAND PHARMA LTD	<u>0.2MG/ML</u>	<u>A212612</u>	<u>001</u>	Sep 30, 2019
<u>AP</u>	!	HIKMA FARMACEUTICA	<u>0.2MG/ML</u>	<u>A090963</u>	<u>001</u>	Sep 21, 2011
<u>AP</u>		MEITHEAL	<u>0.2MG/ML</u>	<u>A212802</u>	<u>001</u>	Jul 06, 2021
<u>AP</u>		NIVAGEN PHARMS INC	<u>0.2MG/ML</u>	<u>A212591</u>	<u>001</u>	Oct 13, 2021
<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>		SANDOZ INC	<u>0.2MG/ML</u>	<u>A211334</u>	<u>001</u>	May 14, 2019
<u>AP</u>		SOMERSET THERAPS	<u>0.2MG/ML</u>	<u>A207639</u>	<u>001</u>	Jun 23, 2017
		LLC				
<u>AP</u>		XIROMED	<u>0.2MG/ML</u>	<u>A212227</u>	<u>001</u>	Mar 04, 2021
<u>AP</u>		ZYDUS PHARMS	<u>0.2MG/ML</u>	<u>A214213</u>	<u>001</u>	Nov 09, 2021

SOLUTION; INHALATION

LONHALA MAGNAIR KIT

	+	SUNOVION RESP	25MCG/ML	N208437	001	Dec 05, 2017
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PRESCRIPTION DRUG PRODUCT LIST

GLYCOPYRROLATE

SOLUTION;INTRAMUSCULAR, INTRAVENOUS

GLYRX-PF

+	!	EXELA PHARMA	0.2MG/ML (0.2MG/ML)	N210997	001	Jul 11, 2018
+	!		0.4MG/2ML (0.2MG/ML)	N210997	002	Jul 11, 2018
+	!		0.6MG/3ML (0.2MG/ML)	N210997	004	Dec 14, 2020
+	!		1MG/5ML (0.2MG/ML)	N210997	003	Apr 09, 2020

SOLUTION;ORAL

CUVPOSA

<u>AA</u>	+	!	MERZ PHARMS	<u>1MG/5ML</u>	<u>N022571</u>	<u>001</u>	Jul 28, 2010
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GLYCOPYRROLATE

<u>AA</u>			PAR PHARM INC	<u>1MG/5ML</u>	<u>A204438</u>	<u>001</u>	Aug 09, 2021
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TABLET;ORAL

GLYCOPYRROLATE

<u>AA</u>			ALEMBIC LABS	<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>			AUROLIFE PHARMA LLC	<u>1MG</u>	<u>A202675</u>	<u>001</u>	Apr 15, 2013
<u>AA</u>				<u>2MG</u>	<u>A202675</u>	<u>002</u>	Oct 30, 2018
<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>			KENTON	<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>			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	<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>			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	<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

ROBINUL

<u>AA</u>	+		CASPER PHARMA LLC	<u>1MG</u>	<u>N012827</u>	<u>001</u>	
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ROBINUL FORTE

<u>AA</u>	+		CASPER PHARMA LLC	<u>2MG</u>	<u>N012827</u>	<u>002</u>	
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GLYCOPYRROLATE

			LGM PHARMA	1.5MG	A091522	001	Mar 12, 2012
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TABLET, ORALLY DISINTEGRATING;ORAL

DARTISLA ODT

+	!	EDENBRIDGE PHARMS	1.7MG	N215019	001	Dec 16, 2021
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GLYCOPYRROLATE TOSYLATE

CLOTH;TOPICAL

QBREXZA

+	!	JOURNEY	EQ 2.4% BASE	N210361	001	Jun 28, 2018
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GOLODIRSEN

SOLUTION;INTRAVENOUS

VYONDYS 53

+	!	SAREPTA THERAPS INC	100MG/2ML (50MG/ML)	N211970	001	Dec 12, 2019
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GOSERELIN ACETATE

IMPLANT;IMPLANTATION

ZOLADEX

+	!	TERSERA	EQ 3.6MG BASE	N019726	001	Dec 29, 1989
+	!		EQ 10.8MG BASE	N020578	001	Jan 11, 1996

GRAMICIDIN; 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>	
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PRESCRIPTION DRUG PRODUCT LIST

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	<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>	BIONPHARMA INC	<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>	! DR REDDYS	<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>	EUGIA PHARMA	<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>	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	<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>	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>	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>	PUNISKA	<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>	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>	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>	BIONPHARMA INC	<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>	! HIKMA	<u>EQ 1MG BASE</u>	<u>A077842</u>	<u>001</u>	Dec 31, 2007
<u>AB</u>	NATCO PHARMA	<u>EQ 1MG BASE</u>	<u>A078969</u>	<u>001</u>	Jun 22, 2009
<u>AB</u>	ORBION PHARMS	<u>EQ 1MG BASE</u>	<u>A078678</u>	<u>001</u>	Feb 13, 2008
<u>AB</u>	TARO	<u>EQ 1MG BASE</u>	<u>A090817</u>	<u>001</u>	May 28, 2010

GRISEOFULVIN, MICROCRYSTALLINE

TABLET;ORAL

FULVICIN-U/F

<u>AB</u>	CHARTWELL RX	<u>250MG</u>	<u>A060569</u>	<u>002</u>	
<u>AB</u>		<u>500MG</u>	<u>A060569</u>	<u>001</u>	

GRISEOFULVIN, MICROSIZE

SUSPENSION;ORAL

GRISEOFULVIN

<u>AB</u>	! ACTAVIS MID ATLANTIC	<u>125MG/5ML</u>	<u>A065394</u>	<u>001</u>	Jul 06, 2007
<u>AB</u>	CHARTWELL RX	<u>125MG/5ML</u>	<u>A065200</u>	<u>001</u>	Mar 02, 2005
<u>AB</u>	CIPLA	<u>125MG/5ML</u>	<u>A065354</u>	<u>001</u>	Sep 10, 2007
<u>AB</u>	COSETTE	<u>125MG/5ML</u>	<u>A065438</u>	<u>001</u>	Oct 08, 2010

TABLET;ORAL

GRISEOFULVIN

<u>AB</u>	! SANDOZ INC	<u>500MG</u>	<u>A091592</u>	<u>002</u>	Aug 07, 2013
<u>AB</u>	SIGMAPHARM LABS LLC	<u>500MG</u>	<u>A202482</u>	<u>001</u>	Oct 22, 2012
	SANDOZ INC	<u>250MG</u>	<u>A091592</u>	<u>001</u>	Aug 07, 2013

PRESCRIPTION DRUG PRODUCT LIST

GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET;ORAL

FULVICIN P/G

<u>AB</u>	CHARTWELL RX	<u>125MG</u>	<u>A061996</u>	<u>001</u>	
<u>AB</u>		<u>250MG</u>	<u>A061996</u>	<u>002</u>	
<u>FULVICIN P/G 165</u>					
<u>AB</u>	CHARTWELL RX	<u>165MG</u>	<u>A061996</u>	<u>003</u>	Apr 06, 1982
<u>FULVICIN P/G 330</u>					
<u>AB</u>	CHARTWELL RX	<u>330MG</u>	<u>A061996</u>	<u>004</u>	Apr 06, 1982

GRISEOFULVIN, ULTRAMICROSIZED

TABLET;ORAL

GRIS-PEG

<u>AB</u>	+ BAUSCH	<u>125MG</u>	<u>N050475</u>	<u>001</u>	
<u>AB</u>	+	<u>250MG</u>	<u>N050475</u>	<u>002</u>	
<u>GRISEOFULVIN, ULTRAMICROSIZED</u>					
<u>AB</u>	MOUNTAIN	<u>125MG</u>	<u>A204371</u>	<u>001</u>	Jan 09, 2014
<u>AB</u>	!	<u>250MG</u>	<u>A204371</u>	<u>002</u>	Jan 09, 2014
<u>AB</u>	SANDOZ INC	<u>125MG</u>	<u>A202805</u>	<u>001</u>	Dec 26, 2018
<u>AB</u>		<u>250MG</u>	<u>A202805</u>	<u>002</u>	Dec 26, 2018

GRISEOFULVIN, ULTRAMICROSIZED

<u>AB</u>	SIGMAPHARM LABS LLC	<u>125MG</u>	<u>A202545</u>	<u>001</u>	Oct 22, 2012
<u>AB</u>		<u>250MG</u>	<u>A202545</u>	<u>002</u>	Oct 22, 2012

GUANFACINE HYDROCHLORIDE

TABLET;ORAL

GUANFACINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARM	<u>EQ 1MG BASE</u>	<u>A075109</u>	<u>001</u>	Nov 25, 1998
<u>AB</u>	!	<u>EQ 2MG BASE</u>	<u>A075109</u>	<u>002</u>	Nov 25, 1998
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 1MG BASE</u>	<u>A074673</u>	<u>001</u>	Feb 28, 1997
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A074673</u>	<u>002</u>	Feb 28, 1997
<u>AB</u>	UNICHEM	<u>EQ 1MG BASE</u>	<u>A214689</u>	<u>001</u>	Mar 03, 2021
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A214689</u>	<u>002</u>	Mar 03, 2021
<u>AB</u>	WATSON LABS	<u>EQ 1MG BASE</u>	<u>A074145</u>	<u>001</u>	Oct 17, 1995
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A074145</u>	<u>002</u>	Oct 17, 1995

TABLET, EXTENDED RELEASE;ORAL

GUANFACINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 1MG BASE</u>	<u>A200881</u>	<u>001</u>	Oct 05, 2012
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A200881</u>	<u>002</u>	Oct 05, 2012
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A200881</u>	<u>003</u>	Oct 05, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A200881</u>	<u>004</u>	Oct 05, 2012
<u>AB</u>	APOTEX	<u>EQ 1MG BASE</u>	<u>A205430</u>	<u>001</u>	Jul 25, 2018
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A205430</u>	<u>002</u>	Jul 25, 2018
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A205430</u>	<u>003</u>	Jul 25, 2018
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A205430</u>	<u>004</u>	Jul 25, 2018
<u>AB</u>	SANDOZ INC	<u>EQ 1MG BASE</u>	<u>A202568</u>	<u>001</u>	Jun 03, 2015
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A202568</u>	<u>002</u>	Jun 03, 2015
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A202568</u>	<u>003</u>	Jun 03, 2015
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A202568</u>	<u>004</u>	Jun 03, 2015
<u>AB</u>	SUN PHARM	<u>EQ 1MG BASE</u>	<u>A205689</u>	<u>001</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A205689</u>	<u>002</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A205689</u>	<u>003</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A205689</u>	<u>004</u>	Nov 16, 2017
<u>AB</u>	TEVA PHARMS USA	<u>EQ 1MG BASE</u>	<u>A201382</u>	<u>001</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A201382</u>	<u>002</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A201382</u>	<u>003</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A201382</u>	<u>004</u>	Jun 02, 2015
<u>AB</u>	TWI PHARMS	<u>EQ 1MG BASE</u>	<u>A201408</u>	<u>001</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A201408</u>	<u>002</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A201408</u>	<u>003</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A201408</u>	<u>004</u>	Jun 02, 2015
<u>INTUNIV</u>					
<u>AB</u>	+ TAKEDA PHARMS USA	<u>EQ 1MG BASE</u>	<u>N022037</u>	<u>001</u>	Sep 02, 2009
<u>AB</u>	+	<u>EQ 2MG BASE</u>	<u>N022037</u>	<u>002</u>	Sep 02, 2009
<u>AB</u>	+	<u>EQ 3MG BASE</u>	<u>N022037</u>	<u>003</u>	Sep 02, 2009
<u>AB</u>	+	<u>EQ 4MG BASE</u>	<u>N022037</u>	<u>004</u>	Sep 02, 2009

HALCINONIDE

CREAM;TOPICAL

HALCINONIDE

<u>AB</u>	GLASSHOUSE PHARMS	<u>0.1%</u>	<u>A214723</u>	<u>001</u>	Sep 08, 2021
<u>AB</u>	MYLAN	<u>0.1%</u>	<u>A211027</u>	<u>001</u>	Aug 12, 2019

PRESCRIPTION DRUG PRODUCT LIST

HALCINONIDE

CREAM; TOPICAL

HALOG**AB** +! SUN PHARM INDS INC **0.1%** **N017556 001**

OINTMENT; TOPICAL

HALOG

+! SUN PHARM INDS INC 0.1% N017824 001

SOLUTION; TOPICAL

HALOG

+! SUN PHARM INDS INC 0.1% N017823 001

HALOBETASOL PROPIONATE

AEROSOL, FOAM; TOPICAL

LEXETTE

+! MAYNE PHARMA 0.05% N210566 001 May 24, 2018

CREAM; TOPICAL

HALOBETASOL PROPIONATE**AB** ! COSETTE **0.05%** **A078162 001** Apr 24, 2007**AB** FOUGERA PHARMS **0.05%** **A077001 001** Dec 16, 2004**AB** PADAGIS ISRAEL **0.05%** **A077123 001** Dec 16, 2004**AB** TARO **0.05%** **A077227 001** Aug 04, 2005

LOTION; TOPICAL

BRYHALI

+! BAUSCH 0.01% N209355 001 Nov 06, 2018

ULTRAVATE

+! SUN PHARM INDUSTRIES 0.05% N208183 001 Nov 06, 2015

OINTMENT; TOPICAL

HALOBETASOL PROPIONATE**AB** COSETTE **0.05%** **A077109 001** Jun 14, 2005**AB** ! PADAGIS ISRAEL **0.05%** **A076872 001** Dec 16, 2004**AB** QUAGEN **0.05%** **A213560 001** Oct 06, 2020**AB** TARO **0.05%** **A076994 001** Dec 16, 2004**AB** TELIGENT **0.05%** **A209978 001** Mar 20, 2018HALOBETASOL PROPIONATE; TAZAROTENE

LOTION; TOPICAL

DUOBRII

+! BAUSCH 0.01%; 0.045% N209354 001 Apr 25, 2019

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL**AB** CYCLE PHARMS LTD **2MG** **A071130 001** Feb 17, 1987**AB** **5MG** **A071131 001** Feb 17, 1987**AB** **10MG** **A071132 001** May 12, 1987**AB** INNOGENIX **0.5MG** **A071173 002** Jan 02, 1987**AB** **1MG** **A071173 003** Jan 02, 1987**AB** **2MG** **A071173 004** Jan 02, 1987**AB** **5MG** **A071173 005** Jan 07, 1988**AB** **10MG** **A071173 001** Jan 07, 1988**AB** **20MG** **A071173 006** Jan 07, 1988**AB** MYLAN **0.5MG** **A070278 006** Jun 10, 1986**AB** **1MG** **A070278 004** Jun 10, 1986**AB** ! **2MG** **A070278 001** Jun 10, 1986**AB** **5MG** **A070278 005** Jun 10, 1986**AB** **10MG** **A070278 002** Jul 16, 2009**AB** **20MG** **A070278 003** Jul 16, 2009**AB** SANDOZ **0.5MG** **A071206 001** Nov 17, 1986**AB** **1MG** **A071207 001** Nov 17, 1986**AB** **5MG** **A071209 001** Nov 17, 1986**AB** **10MG** **A071210 001** Mar 11, 1988**AB** **20MG** **A071211 001** Mar 11, 1988**AB** UPSHER SMITH LABS **0.5MG** **A211061 001** Jan 08, 2020**AB** **1MG** **A211061 002** Jan 08, 2020**AB** **2MG** **A211061 003** Jan 08, 2020**AB** **5MG** **A211061 004** Jan 08, 2020**AB** **10MG** **A211061 005** Jan 08, 2020**AB** **20MG** **A211061 006** Jan 08, 2020**AB** ZYDUS PHARMS USA **5MG** **A077580 003** Nov 29, 2007**AB** **10MG** **A077580 004** Nov 29, 2007**AB** **20MG** **A077580 005** Nov 29, 2007

PRESCRIPTION DRUG PRODUCT LIST

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALDOL

<u>AO</u>	<u>+!</u>	JANSSEN PHARMS	<u>EQ 50MG BASE/ML</u>	<u>N018701 001</u>	Jan 14, 1986
<u>AO</u>	<u>+!</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>		HIKMA	<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
<u>AO</u>		MEITHEAL	<u>EQ 50MG BASE/ML</u>	<u>A214507 001</u>	Jul 26, 2021
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A214507 002</u>	Jul 26, 2021
<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>		ZYDUS PHARMS	<u>EQ 50MG BASE/ML</u>	<u>A211180 001</u>	Oct 22, 2019
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A211180 002</u>	Oct 22, 2019

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>	<u>!</u>	PHARM ASSOC	<u>EQ 2MG BASE/ML</u>	<u>A073037 001</u>	Feb 26, 1993

INJECTABLE; INJECTION

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>			<u>EQ 5MG BASE/ML</u>	<u>A210356 001</u>	Jul 01, 2019
<u>AP</u>		GLAND PHARMA LTD	<u>EQ 5MG BASE/ML</u>	<u>A076774 001</u>	Aug 25, 2004
<u>AP</u>		HIKMA	<u>EQ 5MG BASE/ML</u>	<u>A075858 001</u>	Jun 18, 2001
<u>AP</u>	<u>!</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

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

<u>AP</u>		BE PHARMS	<u>1,000 UNITS/ML</u>	<u>A214804 001</u>	Dec 29, 2020
<u>AP</u>			<u>10,000 UNITS/ML</u>	<u>A214839 001</u>	Dec 29, 2020
<u>AP</u>	<u>+!</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>+!</u>		<u>5,000 UNITS/ML</u>	<u>N017651 006</u>	
<u>AP</u>	<u>+!</u>		<u>10,000 UNITS/ML</u>	<u>N017029 003</u>	
<u>AP</u>	<u>+!</u>		<u>20,000 UNITS/ML</u>	<u>N017029 004</u>	
<u>AP</u>		GLAND	<u>1,000 UNITS/ML</u>	<u>A205323 002</u>	Nov 18, 2019
<u>AP</u>			<u>5,000 UNITS/ML</u>	<u>A205323 001</u>	Feb 06, 2017
<u>AP</u>	<u>+!</u>	HIKMA	<u>1,000 UNITS/ML</u>	<u>N017037 001</u>	
<u>AP</u>	<u>+!</u>		<u>5,000 UNITS/ML</u>	<u>N017037 002</u>	
<u>AP</u>	<u>+!</u>		<u>10,000 UNITS/ML</u>	<u>N017037 003</u>	
<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>			<u>1,000 UNITS/ML</u>	<u>A211007 001</u>	May 28, 2019
<u>AP</u>			<u>5,000 UNITS/ML</u>	<u>A211007 002</u>	May 28, 2019
<u>AP</u>			<u>10,000 UNITS/ML</u>	<u>A211007 003</u>	May 28, 2019
<u>AP</u>			<u>20,000 UNITS/ML</u>	<u>A211004 001</u>	Feb 24, 2020
<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
<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

PRESCRIPTION DRUG PRODUCT LIST

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

<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>HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
<u>AP</u>		BAXTER HLTHCARE	<u>200 UNITS/100ML</u>	<u>N018609 001</u>	Apr 28, 1982
		<u>HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
<u>AP</u>	+	B BRAUN	<u>200 UNITS/100ML</u>	<u>N019953 001</u>	Jul 20, 1992
<u>AP</u>		FRESENIUS KABI USA	<u>200 UNITS/100ML</u>	<u>A212441 001</u>	Jul 24, 2020
<u>AP</u>	+	HOSPIRA	<u>200 UNITS/100ML</u>	<u>N018916 010</u>	Jun 23, 1989
		<u>HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u>		HOSPIRA	<u>10,000 UNITS/100ML</u>	<u>N019339 003</u>	Mar 27, 1985
		<u>HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
<u>AP</u>		BAXTER HLTHCARE	<u>200 UNITS/100ML</u>	<u>N018609 002</u>	Apr 28, 1982
		<u>HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
<u>AP</u>		FRESENIUS KABI USA	<u>200 UNITS/100ML</u>	<u>A212441 002</u>	Jul 24, 2020
<u>AP</u>	+	HOSPIRA	<u>200 UNITS/100ML</u>	<u>N018916 011</u>	Jun 23, 1989
		<u>HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<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
		<u>HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<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
		<u>HEPARIN SODIUM IN PLASTIC CONTAINER</u>			
<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
		<u>HEPARIN SODIUM PRESERVATIVE FREE</u>			
<u>AP</u>	+	FRESENIUS KABI USA	<u>1,000 UNITS/ML</u>	<u>N017029 010</u>	Apr 28, 1986
<u>AP</u>	+		<u>10,000 UNITS/ML</u>	<u>N017029 019</u>	Nov 22, 2010
<u>AP</u>		HOSPIRA	<u>10,000 UNITS/ML</u>	<u>A089522 001</u>	May 04, 1987
<u>AP</u>		SAGENT PHARMS	<u>1,000 UNITS/ML</u>	<u>A090810 001</u>	Jun 30, 2010
		<u>HEPARIN SODIUM</u>			
		B BRAUN MEDICAL INC	5,000 UNITS/0.5ML	A208827 001	Nov 19, 2018
	+	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
		<u>HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
		HOSPIRA	5,000 UNITS/100ML	N019339 001	Mar 27, 1985
		<u>HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
		HOSPIRA	5,000 UNITS/100ML	N018916 006	Jan 31, 1984
		<u>HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
		BAXTER HLTHCARE	2,000 UNITS/1000ML	N018609 004	Apr 23, 2020
		<u>HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
		HOSPIRA	5,000 UNITS/100ML	N018916 007	Jan 31, 1984
			10,000 UNITS/100ML	N018916 008	Jan 31, 1984
		<u>HEPARIN SODIUM PRESERVATIVE FREE</u>			
	+	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>	
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PRE-OP II

<u>AT</u>	+	DAVIS AND GECK	<u>480MG</u>	<u>N017433 002</u>	
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HEXAMINOLEVULINATE HYDROCHLORIDE

FOR SOLUTION; INTRAVESICAL

CYSVIEW KIT

	+	PHOTOCURE ASA	100MG/VIAL	N022555 001	May 28, 2010
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PRESCRIPTION DRUG PRODUCT LIST

HISTRELIN ACETATE

IMPLANT; SUBCUTANEOUS

SUPPRELIN LA

+! ENDO PHARM

50MG

N022058 001 May 03, 2007

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYCODANAA + GENUS1.5MG/5ML; 5MG/5MLN005213 002 Jul 26, 1988HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDEAA ABHAI LLC1.5MG/5ML; 5MG/5MLA207487 001 Feb 21, 2017AA ACTAVIS MID1.5MG/5ML; 5MG/5MLA088017 001 Jul 05, 1983

ATLANTIC

AA ! AKORN1.5MG/5ML; 5MG/5MLA040613 001 Feb 08, 2008AA NOVEL LABS INC1.5MG/5ML; 5MG/5MLA203535 001 Feb 13, 2017AA PADAGIS US1.5MG/5ML; 5MG/5MLA205731 001 Feb 15, 2017AA WOCKHARDT BIO AG1.5MG/5ML; 5MG/5MLA088008 001 Mar 03, 1983

TABLET; ORAL

HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATEAA AVANTHI INC1.5MG; 5MGA207176 001 Aug 07, 2017AA ! NOVEL LABS INC1.5MG; 5MGA091528 001 Apr 20, 2011HYCODANAA + GENUS1.5MG; 5MGN005213 001 Jul 26, 1988HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDRALAZINE HYDROCHLORIDEAP ! AKORN20MG/MLA040730 001 Apr 21, 2009AP AM REGENT20MG/MLA040136 001 Jun 30, 1997AP FRESENIUS KABI USA20MG/MLA040388 001 Mar 13, 2001AP HIKMA20MG/MLA213667 001 Dec 18, 2020AP I3 PHARMS20MG/MLA203110 001 Jun 29, 2015AP MYLAN INSTITUTIONAL20MG/MLA204680 001 Apr 28, 2016AP NAVINTA LLC20MG/MLA202938 001 Mar 28, 2013

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDEAA ALKEM LABS LTD10MGA200737 001 Dec 07, 2012AA25MGA200737 002 Dec 07, 2012AA50MGA200737 003 Dec 07, 2012AA100MGA200737 004 Dec 07, 2012AA

CADILA PHARMS LTD

25MGA203845 001 Sep 18, 2014AA50MGA203845 002 Sep 18, 2014AA100MGA203845 003 Sep 18, 2014AA

GLENMARK PHARMS LTD

10MGA090527 001 May 27, 2009AA25MGA090527 002 May 27, 2009AA50MGA090527 003 May 27, 2009AA100MGA090527 004 May 27, 2009AA

HERITAGE PHARMS INC

10MGA086242 001 Feb 04, 2010AA25MGA086242 003AA50MGA086242 002AA100MGA086242 004 Feb 04, 2010AA

HETERO LABS LTD III

10MGA040901 001 Sep 12, 2008AA25MGA040901 002 Sep 12, 2008AA50MGA040901 003 Sep 12, 2008AA100MGA040901 004 Sep 12, 2008AA

INVAGEN PHARMS

10MGA090255 001 Dec 15, 2008AA25MGA090255 002 Dec 15, 2008AA50MGA090255 003 Dec 15, 2008AA100MGA090255 004 Dec 15, 2008AA ! PLIVA10MGA089097 001 Dec 18, 1985AA +!25MGA088467 001 May 01, 1984AA +!50MGA088468 001 May 01, 1984AA !100MGA089098 001 Dec 18, 1985AA

SCIEGEN PHARMS INC

10MGA205236 001 May 26, 2017AA25MGA205236 002 May 26, 2017AA50MGA205236 003 May 26, 2017AA100MGA205236 004 May 26, 2017AA

STRIDES PHARMA

10MGA087836 001 Oct 05, 1982AA25MGA086961 002AA25MGA200770 001 May 03, 2013AA50MGA086962 001AA50MGA200770 002 May 03, 2013AA100MGA088391 001 Sep 27, 1983AA100MGA200770 003 May 03, 2013

PRESCRIPTION DRUG PRODUCT LIST

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDRA-ZIDE

STRIDES PHARMA	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
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HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

<u>AB</u>	AUROBINDO PHARMA	<u>12.5MG</u>	<u>A078164</u>	<u>001</u>	Sep 18, 2007
<u>AB</u>	IPCA LABS LTD	<u>12.5MG</u>	<u>A079237</u>	<u>001</u>	Apr 02, 2009
<u>AB</u>	JUBILANT CADISTA	<u>12.5MG</u>	<u>A078391</u>	<u>001</u>	Feb 11, 2008
<u>AB</u>	MYLAN	<u>12.5MG</u>	<u>A075640</u>	<u>001</u>	Jan 28, 2000
<u>AB</u>	PRINSTON INC	<u>12.5MG</u>	<u>A075907</u>	<u>001</u>	Sep 17, 2002
<u>AB</u>	SCIEGEN PHARMS INC	<u>12.5MG</u>	<u>A203561</u>	<u>001</u>	Jan 14, 2019
<u>AB</u>	SUN PHARM INDS INC	<u>12.5MG</u>	<u>A090651</u>	<u>001</u>	Apr 07, 2014
<u>AB</u>	UNICHEM	<u>12.5MG</u>	<u>A090510</u>	<u>001</u>	Jan 19, 2010

MICROZIDE

<u>AB</u>	+! TEVA BRANDED PHARM	<u>12.5MG</u>	<u>N020504</u>	<u>001</u>	Dec 27, 1996
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TABLET; ORAL

HYDROCHLOROTHIAZIDE

<u>AB</u>	!	ACCORD HLTHCARE	<u>12.5MG</u>	<u>A202556</u>	<u>001</u>	Sep 24, 2012
<u>AB</u>			<u>25MG</u>	<u>A202556</u>	<u>002</u>	Sep 24, 2012
<u>AB</u>			<u>50MG</u>	<u>A202556</u>	<u>003</u>	Sep 24, 2012
<u>AB</u>		ACTAVIS ELIZABETH	<u>12.5MG</u>	<u>A040707</u>	<u>001</u>	Feb 27, 2007
<u>AB</u>		AUROBINDO PHARMA	<u>25MG</u>	<u>A040780</u>	<u>001</u>	Jul 20, 2007
<u>AB</u>			<u>50MG</u>	<u>A040780</u>	<u>002</u>	Jul 20, 2007
<u>AB</u>		HERITAGE PHARMS INC	<u>25MG</u>	<u>A085182</u>	<u>002</u>	
<u>AB</u>			<u>50MG</u>	<u>A085182</u>	<u>001</u>	
<u>AB</u>		IPCA LABS LTD	<u>12.5MG</u>	<u>A040807</u>	<u>001</u>	Jul 20, 2007
<u>AB</u>			<u>25MG</u>	<u>A040807</u>	<u>002</u>	Jul 20, 2007
<u>AB</u>			<u>50MG</u>	<u>A040807</u>	<u>003</u>	Jul 20, 2007
<u>AB</u>	+	IVAX SUB TEVA PHARMS	<u>25MG</u>	<u>A083177</u>	<u>001</u>	
<u>AB</u>	+!		<u>50MG</u>	<u>A083177</u>	<u>002</u>	
<u>AB</u>		LEADING PHARMA LLC	<u>12.5MG</u>	<u>A040702</u>	<u>003</u>	May 10, 2017
<u>AB</u>			<u>25MG</u>	<u>A040702</u>	<u>001</u>	Mar 16, 2007
<u>AB</u>			<u>50MG</u>	<u>A040702</u>	<u>002</u>	Mar 16, 2007
<u>AB</u>		OXFORD PHARMS	<u>25MG</u>	<u>A087059</u>	<u>001</u>	
<u>AB</u>			<u>50MG</u>	<u>A087068</u>	<u>001</u>	
<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>		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>	+!	SANOFI AVENTIS US	<u>12.5MG; 150MG</u>	<u>N020758</u>	<u>002</u>	Sep 30, 1997
<u>AB</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>		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>		HIKMA	<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
<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

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

IRBESARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>		<u>12.5MG;300MG</u>	<u>A201524 002</u>	Feb 27, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>12.5MG;150MG</u>	<u>A202414 001</u>	Sep 27, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A202414 002</u>	Sep 27, 2012
<u>AB</u>	PRINSTON INC	<u>12.5MG;150MG</u>	<u>A203072 001</u>	May 09, 2014
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A203072 002</u>	May 09, 2014
<u>AB</u>	SANDOZ	<u>12.5MG;150MG</u>	<u>A077446 001</u>	Sep 27, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A077446 002</u>	Sep 27, 2012
<u>AB</u>	TEVA	<u>12.5MG;150MG</u>	<u>A077369 001</u>	Mar 30, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A077369 002</u>	Mar 30, 2012
<u>AB</u>	UNICHEM	<u>12.5MG;150MG</u>	<u>A207018 001</u>	Sep 19, 2017
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A207018 002</u>	Sep 19, 2017

HYDROCHLOROTHIAZIDE; LISINAPRIL

TABLET; ORAL

LISINAPRIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>	AUROBINDO	<u>12.5MG;10MG</u>	<u>A077606 001</u>	Mar 14, 2006
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A077606 002</u>	Mar 14, 2006
<u>AB</u>		<u>25MG;20MG</u>	<u>A077606 003</u>	Mar 14, 2006
<u>AB</u>	INVAGEN PHARMS	<u>12.5MG;10MG</u>	<u>A204058 001</u>	May 23, 2017
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A204058 002</u>	May 23, 2017
<u>AB</u>		<u>25MG;20MG</u>	<u>A204058 003</u>	May 23, 2017
<u>AB</u>	LUPIN	<u>12.5MG;10MG</u>	<u>A077912 001</u>	Sep 27, 2006
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A077912 002</u>	Sep 27, 2006
<u>AB</u>		<u>25MG;20MG</u>	<u>A077912 003</u>	Sep 27, 2006
<u>AB</u>	PRINSTON INC	<u>12.5MG;10MG</u>	<u>A076230 001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076230 002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076230 003</u>	Jul 01, 2002
<u>AB</u>	SANDOZ	<u>12.5MG;10MG</u>	<u>A076262 001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076262 002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076262 003</u>	Jul 01, 2002
<u>AB</u>	SUN PHARM INDS LTD	<u>12.5MG;10MG</u>	<u>A076007 001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076007 002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076007 003</u>	Jul 01, 2002
<u>AB</u>	WATSON LABS	<u>12.5MG;10MG</u>	<u>A076194 003</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076194 001</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076194 002</u>	Jul 01, 2002

ZESTORETIC

<u>AB</u>	+ ALMATICA	<u>12.5MG;10MG</u>	<u>N019888 003</u>	Nov 18, 1993
<u>AB</u>	+!	<u>12.5MG;20MG</u>	<u>N019888 001</u>	Sep 20, 1990
<u>AB</u>	+!	<u>25MG;20MG</u>	<u>N019888 002</u>	Jul 20, 1989

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET; ORAL

HYZAAR

<u>AB</u>	+ ORGANON	<u>12.5MG;50MG</u>	<u>N020387 001</u>	Apr 28, 1995
<u>AB</u>	+	<u>12.5MG;100MG</u>	<u>N020387 003</u>	Oct 20, 2005
<u>AB</u>	+	<u>25MG;100MG</u>	<u>N020387 002</u>	Nov 10, 1998

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>12.5MG;50MG</u>	<u>A091617 001</u>	Feb 17, 2012
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A091617 002</u>	Feb 17, 2012
<u>AB</u>		<u>25MG;100MG</u>	<u>A091617 003</u>	Feb 17, 2012
<u>AB</u>	AUROBINDO PHARMA	<u>12.5MG;50MG</u>	<u>A091629 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A091629 002</u>	Oct 06, 2010
<u>AB</u>	!	<u>25MG;100MG</u>	<u>A091629 003</u>	Jan 06, 2010
<u>AB</u>	IPCA LABS LTD	<u>12.5MG;50MG</u>	<u>A201682 001</u>	Mar 01, 2013
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A201682 002</u>	Mar 01, 2013
<u>AB</u>		<u>25MG;100MG</u>	<u>A201682 003</u>	Mar 01, 2013
<u>AB</u>	JUBILANT CADISTA	<u>12.5MG;50MG</u>	<u>A201845 001</u>	Sep 18, 2012
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A201845 002</u>	Sep 18, 2012
<u>AB</u>		<u>25MG;100MG</u>	<u>A201845 003</u>	Sep 18, 2012
<u>AB</u>	LUPIN LTD	<u>12.5MG;50MG</u>	<u>A078245 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A078245 002</u>	May 21, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A078245 003</u>	Oct 06, 2010
<u>AB</u>	MACLEODS PHARMS LTD	<u>12.5MG;50MG</u>	<u>A202289 001</u>	Aug 09, 2012
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A202289 002</u>	Aug 09, 2012
<u>AB</u>		<u>25MG;100MG</u>	<u>A202289 003</u>	Aug 09, 2012
<u>AB</u>	PRINSTON INC	<u>12.5MG;50MG</u>	<u>A204901 001</u>	Nov 06, 2017
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A204901 002</u>	Nov 06, 2017
<u>AB</u>		<u>25MG;100MG</u>	<u>A204901 003</u>	Nov 06, 2017
<u>AB</u>	SANDOZ	<u>12.5MG;50MG</u>	<u>A077948 001</u>	Oct 06, 2010

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE

<u>AB</u>		<u>12.5MG;100MG</u>	<u>A077948 003</u>	Aug 19, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A077948 002</u>	Oct 06, 2010
<u>AB</u>	TEVA PHARMS	<u>12.5MG;50MG</u>	<u>A077157 001</u>	Apr 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A077157 002</u>	Apr 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A077157 003</u>	Apr 06, 2010
<u>AB</u>	UNICHEM	<u>12.5MG;50MG</u>	<u>A204832 001</u>	Jul 21, 2017
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A204832 002</u>	Jul 21, 2017
<u>AB</u>		<u>25MG;100MG</u>	<u>A204832 003</u>	Jul 21, 2017
<u>AB</u>	ZYDUS PHARMS USA INC	<u>12.5MG;50MG</u>	<u>A078385 001</u>	Oct 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A078385 002</u>	Oct 06, 2010

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

LOPRESSOR HCT

<u>AB</u>	+	VALIDUS PHARMS	<u>25MG;50MG</u>	<u>N018303 001</u>	Dec 31, 1984
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METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>25MG;50MG</u>	<u>A202870 001</u>	Nov 06, 2013
<u>AB</u>	!		<u>25MG;100MG</u>	<u>A202870 002</u>	Nov 06, 2013
<u>AB</u>			<u>50MG;100MG</u>	<u>A202870 003</u>	Nov 06, 2013
<u>AB</u>		MYLAN	<u>25MG;50MG</u>	<u>A076792 001</u>	Aug 20, 2004
<u>AB</u>			<u>25MG;100MG</u>	<u>A076792 002</u>	Aug 20, 2004
<u>AB</u>			<u>50MG;100MG</u>	<u>A076792 003</u>	Aug 20, 2004
<u>AB</u>		SUN PHARM INDS	<u>25MG;50MG</u>	<u>A090654 001</u>	Jan 19, 2012
<u>AB</u>			<u>25MG;100MG</u>	<u>A090654 002</u>	Jan 19, 2012
<u>AB</u>			<u>50MG;100MG</u>	<u>A090654 003</u>	Jan 19, 2012

HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>		GLENMARK PHARMS	<u>12.5MG;7.5MG</u>	<u>A090718 001</u>	Mar 17, 2010
<u>AB</u>			<u>12.5MG;15MG</u>	<u>A090718 002</u>	Mar 17, 2010
<u>AB</u>			<u>25MG;15MG</u>	<u>A090718 003</u>	Mar 17, 2010
<u>AB</u>		TEVA	<u>12.5MG;7.5MG</u>	<u>A076980 001</u>	Mar 07, 2007
<u>AB</u>			<u>12.5MG;15MG</u>	<u>A076980 003</u>	Mar 07, 2007
<u>AB</u>	!		<u>25MG;15MG</u>	<u>A076980 002</u>	Mar 07, 2007

HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

BENICAR HCT

<u>AB</u>	+	DAIICHI SANKYO	<u>12.5MG;20MG</u>	<u>N021532 002</u>	Jun 05, 2003
<u>AB</u>	+		<u>12.5MG;40MG</u>	<u>N021532 003</u>	Jun 05, 2003
<u>AB</u>	+	!	<u>25MG;40MG</u>	<u>N021532 005</u>	Jun 05, 2003

OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>		ACCORD HLTHCARE	<u>12.5MG;20MG</u>	<u>A209281 001</u>	Feb 07, 2019
<u>AB</u>			<u>12.5MG;40MG</u>	<u>A209281 002</u>	Feb 07, 2019
<u>AB</u>			<u>25MG;40MG</u>	<u>A209281 003</u>	Feb 07, 2019
<u>AB</u>		ALEMBIC PHARMS LTD	<u>12.5MG;20MG</u>	<u>A204233 001</u>	Apr 24, 2017
<u>AB</u>			<u>12.5MG;40MG</u>	<u>A204233 002</u>	Apr 24, 2017
<u>AB</u>			<u>25MG;40MG</u>	<u>A204233 003</u>	Apr 24, 2017
<u>AB</u>		AUROBINDO PHARMA LTD	<u>12.5MG;20MG</u>	<u>A205391 001</u>	Apr 24, 2017
<u>AB</u>			<u>12.5MG;40MG</u>	<u>A205391 002</u>	Apr 24, 2017
<u>AB</u>			<u>25MG;40MG</u>	<u>A205391 003</u>	Apr 24, 2017
<u>AB</u>		PRINSTON INC	<u>12.5MG;20MG</u>	<u>A207804 001</u>	Apr 24, 2017
<u>AB</u>			<u>12.5MG;40MG</u>	<u>A207804 002</u>	Apr 24, 2017
<u>AB</u>			<u>25MG;40MG</u>	<u>A207804 003</u>	Apr 24, 2017
<u>AB</u>		TEVA PHARMS USA	<u>12.5MG;40MG</u>	<u>A200532 002</u>	Apr 24, 2017
<u>AB</u>		TORRENT	<u>12.5MG;20MG</u>	<u>A206515 001</u>	May 03, 2017
<u>AB</u>			<u>12.5MG;40MG</u>	<u>A206515 002</u>	May 03, 2017
<u>AB</u>			<u>25MG;40MG</u>	<u>A206515 003</u>	May 03, 2017
<u>AB</u>		UMEDICA LABS PVT LTD	<u>12.5MG;20MG</u>	<u>A208847 001</u>	Sep 17, 2019
<u>AB</u>			<u>12.5MG;40MG</u>	<u>A208847 003</u>	Sep 17, 2019
<u>AB</u>			<u>25MG;40MG</u>	<u>A208847 002</u>	Sep 17, 2019

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

ACCURETIC

<u>AB</u>	+	PFIZER PHARMS	<u>12.5MG;EQ 10MG BASE</u>	<u>N020125 001</u>	Dec 28, 1999
<u>AB</u>	+		<u>12.5MG;EQ 20MG BASE</u>	<u>N020125 002</u>	Dec 28, 1999
<u>AB</u>	+	!	<u>25MG;EQ 20MG BASE</u>	<u>N020125 003</u>	Dec 28, 1999

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>		APOTEX CORP	<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>		AUROBINDO PHARMA	<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>		INVAGEN PHARMS	<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

QUINARETIC

<u>AB</u>		LUPIN	<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>	+	PFIZER	<u>25MG;25MG</u>	<u>N012616 004</u>	Dec 30, 1982
<u>SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE</u>					
<u>AB</u>		MYLAN	<u>25MG;25MG</u>	<u>A086513 001</u>	
<u>AB</u>		SUN PHARM INDUSTRIES	<u>25MG;25MG</u>	<u>A089534 001</u>	Jul 02, 1987
<u>ALDACTAZIDE</u>					
	+	PFIZER	<u>50MG;50MG</u>	<u>N012616 005</u>	Dec 30, 1982

HYDROCHLOROTHIAZIDE; TELMISARTAN

TABLET; ORAL

MICARDIS HCT

<u>AB</u>	+	BOEHRINGER INGELHEIM	<u>12.5MG;40MG</u>	<u>N021162 001</u>	Nov 17, 2000
<u>AB</u>	+		<u>12.5MG;80MG</u>	<u>N021162 002</u>	Nov 17, 2000
<u>AB</u>	+	!	<u>25MG;80MG</u>	<u>N021162 003</u>	Apr 19, 2004

TELMISARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>		ALEMBIC PHARMS LTD	<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>		AUROBINDO PHARMA LTD	<u>12.5MG;40MG</u>	<u>A208727 001</u>	Dec 15, 2016
<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>		GLENMARK PHARMS LTD	<u>12.5MG;40MG</u>	<u>A202544 001</u>	Mar 04, 2019
<u>AB</u>			<u>12.5MG;80MG</u>	<u>A202544 002</u>	Mar 04, 2019
<u>AB</u>			<u>25MG;80MG</u>	<u>A202544 003</u>	Mar 04, 2019
<u>AB</u>		LUPIN LTD	<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>		PRINSTON INC	<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>		ZYDUS PHARMS	<u>12.5MG;40MG</u>	<u>A204221 001</u>	Aug 15, 2017
<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

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

<u>AB</u>		CADILA	<u>25MG;37.5MG</u>	<u>A208358 001</u>	Feb 11, 2019
<u>AB</u>		LANNETT CO INC	<u>25MG;37.5MG</u>	<u>A201407 001</u>	Dec 09, 2011
<u>AB</u>		MYLAN	<u>25MG;37.5MG</u>	<u>A074701 001</u>	Jun 07, 1996
<u>AB</u>	!	SANDOZ	<u>25MG;37.5MG</u>	<u>A074821 001</u>	Jun 05, 1997

TABLET; ORAL

MAXZIDE

<u>AB</u>	+	!	MYLAN PHARMS INC	<u>50MG;75MG</u>	<u>N019129 001</u>	Oct 22, 1984
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MAXZIDE-25

<u>AB</u>	+	MYLAN PHARMS INC	<u>25MG;37.5MG</u>	<u>N019129 003</u>	May 13, 1988
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TRIAMTERENE AND HYDROCHLOROTHIAZIDE

<u>AB</u>		APOTEX INC	<u>25MG;37.5MG</u>	<u>A071251 002</u>	May 05, 1998
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PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; TRIAMTERENE

TABLET; ORAL

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

<u>AB</u>		<u>50MG; 75MG</u>	<u>A071251 001</u>	Apr 17, 1988
<u>AB</u>	SANDOZ	<u>25MG; 37.5MG</u>	<u>A073281 001</u>	Apr 30, 1992
<u>AB</u>		<u>50MG; 75MG</u>	<u>A072011 001</u>	Jun 17, 1988
<u>AB</u>	WATSON LABS	<u>25MG; 37.5MG</u>	<u>A073449 001</u>	Sep 23, 1993
<u>AB</u>		<u>50MG; 75MG</u>	<u>A071851 001</u>	Nov 30, 1988
<u>AB</u>	ZYDUS PHARMS	<u>25MG; 37.5MG</u>	<u>A208360 001</u>	Jun 29, 2018
<u>AB</u>		<u>50MG; 75MG</u>	<u>A208360 002</u>	Jun 29, 2018

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

DIOVAN HCT

<u>AB</u>	+	NOVARTIS	<u>12.5MG; 80MG</u>	<u>N020818 001</u>	Mar 06, 1998
<u>AB</u>	+		<u>12.5MG; 160MG</u>	<u>N020818 002</u>	Mar 06, 1998
<u>AB</u>	+		<u>12.5MG; 320MG</u>	<u>N020818 004</u>	Apr 28, 2006
<u>AB</u>	+		<u>25MG; 160MG</u>	<u>N020818 003</u>	Jan 17, 2002
<u>AB</u>	+	!	<u>25MG; 320MG</u>	<u>N020818 005</u>	Apr 28, 2006

VALSARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>12.5MG; 80MG</u>	<u>A201662 001</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG; 160MG</u>	<u>A201662 002</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG; 320MG</u>	<u>A201662 003</u>	Mar 21, 2013
<u>AB</u>			<u>25MG; 160MG</u>	<u>A201662 004</u>	Mar 21, 2013
<u>AB</u>			<u>25MG; 320MG</u>	<u>A201662 005</u>	Mar 21, 2013
<u>AB</u>		AUROBINDO PHARMA LTD	<u>12.5MG; 80MG</u>	<u>A202519 001</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG; 160MG</u>	<u>A202519 002</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG; 320MG</u>	<u>A202519 003</u>	Mar 21, 2013
<u>AB</u>			<u>25MG; 160MG</u>	<u>A202519 004</u>	Mar 21, 2013
<u>AB</u>			<u>25MG; 320MG</u>	<u>A202519 005</u>	Mar 21, 2013
<u>AB</u>		LUPIN LTD	<u>12.5MG; 80MG</u>	<u>A078946 003</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG; 160MG</u>	<u>A078946 004</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG; 320MG</u>	<u>A078946 001</u>	Mar 21, 2013
<u>AB</u>			<u>25MG; 160MG</u>	<u>A078946 005</u>	Mar 21, 2013
<u>AB</u>			<u>25MG; 320MG</u>	<u>A078946 002</u>	Mar 21, 2013
<u>AB</u>		MACLEODS PHARMS LTD	<u>12.5MG; 80MG</u>	<u>A203145 001</u>	Apr 19, 2013
<u>AB</u>			<u>12.5MG; 160MG</u>	<u>A203145 002</u>	Apr 19, 2013
<u>AB</u>			<u>12.5MG; 320MG</u>	<u>A203145 003</u>	Apr 19, 2013
<u>AB</u>			<u>25MG; 160MG</u>	<u>A203145 004</u>	Apr 19, 2013
<u>AB</u>			<u>25MG; 320MG</u>	<u>A203145 005</u>	Apr 19, 2013
<u>AB</u>		MYLAN PHARMS INC	<u>12.5MG; 80MG</u>	<u>A078020 001</u>	Sep 21, 2012
<u>AB</u>			<u>12.5MG; 160MG</u>	<u>A078020 002</u>	Sep 21, 2012
<u>AB</u>			<u>12.5MG; 320MG</u>	<u>A078020 004</u>	Sep 21, 2012
<u>AB</u>			<u>25MG; 160MG</u>	<u>A078020 003</u>	Sep 21, 2012
<u>AB</u>			<u>25MG; 320MG</u>	<u>A078020 005</u>	Sep 21, 2012
<u>AB</u>		PRINSTON INC	<u>12.5MG; 80MG</u>	<u>A206083 001</u>	Feb 08, 2016
<u>AB</u>			<u>12.5MG; 160MG</u>	<u>A206083 002</u>	Feb 08, 2016
<u>AB</u>			<u>12.5MG; 320MG</u>	<u>A206083 003</u>	Feb 08, 2016
<u>AB</u>			<u>25MG; 160MG</u>	<u>A206083 004</u>	Feb 08, 2016
<u>AB</u>			<u>25MG; 320MG</u>	<u>A206083 005</u>	Feb 08, 2016

HYDROCODONE BITARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

HYDROCODONE BITARTRATE

! ALVOGEN

10MG	A206986 001	Jan 21, 2020
15MG	A206986 002	Jan 21, 2020
20MG	A206986 003	Jan 21, 2020
30MG	A206986 004	Jan 21, 2020
40MG	A206986 005	Jan 21, 2020
50MG	A206986 006	Jan 21, 2020

TABLET, EXTENDED RELEASE; ORAL

HYDROCODONE BITARTRATE

<u>AB</u>		ALVOGEN PINE BROOK	<u>20MG</u>	<u>A208269 001</u>	Mar 01, 2021
<u>AB</u>			<u>30MG</u>	<u>A208269 002</u>	Mar 01, 2021
<u>AB</u>			<u>40MG</u>	<u>A208269 003</u>	Mar 01, 2021
<u>AB</u>			<u>60MG</u>	<u>A208269 004</u>	Mar 01, 2021
<u>AB</u>			<u>80MG</u>	<u>A208269 005</u>	Mar 01, 2021
<u>AB</u>			<u>100MG</u>	<u>A208269 006</u>	Mar 01, 2021
<u>AB</u>			<u>120MG</u>	<u>A208269 007</u>	Mar 01, 2021

HYSINGLA ER

<u>AB</u>	+	!	PURDUE PHARMA LP	<u>20MG</u>	<u>N206627 001</u>	Nov 20, 2014
<u>AB</u>	+			<u>30MG</u>	<u>N206627 002</u>	Nov 20, 2014
<u>AB</u>	+			<u>40MG</u>	<u>N206627 003</u>	Nov 20, 2014

PRESCRIPTION DRUG PRODUCT LIST

HYDROCODONE BITARTRATE

TABLET, EXTENDED RELEASE;ORAL

HYSINGLA ER

<u>AB</u>	+	<u>60MG</u>	<u>N206627</u>	<u>004</u>	Nov 20, 2014
<u>AB</u>	+	<u>80MG</u>	<u>N206627</u>	<u>005</u>	Nov 20, 2014
<u>AB</u>	+	<u>100MG</u>	<u>N206627</u>	<u>006</u>	Nov 20, 2014
<u>AB</u>	+	<u>120MG</u>	<u>N206627</u>	<u>007</u>	Nov 20, 2014

HYDROCODONE BITARTRATE; IBUPROFEN

TABLET;ORAL

HYDROCODONE

<u>AB</u>		NOSTRUM LABS INC	<u>5MG;200MG</u>	<u>A077723</u>	<u>003</u>	Nov 06, 2006
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HYDROCODONE BITARTRATE AND IBUPROFEN

<u>AB</u>		ACTAVIS LABS FL INC	<u>7.5MG;200MG</u>	<u>A076604</u>	<u>001</u>	Dec 31, 2003
<u>AB</u>		AMNEAL PHARMS NY	<u>5MG;200MG</u>	<u>A076642</u>	<u>002</u>	Mar 18, 2004
<u>AB</u>	!		<u>7.5MG;200MG</u>	<u>A076642</u>	<u>001</u>	Oct 12, 2004
<u>AB</u>		AUROLIFE PHARMA LLC	<u>7.5MG;200MG</u>	<u>A204575</u>	<u>001</u>	Jun 02, 2016
<u>AB</u>		NOSTRUM LABS INC	<u>7.5MG;200MG</u>	<u>A077723</u>	<u>001</u>	Nov 06, 2006
			10MG;200MG	A077723	002	Nov 06, 2006

HYDROCORTISONE

CREAM;TOPICAL

ALA-CORT

<u>AT</u>		CROWN LABS	<u>2.5%</u>	<u>A080706</u>	<u>007</u>	Jan 05, 2016
<u>AT</u>			<u>1%</u>	<u>A080706</u>	<u>006</u>	

ANUSOL HC

<u>AT</u>		SALIX PHARMS	<u>2.5%</u>	<u>A088250</u>	<u>001</u>	Jun 06, 1984
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HYDROCORTISONE

<u>AT</u>		ACTAVIS MID ATLANTIC	<u>1%</u>	<u>A087795</u>	<u>001</u>	May 03, 1983
<u>AT</u>			<u>2.5%</u>	<u>A089682</u>	<u>001</u>	Mar 10, 1988
<u>AT</u>	!	FOUGERA PHARMS INC	<u>1%</u>	<u>A080693</u>	<u>003</u>	
<u>AT</u>	!		<u>2.5%</u>	<u>A089414</u>	<u>001</u>	Dec 16, 1986
<u>AT</u>		LANNETT CO INC	<u>2.5%</u>	<u>A040503</u>	<u>001</u>	Mar 12, 2004
<u>AT</u>		PADAGIS US	<u>2.5%</u>	<u>A085025</u>	<u>001</u>	
<u>AT</u>		RISING	<u>2.5%</u>	<u>A040879</u>	<u>001</u>	Aug 20, 2010
<u>AT</u>		TARO PHARM INDS LTD	<u>2.5%</u>	<u>A088799</u>	<u>001</u>	Nov 09, 1984

ENEMA;RECTAL

COLOCORT

<u>AB</u>		CHARTWELL	<u>100MG/60ML</u>	<u>A075172</u>	<u>001</u>	Dec 03, 1999
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CORTENEMA

<u>AB</u>	+	ANI PHARMS	<u>100MG/60ML</u>	<u>N016199</u>	<u>001</u>	
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GRANULE;ORAL

ALKINDI SPRINKLE

	+	ETON	0.5MG	N213876	001	Sep 29, 2020
	+		1MG	N213876	002	Sep 29, 2020
	+		2MG	N213876	003	Sep 29, 2020
	+	!	5MG	N213876	004	Sep 29, 2020

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

STIE-CORT

<u>AT</u>	!	PADAGIS US	<u>2.5%</u>	<u>A089074</u>	<u>001</u>	Nov 26, 1985
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ALA-SCALP

		MARNEL PHARMS	2%	A083231	001	
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OINTMENT;TOPICAL

HYDROCORTISONE

<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>		PADAGIS US	<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
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SOLUTION;TOPICAL

TEXACORT

	!	MISSION PHARMA	2.5%	A081271	001	Apr 17, 1992
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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>	

PRESCRIPTION DRUG PRODUCT LIST

HYDROCORTISONE

TABLET; ORAL

HYDROCORTISONE

AB	IMPAX LABS INC	5MG	A040646 001	Mar 30, 2007
AB		10MG	A040646 002	Mar 30, 2007
AB		20MG	A040646 003	Mar 30, 2007
AB	STRIDES PHARMA	5MG	A207029 001	Apr 27, 2017
AB		10MG	A207029 002	Apr 27, 2017
AB		20MG	A207029 003	Apr 27, 2017

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

U-CORT

TARO 1%

A089472 001 Jun 13, 1988

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

HYDROCORTISONE BUTYRATE

CREAM; TOPICAL

HYDROCORTISONE BUTYRATE**AB1** TARO PHARM INDS **0.1%****A076654 001** Aug 03, 2005LOCOID**AB1** +! BAUSCH **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 **0.1%****A209556 001** Nov 21, 2017LOCOID**AB** +! BAUSCH **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** +! BAUSCH **0.1%****N019116 001** Feb 25, 1987HYDROCORTISONE PROBUTATE

CREAM; TOPICAL

PANDEL

+! ANI PHARMS 0.1%

N020453 001 Feb 28, 1997

PRESCRIPTION DRUG PRODUCT LIST

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION

SOLU-CORTEF

+	!	PHARMACIA AND UPJOHN	EQ 100MG BASE/VIAL	N009866	001	
+	!		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		COSETTE	<u>0.2%</u>	<u>A213724</u>	<u>001</u>	Feb 11, 2021
AB		ENCUBE ETHICALS	<u>0.2%</u>	<u>A211047</u>	<u>001</u>	Nov 04, 2021
AB		GLENMARK PHARMS LTD	<u>0.2%</u>	<u>A211129</u>	<u>001</u>	Oct 12, 2018
AB		LUPIN LTD	<u>0.2%</u>	<u>A210307</u>	<u>001</u>	Aug 15, 2019
AB		PADAGIS ISRAEL	<u>0.2%</u>	<u>A075666</u>	<u>001</u>	May 24, 2000
AB	!	TARO	<u>0.2%</u>	<u>A075042</u>	<u>001</u>	Aug 25, 1998

OINTMENT; TOPICAL

HYDROCORTISONE VALERATE

AB		COSETTE	<u>0.2%</u>	<u>A211764</u>	<u>001</u>	Mar 04, 2020
AB		GLENMARK PHARMS LTD	<u>0.2%</u>	<u>A211750</u>	<u>001</u>	Dec 14, 2018
AB	!	TARO	<u>0.2%</u>	<u>A075043</u>	<u>001</u>	Aug 25, 1998

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

AT	!	BAUSCH AND LOMB	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A064053</u>	<u>001</u>	Dec 29, 1995
AT		SANDOZ INC	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062423</u>	<u>001</u>	Aug 25, 1983

SUSPENSION/DROPS; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

!		SANDOZ INC	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	A062874	001	May 11, 1988
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SUSPENSION/DROPS; OTIC

CASPORYN HC

AT	+	!	CASPER PHARMA LLC	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N060613</u>	<u>001</u>
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NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

AT		AMRING PHARMS	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A065219</u>	<u>001</u>	May 01, 2006
AT		SANDOZ INC	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062488</u>	<u>001</u>	Nov 06, 1985

OTICAIR

AT		BAUSCH AND LOMB	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A064065</u>	<u>001</u>	Aug 28, 1996
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HYDROMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION

DILAUDID

AP	+	!	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>N019034</u>	<u>003</u>	Apr 30, 2009
AP	+	!		<u>2MG/ML</u>	<u>N019034</u>	<u>004</u>	Apr 30, 2009

HYDROMORPHONE HYDROCHLORIDE

AP		AKORN	<u>10MG/ML</u>	<u>A078228</u>	<u>001</u>	Apr 14, 2010
AP			<u>10MG/ML</u>	<u>A078261</u>	<u>001</u>	Apr 14, 2010
AP		EUROHLTH INTL SARL	<u>2MG/ML</u>	<u>A202159</u>	<u>001</u>	Apr 27, 2018
AP		HOSPIRA INC	<u>1MG/ML</u>	<u>N200403</u>	<u>001</u>	Dec 01, 2011
AP			<u>2MG/ML</u>	<u>N200403</u>	<u>002</u>	Dec 01, 2011
AP			<u>4MG/ML</u>	<u>N200403</u>	<u>003</u>	Dec 01, 2011
AP			<u>10MG/ML</u>	<u>A078591</u>	<u>001</u>	Jun 17, 2008

DILAUDID

+	!	FRESENIUS KABI USA	0.2MG/ML	N019034	006	Jan 16, 2020
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SOLUTION; ORAL

DILAUDID

AA	+	!	RHODES PHARMS	<u>5MG/5ML</u>	<u>N019891</u>	<u>001</u>	Dec 07, 1992
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HYDROMORPHONE HYDROCHLORIDE

AA		ASCENT PHARMS INC	<u>5MG/5ML</u>	<u>A210176</u>	<u>001</u>	Oct 27, 2017
AA		HIKMA	<u>5MG/5ML</u>	<u>A074653</u>	<u>001</u>	Jul 29, 1998

TABLET; ORAL

DILAUDID

AB	+		RHODES PHARMS	<u>2MG</u>	<u>N019892</u>	<u>003</u>	Nov 09, 2007
AB	+			<u>4MG</u>	<u>N019892</u>	<u>002</u>	Nov 09, 2007
AB	+	!		<u>8MG</u>	<u>N019892</u>	<u>001</u>	Dec 07, 1992

HYDROMORPHONE HYDROCHLORIDE

AB		ASCENT PHARMS INC	<u>2MG</u>	<u>A210506</u>	<u>001</u>	Jan 17, 2018
AB			<u>4MG</u>	<u>A210506</u>	<u>002</u>	Jan 17, 2018
AB			<u>8MG</u>	<u>A210506</u>	<u>003</u>	Jan 17, 2018
AB		AUROLIFE PHARMA LLC	<u>2MG</u>	<u>A205814</u>	<u>001</u>	May 13, 2016
AB			<u>4MG</u>	<u>A205814</u>	<u>002</u>	May 13, 2016
AB			<u>8MG</u>	<u>A205814</u>	<u>003</u>	May 13, 2016

PRESCRIPTION DRUG PRODUCT LIST

HYDROMORPHONE HYDROCHLORIDE

TABLET; ORAL

HYDROMORPHONE HYDROCHLORIDE

<u>AB</u>	HIKMA	<u>4MG</u>	<u>A074597</u>	<u>003</u>	May 29, 2009
<u>AB</u>		<u>8MG</u>	<u>A074597</u>	<u>001</u>	Jul 29, 1998
<u>AB</u>	SPECGX LLC	<u>2MG</u>	<u>A076855</u>	<u>002</u>	Sep 19, 2007
<u>AB</u>		<u>4MG</u>	<u>A076855</u>	<u>003</u>	Sep 19, 2007
<u>AB</u>		<u>8MG</u>	<u>A076855</u>	<u>001</u>	Dec 23, 2004

TABLET, EXTENDED RELEASE; ORAL

HYDROMORPHONE HYDROCHLORIDE

<u>AB</u>	ASCENT PHARMS INC	<u>8MG</u>	<u>A212133</u>	<u>001</u>	Sep 23, 2020
<u>AB</u>		<u>12MG</u>	<u>A212133</u>	<u>002</u>	Sep 23, 2020
<u>AB</u>		<u>16MG</u>	<u>A212133</u>	<u>003</u>	Sep 23, 2020
<u>AB</u>		<u>32MG</u>	<u>A212133</u>	<u>004</u>	Sep 23, 2020
<u>AB</u>	OSMOTICA PHARM US	<u>8MG</u>	<u>A205629</u>	<u>001</u>	Jul 07, 2016
<u>AB</u>		<u>12MG</u>	<u>A205629</u>	<u>002</u>	Jul 07, 2016
<u>AB</u>		<u>16MG</u>	<u>A205629</u>	<u>003</u>	Jul 07, 2016
<u>AB</u>	!	<u>32MG</u>	<u>A205629</u>	<u>004</u>	Jul 07, 2016
<u>AB</u>	PADAGIS US	<u>8MG</u>	<u>A204278</u>	<u>001</u>	Apr 06, 2015
<u>AB</u>		<u>12MG</u>	<u>A204278</u>	<u>002</u>	Apr 06, 2015
<u>AB</u>		<u>16MG</u>	<u>A204278</u>	<u>003</u>	Apr 06, 2015
<u>AB</u>		<u>32MG</u>	<u>A204278</u>	<u>004</u>	Sep 20, 2017

HYDROXOCOBALAMIN

INJECTABLE; INJECTION

CYANOKIT

+! SERB SA 5GM/VIAL (5GM/KIT) N022041 001 Apr 08, 2011

HYDROXOCOBALAMIN

! ACTAVIS LLC 1MG/ML A085998 001

HYDROXYAMPHETAMINE HYDROBROMIDE; TROPICAMIDE

SOLUTION/DROPS; OPHTHALMIC

PAREMYD

+! AKORN 1%;0.25% N019261 001 Jan 30, 1992

HYDROXYCHLOROQUINE SULFATE

TABLET; ORAL

HYDROXYCHLOROQUINE SULFATE

<u>AB</u>	ACCORD HLTHCARE	<u>200MG</u>	<u>A213342</u>	<u>001</u>	Apr 07, 2020
<u>AB</u>	ALKALOIDA ZRT	<u>200MG</u>	<u>A201691</u>	<u>001</u>	May 08, 2018
<u>AB</u>	AMNEAL PHARMS CO	<u>200MG</u>	<u>A210577</u>	<u>001</u>	May 15, 2018
<u>AB</u>	APPCO	<u>200MG</u>	<u>A210441</u>	<u>001</u>	May 01, 2018
<u>AB</u>	IPCA LABS LTD	<u>200MG</u>	<u>A040766</u>	<u>001</u>	Jun 14, 2007
<u>AB</u>	LAURUS	<u>200MG</u>	<u>A210959</u>	<u>001</u>	Jan 15, 2019
<u>AB</u>	LUPIN LTD	<u>200MG</u>	<u>A210543</u>	<u>001</u>	Jul 06, 2018
<u>AB</u>	MYLAN	<u>200MG</u>	<u>A040274</u>	<u>001</u>	May 29, 1998
<u>AB</u>	SANDOZ	<u>200MG</u>	<u>A040104</u>	<u>001</u>	Nov 30, 1995
<u>AB</u>	SENORES PHARMS	<u>200MG</u>	<u>A212902</u>	<u>001</u>	May 14, 2020
<u>AB</u>	TEVA PHARMS	<u>200MG</u>	<u>A040081</u>	<u>001</u>	Sep 30, 1994
<u>AB</u>	WATSON LABS	<u>200MG</u>	<u>A040133</u>	<u>001</u>	Nov 30, 1995
<u>AB</u>	ZYDUS PHARMS USA INC	<u>200MG</u>	<u>A040657</u>	<u>001</u>	Sep 21, 2007

PLAQUENIL

<u>AB</u>	+! CONCORDIA	<u>200MG</u>	<u>N009768</u>	<u>001</u>	
	HYDROXYCHLOROQUINE SULFATE				
	ACCORD HLTHCARE	100MG	A213342	002	Aug 18, 2021
		300MG	A213342	003	Aug 18, 2021
		400MG	A213342	004	Aug 18, 2021

HYDROXYPROGESTERONE CAPROATE

SOLUTION; INTRAMUSCULAR

HYDROXYPROGESTERONE CAPROATE

<u>AP</u>	AM REGENT	<u>250MG/ML (250MG/ML)</u>	<u>A210723</u>	<u>001</u>	Jun 21, 2018
<u>AP</u>	ASPEN	<u>250MG/ML (250MG/ML)</u>	<u>A211777</u>	<u>001</u>	Aug 08, 2019
<u>AP</u>	EUGIA PHARMA	<u>250MG/ML (250MG/ML)</u>	<u>A211071</u>	<u>001</u>	Apr 16, 2019
<u>AP</u>	SLAYBACK PHARMA LLC	<u>250MG/ML (250MG/ML)</u>	<u>A210877</u>	<u>001</u>	Mar 22, 2019

MAKENA PRESERVATIVE FREE

<u>AP</u>	+! COVIS	<u>250MG/ML (250MG/ML)</u>	<u>N021945</u>	<u>002</u>	Feb 19, 2016
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HYDROXYPROGESTERONE CAPROATE

<u>AP1</u>	EUGIA PHARMA	<u>1250MG/5ML (250MG/ML)</u>	<u>A211070</u>	<u>001</u>	Apr 16, 2019
<u>AP1</u>	SLAYBACK PHARMA LLC	<u>1250MG/5ML (250MG/ML)</u>	<u>A210618</u>	<u>001</u>	Dec 28, 2018
<u>AP1</u>	SUN PHARM	<u>1250MG/5ML (250MG/ML)</u>	<u>A208381</u>	<u>001</u>	Apr 09, 2019

MAKENA

<u>AP1</u>	+! COVIS	<u>1250MG/5ML (250MG/ML)</u>	<u>N021945</u>	<u>001</u>	Feb 03, 2011
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PRESCRIPTION DRUG PRODUCT LIST

HYDROXYPROGESTERONE CAPROATE

SOLUTION; INTRAMUSCULAR

HYDROXYPROGESTERONE CAPROATE

! ASPEN GLOBAL INC 1250MG/5ML (250MG/ML)

A200271 001 Aug 24, 2015

SOLUTION; SUBCUTANEOUS

MAKENA (AUTOINJECTOR)

+! COVIS 275MG/1.1ML (250MG/ML)

N021945 004 Feb 14, 2018

HYDROXYPROPYL CELLULOSE

INSERT; OPHTHALMIC

LACRISERT

+! VALEANT PHARMS INTL 5MG

N018771 001

HYDROXYUREA

CAPSULE; ORAL

HYDREA**AB** +! BRISTOL MYERS
SQUIBB **500MG****N016295 001**HYDROXYUREA**AB** BARR **500MG** **A075143 001** Oct 16, 1998**AB** LEADING PHARMA LLC **500MG** **A213438 001** Apr 08, 2020**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

N208843 001 Dec 21, 2017

+! 1GM

N208843 002 Dec 21, 2017

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE HYDROCHLORIDE

! AM REGENT 25MG/ML

A087408 001

! 50MG/ML

A087408 002

SYRUP; ORAL

HYDROXYZINE HYDROCHLORIDE**AA** ! AKORN **10MG/5ML****A040010 001** Oct 28, 1994**AA** LANNETT CO INC **10MG/5ML****A201674 001** Aug 21, 2013**AA** +! WOCHKHARDT BIO AG **10MG/5ML****A087294 001** Apr 12, 1982

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE**AB** AMNEAL PHARM **10MG** **A040808 001** Sep 24, 2008**AB** **25MG** **A040808 002** Sep 24, 2008**AB** **50MG** **A040808 003** Sep 24, 2008**AB** BIOPHARM **10MG** **A040804 001** Jun 30, 2008**AB** **25MG** **A040804 002** Jun 30, 2008**AB** **50MG** **A040804 003** Jun 30, 2008**AB** EPIC PHARMA LLC **10MG** **A040604 002** Dec 28, 2004**AB** **25MG** **A040604 003** Dec 28, 2004**AB** **50MG** **A040604 001** Dec 28, 2004**AB** HERITAGE PHARMA **10MG** **A204279 001** Aug 20, 2014**AB** **25MG** **A204279 002** Aug 20, 2014**AB** **50MG** **A204279 003** Aug 20, 2014**AB** HETERO LABS LTD III **10MG** **A040805 001** May 29, 2008**AB** **25MG** **A040805 002** May 29, 2008**AB** **50MG** **A040805 003** May 29, 2008**AB** INVAGEN PHARMS **10MG** **A040812 001** Mar 12, 2008**AB** **25MG** **A040812 002** Mar 12, 2008**AB** **50MG** **A040812 003** Mar 12, 2008**AB** KVK TECH **10MG** **A040786 001** Mar 20, 2007**AB** **25MG** **A040786 002** Mar 20, 2007**AB** **50MG** **A040786 003** Mar 20, 2007**AB** NORTHSTAR HLTHCARE **10MG** **A040840 002** Mar 31, 2008**AB** **25MG** **A040840 003** Mar 31, 2008**AB** **50MG** **A040840 001** Mar 31, 2008**AB** NUVO PHARMS INC **10MG** **A207121 002** Mar 29, 2017**AB** **25MG** **A207121 001** Mar 29, 2017**AB** **50MG** **A207121 003** Mar 29, 2017**AB** +! PLIVA **10MG** **A088617 001** Jan 10, 1986**AB** +! **25MG** **A088618 001** Jan 10, 1986**AB** +! **50MG** **A088619 001** Jan 10, 1986

PRESCRIPTION DRUG PRODUCT LIST

HYDROXYZINE HYDROCHLORIDE

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE

<u>AB</u>	PRINSTON INC	<u>10MG</u>	<u>A040580</u>	<u>003</u>	May 27, 2005
<u>AB</u>		<u>25MG</u>	<u>A040580</u>	<u>002</u>	May 27, 2005
<u>AB</u>		<u>50MG</u>	<u>A040580</u>	<u>001</u>	May 27, 2005

HYDROXYZINE PAMOATE

CAPSULE; ORAL

HYDROXYZINE PAMOATE

<u>AB</u>	BARR	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A088496</u>	<u>001</u>	Jun 15, 1984
<u>AB</u>		<u>EQ 50MG HYDROCHLORIDE</u>	<u>A088487</u>	<u>001</u>	Jun 15, 1984
<u>AB</u>	IMPAX LABS INC	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A040156</u>	<u>001</u>	Jul 15, 1996
<u>AB</u>		<u>EQ 50MG HYDROCHLORIDE</u>	<u>A040156</u>	<u>002</u>	Jul 15, 1996
<u>AB</u>	SANDOZ	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A087479</u>	<u>001</u>	
<u>AB</u>	!	<u>EQ 50MG HYDROCHLORIDE</u>	<u>A086183</u>	<u>001</u>	

VISTARIL

<u>AB</u>	+ PFIZER	<u>EQ 25MG HYDROCHLORIDE</u>	<u>N011459</u>	<u>002</u>	
<u>AB</u>	+ PFIZER	<u>EQ 50MG HYDROCHLORIDE</u>	<u>N011459</u>	<u>004</u>	
	HYDROXYZINE PAMOATE				
BX	HERITAGE PHARMA	EQ 25MG HYDROCHLORIDE	A201507	001	Jun 03, 2013
BX		EQ 50MG HYDROCHLORIDE	A201507	002	Jun 03, 2013
	BARR	EQ 100MG HYDROCHLORIDE	A088488	001	Jun 15, 1984

IBANDRONATE SODIUM

INJECTABLE; INTRAVENOUS

IBANDRONATE SODIUM

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 3MG BASE/3ML</u>	<u>A206058</u>	<u>001</u>	Feb 05, 2016
<u>AP</u>	! APOTEX	<u>EQ 3MG BASE/3ML</u>	<u>A204222</u>	<u>001</u>	Oct 16, 2015
<u>AP</u>	EUGIA PHARMA	<u>EQ 3MG BASE/3ML</u>	<u>A205332</u>	<u>001</u>	Aug 19, 2015
<u>AP</u>	MYLAN LABS LTD	<u>EQ 3MG BASE/3ML</u>	<u>A202671</u>	<u>001</u>	Sep 02, 2014
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 3MG BASE/3ML</u>	<u>A202235</u>	<u>001</u>	Sep 02, 2014
<u>AP</u>	SUN PHARM	<u>EQ 3MG BASE/3ML</u>	<u>A090853</u>	<u>001</u>	Feb 14, 2014

TABLET; ORAL

IBANDRONATE SODIUM

<u>AB</u>	APOTEX INC	<u>EQ 150MG BASE</u>	<u>A078948</u>	<u>001</u>	Mar 19, 2012
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 150MG BASE</u>	<u>A204502</u>	<u>001</u>	Mar 11, 2016
<u>AB</u>	! DR REDDYS LABS LTD	<u>EQ 150MG BASE</u>	<u>A078997</u>	<u>001</u>	Apr 30, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 150MG BASE</u>	<u>A206887</u>	<u>001</u>	Oct 31, 2017
<u>AB</u>	ORBION PHARMS	<u>EQ 150MG BASE</u>	<u>A078998</u>	<u>001</u>	Mar 19, 2012
<u>AB</u>	WATSON LABS TEVA	<u>EQ 150MG BASE</u>	<u>A079003</u>	<u>001</u>	Mar 20, 2012

IBREXAFUNGERP CITRATE

TABLET; ORAL

BREXAFEMME

+!	SCYNEXIS	EQ 150MG BASE	N214900	001	Jun 01, 2021
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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
+!		800MG/200ML (4MG/ML)	N022348	003	Jan 25, 2019

SUSPENSION; ORAL

IBUPROFEN

<u>AB</u>	! ACTAVIS MID ATLANTIC	<u>100MG/5ML</u>	<u>A074978</u>	<u>001</u>	Mar 25, 1998
<u>AB</u>	AKORN	<u>100MG/5ML</u>	<u>A205647</u>	<u>001</u>	Nov 03, 2016
<u>AB</u>	PADAGIS US	<u>100MG/5ML</u>	<u>A076925</u>	<u>001</u>	Sep 23, 2004
<u>AB</u>	TARO	<u>100MG/5ML</u>	<u>A209204</u>	<u>001</u>	Jun 23, 2017
	AUROBINDO PHARMA LTD	100MG/5ML	A209178	001	Feb 16, 2018

PRESCRIPTION DRUG PRODUCT LIST

IBUPROFEN

TABLET; ORAL

IBUPROFEN

<u>AB</u>	ALKEM LABS LTD	<u>400MG</u>	<u>A214699 001</u>	Sep 13, 2021
<u>AB</u>		<u>600MG</u>	<u>A214699 002</u>	Sep 13, 2021
<u>AB</u>		<u>800MG</u>	<u>A214699 003</u>	Sep 13, 2021
<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>	AUROBINDO PHARMA LTD	<u>400MG</u>	<u>A213794 001</u>	May 08, 2020
<u>AB</u>		<u>600MG</u>	<u>A213794 002</u>	May 08, 2020
<u>AB</u>		<u>800MG</u>	<u>A213794 003</u>	May 08, 2020
<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	<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>	L PERRIGO CO	<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>	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>	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
<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

IBUPROFEN LYISINE

INJECTABLE; INTRAVENOUS

IBUPROFEN LYISINE

<u>AP</u>	XGEN PHARMS	<u>EQ 20MG BASE/2ML (EQ 10MG BASE/ML)</u>	<u>A202402 001</u>	Mar 30, 2016
<u>AP</u>	+	<u>EQ 20MG BASE/2ML (EQ 10MG BASE/ML)</u>	<u>N021903 001</u>	Apr 13, 2006

IBUTILIDE FUMARATE

INJECTABLE; INJECTION

CORVERT

<u>AP</u>	+	<u>PFIZER</u>	<u>0.1MG/ML</u>	<u>N020491 001</u>	Dec 28, 1995
<u>AP</u>		<u>MYLAN INSTITUTIONAL</u>	<u>0.1MG/ML</u>	<u>A090643 001</u>	Jan 11, 2010

ICATIBANT ACETATE

INJECTABLE; SUBCUTANEOUS

FIRAZYR

<u>AP</u>	+	<u>SHIRE ORPHAN THERAP</u>	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>N022150 001</u>	Aug 25, 2011
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ICATIBANT ACETATE

<u>AP</u>		<u>CIPLA</u>	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A212446 001</u>	Jul 13, 2020
<u>AP</u>		<u>FRESENIUS KABI USA</u>	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A208317 001</u>	Jun 18, 2020
<u>AP</u>		<u>GLENMARK PHARMS LTD</u>	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A213222 001</u>	May 21, 2021
<u>AP</u>		<u>JIANGSU HANSOH PHARM</u>	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A211021 001</u>	Mar 09, 2020
<u>AP</u>		<u>NANG KUANG PHARM CO</u>	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A212081 001</u>	Dec 16, 2020
<u>AP</u>		<u>SLAYBACK PHARMA LLC</u>	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A211501 001</u>	Sep 01, 2020
<u>AP</u>		<u>TEVA PHARMS USA</u>	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A210118 001</u>	Jul 15, 2019

PRESCRIPTION DRUG PRODUCT LIST

ICODEXTRINSOLUTION; INTRAPERITONEAL
EXTRANEAL

+! BAXTER HLTHCARE 7.5GM/100ML N021321 001 Dec 20, 2002

ICOSAPENT ETHYL

CAPSULE; ORAL

ICOSAPENT ETHYL**AB** APOTEX **1GM** **A209437 001** Jun 30, 2021**AB** DR REDDYS **1GM** **A209499 001** Aug 07, 2020**AB** HIKMA **1GM** **A209457 001** May 21, 2020VASCEPA**AB** +! AMARIN PHARMS **1GM** **N202057 001** Jul 26, 2012

+ 500MG N202057 002 Feb 16, 2017

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDAMYCIN PFS**AP** +! PFIZER **1MG/ML** **N050734 001** Feb 17, 1997IDARUBICIN HYDROCHLORIDE**AP** HIKMA **1MG/ML** **A065275 001** Dec 14, 2006**AP** **1MG/ML** **A065288 001** May 15, 2007IDARUBICIN HYDROCHLORIDE PFS**AP** TEVA PHARMS USA **1MG/ML** **A065036 001** May 01, 2002IDELALISIB

TABLET; ORAL

ZYDELIG

+ GILEAD SCIENCES INC 100MG N205858 001 Jul 23, 2014

+! 150MG N205858 002 Jul 23, 2014

IFOSFAMIDE

INJECTABLE; INJECTION

IFEX**AP** + BAXTER HLTHCARE **1GM/VIAL** **N019763 001** Dec 30, 1988**AP** + **3GM/VIAL** **N019763 002** Dec 30, 1988IFOSFAMIDE**AP** ! FRESENIUS KABI USA **1GM/VIAL** **A076078 001** May 28, 2002**AP** ! **3GM/VIAL** **A076078 002** May 28, 2002**AP** HIKMA **1GM/20ML (50MG/ML)** **A076619 001** Jun 29, 2011**AP** **3GM/60ML (50MG/ML)** **A076619 002** Jun 29, 2011**AP** ! TEVA PHARMS USA **1GM/20ML (50MG/ML)** **A076657 001** Apr 04, 2007**AP** ! **3GM/60ML (50MG/ML)** **A076657 002** Apr 04, 2007ILOPERIDONE

TABLET; ORAL

FANAPT

+! VANDA PHARMS INC 1MG N022192 001 May 06, 2009

+ 2MG N022192 002 May 06, 2009

+ 4MG N022192 003 May 06, 2009

+ 6MG N022192 004 May 06, 2009

+ 8MG N022192 005 May 06, 2009

+ 10MG N022192 006 May 06, 2009

+ 12MG N022192 007 May 06, 2009

ILOPROST

SOLUTION; INHALATION

VENTAVIS

+! ACTELION 10MCG/ML (10MCG/ML) N021779 002 Dec 08, 2005

+! 20MCG/ML (20MCG/ML) N021779 003 Aug 07, 2009

IMATINIB MESYLATE

TABLET; ORAL

GLEEVEC**AB** + NOVARTIS **EQ 100MG BASE** **N021588 001** Apr 18, 2003**AB** +! **EQ 400MG BASE** **N021588 002** Apr 18, 2003IMATINIB MESYLATE**AB** APOTEX **EQ 100MG BASE** **A079179 001** Aug 05, 2016**AB** **EQ 400MG BASE** **A079179 002** Aug 05, 2016**AB** BRECKENRIDGE **EQ 100MG BASE** **A205990 001** Feb 08, 2019**AB** **EQ 400MG BASE** **A205990 002** Feb 08, 2019**AB** DR REDDYS **EQ 100MG BASE** **A206547 001** Aug 13, 2018**AB** **EQ 400MG BASE** **A206547 002** Aug 13, 2018**AB** EUGIA PHARMA **EQ 100MG BASE** **A212773 001** Jul 23, 2020**AB** **EQ 400MG BASE** **A212773 002** Jul 23, 2020**AB** HIKMA **EQ 100MG BASE** **A207586 001** Jul 13, 2018

PRESCRIPTION DRUG PRODUCT LIST

IMATINIB MESYLATE

TABLET; ORAL

IMATINIB MESYLATE

<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A207586 002</u>	Jul 13, 2018
<u>AB</u>	MYLAN	<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>	NATCO PHARMA LTD	<u>EQ 100MG BASE</u>	<u>A207818 001</u>	Mar 01, 2019
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A207818 002</u>	Mar 01, 2019
<u>AB</u>	SHILPA	<u>EQ 100MG BASE</u>	<u>A208302 001</u>	Jan 17, 2019
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A208302 002</u>	Jan 17, 2019
<u>AB</u>	SUN PHARM	<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>	WOCKHARDT BIO AG	<u>EQ 100MG BASE</u>	<u>A208429 001</u>	Jan 17, 2019
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A208429 002</u>	Jan 17, 2019
<u>AB</u>	ZYDUS PHARMS	<u>EQ 100MG BASE</u>	<u>A210658 001</u>	Apr 08, 2020
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A210658 002</u>	Apr 08, 2020

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>	OXFORD PHARMS	<u>10MG</u>	<u>A040751 003</u>	Feb 28, 2008
<u>AB</u>		<u>25MG</u>	<u>A040751 002</u>	Feb 28, 2008
<u>AB</u>		<u>50MG</u>	<u>A040751 001</u>	Feb 28, 2008
<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
<u>AB</u>	STRIDES PHARMA	<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>	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 PAMOATE

CAPSULE; ORAL

IMIPRAMINE PAMOATE

<u>AB</u>	! HIKMA	<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
<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

IMIQUIMOD

CREAM; TOPICAL

ALDARA

<u>AB</u>	+! BAUSCH	<u>5%</u>	<u>N020723 001</u>	Feb 27, 1997
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IMIQUIMOD

<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>	PADAGIS ISRAEL	<u>5%</u>	<u>A078837 001</u>	Sep 07, 2010
<u>AB</u>	TARO	<u>3.75%</u>	<u>A205971 001</u>	Jan 26, 2021
<u>AB</u>		<u>5%</u>	<u>A200173 001</u>	Apr 15, 2011

ZYCLARA

<u>AB</u>	+! BAUSCH	<u>3.75%</u>	<u>N022483 001</u>	Mar 25, 2010
	+!	<u>2.5%</u>	<u>N022483 002</u>	Jul 15, 2011

PRESCRIPTION DRUG PRODUCT LIST

INAMRINONE LACTATE

INJECTABLE; INJECTION

AMRINONE LACTATE

! HIKMA

EQ 5MG BASE/ML

A075513 001 May 09, 2000

INCLISIRAN SODIUM

SOLUTION; SUBCUTANEOUS

LEQVIO

+! NOVARTIS

EQ 284MG BASE/1.5ML (EQ 189MG BASE/ML)

N214012 001 Dec 22, 2021

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE**AB** ACTAVIS ELIZABETH**1.25MG****A074722 001** Jun 17, 1996**AB****2.5MG****A074722 002** Jun 17, 1996**AB** ANI PHARMS**1.25MG****A074299 002** Apr 29, 1996**AB** !**2.5MG****A074299 001** Jul 27, 1995INDIUM IN-111 CHLORIDE

INJECTABLE; INJECTION

INDIUM IN 111 CHLORIDE

+! CURIUM

5mCi/0.5ML

N019841 001 Sep 27, 1994

INDIUM IN-111 OXYQUINOLINE

INJECTABLE; INJECTION

INDIUM IN 111 OXYQUINOLINE**AP** BWXT ITG**1mCi/ML****A202586 001** Jul 25, 2018**AP** +!

GE HEALTHCARE

1mCi/ML**N019044 001** Dec 24, 1985INDIUM 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

+! CURIUM

3mCi/ML

N020314 001 Jun 02, 1994

INDOCYANINE GREEN

INJECTABLE; INJECTION

IC-GREEN**AP** +! AKORN**25MG/VIAL****N011525 001**INDOCYANINE GREEN**AP** RENEW PHARMS**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** CADILA**25MG****A090403 001** Nov 15, 2010**AB****50MG****A090403 002** Nov 15, 2010**AB** GLENMARK GENERICS**25MG****A091276 001** Dec 22, 2010**AB** !**50MG****A091276 002** Dec 22, 2010**AB** HETERO LABS LTD III**25MG****A091240 001** Apr 12, 2011**AB****50MG****A091240 002** Apr 12, 2011**AB** SANDOZ**25MG****A070673 001** Apr 29, 1987**AB****50MG****A070674 001** Apr 29, 1987

CAPSULE, EXTENDED RELEASE; ORAL

INDOMETHACIN**AB** AMNEAL PHARMS**75MG****A091549 001** Dec 01, 2010**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**

NOVAST LABS

75MG**A204853 001** May 08, 2017**AB** !

SANDOZ

75MG**A074464 001** May 28, 1998**AB**

ZYDUS PHARMS

75MG**A202711 001** Sep 25, 2017

INJECTABLE; INJECTION

INDOMETHACIN

+! FRESENIUS KABI USA

EQ 1MG BASE/VIAL

N022536 001 Mar 17, 2010

SUPPOSITORY; RECTAL

INDOMETHACIN

! COSETTE

50MG

A073314 001 Aug 31, 1992

PRESCRIPTION DRUG PRODUCT LIST

INDOMETHACIN

SUSPENSION; ORAL

INDOCIN

+! EGALET

25MG/5ML

N018332 001 Oct 10, 1985

INDOMETHACIN SODIUM

INJECTABLE; INJECTION

INDOMETHACIN SODIUMAP HOSPIRA INCEQ 1MG BASE/VIALA204118 001 Apr 19, 2016AP ! NAVINTA LLCEQ 1MG BASE/VIALA206561 001 Jul 19, 2017AP WEST-WARD PHARMS
INTEQ 1MG BASE/VIALA078713 001 Jul 16, 2008INFIGRATINIB PHOSPHATE

CAPSULE; ORAL

TRUSELTIQ

+ HELSINN HLTHCARE

25MG

N214622 001 May 28, 2021

+!

100MG

N214622 002 May 28, 2021

INOTERSEN SODIUM

SOLUTION; SUBCUTANEOUS

TEGSEDI

+! AKCEA THERAPS

EQ 284MG BASE/1.5ML (EQ 189.3MG
BASE/ML)

N211172 001 Oct 05, 2018

IOBENGUANE I-131

SOLUTION; INTRAVENOUS

AZEDRA

+! PROGENICS PHARMS
INC

15mCi/ML

N209607 001 Jul 30, 2018

IOBENGUANE 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

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

PRESCRIPTION DRUG PRODUCT LIST

IOPAMIDOL

INJECTABLE; INJECTION

ISOVUE-200

+! BRACCO 41% N018735 006 Jul 07, 1987

ISOVUE-250

+! BRACCO 51% N018735 007 Jul 06, 1992

+! 51% N020327 002 Oct 12, 1994

ISOVUE-300

+! BRACCO 61% N018735 002 Dec 31, 1985

+! 61% N020327 003 Oct 12, 1994

ISOVUE-370

+! BRACCO 76% N018735 003 Dec 31, 1985

+! 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 62.3% N021425 001 Sep 20, 2002

+! 76.9% N021425 002 Sep 20, 2002

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

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 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** LUOXIN AUROVITAS **0.02%** **A206543 001** Oct 27, 2016**AN** NEPHRON **0.02%** **A075562 001** Sep 27, 2001**AN** ! RITEDOSE CORP **0.02%** **A075693 001** Jan 26, 2001**AN** SUN PHARM **0.02%** **A207903 001** Jan 03, 2017

SPRAY, METERED; NASAL

IPRATROPIUM BROMIDE**AB** APOTEX INC **0.021MG/SPRAY** **A076156 001** Apr 18, 2003**AB** **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** ! HIKMA **0.021MG/SPRAY** **A076664 001** Nov 05, 2003**AB** ! **0.042MG/SPRAY** **A076598 001** Nov 05, 2003

PRESCRIPTION DRUG PRODUCT LIST

IRBESARTAN

TABLET; ORAL

AVAPRO

<u>AB</u>	+	SANOFI AVENTIS US	<u>75MG</u>	<u>N020757</u>	<u>001</u>	Sep 30, 1997
<u>AB</u>	+		<u>150MG</u>	<u>N020757</u>	<u>002</u>	Sep 30, 1997
<u>AB</u>	+	!	<u>300MG</u>	<u>N020757</u>	<u>003</u>	Sep 30, 1997

IRBESARTAN

<u>AB</u>		ALEMBIC PHARMS LTD	<u>75MG</u>	<u>A091236</u>	<u>001</u>	Oct 15, 2012
<u>AB</u>			<u>150MG</u>	<u>A091236</u>	<u>002</u>	Oct 15, 2012
<u>AB</u>			<u>300MG</u>	<u>A091236</u>	<u>003</u>	Oct 15, 2012
<u>AB</u>		AMNEAL PHARMS	<u>75MG</u>	<u>A204740</u>	<u>001</u>	Apr 17, 2018
<u>AB</u>			<u>150MG</u>	<u>A204740</u>	<u>002</u>	Apr 17, 2018
<u>AB</u>			<u>300MG</u>	<u>A204740</u>	<u>003</u>	Apr 17, 2018
<u>AB</u>		AUROBINDO PHARMA LTD	<u>75MG</u>	<u>A203081</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>			<u>150MG</u>	<u>A203081</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>			<u>300MG</u>	<u>A203081</u>	<u>003</u>	Sep 27, 2012
<u>AB</u>		CHARTWELL MOLECULAR	<u>75MG</u>	<u>A077205</u>	<u>001</u>	Nov 14, 2012
<u>AB</u>			<u>150MG</u>	<u>A077205</u>	<u>002</u>	Nov 14, 2012
<u>AB</u>			<u>300MG</u>	<u>A077205</u>	<u>003</u>	Nov 14, 2012
<u>AB</u>		HETERO LABS LTD V	<u>75MG</u>	<u>A202910</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>			<u>150MG</u>	<u>A202910</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>			<u>300MG</u>	<u>A202910</u>	<u>003</u>	Sep 27, 2012
<u>AB</u>		HISUN PHARM HANGZHOU	<u>75MG</u>	<u>A206194</u>	<u>001</u>	Jun 14, 2016
<u>AB</u>			<u>150MG</u>	<u>A206194</u>	<u>002</u>	Jun 14, 2016
<u>AB</u>			<u>300MG</u>	<u>A206194</u>	<u>003</u>	Jun 14, 2016
<u>AB</u>		JUBILANT GENERICS	<u>75MG</u>	<u>A203534</u>	<u>001</u>	Feb 23, 2015
<u>AB</u>			<u>150MG</u>	<u>A203534</u>	<u>002</u>	Feb 23, 2015
<u>AB</u>			<u>300MG</u>	<u>A203534</u>	<u>003</u>	Feb 23, 2015
<u>AB</u>		LUPIN LTD	<u>75MG</u>	<u>A201531</u>	<u>001</u>	Oct 15, 2012
<u>AB</u>			<u>150MG</u>	<u>A201531</u>	<u>002</u>	Oct 15, 2012
<u>AB</u>			<u>300MG</u>	<u>A201531</u>	<u>003</u>	Oct 15, 2012
<u>AB</u>		MACLEODS PHARMS LTD	<u>75MG</u>	<u>A202254</u>	<u>001</u>	Oct 03, 2012
<u>AB</u>			<u>150MG</u>	<u>A202254</u>	<u>002</u>	Oct 03, 2012
<u>AB</u>			<u>300MG</u>	<u>A202254</u>	<u>003</u>	Oct 03, 2012
<u>AB</u>		PRINSTON INC	<u>75MG</u>	<u>A203071</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>			<u>150MG</u>	<u>A203071</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>			<u>300MG</u>	<u>A203071</u>	<u>003</u>	Sep 27, 2012
<u>AB</u>		SANDOZ	<u>75MG</u>	<u>A077466</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>			<u>150MG</u>	<u>A077466</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>			<u>300MG</u>	<u>A077466</u>	<u>003</u>	Sep 27, 2012
<u>AB</u>		SCIEGEN PHARMS INC	<u>75MG</u>	<u>A204774</u>	<u>001</u>	Dec 07, 2015
<u>AB</u>			<u>150MG</u>	<u>A204774</u>	<u>002</u>	Dec 07, 2015
<u>AB</u>			<u>300MG</u>	<u>A204774</u>	<u>003</u>	Dec 07, 2015
<u>AB</u>		TEVA PHARMS	<u>75MG</u>	<u>A077159</u>	<u>001</u>	Mar 30, 2012
<u>AB</u>			<u>150MG</u>	<u>A077159</u>	<u>002</u>	Mar 30, 2012
<u>AB</u>			<u>300MG</u>	<u>A077159</u>	<u>003</u>	Mar 30, 2012
<u>AB</u>		UNICHEM	<u>75MG</u>	<u>A203020</u>	<u>001</u>	Dec 07, 2015
<u>AB</u>			<u>150MG</u>	<u>A203020</u>	<u>002</u>	Dec 07, 2015
<u>AB</u>			<u>300MG</u>	<u>A203020</u>	<u>003</u>	Dec 07, 2015
<u>AB</u>		ZYDUS PHARMS USA INC	<u>75MG</u>	<u>A079213</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>			<u>150MG</u>	<u>A079213</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>			<u>300MG</u>	<u>A079213</u>	<u>003</u>	Sep 27, 2012

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

CAMPTOSAR

<u>AP</u>	+	PFIZER INC	<u>40MG/2ML (20MG/ML)</u>	<u>N020571</u>	<u>001</u>	Jun 14, 1996
<u>AP</u>	+		<u>100MG/5ML (20MG/ML)</u>	<u>N020571</u>	<u>002</u>	Jun 14, 1996
<u>AP</u>	+	!	<u>300MG/15ML (20MG/ML)</u>	<u>N020571</u>	<u>003</u>	Aug 05, 2010

IRINOTECAN HYDROCHLORIDE

<u>AP</u>		ACCORD HLTHCARE	<u>40MG/2ML (20MG/ML)</u>	<u>A079068</u>	<u>001</u>	Nov 21, 2008
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A079068</u>	<u>002</u>	Nov 21, 2008
<u>AP</u>		ACTAVIS TOTOWA	<u>40MG/2ML (20MG/ML)</u>	<u>A078589</u>	<u>001</u>	Feb 27, 2008
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A078589</u>	<u>002</u>	Feb 27, 2008
<u>AP</u>			<u>500MG/25ML (20MG/ML)</u>	<u>A078589</u>	<u>003</u>	Nov 18, 2015
<u>AP</u>		AKORN	<u>40MG/2ML (20MG/ML)</u>	<u>A090726</u>	<u>001</u>	Sep 16, 2009
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A090726</u>	<u>002</u>	Sep 16, 2009
<u>AP</u>		EUGIA PHARMA	<u>40MG/2ML (20MG/ML)</u>	<u>A213278</u>	<u>001</u>	Nov 02, 2020
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A213278</u>	<u>002</u>	Nov 02, 2020
<u>AP</u>			<u>300MG/15ML (20MG/ML)</u>	<u>A213278</u>	<u>003</u>	Nov 02, 2020

PRESCRIPTION DRUG PRODUCT LIST

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

<u>AP</u>	FRESENIUS KABI USA	<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>	GLAND PHARMA LTD	<u>40MG/2ML (20MG/ML)</u>	<u>A212993 001</u>	Nov 18, 2019
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A212993 002</u>	Nov 18, 2019
<u>AP</u>	HIKMA FARMACEUTICA	<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>	HISUN PHARM HANGZHOU	<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>	HOSPIRA	<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>500MG/25ML (20MG/ML)</u>	<u>A078796 001</u>	Feb 27, 2008
<u>AP</u>	INTAS PHARMS USA	<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>	JIANGSU PHARMS	<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>	NOVAST LABS	<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>	QILU PHARM HAINAN	<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>	SHILPA	<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>300MG/15ML (20MG/ML)</u>	<u>A208718 003</u>	Aug 16, 2019
<u>AP</u>	TEVA PHARMS USA	<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>	WEST-WARD PHARMS INT	<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
<u>AP</u>	ZENNOVA	<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

INJECTABLE, LIPOSOMAL; INTRAVENOUS

ONIVYDE

+! IPSEN INC EQ 43MG BASE/10ML (EQ 4.3MG BASE/ML) N207793 001 Oct 22, 2015

ISAVUCONAZONIUM SULFATE

CAPSULE; ORAL

CRESEMBA

+! ASTELLAS 186MG N207500 001 Mar 06, 2015

POWDER; INTRAVENOUS

CRESEMBA

+! ASTELLAS 372MG N207501 001 Mar 06, 2015

ISOCARBOXAZID

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 PHARMA 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 ANDA REPOSITORY 100MG A040090 001 Jun 26, 1997AA 300MG A040090 002 Jun 26, 1997AA ! BARR 100MG A080936 001AA ! 300MG A080937 002AA THEPHARMANETWORK 100MG A202610 001 Oct 29, 2014

PRESCRIPTION DRUG PRODUCT LIST

ISONIAZID

TABLET; ORAL

ISONIAZID

LLC

AA		300MG	A202610 002	Oct 29, 2014
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ISOPROTERENOL HYDROCHLORIDE

INJECTABLE; INJECTION

ISOPROTERENOL HYDROCHLORIDE

AP	AMNEAL	0.2MG/ML	A210576 001	Oct 17, 2018
AP	AMPHASTAR PHARMS INC	0.2MG/ML	A210106 001	Jun 18, 2018
AP	AMRING PHARMS	0.2MG/ML	A211237 001	May 19, 2021
AP	AVET LIFESCIENCES	0.2MG/ML	A212189 001	Nov 19, 2021
AP	CIPLA	0.2MG/ML	A210322 001	Jun 12, 2018
AP	EUGIA PHARMA	0.2MG/ML	A211864 001	Feb 09, 2021
AP	MICRO LABS	0.2MG/ML	A210845 001	Feb 16, 2021
AP	NEXUS PHARMS	0.2MG/ML	A206961 001	Aug 02, 2017

ISUPREL

AP	+!	BAUSCH	0.2MG/ML	N010515 001
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ISOSORBIDE DINITRATE

TABLET; ORAL

ISORDIL

AB	+	BAUSCH	5MG	N012093 007	Jul 29, 1988
AB	+	!	40MG	N012093 001	Jul 29, 1988

ISOSORBIDE DINITRATE

AB		HIKMA INTL PHARMS	5MG	A086067 001	Oct 29, 1987
AB			10MG	A086066 001	Oct 29, 1987
AB			20MG	A088088 001	Nov 02, 1987
AB		PAR PHARM	5MG	A086923 001	Mar 12, 1987
AB			10MG	A086925 001	Mar 12, 1987
AB			20MG	A087537 001	Oct 02, 1987
AB	!		30MG	A087946 001	Jan 12, 1988
AB		PAR PHARM INC	40MG	A211290 001	Nov 23, 2021
AB		SANDOZ	5MG	A086221 001	Jan 07, 1988
AB			10MG	A086223 001	Jan 07, 1988
AB			20MG	A089367 001	Apr 07, 1988
AB		ZYDUS	5MG	A213057 001	Nov 20, 2019
AB			10MG	A213057 002	Nov 20, 2019
AB			20MG	A213057 003	Nov 20, 2019
AB			30MG	A213057 004	Nov 20, 2019
AB			40MG	A213057 005	Nov 20, 2019

TABLET, EXTENDED RELEASE; ORAL

ISOSORBIDE DINITRATE

!	SUN PHARM INDS INC	40MG	A040009 001	Dec 30, 1998
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ISOSORBIDE MONONITRATE

TABLET; ORAL

ISOSORBIDE MONONITRATE

AB		ACTAVIS ELIZABETH	10MG	A075037 002	Oct 30, 1998
AB			20MG	A075037 001	Oct 30, 1998

MONOKET

AB	+	ECI PHARMS LLC	10MG	N020215 002	Jun 30, 1993
AB	+	!	20MG	N020215 001	Jun 30, 1993

TABLET, EXTENDED RELEASE; ORAL

ISOSORBIDE MONONITRATE

AB		DEXCEL LTD	30MG	A075522 002	Sep 20, 2016
AB			60MG	A075522 001	Apr 17, 2000
AB			120MG	A210822 001	Aug 29, 2018
AB		LANNETT CO INC	30MG	A075155 002	Jan 13, 2000
AB			60MG	A075155 001	Oct 30, 1998
AB	!		120MG	A075155 003	Aug 04, 2000
AB		NESHER PHARMS	30MG	A075395 001	Mar 16, 2000
AB			60MG	A075395 002	Mar 16, 2000
AB			120MG	A075395 003	Mar 16, 2000
AB		RICONPHARMA LLC	30MG	A210918 001	Nov 05, 2018
AB			60MG	A210918 002	Nov 05, 2018
AB			120MG	A210918 003	Nov 05, 2018
AB		TORRENT PHARMS	30MG	A200270 001	Jun 03, 2011
AB			60MG	A200495 001	Jun 03, 2011
AB			120MG	A200495 002	Jun 03, 2011

PRESCRIPTION DRUG PRODUCT LIST

ISOSULFAN BLUE

INJECTABLE; INJECTION

ISOSULFAN BLUE

<u>AP</u>	EUGIA PHARMA	<u>1%</u>	<u>A206831</u>	<u>001</u>	Feb 02, 2016
<u>AP</u>	MEITHEAL	<u>1%</u>	<u>A213130</u>	<u>001</u>	Nov 03, 2021
<u>AP</u>	! MYLAN INSTITUTIONAL	<u>1%</u>	<u>A090874</u>	<u>001</u>	Jul 20, 2010

ISOTRETINOIN

CAPSULE; ORAL

AMNESTEEM

<u>AB1</u>	MYLAN PHARMS INC	<u>10MG</u>	<u>A075945</u>	<u>001</u>	Nov 08, 2002
<u>AB1</u>		<u>20MG</u>	<u>A075945</u>	<u>002</u>	Nov 08, 2002
<u>AB1</u>		<u>40MG</u>	<u>A075945</u>	<u>003</u>	Nov 08, 2002

CLARAVIS

<u>AB1</u>	TEVA PHARMS USA	<u>10MG</u>	<u>A076356</u>	<u>001</u>	Apr 11, 2003
<u>AB1</u>		<u>20MG</u>	<u>A076135</u>	<u>002</u>	Apr 11, 2003
<u>AB1</u>		<u>30MG</u>	<u>A076135</u>	<u>003</u>	May 11, 2006
<u>AB1</u>	!	<u>40MG</u>	<u>A076135</u>	<u>001</u>	Apr 11, 2003

ISOTRETINOIN

<u>AB1</u>	AMNEAL PHARMS NY	<u>10MG</u>	<u>A207792</u>	<u>001</u>	Sep 29, 2017
<u>AB1</u>		<u>20MG</u>	<u>A207792</u>	<u>002</u>	Sep 29, 2017
<u>AB1</u>		<u>30MG</u>	<u>A207792</u>	<u>003</u>	Sep 29, 2017
<u>AB1</u>		<u>40MG</u>	<u>A207792</u>	<u>004</u>	Sep 29, 2017

MYORISAN

<u>AB1</u>	DOUGLAS PHARMS	<u>10MG</u>	<u>A076485</u>	<u>001</u>	Jan 19, 2012
<u>AB1</u>		<u>20MG</u>	<u>A076485</u>	<u>002</u>	Jan 19, 2012
<u>AB1</u>		<u>30MG</u>	<u>A076485</u>	<u>004</u>	Aug 25, 2015
<u>AB1</u>		<u>40MG</u>	<u>A076485</u>	<u>003</u>	Jan 19, 2012

ZENATANE

<u>AB1</u>	DR REDDYS LABS LTD	<u>10MG</u>	<u>A202099</u>	<u>001</u>	Mar 25, 2013
<u>AB1</u>		<u>20MG</u>	<u>A202099</u>	<u>002</u>	Mar 25, 2013
<u>AB1</u>		<u>30MG</u>	<u>A202099</u>	<u>004</u>	Feb 23, 2015
<u>AB1</u>		<u>40MG</u>	<u>A202099</u>	<u>003</u>	Mar 25, 2013

ABSORICA

<u>AB2</u>	+ SUN PHARM INDS INC	<u>10MG</u>	<u>N021951</u>	<u>001</u>	May 25, 2012
<u>AB2</u>	+	<u>20MG</u>	<u>N021951</u>	<u>002</u>	May 25, 2012
<u>AB2</u>	+	<u>25MG</u>	<u>N021951</u>	<u>005</u>	Aug 15, 2014
<u>AB2</u>	+	<u>30MG</u>	<u>N021951</u>	<u>003</u>	May 25, 2012
<u>AB2</u>	+	<u>35MG</u>	<u>N021951</u>	<u>006</u>	Aug 15, 2014
<u>AB2</u>	+	<u>40MG</u>	<u>N021951</u>	<u>004</u>	May 25, 2012

ISOTRETINOIN

<u>AB2</u>	ACTAVIS LABS FL	<u>10MG</u>	<u>A205063</u>	<u>001</u>	Mar 31, 2021
<u>AB2</u>		<u>20MG</u>	<u>A205063</u>	<u>002</u>	Mar 31, 2021
<u>AB2</u>		<u>25MG</u>	<u>A205063</u>	<u>003</u>	Mar 31, 2021
<u>AB2</u>		<u>30MG</u>	<u>A205063</u>	<u>004</u>	Mar 31, 2021
<u>AB2</u>		<u>35MG</u>	<u>A205063</u>	<u>005</u>	Mar 31, 2021
<u>AB2</u>		<u>40MG</u>	<u>A205063</u>	<u>006</u>	Mar 31, 2021
<u>AB2</u>	UPSHER SMITH LABS	<u>10MG</u>	<u>A212333</u>	<u>001</u>	Sep 21, 2021
<u>AB2</u>		<u>20MG</u>	<u>A212333</u>	<u>002</u>	Sep 21, 2021
<u>AB2</u>		<u>30MG</u>	<u>A212333</u>	<u>003</u>	Sep 21, 2021
<u>AB2</u>		<u>40MG</u>	<u>A213571</u>	<u>001</u>	Apr 12, 2021

ABSORICA LD

+	SUN PHARM	8MG	N211913	001	Nov 05, 2019
+		16MG	N211913	002	Nov 05, 2019
+		24MG	N211913	004	Nov 05, 2019
+	!	32MG	N211913	006	Nov 05, 2019

ISRADIPINE

CAPSULE; ORAL

ISRADIPINE

<u>AB</u>	ELITE LABS INC	<u>2.5MG</u>	<u>A077169</u>	<u>001</u>	Apr 24, 2006
<u>AB</u>		<u>5MG</u>	<u>A077169</u>	<u>002</u>	Apr 24, 2006
<u>AB</u>	WATSON LABS TEVA	<u>2.5MG</u>	<u>A077317</u>	<u>001</u>	Jan 05, 2006
<u>AB</u>	!	<u>5MG</u>	<u>A077317</u>	<u>002</u>	Jan 05, 2006

ISTRADIFYLLINE

TABLET; ORAL

NOURIANZ

+	KYOWA KIRIN	20MG	N022075	001	Aug 27, 2019
+	!	40MG	N022075	002	Aug 27, 2019

PRESCRIPTION DRUG PRODUCT LIST

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	PAR PHARM INC	100MG	A205724 001	Dec 13, 2016
AB	SANDOZ	100MG	A076104 001	May 28, 2004
AB	TORRENT	100MG	A209460 001	Aug 24, 2018
AB	ZYDUS PHARMS	100MG	A204672 001	Sep 19, 2017

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
AA	ANNORA PHARMA	10MG/ML	A212239 001	Sep 01, 2020

SPORANOX

AA	+ ! JANSSEN PHARMS	10MG/ML	N020657 001	Feb 21, 1997
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IVABRADINE

SOLUTION; ORAL

CORLANOR

	+ ! AMGEN INC	5MG/5ML (1MG/ML)	N209964 001	Apr 22, 2019
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IVABRADINE HYDROCHLORIDE

TABLET; ORAL

CORLANOR

AB	+ AMGEN INC	EQ 5MG BASE	N206143 001	Apr 15, 2015
AB	+ !	EQ 7.5MG BASE	N206143 002	Apr 15, 2015

IVABRADINE HYDROCHLORIDE

AB	CENTAUR PHARMS PVT	EQ 5MG BASE	A214051 001	Dec 30, 2021
AB		EQ 7.5MG BASE	A214051 002	Dec 30, 2021

IVACAFTOR

GRANULE; ORAL

KALYDECO

	+ VERTEX PHARMS INC	25MG/PACKET	N207925 003	Apr 29, 2019
	+	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
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IVACAFTOR; IVACAFTOR, TEZACAFTOR

TABLET; ORAL

SYMDEKO (COPACKAGED)

	+ VERTEX PHARMS INC	75MG;75MG, 50MG	N210491 002	Jun 21, 2019
	+ !	150MG;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

IVERMECTIN

AB	TEVA PHARMS USA	1%	A210019 001	Sep 13, 2019
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SOOLANTRA

AB	+ ! GALDERMA LABS LP	1%	N206255 001	Dec 19, 2014
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TABLET; ORAL

IVERMECTIN

AB	EDENBRIDGE PHARMS	3MG	A204154 001	Oct 24, 2014
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STROMEKTOL

AB	+ ! MERCK SHARP DOHME	3MG	N050742 002	Oct 08, 1998
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PRESCRIPTION DRUG PRODUCT LIST

IVOSIDENIB

TABLET; ORAL

TIBSOVO

+! SERVIER 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

+ TAKEDA PHARMS USA 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

KETALARAP +! PAR STERILE EQ 10MG BASE/ML N016812 001
PRODUCTSAP +! EQ 50MG BASE/ML N016812 002AP +! EQ 100MG BASE/ML N016812 003KETAMINE HYDROCHLORIDEAP EUGIA PHARMA EQ 10MG BASE/ML A076092 001 Sep 30, 2008AP EQ 50MG BASE/ML A076092 002 Dec 28, 2001AP EQ 100MG BASE/ML A076092 003 Oct 25, 2002AP HIKMA EQ 50MG BASE/ML A074524 001 Mar 22, 1996AP EQ 100MG BASE/ML A074524 002 Mar 22, 1996AP HOSPIRA EQ 50MG BASE/ML A074549 001 Jun 27, 1996AP EQ 100MG BASE/ML A074549 002 Jun 27, 1996KETOCONAZOLE

AEROSOL, FOAM; TOPICAL

EXTINAAB +! MYLAN 2% N021738 001 Jun 12, 2007KETOCONAZOLEAB PADAGIS ISRAEL 2% A091550 001 Aug 25, 2011AB XIROMED 2% A213601 001 May 21, 2021

CREAM; TOPICAL

KETOCONAZOLEAB ENCUBE 2% A212443 001 May 20, 2021AB FOUGERA PHARMS 2% A076294 001 Apr 28, 2004AB TASMAN PHARMA 2% A215185 001 Nov 17, 2021AB ! TEVA 2% A075581 001 Apr 25, 2000KETOZOLEAB TARO 2% A075638 001 Dec 18, 2002

GEL; TOPICAL

XOLEGEL

+! ALMIRALL 2% N021946 001 Jul 28, 2006

SHAMPOO; TOPICAL

KETOCONAZOLEAB ! PADAGIS ISRAEL 2% A076419 001 Jan 07, 2004AB TOLMAR 2% A076942 001 Apr 11, 2005

TABLET; ORAL

KETOCONAZOLEAB SENORES PHARMS 200MG A075912 001 Jan 10, 2002AB STRIDES PHARMA 200MG A210457 001 Jun 18, 2018AB ! TARO 200MG A075319 001 Jun 15, 1999KETOPROFEN

CAPSULE; ORAL

KETOPROFENAB MISEMER 50MG A074014 002 Jan 29, 1993AB TEVA 50MG A073516 001 Dec 22, 1992

MISEMER 25MG A074014 001 Jan 29, 1993

! 75MG A074014 003 Jan 29, 1993

CAPSULE, EXTENDED RELEASE; ORAL

KETOPROFEN

! MYLAN 200MG A075679 001 Feb 20, 2002

PRESCRIPTION DRUG PRODUCT LIST

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

AP	BAXTER HLTHCARE CORP	15MG/ML	A209900 002	Jul 25, 2018
AP		30MG/ML	A209900 001	Sep 15, 2017
AP	FRESENIUS KABI USA	15MG/ML	A075784 001	Jan 11, 2002
AP		15MG/ML	A203242 001	Oct 07, 2015
AP		30MG/ML	A075784 002	Jan 11, 2002
AP		30MG/ML	A203242 002	Oct 07, 2015
AP	GLAND	15MG/ML	A204216 001	Nov 01, 2016
AP		30MG/ML	A204216 002	Nov 01, 2016
AP	HIKMA	15MG/ML	A075772 001	Jul 21, 2004
AP		30MG/ML	A075772 002	Jul 21, 2004
AP	! HOSPIRA	15MG/ML	A074802 001	Jun 05, 1997
AP	!	30MG/ML	A074802 002	Jun 05, 1997
AP		30MG/ML	A074993 002	Jan 27, 1999
AP	NEPHRON	30MG/ML	A211445 001	Aug 20, 2020
AP	SAGENT PHARMS INC	15MG/ML	A091065 001	Nov 27, 2013
AP		30MG/ML	A091065 002	Nov 27, 2013
AP	SANDOZ INC	30MG/ML	A076271 002	Oct 06, 2004
AP	WOCKHARDT	15MG/ML	A077942 001	Mar 27, 2007
AP		30MG/ML	A077942 002	Mar 27, 2007

SOLUTION/DROPS; OPHTHALMIC

ACULAR

AT	+! ALLERGAN	0.5%	N019700 001	Nov 09, 1992
	<u>ACULAR LS</u>			
AT	+! ALLERGAN	0.4%	N021528 001	May 30, 2003

KETOROLAC TROMETHAMINE

AT	AKORN	0.4%	A078399 001	Nov 05, 2009
AT		0.5%	A078434 001	Nov 05, 2009
AT	APOTEX INC	0.4%	A077308 001	Nov 05, 2009
AT		0.5%	A076109 001	Nov 05, 2009
AT	MICRO LABS LTD INDIA	0.5%	A203410 001	Apr 05, 2019
AT	SANDOZ INC	0.5%	A076583 001	Nov 05, 2009
AT	SUN PHARM	0.5%	A090017 001	Nov 05, 2009
	ACUVAIL			
	+! ALLERGAN	0.45%	N022427 001	Jul 22, 2009
	SPRAY, METERED; NASAL			
	SPRIX			
	+! ZYLA	15.75MG/SPRAY	N022382 001	May 14, 2010

TABLET; ORAL

KETOROLAC TROMETHAMINE

AB	CYCLE PHARMS LTD	10MG	A210616 001	Aug 16, 2018
AB	MYLAN	10MG	A074761 001	May 16, 1997
AB	! TEVA	10MG	A074754 001	May 16, 1997

KETOROLAC TROMETHAMINE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION; IRRIGATION

KETOROLAC TROMETHAMINE AND PHENYLEPHRINE HYDROCHLORIDE

AT	LUPIN LTD	EQ 0.3% BASE:EQ 1% BASE	A210183 001	Jul 01, 2019
	<u>OMIDRIA</u>			
AT	+! 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
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LABETALOL HYDROCHLORIDE

INJECTABLE; INJECTION

LABETALOL HYDROCHLORIDE

AP	AKORN	5MG/ML	A075431 001	Nov 29, 1999
AP	BAXTER HLTHCARE CORP	5MG/ML	A076051 001	Jul 05, 2002
AP	CAPLIN	5MG/ML	A214533 001	Sep 07, 2021
AP	GLAND PHARMA LTD	5MG/ML	A090699 001	Apr 03, 2012
AP	HIKMA	5MG/ML	A075303 001	May 28, 1999
AP	! HOSPIRA	5MG/ML	A075239 001	Nov 29, 1999
AP	!	5MG/ML	A075240 001	Nov 29, 1999

SOLUTION; INTRAVENOUS

LABETALOL HYDROCHLORIDE IN DEXTROSE

	+! HIKMA	200MG/200ML (1MG/ML)	N213330 001	Nov 09, 2020
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PRESCRIPTION DRUG PRODUCT LIST

LABETALOL HYDROCHLORIDE

SOLUTION;INTRAVENOUS

LABETALOL HYDROCHLORIDE IN SODIUM CHLORIDE

+	!	HIKMA	100MG/100ML (1MG/ML)	N213330	002	Nov 09, 2020
+	!		200MG/200ML (1MG/ML)	N213330	003	Nov 09, 2020
+	!		300MG/300ML (1MG/ML)	N213330	004	Nov 09, 2020

TABLET;ORAL

LABETALOL HYDROCHLORIDE

<u>AB</u>		APPCO	<u>100MG</u>	<u>A209603</u>	<u>001</u>	Jun 20, 2018
<u>AB</u>			<u>200MG</u>	<u>A209603</u>	<u>002</u>	Jun 20, 2018
<u>AB</u>			<u>300MG</u>	<u>A209603</u>	<u>003</u>	Jun 20, 2018
<u>AB</u>		CADILA PHARMS LTD	<u>100MG</u>	<u>A211325</u>	<u>001</u>	May 13, 2019
<u>AB</u>			<u>200MG</u>	<u>A211325</u>	<u>002</u>	May 13, 2019
<u>AB</u>			<u>300MG</u>	<u>A211325</u>	<u>003</u>	May 13, 2019
<u>AB</u>		EPIC PHARMA LLC	<u>100MG</u>	<u>A212990</u>	<u>001</u>	Sep 30, 2020
<u>AB</u>			<u>200MG</u>	<u>A212990</u>	<u>002</u>	Sep 30, 2020
<u>AB</u>			<u>300MG</u>	<u>A212990</u>	<u>003</u>	Sep 30, 2020
<u>AB</u>		EYWA	<u>100MG</u>	<u>A207863</u>	<u>001</u>	Feb 04, 2019
<u>AB</u>			<u>200MG</u>	<u>A207863</u>	<u>002</u>	Feb 04, 2019
<u>AB</u>			<u>300MG</u>	<u>A207863</u>	<u>003</u>	Feb 04, 2019
<u>AB</u>		HERITAGE PHARMA	<u>100MG</u>	<u>A074787</u>	<u>001</u>	Aug 03, 1998
<u>AB</u>			<u>200MG</u>	<u>A074787</u>	<u>002</u>	Aug 03, 1998
<u>AB</u>			<u>300MG</u>	<u>A074787</u>	<u>003</u>	Aug 03, 1998
<u>AB</u>		INNOGENIX	<u>100MG</u>	<u>A075215</u>	<u>001</u>	Jul 29, 1999
<u>AB</u>			<u>200MG</u>	<u>A075215</u>	<u>002</u>	Jul 29, 1999
<u>AB</u>			<u>300MG</u>	<u>A075215</u>	<u>003</u>	Jul 29, 1999
<u>AB</u>		PAR FORM	<u>100MG</u>	<u>A200908</u>	<u>001</u>	Jul 10, 2012
<u>AB</u>	!		<u>200MG</u>	<u>A200908</u>	<u>002</u>	Jul 10, 2012
<u>AB</u>			<u>300MG</u>	<u>A200908</u>	<u>003</u>	Jul 10, 2012
<u>AB</u>		RUBICON	<u>100MG</u>	<u>A211953</u>	<u>001</u>	Aug 18, 2021
<u>AB</u>			<u>200MG</u>	<u>A211953</u>	<u>002</u>	Aug 18, 2021
<u>AB</u>			<u>300MG</u>	<u>A211953</u>	<u>003</u>	Aug 18, 2021
<u>AB</u>		SANDOZ	<u>100MG</u>	<u>A075113</u>	<u>001</u>	Aug 04, 1998
<u>AB</u>	!		<u>200MG</u>	<u>A075113</u>	<u>002</u>	Aug 04, 1998
<u>AB</u>			<u>300MG</u>	<u>A075113</u>	<u>003</u>	Aug 04, 1998
<u>AB</u>		WATSON LABS	<u>100MG</u>	<u>A075133</u>	<u>001</u>	Aug 03, 1998
<u>AB</u>			<u>200MG</u>	<u>A075133</u>	<u>002</u>	Aug 03, 1998
<u>AB</u>			<u>300MG</u>	<u>A075133</u>	<u>003</u>	Aug 03, 1998
<u>AB</u>		ZYDUS PHARMS	<u>100MG</u>	<u>A207743</u>	<u>001</u>	Sep 19, 2017
<u>AB</u>			<u>200MG</u>	<u>A207743</u>	<u>002</u>	Sep 19, 2017
<u>AB</u>			<u>300MG</u>	<u>A207743</u>	<u>003</u>	Sep 19, 2017

TRANDATE

<u>AB</u>	+	ALVOGEN	<u>200MG</u>	<u>N018716</u>	<u>002</u>	Aug 01, 1984
<u>AB</u>	+		<u>300MG</u>	<u>N018716</u>	<u>003</u>	Aug 01, 1984

LACOSAMIDE

SOLUTION;INTRAVENOUS

VIMPAT

+	!	UCB INC	200MG/20ML (10MG/ML)	N022254	001	Oct 28, 2008
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SOLUTION;ORAL

VIMPAT

+	!	UCB INC	10MG/ML	N022255	001	Apr 20, 2010
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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

LACTULOSE

FOR SOLUTION;ORAL

LACTULOSE

!		CUMBERLAND PHARMS	10GM/PACKET	A074712	001	Dec 10, 1997
!			20GM/PACKET	A074712	002	Dec 10, 1997

SOLUTION;ORAL

LACTULOSE

<u>AA</u>	!	AKORN	<u>10GM/15ML</u>	<u>A074076</u>	<u>001</u>	Jul 03, 1995
<u>AA</u>		CHARTWELL RX	<u>10GM/15ML</u>	<u>A209517</u>	<u>001</u>	Nov 23, 2018
<u>AA</u>		FRESENIUS KABI	<u>10GM/15ML</u>	<u>A090503</u>	<u>001</u>	Jan 25, 2012
<u>AA</u>		LANNETT CO INC	<u>10GM/15ML</u>	<u>A075993</u>	<u>001</u>	Jul 26, 2001
<u>AA</u>		PHARM ASSOC	<u>10GM/15ML</u>	<u>A074623</u>	<u>001</u>	Jul 30, 1996
<u>AA</u>		TORRENT	<u>10GM/15ML</u>	<u>A207786</u>	<u>001</u>	Jun 11, 2018
<u>AA</u>		VISTAPHARM	<u>10GM/15ML</u>	<u>A074138</u>	<u>001</u>	Sep 30, 1992

PRESCRIPTION DRUG PRODUCT LIST

LACTULOSE

SOLUTION;ORAL

LACTULOSE

AA	WOCKHARDT BIO AG	10GM/15ML	A074602 001	Nov 14, 1996
AA	XTRTRIUM LABS INC	10GM/15ML	A075911 001	Feb 21, 2002

SOLUTION;ORAL, RECTAL

GENERLAC

AA	WOCKHARDT BIO AG	10GM/15ML	A074603 001	Oct 31, 1996
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LACTULOSE

AA	AKORN	10GM/15ML	A074077 001	Jul 03, 1995
AA	! FRESINIUS KABI	10GM/15ML	A090502 001	Jan 25, 2012
AA	TORRENT	10GM/15ML	A203762 001	Mar 27, 2015

LAMIVUDINE

SOLUTION;ORAL

EPIVIR

AA	+! VIIV HLTHCARE	10MG/ML	N020596 001	Nov 17, 1995
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LAMIVUDINE

AA	AUROBINDO PHARMA LTD	10MG/ML	A077695 001	Nov 21, 2016
AA	LANNETT CO INC	10MG/ML	A203564 001	Oct 31, 2014

EPIVIR-HBV

+!	GLAXOSMITHKLINE	5MG/ML	N021004 001	Dec 08, 1998
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TABLET;ORAL

EPIVIR

AB	+ VIIV HLTHCARE	150MG	N020564 001	Nov 17, 1995
AB	+!	300MG	N020564 003	Jun 24, 2002

EPIVIR-HBV

AB	+! GLAXOSMITHKLINE	100MG	N021003 001	Dec 08, 1998
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LAMIVUDINE

AB	ANNORA	100MG	A211306 001	Mar 21, 2019
AB	APOTEX	100MG	A202941 001	Jan 02, 2014
AB		150MG	A091606 001	Dec 02, 2011
AB		300MG	A091606 002	Dec 02, 2011
AB	APPCO	150MG	A206974 001	Nov 21, 2016
AB		300MG	A206974 002	Nov 21, 2016
AB	AUROBINDO PHARMA LTD	150MG	A077464 001	Nov 21, 2016
AB		150MG	A202032 001	Nov 17, 2011
AB		300MG	A077464 002	Nov 21, 2016
AB		300MG	A202032 002	Nov 17, 2011
AB	BRECKENRIDGE	150MG	A203586 001	Nov 21, 2016
AB	CIPLA	150MG	A077221 001	Mar 03, 2017
AB		300MG	A077221 002	Mar 03, 2017
AB	HETERO LABS LTD V	100MG	A203260 001	Jan 02, 2014
AB		150MG	A203277 001	Jan 06, 2014
AB		300MG	A203277 002	Jan 06, 2014
AB	LUPIN LTD	150MG	A205217 001	Dec 18, 2014
AB		300MG	A205217 002	Dec 18, 2014
AB	MACLEODS PHARMS LTD	150MG	A090198 001	May 01, 2019
AB		300MG	A090198 002	May 01, 2019
AB	MYLAN LABS LTD	150MG	A078545 001	Mar 05, 2019
AB		300MG	A078545 002	Mar 05, 2019
AB	STRIDES PHARMA	150MG	A090457 001	Apr 19, 2018
AB		300MG	A090457 002	Apr 19, 2018

LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

CIMDUO

+!	MYLAN LABS LTD	300MG;300MG	N022141 001	Feb 28, 2018
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LAMIVUDINE; ZIDOVUDINE

TABLET;ORAL

COMBIVIR

AB	+! VIIV HLTHCARE	150MG;300MG	N020857 001	Sep 26, 1997
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LAMIVUDINE AND ZIDOVUDINE

AB	ANDA REPOSITORY	150MG;300MG	A206375 001	Apr 10, 2018
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

PRESCRIPTION DRUG PRODUCT LIST

LAMIVUDINE; ZIDOVUDINE

TABLET;ORAL

LAMIVUDINE AND ZIDOVUDINE

<u>AB</u>	MACLEODS PHARMS LTD	<u>150MG;300MG</u>	<u>A090679 001</u>	Aug 29, 2018
<u>AB</u>	STRIDES PHARMA	<u>150MG;300MG</u>	<u>A079128 001</u>	May 13, 2015

LAMOTRIGINE

TABLET;ORAL

LAMICTAL

<u>AB</u>	+!	GLAXOSMITHKLINE LLC	<u>25MG</u>	<u>N020241 005</u>	Dec 27, 1994
<u>AB</u>	+		<u>100MG</u>	<u>N020241 001</u>	Dec 27, 1994
<u>AB</u>	+		<u>150MG</u>	<u>N020241 002</u>	Dec 27, 1994
<u>AB</u>	+		<u>200MG</u>	<u>N020241 003</u>	Dec 27, 1994

LAMOTRIGINE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A090607 001</u>	Jan 13, 2011
<u>AB</u>			<u>100MG</u>	<u>A090607 002</u>	Jan 13, 2011
<u>AB</u>			<u>150MG</u>	<u>A090607 003</u>	Jan 13, 2011
<u>AB</u>			<u>200MG</u>	<u>A090607 004</u>	Jan 13, 2011
<u>AB</u>		ALKEM LABS LTD	<u>25MG</u>	<u>A200694 001</u>	Jun 14, 2013
<u>AB</u>			<u>100MG</u>	<u>A200694 002</u>	Jun 14, 2013
<u>AB</u>			<u>150MG</u>	<u>A200694 003</u>	Jun 14, 2013
<u>AB</u>			<u>200MG</u>	<u>A200694 004</u>	Jun 14, 2013
<u>AB</u>		AUROBINDO PHARMA	<u>25MG</u>	<u>A078956 001</u>	Jan 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A078956 002</u>	Jan 27, 2009
<u>AB</u>			<u>150MG</u>	<u>A078956 003</u>	Jan 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A078956 004</u>	Jan 27, 2009
<u>AB</u>		DR REDDYS LABS LTD	<u>25MG</u>	<u>A076708 001</u>	Jan 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A076708 002</u>	Jan 27, 2009
<u>AB</u>			<u>150MG</u>	<u>A076708 003</u>	Jan 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A076708 004</u>	Jan 27, 2009
<u>AB</u>		GLENMARK GENERICS	<u>25MG</u>	<u>A090169 001</u>	May 12, 2012
<u>AB</u>			<u>100MG</u>	<u>A090169 002</u>	May 12, 2012
<u>AB</u>			<u>150MG</u>	<u>A090169 003</u>	May 12, 2012
<u>AB</u>			<u>200MG</u>	<u>A090169 004</u>	May 12, 2012
<u>AB</u>		JUBILANT CADISTA	<u>25MG</u>	<u>A079132 001</u>	Jan 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A079132 002</u>	Jan 27, 2009
<u>AB</u>			<u>150MG</u>	<u>A079132 003</u>	Jan 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A079132 004</u>	Jan 27, 2009
<u>AB</u>		LUPIN LTD	<u>25MG</u>	<u>A078691 001</u>	Jun 01, 2010
<u>AB</u>			<u>100MG</u>	<u>A078691 002</u>	Jun 01, 2010
<u>AB</u>			<u>150MG</u>	<u>A078691 003</u>	Jun 01, 2010
<u>AB</u>			<u>200MG</u>	<u>A078691 004</u>	Jun 01, 2010
<u>AB</u>		RUBICON	<u>25MG</u>	<u>A078625 001</u>	Jan 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A078625 002</u>	Jan 27, 2009
<u>AB</u>			<u>150MG</u>	<u>A078625 003</u>	Jan 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A078625 004</u>	Jan 27, 2009
<u>AB</u>		TARO PHARM INDS	<u>25MG</u>	<u>A078525 001</u>	Jan 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A078525 002</u>	Jan 27, 2009
<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>		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

PRESCRIPTION DRUG PRODUCT LIST

LAMOTRIGINE

TABLET, CHEWABLE;ORAL

LAMOTRIGINE

<u>AB</u>		<u>25MG</u>	<u>A090401</u>	<u>003</u>	Nov 04, 2009
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A076701</u>	<u>001</u>	Jan 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A076701</u>	<u>002</u>	Jan 22, 2009
<u>AB</u>	GLENMARK PHARMS LTD	<u>5MG</u>	<u>A079099</u>	<u>001</u>	Feb 19, 2009
<u>AB</u>		<u>25MG</u>	<u>A079099</u>	<u>002</u>	Feb 19, 2009
<u>AB</u>	TARO	<u>5MG</u>	<u>A079204</u>	<u>001</u>	Feb 04, 2009
<u>AB</u>		<u>25MG</u>	<u>A079204</u>	<u>002</u>	Feb 04, 2009
<u>AB</u>	WATSON LABS	<u>2MG</u>	<u>A076928</u>	<u>001</u>	Jan 22, 2009
<u>AB</u>		<u>5MG</u>	<u>A076928</u>	<u>002</u>	Jan 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A076928</u>	<u>003</u>	Jan 22, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>5MG</u>	<u>A078009</u>	<u>002</u>	Jan 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A078009</u>	<u>003</u>	Jan 22, 2009

TABLET, EXTENDED RELEASE;ORAL

LAMICTAL XR

<u>AB</u>	+	GLAXOSMITHKLINE LLC	<u>25MG</u>	<u>N022115</u>	<u>001</u>	May 29, 2009
<u>AB</u>	+		<u>50MG</u>	<u>N022115</u>	<u>002</u>	May 29, 2009
<u>AB</u>	+		<u>100MG</u>	<u>N022115</u>	<u>003</u>	May 29, 2009
<u>AB</u>	+		<u>200MG</u>	<u>N022115</u>	<u>004</u>	May 29, 2009
<u>AB</u>	+		<u>250MG</u>	<u>N022115</u>	<u>006</u>	Jun 21, 2011
<u>AB</u>	+		<u>300MG</u>	<u>N022115</u>	<u>005</u>	Apr 14, 2010

LAMOTRIGINE

<u>AB</u>		ACTAVIS ELIZABETH	<u>100MG</u>	<u>A200672</u>	<u>003</u>	Oct 17, 2013
<u>AB</u>			<u>200MG</u>	<u>A200672</u>	<u>004</u>	Oct 17, 2013
<u>AB</u>			<u>25MG</u>	<u>A200672</u>	<u>001</u>	Oct 17, 2013
<u>AB</u>			<u>50MG</u>	<u>A200672</u>	<u>002</u>	Oct 17, 2013
<u>AB</u>			<u>250MG</u>	<u>A200672</u>	<u>006</u>	Nov 13, 2013
<u>AB</u>			<u>300MG</u>	<u>A200672</u>	<u>005</u>	Oct 17, 2013
<u>AB</u>		AMNEAL PHARMS	<u>25MG</u>	<u>A207497</u>	<u>001</u>	Nov 30, 2018
<u>AB</u>			<u>50MG</u>	<u>A207497</u>	<u>002</u>	Nov 30, 2018
<u>AB</u>			<u>100MG</u>	<u>A207497</u>	<u>003</u>	Nov 30, 2018
<u>AB</u>			<u>200MG</u>	<u>A207497</u>	<u>004</u>	Nov 30, 2018
<u>AB</u>			<u>250MG</u>	<u>A207497</u>	<u>005</u>	Nov 30, 2018
<u>AB</u>			<u>300MG</u>	<u>A207497</u>	<u>006</u>	Nov 30, 2018
<u>AB</u>		ANCHEN PHARMS	<u>25MG</u>	<u>A201374</u>	<u>001</u>	Dec 26, 2012
<u>AB</u>			<u>50MG</u>	<u>A201374</u>	<u>002</u>	Dec 26, 2012
<u>AB</u>			<u>100MG</u>	<u>A201374</u>	<u>003</u>	Dec 26, 2012
<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>		ATLANTIDE	<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
<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>250MG</u>	<u>A202383</u>	<u>006</u>	Sep 06, 2018
<u>AB</u>			<u>300MG</u>	<u>A202383</u>	<u>005</u>	Jun 19, 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>		RUBICON	<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>		TORRENT	<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>		YILING	<u>25MG</u>	<u>A213949</u>	<u>001</u>	Dec 08, 2021
<u>AB</u>			<u>50MG</u>	<u>A213949</u>	<u>002</u>	Dec 08, 2021
<u>AB</u>			<u>100MG</u>	<u>A213949</u>	<u>003</u>	Dec 08, 2021
<u>AB</u>			<u>200MG</u>	<u>A213949</u>	<u>004</u>	Dec 08, 2021
<u>AB</u>			<u>250MG</u>	<u>A213949</u>	<u>005</u>	Dec 08, 2021
<u>AB</u>			<u>300MG</u>	<u>A213949</u>	<u>006</u>	Dec 08, 2021
<u>AB</u>		ZYDUS PHARMS	<u>25MG</u>	<u>A207763</u>	<u>001</u>	Apr 01, 2020

PRESCRIPTION DRUG PRODUCT LIST

LAMOTRIGINE

TABLET, EXTENDED RELEASE;ORAL

LAMOTRIGINE

<u>AB</u>		<u>50MG</u>	<u>A207763</u>	<u>002</u>	Apr 01, 2020
<u>AB</u>		<u>100MG</u>	<u>A207763</u>	<u>003</u>	Apr 01, 2020
<u>AB</u>		<u>200MG</u>	<u>A207763</u>	<u>004</u>	Apr 01, 2020
<u>AB</u>		<u>250MG</u>	<u>A207763</u>	<u>005</u>	Apr 01, 2020
<u>AB</u>		<u>300MG</u>	<u>A207763</u>	<u>006</u>	Apr 01, 2020

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>		AJANTA PHARMA LTD	<u>25MG</u>	<u>A213271</u>	<u>001</u>	Jan 19, 2021
<u>AB</u>			<u>50MG</u>	<u>A213271</u>	<u>002</u>	Jan 19, 2021
<u>AB</u>			<u>100MG</u>	<u>A213271</u>	<u>003</u>	Jan 19, 2021
<u>AB</u>			<u>200MG</u>	<u>A213271</u>	<u>004</u>	Jan 19, 2021
<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

LANREOTIDE ACETATE

+	!	INVAGEN PHARMS	EQ 60MG BASE/0.2ML (EQ 60MG BASE/0.2ML)	N215395	001	Dec 17, 2021
+	!		EQ 90MG BASE/0.3ML (EQ 90MG BASE/0.3ML)	N215395	002	Dec 17, 2021
+	!		EQ 120MG BASE/0.5ML (EQ 120MG BASE/0.5ML)	N215395	003	Dec 17, 2021

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>		ALKEM LABS LTD	<u>15MG</u>	<u>A207394</u>	<u>001</u>	Jan 18, 2019
<u>AB</u>			<u>30MG</u>	<u>A207394</u>	<u>002</u>	Jan 18, 2019
<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>		HETERO LABS LTD III	<u>15MG</u>	<u>A203083</u>	<u>001</u>	May 18, 2020
<u>AB</u>			<u>30MG</u>	<u>A203083</u>	<u>002</u>	May 18, 2020
<u>AB</u>		INVENTIA	<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>		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
<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	<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>		XIROMED	<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>		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

PRESCRIPTION DRUG PRODUCT LIST

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

PREVACID

AB	+ !	TAKEDA PHARMS USA	30MG	N020406	002	May 10, 1995
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TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

AB		DR REDDYS LABS LTD	15MG	A210465	001	Feb 01, 2021
AB			30MG	A210465	002	Feb 01, 2021
AB		MYLAN	15MG	A202396	001	Nov 28, 2018
AB			30MG	A202396	002	Nov 28, 2018
AB		TEVA PHARMS USA	15MG	A208784	001	Sep 21, 2017
AB			30MG	A208784	002	Sep 21, 2017
AB		ZYDUS PHARMS	15MG	A200816	001	Nov 27, 2018
AB			30MG	A200816	002	Nov 27, 2018

PREVACID

AB	+	TAKEDA PHARMS USA	15MG	N021428	001	Aug 30, 2002
AB	+ !		30MG	N021428	002	Aug 30, 2002

LANTHANUM CARBONATE

POWDER;ORAL

FOSRENOL

+		TAKEDA PHARMS USA	EQ 750MG BASE	N204734	001	Sep 24, 2014
+ !			EQ 1GM BASE	N204734	002	Sep 24, 2014

TABLET, CHEWABLE;ORAL

FOSRENOL

AB	+	TAKEDA PHARMS USA	EQ 500MG BASE	N021468	002	Oct 26, 2004
AB	+		EQ 750MG BASE	N021468	003	Nov 23, 2005
AB	+ !		EQ 1GM BASE	N021468	004	Nov 23, 2005

LANTHANUM CARBONATE

AB		NATCO PHARMA LTD	EQ 500MG BASE	A090978	001	Aug 11, 2017
AB			EQ 750MG BASE	A090978	002	Aug 11, 2017
AB			EQ 1GM BASE	A090978	003	Aug 11, 2017

LAPATINIB DITOSYLATE

TABLET;ORAL

LAPATINIB DITOSYLATE

AB		NATCO PHARMA LTD	EQ 250MG BASE	A203007	001	Sep 29, 2020
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TYKERB

AB	+ !	NOVARTIS	EQ 250MG BASE	N022059	001	Mar 13, 2007
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LAROTRECTINIB SULFATE

CAPSULE;ORAL

VITRAKVI

+		BAYER HLTHCARE	EQ 25MG BASE	N210861	001	Nov 26, 2018
+ !			EQ 100MG BASE	N210861	002	Nov 26, 2018

SOLUTION;ORAL

VITRAKVI

+ !		BAYER HEALTHCARE	EQ 20MG BASE/ML	N211710	001	Nov 26, 2018
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LASMIDITAN SUCCINATE

TABLET;ORAL

REYVOW

+		ELI LILLY AND CO	EQ 50MG BASE	N211280	001	Jan 31, 2020
+ !			EQ 100MG BASE	N211280	002	Jan 31, 2020

LATANOPROST

EMULSION;OPHTHALMIC

XELPROS

+ !		SUN PHARM	0.005%	N206185	001	Sep 12, 2018
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SOLUTION/DROPS;OPHTHALMIC

LATANOPROST

AT		AKORN	0.005%	A090887	001	Jul 19, 2011
AT		AMRING PHARMS	0.005%	A200925	001	Mar 22, 2011
AT		BAUSCH AND LOMB	0.005%	A201006	001	Mar 22, 2011
AT		DR REDDYS LABS LTD	0.005%	A202077	001	Feb 11, 2013
AT		FDC LTD	0.005%	A202442	001	Apr 22, 2016
AT		SANDOZ INC	0.005%	A091449	001	Mar 22, 2011
AT		SOMERSET	0.005%	A201786	001	Mar 22, 2011

XALATAN

AT	+ !	UPJOHN	0.005%	N020597	001	Jun 05, 1996
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PRESCRIPTION DRUG PRODUCT LIST

LATANOPROST; NETARSUDIL DIMESYLATE

SOLUTION/DROPS;OPHTHALMIC

ROCKLATAN

+! AERIE PHARMS INC 0.005%;EQ 0.02% BASE N208259 001 Mar 12, 2019

LATANOPROSTENE BUNOD

SOLUTION/DROPS;OPHTHALMIC

VYZULTA

+! BAUSCH AND LOMB 0.024% N207795 001 Nov 02, 2017

LEDIPASVIR; SOFOSBUVIR

PELLETS;ORAL

HARVONI

+ GILEAD SCIENCES INC 33.75MG;150MG/PACKET N212477 001 Aug 28, 2019

+! 45MG;200MG/PACKET N212477 002 Aug 28, 2019

TABLET;ORAL

HARVONI

+ GILEAD SCIENCES INC 45MG;200MG N205834 002 Aug 28, 2019

+! 90MG;400MG N205834 001 Oct 10, 2014

LEFAMULIN ACETATE

SOLUTION;INTRAVENOUS

XENLETA

+! NABRIVA EQ 150MG BASE/15ML (EQ 10MG BASE/ML) N211673 001 Aug 19, 2019

TABLET;ORAL

XENLETA

+! NABRIVA EQ 600MG BASE N211672 001 Aug 19, 2019

LEFLUNOMIDE

TABLET;ORAL

ARAVA**AB** + SANOFI AVENTIS US **10MG** **N020905 001** Sep 10, 1998**AB** +! **20MG** **N020905 002** Sep 10, 1998LEFLUNOMIDE**AB** ABHAI LLC **10MG** **A212453 001** Jun 03, 2019**AB** **20MG** **A212453 002** Jun 03, 2019**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** AUROBINDO PHARMA LTD **10MG** **A213652 001** Mar 29, 2021**AB** **20MG** **A213652 002** Mar 29, 2021**AB** FOSUN PHARMA **10MG** **A077087 001** Sep 13, 2005**AB** **20MG** **A077087 002** Sep 13, 2005**AB** HERITAGE PHARMS INC **10MG** **A077086 001** Sep 13, 2005**AB** **20MG** **A077086 002** Sep 13, 2005**AB** LUPIN LTD **10MG** **A211863 001** Feb 04, 2020**AB** **20MG** **A211863 002** Feb 04, 2020**AB** ZYDUS **10MG** **A212308 001** Apr 24, 2019**AB** **20MG** **A212308 002** Apr 24, 2019

ARAVA

+! SANOFI AVENTIS US 100MG N020905 003 Sep 10, 1998

LEMBOREXANT

TABLET;ORAL

DAYVIGO

+ EISAI INC 5MG N212028 001 Apr 07, 2020

+! 10MG N212028 002 Apr 07, 2020

LENALIDOMIDE

CAPSULE;ORAL

LENALIDOMIDE**AB** ARROW INTL **5MG** **A201452 001** May 21, 2021**AB** **10MG** **A201452 002** May 21, 2021**AB** **15MG** **A201452 003** May 21, 2021**AB** **25MG** **A201452 004** May 21, 2021**AB** DR REDDYS **2.5MG** **A209348 001** Oct 14, 2021**AB** **20MG** **A209348 002** Oct 14, 2021REVLIMID**AB** + CELGENE **2.5MG** **N021880 005** Dec 21, 2011**AB** + **5MG** **N021880 001** Dec 27, 2005**AB** + **10MG** **N021880 002** Dec 27, 2005**AB** + **15MG** **N021880 003** Jun 29, 2006**AB** + **20MG** **N021880 006** Jun 05, 2013**AB** +! **25MG** **N021880 004** Jun 29, 2006

PRESCRIPTION DRUG PRODUCT LIST

LENVATINIB MESYLATE

CAPSULE;ORAL

LENVIMA

+ EISAI INC

EQ 4MG BASE

N206947 001 Feb 13, 2015

+!

EQ 10MG BASE

N206947 002 Feb 13, 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

FEMARAAB +! NOVARTIS PHARMS2.5MGN020726 001 Jul 25, 1997LETROZOLEAB ACCORD HLTHCARE2.5MGA090934 001 Jun 03, 2011AB BEIJING YILING2.5MGA205869 001 Nov 14, 2018AB DR REDDYS LABS LTD2.5MGA091191 001 Jun 03, 2011AB EUGIA PHARMA2.5MGA211717 001 Jan 11, 2019AB INDICUS PHARMA2.5MGA201804 001 Jun 03, 2011AB NATCO PHARMA LTD2.5MGA200161 001 Jun 03, 2011AB TEVA PHARMS2.5MGA090289 001 Jun 03, 2011LETROZOLE; RIBOCICLIB SUCCINATE

TABLET;ORAL

KISQALI FEMARA CO-PACK (COPACKAGED)

+! NOVARTIS

2.5MG;EQ 200MG BASE

N209935 001 May 04, 2017

LEUCOVORIN CALCIUM

INJECTABLE;INJECTION

LEUCOVORIN CALCIUMAP FRESENIUS KABI USAEQ 10MG BASE/MLA207226 001 Jul 27, 2018AP ! HIKMAEQ 50MG BASE/VIALA089384 001 Sep 14, 1987AP !EQ 100MG BASE/VIALA089717 001 Mar 28, 1988AP TEVA PHARMS USAEQ 100MG BASE/VIALA081277 001 Sep 28, 1993APEQ 350MG BASE/VIALA040174 001 Jun 12, 1997LEUCOVORIN CALCIUM PRESERVATIVE FREEAP FRESENIUS KABI USAEQ 200MG BASE/VIALA040258 001 Feb 26, 1999AP !EQ 500MG BASE/VIALA040286 001 Feb 26, 1999AP ! HIKMAEQ 10MG BASE/MLA040347 001 Apr 25, 2000AP !EQ 200MG BASE/VIALA040056 001 May 23, 1995AP !EQ 350MG BASE/VIALA040335 001 Apr 20, 2000AP MYLAN LABS LTDEQ 100MG BASE/VIALA203800 001 May 19, 2017APEQ 200MG BASE/VIALA203800 002 May 19, 2017APEQ 350MG BASE/VIALA203800 003 May 19, 2017AP SAGENT PHARMSEQ 50MG BASE/VIALA200753 001 Sep 06, 2012APEQ 100MG BASE/VIALA200753 002 Sep 06, 2012APEQ 200MG BASE/VIALA200753 003 Sep 06, 2012APEQ 350MG BASE/VIALA200855 001 Sep 06, 2012AP SAGENT PHARMS INCEQ 500MG BASE/VIALA209110 001 Oct 26, 2017

LEUCOVORIN CALCIUM

FRESENIUS KABI USA

EQ 10MG BASE/ML

A207241 001 Mar 14, 2018

TABLET;ORAL

LEUCOVORIN CALCIUMAB BARREQ 5MG BASEA071198 001 Sep 24, 1987ABEQ 25MG BASEA071199 001 Sep 24, 1987AB EPIC PHARMA LLCEQ 5MG BASEA074544 001 Aug 28, 1997ABEQ 10MG BASEA074544 003 May 19, 2021ABEQ 15MG BASEA074544 004 May 19, 2021ABEQ 25MG BASEA074544 002 Aug 28, 1997ABEQ 5MG BASEA072733 001 Feb 22, 1993ABEQ 10MG BASEA072734 001 Feb 22, 1993ABEQ 15MG BASEA072735 001 Feb 22, 1993AB !EQ 25MG BASEA072736 001 Feb 22, 1993ABEQ 5MG BASEA213929 001 Oct 22, 2020ABEQ 10MG BASEA213929 002 Oct 22, 2020ABEQ 15MG BASEA213929 003 Oct 22, 2020ABEQ 25MG BASEA213929 004 Oct 22, 2020

PRESCRIPTION DRUG PRODUCT LIST

LEUCOVORIN CALCIUM

TABLET; ORAL

LEUCOVORIN CALCIUM

<u>AB</u>	NOVAST LABS	<u>EQ 5MG BASE</u>	<u>A211132 001</u>	Jul 30, 2020
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A211132 002</u>	Jul 30, 2020
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A211132 003</u>	Jul 30, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A211132 004</u>	Jul 30, 2020

LEUPROLIDE ACETATE

INJECTABLE; INJECTION

LEUPROLIDE ACETATE

<u>AP</u>	! SANDOZ	<u>1MG/0.2ML</u>	<u>A074728 001</u>	Aug 04, 1998
<u>AP</u>	SUN PHARM	<u>1MG/0.2ML</u>	<u>A078885 001</u>	Mar 09, 2009
<u>AP</u>	TEVA PHARMS USA	<u>1MG/0.2ML</u>	<u>A075471 001</u>	Oct 25, 2000
<u>AP</u>	VGYAAN	<u>1MG/0.2ML</u>	<u>A213829 001</u>	Aug 13, 2021

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

POWDER; INTRAMUSCULAR

LUPRON DEPOT-PED KIT

+!	ABBVIE ENDOCRINE INC	7.5MG	N020263 002	Apr 16, 1993
+!		11.25MG	N020263 005	Jan 21, 1994
+!		11.25MG	N020263 007	Aug 15, 2011
+!		15MG	N020263 006	Jan 21, 1994
+!		30MG	N020263 008	Aug 15, 2011

POWDER; SUBCUTANEOUS

ELIGARD KIT

+!	TOLMAR THERAP	7.5MG	N021343 001	Jan 23, 2002
+!		22.5MG	N021379 001	Jul 24, 2002
+!		30MG	N021488 001	Feb 13, 2003
+!		45MG	N021731 001	Dec 14, 2004

FENSOLVI KIT

+!	TOLMAR	45MG	N213150 001	May 01, 2020
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LEUPROLIDE MESYLATE

EMULSION; SUBCUTANEOUS

CAMCEVI KIT

+!	FORESEE PHARMS	EQ 42MG BASE	N211488 001	May 25, 2021
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LEVALBUTEROL HYDROCHLORIDE

SOLUTION; INHALATION

LEVALBUTEROL HYDROCHLORIDE

<u>AN</u>	CIPLA	<u>EQ 0.021% BASE</u>	<u>A078171 002</u>	Dec 13, 2013
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A078171 003</u>	Dec 13, 2013
<u>AN</u>		<u>EQ 0.0103% BASE</u>	<u>A078171 001</u>	Dec 13, 2013
<u>AN</u>	IMPAX LABS INC	<u>EQ 0.0103% BASE</u>	<u>A077756 003</u>	Apr 09, 2008
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A077756 001</u>	Apr 09, 2008
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A077756 002</u>	Apr 09, 2008
<u>AN</u>	LUOXIN AUROVITAS	<u>EQ 0.25% BASE</u>	<u>A207628 001</u>	Jan 31, 2017
<u>AN</u>		<u>EQ 0.0103% BASE</u>	<u>A207625 001</u>	Dec 30, 2016
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A207625 002</u>	Dec 30, 2016
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A207625 003</u>	Dec 30, 2016
<u>AN</u>	MYLAN SPECIALITY LP	<u>EQ 0.25% BASE</u>	<u>A078309 001</u>	Mar 20, 2009
<u>AN</u>	RITEDOSE CORP	<u>EQ 0.0103% BASE</u>	<u>A203653 001</u>	Mar 22, 2016
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A203653 002</u>	Mar 22, 2016
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A203653 003</u>	Mar 22, 2016
<u>AN</u>	SUN PHARM	<u>EQ 0.0103% BASE</u>	<u>A207820 001</u>	Nov 05, 2018
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A207820 002</u>	Nov 05, 2018
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A207820 003</u>	Nov 05, 2018
<u>AN</u>	TEVA PARENTERAL	<u>EQ 0.25% BASE</u>	<u>A200875 001</u>	Sep 11, 2014
<u>AN</u>	TEVA PHARMS USA	<u>EQ 0.0103% BASE</u>	<u>A090297 001</u>	Apr 26, 2013
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A090297 002</u>	Apr 26, 2013
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A090297 003</u>	Apr 26, 2013

XOPENEX

<u>AN</u>	+! AKORN	<u>EQ 0.0103% BASE</u>	<u>N020837 003</u>	Jan 30, 2002
<u>AN</u>	+!	<u>EQ 0.021% BASE</u>	<u>N020837 001</u>	Mar 25, 1999
<u>AN</u>	+!	<u>EQ 0.042% BASE</u>	<u>N020837 002</u>	Mar 25, 1999
<u>AN</u>	+!	<u>EQ 0.25% BASE</u>	<u>N020837 004</u>	Jul 18, 2003

PRESCRIPTION DRUG PRODUCT LIST

LEVALBUTEROL TARTRATE

AEROSOL, METERED; INHALATION

XOPENEX HFA

+! SUNOVION

EQ 0.045MG BASE/INH

N021730 001 Mar 11, 2005

LEVAMLODIPINE MALEATE

TABLET; ORAL

CONJUPRI

+ CSPEC OUYI

EQ 2.5MG BASE

N212895 002 Dec 19, 2019

+!

EQ 5MG BASE

N212895 003 Dec 19, 2019

LEVETIRACETAM

INJECTABLE; INTRAVENOUS

KEPPRAAP +! UCB INC500MG/5ML (100MG/ML)N021872 001 Jul 31, 2006LEVETIRACETAMAP EUGIA PHARMA500MG/5ML (100MG/ML)A204312 001 Feb 01, 2016AP FRESenius KABI USA500MG/5ML (100MG/ML)A090876 001 Aug 13, 2015AP HAINAN POLY PHARM500MG/5ML (100MG/ML)A209781 001 Mar 20, 2018AP HIKMA FARMACEUTICA500MG/5ML (100MG/ML)A090981 001 Oct 13, 2011AP HOSPIRA INC500MG/5ML (100MG/ML)A202869 001 Apr 06, 2012AP MICRO LABS500MG/5ML (100MG/ML)A211954 001 Aug 09, 2019AP MYLAN LABS LTD500MG/5ML (100MG/ML)A203308 001 Sep 16, 2016AP SAGENT PHARMS500MG/5ML (100MG/ML)A091627 001 Jun 26, 2013AP SUN PHARM INDS LTD500MG/5ML (100MG/ML)A090754 001 Jun 16, 2010AP XGEN PHARMS500MG/5ML (100MG/ML)A091485 001 Aug 05, 2011LEVETIRACETAM IN SODIUM CHLORIDEAP EUGIA PHARMA500MG/100ML (5MG/ML)A207160 001 Jan 04, 2017AP1GM/100ML (10MG/ML)A207160 002 Jan 04, 2017AP1.5GM/100ML (15MG/ML)A207160 003 Jan 04, 2017AP GLAND PHARMA LTD500MG/100ML (5MG/ML)A206880 001 Oct 25, 2017AP1GM/100ML (10MG/ML)A206880 002 Oct 25, 2017AP1.5GM/100ML (15MG/ML)A206880 003 Oct 25, 2017AP +! HQ SPECIALITY500MG/100ML (5MG/ML)N202543 001 Nov 09, 2011

PHARMA

AP +!1GM/100ML (10MG/ML)N202543 002 Nov 09, 2011AP +!1.5GM/100ML (15MG/ML)N202543 003 Nov 09, 2011AP NEXUS PHARMS500MG/100ML (5MG/ML)A213532 001 Jul 06, 2020AP1GM/100ML (10MG/ML)A213532 002 Jul 06, 2020AP1.5GM/100ML (15MG/ML)A213532 003 Jul 06, 2020

+! HQ SPECIALITY

250MG/50ML (5MG/ML)

N202543 004 Dec 14, 2020

PHARMA

SOLUTION; ORAL

KEPPRAAA +! UCB INC100MG/MLN021505 001 Jul 15, 2003LEVETIRACETAMAA ACI100MG/MLA078582 001 Jan 15, 2009AA ACTAVIS MID100MG/MLA078976 001 Jan 15, 2009

ATLANTIC

AA AKORN100MG/MLA090601 001 Feb 28, 2012AA ALEMbic LABS100MG/MLA203067 001 May 09, 2013AA AMNEAL PHARMS100MG/MLA090992 001 Oct 27, 2009AA AUROBINDO PHARMA100MG/MLA079063 001 Jan 15, 2009

LTD

AA BRECKENRIDGE100MG/MLA079120 001 Jan 16, 2009AA GENUS100MG/MLA090079 001 Apr 11, 2012AA HETERO LABS LTD III100MG/MLA203052 001 Feb 28, 2013AA LANNETT CO INC100MG/MLA090263 001 Apr 03, 2009AA LUPIN LTD100MG/MLA090893 001 Oct 17, 2011AA PHARM ASSOC100MG/MLA201157 001 Jun 04, 2015AA TARO100MG/MLA078774 001 Feb 10, 2009AA TRIS PHARMA INC100MG/MLA090461 001 Sep 30, 2010AA WOCKHARDT BIO AG100MG/MLA090028 001 Mar 03, 2010

TABLET; ORAL

KEPPRAAB + UCB INC250MGN021035 001 Nov 30, 1999AB +500MGN021035 002 Nov 30, 1999AB +750MGN021035 003 Nov 30, 1999AB +!1GMN021035 004 Jan 06, 2006LEVETIRACETAMAB ACCORD HLTHCARE250MGA090843 001 Feb 14, 2011AB500MGA090843 002 Feb 14, 2011AB750MGA090843 003 Feb 14, 2011AB1GMA090843 004 Feb 14, 2011

PRESCRIPTION DRUG PRODUCT LIST

LEVETIRACETAM

TABLET; ORAL

LEVETIRACETAM

<u>AB</u>	ACI	<u>250MG</u>	<u>A078042</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078042</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078042</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A078042</u>	<u>004</u>	Jan 15, 2009
<u>AB</u>	AUROBINDO PHARMA	<u>250MG</u>	<u>A078993</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078993</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078993</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A078993</u>	<u>004</u>	Jan 15, 2009
<u>AB</u>	CHARTWELL RX	<u>250MG</u>	<u>A201293</u>	<u>001</u>	Jun 14, 2011
<u>AB</u>		<u>500MG</u>	<u>A201293</u>	<u>002</u>	Jun 14, 2011
<u>AB</u>		<u>750MG</u>	<u>A201293</u>	<u>003</u>	Jun 14, 2011
<u>AB</u>		<u>1GM</u>	<u>A201293</u>	<u>004</u>	Jun 14, 2011
<u>AB</u>	DR REDDYS LABS LTD	<u>250MG</u>	<u>A076920</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A076920</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A076920</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A078904</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>	HETERO LABS LTD III	<u>250MG</u>	<u>A090515</u>	<u>001</u>	Oct 08, 2010
<u>AB</u>		<u>500MG</u>	<u>A090515</u>	<u>002</u>	Oct 08, 2010
<u>AB</u>		<u>750MG</u>	<u>A090515</u>	<u>003</u>	Oct 08, 2010
<u>AB</u>		<u>1GM</u>	<u>A090515</u>	<u>004</u>	Oct 08, 2010
<u>AB</u>	INVAGEN PHARMS	<u>250MG</u>	<u>A078234</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078234</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078234</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>	LOTUS PHARM CO LTD	<u>500MG</u>	<u>A090906</u>	<u>001</u>	Nov 05, 2010
<u>AB</u>	LUPIN	<u>250MG</u>	<u>A078154</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078154</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078154</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A090025</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>	MYLAN	<u>250MG</u>	<u>A076919</u>	<u>001</u>	Nov 04, 2008
<u>AB</u>		<u>500MG</u>	<u>A076919</u>	<u>002</u>	Nov 04, 2008
<u>AB</u>		<u>750MG</u>	<u>A076919</u>	<u>003</u>	Nov 04, 2008
<u>AB</u>		<u>1GM</u>	<u>A090261</u>	<u>001</u>	Dec 08, 2009
<u>AB</u>	ORBION PHARMS	<u>250MG</u>	<u>A078526</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078526</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078526</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A090484</u>	<u>001</u>	Aug 05, 2010
<u>AB</u>	OXFORD PHARMS	<u>250MG</u>	<u>A077319</u>	<u>001</u>	Mar 20, 2009
<u>AB</u>		<u>500MG</u>	<u>A077319</u>	<u>002</u>	Mar 20, 2009
<u>AB</u>		<u>750MG</u>	<u>A077319</u>	<u>003</u>	Mar 20, 2009
<u>AB</u>	PRINSTON INC	<u>250MG</u>	<u>A078106</u>	<u>001</u>	Feb 10, 2009
<u>AB</u>		<u>500MG</u>	<u>A078106</u>	<u>002</u>	Feb 10, 2009
<u>AB</u>		<u>750MG</u>	<u>A078106</u>	<u>003</u>	Feb 10, 2009
<u>AB</u>		<u>1GM</u>	<u>A078106</u>	<u>004</u>	Feb 10, 2009
<u>AB</u>	SCIEGEN PHARMS INC	<u>750MG</u>	<u>A215069</u>	<u>001</u>	Jun 11, 2021
<u>AB</u>		<u>1GM</u>	<u>A215069</u>	<u>002</u>	Jun 11, 2021
<u>AB</u>	SECAN PHARMS	<u>500MG</u>	<u>A205102</u>	<u>004</u>	Dec 16, 2015
<u>AB</u>		<u>1GM</u>	<u>A205102</u>	<u>003</u>	Dec 16, 2015
<u>AB</u>	TARO	<u>250MG</u>	<u>A078960</u>	<u>004</u>	Feb 01, 2010
<u>AB</u>		<u>500MG</u>	<u>A078960</u>	<u>003</u>	Feb 01, 2010
<u>AB</u>		<u>750MG</u>	<u>A078960</u>	<u>002</u>	Feb 01, 2010
<u>AB</u>		<u>1GM</u>	<u>A078960</u>	<u>001</u>	Feb 01, 2010
<u>AB</u>	TORRENT PHARMS	<u>250MG</u>	<u>A078858</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078858</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078858</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A078858</u>	<u>004</u>	Jan 15, 2009
<u>AB</u>	VIWIT PHARM	<u>250MG</u>	<u>A078869</u>	<u>001</u>	Mar 13, 2009
<u>AB</u>		<u>500MG</u>	<u>A078869</u>	<u>002</u>	Mar 13, 2009
<u>AB</u>		<u>750MG</u>	<u>A078869</u>	<u>003</u>	Mar 13, 2009
<u>AB</u>		<u>1GM</u>	<u>A078869</u>	<u>004</u>	Mar 13, 2009
<u>AB</u>	VKT PHARMA	<u>250MG</u>	<u>A090767</u>	<u>001</u>	Jul 28, 2010
<u>AB</u>		<u>500MG</u>	<u>A090767</u>	<u>002</u>	Jul 28, 2010
<u>AB</u>		<u>750MG</u>	<u>A090767</u>	<u>003</u>	Jul 28, 2010
<u>AB</u>		<u>1GM</u>	<u>A090767</u>	<u>004</u>	Jul 28, 2010
<u>AB</u>	WOCKHARDT	<u>250MG</u>	<u>A079042</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A079042</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A079042</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A079042</u>	<u>004</u>	Jan 15, 2009
<u>AB</u>	ZHEJIANG JINGXIN	<u>250MG</u>	<u>A091491</u>	<u>001</u>	Dec 14, 2010
<u>AB</u>		<u>500MG</u>	<u>A091491</u>	<u>002</u>	Dec 14, 2010

PRESCRIPTION DRUG PRODUCT LIST

LEVETIRACETAM

TABLET; ORAL

LEVETIRACETAM

<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>	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

TABLET, EXTENDED RELEASE; ORAL

KEPPRA XR

<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

LEVETIRACETAM

<u>AB</u>	ACTAVIS LABS FL INC	<u>500MG</u>	<u>A091093 001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091093 002</u>	Sep 12, 2011
<u>AB</u>	ANCHEN PHARMS	<u>500MG</u>	<u>A091360 001</u>	Oct 04, 2011
<u>AB</u>		<u>750MG</u>	<u>A091360 002</u>	Oct 04, 2011
<u>AB</u>	ANDA REPOSITORY	<u>500MG</u>	<u>A204511 001</u>	Feb 23, 2016
<u>AB</u>		<u>750MG</u>	<u>A204511 002</u>	Feb 23, 2016
<u>AB</u>	APOTEX INC	<u>500MG</u>	<u>A091261 001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091261 002</u>	Sep 12, 2011
<u>AB</u>	FLORIDA	<u>500MG</u>	<u>A202167 001</u>	Sep 04, 2015
<u>AB</u>		<u>750MG</u>	<u>A202167 002</u>	Sep 04, 2015
<u>AB</u>	GENUS LIFESCIENCES	<u>500MG</u>	<u>A204754 001</u>	Aug 26, 2016
<u>AB</u>		<u>750MG</u>	<u>A204754 002</u>	Aug 26, 2016
<u>AB</u>	HISUN PHARM HANGZHOU	<u>500MG</u>	<u>A207175 001</u>	Sep 28, 2017
<u>AB</u>		<u>750MG</u>	<u>A207175 002</u>	Sep 28, 2017
<u>AB</u>	LUPIN LTD	<u>500MG</u>	<u>A091399 001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091399 002</u>	Sep 12, 2011
<u>AB</u>	OVERSEAS	<u>750MG</u>	<u>A212688 001</u>	Jun 11, 2020
<u>AB</u>	PHARMADAX INC	<u>500MG</u>	<u>A201464 001</u>	May 25, 2012
<u>AB</u>		<u>750MG</u>	<u>A201464 002</u>	May 25, 2012
<u>AB</u>	PRINSTON INC	<u>500MG</u>	<u>A202533 001</u>	Jul 20, 2012
<u>AB</u>		<u>500MG</u>	<u>A203468 001</u>	May 21, 2015
<u>AB</u>		<u>750MG</u>	<u>A202533 002</u>	Jul 20, 2012
<u>AB</u>		<u>750MG</u>	<u>A203468 002</u>	May 21, 2015
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A205130 001</u>	Nov 27, 2020
<u>AB</u>		<u>750MG</u>	<u>A205130 002</u>	Nov 27, 2020
<u>AB</u>	SUN PHARM	<u>500MG</u>	<u>A203059 001</u>	Sep 09, 2013
<u>AB</u>		<u>750MG</u>	<u>A203059 002</u>	Sep 09, 2013

ELEPSIA XR

	+ TRIPOINT	1GM	N204417 001	Dec 20, 2018
	+!	1.5GM	N204417 002	Dec 20, 2018

LEVETIRACETAM

	APOTEX	1GM	A202958 001	Feb 25, 2015
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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

BETAGAN

<u>AT</u>	+! ALLERGAN	<u>0.5%</u>	<u>N019219 002</u>	Dec 19, 1985
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LEVOBUNOLOL HYDROCHLORIDE

<u>AT</u>	BAUSCH AND LOMB	<u>0.5%</u>	<u>A074326 001</u>	Mar 04, 1994
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LEVOCARNITINE

INJECTABLE; INJECTION

CARNITOR

<u>AP</u>	+! LEADIANT BIOSCI INC	<u>200MG/ML</u>	<u>N020182 001</u>	Dec 16, 1992
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LEVOCARNITINE

<u>AP</u>	AM REGENT	<u>200MG/ML</u>	<u>A075861 001</u>	Jun 22, 2001
<u>AP</u>	HIKMA	<u>200MG/ML</u>	<u>A075567 001</u>	Mar 29, 2001

SOLUTION; ORAL

CARNITOR

<u>AA</u>	+! LEADIANT BIOSCI INC	<u>1GM/10ML</u>	<u>N019257 001</u>	Apr 10, 1986
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CARNITOR SF

<u>AA</u>	+ LEADIANT BIOSCI INC	<u>1GM/10ML</u>	<u>N019257 002</u>	Mar 28, 2007
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PRESCRIPTION DRUG PRODUCT LIST

LEVOCARNITINE

SOLUTION;ORAL

LEVOCARNITINE

<u>AA</u>	AKORN	<u>1GM/10ML</u>	<u>A077399 001</u>	Oct 25, 2007
<u>AA</u>	LYNE	<u>1GM/10ML</u>	<u>A076851 001</u>	Aug 10, 2004
<u>AA</u>	NOVITIUM PHARMA	<u>1GM/10ML</u>	<u>A211676 001</u>	Aug 14, 2019
<u>AA</u>	SAPTALIS PHARMS	<u>1GM/10ML</u>	<u>A212533 001</u>	Nov 10, 2021

LEVOCARNITINE SF

<u>AA</u>	NOVITIUM PHARMA	<u>1GM/10ML</u>	<u>A211676 002</u>	Aug 14, 2019
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TABLET;ORAL

CARNITOR

<u>AB</u>	+! LEADIANT BIOSCI INC	<u>330MG</u>	<u>N018948 001</u>	Dec 27, 1985
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LEVOCARNITINE

<u>AB</u>	RISING	<u>330MG</u>	<u>A076858 001</u>	Sep 20, 2004
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LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION;ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

<u>AA</u>	LANNETT CO INC	<u>2.5MG/5ML</u>	<u>A204599 001</u>	May 15, 2017
<u>AA</u>	! PADAGIS US	<u>2.5MG/5ML</u>	<u>A091263 001</u>	Nov 07, 2011
<u>AA</u>	TARO PHARM INDS LTD	<u>2.5MG/5ML</u>	<u>A202673 001</u>	Jul 26, 2013

LEVOCETIRIZINE HYDROCHLORIDE

<u>AA</u>	HETERO LABS LTD III	<u>2.5MG/5ML</u>	<u>A210914 001</u>	Apr 01, 2019
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TABLET;ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A090392 001</u>	Feb 24, 2011
<u>AB</u>	! GLENMARK GENERICS	<u>5MG</u>	<u>A090385 001</u>	Feb 24, 2011
<u>AB</u>	HETERO LABS LTD III	<u>5MG</u>	<u>A091264 001</u>	Jun 29, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A205564 001</u>	Jan 11, 2016
<u>AB</u>	MICRO LABS LTD	<u>5MG</u>	<u>A202046 001</u>	Sep 17, 2013
	INDIA			
<u>AB</u>	SCIEGEN PHARMS INC	<u>5MG</u>	<u>A203646 001</u>	Sep 09, 2014
<u>AB</u>	SUN PHARM	<u>5MG</u>	<u>A090362 001</u>	Jan 31, 2013
<u>AB</u>	SYNTHON PHARMS	<u>5MG</u>	<u>A090229 001</u>	Nov 26, 2010
<u>AB</u>	TEVA PHARMS	<u>5MG</u>	<u>A090199 001</u>	Aug 22, 2011

LEVODOPA

POWDER; INHALATION

INBRIJA

	+! ACORDA	42MG	N209184 001	Dec 21, 2018
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LEVOFLOXACIN

INJECTABLE; INJECTION

LEVOFLOXACIN

<u>AP</u>	AKORN	<u>EQ 500MG/20ML (EQ 25MG/ML)</u>	<u>A091644 001</u>	Jun 20, 2011
<u>AP</u>		<u>EQ 750MG/30ML (EQ 25MG/ML)</u>	<u>A091644 002</u>	Jun 20, 2011
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 500MG/20ML (EQ 25MG/ML)</u>	<u>A091436 001</u>	Jun 05, 2013
<u>AP</u>	! EUGIA PHARMA	<u>EQ 500MG/20ML (EQ 25MG/ML)</u>	<u>A202328 001</u>	Jan 24, 2013
<u>AP</u>	! EUGIA PHARMA	<u>EQ 750MG/30ML (EQ 25MG/ML)</u>	<u>A202328 002</u>	Jan 24, 2013
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 500MG/20ML (EQ 25MG/ML)</u>	<u>A205540 001</u>	Apr 22, 2020
<u>AP</u>		<u>EQ 750MG/30ML (EQ 25MG/ML)</u>	<u>A205540 002</u>	Apr 22, 2020
	<u>LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A091397 001</u>	Aug 08, 2013
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A091397 002</u>	Aug 08, 2013
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A091397 003</u>	Aug 08, 2013
<u>AP</u>	EUGIA PHARMA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A206919 001</u>	Feb 10, 2016
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A206919 002</u>	Feb 10, 2016
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A206919 003</u>	Feb 10, 2016
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A200674 001</u>	Jun 19, 2013
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A200674 002</u>	Jun 19, 2013
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A200674 003</u>	Jun 19, 2013
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A206908 001</u>	Dec 30, 2020
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A206908 002</u>	Dec 30, 2020
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A206908 003</u>	Dec 30, 2020
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A091375 001</u>	Sep 16, 2011
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A091375 002</u>	Sep 16, 2011
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A091375 003</u>	Sep 16, 2011
<u>AP</u>	HOSPIRA INC	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A078579 001</u>	Sep 03, 2015
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A078579 002</u>	Sep 03, 2015
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A078579 003</u>	Sep 03, 2015
<u>AP</u>	! INFORLIFE	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A090343 001</u>	Jul 07, 2011
<u>AP</u>	! INFORLIFE	<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A090343 002</u>	Jul 07, 2011

PRESCRIPTION DRUG PRODUCT LIST

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	!	<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A090343</u>	<u>003</u>	Jul 07, 2011
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SOLUTION; ORAL

LEVOFLOXACIN

<u>AA</u>	!	AKORN	<u>250MG/10ML</u>	<u>A091678</u>	<u>001</u>	Jun 20, 2011
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<u>AA</u>		LANNETT CO INC	<u>250MG/10ML</u>	<u>A205222</u>	<u>001</u>	May 25, 2018
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SOLUTION/DROPS; OPHTHALMIC

LEVOFLOXACIN

<u>AT</u>		AKORN	<u>0.5%</u>	<u>A090268</u>	<u>001</u>	Dec 20, 2010
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<u>AT</u>	!	RISING	<u>0.5%</u>	<u>A077700</u>	<u>001</u>	Dec 20, 2010
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		MICRO LABS LTD	1.5%	A205600	001	Feb 27, 2019
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INDIA

TABLET; ORAL

LEVOFLOXACIN

<u>AB</u>		APOTEX INC	<u>250MG</u>	<u>A090787</u>	<u>001</u>	Sep 29, 2011
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<u>AB</u>			<u>500MG</u>	<u>A090787</u>	<u>002</u>	Sep 29, 2011
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<u>AB</u>			<u>750MG</u>	<u>A090787</u>	<u>003</u>	Sep 29, 2011
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<u>AB</u>		AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A201043</u>	<u>001</u>	Jun 20, 2011
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<u>AB</u>			<u>500MG</u>	<u>A201043</u>	<u>002</u>	Jun 20, 2011
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<u>AB</u>	!		<u>750MG</u>	<u>A201043</u>	<u>003</u>	Jun 20, 2011
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<u>AB</u>		CADILA	<u>250MG</u>	<u>A077652</u>	<u>001</u>	Sep 07, 2012
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<u>AB</u>			<u>500MG</u>	<u>A077652</u>	<u>002</u>	Sep 07, 2012
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<u>AB</u>			<u>750MG</u>	<u>A077652</u>	<u>003</u>	Sep 07, 2012
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<u>AB</u>		CELLTRION	<u>250MG</u>	<u>A090367</u>	<u>001</u>	Jun 20, 2011
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<u>AB</u>			<u>500MG</u>	<u>A090367</u>	<u>002</u>	Jun 20, 2011
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<u>AB</u>			<u>750MG</u>	<u>A090367</u>	<u>003</u>	Jun 20, 2011
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<u>AB</u>		CIPLA LTD	<u>250MG</u>	<u>A076890</u>	<u>001</u>	Mar 30, 2012
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<u>AB</u>			<u>500MG</u>	<u>A076890</u>	<u>002</u>	Mar 30, 2012
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<u>AB</u>			<u>750MG</u>	<u>A076890</u>	<u>003</u>	Mar 30, 2012
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<u>AB</u>		DR REDDYS LABS INC	<u>250MG</u>	<u>A076710</u>	<u>001</u>	Jun 20, 2011
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<u>AB</u>			<u>500MG</u>	<u>A076710</u>	<u>002</u>	Jun 20, 2011
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<u>AB</u>			<u>750MG</u>	<u>A076710</u>	<u>003</u>	Jun 20, 2011
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<u>AB</u>		GLENMARK GENERICS	<u>250MG</u>	<u>A200250</u>	<u>001</u>	Jun 20, 2011
-----------	--	-------------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>A200250</u>	<u>002</u>	Jun 20, 2011
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<u>AB</u>			<u>750MG</u>	<u>A200250</u>	<u>003</u>	Jun 20, 2011
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<u>AB</u>		HEC PHARM	<u>250MG</u>	<u>A204968</u>	<u>001</u>	Feb 05, 2019
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<u>AB</u>			<u>500MG</u>	<u>A204968</u>	<u>002</u>	Feb 05, 2019
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<u>AB</u>			<u>750MG</u>	<u>A204968</u>	<u>003</u>	Feb 05, 2019
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<u>AB</u>		HETERO LABS LTD V	<u>250MG</u>	<u>A202801</u>	<u>001</u>	Jan 08, 2015
-----------	--	-------------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>A202801</u>	<u>002</u>	Jan 08, 2015
-----------	--	--	--------------	----------------	------------	--------------

<u>AB</u>			<u>750MG</u>	<u>A202801</u>	<u>003</u>	Jan 08, 2015
-----------	--	--	--------------	----------------	------------	--------------

<u>AB</u>		LUPIN	<u>250MG</u>	<u>A078424</u>	<u>001</u>	Jun 20, 2011
-----------	--	-------	--------------	----------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>A078424</u>	<u>002</u>	Jun 20, 2011
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<u>AB</u>			<u>750MG</u>	<u>A078424</u>	<u>003</u>	Jun 20, 2011
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<u>AB</u>		MACLEODS PHARMS LTD	<u>250MG</u>	<u>A200839</u>	<u>001</u>	Mar 22, 2012
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<u>AB</u>			<u>500MG</u>	<u>A200839</u>	<u>002</u>	Mar 22, 2012
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<u>AB</u>			<u>750MG</u>	<u>A200839</u>	<u>003</u>	Mar 22, 2012
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<u>AB</u>		ORBION PHARMS	<u>250MG</u>	<u>A202200</u>	<u>001</u>	Jan 30, 2012
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<u>AB</u>			<u>500MG</u>	<u>A202200</u>	<u>002</u>	Jan 30, 2012
-----------	--	--	--------------	----------------	------------	--------------

<u>AB</u>			<u>750MG</u>	<u>A202200</u>	<u>003</u>	Jan 30, 2012
-----------	--	--	--------------	----------------	------------	--------------

<u>AB</u>		SANDOZ	<u>250MG</u>	<u>A077438</u>	<u>001</u>	Jun 20, 2011
-----------	--	--------	--------------	----------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>A077438</u>	<u>002</u>	Jun 20, 2011
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<u>AB</u>			<u>750MG</u>	<u>A077438</u>	<u>003</u>	Jun 20, 2011
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<u>AB</u>		TEVA	<u>250MG</u>	<u>A076361</u>	<u>001</u>	Jun 20, 2011
-----------	--	------	--------------	----------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>A076361</u>	<u>002</u>	Jun 20, 2011
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<u>AB</u>			<u>750MG</u>	<u>A076361</u>	<u>003</u>	Jun 20, 2011
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LEVOKETOCONAZOLE

TABLET; ORAL

RECORLEV

+	!	STRONGBRIDGE	150MG	N214133	001	Dec 30, 2021
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LEVOLEUCOVORIN

POWDER; INTRAVENOUS

KHAPZORY

+	!	ACROTECH	175MG/VIAL	N211226	001	Oct 19, 2018
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+	!		300MG/VIAL	N211226	002	Oct 19, 2018
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PRESCRIPTION DRUG PRODUCT LIST

LEVOLEUCOVORIN CALCIUM

POWDER; INTRAVENOUS

FUSILEV

AP	+!	ACROTECH	<u>EQ 50MG BASE/VIAL</u>	<u>N020140 001</u>	Mar 07, 2008
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LEVOLEUCOVORIN CALCIUM

AP		HIKMA	<u>EQ 50MG BASE/VIAL</u>	<u>A206263 001</u>	Jun 16, 2016
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AP		MEITHEAL	<u>EQ 50MG BASE/VIAL</u>	<u>A211003 001</u>	Aug 22, 2019
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SOLUTION; INTRAVENOUS

LEVOLEUCOVORIN CALCIUM

AP		AMNEAL	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A207548 001</u>	Sep 08, 2017
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AP	!	GLAND PHARMA LTD	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A210892 001</u>	Sep 14, 2018
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AP	!		<u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u>	<u>A210892 002</u>	Sep 14, 2018
-----------	----------	--	--	---------------------------	--------------

AP		MEITHEAL	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A211002 001</u>	Aug 16, 2019
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AP			<u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u>	<u>A211002 002</u>	Aug 16, 2019
-----------	--	--	--	---------------------------	--------------

AP		PRAXGEN	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A213797 001</u>	Nov 02, 2021
-----------	--	---------	--	---------------------------	--------------

AP		SANDOZ INC	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A203563 001</u>	Mar 09, 2015
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LEVOMILNACIPRAN HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

FETZIMA

+		ALLERGAN	EQ 20MG BASE	N204168 001	Jul 25, 2013
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+			EQ 40MG BASE	N204168 002	Jul 25, 2013
----------	--	--	--------------	-------------	--------------

+			EQ 80MG BASE	N204168 003	Jul 25, 2013
----------	--	--	--------------	-------------	--------------

+!			EQ 120MG BASE	N204168 004	Jul 25, 2013
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LEVONORDEFIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

SCANDONEST L

!		DEPROCO	0.05MG/ML; 2%	A088388 001	Oct 10, 1984
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LEVONORGESTREL

INTRAUTERINE DEVICE; INTRAUTERINE

KYLEENA

+!		BAYER HLTHCARE	19.5MG	N208224 001	Sep 16, 2016
-----------	--	----------------	--------	-------------	--------------

LILETTA

+!		MEDICINES360	52MG	N206229 001	Feb 26, 2015
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MIRENA

+!		BAYER HLTHCARE	52MG	N021225 001	Dec 06, 2000
-----------	--	----------------	------	-------------	--------------

SKYLA

+!		BAYER HLTHCARE	13.5MG	N203159 001	Jan 09, 2013
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TABLET; ORAL

LEVONORGESTREL

!		L PERRIGO CO	0.75MG	A090740 001	Dec 30, 2010
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LEVORPHANOL TARTRATE

TABLET; ORAL

LEVORPHANOL TARTRATE

AB		NOVITIUM PHARMA	<u>2MG</u>	<u>A213479 001</u>	Jul 01, 2020
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AB			<u>3MG</u>	<u>A213479 002</u>	Jan 12, 2021
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AB	!	SENTYNL THERAPS INC	<u>2MG</u>	<u>A074278 001</u>	Mar 31, 2000
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AB	!		<u>3MG</u>	<u>A074278 003</u>	Jun 18, 2018
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AB		SPECGX LLC	<u>2MG</u>	<u>A212024 001</u>	Dec 13, 2019
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AB		SUN PHARM INDS INC	<u>2MG</u>	<u>A213906 001</u>	Jun 17, 2021
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AB		VIRTUS PHARMS	<u>2MG</u>	<u>A211484 001</u>	Dec 13, 2018
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		NOVITIUM PHARMA	1MG	A213479 003	Jul 20, 2021
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LEVOTHYROXINE SODIUM

CAPSULE; ORAL

LEVOTHYROXINE SODIUM

AB		TEVA PHARMS USA INC	<u>0.112MG</u>	<u>A211369 003</u>	Apr 16, 2021
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TIROSINT

AB	+	INSTITUT	<u>0.112MG</u>	<u>N021924 008</u>	Oct 02, 2009
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		BIOCHIMIQUE			
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+			0.013MG	N021924 013	Aug 01, 2007
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+			0.025MG	N021924 002	Oct 13, 2006
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+			0.05MG	N021924 003	Oct 13, 2006
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+			0.075MG	N021924 004	Oct 13, 2006
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+			0.088MG	N021924 010	Oct 02, 2009
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+			0.1MG	N021924 005	Oct 13, 2006
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+			0.125MG	N021924 006	Oct 13, 2006
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+			0.137MG	N021924 009	Oct 02, 2009
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+			0.15MG	N021924 007	Oct 13, 2006
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+			0.175MG	N021924 011	Apr 25, 2017
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+!			0.200MG	N021924 012	Apr 25, 2017
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PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM

POWDER; INTRAVENOUS

LEVOTHYROXINE SODIUM

AP	+ !	FRESENIUS KABI USA	100MCG/VIAL	N202231 001	Jun 24, 2011
AP	+ !		200MCG/VIAL	N202231 002	Jun 24, 2011
AP	+ !		500MCG/VIAL	N202231 003	Jun 24, 2011
AP		MAIA PHARMS INC	100MCG/VIAL	A208749 001	Dec 21, 2018
AP			200MCG/VIAL	A208749 002	Dec 21, 2018
AP			500MCG/VIAL	A208749 003	Dec 21, 2018
AP		PIRAMAL CRITICAL	100MCG/VIAL	A206163 001	Jun 29, 2016
AP			500MCG/VIAL	A206163 002	Jun 29, 2016

SOLUTION; INTRAVENOUS

LEVOTHYROXINE SODIUM

+ !	CUSTOPHARM INC	100MCG/ML	N214253 001	May 17, 2021
+ !	FRESENIUS KABI USA	100MCG/5ML (20MCG/ML)	N210632 001	Apr 11, 2019
+ !		200MCG/5ML (40MCG/ML)	N210632 002	Apr 11, 2019
+ !		500MCG/5ML (100MCG/ML)	N210632 003	Apr 11, 2019

SOLUTION; ORAL

THYQUIDITY

+ !	VISTAPHARM	100MCG/5ML	N214047 001	Nov 30, 2020
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TIROSINT-SOL

+	INSTITUT BIOCHIMIQUE	13MCG/ML	N206977 001	Dec 15, 2016
+		25MCG/ML	N206977 002	Dec 15, 2016
+		37.5MCG/ML	N206977 013	Jan 13, 2021
+		44MCG/ML	N206977 014	Jan 13, 2021
+		50MCG/ML	N206977 003	Dec 15, 2016
+		62.5MCG/ML	N206977 015	Jan 13, 2021
+		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

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET; ORAL

SYNTHROID

-->	+	ABBVIE	--> AB1, AB2	0.025MG	N021402 001	Jul 24, 2002
-->	+		--> AB1, AB2	0.05MG	N021402 002	Jul 24, 2002
-->	+		--> AB1, AB2	0.075MG	N021402 003	Jul 24, 2002
-->	+		--> AB1, AB2	0.088MG	N021402 004	Jul 24, 2002
-->	+		--> AB1, AB2	0.1MG	N021402 005	Jul 24, 2002
-->	+		--> AB1, AB2	0.112MG	N021402 006	Jul 24, 2002
-->	+		--> AB1, AB2	0.125MG	N021402 007	Jul 24, 2002
-->	+		--> AB1, AB2	0.137MG	N021402 008	Jul 24, 2002
-->	+		--> AB1, AB2	0.15MG	N021402 009	Jul 24, 2002
-->	+		--> AB1, AB2	0.175MG	N021402 010	Jul 24, 2002
-->	+		--> AB1, AB2	0.2MG	N021402 012	Jul 24, 2002
-->	+ !		--> AB1, AB2	0.3MG	N021402 011	Jul 24, 2002

LEVO-T

-->		CEDIPROF INC	--> AB1, AB2, AB3	0.025MG	N021342 001	Mar 01, 2002
-->			--> AB1, AB2, AB3	0.05MG	N021342 002	Mar 01, 2002
-->			--> AB1, AB2, AB3	0.075MG	N021342 003	Mar 01, 2002
-->			--> AB1, AB2, AB3	0.088MG	N021342 004	Mar 01, 2002
-->			--> AB1, AB2, AB3	0.1MG	N021342 005	Mar 01, 2002
-->			--> AB1, AB2, AB3	0.112MG	N021342 006	Mar 01, 2002
-->			--> AB1, AB2, AB3	0.125MG	N021342 007	Mar 01, 2002
-->			--> AB1, AB2, AB3	0.137MG	N021342 012	Dec 08, 2003
-->			--> AB1, AB2, AB3	0.15MG	N021342 008	Mar 01, 2002
-->			--> AB1, AB2, AB3	0.175MG	N021342 009	Mar 01, 2002

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

LEVO-T--> --> AB1,AB2,AB3 0.2MG N021342 010 Mar 01, 2002--> --> AB1,AB2,AB3 0.3MG N021342 011 Mar 01, 2002LEVOTHYROXINE SODIUM--> LUPIN --> AB1,AB2,AB3 0.025MG A209713 001 Jan 18, 2019--> --> AB1,AB2,AB3 0.05MG A209713 002 Jan 18, 2019--> --> AB1,AB2,AB3 0.075MG A209713 003 Jan 18, 2019--> --> AB1,AB2,AB3 0.088MG A209713 004 Jan 18, 2019--> --> AB1,AB2,AB3 0.1MG A209713 005 Jan 18, 2019--> --> AB1,AB2,AB3 0.112MG A209713 006 Jan 18, 2019--> --> AB1,AB2,AB3 0.125MG A209713 007 Jan 18, 2019--> --> AB1,AB2,AB3 0.137MG A209713 008 Jan 18, 2019--> --> AB1,AB2,AB3 0.15MG A209713 009 Jan 18, 2019--> --> AB1,AB2,AB3 0.175MG A209713 010 Jan 18, 2019--> --> AB1,AB2,AB3 0.2MG A209713 011 Jan 18, 2019--> --> AB1,AB2,AB3 0.3MG A209713 012 Jan 18, 2019UNITHROID--> + STEVENS J --> AB1,AB2,AB3 0.025MG N021210 001 Aug 21, 2000--> + --> AB1,AB2,AB3 0.05MG N021210 002 Aug 21, 2000--> + --> AB1,AB2,AB3 0.075MG N021210 003 Aug 21, 2000--> + --> AB1,AB2,AB3 0.088MG N021210 004 Aug 21, 2000--> + --> AB1,AB2,AB3 0.1MG N021210 005 Aug 21, 2000--> + --> AB1,AB2,AB3 0.112MG N021210 006 Aug 21, 2000--> + --> AB1,AB2,AB3 0.125MG N021210 007 Aug 21, 2000--> + --> AB1,AB2,AB3 0.137MG N021210 012 Feb 08, 2008--> + --> AB1,AB2,AB3 0.15MG N021210 008 Aug 21, 2000--> + --> AB1,AB2,AB3 0.175MG N021210 009 Aug 21, 2000--> + --> AB1,AB2,AB3 0.2MG N021210 010 Aug 21, 2000--> +! --> AB1,AB2,AB3 0.3MG N021210 011 Aug 21, 2000LEVOTHYROXINE 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, 0.2MG A076187 010 Jun 05, 2002

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

LEVOTHYROXINE SODIUM

		<u>AB3, AB4</u>	<u>0.2MG</u>				
-->	!	-->	<u>0.3MG</u>	A076187	011	Jun 05,	2002
		<u>AB1, AB2, AB3, AB4</u>					
<u>THYRO-TABS</u>							
-->	+	ALVOGEN	--> <u>AB1, AB2, AB4</u>	<u>0.025MG</u>	N021116	001	Oct 24, 2002
-->	+		--> <u>AB1, AB2, AB4</u>	<u>0.05MG</u>	N021116	002	Oct 24, 2002
-->	+		--> <u>AB1, AB2, AB4</u>	<u>0.075MG</u>	N021116	003	Oct 24, 2002
-->	+		--> <u>AB1, AB2, AB4</u>	<u>0.088MG</u>	N021116	010	Oct 24, 2002
-->	+		--> <u>AB1, AB2, AB4</u>	<u>0.1MG</u>	N021116	004	Oct 24, 2002
-->	+		--> <u>AB1, AB2, AB4</u>	<u>0.112MG</u>	N021116	011	Oct 24, 2002
-->	+		--> <u>AB1, AB2, AB4</u>	<u>0.125MG</u>	N021116	005	Oct 24, 2002
-->	+		--> <u>AB1, AB2, AB4</u>	<u>0.137MG</u>	N021116	012	Dec 07, 2004
-->	+		--> <u>AB1, AB2, AB4</u>	<u>0.15MG</u>	N021116	006	Oct 24, 2002
-->	+		--> <u>AB1, AB2, AB4</u>	<u>0.175MG</u>	N021116	007	Oct 24, 2002
-->	+		--> <u>AB1, AB2, AB4</u>	<u>0.2MG</u>	N021116	008	Oct 24, 2002
-->	+		--> <u>AB1, AB2, AB4</u>	<u>0.3MG</u>	N021116	009	Oct 24, 2002

LEVOXYL

-->	+	KING PHARMS	--> <u>AB1, AB3</u>	<u>0.025MG</u>	N021301	001	May 25, 2001
-->	+		--> <u>AB1, AB3</u>	<u>0.05MG</u>	N021301	002	May 25, 2001
-->	+		--> <u>AB1, AB3</u>	<u>0.075MG</u>	N021301	003	May 25, 2001
-->	+		--> <u>AB1, AB3</u>	<u>0.088MG</u>	N021301	004	May 25, 2001
-->	+		--> <u>AB1, AB3</u>	<u>0.1MG</u>	N021301	005	May 25, 2001
-->	+		--> <u>AB1, AB3</u>	<u>0.112MG</u>	N021301	006	May 25, 2001
-->	+		--> <u>AB1, AB3</u>	<u>0.125MG</u>	N021301	007	May 25, 2001
-->	+		--> <u>AB1, AB3</u>	<u>0.137MG</u>	N021301	008	May 25, 2001
-->	+		--> <u>AB1, AB3</u>	<u>0.15MG</u>	N021301	009	May 25, 2001
-->	+		--> <u>AB1, AB3</u>	<u>0.175MG</u>	N021301	010	May 25, 2001
-->	+		--> <u>AB1, AB3</u>	<u>0.2MG</u>	N021301	011	May 25, 2001

EUTHYROX

<u>AB2</u>	PROVELL	<u>0.025MG</u>	<u>N021292</u>	<u>001</u>	May 31,	2002
<u>AB2</u>		<u>0.05MG</u>	<u>N021292</u>	<u>002</u>	May 31,	2002
<u>AB2</u>		<u>0.075MG</u>	<u>N021292</u>	<u>003</u>	May 31,	2002
<u>AB2</u>		<u>0.088MG</u>	<u>N021292</u>	<u>004</u>	May 31,	2002
<u>AB2</u>		<u>0.1MG</u>	<u>N021292</u>	<u>005</u>	May 31,	2002
<u>AB2</u>		<u>0.112MG</u>	<u>N021292</u>	<u>006</u>	May 31,	2002
<u>AB2</u>		<u>0.125MG</u>	<u>N021292</u>	<u>007</u>	May 31,	2002
<u>AB2</u>		<u>0.137MG</u>	<u>N021292</u>	<u>008</u>	May 31,	2002
<u>AB2</u>		<u>0.15MG</u>	<u>N021292</u>	<u>009</u>	May 31,	2002
<u>AB2</u>		<u>0.175MG</u>	<u>N021292</u>	<u>010</u>	May 31,	2002
<u>AB2</u>		<u>0.2MG</u>	<u>N021292</u>	<u>011</u>	May 31,	2002

LEVOTHYROXINE SODIUM

<u>AB2</u>	ACCORD HLTHCARE	<u>0.025MG</u>	<u>A212399</u>	<u>001</u>	Oct 19,	2020
<u>AB2</u>		<u>0.05MG</u>	<u>A212399</u>	<u>002</u>	Oct 19,	2020
<u>AB2</u>		<u>0.075MG</u>	<u>A212399</u>	<u>003</u>	Oct 19,	2020
<u>AB2</u>		<u>0.088MG</u>	<u>A212399</u>	<u>004</u>	Oct 19,	2020
<u>AB2</u>		<u>0.1MG</u>	<u>A212399</u>	<u>005</u>	Oct 19,	2020
<u>AB2</u>		<u>0.112MG</u>	<u>A212399</u>	<u>006</u>	Oct 19,	2020
<u>AB2</u>		<u>0.125MG</u>	<u>A212399</u>	<u>007</u>	Oct 19,	2020
<u>AB2</u>		<u>0.137MG</u>	<u>A212399</u>	<u>008</u>	Oct 19,	2020
<u>AB2</u>		<u>0.15MG</u>	<u>A212399</u>	<u>009</u>	Oct 19,	2020
<u>AB2</u>		<u>0.175MG</u>	<u>A212399</u>	<u>010</u>	Oct 19,	2020
<u>AB2</u>		<u>0.2MG</u>	<u>A212399</u>	<u>011</u>	Oct 19,	2020
<u>AB2</u>		<u>0.3MG</u>	<u>A212399</u>	<u>012</u>	Oct 19,	2020

PRESCRIPTION DRUG PRODUCT LIST

LIDOCAINE

OINTMENT; TOPICAL

LIDOCAINE

<u>AT</u>	ALEOR	<u>5%</u>	<u>A211469</u>	<u>001</u>	Nov 23, 2018
	DERMACEUTICALS				
<u>AT</u>	ALKEM LABS LTD	<u>5%</u>	<u>A207810</u>	<u>001</u>	Mar 10, 2017
<u>AT</u>	AMNEAL PHARMS	<u>5%</u>	<u>A206297</u>	<u>001</u>	Aug 07, 2015
<u>AT</u>	COSETTE	<u>5%</u>	<u>A211019</u>	<u>001</u>	Dec 12, 2018
<u>AT</u>	DR REDDYS	<u>5%</u>	<u>A208660</u>	<u>001</u>	Jan 05, 2021
<u>AT</u>	+! FOUGERA PHARMS INC	<u>5%</u>	<u>A080198</u>	<u>001</u>	
<u>AT</u>	GLENMARK PHARMS LTD	<u>5%</u>	<u>A206498</u>	<u>001</u>	Sep 09, 2016
<u>AT</u>	MACLEODS PHARMS LTD	<u>5%</u>	<u>A211697</u>	<u>001</u>	Mar 16, 2020
<u>AT</u>	QUAGEN	<u>5%</u>	<u>A212695</u>	<u>001</u>	Apr 20, 2021
<u>AT</u>	SEPTODONT INC	<u>5%</u>	<u>A040911</u>	<u>001</u>	May 23, 2011
<u>AT</u>	STRIDES PHARMA	<u>5%</u>	<u>A210958</u>	<u>001</u>	Dec 11, 2018
<u>AT</u>	TARO	<u>5%</u>	<u>A086724</u>	<u>001</u>	
<u>AT</u>	TELIGENT	<u>5%</u>	<u>A205318</u>	<u>001</u>	Feb 01, 2016
<u>AT</u>	VITRUVIAS THERAP	<u>5%</u>	<u>A208822</u>	<u>001</u>	Sep 25, 2017

PATCH; TOPICAL

LIDOCAINE

<u>AB</u>	ACTAVIS LABS UT INC	<u>5%</u>	<u>A200675</u>	<u>001</u>	Aug 23, 2012
<u>AB</u>	AMNEAL	<u>5%</u>	<u>A206463</u>	<u>001</u>	Aug 24, 2020
<u>AB</u>	MYLAN TECHNOLOGIES	<u>5%</u>	<u>A202346</u>	<u>001</u>	Aug 07, 2015
<u>AB</u>	NAL PHARM	<u>5%</u>	<u>A205882</u>	<u>001</u>	Apr 29, 2021
<u>AB</u>	RHODES PHARMS	<u>5%</u>	<u>A209190</u>	<u>001</u>	Apr 30, 2020

LIDODERM

<u>AB</u>	+! TEIKOKU PHARMA USA	<u>5%</u>	<u>N020612</u>	<u>001</u>	Mar 19, 1999
	ZTLIDO				
	+! SCILEX PHARMS INC	<u>1.8%</u>	<u>N207962</u>	<u>001</u>	Feb 28, 2018

LIDOCAINE HYDROCHLORIDE

GEL; OPHTHALMIC

AKTEN

	+! AKORN	<u>3.5%</u>	<u>N022221</u>	<u>001</u>	Oct 07, 2008
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INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE

<u>AP</u>	ASPIRO	<u>1%</u>	<u>A214336</u>	<u>001</u>	Nov 08, 2021
<u>AP</u>		<u>1%</u>	<u>A214339</u>	<u>001</u>	Nov 08, 2021
<u>AP</u>		<u>2%</u>	<u>A214339</u>	<u>002</u>	Nov 08, 2021
<u>AP</u>	B BRAUN MEDICAL INC	<u>1%</u>	<u>A208474</u>	<u>001</u>	Aug 03, 2018
<u>AP</u>	EUGIA PHARMA	<u>1%</u>	<u>A207182</u>	<u>001</u>	Oct 30, 2017
<u>AP</u>		<u>2%</u>	<u>A207182</u>	<u>002</u>	Oct 30, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A080404</u>	<u>002</u>	
<u>AP</u>		<u>2%</u>	<u>A080404</u>	<u>003</u>	
<u>AP</u>	HOSPIRA	<u>0.5%</u>	<u>A088328</u>	<u>001</u>	May 17, 1984
<u>AP</u>	!	<u>1%</u>	<u>A083158</u>	<u>001</u>	
<u>AP</u>		<u>1%</u>	<u>A088329</u>	<u>001</u>	May 17, 1984
<u>AP</u>		<u>2%</u>	<u>A040078</u>	<u>001</u>	Jun 23, 1995
<u>AP</u>	!	<u>2%</u>	<u>A083158</u>	<u>002</u>	
<u>AP</u>		<u>2%</u>	<u>A088294</u>	<u>001</u>	May 17, 1984
<u>AP</u>	HUONS	<u>1%</u>	<u>A212821</u>	<u>001</u>	May 07, 2020
<u>AP</u>	INTL MEDICATION	<u>1%</u>	<u>A083173</u>	<u>001</u>	
<u>AP</u>		<u>2%</u>	<u>A083173</u>	<u>002</u>	
<u>AP</u>	SPECTRA MDCL	<u>1%</u>	<u>A208017</u>	<u>001</u>	Apr 18, 2018
	DEVICES				
<u>AP</u>	WEST-WARD PHARMS	<u>1%</u>	<u>A080407</u>	<u>001</u>	
	INT				
<u>AP</u>		<u>2%</u>	<u>A080407</u>	<u>002</u>	

LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>200MG/100ML</u>	<u>N019830</u>	<u>002</u>	Apr 08, 1992
<u>AP</u>	BAXTER HLHCARE	<u>200MG/100ML</u>	<u>N018461</u>	<u>002</u>	

LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>400MG/100ML</u>	<u>N019830</u>	<u>003</u>	Apr 08, 1992
<u>AP</u>	BAXTER HLHCARE	<u>400MG/100ML</u>	<u>N018461</u>	<u>003</u>	

LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>800MG/100ML</u>	<u>N019830</u>	<u>004</u>	Apr 08, 1992
<u>AP</u>	BAXTER HLHCARE	<u>800MG/100ML</u>	<u>N018461</u>	<u>004</u>	Feb 22, 1982

LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER

<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A088586</u>	<u>001</u>	Jul 24, 1985
<u>AP</u>	HOSPIRA	<u>0.5%</u>	<u>A088325</u>	<u>001</u>	Jul 31, 1984
<u>AP</u>		<u>1%</u>	<u>A088299</u>	<u>001</u>	Jul 31, 1984
<u>AP</u>		<u>2%</u>	<u>A088327</u>	<u>001</u>	Jul 31, 1984

PRESCRIPTION DRUG PRODUCT LIST

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	EUGIA PHARMA	<u>1%</u>	<u>A203040 001</u>	Mar 14, 2013
<u>AP</u>		<u>1%</u>	<u>A203082 001</u>	Mar 14, 2013
<u>AP</u>		<u>2%</u>	<u>A203040 002</u>	Mar 14, 2013
<u>AP</u>		<u>2%</u>	<u>A203082 002</u>	Mar 14, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>2%</u>	<u>N017584 001</u>	
<u>AP</u>		<u>4%</u>	<u>N017584 002</u>	
<u>AP</u>	HOSPIRA	<u>1%</u>	<u>A080408 001</u>	
<u>AP</u>		<u>1.5%</u>	<u>A080408 002</u>	
<u>AP</u>	WEST-WARD PHARMS INT	<u>1%</u>	<u>A084625 001</u>	

<u>AP</u>		<u>2%</u>	<u>A084625 002</u>	
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LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>1%</u>	<u>A040302 001</u>	Sep 28, 1998
<u>AP</u>		<u>2%</u>	<u>A040302 002</u>	Sep 28, 1998

XYLOCAINE

<u>AP</u>	+! FRESENIUS KABI USA	<u>0.5%</u>	<u>N006488 008</u>	
<u>AP</u>	+!	<u>1%</u>	<u>N006488 007</u>	
<u>AP</u>	+!	<u>1.5%</u>	<u>N006488 010</u>	
<u>AP</u>	+!	<u>2%</u>	<u>N006488 002</u>	

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

<u>AT</u>	SAGENT PHARMS INC	<u>2%</u>	<u>A201094 001</u>	Apr 28, 2014
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LIDOCAINE HYDROCHLORIDE

<u>AT</u>	! AKORN	<u>2%</u>	<u>A040433 001</u>	Feb 12, 2003
<u>AT</u>	INTL MEDICATION	<u>2%</u>	<u>A086283 001</u>	

SOLUTION; ORAL

LIDOCAINE HYDROCHLORIDE

<u>AT</u>	! AKORN	<u>2%</u>	<u>A040014 001</u>	Jul 10, 1995
<u>AT</u>	WOCKHARDT BIO AG	<u>2%</u>	<u>A087872 001</u>	Nov 18, 1982

LIDOCAINE HYDROCHLORIDE VISCOUS

<u>AT</u>	LANNETT CO INC	<u>2%</u>	<u>A040708 001</u>	Feb 27, 2007
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LIDOCAINE VISCOUS

<u>AT</u>	HIKMA	<u>2%</u>	<u>A088802 001</u>	Apr 26, 1985
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SOLUTION; TOPICAL

LARYNG-O-JET KIT

<u>AT</u>	INTL MEDICATION	<u>4%</u>	<u>A086364 001</u>	
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LIDOCAINE HYDROCHLORIDE

<u>AT</u>	! HIKMA	<u>4%</u>	<u>A088803 001</u>	Apr 03, 1985
<u>AT</u>	LANNETT CO INC	<u>4%</u>	<u>A040710 001</u>	Feb 27, 2007
<u>AT</u>	TELGENT	<u>4%</u>	<u>A204494 001</u>	Mar 12, 2014
<u>AT</u>	WOCKHARDT BIO AG	<u>4%</u>	<u>A087881 001</u>	Nov 18, 1982

SYSTEM; INTRADERMAL

ZINGO

POWDER PHARMS 0.5MG N022114 001 Aug 16, 2007

LIDOCAINE; PRILUCAINE

CREAM; TOPICAL

EMLA

<u>AB</u>	+! TEVA BRANDED PHARM	<u>2.5%;2.5%</u>	<u>N019941 001</u>	Dec 30, 1992
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LIDOCAINE AND PRILUCAINE

<u>AB</u>	ACRUX DDS PTY	<u>2.5%;2.5%</u>	<u>A212482 001</u>	Jul 27, 2021
<u>AB</u>	AKORN	<u>2.5%;2.5%</u>	<u>A076290 001</u>	Sep 25, 2003
<u>AB</u>	FOUGERA PHARMS	<u>2.5%;2.5%</u>	<u>A076453 001</u>	Aug 18, 2003
<u>AB</u>	TELGENT	<u>2.5%;2.5%</u>	<u>A205887 001</u>	Jun 29, 2018
<u>AB</u>	TOLMAR	<u>2.5%;2.5%</u>	<u>A076320 001</u>	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

PRESCRIPTION DRUG PRODUCT LIST

LIDOCAINE; TETRACAINE

PATCH; TOPICAL

SYNERA

+! GALEN SPECIALTY 70MG; 70MG N021623 001 Jun 23, 2005

LIFITEGRAST

SOLUTION/DROPS; OPHTHALMIC

XIIDRA

+! NOVARTIS 5% N208073 001 Jul 11, 2016

LINACLOTIDE

CAPSULE; ORAL

LINZESS

+! ALLERGAN 72MCG N202811 003 Jan 25, 2017

+! 145MCG N202811 001 Aug 30, 2012

+ 290MCG N202811 002 Aug 30, 2012

LINAGLIPTIN

TABLET; ORAL

LINAGLIPTIN**AB** SUNSHINE **5MG** **A208335 001** Aug 31, 2021TRADJENTA**AB** +! BOEHRINGER **5MG** **N201280 001** May 02, 2011

INGELHEIM

LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

JENTADUETO**AB** + BOEHRINGER **2.5MG; 500MG** **N201281 001** Jan 30, 2012

INGELHEIM

AB + **2.5MG; 850MG** **N201281 002** Jan 30, 2012**AB** +! **2.5MG; 1GM** **N201281 003** Jan 30, 2012LINAGLIPTIN AND METFORMIN HYDROCHLORIDE**AB** SUNSHINE **2.5MG; 500MG** **A208336 001** Aug 30, 2021**AB** **2.5MG; 850MG** **A208336 002** Aug 30, 2021**AB** **2.5MG; 1GM** **A208336 003** Aug 30, 2021

TABLET, EXTENDED RELEASE; ORAL

JENTADUETO XR

+ BOEHRINGER 2.5MG; 1GM N208026 001 May 27, 2016

INGELHEIM

+! 5MG; 1GM N208026 002 May 27, 2016

LINCOMYCIN

INJECTABLE; INJECTION

LINCOMYCIN**AP** MICRO LABS **EQ 300MG BASE/ML** **A215082 001** Nov 08, 2021LINCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

LINCOCIN**AP** +! PFIZER **EQ 300MG BASE/ML** **N050317 001**LINCOMYCIN**AP** XGEN PHARMS **EQ 300MG BASE/ML** **A201746 001** Jun 04, 2015LINDANE

SHAMPOO; TOPICAL

LINDANE

! WOCKHARDT BIO AG 1% A088191 001 Sep 18, 1984

LINEZOLID

FOR SUSPENSION; ORAL

LINEZOLID**AB** HIKMA **100MG/5ML** **A200068 001** Jun 03, 2015ZYVOX**AB** +! PFIZER **100MG/5ML** **N021132 001** Apr 18, 2000

SOLUTION; INTRAVENOUS

LINEZOLID**AP** EUGIA PHARMA **600MG/300ML (2MG/ML)** **A206917 001** Aug 04, 2016**AP** FRESENIUS KABI USA **600MG/300ML (2MG/ML)** **A204764 001** Mar 15, 2016**AP** HOSPIRA INC **600MG/300ML (2MG/ML)** **A205442 001** Jul 07, 2015**AP** HQ SPCLT PHARMA **200MG/100ML (2MG/ML)** **A207001 001** Jul 07, 2017**AP** **600MG/300ML (2MG/ML)** **A207001 002** Jul 07, 2017**AP** MYLAN LABS LTD **200MG/100ML (2MG/ML)** **A205154 001** Dec 06, 2017**AP** **600MG/300ML (2MG/ML)** **A205154 002** Dec 06, 2017**AP** NANG KUANG PHARM CO **200MG/100ML (2MG/ML)** **A207354 001** Dec 20, 2016**AP** **600MG/300ML (2MG/ML)** **A207354 002** Dec 20, 2016**AP** SAGENT PHARMS INC **200MG/100ML (2MG/ML)** **A204696 001** Mar 02, 2017

PRESCRIPTION DRUG PRODUCT LIST

LINEZOLID

SOLUTION; INTRAVENOUS

LINEZOLID

<u>AP</u>		<u>600MG/300ML (2MG/ML)</u>	<u>A204696 002</u>	Mar 02, 2017
<u>AP</u>	SANDOZ INC	<u>200MG/100ML (2MG/ML)</u>	<u>A200904 001</u>	Jul 16, 2015
<u>AP</u>		<u>600MG/300ML (2MG/ML)</u>	<u>A200904 002</u>	Jul 16, 2015
<u>AP</u>	TEVA PHARMS	<u>600MG/300ML (2MG/ML)</u>	<u>A200222 001</u>	Jun 27, 2012

ZYVOX

<u>AP</u>	+	PFIZER	<u>200MG/100ML (2MG/ML)</u>	<u>N021131 001</u>	Apr 18, 2000
<u>AP</u>	+	!	<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

<u>AB</u>		ALEMbic PHARMS LTD	<u>600MG</u>	<u>A205233 001</u>	Dec 21, 2015
<u>AB</u>		ALKEM LABS LTD	<u>600MG</u>	<u>A205517 001</u>	Dec 21, 2015
<u>AB</u>		AMNEAL PHARMS	<u>600MG</u>	<u>A204536 001</u>	Dec 21, 2015
<u>AB</u>		CELLTRION	<u>600MG</u>	<u>A210702 001</u>	Apr 25, 2019
<u>AB</u>		GLENMARK PHARMS	<u>600MG</u>	<u>A078987 001</u>	Dec 21, 2015
<u>AB</u>		HETERO LABS LTD V	<u>600MG</u>	<u>A204239 001</u>	Dec 21, 2015
<u>AB</u>		NOVEL LABS INC	<u>600MG</u>	<u>A207526 001</u>	Aug 22, 2016
<u>AB</u>		ZYDUS PHARMS	<u>600MG</u>	<u>A206097 001</u>	Feb 22, 2017

ZYVOX

<u>AB</u>	+	!	PFIZER	<u>600MG</u>	<u>N021130 002</u>	Apr 18, 2000
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LIOTHYRONINE SODIUM

INJECTABLE; INJECTION

LIOTHYRONINE SODIUM

<u>AP</u>		XGEN PHARMS	<u>EQ 0.01MG BASE/ML</u>	<u>A076923 001</u>	Aug 17, 2005	
<u>AP</u>	+	!	PAR STERILE PRODUCTS	<u>EQ 0.01MG BASE/ML</u>	<u>N020105 001</u>	Dec 31, 1991

TABLET; ORAL

CYTOMEL

<u>AB</u>	+	KING PHARMS	<u>EQ 0.005MG BASE</u>	<u>N010379 001</u>	
<u>AB</u>	+		<u>EQ 0.025MG BASE</u>	<u>N010379 002</u>	
<u>AB</u>	+		<u>EQ 0.05MG BASE</u>	<u>N010379 003</u>	

LIOTHYRONINE SODIUM

<u>AB</u>		MAYNE PHARMA INC	<u>EQ 0.005MG BASE</u>	<u>A090097 001</u>	Mar 20, 2009
<u>AB</u>			<u>EQ 0.025MG BASE</u>	<u>A090097 002</u>	Mar 20, 2009
<u>AB</u>			<u>EQ 0.05MG BASE</u>	<u>A090097 003</u>	Mar 20, 2009
<u>AB</u>		SIGMAPHARM LABS LLC	<u>EQ 0.005MG BASE</u>	<u>A200295 001</u>	Nov 29, 2012
<u>AB</u>			<u>EQ 0.025MG BASE</u>	<u>A200295 002</u>	Nov 29, 2012
<u>AB</u>	!		<u>EQ 0.05MG BASE</u>	<u>A200295 003</u>	Nov 29, 2012
<u>AB</u>		SUN PHARM	<u>EQ 0.005MG BASE</u>	<u>A091382 001</u>	Apr 20, 2016
<u>AB</u>			<u>EQ 0.025MG BASE</u>	<u>A091382 002</u>	Apr 20, 2016
<u>AB</u>			<u>EQ 0.05MG BASE</u>	<u>A091382 003</u>	Apr 20, 2016
<u>AB</u>		TEVA PHARMS USA	<u>EQ 0.005MG BASE</u>	<u>A211510 001</u>	Oct 26, 2018
<u>AB</u>			<u>EQ 0.025MG BASE</u>	<u>A211510 002</u>	Oct 26, 2018
<u>AB</u>			<u>EQ 0.05MG BASE</u>	<u>A211510 003</u>	Oct 26, 2018
<u>AB</u>		ZYDUS	<u>EQ 0.005MG BASE</u>	<u>A214803 001</u>	Jan 22, 2021
<u>AB</u>			<u>EQ 0.025MG BASE</u>	<u>A214803 002</u>	Jan 22, 2021
<u>AB</u>			<u>EQ 0.05MG BASE</u>	<u>A214803 003</u>	Jan 22, 2021

LIRAGLUTIDE RECOMBINANT

SOLUTION; SUBCUTANEOUS

SAXENDA

	+	NOVO	18MG/3ML (6MG/ML)	N206321 001	Dec 23, 2014
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VICTOZA

	+	NOVO NORDISK INC	18MG/3ML (6MG/ML)	N022341 001	Jan 25, 2010
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LISDEXAMFETAMINE DIMESYLATE

CAPSULE; ORAL

VYVANSE

	+	TAKEDA PHARMS USA	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

	+	TAKEDA PHARMS USA	10MG	N208510 001	Jan 28, 2017
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PRESCRIPTION DRUG PRODUCT LIST

LISDEXAMFETAMINE DIMESYLATE

TABLET, CHEWABLE;ORAL

VYVANSE

+	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

+	AZURITY	1MG/ML	N208401	001	Jul 29, 2016
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TABLET;ORAL

LISINOPRIL

<u>AB</u>	ACCORD HLTHCARE	<u>2.5MG</u>	<u>A202554</u>	<u>001</u>	Jul 30, 2013
<u>AB</u>		<u>5MG</u>	<u>A202554</u>	<u>002</u>	Jul 30, 2013
<u>AB</u>		<u>10MG</u>	<u>A202554</u>	<u>003</u>	Jul 30, 2013
<u>AB</u>		<u>20MG</u>	<u>A202554</u>	<u>004</u>	Jul 30, 2013
<u>AB</u>		<u>30MG</u>	<u>A202554</u>	<u>005</u>	Jul 30, 2013
<u>AB</u>		<u>40MG</u>	<u>A202554</u>	<u>006</u>	Jul 30, 2013
<u>AB</u>	ASCENT PHARMS INC	<u>2.5MG</u>	<u>A075903</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A075903</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A075903</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075903</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A075903</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A075903</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>	ATLANTIDE	<u>2.5MG</u>	<u>A078402</u>	<u>001</u>	Apr 19, 2007
<u>AB</u>		<u>5MG</u>	<u>A078402</u>	<u>002</u>	Apr 19, 2007
<u>AB</u>		<u>10MG</u>	<u>A078402</u>	<u>003</u>	Apr 19, 2007
<u>AB</u>		<u>20MG</u>	<u>A078402</u>	<u>004</u>	Apr 19, 2007
<u>AB</u>		<u>30MG</u>	<u>A078402</u>	<u>005</u>	Apr 19, 2007
<u>AB</u>		<u>40MG</u>	<u>A078402</u>	<u>006</u>	Apr 19, 2007
<u>AB</u>	AUROBINDO	<u>2.5MG</u>	<u>A077622</u>	<u>001</u>	Feb 22, 2006
<u>AB</u>		<u>5MG</u>	<u>A077622</u>	<u>002</u>	Feb 22, 2006
<u>AB</u>		<u>10MG</u>	<u>A077622</u>	<u>003</u>	Feb 22, 2006
<u>AB</u>		<u>20MG</u>	<u>A077622</u>	<u>004</u>	Feb 22, 2006
<u>AB</u>		<u>30MG</u>	<u>A077622</u>	<u>005</u>	Feb 22, 2006
<u>AB</u>		<u>40MG</u>	<u>A077622</u>	<u>006</u>	Feb 22, 2006
<u>AB</u>	CHARTWELL RX	<u>2.5MG</u>	<u>A075994</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A075994</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A075994</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075994</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A075994</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A075994</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>	COREPHARMA	<u>2.5MG</u>	<u>A076102</u>	<u>001</u>	Sep 30, 2002
<u>AB</u>		<u>5MG</u>	<u>A076102</u>	<u>002</u>	Sep 30, 2002
<u>AB</u>		<u>10MG</u>	<u>A076102</u>	<u>003</u>	Sep 30, 2002
<u>AB</u>		<u>20MG</u>	<u>A076102</u>	<u>004</u>	Sep 30, 2002
<u>AB</u>		<u>30MG</u>	<u>A076102</u>	<u>005</u>	Sep 30, 2002
<u>AB</u>		<u>40MG</u>	<u>A076102</u>	<u>006</u>	Sep 30, 2002
<u>AB</u>	INVAGEN PHARMS	<u>2.5MG</u>	<u>A203508</u>	<u>001</u>	Oct 29, 2013
<u>AB</u>		<u>5MG</u>	<u>A203508</u>	<u>002</u>	Oct 29, 2013
<u>AB</u>		<u>10MG</u>	<u>A203508</u>	<u>003</u>	Oct 29, 2013
<u>AB</u>		<u>20MG</u>	<u>A203508</u>	<u>004</u>	Oct 29, 2013
<u>AB</u>		<u>30MG</u>	<u>A203508</u>	<u>005</u>	Oct 29, 2013
<u>AB</u>		<u>40MG</u>	<u>A203508</u>	<u>006</u>	Oct 29, 2013
<u>AB</u>	LUPIN	<u>2.5MG</u>	<u>A077321</u>	<u>001</u>	Sep 09, 2005
<u>AB</u>		<u>5MG</u>	<u>A077321</u>	<u>002</u>	Sep 09, 2005
<u>AB</u>		<u>10MG</u>	<u>A077321</u>	<u>003</u>	Sep 09, 2005
<u>AB</u>		<u>20MG</u>	<u>A077321</u>	<u>004</u>	Sep 09, 2005
<u>AB</u>		<u>30MG</u>	<u>A077321</u>	<u>005</u>	Sep 09, 2005
<u>AB</u>		<u>40MG</u>	<u>A077321</u>	<u>006</u>	Sep 09, 2005
<u>AB</u>	PRINSTON INC	<u>2.5MG</u>	<u>A075743</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>2.5MG</u>	<u>A076164</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A075743</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A076164</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A075743</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A076164</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075743</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A076164</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A075743</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A076164</u>	<u>002</u>	Jul 01, 2002

PRESCRIPTION DRUG PRODUCT LIST

LISINOPRIL

TABLET; ORAL

LISINOPRIL

<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
<u>AB</u>		<u>40MG</u>	<u>A076059 006</u>	Jul 01, 2002
<u>AB</u>	YILING	<u>2.5MG</u>	<u>A212041 001</u>	Sep 15, 2020
<u>AB</u>		<u>5MG</u>	<u>A212041 002</u>	Sep 15, 2020
<u>AB</u>		<u>10MG</u>	<u>A212041 003</u>	Sep 15, 2020
<u>AB</u>		<u>20MG</u>	<u>A208920 001</u>	Mar 04, 2021
<u>AB</u>		<u>30MG</u>	<u>A208920 002</u>	Mar 04, 2021
<u>AB</u>		<u>40MG</u>	<u>A208920 003</u>	Mar 04, 2021

PRINIVIL

<u>AB</u>	MERCK	<u>5MG</u>	<u>N019558 001</u>	Dec 29, 1987
<u>AB</u>		<u>40MG</u>	<u>N019558 004</u>	Oct 25, 1988

ZESTRIL

<u>AB</u>	+ ALMATICA	<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>	+ HIKMA	<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>	+	HIKMA	<u>300MG</u>	<u>N018558 001</u>	Jan 29, 1982
<u>AB</u>		SUN PHARM INDS INC	<u>300MG</u>	<u>A091027 001</u>	Jun 24, 2010

TABLET, EXTENDED RELEASE; ORAL

LITHIUM CARBONATE

<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>	HIKMA	<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>	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

LITHOBID

<u>AB</u>	+	ANI PHARMS	<u>300MG</u>	<u>N018027 001</u>	
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PRESCRIPTION DRUG PRODUCT LIST

LITHIUM CITRATE

SYRUP; ORAL

LITHIUM CITRATE

AA	+ !	HIKMA	EQ 300MG CARBONATE/5ML	N018421 001	
AA		WOCKHARDT BIO AG	EQ 300MG CARBONATE/5ML	A070755 001	May 21, 1986

LODOXAMIDE TROMETHAMINESOLUTION/DROPS; OPHTHALMIC
ALOMIDE

+ !	NOVARTIS	EQ 0.1% BASE	N020191 001	Sep 23, 1993
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LOFEXIDINE HYDROCHLORIDE

TABLET; ORAL

LUCEMYRA

+ !	USWM	EQ 0.18MG BASE	N209229 001	May 16, 2018
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LOMITAPIDE MESYLATE

CAPSULE; ORAL

JUXTAPID

+	AMRYT	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

LOMUSTINE

CAPSULE; ORAL

GLEOSTINE

+	CORDEN PHARMA	10MG	N017588 001	
+		40MG	N017588 002	
+ !		100MG	N017588 003	

LONAFARNIB

CAPSULE; ORAL

ZOKINVY

+	EIGER BIOPHARMS	50MG	N213969 001	Nov 20, 2020
+ !		75MG	N213969 002	Nov 20, 2020

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL

LOPERAMIDE HYDROCHLORIDE

AB		BIONPHARMA INC	2MG	A215579 001	Oct 22, 2021
AB		EDENBRIDGE PHARMS	2MG	A215001 001	Oct 06, 2021
AB	!	MYLAN	2MG	A072741 001	Sep 18, 1991
AB		TEVA	2MG	A073192 001	Apr 30, 1992

LOPINAVIR; RITONAVIR

SOLUTION; ORAL

KALETRA

AA	+ !	ABBVIE	80MG/ML; 20MG/ML	N021251 001	Sep 15, 2000
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LOPINAVIR AND RITONAVIR

AA		LANNETT CO INC	80MG/ML; 20MG/ML	A207407 001	Dec 27, 2016
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TABLET; ORAL

KALETRA

AB	+	ABBVIE	100MG; 25MG	N021906 002	Nov 09, 2007
AB	+ !		200MG; 50MG	N021906 001	Oct 28, 2005

LOPINAVIR AND RITONAVIR

AB		HETERO LABS LTD III	100MG; 25MG	A091677 001	Jun 04, 2021
AB			200MG; 50MG	A091677 002	Jun 04, 2021

LORAZEPAM

CAPSULE, EXTENDED RELEASE; ORAL

LOREEV XR

+	ALMATICA	1MG	N214826 001	Aug 27, 2021
+		2MG	N214826 002	Aug 27, 2021
+ !		3MG	N214826 003	Aug 27, 2021

CONCENTRATE; ORAL

LORAZEPAM

AA		AKORN	2MG/ML	A200169 001	Jan 30, 2012
AA		AMNEAL PHARMS	2MG/ML	A091383 001	Dec 23, 2009
AA		LANNETT CO INC	2MG/ML	A079244 001	Apr 28, 2009
AA		LUPIN LTD	2MG/ML	A091407 001	Feb 19, 2013
AA		PHARM ASSOC	2MG/ML	A090260 001	Jun 15, 2010
AA	!	HIKMA	2MG/ML	A072755 001	Jun 28, 1991

PRESCRIPTION DRUG PRODUCT LIST

LORAZEPAM

INJECTABLE; INJECTION

ATIVAN

<u>AP</u>	<u>+!</u>	HIKMA	<u>2MG/ML</u>	<u>N018140</u>	<u>001</u>	
<u>AP</u>	<u>+!</u>		<u>4MG/ML</u>	<u>N018140</u>	<u>002</u>	

LORAZEPAM

<u>AP</u>		AKORN	<u>2MG/ML</u>	<u>A075025</u>	<u>001</u>	Jul 23, 1998
<u>AP</u>		HOSPIRA	<u>2MG/ML</u>	<u>A074243</u>	<u>001</u>	Apr 12, 1994
<u>AP</u>			<u>2MG/ML</u>	<u>A074282</u>	<u>001</u>	May 27, 1994
<u>AP</u>			<u>4MG/ML</u>	<u>A074243</u>	<u>002</u>	Apr 12, 1994
<u>AP</u>			<u>4MG/ML</u>	<u>A074282</u>	<u>002</u>	May 27, 1994
<u>AP</u>		INTL MEDICATION SYS	<u>2MG/ML</u>	<u>A076150</u>	<u>001</u>	Nov 15, 2004

TABLET; ORAL

ATIVAN

<u>AB</u>	<u>+</u>	BAUSCH	<u>0.5MG</u>	<u>N017794</u>	<u>001</u>	
<u>AB</u>	<u>+</u>		<u>1MG</u>	<u>N017794</u>	<u>002</u>	
<u>AB</u>	<u>+!</u>		<u>2MG</u>	<u>N017794</u>	<u>003</u>	

LORAZEPAM

<u>AB</u>		AMNEAL PHARMS	<u>0.5MG</u>	<u>A078826</u>	<u>001</u>	Jun 23, 2010
<u>AB</u>			<u>1MG</u>	<u>A078826</u>	<u>002</u>	Jun 23, 2010
<u>AB</u>			<u>2MG</u>	<u>A078826</u>	<u>003</u>	Jun 23, 2010
<u>AB</u>		AUROLIFE PHARMA LLC	<u>0.5MG</u>	<u>A203572</u>	<u>001</u>	Dec 22, 2017
<u>AB</u>			<u>1MG</u>	<u>A203572</u>	<u>002</u>	Dec 22, 2017
<u>AB</u>			<u>2MG</u>	<u>A203572</u>	<u>003</u>	Dec 22, 2017
<u>AB</u>		LEADING PHARMA LLC	<u>0.5MG</u>	<u>A078203</u>	<u>001</u>	Jul 30, 2007
<u>AB</u>			<u>1MG</u>	<u>A078203</u>	<u>002</u>	Jul 30, 2007
<u>AB</u>			<u>2MG</u>	<u>A078203</u>	<u>003</u>	Jul 30, 2007
<u>AB</u>		OXFORD PHARMS	<u>0.5MG</u>	<u>A077754</u>	<u>001</u>	May 10, 2006
<u>AB</u>			<u>1MG</u>	<u>A077754</u>	<u>002</u>	May 10, 2006
<u>AB</u>			<u>2MG</u>	<u>A077754</u>	<u>003</u>	May 10, 2006
<u>AB</u>		SANDOZ	<u>0.5MG</u>	<u>A071141</u>	<u>002</u>	Apr 21, 1987
<u>AB</u>			<u>1MG</u>	<u>A071141</u>	<u>003</u>	Apr 21, 1987
<u>AB</u>			<u>2MG</u>	<u>A071141</u>	<u>001</u>	Apr 21, 1987
<u>AB</u>		WATSON LABS	<u>0.5MG</u>	<u>A072926</u>	<u>001</u>	Oct 31, 1991
<u>AB</u>			<u>1MG</u>	<u>A072927</u>	<u>001</u>	Oct 31, 1991
<u>AB</u>			<u>2MG</u>	<u>A072928</u>	<u>001</u>	Oct 31, 1991

LORLATINIB

TABLET; ORAL

LORBRENA

<u>+</u>	PFIZER	25MG	<u>N210868</u>	<u>001</u>	Nov 02, 2018
<u>+!</u>		100MG	<u>N210868</u>	<u>002</u>	Nov 02, 2018

LOSARTAN POTASSIUM

TABLET; ORAL

COZAAR

<u>AB</u>	<u>+</u>	ORGANON	<u>25MG</u>	<u>N020386</u>	<u>001</u>	Apr 14, 1995
<u>AB</u>	<u>+</u>		<u>50MG</u>	<u>N020386</u>	<u>002</u>	Apr 14, 1995
<u>AB</u>	<u>+!</u>		<u>100MG</u>	<u>N020386</u>	<u>003</u>	Oct 13, 1998

LOSARTAN POTASSIUM

<u>AB</u>		ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A090428</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>			<u>50MG</u>	<u>A090428</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>			<u>100MG</u>	<u>A090428</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>		AUROBINDO PHARMA	<u>25MG</u>	<u>A090083</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>			<u>50MG</u>	<u>A090083</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>			<u>100MG</u>	<u>A090083</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>		HETERO LABS LTD V	<u>25MG</u>	<u>A203835</u>	<u>001</u>	Aug 12, 2015
<u>AB</u>			<u>50MG</u>	<u>A203835</u>	<u>002</u>	Aug 12, 2015
<u>AB</u>			<u>100MG</u>	<u>A203835</u>	<u>003</u>	Aug 12, 2015
<u>AB</u>		HIKMA	<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>		IPCA LABS LTD	<u>25MG</u>	<u>A200290</u>	<u>001</u>	Aug 30, 2013
<u>AB</u>			<u>50MG</u>	<u>A200290</u>	<u>002</u>	Aug 30, 2013
<u>AB</u>			<u>100MG</u>	<u>A200290</u>	<u>003</u>	Aug 30, 2013
<u>AB</u>		JUBILANT CADISTA	<u>25MG</u>	<u>A201170</u>	<u>001</u>	Sep 18, 2012
<u>AB</u>			<u>50MG</u>	<u>A201170</u>	<u>002</u>	Sep 18, 2012
<u>AB</u>			<u>100MG</u>	<u>A201170</u>	<u>003</u>	Sep 18, 2012
<u>AB</u>		LUPIN LTD	<u>25MG</u>	<u>A078232</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>			<u>50MG</u>	<u>A078232</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>			<u>100MG</u>	<u>A078232</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>		MACLEODS PHARMS LTD	<u>25MG</u>	<u>A202230</u>	<u>001</u>	May 30, 2012
<u>AB</u>			<u>50MG</u>	<u>A202230</u>	<u>002</u>	May 30, 2012

PRESCRIPTION DRUG PRODUCT LIST

LOVASTATIN

TABLET; ORAL

LOVASTATIN

AB	SANDOZ	10MG	A075636 001	Dec 17, 2001
AB		20MG	A075636 002	Dec 17, 2001
AB		40MG	A075636 003	Dec 17, 2001
AB	TEVA	10MG	A075551 003	Dec 17, 2001
AB		20MG	A075551 002	Dec 17, 2001
AB		40MG	A075551 001	Dec 17, 2001

TABLET, EXTENDED RELEASE; ORAL

ALTOPREV

+	COVIS	20MG	N021316 002	Jun 26, 2002
+		40MG	N021316 003	Jun 26, 2002
+	!	60MG	N021316 004	Jun 26, 2002

LOXAPINE

POWDER; INHALATION

ADASUVE

+	!	ALEXZA PHARMS	10MG	N022549 001	Dec 21, 2012
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LOXAPINE SUCCINATE

CAPSULE; ORAL

LOXAPINE SUCCINATE

AB	ELITE LABS INC	EQ 5MG BASE	A076868 001	Aug 04, 2005
AB		EQ 10MG BASE	A076868 002	Aug 04, 2005
AB		EQ 25MG BASE	A076868 003	Aug 04, 2005
AB		EQ 50MG BASE	A076868 004	Aug 04, 2005
AB	LANNETT CO INC	EQ 5MG BASE	A090695 001	Sep 26, 2011
AB		EQ 10MG BASE	A090695 002	Sep 26, 2011
AB		EQ 25MG BASE	A090695 003	Sep 26, 2011
AB		EQ 50MG BASE	A090695 004	Sep 26, 2011
AB	WATSON LABS	EQ 5MG BASE	A072204 001	Jun 15, 1988
AB		EQ 10MG BASE	A072205 001	Jun 15, 1988
AB	!	EQ 25MG BASE	A072206 001	Jun 15, 1988
AB		EQ 50MG BASE	A072062 001	Jun 15, 1988

LUBIPROSTONE

CAPSULE; ORAL

AMITIZA

AB	+	SUCAMPO PHARMA LLC	8MCG	N021908 002	Apr 29, 2008
AB	+	!	24MCG	N021908 001	Jan 31, 2006

LUBIPROSTONE

AB	AMNEAL	8MCG	A209450 001	Nov 30, 2021
AB		24MCG	A209450 002	Nov 30, 2021

LULICONAZOLE

CREAM; TOPICAL

LUZU

+	!	BAUSCH	1%	N204153 001	Nov 14, 2013
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LUMASIRAN SODIUM

SOLUTION; SUBCUTANEOUS

OXLUMO

+	!	ALNYLAM PHARMS INC	EQ 94.5MG/0.5ML BASE (EQ 94.5MG/0.5ML BASE)	N214103 001	Nov 23, 2020
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LUMATEPERONE TOSYLATE

CAPSULE; ORAL

CAPLYTA

+	!	INTRA-CELLULAR	EQ 42MG BASE	N209500 001	Dec 20, 2019
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LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LATUDA

AB	+	SUNOVION PHARMS INC	20MG	N200603 003	Dec 07, 2011
AB	+	!	40MG	N200603 001	Oct 28, 2010
AB	+		60MG	N200603 005	Jul 12, 2013
AB	+		80MG	N200603 002	Oct 28, 2010
AB	+		120MG	N200603 004	Apr 26, 2012

LURASIDONE HYDROCHLORIDE

AB	ACCORD HLTHCARE	20MG	A208049 001	Jan 03, 2019
AB		40MG	A208049 002	Jan 03, 2019
AB		60MG	A208049 003	Jan 03, 2019
AB		80MG	A208049 004	Jan 03, 2019
AB		120MG	A208049 005	Jan 03, 2019
AB	ALEMBIC PHARMS LTD	20MG	A213248 001	May 13, 2021

PRESCRIPTION DRUG PRODUCT LIST

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LURASIDONE HYDROCHLORIDE

<u>AB</u>		<u>40MG</u>	<u>A213248 002</u>	May 13, 2021
<u>AB</u>		<u>60MG</u>	<u>A213248 003</u>	May 13, 2021
<u>AB</u>		<u>80MG</u>	<u>A213248 004</u>	May 13, 2021
<u>AB</u>		<u>120MG</u>	<u>A213248 005</u>	May 13, 2021
<u>AB</u>	DR REDDYS LABS LTD	<u>20MG</u>	<u>A208047 001</u>	Aug 24, 2021
<u>AB</u>		<u>40MG</u>	<u>A208047 002</u>	Aug 24, 2021
<u>AB</u>		<u>60MG</u>	<u>A208047 003</u>	Aug 24, 2021
<u>AB</u>		<u>80MG</u>	<u>A208047 004</u>	Aug 24, 2021
<u>AB</u>	SUN PHARM	<u>20MG</u>	<u>A208066 001</u>	Jan 04, 2019
<u>AB</u>		<u>40MG</u>	<u>A208066 002</u>	Jan 04, 2019
<u>AB</u>		<u>60MG</u>	<u>A208066 003</u>	Jan 04, 2019
<u>AB</u>		<u>80MG</u>	<u>A208066 004</u>	Jan 04, 2019
<u>AB</u>		<u>120MG</u>	<u>A208066 005</u>	Jan 04, 2019

LURBINECTEDIN

POWDER; INTRAVENOUS

ZEPZELCA

+! JAZZ

4MG/VIAL

N213702 001 Jun 15, 2020

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

10MG

N204410 001 Oct 18, 2013

MAFENIDE ACETATE

CREAM; TOPICAL

SULFAMYLN

+! RISING

EQ 85MG BASE/GM

N016763 001

FOR SOLUTION; TOPICAL

MAFENIDE ACETATE

<u>AT</u>	NOVAST LABS	<u>5%</u>	<u>A206716 001</u>	Jul 31, 2017
<u>AT</u>	PAR FORM	<u>5%</u>	<u>A201511 001</u>	Feb 12, 2013

SULFAMYLN

<u>AT</u>	+! MYLAN INSTITUTIONAL	<u>5%</u>	<u>N019832 003</u>	Jun 05, 1998
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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

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

PRESCRIPTION DRUG PRODUCT LIST

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

SOLUTION;IRRIGATION

PHYSIOLYTE IN PLASTIC CONTAINER

B BRAUN	30MG/100ML;37MG/100ML;370MG/100ML;530MG	N019024	001	Jun 08, 1984
	/100ML;500MG/100ML			

PHYSIOSOL IN PLASTIC CONTAINER

ICU MEDICAL INC	30MG/100ML;37MG/100ML;222MG/100ML;526MG	N017637	002	Jul 08, 1982
	/100ML;502MG/100ML			

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
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NORMOCARB HF 35

+! DIALYSIS SUPS	0.21GM/100ML;3.97GM/100ML;8.3GM/100ML	N021910	002	Jul 26, 2006
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MAGNESIUM SULFATE

INJECTABLE;INJECTION

MAGNESIUM SULFATE

<u>AP</u>	B BRAUN MEDICAL INC	<u>2GM/50ML (40MG/ML)</u>	<u>A207967</u>	<u>001</u>	Apr 26, 2021
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<u>AP</u>		<u>4GM/100ML (40MG/ML)</u>	<u>A207967</u>	<u>002</u>	Apr 26, 2021
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<u>AP</u>		<u>4GM/50ML (80MG/ML)</u>	<u>A207967</u>	<u>003</u>	Apr 26, 2021
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MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN MEDICAL INC	<u>1GM/100ML</u>	<u>A207966</u>	<u>001</u>	Oct 27, 2020
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<u>AP</u>	BAXTER HLTHCARE	<u>1GM/100ML</u>	<u>A211965</u>	<u>001</u>	Aug 11, 2020
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<u>AP</u>	FRESENIUS KABI USA	<u>1GM/100ML</u>	<u>A206486</u>	<u>001</u>	Mar 07, 2016
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<u>AP</u>	+! HOSPIRA	<u>1GM/100ML</u>	<u>N020488</u>	<u>001</u>	Jul 11, 1995
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<u>AP</u>	HQ SPCLT PHARMA	<u>1GM/100ML</u>	<u>A207349</u>	<u>001</u>	Mar 02, 2016
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<u>AP</u>	MYLAN LABS LTD	<u>1GM/100ML</u>	<u>A209932</u>	<u>001</u>	Sep 10, 2018
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MAGNESIUM SULFATE IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>2GM/50ML (40MG/ML)</u>	<u>A211966</u>	<u>001</u>	Jun 01, 2020
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<u>AP</u>	CORP				
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<u>AP</u>		<u>4GM/50ML (80MG/ML)</u>	<u>A211966</u>	<u>002</u>	Jun 01, 2020
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<u>AP</u>		<u>4GM/100ML (40MG/ML)</u>	<u>A211966</u>	<u>003</u>	Jun 01, 2020
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<u>AP</u>	FRESENIUS KABI USA	<u>4GM/100ML (40MG/ML)</u>	<u>A206485</u>	<u>001</u>	Mar 15, 2016
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<u>AP</u>		<u>4GM/50ML (80MG/ML)</u>	<u>A206485</u>	<u>002</u>	Mar 15, 2016
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<u>AP</u>		<u>2GM/50ML (40MG/ML)</u>	<u>A206485</u>	<u>003</u>	Mar 15, 2016
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<u>AP</u>		<u>20GM/500ML (40MG/ML)</u>	<u>A206485</u>	<u>004</u>	Mar 15, 2016
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<u>AP</u>		<u>40GM/1000ML (40MG/ML)</u>	<u>A206485</u>	<u>005</u>	Mar 15, 2016
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<u>AP</u>	GLAND PHARMA LTD	<u>2GM/50ML (40MG/ML)</u>	<u>A213917</u>	<u>001</u>	Jul 10, 2020
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<u>AP</u>		<u>4GM/100ML (40MG/ML)</u>	<u>A213917</u>	<u>002</u>	Jul 10, 2020
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<u>AP</u>	+ HOSPIRA	<u>2GM/50ML (40MG/ML)</u>	<u>N020309</u>	<u>003</u>	Jan 26, 2007
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<u>AP</u>	+!	<u>4GM/100ML (40MG/ML)</u>	<u>N020309</u>	<u>001</u>	Jun 24, 1994
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<u>AP</u>	+!	<u>4GM/50ML (80MG/ML)</u>	<u>N020309</u>	<u>002</u>	Jun 24, 1994
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<u>AP</u>	+	<u>20GM/500ML (40MG/ML)</u>	<u>N020309</u>	<u>004</u>	Jan 18, 1995
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<u>AP</u>	+	<u>40GM/1000ML (40MG/ML)</u>	<u>N020309</u>	<u>005</u>	Jan 18, 1995
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<u>AP</u>	HQ SPCLT PHARMA	<u>2GM/50ML (40MG/ML)</u>	<u>A207350</u>	<u>001</u>	Dec 06, 2017
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<u>AP</u>		<u>4GM/100ML (40MG/ML)</u>	<u>A207350</u>	<u>002</u>	Dec 06, 2017
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<u>AP</u>		<u>4GM/50ML (80MG/ML)</u>	<u>A207350</u>	<u>003</u>	Dec 06, 2017
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<u>AP</u>		<u>20GM/500ML (40MG/ML)</u>	<u>A207350</u>	<u>004</u>	Dec 06, 2017
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<u>AP</u>		<u>40GM/1000ML (40MG/ML)</u>	<u>A207350</u>	<u>005</u>	Dec 06, 2017
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<u>AP</u>	MYLAN LABS LTD	<u>2GM/50ML (40MG/ML)</u>	<u>A209911</u>	<u>001</u>	Sep 14, 2018
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<u>AP</u>		<u>4GM/100ML (40MG/ML)</u>	<u>A209911</u>	<u>002</u>	Sep 14, 2018
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<u>AP</u>	PTS CONSULTING	<u>2GM/50ML (40MG/ML)</u>	<u>A209642</u>	<u>001</u>	Nov 08, 2021
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<u>AP</u>		<u>4GM/100ML (40MG/ML)</u>	<u>A209642</u>	<u>002</u>	Nov 08, 2021
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<u>AP</u>		<u>4GM/50ML (80MG/ML)</u>	<u>A209642</u>	<u>003</u>	Nov 08, 2021
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MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER

+! HOSPIRA	2GM/100ML	N020488	002	Jul 11, 1995
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SOLUTION;INTRAMUSCULAR, INTRAVENOUS

MAGNESIUM SULFATE

<u>AP</u>	EXELA PHARMA	<u>5GM/10ML (500MG/ML)</u>	<u>A206039</u>	<u>001</u>	Dec 18, 2014
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<u>AP</u>	+! FRESENIUS KABI USA	<u>5GM/10ML (500MG/ML)</u>	<u>N019316</u>	<u>001</u>	Sep 08, 1986
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<u>AP</u>	+!	<u>10GM/20ML (500MG/ML)</u>	<u>N019316</u>	<u>003</u>	Jan 29, 2016
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<u>AP</u>	! HOSPIRA	<u>5GM/10ML (500MG/ML)</u>	<u>A075151</u>	<u>001</u>	Apr 25, 2000
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<u>AP</u>	HOSPIRA INC	<u>10GM/20ML (500MG/ML)</u>	<u>A202411</u>	<u>001</u>	May 14, 2015
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<u>AP</u>	+! FRESENIUS KABI USA	1GM/2ML (500MG/ML)	N019316	002	Sep 08, 1986
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<u>AP</u>	+!	25GM/50ML (500MG/ML)	N019316	004	Jan 29, 2016
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PRESCRIPTION DRUG PRODUCT LIST

MAGNESIUM SULFATE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION;IRRIGATION

TIS-U-SOL

AT	BAXTER HLTHCARE	20MG/100ML;40MG/100ML;6.25MG/100ML;800MG/100ML;8.75MG/100ML	N018508 001	Feb 19, 1982
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TIS-U-SOL IN PLASTIC CONTAINER

AT	BAXTER HLTHCARE	20MG/100ML;40MG/100ML;6.25MG/100ML;800MG/100ML;8.75MG/100ML	N018336 001	
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MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM SULFATE

TABLET;ORAL

SUTAB

+	BRAINTREE LABS	0.225GM;0.188GM;1.479GM	N213135 001	Nov 10, 2020
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MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE

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, 2017
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SUPREP BOWEL PREP KIT

AA	+	BRAINTREE LABS	1.6GM/BOT;3.13GM/BOT;17.5GM/BOT	N022372 001	Aug 05, 2010
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MALATHION

LOTION;TOPICAL

MALATHION

!	SUVEN PHARMS	0.5%	A091559 001	May 23, 2012
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MANGANESE CHLORIDE

INJECTABLE;INJECTION

MANGANESE CHLORIDE IN PLASTIC CONTAINER

+	HOSPIRA	EQ 0.1MG MANGANESE/ML	N018962 001	Jun 26, 1986
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MANNITOL

INJECTABLE;INJECTION

MANNITOL 10% IN PLASTIC CONTAINER

AP	B BRAUN	10GM/100ML	N020006 002	Jul 26, 1993
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MANNITOL 15% IN PLASTIC CONTAINER

AP	B BRAUN	15GM/100ML	N020006 003	Jul 26, 1993
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MANNITOL 20% IN PLASTIC CONTAINER

AP	B BRAUN	20GM/100ML	N020006 004	Jul 26, 1993
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AP	ICU MEDICAL INC	20GM/100ML	N019603 004	Jan 08, 1990
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MANNITOL 25%

AP	FRESENIUS KABI USA	12.5GM/50ML	A080677 001	
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AP	HOSPIRA	12.5GM/50ML	N016269 006	Aug 25, 1994
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MANNITOL 5% IN PLASTIC CONTAINER

AP	B BRAUN	5GM/100ML	N020006 001	Jul 26, 1993
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OSMITROL 10% IN WATER

AP	BAXTER HLTHCARE	10GM/100ML	N013684 002	
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OSMITROL 10% IN WATER IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	10GM/100ML	N013684 006	
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OSMITROL 15% IN WATER

AP	BAXTER HLTHCARE	15GM/100ML	N013684 004	
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OSMITROL 15% IN WATER IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	15GM/100ML	N013684 008	
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OSMITROL 20% IN WATER

AP	BAXTER HLTHCARE	20GM/100ML	N013684 003	
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OSMITROL 20% IN WATER IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	20GM/100ML	N013684 007	
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OSMITROL 5% IN WATER

AP	BAXTER HLTHCARE	5GM/100ML	N013684 001	
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OSMITROL 5% IN WATER IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	5GM/100ML	N013684 005	
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POWDER;INHALATION

ARIDOL KIT

+	PHARMAXIS LTD	N/A, 5MG, 10MG, 20MG, 40MG	N022368 001	Oct 05, 2010
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BRONCHITOL

+	CHIESI	40MG	N202049 001	Oct 30, 2020
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SOLUTION;IRRIGATION

RESECTISOL IN PLASTIC CONTAINER

	B BRAUN	5GM/100ML	N016772 002	
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PRESCRIPTION DRUG PRODUCT LIST

MANNITOL; SORBITOL

SOLUTION;IRRIGATION

SORBITOL-MANNITOL IN PLASTIC CONTAINER

+! ICU MEDICAL INC 540MG/100ML;2.7GM/100ML N018316 001

MARALIXIBAT CHLORIDE

SOLUTION;ORAL

LIVMARLI

+! MIRUM EQ 9.5MG BASE/ML N214662 001 Sep 29, 2021

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

MARIBAVIR

TABLET;ORAL

LIVTENCITY

+! TAKEDA PHARMS USA 200MG N215596 001 Nov 23, 2021

MEBENDAZOLE

TABLET, CHEWABLE;ORAL

EMVERM

! IMPAX LABS INC 100MG A073580 001 Jan 04, 1995

MECAMYLAMINE HYDROCHLORIDE

TABLET;ORAL

MECAMYLAMINE HYDROCHLORIDE

! LGM PHARMA 2.5MG A204054 001 Mar 19, 2013

MECHLORETHAMINE HYDROCHLORIDE

GEL;TOPICAL

VALCHLOR

+! HELSINN EQ 0.016% BASE N202317 001 Aug 23, 2013

MECLIZINE HYDROCHLORIDE

TABLET;ORAL

ANTIVERT**AA + CASPER PHARMA LLC****12.5MG****N010721 006****AA +****25MG****N010721 004****AA +****50MG****N010721 001** Jan 20, 1982**MECLIZINE HYDROCHLORIDE****AA ! AMNEAL PHARMS****12.5MG****A201451 001** Feb 23, 2011**AA !****25MG****A201451 002** Feb 23, 2011**AA ANI PHARMS****12.5MG****A084657 002****AA****25MG****A084657 001****AA ANNORA PHARMA****12.5MG****A087128 002****AA****25MG****A087128 001****AA EPIC PHARMA LLC****12.5MG****A200294 001** Apr 13, 2012**AA****25MG****A200294 002** Apr 13, 2012**AA INDICUS PHARMA****12.5MG****A205136 001** Feb 22, 2019**AA****25MG****A205136 002** Feb 22, 2019**AA****50MG****A205136 003** Feb 22, 2019**AA INVATECH****25MG****A084092 003** May 22, 1989**AA JUBILANT CADISTA****12.5MG****A040659 001** Jun 04, 2010**AA****25MG****A040659 002** Jun 04, 2010**AA SANDOZ****12.5MG****A084843 002** May 22, 1989**AA ZYDUS****12.5MG****A213957 001** Jun 23, 2020**AA****25MG****A213957 002** Jun 23, 2020

TABLET, CHEWABLE;ORAL

ANTIVERT**AA + CASPER PHARMA LLC****25MG****N010721 005**MECLOFENAMATE SODIUM

CAPSULE;ORAL

MECLOFENAMATE SODIUM

MYLAN

EQ 50MG BASE

A071081 002 Sep 03, 1986

! EQ 100MG BASE

A071081 001 Sep 03, 1986

PRESCRIPTION DRUG PRODUCT LIST

MEDROXYPROGESTERONE ACETATE

INJECTABLE; INJECTION

DEPO-PROVERA

AB	+ !	PFIZER	150MG/ML	N020246	001	Oct 29, 1992
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MEDROXYPROGESTERONE ACETATE

AB		AMPHASTAR PHARMS INC	150MG/ML	A077235	001	Nov 28, 2017
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AB			150MG/ML	A077334	001	Nov 28, 2017
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AB		MYLAN LABS LTD	150MG/ML	A210227	001	Oct 12, 2018
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AB		SUN PHARM	150MG/ML	A210760	001	May 01, 2019
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AB			150MG/ML	A210761	001	Apr 24, 2019
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AB		TEVA PHARMS USA	150MG/ML	A076553	001	Jul 28, 2004
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INJECTABLE; SUBCUTANEOUS

DEPO-SUBQ PROVERA 104

+ !	PFIZER	104MG/0.65ML	N021583	001	Dec 17, 2004
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TABLET; ORAL

MEDROXYPROGESTERONE ACETATE

AB		BARR	2.5MG	A040159	001	Aug 09, 1996
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AB			5MG	A040159	002	Aug 09, 1996
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AB			10MG	A040159	003	Aug 09, 1996
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PROVERA

AB	+	PFIZER	2.5MG	N011839	001	
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AB	+		5MG	N011839	003	
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AB	+ !		10MG	N011839	004	
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MEFENAMIC ACID

CAPSULE; ORAL

MEFENAMIC ACID

AB		BELCHER	250MG	A091608	001	Jun 02, 2014
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AB		LUPIN LTD	250MG	A091322	001	Jul 22, 2011
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AB		MICRO LABS	250MG	A090562	001	Nov 19, 2010
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PONSTEL

AB	+ !	AVION PHARMS	250MG	N015034	003	
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MEFLOQUINE HYDROCHLORIDE

TABLET; ORAL

MEFLOQUINE HYDROCHLORIDE

AB	!	BARR	250MG	A076392	001	Dec 29, 2003
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AB		HIKMA	250MG	A076523	001	Oct 01, 2004
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MEGESTROL ACETATE

SUSPENSION; ORAL

MEGACE ES

AB	+ !	ENDO PHARMS INC	125MG/ML	N021778	001	Jul 05, 2005
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MEGESTROL ACETATE

AB		AKORN	40MG/ML	A203960	001	Jun 09, 2017
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AB		BRECKENRIDGE	125MG/ML	A204688	001	Dec 01, 2017
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AB	!	STRIDES PHARMA	40MG/ML	A075671	001	Jul 25, 2001
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AB		TWI PHARMS	125MG/ML	A203139	001	Aug 27, 2014
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AB		WOCKHARDT BIO AG	40MG/ML	A076721	001	Nov 01, 2004
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TABLET; ORAL

MEGESTROL ACETATE

AB		BARR	20MG	A074621	002	Aug 16, 1996
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AB			40MG	A074621	001	Nov 30, 1995
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AB		STRIDES PHARMA	20MG	A072422	001	Aug 08, 1988
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AB	!		40MG	A072423	001	Aug 08, 1988
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MELOXICAM

CAPSULE; ORAL

MELOXICAM

AB		LUPIN LTD	5MG	A209487	001	Jun 01, 2020
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AB	!		10MG	A209487	002	Jun 01, 2020
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AB		NOVITIUM PHARMA	5MG	A211398	001	Mar 09, 2021
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AB			10MG	A211398	002	Mar 09, 2021
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SOLUTION; INTRAVENOUS

ANJESO

+ !	BAUDAX	30MG/ML (30MG/ML)	N210583	001	Feb 20, 2020
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TABLET; ORAL

MELOXICAM

AB		APOTEX INC	7.5MG	A077882	001	Jul 20, 2006
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AB			15MG	A077882	002	Jul 20, 2006
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AB		AUROBINDO PHARMA	7.5MG	A078008	001	Oct 02, 2006
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AB			15MG	A078008	002	Oct 02, 2006
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AB		CIPLA	7.5MG	A077929	001	Jul 19, 2006
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AB			15MG	A077929	002	Jul 19, 2006
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PRESCRIPTION DRUG PRODUCT LIST

MELOXICAM

TABLET; ORAL

MELOXICAM

<u>AB</u>	DR REDDYS LABS INC	<u>7.5MG</u>	<u>A077931 001</u>	Jul 25, 2006
<u>AB</u>		<u>15MG</u>	<u>A077931 002</u>	Jul 25, 2006
<u>AB</u>	GLENMARK GENERICS	<u>7.5MG</u>	<u>A077932 001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077932 002</u>	Jul 19, 2006
<u>AB</u>	LUPIN PHARMS	<u>7.5MG</u>	<u>A077944 001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077944 002</u>	Jul 19, 2006
<u>AB</u>	PURACAP PHARM	<u>7.5MG</u>	<u>A077938 001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077938 002</u>	Jul 19, 2006
<u>AB</u>	STRIDES PHARMA	<u>7.5MG</u>	<u>A077928 001</u>	May 13, 2009
<u>AB</u>		<u>15MG</u>	<u>A077928 002</u>	May 13, 2009
<u>AB</u>	TARO	<u>7.5MG</u>	<u>A078102 001</u>	Nov 07, 2006
<u>AB</u>		<u>15MG</u>	<u>A078102 002</u>	Nov 07, 2006
<u>AB</u>	UNICHEM	<u>7.5MG</u>	<u>A077927 001</u>	Dec 20, 2006
<u>AB</u>		<u>15MG</u>	<u>A077927 002</u>	Dec 20, 2006
<u>AB</u>	YUNG SHIN PHARM	<u>7.5MG</u>	<u>A077918 001</u>	Dec 07, 2006
<u>AB</u>		<u>15MG</u>	<u>A077918 002</u>	Dec 07, 2006
<u>AB</u>	ZYDUS PHARMS USA	<u>7.5MG</u>	<u>A077921 001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077921 002</u>	Jul 19, 2006

MOBIC

<u>AB</u>	+ BOEHRINGER INGELHEIM	<u>7.5MG</u>	<u>N020938 001</u>	Apr 13, 2000
<u>AB</u>	+!	<u>15MG</u>	<u>N020938 002</u>	Aug 23, 2000

MELPHALAN

TABLET; ORAL

MELPHALAN

!	ALVOGEN	2MG	A207809 001	Mar 22, 2017
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MELPHALAN HYDROCHLORIDE

INJECTABLE; INJECTION

MELPHALAN HYDROCHLORIDE

<u>AP</u>	ACTAVIS LLC	<u>EQ 50MG BASE/VIAL</u>	<u>A206018 001</u>	Dec 19, 2016
<u>AP</u>	ALMAJECT	<u>EQ 50MG BASE/VIAL</u>	<u>A204817 001</u>	May 17, 2019
<u>AP</u>	BPI LABS LLC	<u>EQ 50MG BASE/VIAL</u>	<u>A209197 001</u>	May 08, 2020
<u>AP</u>	DR REDDYS LABS LTD	<u>EQ 50MG BASE/VIAL</u>	<u>A203655 001</u>	Dec 08, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 50MG BASE/VIAL</u>	<u>A203393 001</u>	Dec 22, 2017
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 50MG BASE/VIAL</u>	<u>A209826 001</u>	May 28, 2019
<u>AP</u>	HIKMA	<u>EQ 50MG BASE/VIAL</u>	<u>A090303 001</u>	Oct 28, 2010
<u>AP</u>	INGENUS PHARMS LLC	<u>EQ 50MG BASE/VIAL</u>	<u>A210947 001</u>	Feb 18, 2020
<u>AP</u>	MEITHEAL	<u>EQ 50MG BASE/VIAL</u>	<u>A212960 001</u>	May 28, 2021
<u>AP</u>	! MYLAN INSTITUTIONAL	<u>EQ 50MG BASE/VIAL</u>	<u>A090270 001</u>	Jun 09, 2009
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 50MG BASE/VIAL</u>	<u>A201379 001</u>	Feb 28, 2017

POWDER; INTRAVENOUS

EVOMELA

+	ACROTECH	EQ 50MG BASE/VIAL	N207155 001	Mar 10, 2016
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MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MEMANTINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS	<u>7MG</u>	<u>A205825 001</u>	Oct 12, 2016
<u>AB</u>		<u>14MG</u>	<u>A205825 002</u>	Oct 12, 2016
<u>AB</u>		<u>21MG</u>	<u>A205825 003</u>	Oct 12, 2016
<u>AB</u>		<u>28MG</u>	<u>A205825 004</u>	Oct 12, 2016
<u>AB</u>	ANCHEN PHARMS	<u>7MG</u>	<u>A205784 001</u>	Jun 09, 2017
<u>AB</u>		<u>14MG</u>	<u>A205784 002</u>	Jun 09, 2017
<u>AB</u>		<u>21MG</u>	<u>A205784 003</u>	Jun 09, 2017
<u>AB</u>		<u>28MG</u>	<u>A205784 004</u>	Jun 09, 2017
<u>AB</u>	ANI PHARMS	<u>7MG</u>	<u>A205365 001</u>	Feb 28, 2020
<u>AB</u>		<u>14MG</u>	<u>A205365 002</u>	Feb 28, 2020
<u>AB</u>		<u>21MG</u>	<u>A205365 003</u>	Feb 28, 2020
<u>AB</u>		<u>28MG</u>	<u>A205365 004</u>	Feb 28, 2020
<u>AB</u>	APOTEX	<u>7MG</u>	<u>A206135 001</u>	Nov 22, 2016
<u>AB</u>		<u>14MG</u>	<u>A206135 002</u>	Nov 22, 2016
<u>AB</u>		<u>21MG</u>	<u>A206135 003</u>	Nov 22, 2016
<u>AB</u>		<u>28MG</u>	<u>A206135 004</u>	Nov 22, 2016
<u>AB</u>	AUROBINDO PHARMA LTD	<u>7MG</u>	<u>A214651 001</u>	Aug 09, 2021
<u>AB</u>		<u>14MG</u>	<u>A214651 002</u>	Aug 09, 2021
<u>AB</u>		<u>21MG</u>	<u>A214651 003</u>	Aug 09, 2021
<u>AB</u>		<u>28MG</u>	<u>A214651 004</u>	Aug 09, 2021
<u>AB</u>	LUPIN LTD	<u>7MG</u>	<u>A206028 001</u>	Sep 28, 2016

PRESCRIPTION DRUG PRODUCT LIST

MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

MEMANTINE HYDROCHLORIDE

<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>	YICHANG HUMANWELL	<u>7MG</u>	<u>A211100</u>	<u>001</u>	Apr 02, 2021
<u>AB</u>		<u>14MG</u>	<u>A211100</u>	<u>002</u>	Apr 02, 2021
<u>AB</u>		<u>21MG</u>	<u>A211100</u>	<u>003</u>	Apr 02, 2021
<u>AB</u>		<u>28MG</u>	<u>A211100</u>	<u>004</u>	Apr 02, 2021
<u>AB</u>	ZYDUS PHARMS	<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

NAMENDA XR

<u>AB</u>	+ ALLERGAN	<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	<u>2MG/ML</u>	<u>A209955</u>	<u>001</u>	Feb 09, 2018
<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
<u>AA</u>	SETON PHARMS	<u>2MG/ML</u>	<u>A210319</u>	<u>001</u>	Aug 31, 2020

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>	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>	CADILA	<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>AB</u>	CELLTRION	<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>	CSPC OUYI	<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>	HIKMA	<u>5MG</u>	<u>A208173</u>	<u>001</u>	Feb 28, 2020
<u>AB</u>		<u>10MG</u>	<u>A208173</u>	<u>002</u>	Feb 28, 2020
<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>	POLYGEN PHARMS	<u>5MG</u>	<u>A210587</u>	<u>001</u>	Dec 11, 2020
<u>AB</u>		<u>10MG</u>	<u>A210587</u>	<u>002</u>	Dec 11, 2020
<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 PHARM	<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>	UNICHEM	<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

NAMENDA

<u>AB</u>	+ ALLERGAN	<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

PRESCRIPTION DRUG PRODUCT LIST

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DEMEROL

<u>AP</u>	<u>+</u> !	HOSPIRA	<u>25MG/ML</u>	<u>N021171</u>	<u>001</u>	
<u>AP</u>	<u>+</u> !		<u>50MG/ML</u>	<u>N021171</u>	<u>002</u>	
<u>AP</u>	<u>+</u> !		<u>75MG/ML</u>	<u>N021171</u>	<u>003</u>	
<u>AP</u>	<u>+</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
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SYRUP; ORAL

MEPERIDINE HYDROCHLORIDE

!	HIKMA	50MG/5ML	A088744	001	Jan 30, 1985
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TABLET; ORAL

MEPERIDINE HYDROCHLORIDE

	EPIC PHARMA	50MG	A040331	001	May 28, 1999
		100MG	A040331	002	May 28, 1999

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CARBOCAINE

<u>AP</u>	<u>+</u> !	HOSPIRA	<u>1%</u>	<u>N012250</u>	<u>001</u>	
<u>AP</u>	<u>+</u> !		<u>1.5%</u>	<u>N012250</u>	<u>005</u>	
<u>AP</u>	<u>+</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
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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

MERCAPTOPYRINE

SUSPENSION; ORAL

PURIXAN

+!	NOVA LABS LTD	20MG/ML	N205919	001	Apr 28, 2014
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TABLET; ORAL

MERCAPTOPYRINE

<u>AB</u>		DR REDDYS LABS SA	<u>50MG</u>	<u>A040461</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>	!	HIKMA	<u>50MG</u>	<u>A040528</u>	<u>001</u>	Feb 13, 2004
<u>AB</u>		MYLAN	<u>50MG</u>	<u>A040594</u>	<u>001</u>	Jul 01, 2005

PURINETHOL

<u>AB</u>	+	STASON PHARMS	<u>50MG</u>	<u>N009053</u>	<u>002</u>	
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MEROPENEM

INJECTABLE; INJECTION

MEROPENEM

<u>AP</u>		ACS DOBFAR	<u>500MG/VIAL</u>	<u>A091404</u>	<u>001</u>	Oct 26, 2011
<u>AP</u>			<u>1GM/VIAL</u>	<u>A091404</u>	<u>002</u>	Oct 26, 2011
<u>AP</u>		ACS DOBFAR SPA	<u>500MG/VIAL</u>	<u>A204139</u>	<u>001</u>	Jun 09, 2016
<u>AP</u>			<u>1GM/VIAL</u>	<u>A204139</u>	<u>002</u>	Jun 09, 2016
<u>AP</u>		AMNEAL PHARMS	<u>500MG/VIAL</u>	<u>A205883</u>	<u>001</u>	Apr 12, 2016
<u>AP</u>			<u>1GM/VIAL</u>	<u>A205883</u>	<u>002</u>	Apr 12, 2016
<u>AP</u>		DAEWOONG PHARM CO	<u>500MG/VIAL</u>	<u>A204854</u>	<u>001</u>	Dec 18, 2015
<u>AP</u>			<u>1GM/VIAL</u>	<u>A204854</u>	<u>002</u>	Dec 18, 2015

PRESCRIPTION DRUG PRODUCT LIST

MEROPENEM

INJECTABLE; INJECTION

MEROPENEM

<u>AP</u>	EUGIA PHARMA	<u>500MG/VIAL</u>	<u>A205835</u>	<u>001</u>	Mar 27, 2017
<u>AP</u>		<u>1GM/VIAL</u>	<u>A205835</u>	<u>002</u>	Mar 27, 2017
<u>AP</u>	GLAND	<u>500MG/VIAL</u>	<u>A206141</u>	<u>001</u>	Jun 08, 2016
<u>AP</u>		<u>1GM/VIAL</u>	<u>A206141</u>	<u>002</u>	Jun 08, 2016
<u>AP</u>	HQ SPCLT PHARMA	<u>500MG/VIAL</u>	<u>A210773</u>	<u>001</u>	Aug 16, 2019
<u>AP</u>		<u>1GM/VIAL</u>	<u>A210773</u>	<u>002</u>	Aug 16, 2019
<u>AP</u>	SAVIOR LIFETEC CORP	<u>500MG/VIAL</u>	<u>A206086</u>	<u>001</u>	Apr 19, 2016
<u>AP</u>		<u>1GM/VIAL</u>	<u>A206086</u>	<u>002</u>	Apr 19, 2016

MERREM

<u>AP</u>	+! PFIZER	<u>500MG/VIAL</u>	<u>N050706</u>	<u>003</u>	Jun 21, 1996
<u>AP</u>	+!	<u>1GM/VIAL</u>	<u>N050706</u>	<u>001</u>	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	1GM/VIAL;1GM/VIAL	N209776	001	Aug 29, 2017
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MESALAMINE

CAPSULE, DELAYED RELEASE; ORAL

DELZICOL

<u>AB</u>	+! APIL	<u>400MG</u>	<u>N204412</u>	<u>001</u>	Feb 01, 2013
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MESALAMINE

<u>AB</u>	TEVA PHARMS USA	<u>400MG</u>	<u>A207873</u>	<u>001</u>	May 09, 2019
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CAPSULE, EXTENDED RELEASE; ORAL

APRISO

<u>AB</u>	+! VALEANT PHARMS INTL	<u>375MG</u>	<u>N022301</u>	<u>001</u>	Oct 31, 2008
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MESALAMINE

<u>AB</u>	ALKEM LABS LTD	<u>375MG</u>	<u>A214242</u>	<u>001</u>	Jul 15, 2021
<u>AB</u>	MYLAN	<u>375MG</u>	<u>A207271</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>	ZYDUS PHARMS	<u>375MG</u>	<u>A208954</u>	<u>001</u>	Aug 12, 2021

PENTASA

	+ TAKEDA PHARMS USA	250MG	N020049	001	May 10, 1993
	+!	500MG	N020049	002	Jul 08, 2004

ENEMA; RECTAL

MESALAMINE

<u>AB</u>	PADAGIS ISRAEL	<u>4GM/60ML</u>	<u>A076751</u>	<u>001</u>	Sep 17, 2004
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ROWASA

<u>AB</u>	+! MYLAN SPECIALITY LP	<u>4GM/60ML</u>	<u>N019618</u>	<u>001</u>	Dec 24, 1987
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SFROWASA

<u>AB</u>	+ MYLAN SPECIALITY LP	<u>4GM/60ML</u>	<u>N019618</u>	<u>002</u>	Jun 20, 2008
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SUPPOSITORY; RECTAL

CANASA

<u>AB</u>	+! ALLERGAN	<u>1GM</u>	<u>N021252</u>	<u>002</u>	Nov 05, 2004
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MESALAMINE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>1GM</u>	<u>A205654</u>	<u>001</u>	Aug 14, 2020
<u>AB</u>	AMNEAL	<u>1GM</u>	<u>A210509</u>	<u>001</u>	Jan 02, 2020
<u>AB</u>	AMRING PHARMS	<u>1GM</u>	<u>A208362</u>	<u>001</u>	Jun 21, 2019
<u>AB</u>	ANNORA PHARMA	<u>1GM</u>	<u>A213377</u>	<u>001</u>	Mar 19, 2020
<u>AB</u>	MYLAN	<u>1GM</u>	<u>A204354</u>	<u>001</u>	Nov 24, 2015
<u>AB</u>	PHARM SOURCING	<u>1GM</u>	<u>A207448</u>	<u>001</u>	Apr 19, 2019
<u>AB</u>	SANDOZ INC	<u>1GM</u>	<u>A202065</u>	<u>001</u>	Jun 12, 2019
<u>AB</u>	ZYDUS PHARMS	<u>1GM</u>	<u>A208953</u>	<u>001</u>	Feb 12, 2020

TABLET, DELAYED RELEASE; ORAL

ASACOL HD

<u>AB</u>	+! APIL	<u>800MG</u>	<u>N021830</u>	<u>001</u>	May 29, 2008
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LIALDA

<u>AB</u>	+! TAKEDA PHARMS USA	<u>1.2GM</u>	<u>N022000</u>	<u>001</u>	Jan 16, 2007
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MESALAMINE

<u>AB</u>	ACTAVIS LABS FL	<u>1.2GM</u>	<u>A203817</u>	<u>001</u>	Mar 23, 2018
<u>AB</u>	SUN PHARM	<u>1.2GM</u>	<u>A211858</u>	<u>001</u>	Jan 25, 2019
<u>AB</u>	ZYDUS PHARMS	<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

PRESCRIPTION DRUG PRODUCT LIST

MESNA

INJECTABLE; INTRAVENOUS

MESNA

AP	FRESENIUS KABI USA	100MG/ML	A075811 001	Apr 26, 2001
AP	GLAND	100MG/ML	A206992 001	Dec 18, 2017
AP	HIKMA	100MG/ML	A075739 001	Jan 09, 2004
AP	SAGENT PHARMS INC	100MG/ML	A090913 001	Apr 13, 2010

MESNEX

AP	+ ! BAXTER HLTHCARE	100MG/ML	N019884 001	Dec 30, 1988
TABLET; ORAL				
MESNEX				
+ !	BAXTER HLTHCARE	400MG	N020855 001	Mar 21, 2002

METAPROTERENOL SULFATE

SYRUP; ORAL

METAPROTERENOL SULFATE

! GENUS

10MG/5ML

A073632 001 Jul 22, 1992

METARAMINOL BITARTRATE

INJECTABLE; INJECTION

METARAMINOL BITARTRATE

SLAYBACK PHARMA LLC EQ 10MG BASE/ML

A211304 001 Aug 24, 2021

METAXALONE

TABLET; ORAL

METAXALONE

AB	ACTAVIS LABS FL INC	800MG	A203695 001	Jun 15, 2017
AB	AMNEAL PHARMS	800MG	A203399 001	Jun 21, 2013
AB	LANNETT CO INC	800MG	A204770 001	Nov 22, 2016
AB	MOUNTAIN	400MG	A040486 001	Feb 27, 2015
AB	SANDOZ	800MG	A040445 001	Mar 31, 2010
AB	SCIEGEN PHARMS INC	400MG	A207466 002	Mar 13, 2020
AB		800MG	A207466 001	Aug 31, 2017

SKELAXIN

AB	+ ! KING PHARMS	800MG	N013217 003	Aug 30, 2002
METAXALONE				
EX	APPCO	800MG	A208774 001	Sep 24, 2018

METFORMIN HYDROCHLORIDE

SOLUTION; ORAL

METFORMIN HYDROCHLORIDE

AB	SAPTALIS PHARMS	500MG/5ML	A211309 001	Mar 03, 2020
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RIOMET

AB	+ ! RANBAXY	500MG/5ML	N021591 001	Sep 11, 2003
TABLET; ORAL				

METFORMIN HYDROCHLORIDE

AB	ALKEM	500MG	A091184 001	Nov 01, 2010
AB		850MG	A091184 002	Nov 01, 2010
AB		1GM	A091184 003	Nov 01, 2010
AB	AMNEAL PHARMS NY	500MG	A077880 001	Jun 05, 2006
AB		850MG	A077880 002	Jun 05, 2006
AB		1GM	A077880 003	Jun 05, 2006
AB	APOTEX	500MG	A075984 001	Apr 23, 2002
AB		500MG	A090666 001	Dec 07, 2011
AB		850MG	A075984 002	Apr 23, 2002
AB		850MG	A090666 002	Dec 07, 2011
AB		1GM	A075984 003	Apr 23, 2002
AB		1GM	A090666 003	Dec 07, 2011
AB	ATLAS PHARMS LLC	500MG	A076033 001	Jan 24, 2002
AB		850MG	A076033 002	Jan 24, 2002
AB		1GM	A076033 003	Jan 24, 2002
AB	AUROBINDO	500MG	A077095 001	Jan 14, 2005
AB		850MG	A077095 002	Jan 14, 2005
AB		1GM	A077095 003	Jan 14, 2005
AB	CHARTWELL	500MG	A075972 001	Jan 24, 2002
AB		850MG	A075972 002	Jan 24, 2002
AB		1GM	A075972 003	Jan 24, 2002
AB	CSPC OUYI	500MG	A205096 001	Jul 11, 2016
AB		850MG	A205096 002	Jul 11, 2016
AB		1GM	A205096 003	Jul 11, 2016
AB	DR REDDYS LABS INC	500MG	A077787 001	Aug 23, 2006
AB		850MG	A077787 002	Aug 23, 2006
AB		1GM	A077787 003	Aug 23, 2006
AB	GLENMARK GENERICS	500MG	A078170 001	May 23, 2008

PRESCRIPTION DRUG PRODUCT LIST

METFORMIN HYDROCHLORIDE

TABLET; ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>		<u>850MG</u>	<u>A078170</u>	<u>002</u>	May 23, 2008
<u>AB</u>		<u>1GM</u>	<u>A078170</u>	<u>003</u>	May 23, 2008
<u>AB</u>	GRANULES INDIA	<u>500MG</u>	<u>A090564</u>	<u>001</u>	Apr 22, 2010
<u>AB</u>		<u>850MG</u>	<u>A090564</u>	<u>002</u>	Apr 22, 2010
<u>AB</u>	!	<u>1GM</u>	<u>A090564</u>	<u>003</u>	Apr 22, 2010
<u>AB</u>	HARMAN FINOCHEM	<u>500MG</u>	<u>A213320</u>	<u>001</u>	Dec 03, 2021
<u>AB</u>		<u>850MG</u>	<u>A213320</u>	<u>002</u>	Dec 03, 2021
<u>AB</u>		<u>1GM</u>	<u>A213320</u>	<u>003</u>	Dec 03, 2021
<u>AB</u>	LAURUS	<u>500MG</u>	<u>A209882</u>	<u>001</u>	Aug 27, 2018
<u>AB</u>		<u>850MG</u>	<u>A209882</u>	<u>002</u>	Aug 27, 2018
<u>AB</u>		<u>1GM</u>	<u>A209882</u>	<u>003</u>	Aug 27, 2018
<u>AB</u>	MARKSANS PHARMA	<u>500MG</u>	<u>A090888</u>	<u>001</u>	Mar 12, 2012
<u>AB</u>		<u>850MG</u>	<u>A090888</u>	<u>002</u>	Mar 12, 2012
<u>AB</u>		<u>1GM</u>	<u>A090888</u>	<u>003</u>	Mar 12, 2012
<u>AB</u>	MYLAN	<u>500MG</u>	<u>A075973</u>	<u>001</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075973</u>	<u>002</u>	Jan 25, 2002
<u>AB</u>		<u>1GM</u>	<u>A075973</u>	<u>003</u>	Jan 25, 2002
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>A075965</u>	<u>001</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075965</u>	<u>002</u>	Jan 25, 2002
<u>AB</u>		<u>1GM</u>	<u>A075965</u>	<u>003</u>	Jan 25, 2002
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A203769</u>	<u>001</u>	Sep 11, 2013
<u>AB</u>		<u>850MG</u>	<u>A203769</u>	<u>002</u>	Sep 11, 2013
<u>AB</u>		<u>1GM</u>	<u>A203769</u>	<u>003</u>	Sep 11, 2013
<u>AB</u>	SUN PHARM INDS INC	<u>500MG</u>	<u>A075967</u>	<u>001</u>	Jan 29, 2002
<u>AB</u>		<u>850MG</u>	<u>A075967</u>	<u>002</u>	Jan 29, 2002
<u>AB</u>		<u>1GM</u>	<u>A075967</u>	<u>003</u>	Jan 29, 2002
<u>AB</u>	ZYDUS HLTHCARE	<u>500MG</u>	<u>A203686</u>	<u>001</u>	Aug 28, 2014
<u>AB</u>		<u>850MG</u>	<u>A203686</u>	<u>002</u>	Aug 28, 2014
<u>AB</u>		<u>1GM</u>	<u>A203686</u>	<u>003</u>	Aug 28, 2014
<u>AB</u>	ZYDUS PHARMS USA	<u>500MG</u>	<u>A077064</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>850MG</u>	<u>A077064</u>	<u>002</u>	Apr 18, 2005
<u>AB</u>		<u>1GM</u>	<u>A077064</u>	<u>003</u>	Apr 18, 2005
<u>AB</u>	CHARTWELL	625MG	A075972	005	Jan 24, 2002
		750MG	A075972	004	Jan 24, 2002

TABLET, EXTENDED RELEASE; ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>750MG</u>	<u>A076869</u>	<u>001</u>	Apr 12, 2005
<u>AB</u>	!	<u>750MG</u>	<u>A206145</u>	<u>002</u>	Oct 22, 2018
<u>AB</u>	AUROBINDO PHARMA LTD	<u>750MG</u>	<u>A079118</u>	<u>002</u>	Jul 20, 2012
<u>AB</u>	BEXIMCO PHARMS USA	<u>750MG</u>	<u>A207427</u>	<u>002</u>	Dec 13, 2016
<u>AB</u>	CADILA	<u>750MG</u>	<u>A077078</u>	<u>001</u>	Apr 21, 2005
<u>AB</u>	CSPC OUYI	<u>750MG</u>	<u>A078321</u>	<u>002</u>	Apr 17, 2008
<u>AB</u>	GRANULES	<u>750MG</u>	<u>A209313</u>	<u>002</u>	Mar 16, 2018
<u>AB</u>	INTELLIPHARMACEUTICS	<u>750MG</u>	<u>A202306</u>	<u>002</u>	Feb 23, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>750MG</u>	<u>A206955</u>	<u>002</u>	Dec 07, 2016
<u>AB</u>	MARKSANS PHARMA	<u>750MG</u>	<u>A090295</u>	<u>002</u>	Apr 29, 2016
<u>AB</u>	NOSTRUM PHARMS LLC	<u>750MG</u>	<u>A076756</u>	<u>002</u>	Dec 12, 2011
<u>AB</u>	PRINSTON INC	<u>750MG</u>	<u>A208880</u>	<u>002</u>	Sep 10, 2018
<u>AB</u>	SUN PHARM INDS (IN)	<u>750MG</u>	<u>A077336</u>	<u>002</u>	Feb 09, 2006
<u>AB</u>	TEVA	<u>750MG</u>	<u>A076864</u>	<u>001</u>	Apr 12, 2005
<u>AB</u>	UNICHEM	<u>750MG</u>	<u>A213359</u>	<u>002</u>	Aug 11, 2021
<u>AB</u>	YICHANG HUMANWELL	<u>750MG</u>	<u>A211052</u>	<u>002</u>	Sep 24, 2018
<u>AB1</u>	ALIGNSCIENCE PHARMA	<u>500MG</u>	<u>A209303</u>	<u>001</u>	Mar 19, 2018
<u>AB1</u>	ALKEM LABS LTD	<u>500MG</u>	<u>A206145</u>	<u>001</u>	Oct 22, 2018
<u>AB1</u>	AUROBINDO PHARMA LTD	<u>500MG</u>	<u>A079118</u>	<u>001</u>	Jul 20, 2012
<u>AB1</u>	BEXIMCO PHARMS USA	<u>500MG</u>	<u>A207427</u>	<u>001</u>	Dec 13, 2016
<u>AB1</u>	CADILA	<u>500MG</u>	<u>A077060</u>	<u>001</u>	Apr 20, 2005
<u>AB1</u>	CSPC OUYI	<u>500MG</u>	<u>A078321</u>	<u>001</u>	Apr 17, 2008
<u>AB1</u>	GRANULES	<u>500MG</u>	<u>A209313</u>	<u>001</u>	Mar 16, 2018
<u>AB1</u>	INTELLIPHARMACEUTICS	<u>500MG</u>	<u>A202306</u>	<u>001</u>	Feb 23, 2017
<u>AB1</u>	INVENTIA	<u>500MG</u>	<u>A201991</u>	<u>001</u>	Jan 18, 2012
<u>AB1</u>	MACLEODS PHARMS LTD	<u>500MG</u>	<u>A206955</u>	<u>001</u>	Dec 07, 2016
<u>AB1</u>	MARKSANS PHARMA	<u>500MG</u>	<u>A090295</u>	<u>001</u>	Apr 29, 2016
<u>AB1</u>	NOSTRUM PHARMS LLC	<u>500MG</u>	<u>A076756</u>	<u>001</u>	Jul 26, 2006
<u>AB1</u>	PRINSTON INC	<u>500MG</u>	<u>A208880</u>	<u>001</u>	Sep 10, 2018
<u>AB1</u>	SANDOZ	<u>500MG</u>	<u>A076873</u>	<u>001</u>	Dec 14, 2004

PRESCRIPTION DRUG PRODUCT LIST

METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METFORMIN HYDROCHLORIDE

AB1	SUN PHARM INDS (IN)	500MG	A077336 001	Feb 09, 2006
AB1	TEVA	500MG	A076269 001	Jun 18, 2004
AB1	UNICHEM	500MG	A213359 001	Aug 11, 2021
AB1	YICHANG HUMANWELL	500MG	A211052 001	Sep 24, 2018

FORTAMET

AB2	+ ANDRX LABS LLC	1GM	N021574 002	Apr 27, 2004
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METFORMIN HYDROCHLORIDE

AB2	AJANTA PHARMA LTD	500MG	A213651 001	Apr 09, 2020
AB2		1GM	A213651 002	Apr 09, 2020
AB2	AMTA	500MG	A213394 001	Aug 03, 2021
AB2		1GM	A213394 002	Aug 03, 2021
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
AB2	TWI PHARMS	500MG	A213247 001	Sep 29, 2021
AB2		1GM	A213247 002	Sep 29, 2021

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
AB3	AJANTA PHARMA LTD	500MG	A213962 001	Mar 09, 2021
AB3		1GM	A213962 002	Mar 09, 2021
AB3	APOTEX	500MG	A213356 001	Dec 13, 2021
AB3		1GM	A213356 002	Dec 13, 2021
AB3	GLENMARK PHARMS LTD	500MG	A212969 001	Nov 25, 2019
AB3		1GM	A212969 002	Nov 25, 2019
AB3	GRANULES	500MG	A213344 001	Jan 12, 2021
AB3		1GM	A213344 002	Jan 12, 2021
AB3	LUPIN LTD	500MG	A091664 001	Jul 19, 2013
AB3		1GM	A091664 002	Jul 19, 2013
AB3	SCIEGEN PHARMS INC	500MG	A213334 001	Apr 16, 2021
AB3		1GM	A213334 002	Apr 16, 2021
AB3	SUN PHARM	500MG	A202917 001	Aug 01, 2016
AB3		1GM	A202917 002	Aug 01, 2016
BX	AMNEAL PHARMS NY	500MG	A078596 001	Jan 03, 2008
BX		750MG	A078596 002	Jan 03, 2008

METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

ACTOPLUS MET

AB	+ TAKEDA PHARMS USA	500MG;EQ 15MG BASE	N021842 001	Aug 29, 2005
AB	+	850MG;EQ 15MG BASE	N021842 002	Aug 29, 2005

PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE

AB	AUROBINDO PHARMA LTD	500MG;EQ 15MG BASE	A200823 001	Feb 13, 2013
AB		850MG;EQ 15MG BASE	A200823 002	Feb 13, 2013
AB	MACLEODS PHARMS LTD	500MG;EQ 15MG BASE	A204802 001	Nov 05, 2015
AB		850MG;EQ 15MG BASE	A204802 002	Nov 05, 2015
AB	TEVA PHARMS USA	500MG;EQ 15MG BASE	A091155 001	Mar 10, 2014
AB		850MG;EQ 15MG BASE	A091155 002	Mar 10, 2014

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

PRESCRIPTION DRUG PRODUCT LIST

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
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METHADONE HYDROCHLORIDE

CONCENTRATE; ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>		HIKMA	<u>10MG/ML</u>	<u>A040180</u>	<u>001</u>	Apr 30, 1998
<u>AA</u>		LANNETT CO INC	<u>10MG/ML</u>	<u>A212093</u>	<u>001</u>	Nov 02, 2020
<u>AA</u>		SPECGX LLC	<u>10MG/ML</u>	<u>A207368</u>	<u>001</u>	Aug 22, 2019
<u>AA</u>		VISTAPHARM	<u>10MG/ML</u>	<u>A040088</u>	<u>001</u>	Nov 30, 1994

METHADONE HYDROCHLORIDE INTENSOL

<u>AA</u>		HIKMA	<u>10MG/ML</u>	<u>A089897</u>	<u>001</u>	Sep 06, 1988
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METHADOSE

<u>AA</u>	+	!	SPECGX LLC	<u>10MG/ML</u>	<u>N017116</u>	<u>002</u>
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INJECTABLE; INJECTION

METHADONE HYDROCHLORIDE

<u>AP</u>		AKORN	<u>10MG/ML</u>	<u>A208306</u>	<u>001</u>	Oct 27, 2017
<u>AP</u>	+	!	MYLAN INSTITUTIONAL	<u>10MG/ML</u>	<u>N021624</u>	<u>001</u>

SOLUTION; ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>	+	!	HIKMA	<u>5MG/5ML</u>	<u>A087393</u>	<u>001</u>	
<u>AA</u>	+	!		<u>10MG/5ML</u>	<u>A087997</u>	<u>001</u>	Aug 30, 1982
<u>AA</u>			SPECGX LLC	<u>5MG/5ML</u>	<u>A207537</u>	<u>001</u>	Oct 02, 2019
<u>AA</u>				<u>10MG/5ML</u>	<u>A207537</u>	<u>002</u>	Oct 01, 2019
<u>AA</u>			VISTAPHARM	<u>5MG/5ML</u>	<u>A090707</u>	<u>001</u>	Jun 30, 2010
<u>AA</u>				<u>10MG/5ML</u>	<u>A090707</u>	<u>002</u>	Jun 30, 2010

TABLET; ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>		ASCENT PHARMS INC	<u>5MG</u>	<u>A211228</u>	<u>001</u>	Jan 03, 2019
<u>AA</u>			<u>10MG</u>	<u>A211228</u>	<u>002</u>	Jan 03, 2019
<u>AA</u>		AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A203502</u>	<u>001</u>	Aug 31, 2015
<u>AA</u>			<u>10MG</u>	<u>A203502</u>	<u>002</u>	Aug 31, 2015
<u>AA</u>		EPIC PHARMA LLC	<u>5MG</u>	<u>A090065</u>	<u>001</u>	Aug 18, 2015
<u>AA</u>			<u>10MG</u>	<u>A090065</u>	<u>002</u>	Aug 18, 2015
<u>AA</u>		NOSTRUM LABS INC	<u>5MG</u>	<u>A210484</u>	<u>001</u>	Aug 02, 2018
<u>AA</u>			<u>10MG</u>	<u>A210484</u>	<u>002</u>	Aug 02, 2018
<u>AA</u>		ROXANE	<u>5MG</u>	<u>A088108</u>	<u>001</u>	Mar 08, 1983
<u>AA</u>			<u>10MG</u>	<u>A088109</u>	<u>001</u>	Mar 08, 1983
<u>AA</u>	!		<u>5MG</u>	<u>A040517</u>	<u>001</u>	Apr 27, 2004
<u>AA</u>	!		<u>10MG</u>	<u>A040517</u>	<u>002</u>	Apr 27, 2004
<u>AA</u>		SUN PHARM INDUSTRIES	<u>5MG</u>	<u>A208305</u>	<u>001</u>	Mar 30, 2018
<u>AA</u>			<u>10MG</u>	<u>A208305</u>	<u>002</u>	Mar 30, 2018
<u>AA</u>		THEPHARMANETWORK LLC	<u>5MG</u>	<u>A090635</u>	<u>002</u>	Sep 22, 2020
<u>AA</u>			<u>10MG</u>	<u>A090635</u>	<u>001</u>	Nov 25, 2009
<u>AA</u>		VISTAPHARM	<u>10MG</u>	<u>A040241</u>	<u>002</u>	May 29, 1998
<u>AA</u>			<u>10MG</u>	<u>A204166</u>	<u>001</u>	Mar 16, 2020

TABLET, FOR SUSPENSION; ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>	+	!	HIKMA	<u>40MG</u>	<u>N017058</u>	<u>001</u>	
<u>AA</u>			SPECGX LLC	<u>40MG</u>	<u>A077142</u>	<u>001</u>	Jul 12, 2005
<u>AA</u>			VISTAPHARM	<u>40MG</u>	<u>A075082</u>	<u>001</u>	Mar 25, 1998

METHADOSE

<u>AA</u>			SPECGX LLC	<u>40MG</u>	<u>A074184</u>	<u>001</u>	Apr 29, 1993
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PRESCRIPTION DRUG PRODUCT LIST

METHAMPHETAMINE HYDROCHLORIDE

TABLET; ORAL

DESOXYN

AA	+	RECORDATI RARE	5MG	N005378	002	
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METHAMPHETAMINE HYDROCHLORIDE

AA		HIKMA	5MG	A203846	001	Nov 17, 2015
AA		MAYNE PHARMA INC	5MG	A091189	001	Apr 21, 2010

METHAZOLAMIDE

TABLET; ORAL

METHAZOLAMIDE

AB		ANI PHARMS	25MG	A040001	001	Jun 30, 1993
AB			50MG	A040001	002	Jun 30, 1993
AB		BAUSCH	25MG	A207438	001	Oct 05, 2018
AB			50MG	A207438	002	Oct 05, 2018
AB		MIKART	25MG	A040062	001	Jan 27, 1994
AB	!		50MG	A040062	002	Jan 27, 1994
AB		SANDOZ	25MG	A040036	001	Jun 30, 1993
AB			50MG	A040036	002	Jun 30, 1993

METHENAMINE HIPPURATE

TABLET; ORAL

METHENAMINE HIPPURATE

AB	!	AUROBINDO PHARMA LTD	1GM	A205661	001	Jul 05, 2016
AB		MICRO LABS	1GM	A212172	001	Aug 01, 2019
AB		NOVAST LABS	1GM	A210068	001	Nov 27, 2020

UREX

AB		ALVOGEN	1GM	N016151	001	
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METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

AB		BIOPHARM	5MG	A040547	001	Feb 18, 2005
AB			10MG	A040547	002	Feb 18, 2005
AB		CHARTWELL RX	5MG	A040411	001	Mar 27, 2001
AB			10MG	A040411	002	Mar 27, 2001
AB		HERITAGE PHARMA	5MG	A040734	001	Dec 14, 2007
AB			10MG	A040734	002	Dec 14, 2007
AB		MYLAN	5MG	A040350	001	Mar 29, 2000
AB	!		10MG	A040350	002	Mar 29, 2000
AB		RISING	5MG	A202068	001	Mar 07, 2012
AB			10MG	A202068	002	Mar 07, 2012

METHOCARBAMOL

SOLUTION; IM-IV

METHOCARBAMOL

AP		AM REGENT	1GM/10ML (100MG/ML)	A207496	001	Jun 22, 2017
AP		EUGIA PHARMA	1GM/10ML (100MG/ML)	A206128	001	May 27, 2016
AP		FRESENIUS KABI USA	1GM/10ML (100MG/ML)	A209331	001	Apr 17, 2018
AP		GLAND PHARMA LTD	1GM/10ML (100MG/ML)	A211504	001	Oct 26, 2018
AP		MONTEREY PHARMS LLC	1GM/10ML (100MG/ML)	A205354	001	Oct 27, 2016
AP		MYLAN INSTITUTIONAL	1GM/10ML (100MG/ML)	A204404	001	Dec 05, 2014
AP		NAVINTA LLC	1GM/10ML (100MG/ML)	A206071	001	Nov 24, 2017
AP		SAGENT PHARMS INC	1GM/10ML (100MG/ML)	A205404	001	Jul 18, 2017
AP		SLATE	1GM/10ML (100MG/ML)	A208116	001	Jan 19, 2017
AP		SOMERSET THERAPS LLC	1GM/10ML (100MG/ML)	A207522	001	Jul 31, 2017

ROBAXIN

AP	+	HIKMA	1GM/10ML (100MG/ML)	N011790	001	
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TABLET; ORAL

METHOCARBAMOL

AA		AUSTARPHARMA LLC	500MG	A200958	001	Oct 21, 2011
AA			750MG	A200958	002	Oct 21, 2011
AA		BEXIMCO PHARMS USA	500MG	A208507	001	Jul 21, 2017
AA			750MG	A208507	002	Jul 21, 2017
AA		DBL PHARMS	500MG	A203550	001	Feb 08, 2017
AA			750MG	A203550	002	Feb 08, 2017
AA	!	GRANULES	500MG	A209312	001	May 07, 2018
AA	!		750MG	A209312	002	May 07, 2018
AA		HETERO LABS LTD III	500MG	A090200	001	Nov 06, 2009
AA			750MG	A090200	002	Nov 06, 2009
AA		MLV	500MG	A212623	001	Apr 30, 2021
AA			750MG	A212623	002	Apr 30, 2021
AA		OXFORD PHARMS	500MG	A040489	001	Jan 29, 2003

PRESCRIPTION DRUG PRODUCT LIST

METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL

<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>	
!	AUSTARPHARMA LLC	1GM	A200958	003	Dec 06, 2021

METHOHEXITAL SODIUM

INJECTABLE; INJECTION

BREVITAL SODIUM

+	!	PAR STERILE PRODUCTS	500MG/VIAL	N011559	001	
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METHOTREXATE

SOLUTION; SUBCUTANEOUS

OTREXUP

+	!	OTTER PHARMS	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

+	!	MEDEXUS	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

REDITREX

+	!	CUMBERLAND PHARMS	7.5MG/0.3ML (7.5MG/0.3ML)	N210737	001	Nov 27, 2019
+	!		10MG/0.4ML (10MG/0.4ML)	N210737	002	Nov 27, 2019
+	!		12.5MG/0.5ML (12.5MG/0.5ML)	N210737	003	Nov 27, 2019
+	!		15MG/0.6ML (15MG/0.6ML)	N210737	004	Nov 27, 2019
+	!		17.5MG/0.7ML (17.5MG/0.7ML)	N210737	005	Nov 27, 2019
+	!		20MG/0.8ML (20MG/0.8ML)	N210737	006	Nov 27, 2019
+	!		22.5MG/ML (22.5MG/ML)	N210737	007	Nov 27, 2019
+	!		25MG/1ML (25MG/1ML)	N210737	008	Nov 27, 2019

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE PRESERVATIVE FREE

<u>AP</u>	!	FRESENIUS KABI USA	<u>EQ 1GM BASE/VIAL</u>	<u>A040266</u>	<u>001</u>	Feb 26, 1999
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METHOTREXATE SODIUM

<u>AP</u>	!	FRESENIUS KABI USA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A040263</u>	<u>001</u>	Feb 26, 1999
<u>AP</u>	!	HIKMA	<u>EQ 100MG BASE/4ML (EQ 25MG BASE/ML)</u>	<u>A089341</u>	<u>001</u>	Sep 16, 1986
<u>AP</u>	+	HOSPIRA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>N011719</u>	<u>010</u>	Dec 15, 2004

METHOTREXATE SODIUM PRESERVATIVE FREE

<u>AP</u>	!	ACCORD HLTHCARE	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A040767</u>	<u>001</u>	Apr 30, 2007
<u>AP</u>	!		<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A040768</u>	<u>001</u>	Apr 30, 2007
<u>AP</u>	!		<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A040716</u>	<u>001</u>	Apr 30, 2007
<u>AP</u>	!	HIKMA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A089340</u>	<u>001</u>	Sep 16, 1986
<u>AP</u>	!		<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A089343</u>	<u>001</u>	Sep 16, 1986
<u>AP</u>	+	HOSPIRA	<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>N011719</u>	<u>012</u>	Apr 13, 2005
<u>AP</u>		PHARMACHEMIE BV	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A040843</u>	<u>002</u>	Jan 11, 2010
<u>AP</u>			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A040843</u>	<u>004</u>	Jan 11, 2010
<u>AP</u>			<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A040843</u>	<u>001</u>	Jan 11, 2010
<u>AP</u>		SAGENT PHARMS INC	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A203407</u>	<u>001</u>	Aug 09, 2018
<u>AP</u>			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A203407</u>	<u>002</u>	Aug 09, 2018
<u>AP</u>			<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A203407</u>	<u>003</u>	Aug 09, 2018
<u>AP</u>		SANDOZ INC	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A090039</u>	<u>001</u>	Mar 31, 2009
<u>AP</u>			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A090039</u>	<u>002</u>	Mar 31, 2009
<u>AP</u>			<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A090029</u>	<u>001</u>	Mar 31, 2009

METHOTREXATE SODIUM

!	!	HIKMA	EQ 200MG BASE/8ML (EQ 25MG BASE/ML)	A089342	001	Sep 16, 1986
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METHOTREXATE SODIUM PRESERVATIVE FREE

!	!	HIKMA	EQ 1GM BASE/VIAL	A040632	001	Aug 12, 2005
		PHARMACHEMIE BV	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	A040843	003	Feb 27, 2012

PRESCRIPTION DRUG PRODUCT LIST

METHOTREXATE SODIUM

SOLUTION;ORAL

XATMEP

+! AZURITY

EQ 2.5MG BASE/ML

N208400 001 Apr 25, 2017

TABLET;ORAL

METHOTREXATE SODIUM

AB	ACCORD HLTHCARE	<u>EQ 2.5MG BASE</u>	<u>A213343 001</u>	Jan 24, 2020
AB	AMNEAL PHARMS	<u>EQ 2.5MG BASE</u>	<u>A210040 001</u>	Dec 22, 2017
AB	BARR	<u>EQ 2.5MG BASE</u>	<u>A081099 001</u>	Oct 15, 1990
AB	EUGIA PHARMA	<u>EQ 2.5MG BASE</u>	<u>A210454 001</u>	Jan 30, 2020
AB	! HIKMA	<u>EQ 2.5MG BASE</u>	<u>A040054 001</u>	Aug 01, 1994
AB	LOTUS PHARM CO LTD	<u>EQ 2.5MG BASE</u>	<u>A209787 001</u>	Apr 23, 2021
AB	MYLAN	<u>EQ 2.5MG BASE</u>	<u>A081235 001</u>	May 15, 1992
AB	SUN PHARM	<u>EQ 2.5MG BASE</u>	<u>A201749 001</u>	May 21, 2015
AB	ZYDUS PHARMS	<u>EQ 2.5MG BASE</u>	<u>A207812 001</u>	Jan 13, 2017
	TREXALL			
	BARR	EQ 5MG BASE	A040385 001	Mar 21, 2001
		EQ 7.5MG BASE	A040385 002	Mar 21, 2001
		EQ 10MG BASE	A040385 003	Mar 21, 2001
	!	EQ 15MG BASE	A040385 004	Mar 21, 2001

METHOXSALEN

CAPSULE;ORAL

METHOXSALEN

AB	STRIDES PHARMA	<u>10MG</u>	<u>A202687 001</u>	Jun 05, 2014
	<u>OXSORALEN-ULTRA</u>			
AB	+! BAUSCH	<u>10MG</u>	<u>N019600 001</u>	Oct 30, 1986
	INJECTABLE;INJECTION			
	UVADEX			
	+! MALLINCKRODT HOSP	0.02MG/ML	N020969 001	Feb 25, 1999

METHSCOPOLAMINE BROMIDE

TABLET;ORAL

METHSCOPOLAMINE BROMIDE

AA	BAYSHORE PHARMS LLC	<u>2.5MG</u>	<u>A200602 001</u>	Sep 24, 2012
AA	!	<u>5MG</u>	<u>A200602 002</u>	Sep 24, 2012
AA	BRECKENRIDGE PHARM	<u>2.5MG</u>	<u>A040642 001</u>	Dec 06, 2011
AA		<u>5MG</u>	<u>A040642 002</u>	Dec 06, 2011

METHSUXIMIDE

CAPSULE;ORAL

CELONTIN

+! PARKE DAVIS

300MG

N010596 008

METHYLDOPA

TABLET;ORAL

METHYLDOPA

+! ACCORD HLTHCARE

250MG

A070084 001 Oct 15, 1985

!

A070085 001 Oct 15, 1985

METHYLENE BLUE

SOLUTION;INTRAVENOUS

PROVAYBLUE

+! PROVEPHARM SAS

10MG/2ML (5MG/ML)

N204630 002 Jul 18, 2019

+!

50MG/10ML (5MG/ML)

N204630 001 Apr 08, 2016

METHYLERGONOVINE MALEATE

INJECTABLE;INJECTION

METHERGINE

AP	+! EDISON THERAPS LLC	<u>0.2MG/ML</u>	<u>N006035 004</u>	
	<u>METHYLERGONOVINE MALEATE</u>			
AP	AM REGENT	<u>0.2MG/ML</u>	<u>A090193 001</u>	Nov 24, 2008
AP	BRECKENRIDGE	<u>0.2MG/ML</u>	<u>A040889 001</u>	Sep 13, 2010
	TABLET;ORAL			
	<u>METHYLERGONOVINE MALEATE</u>			
AB	AMNEAL PHARMS	<u>0.2MG</u>	<u>A211483 001</u>	Sep 10, 2018
AB	GRANULES	<u>0.2MG</u>	<u>A210424 001</u>	May 15, 2018
AB	! NOVEL LABS INC	<u>0.2MG</u>	<u>A091577 001</u>	May 02, 2011
AB	RISING	<u>0.2MG</u>	<u>A211919 001</u>	Jan 15, 2021
AB	TEVA PHARMS USA	<u>0.2MG</u>	<u>A211455 001</u>	Mar 20, 2019
AB	TULEX PHARMS INC	<u>0.2MG</u>	<u>A212233 001</u>	May 01, 2020

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	150MG	N208271	001	Jul 19, 2016
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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>		GRANULES	<u>10MG</u>	<u>A211796</u>	<u>001</u>	May 23, 2019
<u>AB1</u>			<u>20MG</u>	<u>A211796</u>	<u>002</u>	May 23, 2019
<u>AB1</u>			<u>30MG</u>	<u>A211796</u>	<u>003</u>	May 23, 2019
<u>AB1</u>			<u>40MG</u>	<u>A211796</u>	<u>004</u>	May 23, 2019
<u>AB1</u>			<u>60MG</u>	<u>A211796</u>	<u>005</u>	May 23, 2019
<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
<u>AB1</u>	!		<u>60MG</u>	<u>A078458</u>	<u>004</u>	Jun 23, 2016

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

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

APTENSIO XR

<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

METHYLPHENIDATE HYDROCHLORIDE

<u>AB3</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A208861</u>	<u>001</u>	Dec 13, 2018
<u>AB3</u>		<u>15MG</u>	<u>A208861</u>	<u>002</u>	Dec 13, 2018
<u>AB3</u>		<u>20MG</u>	<u>A208861</u>	<u>003</u>	Dec 13, 2018
<u>AB3</u>		<u>30MG</u>	<u>A208861</u>	<u>004</u>	Dec 13, 2018
<u>AB3</u>		<u>40MG</u>	<u>A208861</u>	<u>005</u>	Dec 13, 2018
<u>AB3</u>		<u>50MG</u>	<u>A208861</u>	<u>006</u>	Dec 13, 2018
<u>AB3</u>		<u>60MG</u>	<u>A208861</u>	<u>007</u>	Dec 13, 2018

ADHANSIA XR

+	PURDUE PHARMA LP	25MG	N212038	001	Feb 27, 2019
+		35MG	N212038	002	Feb 27, 2019
+		45MG	N212038	003	Feb 27, 2019
+		55MG	N212038	004	Feb 27, 2019
+		70MG	N212038	005	Feb 27, 2019
+!		85MG	N212038	006	Feb 27, 2019

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

FOR SUSPENSION, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>5MG/ML</u>	<u>A206049</u>	<u>001</u>	May 17, 2018
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QUILLIVANT XR

<u>AB</u>	+! NEXTWAVE	<u>5MG/ML</u>	<u>N202100</u>	<u>001</u>	Sep 27, 2012
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SOLUTION;ORAL

METHYLIN

<u>AA</u>	+! SPECGX LLC	<u>5MG/5ML</u>	<u>N021419</u>	<u>001</u>	Dec 19, 2002
<u>AA</u>	+!	<u>10MG/5ML</u>	<u>N021419</u>	<u>002</u>	Dec 19, 2002

METHYLPHENIDATE HYDROCHLORIDE

<u>AA</u>	ABHAI LLC	<u>5MG/5ML</u>	<u>A207485</u>	<u>001</u>	Nov 18, 2016
<u>AA</u>		<u>10MG/5ML</u>	<u>A207485</u>	<u>002</u>	Nov 18, 2016
<u>AA</u>	ALKEM LABS LTD	<u>5MG/5ML</u>	<u>A211647</u>	<u>001</u>	Mar 30, 2020
<u>AA</u>		<u>10MG/5ML</u>	<u>A211647</u>	<u>002</u>	Mar 30, 2020
<u>AA</u>	ANDA REPOSITORY	<u>5MG/5ML</u>	<u>A210764</u>	<u>001</u>	Apr 10, 2020
<u>AA</u>		<u>10MG/5ML</u>	<u>A210764</u>	<u>002</u>	Apr 10, 2020
<u>AA</u>	ASCENT PHARMS INC	<u>5MG/5ML</u>	<u>A207417</u>	<u>001</u>	Jan 29, 2021
<u>AA</u>		<u>10MG/5ML</u>	<u>A207417</u>	<u>002</u>	Jan 29, 2021
<u>AA</u>	BRECKENRIDGE	<u>5MG/5ML</u>	<u>A201466</u>	<u>001</u>	Nov 12, 2013
<u>AA</u>		<u>10MG/5ML</u>	<u>A201466</u>	<u>002</u>	Nov 12, 2013
<u>AA</u>	EYWA	<u>5MG/5ML</u>	<u>A210139</u>	<u>001</u>	Oct 03, 2018
<u>AA</u>		<u>10MG/5ML</u>	<u>A210139</u>	<u>002</u>	Oct 03, 2018
<u>AA</u>	NOVEL LABS INC	<u>5MG/5ML</u>	<u>A204602</u>	<u>001</u>	Aug 14, 2015
<u>AA</u>		<u>10MG/5ML</u>	<u>A204602</u>	<u>002</u>	Aug 14, 2015
<u>AA</u>	QUAGEN	<u>5MG/5ML</u>	<u>A213567</u>	<u>001</u>	Jun 04, 2020
<u>AA</u>		<u>10MG/5ML</u>	<u>A213567</u>	<u>002</u>	Jun 04, 2020
<u>AA</u>	TRIS PHARMA INC	<u>5MG/5ML</u>	<u>A091601</u>	<u>001</u>	Jul 23, 2010
<u>AA</u>		<u>10MG/5ML</u>	<u>A091601</u>	<u>002</u>	Jul 23, 2010

TABLET;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	ABHAI INC	<u>5MG</u>	<u>A206932</u>	<u>001</u>	May 11, 2017
<u>AB</u>		<u>10MG</u>	<u>A206932</u>	<u>002</u>	May 11, 2017
<u>AB</u>		<u>20MG</u>	<u>A206932</u>	<u>003</u>	May 11, 2017
<u>AB</u>	ACCORD HLTHCARE	<u>5MG</u>	<u>A213936</u>	<u>001</u>	Oct 28, 2020
<u>AB</u>		<u>10MG</u>	<u>A213936</u>	<u>002</u>	Oct 28, 2020
<u>AB</u>		<u>20MG</u>	<u>A213936</u>	<u>003</u>	Oct 28, 2020
<u>AB</u>	ALKEM LABS LTD	<u>5MG</u>	<u>A211779</u>	<u>001</u>	Oct 04, 2019
<u>AB</u>		<u>10MG</u>	<u>A211779</u>	<u>002</u>	Oct 04, 2019
<u>AB</u>		<u>20MG</u>	<u>A211779</u>	<u>003</u>	Oct 04, 2019
<u>AB</u>	ASCENT PHARMS INC	<u>5MG</u>	<u>A207416</u>	<u>001</u>	Sep 22, 2015
<u>AB</u>		<u>10MG</u>	<u>A207416</u>	<u>002</u>	Sep 22, 2015
<u>AB</u>		<u>20MG</u>	<u>A207416</u>	<u>003</u>	Sep 22, 2015
<u>AB</u>	BIONPHARMA INC	<u>5MG</u>	<u>A209753</u>	<u>001</u>	Mar 02, 2018
<u>AB</u>		<u>10MG</u>	<u>A209753</u>	<u>002</u>	Mar 02, 2018
<u>AB</u>		<u>20MG</u>	<u>A209753</u>	<u>003</u>	Mar 02, 2018
<u>AB</u>	MOUNTAIN	<u>5MG</u>	<u>A091159</u>	<u>001</u>	Mar 12, 2014

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

TABLET; ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>		<u>10MG</u>	<u>A091159</u>	<u>002</u>	Mar 12, 2014
<u>AB</u>		<u>20MG</u>	<u>A091159</u>	<u>003</u>	Mar 12, 2014
<u>AB</u>	NOVEL LABS INC	<u>5MG</u>	<u>A207884</u>	<u>001</u>	Nov 13, 2015
<u>AB</u>		<u>10MG</u>	<u>A207884</u>	<u>002</u>	Nov 13, 2015
<u>AB</u>		<u>20MG</u>	<u>A207884</u>	<u>003</u>	Nov 13, 2015
<u>AB</u>	OXFORD PHARMS	<u>5MG</u>	<u>A202892</u>	<u>001</u>	Sep 23, 2014
<u>AB</u>		<u>10MG</u>	<u>A202892</u>	<u>002</u>	Sep 23, 2014
<u>AB</u>		<u>20MG</u>	<u>A202892</u>	<u>003</u>	Sep 23, 2014
<u>AB</u>	PRINSTON INC	<u>5MG</u>	<u>A212697</u>	<u>001</u>	Jul 23, 2020
<u>AB</u>		<u>10MG</u>	<u>A212697</u>	<u>002</u>	Jul 23, 2020
<u>AB</u>		<u>20MG</u>	<u>A212697</u>	<u>003</u>	Jul 23, 2020
<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>	RISING	<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

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>	ALKEM LABS LTD	<u>20MG</u>	<u>A212288</u>	<u>001</u>	Oct 06, 2020
<u>AB</u>	ALVOGEN	<u>10MG</u>	<u>A204772</u>	<u>001</u>	Feb 29, 2016
<u>AB</u>		<u>18MG</u>	<u>A210818</u>	<u>001</u>	Nov 30, 2018
<u>AB</u>		<u>20MG</u>	<u>A204772</u>	<u>002</u>	Feb 29, 2016
<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>	ANDOR PHARMS	<u>18MG</u>	<u>A211918</u>	<u>001</u>	Apr 24, 2019
<u>AB</u>		<u>27MG</u>	<u>A211918</u>	<u>002</u>	Apr 24, 2019
<u>AB</u>		<u>36MG</u>	<u>A211918</u>	<u>003</u>	Apr 24, 2019
<u>AB</u>		<u>54MG</u>	<u>A211918</u>	<u>004</u>	Apr 24, 2019
<u>AB</u>	ASCENT PHARMS INC	<u>18MG</u>	<u>A211009</u>	<u>001</u>	Sep 03, 2019
<u>AB</u>		<u>27MG</u>	<u>A211009</u>	<u>002</u>	Sep 03, 2019
<u>AB</u>		<u>36MG</u>	<u>A211009</u>	<u>003</u>	Sep 03, 2019
<u>AB</u>		<u>54MG</u>	<u>A211009</u>	<u>004</u>	Sep 03, 2019
<u>AB</u>	AUROLIFE PHARMA LLC	<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>	DR REDDYS	<u>18MG</u>	<u>A213473</u>	<u>001</u>	Jul 29, 2020
<u>AB</u>		<u>27MG</u>	<u>A213473</u>	<u>002</u>	Jul 29, 2020
<u>AB</u>		<u>36MG</u>	<u>A213473</u>	<u>003</u>	Jul 29, 2020
<u>AB</u>		<u>54MG</u>	<u>A213473</u>	<u>004</u>	Jul 29, 2020
<u>AB</u>	GRANULES	<u>10MG</u>	<u>A210992</u>	<u>001</u>	Nov 21, 2018

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>		<u>20MG</u>	<u>A210992</u>	<u>002</u>	Nov 21, 2018
<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
<u>AB</u>	SUN PHARM INDS INC	<u>18MG</u>	<u>A205135</u>	<u>001</u>	Aug 19, 2020
<u>AB</u>		<u>27MG</u>	<u>A205135</u>	<u>002</u>	Aug 19, 2020
<u>AB</u>		<u>36MG</u>	<u>A205135</u>	<u>003</u>	Aug 19, 2020
<u>AB</u>		<u>54MG</u>	<u>A205135</u>	<u>004</u>	Aug 19, 2020
BX	AMNEAL PHARMS	18MG	A207515	001	Feb 01, 2018
BX		27MG	A207515	002	Feb 01, 2018
BX		36MG	A207515	003	Feb 01, 2018
BX		54MG	A207515	004	Feb 01, 2018
BX	LANNETT CO INC	18MG	A091695	001	Jul 09, 2013
BX		27MG	A091695	002	Jul 09, 2013
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

<u>AB</u>	+ PFIZER	<u>2MG</u>	<u>N011153</u>	<u>002</u>	
<u>AB</u>	+	<u>4MG</u>	<u>N011153</u>	<u>001</u>	
<u>AB</u>	+	<u>8MG</u>	<u>N011153</u>	<u>004</u>	
<u>AB</u>	+	<u>16MG</u>	<u>N011153</u>	<u>003</u>	
<u>AB</u>	+!	<u>32MG</u>	<u>N011153</u>	<u>006</u>	

METHYLPREDNISOLONE

<u>AB</u>	AMNEAL	<u>4MG</u>	<u>A207481</u>	<u>001</u>	Sep 21, 2021
<u>AB</u>		<u>8MG</u>	<u>A207481</u>	<u>002</u>	Sep 21, 2021
<u>AB</u>		<u>16MG</u>	<u>A207481</u>	<u>003</u>	Sep 21, 2021
<u>AB</u>		<u>32MG</u>	<u>A207481</u>	<u>004</u>	Sep 21, 2021
<u>AB</u>	JUBILANT CADISTA	<u>4MG</u>	<u>A040189</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>		<u>8MG</u>	<u>A040189</u>	<u>002</u>	Oct 31, 1997
<u>AB</u>		<u>16MG</u>	<u>A040189</u>	<u>003</u>	Jul 20, 2007
<u>AB</u>		<u>32MG</u>	<u>A040189</u>	<u>004</u>	Jul 20, 2007
<u>AB</u>	PRAXGEN	<u>4MG</u>	<u>A212262</u>	<u>001</u>	Jun 27, 2019
<u>AB</u>	SANDOZ	<u>4MG</u>	<u>A040194</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>	TIANJIN TIANYAO	<u>4MG</u>	<u>A204072</u>	<u>001</u>	May 14, 2018
<u>AB</u>	VINTAGE PHARMS	<u>4MG</u>	<u>A040183</u>	<u>001</u>	Dec 22, 1998
<u>AB</u>	WATSON LABS	<u>4MG</u>	<u>A040232</u>	<u>001</u>	Oct 16, 1997
<u>AB</u>	ZYDUS PHARMS	<u>4MG</u>	<u>A206751</u>	<u>001</u>	Apr 23, 2018
<u>AB</u>		<u>8MG</u>	<u>A206751</u>	<u>002</u>	Apr 23, 2018
<u>AB</u>		<u>16MG</u>	<u>A206751</u>	<u>003</u>	Apr 23, 2018
<u>AB</u>		<u>32MG</u>	<u>A206751</u>	<u>004</u>	Apr 23, 2018

METHYLPREDNISOLONE ACETATE

INJECTABLE;INJECTION

DEPO-MEDROL

<u>AB</u>	+! PFIZER	<u>20MG/ML</u>	<u>N011757</u>	<u>002</u>	
<u>AB</u>	+!	<u>40MG/ML</u>	<u>N011757</u>	<u>001</u>	
<u>AB</u>	+!	<u>80MG/ML</u>	<u>N011757</u>	<u>004</u>	

METHYLPREDNISOLONE ACETATE

<u>AB</u>	AMNEAL	<u>40MG/ML</u>	<u>A210043</u>	<u>001</u>	May 20, 2019
<u>AB</u>		<u>80MG/ML</u>	<u>A210043</u>	<u>002</u>	May 20, 2019
<u>AB</u>	SAGENT PHARMS INC	<u>20MG/ML</u>	<u>A201835</u>	<u>001</u>	Jun 27, 2018
<u>AB</u>		<u>40MG/ML</u>	<u>A201835</u>	<u>002</u>	Jun 27, 2018
<u>AB</u>		<u>80MG/ML</u>	<u>A201835</u>	<u>003</u>	Jun 27, 2018
<u>AB</u>	SANDOZ INC	<u>40MG/ML</u>	<u>A040719</u>	<u>001</u>	Jan 29, 2009
<u>AB</u>		<u>40MG/ML</u>	<u>A040794</u>	<u>001</u>	Mar 05, 2009
<u>AB</u>		<u>80MG/ML</u>	<u>A040719</u>	<u>002</u>	Jan 29, 2009
<u>AB</u>		<u>80MG/ML</u>	<u>A040794</u>	<u>002</u>	Mar 05, 2009

PRESCRIPTION DRUG PRODUCT LIST

METOCLOPRAMIDE HYDROCHLORIDE

TABLET;ORAL

REGLAN

AB	+	ANI PHARMS	<u>EQ 5MG BASE</u>	<u>N017854</u>	<u>002</u>	May 05, 1987
AB	+	!	<u>EQ 10MG BASE</u>	<u>N017854</u>	<u>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

METOLAZONE

TABLET;ORAL

METOLAZONE

AB		ALEMBIC PHARMS LTD	<u>2.5MG</u>	<u>A213251</u>	<u>001</u>	Dec 02, 2020
AB			<u>5MG</u>	<u>A213251</u>	<u>002</u>	Dec 02, 2020
AB			<u>10MG</u>	<u>A213251</u>	<u>003</u>	Dec 02, 2020
AB		BAYSHORE PHARMS LLC	<u>2.5MG</u>	<u>A214799</u>	<u>001</u>	Mar 30, 2021
AB			<u>5MG</u>	<u>A214799</u>	<u>002</u>	Mar 30, 2021
AB			<u>10MG</u>	<u>A214799</u>	<u>003</u>	Mar 30, 2021
AB		INNOGENIX	<u>2.5MG</u>	<u>A213827</u>	<u>001</u>	Mar 30, 2021
AB			<u>5MG</u>	<u>A213827</u>	<u>002</u>	Mar 30, 2021
AB			<u>10MG</u>	<u>A213827</u>	<u>003</u>	Mar 30, 2021
AB		MYLAN	<u>2.5MG</u>	<u>A076698</u>	<u>001</u>	Dec 23, 2003
AB			<u>5MG</u>	<u>A076698</u>	<u>002</u>	Oct 19, 2004
AB			<u>10MG</u>	<u>A076698</u>	<u>003</u>	Oct 19, 2004
AB		RUBICON	<u>2.5MG</u>	<u>A215184</u>	<u>001</u>	Aug 20, 2021
AB			<u>5MG</u>	<u>A215184</u>	<u>002</u>	Aug 20, 2021
AB			<u>10MG</u>	<u>A215184</u>	<u>003</u>	Aug 20, 2021
AB		SANDOZ	<u>2.5MG</u>	<u>A076732</u>	<u>001</u>	Dec 19, 2003
AB		!	<u>5MG</u>	<u>A076466</u>	<u>001</u>	Dec 19, 2003
AB		!	<u>10MG</u>	<u>A076466</u>	<u>002</u>	Dec 19, 2003

METOPROLOL SUCCINATE

CAPSULE, EXTENDED RELEASE;ORAL

KAPSPARGO SPRINKLE

	+	SPII	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

AB		ACTAVIS ELIZABETH	<u>EQ 25MG TARTRATE</u>	<u>A204161</u>	<u>001</u>	Nov 25, 2016
AB			<u>EQ 50MG TARTRATE</u>	<u>A204161</u>	<u>002</u>	Nov 25, 2016
AB			<u>EQ 100MG TARTRATE</u>	<u>A204161</u>	<u>003</u>	Nov 25, 2016
AB			<u>EQ 200MG TARTRATE</u>	<u>A204161</u>	<u>004</u>	Nov 25, 2016
AB		ACTAVIS LABS FL INC	<u>EQ 50MG TARTRATE</u>	<u>A076862</u>	<u>001</u>	Aug 03, 2009
AB		ALKEM LABS LTD	<u>EQ 25MG TARTRATE</u>	<u>A211143</u>	<u>001</u>	Nov 25, 2020
AB			<u>EQ 50MG TARTRATE</u>	<u>A211143</u>	<u>002</u>	Nov 25, 2020
AB			<u>EQ 100MG TARTRATE</u>	<u>A211143</u>	<u>003</u>	Nov 25, 2020
AB			<u>EQ 200MG TARTRATE</u>	<u>A211143</u>	<u>004</u>	Nov 25, 2020
AB		CIPLA	<u>EQ 50MG TARTRATE</u>	<u>A207465</u>	<u>001</u>	Oct 26, 2018
AB			<u>EQ 100MG TARTRATE</u>	<u>A207465</u>	<u>002</u>	Oct 26, 2018
AB			<u>EQ 200MG TARTRATE</u>	<u>A207465</u>	<u>003</u>	Oct 26, 2018
AB		DR REDDYS LABS LTD	<u>EQ 25MG TARTRATE</u>	<u>A090617</u>	<u>001</u>	Aug 01, 2012
AB			<u>EQ 50MG TARTRATE</u>	<u>A090617</u>	<u>002</u>	Aug 01, 2012
AB		HETERO LABS LTD III	<u>EQ 25MG TARTRATE</u>	<u>A205541</u>	<u>001</u>	Nov 06, 2020
AB			<u>EQ 50MG TARTRATE</u>	<u>A205541</u>	<u>002</u>	Nov 06, 2020
AB			<u>EQ 100MG TARTRATE</u>	<u>A205541</u>	<u>003</u>	Nov 06, 2020
AB			<u>EQ 200MG TARTRATE</u>	<u>A205541</u>	<u>004</u>	Nov 06, 2020
AB		MYLAN PHARMS INC	<u>EQ 25MG TARTRATE</u>	<u>A202033</u>	<u>001</u>	Dec 15, 2011
AB			<u>EQ 50MG TARTRATE</u>	<u>A202033</u>	<u>002</u>	Dec 15, 2011
AB			<u>EQ 100MG TARTRATE</u>	<u>A202033</u>	<u>003</u>	Dec 15, 2011
AB			<u>EQ 200MG TARTRATE</u>	<u>A202033</u>	<u>004</u>	Dec 15, 2011
AB		NOVAST LABS	<u>EQ 25MG TARTRATE</u>	<u>A204106</u>	<u>001</u>	Feb 06, 2018
AB			<u>EQ 50MG TARTRATE</u>	<u>A204106</u>	<u>002</u>	Feb 06, 2018
AB			<u>EQ 100MG TARTRATE</u>	<u>A204106</u>	<u>003</u>	Feb 06, 2018
AB			<u>EQ 200MG TARTRATE</u>	<u>A204106</u>	<u>004</u>	Feb 06, 2018
AB		PHARMADAX INC	<u>EQ 25MG TARTRATE</u>	<u>A203028</u>	<u>001</u>	Mar 31, 2020
AB			<u>EQ 50MG TARTRATE</u>	<u>A203028</u>	<u>002</u>	Mar 31, 2020
AB			<u>EQ 100MG TARTRATE</u>	<u>A203699</u>	<u>001</u>	Mar 30, 2020
AB			<u>EQ 200MG TARTRATE</u>	<u>A203699</u>	<u>002</u>	Mar 30, 2020
AB		PRINSTON INC	<u>EQ 100MG TARTRATE</u>	<u>A210597</u>	<u>001</u>	Jan 04, 2022
AB			<u>EQ 200MG TARTRATE</u>	<u>A210597</u>	<u>002</u>	Jan 04, 2022

PRESCRIPTION DRUG PRODUCT LIST

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE;ORAL

METOPROLOL SUCCINATE

<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>	VISUM PHARM	<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>	YICHANG HUMANWELL	<u>EQ 50MG TARTRATE</u>	<u>A213854 003</u>	Nov 08, 2021
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A213854 001</u>	Feb 12, 2021
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A213854 002</u>	Feb 12, 2021
<u>AB</u>	ZYDUS PHARMS	<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
<u>TOPROL-XL</u>				
<u>AB</u>	+ TOPROL	<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

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
<u>AP</u>	HIKMA	<u>1MG/ML</u>	<u>A076495 001</u>	Jul 07, 2003
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A077761 001</u>	May 30, 2007
<u>AP</u>	HOSPIRA	<u>1MG/ML</u>	<u>A074133 001</u>	Dec 21, 1993
<u>AP</u>		<u>1MG/ML</u>	<u>A075160 001</u>	Jul 06, 1998
<u>AP</u>	!	<u>1MG/ML</u>	<u>A078085 001</u>	Apr 29, 2008
<u>AP</u>	SANDOZ INC	<u>1MG/ML</u>	<u>A077360 001</u>	Oct 02, 2007

TABLET;ORAL

LOPRESSOR

<u>AB</u>	+ VALIDUS PHARMS	<u>50MG</u>	<u>N017963 001</u>	
<u>AB</u>	+	<u>100MG</u>	<u>N017963 002</u>	

METOPROLOL TARTRATE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A202871 001</u>	May 28, 2013
<u>AB</u>		<u>50MG</u>	<u>A202871 002</u>	May 28, 2013
<u>AB</u>		<u>100MG</u>	<u>A202871 003</u>	May 28, 2013
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A077739 001</u>	Sep 11, 2007
<u>AB</u>		<u>50MG</u>	<u>A077739 002</u>	Sep 11, 2007
<u>AB</u>		<u>100MG</u>	<u>A077739 003</u>	Sep 11, 2007
<u>AB</u>	IPCA LABS LTD	<u>25MG</u>	<u>A078459 001</u>	Jun 17, 2008
<u>AB</u>		<u>50MG</u>	<u>A078459 002</u>	Jun 17, 2008
<u>AB</u>		<u>100MG</u>	<u>A078459 003</u>	Jun 17, 2008
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A076704 001</u>	Jan 16, 2004
<u>AB</u>		<u>37.5MG</u>	<u>A076704 004</u>	Mar 18, 2015
<u>AB</u>		<u>50MG</u>	<u>A076704 002</u>	Jan 16, 2004
<u>AB</u>		<u>75MG</u>	<u>A076704 005</u>	Mar 18, 2015
<u>AB</u>	!	<u>100MG</u>	<u>A076704 003</u>	Jan 16, 2004
<u>AB</u>	RUBICON	<u>25MG</u>	<u>A200981 001</u>	Oct 28, 2014
<u>AB</u>		<u>37.5MG</u>	<u>A200981 004</u>	Aug 21, 2019
<u>AB</u>		<u>50MG</u>	<u>A200981 002</u>	Oct 28, 2014
<u>AB</u>		<u>75MG</u>	<u>A200981 005</u>	Aug 21, 2019
<u>AB</u>		<u>100MG</u>	<u>A200981 003</u>	Oct 28, 2014
<u>AB</u>	SUN PHARM INDS INC	<u>25MG</u>	<u>A076670 001</u>	Jan 15, 2004
<u>AB</u>		<u>50MG</u>	<u>A074644 001</u>	Dec 10, 1996
<u>AB</u>		<u>100MG</u>	<u>A074644 002</u>	Dec 10, 1996
<u>AB</u>	YOUNGTECH PHARMS INC	<u>25MG</u>	<u>A208955 001</u>	Feb 05, 2020
<u>AB</u>		<u>50MG</u>	<u>A208955 002</u>	Feb 05, 2020
<u>AB</u>		<u>100MG</u>	<u>A208955 003</u>	Feb 05, 2020

PRESCRIPTION DRUG PRODUCT LIST

METRONIDAZOLE

CAPSULE; ORAL

FLAGYL

<u>AB</u>	<u>+</u> !	PFIZER	<u>375MG</u>	<u>N020334</u>	<u>001</u>	May 03, 1995
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METRONIDAZOLE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>375MG</u>	<u>A079065</u>	<u>001</u>	Jun 23, 2009
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<u>AB</u>		STRIDES PHARMA	<u>375MG</u>	<u>A076522</u>	<u>001</u>	Jan 29, 2004
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CREAM; TOPICAL

METROCREAM

<u>AB</u>	<u>+</u> !	GALDERMA LABS LP	<u>0.75%</u>	<u>N020531</u>	<u>001</u>	Sep 20, 1995
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METRONIDAZOLE

<u>AB</u>		COSETTE	<u>0.75%</u>	<u>A077549</u>	<u>001</u>	Dec 19, 2007
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<u>AB</u>		FOUGERA PHARMS	<u>0.75%</u>	<u>A076408</u>	<u>001</u>	May 28, 2004
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NORITATE

<u>+</u> !		VALEANT PHARMS NORTH	1%	N020743	001	Sep 26, 1997
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GEL; TOPICAL

METROGEL

<u>AB</u>	<u>+</u> !	GALDERMA LABS LP	<u>0.75%</u>	<u>N019737</u>	<u>001</u>	Nov 22, 1988
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<u>AB</u>	<u>+</u> !		<u>1%</u>	<u>N021789</u>	<u>001</u>	Jun 30, 2005
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METRONIDAZOLE

<u>AB</u>		ALEOR DERMACEUTICALS	<u>1%</u>	<u>A212646</u>	<u>001</u>	Sep 03, 2021
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<u>AB</u>		COSETTE	<u>0.75%</u>	<u>A078178</u>	<u>001</u>	Jan 19, 2011
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<u>AB</u>		FOUGERA PHARMS	<u>0.75%</u>	<u>A077018</u>	<u>001</u>	Jun 06, 2006
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<u>AB</u>		TARO	<u>0.75%</u>	<u>A077819</u>	<u>001</u>	Jul 18, 2006
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<u>AB</u>			<u>1%</u>	<u>A204651</u>	<u>001</u>	Mar 14, 2017
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<u>AB</u>		TOLMAR	<u>0.75%</u>	<u>A077547</u>	<u>001</u>	Jul 13, 2006
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<u>AB</u>			<u>1%</u>	<u>A090903</u>	<u>001</u>	Jul 22, 2011
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GEL; VAGINAL

METROGEL-VAGINAL

<u>AB</u>	<u>+</u> !	BAUSCH	<u>0.75%</u>	<u>N020208</u>	<u>001</u>	Aug 17, 1992
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METRONIDAZOLE

<u>AB</u>		PADAGIS ISRAEL	<u>0.75%</u>	<u>A211786</u>	<u>001</u>	Jul 02, 2019
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<u>AB</u>		SOLARIS PHARMA CORP	<u>0.75%</u>	<u>A213648</u>	<u>001</u>	Oct 14, 2021
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<u>AB</u>		TOLMAR	<u>0.75%</u>	<u>A077264</u>	<u>001</u>	Oct 31, 2006
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VANDAZOLE

BX	!	TEVA PHARMS	0.75%	N021806	001	May 20, 2005
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NUVESSA

<u>+</u> !		CHEMO RESEARCH SL	1.3%	N205223	001	Mar 24, 2014
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INJECTABLE; INJECTION

FLAGYL I.V. RTU IN PLASTIC CONTAINER

<u>AP</u>	<u>+</u> !	BAXTER HLTHCARE	<u>500MG/100ML</u>	<u>N018657</u>	<u>001</u>	
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METRO I.V. IN PLASTIC CONTAINER

<u>AP</u>	<u>+</u> !	B BRAUN	<u>500MG/100ML</u>	<u>N018900</u>	<u>001</u>	Sep 29, 1983
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METRONIDAZOLE IN PLASTIC CONTAINER

<u>AP</u>		BAXTER HLTHCARE CORP	<u>500MG/100ML</u>	<u>A078084</u>	<u>001</u>	Mar 31, 2008
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<u>AP</u>		GLAND PHARMA LTD	<u>500MG/100ML</u>	<u>A212435</u>	<u>001</u>	Aug 03, 2020
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<u>AP</u>	<u>+</u> !	HOSPIRA	<u>500MG/100ML</u>	<u>N018890</u>	<u>002</u>	Nov 18, 1983
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<u>AP</u>		INFORLIFE	<u>500MG/100ML</u>	<u>A206191</u>	<u>001</u>	Feb 25, 2019
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LOTION; TOPICAL

METROLOTION

<u>AB</u>	<u>+</u> !	GALDERMA LABS LP	<u>0.75%</u>	<u>N020901</u>	<u>001</u>	Nov 24, 1998
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METRONIDAZOLE

<u>AB</u>		FOUGERA PHARMS	<u>0.75%</u>	<u>A077197</u>	<u>001</u>	May 24, 2006
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TABLET; ORAL

FLAGYL

<u>AB</u>	<u>+</u>	PFIZER	<u>250MG</u>	<u>N012623</u>	<u>001</u>	
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<u>AB</u>	<u>+</u> !		<u>500MG</u>	<u>N012623</u>	<u>003</u>	
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METRONIDAZOLE

<u>AB</u>		ALEMBIC LABS	<u>250MG</u>	<u>A208681</u>	<u>001</u>	Jun 20, 2017
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<u>AB</u>			<u>500MG</u>	<u>A208681</u>	<u>002</u>	Jun 20, 2017
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<u>AB</u>		ALEMBIC PHARMS LTD	<u>250MG</u>	<u>A079067</u>	<u>001</u>	Mar 13, 2009
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<u>AB</u>			<u>500MG</u>	<u>A079067</u>	<u>002</u>	Mar 13, 2009
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<u>AB</u>		AMICI	<u>250MG</u>	<u>A070772</u>	<u>001</u>	Jul 16, 1986
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<u>AB</u>			<u>500MG</u>	<u>A070772</u>	<u>002</u>	Jul 16, 1986
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<u>AB</u>		AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A203974</u>	<u>001</u>	May 29, 2015
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<u>AB</u>			<u>500MG</u>	<u>A203974</u>	<u>002</u>	May 29, 2015
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<u>AB</u>		CADILA	<u>250MG</u>	<u>A206560</u>	<u>001</u>	Nov 16, 2016
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<u>AB</u>			<u>500MG</u>	<u>A206560</u>	<u>002</u>	Nov 16, 2016
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<u>AB</u>		CADILA PHARMS LTD	<u>250MG</u>	<u>A209794</u>	<u>001</u>	Dec 12, 2017
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PRESCRIPTION DRUG PRODUCT LIST

MICONAZOLE

TABLET;BUCCAL

ORAVIG

+! GALT PHARMS

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

0.25%;81.35%;15%

N021026 001 Feb 16, 2006

MIDAZOLAM

SOLUTION;INTRAVENOUS

MIDAZOLAM IN 0.9% SODIUM CHLORIDE

+! INFORLIFE

50MG/50ML (1MG/ML)

N211844 001 Mar 22, 2021

+!

100MG/100ML (1MG/ML)

N211844 002 Mar 22, 2021

SPRAY;NASAL

NAYZILAM

+! UCB INC

5MG/SPRAY

N211321 001 May 17, 2019

MIDAZOLAM HYDROCHLORIDE

INJECTABLE;INJECTION

MIDAZOLAM HYDROCHLORIDEAP AKORNEQ 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 HIKMAEQ 1MG BASE/MLA075243 001 Jun 20, 2000APEQ 1MG BASE/MLA075247 002 Jun 23, 2000APEQ 1MG BASE/MLA075324 001 Jun 20, 2000APEQ 1MG BASE/MLA075421 002 Jun 20, 2000APEQ 1MG BASE/MLA212847 001 Dec 11, 2020APEQ 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, 2000APEQ 5MG BASE/MLA212847 002 Dec 11, 2020AP ! HOSPIRAEQ 1MG BASE/MLA075293 001 Jun 20, 2000AP !EQ 5MG BASE/MLA075293 002 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, 2002

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

SYRUP;ORAL

MIDAZOLAM HYDROCHLORIDEAA AKORNEQ 2MG BASE/MLA075958 001 Sep 04, 2003AA ! HIKMAEQ 2MG BASE/MLA075873 001 Apr 30, 2002AA PADAGIS USEQ 2MG BASE/MLA076379 001 May 02, 2005MIDODRINE HYDROCHLORIDE

TABLET;ORAL

MIDODRINE HYDROCHLORIDEAB ALEMBIC PHARMS LTD2.5MGA214734 001 Jan 21, 2021AB5MGA214734 002 Jan 21, 2021AB10MGA214734 003 Jan 21, 2021AB APOTEX2.5MGA077746 001 Sep 12, 2006AB5MGA077746 002 Sep 12, 2006AB10MGA077746 003 Sep 12, 2006AB AUROBINDO PHARMA2.5MGA212774 001 Aug 10, 2020

LTD

PRESCRIPTION DRUG PRODUCT LIST

MIDODRINE HYDROCHLORIDE

TABLET; ORAL

MIDODRINE HYDROCHLORIDE

<u>AB</u>		<u>5MG</u>	<u>A212774 002</u>	Aug 10, 2020
<u>AB</u>		<u>10MG</u>	<u>A212774 003</u>	Aug 10, 2020
<u>AB</u>	IMPAX PHARMS	<u>2.5MG</u>	<u>A076449 001</u>	May 27, 2004
<u>AB</u>		<u>5MG</u>	<u>A076449 002</u>	May 27, 2004
<u>AB</u>		<u>10MG</u>	<u>A076449 003</u>	Dec 16, 2005
<u>AB</u>	MYLAN PHARMS INC	<u>2.5MG</u>	<u>A076577 001</u>	Sep 10, 2003
<u>AB</u>		<u>5MG</u>	<u>A076577 002</u>	Sep 10, 2003
<u>AB</u>		<u>10MG</u>	<u>A076577 003</u>	Sep 10, 2003
<u>AB</u>	PAR PHARM INC	<u>2.5MG</u>	<u>A207169 001</u>	Oct 29, 2018
<u>AB</u>		<u>5MG</u>	<u>A207169 002</u>	Oct 29, 2018
<u>AB</u>		<u>10MG</u>	<u>A207169 003</u>	Oct 29, 2018
<u>AB</u>	RUBICON	<u>2.5MG</u>	<u>A212543 001</u>	Aug 19, 2019
<u>AB</u>		<u>5MG</u>	<u>A212543 002</u>	Aug 19, 2019
<u>AB</u>		<u>10MG</u>	<u>A212543 003</u>	Aug 19, 2019
<u>AB</u>	UNIQUE PHARM LABS	<u>2.5MG</u>	<u>A207613 001</u>	Nov 02, 2018
<u>AB</u>		<u>5MG</u>	<u>A207613 002</u>	Nov 02, 2018
<u>AB</u>		<u>10MG</u>	<u>A207613 003</u>	Nov 02, 2018
<u>AB</u>	XIROMED	<u>2.5MG</u>	<u>A207849 001</u>	Oct 01, 2020
<u>AB</u>		<u>5MG</u>	<u>A207849 002</u>	Oct 01, 2020
<u>AB</u>		<u>10MG</u>	<u>A207849 003</u>	Oct 01, 2020
<u>AB</u>	ZYDUS	<u>2.5MG</u>	<u>A213055 001</u>	Sep 01, 2020
<u>AB</u>		<u>5MG</u>	<u>A213055 002</u>	Sep 01, 2020
<u>AB</u>		<u>10MG</u>	<u>A213055 003</u>	Sep 01, 2020
<u>ORVATEN</u>				
<u>AB</u>	UPSHER SMITH LABS	<u>2.5MG</u>	<u>A076725 001</u>	Nov 03, 2004
<u>AB</u>	!	<u>5MG</u>	<u>A076725 002</u>	Nov 03, 2004
<u>AB</u>		<u>10MG</u>	<u>A076725 003</u>	Nov 03, 2004

MIDOSTAURIN

CAPSULE; ORAL

RYDAPT

+! NOVARTIS 25MG N207997 001 Apr 28, 2017

MIFEPRISTONE

TABLET; ORAL

KORLYMAB +! CORCEPT THERAP 300MG N202107 001 Feb 17, 2012MIFEPREXAB +! DANCO LABS LLC 200MG N020687 001 Sep 28, 2000MIFEPRISTONEAB GENBIOPRO 200MG A091178 001 Apr 11, 2019AB TEVA PHARMS USA INC 300MG A211436 001 Aug 03, 2020MIGALASTAT HYDROCHLORIDE

CAPSULE; ORAL

GALAFOLD

+! AMICUS THERAP US EQ 123MG BASE N208623 001 Aug 10, 2018

MIGLITOL

TABLET; ORAL

GLYSETAB +! PFIZER 25MG N020682 001 Dec 18, 1996AB + 50MG N020682 002 Dec 18, 1996AB + 100MG N020682 003 Dec 18, 1996MIGLITOLAB ORIENT PHARMA CO 25MG A203965 001 Feb 24, 2015AB LTD 50MG A203965 002 Feb 24, 2015AB 100MG A203965 003 Feb 24, 2015MIGLUSTAT

CAPSULE; ORAL

MIGLUSTATAB ANI PHARMS 100MG A208342 001 Apr 17, 2018AB EDENBRIDGE PHARMS 100MG A209821 001 Aug 06, 2020ZAVESCAAB +! ACTELION 100MG N021348 001 Jul 31, 2003

PRESCRIPTION DRUG PRODUCT LIST

MILNACIPRAN HYDROCHLORIDE

TABLET; ORAL

SAVELLA

+	ALLERGAN	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>		<u>EQ 1MG BASE/ML</u>	<u>A214380</u>	<u>001</u>	Apr 16, 2021
<u>AP</u>	CAPLIN	<u>EQ 1MG BASE/ML</u>	<u>A075936</u>	<u>001</u>	May 28, 2002
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 1MG BASE/ML</u>	<u>A075530</u>	<u>001</u>	May 28, 2002
<u>AP</u>	HIKMA	<u>EQ 1MG BASE/ML</u>	<u>A075660</u>	<u>001</u>	May 28, 2002
<u>AP</u>	!	<u>EQ 1MG BASE/ML</u>	<u>A077966</u>	<u>001</u>	Dec 03, 2010
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 1MG BASE/ML</u>	<u>A203280</u>	<u>001</u>	Sep 03, 2014
<u>AP</u>	HOSPIRA INC	<u>EQ 1MG BASE/ML</u>	<u>A211671</u>	<u>001</u>	Mar 24, 2020
<u>AP</u>	MEITHEAL	<u>EQ 1MG BASE/ML</u>			

MILRINONE LACTATE IN DEXTROSE 5%

<u>AP</u>	EUGIA PHARMA	<u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u>	<u>A209666</u>	<u>001</u>	Sep 03, 2020
<u>AP</u>		<u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u>	<u>A209666</u>	<u>002</u>	Sep 03, 2020
<u>AP</u>	WOODWARD	<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>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>	GLAND PHARMA LTD	<u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u>	<u>A213585</u>	<u>001</u>	Jul 16, 2020
<u>AP</u>		<u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u>	<u>A213585</u>	<u>002</u>	Jul 16, 2020
<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	<u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u>	<u>A078113</u>	<u>001</u>	May 21, 2008
<u>AP</u>	INT	<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
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MINOCYCLINE HYDROCHLORIDE

AEROSOL, FOAM; TOPICAL

AMZEEQ

+	!	VYNE	EQ 4% BASE	N212379	001	Oct 18, 2019
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ZILXI

+	!	VYNE	EQ 1.5% BASE	N213690	001	May 28, 2020
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CAPSULE; ORAL

DYNACIN

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A063067</u>	<u>003</u>	Aug 14, 1990
<u>AB</u>	ALVOGEN	<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>	+	BAUSCH	<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>		<u>EQ 50MG BASE</u>	<u>A065470</u>	<u>001</u>	Mar 11, 2008
<u>AB</u>	AUROBINDO PHARMA	<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	<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	<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

PRESCRIPTION DRUG PRODUCT LIST

MINOCYCLINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

XIMINO

JOURNEY	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

+! REMPEX	EQ 100MG BASE/VIAL	N050444 001	
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POWDER, EXTENDED RELEASE;DENTAL

ARESTIN

+! ORAPHARMA	EQ 1MG BASE	N050781 001	Feb 16, 2001
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TABLET;ORAL

MINOCYCLINE HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 50MG BASE</u>	<u>A213662 001</u>	May 01, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A213662 002</u>	May 01, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A213662 003</u>	May 01, 2020
<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>	STRIDES PHARMA	<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	<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 55MG BASE</u>	<u>A204453 008</u>	Dec 19, 2019
<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 55MG BASE</u>	<u>A202261 008</u>	Aug 21, 2019
<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>	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>	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 65MG BASE</u>	<u>A091118 003</u>	Dec 03, 2019
<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 115MG BASE</u>	<u>A091118 007</u>	Dec 03, 2019
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A091118 008</u>	Sep 25, 2014
<u>AB</u>	ZYDUS PHARMS	<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

PRESCRIPTION DRUG PRODUCT LIST

MINOCYCLINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

SOLODYN

AB	+	BAUSCH	EQ 55MG BASE	N050808 008	Aug 27, 2010
AB	+		EQ 65MG BASE	N050808 004	Jul 23, 2009
AB	+		EQ 80MG BASE	N050808 007	Aug 27, 2010
AB	+		EQ 105MG BASE	N050808 006	Aug 27, 2010
AB	+	!	EQ 115MG BASE	N050808 005	Jul 23, 2009
MINOLIRA					
		EPI HLTH	EQ 105MG BASE	N209269 001	May 08, 2017
		!	EQ 135MG BASE	N209269 002	May 08, 2017

MINOXIDIL

TABLET;ORAL

MINOXIDIL

AB		PAR PHARM	2.5MG	A071826 001	Nov 14, 1988
AB			10MG	A071839 001	Nov 14, 1988
AB		SUN PHARM INDUSTRIES	2.5MG	A072709 002	Dec 14, 1995
AB			10MG	A072709 001	Dec 14, 1995
AB		WATSON LABS	2.5MG	A071344 001	Mar 03, 1987
AB		!	10MG	A071345 001	Mar 03, 1987

MIRABEGRON

FOR SUSPENSION, EXTENDED RELEASE;ORAL

MYRBETRIQ GRANULES

+	!	APGDI	8MG/ML	N213801 001	Mar 25, 2021
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TABLET, EXTENDED RELEASE;ORAL

MYRBETRIQ

+	!	APGDI	25MG	N202611 001	Jun 28, 2012
+	!		50MG	N202611 002	Jun 28, 2012

MIRTAZAPINE

TABLET;ORAL

MIRTAZAPINE

AB		APOTEX INC	15MG	A077666 001	Aug 22, 2007
AB			30MG	A077666 002	Aug 22, 2007
AB			45MG	A077666 003	Aug 22, 2007
AB		AUROBINDO	7.5MG	A076921 001	Oct 22, 2004
AB			15MG	A076921 002	Oct 22, 2004
AB			30MG	A076921 003	Oct 22, 2004
AB			45MG	A076921 004	Oct 22, 2004
AB		MYLAN	15MG	A076122 001	Jun 19, 2003
AB			30MG	A076122 002	Jun 19, 2003
AB			45MG	A076122 003	Jun 19, 2003
AB		SUN PHARM INDS INC	7.5MG	A076541 004	Apr 22, 2004
AB			15MG	A076541 001	Apr 22, 2004
AB			30MG	A076541 002	Apr 22, 2004
AB			45MG	A076541 003	Apr 22, 2004
AB		TEVA	15MG	A076119 001	Jan 24, 2003
AB			30MG	A076119 002	Jan 24, 2003
AB			45MG	A076119 003	Jun 19, 2003
AB		UPSHER SMITH LABS	15MG	A076219 001	Jun 19, 2003
AB			30MG	A076219 002	Jun 19, 2003
AB			45MG	A076219 003	Jun 19, 2003

REMERON

AB	+	!	ORGANON USA INC	15MG	N020415 001	Jun 14, 1996
AB	+			30MG	N020415 002	Jun 14, 1996

TABLET, ORALLY DISINTEGRATING;ORAL

MIRTAZAPINE

AB		AUROBINDO PHARMA LTD	15MG	A077376 002	Dec 08, 2005
AB			30MG	A077376 003	Dec 08, 2005
AB			45MG	A077376 004	Feb 28, 2006
AB		ZYDUS PHARMS	15MG	A205798 001	Jun 01, 2017
AB			30MG	A205798 002	Jun 01, 2017
AB			45MG	A205798 003	Jun 01, 2017

REMERON SOLTAB

AB	+	!	ORGANON USA INC	15MG	N021208 001	Jan 12, 2001
AB	+			30MG	N021208 002	Jan 12, 2001
AB	+			45MG	N021208 003	Jan 12, 2001

PRESCRIPTION DRUG PRODUCT LIST

MISOPROSTOL

TABLET; ORAL

MISOPROSTOL

BX	NOVEL LABS INC	0.1MG	A091667 001	Jul 25, 2012
BX		0.2MG	A091667 002	Jul 25, 2012
	CYTOTEK			
	+ PFIZER	0.1MG	N019268 003	Sep 21, 1990
	+!	0.2MG	N019268 001	Dec 27, 1988

MITOMYCIN

FOR SOLUTION; TOPICAL

MITOSOL

+! MOBIUS THERAP

INJECTABLE; INJECTION

MITOMYCIN

AP	!	ACCORD HLTHCARE	<u>5MG/VIAL</u>	<u>A064144 001</u>	Apr 30, 1998
AP	!		<u>20MG/VIAL</u>	<u>A064144 002</u>	Apr 30, 1998
AP	!		<u>40MG/VIAL</u>	<u>A064144 003</u>	Aug 11, 2009
AP		GLAND PHARMA LTD	<u>5MG/VIAL</u>	<u>A215687 001</u>	Oct 20, 2021
AP			<u>20MG/VIAL</u>	<u>A215687 002</u>	Oct 20, 2021
AP		HIKMA	<u>5MG/VIAL</u>	<u>A064180 001</u>	Dec 23, 1999
AP			<u>20MG/VIAL</u>	<u>A064117 002</u>	Apr 19, 1995
AP			<u>20MG/VIAL</u>	<u>A064180 002</u>	Dec 23, 1999
AP			<u>40MG/VIAL</u>	<u>A064117 003</u>	Jun 02, 1999
AP		RK PHARMA	<u>5MG/VIAL</u>	<u>A202670 001</u>	Oct 13, 2017
AP			<u>20MG/VIAL</u>	<u>A202670 002</u>	Oct 13, 2017
AP			<u>40MG/VIAL</u>	<u>A203386 001</u>	Oct 13, 2017
		POWDER; PYELOCALYCEAL			
		JELMYTO			
		+! UROGEN PHARMA	40MG/VIAL	N211728 001	Apr 15, 2020

MITOTANE

TABLET; ORAL

LYSODREN

+! HRA PHARMA

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

MITOXANTRONE HYDROCHLORIDE

AP		FRESENIUS KABI USA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A077496 001</u>	Apr 11, 2006
AP			<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A077496 002</u>	Apr 11, 2006
AP			<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A077496 003</u>	Apr 11, 2006
AP		HIKMA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A076611 001</u>	Apr 11, 2006
AP			<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A076611 002</u>	Apr 11, 2006
AP			<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A076611 003</u>	Apr 11, 2006
AP	!	HOSPIRA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A076871 001</u>	Apr 11, 2006
AP	!		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A076871 002</u>	Apr 11, 2006
AP	!		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A076871 003</u>	Apr 11, 2006
AP		TEVA PHARMS USA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A077356 001</u>	Apr 11, 2006
AP			<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A077356 002</u>	Apr 11, 2006
AP			<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A077356 003</u>	Apr 11, 2006

MIVACURIUM CHLORIDE

SOLUTION; INTRAVENOUS

MIVACURIUM CHLORIDE

WOODWARD

EQ 10MG BASE/5ML (EQ 2MG BASE/ML)
EQ 20MG BASE/10ML (EQ 2MG BASE/ML)A209708 001 Oct 12, 2021
A209708 002 Oct 12, 2021MOBOCERTINIB SUCCINATE

CAPSULE; ORAL

EXKIVITY

+! TAKEDA PHARMS USA

EQ 40MG BASE

N215310 001 Sep 15, 2021

MODAFINIL

TABLET; ORAL

MODAFINIL

AB		ALEMBIC PHARMS LTD	<u>100MG</u>	<u>A202700 001</u>	Oct 18, 2012
AB			<u>200MG</u>	<u>A202700 002</u>	Oct 18, 2012
AB		APOTEX INC	<u>100MG</u>	<u>A077667 001</u>	Feb 03, 2014
AB			<u>200MG</u>	<u>A077667 002</u>	Feb 03, 2014
AB		APPCO	<u>100MG</u>	<u>A207196 001</u>	Aug 16, 2017
AB			<u>200MG</u>	<u>A207196 002</u>	Aug 16, 2017
AB		AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A202566 001</u>	Sep 27, 2012
AB			<u>200MG</u>	<u>A202566 002</u>	Sep 27, 2012
AB		CADILA	<u>100MG</u>	<u>A209966 001</u>	Sep 14, 2017

PRESCRIPTION DRUG PRODUCT LIST

MODAFINIL

TABLET; ORAL

MODAFINIL

<u>AB</u>		<u>200MG</u>	<u>A209966 002</u>	Sep 14, 2017
<u>AB</u>	ORBION PHARMS	<u>100MG</u>	<u>A078963 001</u>	Sep 26, 2012
<u>AB</u>		<u>200MG</u>	<u>A078963 002</u>	Sep 26, 2012
<u>AB</u>	WATSON LABS INC	<u>100MG</u>	<u>A076715 001</u>	Nov 01, 2012
<u>AB</u>		<u>200MG</u>	<u>A076715 002</u>	Nov 01, 2012

PROVIGIL

<u>AB</u>	+ CEPHALON	<u>100MG</u>	<u>N020717 001</u>	Dec 24, 1998
<u>AB</u>	+	<u>200MG</u>	<u>N020717 002</u>	Dec 24, 1998

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>7.5MG</u>	<u>A078454 001</u>	Jun 02, 2008
<u>AB</u>		<u>15MG</u>	<u>A078454 002</u>	Jun 02, 2008
<u>AB</u>	CHARTWELL RX	<u>7.5MG</u>	<u>A077536 001</u>	Nov 30, 2006
<u>AB</u>		<u>15MG</u>	<u>A077536 002</u>	Nov 30, 2006
<u>AB</u>	GLENMARK GENERICS	<u>7.5MG</u>	<u>A090416 001</u>	Mar 30, 2010
<u>AB</u>		<u>15MG</u>	<u>A090416 002</u>	Mar 30, 2010
<u>AB</u>	TEVA	<u>7.5MG</u>	<u>A076204 001</u>	May 08, 2003
<u>AB</u>	!	<u>15MG</u>	<u>A076204 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

+ ORGANON

0.05MG/INH

N205641 003 Aug 12, 2019

+

0.10MG/INH

N205641 001 Apr 25, 2014

+!

0.20MG/INH

N205641 002 Apr 25, 2014

CREAM; TOPICAL

MOMETASONE FUROATE

<u>AB</u>	COSETTE	<u>0.1%</u>	<u>A077447 001</u>	May 22, 2006
<u>AB</u>	! GLENMARK GENERICS	<u>0.1%</u>	<u>A078541 001</u>	May 28, 2008
<u>AB</u>	TARO	<u>0.1%</u>	<u>A076679 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 001</u>	Mar 30, 1989
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MOMETASONE FUROATE

<u>AB</u>	COSETTE	<u>0.1%</u>	<u>A077678 001</u>	Nov 21, 2007
<u>AB</u>	FOUGERA PHARMS	<u>0.1%</u>	<u>A075919 001</u>	Nov 29, 2007
<u>AB</u>	GLENMARK GENERICS	<u>0.1%</u>	<u>A090506 001</u>	Aug 09, 2010
<u>AB</u>	PADAGIS ISRAEL	<u>0.1%</u>	<u>A077180 001</u>	Apr 06, 2005
<u>AB</u>	TARO	<u>0.1%</u>	<u>A076788 001</u>	Mar 15, 2006

OINTMENT; TOPICAL

MOMETASONE FUROATE

<u>AB</u>	COSETTE	<u>0.1%</u>	<u>A077401 001</u>	Jun 20, 2006
<u>AB</u>	FOUGERA PHARMS	<u>0.1%</u>	<u>A077061 001</u>	Mar 28, 2005
<u>AB</u>	! GLENMARK GENERICS	<u>0.1%</u>	<u>A078571 001</u>	May 28, 2008
<u>AB</u>	PADAGIS US	<u>0.1%</u>	<u>A076067 001</u>	Mar 18, 2002

POWDER; INHALATION

ASMANEX TWISTHALER

+ ORGANON

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>0.05MG/SPRAY</u>	<u>A207989 001</u>	Apr 03, 2017
<u>AB</u>	! APOTEX INC	<u>0.05MG/SPRAY</u>	<u>A091161 001</u>	Mar 22, 2016

PRESCRIPTION DRUG PRODUCT LIST

MONOMETHYL FUMARATE

CAPSULE, DELAYED RELEASE;ORAL

BAFIERTAM

+	BANNER LIFE SCIENCES	95MG	N210296	001	Apr 28, 2020
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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>	AUROBINDO PHARMA LTD	<u>EQ 4MG BASE/PACKET</u>	<u>A213471</u>	<u>001</u>	Feb 18, 2020
<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>	TEVA PHARMS	<u>EQ 4MG BASE/PACKET</u>	<u>A090955</u>	<u>001</u>	Aug 03, 2012
<u>AB</u>	TORRENT	<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
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TABLET;ORAL

MONTELUKAST SODIUM

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 10MG BASE</u>	<u>A202717</u>	<u>001</u>	Sep 21, 2012
<u>AB</u>	ADAPTIS	<u>EQ 10MG BASE</u>	<u>A209012</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>	AMNEAL PHARMS	<u>EQ 10MG BASE</u>	<u>A204604</u>	<u>001</u>	Sep 04, 2015
<u>AB</u>	ANBISON LAB	<u>EQ 10MG BASE</u>	<u>A205683</u>	<u>001</u>	Jan 12, 2016
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 10MG BASE</u>	<u>A202468</u>	<u>001</u>	Aug 03, 2012
<u>AB</u>	CIPLA	<u>EQ 10MG BASE</u>	<u>A207463</u>	<u>001</u>	Oct 28, 2016
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 10MG BASE</u>	<u>A201582</u>	<u>001</u>	Aug 06, 2012
<u>AB</u>	GLENMARK GENERICS	<u>EQ 10MG BASE</u>	<u>A090926</u>	<u>001</u>	Aug 03, 2012
<u>AB</u>	HETERO LABS LTD V	<u>EQ 10MG BASE</u>	<u>A202843</u>	<u>001</u>	Sep 10, 2014
<u>AB</u>	L FERRIGO CO	<u>EQ 10MG BASE</u>	<u>A206112</u>	<u>001</u>	Apr 26, 2017
<u>AB</u>	LANNETT CO INC	<u>EQ 10MG BASE</u>	<u>A201522</u>	<u>001</u>	Aug 03, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A203366</u>	<u>001</u>	Sep 11, 2014
<u>AB</u>	SANDOZ INC	<u>EQ 10MG BASE</u>	<u>A200889</u>	<u>001</u>	Aug 03, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 10MG BASE</u>	<u>A078605</u>	<u>001</u>	Aug 03, 2012
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A201515</u>	<u>001</u>	Aug 03, 2012
<u>AB</u>	UNICHEM	<u>EQ 10MG BASE</u>	<u>A204290</u>	<u>001</u>	Oct 08, 2015
<u>AB</u>	UNIMARK REMEDIES LTD	<u>EQ 10MG BASE</u>	<u>A202859</u>	<u>001</u>	Oct 30, 2014

SINGULAIR

<u>AB</u>	+	MSD MERCK CO	<u>EQ 10MG BASE</u>	<u>N020829</u>	<u>002</u>	Feb 20, 1998
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TABLET, CHEWABLE;ORAL

MONTELUKAST SODIUM

<u>AB</u>	ADAPTIS	<u>EQ 4MG BASE</u>	<u>A209011</u>	<u>001</u>	Apr 18, 2017
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A209011</u>	<u>002</u>	Apr 18, 2017
<u>AB</u>	AMNEAL PHARMS	<u>EQ 4MG BASE</u>	<u>A205107</u>	<u>001</u>	Sep 04, 2020
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A205107</u>	<u>002</u>	Sep 04, 2020
<u>AB</u>	ANBISON LAB	<u>EQ 4MG BASE</u>	<u>A205695</u>	<u>001</u>	Nov 05, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A205695</u>	<u>002</u>	Nov 05, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 4MG BASE</u>	<u>A202096</u>	<u>001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A202096</u>	<u>002</u>	Aug 03, 2012
<u>AB</u>	CIPLA	<u>EQ 4MG BASE</u>	<u>A207464</u>	<u>001</u>	Dec 06, 2018
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A207464</u>	<u>002</u>	Dec 06, 2018
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE</u>	<u>A201581</u>	<u>001</u>	Aug 06, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A201581</u>	<u>002</u>	Aug 06, 2012
<u>AB</u>	HETERO LABS LTD V	<u>EQ 4MG BASE</u>	<u>A204093</u>	<u>001</u>	May 22, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A204093</u>	<u>002</u>	May 22, 2015
<u>AB</u>	LANNETT CO INC	<u>EQ 4MG BASE</u>	<u>A200405</u>	<u>001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A200405</u>	<u>002</u>	Aug 03, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 4MG BASE</u>	<u>A203582</u>	<u>001</u>	Mar 12, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203582</u>	<u>002</u>	Mar 12, 2015
<u>AB</u>	SANDOZ INC	<u>EQ 4MG BASE</u>	<u>A091414</u>	<u>001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A091414</u>	<u>002</u>	Aug 03, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 4MG BASE</u>	<u>A078723</u>	<u>001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078723</u>	<u>002</u>	Aug 03, 2012
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 4MG BASE</u>	<u>A090984</u>	<u>001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A090984</u>	<u>002</u>	Aug 03, 2012
<u>AB</u>	UNICHEM	<u>EQ 4MG BASE</u>	<u>A208621</u>	<u>001</u>	Jul 02, 2018
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A208621</u>	<u>002</u>	Jul 02, 2018
<u>AB</u>	UNIMARK REMEDIES LTD	<u>EQ 4MG BASE</u>	<u>A203037</u>	<u>001</u>	Oct 30, 2014
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203037</u>	<u>002</u>	Oct 30, 2014

SINGULAIR

<u>AB</u>	+	MSD MERCK CO	<u>EQ 4MG BASE</u>	<u>N020830</u>	<u>002</u>	Mar 03, 2000
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PRESCRIPTION DRUG PRODUCT LIST

MONTELUKAST SODIUM

TABLET, CHEWABLE;ORAL

SINGULAIR

<u>AB</u>	+		<u>EO 5MG BASE</u>	<u>N020830</u>	<u>001</u>	Feb 20, 1998
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MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL

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>		NORTEC DEV ASSOC	<u>20MG</u>	<u>A203158</u>	<u>001</u>	Aug 04, 2021
<u>AB</u>			<u>30MG</u>	<u>A203158</u>	<u>002</u>	Aug 04, 2021
<u>AB</u>			<u>50MG</u>	<u>A203158</u>	<u>003</u>	Aug 04, 2021
<u>AB</u>			<u>60MG</u>	<u>A203158</u>	<u>004</u>	Aug 04, 2021
<u>AB</u>			<u>80MG</u>	<u>A203158</u>	<u>005</u>	Aug 04, 2021
<u>AB</u>			<u>100MG</u>	<u>A203158</u>	<u>006</u>	Aug 04, 2021
<u>AB</u>		UPSHER SMITH LABS	<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
		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
		IMPAX LABS INC	40MG	A200411	007	Jul 25, 2018
	!	UPSHER SMITH LABS	10MG	A202104	001	Jun 03, 2013

INJECTABLE; INJECTION

DURAMORPH PF

<u>AP</u>	+	HIKMA	<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>	+	HIKMA	<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>	!	HIKMA	<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
<u>AP</u>			<u>0.5MG/ML</u>	<u>A073509</u>	<u>001</u>	Sep 30, 1992
<u>AP</u>			<u>1MG/ML</u>	<u>A073510</u>	<u>001</u>	Sep 30, 1992
<u>AP</u>	+	HOSPIRA INC	<u>4MG/ML</u>	<u>N202515</u>	<u>002</u>	Nov 14, 2011
<u>AP</u>	+		<u>8MG/ML</u>	<u>N202515</u>	<u>003</u>	Nov 14, 2011
<u>AP</u>	+		<u>10MG/ML</u>	<u>N202515</u>	<u>004</u>	Nov 14, 2011
	+		2MG/ML	N202515	001	Nov 14, 2011
	+		50MG/ML	N202515	006	Apr 29, 2021

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

MORPHINE SULFATE

	+	FRESENIUS KABI USA	2MG/ML (2MG/ML)	N204223	001	Oct 30, 2013
	+		4MG/ML (4MG/ML)	N204223	002	Oct 30, 2013
	+		5MG/ML (5MG/ML)	N204223	003	Oct 30, 2013
	+		8MG/ML (8MG/ML)	N204223	004	Oct 30, 2013
	+		10MG/ML (10MG/ML)	N204223	005	Oct 30, 2013

SOLUTION; ORAL

MORPHINE SULFATE

<u>AA</u>		AKORN	<u>100MG/5ML</u>	<u>A208809</u>	<u>001</u>	Jul 06, 2017
<u>AA</u>	+	HIKMA	<u>10MG/5ML</u>	<u>N022195</u>	<u>001</u>	Mar 17, 2008
<u>AA</u>	+		<u>20MG/5ML</u>	<u>N022195</u>	<u>002</u>	Mar 17, 2008
<u>AA</u>	+		<u>100MG/5ML</u>	<u>N022195</u>	<u>003</u>	Jan 25, 2010
<u>AA</u>		PADAGIS US	<u>100MG/5ML</u>	<u>A201574</u>	<u>001</u>	Aug 06, 2012
<u>AA</u>		PHARM ASSOC	<u>100MG/5ML</u>	<u>A206573</u>	<u>001</u>	Nov 14, 2016
<u>AA</u>		RHODES PHARMS	<u>10MG/5ML</u>	<u>A206308</u>	<u>001</u>	Jun 22, 2017
<u>AA</u>			<u>20MG/5ML</u>	<u>A206420</u>	<u>001</u>	Jul 12, 2016

PRESCRIPTION DRUG PRODUCT LIST

MORPHINE SULFATE

SOLUTION;ORAL

MORPHINE SULFATE

<u>AA</u>		<u>100MG/5ML</u>	<u>A206308 002</u>	Jun 22, 2017
<u>AA</u>	SPECGX LLC	<u>100MG/5ML</u>	<u>A202348 001</u>	Jul 15, 2011
<u>AA</u>	TRIS PHARMA INC	<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

TABLET;ORAL

MORPHINE SULFATE

<u>AB</u>	ALKEM LABS LTD	<u>15MG</u>	<u>A212451 001</u>	Dec 03, 2020
<u>AB</u>		<u>30MG</u>	<u>A212451 002</u>	Dec 03, 2020
<u>AB</u>	+ HIKMA	<u>15MG</u>	<u>N022207 001</u>	Mar 17, 2008
<u>AB</u>	+!	<u>30MG</u>	<u>N022207 002</u>	Mar 17, 2008
<u>AB</u>	MAYNE PHARMA INC	<u>15MG</u>	<u>A207270 001</u>	Jan 12, 2022
<u>AB</u>		<u>30MG</u>	<u>A207270 002</u>	Jan 12, 2022
<u>AB</u>	UPSHER SMITH LABS	<u>15MG</u>	<u>A210610 001</u>	Jul 22, 2019
<u>AB</u>		<u>30MG</u>	<u>A210610 002</u>	Jul 22, 2019

TABLET, EXTENDED RELEASE;ORAL

MORPHINE SULFATE

<u>AB</u>	ACTAVIS ELIZABETH	<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>	DAVA PHARMS INC	<u>15MG</u>	<u>A075407 001</u>	Jan 28, 2000
<u>AB</u>	MAYNE PHARMA INC	<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>	NESHER PHARMS	<u>30MG</u>	<u>A076720 002</u>	Dec 23, 2005
<u>AB</u>		<u>60MG</u>	<u>A076720 001</u>	May 19, 2004
<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>	STRIDES PHARMA	<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
<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

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

MOXIDECTIN

TABLET;ORAL

MOXIDECTIN

	+! MDGH	2MG	N210867 001	Jun 13, 2018
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PRESCRIPTION DRUG PRODUCT LIST

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>	ALEMBIC PHARMS LTD	<u>EQ 0.5% BASE</u>	<u>A209469 001</u>	Feb 13, 2019
<u>AT1</u>	APOTEX	<u>EQ 0.5% BASE</u>	<u>A090080 001</u>	Jun 30, 2017
<u>AT1</u>	EUGIA PHARMA	<u>EQ 0.5% BASE</u>	<u>A206242 001</u>	Oct 04, 2017
<u>AT1</u>	GLAND PHARMA LTD	<u>EQ 0.5% BASE</u>	<u>A208778 001</u>	Mar 30, 2020
<u>AT1</u>	LUPIN LTD	<u>EQ 0.5% BASE</u>	<u>A202867 001</u>	Sep 04, 2014
<u>AT1</u>	MYLAN	<u>EQ 0.5% BASE</u>	<u>A206447 001</u>	Mar 30, 2020
<u>AT1</u>	UPSHER SMITH LABS	<u>EQ 0.5% BASE</u>	<u>A212616 001</u>	Feb 10, 2021
<u>AT1</u>	WATSON LABS INC	<u>EQ 0.5% BASE</u>	<u>A202525 001</u>	Mar 06, 2015

VIGAMOXAT1 +! NOVARTIS EQ 0.5% BASE N021598 001 Apr 15, 2003MOXEZAAT2 +! NOVARTIS EQ 0.5% BASE N022428 001 Nov 19, 2010MOXIFLOXACIN HYDROCHLORIDEAT2 LUPIN LTD EQ 0.5% BASE A204079 001 May 28, 2015

TABLET; ORAL

MOXIFLOXACIN HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 400MG BASE</u>	<u>A202632 001</u>	Mar 04, 2014
<u>AB</u>	CROSSMEDIKA SA	<u>EQ 400MG BASE</u>	<u>A205348 001</u>	Jan 14, 2016
<u>AB</u>	! DR REDDYS LABS LTD	<u>EQ 400MG BASE</u>	<u>A076938 001</u>	Mar 04, 2014
<u>AB</u>	MSN	<u>EQ 400MG BASE</u>	<u>A208682 001</u>	Sep 22, 2017
<u>AB</u>	NOVEL LABS INC	<u>EQ 400MG BASE</u>	<u>A207285 001</u>	Feb 13, 2017
<u>AB</u>	TEVA PHARMS USA	<u>EQ 400MG BASE</u>	<u>A077437 001</u>	Feb 18, 2014
<u>AB</u>	TORRENT	<u>EQ 400MG BASE</u>	<u>A200160 001</u>	Apr 03, 2014

MUPIROCIN

OINTMENT; TOPICAL

MUPIROCIN

<u>AB</u>	FOUGERA PHARMS	<u>2%</u>	<u>A065192 001</u>	Nov 30, 2005
<u>AB</u>	GLENMARK PHARMS	<u>2%</u>	<u>A090480 001</u>	Jun 08, 2011
<u>AB</u>	! PADAGIS ISRAEL	<u>2%</u>	<u>A065123 001</u>	Nov 07, 2003
<u>AB</u>	TARO	<u>2%</u>	<u>A065170 001</u>	Sep 23, 2005
<u>AB</u>	TEVA	<u>2%</u>	<u>A065085 001</u>	Nov 07, 2003

CENTANY

EX PADAGIS US 2% N050788 001 Dec 04, 2002

MUPIROCIN CALCIUM

CREAM; TOPICAL

MUPIROCIN

<u>AB</u>	ALBOR DERMACEUTICALS	<u>EQ 2% BASE</u>	<u>A213053 001</u>	Nov 16, 2021
<u>AB</u>	ENCUBE	<u>EQ 2% BASE</u>	<u>A213076 001</u>	Aug 31, 2021
<u>AB</u>	! GLENMARK PHARMS INC	<u>EQ 2% BASE</u>	<u>A201587 001</u>	Jan 24, 2013
<u>AB</u>	TARO	<u>EQ 2% BASE</u>	<u>A207116 001</u>	Apr 27, 2020

MYCOPHENOLATE MOFETIL

CAPSULE; ORAL

CELLCEPTAB +! ROCHE PALO 250MG N050722 001 May 03, 1995MYCOPHENOLATE MOFETIL

<u>AB</u>	ACCORD HLTHCARE	<u>250MG</u>	<u>A090253 001</u>	May 04, 2009
<u>AB</u>	ALKEM LABS LTD	<u>250MG</u>	<u>A200197 001</u>	Jun 13, 2013
<u>AB</u>	CONCORD BIOTECH LTD	<u>250MG</u>	<u>A210181 001</u>	Jan 08, 2019
<u>AB</u>	HIKMA	<u>250MG</u>	<u>A065410 001</u>	Jul 29, 2008
<u>AB</u>	MYLAN	<u>250MG</u>	<u>A065520 001</u>	May 04, 2009
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A065379 001</u>	Oct 15, 2008
<u>AB</u>	STRIDES PHARMA	<u>250MG</u>	<u>A090055 001</u>	Jun 10, 2010
<u>AB</u>	TEVA PHARMS	<u>250MG</u>	<u>A065491 001</u>	May 06, 2009
<u>AB</u>	ZHEJIANG HISUN PHARM	<u>250MG</u>	<u>A204077 001</u>	Nov 13, 2017

FOR SUSPENSION; ORAL

CELLCEPTAB +! ROCHE PALO 200MG/ML N050759 001 Oct 01, 1998MYCOPHENOLATE MOFETILAB ALKEM LABS LTD 200MG/ML A203005 001 Nov 14, 2014

PRESCRIPTION DRUG PRODUCT LIST

MYCOPHENOLATE MOFETIL

FOR SUSPENSION;ORAL

MYCOPHENOLATE MOFETIL

AB	AMNEAL	<u>200MG/ML</u>	<u>A214871</u>	<u>001</u>	Nov 02, 2021
AB	LANNETT CO INC	<u>200MG/ML</u>	<u>A214525</u>	<u>001</u>	Jul 29, 2021
AB	VISTAPHARM	<u>200MG/ML</u>	<u>A210370</u>	<u>001</u>	Feb 12, 2019

TABLET;ORAL

CELLCEPT

AB	+ ! ROCHE PALO	<u>500MG</u>	<u>N050723</u>	<u>001</u>	Jun 19, 1997
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MYCOPHENOLATE MOFETIL

AB	ACCORD HLTHCARE	<u>500MG</u>	<u>A065416</u>	<u>001</u>	May 04, 2009
AB	ALKEM LABS LTD	<u>500MG</u>	<u>A091249</u>	<u>001</u>	Nov 04, 2011
AB	CONCORD BIOTECH LTD	<u>500MG</u>	<u>A212087</u>	<u>001</u>	Jul 31, 2020
AB	HIKMA	<u>500MG</u>	<u>A065413</u>	<u>001</u>	Jul 29, 2008
AB	MYLAN	<u>500MG</u>	<u>A065521</u>	<u>001</u>	May 04, 2009
AB	SANDOZ	<u>500MG</u>	<u>A065451</u>	<u>001</u>	Oct 15, 2008
AB	STRIDES PHARMA	<u>500MG</u>	<u>A090456</u>	<u>001</u>	Jun 10, 2010
AB	ZHEJIANG HISUN PHARM	<u>500MG</u>	<u>A204076</u>	<u>001</u>	Nov 16, 2017

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

INJECTABLE;INJECTION

CELLCEPT

AP	+ ! ROCHE PALO	<u>500MG/VIAL</u>	<u>N050758</u>	<u>001</u>	Aug 12, 1998
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MYCOPHENOLATE MOFETIL HYDROCHLORIDE

AP	AKORN	<u>500MG/VIAL</u>	<u>A204043</u>	<u>001</u>	Feb 28, 2017
AP	MEITHEAL	<u>500MG/VIAL</u>	<u>A212130</u>	<u>001</u>	Jan 15, 2021
AP	MYLAN LABS LTD	<u>500MG/VIAL</u>	<u>A203859</u>	<u>001</u>	Mar 31, 2017
AP	PAR STERILE PRODUCTS	<u>500MG/VIAL</u>	<u>A203575</u>	<u>001</u>	Oct 28, 2016
AP	ZYDUS PHARMS	<u>500MG/VIAL</u>	<u>A204473</u>	<u>001</u>	Aug 31, 2017

MYCOPHENOLIC ACID

TABLET, DELAYED RELEASE;ORAL

MYCOPHENOLIC ACID

AB	BIOCON PHARMA	<u>EQ 180MG BASE</u>	<u>A214630</u>	<u>001</u>	Nov 29, 2021
AB		<u>EQ 360MG BASE</u>	<u>A214630</u>	<u>002</u>	Nov 29, 2021

MYCOPHENOLIC SODIUM

TABLET, DELAYED RELEASE;ORAL

MYCOPHENOLIC SODIUM

AB	ACCORD HLTHCARE	<u>EQ 180MG BASE</u>	<u>A202555</u>	<u>001</u>	Aug 23, 2017
AB		<u>EQ 360MG BASE</u>	<u>A202555</u>	<u>002</u>	Aug 23, 2017
AB	ALKEM LABS LTD	<u>EQ 180MG BASE</u>	<u>A208315</u>	<u>001</u>	Sep 23, 2021
AB		<u>EQ 360MG BASE</u>	<u>A208315</u>	<u>002</u>	Sep 23, 2021
AB	AMTA	<u>EQ 180MG BASE</u>	<u>A214376</u>	<u>001</u>	Feb 10, 2021
AB		<u>EQ 360MG BASE</u>	<u>A214376</u>	<u>002</u>	Feb 10, 2021
AB	APOTEX INC	<u>EQ 180MG BASE</u>	<u>A091558</u>	<u>001</u>	Aug 21, 2012
AB		<u>EQ 360MG BASE</u>	<u>A091558</u>	<u>002</u>	Aug 19, 2014
AB	CONCORD BIOTECH LTD	<u>EQ 180MG BASE</u>	<u>A211173</u>	<u>001</u>	Dec 13, 2019
AB		<u>EQ 360MG BASE</u>	<u>A211173</u>	<u>002</u>	Dec 13, 2019
AB	RK PHARMA	<u>EQ 180MG BASE</u>	<u>A091248</u>	<u>002</u>	Jan 08, 2014
AB		<u>EQ 360MG BASE</u>	<u>A091248</u>	<u>001</u>	Jan 08, 2014
AB	TWI PHARMS	<u>EQ 180MG BASE</u>	<u>A214289</u>	<u>001</u>	Nov 03, 2021
AB		<u>EQ 360MG BASE</u>	<u>A214289</u>	<u>002</u>	Nov 03, 2021

MYFORTIC

AB	+ NOVARTIS	<u>EQ 180MG BASE</u>	<u>N050791</u>	<u>001</u>	Feb 27, 2004
AB	+ !	<u>EQ 360MG BASE</u>	<u>N050791</u>	<u>002</u>	Feb 27, 2004

NABILONE

CAPSULE;ORAL

CESAMET

+ !	BAUSCH	1MG	N018677	001	Dec 26, 1985
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NABUMETONE

TABLET;ORAL

NABUMETONE

AB	ANNORA PHARMA	<u>500MG</u>	<u>A090445</u>	<u>001</u>	Jan 12, 2011
AB		<u>750MG</u>	<u>A090445</u>	<u>002</u>	Jan 12, 2011
AB	CHARTWELL MOLECULES	<u>500MG</u>	<u>A076009</u>	<u>001</u>	Jan 24, 2003
AB		<u>750MG</u>	<u>A076009</u>	<u>002</u>	Jan 24, 2003
AB	CHARTWELL RX	<u>500MG</u>	<u>A075280</u>	<u>001</u>	Feb 25, 2002
AB		<u>750MG</u>	<u>A075280</u>	<u>002</u>	Feb 25, 2002
AB	INVAGEN PHARMS	<u>500MG</u>	<u>A078671</u>	<u>001</u>	Mar 07, 2008
AB	!	<u>750MG</u>	<u>A078671</u>	<u>002</u>	Mar 07, 2008

PRESCRIPTION DRUG PRODUCT LIST

NABUMETONE

TABLET; ORAL

NABUMETONE

<u>AB</u>	LGM PHARMA	<u>500MG</u>	<u>A203166 001</u>	Aug 30, 2019
<u>AB</u>		<u>750MG</u>	<u>A203166 002</u>	Aug 30, 2019
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A078420 001</u>	Sep 24, 2008
<u>AB</u>		<u>750MG</u>	<u>A078420 002</u>	Sep 24, 2008
<u>AB</u>	WATSON LABS	<u>500MG</u>	<u>A091083 001</u>	Jun 13, 2011
<u>AB</u>		<u>750MG</u>	<u>A091083 002</u>	Jun 13, 2011
	LGM PHARMA	1GM	A203166 003	Aug 30, 2019

NADOLOL

TABLET; ORAL

CORGARD

<u>AB</u>	+	USWM	<u>20MG</u>	<u>N018063 005</u>	Oct 28, 1986
<u>AB</u>	+		<u>40MG</u>	<u>N018063 001</u>	
<u>AB</u>	+	!	<u>80MG</u>	<u>N018063 002</u>	

NADOLOL

<u>AB</u>		AMNEAL PHARMS CO	<u>20MG</u>	<u>A208832 001</u>	Jun 02, 2017
<u>AB</u>			<u>40MG</u>	<u>A208832 002</u>	Jun 02, 2017
<u>AB</u>			<u>80MG</u>	<u>A208832 003</u>	Jun 02, 2017
<u>AB</u>		AUROBINDO PHARMA LTD	<u>40MG</u>	<u>A201893 001</u>	Sep 16, 2015
<u>AB</u>			<u>80MG</u>	<u>A201893 002</u>	Sep 16, 2015
<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>		HERITAGE PHARMA	<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>		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>		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>		RISING	<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>		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>		VGYAAN	<u>20MG</u>	<u>A212856 001</u>	Sep 13, 2019
<u>AB</u>			<u>40MG</u>	<u>A212856 002</u>	Sep 13, 2019
<u>AB</u>			<u>80MG</u>	<u>A212856 003</u>	Sep 13, 2019
<u>AB</u>		ZYDUS PHARMS	<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

+! PFIZER

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>		FRESENIUS	<u>EQ 10GM BASE/VIAL</u>	<u>A206761 001</u>	Jun 02, 2020
<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>		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

PRESCRIPTION DRUG PRODUCT LIST

NAFCILLIN SODIUM

INJECTABLE; INJECTION

NAFCILLIN SODIUM

<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A062732</u>	<u>002</u>	Dec 23, 1986
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A062527</u>	<u>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</u>	<u>001</u> Jan 06, 2016
<u>AB</u>		TOLMAR	<u>2%</u>	<u>A206960</u>	<u>001</u> Apr 10, 2017
<u>AB</u>		XIROMED	<u>2.0%</u>	<u>A210038</u>	<u>001</u> Sep 22, 2020

NAFTIN

<u>AB</u>	+!	SEBELA IRELAND LTD	<u>2%</u>	<u>N019599</u>	<u>002</u> Jan 13, 2012
NAFTIFINE HYDROCHLORIDE					
	!	TARO PHARMS	1%	A205975	001 Sep 08, 2016

GEL; TOPICAL

NAFTIFINE HYDROCHLORIDE

<u>AB</u>		TOLMAR	<u>1%</u>	<u>A206165</u>	<u>001</u> Mar 20, 2019
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NAFTIN

<u>AB</u>	+!	SEBELA IRELAND LTD	<u>1%</u>	<u>N019356</u>	<u>001</u> Jun 18, 1990
<u>AB</u>	+!		<u>2%</u>	<u>N204286</u>	<u>001</u> Jun 27, 2013

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE HYDROCHLORIDE

<u>AP</u>	!	HOSPIRA	<u>10MG/ML</u>	<u>A070914</u>	<u>001</u> Feb 03, 1989
<u>AP</u>	!		<u>10MG/ML</u>	<u>A070915</u>	<u>001</u> Feb 03, 1989
<u>AP</u>	!		<u>20MG/ML</u>	<u>A070916</u>	<u>001</u> Feb 03, 1989
<u>AP</u>	!		<u>20MG/ML</u>	<u>A070918</u>	<u>001</u> Feb 03, 1989

NALDEMEDINE TOSYLATE

TABLET; ORAL

SYMPROIC

	+!	BDSI	EQ 0.2MG BASE	N208854	001 Mar 23, 2017
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NALOXEGOL OXALATE

TABLET; ORAL

MOVANTIK

	+	REDHILL	EQ 12.5MG BASE	N204760	001 Sep 16, 2014
	+!		EQ 25MG BASE	N204760	002 Sep 16, 2014

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

<u>AP</u>		HIKMA	<u>0.4MG/ML</u>	<u>A070299</u>	<u>001</u> Sep 24, 1986
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NALOXONE HYDROCHLORIDE

<u>AP</u>		AKORN	<u>0.4MG/ML</u>	<u>A208871</u>	<u>001</u> Feb 28, 2017
<u>AP</u>			<u>0.4MG/ML</u>	<u>A208872</u>	<u>001</u> Mar 14, 2017
<u>AP</u>		BAXTER HLTHCARE	<u>0.4MG/ML</u>	<u>A214785</u>	<u>001</u> Jan 29, 2021
CORP					
<u>AP</u>		DR REDDYS	<u>1MG/ML</u>	<u>A213209</u>	<u>001</u> Mar 16, 2020
<u>AP</u>		EUGIA PHARMA	<u>0.4MG/ML</u>	<u>A212455</u>	<u>001</u> Oct 15, 2019
<u>AP</u>			<u>1MG/ML</u>	<u>A213279</u>	<u>001</u> Jan 14, 2021
<u>AP</u>			<u>0.4MG/ML</u>	<u>A212456</u>	<u>001</u> Nov 04, 2019
<u>AP</u>	!	HOSPIRA	<u>0.4MG/ML</u>	<u>A070172</u>	<u>001</u> Sep 24, 1986
<u>AP</u>	!		<u>0.4MG/ML</u>	<u>A070256</u>	<u>001</u> Jan 07, 1987
<u>AP</u>	!		<u>0.4MG/ML</u>	<u>A070257</u>	<u>001</u> Jan 07, 1987
<u>AP</u>	!	INTL MEDICATION	<u>1MG/ML</u>	<u>A072076</u>	<u>001</u> Mar 24, 1988
<u>AP</u>		MYLAN INSTITUTIONAL	<u>0.4MG/ML</u>	<u>A204997</u>	<u>001</u> Mar 06, 2014
<u>AP</u>			<u>0.4MG/ML</u>	<u>A205014</u>	<u>001</u> Jun 29, 2016
<u>AP</u>		SOMERSET THERAPS	<u>0.4MG/ML</u>	<u>A207633</u>	<u>001</u> Aug 08, 2017
LLC					
<u>AP</u>			<u>0.4MG/ML</u>	<u>A207634</u>	<u>001</u> Jul 26, 2017
SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS					
ZIMHI					
	+!	ADAMIS PHARMS CORP	5MG/0.5ML (5MG/0.5ML)	N212854	001 Oct 15, 2021
SPRAY; NASAL					
KLOXXADO					
	+!	HIKMA	8MG/SPRAY	N212045	001 Apr 29, 2021

PRESCRIPTION DRUG PRODUCT LIST

NALOXONE HYDROCHLORIDE

SPRAY, METERED;NASAL

NALOXONE HYDROCHLORIDE

AB	TEVA PHARMS USA	4MG/SPRAY	A209522 001	Apr 19, 2019
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NARCAN

AB	+! EMERGENT	4MG/SPRAY	N208411 001	Nov 18, 2015
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NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET;ORAL

NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE

AB	LUPIN	EQ 0.5MG BASE;EQ 50MG BASE	A075735 001	Jul 11, 2001
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AB	SUN PHARM INDS LTD	EQ 0.5MG BASE;EQ 50MG BASE	A075523 001	Mar 17, 2000
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AB	! WATSON LABS	EQ 0.5MG BASE;EQ 50MG BASE	A074736 001	Jan 21, 1997
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NALTREXONE

FOR SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

VIVITROL

+!	ALKERMES	380MG/VIAL	N021897 001	Apr 13, 2006
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NALTREXONE HYDROCHLORIDE

TABLET;ORAL

NALTREXONE HYDROCHLORIDE

AB	ACCORD HLTHCARE	50MG	A091205 001	Aug 17, 2011
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AB	BARR	50MG	A074918 001	May 08, 1998
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AB	CHARTWELL	50MG	A207905 001	Jul 21, 2017
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AB	ELITE LABS	50MG	A075274 001	May 26, 1999
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AB	! SPECGX LLC	50MG	A076264 002	Mar 22, 2002
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AB	SUN PHARM	50MG	A090356 001	Feb 24, 2012
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	SPECGX LLC	25MG	A076264 001	Mar 22, 2002
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		100MG	A076264 003	Mar 22, 2002
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NAPROXEN

SUSPENSION;ORAL

NAPROSYN

AB	+ ATNAHS PHARMA US	25MG/ML	N018965 001	Mar 23, 1987
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NAPROXEN

AB	AMNEAL	25MG/ML	A212705 001	Jul 31, 2020
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AB	! HIKMA	25MG/ML	A074190 001	Mar 30, 1994
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AB	NOVITIUM PHARMA	25MG/ML	A211910 001	Mar 10, 2021
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TABLET;ORAL

NAPROSYN

AB	+! ATNAHS PHARMA US	500MG	N017581 004	Apr 15, 1982
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NAPROXEN

AB	AMNEAL PHARMS NY	250MG	A075927 001	Dec 18, 2001
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AB		375MG	A075927 002	Dec 18, 2001
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AB		500MG	A075927 003	Dec 18, 2001
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AB	AUROBINDO PHARMA LTD	250MG	A200429 001	Nov 08, 2011
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AB		375MG	A200429 002	Nov 08, 2011
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AB		500MG	A200429 003	Nov 08, 2011
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AB	GLENMARK GENERICS	250MG	A078250 001	Mar 28, 2007
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AB		375MG	A078250 002	Mar 28, 2007
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AB		500MG	A078250 003	Mar 28, 2007
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AB	GRANULES	250MG	A074140 001	Dec 21, 1993
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AB		375MG	A074140 002	Dec 21, 1993
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AB		500MG	A074140 003	Dec 21, 1993
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AB	INVAGEN PHARMS	250MG	A091305 001	Aug 24, 2011
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AB		375MG	A091305 002	Aug 24, 2011
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AB		500MG	A091305 003	Aug 24, 2011
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AB	L PERRIGO CO	250MG	A077339 001	Apr 27, 2005
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AB		375MG	A077339 002	Apr 27, 2005
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AB		500MG	A077339 003	Apr 27, 2005
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AB	MARKSANS PHARMA	250MG	A091416 001	Feb 14, 2011
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AB		375MG	A091416 002	Feb 14, 2011
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AB		500MG	A091416 003	Feb 14, 2011
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AB	SCIEGEN PHARMS INC	250MG	A212517 001	Feb 21, 2020
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AB		375MG	A212517 002	Feb 21, 2020
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AB		500MG	A212517 003	Feb 21, 2020
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AB	TEVA	250MG	A074201 001	Dec 21, 1993
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AB		375MG	A074201 002	Dec 21, 1993
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AB		500MG	A074201 003	Dec 21, 1993
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AB	ZYDUS PHARMS USA	250MG	A078620 001	Jun 07, 2007
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AB		375MG	A078620 002	Jun 07, 2007
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AB		500MG	A078620 003	Jun 07, 2007
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PRESCRIPTION DRUG PRODUCT LIST

NAPROXEN

TABLET, DELAYED RELEASE;ORAL

EC-NAPROSYN

<u>AB</u>	<u>+</u> !	ATNAHS PHARMA US	<u>375MG</u>	<u>N020067</u>	<u>002</u>	Oct 14, 1994
<u>AB</u>	<u>+</u> !		<u>500MG</u>	<u>N020067</u>	<u>003</u>	Oct 14, 1994

NAPROXEN

<u>AB</u>		NUVO PHARMS INC	<u>375MG</u>	<u>A091432</u>	<u>001</u>	Sep 19, 2011
<u>AB</u>			<u>500MG</u>	<u>A091432</u>	<u>002</u>	Sep 19, 2011
<u>AB</u>		TEVA	<u>375MG</u>	<u>A075227</u>	<u>001</u>	Jun 30, 1998
<u>AB</u>			<u>500MG</u>	<u>A075227</u>	<u>002</u>	Jun 30, 1998

NAPROXEN SODIUM

TABLET;ORAL

ANAPROX DS

<u>AB</u>	<u>+</u> !	ATNAHS PHARMA US	<u>EQ 500MG BASE</u>	<u>N018164</u>	<u>003</u>	Sep 30, 1987
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NAPROXEN SODIUM

<u>AB</u>		AMNEAL PHARMS NY	<u>EQ 250MG BASE</u>	<u>A078432</u>	<u>001</u>	Apr 25, 2007
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A078432</u>	<u>002</u>	Apr 25, 2007
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 250MG BASE</u>	<u>A200629</u>	<u>001</u>	Oct 31, 2011
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A200629</u>	<u>002</u>	Oct 31, 2011
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 250MG BASE</u>	<u>A078486</u>	<u>001</u>	Jul 26, 2007
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A078486</u>	<u>002</u>	Jul 26, 2007
<u>AB</u>		GLENMARK PHARMS LTD	<u>EQ 250MG BASE</u>	<u>A078314</u>	<u>001</u>	Apr 27, 2007
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A078314</u>	<u>002</u>	Apr 27, 2007
<u>AB</u>		SCIEGEN PHARMS INC	<u>EQ 250MG BASE</u>	<u>A212199</u>	<u>001</u>	Oct 30, 2019
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A212199</u>	<u>002</u>	Oct 30, 2019

TABLET, EXTENDED RELEASE;ORAL

NAPRELAN

<u>AB</u>	<u>+</u>	ALMATICA	<u>EQ 375MG BASE</u>	<u>N020353</u>	<u>001</u>	Jan 05, 1996
<u>AB</u>	<u>+</u>		<u>EQ 500MG BASE</u>	<u>N020353</u>	<u>002</u>	Jan 05, 1996
<u>AB</u>	<u>+</u> !		<u>EQ 750MG BASE</u>	<u>N020353</u>	<u>003</u>	Jan 05, 1996

NAPROXEN SODIUM

<u>AB</u>		ACTAVIS LABS FL INC	<u>EQ 375MG BASE</u>	<u>A075416</u>	<u>002</u>	Apr 23, 2003
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A075416</u>	<u>001</u>	Aug 27, 2002
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A075416</u>	<u>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</u>	<u>001</u>	Feb 15, 2018
<u>AB</u>		RISING	<u>500MG;EQ 85MG BASE</u>	<u>A090872</u>	<u>001</u>	Sep 04, 2018
<u>AB</u>		SUN PHARM	<u>500MG;EQ 85MG BASE</u>	<u>A202803</u>	<u>001</u>	Jul 20, 2018

TREXIMET

<u>AB</u>	<u>+</u> !	CURRAX	<u>500MG;EQ 85MG BASE</u>	<u>N021926</u>	<u>001</u>	Apr 15, 2008
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NARATRIPTAN HYDROCHLORIDE

TABLET;ORAL

AMERGE

<u>AB</u>	<u>+</u>	GLAXOSMITHKLINE LLC	<u>EQ 1MG BASE</u>	<u>N020763</u>	<u>002</u>	Feb 10, 1998
<u>AB</u>	<u>+</u> !		<u>EQ 2.5MG BASE</u>	<u>N020763</u>	<u>001</u>	Feb 10, 1998

NARATRIPTAN

<u>AB</u>		HERITAGE PHARMS INC	<u>EQ 1MG BASE</u>	<u>A200502</u>	<u>001</u>	Feb 28, 2011
<u>AB</u>			<u>EQ 2.5MG BASE</u>	<u>A200502</u>	<u>002</u>	Feb 28, 2011
<u>AB</u>		HIKMA	<u>EQ 1MG BASE</u>	<u>A090381</u>	<u>001</u>	Jul 07, 2010
<u>AB</u>			<u>EQ 2.5MG BASE</u>	<u>A090381</u>	<u>002</u>	Jul 07, 2010
<u>AB</u>		ORBION PHARMS	<u>EQ 1MG BASE</u>	<u>A091441</u>	<u>001</u>	Apr 30, 2012
<u>AB</u>			<u>EQ 2.5MG BASE</u>	<u>A091441</u>	<u>002</u>	Apr 30, 2012
<u>AB</u>		PADAGIS US	<u>EQ 1MG BASE</u>	<u>A091326</u>	<u>001</u>	Jul 08, 2010
<u>AB</u>			<u>EQ 2.5MG BASE</u>	<u>A091326</u>	<u>002</u>	Jul 08, 2010
<u>AB</u>		SUN PHARM INDS LTD	<u>EQ 2.5MG BASE</u>	<u>A091552</u>	<u>001</u>	Feb 14, 2011

NATAMYCIN

SUSPENSION;OPHTHALMIC

NATACYN

<u>+</u> !	EYEVANCE	5%	<u>N050514</u>	<u>001</u>	
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NATEGLINIDE

TABLET;ORAL

NATEGLINIDE

<u>AB</u>		CADILA PHARMS LTD	<u>60MG</u>	<u>A206432</u>	<u>001</u>	Apr 19, 2019
<u>AB</u>			<u>120MG</u>	<u>A206432</u>	<u>002</u>	Apr 19, 2019
<u>AB</u>		DR REDDYS LABS LTD	<u>60MG</u>	<u>A077461</u>	<u>001</u>	Sep 09, 2009
<u>AB</u>			<u>120MG</u>	<u>A077461</u>	<u>002</u>	Sep 09, 2009

PRESCRIPTION DRUG PRODUCT LIST

NATEGLINIDE

TABLET; ORAL

NATEGLINIDE

<u>AB</u>	RISING	<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>	STRIDES PHARMA	<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>	ZYDUS PHARMS	<u>60MG</u>	<u>A205248 001</u>	Jul 06, 2016
<u>AB</u>	!	<u>120MG</u>	<u>A205248 002</u>	Jul 06, 2016

NEBIVOLOL HYDROCHLORIDE

TABLET; ORAL

BYSTOLIC

<u>AB</u>	+	ALLERGAN	<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

<u>AB</u>		ANI PHARMS	<u>EQ 2.5MG BASE</u>	<u>A203659 001</u>	Apr 16, 2015
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A203659 002</u>	Apr 16, 2015
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A203659 003</u>	Apr 16, 2015
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A203659 004</u>	Apr 16, 2015
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 2.5MG BASE</u>	<u>A211053 001</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A211053 002</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A211053 003</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A211053 004</u>	Dec 17, 2021
<u>AB</u>		CADILA PHARMS LTD	<u>EQ 2.5MG BASE</u>	<u>A208717 001</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A208717 002</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A208717 003</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A208717 004</u>	Dec 17, 2021
<u>AB</u>		HETERO LABS LTD III	<u>EQ 2.5MG BASE</u>	<u>A203825 001</u>	Nov 03, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A203825 002</u>	Nov 03, 2020
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A203825 003</u>	Nov 03, 2020
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A203825 004</u>	Nov 03, 2020
<u>AB</u>		INDCHEMIE HEALTH	<u>EQ 2.5MG BASE</u>	<u>A203828 001</u>	Jul 29, 2015
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A203828 002</u>	Jul 29, 2015
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A203828 003</u>	Jul 29, 2015
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A203828 004</u>	Jul 29, 2015
<u>AB</u>		REYOUNG	<u>EQ 2.5MG BASE</u>	<u>A212917 001</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A212917 002</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A212917 003</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A212917 004</u>	Dec 17, 2021
<u>AB</u>		TORRENT	<u>EQ 2.5MG BASE</u>	<u>A203966 001</u>	Mar 02, 2018
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A203966 002</u>	Mar 02, 2018
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A203966 003</u>	Mar 02, 2018
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A203966 004</u>	Mar 02, 2018

NEDOCROMIL SODIUM

SOLUTION/DROPS; OPHTHALMIC

ALOCRIIL

<u>AT</u>	+	!	ALLERGAN	<u>2%</u>	<u>N021009 001</u>	Dec 08, 1999
<u>AT</u>			AKORN	<u>2%</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

<u>AP</u>	+	!	NOVARTIS	<u>250MG/50ML (5MG/ML)</u>	<u>N021877 001</u>	Oct 28, 2005
<u>AP</u>			ZYDUS PHARMS	<u>250MG/50ML (5MG/ML)</u>	<u>A215037 001</u>	Nov 17, 2021

PRESCRIPTION DRUG PRODUCT LIST

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

AA	!	TEVA	500MG	A060304	001	
AA		XGEN PHARMS	500MG	A065220	001	Jul 28, 2006

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION; IRRIGATION

NEOMYCIN AND POLYMYXIN B SULFATE

AT		WATSON LABS	EQ 40MG BASE/ML; 200,000 UNITS/ML	A062664	001	Apr 08, 1986
AT		XGEN PHARMS	EQ 40MG BASE/ML; 200,000 UNITS/ML	A065106	001	Jan 31, 2006
AT			EQ 40MG BASE/ML; 200,000 UNITS/ML	A065108	001	Jan 31, 2006

NEOSPORIN G.U. IRRIGANT

AT	!	MONARCH PHARMS	EQ 40MG BASE/ML; 200,000 UNITS/ML	A060707	001	
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NEOSTIGMINE METHYLSULFATE

SOLUTION; INTRAVENOUS

BLOXIVERZ

AP	+	!	EXELA PHARMA	5MG/10ML (0.5MG/ML)	N204078	001	May 31, 2013
AP	+	!		10MG/10ML (1MG/ML)	N204078	002	May 31, 2013

NEOSTIGMINE METHYLSULFATE

AP		AM REGENT	5MG/10ML (0.5MG/ML)	A209182	001	May 04, 2018	
AP			10MG/10ML (1MG/ML)	A209182	002	May 04, 2018	
AP		AMNEAL	5MG/10ML (0.5MG/ML)	A210051	001	Jun 15, 2018	
AP			10MG/10ML (1MG/ML)	A210051	002	Jun 15, 2018	
AP		AMPHASTAR PHARMS INC	5MG/10ML (0.5MG/ML)	A209933	001	Sep 25, 2017	
AP			10MG/10ML (1MG/ML)	A209933	002	Sep 25, 2017	
AP		AMRING PHARMS	5MG/10ML (0.5MG/ML)	A210989	001	Aug 22, 2018	
AP			10MG/10ML (1MG/ML)	A210989	002	Aug 22, 2018	
AP		BE PHARMS	5MG/10ML (0.5MG/ML)	A212512	001	May 13, 2019	
AP			10MG/10ML (1MG/ML)	A212512	002	May 13, 2019	
AP		CAPLIN	5MG/10ML (0.5MG/ML)	A213074	001	Apr 20, 2021	
AP			10MG/10ML (1MG/ML)	A213074	002	Apr 20, 2021	
AP		DR REDDYS	5MG/10ML (0.5MG/ML)	A209135	001	Jul 10, 2018	
AP			10MG/10ML (1MG/ML)	A209135	002	Jul 10, 2018	
AP		EUROHLTH INTL SARL	5MG/10ML (0.5MG/ML)	A207042	001	Dec 28, 2015	
AP			10MG/10ML (1MG/ML)	A207042	002	Dec 28, 2015	
AP		GLAND PHARMA LTD	5MG/10ML (0.5MG/ML)	A212968	001	Oct 16, 2019	
AP			10MG/10ML (1MG/ML)	A212968	002	Oct 16, 2019	
AP		INDOCO	5MG/10ML (0.5MG/ML)	A210652	001	Oct 01, 2021	
AP			10MG/10ML (1MG/ML)	A210652	002	Oct 01, 2021	
AP		MEITHEAL	5MG/10ML (0.5MG/ML)	A212804	001	Apr 05, 2021	
AP			10MG/10ML (1MG/ML)	A212804	002	Apr 05, 2021	
AP		PAR STERILE PRODUCTS	5MG/10ML (0.5MG/ML)	A208405	001	Apr 26, 2017	
AP			10MG/10ML (1MG/ML)	A208405	002	Apr 26, 2017	
	+	!	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	0.3%	N203491	001	Oct 16, 2012
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NEVANAC

+	!	NOVARTIS	0.1%	N021862	001	Aug 19, 2005
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NERATINIB MALEATE

TABLET; ORAL

NERLYNX

+	!	PUMA BIOTECH	EQ 40MG BASE	N208051	001	Jul 17, 2017
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NETARSUDIL MESYLATE

SOLUTION/DROPS; OPHTHALMIC

RHOPRESSA

+	!	AERIE PHARMS INC	EQ 0.02% BASE	N208254	001	Dec 18, 2017
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PRESCRIPTION DRUG PRODUCT LIST

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, 2017VIRAMUNE**AA** +! BOEHRINGER **50MG/5ML** **N020933 001** Sep 11, 1998
INGELHEIM

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 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, 2012VIRAMUNE**AB** +! BOEHRINGER **200MG** **N020636 001** Jun 21, 1996
INGELHEIM

TABLET, EXTENDED RELEASE; ORAL

NEVIRAPINE**AB** ALVOGEN **100MG** **A204621 002** Nov 09, 2015**AB** **400MG** **A204621 001** Jul 10, 2015**AB** AUROBINDO PHARMA **100MG** **A208616 001** Nov 23, 2016
LTD**AB** **400MG** **A207698 001** Feb 28, 2017**AB** MACLEODS PHARMS LTD **400MG** **A206879 001** Oct 06, 2017**AB** MYLAN **400MG** **A205651 001** Oct 27, 2014**AB** SANDOZ INC **400MG** **A203411 001** Apr 03, 2014VIRAMUNE XR**AB** + BOEHRINGER **100MG** **N201152 002** Nov 08, 2012
INGELHEIM**AB** +! **400MG** **N201152 001** Mar 25, 2011NIACIN

TABLET; ORAL

NIACOR

! 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 **500MG** **A209236 001** Feb 01, 2018
LTD**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** BEIJING **500MG** **A214428 001** Nov 22, 2021**AB** **1GM** **A214428 002** Nov 22, 2021**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 PHARM **500MG** **A200484 001** Apr 23, 2014**AB** ! **750MG** **A201273 001** Apr 23, 2014**AB** ! **1GM** **A200484 002** Apr 23, 2014NICARDIPINE HYDROCHLORIDE

CAPSULE; ORAL

NICARDIPINE HYDROCHLORIDE**AB** ANI PHARMS **20MG** **A074670 001** Oct 28, 1996**AB** **30MG** **A074670 002** Oct 28, 1996**AB** EPIC PHARMA **20MG** **A074928 001** Mar 19, 1998**AB** ! **30MG** **A074928 002** Mar 19, 1998

PRESCRIPTION DRUG PRODUCT LIST

NICARDIPINE HYDROCHLORIDE

INJECTABLE; INJECTION

NICARDIPINE HYDROCHLORIDE

AP	!	AM REGENT	<u>25MG/10ML (2.5MG/ML)</u>	<u>A090534</u>	<u>001</u>	Nov 17, 2009
AP		EUGIA PHARMA	<u>25MG/10ML (2.5MG/ML)</u>	<u>A211121</u>	<u>001</u>	Apr 08, 2021
AP		SUN PHARM	<u>25MG/10ML (2.5MG/ML)</u>	<u>N078405</u>	<u>001</u>	Nov 17, 2009
	+	EXELA PHARMA SCIENCE	25MG/10ML (2.5MG/ML)	N022276	001	Jul 24, 2008

INJECTABLE; INTRAVENOUS

CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER

+	!	CHIESI	40MG/200ML (0.2MG/ML)	N019734	004	Nov 07, 2008
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CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER

+	!	CHIESI	20MG/200ML (0.1MG/ML)	N019734	003	Jul 31, 2008
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CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER

+	!	CHIESI	20MG/200ML (0.1MG/ML)	N019734	002	Jul 31, 2008
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NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE

+	!	EXELA PHARMA SCIENCE	20MG/200ML (0.1MG/ML)	N022276	002	Apr 07, 2016
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+	!		40MG/200ML (0.2MG/ML)	N022276	003	Apr 07, 2016
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NICOTINE

INHALANT; ORAL

NICOTROL

+	!	PFIZER	4MG/CARTRIDGE	N020714	001	May 02, 1997
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SPRAY, METERED; NASAL

NICOTROL

+	!	PFIZER INC	0.5MG/SPRAY	N020385	001	Mar 22, 1996
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NIFEDIPINE

CAPSULE; ORAL

NIFEDIPINE

AB		ACELLA	<u>10MG</u>	<u>A072781</u>	<u>001</u>	Jul 30, 1993
AB		ACTAVIS ELIZABETH	<u>10MG</u>	<u>A072579</u>	<u>001</u>	Jan 08, 1991
AB			<u>20MG</u>	<u>A072556</u>	<u>001</u>	Sep 20, 1990
AB		HERITAGE PHARMA	<u>10MG</u>	<u>A202644</u>	<u>001</u>	Apr 25, 2013
AB			<u>20MG</u>	<u>A202644</u>	<u>002</u>	Apr 25, 2013
AB		LEADING PHARMA LLC	<u>10MG</u>	<u>A073250</u>	<u>001</u>	Oct 08, 1991
AB			<u>20MG</u>	<u>A074045</u>	<u>001</u>	Apr 30, 1992

PROCARDIA

AB	+	!	PFIZER	<u>10MG</u>	<u>N018482</u>	<u>001</u>
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TABLET, EXTENDED RELEASE; ORAL

NIFEDIPINE

AB1		AUROBINDO PHARMA LTD	<u>30MG</u>	<u>A213361</u>	<u>001</u>	Jul 19, 2021
AB1			<u>60MG</u>	<u>A213361</u>	<u>002</u>	Jul 19, 2021
AB1			<u>90MG</u>	<u>A213361</u>	<u>003</u>	Jul 19, 2021
AB1		NOVAST LABS	<u>30MG</u>	<u>A202987</u>	<u>001</u>	Aug 25, 2016
AB1	!		<u>60MG</u>	<u>A202987</u>	<u>002</u>	Aug 25, 2016
AB1	!		<u>90MG</u>	<u>A202987</u>	<u>003</u>	Aug 25, 2016
AB1		VALEANT PHARMS NORTH	<u>30MG</u>	<u>A075269</u>	<u>001</u>	Dec 04, 2000
AB1			<u>60MG</u>	<u>A075269</u>	<u>002</u>	Dec 04, 2000
AB1			<u>90MG</u>	<u>A076070</u>	<u>001</u>	Aug 16, 2002
AB1		ZYDUS PHARMS	<u>30MG</u>	<u>A210184</u>	<u>001</u>	Jun 29, 2018
AB1			<u>60MG</u>	<u>A210184</u>	<u>002</u>	Jun 29, 2018
AB1			<u>90MG</u>	<u>A210184</u>	<u>003</u>	Jun 29, 2018
AB2		ELITE PHARM SOLUTION	<u>90MG</u>	<u>A212016</u>	<u>001</u>	Nov 18, 2020
AB2		NOVAST LABS	<u>30MG</u>	<u>A210614</u>	<u>001</u>	Mar 12, 2019
AB2			<u>60MG</u>	<u>A210614</u>	<u>002</u>	Mar 12, 2019
AB2			<u>90MG</u>	<u>A210614</u>	<u>003</u>	Mar 12, 2019
AB2		OSMOTICA PHARM US	<u>30MG</u>	<u>A077127</u>	<u>001</u>	Nov 21, 2005
AB2			<u>60MG</u>	<u>A077127</u>	<u>002</u>	Nov 21, 2005
AB2			<u>90MG</u>	<u>A077127</u>	<u>003</u>	Oct 03, 2007
AB2		SPIIL	<u>30MG</u>	<u>A210838</u>	<u>001</u>	Apr 16, 2019
AB2			<u>60MG</u>	<u>A210838</u>	<u>002</u>	Apr 16, 2019
AB2			<u>90MG</u>	<u>A210838</u>	<u>003</u>	Apr 16, 2019
AB2		TWI PHARMS	<u>30MG</u>	<u>A203126</u>	<u>001</u>	Apr 03, 2014
AB2			<u>60MG</u>	<u>A203126</u>	<u>002</u>	Apr 03, 2014
AB2			<u>90MG</u>	<u>A203126</u>	<u>003</u>	Apr 03, 2014
AB2		VALEANT PHARMS NORTH	<u>30MG</u>	<u>A075289</u>	<u>002</u>	Feb 06, 2001
AB2			<u>60MG</u>	<u>A075289</u>	<u>001</u>	Sep 27, 2000
AB2		ZYDUS PHARMS	<u>30MG</u>	<u>A210012</u>	<u>001</u>	Dec 19, 2017
AB2			<u>60MG</u>	<u>A210012</u>	<u>002</u>	Dec 19, 2017

PRESCRIPTION DRUG PRODUCT LIST

NIFEDIPINE

TABLET, EXTENDED RELEASE;ORAL

NIFEDIPINE

AB2		90MG	A210012 003	Dec 19, 2017
	<u>PROCARDIA XL</u>			
AB2	+	PFIZER	N019684 001	Sep 06, 1989
AB2	+		N019684 002	Sep 06, 1989
AB2	+		N019684 003	Sep 06, 1989

NIFURTIMOX

TABLET;ORAL

LAMPIT

+	BAYER HEALTHCARE	30MG	N213464 001	Aug 06, 2020
+	!	120MG	N213464 002	Aug 06, 2020

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	150MG	N020169 002	Apr 30, 1999
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NILUTAMIDE

AB		ANI PHARMS	150MG	A207631 001	Jul 15, 2016
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NIMODIPINE

CAPSULE;ORAL

NIMODIPINE

AB	!	BIONPHARMA INC	30MG	A076740 001	Jan 17, 2008
AB		HERITAGE PHARMS INC	30MG	A077811 001	May 02, 2007
AB		THEPHARMANETWORK LLC	30MG	A090103 001	Apr 07, 2014

SOLUTION;ORAL

NYMALIZE

+	ARBOR PHARMS LLC	6MG/ML	N203340 002	Apr 08, 2020
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NINTEDANIB ESYLATE

CAPSULE;ORAL

OFEV

+	BOEHRINGER INGELHEIM	EQ 100MG BASE	N205832 001	Oct 15, 2014
+	!	EQ 150MG BASE	N205832 002	Oct 15, 2014

NIRAPARIB TOSYLATE

CAPSULE;ORAL

ZEJULA

+	GLAXOSMITHKLINE	EQ 100MG BASE	N208447 001	Mar 27, 2017
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NISOLDIPINE

TABLET, EXTENDED RELEASE;ORAL

NISOLDIPINE

AB		MYLAN	8.5MG	A091001 001	Jan 26, 2011
AB			17MG	A091001 002	Jan 26, 2011
AB			34MG	A091001 004	Jan 26, 2011

SULAR

AB	+	COVIS	8.5MG	N020356 008	Jan 02, 2008
AB	+		17MG	N020356 007	Jan 02, 2008
AB	+		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
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TABLET;ORAL

ALINIA

AB	+	ROMARK	500MG	N021497 001	Jul 21, 2004
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PRESCRIPTION DRUG PRODUCT LIST

NITAZOXANIDE

TABLET; ORAL

NITAZOXANIDE

AB	RISING	500MG	A213820 001	Nov 27, 2020
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NITISINONE

CAPSULE; ORAL

NITISINONE

AB	NOVITIUM PHARMA	2MG	A211041 001	Aug 26, 2019
AB		5MG	A211041 002	Aug 26, 2019
AB		10MG	A211041 003	Aug 26, 2019

ORFADIN

AB	+	SWEDISH ORPHAN	2MG	N021232 001	Jan 18, 2002
AB	+		5MG	N021232 002	Jan 18, 2002
AB	+		10MG	N021232 003	Jan 18, 2002
	+	!	20MG	N021232 004	Jun 13, 2016

SUSPENSION; ORAL

ORFADIN

	+	!	SWEDISH ORPHAN	4MG/ML	N206356 001	Apr 22, 2016
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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

INOMAX

AA	+	!	MALLINCKRODT HOSP	800PPM	N020845 003	Dec 23, 1999
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NOXIVENT

AA			PRAXAIR	800PPM	A207141 002	Oct 02, 2018
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DISTRIBUTION

GENOSYL

	+	!	VERO BIOTECH	800PPM	N202860 001	Dec 20, 2019
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NOXIVENT

			PRAXAIR	100PPM	A207141 001	Oct 02, 2018
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DISTRIBUTION

NITROFURANTOIN

SUSPENSION; ORAL

FURADANTIN

AB	+	!	CASPER PHARMA LLC	25MG/5ML	N009175 001	
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NITROFURANTOIN

AB			ACTAVIS MID	25MG/5ML	A205180 001	May 03, 2016
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ATLANTIC

AB			AMNEAL PHARMS	25MG/5ML	A201679 001	May 11, 2011
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AB			NOSTRUM LABS INC	25MG/5ML	A201355 001	Aug 14, 2013
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AB			NOVEL LABS INC	25MG/5ML	A201693 001	Sep 08, 2014
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NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACRODANTIN

AB	+		ALMATICA	25MG	N016620 003	
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AB	+			50MG	N016620 001	
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AB	+	!		100MG	N016620 002	
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NITROFURANTOIN

AB			ACTAVIS LABS FL INC	25MG	A091095 001	Jun 18, 2015
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AB				50MG	A091095 002	Jun 18, 2015
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AB				100MG	A091095 003	Jun 18, 2015
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AB			ALEMBIC PHARMS LTD	25MG	A211935 001	Jun 25, 2021
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AB				50MG	A211935 002	Jun 25, 2021
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AB				100MG	A211935 003	Jun 25, 2021
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AB			IMPAX LABS INC	50MG	A073671 001	Jan 28, 1993
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AB				100MG	A073652 001	Jan 28, 1993
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AB			NOVEL LABS INC	50MG	A203233 001	Jul 09, 2018
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AB				100MG	A203233 002	Jul 09, 2018
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AB			SUN PHARM	25MG	A201722 001	Feb 16, 2016
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INDUSTRIES

AB				50MG	A201722 002	Feb 16, 2016
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AB				100MG	A201722 003	Feb 16, 2016
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AB			ZYDUS PHARMS	50MG	A205005 001	Dec 12, 2017
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AB				100MG	A205005 002	Dec 12, 2017
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PRESCRIPTION DRUG PRODUCT LIST

NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACROBID

<u>AB</u>	<u>+!</u>	<u>ALMATICA</u>	<u>75MG;25MG</u>	<u>N020064</u>	<u>001</u>	Dec 24, 1991
<u>NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)</u>						
<u>AB</u>		<u>AMNEAL PHARMS</u>	<u>75MG;25MG</u>	<u>A207372</u>	<u>001</u>	May 15, 2017
<u>AB</u>		<u>SANDOZ</u>	<u>75MG;25MG</u>	<u>A077066</u>	<u>001</u>	Apr 05, 2005
<u>AB</u>		<u>SUNNY</u>	<u>75MG;25MG</u>	<u>A208516</u>	<u>001</u>	May 24, 2018
<u>AB</u>		<u>WATSON LABS INC</u>	<u>75MG;25MG</u>	<u>A202250</u>	<u>001</u>	Jul 08, 2015

NITROGLYCERIN

AEROSOL, METERED; SUBLINGUAL

NITROMIST+! EVUS 0.4MG/SPRAYN021780 001 Nov 02, 2006

FILM, EXTENDED RELEASE; TRANSDERMAL

NITROGLYCERIN

<u>AB2</u>		<u>HERCON PHARM</u>	<u>0.1MG/HR</u>	<u>A089885</u>	<u>002</u>	Oct 30, 2017
<u>AB2</u>			<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>	<u>!</u>	<u>MYLAN TECHNOLOGIES</u>	<u>0.1MG/HR</u>	<u>A074559</u>	<u>004</u>	Feb 06, 1998
<u>AB2</u>	<u>!</u>		<u>0.2MG/HR</u>	<u>A074559</u>	<u>003</u>	Aug 30, 1996
<u>AB2</u>	<u>!</u>		<u>0.4MG/HR</u>	<u>A074559</u>	<u>002</u>	Aug 30, 1996
<u>AB2</u>	<u>!</u>		<u>0.6MG/HR</u>	<u>A074559</u>	<u>001</u>	Aug 30, 1996
<u>NITRO-DUR</u>						
	<u>+!</u>	<u>USPHARMA</u>	<u>0.1MG/HR</u>	<u>N020145</u>	<u>001</u>	Apr 04, 1995
	<u>+!</u>		<u>0.2MG/HR</u>	<u>N020145</u>	<u>002</u>	Apr 04, 1995
	<u>+!</u>		<u>0.3MG/HR</u>	<u>N020145</u>	<u>003</u>	Apr 04, 1995
	<u>+!</u>		<u>0.4MG/HR</u>	<u>N020145</u>	<u>004</u>	Apr 04, 1995
	<u>+!</u>		<u>0.6MG/HR</u>	<u>N020145</u>	<u>005</u>	Apr 04, 1995
	<u>+!</u>		<u>0.8MG/HR</u>	<u>N020145</u>	<u>006</u>	Apr 04, 1995

INJECTABLE; INJECTION

NITROGLYCERIN IN DEXTROSE 5%

<u>AP</u>	<u>+!</u>	<u>BAXTER HLTHCARE</u>	<u>10MG/100ML</u>	<u>N019970</u>	<u>001</u>	Dec 29, 1989
<u>AP</u>	<u>+!</u>		<u>20MG/100ML</u>	<u>N019970</u>	<u>002</u>	Dec 29, 1989
<u>AP</u>	<u>+!</u>		<u>40MG/100ML</u>	<u>N019970</u>	<u>003</u>	Dec 29, 1989
<u>NITROGLYCERIN</u>						
	<u>!</u>	<u>AM REGENT</u>	<u>5MG/ML</u>	<u>A072034</u>	<u>001</u>	May 24, 1988
<u>OINTMENT; INTRA-ANAL</u>						
<u>RECTIV</u>						
	<u>+!</u>	<u>ALLERGAN</u>	<u>0.4%</u>	<u>N021359</u>	<u>001</u>	Jun 21, 2011
<u>OINTMENT; TRANSDERMAL</u>						
<u>NITROGLYCERIN</u>						
	<u>!</u>	<u>FOUGERA PHARMS INC</u>	<u>2%</u>	<u>A087355</u>	<u>001</u>	Jul 08, 1988
<u>SPRAY, METERED; SUBLINGUAL</u>						

NITROGLYCERIN

<u>AB</u>		<u>PADAGIS ISRAEL</u>	<u>0.4MG/SPRAY</u>	<u>A091496</u>	<u>001</u>	Sep 20, 2013
<u>NITROLINGUAL PUMPSPRAY</u>						
<u>AB</u>	<u>+!</u>	<u>POHL BOSKAMP</u>	<u>0.4MG/SPRAY</u>	<u>N018705</u>	<u>002</u>	Jan 10, 1997

TABLET; SUBLINGUAL

NITROGLYCERIN

<u>AB</u>		<u>ALVOGEN</u>	<u>0.3MG</u>	<u>A211604</u>	<u>001</u>	Apr 30, 2019
<u>AB</u>			<u>0.4MG</u>	<u>A211604</u>	<u>002</u>	Apr 30, 2019
<u>AB</u>			<u>0.6MG</u>	<u>A211604</u>	<u>003</u>	Apr 30, 2019
<u>AB</u>		<u>DR REDDYS</u>	<u>0.3MG</u>	<u>A208191</u>	<u>001</u>	Aug 26, 2016
<u>AB</u>			<u>0.4MG</u>	<u>A208191</u>	<u>002</u>	Aug 26, 2016
<u>AB</u>			<u>0.6MG</u>	<u>A208191</u>	<u>003</u>	Aug 26, 2016
<u>AB</u>		<u>GLENMARK PHARMS SA</u>	<u>0.3MG</u>	<u>A206391</u>	<u>001</u>	Sep 19, 2017
<u>AB</u>			<u>0.4MG</u>	<u>A206391</u>	<u>002</u>	Sep 19, 2017
<u>AB</u>			<u>0.6MG</u>	<u>A206391</u>	<u>003</u>	Sep 19, 2017
<u>AB</u>		<u>RUBICON</u>	<u>0.3MG</u>	<u>A209779</u>	<u>001</u>	May 03, 2021
<u>AB</u>			<u>0.4MG</u>	<u>A209779</u>	<u>002</u>	May 03, 2021
<u>AB</u>			<u>0.6MG</u>	<u>A209779</u>	<u>003</u>	May 03, 2021

NITROSTAT

<u>AB</u>	<u>+</u>	<u>UPJOHN</u>	<u>0.3MG</u>	<u>N021134</u>	<u>001</u>	May 01, 2000
<u>AB</u>	<u>+</u>		<u>0.4MG</u>	<u>N021134</u>	<u>002</u>	May 01, 2000
<u>AB</u>	<u>+!</u>		<u>0.6MG</u>	<u>N021134</u>	<u>003</u>	May 01, 2000

PRESCRIPTION DRUG PRODUCT LIST

NIZATIDINE

CAPSULE; ORAL

NIZATIDINE

<u>AB</u>	DR REDDYS LABS LTD	<u>150MG</u>	<u>A077314 001</u>	Sep 15, 2005
<u>AB</u>		<u>300MG</u>	<u>A077314 002</u>	Sep 15, 2005
<u>AB</u>	GLENMARK GENERICS	<u>150MG</u>	<u>A090618 001</u>	Jul 15, 2011
<u>AB</u>		<u>300MG</u>	<u>A090618 002</u>	Jul 15, 2011
<u>AB</u>	SANDOZ	<u>150MG</u>	<u>A076178 001</u>	Jul 05, 2002
<u>AB</u>		<u>300MG</u>	<u>A076178 002</u>	Jul 05, 2002
<u>AB</u>	WATSON LABS	<u>150MG</u>	<u>A075616 001</u>	Jul 09, 2002
<u>AB</u>	!	<u>300MG</u>	<u>A075616 002</u>	Jul 09, 2002

SOLUTION; ORAL

NIZATIDINE

!	AMNEAL PHARMS	15MG/ML	A090576 001	Nov 18, 2009
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NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

LEVOPHED

<u>AP</u>	+!	HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>N007513 001</u>	
<u>NOREPINEPHRINE BITARTRATE</u>					
<u>AP</u>		AMNEAL	<u>EQ 1MG BASE/ML</u>	<u>A210839 001</u>	Dec 17, 2018
<u>AP</u>		BAXTER HLTHCARE CORP	<u>EQ 1MG BASE/ML</u>	<u>A040859 001</u>	Mar 27, 2012
<u>AP</u>		BRECKENRIDGE	<u>EQ 1MG BASE/ML</u>	<u>A214455 001</u>	Jan 22, 2021
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 1MG BASE/ML</u>	<u>A211382 001</u>	Nov 03, 2020
<u>AP</u>		GLAND PHARMA LTD	<u>EQ 1MG BASE/ML</u>	<u>A214323 001</u>	May 06, 2021
<u>AP</u>		HIKMA	<u>EQ 1MG BASE/ML</u>	<u>A040462 001</u>	Oct 31, 2003
<u>AP</u>			<u>EQ 1MG BASE/ML</u>	<u>A203662 001</u>	Nov 07, 2018
<u>AP</u>		MYLAN LABS LTD	<u>EQ 1MG BASE/ML</u>	<u>A211242 001</u>	Oct 04, 2018
<u>AP</u>		SANDOZ INC	<u>EQ 1MG BASE/ML</u>	<u>A211359 001</u>	Oct 18, 2018
<u>AP</u>		SUN PHARM	<u>EQ 1MG BASE/ML</u>	<u>A211980 001</u>	Jan 29, 2021
<u>AP</u>		TEVA PHARMS USA	<u>EQ 1MG BASE/ML</u>	<u>A040455 001</u>	Mar 03, 2003

SOLUTION; INTRAVENOUS

NOREPINEPHRINE BITARTRATE IN 5% DEXTROSE

+!	BAXTER HLTHCARE CORP	EQ 4MG BASE/250ML (EQ 16MCG BASE/ML)	N214313 001	Jan 15, 2021
+!		EQ 8MG BASE/250ML (EQ 32MCG BASE/ML)	N214313 002	Jan 15, 2021

NORETHINDRONE

TABLET; ORAL-28

CAMILA

<u>AB1</u>	MAYNE PHARMA	<u>0.35MG</u>	<u>A076177 001</u>	Oct 21, 2002
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HEATHER

<u>AB1</u>	GLENMARK GENERICS	<u>0.35MG</u>	<u>A090454 001</u>	Apr 23, 2010
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INCASSIA

<u>AB1</u>	AUROBINDO PHARMA LTD	<u>0.35MG</u>	<u>A207304 001</u>	Sep 23, 2016
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NOR-QD

<u>AB1</u>	+!	TEVA BRANDED PHARM	<u>0.35MG</u>	<u>N017060 001</u>	
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NORETHINDRONE

<u>AB1</u>	AMNEAL PHARMS	<u>0.35MG</u>	<u>A202260 001</u>	Aug 01, 2013
<u>AB1</u>	LUPIN LTD	<u>0.35MG</u>	<u>A091325 001</u>	Sep 19, 2011
<u>AB1</u>	MYLAN LABS LTD	<u>0.35MG</u>	<u>A201483 001</u>	Jun 24, 2013
<u>AB1</u>	NAARI PTE LTD	<u>0.35MG</u>	<u>A206807 001</u>	Dec 13, 2016
<u>AB1</u>	NOVAST LABS	<u>0.35MG</u>	<u>A202014 001</u>	Sep 13, 2013

ERRIN

<u>AB2</u>	MAYNE PHARMA	<u>0.35MG</u>	<u>A076225 001</u>	Oct 21, 2002
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JENCYCLA

<u>AB2</u>	LUPIN LTD	<u>0.35MG</u>	<u>A091323 001</u>	Mar 28, 2013
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NORETHINDRONE

<u>AB2</u>	GLENMARK GENERICS	<u>0.35MG</u>	<u>A091209 001</u>	Jul 22, 2010	
<u>AB2</u>	!	MYLAN LABS LTD	<u>0.35MG</u>	<u>A200980 001</u>	Jun 12, 2013
<u>AB2</u>	NOVAST LABS	<u>0.35MG</u>	<u>A200961 001</u>	Sep 13, 2013	

NORETHINDRONE ACETATE

TABLET; ORAL

NORETHINDRONE ACETATE

<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A200275 001</u>	Jul 30, 2012	
<u>AB</u>	!	BARR	<u>5MG</u>	<u>A075951 001</u>	May 25, 2001
<u>AB</u>	GLENMARK GENERICS	<u>5MG</u>	<u>A091090 001</u>	Jul 21, 2010	
<u>AB</u>	MYLAN LABS LTD	<u>5MG</u>	<u>A205278 001</u>	Nov 10, 2016	
<u>AB</u>	NOVAST LABS	<u>5MG</u>	<u>A206490 001</u>	Nov 05, 2018	

PRESCRIPTION DRUG PRODUCT LIST

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

NORTRIPTYLINE HYDROCHLORIDE

<u>AB</u>	MAYNE PHARMA	<u>EQ 10MG BASE</u>	<u>A073556 002</u>	Mar 30, 1992
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A073556 003</u>	Mar 30, 1992
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A073556 004</u>	Mar 30, 1992
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A073556 001</u>	Mar 30, 1992
<u>AB</u>	TARO	<u>EQ 10MG BASE</u>	<u>A075520 004</u>	May 08, 2000
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A075520 003</u>	May 08, 2000
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A075520 001</u>	May 08, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A075520 002</u>	May 08, 2000
<u>AB</u>	TEVA	<u>EQ 10MG BASE</u>	<u>A074132 001</u>	Mar 27, 1995
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A074132 002</u>	Mar 27, 1995
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A074132 003</u>	Mar 27, 1995
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A074132 004</u>	Mar 27, 1995
<u>PAMELOR</u>				
<u>AB</u>	+ SPECGX LLC	<u>EQ 10MG BASE</u>	<u>N018013 001</u>	
<u>AB</u>	+	<u>EQ 25MG BASE</u>	<u>N018013 002</u>	
<u>AB</u>	+	<u>EQ 50MG BASE</u>	<u>N018013 004</u>	
<u>AB</u>	+	<u>EQ 75MG BASE</u>	<u>N018013 003</u>	

SOLUTION; ORAL

NORTRIPTYLINE HYDROCHLORIDE

<u>AA</u>	! PHARM ASSOC	<u>EQ 10MG BASE/5ML</u>	<u>A075606 001</u>	Aug 28, 2000
<u>AA</u>	TARO	<u>EQ 10MG BASE/5ML</u>	<u>A077965 001</u>	Jun 20, 2006

NUSINERSEN SODIUMSOLUTION; INTRATHECAL
SPINRAZA

+	! BIOGEN IDEC	EQ 12MG BASE/5ML (EQ 2.4MG BASE/ML)	N209531 001	Dec 23, 2016
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NYSTATIN

CREAM; TOPICAL

NYSTATIN

<u>AT</u>	ACTAVIS MID ATLANTIC	<u>100,000 UNITS/GM</u>	<u>A062949 001</u>	Jun 13, 1988
<u>AT</u>	COSETTE	<u>100,000 UNITS/GM</u>	<u>A061966 001</u>	
<u>AT</u>	CROWN LABS INC	<u>100,000 UNITS/GM</u>	<u>A207733 001</u>	Sep 26, 2017
<u>AT</u>	FOUGERA PHARMS	<u>100,000 UNITS/GM</u>	<u>A062129 001</u>	
<u>AT</u>	MACLEODS PHARMS LTD	<u>100,000 UNITS/GM</u>	<u>A213566 001</u>	Aug 10, 2021
<u>AT</u>	PADAGIS US	<u>100,000 UNITS/GM</u>	<u>A062225 001</u>	
<u>AT</u>	! TARO	<u>100,000 UNITS/GM</u>	<u>A064022 001</u>	Jan 29, 1993
<u>AT</u>	TORRENT	<u>100,000 UNITS/GM</u>	<u>A212557 001</u>	Jul 24, 2019

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>	CADILA	<u>100,000 UNITS/GM</u>	<u>A207767 001</u>	May 25, 2018
<u>AT</u>	COSETTE	<u>100,000 UNITS/GM</u>	<u>A209114 001</u>	Oct 06, 2017
<u>AT</u>	! FOUGERA PHARMS	<u>100,000 UNITS/GM</u>	<u>A062124 002</u>	Sep 23, 1982
<u>AT</u>	LYNE	<u>100,000 UNITS/GM</u>	<u>A209082 001</u>	May 21, 2018
<u>AT</u>	MACLEODS PHARMS LTD	<u>100,000 UNITS/GM</u>	<u>A213826 001</u>	Jan 14, 2021
<u>AT</u>	PADAGIS US	<u>100,000 UNITS/GM</u>	<u>A062472 001</u>	Feb 13, 1984
<u>AT</u>	TORRENT	<u>100,000 UNITS/GM</u>	<u>A211838 001</u>	Jan 28, 2019

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>	LUPIN	<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>	UPSHER SMITH LABS	<u>100,000 UNITS/GM</u>	<u>A065183 001</u>	May 03, 2005
<u>AT</u>	XGEN PHARMS	<u>100,000 UNITS/GM</u>	<u>A065175 001</u>	Dec 17, 2004
<u>AT</u>	ZYDUS PHARMS	<u>100,000 UNITS/GM</u>	<u>A208581 001</u>	Jun 08, 2017

NYSTOP

<u>AT</u>	PADAGIS US	<u>100,000 UNITS/GM</u>	<u>A064118 001</u>	Aug 16, 1996
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SUSPENSION; ORAL

NYSTATIN

<u>AA</u>	AKORN	<u>100,000 UNITS/ML</u>	<u>A064042 001</u>	Feb 28, 1994
<u>AA</u>	DASTECH GENERICS	<u>100,000 UNITS/ML</u>	<u>A062832 001</u>	Dec 27, 1991
<u>AA</u>	FOUGERA PHARMS INC	<u>100,000 UNITS/ML</u>	<u>A062517 001</u>	Jun 07, 1984
<u>AA</u>	GENUS	<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	<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>		<u>100,000 UNITS/ML</u>	<u>A065422 001</u>	Mar 07, 2011

PRESCRIPTION DRUG PRODUCT LIST

NYSTATIN

SUSPENSION; ORAL

NYSTATIN

AA	!	WOCKHARDT BIO AG	100,000 UNITS/ML	A062512	001	Oct 29, 1984
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TABLET; ORAL

NYSTATIN

AA		HERITAGE PHARMS INC	500,000 UNITS	A062474	001	Dec 22, 1983
AA		SUN PHARM	500,000 UNITS	A062838	001	Dec 22, 1988
		INDUSTRIES				
AA	!	TEVA	500,000 UNITS	A062506	001	Jan 16, 1984

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYKACET

AT		COSETTE	100,000 UNITS/GM;0.1%	A062367	001	May 28, 1985
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NYSTATIN AND TRIAMCINOLONE ACETONIDE

AT		ALEOR	100,000 UNITS/GM;0.1%	A214090	001	Mar 31, 2021
		DERMACEUTICALS				
AT		AMNEAL	100,000 UNITS/GM;0.1%	A209990	001	Feb 15, 2018
AT		CROWN LABS INC	100,000 UNITS/GM;0.1%	A207730	001	Dec 26, 2017
AT		DR REDDYS	100,000 UNITS/GM;0.1%	A208326	001	Oct 26, 2016
AT		FOUGERA PHARMS INC	100,000 UNITS/GM;0.1%	A062599	001	Oct 08, 1985
AT		GLENMARK PHARMS LTD	100,000 UNITS/GM;0.1%	A208136	001	Oct 24, 2016
AT		LUPIN LTD	100,000 UNITS/GM;0.1%	A208205	001	May 31, 2018
AT		PADAGIS ISRAEL	100,000 UNITS/GM;0.1%	A208479	001	Aug 14, 2017
AT	!	TARO	100,000 UNITS/GM;0.1%	A062364	001	Dec 22, 1987
AT		TORRENT	100,000 UNITS/GM;0.1%	A213142	001	Jul 14, 2020

OINTMENT; TOPICAL

MYKACET

AT		COSETTE	100,000 UNITS/GM;0.1%	A062733	001	Mar 06, 1987
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NYSTATIN AND TRIAMCINOLONE ACETONIDE

AT		AKORN	100,000 UNITS/GM;0.1%	A207217	001	Aug 04, 2017
AT		CADILA	100,000 UNITS/GM;0.1%	A207764	001	Nov 08, 2018
AT		DR REDDYS	100,000 UNITS/GM;0.1%	A207741	001	Jan 31, 2017
AT		FOUGERA PHARMS INC	100,000 UNITS/GM;0.1%	A062602	001	Oct 09, 1985
AT		GLENMARK PHARMS LTD	100,000 UNITS/GM;0.1%	A208300	001	Jun 23, 2016
AT		MACLEODS PHARMS LTD	100,000 UNITS/GM;0.1%	A214751	001	Aug 17, 2021
AT		PADAGIS ISRAEL	100,000 UNITS/GM;0.1%	A207380	001	Dec 20, 2016
AT		RISING	100,000 UNITS/GM;0.1%	A206785	001	Dec 29, 2016
AT		STRIDES PHARMA	100,000 UNITS/GM;0.1%	A210077	001	Jan 29, 2018
AT	!	TARO	100,000 UNITS/GM;0.1%	A063305	001	Mar 29, 1993
AT		TEBIGENT	100,000 UNITS/GM;0.1%	A208287	001	Dec 30, 2016

OBETICHOIC ACID

TABLET; ORAL

OCALIVA

	+	INTERCEPT PHARMS	5MG	N207999	001	May 27, 2016
		INC				
	+!		10MG	N207999	002	May 27, 2016

OCTREOTIDE ACETATE

CAPSULE, DELAYED RELEASE; ORAL

MYCAPSSA

	+!	CHIASMA	EQ 20MG BASE	N208232	001	Jun 26, 2020
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INJECTABLE; INJECTION

OCTREOTIDE ACETATE

AP		FRESENIUS KABI USA	EQ 0.2MG BASE/ML	A077450	001	Feb 10, 2006
AP			EQ 1MG BASE/ML	A077450	002	Feb 10, 2006
AP		SAGENT PHARMS INC	EQ 0.2MG BASE/ML	A091041	001	Nov 12, 2013
AP			EQ 1MG BASE/ML	A091041	002	Nov 12, 2013
AP		TEVA PHARMS USA	EQ 0.05MG BASE/ML	A075957	001	Oct 03, 2005
AP			EQ 0.1MG BASE/ML	A075957	002	Oct 03, 2005
AP			EQ 0.2MG BASE/ML	A075959	001	Nov 21, 2005
AP			EQ 0.5MG BASE/ML	A075957	003	Oct 03, 2005
AP			EQ 1MG BASE/ML	A075959	002	Nov 21, 2005
AP	!	WEST-WARD PHARMS	EQ 0.2MG BASE/ML	A076330	001	Apr 08, 2005
		INT				
AP	!		EQ 1MG BASE/ML	A076330	002	Apr 08, 2005

OCTREOTIDE ACETATE (PRESERVATIVE FREE)

AP		FRESENIUS KABI USA	EQ 0.05MG BASE/ML	A077457	001	Feb 10, 2006
AP			EQ 0.1MG BASE/ML	A077457	002	Feb 10, 2006
AP			EQ 0.5MG BASE/ML	A077457	003	Feb 10, 2006
AP		MYLAN INSTITUTIONAL	EQ 0.05MG BASE/ML	A079198	001	Feb 10, 2011
AP			EQ 0.1MG BASE/ML	A079198	002	Feb 10, 2011
AP			EQ 0.5MG BASE/ML	A079198	003	Feb 10, 2011

PRESCRIPTION DRUG PRODUCT LIST

OCTREOTIDE ACETATE

INJECTABLE; INJECTION

OCTREOTIDE ACETATE (PRESERVATIVE FREE)

<u>AP</u>	SAGENT PHARMS INC	<u>EQ 0.05MG BASE/ML</u>	<u>A090834 001</u>	Nov 12, 2013
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A090834 002</u>	Nov 12, 2013
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A090834 003</u>	Nov 12, 2013
<u>AP</u>	! WEST-WARD PHARMS INT	<u>EQ 0.05MG BASE/ML</u>	<u>A076313 001</u>	Mar 28, 2005
<u>AP</u>	!	<u>EQ 0.1MG BASE/ML</u>	<u>A076313 003</u>	Mar 28, 2005
<u>AP</u>	!	<u>EQ 0.5MG BASE/ML</u>	<u>A076313 002</u>	Mar 28, 2005

SANDOSTATIN

<u>AP</u>	+! NOVARTIS	<u>EQ 0.05MG BASE/ML</u>	<u>N019667 001</u>	Oct 21, 1988
<u>AP</u>	+!	<u>EQ 0.1MG BASE/ML</u>	<u>N019667 002</u>	Oct 21, 1988
<u>AP</u>	+!	<u>EQ 0.5MG BASE/ML</u>	<u>N019667 003</u>	Oct 21, 1988
	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

ODEVIXIBAT

CAPSULE; ORAL

BYLVAY

+	ALBIREO	0.4MG	N215498 002	Jul 20, 2021
+	!	1.2MG	N215498 004	Jul 20, 2021

CAPSULE, PELLETS; ORAL

BYLVAY

+	ALBIREO	0.2MG	N215498 001	Jul 20, 2021
+	!	0.6MG	N215498 003	Jul 20, 2021

OFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

OCUFLOX

<u>AT</u>	+! ALLERGAN	<u>0.3%</u>	<u>N019921 001</u>	Jul 30, 1993
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OFLOXACIN

<u>AT</u>	AKORN	<u>0.3%</u>	<u>A076407 001</u>	Apr 15, 2008
<u>AT</u>		<u>0.3%</u>	<u>A076615 001</u>	May 14, 2004
<u>AT</u>	ALTAIRE PHARMS INC	<u>0.3%</u>	<u>A202692 001</u>	Apr 29, 2013
<u>AT</u>	APOTEX INC	<u>0.3%</u>	<u>A076513 001</u>	May 14, 2004
<u>AT</u>	BAUSCH AND LOMB	<u>0.3%</u>	<u>A076622 001</u>	May 14, 2004
<u>AT</u>	FDC LTD	<u>0.3%</u>	<u>A078559 001</u>	Feb 25, 2009

SOLUTION/DROPS; OTIC

OFLOXACIN

<u>AT</u>	AKORN	<u>0.3%</u>	<u>A076616 001</u>	Mar 17, 2008
<u>AT</u>	AMNEAL	<u>0.3%</u>	<u>A211525 001</u>	Aug 30, 2019
<u>AT</u>	APOTEX INC	<u>0.3%</u>	<u>A076527 001</u>	Sep 28, 2007
<u>AT</u>	! BAUSCH AND LOMB	<u>0.3%</u>	<u>A076128 001</u>	Mar 17, 2008

TABLET; ORAL

OFLOXACIN

<u>AB</u>	CADILA PHARMS LTD	<u>200MG</u>	<u>A091656 001</u>	Sep 18, 2014
<u>AB</u>		<u>300MG</u>	<u>A091656 002</u>	Sep 18, 2014
<u>AB</u>		<u>400MG</u>	<u>A091656 003</u>	Sep 18, 2014
<u>AB</u>	DR REDDYS LABS LTD	<u>200MG</u>	<u>A077098 001</u>	Feb 10, 2006
<u>AB</u>		<u>300MG</u>	<u>A077098 002</u>	Feb 10, 2006
<u>AB</u>		<u>400MG</u>	<u>A077098 003</u>	Feb 10, 2006
<u>AB</u>	LARKEN LABS	<u>400MG</u>	<u>A076093 003</u>	Sep 02, 2003
<u>AB</u>	TEVA	<u>200MG</u>	<u>A076182 001</u>	Sep 02, 2003
<u>AB</u>		<u>300MG</u>	<u>A076182 002</u>	Sep 02, 2003
<u>AB</u>	!	<u>400MG</u>	<u>A076182 003</u>	Sep 02, 2003

OLANZAPINE

INJECTABLE; INTRAMUSCULAR

OLANZAPINE

<u>AP</u>	AM REGENT	<u>10MG/VIAL</u>	<u>A201741 001</u>	Mar 20, 2012
<u>AP</u>	EUGIA PHARMA	<u>10MG/VIAL</u>	<u>A210968 001</u>	Oct 22, 2020
<u>AP</u>	SANDOZ INC	<u>10MG/VIAL</u>	<u>A201588 001</u>	Oct 24, 2011

ZYPREXA

<u>AP</u>	+! LILLY	<u>10MG/VIAL</u>	<u>N021253 001</u>	Mar 29, 2004
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TABLET; ORAL

OLANZAPINE

<u>AB</u>	ALKEM LABS LTD	<u>2.5MG</u>	<u>A202295 001</u>	Oct 20, 2015
<u>AB</u>		<u>5MG</u>	<u>A202295 002</u>	Oct 20, 2015
<u>AB</u>		<u>7.5MG</u>	<u>A202295 003</u>	Oct 20, 2015
<u>AB</u>		<u>10MG</u>	<u>A202295 004</u>	Oct 20, 2015
<u>AB</u>		<u>15MG</u>	<u>A202295 005</u>	Oct 20, 2015

PRESCRIPTION DRUG PRODUCT LIST

OLANZAPINE

TABLET; ORAL

OLANZAPINE

<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>	INDOCO	<u>2.5MG</u>	<u>A206155</u>	<u>001</u>	Jul 31, 2020
<u>AB</u>		<u>5MG</u>	<u>A206155</u>	<u>002</u>	Jul 31, 2020
<u>AB</u>		<u>7.5MG</u>	<u>A206155</u>	<u>003</u>	Jul 31, 2020
<u>AB</u>		<u>10MG</u>	<u>A206155</u>	<u>004</u>	Jul 31, 2020
<u>AB</u>		<u>15MG</u>	<u>A206155</u>	<u>005</u>	Jul 31, 2020
<u>AB</u>		<u>20MG</u>	<u>A206155</u>	<u>006</u>	Jul 31, 2020
<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>	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
<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>	ORBION PHARMS	<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	<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>	ZYDUS PHARMS	<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
<u>ZYPREXA</u>					
<u>AB</u>	+	<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

PRESCRIPTION DRUG PRODUCT LIST

OLANZAPINE

TABLET, ORALLY DISINTEGRATING;ORAL

OLANZAPINE

<u>AB</u>	APOTEX INC	<u>5MG</u>	<u>A091265 001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A091265 002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A091265 003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A091265 004</u>	Oct 24, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A203708 001</u>	May 15, 2014
<u>AB</u>		<u>10MG</u>	<u>A203708 002</u>	May 15, 2014
<u>AB</u>		<u>15MG</u>	<u>A203708 003</u>	May 15, 2014
<u>AB</u>		<u>20MG</u>	<u>A203708 004</u>	May 15, 2014
<u>AB</u>	BARR LABS INC	<u>5MG</u>	<u>A077243 001</u>	Jan 30, 2012
<u>AB</u>		<u>10MG</u>	<u>A077243 002</u>	Jan 30, 2012
<u>AB</u>		<u>15MG</u>	<u>A077243 003</u>	Jan 30, 2012
<u>AB</u>		<u>20MG</u>	<u>A077243 004</u>	Jan 30, 2012
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A076534 001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A076534 002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A076534 003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A076534 004</u>	Oct 24, 2011
<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	<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>	ORBION PHARMS	<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>	STRIDES PHARMA	<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	<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 RELPREVV

+	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

PRESCRIPTION DRUG PRODUCT LIST

OLANZAPINE; SAMIDORPHAN L-MALATE

TABLET; ORAL

LYBALVI

+	ALKERMES INC	5MG;EQ 10MG BASE	N213378 001	May 28, 2021
+		10MG;EQ 10MG BASE	N213378 002	May 28, 2021
+		15MG;EQ 10MG BASE	N213378 003	May 28, 2021
+		20MG;EQ 10MG BASE	N213378 004	May 28, 2021

OLAPARIB

TABLET; ORAL

LYNPARZA

+	ASTRAZENECA	100MG	N208558 001	Aug 17, 2017
+		150MG	N208558 002	Aug 17, 2017

OLICERIDINE

SOLUTION; INTRAVENOUS

OLINVIK

+	TREVENA	1MG/ML (1MG/ML)	N210730 001	Oct 30, 2020
+		2MG/2ML (1MG/ML)	N210730 002	Oct 30, 2020
+		30MG/30ML (1MG/ML)	N210730 003	Oct 30, 2020

OLIVE OIL; SOYBEAN OIL

EMULSION; INTRAVENOUS

CLINOLIPID 20%

+	BAXTER HLTHCARE CORP	16% (160GM/1000ML); 4% (40GM/1000ML)	N204508 001	Oct 03, 2013
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OLMESARTAN MEDOXOMIL

TABLET; ORAL

BENICAR

AB	+	DAIICHI SANKYO	5MG	N021286 001	Apr 25, 2002
AB	+		20MG	N021286 003	Apr 25, 2002
AB	+		40MG	N021286 004	Apr 25, 2002

OLMESARTAN MEDOXOMIL

AB		ACCORD HLTHCARE	5MG	A207662 001	Apr 24, 2017
AB			20MG	A207662 002	Apr 24, 2017
AB			40MG	A207662 003	Apr 24, 2017
AB		ALEMBIC PHARMS LTD	5MG	A203012 001	Apr 24, 2017
AB			20MG	A203012 002	Apr 24, 2017
AB			40MG	A203012 003	Apr 24, 2017
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		MACLEODS PHARMS LTD	5MG	A204814 001	Apr 24, 2017
AB			20MG	A204814 002	Apr 24, 2017
AB			40MG	A204814 003	Apr 24, 2017
AB		MICRO LABS	5MG	A206372 001	Sep 17, 2019
AB			20MG	A206372 002	Sep 17, 2019
AB			40MG	A206372 003	Sep 17, 2019
AB		QILU	5MG	A210552 001	Jan 10, 2019
AB			20MG	A210552 002	Jan 10, 2019
AB			40MG	A210552 003	Jan 10, 2019
AB		SANDOZ INC	5MG	A090237 001	Apr 13, 2020
AB			20MG	A090237 002	Apr 13, 2020
AB			40MG	A090237 003	Apr 13, 2020
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		SUNSHINE	5MG	A211049 001	Feb 22, 2019
AB			20MG	A211049 002	Feb 22, 2019
AB			40MG	A211049 003	Feb 22, 2019
AB		TORRENT	5MG	A202375 001	Apr 24, 2017
AB			20MG	A202375 002	Apr 24, 2017
AB			40MG	A202375 003	Apr 24, 2017
AB		UMEDICA LABS PVT LTD	5MG	A207135 001	Jul 18, 2019
AB			20MG	A207135 002	Jul 18, 2019
AB			40MG	A207135 003	Jul 18, 2019

PRESCRIPTION DRUG PRODUCT LIST

OLMESARTAN MEDOXOMIL

TABLET;ORAL

OLMESARTAN MEDOXOMIL

AB	ZYDUS PHARMS	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
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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
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OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

OLOPATADINE HYDROCHLORIDE

SOMERSET THERAPS

LLC

SPRAY, METERED;NASAL

OLOPATADINE HYDROCHLORIDE

AB	AKORN	0.665MG/SPRAY	A213757 001	Aug 19, 2020
AB	AMNEAL	0.665MG/SPRAY	A210901 001	Jan 28, 2020
AB	APOTEX INC	0.665MG/SPRAY	A091572 001	Oct 08, 2014
AB	PADAGIS ISRAEL	0.665MG/SPRAY	A202853 001	Jan 31, 2017

PATANASE

AB	+	NOVARTIS	0.665MG/SPRAY	N021861 001	Apr 15, 2008
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OLSALAZINE SODIUM

CAPSULE;ORAL

DIPENTUM

+	MYLAN SPCLT VIATRIS	250MG	N019715 001	Jul 31, 1990
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OMACETAXINE MEPESUCCINATE

POWDER;SUBCUTANEOUS

SYNRIBO

+	TEVA PHARMS INTL	3.5MG/VIAL	N203585 001	Oct 26, 2012
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OMADACYCLINE TOSYLATE

POWDER;INTRAVENOUS

NUZYRA

+	PARATEK PHARMS INC	EQ 100MG BASE/VIAL	N209817 001	Oct 02, 2018
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TABLET;ORAL

NUZYRA

+	PARATEK PHARMS INC	EQ 150MG BASE	N209816 001	Oct 02, 2018
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OMEGA-3-ACID ETHYL ESTERS

CAPSULE;ORAL

LOVAZA

AB	+	WOODWARD	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	N021654 001	Nov 10, 2004
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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	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A090973 001	Sep 30, 2014
AB	ASCENT PHARMS INC	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A207420 001	Feb 25, 2019
AB	BIONPHARMA INC	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A206455 001	Aug 07, 2019
AB	CSPC-NBP PHARM	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A211979 001	May 12, 2020
AB	GLW	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A212504 001	Aug 19, 2020
AB	MANKIND PHARMA	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A215458 001	Nov 15, 2021
AB	PURACAP PHARM LLC	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A210093 001	Jun 15, 2020
AB	SOFGEN PHARMS	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A211355 001	Jul 10, 2019
AB	STRIDES PHARMA	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A203893 001	Sep 19, 2017
AB	SUN PHARM	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A210834 001	Jan 09, 2020

PRESCRIPTION DRUG PRODUCT LIST

OMEGA-3-ACID ETHYL ESTERS

CAPSULE;ORAL

OMEGA-3-ACID ETHYL ESTERS

AB	TEVA PHARMS USA	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A091028 001	Apr 07, 2014
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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	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
AB		40MG	A078490 001	Apr 17, 2009
AB	GLENMARK GENERICS	10MG	A091672 001	Oct 31, 2014
AB		20MG	A091672 002	Oct 31, 2014
AB		40MG	A091672 003	Oct 31, 2014
AB	HETERO LABS LTD III	10MG	A204012 001	Sep 26, 2019
AB		20MG	A204012 002	Sep 26, 2019
AB	IMPAX LABS	10MG	A075785 001	Oct 22, 2007
AB		20MG	A075785 002	Oct 22, 2007
AB		40MG	A075785 003	Jan 21, 2009
AB	LANNETT CO INC	10MG	A075410 001	Nov 01, 2002
AB		20MG	A075410 002	Nov 01, 2002
AB		40MG	A075410 003	Jan 23, 2009
AB	SANDOZ	10MG	A075757 001	Jan 28, 2003
AB	!	20MG	A075757 002	Jan 28, 2003
AB	!	40MG	A076515 001	Jan 21, 2009
AB	TEVA PHARMS USA	20MG	A204661 001	Jun 13, 2017
AB	XIROMED	10MG	A212977 001	Dec 10, 2020
AB		20MG	A212977 002	Dec 10, 2020
AB	ZYDUS PHARMS USA INC	10MG	A091352 001	Nov 19, 2012
AB		20MG	A091352 002	Nov 19, 2012
AB		40MG	A091352 003	Nov 19, 2012

OMEPRAZOLE MAGNESIUM

FOR SUSPENSION, DELAYED RELEASE;ORAL

PRILOSEC

+ COVIS

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

AB	AJANTA PHARMA LTD	20MG;1.1GM	A204228 001	Jul 15, 2016
AB		40MG;1.1GM	A204228 002	Jul 15, 2016
AB	AUROLIFE PHARMA LLC	20MG;1.1GM	A204922 001	Aug 19, 2016
AB		40MG;1.1GM	A204922 002	Aug 19, 2016
AB	DR REDDYS LABS LTD	20MG;1.1GM	A204068 001	Jul 15, 2016
AB		40MG;1.1GM	A204068 002	Jul 15, 2016
AB	GUARDIAN DRUG	20MG;1.1GM	A212587 001	Apr 30, 2020
AB		40MG;1.1GM	A212587 002	Apr 30, 2020
AB	SCIEGEN PHARMS INC	20MG;1.1GM	A207476 001	Dec 06, 2016
AB		40MG;1.1GM	A207476 002	Dec 06, 2016
AB	ZYDUS PHARMS	20MG;1.1GM	A203290 001	May 25, 2018
AB		40MG;1.1GM	A203290 002	May 25, 2018

ZEGERID

AB	+ SALIX	20MG;1.1GM	N021849 001	Feb 27, 2006
AB	+!	40MG;1.1GM	N021849 002	Feb 27, 2006

PRESCRIPTION DRUG PRODUCT LIST

OMEPRAZOLE; SODIUM BICARBONATE

FOR SUSPENSION;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

<u>AB</u>	AJANTA PHARMA LTD	<u>20MG/PACKET;1.68GM/PACKET</u>	<u>A205545 001</u>	Jul 27, 2016
<u>AB</u>		<u>40MG/PACKET;1.68GM/PACKET</u>	<u>A205545 002</u>	Jul 27, 2016
<u>AB</u>	STRIDES PHARMA	<u>20MG/PACKET;1.68GM/PACKET</u>	<u>A079182 001</u>	Apr 19, 2013
<u>AB</u>		<u>40MG/PACKET;1.68GM/PACKET</u>	<u>A079182 002</u>	Apr 19, 2013
<u>ZEGERID</u>				
<u>AB</u>	+ SALIX	<u>20MG/PACKET;1.68GM/PACKET</u>	<u>N021636 001</u>	Jun 15, 2004
<u>AB</u>	+!	<u>40MG/PACKET;1.68GM/PACKET</u>	<u>N021636 002</u>	Dec 21, 2004

ONDANSETRON

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

<u>AB</u>	AUROBINDO PHARMA	<u>4MG</u>	<u>A090469 001</u>	Apr 12, 2010
<u>AB</u>	!	<u>8MG</u>	<u>A090469 002</u>	Apr 12, 2010
<u>AB</u>	GLENMARK GENERICS	<u>4MG</u>	<u>A078152 001</u>	Jun 27, 2007
<u>AB</u>		<u>8MG</u>	<u>A078152 002</u>	Jun 27, 2007
<u>AB</u>	MYLAN	<u>4MG</u>	<u>A078139 001</u>	Jun 25, 2007
<u>AB</u>		<u>8MG</u>	<u>A078139 002</u>	Jun 25, 2007
<u>AB</u>	SANDOZ	<u>4MG</u>	<u>A078050 001</u>	Aug 13, 2007
<u>AB</u>		<u>8MG</u>	<u>A078050 002</u>	Aug 13, 2007
<u>AB</u>	SUN PHARM INDS	<u>4MG</u>	<u>A077557 001</u>	Aug 02, 2007
<u>AB</u>		<u>8MG</u>	<u>A077557 002</u>	Aug 02, 2007
<u>AB</u>	SUN PHARM INDS LTD	<u>4MG</u>	<u>A078602 001</u>	Feb 24, 2011
<u>AB</u>		<u>8MG</u>	<u>A078602 002</u>	Feb 24, 2011

ONDANSETRON HYDROCHLORIDE

INJECTABLE;INJECTION

ONDANSETRON HYDROCHLORIDE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 2MG BASE/ML</u>	<u>A206846 001</u>	Jul 13, 2015
<u>AP</u>	EUGIA PHARMA	<u>EQ 2MG BASE/ML</u>	<u>A202599 001</u>	Dec 21, 2012
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A076974 001</u>	Dec 26, 2006
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 2MG BASE/ML</u>	<u>A079224 001</u>	Sep 25, 2009
<u>AP</u>	!	<u>EQ 2MG BASE/ML</u>	<u>A090648 001</u>	Jun 15, 2012
<u>AP</u>	HIKMA	<u>EQ 2MG BASE/ML</u>	<u>A076967 001</u>	Dec 26, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A077365 001</u>	Dec 26, 2006
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 2MG BASE/ML</u>	<u>A076781 001</u>	Dec 26, 2006
<u>AP</u>	HOSPIRA	<u>EQ 2MG BASE/ML</u>	<u>A077473 001</u>	Dec 26, 2006
<u>AP</u>	!	<u>EQ 2MG BASE/ML</u>	<u>A203711 001</u>	Sep 08, 2014
<u>AP</u>	SANDOZ INC	<u>EQ 2MG BASE/ML</u>	<u>A077430 001</u>	Jun 27, 2007
<u>AP</u>	TEVA	<u>EQ 2MG BASE/ML</u>	<u>A076876 001</u>	Nov 22, 2006
<u>AP</u>	WOCKHARDT	<u>EQ 2MG BASE/ML</u>	<u>A077577 001</u>	Dec 26, 2006

ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 2MG BASE/ML</u>	<u>A206845 001</u>	Mar 10, 2016
<u>AP</u>	!	<u>EQ 2MG BASE/ML</u>	<u>A078287 001</u>	Feb 22, 2013
<u>AP</u>	EUGIA PHARMA	<u>EQ 2MG BASE/ML</u>	<u>A202600 001</u>	Dec 21, 2012
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A076972 001</u>	Dec 26, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A202253 001</u>	Jul 19, 2013
<u>AP</u>	HIKMA	<u>EQ 2MG BASE/ML</u>	<u>A077011 001</u>	Dec 26, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A077541 001</u>	Dec 26, 2006
<u>AP</u>	HOSPIRA	<u>EQ 2MG BASE/ML</u>	<u>A077548 001</u>	Dec 26, 2006
<u>AP</u>	SANDOZ INC	<u>EQ 2MG BASE/ML</u>	<u>A077551 001</u>	Jun 27, 2007
<u>AP</u>	WOCKHARDT	<u>EQ 2MG BASE/ML</u>	<u>A077716 001</u>	Dec 26, 2006

SOLUTION;ORAL

ONDANSETRON HYDROCHLORIDE

<u>AA</u>	AMNEAL PHARMS	<u>EQ 4MG BASE/5ML</u>	<u>A091483 001</u>	Jan 31, 2011
<u>AA</u>	AUROBINDO PHARMA	<u>EQ 4MG BASE/5ML</u>	<u>A078776 001</u>	Nov 28, 2007
<u>AA</u>	HIKMA	<u>EQ 4MG BASE/5ML</u>	<u>A076960 001</u>	Dec 26, 2006
<u>AA</u>	LANNETT CO INC	<u>EQ 4MG BASE/5ML</u>	<u>A091342 001</u>	Jan 27, 2011
<u>AA</u>	PHARM ASSOC	<u>EQ 4MG BASE/5ML</u>	<u>A078127 001</u>	Jun 25, 2007
<u>AA</u>	TARO	<u>EQ 4MG BASE/5ML</u>	<u>A077009 001</u>	Nov 30, 2007
<u>ZOFRAN</u>				
<u>AA</u>	+! NOVARTIS	<u>EQ 4MG BASE/5ML</u>	<u>N020605 001</u>	Jan 24, 1997

TABLET;ORAL

ONDANSETRON HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 4MG BASE</u>	<u>A077306 001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077306 002</u>	Jun 25, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 4MG BASE</u>	<u>A078539 001</u>	Jul 31, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A078539 002</u>	Jul 31, 2007
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A078539 003</u>	Jul 31, 2007
<u>AB</u>	CASI PHARMS INC	<u>EQ 4MG BASE</u>	<u>A077517 001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077517 002</u>	Jun 25, 2007

PRESCRIPTION DRUG PRODUCT LIST

ONDANSETRON HYDROCHLORIDE

TABLET; ORAL

ONDANSETRON HYDROCHLORIDE

<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A077517 003</u>	Jun 25, 2007
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE</u>	<u>A076183 003</u>	Dec 26, 2006
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A076183 002</u>	Dec 26, 2006
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A076183 001</u>	Dec 26, 2006
<u>AB</u>	GLENMARK GENERICS	<u>EQ 4MG BASE</u>	<u>A077535 001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077535 002</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A077535 003</u>	Jun 25, 2007
<u>AB</u>	IPCA LABS LTD	<u>EQ 4MG BASE</u>	<u>A203761 001</u>	Jan 23, 2014
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A203761 002</u>	Jan 23, 2014
<u>AB</u>	NATCO PHARMA LTD	<u>EQ 4MG BASE</u>	<u>A077851 001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077851 002</u>	Jun 25, 2007
<u>AB</u>	SUN PHARM INDS (IN)	<u>EQ 4MG BASE</u>	<u>A077050 001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077050 002</u>	Jun 25, 2007
<u>AB</u>	TEVA	<u>EQ 4MG BASE</u>	<u>A076252 001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A076252 002</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A076252 003</u>	Jun 25, 2007

ZOFRAN

<u>AB</u>	+	NOVARTIS	<u>EQ 4MG BASE</u>	<u>N020103 001</u>	Dec 31, 1992
<u>AB</u>	+		<u>EQ 8MG BASE</u>	<u>N020103 002</u>	Dec 31, 1992
<u>AB</u>	+		<u>EQ 24MG BASE</u>	<u>N020103 003</u>	Aug 27, 1999
		ONDANSETRON HYDROCHLORIDE			
		DR REDDYS LABS LTD	EQ 16MG BASE	A076183 004	Dec 26, 2006

OPICAPONE

CAPSULE; ORAL

ONGENTYS

+	NEUROCRINE	25MG	N212489 001	Apr 24, 2020
+		50MG	N212489 002	Apr 24, 2020

ORITAVANCIN DIPHOSPHATE

POWDER; INTRAVENOUS

KIMYRSA

+	MELINTA THERAP	EQ 1.2GM BASE/VIAL	N214155 001	Mar 12, 2021
	ORBACTIV			
+	MELINTA THERAP	EQ 400MG BASE/VIAL	N206334 001	Aug 06, 2014

ORLISTAT

CAPSULE; ORAL

XENICAL

+	CHEPLAPHARM	120MG	N020766 001	Apr 23, 1999
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ORPHENADRINE CITRATE

INJECTABLE; INJECTION

ORPHENADRINE CITRATE

<u>AP</u>	!	AKORN	<u>30MG/ML</u>	<u>A040484 001</u>	May 24, 2006
<u>AP</u>		HIKMA	<u>30MG/ML</u>	<u>A040463 001</u>	Mar 04, 2003
<u>AP</u>		SAGENT PHARMS	<u>30MG/ML</u>	<u>A090585 001</u>	Aug 30, 2011
<u>AP</u>		WATSON LABS	<u>30MG/ML</u>	<u>A084779 001</u>	Mar 15, 1982

TABLET, EXTENDED RELEASE; ORAL

ORPHENADRINE CITRATE

<u>AB</u>		ANDA REPOSITORY	<u>100MG</u>	<u>A040249 001</u>	Jan 29, 1999
<u>AB</u>		INVAGEN PHARMS	<u>100MG</u>	<u>A091158 001</u>	Jul 27, 2012
<u>AB</u>		LUPIN	<u>100MG</u>	<u>A040284 001</u>	Jun 19, 1998
<u>AB</u>	!	SANDOZ	<u>100MG</u>	<u>A040327 001</u>	Feb 15, 2000

OSELTAMIVIR PHOSPHATE

CAPSULE; ORAL

OSELTAMIVIR PHOSPHATE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>EQ 30MG BASE</u>	<u>A211823 001</u>	Jun 24, 2019
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A211823 002</u>	Jun 24, 2019
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A211823 003</u>	Jun 24, 2019
<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>		EPIC PHARMA LLC	<u>EQ 30MG BASE</u>	<u>A215208 001</u>	Oct 01, 2021
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A215208 002</u>	Oct 01, 2021
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A215208 003</u>	Oct 01, 2021
<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	<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

PRESCRIPTION DRUG PRODUCT LIST

OSELTAMIVIR PHOSPHATE

CAPSULE;ORAL

OSELTAMIVIR PHOSPHATE

<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>	MSN	<u>EQ 30MG BASE</u>	<u>A212544 001</u>	May 20, 2020
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A212544 002</u>	May 20, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A212544 003</u>	May 20, 2020
<u>AB</u>	NATCO	<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>	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
<u>AB</u>	ZYDUS 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

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>	AJANTA PHARMA LTD	<u>EQ 6MG BASE/ML</u>	<u>A212784 001</u>	May 27, 2020
<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>	APTAPHARMA INC	<u>EQ 6MG BASE/ML</u>	<u>A212858 001</u>	Aug 30, 2021
<u>AB</u>	LUPIN	<u>EQ 6MG BASE/ML</u>	<u>A208347 001</u>	Feb 20, 2018
<u>AB</u>	SUNSHINE	<u>EQ 6MG BASE/ML</u>	<u>A213594 001</u>	Jan 05, 2022
<u>AB</u>	TEVA PHARMS USA	<u>EQ 6MG BASE/ML</u>	<u>A211125 001</u>	Feb 27, 2019
<u>AB</u>	ZYDUS 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
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OSILODROSTAT PHOSPHATE

TABLET;ORAL

ISTURISA

+	RECORDATI RARE	EQ 1MG BASE	N212801 001	Mar 06, 2020
+		EQ 5MG BASE	N212801 002	Mar 06, 2020
+		EQ 10MG BASE	N212801 003	Mar 06, 2020

OSIMERTINIB MESYLATE

TABLET;ORAL

TAGRISSO

+	ASTRAZENECA	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
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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>	FRESENIUS KABI USA	<u>EQ 1GM BASE/VIAL</u>	<u>A206198 001</u>	Jul 20, 2020
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A206198 002</u>	Jul 20, 2020
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A206199 001</u>	Jul 27, 2020
<u>AP</u>	PIRAMAL CRITICAL	<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

PRESCRIPTION DRUG PRODUCT LIST

OXACILLIN SODIUM

INJECTABLE; INJECTION

BACTOCILL IN PLASTIC CONTAINER

+	!	BAXTER HLTHCARE	EQ 20MG BASE/ML	N050640	001	Oct 26, 1989
+	!		EQ 40MG BASE/ML	N050640	002	Oct 26, 1989

OXALIPLATIN

INJECTABLE; INTRAVENOUS

ELOXATIN

AP	+	!	SANOFI AVENTIS US	<u>50MG/10ML (5MG/ML)</u>	<u>N021759</u>	<u>001</u>	Jan 31, 2005
AP	+	!		<u>100MG/20ML (5MG/ML)</u>	<u>N021759</u>	<u>002</u>	Jan 31, 2005

OXALIPLATIN

AP			ACCORD HLTHCARE	<u>50MG/10ML (5MG/ML)</u>	<u>A207474</u>	<u>001</u>	Mar 21, 2017
AP				<u>100MG/20ML (5MG/ML)</u>	<u>A207474</u>	<u>002</u>	Mar 21, 2017
AP			ACTAVIS LLC	<u>50MG/10ML (5MG/ML)</u>	<u>A204880</u>	<u>001</u>	Mar 05, 2018
AP				<u>100MG/20ML (5MG/ML)</u>	<u>A204880</u>	<u>002</u>	Mar 05, 2018
AP	!		ACTAVIS TOTOWA	<u>50MG/VIAL</u>	<u>A078803</u>	<u>001</u>	Aug 08, 2012
AP	!			<u>100MG/VIAL</u>	<u>A078803</u>	<u>002</u>	Aug 08, 2012
AP			EUGIA PHARMA	<u>50MG/10ML (5MG/ML)</u>	<u>A205529</u>	<u>001</u>	Sep 06, 2017
AP				<u>100MG/20ML (5MG/ML)</u>	<u>A205529</u>	<u>002</u>	Sep 06, 2017
AP			FRESENIUS KABI USA	<u>50MG/10ML (5MG/ML)</u>	<u>A078811</u>	<u>001</u>	Jun 10, 2010
AP				<u>100MG/20ML (5MG/ML)</u>	<u>A078811</u>	<u>002</u>	Jun 10, 2010
AP				<u>50MG/VIAL</u>	<u>A078819</u>	<u>001</u>	Jun 02, 2010
AP				<u>50MG/10ML (5MG/ML)</u>	<u>A090030</u>	<u>001</u>	Jan 31, 2017
AP				<u>100MG/VIAL</u>	<u>A078819</u>	<u>002</u>	Jun 02, 2010
AP				<u>100MG/20ML (5MG/ML)</u>	<u>A090030</u>	<u>002</u>	Jan 31, 2017
AP			GLAND	<u>50MG/10ML (5MG/ML)</u>	<u>A207325</u>	<u>001</u>	Feb 10, 2017
AP				<u>100MG/20ML (5MG/ML)</u>	<u>A207325</u>	<u>002</u>	Feb 10, 2017
AP			GLAND PHARMA LTD	<u>50MG/VIAL</u>	<u>A207385</u>	<u>001</u>	May 23, 2017
AP				<u>100MG/VIAL</u>	<u>A207385</u>	<u>002</u>	May 23, 2017
AP			HOSPIRA WORLDWIDE	<u>50MG/10ML (5MG/ML)</u>	<u>A078813</u>	<u>001</u>	Aug 07, 2009
AP				<u>100MG/20ML (5MG/ML)</u>	<u>A078813</u>	<u>002</u>	Aug 07, 2009
AP			JIANGSU PHARMS	<u>50MG/10ML (5MG/ML)</u>	<u>A203869</u>	<u>001</u>	Jun 18, 2014
AP				<u>100MG/20ML (5MG/ML)</u>	<u>A203869</u>	<u>002</u>	Jun 18, 2014
AP			MYLAN LABS LTD	<u>50MG/10ML (5MG/ML)</u>	<u>A091358</u>	<u>001</u>	Aug 07, 2012
AP				<u>100MG/20ML (5MG/ML)</u>	<u>A091358</u>	<u>002</u>	Aug 07, 2012
AP			NOVAST LABS	<u>50MG/10ML (5MG/ML)</u>	<u>A207562</u>	<u>001</u>	Oct 16, 2018
AP				<u>100MG/20ML (5MG/ML)</u>	<u>A207562</u>	<u>002</u>	Oct 16, 2018
AP			QILU PHARM HAINAN	<u>50MG/10ML (5MG/ML)</u>	<u>A204368</u>	<u>001</u>	Jun 07, 2016
AP				<u>50MG/VIAL</u>	<u>A204616</u>	<u>001</u>	May 11, 2016
AP				<u>100MG/20ML (5MG/ML)</u>	<u>A204368</u>	<u>002</u>	Jun 07, 2016
AP				<u>100MG/VIAL</u>	<u>A204616</u>	<u>002</u>	May 11, 2016
AP			SANDOZ	<u>50MG/10ML (5MG/ML)</u>	<u>A078817</u>	<u>001</u>	Jan 24, 2011
AP				<u>100MG/20ML (5MG/ML)</u>	<u>A078817</u>	<u>002</u>	Jan 24, 2011
AP	+	!	TEVA PHARMS	<u>50MG/10ML (5MG/ML)</u>	<u>N022160</u>	<u>001</u>	Aug 07, 2009
AP	+	!		<u>100MG/20ML (5MG/ML)</u>	<u>N022160</u>	<u>002</u>	Aug 07, 2009
	!		QILU PHARM HAINAN	200MG/40ML (5MG/ML)	A204368	003	Jun 07, 2016

OXANDROLONE

TABLET; ORAL

OXANDROLONE

AB			PAR PHARM	<u>2.5MG</u>	<u>A077827</u>	<u>001</u>	Jun 22, 2007
AB	!			<u>10MG</u>	<u>A077827</u>	<u>002</u>	Jun 22, 2007
AB			UPSHER SMITH LABS	<u>2.5MG</u>	<u>A076761</u>	<u>001</u>	Dec 01, 2006
AB				<u>10MG</u>	<u>A078033</u>	<u>001</u>	Mar 22, 2007

OXAPROZIN

TABLET; ORAL

DAYPRO

AB	+	!	PFIZER	<u>600MG</u>	<u>N018841</u>	<u>004</u>	Oct 29, 1992
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OXAPROZIN

AB			AMNEAL PHARMS CO	<u>600MG</u>	<u>A208633</u>	<u>001</u>	May 04, 2017
AB			CHARTWELL	<u>600MG</u>	<u>A075987</u>	<u>001</u>	Sep 02, 2004
AB			DR REDDYS LABS LTD	<u>600MG</u>	<u>A075855</u>	<u>001</u>	Jan 31, 2001
AB			SANDOZ	<u>600MG</u>	<u>A075845</u>	<u>001</u>	Jan 31, 2001
AB			TEVA	<u>600MG</u>	<u>A075849</u>	<u>001</u>	Jul 03, 2002

OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

AB			ACTAVIS ELIZABETH	<u>10MG</u>	<u>A072253</u>	<u>002</u>	Apr 14, 1988
AB				<u>15MG</u>	<u>A072253</u>	<u>003</u>	Apr 14, 1988
AB	!			<u>30MG</u>	<u>A072253</u>	<u>001</u>	Apr 14, 1988
AB			FRONTIDA BIOPHARM	<u>10MG</u>	<u>A071026</u>	<u>002</u>	Aug 10, 1987

PRESCRIPTION DRUG PRODUCT LIST

OXAZEPAM

CAPSULE;ORAL

OXAZEPAM

<u>AB</u>		<u>15MG</u>	<u>A071026</u>	<u>003</u>	Aug 10, 1987
<u>AB</u>		<u>30MG</u>	<u>A071026</u>	<u>001</u>	Aug 10, 1987
<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>	AKORN	<u>300MG/5ML</u>	<u>A211420</u>	<u>001</u>	Jul 09, 2021
<u>AB</u>	AMNEAL PHARMS	<u>300MG/5ML</u>	<u>A202961</u>	<u>001</u>	Sep 17, 2012
<u>AB</u>	GLASSHOUSE PHARMS	<u>300MG/5ML</u>	<u>A212428</u>	<u>001</u>	Jun 21, 2021
<u>AB</u>	HIKMA	<u>300MG/5ML</u>	<u>A201193</u>	<u>001</u>	Oct 03, 2012
<u>AB</u>	SUN PHARM INDS LTD	<u>300MG/5ML</u>	<u>A078734</u>	<u>001</u>	Jun 26, 2009

TRILEPTAL

<u>AB</u>	+!	NOVARTIS	<u>300MG/5ML</u>	<u>N021285</u>	<u>001</u>	May 25, 2001
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TABLET;ORAL

OXCARBAZEPINE

<u>AB</u>	ANNORA PHARMA	<u>150MG</u>	<u>A215939</u>	<u>001</u>	Jan 11, 2022
<u>AB</u>		<u>300MG</u>	<u>A215939</u>	<u>002</u>	Jan 11, 2022
<u>AB</u>		<u>600MG</u>	<u>A215939</u>	<u>003</u>	Jan 11, 2022
<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>	RUBICON	<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>	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

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
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OXISTAT

<u>AB</u>	+!	FOUGERA PHARMS	<u>EQ 1% BASE</u>	<u>N019828</u>	<u>001</u>	Dec 30, 1988
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LOTION;TOPICAL

OXISTAT

+	ANI PHARMS	EQ 1% BASE	N020209	001	Sep 30, 1992
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OXYBUTYNIN

FILM, EXTENDED RELEASE;TRANSDERMAL

OXYTROL

+	ALLERGAN	3.9MG/24HR	N021351	002	Feb 26, 2003
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OXYBUTYNIN CHLORIDE

GEL;TRANSDERMAL

GELNIQUE

+	ALLERGAN	10% (100MG/PACKET)	N022204	001	Jan 27, 2009
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SYRUP;ORAL

OXYBUTYNIN CHLORIDE

<u>AA</u>	LANNETT CO INC	<u>5MG/5ML</u>	<u>A074520</u>	<u>001</u>	Mar 29, 1996	
<u>AA</u>	PHARM ASSOC	<u>5MG/5ML</u>	<u>A075137</u>	<u>001</u>	Dec 18, 1998	
<u>AA</u>	!	WOCKHARDT BIO AG	<u>5MG/5ML</u>	<u>A074868</u>	<u>001</u>	Feb 12, 1997

PRESCRIPTION DRUG PRODUCT LIST

OXYBUTYNYN CHLORIDE

TABLET;ORAL

OXYBUTYNYN CHLORIDE

<u>AB</u>	ABHAI LLC	<u>5MG</u>	<u>A209335</u>	<u>001</u>	Dec 22, 2017
<u>AB</u>	APPCO	<u>5MG</u>	<u>A209025</u>	<u>001</u>	Dec 21, 2017
<u>AB</u>	LEADING PHARMA LLC	<u>5MG</u>	<u>A212798</u>	<u>001</u>	Apr 06, 2020
<u>AB</u>	NOVAST LABS	<u>5MG</u>	<u>A210611</u>	<u>001</u>	Oct 30, 2019
<u>AB</u>	NOVITIUM PHARMA	<u>5MG</u>	<u>A209823</u>	<u>001</u>	Oct 23, 2017
<u>AB</u>	! STRIDES PHARMA	<u>5MG</u>	<u>A075079</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>	TEVA PHARMS USA	<u>5MG</u>	<u>A071655</u>	<u>001</u>	Nov 14, 1988
<u>AB</u>	TULEX PHARMS INC	<u>5MG</u>	<u>A210125</u>	<u>001</u>	Sep 06, 2018
<u>AB</u>	UPSHER SMITH LABS	<u>5MG</u>	<u>A074625</u>	<u>001</u>	Jul 31, 1996
BX	EMCURE PHARMS LTD	5MG	A211682	001	May 10, 2019
BX	EYWA	5MG	A211062	001	Feb 06, 2019

TABLET, EXTENDED RELEASE;ORAL

DITROPAN XL

<u>AB</u>	+ JANSSEN PHARMS	<u>5MG</u>	<u>N020897</u>	<u>001</u>	Dec 16, 1998
<u>AB</u>	+	<u>10MG</u>	<u>N020897</u>	<u>002</u>	Dec 16, 1998

OXYBUTYNYN CHLORIDE

<u>AB</u>	ACCORD HLTHCARE	<u>5MG</u>	<u>A207138</u>	<u>001</u>	Feb 29, 2016
<u>AB</u>		<u>10MG</u>	<u>A207138</u>	<u>002</u>	Feb 29, 2016
<u>AB</u>	!	<u>15MG</u>	<u>A207138</u>	<u>003</u>	Feb 29, 2016
<u>AB</u>	AJANTA PHARMA LTD	<u>5MG</u>	<u>A211655</u>	<u>001</u>	Feb 28, 2019
<u>AB</u>		<u>10MG</u>	<u>A211655</u>	<u>002</u>	Feb 28, 2019
<u>AB</u>		<u>15MG</u>	<u>A211655</u>	<u>003</u>	Feb 28, 2019
<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A204010</u>	<u>001</u>	Nov 23, 2015
<u>AB</u>		<u>10MG</u>	<u>A204010</u>	<u>002</u>	Nov 23, 2015
<u>AB</u>		<u>15MG</u>	<u>A204010</u>	<u>003</u>	Nov 23, 2015
<u>AB</u>	BIONPHARMA INC	<u>5MG</u>	<u>A210717</u>	<u>001</u>	Dec 17, 2019
<u>AB</u>		<u>10MG</u>	<u>A210717</u>	<u>002</u>	Dec 17, 2019
<u>AB</u>		<u>15MG</u>	<u>A210717</u>	<u>003</u>	Dec 17, 2019
<u>AB</u>	OSMOTICA PHARM US	<u>5MG</u>	<u>A078503</u>	<u>001</u>	Feb 04, 2009
<u>AB</u>		<u>10MG</u>	<u>A078503</u>	<u>002</u>	Feb 04, 2009
<u>AB</u>		<u>15MG</u>	<u>A078503</u>	<u>003</u>	Feb 04, 2009
<u>AB</u>	RUBICON	<u>5MG</u>	<u>A214415</u>	<u>001</u>	Oct 27, 2020
<u>AB</u>		<u>10MG</u>	<u>A214415</u>	<u>002</u>	Oct 27, 2020
<u>AB</u>		<u>15MG</u>	<u>A214415</u>	<u>003</u>	Oct 27, 2020
<u>AB</u>	UNIQUE PHARM LABS	<u>5MG</u>	<u>A206121</u>	<u>001</u>	May 27, 2016
<u>AB</u>		<u>10MG</u>	<u>A206121</u>	<u>002</u>	May 27, 2016
<u>AB</u>		<u>15MG</u>	<u>A206121</u>	<u>003</u>	May 27, 2016
<u>AB</u>	ZYDUS PHARMS	<u>5MG</u>	<u>A202332</u>	<u>001</u>	Jun 26, 2017
<u>AB</u>		<u>10MG</u>	<u>A202332</u>	<u>002</u>	Jun 26, 2017
<u>AB</u>		<u>15MG</u>	<u>A202332</u>	<u>003</u>	Jun 26, 2017

OXYCODONE

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

<u>AB</u>	ANI PHARMS	<u>5MG</u>	<u>A205177</u>	<u>001</u>	Mar 31, 2016
<u>AB</u>	AVANTHI INC	<u>5MG</u>	<u>A202773</u>	<u>001</u>	Aug 17, 2015
<u>AB</u>	+! GENUS LIFESCIENCES	<u>5MG</u>	<u>N200534</u>	<u>001</u>	Oct 20, 2010
<u>AB</u>	MAYNE PHARMA INC	<u>5MG</u>	<u>A203107</u>	<u>001</u>	Jul 26, 2012
<u>AB</u>	NOVEL LABS INC	<u>5MG</u>	<u>A204752</u>	<u>001</u>	Aug 24, 2015

SOLUTION;ORAL

OXYCODONE HYDROCHLORIDE

<u>AA</u>	ABHAI LLC	<u>5MG/5ML</u>	<u>A208593</u>	<u>001</u>	Jul 21, 2017
<u>AA</u>		<u>100MG/5ML</u>	<u>A208593</u>	<u>002</u>	Jul 21, 2017
<u>AA</u>	AKORN	<u>5MG/5ML</u>	<u>A208817</u>	<u>001</u>	Aug 10, 2017
<u>AA</u>		<u>100MG/5ML</u>	<u>A208795</u>	<u>001</u>	Aug 07, 2017
<u>AA</u>	ALKEM LABS LTD	<u>5MG/5ML</u>	<u>A211748</u>	<u>001</u>	Feb 07, 2019
<u>AA</u>		<u>100MG/5ML</u>	<u>A211749</u>	<u>001</u>	Feb 04, 2019
<u>AA</u>	ANI PHARMS	<u>5MG/5ML</u>	<u>A204979</u>	<u>001</u>	Jun 01, 2015
<u>AA</u>	ASCENT PHARMS INC	<u>5MG/5ML</u>	<u>A209021</u>	<u>001</u>	Nov 09, 2017
<u>AA</u>		<u>100MG/5ML</u>	<u>A209021</u>	<u>002</u>	Nov 09, 2017
<u>AA</u>	EYWA	<u>5MG/5ML</u>	<u>A207511</u>	<u>001</u>	Nov 23, 2016

PRESCRIPTION DRUG PRODUCT LIST

OXYCODONE HYDROCHLORIDE

SOLUTION;ORAL

OXYCODONE HYDROCHLORIDE

<u>AA</u>		<u>100MG/5ML</u>	<u>A209897 001</u>	Sep 06, 2017
<u>AA</u>	+	<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>		<u>5MG/5ML</u>	<u>A204037 001</u>	Jul 15, 2013
<u>AA</u>		<u>100MG/5ML</u>	<u>A204092 001</u>	Jun 05, 2014
<u>AA</u>		<u>100MG/5ML</u>	<u>A204603 001</u>	Apr 29, 2015
<u>AA</u>		<u>5MG/5ML</u>	<u>A206914 001</u>	Feb 01, 2019
<u>AA</u>		<u>100MG/5ML</u>	<u>A206822 001</u>	Aug 15, 2017
<u>AA</u>		<u>5MG/5ML</u>	<u>A213761 001</u>	Jun 02, 2021
<u>AA</u>		<u>100MG/5ML</u>	<u>A213761 002</u>	Jun 02, 2021
<u>AA</u>		<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>	+	<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>		<u>5MG/5ML</u>	<u>A206456 001</u>	Jun 16, 2015

TABLET;ORAL

OXYCODONE HYDROCHLORIDE

<u>AB</u>		<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>		<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>		<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>		<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>		<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>		<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>		<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
<u>AB</u>		<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>		<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>		<u>5MG</u>	<u>A076758 003</u>	Mar 19, 2007
<u>AB</u>		<u>10MG</u>	<u>A076758 004</u>	Oct 12, 2021
<u>AB</u>		<u>15MG</u>	<u>A076758 001</u>	Jun 30, 2004
<u>AB</u>		<u>20MG</u>	<u>A076758 005</u>	Oct 12, 2021
<u>AB</u>		<u>30MG</u>	<u>A076758 002</u>	Jun 30, 2004
<u>AB</u>		<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

PRESCRIPTION DRUG PRODUCT LIST

OXYCODONE HYDROCHLORIDE

TABLET; ORAL

ROXICODONE

<u>AB</u>	+	SPECGX LLC	<u>5MG</u>	<u>N021011</u>	<u>003</u>	May 15, 2009
<u>AB</u>	+	!	<u>15MG</u>	<u>N021011</u>	<u>001</u>	Aug 31, 2000
<u>AB</u>	+		<u>30MG</u>	<u>N021011</u>	<u>002</u>	Aug 31, 2000

OXAYDO

+	ZYLA	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

+	EPI HLTH	1%	N208552	001	Jan 18, 2017
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SOLUTION/DROPS; OPHTHALMIC

UPNEEQ

+	RVL PHARMS	0.1%	N212520	001	Jul 08, 2020
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OXYMETAZOLINE HYDROCHLORIDE; TETRACAINE HYDROCHLORIDE

SPRAY, METERED; NASAL

KOVANAZE

+	ST RENATUS	0.1MG/SPRAY; 6MG/SPRAY	N208032	001	Jun 29, 2016
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OXYMORPHONE HYDROCHLORIDE

TABLET; ORAL

OXYMORPHONE HYDROCHLORIDE

<u>AB</u>		ASCENT PHARMS INC	<u>5MG</u>	<u>A210175</u>	<u>001</u>	Feb 02, 2018
<u>AB</u>			<u>10MG</u>	<u>A210175</u>	<u>002</u>	Feb 02, 2018
<u>AB</u>		AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A204459</u>	<u>001</u>	Apr 26, 2016
<u>AB</u>			<u>10MG</u>	<u>A204459</u>	<u>002</u>	Apr 26, 2016
<u>AB</u>		AVANTHI INC	<u>5MG</u>	<u>A203601</u>	<u>001</u>	Jan 30, 2013
<u>AB</u>	!		<u>10MG</u>	<u>A203601</u>	<u>002</u>	Jan 30, 2013
<u>AB</u>		EPIC PHARMA LLC	<u>5MG</u>	<u>A201187</u>	<u>001</u>	Dec 15, 2014
<u>AB</u>			<u>10MG</u>	<u>A201187</u>	<u>002</u>	Dec 15, 2014
<u>AB</u>		HIKMA	<u>5MG</u>	<u>A090964</u>	<u>001</u>	Sep 27, 2010
<u>AB</u>			<u>10MG</u>	<u>A090964</u>	<u>002</u>	Sep 27, 2010
<u>AB</u>		SPECGX LLC	<u>5MG</u>	<u>A202321</u>	<u>001</u>	Apr 25, 2013
<u>AB</u>			<u>10MG</u>	<u>A202321</u>	<u>002</u>	Apr 25, 2013
<u>AB</u>		TEVA	<u>5MG</u>	<u>A091443</u>	<u>002</u>	Feb 15, 2011
<u>AB</u>			<u>10MG</u>	<u>A091443</u>	<u>001</u>	Feb 15, 2011

TABLET, EXTENDED RELEASE; ORAL

OXYMORPHONE HYDROCHLORIDE

IMPAX LABS

		5MG	A079087	001	Jun 14, 2010
		7.5MG	A079087	002	Dec 21, 2010
		10MG	A079087	003	Jun 14, 2010
		15MG	A079087	004	Dec 21, 2010
		20MG	A079087	005	Jun 14, 2010
		30MG	A079087	006	Jul 22, 2010
	!	40MG	A079087	007	Jun 14, 2010

OXYTOCIN

INJECTABLE; INJECTION

OXYTOCIN

<u>AP</u>	+	FRESENIUS KABI USA	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018248</u>	<u>001</u>	
<u>AP</u>	+		<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018248</u>	<u>002</u>	
<u>AP</u>	+	HIKMA	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018243</u>	<u>001</u>	
<u>AP</u>	+		<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018243</u>	<u>002</u>	Jan 10, 2007
<u>AP</u>		HIKMA FARMACEUTICA	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>A200219</u>	<u>001</u>	Feb 13, 2013
<u>AP</u>		SAGENT PHARMS INC	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>A091676</u>	<u>001</u>	Jul 13, 2018
<u>AP</u>			<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>A091676</u>	<u>002</u>	Jul 13, 2018

PITOCIN

<u>AP</u>	+	PAR STERILE PRODUCTS	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018261</u>	<u>001</u>	
<u>AP</u>	+		<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018261</u>	<u>002</u>	Jul 27, 2007

PRESCRIPTION DRUG PRODUCT LIST

OXYTOCIN

INJECTABLE; INJECTION

OXYTOCIN

+! FRESENIUS KABI USA 300USP UNITS/30ML (10USP UNITS/ML) N018248 003 Jul 27, 2007

PITOCIN

+ PAR STERILE PRODUCTS 500USP UNITS/50ML (10USP UNITS/ML) N018261 003 Sep 05, 2012

OZANIMOD HYDROCHLORIDE

CAPSULE; ORAL

ZEPOSIA

+ CELGENE INTL EQ 0.23MG BASE N209899 001 Mar 25, 2020

+ EQ 0.46MG BASE N209899 002 Mar 25, 2020

+! EQ 0.92MG BASE N209899 003 Mar 25, 2020

OZENOXACIN

CREAM; TOPICAL

XEPI

+! FERRER INTERNACIONAL 1% N208945 001 Dec 11, 2017

PACLITAXEL

INJECTABLE; INJECTION

PACITAXEL**AP** GLAND PHARMA LTD **6MG/ML** **A207326 001** Aug 23, 2016PACLITAXEL**AP** ACCORD HLTHCARE **6MG/ML** **A205720 001** Aug 17, 2018**AP** ACTAVIS TOTOWA **6MG/ML** **A090130 001** Dec 09, 2009**AP** FRESENIUS KABI USA **6MG/ML** **A077574 001** Nov 27, 2006**AP** HIKMA **6MG/ML** **A075190 001** Jan 28, 2002**AP** ! HOSPIRA **6MG/ML** **A076131 001** May 08, 2002**AP** MSN **6MG/ML** **A213434 001** Aug 24, 2020**AP** TEVA PHARMS **6MG/ML** **A075184 001** Jan 25, 2002

POWDER; INTRAVENOUS

ABRAXANE

+! ABRAXIS BIOSCIENCE 100MG/VIAL N021660 001 Jan 07, 2005

PAFOLACIANINE SODIUM

SOLUTION; INTRAVENOUS

CYTALUX

+! ON TARGET LABS EQ 3.2MG BASE/1.6ML (EQ 2MG BASE/ML) N214907 001 Nov 29, 2021

PALBOCICLIB

CAPSULE; ORAL

IBRANCE

+ PFIZER 75MG N207103 001 Feb 03, 2015

+ 100MG N207103 002 Feb 03, 2015

+! 125MG N207103 003 Feb 03, 2015

TABLET; ORAL

IBRANCE

+ PFIZER 75MG N212436 001 Nov 01, 2019

+ 100MG N212436 002 Nov 01, 2019

+! 125MG N212436 003 Nov 01, 2019

PALIPERIDONE

TABLET, EXTENDED RELEASE; ORAL

INVEGA**AB** + JANSSEN PHARMS **1.5MG** **N021999 006** Aug 26, 2008**AB** + **3MG** **N021999 001** Dec 19, 2006**AB** +! **6MG** **N021999 002** Dec 19, 2006**AB** + **9MG** **N021999 003** Dec 19, 2006PALIPERIDONE**AB** ACTAVIS LABS FL INC **1.5MG** **A202645 001** Aug 03, 2015**AB** **3MG** **A202645 002** Aug 03, 2015**AB** **6MG** **A202645 003** Aug 03, 2015**AB** **9MG** **A202645 004** Aug 03, 2015**AB** CSPC OUYI **1.5MG** **A212807 001** Oct 29, 2020**AB** **3MG** **A212807 002** Oct 29, 2020**AB** **6MG** **A212807 003** Oct 29, 2020**AB** **9MG** **A212807 004** Oct 29, 2020**AB** INVENTIA **1.5MG** **A204452 001** Jun 12, 2019**AB** **3MG** **A204452 002** Jun 12, 2019**AB** **6MG** **A204452 003** Jun 12, 2019**AB** **9MG** **A204452 004** Jun 12, 2019**AB** RK PHARMA **1.5MG** **A203802 001** Sep 24, 2015**AB** **3MG** **A203802 002** Sep 24, 2015

PRESCRIPTION DRUG PRODUCT LIST

PALIPERIDONE

TABLET, EXTENDED RELEASE;ORAL

PALIPERIDONE

<u>AB</u>		<u>6MG</u>	<u>A203802</u>	<u>003</u>	Sep 24, 2015
<u>AB</u>		<u>9MG</u>	<u>A203802</u>	<u>004</u>	Sep 24, 2015
<u>AB</u>	SUN PHARM	<u>1.5mg</u>	<u>A205618</u>	<u>001</u>	Apr 06, 2018
<u>AB</u>		<u>3MG</u>	<u>A205618</u>	<u>002</u>	Apr 06, 2018
<u>AB</u>		<u>6MG</u>	<u>A205618</u>	<u>003</u>	Apr 06, 2018
<u>AB</u>		<u>9MG</u>	<u>A205618</u>	<u>004</u>	Apr 06, 2018
BX	AMNEAL PHARMS	1.5MG	A204707	001	Sep 23, 2019
BX		3MG	A204707	002	Sep 23, 2019
BX		6MG	A204707	003	Sep 23, 2019
BX		9MG	A204707	004	Sep 23, 2019

PALIPERIDONE PALMITATE

SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

INVEGA HAFYERA

+	JANSSEN PHARMS	1.092GM/3.5ML (312MG/ML)	N207946	005	Aug 30, 2021
+		1.560GM/5ML (312MG/ML)	N207946	006	Aug 30, 2021

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

PALONOSETRON HYDROCHLORIDE

<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A206916</u>	<u>001</u>	Nov 12, 2021
<u>AP</u>	CIPLA	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A206396</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>	! DR REDDYS LABS LTD	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A201533</u>	<u>002</u>	Apr 21, 2016
<u>AP</u>	EMCURE PHARMS LTD	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A202951</u>	<u>002</u>	Jun 29, 2021
<u>AP</u>	EUGIA PHARMA	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A204702</u>	<u>001</u>	Nov 06, 2018
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A206801</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>		<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A206802</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>	HOSPIRA INC	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A207005</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>	INGENUS PHARMS LLC	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A208789</u>	<u>001</u>	May 22, 2020
<u>AP</u>	MYLAN INSTITUTIONAL	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A206416</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>	QILU PHARM HAINAN	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A205648</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A204289</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>		<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A205870</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>	SANDOZ INC	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A202521</u>	<u>001</u>	Oct 13, 2015
<u>AP</u>	TEVA PHARMS USA	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A090713</u>	<u>001</u>	Mar 23, 2018
<u>AP</u>	VIRTUS PHARM	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A209287</u>	<u>001</u>	Sep 19, 2018
	EMCURE PHARMS LTD	EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)	A202951	001	Jun 29, 2021

SOLUTION;INTRAVENOUS

PALONOSETRON HYDROCHLORIDE

	HIKMA	EQ 0.25MG BASE/2ML (EQ 0.125MG BASE/ML)	N207963	001	Aug 22, 2016
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PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

PAMIDRONATE DISODIUM

<u>AP</u>	AREVA PHARMS	<u>30MG/VIAL</u>	<u>A077433</u>	<u>001</u>	Nov 26, 2008
<u>AP</u>		<u>90MG/VIAL</u>	<u>A077433</u>	<u>003</u>	Nov 26, 2008
<u>AP</u>	DR REDDYS	<u>30MG/10ML (3MG/ML)</u>	<u>A078156</u>	<u>001</u>	Aug 19, 2008
<u>AP</u>		<u>60MG/10ML (6MG/ML)</u>	<u>A078156</u>	<u>002</u>	Aug 19, 2008
<u>AP</u>		<u>90MG/10ML (9MG/ML)</u>	<u>A078156</u>	<u>003</u>	Aug 19, 2008
<u>AP</u>	HIKMA	<u>30MG/VIAL</u>	<u>A075290</u>	<u>001</u>	Apr 30, 2001
<u>AP</u>	+	<u>30MG/10ML (3MG/ML)</u>	<u>N021113</u>	<u>001</u>	Mar 04, 2002
<u>AP</u>		<u>90MG/VIAL</u>	<u>A075290</u>	<u>003</u>	Apr 30, 2001
<u>AP</u>	+	<u>90MG/10ML (9MG/ML)</u>	<u>N021113</u>	<u>002</u>	Mar 04, 2002
<u>AP</u>	!	<u>30MG/10ML (3MG/ML)</u>	<u>A075841</u>	<u>001</u>	Jun 27, 2002
<u>AP</u>	!	<u>60MG/10ML (6MG/ML)</u>	<u>A075841</u>	<u>002</u>	Jun 27, 2002
<u>AP</u>	!	<u>90MG/10ML (9MG/ML)</u>	<u>A075841</u>	<u>003</u>	Jun 27, 2002
<u>AP</u>	MYLAN LABS LTD	<u>30MG/10ML (3MG/ML)</u>	<u>A078520</u>	<u>001</u>	Oct 31, 2008
<u>AP</u>		<u>90MG/10ML (9MG/ML)</u>	<u>A078520</u>	<u>002</u>	Oct 31, 2008

PRESCRIPTION DRUG PRODUCT LIST

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

PAMIDRONATE DISODIUM

<u>AP</u>	SAGENT PHARMS INC	<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
	AREVA PHARMS	60MG/VIAL	A077433 002	Nov 26, 2008

PANCURONIUM BROMIDE

INJECTABLE; INJECTION

PANCURONIUM BROMIDE

<u>AP</u>	! DR REDDYS	<u>1MG/ML</u>	<u>A072759 001</u>	Jul 31, 1990
<u>AP</u>	HOSPIRA	<u>1MG/ML</u>	<u>A072320 001</u>	Jan 19, 1989
	! DR REDDYS	2MG/ML	A072760 001	Jul 31, 1990

PANTOPRAZOLE SODIUM

FOR SUSPENSION, DELAYED RELEASE; ORAL

PANTOPRAZOLE SODIUM

<u>AB</u>	SUN PHARM	<u>EQ 40MG BASE</u>	<u>A213725 001</u>	Jun 30, 2020
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PROTONIX

<u>AB</u>	+! WYETH PHARMS	<u>EQ 40MG BASE</u>	<u>N022020 001</u>	Nov 14, 2007
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INJECTABLE; INTRAVENOUS

PANTOPRAZOLE SODIUM

<u>AP</u>	ACIC PHARMS	<u>EQ 40MG BASE/VIAL</u>	<u>A209524 001</u>	Aug 30, 2021
<u>AP</u>	AKORN	<u>EQ 40MG BASE/VIAL</u>	<u>A079197 001</u>	Nov 08, 2012
<u>AP</u>	EUGIA PHARMA	<u>EQ 40MG BASE/VIAL</u>	<u>A205675 001</u>	Mar 30, 2016
<u>AP</u>	SANDOZ INC	<u>EQ 40MG BASE/VIAL</u>	<u>A090296 001</u>	Jul 14, 2015
<u>AP</u>	SUN PHARM	<u>EQ 40MG BASE/VIAL</u>	<u>A077674 001</u>	Aug 19, 2019

PROTONIX IV

<u>AP</u>	+! WYETH PHARMS	<u>EQ 40MG BASE/VIAL</u>	<u>N020988 001</u>	Mar 22, 2001
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POWDER; INTRAVENOUS

PANTOPRAZOLE SODIUM

+! HIKMA

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>	ATLANTIDE	<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
<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
<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>	INGENUS PHARMS LLC	<u>EQ 40MG BASE</u>	<u>A211368 001</u>	Mar 01, 2019
<u>AB</u>	L PERRIGO CO	<u>EQ 20MG BASE</u>	<u>A203024 001</u>	May 07, 2014
<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>	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>	ORBION PHARMS	<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>	RUBICON	<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>	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

PROTONIX

<u>AB</u>	+ WYETH PHARMS	<u>EQ 20MG BASE</u>	<u>N020987 002</u>	Jun 12, 2001
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<u>AB</u>	+!	<u>EQ 40MG BASE</u>	<u>N020987 001</u>	Feb 02, 2000
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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

PRESCRIPTION DRUG PRODUCT LIST

PARICALCITOL

CAPSULE; ORAL

PARICALCITOL

<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>	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	<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>	DR REDDYS	<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
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A204910 003</u>	Aug 17, 2016
<u>AP</u>	EUGIA PHARMA	<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>	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>	RISING	<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>	+! ABBVIE	<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>	!	<u>EQ 250MG BASE</u>	<u>A064171 001</u>	Jun 30, 1997

PAROXETINE HYDROCHLORIDE

SUSPENSION; ORAL

PAROXETINE HYDROCHLORIDE

<u>AB</u>	NOVITIUM PHARMA	<u>EQ 10MG BASE/5ML</u>	<u>A215003 001</u>	Sep 03, 2021
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PAXIL

<u>AB</u>	+! APOTEX TECHNOLOGIES	<u>EQ 10MG BASE/5ML</u>	<u>N020710 001</u>	Jun 25, 1997
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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

PRESCRIPTION DRUG PRODUCT LIST

PAROXETINE HYDROCHLORIDE

TABLET; ORAL

PAROXETINE HYDROCHLORIDE

<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>	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>	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>	YILING	<u>EQ 10MG BASE</u>	<u>A211248 001</u>	Nov 02, 2021
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A211248 002</u>	Nov 02, 2021
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A211248 003</u>	Nov 02, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A211248 004</u>	Nov 02, 2021
<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
<u>PAXIL</u>				
<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
<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>	AUROBINDO PHARMA USA	<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>	CADILA PHARMS LTD	<u>EQ 12.5MG BASE</u>	<u>A212645 001</u>	Aug 27, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A212645 002</u>	Aug 27, 2021
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A212645 003</u>	Aug 27, 2021
<u>AB</u>	CSPC OUYI	<u>EQ 12.5MG BASE</u>	<u>A213485 001</u>	Feb 16, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213485 002</u>	Feb 16, 2021
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A213485 003</u>	Feb 16, 2021
<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>	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
<u>AB</u>	SINOTHERAPEUTICS INC	<u>EQ 12.5MG BASE</u>	<u>A213612 001</u>	Aug 11, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213612 002</u>	Aug 11, 2021
<u>PAXIL CR</u>				
<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
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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
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PRESCRIPTION DRUG PRODUCT LISTPAROXETINE MESYLATE

TABLET; ORAL

PEXEVA

+		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

+	RECORDATI RARE	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

+	RECORDATI RARE	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

PATIOROMER SORBITEX CALCIUM

POWDER; ORAL

VELTASSA

+	VIFOR PHARMA	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
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PAZOPANIB HYDROCHLORIDE

TABLET; ORAL

VOTRIENT

+	!	NOVARTIS	EQ 200MG BASE	N022465 001	Oct 19, 2009
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PEGCETACOPLAN

SOLUTION; SUBCUTANEOUS

EMPAVELI

+	!	APELLIS PHARMS	1080MG/20ML (54MG/ML)	N215014 001	May 14, 2021
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PEMETREXED

SOLUTION; INTRAVENOUS

PEMFEXY

+	!	EAGLE PHARMS	500MG/20ML (25MG/ML)	N209472 001	Feb 08, 2020
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PEMETREXED DISODIUM

POWDER; INTRAVENOUS

ALIMTA

+	!	LILLY	EQ 100MG BASE/VIAL	N021462 002	Sep 07, 2007
+	!		EQ 500MG BASE/VIAL	N021462 001	Feb 04, 2004

PEMIGATINIB

TABLET; ORAL

PEMAZYRE

+		INCYTE CORP	4.5MG	N213736 001	Apr 17, 2020
+			9MG	N213736 002	Apr 17, 2020
+	!		13.5MG	N213736 003	Apr 17, 2020

PENCICLOVIR

CREAM; TOPICAL

DENA VIR

+	!	MYLAN	1%	N020629 001	Sep 24, 1996
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PENICILLAMINE

CAPSULE; ORAL

CUPRIMINE

AB	+	!	VALEANT PHARMS INTL	250MG	N019853 001
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PENICILLAMINE

AB			ANI PHARMS	250MG	A209921 001	May 07, 2019
AB			APOTEX	250MG	A213310 001	Apr 28, 2020
AB			BRECKENRIDGE	250MG	A215409 001	Aug 23, 2021
AB			DR REDDYS	250MG	A211867 001	Aug 04, 2020
AB			GRANULES	250MG	A211735 001	Dec 02, 2020

PRESCRIPTION DRUG PRODUCT LIST

PENICILLAMINE

CAPSULE; ORAL

PENICILLAMINE

AB	INVAGEN PHARMS	250MG	A213293 001	Aug 19, 2021
AB	NAVINTA LLC	250MG	A214363 001	Oct 08, 2021
AB	PAR PHARM INC	250MG	A211231 001	Dec 23, 2019
AB	WATSON LABS INC	250MG	A210976 001	Jun 24, 2019

TABLET; ORAL

DEPEN

AB	+ !	MYLAN SPECIALITY LP	250MG	N019854 001
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PENICILLAMINE

AB	LUPIN LTD	250MG	A212933 001	Nov 30, 2020
AB	PAR PHARM INC	250MG	A211196 001	Dec 23, 2019

PENICILLIN G BENZATHINE

INJECTABLE; INJECTION

BICILLIN L-A

BC	+ !	KING PHARMS LLC	600,000 UNITS/ML	N050141 001
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PENICILLIN G BENZATHINE; PENICILLIN G PROCAINE

INJECTABLE; INJECTION

BICILLIN C-R

	+ !	KING PHARMS LLC	300,000 UNITS/ML; 300,000 UNITS/ML	N050138 001
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BICILLIN C-R 900/300

	+ !	KING PHARMS LLC	900,000 UNITS/2ML; 300,000 UNITS/2ML	N050138 003
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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	HQ SPECLT PHARMA	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, 2002

PFIZERPEN

AP	!	PFIZER	5,000,000 UNITS/VIAL	A060657 002
AP	!		20,000,000 UNITS/VIAL	A060657 003
		PENICILLIN G POTASSIUM		
		HQ SPECLT PHARMA	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

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
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PENICILLIN V POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN-VK

TEVA

			EQ 125MG BASE/5ML	A060456 001
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!

			EQ 250MG BASE/5ML	A060456 002
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TABLET; ORAL

PENICILLIN V POTASSIUM

AB	AUROBINDO PHARMA	EQ 250MG BASE	A065435 001	Apr 29, 2008
AB		EQ 500MG BASE	A065435 002	Apr 29, 2008
AB	HIKMA PHARMS	EQ 250MG BASE	A090549 001	Oct 11, 2013
AB		EQ 500MG BASE	A090549 002	Oct 11, 2013
AB	SANDOZ	EQ 250MG BASE	A064071 001	Nov 30, 1995
AB	!		EQ 500MG BASE	A064071 002
AB		EQ 250MG BASE	A062936 001	Nov 25, 1988
AB	STRIDES PHARMA	EQ 500MG BASE	A062935 001	Nov 23, 1988

PRESCRIPTION DRUG PRODUCT LIST

PENICILLIN V POTASSIUM

TABLET; ORAL

PENICILLIN-VK

<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>A060711</u>	<u>002</u>	
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A060711</u>	<u>003</u>	

PENTAMIDINE ISETHIONATE

FOR SOLUTION; INHALATION

NEBUPENT

<u>AN</u>	+!	FRESENIUS KABI USA	<u>300MG/VIAL</u>	<u>N019887</u>	<u>001</u>	Jun 15, 1989
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PENTAMIDINE ISETHIONATE

<u>AN</u>		SETON PHARMS	<u>300MG/VIAL</u>	<u>A206667</u>	<u>001</u>	Apr 24, 2019
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INJECTABLE; INJECTION

PENTAM

<u>AP</u>	+!	FRESENIUS KABI USA	<u>300MG/VIAL</u>	<u>N019264</u>	<u>001</u>	Oct 16, 1984
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PENTAMIDINE ISETHIONATE

<u>AP</u>		EMCURE PHARMS LTD	<u>300MG/VIAL</u>	<u>A213806</u>	<u>001</u>	Jan 07, 2021
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<u>AP</u>		SETON PHARMS	<u>300MG/VIAL</u>	<u>A206666</u>	<u>001</u>	Sep 28, 2017
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PENTOBARBITAL SODIUM

INJECTABLE; INJECTION

NEMBUTAL SODIUM

<u>AP</u>	+!	AKORN	<u>50MG/ML</u>	<u>A083246</u>	<u>001</u>	
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PENTOBARBITAL SODIUM

<u>AP</u>		ATHENEX	<u>50MG/ML</u>	<u>A206677</u>	<u>001</u>	Nov 27, 2017
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<u>AP</u>		CUSTOPHARM INC	<u>50MG/ML</u>	<u>A203619</u>	<u>001</u>	Nov 13, 2017
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<u>AP</u>		SAGENT PHARMS INC	<u>50MG/ML</u>	<u>A206404</u>	<u>001</u>	May 23, 2016
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PENTOSAN POLYSULFATE SODIUM

CAPSULE; ORAL

ELMIRON

	+!	JANSSEN PHARMS	100MG	N020193	001	Sep 26, 1996
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PENTOSTATIN

INJECTABLE; INJECTION

NIPENT

<u>AP</u>	+!	HOSPIRA INC	<u>10MG/VIAL</u>	<u>N020122</u>	<u>001</u>	Oct 11, 1991
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PENTOSTATIN

<u>AP</u>		WEST-WARD PHARMS INT	<u>10MG/VIAL</u>	<u>A077841</u>	<u>001</u>	Aug 07, 2007
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PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL

PENTOXIFYLLINE

<u>AB</u>	!	APOTEX	<u>400MG</u>	<u>A075191</u>	<u>001</u>	Jun 09, 1999
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<u>AB</u>		VALEANT PHARMS	<u>400MG</u>	<u>A075028</u>	<u>001</u>	Jul 20, 1998
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PERAMIVIR

SOLUTION; INTRAVENOUS

RAPIVAB

	+!	BIOCRIST	200MG/20ML (10MG/ML)	N206426	001	Dec 19, 2014
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PERAMPANEL

SUSPENSION; ORAL

FYCOMPA

	+!	EISAI INC	0.5MG/ML	N208277	001	Apr 29, 2016
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TABLET; ORAL

FYCOMPA

	+	EISAI INC	2MG	N202834	001	Oct 22, 2012
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	+		4MG	N202834	002	Oct 22, 2012
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	+		6MG	N202834	003	Oct 22, 2012
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	+		8MG	N202834	004	Oct 22, 2012
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	+		10MG	N202834	005	Oct 22, 2012
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	+!		12MG	N202834	006	Oct 22, 2012
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PERFLUTREN

INJECTABLE; INTRAVENOUS

DEFINITY

	+!	LANTHEUS MEDCL	6.52MG/ML	N021064	001	Jul 31, 2001
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DEFINITY RT

	+!	LANTHEUS MEDCL	6.52MG/ML	N021064	002	Nov 17, 2020
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PRESCRIPTION DRUG PRODUCT LIST

PERINDOPRIL ERBUMINE

TABLET; ORAL

PERINDOPRIL ERBUMINE

<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>	HIKMA	<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

PERMETHRIN

<u>AB</u>	ACTAVIS LABS	<u>5%</u>	<u>A074806</u>	<u>001</u>	Jan 23, 1998
<u>AB</u>	ENCUBE ETHICALS	<u>5%</u>	<u>A211303</u>	<u>001</u>	Apr 03, 2019
<u>AB</u>	! PADAGIS ISRAEL	<u>5%</u>	<u>A076369</u>	<u>001</u>	Apr 21, 2003

PERPHENAZINE

TABLET; ORAL

PERPHENAZINE

<u>AB</u>	MYLAN	<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>	RISING	<u>2MG</u>	<u>A205056</u>	<u>001</u>	Mar 01, 2019
<u>AB</u>		<u>4MG</u>	<u>A205056</u>	<u>002</u>	Mar 01, 2019
<u>AB</u>		<u>8MG</u>	<u>A205056</u>	<u>003</u>	Mar 01, 2019
<u>AB</u>		<u>16MG</u>	<u>A205056</u>	<u>004</u>	Mar 01, 2019
<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
<u>AB</u>		<u>16MG</u>	<u>A205973</u>	<u>004</u>	Dec 17, 2015
<u>AB</u>	ZYDUS PHARMS	<u>2MG</u>	<u>A205232</u>	<u>001</u>	Apr 06, 2020
<u>AB</u>		<u>4MG</u>	<u>A205232</u>	<u>002</u>	Apr 06, 2020
<u>AB</u>		<u>8MG</u>	<u>A205232</u>	<u>003</u>	Apr 06, 2020
<u>AB</u>		<u>16MG</u>	<u>A205232</u>	<u>004</u>	Apr 06, 2020

PEXIDARTINIB HYDROCHLORIDE

CAPSULE; ORAL

TURALIO

+	!	DAIICHI SANKYO INC	EQ 200MG BASE	N211810	001	Aug 02, 2019
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PHENDIMETRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

PHENDIMETRAZINE TARTRATE

+	!	VIRTUS PHARMS	105MG	N018074	001	
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TABLET; ORAL

BONTRIL PDM

<u>AA</u>	+	!	VALEANT	<u>35MG</u>	<u>A085272</u>	<u>001</u>
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PHENDIMETRAZINE TARTRATE

<u>AA</u>			ELITE LABS INC	<u>35MG</u>	<u>A040762</u>	<u>001</u>	Jan 28, 2008
<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>
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PHENELZINE SULFATE

<u>AB</u>			NOVEL LABS INC	<u>EQ 15MG BASE</u>	<u>A200181</u>	<u>001</u>	Dec 08, 2010
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PRESCRIPTION DRUG PRODUCT LIST

PHENOXYBENZAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIBENZYLINE

<u>AB</u>	+	CONCORDIA	<u>10MG</u>	<u>N008708</u>	<u>001</u>	
<u>PHENOXYBENZAMINE HYDROCHLORIDE</u>						
<u>AB</u>		AMNEAL	<u>10MG</u>	<u>A212568</u>	<u>001</u>	Oct 27, 2020
<u>AB</u>		HIKMA	<u>10MG</u>	<u>A201050</u>	<u>001</u>	Jul 16, 2012
<u>AB</u>	!	PAR PHARM INC	<u>10MG</u>	<u>A204522</u>	<u>001</u>	Jan 24, 2017

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

ADIPEX-P

<u>AA</u>	+	TEVA	<u>37.5MG</u>	<u>A088023</u>	<u>001</u>	Aug 02, 1983
<u>PHENTERMINE HYDROCHLORIDE</u>						
<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>		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>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>	
<u>LOMAIRA</u>						
<u>AA</u>	!	AVANTHI INC	<u>8MG</u>	<u>A203495</u>	<u>001</u>	Sep 12, 2016
<u>PHENTERMINE HYDROCHLORIDE</u>						
<u>AA</u>		AUROLIFE PHARMA LLC	<u>37.5MG</u>	<u>A203068</u>	<u>001</u>	Aug 06, 2014
<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>		MERRO PHARM USA	<u>37.5MG</u>	<u>A206342</u>	<u>001</u>	Nov 18, 2016
<u>AA</u>		NUVO PHARM	<u>37.5MG</u>	<u>A205008</u>	<u>001</u>	Sep 25, 2014
<u>AA</u>		PRINSTON INC	<u>37.5MG</u>	<u>A040377</u>	<u>001</u>	Jan 04, 2002
<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	15MG	A204663	001	Jun 28, 2017
		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

PRESCRIPTION DRUG PRODUCT LIST

PHENTOLAMINE MESYLATE

INJECTABLE; INJECTION

PHENTOLAMINE MESYLATE

<u>AP</u>	!	HIKMA	<u>5MG/VIAL</u>	<u>A040235</u>	<u>001</u>	Mar 11, 1998
<u>AP</u>		PRECISION DOSE INC	<u>5MG/VIAL</u>	<u>A207686</u>	<u>001</u>	Jul 14, 2017
		ORAVERSE				
	+	SEPTODONT HOLDING	0.4MG/1.7ML	N022159	001	May 09, 2008

PHENYLEPHRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

PHENYLEPHRINE HYDROCHLORIDE

<u>AP1</u>		AMNEAL	<u>10MG/ML (10MG/ML)</u>	<u>A211079</u>	<u>001</u>	Jul 05, 2018
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A211078</u>	<u>001</u>	Jul 19, 2018
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A211078</u>	<u>002</u>	Jul 19, 2018
<u>AP1</u>		CAPLIN	<u>10MG/ML (10MG/ML)</u>	<u>A213318</u>	<u>001</u>	Jun 11, 2020
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A213318</u>	<u>002</u>	Jun 11, 2020
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A213318</u>	<u>003</u>	Jun 11, 2020
<u>AP1</u>		EUGIA PHARMA	<u>50MG/5ML (10MG/ML)</u>	<u>A210697</u>	<u>001</u>	Nov 13, 2020
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A210697</u>	<u>002</u>	Nov 13, 2020
<u>AP1</u>		FRESENIUS KABI USA	<u>50MG/5ML (10MG/ML)</u>	<u>A210666</u>	<u>001</u>	Jan 30, 2019
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A210666</u>	<u>002</u>	Jan 30, 2019
<u>AP1</u>		GLAND PHARMA LTD	<u>10MG/ML (10MG/ML)</u>	<u>A211920</u>	<u>003</u>	Apr 08, 2021
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A211920</u>	<u>001</u>	Jun 05, 2020
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A211920</u>	<u>002</u>	Jun 05, 2020
<u>AP1</u>		MEITHEAL	<u>50MG/5ML (10MG/ML)</u>	<u>A210333</u>	<u>001</u>	Apr 27, 2018
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A210333</u>	<u>002</u>	Apr 27, 2018
<u>AP1</u>		PAR STERILE PRODUCTS	<u>10MG/ML (10MG/ML)</u>	<u>A210025</u>	<u>001</u>	Dec 21, 2018
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A210025</u>	<u>002</u>	Dec 21, 2018
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A210025</u>	<u>003</u>	Dec 21, 2018
<u>AP1</u>		PROVEPHARM SAS	<u>10MG/ML (10MG/ML)</u>	<u>A211081</u>	<u>001</u>	Jul 17, 2020
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A211081</u>	<u>002</u>	Jul 17, 2020
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A211081</u>	<u>003</u>	Jul 17, 2020
<u>AP1</u>		SAGENT PHARMS INC	<u>10MG/ML (10MG/ML)</u>	<u>A209967</u>	<u>001</u>	Jan 16, 2020
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A209967</u>	<u>002</u>	Jan 16, 2020
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A209967</u>	<u>003</u>	Jan 16, 2020
<u>AP1</u>		SANDOZ INC	<u>10MG/ML (10MG/ML)</u>	<u>A208905</u>	<u>001</u>	Jan 31, 2019
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A208905</u>	<u>002</u>	Jan 31, 2019
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A208905</u>	<u>003</u>	Jan 31, 2019

VAZCULEP

<u>AP1</u>	+	EXELA PHARMA	<u>10MG/ML (10MG/ML)</u>	<u>N204300</u>	<u>001</u>	Jun 27, 2014
<u>AP1</u>	+		<u>50MG/5ML (10MG/ML)</u>	<u>N204300</u>	<u>002</u>	Jun 27, 2014
<u>AP1</u>	+		<u>100MG/10ML (10MG/ML)</u>	<u>N204300</u>	<u>003</u>	Jun 27, 2014

PHENYLEPHRINE HYDROCHLORIDE

<u>AP2</u>		EUGIA PHARMA	<u>10MG/ML (10MG/ML)</u>	<u>A210696</u>	<u>001</u>	Jan 07, 2021
<u>AP2</u>		FRESENIUS KABI USA	<u>10MG/ML (10MG/ML)</u>	<u>A210665</u>	<u>001</u>	Jan 29, 2019
<u>AP2</u>	+	HIKMA	<u>10MG/ML (10MG/ML)</u>	<u>N203826</u>	<u>001</u>	Dec 20, 2012
<u>AP2</u>	+		<u>50MG/5ML (10MG/ML)</u>	<u>N203826</u>	<u>002</u>	Jun 19, 2019
<u>AP2</u>	+		<u>100MG/10ML (10MG/ML)</u>	<u>N203826</u>	<u>003</u>	Jun 19, 2019
<u>AP2</u>		MEITHEAL	<u>10MG/ML (10MG/ML)</u>	<u>A210334</u>	<u>001</u>	Apr 27, 2018
		BIORPHEN				
	+	ETON PHARMS	0.5MG/5ML (0.1MG/ML)	N212909	001	Oct 21, 2019

SOLUTION/DROPS; OPHTHALMIC

PHENYLEPHRINE HYDROCHLORIDE

<u>AT</u>	+	AKORN	<u>2.5%</u>	<u>N207926</u>	<u>001</u>	Jan 15, 2015
<u>AT</u>	+		<u>10%</u>	<u>N207926</u>	<u>002</u>	Jan 15, 2015
<u>AT</u>	+	PARAGON BIOTECK	<u>2.5%</u>	<u>N203510</u>	<u>001</u>	Mar 21, 2013
<u>AT</u>	+		<u>10%</u>	<u>N203510</u>	<u>002</u>	Mar 21, 2013

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE

<u>AA</u>		AKORN	<u>5MG/5ML; 6.25MG/5ML</u>	<u>A040675</u>	<u>001</u>	Dec 23, 2014
<u>AA</u>	!	PHARM ASSOC	<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

PHENYTOIN

SUSPENSION; ORAL

DILANTIN-125

<u>AB</u>	+	UPJOHN	<u>125MG/5ML</u>	<u>N008762</u>	<u>001</u>	
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PHENYTOIN

<u>AB</u>		TARO	<u>125MG/5ML</u>	<u>A040521</u>	<u>001</u>	Mar 08, 2004
<u>AB</u>		VISTAPHARM	<u>125MG/5ML</u>	<u>A040342</u>	<u>001</u>	Jan 31, 2001

PRESCRIPTION DRUG PRODUCT LIST

PHENYTOIN

SUSPENSION; ORAL

PHENYTOIN

AB		125MG/5ML	A040610 001	Aug 18, 2005
AB	WOCKHARDT BIO AG	125MG/5ML	A040420 001	Apr 19, 2002

TABLET, CHEWABLE; ORAL

DILANTIN

AB	+ !	PHARMACIA	50MG	A084427 001
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PHENYTOIN

AB		EPIC PHARMA LLC	50MG	A040884 001	Nov 28, 2014
AB		RISING	50MG	A200691 001	Dec 26, 2012
AB		TARO	50MG	A200565 001	Apr 17, 2014

PHENYTOIN SODIUM

CAPSULE; ORAL

DILANTIN

AB	+ !	UPJOHN	100MG EXTENDED	A084349 002
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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		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
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DILANTIN

+!

UPJOHN

30MG EXTENDED

A084349 001

INJECTABLE; INJECTION

PHENYTOIN SODIUM

AP		ACELLA	50MG/ML	A040573 001	Sep 13, 2006
AP	+ !	WEST-WARD PHARMS INT	50MG/ML	A084307 001	

PHYTONADIONE

INJECTABLE; INJECTION

PHYTONADIONE

AB		DR REDDYS LABS LTD	10MG/ML	A207719 001	May 22, 2019
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VITAMIN K1

AB	+ !	HOSPIRA	10MG/ML	A087955 001	Jul 25, 1983
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PHYTONADIONE

BP	+ !	INTL MEDICATION	1MG/0.5ML	A083722 001	
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VITAMIN K1

BP	+ !	HOSPIRA	1MG/0.5ML	A087954 001	Jul 25, 1983
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TABLET; ORAL

MEPHYTON

AB	+ !	BAUSCH	5MG	N010104 003
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PHYTONADIONE

AB		AMNEAL PHARMS CO	5MG	A209373 001	May 11, 2018
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AB		ZYDUS	5MG	A210189 001	Feb 20, 2019
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PIFLUFOLASTAT F-18

SOLUTION; INTRAVENOUS

PYLARIFY

+!

PROGENICS PHARMS INC

50ML (1-80mCi/ML)

N214793 001

May 26, 2021

PILOCARPINE HYDROCHLORIDE

SOLUTION; OPHTHALMIC

ISOPTO CARPINE

AT	+	NOVARTIS	1%	N200890 001	Jun 22, 2010
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AT	+		2%	N200890 002	Jun 22, 2010
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AT	+ !		4%	N200890 003	Jun 22, 2010
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PILOCARPINE HYDROCHLORIDE

AT		AKORN	1%	A204398 001	Sep 27, 2017
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AT			2%	A204398 002	Sep 27, 2017
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AT			4%	A204398 003	Sep 27, 2017
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AT		AMNEAL	1%	A214193 001	Sep 21, 2020
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AT			2%	A214193 002	Sep 21, 2020
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AT			4%	A214193 003	Sep 21, 2020
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AT		SOMERSET THERAPS LLC	1%	A210384 001	Nov 25, 2019
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PRESCRIPTION DRUG PRODUCT LIST

PILOCARPINE HYDROCHLORIDE

SOLUTION;OPHTHALMIC

PILOCARPINE HYDROCHLORIDE

AT		2%	A210384 002	Nov 25, 2019
AT		4%	A210384 003	Nov 25, 2019

VUITY

+	ABBVIE INC	1.25%	N214028 001	Oct 28, 2021
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TABLET;ORAL

PILOCARPINE HYDROCHLORIDE

AB	AMICI	5MG	A076963 001	Dec 22, 2004
AB		7.5MG	A076963 002	Feb 27, 2007
AB	AUROBINDO PHARMA LTD	5MG	A212377 001	Aug 13, 2019
AB		7.5MG	A212377 002	Aug 13, 2019
AB	IMPAX LABS	5MG	A077248 001	Mar 31, 2006
AB		7.5MG	A077248 002	Mar 31, 2006
AB	LANNETT CO INC	5MG	A077220 001	Oct 14, 2005
AB		7.5MG	A077220 002	May 06, 2009
AB	PADAGIS US	5MG	A076746 001	Nov 16, 2004

SALAGEN

AB	+	CONCORDIA	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
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TABLET;ORAL

NUPLAZID

+	ACADIA PHARMS INC	EQ 10MG BASE	N207318 002	Jun 28, 2018
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PIMECROLIMUS

CREAM;TOPICAL

ELIDEL

AB	+	BAUSCH	1%	N021302 001	Dec 13, 2001
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PIMECROLIMUS

AB		ACTAVIS LABS UT INC	1%	A209345 001	Dec 27, 2018
AB		GLENMARK PHARMS LTD	1%	A211769 001	Aug 29, 2019

PIMOZIDE

TABLET;ORAL

PIMOZIDE

	PAR PHARM	1MG	A204521 001	Sep 28, 2015
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!		2MG	A204521 002	Sep 28, 2015
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PINDOLOL

TABLET;ORAL

PINDOLOL

AB		ANI PHARMS	5MG	A073609 002	Mar 29, 1993
AB			10MG	A073609 001	Mar 29, 1993
AB		BAYSHORE PHARMS LLC	5MG	A211712 001	Aug 01, 2019
AB			10MG	A211712 002	Aug 01, 2019
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

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		ANNORA PHARMA	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		AUROBINDO PHARMA LTD	EQ 15MG BASE	A200268 001	Feb 13, 2013

PRESCRIPTION DRUG PRODUCT LIST

PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

PIOGLITAZONE HYDROCHLORIDE

<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A200268 002</u>	Feb 13, 2013
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A200268 003</u>	Feb 13, 2013
<u>AB</u>	CELLTRION	<u>EQ 15MG BASE</u>	<u>A076798 001</u>	Oct 26, 2012
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A076798 002</u>	Oct 26, 2012
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A076798 003</u>	Oct 26, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 15MG BASE</u>	<u>A202467 001</u>	Feb 06, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A202467 002</u>	Feb 06, 2013
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A202467 003</u>	Feb 06, 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>	SHUANGCHENG	<u>EQ 15MG BASE</u>	<u>A210165 001</u>	Jan 22, 2021
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A210165 002</u>	Jan 22, 2021
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A210165 003</u>	Jan 22, 2021
<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>	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>	ASTRAL	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A212287 001</u>	Jul 29, 2019
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A212287 002</u>	Jul 29, 2019
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A212287 003</u>	Jul 29, 2019
<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>	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>	PROVEPHARM SAS	<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>	QILU	<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>	SAGENT PHARMS INC	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A208674 001</u>	Feb 16, 2021
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A208674 002</u>	Feb 16, 2021
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A208674 003</u>	Feb 16, 2021
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A208675 001</u>	Feb 16, 2021
<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>A065363 001</u>	Oct 21, 2010

PRESCRIPTION DRUG PRODUCT LIST

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

PIPERACILLIN AND TAZOBACTAM

AP		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065362 002</u>	Oct 21, 2010
AP		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065363 002</u>	Oct 21, 2010
AP		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065362 003</u>	Oct 21, 2010
AP		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065363 003</u>	Oct 21, 2010
AP	SANDOZ INC	<u>EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL</u>	<u>A203557 001</u>	Oct 29, 2014
AP		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A203557 002</u>	Jul 08, 2021
AP	! WOCKHARDT BIO AG	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A206996 001</u>	Mar 22, 2017
AP	!	<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A206996 002</u>	Mar 22, 2017
AP	!	<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A206996 003</u>	Mar 22, 2017
AP	!	<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A207146 001</u>	Mar 17, 2017
	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

AB	+! GENENTECH INC	<u>267MG</u>	<u>N022535 001</u>	Oct 15, 2014
	<u>PIRFENIDONE</u>			
AB	AMNEAL	<u>267MG</u>	<u>A212569 001</u>	Jan 03, 2022
	TABLET; ORAL			
	ESBRIET			
	+ GENENTECH INC	267MG	N208780 001	Jan 11, 2017
	+!	801MG	N208780 003	Jan 11, 2017

PIROXICAM

CAPSULE; ORAL

FELDENE

AB	+ PFIZER	<u>10MG</u>	<u>N018147 002</u>	Apr 06, 1982
AB	+!	<u>20MG</u>	<u>N018147 003</u>	Apr 06, 1982

PIROXICAM

AB	CADILA	<u>10MG</u>	<u>A205585 001</u>	Jul 17, 2018
AB		<u>20MG</u>	<u>A205585 002</u>	Jul 17, 2018
AB	FLAMINGO PHARMS	<u>10MG</u>	<u>A207938 001</u>	Sep 09, 2016
AB		<u>20MG</u>	<u>A207938 002</u>	Sep 09, 2016
AB	HIKMA	<u>10MG</u>	<u>A209256 001</u>	Aug 11, 2017
AB		<u>20MG</u>	<u>A209256 002</u>	Aug 11, 2017
AB	MICRO LABS	<u>10MG</u>	<u>A206152 001</u>	Dec 29, 2017
AB		<u>20MG</u>	<u>A206152 002</u>	Dec 29, 2017
AB	NOSTRUM LABS INC	<u>10MG</u>	<u>A074118 002</u>	Jun 15, 1993
AB		<u>20MG</u>	<u>A074118 001</u>	Jun 15, 1993
AB	STRIDES PHARMA	<u>10MG</u>	<u>A206136 001</u>	Jun 20, 2017
AB		<u>10MG</u>	<u>A210347 001</u>	Jan 26, 2018
AB		<u>20MG</u>	<u>A206136 002</u>	Jun 20, 2017
AB		<u>20MG</u>	<u>A210347 002</u>	Jan 26, 2018
AB	TEVA	<u>10MG</u>	<u>A074131 001</u>	Dec 11, 1992
AB		<u>20MG</u>	<u>A074131 002</u>	Dec 11, 1992
BX	UNICHEM	10MG	A208340 001	Apr 13, 2017
BX		20MG	A208340 002	Apr 13, 2017

PITAVASTATIN CALCIUM

TABLET; ORAL

LIVALO

AB	+ KOWA CO	<u>EQ 1MG BASE</u>	<u>N022363 001</u>	Aug 03, 2009
AB	+	<u>EQ 2MG BASE</u>	<u>N022363 002</u>	Aug 03, 2009
AB	+!	<u>EQ 4MG BASE</u>	<u>N022363 003</u>	Aug 03, 2009

PITAVASTATIN CALCIUM

AB	AUROBINDO PHARMA LTD	<u>EQ 1MG BASE</u>	<u>A206015 001</u>	Dec 20, 2016
AB		<u>EQ 2MG BASE</u>	<u>A206015 002</u>	Dec 20, 2016
AB		<u>EQ 4MG BASE</u>	<u>A206015 003</u>	Dec 20, 2016
AB	ORIENT PHARMA CO LTD	<u>EQ 1MG BASE</u>	<u>A205932 001</u>	Feb 03, 2017
AB		<u>EQ 2MG BASE</u>	<u>A205932 002</u>	Feb 03, 2017
AB		<u>EQ 4MG BASE</u>	<u>A205932 003</u>	Feb 03, 2017
AB	SAWAI USA	<u>EQ 1MG BASE</u>	<u>A205955 001</u>	Feb 03, 2017
AB		<u>EQ 2MG BASE</u>	<u>A205955 002</u>	Feb 03, 2017
AB		<u>EQ 4MG BASE</u>	<u>A205955 003</u>	Feb 03, 2017

PRESCRIPTION DRUG PRODUCT LIST

PITAVASTATIN MAGNESIUM

TABLET; ORAL

ZYPITAMAG

+ MEDICURE

EQ 2MG BASE

N208379 002 Jul 14, 2017

+!

EQ 4MG BASE

N208379 003 Jul 14, 2017

PITOLISANT HYDROCHLORIDE

TABLET; ORAL

WAKIX

+ HARMONY

EQ 4.45MG BASE

N211150 001 Aug 14, 2019

+!

EQ 17.8MG BASE

N211150 002 Aug 14, 2019

PLAZOMICIN SULFATE

SOLUTION; INTRAVENOUS

ZEMDRI

+! CIPLA USA

EQ 500MG BASE/10ML (EQ 50MG BASE/ML)

N210303 001 Jun 25, 2018

PLECANATIDE

TABLET; ORAL

TRULANCE

+! SALIX

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

0.5%

N020529 001 Mar 13, 1997

SOLUTION; TOPICAL

CONDYLOX**AT** +! TEVA BRANDED PHARM**0.5%****N019795 001** Dec 13, 1990PODOFILOX**AT** PADAGIS US**0.5%****A075600 001** Jan 29, 2002POLIDOCANOL

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; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION; ORAL

LAX-LYTE WITH FLAVOR PACKS**AA** L PERRIGO CO**420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/
BOT****A079232 001** Feb 25, 2010NULYTELY**AA** +! BRAINTREE**420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/
BOT****N019797 001** Apr 22, 1991NULYTELY-FLAVORED**AA** +! BRAINTREE**420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/
BOT****N019797 002** Nov 18, 1994PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE**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, 2015POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SOLUTION; ORAL

COLYTE WITH FLAVOR PACKS**AA** MYLAN SPECIALITY LP**240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/
BOT; 22.72GM/BOT****N018983 012** Oct 08, 1998GOLYTELY**AA** +! BRAINTREE**236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/
BOT; 22.74GM/BOT****N019011 001** Jul 13, 1984PEG 3350 AND ELECTROLYTES**AA** NOVEL LABS INC**236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/
BOT; 22.74GM/BOT****A090231 001** Jun 01, 2009**AA****240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/
BOT; 22.72GM/BOT****A090186 001** Jun 01, 2009

PRESCRIPTION DRUG PRODUCT LIST

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SOLUTION;ORAL

POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES

AA	STRIDES PHARMA	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	A204558 001	Dec 21, 2018
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POLYMYXIN B SULFATE

INJECTABLE;INJECTION

POLYMYXIN B SULFATE

AP	EUGIA PHARMA	EQ 500,000 UNITS BASE/VIAL	A206589 001	Apr 04, 2016
AP	FRESENIUS KABI USA	EQ 500,000 UNITS BASE/VIAL	A065372 001	Jan 10, 2008
AP	! XELLIA PHARMS APS	EQ 500,000 UNITS BASE/VIAL	A202766 001	Jan 15, 2014
AP	XGEN PHARMS	EQ 500,000 UNITS BASE/VIAL	A063000 001	Sep 30, 1994

POLYMYXIN B SULFATE; TRIMETHOPRIM SULFATE

SOLUTION/DROPS;OPHTHALMIC

POLYTRIM

AT	+! ALLERGAN	10,000 UNITS/ML;EQ 1MG BASE/ML	N050567 001	Oct 20, 1988
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TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE

AT	AKORN	10,000 UNITS/ML;EQ 1MG BASE/ML	A065006 001	Dec 17, 1998
AT	BAUSCH AND LOMB	10,000 UNITS/ML;EQ 1MG BASE/ML	A064120 001	Feb 14, 1997
AT	SANDOZ INC	10,000 UNITS/ML;EQ 1MG BASE/ML	A064211 001	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

+	TAKEDA PHARMS USA	EQ 10MG BASE	N203469 004	Dec 18, 2020
+		EQ 15MG BASE	N203469 001	Dec 14, 2012
+		EQ 30MG BASE	N203469 003	Apr 23, 2015
+!		EQ 45MG BASE	N203469 002	Dec 14, 2012

PONESIMOD

TABLET;ORAL

PONVORY

+	JANSSEN PHARMS	2MG	N213498 001	Mar 18, 2021
+		3MG	N213498 002	Mar 18, 2021
+		4MG	N213498 003	Mar 18, 2021
+		5MG	N213498 004	Mar 18, 2021
+		6MG	N213498 005	Mar 18, 2021
+		7MG	N213498 006	Mar 18, 2021
+		8MG	N213498 007	Mar 18, 2021
+		9MG	N213498 008	Mar 18, 2021
+		10MG	N213498 009	Mar 18, 2021
+!		20MG	N213498 010	Mar 18, 2021

PORFIMER SODIUM

INJECTABLE;INJECTION

PHOTOFRIN

	PINNACLE BIOLGS	75MG/VIAL	N020451 001	Dec 27, 1995
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POSACONAZOLE

FOR SUSPENSION, DELAYED RELEASE;ORAL

NOXAFIL POWDERMIX KIT

+	MSD MERCK CO	300MG	N214770 001	May 31, 2021
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SOLUTION;INTRAVENOUS

NOXAFIL

+	MERCK SHARP DOHME	300MG/16.7ML (18MG/ML)	N205596 001	Mar 13, 2014
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SUSPENSION;ORAL

NOXAFIL

+	SCHERING	40MG/ML	N022003 001	Sep 15, 2006
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TABLET, DELAYED RELEASE;ORAL

NOXAFIL

AB	+! MERCK SHARP DOHME	100MG	N205053 001	Nov 25, 2013
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POSACONAZOLE

AB	AET PHARMA	100MG	A213454 001	Feb 01, 2021
AB	SINOTHERAPEUTICS INC	100MG	A212411 001	Aug 21, 2019

PRESCRIPTION DRUG PRODUCT LIST

POTASSIUM ACETATE

INJECTABLE; INJECTION

POTASSIUM ACETATE

<u>AP</u>	EXELA PHARMA	<u>2MEQ/ML</u>	<u>A206203</u>	<u>001</u>	Dec 29, 2015
<u>AP</u>		<u>2MEQ/ML</u>	<u>A212692</u>	<u>001</u>	Oct 20, 2021
<u>AP</u>	+! HOSPIRA	<u>2MEQ/ML</u>	<u>N018896</u>	<u>001</u>	Jul 20, 1984

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

KLOR-CON

<u>AB</u>	UPSHER SMITH LABS	<u>8MEQ</u>	<u>A203106</u>	<u>001</u>	Jul 10, 2015
<u>AB</u>		<u>10MEQ</u>	<u>A203106</u>	<u>002</u>	Jul 10, 2015

POTASSIUM CHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>8MEQ</u>	<u>A077419</u>	<u>001</u>	Jun 02, 2008
<u>AB</u>	!	<u>10MEQ</u>	<u>A077419</u>	<u>002</u>	Jun 02, 2008
<u>AB</u>	ADARE PHARMS INC	<u>8MEQ</u>	<u>A208864</u>	<u>001</u>	Mar 17, 2017
<u>AB</u>		<u>10MEQ</u>	<u>A208864</u>	<u>002</u>	Mar 17, 2017
<u>AB</u>	AMNEAL PHARMS	<u>10MEQ</u>	<u>A202128</u>	<u>001</u>	Feb 22, 2013
<u>AB</u>	ANCHEN PHARMS	<u>8MEQ</u>	<u>A202886</u>	<u>001</u>	Dec 26, 2013
<u>AB</u>		<u>10MEQ</u>	<u>A202886</u>	<u>002</u>	Dec 26, 2013
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MEQ</u>	<u>A202868</u>	<u>001</u>	Jan 19, 2016
<u>AB</u>	GRANULES	<u>8MEQ</u>	<u>A214686</u>	<u>001</u>	Feb 16, 2021
<u>AB</u>		<u>10MEQ</u>	<u>A214686</u>	<u>002</u>	Feb 16, 2021
<u>AB</u>	LANNETT CO INC	<u>8MEQ</u>	<u>A204210</u>	<u>001</u>	Mar 28, 2016
<u>AB</u>		<u>10MEQ</u>	<u>A204210</u>	<u>002</u>	Mar 28, 2016
<u>AB</u>	LUPIN LTD	<u>8MEQ</u>	<u>A203002</u>	<u>001</u>	Dec 18, 2015
<u>AB</u>		<u>10MEQ</u>	<u>A203002</u>	<u>002</u>	Dec 18, 2015
<u>AB</u>	NOVEL LABS INC	<u>8MEQ</u>	<u>A204828</u>	<u>001</u>	Aug 16, 2016
<u>AB</u>		<u>10MEQ</u>	<u>A204828</u>	<u>002</u>	Aug 16, 2016
<u>AB</u>	PADAGIS US	<u>8MEQ</u>	<u>A200185</u>	<u>001</u>	May 18, 2011
<u>AB</u>		<u>10MEQ</u>	<u>A200185</u>	<u>002</u>	May 18, 2011
<u>AB</u>	PRINSTON INC	<u>8MEQ</u>	<u>A209026</u>	<u>001</u>	Jun 11, 2019
<u>AB</u>		<u>10MEQ</u>	<u>A209026</u>	<u>002</u>	Jun 11, 2019
<u>AB</u>	STRIDES PHARMA	<u>8MEQ</u>	<u>A205549</u>	<u>001</u>	Dec 08, 2015
<u>AB</u>		<u>10MEQ</u>	<u>A205549</u>	<u>002</u>	Dec 08, 2015
<u>AB</u>	TRIS PHARMA INC	<u>8MEQ</u>	<u>A201944</u>	<u>001</u>	Mar 04, 2016
<u>AB</u>		<u>10MEQ</u>	<u>A201944</u>	<u>002</u>	Mar 04, 2016
<u>AB</u>	ZYDUS PHARMS	<u>8MEQ</u>	<u>A208445</u>	<u>001</u>	Mar 11, 2019
<u>AB</u>		<u>10MEQ</u>	<u>A208445</u>	<u>002</u>	Mar 11, 2019

FOR SOLUTION; ORAL

KLOR-CON

<u>AA</u>	UPSHER SMITH LABS	<u>20MEQ</u>	<u>A209662</u>	<u>001</u>	Oct 23, 2017
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POTASSIUM CHLORIDE

<u>AA</u>	AMNEAL	<u>20MEQ</u>	<u>A210902</u>	<u>001</u>	May 23, 2019
<u>AA</u>	BELCHER	<u>20MEQ</u>	<u>A212183</u>	<u>001</u>	May 06, 2019
<u>AA</u>	EPIC PHARMA LLC	<u>20MEQ</u>	<u>A210200</u>	<u>001</u>	Nov 23, 2018
<u>AA</u>	NOVEL LABS INC	<u>20MEQ</u>	<u>A210241</u>	<u>001</u>	Nov 21, 2018
<u>AA</u>	+! PHARMA RES SOFTWARE	<u>20MEQ</u>	<u>N208019</u>	<u>001</u>	Aug 19, 2015

INJECTABLE; INJECTION

POTASSIUM CHLORIDE

<u>AP</u>	B BRAUN	<u>2MEQ/ML</u>	<u>A085870</u>	<u>001</u>	
<u>AP</u>	+ FRESENIUS KABI USA	<u>2MEQ/ML</u>	<u>A080225</u>	<u>001</u>	
<u>AP</u>	! HOSPIRA	<u>2MEQ/ML</u>	<u>A080205</u>	<u>001</u>	

POTASSIUM CHLORIDE 10MEQ

<u>AP</u>	NEXUS PHARMS	<u>14.9MG/ML</u>	<u>A214727</u>	<u>001</u>	Mar 18, 2021
<u>AP</u>		<u>745MG/100ML</u>	<u>A214727</u>	<u>002</u>	Mar 18, 2021

POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER

<u>AP</u>	+! BAXTER HLTHCARE	<u>14.9MG/ML</u>	<u>N019904</u>	<u>001</u>	Dec 26, 1989
<u>AP</u>	+!	<u>746MG/100ML</u>	<u>N019904</u>	<u>005</u>	Dec 17, 1990
<u>AP</u>	+! ICU MEDICAL INC	<u>14.9MG/ML</u>	<u>N020161</u>	<u>005</u>	Nov 30, 1992
<u>AP</u>	+!	<u>745MG/100ML</u>	<u>N020161</u>	<u>001</u>	Nov 30, 1992

POTASSIUM CHLORIDE 20MEQ

<u>AP</u>	NEXUS PHARMS	<u>29.8MG/ML</u>	<u>A214727</u>	<u>003</u>	Mar 18, 2021
<u>AP</u>		<u>1.49GM/100ML</u>	<u>A214727</u>	<u>004</u>	Mar 18, 2021

POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER

<u>AP</u>	+! BAXTER HLTHCARE	<u>29.8MG/ML</u>	<u>N019904</u>	<u>002</u>	Dec 26, 1989
<u>AP</u>	+!	<u>1.49GM/100ML</u>	<u>N019904</u>	<u>006</u>	Dec 17, 1990
<u>AP</u>	+! ICU MEDICAL INC	<u>29.8MG/ML</u>	<u>N020161</u>	<u>006</u>	Aug 11, 1998
<u>AP</u>	+!	<u>1.49GM/100ML</u>	<u>N020161</u>	<u>002</u>	Nov 30, 1992

POTASSIUM CHLORIDE 40MEQ

<u>AP</u>	NEXUS PHARMS	<u>2.98GM/100ML</u>	<u>A214727</u>	<u>005</u>	Mar 18, 2021
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PRESCRIPTION DRUG PRODUCT LIST

POTASSIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER

<u>AP</u>	<u>+</u>	<u>BAXTER HLTHCARE</u>	<u>2.98GM/100ML</u>	<u>N019904 004</u>	Dec 26, 1989
<u>AP</u>	<u>+</u>	<u>ICU MEDICAL INC</u>	<u>2.98GM/100ML</u>	<u>N020161 004</u>	Aug 11, 1998

POTASSIUM CHLORIDE IN PLASTIC CONTAINER

<u>AP</u>		<u>FRESENIUS KABI USA</u>	<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>		<u>AMNEAL</u>	<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>		<u>APOTEX</u>	<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>	<u>+</u>	<u>GENUS LIFESCIENCES</u>	<u>20MEQ/15ML</u>	<u>N206814 001</u>	Dec 22, 2014
<u>AA</u>	<u>+</u>		<u>40MEQ/15ML</u>	<u>N206814 002</u>	Dec 22, 2014
<u>AA</u>		<u>GRANULES</u>	<u>20MEQ/15ML</u>	<u>A213392 001</u>	Jan 29, 2021
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A213392 002</u>	Jan 29, 2021
<u>AA</u>		<u>NOVEL LABS INC</u>	<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
<u>AA</u>		<u>PHARM ASSOC</u>	<u>20MEQ/15ML</u>	<u>A210766 001</u>	Mar 29, 2019
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A210766 002</u>	Mar 29, 2019
<u>AA</u>		<u>SAPTALIS PHARMS</u>	<u>20MEQ/15ML</u>	<u>A211648 001</u>	May 21, 2021
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A211648 002</u>	May 21, 2021

TABLET, EXTENDED RELEASE; ORAL

KLOR-CON M10

<u>AB1</u>		<u>UPSHER SMITH LABS</u>	<u>10MEQ</u>	<u>A074726 002</u>	Aug 09, 2000
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KLOR-CON M15

<u>AB1</u>		<u>UPSHER SMITH LABS</u>	<u>15MEQ</u>	<u>A074726 003</u>	Jun 06, 2003
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KLOR-CON M20

<u>AB1</u>	<u>!</u>	<u>UPSHER SMITH LABS</u>	<u>20MEQ</u>	<u>A074726 001</u>	Nov 20, 1998
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POTASSIUM CHLORIDE

<u>AB1</u>		<u>ACTAVIS LABS FL INC</u>	<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>		<u>ADARE PHARMS INC</u>	<u>10MEQ</u>	<u>A076368 002</u>	Jun 05, 2019
<u>AB1</u>			<u>20MEQ</u>	<u>A076368 001</u>	Aug 18, 2004
<u>AB1</u>		<u>AMNEAL</u>	<u>10MEQ</u>	<u>A212861 001</u>	May 08, 2020
<u>AB1</u>			<u>20MEQ</u>	<u>A212861 003</u>	May 08, 2020
<u>AB1</u>		<u>ASCENT PHARMS INC</u>	<u>10MEQ</u>	<u>A214422 001</u>	Dec 29, 2020
<u>AB1</u>			<u>15MEQ</u>	<u>A214422 002</u>	Dec 29, 2020
<u>AB1</u>			<u>20MEQ</u>	<u>A214422 003</u>	Dec 29, 2020
<u>AB1</u>		<u>GLENMARK PHARMS LTD</u>	<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>		<u>GRANULES</u>	<u>10MEQ</u>	<u>A214452 001</u>	Oct 21, 2020
<u>AB1</u>			<u>20MEQ</u>	<u>A214452 002</u>	Oct 21, 2020
<u>AB1</u>		<u>NOVEL LABS INC</u>	<u>10MEQ</u>	<u>A206347 001</u>	Jan 21, 2016
<u>AB1</u>			<u>20MEQ</u>	<u>A206347 002</u>	Jan 21, 2016
<u>AB1</u>		<u>PRINSTON INC</u>	<u>10MEQ</u>	<u>A209922 001</u>	Apr 30, 2019
<u>AB1</u>			<u>15MEQ</u>	<u>A209922 002</u>	Apr 30, 2019
<u>AB1</u>			<u>20MEQ</u>	<u>A209922 003</u>	Apr 30, 2019
<u>AB1</u>		<u>ZYDUS PHARMS</u>	<u>10MEQ</u>	<u>A210395 001</u>	Sep 17, 2020
<u>AB1</u>			<u>20MEQ</u>	<u>A210395 002</u>	Sep 17, 2020

KLOR-CON

<u>AB2</u>	<u>+</u>	<u>UPSHER SMITH LABS</u>	<u>8MEQ</u>	<u>N019123 001</u>	Apr 17, 1986
<u>AB2</u>	<u>+</u>		<u>10MEQ</u>	<u>N019123 002</u>	Apr 17, 1986

POTASSIUM CHLORIDE

<u>AB2</u>		<u>AUROBINDO PHARMA LTD</u>	<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>		<u>GRANULES</u>	<u>8MEQ</u>	<u>A211797 001</u>	Mar 04, 2020
<u>AB2</u>			<u>10MEQ</u>	<u>A211797 002</u>	Mar 04, 2020
<u>AB2</u>		<u>MYLAN</u>	<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>		<u>NOVEL LABS INC</u>	<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>		<u>PADAGIS US</u>	<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>		<u>STRIDES PHARMA</u>	<u>8MEQ</u>	<u>A210733 001</u>	Aug 31, 2018
<u>AB2</u>			<u>10MEQ</u>	<u>A210733 002</u>	Aug 31, 2018

PRESCRIPTION DRUG PRODUCT LIST

POTASSIUM CHLORIDE

TABLET, EXTENDED RELEASE;ORAL

POTASSIUM CHLORIDE

<u>AB2</u>	YICHANG HUMANWELL	<u>8MEQ</u>	<u>A209314</u>	<u>001</u>	Jun 22, 2018
<u>AB2</u>		<u>10MEQ</u>	<u>A209314</u>	<u>002</u>	Jun 22, 2018

K-TAB

<u>AB3</u>	+ ABBVIE	<u>10MEQ</u>	<u>N018279</u>	<u>001</u>	
<u>AB3</u>	+!	<u>20MEQ</u>	<u>N018279</u>	<u>003</u>	Nov 25, 2013

POTASSIUM CHLORIDE

<u>AB3</u>	VITRUVIAS THERAP	<u>10MEQ</u>	<u>A209688</u>	<u>001</u>	Jan 12, 2018
<u>AB3</u>		<u>20MEQ</u>	<u>A209688</u>	<u>002</u>	Jan 12, 2018
<u>AB3</u>	YICHANG HUMANWELL	<u>10MEQ</u>	<u>A212561</u>	<u>001</u>	Sep 30, 2019
<u>AB3</u>		<u>20MEQ</u>	<u>A212561</u>	<u>002</u>	Sep 30, 2019

K-TAB

BC	+ ABBVIE	8MEQ	N018279	002	Aug 01, 1988
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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</u>	<u>001</u>	Sep 10, 2008
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POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9%

<u>AP</u>	+! BAXTER HLTHCARE	<u>150MG/100ML;900MG/100ML</u>	<u>N017648</u>	<u>001</u>	
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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</u>	<u>004</u>	Sep 29, 1989
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POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	ICU MEDICAL INC	<u>149MG/100ML;900MG/100ML</u>	<u>N019686</u>	<u>001</u>	Oct 17, 1988
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POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	ICU MEDICAL INC	<u>298MG/100ML;900MG/100ML</u>	<u>N019686</u>	<u>002</u>	Oct 17, 1988
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POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45%

+!	BAXTER HLTHCARE	150MG/100ML;450MG/100ML	N017648	005	Nov 26, 2002
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POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9%

+!	BAXTER HLTHCARE	300MG/100ML;900MG/100ML	N017648	002	
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POTASSIUM CITRATE

TABLET, EXTENDED RELEASE;ORAL

POTASSIUM CITRATE

<u>AB</u>	ANI PHARMS	<u>10MEQ</u>	<u>A212779</u>	<u>001</u>	Jan 14, 2020
<u>AB</u>		<u>15MEQ</u>	<u>A212779</u>	<u>002</u>	Jan 14, 2020
<u>AB</u>	ASCENT PHARMS INC	<u>5MEQ</u>	<u>A214420</u>	<u>001</u>	Feb 05, 2021
<u>AB</u>		<u>10MEQ</u>	<u>A214420</u>	<u>002</u>	Feb 05, 2021
<u>AB</u>		<u>15MEQ</u>	<u>A214420</u>	<u>003</u>	Feb 05, 2021
<u>AB</u>	BIONPHARMA INC	<u>10MEQ</u>	<u>A212799</u>	<u>001</u>	Jun 29, 2020
<u>AB</u>		<u>15MEQ</u>	<u>A212799</u>	<u>002</u>	Jun 29, 2020
<u>AB</u>	EYWA	<u>5MEQ</u>	<u>A214426</u>	<u>001</u>	Feb 19, 2021
<u>AB</u>		<u>10MEQ</u>	<u>A214426</u>	<u>002</u>	Feb 19, 2021
<u>AB</u>		<u>15MEQ</u>	<u>A214426</u>	<u>003</u>	Feb 19, 2021
<u>AB</u>	RISING	<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	<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

UROCIIT-K

<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

POTASSIUM PHOSPHATE, DIBASIC; POTASSIUM PHOSPHATE, MONOBASIC

SOLUTION;INTRAVENOUS

POTASSIUM PHOSPHATES

+!	CMP DEV LLC	4.5GM/15ML (300MG/ML);2.65GM/15ML (175MG/ML)	N212121	001	Sep 19, 2019
+!	FRESENIUS KABI USA	1.18GM/5ML (236MG/ML);1.12GM/5ML (224MG/ML)	N212832	001	Nov 26, 2019
+!		3.54GM/15ML (236MG/ML);3.36GM/15ML (224MG/ML)	N212832	002	Nov 26, 2019
+!		11.8GM/50ML (236MG/ML);11.2GM/50ML (224MG/ML)	N212832	003	Nov 26, 2019

PRESCRIPTION DRUG PRODUCT LIST

POVIDONE-IODINE

SOLUTION/DROPS;OPHTHALMIC

BETADINE

+! ALCON PHARMS LTD 5%

N018634 001 Dec 17, 1986

PRALATREXATE

SOLUTION;INTRAVENOUS

FOLOTYN

+ ACROTECH 20MG/ML (20MG/ML)

N022468 001 Sep 24, 2009

+! 40MG/2ML (20MG/ML)

N022468 002 Sep 24, 2009

PRALIDOXIME CHLORIDE

INJECTABLE;INJECTION

PROTOPAM CHLORIDE

+! BAXTER HLTHCARE 1GM/VIAL

N014134 001

CORP

PRALSETINIB

CAPSULE;ORAL

GAVRETO

+! GENENTECH INC 100MG

N213721 001 Sep 04, 2020

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET;ORAL

MIRAPEXAB + BOEHRINGER
INGELHEIM0.125MGN020667 001 Jul 01, 1997AB +!0.25MGN020667 002 Jul 01, 1997AB +0.5MGN020667 006 Feb 12, 1998AB +0.75MGN020667 007 Jul 30, 2007AB +1MGN020667 003 Jul 01, 1997AB +1.5MGN020667 005 Jul 01, 1997PRAMIPEXOLE DIHYDROCHLORIDEAB AUROBINDO PHARMA
LTD0.125MGA202633 001 Oct 26, 2012AB0.25MGA202633 002 Oct 26, 2012AB0.5MGA202633 003 Oct 26, 2012AB0.75MGA202633 004 Oct 26, 2012AB1MGA202633 005 Oct 26, 2012AB1.5MGA202633 006 Oct 26, 2012AB GLENMARK GENERICS0.125MGA090781 001 Oct 08, 2010AB0.25MGA090781 002 Oct 08, 2010AB0.5MGA090781 003 Oct 08, 2010AB0.75MGA090781 006 Sep 11, 2015AB1MGA090781 004 Oct 08, 2010AB1.5MGA090781 005 Oct 08, 2010AB GRAVITI PHARMS0.25MGA211088 001 Oct 03, 2018AB0.5MGA211088 002 Oct 03, 2018AB0.75MGA211088 003 Oct 03, 2018AB1MGA211088 004 Oct 03, 2018AB1.5MGA211088 005 Oct 03, 2018AB SCIEGEN PHARMS INC0.125MGA203855 001 Oct 28, 2014AB0.25MGA203855 002 Oct 28, 2014AB0.5MGA203855 003 Oct 28, 2014AB0.75MGA203855 004 Oct 28, 2014AB1MGA203855 005 Oct 28, 2014AB1.5MGA203855 006 Oct 28, 2014AB STRIDES PHARMA0.125MGA202702 001 Jun 03, 2014AB0.25MGA202702 002 Jun 03, 2014AB0.5MGA202702 003 Jun 03, 2014AB0.75MGA202702 004 Jun 03, 2014AB1MGA202702 005 Jun 03, 2014AB1.5MGA202702 006 Jun 03, 2014AB TORRENT PHARMS0.125MGA090865 001 Oct 08, 2010AB0.25MGA090865 002 Oct 08, 2010AB0.5MGA090865 003 Oct 08, 2010AB0.75MGA090865 004 Oct 08, 2010AB1MGA090865 005 Oct 08, 2010AB1.5MGA090865 006 Oct 08, 2010AB ZENNOVA0.125MGA090151 001 Apr 30, 2012AB0.25MGA090151 002 Apr 30, 2012AB0.5MGA090151 003 Apr 30, 2012AB0.75MGA090151 006 Apr 30, 2012AB1MGA090151 004 Apr 30, 2012AB1.5MGA090151 005 Apr 30, 2012

PRESCRIPTION DRUG PRODUCT LIST

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET;ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

<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>	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	<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>	XIAMEN LP PHARM CO	<u>0.375MG</u>	<u>A212797 001</u>	Jun 11, 2021
<u>AB</u>		<u>0.75MG</u>	<u>A212797 002</u>	Jun 11, 2021
<u>AB</u>	ZYDUS PHARMS	<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

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

+	MILLICENT	6.5MG	N208470 001	Nov 16, 2016
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PRESCRIPTION DRUG PRODUCT LIST

PRASUGREL HYDROCHLORIDE

TABLET;ORAL

EFFIENT

<u>AB</u>	+	DAIICHI SANKYO INC	<u>EQ 5MG BASE</u>	<u>N022307 001</u>	Jul 10, 2009
<u>AB</u>	+	!	<u>EQ 10MG BASE</u>	<u>N022307 002</u>	Jul 10, 2009

PRASUGREL

<u>AB</u>		ACCORD HLTHCARE	<u>EQ 5MG BASE</u>	<u>A205987 001</u>	Feb 02, 2018
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A205987 002</u>	Feb 02, 2018
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A205888 001</u>	Oct 16, 2017
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A205888 002</u>	Oct 16, 2017
<u>AB</u>		HEC PHARM	<u>EQ 5MG BASE</u>	<u>A206021 001</u>	Jan 16, 2019
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A206021 002</u>	Jan 16, 2019
<u>AB</u>		MYLAN	<u>EQ 5MG BASE</u>	<u>A205927 001</u>	Jul 12, 2017
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A205927 002</u>	Jul 12, 2017
<u>AB</u>		PANACEA	<u>EQ 5MG BASE</u>	<u>A205897 001</u>	Oct 16, 2017
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A205897 002</u>	Oct 16, 2017
<u>AB</u>		USPHARMA WINDLAS	<u>EQ 5MG BASE</u>	<u>A205790 001</u>	Oct 16, 2017
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A205790 002</u>	Oct 16, 2017
BX		AMNEAL PHARMS	EQ 5MG BASE	A205913 001	Jun 19, 2018
BX			EQ 10MG BASE	A205913 002	Jun 19, 2018

PRAVASTATIN SODIUM

TABLET;ORAL

PRAVASTATIN SODIUM

<u>AB</u>		ACCORD HLTHCARE	<u>10MG</u>	<u>A207068 001</u>	Nov 17, 2016
<u>AB</u>			<u>20MG</u>	<u>A207068 002</u>	Nov 17, 2016
<u>AB</u>			<u>40MG</u>	<u>A207068 003</u>	Nov 17, 2016
<u>AB</u>			<u>80MG</u>	<u>A207068 004</u>	Nov 17, 2016
<u>AB</u>		APNAR PHARMA LP	<u>10MG</u>	<u>A077491 002</u>	Oct 23, 2006
<u>AB</u>			<u>20MG</u>	<u>A077491 003</u>	Oct 23, 2006
<u>AB</u>			<u>40MG</u>	<u>A077491 004</u>	Oct 23, 2006
<u>AB</u>			<u>80MG</u>	<u>A077491 001</u>	Feb 11, 2008
<u>AB</u>		APOTEX	<u>10MG</u>	<u>A076341 001</u>	Oct 23, 2006
<u>AB</u>			<u>20MG</u>	<u>A076341 002</u>	Oct 23, 2006
<u>AB</u>			<u>40MG</u>	<u>A076341 003</u>	Oct 23, 2006
<u>AB</u>			<u>80MG</u>	<u>A076341 004</u>	Dec 28, 2007
<u>AB</u>		AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A203367 001</u>	Feb 02, 2017
<u>AB</u>			<u>20MG</u>	<u>A203367 002</u>	Feb 02, 2017
<u>AB</u>			<u>40MG</u>	<u>A203367 003</u>	Feb 02, 2017
<u>AB</u>			<u>80MG</u>	<u>A203367 004</u>	Feb 02, 2017
<u>AB</u>		BIOCON PHARMA	<u>10MG</u>	<u>A209869 001</u>	Apr 13, 2018
<u>AB</u>			<u>20MG</u>	<u>A209869 002</u>	Apr 13, 2018
<u>AB</u>			<u>40MG</u>	<u>A209869 003</u>	Apr 13, 2018
<u>AB</u>			<u>80MG</u>	<u>A209869 004</u>	Apr 13, 2018
<u>AB</u>		CIPLA	<u>10MG</u>	<u>A077904 001</u>	Oct 23, 2006
<u>AB</u>			<u>20MG</u>	<u>A077904 002</u>	Oct 23, 2006
<u>AB</u>			<u>40MG</u>	<u>A077904 003</u>	Oct 23, 2006
<u>AB</u>			<u>80MG</u>	<u>A077904 004</u>	Mar 22, 2016
<u>AB</u>		DR REDDYS LABS INC	<u>10MG</u>	<u>A076714 001</u>	Oct 23, 2006
<u>AB</u>			<u>20MG</u>	<u>A076714 002</u>	Oct 23, 2006
<u>AB</u>			<u>40MG</u>	<u>A076714 003</u>	Oct 23, 2006
<u>AB</u>			<u>80MG</u>	<u>A076714 004</u>	Dec 28, 2007
<u>AB</u>		GLENMARK GENERICS	<u>10MG</u>	<u>A077987 001</u>	May 11, 2007
<u>AB</u>			<u>20MG</u>	<u>A077987 002</u>	May 11, 2007
<u>AB</u>			<u>40MG</u>	<u>A077987 003</u>	May 11, 2007
<u>AB</u>			<u>80MG</u>	<u>A077987 004</u>	Dec 28, 2007
<u>AB</u>		LUPIN PHARMS	<u>10MG</u>	<u>A077917 001</u>	Jan 08, 2008
<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>		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>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

PRESCRIPTION DRUG PRODUCT LIST

PRAZIOUANTEL

TABLET;ORAL

BILTRICIDE

AB	+	BAYER HLTHCARE	600MG	<u>N018714</u>	<u>001</u>	Dec 29, 1982
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PRAZIOUANTEL

AB		PAR PHARM INC	600MG	<u>A208820</u>	<u>001</u>	Nov 27, 2017
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PRAZOSIN HYDROCHLORIDE

CAPSULE;ORAL

MINIPRESS

AB	+	PFIZER	EQ 1MG BASE	<u>N017442</u>	<u>002</u>	
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AB	+		EQ 2MG BASE	<u>N017442</u>	<u>003</u>	
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AB	+		EQ 5MG BASE	<u>N017442</u>	<u>001</u>	
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PRAZOSIN HYDROCHLORIDE

AB		ANI PHARMS	EQ 1MG BASE	<u>A072577</u>	<u>002</u>	May 16, 1989
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AB			EQ 2MG BASE	<u>A072577</u>	<u>001</u>	May 16, 1989
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AB			EQ 5MG BASE	<u>A072577</u>	<u>003</u>	May 16, 1989
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AB		GRANULES	EQ 1MG BASE	<u>A214608</u>	<u>001</u>	Dec 23, 2021
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AB			EQ 2MG BASE	<u>A214608</u>	<u>002</u>	Dec 23, 2021
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AB			EQ 5MG BASE	<u>A214608</u>	<u>003</u>	Dec 23, 2021
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AB		MYLAN	EQ 1MG BASE	<u>A072575</u>	<u>003</u>	May 16, 1989
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AB			EQ 2MG BASE	<u>A072575</u>	<u>002</u>	May 16, 1989
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AB			EQ 5MG BASE	<u>A072575</u>	<u>001</u>	May 16, 1989
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AB		NOVITIUM PHARMA	EQ 1MG BASE	<u>A210971</u>	<u>001</u>	Oct 03, 2018
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AB			EQ 2MG BASE	<u>A210971</u>	<u>002</u>	Oct 03, 2018
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AB			EQ 5MG BASE	<u>A210971</u>	<u>003</u>	Oct 03, 2018
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AB		TEVA PHARMS	EQ 1MG BASE	<u>A071745</u>	<u>002</u>	Sep 12, 1988
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AB			EQ 2MG BASE	<u>A071745</u>	<u>003</u>	Sep 12, 1988
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AB			EQ 5MG BASE	<u>A071745</u>	<u>001</u>	Sep 12, 1988
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PREDNICARBATE

OINTMENT;TOPICAL

PREDNICARBATE

!	FOUGERA PHARMS	0.1%	<u>A077236</u>	<u>001</u>	Mar 09, 2007
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PREDNISOLONE

SYRUP;ORAL

PREDNISOLONE

AA	!	AKORN	15MG/5ML	<u>A040401</u>	<u>001</u>	Feb 27, 2003
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AA		CHARTWELL RX	15MG/5ML	<u>A040323</u>	<u>001</u>	May 13, 1999
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AA		LANNETT CO INC	15MG/5ML	<u>A040775</u>	<u>001</u>	Sep 21, 2007
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AA		PHARM ASSOC	15MG/5ML	<u>A040399</u>	<u>001</u>	Mar 05, 2003
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AA		WOCKHARDT BIO AG	15MG/5ML	<u>A040313</u>	<u>001</u>	Sep 10, 2003
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PRELONE

AA		TEVA	15MG/5ML	<u>A089081</u>	<u>001</u>	Feb 04, 1986
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TABLET;ORAL

PREDNISOLONE

!	WATSON LABS	5MG	<u>A080354</u>	<u>001</u>	
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PREDNISOLONE ACETATE

SUSPENSION/DROPS;OPHTHALMIC

OMNIPRED

AB	+	NOVARTIS	1%	<u>N017469</u>	<u>001</u>	
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PRED FORTE

AB	+	ALLERGAN	1%	<u>N017011</u>	<u>001</u>	
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PRED MILD

+	ALLERGAN	0.12%	<u>N017100</u>	<u>001</u>	
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PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT;OPHTHALMIC

BLEPHAMIDE S.O.P.

!	ALLERGAN	0.2%;10%	<u>A087748</u>	<u>001</u>	Dec 03, 1986
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SUSPENSION;OPHTHALMIC

BLEPHAMIDE

+	ALLERGAN	0.2%;10%	<u>N012813</u>	<u>002</u>	
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PREDNISOLONE SODIUM PHOSPHATE

SOLUTION;ORAL

PEDIAPRED

AA	+	SETON PHARM	EQ 5MG BASE/5ML	<u>N019157</u>	<u>001</u>	May 28, 1986
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PREDNISOLONE SODIUM PHOSPHATE

AA		AKORN	EQ 5MG BASE/5ML	<u>A075183</u>	<u>001</u>	Mar 26, 2003
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AA		CHARTWELL RX	EQ 5MG BASE/5ML	<u>A075988</u>	<u>001</u>	May 25, 2004
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AA		EDENBRIDGE PHARMS	EQ 10MG BASE/5ML	<u>A203559</u>	<u>001</u>	Dec 20, 2016
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AA			EQ 20MG BASE/5ML	<u>A203559</u>	<u>002</u>	Dec 20, 2016
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PRESCRIPTION DRUG PRODUCT LIST

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION;ORAL

PREDNISOLONE SODIUM PHOSPHATE

<u>AA</u>	!	PHARM ASSOC	<u>EQ 10MG BASE/5ML</u>	<u>A078465 001</u>	Mar 07, 2008
<u>AA</u>			<u>EQ 15MG BASE/5ML</u>	<u>A076913 001</u>	Apr 25, 2005
<u>AA</u>	!		<u>EQ 20MG BASE/5ML</u>	<u>A078988 001</u>	Jun 09, 2008
<u>AA</u>		WOCKHARDT BIO AG	<u>EQ 5MG BASE/5ML</u>	<u>A075099 001</u>	Jun 28, 2002
<u>AA</u>	!		<u>EQ 15MG BASE/5ML</u>	<u>A076895 001</u>	Oct 04, 2004
	!	MISSION PHARMA	EQ 25MG BASE/5ML	A091396 001	Sep 13, 2010
		PHARM ASSOC	EQ 30MG BASE/5ML	A204962 001	Mar 11, 2020

SOLUTION/DROPS;OPHTHALMIC

PREDNISOLONE SODIUM PHOSPHATE

! BAUSCH AND LOMB EQ 0.9% PHOSPHATE A040070 001 Jul 29, 1994

TABLET, ORALLY DISINTEGRATING;ORAL

ORAPRED ODT

+ CONCORDIA PHARMS EQ 10MG BASE N021959 001 Jun 01, 2006
INC

+ EQ 15MG BASE N021959 002 Jun 01, 2006

+! EQ 30MG BASE N021959 003 Jun 01, 2006

PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC

SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE

! BAUSCH AND LOMB EQ 0.23% PHOSPHATE;10% A074449 001 Dec 29, 1995

PREDNISON

SOLUTION;ORAL

PREDNISON

! HIKMA 5MG/5ML A088703 001 Nov 08, 1984

PREDNISON INTENSOL

! HIKMA 5MG/ML A088810 001 Feb 20, 1985

TABLET;ORAL

PREDNISON

<u>AB</u>		AMNEAL	<u>1MG</u>	<u>A213385 001</u>	Jun 16, 2020
<u>AB</u>			<u>5MG</u>	<u>A213385 002</u>	Jun 16, 2020
<u>AB</u>			<u>10MG</u>	<u>A213386 001</u>	Jun 24, 2020
<u>AB</u>			<u>20MG</u>	<u>A213386 002</u>	Jun 24, 2020
<u>AB</u>		AUROBINDO PHARMA LTD	<u>1MG</u>	<u>A215671 001</u>	Nov 16, 2021
<u>AB</u>		GENEYORK PHARMS	<u>1MG</u>	<u>A211496 001</u>	Dec 28, 2018
<u>AB</u>			<u>2.5MG</u>	<u>A211495 001</u>	Dec 07, 2018
<u>AB</u>			<u>5MG</u>	<u>A211495 002</u>	Dec 07, 2018
<u>AB</u>			<u>10MG</u>	<u>A210525 001</u>	Dec 04, 2018
<u>AB</u>			<u>20MG</u>	<u>A210525 002</u>	Dec 04, 2018
<u>AB</u>			<u>50MG</u>	<u>A210525 003</u>	Dec 04, 2018
<u>AB</u>	+!	HIKMA	<u>1MG</u>	<u>A087800 001</u>	Apr 22, 1982
<u>AB</u>	!		<u>2.5MG</u>	<u>A087801 001</u>	Apr 22, 1982
<u>AB</u>	!		<u>5MG</u>	<u>A080352 001</u>	
<u>AB</u>	+!		<u>10MG</u>	<u>A084122 001</u>	
<u>AB</u>	+!		<u>20MG</u>	<u>A087342 001</u>	
<u>AB</u>	!		<u>50MG</u>	<u>A084283 001</u>	
<u>AB</u>		JUBILANT CADISTA	<u>1MG</u>	<u>A040611 001</u>	Jun 06, 2005
<u>AB</u>			<u>5MG</u>	<u>A040362 002</u>	Aug 29, 2001
<u>AB</u>			<u>10MG</u>	<u>A040362 001</u>	Aug 29, 2001
<u>AB</u>			<u>20MG</u>	<u>A040362 003</u>	Jun 29, 2005
<u>AB</u>		MYLAN	<u>5MG</u>	<u>A080292 001</u>	
<u>AB</u>			<u>10MG</u>	<u>A088832 001</u>	Dec 04, 1985
<u>AB</u>			<u>20MG</u>	<u>A083677 001</u>	
<u>AB</u>		NOVITIUM PHARMA	<u>1MG</u>	<u>A215246 001</u>	Jul 06, 2021
<u>AB</u>			<u>2.5MG</u>	<u>A211575 001</u>	Nov 15, 2019
<u>AB</u>			<u>5MG</u>	<u>A211575 002</u>	Nov 15, 2019
<u>AB</u>			<u>10MG</u>	<u>A211575 003</u>	Nov 15, 2019
<u>AB</u>			<u>20MG</u>	<u>A211575 004</u>	Nov 15, 2019
<u>AB</u>			<u>50MG</u>	<u>A211575 005</u>	Nov 15, 2019
<u>AB</u>		STRIDES PHARMA	<u>1MG</u>	<u>A210785 001</u>	Sep 02, 2020
<u>AB</u>			<u>2.5MG</u>	<u>A209727 001</u>	Nov 20, 2020
<u>AB</u>			<u>5MG</u>	<u>A209727 002</u>	Nov 20, 2020
<u>AB</u>			<u>10MG</u>	<u>A208412 001</u>	Feb 11, 2021
<u>AB</u>			<u>20MG</u>	<u>A208412 002</u>	Feb 11, 2021
<u>AB</u>			<u>50MG</u>	<u>A208412 003</u>	Jan 11, 2022
<u>AB</u>		SUN PHARM INDUSTRIES	<u>5MG</u>	<u>A089247 002</u>	Dec 04, 1985
<u>AB</u>			<u>10MG</u>	<u>A089247 003</u>	Dec 04, 1985
<u>AB</u>			<u>20MG</u>	<u>A089247 001</u>	Dec 04, 1985

PRESCRIPTION DRUG PRODUCT LIST

PREDNISON

TABLET; ORAL

PREDNISON

<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>	

TABLET, DELAYED RELEASE; ORAL

RAYOS

+	HORIZON	1MG	N202020	001	Jul 26, 2012
+		2MG	N202020	002	Jul 26, 2012
+	!	5MG	N202020	003	Jul 26, 2012

PREGABALIN

CAPSULE; ORAL

LYRICA

<u>AB</u>	+	UPJOHN	<u>25MG</u>	<u>N021446</u>	<u>001</u>	Dec 30, 2004
<u>AB</u>	+		<u>50MG</u>	<u>N021446</u>	<u>002</u>	Dec 30, 2004
<u>AB</u>	+		<u>75MG</u>	<u>N021446</u>	<u>003</u>	Dec 30, 2004
<u>AB</u>	+		<u>100MG</u>	<u>N021446</u>	<u>004</u>	Dec 30, 2004
<u>AB</u>	+		<u>150MG</u>	<u>N021446</u>	<u>005</u>	Dec 30, 2004
<u>AB</u>	+		<u>200MG</u>	<u>N021446</u>	<u>006</u>	Dec 30, 2004
<u>AB</u>	+		<u>225MG</u>	<u>N021446</u>	<u>007</u>	Dec 30, 2004
<u>AB</u>	+	!	<u>300MG</u>	<u>N021446</u>	<u>008</u>	Dec 30, 2004

PREGABALIN

<u>AB</u>	ACTAVIS ELIZABETH	<u>25MG</u>	<u>A091025</u>	<u>001</u>	Jul 09, 2020
<u>AB</u>		<u>50MG</u>	<u>A091025</u>	<u>002</u>	Jul 09, 2020
<u>AB</u>		<u>75MG</u>	<u>A091025</u>	<u>003</u>	Jul 09, 2020
<u>AB</u>		<u>100MG</u>	<u>A091025</u>	<u>004</u>	Jul 09, 2020
<u>AB</u>		<u>150MG</u>	<u>A091025</u>	<u>005</u>	Jul 09, 2020
<u>AB</u>		<u>200MG</u>	<u>A091025</u>	<u>006</u>	Jul 09, 2020
<u>AB</u>		<u>225MG</u>	<u>A091025</u>	<u>007</u>	Jul 09, 2020
<u>AB</u>		<u>300MG</u>	<u>A091025</u>	<u>008</u>	Jul 09, 2020
<u>AB</u>	ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A203459</u>	<u>001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A203459</u>	<u>002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A203459</u>	<u>003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A203459</u>	<u>004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A203459</u>	<u>005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A203459</u>	<u>006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A203459</u>	<u>007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A203459</u>	<u>008</u>	Jul 19, 2019
<u>AB</u>	ALKEM LABS LTD	<u>25MG</u>	<u>A207799</u>	<u>007</u>	Sep 30, 2019
<u>AB</u>		<u>50MG</u>	<u>A207799</u>	<u>008</u>	Sep 30, 2019
<u>AB</u>		<u>75MG</u>	<u>A207799</u>	<u>001</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A207799</u>	<u>002</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A207799</u>	<u>003</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A207799</u>	<u>004</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A207799</u>	<u>005</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A207799</u>	<u>006</u>	Jul 19, 2019
<u>AB</u>	AMNEAL PHARMS CO	<u>25MG</u>	<u>A209743</u>	<u>001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A209743</u>	<u>002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A209743</u>	<u>003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A209743</u>	<u>004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A209743</u>	<u>005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A209743</u>	<u>006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A209743</u>	<u>007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A209743</u>	<u>008</u>	Jul 19, 2019
<u>AB</u>	APOTEX	<u>25MG</u>	<u>A211685</u>	<u>001</u>	Jul 07, 2021
<u>AB</u>		<u>50MG</u>	<u>A211685</u>	<u>002</u>	Jul 07, 2021
<u>AB</u>		<u>75MG</u>	<u>A211685</u>	<u>003</u>	Jul 07, 2021
<u>AB</u>		<u>100MG</u>	<u>A211685</u>	<u>004</u>	Jul 07, 2021
<u>AB</u>		<u>150MG</u>	<u>A211685</u>	<u>005</u>	Jul 07, 2021
<u>AB</u>		<u>200MG</u>	<u>A211685</u>	<u>006</u>	Jul 07, 2021
<u>AB</u>		<u>225MG</u>	<u>A211685</u>	<u>007</u>	Jul 07, 2021
<u>AB</u>		<u>300MG</u>	<u>A211685</u>	<u>008</u>	Jul 07, 2021
<u>AB</u>	CELLTRION	<u>50MG</u>	<u>A212865</u>	<u>001</u>	Mar 20, 2020
<u>AB</u>		<u>75MG</u>	<u>A212865</u>	<u>002</u>	Mar 20, 2020
<u>AB</u>		<u>100MG</u>	<u>A212865</u>	<u>003</u>	Mar 20, 2020
<u>AB</u>		<u>150MG</u>	<u>A212865</u>	<u>004</u>	Mar 20, 2020

PRESCRIPTION DRUG PRODUCT LIST

PREGABALIN

CAPSULE; ORAL

PREGABALIN

<u>AB</u>	CHANGZHOU PHARM	<u>50MG</u>	<u>A214322 001</u>	Jul 15, 2021
<u>AB</u>		<u>75MG</u>	<u>A214322 002</u>	Jul 15, 2021
<u>AB</u>		<u>100MG</u>	<u>A214322 003</u>	Jul 15, 2021
<u>AB</u>		<u>150MG</u>	<u>A214322 004</u>	Jul 15, 2021
<u>AB</u>		<u>200MG</u>	<u>A214322 005</u>	Jul 15, 2021
<u>AB</u>		<u>225MG</u>	<u>A214322 006</u>	Jul 15, 2021
<u>AB</u>		<u>300MG</u>	<u>A214322 007</u>	Jul 15, 2021
<u>AB</u>	CIPLA	<u>25MG</u>	<u>A212280 001</u>	Jan 10, 2020
<u>AB</u>		<u>50MG</u>	<u>A212280 002</u>	Jan 10, 2020
<u>AB</u>		<u>75MG</u>	<u>A212280 003</u>	Jan 10, 2020
<u>AB</u>		<u>100MG</u>	<u>A212280 004</u>	Jan 10, 2020
<u>AB</u>		<u>150MG</u>	<u>A212280 005</u>	Jan 10, 2020
<u>AB</u>		<u>200MG</u>	<u>A212280 006</u>	Jan 10, 2020
<u>AB</u>		<u>225MG</u>	<u>A212280 007</u>	Jan 10, 2020
<u>AB</u>		<u>300MG</u>	<u>A212280 008</u>	Jan 10, 2020
<u>AB</u>	CSPC OUYI	<u>50MG</u>	<u>A210585 001</u>	Dec 26, 2019
<u>AB</u>		<u>75MG</u>	<u>A210585 002</u>	Dec 26, 2019
<u>AB</u>		<u>100MG</u>	<u>A210585 003</u>	Dec 26, 2019
<u>AB</u>		<u>150MG</u>	<u>A210585 004</u>	Dec 26, 2019
<u>AB</u>		<u>200MG</u>	<u>A210585 005</u>	Dec 26, 2019
<u>AB</u>		<u>300MG</u>	<u>A210585 006</u>	Dec 26, 2019
<u>AB</u>	DR REDDYS	<u>25MG</u>	<u>A209664 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A209664 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A209664 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A209664 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A209664 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A209664 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A209664 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A209664 008</u>	Jul 19, 2019
<u>AB</u>	HETERO LABS LTD III	<u>25MG</u>	<u>A206912 001</u>	Oct 08, 2019
<u>AB</u>		<u>50MG</u>	<u>A206912 002</u>	Oct 08, 2019
<u>AB</u>		<u>75MG</u>	<u>A206912 003</u>	Oct 08, 2019
<u>AB</u>		<u>100MG</u>	<u>A206912 004</u>	Oct 08, 2019
<u>AB</u>		<u>150MG</u>	<u>A206912 005</u>	Oct 08, 2019
<u>AB</u>		<u>200MG</u>	<u>A206912 006</u>	Oct 08, 2019
<u>AB</u>		<u>225MG</u>	<u>A206912 007</u>	Oct 08, 2019
<u>AB</u>		<u>300MG</u>	<u>A206912 008</u>	Oct 08, 2019
<u>AB</u>	HIKAL	<u>25MG</u>	<u>A213423 001</u>	Mar 23, 2020
<u>AB</u>		<u>50MG</u>	<u>A213423 002</u>	Mar 23, 2020
<u>AB</u>		<u>75MG</u>	<u>A213423 003</u>	Mar 23, 2020
<u>AB</u>		<u>100MG</u>	<u>A213423 004</u>	Mar 23, 2020
<u>AB</u>		<u>150MG</u>	<u>A213423 005</u>	Mar 23, 2020
<u>AB</u>		<u>200MG</u>	<u>A213423 006</u>	Mar 23, 2020
<u>AB</u>		<u>225MG</u>	<u>A213423 007</u>	Mar 23, 2020
<u>AB</u>		<u>300MG</u>	<u>A213423 008</u>	Mar 23, 2020
<u>AB</u>	INVAGEN PHARMS	<u>25MG</u>	<u>A211384 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A211384 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A211384 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A211384 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A211384 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A211384 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A211384 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A211384 008</u>	Jul 19, 2019
<u>AB</u>	MSN	<u>25MG</u>	<u>A209357 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A209357 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A209357 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A209357 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A209357 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A209357 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A209357 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A209357 008</u>	Jul 19, 2019
<u>AB</u>	RISING	<u>25MG</u>	<u>A210432 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A210432 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A210432 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A210432 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A210432 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A210432 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A210432 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A210432 008</u>	Jul 19, 2019

PRESCRIPTION DRUG PRODUCT LIST

PREGABALIN

CAPSULE; ORAL

PREGABALIN

<u>AB</u>	SCIEGEN PHARMS INC	<u>25MG</u>	<u>A208677 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A208677 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A208677 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A208677 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A208677 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A208677 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A208677 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A208677 008</u>	Jul 19, 2019
<u>AB</u>	SHUANGCHENG	<u>75MG</u>	<u>A210891 001</u>	Sep 30, 2019
<u>AB</u>		<u>300MG</u>	<u>A210891 002</u>	Sep 30, 2019
<u>AB</u>	SUN PHARM	<u>25MG</u>	<u>A091157 001</u>	Nov 29, 2019
<u>AB</u>		<u>50MG</u>	<u>A091157 002</u>	Nov 29, 2019
<u>AB</u>		<u>75MG</u>	<u>A091157 003</u>	Nov 29, 2019
<u>AB</u>		<u>100MG</u>	<u>A091157 004</u>	Nov 29, 2019
<u>AB</u>		<u>150MG</u>	<u>A091157 005</u>	Nov 29, 2019
<u>AB</u>		<u>200MG</u>	<u>A091157 006</u>	Nov 29, 2019
<u>AB</u>		<u>225MG</u>	<u>A091157 007</u>	Nov 29, 2019
<u>AB</u>		<u>300MG</u>	<u>A091157 008</u>	Nov 29, 2019
<u>AB</u>	TEVA PHARMS	<u>25MG</u>	<u>A091219 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A091219 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A091224 001</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A091224 002</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A091224 003</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A091224 004</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A091224 005</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A091224 006</u>	Jul 19, 2019

SOLUTION; ORAL

LYRICA

<u>AA</u>	+! UPJOHN	<u>20MG/ML</u>	<u>N022488 001</u>	Jan 04, 2010
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PREGABALIN

<u>AA</u>	ALKEM LABS LTD	<u>20MG/ML</u>	<u>A207623 001</u>	Jul 19, 2019
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TABLET, EXTENDED RELEASE; ORAL

LYRICA CR

<u>AB</u>	+ UPJOHN	<u>82.5MG</u>	<u>N209501 001</u>	Oct 11, 2017
<u>AB</u>	+	<u>165MG</u>	<u>N209501 002</u>	Oct 11, 2017
<u>AB</u>	+!	<u>330MG</u>	<u>N209501 003</u>	Oct 11, 2017

PREGABALIN

<u>AB</u>	ALVOGEN	<u>82.5MG</u>	<u>A211593 001</u>	Apr 13, 2021
<u>AB</u>		<u>165MG</u>	<u>A211593 002</u>	Apr 13, 2021
<u>AB</u>		<u>330MG</u>	<u>A211593 003</u>	Apr 13, 2021
<u>AB</u>	APOTEX	<u>165MG</u>	<u>A213313 001</u>	Apr 13, 2021
<u>AB</u>		<u>330MG</u>	<u>A213313 002</u>	Apr 13, 2021
<u>AB</u>	MSN	<u>82.5MG</u>	<u>A213226 001</u>	Apr 13, 2021
<u>AB</u>		<u>165MG</u>	<u>A213226 002</u>	Apr 13, 2021
<u>AB</u>		<u>330MG</u>	<u>A213226 003</u>	Apr 13, 2021
<u>AB</u>	MYLAN	<u>82.5MG</u>	<u>A211948 001</u>	Apr 13, 2021
<u>AB</u>		<u>165MG</u>	<u>A211967 001</u>	Nov 04, 2021
<u>AB</u>	SUN PHARM	<u>82.5MG</u>	<u>A211889 001</u>	Apr 13, 2021
<u>AB</u>		<u>165MG</u>	<u>A211889 002</u>	Apr 13, 2021
<u>AB</u>		<u>330MG</u>	<u>A211889 003</u>	Apr 13, 2021

PRETOMANID

TABLET; ORAL

PRETOMANID

+	MYLAN IRELAND LTD	200MG	N212862 001	Aug 14, 2019
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PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PRILOCAINE HYDROCHLORIDE

!	SEPTODONT INC	4%	A079235 001	Sep 29, 2010
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PRIMAQUINE PHOSPHATE

TABLET; ORAL

PRIMAQUINE

<u>AB</u>	+! SANOFI AVENTIS US	<u>EQ 15MG BASE</u>	<u>N008316 001</u>	
<u>AB</u>	BAYSHORE PHARMS LLC	<u>EQ 15MG BASE</u>	<u>A204476 001</u>	Feb 25, 2014
<u>AB</u>	NOVAST LABS	<u>EQ 15MG BASE</u>	<u>A206043 001</u>	Jun 23, 2016

PRESCRIPTION DRUG PRODUCT LIST

PRIMIDONE

TABLET; ORAL

MYSOLINE

AB	+ !	VALEANT	50MG	N009170	003	
AB	+		250MG	N009170	002	

PRIMIDONE

AB		AMNEAL PHARM	50MG	A040866	001	Apr 23, 2008
AB			250MG	A040866	002	Apr 23, 2008
AB		ANDA REPOSITORY	50MG	A040626	001	Sep 29, 2005
AB			250MG	A040626	002	Sep 29, 2005
AB		LANNETT	50MG	A084903	002	May 24, 2001
AB			250MG	A084903	001	
AB		OXFORD PHARMS	50MG	A040586	001	Feb 24, 2005
AB			250MG	A040586	002	Feb 24, 2005
AB		WATSON LABS	250MG	A083551	001	

PROBENECID

TABLET; ORAL

PROBALAN

AB		LANNETT	500MG	A080966	001	
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PROBENECID

AB	!	WATSON LABS TEVA	500MG	A084442	004	Mar 29, 1983
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PROCAINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

AP	!	HOSPIRA	100MG/ML	A089069	001	Feb 12, 1986
AP		INTL MEDICATION	100MG/ML	A088636	001	Jul 31, 1984
AP		NEXUS PHARMS	100MG/ML	A206332	001	Oct 13, 2017
AP			500MG/ML	A206332	002	Oct 13, 2017
	!	HOSPIRA	500MG/ML	A089070	001	Feb 12, 1986

PROCARBAZINE HYDROCHLORIDE

CAPSULE; ORAL

MATULANE

+ !	LEADIANT BIOSCI INC	EQ 50MG BASE	N016785	001	
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PROCHLORPERAZINE

SUPPOSITORY; RECTAL

COMPRO

AB		PADAGIS US	25MG	A040246	001	Jun 28, 2000
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PROCHLORPERAZINE

AB	!	COSETTE	25MG	A040058	001	Nov 24, 1993
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PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

AP		ATHENEX INC	EQ 5MG BASE/ML	A040540	001	May 28, 2004
AP		CAPLIN	EQ 5MG BASE/ML	A214379	001	Apr 22, 2021
AP	!	EMCURE PHARMS LTD	EQ 5MG BASE/ML	A204147	001	Oct 15, 2013
AP		GLAND PHARMA LTD	EQ 5MG BASE/ML	A214107	001	Sep 22, 2021
AP		HIKMA	EQ 5MG BASE/ML	A089903	001	Aug 29, 1989
AP		MYLAN LABS LTD	EQ 5MG BASE/ML	A210710	001	Oct 25, 2018
AP		NEXUS PHARMS	EQ 5MG BASE/ML	A204860	001	Feb 15, 2019
AP		VIWIT PHARM	EQ 5MG BASE/ML	A213626	001	Sep 28, 2021

PROCHLORPERAZINE MALEATE

TABLET; ORAL

PROCHLORPERAZINE MALEATE

AB		MYLAN	EQ 5MG BASE	A040185	002	Oct 28, 1996
AB			EQ 10MG BASE	A040185	001	Oct 28, 1996

PROCOMP

AB		JUBILANT CADISTA	EQ 5MG BASE	A040268	001	Feb 27, 1998
AB	!		EQ 10MG BASE	A040268	002	Feb 27, 1998

PROGESTERONE

CAPSULE; ORAL

PROGESTERONE

AB		AMNEAL PHARMS NY	100MG	A207724	001	Sep 07, 2017
AB			200MG	A207724	002	Sep 07, 2017
AB		BIONPHARMA INC	100MG	A200900	001	Aug 16, 2013
AB			200MG	A200900	002	Aug 16, 2013
AB		DR REDDYS	100MG	A208801	001	Feb 28, 2017
AB			200MG	A208801	002	Feb 28, 2017
AB		EUGIA PHARMA	100MG	A211285	001	Oct 26, 2018
AB			200MG	A211285	002	Oct 26, 2018

PRESCRIPTION DRUG PRODUCT LIST

PROGESTERONE

CAPSULE; ORAL

PROGESTERONE

<u>AB</u>	XIROMED	<u>100MG</u>	<u>A205229 001</u>	Oct 20, 2017
<u>AB</u>		<u>200MG</u>	<u>A205229 002</u>	Oct 20, 2017

PROMETRIUM

<u>AB</u>	+ VIRTUS PHARMS	<u>100MG</u>	<u>N019781 001</u>	May 14, 1998
<u>AB</u>	+!	<u>200MG</u>	<u>N019781 002</u>	Oct 15, 1999

PROGESTERONE

BX	SOFGEN PHARMS	100MG	A200456 001	Sep 28, 2012
BX		200MG	A200456 002	Sep 28, 2012

GEL; VAGINAL

CRINONE

+!	ALLERGAN	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 002</u>	
<u>AO</u>	EUGIA PHARMA	<u>50MG/ML</u>	<u>A210965 001</u>	Dec 06, 2018
<u>AO</u>	FRESENIUS KABI USA	<u>50MG/ML</u>	<u>A075906 001</u>	Apr 25, 2001
<u>AO</u>	HIKMA FARMACEUTICA	<u>50MG/ML</u>	<u>A091033 001</u>	Oct 28, 2010
<u>AO</u>	XIROMED	<u>50MG/ML</u>	<u>A215634 001</u>	Jan 05, 2022

INSERT; VAGINAL

ENDOMETRIN

+!	FERRING	100MG	N022057 001	Jun 21, 2007
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PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROMETHAZINE HYDROCHLORIDE

<u>AP</u>	! WEST-WARD PHARMS	<u>25MG/ML</u>	<u>A083312 001</u>	
	INT			
<u>AP</u>	!	<u>50MG/ML</u>	<u>A083312 002</u>	
<u>AP</u>	XGEN PHARMS	<u>25MG/ML</u>	<u>A040737 001</u>	Apr 24, 2008
<u>AP</u>		<u>50MG/ML</u>	<u>A040737 002</u>	Apr 24, 2008

SUPPOSITORY; RECTAL

PROMETHAZINE HYDROCHLORIDE

<u>AB</u>	COSETTE	<u>12.5MG</u>	<u>A040428 002</u>	Mar 31, 2003
<u>AB</u>	!	<u>25MG</u>	<u>A040428 001</u>	Feb 05, 2002
<u>AB</u>	PADAGIS ISRAEL	<u>12.5MG</u>	<u>A040500 001</u>	Jun 30, 2003
<u>AB</u>		<u>25MG</u>	<u>A040500 002</u>	Jun 30, 2003
<u>AB</u>	TARO	<u>12.5MG</u>	<u>A040603 001</u>	Oct 26, 2006
<u>AB</u>		<u>25MG</u>	<u>A040603 002</u>	Oct 26, 2006

PROMETHEGAN

!	COSETTE	50MG	A087165 001	Aug 14, 1987
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SYRUP; ORAL

PROMETHAZINE HYDROCHLORIDE

<u>AA</u>	AKORN	<u>6.25MG/5ML</u>	<u>A040026 001</u>	Sep 25, 1998
<u>AA</u>	AMNEAL PHARMS	<u>6.25MG/5ML</u>	<u>A040882 001</u>	Dec 30, 2009
<u>AA</u>	NOSTRUM LABS INC	<u>6.25MG/5ML</u>	<u>A040891 001</u>	Mar 13, 2009
<u>AA</u>	PHARM ASSOC	<u>6.25MG/5ML</u>	<u>A040643 001</u>	Apr 26, 2006
<u>AA</u>	QUAGEN	<u>6.25MG/5ML</u>	<u>A213890 001</u>	Jul 12, 2021
<u>AA</u>	TARO	<u>6.25MG/5ML</u>	<u>A040718 001</u>	Apr 04, 2007
<u>AA</u>	TRIS PHARMA INC	<u>6.25MG/5ML</u>	<u>A091675 001</u>	Jun 28, 2012

PROMETHAZINE PLAIN

<u>AA</u>	+! WOCKHARDT BIO AG	<u>6.25MG/5ML</u>	<u>A087953 001</u>	Nov 15, 1982
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TABLET; ORAL

PROMETHAZINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS NY	<u>12.5MG</u>	<u>A091179 001</u>	Dec 13, 2010
<u>AB</u>		<u>25MG</u>	<u>A091179 002</u>	Dec 13, 2010
<u>AB</u>		<u>50MG</u>	<u>A091179 003</u>	Dec 13, 2010
<u>AB</u>	KVK TECH	<u>12.5MG</u>	<u>A040712 002</u>	May 04, 2007
<u>AB</u>		<u>25MG</u>	<u>A040712 001</u>	Jul 31, 2006
<u>AB</u>		<u>50MG</u>	<u>A040712 003</u>	Jul 31, 2006
<u>AB</u>	PRINSTON INC	<u>12.5MG</u>	<u>A040622 001</u>	Jul 18, 2006
<u>AB</u>		<u>25MG</u>	<u>A040622 002</u>	Jul 18, 2006
<u>AB</u>		<u>50MG</u>	<u>A040622 003</u>	Jul 18, 2006
<u>AB</u>	QUAGEN	<u>12.5MG</u>	<u>A040673 001</u>	Mar 05, 2008
<u>AB</u>		<u>25MG</u>	<u>A040673 002</u>	Mar 05, 2008
<u>AB</u>		<u>50MG</u>	<u>A040673 003</u>	Mar 05, 2008
<u>AB</u>	+ SANDOZ	<u>25MG</u>	<u>A084176 003</u>	
<u>AB</u>	+!	<u>50MG</u>	<u>A084176 001</u>	
<u>AB</u>	STRIDES PHARMA	<u>12.5MG</u>	<u>A209177 001</u>	Jun 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A209177 002</u>	Jun 30, 2017

PRESCRIPTION DRUG PRODUCT LIST

PROMETHAZINE HYDROCHLORIDE

TABLET; ORAL

PROMETHAZINE HYDROCHLORIDE

<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

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>	RISING	<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>AB</u>	STRIDES PHARMA	<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>	TWI PHARMS	<u>225MG</u>	<u>A212928</u>	<u>001</u>	Jun 18, 2020
<u>AB</u>		<u>325MG</u>	<u>A212928</u>	<u>002</u>	Jun 18, 2020
<u>AB</u>		<u>425MG</u>	<u>A212928</u>	<u>003</u>	Jun 18, 2020
<u>AB</u>	UPSHER SMITH LABS	<u>225MG</u>	<u>A212744</u>	<u>001</u>	Jun 25, 2020
<u>AB</u>		<u>325MG</u>	<u>A212744</u>	<u>002</u>	Jun 25, 2020
<u>AB</u>		<u>425MG</u>	<u>A212744</u>	<u>003</u>	Jun 25, 2020
<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>	ZYDUS	<u>225MG</u>	<u>A214184</u>	<u>001</u>	Apr 21, 2021
<u>AB</u>		<u>325MG</u>	<u>A214184</u>	<u>002</u>	Apr 21, 2021
<u>AB</u>		<u>425MG</u>	<u>A214184</u>	<u>003</u>	Apr 21, 2021

RYTHMOL SR

<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

PROPAFENONE HYDROCHLORIDE

BX	SINOTHERAPEUTICS INC	225MG	A210339	001	Jan 04, 2019
BX		325MG	A210339	002	Jan 04, 2019
BX		425MG	A210339	003	Jan 04, 2019

TABLET; ORAL

PROPAFENONE HYDROCHLORIDE

<u>AB</u>	ANI PHARMS	<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>	STRIDES PHARMA	<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>	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>	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

PROPARACAINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ALCAINE

<u>AT</u>	ALCON LABS INC	<u>0.5%</u>	<u>A080027</u>	<u>001</u>	
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PROPARACAINE HYDROCHLORIDE

<u>AT</u>	!	AKORN	<u>0.5%</u>	<u>A040277</u>	<u>001</u>	Mar 16, 2000
<u>AT</u>		BAUSCH AND LOMB	<u>0.5%</u>	<u>A040074</u>	<u>001</u>	Sep 29, 1995

PRESCRIPTION DRUG PRODUCT LIST

PROPOFOL

INJECTABLE; INJECTION

DIPRIVAN

AB	+!	FRESENIUS KABI USA	10MG/ML	N019627 002	Jun 11, 1996
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PROPOFOL

AB		DR REDDYS	10MG/ML	A205067 001	Nov 15, 2018
AB		EMCURE PHARMS LTD	10MG/ML	A206408 001	Oct 12, 2021
AB		HIKMA	10MG/ML	A074848 001	Apr 19, 2005
AB		HOSPIRA	10MG/ML	A077908 001	Mar 17, 2006
AB		INNOPHARMA	10MG/ML	A205576 001	Sep 16, 2020
AB		SAGENT PHARMS INC	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	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		ADARE PHARMS INC	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		AMTA	60MG	A212026 001	Jan 06, 2020
AB			80MG	A212026 002	Jan 06, 2020
AB			120MG	A212026 003	Jan 06, 2020
AB			160MG	A212026 004	Jan 06, 2020
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		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	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			
	!	HIKMA	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		INNOGENIX	10MG	A070322 002	Oct 22, 1985
AB			20MG	A070322 003	Oct 22, 1985
AB			40MG	A070322 004	Oct 22, 1985
AB			60MG	A070322 005	Sep 24, 1986
AB			80MG	A070322 001	Aug 04, 1986
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

PRESCRIPTION DRUG PRODUCT LIST

PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

AB		20MG	A070213 003	Nov 19, 1985
AB		40MG	A070213 001	Nov 19, 1985
AB		60MG	A070213 005	Apr 08, 2011
AB		80MG	A070213 004	Nov 19, 1985
AB	NORTHSTAR HLTHCARE	10MG	A078213 001	Jan 10, 2008
AB		20MG	A078213 002	Jan 10, 2008
AB		40MG	A078213 003	Jan 10, 2008
AB		60MG	A078213 004	Jan 10, 2008
AB		80MG	A078213 005	Jan 10, 2008
AB	VINTAGE PHARMS	10MG	A070221 002	Aug 01, 1986
AB		20MG	A070221 003	Aug 01, 1986
AB		40MG	A070221 004	Aug 01, 1986
AB		60MG	A070221 005	Sep 24, 1986
AB		80MG	A070221 001	Apr 14, 1986
AB	WATSON LABS	10MG	A070175 001	May 13, 1986
AB		20MG	A070176 001	May 13, 1986
AB		40MG	A070177 001	May 13, 1986
AB	!	80MG	A070178 001	May 13, 1986

PROPYLTHIOURACIL

TABLET; ORAL

PROPYLTHIOURACIL

BD	ACTAVIS ELIZABETH	50MG	A080172 001	
BD	+! DAVA PHARMS INC	50MG	N006188 001	
BD	QUAGEN	50MG	A080154 001	

PROTAMINE SULFATE

INJECTABLE; INJECTION

PROTAMINE SULFATE

!	FRESENIUS KABI USA	10MG/ML	A089454 001	Apr 07, 1987
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PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

PROTRIPTYLINE HYDROCHLORIDE

AB	EPIC PHARMA LLC	5MG	A202220 001	Nov 19, 2012
AB		10MG	A202220 002	Nov 19, 2012
AB	HIKMA	5MG	A078913 001	Sep 16, 2008
AB	!	10MG	A078913 002	Sep 16, 2008
AB	SIGMAPHARM LABS LLC	5MG	A090462 001	May 03, 2010
AB		10MG	A090462 002	May 03, 2010

PRUCALOPRIDE SUCCINATE

TABLET; ORAL

MOTEGRITY

+	TAKEDA PHARMS USA	EQ 1MG BASE	N210166 001	Dec 14, 2018
+!		EQ 2MG BASE	N210166 002	Dec 14, 2018

PYRAZINAMIDE

TABLET; ORAL

PYRAZINAMIDE

AB	!	AKORN	500MG	A081319 001	Jun 30, 1992
AB		MACLEODS PHARMS LTD	500MG	A212541 001	Jul 27, 2020
AB	+	NOVITIUM PHARMA	500MG	A080157 001	

PYRIDOSTIGMINE BROMIDE

INJECTABLE; INJECTION

MESTINON

AP	+!	BAUSCH	5MG/ML	N009830 001	
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REGONOL

AP		SANDOZ INC	5MG/ML	N017398 001	
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SYRUP; ORAL

MESTINON

AA	+!	BAUSCH	60MG/5ML	N015193 001	
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PYRIDOSTIGMINE BROMIDE

AA		AMNEAL	60MG/5ML	A212702 001	Jan 10, 2020
AA		ELYSIUM	60MG/5ML	A208797 001	Jan 09, 2020
AA		NOVITIUM PHARMA	60MG/5ML	A211694 001	Mar 08, 2019

TABLET; ORAL

MESTINON

AB	+!	BAUSCH	60MG	N009829 002	
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PYRIDOSTIGMINE BROMIDE

AB		EYWA	60MG	A211181 001	Jul 20, 2018
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PRESCRIPTION DRUG PRODUCT LIST

PYRIDOSTIGMINE BROMIDE

TABLET;ORAL

PYRIDOSTIGMINE BROMIDE

AB	IMPAX LABS	60MG	A040502 001	Apr 24, 2003
AB	ZYDUS PHARMS	60MG	A205650 001	Jun 22, 2015
	EYWA	30MG	A211181 002	May 14, 2019

TABLET, EXTENDED RELEASE;ORAL

MESTINON

AB	+ ! BAUSCH	180MG	N011665 001	
	<u>PYRIDOSTIGMINE BROMIDE</u>			
AB	ALVOGEN	180MG	A204737 001	Jun 26, 2015
AB	IMPAX LABS INC	180MG	A203184 001	Sep 15, 2015
AB	RISING	180MG	A205464 001	Aug 15, 2017

PYRIDOXINE HYDROCHLORIDE

INJECTABLE;INJECTION

PYRIDOXINE HYDROCHLORIDE

+ !	FRESENIUS KABI USA	100MG/ML	A080618 001	
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PYRIMETHAMINE

TABLET;ORAL

DARAPRIM

AB	+ ! VYERA PHARMS LLC	25MG	N008578 001	
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PYRIMETHAMINE

AB	ALVOGEN	25MG	A211271 001	Jul 27, 2021
AB	CEROVENE INC	25MG	A207127 001	Feb 28, 2020
AB	TEVA PHARMS	25MG	A215506 001	Aug 13, 2021

QUAZEPAM

TABLET;ORAL

DORAL

+ !	GALT PHARMS	15MG	N018708 001	Dec 27, 1985
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QUETIAPINE FUMARATE

TABLET;ORAL

QUETIAPINE FUMARATE

AB	ACCORD HLTHCARE	EQ 25MG BASE	A202152 001	Mar 27, 2012
AB		EQ 50MG BASE	A202152 002	Mar 27, 2012
AB		EQ 100MG BASE	A202152 003	Mar 27, 2012
AB		EQ 200MG BASE	A202152 004	Mar 27, 2012
AB		EQ 300MG BASE	A202152 005	Mar 27, 2012
AB		EQ 400MG BASE	A202152 006	Mar 27, 2012
AB	ALKEM LABS LTD	EQ 25MG BASE	A201504 001	Feb 12, 2013
AB		EQ 50MG BASE	A201504 002	Feb 12, 2013
AB		EQ 100MG BASE	A201504 003	Feb 12, 2013
AB		EQ 150MG BASE	A201504 004	Feb 12, 2013
AB		EQ 200MG BASE	A201504 005	Feb 12, 2013
AB		EQ 300MG BASE	A201504 006	Feb 12, 2013
AB		EQ 400MG BASE	A201504 007	Feb 12, 2013
AB	AUROBINDO PHARMA LTD	EQ 25MG BASE	A091388 001	Mar 27, 2012
AB		EQ 50MG BASE	A091388 002	Mar 27, 2012
AB		EQ 100MG BASE	A091388 003	Mar 27, 2012
AB		EQ 150MG BASE	A091388 004	Mar 27, 2012
AB		EQ 200MG BASE	A091388 005	Mar 27, 2012
AB		EQ 300MG BASE	A091388 006	Mar 27, 2012
AB		EQ 400MG BASE	A091388 007	Mar 27, 2012
AB	DR REDDYS LABS LTD	EQ 25MG BASE	A077380 001	Mar 27, 2012
AB		EQ 50MG BASE	A077380 002	Mar 27, 2012
AB		EQ 100MG BASE	A077380 003	Mar 27, 2012
AB		EQ 150MG BASE	A077380 004	Mar 27, 2012
AB		EQ 200MG BASE	A077380 005	Mar 27, 2012
AB		EQ 300MG BASE	A077380 006	Mar 27, 2012
AB		EQ 400MG BASE	A077380 007	Mar 27, 2012
AB	HIKMA	EQ 25MG BASE	A090120 001	Mar 27, 2012
AB		EQ 50MG BASE	A090749 001	Mar 27, 2012
AB		EQ 100MG BASE	A090749 002	Mar 27, 2012
AB		EQ 200MG BASE	A090749 003	Mar 27, 2012
AB		EQ 300MG BASE	A090749 004	Mar 27, 2012
AB		EQ 400MG BASE	A090749 005	Mar 27, 2012
AB	LUPIN LTD	EQ 25MG BASE	A201109 001	Mar 27, 2012
AB		EQ 50MG BASE	A201109 002	Mar 27, 2012
AB		EQ 100MG BASE	A201109 003	Mar 27, 2012
AB		EQ 200MG BASE	A201109 004	Mar 27, 2012
AB		EQ 300MG BASE	A201109 005	Mar 27, 2012

PRESCRIPTION DRUG PRODUCT LIST

QUETIAPINE FUMARATE

TABLET;ORAL

QUETIAPINE FUMARATE

<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A201109 006</u>	Mar 27, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A203359 001</u>	May 17, 2016
<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 PHARM	<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>	UNICHEM	<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>	ZENNOVA	<u>EQ 25MG BASE</u>	<u>A090960 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A090960 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090960 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A090960 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A090960 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A090960 006</u>	Mar 27, 2012
<u>SEROQUEL</u>				
<u>AB</u>	+! ASTRAZENECA	<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

QUETIAPINE FUMARATE

<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>	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
<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

PRESCRIPTION DRUG PRODUCT LIST

QUETIAPINE FUMARATE

TABLET, EXTENDED RELEASE;ORAL

QUETIAPINE FUMARATE

<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

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

QUETIAPINE FUMARATE

BX		SCIEGEN PHARMS INC	EQ 50MG BASE	A209635 005	Nov 16, 2018
BX			EQ 150MG BASE	A209635 001	Nov 29, 2017
BX			EQ 200MG BASE	A209635 002	Nov 29, 2017
BX			EQ 300MG BASE	A209635 003	Nov 29, 2017
BX			EQ 400MG BASE	A209635 004	Nov 29, 2017

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>		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
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A205823 003</u>	Sep 15, 2016
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A205823 004</u>	Sep 15, 2016

QUINIDINE GLUCONATE

TABLET, EXTENDED RELEASE;ORAL

QUINIDINE GLUCONATE

<u>AB</u>		EYWA	<u>324MG</u>	<u>A212589 001</u>	Sep 17, 2021
<u>AB</u>	!	SUN PHARM INDUSTRIES	<u>324MG</u>	<u>A089338 001</u>	Feb 11, 1987

PRESCRIPTION DRUG PRODUCT LIST

QUINIDINE SULFATE

TABLET;ORAL

QUINIDINE SULFATE

! SANDOZ

200MG

A088072 002

!

300MG

A088072 001 Sep 26, 1983

QUININE SULFATE

CAPSULE;ORAL

QUALAQUIN**AB** +! SUN PHARM
INDUSTRIES**324MG****N021799 001** Aug 12, 2005QUININE SULFATE**AB** AMNEAL PHARMS**324MG****A203729 001** Jul 15, 2015**AB** LUPIN LTD**324MG****A203112 001** Apr 24, 2015**AB** NOVAST LABS**324MG****A204372 001** Jul 22, 2015**AB** TEVA PHARMS**324MG****A091661 001** Sep 28, 2012RABEPRAZOLE SODIUM

TABLET, DELAYED RELEASE;ORAL

ACIPHEX**AB** +! WOODWARD**20MG****N020973 002** Aug 19, 1999RABEPRAZOLE 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** DR REDDYS**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** RUBICON**20MG****A204237 001** Nov 18, 2015**AB** TEVA PHARMS USA**20MG****A076822 001** Nov 08, 2013**AB** TORRENT**20MG****A202376 001** Nov 08, 2013RADIUM 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, 1997RALOXIFENE HYDROCHLORIDE**AB** AMNEAL PHARMS**60MG****A208206 001** Apr 08, 2016**AB** AUROBINDO PHARMA
LTD**60MG****A204310 001** Aug 28, 2015**AB** CADILA PHARMS LTD**60MG****A211324 001** Aug 18, 2020**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**AB** TEVA PHARMS USA**60MG****A078193 001** Mar 04, 2014**AB** WATSON LABS INC**60MG****A200825 001** Jan 21, 2015RALTEGRAVIR 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**AB** ACTAVIS LABS FL INC**8MG****A091610 001** Aug 19, 2015**AB** APPCO**8MG****A213815 001** Oct 26, 2020**AB** DR REDDYS LABS SA**8MG****A091693 001** Jul 26, 2013**AB** GRANULES**8MG****A213186 001** Aug 21, 2020**AB** I3 PHARMS**8MG****A212650 001** Apr 10, 2020**AB** ZYDUS PHARMS**8MG****A211567 001** Jul 22, 2019

PRESCRIPTION DRUG PRODUCT LIST

RAMELTEON

TABLET; ORAL

ROZEREM

<u>AB</u>	<u>+</u> !	TAKEDA PHARMS USA	<u>8MG</u>	<u>N021782</u>	<u>001</u>	Jul 22, 2005
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RAMIPRIL

CAPSULE; ORAL

ALTACE

<u>AB</u>	<u>+</u>	KING PHARMS LLC	<u>1.25MG</u>	<u>N019901</u>	<u>001</u>	Jan 28, 1991
<u>AB</u>	<u>+</u>		<u>2.5MG</u>	<u>N019901</u>	<u>002</u>	Jan 28, 1991
<u>AB</u>	<u>+</u>		<u>5MG</u>	<u>N019901</u>	<u>003</u>	Jan 28, 1991
<u>AB</u>	<u>+</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>		HIKMA	<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>		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>		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>		ZYDUS PHARMS USA	<u>1.25MG</u>	<u>A078832</u>	<u>001</u>	Sep 02, 2008
<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 HYDROCHLORIDE

CAPSULE; ORAL

RANITIDINE HYDROCHLORIDE

<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>!</u>		<u>EQ 300MG BASE</u>	<u>A074655</u>	<u>002</u>	Oct 22, 1997

INJECTABLE; INJECTION

RANITIDINE HYDROCHLORIDE

<u>AP</u>		HIKMA	<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>		MYLAN LABS LTD	<u>EQ 25MG BASE/ML</u>	<u>A079076</u>	<u>001</u>	Jun 09, 2016
<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>	<u>+</u> !	TELIGENT	<u>EQ 25MG BASE/ML</u>	<u>N019090</u>	<u>001</u>	Oct 19, 1984
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SYRUP; ORAL

RANITIDINE HYDROCHLORIDE

<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>	<u>!</u>	PHARM ASSOC	<u>EQ 15MG BASE/ML</u>	<u>A077405</u>	<u>001</u>	Sep 21, 2007

PRESCRIPTION DRUG PRODUCT LIST

RANITIDINE HYDROCHLORIDE

TABLET;ORAL

RANITIDINE HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 150MG BASE</u>	<u>A074680 001</u>	Sep 12, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A074680 002</u>	Sep 12, 1997
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 150MG BASE</u>	<u>A076705 001</u>	Jul 27, 2005
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A076705 002</u>	Jul 27, 2005
<u>AB</u>	GLENMARK PHARMS INC	<u>EQ 150MG BASE</u>	<u>A078542 001</u>	Nov 19, 2008
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A078542 002</u>	Nov 19, 2008
<u>AB</u>	PAR PHARM	<u>EQ 150MG BASE</u>	<u>A075180 001</u>	Jan 28, 1999
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A075180 002</u>	Jan 28, 1999
<u>AB</u>	SANDOZ	<u>EQ 150MG BASE</u>	<u>A074467 001</u>	Aug 29, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A074467 002</u>	Aug 29, 1997
<u>AB</u>	VKT PHARMA	<u>EQ 150MG BASE</u>	<u>A211289 001</u>	Jan 31, 2019
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A211289 002</u>	Jan 31, 2019

RANOLAZINE

TABLET, EXTENDED RELEASE;ORAL

RANEXA

<u>AB</u>	+ GILEAD	<u>500MG</u>	<u>N021526 002</u>	Jan 27, 2006
<u>AB</u>	+!	<u>1GM</u>	<u>N021526 001</u>	Feb 12, 2007

RANOLAZINE

<u>AB</u>	ACTAVIS ELIZABETH	<u>500MG</u>	<u>A208862 001</u>	May 28, 2019
<u>AB</u>		<u>1GM</u>	<u>A208862 002</u>	May 28, 2019
<u>AB</u>	AJANTA PHARMA LTD	<u>500MG</u>	<u>A210054 001</u>	May 28, 2019
<u>AB</u>		<u>1GM</u>	<u>A210054 002</u>	May 28, 2019
<u>AB</u>	ALKEM LABS LTD	<u>500MG</u>	<u>A209953 001</u>	Nov 30, 2020
<u>AB</u>		<u>1GM</u>	<u>A209953 002</u>	Nov 30, 2020
<u>AB</u>	CADILA	<u>500MG</u>	<u>A210188 001</u>	Aug 19, 2019
<u>AB</u>		<u>1GM</u>	<u>A210188 002</u>	Aug 19, 2019
<u>AB</u>	GLENMARK PHARMS LTD	<u>500MG</u>	<u>A211082 001</u>	Jul 05, 2019
<u>AB</u>		<u>1GM</u>	<u>A211082 002</u>	Jul 05, 2019
<u>AB</u>	GRAVITI PHARMS	<u>500MG</u>	<u>A212889 001</u>	Jan 28, 2021
<u>AB</u>		<u>1GM</u>	<u>A212889 002</u>	Jan 28, 2021
<u>AB</u>	LUPIN LTD	<u>500MG</u>	<u>A201046 001</u>	Jul 29, 2013
<u>AB</u>		<u>1GM</u>	<u>A201046 002</u>	Jul 29, 2013
<u>AB</u>	MANKIND PHARMA	<u>500MG</u>	<u>A212284 001</u>	Feb 12, 2020
<u>AB</u>		<u>1GM</u>	<u>A212284 002</u>	Feb 12, 2020
<u>AB</u>	MICRO LABS	<u>500MG</u>	<u>A211745 001</u>	Feb 27, 2020
<u>AB</u>		<u>1GM</u>	<u>A211745 002</u>	Feb 27, 2020
<u>AB</u>	PRAXGEN	<u>500MG</u>	<u>A212781 001</u>	Mar 23, 2020
<u>AB</u>		<u>1GM</u>	<u>A212781 002</u>	Mar 23, 2020
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A211829 001</u>	Jun 04, 2019
<u>AB</u>		<u>1GM</u>	<u>A211829 002</u>	Jun 04, 2019
<u>AB</u>	SUN PHARM	<u>500MG</u>	<u>A211707 001</u>	May 28, 2019
<u>AB</u>		<u>1GM</u>	<u>A211707 002</u>	May 28, 2019
<u>AB</u>	SUNSHINE	<u>500MG</u>	<u>A211865 001</u>	Mar 23, 2020
<u>AB</u>		<u>1GM</u>	<u>A211865 002</u>	Mar 23, 2020

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>	AUROBINDO PHARMA USA	<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>	DR REDDYS	<u>EQ 0.5MG BASE</u>	<u>A201942 001</u>	Nov 18, 2021
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A201942 002</u>	Nov 18, 2021
<u>AB</u>	INDOCO	<u>EQ 0.5MG BASE</u>	<u>A206153 001</u>	Oct 04, 2019
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A206153 002</u>	Oct 04, 2019
<u>AB</u>	MICRO LABS	<u>EQ 0.5MG BASE</u>	<u>A207004 001</u>	Mar 29, 2019
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A207004 002</u>	Mar 29, 2019
<u>AB</u>	ORBION PHARMS	<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

PRESCRIPTION DRUG PRODUCT LIST

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

RELUGOLIX

TABLET; ORAL

ORGOVYX

+! MYOVANT SCIENCES 120MG N214621 001 Dec 18, 2020

REMDESIVIR

POWDER; INTRAVENOUS

VEKLURY

+! GILEAD SCIENCES INC 100MG/VIAL N214787 001 Oct 22, 2020

SOLUTION; INTRAVENOUS

VEKLURY

+! GILEAD SCIENCES INC 100MG/20ML (5MG/ML) N214787 002 Oct 22, 2020

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
<u>AP</u>	NAVINTA LLC	<u>EQ 1MG BASE/VIAL</u>	<u>A210594 001</u>	Oct 13, 2020
<u>AP</u>		<u>EQ 2MG BASE/VIAL</u>	<u>A210594 002</u>	Oct 13, 2020
<u>AP</u>		<u>EQ 5MG BASE/VIAL</u>	<u>A210594 003</u>	Oct 13, 2020

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

REMIMAZOLAM BESYLATE

POWDER; INTRAVENOUS

BYFAVO

+! ACACIA EQ 20MG BASE/VIAL N212295 001 Oct 06, 2020

REPAGLINIDE

TABLET; ORAL

REPAGLINIDE

<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>	CHARTWELL RX	<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>	PADAGIS US	<u>0.5MG</u>	<u>A201189 001</u>	Jul 17, 2013
<u>AB</u>		<u>1MG</u>	<u>A201189 002</u>	Jan 22, 2014
<u>AB</u>		<u>2MG</u>	<u>A201189 003</u>	Jan 22, 2014
<u>AB</u>	SUN PHARM INDS INC	<u>1MG</u>	<u>A077571 002</u>	Jul 11, 2013
<u>AB</u>		<u>2MG</u>	<u>A077571 003</u>	Jul 11, 2013

RETAPAMULIN

OINTMENT; TOPICAL

ALTABAX

+! ALMIRALL 1% N022055 001 Apr 12, 2007

REVEFENACIN

SOLUTION; INHALATION

YUPELRI

+! MYLAN IRELAND LTD 175MCG/3ML N210598 001 Nov 09, 2018

PRESCRIPTION DRUG PRODUCT LIST

RIBAVIRIN

CAPSULE;ORAL

REBETOL

AB	+ !	MERCK SHARP DOHME	200MG	N020903	002	Jul 25, 2001
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RIBAVIRIN

AB		AUROBINDO PHARMA	200MG	A079117	001	Sep 17, 2009
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AB		ZYDUS PHARMS USA	200MG	A077224	001	Oct 28, 2005
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FOR SOLUTION;INHALATION

RIBAVIRIN

AN		NAVINTA LLC	6GM/VIAL	A207366	001	Oct 06, 2016
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VIRAZOLE

AN	+ !	BAUSCH	6GM/VIAL	N018859	001	Dec 31, 1985
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TABLET;ORAL

RIBAVIRIN

AB		AUROBINDO PHARMA	200MG	A079111	001	Sep 17, 2009
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AB		SANDOZ	200MG	A077743	001	Oct 03, 2006
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AB		ZYDUS PHARMS USA	200MG	A077094	001	Dec 05, 2005
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RIBOCICLIB SUCCINATE

TABLET;ORAL

KISQALI

+ !	NOVARTIS	EQ 200MG BASE		N209092	001	Mar 13, 2017
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RIBOFLAVIN 5'-PHOSPHATE SODIUM

SOLUTION/DROPS;OPHTHALMIC

PHOTREXA

+ !	GLAUKOS	0.146%		N203324	001	Apr 15, 2016
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PHOTREXA VISCOUS IN DEXTRAN 20%

+ !	GLAUKOS	0.146%		N203324	002	Apr 15, 2016
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RIFABUTIN

CAPSULE;ORAL

MYCOBUTIN

AB	+ !	PFIZER	150MG	N050689	001	Dec 23, 1992
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RIFABUTIN

AB		LUPIN LTD	150MG	A090033	001	Feb 24, 2014
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AB		NOVITIUM PHARMA	150MG	A215041	001	Dec 17, 2021
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RIFAMPIN

CAPSULE;ORAL

RIFAMPIN

AB		AKORN	150MG	A065028	001	Mar 14, 2001
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AB			300MG	A065028	002	Mar 14, 2001
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AB		LANNETT CO INC	150MG	A065390	001	Mar 28, 2008
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AB			300MG	A065390	002	Mar 28, 2008
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AB		LUPIN PHARMS	150MG	A090034	001	Aug 21, 2013
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AB	!		300MG	A090034	002	Aug 21, 2013
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AB		SANDOZ	150MG	A064150	002	Jan 02, 1998
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AB			300MG	A064150	001	May 28, 1997
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RIMACTANE

AB		OXFORD PHARMS	300MG	N050429	001	
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INJECTABLE;INJECTION

RIFADIN

AP	+ !	SANOFI AVENTIS US	600MG/VIAL	N050627	001	May 25, 1989
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RIFAMPIN

AP		AKORN	600MG/VIAL	A065502	001	Sep 21, 2010
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AP		FRESENIUS KABI USA	600MG/VIAL	A091181	001	Aug 21, 2014
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AP		HIKMA	600MG/VIAL	A064217	001	Oct 29, 1999
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AP		HIKMA PHARMS	600MG/VIAL	A205039	001	Mar 03, 2016
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AP		MYLAN LABS LTD	600MG/VIAL	A065421	001	May 22, 2008
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RIFAMYCIN SODIUM

TABLET, DELAYED RELEASE;ORAL

AEMCOLO

+ !	REDHILL	EQ 194MG BASE		N210910	001	Nov 16, 2018
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RIFAPENTINE

TABLET;ORAL

PRIFTIN

+ !	SANOFI AVENTIS US	150MG		N021024	001	Jun 22, 1998
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PRESCRIPTION DRUG PRODUCT LIST

RIFAXIMIN

TABLET;ORAL

XIFAXAN

+	!	SALIX PHARMS	200MG	N021361	001	May 25, 2004
+	!		550MG	N021361	002	Mar 24, 2010

RILPIVIRINE HYDROCHLORIDE

TABLET;ORAL

EDURANT

+	!	JANSSEN PRODS	EQ 25MG BASE	N202022	001	May 20, 2011
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RILUZOLE

FILM;ORAL

EXSERVAN

+	!	mitsubishi holdings	50MG	N212640	001	Nov 22, 2019
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SUSPENSION;ORAL

TIGLUTIK KIT

+	!	ITALFARMACO SPA	50MG/10ML	N209080	001	Sep 05, 2018
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TABLET;ORAL

RILUTEK

AB	+	!	COVIS	50MG	N020599	001	Dec 12, 1995
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RILUZOLE

AB			ALKEM LABS LTD	50MG	A204048	001	Mar 30, 2016
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AB			GLENMARK PHARMS LTD	50MG	A091394	001	Jun 18, 2013
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AB			IMPAX LABS	50MG	A076173	001	Jan 29, 2003
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AB			KENTON	50MG	A206045	001	Dec 09, 2019
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AB			MYLAN PHARMS INC	50MG	A203042	001	Jul 01, 2013
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AB			SUN PHARM INDS LTD	50MG	A091417	001	Jun 18, 2013
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RIMANTADINE HYDROCHLORIDE

TABLET;ORAL

FLUMADINE

AB	+	!	SUN PHARM INDS INC	100MG	N019649	001	Sep 17, 1993
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RIMANTADINE HYDROCHLORIDE

AB			IMPAX LABS	100MG	A076132	001	Aug 30, 2002
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RIMEGEPANT SULFATE

TABLET, ORALLY DISINTEGRATING;ORAL

NURTEC ODT

+	!	BIOHAVEN IRELAND	EQ 75MG BASE	N212728	001	Feb 27, 2020
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RIOCIGUAT

TABLET;ORAL

ADEMPAS

+			BAYER HLTHCARE	0.5MG	N204819	001	Oct 08, 2013
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+				1MG	N204819	002	Oct 08, 2013
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+				1.5MG	N204819	003	Oct 08, 2013
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+				2MG	N204819	004	Oct 08, 2013
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+	!			2.5MG	N204819	005	Oct 08, 2013
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RIPRETINIB

TABLET;ORAL

QINLOCK

+	!	DECIPHERA PHARMS	50MG	N213973	001	May 15, 2020
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RISDIPLAM

FOR SOLUTION;ORAL

EVRYSDI

+	!	GENENTECH INC	0.75MG/ML	N213535	001	Aug 07, 2020
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RISEDRONATE SODIUM

TABLET;ORAL

ACTONEL

AB	+		APIL	5MG	N020835	002	Apr 14, 2000
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AB	+			30MG	N020835	001	Mar 27, 1998
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AB	+	!		35MG	N020835	003	May 25, 2002
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AB	+	!		150MG	N020835	005	Apr 22, 2008
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RISEDRONATE SODIUM

AB			APOTEX INC	35MG	A090877	001	Nov 30, 2015
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AB				75MG	A090877	002	Jun 10, 2014
-----------	--	--	--	-------------	----------------	------------	--------------

AB				150MG	A090877	003	Jun 10, 2014
-----------	--	--	--	--------------	----------------	------------	--------------

AB			AUROBINDO PHARMA LTD	5MG	A200296	001	Nov 30, 2015
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AB				30MG	A200296	002	Nov 30, 2015
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AB				35MG	A200296	003	Nov 30, 2015
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AB				150MG	A206768	001	Oct 21, 2016
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AB			MACLEODS PHARMS LTD	5MG	A203533	001	Dec 09, 2015
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PRESCRIPTION DRUG PRODUCT LIST

RISEDRONATE SODIUM

TABLET; ORAL

RISEDRONATE SODIUM

<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>	ORBION PHARMS	<u>30MG</u>	<u>A205280 001</u>	May 13, 2019
<u>AB</u>		<u>35MG</u>	<u>A205280 002</u>	May 13, 2019
<u>AB</u>	SUN PHARM	<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
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RISEDRONATE SODIUM

<u>AB</u>		SUN PHARM	<u>35MG</u>	<u>A203925 001</u>	Jul 09, 2019
<u>AB</u>		TEVA PHARMS USA	<u>35MG</u>	<u>A203217 001</u>	May 18, 2015

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
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RISPERIDONE

<u>AA</u>		AMNEAL PHARMS	<u>1MG/ML</u>	<u>A091384 001</u>	May 25, 2011
<u>AA</u>		AUROBINDO PHARMA LTD	<u>1MG/ML</u>	<u>A078452 001</u>	Sep 04, 2009
<u>AA</u>		HIKMA	<u>1MG/ML</u>	<u>A076904 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>		PHARM ASSOC	<u>1MG/ML</u>	<u>A077719 001</u>	Jul 29, 2009
<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

TABLET; ORAL

RISPERDAL

<u>AB</u>	+	JANSSEN PHARMS	<u>0.25MG</u>	<u>N020272 008</u>	May 10, 1999
<u>AB</u>	+		<u>0.5MG</u>	<u>N020272 007</u>	Jan 27, 1999
<u>AB</u>	+!		<u>1MG</u>	<u>N020272 001</u>	Dec 29, 1993
<u>AB</u>	+		<u>2MG</u>	<u>N020272 002</u>	Dec 29, 1993
<u>AB</u>	+		<u>3MG</u>	<u>N020272 003</u>	Dec 29, 1993
<u>AB</u>	+		<u>4MG</u>	<u>N020272 004</u>	Dec 29, 1993

RISPERIDONE

<u>AB</u>		AJANTA PHARMA LTD	<u>0.25MG</u>	<u>A201003 001</u>	Aug 24, 2011
<u>AB</u>			<u>0.5MG</u>	<u>A201003 002</u>	Aug 24, 2011
<u>AB</u>			<u>1MG</u>	<u>A201003 003</u>	Aug 24, 2011
<u>AB</u>			<u>2MG</u>	<u>A201003 004</u>	Aug 24, 2011
<u>AB</u>			<u>3MG</u>	<u>A201003 005</u>	Aug 24, 2011
<u>AB</u>			<u>4MG</u>	<u>A201003 006</u>	Aug 24, 2011
<u>AB</u>		AMNEAL	<u>0.25MG</u>	<u>A078071 001</u>	Jun 17, 2009
<u>AB</u>			<u>0.5MG</u>	<u>A078071 002</u>	Jun 17, 2009
<u>AB</u>			<u>1MG</u>	<u>A078071 003</u>	Jun 17, 2009
<u>AB</u>			<u>2MG</u>	<u>A078071 004</u>	Jun 17, 2009
<u>AB</u>			<u>3MG</u>	<u>A078071 005</u>	Jun 17, 2009
<u>AB</u>			<u>4MG</u>	<u>A078071 006</u>	Jun 17, 2009
<u>AB</u>		APOTEX INC	<u>0.25MG</u>	<u>A077953 001</u>	Sep 15, 2008
<u>AB</u>			<u>0.5MG</u>	<u>A077953 002</u>	Sep 15, 2008
<u>AB</u>			<u>1MG</u>	<u>A077953 003</u>	Sep 15, 2008
<u>AB</u>			<u>2MG</u>	<u>A077953 004</u>	Sep 15, 2008
<u>AB</u>			<u>3MG</u>	<u>A077953 005</u>	Sep 15, 2008
<u>AB</u>			<u>4MG</u>	<u>A077953 006</u>	Sep 15, 2008

PRESCRIPTION DRUG PRODUCT LIST

RISPERIDONE

TABLET; ORAL

RISPERIDONE

<u>AB</u>	CELLTRION	<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>	CHARTWELL MOLECULAR	<u>0.25MG</u>	<u>A077543 001</u>	May 18, 2011
<u>AB</u>		<u>0.5MG</u>	<u>A077543 002</u>	May 18, 2011
<u>AB</u>		<u>1MG</u>	<u>A077543 003</u>	May 18, 2011
<u>AB</u>		<u>2MG</u>	<u>A077543 004</u>	May 18, 2011
<u>AB</u>		<u>3MG</u>	<u>A077543 005</u>	May 18, 2011
<u>AB</u>		<u>4MG</u>	<u>A077543 006</u>	May 18, 2011
<u>AB</u>	DR REDDYS LABS LTD	<u>0.25MG</u>	<u>A076879 001</u>	Oct 24, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A076879 002</u>	Oct 24, 2008
<u>AB</u>		<u>1MG</u>	<u>A076879 003</u>	Oct 24, 2008
<u>AB</u>		<u>2MG</u>	<u>A076879 004</u>	Oct 24, 2008
<u>AB</u>		<u>3MG</u>	<u>A076879 005</u>	Oct 24, 2008
<u>AB</u>		<u>4MG</u>	<u>A076879 006</u>	Oct 24, 2008
<u>AB</u>	PRINSTON INC	<u>0.25MG</u>	<u>A077493 001</u>	Nov 29, 2011
<u>AB</u>		<u>0.5MG</u>	<u>A077493 002</u>	Nov 29, 2011
<u>AB</u>		<u>1MG</u>	<u>A077493 003</u>	Nov 29, 2011
<u>AB</u>		<u>2MG</u>	<u>A077493 004</u>	Nov 29, 2011
<u>AB</u>		<u>3MG</u>	<u>A077493 005</u>	Nov 29, 2011
<u>AB</u>		<u>4MG</u>	<u>A077493 006</u>	Nov 29, 2011
<u>AB</u>	RENATA	<u>0.25MG</u>	<u>A078707 001</u>	Dec 29, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A078707 002</u>	Dec 29, 2008
<u>AB</u>		<u>1MG</u>	<u>A078707 003</u>	Dec 29, 2008
<u>AB</u>		<u>2MG</u>	<u>A078707 004</u>	Dec 29, 2008
<u>AB</u>		<u>3MG</u>	<u>A078707 005</u>	Dec 29, 2008
<u>AB</u>		<u>4MG</u>	<u>A078707 006</u>	Dec 29, 2008
<u>AB</u>	RISING	<u>0.25MG</u>	<u>A078269 001</u>	Oct 08, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A078269 002</u>	Oct 08, 2008
<u>AB</u>		<u>1MG</u>	<u>A078269 003</u>	Oct 08, 2008
<u>AB</u>		<u>2MG</u>	<u>A078269 004</u>	Oct 08, 2008
<u>AB</u>		<u>3MG</u>	<u>A078269 005</u>	Oct 08, 2008
<u>AB</u>		<u>4MG</u>	<u>A078269 006</u>	Oct 08, 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>	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>	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

RISPERIDONE

<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

PRESCRIPTION DRUG PRODUCT LIST

RISPERIDONE

TABLET, ORALLY DISINTEGRATING;ORAL

RISPERIDONE

<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
<u>AB</u>		<u>4MG</u>	<u>A078116 005</u>	Dec 22, 2009
<u>AB</u>	SUN PHARM INDS LTD	<u>0.5MG</u>	<u>A077542 001</u>	Aug 06, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A078464 001</u>	Apr 08, 2013
<u>AB</u>		<u>1MG</u>	<u>A077542 002</u>	Aug 06, 2010
<u>AB</u>		<u>1MG</u>	<u>A078464 002</u>	Apr 08, 2013
<u>AB</u>		<u>2MG</u>	<u>A077542 003</u>	Aug 06, 2010
<u>AB</u>		<u>2MG</u>	<u>A078464 003</u>	Apr 08, 2013
<u>AB</u>		<u>3MG</u>	<u>A078464 004</u>	Apr 08, 2013
<u>AB</u>		<u>3MG</u>	<u>A078474 001</u>	Aug 06, 2010
<u>AB</u>		<u>4MG</u>	<u>A078464 005</u>	Apr 08, 2013
<u>AB</u>		<u>4MG</u>	<u>A078474 002</u>	Aug 06, 2010
<u>AB</u>	ZYDUS PHARMS USA	<u>0.5MG</u>	<u>A078516 001</u>	May 01, 2009
<u>AB</u>		<u>2MG</u>	<u>A078516 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

NORVIR

<u>AB</u>	+! ABBVIE	<u>100MG</u>	<u>N022417 001</u>	Feb 10, 2010
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RITONAVIR

<u>AB</u>	AMNEAL	<u>100MG</u>	<u>A208890 001</u>	Sep 17, 2018
<u>AB</u>	AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A206614 001</u>	Sep 17, 2018
<u>AB</u>	HETERO LABS LTD III	<u>100MG</u>	<u>A204587 001</u>	Sep 17, 2018
<u>AB</u>	HIKMA	<u>100MG</u>	<u>A202573 001</u>	Jan 15, 2015

RIVAROXABAN

FOR SUSPENSION;ORAL

XARELTO

+! JANSSEN PHARMS

1MG/ML

N215859 001 Dec 20, 2021

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

EXELON

<u>AB</u>	+ NOVARTIS	<u>4.6MG/24HR</u>	<u>N022083 001</u>	Jul 06, 2007
<u>AB</u>	+!	<u>9.5MG/24HR</u>	<u>N022083 002</u>	Jul 06, 2007
<u>AB</u>	+	<u>13.3MG/24HR</u>	<u>N022083 005</u>	Aug 31, 2012

RIVASTIGMINE

<u>AB</u>	ALVOGEN	<u>4.6MG/24HR</u>	<u>A204403 001</u>	Sep 03, 2015
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A204403 002</u>	Sep 03, 2015
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A204403 003</u>	Aug 31, 2015
<u>AB</u>	AMNEAL PHARMS	<u>4.6MG/24HR</u>	<u>A207308 001</u>	Jan 08, 2019
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A207308 002</u>	Jan 08, 2019
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A207308 003</u>	Jan 08, 2019
<u>AB</u>	BRECKENRIDGE	<u>4.6MG/24HR</u>	<u>A209063 001</u>	Nov 26, 2019
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A209063 002</u>	Nov 26, 2019
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A209063 003</u>	Nov 26, 2019
<u>AB</u>	MYLAN TECHNOLOGIES	<u>4.6MG/24HR</u>	<u>A205622 001</u>	Jun 20, 2018
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A205622 002</u>	Jun 20, 2018
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A205622 003</u>	Jun 20, 2018
<u>AB</u>	ZYDUS NOVELTECH INC	<u>4.6MG/24HR</u>	<u>A206318 001</u>	Mar 04, 2019
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A206318 002</u>	Mar 04, 2019
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A206318 003</u>	Mar 04, 2019

PRESCRIPTION DRUG PRODUCT LIST

RIVASTIGMINE TARTRATE

CAPSULE;ORAL

RIVASTIGMINE TARTRATE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 1.5MG BASE</u>	<u>A091689 001</u>	Jun 12, 2012
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A091689 002</u>	Jun 12, 2012
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A091689 003</u>	Jun 12, 2012
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A091689 004</u>	Jun 12, 2012
<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>	ORBION PHARMS	<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	<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
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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>	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>	PHARMA LIFE	<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>	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
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077263 002</u>	Dec 31, 2012

PRESCRIPTION DRUG PRODUCT LIST

RIZATRIPTAN BENZOATE

TABLET, ORALLY DISINTEGRATING;ORAL

MAXALT-MLT

AB	+ !	MERCK	EQ 10MG BASE	N020865 002	Jun 29, 1998
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RIZATRIPTAN BENZOATE

AB		AUROBINDO PHARMA LTD	EQ 5MG BASE	A203062 001	Jul 01, 2013
AB			EQ 10MG BASE	A203062 002	Jul 01, 2013
AB		GLENMARK PHARMS LTD	EQ 5MG BASE	A201914 001	Jul 01, 2013
AB			EQ 10MG BASE	A201914 002	Jul 01, 2013
AB		MACLEODS PHARMS LTD	EQ 5MG BASE	A203146 001	Sep 19, 2014
AB			EQ 10MG BASE	A203146 002	Sep 19, 2014
AB		NATCO PHARMA LTD	EQ 5MG BASE	A203478 001	Jul 01, 2013
AB			EQ 10MG BASE	A203478 002	Jul 01, 2013
AB		PANACEA	EQ 5MG BASE	A204722 001	Jan 11, 2017
AB			EQ 10MG BASE	A204722 002	Jan 11, 2017
AB		SANDOZ	EQ 5MG BASE	A078739 001	Jul 01, 2013
AB			EQ 10MG BASE	A078739 002	Jul 01, 2013
AB		UNICHEM	EQ 5MG BASE	A207835 001	Mar 07, 2017
AB			EQ 10MG BASE	A207835 002	Mar 07, 2017

ROCURONIUM BROMIDE

INJECTABLE; INJECTION

ROCURONIUM BROMIDE

AP		EUGIA PHARMA	50MG/5ML (10MG/ML)	A206206 001	Apr 12, 2017
AP			100MG/10ML (10MG/ML)	A206206 002	Apr 12, 2017
AP		FRESENIUS KABI USA	50MG/5ML (10MG/ML)	A078651 001	Dec 29, 2008
AP			100MG/10ML (10MG/ML)	A078651 002	Dec 29, 2008
AP		GLAND PHARMA LTD	50MG/5ML (10MG/ML)	A205656 001	Apr 04, 2018
AP			100MG/10ML (10MG/ML)	A205656 002	Apr 04, 2018
AP		HOSPIRA	50MG/5ML (10MG/ML)	A078519 001	Nov 26, 2008
AP			100MG/10ML (10MG/ML)	A078519 002	Nov 26, 2008
AP		MYLAN INSTITUTIONAL	50MG/5ML (10MG/ML)	A079199 001	Nov 26, 2008
AP			100MG/10ML (10MG/ML)	A079199 002	Nov 26, 2008
AP		PIRAMAL CRITICAL	50MG/5ML (10MG/ML)	A210437 001	Aug 13, 2019
AP			100MG/10ML (10MG/ML)	A210437 002	Aug 13, 2019
AP		SAGENT PHARMS INC	50MG/5ML (10MG/ML)	A091458 001	Jul 28, 2010
AP			100MG/10ML (10MG/ML)	A091458 002	Jul 28, 2010
AP	!	SANDOZ INC	50MG/5ML (10MG/ML)	A079195 001	Dec 05, 2008
AP	!		100MG/10ML (10MG/ML)	A079195 002	Dec 05, 2008
AP		TAMARANG	50MG/5ML (10MG/ML)	A091115 001	Aug 27, 2012
AP			100MG/10ML (10MG/ML)	A091115 002	Aug 27, 2012
AP		WEST WARD PHARM CORP	50MG/5ML (10MG/ML)	A204679 001	Feb 28, 2017
AP			100MG/10ML (10MG/ML)	A204679 002	Feb 28, 2017

ROFLUMILAST

TABLET; ORAL

DALIRESP

AB	+ !	ASTRAZENECA	500MCG	N022522 001	Feb 28, 2011
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ROFLUMILAST

AB		HETERO LABS LTD III	500MCG	A208213 001	Nov 23, 2018
AB		STRIDES PHARMA	500MCG	A208247 001	Mar 30, 2020
AB		TORRENT	500MCG	A208272 001	Aug 06, 2018
		DALIRESP			
	+	ASTRAZENECA	250MCG	N022522 002	Jan 23, 2018

ROLAPITANT HYDROCHLORIDE

TABLET; ORAL

VARUBI

	+ !	TERSERA	EQ 90MG BASE	N206500 001	Sep 01, 2015
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ROMIDEPSIN

POWDER; INTRAVENOUS

ISTODAX

AP	+ !	CELGENE	10MG/VIAL	N022393 001	Nov 05, 2009
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ROMIDEPSIN

AP		FRESENIUS KABI USA	10MG/VIAL	A206254 001	Oct 12, 2021
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SOLUTION; INTRAVENOUS

ROMIDEPSIN

	+ !	TEVA PHARMS USA INC	27.5MG/5.5ML (5MG/ML)	N208574 002	Mar 13, 2020
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PRESCRIPTION DRUG PRODUCT LIST

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

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>	CADILA	<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
<u>AB</u>	CELLTRION	<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>	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>	MLV	<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>	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>	ORBION PHARMS	<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

TABLET, EXTENDED RELEASE; ORAL

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

PRESCRIPTION DRUG PRODUCT LIST

ROPINIROLE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ROPINIROLE HYDROCHLORIDE

<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>	CELLTRION	<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
<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

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
<u>AP</u>			<u>400MG/200ML (2MG/ML)</u>	<u>A204636 002</u>	Mar 16, 2018
<u>AP</u>			<u>150MG/30ML (5MG/ML)</u>	<u>A203955 001</u>	Apr 11, 2016
<u>AP</u>		CAPLIN	<u>40MG/20ML (2MG/ML)</u>	<u>A212808 001</u>	Apr 09, 2020
<u>AP</u>			<u>100MG/20ML (5MG/ML)</u>	<u>A212808 002</u>	Apr 09, 2020
<u>AP</u>			<u>150MG/30ML (5MG/ML)</u>	<u>A212808 003</u>	Apr 09, 2020
<u>AP</u>			<u>200MG/20ML (10MG/ML)</u>	<u>A212808 004</u>	Apr 09, 2020
<u>AP</u>		EUGIA PHARMA	<u>40MG/20ML (2MG/ML)</u>	<u>A205612 001</u>	Jul 13, 2016
<u>AP</u>			<u>100MG/20ML (5MG/ML)</u>	<u>A205612 003</u>	Jul 13, 2016
<u>AP</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A205612 006</u>	Jul 13, 2016
<u>AP</u>			<u>150MG/30ML (5MG/ML)</u>	<u>A205612 004</u>	Jul 13, 2016
<u>AP</u>			<u>150MG/20ML (7.5MG/ML)</u>	<u>A205612 005</u>	Jul 13, 2016
<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A205612 002</u>	Jul 13, 2016
<u>AP</u>			<u>200MG/20ML (10MG/ML)</u>	<u>A205612 007</u>	Jul 13, 2016
<u>AP</u>		HIKMA	<u>40MG/20ML (2MG/ML)</u>	<u>A214074 001</u>	Jul 20, 2020
<u>AP</u>			<u>150MG/30ML (5MG/ML)</u>	<u>A214074 002</u>	Jul 20, 2020
<u>AP</u>			<u>150MG/20ML (7.5MG/ML)</u>	<u>A214074 003</u>	Jul 20, 2020
<u>AP</u>			<u>200MG/20ML (10MG/ML)</u>	<u>A214074 004</u>	Jul 20, 2020
<u>AP</u>		HOSPIRA	<u>20MG/10ML (2MG/ML)</u>	<u>A090194 001</u>	Sep 23, 2014
<u>AP</u>			<u>40MG/20ML (2MG/ML)</u>	<u>A090194 005</u>	Sep 23, 2014
<u>AP</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A090194 004</u>	Sep 23, 2014
<u>AP</u>			<u>150MG/30ML (5MG/ML)</u>	<u>A090194 002</u>	Sep 23, 2014
<u>AP</u>			<u>150MG/20ML (7.5MG/ML)</u>	<u>A090194 003</u>	Sep 23, 2014
<u>AP</u>			<u>200MG/20ML (10MG/ML)</u>	<u>A090194 006</u>	Sep 23, 2014
<u>AP</u>		INFORLIFE	<u>200MG/100ML (2MG/ML)</u>	<u>A206166 001</u>	Jun 11, 2018
<u>AP</u>			<u>400MG/200ML (2MG/ML)</u>	<u>A206166 002</u>	Jun 11, 2018
<u>AP</u>			<u>500MG/100ML (5MG/ML)</u>	<u>A206166 003</u>	Jun 11, 2018
<u>AP</u>			<u>1GM/200ML (5MG/ML)</u>	<u>A206166 004</u>	Jun 11, 2018
<u>AP</u>		NAVINTA LLC	<u>150MG/30ML (5MG/ML)</u>	<u>A078601 002</u>	Jul 17, 2014
<u>AP</u>			<u>200MG/20ML (10MG/ML)</u>	<u>A078601 003</u>	Jul 17, 2014
<u>AP</u>		SOMERSET THERAPS LLC	<u>20MG/10ML (2MG/ML)</u>	<u>A207636 001</u>	Jun 15, 2018
<u>AP</u>			<u>40MG/20ML (2MG/ML)</u>	<u>A207636 002</u>	Jun 15, 2018

PRESCRIPTION DRUG PRODUCT LIST

ROPIVACAINE HYDROCHLORIDE

SOLUTION; INJECTION

ROPIVACAINE HYDROCHLORIDE

<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

+ WOODWARD

EQ 2MG BASE

N021071 002

May 25, 1999

+

EQ 4MG BASE

N021071 003

May 25, 1999

ROSUVASTATIN CALCIUM

CAPSULE; ORAL

EZALLOR SPRINKLE

+ SUN PHARM

EQ 5MG BASE

N208647 001

Dec 18, 2018

+

EQ 10MG BASE

N208647 002

Dec 18, 2018

+

EQ 20MG BASE

N208647 003

Dec 18, 2018

+!

EQ 40MG BASE

N208647 004

Dec 18, 2018

TABLET; ORAL

CRESTOR

<u>AB</u>	+	IPR	<u>EQ 5MG BASE</u>	<u>N021366</u>	<u>002</u>	Aug 12, 2003
<u>AB</u>	+		<u>EQ 10MG BASE</u>	<u>N021366</u>	<u>003</u>	Aug 12, 2003
<u>AB</u>	+		<u>EQ 20MG BASE</u>	<u>N021366</u>	<u>004</u>	Aug 12, 2003
<u>AB</u>	+!		<u>EQ 40MG BASE</u>	<u>N021366</u>	<u>005</u>	Aug 12, 2003

ROSUVASTATIN CALCIUM

<u>AB</u>		ACCORD HLTHCARE	<u>EQ 5MG BASE</u>	<u>A206434</u>	<u>001</u>	Oct 31, 2016
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A206434</u>	<u>002</u>	Oct 31, 2016
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A206434</u>	<u>003</u>	Oct 31, 2016
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A206434</u>	<u>004</u>	Oct 31, 2016
<u>AB</u>		ALKEM LABS LTD	<u>EQ 5MG BASE</u>	<u>A206465</u>	<u>001</u>	Mar 21, 2017
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A206465</u>	<u>002</u>	Mar 21, 2017
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A206465</u>	<u>003</u>	Mar 21, 2017
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A206465</u>	<u>004</u>	Mar 21, 2017
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A079170</u>	<u>001</u>	Jul 19, 2016
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A079170</u>	<u>002</u>	Jul 19, 2016
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A079170</u>	<u>003</u>	Jul 19, 2016
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A079170</u>	<u>004</u>	Jul 19, 2016
<u>AB</u>		BIOCON PHARMA	<u>EQ 5MG BASE</u>	<u>A207752</u>	<u>001</u>	Oct 31, 2016
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A207752</u>	<u>002</u>	Oct 31, 2016
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A207752</u>	<u>003</u>	Oct 31, 2016
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A207752</u>	<u>004</u>	Oct 31, 2016
<u>AB</u>		CADILA PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A207453</u>	<u>001</u>	Nov 23, 2016
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A207453</u>	<u>002</u>	Nov 23, 2016
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A207453</u>	<u>003</u>	Nov 23, 2016
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A207453</u>	<u>004</u>	Nov 23, 2016
<u>AB</u>		CHANGZHOU PHARM	<u>EQ 5MG BASE</u>	<u>A207408</u>	<u>001</u>	Oct 31, 2016
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A207408</u>	<u>002</u>	Oct 31, 2016
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A207408</u>	<u>003</u>	Oct 31, 2016
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A207408</u>	<u>004</u>	Oct 31, 2016
<u>AB</u>		CHARTWELL RX	<u>EQ 5MG BASE</u>	<u>A079168</u>	<u>001</u>	Jul 19, 2016
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A079168</u>	<u>002</u>	Jul 19, 2016
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A079168</u>	<u>003</u>	Jul 19, 2016
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A079168</u>	<u>004</u>	Jul 19, 2016
<u>AB</u>		GLENMARK PHARMS	<u>EQ 5MG BASE</u>	<u>A079172</u>	<u>001</u>	Jul 19, 2016
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A079172</u>	<u>002</u>	Jul 19, 2016
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A079172</u>	<u>003</u>	Jul 19, 2016
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A079172</u>	<u>004</u>	Jul 19, 2016
<u>AB</u>		HETERO LABS LTD V	<u>EQ 5MG BASE</u>	<u>A207616</u>	<u>001</u>	Oct 31, 2016
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A207616</u>	<u>002</u>	Oct 31, 2016
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A207616</u>	<u>003</u>	Oct 31, 2016
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A207616</u>	<u>004</u>	Oct 31, 2016
<u>AB</u>		LUPIN	<u>EQ 5MG BASE</u>	<u>A205587</u>	<u>001</u>	Jul 31, 2017
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A205587</u>	<u>002</u>	Jul 31, 2017
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A205587</u>	<u>003</u>	Jul 31, 2017
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A205587</u>	<u>004</u>	Jul 31, 2017
<u>AB</u>		MSN	<u>EQ 5MG BASE</u>	<u>A208898</u>	<u>001</u>	Nov 22, 2017
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A208898</u>	<u>002</u>	Nov 22, 2017
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A208898</u>	<u>003</u>	Nov 22, 2017
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A208898</u>	<u>004</u>	Nov 22, 2017

PRESCRIPTION DRUG PRODUCT LIST

ROSUVASTATIN CALCIUM

TABLET; ORAL

ROSUVASTATIN CALCIUM

<u>AB</u>	RENATA	<u>EQ 5MG BASE</u>	<u>A207062 001</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207062 002</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207062 003</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207062 004</u>	Oct 31, 2016
<u>AB</u>	SANDOZ INC	<u>EQ 5MG BASE</u>	<u>A079171 001</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079171 002</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A079171 003</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079171 004</u>	Jul 19, 2016
<u>AB</u>	SHANDONG	<u>EQ 5MG BASE</u>	<u>A207375 001</u>	May 07, 2019
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207375 002</u>	May 07, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207375 003</u>	May 07, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207375 004</u>	May 07, 2019
<u>AB</u>	SUN PHARM	<u>EQ 5MG BASE</u>	<u>A079169 001</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079169 002</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A079169 003</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079169 004</u>	Jul 19, 2016
<u>AB</u>	TORRENT	<u>EQ 5MG BASE</u>	<u>A201619 001</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A201619 002</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A201619 003</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A201619 004</u>	Oct 31, 2016
<u>AB</u>	WATSON LABS INC	<u>EQ 5MG BASE</u>	<u>A079167 001</u>	Apr 29, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079167 002</u>	Apr 29, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A079167 003</u>	Apr 29, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079167 004</u>	Apr 29, 2016
<u>AB</u>	ZHEJIANG YONGTAI	<u>EQ 5MG BASE</u>	<u>A212059 001</u>	Nov 04, 2019
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A212059 002</u>	Nov 04, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A212059 003</u>	Nov 04, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A212059 004</u>	Nov 04, 2019

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 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

BANZELAB +! EISAI INC 40MG/ML N201367 001 Mar 03, 2011RUFINAMIDE

<u>AB</u>	ALKEM LABS LTD	<u>40MG/ML</u>	<u>A213410 001</u>	Feb 23, 2021
<u>AB</u>	BIONPHARMA INC	<u>40MG/ML</u>	<u>A211388 001</u>	Apr 23, 2019
<u>AB</u>	HIKMA	<u>40MG/ML</u>	<u>A207363 001</u>	Apr 23, 2019
<u>AB</u>	LUPIN LTD	<u>40MG/ML</u>	<u>A213457 001</u>	Dec 18, 2020

TABLET; ORAL

BANZEL

<u>AB</u>	+	EISAI INC	<u>200MG</u>	<u>N021911 002</u>	Nov 14, 2008
<u>AB</u>	+	!	<u>400MG</u>	<u>N021911 003</u>	Nov 14, 2008

RUFINAMIDE

<u>AB</u>	GLENMARK PHARMS LTD	<u>200MG</u>	<u>A205075 001</u>	May 16, 2016
<u>AB</u>		<u>400MG</u>	<u>A205075 002</u>	May 16, 2016
<u>AB</u>	HETERO LABS LTD III	<u>200MG</u>	<u>A204993 001</u>	May 11, 2021

PRESCRIPTION DRUG PRODUCT LIST

RUFINAMIDE

TABLET; ORAL

RUFINAMIDE

AB		400MG	A204993 002	May 11, 2021
AB	HIKMA	200MG	A204988 001	May 16, 2016
AB		400MG	A204988 002	May 16, 2016
AB	MYLAN	200MG	A205095 001	May 16, 2016
AB		400MG	A205095 002	May 16, 2016

RUXOLITINIB PHOSPHATE

CREAM; TOPICAL

OPZELURA

+! INCYTE CORP EQ 1.5% BASE N215309 001 Sep 21, 2021

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

SACUBITRIL; VALSARTAN

TABLET; ORAL

ENTRESTO

+ NOVARTIS PHARMS 24MG; 26MG N207620 001 Jul 07, 2015

+ CORP 49MG; 51MG N207620 002 Jul 07, 2015

+! 97MG; 103MG N207620 003 Jul 07, 2015

SAFINAMIDE MESYLATE

TABLET; ORAL

XADAGO

+ MDD US EQ 50MG BASE N207145 001 Mar 21, 2017

+! EQ 100MG BASE N207145 002 Mar 21, 2017

SALMETEROL XINAFOATE

POWDER; INHALATION

SEREVENT

+! GLAXOSMITHKLINE EQ 0.05MG BASE/INH N020692 001 Sep 19, 1997

SAMARIUM SM-153 LEXIDRONAM PENTASODIUM

INJECTABLE; INJECTION

QUADRAMET

+! LANTHEUS MEDICAL 50mCi/ML N020570 001 Mar 28, 1997

SAPROPTERIN DIHYDROCHLORIDE

POWDER; ORAL

KUVAN**AB** +! BIOMARIN PHARM **100MG/PACKET** **N205065 001** Dec 19, 2013**AB** + **500MG/PACKET** **N205065 002** Oct 27, 2015SAPROPTERIN DIHYDROCHLORIDE**AB** DR REDDYS **100MG/PACKET** **A209452 001** Mar 30, 2021**AB** PAR PHARM INC **100MG/PACKET** **A207207 001** Aug 20, 2019**AB** **500MG/PACKET** **A210027 001** Aug 20, 2019

TABLET; ORAL

KUVAN**AB** +! BIOMARIN PHARM **100MG** **N022181 001** Dec 13, 2007SAPROPTERIN DIHYDROCHLORIDE**AB** DR REDDYS **100MG** **A207685 001** Jun 15, 2021**AB** PAR PHARM INC **100MG** **A207200 001** May 10, 2019SARECYCLINE 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

PRESCRIPTION DRUG PRODUCT LIST

SCOPOLAMINE

SYSTEM; TRANSDERMAL

SCOPOLAMINE

AB	ACTAVIS LABS UT INC	1MG/72HR	A208769 001	Jan 10, 2022
AB	MYLAN TECHNOLOGIES	1MG/72HR	A203753 001	Jun 19, 2019
AB	PADAGIS US	1MG/72HR	A078830 001	Jan 30, 2015
AB	RICONPHARMA LLC	1MG/72HR	A212342 001	Nov 24, 2020

TRANSDERM SCOP

AB	+ !	BAXTER HLTHCARE CORP	1MG/72HR	N017874 001
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SECNIDAZOLE

GRANULE; ORAL

SOLOSEC

+ !	LUPIN	2GM/PACKET	N209363 001	Sep 15, 2017
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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		NOVITIUM PHARMA	5MG	A075352 001	Nov 30, 1998
AB		RISING	5MG	A206803 001	Apr 02, 2019

TABLET; ORAL

SELEGILINE HYDROCHLORIDE

AB	!	APOTEX INC	5MG	A074871 001	Jun 06, 1997
AB		BOSCOGEN	5MG	A074912 001	Apr 30, 1998
AB		I3 PHARMS	5MG	A074672 001	Apr 01, 1997

TABLET, ORALLY DISINTEGRATING; ORAL

ZELAPAR

+ !	BAUSCH	1.25MG	N021479 001	Jun 14, 2006
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SELENIOS ACID

SOLUTION; INTRAVENOUS

SELENIOS ACID

+ !	AM REGENT	EQ 12MCG SELENIUM/2ML (EQ 6MCG SELENIUM/ML)	N209379 003	Aug 30, 2021
+ !		EQ 60MCG SELENIUM/ML (EQ 60MCG SELENIUM/ML)	N209379 002	Jan 25, 2021
+ !		EQ 600MCG SELENIUM/10ML (EQ 60MCG SELENIUM/ML)	N209379 001	Apr 30, 2019

SELENIUM SULFIDE

LOTION; SHAMPOO; TOPICAL

SELENIUM SULFIDE

AT	!	PADAGIS US	2.5%	A089996 001	Jan 10, 1991
AT		WOCKHARDT BIO AG	2.5%	A088228 001	Sep 01, 1983

SELEXIPAG

POWDER; INTRAVENOUS

UPTRAVI

+ !	ACTELION	1.8MG/VIAL	N214275 001	Jul 29, 2021
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TABLET; ORAL

UPTRAVI

+	ACTELION	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

PRESCRIPTION DRUG PRODUCT LIST

SELINEXOR

TABLET; ORAL

XPOVIO

+	!	KARYOPHARM THERAPS	20MG	N212306	001	Jul 03, 2019
+			40MG	N212306	002	Apr 15, 2021
+			50MG	N212306	003	Apr 15, 2021
+			60MG	N212306	004	Apr 15, 2021

SELPERCATINIB

CAPSULE; ORAL

RETEVMO

+		LOXO ONCOLOGY INC	40MG	N213246	001	May 08, 2020
+	!		80MG	N213246	002	May 08, 2020

SELUMETINIB SULFATE

CAPSULE; ORAL

KOSELUGO

+		ASTRAZENECA	EQ 10MG BASE	N213756	001	Apr 10, 2020
+	!		EQ 25MG BASE	N213756	002	Apr 10, 2020

SEMAGLUTIDE

SOLUTION; SUBCUTANEOUS

OZEMPIC

+	!	NOVO	2MG/1.5ML (1.34MG/ML)	N209637	001	Dec 05, 2017
+	!		4MG/3ML (1.34MG/ML)	N209637	002	Apr 09, 2019

WEGOVY

+	!	NOVO	0.25MG/0.5ML (0.25MG/0.5ML)	N215256	001	Jun 04, 2021
+	!		0.5MG/0.5ML (0.5MG/0.5ML)	N215256	002	Jun 04, 2021
+	!		1MG/0.5ML (1MG/0.5ML)	N215256	003	Jun 04, 2021
+	!		1.7MG/0.75ML (1.7MG/0.75ML)	N215256	004	Jun 04, 2021
+	!		2.4MG/0.75ML (2.4MG/0.75ML)	N215256	005	Jun 04, 2021

TABLET; ORAL

RYBELSUS

+		NOVO	3MG	N213051	001	Sep 20, 2019
+			7MG	N213051	002	Sep 20, 2019
+	!		14MG	N213051	003	Sep 20, 2019

SERTACONAZOLE NITRATE

CREAM; TOPICAL

ERTACZO

+	!	BAUSCH	2%	N021385	001	Dec 10, 2003
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SERTRALINE HYDROCHLORIDE

CAPSULE; ORAL

SERTRALINE HYDROCHLORIDE

+		ALMATICA	EQ 150MG BASE	N215133	001	Oct 04, 2021
+	!		EQ 200MG BASE	N215133	002	Oct 04, 2021

CONCENTRATE; ORAL

SERTRALINE HYDROCHLORIDE

AA		ACI	<u>EQ 20MG BASE/ML</u>	<u>A076934</u>	<u>001</u>	Jun 30, 2006
AA		AUROBINDO PHARMA	<u>EQ 20MG BASE/ML</u>	<u>A078861</u>	<u>001</u>	Oct 31, 2008

ZOLOFT

AA	+	!	UPJOHN	<u>EQ 20MG BASE/ML</u>	<u>N020990</u>	<u>001</u>	Dec 07, 1999
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TABLET; ORAL

SERTRALINE HYDROCHLORIDE

AB		ACCORD HLTHCARE	<u>EQ 25MG BASE</u>	<u>A202825</u>	<u>001</u>	Nov 07, 2014
AB			<u>EQ 50MG BASE</u>	<u>A202825</u>	<u>002</u>	Nov 07, 2014
AB			<u>EQ 100MG BASE</u>	<u>A202825</u>	<u>003</u>	Nov 07, 2014
AB		ACI	<u>EQ 25MG BASE</u>	<u>A076881</u>	<u>001</u>	Feb 06, 2007
AB			<u>EQ 50MG BASE</u>	<u>A076881</u>	<u>002</u>	Feb 06, 2007
AB			<u>EQ 100MG BASE</u>	<u>A076881</u>	<u>003</u>	Feb 06, 2007
AB		ASCENT PHARMS INC	<u>EQ 25MG BASE</u>	<u>A214790</u>	<u>001</u>	May 03, 2021
AB			<u>EQ 50MG BASE</u>	<u>A214790</u>	<u>002</u>	May 03, 2021
AB			<u>EQ 100MG BASE</u>	<u>A214790</u>	<u>003</u>	May 03, 2021
AB		AUROBINDO PHARMA	<u>EQ 25MG BASE</u>	<u>A077206</u>	<u>001</u>	Feb 06, 2007
AB			<u>EQ 50MG BASE</u>	<u>A077206</u>	<u>002</u>	Feb 06, 2007
AB			<u>EQ 100MG BASE</u>	<u>A077206</u>	<u>003</u>	Feb 06, 2007
AB		GRANULES	<u>EQ 25MG BASE</u>	<u>A078403</u>	<u>001</u>	Jan 08, 2008
AB			<u>EQ 50MG BASE</u>	<u>A078403</u>	<u>002</u>	Jan 08, 2008
AB			<u>EQ 100MG BASE</u>	<u>A078403</u>	<u>003</u>	Jan 08, 2008
AB		INVAGEN PHARMS	<u>EQ 25MG BASE</u>	<u>A077397</u>	<u>001</u>	Feb 06, 2007
AB			<u>EQ 50MG BASE</u>	<u>A077397</u>	<u>002</u>	Feb 06, 2007
AB			<u>EQ 100MG BASE</u>	<u>A077397</u>	<u>003</u>	Feb 06, 2007
AB		LUPIN	<u>EQ 25MG BASE</u>	<u>A077670</u>	<u>001</u>	Feb 06, 2007
AB			<u>EQ 50MG BASE</u>	<u>A077670</u>	<u>002</u>	Feb 06, 2007

PRESCRIPTION DRUG PRODUCT LIST

SERTRALINE HYDROCHLORIDE

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077670 003</u>	Feb 06, 2007
<u>AB</u>	OXFORD PHARMS	<u>EQ 25MG BASE</u>	<u>A078175 001</u>	Jul 21, 2010
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078175 002</u>	Jul 21, 2010
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078175 003</u>	Jul 21, 2010
<u>AB</u>	REYOUNG	<u>EQ 25MG BASE</u>	<u>A078677 001</u>	Mar 04, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078677 002</u>	Mar 04, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078677 003</u>	Mar 04, 2009
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 25MG BASE</u>	<u>A076442 001</u>	Apr 30, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076442 002</u>	Apr 30, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076442 003</u>	Apr 30, 2007
<u>AB</u>	VIWIT PHARM	<u>EQ 25MG BASE</u>	<u>A076882 001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076882 002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076882 003</u>	Feb 06, 2007

ZOLOFT

<u>AB</u>	+	UPJOHN	<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

SETMELANOTIDE ACETATE

SOLUTION; SUBCUTANEOUS

IMCIVREE

+	!	RHYTHM	EQ 10MG/ML BASE (EQ 10MG/ML BASE)	N213793 001	Nov 25, 2020
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SEVELAMER CARBONATE

FOR SUSPENSION; ORAL

REVELA

<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>		BIONPHARMA INC	<u>800MG/PACKET</u>	<u>A209374 001</u>	May 17, 2021
<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
<u>AB</u>		IMPAX	<u>800MG/PACKET</u>	<u>A211316 001</u>	Nov 20, 2020
<u>AB</u>			<u>2.4GM/PACKET</u>	<u>A211316 002</u>	Nov 20, 2020
<u>AB</u>		LUPIN LTD	<u>800MG/PACKET</u>	<u>A201513 001</u>	Dec 23, 2021
<u>AB</u>			<u>2.4GM/PACKET</u>	<u>A201513 002</u>	Dec 23, 2021

TABLET; ORAL

REVELA

<u>AB</u>	+	SANOFI	<u>800MG</u>	<u>N022127 001</u>	Oct 19, 2007
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SEVELAMER CARBONATE

<u>AB</u>		AMNEAL PHARMS CO	<u>800MG</u>	<u>A207288 001</u>	Nov 28, 2017
<u>AB</u>		ANXIN	<u>800MG</u>	<u>A212970 001</u>	Apr 09, 2020
<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>		INVAGEN PHARMS	<u>800MG</u>	<u>A203860 001</u>	Oct 26, 2017
<u>AB</u>		LUPIN LTD	<u>800MG</u>	<u>A204600 001</u>	Jan 14, 2021
<u>AB</u>		RISING	<u>800MG</u>	<u>A204451 001</u>	Nov 29, 2018
<u>AB</u>		TWI PHARMS	<u>800MG</u>	<u>A200959 001</u>	Mar 20, 2018
<u>AB</u>		ZYDUS PHARMS	<u>800MG</u>	<u>A207759 001</u>	Aug 11, 2020

SEVELAMER HYDROCHLORIDE

TABLET; ORAL

RENAGEL

<u>AB</u>	+	GENZYME	<u>400MG</u>	<u>N021179 001</u>	Jul 12, 2000
<u>AB</u>	+	!	<u>800MG</u>	<u>N021179 002</u>	Jul 12, 2000

SEVELAMER HYDROCHLORIDE

<u>AB</u>		GLENMARK PHARMS LTD	<u>400MG</u>	<u>A204724 001</u>	Feb 08, 2019
<u>AB</u>			<u>800MG</u>	<u>A204724 002</u>	Feb 08, 2019
<u>AB</u>		LUPIN LTD	<u>400MG</u>	<u>A213145 001</u>	Jun 16, 2021
<u>AB</u>			<u>800MG</u>	<u>A213145 002</u>	Jun 16, 2021

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

PRESCRIPTION DRUG PRODUCT LIST

SEVOFLURANE

LIQUID; INHALATION

SOJOURN

AN	PIRAMAL CRITICAL	100%	A077867 001	May 02, 2007
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ULTANE

AN	+! ABEVIE	100%	N020478 001	Jun 07, 1995
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SILDENAFIL CITRATE

FOR SUSPENSION; ORAL

REVATIO

AB	+! UPJOHN	EQ 10MG BASE/ML	N203109 001	Aug 30, 2012
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SILDENAFIL CITRATE

AB	AJANTA PHARMA LTD	EQ 10MG BASE/ML	A212883 001	Nov 27, 2019
AB	ALKEM LABS LTD	EQ 10MG BASE/ML	A212440 001	Nov 27, 2019
AB	AMNEAL PHARMS	EQ 10MG BASE/ML	A211092 001	Nov 27, 2019
AB	APPCO	EQ 10MG BASE/ML	A213814 001	Apr 29, 2021
AB	APTAPHARMA INC	EQ 10MG BASE/ML	A213041 001	Aug 06, 2020
AB	HETERO LABS LTD V	EQ 10MG BASE/ML	A213014 001	May 11, 2021
AB	NOVITIUM PHARMA	EQ 10MG BASE/ML	A212260 001	May 31, 2019
AB	TARO PHARM INDS LTD	EQ 10MG BASE/ML	A215522 001	Nov 16, 2021
AB	TEVA PHARMS USA	EQ 10MG BASE/ML	A211534 001	May 29, 2020
AB	TRIS PHARMA INC	EQ 10MG BASE/ML	A212312 001	Nov 17, 2021

SOLUTION; INTRAVENOUS

REVATIO

AP	+! UPJOHN	EQ 10MG BASE/12.5ML (EQ 0.8MG BASE/ML)	N022473 001	Nov 18, 2009
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SILDENAFIL CITRATE

AP	EUGIA PHARMA	EQ 10MG BASE/12.5ML (EQ 0.8MG BASE/ML)	A203988 001	Apr 01, 2015
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TABLET; ORAL

REVATIO

AB	+! UPJOHN	EQ 20MG BASE	N021845 001	Jun 03, 2005
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SILDENAFIL CITRATE

AB	AJANTA PHARMA LTD	EQ 20MG BASE	A210394 001	May 04, 2018
AB		EQ 25MG BASE	A206401 001	Oct 12, 2018
AB		EQ 50MG BASE	A206401 002	Oct 12, 2018
AB		EQ 100MG BASE	A206401 003	Oct 12, 2018
AB	AMNEAL PHARMS	EQ 20MG BASE	A202025 001	Feb 28, 2013
AB	AMNEAL PHARMS NY	EQ 25MG BASE	A202023 001	Jun 27, 2018
AB		EQ 50MG BASE	A202023 002	Jun 27, 2018
AB		EQ 100MG BASE	A202023 003	Jun 27, 2018
AB	APPCO	EQ 25MG BASE	A207178 001	Mar 02, 2020
AB		EQ 50MG BASE	A207178 002	Mar 02, 2020
AB		EQ 100MG BASE	A207178 003	Mar 02, 2020
AB	AUROBINDO PHARMA LTD	EQ 20MG BASE	A203963 001	Nov 18, 2015
AB		EQ 25MG BASE	A203962 001	Jun 11, 2018
AB		EQ 50MG BASE	A203962 002	Jun 11, 2018
AB		EQ 100MG BASE	A203962 003	Jun 11, 2018
AB	CHARTWELL RX	EQ 20MG BASE	A202598 001	Nov 06, 2012
AB	HETERO LABS LTD V	EQ 20MG BASE	A203623 001	Nov 26, 2014
AB		EQ 25MG BASE	A202659 001	Jun 11, 2018
AB		EQ 50MG BASE	A202659 002	Jun 11, 2018
AB		EQ 100MG BASE	A202659 003	Jun 11, 2018
AB	LUPIN LTD	EQ 25MG BASE	A212051 001	Mar 22, 2019
AB		EQ 50MG BASE	A212051 002	Mar 22, 2019
AB		EQ 100MG BASE	A212051 003	Mar 22, 2019
AB	MACLEODS PHARMS LTD	EQ 20MG BASE	A203814 001	Dec 17, 2013
AB	MYLAN	EQ 25MG BASE	A201171 001	Mar 25, 2019
AB		EQ 50MG BASE	A201171 002	Mar 25, 2019
AB		EQ 100MG BASE	A201171 003	Mar 25, 2019
AB	MYLAN PHARMS INC	EQ 20MG BASE	A201150 001	Nov 09, 2012
AB	REYOUNG	EQ 25MG BASE	A208494 001	Jun 12, 2020
AB		EQ 50MG BASE	A208494 002	Jun 12, 2020
AB		EQ 100MG BASE	A208494 003	Jun 12, 2020
AB	RUBICON	EQ 20MG BASE	A204883 001	Jun 20, 2016
AB		EQ 25MG BASE	A204882 001	Jun 11, 2018
AB		EQ 50MG BASE	A204882 002	Jun 11, 2018
AB		EQ 100MG BASE	A204882 003	Jun 11, 2018
AB	SUNSHINE	EQ 25MG BASE	A213032 001	Jun 11, 2020
AB		EQ 50MG BASE	A213032 002	Jun 11, 2020
AB		EQ 100MG BASE	A213032 003	Jun 11, 2020
AB	TEVA	EQ 25MG BASE	A077342 001	Mar 09, 2016
AB		EQ 50MG BASE	A077342 002	Mar 09, 2016
AB		EQ 100MG BASE	A077342 003	Mar 09, 2016

PRESCRIPTION DRUG PRODUCT LIST

SILDENAFIL CITRATE

TABLET;ORAL

SILDENAFIL CITRATE

AB	TEVA PHARMS	EQ 20MG BASE	A078380 001	Jan 07, 2013
AB	TORRENT	EQ 25MG BASE	A091448 001	Jun 11, 2018
AB		EQ 50MG BASE	A091448 002	Jun 11, 2018
AB		EQ 100MG BASE	A091448 003	Jun 11, 2018
AB	TORRENT PHARMS LTD	EQ 20MG BASE	A091479 001	Nov 06, 2012
AB	UMEDICA LABS PVT LTD	EQ 25MG BASE	A209302 003	Jun 15, 2021
AB		EQ 50MG BASE	A209302 001	Aug 25, 2020
AB		EQ 100MG BASE	A209302 002	Aug 25, 2020
AB	WATSON LABS INC	EQ 20MG BASE	A202503 001	Nov 06, 2012
VIAGRA				
AB	+ UPJOHN	EQ 25MG BASE	N020895 001	Mar 27, 1998
AB	+	EQ 50MG BASE	N020895 002	Mar 27, 1998
AB	+	EQ 100MG BASE	N020895 003	Mar 27, 1998

SILODOSIN

CAPSULE;ORAL

RAPAFLO

AB	+	ALLERGAN	4MG	N022206 001	Oct 08, 2008
AB	+		8MG	N022206 002	Oct 08, 2008

SILODOSIN

AB		AJANTA PHARMA LTD	4MG	A211060 001	Dec 03, 2018
AB			8MG	A211060 002	Dec 03, 2018
AB		AMNEAL PHARMS CO	4MG	A209745 001	Dec 03, 2018
AB			8MG	A209745 002	Dec 03, 2018
AB		AUROBINDO PHARMA LTD	4MG	A210626 001	Dec 10, 2018
AB			8MG	A210626 002	Dec 10, 2018
AB		HETERO LABS LTD V	4MG	A204793 001	Feb 14, 2020
AB			8MG	A204793 002	Feb 14, 2020
AB		LUPIN LTD	4MG	A206541 001	Dec 03, 2018
AB			8MG	A206541 002	Dec 03, 2018
AB		MACLEODS PHARMS LTD	4MG	A211166 001	Dec 03, 2018
AB			8MG	A211166 002	Dec 03, 2018
AB		MSN	4MG	A210687 001	Dec 03, 2018
AB			8MG	A210687 002	Dec 03, 2018
AB		PRINSTON INC	4MG	A209029 001	Jan 04, 2022
AB			8MG	A209029 002	Jan 04, 2022
AB		SANDOZ INC	4MG	A204726 001	Mar 31, 2017
AB			8MG	A204726 002	Mar 31, 2017
AB		UPSHER SMITH LABS	4MG	A213230 001	Jan 03, 2022
AB			8MG	A213230 002	Jan 03, 2022

SILVER SULFADIAZINE

CREAM;TOPICAL

SILVADENE

AB	+	KING PHARMS LLC	1%	N017381 001	
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SSD

AB		DR REDDYS LA	1%	N018578 001	Feb 25, 1982
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THERMAZENE

AB		THEPHARMANETWORK LLC	1%	N018810 001	Dec 23, 1985
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SIMVASTATIN

SUSPENSION;ORAL

FLOLIPID

	+	TCG FLUENT PHARMA	20MG/5ML	N206679 001	Apr 21, 2016
	+		40MG/5ML	N206679 002	Apr 21, 2016

TABLET;ORAL

SIMVASTATIN

AB		ACCORD HLTHCARE	5MG	A078155 005	Apr 05, 2013
AB			10MG	A078155 002	Feb 26, 2008
AB			20MG	A078155 003	Feb 26, 2008
AB			40MG	A078155 004	Feb 26, 2008
AB			80MG	A078155 001	Feb 26, 2008
AB		AUROBINDO PHARMA	5MG	A077691 001	Dec 20, 2006
AB			10MG	A077691 002	Dec 20, 2006
AB			20MG	A077691 003	Dec 20, 2006
AB			40MG	A077691 004	Dec 20, 2006
AB			80MG	A077691 005	Dec 20, 2006
AB		BIOCON PHARMA	5MG	A078034 001	Dec 20, 2006

PRESCRIPTION DRUG PRODUCT LIST

SIMVASTATIN

TABLET; ORAL

SIMVASTATIN

<u>AB</u>		<u>10MG</u>	<u>A078034</u>	<u>002</u>	Dec 20, 2006
<u>AB</u>		<u>20MG</u>	<u>A078034</u>	<u>003</u>	Dec 20, 2006
<u>AB</u>		<u>40MG</u>	<u>A078034</u>	<u>004</u>	Dec 20, 2006
<u>AB</u>		<u>80MG</u>	<u>A078034</u>	<u>005</u>	Dec 20, 2006
<u>AB</u>	DR REDDYS LABS INC	<u>5MG</u>	<u>A077752</u>	<u>005</u>	Jan 23, 2008
<u>AB</u>		<u>10MG</u>	<u>A077752</u>	<u>001</u>	Dec 20, 2006
<u>AB</u>		<u>20MG</u>	<u>A077752</u>	<u>002</u>	Dec 20, 2006
<u>AB</u>		<u>40MG</u>	<u>A077752</u>	<u>003</u>	Dec 20, 2006
<u>AB</u>		<u>80MG</u>	<u>A077752</u>	<u>004</u>	Dec 20, 2006
<u>AB</u>	HETERO LABS LTD III	<u>5MG</u>	<u>A200895</u>	<u>001</u>	Nov 25, 2014
<u>AB</u>		<u>10MG</u>	<u>A200895</u>	<u>002</u>	Nov 25, 2014
<u>AB</u>		<u>20MG</u>	<u>A200895</u>	<u>003</u>	Nov 25, 2014
<u>AB</u>		<u>40MG</u>	<u>A200895</u>	<u>004</u>	Nov 25, 2014
<u>AB</u>		<u>80MG</u>	<u>A200895</u>	<u>005</u>	Nov 25, 2014
<u>AB</u>	HISUN PHARM HANGZHOU	<u>10MG</u>	<u>A206557</u>	<u>001</u>	Nov 13, 2017
<u>AB</u>		<u>20MG</u>	<u>A206557</u>	<u>002</u>	Nov 13, 2017
<u>AB</u>		<u>40MG</u>	<u>A206557</u>	<u>003</u>	Nov 13, 2017
<u>AB</u>		<u>80MG</u>	<u>A206557</u>	<u>004</u>	Nov 13, 2017
<u>AB</u>	LUPIN	<u>5MG</u>	<u>A078103</u>	<u>005</u>	Apr 14, 2009
<u>AB</u>		<u>10MG</u>	<u>A078103</u>	<u>001</u>	May 11, 2007
<u>AB</u>		<u>20MG</u>	<u>A078103</u>	<u>002</u>	May 11, 2007
<u>AB</u>		<u>40MG</u>	<u>A078103</u>	<u>003</u>	May 11, 2007
<u>AB</u>	!	<u>80MG</u>	<u>A078103</u>	<u>004</u>	May 11, 2007
<u>AB</u>	MICRO LABS	<u>5MG</u>	<u>A090383</u>	<u>001</u>	Sep 16, 2011
<u>AB</u>		<u>10MG</u>	<u>A090383</u>	<u>002</u>	Sep 16, 2011
<u>AB</u>		<u>20MG</u>	<u>A090383</u>	<u>003</u>	Sep 16, 2011
<u>AB</u>		<u>40MG</u>	<u>A090383</u>	<u>004</u>	Sep 16, 2011
<u>AB</u>		<u>80MG</u>	<u>A090383</u>	<u>005</u>	Sep 16, 2011
<u>AB</u>	OXFORD PHARMS	<u>5MG</u>	<u>A078735</u>	<u>001</u>	Aug 30, 2010
<u>AB</u>		<u>10MG</u>	<u>A078735</u>	<u>002</u>	Aug 30, 2010
<u>AB</u>		<u>20MG</u>	<u>A078735</u>	<u>003</u>	Aug 30, 2010
<u>AB</u>		<u>40MG</u>	<u>A078735</u>	<u>004</u>	Aug 30, 2010
<u>AB</u>		<u>80MG</u>	<u>A078735</u>	<u>005</u>	Aug 30, 2010
<u>AB</u>	WATSON LABS TEVA	<u>5MG</u>	<u>A076685</u>	<u>001</u>	Dec 20, 2006
<u>AB</u>		<u>10MG</u>	<u>A076685</u>	<u>002</u>	Dec 20, 2006
<u>AB</u>		<u>20MG</u>	<u>A076685</u>	<u>003</u>	Dec 20, 2006
<u>AB</u>		<u>40MG</u>	<u>A076685</u>	<u>004</u>	Dec 20, 2006
<u>AB</u>		<u>80MG</u>	<u>A076685</u>	<u>005</u>	Dec 20, 2006
<u>AB</u>	ZYDUS PHARMS USA	<u>5MG</u>	<u>A077837</u>	<u>001</u>	Dec 20, 2006
<u>AB</u>		<u>10MG</u>	<u>A077837</u>	<u>002</u>	Dec 20, 2006
<u>AB</u>		<u>20MG</u>	<u>A077837</u>	<u>003</u>	Dec 20, 2006
<u>AB</u>		<u>40MG</u>	<u>A077837</u>	<u>004</u>	Dec 20, 2006
<u>AB</u>		<u>80MG</u>	<u>A077837</u>	<u>005</u>	Dec 20, 2006
<u>ZOCOR</u>					
<u>AB</u>	+	ORGANON	<u>5MG</u>	<u>N019766</u>	<u>001</u> Dec 23, 1991
<u>AB</u>	+		<u>10MG</u>	<u>N019766</u>	<u>002</u> Dec 23, 1991
<u>AB</u>	+		<u>20MG</u>	<u>N019766</u>	<u>003</u> Dec 23, 1991
<u>AB</u>	+		<u>40MG</u>	<u>N019766</u>	<u>004</u> Dec 23, 1991

SINCALIDE

INJECTABLE; INJECTION

KINEVAC

+! BRACCO

0.005MG/VIAL

N017697 001

SINECATECHINS

OINTMENT; TOPICAL

VEREGEN

+! ANI PHARMS

15%

N021902 001 Oct 31, 2006

SIPONIMOD FUMARIC ACID

TABLET; ORAL

MAYZENT

+ NOVARTIS

EQ 0.25MG BASE

N209884 001 Mar 26, 2019

+

EQ 1MG BASE

N209884 003 Aug 24, 2021

+!

EQ 2MG BASE

N209884 002 Mar 26, 2019

PRESCRIPTION DRUG PRODUCT LIST

SIROLIMUS

POWDER; INTRAVENOUS

FYARRO

+! AADI

100MG/VIAL

N213312 001 Nov 22, 2021

SOLUTION; ORAL

RAPAMUNEAA +! PF PRISM CV1MG/MLN021083 001 Sep 15, 1999SIROLIMUSAA AMNEAL1MG/MLA211212 001 Oct 18, 2019AA APOTEX1MG/MLA211406 001 Oct 22, 2019AA NOVITIUM PHARMA1MG/MLA211040 001 Jan 28, 2019AA TORRENT1MG/MLA215016 001 Dec 27, 2021

TABLET; ORAL

RAPAMUNEAB + PF PRISM CV0.5MGN021110 004 Jan 25, 2010AB +1MGN021110 001 Aug 25, 2000AB +!2MGN021110 002 Aug 22, 2002SIROLIMUSAB ALKEM LABS LTD0.5MGA214753 001 Mar 12, 2021AB1MGA214753 002 Mar 12, 2021AB2MGA214753 003 Mar 12, 2021AB DR REDDYS LABS LTD1MGA201578 001 Oct 27, 2014AB2MGA201578 002 Oct 27, 2014AB GLENMARK PHARMS LTD0.5MGA208691 001 Oct 16, 2020AB1MGA208691 002 Oct 16, 2020AB2MGA208691 003 Oct 16, 2020AB ZYDUS PHARMS0.5MGA201676 003 Jan 08, 2014SITAGLIPTIN 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 ACETATEAP +! HOSPIRA2MEQ/MLN018893 001 May 04, 1983AP MILLA PHARMS2MEQ/MLA214805 001 May 04, 2021

FRESENIUS KABI USA

4MEQ/ML

A206687 001 Oct 30, 2017

SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION; INTRAVENOUS

AMMONULAP +! BAUSCH10%;10% (5GM/50ML;5GM/50ML)N020645 001 Feb 17, 2005SODIUM PHENYLACETATE AND SODIUM BENZOATEAP AILEX PHARMS LLC10%;10% (5GM/50ML;5GM/50ML)A207096 001 Feb 24, 2016AP MAIA PHARMS INC10%;10% (5GM/50ML;5GM/50ML)A208521 001 May 08, 2017AP NAVINTA LLC10%;10% (5GM/50ML;5GM/50ML)A205880 001 Aug 04, 2016

+! MAIA PHARMS INC

10%;10% (2GM/20ML;2GM/20ML)

N215025 001 Jun 10, 2021

SODIUM BICARBONATE

INJECTABLE; INJECTION

SODIUM BICARBONATEAP ! EXELA PHARMA0.5MEQ/MLA211091 001 Jun 20, 2019AP !0.9MEQ/MLA211091 002 Jun 20, 2019AP !1MEQ/MLA211091 003 Jun 20, 2019AP HOSPIRA INC0.5MEQ/MLA202679 001 Mar 07, 2017AP0.5MEQ/MLA202981 001 Mar 04, 2016AP0.9MEQ/MLA202494 001 Mar 06, 2017AP1MEQ/MLA202432 001 Sep 26, 2017AP1MEQ/MLA202494 002 Mar 06, 2017AP INTL MEDICATION SYS1MEQ/MLA203449 001 Sep 19, 2017

HOSPIRA INC

1MEQ/ML

A202495 001 Mar 06, 2017

SODIUM CHLORIDE

INJECTABLE; INJECTION

BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINERAP FRESENIUS KABI USA9MG/MLA088911 001 Feb 07, 1985AP +! HOSPIRA9MG/MLN018800 001 Oct 29, 1982SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINERAP + B BRAUN450MG/100MLN019635 001 Mar 09, 1988AP +! BAXTER HLTHCARE450MG/100MLN018016 001AP FRESENIUS KABI USA450MG/100MLA208122 001 Jul 23, 2018

PRESCRIPTION DRUG PRODUCT LIST

SODIUM CHLORIDE

INJECTABLE; INJECTION

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>450MG/100ML</u>	<u>N019759 001</u>	Jun 08, 1988
<u>AP</u>	+! ICU MEDICAL INC	<u>450MG/100ML</u>	<u>N018090 001</u>	

SODIUM CHLORIDE 0.9%

<u>AP</u>	HIKMA	<u>9MG/ML</u>	<u>A201850 001</u>	Jan 20, 2012
<u>AP</u>	SPECTRA MDCL DEVICES	<u>9MG/ML</u>	<u>A206171 001</u>	Jul 21, 2017

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	+! B BRAUN	<u>900MG/100ML</u>	<u>N017464 001</u>	
<u>AP</u>	+! BAXTER HLTHCARE	<u>900MG/100ML</u>	<u>N019635 002</u>	Mar 09, 1988
<u>AP</u>	+! BAXTER HLTHCARE	<u>9MG/ML</u>	<u>N016677 004</u>	Oct 30, 1985
<u>AP</u>	+! BAXTER HLTHCARE	<u>9MG/ML</u>	<u>N020178 002</u>	Dec 07, 1992
<u>AP</u>	+! BAXTER HLTHCARE	<u>900MG/100ML</u>	<u>N016677 001</u>	
<u>AP</u>	+! BAXTER HLTHCARE	<u>900MG/100ML</u>	<u>N020178 001</u>	Dec 07, 1992
<u>AP</u>	! FRESenius KABI USA	<u>9MG/ML</u>	<u>A088912 001</u>	Jan 10, 1985
<u>AP</u>	FRESenius KABI USA	<u>900MG/100ML</u>	<u>A207310 001</u>	Sep 19, 2017
<u>AP</u>	FRESenius MEDCL	<u>900MG/100ML</u>	<u>A078177 001</u>	Apr 12, 2007
<u>AP</u>	HAEMONETICS	<u>900MG/100ML</u>	<u>A076316 001</u>	Oct 27, 2004
<u>AP</u>	+! HOSPIRA	<u>9MG/ML</u>	<u>N018803 001</u>	Oct 29, 1982
<u>AP</u>	+! HOSPIRA	<u>9MG/ML</u>	<u>N019465 002</u>	Jul 15, 1985
<u>AP</u>	+! HOSPIRA	<u>900MG/100ML</u>	<u>N019465 001</u>	Jul 15, 1985
<u>AP</u>	+! HOSPIRA	<u>900MG/100ML</u>	<u>N019480 001</u>	Sep 17, 1985
<u>AP</u>	+! ICU MEDICAL INC	<u>900MG/100ML</u>	<u>N016366 001</u>	
<u>AP</u>	LABORATORIOS GRIFOLS	<u>900MG/100ML</u>	<u>A207956 001</u>	May 25, 2017
<u>AP</u>	NEPHRON	<u>9MG/ML</u>	<u>A211968 001</u>	Apr 23, 2020
<u>AP</u>	! TARO	<u>9MG/ML</u>	<u>A077407 001</u>	Aug 11, 2006

SODIUM CHLORIDE 3% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>3GM/100ML</u>	<u>N019635 003</u>	Mar 09, 1988
<u>AP</u>	+! BAXTER HLTHCARE	<u>3GM/100ML</u>	<u>N019022 001</u>	Nov 01, 1983
<u>AP</u>	FRESenius KABI USA	<u>3GM/100ML</u>	<u>A209476 001</u>	Mar 13, 2019

SODIUM CHLORIDE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>5GM/100ML</u>	<u>N019635 004</u>	Mar 09, 1988
<u>AP</u>	+ BAXTER HLTHCARE	<u>5GM/100ML</u>	<u>N019022 002</u>	Nov 01, 1983

SODIUM CHLORIDE 0.9%

+	B BRAUN	9MG/10ML	N019635 005	Aug 11, 2016
	HIKMA	9MG/ML	A201833 001	Sep 24, 2013
+	MEDEFIL INC	90MG/10ML (9MG/ML)	N202832 006	Jan 06, 2012

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+	LIEBEL-FLARSHEIM	1012.5MG/125ML (9MG/ML)	N021569 002	Jul 27, 2006
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SODIUM CHLORIDE 23.4%

	FRESenius KABI USA	234MG/ML	A212248 001	Apr 28, 2021
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SOLUTION; INTRAVENOUS

SODIUM CHLORIDE 14.6%

<u>AP</u>	FRESenius KABI USA	<u>100MEQ/40ML (2.5MEQ/ML)</u>	<u>A212070 002</u>	Apr 28, 2021
<u>AP</u>	+! HOSPIRA	<u>100MEQ/40ML (2.5MEQ/ML)</u>	<u>N018897 002</u>	Jul 20, 1984
	FRESenius KABI USA	50MEQ/20ML (2.5MEQ/ML)	A212070 001	Apr 28, 2021
	SODIUM CHLORIDE 23.4%			
+	HOSPIRA	400MEQ/100ML (4MEQ/ML)	N018897 003	Jun 18, 2020

SOLUTION; IRRIGATION

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AT</u>	+! B BRAUN	<u>900MG/100ML</u>	<u>N016733 001</u>	
<u>AT</u>	BAXTER HLTHCARE	<u>900MG/100ML</u>	<u>N017427 001</u>	
<u>AT</u>	BAXTER HLTHCARE	<u>900MG/100ML</u>	<u>N017867 001</u>	
<u>AT</u>	+ ICU MEDICAL INC	<u>900MG/100ML</u>	<u>N017514 001</u>	
<u>AT</u>	BAXTER HLTHCARE	<u>900MG/100ML</u>	<u>N018314 001</u>	

SOLUTION FOR SLUSH; IRRIGATION

SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER

+	BAXTER HLTHCARE	900MG/100ML	N019319 002	May 17, 1985
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SODIUM FLUORIDE F-18

INJECTABLE; INTRAVENOUS

SODIUM FLUORIDE F-18

<u>AP</u>	3D IMAGING DRUG	<u>10-200mCi/ML</u>	<u>A203777 001</u>	Oct 19, 2015
<u>AP</u>	BIOMEDCL RES FDN	<u>10-200mCi/ML</u>	<u>A204351 001</u>	Jan 09, 2015
<u>AP</u>	CARDINAL HEALTH 414	<u>10-200mCi/ML</u>	<u>A203780 001</u>	Jul 30, 2015
<u>AP</u>	ESSENTIAL ISOTOPE	<u>10-200mCi/ML</u>	<u>A204541 001</u>	Oct 29, 2014
<u>AP</u>	HOT SHOTS NM LLC	<u>10-200mCi/ML</u>	<u>A204530 001</u>	Jul 29, 2015
<u>AP</u>	JUBILANT DRAXIMAGE	<u>10-200mCi/ML</u>	<u>A203968 001</u>	Oct 23, 2015
<u>AP</u>	KREITCHMAN PET CTR	<u>10-200mCi/ML</u>	<u>A203936 001</u>	May 19, 2016
<u>AP</u>	MIDWEST MEDCL	<u>10-200mCi/ML</u>	<u>A204440 001</u>	Nov 17, 2015

PRESCRIPTION DRUG PRODUCT LIST

SODIUM FLUORIDE F-18

INJECTABLE; INTRAVENOUS

SODIUM FLUORIDE F-18

AP	MIPS CRF	<u>10-200mCi/ML</u>	<u>A204517</u>	<u>001</u>	Jul 21, 2015
AP	NCM USA BRONX LLC	<u>10-200mCi/ML</u>	<u>A204513</u>	<u>001</u>	Nov 28, 2014
AP	NUKEMED	<u>10-200mCi/ML</u>	<u>A203912</u>	<u>001</u>	Apr 22, 2015
AP	PETNET	<u>10-200mCi/ML</u>	<u>A203890</u>	<u>001</u>	Sep 28, 2015
AP	PRECISION NUCLEAR	<u>10-200mCi/ML</u>	<u>A204542</u>	<u>001</u>	Feb 27, 2015
AP	SHERTECH LABS LLC	<u>10-200mCi/ML</u>	<u>A204315</u>	<u>001</u>	Sep 22, 2014
AP	SOFIE	<u>10-200mCi/ML</u>	<u>A203592</u>	<u>001</u>	Aug 18, 2015
AP	!	<u>10-200mCi/ML</u>	<u>A203544</u>	<u>001</u>	Dec 26, 2012
AP	UNIV UTAH CYCLOTRON	<u>10-200mCi/ML</u>	<u>A204497</u>	<u>001</u>	Apr 20, 2015
	!	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

AA	+	CARDINAL HEALTH 418	<u>100uCi</u>	<u>N018671</u>	<u>001</u>	May 27, 1982
AA	+		<u>200uCi</u>	<u>N018671</u>	<u>002</u>	May 27, 1982
AA		CURIUM	<u>100uCi</u>	<u>A071909</u>	<u>001</u>	Feb 28, 1989
AA			<u>200uCi</u>	<u>A071910</u>	<u>001</u>	Feb 28, 1989

SODIUM IODIDE I-131

CAPSULE; ORAL

SODIUM IODIDE I 131

+ JUBILANT

0.009-0.1mCi

N021305 006 May 19, 2005

SOLUTION; ORAL

HICON

AA	+	JUBILANT	<u>250-1000mCi</u>	<u>N021305</u>	<u>007</u>	Dec 05, 2011
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SODIUM IODIDE I 131

AA		INTL ISOTOPES	<u>250-1000mCi</u>	<u>A209166</u>	<u>001</u>	Feb 05, 2020
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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

NITHIODE

+! HOPE PHARMS

300MG/10ML (30MG/ML), N/A; N/A, 12.5GM/50ML (250MG/ML)

N201444 001 Jan 14, 2011

SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION

NITROPRESS

AP	!	HOSPIRA	<u>25MG/ML</u>	<u>A071961</u>	<u>001</u>	Aug 01, 1988
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SODIUM NITROPRUSSIDE

AP		AKORN	<u>25MG/ML</u>	<u>A208635</u>	<u>001</u>	May 04, 2017
AP		AMNEAL	<u>25MG/ML</u>	<u>A209493</u>	<u>001</u>	Nov 07, 2017
AP		BE PHARMS	<u>25MG/ML</u>	<u>A214971</u>	<u>001</u>	Jul 12, 2021
AP		DR REDDYS LABS LTD	<u>25MG/ML</u>	<u>A210114</u>	<u>001</u>	Apr 10, 2019
AP		HAINAN POLY PHARM	<u>25MG/ML</u>	<u>A214199</u>	<u>001</u>	Aug 25, 2020
AP		HONG KONG	<u>25MG/ML</u>	<u>A211016</u>	<u>001</u>	Nov 29, 2019
AP		MEDICURE	<u>25MG/ML</u>	<u>A209584</u>	<u>001</u>	Aug 10, 2018
AP		MICRO LABS	<u>25MG/ML</u>	<u>A209352</u>	<u>001</u>	Dec 08, 2017
AP		MYLAN LABS LTD	<u>25MG/ML</u>	<u>A210763</u>	<u>001</u>	Apr 17, 2018
AP		NEXUS PHARMS	<u>25MG/ML</u>	<u>A207499</u>	<u>001</u>	May 25, 2017
AP		SAGENT PHARMS INC	<u>25MG/ML</u>	<u>A207426</u>	<u>001</u>	Dec 08, 2016
AP		SOMERSET THERAPS LLC	<u>25MG/ML</u>	<u>A210882</u>	<u>001</u>	Aug 17, 2018
AP		XIROMED	<u>25MG/ML</u>	<u>A211277</u>	<u>001</u>	Oct 29, 2020
		SOLUTION; INTRAVENOUS				
		NIPRIDE RTU IN SODIUM CHLORIDE 0.9%				
		+	EXELA PHARMA	20MG/100ML (0.2MG/ML)	N209387	003 Jul 13, 2018
		+		50MG/100ML (0.5MG/ML)	N209387	001 Mar 08, 2017

SODIUM OXYBATE

SOLUTION; ORAL

XYREM

+! JAZZ PHARMS

0.5GM/ML

N021196 001 Jul 17, 2002

PRESCRIPTION DRUG PRODUCT LIST

SODIUM PHENYL BUTYRATE

POWDER; ORAL

BUPHENYL

AB	+!	HORIZON THERAP	3GM/TEASPOONFUL	N020573 001	Apr 30, 1996
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SODIUM PHENYL BUTYRATE

AB		PAR PHARM	3GM/TEASPOONFUL	A203918 001	Jun 15, 2016
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AB		SIGMAPHARM LABS LLC	3GM/TEASPOONFUL	A202819 001	Mar 22, 2013
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TABLET; ORAL

BUPHENYL

AB	+!	HORIZON THERAP	500MG	N020572 001	May 13, 1996
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SODIUM PHENYL BUTYRATE

AB		PAR PHARM	500MG	A204395 001	Apr 15, 2016
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SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL

MONOBASIC SODIUM PHOSPHATE AND DIBASIC SODIUM PHOSPHATE

AB		NOVEL LABS INC	0.398GM;1.102GM	A079247 001	Dec 30, 2011
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OSMOPREP

AB	+!	SALIX PHARMS	0.398GM;1.102GM	N021892 001	Mar 16, 2006
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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
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SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

KALEXATE

AA	!	KVK TECH	454GM/BOT	A040905 001	Mar 30, 2009
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KIONEX

AA		ANI PHARMS	454GM/BOT	A040029 001	Feb 06, 1998
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SODIUM POLYSTYRENE SULFONATE

AA		BELCHER	454GM/BOT	A205727 001	Feb 23, 2016
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AA		BIOPHARM	453.6GM/BOT	A090313 001	Dec 21, 2011
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AA		CHARTWELL RX	454GM/BOT	A206815 001	Feb 18, 2016
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AA		CMP PHARMA INC	454GM/BOT	A089910 001	Jan 19, 1989
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AA		EPIC PHARMA LLC	453.6GM/BOT	A202333 001	Mar 19, 2014
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AA		NUVO PHARMS INC	454GM/BOT	A204071 001	Nov 28, 2014
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KALEXATE

	KVK TECH	15GM/BOT	A040905 002	Apr 03, 2015
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SODIUM POLYSTYRENE SULFONATE

	NUVO PHARMS INC	15GM/BOT	A204071 002	Nov 28, 2014
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SUSPENSION; ORAL, RECTAL

SPS

+!	CMP PHARMA INC	15GM/60ML	A087859 001	Dec 08, 1982
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SODIUM TETRADECYL SULFATE

INJECTABLE; INJECTION

SODIUM TETRADECYL SULFATE

AP		CUSTOPHARM INC	60MG/2ML (30MG/ML)	A209937 001	Dec 09, 2019
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SOTRADECOL

AP	!	MYLAN INSTITUTIONAL	60MG/2ML (30MG/ML)	A040541 002	Nov 12, 2004
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!			20MG/2ML (10MG/ML)	A040541 001	Nov 12, 2004
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SODIUM THIOSULFATE

SOLUTION; INTRAVENOUS

SODIUM THIOSULFATE

+!	HOPE PHARMS	12.5GM/50ML (250MG/ML)	N203923 001	Feb 14, 2012
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SODIUM ZIRCONIUM CYCLOSILICATE

FOR SUSPENSION; ORAL

LOKELMA

+	ASTRAZENECA	5GM/PACKET	N207078 001	May 18, 2018
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+!		10GM/PACKET	N207078 002	May 18, 2018
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SOFOSBUVIR

PELLETS; ORAL

SOVALDI

+	GILEAD SCIENCES INC	150MG/PACKET	N212480 001	Aug 28, 2019
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+!		200MG/PACKET	N212480 002	Aug 28, 2019
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TABLET; ORAL

SOVALDI

+	GILEAD SCIENCES INC	200MG	N204671 002	Aug 28, 2019
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+!		400MG	N204671 001	Dec 06, 2013
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PRESCRIPTION DRUG PRODUCT LIST

SOFOSBUVIR; VELPATASVIR

PELLETS; ORAL

EPCLUSA

+	GILEAD SCIENCES INC	150MG;37.5MG/PACKET	N214187	001	Jun 10, 2021
+	!	200MG;50MG/PACKET	N214187	002	Jun 10, 2021

TABLET; ORAL

EPCLUSA

+	GILEAD SCIENCES INC	200MG;50MG	N208341	002	Mar 19, 2020
+	!	400MG;100MG	N208341	001	Jun 28, 2016

SOFOSBUVIR; VELPATASVIR; VOXILAPREVIR

TABLET; ORAL

VOSEVI

+	GILEAD SCIENCES INC	400MG;100MG;100MG	N209195	001	Jul 18, 2017
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SOLIFENACIN SUCCINATE

SUSPENSION; ORAL

VESICARE LS

+	ASTELLAS	1MG/ML	N209529	001	May 26, 2020
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TABLET; ORAL

SOLIFENACIN SUCCINATE

AB	ACCORD HLTHCARE	5MG	A207477	001	Jan 04, 2022
AB		10MG	A207477	002	Jan 04, 2022
AB	ALEMBIC PHARMS LTD	5MG	A205575	001	May 20, 2019
AB		10MG	A205575	002	May 20, 2019
AB	ALKEM LABS LTD	5MG	A210224	001	May 20, 2019
AB		10MG	A210224	002	May 20, 2019
AB	AMNEAL PHARMS CO	5MG	A209719	001	May 20, 2019
AB		10MG	A209719	002	May 20, 2019
AB	AUSTARPHARMA	5MG	A210281	001	May 20, 2019
AB		10MG	A210281	002	May 20, 2019
AB	CELLTRION	5MG	A210582	001	May 20, 2019
AB		10MG	A210582	002	May 20, 2019
AB	CIPLA	5MG	A209839	001	May 20, 2019
AB		10MG	A209839	002	May 20, 2019
AB	GLENMARK PHARMS INC	5MG	A209239	001	May 20, 2019
AB		10MG	A209239	002	May 20, 2019
AB	GRAVITI PHARMS	5MG	A211423	001	Dec 11, 2019
AB		10MG	A211423	002	Dec 11, 2019
AB	MSN	5MG	A210688	001	May 20, 2019
AB		10MG	A210688	002	May 20, 2019
AB	QILU	5MG	A209333	001	May 20, 2019
AB		10MG	A209333	002	May 20, 2019
AB	SCIEGEN PHARMS INC	5MG	A211657	001	May 20, 2019
AB		10MG	A211657	002	May 20, 2019
AB	STRIDES PHARMA	5MG	A212214	001	Sep 26, 2019
AB		10MG	A212214	002	Sep 26, 2019
AB	TEVA PHARMS USA	5MG	A091464	001	Apr 02, 2014
AB		10MG	A091464	002	Apr 02, 2014
AB	UNICHEM	5MG	A211701	001	Aug 27, 2019
AB		10MG	A211701	002	Aug 27, 2019
AB	WATSON LABS INC	5MG	A202551	001	May 20, 2019
AB		10MG	A202551	002	May 20, 2019

VESICARE

AB	+	ASTELLAS	5MG	N021518	001	Nov 19, 2004
AB	+	!	10MG	N021518	002	Nov 19, 2004

SOLRIAMFETOL HYDROCHLORIDE

TABLET; ORAL

SUNOSI

+	JAZZ	EQ 75MG BASE	N211230	001	Jun 17, 2019
+	!	EQ 150MG BASE	N211230	002	Jun 17, 2019

SONIDEGIB PHOSPHATE

CAPSULE; ORAL

ODOMZO

+	SUN PHARMA GLOBAL	EQ 200MG BASE	N205266	001	Jul 24, 2015
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SORAFENIB TOSYLATE

TABLET; ORAL

NEXAVAR

+	BAYER HLTHCARE	EQ 200MG BASE	N021923	001	Dec 20, 2005
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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

BETAPACE**AB** + COVIS**80MG****N019865 001** Oct 30, 1992**AB** +**120MG****N019865 005** Apr 20, 1994**AB** +!**160MG****N019865 002** Oct 30, 1992**AB** +**240MG****N019865 003** Oct 30, 1992BETAPACE AF**AB** + COVIS**80MG****N021151 001** Feb 22, 2000**AB** +**120MG****N021151 002** Feb 22, 2000**AB** +!**160MG****N021151 003** Feb 22, 2000SORINE**AB** UPSHER SMITH LABS**80MG****A075500 001** Apr 27, 2001**AB****120MG****A075500 004** Apr 27, 2001**AB****160MG****A075500 002** Apr 27, 2001**AB****240MG****A075500 003** Apr 27, 2001SOTALOL HYDROCHLORIDE**AB** APOTEX**80MG****A076140 001** Sep 26, 2002**AB****80MG****A076214 001** Aug 27, 2003**AB****120MG****A076140 002** Sep 26, 2002**AB****120MG****A076214 002** Aug 27, 2003**AB****160MG****A076140 003** Sep 26, 2002**AB****160MG****A076214 003** Aug 27, 2003**AB****240MG****A076140 004** Sep 26, 2002**AB** BEXIMCO PHARMS USA**80MG****A207428 001** Oct 21, 2016**AB****80MG****A207429 001** Nov 02, 2018**AB****120MG****A207428 002** Oct 21, 2016**AB****120MG****A207429 002** Nov 02, 2018**AB****160MG****A207428 003** Oct 21, 2016**AB****160MG****A207429 003** Nov 02, 2018**AB** EPIC PHARMA INC**80MG****A077070 001** Nov 04, 2005**AB****120MG****A077070 002** Nov 04, 2005**AB****160MG****A077070 003** Nov 04, 2005**AB** OXFORD PHARMS**80MG****A075563 001** Nov 07, 2003**AB****120MG****A075563 002** Nov 07, 2003**AB****160MG****A075563 003** Nov 07, 2003**AB****240MG****A075563 004** Nov 07, 2003**AB** TEVA**80MG****A075429 001** May 01, 2000**AB****120MG****A075429 002** May 01, 2000**AB****160MG****A075429 003** May 01, 2000**AB****240MG****A075429 004** May 01, 2000SOTORASIB

TABLET;ORAL

LUMAKRAS

+! AMGEN INC 120MG

N214665 001 May 28, 2021

SOYBEAN OIL

INJECTABLE; INJECTION

INTRALIPID 10%**AP** +! FRESENIUS**10%****N017643 001**INTRALIPID 20%**AP** +! FRESENIUS**20%****N018449 001****AP** +!**20%****N020248 001** Aug 07, 1996NUTRILIPID 10%**AP** +! B BRAUN**10%****N019531 001** May 28, 1993NUTRILIPID 20%**AP** +! B BRAUN**20%****N019531 002** May 28, 1993

INTRALIPID 30%

+! FRESENIUS 30%

N019942 001 Dec 30, 1993

PRESCRIPTION DRUG PRODUCT LIST

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

ALDACTONEAB + PFIZER 25MG N012151 009 Dec 30, 1983AB + 50MG N012151 008 Dec 30, 1982AB +! 100MG N012151 010 Dec 30, 1983SPIRONOLACTONEAB ACCORD HLTHCARE 25MG A203512 001 Sep 19, 2016AB 50MG A203512 002 Sep 19, 2016AB 100MG A203512 003 Sep 19, 2016AB AMNEAL PHARMS 25MG A091426 001 Jul 02, 2010AB 50MG A091426 002 Jul 02, 2010AB 100MG A091426 003 Jul 02, 2010AB AUROBINDO PHARMA 25MG A202187 001 Mar 06, 2014

LTD

AB 50MG A202187 002 Mar 06, 2014AB 100MG A202187 003 Mar 06, 2014AB JUBILANT GENERICS 25MG A203253 001 Apr 23, 2014AB 50MG A203253 002 Apr 23, 2014AB 100MG A203253 003 Apr 23, 2014AB MYLAN 25MG A040424 001 Aug 20, 2001AB 50MG A040424 002 Aug 20, 2001AB 100MG A040424 003 Aug 20, 2001AB SUN PHARM 25MG A089424 001 Jul 23, 1986

INDUSTRIES

AB 50MG A089424 002 Aug 11, 1999AB 100MG A089424 003 Aug 11, 1999AB ZYDUS PHARMS 25MG A205936 001 Jul 18, 2018AB 50MG A205936 002 Jul 18, 2018AB 100MG A205936 003 Jul 18, 2018STERILE WATER FOR INJECTION

LIQUID;N/A

BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINERAP +! HOSPIRA 100% N018802 001 Oct 27, 1982STERILE WATER FOR INJECTIONAP FRESENIUS KABI USA 100% A209689 001 Nov 24, 2017AP HIKMA 100% (10ML) A206369 001 Sep 02, 2015AP +! HOSPIRA 100% (10ML) N018801 002 Oct 27, 1982AP MEDEFIL INC 100% (10ML) A211188 005 Dec 02, 2019STERILE WATER FOR INJECTION IN PLASTIC CONTAINERAP +! B BRAUN 100% N019633 001 Feb 29, 1988AP +! BAXTER HLTHCARE 100% N018632 001 Jun 30, 1982AP +! 100% N018632 002 Apr 19, 1988AP FRESENIUS KABI USA 100% A088400 001 Jan 16, 1984AP +! ICU MEDICAL INC 100% N018233 001AP +! 100% N019869 001 Dec 26, 1989AP TARO 100% A077393 001 Aug 11, 2006

STERILE WATER FOR INJECTION

+! HOSPIRA 100% (20ML) N018801 003 Oct 27, 1982

+! 100% (50ML) N018801 004 Oct 27, 1982

+! 100% (100ML) N018801 005 Oct 27, 1982

MEDEFIL INC 100% (1ML) A211188 001 Dec 02, 2019

! 100% (2.5ML) A211188 002 Dec 02, 2019

! 100% (3ML) A211188 003 Dec 02, 2019

! 100% (5ML) A211188 004 Dec 02, 2019

STERILE WATER FOR IRRIGATION

LIQUID;IRRIGATION

STERILE WATERAT + BAXTER HLTHCARE 100% N017428 001STERILE WATER IN PLASTIC CONTAINERAT + B BRAUN 100% N016734 001AT BAXTER HLTHCARE 100% N017866 001AT ICU MEDICAL INC 100% N017513 001AT 100% N018313 001

PRESCRIPTION DRUG PRODUCT LIST

STIRIPENTOL

CAPSULE; ORAL

DIACOMIT

+ BIOCDEX SA

250MG

N206709 001 Aug 20, 2018

+!

500MG

N206709 002 Aug 20, 2018

FOR SUSPENSION; ORAL

DIACOMIT

+ BIOCDEX SA

250MG/PACKET

N207223 001 Aug 20, 2018

+!

500MG/PACKET

N207223 002 Aug 20, 2018

STREPTOMYCIN SULFATE

INJECTABLE; INJECTION

STREPTOMYCIN SULFATE

! XGEN PHARMS

EQ 1GM BASE/VIAL

A064210 001 Jun 30, 1998

STREPTOZOCIN

INJECTABLE; INJECTION

ZANOSAR

+! TEVA PHARMS USA

1GM/VIAL

N050577 001 May 07, 1982

STRONTIUM CHLORIDE SR-89

INJECTABLE; INJECTION

METASTRON**AP** +! Q BIOMED**1mCi/ML****N020134 001** Jun 18, 1993STRONTIUM CHLORIDE SR-89**AP** Q BIOMED**1mCi/ML****A075941 001** Jan 06, 2003SUCCIMER

CAPSULE; ORAL

CHEMET

+! RECORDATI RARE

100MG

N019998 002 Jan 30, 1991

SUCCINYLCHOLINE CHLORIDE

INJECTABLE; INJECTION

ANECTINE**AP** +! SANDOZ INC**20MG/ML****N008453 002**QUELICIN**AP** +! HOSPIRA**20MG/ML****N008845 006**SUCCINYLCHOLINE CHLORIDE**AP** ACCORD HLTHCARE**20MG/ML****A213705 001** May 20, 2020**AP** AMNEAL**20MG/ML****A211432 001** Nov 16, 2018**AP** AMRING PHARMS**20MG/ML****A210231 001** Jun 04, 2018**AP** ASPIRO**20MG/ML****A213810 001** May 04, 2020**AP** BRECKENRIDGE**20MG/ML****A212638 001** Oct 09, 2019**AP** DEVA HOLDING AS**20MG/ML****A214491 001** Dec 21, 2020**AP** DR REDDYS**20MG/ML****A210698 001** Aug 02, 2019**AP** FRESENIUS KABI USA**20MG/ML****A211346 001** Nov 20, 2020**AP** GLAND PHARMA LTD**20MG/ML****A214246 001** Jun 16, 2020**AP** HIKMA**20MG/ML****A213229 001** Jun 12, 2020**AP** HONG KONG**20MG/ML****A214514 001** Oct 19, 2021**AP** INDOCO**20MG/ML****A214308 001** May 22, 2020**AP** MICRO LABS**20MG/ML****A214879 001** Nov 24, 2020**AP** NEXUS PHARMS**20MG/ML****A213552 001** Oct 27, 2020**AP** NIVAGEN PHARMS INC**20MG/ML****A211625 001** May 19, 2020**AP** SAGENT PHARMS INC**20MG/ML****A215022 001** Mar 29, 2021**AP** SOMERSET THERAPS**20MG/ML****A211589 001** Jan 15, 2020**AP** ! ZYDUS PHARMS**20MG/ML****A209467 001** May 04, 2018

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

SUCCINYLCHOLINE CHLORIDE

+! HIKMA

100MG/5ML (20MG/ML)

N215143 001 Aug 20, 2021

SUCRALFATE

SUSPENSION; ORAL

CARAFATE**AB** +! ALLERGAN**1GM/10ML****N019183 001** Dec 16, 1993SUCRALFATE**AB** AMNEAL**1GM/10ML****A209356 001** Dec 02, 2019

TABLET; ORAL

CARAFATE**AB** +! ALLERGAN**1GM****N018333 001**SUCRALFATE**AB** NOSTRUM LABS INC**1GM****A074415 001** Jun 08, 1998**AB** TEVA**1GM****A070848 001** Mar 29, 1996

PRESCRIPTION DRUG PRODUCT LIST

SUFENTANIL CITRATE

INJECTABLE; INJECTION

SUFENTA PRESERVATIVE FREE

AP	+ !	AKORN	EQ 0.05MG BASE/ML	N019050 001	May 04, 1984
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SUFENTANIL CITRATE

AP		HIKMA	EQ 0.05MG BASE/ML	A074413 001	Dec 15, 1995
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AP		HOSPIRA	EQ 0.05MG BASE/ML	A074534 001	Dec 11, 1996
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TABLET; SUBLINGUAL

DSUVIA

+ !	ACELRX PHARMS	EQ 0.03MG BASE	N209128 001	Nov 02, 2018
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SUGAMMADEX SODIUM

SOLUTION; INTRAVENOUS

BRIDION

+	ORGANON SUB MERCK	EQ 200MG BASE/2ML (EQ 100MG BASE/ML)	N022225 002	Dec 15, 2015
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+ !		EQ 500MG BASE/5ML (EQ 100MG BASE/ML)	N022225 001	Dec 15, 2015
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SULCONAZOLE NITRATE

CREAM; TOPICAL

EXELDERM

+ !	JOURNEY	1%	N018737 001	Feb 28, 1989
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SOLUTION; TOPICAL

EXELDERM

+ !	JOURNEY	1%	N018738 001	Aug 30, 1985
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SULFACETAMIDE SODIUM

LOTION; TOPICAL

KLARON

AB	+ !	BAUSCH	10%	N019931 001	Dec 23, 1996
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SULFACETAMIDE SODIUM

AB		FOUGERA PHARMS	10%	A077015 001	Nov 17, 2006
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AB		PADAGIS US	10%	A078649 001	Mar 23, 2009
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AB		TARO	10%	A078668 001	May 20, 2009
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OINTMENT; OPHTHALMIC

SULFACETAMIDE SODIUM

!	PADAGIS US	10%	A080029 001	
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SOLUTION/DROPS; OPHTHALMIC

BLEPH-10

AT	+ !	ALLERGAN	10%	A080028 001	
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SULFACETAMIDE SODIUM

AT		AKORN	10%	A040215 001	May 25, 1999
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AT		BAUSCH AND LOMB	10%	A040066 001	Dec 28, 1994
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AT		SANDOZ INC	10%	A089560 001	Oct 18, 1988
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SULFADIAZINE

TABLET; ORAL

SULFADIAZINE

!	SANDOZ	500MG	A040091 001	Jul 29, 1994
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SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AP	!	MYLAN LABS LTD	80MG/ML; 16MG/ML	A206607 001	Aug 30, 2017
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AP		SOMERSET	80MG/ML; 16MG/ML	A212231 001	Jun 26, 2019
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AP		TEVA PHARMS USA	80MG/ML; 16MG/ML	A073303 001	Oct 31, 1991
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SUSPENSION; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AB	!	AKORN	200MG/5ML; 40MG/5ML	A074650 001	Dec 29, 1997
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AB		AUROBINDO PHARMA	200MG/5ML; 40MG/5ML	A091348 001	Jun 08, 2010
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AB		LANNETT CO INC	200MG/5ML; 40MG/5ML	A077785 001	Jan 24, 2007
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AB		LUPIN LTD	200MG/5ML; 40MG/5ML	A212699 001	Jan 05, 2021
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AB		PRASCO	200MG/5ML; 40MG/5ML	A077612 001	Nov 13, 2006
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SULFATRIM PEDIATRIC

AB		PHARM ASSOC	200MG/5ML; 40MG/5ML	N018615 001	Jan 07, 1983
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TABLET; ORAL

BACTRIM

AB	+	SUN PHARM INDUSTRIES	400MG; 80MG	N017377 001	
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BACTRIM DS

AB	+ !	SUN PHARM INDUSTRIES	800MG; 160MG	N017377 002	
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SEPTRA

AB		MONARCH PHARMS	400MG; 80MG	N017376 001	
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SEPTRA DS

AB		MONARCH PHARMS	800MG; 160MG	N017376 002	
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PRESCRIPTION DRUG PRODUCT LIST

SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

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

SULFASALAZINE

TABLET; ORAL

AZULFIDINE

AB	+ ! PFIZER	500MG	N007073 001	
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SULFASALAZINE

AB	CHARTWELL	500MG	A080197 001	
AB	NUVO PHARMS INC	500MG	A040349 001	Jan 11, 2002
AB	WATSON LABS	500MG	A085828 001	

TABLET, DELAYED RELEASE; ORAL

AZULFIDINE EN-TABS

AB	+ ! PFIZER	500MG	N007073 002	Apr 06, 1983
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SULFASALAZINE

AB	NUVO PHARMS INC	500MG	A075339 001	Jan 11, 2002
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SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES

FOR SUSPENSION; INTRAVENOUS

LUMASON

+ !	BRACCO	60.7MG/25MG	N203684 001	Oct 15, 2014
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SULINDAC

TABLET; ORAL

SULINDAC

AB	EPIC PHARMA	150MG	A072710 001	Mar 25, 1991
AB		200MG	A072711 001	Mar 25, 1991
AB	SUN PHARM INDUSTRIES	150MG	A072050 001	Apr 17, 1991
AB		200MG	A072051 001	Apr 17, 1991
AB	WATSON LABS	150MG	A071891 001	Apr 03, 1990
AB	!	200MG	A071795 001	Apr 03, 1990

SUMATRIPTAN

SPRAY; NASAL

IMITREX

AB	+ ! GLAXOSMITHKLINE	5MG/SPRAY	N020626 001	Aug 26, 1997
AB	+ !	20MG/SPRAY	N020626 003	Aug 26, 1997

SUMATRIPTAN

AB	CIPLA	20MG/SPRAY	A214209 001	Feb 22, 2021
AB	FLORIDA	20MG/SPRAY	A208967 001	Feb 17, 2021
AB	LANNETT CO INC	5MG/SPRAY	A204841 001	Feb 19, 2016
AB		20MG/SPRAY	A204841 002	Feb 19, 2016
AB	PADAGIS ISRAEL	20MG/SPRAY	A213465 001	Sep 21, 2020

TOSYMRA

+ !	UPSHER SMITH LABS	10MG/SPRAY	N210884 001	Jan 25, 2019
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SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS

IMITREX STATDOSE

AB	+ ! GLAXOSMITHKLINE	EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)	N020080 002	Feb 01, 2006
AB	+ !	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N020080 003	Dec 23, 1996

SUMATRIPTAN SUCCINATE

AB	ANTARES PHARMA INC	EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)	A078319 001	Dec 10, 2015
AB		EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A078319 002	Dec 10, 2015
AB	DR REDDYS	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A090495 001	Jan 29, 2014
AB	SUN PHARM	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A090358 001	Jun 21, 2011

IMITREX

AP	+ ! GLAXOSMITHKLINE	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N020080 001	Dec 28, 1992
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SUMATRIPTAN SUCCINATE

AP	CAPLIN	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A213998 001	Jul 13, 2021
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PRESCRIPTION DRUG PRODUCT LIST

SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS

SUMATRIPTAN SUCCINATE

<u>AP</u>	EUGIA PHARMA	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A202758 001</u>	Apr 23, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A079242 001</u>	Mar 02, 2009
<u>AP</u>	HIKMA	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A200183 001</u>	Sep 16, 2013
<u>AP</u>	PAR PHARM	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A077332 001</u>	Oct 09, 2009
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A079123 001</u>	Feb 06, 2009
<u>AP</u>	WOCKHARDT	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A078593 001</u>	Feb 06, 2009

POWDER; NASAL

ONZETRA XSAIL

+! CURRAX

EQ 11MG BASE

N206099 001 Jan 27, 2016

SOLUTION; SUBCUTANEOUS

ZEMBRACE SYMTOUCH

+ UPSHER SMITH LABS

EQ 3MG BASE/0.5ML (EQ 3MG BASE/0.5ML)

N208223 001 Jan 28, 2016

SPRAY; NASAL

SUMATRIPTAN

<u>AB</u>	PADAGIS ISRAEL	<u>5MG/SPRAY</u>	<u>A213465 002</u>	Sep 21, 2020
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TABLET; ORAL

IMITREX

<u>AB</u>	+ GLAXOSMITHKLINE	<u>EQ 25MG BASE</u>	<u>N020132 002</u>	Jun 01, 1995
<u>AB</u>	+	<u>EQ 50MG BASE</u>	<u>N020132 003</u>	Jun 01, 1995
<u>AB</u>	+!	<u>EQ 100MG BASE</u>	<u>N020132 001</u>	Jun 01, 1995

SUMATRIPTAN SUCCINATE

<u>AB</u>	AUROBINDO PHARMA	<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>	COREPHARMA	<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>	DR REDDYS LABS INC	<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>	MYLAN	<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>	ORBION PHARMS	<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>	SUN PHARM INDS	<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>	SUN PHARM INDS LTD	<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>	WATSON LABS	<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

SUNITINIB MALATE

CAPSULE; ORAL

SUNITINIB MALATE

<u>AB</u>	MYLAN	<u>EQ 12.5MG BASE</u>	<u>A201275 001</u>	Dec 06, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A201275 002</u>	Dec 06, 2021
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A201275 003</u>	Dec 06, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A201275 004</u>	Dec 06, 2021
<u>AB</u>	SUN PHARM	<u>EQ 12.5MG BASE</u>	<u>A213914 001</u>	Aug 16, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213914 002</u>	Aug 16, 2021
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A213914 003</u>	Aug 16, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A213914 004</u>	Aug 16, 2021
<u>AB</u>	TEVA PHARMS USA	<u>EQ 12.5MG BASE</u>	<u>A213803 001</u>	Nov 30, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213803 002</u>	Nov 30, 2021
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A213803 003</u>	Nov 30, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A213803 004</u>	Nov 30, 2021

SUTENT

<u>AB</u>	+ CPPI CV	<u>EQ 12.5MG BASE</u>	<u>N021938 001</u>	Jan 26, 2006
<u>AB</u>	+	<u>EQ 25MG BASE</u>	<u>N021938 002</u>	Jan 26, 2006
<u>AB</u>	+	<u>EQ 37.5MG BASE</u>	<u>N021938 004</u>	Mar 31, 2009
<u>AB</u>	+!	<u>EQ 50MG BASE</u>	<u>N021938 003</u>	Jan 26, 2006

PRESCRIPTION DRUG PRODUCT LIST

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

<u>AB</u>	+	ASTELLAS	<u>EQ 0.5MG BASE</u>	<u>N050708</u>	<u>003</u>	Aug 24, 1998
<u>AB</u>	+		<u>EQ 1MG BASE</u>	<u>N050708</u>	<u>001</u>	Apr 08, 1994
<u>AB</u>	+		<u>EQ 5MG BASE</u>	<u>N050708</u>	<u>002</u>	Apr 08, 1994

TACROLIMUS

<u>AB</u>		ACCORD HLTHCARE	<u>EQ 0.5MG BASE</u>	<u>A091195</u>	<u>001</u>	Aug 31, 2011
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A091195</u>	<u>002</u>	Aug 31, 2011
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A091195</u>	<u>003</u>	Aug 31, 2011
<u>AB</u>		ALKEM LABS LTD	<u>EQ 0.5MG BASE</u>	<u>A203740</u>	<u>001</u>	Nov 12, 2020
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A203740</u>	<u>002</u>	Nov 12, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A203740</u>	<u>003</u>	Nov 12, 2020
<u>AB</u>		BELCHER	<u>EQ 0.5MG BASE</u>	<u>A206651</u>	<u>001</u>	Nov 30, 2017
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A206651</u>	<u>002</u>	Nov 30, 2017
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A206651</u>	<u>003</u>	Nov 30, 2017
<u>AB</u>		BIOCON PHARMA	<u>EQ 0.5MG BASE</u>	<u>A212297</u>	<u>001</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A212297</u>	<u>002</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A212297</u>	<u>003</u>	Nov 10, 2020
<u>AB</u>		CONCORD BIOTECH LTD	<u>EQ 1MG BASE</u>	<u>A213112</u>	<u>002</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A213112</u>	<u>003</u>	Nov 10, 2020
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 0.5MG BASE</u>	<u>A090509</u>	<u>001</u>	May 12, 2010
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090509</u>	<u>002</u>	May 12, 2010
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090509</u>	<u>003</u>	May 12, 2010
<u>AB</u>		GLENMARK PHARMS LTD	<u>EQ 0.5MG BASE</u>	<u>A206662</u>	<u>001</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A206662</u>	<u>002</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A206662</u>	<u>003</u>	Nov 10, 2020
<u>AB</u>		MYLAN	<u>EQ 0.5MG BASE</u>	<u>A090596</u>	<u>001</u>	Sep 17, 2010
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090596</u>	<u>002</u>	Sep 17, 2010
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090596</u>	<u>003</u>	Sep 17, 2010
<u>AB</u>		PANACEA	<u>EQ 0.5MG BASE</u>	<u>A090802</u>	<u>001</u>	Sep 28, 2012
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090802</u>	<u>002</u>	Sep 28, 2012
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090802</u>	<u>003</u>	Sep 28, 2012
<u>AB</u>		SANDOZ	<u>EQ 0.5MG BASE</u>	<u>A065461</u>	<u>001</u>	Aug 10, 2009
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A065461</u>	<u>002</u>	Aug 10, 2009
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A065461</u>	<u>003</u>	Aug 10, 2009
<u>AB</u>		STRIDES PHARMA	<u>EQ 0.5MG BASE</u>	<u>A090687</u>	<u>001</u>	Jul 22, 2014
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090687</u>	<u>002</u>	Jul 22, 2014
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090687</u>	<u>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

<u>AP</u>	+	ASTELLAS	<u>EQ 5MG BASE/ML</u>	<u>N050709</u>	<u>001</u>	Apr 08, 1994
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TACROLIMUS

<u>AP</u>		HOSPIRA INC	<u>EQ 5MG BASE/ML</u>	<u>A203900</u>	<u>001</u>	Aug 25, 2017
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OINTMENT; TOPICAL

PROTOPIC

<u>AB</u>	+	LEO PHARMA AS	<u>0.03%</u>	<u>N050777</u>	<u>001</u>	Dec 08, 2000
<u>AB</u>	+		<u>0.1%</u>	<u>N050777</u>	<u>002</u>	Dec 08, 2000

TACROLIMUS

<u>AB</u>		ACCORD HLTHCARE	<u>0.03%</u>	<u>A211688</u>	<u>001</u>	Jan 31, 2019
<u>AB</u>			<u>0.1%</u>	<u>A211688</u>	<u>002</u>	Jan 31, 2019
<u>AB</u>		FOUGERA PHARMS INC	<u>0.03%</u>	<u>A200744</u>	<u>001</u>	Sep 09, 2014
<u>AB</u>			<u>0.1%</u>	<u>A200744</u>	<u>002</u>	Sep 09, 2014
<u>AB</u>		GLENMARK PHARMS LTD	<u>0.1%</u>	<u>A210393</u>	<u>001</u>	Apr 16, 2018

PRESCRIPTION DRUG PRODUCT LIST

TACROLIMUS

TABLET, EXTENDED RELEASE;ORAL

ENVARUSUS 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

TADALAFIL

TABLET;ORAL

CTIALIS

<u>AB1</u>	+	LILLY	<u>2.5MG</u>	<u>N021368</u>	<u>004</u>	Jan 07, 2008
<u>AB1</u>	+		<u>5MG</u>	<u>N021368</u>	<u>001</u>	Nov 21, 2003
<u>AB1</u>	+		<u>10MG</u>	<u>N021368</u>	<u>002</u>	Nov 21, 2003
<u>AB1</u>	+		<u>20MG</u>	<u>N021368</u>	<u>003</u>	Nov 21, 2003

TADALAFIL

<u>AB1</u>		ACCORD HLTHCARE	<u>2.5MG</u>	<u>A209167</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A209167</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A209167</u>	<u>003</u>	Oct 23, 2019
<u>AB1</u>			<u>20MG</u>	<u>A209167</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>		AJANTA PHARMA LTD	<u>2.5MG</u>	<u>A209654</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A209654</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A209654</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A209654</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>		ALEMBIC PHARMS LTD	<u>2.5MG</u>	<u>A204809</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A204809</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A204809</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A204809</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>		AMNEAL PHARMS CO	<u>2.5MG</u>	<u>A209744</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A209744</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A209744</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A209744</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>		AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A206285</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A206285</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A206285</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A206285</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>		CIPLA	<u>2.5MG</u>	<u>A209539</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A209539</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A209539</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A209539</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>		DR REDDYS LABS LTD	<u>2.5MG</u>	<u>A210069</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A210069</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A210069</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A210069</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>		HETERO LABS LTD III	<u>2.5MG</u>	<u>A209908</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A209908</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A209908</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A209908</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>		LUPIN LTD	<u>2.5MG</u>	<u>A210567</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A210567</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A210567</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A210567</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>		MACLEODS PHARMS LTD	<u>2.5MG</u>	<u>A207244</u>	<u>001</u>	Oct 07, 2019
<u>AB1</u>			<u>5MG</u>	<u>A207244</u>	<u>002</u>	Oct 07, 2019
<u>AB1</u>			<u>10MG</u>	<u>A207244</u>	<u>003</u>	Oct 07, 2019
<u>AB1</u>			<u>20MG</u>	<u>A207244</u>	<u>004</u>	Oct 07, 2019
<u>AB1</u>		QILU PHARM HAINAN	<u>2.5MG</u>	<u>A210420</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A210420</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A210420</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A210420</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>		SUN PHARM	<u>2.5MG</u>	<u>A208934</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A208934</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A208934</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A208934</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>		SUNSHINE	<u>2.5MG</u>	<u>A211335</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A211335</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A211335</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A211335</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>		TEVA PHARMS USA	<u>2.5MG</u>	<u>A090141</u>	<u>001</u>	May 22, 2018
<u>AB1</u>			<u>5MG</u>	<u>A090141</u>	<u>002</u>	May 22, 2018
<u>AB1</u>			<u>10MG</u>	<u>A090141</u>	<u>003</u>	May 22, 2018
<u>AB1</u>			<u>20MG</u>	<u>A090141</u>	<u>004</u>	May 22, 2018
<u>AB1</u>		TORRENT	<u>2.5MG</u>	<u>A211839</u>	<u>001</u>	Mar 26, 2019

PRESCRIPTION DRUG PRODUCT LIST

TADALAFIL

TABLET; ORAL

TADALAFIL

<u>AB1</u>		<u>5MG</u>	<u>A211839 002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A211839 003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A211839 004</u>	Mar 26, 2019
<u>AB1</u>	UMEDICA LABS PVT LTD	<u>2.5MG</u>	<u>A211298 001</u>	Oct 23, 2020
<u>AB1</u>		<u>5MG</u>	<u>A211298 002</u>	Oct 23, 2020
<u>AB1</u>		<u>10MG</u>	<u>A211298 003</u>	Oct 23, 2020
<u>AB1</u>		<u>20MG</u>	<u>A211298 004</u>	Oct 23, 2020
<u>AB1</u>	VKT PHARMA	<u>2.5MG</u>	<u>A215556 001</u>	Nov 04, 2021
<u>AB1</u>		<u>5MG</u>	<u>A215556 002</u>	Nov 04, 2021
<u>AB1</u>		<u>10MG</u>	<u>A215556 003</u>	Nov 04, 2021
<u>AB1</u>		<u>20MG</u>	<u>A215556 004</u>	Nov 04, 2021
<u>AB1</u>	YANGLING BUCHANG	<u>2.5MG</u>	<u>A208824 001</u>	Oct 27, 2020
<u>AB1</u>		<u>5MG</u>	<u>A208824 002</u>	Oct 27, 2020
<u>AB1</u>		<u>10MG</u>	<u>A208824 003</u>	Oct 27, 2020
<u>AB1</u>		<u>20MG</u>	<u>A208824 004</u>	Oct 27, 2020
<u>AB1</u>	ZYDUS PHARMS	<u>2.5MG</u>	<u>A206693 001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A206693 002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A206693 003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A206693 004</u>	Mar 26, 2019

ADCIRCA

<u>AB2</u>	+!	ELI LILLY CO	<u>20MG</u>	<u>N022332 001</u>	May 22, 2009
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ALYO

<u>AB2</u>		TEVA PHARMS USA	<u>20MG</u>	<u>A209942 001</u>	Feb 05, 2019
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TADALAFIL

<u>AB2</u>		AJANTA PHARMA LTD	<u>20MG</u>	<u>A210392 001</u>	Feb 05, 2019
<u>AB2</u>		AUROBINDO PHARMA LTD	<u>20MG</u>	<u>A206286 001</u>	Feb 05, 2019
<u>AB2</u>		CIPLA	<u>20MG</u>	<u>A210255 001</u>	Feb 05, 2019
<u>AB2</u>		DR REDDYS LABS LTD	<u>20MG</u>	<u>A210145 001</u>	Feb 05, 2019
<u>AB2</u>		HETERO LABS LTD III	<u>20MG</u>	<u>A209907 001</u>	Feb 05, 2019
<u>AB2</u>		LUPIN LTD	<u>20MG</u>	<u>A210572 001</u>	Feb 05, 2019
<u>AB2</u>		MACLEODS PHARMS LTD	<u>20MG</u>	<u>A207290 001</u>	Oct 16, 2019
<u>AB2</u>		SUNSHINE	<u>20MG</u>	<u>A213496 001</u>	Nov 23, 2020
<u>AB2</u>		TORRENT	<u>20MG</u>	<u>A212062 001</u>	Mar 26, 2019
BX		UNICHEM	2.5MG	A209250 001	Mar 26, 2019
BX			5MG	A209250 002	Mar 26, 2019
BX			10MG	A209250 003	Mar 26, 2019
BX			20MG	A209250 004	Mar 26, 2019

TAFAMIDIS

CAPSULE; ORAL

VYNDAMAX

+!	FOLDRX PHARMS	61MG	N212161 001	May 03, 2019
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TAFAMIDIS MEGLUMINE

CAPSULE; ORAL

VYNDAQEL

+!	FOLDRX PHARMS	20MG	N211996 001	May 03, 2019
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TAFENOQUINE SUCCINATE

TABLET; ORAL

ARAKODA

+!	60 DEGREES PHARMS	EQ 100MG BASE	N210607 001	Aug 08, 2018
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KRINTAFEL

+!	GLAXOSMITHKLINE	EQ 150MG BASE	N210795 001	Jul 20, 2018
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TAFLUPROST

SOLUTION/DROPS; OPHTHALMIC

TAFLUPROST

<u>AT</u>		MICRO LABS	<u>0.0015%</u>	<u>A209051 001</u>	Aug 19, 2019
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ZIOPTAN

<u>AT</u>	+!	AKORN	<u>0.0015%</u>	<u>N202514 001</u>	Feb 10, 2012
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TALAZOPARIB TOSYLATE

CAPSULE; ORAL

TALZENNA

+	PFIZER	EQ 0.25MG BASE	N211651 001	Oct 16, 2018
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+		EQ 0.5MG BASE	N211651 003	Sep 20, 2021
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+		EQ 0.75MG BASE	N211651 004	Sep 20, 2021
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+!		EQ 1MG BASE	N211651 002	Oct 16, 2018
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PRESCRIPTION DRUG PRODUCT LIST

TALC

AEROSOL; INTRAPLEURAL

SCLEROSOL

+! SCIARRA LABS

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

+! SCIARRA LABS

5GM/BOT

N021388 001 Dec 15, 2003

TAMOXIFEN CITRATE

SOLUTION; ORAL

SOLTAMOX

+! MAYNE PHARMA INC

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** EUGIA PHARMA**EQ 10MG BASE****A213358 001** Aug 14, 2020**AB****EQ 20MG BASE****A213358 002** Aug 14, 2020**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**EQ 10MG BASE****A206694 001** Oct 27, 2017**AB****EQ 20MG BASE****A206694 002** Oct 27, 2017TAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL

FLOMAX**AB** +! SANOFI**0.4MG****N020579 001** Apr 15, 1997TAMSULOSIN HYDROCHLORIDE**AB** ALKEM LABS LTD**0.4MG****A207405 001** Aug 11, 2017**AB** ANBISON LAB**0.4MG****A211885 001** Oct 17, 2019**AB** AUROBINDO PHARMA**0.4MG****A202433 001** Apr 30, 2013

LTD

AB IMPAX LABS**0.4MG****A090377 001** Mar 02, 2010**AB** MACLEODS PHARMS LTD**0.4MG****A204645 001** Jan 20, 2017**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**0.4MG****A078225 001** Apr 27, 2010

INC

TAPENTADOL HYDROCHLORIDE

TABLET; ORAL

NUCYNTA

+ COLLEGIUM PHARM INC

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

+ COLLEGIUM PHARM INC

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

SUSPENSION; ORAL

HETLIOZ LQ

+! VANDA PHARMS INC

4MG/ML

N214517 001 Dec 01, 2020

PRESCRIPTION DRUG PRODUCT LIST

TAVABOROLE

SOLUTION; TOPICAL

KERYDIN

AB	+	!	ANACOR PHARMS INC	5%	N204427	001	Jul 07, 2014
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TAVABOROLE

AB			ALEOR	5%	A212188	001	Oct 21, 2020
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DERMACEUTICALS

AB			AMNEAL	5%	A212256	001	Nov 25, 2020
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AB			CIPLA	5%	A212224	001	Feb 09, 2021
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AB			ENCUBE	5%	A211297	001	Oct 13, 2020
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AB			LUPIN LTD	5%	A212168	001	Feb 08, 2021
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AB			PADAGIS US	5%	A211848	001	Oct 13, 2020
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AB			TARO PHARMS	5%	A212215	001	May 07, 2021
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TAZAROTENE

AEROSOL, FOAM; TOPICAL

FABIOR

+	!	MAYNE PHARMA	0.1%	N202428	001	May 11, 2012
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CREAM; TOPICAL

AVAGE

AB	+	!	ALLERGAN	0.1%	N021184	003	Sep 30, 2002
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TAZAROTENE

AB			COSETTE	0.1%	A208662	001	Dec 22, 2017
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AB			TARO PHARMS	0.1%	A208258	001	Apr 03, 2017
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TAZORAC

AB	+	!	ALLERGAN	0.1%	N021184	002	Sep 29, 2000
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+	!			0.05%	N021184	001	Sep 29, 2000
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GEL; TOPICAL

TAZORAC

+	!	ALLERGAN	0.05%	N020600	001	Jun 13, 1997
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+	!			0.1%	N020600	002	Jun 13, 1997
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LOTION; TOPICAL

ARAZLO

+	!	BAUSCH	0.045%	N211882	001	Dec 18, 2019
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TAZEMETOSTAT HYDROBROMIDE

TABLET; ORAL

TAZVERIK

+	!	EPIZYME INC	EQ 200MG BASE	N211723	001	Jan 23, 2020
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TECHNETIUM TC-99M BICISATE KIT

INJECTABLE; INJECTION

NEUROLITE

+	!	LANTHEUS MEDCL	N/A	N020256	001	Nov 23, 1994
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TECHNETIUM TC-99M EXAMETAZIME KIT

INJECTABLE; INJECTION

CERETEC

+	!	GE HEALTHCARE	N/A	N019829	001	Dec 30, 1988
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POWDER; INTRAVENOUS

DRAX EXAMETAZIME

		JUBILANT	N/A	N208870	001	Aug 17, 2017
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TECHNETIUM TC-99M MEBROFENIN KIT

INJECTABLE; INJECTION

CHOLETEC

AP	+	!	BRACCO	N/A	N018963	001	Jan 21, 1987
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TECHNETIUM TC-99M MEBROFENIN

AP			SUN PHARM INDS INC	N/A	A078242	001	Jan 29, 2008
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TECHNETIUM TC-99M MEDRONATE

INJECTABLE; INJECTION

DRAXIMAGE MDP-25

+	!	JUBILANT	N/A	N018035	002	Feb 27, 2004
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TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION

CIS-MDP

AP			SUN PHARM INDS INC	N/A	N018124	001	
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TECHNETIUM TC-99M MEDRONATE KIT

AP			CARDINAL HEALTH 414	N/A	N018107	001	
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PRESCRIPTION DRUG PRODUCT LIST

TECHNETIUM TC-99M MERTIATIDE KIT

INJECTABLE; INJECTION

TECHNESCAN MAG3

AP	+ !	CURIUM	N/A	N019882	001	Jun 15, 1990
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TECHNETIUM TC99M MERTIATIDE KIT

AP		SOMMER PHARMS II LLC	N/A	A206489	001	Feb 06, 2020
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AP		SUN PHARM INDS INC	N/A	A208994	001	Jul 12, 2019
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TECHNETIUM TC-99M OXIDRONATE KIT

INJECTABLE; INJECTION

TECHNESCAN

+ !	CURIUM	N/A	N018321	001		
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TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION

DRAXIMAGE DTPA

+ !	JUBILANT	N/A	N018511	001	Dec 29, 1989	
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TECHNETIUM TC-99M PYROPHOSPHATE KIT

INJECTABLE; INJECTION

CIS-PYRO

AP		SUN PHARM INDS INC	N/A	N019039	001	Jun 30, 1987
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TECHNESCAN PYP KIT

AP		CURIUM	N/A	N017538	001	
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TECHNETIUM TC-99M RED BLOOD CELL KIT

INJECTABLE; INJECTION

ULTRATAG

+ !	CURIUM	N/A	N019981	001	Jun 10, 1991	
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TECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE; INJECTION

CARDIOLITE

AP	+ !	LANTHEUS MEDCL	N/A	N019785	001	Dec 21, 1990
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TECHNETIUM TC 99M SESTAMIBI

AP		CARDINAL HEALTH 414	N/A	A078809	001	Apr 28, 2009
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AP		CURIUM	N/A	A078098	001	Sep 22, 2008
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AP		JUBILANT DRAXIMAGE	N/A	A078806	001	Apr 29, 2009
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AP		SUN PHARM INDS INC	10-30mCi	A079157	001	Jul 10, 2009
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TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION; INTRAVENOUS

TECHNELITE

+ !	LANTHEUS MEDCL	1-20 CI/GENERATOR	N017771	002	Feb 12, 2014	
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ULTRA-TECHNEKOW FM

+ !	CURIUM	1-19 CI/GENERATOR	N017243	003	Feb 18, 2014	
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SOLUTION; INTRAVENOUS, INTRAVESICULAR, OPHTHALMIC

RADIOGENIX SYSTEM

+	NORTHSTAR MEDICAL	30-1153mCi/GENERATOR	N202158	001	Feb 08, 2018	
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TECHNETIUM TC-99M SULFUR COLLOID KIT

SOLUTION; INJECTION, ORAL

AN-SULFUR COLLOID

+ !	SUN PHARM INDS INC	N/A	N017858	001		
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TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE; INJECTION

MYOVIEW 30ML

+ !	GE HEALTHCARE	N/A	N020372	002	Jul 07, 2005	
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TECHNETIUM TC-99M TILMANOCEPT

INJECTABLE; INJECTION

LYMPHOSEEK KIT

+ !	CARDINAL HEALTH 414	N/A	N202207	001	Mar 13, 2013	
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TECOVIRIMAT

CAPSULE; ORAL

TPOXX

+ !	SIGA TECHNOLOGIES	200MG	N208627	001	Jul 13, 2018	
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TEDIZOLID PHOSPHATE

POWDER; INTRAVENOUS

SIVEXTRO

+ !	CUBIST PHARMS LLC	200MG/VIAL	N205436	001	Jun 20, 2014	
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TABLET; ORAL

SIVEXTRO

+ !	CUBIST PHARMS LLC	200MG	N205435	001	Jun 20, 2014	
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PRESCRIPTION DRUG PRODUCT LIST

TEDUGLUTIDE RECOMBINANT

POWDER; SUBCUTANEOUS

GATTEX KIT

+! NPS PHARMS INC

5MG/VIAL

N203441 001 Dec 21, 2012

TEGASEROD MALEATE

TABLET; ORAL

ZELNORM

+! ALFASIGMA

EQ 6MG BASE

N021200 002 Jul 24, 2002

TELAVANCIN HYDROCHLORIDE

POWDER; INTRAVENOUS

VIBATIV

+! CUMBERLAND PHARMS

EQ 750MG BASE/VIAL

N022110 002 Sep 11, 2009

TELMISARTAN

TABLET; ORAL

MICARDISAB + BOEHRINGER
INGELHEIM20MGN020850 003 Apr 04, 2000AB +40MGN020850 001 Nov 10, 1998AB +!80MGN020850 002 Nov 10, 1998TELMISARTANAB ALEMBIC PHARMS LTD20MGA202130 001 Jul 07, 2014AB40MGA202130 002 Jul 07, 2014AB80MGA202130 003 Jul 07, 2014AB AMNEAL PHARMS20MGA204415 001 Sep 08, 2015AB40MGA204415 002 Sep 08, 2015AB80MGA204415 003 Sep 08, 2015AB AUROBINDO PHARMA
LTD20MGA206511 001 Sep 03, 2015AB40MGA206511 002 Sep 03, 2015AB80MGA206511 003 Sep 03, 2015AB CADILA PHARMS LTD20MGA208605 001 Jul 25, 2017AB40MGA208605 002 Jul 25, 2017AB80MGA208605 003 Jul 25, 2017AB CELLTRION20MGA078710 001 Jan 08, 2014AB40MGA078710 002 Jan 08, 2014AB80MGA078710 003 Jan 08, 2014AB GLENMARK PHARMS LTD20MGA090032 001 Jul 07, 2014AB40MGA090032 002 Jul 07, 2014AB80MGA090032 003 Jul 07, 2014AB HETERO LABS LTD V20MGA205901 001 Apr 22, 2016AB40MGA205901 002 Apr 22, 2016AB80MGA205901 003 Apr 22, 2016AB INVENTIA20MGA205150 001 Oct 30, 2015AB40MGA205150 002 Oct 30, 2015AB80MGA205150 003 Oct 30, 2015AB MICRO LABS20MGA207016 001 Oct 03, 2017AB40MGA207016 002 Oct 03, 2017AB80MGA207016 003 Oct 03, 2017AB MYLAN20MGA202397 001 Jul 07, 2014AB40MGA202397 002 Jul 07, 2014AB80MGA202397 003 Jul 07, 2014AB PRINSTON INC20MGA207882 001 May 03, 2017AB40MGA207882 002 May 03, 2017AB80MGA207882 003 May 03, 2017AB SANDOZ INC20MGA203867 001 Nov 03, 2014AB40MGA203867 002 Nov 03, 2014AB80MGA203867 003 Nov 03, 2014AB ZYDUS PHARMS20MGA203325 001 Aug 26, 2014AB40MGA203325 002 Aug 26, 2014AB80MGA203325 003 Aug 26, 2014TELOTRISTAT ETIPRATE

TABLET; ORAL

XERMELO

+! TERSERA

EQ 250MG BASE

N208794 001 Feb 28, 2017

PRESCRIPTION DRUG PRODUCT LIST

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>		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

TEMOZOLOMIDE

<u>AB</u>		ACCORD HLTHCARE	<u>5MG</u>	<u>A201528</u>	<u>001</u>	Feb 27, 2017
<u>AB</u>			<u>20MG</u>	<u>A201528</u>	<u>002</u>	Feb 27, 2017
<u>AB</u>			<u>100MG</u>	<u>A201528</u>	<u>003</u>	Feb 27, 2017
<u>AB</u>			<u>140MG</u>	<u>A201528</u>	<u>004</u>	Feb 27, 2017
<u>AB</u>			<u>180MG</u>	<u>A201528</u>	<u>005</u>	Feb 27, 2017
<u>AB</u>			<u>250MG</u>	<u>A201528</u>	<u>006</u>	Feb 27, 2017
<u>AB</u>		AMNEAL PHARMS	<u>5MG</u>	<u>A203691</u>	<u>001</u>	May 08, 2015
<u>AB</u>			<u>20MG</u>	<u>A203691</u>	<u>002</u>	May 08, 2015
<u>AB</u>			<u>100MG</u>	<u>A203691</u>	<u>003</u>	May 08, 2015
<u>AB</u>			<u>140MG</u>	<u>A203691</u>	<u>004</u>	May 08, 2015
<u>AB</u>			<u>180MG</u>	<u>A203691</u>	<u>005</u>	May 08, 2015
<u>AB</u>			<u>250MG</u>	<u>A203691</u>	<u>006</u>	May 08, 2015
<u>AB</u>		ANI PHARMS	<u>5MG</u>	<u>A203490</u>	<u>001</u>	Jul 13, 2016
<u>AB</u>			<u>20MG</u>	<u>A203490</u>	<u>002</u>	Jul 13, 2016
<u>AB</u>			<u>100MG</u>	<u>A203490</u>	<u>003</u>	Jul 13, 2016
<u>AB</u>			<u>140MG</u>	<u>A203490</u>	<u>004</u>	Jul 13, 2016
<u>AB</u>			<u>180MG</u>	<u>A203490</u>	<u>005</u>	Jul 13, 2016
<u>AB</u>			<u>250MG</u>	<u>A203490</u>	<u>006</u>	Jul 13, 2016
<u>AB</u>		CHARTWELL	<u>5MG</u>	<u>A206413</u>	<u>001</u>	Apr 12, 2016
<u>AB</u>			<u>20MG</u>	<u>A206413</u>	<u>002</u>	Apr 12, 2016
<u>AB</u>			<u>100MG</u>	<u>A206413</u>	<u>003</u>	Apr 12, 2016
<u>AB</u>			<u>140MG</u>	<u>A206413</u>	<u>004</u>	Apr 12, 2016
<u>AB</u>			<u>180MG</u>	<u>A206413</u>	<u>005</u>	Apr 12, 2016
<u>AB</u>			<u>250MG</u>	<u>A206413</u>	<u>006</u>	Apr 12, 2016
<u>AB</u>		CHEMI SPA	<u>5MG</u>	<u>A204639</u>	<u>001</u>	Nov 23, 2016
<u>AB</u>			<u>20MG</u>	<u>A204639</u>	<u>002</u>	Nov 23, 2016
<u>AB</u>			<u>100MG</u>	<u>A204639</u>	<u>003</u>	Nov 23, 2016
<u>AB</u>			<u>140MG</u>	<u>A204639</u>	<u>004</u>	Nov 23, 2016
<u>AB</u>			<u>180MG</u>	<u>A204639</u>	<u>005</u>	Nov 23, 2016
<u>AB</u>			<u>250MG</u>	<u>A204639</u>	<u>006</u>	Nov 23, 2016
<u>AB</u>		DEVA HOLDING AS	<u>5MG</u>	<u>A207658</u>	<u>001</u>	Apr 26, 2017

PRESCRIPTION DRUG PRODUCT LIST

TEMOZOLOMIDE

CAPSULE; ORAL

TEMOZOLOMIDE

<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>	NIVAGEN PHARMS INC	<u>5MG</u>	<u>A213328 001</u>	Nov 23, 2021
<u>AB</u>		<u>20MG</u>	<u>A213328 002</u>	Nov 23, 2021
<u>AB</u>		<u>100MG</u>	<u>A213328 003</u>	Nov 23, 2021
<u>AB</u>		<u>140MG</u>	<u>A213328 004</u>	Nov 23, 2021
<u>AB</u>		<u>180MG</u>	<u>A213328 005</u>	Nov 23, 2021
<u>AB</u>		<u>250MG</u>	<u>A213328 006</u>	Nov 23, 2021
<u>AB</u>	RISING	<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 PHARM	<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>	ZYDUS PHARMS	<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>	GLAND PHARMA LTD	<u>25MG/ML (25MG/ML)</u>	<u>A207383 001</u>	Aug 16, 2019
<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>	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>	QILU	<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>AB</u>	+ GILEAD SCIENCES INC	<u>150MG</u>	<u>N021356 002</u>	Jan 18, 2012

PRESCRIPTION DRUG PRODUCT LIST

TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

VIREAD

<u>AB</u>	+		<u>200MG</u>	<u>N021356</u>	<u>003</u>	Jan 18, 2012
<u>AB</u>	+		<u>250MG</u>	<u>N021356</u>	<u>004</u>	Jan 18, 2012
<u>AB</u>	+	!	<u>300MG</u>	<u>N021356</u>	<u>001</u>	Oct 26, 2001

TEPOTINIB HYDROCHLORIDE

TABLET; ORAL

TEPMETKO

+	!	EMD SERONO INC	EQ 225MG BASE	N214096	001	Feb 03, 2021
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TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

TERAZOSIN HYDROCHLORIDE

<u>AB</u>		APNAR PHARMA LP	<u>EQ 1MG BASE</u>	<u>A074823</u>	<u>001</u>	Mar 30, 1998
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A074823</u>	<u>002</u>	Mar 30, 1998
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A074823</u>	<u>003</u>	Mar 30, 1998
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A074823</u>	<u>004</u>	Mar 30, 1998
<u>AB</u>		HERITAGE PHARMA AVET	<u>EQ 1MG BASE</u>	<u>A075614</u>	<u>002</u>	Jan 30, 2001
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A075614</u>	<u>001</u>	Jan 30, 2001
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A075614</u>	<u>003</u>	Jan 30, 2001
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A075614</u>	<u>004</u>	Jan 30, 2001
<u>AB</u>	!	JUBILANT CADISTA	<u>EQ 1MG BASE</u>	<u>A075317</u>	<u>001</u>	Dec 20, 2004
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A075317</u>	<u>002</u>	Dec 20, 2004
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A075317</u>	<u>003</u>	Dec 20, 2004
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A075317</u>	<u>004</u>	Dec 20, 2004

TERBINAFINE HYDROCHLORIDE

TABLET; ORAL

TERBINAFINE HYDROCHLORIDE

<u>AB</u>	!	AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A078297</u>	<u>001</u>	Jul 02, 2007
<u>AB</u>		BRECKENRIDGE PHARM	<u>EQ 250MG BASE</u>	<u>A077714</u>	<u>001</u>	Jun 04, 2010
<u>AB</u>		CIPLA	<u>EQ 250MG BASE</u>	<u>A077137</u>	<u>001</u>	Jul 02, 2007
<u>AB</u>		DR REDDYS LABS INC	<u>EQ 250MG BASE</u>	<u>A076390</u>	<u>001</u>	Jul 02, 2007
<u>AB</u>		EMED MEDCL	<u>EQ 250MG BASE</u>	<u>A077919</u>	<u>001</u>	Jul 02, 2007
<u>AB</u>		GLENMARK GENERICS	<u>EQ 250MG BASE</u>	<u>A078157</u>	<u>001</u>	Jul 02, 2007
<u>AB</u>		HERITAGE PHARMA AVET	<u>EQ 250MG BASE</u>	<u>A076377</u>	<u>001</u>	Jul 02, 2007
<u>AB</u>		INVAGEN PHARMS	<u>EQ 250MG BASE</u>	<u>A077533</u>	<u>001</u>	Jul 02, 2007
<u>AB</u>		ORBION PHARMS	<u>EQ 250MG BASE</u>	<u>A078163</u>	<u>001</u>	Jul 02, 2007

TERBUTALINE SULFATE

INJECTABLE; INJECTION

TERBUTALINE SULFATE

<u>AP</u>		AKORN	<u>1MG/ML</u>	<u>A078151</u>	<u>001</u>	Jan 07, 2008
<u>AP</u>	!	ATHENEX INC	<u>1MG/ML</u>	<u>A076770</u>	<u>001</u>	Apr 23, 2004
<u>AP</u>		FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A076887</u>	<u>001</u>	May 26, 2004
<u>AP</u>		HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A078630</u>	<u>001</u>	May 20, 2009
<u>AP</u>		UNITED BIOMEDCL	<u>1MG/ML</u>	<u>A200122</u>	<u>001</u>	Nov 08, 2013

TABLET; ORAL

BRETHINE

<u>AB</u>	+	ANI PHARMS	<u>2.5MG</u>	<u>N017849</u>	<u>001</u>	
<u>AB</u>	+		<u>5MG</u>	<u>N017849</u>	<u>002</u>	
<u>AB</u>			<u>2.5MG</u>	<u>A075877</u>	<u>001</u>	Jun 26, 2001
<u>AB</u>			<u>5MG</u>	<u>A075877</u>	<u>002</u>	Jun 26, 2001
<u>AB</u>		LANNETT CO INC	<u>2.5MG</u>	<u>A077152</u>	<u>001</u>	Mar 25, 2005
<u>AB</u>	!		<u>5MG</u>	<u>A077152</u>	<u>002</u>	Mar 25, 2005
<u>AB</u>		TWI PHARMS	<u>2.5MG</u>	<u>A211832</u>	<u>001</u>	Jun 19, 2020
<u>AB</u>			<u>5MG</u>	<u>A211832</u>	<u>002</u>	Jun 19, 2020

TERCONAZOLE

CREAM; VAGINAL

TERCONAZOLE

<u>AB</u>		FOUGERA PHARMS	<u>0.4%</u>	<u>A076712</u>	<u>001</u>	Feb 18, 2005
<u>AB</u>	!	TARO	<u>0.4%</u>	<u>A076043</u>	<u>001</u>	Jan 19, 2005
BX	+	FOUGERA PHARMS INC	0.8%	N021735	001	Oct 01, 2004
BX	!	TARO	0.8%	A075953	001	Apr 06, 2004

SUPPOSITORY; VAGINAL

TERCONAZOLE

<u>AB</u>	!	PADAGIS ISRAEL	<u>80MG</u>	<u>A077149</u>	<u>001</u>	Mar 17, 2006
<u>AB</u>		TARO	<u>80MG</u>	<u>A077553</u>	<u>001</u>	Mar 09, 2007

PRESCRIPTION DRUG PRODUCT LIST

TERIFLUNOMIDE

TABLET; ORAL

AUBAGIO

AB	+	SANOFI AVENTIS US	7MG	N202992 001	Sep 12, 2012
AB	+	!	14MG	N202992 002	Sep 12, 2012

TERIFLUNOMIDE

AB		ACCORD HLTHCARE	7MG	A209690 001	Jan 07, 2019
AB			14MG	A209690 002	Jan 07, 2019
AB		ALEMBIC PHARMS LTD	7MG	A209572 001	Apr 19, 2019
AB			14MG	A209572 002	Apr 19, 2019
AB		GLENMARK PHARMS	7MG	A209663 001	Nov 15, 2018
AB			14MG	A209663 002	Nov 15, 2018
AB		SANDOZ INC	7MG	A209710 001	Jan 03, 2019
AB			14MG	A209710 002	Jan 03, 2019

TERIPARATIDE

SOLUTION; SUBCUTANEOUS

BONSITY

		ALVOGEN	0.62MG/2.48ML (0.25MG/ML)	N211939 001	Oct 04, 2019
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FORTEO

		!	LILLY	0.6MG/2.4ML (0.25MG/ML)	N021318 002	Jun 25, 2008
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TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL

ANDRODERM

		!	ALLERGAN	2MG/24HR	N020489 003	Oct 20, 2011
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		!		4MG/24HR	N020489 004	Oct 20, 2011
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GEL; TRANSDERMAL

ANDROGEL

AB1	+	ABEVIE	25MG/2.5GM PACKET	N021015 001	Feb 28, 2000
AB1	+	!	50MG/5GM PACKET	N021015 002	Feb 28, 2000

TESTOSTERONE

AB1		ACTAVIS LABS UT INC	25MG/2.5GM PACKET	A076737 001	Jan 27, 2006
AB1			50MG/5GM PACKET	A076737 002	Jan 27, 2006
AB1		ENCUBE	50MG/5GM PACKET	A212984 001	Nov 09, 2021
AB1		STRIDES PHARMA	25MG/2.5GM PACKET	A076744 001	May 23, 2007
AB1			50MG/5GM PACKET	A076744 002	May 23, 2007

ANDROGEL

AB2	+	ABEVIE	1.62% (20.25MG/1.25GM PACKET)	N022309 002	Sep 07, 2012
AB2	+	!	1.62% (40.5MG/2.5GM PACKET)	N022309 003	Sep 07, 2012

TESTIM

AB2	+	AUXILIUM PHARMS LLC	50MG/5GM PACKET	N021454 001	Oct 31, 2002
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TESTOSTERONE

AB2		ACTAVIS LABS UT INC	1.62% (20.25MG/1.25GM PACKET)	A204570 002	Jul 17, 2020
AB2			1.62% (40.5MG/2.5GM PACKET)	A204570 003	Jul 17, 2020
AB2			50MG/5GM PACKET	A091073 001	Sep 18, 2017
AB2		PADAGIS ISRAEL	1.62% (20.25MG/1.25GM PACKET)	A205781 001	Jul 12, 2017
AB2			1.62% (40.5MG/2.5GM PACKET)	A205781 002	Jul 12, 2017

VOGELXO

AB2		UPSHER SMITH LABS	50MG/5GM PACKET	N204399 002	Jun 04, 2014
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GEL, METERED; NASAL

NATESTO

		ACERUS	5.5MG/0.122GM ACTUATION	N205488 001	May 28, 2014
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GEL, METERED; TRANSDERMAL

ANDROGEL

AB	+	!	ABEVIE	1.62% (20.25MG/1.25GM ACTUATION)	N022309 001	Apr 29, 2011
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FORTESTA

AB	+	!	ENDO PHARMS	10MG/0.5GM ACTUATION	N021463 001	Dec 29, 2010
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TESTOSTERONE

AB		ACTAVIS LABS UT INC	1.62% (20.25MG/1.25GM ACTUATION)	A204570 001	Apr 10, 2019
AB			10MG/0.5GM ACTUATION	A204571 001	Aug 05, 2015
AB		AMNEAL	1.62% (20.25MG/1.25GM ACTUATION)	A207373 001	Apr 10, 2019
AB		DR REDDYS	1.62% (20.25MG/1.25GM ACTUATION)	A208620 001	Apr 10, 2019
AB		LUPIN	1.62% (20.25MG/1.25GM ACTUATION)	A208560 001	Apr 10, 2019
AB		PADAGIS ISRAEL	1.62% (20.25MG/1.25GM ACTUATION)	A204268 001	Aug 04, 2015
AB		TWI PHARMS	1.62% (20.25MG/1.25GM ACTUATION)	A209390 001	Sep 23, 2019
AB		XIROMED	1.62% (20.25MG/1.25GM ACTUATION)	A210835 001	Apr 16, 2020
BX	!	ACTAVIS LABS UT INC	12.5MG/1.25GM ACTUATION	A076737 003	Mar 09, 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	

PRESCRIPTION DRUG PRODUCT LIST

TESTOSTERONE

SOLUTION, METERED; TRANSDERMAL

TESTOSTERONE

AT	ACTAVIS LABS UT INC	30MG/1.5ML ACTUATION	A205328 001	Aug 07, 2017
AT	CIPLA	30MG/1.5ML ACTUATION	A209533 001	Jan 29, 2018
AT	DASH PHARMS	30MG/1.5ML ACTUATION	A212301 001	Jan 11, 2021
AT	LUPIN LTD	30MG/1.5ML ACTUATION	A208061 001	Oct 23, 2017
AT	! PADAGIS ISRAEL	30MG/1.5ML ACTUATION	A204255 001	Feb 28, 2017
AT	TWI PHARMS	30MG/1.5ML ACTUATION	A209836 001	Sep 03, 2021

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTOSTERONE

AO	! PFIZER	100MG/ML	A085635 002	
AO	! PFIZER	200MG/ML	A085635 003	

TESTOSTERONE CYPIONATE

AO	AM REGENT	200MG/ML	A207742 001	Jun 16, 2017
AO	CIPLA	100MG/ML	A210362 001	Jun 19, 2018
AO		200MG/ML	A210362 002	Jun 19, 2018
AO	HIKMA	100MG/ML	A090387 001	Jul 15, 2010
AO		200MG/ML	A090387 002	Jul 15, 2010
AO	HIKMA FARMACEUTICA	200MG/ML	A091244 001	May 01, 2012
AO	PADAGIS US	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	WILSHIRE PHARMS INC	200MG/ML	A206368 001	Apr 24, 2019

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
	! ANTARES PHARMA INC	75MG/0.5ML (75MG/0.5ML)	N209863 002	Sep 28, 2018
	! ANTARES PHARMA INC	100MG/0.5ML (100MG/0.5ML)	N209863 003	Sep 28, 2018

TESTOSTERONE UNDECANOATE

CAPSULE; ORAL

JATENZO

	+ CLARUS	158MG	N206089 001	Mar 27, 2019
	+ CLARUS	198MG	N206089 002	Mar 27, 2019
	! CLARUS	237MG	N206089 003	Mar 27, 2019

INJECTABLE; INTRAMUSCULAR

AVEED

	! ENDO PHARMS INC	750MG/3ML (250MG/ML)	N022219 001	Mar 05, 2014
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TETRABENAZINE

TABLET; ORAL

TETRABENAZINE

AB	ACTAVIS LABS FL INC	25MG	A206686 001	Jul 07, 2017
AB	APOTEX	12.5MG	A206093 001	Mar 17, 2020
AB		25MG	A206093 002	Mar 17, 2020
AB	BIONPHARMA INC	12.5MG	A208826 001	Dec 18, 2017
AB		25MG	A208826 002	Dec 18, 2017
AB	DR REDDYS	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	HIKMA	12.5MG	A209739 001	Apr 08, 2019
AB		25MG	A209739 002	Apr 08, 2019
AB	LUPIN LTD	12.5MG	A210544 001	Apr 20, 2018
AB		25MG	A210544 002	Apr 20, 2018
AB	MYLAN	12.5MG	A207682 001	Jan 31, 2017
AB		25MG	A207682 002	Jan 31, 2017
AB	PIRAMAL HLTHCARE UK	12.5MG	A213316 001	Jan 22, 2020
AB		25MG	A213316 002	Jan 22, 2020
AB	SUN PHARM	12.5MG	A206129 001	Aug 17, 2015
AB		25MG	A206129 002	Aug 17, 2015

PRESCRIPTION DRUG PRODUCT LIST

TETRABENAZINE

TABLET; ORAL

XENAZINE

AB	+	BAUSCH	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
+	!	BAUSCH LOMB IRELAND	0.5%	N210821 001	Mar 12, 2019

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

ACHROMYCIN V

AB	+	AVET	250MG	N050278 003	
AB	+	!	500MG	N050278 001	

TETRACYCLINE HYDROCHLORIDE

AB		AMNEAL PHARMS NY	250MG	A210674 001	Sep 18, 2018
AB			500MG	A210674 002	Sep 18, 2018
AB		BRECKENRIDGE	250MG	A210662 001	Nov 07, 2018
AB			500MG	A210662 002	Nov 07, 2018
AB		CHARTWELL TETRA	250MG	A062752 001	Aug 12, 1988
AB			500MG	A062752 002	Aug 12, 1988
AB		STRIDES PHARMA	250MG	A212635 001	Mar 03, 2020
AB			500MG	A212635 002	Mar 03, 2020
AB		WATSON LABS	250MG	A061837 001	
AB			500MG	A061837 002	

TETRAHYDROZOLINE HYDROCHLORIDE

SOLUTION; NASAL

TYZINE

!		FOUGERA PHARMS	0.05%	A086576 002	
			0.1%	A086576 001	

SPRAY; NASAL

TYZINE

!		FOUGERA PHARMS	0.1%	A086576 003	
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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

AP	+	!	CURIUM	1mCi/ML	N018150 001
AP	+	!	LANTHEUS MEDCL	1mCi/ML	N017806 001

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
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THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER

+	!	B BRAUN	80MG/100ML	N019826 002	Aug 14, 1992
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THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER

+	!	B BRAUN	160MG/100ML	N019826 003	Aug 14, 1992
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THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER

+	!	B BRAUN	320MG/100ML	N019826 006	Aug 14, 1992
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SOLUTION; ORAL

THEOPHYLLINE

AA	!	LANNETT CO INC	80MG/15ML	A091156 001	Apr 13, 2011
AA	!	TRIS PHARMA INC	80MG/15ML	A091586 001	Jun 15, 2012

SOLUTION, ELIXIR; ORAL

ELIXOPHYLLIN

AA	+	!	NOSTRUM LABS INC	80MG/15ML	A085186 001
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PRESCRIPTION DRUG PRODUCT LIST

THEOPHYLLINE

SOLUTION, ELIXIR;ORAL

THEOPHYLLINE

AA	PHARM ASSOC	80MG/15ML	A206344 001	Dec 16, 2016
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TABLET, EXTENDED RELEASE;ORAL

THEOPHYLLINE

AB	ALEMBIC PHARMS LTD	300MG	A090430 001	Oct 27, 2010
AB	!	450MG	A090430 002	Oct 27, 2010
AB	GLENMARK GENERICS	400MG	A090355 001	Jul 13, 2010
AB		600MG	A090355 002	Jul 13, 2010
AB	GLENMARK PHARMS LTD	300MG	A212184 001	Jun 03, 2021
AB		450MG	A212184 002	Jun 03, 2021
AB	HERITAGE PHARMA AVET	300MG	A089763 001	Apr 30, 1990
AB		450MG	A081236 001	Nov 09, 1992
AB	NOSTRUM LABS INC	400MG	A040560 003	Apr 21, 2006
AB	!	600MG	A040560 002	Apr 21, 2006
AB	+ RHODES PHARMS	400MG	A040086 002	Sep 01, 1982
AB		600MG	A040086 001	Apr 15, 1996

THIAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

THIAMINE HYDROCHLORIDE

AP	+!	FRESENIUS KABI USA	100MG/ML	A080556 001
AP		MYLAN INSTITUTIONAL	100MG/ML	A091623 001
AP		SAGENT PHARMS INC	100MG/ML	A206106 001
AP		WEST-WARD PHARMS INT	100MG/ML	A080575 001

THIOGUANINE

TABLET;ORAL

THIOGUANINE

+	!	ASPEN GLOBAL INC	40MG	N012429 001
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THIORIDAZINE HYDROCHLORIDE

TABLET;ORAL

THIORIDAZINE HYDROCHLORIDE

		MYLAN	10MG	A088004 002	Mar 15, 1983
			25MG	A088004 003	Mar 15, 1983
			50MG	A088004 004	Mar 15, 1983
	!		100MG	A088004 001	Nov 18, 1983

THIOTEPA

INJECTABLE; INJECTION

THIOTEPA

AP		BELOTECA INC	15MG/VIAL	A211831 001	Dec 08, 2021
AP		DR REDDYS	15MG/VIAL	A210337 001	May 04, 2018
AP		JIANGSU PHARMS	15MG/VIAL	A209150 001	May 04, 2018
AP		STI PHARMA LLC	15MG/VIAL	A208242 001	Jan 10, 2020
AP	!	WEST-WARD PHARMS INT	15MG/VIAL	A075547 001	Apr 02, 2001

POWDER; INTRACAVITARY, INTRAVENOUS, INTRAVESICAL

TEPADINA

AP	+!	ADIENNE SA	15MG/VIAL	N208264 001	Jan 26, 2017
AP	+!		100MG/VIAL	N208264 002	Jan 26, 2017

THIOTEPA

AP		GLAND PHARMA LTD	15MG/VIAL	A214222 001	Mar 08, 2021
AP			100MG/VIAL	A214222 002	Jan 03, 2022
AP		MSN	15MG/VIAL	A213049 001	Mar 04, 2020
AP			100MG/VIAL	A213049 002	Mar 04, 2020

THIOTHIXENE

CAPSULE;ORAL

THIOTHIXENE

AB		MYLAN	1MG	A071093 002	Jun 23, 1987
AB			2MG	A071093 003	Jun 23, 1987
AB			5MG	A071093 004	Jun 23, 1987
AB			10MG	A071093 001	Jun 23, 1987
AB		NOVITIUM PHARMA	1MG	A211642 001	Apr 05, 2019
AB			2MG	A211642 002	Apr 05, 2019
AB	!		5MG	A211642 003	Apr 05, 2019
AB			10MG	A211642 004	Apr 05, 2019

PRESCRIPTION DRUG PRODUCT LIST

TIAGABINE HYDROCHLORIDE

TABLET; ORAL

GABITRIL

<u>AB</u>	+	CEPHALON	<u>2MG</u>	<u>N020646</u>	<u>005</u>	Apr 16, 1999
<u>AB</u>	+	!	<u>4MG</u>	<u>N020646</u>	<u>001</u>	Sep 30, 1997
<u>AB</u>	+		<u>12MG</u>	<u>N020646</u>	<u>002</u>	Sep 30, 1997
<u>AB</u>	+		<u>16MG</u>	<u>N020646</u>	<u>003</u>	Sep 30, 1997

TIAGABINE HYDROCHLORIDE

<u>AB</u>		AMNEAL PHARMS CO	<u>2MG</u>	<u>A208181</u>	<u>001</u>	Dec 08, 2017
<u>AB</u>			<u>4MG</u>	<u>A208181</u>	<u>002</u>	Dec 08, 2017
<u>AB</u>			<u>12MG</u>	<u>A208181</u>	<u>003</u>	Dec 08, 2017
<u>AB</u>			<u>16MG</u>	<u>A208181</u>	<u>004</u>	Dec 08, 2017
<u>AB</u>		MSN	<u>2MG</u>	<u>A214816</u>	<u>001</u>	Nov 16, 2021
<u>AB</u>			<u>4MG</u>	<u>A214816</u>	<u>002</u>	Nov 16, 2021
<u>AB</u>			<u>12MG</u>	<u>A214816</u>	<u>003</u>	Nov 16, 2021
<u>AB</u>			<u>16MG</u>	<u>A214816</u>	<u>004</u>	Nov 16, 2021
<u>AB</u>		SUN PHARM INDS	<u>2MG</u>	<u>A077555</u>	<u>001</u>	Nov 04, 2011
<u>AB</u>			<u>4MG</u>	<u>A077555</u>	<u>002</u>	Nov 04, 2011

TICAGRELOR

TABLET; ORAL

BRILINTA

<u>AB</u>	+	ASTRAZENECA	<u>60MG</u>	<u>N022433</u>	<u>002</u>	Sep 03, 2015
<u>AB</u>	+	!	<u>90MG</u>	<u>N022433</u>	<u>001</u>	Jul 20, 2011

TICAGRELOR

<u>AB</u>		HISUN PHARM HANGZHOU	<u>90MG</u>	<u>A208575</u>	<u>001</u>	Jan 23, 2019
<u>AB</u>		MYLAN	<u>60MG</u>	<u>A208597</u>	<u>001</u>	Jul 09, 2021
<u>AB</u>			<u>90MG</u>	<u>A208597</u>	<u>002</u>	Jul 09, 2021

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLOPIDINE HYDROCHLORIDE

<u>AB</u>		APOTEX	<u>250MG</u>	<u>A075089</u>	<u>001</u>	Jul 01, 1999
<u>AB</u>		!	<u>250MG</u>	<u>A075149</u>	<u>001</u>	Aug 20, 1999

TIGECYCLINE

POWDER; INTRAVENOUS

TIGECYCLINE

<u>AP</u>		AMNEAL	<u>50MG/VIAL</u>	<u>N211158</u>	<u>001</u>	Aug 02, 2018
<u>AP</u>		APOTEX	<u>50MG/VIAL</u>	<u>A204439</u>	<u>001</u>	Dec 21, 2018
<u>AP</u>		EUGIA PHARMA	<u>50MG/VIAL</u>	<u>A206335</u>	<u>001</u>	Jun 11, 2019
<u>AP</u>	+	FRESENIUS KABI USA	<u>50MG/VIAL</u>	<u>N205645</u>	<u>001</u>	Dec 01, 2016
<u>AP</u>		MEITHEAL	<u>50MG/VIAL</u>	<u>A214020</u>	<u>001</u>	May 13, 2021
<u>AP</u>		SANDOZ INC	<u>50MG/VIAL</u>	<u>A091620</u>	<u>001</u>	May 27, 2015
<u>AP</u>		XELLIA PHARMS APS	<u>50MG/VIAL</u>	<u>A205722</u>	<u>001</u>	Oct 18, 2019

TYGACIL

<u>AP</u>	+	PF PRISM CV	<u>50MG/VIAL</u>	<u>N021821</u>	<u>001</u>	Jun 15, 2005
		TIGECYCLINE				
		ACCORD HLTHCARE INC	<u>50MG/VIAL</u>	<u>N208744</u>	<u>001</u>	Jan 18, 2018

TIMOLOL

SOLUTION/DROPS; OPHTHALMIC

BETIMOL

<u>AT</u>	+	AKORN	<u>EQ 0.25% BASE</u>	<u>N020439</u>	<u>001</u>	Mar 31, 1995
<u>AT</u>	+	!	<u>EQ 0.5% BASE</u>	<u>N020439</u>	<u>002</u>	Mar 31, 1995

TIMOLOL

<u>AT</u>		AKORN	<u>EQ 0.25% BASE</u>	<u>A205309</u>	<u>001</u>	Sep 30, 2016
<u>AT</u>			<u>EQ 0.5% BASE</u>	<u>A205309</u>	<u>002</u>	Sep 30, 2016

TIMOLOL MALEATE

SOLUTION, GEL FORMING/DROPS; OPHTHALMIC

TIMOLOL MALEATE

<u>AB</u>	!	ALEMBIC PHARMS LTD	<u>EQ 0.25% BASE</u>	<u>A212942</u>	<u>001</u>	Oct 22, 2020
<u>AB</u>	!		<u>EQ 0.5% BASE</u>	<u>A212942</u>	<u>002</u>	Oct 22, 2020
<u>AB</u>	+	SANDOZ INC	<u>EQ 0.25% BASE</u>	<u>N020963</u>	<u>001</u>	Oct 21, 1998
<u>AB</u>	+	!	<u>EQ 0.5% BASE</u>	<u>N020963</u>	<u>002</u>	Oct 21, 1998

TIMOPTIC-XE

<u>AB</u>	+	VALEANT PHARMS LLC	<u>EQ 0.25% BASE</u>	<u>N020330</u>	<u>001</u>	Nov 04, 1993
<u>AB</u>	+		<u>EQ 0.5% BASE</u>	<u>N020330</u>	<u>002</u>	Nov 04, 1993

SOLUTION/DROPS; OPHTHALMIC

TIMOLOL MALEATE

<u>AT1</u>		AKORN	<u>EQ 0.5% BASE</u>	<u>A074466</u>	<u>001</u>	Mar 25, 1997
<u>AT1</u>			<u>EQ 0.5% BASE</u>	<u>A074516</u>	<u>001</u>	Mar 25, 1997
<u>AT1</u>			<u>EQ 0.5% BASE</u>	<u>A075163</u>	<u>001</u>	Sep 10, 2002

PRESCRIPTION DRUG PRODUCT LIST

TIMOLOL MALEATE

SOLUTION/DROPS;OPHTHALMIC

TIMOLOL MALEATE

<u>AT1</u>	BAUSCH AND LOMB	<u>EQ 0.25% BASE</u>	<u>A074778 001</u>	Mar 25, 1997
<u>AT1</u>		<u>EQ 0.5% BASE</u>	<u>A074776 001</u>	Mar 25, 1997
<u>AT1</u>	FDC LTD	<u>EQ 0.25% BASE</u>	<u>A077259 001</u>	Apr 30, 2008
<u>AT1</u>		<u>EQ 0.5% BASE</u>	<u>A077259 002</u>	Apr 30, 2008
<u>AT1</u>	MANKIND PHARMA	<u>EQ 0.25% BASE</u>	<u>A078771 001</u>	Sep 28, 2009
<u>AT1</u>		<u>EQ 0.5% BASE</u>	<u>A078771 002</u>	Sep 28, 2009
<u>AT1</u>	PACIFIC PHARMA	<u>EQ 0.25% BASE</u>	<u>A074746 001</u>	Mar 25, 1997
<u>AT1</u>		<u>EQ 0.5% BASE</u>	<u>A074747 001</u>	Mar 25, 1997
<u>AT1</u>	! SANDOZ INC	<u>EQ 0.25% BASE</u>	<u>A074261 001</u>	Apr 28, 1995
<u>AT1</u>	!	<u>EQ 0.5% BASE</u>	<u>A074262 001</u>	Apr 28, 1995

TIMOPTIC

<u>AT1</u>	+ VALEANT PHARMS INTL	<u>EQ 0.25% BASE</u>	<u>N018086 001</u>	
<u>AT1</u>	+	<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
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TIMOLOL MALEATE

<u>AT2</u>	APOTEX	<u>EQ 0.5% BASE</u>	<u>A204936 001</u>	Apr 17, 2015
<u>AT3</u>	AKORN	<u>EQ 0.5% BASE</u>	<u>A212291 001</u>	Sep 11, 2020
<u>AT3</u>	AMRING PHARMS	<u>EQ 0.25% BASE</u>	<u>A212592 001</u>	Dec 13, 2021
<u>AT3</u>		<u>EQ 0.5% BASE</u>	<u>A212592 002</u>	Dec 13, 2021

TIMOPTIC IN OCULOSE

<u>AT3</u>	+! BAUSCH	<u>EQ 0.25% BASE</u>	<u>N019463 001</u>	Nov 05, 1986
<u>AT3</u>	+	<u>EQ 0.5% BASE</u>	<u>N019463 002</u>	Nov 05, 1986

TABLET;ORAL

TIMOLOL MALEATE

<u>AB</u>	ELYSIUM	<u>5MG</u>	<u>A207556 001</u>	May 02, 2019
<u>AB</u>		<u>10MG</u>	<u>A207556 002</u>	May 02, 2019
<u>AB</u>		<u>20MG</u>	<u>A207556 003</u>	May 02, 2019
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A072668 002</u>	Jun 08, 1990
<u>AB</u>		<u>10MG</u>	<u>A072668 003</u>	Jun 08, 1990
<u>AB</u>	!	<u>20MG</u>	<u>A072668 001</u>	Jun 08, 1990

TINIDAZOLE

TABLET;ORAL

TINDAMAX

<u>AB</u>	+ MISSION PHARMA	<u>250MG</u>	<u>N021618 001</u>	May 17, 2004
<u>AB</u>	+	<u>500MG</u>	<u>N021618 002</u>	May 17, 2004

TINIDAZOLE

<u>AB</u>	EDENBRIDGE PHARMS	<u>250MG</u>	<u>A203808 001</u>	Aug 04, 2015
<u>AB</u>		<u>500MG</u>	<u>A203808 002</u>	Aug 04, 2015
<u>AB</u>	HIKMA	<u>250MG</u>	<u>A201172 001</u>	Apr 30, 2012
<u>AB</u>		<u>500MG</u>	<u>A201172 002</u>	Apr 30, 2012
<u>AB</u>	NOVEL LABS INC	<u>250MG</u>	<u>A202044 001</u>	Apr 30, 2012
<u>AB</u>		<u>500MG</u>	<u>A202044 002</u>	Apr 30, 2012
<u>AB</u>	UNIQUE PHARM LABS	<u>250MG</u>	<u>A202489 001</u>	Oct 09, 2013
<u>AB</u>		<u>500MG</u>	<u>A202489 002</u>	Oct 09, 2013

TIOPRONIN

TABLET;ORAL

THIOLA

<u>AB</u>	+! MISSION PHARMA	<u>100MG</u>	<u>N019569 001</u>	Aug 11, 1988
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TIOPRONIN

<u>AB</u>	TEVA PHARMS USA INC	<u>100MG</u>	<u>A214326 001</u>	Apr 26, 2021
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TABLET, DELAYED RELEASE;ORAL

THIOLA EC

+	MISSION PHARMACAL	100MG	N211843 001	Jun 28, 2019
+	CO	300MG	N211843 002	Jun 28, 2019

TIOTROPIUM BROMIDE

POWDER; INHALATION

SPIRIVA

+	BOEHRINGER	EQ 0.018MG BASE/INH	N021395 001	Jan 30, 2004
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SPRAY, METERED; INHALATION

SPIRIVA RESPIMAT

+	BOEHRINGER	EQ 0.00125MG BASE/INH	N021936 002	Sep 15, 2015
+	INGELHEIM	EQ 0.0025MG BASE/INH	N021936 001	Sep 24, 2014

PRESCRIPTION DRUG PRODUCT LIST

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

TIRBANIBULIN

OINTMENT; TOPICAL

KLISYRI

+! ALMIRALL

1%

N213189 001 Dec 14, 2020

TIROFIBAN HYDROCHLORIDE

SOLUTION; INJECTION

AGGRASTAT

+! MEDICURE

EQ 3.75MG BASE/15ML (EQ 0.25MG BASE/ML)

N020912 002 Aug 31, 2016

SOLUTION; INTRAVENOUS

AGGRASTAT**AP +! MEDICURE****EQ 12.5MG BASE/250ML (EQ 0.05MG
BASE/ML)****N020913 003** Apr 20, 2000**TIROFIBAN HYDROCHLORIDE****AP GLAND PHARMA LTD****EQ 12.5MG BASE/250ML (EQ 0.05MG
BASE/ML)****A206888 001** Apr 08, 2021

AGGRASTAT

+ MEDICURE

EQ 5MG BASE/100ML (EQ 0.05MG BASE/ML)

N020913 002 May 17, 2002

TIVOZANIB HYDROCHLORIDE

CAPSULE; ORAL

FOTIVDA

+ AVEO PHARMS

EQ 0.89MG BASE

N212904 001 Mar 10, 2021

+!

EQ 1.34MG BASE

N212904 002 Mar 10, 2021

TIZANIDINE HYDROCHLORIDE

CAPSULE; ORAL

TIZANIDINE HYDROCHLORIDE**AB ALEMBIC PHARMS LTD****EQ 2MG BASE****A213223 001** Jan 13, 2020**AB****EQ 4MG BASE****A213223 002** Jan 13, 2020**AB****EQ 6MG BASE****A213223 003** Jan 13, 2020**AB**

ALKEM LABS LTD

EQ 2MG BASE**A212196 001** Mar 27, 2019**AB****EQ 4MG BASE****A212196 002** Mar 27, 2019**AB****EQ 6MG BASE****A212196 003** Mar 27, 2019**AB**

APOTEX INC

EQ 2MG BASE**A078868 001** Feb 03, 2012**AB****EQ 4MG BASE****A078868 002** Feb 03, 2012**AB****EQ 6MG BASE****A078868 003** Feb 03, 2012**AB**AUROBINDO PHARMA
LTD**EQ 2MG BASE****A213544 001** Mar 24, 2020**AB****EQ 4MG BASE****A213544 002** Mar 24, 2020**AB****EQ 6MG BASE****A213544 003** Mar 24, 2020**AB**

CADILA PHARMS LTD

EQ 2MG BASE**A210021 001** Mar 26, 2019**AB****EQ 4MG BASE****A210021 002** Mar 26, 2019**AB****EQ 6MG BASE****A210021 003** Mar 26, 2019**AB**

JUBILANT GENERICS

EQ 2MG BASE**A209605 001** Aug 04, 2017**AB****EQ 4MG BASE****A209605 002** Aug 04, 2017**AB****EQ 6MG BASE****A209605 003** Aug 04, 2017**AB**

NOVAST LABS

EQ 2MG BASE**A210267 001** Mar 12, 2019**AB****EQ 4MG BASE****A210267 002** Mar 12, 2019**AB****EQ 6MG BASE****A210267 003** Mar 12, 2019**AB**

RUBICON

EQ 2MG BASE**A213798 001** May 27, 2020**AB****EQ 4MG BASE****A213798 002** May 27, 2020**AB****EQ 6MG BASE****A213798 003** May 27, 2020**AB**

ZYDUS PHARMS

EQ 2MG BASE**A208622 001** Mar 03, 2017**AB****EQ 4MG BASE****A208622 002** Mar 03, 2017**AB****EQ 6MG BASE****A208622 003** Mar 03, 2017**ZANAFLEX****AB + COVIS****EQ 2MG BASE****N021447 001** Aug 29, 2002**AB +****EQ 4MG BASE****N021447 002** Aug 29, 2002**AB +!****EQ 6MG BASE****N021447 003** Aug 29, 2002

PRESCRIPTION DRUG PRODUCT LIST

TIZANIDINE HYDROCHLORIDE

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE

AB	ALKEM LABS LTD	EQ 2MG BASE	A211798 001	Jan 25, 2019
AB		EQ 4MG BASE	A211798 002	Jan 25, 2019
AB	APOTEX	EQ 2MG BASE	A076533 001	Jan 16, 2004
AB		EQ 4MG BASE	A076533 002	Jan 16, 2004
AB	CADILA	EQ 2MG BASE	A208187 001	Mar 09, 2018
AB		EQ 4MG BASE	A208187 002	Mar 09, 2018
AB	CASI PHARMS INC	EQ 2MG BASE	A076280 001	Nov 26, 2002
AB		EQ 4MG BASE	A076280 002	Jun 27, 2002
AB	DR REDDYS LABS INC	EQ 2MG BASE	A076286 001	Jul 03, 2002
AB		EQ 4MG BASE	A076286 002	Jul 03, 2002
AB	EPIC PHARMA LLC	EQ 2MG BASE	A076347 001	Oct 11, 2002
AB		EQ 4MG BASE	A076347 002	Oct 11, 2002
AB	OXFORD PHARMS	EQ 2MG BASE	A076281 001	Oct 20, 2003
AB		EQ 4MG BASE	A076281 002	Oct 20, 2003
AB	SUN PHARM INDS INC	EQ 2MG BASE	A076416 001	Sep 29, 2003
AB		EQ 4MG BASE	A076416 002	Sep 29, 2003
AB	UNICHEM LABS LTD	EQ 2MG BASE	A091283 001	Nov 28, 2012
AB		EQ 4MG BASE	A091283 002	Nov 28, 2012

ZANAFLEX

AB	+ ! COVIS	EQ 4MG BASE	N020397 001	Nov 27, 1996
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TOBRAMYCIN

OINTMENT; OPHTHALMIC

TOBREX

+! NOVARTIS

0.3%

N050555 001

POWDER; INHALATION

TOBI PODHALER

+! MYLAN SPECIALITY LP

28MG

N201688 001 Mar 22, 2013

SOLUTION; INHALATION

BETHKIS

AN	+ ! CHIESI	300MG/4ML	N201820 001	Oct 12, 2012
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KITABIS PAK

AN	PULMOFLOW INC	300MG/5ML	N205433 001	Dec 02, 2014
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TOBI

AN	+ ! MYLAN SPECIALITY LP	300MG/5ML	N050753 001	Dec 22, 1997
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TOBRAMYCIN

AN	AKORN	300MG/5ML	A201422 001	May 28, 2014
AN	ALKEM LABS LTD	300MG/5ML	A212848 001	Apr 01, 2021
AN	AMNEAL PHARMS	300MG/5ML	A205501 001	Jul 13, 2015
AN	DR REDDYS LABS SA	300MG/5ML	A207080 001	Jul 09, 2018
AN	LUPIN	300MG/5ML	A208964 001	Mar 22, 2017
AN	SUN PHARM	300MG/5ML	A207136 001	Dec 26, 2019
AN	TEVA PHARMS USA	300MG/5ML	A091589 001	Oct 10, 2013
AN		300MG/4ML	A210915 001	Jun 26, 2019

SOLUTION/DROPS; OPHTHALMIC

AKTOB

AT	AKORN	0.3%	A064096 001	Jan 31, 1996
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TOBRAMYCIN

AT	ALEMBIC PHARMS LTD	0.3%	A211847 001	Apr 19, 2019
AT	BAUSCH AND LOMB	0.3%	A064052 001	Nov 29, 1993
AT	FERA PHARMS	0.3%	A065026 001	Sep 11, 2001
AT	GLAND PHARMA LTD	0.3%	A212628 001	Mar 09, 2021
AT	SOMERSET THERAPS LLC	0.3%	A207444 001	Jun 28, 2017

TOBREX

AT	+ ! NOVARTIS	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	GLAND PHARMA LTD	EQ 10MG BASE/ML	A209621 001	Feb 11, 2021
AP		EQ 40MG BASE/ML	A209621 002	Feb 11, 2021
AP		EQ 1.2GM BASE/VIAL	A211189 001	May 18, 2021
AP	HIKMA	EQ 40MG BASE/ML	A063117 001	Apr 26, 1991

PRESCRIPTION DRUG PRODUCT LIST

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

TOBRAMYCIN SULFATE

AP	!	HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>A063112 001</u>	Apr 30, 1991
AP			<u>EQ 40MG BASE/ML</u>	<u>A063111 001</u>	Apr 30, 1991
AP		MYLAN LABS LTD	<u>EQ 40MG BASE/ML</u>	<u>A065407 001</u>	Mar 11, 2008
AP		TEVA PHARMS USA	<u>EQ 40MG BASE/ML</u>	<u>A063100 001</u>	Jan 30, 1992
AP		XELLIA PHARMS APS	<u>EQ 1.2GM BASE/VIAL</u>	<u>A205685 001</u>	Sep 16, 2014
AP	!	XGEN PHARMS	<u>EQ 1.2GM BASE/VIAL</u>	<u>A065013 001</u>	Aug 17, 2001
<u>TOBRAMYCIN SULFATE (PHARMACY BULK)</u>					
AP	!	FRESENIUS KABI USA	<u>EQ 40MG BASE/ML</u>	<u>A065120 001</u>	Nov 29, 2002
AP		GLAND PHARMA LTD	<u>EQ 40MG BASE/ML</u>	<u>A209346 001</u>	Feb 09, 2021
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

SOLUTION; ORAL

XELJANZ

+	!	PFIZER	EQ 1MG/ML BASE	N213082 001	Sep 25, 2020
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TABLET; ORAL

XELJANZ

+		PF PRISM CV	EQ 5MG BASE	N203214 001	Nov 06, 2012
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+	!		EQ 10MG BASE	N203214 002	May 30, 2018
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TABLET, EXTENDED RELEASE; ORAL

TOFACITINIB

AB		ZYDUS PHARMS	<u>EQ 11MG BASE</u>	<u>A214264 001</u>	Aug 19, 2021
AB			<u>EQ 22MG BASE</u>	<u>A214264 002</u>	Aug 19, 2021
<u>XELJANZ XR</u>					
AB	+	PFIZER	<u>EQ 11MG BASE</u>	<u>N208246 001</u>	Feb 23, 2016
AB	+	!	<u>EQ 22MG BASE</u>	<u>N208246 002</u>	Dec 12, 2019

TOLCAPONE

TABLET; ORAL

TASMAR

AB	+	!	BAUSCH	<u>100MG</u>	<u>N020697 001</u>	Jan 29, 1998
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TOLCAPONE

AB		NOVAST LABS	<u>100MG</u>	<u>A208937 001</u>	Aug 07, 2018
AB		PAR PHARM INC	<u>100MG</u>	<u>A204584 001</u>	Mar 26, 2015

TOLTERODINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

DETROL LA

AB	+	UPJOHN	<u>2MG</u>	<u>N021228 001</u>	Dec 22, 2000
AB	+	!	<u>4MG</u>	<u>N021228 002</u>	Dec 22, 2000

TOLTERODINE TARTRATE

AB		AJANTA PHARMA LTD	<u>2MG</u>	<u>A213397 001</u>	May 19, 2020
AB			<u>4MG</u>	<u>A213397 002</u>	May 19, 2020
AB		AMTA	<u>2MG</u>	<u>A213858 001</u>	Feb 02, 2021
AB			<u>4MG</u>	<u>A213858 002</u>	Feb 02, 2021
AB		HETERO LABS LTD III	<u>2MG</u>	<u>A206419 001</u>	Dec 12, 2017
AB			<u>4MG</u>	<u>A206419 002</u>	Dec 12, 2017
AB		INVENTIA HLTHCARE	<u>2MG</u>	<u>A204562 001</u>	Aug 19, 2019
AB			<u>4MG</u>	<u>A204562 002</u>	Aug 19, 2019
AB		TEVA PHARMS USA	<u>2MG</u>	<u>A079141 001</u>	Nov 22, 2016
AB			<u>4MG</u>	<u>A079141 002</u>	Nov 22, 2016
AB		TORRENT	<u>2MG</u>	<u>A203016 001</u>	Aug 11, 2015
AB			<u>4MG</u>	<u>A203016 002</u>	Aug 11, 2015

TABLET; ORAL

DETROL

AB	+	UPJOHN	<u>1MG</u>	<u>N020771 001</u>	Mar 25, 1998
AB	+	!	<u>2MG</u>	<u>N020771 002</u>	Mar 25, 1998

TOLTERODINE TARTRATE

AB		ATHEM	<u>1MG</u>	<u>A210775 001</u>	Dec 30, 2019
AB			<u>2MG</u>	<u>A210775 002</u>	Dec 30, 2019
AB		HETERO LABS LTD V	<u>1MG</u>	<u>A204397 001</u>	Aug 02, 2021
AB			<u>2MG</u>	<u>A204397 002</u>	Aug 02, 2021
AB		IVAX SUB TEVA PHARMS	<u>1MG</u>	<u>A077006 001</u>	Feb 23, 2015
AB			<u>2MG</u>	<u>A077006 002</u>	Feb 23, 2015
AB		MACLEODS PHARMS LTD	<u>1MG</u>	<u>A203409 001</u>	Aug 31, 2015
AB			<u>2MG</u>	<u>A203409 002</u>	Aug 31, 2015
AB		UNICHEM	<u>1MG</u>	<u>A205399 001</u>	Aug 05, 2020

PRESCRIPTION DRUG PRODUCT LIST

TOLTERODINE TARTRATE

TABLET; ORAL

TOLTERODINE TARTRATE

<u>AB</u>		<u>2MG</u>	<u>A205399 002</u>	Aug 05, 2020
<u>AB</u>	UNIQUE PHARM LABS	<u>1MG</u>	<u>A204721 001</u>	Jan 24, 2020
<u>AB</u>		<u>2MG</u>	<u>A204721 002</u>	Jan 24, 2020

TOLVAPTAN

TABLET; ORAL

SAMSCA

<u>AB</u>	+! OTSUKA	<u>30MG</u>	<u>N022275 002</u>	May 19, 2009
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TOLVAPTAN

<u>AB</u>	ALKEM LABS LTD	<u>30MG</u>	<u>A211891 001</u>	May 19, 2020
<u>AB</u>	APOTEX	<u>30MG</u>	<u>A207605 001</u>	May 19, 2020
<u>AB</u>	HETERO LABS LTD V	<u>30MG</u>	<u>A205646 001</u>	Jul 16, 2021

JYNARQUE

	+ OTSUKA	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	15MG	N022275 001	May 19, 2009
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TOLVAPTAN

	ALKEM LABS LTD	60MG	A211891 002	May 19, 2020
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TOPIRAMATE

CAPSULE; ORAL

TOPAMAX

<u>AB</u>	+ JANSSEN PHARMS	<u>15MG</u>	<u>N020844 001</u>	Oct 26, 1998
<u>AB</u>	+!	<u>25MG</u>	<u>N020844 002</u>	Oct 26, 1998

TOPIRAMATE

<u>AB</u>	TEVA	<u>15MG</u>	<u>A076575 001</u>	Apr 17, 2009
<u>AB</u>		<u>25MG</u>	<u>A076575 002</u>	Apr 17, 2009
<u>AB</u>	WATSON LABS	<u>15MG</u>	<u>A077868 001</u>	Apr 15, 2009
<u>AB</u>		<u>25MG</u>	<u>A077868 002</u>	Apr 15, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>15MG</u>	<u>A078877 001</u>	Oct 14, 2009
<u>AB</u>		<u>25MG</u>	<u>A078877 002</u>	Oct 14, 2009

CAPSULE, EXTENDED RELEASE; ORAL

TOPIRAMATE

<u>AB1</u>	ZYDUS PHARMS	<u>25MG</u>	<u>A207382 001</u>	Nov 24, 2017
<u>AB1</u>		<u>50MG</u>	<u>A207382 002</u>	Nov 24, 2017
<u>AB1</u>		<u>100MG</u>	<u>A207382 003</u>	Nov 24, 2017

TROKENDI XR

<u>AB1</u>	+ SUPERNUS PHARMS	<u>25MG</u>	<u>N201635 001</u>	Aug 16, 2013
<u>AB1</u>	+	<u>50MG</u>	<u>N201635 002</u>	Aug 16, 2013
<u>AB1</u>	+	<u>100MG</u>	<u>N201635 003</u>	Aug 16, 2013

OUDEXY XR

<u>AB2</u>	+ UPSHER SMITH LABS	<u>25MG</u>	<u>N205122 001</u>	Mar 11, 2014
<u>AB2</u>	+	<u>50MG</u>	<u>N205122 002</u>	Mar 11, 2014
<u>AB2</u>	+	<u>100MG</u>	<u>N205122 003</u>	Mar 11, 2014
<u>AB2</u>	+	<u>150MG</u>	<u>N205122 004</u>	Mar 11, 2014
<u>AB2</u>	+!	<u>200MG</u>	<u>N205122 005</u>	Mar 11, 2014

TOPIRAMATE

<u>AB2</u>	GLENMARK PHARMS LTD	<u>25MG</u>	<u>A210278 001</u>	Feb 01, 2021
<u>AB2</u>		<u>50MG</u>	<u>A210278 002</u>	Feb 01, 2021
<u>AB2</u>		<u>100MG</u>	<u>A210278 003</u>	Feb 01, 2021
<u>AB2</u>		<u>150MG</u>	<u>A210278 004</u>	Feb 01, 2021
<u>AB2</u>		<u>200MG</u>	<u>A210278 005</u>	Feb 01, 2021

TROKENDI XR

BC	+! SUPERNUS PHARMS	200MG	N201635 004	Aug 16, 2013
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SOLUTION; ORAL

EPRONTIA

	+! AZURITY	25MG/ML	N214679 001	Nov 05, 2021
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TABLET; ORAL

TOPAMAX

<u>AB</u>	+ JANSSEN PHARMS	<u>25MG</u>	<u>N020505 004</u>	Dec 24, 1996
<u>AB</u>	+	<u>50MG</u>	<u>N020505 005</u>	Dec 24, 1996
<u>AB</u>	+!	<u>100MG</u>	<u>N020505 001</u>	Dec 24, 1996
<u>AB</u>	+	<u>200MG</u>	<u>N020505 002</u>	Dec 24, 1996

TOPIRAMATE

<u>AB</u>	ACCORD HLTHCARE	<u>25MG</u>	<u>A076311 001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A076311 002</u>	Mar 27, 2009

PRESCRIPTION DRUG PRODUCT LIST

TOPIRAMATE

TABLET; ORAL

TOPIRAMATE

<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>	ASCENT PHARMS INC	<u>25MG</u>	<u>A215414</u>	<u>001</u>	Aug 26, 2021
<u>AB</u>		<u>50MG</u>	<u>A215414</u>	<u>002</u>	Aug 26, 2021
<u>AB</u>		<u>100MG</u>	<u>A215414</u>	<u>003</u>	Aug 26, 2021
<u>AB</u>		<u>200MG</u>	<u>A215414</u>	<u>004</u>	Aug 26, 2021
<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>	SUN PHARM	<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>	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>	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>	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>	VIWIT PHARM	<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>	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

HYCAMTIN

+ NOVARTIS

EQ 0.25MG BASE

N020981 001 Oct 11, 2007

+!

EQ 1MG BASE

N020981 002 Oct 11, 2007

INJECTABLE; INJECTION

HYCAMTINAP +! NOVARTISEQ 4MG BASE/VIALN020671 001 May 28, 1996TOPOTECAN HYDROCHLORIDEAP ACCORD HLTHCAREEQ 4MG BASE/VIALA202351 001 Jun 26, 2013AP ACTAVIS TOTOWAEQ 4MG BASE/VIALA090620 001 Dec 02, 2010AP CIPLAEQ 4MG BASE/VIALA091199 001 Dec 01, 2010AP DR REDDYS LABS LTDEQ 4MG BASE/VIALA201191 001 Mar 09, 2011AP FRESENIUS KABI USAEQ 4MG BASE/VIALA091089 001 Nov 29, 2010AP NOVAST LABSEQ 4MG BASE/VIALA206962 001 Nov 30, 2016AP SAGENT PHARMSEQ 4MG BASE/VIALA091284 001 Jan 26, 2011

SOLUTION; INTRAVENOUS

TOPOTECAN HYDROCHLORIDEAP ACCORD HLTHCAREEQ 4MG BASE/4ML (EQ 1MG BASE/ML)A204406 002 Jul 06, 2017

PRESCRIPTION DRUG PRODUCT LIST

TOPOTECAN HYDROCHLORIDE

SOLUTION; INTRAVENOUS

TOPOTECAN HYDROCHLORIDE

<u>AP</u>	+	HOSPIRA INC	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>N200582</u>	<u>001</u>	Feb 02, 2011
<u>AP</u>		MYLAN LABS LTD	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A206074</u>	<u>001</u>	Nov 24, 2017
<u>AP</u>		TEVA PHARMS USA	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>N022453</u>	<u>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>	+	KYOWA KIRIN	<u>EQ 60MG BASE</u>	<u>N020497</u>	<u>001</u>	May 29, 1997
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TOREMIFENE CITRATE

<u>AB</u>		MSN	<u>EQ 60MG BASE</u>	<u>A212818</u>	<u>001</u>	Aug 18, 2020
<u>AB</u>		RISING	<u>EQ 60MG BASE</u>	<u>A208813</u>	<u>001</u>	Dec 04, 2018

TORSEMIDE

TABLET; ORAL

TORSEMIDE

<u>AB</u>		AUROBINDO PHARMA	<u>5MG</u>	<u>A078249</u>	<u>001</u>	Oct 17, 2007
<u>AB</u>			<u>10MG</u>	<u>A078249</u>	<u>002</u>	Oct 17, 2007
<u>AB</u>			<u>20MG</u>	<u>A078249</u>	<u>003</u>	Oct 17, 2007
<u>AB</u>			<u>100MG</u>	<u>A078249</u>	<u>004</u>	Oct 17, 2007
<u>AB</u>		COREPHARMA	<u>5MG</u>	<u>A076894</u>	<u>001</u>	May 31, 2005
<u>AB</u>			<u>10MG</u>	<u>A076894</u>	<u>002</u>	May 31, 2005
<u>AB</u>			<u>20MG</u>	<u>A076894</u>	<u>003</u>	May 31, 2005
<u>AB</u>			<u>100MG</u>	<u>A076894</u>	<u>004</u>	May 31, 2005
<u>AB</u>		HETERO LABS LTD III	<u>5MG</u>	<u>A079234</u>	<u>001</u>	Jan 27, 2009
<u>AB</u>			<u>10MG</u>	<u>A079234</u>	<u>002</u>	Jan 27, 2009
<u>AB</u>			<u>20MG</u>	<u>A079234</u>	<u>003</u>	Jan 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A079234</u>	<u>004</u>	Jan 27, 2009
<u>AB</u>		HIKMA	<u>5MG</u>	<u>A076943</u>	<u>001</u>	Mar 01, 2005
<u>AB</u>			<u>10MG</u>	<u>A076943</u>	<u>002</u>	Mar 01, 2005
<u>AB</u>			<u>20MG</u>	<u>A076943</u>	<u>003</u>	Mar 01, 2005
<u>AB</u>		PLIVA PHARM IND	<u>5MG</u>	<u>A076346</u>	<u>001</u>	May 30, 2003
<u>AB</u>			<u>10MG</u>	<u>A076346</u>	<u>002</u>	May 30, 2003
<u>AB</u>	!		<u>20MG</u>	<u>A076346</u>	<u>003</u>	May 30, 2003
<u>AB</u>			<u>100MG</u>	<u>A076346</u>	<u>004</u>	Oct 19, 2004
<u>AB</u>		STRIDES PHARMA	<u>5MG</u>	<u>A076226</u>	<u>001</u>	May 27, 2003
<u>AB</u>			<u>5MG</u>	<u>A090613</u>	<u>001</u>	Mar 22, 2011
<u>AB</u>			<u>10MG</u>	<u>A076226</u>	<u>002</u>	May 27, 2003
<u>AB</u>			<u>10MG</u>	<u>A090613</u>	<u>002</u>	Mar 22, 2011
<u>AB</u>			<u>20MG</u>	<u>A076226</u>	<u>003</u>	May 27, 2003
<u>AB</u>			<u>20MG</u>	<u>A090613</u>	<u>003</u>	Mar 22, 2011
<u>AB</u>			<u>100MG</u>	<u>A076226</u>	<u>004</u>	May 27, 2003
<u>AB</u>			<u>100MG</u>	<u>A090613</u>	<u>004</u>	Mar 22, 2011
<u>AB</u>		TEVA	<u>5MG</u>	<u>A076110</u>	<u>001</u>	May 14, 2002
<u>AB</u>			<u>10MG</u>	<u>A076110</u>	<u>002</u>	May 14, 2002
<u>AB</u>			<u>20MG</u>	<u>A076110</u>	<u>003</u>	May 14, 2002
<u>AB</u>			<u>100MG</u>	<u>A076110</u>	<u>004</u>	May 14, 2002
		SOAANZ				
	+	SARFE PHARMS	20MG	N213218	001	Jun 14, 2021
	+		40MG	N213218	003	Nov 17, 2021
	+		60MG	N213218	002	Jun 14, 2021

TRABECTEDIN

POWDER; INTRAVENOUS

YONDELIS

+	JANSSEN PRODS	1MG/VIAL	N207953	001	Oct 23, 2015
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TRAMADOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CONZIP

+	CIPHER PHARMS INC	100MG	N022370	001	May 07, 2010
+		200MG	N022370	002	May 07, 2010
+		300MG	N022370	003	May 07, 2010

SOLUTION; ORAL

QDOLO

+	ATHENA	5MG/ML	N214044	001	Sep 01, 2020
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TABLET; ORAL

TRAMADOL HYDROCHLORIDE

<u>AB</u>		ACI	<u>50MG</u>	<u>A202075</u>	<u>001</u>	Nov 28, 2011
<u>AB</u>		AMNEAL PHARMS	<u>50MG</u>	<u>A076003</u>	<u>001</u>	Jun 20, 2002
<u>AB</u>		APOTEX	<u>50MG</u>	<u>A075981</u>	<u>001</u>	Jul 10, 2002

PRESCRIPTION DRUG PRODUCT LIST

TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE

AB	AUROBINDO PHARMA LTD	50MG	A203494 001	Mar 31, 2014
AB	CSPC OUYI PHARM CO	50MG	A091498 001	Mar 29, 2013
AB	IPCA LABS LTD	50MG	A201973 001	Nov 16, 2012
AB	MERRO PHARM USA	50MG	A206706 001	Jul 02, 2019
AB	MYLAN	50MG	A075986 001	Jun 21, 2002
AB	PLIVA	50MG	A075982 001	Jul 01, 2002
AB	RUBICON	50MG	A208708 001	Jun 28, 2019
AB		100MG	A208708 002	Jun 28, 2019
AB	SUN PHARM INDS INC	50MG	A075964 001	Jun 19, 2002
AB	TEVA	50MG	A075977 001	Jun 19, 2002
AB	UNICHEM	50MG	A211825 001	Aug 09, 2019
AB	ZYDUS PHARMS USA INC	50MG	A090404 001	Jan 31, 2011

ULTRAM

AB	+ ! JANSSEN PHARMS	50MG	N020281 002	Mar 03, 1995
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TABLET, EXTENDED RELEASE; ORAL

TRAMADOL HYDROCHLORIDE

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	100MG	A205257 001	Dec 22, 2015
AB1		200MG	A205257 002	Dec 22, 2015
AB1		300MG	A205257 003	Dec 22, 2015
AB1	SUN PHARM	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	! SUN PHARM	100MG	A091607 001	Dec 30, 2011
AB2		200MG	A091607 002	Dec 30, 2011
AB2		300MG	A091607 003	Dec 30, 2011

TRAMETINIB DIMETHYL SULFOXIDE

TABLET; ORAL

MEKINIST

+	NOVARTIS	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
AB		2MG	A078508 001	Jun 18, 2008
AB		4MG	A078508 002	Jun 18, 2008
AB	LUPIN	1MG	A077522 001	Jun 12, 2007
AB		2MG	A077522 002	Jun 12, 2007
AB	!	4MG	A077522 003	Jun 12, 2007
AB	TEVA PHARMS	1MG	A077489 001	Dec 12, 2006
AB		2MG	A077489 002	Dec 12, 2006
AB		4MG	A077489 003	Dec 12, 2006

TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE

	GLENMARK GENERICS	1MG; 240MG	A079135 004	Aug 30, 2010
		2MG; 180MG	A079135 001	May 26, 2010
		2MG; 240MG	A079135 002	May 26, 2010
!		4MG; 240MG	A079135 003	May 05, 2010

TRANEXAMIC ACID

INJECTABLE; INJECTION

CYKLOKAPRON

AP	+ ! PFIZER	100MG/ML	N019281 001	Dec 30, 1986
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AP	<u>TRANEXAMIC ACID</u>			
AP	ACIC PHARMS	100MG/ML	A202436 001	Feb 11, 2014
AP	AKORN	100MG/ML	A202373 001	Nov 17, 2011
AP		100MG/ML	A206594 001	Sep 28, 2017

PRESCRIPTION DRUG PRODUCT LIST

TRANEXAMIC ACID

INJECTABLE; INJECTION

TRANEXAMIC ACID

<u>AP</u>		<u>100MG/ML</u>	<u>A206634 001</u>	Jun 09, 2016
<u>AP</u>	AM REGENT	<u>100MG/ML</u>	<u>A201885 001</u>	Aug 10, 2011
<u>AP</u>	AMNEAL PHARMS CO	<u>100MG/ML</u>	<u>A208840 001</u>	Feb 28, 2017
<u>AP</u>	APOTEX	<u>100MG/ML</u>	<u>A209860 001</u>	Jan 14, 2020
<u>AP</u>	CAPLIN	<u>100MG/ML</u>	<u>A212360 001</u>	Jul 17, 2019
<u>AP</u>	EMCURE PHARMS LTD	<u>100MG/ML</u>	<u>A203521 001</u>	Aug 12, 2014
<u>AP</u>	EUGIA PHARMA	<u>100MG/ML</u>	<u>A205035 001</u>	Jan 14, 2016
<u>AP</u>	FRESENIUS KABI USA	<u>100MG/ML</u>	<u>A091596 001</u>	Mar 02, 2012
<u>AP</u>	GLAND	<u>100MG/ML</u>	<u>A207239 001</u>	Feb 13, 2017
<u>AP</u>	MICRO LABS	<u>100MG/ML</u>	<u>A206713 001</u>	Jun 27, 2017
<u>AP</u>	MYLAN INSTITUTIONAL	<u>100MG/ML</u>	<u>A091657 001</u>	Nov 03, 2011
<u>AP</u>	PROVEPHARM SAS	<u>100MG/ML</u>	<u>A212676 001</u>	Jul 17, 2019
<u>AP</u>	XGEN PHARMS	<u>100MG/ML</u>	<u>A201580 001</u>	Jun 14, 2013

SOLUTION; INTRAVENOUS

TRANEXAMIC ACID

+! EXELA PHARMA

1GM/100ML (10MG/ML)

N212020 001 Apr 15, 2019

TABLET; ORAL

LYSTEDA

<u>AB</u>	+! AMRING PHARMS	<u>650MG</u>	<u>N022430 001</u>	Nov 13, 2009
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TRANEXAMIC ACID

<u>AB</u>	ACTAVIS LABS FL INC	<u>650MG</u>	<u>A202093 001</u>	Dec 27, 2012
<u>AB</u>	ANI PHARMS	<u>650MG</u>	<u>A203256 001</u>	Jul 25, 2016

TRANLYCYPROMINE SULFATE

TABLET; ORAL

PARNATE

<u>AB</u>	+! CONCORDIA	<u>EQ 10MG BASE</u>	<u>N012342 003</u>	Aug 16, 1985
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TRANLYCYPROMINE SULFATE

<u>AB</u>	NOVITIUM PHARMA	<u>EQ 10MG BASE</u>	<u>A206856 001</u>	Apr 17, 2018
<u>AB</u>	STRIDES PHARMA	<u>EQ 10MG BASE</u>	<u>A040640 001</u>	Jun 29, 2006

TRAVOPROST

SOLUTION/DROPS; OPHTHALMIC

TRAVATAN Z

<u>AT</u>	+! NOVARTIS	<u>0.004%</u>	<u>N021994 001</u>	Sep 21, 2006
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TRAVOPROST

<u>AT</u>	APOTEX	<u>0.004%</u>	<u>A203431 001</u>	Jul 10, 2015
<u>AT</u>	MICRO LABS	<u>0.004%</u>	<u>A203767 001</u>	Mar 19, 2021
<u>AT</u>	MYLAN	<u>0.004%</u>	<u>A205050 001</u>	Jul 07, 2017
<u>AT1</u>	ALEMBIC PHARMS LTD	<u>0.004%</u>	<u>A210458 001</u>	Dec 20, 2019
<u>AT1</u>	! STRIDES PHARMA	<u>0.004%</u>	<u>A091340 001</u>	Mar 01, 2013

TRAZODONE HYDROCHLORIDE

TABLET; ORAL

TRAZODONE HYDROCHLORIDE

<u>AB</u>	ACCORD HLTHCARE	<u>50MG</u>	<u>A206923 001</u>	Sep 08, 2017
<u>AB</u>		<u>100MG</u>	<u>A206923 002</u>	Sep 08, 2017
<u>AB</u>		<u>150MG</u>	<u>A206923 003</u>	Sep 08, 2017
<u>AB</u>		<u>300MG</u>	<u>A206923 004</u>	Sep 08, 2017
<u>AB</u>	APOTEX	<u>50MG</u>	<u>A071258 001</u>	Mar 25, 1987
<u>AB</u>	! APOTEX INC	<u>100MG</u>	<u>A071196 001</u>	Mar 25, 1987
<u>AB</u>		<u>150MG</u>	<u>A071196 002</u>	Apr 26, 1999
<u>AB</u>		<u>300MG</u>	<u>A071196 003</u>	Apr 26, 1999
<u>AB</u>	AUROLIFE PHARMA LLC	<u>50MG</u>	<u>A204852 001</u>	Feb 05, 2020
<u>AB</u>		<u>100MG</u>	<u>A204852 002</u>	Feb 05, 2020
<u>AB</u>		<u>150MG</u>	<u>A204852 003</u>	Feb 05, 2020
<u>AB</u>		<u>300MG</u>	<u>A204852 004</u>	Feb 05, 2020
<u>AB</u>	OXFORD PHARMS	<u>50MG</u>	<u>A072192 001</u>	Feb 02, 1989
<u>AB</u>		<u>100MG</u>	<u>A072193 001</u>	Feb 02, 1989
<u>AB</u>	PLIVA	<u>150MG</u>	<u>A071525 001</u>	Mar 09, 1988
<u>AB</u>	SUN PHARM INDUSTRIES	<u>50MG</u>	<u>A073137 002</u>	Mar 24, 1993
<u>AB</u>		<u>100MG</u>	<u>A073137 001</u>	Mar 24, 1993
<u>AB</u>		<u>150MG</u>	<u>A073137 003</u>	Dec 22, 1995
<u>AB</u>	TEVA PHARMS USA	<u>50MG</u>	<u>A071523 001</u>	Dec 11, 1987
<u>AB</u>		<u>100MG</u>	<u>A071524 001</u>	Dec 11, 1987
<u>AB</u>	TORRENT	<u>50MG</u>	<u>A202180 001</u>	Nov 27, 2013
<u>AB</u>		<u>100MG</u>	<u>A202180 002</u>	Nov 27, 2013
<u>AB</u>		<u>150MG</u>	<u>A202180 003</u>	Nov 27, 2013
<u>AB</u>		<u>300MG</u>	<u>A202180 004</u>	Nov 27, 2013
<u>AB</u>	ZYDUS PHARMS	<u>50MG</u>	<u>A205253 001</u>	Oct 10, 2017

PRESCRIPTION DRUG PRODUCT LIST

TRAZODONE HYDROCHLORIDE

TABLET; ORAL

TRAZODONE HYDROCHLORIDE

<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

REMODULIN

<u>AP</u>	<u>+</u> !	UNITED THERAP	<u>1MG/ML</u>	<u>N021272</u>	<u>001</u>	May 21, 2002
<u>AP</u>	<u>+</u> !		<u>2.5MG/ML</u>	<u>N021272</u>	<u>002</u>	May 21, 2002
<u>AP</u>	<u>+</u> !		<u>5MG/ML</u>	<u>N021272</u>	<u>003</u>	May 21, 2002
<u>AP</u>	<u>+</u> !		<u>10MG/ML</u>	<u>N021272</u>	<u>004</u>	May 21, 2002

TREPROSTINIL

<u>AP</u>		ALEMBIC GLOBAL	<u>1MG/ML</u>	<u>A211574</u>	<u>001</u>	Feb 11, 2021
<u>AP</u>			<u>2.5MG/ML</u>	<u>A211574</u>	<u>002</u>	Feb 11, 2021
<u>AP</u>			<u>5MG/ML</u>	<u>A211574</u>	<u>003</u>	Feb 11, 2021
<u>AP</u>			<u>10MG/ML</u>	<u>A211574</u>	<u>004</u>	Feb 11, 2021
<u>AP</u>		PAR STERILE PRODUCTS	<u>1MG/ML</u>	<u>A209382</u>	<u>001</u>	Sep 24, 2019
<u>AP</u>			<u>2.5MG/ML</u>	<u>A209382</u>	<u>002</u>	Sep 24, 2019
<u>AP</u>			<u>5MG/ML</u>	<u>A209382</u>	<u>003</u>	Sep 24, 2019
<u>AP</u>			<u>10MG/ML</u>	<u>A209382</u>	<u>004</u>	Sep 24, 2019
<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
<u>AP</u>		TEVA PHARMS USA	<u>1MG/ML</u>	<u>A206648</u>	<u>001</u>	Sep 26, 2019
<u>AP</u>			<u>2.5MG/ML</u>	<u>A206648</u>	<u>002</u>	Sep 26, 2019
<u>AP</u>			<u>5MG/ML</u>	<u>A206648</u>	<u>003</u>	Sep 26, 2019
<u>AP</u>			<u>10MG/ML</u>	<u>A206648</u>	<u>004</u>	Sep 26, 2019

REMODULIN

<u>+</u> !	UNITED THERAP	20MG/ML	N021272	005	Jul 30, 2021
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SOLUTION; INHALATION

TYVASO

<u>+</u> !	UNITED THERAP	0.6MG/ML	N022387	001	Jul 30, 2009
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TREPROSTINIL DIOLAMINE

TABLET, EXTENDED RELEASE; ORAL

ORENITRAM

<u>+</u>	UNITED THERAP	EQ 0.125MG BASE	N203496	001	Dec 20, 2013
<u>+</u>		EQ 0.25MG BASE	N203496	002	Dec 20, 2013
<u>+</u> !		EQ 1MG BASE	N203496	003	Dec 20, 2013
<u>+</u>		EQ 2.5MG BASE	N203496	004	Dec 20, 2013
<u>+</u>		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>	<u>!</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
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RETIN-A

<u>AB</u>	<u>+</u> !	VALEANT BERMUDA	<u>0.025%</u>	<u>N019049</u>	<u>001</u>	Sep 16, 1988
<u>AB</u>	<u>+</u> !	VALEANT PHARMS NORTH	<u>0.1%</u>	<u>N017340</u>	<u>001</u>	

TRETINOIN

<u>AB</u>		PADAGIS US	<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
<u>AB</u>		TARO PHARMS	<u>0.1%</u>	<u>A211645</u>	<u>001</u>	Jan 22, 2019

RETIN-A

<u>AB1</u>	<u>+</u> !	VALEANT BERMUDA	<u>0.05%</u>	<u>N017522</u>	<u>001</u>	
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TRETINOIN

<u>AB1</u>		PADAGIS US	<u>0.05%</u>	<u>A075265</u>	<u>001</u>	Dec 24, 1998
<u>AB1</u>		TARO PHARMS	<u>0.05%</u>	<u>A211644</u>	<u>001</u>	Jan 25, 2019

RENOVA

<u>+</u> !	VALEANT PHARMS NORTH	0.02%	N021108	001	Aug 31, 2000
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PRESCRIPTION DRUG PRODUCT LIST

TRETINOIN

GEL; TOPICAL

ATRALIN

AB	+ !	DOW PHARM	0.05%	N022070	001	Jul 26, 2007
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RETIN-A

AB	+ !	VALEANT INTL	0.01%	N017955	001	
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AB	+ !		0.025%	N017579	002	
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TRETINOIN

AB		MYLAN	0.05%	A207955	001	Aug 13, 2015
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AB		PADAGIS US	0.01%	A075589	001	Jun 11, 2002
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AB			0.025%	A075529	001	Feb 22, 2000
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AVITA

BT		MYLAN	0.025%	N020400	001	Jan 29, 1998
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RETIN-A MICRO

	+ !	VALEANT INTL	0.04%	N020475	002	May 10, 2002
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	+ !		0.1%	N020475	001	Feb 07, 1997
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RETIN-A-MICRO

	+ !	VALEANT INTL	0.06%	N020475	004	Oct 23, 2017
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	+ !		0.08%	N020475	003	Jan 28, 2014
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LOTION; TOPICAL

ALTRENO

	+ !	DOW PHARM	0.05%	N209353	001	Aug 23, 2018
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TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

TRIAMCINOLONE ACETONIDE

AT		ALKEM LABS LTD	0.025%	A207651	001	Dec 26, 2017
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AT			0.1%	A207651	002	Dec 26, 2017
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AT			0.5%	A207651	003	Dec 26, 2017
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AT		COSETTE	0.025%	A089797	001	May 31, 1991
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AT			0.1%	A089798	001	May 31, 1991
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AT	+ !	FOUGERA PHARMS	0.025%	A085692	001	
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AT	+ !		0.1%	A085692	003	
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AT	+ !		0.5%	A085692	002	
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AT		GLENMARK PHARMS LTD	0.1%	A207117	001	Aug 05, 2016
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AT		LUPIN	0.025%	A208763	001	Feb 01, 2017
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AT			0.1%	A208763	002	Feb 01, 2017
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AT			0.5%	A208763	003	Feb 01, 2017
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AT		MACLEODS PHARMS LTD	0.025%	A209535	001	May 18, 2018
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AT			0.1%	A209535	002	May 18, 2018
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AT			0.5%	A209535	003	May 18, 2018
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AT		MICRO LABS	0.025%	A040671	001	Jun 09, 2006
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AT			0.1%	A040671	002	Jun 09, 2006
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AT		PADAGIS US	0.025%	A086413	002	
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AT			0.1%	A086413	003	
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AT			0.5%	A086413	001	
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AT		STRIDES PHARMA	0.025%	A210346	001	Feb 11, 2019
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AT			0.1%	A210346	002	Feb 11, 2019
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AT			0.5%	A210346	003	Feb 11, 2019
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AT		TARO PHARM INDS LTD	0.1%	A040039	001	Nov 26, 1997
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AT		TELLIGENT	0.1%	A208848	001	Sep 18, 2017
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TRIDERM

AT		CROWN LABS	0.025%	A088042	002	Mar 25, 2015
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AT			0.1%	A088042	001	Mar 19, 1984
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AT			0.5%	A088042	003	Mar 25, 2015
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FOR SUSPENSION, EXTENDED RELEASE; INTRA-ARTICULAR

ZILRETTA

	+ !	FLEXION THERAPS INC	32MG/VIAL	N208845	001	Oct 06, 2017
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INJECTABLE; INJECTION

KENALOG-40

AB	+ !	APOTHECON	40MG/ML	N014901	001	
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TRIAMCINOLONE ACETONIDE

AB		TEVA PHARMS USA	40MG/ML	A209852	001	Oct 05, 2018
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BX		AMNEAL	40MG/ML	A207550	001	Dec 11, 2017
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KENALOG-10

	+	APOTHECON	10MG/ML	N012041	001	
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KENALOG-80

	+ !	APOTHECON	80MG/ML	N014901	002	Apr 12, 2019
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INJECTABLE; INTRAVITREAL

TRIESENCE

	+ !	NOVARTIS	40MG/ML (40MG/ML)	N022048	001	Nov 29, 2007
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PRESCRIPTION DRUG PRODUCT LIST

TRIAMCINOLONE ACETONIDE

LOTION; TOPICAL

TRIAMCINOLONE ACETONIDE

<u>AT</u>	AKORN	<u>0.025%</u>	<u>A202374</u>	<u>001</u>	May 08, 2013
<u>AT</u>		<u>0.1%</u>	<u>A202374</u>	<u>002</u>	May 08, 2013
<u>AT</u>	COSETTE	<u>0.1%</u>	<u>A089129</u>	<u>001</u>	Aug 14, 1986
<u>AT</u>	FOUGERA PHARMS	<u>0.025%</u>	<u>A040467</u>	<u>001</u>	Apr 21, 2003
<u>AT</u>		<u>0.1%</u>	<u>A040467</u>	<u>002</u>	Apr 21, 2003
<u>AT</u>	MICRO LABS	<u>0.1%</u>	<u>A040672</u>	<u>002</u>	Dec 13, 2006
<u>AT</u>	QUAGEN	<u>0.025%</u>	<u>A213559</u>	<u>001</u>	Jul 01, 2020
<u>AT</u>		<u>0.1%</u>	<u>A213559</u>	<u>002</u>	Jul 01, 2020
<u>AT</u>	TELIGENT	<u>0.025%</u>	<u>A204608</u>	<u>001</u>	Jul 07, 2016
<u>AT</u>		<u>0.1%</u>	<u>A204606</u>	<u>001</u>	Jul 07, 2016
<u>AT</u>	! WOCKHARDT BIO AG	<u>0.025%</u>	<u>A088450</u>	<u>001</u>	Apr 01, 1985
<u>AT</u>	!	<u>0.1%</u>	<u>A088451</u>	<u>001</u>	Apr 03, 1985

OINTMENT; TOPICAL

TRIAMCINOLONE ACETONIDE

<u>AT</u>	CINTEX SVCS	<u>0.05%</u>	<u>A213619</u>	<u>001</u>	Nov 10, 2020
<u>AT</u>	COSETTE	<u>0.025%</u>	<u>A089795</u>	<u>001</u>	Dec 23, 1988
<u>AT</u>		<u>0.1%</u>	<u>A089796</u>	<u>001</u>	Dec 23, 1988
<u>AT</u>		<u>0.5%</u>	<u>A208925</u>	<u>001</u>	Oct 06, 2017
<u>AT</u>	ENCUBE	<u>0.05%</u>	<u>A212384</u>	<u>001</u>	Nov 29, 2019
<u>AT</u>	FOUGERA PHARMS	<u>0.025%</u>	<u>A085691</u>	<u>001</u>	
<u>AT</u>		<u>0.1%</u>	<u>A085691</u>	<u>003</u>	
<u>AT</u>		<u>0.5%</u>	<u>A085691</u>	<u>002</u>	
<u>AT</u>	GLENMARK PHARMS	<u>0.1%</u>	<u>A208320</u>	<u>001</u>	Aug 22, 2017
<u>AT</u>	GLENMARK PHARMS LTD	<u>0.5%</u>	<u>A206379</u>	<u>001</u>	Jul 22, 2016
<u>AT</u>	MACLEODS PHARMS LTD	<u>0.025%</u>	<u>A209828</u>	<u>001</u>	Nov 23, 2018
<u>AT</u>		<u>0.1%</u>	<u>A209828</u>	<u>002</u>	Nov 23, 2018
<u>AT</u>		<u>0.5%</u>	<u>A209828</u>	<u>003</u>	Nov 23, 2018
<u>AT</u>	NOVEL LABS INC	<u>0.1%</u>	<u>A207365</u>	<u>001</u>	Oct 12, 2018
<u>AT</u>	PADAGIS ISRAEL	<u>0.05%</u>	<u>A212460</u>	<u>001</u>	Feb 05, 2021
<u>AT</u>	+! PADAGIS US	<u>0.025%</u>	<u>A087385</u>	<u>002</u>	
<u>AT</u>	+!	<u>0.1%</u>	<u>A087385</u>	<u>003</u>	
<u>AT</u>	+!	<u>0.5%</u>	<u>A087385</u>	<u>001</u>	
<u>AT</u>	STRIDES PHARMA	<u>0.05%</u>	<u>A212356</u>	<u>001</u>	Jun 01, 2020
<u>AT</u>	TARO PHARM INDS LTD	<u>0.1%</u>	<u>A040037</u>	<u>001</u>	Sep 30, 1994
<u>AT</u>	TELIGENT	<u>0.1%</u>	<u>A205373</u>	<u>001</u>	May 13, 2016
<u>AT</u>		<u>0.5%</u>	<u>A208590</u>	<u>001</u>	Mar 03, 2017

TRIANEX

<u>AT</u>	! CMP PHARMA INC	<u>0.05%</u>	<u>A089595</u>	<u>001</u>	Mar 23, 1995
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PASTE; DENTAL

TRIAMCINOLONE ACETONIDE

<u>AT</u>	AKORN	<u>0.1%</u>	<u>A206312</u>	<u>001</u>	Aug 11, 2016
<u>AT</u>	COSETTE	<u>0.1%</u>	<u>A205592</u>	<u>001</u>	Jan 12, 2017
<u>AT</u>	LYNE	<u>0.1%</u>	<u>A040771</u>	<u>001</u>	Jul 01, 2010
<u>AT</u>	! TARO	<u>0.1%</u>	<u>A070730</u>	<u>001</u>	Oct 01, 1986

SPRAY; TOPICAL

KENALOG

<u>AT</u>	+! SUN PHARM INDS INC	<u>0.147MG/GM</u>	<u>N012104</u>	<u>001</u>	
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TRIAMCINOLONE ACETONIDE

<u>AT</u>	AKORN	<u>0.147MG/GM</u>	<u>A207094</u>	<u>001</u>	Dec 07, 2016
<u>AT</u>	PADAGIS ISRAEL	<u>0.147MG/GM</u>	<u>A205782</u>	<u>001</u>	Apr 13, 2015
<u>AT</u>	RISING	<u>0.147MG/GM</u>	<u>A206786</u>	<u>001</u>	Sep 08, 2017

SPRAY, METERED; NASAL

NASACORT AQ

<u>AB</u>	+ SANOFI AVENTIS US	<u>0.055MG/SPRAY</u>	<u>N020468</u>	<u>001</u>	May 20, 1996
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SUSPENSION; INJECTION

XIPERE

+!	BAUSCH AND LOMB INC	40MG/ML	N211950	001	Oct 22, 2021
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TRIAMCINOLONE HEXACETONIDE

INJECTABLE; INJECTION

ARISTOSPAN

+!	SANDOZ INC	5MG/ML	N016466	001	
+!		20MG/ML	N016466	002	

PRESCRIPTION DRUG PRODUCT LIST

TRIAMTERENE

CAPSULE; ORAL

DYRENIUM

<u>AB</u>	+	CONCORDIA	<u>50MG</u>	<u>N013174</u>	<u>001</u>	
<u>AB</u>	+	!	<u>100MG</u>	<u>N013174</u>	<u>002</u>	

TRIAMTERENE

<u>AB</u>		AGNITIO	<u>50MG</u>	<u>A211581</u>	<u>001</u>	Aug 19, 2019
<u>AB</u>			<u>100MG</u>	<u>A211581</u>	<u>002</u>	Aug 19, 2019

TRIAZOLAM

TABLET; ORAL

HALCION

<u>AB</u>	+	PFIZER	<u>0.125MG</u>	<u>N017892</u>	<u>003</u>	Apr 26, 1985
<u>AB</u>	+	!	<u>0.25MG</u>	<u>N017892</u>	<u>001</u>	Nov 15, 1982

TRIAZOLAM

<u>AB</u>		HIKMA	<u>0.125MG</u>	<u>A074224</u>	<u>001</u>	Jun 01, 1994
<u>AB</u>			<u>0.25MG</u>	<u>A074224</u>	<u>002</u>	Jun 01, 1994
<u>AB</u>		NOVAST LABS	<u>0.125MG</u>	<u>A214219</u>	<u>001</u>	Oct 20, 2020
<u>AB</u>			<u>0.25MG</u>	<u>A214219</u>	<u>002</u>	Oct 20, 2020

TRICLABENDAZOLE

TABLET; ORAL

EGATEN

	+	NOVARTIS	250MG	N208711	001	Feb 13, 2019
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TRIENTINE HYDROCHLORIDE

CAPSULE; ORAL

SYPRINE

<u>AB</u>	+	BAUSCH	<u>250MG</u>	<u>N019194</u>	<u>001</u>	Nov 08, 1985
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TRIENTINE HYDROCHLORIDE

<u>AB</u>		ACCORD HLTHCARE	<u>250MG</u>	<u>A212929</u>	<u>001</u>	Aug 30, 2021
<u>AB</u>		DR REDDYS LABS LTD	<u>250MG</u>	<u>A211076</u>	<u>001</u>	Jul 03, 2019
<u>AB</u>		LUPIN LTD	<u>250MG</u>	<u>A211637</u>	<u>001</u>	May 21, 2020
<u>AB</u>		NAVINTA LLC	<u>250MG</u>	<u>A211251</u>	<u>001</u>	Jan 16, 2019
<u>AB</u>		PAR PHARM INC	<u>250MG</u>	<u>A210096</u>	<u>001</u>	Sep 25, 2019
<u>AB</u>		PHARMA LIFE	<u>250MG</u>	<u>A209945</u>	<u>001</u>	Aug 13, 2021
<u>AB</u>		RISING	<u>250MG</u>	<u>A212238</u>	<u>001</u>	Feb 20, 2020
<u>AB</u>		WATSON LABS TEVA	<u>250MG</u>	<u>A207567</u>	<u>001</u>	Feb 07, 2018
<u>AB</u>		ZYDUS PHARMS	<u>250MG</u>	<u>A211554</u>	<u>001</u>	Apr 26, 2019
EX		MSN	250MG	A211134	001	May 22, 2019

TRIFAROTENE

CREAM; TOPICAL

AKLIEF

	+	GALDERMA LABS LP	0.005%	N211527	001	Oct 04, 2019
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TRIFLUOPERAZINE HYDROCHLORIDE

TABLET; ORAL

TRIFLUOPERAZINE HYDROCHLORIDE

<u>AB</u>		MYLAN	<u>EQ 1MG BASE</u>	<u>A040209</u>	<u>001</u>	Jul 07, 1997
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A040209</u>	<u>002</u>	Jul 07, 1997
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A040209</u>	<u>003</u>	Jul 07, 1997
<u>AB</u>	!		<u>EQ 10MG BASE</u>	<u>A040209</u>	<u>004</u>	Jul 07, 1997
<u>AB</u>		SANDOZ	<u>EQ 1MG BASE</u>	<u>A085785</u>	<u>001</u>	
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A085786</u>	<u>001</u>	
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A085789</u>	<u>001</u>	
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A085788</u>	<u>001</u>	

TRIFLURIDINE

SOLUTION/DROPS; OPHTHALMIC

TRIFLURIDINE

<u>AT</u>		AKORN	<u>1%</u>	<u>A205438</u>	<u>001</u>	Jul 28, 2017
<u>AT</u>		SANDOZ INC	<u>1%</u>	<u>A074311</u>	<u>001</u>	Oct 06, 1995
<u>AT</u>	+	MONARCH PHARMS	<u>1%</u>	<u>N018299</u>	<u>001</u>	

TRIHPTANOIN

LIQUID; ORAL

DOJOLVI

	+	ULTRAGENYX PHARM INC	100% w/w	N213687	001	Jun 30, 2020
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PRESCRIPTION DRUG PRODUCT LIST

TRIHEXYPHENIDYL HYDROCHLORIDE

ELIXIR; ORAL

TRIHEXYPHENIDYL HYDROCHLORIDE

AA	MIKART	2MG/5ML	A040251 001	Sep 27, 1999
AA	! PHARM ASSOC	2MG/5ML	A040177 001	Apr 17, 1997

TABLET; ORAL

TRIHEXYPHENIDYL HYDROCHLORIDE

AA	NATCO PHARMA LTD	2MG	A091630 001	Nov 17, 2010
AA		5MG	A091630 002	Nov 17, 2010
AA	NOVITIUM PHARMA	2MG	A040254 001	Dec 24, 1998
AA		5MG	A040254 002	Dec 24, 1998
AA	! WATSON LABS	2MG	A084363 001	
AA	!	5MG	A084364 001	

TRILACICLIB DIHYDROCHLORIDE

POWDER; INTRAVENOUS

COSELA

+	! G1 THERAP	EQ 300MG BASE/VIAL	N214200 001	Feb 12, 2021
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TRIMETHOBENZAMIDE HYDROCHLORIDE

CAPSULE; ORAL

TRIMETHOBENZAMIDE HYDROCHLORIDE

AB	LUPIN	300MG	A076546 001	Aug 20, 2003
AB	! SUN PHARM INDUSTRIES	300MG	A076570 001	Aug 28, 2003

INJECTABLE; INJECTION

TIGAN

+	! PAR STERILE PRODUCTS	100MG/ML	N017530 001	
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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

TRIMIPRAMINE MALEATE

CAPSULE; ORAL

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
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INJECTABLE; INTRAMUSCULAR

TRELSTAR

+	! VERITY	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

TROMETHAMINE

SOLUTION; INJECTION

THAM

+	! HOSPIRA	18GM/500ML (3.6GM/100ML)	N013025 002	
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TROPICAMIDE

SOLUTION/DROPS; OPHTHALMIC

MYDRIACYL

AT	! ALCON LABS INC	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
AT	SOMERSET THERAPS LLC	1%	A207524 001	Dec 12, 2019

PRESCRIPTION DRUG PRODUCT LIST

TROSPIUM CHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

TROSPIUM CHLORIDE

AB	!	ACTAVIS LABS FL INC	60MG	A091289 001	Oct 12, 2012
AB		AMTA	60MG	A214760 001	Apr 30, 2021
AB		GRANULES	60MG	A213185 001	Apr 23, 2020
AB		PADAGIS US	60MG	A201291 001	May 24, 2013

TABLET;ORAL

TROSPIUM 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
AB		INVAGEN PHARMS	20MG	A091688 001	Aug 23, 2016
AB		PADAGIS US	20MG	A091573 001	Nov 17, 2010

TRYPAN BLUE

SOLUTION;OPHTHALMIC

MEMBRANEBLUE

+	!	DORC	0.15%	N022278 001	Feb 20, 2009
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VISIONBLUE

+	!	DORC	0.06%	N021670 001	Dec 16, 2004
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TUCATINIB

TABLET;ORAL

TUKYSA

+		SEAGEN	50MG	N213411 001	Apr 17, 2020
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+	!		150MG	N213411 002	Apr 17, 2020
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UBROGEPANT

TABLET;ORAL

UBRELVY

+		ALLERGAN	50MG	N211765 001	Dec 23, 2019
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+	!		100MG	N211765 002	Dec 23, 2019
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ULIPRISTAL ACETATE

TABLET;ORAL

ELLA

AB	+	!	LAB HRA PHARMA	30MG	N022474 001	Aug 13, 2010
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LOGILIA

AB			TEVA PHARMS USA	30MG	A207952 001	Feb 13, 2017
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UMBRALISIB TOSYLATE

TABLET;ORAL

UKONIQ

+	!	TG THERAPS	EQ 200MG BASE	N213176 001	Feb 05, 2021
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UMECLIDINIUM BROMIDE

POWDER;INHALATION

INCRUSE ELLIPTA

+	!	GLAXO GRP ENGLAND	EQ 62.5MCG BASE/INH	N205382 001	Apr 30, 2014
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UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE

POWDER;INHALATION

ANORO ELLIPTA

+	!	GLAXOSMITHKLINE	EQ 0.0625MG BASE/INH;EQ 0.025MG	N203975 001	Dec 18, 2013
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BASE/INH

UPADACITINIB

TABLET, EXTENDED RELEASE;ORAL

RINVOQ

+	!	ABBVIE INC	15MG	N211675 001	Aug 16, 2019
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UREA, C-14

CAPSULE;ORAL

PYTEST

+	!	AVENT	1uCi	N020617 001	May 09, 1997
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PYTEST KIT

+	!	AVENT	1uCi	N020617 002	May 09, 1997
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URIDINE TRIACETATE

GRANULE;ORAL

VISTOGARD

+	!	WELLSTAT THERAP	10GM/PACKET	N208159 001	Dec 11, 2015
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XURIDEN

+	!	WELLSTAT THERAP	2GM/PACKET	N208169 001	Sep 04, 2015
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PRESCRIPTION DRUG PRODUCT LIST

URSODIOL

CAPSULE; ORAL

ACTIGALL

AB	+!	ALLERGAN	300MG	<u>N019594</u>	<u>002</u>	Dec 31, 1987
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URSODIOL

AB		ABHAI LLC	300MG	<u>A210707</u>	<u>001</u>	May 17, 2018
AB		AMNEAL PHARMS CO	300MG	<u>A211301</u>	<u>001</u>	Oct 16, 2018
AB		CHARTWELL RX	300MG	<u>A213555</u>	<u>001</u>	Aug 17, 2020
AB		EPIC PHARMA	300MG	<u>A075517</u>	<u>001</u>	Mar 14, 2000
AB		EYWA PHARMA	300MG	<u>A212452</u>	<u>001</u>	Oct 30, 2019
AB		LANNETT CO INC	300MG	<u>A079082</u>	<u>001</u>	Dec 15, 2008
AB		MYLAN	300MG	<u>A090530</u>	<u>001</u>	Feb 17, 2010
AB		RISING	300MG	<u>A213200</u>	<u>001</u>	Feb 12, 2020
AB		STRIDES PHARMA	300MG	<u>A210344</u>	<u>001</u>	Jan 22, 2021
AB		TEVA PHARMS	300MG	<u>A075592</u>	<u>001</u>	May 25, 2000
AB		VGYAAN	300MG	<u>A214329</u>	<u>001</u>	Jul 28, 2021
AB		ZYDUS	300MG	<u>A214295</u>	<u>001</u>	Oct 16, 2020
		LGM PHARMA	200MG	A205789	001	May 08, 2020
		!	400MG	A205789	002	May 08, 2020

TABLET; ORAL

URSO 250

AB	+	ALLERGAN	250MG	<u>N020675</u>	<u>001</u>	Dec 10, 1997
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URSO FORTE

AB	+!	ALLERGAN	500MG	<u>N020675</u>	<u>002</u>	Jul 21, 2004
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URSODIOL

AB		GLENMARK GENERICS	250MG	<u>A090801</u>	<u>001</u>	Jul 12, 2011
AB			500MG	<u>A090801</u>	<u>002</u>	Jul 12, 2011
AB		PAR PHARM	250MG	<u>A202540</u>	<u>001</u>	Feb 14, 2013
AB			500MG	<u>A202540</u>	<u>002</u>	Feb 14, 2013
AB		STRIDES PHARMA	250MG	<u>A213504</u>	<u>001</u>	Aug 20, 2020
AB			500MG	<u>A213504</u>	<u>002</u>	Aug 20, 2020
AB		ZYDUS	250MG	<u>A211145</u>	<u>001</u>	Oct 30, 2018
AB			500MG	<u>A211145</u>	<u>002</u>	Oct 30, 2018

VALACYCLOVIR HYDROCHLORIDE

TABLET; ORAL

VALACYCLOVIR HYDROCHLORIDE

AB		APOTEX	EQ 500MG BASE	<u>A090500</u>	<u>001</u>	Apr 04, 2014
AB			EQ 1GM BASE	<u>A090500</u>	<u>002</u>	Apr 04, 2014
AB		ATLANTIDE	EQ 500MG BASE	<u>A090216</u>	<u>001</u>	May 24, 2010
AB			EQ 1GM BASE	<u>A090216</u>	<u>002</u>	May 24, 2010
AB		AUROBINDO PHARMA	EQ 500MG BASE	<u>A090682</u>	<u>001</u>	May 24, 2010
AB			EQ 1GM BASE	<u>A090682</u>	<u>002</u>	May 24, 2010
AB		CADILA	EQ 500MG BASE	<u>A079137</u>	<u>001</u>	Dec 29, 2017
AB			EQ 1GM BASE	<u>A079137</u>	<u>002</u>	Dec 29, 2017
AB		CIPLA	EQ 500MG BASE	<u>A077135</u>	<u>001</u>	May 24, 2010
AB			EQ 1GM BASE	<u>A077135</u>	<u>002</u>	May 24, 2010
AB		HETERO LABS LTD V	EQ 500MG BASE	<u>A203047</u>	<u>001</u>	Apr 08, 2015
AB			EQ 1GM BASE	<u>A203047</u>	<u>002</u>	Apr 08, 2015
AB		JUBILANT GENERICS	EQ 500MG BASE	<u>A201506</u>	<u>001</u>	Apr 03, 2012
AB			EQ 1GM BASE	<u>A201506</u>	<u>002</u>	Apr 03, 2012
AB		MYLAN PHARMS INC	EQ 500MG BASE	<u>A078518</u>	<u>001</u>	May 24, 2010
AB			EQ 1GM BASE	<u>A078518</u>	<u>002</u>	May 24, 2010
AB		SANDOZ	EQ 500MG BASE	<u>A077478</u>	<u>001</u>	May 24, 2010
AB			EQ 1GM BASE	<u>A077478</u>	<u>002</u>	May 24, 2010
AB		SUN PHARM INDS LTD	EQ 500MG BASE	<u>A076588</u>	<u>001</u>	Jan 31, 2007
AB			EQ 1GM BASE	<u>A076588</u>	<u>002</u>	Jan 31, 2007
AB		TIME-CAP LABS INC	EQ 500MG BASE	<u>A079012</u>	<u>001</u>	May 24, 2010
AB			EQ 1GM BASE	<u>A079012</u>	<u>002</u>	May 24, 2010
AB		YILING	EQ 500MG BASE	<u>A209553</u>	<u>001</u>	Mar 18, 2020
AB			EQ 1GM BASE	<u>A209553</u>	<u>002</u>	Mar 18, 2020

VALTREX

AB	+	GLAXOSMITHKLINE	EQ 500MG BASE	<u>N020487</u>	<u>001</u>	Jun 23, 1995
AB	+!		EQ 1GM BASE	<u>N020487</u>	<u>002</u>	Jun 23, 1995

VALBENAZINE TOSYLATE

CAPSULE; ORAL

INGREZZA

	+	NEUROCRINE	EQ 40MG BASE	N209241	001	Apr 11, 2017
	+		EQ 60MG BASE	N209241	003	Apr 23, 2021
	+!		EQ 80MG BASE	N209241	002	Oct 04, 2017

PRESCRIPTION DRUG PRODUCT LIST

VALGANCICLOVIR HYDROCHLORIDE

FOR SOLUTION; ORAL

VALCYTE

AB	+ !	HOFFMANN LA ROCHE	50MG/ML	<u>N022257</u>	<u>001</u>	Aug 28, 2009
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VALGANCICLOVIR HYDROCHLORIDE

AB		ACTAVIS LABS FL INC	50MG/ML	<u>A205220</u>	<u>001</u>	Jul 18, 2016
AB		AJANTA PHARMA LTD	50MG/ML	<u>A212890</u>	<u>001</u>	Jan 13, 2020
AB		GRANULES	50MG/ML	<u>A213306</u>	<u>001</u>	Jan 31, 2020

TABLET; ORAL

VALCYTE

AB	+ !	HOFFMANN LA ROCHE	EQ 450MG BASE	<u>N021304</u>	<u>001</u>	Mar 29, 2001
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VALGANCICLOVIR HYDROCHLORIDE

AB		AJANTA PHARMA LTD	EQ 450MG BASE	<u>A212234</u>	<u>001</u>	Dec 26, 2019
AB		AUROBINDO PHARMA LTD	EQ 450MG BASE	<u>A204750</u>	<u>001</u>	Mar 31, 2016
AB		CIPLA	EQ 450MG BASE	<u>A209672</u>	<u>001</u>	Nov 09, 2018
AB		DR REDDYS	EQ 450MG BASE	<u>A206876</u>	<u>001</u>	Dec 12, 2017
AB		DR REDDYS LABS LTD	EQ 450MG BASE	<u>A203511</u>	<u>001</u>	Nov 04, 2014
AB		HETERO LABS LTD V	EQ 450MG BASE	<u>A205166</u>	<u>001</u>	Mar 18, 2016
AB		MYLAN	EQ 450MG BASE	<u>A205151</u>	<u>001</u>	Mar 03, 2021
AB		STRIDES PHARMA	EQ 450MG BASE	<u>A200790</u>	<u>001</u>	Nov 04, 2014

VALPROATE SODIUM

INJECTABLE; INJECTION

VALPROATE SODIUM

AP		ATHENEX INC	EQ 100MG BASE/ML	<u>A076295</u>	<u>001</u>	Nov 14, 2002
AP		FRESENIUS KABI USA	EQ 100MG BASE/ML	<u>A076539</u>	<u>001</u>	Jun 26, 2003
AP	!	HIKMA FARMACEUTICA	EQ 100MG BASE/ML	<u>A078523</u>	<u>001</u>	Feb 17, 2010
AP		MYLAN	EQ 100MG BASE/ML	<u>A208120</u>	<u>001</u>	Dec 22, 2021

VALPROIC ACID

CAPSULE; ORAL

VALPROIC ACID

AB	!	BIONPHARMA INC	250MG	<u>A073484</u>	<u>001</u>	Jun 29, 1993
AB		CATALENT	250MG	<u>A073229</u>	<u>001</u>	Oct 29, 1991
AB		SUN PHARM INDS LTD	250MG	<u>A091037</u>	<u>001</u>	Feb 22, 2013
BX		EYWA	250MG	<u>A207611</u>	<u>001</u>	Aug 05, 2019

SYRUP; ORAL

VALPROIC ACID

AA		BIOPHARM	250MG/5ML	<u>A090517</u>	<u>001</u>	May 28, 2010
AA		CHARTWELL RX	250MG/5ML	<u>A075782</u>	<u>001</u>	Dec 22, 2000
AA		HIGH TECH PHARMA	250MG/5ML	<u>A074060</u>	<u>001</u>	Jan 13, 1995
AA		LANNETT CO INC	250MG/5ML	<u>A077960</u>	<u>001</u>	Oct 13, 2006
AA	!	PHARM ASSOC	250MG/5ML	<u>A075379</u>	<u>001</u>	Dec 15, 2000
AA		WOCKHARDT BIO AG	250MG/5ML	<u>A070868</u>	<u>001</u>	Jul 01, 1986

VALRUBICIN

SOLUTION; INTRAVESICAL

VALRUBICIN

AO		CUSTOPHARM INC	40MG/ML	<u>A206430</u>	<u>001</u>	Apr 19, 2019
AO	+ !	ENDO PHARM	40MG/ML	<u>N020892</u>	<u>001</u>	Sep 25, 1998

VALSARTAN

SOLUTION; ORAL

VALSARTAN

		NOVITIUM PHARMA	20MG/5ML	<u>A214102</u>	<u>001</u>	Nov 02, 2021
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TABLET; ORAL

DIOVAN

AB	+	NOVARTIS	40MG	<u>N021283</u>	<u>004</u>	Aug 14, 2002
AB	+		80MG	<u>N021283</u>	<u>001</u>	Jul 18, 2001
AB	+		160MG	<u>N021283</u>	<u>002</u>	Jul 18, 2001
AB	+ !		320MG	<u>N021283</u>	<u>003</u>	Jul 18, 2001

VALSARTAN

AB		ALEMBIC PHARMS LTD	40MG	<u>A091367</u>	<u>001</u>	Jan 05, 2015
AB			80MG	<u>A091367</u>	<u>002</u>	Jan 05, 2015
AB			160MG	<u>A091367</u>	<u>003</u>	Jan 05, 2015
AB			320MG	<u>A091367</u>	<u>004</u>	Jan 05, 2015
AB		ALKEM LABS LTD	40MG	<u>A205536</u>	<u>001</u>	Mar 12, 2019
AB			80MG	<u>A205536</u>	<u>002</u>	Mar 12, 2019
AB			160MG	<u>A205536</u>	<u>003</u>	Mar 12, 2019
AB			320MG	<u>A205536</u>	<u>004</u>	Mar 12, 2019
AB		AMNEAL PHARMS	40MG	<u>A204011</u>	<u>001</u>	Jan 11, 2016
AB			80MG	<u>A204011</u>	<u>002</u>	Jan 11, 2016
AB			160MG	<u>A204011</u>	<u>003</u>	Jan 11, 2016

PRESCRIPTION DRUG PRODUCT LIST

VANCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

VANCOMYCIN HYDROCHLORIDE

<u>AB</u>	WATSON LABS	<u>EQ 125MG BASE</u>	<u>A065510</u>	<u>001</u>	Apr 09, 2012
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065510</u>	<u>002</u>	Apr 09, 2012

FOR SOLUTION; ORAL

FIRVANQ KIT

+	!	AZURITY	EQ 25MG BASE/ML	N208910	001	Jan 26, 2018
+	!		EQ 50MG BASE/ML	N208910	002	Jan 26, 2018

VANCOMYCIN HYDROCHLORIDE

!		ANI PHARMS	EQ 250MG BASE/5ML	A061667	002	Jul 13, 1983
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INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

<u>AP</u>		EUGIA PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A205780</u>	<u>001</u>	Mar 31, 2016	
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A205780</u>	<u>002</u>	Mar 31, 2016	
<u>AP</u>			<u>EQ 5GM BASE/VIAL</u>	<u>A205779</u>	<u>001</u>	Mar 29, 2016	
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A205779</u>	<u>002</u>	Mar 29, 2016	
<u>AP</u>	!	FRESENIUS KABI USA	<u>EQ 500MG BASE/VIAL</u>	<u>A062663</u>	<u>001</u>	Mar 17, 1987	
<u>AP</u>			<u>EQ 750MG BASE/VIAL</u>	<u>A062663</u>	<u>005</u>	Aug 17, 2016	
<u>AP</u>	!		<u>EQ 1GM BASE/VIAL</u>	<u>A062663</u>	<u>002</u>	Jul 31, 1987	
<u>AP</u>	!		<u>EQ 5GM BASE/VIAL</u>	<u>A062663</u>	<u>003</u>	Jun 03, 1988	
<u>AP</u>	!		<u>EQ 10GM BASE/VIAL</u>	<u>A062663</u>	<u>004</u>	Nov 28, 1997	
<u>AP</u>		GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A205694</u>	<u>001</u>	Jan 21, 2016	
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A205694</u>	<u>002</u>	Jan 21, 2016	
<u>AP</u>		HAINAN POLY	<u>EQ 5GM BASE/VIAL</u>	<u>A215821</u>	<u>001</u>	Nov 18, 2021	
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A215821</u>	<u>002</u>	Nov 18, 2021	
<u>AP</u>		HAINAN POLY PHARM	<u>EQ 500MG BASE/VIAL</u>	<u>A212332</u>	<u>001</u>	Jun 12, 2019	
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A212332</u>	<u>002</u>	Jun 12, 2019	
<u>AP</u>		HIKMA	<u>EQ 5GM BASE/VIAL</u>	<u>A204360</u>	<u>001</u>	Oct 15, 2018	
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A204360</u>	<u>002</u>	Oct 15, 2018	
<u>AP</u>		HIKMA PHARMS	<u>EQ 750MG BASE/VIAL</u>	<u>A206616</u>	<u>001</u>	Oct 03, 2018	
<u>AP</u>	!	HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>A062911</u>	<u>001</u>	Aug 04, 1988	
<u>AP</u>	!		<u>EQ 500MG BASE/VIAL</u>	<u>A062931</u>	<u>001</u>	Oct 29, 1992	
<u>AP</u>	!		<u>EQ 750MG BASE/VIAL</u>	<u>A062912</u>	<u>002</u>	Jan 07, 2009	
<u>AP</u>	!		<u>EQ 750MG BASE/VIAL</u>	<u>A062933</u>	<u>002</u>	May 27, 2009	
<u>AP</u>	!		<u>EQ 1GM BASE/VIAL</u>	<u>A062912</u>	<u>001</u>	Aug 04, 1988	
<u>AP</u>	!		<u>EQ 1GM BASE/VIAL</u>	<u>A062933</u>	<u>001</u>	Oct 29, 1992	
<u>AP</u>	!		<u>EQ 5GM BASE/VIAL</u>	<u>A063076</u>	<u>001</u>	Dec 21, 1990	
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A063076</u>	<u>002</u>	Sep 14, 2020	
<u>AP</u>		HOSPIRA INC	<u>EQ 10GM BASE/VIAL</u>	<u>A065455</u>	<u>001</u>	Apr 29, 2009	
<u>AP</u>		MYLAN LABS LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A065397</u>	<u>001</u>	Dec 30, 2008	
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A065397</u>	<u>002</u>	Dec 30, 2008	
<u>AP</u>			<u>EQ 5GM BASE/VIAL</u>	<u>A065432</u>	<u>001</u>	Dec 30, 2008	
<u>AP</u>	!		<u>EQ 10GM BASE/VIAL</u>	<u>A091554</u>	<u>001</u>	Sep 19, 2011	
<u>AP</u>		PHARM ASSOC	<u>EQ 500MG BASE/VIAL</u>	<u>A065401</u>	<u>001</u>	Jun 30, 2008	
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A065401</u>	<u>002</u>	Jun 30, 2008	
<u>AP</u>		SAGENT PHARMS	<u>EQ 5GM BASE/VIAL</u>	<u>A200837</u>	<u>001</u>	Aug 10, 2012	
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A200837</u>	<u>002</u>	Sep 02, 2014	
<u>AP</u>		SANDOZ	<u>EQ 500MG BASE/VIAL</u>	<u>A090250</u>	<u>001</u>	Apr 27, 2010	
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A090250</u>	<u>002</u>	Apr 27, 2010	
<u>AP</u>		SANDOZ INC	<u>EQ 5GM BASE/VIAL</u>	<u>A201048</u>	<u>001</u>	Aug 10, 2012	
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A201048</u>	<u>002</u>	Aug 10, 2012	
<u>AP</u>		XELLIA PHARMS APS	<u>EQ 5GM BASE/VIAL</u>	<u>A204125</u>	<u>001</u>	Dec 28, 2015	
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A204125</u>	<u>002</u>	Dec 28, 2015	
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A204107</u>	<u>001</u>	Dec 28, 2015	
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A204107</u>	<u>002</u>	Dec 28, 2015	
		VANCOMYCIN 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
		VANCOMYCIN HYDROCHLORIDE					
	!	HOSPIRA	EQ 1.5GM BASE/VIAL	A062912	003	Jul 10, 2020	
		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	

PRESCRIPTION DRUG PRODUCT LIST

VANCOMYCIN HYDROCHLORIDE

SOLUTION;INTRAVENOUS

VANCOMYCIN HYDROCHLORIDE

+	!	XELLIA PHARMS APS	EQ 500MG BASE/100ML (EQ 5MG BASE/ML)	N211962	001	Feb 15, 2019
+	!		EQ 750MG BASE/150ML (EQ 5MG BASE/ML)	N211962	005	May 13, 2020
+	!		EQ 1GM BASE/200ML (EQ 5MG BASE/ML)	N211962	002	Feb 15, 2019
+	!		EQ 1.25GM BASE/250ML (EQ 5MG BASE/ML)	N211962	006	May 13, 2020
+	!		EQ 1.5GM BASE/300ML (EQ 5MG BASE/ML)	N211962	003	Feb 15, 2019
+	!		EQ 1.75GM BASE/350ML (EQ 5MG BASE/ML)	N211962	007	May 13, 2020
+	!		EQ 2GM BASE/400ML (EQ 5MG BASE/ML)	N211962	004	Feb 15, 2019

VANDETANIB

TABLET;ORAL

CAPRELSA

+		GENZYME CORP	100MG	N022405	001	Apr 06, 2011
+	!		300MG	N022405	002	Apr 06, 2011

VARDENAFIL HYDROCHLORIDE

TABLET;ORAL

LEVITRA

AB	+	!	BAYER HLTHCARE	EQ 20MG BASE	N021400	004	Aug 19, 2003
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VARDENAFIL HYDROCHLORIDE

AB			ALEMBIC PHARMS LTD	EQ 2.5MG BASE	A214031	001	Aug 04, 2020
AB				EQ 5MG BASE	A214031	002	Aug 04, 2020
AB				EQ 10MG BASE	A214031	003	Aug 04, 2020
AB				EQ 20MG BASE	A214031	004	Aug 04, 2020
AB			CROSSMEDIKA SA	EQ 2.5MG BASE	A209057	001	Oct 31, 2018
AB				EQ 5MG BASE	A209057	002	Oct 31, 2018
AB				EQ 10MG BASE	A209057	003	Oct 31, 2018
AB				EQ 20MG BASE	A209057	004	Oct 31, 2018
AB			MACLEODS PHARMS LTD	EQ 2.5MG BASE	A204632	001	Oct 22, 2019
AB				EQ 5MG BASE	A204632	002	Oct 22, 2019
AB				EQ 10MG BASE	A204632	003	Oct 22, 2019
AB				EQ 20MG BASE	A204632	004	Oct 22, 2019
AB			TEVA PHARMS	EQ 2.5MG BASE	A091347	001	May 03, 2012
AB				EQ 5MG BASE	A091347	002	May 03, 2012
AB				EQ 10MG BASE	A091347	003	May 03, 2012
AB				EQ 20MG BASE	A091347	004	May 03, 2012
AB			ZYDUS PHARMS	EQ 2.5MG BASE	A208960	001	Oct 31, 2018
AB				EQ 5MG BASE	A208960	002	Oct 31, 2018
AB				EQ 10MG BASE	A208960	003	Oct 31, 2018
AB				EQ 20MG BASE	A208960	004	Oct 31, 2018

TABLET, ORALLY DISINTEGRATING;ORAL

STAXYN

AB	+	!	BAYER HLTHCARE	10MG	N200179	001	Jun 17, 2010
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VARDENAFIL HYDROCHLORIDE

AB			ALEMBIC PHARMS LTD	10MG	A208324	001	Nov 16, 2018
AB			MACLEODS PHARMS LTD	10MG	A205988	001	Mar 10, 2020

VARENICLINE TARTRATE

SOLUTION;NASAL

TYRVAYA

+	!	OYSTER POINT PHARMA	EQ 0.03MG BASE/SPRAY	N213978	001	Oct 15, 2021
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TABLET;ORAL

CHANTIX

AB	+		PF PRISM CV	EQ 0.5MG BASE	N021928	001	May 10, 2006
AB	+	!		EQ 1MG BASE	N021928	002	May 10, 2006

VARENICLINE TARTRATE

AB			PAR PHARM INC	EQ 0.5MG BASE	A201785	001	Aug 11, 2021
AB				EQ 1MG BASE	A201785	002	Aug 11, 2021

VASOPRESSIN

SOLUTION;INTRAVENOUS

VASOPRESSIN

AP			EAGLE PHARMS	20UNITS/ML (20UNITS/ML)	A211538	001	Dec 15, 2021
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VASOSTRICT

AP	+	!	PAR STERILE PRODUCTS	20UNITS/ML (20UNITS/ML)	N204485	001	Apr 17, 2014
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VASOPRESSIN

+	!	AM REGENT	20UNITS/ML (20UNITS/ML)	N212593	001	Aug 03, 2020
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VASOSTRICT

+	!	PAR STERILE PRODUCTS	20UNITS/100ML (0.2UNITS/ML)	N204485	005	Apr 21, 2021
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+	!		40UNITS/100ML (0.4UNITS/ML)	N204485	003	Apr 15, 2020
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+	!		60UNITS/100ML (0.6UNITS/ML)	N204485	004	Apr 15, 2020
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PRESCRIPTION DRUG PRODUCT LIST

VASOPRESSIN

SOLUTION; INTRAVENOUS

VASOSTRICT

+!

200UNITS/10ML (20UNITS/ML)

N204485 002 Dec 17, 2016

VECURONIUM BROMIDE

INJECTABLE; INJECTION

VECURONIUM BROMIDE

<u>AP</u>	EUGIA PHARMA	<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	<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>	HIKMA	<u>10MG/VIAL</u>	<u>A075549 001</u>	Jun 13, 2000
<u>AP</u>		<u>10MG/VIAL</u>	<u>A203725 001</u>	Jul 30, 2019
<u>AP</u>		<u>20MG/VIAL</u>	<u>A075549 002</u>	Jun 13, 2000
<u>AP</u>		<u>20MG/VIAL</u>	<u>A203725 002</u>	Jul 30, 2019
<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 INC	<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 PHARM	<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

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>	+ UPJOHN	<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
	<u>VENLAFAXINE HYDROCHLORIDE</u>			
<u>AB</u>	ANNORA PHARMA	<u>EQ 37.5MG BASE</u>	<u>A212277 001</u>	Jul 08, 2019
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A212277 002</u>	Jul 08, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A212277 003</u>	Jul 08, 2019
<u>AB</u>	ATLANTIDE	<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
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A078865 003</u>	Apr 14, 2011
<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>	INTELLIPHARMACEUTICS	<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>	INVENTIA HLTHCARE	<u>EQ 37.5MG BASE</u>	<u>A203332 001</u>	Mar 12, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A203332 002</u>	Mar 12, 2020
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A203332 003</u>	Mar 12, 2020
<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>	ORBION PHARMS	<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

PRESCRIPTION DRUG PRODUCT LIST

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>	TORRENT	<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>	YICHANG HUMANWELL	<u>EQ 37.5MG BASE</u>	<u>A214654 001</u>	Aug 06, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214654 002</u>	Aug 06, 2021
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214654 003</u>	Aug 06, 2021
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 37.5MG BASE</u>	<u>A090174 001</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090174 002</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A090174 003</u>	Apr 14, 2011

TABLET;ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A078932 001</u>	Dec 14, 2010
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078932 002</u>	Dec 14, 2010
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078932 003</u>	Dec 14, 2010
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078932 004</u>	Dec 14, 2010
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078932 005</u>	Dec 14, 2010
<u>AB</u>	AMNEAL PHARMS	<u>EQ 25MG BASE</u>	<u>A079098 001</u>	May 11, 2010
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A079098 002</u>	May 11, 2010
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A079098 003</u>	May 11, 2010
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A079098 004</u>	May 11, 2010
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A079098 005</u>	May 11, 2010
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 25MG BASE</u>	<u>A090555 001</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A090555 002</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A090555 003</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090555 004</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090555 005</u>	Apr 07, 2010
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A206250 001</u>	Nov 21, 2018
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A206250 002</u>	Nov 21, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A206250 003</u>	Nov 21, 2018
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A206250 004</u>	Nov 21, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A206250 005</u>	Nov 21, 2018
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 25MG BASE</u>	<u>A078301 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078301 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078301 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078301 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078301 005</u>	Jun 13, 2008
<u>AB</u>	HERITAGE PHARMS INC	<u>EQ 25MG BASE</u>	<u>A078554 001</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078554 002</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078554 003</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078554 004</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078554 005</u>	Jan 09, 2009
<u>AB</u>	SUN PHARM INDS INC	<u>EQ 25MG BASE</u>	<u>A078627 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078627 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078627 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078627 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078627 005</u>	Jun 13, 2008
<u>AB</u>	TEVA	<u>EQ 25MG BASE</u>	<u>A076690 001</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A076690 002</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076690 003</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A076690 004</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076690 005</u>	Aug 03, 2006
<u>AB</u>	YAOPHARMA CO LTD	<u>EQ 25MG BASE</u>	<u>A202036 001</u>	May 28, 2015
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A202036 002</u>	May 28, 2015
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202036 003</u>	May 28, 2015
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A202036 004</u>	May 28, 2015
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202036 005</u>	May 28, 2015
<u>AB</u>	ZYDUS PHARMS USA	<u>EQ 25MG BASE</u>	<u>A077653 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A077653 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077653 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A077653 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077653 005</u>	Jun 13, 2008

TABLET, EXTENDED RELEASE;ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>	ALKEM LABS LTD	<u>EQ 37.5MG BASE</u>	<u>A214127 001</u>	Jun 10, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214127 002</u>	Jun 10, 2021

PRESCRIPTION DRUG PRODUCT LIST

VENLAFAXINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214127 003</u>	Jun 10, 2021
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A214127 004</u>	Jun 10, 2021
<u>AB</u>	APPCO	<u>EQ 150MG BASE</u>	<u>A214609 001</u>	Jun 30, 2021
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A214609 002</u>	Jun 30, 2021
<u>AB</u>	ASCENT PHARMS INC	<u>EQ 37.5MG BASE</u>	<u>A214419 001</u>	Oct 21, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214419 002</u>	Oct 21, 2020
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214419 003</u>	Oct 21, 2020
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A214419 004</u>	Oct 21, 2020
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A211323 001</u>	Aug 29, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A211323 002</u>	Aug 29, 2019
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A211323 003</u>	Aug 29, 2019
<u>AB</u>	DEXCEL PHARMA	<u>EQ 75MG BASE</u>	<u>A213927 001</u>	Jan 21, 2021
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A209193 001</u>	Oct 31, 2019
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A209193 002</u>	Oct 31, 2019
<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>	PHARMADAX INC	<u>EQ 75MG BASE</u>	<u>A214423 001</u>	Jan 04, 2022
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214423 002</u>	Jan 04, 2022
<u>AB</u>	SUN PHARM	<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>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>240MG</u>	<u>A075138 003</u>	Apr 20, 1999
<u>VERELAN</u>				
<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
	+!	360MG	N019614 004	May 10, 1996
VERELAN PM				
	+ RECRO GAINESVILLE	100MG	N020943 001	Nov 25, 1998
	+	200MG	N020943 002	Nov 25, 1998
	+!	300MG	N020943 003	Nov 25, 1998

INJECTABLE; INJECTION

VERAPAMIL HYDROCHLORIDE

<u>AP</u>	! EUGIA PHARMA	<u>2.5MG/ML</u>	<u>A212965 001</u>	Jul 06, 2020
<u>AP</u>	GLAND PHARMA LTD	<u>2.5MG/ML</u>	<u>A214361 001</u>	Oct 15, 2020
<u>AP</u>	ZYDUS PHARMS	<u>2.5MG/ML</u>	<u>A214215 001</u>	Oct 15, 2020

SOLUTION; INTRAVENOUS

VERAPAMIL HYDROCHLORIDE

<u>AP</u>	AMNEAL	<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>	AREVA PHARMS	<u>5MG/2ML (2.5MG/ML)</u>	<u>A213352 002</u>	Sep 16, 2020
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A213352 001</u>	Mar 17, 2020
<u>AP</u>	CAPLIN	<u>5MG/2ML (2.5MG/ML)</u>	<u>A213232 001</u>	Mar 25, 2020
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A213232 002</u>	Mar 25, 2020
<u>AP</u>	EXELA PHARMA	<u>5MG/2ML (2.5MG/ML)</u>	<u>N018925 001</u>	Mar 30, 1984
<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	<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>A211035 002</u>	Jun 18, 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>10MG/4ML (2.5MG/ML)</u>	<u>A211015 002</u>	Jun 18, 2018

TABLET; ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	HERITAGE PHARMS INC	<u>40MG</u>	<u>A071881 002</u>	Oct 14, 2015
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PRESCRIPTION DRUG PRODUCT LIST

VERAPAMIL HYDROCHLORIDE

TABLET;ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>		<u>80MG</u>	<u>A071881</u>	<u>003</u>	Apr 05, 1988
<u>AB</u>	!	<u>120MG</u>	<u>A071881</u>	<u>001</u>	Apr 05, 1988
<u>AB</u>	WATSON LABS	<u>40MG</u>	<u>A072924</u>	<u>001</u>	Jun 29, 1993
<u>AB</u>		<u>80MG</u>	<u>A070995</u>	<u>001</u>	Oct 01, 1986
<u>AB</u>		<u>120MG</u>	<u>A070994</u>	<u>001</u>	Oct 01, 1986

TABLET, EXTENDED RELEASE;ORAL

CALAN SR

<u>AB</u>	+!	PFIZER	<u>120MG</u>	<u>N019152</u>	<u>003</u>	Mar 06, 1991
<u>AB</u>	+!		<u>240MG</u>	<u>N019152</u>	<u>001</u>	Dec 16, 1986

VERAPAMIL HYDROCHLORIDE

<u>AB</u>		CADILA PHARMS LTD	<u>180MG</u>	<u>A206173</u>	<u>001</u>	May 05, 2017
<u>AB</u>			<u>240MG</u>	<u>A206173</u>	<u>002</u>	May 05, 2017
<u>AB</u>		GLENMARK GENERICS	<u>120MG</u>	<u>A090700</u>	<u>001</u>	Aug 03, 2011
<u>AB</u>	!		<u>180MG</u>	<u>A090700</u>	<u>002</u>	Aug 03, 2011
<u>AB</u>			<u>240MG</u>	<u>A078906</u>	<u>001</u>	Sep 17, 2009
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>120MG</u>	<u>A073568</u>	<u>002</u>	Oct 10, 1997
<u>AB</u>			<u>180MG</u>	<u>A074330</u>	<u>001</u>	Jan 31, 1994
<u>AB</u>			<u>240MG</u>	<u>A073568</u>	<u>001</u>	Jul 31, 1992
<u>AB</u>		STRIDES PHARMA	<u>120MG</u>	<u>A075072</u>	<u>001</u>	May 25, 1999
<u>AB</u>			<u>240MG</u>	<u>A075072</u>	<u>003</u>	May 25, 1999
<u>AB</u>		SUN PHARM INDS INC	<u>120MG</u>	<u>A090529</u>	<u>001</u>	Dec 30, 2011
<u>AB</u>			<u>180MG</u>	<u>A090529</u>	<u>002</u>	Dec 30, 2011
<u>AB</u>			<u>240MG</u>	<u>A090529</u>	<u>003</u>	Dec 30, 2011

VERICIGUAT

TABLET;ORAL

VERQUVO

+	MERCK SHARP DOHME	2.5MG	N214377	001	Jan 19, 2021
+		5MG	N214377	002	Jan 19, 2021
+	!	10MG	N214377	003	Jan 19, 2021

VERTEPORFIN

INJECTABLE;INJECTION

VISUDYNE

+	VALEANT LUXEMBOURG	15MG/VIAL	N021119	001	Apr 12, 2000
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VIBEGRON

TABLET;ORAL

GEMTESA

+	UROVANT	75MG	N213006	001	Dec 23, 2020
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VIGABATRIN

FOR SOLUTION;ORAL

SABRIL

<u>AA</u>	+	LUNDBECK PHARMS LLC	<u>500MG/PACKET</u>	<u>N022006</u>	<u>001</u>	Aug 21, 2009
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VIGABATRIN

<u>AA</u>		ACCORD HLTHCARE	<u>500MG/PACKET</u>	<u>A214425</u>	<u>001</u>	Nov 13, 2020
<u>AA</u>		ALKEM LABS LTD	<u>500MG/PACKET</u>	<u>A213375</u>	<u>001</u>	Dec 02, 2020
<u>AA</u>		AMNEAL PHARMS	<u>500MG/PACKET</u>	<u>A210155</u>	<u>001</u>	Mar 13, 2018
<u>AA</u>		ANNORA PHARMA	<u>500MG/PACKET</u>	<u>A213519</u>	<u>001</u>	Jan 26, 2021
<u>AA</u>		AUROBINDO PHARMA LTD	<u>500MG/PACKET</u>	<u>A213899</u>	<u>001</u>	Sep 29, 2021
<u>AA</u>		DEXCEL PHARMA	<u>500MG/PACKET</u>	<u>A214992</u>	<u>001</u>	May 13, 2021
<u>AA</u>		DR REDDYS LABS LTD	<u>500MG/PACKET</u>	<u>A211481</u>	<u>001</u>	Nov 20, 2018
<u>AA</u>		GRANULES	<u>500MG/PACKET</u>	<u>A213469</u>	<u>001</u>	Apr 24, 2020
<u>AA</u>		INVAGEN PHARMS	<u>500MG/PACKET</u>	<u>A211592</u>	<u>001</u>	Dec 03, 2019
<u>AA</u>		PAR PHARM INC	<u>500MG/PACKET</u>	<u>A208218</u>	<u>001</u>	Apr 27, 2017
<u>AA</u>		PROPEL PHARMA	<u>500MG/PACKET</u>	<u>A213390</u>	<u>001</u>	Jul 29, 2021
<u>AA</u>		TEVA PHARMS USA	<u>500MG/PACKET</u>	<u>A209824</u>	<u>001</u>	Apr 23, 2018

VIGADRONE

<u>AA</u>		AUCTA	<u>500MG/PACKET</u>	<u>A210196</u>	<u>001</u>	Jun 21, 2018
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TABLET;ORAL

SABRIL

<u>AB</u>	+	LUNDBECK PHARMS LLC	<u>500MG</u>	<u>N020427</u>	<u>001</u>	Aug 21, 2009
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VIGABATRIN

<u>AB</u>		DEXCEL PHARMA	<u>500MG</u>	<u>A215109</u>	<u>001</u>	Sep 23, 2021
<u>AB</u>		DR REDDYS	<u>500MG</u>	<u>A211539</u>	<u>001</u>	Jan 29, 2021
<u>AB</u>		TEVA PHARMS USA	<u>500MG</u>	<u>A209822</u>	<u>001</u>	Jan 14, 2019

PRESCRIPTION DRUG PRODUCT LIST

VILAZODONE HYDROCHLORIDE

TABLET; ORAL

VIIBRYD

<u>AB</u>	+	ALLERGAN	<u>10MG</u>	<u>N022567</u>	<u>001</u>	Jan 21, 2011
<u>AB</u>	+		<u>20MG</u>	<u>N022567</u>	<u>002</u>	Jan 21, 2011
<u>AB</u>	+		<u>40MG</u>	<u>N022567</u>	<u>003</u>	Jan 21, 2011

VILAZODONE HYDROCHLORIDE

<u>AB</u>		ACCORD HLTHCARE	<u>10MG</u>	<u>A208209</u>	<u>001</u>	Apr 27, 2021
<u>AB</u>			<u>20MG</u>	<u>A208209</u>	<u>002</u>	Apr 27, 2021
<u>AB</u>			<u>40MG</u>	<u>A208209</u>	<u>003</u>	Apr 27, 2021
<u>AB</u>		ALEMBIC PHARMS LTD	<u>10MG</u>	<u>A208202</u>	<u>001</u>	Jan 10, 2020
<u>AB</u>			<u>20MG</u>	<u>A208202</u>	<u>002</u>	Jan 10, 2020
<u>AB</u>			<u>40MG</u>	<u>A208202</u>	<u>003</u>	Jan 10, 2020
<u>AB</u>		INVAGEN PHARMS	<u>10MG</u>	<u>A208200</u>	<u>001</u>	Apr 07, 2021
<u>AB</u>			<u>40MG</u>	<u>A208200</u>	<u>002</u>	Apr 07, 2021

VILOXAZINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

QELBREE

	+	SUPERNUS PHARMS	EQ 100MG BASE	N211964	001	Apr 02, 2021
	+		EQ 150MG BASE	N211964	002	Apr 02, 2021
	+		EQ 200MG BASE	N211964	003	Apr 02, 2021

VILTOLARSEN

SOLUTION; INTRAVENOUS

VILTEPSO

	+	NIPPON SHINYAKU	250MG/5ML (50MG/ML)	N212154	001	Aug 12, 2020
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VINBLASTINE SULFATE

INJECTABLE; INJECTION

VINBLASTINE SULFATE

	!	FRESENIUS KABI USA	1MG/ML	A089515	001	Apr 29, 1987
	!	HIKMA	10MG/VIAL	A089395	001	Apr 09, 1987

VINCRIStINE SULFATE

INJECTABLE; INJECTION

VINCRIStINE SULFATE PFS

<u>AP</u>	!	HOSPIRA	<u>1MG/ML</u>	<u>A071484</u>	<u>001</u>	Apr 19, 1988
<u>AP</u>		TEVA PHARMS USA	<u>1MG/ML</u>	<u>A075493</u>	<u>001</u>	Sep 01, 1999

VINORELBINE TARTRATE

INJECTABLE; INJECTION

VINORELBINE TARTRATE

<u>AP</u>		ACTAVIS TOTOWA	<u>EQ 10MG BASE/ML</u>	<u>A078011</u>	<u>001</u>	Jul 22, 2009
<u>AP</u>		DR REDDYS	<u>EQ 10MG BASE/ML</u>	<u>A202017</u>	<u>001</u>	Sep 12, 2013
<u>AP</u>		HIKMA	<u>EQ 10MG BASE/ML</u>	<u>A075992</u>	<u>001</u>	Jun 10, 2003
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A076461</u>	<u>001</u>	Dec 11, 2003
<u>AP</u>		JIANGSU HANSON PHARM	<u>EQ 10MG BASE/ML</u>	<u>A091106</u>	<u>001</u>	Sep 26, 2012
<u>AP</u>		TEVA PHARMS USA	<u>EQ 10MG BASE/ML</u>	<u>A076028</u>	<u>001</u>	Feb 03, 2003

VISMODEGIB

CAPSULE; ORAL

ERIVEDGE

	+	GENENTECH	150MG	N203388	001	Jan 30, 2012
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VITAMIN A PALMITATE

INJECTABLE; INJECTION

AQUASOL A

	+	CASPER PHARMA LLC	EQ 50,000 UNITS BASE/ML	N006823	001	
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VOCLOSPORIN

CAPSULE; ORAL

LUPKYNIS

	+	AURINIA	7.9MG	N213716	001	Jan 22, 2021
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VORAPAXAR SULFATE

TABLET; ORAL

ZONTIVITY

	+	XSPIRE PHARMA	EQ 2.08MG BASE	N204886	001	May 08, 2014
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VORICONAZOLE

FOR SUSPENSION; ORAL

VFEND

<u>AB</u>	+	PF PRISM CV	<u>200MG/5ML</u>	<u>N021630</u>	<u>001</u>	Dec 19, 2003
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VORICONAZOLE

<u>AB</u>		AMNEAL PHARMS	<u>200MG/5ML</u>	<u>A205034</u>	<u>001</u>	Apr 13, 2016
<u>AB</u>		NOVEL LABS INC	<u>200MG/5ML</u>	<u>A206799</u>	<u>001</u>	May 31, 2016

PRESCRIPTION DRUG PRODUCT LIST

VORICONAZOLE

INJECTABLE; INTRAVENOUS

VFEND

AP	+	PF PRISM CV	200MG/VIAL	N021267	001	May 24, 2002
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VORICONAZOLE

AP		ALMAJECT	200MG/VIAL	A206398	001	Mar 23, 2016
AP		GLAND PHARMA LTD	200MG/VIAL	A211099	001	Mar 31, 2020
AP		HAINAN POLY PHARM	200MG/VIAL	A211661	001	Nov 30, 2018
AP		SANDOZ INC	200MG/VIAL	A090862	001	May 30, 2012
AP		ZYDUS PHARMS	200MG/VIAL	A208983	001	Jul 16, 2018

POWDER; INTRAVENOUS

VORICONAZOLE

AP		XELLIA PHARMS APS	200MG/VIAL	N208562	001	Mar 09, 2017
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TABLET; ORAL

VFEND

AB	+	PF PRISM CV	50MG	N021266	001	May 24, 2002
AB	+	!	200MG	N021266	002	May 24, 2002

VORICONAZOLE

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 LTD	50MG	A206837	001	Jan 22, 2016
AB			200MG	A206837	002	Jan 22, 2016
AB		CADILA	50MG	A206747	001	May 24, 2016
AB			200MG	A206747	002	May 24, 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	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

VORINOSTAT

CAPSULE; ORAL

ZOLINZA

+	!	MERCK	100MG	N021991	001	Oct 06, 2006
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VORTIOXETINE HYDROBROMIDE

TABLET; ORAL

TRINTELLIX

AB	+	TAKEDA PHARMS USA	EQ 5MG BASE	N204447	001	Sep 30, 2013
AB	+		EQ 10MG BASE	N204447	002	Sep 30, 2013
AB	+	!	EQ 20MG BASE	N204447	004	Sep 30, 2013

VORTIOXETINE HYDROBROMIDE

AB		ZYDUS PHARMS	EQ 5MG BASE	A211146	001	Sep 17, 2021
AB			EQ 10MG BASE	A211146	002	Sep 17, 2021
AB			EQ 20MG BASE	A211146	003	Sep 17, 2021

VOSORITIDE

POWDER; SUBCUTANEOUS

VOXZOGO

+	!	BIOMARIN PHARM	0.4MG/VIAL	N214938	001	Nov 19, 2021
+	!		0.56MG/VIAL	N214938	002	Nov 19, 2021
+	!		1.2MG/VIAL	N214938	003	Nov 19, 2021

VOXELOTOR

TABLET; ORAL

OXBRYTA

+	!	GLOBAL BLOOD THERAPS	500MG	N213137	001	Nov 25, 2019
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TABLET, FOR SUSPENSION; ORAL

OXBRYTA

+	!	GLOBAL BLOOD THERAPS	300MG	N216157	001	Dec 17, 2021
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PRESCRIPTION DRUG PRODUCT LIST

WARFARIN SODIUM

TABLET; ORAL

JANTOVEN

<u>AB</u>	UPSHER SMITH LABS	<u>1MG</u>	<u>A040416 001</u>	Oct 02, 2003
<u>AB</u>		<u>2MG</u>	<u>A040416 002</u>	Oct 02, 2003
<u>AB</u>		<u>2.5MG</u>	<u>A040416 003</u>	Oct 02, 2003
<u>AB</u>		<u>3MG</u>	<u>A040416 004</u>	Oct 02, 2003
<u>AB</u>		<u>4MG</u>	<u>A040416 005</u>	Oct 02, 2003
<u>AB</u>		<u>5MG</u>	<u>A040416 006</u>	Oct 02, 2003
<u>AB</u>		<u>6MG</u>	<u>A040416 007</u>	Oct 02, 2003
<u>AB</u>		<u>7.5MG</u>	<u>A040416 008</u>	Oct 02, 2003
<u>AB</u>		<u>10MG</u>	<u>A040416 009</u>	Oct 02, 2003

WARFARIN SODIUM

<u>AB</u>	AMNEAL PHARMS	<u>1MG</u>	<u>A202202 001</u>	Mar 04, 2013
<u>AB</u>		<u>2MG</u>	<u>A202202 002</u>	Mar 04, 2013
<u>AB</u>		<u>2.5MG</u>	<u>A202202 003</u>	Mar 04, 2013
<u>AB</u>		<u>3MG</u>	<u>A202202 004</u>	Mar 04, 2013
<u>AB</u>		<u>4MG</u>	<u>A202202 005</u>	Mar 04, 2013
<u>AB</u>		<u>5MG</u>	<u>A202202 006</u>	Mar 04, 2013
<u>AB</u>		<u>6MG</u>	<u>A202202 007</u>	Mar 04, 2013
<u>AB</u>		<u>7.5MG</u>	<u>A202202 008</u>	Mar 04, 2013
<u>AB</u>		<u>10MG</u>	<u>A202202 009</u>	Mar 04, 2013
<u>AB</u>	BARR	<u>1MG</u>	<u>A040145 001</u>	Mar 26, 1997
<u>AB</u>		<u>2MG</u>	<u>A040145 002</u>	Mar 26, 1997
<u>AB</u>		<u>2.5MG</u>	<u>A040145 003</u>	Mar 26, 1997
<u>AB</u>		<u>3MG</u>	<u>A040145 008</u>	Nov 05, 1998
<u>AB</u>		<u>4MG</u>	<u>A040145 004</u>	Mar 26, 1997
<u>AB</u>		<u>5MG</u>	<u>A040145 005</u>	Mar 26, 1997
<u>AB</u>		<u>6MG</u>	<u>A040145 009</u>	Nov 05, 1998
<u>AB</u>		<u>7.5MG</u>	<u>A040145 006</u>	Mar 26, 1997
<u>AB</u>		<u>10MG</u>	<u>A040145 007</u>	Mar 26, 1997
<u>AB</u>	INVAGEN PHARMS	<u>1MG</u>	<u>A090935 001</u>	May 25, 2011
<u>AB</u>		<u>2MG</u>	<u>A090935 002</u>	May 25, 2011
<u>AB</u>		<u>2.5MG</u>	<u>A090935 003</u>	May 25, 2011
<u>AB</u>		<u>3MG</u>	<u>A090935 004</u>	May 25, 2011
<u>AB</u>		<u>4MG</u>	<u>A090935 005</u>	May 25, 2011
<u>AB</u>		<u>5MG</u>	<u>A090935 006</u>	May 25, 2011
<u>AB</u>		<u>6MG</u>	<u>A090935 007</u>	May 25, 2011
<u>AB</u>		<u>7.5MG</u>	<u>A090935 008</u>	May 25, 2011
<u>AB</u>		<u>10MG</u>	<u>A090935 009</u>	May 25, 2011
<u>AB</u>	IPCA LABS LTD	<u>1MG</u>	<u>A200104 001</u>	Jun 27, 2013
<u>AB</u>		<u>2MG</u>	<u>A200104 002</u>	Jun 27, 2013
<u>AB</u>		<u>2.5MG</u>	<u>A200104 003</u>	Jun 27, 2013
<u>AB</u>		<u>3MG</u>	<u>A200104 004</u>	Jun 27, 2013
<u>AB</u>		<u>4MG</u>	<u>A200104 005</u>	Jun 27, 2013
<u>AB</u>		<u>5MG</u>	<u>A200104 006</u>	Jun 27, 2013
<u>AB</u>		<u>6MG</u>	<u>A200104 007</u>	Jun 27, 2013
<u>AB</u>		<u>7.5MG</u>	<u>A200104 008</u>	Jun 27, 2013
<u>AB</u>		<u>10MG</u>	<u>A200104 009</u>	Jun 27, 2013
<u>AB</u>	PLIVA	<u>1MG</u>	<u>A040616 009</u>	Jul 05, 2006
<u>AB</u>		<u>2MG</u>	<u>A040616 001</u>	Jul 05, 2006
<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	<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

PRESCRIPTION DRUG PRODUCT LIST

WARFARIN SODIUM

TABLET; ORAL

WARFARIN SODIUM

<u>AB</u>		<u>5MG</u>	<u>A040663</u>	<u>006</u>	May 30, 2006
<u>AB</u>		<u>6MG</u>	<u>A040663</u>	<u>007</u>	May 30, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A040663</u>	<u>008</u>	May 30, 2006
<u>AB</u>		<u>10MG</u>	<u>A040663</u>	<u>009</u>	May 30, 2006

XENON XE-133

GAS; INHALATION

XENON XE 133

CURIUM

10mCi/VIAL

N018327 001 Mar 09, 1982

20mCi/VIAL

N018327 002 Mar 09, 1982

LANTHEUS MEDCL

10mCi/VIAL

N017284 001

20mCi/VIAL

N017284 002

ZAFIRLUKAST

TABLET; ORAL

ACCOLATE

<u>AB</u>	+	STRIDES PHARMA	<u>10MG</u>	<u>N020547</u>	<u>003</u>	Sep 17, 1999
<u>AB</u>	+	!	<u>20MG</u>	<u>N020547</u>	<u>001</u>	Sep 26, 1996

ZAFIRLUKAST

<u>AB</u>		ANNORA PHARMA	<u>10MG</u>	<u>A212475</u>	<u>001</u>	Sep 10, 2020
<u>AB</u>			<u>20MG</u>	<u>A212475</u>	<u>002</u>	Sep 10, 2020
<u>AB</u>		DR REDDYS LABS LTD	<u>10MG</u>	<u>A090372</u>	<u>001</u>	Nov 18, 2010
<u>AB</u>			<u>20MG</u>	<u>A090372</u>	<u>002</u>	Nov 18, 2010

ZALEPLON

CAPSULE; ORAL

SONATA

<u>AB</u>	+	PFIZER	<u>5MG</u>	<u>N020859</u>	<u>001</u>	Aug 13, 1999
<u>AB</u>	+	!	<u>10MG</u>	<u>N020859</u>	<u>002</u>	Aug 13, 1999

ZALEPLON

<u>AB</u>		AUROBINDO PHARMA	<u>5MG</u>	<u>A078829</u>	<u>001</u>	Jun 06, 2008
<u>AB</u>			<u>10MG</u>	<u>A078829</u>	<u>002</u>	Jun 06, 2008
<u>AB</u>		CIPLA LTD	<u>5MG</u>	<u>A077505</u>	<u>001</u>	Jun 20, 2008
<u>AB</u>			<u>10MG</u>	<u>A077505</u>	<u>002</u>	Jun 20, 2008
<u>AB</u>		HIKMA	<u>5MG</u>	<u>A077237</u>	<u>001</u>	Jun 06, 2008
<u>AB</u>			<u>10MG</u>	<u>A077237</u>	<u>002</u>	Jun 06, 2008
<u>AB</u>		ORBION PHARMS	<u>5MG</u>	<u>A090374</u>	<u>001</u>	Sep 17, 2009
<u>AB</u>			<u>10MG</u>	<u>A090374</u>	<u>002</u>	Sep 17, 2009
<u>AB</u>		UNICHEM	<u>5MG</u>	<u>A078989</u>	<u>001</u>	Jun 06, 2008
<u>AB</u>			<u>10MG</u>	<u>A078989</u>	<u>002</u>	Jun 06, 2008

ZANAMIVIR

POWDER; INHALATION

RELENZA

+! GLAXOSMITHKLINE

5MG

N021036 001 Jul 26, 1999

ZANUBRUTINIB

CAPSULE; ORAL

BRUKINSA

+! BEIGENE

80MG

N213217 001 Nov 14, 2019

ZICONOTIDE ACETATE

INJECTABLE; INTRATHECAL

PRIALT

+! TERSERA

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

<u>AB</u>	+	VIIV HLTHCARE	<u>100MG</u>	<u>N019655</u>	<u>001</u>	Mar 19, 1987
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ZIDOVUDINE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A078128</u>	<u>001</u>	Mar 27, 2006
<u>AB</u>		CIPLA LTD	<u>100MG</u>	<u>A078349</u>	<u>001</u>	May 23, 2007

INJECTABLE; INJECTION

RETROVIR

+! VIIV HLTHCARE

10MG/ML

N019951 001 Feb 02, 1990

SOLUTION; ORAL

RETROVIR

<u>AA</u>	+	VIIV HLTHCARE	<u>50MG/5ML</u>	<u>N019910</u>	<u>001</u>	Sep 28, 1989
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PRESCRIPTION DRUG PRODUCT LIST

ZIDOVUDINE

SOLUTION;ORAL

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

ZILEUTON

TABLET;ORAL

ZYFLO

+	!	CHIESI	600MG	N020471 003	Dec 09, 1996
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TABLET, EXTENDED RELEASE;ORAL

ZILEUTON

AB	DASH PHARMS	600MG	A211390 001	Oct 23, 2020
AB	RISING	600MG	A204929 001	Mar 17, 2017
AB	STRIDES PHARMA	600MG	A212670 001	Dec 16, 2019

ZYFLO CR

AB	+	!	CHIESI	600MG	N022052 001	May 30, 2007
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ZILEUTON

BX	LUPIN LTD	600MG	A211972 001	Nov 05, 2019
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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

AP	EXELA PHARMA	EQ 1MG ZINC/ML	A212007 001	May 21, 2021
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ZINC CHLORIDE IN PLASTIC CONTAINER

AP	+	!	HOSPIRA	EQ 1MG ZINC/ML	N018959 001	Jun 26, 1986
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ZINC SULFATE

SOLUTION;INTRAVENOUS

ZINC SULFATE

+	!	AM REGENT	EQ 10MG BASE/10ML (EQ 1MG BASE/ML)	N209377 003	Apr 15, 2020
+	!		EQ 25MG BASE/5ML (EQ 5MG BASE/ML)	N209377 002	Jul 18, 2019
+	!		EQ 30MG BASE/10ML (EQ 3MG BASE/ML)	N209377 001	Jul 18, 2019

ZIPRASIDONE HYDROCHLORIDE

CAPSULE;ORAL

GEODON

AB	+	!	UPJOHN	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	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	CADILA	EQ 20MG BASE	A208988 001	Aug 22, 2017
AB		EQ 40MG BASE	A208988 002	Aug 22, 2017
AB		EQ 60MG BASE	A208988 003	Aug 22, 2017
AB		EQ 80MG BASE	A208988 004	Aug 22, 2017
AB	CHARTWELL RX	EQ 20MG BASE	A090348 001	Sep 05, 2012
AB		EQ 40MG BASE	A090348 002	Sep 05, 2012
AB		EQ 60MG BASE	A090348 003	Sep 05, 2012
AB		EQ 80MG BASE	A090348 004	Sep 05, 2012
AB	DR REDDYS LABS INC	EQ 20MG BASE	A077565 001	Mar 02, 2012
AB		EQ 40MG BASE	A077565 002	Mar 02, 2012
AB		EQ 60MG BASE	A077565 003	Mar 02, 2012
AB		EQ 80MG BASE	A077565 004	Mar 02, 2012

PRESCRIPTION DRUG PRODUCT LIST

ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

ZIPRASIDONE HYDROCHLORIDE

<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

ZIPRASIDONE MESYLATE

INJECTABLE; INTRAMUSCULAR

GEODON

<u>AP</u>	+! UPJOHN	<u>EQ 20MG BASE/ML</u>	<u>N020919 001</u>	Jun 21, 2002
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ZIPRASIDONE MESYLATE

<u>AP</u>	GLAND PHARMA LTD	<u>EQ 20MG BASE/ML</u>	<u>A211908 001</u>	Dec 26, 2019
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ZOLEDRONIC ACID

INJECTABLE; INTRAVENOUS

RECLAST

<u>AP</u>	+! NOVARTIS	<u>EQ 5MG BASE/100ML</u>	<u>N021817 001</u>	Apr 16, 2007
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ZOLEDRONIC

<u>AP</u>	GLAND PHARMA LTD	<u>EQ 4MG BASE/100ML</u>	<u>A205749 001</u>	Jun 29, 2018
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ZOLEDRONIC ACID

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 4MG BASE/5ML</u>	<u>A205279 001</u>	Nov 28, 2016
<u>AP</u>	AKORN	<u>EQ 4MG BASE/5ML</u>	<u>A202548 001</u>	May 22, 2014
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A200918 001</u>	Aug 21, 2014
<u>AP</u>	APOTEX	<u>EQ 5MG BASE/100ML</u>	<u>A204367 001</u>	Dec 24, 2015
<u>AP</u>	BPI LABS LLC	<u>EQ 4MG BASE/5ML</u>	<u>A207341 001</u>	Dec 29, 2017
<u>AP</u>	BRECKENRIDGE	<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>	DR REDDYS LABS LTD	<u>EQ 4MG BASE/5ML</u>	<u>A091186 001</u>	Mar 04, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A091363 001</u>	Mar 29, 2013
<u>AP</u>	EUGIA PHARMA	<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>	FRESENIUS KABI USA	<u>EQ 4MG BASE/5ML</u>	<u>A091516 001</u>	Apr 23, 2015
<u>AP</u>	GLAND	<u>EQ 5MG BASE/100ML</u>	<u>A204217 001</u>	Aug 18, 2016
<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>A209578 001</u>	Aug 08, 2019
<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>	INGENUS PHARMS LLC	<u>EQ 4MG BASE/5ML</u>	<u>A208968 001</u>	Feb 19, 2020
<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>	PUNISKA	<u>EQ 4MG BASE/100ML</u>	<u>A210174 001</u>	Oct 27, 2017
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 4MG BASE/5ML</u>	<u>A091493 001</u>	Nov 24, 2014
<u>AP</u>	USV	<u>EQ 4MG BASE/5ML</u>	<u>A202923 001</u>	Sep 04, 2014

ZOMETA

<u>AP</u>	+! NOVARTIS	<u>EQ 4MG BASE/5ML</u>	<u>N021223 002</u>	Mar 07, 2003
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<u>AP</u>	+!	<u>EQ 4MG BASE/100ML</u>	<u>N021223 003</u>	Jun 17, 2011
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SOLUTION; INTRAVENOUS

ZOLEDRONIC ACID

	HOSPIRA INC	<u>EQ 4MG BASE/100ML (EQ 0.04MG BASE/ML)</u>	<u>N204016 001</u>	Dec 28, 2015
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ZOLMITRIPTAN

SPRAY; NASAL

ZOLMITRIPTAN

<u>AB</u>	PADAGIS ISRAEL	<u>2.5MG/SPRAY</u>	<u>A212469 001</u>	Sep 30, 2021
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<u>AB</u>		<u>5MG/SPRAY</u>	<u>A212469 002</u>	Sep 30, 2021
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ZOMIG

<u>AB</u>	+ ASTRAZENECA	<u>2.5MG/SPRAY</u>	<u>N021450 003</u>	Sep 16, 2013
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<u>AB</u>	+!	<u>5MG/SPRAY</u>	<u>N021450 004</u>	Sep 30, 2003
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PRESCRIPTION DRUG PRODUCT LIST

ZOLMITRIPTAN

TABLET; ORAL

ZOLMITRIPTAN

<u>AB</u>	AJANTA PHARMA LTD	<u>2.5MG</u>	<u>A204041 001</u>	May 20, 2016
<u>AB</u>		<u>5MG</u>	<u>A204041 002</u>	May 20, 2016
<u>AB</u>	ALEMBIC PHARMS LTD	<u>2.5MG</u>	<u>A204232 001</u>	Sep 30, 2015
<u>AB</u>		<u>5MG</u>	<u>A204232 002</u>	Sep 30, 2015
<u>AB</u>	APPCO	<u>2.5MG</u>	<u>A206973 001</u>	Jun 30, 2017
<u>AB</u>		<u>5MG</u>	<u>A206973 002</u>	Jun 30, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A207021 001</u>	May 11, 2016
<u>AB</u>		<u>5MG</u>	<u>A207021 002</u>	May 11, 2016
<u>AB</u>	GLENMARK GENERICS	<u>2.5MG</u>	<u>A201779 001</u>	May 14, 2013
<u>AB</u>		<u>5MG</u>	<u>A201779 002</u>	May 14, 2013
<u>AB</u>	INVAGEN PHARMS	<u>2.5MG</u>	<u>A204284 001</u>	Apr 09, 2014
<u>AB</u>		<u>5MG</u>	<u>A204284 002</u>	Apr 09, 2014
<u>AB</u>	JUBILANT GENERICS	<u>2.5MG</u>	<u>A202279 001</u>	Nov 20, 2014
<u>AB</u>		<u>5MG</u>	<u>A202279 002</u>	Nov 20, 2014
<u>AB</u>	ORBION PHARMS	<u>2.5MG</u>	<u>A203726 001</u>	Oct 21, 2019
<u>AB</u>		<u>5MG</u>	<u>A203726 002</u>	Oct 21, 2019
<u>AB</u>	PLD ACQUISITIONS LLC	<u>2.5MG</u>	<u>A207867 001</u>	Feb 27, 2017
<u>AB</u>		<u>5MG</u>	<u>A207867 002</u>	Feb 27, 2017
<u>AB</u>	ZYDUS PHARMS	<u>2.5MG</u>	<u>A203019 001</u>	Jul 11, 2018
<u>AB</u>		<u>5MG</u>	<u>A203019 002</u>	Jul 11, 2018

ZOMIG

<u>AB</u>	+ IPR	<u>2.5MG</u>	<u>N020768 001</u>	Nov 25, 1997
<u>AB</u>	+!	<u>5MG</u>	<u>N020768 002</u>	Nov 25, 1997

TABLET, ORALLY DISINTEGRATING; ORAL

ZOLMITRIPTAN

<u>AB</u>	ALEMBIC PHARMS LTD	<u>2.5MG</u>	<u>A205074 001</u>	Dec 01, 2016
<u>AB</u>		<u>5MG</u>	<u>A205074 002</u>	Dec 01, 2016
<u>AB</u>	GLENMARK GENERICS	<u>2.5MG</u>	<u>A202560 001</u>	May 14, 2013
<u>AB</u>		<u>5MG</u>	<u>A202560 002</u>	May 14, 2013
<u>AB</u>	JUBILANT GENERICS	<u>2.5MG</u>	<u>A202956 001</u>	Sep 17, 2015
<u>AB</u>		<u>5MG</u>	<u>A202956 002</u>	Sep 17, 2015
<u>AB</u>	ZYDUS PHARMS USA INC	<u>2.5MG</u>	<u>A202890 001</u>	May 15, 2013
<u>AB</u>		<u>5MG</u>	<u>A202890 002</u>	May 15, 2013

ZOMIG-ZMT

<u>AB</u>	+ ASTRAZENECA	<u>2.5MG</u>	<u>N021231 001</u>	Feb 13, 2001
<u>AB</u>	+!	<u>5MG</u>	<u>N021231 002</u>	Sep 17, 2001

ZOLPIDEM TARTRATE

SPRAY, METERED; ORAL

ZOLPIMIST

+! AYTU

5MG/SPRAY

N022196 001 Dec 19, 2008

TABLET; ORAL

AMBIEN

<u>AB</u>	+ SANOFI AVENTIS US	<u>5MG</u>	<u>N019908 001</u>	Dec 16, 1992
<u>AB</u>	+!	<u>10MG</u>	<u>N019908 002</u>	Dec 16, 1992

ZOLPIDEM TARTRATE

<u>AB</u>	ACME LABS	<u>5MG</u>	<u>A077214 001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A077214 002</u>	Apr 23, 2007
<u>AB</u>	APOTEX INC	<u>5MG</u>	<u>A077884 001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A077884 002</u>	Apr 23, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A078413 001</u>	May 04, 2007
<u>AB</u>		<u>10MG</u>	<u>A078413 002</u>	May 04, 2007
<u>AB</u>	CIPLA LTD	<u>5MG</u>	<u>A077388 001</u>	Jul 30, 2012
<u>AB</u>		<u>10MG</u>	<u>A077388 002</u>	Jul 30, 2012
<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>	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>	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

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

PRESCRIPTION DRUG PRODUCT LIST

ZOLPIDEM TARTRATE

TABLET;SUBLINGUAL

ZOLPIDEM TARTRATE

<u>AB</u>	DR REDDYS	<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>	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 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>	BRECKENRIDGE	<u>6.25MG</u>	<u>A213592 001</u>	Jun 04, 2020
<u>AB</u>		<u>12.5MG</u>	<u>A213592 002</u>	Jun 04, 2020
<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 PHARM	<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>	+	CONCORDIA	<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>	AUROBINDO PHARMA LTD	<u>25MG</u>	<u>A077645 002</u>	Sep 29, 2006
<u>AB</u>		<u>50MG</u>	<u>A077645 003</u>	Sep 29, 2006
<u>AB</u>		<u>100MG</u>	<u>A077645 001</u>	Dec 22, 2005
<u>AB</u>	BIONPHARMA INC	<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>	CADILA	<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
<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
<u>AB</u>		<u>100MG</u>	<u>A077651 003</u>	Jan 30, 2006
<u>AB</u>	GRANULES	<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>	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>	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>	UNICHEM	<u>25MG</u>	<u>A214492 001</u>	Jan 26, 2021
<u>AB</u>		<u>50MG</u>	<u>A214492 002</u>	Jan 26, 2021
<u>AB</u>		<u>100MG</u>	<u>A214492 003</u>	Jan 26, 2021

OTC DRUG PRODUCT LIST

ACETAMINOPHEN

SUPPOSITORY; RECTAL

ACEPHEN

COSETTE	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
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TABLET, EXTENDED RELEASE; ORAL

ACETAMINOPHEN

AUROBINDO PHARMA LTD	650MG	A207229 001	Nov 09, 2016
GRANULES	650MG	A211544 001	Apr 16, 2019
HERITAGE PHARMA	650MG	A207035 001	May 31, 2018
MARKSANS PHARMA	650MG	A215486 001	Aug 25, 2021
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

GRANULES	250MG; 250MG; 65MG	A214039 001	Feb 23, 2021
PERRIGO	250MG; 250MG; 65MG	A075794 001	Nov 26, 2001

EXCEDRIN (MIGRAINE)

+! GLAXOSMITHKLINE CONS	250MG; 250MG; 65MG	N020802 001	Jan 14, 1998
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ACETAMINOPHEN; IBUPROFEN

TABLET; ORAL

ADVIL DUAL ACTION WITH ACETAMINOPHEN

+! GLAXOSMITHKLINE	250MG; 125MG	N211733 001	Feb 28, 2020
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ADAPALENE

GEL; TOPICAL

ADAPALENE

GLENMARK GENERICS	0.1%	A091314 001	Jul 01, 2010
P AND L	0.1%	A090962 001	Jun 02, 2010

DIFFERIN

+! GALDERMA LABS LP	0.1%	N020380 002	Jul 08, 2016
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ALCAFTADINE

SOLUTION/DROPS; OPHTHALMIC

LASTACFT

+! ALLERGAN	0.25%	N022134 001	Jul 28, 2010
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ALCOHOL; CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL

AVAGARD

+! 3M	61%; 1%	N021074 001	Jun 07, 2001
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ALUMINUM HYDROXIDE; MAGNESIUM TRISILICATE

TABLET, CHEWABLE; ORAL

GAVISCON

+ CHATTEM SANOFI	80MG; 20MG	N018685 001	Dec 09, 1983
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ASPIRIN

CAPSULE; ORAL

VAZALORE

+ PLX PHARMA	81MG	N203697 002	Feb 26, 2021
+!	325MG	N203697 001	Jan 14, 2013

OTC DRUG PRODUCT LIST

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

AZELASTINE HYDROCHLORIDE

SPRAY, METERED;NASAL

ASTEPRO ALLERGY

+! BAYER HLTHCARE 0.2055MG/SPRAY N213872 001 Jun 17, 2021

CHILDREN'S ASTEPRO ALLERGY

+! BAYER HLTHCARE 0.2055MG/SPRAY N213872 002 Jun 17, 2021

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

+! J AND J CONSUMER 0.032MG/SPRAY N020746 003 Mar 23, 2015

INC

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 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

CATALENT 10MG A213105 001 Sep 21, 2020

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

AUROBINDO PHARMA 5MG/5ML A090750 002 Feb 02, 2010

HETERO LABS LTD III 5MG/5ML A210622 001 Mar 13, 2019

OTC DRUG PRODUCT LIST

CETIRIZINE HYDROCHLORIDE

SYRUP; ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

LANNETT CO INC	5MG/5ML	A091130	001	Apr 22, 2011
MICRO LABS	5MG/5ML	A091327	001	Oct 17, 2011
PERRIGO R AND D	5MG/5ML	A204226	001	Sep 09, 2013
	5MG/5ML	A090254	002	Apr 09, 2008
QUAGEN	5MG/5ML	A212266	001	May 16, 2019
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
AUROBINDO PHARMA	5MG/5ML	A090750	001	Feb 02, 2010
LANNETT CO INC	5MG/5ML	A091130	002	Apr 22, 2011
MICRO LABS	5MG/5ML	A091327	002	Oct 17, 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 INC	5MG/5ML	N022155	002	Nov 16, 2007
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CHILDREN'S ZYRTEC HIVES RELIEF

+! J AND J CONSUMER INC	5MG/5ML	N022155	001	Nov 16, 2007
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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
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	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
ORBION PHARMS	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 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
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
ORBION PHARMS	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

OTC DRUG PRODUCT LIST

CETIRIZINE HYDROCHLORIDE

TABLET;ORAL

CETIRIZINE HYDROCHLORIDE HIVES

SUN PHARM INDS LTD	5MG	A077498	003	Dec 27, 2007
	10MG	A077498	004	Dec 27, 2007
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

ZYRTEC ALLERGY

+! J AND J CONSUMER INC	10MG	N019835	004	Nov 16, 2007
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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 PHARM	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 PHARM	5MG	A090142	003	Aug 30, 2011
	10MG	A090142	004	Aug 30, 2011

TABLET, ORALLY DISINTEGRATING;ORAL

CETIRIZINE HYDROCHLORIDE

AUROBINDO PHARMA LTD	10MG	A213557	001	Sep 11, 2020
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CETIRIZINE HYDROCHLORIDE ALLERGY

PERRIGO R AND D	10MG	A205490	001	Sep 02, 2015
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ZYRTEC ALLERGY

+! J AND J CONSUMER INC	10MG	N022578	001	Sep 03, 2010
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CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

IVAX SUB TEVA PHARMS	5MG;120MG	A077170	001	Feb 25, 2008
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	5MG;120MG	N021150	002	Nov 09, 2007

CHLORHEXIDINE GLUCONATE

AEROSOL, METERED;TOPICAL

EXIDINE

+! XTTRIUM	4%	N019127	001	Dec 24, 1984
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CLOTH;TOPICAL

CHLORHEXIDINE GLUCONATE

+! SAGE PRODS	2%	N021669	001	Apr 25, 2005
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READYPREP CHG

MEDLINE INDUSTRIES	2%	N207964	001	Nov 20, 2018
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SOLUTION;TOPICAL

CHG SCRUB

ECOLAB	4%	N019258	002	Jul 22, 1986
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CHLORHEXIDINE GLUCONATE

BAJAJ	0.75%	N020111	001	Sep 11, 1997
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CIDA-STAT

ECOLAB	2%	N019258	001	Jul 22, 1986
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EXIDINE

+! XTTRIUM	2%	N019422	001	Dec 17, 1985
	4%	N019125	001	Dec 24, 1984

OTC DRUG PRODUCT LIST

CHLORHEXIDINE GLUCONATE

SOLUTION;TOPICAL

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

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

CHLORPHENIRAMINE MALEATE

! AVANTHI INC 12MG A040829 001 May 13, 2009

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PHENYLEPHRINE HYDROCHLORIDE

TABLET;ORAL

ADVIL ALLERGY AND CONGESTION RELIEF

+! GLAXOSMITHKLINE 4MG;200MG;10MG N022113 001 Dec 21, 2011

ADVIL MULTI-SYMPOM COLD & FLU

+! GLAXOSMITHKLINE 4MG;200MG;10MG N022113 002 Apr 28, 2017

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

SUSPENSION;ORAL

CHILDREN'S ADVIL ALLERGY SINUS

+! GLAXOSMITHKLINE 1MG/5ML;100MG/5ML;15MG/5ML N021587 001 Feb 24, 2004

TABLET;ORAL

ADVIL ALLERGY SINUS

+! GLAXOSMITHKLINE 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

OTC DRUG PRODUCT LIST

CLEMASTINE FUMARATE

TABLET;ORAL

CLEMASTINE FUMARATE

! L PERRIGO CO 1.34MG A074512 001 Nov 22, 1995

CLOTRIMAZOLE

CREAM;VAGINAL

CLOTRIMAZOLE

! P AND L 1% A074165 001 Jul 16, 1993

TARO 1% A072641 001 Dec 04, 1995

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

SUN PHARM 30MG;600MG A214781 001 Jul 01, 2021

60MG;1.2GM A214781 002 Jul 01, 2021

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 EQ 30MG HYDROBROMIDE/5ML A203133 001 Jul 28, 2017

TRIS PHARMA INC EQ 30MG HYDROBROMIDE/5ML A091135 001 May 25, 2012

DICLOFENAC SODIUM

GEL;TOPICAL

DICLOFENAC SODIUM

AMNEAL PHARMS 1% A208077 001 Mar 18, 2016

AUROLIFE PHARMA LLC 1% A204306 001 May 06, 2019

ENCUBE 1% A210986 001 Jan 27, 2020

PERRIGO PHARMA INTL 1% A211253 001 May 16, 2019

VOLTAREN ARTHRITIS PAIN

+! GLAXOSMITHKLINE 1% N022122 001 Oct 17, 2007

CONS

DIPHENHYDRAMINE CITRATE; IBUPROFEN

TABLET;ORAL

ADVIL PM

+! GLAXOSMITHKLINE 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

+! GLAXOSMITHKLINE 25MG;EQ 200MG FREE ACID AND POTASSIUM SALT N021393 001 Dec 21, 2005

IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE

AUROBINDO PHARMA 25MG;EQ 200MG FREE ACID AND POTASSIUM SALT A210676 001 Feb 14, 2019

LTD 25MG;EQ 200MG FREE ACID AND POTASSIUM SALT A090397 001 Nov 22, 2010

BIONPHARMA INC 25MG;EQ 200MG FREE ACID AND POTASSIUM SALT A200888 001 Mar 05, 2012

STRIDES PHARMA 25MG;EQ 200MG FREE ACID AND POTASSIUM SALT

OTC DRUG PRODUCT LIST

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

APOTEX 25MG; 220MG A211830 001 Aug 22, 2019

GRANULES 25MG; 220MG A213663 001 Sep 24, 2020

PERRIGO R AND D 25MG; 220MG A208499 001 May 10, 2019

DOCOSANOL

CREAM; TOPICAL

ABREVA

+! GLAXOSMITHKLINE 10% N020941 001 Jul 25, 2000

DOCOSANOL

P AND L 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

EPINEPHRINE

AEROSOL, METERED; INHALATION

PRIMATENE MIST

+! ARMSTRONG PHARMS 0.125MG/INH N205920 001 Nov 07, 2018

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

AMNEAL PHARMS NY EQ 20MG BASE A209716 001 Jun 05, 2019

AUROBINDO PHARMA EQ 20MG BASE A209339 001 Oct 16, 2017

LTD

DR REDDYS LABS LTD EQ 20MG BASE A207673 001 May 15, 2018

HETERO LABS LTD III EQ 20MG BASE A208939 001 May 28, 2020

EQ 20MG BASE A212507 001 Jun 02, 2020

PERRIGO R AND D EQ 20MG BASE A207193 001 Aug 18, 2017

SUN PHARM EQ 20MG BASE A212866 001 May 04, 2020

NEXIUM 24HR

+! ASTRAZENECA LP EQ 20MG BASE N204655 001 Mar 28, 2014

TABLET, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

DR REDDYS LABS LTD EQ 20MG BASE A211571 001 May 14, 2020

MYLAN EQ 20MG BASE A212088 001 Jun 25, 2020

NEXIUM 24HR

+! ASTRAZENECA LP EQ 20MG BASE N207920 001 Nov 23, 2015

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

+! DEXCEL PHARMA EQ 20MG BASE N214278 001 Oct 20, 2020

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

ANNORA PHARMA 10MG A215766 001 Nov 08, 2021

20MG A215766 002 Nov 08, 2021

ASCENT PHARMS INC 10MG A216030 001 Nov 03, 2021

20MG A216030 002 Nov 03, 2021

AUROBINDO PHARMA 10MG A206531 001 Apr 26, 2016

LTD

20MG A206531 002 Apr 26, 2016

DR REDDYS LABS LTD 10MG A077367 002 Aug 17, 2001

20MG A077367 001 Sep 25, 2006

P AND L 10MG A075512 001 Jul 26, 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

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

OTC DRUG PRODUCT LIST

FAMOTIDINE

TABLET;ORAL

PEPCID AC

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

+! CHATTEM SANOFI 30MG/5ML

N201373 001 Jan 24, 2011

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

P AND L 30MG/5ML

A203330 001 Nov 18, 2014

TARO 30MG/5ML

A208123 001 Nov 09, 2017

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

! P AND L 30MG/5ML

A203330 002 Nov 18, 2014

TARO 30MG/5ML

A208123 002 Nov 09, 2017

TABLET;ORAL

ALLEGRA ALLERGY

+ CHATTEM SANOFI 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

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

SUN PHARM INDS 30MG

A091567 001 Feb 06, 2012

TEVA 30MG

A076447 005 Apr 13, 2011

WOCKHARDT LTD 30MG

A079112 001 Feb 08, 2012

FEXOFENADINE HYDROCHLORIDE

L PERRIGO CO 60MG

A212971 001 Feb 24, 2020

180MG

A212971 002 Feb 24, 2020

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

GRANULES 60MG

A211075 001 Oct 18, 2019

180MG

A211075 002 Oct 18, 2019

HETERO LABS LTD V 60MG

A204097 002 Aug 19, 2016

180MG

A204097 003 Aug 19, 2016

RISING 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

RISING 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

OTC DRUG PRODUCT LIST

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 PHARM 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
CONSFLUTICASONE PROPIONATE

SPRAY, METERED;NASAL

FLONASE ALLERGY RELIEF

+ GLAXOSMITHKLINE 0.05MG/SPRAY N205434 001 Jul 23, 2014
CONS

FLUTICASONE PROPIONATE

AKORN 0.05MG/SPRAY A208024 001 Apr 17, 2019

APOTEX 0.05MG/SPRAY A208150 001 Feb 29, 2016

! HIKMA 0.05MG/SPRAY A207957 001 May 26, 2016

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

AUROBINDO PHARMA 600MG A210453 001 Oct 21, 2019

LTD

1.2GM A210453 002 Oct 21, 2019

GRANULES 600MG A213420 001 May 08, 2020

1.2GM A213420 002 May 08, 2020

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

1.2GM A078912 002 Nov 05, 2020

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

AUROBINDO PHARMA 600MG;60MG A213203 001 Mar 25, 2020

LTD

1.2GM;120MG A213203 002 Mar 25, 2020

DR REDDYS LABS LTD 600MG;60MG A208369 001 Dec 29, 2017

1.2GM;120MG A208369 002 Dec 29, 2017

SUN PHARM INDS INC 600MG;60MG A212542 001 Apr 28, 2020

1.2GM;120MG A212542 002 Apr 28, 2020

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

+! GLAXOSMITHKLINE EQ 200MG FREE ACID AND POTASSIUM SALT N020402 001 Apr 20, 1995

ADVIL MIGRAINE LIQUI-GELS

+! GLAXOSMITHKLINE 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 EQ 200MG FREE ACID AND POTASSIUM SALT A207753 001 Jun 29, 2018

OTC DRUG PRODUCT LIST

IBUPROFEN

CAPSULE;ORAL

IBUPROFEN

LTD

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

MIDOL LIQUID GELS

+! BIONPHARMA INC	200MG	N021472 001	Oct 18, 2002
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SUSPENSION;ORAL

CHILDREN'S ADVIL

GLAXOSMITHKLINE	100MG/5ML	N020589 001	Jun 27, 1996
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CHILDREN'S ADVIL-FLAVORED

GLAXOSMITHKLINE	100MG/5ML	N020589 002	Nov 07, 1997
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CHILDREN'S IBUPROFEN

PERRIGO	100MG/5ML	A074937 001	Dec 22, 1998
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CHILDREN'S MOTRIN

+! J AND J CONSUMER INC	100MG/5ML	N020516 001	Jun 16, 1995
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IBUPROFEN

APTAPHARMA INC	100MG/5ML	A210602 001	Nov 23, 2018
ARISE	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
P AND L	100MG/5ML	A074916 001	Apr 30, 1999
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
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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

INFANT'S ADVIL

+! GLAXOSMITHKLINE	50MG/1.25ML	N020812 002	Jan 12, 2000
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TABLET;ORAL

ADVIL

+! GLAXOSMITHKLINE	200MG	N018989 001	May 18, 1984
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IBUPROFEN

AMNEAL PHARMS	200MG	A079233 001	Mar 18, 2014
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	200MG	A202312 001	Oct 07, 2016
GRANULES INDIA	200MG	A079174 001	Dec 10, 2010
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
OHM	200MG	A071163 001	Jul 15, 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	A207095 001	May 05, 2017
	200MG	A207095 002	Aug 21, 2018
STRIDES PHARMA	200MG	A070481 001	Sep 24, 1986
	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

JUNIOR STRENGTH ADVIL

GLAXOSMITHKLINE	100MG	N020267 002	Dec 13, 1996
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OTC DRUG PRODUCT LIST

IBUPROFEN

TABLET; ORAL

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 GLAXOSMITHKLINE	50MG	N020944 001	Dec 18, 1998
IBUPROFEN ! PERRIGO	100MG	A076359 002	Jan 16, 2004
JUNIOR STRENGTH ADVIL GLAXOSMITHKLINE	100MG	N020944 002	Dec 18, 1998

IBUPROFEN SODIUM

TABLET; ORAL

ADVIL +! GLAXOSMITHKLINE	EQ 200MG BASE	N201803 001	Jun 12, 2012
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IBUPROFEN; PHENYLEPHRINE HYDROCHLORIDE

TABLET; ORAL

ADVIL CONGESTION RELIEF +! GLAXOSMITHKLINE	200MG; 10MG	N022565 001	May 27, 2010
IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE PERRIGO R AND D	200MG; 10MG	A203200 001	Jul 03, 2014

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE; ORAL

ADVIL COLD AND SINUS +! GLAXOSMITHKLINE	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 GLAXOSMITHKLINE	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 +! GLAXOSMITHKLINE	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

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

IVERMECTIN

LOTION; TOPICAL

IVERMECTIN TARO	0.5%	A210720 001	May 06, 2020
SKLICE +! ARBOR PHARMS LLC	0.5%	N202736 001	Feb 07, 2012

OTC DRUG PRODUCT LIST

KETOCONAZOLE

SHAMPOO; TOPICAL

NIZORAL ANTI-DANDRUFF

+! KRAMER 1% N020310 001 Oct 10, 1997

KETOTIFEN FUMARATE

SOLUTION/DROPS; OPHTHALMIC

ALAWAY

BAUSCH EQ 0.035% BASE A208158 001 Sep 24, 2020

+! 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

APOTEX INC EQ 0.025% BASE A077354 001 May 09, 2006

BAYSHORE PHARMS LLC EQ 0.025% BASE A204059 001 Jun 01, 2020

ZADITOR

! ALCON PHARMS LTD EQ 0.025% BASE A077200 001 Sep 02, 2008

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 15MG A203187 001 Jun 01, 2016

NATCO 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

+! PERRIGO PHARMA INTL 15MG N022327 001 May 18, 2009

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE; ORAL

LANSOPRAZOLE

DEXCEL PHARMA 15MG N208025 001 Jun 07, 2016

LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION; ORAL

XYZAL ALLERGY 24HR

+! SANOFI 2.5MG/5ML N209090 001 Jan 31, 2017

TABLET; ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

APOTEX 5MG A211443 001 Apr 21, 2021

DR REDDYS LABS LTD 5MG A210375 001 Jan 19, 2018

HETERO LABS LTD III 5MG A213513 001 Apr 29, 2020

MICRO LABS 5MG A211551 001 Nov 20, 2018

PERRIGO R AND D 5MG A211983 001 Mar 28, 2019

XYZAL ALLERGY 24HR

+! SANOFI 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

GLENMARK PHARMS LTD 1.5MG A207044 001 Mar 25, 2016

! L PERRIGO CO 0.75MG A090740 001 Dec 30, 2010

LABORATOIRE HRA 1.5MG A204044 001 Jul 03, 2018

MYLAN LABS LTD 1.5MG A202739 001 Oct 31, 2014

NAARI PTE LTD 1.5MG A202380 001 May 29, 2015

NOVEL LABS INC 1.5MG A202508 001 Feb 22, 2013

PERRIGO R AND D 1.5MG A202334 001 Aug 20, 2014

XIROMED 1.5MG A205329 001 Sep 18, 2018

OPCICON ONE-STEP

SUN PHARM 1.5MG A202635 001 Sep 11, 2014

PLAN B ONE-STEP

+! FDN CONSUMER 1.5MG N021998 001 Jul 10, 2009

OTC DRUG PRODUCT LIST

LOPERAMIDE HYDROCHLORIDE

CAPSULE;ORAL

LOPERAMIDE HYDROCHLORIDE

+ BIONPHARMA INC

1MG

N021855 001 Aug 04, 2005

+!

2MG

N021855 002 Aug 04, 2005

STRIDES PHARMA

2MG

A213070 001 Aug 11, 2021

SOLUTION;ORAL

IMODIUM A-D

+! J AND J CONSUMER

1MG/5ML

N019487 001 Mar 01, 1988

INC

LOPERAMIDE HYDROCHLORIDE

AKORN

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

1MG/7.5ML

N019487 002 Jul 08, 2004

INC

LOPERAMIDE HYDROCHLORIDE

PERRIGO R AND D

1MG/7.5ML

A091292 001 May 20, 2011

TABLET;ORAL

IMODIUM A-D

+! J AND J CONSUMER

2MG

N019860 001 Nov 22, 1989

INC

LOPERAMIDE HYDROCHLORIDE

AUROBINDO PHARMA

2MG

A206548 001 Dec 15, 2015

LTD

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

2MG;125MG

N021140 001 Nov 30, 2000

INC

LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE

AUROBINDO PHARMA

2MG;125MG

A211059 001 Dec 14, 2020

LTD

BIONPHARMA INC

2MG;125MG

A213484 001 Sep 10, 2021

GUARDIAN DRUG

2MG;125MG

A214541 001 May 27, 2021

HETERO LABS LTD V

2MG;125MG

A211438 001 Jun 17, 2021

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

10MG

N021952 001 Jun 16, 2008

LLC

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

1MG/ML

N020641 002 Nov 27, 2002

LLC

LORATADINE

AUROBINDO PHARMA

1MG/ML

A208931 001 Jun 29, 2018

LTD

HETERO LABS LTD III

1MG/ML

A210409 001 May 07, 2021

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

WOCKHARDT BIO AG

1MG/ML

A201865 001 Jul 31, 2015

WOCKHARDT BIO AG

1MG/ML

A075815 001 Aug 20, 2004

TABLET;ORAL

CLARITIN

+! BAYER HEALTHCARE

10MG

N019658 002 Nov 27, 2002

OTC DRUG PRODUCT LIST

LORATADINE

TABLET; ORAL

CLARITIN

LLC

CLARITIN HIVES RELIEF

+! BAYER HEALTHCARE 10MG

N019658 003 Nov 19, 2003

LLC

LORATADINE

APOTEX INC 10MG

A076471 001 Feb 14, 2006

APPCO 10MG

A207569 001 Mar 12, 2019

AUROBINDO PHARMA LTD

A208314 001 Apr 16, 2018

GRANULES 10MG

A210722 001 Dec 23, 2019

MYLAN 10MG

A076154 001 Aug 20, 2003

PERRIGO 10MG

A076301 001 Jun 25, 2004

PLD ACQUISITIONS LLC

A075209 001 Jan 21, 2003

SUN PHARM INDS LTD 10MG

A076134 001 Aug 18, 2003

UNIQUE PHARM 10MG

A214684 001 Jan 07, 2021

TABLET, CHEWABLE; ORAL

CHILDREN'S CLARITIN

+! BAYER HEALTHCARE 5MG

N021891 001 Aug 23, 2006

LLC

CLARITIN

+ BAYER HEALTHCARE 10MG

N021891 002 Nov 21, 2018

LLC

LORATADINE

PERRIGO PHARMA INTL 5MG

A210033 001 Jun 12, 2019

SUN PHARM 5MG

A210088 001 Apr 16, 2018

TABLET, ORALLY DISINTEGRATING; ORAL

ALAVERT

FDN CONSUMER 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

AUROBINDO PHARMA LTD

A208477 001 Apr 11, 2018

PERRIGO PHARMA INTL 10MG

A076011 001 Sep 29, 2003

RUBICON 10MG

A214280 001 Sep 10, 2020

TENSHI 5MG

A212795 001 Sep 18, 2020

10MG

A213294 001 Oct 30, 2020

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

P AND L 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

OTC DRUG PRODUCT LIST

MICONAZOLE NITRATE

CREAM;TOPICAL, VAGINAL

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

P AND L 2% A074164 001 Mar 29, 1996

MICONAZOLE NITRATE

COSETTE 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, INSERT;TOPICAL, VAGINAL

MICONAZOLE NITRATE

PERRIGO R AND D 2%,1.2GM A079114 001 Jun 02, 2010

MONISTAT 1 COMBINATION PACK

+! MEDTECH PRODUCTS 2%,1.2GM N021308 001 Jun 29, 2001

CREAM, SUPPOSITORY;TOPICAL, VAGINAL

M-ZOLE 3 COMBINATION PACK

P AND L 2%,200MG A074926 001 Apr 16, 1999

MICONAZOLE NITRATE COMBINATION PACK

L PERRIGO CO 2%,200MG A075329 001 Apr 20, 1999

MONISTAT 3 COMBINATION PACK

+! MEDTECH PRODUCTS 2%,200MG N020670 002 Apr 16, 1996

MONISTAT 7 COMBINATION PACK

+! MEDTECH PRODUCTS 2%,100MG N020288 002 Apr 26, 1993

SUPPOSITORY;VAGINAL

MICONAZOLE NITRATE

COSETTE 100MG A074414 001 Apr 30, 1997

P AND L 100MG A073507 001 Nov 19, 1993

! 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 5% A091344 001 Apr 28, 2011

MINOXIDIL (FOR MEN)

P AND L 5% A208092 001 Feb 17, 2017

TARO 5% A209074 001 Dec 31, 2018

MINOXIDIL (FOR WOMEN)

P AND L 5% A208092 002 Jul 27, 2017

TARO 5% A209074 002 Apr 22, 2019

WOMEN'S ROGAINE

+! JOHNSON AND JOHNSON 5% N021812 002 Feb 28, 2014

SOLUTION;TOPICAL

MINOXIDIL (FOR MEN)

AKORN 2% A074731 001 Dec 24, 1996

L PERRIGO CO 2% A075357 001 Jul 30, 1999

P AND L 2% A074588 001 Apr 05, 1996

WOCKHARDT BIO AG 2% A074767 001 Feb 28, 1997

MINOXIDIL (FOR WOMEN)

AKORN 2% A074731 002 May 11, 2005

L PERRIGO CO 2% A075357 002 Jul 30, 1999

MINOXIDIL EXTRA STRENGTH (FOR MEN)

P AND L 5% A075518 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

OTC DRUG PRODUCT LIST

MINOXIDIL

SOLUTION;TOPICAL

ROGAINE EXTRA STRENGTH (FOR MEN)

+!	JOHNSON AND JOHNSON	5%	N020834	001	Nov 14, 1997
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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	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
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OPCON-A

+!	BAUSCH AND LOMB	0.02675%;0.315%	N020065	001	Jun 08, 1994
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VISINE

+!	JOHNSON AND JOHNSON	0.025%;0.3%	N020485	001	Jan 31, 1996
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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	220MG	N020204	002	Jan 11, 1994
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NAPROXEN SODIUM

	AMNEAL PHARMS NY	220MG	A079096	001	Dec 16, 2008
	AUROBINDO PHARMA LTD	220MG	A205497	001	Mar 18, 2016
	CONTRACT PHARMACAL	220MG	A074635	001	Jan 13, 1997
	DR REDDYS LABS INC	220MG	A075168	001	Jul 28, 1998
	GRANULES INDIA	220MG	A091353	001	Sep 20, 2011
	LNK INTL INC	220MG	A204872	001	Jan 23, 2017
	MARKSANS PHARMA	220MG	A090545	001	Mar 16, 2011
	NOVELGENIX THERAPS	220MG	A207612	001	Nov 16, 2018
	PERRIGO	220MG	A074661	001	Jan 13, 1997
	SUN PHARM INDS LTD	220MG	A091183	001	May 20, 2011

NAPROXEN SODIUM; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ALEVE-D SINUS & COLD

+!	BAYER	220MG;120MG	N021076	001	Nov 29, 1999
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NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE

	DR REDDYS LABS INC	220MG;120MG	A077381	001	Sep 27, 2006
	PERRIGO	220MG;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

+	CHATTEM SANOFI	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

OTC DRUG PRODUCT LIST

NICOTINE POLACRILEX

GUM, CHEWING;BUCCAL

NICOTINE POLACRILEX

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
P AND L	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	A079044	001 Jul 08, 2009
	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
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
	EQ 4MG BASE	A091354	001 Jul 20, 2011
	EQ 4MG BASE	A206393	001 Dec 15, 2016

TROCHE/LOZENGE;ORAL

NICORETTE

+ GLAXOSMITHKLINE

+!

+ GLAXOSMITHKLINE

CONS

+!

NICOTINE POLACRILEX

AUROBINDO PHARMA
LTD

DR REDDYS LABS SA

P AND L

PERRIGO R AND D

PLD ACQUISITIONS

EQ 2MG BASE	N022360	001	May 18, 2009
EQ 4MG BASE	N022360	002	May 18, 2009
EQ 2MG BASE	N021330	001	Oct 31, 2002
EQ 4MG BASE	N021330	002	Oct 31, 2002
EQ 2MG BASE	A213266	001	Aug 03, 2021
EQ 4MG BASE	A213266	002	Aug 03, 2021
EQ 2MG BASE	A212796	001	Jan 08, 2020
EQ 2MG BASE	A212983	001	Feb 21, 2020
EQ 2MG BASE	A213233	001	Aug 04, 2020
EQ 4MG BASE	A212796	002	Jan 08, 2020
EQ 4MG BASE	A212983	002	Feb 21, 2020
EQ 4MG BASE	A213233	002	Aug 04, 2020
EQ 2MG BASE	A208875	001	Oct 31, 2019
EQ 2MG BASE	A209206	001	Jun 26, 2018
EQ 2MG BASE	A209519	001	Jul 02, 2018
EQ 2MG BASE	A209520	001	Oct 31, 2019
EQ 2MG BASE	A210711	001	Oct 31, 2019
EQ 2MG BASE	A210712	001	Sep 06, 2019
EQ 2MG BASE	A212056	001	Jul 26, 2019
EQ 2MG BASE	A212057	001	May 14, 2020
EQ 4MG BASE	A208875	002	Oct 31, 2019
EQ 4MG BASE	A209206	002	Jun 26, 2018
EQ 4MG BASE	A209519	002	Jul 02, 2018
EQ 4MG BASE	A209520	002	Oct 31, 2019
EQ 4MG BASE	A210711	002	Oct 31, 2019
EQ 4MG BASE	A210712	002	Sep 06, 2019
EQ 4MG BASE	A212056	002	Jul 26, 2019
EQ 4MG BASE	A212057	002	May 14, 2020
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
EQ 2MG BASE	A207868	001	Feb 07, 2019

OTC DRUG PRODUCT LIST

NICOTINE POLACRILEX

TROCHE/LOZENGE;ORAL

NICOTINE POLACRILEX

EQ 4MG BASE

A207868 002 Feb 07, 2019

NIZATIDINE

TABLET;ORAL

AXID AR

+! GLAXOSMITHKLINE

75MG

N020555 001 May 09, 1996

OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

OLOPATADINE HYDROCHLORIDE

AKORN

EQ 0.1% BASE

A204532 001 Jan 10, 2017

EQ 0.2% BASE

A204723 001 Dec 05, 2017

ALEMBIC PHARMS LTD

EQ 0.1% BASE

A209919 001 Dec 07, 2018

EQ 0.2% BASE

A209420 001 Apr 29, 2019

APOTEX

EQ 0.1% BASE

A078350 001 Dec 07, 2015

APOTEX INC

EQ 0.2% BASE

A090918 001 Dec 05, 2017

BARR LABS INC

EQ 0.2% BASE

A090848 001 Jul 13, 2015

EUGIA PHARMA

EQ 0.1% BASE

A204812 001 Dec 18, 2015

EQ 0.2% BASE

A209995 001 Apr 04, 2019

GLAND PHARMA LTD

EQ 0.1% BASE

A209619 001 Aug 02, 2019

EQ 0.2% BASE

A209752 001 May 20, 2020

MYLAN

EQ 0.2% BASE

A204620 001 Jun 16, 2020

USV

EQ 0.1% BASE

A203152 001 Dec 07, 2015

PATADAY ONCE DAILY RELIEF

+! ALCON LABS INC

EQ 0.2% BASE

N021545 001 Dec 22, 2004

+!

EQ 0.7% BASE

N206276 001 Jan 30, 2015

PATADAY TWICE DAILY RELIEF

+! ALCON LABS INC

EQ 0.1% BASE

N020688 001 Dec 18, 1996

OMEPRAZOLE

TABLET, DELAYED RELEASE;ORAL

OMEPRAZOLE

APOTEX

20MG

A210070 001 Feb 11, 2019

+! DEXCEL PHARMA

20MG

N022032 001 Dec 04, 2007

DR REDDYS

20MG

A207740 001 Nov 05, 2018

SUN PHARM

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

HETERO LABS LTD III

EQ 20MG BASE

A211732 001 Mar 25, 2020

P AND L

EQ 20MG BASE

A206582 001 Jun 01, 2020

PERRIGO R AND D

EQ 20MG BASE

A204152 001 Jul 30, 2015

PRILOSEC OTC

+! ASTRAZENECA

EQ 20MG BASE

N021229 001 Jun 20, 2003

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

AUROLIFE PHARMA LLC

20MG;1.1GM

A204923 001 Nov 07, 2016

PERRIGO R AND D

20MG;1.1GM

A201361 001 Jul 15, 2016

ZYDUS PHARMS

20MG;1.1GM

A203345 001 Mar 16, 2018

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	3.9MG/24HR	N202211	001	Jan 25, 2013
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OXYMETAZOLINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

VISINE L.R.

+	!	JOHNSON AND JOHNSON	0.025%	N019407	001	Mar 31, 1989
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PERMETHRIN

LOTION;TOPICAL

NIX

+	!	MEDTECH PRODUCTS	1%	N019918	001	May 02, 1990
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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

MIRALAX

+	!	BAYER HEALTHCARE	17GM/SCOOPFUL	N022015	001	Oct 06, 2006
		LLC				

POLYETHYLENE GLYCOL 3350

ANI PHARMS

			17GM/SCOOPFUL	A202850	001	Dec 15, 2015
--	--	--	---------------	---------	-----	--------------

ANNORA PHARMA

			17GM/SCOOPFUL	A214990	001	Apr 14, 2021
--	--	--	---------------	---------	-----	--------------

AUROBINDO PHARMA

			17GM/SCOOPFUL	A209017	001	Apr 09, 2018
--	--	--	---------------	---------	-----	--------------

LTD

LGM PHARMA

			17GM/SCOOPFUL	A090812	001	Oct 07, 2009
--	--	--	---------------	---------	-----	--------------

MYLAN

			17GM/PACKET	A078915	001	Oct 06, 2009
--	--	--	-------------	---------	-----	--------------

			17GM/SCOOPFUL	A078915	002	Oct 06, 2009
--	--	--	---------------	---------	-----	--------------

NOVEL LABS INC

			17GM/SCOOPFUL	A091077	001	Oct 06, 2009
--	--	--	---------------	---------	-----	--------------

NOVELGENIX THERAPS

			17GM/SCOOPFUL	A202071	001	Dec 28, 2012
--	--	--	---------------	---------	-----	--------------

NUVO PHARMS INC

			17GM/SCOOPFUL	A206105	001	Oct 28, 2016
--	--	--	---------------	---------	-----	--------------

PERRIGO R AND D

			17GM/PACKET	A090685	001	Oct 06, 2009
--	--	--	-------------	---------	-----	--------------

			17GM/SCOOPFUL	A090685	002	Oct 06, 2009
--	--	--	---------------	---------	-----	--------------

STRIDES PHARMA

			17GM/SCOOPFUL	A079214	001	Jan 31, 2013
--	--	--	---------------	---------	-----	--------------

			17GM/SCOOPFUL	A203928	001	Aug 24, 2016
--	--	--	---------------	---------	-----	--------------

			17GM/PACKET	A203928	002	Aug 24, 2016
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POTASSIUM IODIDE

SOLUTION;ORAL

POTASSIUM IODIDE

!	!	MISSION PHARMACAL	65MG/ML	A206211	001	Mar 24, 2016
		CO				

TABLET;ORAL

IOSAT

+		ANBEX	65MG	N018664	002	May 12, 2011
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+	!		130MG	N018664	001	Oct 14, 1982
---	---	--	-------	---------	-----	--------------

THYROSAFE

!	!	BTG INTL	65MG	A076350	001	Sep 10, 2002
---	---	----------	------	---------	-----	--------------

POVIDONE-IODINE

SOLUTION;TOPICAL

POVIDONE IODINE

+	!	ALLEGIANCE HLTHCARE	1%	N019522	001	Mar 31, 1989
---	---	---------------------	----	---------	-----	--------------

SPONGE;TOPICAL

E-Z SCRUB 201

+	!	BECTON DICKINSON	20%	N019240	001	Nov 29, 1985
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E-Z SCRUB 241

+	!	BECTON DICKINSON	10%	N019476	001	Jan 07, 1987
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PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

PSEUDOEPHEDRINE HYDROCHLORIDE

AUROBINDO PHARMA

			120MG	A209008	001	Jun 09, 2017
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LTD

L PERRIGO CO

			120MG	A075153	001	Feb 26, 1999
--	--	--	-------	---------	-----	--------------

SUN PHARM INDS LTD

			120MG	A077442	001	Sep 28, 2005
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OTC DRUG PRODUCT LIST

PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

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

DR REDDYS LABS LTD

EQ 75MG BASE

A075294 001 Mar 28, 2000

EQ 150MG BASE

A078192 001 Aug 31, 2007

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

UNIQUE PHARM LABS

EQ 75MG BASE

A210250 001 Aug 30, 2019

EQ 150MG BASE

A210228 001 Aug 30, 2019

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

TIOCONAZOLE

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

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**

0.9% SODIUM CHLORIDE INJECTION USP SOLUTION

INJECTABLE; INJECTION

NONE

FRESENIUS KABI AG

N125695

Sep 05, 2019

ANTICOAGULANT 4% SODIUM CITRATE SOLUTION USP

INJECTABLE; INJECTION

NONE

HAEMONETICS
MANUFACTURING INC

N760305

Jun 30, 1978

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

ANTICOAGULANT CITRATE DEXTROSE SOLUTION USP

INJECTABLE; INJECTION

NONE

FENWAL INC

N160918

Mar 17, 1978

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

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE-1 SOLUTION

INJECTABLE; INJECTION

NONE

HAEMONETICS

N800077

Nov 06, 1980

MANUFACTURING INC

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

MACOPHARMA

N040083

Nov 21, 2005

NONE

TERUMO BCT INC

A070025

Jan 06, 2009

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP

INJECTABLE; INJECTION

NONE

FENWAL INC

N170401

Dec 06, 1977

N811012

Jun 28, 1983

PALL CORP

N800222

Aug 23, 1982

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

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

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; 0.718GM/100ML;0.017GM/100ML; 0.396GM/100ML;0.588GM/100ML	N820915	Sep 22, 1983
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ANTICOAGULANT SODIUM CITRATE 4% SOLUTION

INJECTABLE; INJECTION

NONE

N980123 Mar 03, 2000

HAEMONETICS
CORPORATION
LABORATORIES
GRIFOIS, S.A.

A125697 Oct 25, 2019

TERUMO BCT

A125608 Jun 26, 2018

ANTICOAGULANT SODIUM CITRATE SOLUTION

INJECTABLE; INJECTION

TRICITRASOL

CITRA LABS, LLC

N010409 Jul 10, 2003

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

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

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 PHARMACEUTICALS USA INC	6GM/100ML;0.9GM/100ML	A740592 Nov 12, 1998

ISOPLATE SOLUTION IN THE 500 ML EXCEL CONTAINERSTORAGE/PROCESSING SOLUTION ONLY; SHOULD NEVER BE
INFUSED DIRECTLY TO THE PATIENT.

ISOPLATE SOLUTION		
TERUMO BCT		N90067 Mar 05, 2013

LEUKOCYTE REDUCTION FILTRATION SYSTEM FOR WHOLE BLOOD WITH CPD ANTICOAGULANT AND SOLX ADDITIVE

INJECTABLE; INJECTION

LEUKOSEP HWB-600-XL		
HEMERUS MEDICAL, LLC		N110059 Apr 25, 2013

RED BLOOD CELL PROCESSING SOLUTIONSTORAGE/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, MONOHYDRATESTORAGE/PROCESSING SOLUTION ONLY; SHOULD NEVER BE
INFUSED DIRECTLY TO THE PATIENT.

INTERSOL SOLUTION		
FRESENIUS KABI USA, LLC	2.26GM/500ML;2.21GM/500ML;1.59GM/500ML;1.53GM/500ML;0.465GM/500ML	N080041 Dec 09, 2009

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ABACAIVIR SULFATE

TABLET;ORAL

ABACAIVIR SULFATE

APOTEX INC

EQ 300MG BASE

A201570 001 Dec 17, 2012

ABACAIVIR SULFATE; LAMIVUDINE

TABLET;ORAL

ABACAIVIR SULFATE AND LAMIVUDINE

ZYDUS PHARMS

EQ 600MG BASE;300MG

A208990 001 Nov 15, 2018

ABAMETAPIR

LOTION;TOPICAL

XEGLYZE

+ DR REDDYS LABS SA

0.74%

N206966 001 Jul 24, 2020

ABARELIX

INJECTABLE;INTRAMUSCULAR

PLENAXIS

SPECIALITY EUROPEAN

100MG/VIAL

N021320 001 Nov 25, 2003

ACAMPROSATE CALCIUM

TABLET, DELAYED RELEASE;ORAL

ACAMPROSATE CALCIUM

BARR LABS DIV TEVA

333MG

A200143 001 Nov 18, 2013

CAMPRAL

+ FOREST LABS

333MG **

N021431 001 Jul 29, 2004

ACARBOSE

TABLET;ORAL

ACARBOSE

HANGZHOU ZHONGMEI

25MG

A213821 001 Aug 18, 2020

50MG

A213821 002 Aug 18, 2020

100MG

A213821 003 Aug 18, 2020

MYLAN

25MG

A091053 001 Jan 06, 2011

50MG

A091053 002 Jan 06, 2011

100MG

A091053 003 Jan 06, 2011

PRECOSE

+ BAYER HLTHCARE

25MG **

N020482 004 May 29, 1997

+

50MG **

N020482 001 Sep 06, 1995

+

100MG **

N020482 002 Sep 06, 1995

ACEBUTOLOL HYDROCHLORIDE

CAPSULE;ORAL

ACEBUTOLOL HYDROCHLORIDE

ANI PHARMS

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

POWDER;INTRAVENOUS

ACETAMINOPHEN

+ RISING PHARMA

1GM/VIAL

N206610 001 Jan 15, 2021

SUPPOSITORY;RECTAL

ACEPHEN

COSETTE

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

NEOPAP

POLYMEDICA

120MG

N016401 001

TYLENOL

J AND J CONSUMER INC

120MG

N017756 002

650MG

N017756 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN

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

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 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

GRANULES 325MG;50MG;40MG

A040864 001 Dec 01, 2008

HIKMA 500MG;50MG;40MG

A040336 001 Aug 18, 1999

HIKMA PHARMS 325MG;50MG;40MG

A089718 001 Jun 12, 1995

MIKART 750MG;50MG;40MG

A040496 001 Dec 23, 2003

MIRROR PHARMS LLC 500MG;50MG;40MG

A040883 001 Dec 23, 2008

SPECGX LLC 325MG;50MG;40MG

A087804 001 Jan 24, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

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
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ESGIC-PLUS

MIKART	500MG;50MG;40MG	A089451	001	May 23, 1988
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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

ACETAMINOPHEN; CLEMASTINE FUMARATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET; ORAL

TAVIST ALLERGY/SINUS/HEADACHE

NOVARTIS	500MG;EQ 0.25MG BASE;30MG	N021082	001	Mar 01, 2001
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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				
ACTAVIS MID ATLANTIC	120MG/5ML;12MG/5ML	A086366	001	
DAVA PHARMS INC	120MG/5ML;12MG/5ML	A085861	001	
LANNETT CO INC	120MG/5ML;12MG/5ML	A040098	001	Sep 20, 1996
WOCKHARDT BIO AG	120MG/5ML;12MG/5ML	A091238	001	Nov 10, 2011
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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; CODEINE PHOSPHATE

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	
	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	
STRIDES PHARMA	300MG;15MG	A089990	001	Sep 30, 1988
	300MG;30MG	A089805	001	Sep 30, 1988
	300MG;60MG	A089828	001	Sep 30, 1988
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

	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	
TYLENOL W/ CODEINE NO. 3				
+ JANSSEN PHARMS	300MG;30MG	A085055	003	
TYLENOL W/ CODEINE NO. 4				
+ JANSSEN PHARMS	300MG;60MG	A085055	004	

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

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
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CO-GESIC

UCB INC	500MG;5MG	A087757	001	May 03, 1982
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DURADYNE DHC

FOREST PHARMS	500MG;5MG	A087809	001	Mar 17, 1983
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HY-PHEN

ASCHER	500MG;5MG	A087677	001	May 03, 1982
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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
ACTAVIS LABS FL INC	300MG;5MG	A206470	001	Jun 02, 2016
	300MG;7.5MG	A206470	002	Jun 02, 2016
	300MG;10MG	A206470	003	Jun 02, 2016
ALVOGEN	300MG;5MG	A208540	001	Nov 08, 2018
	300MG;7.5MG	A208540	002	Nov 08, 2018
	300MG;10MG	A208540	003	Nov 08, 2018
	325MG;2.5MG	A209958	001	Oct 24, 2018
	325MG;5MG	A209958	002	Oct 24, 2018
	325MG;7.5MG	A209958	003	Oct 24, 2018
	325MG;10MG	A209958	004	Oct 24, 2018
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
	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
CEROVENE INC	325MG;5MG	A211690	001	Feb 07, 2020
	325MG;7.5MG	A211690	002	Feb 07, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

	325MG;10MG	A211690	003	Feb 07, 2020
GRANULES	325MG;5MG	A211729	001	Jan 03, 2020
	325MG;7.5MG	A211729	002	Jan 03, 2020
	325MG;10MG	A211729	003	Jan 03, 2020
HALSEY	500MG;5MG	A089554	001	Jun 12, 1987
IVAX PHARMS	500MG;5MG	A089696	001	Apr 21, 1988
LANNETT CO INC	300MG;5MG	A207171	001	Jun 20, 2017
	300MG;7.5MG	A207171	002	Jun 20, 2017
	300MG;10MG	A207171	003	Jun 20, 2017
	325MG;5MG	A207172	001	Jun 22, 2017
	325MG;7.5MG	A207172	002	Jun 22, 2017
	325MG;10MG	A207172	003	Jun 22, 2017
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
NOSTRUM LABS INC	325MG;2.5MG	A209924	001	Nov 16, 2018
	325MG;5MG	A209924	002	Nov 16, 2018
	325MG;7.5MG	A209924	003	Nov 16, 2018
	325MG;10MG	A209924	004	Nov 16, 2018
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
STRIDES PHARMA	300MG;5MG	A205001	001	Jul 05, 2016
	300MG;7.5MG	A205001	002	Jul 05, 2016
	300MG;10MG	A205001	003	Jul 05, 2016
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
UPSHER SMITH LABS	325MG;5MG	A206484	001	Mar 24, 2017
	325MG;7.5MG	A206484	002	Mar 24, 2017
	325MG;10MG	A206484	003	Mar 24, 2017
USL PHARMA	500MG;5MG	A089290	001	May 29, 1987
	500MG;5MG	A089291	001	May 29, 1987
VINTAGE PHARMS	300MG;5MG	A090415	001	Jan 24, 2011
	300MG;7.5MG	A090415	002	Jan 24, 2011
	300MG;10MG	A090415	003	Jan 24, 2011
	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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

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
	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
NORCO			
APIL	325MG; 2.5MG	A040148 004	Jul 07, 2014
	325MG; 5MG	A040099 001	Jun 25, 1997
	325MG; 5MG	A040148 005	Jul 07, 2014
	325MG; 7.5MG	A040148 003	Sep 12, 2000
	325MG; 10MG	A040148 001	Feb 14, 1997
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

SOLUTION; ORAL

OXYCODONE HYDROCHLORIDE AND ACETAMINOPHEN

VINTAGE PHARMS 325MG/5ML; 5MG/5ML A203573 001 Dec 18, 2014

ROXICET

HIKMA 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

CEROVENE INC 325MG; 5MG A207574 001 Dec 13, 2016

DURAMED PHARMS BARR 325MG; 5MG A040272 001 Jun 30, 1998

GRANULES 325MG; 2.5MG A211708 001 Oct 31, 2019

325MG; 5MG A211708 002 Oct 31, 2019

325MG; 7.5MG A211708 003 Oct 31, 2019

325MG; 10MG A211708 004 Oct 31, 2019

LANNETT CO INC 325MG; 5MG A207333 001 Sep 25, 2017

325MG; 10MG A207333 002 Sep 25, 2017

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

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

NOSTRUM LABS INC 325MG; 5MG A209385 001 Jul 02, 2018

325MG; 7.5MG A209385 002 Jul 02, 2018

325MG; 10MG A209385 003 Jul 02, 2018

VINTAGE PHARMS 325MG; 2.5MG A090733 001 Jul 11, 2013

325MG; 5MG A040105 001 Jul 30, 1996

325MG; 7.5MG A090734 001 Jul 11, 2013

325MG; 10MG A090734 002 Jul 11, 2013

WATSON LABS 325MG; 5MG A040171 001 Oct 30, 1997

325MG; 7.5MG A040535 001 Sep 05, 2003

325MG; 10MG A040535 002 Sep 05, 2003

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

+ HIKMA 325MG; 5MG A087003 001

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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	
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DOLENE AP-65

LEDERLE	650MG;65MG	A085100	001	
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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	
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ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET; ORAL

DARVOCET A500

XANODYNE PHARM	500MG;100MG	A076429	001	Sep 10, 2003
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DARVOCET-N 100

XANODYNE PHARM	650MG;100MG	N017122	002	
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DARVOCET-N 50

XANODYNE PHARM	325MG;50MG	N017122	001	
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PROPACET 100

TEVA	650MG;100MG	A070107	001	Jun 12, 1985
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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
	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

GRAVITI PHARMS	325MG;37.5MG	A076914	001	Jul 26, 2006
MACLEODS PHARMS LTD	325MG;37.5MG	A206885	001	May 02, 2017
NOSTRUM LABS INC	325MG;37.5MG	A078778	001	Apr 07, 2014
STRIDES PHARMA	325MG;37.5MG	A076475	001	Apr 21, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAZOLAMIDE

CAPSULE, EXTENDED RELEASE;ORAL

ACETAZOLAMIDE

RISING 500MG A203917 001 Jun 18, 2019

DIAMOX

+ TEVA BRANDED PHARM 500MG ** N012945 001

TABLET;ORAL

ACETAZOLAMIDE

ALRA 250MG A083320 001

ASCOT 250MG A087686 001 Oct 20, 1982

HERITAGE PHARMA 250MG A088882 001 Oct 22, 1985

SUN PHARM INDUSTRIES 125MG A089753 002 Jun 22, 1988

250MG A089753 001 Jun 22, 1988

VANGARD 250MG A087654 001 Feb 05, 1982

WATSON LABS 250MG A084498 002

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

PAR STERILE PRODUCTS EQ 500MG BASE/VIAL A205358 001 Jun 20, 2017

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

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS;OTIC

ACETASOL HC

ACTAVIS MID ATLANTIC 2%;1% A087143 001 Jan 13, 1982

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETOHEXAMIDE

TABLET; ORAL

ACETOHEXAMIDE

ANI PHARMS	250MG	A070870 002	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	
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ACETRIZOATE SODIUM

SOLUTION; INTRAUTERINE

SALPIX

ORTHO MCNEIL PHARM	53%	N009008 001	
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ACETYLCHOLINE CHLORIDE

FOR SOLUTION; OPHTHALMIC

MIOCHOL

+ NOVARTIS	20MG/VIAL **	N016211 001	
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ACETYLCYSTEINE

INJECTABLE; INTRAVENOUS

ACETYLCYSTEINE

RISING	6GM/30ML (200MG/ML)	A203624 001	Jun 19, 2015
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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	
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+	20% **	N013601 001	
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MUCOSIL-10

DEY	10%	A070575 001	Oct 14, 1986
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MUCOSIL-20

DEY	20%	A070576 001	Oct 14, 1986
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TABLET, EFFERVESCENT; ORAL

CETYLEV

+ ARBOR PHARMS LLC	500MG	N207916 001	Jan 29, 2016
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+	2.5GM	N207916 002	Jan 29, 2016
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ACETYLCYSTEINE; ISOPROTERENOL HYDROCHLORIDE

SOLUTION; INHALATION

MUCOMYST W/ ISOPROTERENOL

MEAD JOHNSON	10%; 0.05%	N017366 001	
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ACETYLDIGITOXIN

TABLET; ORAL

ACYLANID

NOVARTIS	0.1MG	N009436 001	
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ACITRETIN

CAPSULE; ORAL

ACITRETIN

MYLAN	17.5MG	A203707 001	Sep 10, 2015
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	22.5MG	A203707 002	Sep 10, 2015
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SORIATANE

+ STIEFEL LABS INC	10MG	N019821 001	Oct 28, 1996
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+	17.5MG	N019821 003	Aug 06, 2009
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+	22.5MG	N019821 004	Aug 06, 2009
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+	25MG	N019821 002	Oct 28, 1996
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACRISORCIN

CREAM;TOPICAL

AKRINOL

SCHERING

2MG/GM

N012470 001

ACRIVASTINE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE;ORAL

SEMPREX-D

+

AUXILIUM PHARMS LLC

8MG;60MG

N019806 001 Mar 25, 1994

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

MYLAN

200MG

A074977 001 Apr 13, 1998

RANBAXY

200MG

A074975 001 Sep 30, 1998

ROXANE

200MG

A074570 002 Apr 22, 1997

STRIDES PHARMA

200MG

A074833 001 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

ZOVIRAX

+

MYLAN

200MG **

N018828 001 Jan 25, 1985

OINTMENT;OPHTHALMIC

AVACLYR

+

FERA PHARMS LLC

3% **

N202408 001 Mar 29, 2019

OINTMENT;TOPICAL

ACYCLOVIR

ANDA REPOSITORY

5%

A206437 001 Jul 31, 2017

PADAGIS ISRAEL

5%

A205659 001 Feb 20, 2019

TABLET;ORAL

ACYCLOVIR

ACTAVIS ELIZABETH

400MG

A074870 001 Jun 05, 1997

ACTAVIS ELIZABETH

800MG

A074870 002 Jun 05, 1997

CHARTWELL MOLECULES

400MG

A074834 001 Apr 24, 1997

CHARTWELL MOLECULES

800MG

A074834 002 Apr 24, 1997

IVAX SUB TEVA PHARMS

400MG

A074836 001 Apr 22, 1997

IVAX SUB TEVA PHARMS

800MG

A074836 002 Apr 22, 1997

LEK PHARM

400MG

A074658 001 Apr 22, 1997

LEK PHARM

800MG

A074658 002 Apr 22, 1997

MYLAN

400MG

A074976 001 Apr 13, 1998

MYLAN

400MG

A075211 001 Sep 28, 1998

MYLAN

800MG

A074976 002 Apr 13, 1998

MYLAN

800MG

A075211 002 Sep 28, 1998

SUN PHARM INDS LTD

400MG

A074980 001 Sep 30, 1998

SUN PHARM INDS LTD

800MG

A074980 002 Sep 30, 1998

TEVA

200MG **

A074556 001 Apr 22, 1997

TEVA PHARMS

400MG

A075021 001 Mar 18, 1998

TEVA PHARMS

800MG

A075021 002 Mar 18, 1998

ZOVIRAX

+

MYLAN

400MG **

N020089 001 Apr 30, 1991

+

MYLAN

800MG **

N020089 002 Apr 30, 1991

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

EUROHLTH INTL SARL

EQ 1GM BASE/VIAL

A074885 002 Dec 19, 1997

ACYCLOVIR SODIUM

APOTHECON

EQ 500MG BASE/VIAL

A074897 001 Feb 27, 1998

APOTHECON

EQ 1GM BASE/VIAL

A074897 002 Feb 27, 1998

ATHENEX INC

EQ 500MG BASE/VIAL

A074596 002 Apr 22, 1997

ATHENEX INC

EQ 1GM BASE/VIAL

A074596 001 Apr 22, 1997

EUROHLTH INTL SARL

EQ 500MG BASE/VIAL

A074913 001 Oct 15, 1997

EUROHLTH INTL SARL

EQ 1GM BASE/VIAL

A074913 002 Oct 15, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR SODIUM

FRESENIUS KABI USA	EQ 500MG BASE/VIAL	A075015 001	Apr 30, 1998
HIKMA	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
	EQ 500MG BASE/VIAL	A074969 001	Aug 26, 1997
	EQ 1GM BASE/VIAL	A074969 002	Aug 26, 1997
ZYDUS PHARMS	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

GEL; TOPICAL

ADAPALENE

ENCUBE ETHICALS	0.3%	A200298 001	Jun 14, 2012
SOLUTION; TOPICAL			
DIFFERIN			
+ GALDERMA LABS LP	0.1% **	N020338 001	May 31, 1996

ADAPALENE; BENZOYL PEROXIDE

GEL; TOPICAL

ADAPALENE AND BENZOYL PEROXIDE

ENCUBE	0.1%; 2.5%	A206164 001	May 23, 2018
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ADENOSINE

INJECTABLE; INJECTION

ADENOCARD

+ ASTELLAS	3MG/ML **	N019937 002	Oct 30, 1989
ADENOSINE			
AM REGENT	3MG/ML	A090010 001	Apr 28, 2009
HIKMA	3MG/ML	A076501 001	Jun 16, 2004
MYLAN LABS LTD	3MG/ML	A078640 001	Mar 21, 2014
TEVA PHARMS USA	3MG/ML	A076564 001	Jun 16, 2004
	3MG/ML	A078676 001	Jul 31, 2008
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; ORAL

ALBENDAZOLE

LUPIN LTD	200MG	A211636 001	Jun 10, 2020
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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 effectiveness 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

AEROSOL, METERED; INHALATION

ALBUTEROL SULFATE

PADAGIS US

EQ 0.09MG BASE/INH

A203760 001 Feb 24, 2020

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

AKORN

EQ 0.083% BASE

A075063 001 Feb 09, 1999

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

EQ 0.5% BASE

A075050 001 Jun 18, 1998

COPLEY PHARM

EQ 0.083% BASE

A073495 001 May 28, 1993

EQ 0.5% BASE

A073307 001 Nov 27, 1991

LANDELA PHARM

EQ 0.083% BASE

A077569 001 Apr 04, 2006

LUOXIN AUROVITAS

EQ 0.021% BASE

A211888 001 Apr 20, 2020

EQ 0.042% BASE

A211888 002 Apr 20, 2020

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

CHARTWELL RX

EQ 2MG BASE/5ML

A077788 001 Jun 26, 2007

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

AUROBINDO PHARMA LTD

EQ 2MG BASE

A213657 001 May 14, 2020

EQ 4MG BASE

A213657 002 May 14, 2020

CHARTWELL RX

EQ 2MG BASE

A072151 001 Dec 05, 1989

EQ 4MG BASE

A072152 001 Dec 05, 1989

COPLEY PHARM

EQ 2MG BASE

A072966 001 Nov 22, 1991

EQ 4MG BASE

A072967 001 Nov 22, 1991

EYWA

EQ 2MG BASE

A213524 001 Oct 08, 2020

EQ 4MG BASE

A213524 002 Oct 08, 2020

PLIVA

EQ 2MG BASE

A072316 001 Dec 05, 1989

EQ 4MG BASE

A072317 001 Dec 05, 1989

STRIDES PHARMA

EQ 2MG BASE

A072860 002 Dec 20, 1989

EQ 4MG BASE

A072860 001 Dec 20, 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALBUTEROL SULFATE

TABLET;ORAL

ALBUTEROL SULFATE

	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
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

ALBUTEROL SULFATE

RISING PHARMA	EQ 4MG BASE	A078092 002	Jan 29, 2007
	EQ 8MG BASE	A078092 001	Jan 29, 2007
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			
STRIDES PHARMA	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
LUOXIN AUROVITAS	EQ 0.083% BASE;0.017%	A206532 001	Jul 08, 2020
TEVA PHARMS	EQ 0.083% BASE;0.017%	A076724 001	Dec 31, 2007
WATSON LABS TEVA	EQ 0.083% BASE;0.017%	A077063 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

ALENDRONATE SODIUM

SOLUTION;ORAL

FOSAMAX

+ MERCK EQ 70MG BASE/75ML ** N021575 001 Sep 17, 2003

TABLET;ORAL

ALENDRONATE SODIUM

CHARTWELL RX	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
IMPAX LABS INC	EQ 5MG BASE	A075710 001	Feb 06, 2008
	EQ 10MG BASE	A075710 002	Feb 06, 2008
	EQ 35MG BASE	A075710 003	Feb 06, 2008
	EQ 40MG BASE	A075710 004	Feb 06, 2008
	EQ 70MG BASE	A075710 005	Feb 06, 2008
JUBILANT CADISTA	EQ 5MG BASE	A090557 001	Feb 18, 2010
	EQ 10MG BASE	A090557 002	Feb 18, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALENDRONATE SODIUM

TABLET;ORAL

ALENDRONATE SODIUM

	EQ 35MG BASE	A090557 003	Feb 18, 2010
	EQ 70MG BASE	A090557 004	Feb 18, 2010
MYLAN	EQ 35MG BASE	A078638 001	Aug 04, 2008
	EQ 70MG BASE	A078638 002	Aug 04, 2008
PICKET PHARMS	EQ 5MG BASE	A079049 003	Aug 04, 2008
	EQ 10MG BASE	A079049 004	Aug 04, 2008
	EQ 35MG BASE	A079049 001	Aug 04, 2008
	EQ 70MG BASE	A079049 002	Aug 04, 2008
RISING PHARMA	EQ 5MG BASE	A076584 001	Aug 04, 2008
	EQ 10MG BASE	A076584 002	Aug 04, 2008
	EQ 35MG BASE	A076584 003	Aug 04, 2008
	EQ 70MG BASE	A076584 004	Aug 04, 2008
TEVA PHARMS	EQ 35MG BASE	A076184 002	Aug 04, 2008
	EQ 70MG BASE	A076184 001	Feb 06, 2008
FOSAMAX			
+	MERCK AND CO INC	EQ 5MG BASE **	N020560 003
+		EQ 10MG BASE **	N020560 001
+		EQ 35MG BASE **	N020560 004
+		EQ 40MG BASE **	N020560 002

ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ALFUZOSIN HYDROCHLORIDE

HERITAGE PHARMA	10MG	A079056 001	Jul 18, 2011
MYLAN	10MG	A079014 001	Jul 18, 2011
TORRENT PHARMS	10MG	A079054 001	Jul 18, 2011
WOCKHARDT LTD	10MG	A090221 001	Aug 10, 2012

ALISKIREN HEMIFUMARATE

CAPSULE, PELLETT;ORAL

TEKTURNA

+	NODEN PHARMA	EQ 37.5MG BASE	N210709 001	Nov 14, 2017
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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; 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

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	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
SUN PHARM INDS INC	100MG	A078390 001	Aug 30, 2007
	300MG	A078390 002	Aug 30, 2007
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

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

AXERT

+ JANSSEN PHARMS	EQ 6.25MG BASE **	N021001 001	May 07, 2001
+	EQ 12.5MG BASE **	N021001 002	May 07, 2001

ALOSETRON HYDROCHLORIDE

TABLET; ORAL

ALOSETRON HYDROCHLORIDE

EYWA PHARMA	EQ 0.5MG BASE	A211621 001	Sep 16, 2019
	EQ 1MG BASE	A211621 002	Sep 16, 2019

ALPRAZOLAM

SOLUTION; ORAL

ALPRAZOLAM

ROXANE	0.5MG/5ML	A074314 001	Oct 31, 1993
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TABLET; ORAL

ALPRAZOLAM

ANI PHARMS	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	0.25MG	A074215 001	Jan 27, 1994
	0.5MG	A074215 002	Jan 27, 1994
	1MG	A074215 003	Jan 27, 1994
	2MG	A074215 004	Jan 27, 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
OXFORD PHARMS	0.25MG	A078491 001	Sep 25, 2008
	0.5MG	A078491 002	Sep 25, 2008
	1MG	A078491 003	Sep 25, 2008
	2MG	A078491 004	Dec 12, 2008
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALPRAZOLAMTABLET; ORAL
ALPRAZOLAM

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	0.5MG	A077725 001	Jul 31, 2006
	0.5MG	A077979 001	Feb 28, 2007
	1MG	A077725 002	Jul 31, 2006
	1MG	A077979 002	Feb 28, 2007
	2MG	A077725 004	Jul 31, 2006
	2MG	A077979 003	Feb 28, 2007
	3MG	A077725 003	Jul 31, 2006
	3MG	A077979 004	Feb 28, 2007
HERITAGE PHARMS INC	0.5MG	A078489 001	Oct 17, 2008
	1MG	A078489 002	Oct 17, 2008
	2MG	A078489 003	Oct 17, 2008
	3MG	A078489 004	Oct 17, 2008
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

ALPROSTADIL

INJECTABLE; INJECTION

CAVERJECT

+	PFIZER	0.005MG/VIAL	N020379 003	Jun 27, 1996
		0.005MG/ML	N020755 001	Oct 31, 1997
		0.01MG/ML	N020755 002	Oct 01, 1997
		0.02MG/ML	N020755 003	Oct 01, 1997

EDEX

AUXILIUM PHARMS LLC	0.005MG/VIAL	N020649 001	Jun 12, 1997
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ALSEROXYLON

TABLET; ORAL

RAUTENSIN

NOVARTIS	2MG	N009215 001
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RAUWILOID

3M	2MG	N008867 001
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALTRETAMINE

CAPSULE;ORAL

HEXALEN

+ EISAI INC 50MG N019926 001 Dec 26, 1990

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

+ CHATTEM SANOFI 160MG;40MG N018685 002 Dec 09, 1983

AMANTADINE HYDROCHLORIDE

CAPSULE;ORAL

AMANTADINE HYDROCHLORIDE

ACTAVIS ELIZABETH 100MG A077659 001 Feb 23, 2006

INVAGEN PHARMS 100MG A207570 001 Sep 30, 2016

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

GENUS 50MG/5ML A076352 001 Sep 10, 2004

TEVA PHARMS 50MG/5ML A073115 001 Aug 23, 1991

VINTAGE 50MG/5ML A077992 001 Dec 12, 2006

WOCKHARDT BIO AG 50MG/5ML A075060 001 Dec 24, 1998

SYMMETREL

+ ENDO PHARMS 50MG/5ML ** N016023 002

TABLET;ORAL

AMANTADINE HYDROCHLORIDE

INVAGEN PHARMS 100MG A207571 001 Jan 31, 2017

SYMMETREL

+ ENDO PHARMS 100MG ** N018101 001

TABLET, EXTENDED RELEASE;ORAL

OSMOLEX ER

+ ADAMAS PHARMA EQ 161MG BASE N209410 004 Apr 22, 2020

+ EQ 258MG BASE N209410 003 Feb 16, 2018

AMBENONIUM CHLORIDE

TABLET;ORAL

MYTELASE

SANOFI AVENTIS US 10MG N010155 002

AMCINONIDE

CREAM;TOPICAL

CYCLOCORT

+ ASTELLAS 0.025% N018116 001

+ 0.1% N018116 002

LOTION;TOPICAL

CYCLOCORT

+ ASTELLAS 0.1% N019729 001 Jun 13, 1988

OINTMENT;TOPICAL

CYCLOCORT

+ ASTELLAS 0.1% N018498 001

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMIFOSTINE

INJECTABLE; INJECTION

AMIFOSTINE

AUROMEDICS PHARMA 500MG/VIAL A204363 001 Jul 17, 2017

ETHYOL

CLINIGEN 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

HIKMA

EQ 50MG BASE/ML A063274 001 May 18, 1992

EQ 250MG BASE/ML A063275 001 May 18, 1992

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

EQ 250MG BASE/ML A064099 001 Jun 20, 1995

IGI LABS INC

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

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

CHARTWELL RX EQ 5MG ANHYDROUS; 50MG A073357 001 Nov 27, 1991

TEVA EQ 5MG ANHYDROUS; 50MG A070795 001 Apr 17, 1988

WATSON LABS EQ 5MG ANHYDROUS; 50MG A073334 001 Jul 19, 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%

ICU MEDICAL INC 10% (10GM/100ML) N017673 003

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

AMINOSYN 8.5%

ICU MEDICAL INC 8.5% (8.5GM/100ML) N017673 004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

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 effectiveness 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
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AMINOSYN 3.5% W/ DEXTROSE 5% IN PLASTIC CONTAINER

ABBOTT	3.5%;5GM/100ML	N019120 001	Oct 11, 1984
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AMINOSYN 4.25% W/ DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT	4.25%;25GM/100ML	N019119 001	Oct 11, 1984
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AMINOSYN II 3.5% IN DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT	3.5%;25GM/100ML	N019505 002	Nov 07, 1986
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	3.5%;25GM/100ML	N019713 006	Sep 09, 1988
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HOSPIRA	3.5%;25GM/100ML	N019681 001	Nov 01, 1988
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AMINOSYN II 3.5% IN DEXTROSE 5% IN PLASTIC CONTAINER

ABBOTT	3.5%;5GM/100ML	N019506 001	Nov 07, 1986
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	3.5%;5GM/100ML	N019713 002	Sep 09, 1988
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HOSPIRA	3.5%;5GM/100ML	N019681 002	Nov 01, 1988
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AMINOSYN II 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER

ABBOTT	4.25%;10GM/100ML	N019713 001	Sep 09, 1988
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HOSPIRA	4.25%;10GM/100ML	N019681 004	Nov 01, 1988
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AMINOSYN II 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER

ABBOTT	4.25%;20GM/100ML	N019713 004	Sep 09, 1988
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HOSPIRA	4.25%;20GM/100ML	N019681 005	Nov 01, 1988
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AMINOSYN II 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT	4.25%;25GM/100ML	N019504 002	Nov 07, 1986
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	4.25%;25GM/100ML	N019713 005	Sep 09, 1988
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HOSPIRA	4.25%;25GM/100ML	N019681 003	Nov 01, 1988
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AMINOSYN II 5% IN DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT	5%;25GM/100ML	N019565 001	Dec 17, 1986
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	5%;25GM/100ML	N019713 003	Sep 09, 1988
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HOSPIRA	5%;25GM/100ML	N019681 006	Nov 01, 1988
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TRAVASOL 2.75% IN DEXTROSE 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE	2.75%;10GM/100ML	N019520 002	Sep 23, 1988
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TRAVASOL 2.75% IN DEXTROSE 15% IN PLASTIC CONTAINER

BAXTER HLTHCARE	2.75%;15GM/100ML	N019520 003	Sep 23, 1988
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TRAVASOL 2.75% IN DEXTROSE 20% IN PLASTIC CONTAINER

BAXTER HLTHCARE	2.75%;20GM/100ML	N019520 004	Sep 23, 1988
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TRAVASOL 2.75% IN DEXTROSE 25% IN PLASTIC CONTAINER

BAXTER HLTHCARE	2.75%;25GM/100ML	N019520 005	Sep 23, 1988
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TRAVASOL 2.75% IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE	2.75%;5GM/100ML	N019520 001	Sep 23, 1988
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TRAVASOL 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE	4.25%;10GM/100ML	N019520 007	Sep 23, 1988
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TRAVASOL 4.25% IN DEXTROSE 15% IN PLASTIC CONTAINER

BAXTER HLTHCARE	4.25%;15GM/100ML	N019520 008	Sep 23, 1988
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TRAVASOL 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER

BAXTER HLTHCARE	4.25%;20GM/100ML	N019520 009	Sep 23, 1988
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TRAVASOL 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER

BAXTER HLTHCARE	4.25%;25GM/100ML	N019520 010	Sep 23, 1988
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TRAVASOL 4.25% IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE	4.25%;5GM/100ML	N019520 006	Sep 23, 1988
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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/100ML;22.4MG/100ML;104.5MG/100ML;205MG/100ML	N019712 002	Sep 08, 1988
HOSPIRA INC			
	4.25%;10GM/100ML;51MG/100ML;176.5MG/100ML;22.4MG/100ML;104.5MG/100ML;205MG/100ML	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/100ML;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;120MG/100ML;49.3MG/100ML	N019564 001	Dec 16, 1986
	3.5%;5GM/100ML;30MG/100ML;97MG/100ML;120MG/100ML;49.3MG/100ML	N019712 001	Sep 08, 1988
HOSPIRA INC			
	3.5%;5GM/100ML;30MG/100ML;97MG/100ML;120MG/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;120MG/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

ICU MEDICAL INC	3.5%; 21MG/100ML; 40MG/100ML; 128MG/100ML; 234MG/100ML	N017789 003	
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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	
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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
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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	
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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
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AMINOSYN II 7% W/ ELECTROLYTES

ICU MEDICAL INC	7%; 102MG/100ML; 45MG/100ML; 522MG/100ML; 4 10MG/100ML	N019437 006	Apr 03, 1986
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AMINOSYN II 8.5% W/ ELECTROLYTES

ICU MEDICAL INC	8.5%; 102MG/100ML; 45MG/100ML; 522MG/100ML ; 410MG/100ML	N019437 005	Apr 03, 1986
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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
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AMINOSYN II 8.5% W/ELECTROLYTES

ICU MEDICAL INC	8.5%; 102MG/100ML; 492MG/100ML; 60MG/100ML ; 425MG/100ML	N019437 008	Oct 25, 2002
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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
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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
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TRAVASOL 3.5% W/ ELECTROLYTES

BAXTER HLTHCARE	3.5%; 51MG/100ML; 131MG/100ML; 218MG/100ML ; 35MG/100ML	N017493 003	
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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
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TRAVASOL 5.5% W/ ELECTROLYTES

BAXTER HLTHCARE	5.5%; 102MG/100ML; 522MG/100ML; 431MG/100M L; 224MG/100ML	N017493 001	
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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
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TRAVASOL 8.5% W/ ELECTROLYTES

BAXTER HLTHCARE	8.5%; 102MG/100ML; 522MG/100ML;	N017493 002	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

INJECTABLE; INJECTION

TRAVASOL 8.5% W/ ELECTROLYTES

594MG/100ML; 154MG/100ML

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

+ AKORN 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

SOLUTION; ORAL

AMINOCAPROIC ACID

AKORN 0.25GM/ML A074759 001 Sep 02, 1998

TABLET; ORAL

AMINOCAPROIC

AKORN 500MG A075602 001 May 24, 2001

AMINOCAPROIC ACID

EDENBRIDGE PHARMS 500MG A212110 001 Jun 14, 2021

AMINOGLUTETHIMIDE

TABLET; ORAL

CYTADREN

NOVARTIS 250MG N018202 001

AMINOHIPPURATE SODIUM

INJECTABLE; INJECTION

AMINOHIPPURATE SODIUM

MERCK 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

AM REGENT 25MG/ML A087600 001

ELKINS SINN 25MG/ML A087239 001

HOSPIRA 25MG/ML A087601 001 Jul 23, 1982

INTL MEDICATION 25MG/ML A087209 001 Feb 01, 1982

25MG/ML A087867 001 Nov 10, 1983

25MG/ML A087868 001 Nov 10, 1983

KING PHARMS 25MG/ML A086606 001

LUITPOLD 25MG/ML A087240 001

LYPHOMED 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINOPHYLLINE

INJECTABLE; INJECTION

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
ROXANE	105MG/5ML	A088126 001	Aug 19, 1983

AMINOPHYLLINE DYE FREE

ACTAVIS MID ATLANTIC	105MG/5ML	A087727 001	Apr 16, 1982
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SOMOPHYLLIN

FISONS	105MG/5ML	A086466 001	
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SOMOPHYLLIN-DF

FISONS	105MG/5ML	A087045 001	
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SUPPOSITORY; RECTAL

TRUPHYLLINE

COSETTE	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	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	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINOSALICYLATE SODIUM

POWDER; ORAL

P.A.S. SODIUM

CENTURY PHARMS 4GM/PACKET A080947 001

SODIUM AMINOSALICYLATE

HEXCEL 100% A080097 001

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

DR REDDYS 50MG/ML A076163 001 Sep 05, 2003

HOSPIRA 50MG/ML A075955 001 Oct 18, 2002

50MG/ML A076108 001 Oct 15, 2002

HOSPIRA INC 50MG/ML A203885 001 Nov 25, 2013

INTL MEDICATION SYS 50MG/ML N021594 001 Feb 04, 2004

PAR STERILE PRODUCTS 50MG/ML A076394 001 Apr 25, 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIL

WARNER CHILCOTT	10MG	A083939	001	
	25MG	A083937	001	
	50MG	A083938	002	
	75MG	A084957	001	
	100MG	A085093	001	
	150MG	A086295	001	
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	10MG	A085031	002	
	25MG	A085031	001	
	50MG	A085031	003	
	75MG	A085031	004	
AUROBINDO PHARMA USA	10MG	A086009	002	
	25MG	A086009	003	
	50MG	A086009	001	
	75MG	A086009	004	
	100MG	A086009	005	
	150MG	A086009	006	
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

AMITRIPTYLINE HYDROCHLORIDE

	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	
	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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

ELAVIL

+

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

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

HERITAGE PHARMA

EQ 12.5MG BASE;5MG

A072052 001 Dec 16, 1988

EQ 25MG BASE;10MG

A072053 001 Dec 16, 1988

STRIDES PHARMA

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

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

HERITAGE PHARMA

10MG;2MG

A073007 001 Oct 17, 1991

10MG;4MG

A073009 001 Oct 17, 1991

25MG;2MG

A073008 001 Oct 17, 1991

25MG;4MG

A073010 001 Oct 17, 1991

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;4MG

A072539 001 Feb 15, 1989

10MG;4MG

A070375 001 Aug 25, 1986

10MG;4MG

A072540 001 Feb 15, 1989

25MG;2MG

A070374 001 Aug 25, 1986

25MG;2MG

A072541 001 Feb 15, 1989

25MG;4MG

A070376 001 Aug 25, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET; ORAL

PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE

	25MG; 4MG	A072134	001	Feb 15, 1989
	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	
TRIAVIL 4-50				
NEW RIVER	50MG; 4MG **	N014715	006	

AMLEXANOX

PASTE; DENTAL

APHTHASOL

ULURU	5%	N020511	001	Dec 17, 1996
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PATCH; TOPICAL

AMLEXANOX

ULURU	2MG	N021727	001	Sep 29, 2004
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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
HIKMA	EQ 2.5MG BASE	A077262	001	Jul 09, 2007
	EQ 5MG BASE	A077262	002	Jul 09, 2007
	EQ 10MG BASE	A077262	003	Jul 09, 2007
HIKMA PHARMS	EQ 2.5MG BASE	A077771	001	Apr 12, 2011
	EQ 5MG BASE	A077771	002	Apr 12, 2011
	EQ 10MG BASE	A077771	003	Apr 12, 2011
MACLEODS PHARMS LTD	EQ 5MG BASE	A201380	001	Apr 13, 2012
	EQ 10MG BASE	A201380	002	Apr 13, 2012
MYLAN	EQ 2.5MG BASE	A076418	001	Oct 03, 2005
	EQ 2.5MG BASE	A078224	001	Feb 27, 2008
	EQ 5MG BASE	A076418	002	Oct 03, 2005
	EQ 5MG BASE	A078224	002	Feb 27, 2008
	EQ 10MG BASE	A076418	003	Oct 03, 2005
	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 INDS INC	EQ 2.5MG BASE	A078231	001	Nov 30, 2007
	EQ 5MG BASE	A078231	002	Nov 30, 2007
	EQ 10MG BASE	A078231	003	Nov 30, 2007
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	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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMLODIPINE BESYLATE

TABLET;ORAL

AMLODIPINE BESYLATE

TEVA	EQ 2.5MG BASE	A076846 001	Jun 28, 2007
	EQ 5MG BASE	A076846 002	Jun 28, 2007
	EQ 10MG BASE	A076846 003	Jun 28, 2007
TORRENT PHARMS	EQ 2.5MG BASE	A078573 001	Sep 22, 2008
	EQ 5MG BASE	A078573 002	Sep 22, 2008
	EQ 10MG BASE	A078573 003	Sep 22, 2008
UPSHER SMITH LABS	EQ 2.5MG BASE	A077759 001	Jul 09, 2007
	EQ 5MG BASE	A077759 002	Jul 09, 2007
	EQ 10MG BASE	A077759 003	Jul 09, 2007
WATSON LABS	EQ 2.5MG BASE	A077671 001	Jul 19, 2007
	EQ 5MG BASE	A077671 002	Jul 19, 2007
	EQ 10MG BASE	A077671 003	Jul 19, 2007
WOCKHARDT	EQ 2.5MG BASE	A078500 001	Sep 06, 2007
	EQ 5MG BASE	A078500 002	Sep 06, 2007
	EQ 10MG BASE	A078500 003	Sep 06, 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 BESYLATE; ATORVASTATIN CALCIUM

TABLET;ORAL

AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM

MYLAN	EQ 2.5MG BASE;EQ 10MG BASE	A200465 001	Nov 29, 2013
	EQ 2.5MG BASE;EQ 20MG BASE	A200465 002	Nov 29, 2013
	EQ 2.5MG BASE;EQ 40MG BASE	A200465 003	Nov 29, 2013

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE;ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

CIPLA	EQ 2.5MG BASE;10MG	A077215 001	Dec 07, 2018
	EQ 5MG BASE;10MG	A077215 002	Dec 07, 2018
	EQ 5MG BASE;20MG	A077215 003	Dec 07, 2018
	EQ 10MG BASE;20MG	A077215 004	Dec 07, 2018
MYLAN	EQ 2.5MG BASE;10MG	A077375 001	May 21, 2010
	EQ 5MG BASE;10MG	A077375 002	May 21, 2010
	EQ 5MG BASE;20MG	A077375 003	May 21, 2010
	EQ 5MG BASE;40MG	A079047 001	Jul 05, 2011
	EQ 10MG BASE;20MG	A077375 004	May 21, 2010
	EQ 10MG BASE;40MG	A079047 002	Jul 05, 2011
STRIDES PHARMA	EQ 2.5MG BASE;10MG	A078381 001	Jul 29, 2010
	EQ 5MG BASE;10MG	A078381 002	Jul 29, 2010
	EQ 5MG BASE;20MG	A078381 003	Jul 29, 2010
	EQ 5MG BASE;40MG	A078381 005	Jul 29, 2010
	EQ 10MG BASE;20MG	A078381 004	Jul 29, 2010
	EQ 10MG BASE;40MG	A078381 006	Jul 29, 2010
TEVA PHARMS	EQ 2.5MG BASE;10MG	A077179 001	May 18, 2007
	EQ 5MG BASE;10MG	A077179 002	May 18, 2007
	EQ 5MG BASE;20MG	A077179 003	May 18, 2007
	EQ 5MG BASE;40MG	A077179 005	Jul 05, 2011
	EQ 10MG BASE;20MG	A077179 004	May 18, 2007
	EQ 10MG BASE;40MG	A077179 006	Jul 05, 2011

AMLODIPINE BESYLATE; CELECOXIB

TABLET;ORAL

CONSENSI

+	PURPLE BIOTECH	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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET;ORAL

AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE

STRIDES PHARMA	EQ 5MG BASE;12.5MG;160MG	A201087 001	Jun 01, 2015
	EQ 5MG BASE;25MG;160MG	A201087 002	Jun 01, 2015
	EQ 10MG BASE;12.5MG;160MG	A201087 003	Jun 01, 2015
	EQ 10MG BASE;25MG;160MG	A201087 004	Jun 01, 2015
TEVA PHARMS	EQ 10MG BASE;25MG;320MG	A201087 005	Jun 01, 2015
	EQ 5MG BASE;12.5MG;160MG	A200435 001	Sep 25, 2012
	EQ 5MG BASE;25MG;160MG	A200435 002	Sep 25, 2012
	EQ 10MG BASE;12.5MG;160MG	A200435 005	Sep 25, 2012
	EQ 10MG BASE;25MG;160MG	A200435 003	Sep 25, 2012
TORRENT	EQ 10MG BASE;25MG;320MG	A200435 004	Sep 25, 2012
	EQ 5MG BASE;12.5MG;160MG	A201593 001	Jun 03, 2015
	EQ 5MG BASE;25MG;160MG	A201593 002	Jun 03, 2015
	EQ 10MG BASE;12.5MG;160MG	A201593 003	Jun 03, 2015
	EQ 10MG BASE;25MG;160MG	A201593 004	Jun 03, 2015
	EQ 10MG BASE;25MG;320MG	A201593 005	Jun 03, 2015

AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL

TABLET;ORAL

AMLODIPINE AND OLMESARTAN MEDOXOMIL

ACCORD HLTHCARE INC	EQ 5MG BASE;20MG	A209600 001	Aug 30, 2018
	EQ 5MG BASE;40MG	A209600 003	Aug 30, 2018
	EQ 10MG BASE;20MG	A209600 002	Aug 30, 2018
	EQ 10MG BASE;40MG	A209600 004	Aug 30, 2018
JUBILANT GENERICS	EQ 5MG BASE;20MG	A207450 001	May 15, 2017
	EQ 5MG BASE;40MG	A207450 002	May 15, 2017
	EQ 10MG BASE;20MG	A207450 003	May 15, 2017
	EQ 10MG BASE;40MG	A207450 004	May 15, 2017
SCIEGEN PHARMS INC	EQ 5MG BASE;20MG	A209010 001	Dec 03, 2018
	EQ 5MG BASE;40MG	A209010 002	Dec 03, 2018
	EQ 10MG BASE;20MG	A209010 003	Dec 03, 2018
	EQ 10MG BASE;40MG	A209010 004	Dec 03, 2018
TEVA PHARMS USA	EQ 5MG BASE;20MG	A091154 001	Oct 26, 2016
	EQ 5MG BASE;40MG	A091154 002	Oct 26, 2016
	EQ 10MG BASE;20MG	A091154 003	Oct 26, 2016
	EQ 10MG BASE;40MG	A091154 004	Oct 26, 2016

AMLODIPINE BESYLATE; TELMISARTAN

TABLET;ORAL

TELMISARTAN AND AMLODIPINE

ALEMBIC PHARMS LTD	EQ 5MG BASE;40MG	A205234 001	Nov 17, 2016	
	EQ 5MG BASE;80MG	A205234 003	Nov 17, 2016	
	EQ 10MG BASE;40MG	A205234 002	Nov 17, 2016	
	EQ 10MG BASE;80MG	A205234 004	Nov 17, 2016	
TWINSTA				
	+ BOEHRINGER INGELHEIM	EQ 5MG BASE;40MG	N022401 001	Oct 16, 2009
	+	EQ 5MG BASE;80MG	N022401 003	Oct 16, 2009
	+	EQ 10MG BASE;40MG	N022401 002	Oct 16, 2009
+	EQ 10MG BASE;80MG	N022401 004	Oct 16, 2009	

AMLODIPINE BESYLATE; VALSARTAN

TABLET;ORAL

AMLODIPINE BESYLATE AND VALSARTAN

STRIDES PHARMA	EQ 5MG BASE;160MG	A090011 001	Mar 28, 2013
	EQ 5MG BASE;320MG	A090011 003	Mar 28, 2013
	EQ 10MG BASE;160MG	A090011 002	Mar 28, 2013
	EQ 10MG BASE;320MG	A090011 004	Mar 28, 2013
TEVA PHARMS USA	EQ 5MG BASE;160MG	A091235 001	Mar 30, 2015
	EQ 5MG BASE;320MG	A091235 003	Mar 30, 2015
	EQ 10MG BASE;160MG	A091235 002	Mar 30, 2015
	EQ 10MG BASE;320MG	A091235 004	Mar 30, 2015
TORRENT	EQ 5MG BASE;160MG	A202377 001	Mar 30, 2015
	EQ 5MG BASE;320MG	A202377 002	Mar 30, 2015
	EQ 10MG BASE;160MG	A202377 003	Mar 30, 2015
	EQ 10MG BASE;320MG	A202377 004	Mar 30, 2015

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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	
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

AMMONIUM LACTATE

WATSON LABS INC	EQ 12% BASE	A076829 001	Feb 07, 2006
LAC-HYDRIN			
SUN PHARM INDS INC	EQ 12% BASE **	N020508 001	Aug 29, 1996

LOTION; TOPICAL

AMMONIUM LACTATE

WATSON LABS INC	EQ 12% BASE	A075575 001	Jun 11, 2002
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	
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AMOXAPINE

TABLET; ORAL

AMOXAPINE

CHARTWELL RX	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	Aug 01, 1989
	50MG	A072419 001	Aug 01, 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 004	

AMOXICILLIN

CAPSULE; ORAL

AMOXICILLIN

LABS ATRAL	250MG	A062528 001	Aug 07, 1985
	500MG	A062528 002	Aug 07, 1985
MYLAN	250MG	A062067 001	
	500MG	A062067 002	
STRIDES PHARMA	250MG	A062884 001	Feb 25, 1988
	500MG	A062881 001	Feb 25, 1988
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMOXICILLIN

CAPSULE; ORAL

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

BELCHER PHARMS	125MG/5ML	A062059 001	
	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	A061931 001	
	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	
+	US ANTIBIOTICS	200MG/5ML **	N050760 001	Apr 15, 1999
+		400MG/5ML **	N050760 002	Apr 15, 1999

LAROTID

+	GLAXOSMITHKLINE	50MG/ML **	N050460 006
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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

STRIDES PHARMA	875MG	A065344 001	Jan 15, 2009
SUN PHARM INDS LTD	500MG	A065059 001	Nov 24, 2000
	875MG	A065059 002	Nov 24, 2000

AMOXIL

+	US ANTIBIOTICS	500MG **	N050754 002	Jul 10, 1998
+		875MG **	N050754 001	Jul 10, 1998

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMOXICILLIN

TABLET, CHEWABLE;ORAL

AMOXICILLIN

APOTHECON	125MG	A064131 001	May 06, 1996
	250MG	A064131 002	May 06, 1996
HIKMA	125MG	A205274 001	Jun 25, 2020
	250MG	A205274 002	Jun 25, 2020
STRIDES PHARMA	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

+ US ANTIBIOTICS	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
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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, TABLET, CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED)

ANI PHARMS	500MG;500MG;30MG	A200218 001	Aug 30, 2013
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PREVPAC (COPACKAGED)

+ TAKEDA PHARMS USA	500MG;500MG;30MG **	N050757 001	Dec 02, 1997
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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
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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'

+ US ANTIBIOTICS	200MG/5ML;EQ 28.5MG BASE/5ML	N050725 001	May 31, 1996
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AUGMENTIN '400'

+ US ANTIBIOTICS	400MG/5ML;EQ 57MG BASE/5ML	N050725 002	May 31, 1996
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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'

+ US ANTIBIOTICS	250MG;EQ 125MG BASE **	N050564 001	Aug 06, 1984
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AUGMENTIN '500'

+ US ANTIBIOTICS	500MG;EQ 125MG BASE **	N050564 002	Aug 06, 1984
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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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMOXICILLIN; CLAVULANATE POTASSIUM

TABLET, CHEWABLE;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

	400MG;EQ 57MG BASE	A065161 002	Dec 03, 2003
AUGMENTIN '125'			
+ US ANTIBIOTICS	125MG;EQ 31.25MG BASE **	N050597 001	Jul 22, 1985
AUGMENTIN '200'			
+ US ANTIBIOTICS	200MG;EQ 28.5MG BASE	N050726 001	May 31, 1996
AUGMENTIN '250'			
+ US ANTIBIOTICS	250MG;EQ 62.5MG BASE **	N050597 002	Jul 22, 1985
AUGMENTIN '400'			
+ US ANTIBIOTICS	400MG;EQ 57MG BASE	N050726 002	May 31, 1996

TABLET, EXTENDED RELEASE;ORAL

AUGMENTIN XR

+ US ANTIBIOTICS	1GM;EQ 62.5MG BASE	N050785 001	Sep 25, 2002
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AMPHETAMINE

SUSPENSION, EXTENDED RELEASE;ORAL

ADZENYS ER

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

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AUROLIFE PHARMA LLC	1.25MG;1.25MG;1.25MG;1.25MG	A211876 001	Jun 24, 2020
	2.5MG;2.5MG;2.5MG;2.5MG	A211876 002	Jun 24, 2020
	3.75MG;3.75MG;3.75MG;3.75MG	A211876 003	Jun 24, 2020
	5MG;5MG;5MG;5MG	A211876 004	Jun 24, 2020
	6.25MG;6.25MG;6.25MG;6.25MG	A211876 005	Jun 24, 2020
	7.5MG;7.5MG;7.5MG;7.5MG	A211876 006	Jun 24, 2020
NESHER PHARMS	1.25MG;1.25MG;1.25MG;1.25MG	A210080 001	Jul 17, 2019
	2.5MG;2.5MG;2.5MG;2.5MG	A210080 002	Jul 17, 2019
	3.75MG;3.75MG;3.75MG;3.75MG	A210080 003	Jul 17, 2019
	5MG;5MG;5MG;5MG	A210080 004	Jul 17, 2019
	6.25MG;6.25MG;6.25MG;6.25MG	A210080 005	Jul 17, 2019
	7.5MG;7.5MG;7.5MG;7.5MG	A210080 006	Jul 17, 2019
TEVA	1.25MG;1.25MG;1.25MG;1.25MG	A077488 001	Apr 29, 2013
	2.5MG;2.5MG;2.5MG;2.5MG	A077488 002	Apr 29, 2013
	3.75MG;3.75MG;3.75MG;3.75MG	A077488 003	Apr 29, 2013
	5MG;5MG;5MG;5MG	A077488 004	Apr 29, 2013
	6.25MG;6.25MG;6.25MG;6.25MG	A077488 005	Apr 29, 2013
	7.5MG;7.5MG;7.5MG;7.5MG	A077488 006	Apr 29, 2013
DEXTROAMP SACCHARATE,AMP ASPARTATE,DEXTROAMP SULFATE AND AMP SULFATE			
BARR LABS INC	1.25MG;1.25MG;1.25MG;1.25MG	A076536 001	Feb 12, 2013
	2.5MG;2.5MG;2.5MG;2.5MG	A076536 002	Feb 12, 2013
	3.75MG;3.75MG;3.75MG;3.75MG	A076536 003	Feb 12, 2013
	5MG;5MG;5MG;5MG	A076536 004	Feb 12, 2013
	6.25MG;6.25MG;6.25MG;6.25MG	A076536 005	Feb 12, 2013
	7.5MG;7.5MG;7.5MG;7.5MG	A076536 006	Feb 12, 2013

TABLET;ORAL

ADDERALL 10

+ TEVA WOMENS	2.5MG;2.5MG;2.5MG;2.5MG **	N011522 007	Feb 13, 1996
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ADDERALL 12.5

+ TEVA WOMENS	3.125MG;3.125MG;3.125MG;3.125MG **	N011522 012	Aug 31, 2000
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

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	
MYLAN	1.25MG; 1.25MG; 1.25MG; 1.25MG			A206721	001	Nov 10,	2015	
	1.875MG; 1.875MG; 1.875MG; 1.875MG			A206721	002	Nov 10,	2015	
	2.5MG; 2.5MG; 2.5MG; 2.5MG			A206721	003	Nov 10,	2015	
	3.125MG; 3.125MG; 3.125MG; 3.125MG			A206721	004	Nov 10,	2015	
	3.75MG; 3.75MG; 3.75MG; 3.75MG			A206721	005	Nov 10,	2015	
	5MG; 5MG; 5MG; 5MG			A206721	006	Nov 10,	2015	
	7.5MG; 7.5MG; 7.5MG; 7.5MG			A206721	007	Nov 10,	2015	
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								
GLENMARK PHARMS LTD	5MG			A212186	001	Jan 27,	2021	
	10MG			A212186	002	Jan 27,	2021	
+ LANNETT	5MG			A083901	001	Aug 31,	1984	
+	10MG			A083901	002	Aug 31,	1984	
MAYNE PHARMA	5MG			A213898	001	Jul 14,	2020	
	10MG			A213898	002	Jul 14,	2020	
NOVAST LABS	5MG			A213763	001	Aug 24,	2020	
	10MG			A213763	002	Aug 24,	2020	
TABLET, ORALLY DISINTEGRATING; ORAL								
EVEKEO ODT								
+ ARBOR PHARMS LLC	2.5MG			N209905	005	Apr 16,	2021	

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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPHOTERICIN BLOTION; TOPICAL
FUNGIZONE

APOTHECON 3% A060570 001

OINTMENT; TOPICAL
FUNGIZONE

APOTHECON 3% N050313 001

SUSPENSION; ORAL
FUNGIZONE

BRISTOL MYERS SQUIBB 100MG/ML N050341 003

AMPICILLIN SODIUMINJECTABLE; 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

HIKMA 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

HOSPIRA INC EQ 250MG BASE/VIAL A202864 001 Sep 04, 2015

EQ 500MG BASE/VIAL A202864 002 Sep 04, 2015

EQ 1GM BASE/VIAL A202864 003 Sep 04, 2015

EQ 2GM BASE/VIAL A202864 004 Sep 04, 2015

EQ 10GM BASE/VIAL A202865 001 Sep 04, 2015

HQ SPECLT PHARMA 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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPICILLIN SODIUM

INJECTABLE; INJECTION

PENBRITIN-S

+	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
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

AMPICILLIN AND SULBACTAM

HOSPIRA INC

EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A090375	001	Dec 21, 2011
EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A090653	001	Dec 21, 2011
EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A090375	002	Dec 21, 2011
EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A090653	002	Dec 21, 2011
EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL	A090646	001	Dec 21, 2011
EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A065316	001	Jun 29, 2007
EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A065316	002	Jun 29, 2007

PHARM ASSOC

UNASYN

PFIZER

EQ 500MG BASE/VIAL;EQ 250MG BASE/VIAL	N050608	003	Dec 31, 1986
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AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMCILL

PARKE DAVIS

EQ 250MG BASE	A062041	001
EQ 500MG BASE	A062041	002

AMPICILLIN TRIHYDRATE

BELCHER PHARMS

EQ 250MG BASE	A061602	001
EQ 500MG BASE	A061602	002

HERITAGE PHARMA

EQ 250MG BASE	A061502	001
EQ 500MG BASE	A061502	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

SANDOZ

EQ 250MG BASE	A064082	001	Aug 29, 1995
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STRIDES PHARMA

EQ 250MG BASE	A062883	001	Feb 25, 1988
EQ 500MG BASE	A062882	001	Feb 25, 1988

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

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

BELCHER PHARMS	EQ 125MG BASE/5ML	A061601 001	
	EQ 250MG BASE/5ML	A061601 002	
DAVA PHARMS INC	EQ 125MG BASE/5ML	A062982 001	Feb 10, 1989
	EQ 250MG BASE/5ML	A062982 002	Feb 10, 1989
MYLAN	EQ 125MG BASE/5ML	A061829 002	
	EQ 250MG BASE/5ML	A061829 001	
PUREPAC PHARM	EQ 125MG BASE/5ML	A061980 001	
	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	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPICILLIN/AMPICILLIN TRIHYDRATE

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

COSETTE

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

ANAGRELIDE HYDROCHLORIDE

CAPSULE;ORAL

AGRYLIN

+

TAKEDA PHARMS USA

EQ 1MG BASE **

N020333 002 Mar 14, 1997

ANAGRELIDE HYDROCHLORIDE

BARR

EQ 0.5MG BASE

A076530 001 Apr 18, 2005

EQ 1MG BASE

A076530 002 Apr 18, 2005

CHARTWELL RX

EQ 0.5MG BASE

A076683 001 Apr 18, 2005

EQ 1MG BASE

A076683 002 Apr 18, 2005

MYLAN

EQ 0.5MG BASE

A076811 001 Apr 18, 2005

EQ 1MG BASE

A076811 002 Apr 18, 2005

MYLAN PHARMS INC

EQ 0.5MG BASE

A077613 001 Jun 27, 2006

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

WATSON LABS

EQ 0.5MG BASE

A076417 001 Apr 18, 2005

EQ 1MG BASE

A076417 002 Apr 18, 2005

ANASTROZOLE

TABLET;ORAL

ANASTROZOLE

APOTEX INC

1MG

A200654 001 May 11, 2012

FRESENIUS KABI USA

1MG

A090088 001 Jun 28, 2010

HIKMA

1MG

A078485 001 Jun 28, 2010

IMPAX LABS INC

1MG

A091242 001 May 31, 2012

LANNETT CO INC

1MG

A091331 001 Jan 05, 2011

MYLAN

1MG

A091051 001 Jun 28, 2010

PICKET PHARMS

1MG

A090732 001 Jun 28, 2010

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

APIXABAN

TABLET; ORAL

APIXABAN

BIONPHARMA INC

2.5MG

A210152 001 Apr 08, 2020

5MG

A210152 002 Apr 08, 2020

BRECKENRIDGE

2.5MG

A209845 001 Jul 26, 2021

5MG

A209845 002 Jul 26, 2021

MICRO LABS

2.5MG

A210013 001 Dec 23, 2019

5MG

A210013 002 Dec 23, 2019

MYLAN

2.5MG

A210128 001 Dec 23, 2019

5MG

A210128 002 Dec 23, 2019

APOMORPHINE HYDROCHLORIDE

INJECTABLE; SUBCUTANEOUS

APOKYN

MDD US

20MG/2ML (10MG/ML)

N021264 001 Apr 20, 2004

APREMILAST

TABLET; ORAL

APREMILAST

UNICHEM

10MG

A211819 001 Feb 17, 2021

20MG

A211819 002 Feb 17, 2021

30MG

A211819 003 Feb 17, 2021

APREPITANT

CAPSULE; ORAL

EMEND

+ MERCK

40MG **

N021549 003 Jun 30, 2006

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

INJECTABLE; INTRAVENOUS

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 DEXTROSE

SANDOZ

125MG/125ML (1MG/ML)

N201743 001 May 09, 2011

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ARIPIPRAZOLE

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

ARIPIPRAZOLE

MYLAN 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

TEVA PHARMS USA 2MG A078607 001 Apr 28, 2015

5MG A078607 002 Apr 28, 2015

10MG A078608 001 Apr 28, 2015

15MG A078708 001 Apr 28, 2015

20MG A078708 002 Apr 28, 2015

30MG A078708 003 Apr 28, 2015

ZYDUS PHARMS 2MG A090472 001 Jan 07, 2019

5MG A090472 002 Jan 07, 2019

10MG A090472 003 Jan 07, 2019

15MG A090472 004 Jan 07, 2019

20MG A090472 005 Jan 07, 2019

30MG A090472 006 Jan 07, 2019

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

APOTEX INC 50MG A201514 001 Mar 25, 2019

150MG A201514 002 Mar 25, 2019

250MG A201514 003 Mar 25, 2019

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

ARSENIC TRIOXIDE

INGENUS PHARMS LLC 2MG/ML A209315 002 Jan 14, 2021

TRISENOX

+ CEPHALON 1MG/ML ** N021248 001 Sep 25, 2000

ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

ARTICAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE

HOSPIRA 4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML) A079138 001 Jun 18, 2010

ULTACAN

HANSAMED INC 4%;EQ 0.0085MG BASE/1.7ML (4%; EQ 0.005MG BASE/ML) A201751 001 Jul 11, 2017

ULTACAN FORTE

HANSAMED INC 4%;EQ 0.017MG BASE/1.7ML (4%; EQ 0.01MG BASE/ML) A201750 001 Jul 11, 2017

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;

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness 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.C. 9+3

ABRAXIS PHARM	10MG/ML;0.006MG/ML;0.5MCG/ML;1.5MG/ML;2 0	N018440 002	Aug 08, 1985
	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 0	N008809 004	Aug 08, 1985
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	IU/ML;0.04MG/ML;4MG/ML;0.4MG/ML;0.36MG/		
	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;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 0	N018439 002	Aug 08, 1985
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	IU/ML;0.04MG/ML;4MG/ML;0.4MG/ML;0.36MG/		
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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 0	N008809 005	Apr 22, 2004
	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	100MG/VIAL;0.06MG/VIAL;0.005MG/VIAL;15M G/VIAL;5MCG/VIAL;0.4MG/VIAL;40MG/VIAL;4 MG/VIAL;3.6MG/VIAL;3MG/VIAL;1MG/VIAL;10 MG/VIAL	N018933 002	Aug 08, 1985
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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 MG/VIAL;400	N020176 001	Dec 29, 1993
	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		

ASPIRIN

TABLET; ORAL

BAYER EXTRA STRENGTH ASPIRIN FOR MIGRAINE PAIN

BAYER	500MG	N021317 001	Oct 18, 2001
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TABLET, EXTENDED RELEASE; ORAL

8-HOUR BAYER

BAYER	650MG	N016030 001	
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MEASURIN

BAYER	650MG	N016030 002	
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ASPIRIN; BUTALBITAL

TABLET; ORAL

AXOTAL

SAVAGE LABS	650MG;50MG	A088305 001	Oct 13, 1983
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ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

NOSTRUM LABS INC	325MG;50MG;40MG	A078149 001	Jun 13, 2007
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WATSON LABS	325MG;50MG;40MG	A086231 002	Feb 12, 1985
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TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

ACTAVIS ELIZABETH	325MG;50MG;40MG	A086710 002	Aug 23, 1983
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FOSUN PHARMA	325MG;50MG;40MG	A086398 002	Apr 06, 1984
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HALSEY	325MG;50MG;40MG	A089448 001	Dec 01, 1986
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HIKMA INTL PHARMS	325MG;50MG;40MG	A086162 002	Feb 16, 1984
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IVAX PHARMS	325MG;50MG;40MG	A085441 002	Oct 31, 1984
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PURACAP PHARM	325MG;50MG;40MG	A087048 002	Dec 09, 1983
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASPIRIN; BUTALBITAL; CAFFEINE

TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

QUANTUM PHARMICS 325MG; 50MG; 40MG

A088972 001 Jun 18, 1985

WATSON LABS 325MG; 50MG; 40MG

A086237 002 Mar 23, 1984

FIORINAL

+ ALLERGAN 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

STRIDES PHARMA 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

+ BAUSCH 385MG; 30MG; 25MG **

N013416 003 Oct 27, 1982

NORGESIC FORTE

+ BAUSCH 770MG; 60MG; 50MG **

N013416 004 Oct 27, 1982

ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE

SANDOZ 385MG; 30MG; 25MG

A074654 001 Dec 31, 1996

770MG; 60MG; 50MG

A074654 002 Dec 31, 1996

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

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

HERITAGE PHARMS INC 325MG; 200MG

A089594 001 Mar 31, 1989

NOVAST LABS 325MG; 200MG

A040832 001 Jan 07, 2010

OXFORD PHARMS 325MG; 200MG

A040252 001 Dec 10, 1997

SANDOZ 325MG; 200MG

A040116 001 Apr 25, 1996

CARISOPRODOL COMPOUND

WATSON LABS 325MG; 200MG

A088809 001 Oct 03, 1985

SOMA COMPOUND

MEDA PHARMS 325MG; 200MG **

N012365 005 Jul 11, 1983

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASPIRIN; CARISOPRODOL; CODEINE PHOSPHATE

TABLET;ORAL

CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE

OXFORD PHARMS 325MG;200MG;16MG A040283 001 Dec 29, 1998

SANDOZ 325MG;200MG;16MG A040118 001 Apr 16, 1996

SOMA COMPOUND W/ CODEINE

MEDA PHARMS 325MG;200MG;16MG ** N012366 002 Jul 11, 1983

ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE;ORAL

ASPIRIN AND DIPYRIDAMOLE

ANI PHARMS 25MG;200MG A206964 001 Jan 18, 2017

LANNETT CO INC 25MG;200MG A204552 001 Mar 20, 2019

SUN PHARM 25MG;200MG A208572 001 Aug 21, 2018

ASPIRIN; HYDROCODONE BITARTRATE

TABLET;ORAL

AZDONE

SCHWARZ PHARMA 500MG;5MG ** A089420 001 Jan 25, 1988

HYDROCODONE BITARTRATE AND ASPIRIN

LGM PHARMA 500MG;5MG A205479 001 May 28, 2021

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

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

ROBAXISAL

ROBINS AH 325MG;400MG N012281 001

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

ACTAVIS LABS FL INC 325MG;4.8355MG A090084 001 Mar 22, 2011

MAYNE PHARMA INC 325MG;4.8355MG A091670 001 Mar 16, 2011

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET;ORAL

CODOXY

HALSEY 325MG;4.5MG;0.38MG A087464 001 Jul 01, 1982

OXYCODONE AND ASPIRIN

ANI PHARMS 325MG;4.5MG;0.38MG A040255 001 Feb 27, 1998

SUN PHARM INDUSTRIES 325MG;4.5MG;0.38MG A040260 001 Jul 17, 1998

325MG;4.5MG;0.38MG A087794 001 May 26, 1982

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET; ORAL

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; ORAL

PRAVIGARD PAC (COPACKAGED)

BRISTOL MYERS SQUIBB

81MG; 20MG

N021387 001 Jun 24, 2003

81MG; 40MG

N021387 002 Jun 24, 2003

81MG; 80MG

N021387 003 Jun 24, 2003

325MG; 20MG

N021387 004 Jun 24, 2003

325MG; 40MG

N021387 005 Jun 24, 2003

325MG; 80MG

N021387 006 Jun 24, 2003

ASPIRIN; PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DARVON W/ ASA

XANODYNE PHARM

325MG; 65MG

N010996 005

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

ATAZANAVIR SULFATE

AMNEAL

EQ 100MG BASE

A209717 001 Jun 01, 2020

MYLAN

EQ 150MG BASE

A208177 001 Sep 24, 2018

EQ 200MG BASE

A208177 002 Sep 24, 2018

EQ 300MG BASE

A208177 003 Sep 24, 2018

ZYDUS PHARMS

EQ 150MG BASE

A210575 001 Jun 04, 2020

EQ 200MG BASE

A210575 002 Jun 04, 2020

EQ 300MG BASE

A210575 003 Jun 04, 2020

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

INVATECH

25MG

A074265 001 Feb 28, 1994

50MG

A074265 002 Feb 28, 1994

100MG

A074265 003 Feb 28, 1994

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ATENOLOL

TABLET; ORAL

ATENOLOL

	100MG	A074101 003	Jul 17, 1997
SCS	50MG	A073676 001	Oct 30, 1992
	100MG	A073676 002	Oct 30, 1992
STRIDES PHARMA	25MG	A074099 001	Apr 28, 1992
	50MG	A073542 001	Dec 19, 1991
	100MG	A073543 001	Dec 19, 1991
SUN PHARM INDS INC	25MG	A078210 001	Jul 10, 2007
	50MG	A078210 002	Jul 10, 2007
	100MG	A078210 003	Jul 10, 2007
SUN PHARM INDUSTRIES	25MG	A074499 001	Jul 30, 1997
	50MG	A073475 001	Mar 30, 1993
	100MG	A073476 001	Mar 30, 1993
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

MYLAN	50MG; 25MG	A074203 001	Oct 31, 1993
	100MG; 25MG	A074203 002	Oct 31, 1993
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
SUN PHARM INDUSTRIES	50MG; 25MG	A073582 002	Apr 29, 1993
	100MG; 25MG	A073582 001	Apr 29, 1993

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

ATOMOXETINE HYDROCHLORIDE

MYLAN	10MG	A079021 001	Feb 18, 2021
	18MG	A079021 002	Feb 18, 2021
	25MG	A079021 003	Feb 18, 2021
	40MG	A079021 004	Feb 18, 2021
	60MG	A079021 005	Feb 18, 2021
	80MG	A079021 006	Feb 18, 2021
	100MG	A079021 007	Feb 18, 2021
STRATTERA			
LILLY	5MG	N021411 001	Nov 26, 2002

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

DR REDDYS LABS LTD	EQ 10MG BASE	A214659 001	Jul 14, 2021
	EQ 20MG BASE	A214659 002	Jul 14, 2021
	EQ 40MG BASE	A214659 003	Jul 14, 2021
	EQ 80MG BASE	A214659 004	Jul 14, 2021
PERRIGO R AND D	EQ 10MG BASE	A208478 001	Jun 23, 2020
	EQ 20MG BASE	A208478 002	Jun 23, 2020
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

EZETIMIBE AND ATORVASTATIN CALCIUM

SCOV3	EQ 10MG BASE; 10MG	A206084 001	Apr 26, 2017
	EQ 20MG BASE; 10MG	A206084 002	Apr 26, 2017
	EQ 40MG BASE; 10MG	A206084 003	Apr 26, 2017
	EQ 80MG BASE; 10MG	A206084 004	Apr 26, 2017
LIPTRUZET			
+ ORGANON	EQ 10MG BASE; 10MG **	N200153 001	May 03, 2013
+	EQ 20MG BASE; 10MG **	N200153 002	May 03, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ATORVASTATIN CALCIUM; EZETIMIBE

TABLET; ORAL

LIPTRUZET

+	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
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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
MYLAN LABS LTD	10MG/ML	A206096 001	Jun 22, 2017
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
MYLAN LABS LTD	10MG/ML	A206001 001	Apr 07, 2017
WATSON LABS INC	10MG/ML	A074944 001	Jul 28, 1998

TRACRIUM

+	HOSPIRA	10MG/ML **	N018831 002	Jun 20, 1985
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TRACRIUM PRESERVATIVE FREE

+	HOSPIRA	10MG/ML **	N018831 001	Nov 23, 1983
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ATROPINE

SOLUTION; INTRAMUSCULAR

ATROPINE

ABBVIE	EQ 2MG SULFATE/0.7ML	A071295 001	Jan 30, 1987
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ATROPINE (AUTOINJECTOR)

RAFA LABS LTD	EQ 2MG SULFATE/0.7ML (EQ 2MG SULFATE/0.7ML)	N212319 001	Jul 09, 2018
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ATROPINE SULFATE

AEROSOL, METERED; INHALATION

ATROPINE SULFATE

US ARMY	EQ 0.36MG BASE/INH	N020056 001	Sep 19, 1990
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SOLUTION; INTRAVENOUS

ATROPINE SULFATE

+	HOSPIRA	0.5MG/5ML (0.1MG/ML) **	N021146 001	Jul 09, 2001
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ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL

MOTOFEN HALF-STRENGTH

SEBELA IRELAND LTD	0.025MG;0.5MG	N017744 001
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ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENOXYLATE HYDROCHLORIDE W/ ATROPINE SULFATE

SCHERER RP	0.025MG;2.5MG	A086440 001
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SOLUTION; ORAL

COLONOID

MEDPOINTE PHARM HLC	0.025MG/5ML;2.5MG/5ML	A085735 001
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LOMANATE

ALPHARMA US PHARMS	0.025MG/5ML;2.5MG/5ML	A085746 001
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LOMOTIL

GD SEARLE LLC	0.025MG/5ML;2.5MG/5ML	N012699 001
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TABLET; ORAL

COLONOID

MEDPOINTE PHARM HLC	0.025MG;2.5MG	A085737 001
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DI-ATRO

MD PHARM	0.025MG;2.5MG	A085266 001
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

TABLET; ORAL

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	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	
STRIDES PHARMA	0.025MG;2.5MG	A040357 001	May 02, 2000
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	0.14MG/ML;10MG/ML	N019678 001	Nov 06, 1991
	MYLAN INSTITUTIONAL	0.14MG/ML;10MG/ML	N019677 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
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AVOBENZONE; OCTINOXATE; OXYBENZONE

LOTION; TOPICAL

SHADE UVAGUARD

+	BAYER HEALTHCARE LLC	3%;7.5%;3%	N020045 001	Dec 07, 1992
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AZACITIDINE

POWDER; INTRAVENOUS, SUBCUTANEOUS

AZACITIDINE

LUPIN LTD	100MG/VIAL	A210748 001	Feb 27, 2019
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AZATADINE MALEATE

TABLET; ORAL

OPTIMINE

SCHERING	1MG	N017601 001	
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AZATADINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

TRINALIN

SCHERING	1MG;120MG	N018506 001	Mar 23, 1982
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AZATHIOPRINE

TABLET; ORAL

IMURAN

+ SEBELA IRELAND LTD 25MG ** N016324 002

AZATHIOPRINE SODIUM

INJECTABLE; INJECTION

IMURAN

+ CASPER PHARMA LLC EQ 100MG BASE/VIAL ** N017391 001

AZELAIC ACID

AEROSOL, FOAM; TOPICAL

AZELAIC ACID

TEVA PHARMS USA 15% A210928 001 Oct 07, 2020

AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

OPTIVAR

+ MYLAN SPECIALITY LP 0.05% ** N021127 001 May 22, 2000

SPRAY, METERED; NASAL

ASTELIN

+ MYLAN SPCLT VIATRIS 0.137MG/SPRAY N020114 001 Nov 01, 1996

ASTEPRO

+ MYLAN SPECIALITY LP 0.137MG/SPRAY N022203 001 Oct 15, 2008

0.2055MG/SPRAY N022203 002 Aug 31, 2009

AZITHROMYCIN

CAPSULE; ORAL

ZITHROMAX

+ PFIZER EQ 250MG BASE ** N050670 001 Nov 01, 1991

FOR SUSPENSION; ORAL

AZITHROMYCIN

LUPIN LTD

EQ 100MG BASE/5ML A065488 001 May 15, 2015

EQ 200MG BASE/5ML A065488 002 May 15, 2015

SANDOZ

EQ 100MG BASE/5ML A065297 001 Sep 18, 2006

EQ 200MG BASE/5ML A065297 002 Sep 18, 2006

TEVA PHARMS

EQ 100MG BASE/5ML A065419 001 Jun 24, 2008

EQ 200MG BASE/5ML A065419 002 Jun 24, 2008

FOR SUSPENSION, EXTENDED RELEASE; ORAL

ZMAX

+ PF PRISM CV EQ 2GM BASE/BOT N050797 001 Jun 10, 2005

INJECTABLE; INJECTION

AZITHROMYCIN

HIKMA

EQ 500MG BASE/VIAL A065265 001 Jan 18, 2007

MYLAN ASI

EQ 500MG BASE/VIAL A065506 001 Mar 24, 2009

RISING PHARMA

EQ 500MG BASE/VIAL A204732 001 Jan 26, 2017

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

EQ 600MG BASE A065360 001 Jan 08, 2007

TEVA

EQ 250MG BASE A065153 001 Nov 14, 2005

EQ 600MG BASE A065150 001 Nov 14, 2005

YUNG SHIN PHARM

EQ 500MG BASE A211318 001 Mar 17, 2021

ZITHROMAX

+ PFIZER EQ 600MG BASE ** N050730 001 Jun 12, 1996

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

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
+	20MG/ML	N050632 002	May 24, 1989
+	40MG/ML	N050632 001	May 24, 1989
AZTREONAM			
HIKMA	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
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TABLET; ORAL

SPECTROBID

PFIZER	400MG	N050520 001	
	800MG	N050520 002	Sep 12, 1983

BACITRACIN

INJECTABLE; INJECTION

BACIIM

X GEN PHARMS	50,000 UNITS/VIAL	A064153 001	May 09, 1997
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BACITRACIN

AKORN	50,000 UNITS/VIAL	A206719 001	Oct 20, 2017
FRESENIUS KABI USA	50,000 UNITS/VIAL	A065116 001	Dec 03, 2002
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	
	50,000 UNITS/VIAL	A060733 002	
XELLIA PHARMS APS	50,000 UNITS/VIAL	A203177 001	Aug 25, 2014

OINTMENT; OPHTHALMIC

BACIGUENT

PHARMACIA AND UPJOHN	500 UNITS/GM	A060734 001	
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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	
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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	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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	
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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
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OINTMENT;TOPICAL

CORTISPORIN

+	MONARCH PHARMS	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;5,000 UNITS/GM	N050168 002	May 04, 1984
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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
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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
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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
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BACITRACIN-NEOMYCIN-POLYMYXIN

	PHARMADERM	400 UNITS/GM;EQ 3.5MG BASE/GM;5,000 UNITS/GM	A062167 001	
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NEO-POLYCIN

	DOW PHARM	500 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A060647 001	
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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
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BACITRACIN ZINC; POLYMYXIN B SULFATE

AEROSOL;TOPICAL

POLYSPORIN

	GLAXOSMITHKLINE	10,000 UNITS/GM;2,000,000 UNITS/GM	N050167 002	Mar 01, 1985
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OINTMENT;OPHTHALMIC

OCUMYCIN

	PHARMAFAIR	500 UNITS/GM;10,000 UNITS/GM	A062430 001	Apr 08, 1983
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POLYSPORIN

	MONARCH PHARMS	500 UNITS/GM;10,000 UNITS/GM **	A061229 001	
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OINTMENT;TOPICAL

BACITRACIN ZINC-POLYMYXIN B SULFATE

	NASKA	500 UNITS/GM;10,000 UNITS/GM	A062849 001	Nov 13, 1987
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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	
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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	
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BACITRACIN; POLYMYXIN B SULFATE

DISC;TOPICAL

LANABIOTIC

	COMBE	500 UNITS/GM;5,000 UNITS/GM	N050598 001	Sep 22, 1986
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BACLOFEN

TABLET;ORAL

BACLOFEN

	MYLAN	10MG	A077181 001	Jul 29, 2005
		20MG	A077121 002	Jul 29, 2005
	MYLAN PHARMS INC	10MG	A090334 001	Feb 18, 2010
		20MG	A090334 002	Feb 18, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BACLOFEN

TABLET; ORAL

BACLOFEN

SUN PHARM INDS INC	10MG	A077862 001	Aug 14, 2006
	20MG	A077862 002	Aug 14, 2006
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

BALSALAZIDE DISODIUM

CAPSULE; ORAL

BALSALAZIDE DISODIUM

MYLAN

750MG

A077807 001 Dec 28, 2007

TABLET; ORAL

BALSALAZIDE DISODIUM

STRIDES PHARMA

1.1GM

A206336 001 Sep 08, 2015

GIAZO

+ VALEANT PHARMS INTL

1.1GM **

N022205 001 Feb 03, 2012

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

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

ANI PHARMS

5MG

A076333 001 Feb 11, 2004

10MG

A076333 002 Feb 11, 2004

20MG

A076333 003 Feb 11, 2004

40MG

A076333 004 Feb 11, 2004

GENPHARM

5MG

A076476 001 Feb 11, 2004

10MG

A076476 002 Feb 11, 2004

20MG

A076476 003 Feb 11, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

	40MG	A076476 004	Feb 11, 2004
HERITAGE PHARMA	5MG	A076267 001	Feb 11, 2004
	10MG	A076267 002	Feb 11, 2004
	20MG	A076267 003	Feb 11, 2004
	40MG	A076267 004	Feb 11, 2004
RISING PHARMA	5MG	A076430 001	Feb 11, 2004
	10MG	A076430 002	Feb 11, 2004
	20MG	A076430 003	Feb 11, 2004
	40MG	A076430 004	Feb 11, 2004

BENAZEPRIL HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

ANI PHARMS	5MG; 6.25MG	A076342 001	Feb 11, 2004
	5MG; 6.25MG	A076348 001	Feb 11, 2004
	10MG; 12.5MG	A076342 002	Feb 11, 2004
	10MG; 12.5MG	A076348 002	Feb 11, 2004
	20MG; 12.5MG	A076342 003	Feb 11, 2004
	20MG; 12.5MG	A076348 003	Feb 11, 2004
	20MG; 25MG	A076342 004	Feb 11, 2004
	20MG; 25MG	A076348 004	Feb 11, 2004
APOTEX	5MG; 6.25MG	A078794 001	Aug 21, 2014
	10MG; 12.5MG	A078794 002	Aug 21, 2014
	20MG; 12.5MG	A078794 003	Aug 21, 2014
	20MG; 25MG	A078794 004	Aug 21, 2014
AUROBINDO PHARMA USA	5MG; 6.25MG	A076688 001	Feb 11, 2004
	10MG; 12.5MG	A076688 002	Feb 11, 2004
	20MG; 12.5MG	A076688 003	Feb 11, 2004
	20MG; 25MG	A076688 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 + VALIDUS PHARMS	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

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

CORZIDE

+ KING PHARMS LLC	5MG; 40MG	N018647 001	May 25, 1983
+	5MG; 80MG	N018647 002	May 25, 1983

NADOLOL AND BENDROFLUMETHIAZIDE

IMPAX LABS	5MG; 40MG	A077833 001	Mar 30, 2007
	5MG; 80MG	A077833 002	Mar 30, 2007
MYLAN	5MG; 40MG	A078688 001	Feb 15, 2008
	5MG; 80MG	A078688 002	Feb 15, 2008

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

SUN PHARM INDS INC

100MG

A040587 001 Mar 19, 2008

200MG

A040587 002 Mar 19, 2008

TESSALON

+ PFIZER

200MG **

N011210 003 Jun 25, 1999

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL;TOPICAL

BENZAFLIN

BAUSCH

5%;EQ 1% BASE

N050756 002 Apr 20, 2007

CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE

ANDA REPOSITORY

2.5%;EQ 1.2% BASE

A207194 001 Aug 19, 2019

ENCUBE

5%;EQ 1% BASE

A204087 001 Jun 27, 2017

TARO

3.75%;EQ 1.2% BASE

A208683 001 Jun 05, 2018

TOLMAR

5%;1.2%

A203688 001 Aug 25, 2016

BENZOYL PEROXIDE; ERYTHROMYCIN

GEL;TOPICAL

AKTIPAK

+ BIOFRONTERA

5%;3%

N050769 001 Nov 27, 2000

ERYTHROMYCIN AND BENZOYL PEROXIDE

TOLMAR

5%;3%

A065112 001 Mar 29, 2004

BENZPHETAMINE HYDROCHLORIDE

TABLET;ORAL

BENZPHETAMINE HYDROCHLORIDE

EMCURE PHARMS LTD

50MG

A202061 001 Jan 27, 2012

EPIC PHARMA LLC

50MG

A040714 001 Oct 29, 2007

IMPAX LABS

50MG

A040845 001 Nov 18, 2008

SCINOPHARM TAIWAN

50MG

A040578 001 Apr 17, 2006

SPECGX LLC

50MG

A040773 001 Apr 25, 2007

TEDOR PHARM

25MG

A040747 002 Nov 20, 2015

50MG

A040747 001 Mar 30, 2007

DIDREX

+ PFIZER

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BENZTROPINE MESYLATE

INJECTABLE; INJECTION

BENZTROPINE MESYLATE

LUITPOLD

1MG/ML

A091152 001 Mar 29, 2010

TABLET; ORAL

BENZTROPINE MESYLATE

LANNETT CO INC

0.5MG **

A088877 001 Apr 11, 1985

1MG **

A088894 001 Apr 11, 1985

2MG **

A088895 001 Apr 11, 1985

OXFORD PHARMS

0.5MG

A040706 002 Feb 14, 2008

1MG

A040706 003 Feb 14, 2008

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

UPSHER SMITH LABS

0.5MG

A040103 001 Dec 12, 1996

1MG

A040103 002 Dec 12, 1996

2MG

A040103 003 Dec 12, 1996

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 ALCOHOL

LOTION; TOPICAL

ULESFIA

+ SHIONOGI INC

5%

N022129 001 Apr 09, 2009

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BETAMETHASONE BENZOATE

LOTION; TOPICAL

UTICORT

PARKE DAVIS 0.025% N017528 001

OINTMENT; TOPICAL

UTICORT

PARKE DAVIS 0.025% N018089 001

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

BETAMETHASONE DIPROPIONATE

ANDA REPOSITORY EQ 0.05% BASE A076603 001 Jan 23, 2004

DIPROLENE

SCHERING EQ 0.05% BASE N019408 001 Jan 31, 1986

DIPROLENE AF

+ MERCK SHARP DOHME EQ 0.05% BASE ** N019555 001 Apr 27, 1987

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

ACTAVIS MID ATLANTIC EQ 0.05% BASE A070281 001 Jul 31, 1985

ALPHARMA US PHARMS EQ 0.05% BASE A071085 001 Feb 03, 1987

COSETE EQ 0.05% BASE A071882 001 Jun 06, 1988

PHARMGEN 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

SPRAY; TOPICAL

BETAMETHASONE DIPROPIONATE

TARO PHARMS EQ 0.05% BASE/SPRAY A211722 001 Jun 17, 2020

BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

LOTION; TOPICAL

LOTRISONE

+ MERCK SHARP DOHME EQ 0.05% BASE; 1% N020010 001 Dec 08, 2000

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

AEROSOL, FOAM; TOPICAL

BETAMETHASONE VALERATE

RICONPHARMA LLC 0.12% A207144 001 May 24, 2017

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

COSETTE EQ 0.1% BASE A070072 001 Jun 27, 1985

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
HERITAGE PHARMA	5MG	A091256	001	May 04, 2010
	10MG	A091256	002	May 04, 2010
	25MG	A091256	003	May 04, 2010
	50MG	A091256	004	May 04, 2010
IMPAX LABS	5MG	A040721	001	Nov 01, 2006
	10MG	A040721	002	Nov 01, 2006
	25MG	A040721	003	Nov 01, 2006
	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	
	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	
WOCKHARDT	5MG	A040532	001	Sep 29, 2003
	10MG	A040533	001	Sep 29, 2003
	25MG	A040534	001	Sep 29, 2003
	50MG	A040518	001	Sep 29, 2003
MYOTONACHOL				
GLENWOOD	5MG	A084188	001	
	10MG	A084188	003	
	25MG	A084188	004	
URECHOLINE				
ODYSSEY PHARMS	5MG	A089095	001	Dec 19, 1985
+	5MG **	N006536	003	
+	10MG	A088440	001	May 29, 1984
+	10MG **	N006536	002	
+	25MG	A088441	001	May 29, 1984
+	25MG **	N006536	004	
	50MG	A089096	001	Dec 19, 1985
+	50MG **	N006536	005	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BETHANIDINE SULFATE

TABLET; ORAL

TENATHAN

ROBINS AH

10MG

N017675 001

25MG

N017675 002

BETRIXABAN

CAPSULE; ORAL

BEVYXXA

+ PORTOLA PHARMS INC

40MG

N208383 001 Jun 23, 2017

+

80MG

N208383 002 Jun 23, 2017

BICALUTAMIDE

TABLET; ORAL

BICALUTAMIDE

FRESENIUS KABI USA

50MG

A079045 001 May 13, 2010

KUDCO IRELAND

50MG

A077995 001 Jul 06, 2009

MYLAN

50MG

A079185 001 Jul 06, 2009

ROXANE

50MG

A078285 001 Mar 24, 2011

SYNTHON PHARMS

50MG

A077973 001 Jul 06, 2009

TEVA

50MG

A076932 001 Jul 06, 2009

BIMATOPROST

SOLUTION/DROPS; OPHTHALMIC

BIMATOPROST

AKORN

0.03%

A203299 001 Nov 08, 2018

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 BISACODYL

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

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, 5
00MG **

N050719 001 Aug 15, 1996

BISOPROLOL FUMARATE

TABLET; ORAL

BISOPROLOL FUMARATE

MYLAN

5MG

A075831 001 Dec 14, 2005

10MG

A075831 002 Dec 14, 2005

TEVA PHARMS

5MG

A075644 001 Jun 26, 2001

10MG

A075644 002 Jun 26, 2001

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

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
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SOLUTION; INHALATION

TORNALATE

SANOFI AVENTIS US	0.2%	N019548	001	Feb 19, 1992
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BIVALIRUDIN

INJECTABLE; INTRAVENOUS

BIVALIRUDIN

APOTEX	250MG/VIAL	A204876	001	Jul 06, 2017
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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

CIPLA	EQ 15 UNITS BASE/VIAL	A209439	001	Mar 11, 2019
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
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BOSENTAN

TABLET; ORAL

BOSENTAN

ALEMBIC PHARMS LTD	62.5MG	A211461	001	Jan 23, 2020
	125MG	A211461	002	Jan 23, 2020
ALVOGEN PINE BROOK	62.5MG	A206002	001	Apr 26, 2019
	125MG	A206002	002	Apr 26, 2019
AMNEAL PHARMS CO	62.5MG	A209742	001	Apr 26, 2019
	125MG	A209742	002	Apr 26, 2019
CIPLA	62.5MG	A210342	001	Jan 03, 2020
	125MG	A210342	002	Jan 03, 2020
MYLAN	62.5MG	A205173	001	Jan 15, 2020
	125MG	A205173	002	Jan 15, 2020
NATCO PHARMA LTD	62.5MG	A206987	001	Apr 26, 2019
	125MG	A206987	002	Apr 26, 2019

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
HIKMA	50MG/ML	A070545	001	May 14, 1986
	50MG/ML	A070546	001	May 14, 1986
+ HOSPIRA	50MG/ML **	N019030	001	Apr 29, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION

BRETYLIUM TOSYLATE

	50MG/ML	N019033 001	Apr 29, 1986
INTL MEDICATION	50MG/ML	A070119 001	Apr 29, 1986
LUITPOLD	50MG/ML	A070891 001	Jul 26, 1988
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

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

NOVITIUM PHARMA

EQ 0.09% ACID

A201941 001 Feb 10, 2015

RISING PHARMA

EQ 0.09% ACID

A203368 001 Jun 03, 2019

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

TABLET; ORAL

BROMOCRIPTINE MESYLATE

MYLAN

EQ 2.5MG BASE

A076962 001 Sep 24, 2004

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

INJECTABLE; INJECTION

BROMPHENIRAMINE MALEATE

WATSON LABS	10MG/ML	A083821	001	
	100MG/ML	A083820	001	

DIMETANE-TEN

WYETH AYERST	10MG/ML	N011418	002	
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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	
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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
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BROMFED-DM

WOCKHARDT	2MG/5ML; 10MG/5ML; 30MG/5ML	A089681	001	Dec 22, 1988
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BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

PHARM ASSOC	2MG/5ML; 10MG/5ML; 30MG/5ML	A202940	001	Jul 21, 2014
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DIMETANE-DX

+ ROBINS AH	2MG/5ML; 10MG/5ML; 30MG/5ML **	N019279	001	Aug 24, 1984
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BROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

EFIDAC 24 PSEUDOEPHEDRINE HYDROCHLORIDE/BROMPHENIRAMINE MALEATE

ALZA	16MG; 240MG	N019672	001	Mar 29, 1996
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BUCLIZINE HYDROCHLORIDE

TABLET; ORAL

BUCLADIN-S

STUART PHARMS	50MG	N010911	006	
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BUDESONIDE

AEROSOL, METERED; NASAL

RHINOCORT

ASTRAZENECA	0.032MG/INH	N020233	001	Feb 14, 1994
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CAPSULE, DELAYED RELEASE; ORAL

BUDESONIDE

ALVOGEN	3MG	A206724	001	Nov 23, 2016
BARR LABS DIV TEVA	3MG	A090379	001	Apr 02, 2014

POWDER, METERED; INHALATION

PULMICORT

ASTRAZENECA	0.16MG/INH	N020441	002	Jun 24, 1997
	0.32MG/INH	N020441	003	Jun 24, 1997

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

SOLUTION, EXTENDED RELEASE; INFILTRATION

POSIMIR

+ DURECT	660MG/5ML (132MG/ML)	N204803	001	Feb 01, 2021
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BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE

HOSPIRA	0.25%	A070586	001	Mar 03, 1987
	0.25%	N018053	002	
	0.5%	N018053	001	
	0.75%	A070587	001	Mar 03, 1987
	0.75%	N018053	003	
MYLAN ASI	0.25%	A091503	001	Oct 18, 2011
	0.5%	A091503	002	Oct 18, 2011
NOVOCOL HEALTHCARE	0.5%	A211096	001	Feb 19, 2019
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
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
MYLAN ASI	0.25%	A091487	002	Oct 18, 2011
	0.5%	A091487	001	Oct 18, 2011
	0.75%	A091487	003	Oct 18, 2011

INJECTABLE; SPINAL

SENSORCAINE

FRESENIUS KABI USA	0.75%	A071202	001	Apr 15, 1987
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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; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

+ HOSPIRA	0.5%; 0.0091MG/ML	N022046	001	Jul 13, 1983
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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
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BUPRENORPHINE

FILM, EXTENDED RELEASE; TRANSDERMAL

BUPRENORPHINE

MYLAN TECH VIATRIS	5MCG/HR	A210162	001	May 03, 2021
	7.5MCG/HR	A210162	002	May 03, 2021
	10MCG/HR	A210162	003	May 03, 2021
	15MCG/HR	A210162	004	May 03, 2021
	20MCG/HR	A210162	005	May 03, 2021

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUPRENORPHINE HYDROCHLORIDE

IMPLANT; IMPLANTATION

PROBUPHINE

+	TITAN PHARMS	EQ 80MG BASE/IMPLANT	N204442	001	May 26, 2016
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TABLET; SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE

	BARR	EQ 2MG BASE	A090360	001	May 07, 2010
		EQ 8MG BASE	A090360	002	May 07, 2010
	MYLAN	EQ 2MG BASE	A201066	001	Mar 06, 2015
		EQ 8MG BASE	A201066	002	Mar 06, 2015

SUBUTEX

+	INDIVIOR INC	EQ 2MG BASE **	N020732	002	Oct 08, 2002
+		EQ 8MG BASE **	N020732	003	Oct 08, 2002

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; SUBLINGUAL

CASSIPA

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

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

	MAYNE PHARMA INC	EQ 2MG BASE;EQ 0.5MG BASE	A206953	001	Jul 17, 2020
		EQ 8MG BASE;EQ 2MG BASE	A206953	002	Jul 17, 2020
	TEVA PHARMS USA	EQ 2MG BASE;EQ 0.5MG BASE	A091149	001	Sep 08, 2014
		EQ 8MG BASE;EQ 2MG BASE	A091149	002	Sep 08, 2014

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

	AUROBINDO PHARMA USA	75MG	A075491	001	Apr 17, 2000
		100MG	A075491	002	Apr 17, 2000
	HERITAGE PHARMA	75MG	A075310	001	Nov 29, 1999
		100MG	A075310	002	Nov 29, 1999
	INVATECH	75MG	A075613	002	Oct 10, 2000
		100MG	A075613	001	Oct 10, 2000

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
	AUROBINDO PHARMA USA	100MG	A090325	001	Apr 08, 2010
		150MG	A090325	002	Apr 08, 2010
		150MG	A090942	001	Jul 14, 2010
		200MG	A090325	003	Apr 08, 2010
		300MG	A090942	002	Jul 14, 2010
	IMPAX LABS	150MG	A077415	001	Nov 26, 2008
		200MG	A076711	001	Dec 03, 2004
		300MG	A077415	002	Dec 15, 2006
	JUBILANT GENERICS	100MG	A202774	001	Oct 11, 2013
		150MG	A202774	002	Oct 11, 2013
		150MG	A202775	001	Oct 11, 2013
		150MG	A207459	001	Jun 30, 2017
		200MG	A202774	003	Oct 11, 2013
		300MG	A207459	002	Jun 30, 2017
	MYLAN	150MG	A090941	001	May 03, 2010
	SANDOZ	100MG	A076845	001	Jul 14, 2005
		150MG	A076834	001	Jul 14, 2005
		150MG	A076845	002	Jul 14, 2005
	SUN PHARM	150MG	A200695	001	Dec 18, 2014
	TORRENT	100MG	A203969	001	Oct 31, 2014
		150MG	A203969	002	Oct 31, 2014

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

BUPROPION HYDROCHLORIDE

	200MG	A203969 003	Oct 31, 2014
WATSON LABS INC	100MG	A077455 001	Jul 19, 2010
	150MG	A077455 002	Mar 12, 2008
	200MG	A077455 003	Jul 19, 2010
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
+	150MG	N020711 003	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
+		15MG **	N018731 003	Apr 22, 1996
+		30MG **	N018731 004	Apr 22, 1996

BUSPIRONE HYDROCHLORIDE

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

BUSULFAN

INJECTABLE; INJECTION

BUSULFAN

ACTAVIS LLC	6MG/ML	A205139 001	Dec 08, 2017
MYLAN LABS LTD	6MG/ML	A205184 001	Jul 13, 2018

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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUTABARBITAL SODIUM

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

+ 30MG N000793 004

50MG ** N000793 003

100MG ** N000793 005

SARISOL NO. 1

HALSEY 15MG A084719 001

SARISOL NO. 2

HALSEY 30MG A084719 002

SODIUM BUTABARBITAL

HIKMA 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

PADAGIS US 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATE PRESERVATIVE FREE
2MG/ML

A075170 002 Sep 28, 1998

STADOL

+ APOTHECON 2MG/ML **

N017857 004

STADOL PRESERVATIVE FREE

+ APOTHECON 1MG/ML **

N017857 001

+ 2MG/ML **

N017857 002

SPRAY, METERED; NASAL

STADOL

BRISTOL MYERS SQUIBB 1MG/SPRAY **

N019890 001 Dec 12, 1991

CABERGOLINE

TABLET; ORAL

CABERGOLINE

ACTAVIS LABS FL INC 0.5MG

A078035 001 Apr 21, 2008

APOTEX CORP 0.5MG

A201503 001 Mar 08, 2013

IMPAX LABS INC 0.5MG

A077843 001 Jul 03, 2007

MYLAN 0.5MG

A202947 001 Dec 02, 2013

STRIDES PHARMA 0.5MG

A076310 001 Dec 29, 2005

DOSTINEX

+ PFIZER 0.5MG **

N020664 001 Dec 23, 1996

CAFFEINE CITRATE

SOLUTION; INTRAVENOUS

CAFFEINE CITRATE

SUN PHARM EQ 30MG BASE/3ML (EQ 10MG BASE/ML)

A090077 001 Sep 30, 2009

SOLUTION; ORAL

CAFFEINE CITRATE

AM REGENT EQ 30MG BASE/3ML (EQ 10MG BASE/ML)

A090064 001 Nov 20, 2009

CAFFEINE; ERGOTAMINE TARTRATE

SUPPOSITORY; RECTAL

CAFERGOT

+ NOVARTIS 100MG; 2MG **

N009000 002

TABLET; ORAL

CAFERGOT

NOVARTIS 100MG; 1MG

N006620 001

+ SANDOZ 100MG; 1MG

A084294 001

ERGOTAMINE TARTRATE AND CAFFEINE

HIKMA INTL PHARMS 100MG; 1MG

A040510 001 Sep 17, 2004

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCITONIN SALMON

INJECTABLE; INJECTION

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

AM REGENT 0.001MG/ML A075746 001 Sep 26, 2003

0.002MG/ML A075746 002 Sep 26, 2003

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

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

CALCIUM ACETATE

CAPSULE; ORAL

CALCIUM ACETATE

AMNEAL PHARMS 667MG A201658 001 Oct 06, 2014

LOTUS PHARM CO LTD 667MG A203298 001 Jul 26, 2016

PHOSLO

FRESENIUS MEDCL 333.5MG N021160 001 Apr 02, 2001

667MG N021160 002 Apr 02, 2001

TABLET; ORAL

CALCIUM ACETATE

HIKMA 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; ORAL

ACTONEL WITH CALCIUM (COPACKAGED)

+ WARNER CHILCOTT EQ 500MG BASE; 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

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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; 3.0 5GM/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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

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;3.09GM/1000ML; 0ML;7.07GM/1000ML (5000ML)	N021703 013	Oct 10, 2008
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PRISMASOL BGK 4/0 IN PLASTIC CONTAINER

BAXTER HLTHCARE CORP	N/A/1000ML;20GM/1000ML;5.4GM/1000ML;3.0 5GM/1000ML;0.314GM/1000ML;3.09GM/1000ML ;6.46GM/1000ML (5000ML)	N021703 005	Oct 25, 2006
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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/100 0ML;6.46GM/1000ML (5000ML)	N021703 008	Oct 25, 2006
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PRISMASOL BK 0/0 IN PLASTIC CONTAINER

BAXTER HLTHCARE CORP	N/A/1000ML;N/A/1000ML;5.4GM/1000ML;3.05 GM/1000ML;N/A/1000ML;3.09GM/1000ML;6.46 GM/1000ML (5000ML)	N021703 007	Oct 25, 2006
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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
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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/1000 ML;6.46GM/1000ML (5000ML)	N021703 009	Oct 25, 2006
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CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; OXIGLUTATONE; 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/M L;0.38MG/ML;2.1MG/ML;7.14MG/ML;0.42MG/M L	N022193 001	Jul 24, 2008
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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/1 00ML;330MG/100ML;88MG/100ML	N019864 001	Jun 10, 1993
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ISOLYTE R W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	37MG/100ML;5GM/100ML;31MG/100ML;120MG/1 00ML;330MG/100ML;88MG/100ML	N018271 001	
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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/10 0ML;640MG/100ML;500MG/100ML;74MG/100ML	N019867 001	Dec 20, 1993
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ISOLYTE E W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	35MG/100ML;5GM/100ML;30MG/100ML;74MG/10 0ML;640MG/100ML;500MG/100ML;74MG/100ML	N018269 002	Jan 17, 1983
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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/1 00ML;161MG/100ML;94MG/100ML;138MG/100ML	N017390 001	
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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;9.2G M/100ML;9.6GM/100ML	N018807 001	Aug 26, 1983
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B BRAUN	510MG/100ML;30GM/100ML;200MG/100ML;9.4G M/100ML;11GM/100ML	N018807 003	Aug 26, 1983
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DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER

B BRAUN	510MG/100ML;50GM/100ML;200MG/100ML;9.2G M/100ML;9.6GM/100ML	N018807 002	Aug 26, 1983
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B BRAUN	510MG/100ML;50GM/100ML;200MG/100ML;9.4G	N018807 004	Aug 26, 1983
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

SOLUTION;INTRAPERITONEAL

DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER				
	GM/100ML;11GM/100ML			
DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER				
B BRAUN	29MG/100ML;2.5GM/100ML;15MG/100ML;610MG	N018460	006	Jan 29, 1986
	/100ML;560MG/100ML			
DIALYTE W/ DEXTROSE 1.5% IN PLASTIC CONTAINER				
B BRAUN	29MG/100ML;1.5GM/100ML;15MG/100ML;610MG	N018460	001	
	/100ML;560MG/100ML			
DIALYTE W/ DEXTROSE 4.25% IN PLASTIC CONTAINER				
B BRAUN	29MG/100ML;4.25GM/100ML;15MG/100ML;610M	N018460	003	
	G/100ML;560MG/100ML			

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;5	N018379	002	
	67MG/100ML;392MG/100ML			
DELFLX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER				
FRESENIUS MEDCL	25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5	N018379	003	
	67MG/100ML;392MG/100ML			
DELFLX W/ DEXTROSE 3.5% IN PLASTIC CONTAINER				
FRESENIUS MEDCL	25.7MG/100ML;3.5GM/100ML;15.2MG/100ML;5	N018379	007	Jun 24, 1988
	67MG/100ML;392MG/100ML			
DELFLX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER				
FRESENIUS MEDCL	25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;	N018379	001	
	567MG/100ML;392MG/100ML			
DELFLX-LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER				
FRESENIUS MEDCL	25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;5	N018379	004	Jul 07, 1982
	38MG/100ML;448MG/100ML			
DELFLX-LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER				
FRESENIUS MEDCL	25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;5	N018379	005	Jul 07, 1982
	38MG/100ML;448MG/100ML			
DELFLX-LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER				
FRESENIUS MEDCL	25.7MG/100ML;3.5GM/100ML;5.08MG/100ML;5	N018379	008	Jun 24, 1988
	38MG/100ML;448MG/100ML			
DELFLX-LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER				
FRESENIUS MEDCL	25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;	N018379	006	Jul 07, 1982
	538MG/100ML;448MG/100ML			
DIALYTE LM/ DEXTROSE 1.5% IN PLASTIC CONTAINER				
B BRAUN	26MG/100ML;1.5GM/100ML;5MG/100ML;530MG/	N018460	007	Jan 29, 1986
	100ML;450MG/100ML			
	26MG/100ML;1.5GM/100ML;15MG/100ML;560MG	N018460	002	
	/100ML;390MG/100ML			
DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER				
B BRAUN	26MG/100ML;2.5GM/100ML;5MG/100ML;530MG/	N018460	005	Nov 02, 1983
	100ML;450MG/100ML			
	26MG/100ML;5GM/100ML;5MG/100ML;530MG/10	N018460	008	Jan 29, 1986
	0ML;450MG/100ML			
DIALYTE LM/ DEXTROSE 4.25% IN PLASTIC CONTAINER				
B BRAUN	26MG/100ML;4.25GM/100ML;5MG/100ML;530MG	N018460	009	Jan 29, 1986
	/100ML;450MG/100ML			
	26MG/100ML;4.25GM/100ML;15MG/100ML;560M	N018460	004	
	G/100ML;390MG/100ML			
DIANEAL 137 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;5	N017512	001	
	67MG/100ML;392MG/100ML			
DIANEAL 137 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5	N017512	003	
	67MG/100ML;392MG/100ML			
DIANEAL 137 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;	N017512	002	
	567MG/100ML;392MG/100ML			
DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER				
+ BAXTER HLTHCARE	18.3MG/100ML;3.5GM/100ML;5.08MG/100ML;5	N020183	003	Dec 04, 1992
	38MG/100ML;448MG/100ML			
DIANEAL PD-1 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;5	N017512	007	Jul 09, 1984
	67MG/100ML;392MG/100ML			
DIANEAL PD-1 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5	N017512	008	Jul 09, 1984
	67MG/100ML;392MG/100ML			

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DIANEAL PD-1 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	25.7MG/100ML; 3.5GM/100ML; 15.2MG/100ML; 5	N017512	010	Nov 18, 1985
	67MG/100ML; 392MG/100ML			
DIANEAL PD-1 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 5	N017512	009	Jul 09, 1984
	567MG/100ML; 392MG/100ML			
DIANEAL PD-2 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER				
+ BAXTER HLTHCARE	25.7MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 5	N017512	011	Nov 18, 1985
	38MG/100ML; 448MG/100ML			
INPERSOL-IC/LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER				
FRESENIUS	18.4MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5	A020374	001	Jun 13, 1994
	38MG/100ML; 448MG/100ML			
INPERSOL-LC/LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER				
FRESENIUS	18.4MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5	A020374	002	Jun 13, 1994
	38MG/100ML; 448MG/100ML			
INPERSOL-LC/LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER				
FRESENIUS	18.4MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 5	A020374	003	Jun 13, 1994
	38MG/100ML; 448MG/100ML			
INPERSOL-IC/LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER				
FRESENIUS	18.4MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5	A020374	004	Jun 13, 1994
	538MG/100ML; 448MG/100ML			

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/1	N018258	001	
	00ML; 600MG/100ML			

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/1	N018254	001	
	00ML			
DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER				
B BRAUN	33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/1	N018256	001	
	00ML			
	33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/1	N020000	001	Apr 17, 1992
	00ML			
BAXTER HLTHCARE	33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/1	N016695	001	
	00ML			

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/100	N019634	002	Feb 24, 1988
	ML; 62MG/100ML			
DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER				
B BRAUN	20MG/100ML; 5GM/100ML; 30MG/100ML; 600MG/1	N017510	001	
	00ML; 310MG/100ML			
MILES	20MG/100ML; 5GM/100ML; 30MG/100ML; 600MG/1	N018499	001	
	00ML; 310MG/100ML			
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 104MG/100ML; 600MG/	N019685	005	Oct 17, 1988
	100ML; 310MG/100ML			
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 179MG/100ML; 600MG/	N019685	006	Oct 17, 1988
	100ML; 310MG/100ML			
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 254MG/100ML; 600MG/	N019685	007	Oct 17, 1988
	100ML; 310MG/100ML			
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 328MG/100ML; 600MG/	N019685	008	Oct 17, 1988
	100ML; 310MG/100ML			
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 254MG/100ML; 600MG/	N019685	003	Oct 17, 1988
	100ML; 310MG/100ML			
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 328MG/100ML; 600MG/	N019685	004	Oct 17, 1988
	100ML; 310MG/100ML			
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 104MG/100ML; 600MG/	N019685	001	Oct 17, 1988
	100ML; 310MG/100ML			

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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;44 8MG/100ML	N019395 001	Mar 26, 1986
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INPERSOL-ZM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

FRESENIUS MEDCL	25.7MG/100ML;2.5GM/100ML;538MG/100ML;44 8MG/100ML	N019395 002	Mar 26, 1986
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INPERSOL-ZM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

FRESENIUS MEDCL	25.7MG/100ML;4.25GM/100ML;538MG/100ML;4 48MG/100ML	N019395 003	Mar 26, 1986
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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
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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
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	35MG/100ML;30MG/100ML;74MG/100ML;640MG/ 100ML;500MG/100ML;74MG/100ML	N019718 001	Sep 29, 1989
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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	
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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
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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
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SOLUTION;IRRIGATION

RINGER'S IN PLASTIC CONTAINER

ABBOTT	33MG/100ML;30MG/100ML;860MG/100ML	N018462 001	
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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
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B BRAUN	20MG/100ML;30MG/100ML;600MG/100ML;310MG /100ML	N018023 001	
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MILES	20MG/100ML;30MG/100ML;600MG/100ML;310MG /100ML	N018417 001	
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SOLUTION;IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER

BAXTER HLTHCARE	20MG/100ML;30MG/100ML;600MG/100ML;310MG /100ML	N019933 001	Aug 29, 1989
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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
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LILLY	EQ 90MG CALCIUM/5ML	N006470 001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

CAPREOMYCIN SULFATE

INJECTABLE; INJECTION

CAPASTAT SULFATE

+ AKORN

EQ 1GM BASE/VIAL N050095 001

CAPREOMYCIN SULFATE

HISUN PHARM HANGZHOU	EQ 1GM BASE/VIAL	A204796 001	Oct 18, 2018
MYLAN LABS LTD	EQ 1GM BASE/VIAL	A202634 001	Nov 27, 2017

CAPTOPRIL

TABLET; ORAL

CAPOTEN

+ STRIDES PHARMA

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

CAPTOPRIL

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

CHARTWELL RX

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

100MG

A074363 004 Nov 09, 1995

100MG

A074519 004 Feb 13, 1996

COSETTE

12.5MG

A074433 001 Feb 13, 1996

12.5MG

A074462 001 Feb 13, 1996

12.5MG

A074483 001 Feb 13, 1996

25MG

A074433 002 Feb 13, 1996

25MG

A074462 002 Feb 13, 1996

25MG

A074483 002 Feb 13, 1996

50MG

A074433 003 Feb 13, 1996

50MG

A074462 003 Feb 13, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CAPTOPRILTABLET; ORAL
CAPTOPRIL

	50MG	A074483	003	Feb 13, 1996
	100MG	A074433	004	Feb 13, 1996
	100MG	A074462	004	Feb 13, 1996
	100MG	A074483	004	Feb 13, 1996
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	A074590	004	Aug 30, 1996
	25MG	A074590	002	Aug 30, 1996
	50MG	A074590	001	Aug 30, 1996
	100MG	A074590	003	Aug 30, 1996
MYLAN	12.5MG	A074434	001	Feb 13, 1996
	25MG	A074434	002	Feb 13, 1996
	50MG	A074434	003	Feb 13, 1996
	100MG	A074434	004	Feb 13, 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
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
SETON PHARMS	12.5MG	A212223	001	Oct 30, 2019
	25MG	A212223	002	Oct 30, 2019
	50MG	A212223	003	Oct 30, 2019
	100MG	A212223	004	Oct 30, 2019
STRIDES PHARMA	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
TEVA	12.5MG	A074322	001	Feb 13, 1996
	25MG	A074322	002	Feb 13, 1996
	50MG	A074322	003	Feb 13, 1996
	100MG	A074322	004	Feb 13, 1996
WATSON LABS	12.5MG	A074386	001	May 23, 1996
	12.5MG	A074451	001	Feb 13, 1996
	12.5MG	A074576	001	Apr 23, 1996
	25MG	A074386	002	May 23, 1996
	25MG	A074451	002	Feb 13, 1996
	25MG	A074576	002	Apr 23, 1996
	50MG	A074386	003	May 23, 1996
	50MG	A074451	003	Feb 13, 1996
	50MG	A074576	003	Apr 23, 1996
	100MG	A074386	004	May 23, 1996
	100MG	A074451	004	Feb 13, 1996
	100MG	A074576	004	Apr 23, 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CAPTOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CAPOZIDE 50/25

+ APOTHECON 50MG;25MG ** N018709 003 Oct 12, 1984

CAPTOPRIL AND HYDROCHLOROTHIAZIDE

COSETTE 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

RISING PHARMA 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

STRIDES PHARMA 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

TEGRETOL

+ NOVARTIS 100MG N018281 001

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

EQ 20GM BASE/VIAL N050298 007

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARBENICILLIN INDANYL SODIUM

TABLET; ORAL

GEOCILLIN

PFIZER

EQ 382MG BASE

N050435 001

CARBIDOPA; ENTACAPONE; LEVODOPA

TABLET; ORAL

CARBIDOPA, LEVODOPA AND ENTACAPONE

MYLAN

12.5MG; 200MG; 50MG

A203424 001 Aug 13, 2020

18.75MG; 200MG; 75MG

A203424 002 Aug 13, 2020

25MG; 200MG; 100MG

A203424 003 Aug 13, 2020

31.25MG; 200MG; 125MG

A203424 004 Aug 13, 2020

37.5MG; 200MG; 150MG

A203424 005 Aug 13, 2020

50MG; 200MG; 200MG

A203424 006 Aug 13, 2020

WOCKHARDT LTD

12.5MG; 200MG; 50MG

A090786 001 Nov 20, 2012

18.75MG; 200MG; 75MG

A090833 001 Nov 20, 2012

25MG; 200MG; 100MG

A090833 002 Nov 20, 2012

31.25MG; 200MG; 125MG

A090833 003 Nov 20, 2012

37.5MG; 200MG; 150MG

A090833 004 Nov 20, 2012

50MG; 200MG; 200MG

A090833 005 Nov 20, 2012

CARBIDOPA; LEVODOPA

TABLET; ORAL

CARBIDOPA AND LEVODOPA

ANI PHARMS

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

SINEMET CR

+ ORGANON

25MG; 100MG

N019856 002 Dec 24, 1992

+

50MG; 200MG

N019856 001 May 30, 1991

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

RISING PHARMA

10MG; 100MG

A078893 001 Sep 18, 2008

25MG; 100MG

A078893 002 Sep 18, 2008

25MG; 250MG

A078893 003 Sep 18, 2008

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

VINTAGE PHARMS

4MG/5ML

A040814 001 Feb 26, 2008

TABLET; ORAL

CARBINOXAMINE MALEATE

CYPRESS PHARM

4MG

A090417 001 Aug 23, 2010

STRIDES PHARMA

4MG

A040639 002 May 30, 2008

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARBINOXAMINE MALEATE

TABLET; ORAL

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
HIKMA	50MG/VIAL	A076099 001	Oct 14, 2004
	150MG/VIAL	A076099 002	Oct 14, 2004
	450MG/VIAL	A076099 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
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
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; INTRAVENOUS			
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)	A077432 001	Sep 29, 2006
	150MG/15ML (10MG/ML)	A077432 002	Sep 29, 2006
	450MG/45ML (10MG/ML)	A077432 003	Sep 29, 2006
	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
	150MG/15ML (10MG/ML)	A077266 002	Feb 15, 2006
MEITHEAL	50MG/5ML (10MG/ML)	A077096 001	Jun 14, 2005
	150MG/15ML (10MG/ML)	A077096 002	Jun 14, 2005
	450MG/45ML (10MG/ML)	A077096 003	Jun 14, 2005
	600MG/60ML (10MG/ML)	A077096 004	Jun 03, 2013
MYLAN INSTITUTIONAL	50MG/5ML (10MG/ML)	A077998 001	Apr 24, 2007
	150MG/15ML (10MG/ML)	A077998 002	Apr 24, 2007
	450MG/45ML (10MG/ML)	A077998 003	Apr 24, 2007
MYLAN LABS LTD	50MG/5ML (10MG/ML)	A091063 001	Nov 09, 2011
	150MG/15ML (10MG/ML)	A091063 002	Nov 09, 2011
	450MG/45ML (10MG/ML)	A091063 003	Nov 09, 2011
	600MG/60ML (10MG/ML)	A091063 004	Nov 09, 2011
	1GM/100ML (10MG/ML)	A091478 001	Nov 23, 2011
NOVAST LABS	450MG/45ML (10MG/ML)	A208487 003	Apr 26, 2017
	600MG/60ML (10MG/ML)	A208487 004	Apr 26, 2017
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
PLIVA LACHEMA	50MG/5ML (10MG/ML)	A078631 001	Dec 02, 2008
	150MG/15ML (10MG/ML)	A078631 002	Dec 02, 2008
	450MG/45ML (10MG/ML)	A078631 003	Dec 02, 2008
	600MG/60ML (10MG/ML)	A078631 004	Dec 02, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARBOPLATIN

INJECTABLE; INTRAVENOUS

CARBOPLATIN

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
TEVA PHARMS USA	50MG/5ML (10MG/ML)	A077139 001	Sep 21, 2005
	150MG/15ML (10MG/ML)	A077139 002	Sep 21, 2005
	450MG/45ML (10MG/ML)	A077139 003	Sep 21, 2005
	600MG/60ML (10MG/ML)	A077139 004	Sep 21, 2005
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

CARFILZOMIB

POWDER; INTRAVENOUS

CARFILZOMIB

APOTEX	30MG/VIAL	A211185 001	Mar 20, 2020
	60MG/VIAL	A209425 001	Mar 16, 2020
BRECKENRIDGE	10MG/VIAL	A209330 001	Jun 11, 2021
	60MG/VIAL	A209330 002	Jun 11, 2021

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
HIKMA INTL PHARMS	350MG	A040124 001	Jan 24, 1996
PIONEER PHARMS	350MG	A089390 001	Oct 13, 1988
SANDOZ	350MG	A089566 001	Aug 30, 1988
STRIDES PHARMA	250MG	A205513 001	Nov 12, 2015
	350MG	A205513 002	Nov 12, 2015
SUN PHARM INDS LTD	350MG	A040755 001	Feb 27, 2007
SUN PHARM INDUSTRIES	350MG	A089346 001	Oct 17, 1991
WATSON LABS	350MG	A040152 001	Dec 03, 1996
	350MG	A085433 001	
WATSON LABS TEVA	350MG	A086179 001	

RELA

SCHERING 350MG N012155 001

CARPENAZINE 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	

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
BAUSCH AND LOMB	1%	A075546 001	Jan 20, 2000

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARTEOLOL HYDROCHLORIDETABLET; ORAL
CARTROL

10MG

N019204 003 Dec 28, 1988

CARVEDILOLTABLET; ORAL
CARVEDILOL

ACI

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

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

SUN PHARM INDS INC

3.125MG

A077346 004 Sep 05, 2007

6.25MG

A077346 001 Sep 05, 2007

12.5MG

A077346 002 Sep 05, 2007

25MG

A077346 003 Sep 05, 2007

CASPOFUNGIN ACETATEPOWDER; INTRAVENOUS
CASPOFUNGIN ACETATE

CIPLA

50MG/VIAL

A209489 001 Jul 12, 2018

70MG/VIAL

A209489 002 Jul 12, 2018

MYLAN LABS LTD

50MG/VIAL

A207650 001 Sep 29, 2017

70MG/VIAL

A207650 002 Sep 29, 2017

CEFACLORCAPSULE; 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

HERITAGE PHARMA

EQ 250MG BASE

A064148 001 May 23, 1996

EQ 500MG BASE

A064148 002 May 23, 1996

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

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

RANBAXY

EQ 125MG BASE/5ML

A064166 001 Oct 02, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFACTOR

FOR SUSPENSION;ORAL

CEFACTOR

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
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CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE;ORAL

CEFADROXIL

CSPC OUYI	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
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FOR SUSPENSION;ORAL

CEFADROXIL

ANI PHARMS	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
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DURICEF

+	WARNER CHILCOTT	EQ 1GM BASE **	N050528	001
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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
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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
DR REDDYS	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
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
HIKMA	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
	EQ 10GM BASE/VIAL	A062807 005	Jan 12, 1988
	EQ 20GM BASE/VIAL	A062807 006	Jan 12, 1988
HOSPIRA INC	EQ 500MG BASE/VIAL	A065226 001	Apr 21, 2005
	EQ 1GM BASE/VIAL	A065226 002	Apr 21, 2005
	EQ 1GM BASE/VIAL	A065244 001	Aug 12, 2005
	EQ 1GM BASE/VIAL	A201654 001	Feb 03, 2016
	EQ 10GM BASE/VIAL	A065247 001	Aug 12, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

CEFAZOLIN SODIUM

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

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

LILLY

EQ 500MG BASE/VIAL

A062557 001 Sep 10, 1985

EQ 1GM BASE/VIAL

A062557 002 Sep 10, 1985

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

ASTRAL

EQ 1GM BASE/VIAL

A212721 001 Jul 21, 2020

EQ 2GM BASE/VIAL

A212721 002 Jul 21, 2020

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

HOSPIRA INC

EQ 500MG BASE/VIAL

A065369 001 Jun 18, 2007

EQ 1GM BASE/VIAL

A065369 002 Jun 18, 2007

EQ 1GM BASE/VIAL

A202268 001 Jul 30, 2012

EQ 2GM BASE/VIAL

A065369 003 Jun 18, 2007

EQ 2GM BASE/VIAL

A202268 002 Jul 30, 2012

CEFIXIME

CAPSULE; ORAL

SUPRAX

+ LUPIN LTD

400MG **

N203195 001 Jun 01, 2012

FOR SUSPENSION; ORAL

CEFIXIME

SANDOZ INC

100MG/5ML

A206144 001 Nov 17, 2017

200MG/5ML

A206144 002 Nov 17, 2017

SUPRAX

+ LEDERLE

100MG/5ML **

N050622 001 Apr 28, 1989

LUPIN PHARMS

100MG/5ML

A065129 001 Feb 23, 2004

TABLET; ORAL

SUPRAX

+ LEDERLE

200MG **

N050621 001 Apr 28, 1989

+

400MG **

N050621 002 Apr 28, 1989

LUPIN PHARMS

400MG

A065130 001 Feb 12, 2004

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

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	
WOCKHARDT	EQ 1GM BASE/VIAL	A065197	001	Aug 29, 2006	
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	
HOSPIRA INC	EQ 500MG BASE/VIAL	A065290	001	Aug 11, 2006	
	EQ 1GM BASE/VIAL	A065290	002	Aug 11, 2006	
	EQ 1GM BASE/VIAL	A065293	001	Aug 10, 2006	
	EQ 1GM BASE/VIAL	A203132	001	Feb 19, 2016	
	EQ 2GM BASE/VIAL	A065290	003	Aug 11, 2006	
	EQ 2GM BASE/VIAL	A065293	002	Aug 10, 2006	
	EQ 2GM BASE/VIAL	A203132	002	Feb 19, 2016	
	EQ 10GM BASE/VIAL	A065292	001	Aug 10, 2006	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFOTAXIME SODIUM

INJECTABLE; INJECTION

CEFOTAXIME SODIUM

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
WOCKHARDT	EQ 500MG BASE/VIAL	A065197 002	Jun 20, 2008
	EQ 2GM BASE/VIAL	A065197 003	Jun 20, 2008
CLAFORAN			
SANOFI AVENTIS US	EQ 1GM BASE/VIAL	A062659 001	Jan 13, 1987
	EQ 2GM BASE/VIAL	A062659 002	Jan 13, 1987
+ VALIDUS PHARMS	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			
VALIDUS PHARMS	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			
VALIDUS PHARMS	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 1GM BASE/VIAL	A063293 001	Apr 29, 1993
	EQ 2GM BASE/VIAL	A063293 002	Apr 29, 1993
	EQ 10GM BASE/VIAL	N050588 003	Apr 25, 1988
CEFOTAN IN PLASTIC CONTAINER			
TELIGENT	EQ 20MG BASE/ML	N050694 002	Jul 30, 1993
	EQ 40MG BASE/ML	N050694 001	Jul 30, 1993
CEFOTETAN			
FRESENIUS KABI USA	EQ 10GM BASE/VIAL	A065375 001	Aug 09, 2007
HIKMA	EQ 1GM BASE/VIAL	A091031 001	Oct 26, 2011
	EQ 2GM BASE/VIAL	A091031 002	Oct 26, 2011
WEST-WARD PHARM CORP	EQ 10GM BASE/VIAL	A091030 001	Oct 26, 2011

CEFOTIAM HYDROCHLORIDE

INJECTABLE; INJECTION

CERADON

TAKEDA	EQ 1GM BASE/VIAL	N050601 001	Dec 30, 1988
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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
HOSPIRA INC	EQ 1GM BASE/VIAL	A065313 001	Jan 23, 2006
	EQ 2GM BASE/VIAL	A065313 002	Jan 23, 2006
	EQ 10GM BASE/VIAL	A065312 001	Feb 13, 2006
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			
+ MERCK	EQ 20MG BASE/ML **	N050581 003	Sep 20, 1984
+	EQ 40MG BASE/ML **	N050581 004	Sep 20, 1984
MEFOXIN IN PLASTIC CONTAINER			
MYLAN INSTITUTIONAL	EQ 20MG BASE/ML	A063182 001	Jan 25, 1993
	EQ 40MG BASE/ML	A063182 002	Jan 25, 1993
MEFOXIN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
+ MERCK	EQ 20MG BASE/ML **	N050581 002	Sep 20, 1984
+	EQ 40MG BASE/ML **	N050581 001	Sep 20, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFPARAMIDE SODIUM

INJECTABLE; INJECTION

CEFPARAMIDE 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

SANDOZ

EQ 50MG BASE/5ML

A090031 001 Jan 14, 2009

EQ 100MG BASE/5ML

A090031 002 Jan 14, 2009

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

TEVA PHARMS

125MG/5ML

A065236 001 Dec 08, 2005

250MG/5ML

A065236 002 Dec 08, 2005

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

WOCKHARDT

250MG

A065428 001 Jun 14, 2007

500MG

A065428 002 Jun 14, 2007

CEFZIL

+ CORDEN PHARMA

250MG **

N050664 001 Dec 23, 1991

+

500MG **

N050664 002 Dec 23, 1991

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

WOCKHARDT

1GM/VIAL

A065196 001 Oct 15, 2008

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFTAZIDIME

INJECTABLE; INJECTION

PENTACEF

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

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

EQ 10GM BASE/VIAL

A065232 001 Aug 02, 2005

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFTRIAXONE SODIUM

INJECTABLE; INJECTION

ROCEPHIN

	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
HOSPIRA INC	EQ 250MG BASE/VIAL	A065230 001	Aug 02, 2005
	EQ 500MG BASE/VIAL	A065230 002	Aug 02, 2005
	EQ 1GM BASE/VIAL	A065230 003	Aug 02, 2005
	EQ 2GM BASE/VIAL	A065230 004	Aug 02, 2005
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

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

ANI PHARMS	EQ 250MG BASE	A065190 001	Oct 18, 2004
	EQ 500MG BASE	A065190 002	Oct 18, 2004
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFUROXIME AXETIL

TABLET; ORAL

CEFUROXIME AXETIL

	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
HOSPIRA INC	EQ 1.5GM BASE/VIAL	A065483 002	Oct 15, 2008
	EQ 1.5GM BASE/VIAL	A065503 001	Oct 15, 2008
	EQ 7.5GM BASE/VIAL	A065484 001	Oct 15, 2008
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
HOSPIRA INC	EQ 750MG BASE/VIAL	A065483 001	Oct 15, 2008
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

CELECOXIB

CAPSULE; ORAL

CELECOXIB

ACIC PHARMS	200MG	A212925 001	Dec 09, 2020
	400MG	A212925 002	Dec 09, 2020
JUBILANT GENERICS	50MG	A207061 001	Apr 04, 2017
	100MG	A207061 002	Apr 04, 2017
	200MG	A207061 003	Apr 04, 2017
	400MG	A207061 004	Apr 04, 2017

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

	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	
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 (IN)	EQ 250MG BASE	A062791 001	Jun 11, 1987
	EQ 500MG BASE	A062791 002	Jun 11, 1987
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	EQ 250MG BASE	N050405 002	
+	EQ 333MG BASE **	N050405 004	May 12, 2006
+	EQ 500MG BASE	N050405 003	
+	EQ 750MG BASE	N050405 005	May 12, 2006

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	EQ 100MG BASE/ML **	N050406 003	
+	EQ 125MG BASE/5ML **	N050406 001	
+	EQ 250MG BASE/5ML **	N050406 002	

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

CEPHALOGLYCIN

CAPSULE; ORAL

KAFOCIN

LILLY	250MG	N050219 001	
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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	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEPHAPIRIN SODIUM

INJECTABLE; INJECTION

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

HIKMA

EQ 500MG BASE/VIAL

A062720 001 Jul 02, 1987

EQ 1GM BASE/VIAL

A062720 002 Jul 02, 1987

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

CERITINIB

CAPSULE; ORAL

ZYKADIA

+ NOVARTIS

150MG

N205755 001 Apr 29, 2014

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CERIVASTATIN SODIUMTABLET; ORAL
BAYCOL

0.8MG

N020740 006 Jul 24, 2000

CERULETIDE DIETHYLAMINEINJECTABLE; INJECTION
TYMTRAN

PHARMACIA AND UPJOHN 0.02MG/ML

N018296 001

CETIRIZINE HYDROCHLORIDE

SYRUP; ORAL

CETIRIZINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC 5MG/5ML
AUROBINDO PHARMA LTD 5MG/5ML
LANNETT CO INC 5MG/5ML
PHARM ASSOC 5MG/5ML
RANBAXY LABS LTD 5MG/5ML
TORRENT 5MG/5ML
WOCKHARDT 5MG/5MLA078617 001 Feb 02, 2010
A090751 001 Dec 16, 2009
A078496 001 Sep 25, 2009
A078412 001 Jun 18, 2008
A077472 001 Jun 18, 2008
A078870 001 Apr 27, 2009
A078757 001 Aug 28, 2009

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

ACTAVIS MID ATLANTIC 5MG/5ML
CYPRESS PHARM 5MG/5ML
PHARM ASSOC 5MG/5ML
RANBAXY LABS LTD 5MG/5ML
TORRENT 5MG/5MLA090378 002 May 09, 2008
A090300 001 Oct 10, 2008
A090188 002 Apr 22, 2008
A090183 002 Apr 24, 2008
A090474 002 Mar 30, 2009

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

ACTAVIS MID ATLANTIC 5MG/5ML
CYPRESS PHARM 5MG/5ML
PHARM ASSOC 5MG/5ML
RANBAXY LABS LTD 5MG/5ML
TORRENT 5MG/5MLA090378 001 May 09, 2008
A090300 002 Oct 10, 2008
A090188 001 Apr 22, 2008
A090183 001 Apr 24, 2008
A090474 001 Mar 30, 2009

ZYRTEC

J AND J CONSUMER INC 5MG/5ML **

N020346 001 Sep 27, 1996

TABLET; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

CIPLA LTD 5MG
10MG
HERITAGE PHARMA 5MG
10MG
SUN PHARM INDS INC 5MG
10MG
TORRENT PHARMS LLC 5MG
10MG
UNICHEM 5MG
10MGA077318 001 Jul 25, 2013
A077318 002 Jul 25, 2013
A078615 003 Dec 28, 2007
A078615 004 Dec 28, 2007
A077499 001 Dec 27, 2007
A077499 002 Dec 27, 2007
A079191 001 Apr 15, 2010
A079191 004 Apr 15, 2010
A078680 003 Jun 26, 2009
A078680 004 Jun 26, 2009

CETIRIZINE HYDROCHLORIDE HIVES

SUN PHARM INDS INC 5MG
10MG
UNICHEM 5MG
10MGA077499 003 Dec 27, 2007
A077499 004 Dec 27, 2007
A078680 001 Jun 26, 2009
A078680 002 Jun 26, 2009

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

TORRENT PHARMS LLC 5MG
10MGA079191 003 Apr 15, 2010
A079191 002 Apr 15, 2010

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

+ 10MG

N019835 006 Nov 16, 2007

TABLET, CHEWABLE; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

SUN PHARM INDS INC 5MG
10MGA077631 004 Jan 11, 2008
A077631 003 Jan 11, 2008

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

SUN PHARM INDS INC 5MG
10MGA077631 001 Jan 11, 2008
A077631 002 Jan 11, 2008

CHILDREN'S ZYRTEC ALLERGY

+ J AND J CONSUMER INC 2.5MG

N021621 007 Nov 30, 2020

+ 5MG **

N021621 003 Nov 16, 2007

+ 10MG **

N021621 004 Nov 16, 2007

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CETIRIZINE HYDROCHLORIDE

TABLET, CHEWABLE;ORAL

CHILDREN'S ZYRTEC HIVES RELIEF

+ J AND J CONSUMER INC 5MG **

N021621 005 Nov 16, 2007

+ 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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

+ PARKE DALE EQ 1GM BASE/VIAL

N050155 001

MYCHEL-S

ANGUS EQ 1GM BASE/VIAL

A060132 001

CHLORAMPHENICOL; HYDROCORTISONE ACETATE

FOR SUSPENSION; OPHTHALMIC

CHLOROMYCETIN HYDROCORTISONE

PARKE DALE 12.5MG/VIAL; 25MG/VIAL

N050202 001

CHLORAMPHENICOL; HYDROCORTISONE ACETATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

OPHTHOCORT

PARKE DALE 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

CHLORAMPHENICOL; PREDNISOLONE

OINTMENT; OPHTHALMIC

CHLOROPTIC-P S.O.P.

ALLERGAN 1%; 0.5%

A061188 001

CHLORDIAZEPOXIDE

CAPSULE, EXTENDED RELEASE; ORAL

LIBRELEAS

VALEANT PHARM INTL 30MG

N017813 001 Sep 12, 1983

TABLET; ORAL

LIBRITABS

VALEANT PHARM INTL 5MG
10MG
25MGA085482 001
A085481 001
A085488 001CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

A-POXIDE

ABBOTT 5MG
5MG
10MG
10MG
25MG
25MGA085447 001
A085517 001
A085447 002
A085518 001
A085447 003
A085513 001

CHLORDIAZACHEL

RACHELLE 5MG
10MG
25MGA085086 001
A084639 001
A085087 001

CHLORDIAZEPOXIDE HYDROCHLORIDE

ASCOT 5MG
10MG
25MGA087525 001 Jan 07, 1982
A087524 001 Jan 07, 1982
A087512 001 Jan 07, 1982CHARTWELL RX 5MG
10MG
25MGA084678 001
A084041 001
A084679 002FERRANTE 5MG
10MG
25MGA085118 001
A085119 001
A085120 001HALSEY 5MG
10MG
25MGA085340 001
A085339 001
A084685 001IMPAX LABS 5MG
10MG
25MGA086213 001
A085113 001
A086212 001IVAX SUB TEVA PHARMS 5MG
10MGA083741 001
A083742 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE

	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
	25MG	A086494	001	
	25MG	A088707	001	Jan 18, 1985
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

BAUSCH 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	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL

BRIAN CARE

SOAPCO

4%

A071419 001 Dec 17, 1987

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

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

CHLOROPROCAINE HYDROCHLORIDE

HOSPIRA

2%

A087447 001 Apr 16, 1982

3%

A087446 001 Apr 16, 1982

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

500MG **

N006002 001

CHLOROQUINE PHOSPHATE

HIKMA PHARMS

250MG

A083082 001

500MG

A083082 002 Sep 17, 1999

IMPAX LABS

250MG

A080880 001

500MG

A040516 001 Aug 29, 2003

MD PHARM

250MG

A087228 001

PUREPAC PHARM

250MG

A080886 001

TEVA

250MG

A087504 001 Jan 13, 1982

WATSON LABS

250MG

A087979 001 Dec 21, 1982

500MG

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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	
MYLAN	250MG	A084217	002	
	500MG	A084217	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				
+ AKORN	250MG **	N011145	004	
+	500MG **	N011145	002	

CHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

ALDOCLOR-150

MERCK	150MG;250MG	N016016	001	
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ALDOCLOR-250

MERCK	250MG;250MG	N016016	002	
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METHYLDOPA AND CHLOROTHIAZIDE

PAR PHARM	150MG;250MG	A070783	001	Nov 06, 1987
	250MG;250MG	A070654	001	Nov 06, 1987

CHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CHLOROTHIAZIDE AND RESERPINE

HIKMA	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
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DIUPRES-500

MERCK	500MG;0.125MG	N011635	006	Aug 26, 1987
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CHLOROTRIANISENE

CAPSULE; ORAL

CHLOROTRIANISENE

BANNER PHARMACAPS	12MG	A084652	001	
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TACE

SANOFI AVENTIS US	12MG	N008102	004	
	25MG	N011444	001	
	72MG	N016235	001	

CHLOROXYLINE

SHAMPOO; TOPICAL

CAPITROL

WESTWOOD SQUIBB	2%	N017594	001	
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CHLORPHENESIN CARBAMATE

TABLET; ORAL

MAOLATE

PAMLAB LLC	400MG	N014217	002	
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CHLORPHENIRAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

CHLORPHENIRAMINE MALEATE

AUROLIFE PHARMA LLC	12MG	A070797	001	Aug 12, 1988
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TELDRIN

GLAXOSMITHKLINE	8MG	N017369	001	
	12MG	N017369	002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPHENIRAMINE MALEATE

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	
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SYRUP; ORAL

CHLOR-TRIMETON

SCHERING	2MG/5ML	N006921 006	
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CHLORPHENIRAMINE MALEATE

PHARM ASSOC	2MG/5ML	A087520 001	Feb 10, 1982
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TABLET; ORAL

ANTAGONATE

BAYER PHARMS	4MG	A083381 001	
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CHLOR-TRIMETON

SCHERING	4MG	N006921 002	
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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	
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	

KLOROMIN

HALSEY	4MG	A083629 001	
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PHENETRON

LANNETT	4MG	A080846 001	
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TABLET, EXTENDED RELEASE; ORAL

CHLOR-TRIMETON

+ BAYER HEALTHCARE LLC	8MG **	N007638 001	
+	12MG **	N007638 002	

EFIDAC 24 CHLORPHENIRAMINE MALEATE

ALZA	16MG	N019746 002	Nov 18, 1994
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CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND CHLORPHENIRAMINE MALEATE

ACELLA	4MG/5ML; 5MG/5ML	A206891 001	Jun 09, 2017
TRIS PHARMA INC	4MG/5ML; 5MG/5ML	A206438 001	Jan 27, 2015

VITUZ

+ PERSION	4MG/5ML; 5MG/5ML	N204307 001	Feb 20, 2013
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION;ORAL

HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

MAYNE PHARMA INC	4MG/5ML;5MG/5ML;60MG/5ML	A205657 001	Aug 03, 2015
TORRENT	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
ZUTRIPRO			
+ PERSION	4MG/5ML;5MG/5ML;60MG/5ML **	N022439 001	Jun 08, 2011

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	
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CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

CODIMAL-L.A. 12

SCHWARZ PHARMA	12MG;120MG	N018935 001	Apr 15, 1985
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ISOCOLOR

FISONS	8MG;120MG	N018747 001	Mar 06, 1986
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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	
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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
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CHLORPHERMINE HYDROCHLORIDE

TABLET;ORAL

PRE-SATE

PARKE DAVIS	EQ 65MG BASE	N014696 001	
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CHLORPROMAZINE

SUPPOSITORY;RECTAL

THORAZINE

+ GLAXOSMITHKLINE	25MG **	N009149 024	
+ GLAXOSMITHKLINE	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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPROMAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

CHLORPROMAZINE HYDROCHLORIDE

WOCKHARDT	30MG/ML	A087032	001	Jul 08, 1982
	100MG/ML	A087053	001	

CHLORPROMAZINE HYDROCHLORIDE INTENSOL

HIKMA	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	
DR REDDYS	25MG/ML	A080365	001	
MARSAM PHARMS LLC	25MG/ML	A089563	001	Apr 15, 1988
WATSON LABS	25MG/ML	A085591	001	
WYETH AYERST	25MG/ML	A080370	001	

THORAZINE

+	GLAXOSMITHKLINE	25MG/ML **	N009149	011
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SYRUP; ORAL

CHLORPROMAZINE HYDROCHLORIDE

ALPHARMA US PHARMS	10MG/5ML	A086712	001	
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SONAZINE

FOSUN PHARMA	10MG/5ML	A083040	001	
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THORAZINE

+	GLAXOSMITHKLINE	10MG/5ML **	N009149	022
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TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

ABBOTT	10MG	A084414	001	
	25MG	A084415	001	
	50MG	A084411	001	
	100MG	A084412	001	
	200MG	A084413	001	
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	
MAYNE PHARMA	10MG	A213590	001	Aug 31, 2020
	25MG	A213590	002	Aug 31, 2020
	50MG	A213590	003	Aug 31, 2020
	100MG	A213590	004	Aug 31, 2020
	200MG	A213590	005	Aug 31, 2020
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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPROMAZINE HYDROCHLORIDE

TABLET;ORAL

CHLORPROMAZINE HYDROCHLORIDE

	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	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
	100MG	A089446	001	Nov 17, 1986
	250MG	A087353	001	
	250MG	A088813	001	Oct 19, 1984
	250MG	A088826	001	Sep 26, 1984
	250MG	A088919	001	Oct 16, 1984
	250MG	A088922	001	Apr 12, 1985
	250MG	A089447	001	Nov 17, 1986
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
MYLAN	100MG	A088549	002	Jun 01, 1984
	250MG	A088549	001	Jun 01, 1984
PAR PHARM	100MG	A088175	001	Feb 27, 1984
	250MG	A088176	001	Feb 27, 1984
RISING	100MG	A088725	001	Aug 31, 1984
	250MG	A088726	001	Aug 31, 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPROPAMIDE

TABLET; ORAL

DIABINESE

+ PFIZER

100MG

N011641 003

+

250MG

N011641 006

GLUCAMIDE

ANI PHARMS

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

ANI PHARMS

25MG

A087296 001

25MG

A087706 001

25MG

A088164 001 Jan 09, 1984

50MG

A087689 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

COSETTE

50MG

A088651 001 May 30, 1985

DAVA PHARMS INC

25MG

A087451 001

50MG

A087450 001

IVAX PHARMS

25MG

A087555 001

50MG

A087176 001

50MG

A087947 001 Feb 27, 1984

KV PHARM

25MG

A087311 001

50MG

A087312 001

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

50MG

A087029 001

50MG

A087082 001

50MG

A087521 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORTHALIDONE

TABLET; ORAL

HYGROTON

+	SANOFI AVENTIS US	25MG **	N012283	004	
+		50MG **	N012283	003	

THALITONE

MONARCH PHARMS	25MG	A088051	001	Nov 12, 1982
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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	
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REGROTON

SANOFI AVENTIS US	50MG; 0.25MG	N015103	001	
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CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

ACTAVIS ELIZABETH	250MG	A088928	001	May 08, 1987
	500MG	A040113	001	Sep 29, 1995
BARR	500MG	A089895	001	May 04, 1988
OHM LABS	250MG	A081298	001	Dec 29, 1993
	500MG	A081299	001	Dec 29, 1993
PIONEER PHARMS	250MG	A089592	001	Jan 06, 1989
	500MG	A089948	001	Jan 06, 1989
RISING	250MG	A089852	001	May 04, 1988
STRIDES PHARMA	250MG	A087981	001	Sep 20, 1983
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	
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PARAFON FORTE DSC

+	JANSSEN R AND D	500MG **	N011529	002	Jun 15, 1987
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STRIFON FORTE DSC

FERNDAL LABS	500MG	A081008	001	Dec 23, 1988
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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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHOLESTYRAMINE

POWDER; ORAL

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

CHARTWELL RX

EQ 4GM RESIN/PACKET

A074561 001 Aug 15, 1996

EQ 4GM RESIN/SCOOPFUL

A074561 002 Aug 15, 1996

LOCHOLEST LIGHT

CHARTWELL RX

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

+

EQ 4GM RESIN/SCOOPFUL **

N016640 003

QUESTRAN LIGHT

+ BRISTOL MYERS

EQ 4GM RESIN/PACKET **

N019669 001 Dec 05, 1988

+

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

CHOLINE C-11

INJECTABLE; INTRAVENOUS

CHOLINE C-11

UCSF RODIOPHARM

4-33.1mCi/ML

A208444 001 Nov 20, 2017

CHOLINE FENOFIBRATE

CAPSULE, DELAYED RELEASE; ORAL

FENOFIBRIC ACID

MYLAN PHARMS INC

EQ 45MG FENOFIBRIC ACID

A200913 001 Mar 25, 2013

EQ 135MG FENOFIBRIC ACID

A200913 002 Mar 25, 2013

CHROMIC CHLORIDE

INJECTABLE; INJECTION

CHROMIC CHLORIDE

ABRAXIS PHARM

EQ 0.004MG CHROMIUM/ML

N019271 001 May 05, 1987

CHROMIC PHOSPHATE P-32

INJECTABLE; INJECTION

PHOSPHOCOL P32

CURIUM

5mCi/ML

N017084 001

CICLOPIROX

SOLUTION; TOPICAL

CICLOPIROX

AKORN

8%

A078975 001 Feb 17, 2010

ENCUBE

8%

A077687 001 Sep 18, 2007

MYLAN PHARMS INC

8%

A078567 001 Sep 18, 2007

TEVA PHARMS

8%

A078079 001 Sep 18, 2007

CIDOFIVIR

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

EQ 500MG BASE/VIAL; 500MG/VIAL

A090825 002 Nov 16, 2011

EQ 500MG BASE/VIAL; 500MG/VIAL

A091007 001 Nov 16, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CILASTATIN SODIUM; IMIPENEM

POWDER; INTRAVENOUS

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

A077022 002 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

NOSTRUM LABS INC

50MG

A077708 001 Sep 28, 2009

100MG

A077708 002 Sep 28, 2009

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

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

HIKMA

200MG

A074890 001 Dec 18, 1998

300MG

A074890 002 Dec 18, 1998

400MG

A074890 003 Dec 18, 1998

800MG

A074890 004 Dec 18, 1998

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

NOVITIUM PHARMA

300MG

A074340 001 Jun 23, 1995

400MG

A074340 002 Jun 23, 1995

800MG

A074339 001 Jun 23, 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CIMETIDINETABLET; ORAL
CIMETIDINE

	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			
DAVA PHARMS INC	EQ 300MG BASE/2ML	A074428 001	Apr 25, 1996
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
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	EQ 300MG BASE/5ML	A074610 001	Sep 26, 1996
	EQ 300MG BASE/5ML	A074859 001	Jul 09, 1998
	EQ 300MG BASE/5ML	A075110 001	Jun 18, 1998
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
PHARM ASSOC	EQ 300MG BASE/5ML	A075560 001	Mar 15, 2000
TAGAMET			
GLAXOSMITHKLINE	EQ 300MG BASE/5ML **	N017924 001	

CINACALCET HYDROCHLORIDE

TABLET; ORAL

CINACALCET HYDROCHLORIDE			
LUPIN LTD	EQ 30MG BASE	A210548 001	Jun 28, 2019
	EQ 60MG BASE	A210548 002	Jun 28, 2019
	EQ 90MG BASE	A210548 003	Jun 28, 2019
MYLAN	EQ 30MG BASE	A203422 001	Oct 16, 2018
	EQ 60MG BASE	A203422 002	Oct 16, 2018
	EQ 90MG BASE	A203422 003	Oct 16, 2018

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

FOR SUSPENSION; ORAL

CIPROFLOXACIN

CHARTWELL	250MG/5ML	A200563 001	Mar 05, 2014
	500MG/5ML	A200563 002	Mar 05, 2014

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
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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
DR REDDYS	200MG/20ML (10MG/ML)	A077782 001	Aug 28, 2006
	400MG/40ML (10MG/ML)	A077782 002	Aug 28, 2006
FRESENIUS KABI USA	200MG/20ML (10MG/ML)	A076484 001	Aug 28, 2006
	400MG/40ML (10MG/ML)	A076484 002	Aug 28, 2006
HOSPIRA	200MG/20ML (10MG/ML)	A077245 001	Aug 28, 2006
	400MG/40ML (10MG/ML)	A077245 002	Aug 28, 2006

CIPROFLOXACIN IN DEXTROSE 5%

HIKMA FARMACEUTICA	200MG/100ML	A076757 001	Apr 21, 2008
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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
RUBICON	EQ 0.3% BASE	A075928 001	Jun 09, 2004

TABLET; ORAL

CIPRO

+	BAYER HLTHCARE	EQ 100MG BASE **	N019537 001	Apr 08, 1996
+		EQ 750MG BASE **	N019537 004	Oct 22, 1987

CIPROFLOXACIN HYDROCHLORIDE

AMNEAL	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 250MG BASE	A075685 002	Jun 09, 2004
	EQ 500MG BASE	A075685 003	Jun 09, 2004
	EQ 750MG BASE	A075685 001	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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CIPROFLOXACIN HYDROCHLORIDE

TABLET; ORAL

CIPROFLOXACIN HYDROCHLORIDE

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
RISING PHARMA	EQ 100MG BASE	A075817 001	Jun 25, 2007
	EQ 250MG BASE	A075817 002	Jun 09, 2004
	EQ 750MG BASE	A075817 004	Jun 09, 2004
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
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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

ANI PHARMS	212.6MG;EQ 287.5MG BASE	A077809 002	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
MYLAN PHARMS INC	212.6MG;EQ 287.5MG BASE	A078183 001	Mar 22, 2007
	425.2MG;EQ 574.9MG BASE	A078183 002	Mar 22, 2007

CISAPRIDE MONOHYDRATE

SUSPENSION; ORAL

PROPULSID

JANSSEN PHARMS	EQ 1MG BASE/ML	N020398 001	Sep 15, 1995
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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
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CISATRACURIUM BESYLATE

INJECTABLE; INJECTION

CISATRACURIUM BESYLATE

ACCORD HLTHCARE	EQ 2MG BASE/ML	A205873 001	Jun 16, 2017
PIRAMAL HLTHCARE UK	EQ 2MG BASE/ML	A212190 001	May 04, 2020
SAGENT PHARMS INC	EQ 2MG BASE/ML	A201851 001	Nov 06, 2020
ZYDUS PHARMS	EQ 2MG BASE/ML	A213527 001	Aug 31, 2020

CISATRACURIUM BESYLATE PRESERVATIVE FREE

ACCORD HLTHCARE	EQ 2MG BASE/ML	A205872 001	Jun 16, 2017
	EQ 10MG BASE/ML	A205872 002	Jun 16, 2017

CISPLATIN

INJECTABLE; INJECTION

CISPLATIN

BEDFORD	10MG/VIAL	A074713 001	Nov 14, 2000
	50MG/VIAL	A074713 002	Nov 14, 2000
MYLAN LABS LTD	1MG/ML	A091062 001	Apr 18, 2012
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
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

PHARM ASSOC

EQ 10MG BASE/5ML

A077601 001 Nov 15, 2005

TABLET;ORAL

CELEXA

ALLERGAN

EQ 60MG BASE

N020822 004 Jul 17, 1998

CITALOPRAM HYDROBROMIDE

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

HERITAGE PHARMA

EQ 10MG BASE

A077033 001 Oct 28, 2004

EQ 10MG BASE

A077034 001 Jun 30, 2005

EQ 10MG BASE

A077213 001 Mar 31, 2006

EQ 10MG BASE

A077232 001 Oct 31, 2005

EQ 20MG BASE

A077033 002 Oct 28, 2004

EQ 20MG BASE

A077034 002 Jun 30, 2005

EQ 20MG BASE

A077213 002 Mar 31, 2006

EQ 20MG BASE

A077232 002 Oct 31, 2005

EQ 40MG BASE

A077033 003 Oct 28, 2004

EQ 40MG BASE

A077034 003 Jun 30, 2005

EQ 40MG BASE

A077213 003 Mar 31, 2006

EQ 40MG BASE

A077232 003 Oct 31, 2005

JUBILANT GENERICS

EQ 10MG BASE

A205407 001 Dec 23, 2015

EQ 20MG BASE

A205407 002 Dec 23, 2015

EQ 40MG BASE

A205407 003 Dec 23, 2015

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 INDS INC

EQ 10MG BASE

A077032 001 Nov 12, 2004

EQ 20MG BASE

A077032 002 Nov 12, 2004

EQ 40MG BASE

A077032 003 Nov 12, 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

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

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

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

AJANTA PHARMA LTD 250MG

A206714 001 Apr 25, 2019

500MG

A206714 002 Apr 25, 2019

HIKMA 250MG

A065178 002 May 25, 2004

500MG

A065178 001 May 25, 2004

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

WOCKHARDT 250MG

A065266 001 May 31, 2006

500MG

A065266 002 May 31, 2006

TABLET, EXTENDED RELEASE; ORAL

BIAXIN XL

+ ABBVIE 500MG **

N050775 001 Mar 03, 2000

CLARITHROMYCIN

ANI PHARMS 500MG

A065250 001 Aug 25, 2005

LUPIN LTD 500MG

A202532 001 Sep 15, 2015

NOSTRUM LABS INC 500MG

A203243 001 Feb 29, 2016

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLEMASTINE FUMARATE

SYRUP;ORAL

TAVIST

+ NOVARTIS

EQ 0.5MG BASE/5ML **

N018675 001 Jun 28, 1985

TABLET;ORAL

CLEMASTINE FUMARATE

ANI PHARMS

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

CLEVIDIPINE

EMULSION;INTRAVENOUS

CLEVIPREX

+ CHIESI

125MG/250ML (0.5MG/ML)

N022156 003 Nov 08, 2013

CLIDINIUM BROMIDE

CAPSULE;ORAL

QUARZAN

ROCHE

2.5MG

N010355 001

5MG

N010355 002

CLINDAMYCIN HYDROCHLORIDE

CAPSULE;ORAL

CLEOCIN

PHARMACIA AND UPJOHN

EQ 75MG BASE

A061809 001

EQ 150MG BASE

A061809 002

CLINDAMYCIN HYDROCHLORIDE

LANNETT CO INC

EQ 150MG BASE

A065243 003 Aug 12, 2005

EQ 300MG BASE

A065243 001 Aug 12, 2005

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, 1991

CLINDAMYCIN PALMITATE HYDROCHLORIDE

FOR SOLUTION;ORAL

CLEOCIN

PHARMACIA AND UPJOHN

EQ 75MG BASE/5ML **

A061827 001

CLINDAMYCIN PALMITATE HYDROCHLORIDE

MYLAN

EQ 75MG BASE/5ML

A203063 001 May 25, 2016

CLINDAMYCIN PHOSPHATE

AEROSOL, FOAM;TOPICAL

CLINDAMYCIN PHOSPHATE

TARO PHARM INDS LTD

1%

A210004 001 Mar 11, 2020

CREAM;VAGINAL

CLEOCIN

PFIZER

EQ 2% BASE

N050680 001 Aug 11, 1992

INJECTABLE;INJECTION

CLEOCIN PHOSPHATE

PHARMACIA AND UPJOHN

EQ 150MG BASE/ML

A061839 001

CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER

+ PFIZER

EQ 6MG BASE/ML

N050639 001 Aug 30, 1989

+

EQ 12MG BASE/ML

N050639 002 Aug 30, 1989

+

EQ 18MG BASE/ML

N050639 003 Apr 10, 1991

CLINDAMYCIN PHOSPHATE

ABRAXIS PHARM

EQ 150MG BASE/ML

A062747 001 Jun 03, 1988

ALMAJECT

EQ 150MG BASE/ML

A062801 001 Jul 24, 1987

BEDFORD

EQ 150MG BASE/ML

A063163 001 Jun 30, 1994

BRISTOL MYERS SQUIBB

EQ 150MG BASE/ML

A062908 001 Feb 01, 1989

HIKMA

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

IGI LABS INC

EQ 150MG BASE/ML

A062928 001 Feb 13, 1989

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLINDAMYCIN PHOSPHATE

INJECTABLE; INJECTION

CLINDAMYCIN PHOSPHATE

LOCH	EQ 150MG BASE/ML	A062905 001	May 09, 1988
MARSAM PHARMS LLC	EQ 150MG BASE/ML	A062913 001	Oct 20, 1988
RISING PHARMA	EQ 150MG BASE/ML	A204748 001	Oct 10, 2017
	EQ 150MG BASE/ML	A204749 001	Oct 10, 2017
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

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%

ABRAXIS PHARM	EQ 12MG BASE/ML	N050636 001	Dec 22, 1989
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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

+ PFIZER	EQ 1% BASE	N050537 001	
PHARMACIA AND UPJOHN	EQ 1% BASE	A062363 001	Feb 08, 1982

CLINDAMYCIN PHOSPHATE

BOCA PHARMA LLC	EQ 1% BASE	A062944 001	Jan 11, 1989
FOUGERA PHARMS	EQ 1% BASE	A065254 001	Feb 14, 2006
G AND W LABS INC	EQ 1% BASE	A062811 001	Sep 01, 1988
NOVAST LABS	EQ 1% BASE	A064108 001	Sep 27, 1996
VINTAGE PHARMS	EQ 1% BASE	A062930 001	Jun 28, 1989
	EQ 1% BASE	A203343 001	May 29, 2015
WOCKHARDT BIO AG	EQ 1% BASE	A063304 001	Jul 15, 1997

SWAB; TOPICAL

CLEOCIN

+ PFIZER	EQ 1% BASE	N050537 002	Feb 22, 1994
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CLIOQUINOL; NYSTATIN

OINTMENT; TOPICAL

NYSTAFORM

BAYER PHARMS	10MG/GM; 100,000 UNITS/GM	N050235 001	
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CLOBAZAM

SUSPENSION; ORAL

CLOBAZAM

TEVA PHARMS USA	2.5MG/ML	A211032 001	Jan 31, 2020
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TABLET; ORAL

CLOBAZAM

ACCORD HLTHCARE	10MG	A212398 001	May 23, 2019
	20MG	A212398 002	May 23, 2019
APOTEX	10MG	A209853 001	Jun 09, 2020
	20MG	A209853 002	Jun 09, 2020
BRECKENRIDGE	5MG	A209308 003	Oct 19, 2021
CELLTRION	10MG	A211959 001	Dec 09, 2020
	20MG	A211959 002	Dec 09, 2020
LANNETT CO INC	10MG	A212092 001	Oct 30, 2019
	20MG	A212092 002	Oct 30, 2019
MSN	10MG	A213404 001	May 11, 2021
	20MG	A213404 002	May 11, 2021
TARO	10MG	A209440 001	Oct 22, 2018
	20MG	A209440 002	Oct 22, 2018

ONFI

+ LUNDBECK PHARMS LLC	5MG **	N202067 001	Oct 21, 2011
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLOBETASOL PROPIONATE

AEROSOL, FOAM;TOPICAL

CLOBETASOL PROPIONATE

INGENUS PHARMS LLC

0.05%

A206805 001 Jul 31, 2017

CREAM;TOPICAL

CLOBETASOL PROPIONATE

PAI HOLDINGS PHARM

0.05%

A211207 001 Mar 26, 2021

TEVA PHARMS USA

0.05%

A074087 001 Feb 16, 1994

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

LOTION;TOPICAL

CLOBETASOL PROPIONATE

AKORN

0.05%

A211348 001 Oct 26, 2018

OINTMENT;TOPICAL

CLOBETASOL PROPIONATE

ACTAVIS MID ATLANTIC

0.05%

A074128 001 Aug 03, 1994

AMNEAL

0.05%

A210551 001 Aug 21, 2018

TORRENT

0.05%

A212926 001 Oct 25, 2019

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

AKORN

0.05%

A207218 001 Apr 28, 2017

APOTEX

0.05%

A210446 001 Apr 17, 2018

CLOFARABINE

SOLUTION;INTRAVENOUS

CLOFARABINE

HOSPIRA INC

20MG/20ML (1MG/ML)

A210283 001 Dec 27, 2018

NOVAST LABS

20MG/20ML (1MG/ML)

A210270 001 Sep 14, 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

GRANATA BIO

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

25MG

A074958 001 Aug 26, 1997

50MG

A074849 002 Apr 04, 1997

50MG

A074958 002 Aug 26, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

CLOMIPRAMINE HYDROCHLORIDE

	75MG	A074849	003	Apr 04, 1997
	75MG	A074958	003	Aug 26, 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
	75MG	A074751	003	Sep 30, 1998

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

AUROBINDO PHARMA USA	0.5MG	A075150	001	Oct 05, 1998
	1MG	A075150	002	Oct 05, 1998
	2MG	A075150	003	Oct 05, 1998
MYLAN PHARMS INC	0.5MG	A074940	001	Oct 30, 1997
	1MG	A074940	002	Oct 30, 1997
	2MG	A074940	003	Oct 30, 1997
RUBICON	0.5MG	A075468	001	Oct 06, 2000
	1MG	A075468	002	Oct 06, 2000
	2MG	A075468	003	Oct 06, 2000
SANDOZ	0.5MG	A074925	001	Sep 30, 1997
	1MG	A074925	002	Sep 30, 1997
	2MG	A074925	003	Sep 30, 1997
SUN PHARM INDS INC	0.5MG	A075423	001	Apr 27, 2001
	1MG	A075423	002	Apr 27, 2001
	2MG	A075423	003	Apr 27, 2001
TEVA	0.5MG	A074920	001	Aug 04, 1998
	1MG	A074920	002	Aug 04, 1998
	2MG	A074920	003	Aug 04, 1998
WATSON LABS	0.5MG	A074964	001	Dec 30, 1997
	1MG	A074964	002	Dec 30, 1997
	2MG	A074964	003	Dec 30, 1997

KLONOPIN

CHEPLAPHARM	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
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TABLET, EXTENDED RELEASE; ORAL

NEXICLON XR

TRIS PHARMA INC	EQ 0.17MG BASE	N022500	001	Dec 03, 2009
	EQ 0.26MG BASE	N022500	002	Dec 03, 2009

CLONIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CLONIDINE HYDROCHLORIDE

AM REGENT	1MG/10ML (0.1MG/ML)	A091104	001	Oct 08, 2009
	5MG/10ML (0.5MG/ML)	A091104	002	Oct 08, 2009

DURACLON

+	MYLAN INSTITUTIONAL	5MG/10ML (0.5MG/ML)	N020615	002	Apr 27, 1999
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TABLET; ORAL

CATAPRES

+	BOEHRINGER INGELHEIM	0.1MG	N017407	001
+		0.2MG	N017407	002
+		0.3MG	N017407	003

CLONIDINE HYDROCHLORIDE

AM THERAP	0.1MG	A070881	001	Jul 08, 1986
	0.2MG	A070882	001	Jul 08, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLONIDINE HYDROCHLORIDE

TABLET;ORAL

CLONIDINE HYDROCHLORIDE

	0.3MG	A070883	001	Jul 08, 1986
AUROBINDO PHARMA LTD	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
RISING PHARMA	0.1MG	A070317	002	Jul 09, 1987
	0.2MG	A070317	003	Jun 09, 1987
	0.3MG	A070317	001	Jun 09, 1987
SUN PHARM INDS INC	0.1MG	A090329	001	Jul 03, 2014
	0.2MG	A090329	002	Jul 03, 2014
	0.3MG	A090329	003	Jul 03, 2014
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
	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
UPSHER SMITH LABS	0.1MG	A211433	001	Oct 12, 2018
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
ANI PHARMS	EQ 300MG BASE	A090625	001	May 17, 2012
	EQ 300MG BASE	A091216	001	May 17, 2012
CADILA	EQ 75MG BASE	A201686	001	Oct 10, 2012
CELLTRION	EQ 75MG BASE	A202266	001	Aug 14, 2012
	EQ 300MG BASE	A202266	002	Nov 20, 2012
RISING	EQ 75MG BASE	A077665	001	May 17, 2012
	EQ 300MG BASE	A077665	002	May 17, 2012
SUN PHARM INDUSTRIES	EQ 75MG BASE	A078133	001	Jun 10, 2013

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
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
RISING	3.75MG	A072112	002	Aug 26, 1988
	7.5MG	A072112	003	Aug 26, 1988
	15MG	A072112	001	Aug 26, 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
	15MG	A071749	001	Jun 23, 1987
AUROLIFE PHARMA LLC	3.75MG	A072514	002	May 11, 1990
	7.5MG	A072514	003	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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLORAZEPATE DIPOTASSIUM

TABLET;ORAL

GEN-XENE

ALRA

3.75MG

A071787 001 Apr 26, 1988

7.5MG

A071788 001 Apr 26, 1988

15MG

A071789 001 Apr 26, 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

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLOXACILLIN SODIUM

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	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	12.5MG **	N021590	004	May 30, 2007
+	25MG **	N021590	001	Feb 10, 2004
	50MG **	N021590	003	Jun 03, 2005
+	100MG **	N021590	002	Feb 10, 2004
+	150MG **	N021590	005	Jul 09, 2010
+	200MG **	N021590	006	Jul 09, 2010

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
PROMETH VC W/ CODEINE				
NOSTRUM LABS INC	10MG/5ML;5MG/5ML;6.25MG/5ML	A088764	001	Oct 31, 1984
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
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

HIKMA	30MG/5ML	N202245	001	Jun 30, 2011
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

COLCHICINE

CAPSULE; ORAL

COLCHICINE

PAR PHARM INC

0.6MG

A208678 001 Nov 29, 2018

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

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

BAR, CHEWABLE; ORAL

WELCHOL

+ DAIICHI SANKYO INC

3.75GM

N210895 001 Apr 03, 2019

CAPSULE; ORAL

WELCHOL

DAIICHI SANKYO

375MG

N021141 001 May 26, 2000

FOR SUSPENSION; ORAL

COLESEVELAM HYDROCHLORIDE

WATSON LABS INC

1.875GM/PACKET

A202178 001 Sep 01, 2020

3.75GM/PACKET

A202178 002 Sep 01, 2020

WELCHOL

+ DAIICHI SANKYO

1.875GM/PACKET **

N022362 001 Oct 02, 2009

TABLET; ORAL

COLESEVELAM HYDROCHLORIDE

INVAGEN PHARMS

625MG

A212602 001 Apr 20, 2020

WATSON LABS INC

625MG

A200830 001 Sep 02, 2020

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

ACTHAR GEL

MALLINCKRODT ARD

40 UNITS/ML

N008372 006

CORTICOTROPIN

ORGANICS LAGRANGE

40 UNITS/ML

N010831 001

80 UNITS/ML

N010831 002

WATSON LABS

40 UNITS/VIAL

A088772 001 Nov 21, 1984

PURIFIED CORTROPHIN GEL

ANI PHARMS

40 UNITS/ML

N008975 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CORTICOTROPIN-ZINC HYDROXIDE

INJECTABLE; INJECTION

CORTROPHIN-ZINC

ANI PHARMS

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

BAUSCH AND LOMB

10MG/ML

A075585 001 Dec 21, 2000

FERA PHARMS LLC

10MG/ML

A075437 001 Apr 21, 2000

HIKMA

10MG/ML

A075333 001 Apr 30, 2002

MYLAN SPECIALITY LP

10MG/ML

A074209 001 Apr 26, 1994

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CROMOLYN SODIUM

SOLUTION/DROPS;OPHTHALMIC

CROLOM					
BAUSCH AND LOMB	4%	A074443	001	Jan 30, 1995	
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	

CROTAMITON

CREAM;TOPICAL

EURAX					
+ JOURNEY	10%	N006927	001		

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		
DR REDDYS	0.1MG/ML	A080573	002		
	1MG/ML	A080573	001		
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	A083120	001		
	1MG/ML	A083120	002		
WYETH AYERST	0.1MG/ML	A080554	001		
	1MG/ML	A080554	002		
XIROMED	1MG/ML	A215046	001	Aug 20, 2021	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CYANOCOBALAMIN

INJECTABLE; INJECTION

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
	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
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
CYANOCOBALAMIN				
PADAGIS ISRAEL	0.5MG/SPRAY		A212458	001
				Sep 09, 2020
TABLET; ORAL				
CYANOCOBALAMIN				
WEST WARD	1MG		A084264	001

CYANOCOBALAMIN; CYANOCOBALAMIN CO-57; CYANOCOBALAMIN CO-58

N/A; N/A

DICOPAC KIT

GE HEALTHCARE	N/A; N/A; N/A		N017406	001
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CYANOCOBALAMIN; TANNIC ACID; ZINC ACETATE

INJECTABLE; INJECTION

DEPINAR

ARMOUR PHARM	0.5MG/ML; 2.3MG/ML; 1MG/ML		N011208	001
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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
	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
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CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

CHARTWELL RX	5MG		A072854	002
	10MG		A072854	001
				Feb 03, 2006
PLIVA	10MG		A074421	001
				Nov 19, 1991
RISING PHARMA	5MG		A073144	002
				Sep 29, 1995
	7.5MG		A073144	003
				Feb 03, 2006
	10MG		A073144	001
				Mar 25, 2013
	10MG		A073144	001
				May 30, 1991
SANDOZ	10MG		A073683	001
				Feb 26, 1993
WATSON LABS	10MG		A073143	001
				Nov 27, 1991
	10MG		A074436	001
				Nov 30, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

FLEXERIL

+	JANSSEN RES AND DEV	5MG **	N017821	001	
+		10MG **	N017821	002	

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AK-PENTOLATE

	AKORN	1%	A085555	001	
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CYCLOPENTOLATE HYDROCHLORIDE

	ALCON PHARMS LTD	1%	A089162	001	Jan 24, 1991
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	SOLA BARNES HIND	1%	A084150	001	
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		1%	A084863	001	
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PENTOLAIR

	PHARMAFAIR	0.5%	A088643	001	Feb 09, 1987
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		1%	A088150	001	Feb 25, 1983
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CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

	BAXTER HLTHCARE	100MG/VIAL	A088371	001	Jul 03, 1986
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		200MG/VIAL	A088372	001	Jul 03, 1986
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		500MG/VIAL	A088373	001	Jul 03, 1986
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		1GM/VIAL	A088374	001	Sep 24, 1986
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CYTOXAN

+	BAXTER HLTHCARE	100MG/VIAL **	N012142	001	
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+		200MG/VIAL **	N012142	002	
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CYTOXAN (LYOPHILIZED)

+	BAXTER HLTHCARE	500MG/VIAL	N012142	003	
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+		500MG/VIAL **	N012142	008	Jan 04, 1984
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+		1GM/VIAL	N012142	004	Aug 30, 1982
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+		1GM/VIAL **	N012142	010	Sep 24, 1985
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+		2GM/VIAL	N012142	005	Aug 30, 1982
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+		2GM/VIAL **	N012142	009	Dec 10, 1985
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LYOPHILIZED CYTOXAN

+	BAXTER HLTHCARE	100MG/VIAL **	N012142	006	Dec 05, 1985
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+		200MG/VIAL **	N012142	007	Dec 10, 1985
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NEOSAR

	BEDFORD	100MG/VIAL	A087442	001	Feb 16, 1982
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		200MG/VIAL	A087442	002	Feb 16, 1982
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		500MG/VIAL	A087442	003	Feb 16, 1982
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		1GM/VIAL	A087442	004	Jul 08, 1983
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		2GM/VIAL	A087442	005	Mar 30, 1989
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	TEVA PARENTERAL	100MG/VIAL	A040015	001	Apr 29, 1993
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		200MG/VIAL	A040015	002	Apr 29, 1993
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		500MG/VIAL	A040015	003	Apr 29, 1993
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		1GM/VIAL	A040015	004	Apr 29, 1993
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		2GM/VIAL	A040015	005	Apr 29, 1993
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TABLET; ORAL

CYCLOPHOSPHAMIDE

	ROXANE	25MG	A040032	001	Aug 17, 1999
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		50MG	A040032	002	Aug 17, 1999
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CYCLOSPORINE

CAPSULE; ORAL

GENGRAF

	ABBVIE	50MG	A065003	002	May 12, 2000
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NEORAL

+	NOVARTIS	50MG **	N050715	003	Jul 14, 1995
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SOLUTION; ORAL

CYCLOSPORINE

	PHARM ASSOC	100MG/ML	A065167	001	Jan 05, 2005
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CYCLOTHIAZIDE

TABLET; ORAL

ANHYDRON

	LILLY	2MG	N013157	002	
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FLUIDIL

	PHARMACIA AND UPJOHN	2MG	N018173	001	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
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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
	STRIDES PHARMA	4MG	A087129	001
	SUPERPHARM	4MG	A087405	001
	VITARINE	4MG	A087284	001
	WATSON LABS	4MG	A085245	001
		4MG	A086165	001
		4MG	A086580	001

PERIACTIN

+	MERCK	4MG **	N012649	001
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CYSTEINE HYDROCHLORIDE

INJECTABLE; INJECTION

CYSTEINE HYDROCHLORIDE

+	HOSPIRA	7.25% **	N019523	001 Oct 22, 1986
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SOLUTION; INTRAVENOUS

NOURESS

+	BAXTER HLTHCARE CORP	500MG/10ML (50MG/ML)	N212535	001 Dec 13, 2019
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CYTARABINE

INJECTABLE; INJECTION

CYTARABINE

	HONG KONG	20MG/ML	A206189	001 Jun 26, 2020
	MEITHEAL	20MG/ML	A206190	001 Nov 09, 2017
	MYLAN LABS LTD	20MG/ML	A200916	001 Dec 13, 2011
	RISING PHARMA	20MG/ML	A200914	001 Dec 13, 2011
+	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
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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

DACLATASVIR DIHYDROCHLORIDE

TABLET; ORAL

DAKLINZA

+ BRISTOL-MYERS SQUIBB

EQ 30MG BASE

N206843 001 Jul 24, 2015

+

EQ 60MG BASE

N206843 002 Jul 24, 2015

+

EQ 90MG BASE

N206843 003 Apr 13, 2016

DACTINOMYCIN

INJECTABLE; INJECTION

DACTINOMYCIN

AM REGENT

0.5MG/VIAL

A202562 001 Aug 23, 2013

HIKMA

0.5MG/VIAL

A090304 001 Mar 16, 2010

DALFAMPRIDINE

TABLET, EXTENDED RELEASE; ORAL

DALFAMPRIDINE

RISING PHARMA

10MG

A206858 001 Jul 06, 2020

DALFOPRISTIN; QUINUPRISTIN

INJECTABLE; INTRAVENOUS

SYNERCID

KING PHARMS

420MG/VIAL; 180MG/VIAL

N050748 002 Aug 24, 2000

DALTEPARIN SODIUM

INJECTABLE; INJECTION

FRAGMIN

PFIZER

7,500 IU/0.75ML

N020287 008 Apr 04, 2002

INJECTABLE; SUBCUTANEOUS

FRAGMIN

PFIZER

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

DANTROLENE SODIUM

INJECTABLE; INJECTION

DANTROLENE SODIUM

AUROMEDICS PHARMA

20MG/VIAL

A205239 001 Feb 18, 2016

DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

QTERNMET XR

+ ASTRAZENECA AB

2.5MG; 1GM; EQ 2.5MG BASE

N210874 001 May 02, 2019

+

5MG; 1GM; EQ 2.5MG BASE

N210874 002 May 02, 2019

+

5MG; 1GM; EQ 5MG BASE

N210874 003 May 02, 2019

+

10MG; 1GM; EQ 5MG BASE

N210874 004 May 02, 2019

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DAPIPIRAZOLE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

DAPIPIRAZOLE HYDROCHLORIDE

+	FERA PHARMS	0.5% **	N019849	001	Dec 31, 1990
	WOODWARD	0.5%	A204902	001	May 30, 2019

DAPSONE

TABLET;ORAL

DAPSONE

	TARO	25MG	A209430	001	Mar 01, 2019
		100MG	A209430	002	Mar 01, 2019

DAPTOMYCIN

POWDER;INTRAVENOUS

CUBICIN

	CUBIST PHARMS LLC	250MG/VIAL	N021572	001	Sep 12, 2003
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DAPTOMYCIN

	HISUN PHARM HANGZHOU	500MG/VIAL	A212250	001	Apr 21, 2021
	HOSPIRA INC	500MG/VIAL	A202857	001	Sep 12, 2014

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE;ORAL

DARIFENACIN HYDROBROMIDE

	ANCHEN PHARMS	EQ 7.5MG BASE	A091190	001	Mar 13, 2015
		EQ 15MG BASE	A091190	002	Mar 13, 2015
	JUBILANT GENERICS	EQ 7.5MG BASE	A205550	001	Oct 12, 2016
		EQ 15MG BASE	A205550	002	Oct 12, 2016

ENABLEX

+	APIL	EQ 7.5MG BASE	N021513	001	Dec 22, 2004
+		EQ 15MG BASE	N021513	002	Dec 22, 2004

DARUNAVIR

TABLET;ORAL

PREZISTA

+	JANSSEN PRODS	300MG **	N021976	001	Jun 23, 2006
+		400MG **	N021976	003	Oct 21, 2008

DASABUVIR SODIUM; OMBITASVIR, PARITAPREVIR, RITONAVIR

TABLET;ORAL

VIEKIRA PAK (COPACKAGED)

+	ABBVIE INC	EQ 250MG BASE;12.5MG, 75MG, 50MG	N206619	001	Dec 19, 2014
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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
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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
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DAUNORUBICIN HYDROCHLORIDE

INJECTABLE;INJECTION

CERUBIDINE

	SANOFI AVENTIS US	EQ 20MG BASE/VIAL	A061876	001	
	WYETH AYERST	EQ 20MG BASE/VIAL **	N050484	001	

DAUNORUBICIN HYDROCHLORIDE

	HISUN PHARM HANGZHOU	EQ 20MG BASE/VIAL	A206195	001	Apr 25, 2019
	TEVA PARENTERAL	EQ 20MG BASE/VIAL	A064212	001	Jun 23, 1998
		EQ 50MG BASE/VIAL	A064212	002	May 03, 1999

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DECAMETHONIUM BROMIDE

INJECTABLE; INJECTION

SYNCURINE

GLAXOSMITHKLINE

1MG/ML

N006931 002

DECITABINE

INJECTABLE; INTRAVENOUS

DECITABINE

CIPLA

50MG/VIAL

A208601 001 Nov 16, 2017

DEFERASIROX

TABLET; ORAL

DEFERASIROX

AMNEAL

90MG

A210727 001 Dec 27, 2019

360MG

A210727 002 Dec 27, 2019

TABLET, FOR SUSPENSION; ORAL

DEFERASIROX

ZYDUS PHARMS

125MG

A208882 001 May 05, 2020

250MG

A208882 002 May 05, 2020

500MG

A208882 003 May 05, 2020

DEFERIPRONE

SOLUTION; ORAL

FERRIPROX

+ CHIESI

80MG/ML

N208030 002 Apr 20, 2018

DEFEROXAMINE MESYLATE

INJECTABLE; INJECTION

DEFEROXAMINE MESYLATE

DR REDDYS

500MG/VIAL

A076806 001 Mar 31, 2006

2GM/VIAL

A076806 002 Mar 31, 2006

DESFERAL

+ NOVARTIS

2GM/VIAL

N016267 002 May 25, 2000

DELAVIRDINE MESYLATE

TABLET; ORAL

RESCRIPTOR

+ VIIV HLTHCARE

100MG

N020705 001 Apr 04, 1997

+

200MG

N020705 002 Jul 14, 1999

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

BARR

150MG

A065171 001 Dec 13, 2004

300MG

A065171 002 Dec 13, 2004

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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; METHYLCLOTHIAZIDE

TABLET; ORAL

ENDURONYL

ABBOTT

0.25MG; 5MG

N012775 001

ENDURONYL FORTE

ABBOTT

0.5MG; 5MG

N012775 002

METHYLCLOTHIAZIDE AND DESERPIDINE

WATSON LABS

0.25MG; 5MG

A088486 001 Aug 10, 1984

0.5MG; 5MG

A088452 001 Aug 10, 1984

DESIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

PERTOFRANE

SANOFI AVENTIS US

25MG

N013621 001

50MG

N013621 002

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

ANI PHARMS

10MG

A205153 001 Oct 28, 2016

25MG

A071803 002 Dec 08, 1987

25MG

A205153 002 Oct 28, 2016

50MG

A071803 003 Dec 08, 1987

50MG

A205153 003 Oct 28, 2016

75MG

A071803 004 Dec 08, 1987

75MG

A205153 004 Oct 28, 2016

100MG

A071803 001 May 29, 1997

100MG

A205153 005 Oct 28, 2016

150MG

A071803 005 May 29, 1997

150MG

A205153 006 Oct 28, 2016

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

DESLANOSIDE

INJECTABLE; INJECTION

CEDILANID-D

NOVARTIS

0.2MG/ML

N009282 002

DESLORATADINE

SOLUTION; ORAL

CLARINEX

+ MERCK SHARP DOHME

0.5MG/ML

N021300 001 Sep 01, 2004

DESLORATADINE

TARO

0.5MG/ML

A202936 001 May 26, 2016

TARO PHARM INDS LTD

0.5MG/ML

A202592 001 Jun 30, 2015

TABLET; ORAL

DESLORATADINE

MYLAN PHARMS INC

5MG

A078351 001 Feb 10, 2012

SUN PHARM INDS

5MG

A078359 001 Nov 16, 2010

TABLET, ORALLY DISINTEGRATING; ORAL

CLARINEX

+ MERCK SHARP DOHME

2.5MG

N021312 002 Jul 14, 2005

+

5MG

N021312 001 Jun 26, 2002

DESLORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

CLARINEX D 24 HOUR

+ MERCK SHARP DOHME

5MG; 240MG

N021605 001 Mar 03, 2005

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

DESMOPRESSIN ACETATE PRESERVATIVE FREE

BEDFORD 0.004MG/ML A074574 001 Feb 18, 2000

SOLUTION; NASAL

CONCENTRAID

FERRING 0.01% N019776 001 Dec 26, 1990

DDAVP

+ FERRING PHARMS INC 0.01% N017922 001

DESMOPRESSIN ACETATE

SUN PHARM INDS 0.01% A077212 001 Apr 12, 2012

SPRAY, METERED; NASAL

DDAVP

+ FERRING PHARMS INC 0.01MG/SPRAY ** N017922 002 Feb 06, 1989

DDAVP (NEEDS NO REFRIGERATION)

+ FERRING PHARMS INC 0.01MG/SPRAY N017922 003 Aug 07, 1996

MINIRIN

+ FERRING 0.01MG/SPRAY N021333 001 Sep 16, 2002

NOCTIVA

+ SERENITY PHARMS LLC 0.00083MG/SPRAY N201656 001 Mar 03, 2017

+ 0.00166MG/SPRAY N201656 002 Mar 03, 2017

STIMATE

FERRING PHARMS INC 0.15MG/SPRAY N020355 001 Mar 07, 1994

STIMATE (NEEDS NO REFRIGERATION)

+ FERRING PHARMS INC 0.15MG/SPRAY N020355 002 Oct 24, 2007

TABLET; ORAL

DESMOPRESSIN ACETATE

FERRING 0.1MG N021795 001 May 08, 2008

0.2MG N021795 002 May 08, 2008

IMPAX LABS INC 0.1MG A077122 001 Jan 25, 2006

0.2MG A077122 002 Jan 25, 2006

MYLAN 0.1MG A200653 001 Jun 27, 2014

0.2MG A200653 002 Jun 27, 2014

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

DESOGEN

ORGANON USA INC 0.15MG;0.03MG N020071 002 Dec 10, 1992

DESOGESTREL AND ETHINYL ESTRADIOL

MYLAN LABS LTD 0.15MG;0.03MG A202085 001 May 20, 2015

NAARI PTE LTD 0.15MG;0.03MG A207067 001 Sep 13, 2018

EMOQUETTE

VINTAGE PHARMS LLC 0.15MG;0.03MG A076675 001 Feb 25, 2011

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

DESONIDE

GEL; TOPICAL

DESONATE

+ LEO PHARMA AS 0.05% N021844 001 Oct 20, 2006

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DESOXIMETASONE

CREAM; TOPICAL

DESOXIMETASONE

AKORN

0.05%

A203787 001 Jan 06, 2017

0.25%

A203234 001 Jun 12, 2015

TOPICORT

+ TARO PHARM INDS LTD 0.25% **

N017856 001

TOPICORT LP

+ TARO PHARM INDS LTD 0.05% **

N018309 001

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%

A074286 001 Jun 07, 1996

+ 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

TABLET, EXTENDED RELEASE; ORAL

KHEDEZLA

OSMOTICA PHARM CORP 50MG

N204683 001 Jul 10, 2013

100MG

N204683 002 Jul 10, 2013

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

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

DESVENLAFAXINE SUCCINATE

MYLAN EQ 50MG BASE

A204095 001 Jun 29, 2015

EQ 100MG BASE

A204095 002 Jun 29, 2015

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

+ WOCKHARDT BIO AG 0.5MG/5ML

A088254 001 Jul 27, 1983

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXAMETHASONE

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

	BAUSCH	1.5MG	A040700 001	Aug 15, 2008
	HERITAGE PHARMA	1.5MG	A085456 001	
	IMPAX LABS	0.75MG	A085376 001	
	PHOENIX LABS NY	0.75MG	A083806 001	
	PRASCO	0.75MG	A080399 001	
	PVT FORM	0.75MG	A083420 001	
	ROXANE	0.25MG	A084614 001	
	STRIDES PHARMA	0.25MG	A088149 001	Apr 28, 1983
	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	
		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	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	
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AEROSOL, METERED; INHALATION

DEXACORT

	UCB INC	EQ 0.1MG PHOSPHATE/INH	N013413 001	
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CREAM; TOPICAL

DECADRON

	MERCK	EQ 0.1% PHOSPHATE	N011983 002	
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INJECTABLE; INJECTION

DECADRON

+	MERCK	EQ 4MG PHOSPHATE/ML **	N012071 002	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

DECADRON				
+	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
DR REDDYS	EQ 4MG PHOSPHATE/ML		A089169	001 Apr 09, 1986
INTL MEDICATION	EQ 20MG PHOSPHATE/ML		A088522	001 Feb 17, 1984
+	EQ 4MG PHOSPHATE/ML		A087440	001 Jul 21, 1982
LUITPOLD	EQ 4MG PHOSPHATE/ML		A087065	001
LYPHOMED	EQ 4MG PHOSPHATE/ML		A081125	001 Aug 31, 1990
TEVA PARENTERAL	EQ 10MG PHOSPHATE/ML		A081126	001 Aug 31, 1990
	EQ 4MG PHOSPHATE/ML		A083702	001
WATSON LABS	EQ 4MG PHOSPHATE/ML		A084355	001
	EQ 10MG PHOSPHATE/ML		A087668	001 Jul 01, 1982
	EQ 24MG PHOSPHATE/ML		A085606	001
WYETH AYERST	EQ 4MG PHOSPHATE/ML		A085641	001
HEXADROL				
+	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				
+	EQ 0.1% PHOSPHATE		N011984	001
DEXAMETHASONE SODIUM PHOSPHATE				
SANDOZ INC	EQ 0.1% PHOSPHATE		A088771	001 Jan 16, 1985
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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

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 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

DEXMEDETOMIDINE HYDROCHLORIDE

INJECTABLE;INJECTION

DEXMEDETOMIDINE HYDROCHLORIDE

AM REGENT EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML) A203773 001 May 12, 2017

SLAYBACK PHARMA LLC EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML) A212791 003 May 08, 2020

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

AUROLIFE PHARMA LLC 5MG A204266 001 Aug 25, 2015

10MG A204266 002 Aug 25, 2015

15MG A204266 003 Aug 25, 2015

20MG A204266 004 Dec 21, 2015

25MG A204266 005 May 04, 2021

30MG A202580 001 Aug 28, 2013

35MG A204266 006 May 04, 2021

40MG A204266 007 Aug 25, 2015

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXMETHYLPHENIDATE HYDROCHLORIDE

TABLET; ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

BIONPHARMA INC	5MG	A209754 001	Mar 24, 2020
	10MG	A209754 002	Mar 24, 2020
LANNETT CO INC	2.5MG	A209468 001	Sep 25, 2017
	5MG	A209468 002	Sep 25, 2017
	10MG	A209468 003	Sep 25, 2017
TEVA PHARMS	2.5MG	A077107 003	Jan 29, 2007
	5MG	A077107 001	Jan 29, 2007
	10MG	A077107 002	Jan 29, 2007

DEXRAZOXANE HYDROCHLORIDE

INJECTABLE; INJECTION

TOTECT

+ CLINIGEN EQ 500MG BASE/VIAL N022025 001 Sep 06, 2007

ZINECARD

+ PFIZER EQ 250MG BASE/VIAL N020212 001 May 26, 1995

+ EQ 500MG BASE/VIAL N020212 002 May 26, 1995

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
MYLAN	5MG	A206735 001	Jan 27, 2016
	10MG	A206735 002	Jan 27, 2016
	15MG	A206735 003	Jan 27, 2016
STRIDES PHARMA	5MG	A205077 001	Jun 21, 2019
	10MG	A205077 002	Jun 21, 2019
	15MG	A205077 003	Jun 21, 2019

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	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	
STRIDES PHARMA	5MG	A040299 001	May 13, 1999
VITARINE	5MG	A084986 001	
	10MG	A085892 001	
DEXTROSTAT			
+ SHIRE	5MG **	A084051 001	
+	10MG **	A084051 002	
FERNDEX			
FERNDALE LABS	5MG	A084001 001	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

AKORN 15MG/5ML; 6.25MG/5ML A040027 001 Jul 31, 1996

+ ANI PHARMS 15MG/5ML; 6.25MG/5ML ** N011265 002 Apr 02, 1984

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 70% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 70GM/100ML ** N017521 006 Mar 26, 1982

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

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075%				
FRESENIUS KABI USA	5GM/100ML; 75MG/100ML	A212346	001	Sep 10, 2020
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15%				
FRESENIUS KABI USA	5GM/100ML; 150MG/100ML	A212346	002	Sep 10, 2020
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
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER				
ICU MEDICAL INC	5GM/100ML; 224MG/100ML	N018371	003	
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER				
ICU MEDICAL INC	5GM/100ML; 298MG/100ML	N018371	002	

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%				
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

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 0.225% IN PLASTIC CONTAINER			
FRESENIUS KABI USA	5GM/100ML;149MG/100ML;225MG/100ML	A212348 001	Jul 30, 2021
+ ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;225MG/100ML **	N018365 002	Jul 05, 1983
+ ICU MEDICAL INC	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
ICU MEDICAL INC	5GM/100ML;149MG/100ML;300MG/100ML	N018876 006	Mar 28, 1988
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%			
FRESENIUS KABI USA	5GM/100ML;74.5MG/100ML;450MG/100ML	A213523 001	Mar 09, 2021
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 0.225% 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 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 0.225% IN PLASTIC CONTAINER			
FRESENIUS KABI USA	5GM/100ML;149MG/100ML;225MG/100ML	A212348 002	Jul 30, 2021
+ ICU MEDICAL INC	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 0.45%			
FRESENIUS KABI USA	5GM/100ML;149MG/100ML;450MG/100ML	A213523 002	Mar 09, 2021
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%			
FRESENIUS KABI USA	5GM/100ML;149MG/100ML;900MG/100ML	A213445 001	Mar 09, 2021
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 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 0.45%			
FRESENIUS KABI USA	5GM/100ML;224MG/100ML;450MG/100ML	A213523 003	Mar 09, 2021
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 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%				
FRESENIUS KABI USA	5GM/100ML;298MG/100ML;450MG/100ML	A213523	004	Mar 09, 2021
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9%				
FRESENIUS KABI USA	5GM/100ML;298MG/100ML;900MG/100ML	A213445	002	Mar 09, 2021
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				
+ 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	
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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	
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	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

CONCENTRATE;ORAL

DIAZEPAM

LANNETT CO INC 5MG/ML A204433 001 Apr 14, 2014

GEL;RECTAL

DIASTAT

+ BAUSCH 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

HIKMA 5MG/ML A070311 001 Dec 16, 1985

5MG/ML A070312 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

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

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

BARR 2MG A070152 001 Nov 01, 1985

10MG A070154 001 Nov 01, 1985

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

FERNDALE LABS 2MG A070903 001 Apr 01, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIAZEPAM

TABLET; ORAL

DIAZEPAM

	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
NUVO PHARM	10MG	A070464 001	Feb 25, 1986
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
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
HYPERSTAT			
SCHERING	15MG/ML	N016996 001	

DIBUCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

HEAVY SOLUTION NUPERCAINE

NOVARTIS	2.5MG/ML	N006203 001	
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DICHLORPHENAMIDE

TABLET; ORAL

DARANIDE

+ XERIS	50MG **	N011366 001	
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DICLOFENAC

CAPSULE; ORAL

ZORVOLEX

+ ZYLA	18MG	N204592 001	Oct 18, 2013
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DICLOFENAC POTASSIUM

CAPSULE; ORAL

DICLOFENAC POTASSIUM

STRIDES PHARMA	25MG	A210078 001	Dec 03, 2019
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

GEL; TOPICAL

SOLARAZE

+ FOUGERA PHARMS	3%	N021005 001	Oct 16, 2000
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SOLUTION; INTRAVENOUS

DICLOFENAC SODIUM

RISING PHARMA	37.5MG/ML (37.5MG/ML)	A208786 001	Jun 18, 2019
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DYLOJECT

+ JAVELIN PHARMS INC	37.5MG/ML (37.5MG/ML) **	N022396 001	Dec 23, 2014
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SOLUTION; TOPICAL

DICLOFENAC SODIUM

APOTEX INC	1.5%	A202027 001	May 27, 2014
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RISING	1.5%	A206715 001	Aug 07, 2017
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PENNSAID

+ NUVO PHARMS INC	1.5% **	N020947 001	Nov 04, 2009
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SOLUTION/DROPS; OPHTHALMIC

DICLOFENAC SODIUM

FALCON PHARMS	0.1%	N020809 001	May 04, 1998
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RUBICON	0.1%	A077600 001	Nov 13, 2008
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VOLTAREN

+ NOVARTIS	0.1%	N020037 001	Mar 28, 1991
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TABLET, DELAYED RELEASE; ORAL

DICLOFENAC SODIUM

AUROBINDO PHARMA USA	50MG	A075281 002	Feb 12, 2002
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	75MG	A075281 003	Feb 12, 2002
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CHARTWELL RX	25MG	A074376 001	Sep 28, 1995
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	50MG	A074376 002	Sep 28, 1995
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	75MG	A074394 001	Nov 30, 1995
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MICRO LABS	50MG	A074986 001	Feb 26, 1999
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	75MG	A074986 002	Feb 26, 1999
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PLIVA	50MG	A074432 002	Jul 29, 1999
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	75MG	A074432 003	Jul 29, 1999
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ROXANE	25MG	A074391 001	Jun 29, 1995
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	50MG	A074391 002	Jun 29, 1995
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	75MG	A074391 003	Jun 29, 1995
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TEVA	50MG	A074723 001	Mar 30, 1999
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	75MG	A074390 001	Aug 15, 1996
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TEVA PHARMS	25MG	A074459 001	Jun 25, 1997
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	50MG	A074459 002	Jun 25, 1997
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	75MG	A074459 003	Jun 25, 1997
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VOLTAREN

+ NOVARTIS	25MG **	N019201 001	Jul 28, 1988
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+	50MG **	N019201 002	Jul 28, 1988
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+	75MG **	N019201 003	Jul 28, 1988
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TABLET, EXTENDED RELEASE; ORAL

DICLOFENAC SODIUM

ACTAVIS ELIZABETH	100MG	A075910 001	Jan 07, 2002
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MYLAN	100MG	A076152 001	Dec 13, 2001
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VOLTAREN-XR

+ NOVARTIS	100MG **	N020254 001	Mar 08, 1996
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DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE; ORAL

DICLOFENAC SODIUM AND MISOPROSTOL

EXELA HOLDINGS	50MG; 0.2MG	A200540 001	Mar 14, 2014
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	75MG; 0.2MG	A200540 002	Mar 14, 2014
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DICLOXACILLIN SODIUM

CAPSULE; ORAL

DICLOXACILLIN SODIUM

SANDOZ

EQ 125MG BASE

A061454 002

EQ 250MG BASE

A061454 001

EQ 500MG BASE

A061454 003

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

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

BENTYL

+ ALLERGAN

10MG **

N007409 003 Oct 15, 1984

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

DR REDDYS

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

BENTYL

+ ALLERGAN

20MG **

N007409 001 Oct 15, 1984

DICYCLOMINE HYDROCHLORIDE

AUROBINDO PHARMA USA

20MG

A040317 001 Sep 07, 1999

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

AUROBINDO PHARMA

125MG

A090094 001 Sep 24, 2008

200MG

A090094 002 Sep 24, 2008

250MG

A090094 003 Sep 24, 2008

400MG

A090094 004 Sep 24, 2008

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIRIDANOSINE

CAPSULE, DELAYED REL PELLETS;ORAL

DIRIDANOSINE

400MG

A090788 004 Apr 08, 2010

VIDEX EC

+ BRISTOL MYERS SQUIBB 125MG

N021183 001 Oct 31, 2000

+ 200MG

N021183 002 Oct 31, 2000

+ 250MG

N021183 003 Oct 31, 2000

+ 400MG

N021183 004 Oct 31, 2000

FOR SOLUTION;ORAL

DIRIDANOSINE

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

+ BRISTOL-MYERS SQUIBB 10MG/ML

N020156 001 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

DIRIDANOSINE

AUROBINDO 100MG

A077275 001 Aug 14, 2012

150MG

A077275 002 Aug 14, 2012

200MG

A077275 003 Aug 14, 2012

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

+ NOSTRUM LABS INC 25MG

N011722 002

SANOFI AVENTIS US 25MG

N017668 001

TEPANIL

3M 25MG

N011673 001

TABLET, EXTENDED RELEASE;ORAL

TENUATE

SANOFI AVENTIS US 75MG

N017669 001

TENUATE DOSPAN

+ NOSTRUM LABS INC 75MG

N012546 001

TEPANIL TEN-TAB

3M 75MG

N017956 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
	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
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TABLET; ORAL

STILPHOSTROL

BAYER PHARMS	50MG	N010010 002
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DIFLORASONE DIACETATE

CREAM; TOPICAL

DIFLORASONE DIACETATE

FOUGERA PHARMS	0.05%	A075187 001	Mar 30, 1998
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FLORONE

PFIZER	0.05% **	N017741 001
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FLORONE E

PFIZER	0.05%	N019259 001	Aug 28, 1985
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PSORCON

+ TARO PHARMS NORTH	0.05% **	N020205 001	Nov 20, 1992
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OINTMENT; TOPICAL

DIFLORASONE DIACETATE

AKORN	0.05%	A206572 001	Jul 24, 2015
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PSORCON

+ PFIZER	0.05%	N019260 001	Aug 28, 1985
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIFLORASONE DIACETATE

OINTMENT; TOPICAL

PSORCON E

PFIZER

0.05%

N017994 001

DIFLUNISAL

TABLET; ORAL

DIFLUNISAL

ANI PHARMS

500MG

A074604 001 Jun 10, 1996

DASTECH INTL

250MG

A073562 001 Nov 27, 1992

PUREPAC PHARM

500MG

A073563 001 Nov 27, 1992

PUREPAC PHARM

250MG

A074285 001 May 07, 1996

PUREPAC PHARM

500MG

A074285 002 May 07, 1996

TEVA

250MG

A073679 001 Jul 31, 1992

WATSON LABS

250MG

A074400 001 Jul 17, 1997

WATSON LABS

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

HOSPIRA

0.25MG/ML

A040206 001 Aug 28, 1998

WYETH AYERST

0.25MG/ML

A084386 001

DIGOXIN PEDIATRIC

HOSPIRA

0.1MG/ML

A040092 001 Apr 25, 1996

TABLET; ORAL

LANOXIN

+ CONCORDIA

0.1875MG

N020405 003 Sep 30, 1997

0.375MG

N020405 005 Sep 30, 1997

0.5MG

N020405 006 Sep 30, 1997

DIHYDROERGOTAMINE MESYLATE; HEPARIN SODIUM; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

EMBOLEX

NOVARTIS

0.5MG/0.5ML; 2,500

N018885 001 Nov 30, 1984

UNITS/0.5ML; 5.33MG/0.5ML

0.5MG/0.7ML; 5,000

N018885 002 Nov 30, 1984

UNITS/0.7ML; 7.46MG/0.7ML

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CARDIZEM SR

+ BIOVAIL

60MG **

N019471 001 Jan 23, 1989

+

90MG **

N019471 002 Jan 23, 1989

+

120MG **

N019471 003 Jan 23, 1989

+

180MG **

N019471 004 Jan 23, 1989

DILACOR XR

+ ALLERGAN

120MG **

N020092 001 May 29, 1992

+

180MG **

N020092 002 May 29, 1992

+

240MG **

N020092 003 May 29, 1992

DILT-CD

APOTEX

120MG

A076151 001 May 20, 2004

APOTEX

180MG

A076151 002 May 20, 2004

APOTEX

240MG

A076151 003 May 20, 2004

APOTEX

300MG

A076151 004 May 20, 2004

DILTIAZEM HYDROCHLORIDE

ACTAVIS LABS FL INC

120MG

A074852 001 Oct 10, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DILTIAZEM HYDROCHLORIDECAPSULE, EXTENDED RELEASE;ORAL
DILTIAZEM HYDROCHLORIDE

	180MG		A074852	002	Oct 10, 1997
	240MG		A074852	003	Oct 10, 1997
BIOVAIL	60MG		A074845	001	Sep 15, 1999
	90MG		A074845	002	Sep 15, 1999
	120MG		A074845	003	Sep 15, 1999
	120MG		N020939	001	Jan 28, 2000
	180MG		N020939	002	Jan 28, 2000
	240MG		N020939	003	Jan 28, 2000
	300MG		N020939	004	Jan 28, 2000
	360MG		N020939	005	Sep 14, 2001
	420MG		N020939	006	Sep 14, 2001
MYLAN	120MG		A075124	002	Mar 18, 1998
	180MG		A075124	003	Mar 18, 1998
	240MG		A075124	001	Mar 18, 1998
NESHER PHARMS	120MG		A076563	002	Sep 12, 2006
	180MG		A076563	003	Sep 12, 2006
	240MG		A076563	004	Sep 12, 2006
	300MG		A076563	005	Sep 12, 2006
	360MG		A076563	006	Sep 12, 2006
	420MG		A076563	001	Sep 12, 2006
PAR PHARM	360MG		A209766	001	May 30, 2018
TEVA	60MG		A074079	001	Nov 30, 1993
	90MG		A074079	002	Nov 30, 1993
	120MG		A074079	003	Nov 30, 1993
DILTZAC					
APOTEX	120MG		A076395	001	Feb 01, 2006
	180MG		A076395	002	Feb 01, 2006
	240MG		A076395	003	Feb 01, 2006
	300MG		A076395	004	Feb 01, 2006
	360MG		A076395	005	Feb 01, 2006
INJECTABLE; INJECTION					
CARDIZEM					
BIOVAIL	100MG/VIAL **		N020792	001	Sep 05, 1997
+ BIOVAIL LABS INTL	5MG/ML **		N020027	001	Oct 24, 1991
+	25MG/VIAL **		N020027	003	Aug 18, 1995
DILTIAZEM HYDROCHLORIDE					
DR REDDYS	5MG/ML		A074894	001	Aug 26, 1997
HOSPIRA	5MG/ML		A075004	001	Feb 16, 2000
	5MG/ML		A075106	001	Apr 29, 1999
INTL MEDICATION	5MG/ML		A075749	001	Nov 21, 2001
MYLAN LABS LTD	5MG/ML		A075375	001	Sep 30, 1999
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
RISING PHARMA	30MG		A072838	004	Nov 05, 1992
	60MG		A072838	003	Nov 05, 1992
	90MG		A072838	002	Nov 05, 1992
	120MG		A072838	001	Nov 05, 1992
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DILTIAZEM MALATETABLET, 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 MALEATETABLET, EXTENDED RELEASE;ORAL
TECZEM

BIOVAIL	EQ 180MG HYDROCHLORIDE;5MG	N020507 001	Oct 04, 1996
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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	
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TABLET; ORAL

DIMENHYDRINATE

HEATHER	50MG	A080841 001	
NEXGEN PHARMA INC	50MG	A085985 001	
+ WATSON LABS	50MG	A085166 001	

DIMETHYL FUMARATE

CAPSULE, DELAYED RELEASE;ORAL

DIMETHYL FUMARATE

ZYDUS PHARMS	120MG	A210538 001	Sep 24, 2020
	240MG	A210538 002	Sep 24, 2020

DIMETHYL SULFOXIDE

SOLUTION; INTRAVESICAL

DIMETHYL SULFOXIDE

MYLAN INSTITUTIONAL	50%	A076185 001	Nov 29, 2002
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DIMYRISTOYL LECITHIN; PERFLEXANE

INJECTABLE; INTRAVENOUS

IMAGENT

VESSELON SPV LLC	0.92MG/VIAL;0.092MG/VIAL	N021191 001	May 31, 2002
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DINOPROST TROMETHAMINE

INJECTABLE; INJECTION

PROSTIN F2 ALPHA

PHARMACIA AND UPJOHN	EQ 5MG BASE/ML	N017434 001	
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DINOPROSTONE

SUPPOSITORY; VAGINAL

PROSTIN E2

+ PFIZER	20MG	N017810 001	
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DIPHEMANIL METHYLSULFATE

TABLET; ORAL

PRANTAL

SCHERING	100MG	N008114 004	
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DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

BENADRYL

MCNEIL CONS	25MG	N005845 007	
	50MG	N005845 001	

DIPHENHYDRAMINE HYDROCHLORIDE

ALRA	25MG	A080519 004	
	50MG	A080519 003	
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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

HALSEY	50MG	A087914	001	Jun 04, 1984
HEATHER	25MG	A084524	001	
	50MG	A083953	001	
HERITAGE PHARMA	50MG	A080727	001	
	50MG	A080738	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	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
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BENADRYL

MCNEIL CONS	12.5MG/5ML	N005845	004	
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DIBENIL

CENCI	12.5MG/5ML	A088304	001	Dec 16, 1983
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DIPHEN

USL PHARMA	12.5MG/5ML	A084640	001	
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DIPHENHYDRAMINE HYDROCHLORIDE

BUNDY	12.5MG/5ML	A083674	001	
CENCI	12.5MG/5ML	A087941	001	Dec 17, 1982
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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIPHENHYDRAMINE HYDROCHLORIDE

ELIXIR; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

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	
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INJECTABLE; INJECTION

BENADRYL

MCNEIL CONS	10MG/ML	N006146	001	
+	50MG/ML **	N006146	002	

BENADRYL PRESERVATIVE FREE

+	MCNEIL CONS	50MG/ML **	N009486	001
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DIPHENHYDRAMINE HYDROCHLORIDE

BEL MAR	10MG/ML	A080822	001	
DR REDDYS	10MG/ML	A080873	001	
	50MG/ML	A080873	002	
EUROHLTH INTL SARL	50MG/ML	A083183	001	
LYPHOMED	10MG/ML	A087066	001	
WATSON LABS	10MG/ML	A083533	001	
WYETH AYERST	50MG/ML	A080577	001	

DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE

ABRAXIS PHARM	50MG/ML	A080586	002	
DR REDDYS	50MG/ML	A080873	003	
INTL MEDICATION	50MG/ML	A084094	001	

SYRUP; ORAL

ANTITUSSIVE

PERRIGO	12.5MG/5ML	A071292	001	Apr 10, 1987
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BELDIN

HALSEY	12.5MG/5ML	A089179	001	Jun 05, 1986
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BENYLIN

PARKE DAVIS	12.5MG/5ML	N006514	004	
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DIPHEN

MORTON GROVE	12.5MG/5ML	A070118	001	Oct 01, 1985
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DIPHENHYDRAMINE HYDROCHLORIDE

AKORN	12.5MG/5ML	A072416	001	Sep 28, 1990
ALPHARMA US PHARMS	12.5MG/5ML	A070497	001	Apr 25, 1989
CUMBERLAND SWAN	12.5MG/5ML	A073611	001	Aug 20, 1992

HYDRAMINE

ALPHARMA US PHARMS	12.5MG/5ML	A070205	001	Jan 28, 1986
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SILPHEN

LANNETT CO INC	12.5MG/5ML	A072646	001	Feb 27, 1992
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VICKS FORMULA 44

WARNER CHILCOTT	12.5MG/5ML	A070524	001	Jan 14, 1987
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DIPHENHYDRAMINE HYDROCHLORIDE; NAPROXEN SODIUM

TABLET; ORAL

NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE

P AND L	25MG; 220MG	A207597	001	Jan 25, 2019
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DIPHENHYDRAMINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

BENYLIN

PARKE DAVIS	12.5MG/5ML; 30MG/5ML	N019014	001	Jun 11, 1985
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DIPHENIDOL HYDROCHLORIDE

TABLET; ORAL

VONTROL

GLAXOSMITHKLINE	EQ 25MG BASE	N016033	001	
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DIPHENYLPYRALINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

HISPRIL

GLAXOSMITHKLINE	5MG	N011945	001	
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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

AUROMEDICS PHARMA 5MG/ML A075769 001 Nov 27, 2002

DR REDDYS 5MG/ML A074952 001 Nov 26, 1997

HOSPIRA 5MG/ML A074601 001 Dec 19, 1997

IV PERSANTINE

+ BOEHRINGER INGELHEIM 5MG/ML ** N019817 001 Dec 13, 1990

TABLET; ORAL

DIPYRIDAMOLE

GENUS 25MG A040898 001 Apr 23, 2008

50MG A040898 002 Apr 23, 2008

75MG A040898 003 Apr 23, 2008

GLENMARK GENERICS 25MG A088999 001 Feb 05, 1991

50MG A089000 001 Feb 05, 1991

75MG A089001 001 Feb 05, 1991

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

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

MAYNE PHARMA EQ 100MG BASE A070173 001 May 31, 1985

EQ 150MG BASE A070173 002 May 31, 1985

MYLAN EQ 100MG BASE A070138 001 Jun 14, 1985

EQ 150MG BASE A070139 001 Jun 14, 1985

RISING EQ 100MG BASE A070470 001 Dec 10, 1985

EQ 150MG BASE A070471 001 Dec 10, 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

+ ODYSSEY PHARMS 250MG A088482 001 Dec 08, 1983

+ 500MG A088483 001 Dec 08, 1983

+ TEVA WOMENS 250MG ** N007883 003

+ 500MG ** N007883 002

DISULFIRAM

MYLAN 250MG A203916 001 Mar 04, 2015

500MG A203916 002 Mar 04, 2015

STRIDES PHARMA 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 effectiveness 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

CAPSULE, DELAYED REL PELLETS;ORAL

DIVALPROEX SODIUM

MYLAN	EQ 125MG VALPROIC ACID	A090407	001	Mar 28, 2011
TEVA PHARMS USA	EQ 125MG VALPROIC ACID	A211505	001	Nov 17, 2020

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

ANCHEN PHARMS	EQ 500MG VALPROIC ACID	A078411	001	Nov 03, 2008
MYLAN	EQ 125MG VALPROIC ACID	A077254	001	Jul 29, 2008
	EQ 125MG VALPROIC ACID	A090062	001	Mar 17, 2009
	EQ 250MG VALPROIC ACID	A077254	002	Jul 29, 2008
	EQ 250MG VALPROIC ACID	A090062	002	Mar 17, 2009
	EQ 500MG VALPROIC ACID	A077254	003	Jul 29, 2008
	EQ 500MG VALPROIC ACID	A090062	003	Mar 17, 2009
TEVA	EQ 125MG VALPROIC ACID	A076941	001	Jul 29, 2008
	EQ 250MG VALPROIC ACID	A076941	002	Jul 29, 2008
	EQ 500MG VALPROIC ACID	A076941	003	Jul 29, 2008

TABLET, EXTENDED RELEASE;ORAL

DIVALPROEX SODIUM

AMTA	EQ 250MG VALPROIC ACID	A214462	001	Mar 15, 2021
	EQ 500MG VALPROIC ACID	A214462	002	Mar 15, 2021
ANCHEN PHARMS	EQ 250MG VALPROIC ACID	A078445	001	Feb 26, 2009
	EQ 500MG VALPROIC ACID	A078445	002	Aug 04, 2009
COSETTE	EQ 500MG VALPROIC ACID	A078700	001	Aug 03, 2009
IMPAX LABS	EQ 250MG VALPROIC ACID	A078791	001	May 06, 2009
	EQ 500MG VALPROIC ACID	A078791	002	Aug 04, 2009

DOBUTAMINE HYDROCHLORIDE

INJECTABLE;INJECTION

DOBUTAMINE HYDROCHLORIDE

BAXTER HLTHCARE	EQ 12.5MG BASE/ML	A074381	001	Sep 26, 1996
DR REDDYS	EQ 12.5MG BASE/ML	A074995	001	Mar 31, 1998
HOSPIRA	EQ 12.5MG BASE/ML	A074292	001	Feb 16, 1995
	EQ 1.25GM BASE/100ML	A074634	001	Sep 27, 1996
LUITPOLD	EQ 12.5MG BASE/ML	A074545	001	Jun 25, 1998
TELIGENT	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

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 PHARM

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

DFB ONCOLOGY LTD

20MG/ML (20MG/ML) A206177 001 Jan 20, 2017

80MG/4ML (20MG/ML) A206177 002 Jan 20, 2017

200MG/10ML (20MG/ML) A206177 003 Jan 20, 2017

+ HOSPIRA INC

120MG/6ML (20MG/ML) N022234 006 Jun 23, 2016

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOCETAXEL

INJECTABLE; INJECTION

DOCETAXEL

INGENUS PHARMS LLC	20MG/2ML (10MG/ML)	A207563	001	Aug 31, 2017
JIANGSU PHARMS	40MG/ML	A203170	001	Feb 15, 2017
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
TEVA PHARMS USA	20MG/ML (20MG/ML)	A203877	001	Sep 16, 2015
	80MG/4ML (20MG/ML)	A203877	002	Sep 16, 2015

TAXOTERE

+ SANOFI AVENTIS US	40MG/ML **	N020449	001	May 14, 1996
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DOFETILIDE

CAPSULE; ORAL

DOFETILIDE

NOVAST LABS	0.125MG	A212410	001	Dec 27, 2019
	0.25MG	A212410	002	Dec 27, 2019
	0.5MG	A212410	003	Dec 27, 2019
PRINSTON INC	0.125MG	A211223	001	Dec 17, 2019
	0.25MG	A211223	002	Dec 17, 2019
	0.5MG	A211223	003	Dec 17, 2019

DOLASETRON MESYLATE

INJECTABLE; INJECTION

ANZEMET

+ VALIDUS PHARMS	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

+ VALIDUS PHARMS	100MG	N020623	002	Sep 11, 1997
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DONEPEZIL HYDROCHLORIDE

SOLUTION; ORAL

ARICEPT

EISAI INC	5MG/5ML	N021719	001	Oct 18, 2004
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TABLET; ORAL

DONEPEZIL HYDROCHLORIDE

ACCORD HLTHCARE	5MG	A201335	001	Aug 29, 2011
	10MG	A201335	002	Aug 29, 2011
ACTAVIS ELIZABETH	23MG	A202415	001	Dec 17, 2015
APOTEX	5MG	A078841	001	Jun 02, 2011
	10MG	A078841	002	Jun 02, 2011
HERITAGE PHARMA	5MG	A077344	001	May 31, 2011
	10MG	A077344	002	May 31, 2011
HIKMA PHARMS	5MG	A090247	001	May 31, 2011
	10MG	A090247	002	May 31, 2011
HISUN PHARM HANGZHOU	23MG	A202410	001	Mar 24, 2017
MYLAN PHARMS INC	5MG	A090521	001	May 31, 2011
	10MG	A090521	002	May 31, 2011
OSMOTICA PHARM US	23MG	A203114	001	Jan 26, 2016
PAR PHARM	23MG	A202542	001	Jul 24, 2013
RISING PHARMA	23MG	A202656	001	Oct 22, 2015
SUN PHARM	23MG	A204293	001	Jun 05, 2015
SUN PHARM INDS LTD	5MG	A076786	001	Nov 26, 2010
	10MG	A076786	002	Nov 26, 2010
UNICHEM	5MG	A203656	001	Jun 23, 2016
	10MG	A203656	002	Jun 23, 2016
WOCKHARDT	5MG	A091267	001	May 31, 2011
	10MG	A091267	002	May 31, 2011

TABLET, ORALLY DISINTEGRATING; ORAL

ARICEPT ODT

+ EISAI INC	5MG	N021720	001	Oct 18, 2004
+	10MG	N021720	002	Oct 18, 2004

DONEPEZIL HYDROCHLORIDE

HERITAGE PHARMA	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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DONEPEZIL HYDROCHLORIDE

TABLET, ORALLY DISINTEGRATING;ORAL

DONEPEZIL HYDROCHLORIDE

UNICHEM	5MG	A204831	001	Nov 10, 2016
	10MG	A204831	002	Nov 10, 2016
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
AM REGENT	40MG/ML	A070799	001	Feb 11, 1987
	80MG/ML	A070820	001	Feb 11, 1987
	160MG/ML	A070826	001	Feb 11, 1987
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	
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	

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

AM REGENT	EQ 2% BASE	A079186	001	Nov 18, 2009
RUBICON	EQ 2% BASE	A078395	001	Oct 28, 2008
TEVA PHARMS	EQ 2% BASE	A078756	001	Dec 04, 2008
ZAMBON SPA	EQ 2% BASE	A091034	001	Dec 04, 2013

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS;OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE

LANNETT CO INC	EQ 2% BASE;EQ 0.5% BASE	A201998	001	Dec 17, 2014
RUBICON	EQ 2% BASE;EQ 0.5% BASE	A078201	001	Oct 28, 2008
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
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DOXAPRAM HYDROCHLORIDE

INJECTABLE; INJECTION

DOXAPRAM HYDROCHLORIDE

WATSON LABS	20MG/ML	A073529	001	Jan 30, 1992
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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
ANI PHARMS	EQ 1MG BASE	A075432	001	Oct 18, 2000
	EQ 2MG BASE	A075432	002	Oct 18, 2000
	EQ 4MG BASE	A075432	003	Oct 18, 2000
	EQ 8MG BASE	A075432	004	Oct 18, 2000
AUROBINDO PHARMA USA	EQ 1MG BASE	A075509	001	Oct 19, 2000
	EQ 2MG BASE	A075509	002	Oct 19, 2000
	EQ 4MG BASE	A075509	003	Oct 19, 2000
	EQ 8MG BASE	A075509	004	Oct 19, 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
STRIDES PHARMA	EQ 1MG BASE	A076161	001	Jun 10, 2004
	EQ 2MG BASE	A076161	002	Jun 10, 2004
	EQ 4MG BASE	A076161	003	Jun 10, 2004
	EQ 8MG BASE	A076161	004	Jun 10, 2004
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

CHARTWELL RX	EQ 10MG BASE	A210268	001	Sep 04, 2020
	EQ 25MG BASE	A210268	002	Sep 04, 2020
	EQ 50MG BASE	A210268	003	Sep 04, 2020
	EQ 75MG BASE	A210268	004	Sep 04, 2020
	EQ 100MG BASE	A210268	005	Sep 04, 2020
DAVA PHARMS INC	EQ 10MG BASE	A071685	001	Jan 05, 1988
	EQ 25MG BASE	A071686	001	Jan 05, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIN HYDROCHLORIDE

	EQ 50MG BASE	A071673	001	Jan 05, 1988
	EQ 75MG BASE	A071674	001	Jan 05, 1988
	EQ 100MG BASE	A071675	001	Jan 05, 1988
	EQ 150MG BASE	A071676	001	Jan 05, 1988
LANNETT CO INC	EQ 10MG BASE	A212997	001	Jul 24, 2020
	EQ 25MG BASE	A212997	002	Jul 24, 2020
	EQ 50MG BASE	A212997	003	Jul 24, 2020
	EQ 75MG BASE	A212997	004	Jul 24, 2020
	EQ 100MG BASE	A212997	005	Jul 24, 2020
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
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
TEVA PHARMS	EQ 10MG BASE/ML	A071609	001	Nov 09, 1987
SINEQUAN				
+ PFIZER	EQ 10MG BASE/ML **	N017516	001	
TABLET; ORAL				
DOXEPIN HYDROCHLORIDE				
STRIDES ARCOLAB LTD	EQ 3MG BASE	A202510	001	Jul 24, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXEPIN HYDROCHLORIDE

TABLET; ORAL

DOXEPIN HYDROCHLORIDE

EQ 6MG BASE

A202510 002 Jul 24, 2020

DOXERCALCIFEROL

CAPSULE; ORAL

DOXERCALCIFEROL

HIKMA

0.5MCG

A091433 001 Sep 23, 2011

1MCG

A091433 002 Jan 14, 2014

2.5MCG

A091433 003 Jan 14, 2014

INJECTABLE; INJECTION

DOXERCALCIFEROL

AMNEAL

4MCG/2ML (2MCG/ML)

A208975 001 May 24, 2017

+ HOSPIRA INC

10MCG/5ML (2MCG/ML)

N208614 002 Jul 24, 2018

SUN PHARM

2MCG/ML (2MCG/ML)

A203875 001 Nov 14, 2019

4MCG/2ML (2MCG/ML)

A203875 002 Nov 14, 2019

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ADRIAMYCIN PFS

PFIZER

2MG/ML

A063165 001 Jan 30, 1991

200MG/100ML

A063165 002 Jan 30, 1991

DOXORUBICIN HYDROCHLORIDE

ALMAJECT

2MG/ML

A065515 001 Nov 08, 2012

HISUN PHARM HANGZHOU

20MG/VIAL

A206062 001 May 13, 2019

HLTHCARE

2MG/ML

A200146 001 Jul 18, 2012

MYLAN LABS LTD

2MG/ML

A200901 001 Feb 14, 2012

10MG/VIAL

A200170 001 Oct 28, 2011

PFIZER

10MG/VIAL

N050467 001

20MG/VIAL

N050467 003 May 20, 1985

50MG/VIAL

N050467 002

150MG/VIAL

N050467 004 Jul 22, 1987

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

MYLAN PHARMS INC

EQ 150MG BASE

A202778 001 Jun 08, 2012

RISING PHARMA

EQ 50MG BASE

A208942 001 Jan 21, 2021

EQ 75MG BASE

A208942 002 Jan 21, 2021

EQ 100MG BASE

A208942 003 Jan 21, 2021

SANDOZ INC

EQ 50MG BASE

A065032 001 Jun 30, 2000

EQ 100MG BASE

A065032 002 Jun 30, 2000

STRIDES PHARMA

EQ 75MG BASE

A065055 004 Apr 18, 2005

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

MYLAN

EQ 50MG BASE

A065377 001 Nov 07, 2006

EQ 75MG BASE

A065377 002 Nov 07, 2006

EQ 100MG BASE

A065377 003 Nov 07, 2006

EQ 150MG BASE

A065427 001 Jun 07, 2007

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

STRIDES PHARMA

EQ 50MG BASE

A065070 001 Dec 15, 2000

EQ 75MG BASE

A065070 003 Dec 30, 2002

EQ 100MG BASE

A065070 002 Dec 15, 2000

EQ 150MG BASE

A065070 004 Jul 14, 2005

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXYCYCLINE HYCLATE

CAPSULE;ORAL

ACTICLATE CAP

+ ALMIRALL

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

AJANTA PHARMA LTD

EQ 50MG BASE

A211012 001 Sep 24, 2018

EQ 100MG BASE

A211012 002 Sep 24, 2018

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

MYLAN

EQ 50MG BASE

A062337 001 Mar 29, 1982

EQ 100MG BASE

A062337 002 Mar 29, 1982

NOSTRUM LABS INC

EQ 50MG BASE

A209393 001 Dec 10, 2020

EQ 100MG BASE

A209393 002 Dec 10, 2020

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

ZHEJIANG YONGTAI

EQ 50MG BASE

A212610 001 Mar 31, 2020

EQ 100MG BASE

A212610 002 Mar 31, 2020

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

BAUSCH

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

HIKMA

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

VIBRAMYCIN

+ PFIZER

EQ 100MG BASE/VIAL **

N050442 002

+

EQ 200MG BASE/VIAL **

N050442 001

SYSTEM, EXTENDED RELEASE;PERIODONTAL

ATRIDOX

+ DEN-MAT

50MG

N050751 001 Sep 03, 1998

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXYCYCLINE HYCLATE

TABLET; ORAL

DOXY-LEMMON					
TEVA	EQ 100MG BASE	A062581	001	Mar 15,	1985
DOXYCYCLINE HYCLATE					
ALEMBIC PHARMS LTD	EQ 100MG BASE	A210536	001	May 14,	2020
AMICI	EQ 100MG BASE	A213475	001	Mar 10,	2021
AMNEAL PHARMS CO	EQ 75MG BASE	A209372	001	Oct 06,	2017
	EQ 150MG BASE	A209372	002	Oct 06,	2017
HEATHER	EQ 100MG BASE	A062462	001	May 11,	1983
HERITAGE PHARMA	EQ 20MG BASE	A065163	001	May 13,	2005
INTERPHARM	EQ 100MG BASE	A062764	001	Sep 02,	1988
MUTUAL PHARM	EQ 100MG BASE	A062391	001	Sep 30,	1982
MYLAN	EQ 100MG BASE	A062432	001	Feb 15,	1983
RISING PHARMA	EQ 75MG BASE	A209987	001	Oct 05,	2020
	EQ 150MG BASE	A209987	002	Oct 05,	2020
STRIDES PHARMA	EQ 100MG BASE	A062538	001	Apr 07,	1986
SUPERPHARM	EQ 100MG BASE	A062494	001	Feb 20,	1985
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 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 PHARMS INC	EQ 150MG BASE	A091052	001	Feb 08,	2012
RISING PHARMA	EQ 50MG BASE	A090431	003	May 23,	2016
	EQ 75MG BASE	A090431	001	Dec 28,	2010
	EQ 80MG BASE	A090431	004	Apr 29,	2016
	EQ 100MG BASE	A090431	002	Dec 28,	2010
	EQ 200MG BASE	A090431	005	May 19,	2016
ZYDUS PHARMS	EQ 75MG BASE	A206772	001	Dec 21,	2018
	EQ 100MG BASE	A206772	002	Dec 21,	2018
	EQ 150MG BASE	A206772	003	Dec 21,	2018

DOXYLAMINE SUCCINATE

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
DOXYLAMINE SUCCINATE					
COPELY 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		

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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	2.5MG/ML;EQ 0.05MG BASE/ML	N016049 001	

DROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL

DROSPIRENONE AND ETHINYL ESTRADIOL

JUBILANT CADISTA	3MG;0.02MG	A209423 001	Dec 22, 2017
KYRA			
SUN PHARM	3MG;0.02MG	A202318 001	Jul 23, 2019

TABLET; ORAL-28

DROSPIRENONE AND ETHINYL ESTRADIOL

APOTEX	3MG;0.03MG	A205876 001	Sep 21, 2016
JUBILANT CADISTA	3MG;0.03MG	A210017 001	Sep 10, 2018
KEMEYA			
SUN PHARM	3MG;0.03MG	A202138 001	Mar 13, 2019

DROXIDOPA

CAPSULE; ORAL

DROXIDOPA

TEVA PHARMS USA INC	100MG	A213162 001	Feb 18, 2021
	200MG	A213162 002	Feb 18, 2021
	300MG	A213162 003	Feb 18, 2021

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS; ORAL

DULOXETINE HYDROCHLORIDE

TEVA PHARMS USA	EQ 20MG BASE	A090783 001	Dec 11, 2013
	EQ 30MG BASE	A090783 002	Dec 11, 2013
	EQ 60MG BASE	A090783 003	Dec 11, 2013
YAOPHARMA CO LTD	EQ 20MG BASE	A207219 001	Aug 16, 2019
	EQ 30MG BASE	A207219 002	Aug 16, 2019
	EQ 60MG BASE	A207219 003	Aug 16, 2019

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DUTASTERIDE

CAPSULE; ORAL

DUTASTERIDE

ACTAVIS LABS FL INC	0.5MG	A202808	001	Nov 20, 2015
APOTEX INC	0.5MG	A204292	001	Nov 24, 2015
HERITAGE PHARMS INC	0.5MG	A207935	001	Oct 13, 2017
HIKMA	0.5MG	A202204	001	Nov 23, 2015
MYLAN	0.5MG	A203241	001	Jun 14, 2016
NOSTRUM LABS INC	0.5MG	A204705	001	Nov 20, 2015
RISING	0.5MG	A202530	001	Nov 20, 2015
STRIDES PHARMA	0.5MG	A208227	001	Jun 22, 2018

DUTASTERIDE; TAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL

DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE

ACTAVIS LABS FL INC	0.5MG;0.4MG	A202975	001	Nov 20, 2015
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DYCLONINE HYDROCHLORIDE

SOLUTION; TOPICAL

DYCLONE

+	ASTRAZENECA	0.5% **	N009925	002
+		1% **	N009925	001

DYDROGESTERONE

TABLET; ORAL

GYNOREST

+	SOLVAY	5MG **	N017388	001
+		10MG **	N017388	002

DYPHYLLINE

ELIXIR; ORAL

NEOTHYLLINE

TEVA	160MG/15ML	N007794	003
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INJECTABLE; INJECTION

NEOTHYLLINE

TEVA	250MG/ML	N009088	001
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TABLET; ORAL

DILOR

SAVAGE LABS	200MG	A084514	001
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DILOR-400

SAVAGE LABS	400MG	A084751	001
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LUFYLLIN

MYLAN SPECIALITY LP	200MG	A084566	001
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	400MG	A084566	002
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NEOTHYLLINE

TEVA	200MG	N007794	001
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	400MG	N007794	002
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ECHOTHIOPHATE IODIDE

FOR SOLUTION; OPHTHALMIC

PHOSPHOLINE IODIDE

FERA PHARMS LLC	0.03%	N011963	002
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	0.06%	N011963	004
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	0.25%	N011963	003
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ECONAZOLE NITRATE

CREAM; TOPICAL

ECONAZOLE NITRATE

CHARTWELL RX	1%	A076075	001	Nov 26, 2002
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MYLAN	1%	A210364	001	Apr 18, 2018
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EDETATE CALCIUM DISODIUM

TABLET; ORAL

CALCIUM DISODIUM VERSENATE

BAUSCH	500MG	N008922	002
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EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

EDROPHONIUM CHLORIDE

HOSPIRA	10MG/ML	A040131	001	Feb 24, 1998
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WATSON LABS	10MG/ML	A040044	001	Mar 20, 1996
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EDROPHONIUM CHLORIDE PRESERVATIVE FREE

WATSON LABS	10MG/ML	A040043	001	Mar 20, 1996
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

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

EFAVIRENZ

MYLAN 600MG

A091471 001 Feb 17, 2016

STRIDES PHARMA 600MG

A078509 001 Jun 16, 2021

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

MACLEODS PHARMS LTD 400MG; 300MG; 300MG

N210649 001 Mar 15, 2019

EFINACONAZOLE

SOLUTION; TOPICAL

EFINACONAZOLE

AUROBINDO PHARMA LTD 10%

A212066 001 Mar 29, 2021

PADAGIS US 10%

A211851 001 Dec 16, 2020

EFLORNITHINE HYDROCHLORIDE

INJECTABLE; INJECTION

ORNIDYL

SANOFI AVENTIS US 200MG/ML

N019879 002 Nov 28, 1990

ELETRIPTAN HYDROBROMIDE

TABLET; ORAL

ELETRIPTAN HYDROBROMIDE

AMNEAL PHARMS CO EQ 20MG BASE

A206787 001 May 25, 2018

EQ 40MG BASE

A206787 002 May 25, 2018

YUNG SHIN PHARM EQ 20MG BASE

A209680 001 Jul 13, 2020

EQ 40MG BASE

A209680 002 Jul 13, 2020

ELTROMBOPAG OLAMINE

TABLET; ORAL

PROMACTA

+ NOVARTIS EQ 100MG ACID **

N022291 005 Nov 16, 2012

ELVITEGRAVIR

TABLET; ORAL

VITEKTA

+ GILEAD SCIENCES INC 85MG

N203093 001 Sep 24, 2014

+ 150MG

N203093 002 Sep 24, 2014

EMEDASTINE DIFUMARATE

SOLUTION/DROPS; OPHTHALMIC

EMADINE

+ NOVARTIS 0.05%

N020706 001 Dec 29, 1997

ENALAPRIL MALEATE

FOR SOLUTION; ORAL

EPANED KIT

+ SILVERGATE PHARMS 1MG/ML **

N204308 001 Aug 13, 2013

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

AUROBINDO PHARMA USA 2.5MG

A075480 001 Aug 22, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ENALAPRIL MALEATE

TABLET; ORAL

ENALAPRIL MALEATE

	5MG	A075480 002	Aug 22, 2000
	10MG	A075480 003	Aug 22, 2000
	20MG	A075480 004	Aug 22, 2000
BEXIMCO PHARMS USA	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
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
	5MG	A075472 002	Aug 22, 2000
	10MG	A075472 003	Aug 22, 2000
	20MG	A075472 004	Aug 22, 2000
NOSTRUM LABS INC	2.5MG	A075178 002	Mar 23, 2001
	5MG	A075178 001	Mar 23, 2001
	10MG	A075178 003	Mar 23, 2001
	20MG	A075178 004	Mar 23, 2001
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
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

CHARTWELL RX	5MG; 12.5MG	A076116 001	Sep 19, 2001
	10MG; 25MG	A076116 002	Sep 19, 2001
IVAX SUB TEVA PHARMS	5MG; 12.5MG	A075736 001	Mar 25, 2003
	10MG; 25MG	A075736 002	Mar 25, 2003
NOSTRUM LABS INC	5MG; 12.5MG	A076486 001	Oct 27, 2004
	10MG; 25MG	A076486 002	Oct 27, 2004
RISING PHARMA	5MG; 12.5MG	A075624 001	Sep 18, 2001
	10MG; 25MG	A075624 002	Sep 18, 2001

ENALAPRILAT

INJECTABLE; INJECTION

ENALAPRILAT

DR REDDYS	1.25MG/ML	A075578 001	Aug 22, 2000
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ENCORAFENIB

CAPSULE; ORAL

BRAFTOVI

+ ARRAY BIOPHARMA INC 50MG N210496 001 Jun 27, 2018

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

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

ENTECAVIR

TABLET; ORAL

ENTECAVIR

RISING PHARMA 0.5MG A206226 001 Mar 26, 2019

1MG A206226 002 Mar 26, 2019

STRIDES PHARMA 0.5MG A206294 001 Nov 23, 2016

1MG A206294 002 Nov 23, 2016

SUNSHINE 0.5MG A211978 001 May 20, 2020

1MG A211978 002 May 20, 2020

TEVA PHARMS USA 0.5MG A202122 001 Aug 26, 2014

1MG A202122 002 Aug 26, 2014

YAOPHARMA CO LTD 0.5MG A212201 001 Nov 04, 2019

1MG A212201 002 Nov 04, 2019

EPHEDRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

REZIPRES

+ ETON 47MG/ML (47MG/ML) N213536 002 Jun 14, 2021

+ 47MG/5ML (9.4MG/ML) N213536 003 Jun 14, 2021

EPHEDRINE SULFATE

SOLUTION; INTRAVENOUS

EPHEDRINE SULFATE

+ ENDO VENTURES 50MG/10ML (5MG/ML) N213994 001 Oct 16, 2020

EPINASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

EPINASTINE HYDROCHLORIDE

CHARTWELL RX 0.05% A203384 001 Dec 07, 2016

SUN PHARM INDS 0.05% A091626 001 Oct 31, 2011

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EPINEPHRINE

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
SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS				
EPINEPHRINE				
AM REGENT	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A207568	001	Jul 06, 2018

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; PRILUCAINE 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
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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

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				
ACTAVIS TOTOWA	10MG/5ML (2MG/ML)		A065445	001 Sep 18, 2008
EBEWE PHARMA	50MG/25ML (2MG/ML)		A065339	001 Dec 22, 2009
	200MG/100ML (2MG/ML)		A065339	002 Dec 22, 2009
FRESENIUS KABI USA				
	10MG/5ML (2MG/ML)		A065408	001 Oct 15, 2007
	50MG/25ML (2MG/ML)		A065408	002 Oct 15, 2007
	150MG/75ML (2MG/ML)		A065408	003 Oct 15, 2007
	200MG/100ML (2MG/ML)		A065408	004 Oct 15, 2007
	200MG/100ML (2MG/ML)		A065411	001 Aug 20, 2007
	50MG/25ML (2MG/ML)		A065411	002 Aug 20, 2007
HOSPIRA				
	10MG/5ML (2MG/ML)		A065343	001 Apr 19, 2007
	50MG/25ML (2MG/ML)		A065343	002 Apr 19, 2007
	150MG/75ML (2MG/ML)		A065343	003 Apr 19, 2007
	200MG/100ML (2MG/ML)		A065343	004 Apr 19, 2007
MYLAN INSTITUTIONAL				
	50MG/25ML (2MG/ML)		A065371	001 Nov 28, 2007
	200MG/100ML (2MG/ML)		A065371	002 Nov 28, 2007
MYLAN LABS LTD				
	50MG/25ML (2MG/ML)		A091599	001 Mar 12, 2012
	200MG/100ML (2MG/ML)		A091599	002 Mar 12, 2012
ZENNOVA				
	50MG/25ML (2MG/ML)		A090266	001 Apr 15, 2011
	200MG/100ML (2MG/ML)		A090266	002 Apr 15, 2011
POWDER; INTRAVENOUS				
EPIRUBICIN HYDROCHLORIDE				
HOSPIRA	50MG/VIAL		N050807	001 Sep 15, 2006
	200MG/VIAL		N050807	002 Sep 15, 2006

EPLERENONE

TABLET; ORAL

EPLERENONE				
PIRAMAL HLTHCARE UK	25MG		A212765	001 Aug 10, 2020
	50MG		A212765	002 Aug 10, 2020
INSPRA				
UPJOHN	100MG		N021437	003 Sep 27, 2002

EPROSARTAN MESYLATE

TABLET; ORAL

EPROSARTAN MESYLATE				
MYLAN PHARMS INC	EQ 400MG BASE		A202012	001 Nov 16, 2011
	EQ 600MG BASE		A202012	002 Nov 16, 2011
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

AMNEAL PHARMS

2MG/ML

A205581 001 Dec 08, 2016

75MG/100ML

A205581 002 Dec 08, 2016

HYBIO

2MG/ML

A207864 001 Mar 20, 2020

75MG/100ML

A207864 002 Mar 20, 2020

TEVA PHARMS USA

75MG/100ML

A091555 001 Jun 05, 2015

USV

2MG/ML

A204361 001 Mar 14, 2019

2MG/ML

A204362 001 Mar 11, 2019

75MG/100ML

A204361 002 Mar 14, 2019

INTEGRILIN

+ SCHERING

2MG/ML

N020718 001 May 18, 1998

+

75MG/100ML

N020718 002 May 18, 1998

ERGOCALCIFEROL

CAPSULE; ORAL

DELTALIN

LILLY

50,000 IU

A080884 001

ERGOCALCIFEROL

SIGMAPHARM LABS LLC

50,000 IU

A091004 001 Jul 14, 2010

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERGOLOID MESYLATES

TABLET;SUBLINGUAL

ERGOLOID MESYLATES

	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	
	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

ACCORD HLTHCARE EQ 25MG BASE

A211083 001 Jul 02, 2020

EQ 100MG BASE

A211083 002 Jul 02, 2020

EQ 150MG BASE

A211083 003 Jul 02, 2020

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 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

+ BAUSCH 2%

N050584 001 Jan 10, 1985

POWDER;FOR RX COMPOUNDING

ERYTHROMYCIN

PADDOCK LLC 100%

N050610 001 Nov 07, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN

SOLUTION;TOPICAL

A/T/S				
TARO	2%		A062405 001	Nov 18, 1982
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
ERYTHRO-STATIN				
AKORN	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	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 effectiveness 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

COSETTE 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

ERYTHROMYCIN ETHYLSUCCINATE

PAR PHARM INC EQ 200MG BASE/5ML A211991 001 Oct 23, 2019

EQ 400MG BASE/5ML A211991 002 Oct 23, 2019

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN ETHYLSUCCINATE

TABLET; ORAL

ERYTHROMYCIN ETHYLSUCCINATE

BARR EQ 400MG BASE A062256 001

MYLAN EQ 400MG BASE A062847 001 Sep 14, 1988

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 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN STEARATE

TABLET; ORAL

ERYTHROMYCIN STEARATE

WATSON LABS

EQ 250MG BASE

A062121 002

EQ 500MG BASE

A062121 001

ETHRIL 250

BRISTOL MYERS SQUIBB EQ 250MG BASE

A061605 001

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

SOLUTION; ORAL

ESCITALOPRAM OXALATE

ANTRIM PHARMS LLC

EQ 5MG BASE/5ML

A203967 001 May 26, 2015

LEXAPRO

+ ALLERGAN

EQ 5MG BASE/5ML **

N021365 001 Nov 27, 2002

TABLET; ORAL

ESCITALOPRAM OXALATE

HIKMA PHARMS

EQ 5MG BASE

A078766 001 Sep 11, 2012

EQ 10MG BASE

A078766 002 Sep 11, 2012

EQ 20MG BASE

A078766 003 Sep 11, 2012

MACLEODS PHARMS LTD

EQ 5MG BASE

A202210 001 Sep 11, 2012

EQ 10MG BASE

A202210 002 Sep 11, 2012

EQ 20MG BASE

A202210 003 Sep 11, 2012

MYLAN

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

ESMOLOL HYDROCHLORIDE

AM REGENT

10MG/ML

A201126 001 Feb 20, 2015

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED REL PELLETS; ORAL

ESOMEPRAZOLE MAGNESIUM

AMNEAL PHARMS NY

EQ 20MG BASE

A209647 001 Apr 10, 2019

EQ 40MG BASE

A209647 002 Apr 10, 2019

AMTA

EQ 20MG BASE

A213859 001 Nov 18, 2020

EQ 40MG BASE

A213859 002 Nov 18, 2020

HEC PHARM

EQ 20MG BASE

A207265 002 May 18, 2018

EQ 40MG BASE

A207265 001 May 18, 2018

CAPSULE, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

MYLAN

EQ 20MG BASE

A212376 001 Oct 16, 2019

TABLET, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

P AND L

EQ 20MG BASE

A209202 001 Mar 05, 2019

ESOMEPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS

ESOMEPRAZOLE SODIUM

EUGIA PHARMA

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

EQ 20MG BASE/VIAL **

N021689 001 Mar 31, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESOMEPRAZOLE STRONTIUM

CAPSULE, DELAYED RELEASE;ORAL

ESOMEPRAZOLE STRONTIUM

+	BELCHER	24.65MG	N202342 001	Aug 06, 2013
+		49.3MG	N202342 002	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

ALORA

	ALLERGAN	0.025MG/24HR	N020655 004	Apr 05, 2002
		0.05MG/24HR	N020655 001	Dec 20, 1996
		0.075MG/24HR	N020655 002	Dec 20, 1996
		0.1MG/24HR	N020655 003	Dec 20, 1996

FEMPATCH

	PARKE DAVIS	0.025MG/24HR	N020417 001	Dec 03, 1996
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GEL;TOPICAL

ESTROGEL

	ASCEND THERAPS US	0.06%	N021166 001	Feb 09, 2004
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SYSTEM;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

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

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
	MYLAN	0.5MG	A040326 001	Apr 21, 1999
		1MG	A040326 002	Apr 21, 1999
		2MG	A040326 003	Apr 21, 1999
	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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESTRADIOL

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, 2004

ESTRADIOL CYPIONATE

INJECTABLE; INJECTION

DEPO-ESTRADIOL

PFIZER

1MG/ML

A085470 001

3MG/ML

A085470 002

ESTRADIOL CYPIONATE

DR REDDYS

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

DR REDDYS

20MG/ML

A083547 001

40MG/ML

A083714 001

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

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; NORETHINDRONE ACETATE

TABLET; ORAL

ACTIVELLA

+ AMNEAL 0.5MG; 0.1MG N020907 002 Dec 28, 2006

ESTRADIOL AND NORETHINDRONE ACETATE

NAARI PTE LTD

1MG; 0.5MG

A210233 001 Feb 28, 2018

TEVA PHARMS USA

0.5MG; 0.1MG

A200747 001 Mar 08, 2012

ESTRADIOL; NORGESTIMATE

TABLET; ORAL

PREFEST

+ TEVA WOMENS 1MG, 1MG; N/A, 0.09MG ** N021040 001 Oct 22, 1999

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

+	ASPEN	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

	ASPEN	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
		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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESTRONE

INJECTABLE; INJECTION

ESTROGENIC SUBSTANCE

WYETH AYERST	2MG/ML	A083488	001	
ESTRONE				
DR REDDYS	5MG/ML	A085239	001	
WATSON LABS	2MG/ML	A083397	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	
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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	0.75MG	A040359	001	Aug 26, 1999
	1.5MG	A040359	002	Aug 26, 1999
	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				
+ PFIZER	0.75MG	A083220	001	
OGEN 1.25				
+ PFIZER	1.5MG	A083220	002	
OGEN 2.5				
+ PFIZER	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

HIKMA	1MG	A091153	001	Apr 15, 2014
	2MG	A091153	002	Apr 15, 2014
	3MG	A091153	003	Apr 15, 2014
NOSTRUM LABS INC	1MG	A203087	001	May 08, 2019
	2MG	A203087	002	May 08, 2019
	3MG	A203087	003	May 08, 2019
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

BAUSCH	50MG	N016092	002	
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ETHAMBUTOL HYDROCHLORIDE

TABLET; ORAL

MYAMBUTOL

STI PHARMA LLC	200MG	N016320	002	
	500MG	N016320	004	

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
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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
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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
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DEMULEN 1/50-21

GD SEARLE LLC	0.05MG; 1MG	N016927 001
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ZOVIA 1/35E-21

WATSON PHARMS TEVA	0.035MG; 1MG	A072720 001	Dec 30, 1991
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ZOVIA 1/50E-21

WATSON LABS	0.05MG; 1MG	A072722 001	Dec 30, 1991
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TABLET; ORAL-28

DEMULEN 1/35-28

GD SEARLE LLC	0.035MG; 1MG **	N018160 001
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DEMULEN 1/50-28

GD SEARLE LLC	0.05MG; 1MG **	N016936 001
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ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE

TABLET; ORAL-28

NORQUEST FE

PFIZER	0.035MG; 75MG; 1MG	N018926 001	Jul 18, 1986
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ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE ACETATE

TABLET; ORAL-28

NORLESTRIN FE 1/50

PARKE DAVIS	0.05MG; 75MG; 1MG	N016766 001
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NORLESTRIN FE 2.5/50

PARKE DAVIS	0.05MG; 75MG; 2.5MG	N016854 001
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ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL

FAYOSIM

LUPIN LTD	0.02MG, 0.15MG; 0.025MG, 0.15MG; 0.03MG, 0.15MG; 0.01MG, N/A	A205943 001	Mar 29, 2016
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LYBREL

+ WYETH PHARMS INC	0.02MG; 0.09MG **	N021864 001	May 22, 2007
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PREVEN EMERGENCY CONTRACEPTIVE KIT

TEVA BRANDED PHARM	0.05MG; 0.25MG	N020946 001	Sep 01, 1998
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SYLEVIA

SUN PHARM	0.03MG; 0.15MG	A202988 001	Feb 06, 2019
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TABLET; ORAL-21

ALESSE

+ CADENCE HEALTH	0.02MG; 0.1MG **	N020683 001	Mar 27, 1997
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL-21

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

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

CERINTA

SUN PHARM 0.02MG;0.1MG A202817 001 Jan 07, 2019

ELIFEMME

XIROMED 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG A202507 001 Dec 04, 2015

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

MYLAN LABS LTD 0.02MG;0.1MG A202247 001 Dec 08, 2014

0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG A202970 001 Mar 23, 2018

NORDETTE-28

+ TEVA BRANDED PHARM 0.03MG;0.15MG ** N018782 001 Jul 21, 1982

ORSYTHIA

VINTAGE PHARMS LLC 0.02MG;0.1MG A077099 001 May 11, 2011

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 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21

NORETHINDRONE AND ETHINYL ESTRADIOL				
	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
NORINYL 1+35 21-DAY				
ALLERGAN	0.035MG;1MG	N017565	001	
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
ORTHO-NOVUM 7/7/7-21				
JANSSEN PHARMS	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	N018985	001	Apr 04, 1984
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				
BREVICON 28-DAY				
ALLERGAN	0.035MG;0.5MG	N017743	001	
CYCLAFEM 0.5/35				
VINTAGE PHARMS	0.035MG;0.5MG	A203413	001	Dec 16, 2015
CYCLAFEM 1/35				
VINTAGE PHARMS LLC	0.035MG;1MG	A076337	001	Nov 12, 2010
CYCLAFEM 7/7/7				
VINTAGE PHARMS LLC	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A076338	001	Nov 16, 2010
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
	0.035MG;0.5MG	A200488	001	Oct 21, 2015
	0.035MG;1MG	A200489	001	Oct 21, 2015
	0.05MG;1MG	A203006	001	Aug 05, 2013
WATSON LABS	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A076393	001	Feb 04, 2010
NORETHINDRONE AND ETHINYL ESTRADIOL (7/14)				
WATSON LABS	0.035MG,0.035MG;0.5MG,1MG	A071042	001	Sep 24, 1991
NORINYL 1+35 28-DAY				
ALLERGAN	0.035MG;1MG	N017565	002	
ORTHO-NOVUM 1/35-28				
+ JANSSEN PHARMS	0.035MG;1MG **	N017919	002	
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
ORTHO-NOVUM 7/7/7-28				
+ JANSSEN PHARMS	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	N018985	002	Apr 04, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET;ORAL-28

OVCON-35

+ WARNER CHILCOTT LLC 0.035MG;0.4MG **

N017716 001

OVCON-50

WARNER CHILCOTT LLC 0.05MG;1MG **

N017576 001

TABLET, CHEWABLE;ORAL

FEMCON FE

+ APIL 0.035MG;0.4MG **

N021490 001 Nov 14, 2003

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET;ORAL

FEMHRT

+ APIL 0.0025MG;0.5MG

N021065 001 Jan 14, 2005

+ 0.005MG;1MG **

N021065 002 Oct 15, 1999

LOESTRIN 24 FE

+ TEVA BRANDED PHARM 0.02MG;1MG **

N021871 001 Feb 17, 2006

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

MYLAN LABS LTD 0.0025MG;0.5MG

A207260 001 Feb 02, 2017

0.005MG;1MG

A207259 001 Dec 27, 2016

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 0.02MG;1MG

A208639 001 Mar 21, 2018

MYLAN LABS LTD 0.02MG;1MG

A202742 001 Oct 30, 2014

TABLET;ORAL-21

ESTROSTEP 21

+ APIL 0.02MG,0.03MG,0.035MG;1MG,1MG,1MG **

N020130 001 Oct 09, 1996

GILDESS 1.5/30

VINTAGE PHARMS LLC 0.03MG;1.5MG

A077075 002 Jul 24, 2012

GILDESS 1/20

VINTAGE PHARMS LLC 0.02MG;1MG

A077077 002 Jul 24, 2012

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

ESTROSTEP FE

+ APIL 0.02MG,0.03MG,0.035MG;1MG,1MG,1MG

N020130 002 Oct 09, 1996

GILDESS FE 1.5/30

VINTAGE PHARMS LLC 0.03MG;1.5MG

A077075 001 Apr 28, 2005

GILDESS FE 1/20

VINTAGE PHARMS LLC 0.02MG;1MG

A077077 001 May 20, 2005

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

MYLAN LABS LTD 0.02MG,0.03MG,0.035MG;1MG,1MG,1MG

A205069 001 Jun 22, 2018

NORLESTRIN 28 1/50

PARKE DAVIS 0.05MG;1MG

N016723 001

TABLET, CHEWABLE;ORAL

FINZALA

TEVA PHARMS USA INC 0.02MG;1MG

A210087 001 Apr 07, 2020

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

MYLAN LABS LTD 0.02MG;1MG

A206120 001 Sep 12, 2017

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

TABLET;ORAL-28

NORGESTIMATE AND ETHINYL ESTRADIOL

MYLAN 0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,
0.25MG

A201897 001 Jan 27, 2016

0.035MG;0.25MG

A201896 001 Jan 27, 2016

MYLAN LABS LTD 0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,
0.25MG

A202132 001 Sep 09, 2015

WATSON LABS 0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,

A090479 001 Mar 09, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

NORGESTIMATE AND ETHINYL ESTRADIOL

	0.25MG		
	0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG,	A076626	001 Aug 17, 2006
	0.25MG		
	0.035MG; 0.25MG	A076627	001 Aug 17, 2006
ORTHO CYCLEN-28			
+ JANSSEN PHARMS	0.035MG; 0.25MG **	N019653	002 Dec 29, 1989
ORTHO TRI-CYCLEN			
+ JANSSEN PHARMS	0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG,	N019697	001 Jul 03, 1992
	0.25MG		
ORTHO TRI-CYCLEN LO			
+ JANSSEN PHARMS	0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG,	N021241	001 Aug 22, 2002
	0.25MG		
TRI-PREVIFEM			
VINTAGE PHARMS LLC	0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG,	A076335	001 Mar 26, 2004
	0.25MG		

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
OGESTREL 0.5/50-28			
WATSON LABS	0.05MG; 0.5MG	A075406	002 Dec 15, 1999
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

ETHOSUXIMIDE

SYRUP; ORAL

ETHOSUXIMIDE

TEVA PHARMS	250MG/5ML	A081306	001 Jul 30, 1993
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ETHOTOIN

TABLET; ORAL

PEGANONE

+ RECORDATI RARE	250MG	N010841	001
	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
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHYLESTRENOL

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

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

ETIDRONATE DISODIUM

MYLAN

200MG

A075800 001 Jan 24, 2003

400MG

A075800 002 Jan 24, 2003

ETODOLAC

CAPSULE; ORAL

ETODOLAC

ANI PHARMS

200MG

A074840 001 Aug 29, 1997

200MG

A074844 001 Dec 23, 1997

200MG

A074899 001 Jul 08, 1997

300MG

A074840 002 Aug 29, 1997

300MG

A074844 002 Dec 23, 1997

300MG

A074899 002 Jul 08, 1997

BIOPHARM

300MG

A074929 001 Jan 30, 1998

CHARTWELL MOLECULES

200MG

A074842 001 Jul 17, 1997

300MG

A074842 002 Jul 17, 1997

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

LODINE

+ WYETH PHARMS INC

200MG **

N018922 002 Jan 31, 1991

+

300MG

N018922 003 Jan 31, 1991

TABLET; ORAL

ETODOLAC

BIOPHARM

400MG

A074927 001 Oct 30, 1997

CHARTWELL MOLECULES

400MG

A074841 001 Jun 27, 1997

IVAX SUB TEVA PHARMS

400MG

A074883 001 Feb 28, 1997

500MG

A074883 002 Nov 20, 1998

MYLAN

400MG

A075012 001 Sep 30, 1998

400MG

A075104 001 Feb 06, 1998

500MG

A075012 002 Sep 30, 1998

500MG

A075104 002 Nov 20, 1998

OXFORD PHARMS

400MG

A074819 001 Feb 28, 1997

500MG

A074819 002 Apr 28, 1998

PHARMGEN

400MG

A074839 001 Jul 11, 1997

400MG

A074846 001 Feb 28, 1997

RANBAXY LABS LTD

400MG

A075226 001 Nov 24, 1998

500MG

A075226 002 Nov 24, 1998

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETODOLAC

TABLET; ORAL

ETODOLAC

TEVA	400MG	A074847 001	Apr 23, 1999
	400MG	A075009 001	Nov 26, 1997
	500MG	A074847 002	Apr 23, 1999
	500MG	A075009 002	Dec 28, 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	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
	500MG **	N020584 003	Jan 20, 1998
+	600MG **	N020584 002	Oct 25, 1996

ETOMIDATE

INJECTABLE; INJECTION

ETOMIDATE

LUITPOLD	2MG/ML	A078867 001	Dec 22, 2009
PAR STERILE PRODUCTS	2MG/ML	A091297 001	Jun 20, 2012
RISING PHARMA	2MG/ML	A078289 001	Jan 02, 2009

ETONOGESTREL

IMPLANT; IMPLANTATION

IMPLANON

ORGANON USA INC	68MG/IMPLANT	N021529 001	Jul 17, 2006
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ETOPOSIDE

CAPSULE; ORAL

VEPESID

+ STRIDES PHARMA	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
MYLAN LABS LTD	20MG/ML	A203507 001	Nov 20, 2017
	20MG/ML	A204927 001	Oct 31, 2017
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

ETOPHOS PRESEVATIVE FREE

BRISTOL MYERS SQUIBB	EQ 500MG BASE/VIAL	N020906 001	Feb 27, 1998
	EQ 1GM BASE/VIAL	N020906 002	Feb 27, 1998

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETRAVIRINETABLET; ORAL
ETRAVIRINE

AMNEAL 25MG A214196 001 Jun 14, 2021

ETRETINATECAPSULE; ORAL
TEGISONROCHE 10MG N019369 001 Sep 30, 1986
25MG N019369 002 Sep 30, 1986EVANS BLUEINJECTABLE; INJECTION
EVANS BLUE

PARKE DAVIS 0.5% ** N008041 001

EVEROLIMUSTABLET; ORAL
EVEROLIMUS

TEVA PHARMS USA 10MG A210050 004 Dec 09, 2019

EXEMESTANETABLET; ORAL
EXEMESTANEAMNEAL PHARMS 25MG A206421 001 Dec 28, 2018
MAYNE PHARMA INC 25MG A208764 001 Aug 08, 2019EXENATIDE 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

EZETIMIBETABLET; ORAL
EZETIMIBERISING PHARMA 10MG A201790 001 Apr 26, 2019
TEVA PHARMS USA 10MG A078724 001 Jun 12, 2017EZETIMIBE; SIMVASTATIN

TABLET; ORAL

EZETIMIBE AND SIMVASTATIN

ANI PHARMS 10MG; 10MG A201890 001 Apr 26, 2017
10MG; 20MG A201890 002 Apr 26, 2017
10MG; 40MG A201890 003 Apr 26, 2017
10MG; 80MG A201890 004 Apr 26, 2017
AUROBINDO PHARMA USA 10MG; 10MG A200082 001 Dec 17, 2020
10MG; 20MG A200082 002 Dec 17, 2020
10MG; 40MG A200082 003 Dec 17, 2020
10MG; 80MG A200082 004 Dec 17, 2020EZOGABINE

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, 2011FAMCICLOVIR

TABLET; ORAL

FAMCICLOVIR

HIKMA 125MG A090128 001 Mar 21, 2011
250MG A090128 002 Mar 21, 2011
500MG A090128 003 Mar 21, 2011
MYLAN 125MG A201333 001 Mar 24, 2011
250MG A201333 002 Mar 24, 2011
500MG A201333 003 Mar 24, 2011

FAMVIR

+ NOVARTIS 125MG ** N020363 003 Dec 11, 1995
+ 250MG ** N020363 001 Apr 26, 1996
+ 500MG ** N020363 002 Jun 29, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FAMOTIDINE

FOR SUSPENSION;ORAL

PEPCID

+ SALIX PHARMS

40MG/5ML

N019527 001 Feb 02, 1987

INJECTABLE; INJECTION

FAMOTIDINE

APOTEX INC

10MG/ML

A075942 001 Aug 02, 2002

APOTHECON

10MG/ML

A075707 001 Apr 16, 2001

HIKMA

10MG/ML

A075799 001 Apr 30, 2002

HOSPIRA

10MG/ML

A075705 001 Apr 16, 2001

10MG/ML

A075870 001 Nov 23, 2001

10MG/ML

A075905 001 Nov 23, 2001

FAMOTIDINE PRESERVATIVE FREE

APOTEX INC

10MG/ML

A076324 001 Nov 27, 2002

APOTHECON

10MG/ML

A075708 001 Apr 16, 2001

HIKMA

10MG/ML

A075789 001 Apr 30, 2002

HOSPIRA

10MG/ML

A075669 001 Apr 16, 2001

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

MANKIND PHARMA

20MG

A075302 001 Apr 16, 2001

40MG

A075302 002 Apr 16, 2001

MYLAN

10MG

A075674 001 Dec 21, 2001

MYLAN PHARMS INC

20MG

A075457 001 Apr 18, 2001

40MG

A075457 002 Apr 18, 2001

RISING PHARMA

20MG

A075704 001 Apr 16, 2001

40MG

A075704 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

TEVA

10MG

A075312 001 May 31, 2001

20MG

A075311 001 Apr 16, 2001

40MG

A075311 002 Apr 16, 2001

WATSON LABS

10MG

A075404 001 Nov 28, 2001

20MG

A075062 002 Apr 16, 2001

40MG

A075062 001 Apr 16, 2001

PEPCID

+ BAUSCH

20MG

N019462 001 Oct 15, 1986

+

40MG

N019462 002 Oct 15, 1986

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FELODIPINE

TABLET, EXTENDED RELEASE;ORAL

FELODIPINE

JUBILANT GENERICS	2.5MG	A203983 001	Aug 19, 2016
	5MG	A203983 002	Aug 19, 2016
	10MG	A203983 003	Aug 19, 2016
MYLAN	2.5MG	A078855 001	Apr 17, 2008
	5MG	A078855 002	Apr 17, 2008
	10MG	A078855 003	Apr 17, 2008
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	87MG	N021695 002	Nov 30, 2004
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FENOFIBRATE (MICRONIZED)

MYLAN PHARMS INC	43MG	A202579 001	Jan 10, 2013
	130MG	A202579 002	Jan 10, 2013
NOVAST LABS	67MG	A207564 001	Apr 19, 2019
	134MG	A207564 002	Apr 19, 2019
	200MG	A207564 003	Apr 19, 2019
RISING PHARMA	67MG	A202676 001	Oct 23, 2012
	134MG	A202676 002	Oct 23, 2012
	200MG	A202676 003	Oct 23, 2012

LIPIDIL

ABBVIE	100MG	N019304 001	Dec 31, 1993
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LIPOFEN

CIPHER PHARMS INC	100MG	N021612 002	Jan 11, 2006
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TRICOR (MICRONIZED)

+ ABBVIE	67MG **	N019304 002	Feb 09, 1998
+	134MG **	N019304 003	Jun 30, 1999
+	200MG **	N019304 004	Jun 30, 1999

TABLET;ORAL

FENOFIBRATE

ALEMBIC PHARMS LTD	54MG	A213252 001	Jan 17, 2020
	160MG	A213252 002	Jan 17, 2020
MYLAN	107MG	A076520 002	Dec 29, 2005

TRICOR

+ ABBVIE INC	54MG **	N021203 001	Sep 04, 2001
+	160MG **	N021203 003	Sep 04, 2001

TRIGLIDE

SKYEPHARMA AG	50MG	N021350 001	May 07, 2005
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FENOFIBRIC ACID

TABLET;ORAL

FIBRICOR

+ ATHENA	35MG	N022418 001	Aug 14, 2009
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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
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FENOPROFEN CALCIUM

CAPSULE;ORAL

FENOPROFEN CALCIUM

RISING	EQ 200MG BASE	A072394 001	Oct 17, 1988
	EQ 300MG BASE	A072395 001	Oct 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	
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TABLET;ORAL

FENOPROFEN CALCIUM

AM THERAP	EQ 600MG BASE	A072309 001	Aug 17, 1988
ANI PHARMS	EQ 600MG BASE	A072274 001	May 02, 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
RISING	EQ 600MG BASE	A072396 001	Oct 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	
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FENTANYL

FILM, EXTENDED RELEASE;TRANSDERMAL

DURAGESIC-100

+ JANSSEN PHARMS	100MCG/HR	N019813 001	Aug 07, 1990
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DURAGESIC-12

+ JANSSEN PHARMS	12.5MCG/HR	N019813 005	Feb 04, 2005
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DURAGESIC-25

+ JANSSEN PHARMS	25MCG/HR	N019813 004	Aug 07, 1990
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DURAGESIC-37

+ JANSSEN PHARMS	37.5MCG/HR	N019813 006	Jan 24, 2018
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DURAGESIC-50

+ JANSSEN PHARMS	50MCG/HR	N019813 003	Aug 07, 1990
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DURAGESIC-75

+ JANSSEN PHARMS	75MCG/HR	N019813 002	Aug 07, 1990
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FENTANYL-100

ACTAVIS LABS UT INC	100MCG/HR	A076709 004	Aug 20, 2007
LAVIPHARM LABS	100MCG/HR	A077051 004	Aug 04, 2006
MAYNE PHARMA	100MCG/HR	A077062 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
LAVIPHARM LABS	25MCG/HR	A077051 001	Aug 04, 2006
MAYNE PHARMA	25MCG/HR	A077062 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
LAVIPHARM LABS	50MCG/HR	A077051 002	Aug 04, 2006
MAYNE PHARMA	50MCG/HR	A077062 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
LAVIPHARM LABS	75MCG/HR	A077051 003	Aug 04, 2006
MAYNE PHARMA	75MCG/HR	A077062 003	Aug 20, 2007
NOVEN	75MCG/HR	A077775 003	Oct 16, 2009

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FENTANYL CITRATEFILM;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

DR REDDYS	EQ 0.05MG BASE/ML	A074917 001	Feb 03, 1998
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SPRAY, METERED;NASAL

LAZANDA

+ BTCP PHARMA	EQ 0.3MG BASE	N022569 003	Dec 21, 2015
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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
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TABLET;SUBLINGUAL

ABSTRAL

+ SENTYNL THERAPS INC	EQ 0.1MG BASE **	N022510 001	Jan 07, 2011
+	EQ 0.2MG BASE **	N022510 002	Jan 07, 2011
+	EQ 0.3MG BASE **	N022510 003	Jan 07, 2011
+	EQ 0.4MG BASE **	N022510 004	Jan 07, 2011
+	EQ 0.6MG BASE **	N022510 005	Jan 07, 2011
+	EQ 0.8MG BASE **	N022510 006	Jan 07, 2011

FENTANYL CITRATE

ACTAVIS LABS FL INC	EQ 0.1MG BASE	A207338 001	Nov 17, 2017
	EQ 0.2MG BASE	A207338 002	Nov 17, 2017
	EQ 0.3MG BASE	A207338 003	Nov 17, 2017
	EQ 0.4MG BASE	A207338 004	Nov 17, 2017
	EQ 0.6MG BASE	A207338 005	Nov 17, 2017
	EQ 0.8MG BASE	A207338 006	Nov 17, 2017

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
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FERRIC AMMONIUM CITRATE

FOR SOLUTION;ORAL

FERRISELTZ

OTSUKA	600MG/PACKET	N020292 001	Oct 14, 1997
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FERRIC DERISOMALTOSESOLUTION; INTRAVENOUS
MONOFERRIC

+ PHARMACOSMOS AS	100MG/ML (100MG/ML)	N208171 001	Jan 16, 2020
+	500MG/5ML (100MG/ML)	N208171 002	Jan 16, 2020

FERRIC OXYHYDROXIDEINJECTABLE; INJECTION
DEXFERRUM

AM REGENT	EQ 50MG IRON/ML	N040024 001	Feb 23, 1996
IRON DEXTRAN			
SANOFI AVENTIS US	EQ 50MG IRON/ML	N010787 002	
PROFERDEX			
NEW RIVER	EQ 50MG IRON/ML	N017807 001	
INJECTABLE; INTRAVENOUS			
VENOFER			
AM REGENT	EQ 65MG IRON/3.25ML (EQ 20MG IRON/ML)	N021135 005	Mar 29, 2013
	EQ 75MG IRON/3.75ML (EQ 20MG IRON/ML)	N021135 003	Mar 29, 2005

FERRIC PYROPHOSPHATE CITRATESOLUTION; INTRAVENOUS
TRIFERIC

+ ROCKWELL MEDICAL INC	272MG IRON/50ML (5.44MG IRON/ML)	N206317 002	Sep 04, 2015
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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

FERUMOXSIIL

SUSPENSION; ORAL

GASTROMARK			
AMAG PHARMS INC	EQ 0.175MG IRON/ML	N020410 001	Dec 06, 1996

FESOTERODINE FUMARATE

TABLET, EXTENDED RELEASE; ORAL

FESOTERODINE FUMARATE			
ALKEM LABS LTD	4MG	A204827 001	Dec 10, 2015
	8MG	A204827 002	Dec 10, 2015
DR REDDYS LABS LTD	4MG	A204975 001	Aug 13, 2019
	8MG	A204975 002	Aug 13, 2019

FEXOFENADINE 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			
+ CHATTEM SANOFI	30MG/5ML	N021963 001	Oct 16, 2006
CHILDREN'S ALLEGRA HIVES			
+ CHATTEM SANOFI	30MG/5ML **	N201373 002	Jan 24, 2011
FEXOFENADINE HYDROCHLORIDE			
P AND L	30MG/5ML	A201311 001	Jul 25, 2012

TABLET; ORAL

ALLEGRA HIVES			
+ CHATTEM SANOFI	60MG	N020872 008	Jan 24, 2011
+	180MG	N020872 009	Jan 24, 2011
CHILDREN'S ALLEGRA ALLERGY			
+ CHATTEM SANOFI	30MG **	N020872 005	Jan 24, 2011
CHILDREN'S ALLEGRA HIVES			
+ CHATTEM SANOFI	30MG	N020872 006	Jan 24, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FEXOFENADINE HYDROCHLORIDE

TABLET;ORAL

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY					
RISING	30MG	A077081	004	Jul 21, 2011	
CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES					
RISING	30MG	A077081	005	Jul 21, 2011	
FEXOFENADINE HYDROCHLORIDE					
BARR	30MG	A076191	001	Aug 31, 2005	
	60MG	A076191	002	Aug 31, 2005	
	180MG	A076191	003	Aug 31, 2005	
RISING	30MG	A077081	002	Apr 11, 2008	
TABLET, ORALLY DISINTEGRATING;ORAL					
CHILDREN'S ALLEGRA ALLERGY					
+ CHATTEM SANOFI	30MG	N021909	002	Jan 24, 2011	
CHILDREN'S ALLEGRA HIVES					
+ CHATTEM SANOFI	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	

FINAFLOXACIN

SUSPENSION/DROPS;OTIC

XTORO					
+ MERLION PHARMS GMBH	0.3%	N206307	001	Dec 17, 2014	

FINASTERIDE

TABLET;ORAL

FINASTERIDE					
ACTAVIS TOTOWA	1MG	A078371	001	Nov 05, 2013	
ACTAVIS TOTOWA TEVA	5MG	A077914	001	Mar 28, 2007	
GEDEON RICHTER USA	5MG	A077251	001	Dec 22, 2006	
IVAX SUB TEVA PHARMS	5MG	A076340	001	Jun 19, 2006	
MYLAN	1MG	A078161	001	Nov 05, 2013	
	5MG	A077578	001	Dec 18, 2006	
TEVA	1MG	A076905	001	Nov 05, 2013	

FINGOLIMOD HYDROCHLORIDE

CAPSULE;ORAL

FINGOLIMOD HYDROCHLORIDE					
ALKEM LABS LTD	EQ 0.5MG BASE	A208004	001	Dec 30, 2020	
APOTEX	EQ 0.5MG BASE	A207993	001	Dec 18, 2020	
BIOCON LTD	EQ 0.5MG BASE	A207979	001	Dec 04, 2019	
DR REDDYS	EQ 0.5MG BASE	A208000	001	Mar 05, 2021	
GLENMARK PHARMS LTD	EQ 0.5MG BASE	A207985	001	Jun 18, 2020	
HETERO LABS LTD V	EQ 0.5MG BASE	A207933	001	May 18, 2020	
STRIDES PHARMA	EQ 0.5MG BASE	A207971	001	Jun 29, 2020	
SUN PHARM	EQ 0.5MG BASE	A208014	001	Dec 04, 2019	
TEVA PHARMS USA	EQ 0.5MG BASE	A208008	001	Jul 02, 2020	
ZYDUS PHARMS	EQ 0.5MG BASE	A207994	001	Oct 14, 2020	

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	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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLECAINIDE ACETATE

TABLET; ORAL

FLECAINIDE ACETATE

	150MG	A079164 003	Jul 09, 2009
TAMBOCOR			
+ ALVOGEN	50MG **	N018830 004	Aug 23, 1988
+	100MG **	N018830 001	Oct 31, 1985
+	150MG **	N018830 003	Jun 03, 1988
	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
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FLOXURIDINE

INJECTABLE; INJECTION

FLOXURIDINE

AM REGENT	500MG/VIAL	A203008 001	Nov 22, 2017
FUDR			
+ HOSPIRA	500MG/VIAL **	N016929 001	

FLUCONAZOLE

FOR SUSPENSION; ORAL

FLUCONAZOLE

HIKMA	50MG/5ML	A076246 001	Jul 29, 2004
	200MG/5ML	A076246 002	Jul 29, 2004
IVAX SUB TEVA PHARMS	50MG/5ML	A077523 001	Sep 12, 2007
	200MG/5ML	A077523 002	Sep 12, 2007
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

HIKMA FARMACEUTICA	200MG/100ML (2MG/ML)	A078764 001	Jan 30, 2012
	400MG/200ML (2MG/ML)	A078764 002	Jan 30, 2012
HOSPIRA	200MG/100ML (2MG/ML)	A076304 001	Jul 29, 2004
	400MG/200ML (2MG/ML)	A076304 002	Jul 29, 2004
MYLAN LABS LTD	200MG/100ML (2MG/ML)	A076888 001	Mar 25, 2005
	400MG/200ML (2MG/ML)	A076888 002	Mar 25, 2005
WOODWARD	200MG/100ML (2MG/ML)	A077988 001	May 26, 2010
	400MG/200ML (2MG/ML)	A077988 002	May 26, 2010

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

DR REDDYS	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

DR REDDYS	200MG/100ML (2MG/ML)	A076837 001	Jan 13, 2005
	400MG/200ML (2MG/ML)	A076837 002	Jan 13, 2005
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

TABLET; ORAL

FLUCONAZOLE

ANI PHARMS	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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUCONAZOLE

TABLET; ORAL

FLUCONAZOLE

	200MG	A076432 004	Jul 29, 2004
MYLAN	50MG	A076351 001	Jul 29, 2004
	100MG	A076351 002	Jul 29, 2004
	150MG	A076351 003	Jul 29, 2004
	200MG	A076351 004	Jul 29, 2004
MYLAN PHARMS INC	50MG	A076042 001	Jul 29, 2004
	100MG	A076042 002	Jul 29, 2004
	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
TEVA	50MG	A074681 001	Jul 29, 2004
	100MG	A074681 002	Jul 29, 2004
	150MG	A074681 003	Jul 29, 2004
	200MG	A074681 004	Jul 29, 2004

FLUCYTOSINE

CAPSULE; ORAL

FLUCYTOSINE

STRIDES PHARMA	250MG	A207536 001	Jun 18, 2018
	250MG	A212632 001	Apr 17, 2020
	500MG	A207536 002	Jun 18, 2018
	500MG	A212632 002	Apr 17, 2020

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARA

+ GENZYME CORP	50MG/VIAL **	N020038 001	Apr 18, 1991
FLUDARABINE PHOSPHATE			
HOSPIRA	50MG/VIAL	A077790 001	Apr 06, 2007
MYLAN LABS LTD	50MG/2ML (25MG/ML)	A200647 001	Dec 21, 2011
	50MG/VIAL	A200648 001	Oct 16, 2012

TABLET; ORAL

OFORTA

SANOFI AVENTIS US	10MG	N022273 001	Dec 18, 2008
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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
UNIV TX SW MEDCTR	20-200mCi/ML	A210265 001	Feb 06, 2020
WEILL MEDCL COL	10-100mCi/ML **	N021768 001	Aug 05, 2004

FLUDROCORTISONE ACETATE

TABLET; ORAL

FLORINEF

+ CASPER PHARMA LLC	0.1MG **	N010060 001	
FLUDROCORTISONE ACETATE			
HIKMA PHARMS	0.1MG	A091302 001	Jul 22, 2011

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

DR REDDYS	0.5MG/5ML (0.1MG/ML)	A076589 002	Oct 12, 2004
	1MG/10ML (0.1MG/ML)	A076589 001	Oct 12, 2004
RISING PHARMA	1MG/10ML (0.1MG/ML)	A078595 002	May 13, 2008
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	
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FLUNISOLIDE

AEROSOL, METERED; INHALATION

AEROBID

ROCHE PALO	0.25MG/INH	N018340 001	Aug 17, 1984
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AEROSPAN HFA

+ MYLAN SPECIALITY LP	0.078MG/INH	N021247 001	Jan 27, 2006
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SPRAY, METERED; NASAL

NASALIDE

+ IVAX RES	0.025MG/SPRAY **	N018148 001	
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NASAREL

TEVA BRANDED PHARM	0.029MG/SPRAY	N020409 001	Mar 08, 1995
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FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

FLUCET

ALPHARMA US PHARMS	0.025%	A088360 001	Jan 16, 1984
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FLUOCINOLONE ACETONIDE

ALLIED	0.01%	A088047 001	Dec 16, 1982
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	0.025%	A088045 001	Dec 16, 1982
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ALPHARMA US PHARMS	0.01%	A088361 001	Jan 16, 1984
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COSETTE	0.025%	A089525 001	Jul 26, 1988
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PERRIGO NEW YORK	0.01%	A086810 001	Mar 04, 1982
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	0.025%	A086811 001	Mar 04, 1982
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PHARMAFAIR	0.01%	A088499 001	Aug 02, 1984
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	0.025%	A088506 001	Aug 02, 1984
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TARO	0.01%	A040035 001	Oct 31, 1994
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	0.01%	A087102 001	Apr 27, 1982
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	0.025%	A040042 001	Oct 31, 1994
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USL PHARMA	0.01%	A088757 001	Feb 11, 1985
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	0.025%	A088756 001	Mar 28, 1985
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FLUONID

ALLERGAN HERBERT	0.025%	A087156 002	Sep 06, 1984
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FLUOTREX

SAVAGE LABS	0.01%	A088174 001	May 06, 1983
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	0.025%	A088173 001	Mar 09, 1983
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SYNALAR-HP

MEDIMETRIKS PHARMS	0.2%	N016161 002	
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GEL; TOPICAL

FLUONID

ALLERGAN HERBERT	0.025%	A087300 001	May 27, 1982
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OIL; TOPICAL

FLUOCINOLONE ACETONIDE

AKORN	0.01%	A091514 001	Jun 25, 2015
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OIL/DROPS; OTIC

FLUOCINOLONE ACETONIDE

AKORN	0.01%	A202705 001	Sep 09, 2016
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OINTMENT; TOPICAL

FLUOCINOLONE ACETONIDE

PHARMADERM	0.025%	A088046 001	Dec 16, 1982
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PHARMAFAIR	0.025%	A088507 001	Feb 27, 1984
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USL PHARMA	0.025%	A088742 001	Feb 08, 1985
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FLUONID

ALLERGAN HERBERT	0.025%	A087157 001	Sep 06, 1984
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FLUOTREX

SAVAGE LABS	0.025%	A088172 001	Mar 09, 1983
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOCINOLONE ACETONIDE

SOLUTION; TOPICAL

FLUOCINOLONE ACETONIDE

ACTAVIS LABS UT INC	0.01%	A208386	001	Oct 21, 2016
ALLIED	0.01%	A088048	001	Dec 16, 1982
ALPHARMA US PHARMS	0.01%	A087159	001	Jun 16, 1982
BAUSCH AND LOMB	0.01%	A040059	001	Dec 20, 1993
COSETTE	0.01%	A207441	001	Sep 28, 2016
MORTON GROVE	0.01%	A088312	001	Jan 27, 1984
PHARMAFAIR	0.01%	A088449	001	Feb 08, 1984

FLUONID

ALLERGAN HERBERT	0.01%	A087158	001	Mar 17, 1983
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FLUOTREX

SAVAGE LABS	0.01%	A088171	001	Mar 09, 1983
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FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE

PERRIGO NEW YORK	0.05%	A071790	001	Jul 13, 1988
TELIGENT	0.05%	A211410	001	Oct 16, 2018

LIDEX

+ ALVOGEN 0.05%

N016908 002

SOLUTION; TOPICAL

FLUOCINONIDE

TARO	0.05%	A072857	001	Aug 02, 1989
TEVA	0.05%	A072511	001	Feb 07, 1989
TEVA PHARMS	0.05%	A072522	001	Sep 28, 1990

FLUORESCEIN SODIUM

INJECTABLE; INJECTION

FUNDUSCEIN-25

+ NOVARTIS 25% ** N017869 001

FLUOROMETHOLONE

CREAM; TOPICAL

OXYLONE

PHARMACIA AND UPJOHN	0.025%	N011748	001	
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SUSPENSION/DROPS; OPHTHALMIC

FLUOR-OP

NOVARTIS	0.1%	A070185	001	Feb 27, 1986
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FLUOROMETHOLONE ACETATE; TOBRAMYCIN

SUSPENSION/DROPS; OPHTHALMIC

TOBRASONE

EYEVANCE	0.1%; 0.3%	N050628	001	Jul 21, 1989
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FLUOROMETHOLONE; SULFACETAMIDE SODIUM

SUSPENSION/DROPS; OPHTHALMIC

FML-S

ALLERGAN	0.1%; 10%	N019525	001	Sep 29, 1989
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FLUOROURACIL

CREAM; TOPICAL

FLUOROPLEX

+ ALMIRALL 1% N016988 001

FLUOROURACIL

MYLAN	0.5%	A203122	001	Apr 20, 2015
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INJECTABLE; INJECTION

ADRUCIL

PHARMACIA AND UPJOHN	50MG/ML	A081222	001	Jun 28, 1991
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+ 50MG/ML N017959 001

TEVA PARENTERAL	50MG/ML	A040023	001	Oct 18, 1991
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	50MG/ML	A081225	001	Aug 28, 1991
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FLUOROURACIL

ABIC	50MG/ML	A088929	001	Mar 04, 1986
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ABRAXIS PHARM	50MG/ML	A089152	001	Mar 21, 1986
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	50MG/ML	A089428	001	Jan 12, 1987
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	50MG/ML	A089519	001	Mar 12, 1987
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AUROMEDICS PHARMA	500MG/10ML (50MG/ML)	A202668	001	Jul 17, 2012
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	1GM/20ML (50MG/ML)	A202668	002	Jul 17, 2012
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	2.5GM/50ML (50MG/ML)	A202669	001	Jul 17, 2012
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	5GM/100ML (50MG/ML)	A202669	002	Jul 17, 2012
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOROURACIL

INJECTABLE; INJECTION

FLUOROURACIL

BEDFORD	50MG/ML	A089508	001	Jan 26, 1988
EBEWE PHARMA	500MG/10ML (50MG/ML)	A040772	001	Aug 11, 2008
FRESENIUS KABI USA	50MG/ML	A040291	001	Mar 24, 1999
	50MG/ML	A040379	001	Nov 15, 2000
MARCHAR	50MG/ML	A087791	001	Jan 18, 1983
NOVAST LABS	500MG/10ML (50MG/ML)	A209219	001	Dec 12, 2019
	1GM/20ML (50MG/ML)	A209219	002	Dec 12, 2019
	2.5GM/50ML (50MG/ML)	A209271	002	Dec 11, 2019
	5GM/100ML (50MG/ML)	A209271	001	Dec 11, 2019
SANDOZ	2.5GM/50ML (50MG/ML)	A091299	001	May 02, 2011
	5GM/100ML (50MG/ML)	A091299	002	May 02, 2011
SMITH AND NEPHEW	50MG/ML	A088766	001	Dec 28, 1984
	50MG/ML	A088767	001	Dec 28, 1984
	50MG/ML	A089434	001	Mar 26, 1987
SPECTRUM PHARMS	50MG/ML	A087792	001	Oct 13, 1982
+	500MG/10ML (50MG/ML) **	N012209	001	
+	2.5GM/50ML (50MG/ML) **	N012209	002	Jul 29, 2016
TEVA PHARMS USA	2.5GM/50ML (50MG/ML)	A040334	001	Feb 25, 2000
	5GM/100ML (50MG/ML)	A040334	002	Feb 25, 2000

SOLUTION; TOPICAL

EFUDEX

+	BAUSCH	5% **	N016831	002
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FLUOROPLEX

	ELORAC	1%	N016765	001
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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

ACCORD HLTHCARE	EQ 10MG BASE	A202729	001	Aug 27, 2020
	EQ 20MG BASE	A202729	002	Aug 27, 2020
	EQ 40MG BASE	A202729	003	Aug 27, 2020
ANI PHARMS	EQ 10MG BASE	A076287	001	May 20, 2008
	EQ 20MG BASE	A076287	002	May 20, 2008
BARR	EQ 10MG BASE	A074803	002	Jan 30, 2002
	EQ 20MG BASE	A074803	001	Aug 02, 2001
	EQ 40MG BASE	A076251	001	May 18, 2005
BEXIMCO PHARMS USA	EQ 10MG BASE	A075807	001	Jan 29, 2002
	EQ 20MG BASE	A075807	002	Jan 29, 2002
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
GRANULES	EQ 10MG BASE	A078143	001	Jan 16, 2008
	EQ 20MG BASE	A078143	002	Jan 16, 2008
	EQ 40MG BASE	A078143	003	Jan 16, 2008
MYLAN	EQ 10MG BASE	A075207	001	Jan 30, 2002
	EQ 10MG BASE	A078045	001	Nov 17, 2008
	EQ 20MG BASE	A075207	002	Jan 30, 2002
	EQ 20MG BASE	A078045	002	Nov 17, 2008
	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
SANDOZ	EQ 10MG BASE	A077469	001	Nov 17, 2008
	EQ 20MG BASE	A077469	002	Nov 17, 2008
SPECGX LLC	EQ 10MG BASE	A075658	001	Jan 29, 2002
	EQ 20MG BASE	A075658	002	Jan 29, 2002
STRIDES PHARMA	EQ 10MG BASE	A076922	001	Dec 16, 2004
	EQ 20MG BASE	A076922	002	Dec 16, 2004
	EQ 40MG BASE	A076922	003	Dec 16, 2004

PROZAC

ELI LILLY AND CO	EQ 60MG BASE	N018936	004	Jun 15, 1999
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOXETINE HYDROCHLORIDE

CAPSULE;ORAL

SARAFEM

+ ELI LILLY AND CO	EQ 10MG BASE **	N018936 007	Jul 06, 2000
+	EQ 20MG BASE **	N018936 008	Jul 06, 2000

CAPSULE, DELAYED REL PELLETS;ORAL

FLUOXETINE HYDROCHLORIDE

BARR	EQ 90MG BASE	A076237 001	Mar 24, 2010
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PROZAC WEEKLY

+ LILLY	EQ 90MG BASE **	N021235 001	Feb 26, 2001
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SOLUTION;ORAL

FLUOXETINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC	EQ 20MG BASE/5ML	A075690 001	Jan 31, 2002
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AKORN	EQ 20MG BASE/5ML	A075525 001	Jun 27, 2002
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LANNETT CO INC	EQ 20MG BASE/5ML	A076458 001	May 14, 2004
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NOSTRUM LABS INC	EQ 20MG BASE/5ML	A075292 001	Feb 07, 2002
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SPECGX LLC	EQ 20MG BASE/5ML	A075920 001	Jan 29, 2002
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PROZAC

+ LILLY	EQ 20MG BASE/5ML **	N020101 001	Apr 24, 1991
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TABLET;ORAL

FLUOXETINE HYDROCHLORIDE

BARR	EQ 10MG BASE	A075810 001	Feb 01, 2002
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FOSUN PHARMA	EQ 10MG BASE	A076024 001	Jan 29, 2002
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G AND W LABS INC	EQ 60MG BASE	A212191 001	Jul 05, 2019
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IVAX SUB TEVA PHARMS	EQ 10MG BASE	A075865 001	Feb 28, 2002
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	EQ 40MG BASE	A075865 003	Aug 30, 2004
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RISING PHARMA	EQ 10MG BASE	A075755 001	Aug 02, 2001
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	EQ 20MG BASE	A075755 002	Aug 02, 2001
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PROZAC

+ LILLY	EQ 10MG BASE **	N020974 001	Mar 09, 1999
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+	EQ 20MG BASE **	N020974 002	Mar 09, 1999
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SARAFEM

+ APIL	EQ 10MG BASE	N021860 001	May 19, 2006
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+	EQ 15MG BASE **	N021860 002	May 19, 2006
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+	EQ 20MG BASE	N021860 003	May 19, 2006
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SELFEMRA

TEVA PHARMS USA	EQ 10MG BASE	A200151 001	Feb 03, 2014
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	EQ 15MG BASE	A200151 002	Feb 03, 2014
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	EQ 20MG BASE	A200151 003	Feb 03, 2014
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FLUOXYMESTERONE

TABLET;ORAL

ANDROID-F

VALEANT PHARM INTL	10MG	A087196 001	
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FLUOXYMESTERONE

UPSHER SMITH LABS	10MG	A088342 001	Oct 21, 1983
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VALEANT PHARM INTL	10MG	A088221 001	May 05, 1983
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WATSON LABS	2MG	A088260 001	Dec 06, 1983
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	5MG	A088265 001	Dec 06, 1983
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	10MG	A088309 001	Dec 06, 1983
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HALOTESTIN

PHARMACIA AND UPJOHN	2MG	N010611 002	
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	5MG	N010611 006	
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	10MG	N010611 010	
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ORA-TESTRYL

BRISTOL MYERS SQUIBB	2MG	N011359 001	
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	5MG	N011359 002	
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FLUPHENAZINE DECANOATE

INJECTABLE;INJECTION

FLUPHENAZINE DECANOATE

HOSPIRA	25MG/ML	A074966 001	Apr 16, 1998
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TEVA PARENTERAL	25MG/ML	A074795 001	Sep 10, 1996
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PROLIXIN DECANOATE

+ BRISTOL MYERS SQUIBB	25MG/ML **	N016727 001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUPHENAZINE ENANTHATE

INJECTABLE; INJECTION

PROLIXIN ENANTHATE

APOTHECON

25MG/ML **

N016110 001

FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

FLUPHENAZINE HYDROCHLORIDE

ANI PHARMS

5MG/ML

A073058 001 Aug 30, 1991

PERMITIL

SCHERING

5MG/ML **

N016008 001

PROLIXIN

APOTHECON

5MG/ML

A070533 001 Nov 07, 1985

ELIXIR; ORAL

FLUPHENAZINE HYDROCHLORIDE

ANI PHARMS

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

MYLAN

1MG

A089804 002 Aug 12, 1988

2.5MG

A089804 003 Aug 12, 1988

5MG

A089804 004 Aug 12, 1988

10MG

A089804 001 Aug 12, 1988

SANDOZ

1MG

A089586 002 Oct 16, 1987

2.5MG

A089586 003 Oct 16, 1987

5MG

A089586 004 Oct 16, 1987

10MG

A089586 001 Oct 16, 1987

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

CREAM; TOPICAL

CORDRAN SP

+ ALMIRALL

0.025%

N012806 003

LOTION; TOPICAL

FLURANDRENOLIDE

ALPHARMA US PHARMS

0.05%

A087203 001 Apr 29, 1982

OINTMENT; TOPICAL

CORDRAN

+ ALMIRALL

0.025% **

N012806 004

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

HALSEY

15MG

A071808 001 Jan 07, 1988

30MG

A071809 001 Jan 07, 1988

HERITAGE PHARMA

15MG

A071205 001 Nov 25, 1986

15MG

A072368 001 Mar 30, 1989

30MG

A071068 001 Nov 25, 1986

30MG

A072369 001 Mar 30, 1989

HIKMA

30MG

A071108 001 Dec 08, 1986

HIKMA INTL PHARMS

15MG

A071107 001 Dec 08, 1986

PUREPAC PHARM

15MG

A071927 001 Sep 09, 1987

30MG

A071551 001 Sep 09, 1987

RISING

15MG

A071717 002 Jul 31, 1991

30MG

A071717 001 Jul 31, 1991

STRIDES PHARMA

15MG

A070444 001 Mar 20, 1986

30MG

A070445 001 Mar 20, 1986

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

30MG

A070563 001 Jul 09, 1987

WARNER CHILCOTT

15MG

A071767 001 Dec 04, 1987

30MG

A071768 001 Dec 04, 1987

FLURBIPROFEN

TABLET; ORAL

ANSAID

PHARMACIA AND UPJOHN

50MG

N018766 002 Oct 31, 1988

100MG

N018766 003 Oct 31, 1988

FLURBIPROFEN

IVAX SUB TEVA PHARMS

50MG

A074411 001 May 31, 1995

100MG

A074411 002 May 31, 1995

MYLAN

50MG

A074358 001 Jun 20, 1994

100MG

A074358 002 Jun 20, 1994

PLIVA

50MG

A074647 001 Apr 01, 1997

100MG

A074647 002 Apr 01, 1997

RISING

50MG

A074448 001 Jul 28, 1995

100MG

A074448 002 Jul 28, 1995

SUN PHARM INDS INC

50MG

A075058 001 Apr 27, 2001

100MG

A075058 002 Apr 27, 2001

TEVA

50MG

A074405 002 May 24, 1995

100MG

A074405 001 May 24, 1995

THERAGEN

100MG

A074560 002 May 16, 1997

FLURBIPROFEN SODIUM

SOLUTION/DROPS; OPHTHALMIC

OCUFEN

+

ALLERGAN

0.03%

N019404 001 Dec 31, 1986

FLUTAMIDE

CAPSULE; ORAL

EULEXIN

+

SCHERING

125MG **

N018554 001 Jan 27, 1989

FLUTAMIDE

ACTAVIS LABS FL INC

125MG

A075820 001 Sep 18, 2001

CIPLA

125MG

A075780 001 Sep 19, 2001

MYLAN

125MG

A076224 001 May 09, 2003

YAOPHARMA CO LTD

125MG

A075818 001 Sep 18, 2001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

ANDA REPOSITORY 0.05% A076633 001 May 14, 2004

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

ARMONAIR RESPICLICK

+ TEVA PHARM 0.03MG/INH N208798 007 Jul 09, 2021

+ 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

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

FLUVASTATIN SODIUM

CAPSULE; ORAL

LESCOL

+ NOVARTIS EQ 20MG BASE ** N020261 001 Dec 31, 1993

+ EQ 40MG BASE ** N020261 002 Dec 31, 1993

TABLET, EXTENDED RELEASE; ORAL

FLUVASTATIN SODIUM

MYLAN EQ 80MG BASE A202458 001 Sep 11, 2015

FLUVOXAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

FLUVOXAMINE MALEATE

BIONPHARMA INC 100MG A212182 002 Sep 16, 2020

TORRENT 100MG A203240 001 Oct 31, 2014

150MG A203240 002 Oct 31, 2014

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 25MG A075897 001 Jan 25, 2001

25MG A075898 001 Mar 12, 2001

50MG A075897 002 Jan 25, 2001

50MG A075898 002 Mar 12, 2001

100MG A075897 003 Jan 25, 2001

100MG A075898 003 Mar 12, 2001

HERITAGE PHARMA 25MG A075894 001 Apr 18, 2001

50MG A075894 002 Apr 18, 2001

100MG A075894 003 Apr 18, 2001

MYLAN 25MG A075889 001 Nov 29, 2000

50MG A075889 002 Nov 29, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUVOXAMINE MALEATE

TABLET;ORAL

FLUVOXAMINE MALEATE

	50MG	A075950 001	Oct 15, 2001
	100MG	A075889 003	Nov 29, 2000
	100MG	A075950 002	Oct 15, 2001
PHARMA LIFE	25MG	A075900 001	Feb 23, 2006
	50MG	A075900 002	Feb 23, 2006
	100MG	A075900 003	Feb 23, 2006
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
TEVA	25MG	A075893 001	Sep 10, 2002
	50MG	A075893 002	Sep 10, 2002
	100MG	A075893 003	Sep 10, 2002
UPSHER SMITH LABS	25MG	A075887 001	Jan 05, 2001
	50MG	A075887 002	Jan 05, 2001
	100MG	A075887 003	Jan 05, 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	
HIKMA PHARMS	1MG	A080600 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	
VANGARD	1MG	A088730 001	Mar 23, 1984
VINTAGE	1MG	A040756 001	Jun 04, 2010
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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FOMEPIZOLE

INJECTABLE; INJECTION

ANTIZOL

+ PAR PHARM INC

1.5GM/1.5ML (1GM/ML) **

N020696 001 Dec 04, 1997

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

FOSAPREPITANT DIMEGLUMINE

MYLAN LABS LTD

EQ 115MG BASE/VIAL

A204015 001 Sep 05, 2019

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

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

ANI PHARMS

10MG; 12.5MG

A076608 001 Dec 03, 2004

10MG; 12.5MG

A077144 001 Aug 16, 2005

20MG; 12.5MG

A076608 002 Dec 03, 2004

20MG; 12.5MG

A077144 002 Aug 16, 2005

EMCURE PHARMS LTD

10MG; 12.5MG

A079025 001 Sep 17, 2010

20MG; 12.5MG

A079025 002 Sep 17, 2010

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

MONOPRIL-HCT

+ BRISTOL MYERS SQUIBB

10MG; 12.5MG **

N020286 002 Nov 30, 1994

+ 20MG; 12.5MG **

N020286 001 Nov 30, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

FOSPHENYTOIN SODIUM

AM REGENT

EQ 50MG PHENYTOIN NA/ML

A078277 001 Aug 06, 2007

EQ 50MG PHENYTOIN NA/ML

A090099 001 May 13, 2010

AMNEAL

EQ 50MG PHENYTOIN NA/ML

A078476 001 Mar 18, 2008

APOTEX INC

EQ 50MG PHENYTOIN NA/ML

A078126 001 Aug 06, 2007

DR REDDYS

EQ 50MG PHENYTOIN NA/ML

A076886 001 Aug 06, 2007

HOSPIRA

EQ 50MG PHENYTOIN NA/ML

A078158 001 Aug 06, 2007

SOLUTION; INTRAVENOUS

SESQUIENT

+ LUPIN

EQ 100MG PHENYTOIN NA/2ML (EQ 50MG
PHENYTOIN NA/ML)

N210864 001 Nov 05, 2020

+

EQ 500MG PHENYTOIN NA/10ML (EQ 50MG
PHENYTOIN NA/ML)

N210864 002 Nov 05, 2020

FOSPROPOFOL DISODIUM

SOLUTION; INTRAVENOUS

LUSEDRA

EISAI INC

1050MG/30ML (35MG/ML)

N022244 001 Dec 12, 2008

FULVESTRANT

INJECTABLE; INTRAMUSCULAR

FULVESTRANT

ZYDUS PHARMS

50MG/ML

A215234 001 Jul 29, 2021

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

+ AM REGENT

10MG/ML **

N018579 001 Nov 30, 1983

AMNEAL PHARMS CO

10MG/ML

A207552 001 Jul 20, 2016

ASTRAZENECA

10MG/ML

A070014 001 Sep 09, 1985

HIKMA

10MG/ML

A071439 001 Sep 14, 1990

10MG/ML

N018267 001

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

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

WYETH AYERST

10MG/ML

N018670 001 Jul 20, 1982

LASIX

+ SANOFI AVENTIS US

10MG/ML **

N016363 001

SOLUTION; ORAL

LASIX

SANOFI AVENTIS US

10MG/ML

N017688 001

TABLET; ORAL

FUROSEMIDE

ANI PHARMS

20MG

A071379 001 Jan 02, 1987

40MG

A070413 001 Feb 26, 1986

80MG

A071594 001 Feb 09, 1988

AVET

20MG

N018413 001 Nov 30, 1983

40MG

N018413 002 Nov 30, 1983

INTL MEDICATION

20MG

N018753 001 Feb 28, 1984

40MG

N018753 002 Feb 28, 1984

KALAPHARM

20MG

N018868 001 Jun 28, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FUROSEMIDETABLET;ORAL
FUROSEMIDE

	40MG	N018868 002	Jun 28, 1983
SANDOZ	40MG	N018750 002	Jul 30, 1984
STRIDES PHARMA	20MG	N018415 001	Jul 27, 1982
	40MG	N018415 002	Jul 27, 1982
	80MG	N018415 003	Nov 26, 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	N018369 001	May 14, 1982
	40MG	A070450 001	Nov 22, 1985
	40MG	N018369 002	May 14, 1982
WATSON LABS TEVA	20MG	A070449 001	Nov 22, 1985
	80MG	A070528 001	Jan 07, 1986

GABAPENTINCAPSULE;ORAL
GABAPENTIN

HIKMA	100MG	A078150 001	Sep 25, 2007
	300MG	A078150 002	Sep 25, 2007
	400MG	A078150 003	Sep 25, 2007
JIANGSU PHARMS	100MG	A091008 001	Oct 26, 2017
	300MG	A091008 002	Oct 26, 2017
	400MG	A091008 003	Oct 26, 2017
MYLAN	100MG	A090158 001	Feb 14, 2011
	300MG	A090158 002	Feb 14, 2011
	400MG	A090158 003	Feb 14, 2011
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
TEVA PHARMS	100MG	A075435 001	Oct 08, 2004
	300MG	A075435 002	Oct 08, 2004
	400MG	A075435 003	Oct 08, 2004
WATSON LABS	100MG	A075485 003	May 11, 2007
	300MG	A075485 002	May 11, 2007
	400MG	A075485 001	May 11, 2007

SOLUTION;ORAL

GABAPENTIN

NOVITIUM PHARMA	250MG/5ML	A211330 001	Dec 03, 2019
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TABLET;ORAL

GABAPENTIN

HIKMA PHARMS	600MG	A078782 001	Jul 21, 2011
	800MG	A078782 002	Jul 21, 2011
INVATECH	600MG	A076877 001	Jul 06, 2006
	800MG	A076877 002	Jul 06, 2006
IVAX SUB TEVA PHARMS	600MG	A076017 004	Apr 29, 2005
	800MG	A076017 005	Apr 29, 2005
LUPIN LTD	600MG	A209306 001	Aug 24, 2018
	800MG	A209306 002	Aug 24, 2018
MYLAN PHARMS INC	600MG	A090335 001	Jun 01, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GABAPENTINTABLET; ORAL
GABAPENTIN

	800MG	A090335 002	Jun 01, 2010
RANBAXY	600MG	A076605 001	Sep 14, 2005
	800MG	A076605 002	Sep 14, 2005
SANDOZ	600MG	A076120 001	Jan 27, 2006
	800MG	A076120 002	Jan 27, 2006
TEVA	600MG	A075827 001	Dec 15, 2004
	800MG	A075827 002	Dec 15, 2004
TEVA PHARMS USA	600MG	A205807 001	Mar 10, 2017
	800MG	A205807 002	Mar 10, 2017

GADODIAMIDEINJECTABLE; INJECTION
OMNISCAN

GE HEALTHCARE	14.35GM/50ML (287MG/ML)	N022066 001	Sep 05, 2007
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GADOFOSVESET TRISODIUMSOLUTION; INTRAVENOUS
ABLAVAR

LANTHEUS MEDCL	2440MG/10ML (244MG/ML)	N021711 001	Dec 22, 2008
	3660MG/15ML (244MG/ML)	N021711 002	Dec 22, 2008

GADOPENTETATE DIMEGLUMINEINJECTABLE; INJECTION
MAGNEVIST

+ BAYER HLTHCARE	469.01MG/ML	N019596 001	Jun 02, 1988
+	469.01MG/ML	N021037 001	Mar 10, 2000

GADOVERSETAMIDEINJECTABLE; 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 HYDROBROMIDECAPSULE, 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
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TABLET; ORAL

GALANTAMINE HYDROBROMIDE

APOTEX INC	EQ 4MG BASE	A077781 001	Sep 27, 2011
	EQ 8MG BASE	A077781 002	Sep 27, 2011
	EQ 12MG BASE	A077781 003	Sep 27, 2011
HERITAGE PHARMA	EQ 4MG BASE	A077585 001	Sep 15, 2009
	EQ 4MG BASE	A077587 001	Jul 09, 2009
	EQ 8MG BASE	A077585 002	Sep 15, 2009
	EQ 8MG BASE	A077587 002	Jul 09, 2009
	EQ 12MG BASE	A077585 003	Sep 15, 2009
	EQ 12MG BASE	A077587 003	Jul 09, 2009
HIKMA	EQ 4MG BASE	A077608 001	Feb 11, 2009
	EQ 8MG BASE	A077608 002	Feb 11, 2009
	EQ 12MG BASE	A077608 003	Feb 11, 2009
MYLAN	EQ 4MG BASE	A077590 001	May 29, 2009
	EQ 4MG BASE	A077603 001	Aug 28, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GALANTAMINE HYDROBROMIDE

TABLET; ORAL

GALANTAMINE HYDROBROMIDE

EQ 8MG BASE	A077590 002	May 29, 2009
EQ 8MG BASE	A077603 002	Aug 28, 2008
EQ 12MG BASE	A077590 003	May 29, 2009
EQ 12MG BASE	A077603 003	Aug 28, 2008

RAZADYNE

+ JANSSEN PHARMS	EQ 4MG BASE	N021169 001	Feb 28, 2001
+	EQ 8MG BASE	N021169 002	Feb 28, 2001
+	EQ 12MG BASE	N021169 003	Feb 28, 2001

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
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NEOSCAN

GE HEALTHCARE	2mCi/ML	N017655 001
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GALLIUM NITRATE

INJECTABLE; INJECTION

GANITE

CHAPTER 7 TRUSTEE	25MG/ML **	N019961 002	Jan 17, 1991
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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
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GANCICLOVIR SODIUM

INJECTABLE; INJECTION

CYTOVENE

+ CHEPLAPHARM	EQ 500MG BASE/VIAL	N019661 001	Jun 23, 1989
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GANCICLOVIR SODIUM

AM REGENT	EQ 500MG BASE/VIAL	A202624 001	Sep 18, 2013
CUSTOPHARM INC	EQ 500MG BASE/VIAL	A212001 001	Jun 20, 2019
MYLAN LABS LTD	EQ 500MG BASE/VIAL	A204560 001	Nov 17, 2017

GATIFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

GATIFLOXACIN

APOTEX INC	0.3%	A079084 001	Aug 19, 2011
MYLAN	0.5%	A206446 001	Jun 08, 2018

GEFITINIB

TABLET; ORAL

IRESSA

ASTRAZENECA	250MG	N021399 001	May 05, 2003
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GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

ACTAVIS INC	200MG/5.26ML (38MG/ML)	A204549 001	Apr 11, 2016
	1GM/26.3ML (38MG/ML)	A204549 002	Apr 11, 2016
	2GM/52.6ML (38MG/ML)	A204549 003	Apr 11, 2016
ACTAVIS TOTOWA	EQ 200MG BASE/VIAL	A079160 001	Jul 25, 2011
	EQ 1GM BASE/VIAL	A079160 002	Jul 25, 2011
	EQ 2GM BASE/VIAL	A079160 003	Jul 28, 2016
AM REGENT	EQ 200MG BASE/VIAL	A202031 001	May 07, 2013
	EQ 1GM BASE/VIAL	A202031 002	May 07, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

APOTEX	200MG/5.26ML (38MG/ML)	A206776 001	May 23, 2017
	1GM/26.3ML (38MG/ML)	A206776 002	May 23, 2017
	2GM/52.6ML (38MG/ML)	A206776 003	May 23, 2017
EMCURE PHARMS LTD	EQ 200MG BASE/VIAL	A202063 001	Sep 11, 2012
	EQ 1GM BASE/VIAL	A202063 002	Sep 11, 2012
HAMELN RDS GMBH	EQ 200MG BASE/VIAL	A090663 001	Sep 10, 2012
	EQ 1GM BASE/VIAL	A090663 002	Sep 10, 2012
MYLAN LABS LTD	EQ 200MG BASE/VIAL	A200145 001	Jul 25, 2011
	EQ 1GM BASE/VIAL	A200145 002	Jul 25, 2011
	EQ 2GM BASE/VIAL	A200145 003	Jul 25, 2011
NOVAST LABS	200MG/5.26ML (38MG/ML)	A210383 001	Feb 14, 2019
	1GM/26.3ML (38MG/ML)	A210383 002	Feb 14, 2019
	2GM/52.6ML (38MG/ML)	A210383 003	Feb 14, 2019
SAGENT PHARMS	EQ 200MG BASE/VIAL	A091597 001	May 07, 2013
	EQ 1GM BASE/VIAL	A091597 002	May 07, 2013
TEVA PHARMS	EQ 200MG BASE/VIAL	A077983 002	Jan 25, 2011
	EQ 1GM BASE/VIAL	A077983 001	Jan 25, 2011
GEMZAR			
+ LILLY	EQ 200MG BASE/VIAL	N020509 001	May 15, 1996
+	EQ 1GM BASE/VIAL	N020509 002	May 15, 1996

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

HIKMA PHARMS	600MG	A078599 001	Aug 16, 2010
MYLAN	600MG	A074452 001	Feb 16, 1995
PUREPAC PHARM	600MG	A074360 001	Aug 31, 1994
SUN PHARM INDS INC	600MG	A079239 001	Dec 29, 2008
TEVA	600MG	A074256 001	Oct 31, 1993
WATSON LABS	600MG	A074156 001	Oct 24, 1994
	600MG	A074442 001	Apr 28, 1995
YAOPHARMA CO LTD	600MG	A074615 001	Sep 29, 1995

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GENTAMICIN SULFATE

INJECTABLE; INJECTION

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
HIKMA	EQ 10MG BASE/ML	A062251 002	
	EQ 40MG BASE/ML	A062251 001	
HOSPIRA	EQ 10MG BASE/ML	A062612 004	Feb 20, 1986
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	
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
	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	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GENTAMICIN SULFATE

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

1MG

A077624 001 Nov 28, 2005

2MG

A077486 002 Feb 10, 2006

2MG

A077624 002 Nov 28, 2005

4MG

A077486 003 Feb 10, 2006

4MG

A077624 003 Nov 28, 2005

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

TEVA

1MG

A076802 001 Oct 06, 2005

2MG

A076802 002 Oct 06, 2005

4MG

A076802 003 Oct 06, 2005

WATSON LABS

1MG

A077280 001 Feb 03, 2006

2MG

A077280 002 Feb 03, 2006

4MG

A077280 003 Feb 03, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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	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
	BEXIMCO PHARMS USA	5MG	A074542	001	Jun 20, 1995
		10MG	A074542	002	Jun 20, 1995
	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
	WATSON LABS	5MG	A074370	001	Nov 22, 1994
		10MG	A074370	002	Nov 22, 1994

GLUCOTROL

+	PFIZER	2.5MG	N017783	003	May 11, 1993
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TABLET, EXTENDED RELEASE; ORAL

GLIPIZIDE

	MYLAN	2.5MG	A202298	001	May 19, 2015
		5MG	A202298	002	May 19, 2015
		10MG	A202298	003	May 19, 2015
	PAR PHARM	5MG	A076159	002	Sep 20, 2013
		10MG	A076159	001	Sep 20, 2013

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

	MYLAN	2.5MG;250MG	A078083	001	Apr 12, 2007
		2.5MG;500MG	A078083	002	Apr 12, 2007
		5MG;500MG	A078083	003	Apr 12, 2007
	SUN PHARM INDS INC	2.5MG;250MG	A077620	001	Jan 11, 2008
		2.5MG;500MG	A077620	002	Jan 11, 2008
		5MG;500MG	A077620	003	Jan 11, 2008

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	
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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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GLUTETHIMIDE

TABLET;ORAL

GLUTETHIMIDE

LANNETT	250MG	A083475	001	
	500MG	A085571	001	
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
AUROBINDO PHARMA	1.25MG	A077537	001	Oct 18, 2007
	2.5MG	A077537	002	Oct 18, 2007
	5MG	A077537	003	Oct 18, 2007

GLYBURIDE (MICRONIZED)

MYLAN	1.5MG	A074792	001	Jun 26, 1998
	3MG	A074792	002	Jun 26, 1998
	6MG	A074792	003	Aug 17, 1999
SANOFI AVENTIS US	1.5MG	N020055	001	Apr 17, 1992
	3MG	N020055	002	Apr 17, 1992
	6MG	N020055	003	Mar 08, 2000
STRIDES PHARMA	1.5MG	A074591	001	Dec 22, 1997
	3MG	A074591	002	Dec 22, 1997
	4.5MG	A074591	003	Dec 22, 1997
	6MG	A074591	004	Dec 22, 1997
YAOPHARMA CO LTD	1.5MG	A075174	001	Jun 22, 1998
	3MG	A075174	002	Jun 22, 1998

GLYNASE

+ PFIZER 4.5MG ** N020051 003 Sep 24, 1993

MICRONASE

+ PFIZER 1.25MG ** N017498 001 May 01, 1984

+ PFIZER 2.5MG N017498 002 May 01, 1984

+ PFIZER 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

HERITAGE PHARMS INC 1.25MG;250MG A079009 001 Jun 03, 2009

HERITAGE PHARMS INC 2.5MG;500MG A079009 002 Jun 03, 2009

HERITAGE PHARMS INC 5MG;500MG A079009 003 Jun 03, 2009

IMPAX LABS INC 1.25MG;250MG A076731 001 Nov 19, 2004

IMPAX LABS INC 2.5MG;500MG A076731 002 Nov 19, 2004

IMPAX LABS INC 5MG;500MG A076731 003 Nov 19, 2004

TEVA 1.25MG;250MG A076821 001 Jan 27, 2005

TEVA 2.5MG;500MG A076821 002 Jan 27, 2005

TEVA 5MG;500MG A076821 003 Jan 27, 2005

GLYCEROL PHENYLBUTYRATE

LIQUID;ORAL

GLYCEROL PHENYLBUTYRATE

PAR PHARM INC 1.1GM/ML A205742 001 Dec 02, 2021

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

+ HIKMA	0.2MG/ML **	N017558 001	
ROBINS AH	0.2MG/ML	N014764 001	

POWDER; INHALATION

SEEBRI

+ NOVARTIS	15.6MCG/INH	N207923 001	Oct 29, 2015
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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	

GLYCOPYRROLATE; INDACATEROL MALEATE

POWDER; INHALATION

UTIBRON

+ NOVARTIS	15.6MCG/INH; 27.5MCG/INH	N207930 001	Oct 29, 2015
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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

HIKMA	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

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	
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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
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WATSON LABS	0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML	A062788 001	Jun 11, 1987
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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
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GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE

AM REGENT	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A091274 001	Sep 22, 2010
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
MYLAN LABS LTD	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)	A203454 001	Apr 04, 2017
	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A203454 002	Apr 04, 2017
	EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	A203453 001	Jan 31, 2017
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
YUNG SHIN PHARM	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)	A202647 001	Mar 06, 2020
	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A202648 001	Jun 29, 2020
	EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	A202648 002	Jun 29, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE

DR REDDYS	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

INTRA SANA LABS	EQ 2MG BASE/10ML	A078334 001	Feb 28, 2008
KYTRIL			
+ ROCHE	EQ 2MG BASE/10ML **	N021238 001	Jun 27, 2001

TABLET; ORAL

GRANISETRON HYDROCHLORIDE

BARR	EQ 1MG BASE	A078221 001	Dec 31, 2007
EPIC PHARMA LLC	EQ 1MG BASE	A078260 001	Dec 31, 2007
MYLAN	EQ 1MG BASE	A078725 001	Jan 30, 2008
TEVA PHARMS	EQ 1MG BASE	A078080 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	
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TABLET; ORAL

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	
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GRISEOFULVIN, MICROSIZE

SUSPENSION; ORAL

GRIFULVIN V

VALEANT LUXEMBOURG	125MG/5ML **	A062483 001	Jan 26, 1984
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TABLET; ORAL

GRIFULVIN V

VALEANT LUXEMBOURG	500MG	A062279 003	
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GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET; ORAL

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GUAIFENESIN; HYDROCODONE BITARTRATE

SOLUTION; ORAL

FLOWTUSS

BKK PHARMS 200MG/5ML; 2.5MG/5ML N022424 001 May 14, 2015

OBREDON

+ SOVEREIGN PHARMS 200MG/5ML; 2.5MG/5ML N205474 001 Nov 14, 2014

TABLET; ORAL

XTRELUS

+ ECI PHARMS LLC 400MG; 5MG N208085 001 Apr 25, 2018

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 EQ 4MG BASE A074149 001 Apr 07, 1995

EQ 4MG BASE A074267 001 Jun 01, 1994

EQ 8MG BASE A074149 002 Apr 07, 1995

EQ 8MG BASE A074267 002 Jun 01, 1994

CHARTWELL RX EQ 4MG BASE A074517 001 Sep 30, 1998

EQ 8MG BASE A074517 002 Sep 30, 1998

WATSON LABS EQ 4MG BASE A074025 001 Feb 28, 1994

EQ 8MG BASE A074025 002 Feb 28, 1994

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

AUROBINDO PHARMA USA EQ 1MG BASE A074796 001 Jan 27, 1997

EQ 2MG BASE A074796 002 Jan 27, 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

MYLAN EQ 1MG BASE A202578 001 Jun 02, 2015

EQ 2MG BASE A202578 002 Jun 02, 2015

EQ 3MG BASE A202578 003 Jun 02, 2015

EQ 4MG BASE A202578 004 Jun 02, 2015

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GUANFACINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

GUANFACINE HYDROCHLORIDE

YICHANG HUMANWELL	EQ 1MG BASE	A213428 001	Nov 25, 2020
	EQ 2MG BASE	A213428 002	Nov 25, 2020
	EQ 3MG BASE	A213428 003	Nov 25, 2020
	EQ 4MG BASE	A213428 004	Nov 25, 2020

GUANIDINE HYDROCHLORIDE

TABLET;ORAL

GUANIDINE HYDROCHLORIDE

MERCK SHARP DOHME	125MG	N001546 001	
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HALAZEPAM

TABLET;ORAL

PAXIPAM

SCHERING	20MG	N017736 003	
	40MG	N017736 004	

HALCINONIDE

CREAM;TOPICAL

HALOG

WESTWOOD SQUIBB	0.025%	N017818 001	
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HALOG-E

SUN PHARM INDS INC	0.1%	N018234 001	
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OINTMENT;TOPICAL

HALOG

BRISTOL MYERS SQUIBB	0.025%	N018125 001	
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HALOBETASOL PROPIONATE

CREAM;TOPICAL

ULTRAVATE

+ SUN PHARM INDS INC	0.05% **	N019967 001	Dec 27, 1990
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LOTION;TOPICAL

HALOBETASOL PROPIONATE

PADAGIS ISRAEL	0.05%	A211464 001	Jun 03, 2020
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OINTMENT;TOPICAL

HALOBETASOL PROPIONATE

COSETTE	0.05%	A077721 001	Sep 07, 2006
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FOUGERA PHARMS	0.05%	A076903 001	Dec 16, 2004
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ULTRAVATE

+ SUN PHARM INDS INC	0.05%	N019968 001	Dec 17, 1990
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HALOFANTRINE HYDROCHLORIDE

TABLET;ORAL

HALFAN

GLAXOSMITHKLINE	250MG	N020250 001	Jul 24, 1992
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HALOPERIDOL

TABLET;ORAL

HALDOL

+ ORTHO MCNEIL	0.5MG **	N015921 001	
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	1MG **	N015921 002	
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	2MG **	N015921 003	
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	5MG **	N015921 004	
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	10MG **	N015921 005	
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	20MG **	N015921 006	Feb 02, 1982
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HALDOL SOLUTAB

ORTHO MCNEIL PHARM	1MG	N017079 001	
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HALOPERIDOL

CYCLE PHARMS LTD	0.5MG	A071128 001	Feb 17, 1987
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	1MG	A071129 001	Feb 17, 1987
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	20MG	A071133 001	May 12, 1987
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DURAMED PHARMS BARR	0.5MG	A071216 001	Dec 04, 1986
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	1MG	A071217 001	Dec 04, 1986
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	2MG	A071218 001	Dec 04, 1986
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	5MG	A071219 001	Dec 04, 1986
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	10MG	A071220 001	Jul 07, 1987
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	20MG	A071221 001	Jul 07, 1987
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LEDERLE	0.5MG	A072727 001	Sep 19, 1989
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	1MG	A072728 001	Sep 19, 1989
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	2MG	A072729 001	Sep 19, 1989
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL

	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
SANDOZ	2MG	A071208 001	Nov 17, 1986
SCS	0.5MG	A070720 001	Jun 10, 1986
	1MG	A070721 001	Jun 10, 1986
	2MG	A070722 001	Jun 10, 1986
	5MG	A070723 001	Jun 10, 1986
	10MG	A070724 001	Jun 10, 1986
	20MG	A070725 001	Sep 24, 1986
STRIDES PHARMA	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

HALOPERIDOL

	EQ 2MG BASE/ML **	N015922 001	
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
TEVA PHARMS	EQ 2MG BASE/ML	A071617 001	Dec 01, 1988
HALOPERIDOL INTENSOL			
HIKMA	EQ 2MG BASE/ML	A072045 001	Apr 12, 1988

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HALOPERIDOL LACTATE

INJECTABLE; INJECTION

HALDOL

+ JANSSEN PHARMS EQ 5MG BASE/ML N015923 001

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
TEVA PHARMS USA	EQ 5MG BASE/ML	A076035	001	Aug 29, 2001
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

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
PARKE DAVIS	10 UNITS/ML	N017346	006
SMITH AND NEPHEW	10 UNITS/ML	A087904	001
	10 UNITS/ML	A087958	001
	10 UNITS/ML	A088458	001
	10 UNITS/ML	A088580	001
	100 UNITS/ML	A087906	001
	100 UNITS/ML	A087959	001
	100 UNITS/ML	A088460	001
	100 UNITS/ML	A088581	001
SOLOPAK	10 UNITS/ML	A087903	001
	10 UNITS/ML	A088457	001
	100 UNITS/ML	A087905	001
	100 UNITS/ML	A088459	001
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

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	
CASI PHARMS INC	5,000 UNITS/ML	A091659 001	Jun 08, 2011
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	
DR REDDYS	1,000 UNITS/ML	A040007 001	Jun 07, 1996
	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	
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	
HIKMA	5,000 UNITS/0.5ML	N017037 013	Apr 07, 1986
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
NANJING KING-FRIEND	5,000 UNITS/ML	A212061 001	Jul 15, 2020
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	
	10,000 UNITS/ML	N017346 004	
	20,000 UNITS/ML	N017346 005	
+ PFIZER	10,000 UNITS/ML	N201370 003	Jul 21, 2011
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 INC	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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

+		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
		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
		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			
DR REDDYS		1,000 UNITS/ML	A089464 001	Jun 03, 1986
HOSPIRA		2,000 UNITS/ML	N005264 013	Apr 07, 1986
		2,500 UNITS/ML	N005264 014	Apr 07, 1986
NANJING KING-FRIEND		10,000 UNITS/ML	A212060 001	Apr 02, 2020
PHARMA SERVE NY		1,000 UNITS/ML	A086129 001	
SHENZHEN TECHDOW		1,000 UNITS/ML	A202732 001	Jun 12, 2014

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

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

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

XTTRIUM

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

XTTRIUM

3% N019055 001 Nov 30, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEXACHLOROPHENE

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

HEXOCYCLIUM METHYLSULFATE

TABLET; ORAL			
TRAL			
ABBVIE	25MG		N010599 001

HEXYLCAINE HYDROCHLORIDE

SOLUTION; TOPICAL			
CYCLAINE			
MERCK	5%		N008472 001

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

IMPLANT; SUBCUTANEOUS			
VANTAS			
+	ENDO PHARM	50MG	N021732 001 Oct 12, 2004
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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE

IVAX SUB TEVA PHARMS	1.5MG/5ML; 5MG/5ML	A040285	001	Jul 19, 1999
NOSTRUM LABS INC	1.5MG/5ML; 5MG/5ML	A210663	001	Jun 11, 2019
TORRENT	1.5MG/5ML; 5MG/5ML	A204765	001	Mar 06, 2017

HYDROCODONE

HALSEY	1.5MG/5ML; 5MG/5ML	A088066	001	Jun 28, 1985
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TABLET; ORAL

HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE

ACTAVIS ELIZABETH	1.5MG; 5MG	A040295	001	Dec 01, 2000
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TUSSIGON

KING PHARMS	1.5MG; 5MG	A088508	001	Jul 30, 1985
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HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

APRESOLINE

+ NOVARTIS	20MG/ML **	N008303	003	
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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	
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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
	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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDRALAZINE HYDROCHLORIDE

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDE

	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
STRIDES PHARMA	10MG	A200770	004	Jun 25, 2019
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	
	10MG	A209251	001	Jul 09, 2018
	25MG	A083560	001	
	25MG	A209251	002	Jul 09, 2018
	50MG	A083561	001	
	50MG	A085088	001	
	50MG	A209251	003	Jul 09, 2018
	100MG	A209251	004	Jul 09, 2018
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

STRIDES PHARMA	100MG; 50MG	A088961	001	Oct 21, 1985
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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

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

ALEMBIC PHARMS LTD	12.5MG	A200645	001	Nov 30, 2010
APOTEX	12.5MG	A078389	001	May 16, 2008
HIKMA INTL PHARMS	12.5MG	A077885	001	Nov 26, 2007
IVAX SUB TEVA PHARMS	12.5MG	A077005	001	Jul 13, 2005
LANNETT CO INC	12.5MG	A091662	001	Jan 27, 2012

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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE

TABLET; ORAL

HYDROCHLOROTHIAZIDE

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
	50MG	A084878	001	
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
	25MG	A040735	002	Jan 23, 2007
	50MG	A040735	003	Jan 23, 2007
PVT FORM	50MG	A086597	001	
ROXANE	25MG	A085004	001	
	50MG	A084536	002	
	50MG	A085005	001	
SOLVAY	25MG	A085323	001	
SUN PHARM INDS INC	12.5MG	A040857	001	May 30, 2008
	25MG	A040810	001	Mar 27, 2007
	50MG	A040810	002	Mar 27, 2007
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
	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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE

TABLET; ORAL

HYDRODIURIL

+	MERCK	25MG **	N011835	003
+		50MG **	N011835	006
+		100MG **	N011835	007

ORETIC

	ABBVIE	25MG	N011971	001
		50MG	N011971	002

ZIDE

	SOLVAY	50MG	A083925	001
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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

	APOTEX INC	12.5MG; 150MG	A201505	001	Oct 15, 2012
		12.5MG; 300MG	A201505	002	Oct 15, 2012
	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

	COREPHARMA	12.5MG; 10MG	A076674	001	Oct 05, 2004
		12.5MG; 20MG	A076674	002	Oct 05, 2004
		25MG; 20MG	A076674	003	Oct 05, 2004
	HERITAGE PHARMA	12.5MG; 10MG	A075776	001	Jul 01, 2002
		12.5MG; 20MG	A075776	002	Jul 01, 2002
		25MG; 20MG	A075776	003	Jul 01, 2002
	HIKMA INTL PHARMS	12.5MG; 10MG	A076265	001	Jul 08, 2002
		12.5MG; 20MG	A076265	002	Jul 08, 2002
		25MG; 20MG	A076265	003	Jul 08, 2002
	MYLAN	12.5MG; 10MG	A076113	001	Jul 01, 2002
		12.5MG; 20MG	A076113	002	Jul 01, 2002
		25MG; 20MG	A076113	003	Jul 01, 2002
	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
		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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET;ORAL

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
HIKMA	12.5MG;50MG	A077732	002	Oct 06, 2010
	12.5MG;100MG	A077732	001	Apr 06, 2010
	25MG;100MG	A077732	003	Oct 06, 2010
MYLAN	12.5MG;50MG	A091652	001	Oct 06, 2010
	12.5MG;100MG	A091652	002	Apr 06, 2010
	25MG;100MG	A091652	003	Oct 06, 2010
TORRENT PHARMS	12.5MG;50MG	A090528	001	Oct 06, 2010
	12.5MG;100MG	A090528	003	Apr 06, 2010
	25MG;100MG	A090528	002	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
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ALDORIL 25

MERCK	25MG;250MG	N013402	002
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ALDORIL D30

MERCK	30MG;500MG	N013402	003
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ALDORIL D50

MERCK	50MG;500MG	N013402	004
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METHYLDOPA AND HYDROCHLOROTHIAZIDE

CHARTWELL RX	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
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
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
RISING	15MG;250MG	A070265	002	Jan 23, 1986
	25MG;250MG	A070265	001	Jan 23, 1986
SANDOZ	15MG;250MG	A070829	001	Mar 09, 1987
	25MG;250MG	A070830	001	Mar 09, 1987
STRIDES PHARMA	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
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
WATSON LABS	30MG;500MG	A070367	001	Mar 19, 1986
	30MG;500MG	A071069	001	Jan 19, 1989

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

METHYLDOPA AND HYDROCHLOROTHIAZIDE

30MG; 500MG	A071922	001	Aug 29, 1988
50MG; 500MG	A070368	001	Apr 16, 1986
50MG; 500MG	A070960	001	Feb 06, 1989
50MG; 500MG	A071923	001	Aug 29, 1988

HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

DUTOPROL

+ CONCORDIA	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

+ VALIDUS PHARMS	25MG; 100MG		N018303	002	Dec 31, 1984
+	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
HERITAGE PHARMS INC	12.5MG; 7.5MG		A202150	001	Mar 07, 2014
	12.5MG; 15MG		A202150	002	Mar 07, 2014
	25MG; 15MG		A202150	003	Mar 07, 2014

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; OLMESARTAN MEDOXOMIL

TABLET; ORAL

OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE

MYLAN	12.5MG; 20MG		A078827	001	Oct 26, 2016
	12.5MG; 40MG		A078827	002	Oct 26, 2016
	25MG; 40MG		A078827	003	Oct 26, 2016
TEVA PHARMS USA	12.5MG; 20MG		A200532	001	Apr 24, 2017
	25MG; 40MG		A200532	003	Apr 24, 2017

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
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INDERIDE LA 160/50

WYETH AYERST	50MG; 160MG		N019059	003	Jul 03, 1985
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INDERIDE LA 80/50

WYETH AYERST	50MG; 80MG		N019059	001	Jul 03, 1985
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TABLET; ORAL

INDERIDE-40/25

+ WYETH PHARMS INC	25MG; 40MG	**	N018031	001	
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INDERIDE-80/25

+ WYETH PHARMS INC	25MG; 80MG	**	N018031	002	
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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	25MG; 40MG		A070705	002	Oct 01, 1986
	25MG; 40MG		A072043	002	Mar 14, 1988
	25MG; 80MG		A070705	001	Oct 01, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

	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
MYLAN	25MG;40MG	A070947 002	Mar 04, 1987
	25MG;80MG	A070947 001	Apr 01, 1987
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

MYLAN	12.5MG;EQ 10MG BASE	A077093 001	Mar 28, 2005
	12.5MG;EQ 20MG BASE	A077093 002	Mar 28, 2005
	25MG;EQ 20MG BASE	A077093 003	Mar 28, 2005
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

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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL

SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE

ASCOT	25MG; 25MG	A088025	001	Nov 23, 1984
CHARTWELL RX	25MG; 25MG	A086881	001	
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	

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; TELMISARTAN

TABLET; ORAL

TELMISARTAN AND HYDROCHLOROTHIAZIDE

MACLEODS PHARMS LTD	12.5MG; 40MG	A204169	001	Nov 02, 2015
	12.5MG; 80MG	A204169	002	Nov 02, 2015
	25MG; 80MG	A204169	003	Nov 02, 2015
MYLAN	12.5MG; 40MG	A091648	001	Feb 25, 2014
	12.5MG; 80MG	A091648	002	Feb 25, 2014
	25MG; 80MG	A091648	003	Feb 25, 2014
TORRENT	12.5MG; 40MG	A201192	001	Feb 25, 2014
	12.5MG; 80MG	A201192	002	Feb 25, 2014
	25MG; 80MG	A201192	003	Feb 25, 2014

HYDROCHLOROTHIAZIDE; TIMOLOL MALEATE

TABLET; ORAL

TIMOLIDE 10-25

MERCK	25MG; 10MG	N018061	001	
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HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

DYAZIDE

+ GLAXOSMITHKLINE LLC	25MG; 37.5MG	N016042	003	Mar 03, 1994
	25MG; 50MG	N016042	002	

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

ANI PHARMS	25MG; 37.5MG	A074970	001	Jan 06, 1998
	25MG; 50MG	A074259	001	Mar 30, 1995
CHARTWELL RX	25MG; 50MG	A073191	001	Jul 31, 1991
DURAMED PHARMS BARR	25MG; 37.5MG	A075052	001	Jun 18, 1999
NOVARTIS	25MG; 37.5MG	A074857	001	Sep 09, 1997
VITARINE	25MG; 50MG	A071737	001	Feb 12, 1988

TABLET; ORAL

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

AM THERAP	50MG; 75MG	A072022	001	Apr 17, 1988
ANI PHARMS	50MG; 75MG	A073467	001	Jan 31, 1996
PLIVA	25MG; 37.5MG	A074026	001	Apr 26, 1996
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
CADILA	12.5MG; 80MG	A203000	001	Mar 15, 2019
	12.5MG; 160MG	A203000	002	Mar 15, 2019
	12.5MG; 320MG	A203000	003	Mar 15, 2019
	25MG; 160MG	A203000	004	Mar 15, 2019
	25MG; 320MG	A203000	005	Mar 15, 2019

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

VALSARTAN AND HYDROCHLOROTHIAZIDE

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

CAPSULE, EXTENDED RELEASE; ORAL

ZOHYDRO ER

+ RECRO GAINESVILLE	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

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

ANI PHARMS	5MG;200MG	A077454 001	Jun 23, 2010
SUN PHARM INDS INC	2.5MG;200MG	A091633 001	May 28, 2013
	5MG;200MG	A091633 002	May 28, 2013
	7.5MG;200MG	A091633 003	May 28, 2013
	10MG;200MG	A091633 004	May 28, 2013
TEVA	7.5MG;200MG	A076023 001	Apr 11, 2003
REPREXAIN			
AMNEAL PHARMS NY	2.5MG;200MG	A076642 003	Oct 19, 2007
	10MG;200MG	A076642 004	Oct 19, 2007
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
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HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

MAYNE PHARMA INC	5MG/5ML;60MG/5ML	A205658 001	Nov 17, 2015
PADAGIS US	5MG/5ML;60MG/5ML	A204658 001	Apr 29, 2014
TORRENT	5MG/5ML;60MG/5ML	A206661 001	Jan 23, 2019
TRIS PHARMA INC	5MG/5ML;60MG/5ML	A203839 001	Oct 28, 2014
REZIRA			
+ PERSION	5MG/5ML;60MG/5ML **	N022442 001	Jun 08, 2011

HYDROCORTAMATE HYDROCHLORIDE

OINTMENT; TOPICAL

MAGNACORT

PFIZER	0.5%	N010554 001	
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HYDROCORTISONE

AEROSOL; TOPICAL

AEROSEB-HC

ALLERGAN HERBERT	0.5%	A085805 001	
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CREAM; TOPICAL

CORT-DOME

BAYER PHARMS	0.5%	N009585 003	
	1%	N009585 001	

DERMACORT

MONARCH PHARMS	1%	A083011 002	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE

CREAM;TOPICAL

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

H-CORT

PHARM ASSOC	0.5%	A086823	001	
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HC #1

BAYER PHARMS	0.5%	A080438	001	
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HC #4

BAYER PHARMS	1%	A080438	002	
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HC (HYDROCORTISONE)

C AND M PHARMA	0.5%	A080482	003	
	1%	A080482	004	

HI-COR

C AND M PHARMA	2.5%	A080483	001	
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HYDROCORTISONE

ALPHARMA US PHARMS	2.5%	A089754	001	Feb 01, 1989
ALTANA	0.5%	A080848	002	
	1%	A080848	003	

AMBIX	1%	A086080	001	
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	2.5%	A086271	001	
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EVERYLIFE	0.5%	A080452	001	
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	1%	A080452	002	
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G AND W LABS	1%	A084059	001	
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INGRAM PHARM	0.5%	A080456	002	
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	1%	A080456	003	
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IVAX PHARMS	1%	A085733	001	
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NASKA	1%	A089706	001	Mar 10, 1988
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PERRIGO NEW YORK	0.5%	A084970	002	
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	1%	A085026	001	
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PHARMADERM	1%	A088845	001	Feb 27, 1986
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	2.5%	A089413	001	Dec 16, 1986
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PHARMAFAIR	1%	A087838	001	Jul 28, 1982
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STIEFEL	1%	A086170	001	
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SYOSSET	0.5%	A085527	001	
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TARO	0.5%	A086154	001	
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TARO PHARM INDS LTD	1%	A086155	001	
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TELIGENT	2.5%	A203810	001	Jul 23, 2018
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TEVA	0.5%	A080400	002	
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	1%	A080400	003	
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	1%	A085191	001	
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	2.5%	A080400	004	
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TOPIDERM	1%	A089273	001	Feb 17, 1989
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USL PHARMA	1%	A088027	001	Sep 27, 1983
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	2.5%	A088029	001	Sep 27, 1983
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WHITEWORTH TOWN PLSN	1%	A080496	002	
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HYTONE

+ VALEANT INTL	1% **	A080472	003	
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+	2.5% **	A080472	004	
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NOGENIC HC

IVAX PHARMS	1%	A087427	001	Apr 04, 1988
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NUTRACORT

BAUSCH	0.5%	A080442	002	
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	1%	A080442	003	
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PENECORT

ALLERGAN HERBERT	1%	A088216	001	Jun 06, 1984
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PROTOCORT

MONARCH PHARMS	1%	A083011	001	
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SYNACORT

BAUSCH	0.5%	A087459	001	
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+	1%	A087458	001	
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+	2.5%	A087457	001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE

ENEMA;RECTAL			
HYDROCORTISONE			
TEVA PHARMS	100MG/60ML	A074171 001	May 27, 1994
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	
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			
BAUSCH	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
FOUGERA PHARMS	2.5%	A040351 001	Jul 25, 2000
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
TELIGENT	2.5%	A203804 001	Jul 27, 2018
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			
PADAGIS US	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			
ACTAVIS MID ATLANTIC	1%	A087796 001	Oct 13, 1982
ALTANA	0.5%	A080489 002	
	1%	A080489 003	
AMBIX	1%	A086079 001	
	2.5%	A086272 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE

OINTMENT; TOPICAL

HYDROCORTISONE

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
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POWDER; FOR RX COMPOUNDING

H-CORT

TORCH	100%	A087834 001	Mar 29, 1982
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HYDRO-RX

X GEN PHARMS	100%	A085982 001	
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HYDROCORTISONE

PADDOCK LLC	100%	A088082 001	Apr 08, 1983
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SOLUTION; TOPICAL

PENECORT

+ ALLERGAN HERBERT	1%	A088214 001	Jun 06, 1984
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TEXACORT

MISSION PHARMA	1%	A080425 001	
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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	
HIKMA INTL PHARMS	5MG	A083365 002	Feb 23, 2015
	10MG	A083365 003	Feb 23, 2015
	20MG	A083365 001	
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	
STRIDES PHARMA	5MG	A040761 001	Jul 16, 2007
	10MG	A040761 002	Jul 16, 2007
	20MG	A040761 003	Jul 16, 2007
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	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE ACETATE

CREAM;TOPICAL

CARMOL HC

+ FOUGERA PHARMS 1% A080505 001

HEMSOL-HC

ABLE 1% A081274 001 Jun 19, 1992

HYDROCORTISONE ACETATE

CENCI 1% A080419 001 Jan 25, 1982

IMPERIUM 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

HYDROCORTONE

MERCK 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

MERCK 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE

SUSPENSION/DROPS;OPHTHALMIC

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; 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
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HYDROCORTISONE ACETATE; OXYTETRACYCLINE HYDROCHLORIDE

SUSPENSION;OPHTHALMIC

TERRA-CORTRIL

PFIZER	1.5%;EQ 5MG BASE/ML	A061016	001
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HYDROCORTISONE ACETATE; PRAMOXINE HYDROCHLORIDE

AEROSOL, METERED;TOPICAL

HYDROCORTISONE ACETATE 1% AND PRAMOXINE HYDROCHLORIDE 1%

GENUS	1%;1%	A089440	001	May 17, 1988
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LOTION;TOPICAL

PRAMOSONE

FERNDAL LABS	0.5%;1%	A083213	002
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HYDROCORTISONE BUTYRATE

CREAM;TOPICAL

LOCOID

YAMANOUCHI	0.1%	N018795	001	Jan 07, 1983
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OINTMENT;TOPICAL

LOCOID

YAMANOUCHI	0.1%	N019106	001	Jul 03, 1984
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SOLUTION;TOPICAL

LOCOID

YAMANOUCHI	0.1%	N019819	001	Sep 15, 1988
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HYDROCORTISONE CYPIONATE

SUSPENSION;ORAL

CORTEF

PHARMACIA AND UPJOHN	EQ 10MG BASE/5ML	N009900	001
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HYDROCORTISONE SODIUM PHOSPHATE

INJECTABLE;INJECTION

HYDROCORTONE

+	MERCK	EQ 50MG BASE/ML	N012052	001
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HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE;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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION

HYDROCORTISONE SODIUM SUCCINATE

EQ 1GM BASE/VIAL

A084748 001

HYDROCORTISONE VALERATE

CREAM; TOPICAL

HYDROCORTISONE VALERATE

COSETTE 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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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	
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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	
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SALUTENSIN-DEMI

SHIRE	25MG; 0.125MG	N012359	004	
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HYDROGEN PEROXIDE

SOLUTION; TOPICAL

ESKATA

+ ACLARIS	40%	N209305	001	Dec 14, 2017
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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

INJECTABLE; INJECTION

DILAUDID

+ FRESENIUS KABI USA	4MG/ML	N019034	005	Apr 30, 2009
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DILAUDID-HP

+ FRESENIUS KABI USA	10MG/ML	N019034	001	Jan 11, 1984
	250MG/VIAL	N019034	002	Aug 04, 1994

HYDROMORPHONE HYDROCHLORIDE

BARR	10MG/ML	A076444	001	Apr 25, 2003
HOSPIRA	10MG/ML	A074598	001	Jun 19, 1997
WATSON LABS	10MG/ML	A074317	001	Aug 23, 1995

SOLUTION; ORAL

HYDROMORPHONE HYDROCHLORIDE

GENUS	5MG/5ML	A207108	001	Apr 22, 2020
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TABLET; ORAL

HYDROMORPHONE HYDROCHLORIDE

GENUS	2MG	A077471	002	Dec 09, 2009
	4MG	A077471	003	Dec 09, 2009
	8MG	A077471	001	Dec 09, 2009
NESHER PHARMS	2MG	A077311	001	Nov 09, 2005
	4MG	A077311	002	Nov 09, 2005
	8MG	A077311	003	Nov 09, 2005
NOSTRUM LABS INC	8MG	A076723	001	Oct 18, 2005

TABLET, EXTENDED RELEASE; ORAL

EXALGO

+ SPECGX LLC	8MG **	N021217	001	Mar 01, 2010
	12MG **	N021217	002	Mar 01, 2010
	16MG **	N021217	003	Mar 01, 2010
	32MG **	N021217	004	Aug 24, 2012

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROXOCOBALAMIN

INJECTABLE; INJECTION

ALPHAREDISOL

MERCCK

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

PAREDRIINE

PHARMICS

1%

N000004 004

HYDROXYCHLOROQUINE SULFATE

TABLET; ORAL

HYDROXYCHLOROQUINE SULFATE

HIKMA PHARMS

200MG

A040760 001 Aug 15, 2007

INVATECH

200MG

A040150 001 Jan 27, 1996

HYDROXYPROGESTERONE CAPROATE

INJECTABLE; INJECTION

HYDROXYPROGESTERONE CAPROATE

AKORN

125MG/ML

N018004 001

ALLERGAN

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

HYDROXYPROGESTERONE CAPROATE

AM REGENT

1250MG/5ML (250MG/ML)

A210724 001 Aug 09, 2019

EUGIA PHARMA

1250MG/5ML (250MG/ML)

A211142 001 May 09, 2019

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

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

DR REDDYS

50MG/ML

A085779 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE HYDROCHLORIDE

	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	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
ANIMA	10MG/5ML	A086880	001	
KV PHARM	10MG/5ML	A087730	001	Jul 01, 1982
TORRENT	10MG/5ML	A210634	001	Feb 26, 2019
VINTAGE PHARMS	10MG/5ML	A040391	001	Apr 10, 2002
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
AUROBINDO PHARMA LTD	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
	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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROXYZINE HYDROCHLORIDE

TABLET;ORAL

HYDROXYZINE HYDROCHLORIDE

RISING PHARMA	10MG	A091176 001	Jun 07, 2010
	25MG	A091176 002	Jun 07, 2010
	50MG	A091176 003	Jun 07, 2010
SANDOZ	10MG	A087246 002	
	25MG	A085247 001	
	50MG	A087245 001	
STRIDES PHARMA	10MG	A087602 001	Jan 22, 1982
	25MG	A087603 001	Jan 22, 1982
	50MG	A087604 001	Jan 22, 1982
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
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			
BEXIMCO PHARMS USA	EQ 25MG HYDROCHLORIDE	A081127 001	Jun 28, 1991
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 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
	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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROXYZINE PAMOATE

SUSPENSION; ORAL

VISTARIL

PFIZER

EQ 25MG HYDROCHLORIDE/5ML

N011795 001

IBANDRONATE SODIUM

INJECTABLE; INTRAVENOUS

BONIVA

+ ROCHE

EQ 3MG BASE/3ML

N021858 001 Jan 06, 2006

IBANDRONATE SODIUM

EMCURE PHARMS LTD

EQ 3MG BASE/3ML

A203987 001 Sep 02, 2014

NANG KUANG PHARM CO

EQ 3MG BASE/3ML

A204329 001 Jun 16, 2021

TABLET; ORAL

BONIVA

+ HOFFMANN LA ROCHE

EQ 2.5MG BASE **

N021455 001 May 16, 2003

+

EQ 150MG BASE

N021455 002 Mar 24, 2005

IBANDRONATE SODIUM

MYLAN PHARMS INC

EQ 150MG BASE

A078995 001 Mar 19, 2012

SUN PHARM INDUSTRIES

EQ 150MG BASE

A078996 001 Aug 15, 2012

IBRUTINIB

CAPSULE; ORAL

IBRUTINIB

ZYDUS

70MG

A211344 001 Mar 31, 2021

140MG

A211344 002 Mar 31, 2021

IBUPROFEN

CAPSULE; ORAL

IBUPROFEN

CONTRACT PHARMACAL

200MG

A074782 001 Jul 06, 1998

STRIDES PHARMA

EQ 200MG FREE ACID AND POTASSIUM SALT

A204469 001 Mar 28, 2018

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

GLAXOSMITHKLINE

100MG/5ML

N019833 002 Sep 19, 1989

CHILDREN'S ELIXSURE

MOBERG PHARMA NORTH

100MG/5ML

N021604 001 Jan 07, 2004

IBU

ABBOTT

100MG/5ML

N019784 001 Dec 18, 1989

IBUPROFEN

STRIDES PHARMA

100MG/5ML

A211666 001 Feb 22, 2021

MOTRIN

+ MCNEIL CONSUMER

100MG/5ML **

N019842 001 Sep 19, 1989

SUSPENSION/DROPS; ORAL

MOTRIN

MCNEIL

40MG/ML

N020476 001 May 25, 1995

PEDIATRIC ADVIL

+ GLAXOSMITHKLINE

100MG/2.5ML

N020812 001 Jan 30, 1998

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

400MG

A071058 001 Aug 11, 1988

600MG

A071059 001 Aug 11, 1988

800MG

A071965 001 Aug 11, 1988

IBU-TAB 200

ALRA

200MG

A071057 001 Aug 11, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBUPROFEN

TABLET; ORAL

IBUPRIN

PLIVA	200MG	A071773 001	Jul 16, 1987
IBUPROFEN			
ABBOTT	600MG	A070556 001	Jun 14, 1985
	800MG	A071264 001	Jul 25, 1986
ANI PHARMS	200MG	A071144 001	Jan 20, 1987
	200MG	A072901 001	Dec 19, 1991
	200MG	A072903 001	Dec 19, 1991
CONTRACT PHARMACAL	200MG	A071265 001	Oct 15, 1986
	200MG	A071265 002	Sep 10, 1987
	200MG	A071735 001	Sep 10, 1987
	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
HEC PHARM	400MG	A204062 001	Sep 10, 2018
	600MG	A204062 002	Sep 10, 2018
	800MG	A204062 003	Sep 10, 2018
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
MERRO PHARM	200MG	A070985 001	Oct 02, 1987
MYLAN	200MG	A071870 001	May 05, 1988
	400MG	A070045 001	Sep 24, 1985
	600MG	A070057 001	Sep 24, 1985
	800MG	A071999 001	Dec 03, 1987
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	300MG	A070328 001	Aug 06, 1985
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
RISING	300MG	A070736 002	Jun 12, 1986
	400MG	A070736 003	Jun 12, 1986
	600MG	A070736 001	Jun 12, 1986
	800MG	A071938 001	Jan 14, 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
STRIDES PHARMA	200MG	A071575 001	May 08, 1987
	400MG	A070329 001	Aug 06, 1985
	600MG	A070330 001	Aug 06, 1985
	800MG	A070986 001	Jul 25, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBUPROFEN

TABLET;ORAL

IBUPROFEN

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
ULTRATAB LABS INC	200MG		A209076	001	Jan 06, 2020
VINTAGE PHARMS	200MG		A071639	001	Feb 02, 1988
	200MG		A072249	001	Jan 10, 1989
	300MG		A071230	001	Oct 22, 1986
	400MG		A071231	001	Oct 22, 1986
	400MG		A071644	001	Feb 01, 1988
	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
	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
YICHANG HUMANWELL	200MG		A214003	001	Oct 19, 2020
IBUPROHM					
OHM LABS	200MG		A071214	001	Dec 01, 1986
	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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBUPROFEN SODIUM

TABLET; ORAL

IBUPROFEN SODIUM

PERRIGO R AND D

EQ 200MG BASE

A206581 001 Aug 03, 2015

IBUPROFEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

COMBUNOX

+ FOREST LABS

400MG;5MG **

N021378 001 Nov 26, 2004

OXYCODONE HYDROCHLORIDE AND IBUPROFEN

ACTAVIS ELIZABETH

400MG;5MG

A078769 001 Jan 04, 2008

BARR LABS INC

400MG;5MG

A078316 001 Nov 29, 2007

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

LUITPOLD

0.1MG/ML

A090240 001 Jan 11, 2010

MYLAN INSTITUTIONAL

0.1MG/ML

A090924 001 Jan 11, 2010

ICATIBANT ACETATE

INJECTABLE; SUBCUTANEOUS

ICATIBANT ACETATE

DR REDDYS

EQ 30MG BASE/3ML (EQ 10MG BASE/ML)

A213054 001 Oct 05, 2020

ICOSAPENT ETHYL

CAPSULE; ORAL

ICOSAPENT ETHYL

TEVA PHARMS USA

500MG

A209525 001 Sep 11, 2020

1GM

A209525 002 Sep 11, 2020

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDAMYCIN

PFIZER

5MG/VIAL

N050661 002 Sep 27, 1990

10MG/VIAL

N050661 001 Sep 27, 1990

20MG/VIAL

N050661 003 Apr 25, 1995

IDARUBICIN HYDROCHLORIDE

FRESENIUS KABI USA

1MG/ML

A065440 001 Aug 04, 2009

MYLAN LABS LTD

1MG/ML

A200144 001 Oct 11, 2012

SANDOZ

1MG/ML

A091293 001 Mar 29, 2011

TEVA PARENTERAL

5MG/VIAL

A065037 003 May 01, 2002

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

MYLAN LABS LTD

1GM/20ML (50MG/ML)

A201689 001 Nov 26, 2012

3GM/60ML (50MG/ML)

A201689 002 Nov 26, 2012

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

ILOPERIDONE

TABLET; ORAL

ILOPERIDONE

INVENTIA

1MG

A207231 001 Nov 28, 2016

2MG

A207231 002 Nov 28, 2016

4MG

A207231 003 Nov 28, 2016

6MG

A207231 004 Nov 28, 2016

8MG

A207231 005 Nov 28, 2016

10MG

A207231 006 Nov 28, 2016

12MG

A207231 007 Nov 28, 2016

TARO PHARM INDS LTD

1MG

A207098 001 Jul 22, 2019

2MG

A207098 002 Jul 22, 2019

4MG

A207098 003 Jul 22, 2019

6MG

A207098 004 Jul 22, 2019

8MG

A207098 005 Jul 22, 2019

10MG

A207098 006 Jul 22, 2019

12MG

A207098 007 Jul 22, 2019

ILOPROST

SOLUTION; INHALATION

VENTAVIS

ACTELION

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

TABLET; ORAL

IMATINIB MESYLATE

AMNEAL PHARMS

EQ 100MG BASE

A207495 001 Feb 08, 2019

EQ 400MG BASE

A207495 002 Feb 08, 2019

IMIPRAMINE HYDROCHLORIDE

CONCENTRATE; ORAL

IMIPRAMINE HYDROCHLORIDE

NOVARTIS

25MG/ML

A086765 001

INJECTABLE; INJECTION

TOFRANIL

NOVARTIS

12.5MG/ML

N011838 002

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

CHARTWELL

10MG

A090441 002 Mar 11, 2010

25MG

A090441 003 Mar 11, 2010

50MG

A090441 001 Mar 11, 2010

LEDERLE

10MG

A086269 001

25MG

A086267 001

50MG

A086268 001

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

	25MG	A087619 001	Feb 09, 1982
	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

RISING	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

ANDA REPOSITORY	5%	A091044 001	Feb 28, 2011
COSETTE	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

INDACATEROL MALEATE

POWDER; INHALATION

ARCAPTA NEOHALER

+	NOVARTIS	EQ 75MCG BASE	N022383 001	Jul 01, 2011
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INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

ANI PHARMS	1.25MG	A074498 002	Feb 12, 1998
	1.25MG	A075201 001	Dec 04, 1998
	2.5MG	A074498 001	Oct 31, 1996
	2.5MG	A075201 002	Dec 04, 1998
MYLAN PHARMS INC	1.25MG	A075105 001	Jul 23, 1998
	2.5MG	A075105 002	Jul 23, 1998
RISING	1.25MG	A074461 002	Mar 26, 1997
	2.5MG	A074461 001	Mar 27, 1996
TEVA	1.25MG	A074665 001	Apr 04, 1997
	2.5MG	A074665 002	Apr 04, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

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	Apr 29, 1993
+	2.5MG **	N018538 001	Jul 06, 1983

INDECAINIDE HYDROCHLORIDE

TABLET, 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 200MG BASE	N020685 003	Mar 13, 1996
	EQ 333MG BASE	N020685 005	Dec 17, 1998
+	EQ 400MG BASE	N020685 001	Mar 13, 1996

INDIUM IN-111 CHLORIDE

INJECTABLE; INJECTION

INDICLOR

+ GE HEALTHCARE	2mCi/0.2ML	N019862 001	Dec 29, 1992
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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

+ EGALET	25MG **	N016059 001	
+	50MG **	N016059 002	

INDOMETHACIN

ABLE	25MG	A076666 001	Dec 17, 2003
	50MG	A076666 002	Dec 17, 2003
ANI PHARMS	25MG	A071148 001	Mar 18, 1987
	50MG	A071149 001	Mar 18, 1987
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
HERITAGE	25MG	N018851 001	May 18, 1984
	50MG	N018851 002	May 18, 1984
HERITAGE PHARMA	25MG	A070719 001	Feb 12, 1986
	50MG	A070756 001	Feb 12, 1986
IVAX SUB TEVA PHARMS	25MG	N018730 001	May 04, 1984
	50MG	N018730 002	May 04, 1984
JUBILANT GENERICS	25MG	A205215 001	Aug 25, 2017
	50MG	A205215 002	Aug 25, 2017
MUTUAL PHARM	25MG	A070067 001	Oct 03, 1986
	50MG	A070068 001	Oct 03, 1986
PARKE DAVIS	25MG	N018806 001	Nov 23, 1984
	50MG	N018806 002	Nov 23, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INDOMETHACIN

CAPSULE; ORAL

INDOMETHACIN

PIONEER PHARMS	25MG	A070813 001	Aug 11, 1986
	50MG	A070592 001	Aug 11, 1986
RISING	25MG	N018858 001	Apr 20, 1984
	50MG	A070624 001	Sep 04, 1985
	50MG	N018858 002	Apr 20, 1984
SUN PHARM INDS INC	25MG	A091401 001	Mar 28, 2013
	50MG	A091401 002	Mar 28, 2013
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

TIVORBEX

+ GENUS	20MG	N204768 001	Feb 24, 2014
+	40MG	N204768 002	Feb 24, 2014

CAPSULE, EXTENDED RELEASE; ORAL

INDOCIN SR

+ EGALET	75MG **	N018185 001	Feb 23, 1982
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INDOMETHACIN

ABLE	75MG	A076114 001	Feb 06, 2002
AUROBINDO PHARMA LTD	75MG	A204243 001	Dec 27, 2016
INWOOD LABS	75MG	A072410 001	Mar 15, 1989
JUBILANT GENERICS	75MG	A202706 001	Oct 05, 2015
RISING PHARMA	75MG	A202139 001	Mar 20, 2014
WATSON LABS INC	75MG	A202572 001	Dec 09, 2013

SUPPOSITORY; RECTAL

INDOCIN

+ EGALET	50MG **	N017814 001	Aug 13, 1984
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SUSPENSION; ORAL

INDOMETHACIN

HIKMA	25MG/5ML	A071412 001	Mar 18, 1987
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INDOMETHACIN SODIUM

INJECTABLE; INJECTION

INDOCIN

+ RECORDATI RARE	EQ 1MG BASE/VIAL	N018878 001	Jan 30, 1985
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INGENOL MEBUTATE

GEL; TOPICAL

INGENOL MEBUTATE

PADAGIS ISRAEL	0.015%	A209018 001	Jan 07, 2019
	0.05%	A209019 001	Jan 09, 2019

PICATO

+ LEO LABS	0.015%	N202833 001	Jan 23, 2012
+	0.05%	N202833 002	Jan 23, 2012

INULIN

INJECTABLE; INJECTION

INULIN AND SODIUM CHLORIDE

ISO TEX	100MG/ML	N002282 001	
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INVERT SUGAR

INJECTABLE; INJECTION

TRAVERT 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE	10GM/100ML	N016717 001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

IO DOXAMATE 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

FOR SOLUTION; ORAL

ORALTAG

INTERPHARMA PRAHA AS

9.7GM/BOT

N205383 001 Mar 26, 2015

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
SCANLUX-300			
SANOCHEMIA CORP USA	61%	A090394 001	Jun 18, 2010
SCANLUX-370			
SANOCHEMIA CORP USA	76%	A090394 002	Jun 18, 2010

IOPANOIC ACID

TABLET; ORAL

TELEPAQUE

GE HEALTHCARE	500MG	N008032 001
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IOPHENDYLATE

INJECTABLE; INJECTION

PANTOPAQUE

ALCON	100%	N005319 001
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IOPROMIDE

INJECTABLE; INJECTION

ULTRAVIST (PHARMACY BULK)

+ BAYER HLTHCARE	49.9%	N021425 003	Mar 12, 2004
ULTRAVIST 150			
+ BAYER HLTHCARE	31.2%	N020220 004	May 10, 1995
ULTRAVIST 240			
+ BAYER HLTHCARE	49.9%	N020220 003	May 10, 1995
ULTRAVIST 300 IN PLASTIC CONTAINER			
+ BAYER HLTHCARE	62.3%	N020220 005	Nov 18, 2008

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IOTHALAMATE MEGLUMINE

INJECTABLE; INJECTION

CONRAY 30

+ LIEBEL-FLARSHEIM 30% N016983 001

CONRAY 43

+ LIEBEL-FLARSHEIM 43% N013295 002

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% N019710 002 Dec 30, 1988

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

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

LANDELA PHARM 0.02% A077072 001 Jul 19, 2005

MYLAN SPECIALITY LP 0.02% A074755 001 Jan 10, 1997

ROXANE 0.02% A075867 001 Jul 22, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IPRATROPIUM BROMIDE

SOLUTION; INHALATION

IPRATROPIUM BROMIDE

TEVA PHARMS USA	0.02%	A075313	001	Feb 07, 2000
WATSON LABS	0.02%	A076291	001	May 09, 2005
ZENNOVA	0.02%	A075507	001	Jan 19, 2001

SPRAY, METERED; NASAL

ATROVENT

+ BOEHRINGER INGELHEIM	0.021MG/SPRAY **	N020393	001	Oct 20, 1995
+	0.042MG/SPRAY **	N020394	001	Oct 20, 1995

IPRATROPIUM BROMIDE

MYLAN SPECIALITY LP	0.021MG/SPRAY	A075552	001	Mar 31, 2003
	0.042MG/SPRAY	A075553	001	Mar 31, 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
HIKMA	75MG	A090201	001	Oct 15, 2012
	150MG	A090201	002	Oct 15, 2012
	300MG	A090201	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
PICKET PHARMS	75MG	A203161	001	Sep 27, 2012
	150MG	A203161	002	Sep 27, 2012
	300MG	A203161	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

CIPLA LTD	40MG/2ML (20MG/ML)	A077219	001	Feb 20, 2008
	100MG/5ML (20MG/ML)	A077219	002	Feb 20, 2008
EMCURE PHARMS LTD	40MG/2ML (20MG/ML)	A200771	001	Feb 14, 2012
	100MG/5ML (20MG/ML)	A200771	002	Feb 14, 2012
FRESENIUS KABI USA	40MG/2ML (20MG/ML)	A078188	001	Feb 27, 2008
	100MG/5ML (20MG/ML)	A078188	002	Feb 27, 2008
PICKET PHARMS	40MG/2ML (20MG/ML)	A078953	001	Apr 15, 2010
	100MG/5ML (20MG/ML)	A078953	002	Apr 15, 2010
PLIVA LACHEMA	40MG/2ML (20MG/ML)	A078122	001	Oct 31, 2008
	100MG/5ML (20MG/ML)	A078122	002	Oct 31, 2008
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

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
+	0.167%	A088470	001	Mar 14, 1984
	0.167%	A089616	001	Jun 13, 1991

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOETHARINE HYDROCHLORIDE

SOLUTION; INHALATION

ISOETHARINE HYDROCHLORIDE

+		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 MESYLATEAEROSOL, 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	
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ISONIAZID

INJECTABLE; INJECTION

NYDRAZID

	SANDOZ	100MG/ML **	N008662 001	
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RIMIFON

	ROCHE	25MG/ML	N008420 002	
		100MG/ML	N008420 003	

SYRUP; ORAL

ISONIAZID

	ANDA REPOSITORY	50MG/5ML	A081118 001	Jul 21, 1997
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LANIAZID

	LANNETT	50MG/5ML	A089243 001	Feb 03, 1986
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RIMIFON

	ROCHE	50MG/5ML	N008420 001	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISONIAZID

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

+ SANDOZ 100MG ** N008678 002

+ 300MG ** N008678 003

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

300MG A089776 001 Jun 13, 1988

NYDRAZID

BRISTOL MYERS SQUIBB 100MG N008392 003

STANOZIDE

EVERYLIFE 100MG A080126 001

300MG A080126 002

ISONIAZID; 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

RIFAMPIN AND ISONIAZID

HIKMA INTL PHARMS 150MG; 300MG A065221 001 Jul 29, 2005

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOPROPAMIDE IODIDE

TABLET; ORAL

DARBID

GLAXOSMITHKLINE EQ 5MG BASE N010744 001

ISOPROPYL ALCOHOL

SOLUTION; TOPICAL

ZURAGARD

+ ZUREX PHARMA 70% N210872 001 Apr 26, 2019

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

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

CIPLA 0.2MG/ML A211738 001 Jun 28, 2019

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-MEDIHALER

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOSORBIDE DINITRATE

CAPSULE, EXTENDED RELEASE;ORAL

DILATRATE-SR

+ AUXILIUM PHARMS LLC 40MG N019790 001 Sep 02, 1988

ISORDIL

WYETH AYERST 40MG N012882 002 Jul 29, 1988

TABLET;ORAL

ISORDIL

+ BAUSCH 10MG ** N012093 002 Jul 29, 1988

+ 20MG ** N012093 006 Jul 29, 1988

+ 30MG ** N012093 005 Jul 29, 1988

ISOSORBIDE DINITRATE

ANI PHARMS 10MG A086032 001 Jan 07, 1988

HIKMA INTL PHARMS 30MG A040591 001 Jan 10, 2007

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

SORBITRATE

ASTRAZENECA 5MG N016192 001 Apr 01, 1996

10MG N016192 002 Apr 01, 1996

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

ISOSORBIDE MONONITRATE

ANI PHARMS 20MG A075147 001 Nov 27, 1998

HIKMA PHARMS 20MG A075361 001 Oct 05, 2000

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOSORBIDE MONONITRATE

TABLET, EXTENDED RELEASE;ORAL

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
HIKMA INTL PHARMS	30MG	A076813 002	Mar 30, 2006
	60MG	A076813 001	Jan 07, 2005
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
STRIDES PHARMA	30MG	A090598 001	Aug 11, 2010
	60MG	A090598 002	Aug 11, 2010
	120MG	A090598 003	Aug 11, 2010

ISOSULFAN BLUE

INJECTABLE; INJECTION

ISOSULFAN BLUE

BELOTECA INC	1%	A210714 001	Jan 16, 2019
SOMERSET THERAPS LLC	1%	A210558 001	Jul 12, 2019
LYMPHAZURIN			
+ COVIDIEN	1% **	N018310 001	

ISOTRETINOIN

CAPSULE;ORAL

ABSORICA LD

+ SUN PHARM	20MG	N211913 003	Nov 05, 2019
	28MG	N211913 005	Nov 05, 2019

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

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	5MG	A201067 001	Nov 27, 2015
	10MG	A201067 002	Nov 27, 2015

ITRACONAZOLE

CAPSULE;ORAL

ITRACONAZOLE

MYLAN PHARMS INC	100MG	A200463 001	Jul 20, 2012
STRIDES PHARMA	100MG	A206410 001	Jul 02, 2019

INJECTABLE; INJECTION

SPORANOX

JANSSEN PHARMS	10MG/ML	N020966 001	Mar 30, 1999
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SOLUTION;ORAL

ITRACONAZOLE

APOTEX	10MG/ML	A208481 001	Aug 02, 2019
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TABLET;ORAL

ONMEL

+ SEBELA IRELAND LTD	200MG	N022484 001	Apr 29, 2010
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IVERMECTIN

CREAM; TOPICAL

IVERMECTIN

PADAGIS ISRAEL

1%

A210225 001 Apr 13, 2020

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

HIKMA

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

KETOCONAZOLE

CREAM; TOPICAL

NIZORAL

+ JANSSEN PHARMA

2% **

N019084 001 Dec 31, 1985

SHAMPOO; TOPICAL

NIZORAL

+ JANSSEN PHARMS

2%

N019927 001 Aug 31, 1990

SUSPENSION; ORAL

NIZORAL

JANSSEN PHARMA

100MG/5ML

A070767 001 Nov 07, 1986

TABLET; ORAL

KETOCONAZOLE

AAIPHARMA LLC

200MG

A075341 001 Jul 27, 1999

HERITAGE PHARMA

200MG

A074971 001 Jun 15, 1999

200MG

A075362 001 Jun 15, 1999

MYLAN

200MG

A075597 001 Dec 23, 1999

SUN PHARM INDUSTRIES

200MG

A075314 001 Jun 15, 1999

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

KETOCONAZOLE

TABLET; ORAL

KETOCONAZOLE

TEVA 200MG A075273 001 Jun 15, 1999

NIZORAL

+ JANSSEN PHARMS 200MG ** N018533 001

KETOPROFEN

CAPSULE; ORAL

KETOPROFEN

MYLAN 50MG A074035 002 Dec 31, 1996

75MG A074035 003 Dec 31, 1996

RISING 50MG A074024 001 Dec 29, 1995

75MG A074024 002 Dec 29, 1995

TEVA 25MG A073515 001 Dec 22, 1992

75MG A073517 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

AMPHASTAR PHARM 15MG/ML A076209 001 Jul 21, 2004

30MG/ML A076209 002 Jul 21, 2004

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

ATLANTIDE 30MG/ML A077943 001 Mar 27, 2007

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

EUGIA PHARMA 15MG/ML A212939 001 Oct 20, 2020

30MG/ML A212939 002 Oct 20, 2020

GLAND PHARMA LTD 15MG/ML A076722 001 Jul 27, 2004

30MG/ML A076722 002 Jul 27, 2004

HIKMA 15MG/ML ** A075222 001 Apr 26, 1999

15MG/ML A075299 001 Nov 03, 1999

30MG/ML ** A075222 002 Apr 26, 1999

30MG/ML ** A075228 001 Apr 26, 1999

30MG/ML A075299 002 Nov 03, 1999

HOSPIRA 15MG/ML A074801 001 Jun 05, 1997

15MG/ML A074993 001 Jan 27, 1999

30MG/ML A074801 002 Jun 05, 1997

LUITPOLD 15MG/ML A078145 001 Jan 14, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

	30MG/ML	A078145 002	Jan 14, 2008
MYLAN LABS LTD	15MG/ML	A078299 001	Jul 16, 2007
	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 PHARM	15MG/ML	A078737 001	Oct 06, 2008
	30MG/ML	A078737 002	Oct 06, 2008

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
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KETOROLAC TROMETHAMINE

AKORN	0.45%	A203376 001	Feb 10, 2014
EUGIA PHARMA	0.4%	A205191 001	Nov 15, 2018
	0.5%	A205190 001	Dec 03, 2020
SANDOZ INC	0.4%	A078721 001	Nov 05, 2009

TABLET; ORAL

KETOROLAC TROMETHAMINE

CYCLE PHARMS LTD	10MG	A074790 001	Jun 26, 1997
PLIVA	10MG	A075284 001	Jun 23, 1999
WATSON LABS	10MG	A074955 001	Sep 19, 1997

TORADOL

+ ROCHE PALO	10MG **	N019645 001	Dec 20, 1991
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KETOTIFEN FUMARATE

SOLUTION/DROPS; OPHTHALMIC

ZADITOR

+ ALCON PHARMA	EQ 0.025% BASE **	N021066 002	Oct 19, 2006
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KRYPTON, KR-81M

GAS; INHALATION

MPI KRYPTON 81M GENERATOR

GE HEALTHCARE	N/A	N018088 001	
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L-GLUTAMINE

FOR SOLUTION; ORAL

NUTRESTORE

+ EMMAUS MEDCL	5GM/PACKET	N021667 001	Jun 10, 2004
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LABETALOL HYDROCHLORIDE

INJECTABLE; INJECTION

LABETALOL HYDROCHLORIDE

AKORN	5MG/ML	A075524 001	Nov 29, 1999
APOTHECON	5MG/ML	A075355 001	Nov 29, 1999
HOSPIRA	5MG/ML	A075242 001	Sep 30, 1999
MYLAN ASI	5MG/ML	A079134 001	Feb 03, 2010

NORMODYNE

+ SCHERING	5MG/ML **	N018686 001	Aug 01, 1984
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TRANDATE

+ SEBELA IRELAND LTD	5MG/ML **	N019425 001	Dec 31, 1985
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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

+ ALVOGEN	100MG	N018716 001	May 24, 1985
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LABETALOL HYDROCHLORIDE

TABLET; ORAL

TRANDATE

+

400MG **

N018716 004 Aug 01, 1984

LACTITOL

FOR SOLUTION; ORAL

PIZENSY

+

BRAINTREE LABS

10GM

N211281 001 Feb 12, 2020

LACTULOSE

SOLUTION; ORAL

CHRONULAC

+

SANOFI AVENTIS US

10GM/15ML **

N017884 001

CONSTILAC

ALRA

10GM/15ML

A071054 001 Jul 26, 1988

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

ANI PHARMS

10GM/15ML

A078430 001 Nov 28, 2007

HIKMA

10GM/15ML

A073591 001 May 29, 1992

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

CHOLAC

ALRA

10GM/15ML

A071331 001 Jul 26, 1988

ENULOSE

ACTAVIS MID ATLANTIC

10GM/15ML

A071548 001 Aug 15, 1988

GENERLAC

MORTON GROVE

10GM/15ML

A071842 001 Sep 27, 1988

HEPTALAC

TEVA PHARMS

10GM/15ML

A073504 001 May 28, 1993

LACTULOSE

ANI PHARMS

10GM/15ML

A090426 001 Nov 21, 2008

BAJAJ

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

TABLET; ORAL

LAMIVUDINE

MYLAN

100MG

A204002 001 Dec 31, 2014

150MG

A204528 001 Mar 04, 2016

300MG

A204528 002 Mar 04, 2016

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE

+ AUROBINDO PHARMA LTD 300MG;300MG

N022344 001 May 15, 2018

TEMIXYS

+ CELLTRION 300MG;300MG

N211284 001 Nov 16, 2018

LAMIVUDINE; ZIDOVUDINE

TABLET;ORAL

LAMIVUDINE AND ZIDOVUDINE

MYLAN 150MG;300MG

A204005 001 Aug 28, 2014

MYLAN LABS LTD 150MG;300MG

A079079 001 Aug 12, 2019

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

CIPLA 25MG

A077783 001 Nov 01, 2010

100MG

A077783 002 Nov 01, 2010

150MG

A077783 003 Nov 01, 2010

200MG

A077783 004 Nov 01, 2010

GRANULES 25MG

A078982 001 Jan 27, 2009

100MG

A078982 002 Jan 27, 2009

150MG

A078982 003 Jan 27, 2009

200MG

A078982 004 Jan 27, 2009

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

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

RISING PHARMA 25MG

A077420 001 Jan 27, 2009

100MG

A077420 002 Jan 27, 2009

150MG

A077420 003 Jan 27, 2009

200MG

A077420 004 Jan 27, 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

TEVA 25MG

A076388 001 Aug 30, 2006

100MG

A076388 002 Aug 30, 2006

150MG

A076388 003 Aug 30, 2006

200MG

A076388 004 Aug 30, 2006

ZENNOVA 25MG

A078310 001 Feb 04, 2009

100MG

A078310 002 Feb 04, 2009

150MG

A078310 003 Feb 04, 2009

200MG

A078310 004 Feb 04, 2009

TABLET, CHEWABLE;ORAL

LAMICTAL CD

GLAXOSMITHKLINE LLC 100MG

N020764 003 Aug 24, 1998

LAMOTRIGINE

JUBILANT GENERICS 5MG

A200220 001 Feb 28, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LAMOTRIGINE

TABLET, CHEWABLE;ORAL

LAMOTRIGINE

	25MG	A200220 002	Feb 28, 2011
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
TEVA	5MG	A076420 001	Jun 21, 2006
	25MG	A076420 002	Jun 21, 2006

TABLET, EXTENDED RELEASE;ORAL

LAMOTRIGINE

RUBICON	25MG	A202887 001	Jun 17, 2013
	50MG	A202887 002	Jun 17, 2013

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

AJANTA PHARMA LTD	15MG	A203957 001	Oct 14, 2016
	30MG	A203957 002	Oct 14, 2016
BRECKENRIDGE	15MG	A203964 001	Oct 17, 2018
	30MG	A203964 002	Oct 17, 2018
KRKA TOVARNA ZDRAVIL	15MG	A091212 001	Sep 16, 2013
	30MG	A091212 002	Sep 16, 2013

PREVACID

+ TAKEDA PHARMS USA	15MG	N020406 001	May 10, 1995
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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
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TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

ANI PHARMS	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
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PREVACID NAPRAPAC 375 (COPACKAGED)

TAKEDA PHARMS NA	15MG,N/A;N/A,375MG	N021507 003	Nov 14, 2003
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PREVACID NAPRAPAC 500 (COPACKAGED)

TAKEDA PHARMS NA	15MG,N/A;N/A,500MG	N021507 004	Nov 14, 2003
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LANTHANUM CARBONATE

TABLET, CHEWABLE;ORAL

FOSRENOL

TAKEDA PHARMS USA	EQ 250MG BASE	N021468 001	Oct 26, 2004
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LAPYRIUM CHLORIDE; UNDECOYLIUM CHLORIDE IODINE COMPLEX

SOLUTION;TOPICAL

VIRAC REX

CHESEBROUGH PONDS	0.5%;1.8%	N011914 001	
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LASMIDITAN SUCCINATE

TABLET;ORAL

REYVOW

+ ELI LILLY AND CO	EQ 200MG BASE	N211280 003	Dec 18, 2020
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LATANOPROST

SOLUTION/DROPS;OPHTHALMIC

LATANOPROST

APOTEX INC	0.005%	A077697 001	Mar 22, 2011
EUGIA PHARMA	0.005%	A206519 001	Sep 03, 2019

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEFLUNOMIDE

TABLET; ORAL

LEFLUNOMIDE

AET PHARMA	10MG	A213497 001	May 10, 2021
	20MG	A213497 002	May 10, 2021
BARR	10MG	A077083 001	Sep 13, 2005
	20MG	A077083 002	Sep 13, 2005
SANDOZ	10MG	A077085 001	Sep 13, 2005
	20MG	A077085 002	Sep 13, 2005
TEVA PHARMS	10MG	A077084 001	Sep 13, 2005
	20MG	A077084 002	Sep 13, 2005

LESINURAD

TABLET; ORAL

ZURAMPIC

+ IRONWOOD PHARMS INC	200MG	N207988 001	Dec 22, 2015
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LETROZOLE

TABLET; ORAL

LETROZOLE

ACTAVIS TOTOWA	2.5MG	A090292 001	Jul 13, 2011
APOTEX INC	2.5MG	A091303 001	Apr 19, 2012
FRESENIUS KABI USA	2.5MG	A090491 001	Jun 03, 2011
HIKMA	2.5MG	A090838 001	Jun 03, 2011
HIKMA PHARMS	2.5MG	A203796 001	Jun 03, 2016
IMPAX LABS	2.5MG	A091638 001	Jun 03, 2011
JIANGSU PHARMS	2.5MG	A202716 001	May 16, 2013
LANNETT CO INC	2.5MG	A091098 001	Jun 03, 2011
	2.5MG	A202048 001	Oct 29, 2014
MYLAN	2.5MG	A078190 001	Dec 24, 2008
STRIDES PHARMA	2.5MG	A090789 001	Jun 03, 2011
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
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INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

ABIC	EQ 3MG BASE/ML	A089352 001	Jun 01, 1988
	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
	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
NOVAST LABS	EQ 10MG BASE/ML	A210917 001	Nov 23, 2018
PHARMACHEMIE	EQ 350MG BASE/VIAL	A040262 001	Dec 15, 1999
PHARMACHEMIE USA	EQ 50MG BASE/VIAL	A089628 001	Apr 17, 1997
	EQ 100MG BASE/VIAL	A089915 001	Apr 17, 1997
TEVA PARENTERAL	EQ 50MG BASE/VIAL	A081278 001	Sep 28, 1993

LEUCOVORIN CALCIUM PRESERVATIVE FREE

AM REGENT	EQ 50MG BASE/VIAL	A040338 001	Jan 31, 2001
HOSPIRA	EQ 10MG BASE/ML **	A040147 001	Jun 25, 1997
TEVA PARENTERAL	EQ 10MG BASE/ML	A040332 001	Jun 28, 1999

WELLCOVORIN

GLAXOSMITHKLINE	EQ 5MG BASE/ML	A087439 001	Oct 19, 1982
	EQ 25MG BASE/VIAL	A089833 001	Jan 23, 1989
	EQ 50MG BASE/VIAL	A089465 001	Jan 23, 1989
	EQ 100MG BASE/VIAL	A089834 001	Jan 23, 1989

TABLET; ORAL

LEUCOVORIN CALCIUM

ANI PHARMS	EQ 15MG BASE	A075327 001	Mar 24, 1999
PAR PHARM	EQ 5MG BASE	A071600 001	Oct 14, 1987
	EQ 25MG BASE	A071598 001	Oct 14, 1987
PHARMACHEMIE	EQ 5MG BASE	A073099 001	Mar 28, 1997
	EQ 25MG BASE	A073101 001	Mar 28, 1997
XANODYNE PHARM	EQ 5MG BASE	N018459 001	Jan 30, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEUCOVORIN CALCIUM

TABLET; ORAL

LEUCOVORIN CALCIUM

EQ 10MG BASE

A071962 001 Nov 19, 1987

EQ 15MG BASE

A071104 001 Mar 04, 1987

WELLCOVORIN

+ GLAXOSMITHKLINE

EQ 5MG BASE **

N018342 001 Jul 08, 1983

+

EQ 25MG BASE **

N018342 002 Jul 08, 1983

LEUPROLIDE ACETATE

FOR SUSPENSION; INTRAMUSCULAR

LUTRATE DEPOT KIT

+ GP-PHARM SA

22.5MG/VIAL

N205054 001 Aug 28, 2018

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

POWDER; INTRAMUSCULAR

LUPRON DEPOT-PED KIT

+ ABBVIE ENDOCRINE INC 3.75MG, 7.5MG **

N020263 003 Apr 16, 1993

+ 7.5MG, 7.5MG **

N020263 004 Apr 16, 1993

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

MYLAN SPECIALITY LP EQ 0.0103% BASE

A077800 001 Mar 15, 2013

EQ 0.021% BASE

A077800 002 Mar 15, 2013

EQ 0.042% BASE

A077800 003 Mar 15, 2013

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

LEVAMLODIPINE MALEATE

TABLET; ORAL

CONJUPRI

+ CSPC OUYI

EQ 1.25MG BASE

N212895 001 Dec 19, 2019

LEVETIRACETAM

INJECTABLE; INTRAVENOUS

LEVETIRACETAM

AKORN 500MG/5ML (100MG/ML)

A209934 001 May 04, 2018

AM REGENT 500MG/5ML (100MG/ML)

A202143 001 Jan 31, 2012

FRESENIUS KABI USA 500MG/5ML (100MG/ML)

A090813 001 May 26, 2010

JUBILANT GENERICS 500MG/5ML (100MG/ML)

A206838 001 Jun 02, 2016

SOLUTION; ORAL

LEVETIRACETAM

APOTEX INC 100MG/ML

A090187 001 Aug 05, 2011

TOLMAR 100MG/ML

A079107 001 Jan 15, 2009

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVETIRACETAM

TABLET;ORAL

LEVETIRACETAM

	500MG	A077324 002	Jan 15, 2009
	750MG	A077324 003	Jan 15, 2009
	1GM	A077324 004	Jan 15, 2009
LOTUS PHARM CO LTD	250MG	A090906 002	Oct 31, 2016
	750MG	A090906 003	Oct 31, 2016
	1GM	A090906 004	Oct 31, 2016
MYLAN	250MG	A078731 001	Feb 10, 2009
	500MG	A078731 002	Feb 10, 2009
	750MG	A078731 003	Feb 10, 2009
	1GM	A078731 004	Feb 10, 2009
NOSTRUM LABS INC	250MG	A090511 001	Aug 18, 2011
	500MG	A090511 002	Aug 18, 2011
	750MG	A090511 003	Aug 18, 2011
	1GM	A090511 004	Aug 18, 2011
TEVA PHARMS	250MG	A078101 001	Jan 15, 2009
	500MG	A078101 002	Jan 15, 2009
	750MG	A078101 003	Jan 15, 2009
	1GM	A078101 004	Jan 15, 2009
WATSON LABS INC	250MG	A078797 002	Jan 15, 2009
	500MG	A078797 003	Jan 15, 2009
	750MG	A078797 004	Jan 15, 2009
	1GM	A078797 001	Jan 15, 2009

TABLET, EXTENDED RELEASE;ORAL

LEVETIRACETAM

ACTAVIS ELIZABETH	500MG	A091557 001	Sep 12, 2011
	750MG	A091557 002	Sep 12, 2011
LOTUS PHARM CO LTD	500MG	A202095 002	Jun 06, 2016
	750MG	A202095 001	Jun 06, 2016
MYLAN PHARMS INC	500MG	A200475 001	Dec 19, 2011
	750MG	A200475 002	Dec 19, 2011
	1GM	A200475 003	Dec 07, 2015
ROUSES POINT PHARMS	500MG	A202524 001	Aug 27, 2012
	750MG	A202524 002	Aug 27, 2012
SANDOZ	500MG	A091668 001	Nov 01, 2012
	750MG	A091668 002	Nov 01, 2012
SUN PHARM INDUSTRIES	500MG	A091285 001	Sep 12, 2011
	750MG	A091285 002	Sep 12, 2011
TEVA PHARMS	500MG	A091430 001	Sep 12, 2011
	750MG	A091430 002	Sep 12, 2011
TORRENT PHARMS LTD	500MG	A091338 001	May 29, 2012
	750MG	A091338 002	May 29, 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
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LEVOBUNOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

AKBETA

AKORN	0.25%	A074779 001	Oct 29, 1996
	0.5%	A074780 001	Oct 29, 1996

BETAGAN

+ ALLERGAN	0.25%	N019814 001	Jun 28, 1989
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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
SANDOZ INC	0.5%	A074850 001	Oct 28, 1996

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVOPUIVACAINE 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
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LEVOCARNITINE

INJECTABLE; INJECTION

LEVOCARNITINE

TEVA PHARMS USA	200MG/ML	A075881 001	Mar 29, 2001
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SOLUTION; ORAL

CARNITOR

LEADIANT BIOSCI INC	1GM/10ML	N018948 002	Apr 27, 1988
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LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION; ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

APOTEX	2.5MG/5ML	A202915 001	Aug 21, 2014
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XYZAL

+ SANOFI	2.5MG/5ML	N022157 001	Jan 28, 2008
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TABLET; ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

APOTEX	5MG	A203027 001	Feb 13, 2015
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GRANULES	5MG	A090486 001	Mar 26, 2013
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SUN PHARM INDS LTD	5MG	A201653 001	Jun 26, 2015
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US ANTIBIOTICS	5MG	A204323 001	Dec 20, 2016
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XYZAL

+ SANOFI	5MG	N022064 001	May 25, 2007
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LEVODOPA

CAPSULE; ORAL

BENDOPA

VALEANT PHARM INTL	100MG	N016948 003	
	250MG	N016948 001	
	500MG	N016948 002	

DOPAR

SHIRE	100MG	N016913 003	
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	250MG	N016913 001	
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	500MG	N016913 002	
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LARODOPA

ROCHE	100MG	N016912 002	
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	250MG	N016912 001	
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	500MG	N016912 006	
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TABLET; ORAL

DOPAR

SHIRE	250MG	N016913 004	
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	500MG	N016913 005	
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LARODOPA

ROCHE	100MG	N016912 005	
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	250MG	N016912 003	
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	500MG	N016912 004	
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LEVOFLOXACIN

INJECTABLE; INJECTION

LEVAQUIN

+ JANSSEN PHARMS	EQ 500MG/20ML (EQ 25MG/ML) **	N020635 001	Dec 20, 1996
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+ JANSSEN PHARMS	EQ 750MG/30ML (EQ 25MG/ML) **	N020635 004	Dec 20, 1996
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LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER

+ JANSSEN PHARMS	EQ 250MG/50ML (EQ 5MG/ML) **	N020635 002	Dec 20, 1996
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+ JANSSEN PHARMS	EQ 500MG/100ML (EQ 5MG/ML) **	N020635 003	Dec 20, 1996
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+ JANSSEN PHARMS	EQ 750MG/150ML (EQ 5MG/ML) **	N020635 005	Dec 20, 1996
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LEVOFLOXACIN

EMCURE PHARMS LTD	EQ 500MG/20ML (EQ 25MG/ML)	A202590 001	Jan 24, 2013
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	EQ 750MG/30ML (EQ 25MG/ML)	A202590 002	Jan 24, 2013
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HOSPIRA INC	EQ 500MG/20ML (EQ 25MG/ML)	A078577 001	Aug 12, 2015
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	EQ 750MG/30ML (EQ 25MG/ML)	A078577 002	Aug 12, 2015
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness 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	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
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SOLUTION/DROPS; OPHTHALMIC

IQUIX

+ SANTEN	1.5% **	N021571 001	Mar 01, 2004
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LEVOFLOXACIN

MYLAN LABS LTD	0.5%	A204899 001	Dec 08, 2017
RUBICON	0.5%	A078282 001	Dec 20, 2010
WATSON LABS TEVA	0.5%	A076826 001	Feb 10, 2011

QUIXIN

+ SANTEN	0.5% **	N021199 001	Aug 18, 2000
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TABLET; ORAL

LEVAQUIN

+ JANSSEN PHARMS	250MG **	N020634 001	Dec 20, 1996
	500MG **	N020634 002	Dec 20, 1996
	750MG **	N020634 003	Sep 08, 2000

LEVOFLOXACIN

JUBILANT GENERICS	250MG	A203613 001	Jun 19, 2015
	500MG	A203613 002	Jun 19, 2015
MYLAN	250MG	A076276 001	Jun 20, 2011
	500MG	A076276 002	Jun 20, 2011
	750MG	A077097 001	Jun 20, 2011
TORRENT PHARMS	250MG	A090722 001	Jun 20, 2011
	500MG	A090722 002	Jun 20, 2011
	750MG	A090722 003	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

POWDER; INTRAVENOUS

LEVOLEUCOVORIN CALCIUM

ACTAVIS LLC	EQ 50MG BASE/VIAL	A206516 001	Feb 13, 2017
+ AMNEAL	EQ 175MG BASE/VIAL	N208723 001	Sep 29, 2016
	EQ 50MG BASE/VIAL	A207547 001	Feb 13, 2017

SOLUTION; INTRAVENOUS

FUSILEV

+ ACROTECH	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

LEVOLEUCOVORIN CALCIUM

MYLAN TEORANTA	EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)	A203576 001	Oct 20, 2015
	EQ 250MG BASE/25ML (EQ 10MG BASE/ML)	A203576 002	Oct 20, 2015
NOVAST LABS	EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)	A210623 001	May 03, 2018
	EQ 250MG BASE/25ML (EQ 10MG BASE/ML)	A210623 002	May 03, 2018
SANDOZ INC	EQ 250MG BASE/25ML (EQ 10MG BASE/ML)	A203563 002	Mar 09, 2015

LEVOMEPRMAZINE

INJECTABLE; INJECTION

LEVOPROME

IMMUNEX	20MG/ML	N015865 001	
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LEVOMETHADYL ACETATE HYDROCHLORIDE

CONCENTRATE; ORAL

ORLAAM

+ ROXANE	10MG/ML **	N020315 001	Jul 09, 1993
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVOMILNACIPRAN HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

LEVOMILNACIPRAN HYDROCHLORIDE

AMNEAL PHARMS CO	EQ 20MG BASE	A210790 001	Feb 04, 2019
	EQ 40MG BASE	A210790 002	Feb 04, 2019
	EQ 80MG BASE	A210790 003	Feb 04, 2019
	EQ 120MG BASE	A210790 004	Feb 04, 2019
HIKMA	EQ 20MG BASE	A210732 001	Nov 05, 2020
	EQ 40MG BASE	A210732 002	Nov 05, 2020
	EQ 80MG BASE	A210732 003	Nov 05, 2020
	EQ 120MG BASE	A210732 004	Nov 05, 2020

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	
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LEVONORGESTREL

IMPLANT; IMPLANTATION

JADELLE

+ POPULATION COUNCIL	75MG/IMPLANT **	N020544 001	Nov 01, 1996
LEVONORGESTREL			
WYETH PHARMS INC	75MG/IMPLANT	N020627 001	Aug 15, 1996
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

ALVOGEN	1.5MG	A202246 001	Jun 05, 2015
FDN CONSUMER	0.75MG **	A078665 001	Aug 28, 2009
	1.5MG	A200670 001	Jul 12, 2012
LOTUS PHARM CO LTD	0.75MG	A202684 001	Sep 02, 2016
LUPIN LTD	0.75MG	A091328 001	Jan 23, 2013
MYLAN LABS LTD	0.75MG	A202740 001	Sep 02, 2016
NAARI PTE LTD	1.5MG	A207660 001	May 02, 2019
WATSON LABS	0.75MG	A078666 001	Jun 24, 2009
PLAN B			
+ FDN CONSUMER	0.75MG **	N021045 001	Jul 28, 1999
+ FDN CONSUMER	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	
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LEVORPHANOL TARTRATE

INJECTABLE; INJECTION

LEVO-DROMORAN

VALEANT PHARM INTL	2MG/ML	N008719 001	Dec 19, 1991
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TABLET; ORAL

LEVO-DROMORAN

+ VALEANT PHARM INTL	2MG **	N008720 001	Dec 19, 1991
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVORPHANOL TARTRATE

TABLET;ORAL

LEVORPHANOL TARTRATE

SENTYNL THERAPS INC 1MG

A074278 002 Jun 18, 2018

LEVOTHYROXINE SODIUM

CAPSULE;ORAL

LEVOTHYROXINE SODIUM

TEVA PHARMS USA INC 0.075MG

A211369 001 Oct 28, 2020

0.088MG

A213256 001 Jan 06, 2021

0.1MG

A213256 002 Jan 06, 2021

0.125MG

A213256 003 Jan 06, 2021

0.15MG

A211369 002 Oct 28, 2020

POWDER;INTRAVENOUS

LEVOTHYROXINE SODIUM

DR REDDYS 100MCG/VIAL

A208837 001 Mar 27, 2020

PAR STERILE PRODUCTS 200MCG/VIAL

A205366 001 Dec 07, 2015

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

AMNEAL

0.025MG

A210831 001 Feb 19, 2019

0.05MG

A210831 002 Feb 19, 2019

0.075MG

A210831 003 Feb 19, 2019

0.088MG

A210831 004 Feb 19, 2019

0.1MG

A210831 005 Feb 19, 2019

0.112MG

A210831 006 Feb 19, 2019

0.125MG

A210831 007 Feb 19, 2019

0.137MG

A210831 008 Feb 19, 2019

0.15MG

A210831 009 Feb 19, 2019

0.175MG

A210831 010 Feb 19, 2019

0.2MG

A210831 011 Feb 19, 2019

0.3MG

A210831 012 Feb 19, 2019

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

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LIDOCAINEOINTMENT; TOPICAL
ALPHACAINE5% A084946 001
5% A084947 001

LIDOCAINE

BELMORA LLC 5% A080210 001
GENEYORK PHARMS 5% A212486 001 Oct 17, 2019
RISING 5% A208604 001 Sep 20, 2017
TEVA PHARMS USA 5% A210256 001 Jan 16, 2018

XYLOCAINE

+ ASTRAZENECA 5% ** N008048 001

PATCH; TOPICAL

DENTIPATCH

NOVEN 46.1MG/PATCH N020575 002 May 21, 1996

LIDOCAINE

NOVEN PHARMS INC 5% A203265 001 Dec 01, 2020

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

AM REGENT 1% A080850 001

1% A091564 001 Aug 14, 2015

BEL MAR 1% A080710 001

2% A080760 001

BELMORA LLC 2% A080504 001

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

20% A083158 003

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

RISING 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE				
RISING PHARMA	2%	A202242	001	Apr 11, 2014
WATSON LABS	1%	A080377	001	
	1%	A083627	001	
	2%	A080377	002	
	2%	A083627	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
LIDOCATON				
PHARMATON	2%	A084727	001	Aug 17, 1983
LIDOPEN				
MERIDIAN MEDCL TECHN	10%	N017549	001	
XYLOCAINE				
ASTRAZENECA	1%	N010418	005	
	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	1% **	N016801	005	Jan 19, 1988
+	2% **	N016801	001	
+	4% **	N016801	002	
+	10% **	N016801	003	
+	20% **	N016801	004	
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				
BIONPHARMA INC	2%	A080429	001	
LIDOCAINE HYDROCHLORIDE				
COSETTE	2%	A081318	001	Apr 29, 1993
WATSON LABS INC	2%	A040837	001	Mar 23, 2011
XYLOCAINE				
+ AKORN	2%	N008816	001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LIDOCAINE HYDROCHLORIDE

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

PACO 4%

A089688 001 Jun 30, 1989

LTA II KIT

HOSPIRA 4%

A080409 001

HOSPIRA 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

CREAM;TOPICAL

LIDOCAINE AND PRILOCAINE

RHODES PHARMS 2.5%;2.5%

A213253 001 Sep 21, 2020

DISC;TOPICAL

EMLA

ASTRAZENECA 2.5%;2.5%

N020962 001 Feb 04, 1998

LINACLOTIDE

CAPSULE;ORAL

LINACLOTIDE

MYLAN 145MCG

A209564 001 Feb 09, 2021

290MCG

A209564 002 Feb 09, 2021

LINCOMYCIN HYDROCHLORIDE

CAPSULE;ORAL

LINCOCIN

PHARMACIA AND UPJOHN EQ 250MG BASE

N050316 001

EQ 500MG BASE

N050316 002

INJECTABLE; INJECTION

LINCOMYCIN HYDROCHLORIDE

PRAXGEN EQ 300MG BASE/ML

A212770 001 Mar 12, 2021

WATSON LABS EQ 300MG BASE/ML

A063180 001 Apr 16, 1991

LINDANE

CREAM;TOPICAL

KWELL

REED AND CARNRICK 1%

A084218 001

1%

N006309 001

LOTION;TOPICAL

GAMENE

SOLA BARNES HIND 1%

A084989 001

KWELL

REED AND CARNRICK 1%

A084218 002

1%

N006309 003

LINDANE

OLTA PHARMS 1%

A087313 001

WOCKHARDT BIO AG 1%

A088190 001 Aug 16, 1984

SCABENE

STIEFEL 1%

A086769 001

SHAMPOO;TOPICAL

GAMENE

SOLA BARNES HIND 1%

A084988 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LINDANESHAMPOO; TOPICAL
KWELLREED AND CARNRICK 1%
1%A084219 001
N010718 001

LINDANE

OLTA PHARMS 1%

A087266 001

SCABENE

STIEFEL 1%

A087940 001 Apr 08, 1983

LINEZOLID

SOLUTION; INTRAVENOUS

ZYVOX

+ PFIZER 400MG/200ML (2MG/ML) **

N021131 002 Apr 18, 2000

TABLET; ORAL

LINEZOLID

GATE PHARMS 600MG

A091210 001 Feb 05, 2016

RISING 600MG

A078845 001 Dec 21, 2015

TEVA PHARMS USA 600MG

A078061 001 May 18, 2015

ZYVOX

+ PFIZER 400MG **

N021130 001 Apr 18, 2000

LIOTHYRONINE SODIUM

TABLET; ORAL

LIOTHYRONINE SODIUM

MYLAN

EQ 0.005MG BASE

A090326 001 Jul 14, 2009

EQ 0.025MG BASE

A090326 002 Jul 14, 2009

EQ 0.05MG BASE

A090326 003 Jul 14, 2009

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 0.0125MG;0.0031MG

N016807 001

THYROLAR-0.5

+ ALLERGAN 0.025MG;0.0063MG

N016807 005

THYROLAR-1

+ ALLERGAN 0.05MG;0.0125MG

N016807 004

THYROLAR-2

+ ALLERGAN 0.1MG;0.025MG

N016807 002

THYROLAR-3

+ ALLERGAN 0.15MG;0.0375MG

N016807 003

THYROLAR-5

ALLERGAN 0.25MG;0.0625MG

N016807 006

LISINAPRIL

TABLET; ORAL

LISINAPRIL

HERITAGE PHARMA 2.5MG

A075752 001 Jul 01, 2002

5MG

A075752 002 Jul 01, 2002

10MG

A075752 003 Jul 01, 2002

20MG

A075752 004 Jul 01, 2002

30MG

A075752 005 Jul 01, 2002

40MG

A075752 006 Jul 01, 2002

HIKMA INTL PHARMS 2.5MG

A076063 001 Jul 01, 2002

5MG

A076063 002 Jul 01, 2002

10MG

A076063 003 Jul 01, 2002

20MG

A076063 004 Jul 01, 2002

30MG

A076063 006 Jun 27, 2003

40MG

A076063 005 Jul 01, 2002

MYLAN 2.5MG

A076071 001 Jul 01, 2002

5MG

A076071 002 Jul 01, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LISINAPRILTABLET;ORAL
LISINAPRIL

	10MG	A076071 003	Jul 01, 2002
	20MG	A076071 004	Jul 01, 2002
	30MG	A076071 005	Jul 01, 2002
	40MG	A076071 006	Jul 01, 2002
PHARMGEN	2.5MG	A075999 001	Jul 01, 2002
	5MG	A075999 002	Jul 01, 2002
	10MG	A075999 003	Jul 01, 2002
	20MG	A075999 004	Jul 01, 2002
	30MG	A075999 005	Jul 01, 2002
	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
	10MG	N019558 002	Dec 29, 1987
	20MG	N019558 003	Dec 29, 1987

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

MYLAN

150MG

A076243 002 Feb 24, 2003

300MG

A076243 001 Jun 27, 2002

600MG

A078763 001 Apr 15, 2008

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

ALEMBIC PHARMS LTD

300MG

A204445 001 Jun 10, 2015

HERITAGE PHARMA

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LOMEFLOXACIN HYDROCHLORIDE

TABLET; ORAL

MAXAQUIN

PHARMACIA

EQ 400MG BASE

N020013 001 Feb 21, 1992

LOMITAPIDE MESYLATE

CAPSULE; ORAL

JUXTAPID

+ AMRYT

EQ 40MG BASE

N203858 005 Apr 23, 2015

+

EQ 60MG BASE

N203858 006 Apr 23, 2015

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

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

CAPSULE; ORAL

LORATADINE

STRIDES PHARMA

10MG

A211926 001 Jan 15, 2020

SYRUP; ORAL

CLARITIN HIVES RELIEF

+ BAYER HEALTHCARE LLC

1MG/ML **

N020641 003 Nov 19, 2003

LORATADINE

PHARM ASSOC

1MG/ML

A075565 001 Oct 05, 2004

RANBAXY LABS LTD

1MG/ML

A076529 001 Aug 20, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LORATADINE

SYRUP; ORAL

LORATADINE

TEVA

1MG/ML

A075505 001 Nov 07, 2003

TABLET; ORAL

LORATADINE

MYLAN

10MG

A075790 001 Nov 07, 2008

10MG

A078447 001 Aug 12, 2011

PERRIGO

10MG

N021512 001 Jun 24, 2004

TABLET, ORALLY DISINTEGRATING; ORAL

LORATADINE

ACTAVIS LABS FL INC

10MG

A075990 001 Nov 03, 2003

GLAXOSMITHKLINE

10MG

A075822 001 Feb 10, 2003

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

LORATADINE AND PSEUDOEPHEDRINE SULFATE

HERITAGE PHARMA

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

DR REDDYS

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

HIKMA

2MG/ML

A074496 001 Sep 28, 1998

4MG/ML

A074496 002 Sep 28, 1998

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

RISING

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

4MG/ML

A074276 002 Apr 15, 1994

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

ANI PHARMS

0.5MG

A077396 001 Dec 13, 2006

1MG

A077396 002 Dec 13, 2006

2MG

A077396 003 Dec 13, 2006

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LORAZEPAM

TABLET;ORAL

LORAZEPAM

	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
RISING PHARMA	0.5MG	A077657	001	Mar 16, 2006
	1MG	A077657	002	Mar 16, 2006
	2MG	A077657	003	Mar 16, 2006
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

LOSARTAN POTASSIUM

APOTEX CORP	25MG	A090790	001	Oct 06, 2010
	50MG	A090790	002	Oct 06, 2010
	100MG	A090790	003	Oct 06, 2010
HISUN PHARM HANGZHOU	25MG	A204795	001	Apr 04, 2019
	50MG	A204795	002	Apr 04, 2019
	100MG	A204795	003	Apr 04, 2019
MYLAN	25MG	A091590	001	Oct 06, 2010
	50MG	A091590	002	Oct 06, 2010
	100MG	A091590	003	Oct 06, 2010
TEVA	25MG	A076958	001	Apr 06, 2010
	50MG	A076958	002	Apr 06, 2010
	100MG	A076958	003	Apr 06, 2010
TORRENT PHARMS	25MG	A090467	001	Oct 06, 2010
	50MG	A090467	002	Oct 06, 2010
	100MG	A090467	003	Oct 06, 2010
UPSHER SMITH LABS	25MG	A090544	001	Oct 06, 2010
	50MG	A090544	002	Oct 06, 2010
	100MG	A090544	003	Oct 06, 2010

LOTEPREDNOL ETABONATE

SUSPENSION/DROPS;OPHTHALMIC

LOTEMAX

PHARMOS	0.5%	N020841	001	Mar 09, 1998
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LOVASTATIN

TABLET;ORAL

LOVASTATIN

MYLAN	10MG	A075451	001	Dec 17, 2001
	10MG	A075935	001	Dec 17, 2001
	20MG	A075451	002	Dec 17, 2001
	20MG	A075935	002	Dec 17, 2001
	40MG	A075451	003	Dec 17, 2001
	40MG	A075935	003	Dec 17, 2001
SUN PHARM INDUSTRIES	10MG	A077520	001	Apr 14, 2006
	20MG	A077520	002	Apr 14, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LOVASTATINTABLET; ORAL
LOVASTATIN

	40MG	A077520 003	Apr 14, 2006
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	10MG	N021316 001	Jun 26, 2002
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LOXAPINE HYDROCHLORIDE

CONCENTRATE; ORAL

LOXITANE C

TEVA BRANDED PHARM	EQ 25MG BASE/ML	N017658 001	
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INJECTABLE; INJECTION

LOXITANE IM

ACTAVIS LABS UT INC	EQ 50MG BASE/ML	N018039 001	
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LOXAPINE SUCCINATE

CAPSULE; ORAL

LOXAPINE SUCCINATE

RISING PHARMA	EQ 5MG BASE	A076762 001	Nov 01, 2004
	EQ 10MG BASE	A076762 002	Nov 01, 2004
	EQ 25MG BASE	A076762 003	Nov 01, 2004
	EQ 50MG BASE	A076762 004	Nov 01, 2004

LOXITANE

+ TEVA BRANDED PHARM	EQ 5MG BASE **	N017525 001	
+	EQ 10MG BASE **	N017525 002	
+	EQ 25MG BASE **	N017525 003	
+	EQ 50MG BASE **	N017525 004	

TABLET; ORAL

LOXITANE

+ TEVA BRANDED PHARM	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
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LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LURASIDONE HYDROCHLORIDE

AMNEAL PHARMS CO	20MG	A208002 001	Jan 03, 2019
	40MG	A208002 002	Jan 03, 2019
	60MG	A208002 003	Jan 03, 2019
	80MG	A208002 004	Jan 03, 2019
	120MG	A208002 005	Jan 03, 2019
EMCURE PHARMS LTD	20MG	A208058 001	Sep 04, 2019
	40MG	A208058 002	Sep 04, 2019
	60MG	A208058 003	Sep 04, 2019
	80MG	A208058 004	Sep 04, 2019
INVAGEN PHARMS	20MG	A208028 001	Jan 03, 2019
	40MG	A208028 002	Jan 03, 2019
	60MG	A208028 003	Jan 03, 2019
	80MG	A208028 004	Jan 03, 2019
	120MG	A208028 005	Jan 03, 2019
LUPIN LTD	20MG	A208031 001	Jan 03, 2019
	40MG	A208031 002	Jan 03, 2019
	60MG	A208031 003	Jan 03, 2019
	80MG	A208031 004	Jan 03, 2019
	120MG	A208031 005	Jan 03, 2019
PIRAMAL HLTHCARE UK	20MG	A212091 001	Dec 28, 2020
	40MG	A212091 002	Dec 28, 2020
	60MG	A212091 003	Dec 28, 2020
	80MG	A212091 004	Dec 28, 2020
	120MG	A212091 005	Dec 28, 2020
TEVA PHARMS USA	20MG	A208060 001	May 17, 2019

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LURASIDONE HYDROCHLORIDE

	40MG	A208060 002	May 17, 2019
	60MG	A208060 003	May 17, 2019
	80MG	A208060 004	May 17, 2019
	120MG	A208060 005	May 17, 2019
TORRENT	20MG	A208055 001	Jan 03, 2019
	40MG	A208055 002	Jan 03, 2019
	80MG	A208055 003	Jan 03, 2019
	120MG	A208055 004	Jan 03, 2019
WATSON LABS TEVA	20MG	A208016 001	Feb 02, 2021
	40MG	A208016 002	Feb 02, 2021
	60MG	A208016 003	Feb 02, 2021
	80MG	A208016 004	Feb 02, 2021
	120MG	A208016 005	Feb 02, 2021
ZYDUS PHARMS	20MG	A208052 001	Mar 19, 2019
	40MG	A208052 002	Mar 19, 2019
	60MG	A208052 003	Mar 19, 2019
	80MG	A208052 004	Mar 19, 2019
	120MG	A208052 005	Mar 19, 2019

LYPRESSIN

SOLUTION; NASAL

DIAPID

NOVARTIS	0.185MG/ML	N016755 001	
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MACITENTAN

TABLET; ORAL

MACITENTAN

ZYDUS	10MG	A211224 001	Apr 06, 2021
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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
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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
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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	
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SOLUTION; IRRIGATION

PHYSIOSOL IN PLASTIC CONTAINER

HOSPIRA INC	14MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML	N018406 001	
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PHYSIOSOL PH 7.4 IN PLASTIC CONTAINER

HOSPIRA INC	30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML	N018406 002	Jul 08, 1982
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SYNOVALYTE IN PLASTIC CONTAINER

BAXTER HLTHCARE	30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML	N019326 001	Jan 25, 1985
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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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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,N/A,N/A,210GM,0.74GM,2.86GM,5.6GM **	N203595 001	Jan 18, 2013
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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
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MALATHION

LOTION;TOPICAL

MALATHION

	MYLAN PHARMS INC	0.5%	A078743 001	Mar 06, 2009
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OVIDE

+	TARO PHARM INDS LTD	0.5% **	N018613 001	Aug 02, 1982
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MANGAFODIPIR TRISODIUM

INJECTABLE;INJECTION

TESLASCAN

	IC TARGETS	37.9MG/ML	N020652 001	Nov 26, 1997
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MANGANESE CHLORIDE TETRAHYDRATE

FOR SOLUTION;ORAL

LUMENHANCE

	BRACCO	3.49MG/GM	N020686 001	Dec 19, 1997
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MANGANESE SULFATE

INJECTABLE;INJECTION

MANGANESE SULFATE

+	ABRAXIS PHARM	EQ 0.1MG MANGANESE/ML **	N019228 001	May 05, 1987
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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
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MANNITOL 10% W/ DEXTROSE 5% IN DISTILLED WATER

	B BRAUN	10GM/100ML	N016080 006	
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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
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MANNITOL 15% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.45%

	B BRAUN	15GM/100ML	N016080 005	
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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
	INTL MEDICATION	12.5GM/50ML	A083051 001	
	LUITPOLD	12.5GM/50ML	A087409 001	Jan 21, 1982
	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
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MANNITOL

INJECTABLE; INJECTION

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

HERITAGE PHARMA 25MG A072162 001 Jun 01, 1988

50MG A072163 001 Jun 01, 1988

RISING PHARMA 25MG A072285 002 Oct 03, 1988

50MG A072285 001 Oct 03, 1988

75MG A072285 003 Oct 03, 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

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

+ NOVARTIS 1MG ** N017247 001

+ 2MG ** N017247 002

MEBENDAZOLE

TABLET, CHEWABLE; ORAL

VERMOX

+ JANSSEN PHARMS 100MG ** N017481 001

+ 500MG N208398 001 Oct 19, 2016

MEBUTAMATE

TABLET; ORAL

DORMATE

MEDPOINTE PHARM HLC 600MG N017374 001

MECAMYLAMINE HYDROCHLORIDE

TABLET; ORAL

INVERSINE

+ TARGACEPT 2.5MG ** N010251 001

MECHLORETHAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

MUSTARGEN

+ RECORDATI RARE 10MG/VIAL N006695 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MECLIZINE HYDROCHLORIDE

TABLET;ORAL

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	12.5MG	A085269	001	
	12.5MG	A088732	001	Dec 11, 1985
	25MG	A085740	001	
AUROBINDO PHARMA USA	12.5MG	A202640	001	Sep 17, 2012
	25MG	A202640	002	Sep 17, 2012
	50MG	A202640	003	Sep 17, 2012
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	
PLIVA	25MG	A088734	001	Dec 11, 1985
RISING	12.5MG	A040179	001	Jan 30, 1997
	25MG	A040179	002	Jan 30, 1997
STRIDES PHARMA	50MG	A089674	001	Mar 31, 1988
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	

TABLET, CHEWABLE;ORAL

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	
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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
ANI PHARMS	EQ 50MG BASE	A071469	002	Apr 15, 1987
	EQ 100MG BASE	A071469	001	Apr 15, 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	A071640	001	Aug 11, 1987
	EQ 100MG BASE	A070401	001	Nov 25, 1986
	EQ 100MG BASE	A071641	001	Aug 11, 1987

MECLOMEN

PARKE DAVIS	EQ 50MG BASE	N018006	001	
	EQ 100MG BASE	N018006	002	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEDROXYPROGESTERONE ACETATE

INJECTABLE; INJECTION

DEPO-PROVERA

+	PFIZER	100MG/ML **	N012541	002	
+		400MG/ML	N012541	003	

MEDROXYPROGESTERONE ACETATE

	CIPLA	150MG/ML	A210335	001	Jan 25, 2019
	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	
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CURRETAB

	SOLVAY	10MG	A085686	001	
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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
	UPSHER SMITH LABS	10MG	A088484	001	Jul 26, 1984

MEDRYSONE

SUSPENSION; OPHTHALMIC

HMS

	ALLERGAN	1%	N016624	003	
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MEFENAMIC ACID

CAPSULE; ORAL

MEFENAMIC ACID

	NOSTRUM LABS INC	250MG	A090359	001	Feb 05, 2013
	STRIDES PHARMA	250MG	A209209	001	Sep 18, 2020

MEFLOQUINE HYDROCHLORIDE

TABLET; ORAL

LARIAM

+	ROCHE	250MG **	N019591	001	May 02, 1989
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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
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MEGESTROL ACETATE

	HIKMA	40MG/ML	A075997	001	Feb 15, 2002
	PHARM ASSOC	40MG/ML	A077404	001	Feb 16, 2006
	TEVA PHARMS	40MG/ML	A075681	001	May 05, 2003

TABLET; ORAL

MEGACE

+	BRISTOL MYERS SQUIBB	20MG **	N016979	001	
+		40MG **	N016979	002	

MEGESTROL ACETATE

	HIKMA	20MG	A074458	001	Sep 29, 1995
		40MG	A074458	002	Sep 29, 1995
	TEVA	40MG	A074745	001	Feb 27, 1998
	USL PHARMA	20MG	A070646	001	Oct 02, 1987
		40MG	A070647	001	Oct 02, 1987

MELOXICAM

CAPSULE; ORAL

VIVLODEX

+	ZYLA	5MG **	N207233	001	Oct 22, 2015
+		10MG **	N207233	002	Oct 22, 2015

SUSPENSION; ORAL

MOBIC

+	AVONDALE PHARMS	7.5MG/5ML **	N021530	001	Jun 01, 2004
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MELOXICAM

TABLET; ORAL

MELOXICAM

AIPING PHARM INC	7.5MG	A077920 001	Jul 19, 2006
	15MG	A077920 002	Jul 19, 2006
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
HERITAGE PHARMA	7.5MG	A077936 001	Jul 19, 2006
	15MG	A077936 002	Jul 19, 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

TABLET, ORALLY DISINTEGRATING; ORAL

QMIIZ ODT

+ TERSERA	7.5MG	N211210 001	Oct 19, 2018
+	15MG	N211210 002	Oct 19, 2018

MELPHALAN

TABLET; ORAL

ALKERAN

+ APOTEX	2MG	N014691 002	
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MELPHALAN FLUFENAMIDE HYDROCHLORIDE

POWDER; INTRAVENOUS

PEPAXTO

+ ONCOPEPTIDES AB	EQ 20MG BASE/VIAL	N214383 001	Feb 26, 2021
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MELPHALAN HYDROCHLORIDE

INJECTABLE; INJECTION

ALKERAN

+ APOTEX	EQ 50MG BASE/VIAL **	N020207 001	Nov 18, 1992
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MELPHALAN HYDROCHLORIDE

MYLAN INSTITUTIONAL	EQ 50MG BASE/VIAL	A090299 001	Oct 27, 2009
PAR STERILE PRODUCTS	EQ 50MG BASE/VIAL	A204773 001	Aug 22, 2016
SCINOPHARM TAIWAN	EQ 50MG BASE/VIAL	A211463 001	Sep 13, 2019
USWM	EQ 50MG BASE/VIAL	A207032 001	May 03, 2019

POWDER; INTRAVENOUS

MELPHALAN HYDROCHLORIDE

ACTAVIS LLC	EQ 50MG BASE/VIAL	A209323 001	Mar 06, 2020
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MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MEMANTINE HYDROCHLORIDE

RISING PHARMA	7MG	A206032 001	Sep 28, 2016
	14MG	A206032 002	Sep 28, 2016
	21MG	A206032 003	Sep 28, 2016
	28MG	A206032 004	Sep 28, 2016
SUN PHARM	7MG	A205905 001	Sep 28, 2016
	14MG	A205905 002	Sep 28, 2016
	21MG	A205905 003	Sep 28, 2016
	28MG	A205905 004	Sep 28, 2016

SOLUTION; ORAL

MEMANTINE HYDROCHLORIDE

TORRENT	2MG/ML	A205446 001	Dec 07, 2015
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NAMENDA

+ ALLERGAN	2MG/ML	N021627 001	Apr 18, 2005
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TABLET; ORAL

MEMANTINE HYDROCHLORIDE

CHARTWELL	5MG	A090244 001	Jul 11, 2018
	10MG	A090244 002	Jul 11, 2018

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEMANTINE HYDROCHLORIDE

TABLET; ORAL

MEMANTINE HYDROCHLORIDE

JUBILANT GENERICS	5MG	A091585 001	Oct 13, 2015
	10MG	A091585 002	Oct 13, 2015
MYLAN	5MG	A079225 001	Jan 30, 2015
	10MG	A079225 002	Jan 30, 2015
NINGBO	5MG	A212947 001	Apr 03, 2020
	10MG	A212947 002	Apr 03, 2020
ORBION PHARMS	5MG	A090044 001	Mar 12, 2012
	10MG	A090044 002	Mar 12, 2012
TEVA PHARMS	5MG	A090052 001	Oct 25, 2011
	10MG	A090052 002	Oct 25, 2011
TORRENT	5MG	A200155 001	Oct 13, 2015
	10MG	A200155 002	Oct 13, 2015

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	
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MENADIONE

TABLET; ORAL

MENADIONE

LILLY	5MG	N002139 003	
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MEPENZOLATE BROMIDE

SOLUTION; ORAL

CANTIL

SANOFI AVENTIS US	25MG/5ML	N010679 004	
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TABLET; ORAL

CANTIL

+ SANOFI AVENTIS US	25MG	N010679 003	
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MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DEMEROL

VALIDUS PHARMS	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	
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PARKE DAVIS

50MG/ML	A080364 002	
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75MG/ML	A080364 003	
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100MG/ML	A080364 001	
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WATSON LABS

50MG/ML	A073444 001	Mar 17, 1992
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPERIDINE HYDROCHLORIDE

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

VALIDUS PHARMS

50MG/5ML **

N005010 005

TABLET; ORAL

DEMEROL

+ VALIDUS PHARMS

50MG

N005010 001

+

100MG

N005010 004

MEPERIDINE HYDROCHLORIDE

ANDA REPOSITORY

50MG

A040893 001 Jun 24, 2009

75MG

A040893 002 Jun 24, 2009

100MG

A040893 003 Jun 24, 2009

150MG

A040893 004 Jun 24, 2009

BARR

50MG

A088639 001 Jul 02, 1984

100MG

A088640 001 Sep 19, 1984

DURAMED PHARMS BARR

50MG

A040318 001 Oct 05, 1999

100MG

A040318 002 Oct 05, 1999

HIKMA

50MG

A040110 001 Mar 12, 1997

100MG

A040110 002 Mar 12, 1997

SPECGX LLC

50MG

A040352 001 Jun 13, 2000

100MG

A040352 002 Jun 13, 2000

STRIDES PHARMA

50MG

A040191 001 Dec 17, 1998

100MG

A040191 002 Dec 17, 1998

SUN PHARM INDS INC

50MG

A040446 001 Aug 08, 2002

100MG

A040446 002 Aug 08, 2002

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

HIKMA

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

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
400MG

N011284 001

N011284 002

TABLET; ORAL

AMOSENE

FERNDALE LABS 400MG

A084030 001

BAMATE

ALRA 200MG
400MG

A080380 001

A080380 002

EQUANIL

WYETH AYERST 200MG
400MG

N010028 005

N010028 004

MEPRIAM

TEVA 400MG

N016069 001

MEPROBAMATE

ACELLA 400MG

A084153 001

BARR 600MG

A084230 001

ELKINS SINN 200MG

N015426 002

400MG

N015426 001

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

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

RISING 400MG

A080655 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEPROBAMATE

TABLET; ORAL

MEPROBAMATE

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		A083304 001	
	200MG		A085720 001	
+	400MG		A083308 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	

MEQUINOL; TRETINOIN

SOLUTION; TOPICAL

SOLAGE

ALMIRALL	2%;0.01%		N020922 001	Dec 10, 1999
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MEROPENEM

INJECTABLE; INJECTION

MEROPENEM

HOSPIRA INC	500MG/VIAL		A090940 001	Jun 22, 2010
	1GM/VIAL		A090940 002	Jun 22, 2010
SANDOZ	500MG/VIAL		A091201 001	Mar 29, 2011
	1GM/VIAL		A091201 002	Mar 29, 2011

MERSALYL SODIUM; THEOPHYLLINE

INJECTABLE; INJECTION

MERSALYL-THEOPHYLLINE

WATSON LABS	100MG/ML; 50MG/ML		A084875 001	
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MESALAMINE

ENEMA; RECTAL

MESALAMINE

G AND W LABS INC	4GM/60ML		A076841 001	Sep 30, 2004
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SUPPOSITORY; RECTAL

CANASA

ALLERGAN	500MG		N021252 001	Jan 05, 2001
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ROWASA

+ MEDA PHARMS	500MG **		N019919 001	Dec 18, 1990
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TABLET, DELAYED RELEASE; ORAL

ASACOL

APIL	400MG		N019651 001	Jan 31, 1992
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MESALAMINE

MYLAN	1.2GM		A203574 001	Nov 20, 2018
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MESNA

INJECTABLE; INTRAVENOUS

MESNA

MYLAN INSTITUTIONAL	100MG/ML		A076488 001	Mar 08, 2012
MYLAN LABS LTD	100MG/ML		A203364 001	Jul 18, 2014
TEVA PHARMS USA	100MG/ML		A075764 001	Apr 27, 2001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MESORIDAZINE BESYLATE

CONCENTRATE; 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; NORETHINDRONE

TABLET; ORAL-20

NORINYL

ACTAVIS LABS UT INC

0.1MG; 2MG

N013625 004

TABLET; ORAL-21

NORETHIN 1/50M-21

HERITAGE PHARMA

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

TABLET; ORAL-28

NORETHIN 1/50M-28

HERITAGE PHARMA

0.05MG; 1MG

A071540 001 Apr 12, 1988

NORETHINDRONE AND MESTRANOL

WATSON LABS

0.05MG; 1MG

A070759 001 Jul 01, 1988

NORINYL 1+50 28-DAY

+ ACTAVIS LABS UT INC

0.05MG; 1MG

N016659 001

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
0.15MG; 9.85MG

N010976 008

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%
0.6%

N018761 002 Oct 10, 1986

N018761 001 Jun 30, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METAPROTERENOL SULFATE

SOLUTION; INHALATION

ALUPENT

	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
COSETTE	10MG/5ML	A072761 001	Feb 27, 1992
G AND W LABS INC	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
	20MG	A072055 001	Jun 23, 1988
HERITAGE PHARMA	10MG	A072519 001	Mar 30, 1990
	20MG	A072520 001	Mar 30, 1990
STRIDES PHARMA	10MG	A072024 001	Jun 28, 1988
	20MG	A072025 001	Jun 28, 1988
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

INGENUS PHARMS LLC	800MG	A213836 001	Oct 21, 2020
+ PRIMUS PHARMS	640MG **	N022503 001	Jun 01, 2015
SKELAXIN			
+ KING PHARMS	400MG **	N013217 001	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METFORMIN HYDROCHLORIDE

FOR SUSPENSION, EXTENDED RELEASE;ORAL

RIOMET ER

+ SUN PHARM 500MG/5ML N212595 001 Aug 29, 2019

TABLET;ORAL

GLUCOPHAGE

+ EMD SERONO INC 500MG ** N020357 001 Mar 03, 1995

+ 625MG ** N020357 003 Nov 05, 1998

+ 750MG ** N020357 004 Nov 05, 1998

+ 850MG ** N020357 002 Mar 03, 1995

+ 1GM ** N020357 005 Nov 05, 1998

METFORMIN HYDROCHLORIDE

BARR 500MG A075971 001 Jan 25, 2002

850MG A075971 002 Jan 25, 2002

1GM A075971 003 Jan 25, 2002

HERITAGE PHARMA 500MG A075978 001 Jan 25, 2002

850MG A075978 002 Jan 25, 2002

1GM A075978 003 Nov 05, 2002

INDICUS PHARMA 500MG A079148 001 Nov 25, 2008

850MG A079148 002 Nov 25, 2008

1GM A079148 003 Nov 25, 2008

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

MACLEODS PHARMS LTD 500MG A205330 001 Oct 31, 2017

850MG A205330 002 Oct 31, 2017

1GM A205330 003 Oct 31, 2017

MYLAN 500MG A075976 001 Jan 24, 2002

850MG A075976 002 Jan 24, 2002

1GM A075976 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

SUN PHARM INDUSTRIES 500MG A076038 001 Feb 21, 2002

850MG A076038 002 Feb 21, 2002

1GM A076038 003 Feb 21, 2002

SUNSHINE 500MG A208999 001 Oct 12, 2018

850MG A208999 002 Oct 12, 2018

1GM A208999 003 Oct 12, 2018

TEVA 500MG A076328 001 Dec 16, 2002

850MG A076328 002 Dec 16, 2002

1GM A076328 003 Dec 16, 2002

TORRENT PHARMS 500MG A077711 001 Jan 24, 2007

850MG A077711 002 Jan 24, 2007

1GM A077711 003 Jan 24, 2007

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

FORTAMET

+ ANDRX LABS LLC 500MG ** N021574 001 Apr 27, 2004

GLUCOPHAGE XR

+ EMD SERONO INC 500MG ** N021202 001 Oct 13, 2000

+ 750MG ** N021202 004 Apr 11, 2003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METFORMIN HYDROCHLORIDE

ACTAVIS ELIZABETH	500MG	A076450	001	Oct 01, 2004
	750MG	A076878	001	Apr 13, 2005
ACTAVIS LABS FL INC	500MG	A076172	001	Jun 16, 2004
APOTEX	500MG	A076706	001	Dec 14, 2004
	750MG	A076706	002	Dec 29, 2005
BARR	500MG	A076496	001	Nov 25, 2005
	750MG	A076863	001	Oct 14, 2004
IMPAX LABS	500MG	A076249	001	Jul 30, 2004
	750MG	A076985	001	Sep 13, 2005
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	500MG	A090014	001	Dec 30, 2009
TORRENT PHARMS LTD	750MG	A079226	001	Feb 18, 2010
WATSON LABS INC	500MG	A076818	001	Dec 14, 2004

METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE

MYLAN	500MG;EQ 15MG BASE	A090406	001	Feb 25, 2011
	850MG;EQ 15MG BASE	A090406	002	Feb 25, 2011
SANDOZ	500MG;EQ 15MG BASE	A091273	001	Apr 16, 2013
	850MG;EQ 15MG BASE	A091273	002	Apr 16, 2013
TORRENT PHARMS LTD	500MG;EQ 15MG BASE	A202001	001	Feb 13, 2013
	850MG;EQ 15MG BASE	A202001	002	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; 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
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
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METHADONE HYDROCHLORIDE

CONCENTRATE; ORAL

METHADONE HYDROCHLORIDE

LANNETT CO INC	10MG/ML	A212094	001	Mar 03, 2021
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POWDER; FOR RX COMPOUNDING

METHADONE HYDROCHLORIDE

MALLINCKRODT INC	50GM/BOT	N006383	002
	100GM/BOT	N006383	003
	500GM/BOT	N006383	004

SYRUP; ORAL

DOLOPHINE HYDROCHLORIDE

HIKMA	10MG/30ML	N006134	004
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TABLET; ORAL

DOLOPHINE HYDROCHLORIDE

+ HIKMA	5MG **	N006134	002
+ HIKMA	10MG **	N006134	010

METHADONE HYDROCHLORIDE

ROXANE	40MG	A074081	001	Apr 28, 1995
VISTAPHARM	5MG	A040241	001	May 29, 1998

METHADOSE

SPECGX LLC	5MG	A040050	001	Apr 15, 1993
	10MG	A040050	002	Apr 15, 1993

TABLET, DISPERSIBLE; ORAL

WESTADONE

SANDOZ	2.5MG	N017108	001
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TABLET, EFFERVESCENT; ORAL

WESTADONE

SANDOZ	5MG	N017108	002
	10MG	N017108	003
	40MG	N017108	004

METHAMPHETAMINE HYDROCHLORIDE

TABLET; ORAL

METHAMPEX

TEVA	10MG	A083889	001
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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
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METHARBITAL

TABLET; ORAL

GEMONIL

ABBVIE	100MG	N008322	001
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METHAZOLAMIDE

TABLET; ORAL

METHAZOLAMIDE

APPLIED ANAL	25MG	A040011	001	Jul 17, 1997
	50MG	A040011	002	Jul 17, 1997
ATHEM	25MG	A040102	001	Aug 28, 1996
	50MG	A040102	002	Aug 28, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHAZOLAMIDE

TABLET; ORAL

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

METHENAMINE HIPPURATE

TABLET; ORAL

HIPREX

+ VALIDUS PHARMS

1GM

N017681 001

METHENAMINE HIPPURATE

IMPAX LABS INC

1GM

A076411 001 Jun 20, 2003

METHICILLIN SODIUM

INJECTABLE; INJECTION

STAPHICILLIN

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

METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

BIOPHARM

20MG

A040547 004 Feb 18, 2005

DISCOVERY THERAP

15MG

A040619 003 Jul 12, 2005

MYLAN

20MG

A040350 003 Jun 07, 2001

SUN PHARM INDS INC

5MG

A040870 001 Sep 25, 2007

10MG

A040870 002 Sep 25, 2007

TAPAZOLE

+ KING PHARMS

5MG **

N007517 002

+

10MG **

N007517 004

KING PHARMS LLC

5MG

A040320 001 Mar 31, 2000

10MG

A040320 002 Mar 31, 2000

METHIXENE HYDROCHLORIDE

TABLET; ORAL

TREST

NOVARTIS

1MG

N013420 001

METHOCARBAMOL

INJECTABLE; INJECTION

METHOCARBAMOL

DR REDDYS

100MG/ML

A086459 001

MARSAM PHARMS LLC

100MG/ML

A089849 001 Dec 27, 1991

TABLET; ORAL

DELAXIN

FERNDAL 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

ANI PHARMS

500MG

A084277 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL

	750MG		A084276	002	
ASCOT	500MG		A087660	001	Oct 27, 1982
	750MG		A087661	001	Oct 27, 1982
AUROBINDO PHARMA LTD	750MG		A213967	001	Aug 12, 2020
CLONMEL HLTHCARE	500MG		A085961	001	
	750MG		A085963	001	
FOSUN PHARMA	500MG		A084616	001	
	750MG		A084615	001	
HEATHER	500MG		A084675	001	
	750MG		A084924	001	
HIKMA INTL PHARMS	500MG		A085159	001	
	750MG		A085123	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	
	500MG		A085180	001	
	750MG		A083605	002	
	750MG		A085192	001	
ROBAXIN					
+ AUXILIUM PHARMS LLC	500MG **		N011011	004	
ROBAXIN-750					
+ AUXILIUM PHARMS LLC	750MG **		N011011	006	

METHOHEXITAL SODIUM

INJECTABLE; INJECTION

BREVITAL SODIUM

PAR STERILE PRODUCTS	200MG/VIAL		N011559	004	Dec 21, 2012
+	2.5GM/VIAL		N011559	002	
	5GM/VIAL		N011559	003	

METHOTREXATE

SOLUTION; INTRAVENOUS

METHOTREXATE

+ ACCORD HLTHCARE	5GM/50ML (100MG/ML)		N214121	001	Aug 24, 2020
SOLUTION; SUBCUTANEOUS					
OTREXUP					
+ OTTER PHARMS	7.5MG/0.4ML (7.5MG/0.4ML)		N204824	005	Nov 07, 2014

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHOTREXATE

SOLUTION; SUBCUTANEOUS

OTREXUP PFS

+	OTTER PHARMS	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

+	MEDEXUS	27.5MG/0.55ML (27.5MG/0.55ML)	N205776	009	Jul 10, 2014
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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
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METHOTREXATE PRESERVATIVE FREE

	FRESENIUS KABI USA	EQ 25MG BASE/ML	A040265	001	Feb 26, 1999
+	HOSPIRA	EQ 2.5GM BASE/100ML (EQ 25MG BASE/ML)	N011719	011	Apr 13, 2005
+		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

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

	AUROMEDICS PHARMA	EQ 50MG BASE/2ML (EQ 25MG BASE/ML)	A201529	001	Mar 29, 2012
		EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	A201529	002	Mar 29, 2012
		EQ 200MG BASE/8ML (EQ 25MG BASE/ML)	A201529	003	Mar 29, 2012
		EQ 250MG BASE/10ML (EQ 25MG BASE/ML)	A201529	004	Mar 29, 2012
+	HOSPIRA	EQ 1GM BASE/VIAL	N011719	009	Apr 07, 1988
	MYLAN	EQ 1GM BASE/40ML (EQ 25MG BASE/ML)	A201530	001	Mar 29, 2012

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	

MEXATE-AQ

	BRISTOL MYERS	EQ 25MG BASE/ML	A088760	001	Feb 14, 1985
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MEXATE-AQ PRESERVED

	BRISTOL MYERS SQUIBB	EQ 25MG BASE/ML	A089887	001	Apr 14, 1989
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TABLET; ORAL

METHOTREXATE SODIUM

	DURAMED PHARMS BARR	EQ 2.5MG BASE	A040233	001	Jun 17, 1999
+	STRIDES PHARMA	EQ 2.5MG BASE **	N008085	002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
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METHOXSALEN

ACTAVIS INC	10MG	A202603	001 Jun 09, 2015
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ANI PHARMS	10MG	A087781	001 Jun 08, 1982
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LOTION; TOPICAL

OXSORALEN

+ VALEANT PHARM INTL	1%	N009048	002
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METHSCOPOLAMINE BROMIDE

TABLET; ORAL

METHSCOPOLAMINE BROMIDE

PVT FORM	2.5MG	A080970	001
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STRIDES PHARMA	2.5MG	A040624	001 Dec 28, 2006
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	5MG	A040624	002 Dec 28, 2006
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PAMINE

FOUGERA PHARMS	2.5MG **	N008848	001
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PAMINE FORTE

FOUGERA PHARMS	5MG **	N008848	002 Mar 25, 2003
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METHSUXIMIDE

CAPSULE; ORAL

CELONTIN

+ PARKE DAVIS	150MG	N010596	007
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METHYCLOTHIAZIDE

TABLET; ORAL

AQUATENSEN

MEDPOINTE PHARM HLC	5MG	N017364	001
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ENDURON

+ ABBVIE	2.5MG **	N012524	001
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+	5MG **	N012524	004
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METHYCLOTHIAZIDE

CHARTWELL RX	2.5MG	A089835	001 Aug 18, 1988
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	5MG	A089837	001 Aug 18, 1988
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IVAX PHARMS	2.5MG	A087913	001 Jun 03, 1982
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	5MG	A087786	001 May 18, 1982
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MYLAN	2.5MG	A087671	001 Aug 17, 1982
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MYLAN PHARMS INC	5MG	A087672	001 Aug 17, 1982
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PAR PHARM	2.5MG	A089135	001 Feb 12, 1986
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	5MG	A089136	001 Feb 12, 1986
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USL PHARMA	5MG	A088745	001 Mar 21, 1985
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WATSON LABS	2.5MG	A085487	001 Mar 11, 1982
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	2.5MG	A088750	001 Sep 06, 1984
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	5MG	A085476	001 Mar 11, 1982
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	5MG	A088724	001 Sep 06, 1984
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METHYCLOTHIAZIDE; PARGYLINE HYDROCHLORIDE

TABLET; ORAL

EUTRON

ABBOTT	5MG; 25MG	N016047	001
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METHYCLOTHIAZIDE; RESERPINE

TABLET; ORAL

DIUTENSEN-R

MEDPOINTE PHARM HLC	2.5MG; 0.1MG	N012708	005
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METHYL AMINOLEVULINATE HYDROCHLORIDE

CREAM; TOPICAL

METVIXIA

GALDERMA LABS LP	EQ 16.8% BASE	N021415	001 Jul 27, 2004
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLDOPA

SUSPENSION; ORAL

ALDOMET

MERCK	250MG/5ML	N018389	001
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TABLET; ORAL

ALDOMET

+	MERCK	125MG **	N013400	003
+		250MG **	N013400	001
+		500MG **	N013400	002

METHYLDOPA

ACCORD HLTHCARE	125MG	A070070	003	Oct 15, 1985
CHARTWELL RX	125MG	A071700	001	Mar 02, 1988
	250MG	N018934	001	Jun 29, 1984
	500MG	N018934	002	Jun 29, 1984
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
HERITAGE PHARMA	250MG	A070098	001	Feb 20, 1986
	500MG	A070343	001	Feb 20, 1986
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
RISING PHARMA	250MG	A070076	002	Apr 18, 1985
	500MG	A070076	001	Apr 18, 1985
ROXANE	125MG	A070192	001	Apr 25, 1986
	250MG	A070193	001	Apr 25, 1986
	500MG	A070194	001	Apr 25, 1986
STRIDES PHARMA	125MG	A070535	001	Jan 02, 1987
	250MG	A070536	001	Jan 02, 1987
	500MG	A070537	001	Jan 02, 1987
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
	500MG	A070625	001	Jun 06, 1986

METHYLDOPATE HYDROCHLORIDE

INJECTABLE; INJECTION

ALDOMET

+	MERCK	50MG/ML **	N013401	001
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METHYLDOPATE HYDROCHLORIDE

ABRAXIS PHARM	50MG/ML	A070652	001	Jun 03, 1986
AM REGENT	50MG/ML	A071279	001	Oct 02, 1987
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 effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLERGONOVINE MALEATE

TABLET;ORAL

METHERGINE

+ EDISON THERAPS LLC 0.2MG ** N006035 003

METHYLPHENIDATE

TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE;ORAL

METHYLPHENIDATE

ACTAVIS ELIZABETH	8.6MG	A210924	001	Jun 19, 2020
	17.3MG	A210924	002	Jun 19, 2020
	25.9MG	A210924	003	Jun 19, 2020

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

BARR LABS INC	10MG	A079031	004	Oct 15, 2014
	20MG	A079031	001	Jul 13, 2012
	30MG	A079031	002	Jul 13, 2012
	40MG	A079031	003	Jul 13, 2012

RITALIN LA

+ NOVARTIS 60MG ** N021284 005 Oct 27, 2014

SOLUTION;ORAL

METHYLPHENIDATE HYDROCHLORIDE

LANNETT CO INC	5MG/5ML	A207414	001	Dec 16, 2020
	10MG/5ML	A207414	002	Dec 16, 2020

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
ALVOGEN	5MG	A206840	001	Sep 15, 2016
	10MG	A206840	002	Sep 15, 2016
	20MG	A206840	003	Sep 15, 2016
AUROLIFE PHARMA LLC	5MG	A209276	001	Oct 25, 2018
	10MG	A209276	002	Oct 25, 2018
	20MG	A209276	003	Oct 25, 2018
CEDIPROF INC	5MG	A208737	001	Feb 01, 2019
	10MG	A208737	002	Feb 01, 2019
	20MG	A208737	003	Feb 01, 2019
LANNETT CO INC	5MG	A086429	001	
	10MG	A085799	001	
	20MG	A086428	001	
NOSTRUM LABS INC	5MG	A207587	001	Mar 03, 2017
	10MG	A207587	002	Mar 03, 2017
	20MG	A207587	003	Mar 03, 2017
WATSON LABS	5MG	A040220	001	Aug 29, 1997
	10MG	A040220	002	Aug 29, 1997
	20MG	A040220	003	Aug 29, 1997

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

METHYLPHENIDATE HYDROCHLORIDE

NOSTRUM LABS INC	2.5MG	A204954	001	Jan 26, 2017
	5MG	A204954	002	Jan 26, 2017
	10MG	A204954	003	Jan 26, 2017
NOVEL LABS INC	2.5MG	A204115	001	Feb 25, 2015
	5MG	A204115	002	Feb 25, 2015
	10MG	A204115	003	Feb 25, 2015

TABLET, EXTENDED RELEASE;ORAL

METADATE ER

LANNETT CO INC	10MG	A040306	001	Oct 20, 1999
	20MG	A089601	001	Jun 01, 1988

METHYLPHENIDATE HYDROCHLORIDE

ABLE	20MG	A076032	001	May 09, 2001
ANI PHARMS	18MG	A208607	001	Jul 14, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLPHENIDATE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

	27MG	A208607 002	Jul 14, 2017
	36MG	A208607 003	Jul 14, 2017
	54MG	A208607 004	Jul 14, 2017
HERITAGE PHARMA	20MG	A075450 001	Dec 21, 2001
STRIDES PHARMA	36MG	A204659 001	Jul 15, 2019
	54MG	A204659 002	Jul 15, 2019
WATSON LABS	20MG	A040410 001	Feb 09, 2001
RITALIN-SR			
+ NOVARTIS	20MG **	N018029 001	Mar 30, 1982

METHYLPREDNISOLONE

TABLET;ORAL

MEDROL

PFIZER	24MG	N011153 005	
METHYLPREDNISOLONE			
DURAMED PHARMS BARR	4MG	A088497 001	Feb 21, 1984
HEATHER	4MG	A085650 001	
INVATECH	4MG	A087341 001	
LUPIN LTD	2MG	A209097 001	Feb 22, 2019
	4MG	A209097 002	Feb 22, 2019
	8MG	A209097 003	Feb 22, 2019
	16MG	A209097 004	Feb 22, 2019
	32MG	A209097 005	Feb 22, 2019
NOVAST LABS	4MG	A210985 001	Jan 09, 2019
PAR PHARM	16MG	A089207 001	Apr 25, 1988
	24MG	A089208 001	Apr 25, 1988
	32MG	A089209 001	Apr 25, 1988
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	

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	A040664 001	Dec 20, 2005
	EQ 40MG BASE/VIAL	A085853 001	
	EQ 125MG BASE/VIAL	A040665 001	Dec 20, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

A-METHAPRED

	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

INTL MEDICATION

	EQ 40MG BASE/VIAL	A086906	001	
	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

TIANJIN KINGYORK

	EQ 1GM BASE/VIAL	A212396	004	Apr 20, 2021
	EQ 2GM BASE/VIAL	A212396	005	Apr 20, 2021
	EQ 40MG BASE/VIAL	A212396	001	Apr 20, 2021
	EQ 125MG BASE/VIAL	A212396	002	Apr 20, 2021
	EQ 500MG BASE/VIAL	A212396	003	Apr 20, 2021

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	
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METHYLTESTOSTERONE

CAPSULE; ORAL

METHYLTESTOSTERONE

HEATHER	10MG	A084967	001	
TESTRED				
+ BAUSCH	10MG	A083976	001	
VIRILON				
CHARTWELL	10MG	A087750	001	Nov 24, 1982

TABLET; BUCCAL

ANDROID 5

VALEANT PHARM INTL	5MG	A087222	001	
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ORETON

SCHERING	10MG	A080281	001	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLTESTOSTERONETABLET;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
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METHYLTESTOSTERONE

IMPAX LABS	25MG	A084310 001
INWOOD LABS	10MG	A080839 001
	25MG	A080973 001
KV PHARM	10MG	A084312 001
LANNETT	10MG	A087092 001
	25MG	A087111 001
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

Nov 05, 1982
Jan 27, 1983

ORETON METHYL

SCHERING	10MG	N003158 001
	25MG	N003158 002

METHYPRYLON

CAPSULE;ORAL

NOLUDAR

ROCHE	300MG	N009660 008
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ELIXIR;ORAL

NOLUDAR

ROCHE	50MG/5ML	N009660 007
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TABLET;ORAL

NOLUDAR

ROCHE	50MG	N009660 002
	200MG	N009660 004

METHYSERGIDE MALEATE

TABLET;ORAL

SANSERT

NOVARTIS	2MG	N012516 001
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METIPRANOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

METIPRANOLOL

SANDOZ INC	0.3%	A075720 001	Aug 06, 2001
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OPTIPRANOLOL

+ BAUSCH AND LOMB	0.3%	N019907 001	Dec 29, 1989
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METOCLOPRAMIDE HYDROCHLORIDE

CONCENTRATE; ORAL

METOCLOPRAMIDE INTENSOL

ROXANE	EQ 10MG BASE/ML	A072995 001	Jan 30, 1992
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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

+ HIKMA	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
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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
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MAXOLON

KING PHARMS	EQ 10MG BASE	A070106 001	Mar 04, 1986
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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

ANI PHARMS

2.5MG
5MG
10MGA075543 001 Jan 06, 2004
A075543 002 Mar 01, 2004
A075543 003 Dec 24, 2003

ROXANE

10MG

A076482 002 Apr 29, 2004

WATSON LABS

10MG

A076891 001 Jul 21, 2004

MYKROX

LANNETT CO INC

0.5MG

N019532 001 Oct 30, 1987

ZAROXOLYN

+ LANNETT CO INC

2.5MG

N017386 001

+

5MG

N017386 002

+

10MG

N017386 003

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

ACTAVIS LABS FL INC

EQ 25MG TARTRATE
EQ 100MG TARTRATE
EQ 200MG TARTRATEA076862 002 Aug 03, 2009
A077298 001 Apr 15, 2010
A077298 002 Apr 15, 2010

NESHER PHARMS

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

SANDOZ

EQ 25MG TARTRATE
EQ 50MG TARTRATE
EQ 100MG TARTRATE
EQ 200MG TARTRATEA076969 001 Jul 31, 2006
A076969 002 May 18, 2007
A076969 003 Mar 20, 2008
A076969 004 Mar 20, 2008METOPROLOL TARTRATE

INJECTABLE; INJECTION

LOPRESSOR

+ NOVARTIS

1MG/ML **

N018704 001 Mar 30, 1984

METOPROLOL TARTRATE

AM REGENT

1MG/ML

A090386 001 Sep 30, 2009

LUITPOLD

1MG/ML

A091307 001 Dec 29, 2010

MYLAN ASI

1MG/ML

A090317 001 Apr 19, 2010

WATSON LABS

1MG/ML

A074032 001 Dec 21, 1993

TABLET;ORAL

METOPROLOL TARTRATE

APOTHECON

50MG
100MGA074258 001 Jan 27, 1994
A074258 002 Jan 27, 1994

HERITAGE PHARMA

50MG
100MGA074141 001 Jan 31, 1995
A074141 002 Jan 31, 1995

MYLAN

50MG
100MGA073666 001 Dec 21, 1993
A073666 002 Dec 21, 1993

PUREPAC PHARM

50MG
100MGA074380 001 Jul 29, 1994
A074380 002 Jul 29, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METOPROLOL TARTRATE

TABLET; ORAL

METOPROLOL TARTRATE

RENATA	50MG	A074453 001	Apr 27, 1995
	100MG	A074453 002	Apr 27, 1995
SUN PHARM INDUSTRIES	25MG	A073654 002	Jul 15, 2009
	50MG	A073654 003	Dec 21, 1993
	100MG	A073654 001	Dec 21, 1993
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
WATSON LABS	50MG	A074217 001	May 27, 1994
	100MG	A074217 002	May 27, 1994
YAOPHARMA CO LTD	50MG	A073288 001	Mar 25, 1994
	100MG	A073289 001	Mar 25, 1994

METRIXAMIDE

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
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INJECTABLE; INJECTION

FLAGYL I.V. RTU IN PLASTIC CONTAINER

PFIZER	500MG/100ML	N018353 002	
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METRO I.V.

B BRAUN	500MG/100ML	N018674 001	Aug 31, 1982
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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

METRONIDAZOLE IN PLASTIC CONTAINER

MYLAN LABS LTD	500MG/100ML	A205531 001	May 09, 2017
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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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METRONIDAZOLE

TABLET; ORAL

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

+ PFIZER

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

METYRAPONE

TABLET; ORAL

METOPIRONE

HRA PHARMA 250MG N012911 001

MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

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

MICAFUNGIN SODIUM

POWDER; INTRAVENOUS

MICAFUNGIN

+ TEVA PHARMS USA INC EQ 50MG BASE/VIAL N212125 001 Jul 30, 2021

+ EQ 100MG BASE/VIAL N212125 002 Jul 30, 2021

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

COSETTE 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

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

AKORN 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

ATLANTIDE 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

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 1MG BASE/ML A075856 001 Jun 13, 2002

EQ 5MG BASE/ML A075396 002 Jun 20, 2000

EQ 5MG BASE/ML A075484 001 Jun 20, 2000

EQ 5MG BASE/ML A075856 002 Jun 13, 2002

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

MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE

MYLAN ASI EQ 1MG BASE/ML A090315 001 Nov 29, 2010

EQ 5MG BASE/ML A090315 002 Nov 29, 2010

MIDOZALAM HYDROCHLORIDE

MYLAN ASI EQ 1MG BASE/ML A090316 001 May 04, 2011

EQ 5MG BASE/ML A090316 002 May 04, 2011

VERSED

+ HLR EQ 1MG BASE/ML ** N018654 002 May 26, 1987

+ EQ 5MG BASE/ML ** N018654 001 Dec 20, 1985

SYRUP; ORAL

MIDAZOLAM HYDROCHLORIDE

PHARM ASSOC 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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MIDODRINE HYDROCHLORIDE

TABLET; ORAL

MIDODRINE HYDROCHLORIDE

CHARTWELL RX	2.5MG	A076514 001	Sep 11, 2003
	5MG	A076514 002	Sep 11, 2003
	10MG	A076514 003	Jul 02, 2004

PROAMATINE

+ TAKEDA PHARMS USA	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
GLAND PHARMA LTD	EQ 1MG BASE/ML	A077190 001	Oct 31, 2006
HIKMA	EQ 1MG BASE/ML	A075852 001	May 28, 2002
HOSPIRA	EQ 1MG BASE/ML	A075830 001	May 28, 2002
	EQ 1MG BASE/ML	A075884 001	May 28, 2002
INTL MEDICATED	EQ 1MG BASE/ML	A076013 001	Aug 02, 2002
MYLAN INSTITUTIONAL	EQ 1MG BASE/ML	A076428 001	Jun 16, 2003

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
HIKMA	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A075510 001	May 28, 2002
WOODWARD	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A077151 001	Jul 20, 2005

PRIMACOR

+ SANOFI AVENTIS US	EQ 1MG BASE/ML **	N019436 001	Dec 31, 1987
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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

+ BAUSCH	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

JOURNEY	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	
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SUSPENSION; ORAL

MINOCIN

BAUSCH	EQ 50MG BASE/5ML	N050445 001	
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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
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MINOCYCLINE HYDROCHLORIDETABLET, EXTENDED RELEASE;ORAL
MINOCYCLINE HYDROCHLORIDE

	EQ 65MG BASE	A065485 004	May 18, 2012
	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 115MG BASE	A065485 005	May 18, 2012
	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
RISING	EQ 55MG BASE	A203443 001	Aug 21, 2019
	EQ 65MG BASE	A201467 001	Jul 30, 2019
	EQ 80MG BASE	A203443 002	Aug 22, 2014
	EQ 105MG BASE	A203443 003	Aug 22, 2014
	EQ 115MG BASE	A201467 002	Jul 30, 2019
SOLODYN			
+ BAUSCH	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
COPLBY 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
AVACOR PRODS	5%	A075619 001	Nov 17, 2000

TABLET;ORAL

LONITEN			
+ PFIZER	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
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

MIRABEGRON

TABLET, EXTENDED RELEASE;ORAL

MIRABEGRON			
SAWAI USA	25MG	A209446 001	Dec 27, 2019

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MIRTAZAPINE

TABLET; ORAL

MIRTAZAPINE

	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
WATSON LABS	15MG	A076312 001	Jun 19, 2003
	30MG	A076312 002	Jun 19, 2003
	45MG	A076312 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
IMPAX LABS INC	15MG	A076901 001	Jun 28, 2005
	30MG	A076901 002	Jun 28, 2005
	45MG	A076901 003	Jun 28, 2005

MISOPROSTOL

TABLET; ORAL

MISOPROSTOL

ACQ PHARMA	0.1MG	A210201 001	Jul 02, 2019
	0.2MG	A210201 002	Jul 02, 2019
ANI PHARMS	0.1MG	A076095 001	Jul 10, 2002
	0.2MG	A076095 002	Jul 10, 2002

MITOMYCIN

INJECTABLE; INJECTION

MITOMYCIN

HIKMA	5MG/VIAL	A064117 001	Apr 19, 1995
HOSPIRA	20MG/VIAL	A064106 001	Nov 29, 1995
MITOZYTREX			
+ SUPERGEN	5MG/VIAL **	N050763 001	Nov 14, 2002
MUTAMYCIN			
+ BRISTOL	5MG/VIAL **	N050450 001	
+ BRISTOL MYERS	20MG/VIAL **	N050450 002	
	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
MYLAN LABS LTD	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	A201014 001	Dec 11, 2012
RISING	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	A078980 001	Apr 13, 2009
	EQ 30MG BASE/15ML (EQ 2MG BASE/ML)	A078980 002	Apr 13, 2009
NOVANTRONE			
+ EMD SERONO	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	N019297 001	Dec 23, 1987
+ EMD SERONO	EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML) **	N019297 002	Dec 23, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

NOVANTRONE

+

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

RISING PHARMA

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

MODAFINIL

TABLET; ORAL

MODAFINIL

HIKMA PHARMS

100MG

A090543 001 Sep 26, 2012

200MG

A090543 002 Sep 26, 2012

MYLAN PHARMS INC

100MG

A076594 001 Jul 16, 2012

200MG

A076594 002 Jul 16, 2012

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

+ STRIDES PHARMA

5MG **

N017111 001

+

10MG **

N017111 002

+

25MG **

N017111 003

CONCENTRATE; ORAL

MOBAN

ENDO PHARMS

20MG/ML

N017938 001

TABLET; ORAL

MOBAN

+ STRIDES PHARMA

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

+

0.1% **

N019625 002 Apr 19, 2013

MOMETASONE FUROATE

ANDA REPOSITORY

0.1%

A076591 001 Apr 18, 2007

FOUGERA PHARMS

0.1%

A076171 001 Apr 08, 2005

LOTION; TOPICAL

MOMETASONE FUROATE

ANDA REPOSITORY

0.1%

A076499 001 Nov 21, 2007

OINTMENT; TOPICAL

ELOCON

+ MERCK SHARP DOHME

0.1%

N019543 001 Apr 30, 1987

MOMETASONE FUROATE

ANDA REPOSITORY

0.1%

A076481 001 Nov 14, 2003

TARO

0.1%

A076624 001 Dec 03, 2004

TORRENT

0.1%

A207899 001 Jul 13, 2018

SPRAY, METERED; NASAL

NASONEX

+ ORGANON

0.05MG/SPRAY

N020762 001 Oct 01, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MONOBENZONE

CREAM;TOPICAL

BENOQUIN

VALEANT PHARM INTL 20%

N008173 003

MONOCTANOIN

LIQUID;PERFUSION, BILIARY

MOCTANIN

ETHITEK 100%

N019368 001 Oct 29, 1985

MONTELUKAST SODIUM

GRANULE;ORAL

MONTELUKAST SODIUM

MYLAN PHARMS INC EQ 4MG BASE/PACKET

A202776 001 Dec 18, 2012

TABLET;ORAL

MONTELUKAST SODIUM

AJANTA PHARMA LTD EQ 10MG BASE

A203432 001 Jul 31, 2015

APOTEX CORP EQ 10MG BASE

A201294 001 Aug 03, 2012

BRECKENRIDGE EQ 10MG BASE

A205319 001 Oct 30, 2020

HIKMA EQ 10MG BASE

A090655 001 Aug 03, 2012

MYLAN PHARMS INC EQ 10MG BASE

A079103 001 Aug 03, 2012

STRIDES PHARMA EQ 10MG BASE

A091576 001 Aug 03, 2012

TABLET, CHEWABLE;ORAL

MONTELUKAST SODIUM

AJANTA PHARMA LTD EQ 4MG BASE

A203328 001 Jul 31, 2015

EQ 5MG BASE

A203328 002 Jul 31, 2015

APOTEX INC EQ 4MG BASE

A201508 001 Aug 03, 2012

EQ 5MG BASE

A201508 002 Aug 03, 2012

HIKMA EQ 4MG BASE

A091128 001 Aug 03, 2012

EQ 5MG BASE

A091128 002 Aug 03, 2012

JUBILANT GENERICS EQ 4MG BASE

A203795 001 Feb 27, 2015

EQ 5MG BASE

A203795 002 Feb 27, 2015

MYLAN PHARMS INC EQ 4MG BASE

A079142 001 Aug 03, 2012

EQ 5MG BASE

A079142 002 Aug 03, 2012

STRIDES PHARMA EQ 4MG BASE

A091588 001 Aug 03, 2012

EQ 5MG BASE

A091588 002 Aug 03, 2012

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

KADIAN

+ ALLERGAN 10MG

N020616 008 Apr 20, 2007

+ 20MG

N020616 001 Jul 03, 1996

+ 30MG

N020616 004 Mar 09, 2001

+ 40MG

N020616 009 Jul 09, 2012

+ 50MG

N020616 002 Jul 03, 1996

+ 60MG

N020616 005 Mar 09, 2001

+ 70MG

N020616 010 Jul 09, 2012

+ 80MG

N020616 006 Oct 27, 2006

+ 100MG

N020616 003 Jul 03, 1996

+ 130MG

N020616 011 Jul 09, 2012

+ 150MG

N020616 012 Jul 09, 2012

+ 200MG

N020616 007 Feb 27, 2007

MORPHINE SULFATE

STRIDES PHARMA 20MG

A200812 001 Nov 10, 2011

30MG

A200812 002 Nov 10, 2011

50MG

A200812 003 Nov 10, 2011

60MG

A200812 004 Nov 10, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MORPHINE SULFATECAPSULE, EXTENDED RELEASE;ORAL
MORPHINE SULFATE

	80MG	A200812 005	Nov 10, 2011
	100MG	A200812 006	Nov 10, 2011
TEVA PHARMS USA	20MG	A202718 001	Dec 29, 2014
	30MG	A202718 002	Dec 29, 2014
	40MG	A202718 007	Jun 03, 2015
	50MG	A202718 003	Dec 29, 2014
	60MG	A202718 004	Dec 29, 2014
	70MG	A202718 008	Jun 03, 2015
	80MG	A202718 005	Dec 29, 2014
	100MG	A202718 006	Dec 29, 2014

INJECTABLE; INJECTION

ASTRAMORPH PF

FRESENIUS KABI USA	0.5MG/ML	A071050 001	Oct 07, 1986
	0.5MG/ML	A071051 001	Oct 07, 1986
	1MG/ML	A071052 001	Oct 07, 1986
	1MG/ML	A071053 001	Oct 07, 1986

MORPHINE SULFATE

HOSPIRA	1MG/ML	A071850 001	May 11, 1988
+ HOSPIRA INC	15MG/ML	N202515 005	Nov 14, 2011
ICU MEDICAL INC	0.5MG/ML	N019917 001	Oct 30, 1992
+	1MG/ML **	N019916 001	Oct 30, 1992
+	5MG/ML **	N019916 002	Oct 27, 2006
INTL MEDICATION SYS	1MG/ML	A202861 001	Apr 29, 2021
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

SOLUTION; INTRAMUSCULAR

MORPHINE SULFATE (AUTOINJECTOR)

+ MERIDIAN MEDCL TECHN	10MG/0.7ML (10MG/0.7ML)	N019999 001	Jul 12, 1990
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SOLUTION; ORAL

MORPHINE SULFATE

ANI PHARMS	10MG/5ML	A205509 001	Apr 17, 2018
	20MG/5ML	A205509 002	Apr 17, 2018
	100MG/5ML	A205509 003	Apr 17, 2018
LANNETT CO INC	10MG/5ML	A202309 001	Nov 25, 2015
	20MG/5ML	A202310 001	Oct 30, 2015
	100MG/5ML	N201517 001	Jun 23, 2011
NOSTRUM LABS INC	10MG/5ML	A201011 001	Feb 05, 2014
	20MG/5ML	A201011 002	Feb 05, 2014
	100MG/5ML	A201011 003	Oct 06, 2016
VISTAPHARM	10MG/5ML	A201947 001	Jan 05, 2012
	20MG/5ML	A201947 002	Jan 05, 2012
WINDER LABS LLC	10MG/5ML	A211454 001	Feb 12, 2021
	20MG/5ML	A211454 002	Feb 12, 2021
	100MG/5ML	A211454 003	Feb 12, 2021

TABLET, EXTENDED RELEASE; ORAL

ARYMO ER

+ ZYLA	15MG	N208603 001	Jan 09, 2017
+	30MG	N208603 002	Jan 09, 2017
+	60MG	N208603 003	Jan 09, 2017

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

EPIC PHARMA LLC	15MG	A091357 001	Jun 23, 2016
	30MG	A091357 002	Jun 23, 2016

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MORPHINE SULFATETABLET, EXTENDED RELEASE;ORAL
MORPHINE SULFATE

	60MG	A091357 003	Jun 23, 2016
	100MG	A091357 004	Jun 23, 2016
	200MG	A091357 005	Jun 23, 2016
NESHER PHARMS	15MG	A076733 001	May 19, 2004
	100MG	A077855 001	Sep 27, 2007
	200MG	A077855 002	Sep 27, 2007
RISING	15MG	A200824 001	Oct 18, 2011
	30MG	A200824 002	Oct 18, 2011
	60MG	A200824 003	Oct 18, 2011
	100MG	A200824 004	Oct 18, 2011
	200MG	A200824 005	Oct 18, 2011
SUN PHARM INDUSTRIES	15MG	A205634 001	Aug 25, 2016
	30MG	A205634 002	Aug 25, 2016
	60MG	A205634 003	Aug 25, 2016
	100MG	A205634 004	Aug 25, 2016
	200MG	A205634 005	Aug 25, 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

MORPHINE SULFATE; NALTREXONE HYDROCHLORIDECAPSULE, 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

MOXALACTAM DISODIUMINJECTABLE; 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

TABLET; ORAL

AVELOX				
+	BAYER HLTHCARE	EQ 400MG BASE	N021085 001	Dec 10, 1999
MOXIFLOXACIN HYDROCHLORIDE				
MYLAN	EQ 400MG BASE	A204635 001	Aug 31, 2015	
SUNSHINE	EQ 400MG BASE	A206295 001	Sep 28, 2018	

MUPIROCIIN

OINTMENT; TOPICAL

BACTROBAN				
+	GLAXOSMITHKLINE	2% **	N050591 001	Dec 31, 1987

MUPIROCIIN 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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
STRIDES PHARMA	250MG	A090111	001	Dec 22, 2009
ZYDUS PHARMS USA INC	250MG	A065433	001	May 04, 2009

TABLET; ORAL

MYCOPHENOLATE MOFETIL

AMNEAL	500MG	A090606	001	Jul 16, 2010
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
TEVA PHARMS	500MG	A065457	001	May 04, 2009
ZYDUS PHARMS USA INC	500MG	A065477	001	May 04, 2009

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

INJECTABLE; INJECTION

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

AMNEAL	500MG/VIAL	A211374	001	Mar 05, 2021
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MYCOPHENOLIC SODIUM

TABLET, DELAYED RELEASE; ORAL

MYCOPHENOLIC SODIUM

TEVA PHARMS USA	EQ 180MG BASE	A202720	001	Oct 30, 2014
	EQ 360MG BASE	A202720	002	Oct 30, 2014

NABUMETONE

TABLET; ORAL

NABUMETONE

COPLEY PHARM	750MG	A075179	001	Jun 06, 2000
IMPAX LABS INC	500MG	A075189	001	May 26, 2000
	750MG	A075189	002	Sep 24, 2001
MYLAN PHARMS INC	500MG	A090516	001	Jul 12, 2010
	750MG	A090516	002	Jul 12, 2010
NOSTRUM LABS INC	500MG	A090427	001	Dec 30, 2011
	750MG	A090427	002	Dec 30, 2011
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

RELAFEN

+	SMITHKLINE BEECHAM	500MG **	N019583	001	Dec 24, 1991
+		750MG **	N019583	002	Dec 24, 1991

NADOLOL

TABLET; ORAL

CORGARD

USWM	120MG	N018063	003	
	160MG	N018063	004	

NADOLOL

HERITAGE PHARMA	120MG	A074255	002	Jan 24, 1996
	160MG	A074255	003	Jan 24, 1996
NOVAST LABS	20MG	A210786	001	Jun 01, 2018
	40MG	A210786	002	Jun 01, 2018
	80MG	A210786	003	Jun 01, 2018
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	
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FOR SOLUTION; ORAL

UNIPEN

WYETH AYERST	EQ 250MG BASE/5ML	N050199	001	
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INJECTABLE; INJECTION

NAFCILLIN SODIUM

APOTHECON	EQ 500MG BASE/VIAL	A061984	001	
	EQ 1GM BASE/VIAL	A061984	002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NAFCILLIN SODIUMINJECTABLE; INJECTION
NAFCILLIN SODIUM

	EQ 2GM BASE/VIAL	A061984 003	
	EQ 4GM BASE/VIAL	A061984 005	
FRESENIUS	EQ 1GM BASE/VIAL	A206682 001	Dec 10, 2019
	EQ 2GM BASE/VIAL	A206682 002	Dec 10, 2019
MYLAN LABS LTD	EQ 1GM BASE/VIAL	A200002 001	Apr 07, 2014
	EQ 2GM BASE/VIAL	A200002 002	Apr 07, 2014
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	
	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

GEL; TOPICAL

NAFTIFINE HYDROCHLORIDE

TARO

2%

A208201 001 Apr 10, 2019

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

DR REDDYS

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

RISING

10MG/ML

A206506 001 Feb 06, 2019

10MG/ML

A207595 001 Jan 11, 2019

20MG/ML

A206506 002 Feb 06, 2019

20MG/ML

A207595 002 Jan 11, 2019

NUBAIN

+ PAR PHARM INC

10MG/ML **

N018024 001

+

20MG/ML **

N018024 002 May 27, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

HIKMA 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

HIKMA 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 A070254 001 Jan 07, 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

0.4MG/ML A070639 001 Sep 24, 1986

1MG/ML A072115 001 Apr 27, 1988

MARSAM PHARMS LLC 0.4MG/ML A071811 001 Jul 19, 1988

PAR STERILE PRODUCTS 0.4MG/ML A211286 001 Jan 17, 2020

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

VIRTUS PHARMS 0.4MG/ML A207846 001 Dec 17, 2018

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE HYDROCHLORIDE					
	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	Jun 14, 1982
	BRISTOL MYERS SQUIBB	0.4MG/ML	A071083	001	Jul 28, 1988
		1MG/ML	A071084	001	Jul 28, 1988
		1MG/ML	A071311	001	Jul 28, 1988
SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS					
EVZIO					
+	KALEO INC	0.4MG/0.4ML (0.4MG/0.4ML)	N205787	001	Apr 03, 2014
EVZIO (AUTOINJECTOR)					
+	KALEO INC	2MG/0.4ML (2MG/0.4ML)	N209862	001	Oct 19, 2016
SPRAY, METERED; NASAL					
NARCAN					
+	EMERGENT	2MG/SPRAY **	N208411	002	Jan 24, 2017

NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TARGINIQ					
+	PURDUE PHARMA LP	5MG; 10MG	N205777	001	Jul 23, 2014
+		10MG; 20MG	N205777	002	Jul 23, 2014
+		20MG; 40MG	N205777	003	Jul 23, 2014

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	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
	AM REGENT	200MG/ML	A091252	001	Aug 30, 2010
	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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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	
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NAFAZAIR

BAUSCH AND LOMB	0.1%	A040073	001	May 25, 1994
PHARMAFAIR	0.1%	A088101	001	Apr 15, 1983

NAPHAZOLINE HYDROCHLORIDE

AKORN	0.1%	A083590	001	
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NAPHCN FORTE

+ ALCON	0.1%	A080229	001	
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OPCON

BAUSCH AND LOMB	0.1%	A087506	001	
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VASOCON

NOVARTIS	0.1%	A080235	002	Mar 24, 1983
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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
	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

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

MYLAN	250MG	A074121	001	Dec 21, 1993
	375MG	A074121	002	Dec 21, 1993
	500MG	A074121	003	Dec 21, 1993

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NAPROXEN

TABLET;ORAL

NAPROXEN

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
PLIVA	375MG	A075337 001	May 26, 1999
	500MG	A075337 002	May 26, 1999

NAPROXEN SODIUM

TABLET;ORAL

ANAPROX

+ ATNAHS PHARMA US

EQ 250MG BASE **

N018164 001

NAPROXEN SODIUM

ABLE	EQ 250MG BASE	A076544 001	Aug 22, 2003
	EQ 500MG BASE	A076544 002	Aug 22, 2003
CONTRACT PHARMACAL	220MG	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	220MG	A074646 001	Jan 13, 1997
PLIVA	EQ 250MG BASE	A074242 001	Jun 20, 1996
	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 250MG BASE	A074198 001	Dec 21, 1993
	EQ 500MG BASE	A074142 002	Dec 21, 1993
	EQ 500MG BASE	A074198 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
YICHANG HUMANWELL	220MG	A212033 001	Aug 30, 2019

NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE

TABLET;ORAL

Treximet

+ CURRAX

60MG;EQ 10MG BASE

N021926 002 May 14, 2015

NARATRIPTAN HYDROCHLORIDE

TABLET;ORAL

NARATRIPTAN

ANI PHARMS	EQ 1MG BASE	A078751 001	Jul 07, 2010
	EQ 2.5MG BASE	A078751 002	Jul 07, 2010
APOTEX CORP	EQ 1MG BASE	A091373 001	Apr 22, 2011
	EQ 2.5MG BASE	A091373 002	Apr 22, 2011
CHARTWELL RX	EQ 1MG BASE	A090288 001	Jul 07, 2010
	EQ 2.5MG BASE	A090288 002	Jul 07, 2010
MYLAN PHARMS INC	EQ 1MG BASE	A202431 001	May 31, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NARATRIPTAN HYDROCHLORIDETABLET; ORAL
NARATRIPTAN

EQ 2.5MG BASE A202431 002 May 31, 2012

NATEGLINIDETABLET; ORAL
NATEGLINIDEALVOGEN 60MG A205055 001 Dec 11, 2015
120MG A205055 002 Dec 11, 2015
TEVA PHARMS 60MG A077467 001 Sep 09, 2009
120MG A077467 002 Sep 09, 2009

STARLIX

+ NOVARTIS 60MG ** N021204 001 Dec 22, 2000
+ 120MG ** N021204 002 Dec 22, 2000NEBIVOLOL 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
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
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, 2015NEBIVOLOL HYDROCHLORIDE; VALSARTAN

TABLET; ORAL

BYVALSON

+ ALLERGAN EQ 5MG BASE; 80MG N206302 001 Jun 03, 2016

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 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NEFAZODONE HYDROCHLORIDE

TABLET;ORAL

NEFAZODONE HYDROCHLORIDE

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
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NEOMYCIN SULFATE

SOLUTION;ORAL

MYCIFRADIN

PHARMACIA AND UPJOHN	EQ 87.5MG BASE/5ML	N050285 001	
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NEO-FRADIN

X GEN PHARMS	EQ 87.5MG BASE/5ML	A065010 001	May 23, 2002
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TABLET;ORAL

MYCIFRADIN

PHARMACIA AND UPJOHN	EQ 350MG BASE	A060520 001	
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NEOBIOTIC

PFIZER	EQ 350MG BASE	A060475 001	
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NEOMYCIN SULFATE

BRISTOL MYERS SQUIBB	500MG	A060365 001	
LANNETT	500MG	A060607 001	
LANNETT CO INC	500MG	A204435 001	Jun 10, 2016
LILLY	500MG	A060385 001	
NOSTRUM LABS INC	500MG	A065468 001	Mar 29, 2010
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
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OINTMENT;OPHTHALMIC

STATROL

ALCON	EQ 3.5MG BASE/GM;10,000 UNITS/GM	N050344 002	
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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	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

NESIRITIDE RECOMBINANT

FOR SOLUTION;INTRAVENOUS

NATRECOR

+ SCIOS LLC

1.5MG/VIAL **

N020920 001 Aug 10, 2001

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

NEVIRAPINE

TABLET;ORAL

NEVIRAPINE

APOTEX INC 200MG

A203021 001 May 22, 2012

MYLAN LABS 200MG

A078864 001 May 22, 2012

TECH ORGANIZED 200MG

A203176 001 May 22, 2012

TABLET, EXTENDED RELEASE;ORAL

NEVIRAPINE

APOTEX 400MG

A205258 001 Apr 03, 2014

CIPLA 400MG

A206448 001 Oct 15, 2015

MYLAN 100MG

A206271 001 Nov 09, 2015

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 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NIACIN

TABLET; ORAL

NIACIN

WOCKHARDT 500MG A081134 001 Apr 28, 1992

NICOLAR

+ SANOFI AVENTIS US 500MG A083823 001

TABLET, EXTENDED RELEASE; ORAL

NIACIN

JUBILANT GENERICS 500MG A209156 001 May 14, 2018

750MG A209156 002 May 14, 2018

1GM A209156 003 May 14, 2018

RISING 500MG A203742 001 Feb 22, 2019

750MG A203742 002 Feb 22, 2019

1GM A203742 003 Feb 22, 2019

YICHANG HUMANWELL 500MG A212017 001 Jun 10, 2019

750MG A212017 002 Jun 10, 2019

1GM A212017 003 Jun 10, 2019

NIASPAN

ABBVIE 375MG N020381 001 Jul 28, 1997

+ 500MG N020381 002 Jul 28, 1997

+ 750MG N020381 003 Jul 28, 1997

+ 1GM N020381 004 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 20MG ** N019488 001 Dec 21, 1988

+ 30MG ** N019488 002 Dec 21, 1988

NICARDIPINE HYDROCHLORIDE

ANI PHARMS 20MG A074439 001 Dec 10, 1996

20MG A074540 001 Oct 28, 1996

30MG A074439 002 Dec 10, 1996

30MG A074540 002 Oct 28, 1996

MYLAN 20MG A074642 001 Jul 18, 1996

30MG A074642 002 Jul 18, 1996

CAPSULE, EXTENDED RELEASE; ORAL

CARDENE SR

+ CHIESI 30MG ** N020005 001 Feb 21, 1992

+ 45MG ** N020005 002 Feb 21, 1992

+ 60MG ** N020005 003 Feb 21, 1992

INJECTABLE; INJECTION

CARDENE

+ CHIESI 25MG/10ML (2.5MG/ML) ** N019734 001 Jan 30, 1992

NICARDIPINE HYDROCHLORIDE

NAVINTA LLC 25MG/10ML (2.5MG/ML) A090125 001 Nov 17, 2009

RK PHARMA 25MG/10ML (2.5MG/ML) A090664 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 40MG/200ML (0.2MG/ML) N019734 005 Nov 07, 2008

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NICOTINE POLACRILEX

GUM, CHEWING;BUCCAL

NICOTINE POLACRILEX

IVAX SUB TEVA PHARMS EQ 2MG BASE
EQ 4MG BASEA076880 001 Feb 18, 2009
A077850 001 Feb 18, 2009

THRIVE

GLAXOSMITHKLINE CONS EQ 2MG BASE
EQ 4MG BASEA077658 001 Jun 19, 2007
A077656 001 Jun 19, 2007NIFEDIPINE

CAPSULE;ORAL

ADALAT

BAYER PHARMS 10MG
20MGN019478 001 Nov 27, 1985
N019478 002 Sep 17, 1986

NIFEDIPINE

CHASE LABS NJ 10MG
20MGA072409 001 Jul 04, 1990
A073421 001 Jun 19, 1991

TEVA 10MG

A072651 001 Feb 19, 1992

PROCARDIA

+ PFIZER 20MG **

N018482 002 Jul 24, 1986

TABLET, EXTENDED RELEASE;ORAL

ADALAT CC

+ NORWICH 30MG **

N020198 001 Apr 21, 1993

+ 60MG **

N020198 002 Apr 21, 1993

+ 90MG **

N020198 003 Apr 21, 1993

AFEDITAB CR

WATSON LABS 60MG

A075659 001 Oct 26, 2001

WATSON LABS TEVA 30MG

A075128 001 Mar 10, 2000

NIFEDIPINE

AUROBINDO PHARMA USA 30MG

A090649 001 Jun 21, 2010

60MG

A090649 002 Jun 21, 2010

90MG

A090649 003 Jun 21, 2010

MARTEC USA LLC 90MG

A075414 003 Mar 23, 2004

MYLAN 30MG

A075108 001 Dec 17, 1999

30MG

A201071 001 Dec 03, 2010

60MG

A201071 002 Dec 03, 2010

90MG

A201071 003 Dec 03, 2010

MYLAN LABS LTD 30MG

A090602 001 Sep 13, 2010

60MG

A090602 002 Sep 13, 2010

90MG

A090602 003 Sep 13, 2010

PAR PHARM 30MG

A077899 001 Dec 13, 2006

60MG

A077899 002 Dec 13, 2006

90MG

A077899 003 May 25, 2012

NILUTAMIDE

TABLET;ORAL

NILANDRON

CONCORDIA 50MG

N020169 001 Sep 19, 1996

NIMODIPINE

CAPSULE;ORAL

NIMODIPINE

SOFGEN PHARMS 30MG

A201832 001 Jul 24, 2015

SUN PHARM INDS INC 30MG

A077067 001 Apr 17, 2007

NIMOTOP

+ BAYER PHARMS 30MG **

N018869 001 Dec 28, 1988

SOLUTION;ORAL

NYMALIZE

+ ARBOR PHARMS LLC 3MG/ML **

N203340 001 May 10, 2013

NISOLDIPINE

TABLET, EXTENDED RELEASE;ORAL

SULAR

+ COVIS 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NITRIC OXIDE

GAS; INHALATION

INOMAX

+ MALLINCKRODT HOSP 100PPM ** N020845 002 Dec 23, 1999

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

AUROBINDO PHARMA USA 50MG A074967 001 Jul 09, 1997

100MG A074967 002 Jul 09, 1997

MYLAN 100MG A077025 001 Aug 18, 2004

PHARMGEN 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)

AUROBINDO PHARMA USA 75MG;25MG A076648 001 Mar 22, 2004

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NITROFURAZONE

OINTMENT; TOPICAL

NITROFURAZONE

WENDT 0.2% A086766 001

POWDER; TOPICAL

FURACIN

SHIRE 0.2% A083791 001

SOLUTION; TOPICAL

NITROFURAZONE

PERRIGO NEW YORK 0.2% A085130 001

WENDT 0.2% A087081 001

NITROGLYCERIN

AEROSOL; SUBLINGUAL

NITROLINGUAL

POHL BOSKAMP 0.4MG/SPRAY N018705 001 Oct 31, 1985

FILM, EXTENDED RELEASE; TRANSDERMAL

MINITRAN

BAUSCH 0.4MG/HR A089773 001 Aug 30, 1996

VALEANT PHARMS 0.1MG/HR A089771 001 Aug 30, 1996

VALEANT PHARMS 0.6MG/HR A089774 001 Aug 30, 1996

VALEANT PHARMS NORTH 0.2MG/HR A089772 001 Aug 30, 1996

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NITROGLYCERIN

POWDER; SUBLINGUAL

GONITRO

+ POHL BOSKAMP 0.4MG/PACKET N208424 001 Jun 08, 2016

TABLET; SUBLINGUAL

NITROGLYCERIN

ACTAVIS LABS FL INC	0.3MG	A203693 001	Oct 16, 2017
	0.4MG	A203693 002	Oct 16, 2017
	0.6MG	A203693 003	Oct 16, 2017
SIGMAPHARM LABS LLC	0.3MG	A207745 001	May 07, 2018
	0.4MG	A207745 002	May 07, 2018
	0.6MG	A207745 003	May 07, 2018

NIZATIDINE

CAPSULE; ORAL

AXID

SMITHKLINE BEECHAM	150MG	N019508 001	Apr 12, 1988
	300MG	N019508 002	Apr 12, 1988

NIZATIDINE

ANI PHARMS	150MG	A075461 001	Jul 08, 2002
	150MG	A075668 001	Sep 12, 2002
	300MG	A075461 002	Jul 08, 2002
	300MG	A075668 002	Sep 12, 2002
APOTEX INC	150MG	A076383 001	Jan 23, 2003
	300MG	A076383 002	Jan 23, 2003
MYLAN PHARMS INC	150MG	A075806 001	Jul 05, 2002
	150MG	A075934 001	Jul 09, 2002
	300MG	A075806 002	Jul 05, 2002
	300MG	A075934 002	Jul 09, 2002

SOLUTION; ORAL

AXID

+ BRAINTREE 15MG/ML ** N021494 001 May 25, 2004

NONOXYNOL-9

AEROSOL; VAGINAL

DELFIN

PERSONAL PRODS 12.5% N014349 002

SPONGE; VAGINAL

TODAY

+ MAYER LABS INC 1GM N018683 001 Apr 01, 1983

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

TABLET; ORAL-28

MICRONOR

+ JANSSEN PHARMS 0.35MG N016954 001

NORETHINDRONE ACETATE

TABLET; ORAL

AYGESTIN

+ DURAMED RES 5MG ** N018405 001 Apr 21, 1982

NORETHINDRONE ACETATE

AUROBINDO PHARMA LTD 5MG A204236 001 Jan 08, 2016

NORLUTATE

PARKE DAVIS 5MG N012184 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

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

ZYDUS

EQ 10MG BASE

A213441 001 Feb 24, 2021

EQ 25MG BASE

A213441 002 Feb 24, 2021

EQ 50MG BASE

A213441 003 Feb 24, 2021

EQ 75MG BASE

A213441 004 Feb 24, 2021

SOLUTION;ORAL

AVENTYL

+ RANBAXY

EQ 10MG BASE/5ML **

N014685 001

PAMELOR

SPECGX LLC

EQ 10MG BASE/5ML

N018012 001

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

STRIDES PHARMA

100,000 UNITS/GM

A065315 001 May 31, 2006

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NYSTATIN

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				
	+ STRIDES PHARMA	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
	COSETTE	100,000 UNITS/ML	A062776 001	Dec 17, 1987
	G AND W LABS INC	100,000 UNITS/ML	A062349 001	Jul 14, 1982
	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
	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				
	CHARTWELL RX	500,000 UNITS	A062524 001	Nov 26, 1985
	QUANTUM PHARMICS	500,000 UNITS	A062525 001	Oct 29, 1984
	SANDOZ	500,000 UNITS	A062065 001	
	WATSON LABS	500,000 UNITS	A062402 001	Dec 16, 1982
TABLET;VAGINAL				
KOROSTATIN				
	HOLLAND RANTOS	100,000 UNITS	A061718 001	
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	
<u>NYSTATIN; TRIAMCINOLONE ACETONIDE</u>				
CREAM;TOPICAL				
MYCO-TRIACET 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	100,000 UNITS/GM;0.1% **	A062606 001	May 15, 1985
MYTREX F				
	SAVAGE LABS	100,000 UNITS/GM;0.1%	A062597 001	Oct 08, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

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
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OINTMENT; TOPICAL

MYCO-TRIACET II

TEVA	100,000 UNITS/GM;0.1%	A062045	002	Nov 26, 1985
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MYCOLOG-II

MYLAN	100,000 UNITS/GM;0.1% **	A060572	001	Jun 28, 1985
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MYTREX F

SAVAGE LABS	100,000 UNITS/GM;0.1%	A062601	001	Oct 09, 1985
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NYSTATIN AND TRIAMCINOLONE ACETONIDE

CROWN LABS INC	100,000 UNITS/GM;0.1%	A207731	001	Dec 26, 2017
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
VITRUVIAS THERAP	100,000 UNITS/GM;0.1%	A207316	001	Nov 18, 2019

NYSTATIN-TRIAMCINOLONE ACETONIDE

PHARMADERM	100,000 UNITS/GM;0.1%	A062603	001	Oct 09, 1985
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OCTREOTIDE ACETATE

INJECTABLE; INJECTION

OCTREOTIDE ACETATE

HERITAGE PHARMS INC	EQ 0.05MG BASE/ML	A204669	001	Dec 27, 2018
	EQ 0.1MG BASE/ML	A204669	002	Dec 27, 2018
	EQ 0.2MG BASE/ML	A203765	001	Sep 07, 2018
	EQ 0.5MG BASE/ML	A204669	003	Dec 27, 2018
	EQ 1MG BASE/ML	A203765	002	Sep 07, 2018
SUN PHARM INDS	EQ 0.05MG BASE/ML	A077329	001	Mar 04, 2008
	EQ 0.05MG BASE/ML	A077372	001	Aug 14, 2007
	EQ 0.1MG BASE/ML	A077329	002	Mar 04, 2008
	EQ 0.1MG BASE/ML	A077372	002	Aug 14, 2007
	EQ 0.2MG BASE/ML	A077330	001	Mar 04, 2008
	EQ 0.2MG BASE/ML	A077373	001	Aug 14, 2007
	EQ 0.5MG BASE/ML	A077329	003	Mar 04, 2008
	EQ 0.5MG BASE/ML	A077372	003	Aug 14, 2007
	EQ 1MG BASE/ML	A077331	001	Mar 04, 2008
	EQ 1MG BASE/ML	A077373	002	Aug 14, 2007
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

SANDOSTATIN

+	NOVARTIS	EQ 0.2MG BASE/ML **	N019667	004	Jun 12, 1991
+		EQ 1MG BASE/ML **	N019667	005	Jun 12, 1991

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
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FLOXIN IN DEXTROSE 5% IN PLASTIC CONTAINER

ORTHO MCNEIL PHARM	4MG/ML	N020087	004	Mar 31, 1992
	400MG/100ML	N020087	005	Mar 31, 1992

OFLOXACIN

BEDFORD	40MG/ML	A075762	001	Jan 16, 2002
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SOLUTION/DROPS; OPHTHALMIC

OFLOXACIN

ALVOGEN	0.3%	A076830	001	Aug 31, 2004
SANDOZ	0.3%	A076848	001	Nov 25, 2008
SANDOZ INC	0.3%	A076231	001	May 14, 2004

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OFLOXACIN

SOLUTION/DROPS;OTIC

FLOXIN OTIC

+ DAIICHI	0.3% **	N020799 001	Dec 16, 1997
OFLOXACIN			
ALVOGEN	0.3%	A090395 001	Aug 11, 2009
SANDOZ INC	0.3%	A078222 001	Mar 17, 2008

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
HIKMA	2.5MG	A204866 001	Jun 16, 2017
	5MG	A204866 002	Jun 16, 2017
	7.5MG	A204866 003	Jun 16, 2017
	10MG	A204866 004	Jun 16, 2017
	15MG	A204866 005	Jun 16, 2017
	20MG	A204866 006	Jun 16, 2017
HISUN PHARM HANGZHOU	2.5MG	A206924 001	Dec 31, 2020
	5MG	A206924 002	Dec 31, 2020
	7.5MG	A206924 003	Dec 31, 2020
	10MG	A206924 004	Dec 31, 2020
	15MG	A206924 005	Dec 31, 2020
	20MG	A206924 006	Dec 31, 2020
IVAX PHARMS INC	20MG	A077301 001	Apr 29, 2015
JIANGSU HANSOH PHARM	2.5MG	A209399 001	Sep 24, 2018
	5MG	A209399 002	Sep 24, 2018
	10MG	A209399 003	Sep 24, 2018
MYLAN	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
SUNSHINE	2.5MG	A206238 001	Nov 19, 2018
	5MG	A206238 002	Nov 19, 2018
	7.5MG	A206238 003	Nov 19, 2018
	10MG	A206238 004	Nov 19, 2018
	15MG	A206238 005	Nov 19, 2018
	20MG	A206238 006	Nov 19, 2018
TEVA PHARMS	2.5MG	A076000 001	Oct 24, 2011
	5MG	A076000 002	Oct 24, 2011
	7.5MG	A076000 003	Oct 24, 2011
	10MG	A076000 004	Oct 24, 2011
	15MG	A076000 005	Oct 24, 2011
TORRENT PHARMS LTD	2.5MG	A091434 001	Apr 23, 2012
	5MG	A091434 002	Apr 23, 2012
	7.5MG	A091434 003	Apr 23, 2012
	10MG	A091434 004	Apr 23, 2012
	15MG	A091434 005	Apr 23, 2012
	20MG	A091434 006	Apr 23, 2012

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OLANZAPINETABLET, 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
HISUN PHARM HANGZHOU	5MG	A206892 001	Dec 31, 2020
	10MG	A206892 002	Dec 31, 2020
	15MG	A206892 003	Dec 31, 2020
	20MG	A206892 004	Dec 31, 2020

OLAPARIB

CAPSULE;ORAL

LYNPARZA

+ ASTRAZENECA	50MG	N206162 001	Dec 19, 2014
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OLMESARTAN MEDOXOMIL

TABLET;ORAL

OLMESARTAN MEDOXOMIL

INVENTIA	5MG	A208659 001	May 18, 2020
	20MG	A208659 002	May 18, 2020
	40MG	A208659 003	May 18, 2020
JUBILANT GENERICS	5MG	A205482 001	Apr 24, 2017
	20MG	A205482 002	Apr 24, 2017
	40MG	A205482 003	Apr 24, 2017
LUPIN LTD	5MG	A206631 001	Apr 27, 2017
	20MG	A206631 002	Apr 27, 2017
	40MG	A206631 003	Apr 27, 2017
RISING PHARMA	5MG	A078276 001	Oct 26, 2016
	20MG	A078276 002	Oct 26, 2016
	40MG	A078276 003	Oct 26, 2016
TEVA PHARMS USA	5MG	A091079 001	Apr 24, 2017
	20MG	A091079 002	Apr 24, 2017
	40MG	A091079 003	Apr 24, 2017

OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

OLOPATADINE HYDROCHLORIDE

BAUSCH	EQ 0.1% BASE	A206046 001	Jul 26, 2017
CIPLA	EQ 0.2% BASE	A206087 001	Dec 05, 2017
FDC LTD	EQ 0.1% BASE	A209282 001	Sep 26, 2019
MYLAN	EQ 0.1% BASE	A204392 001	Mar 21, 2018
WATSON LABS INC	EQ 0.7% BASE	A208637 001	Feb 19, 2020
WOCKHARDT LTD	EQ 0.1% BASE	A200810 001	Jun 28, 2017
ZAMBON SPA	EQ 0.1% BASE	A204706 001	Dec 07, 2015

OMBITASVIR; PARITAPREVIR; RITONAVIR

TABLET;ORAL

TECHNIVIE

+ ABBVIE INC	12.5MG;75MG;50MG **	N207931 001	Jul 24, 2015
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OMEGA-3-ACID ETHYL ESTERS

CAPSULE;ORAL

OMEGA-3-ACID ETHYL ESTERS

STRIDES PHARMA	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A091018 001	Jun 24, 2014
ZYDUS	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A210107 001	Jun 14, 2019

OMEGA-3-ACID ETHYL ESTERS TYPE A

CAPSULE;ORAL

OMTRYG

+ OSMOTICA PHARM US	1.2GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	N204977 001	Apr 23, 2014
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OMEGA-3-CARBOXYLIC ACIDS

CAPSULE;ORAL

EPANOVA

+ ASTRAZENECA	1GM CONTAINS AT LEAST 850MG OF POLYUNSATURATED FATTY ACIDS	N205060 001	May 05, 2014
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

OMEPRAZOLE

LUPIN LTD	40MG	A202384	001	Aug 25, 2015
MYLAN	10MG	A075876	001	May 29, 2003
	10MG	A205070	001	Jun 29, 2018
	20MG	A075876	002	May 29, 2003
	20MG	A205070	002	Jun 29, 2018
	40MG	A075876	003	Jan 21, 2009
	40MG	A205070	003	Jun 29, 2018
TEVA PHARMS USA	40MG	A204661	002	Jun 13, 2017

PRILOSEC

+ ASTRAZENECA	10MG **	N019810	003	Oct 05, 1995
+	20MG **	N019810	001	Sep 14, 1989
+	40MG **	N019810	002	Jan 15, 1998

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

STRIDES PHARMA	20MG;1.1GM	A078966	001	May 25, 2010
	20MG;1.1GM	A201946	001	Jul 15, 2016
	40MG;1.1GM	A078966	002	May 25, 2010
UNICORN	20MG;1.1GM	A204137	001	Jul 15, 2016

ONDANSETRON

FILM;ORAL

ZUPLENZ

+ AQUESTIVE	4MG	N022524	001	Jul 02, 2010
+	8MG	N022524	002	Jul 02, 2010

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

BARR	4MG	A076693	001	Jun 25, 2007
	8MG	A076693	002	Jun 25, 2007
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
TEVA	4MG	A076810	001	Jun 25, 2007
	8MG	A076810	002	Jun 25, 2007

ZOFRAN ODT

+ NOVARTIS	4MG	N020781	001	Jan 27, 1999
+	8MG	N020781	002	Jan 27, 1999

ONDANSETRON HYDROCHLORIDE

INJECTABLE;INJECTION

ONDANSETRON HYDROCHLORIDE

APOTEX INC	EQ 2MG BASE/ML	A077368	001	Dec 26, 2006
BAXTER HLTHCARE CORP	EQ 2MG BASE/ML	A078288	001	Feb 22, 2013
EMCURE PHARMS	EQ 2MG BASE/ML	A090424	001	Apr 16, 2010
HOSPIRA	EQ 2MG BASE/ML	A076695	001	Dec 26, 2006
	EQ 2MG BASE/ML	A077840	001	Jan 19, 2007
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
	EQ 2MG BASE/ML	A079039	001	Nov 18, 2008
MYLAN LABS LTD	EQ 2MG BASE/ML	A078257	001	Apr 23, 2008
PLIVA HRVATSKA DOO	EQ 2MG BASE/ML	A077544	001	Dec 26, 2006
RISING PHARMA	EQ 2MG BASE/ML	A204906	001	Jul 31, 2017
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
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ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

AM REGENT	EQ 2MG BASE/ML	A079032	001	Nov 18, 2008
APOTEX INC	EQ 2MG BASE/ML	A077343	001	Dec 26, 2006
EMCURE PHARMS LTD	EQ 2MG BASE/ML	A078945	001	Jan 03, 2013
HIKMA FARMACEUTICA	EQ 2MG BASE/ML	A076780	001	Dec 26, 2006
HOSPIRA	EQ 2MG BASE/ML	A076696	001	Dec 26, 2006
LUITPOLD	EQ 2MG BASE/ML	A077387	001	Dec 26, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

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
TEVA	EQ 2MG BASE/ML	A076759	001	Nov 22, 2006

ZOFRAN

+ NOVARTIS	EQ 2MG BASE/ML **	N020007	001	Jan 04, 1991
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ZOFRAN AND DEXTROSE IN PLASTIC CONTAINER

+ GLAXOSMITHKLINE	EQ 0.64MG BASE/ML **	N020403	001	Jan 31, 1995
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ZOFRAN PRESERVATIVE FREE

+ NOVARTIS	EQ 2MG BASE/ML **	N020007	003	Dec 10, 1993
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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
PLIVA HRVATSKA DOO	EQ 4MG BASE	A077112	001	Jun 25, 2007
	EQ 8MG BASE	A077112	002	Jun 25, 2007
	EQ 24MG BASE	A077112	003	Jun 25, 2007
RISING	EQ 4MG BASE	A076930	001	Jun 25, 2007
	EQ 8MG BASE	A076930	002	Jun 25, 2007
	EQ 24MG BASE	A076930	004	Jun 25, 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				
+ BAUSCH	100MG	N012157	001	
ORPHENADRINE CITRATE				
ASCOT	100MG	A088067	001	Apr 06, 1983
IMPAX PHARMS	100MG	A040368	001	Jun 23, 2000
SANDOZ	100MG	A085046	001	
WATSON LABS	100MG	A084303	001	

ORPHENADRINE HYDROCHLORIDE

TABLET; ORAL

DISIPAL				
3M	50MG	N010653	001	

OSELTAMIVIR PHOSPHATE

CAPSULE; ORAL

OSELTAMIVIR PHOSPHATE

RISING PHARMA	EQ 30MG BASE	A210157	001	Jan 21, 2021
	EQ 45MG BASE	A210157	002	Jan 21, 2021
	EQ 75MG BASE	A210157	003	Jan 21, 2021
SUNSHINE	EQ 30MG BASE	A212739	001	Mar 04, 2020
	EQ 45MG BASE	A212739	002	Mar 04, 2020
	EQ 75MG BASE	A212739	003	Mar 04, 2020

FOR SUSPENSION; ORAL

TAMIFLU

ROCHE	EQ 12MG BASE/ML	N021246	001	Dec 14, 2000
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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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXACILLIN SODIUM

CAPSULE; ORAL

OXACILLIN SODIUM

ANI PHARMS

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

HOSPIRA INC

EQ 1GM BASE/VIAL

A203950 001 Dec 11, 2015

EQ 2GM BASE/VIAL

A203950 002 Dec 11, 2015

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

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; INTRAVENOUS

ELOXATIN

+ SANOFI AVENTIS US

50MG/VIAL **

N021492 001 Aug 09, 2002

+ SANOFI AVENTIS US

100MG/VIAL **

N021492 002 Aug 09, 2002

+ SANOFI AVENTIS US

200MG/40ML (5MG/ML) **

N021759 003 Nov 17, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXALIPLATIN

INJECTABLE; INTRAVENOUS

OXALIPLATIN

ACCORD HLTHCARE	200MG/40ML (5MG/ML)	A207474	003	Mar 21, 2017
AM REGENT	50MG/10ML (5MG/ML)	A204378	001	May 12, 2017
	100MG/20ML (5MG/ML)	A204378	002	May 12, 2017
CIPLA	50MG/10ML (5MG/ML)	A208523	001	Feb 10, 2017
	100MG/20ML (5MG/ML)	A208523	002	Feb 10, 2017
FRESENIUS KABI ONCOL	50MG/VIAL	A078810	001	Aug 07, 2009
	100MG/VIAL	A078810	002	Aug 07, 2009
FRESENIUS KABI USA	200MG/40ML (5MG/ML)	A090030	003	Jan 31, 2017
GLAND	200MG/40ML (5MG/ML)	A207325	003	Oct 18, 2017
HOSPIRA INC	50MG/VIAL	A078815	001	Sep 30, 2009
	100MG/VIAL	A078815	002	Sep 30, 2009
MYLAN LABS LTD	50MG/VIAL	A200979	001	Aug 08, 2012
	100MG/VIAL	A200979	002	Aug 08, 2012
	200MG/40ML (5MG/ML)	A091358	003	Nov 14, 2017
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
SUN PHARM	50MG/VIAL	A078818	001	Aug 07, 2009
	50MG/10ML (5MG/ML)	A202922	001	Apr 08, 2014
	100MG/VIAL	A078818	002	Aug 07, 2009
	100MG/20ML (5MG/ML)	A202922	002	Apr 08, 2014
	200MG/40ML (5MG/ML)	A202922	003	Feb 15, 2019

OXAMNIQUINE

CAPSULE; ORAL

VANSIL

PFIZER	250MG	N018069	001	
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OXANDROLONE

TABLET; ORAL

OXANDRIN

+	GEMINI LABS LLC	2.5MG	N013718	001
+		10MG	N013718	002

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
BEXIMCO PHARMS USA	600MG	A075842	001	Apr 12, 2001
IVAX SUB TEVA PHARMS	600MG	A075846	001	May 13, 2002
MYLAN	600MG	A075851	001	Aug 17, 2001
MYLAN PHARMS INC	600MG	A075847	001	Feb 28, 2001
SANDOZ	600MG	A075850	001	Apr 27, 2001
SUN PHARM INDS INC	600MG	A075844	001	Jan 03, 2002
WATSON LABS	600MG	A075848	001	Feb 09, 2001

OXAPROZIN POTASSIUM

TABLET; ORAL

DAYPRO ALTA

PFIZER	600MG	N020776	001	Oct 17, 2002
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OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

AM THERAP	10MG	A071955	001	Mar 03, 1988
	15MG	A071956	001	Mar 03, 1988
	30MG	A071957	001	Mar 03, 1988
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

WATSON LABS	15MG	A072953	001	Sep 28, 1990
	30MG	A072954	001	Sep 28, 1990
WATSON LABS TEVA	10MG	A072952	001	Sep 28, 1990

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	
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OXCARBAZEPINE

TABLET; ORAL

OXCARBAZEPINE

ANI PHARMS	150MG	A078005	001	Dec 11, 2007
	300MG	A078005	002	Dec 11, 2007
	600MG	A078005	003	Dec 11, 2007
HIKMA	150MG	A077795	001	Oct 09, 2007
	300MG	A077795	002	Oct 09, 2007
	600MG	A077795	003	Oct 09, 2007
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	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXYBUTYNYNIN

FILM, EXTENDED RELEASE;TRANSDERMAL

OXYBUTYNYNIN

BARR LABS DIV TEVA 3.9MG/24HR

A090526 001 Mar 04, 2014

GEL, METERED;TRANSDERMAL

GELNIQUE 3%

+ ALLERGAN 3%

N202513 001 Dec 07, 2011

OXYBUTYNYNIN CHLORIDE

GEL;TRANSDERMAL

OXYBUTYNYNIN CHLORIDE

STRIDES PHARMA 10% (100MG/PACKET)

A207329 001 May 31, 2018

SYRUP;ORAL

DITROPAN

+ ORTHO MCNEIL JANSSEN 5MG/5ML **

N018211 001

OXYBUTYNYNIN CHLORIDE

ANDA REPOSITORY 5MG/5ML

A075039 001 Jan 29, 1999

LANNETT CO INC 5MG/5ML

A076682 001 Dec 28, 2004

PHARM ASSOC 5MG/5ML

A074997 001 Oct 15, 1997

TABLET;ORAL

DITROPAN

+ JANSSEN PHARMS 5MG **

N017577 001

OXYBUTYNYNIN CHLORIDE

QUANTUM PHARMICS 5MG

A072296 001 Dec 08, 1988

STRIDES PHARMA 5MG

A208165 001 Dec 17, 2020

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

OXYBUTYNYNIN CHLORIDE

IMPAX PHARMS 5MG

A076745 002 May 09, 2007

10MG

A076745 003 May 09, 2007

15MG

A076745 001 Nov 09, 2006

MYLAN 5MG

A076702 001 Nov 09, 2006

MYLAN PHARMS INC 10MG

A076644 001 Nov 09, 2006

15MG

A076644 002 May 10, 2007

OXYCODONE HYDROCHLORIDE

CAPSULE;ORAL

OXYCODONE HYDROCHLORIDE

LANNETT CO INC 5MG

A203823 001 Aug 01, 2014

SOLUTION;ORAL

OXYCODONE HYDROCHLORIDE

ANI PHARMS 100MG/5ML

A203447 001 Aug 30, 2017

AUROLIFE PHARMA LLC 5MG/5ML

A212429 001 Jan 27, 2020

100MG/5ML

A212429 002 Jan 27, 2020

HIKMA 100MG/5ML

A203208 001 Jul 12, 2013

LANNETT CO INC 100MG/5ML

A204085 001 Sep 09, 2014

RHODES PHARMS 100MG/5ML

A205853 001 Apr 29, 2020

TABLET;ORAL

OXYCODONE HYDROCHLORIDE

ACTAVIS ELIZABETH 5MG

A076636 003 Apr 07, 2015

15MG

A076636 001 Feb 06, 2004

30MG

A076636 002 Feb 06, 2004

MAYNE PHARMA INC 5MG

A091313 001 Feb 18, 2011

10MG

A091313 004 Apr 29, 2016

15MG

A091313 002 Feb 18, 2011

20MG

A091313 005 Apr 29, 2016

30MG

A091313 003 Feb 18, 2011

NESHER PHARMS 5MG

A077290 001 Dec 08, 2005

10MG

A077290 002 Dec 08, 2005

15MG

A077290 003 Dec 08, 2005

20MG

A077290 004 Dec 08, 2005

30MG

A077290 005 Dec 08, 2005

STRIDES PHARMA 5MG

A077712 003 Mar 02, 2009

10MG

A077712 004 Apr 13, 2015

15MG

A077712 001 Jan 31, 2007

20MG

A077712 005 Apr 13, 2015

30MG

A077712 002 Jan 31, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXYCODONE HYDROCHLORIDE

TABLET; ORAL

ROXYBOND

OHEMO LIFE	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

CREAM; TOPICAL

OXYMETAZOLINE HYDROCHLORIDE

TARO PHARMS	1%	A213584 001	Oct 04, 2021
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SOLUTION/DROPS; OPHTHALMIC

OCUCLEAR

BAYER HEALTHCARE LLC	0.025%	N018471 001	May 30, 1986
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OXYMETHOLONE

TABLET; ORAL

ANADROL-50

+ MYLAN SPECIALITY LP	50MG	N016848 001	
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OXYMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION

OPANA

+ ENDO PHARMS	1MG/ML	N011707 002	
	1.5MG/ML	N011707 001	

SUPPOSITORY; RECTAL

NUMORPHAN

ENDO PHARMS	5MG	N011738 004	
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TABLET; ORAL

OPANA

+ ENDO PHARMS	5MG	N021611 001	Jun 22, 2006
	10MG	N021611 002	Jun 22, 2006

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

ACTAVIS ELIZABETH

	5MG	A079046 003	Jul 11, 2013
	7.5MG	A079046 001	Dec 13, 2010
	10MG	A079046 004	Jul 11, 2013
	15MG	A079046 002	Dec 13, 2010
	20MG	A079046 005	Jul 11, 2013
	30MG	A079046 006	Jul 11, 2013
	40MG	A079046 007	Jul 11, 2013
HIKMA	5MG	A200822 002	Jul 15, 2013
	7.5MG	A200822 003	Jul 15, 2013
	10MG	A200822 004	Jul 15, 2013
	15MG	A200822 005	Jul 15, 2013
	20MG	A200822 006	Jul 15, 2013
	30MG	A200822 007	Jul 15, 2013
	40MG	A200822 001	Jul 15, 2013

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXYMORPHONE HYDROCHLORIDETABLET, EXTENDED RELEASE;ORAL
OXYMORPHONE HYDROCHLORIDE

	20MG	A200792 005	Oct 24, 2014
	30MG	A200792 006	Oct 24, 2014
	40MG	A200792 007	Oct 24, 2014
SPECGX LLC	5MG	A202946 001	Jun 27, 2014
	7.5MG	A202946 002	Jun 27, 2014
	10MG	A202946 003	Jun 27, 2014
	15MG	A202946 004	Jun 27, 2014
	20MG	A202946 005	Jun 27, 2014
	30MG	A202946 006	Jun 27, 2014
	40MG	A202946 007	Jun 27, 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

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXYTETRACYCLINE HYDROCHLORIDE; POLYMYXIN B SULFATE

TABLET;VAGINAL

TERRAMYCIN-POLYMYXIN

PFIZER

EQ 100MG BASE;100,000 UNITS

A061009 001

OXYTOCIN

INJECTABLE;INJECTION

OXYTOCIN

DR REDDYS

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

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

MYLAN LABS LTD

6MG/ML

A091540 001 Sep 29, 2011

PLIVA LACHEMA

6MG/ML

A077413 001 Mar 12, 2008

SANDOZ INC

6MG/ML

A078167 001 Dec 26, 2007

TEVA PHARMS USA

6MG/ML

A075297 001 Jan 25, 2002

TAXOL

+ HQ SPCLT PHARMA

6MG/ML

N020262 001 Dec 29, 1992

PALIPERIDONE

TABLET, EXTENDED RELEASE;ORAL

INVEGA

+ JANSSEN PHARMS

12MG **

N021999 004 Dec 19, 2006

PALIPERIDONE PALMITATE

SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

PALIPERIDONE PALMITATE

TEVA PHARMS USA

39MG/0.25ML (39MG/0.25ML)

A211149 001 Jul 06, 2021

78MG/0.5ML (78MG/0.5ML)

A211149 002 Jul 06, 2021

117MG/0.75ML (117MG/0.75ML)

A211149 003 Jul 06, 2021

156MG/ML (156MG/ML)

A211149 004 Jul 06, 2021

234MG/1.5ML (156MG/ML)

A211149 005 Jul 06, 2021

PALONOSETRON HYDROCHLORIDE

CAPSULE;ORAL

ALOXI

+ HELSINN HLTHCARE

EQ 0.5MG BASE **

N022233 001 Aug 22, 2008

INJECTABLE;INTRAVENOUS

ALOXI

+ HELSINN HLTHCARE

EQ 0.075MG BASE/1.5ML (EQ 0.05MG
BASE/ML)

N021372 002 Feb 29, 2008

+

EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)

N021372 001 Jul 25, 2003

PALONOSETRON HYDROCHLORIDE

ACCORD HLTHCARE

EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)

A204615 001 Mar 15, 2021

DR REDDYS LABS LTD

EQ 0.075MG BASE/1.5ML (EQ 0.05MG
BASE/ML)

A201533 001 Apr 21, 2016

HOSPIRA INC

EQ 0.075MG BASE/1.5ML (EQ 0.05MG
BASE/ML)

A207005 002 Sep 19, 2018

QILU PHARM HAINAN

EQ 0.075MG BASE/1.5ML (EQ 0.05MG
BASE/ML)

A205648 002 Sep 19, 2018

TEVA PHARMS USA

EQ 0.075MG BASE/1.5ML (EQ 0.05MG
BASE/ML)

A090713 002 Mar 23, 2018

SOLUTION;INTRAVENOUS

PALONOSETRON HYDROCHLORIDE

DR REDDYS LABS LTD

EQ 0.075MG BASE/1.5ML (EQ 0.05MG
BASE/ML)

N203050 001 Mar 01, 2016

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PALONOSETRON HYDROCHLORIDE

SOLUTION; INTRAVENOUS

PALONOSETRON HYDROCHLORIDE

	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	N203050 002	Mar 01, 2016
+	FRESENIUS KABI USA	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	N208109 001
			Nov 21, 2017

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

AREZIA

+	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

AM REGENT

	30MG/10ML (3MG/ML)	A078942 001	Jul 25, 2008
	90MG/10ML (9MG/ML)	A078942 002	Jul 25, 2008

FRESENIUS KABI USA

	30MG/VIAL	A075773 001	May 06, 2002
	30MG/10ML (3MG/ML)	A076207 001	May 17, 2002
	90MG/VIAL	A075773 002	May 06, 2002
	90MG/10ML (9MG/ML)	A076207 002	May 17, 2002

MN PHARMS

	30MG/VIAL	A078300 001	Mar 10, 2009
	90MG/VIAL	A078300 002	Mar 10, 2009

SUN PHARMA GLOBAL

	30MG/VIAL	A077703 001	Dec 24, 2008
	90MG/VIAL	A077703 002	Dec 24, 2008

TEVA PHARMS USA

	30MG/10ML (3MG/ML)	A076153 001	Mar 27, 2002
	90MG/10ML (9MG/ML)	A076153 002	Mar 27, 2002

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
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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

PANOBINOSTAT LACTATE

CAPSULE; ORAL

FARYDAK

+	SECURA	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

INJECTABLE; INTRAVENOUS

PANTOPRAZOLE SODIUM

MYLAN LABS LTD

	EQ 40MG BASE/VIAL	A208580 001	May 04, 2018
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TABLET, DELAYED RELEASE; ORAL

PANTOPRAZOLE SODIUM

JUBILANT GENERICS

	EQ 20MG BASE	A090901 001	Aug 30, 2011
	EQ 40MG BASE	A090901 002	Aug 30, 2011

MACLEODS PHARMS LTD

	EQ 20MG BASE	A200821 001	Feb 16, 2012
	EQ 40MG BASE	A200821 002	Feb 16, 2012

SUN PHARM

	EQ 20MG BASE	A077058 001	Sep 10, 2007
	EQ 40MG BASE	A077058 002	Sep 10, 2007

SUN PHARM INDS LTD

	EQ 20MG BASE	A200794 001	May 02, 2012
	EQ 40MG BASE	A200794 002	May 02, 2012

TEVA

	EQ 20MG BASE	A077056 001	Aug 02, 2007
	EQ 40MG BASE	A077056 002	Aug 02, 2007

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

PARICALCITOL

LOTUS PHARM CO LTD

1MCG

A206710 001 Feb 24, 2016

2MCG

A206710 002 Feb 24, 2016

4MCG

A206710 003 Feb 24, 2016

ZEMPLAR

+ ABBVIE

4MCG **

N021606 003 May 26, 2005

PAROMOMYCIN SULFATE

CAPSULE; ORAL

HUMATIN

KING PFIZER

EQ 250MG BASE

A062310 001

PARKE DALE

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

JUBILANT GENERICS

EQ 10MG BASE

A205528 001 Nov 27, 2015

EQ 20MG BASE

A205528 002 Nov 27, 2015

EQ 30MG BASE

A205528 003 Nov 27, 2015

EQ 40MG BASE

A205528 004 Nov 27, 2015

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

SUN PHARM INDS INC

EQ 10MG BASE

A078194 001 Jun 29, 2007

EQ 20MG BASE

A078194 002 Jun 29, 2007

EQ 30MG BASE

A078194 003 Jun 29, 2007

EQ 40MG BASE

A078194 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PAROXETINE HYDROCHLORIDE

TABLET; ORAL

PAROXETINE HYDROCHLORIDE

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
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PAZOPANIB HYDROCHLORIDE

TABLET; ORAL

VOTRIENT

+ NOVARTIS	EQ 400MG BASE **	N022465 002	Oct 19, 2009
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PEGAPTANIB SODIUM

INJECTABLE; INTRAVITREAL

MACUGEN

+ VALEANT PHARMS LLC	EQ 0.3MG ACID/0.09ML	N021756 001	Dec 17, 2004
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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

PEMETREXED

SOLUTION; INTRAVENOUS

PEMETREXED

+ ACTAVIS LLC	100MG/4ML (25MG/ML)	N208419 001	Aug 21, 2020
	500MG/20ML (25MG/ML)	N208419 002	Aug 21, 2020
	1GM/40ML (25MG/ML)	N208419 003	Aug 21, 2020

PEMIROLAST POTASSIUM

SOLUTION/DROPS; OPHTHALMIC

ALAMAST

SANTEN	0.1%	N021079 001	Sep 24, 1999
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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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PEMOLINE

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

VALEANT PHARMS INTL 125MG N019853 002

TABLET;ORAL

PENICILLAMINE

TEVA PHARMS USA 250MG A211497 001 Feb 13, 2020

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLIN G POTASSIUM

INJECTABLE; INJECTION

PENICILLIN G POTASSIUM

	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
	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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
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PENICILLIN V

FOR SUSPENSION; ORAL

V-CILLIN

LILLY	125MG/0.6ML	A060002	001
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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
	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

BELCHER PHARMS	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
STRIDES PHARMA	EQ 125MG BASE/5ML	A062981	001 Feb 10, 1989
	EQ 250MG BASE/5ML	A062981	002 Feb 10, 1989

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLIN V POTASSIUM

TABLET; ORAL

PENAPAR-VK

PARKE DAVIS	EQ 250MG BASE	A062001 001
	EQ 500MG BASE	A062001 002

PENICILLIN V POTASSIUM

BELCHER PHARMS	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

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
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PENTAMIDINE ISETHIONATE

FOR SOLUTION; INHALATION

NEBUPENT

FRESENIUS KABI USA	600MG/VIAL	N019887 002	Mar 22, 1996
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INJECTABLE; INJECTION

PENTACARINAT

ARMOUR PHARM	300MG/VIAL	A073447 001	Apr 28, 1994
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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
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PENTAZOCINE LACTATE

INJECTABLE; INJECTION

TALWIN

+ HOSPIRA	EQ 30MG BASE/ML	N016194 001
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PENTETATE CALCIUM TRISODIUM

SOLUTION; INHALATION, INTRAVENOUS

PENTETATE CALCIUM TRISODIUM

+ HAMELN	EQ 1GM BASE/5ML (EQ 200MG BASE/ML) **	N021749 001	Aug 11, 2004
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENTETATE CALCIUM TRISODIUM YB-169

INJECTABLE; INJECTION

YTTERBIUM YB 169 DTPA

3M

2mCi/ML

N017518 001

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

ELIXIR; ORAL

NEMBUTAL

AKORN

18.2MG/5ML

A083244 001

PENTOBARBITAL SODIUM

CAPSULE; ORAL

NEMBUTAL SODIUM

AKORN

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

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

AKORN

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

PENTOSTATIN

INJECTABLE; INJECTION

PENTOSTATIN

RISING PHARMA

10MG/VIAL

A203554 001 Sep 19, 2014

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE;ORAL

PENTOXIFYLLINE

ANI PHARMS	400MG	A074878	001	Jul 09, 1997
	400MG	A075107	001	Sep 04, 1998
	400MG	A075199	001	Sep 03, 1999
HERITAGE PHARMS INC	400MG	A074877	001	Jul 08, 1997
IMPAX LABS	400MG	A075093	001	Aug 10, 1999
PLIVA	400MG	A074874	001	May 25, 1999
RISING PHARMA	400MG	A074425	001	Jul 08, 1997
PENTOXIL				
UPSHER SMITH LABS	400MG	A074962	001	Mar 31, 1999
TRENENTAL				
+ VALIDUS PHARMS	400MG **	N018631	001	Aug 30, 1984

PERFLUBRON

LIQUID;ORAL

IMAGENT

ALLIANCE PHARM	100%	N020091	001	Aug 13, 1993
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PERFLUOROPOLYMETHYLISOPROPYL ETHER; POLYTETRAFLUOROETHYLENE

PASTE;TOPICAL

SKIN EXPOSURE REDUCTION PASTE AGAINST CHEMICAL WARFARE AGENTS

US ARMY	50%;50%	N021084	001	Feb 17, 2000
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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
STRIDES PHARMA	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

PERINDOPRIL ERBUMINE

ANI PHARMS	2MG	A078138	001	Nov 10, 2009
	4MG	A078138	002	Nov 10, 2009
	8MG	A078138	003	Nov 10, 2009
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

CREAM;TOPICAL

ELIMITE

+ AUROBINDO PHARMA USA	5%	N019855	001	Aug 25, 1989
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LOTION;TOPICAL

NIX

GLAXOSMITHKLINE	1%	N019435	001	Mar 31, 1986
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PERPHENAZINE

CONCENTRATE;ORAL

PERPHENAZINE

PHARM ASSOC	16MG/5ML	A040360	001	May 25, 2001
TRILAFON				
SCHERING	16MG/5ML	N011557	001	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PERPHENAZINE

INJECTABLE; INJECTION

TRILAFON

SCHERING 5MG/ML N011213 002

SYRUP; ORAL

TRILAFON

SCHERING 2MG/5ML N011294 002

TABLET; ORAL

PERPHENAZINE

ANI PHARMS	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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENDIMETRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE;ORAL

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			
FERNDALE 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
FERNDALE LABS	35MG	A086834 001	Sep 15, 1983
INWOOD LABS	35MG	A084740 001	
	35MG	A084741 001	
	35MG	A084742 001	
	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	
NOSTRUM LABS INC	35MG	A203600 001	Dec 27, 2017
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	
UPSHER SMITH LABS	35MG	A084399 001	
USL PHARMA	35MG	A083805 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENDIMETRAZINE TARTRATE

TABLET;ORAL

PHENDIMETRAZINE TARTRATE

	35MG	A084398	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
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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
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PHENSUXIMIDE

CAPSULE;ORAL

MILONTIN

PARKE DAVIS	500MG	N008855	004
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PHENTERMINE HYDROCHLORIDE

CAPSULE;ORAL

FASTIN

GLAXOSMITHKLINE	30MG **	N017352	001
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OBESTIN-30

FERNDAL LABS	30MG	A087144	001
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OBY-TRIM

SHIRE RICHWOOD	30MG	A087764	001	Mar 18, 1982
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ONA-MAST

MAST MM	30MG	A086511	001
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	30MG	A086516	001
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PHENTERMINE HYDROCHLORIDE

ABC HOLDING	30MG	A085411	001
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ABLE	15MG	A040497	001	Mar 13, 2003
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	30MG	A040403	001	Aug 30, 2001
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	30MG	A040427	001	Aug 30, 2001
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BARR	15MG	A090591	001	Mar 18, 2010
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	30MG	A090591	002	Mar 18, 2010
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CAMALL	15MG	A086735	001
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	30MG	A087226	001
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CHARTWELL RX	18.75MG	A088576	001	May 23, 1984
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	30MG	A085417	001
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

PHENTERMINE HYDROCHLORIDE

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	A088948	001	Apr 25, 1986
ELITE LABS INC	A040460	001	Jan 14, 2003
	A040227	001	Jun 18, 1997
	A040448	001	Jan 22, 2003
IVAX PHARMS	A086329	001	
LANNETT CO INC	A091359	001	Jul 16, 2010
SANDOZ	A087208	001	
	A087223	001	
	A088414	001	Oct 19, 1983
SUN PHARM INDUSTRIES	A040527	001	Oct 23, 2003
TEVA	A086911	001	
	A087126	001	
	A087777	001	Nov 01, 1985
	A088612	001	Apr 04, 1984
	A088613	001	Apr 09, 1984
	A088614	001	Apr 09, 1984
TG UNITED INC	A040083	001	Mar 07, 1997
UPSHER SMITH LABS	A084487	001	Apr 09, 1982
	A088430	001	Mar 27, 1984
USL PHARMA	A088797	001	Dec 10, 1984
VITARINE	A087202	001	
	A087235	001	
WATSON LABS	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
BARR	37.5MG	A090470	001 Aug 31, 2009
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
NOVAST LABS	37.5MG	A091451	001 Sep 21, 2012
SANDOZ	8MG	A085671	001
	8MG	A085689	001
SANDOZ INC	30MG	A088605	001 Sep 28, 1987
SUN PHARM INDS INC	37.5MG	A040790	001 Aug 21, 2007
+ 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
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

SOLUTION;INTRAVENOUS

BIORPHEN

+ ETON PHARMS

10MG/ML (10MG/ML)

N212909 002 Mar 11, 2021

PHENYLEPHRINE HYDROCHLORIDE

ACCORD HLTHCARE

10MG/ML (10MG/ML)

A213237 001 Jul 01, 2020

50MG/5ML (10MG/ML)

A213237 002 Jul 01, 2020

100MG/10ML (10MG/ML)

A213237 003 Jul 01, 2020

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

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

UPJOHN 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 100MG EXTENDED A040435 001 Jun 20, 2003

100MG EXTENDED A089441 001 Dec 18, 1986

LUPIN LTD 100MG EXTENDED A211633 001 Sep 30, 2019

SUN PHARM INDS (IN) 100MG EXTENDED A040621 001 Dec 11, 2006

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 100MG PROMPT A080259 001

WATSON LABS 100MG PROMPT A080905 001

INJECTABLE;INJECTION

DILANTIN

PARKE DAVIS 50MG/ML N010151 001

PHENYTOIN SODIUM

AM REGENT 50MG/ML A040781 001 Dec 04, 2007

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

PIMAVANSERIN TARTRATE

TABLET;ORAL

NUPLAZID

+ ACADIA PHARMS INC

EQ 17MG BASE

N207318 001 Apr 29, 2016

PIMOZIDE

TABLET;ORAL

ORAP

+ TEVA

1MG **

N017473 003 Aug 27, 1997

+

2MG **

N017473 001 Jul 31, 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

COSETTE

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

ZYDUS PHARMS

5MG

A209866 001 Aug 18, 2017

10MG

A209866 002 Aug 18, 2017

VISKEN

+ NOVARTIS

5MG **

N018285 001 Sep 03, 1982

+

10MG **

N018285 002 Sep 03, 1982

PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

PIOGLITAZONE HYDROCHLORIDE

MYLAN PHARMS INC

EQ 15MG BASE

A076801 001 Aug 17, 2012

EQ 30MG BASE

A076801 002 Aug 17, 2012

EQ 45MG BASE

A076801 003 Aug 17, 2012

NOSTRUM LABS INC

EQ 15MG BASE

A078472 001 Feb 13, 2013

EQ 30MG BASE

A078472 002 Feb 13, 2013

EQ 45MG BASE

A078472 003 Feb 13, 2013

PICKET PHARMS

EQ 15MG BASE

A078383 001 Mar 12, 2013

EQ 30MG BASE

A078383 002 Mar 12, 2013

EQ 45MG BASE

A078383 003 Mar 12, 2013

TORRENT PHARMS LTD

EQ 15MG BASE

A091298 001 Feb 13, 2013

EQ 30MG BASE

A091298 002 Feb 13, 2013

EQ 45MG BASE

A091298 003 Feb 13, 2013

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

PIPERACILLIN AND TAZOBACTAM

	HOSPIRA INC	EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL	A065386 001	Sep 15, 2009
		EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL	A065386 002	Sep 15, 2009
		EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL	A065386 003	Sep 15, 2009
		EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL	A065446 001	Sep 15, 2009
	MYLAN LABS LTD	EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL	A065458 001	Aug 15, 2014
		EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL	A065458 002	Aug 15, 2014
		EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL	A065458 003	Aug 15, 2014
	ZOSYN			
+	WYETH PHARMS	EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL	N050684 001	Oct 22, 1993
+		EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL	N050684 002	Oct 22, 1993
+		EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL	N050684 003	Oct 22, 1993
+		EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL	N050684 004	Oct 22, 1993

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	
	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
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PIPOBROMAN

TABLET; ORAL

VERCYTE

	ABBOTT	10MG	N016245 001	
		25MG	N016245 002	

PIRBUTEROL ACETATE

AEROSOL, METERED; INHALATION

MAXAIR

	BAUSCH	EQ 0.2MG BASE/INH	N019009 001	Dec 30, 1986
		EQ 0.2MG BASE/INH	N020014 001	Nov 30, 1992

PIRFENIDONE

TABLET; ORAL

ESBRIET

+	GENENTECH INC	534MG **	N208780 002	Jan 11, 2017
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PIROXICAM

CAPSULE; ORAL

PIROXICAM

BRECKENRIDGE	10MG	A208991 001	Feb 21, 2018
	20MG	A208991 002	Feb 21, 2018
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
SUN PHARM INDUSTRIES	10MG	A073536 002	Jan 23, 2008
	20MG	A073536 001	Mar 12, 1993
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 CALCIUM

TABLET; ORAL

PITAVASTATIN CALCIUM

MYLAN

EQ 1MG BASE	A206070 001	Apr 04, 2019
EQ 2MG BASE	A206070 002	Apr 04, 2019
EQ 4MG BASE	A206070 003	Apr 04, 2019

PITAVASTATIN MAGNESIUM

TABLET; ORAL

ZYPITAMAG

+ MEDICURE

EQ 1MG BASE	N208379 001	Jul 14, 2017
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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	
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PODOFILOX

SOLUTION; TOPICAL

PODOFILOX

BAUSCH AND LOMB INC 0.5%

A090184 001	Jul 21, 2010
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POLYESTRADIOL PHOSPHATE

INJECTABLE; INJECTION

ESTRADURIN

WYETH AYERST

40MG/AMP	N010753 001	
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POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL

GLYCOLAX

LANNETT CO INC

17GM/SCOOPFUL	A076652 001	Jul 02, 2004
17GM/PACKET	A090600 001	Oct 06, 2009
17GM/SCOOPFUL	A090600 002	Oct 06, 2009

POLYETHYLENE GLYCOL 3350

BRECKENRIDGE PHARM

NEXGEN PHARMA INC

PADDOCK LLC

17GM/SCOOPFUL	A077736 001	May 26, 2006
17GM/SCOOPFUL	A077706 001	Sep 27, 2006
17GM/SCOOPFUL	A077893 001	May 26, 2006
17GM/SCOOPFUL	A090567 001	Oct 15, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POLYETHYLENE GLYCOL 3350

FOR SOLUTION;ORAL

POLYETHYLENE GLYCOL 3350

TEVA PHARMS

17GM/SCOOPFUL

A077445 001 May 04, 2006

POLYETHYLENE 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

NOSTRUM LABS INC

420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/
BOT

A202060 001 Mar 08, 2017

TRILYTE

AUROBINDO PHARMA USA 420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/
BOT

A076491 001 Feb 05, 2004

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/PACKE
T;2.92GM/PACKET;11.36GM/PACKET

N018983 005 Oct 26, 1984

227.1GM/PACKET;2.82GM/PACKET;6.36GM/PAC
KET;5.53GM/PACKET;21.5GM/PACKET

N018983 004 Oct 26, 1984

227.1GM/BOT;2.82GM/BOT;6.36GM/BOT;5.53G
M/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/PACK
ET;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;5.53G
M/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

GOLYTELY

+ BRAINTREE

227.1GM/PACKET;2.82GM/PACKET;6.36GM/PAC
KET;5.53GM/PACKET;21.5GM/PACKET

N019011 002 Jun 02, 1992

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/PAC
KET;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 UNITS BASE/VIAL A062036 001

POLYMYXIN B SULFATE

GLAND EQ 500,000 UNITS BASE/VIAL A207322 001 Apr 14, 2016

HIKMA EQ 500,000 UNITS BASE/VIAL A060716 001

RISING PHARMA EQ 500,000 UNITS BASE/VIAL A090110 001 Jun 29, 2011

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

POMALIDOMIDE

CAPSULE; ORAL

POMALIDOMIDE

BRECKENRIDGE 1MG A210111 001 Oct 30, 2020

2MG A210111 002 Oct 30, 2020

3MG A210111 003 Oct 30, 2020

4MG A210111 004 Oct 30, 2020

EUGIA PHARMA 1MG A210249 001 Oct 30, 2020

2MG A210249 002 Oct 30, 2020

3MG A210249 003 Oct 30, 2020

4MG A210249 004 Oct 30, 2020

POSACONAZOLE

SUSPENSION; ORAL

POSACONAZOLE

HIKMA 40MG/ML A208773 001 May 15, 2020

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

MICRO-K

+ NESHER PHARMS 8MEQ ** N018238 001

MICRO-K 10

+ NESHER PHARMS 10MEQ ** N018238 002 May 14, 1984

POTASSIUM CHLORIDE

NESHER PHARMS 10MEQ A070980 001 Feb 17, 1987

TEVA 8MEQ A073531 001 Apr 26, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

POTASSIUM CHLORIDE

10MEQ

A073532 001 Apr 26, 1996

FOR SOLUTION;ORAL

POTASSIUM CHLORIDE

STRIDES PHARMA

20MEQ

A211667 001 Mar 11, 2021

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

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 10MEQ

FRESENIUS KABI USA

14.9MG/ML

A211087 001 Sep 09, 2020

746MG/100ML

A211087 002 Sep 09, 2020

POTASSIUM CHLORIDE 20MEQ

FRESENIUS KABI USA

29.8MG/ML

A211087 003 Sep 09, 2020

1.49GM/100ML

A211087 005 May 07, 2021

POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER

+ ICU MEDICAL INC

2.24GM/100ML

N020161 003 Aug 11, 1998

POTASSIUM CHLORIDE 40MEQ

FRESENIUS KABI USA

2.98GM/100ML

A211087 004 Sep 09, 2020

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POTASSIUM CHLORIDE

TABLET, EXTENDED RELEASE;ORAL

KLOTRIX

APOTHECON	10MEQ		N017850	001	
POTASSIUM CHLORIDE					
AMNEAL	15MEQ		A212861	002	May 08, 2020
AMTA	10MEQ		A214395	001	Jan 28, 2021
	20MEQ		A214395	002	Jan 28, 2021
AUROBINDO PHARMA LTD	10MEQ		A214728	001	Mar 31, 2021
	15MEQ		A214728	002	Mar 31, 2021
	20MEQ		A214728	003	Mar 31, 2021
BRECKENRIDGE	10MEQ		A213588	001	Aug 21, 2020
	20MEQ		A213588	002	Aug 21, 2020
COPLEY PHARM	8MEQ		A070618	001	Sep 09, 1987
NESHER PHARMS	20MEQ		A076044	001	Apr 05, 2002
+ SCHERING	10MEQ	**	N019439	002	Jun 13, 1986
+	20MEQ	**	N019439	001	Jun 13, 1986
SIGMAPHARM LABS LLC	8MEQ		A207528	001	Aug 19, 2016
	10MEQ		A207528	002	Aug 19, 2016
STRIDES PHARMA	8MEQ		A206881	001	Jan 22, 2019
	10MEQ		A206630	001	Mar 29, 2019
	10MEQ		A206881	002	Jan 22, 2019
	10MEQ		A210097	001	Jun 17, 2019
	15MEQ		A206630	002	Mar 29, 2019
	20MEQ		A206630	003	Mar 29, 2019
	20MEQ		A210098	001	Apr 26, 2019
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 0.15% IN SODIUM CHLORIDE 0.45%					
FRESENIUS KABI USA	150MG/100ML;450MG/100ML		A212347	001	Sep 17, 2020
POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9%					
FRESENIUS KABI USA	150MG/100ML;900MG/100ML		A212347	003	Jun 02, 2021
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% AND SODIUM CHLORIDE 0.9%					
FRESENIUS KABI USA	300MG/100ML;900MG/100ML		A212347	002	Sep 17, 2020
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%					
BAXTER HLTHCARE	75MG/100ML;900MG/100ML		N017648	004	
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER					
B BRAUN	75MG/100ML;900MG/100ML		N018722	001	Nov 09, 1982
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

POTASSIUM CHLORIDE; SODIUM CHLORIDE; TROMETHAMINE

INJECTABLE; INJECTION

THAM-E

HOSPIRA	370MG/VIAL;1.75GM/VIAL;36GM/VIAL		N013025	001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

THYROSHIELD

ARCO PHARMS LLC

65MG/ML

A077218 001 Jan 12, 2005

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

SOLUTION;INTRAMUSCULAR

PRALIDOXIME CHLORIDE (AUTOINJECTOR)

+ MERIDIAN MEDCL TECHN 600MG/2ML (300MG/ML)

N018986 001 Apr 26, 1983

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

ALEMBIC PHARMS LTD 0.125MG

A078894 001 Oct 08, 2010

0.25MG

A078894 002 Oct 08, 2010

0.5MG

A078894 003 Oct 08, 2010

1MG

A078894 004 Oct 08, 2010

HERITAGE PHARMA AVET 1.5MG

A078894 005 Oct 08, 2010

0.125MG

A077724 001 Feb 19, 2008

0.125MG

A078551 001 Oct 08, 2010

0.125MG

A090241 001 Oct 08, 2010

0.125MG

A091254 001 Nov 30, 2010

0.25MG

A077724 002 Feb 19, 2008

0.25MG

A078551 002 Oct 08, 2010

0.25MG

A090241 002 Oct 08, 2010

0.25MG

A091254 002 Nov 30, 2010

0.5MG

A077724 003 Feb 19, 2008

0.5MG

A078551 003 Oct 08, 2010

0.5MG

A090241 003 Oct 08, 2010

0.5MG

A091254 003 Nov 30, 2010

0.75MG

A090241 004 Oct 08, 2010

0.75MG

A091254 004 Nov 30, 2010

1MG

A077724 004 Feb 19, 2008

1MG

A078551 004 Oct 08, 2010

1MG

A090241 005 Oct 08, 2010

1MG

A091254 005 Nov 30, 2010

1.5MG

A077724 005 Feb 19, 2008

1.5MG

A078551 005 Oct 08, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET;ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

	1.5MG	A090241	006	Oct 08, 2010
	1.5MG	A091254	006	Nov 30, 2010
MACLEODS PHARMS LTD	0.125MG	A202164	001	Sep 20, 2012
	0.25MG	A202164	002	Sep 20, 2012
	0.5MG	A202164	003	Sep 20, 2012
	1MG	A202164	004	Sep 20, 2012
	1.5MG	A202164	005	Sep 20, 2012
MYLAN	0.125MG	A077854	001	Oct 08, 2010
	0.25MG	A077854	002	Oct 08, 2010
	0.5MG	A077854	003	Oct 08, 2010
	0.75MG	A090764	001	Apr 09, 2010
	1MG	A077854	004	Oct 08, 2010
	1.5MG	A077854	005	Oct 08, 2010
NOSTRUM LABS INC	0.125MG	A091450	001	Oct 08, 2010
	0.25MG	A091450	002	Oct 08, 2010
	0.5MG	A091450	003	Oct 08, 2010
	1MG	A091450	004	Oct 08, 2010
	1.5MG	A091450	005	Oct 08, 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
SUN PHARM INDS INC	0.125MG	A091683	001	Mar 27, 2013
	0.25MG	A091683	002	Mar 27, 2013
	0.5MG	A091683	003	Mar 27, 2013
	0.75MG	A091683	004	Mar 27, 2013
	1MG	A091683	005	Mar 27, 2013
	1.5MG	A091683	006	Mar 27, 2013
UNICHEM	0.125MG	A207011	001	Dec 19, 2018
	0.25MG	A207011	002	Dec 19, 2018
	0.5MG	A207011	003	Dec 19, 2018
	0.75MG	A207011	004	Dec 19, 2018
	1MG	A207011	005	Dec 19, 2018
	1.5MG	A207011	006	Dec 19, 2018

TABLET, EXTENDED RELEASE;ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

ALEMBIC PHARMS LTD	0.375MG	A204518	001	Jan 02, 2019
	0.75MG	A204518	002	Jan 02, 2019
	1.5MG	A204518	003	Jan 02, 2019
	2.25MG	A204518	004	Jan 02, 2019
	3MG	A204518	005	Jan 02, 2019
	3.75MG	A204518	006	Jan 02, 2019
	4.5MG	A204518	007	Jan 02, 2019

PRAMLINTIDE ACETATE

INJECTABLE;SUBCUTANEOUS

SYMLIN

ASTRAZENECA AB	EQ 3MG BASE/5ML (EQ 600MCG BASE/ML)	N021332	001	Mar 16, 2005
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PRASUGREL HYDROCHLORIDE

TABLET;ORAL

PRASUGREL

DR REDDYS LABS LTD	EQ 5MG BASE	A205926	001	Jul 07, 2020
	EQ 10MG BASE	A205926	002	Jul 07, 2020

PRAVASTATIN SODIUM

TABLET;ORAL

PRAVACHOL

+	BRISTOL MYERS SQUIBB	10MG **	N019898	002	Oct 31, 1991
+		20MG	N019898	003	Oct 31, 1991
+		40MG	N019898	004	Mar 22, 1993
+		80MG	N019898	008	Dec 18, 2001

PRAVASTATIN SODIUM

HISUN PHARM HANGZHOU	20MG	A206061	001	Nov 23, 2018
	40MG	A206061	002	Nov 23, 2018
	80MG	A206061	003	Nov 23, 2018

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PRAVASTATIN SODIUM

TABLET; ORAL

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
MYLAN PHARMS INC	10MG	A079187 001	May 27, 2010
	20MG	A079187 002	May 27, 2010
	40MG	A079187 003	May 27, 2010
	80MG	A079187 004	May 27, 2010
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
ZYDUS PHARMS USA	10MG	A077751 001	Apr 30, 2008

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	
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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
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

PREDNICARBATE

CREAM; TOPICAL

DERMATOP E EMOLLIENT

+ VALEANT BERMUDA	0.1%	N020279 001	Oct 29, 1993
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PREDNICARBATE

FOUGERA PHARMS	0.1%	A077287 001	Sep 19, 2006
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OINTMENT; TOPICAL

DERMATOP

+ VALEANT PHARMS NORTH	0.1% **	N019568 001	Sep 23, 1991
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PREDNISOLONE

CREAM; TOPICAL

METI-DERM

SCHERING	0.5%	N010209 002	
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SYRUP; ORAL

PREDNISOLONE

IVAX SUB TEVA PHARMS	15MG/5ML	A040287 001	May 28, 1999
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISOLONE

SYRUP; ORAL

PREDNISOLONE

NESHER PHARMS	5MG/5ML	A040423	001	Oct 22, 2001
	15MG/5ML	A040364	001	Apr 10, 2002
PHARM ASSOC	5MG/5ML	A040570	001	Aug 25, 2005
	15MG/5ML	A040571	001	Aug 25, 2005
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
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TABLET; ORAL

CORTALONE

HALSEY	1MG	A080304	003	
	2.5MG	A080304	002	
	5MG	A080304	001	

DELTA-CORTEF

PHARMACIA AND UPJOHN	5MG	N009987	004	
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FERNISOLONE-P

FERNDALE LABS	5MG	A083941	001	
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PREDNISOLONE

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	
RISING	5MG	A084773	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	
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PREDNISOLONE ACETATE

INJECTABLE; INJECTION

METICORTELONE

SCHERING	25MG/ML	N010255	002	
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PREDNISOLONE ACETATE

AKORN	25MG/ML	A083032	001	
	50MG/ML	A084492	001	
BEL MAR	25MG/ML	A083738	001	
	50MG/ML	A083738	002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISOLONE ACETATE

INJECTABLE; INJECTION

PREDNISOLONE ACETATE

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
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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

EYEVANCE	0.125%	N017468	001
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PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC

CETAPRED

ALCON	0.25%;10%	A087771	001	Aug 06, 1993
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METIMYD

SCHERING	0.5%;10%	N010210	002	Sep 09, 1984
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PREDSULFAIR

PHARMAFAIR	0.5%;10%	A088032	001	Apr 15, 1983
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VASOCIDIN

NOVARTIS	0.5%;10%	A088791	001	Oct 05, 1984
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SUSPENSION; OPHTHALMIC

ISOPTO CETAPRED

ALCON	0.25%;10%	A087547	001
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SUSPENSION/DROPS; OPHTHALMIC

METIMYD

SCHERING	0.5%;10%	N010210	001
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PREDAMIDE

AKORN	0.5%;10%	A088059	001	Jul 29, 1983
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PREDSULFAIR

PHARMAFAIR	0.5%;10%	A088007	001	Apr 19, 1983
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PREDSULFAIR II

PHARMAFAIR	0.2%;10%	A088837	001	Dec 24, 1985
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SULPHRIN

BAUSCH AND LOMB	0.5%;10%	A088089	001	Dec 28, 1982
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PREDNISOLONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

HYDELTRASOL

MERCK	EQ 20MG PHOSPHATE/ML	N011583	002
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PREDNISOLONE SODIUM PHOSPHATE

WATSON LABS	EQ 20MG PHOSPHATE/ML	A080517	001
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OINTMENT; OPHTHALMIC, OTIC

HYDELTRASOL

MERCK	EQ 0.25% PHOSPHATE	N011028	001
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SOLUTION; ORAL

ORAPRED

CONCORDIA PHARMS INC	EQ 15MG BASE/5ML **	A075117	001	Dec 14, 2000
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PREDNISOLONE SODIUM PHOSPHATE

AMNEAL PHARMS	EQ 15MG BASE/5ML	A078345	001	Mar 10, 2009
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BAUSCH	EQ 15MG BASE/5ML	A075250	001	Jul 12, 2002
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NESHER PHARMS	EQ 5MG BASE/5ML	A076982	001	May 24, 2005
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	EQ 15MG BASE/5ML	A076988	001	May 24, 2005
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PHARM ASSOC	EQ 5MG BASE/5ML	A076123	001	Dec 23, 2002
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VINTAGE	EQ 15MG BASE/5ML	A079010	001	May 26, 2009
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VINTAGE PHARMS	EQ 5MG BASE/5ML	A078416	001	Oct 31, 2007
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WE PHARMS	EQ 5MG BASE/5ML	A075181	001	Dec 23, 2002
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SOLUTION/DROPS; OPHTHALMIC

INFLAMASE FORTE

+ NOVARTIS	EQ 0.9% PHOSPHATE	A080751	002
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION/DROPS;OPHTHALMIC

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

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

TABLET, ORALLY DISINTEGRATING;ORAL

PREDNISOLONE SODIUM PHOSPHATE

MYLAN PHARMS INC

EQ 10MG BASE

A202179 001 Apr 10, 2013

EQ 15MG BASE

A202179 002 Apr 10, 2013

EQ 30MG BASE

A202179 003 Apr 10, 2013

PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC

SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE

SANDOZ INC

EQ 0.23% PHOSPHATE;10%

A073630 001 May 27, 1993

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

PREDNISON

SOLUTION;ORAL

PREDNISON

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

FERNISON

FERNDAL 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISON

TABLET;ORAL

PARACORT

PARKE DAVIS 5MG N010962 002

PREDNICEN-M

SCHWARZ PHARMA 5MG A084655 001

PREDNISON

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

BUNDY 5MG A083676 001

CHARTWELL RX 5MG A083059 001

CONTRACT PHARMACAL 5MG A080209 001

DURAMED PHARMS BARR 5MG A088394 001 Oct 04, 1983

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

50MG A088465 001 Jun 01, 1984

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

RISING 5MG A084774 001

10MG A089983 001 Jan 12, 1989

20MG A085813 001

50MG A089984 001 Jan 12, 1989

ROXANE 20MG N017109 001

25MG A087833 001 May 04, 1982

SANDOZ 5MG A080336 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISON

TABLET; ORAL

PREDNISON

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	
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

TABLET, DELAYED RELEASE; ORAL

PREDNISON

ACTAVIS LABS FL INC

1MG

A204867 001 Apr 25, 2017

2MG

A204867 002 Apr 25, 2017

5MG

A204867 003 Apr 25, 2017

PREGABALIN

CAPSULE; ORAL

PREGABALIN

MYLAN

25MG

A091228 001 Sep 20, 2019

50MG

A091228 002 Sep 20, 2019

75MG

A091228 003 Sep 20, 2019

100MG

A091228 004 Sep 20, 2019

150MG

A091228 005 Sep 20, 2019

200MG

A091228 006 Sep 20, 2019

225MG

A091228 007 Sep 20, 2019

300MG

A091228 008 Sep 20, 2019

TABLET, EXTENDED RELEASE; ORAL

PREGABALIN

ALVOGEN

82.5MG

A211687 001 Jul 06, 2021

165MG

A211687 002 Jul 06, 2021

330MG

A211687 003 Jul 06, 2021

MYLAN

330MG

A211431 001 Jul 02, 2021

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PRIMAQUINE PHOSPHATE

TABLET; ORAL

PRIMAQUINE PHOSPHATE

ALVOGEN

EQ 15MG BASE

A203924 001 Feb 03, 2014

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

250MG

A040667 002 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

MYLAN

500MG

A084211 002

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

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROCAINAMIDE HYDROCHLORIDE

CAPSULE; ORAL

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	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

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
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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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINE HYDROCHLORIDE

	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
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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

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

HIKMA EQ 5MG BASE/ML A089523 001 May 03, 1988

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

WYETH AYERST EQ 5MG BASE/ML A086348 001

SYRUP; ORAL

COMPAZINE

GLAXOSMITHKLINE EQ 5MG BASE/5ML N011188 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROCHLORPERAZINE EDISYLATE

SYRUP;ORAL

PROCHLORPERAZINE EDISYLATE

ALPHARMA US PHARMS EQ 5MG BASE/5ML
MORTON GROVE EQ 5MG BASE/5MLA087154 001 Sep 01, 1982
A088597 001 Oct 25, 1984PROCHLORPERAZINE MALEATE

CAPSULE, EXTENDED RELEASE;ORAL

COMPAZINE

GLAXOSMITHKLINE EQ 10MG BASE
EQ 10MG BASE
EQ 15MG BASE
EQ 15MG BASE
EQ 30MG BASE
EQ 75MG BASEN011000 001
N021019 001 Oct 06, 1999
N011000 002
N021019 002 Oct 06, 1999
N011000 003
N011000 004

TABLET;ORAL

COMPAZINE

+ GLAXOSMITHKLINE EQ 5MG BASE **
+ EQ 10MG BASE **
+ EQ 25MG BASE **N010571 001
N010571 002
N010571 003

PROCHLORPERAZINE

WATSON LABS EQ 5MG BASE
EQ 10MG BASE
EQ 25MG BASEA085580 001
A085178 001
A085579 001

PROCHLORPERAZINE MALEATE

DURAMED PHARMS BARR EQ 5MG BASE
EQ 5MG BASE
EQ 10MG BASE
EQ 10MG BASE
EQ 25MG BASEA040207 001 May 01, 1997
A089484 001 Jan 20, 1987
A040207 002 May 01, 1997
A089485 001 Jan 20, 1987
A089486 001 Jan 20, 1987IVAX SUB TEVA PHARMS EQ 5MG BASE
EQ 10MG BASEA040162 001 Jan 20, 1998
A040162 002 Jan 20, 1998SANDOZ EQ 5MG BASE
EQ 10MG BASE
EQ 25MG BASEA040101 001 Jul 19, 1996
A040101 002 Jul 19, 1996
A040101 003 Jul 19, 1996TEVA PHARMS EQ 5MG BASE
EQ 10MG BASEA040120 001 Jul 11, 1996
A040120 002 Jul 11, 1996PROCYCLIDINE HYDROCHLORIDE

TABLET;ORAL

KEMADRIN

MONARCH PHARMS 2MG
5MGN009818 005
N009818 003PROGESTERONE

CAPSULE;ORAL

PROGESTERONE

TEVA PHARMS 100MG
200MGA202121 001 Feb 29, 2012
A202121 002 Feb 29, 2012

PROMETRIUM

VIRTUS PHARMS 300MG

N019781 003 Oct 15, 1999

INJECTABLE;INJECTION

PROGESTERONE

AM REGENT 50MG/ML
LILLY 25MG/ML
50MG/MLA090845 001 Jun 22, 2009
N009238 002
N009238 001

INSERT, EXTENDED RELEASE;INTRAUTERINE

PROGESTASERT

ALZA 38MG

N017553 001

SYSTEM;VAGINAL

MILPROSA

+ FERRING PHARMS INC 1.78GM

N201110 001 Apr 29, 2020

PROMAZINE HYDROCHLORIDE

CONCENTRATE;ORAL

SPARINE

WYETH AYERST 30MG/ML
100MG/MLN010942 001
N010942 004

INJECTABLE;INJECTION

PROMAZINE HYDROCHLORIDE

WATSON LABS 25MG/ML

A084510 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROMAZINE HYDROCHLORIDE	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
AM REGENT	25MG/ML	A040515	001 Mar 19, 2003
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
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
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ZIPAN-50

ALTANA	50MG/ML	A083997	002
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SUPPOSITORY; RECTAL

PHENERGAN

+ MYLAN	12.5MG **	N010926	002
+	25MG **	N010926	001
+	50MG **	N011689	001

PROMETHACON

POLYMEDICA	25MG	A084901	001
	50MG	A084902	001

PROMETHAZINE HYDROCHLORIDE

ABLE	12.5MG	A040504	001 Apr 11, 2003
	25MG	A040504	002 Apr 11, 2003
	50MG	A040449	001 Feb 27, 2003
WATSON LABS INC	12.5MG	A040479	001 Jun 24, 2003
	25MG	A040479	002 Jun 24, 2003

SYRUP; ORAL

MYMETHAZINE FORTIS

USL PHARMA	25MG/5ML	A087996	001 Jan 18, 1983
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

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	12.5MG		A040791	002 Feb 12, 2008
	25MG		A040791	003 Feb 12, 2008
	25MG		A084214	002 Jul 07, 1982
	50MG		A040791	001 May 20, 2008
INVATECH	12.5MG		A084233	001
	25MG		A085146	001
	50MG		A085146	002
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
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
REMSSED				
BRISTOL MYERS SQUIBB	25MG		A083176	002
	50MG		A083176	001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPAPENONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

PROPAPENONE HYDROCHLORIDE

MYLAN	225MG	A203803 001	Apr 29, 2016
	325MG	A203803 002	Apr 29, 2016
	425MG	A203803 003	Apr 29, 2016

TABLET;ORAL

PROPAPENONE HYDROCHLORIDE

NESHER PHARMS	150MG	A076193 001	Feb 07, 2002
	225MG	A076193 002	Feb 07, 2002
	300MG	A076193 003	Feb 07, 2002

RYTHMOL

+	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	
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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	
HIKMA	7.5MG	A080927 001	
	15MG	A080927 002	
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	

PROPARACAINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

KAINAIR

PHARMAFAIR	0.5%	A088087 001	Jun 07, 1983
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OPHTHAINE

+	APOTHECON	0.5% **	N008883 001
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OPHTHETIC

+	ALLERGAN	0.5% **	N012583 001
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PARACAINE

OPTOPICS	0.5%	A087681 001	Aug 05, 1982
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PROPARACAINE HYDROCHLORIDE

SOLA BARNES HIND	0.5%	A084144 001	
	0.5%	A084151 001	

PROPIOLACTONE

SOLUTION;IRRIGATION

BETAPRONE

FOREST LABS	N/A	N011657 001	
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PROPIOMAZINE HYDROCHLORIDE

INJECTABLE;INJECTION

LARGON

WEST-WARD PHARMS INT	20MG/ML	N012382 002	
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PROPOFOL

INJECTABLE;INJECTION

DIPRIVAN

FRESENIUS KABI USA	10MG/ML	N019627 001	Oct 02, 1989
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PROPOFOL

TEVA PARENTERAL	10MG/ML	A075392 001	Sep 19, 2000
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DARVON

XANODYNE PHARM	32MG	N010997 001
	65MG	N010997 003

DOLENE

HERITAGE PHARMS INC	65MG	A080530 001
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KESSO-GESIC

MK LABS	65MG	A083544 001
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PROPHENE 65

HALSEY	65MG	A083538 002
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PROPOXYPHENE HYDROCHLORIDE

ALRA	65MG	A083184 001
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IMPAX LABS	65MG	A083317 001
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IVAX SUB TEVA PHARMS	32MG	A083597 001
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MUTUAL PHARM	65MG	A083186 001
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MYLAN	32MG	A083528 001
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	65MG	A040569 001
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	65MG	A083299 001
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NEXGEN PHARMA INC	65MG	A083185 001
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PAR PHARM	65MG	A080269 001
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PUREPAC PHARM	65MG	A083278 001
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PVT FORM	32MG	A083464 001
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	65MG	A083113 001
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ROXANE	32MG	A083089 001
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	65MG	A083089 002
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SANDOZ	32MG	A084014 001
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	65MG	A083125 002
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	65MG	A083688 001
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	65MG	A083870 002
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	65MG	A086495 001
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TEVA	65MG	A088615 001
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VALEANT PHARM INTL	65MG	A080783 001
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VINTAGE PHARMS	65MG	A040908 001
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WATSON LABS	65MG	A080908 002
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	65MG	A085190 001
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WEST WARD	65MG	A083501 001
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WHITEWORTH TOWN PLSN	65MG	A084551 001
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PROPOXYPHENE HYDROCHLORIDE 65

WARNER CHILCOTT	65MG	A083786 001
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PROPOXYPHENE NAPSYLATE

SUSPENSION; ORAL

DARVON-N

AAIPHARMA LLC	50MG/5ML	N016861 001
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TABLET; ORAL

DARVON-N

XANODYNE PHARM	100MG	N016862 002
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PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PROPRANOLOL HYDROCHLORIDE

INWOOD LABS	60MG	A072499 001	Apr 11, 1989
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	80MG	A072500 001	Apr 11, 1989
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	120MG	A072501 001	Apr 11, 1989
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	160MG	A072502 001	Apr 11, 1989
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MYLAN	60MG	A078022 001	Feb 15, 2007
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	80MG	A078022 002	Feb 15, 2007
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	120MG	A078022 003	Feb 15, 2007
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	160MG	A078022 004	Feb 15, 2007
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UPSHER SMITH LABS	60MG	A078311 001	Mar 06, 2009
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	80MG	A078311 002	Mar 06, 2009
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	120MG	A078311 003	Mar 06, 2009
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	160MG	A078311 004	Mar 06, 2009
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CONCENTRATE; ORAL

PROPRANOLOL HYDROCHLORIDE INTENSOL

ROXANE	80MG/ML	A071388 001	May 15, 1987
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INJECTABLE; INJECTION

PROPRANOLOL HYDROCHLORIDE

+ BAXTER HLTHCARE CORP	1MG/ML **	N016419 001
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPRANOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

PROPRANOLOL HYDROCHLORIDE

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

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

ANI PHARMS 60MG A071791 001 Jul 15, 1987

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

IMPAX LABS INC 80MG A071976 001 Apr 06, 1988

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPRANOLOL HYDROCHLORIDE

TABLET;ORAL

PROPRANOLOL HYDROCHLORIDE

SCHERING	10MG	A070120	001	Aug 06, 1985
	20MG	A070121	001	Aug 06, 1985
	40MG	A070122	001	Aug 06, 1985
	60MG	A070123	001	Oct 29, 1986
	80MG	A070124	001	Aug 06, 1985
STRIDES PHARMA	90MG	A071288	001	Oct 22, 1986
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	A070178	002	Apr 23, 2018
	60MG	A070381	001	Mar 19, 1987
	60MG	A071098	001	Oct 06, 1986
	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

PROPYLIODONE

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	50MG	A080215	001
CHARTWELL RX	50MG	A084543	001
HALSEY	50MG	A080015	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

HIKMA	10MG/ML	A089474	001	Nov 05, 1986
	10MG/ML	A089475	001	Nov 05, 1986
+ LILLY	10MG/ML **	N006460	002	
PHARMACIA AND UPJOHN	50MG/VIAL	N007413	001	
	250MG/VIAL	N007413	002	Aug 02, 1984

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	
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PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

VIVACTIL

HERITAGE PHARMA AVET	5MG	A073644	001	Aug 24, 1995
	10MG	A073645	001	Aug 24, 1995
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
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HISTAFED

CENCI	30MG/5ML; 1.25MG/5ML	A088283	001	Apr 20, 1984
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MYFED

USL PHARMA	30MG/5ML; 1.25MG/5ML	A088116	001	Mar 04, 1983
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TRILITRON

NEWTRON PHARMS	30MG/5ML; 1.25MG/5ML	A088474	001	Feb 12, 1985
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TABLET; ORAL

ALLERFED

PVT FORM	60MG; 2.5MG	A088860	001	Jan 31, 1985
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CORPHED

FOSUN PHARMA	60MG; 2.5MG	A088602	001	Apr 11, 1985
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PSEUDOEPHEDRINE HYDROCHLORIDE AND TRIPROLIDINE HYDROCHLORIDE

SANDOZ	60MG; 2.5MG	A088193	001	May 17, 1983
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TRILITRON

NEWTRON PHARMS	60MG; 2.5MG	A088515	001	Jan 09, 1985
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TRIPHED

TEVA	60MG; 2.5MG	A088630	001	May 17, 1984
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TRIPROLIDINE AND PSEUDOEPHEDRINE

WATSON LABS	60MG; 2.5MG	A088318	002	Jan 13, 1984
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WEST WARD	60MG; 2.5MG	A088117	001	Apr 19, 1983
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

TABLET; ORAL

TRIPROLIDINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

IVAX SUB TEVA PHARMS 60MG;2.5MG

A085273 001 Dec 12, 1984

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 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

DR REDDYS 100MG/ML

A080572 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

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
ALEMBIC PHARMS LTD	EQ 25MG BASE	A203390 001	Oct 28, 2014
	EQ 50MG BASE	A203390 002	Oct 28, 2014
	EQ 100MG BASE	A203390 003	Oct 28, 2014
	EQ 200MG BASE	A203390 004	Oct 28, 2014
	EQ 300MG BASE	A203390 005	Oct 28, 2014
	EQ 400MG BASE	A203390 006	Oct 28, 2014
JUBILANT GENERICS	EQ 25MG BASE	A203150 001	Nov 26, 2013
MYLAN PHARMS INC	EQ 25MG BASE	A090323 001	Mar 27, 2012
TORRENT PHARMS LTD	EQ 25MG BASE	A200363 001	Mar 27, 2012
	EQ 50MG BASE	A200363 002	Mar 27, 2012
	EQ 100MG BASE	A200363 003	Mar 27, 2012
	EQ 200MG BASE	A200363 004	Mar 27, 2012
	EQ 300MG BASE	A200363 005	Mar 27, 2012
	EQ 400MG BASE	A200363 006	Mar 27, 2012
SEROQUEL			
+ ASTRAZENECA	EQ 150MG BASE **	N020639 004	Dec 20, 1998
TABLET, EXTENDED RELEASE;ORAL			
QUETIAPINE FUMARATE			
AMNEAL PHARMS	EQ 400MG BASE	A211405 001	Oct 26, 2018
ANCHEN PHARMS	EQ 150MG BASE	A090757 001	Dec 01, 2017
	EQ 200MG BASE	A090757 002	Dec 01, 2017
	EQ 300MG BASE	A090757 003	Dec 01, 2017
	EQ 400MG BASE	A090757 004	Dec 01, 2017
RISING	EQ 50MG BASE	A202228 001	Feb 02, 2021
	EQ 150MG BASE	A202228 002	Feb 02, 2021
	EQ 200MG BASE	A202228 003	Feb 02, 2021
	EQ 300MG BASE	A202228 004	Feb 02, 2021
	EQ 400MG BASE	A202228 005	Feb 02, 2021

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
ANI PHARMS	EQ 5MG BASE	A075504 001	Aug 24, 2007
	EQ 10MG BASE	A075504 002	Aug 24, 2007
	EQ 20MG BASE	A075504 003	Aug 24, 2007
	EQ 40MG BASE	A075504 004	Aug 24, 2007
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 5MG BASE	A076694 001	Dec 23, 2004
	EQ 10MG BASE	A076036 002	Jan 28, 2005
	EQ 10MG BASE	A076694 002	Dec 23, 2004
	EQ 20MG BASE	A076036 003	Jan 28, 2005
	EQ 20MG BASE	A076694 003	Dec 23, 2004
	EQ 40MG BASE	A076036 004	Jan 28, 2005
	EQ 40MG BASE	A076694 004	Dec 23, 2004
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

QUINAPRIL HYDROCHLORIDE

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
	EQ 10MG BASE	A076803 002
	EQ 20MG BASE	A076803 003
	EQ 40MG BASE	A076803 004

QUINESTROL

TABLET; ORAL

ESTROVIS

PARKE DAVIS

0.1MG	N016768 002
0.2MG	N016768 003

QUINETHAZONE

TABLET; ORAL

HYDROMOX

LEDERLE

50MG	N013264 001
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QUINETHAZONE; RESERPINE

TABLET; ORAL

HYDROMOX R

LEDERLE

50MG; 0.125MG	N013927 001
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QUINIDINE GLUCONATE

INJECTABLE; INJECTION

QUINIDINE GLUCONATE

+ LILLY

80MG/ML	N007529 002	Feb 10, 1989
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TABLET; ORAL

QUINACT

BAYER HLTHCARE

266MG	A085978 001
400MG	A086099 001

TABLET, EXTENDED RELEASE; ORAL

DURAQUIN

WARNER CHILCOTT

330MG	N017917 001
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QUINAGLUTE

+ BAYER HLTHCARE

324MG	N016647 001
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QUINALAN

LANNETT

324MG	A088081 001	Feb 10, 1986
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QUINATIME

WATSON LABS

324MG	A087448 001
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QUINIDINE GLUCONATE

ANI PHARMS

324MG	A087810 001	Sep 29, 1982
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ASCOT

324MG	A088582 001	Jun 17, 1985
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CYCLE PHARMS LTD

324MG	A088431 001	Jan 06, 1984
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HALSEY

324MG	A089476 001	Apr 10, 1987
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RISING

324MG	A089894 001	Dec 15, 1988
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SUPERPHARM

324MG	A089164 001	Nov 21, 1985
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WATSON LABS

324MG	A087785 001	Jan 24, 1983
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QUINIDINE POLYGALACTURONATE

TABLET; ORAL

CARDIOQUIN

PHARM RES ASSOC

275MG	N011642 002
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QUINIDINE SULFATE

CAPSULE; ORAL

CIN-QUIN

SOLVAY

200MG	A085296 001
300MG	A085297 001

QUINIDINE SULFATE

LILLY

200MG	A085103 001
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TABLET; ORAL

CIN-QUIN

+ SOLVAY

100MG	A085299 001
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+

200MG	A084932 001
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300MG	A085298 001
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

QUINIDINE SULFATE

TABLET;ORAL

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	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
	200MG	A081030	001	Apr 14, 1989
	300MG	A081031	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	A083288	001	
	200MG	A085140	002	
	300MG	A085583	001	
WHITEWORTH TOWN PLSN	200MG	A085444	001	

QUINORA

KEY PHARMS 200MG A083576 001

+ SCHERING 300MG A085222 001

TABLET, EXTENDED RELEASE;ORAL

QUINIDEX

WYETH PHARMS INC 300MG N012796 002

QUINIDINE SULFATE

COSETTE 300MG A040045 001 Jun 30, 1994

QUININE SULFATE

CAPSULE;ORAL

QUININE SULFATE

MYLAN PHARMS INC 324MG A202581 001 Dec 14, 2012

RABEPRAZOLE SODIUM

CAPSULE, DELAYED RELEASE;ORAL

ACIPHEX SPRINKLE

+ AYTU 5MG N204736 001 Mar 26, 2013

+ 10MG N204736 002 Mar 26, 2013

TABLET, DELAYED RELEASE;ORAL

ACIPHEX

+ WOODWARD 10MG ** N020973 001 May 29, 2002

RABEPRAZOLE SODIUM

MYLAN 20MG A076885 001 Nov 08, 2013

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
	10MG	A077004 004	Aug 07, 2008
RANBAXY LABS LTD	5MG	A078849 001	Mar 06, 2009
	10MG	A078849 002	Mar 06, 2009
TEVA PHARMS	1.25MG	A077470 001	Jun 18, 2008
	2.5MG	A077470 002	Jun 18, 2008
	5MG	A077470 003	Jun 18, 2008
	10MG	A077470 004	Jun 18, 2008
WATSON LABS	5MG	A076549 003	Oct 24, 2005
YAOPHARMA CO LTD	1.25MG	A077514 001	Jun 18, 2008
	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	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
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RANITIDINE HYDROCHLORIDE

CAPSULE; ORAL

RANITIDINE HYDROCHLORIDE

AJANTA PHARMA LTD	EQ 150MG BASE	A209859 001	Sep 27, 2018
	EQ 300MG BASE	A209859 002	Sep 27, 2018
APPCO	EQ 150MG BASE	A211893 001	Apr 05, 2019
	EQ 300MG BASE	A211893 002	Apr 05, 2019
AUROBINDO PHARMA LTD	EQ 150MG BASE	A211058 001	Jul 16, 2018
	EQ 300MG BASE	A211058 002	Jul 16, 2018
MYLAN	EQ 150MG BASE	A075564 001	Oct 27, 2000
	EQ 300MG BASE	A075564 002	Oct 27, 2000
NOVITIUM PHARMA	EQ 150MG BASE	A210681 001	Nov 23, 2018
	EQ 300MG BASE	A210681 002	Nov 23, 2018
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
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ZANTAC 300

+ GLAXOSMITHKLINE	EQ 300MG BASE **	N020095 002	Mar 08, 1994
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GRANULE, EFFERVESCENT; ORAL

ZANTAC 150

GLAXO GRP LTD	EQ 150MG BASE/PACKET	N020251 002	Mar 31, 1994
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RANITIDINE HYDROCHLORIDE

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

ACTAVIS MID ATLANTIC EQ 15MG BASE/ML A076124 001 Feb 21, 2007

AKORN EQ 15MG BASE/ML A091078 001 Mar 22, 2011

AMNEAL PHARMS EQ 15MG BASE/ML A078312 001 Sep 02, 2008

APOTEX INC EQ 15MG BASE/ML A077602 001 Sep 17, 2007

AUROBINDO PHARMA LTD EQ 15MG BASE/ML A090623 001 Jul 28, 2010

NOSTRUM LABS INC EQ 15MG BASE/ML A078684 001 Aug 27, 2009

EQ 15MG BASE/ML A091091 001 Sep 20, 2011

RANBAXY EQ 15MG BASE/ML A078448 001 Dec 13, 2007

TARO EQ 15MG BASE/ML A077476 001 Jun 13, 2011

TOLMAR EQ 15MG BASE/ML A090054 001 Nov 15, 2010

TORRENT EQ 15MG BASE/ML A090102 001 May 26, 2009

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

AMNEAL PHARMS NY EQ 150MG BASE A077824 001 Oct 13, 2006

EQ 300MG BASE A077824 002 Oct 13, 2006

ANI PHARMS EQ 75MG BASE A075212 001 Jan 14, 2000

EQ 75MG BASE A075296 001 Jan 14, 2000

EQ 150MG BASE A074488 001 Jul 31, 1997

EQ 150MG BASE A077426 001 Dec 19, 2005

EQ 300MG BASE A074488 002 Jul 31, 1997

EQ 300MG BASE A077426 002 Dec 19, 2005

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

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

GRANULES EQ 150MG BASE A210243 001 Aug 20, 2018

EQ 150MG BASE A210243 002 Aug 20, 2018

HERITAGE PHARMA AVET EQ 150MG BASE A075165 001 Sep 30, 1998

EQ 300MG BASE A075165 002 Sep 30, 1998

MYLAN EQ 75MG BASE A075497 001 Jan 14, 2000

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

NOSTRUM LABS INC EQ 150MG BASE A203694 001 Nov 30, 2017

EQ 300MG BASE A203694 002 Nov 30, 2017

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

STRIDES PHARMA EQ 75MG BASE A201745 001 Feb 29, 2012

EQ 75MG BASE A209160 001 Mar 05, 2018

EQ 150MG BASE A200536 001 Jun 28, 2011

EQ 150MG BASE A205512 001 Aug 22, 2016

EQ 150MG BASE A209161 001 Feb 22, 2018

EQ 150MG BASE A210010 001 Aug 01, 2018

EQ 300MG BASE A205512 002 Aug 22, 2016

EQ 300MG BASE A210010 002 Aug 01, 2018

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 150MG BASE A074864 001 Oct 20, 1997

EQ 300MG BASE A074864 002 Oct 20, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RANITIDINE HYDROCHLORIDE

TABLET;ORAL

RANITIDINE HYDROCHLORIDE

WOCKHARDT

EQ 75MG BASE

A076760 001 Feb 24, 2006

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

WOCKHARDT LTD

EQ 150MG BASE

A075208 001 Dec 17, 1998

EQ 300MG BASE

A075208 002 Dec 17, 1998

ZANTAC 150

+ GLAXO GRP LTD

EQ 150MG BASE **

N018703 001 Jun 09, 1983

+ SANOFI US

EQ 150MG BASE

N021698 001 Aug 31, 2004

+

EQ 150MG BASE

N021698 002 Mar 13, 2007

ZANTAC 300

+ GLAXO GRP LTD

EQ 300MG BASE **

N018703 002 Dec 09, 1985

ZANTAC 75

+ SANOFI US

EQ 75MG BASE

N020520 001 Dec 19, 1995

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

ACCORD HLTHCARE

500MG

A212930 001 May 18, 2021

1GM

A212930 002 May 18, 2021

AMNEAL

500MG

A207690 001 Mar 11, 2021

1GM

A207690 002 Mar 11, 2021

ANI PHARMS

500MG

A210482 001 Oct 29, 2019

1GM

A210482 002 Oct 29, 2019

CIPLA

500MG

A211291 001 May 28, 2019

1GM

A211291 002 May 28, 2019

RAPACURONIUM BROMIDE

INJECTABLE;INJECTION

RAPLON

ORGANON USA INC

100MG/VIAL

N020984 001 Aug 18, 1999

200MG/VIAL

N020984 002 Aug 18, 1999

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

WATSON LABS INC

EQ 0.5MG BASE

A201823 001 Jul 01, 2013

EQ 1MG BASE

A201823 002 Jul 01, 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

KONGLUCOID

PANRAY

50MG

N009278 001

100MG

N009278 002

RAUDIXIN

APOTHECON

50MG

N008842 001

100MG

N008842 002

RAUSERPIN

FERNDALE LABS

50MG

N009926 002

100MG

N009926 004

RAUVAL

PAL PAK

50MG

N009108 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RAUWOLFIA SERPENTINA ROOT

TABLET; ORAL

RAUVAL

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

REPAGLINIDE

TABLET; ORAL

PRANDIN

+ GEMINI LABS LLC

0.5MG **

N020741 001 Dec 22, 1997

+

1MG **

N020741 002 Dec 22, 1997

+

2MG **

N020741 003 Dec 22, 1997

REPAGLINIDE

ACTAVIS TOTOWA

0.5MG

A090008 001 Jan 22, 2014

1MG

A090008 002 Jan 22, 2014

2MG

A090008 003 Jan 22, 2014

MYLAN

0.5MG

A090252 001 Aug 23, 2013

1MG

A090252 002 Jan 22, 2014

2MG

A090252 003 Jan 22, 2014

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RESERPINE

TABLET; ORAL

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	
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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RESERPINE

TABLET; ORAL

SERPATE

VALE	0.1MG	N009453	001
	0.25MG	N009453	002

SERPIVITE

VITARINE	0.25MG	N009645	002
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RESERPINE; TRICHLORMETHIAZIDE

TABLET; ORAL

METATENSIN #2

SANOFI AVENTIS US	0.1MG;2MG	N012972	001
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METATENSIN #4

SANOFI AVENTIS US	0.1MG;4MG	N012972	002
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NAQUIVAL

SCHERING	0.1MG;4MG	N012265	003
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TRICHLORMETHIAZIDE W/ RESERPINE

WATSON LABS	0.1MG;4MG	A085248	001
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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
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RIBASPHERE

CHARTWELL RX	200MG	A076203	001	Apr 06, 2004
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RIBAVIRIN

CHARTWELL RX	200MG	A076192	001	Apr 06, 2004
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TEVA	200MG	A076277	001	Oct 04, 2004
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SOLUTION; ORAL

REBETOL

+ SCHERING	40MG/ML	N021546	001	Jul 29, 2003
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TABLET; ORAL

COPEGUS

+ ROCHE	200MG **	N021511	001	Dec 03, 2002
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+	400MG **	N021511	002	Jun 21, 2005
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RIBAVIRIN

BEXIMCO PHARMS USA	200MG	A202546	001	Aug 12, 2014
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	400MG	A202546	002	Aug 12, 2014
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	500MG	A202546	003	Aug 12, 2014
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	600MG	A202546	004	Aug 12, 2014
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CHARTWELL RX	200MG	A077456	001	Dec 05, 2005
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	400MG	A077456	002	Dec 05, 2005
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	600MG	A077456	003	Dec 05, 2005
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HERITAGE PHARMA AVET	200MG	A077053	001	Dec 05, 2005
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ZYDUS PHARMS USA	400MG	A077094	002	Mar 16, 2007
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	500MG	A077094	004	Apr 18, 2008
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	600MG	A077094	003	Mar 16, 2007
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RIFAMPIN

CAPSULE; ORAL

RIFADIN

SANOFI AVENTIS US	150MG	A062303	001
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+	300MG	N050420	001
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INJECTABLE; INJECTION

RIFAMPIN

EMCURE PHARMS LTD	600MG/VIAL	A204101	001	Aug 18, 2014
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WATSON PHARMS TEVA	600MG/VIAL	A206736	001	Jan 19, 2016
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RILUZOLE

TABLET; ORAL

RILUZOLE

APOTEX CORP	50MG	A091300	001	Jun 18, 2013
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DAITO PHARMS CO LTD	50MG	A204430	001	Oct 16, 2018
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RIMANTADINE HYDROCHLORIDE

SYRUP;ORAL

FLUMADINE

FOREST LABS

50MG/5ML

N019650 001 Sep 17, 1993

TABLET;ORAL

RIMANTADINE HYDROCHLORIDE

HERITAGE PHARMA AVET

100MG

A076375 001 Jan 14, 2003

IMPAX LABS INC

100MG

A075916 001 Nov 02, 2001

RIMEXOLONE

SUSPENSION/DROPS;OPHTHALMIC

VEXOL

EYEVANCE

1%

N020474 001 Dec 30, 1994

RISEDRONATE SODIUM

TABLET;ORAL

ACTONEL

+ APIL

75MG **

N020835 004 Apr 16, 2007

RISEDRONATE SODIUM

HANGZHOU BINJIANG

35MG

A207516 001 Feb 15, 2019

MYLAN

5MG

A200477 001 Nov 30, 2015

30MG

A200477 002 Nov 30, 2015

35MG

A200477 003 Nov 30, 2015

75MG

A200477 004 Jun 10, 2014

150MG

A200477 005 Jun 10, 2014

TABLET, DELAYED RELEASE;ORAL

RISEDRONATE SODIUM

IMPAX LABS INC

35MG

A205066 001 Jun 29, 2018

ZYDUS PHARMS

35MG

A203822 001 Sep 11, 2018

RISPERIDONE

SOLUTION;ORAL

RISPERIDONE

ANI PHARMS

1MG/ML

A076440 001 Jan 30, 2009

LANNETT CO INC

1MG/ML

A202386 001 Jan 12, 2015

PRECISION DOSE

1MG/ML

A076797 001 Jun 28, 2010

TORRENT

1MG/ML

A078909 001 Jul 29, 2009

WOCKHARDT

1MG/ML

A078744 001 Oct 08, 2009

TABLET;ORAL

RISPERDAL

JANSSEN PHARMS

5MG

N020272 005 Dec 29, 1993

RISPERIDONE

HERITAGE PHARMA AVET

0.25MG

A076228 001 Jun 30, 2008

0.25MG

A077769 001 Oct 16, 2008

0.5MG

A076228 002 Jun 30, 2008

0.5MG

A077769 002 Oct 16, 2008

1MG

A076228 003 Jun 30, 2008

1MG

A077769 003 Oct 16, 2008

2MG

A076228 004 Jun 30, 2008

2MG

A077769 004 Oct 16, 2008

3MG

A076228 005 Jun 30, 2008

3MG

A077769 005 Oct 16, 2008

4MG

A076228 006 Jun 30, 2008

4MG

A077769 006 Oct 16, 2008

JUBILANT CADISTA

0.25MG

A078828 001 Mar 23, 2009

0.5MG

A078828 002 Mar 23, 2009

1MG

A078828 003 Mar 23, 2009

2MG

A078828 004 Mar 23, 2009

3MG

A078828 005 Mar 23, 2009

4MG

A078828 006 Mar 23, 2009

MYLAN

0.25MG

A076288 001 Sep 15, 2008

0.5MG

A076288 002 Sep 15, 2008

1MG

A076288 003 Sep 15, 2008

2MG

A076288 004 Sep 15, 2008

3MG

A076288 005 Sep 15, 2008

4MG

A076288 006 Sep 15, 2008

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RISPERIDONE

TABLET; ORAL

RISPERIDONE

	4MG	A077784	006	Jun 08, 2010
SUN PHARM INDS INC	0.25MG	A078036	001	Mar 10, 2014
	0.5MG	A078036	002	Mar 10, 2014
	1MG	A078036	003	Mar 10, 2014
	2MG	A078036	004	Mar 10, 2014
	3MG	A078036	005	Mar 10, 2014
	4MG	A078036	006	Mar 10, 2014
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

RISPERDAL

+	JANSSEN PHARMS	0.5MG	N021444	001	Apr 02, 2003
+		1MG	N021444	002	Apr 02, 2003
+		2MG	N021444	003	Apr 02, 2003
+		3MG	N021444	004	Dec 23, 2004
+		4MG	N021444	005	Dec 23, 2004

RISPERIDONE

ACTAVIS LABS FL INC	0.5MG	A076996	001	Apr 19, 2011
	1MG	A076996	002	Apr 19, 2011
	2MG	A076996	003	Apr 19, 2011
	3MG	A076996	004	Apr 19, 2011
	4MG	A076996	005	Apr 19, 2011
HERITAGE PHARMA AVET	0.5MG	A076908	001	Mar 12, 2012
	1MG	A076908	002	Mar 12, 2012
	2MG	A076908	003	Mar 12, 2012
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	Sep 27, 1984

TABLET; ORAL

YUTOPAR

ASTRAZENECA	10MG	N018555	001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RITONAVIR

CAPSULE;ORAL

NORVIR

ABBOTT	100MG	N020680	001	Mar 01, 1996
+ ABBVIE	100MG **	N020945	001	Jun 29, 1999
RITONAVIR				
HIKMA	100MG	A205801	001	Dec 03, 2020

RIVASTIGMINE TARTRATE

CAPSULE;ORAL

EXELON

+ NOVARTIS	EQ 1.5MG BASE **	N020823	003	Apr 21, 2000
+	EQ 3MG BASE **	N020823	004	Apr 21, 2000
+	EQ 4.5MG BASE **	N020823	005	Apr 21, 2000
+	EQ 6MG BASE **	N020823	006	Apr 21, 2000

RIVASTIGMINE TARTRATE

APOTEX INC

EQ 1.5MG BASE	A091072	001	May 16, 2013
EQ 3MG BASE	A091072	002	May 16, 2013
EQ 4.5MG BASE	A091072	003	May 16, 2013
EQ 6MG BASE	A091072	004	May 16, 2013

SOLUTION;ORAL

EXELON

NOVARTIS	EQ 2MG BASE/ML	N021025	001	Apr 21, 2000
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RIZATRIPTAN BENZOATE

TABLET;ORAL

MAXALT

+ MERCK	EQ 5MG BASE **	N020864	001	Jun 29, 1998
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RIZATRIPTAN BENZOATE

APOTEX INC

EQ 5MG BASE	A202244	001	Dec 31, 2012
EQ 10MG BASE	A202244	002	Dec 31, 2012

EMCURE PHARMS LTD

EQ 5MG BASE	A204090	001	Nov 26, 2013
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EQ 10MG BASE	A204090	002	Nov 26, 2013
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UNICHEM

EQ 5MG BASE	A207836	001	Mar 07, 2017
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EQ 10MG BASE	A207836	002	Mar 07, 2017
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TABLET, ORALLY DISINTEGRATING;ORAL

MAXALT-MLT

+ MERCK	EQ 5MG BASE **	N020865	001	Jun 29, 1998
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RIZATRIPTAN BENZOATE

APOTEX INC

EQ 5MG BASE	A202477	001	Jul 01, 2013
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EQ 10MG BASE	A202477	002	Jul 01, 2013
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JUBILANT GENERICS

EQ 5MG BASE	A203334	001	Oct 16, 2015
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EQ 10MG BASE	A203334	002	Oct 16, 2015
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MYLAN PHARMS INC

EQ 5MG BASE	A078173	001	Dec 31, 2012
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EQ 10MG BASE	A078173	002	Dec 31, 2012
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ROCURONIUM BROMIDE

INJECTABLE;INJECTION

ROCURONIUM BROMIDE

TEVA PHARMS

50MG/5ML (10MG/ML)	A078717	001	Nov 26, 2008
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100MG/10ML (10MG/ML)	A078717	002	Nov 26, 2008
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ZEMURON

+ ORGANON USA INC	50MG/5ML (10MG/ML) **	N020214	001	Mar 17, 1994
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+	10MG/ML (10MG/ML) **	N020214	002	Mar 17, 1994
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+	100MG/10ML (10MG/ML) **	N020214	003	Mar 17, 1994
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ROFECOXIB

SUSPENSION;ORAL

VIOXX

MERCK

12.5MG/5ML	N021052	001	May 20, 1999
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25MG/5ML	N021052	002	May 20, 1999
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TABLET;ORAL

VIOXX

MERCK

12.5MG	N021042	001	May 20, 1999
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25MG	N021042	002	May 20, 1999
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50MG	N021042	003	Feb 25, 2000
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ROFLUMILAST

TABLET; ORAL

ROFLUMILAST

BRECKENRIDGE	500MCG	A208236	001	Oct 03, 2018
MICRO LABS	500MCG	A208180	001	Mar 22, 2019
MYLAN	500MCG	A208257	001	Jul 13, 2018

ROLAPITANT HYDROCHLORIDE

EMULSION; INTRAVENOUS

VARUBI

+ TERSERA	EQ 166.5MG BASE/92.5ML (EQ 1.8MG BASE/ML)	N208399	001	Oct 25, 2017
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ROMIDEPSIN

SOLUTION; INTRAVENOUS

ROMIDEPSIN

+ TEVA PHARMS USA INC	10MG/2ML (5MG/ML)	N208574	001	Mar 13, 2020
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ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

REQUIP

+ GLAXOSMITHKLINE LLC	EQ 0.25MG BASE	N020658	001	Sep 19, 1997
+	EQ 0.5MG BASE	N020658	002	Sep 19, 1997
+	EQ 1MG BASE	N020658	003	Sep 19, 1997
+	EQ 2MG BASE	N020658	004	Sep 19, 1997
+	EQ 3MG BASE	N020658	006	Jan 27, 1999
+	EQ 4MG BASE	N020658	007	Jan 27, 1999
+	EQ 5MG BASE	N020658	005	Sep 19, 1997

ROPINIROLE HYDROCHLORIDE

COSETTE

	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
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
HIKMA	EQ 0.25MG BASE	A077852	001	May 05, 2008
	EQ 0.5MG BASE	A077852	002	May 05, 2008
	EQ 1MG BASE	A077852	003	May 05, 2008
	EQ 2MG BASE	A077852	004	May 05, 2008
	EQ 3MG BASE	A077852	005	May 05, 2008
	EQ 4MG BASE	A077852	006	May 05, 2008
	EQ 5MG BASE	A077852	007	May 19, 2008

TABLET, EXTENDED RELEASE; ORAL

REQUIP XL

+ GLAXOSMITHKLINE LLC	EQ 2MG BASE	N022008	001	Jun 13, 2008
+	EQ 3MG BASE **	N022008	002	Jun 13, 2008
+	EQ 4MG BASE	N022008	003	Jun 13, 2008
+	EQ 6MG BASE	N022008	006	Apr 10, 2009
+	EQ 8MG BASE	N022008	004	Jun 13, 2008
+	EQ 12MG BASE	N022008	005	Oct 31, 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

WATSON LABS INC

	EQ 2MG BASE	A200431	001	Jun 06, 2012
	EQ 4MG BASE	A200431	002	Jun 06, 2012
	EQ 6MG BASE	A200431	003	Jun 06, 2012
	EQ 8MG BASE	A200431	004	Jun 06, 2012
	EQ 12MG BASE	A200431	005	Jun 06, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

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

ROPIVACAINE HYDROCHLORIDE

RISING PHARMA

		40MG/20ML (2MG/ML)	A090318	001	Sep 23, 2014
		150MG/30ML (5MG/ML)	A090318	002	Sep 23, 2014
		150MG/20ML (7.5MG/ML)	A090318	003	Sep 23, 2014
		200MG/20ML (10MG/ML)	A090318	004	Sep 23, 2014

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	
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ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDIA

+ WOODWARD

		EQ 8MG BASE **	N021071	004	May 25, 1999
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ROSIGLITAZONE MALEATE

ANI PHARMS

		EQ 2MG BASE	A076747	001	Jan 25, 2013
		EQ 4MG BASE	A076747	002	Jan 25, 2013
		EQ 8MG BASE	A076747	003	Jan 25, 2013

ROSUVASTATIN CALCIUM

TABLET; ORAL

ROSUVASTATIN CALCIUM

AMNEAL PHARMS CO

		EQ 5MG BASE	A208850	001	Oct 16, 2018
		EQ 10MG BASE	A208850	002	Oct 16, 2018
		EQ 20MG BASE	A208850	003	Oct 16, 2018
		EQ 40MG BASE	A208850	004	Oct 16, 2018

APOTEX INC

		EQ 5MG BASE	A079145	001	Jul 19, 2016
		EQ 10MG BASE	A079145	002	Jul 19, 2016
		EQ 20MG BASE	A079145	003	Jul 19, 2016
		EQ 40MG BASE	A079145	004	Jul 19, 2016

INVENTIA

		EQ 5MG BASE	A207653	001	Feb 05, 2021
		EQ 10MG BASE	A207653	002	Feb 05, 2021
		EQ 20MG BASE	A207653	003	Feb 05, 2021
		EQ 40MG BASE	A207653	004	Feb 05, 2021

MYLAN

		EQ 5MG BASE	A079161	001	Jul 19, 2016
		EQ 10MG BASE	A079161	002	Jul 19, 2016
		EQ 20MG BASE	A079161	003	Jul 19, 2016
		EQ 40MG BASE	A079161	004	Jul 19, 2016

SCIEGEN PHARMS INC

		EQ 5MG BASE	A206381	001	Apr 24, 2019
		EQ 10MG BASE	A206381	002	Apr 24, 2019
		EQ 20MG BASE	A206381	003	Apr 24, 2019
		EQ 40MG BASE	A206381	004	Apr 24, 2019

SUNSHINE

		EQ 5MG BASE	A210667	001	Apr 01, 2020
		EQ 10MG BASE	A210667	002	Apr 01, 2020
		EQ 20MG BASE	A210667	003	Apr 01, 2020
		EQ 40MG BASE	A210667	004	Apr 01, 2020

TEVA PHARMS USA

		EQ 5MG BASE	A079166	001	Jul 19, 2016
		EQ 10MG BASE	A079166	002	Jul 19, 2016
		EQ 20MG BASE	A079166	003	Jul 19, 2016
		EQ 40MG BASE	A079166	004	Jul 19, 2016

UMEDICA LABS PVT LTD

		EQ 5MG BASE	A207626	001	Apr 09, 2019
		EQ 10MG BASE	A207626	002	Apr 09, 2019
		EQ 20MG BASE	A207626	003	Apr 09, 2019
		EQ 40MG BASE	A207626	004	Apr 09, 2019

ZYDUS PHARMS

		EQ 5MG BASE	A206513	001	Mar 01, 2019
		EQ 10MG BASE	A206513	002	Mar 01, 2019
		EQ 20MG BASE	A206513	003	Mar 01, 2019
		EQ 40MG BASE	A206513	004	Mar 01, 2019

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

TABLET; ORAL

INVIRASE

+ HOFFMANN-LA ROCHE

EQ 500MG BASE

N021785 001 Dec 17, 2004

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

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

SECONAL SODIUM

VALEANT PHARMS NORTH

50MG

A086101 001 Oct 03, 1983

100MG

A086101 002 Oct 03, 1983

INJECTABLE; INJECTION

SECOBARBITAL SODIUM

ELKINS SINN

100MG/VIAL

A083281 001

WYETH AYERST

50MG/ML

A083262 001

SECONAL SODIUM

LILLY

50MG/ML **

N007392 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SECOBARBITAL SODIUMSUPPOSITORY;RECTAL
SECONAL SODIUM

LILLY	30MG	A086530 001
	60MG	A086530 002
	120MG	A086530 003
	200MG	A086530 004

SECRETININJECTABLE; INJECTION
SECRETIN-FERRING

FERRING	75CU/VIAL	N018290 001
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SECRETIN SYNTHETIC PORCINEFOR SOLUTION; INTRAVENOUS
SECREFLO

CHIRHOCLIN	16MCG/VIAL	N021136 001	Apr 04, 2002
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SELEGILINE HYDROCHLORIDE

CAPSULE; ORAL

ELDEPRYL

+ SOMERSET	5MG **	N020647 001	May 15, 1996
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SELEGILINE HYDROCHLORIDE

LANNETT CO INC	5MG	A075145 001	Sep 15, 2003
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TABLET; ORAL

SELEGILINE HYDROCHLORIDE

CHARTWELL MOLECULES	5MG	A074565 001	Aug 02, 1996
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	5MG	A074641 001	Aug 02, 1996
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COSETTE	5MG	A074744 001	Jan 27, 1997
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	5MG	A074756 001	Nov 25, 1998
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G AND W LABS INC	5MG	A074537 001	Aug 02, 1996
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MYLAN	5MG	A074866 001	Nov 26, 1997
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+ SOMERSET	5MG **	N019334 001	Jun 05, 1989
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SELENIUM SULFIDE

LOTION/SHAMPOO; TOPICAL

EXSEL

ALLERGAN HERBERT	2.5%	A083892 001
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SELENIUM SULFIDE

ACTAVIS MID ATLANTIC	2.5%	A084394 001
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COSETTE	2.5%	A086209 001
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IVAX PHARMS	2.5%	A085777 001
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SELSUN

+ CHATTEM	2.5%	N007936 001
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SELENOMETHIONINE SE-75

INJECTABLE; INJECTION

SELENOMETHIONINE SE 75

GE HEALTHCARE	250uCi/ML	N017257 001
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MALLINCKRODT	100uCi/ML	N017098 001
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PHARMALUCENCE	500uCi/ML	N017322 001
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SETHOTOPE

BRACCO	85-550uCi/ML	N017047 001
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SERMORELIN ACETATE

INJECTABLE; INJECTION

GEREF

+ EMD SERONO	EQ 0.05MG BASE/AMP **	N019863 001	Dec 28, 1990
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+ EMD SERONO INC	EQ 0.5MG BASE/VIAL **	N020443 001	Sep 26, 1997
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+	EQ 1MG BASE/VIAL **	N020443 002	Sep 26, 1997
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SERTRALINE HYDROCHLORIDE

CONCENTRATE; ORAL

SERTRALINE HYDROCHLORIDE

RANBAXY LABS LTD	EQ 20MG BASE/ML	A078053 001	Feb 05, 2007
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TABLET; ORAL

SERTRALINE HYDROCHLORIDE

ANDA REPOSITORY	EQ 25MG BASE	A077818 001	Feb 06, 2007
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	EQ 50MG BASE	A077818 002	Feb 06, 2007
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	EQ 100MG BASE	A077818 003	Feb 06, 2007
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CADILA	EQ 25MG BASE	A077106 001	Feb 06, 2007
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	EQ 50MG BASE	A077106 002	Feb 06, 2007
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	EQ 100MG BASE	A077106 003	Feb 06, 2007
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SERTRALINE HYDROCHLORIDE

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

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
HERITAGE PHARMA AVET	EQ 25MG BASE	A076465 001	Aug 11, 2006
	EQ 25MG BASE	A077299 001	Feb 06, 2007
	EQ 25MG BASE	A077345 001	Feb 06, 2007
	EQ 25MG BASE	A077663 001	Feb 06, 2007
	EQ 50MG BASE	A076465 002	Aug 11, 2006
	EQ 50MG BASE	A077299 002	Feb 06, 2007
	EQ 50MG BASE	A077345 002	Feb 06, 2007
	EQ 50MG BASE	A077663 002	Feb 06, 2007
	EQ 100MG BASE	A076465 003	Aug 11, 2006
	EQ 100MG BASE	A077299 003	Feb 06, 2007
	EQ 100MG BASE	A077345 003	Feb 06, 2007
	EQ 100MG BASE	A077663 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 25MG BASE	A078626 001	Jan 31, 2008
	EQ 50MG BASE	A076540 002	Mar 20, 2007
	EQ 50MG BASE	A078626 002	Jan 31, 2008
	EQ 100MG BASE	A076540 003	Mar 20, 2007
	EQ 100MG BASE	A078626 003	Jan 31, 2008
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
SUN PHARM INDS LTD	EQ 25MG BASE	A077977 001	Feb 06, 2007
	EQ 50MG BASE	A077977 002	Feb 06, 2007
	EQ 100MG BASE	A077977 003	Feb 06, 2007
	EQ 150MG BASE	A077977 004	Feb 06, 2007
	EQ 200MG BASE	A077977 005	Feb 06, 2007
TORRENT PHARMS	EQ 25MG BASE	A077765 001	Feb 06, 2007
	EQ 50MG BASE	A077765 002	Feb 06, 2007
	EQ 100MG BASE	A077765 003	Feb 06, 2007
ZOLOFT			
+ UPJOHN	EQ 150MG BASE **	N019839 003	Dec 30, 1991
+	EQ 200MG BASE **	N019839 004	Dec 30, 1991

SEVELAMER CARBONATE

TABLET; ORAL

SEVELAMER CARBONATE

IMPAX LABS INC	800MG	A090975 001	Oct 23, 2017
MYLAN	800MG	A201069 001	Aug 05, 2020

SEVELAMER HYDROCHLORIDE

CAPSULE; ORAL

RENAGEL

GENZYME	403MG	N020926 001	Oct 30, 1998
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TABLET; ORAL

SEVELAMER HYDROCHLORIDE

RISING PHARMA	400MG	A201068 001	Dec 14, 2020
	800MG	A201068 002	Dec 14, 2020

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SIBUTRAMINE HYDROCHLORIDE

CAPSULE;ORAL

MERIDIA

ABBOTT

5MG

N020632 001 Nov 22, 1997

10MG

N020632 002 Nov 22, 1997

15MG

N020632 003 Nov 22, 1997

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

PERRIGO R AND D

EQ 25MG BASE

A205791 001 Apr 23, 2020

EQ 50MG BASE

A205791 002 Apr 23, 2020

WATSON LABS INC

EQ 25MG BASE

A202506 001 Nov 25, 2020

EQ 50MG BASE

A202506 002 Nov 25, 2020

EQ 100MG BASE

A202506 003 Nov 25, 2020

SILODOSIN

CAPSULE;ORAL

SILODOSIN

ALEMBIC PHARMS LTD

4MG

A211731 001 Nov 22, 2019

8MG

A211731 002 Nov 22, 2019

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

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

ZOCOR

+ ORGANON

80MG

N019766 005 Jul 10, 1998

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SIMVASTATINTABLET, ORALLY DISINTEGRATING;ORAL
SIMVASTATIN

80MG

N021961 004 Oct 09, 2007

SIMVASTATIN; SITAGLIPTIN PHOSPHATETABLET;ORAL
JUVISYNC

+	MERCK SHARP DOHME	10MG;EQ 50MG BASE **
+		10MG;EQ 100MG BASE **
+		20MG;EQ 50MG BASE **
+		20MG;EQ 100MG BASE **
+		40MG;EQ 50MG BASE **
+		40MG;EQ 100MG BASE **

N202343	004	Sep 18, 2012
N202343	001	Oct 07, 2011
N202343	005	Sep 18, 2012
N202343	002	Oct 07, 2011
N202343	006	Sep 18, 2012
N202343	003	Oct 07, 2011

SIROLIMUSTABLET;ORAL
RAPAMUNE

+ PF PRISM CV 5MG **

N021110 003 Feb 23, 2004

SODIUM BENZOATE; SODIUM PHENYLACETATESOLUTION;ORAL
UCEPHAN

B BRAUN 100MG/ML;100MG/ML

N019530 001 Dec 23, 1987

SODIUM BICARBONATEINJECTABLE; INJECTION
SODIUM BICARBONATEHOSPIRA 0.9MEQ/ML
1MEQ/MLA077394 001 Nov 09, 2005
A077394 002 Nov 09, 2005SODIUM BICARBONATE IN PLASTIC CONTAINER
+ ABBOTT 0.9MEQ/ML **
+ 1MEQ/ML **N019443 001 Jun 03, 1986
N019443 002 Jun 03, 1986SODIUM BICARBONATE; TARTARIC ACIDGRANULE, EFFERVESCENT;ORAL
BAROS

MALLINCKRODT INC 460MG/GM;420MG/GM

N018509 001 Aug 07, 1985

SODIUM CHLORIDEAEROSOL, METERED; INHALATION
BRONCHO SALINE

+ BLAIREX 0.9%

N019912 001 Sep 03, 1992

INJECTABLE; INJECTION

BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
ABRAXIS PHARM 9MG/ML

A088909 001 Feb 07, 1985

SODIUM CHLORIDE

ABBOTT 20GM/100ML
B BRAUN 20GM/100MLN017013 001
N017038 001

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN 450MG/100ML
MILES 450MG/100MLN018184 001
N018503 001

SODIUM CHLORIDE 0.9%

+	MEDEFIL INC	9MG/ML (9MG/ML)
+		18MG/2ML (9MG/ML)
+		22.5MG/2.5ML (9MG/ML)
+		27MG/3ML (9MG/ML)
+		45MG/5ML (9MG/ML)

N202832	001	Jan 06, 2012
N202832	002	Jan 06, 2012
N202832	003	Jan 06, 2012
N202832	004	Jan 06, 2012
N202832	005	Jan 06, 2012

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

	ABBOTT	9MG/ML
+	ICU MEDICAL INC	9MG/ML
	JUBILANT CADISTA	9MG/ML
+	LIEBEL-FLARSHEIM	405MG/50ML (9MG/ML)
	MILES	900MG/100ML

N019218	001	Jul 13, 1984
N019217	001	Jul 13, 1984
A203352	001	May 18, 2016
N021569	001	Jul 27, 2006
N018502	001	

SODIUM CHLORIDE 23.4% IN PLASTIC CONTAINER

+ ABRAXIS PHARM 234MG/ML **

N019329 001 Apr 22, 1987

SOLUTION;INTRAVENOUS

SODIUM CHLORIDE 14.6%

+ HOSPIRA 50MEQ/20ML (2.5MEQ/ML) **

N018897 001 Jul 20, 1984

SOLUTION;IRRIGATION

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

BAXTER HLTHCARE 450MG/100ML

N017864 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SODIUM CHLORIDE

SOLUTION;IRRIGATION

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

	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

CURIUM	100uCi/ML	N016708 001	
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SODIUM FLUORIDE F-18

INJECTABLE;INTRAVENOUS

FLUORINE F-18

+ GE HEALTHCARE	2mCi/ML **	N017042 001	
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SODIUM FLUORIDE F 18

NIH NCI DCTD	10-200mCi/ML **	N022494 001	Jan 26, 2011
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SODIUM FLUORIDE F-18

DECATUR	10-200mCi/ML	A204464 001	Oct 21, 2014
UCSF RODIOPHARM	10-200mCi/ML	A204437 001	Mar 13, 2014
UIHC PET IMAGING	10-200mCi/ML	A204462 001	Nov 17, 2015
UNIV TX MD ANDERSON	10-200mCi/ML	A203247 001	Dec 23, 2013

SODIUM FLUORIDE; TRICLOSAN

PASTE;DENTAL

COLGATE TOTAL

+ COLGATE PALMOLIVE	0.24%;0.3%	N020231 001	Jul 11, 1997
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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	
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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	

CURIUM	0.8-100mCi	N016515 002	
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+ CURIUM	0.8-100mCi	N016517 001	
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	15-100uCi	N016517 002	
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JUBILANT	2-200mCi	N021305 004	Nov 18, 2004
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SOLUTION;ORAL

HICON

JUBILANT	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	
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SODIUM IODIDE I 131

CIS	50mCi/ML	N017315 001	
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+ CURIUM	3.5-150mCi/VIAL	N016515 001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
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SODIUM LACTATE IN PLASTIC CONTAINER

+ HOSPIRA	5MEQ/ML	N018947	001	Sep 05, 1984
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SODIUM MONOFLUOROPHOSPHATE

GEL; DENTAL

EXTRA-STRENGTH AIM

CHESEBROUGH PONDS	1.2%	N019518	002	Aug 06, 1986
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PASTE; DENTAL

EXTRA-STRENGTH AIM

CHESEBROUGH PONDS	1.2%	N019518	001	Jun 03, 1987
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SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION

NIPRIDE

ROCHE	50MG/VIAL	N017546	001
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NITROPRESS

ABBOTT	50MG/VIAL	A071555	001	Nov 16, 1987
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+ ABBVIE	50MG/VIAL **	N018450	001
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HOSPIRA	50MG/VIAL	A070566	001	Jun 09, 1986
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SODIUM NITROPRUSSIDE

ABRAXIS PHARM	50MG/VIAL	A070031	001	Jan 17, 1985
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AMPHASTAR PHARMS INC	25MG/ML	A209832	001	Dec 18, 2017
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+ BAXTER HLTHCARE	50MG/VIAL **	N018581	001	Jul 28, 1982
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CIPLA	25MG/ML	A210855	001	Jul 16, 2018
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EUGIA PHARMA	25MG/ML	A211934	001	Dec 10, 2020
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SUN PHARM	25MG/ML	A210467	001	Nov 26, 2018
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TEVA PARENTERAL	25MG/ML	A073465	001	Mar 30, 1992
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VIRTUS PHARM	25MG/ML	A209834	001	Jun 26, 2018
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SOLUTION; INTRAVENOUS

NIPRIDE RTU IN SODIUM CHLORIDE 0.9%

+ EXELA PHARMA	10MG/50ML (0.2MG/ML) **	N209387	002	Dec 07, 2017
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SODIUM OXYBATE

SOLUTION; ORAL

SODIUM OXYBATE

HIKMA	0.5GM/ML	A202090	001	Jan 17, 2017
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SODIUM PHENYL BUTYRATE

TABLET; ORAL

SODIUM PHENYL BUTYRATE

ALVOGEN	500MG	A090910	001	Nov 18, 2011
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SODIUM PHOSPHATE P-32

SOLUTION; INJECTION, ORAL

PHOSPHOTOPE

BRACCO	1-8mCi/VIAL	N010927	001
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SODIUM PHOSPHATE P 32

MALLINCKRODT	0.67mCi/ML	N011777	001
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	1.5mCi/VIAL	N011777	002
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SODIUM PHOSPHATE, DIBASIC ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL

VISICOL

SALIX PHARMS	0.398GM; 1.102GM	N021097	001	Sep 21, 2000
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SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

KAYEXALATE

+ CONCORDIA	453.6GM/BOT **	N011287	001
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SODIUM POLYSTYRENE SULFONATE

CITRUSPHARMA	454GM/BOT	A040909	001	Dec 03, 2008
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+ WOCKHARDT	453.6GM/BOT	A088786	001	Sep 11, 1984
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SODIUM POLYSTYRENE SULFONATE

SUSPENSION;ORAL, RECTAL

KIONEX

ANI PHARMS 15GM/60ML A040028 001 Sep 17, 2007

SODIUM POLYSTYRENE SULFONATE

ANI PHARMS 15GM/60ML A090590 001 May 13, 2011

HIKMA 15GM/60ML A089049 001 Nov 17, 1986

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

SOLIFENACIN SUCCINATE

TABLET;ORAL

SOLIFENACIN SUCCINATE

AJANTA PHARMA LTD 5MG A205483 001 May 20, 2019

10MG A205483 002 May 20, 2019

BRECKENRIDGE 5MG A209818 001 May 20, 2019

10MG A209818 002 May 20, 2019

SUNSHINE 5MG A213346 001 Apr 13, 2020

10MG A213346 002 Apr 13, 2020

ZYDUS PHARMS 5MG A207721 001 Oct 19, 2020

10MG A207721 002 Oct 19, 2020

SORAFENIB TOSYLATE

TABLET;ORAL

SORAFENIB TOSYLATE

MYLAN EQ 200MG BASE A207012 001 Sep 10, 2020

TEVA PHARMS USA INC EQ 200MG BASE A209567 001 Nov 12, 2020

SORBITOL

SOLUTION;IRRIGATION

SORBITOL 3% IN PLASTIC CONTAINER

BAXTER HLTHCARE 3GM/100ML N018512 001 May 27, 1982

SOTALOL HYDROCHLORIDE

TABLET;ORAL

BETAPACE

COVIS 320MG N019865 004 Oct 30, 1992

BETAPACE AF

COVIS 40MG N021151 006 Apr 02, 2003

60MG N021151 007 Apr 02, 2003

100MG N021151 005 Mar 14, 2003

SOTALOL HYDROCHLORIDE

AUROBINDO PHARMA USA 80MG A077616 001 Feb 07, 2007

120MG A077616 002 Feb 07, 2007

160MG A077616 003 Feb 07, 2007

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SOTALOL HYDROCHLORIDE

TABLET; ORAL

SOTALOL HYDROCHLORIDE

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
UPSHER SMITH LABS	80MG	A075366 001	May 01, 2000
	120MG	A075366 002	May 01, 2000
	160MG	A075366 003	May 01, 2000
	240MG	A075366 004	May 01, 2000
WATSON LABS	80MG	A075238 001	Jul 13, 2000
	120MG	A075238 002	Jul 13, 2000
	160MG	A075238 003	Jul 13, 2000
	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
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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

ACTAVIS ELIZABETH	25MG	A040353 003	Mar 15, 2006
	50MG	A040353 001	Jul 29, 1999
	100MG	A040353 002	Jul 29, 1999
ASCOT	25MG	A087687 001	Oct 20, 1982
CHARTWELL RX	25MG	A086809 001	
IVAX PHARMS	25MG	A087108 001	
LEDERLE	25MG	A087634 001	
MUTUAL PHARM	25MG	A087265 001	
MYLAN	25MG	A087086 001	
OXFORD PHARMS	25MG	A040750 001	Aug 29, 2006
	50MG	A040750 002	Aug 29, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SPIRONOLACTONE

TABLET;ORAL

SPIRONOLACTONE

	100MG	A040750 003	Aug 29, 2006
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	

STANZOLOL

TABLET;ORAL

WINSTROL

+ LUNDBECK INC

2MG

N012885 001 May 14, 1984

STAVUDINE

CAPSULE;ORAL

STAVUDINE

AUROBINDO PHARMA	15MG	A077672 003	Dec 29, 2008
	20MG	A077672 004	Dec 29, 2008
	30MG	A077672 001	Dec 29, 2008
	40MG	A077672 002	Dec 29, 2008
HETERO LABS LTD III	15MG	A078957 001	Dec 29, 2008
	20MG	A078957 002	Dec 29, 2008
	30MG	A078957 003	Dec 29, 2008
	40MG	A078957 004	Dec 29, 2008
MYLAN	15MG	A079069 001	Dec 29, 2008
	20MG	A079069 002	Dec 29, 2008
	30MG	A079069 003	Dec 29, 2008
	40MG	A079069 004	Dec 29, 2008
MYLAN LABS LTD	30MG	A078775 001	Jan 05, 2009
	40MG	A078775 002	Jan 05, 2009

ZERIT

BRISTOL-MYERS SQUIBB	5MG	N020412 001	Jun 24, 1994
+	15MG	N020412 002	Jun 24, 1994
+	20MG	N020412 003	Jun 24, 1994
+	30MG	N020412 004	Jun 24, 1994
+	40MG	N020412 005	Jun 24, 1994

CAPSULE, EXTENDED RELEASE;ORAL

ZERIT XR

BRISTOL 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, 2002

FOR SOLUTION;ORAL

STAVUDINE

AUROBINDO PHARMA	1MG/ML	A077774 001	Dec 29, 2008
CIPLA LTD	1MG/ML	A078030 001	Mar 20, 2009

ZERIT

+	BRISTOL-MYERS SQUIBB	1MG/ML **	N020413 001	Sep 06, 1996
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STERILE 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

HIKMA	100% (20ML)	A206369 002	Sep 02, 2015	
+	HOSPIRA	100% (1ML)	N018801 001	Oct 27, 1982
NEPHRON	100% (5ML)	A211222 001	Feb 10, 2021	

STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

B BRAUN	100%	N019077 001	Mar 02, 1984
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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 002

SUCCINYLCHOLINE 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

AMPHASTAR PHARMS INC 20MG/ML A213432 001 Jun 08, 2020

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 003

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC

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

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

BARR 500MG A087189 001 Jul 25, 1983

HEATHER 500MG A086163 001

RISING 500MG A085844 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFAMETHOXAZOLE

TABLET; ORAL

SULFAMETHOXAZOLE

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

HIKMA

80MG/ML; 16MG/ML

A070627 001 Dec 29, 1987

80MG/ML; 16MG/ML

A070628 001 Dec 29, 1987

HOSPIRA

80MG/ML; 16MG/ML

A073199 001 Sep 11, 1992

WATSON LABS

80MG/ML; 16MG/ML

A071556 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

200MG/5ML; 40MG/5ML

A070028 001 Jun 02, 1987

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM

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
HERITAGE PHARMA AVET	800MG;160MG	A070037	001	Jun 02, 1987
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

SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH

HERITAGE PHARMA AVET	400MG;80MG	A070030	001	Jun 02, 1987
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SULFATRIM-DS

SUPERPHARM	800MG;160MG	A070066	001	Jun 24, 1985
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SULFATRIM-SS

SUPERPHARM	400MG;80MG	A070065	002	Jun 24, 1985
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UROPLUS DS

SHIONOGI	800MG;160MG	A071816	001	Sep 28, 1987
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UROPLUS SS

SHIONOGI	400MG;80MG	A071815	001	Sep 28, 1987
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SULFANILAMIDE

CREAM; VAGINAL

AVC

+ MYLAN SPECIALITY LP	15%	N006530	003	Jan 27, 1987
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SULFANILAMIDE

COSETTE	15%	A088718	001	Sep 19, 1985
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SUPPOSITORY; VAGINAL

AVC

MYLAN SPECIALITY LP	1.05GM	N006530	004	Jan 27, 1987
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SULFAPHENAZOLE

SUSPENSION; ORAL

SULFABID

PHARM RES ASSOC	500MG/5ML	N013093	001	
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TABLET; ORAL

SULFABID

PURDUE FREDERICK	500MG	N013092	002	
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SULFAPYRIDINE

TABLET; ORAL

SULFAPYRIDINE

LILLY	500MG	N000159	001	
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SULFASALAZINE

SUSPENSION; ORAL

AZULFIDINE

PHARMACIA AND UPJOHN	250MG/5ML	N018605	001	
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TABLET; ORAL

S.A.S.-500

SOLVAY	500MG	A083450	001	
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SULFASALAZINE

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
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SULFINPYRAZONE

CAPSULE; ORAL

ANTURANE

+ NOVARTIS	200MG **	N011556	004	
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SULFINPYRAZONE

BARR	200MG	A087666	001	Sep 17, 1982
IVAX PHARMS	200MG	A087770	001	Nov 19, 1982
PAR PHARM	200MG	A088934	001	Sep 06, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFINPYRAZONE

CAPSULE; ORAL

SULFINPYRAZONE

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

500MG

A080142 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

RISING

500MG

A085628 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

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

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

MYLAN

150MG

A073039 002 Jun 22, 1993

200MG

A073039 001 Jun 22, 1993

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

MYLAN ASI

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A090314 001 Jun 10, 2010

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A090641 001 Jul 28, 2010

MYLAN LABS LTD

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A203322 001 Apr 14, 2014

PAR STERILE PRODUCTS

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A077871 001 Jul 09, 2009

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

TEVA PHARMS USA

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A077907 001 Feb 06, 2009

ZYDUS

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A090310 001 Aug 11, 2010

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SUPROFEN

SOLUTION/DROPS;OPHTHALMIC

PROFENAL

ALCON

1%

N019387 001 Dec 23, 1988

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

CONCORD BIOTECH LTD

EQ 0.5MG BASE

A213112 001 Nov 10, 2020

HERITAGE PHARMA AVET

EQ 5MG BASE

A090402 001 Jul 01, 2010

TADALAFIL

TABLET;ORAL

TADALAFIL

GLENMARK PHARMS LTD

2.5MG

A210716 001 Dec 29, 2020

5MG

A210716 002 Dec 29, 2020

10MG

A210716 003 Dec 29, 2020

20MG

A210716 004 Dec 29, 2020

MYLAN

5MG

A206957 001 Apr 29, 2019

20MG

A200630 001 Aug 03, 2018

RISING PHARMA

2.5MG

A206956 001 Apr 29, 2019

10MG

A206956 002 Apr 29, 2019

20MG

A206956 003 Apr 29, 2019

WATSON LABS INC

2.5MG

A205885 001 Mar 29, 2019

5MG

A205885 002 Mar 29, 2019

10MG

A205885 003 Mar 29, 2019

20MG

A205885 004 Mar 29, 2019

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

TAMSULOSIN HYDROCHLORIDE

CAPSULE;ORAL

TAMSULOSIN HYDROCHLORIDE

ANCHEN PHARMS

0.4MG

A202010 001 Jan 04, 2013

MYLAN

0.4MG

A090408 001 Apr 27, 2010

TAPENTADOL HYDROCHLORIDE

SOLUTION;ORAL

NUCYNTA

+ COLLEGIUM PHARM INC

EQ 20MG BASE/ML

N203794 001 Oct 15, 2012

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TAVABOROLESOLUTION; TOPICAL
TAVABOROLE

IDENTIRX 5% A211963 001 Feb 03, 2021

TAZAROTENECREAM; TOPICAL
TAZAROTENE

FOUGERA PHARMS INC 0.1% A211175 001 Jan 28, 2019

TECHNETIUM TC-99M APCITIDEINJECTABLE; INJECTION
ACUTECT

CIS BIO INTL SA N/A N020887 001 Sep 14, 1998

TECHNETIUM TC-99M DEPREOTIDEINJECTABLE; INJECTION
NEO TECT KIT

CIS BIO INTL SA N/A ** N021012 001 Aug 03, 1999

TECHNETIUM TC-99M DISOFENIN KITINJECTABLE; INJECTION
HEPATOLITE

SUN PHARM INDS INC N/A N018467 001 Mar 16, 1982

TECHNETIUM TC-99M ETIDRONATE KITINJECTABLE; 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 FERSENTETATE KITINJECTABLE; INJECTION
RENOTEC

BRACCO N/A N017045 001

TECHNETIUM TC-99M GLUCEPTATE KITINJECTABLE; INJECTION
GLUCOSCAN

BRISTOL MYERS SQUIBB N/A N017907 001

TECHNESCAN GLUCEPTATE
DRAXIMAGE N/A

N018272 001 Jan 27, 1982

TECHNETIUM TC-99M LIDOFENIN KITINJECTABLE; INJECTION
TECHNESCAN HIDA

DRAXIMAGE N/A N018489 001 Oct 31, 1986

TECHNETIUM TC-99M MEDRONATEINJECTABLE; INJECTION
DRAXIMAGE MDP-10

JUBILANT N/A N018035 001

TECHNETIUM TC-99M MEDRONATE KITINJECTABLE; INJECTION
AMERSCAN MDP KIT

GE HEALTHCARE N/A N018335 001 Aug 05, 1982

OSTEOLITE
PHARMALUCENCE N/A

N017972 001

TECHNETIUM TC 99M MPI MDP
GE HEALTHCARE N/A

N018141 001

N018141 002 Jun 12, 1989

TECHNETIUM TC-99M PENTETATE KITINJECTABLE; INJECTION
AN-DTPA

JUBILANT DRAXIMAGE N/A N017714 001

MPI DTPA KIT - CHELATE
GE HEALTHCARE N/A

N017255 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION

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

CURIUM 0.25-3 CI/GENERATOR

N017243 002

SOLUTION; INTRAVENOUS, ORAL

TECHNETIUM TC 99M GENERATOR

+ GE HEALTHCARE 68-2703mCi/GENERATOR

N017693 002 Dec 13, 2013

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

TECHNECOLL

MALLINCKRODT N/A

N017059 001

TECHNETIUM TC 99M TSC

GE HEALTHCARE N/A

N017784 001

TESULOID

BRACCO N/A

N016923 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TECHNETIUM TC-99M TEBOROXIME KIT

INJECTABLE; INJECTION

CARDIOTEC

BRACCO

N/A

N019928 001 Dec 19, 1990

TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE; INJECTION

MYOVIEW

+ GE HEALTHCARE

N/A

N020372 001 Feb 09, 1996

TEGASEROD MALEATE

TABLET; ORAL

ZELNORM

+ ALFASIGMA

EQ 2MG BASE

N021200 001 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

HISUN PHARM HANGZHOU

20MG

A207843 001 Feb 19, 2019

40MG

A207843 002 Feb 19, 2019

80MG

A207843 003 Feb 19, 2019

JUBILANT GENERICS

20MG

A204164 001 Aug 22, 2016

40MG

A204164 002 Aug 22, 2016

80MG

A204164 003 Aug 22, 2016

TORRENT

20MG

A203171 001 Jul 07, 2014

40MG

A203171 002 Jul 07, 2014

80MG

A203171 003 Jul 07, 2014

TEMAZEPAM

CAPSULE; ORAL

TEMAZ

QUANTUM PHARMICS

15MG

A070564 001 Oct 15, 1985

30MG

A070547 001 Oct 15, 1985

TEMAZEPAM

AUROBINDO PHARMA USA

7.5MG

A070920 002 May 21, 2010

15MG

A070920 004 Jul 07, 1986

22.5MG

A070920 003 Jun 12, 2009

30MG

A070920 001 Jul 10, 1986

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TEMOZOLOMIDE

CAPSULE; ORAL

TEMOZOLOMIDE

APOTEX INC	5MG	A204159 001	Jul 05, 2018
	20MG	A204159 002	Jul 05, 2018
	100MG	A204159 003	Jul 05, 2018
	140MG	A204159 004	Jul 05, 2018
	180MG	A204159 005	Jul 05, 2018
	250MG	A204159 006	Jul 05, 2018
HERITAGE	5MG	A078879 001	Mar 01, 2010
	20MG	A078879 002	Mar 01, 2010
	100MG	A078879 003	Mar 01, 2010
	140MG	A078879 005	Mar 01, 2010
	180MG	A078879 006	Mar 01, 2010
	250MG	A078879 004	Mar 01, 2010
LANNETT CO INC	5MG	A203898 001	Feb 10, 2016
	20MG	A203898 002	Feb 10, 2016
	100MG	A203898 003	Feb 10, 2016
	140MG	A203898 004	Feb 10, 2016
	180MG	A203898 005	Feb 10, 2016
	250MG	A203898 006	Feb 10, 2016
MYLAN	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
WATSON LABS TEVA	5MG	A203959 001	Apr 18, 2017
	20MG	A203959 002	Apr 18, 2017
	100MG	A203959 003	Apr 18, 2017
	140MG	A203959 004	Apr 18, 2017
	250MG	A203959 005	Apr 18, 2017

TENAPANOR HYDROCHLORIDE

TABLET; ORAL

IBSRELA

+ ARDELYX INC EQ 50MG BASE N211801 001 Sep 12, 2019

TENIPOSIDE

INJECTABLE; INJECTION

VUMON

+ HQ SPECLT PHARMA 10MG/ML N020119 001 Jul 14, 1992

TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

TENOFOVIR DISOPROXIL FUMARATE

CHARTWELL	300MG	A206481 001	Jul 26, 2018
MYLAN	150MG	A206569 001	Nov 27, 2018
	200MG	A206569 002	Nov 27, 2018
	250MG	A206569 003	Nov 27, 2018
	300MG	A206569 004	Nov 27, 2018

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

BEXIMCO PHARMS USA	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
HIKMA	EQ 1MG BASE	A075498 001	Apr 12, 2001
	EQ 2MG BASE	A075498 002	Apr 12, 2001
	EQ 5MG BASE	A075498 003	Apr 12, 2001
	EQ 10MG BASE	A075498 004	Apr 12, 2001
MYLAN	EQ 1MG BASE	A075140 002	Feb 11, 2000
	EQ 2MG BASE	A075140 003	Feb 11, 2000
	EQ 5MG BASE	A075140 001	Feb 11, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

TERAZOSIN HYDROCHLORIDE

	EQ 10MG BASE	A075140 004	Feb 11, 2000
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

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
TEVA	EQ 1MG BASE	A074446 001	May 18, 2000
	EQ 2MG BASE	A074446 002	May 18, 2000
	EQ 5MG BASE	A074446 003	May 18, 2000
	EQ 10MG BASE	A074446 004	May 18, 2000

TERBINAFINE

GEL; TOPICAL

LAMISIL

GLAXOSMITHKLINE CONS	1%	N020846 001	Apr 29, 1998
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TERBINAFINE HYDROCHLORIDE

CREAM; TOPICAL

LAMISIL

NOVARTIS	1% **	N020192 001	Dec 30, 1992
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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
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TABLET; ORAL

LAMISIL

+ NOVARTIS	EQ 250MG BASE **	N020539 001	May 10, 1996
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TERBINAFINE HYDROCHLORIDE

CHARTWELL	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
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BRICANYL

SANOFI AVENTIS US	0.2MG/INH	N018000 001	Mar 19, 1985
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TERBUTALINE SULFATE

INJECTABLE; INJECTION

BRETHINE

+ PHARMACARE

1MG/ML **

N018571 001

BRICANYL

SANOFI AVENTIS US

1MG/ML

N017466 001

TERBUTALINE SULFATE

DR REDDYS

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

TERIFLUNOMIDE

TABLET; ORAL

TERIFLUNOMIDE

AMNEAL PHARMS CO

7MG

A209613 001 Sep 28, 2018

14MG

A209613 002 Sep 28, 2018

APOTEX

7MG

A209601 001 Nov 02, 2018

14MG

A209601 002 Nov 02, 2018

AUROBINDO PHARMA LTD

7MG

A209638 001 Oct 26, 2018

14MG

A209638 002 Oct 26, 2018

BRECKENRIDGE

7MG

A209583 001 Sep 24, 2021

14MG

A209583 002 Sep 24, 2021

MSN

7MG

A209623 001 Apr 24, 2019

14MG

A209623 002 Apr 24, 2019

MYLAN

7MG

A209702 001 Feb 28, 2020

14MG

A209702 002 Feb 28, 2020

SOLA PHARMS

14MG

A209677 001 Jun 17, 2020

TEVA PHARMS USA

7MG

A209700 001 Sep 04, 2018

14MG

A209700 002 Sep 04, 2018

WATSON LABS TEVA

7MG

A209549 001 Jul 27, 2018

14MG

A209549 002 Jul 27, 2018

ZYDUS PHARMS

7MG

A209668 001 Nov 30, 2018

14MG

A209668 002 Nov 30, 2018

TERIPARATIDE

SOLUTION; SUBCUTANEOUS

FORTEO

LILLY

0.75MG/3ML (0.25MG/ML)

N021318 001 Nov 26, 2002

TERIPARATIDE ACETATE

INJECTABLE; INJECTION

PARATHAR

SANOFI AVENTIS US

200 UNITS/VIAL

N019498 001 Dec 23, 1987

TESTOLACTONE

INJECTABLE; INJECTION

TESLAC

BRISTOL MYERS SQUIBB

100MG/ML

N016119 001

TABLET; ORAL

TESLAC

BRISTOL MYERS SQUIBB

50MG

N016118 001

250MG

N016118 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TESTOSTERONE

FILM, EXTENDED RELEASE;TRANSDERMAL

ANDRODERM

ALLERGAN	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
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GEL;TRANSDERMAL

TESTOSTERONE

ANI PHARMS	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

ANDROGEL

+ ABBVIE	12.5MG/1.25GM ACTUATION **	N021015 003	Sep 26, 2003
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TESTOSTERONE

ALEOR DERMACEUTICALS	1.62% (20.25MG/1.25GM ACTUATION)	A213922 001	Mar 03, 2021
PERRIGO ISRAEL	12.5MG/1.25GM ACTUATION	N203098 001	Jan 31, 2013

INJECTABLE;INJECTION

TESTOSTERONE

DR REDDYS	100MG/ML	A086417 001	Jul 07, 1983
WATSON LABS	25MG/ML	A086420 001	May 10, 1983
	50MG/ML	A086419 001	Aug 23, 1983

SOLUTION, METERED;TRANSDERMAL

AXIRON

+ ELI LILLY AND CO	30MG/1.5ML ACTUATION **	N022504 001	Nov 23, 2010
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TESTOSTERONE

ALEOR DERMACEUTICALS	30MG/1.5ML ACTUATION	A212882 001	Jun 14, 2021
APOTEX	30MG/1.5ML ACTUATION	A209181 001	Nov 25, 2020

TABLET, EXTENDED RELEASE;BUCCAL

STRIANT

+ AUXILIUM PHARMS LLC	30MG	N021543 001	Jun 19, 2003
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TESTOSTERONE CYPIONATE

INJECTABLE;INJECTION

DEPO-TESTOSTERONE

PFIZER	50MG/ML	A085635 001	
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TESTOSTERONE CYPIONATE

RISING PHARMA	200MG/ML	A040652 001	Dec 11, 2006
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

RISING PHARMA	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	
	50MG/ML	A080188 002	
	50MG/ML	A085490 002	
	100MG/ML	A080188 003	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TESTOSTERONE PROPIONATE

INJECTABLE; INJECTION

TESTOSTERONE PROPIONATE

100MG/ML

A083595 003

TETRABENAZINE

TABLET; ORAL

TETRABENAZINE

AJANTA PHARMA LTD

12.5MG

A213621 001 Dec 04, 2020

25MG

A213621 002 Dec 04, 2020

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

250MG

A061471 001

ELKINS SINN

250MG

A060059 001

FERRANTE

125MG

A060173 001

250MG

A060173 002

HEATHER

250MG

A061148 001

500MG

A061148 002

HIKMA

250MG

A060768 001

500MG

A060768 002

IMPAX LABS

100MG

A060469 002

250MG

A060469 001

500MG

A060469 003

IVAX SUB TEVA PHARMS

250MG

A060704 001

500MG

A060704 002

MAST MM

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

TETRACYCLINE HYDROCHLORIDE

WARNER CHILCOTT	250MG	A062300 001
	500MG	A062300 002
WATSON LABS	250MG	A062103 001
	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
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FOR SOLUTION; TOPICAL

TOPICYCLINE

SHIRE	2.2MG/ML	N050493 001
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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
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SUSPENSION; ORAL

ACHROMYCIN V

LEDERLE	125MG/5ML	N050263 002
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SUMYCIN

PAR PHARM	125MG/5ML	A060400 001
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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
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TETRAMED

IVAX SUB TEVA PHARMS	125MG/5ML	A061468 001
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SUSPENSION/DROPS; OPHTHALMIC

ACHROMYCIN

STORZ	1%	N050268 001
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TABLET; ORAL

PANMYCIN

PHARMACIA AND UPJOHN	250MG	A061705 001
	500MG	A061705 002

SUMYCIN

STRIDES PHARMA	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
+ GE HEALTHCARE	1mCi/ML	N018110	002	Feb 27, 1996
TRACE LIFE	1mCi/ML	A075569	001	Nov 21, 2001

INJECTABLE; INTRAVENOUS

THALLOUS CHLORIDE TL 201

CURIUM	2mCi/ML	A077698	001	Nov 09, 2006
+ LANTHEUS MEDCL	2mCi/ML	N017806	002	Oct 09, 1998

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
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AEROLATE JR

FLEMING PHARMS	130MG	A085075	002	Nov 24, 1986
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AEROLATE SR

FLEMING PHARMS	260MG	A085075	001	Nov 24, 1986
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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
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THEOBID JR.

WHITBY	130MG	A087854	001	Mar 20, 1985
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THEOCLEAR L.A.-130

SCHWARZ PHARMA	130MG	A086569	001	May 27, 1982
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THEOCLEAR L.A.-260

SCHWARZ PHARMA	260MG	A086569	002	May 27, 1982
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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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

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				
	FERNDALE 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	100MG	A087400 003	Feb 21, 1985
		200MG	A087400 004	Feb 21, 1985
		300MG	A087400 002	Jan 11, 1983
THEOLAIR-SR				
	3M	200MG	A088369 001	Jul 16, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THEOPHYLLINETABLET, EXTENDED RELEASE;ORAL
THEOLAIR-SR

250MG	A086363	002	Jul 16, 1987
300MG	A088364	001	Jul 16, 1987
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

HERITAGE PHARMA AVET

100MG	A089807	001	Apr 30, 1990
200MG	A089808	001	Apr 30, 1990

INWOOD LABS

450MG	A040034	001	Apr 28, 1995
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UNI-DUR

SCHERING

400MG	A089822	001	Jan 04, 1995
600MG	A089823	001	Jan 04, 1995

THEOPHYLLINE SODIUM GLYCINATE

ELIXIR;ORAL

SYNOPHYLATE

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

DR REDDYS

100MG/ML A080571 001

200MG/ML A080571 002

HOSPIRA

100MG/ML A040079 001 May 03, 1996

LUITPOLD

100MG/ML A080667 001

PARKE DAVIS

100MG/ML A080770 001

WATSON LABS

100MG/ML A083534 001

200MG/ML A083534 002

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

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

AKORN

30MG/ML

A040125 001 Aug 16, 1996

100MG/ML

A040126 001 Aug 16, 1996

ALPHARMA US PHARMS

30MG/ML

A087766 001 Apr 26, 1983

ANI PHARMS

30MG/ML

A089602 001 Nov 09, 1987

100MG/ML

A089603 001 Nov 09, 1987

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

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

HERITAGE PHARMA AVET

10MG

A088476 001 Nov 08, 1983

25MG

A088478 001 Nov 08, 1983

50MG

A088479 001 Nov 08, 1983

100MG

A088736 001 Jul 24, 1984

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THIORIDAZINE HYDROCHLORIDE

TABLET; ORAL

THIORIDAZINE HYDROCHLORIDE

	25MG	A088336 001	Dec 05, 1983
	50MG	A088322 001	Dec 05, 1983
	100MG	A088480 001	Dec 29, 1983
	150MG	A089764 001	Feb 09, 1988
	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	10MG	A089953 004	Aug 01, 1986
	15MG	A088461 001	Nov 18, 1983
	25MG	A089953 003	Aug 01, 1986
	50MG	A089953 002	Aug 01, 1986
	100MG	A089953 001	Oct 07, 1988
	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	A088561 001	May 11, 1984
	15MG	A088345 001	Jul 28, 1983
	15MG	A088562 001	May 11, 1984
	25MG	A088296 001	Jul 28, 1983
	25MG	A088755 001	Jul 24, 1984
	50MG	A088323 001	Jul 28, 1983
	50MG	A088563 001	May 11, 1984
	100MG	A088284 001	Aug 25, 1983
	100MG	A088564 001	May 11, 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

HERITAGE PHARMA AVET 1MG A070600 001 Jun 05, 1987

2MG A070601 001 Jun 05, 1987

5MG A070602 001 Jun 05, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THIOTHIXENE

CAPSULE; ORAL

THIOTHIXENE

	10MG	A070603 001	Jun 05, 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	2MG	A071626 001	Jun 25, 1987
	5MG	A071627 001	Jun 25, 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

HIKMA

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

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

TIAGABINE HYDROCHLORIDE

WILSHIRE PHARMS INC

2MG

A206857 001

Oct 13, 2017

4MG

A206857 002

Oct 13, 2017

12MG

A206857 003

Oct 13, 2017

16MG

A206857 004

Oct 13, 2017

TICAGRELOR

TABLET; ORAL

TICAGRELOR

AMNEAL

90MG

A208531 001

Jan 23, 2019

SIGMAPHARM LABS LLC

90MG

A208596 001

Apr 07, 2020

SUNSHINE

90MG

A208508 001

Apr 06, 2020

WATSON LABS INC

60MG

A208390 001

Sep 04, 2018

90MG

A208390 002

Sep 04, 2018

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

CHARTWELL RX

250MG

A075318 001

Aug 20, 1999

250MG

A075326 001

Aug 20, 1999

MYLAN

250MG

A075161 001

Sep 13, 1999

250MG

A075316 001

Nov 02, 1999

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLOPIDINE HYDROCHLORIDE

SUN PHARM INDS INC

250MG

A075526 001 Sep 26, 2002

WATSON LABS

250MG

A075309 001 Apr 26, 2000

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

ANI PHARMS

5MG

A072917 001 Jul 31, 1991

10MG

A072918 001 Jul 31, 1991

20MG

A072919 001 Jul 31, 1991

CHARTWELL RX

5MG

A072550 001 Apr 13, 1989

10MG

A072551 001 Apr 13, 1989

20MG

A072552 001 Apr 13, 1989

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

10MG

A072270 001 Apr 11, 1989

20MG

A072271 001 Apr 11, 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

SOLUTION; INTRAVENOUS

AGGRASTAT

MEDICURE

EQ 25MG BASE/500ML (EQ 0.05MG BASE/ML)

N020913 001 May 14, 1998

TIZANIDINE HYDROCHLORIDE

CAPSULE; ORAL

TIZANIDINE HYDROCHLORIDE

MYLAN PHARMS INC

EQ 2MG BASE

A091502 001 Nov 09, 2012

EQ 4MG BASE

A091502 002 Nov 09, 2012

EQ 6MG BASE

A091502 003 Nov 09, 2012

PAR PHARM INC

EQ 2MG BASE

A207199 001 Mar 14, 2017

EQ 4MG BASE

A207199 002 Mar 14, 2017

EQ 6MG BASE

A207199 003 Mar 14, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TIZANIDINE HYDROCHLORIDE

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE

ANI PHARMS

EQ 2MG BASE

A076283 001 Jul 12, 2002

EQ 2MG BASE

A076284 001 Jul 03, 2002

EQ 2MG BASE

A076321 001 Sep 30, 2004

EQ 2MG BASE

A076371 001 Apr 09, 2003

EQ 4MG BASE

A076283 002 Jul 12, 2002

EQ 4MG BASE

A076284 002 Jul 03, 2002

EQ 4MG BASE

A076321 002 Sep 30, 2004

EQ 4MG BASE

A076371 002 Apr 09, 2003

MYLAN PHARMS INC

EQ 2MG BASE

A076282 001 Dec 16, 2003

EQ 4MG BASE

A076282 002 Dec 16, 2003

PAR PHARM INC

EQ 2MG BASE

A207170 001 Jan 26, 2017

EQ 4MG BASE

A207170 002 Jan 26, 2017

RISING PHARMA

EQ 2MG BASE

A076354 001 Mar 28, 2003

EQ 4MG BASE

A076354 002 Mar 28, 2003

ZANAFLEX

+ COVIS

EQ 2MG BASE **

N020397 002 Feb 04, 2000

TOBRAMYCIN

SOLUTION; INHALATION

TOBRAMYCIN

LUOXIN AUROVITAS

300MG/5ML

A210871 001 Jan 22, 2021

MYLAN

300MG/5ML

A209554 001 Oct 13, 2017

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

HIKMA

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

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

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

TOFACITINIB CITRATE

TABLET; ORAL

TOFACITINIB CITRATE

AJANTA PHARMA LTD

EQ 10MG BASE

A212943 001 Jun 01, 2021

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOLAZAMIDE

TABLET; ORAL

TOLAZAMIDE

BARR	100MG		A070162	001	Jan 14, 1986
	250MG		A070163	001	Jan 14, 1986
	500MG		A070164	001	Jan 14, 1986
CHARTWELL RX	100MG		A071633	001	Dec 09, 1987
	250MG		A070289	001	Mar 13, 1986
	500MG		A070290	001	Mar 13, 1986
COSETTE	100MG		N018894	001	Nov 02, 1984
	250MG		N018894	002	Nov 02, 1984
	500MG		N018894	003	Nov 02, 1984
DURAMED PHARMS BARR	100MG		A070165	001	Jan 10, 1986
	250MG		A070166	001	Jan 10, 1986
	500MG		A070167	001	Jan 10, 1986
INTERPHARM	250MG		A071270	001	Sep 23, 1986
	500MG		A071271	001	Sep 23, 1986
MYLAN PHARMS INC	250MG		A070259	001	Jan 02, 1986
	500MG		A070259	003	Mar 17, 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
	250MG		A070243	001	Aug 01, 1986
	250MG		A070514	001	Jan 09, 1986
	500MG		A070244	001	Aug 01, 1986
	500MG		A070515	001	Jan 09, 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
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TOLBUTAMIDE

TABLET; ORAL

ORINASE

PHARMACIA AND UPJOHN	250MG **		N010670	002	
	500MG **		N010670	001	
TOLBUTAMIDE					
ALRA	500MG		A086141	001	
ANI PHARMS	500MG		A087093	001	
ASCOT	500MG		A087541	001	Mar 01, 1983
BARR	500MG		A087121	001	
CHARTWELL RX	500MG		A086574	001	
DAVA PHARMS INC	500MG		A086926	001	
MYLAN PHARMS INC	500MG		A086445	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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOLBUTAMIDE SODIUM

INJECTABLE; INJECTION

ORINASE DIAGNOSTIC

PHARMACIA AND UPJOHN EQ 1GM BASE/VIAL

N012095 001

TOLCAPONE

TABLET; ORAL

TASMAR

BAUSCH 200MG

N020697 002 Jan 29, 1998

TOLCAPONE

ALVOGEN 100MG

A207729 001 Jul 29, 2020

MAYNE PHARMA 100MG

A210095 001 Aug 01, 2019

TOLMETIN SODIUM

CAPSULE; ORAL

TOLECTIN DS

+ ORTHO MCNEIL JANSSEN EQ 400MG BASE

N018084 001

TOLMETIN SODIUM

ANI PHARMS EQ 400MG BASE

A073308 001 Jan 24, 1992

EQ 400MG BASE

A073392 001 Jan 24, 1992

EQ 400MG BASE

A073519 001 May 29, 1992

FOSUN PHARMA EQ 400MG BASE

A073462 001 Apr 30, 1992

RISING EQ 400MG BASE

A073393 001 May 27, 1993

SUN PHARM INDUSTRIES EQ 400MG BASE

A073311 001 Nov 27, 1991

TEVA EQ 400MG BASE

A073290 001 Nov 27, 1991

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

ANI PHARMS EQ 600MG BASE

A073527 001 Jun 30, 1992

COSETTE EQ 600MG BASE

A074399 001 Mar 28, 1996

EQ 600MG BASE

A074729 001 Feb 27, 1997

FOSUN PHARMA EQ 200MG BASE

A073588 001 Jul 31, 1992

EQ 600MG BASE

A074002 001 Sep 27, 1993

RISING EQ 600MG BASE

A074473 001 Aug 30, 1994

SUN PHARM INDUSTRIES EQ 200MG BASE

A073310 001 Nov 27, 1991

TOLTERODINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

TOLTERODINE TARTRATE

AUROBINDO PHARMA USA 2MG

A201486 001 Oct 31, 2013

4MG

A201486 002 Oct 31, 2013

TABLET; ORAL

TOLTERODINE TARTRATE

APOTEX CORP 1MG

A200164 001 Sep 25, 2012

2MG

A200164 002 Sep 25, 2012

MYLAN PHARMS INC 1MG

A202641 001 Nov 27, 2012

2MG

A202641 002 Nov 27, 2012

TOLVAPTAN

TABLET; ORAL

SAMSCA

+ OTSUKA 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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOPIRAMATETABLET; ORAL
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
CHARTWELL	25MG	A078410 001	Sep 11, 2013
	50MG	A078410 002	Sep 11, 2013
	100MG	A078410 003	Sep 11, 2013
	200MG	A078410 004	Sep 11, 2013
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
TORRENT PHARMS	25MG	A079153 001	Mar 27, 2009
	50MG	A079153 002	Mar 27, 2009
	100MG	A079153 003	Mar 27, 2009
	200MG	A079153 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

TOPOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

TOPOTECAN HYDROCHLORIDE

FRESENIUS KABI USA	EQ 4MG BASE/VIAL	A091376 001	Nov 29, 2010
MEITHEAL	EQ 4MG BASE/VIAL	A201166 001	Aug 08, 2012
MYLAN LABS LTD	EQ 4MG BASE/VIAL	A091542 001	Aug 28, 2012
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

AM REGENT	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

DEMADEX

+ MYLAN SPECIALITY LP	5MG **	N020136 001	Aug 23, 1993
+	10MG **	N020136 002	Aug 23, 1993

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TORSEMIDE

TABLET;ORAL

DEMADEX

+		20MG **	N020136 003	Aug 23, 1993
+		100MG **	N020136 004	Aug 23, 1993

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

CAPSULE, EXTENDED RELEASE;ORAL

CONZIP

+	CIPHER PHARMS INC	150MG	N022370 004	Aug 01, 2011
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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
GRAVITI PHARMS	50MG	A075968 001	Jun 25, 2002
IVAX SUB TEVA PHARMS	50MG	A075963 001	Jul 03, 2002
MACLEODS PHARMS LTD	50MG	A205702 001	Sep 25, 2015
MYLAN PHARMS INC	50MG	A075980 001	Nov 21, 2002
NORTHSTAR HLTHCARE	50MG	A078935 001	May 26, 2010
SPECGX LLC	50MG	A075983 001	Jun 25, 2002
SUN PHARM INDUSTRIES	50MG	A076100 001	Jun 20, 2002
WATSON LABS	50MG	A075962 001	Jun 24, 2002

ULTRAM

JANSSEN PHARMS	100MG	N020281 001	Mar 03, 1995
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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

TRAMADOL HYDROCHLORIDE

ANCHEN PHARMS	100MG	A200491 001	Jun 27, 2012
	200MG	A200491 002	Jun 27, 2012
	300MG	A200491 003	Jun 27, 2012
AUROBINDO PHARMA LTD	100MG	A204421 001	Oct 20, 2015
	200MG	A204421 002	Oct 20, 2015
	300MG	A204421 003	Oct 20, 2015
STRIDES PHARMA	100MG	A078783 001	Nov 13, 2009
	200MG	A078783 002	Nov 13, 2009
	300MG	A078783 003	Sep 20, 2011

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
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TRAMETINIB DIMETHYL SULFOXIDE

TABLET;ORAL

MEKINIST

+	NOVARTIS	EQ 1MG	N204114 002	May 29, 2013
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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
	4MG	A078493 003	Aug 25, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRANDOLAPRIL

TABLET; ORAL

TRANDOLAPRIL

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

TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TARKA

+	ABBVIE	1MG;240MG **	N020591 003	Oct 22, 1996
+		2MG;180MG	N020591 001	Oct 22, 1996
+		2MG;240MG	N020591 004	Oct 22, 1996
+		4MG;240MG	N020591 002	Oct 22, 1996

TRANEXAMIC ACID

INJECTABLE; INJECTION

TRANEXAMIC ACID

VIRTUS PHARMS	100MG/ML	A202755 001	Feb 25, 2016
ZYDUS PHARMS	100MG/ML	A205228 001	Jul 17, 2017

TABLET; ORAL

CYKLOKAPRON

PHARMACIA AND UPJOHN	500MG	N019280 001	Dec 30, 1986
TRANEXAMIC ACID			
APOTEX INC	650MG	A202286 001	Jan 27, 2014
AUROBINDO PHARMA USA	650MG	A205133 001	Sep 21, 2015

TRAVOPROST

SOLUTION/DROPS; OPHTHALMIC

IZBA

+	NOVARTIS	0.003% **	N204822 001	May 15, 2014
+	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

ALVOGEN	50MG	A071636 001	Apr 18, 1988
	100MG	A071514 001	Apr 18, 1988
AM THERAP	50MG	A071139 001	Oct 29, 1986
	100MG	A071140 001	Oct 29, 1986
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
RISING	50MG	A072484 001	Apr 30, 1990
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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRAZODONE HYDROCHLORIDE

TABLET;ORAL

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

TREPROSTINIL

INJECTABLE;IV (INFUSION), SUBCUTANEOUS

TREPROSTINIL

DR REDDYS 1MG/ML

A210214 001 May 22, 2020

2.5MG/ML

A210214 002 May 22, 2020

5MG/ML

A210214 003 May 22, 2020

10MG/ML

A210214 004 May 22, 2020

SOLUTION;INTRAVENOUS, SUBCUTANEOUS

REMODULIN

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

TRETINOIN

CAPSULE;ORAL

VESANOID

+ CHEPLAPHARM 10MG **

N020438 001 Nov 22, 1995

CREAM;TOPICAL

RENOVA

+ VALEANT PHARMS NORTH 0.05%

N019963 001 Dec 29, 1995

TRETINOIN

ALLERGAN 0.0375%

A090098 001 Mar 22, 2010

0.075%

A202209 001 Oct 11, 2012

ZO SKIN HEALTH 0.05%

A076498 001 Sep 15, 2005

GEL;TOPICAL

TRETINOIN

MYLAN 0.04%

A202567 001 Jul 17, 2013

0.1%

A202026 001 Jul 17, 2013

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIAMCINOLONE

TABLET; ORAL

TRIAMCINOLONE

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
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AEROSOL, METERED; NASAL

NASACORT

SANOVI AVENTIS US	0.055MG/INH	N019798 001	Jul 11, 1991
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CREAM; TOPICAL

ARISTOCORT

ASTELLAS	0.025%	A083017 003
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	0.1%	A083016 004
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+	0.5%	A083015 002
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ARISTOCORT A

ASTELLAS	0.025%	A083017 004
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	0.025%	A088818 001	Oct 16, 1984
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	0.1%	A083016 005
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	0.1%	A088819 001	Oct 16, 1984
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	0.5%	A083015 003
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	0.5%	A088820 001	Oct 16, 1984
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FLUTEX

IVAX PHARMS	0.025%	A085539 001
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	0.1%	A085539 002
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	0.5%	A085539 003
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KENALOG

+	DELCOR ASSET CORP	0.5%	A083943 001
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KENALOG-H

DELCOR ASSET CORP	0.1%	A086240 001
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TRIACET

TEVA	0.025%	A084908 001
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	0.1%	A084908 002
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	0.5%	A084908 003
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TRIACORT

SOLVAY	0.1%	A087113 001
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TRIAMCINOLONE ACETONIDE

ACTAVIS MID ATLANTIC	0.1%	A087798 001	Jun 04, 1982
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ALPHARMA US PHARMS	0.025%	A087797 001	Jun 07, 1982
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AMBIX	0.025%	A087932 001	May 09, 1983
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MORTON GROVE	0.025%	A088094 001	Sep 01, 1983
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	0.1%	A088095 001	Sep 01, 1983
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	0.5%	A088096 001	Sep 01, 1983
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+	MYLAN	0.025% **	N011601 003
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+		0.1% **	N011601 006
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PHARMADERM	0.025%	A087990 001	Jul 07, 1983
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	0.1%	A087991 001	Jul 07, 1983
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	0.5%	A087992 001	Jul 07, 1983
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PHARMAFAIR	0.025%	A087921 001	Aug 10, 1982
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	0.1%	A087912 001	Aug 10, 1982
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	0.5%	A087922 001	Aug 10, 1982
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TARO	0.025%	A040038 001	Oct 26, 1994
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	0.025%	A086277 001
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	0.1%	A086276 001
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	0.5%	A086275 001
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TOPIDERM	0.025%	A089274 001	Feb 21, 1989
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	0.1%	A089275 001	Feb 21, 1989
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	0.5%	A089276 001	Feb 21, 1989
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

TRIALEX

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	
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INJECTABLE; INJECTION

TRIAMCINOLONE ACETONIDE

PARNELL	3MG/ML	N019503	001	Oct 16, 1987
SANDOZ 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
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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	
	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

+ DELCOR ASSET CORP	0.5% **	A083944	001	
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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
+	MYLAN	0.025% **	N011600	003
+		0.1% **	N011600	001
PHARMADERM	0.025%	A088692	001	Aug 02, 1984
	0.1%	A088690	001	Aug 02, 1984
STRIDES PHARMA	0.1%	A211315	001	Mar 18, 2020
	0.5%	A211315	002	Mar 18, 2020
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

+ DELCOR ASSET CORP	0.1% **	N012097	001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIAMCINOLONE ACETONIDE

PASTE;DENTAL

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

0.055MG/SPRAY

A078104 001 Jul 30, 2009

TRIAMCINOLONE DIACETATE

INJECTABLE; INJECTION

ARISTOCORT

FOSUN PHARMA

25MG/ML

N011685 003

+

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

DELCOR ASSET CORP

EQ 4MG BASE/5ML

N012515 001

TRIAZOLAM

TABLET;ORAL

HALCION

PFIZER

0.5MG

N017892 002 Nov 15, 1982

TRIAZOLAM

MYLAN PHARMS INC

0.125MG

A074031 001 Mar 25, 1994

0.25MG

A074031 002 Mar 25, 1994

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIDIHEXETHYL CHLORIDE

INJECTABLE; INJECTION

PATHILON

LEDERLE

10MG/ML

N009729 001

TABLET; ORAL

PATHILON

LEDERLE

25MG

N009489 005

TRIENTINE HYDROCHLORIDE

CAPSULE; ORAL

CLOVIQUE

CHARTWELL RX

250MG

A209731 001 Oct 21, 2019

TRIENTINE HYDROCHLORIDE

AMNEAL

250MG

A210619 001 Feb 08, 2019

CHARTWELL RX

250MG

A209415 001 Sep 16, 2019

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

ATHEM

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

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

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

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIHEXYPHENIDYL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

ARTANE

LEDERLE	5MG	N006773	010
	5MG	N012947	001

ELIXIR;ORAL

ARTANE

LEDERLE	2MG/5ML	N006773	009
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TRIHEXYPHENIDYL HYDROCHLORIDE

PHARM VENTURES	2MG/5ML	A089514	001	Apr 07, 1989
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TABLET;ORAL

ARTANE

+ LEDERLE	2MG **	N006773	005
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+	5MG **	N006773	003
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TREMIM

SCHERING	2MG	A080381	001
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	5MG	A080381	003
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TRIHEXYPHENIDYL HYDROCHLORIDE

HIKMA	2MG	A040337	002	Feb 16, 2000
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	5MG	A040337	001	Feb 16, 2000
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NYLOS	5MG	A085622	001
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VANGARD	2MG	A088035	001	Jul 30, 1982
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WATSON LABS	2MG	A040184	001	Feb 06, 1998
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	2MG	A085117	001
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	5MG	A040184	002	Feb 06, 1998
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	5MG	A085105	001
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TRILOSTANE

CAPSULE;ORAL

MODRASTANE

BIOENVISION	30MG	N018719	002	Dec 31, 1984
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	60MG	N018719	001	Dec 31, 1984
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TRIMEPRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE;ORAL

TEMARIL

ALLERGAN HERBERT	EQ 5MG BASE	N011316	004
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SYRUP;ORAL

TEMARIL

ALLERGAN HERBERT	EQ 2.5MG BASE/5ML	N011316	003
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TRIMEPRAZINE TARTRATE

ALPHARMA US PHARMS	EQ 2.5MG BASE/5ML	A085015	001	Feb 18, 1982
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MORTON GROVE	EQ 2.5MG BASE/5ML	A088285	001	Apr 11, 1985
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TABLET;ORAL

TEMARIL

ALLERGAN HERBERT	EQ 2.5MG BASE	N011316	001
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TRIMETHADIONE

CAPSULE;ORAL

TRIDIONE

ABBVIE	300MG	N005856	005
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SOLUTION;ORAL

TRIDIONE

ABBVIE	200MG/5ML	N005856	002
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TABLET;ORAL

TRIDIONE

+ ABBVIE	150MG	N005856	009
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TRIMETHAPHAN CAMSYLATE

INJECTABLE; INJECTION

ARFONAD

ROCHE	50MG/ML	N008983	001
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TRIMETHOBENZAMIDE HYDROCHLORIDE

CAPSULE;ORAL

TIGAN

+ KING PHARMS LLC	300MG	N017531	006	Dec 13, 2001
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INJECTABLE; INJECTION

TRIMETHOBENZAMIDE HYDROCHLORIDE

AM REGENT	100MG/ML	A091330	001	Mar 08, 2011
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HOSPIRA	100MG/ML	A088804	001	Apr 03, 1987
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SMITH AND NEPHEW	100MG/ML	A088960	001	Apr 04, 1986
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIMETHOBENZAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

TRIMETHOBENZAMIDE HYDROCHLORIDE

	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
TRIMETHOBENZAMIDE HYDROCHLORIDE PRESERVATIVE FREE			
AM REGENT	100MG/ML	A091329 001	Mar 08, 2011

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	
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TRIMPEX 200

ROCHE	200MG	N017952 002	Nov 09, 1982
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TRIMETHOPRIM HYDROCHLORIDE

SOLUTION; ORAL

PRIMSOL

ALLEGIS	EQ 25MG BASE/5ML	N074374 001	Jun 23, 1995
+	EQ 50MG BASE/5ML	N074973 001	Jan 24, 2000

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

SURMONTIL

+	ODYSSEY PHARMS	EQ 25MG BASE **	N016792 001
+		EQ 50MG BASE **	N016792 002
+		EQ 100MG BASE **	N016792 003

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	
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TRIPLENNAMINE CITRATE

ELIXIR; ORAL

PBZ

NOVARTIS	EQ 25MG HYDROCHLORIDE/5ML	N005914 004	
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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 effectiveness 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

COSETTE

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

PADAGIS US

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

COSETTE

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 effectiveness 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

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

TROVAFLOXACIN MESYLATE

TABLET;ORAL

TROVAN

PFIZER EQ 100MG BASE N020759 001 Dec 18, 1997

EQ 200MG BASE N020759 002 Dec 18, 1997

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
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UNOPROSTONE ISOPROPYL

SOLUTION/DROPS; OPHTHALMIC

RESCULA

+ SUCAMPO PHARMA LLC	0.15% **	N021214	001	Aug 03, 2000
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URACIL MUSTARD

CAPSULE; ORAL

URACIL MUSTARD

SHIRE	1MG	N012892	001
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UREA

INJECTABLE; INJECTION

STERILE UREA

HOSPIRA	40GM/VIAL	N017698	001
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UREAPHIL

HOSPIRA	40GM/VIAL	N012154	001
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UREA C-13

FOR SOLUTION; ORAL

BREATHTEK UBT FOR H-PYLORI

OTSUKA AMERICA	EQ 75MG/POUCH	N020586	002	May 10, 2001
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HELICOSOL

METABOLIC SOLUTIONS	125MG/VIAL	N021092	001	Dec 17, 1999
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MERETEK UBT KIT (W/ PRANACTIN)

OTSUKA AMERICA	125MG/VIAL	N020586	001	Sep 17, 1996
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PYLORI-CHEK BREATH TEST

DXS DEVICES	100MG/VIAL	N020900	001	Feb 04, 1999
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URSODIOL

CAPSULE; ORAL

ACTIGALL

ALLERGAN	150MG	N019594	001	Dec 31, 1987
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URSODIOL

IMPAX LABS INC	300MG	A077895	001	Jul 27, 2006
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TABLET; ORAL

URSODIOL

IMPAX LABS INC	250MG	A200826	001	Dec 23, 2011
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	500MG	A200826	002	Dec 23, 2011
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TEVA PHARMS USA	250MG	A079184	001	May 13, 2009
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	500MG	A079184	002	May 13, 2009
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VALACYCLOVIR HYDROCHLORIDE

TABLET; ORAL

VALACYCLOVIR HYDROCHLORIDE

HIKMA	EQ 500MG BASE	A078656	001	May 24, 2010
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	EQ 1GM BASE	A078656	002	May 24, 2010
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MYLAN	EQ 500MG BASE	A078070	001	May 24, 2010
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	EQ 1GM BASE	A078070	002	May 24, 2010
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TEVA PHARMS	EQ 500MG BASE	A077655	001	May 24, 2010
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	EQ 1GM BASE	A077655	002	May 24, 2010
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WATSON LABS INC	EQ 500MG BASE	A090370	001	Mar 16, 2011
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	EQ 1GM BASE	A090370	002	Mar 16, 2011
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VALDECOXIB

TABLET; ORAL

BEXTRA

GD SEARLE	10MG	N021341	002	Nov 16, 2001
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	20MG	N021341	003	Nov 16, 2001
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VALPROATE SODIUM

INJECTABLE; INJECTION

DEPAACON

+ ABBVIE

EQ 100MG BASE/ML

N020593 001 Dec 30, 1996

VALPROIC ACID

CAPSULE; ORAL

DEPAKENE

+ ABBVIE

250MG

N018081 001

VALPROIC ACID

PAR PHARM

250MG

A070431 001 Feb 28, 1986

SCHERER RP

250MG

A070195 001 Jul 02, 1987

UPSHER SMITH LABS

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

DEPAKENE

+ ABBVIE

250MG/5ML

N018082 001

VALPROIC ACID

ANI PHARMS

250MG/5ML

A073178 001 Aug 25, 1992

NOSTRUM LABS 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

TORRENT

40MG

A202728 001 Jan 05, 2015

80MG

A202728 002 Jan 05, 2015

160MG

A202728 003 Jan 05, 2015

320MG

A202728 004 Jan 05, 2015

UNICHEM

40MG

A209261 001 May 04, 2018

80MG

A209261 002 May 04, 2018

160MG

A209261 003 May 04, 2018

320MG

A209261 004 May 04, 2018

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

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

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 effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

VANCOLED

HIKMA	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

EMCURE PHARMS LTD

EQ 500MG BASE/VIAL	A202275 001	Oct 31, 2013
EQ 1GM BASE/VIAL	A202275 002	Oct 31, 2013
EQ 10GM BASE/VIAL	A202464 001	Oct 09, 2013
EQ 5GM BASE/VIAL	A202274 001	Oct 31, 2013

HIKMA

EQ 500MG BASE/VIAL	A062879 001	Aug 02, 1988
EQ 500MG BASE/VIAL	A203300 001	Aug 11, 2020
EQ 1GM BASE/VIAL	A062879 002	Aug 02, 1988
EQ 1GM BASE/VIAL	A203300 002	Aug 11, 2020

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

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	EQ 2.5MG BASE **	N021400 003	Aug 19, 2003
+	EQ 5MG BASE	N021400 001	Aug 19, 2003
+	EQ 10MG BASE	N021400 002	Aug 19, 2003

VARDENAFIL HYDROCHLORIDE

AMNEAL PHARMS CO

EQ 5MG BASE	A210738 001	Oct 31, 2018
EQ 10MG BASE	A210738 002	Oct 31, 2018
EQ 20MG BASE	A210738 003	Oct 31, 2018

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

HIKMA

10MG/VIAL	A075218 001	Aug 23, 1999
20MG/VIAL	A075218 002	Aug 23, 1999

HOSPIRA

EQ 4MG/VIAL A075558 001 Sep 11, 2001

WATSON LABS

10MG/VIAL	A074334 001	Aug 31, 1995
20MG/VIAL	A074334 002	Aug 31, 1995

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

EFFEXOR XR

UPJOHN 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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VENLAFAXINE HYDROCHLORIDE

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

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

MYLAN

		EQ 25MG BASE	A077166 001	Jun 13, 2008
		EQ 37.5MG BASE	A077166 002	Jun 13, 2008
		EQ 50MG BASE	A077166 003	Jun 13, 2008
		EQ 75MG BASE	A077166 004	Jun 13, 2008
		EQ 100MG BASE	A077166 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

PRINSTON INC

		EQ 25MG BASE	A090027 001	Aug 04, 2010
		EQ 37.5MG BASE	A090027 002	Aug 04, 2010
		EQ 50MG BASE	A090027 003	Aug 04, 2010
		EQ 75MG BASE	A090027 004	Aug 04, 2010
		EQ 100MG BASE	A090027 005	Aug 04, 2010

VERAPAMIL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

VERAPAMIL HYDROCHLORIDE

MYLAN

		100MG	A078306 001	Aug 09, 2007
		200MG	A078306 002	Aug 09, 2007
		300MG	A078306 003	Aug 09, 2007

INJECTABLE; INJECTION

CALAN

GD SEARLE LLC

		2.5MG/ML	N019038 001	Mar 30, 1984
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ISOPTIN

+ MT ADAMS

		2.5MG/ML **	N018485 001	
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VERAPAMIL HYDROCHLORIDE

ABRAXIS PHARM

		2.5MG/ML	A070348 001	May 01, 1986
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BEDFORD

		2.5MG/ML	A072888 001	Jul 28, 1995
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HOSPIRA

		2.5MG/ML	A070577 001	Feb 02, 1987
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		2.5MG/ML	A070739 001	May 06, 1987
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		2.5MG/ML	A070740 001	May 06, 1987
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INTL MEDICATION

		2.5MG/ML	A070451 001	Dec 16, 1985
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LUITPOLD

		2.5MG/ML	A070225 001	Nov 12, 1985
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		2.5MG/ML	A070617 001	Nov 12, 1985
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MARSAM PHARMS LLC

		2.5MG/ML	A072233 001	Feb 26, 1993
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		2.5MG/ML	A073485 001	Sep 27, 1993
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SMITH AND NEPHEW

		2.5MG/ML	A070696 001	Jul 31, 1987
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		2.5MG/ML	A070697 001	Jul 31, 1987
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SOLOPAK

		2.5MG/ML	A070695 001	Jul 31, 1987
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SOLUTION; INTRAVENOUS

VERAPAMIL HYDROCHLORIDE

EXELA PHARMA

		10MG/4ML (2.5MG/ML)	N018925 002	Apr 05, 2018
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TABLET; ORAL

CALAN

PFIZER

		40MG **	N018817 003	Feb 23, 1988
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+

		80MG **	N018817 001	Sep 10, 1984
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+

		120MG **	N018817 002	Sep 10, 1984
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		160MG **	N018817 004	Feb 23, 1988
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ISOPTIN

+ MT ADAMS

		40MG **	N018593 003	Nov 23, 1987
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+

		80MG **	N018593 001	Mar 08, 1982
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+

		120MG **	N018593 002	Mar 08, 1982
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VERAPAMIL HYDROCHLORIDE

TABLET; ORAL

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
RISING PHARMA	80MG	A071483 002	Feb 15, 1989
	120MG	A071483 001	Feb 15, 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

PFIZER 180MG N020552 001 Feb 26, 1996

240MG N020552 002 Feb 26, 1996

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
RISING PHARMA	120MG	A074587 002	Feb 21, 1997
	180MG	A074587 003	Sep 09, 1997
	240MG	A074587 001	Mar 23, 1996

VERATRUM VIRIDE ROOT

TABLET; ORAL

VERTAVIS

MEDPOINTE PHARM HLC 130CSR UNIT N005691 002

VIDARABINE

INJECTABLE; INJECTION

VIRA-A

PARKEDALE EQ 187.4MG BASE/ML N050523 001

OINTMENT; OPHTHALMIC

VIRA-A

PARKEDALE 3% N050486 001

VIGABATRIN

FOR SOLUTION; ORAL

VIGABATRIN

SPECGX LLC 500MG/PACKET A212626 001 Jul 28, 2021

VILAZODONE HYDROCHLORIDE

TABLET; ORAL

VILAZODONE HYDROCHLORIDE

TEVA PHARMS USA 10MG A208212 001 Sep 30, 2019

20MG A208212 002 Sep 30, 2019

40MG A208212 003 Sep 30, 2019

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

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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
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VINCREX

	BRISTOL MYERS SQUIBB	5MG/VIAL	A070867 001	Jul 12, 1988
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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
		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

INJECTABLE, LIPOSOMAL; INTRAVENOUS

MARQIBO KIT

+	ACROTECH	5MG/5ML (1MG/ML)	N202497 001	Aug 09, 2012
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VINORELBINE TARTRATE

INJECTABLE; INJECTION

NAVELBINE

+	PIERRE FABRE	EQ 10MG BASE/ML **	N020388 001	Dec 23, 1994
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VINORELBINE TARTRATE

	EBEWE PHARMA	EQ 10MG BASE/ML	A078408 001	Feb 13, 2008
	FRESENIUS KABI USA	EQ 10MG BASE/ML	A076849 001	Apr 18, 2005
	HOSPIRA	EQ 10MG BASE/ML	A076827 001	Jun 02, 2005
	MYLAN LABS LTD	EQ 10MG BASE/ML	A200148 001	Aug 31, 2012
	NOVAST LABS	EQ 10MG BASE/ML	A208997 001	Aug 05, 2019

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	

VITAMIN A PALMITATE

CAPSULE; ORAL

AFAXIN

	STERLING WINTHROP	EQ 50,000 UNITS BASE	A083187 001	
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ALPHALIN

	LILLY	EQ 50,000 UNITS BASE	A080883 001	
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DEL-VI-A

	DEL RAY LABS	EQ 50,000 UNITS BASE	A080830 001	
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VI-DOM-A

	BAYER PHARMS	EQ 50,000 UNITS BASE	A080972 001	
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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	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VITAMIN A PALMITATE

CAPSULE; ORAL

VITAMIN A

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
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INJECTABLE; INJECTION

VITAMIN A PALMITATE

BEL MAR	EQ 50,000 UNITS BASE/ML	A080819 001
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VORICONAZOLE

FOR SUSPENSION; ORAL

VORICONAZOLE

MYLAN PHARMS INC	200MG/5ML	A202361 001	May 28, 2013
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TABLET; ORAL

VORICONAZOLE

TEVA PHARMS	50MG	A091658 001	Apr 06, 2012
	200MG	A091658 002	Apr 06, 2012

VORTIOXETINE HYDROBROMIDE

TABLET; ORAL

TRINTELLIX

+ TAKEDA PHARMS USA	EQ 15MG BASE **	N204447 003	Sep 30, 2013
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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

COUMADIN

+ BRISTOL MYERS SQUIBB	1MG	N009218 022	Mar 01, 1990
	2MG	N009218 013	
	2.5MG	N009218 018	
	3MG	N009218 025	Nov 18, 1996
	4MG	N009218 023	Aug 24, 1993
	5MG	N009218 007	
	6MG	N009218 026	Nov 18, 1996
	7.5MG	N009218 016	
	10MG	N009218 005	

PANWARFIN

ABBOTT	2MG	N017020 001
	2.5MG	N017020 002
	5MG	N017020 003
	7.5MG	N017020 004
	10MG	N017020 005

WARFARIN SODIUM

CHARTWELL RX	1MG	A040196 001	Sep 30, 1997
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

WARFARIN SODIUM

TABLET; ORAL

WARFARIN SODIUM

MYLAN

USL PHARMA

WATSON LABS

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
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
2MG	A088719 001	Jun 27, 1985
2.5MG	A088720 001	Aug 06, 1985
5MG	A088721 001	Jul 02, 1985
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

XENON XE-127

GAS; INHALATION

XENON XE 127

MALLINCKRODT

5mCi/VIAL	N018536 001	Oct 01, 1982
10mCi/VIAL	N018536 002	Oct 01, 1982

XENON XE-133

GAS; INHALATION

XENON XE 133

GE HEALTHCARE

GEN ELECTRIC

XENON XE 133-V.S.S.

GE HEALTHCARE

INJECTABLE; INJECTION

XENON XE 133

GE HEALTHCARE

LANTHEUS MEDCL

SOLUTION; INHALATION, INJECTION

XENEISOL

MALLINCKRODT

1 CI/AMP	N017256 002	
10mCi/VIAL	N017687 002	
20mCi/VIAL	N017687 003	
5-100 CI/CYLINDER	N017550 001	
0.25-5 CI/AMP	N017550 003	
10mCi/VIAL	N017687 001	
1.3-1.7 CI/AMP	N017256 001	
6.3mCi/ML	N017283 001	
18-25mCi/AMP	N017262 002	

XYLOSE

POWDER; ORAL

XYLO-PFAN

SAVAGE LABS

XYLOSE

LYNE

25GM/BOT	N017605 001	
25GM/BOT	N018856 001	Mar 26, 1987

ZALCITABINE

TABLET; ORAL

HIVID

ROCHE

0.375MG	N020199 001	Jun 19, 1992
0.75MG	N020199 002	Jun 19, 1992

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ZALEPLON

CAPSULE; ORAL

ZALEPLON

HIKMA PHARMS	5MG	A078147 001	Nov 25, 2008
	10MG	A078147 002	Nov 25, 2008
MYLAN	5MG	A077238 001	Jun 06, 2008
	10MG	A077238 002	Jun 06, 2008
TEVA PHARMS	5MG	A077239 001	Jun 06, 2008
	10MG	A077239 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	200MCG/2ML (100MCG/ML)	N021060 003	Dec 28, 2004
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ZIDOVUDINE

INJECTABLE; INJECTION

ZIDOVUDINE

AM REGENT	10MG/ML	A091457 001	May 06, 2010
LIAONING CHENGDA	10MG/ML	A204538 001	Nov 26, 2013

TABLET; ORAL

RETROVIR

VIIV 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
HIKMA	300MG	A076844 001	Sep 19, 2005
MYLAN LABS LTD	100MG	N200732 001	Feb 23, 2011
RANBAXY LABS LTD	300MG	A077327 001	Sep 19, 2005

ZILEUTON

TABLET; ORAL

ZYFLO

CHIESI	300MG	N020471 001	Dec 09, 1996
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ZINC SULFATE

INJECTABLE; INJECTION

ZINC SULFATE

ABRAXIS PHARM	EQ 1MG ZINC/ML	N019229 002	May 05, 1987
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ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

ZIPRASIDONE HYDROCHLORIDE

MYLAN	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
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ZOLEDRONIC ACID

INJECTABLE; INTRAVENOUS

ZOLEDRONIC ACID

ACTAVIS INC	EQ 4MG BASE/5ML	A202472 001	Mar 04, 2013
DR REDDYS LABS LTD	EQ 4MG BASE/100ML	A204344 001	Nov 19, 2018
EMCURE PHARMS LTD	EQ 4MG BASE/5ML	A201783 001	Mar 12, 2013
	EQ 5MG BASE/100ML	A201801 001	Mar 29, 2013
SHILPA	EQ 4MG BASE/5ML	A208513 001	May 15, 2019
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
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ZOLMITRIPTAN

TABLET;ORAL

ZOLMITRIPTAN

ANI PHARMS	2.5MG	A090861 001	Mar 04, 2014
	5MG	A090861 002	Mar 04, 2014
APOTEX INC	2.5MG	A202078 001	May 14, 2013
	5MG	A202078 002	May 14, 2013
MACLEODS PHARMS LTD	2.5MG	A203772 001	Sep 30, 2015
	5MG	A203772 002	Sep 30, 2015
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
MACLEODS PHARMS LTD	2.5MG	A204336 001	Oct 22, 2015
	5MG	A204336 002	Oct 22, 2015
RISING PHARMA	2.5MG	A202855 001	Sep 20, 2019
	5MG	A202855 002	Sep 20, 2019

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
RISING PHARMA	5MG	A076578 001	Apr 23, 2007
	10MG	A076578 002	Apr 23, 2007
STRIDES PHARMA	5MG	A076062 001	Apr 23, 2007
	5MG	A078616 001	Nov 21, 2008
	10MG	A076062 002	Apr 23, 2007
	10MG	A078616 002	Nov 21, 2008
SUN PHARM INDS INC	5MG	A077359 001	Apr 23, 2007
	10MG	A077359 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
WOCKHARDT	5MG	A078426 001	May 15, 2007
	10MG	A078426 002	May 15, 2007
YUNG SHIN PHARM	5MG	A077990 001	Apr 23, 2007
	10MG	A077990 002	Apr 23, 2007

TABLET;SUBLINGUAL

INTERMEZZO

+ PURDUE PHARMA	1.75MG	N022328 001	Nov 23, 2011
+	3.5MG	N022328 002	Nov 23, 2011

ZOLPIDEM TARTRATE

MYLAN	5MG	A202657 001	Aug 08, 2016
	10MG	A202657 002	Aug 08, 2016

TABLET, EXTENDED RELEASE;ORAL

ZOLPIDEM TARTRATE

ACTAVIS ELIZABETH	6.25MG	A078179 002	Oct 13, 2010
	12.5MG	A078179 001	Jun 06, 2011
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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ZONISAMIDE

CAPSULE; ORAL

ZONEGRAN

+ CONCORDIA

50MG **

N020789 002 Aug 22, 2003

ZONISAMIDE

ANI PHARMS

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

EPIC PHARMA LLC

25MG

A077876 001 Feb 21, 2007

50MG

A077876 002 Feb 21, 2007

100MG

A077876 003 Feb 21, 2007

HERITAGE PHARMA AVET

25MG

A077650 001 Apr 20, 2006

50MG

A077650 002 Apr 20, 2006

100MG

A077650 003 Apr 20, 2006

MYLAN PHARMS INC

25MG

A077647 001 Dec 22, 2005

50MG

A077647 002 Dec 22, 2005

100MG

A077647 003 Dec 22, 2005

RISING PHARMA

25MG

A077637 001 Dec 22, 2005

50MG

A077637 002 Dec 22, 2005

100MG

A077637 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

ORPHAN PRODUCTS 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 ****

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
ABSORICA LD, ISOTRETINOIN
ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
ACANYA, BENZOYL PEROXIDE
ACARBOSE, ACARBOSE
ACCOLATE, ZAFIRLUKAST
ACCRUFER, FERRIC MALTOL
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
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
ACITRETIN, ACITRETIN
ACTHAR GEL, CORTICOTROPIN
ACTICLATE, DOXYCYCLINE HYCLATE
ACTIGALL, URSODIOL
ACTIQ, FENTANYL CITRATE
ACTIVELLA, ESTRADIOL
ACTONEL, RISEDRONATE SODIUM
ACTOPLUS MET, METFORMIN HYDROCHLORIDE
ACTOS, PIOGLITAZONE HYDROCHLORIDE
ACULAR, KETOROLAC TROMETHAMINE
ACULAR LS, KETOROLAC TROMETHAMINE
ACUVAIL, KETOROLAC TROMETHAMINE
ACYCLOVIR, ACYCLOVIR
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
ACZONE, DAPSONE
ADAPALENE, ADAPALENE (OTC)
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
ADDERALL XR 5, AMPHETAMINE ASPARTATE
ADDYI, FLIBANSERIN
ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL
ADEMPAS, RIOCIQUAT

APPENDIX A - PRODUCT NAME INDEX**** A ****

ADENOSINE, ADENOSINE
ADHANSIA XR, METHYLPHENIDATE HYDROCHLORIDE
ADIPEX-P, PHENTERMINE HYDROCHLORIDE
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 DUAL ACTION WITH ACETAMINOPHEN, ACETAMINOPHEN (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)
ADZENYS XR-ODT, AMPHETAMINE
AEMCOLO, RIFAMYCIN SODIUM
AFINITOR, EVEROLIMUS
AFINITOR DISPERZ, EVEROLIMUS
AFIRMELLE, ETHINYL ESTRADIOL
AGGRASTAT, TIROFIBAN HYDROCHLORIDE
AGGRENOLX, ASPIRIN
AGRYLIN, ANAGRELIDE HYDROCHLORIDE
AIRDUO DIGIHALER, FLUTICASONE PROPIONATE
AIRDUO RESPICLICK, FLUTICASONE PROPIONATE
AK-FLUOR 10%, FLUORESCEIN SODIUM
AK-FLUOR 25%, FLUORESCEIN SODIUM
AKLIEF, TRIFAROTENE
AKOAZ, EPHEDRINE SULFATE
AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
AKTEN, LIDOCAINE HYDROCHLORIDE
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)
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

APPENDIX A - PRODUCT NAME INDEX**** A ****

ALISKIREN HEMIFUMARATE, ALISKIREN HEMIFUMARATE
 ALKINDI SPRINKLE, HYDROCORTISONE
 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
 ALOCRI, NEDOCROMIL SODIUM
 ALOMIDE, LODOXAMIDE TROMETHAMINE
 ALOPRIM, ALLOPURINOL SODIUM
 ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 ALPHAGAN P, BRIMONIDINE TARTRATE
 ALPRAZOLAM, ALPRAZOLAM
 ALPROSTADIL, ALPROSTADIL
 ALREX, LOTEHPREDNOL ETABONATE
 ALTABAX, RETAPAMULIN
 ALTACE, RAMIPRIL
 ALTAFLUOR BENOX, BENOXINATE HYDROCHLORIDE
 ALTAVERA, ETHINYL ESTRADIOL
 ALTOPREV, LOVASTATIN
 ALTRENO, TRETINOIN
 ALUNBRIG, BRIGATINIB
 ALVESCO, CICLESONIDE
 ALVIMOPAN, ALVIMOPAN
 ALYACEN 1/35, ETHINYL ESTRADIOL
 ALYACEN 7/7/7, ETHINYL ESTRADIOL
 ALYQ, TADALAFIL
 AMABELZ, ESTRADIOL
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMARYL, GLIMEPIRIDE
 AMBIEN, ZOLPIDEM TARTRATE
 AMBIEN CR, ZOLPIDEM TARTRATE
 AMBISOME, AMPHOTERICIN B
 AMBRISENTAN, AMBRISENTAN
 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 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
 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

APPENDIX A - PRODUCT NAME INDEX**** A ****

AMMONIUM CHLORIDE IN PLASTIC CONTAINER, AMMONIUM CHLORIDE
AMMONIUM LACTATE, AMMONIUM LACTATE
AMMONUL, SODIUM BENZOATE
AMNESTEEM, ISOTRETINOIN
AMONDYS 45, CASIMERSSEN
AMOXAPINE, AMOXAPINE
AMOXICILLIN, AMOXICILLIN
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN PEDIATRIC, AMOXICILLIN
AMOXIL, AMOXICILLIN
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
AMZEEQ, MINOCYCLINE HYDROCHLORIDE
AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT
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
ANGIOMAX RTU, BIVALIRUDIN
ANJESO, MELOXICAM
ANNOVERA, ETHINYL ESTRADIOL
ANORO ELLIPTA, UMECLIDINIUM BROMIDE
ANTARA (MICRONIZED), FENOFIBRATE
ANTHELIOS 20, AVOBENZONE (OTC)
ANTHELIOS 40, AVOBENZONE (OTC)
ANTHELIOS SX, AVOBENZONE (OTC)
ANTIVERT, MECLIZINE HYDROCHLORIDE
ANUSOL HC, HYDROCORTISONE
ANZEMET, DOLASETRON MESYLATE
APADAZ, ACETAMINOPHEN
APIXABAN, APIXABAN
APLENZIN, BUPROPION HYDROBROMIDE
APOKYN, APOMORPHINE HYDROCHLORIDE
APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE
APREMILAST, APREMILAST
APREPITANT, APREPITANT
APRETUDE, CABOTEGRAVIR
APRISO, MESALAMINE
APTENSIO XR, METHYLPHENIDATE HYDROCHLORIDE
APTIOM, ESLICARBAZEPINE ACETATE
APTIVUS, TIPRANAVIR
AQUASOL A, VITAMIN A PALMITATE
ARAKODA, TAFENOQUINE SUCCINATE
ARANELLE, ETHINYL ESTRADIOL
ARAVA, LEFLUNOMIDE
ARAZLO, TAZAROTENE
ARESTIN, MINOCYCLINE HYDROCHLORIDE
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE

APPENDIX A - PRODUCT NAME INDEX

** A **

ARGATROBAN, 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 DIGIHALER, FLUTICASONE PROPIONATE
ARNUITY ELLIPTA, FLUTICASONE FUROATE
AROMASIN, EXEMESTANE
ARRANON, NELARABINE
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
ARTESUNATE, ARTESUNATE
ARTHROTEC, DICLOFENAC SODIUM
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
ASTEPRO ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
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
ATRIPLA, EFAVIRENZ
ATROPEN, ATROPINE
ATROPINE SULFATE, ATROPINE SULFATE
ATROVENT HFA, IPRATROPIUM BROMIDE
AUBAGIO, TERIFLUNOMIDE
AUGMENTIN '125', AMOXICILLIN
AUGMENTIN '250', AMOXICILLIN
AUGMENTIN '875', AMOXICILLIN
AUGMENTIN ES-600, AMOXICILLIN
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

APPENDIX A - PRODUCT NAME INDEX**** A ****

AVAPRO, IRBESARTAN
 AVEED, TESTOSTERONE UNDECANOATE
 AVIANE-28, ETHINYL ESTRADIOL
 AVITA, TRETINOIN
 AVODART, DUTASTERIDE
 AVYCAZ, AVIBACTAM SODIUM
 AXID AR, NIZATIDINE (OTC)
 AXUMIN, FLUCICLOVINE F-18
 AYUNA, ETHINYL ESTRADIOL
 AYVAKIT, AVAPRITINIB
 AZACITIDINE, AZACITIDINE
 AZACTAM, AZTREONAM
 AZASAN, AZATHIOPRINE
 AZASITE, AZITHROMYCIN
 AZATHIOPRINE, AZATHIOPRINE
 AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
 AZEDRA, IOBENGUANE I-131
 AZELAIC ACID, AZELAIC ACID
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE, AZELASTINE HYDROCHLORIDE
 AZELEX, AZELAIC ACID
 AZILECT, RASAGILINE MESYLATE
 AZITHROMYCIN, AZITHROMYCIN
 AZOPT, BRINZOLAMIDE
 AZOR, AMLODIPINE BESYLATE
 AZSTARYS, DEXMETHYLPHENIDATE HYDROCHLORIDE
 AZTREONAM, AZTREONAM
 AZULFIDINE, SULFASALAZINE
 AZULFIDINE EN-TABS, SULFASALAZINE

**** B ****

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
 BAFIERTAM, MONOMETHYL FUMARATE
 BAL, DIMERCAPROL
 BALANCED SALT, CALCIUM CHLORIDE
 BALCOLTRA, ETHINYL ESTRADIOL
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 BALVERSA, ERDAFITINIB
 BALZIVA-28, ETHINYL ESTRADIOL
 BANZEL, RUFINAMIDE
 BAQSIMI, GLUCAGON
 BARACLUDE, ENTECAVIR
 BARHEMSYS, AMISULPRIDE
 BAXDELA, DELAFLOXACIN MEGLUMINE
 BECONASE AQ, BECLOMETHASONE DIPROPIONATE MONOHYDRATE
 BEKYREE, DESOGESTREL
 BELBUCA, BUPRENORPHINE HYDROCHLORIDE
 BELEODAQ, BELINOSTAT
 BELRAPZO, BENDAMUSTINE HYDROCHLORIDE
 BELSOMRA, SUVOREXANT
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BENDEKA, BENDAMUSTINE HYDROCHLORIDE
 BENICAR, OLMESARTAN MEDOXOMIL
 BENICAR HCT, HYDROCHLOROTHIAZIDE
 BENTYL, DICYCLOMINE HYDROCHLORIDE
 BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE

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BENZAFLIN, BENZOYL PEROXIDE
BENZAMYCIN, BENZOYL PEROXIDE
BENZNIDAZOLE, BENZNIDAZOLE
BENZONATATE, BENZONATATE
BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
BEPOTASTINE BESILATE, BEPOTASTINE BESILATE
BEPREVE, BEPOTASTINE BESILATE
BESIVANCE, BESIFLOXACIN HYDROCHLORIDE
BETA-VAL, BETAMETHASONE VALERATE
BETADINE, POVIDONE-IODINE
BETAGAN, LEVOBUNOLOL HYDROCHLORIDE
BETAINE, BETAINE
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
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
BIORPHEN, PHENYLEPHRINE HYDROCHLORIDE
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
BONJESTA, DOXYLAMINE SUCCINATE
BONSITY, TERIPARATIDE
BONTRIL PDM, PHENDIMETRAZINE TARTRATE
BORTEZOMIB, BORTEZOMIB
BOSENTAN, BOSENTAN
BOSULIF, BOSUTINIB MONOHYDRATE
BRAFTOVI, ENCORAFENIB
BREO ELLIPTA, FLUTICASONE FUROATE
BRETHINE, TERBUTALINE SULFATE
BRETILIUM TOSYLATE, BRETILIUM TOSYLATE
BREVIBLOC, ESMOLOL HYDROCHLORIDE
BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX**** B ****

BREVIBLOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 BREVITAL SODIUM, METHOHEXITAL SODIUM
 BREXAFEMME, IBREXAFUNGERP CITRATE
 BREZTRI AEROSPHERE, BUDESONIDE
 BRIDION, SUGAMMADEX SODIUM
 BRIELLYN, ETHINYL ESTRADIOL
 BRILINTA, TICAGRELOR
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BRINZOLAMIDE, BRINZOLAMIDE
 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
 BRONCHITOL, MANNITOL
 BROVANA, ARFORMOTEROL TARTRATE
 BRUKINSA, ZANUBRUTINIB
 BRYHALI, HALOBETASOL PROPIONATE
 BSS, CALCIUM CHLORIDE
 BSS PLUS, CALCIUM CHLORIDE
 BUDESONIDE, BUDESONIDE (OTC)
 BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 BUMEX, BUMETANIDE
 BUPHENYL, SODIUM PHENYLBUTYRATE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, 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)
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 BUTRANS, BUPRENORPHINE
 BYDUREON BCISE, EXENATIDE SYNTHETIC
 BYETTA, EXENATIDE SYNTHETIC
 BYFAVO, REMIMAZOLAM BESYLATE
 BYLVAY, ODEVIXIBAT
 BYSTOLIC, NEBIVOLOL HYDROCHLORIDE

**** C ****

CABAZITAXEL, CABAZITAXEL
 CABENUVA KIT, CABOTEGRAVIR
 CABERGOLINE, CABERGOLINE
 CABOMETYX, CABOZANTINIB S-MALATE
 CADUET, AMLODIPINE BESYLATE
 CAFCIT, CAFFEINE CITRATE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALAN SR, VERAPAMIL HYDROCHLORIDE
 CALCIPOTRIENE, CALCIPOTRIENE
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE

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CALCIPOTRIENE AND BETHAMETHASONE DIPROPIONATE, BETHAMETHASONE 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, EDETATE CALCIUM DISODIUM
CALCIUM GLUCONATE, CALCIUM GLUCONATE
CALCIUM GLUCONATE IN SODIUM CHLORIDE, CALCIUM GLUCONATE
CALDOLOR, IBUPROFEN
CALQUENCE, ACALABRUTINIB
CAMBIA, DICLOFENAC POTASSIUM
CAMCEVI KIT, LEUPROLIDE MESYLATE
CAMILA, NORETHINDRONE
CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
CANASA, MESALAMINE
CANCIDAS, CASPOFUNGIN ACETATE
CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
CAPECITABINE, CAPECITABINE
CAPEX, FLUOCINOLONE ACETONIDE
CAPITAL SOLEIL 15, AVOBENZONE (OTC)
CAPLYTA, LUMATEPERONE TOSYLATE
CAPRELSA, VANDETANIB
CAPTOPRIL, 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
CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
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
CARFILZOMIB, CARFILZOMIB
CARGLUMIC ACID, CARGLUMIC ACID
CARISOPRODOL, CARISOPRODOL
CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
CARMUSTINE, CARMUSTINE
CARNITOR, LEVOCARNITINE
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
CATAFLAM, DICLOFENAC POTASSIUM
CATAPRES-TTS-1, CLONIDINE

APPENDIX A - PRODUCT NAME INDEX**** C ****

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
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
CEFTAROLINE FOSAMIL, CEFTAROLINE FOSAMIL
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
CERIANNA, FLUOROESTRADIOL F-18
CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
CERVIDIL, DINOPROSTONE
CESAMET, NABILONE
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE (OTC)
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
CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
CHABELINA FE, ETHINYL ESTRADIOL
CHANTIX, VARENICLINE TARTRATE
CHEMET, SUCCIMER
CHENODIOL, CHENODIOL

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** C **

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 ASTEPRO ALLERGY, AZELASTINE 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 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
 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
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE (OTC)
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CHLORZOXAZONE, CHLORZOXAZONE
 CHOLBAM, CHOLIC ACID
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLETEC, TECHNETIUM TC-99M MEBROFENIN KIT
 CHOLINE C-11, CHOLINE C-11
 CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE
 CIALIS, TADALAFIL
 CICLOPIROX, CICLOPIROX
 CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)
 CIDOFOVIR, CIDOFOVIR
 CILOSTAZOL, CILOSTAZOL
 CILOXAN, CIPROFLOXACIN HYDROCHLORIDE
 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 AND DEXAMETHASONE, CIPROFLOXACIN
 CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT

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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 12 HOUR, DESLORATADINE
 CLARISCAN, GADOTERATE MEGLUMINE
 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 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
 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,

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CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
CLINOLIPID 20%, OLIVE OIL
CLOBAZAM, CLOBAZAM
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
CLOBEX, CLOBETASOL PROPIONATE
CLOCORTOLONE PIVALATE, CLOCORTOLONE PIVALATE
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
CLOTTRIMAZOLE, CLOTTRIMAZOLE (OTC)
CLOTTRIMAZOLE, CLOTTRIMAZOLE
CLOTTRIMAZOLE 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
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
COLOCORT, HYDROCORTISONE
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
CONJUPRI, LEVAMLODIPINE MALEATE
CONRAY, IOTHALAMATE MEGLUMINE
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
CORLANOR, IVABRADINE HYDROCHLORIDE
CORLOPAM, FENOLDOPAM MESYLATE
CORMAX, CLOBETASOL PROPIONATE
CORPHEDRA, EPHEDRINE SULFATE
CORTEF, HYDROCORTISONE

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CORTENEMA, HYDROCORTISONE
 CORTIFOAM, HYDROCORTISONE ACETATE
 CORTISONE ACETATE, CORTISONE ACETATE
 CORTROSYN, COSYNTROPIN
 CORVERT, IBUTILIDE FUMARATE
 COSELA, TRILACICLIB DIHYDROCHLORIDE
 COSMEGEN, DACTINOMYCIN
 COSOPT, DORZOLAMIDE HYDROCHLORIDE
 COSOPT PF, DORZOLAMIDE HYDROCHLORIDE
 COSYNTROPIN, COSYNTROPIN
 COTELLIC, COBIMETINIB FUMARATE
 COTEMPLA XR-ODT, METHYLPHENIDATE
 COZAAR, LOSARTAN POTASSIUM
 CRESEMBA, ISAVUCONAZONIUM SULFATE
 CRESTOR, ROSUVASTATIN CALCIUM
 CRINONE, PROGESTERONE
 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
 CUTIVATE, FLUTICASONE PROPIONATE
 CUVPOSA, GLYCOPYRROLATE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYANOKIT, HYDROXOCOBALAMIN
 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
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 CYSTADANE, BETAINE
 CYSTADROPS, CYSTEAMINE HYDROCHLORIDE
 CYSTAGON, CYSTEAMINE BITARTRATE
 CYSTARAN, CYSTEAMINE HYDROCHLORIDE
 CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
 CYSTOGRAFIN, DIATRIZOATE MEGLUMINE
 CYSTOGRAFIN DILUTE, DIATRIZOATE MEGLUMINE
 CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE
 CYTALUX, PAFOLACIANINE SODIUM
 CYTARABINE, CYTARABINE
 CYTOMEL, LIOTHYRONINE SODIUM
 CYTOTEC, MISOPROSTOL
 CYTOXAN, CYCLOPHOSPHAMIDE

**** D ****

D.H.E. 45, DIHYDROERGOTAMINE MESYLATE
 DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
 DACARBAZINE, DACARBAZINE
 DACOGEN, DECITABINE
 DACTINOMYCIN, DACTINOMYCIN
 DALFAMPRIDINE, DALFAMPRIDINE
 DALIRESP, ROFLUMILAST
 DALVANCE, DALBAVANCIN HYDROCHLORIDE
 DANAZOL, DANAZOL
 DANTRIUM, DANTROLENE SODIUM

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DANTROLENE SODIUM, DANTROLENE SODIUM
DAPSONE, DAPSONE
DAPTOMYCIN, DAPTOMYCIN
DARAPRIM, PYRIMETHAMINE
DARIFENACIN, DARIFENACIN HYDROBROMIDE
DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
DARTISLA ODT, GLYCOPYRROLATE
DARUNAVIR, DARUNAVIR
DASATINIB, DASATINIB
DASETTA 1/35, ETHINYL ESTRADIOL
DASETTA 7/7/7, ETHINYL ESTRADIOL
DATSCAN, IOFLUPANE I-123
DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
DAURISMO, GLASDEGIB MALEATE
DAYPRO, OXAPROZIN
DAYSEE, ETHINYL ESTRADIOL
DAYTRANA, METHYLPHENIDATE
DAYVIGO, LEMBOREXANT
DDAVP, DESMOPRESSIN ACETATE
DECITABINE, DECITABINE
DEFERASIROX, DEFERASIROX
DEFERIPRONE, DEFERIPRONE
DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
DEFINITY, PERFLUTREN
DEFINITY RT, 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
DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
DEMEROL, MEPERIDINE HYDROCHLORIDE
DEMSE, METYROSINE
DENA VIR, PENCICLOVIR
DEOXYCHOLIC ACID, DEOXYCHOLIC 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
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
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
DESONIDE, DESONIDE

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DESOWEN, DESONIDE
DESOXIMETASONE, DESOXIMETASONE
DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
DESVENLAFAXINE, DESVENLAFAXINE
DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
DETECTNET, COPPER DOTATATE CU-64
DETRAMP SACCHARATE, AMP ASPARTATE, DETROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
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
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
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%, 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%, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, 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

APPENDIX A - PRODUCT NAME INDEX

** D **

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
 DHIVY, CARBIDOPA
 DIABETA, GLYBURIDE
 DIACOMIT, STIRIPENTOL
 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 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 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
 DIAZOXIDE, DIAZOXIDE
 DIBENZYLINE, PHENOXYBENZAMINE HYDROCHLORIDE
 DICLEGIS, DOXYLAMINE SUCCINATE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE), DICYCLOMINE HYDROCHLORIDE
 DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
 DIFFERIN, ADAPALENE (OTC)
 DIFFERIN, ADAPALENE
 DIFICID, FIDAXOMICIN
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 DIFLUCAN, FLUCONAZOLE
 DIFLUNISAL, DIFLUNISAL
 DIFLUPREDNATE, DIFLUPREDNATE
 DIGOXIN, DIGOXIN
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DILANTIN, PHENYTOIN
 DILANTIN, PHENYTOIN SODIUM
 DILANTIN-125, PHENYTOIN
 DILAUDID, HYDROMORPHONE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE IN DEXTROSE 5%, DILTIAZEM HYDROCHLORIDE
 DIMENHYDRINATE, DIMENHYDRINATE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE

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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
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
DOCOSANOL, DOCOSANOL (OTC)
DOFETILDE, DOFETILIDE
DOFETILIDE, DOFETILIDE
DOJOLVI, TRIHEPTANOIN
DOLISHALE, ETHINYL ESTRADIOL
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
DOVATO, DOLUTEGRAVIR SODIUM
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
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 DTPA, TECHNETIUM TC-99M PENTETATE KIT
DRAXIMAGE MDP-25, TECHNETIUM TC-99M MEDRONATE
DRISDOL, ERGOCALCIFEROL
DRIZALMA SPRINKLE, DULOXETINE HYDROCHLORIDE
DRONABINOL, DRONABINOL
DROPERIDOL, DROPERIDOL
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE
DROXIA, HYDROXYUREA
DROXIDOPA, DROXIDOPA
DSUVIA, SUFENTANIL CITRATE

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DUAC, BENZOYL PEROXIDE
 DUAKLIR PRESSAIR, ACLIDINIUM BROMIDE
 DUAVEE, BAZEDOXIFENE ACETATE
 DUETACT, GLIMEPIRIDE
 DUEXIS, FAMOTIDINE
 DULERA, FORMOTEROL FUMARATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUOBRII, HALOBETASOL PROPIONATE
 DUODOTE, ATROPINE
 DUOPA, CARBIDOPA
 DURACLON, CLONIDINE HYDROCHLORIDE
 DURAMORPH PF, MORPHINE SULFATE
 DURAPREP, IODINE POVACRYLEX (OTC)
 DUREZOL, DIFLUPREDNATE
 DURLAZA, ASPIRIN
 DURYSTA, BIMATOPROST
 DUTASTERIDE, DUTASTERIDE
 DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
 DUVOID, BETHANECHOL CHLORIDE
 DYANAVAL XR, AMPHETAMINE
 DYANAVAL XR 10, AMPHETAMINE
 DYANAVAL XR 15, AMPHETAMINE
 DYANAVAL XR 20, AMPHETAMINE
 DYANAVAL XR 5, AMPHETAMINE
 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
 EFAVIRENZ, LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE
 EFFIENT, PRASUGREL HYDROCHLORIDE
 EFINACONAZOLE, EFINACONAZOLE
 EFUDEX, FLUOROURACIL
 EGATEN, TRICLABENDAZOLE
 ELCYS, CYSTEINE HYDROCHLORIDE
 ELEPSIA XR, LEVETIRACETAM
 ELESTAT, EPINASTINE HYDROCHLORIDE
 ELESTRIN, ESTRADIOL
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 ELIDEL, PIMECROLIMUS
 ELIGARD KIT, LEUPROLIDE ACETATE
 ELIGLUSTAT TARTRATE, ELIGLUSTAT TARTRATE
 ELINEST, ETHINYL ESTRADIOL
 ELIQUIS, APIXABAN
 ELIXOPHYLLIN, THEOPHYLLINE

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** E **

ELLA, ULIPRISTAL ACETATE
ELLENCEN, EPIRUBICIN HYDROCHLORIDE
ELLIOTTS B SOLUTION, CALCIUM CHLORIDE
ELMIRON, PENTOSAN POLYSULFATE SODIUM
ELOCON, MOMETASONE FUROATE
ELOXATIN, OXALIPLATIN
ELURYNG, ETHINYL ESTRADIOL
ELYXYB, CELECOXIB
EMBELINE, CLOBETASOL PROPIONATE
EMBELINE E, CLOBETASOL PROPIONATE
EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM
EMEND, APREPITANT
EMEND, FOSAPREPITANT DIMEGLUMINE
EMERPHED, EPHEDRINE SULFATE
EMFLAZA, DEFLAZACORT
EMLA, LIDOCAINE
EMPAVELI, PEGCETACOPLAN
EMSAM, SELEGILINE
EMTRICITABINE, EMTRICITABINE
EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
EMTRIVA, EMTRICITABINE
EMVERM, MEBENDAZOLE
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
ENTADFI, FINASTERIDE
ENTECAVIR, ENTECAVIR
ENTEREG, ALVIMOPAN
ENTERO VU 24%, BARIUM SULFATE
ENTOCORT EC, BUDESONIDE
ENTRESTO, SACUBITRIL
ENVARUSUS XR, TACROLIMUS
ENZALUTAMIDE, ENZALUTAMIDE
EOVIST, GADOXETATE DISODIUM
EPANED, ENALAPRIL MALEATE
EPCLUSA, SOFOSBUVIR
EPHEDRINE SULFATE, EPHEDRINE SULFATE
EPIDIOLEX, CANNABIDIOL
EPIDUO, ADAPALENE
EPIDUO FORTE, ADAPALENE
EPIFOAM, HYDROCORTISONE ACETATE
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
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
EPRONTIA, TOPIRAMATE
EPTIFIBATIDE, EPTIFIBATIDE
EPZICOM, ABACAVIR SULFATE
EQUETRO, CARBAMAZEPINE

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ERAXIS, ANIDULAFUNGIN
ERGOCALCIFEROL, ERGOCALCIFEROL
ERGOLOID MESYLATES, ERGOLOID MESYLATES
ERGOMAR, ERGOTAMINE TARTRATE
ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
ERIVEDGE, VISMODEGIB
ERLEADA, APALUTAMIDE
ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
ERRIN, NORETHINDRONE
ERTACZO, SERTACONAZOLE NITRATE
ERTAPENEM SODIUM, ERTAPENEM SODIUM
ERY-TAB, ERYTHROMYCIN
ERYC, ERYTHROMYCIN
ERYGEL, ERYTHROMYCIN
ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHRA-DERM, ERYTHROMYCIN
ERYTHROCIN, ERYTHROMYCIN LACTOBIONATE
ERYTHROCIN STEARATE, ERYTHROMYCIN STEARATE
ERYTHROMYCIN, ERYTHROMYCIN
ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN LACTOBIONATE, ERYTHROMYCIN LACTOBIONATE
ESBRIET, PIRFENIDONE
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
ESLICARBAZEPINE ACETATE, ESLICARBAZEPINE ACETATE
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
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
ESZOPICLONE, ESZOPICLONE
ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
ETHACRYNIC ACID, ETHACRYNIC ACID
ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
ETHAMOLIN, ETHANOLAMINE OLEATE
ETHINYL ESTRADIOL AND NORELGESTROMIN, ETHINYL ESTRADIOL
ETHINYL ESTRADIOL; ETONOGESTREL, ETHINYL ESTRADIOL
ETHOSUXIMIDE, ETHOSUXIMIDE
ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
ETHYOL, AMIFOSTINE
ETODOLAC, ETODOLAC
ETOMIDATE, ETOMIDATE
ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
ETOPOSIDE, ETOPOSIDE
ETRAVIRINE, ETRAVIRINE
EUCRISA, CRISABOROLE
EURAX, CROTAMITON
EUTHYROX, LEVOTHYROXINE SODIUM **
EVAMIST, ESTRADIOL
EVEKEO, AMPHETAMINE SULFATE
EVEKEO ODT, AMPHETAMINE SULFATE
EVEROLIMUS, EVEROLIMUS
EVISTA, RALOXIFENE HYDROCHLORIDE
EVOCLIN, CLINDAMYCIN PHOSPHATE
EVOMELA, MELPHALAN HYDROCHLORIDE

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EVOTAZ, ATAZANAVIR SULFATE
 EVOXAC, CEVIMELINE HYDROCHLORIDE
 EVRYSDI, RISDIPLAM
 EXCEDRIN (MIGRAINE), ACETAMINOPHEN (OTC)
 EXELDERM, SULCONAZOLE NITRATE
 EXELON, RIVASTIGMINE
 EXEM FOAM KIT, AIR POLYMER-TYPE A
 EXEMESTANE, EXEMESTANE
 EXFORGE, AMLODIPINE BESYLATE
 EXFORGE HCT, AMLODIPINE BESYLATE
 EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)
 EXJADE, DEFERASIROX
 EXKIVITY, MOBOCERTINIB SUCCINATE
 EXONDYS 51, ETEPLIRSEN
 EXPAREL, BUPIVACAINE
 EXSERVAN, RILUZOLE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 EXTINA, KETOCONAZOLE
 EXTRANEAL, ICODextrin
 EYSUVIS, LOTEPREDNOL ETABONATE
 EZALLOR SPRINKLE, ROSUVASTATIN CALCIUM
 EZETIMIBE, EZETIMIBE
 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
 FASLODEX, FULVESTRANT
 FEBUXOSTAT, FEBUXOSTAT
 FELBAMATE, FELBAMATE
 FELBATOL, FELBAMATE
 FELDENE, PIROXICAM
 FELODIPINE, FELODIPINE
 FEMARA, LETROZOLE
 FEMRING, ESTRADIOL ACETATE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FENOGLIDE, FENOFIBRATE
 FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
 FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
 FENSOLVI KIT, LEUPROLIDE ACETATE
 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

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** F **

FERRIPROX, DEFERIPRONE
 FERRLECIT, FERRIC OXYHYDROXIDE
 FERUMOXYTOL, FERUMOXYTOL
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FETROJA, CEFIDEROCOL SULFATE TOSYLATE
 FETZIMA, LEVOMILNACIPRAN HYDROCHLORIDE
 FEXINIDAZOLE, FEXINIDAZOLE
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FIBRICOR, FENOFIBRIC ACID
 FINACEA, AZELAIC ACID
 FINASTERIDE, FINASTERIDE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FINTEPLA, FENFLURAMINE HYDROCHLORIDE
 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 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
 FLUOCINONIDE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUORESCIN SODIUM AND BENOXINATE HYDROCHLORIDE, BENOXINATE HYDROCHLORIDE
 FLUORESCITE, FLUORESCIN SODIUM
 FLUORODOPA F18, FLUORODOPA F-18
 FLUOROURACIL, FLUOROURACIL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FLURANDRENOLIDE, FLURANDRENOLIDE

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FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 FLURBIPROFEN, FLURBIPROFEN
 FLURBIPROFEN SODIUM, FLURBIPROFEN SODIUM
 FLUTAMIDE, FLUTAMIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 FLUTICASONE PROPIONATE AND SALMETEROL XINAFOATE, 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
 FOLOTYN, PRALATREXATE
 FOMEPIZOLE, FOMEPIZOLE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 FORANE, ISOFLURANE
 FORFIVO XL, BUPROPION HYDROCHLORIDE
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 FORTAMET, METFORMIN HYDROCHLORIDE
 FORTAZ, CEFTAZIDIME
 FORTEO, TERIPARATIDE
 FORTESTA, TESTOSTERONE
 FOSAMAX, ALENDRONATE SODIUM
 FOSAMAX PLUS D, ALENDRONATE SODIUM
 FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FOSCARNET SODIUM, FOSCARNET SODIUM
 FOSCAVIR, FOSCARNET SODIUM
 FOSFOMYCIN TROMETHAMINE, FOSFOMYCIN TROMETHAMINE
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FOSRENOL, LANTHANUM CARBONATE
 FOTIVDA, TIVOZANIB HYDROCHLORIDE
 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
 FULVESTRANT, FULVESTRANT
 FULVICIN P/G, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 FULVICIN P/G 165, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 FULVICIN P/G 330, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 FULVICIN-U/F, GRISEOFULVIN, MICROCRYSTALLINE
 FURADANTIN, NITROFURANTOIN
 FUROSEMIDE, FUROSEMIDE
 FUSILEV, LEVOLEUCOVORIN CALCIUM
 FUZEON, ENFUVIRTIDE
 FYARRO, SIROLIMUS
 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

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GALLIUM DOTATOC GA 68, GALLIUM DOTATOC GA-68
GALLIUM GA 68 PSMA-11, GALLIUM GA-68 PSMA-11
GALZIN, ZINC ACETATE
GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
GANIRELIX ACETATE, GANIRELIX ACETATE
GANZYK-RTU, GANCICLOVIR
GASTROCROM, CROMOLYN SODIUM
GASTROGRAFIN, DIATRIZOATE MEGLUMINE
GATIFLOXACIN, GATIFLOXACIN
GATTEX KIT, TEDUGLUTIDE RECOMBINANT
GAVISCON, ALUMINUM HYDROXIDE (OTC)
GAVRETO, PRALSETINIB
GELNIQUE, OXYBUTYNIN CHLORIDE
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
GEMFIBROZIL, GEMFIBROZIL
GEMIFLOXACIN MESYLATE, GEMIFLOXACIN MESYLATE
GEMMILY, ETHINYL ESTRADIOL
GEMTESA, VIBEGRON
GENERLAC, LACTULOSE
GENGRAF, CYCLOSPORINE
GENOPTIC, GENTAMICIN SULFATE
GENOSYL, NITRIC OXIDE
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
GILDAGIA, ETHINYL ESTRADIOL
GILDESS 24 FE, ETHINYL ESTRADIOL
GILENYA, FINGOLIMOD HYDROCHLORIDE
GILOTRIF, AFATINIB DIMALEATE
GIMOTI, METOCLOPRAMIDE HYDROCHLORIDE
GIVLAARI, GIVOSIRAN SODIUM
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
GLOPERBA, COLCHICINE
GLUCAGEN, GLUCAGON HYDROCHLORIDE
GLUCAGON, GLUCAGON
GLUCAGON, GLUCAGON 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
GLYCOPYRROLATE, GLYCOPYRROLATE
GLYDO, LIDOCAINE HYDROCHLORIDE
GLYNASE, GLYBURIDE
GLYRX-PF, GLYCOPYRROLATE
GLYSET, MIGLITOL
GLYXAMBI, EMPAGLIFLOZIN
GOCOVRI, AMANTADINE HYDROCHLORIDE
GOLYTELY, POLYETHYLENE GLYCOL 3350
GOPRELTO, COCAINE HYDROCHLORIDE

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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)
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 GVOKE HYPOPEN, GLUCAGON
 GVOKE PFS, GLUCAGON
 GYNAZOLE-1, BUTOCONAZOLE NITRATE

**** H ****

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
 HALCINONIDE, HALCINONIDE
 HALCION, TRIAZOLAM
 HALDOL, HALOPERIDOL DECANOATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HALOG, HALCINONIDE
 HALOPERIDOL, HALOPERIDOL
 HALOPERIDOL, HALOPERIDOL LACTATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HARVONI, LEDIPASVIR
 HEATHER, NORETHINDRONE
 HECTOROL, DOXERCALCIFEROL
 HEMABATE, CARBOPROST TROMETHAMINE
 HEMADY, DEXAMETHASONE
 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
 HEPSERA, ADEFOVIR DIPIVOXIL
 HER STYLE, LEVONORGESTREL (OTC)
 HETLIOZ, TASIMELTEON
 HETLIOZ LQ, TASIMELTEON
 HIBICLENS, CHLORHEXIDINE GLUCONATE (OTC)
 HIBISTAT, CHLORHEXIDINE GLUCONATE (OTC)
 HICON, SODIUM IODIDE I-131
 HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE
 HORIZANT, GABAPENTIN ENACARBIL
 HYCAMTIN, TOPOTECAN HYDROCHLORIDE
 HYCODAN, HOMATROPINE METHYLBROMIDE
 HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDREA, HYDROXYUREA
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROCODONE, HYDROCODONE BITARTRATE

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HYDROCODONE BITARTRATE, HYDROCODONE BITARTRATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROCODONE BITARTRATE AND IBUPROFEN, 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
 HYSINGLA ER, HYDROCODONE BITARTRATE
 HYZAAR, HYDROCHLOROTHIAZIDE

**** I ****

IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IBRANCE, PALBOCICLIB
 IBUPROFEN, IBUPROFEN (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN AND FAMOTIDINE, FAMOTIDINE
 IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN LYSINE, IBUPROFEN LYSINE
 IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)
 IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
 IC-GREEN, INDOCYANINE GREEN
 ICATIBANT ACETATE, ICATIBANT ACETATE
 ICLEVIA, ETHINYL ESTRADIOL
 ICLUSIG, PONATINIB HYDROCHLORIDE
 ICOSAPENT ETHYL, ICOSAPENT ETHYL
 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
 ILLUCCIX, GALLIUM GA-68 GOZETOTIDE
 ILUVIEN, FLUOCINOLONE ACETONIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IMBRUVICA, IBRUTINIB
 IMCIVREE, SETMELANOTIDE ACETATE
 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-SYMPTOM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)
 IMPAVIDO, MILTEFOSINE
 IMPEKLO, CLOBETASOL PROPIONATE
 IMPOYZ, CLOBETASOL PROPIONATE

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IMURAN, AZATHIOPRINE
IMVEXXY, ESTRADIOL
INAPSINE, DROPERIDOL
INBRIJA, LEVODOPA
INCASSIA, NORETHINDRONE
INCRUSE ELLIPTA, UMECLIDINIUM BROMIDE
INDAPAMIDE, INDAPAMIDE
INDERAL LA, PROPRANOLOL HYDROCHLORIDE
INDIUM IN 111 CHLORIDE, INDIUM IN-111 CHLORIDE
INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE
INDOCIN, INDOMETHACIN
INDOCYANINE GREEN, INDOCYANINE GREEN
INDOMETHACIN, INDOMETHACIN
INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
INFANT'S ADVIL, IBUPROFEN (OTC)
INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
INFED, FERRIC OXYHYDROXIDE
INFUGEM, GEMCITABINE HYDROCHLORIDE
INFUMORPH, MORPHINE SULFATE
INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE
INFUVITE PEDIATRIC, ASCORBIC ACID
INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID
INGREZZA, VALBENAZINE TOSYLATE
INJECTAFER, FERRIC CARBOXYMALTOSE
INLYTA, AXITINIB
INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
INOMAX, NITRIC OXIDE
INQOVI, CEDAZURIDINE
INREBIC, FEDRATINIB HYDROCHLORIDE
INSPIRA, EPLERENONE
INTELENCE, ETRAVIRINE
INTRALIPID 10%, SOYBEAN OIL
INTRALIPID 20%, SOYBEAN OIL
INTRALIPID 30%, SOYBEAN OIL
INTRAROSA, PRASTERONE
INTROVALE, ETHINYL ESTRADIOL
INTUNIV, GUANFACINE HYDROCHLORIDE
INVANZ, ERTAPENEM SODIUM
INVEGA, PALIPERIDONE
INVEGA HAFYERA, PALIPERIDONE PALMITATE
INVEGA SUSTENNA, PALIPERIDONE PALMITATE
INVEGA TRINZA, PALIPERIDONE PALMITATE
INVELTYS, LOTEPREDNOL ETABONATE
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

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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
 ISTURISA, OSILODROSTAT PHOSPHATE
 ISUPREL, ISOPROTERENOL HYDROCHLORIDE
 ITRACONAZOLE, ITRACONAZOLE
 IVABRADINE HYDROCHLORIDE, IVABRADINE HYDROCHLORIDE
 IVERMECTIN, IVERMECTIN (OTC)
 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
 JATENZO, TESTOSTERONE UNDECANOATE
 JELMYTO, MITOMYCIN
 JENCYCLA, NORETHINDRONE
 JENTADUETO, LINAGLIPTIN
 JENTADUETO XR, LINAGLIPTIN
 JEVTANA KIT, CABAZITAXEL
 JORNAY PM, METHYLPHENIDATE HYDROCHLORIDE
 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
 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

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KATERZIA, AMLODIPINE BENZOATE
 KAZANO, ALOGLIPTIN BENZOATE
 KELNOR, ETHINYL ESTRADIOL
 KENALOG, TRIAMCINOLONE ACETONIDE
 KENALOG-10, TRIAMCINOLONE ACETONIDE
 KENALOG-40, TRIAMCINOLONE ACETONIDE
 KENALOG-80, TRIAMCINOLONE ACETONIDE
 KENGREAL, CANGRELOR
 KEPPRA, LEVETIRACETAM
 KEPPRA XR, LEVETIRACETAM
 KERENDIA, FINERENONE
 KERYDIN, TAVABOROLE
 KETALAR, KETAMINE HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 KETOCONAZOLE, KETOCONAZOLE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 KETOROLAC TROMETHAMINE AND PHENYLEPHRINE HYDROCHLORIDE, KETOROLAC TROMETHAMINE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 KETOZOLE, KETOCONAZOLE
 KEVEYIS, DICHLORPHENAMIDE
 KHAPZORY, LEVOLEUCOVORIN
 KIMIDESS, DESOGESTREL
 KIMYRSA, ORITAVANCIN DIPHOSPHATE
 KINEVAC, SINCALIDE
 KIONEX, SODIUM POLYSTYRENE SULFONATE
 KISQALI, RIBOCICLIB SUCCINATE
 KISQALI FEMARA CO-PACK (COPACKAGED), LETROZOLE
 KITABIS PAK, TOBRAMYCIN
 KLARON, SULFACETAMIDE SODIUM
 KLISYRI, TIRBANIBULIN
 KLONOPIN, CLONAZEPAM
 KLOR-CON, POTASSIUM CHLORIDE
 KLOR-CON M10, POTASSIUM CHLORIDE
 KLOR-CON M15, POTASSIUM CHLORIDE
 KLOR-CON M20, POTASSIUM CHLORIDE
 KLOXXADO, NALOXONE HYDROCHLORIDE
 KOMBIGLYZE XR, METFORMIN HYDROCHLORIDE
 KORLYM, MIFEPRISTONE
 KORSUVA, DIFELIKEFALIN ACETATE
 KOSELUGO, SELUMETINIB SULFATE
 KOVANAZE, OXYMETAZOLINE HYDROCHLORIDE
 KRINTAFEL, TAFENOQUINE SUCCINATE
 KURVELO, ETHINYL ESTRADIOL
 KUVAN, SAPROPTERIN DIHYDROCHLORIDE
 KYBELLA, DEOXYCHOLIC ACID
 KYLEENA, LEVONORGESTREL
 KYNMOBI, APOMORPHINE HYDROCHLORIDE
 KYPROLIS, CARFILZOMIB

**** L ****

LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE IN DEXTROSE, LABETALOL HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE IN SODIUM CHLORIDE, 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 AT, TERBINAFINE (OTC)
 LAMISIL AT, TERBINAFINE HYDROCHLORIDE (OTC)

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** L **

LAMIVUDINE, LAMIVUDINE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMOTRIGINE, LAMOTRIGINE
 LAMPIT, NIFURTIMOX
 LANORINAL, ASPIRIN
 LANOXIN, DIGOXIN
 LANOXIN PEDIATRIC, DIGOXIN
 LANREOTIDE ACETATE, LANREOTIDE ACETATE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED), AMOXICILLIN
 LANTHANUM CARBONATE, LANTHANUM CARBONATE
 LAPATINIB DITOSYLATE, LAPATINIB DITOSYLATE
 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 (OTC)
 LATANOPROST, LATANOPROST
 LATISSE, BIMATOPROST
 LATUDA, LURASIDONE HYDROCHLORIDE
 LAX-LYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
 LAZANDA, FENTANYL CITRATE
 LEFLUNOMIDE, LEFLUNOMIDE
 LENALIDOMIDE, LENALIDOMIDE
 LENVIMA, LENVATINIB MESYLATE
 LEQVIO, INCLISIRAN SODIUM
 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
 LEVETIRACETAM, LEVETIRACETAM
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVITRA, VARDENAFIL HYDROCHLORIDE
 LEVO-T, LEVOTHYROXINE SODIUM **
 LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
 LEVOCARNITINE, LEVOCARNITINE
 LEVOCARNITINE SF, LEVOCARNITINE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOCETIRIZINE HYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 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 **

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** L **

LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE
LEXAPRO, ESCITALOPRAM OXALATE
LEXETTE, HALOBETASOL PROPIONATE
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
LINAGLIPTIN, LINAGLIPTIN
LINAGLIPTIN AND METFORMIN HYDROCHLORIDE, LINAGLIPTIN
LINCOCIN, LINCOMYCIN HYDROCHLORIDE
LINCOMYCIN, LINCOMYCIN
LINCOMYCIN, LINCOMYCIN HYDROCHLORIDE
LINDANE, LINDANE
LINEZOLID, LINEZOLID
LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, LINEZOLID
LINZESS, LINACLOTIDE
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
LIVMARLI, MARALIXIBAT CHLORIDE
LIVTENCITY, MARIBAVIR
LO LOESTRIN FE, ETHINYL ESTRADIOL
LO SIMPESE, ETHINYL ESTRADIOL
LO-MALMOREDE, ETHINYL ESTRADIOL
LO-ZUMANDIMINE, DROSPIRENONE
LOCOID, HYDROCORTISONE BUTYRATE
LOCOID LIPOCREAM, HYDROCORTISONE BUTYRATE
LODOSYN, CARBIDOPA
LOESTRIN 21 1.5/30, ETHINYL ESTRADIOL
LOESTRIN 21 1/20, ETHINYL ESTRADIOL
LOESTRIN FE 1.5/30, ETHINYL ESTRADIOL
LOESTRIN FE 1/20, ETHINYL ESTRADIOL
LOGILIA, ULIPRISTAL ACETATE
LOKELMA, SODIUM ZIRCONIUM CYCLOSILICATE

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** L **

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
 LOPINAVIR AND RITONAVIR, LOPINAVIR
 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
 LOREEV XR, LORAZEPAM
 LORYNA, DROSPIRENONE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSEASONIQUE, ETHINYL ESTRADIOL
 LOTEMAX, LOTEPIREDNOL ETABONATE
 LOTEMAX SM, LOTEPIREDNOL ETABONATE
 LOTENSIN, BENAZEPRIL HYDROCHLORIDE
 LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
 LOTEPIREDNOL ETABONATE, LOTEPIREDNOL ETABONATE
 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
 LUBIPROSTONE, LUBIPROSTONE
 LUCEMYRA, LOFEXIDINE HYDROCHLORIDE
 LUMAKRAS, SOTORASIB
 LUMASON, SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES
 LUMI-SPORYN, BACITRACIN ZINC
 LUMIFY, BRIMONIDINE TARTRATE (OTC)
 LUMIGAN, BIMATOPROST
 LUNESTA, ESZOPICLONE
 LUPKYNIS, VOCLOSPORIN
 LUPRON DEPOT, LEUPROLIDE ACETATE
 LUPRON DEPOT-PED KIT, LEUPROLIDE ACETATE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 LUTATHERA, LUTETIUM DOTATATE LU-177
 LUVOX, FLUVOXAMINE MALEATE
 LUXIQ, BETAMETHASONE VALERATE
 LUZU, LULICONAZOLE
 LYBALVI, OLANZAPINE
 LYMPHOSEEK KIT, TECHNETIUM TC-99M TILMANOCEPT
 LYNPARZA, OLAPARIB
 LYRICA, PREGABALIN
 LYRICA CR, PREGABALIN
 LYSODREN, MITOTANE
 LYSTEDA, TRANEXAMIC ACID
 LYVISPAH, BACLOFEN

** M **

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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
MAFENIDE ACETATE, MAFENIDE ACETATE
MAGNESIUM SULFATE, MAGNESIUM SULFATE
MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
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
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
MATULANE, PROCARBAZINE HYDROCHLORIDE
MAVENCLAD, CLADRIBINE
MAVYRET, GLECAPREVIR
MAXALT, RIZATRIPTAN BENZOATE
MAXALT-MLT, RIZATRIPTAN BENZOATE
MAXIDEX, DEXAMETHASONE
MAXIPIME, CEFEPIME HYDROCHLORIDE
MAXITROL, DEXAMETHASONE
MAXZIDE, HYDROCHLOROTHIAZIDE
MAXZIDE-25, HYDROCHLOROTHIAZIDE
MAYZENT, SIPONIMOD FUMARIC ACID
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
MEGACE ES, MEGESTROL ACETATE
MEGESTROL ACETATE, MEGESTROL ACETATE
MEKINIST, TRAMETINIB DIMETHYL SULFOXIDE
MEKTOVI, BINIMETINIB
MELAMISA, DROSPIRENONE
MELOXICAM, MELOXICAM
MELPHALAN, MELPHALAN
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
MEMBRANEBLUE, TRYPAN BLUE
MEN'S ROGAINE, MINOXIDIL (OTC)
MENEST, ESTROGENS, ESTERIFIED
MENOSTAR, ESTRADIOL
MENTAX, BUTENAFINE HYDROCHLORIDE

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MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
 MEPHYTON, PHYTONADIONE
 MEPROMAMATE, MEPROMAMATE
 MEPRON, ATOVAQUONE
 MERCAPTOPYRINE, MERCAPTOPYRINE
 MEROPENEM, MEROPENEM
 MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER, MEROPENEM
 MERREM, MEROPENEM
 MERZEE, ETHINYL ESTRADIOL
 MESALAMINE, MESALAMINE
 MESNA, MESNA
 MESNEX, MESNA
 MESTINON, PYRIDOSTIGMINE BROMIDE
 METADATE CD, METHYLPHENIDATE HYDROCHLORIDE
 METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
 METARAMINOL BITARTRATE, METARAMINOL BITARTRATE
 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
 METHOTREXATE PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOXSALEN, METHOXSALEN
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 METHYLDOPA, METHYLDOPA
 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
 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
 METRONIDAZOLE, METRONIDAZOLE
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 METYROSINE, METYROSINE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MIACALCIN, CALCITONIN SALMON
 MIBELAS 24 FE, ETHINYL ESTRADIOL
 MICAFUNGIN, MICAFUNGIN SODIUM
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 MICARDIS, TELMISARTAN
 MICARDIS HCT, HYDROCHLOROTHIAZIDE

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** M **

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
 MICROGESTIN 1.5/30, ETHINYL ESTRADIOL
 MICROGESTIN 1/20, ETHINYL ESTRADIOL
 MICROGESTIN FE 1.5/30, ETHINYL ESTRADIOL
 MICROGESTIN FE 1/20, ETHINYL ESTRADIOL
 MICROZIDE, HYDROCHLOROTHIAZIDE
 MIDAMOR, AMILORIDE HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM IN 0.9% SODIUM CHLORIDE, MIDAZOLAM
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MIDOL LIQUID GELS, IBUPROFEN (OTC)
 MIFEPREX, MIFEPRISTONE
 MIFEPRISTONE, 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
 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
 MIVACURIUM CHLORIDE, MIVACURIUM CHLORIDE
 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)

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MONISTAT 3 COMBINATION PACK (PREFILLED), MICONAZOLE NITRATE (OTC)
 MONISTAT 7, MICONAZOLE NITRATE (OTC)
 MONISTAT 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MONO-LINYAH, ETHINYL ESTRADIOL
 MONOBASIC SODIUM PHOSPHATE AND DIBASIC SODIUM PHOSPHATE, SODIUM PHOSPHATE, DIBASIC,
 MONODOX, DOXYCYCLINE
 MONOFERRIC, FERRIC DERISOMALTOSE
 MONOKET, ISOSORBIDE MONONITRATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MONUROL, FOSFOMYCIN TROMETHAMINE
 MORPHINE SULFATE, MORPHINE SULFATE
 MOTTEGRITY, 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
 MULTRYS, CUPRIC SULFATE
 MUPIROCIN, MUPIROCIN
 MUPIROCIN, MUPIROCIN CALCIUM
 MUSE, ALPROSTADIL
 MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE
 MYCAMINE, MICAFUNGIN SODIUM
 MYCAPSSA, OCTREOTIDE ACETATE
 MYCOBUTIN, RIFABUTIN
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 MYDAYIS, AMPHETAMINE ASPARTATE
 MYDRIACYL, TROPICAMIDE
 MYFEMBREE, ESTRADIOL
 MYFORTIC, MYCOPHENOLIC SODIUM
 MYKACET, NYSTATIN
 MYLERAN, BUSULFAN
 MYORISAN, ISOTRETINOIN
 MYOVIEV 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT
 MYRBETRIQ, MIRABEGRON
 MYRBETRIQ GRANULES, MIRABEGRON
 MYSOLINE, PRIMIDONE
 MYTESI, CROFELEMER
 MYZILRA, ETHINYL ESTRADIOL

**** N ****

NABUMETONE, NABUMETONE
 NADOLOL, NADOLOL
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 NAFTIN, NAFTIFINE HYDROCHLORIDE
 NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE
 NALFON, FENOPROFEN CALCIUM
 NALLPEN IN PLASTIC CONTAINER, NAFCILLIN SODIUM

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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
 NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 NAPHCN-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
 NAPRELAN, NAPROXEN SODIUM
 NAPROSYN, NAPROXEN
 NAPROXEN, NAPROXEN
 NAPROXEN AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 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)
 NASACORT AQ, TRIAMCINOLONE ACETONIDE
 NASCOBAL, CYANOCOBALAMIN
 NATACYN, NATAMYCIN
 NATAZIA, DIENOGEST
 NATEGLINIDE, NATEGLINIDE
 NATESTO, TESTOSTERONE
 NATROBA, SPINOSAD
 NAYZILAM, MIDAZOLAM
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NEBUPENT, PENTAMIDINE ISETHIONATE
 NEDOCROMIL SODIUM, NEDOCROMIL SODIUM
 NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 NELARABINE, NELARABINE
 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
 NEOPROFEN, IBUPROFEN LYSINE
 NEORAL, CYCLOSPORINE
 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
 NEVANAC, NEPAFENAC
 NEVIRAPINE, NEVIRAPINE
 NEXAVAR, SORAFENIB TOSYLATE
 NEXESTA FE, ETHINYL ESTRADIOL
 NEXIUM, ESOMEPRAZOLE MAGNESIUM
 NEXIUM 24HR, ESOMEPRAZOLE MAGNESIUM (OTC)

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NEXIUM IV, ESOMEPRAZOLE SODIUM
 NEXLETOL, BEMPEDOIC ACID
 NEXLIZET, BEMPEDOIC ACID
 NEXPLANON, ETNOGESTREL
 NEXTERONE, AMIODARONE HYDROCHLORIDE
 NEXTSTELLIS, DROSPIRENONE
 NIACIN, NIACIN
 NIACOR, NIACIN
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE, 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
 NITAZOXANIDE, NITAZOXANIDE
 NITHIODOLE, SODIUM NITRITE
 NITISINONE, NITISINONE
 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 ANTI-DANDRUFF, KETOCONAZOLE (OTC)
 NOCDURNA, DESMOPRESSIN ACETATE
 NOR-QD, NORETHINDRONE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 NOREPINEPHRINE BITARTRATE IN 5% DEXTROSE, 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
 NORITATE, METRONIDAZOLE
 NORMOCARB HF 25, MAGNESIUM CHLORIDE
 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

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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
 NOURIANZ, ISTRADefylline
 NOXAFIL, POSACONAZOLE
 NOXAFIL POWDERMIX KIT, POSACONAZOLE
 NOXIVENT, NITRIC OXIDE
 NUBEQA, DAROLUTAMIDE
 NUCYNТА, TAPENTADOL HYDROCHLORIDE
 NUCYNТА ER, TAPENTADOL HYDROCHLORIDE
 NUEDEXТА, DEXTROMETHORPHAN HYDROBROMIDE
 NULIBRY, FOSDENOPTERIN HYDROBROMIDE
 NULYTELY, POLYETHYLENE GLYCOL 3350
 NULYTELY-FLAVORED, POLYETHYLENE GLYCOL 3350
 NUMBRINO, COCAINE HYDROCHLORIDE
 NUPLAZID, PIMAVANSERIN TARTRATE
 NURTEC ODT, RIMEGEPANT SULFATE
 NUTRILIPID 10%, SOYBEAN OIL
 NUTRILIPID 20%, SOYBEAN OIL
 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 **

OCALIVA, OBETICHOlic ACID
 OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCUFLOX, OFLOXACIN
 ODEFSEY, EMTRICITABINE
 ODOMZO, SONIDEGIB PHOSPHATE
 OFEV, NINTEDANIB ESYLATE
 OFIRMEV, ACETAMINOPHEN
 OFLOXACIN, OFLOXACIN
 OGEN 5, ESTROPIPATE
 OLANZAPINE, OLANZAPINE
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OLINVYK, OLICERIDINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 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 SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)

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OMIIDRIA, 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
 ONDANSETRON, ONDANSETRON
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONEXTON, BENZOYL PEROXIDE
 ONFI, CLOBAZAM
 ONGENTYS, OPICAPONE
 ONGLYZA, SAXAGLIPTIN HYDROCHLORIDE
 ONIVYDE, IRINOTECAN HYDROCHLORIDE
 ONPATTRO, PATISIRAN SODIUM
 ONSURA, ETHINYL ESTRADIOL
 ONUREG, AZACITIDINE
 ONZETRA XSAIL, SUMATRIPTAN SUCCINATE
 OPCICON ONE-STEP, LEVONORGESTREL (OTC)
 OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
 OPSUMIT, MACITENTAN
 OPTIRAY 300, IOVERSOL
 OPTIRAY 320, IOVERSOL
 OPTIRAY 350, IOVERSOL
 OPTISON, ALBUMIN HUMAN
 OPZELURA, RUXOLITINIB PHOSPHATE
 ORABLOC, ARTICAIN HYDROCHLORIDE
 ORACEA, DOXYCYCLINE
 ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE
 ORAQIX, LIDOCAINE
 ORAVERSE, PHENTOLAMINE MESYLATE
 ORAVIG, MICONAZOLE
 ORBACTIV, ORITAVANCIN DIPHOSPHATE
 ORENITRAM, TREPROSTINIL DIOLAMINE
 ORFADIN, NITISINONE
 ORGOVYX, RELUGOLIX
 ORIAHNN (COPACKAGED), ELAGOLIX SODIUM, ESTRADIOL, NORETHINDRONE ACETATE
 ORILISSA, ELAGOLIX SODIUM
 ORKAMBI, IVACAFTOR
 ORLADEYO, BEROTRALSTAT HYDROCHLORIDE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 ORPHENGESIC FORTE, ASPIRIN
 ORTIKOS, BUDESONIDE
 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

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OTOVEL, CIPROFLOXACIN HYDROCHLORIDE
 OTREXUP, METHOTREXATE
 OXACILLIN SODIUM, OXACILLIN SODIUM
 OXALIPLATIN, OXALIPLATIN
 OXANDROLONE, OXANDROLONE
 OXAPROZIN, OXAPROZIN
 OXAYDO, OXYCODONE HYDROCHLORIDE
 OXAZEPAM, OXAZEPAM
 OXBRYTA, VOXELOTOR
 OXCARBAZEPINE, OXCARBAZEPINE
 OXICONAZOLE NITRATE, OXICONAZOLE NITRATE
 OXISTAT, OXICONAZOLE NITRATE
 OXLUMO, LUMASIRAN SODIUM
 OXSORALEN-ULTRA, METHOXSALLEN
 OXTELLAR XR, OXCARBAZEPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCET, ACETAMINOPHEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ASPIRIN, ASPIRIN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYCONTIN, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 OXYTOCIN, OXYTOCIN
 OXYTROL, OXYBUTYNIN
 OXYTROL FOR WOMEN, OXYBUTYNIN (OTC)
 OZEMPIC, SEMAGLUTIDE
 OZOBAX, BACLOFEN
 OZURDEX, DEXAMETHASONE

**** P ****

PACERONE, AMIODARONE HYDROCHLORIDE
 PACITAXEL, PACLITAXEL
 PACLITAXEL, PACLITAXEL
 PALIPERIDONE, PALIPERIDONE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMELOR, NORTRIPTYLINE HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 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
 PAROEX, CHLORHEXIDINE GLUCONATE
 PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
 PAROXETINE, PAROXETINE HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PAROXETINE MESYLATE, PAROXETINE MESYLATE
 PARSABIV, ETELICALCETIDE
 PASER, AMINOSALICYLIC ACID
 PATADAY ONCE DAILY RELIEF, OLOPATADINE HYDROCHLORIDE (OTC)
 PATADAY TWICE DAILY RELIEF, OLOPATADINE HYDROCHLORIDE (OTC)
 PATANASE, OLOPATADINE HYDROCHLORIDE
 PAXIL, PAROXETINE HYDROCHLORIDE
 PAXIL CR, PAROXETINE HYDROCHLORIDE
 PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE
 PEG 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
 PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL
 PEG-3350, SODIUM SULFATE, SODIUM CHLORIDE, POTASSIUM CHLORIDE, SODIUM ASCORBATE AND
 PEMAZYRE, PEMIGATINIB
 PEMFEXY, PEMETREXED
 PENICILLAMINE, PENICILLAMINE

APPENDIX A - PRODUCT NAME INDEX

** P **

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
PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE
PENTOSTATIN, PENTOSTATIN
PENTOXIFYLLINE, PENTOXIFYLLINE
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
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
PHENYTOIN SODIUM, PHENYTOIN SODIUM
PHEXXI, CITRIC ACID
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
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

APPENDIX A - PRODUCT NAME INDEX

** P **

PIPERACILLIN, PIPERACILLIN SODIUM
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
PIQRAY, ALPELISIB
PIRFENIDONE, PIRFENIDONE
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 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
POLYTRIM, POLYMYXIN B SULFATE
POMALYST, POMALIDOMIDE
PONSTEL, MEFENAMIC ACID
PONVORY, PONESIMOD
PORTIA-28, ETHINYL ESTRADIOL
POSACONAZOLE, POSACONAZOLE
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,
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 CONTAINER,
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
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 CONTAINER,
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 CONTAINER,
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 0.149% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, POTASSIUM
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 CONTAINER,
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 CONTAINER,
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,

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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 0.45%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 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 CONTAINER,
 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 CONTAINER,
 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% AND SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 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,
 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 10MEQ, POTASSIUM CHLORIDE
 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, POTASSIUM CHLORIDE
 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 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 PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% 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, 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 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 0.45% 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)
 POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC
 POVIDONE IODINE, POVIDONE-IODINE (OTC)
 PRADAXA, DABIGATRAN ETEXILATE MESYLATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRAMOSONE, HYDROCORTISONE ACETATE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PRAZIQUANTEL, PRAZIQUANTEL

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PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
PRE-OP, HEXACHLOROPHENE
PRE-OP II, HEXACHLOROPHENE
PRECEDEX, DEXMEDETOMIDINE HYDROCHLORIDE
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
PREGABALIN, PREGABALIN
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
PRESTALIA, AMLODIPINE BESYLATE
PRETOMANID, PRETOMANID
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
PREVYMIS, LETERMOVIR
PREZCOBIX, COBICISTAT
PREZISTA, DARUNAVIR
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
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 DIGIHALER, ALBUTEROL SULFATE
PROAIR HFA, ALBUTEROL SULFATE
PROAIR RESPICLICK, ALBUTEROL SULFATE
PROBALAN, PROBENECID
PROBENECID, PROBENECID
PROBENECID AND COLCHICINE, COLCHICINE
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
PROCALAMINE, AMINO ACIDS
PROCARDIA, NIFEDIPINE
PROCARDIA XL, NIFEDIPINE
PROCHLORPERAZINE, PROCHLORPERAZINE
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE

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** P **

PROCOMP, PROCHLORPERAZINE MALEATE
 PROCTOFOAM HC, HYDROCORTISONE ACETATE
 PROCYSBI, CYSTEAMINE BITARTRATE
 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
 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
 PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
 PROPECIA, FINASTERIDE
 PROPOFOL, PROPOFOL
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 PROSCAR, FINASTERIDE
 PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
 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
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 PULMICORT FLEXHALER, BUDESONIDE
 PULMICORT RESPULES, BUDESONIDE
 PUR-WASH, PURIFIED WATER (OTC)
 PURIFIED CORTROPHIN GEL, CORTICOTROPIN
 PURINETHOL, MERCAPTOPURINE
 PURIXAN, MERCAPTOPURINE
 PYLARIFY, PIFLUFOLASTAT F-18
 PYLERA, BISMUTH SUBCITRATE POTASSIUM
 PYRAZINAMIDE, PYRAZINAMIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
 PYRIMETHAMINE, PYRIMETHAMINE
 PYTEST, UREA, C-14
 PYTEST KIT, UREA, C-14

** Q **

QBRELIS, LISINAPRIL
 QBREXZA, GLYCOPYRRONIUM TOSYLATE
 QDOLO, TRAMADOL HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** Q **

QELBREE, VILOXAZINE HYDROCHLORIDE
 QINLOCK, RIPRETINIB
 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
 QUILLICHEW 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
 QULIPTA, ATOGEPANT
 QUTENZA, CAPSAICIN
 QUZYTIR, CETIRIZINE HYDROCHLORIDE
 QVAR REDIHALER, BECLOMETHASONE DIPROPIONATE

** R **

R-GENE 10, ARGININE HYDROCHLORIDE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RADICAVA, EDARAVONE
 RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE (II)
 RADIOGENIX SYSTEM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RAMELTEON, RAMELTEON
 RAMIPRIL, RAMIPRIL
 RANEXA, RANOLAZINE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE
 RAPAFLO, SILODOSIN
 RAPAMUNE, SIROLIMUS
 RAPIVAB, PERAMIVIR
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 RASUVO, METHOTREXATE
 RAVICTI, GLYCEROL PHENYL BUTYRATE
 RAYALDEE, CALCIFEDIOL
 RAYOS, PREDNISONE
 RAZADYNE ER, GALANTAMINE HYDROBROMIDE
 READI-CAT 2, BARIUM SULFATE
 READI-CAT 2 SMOOTHIE, BARIUM SULFATE
 READYPREP CHG, CHLORHEXIDINE GLUCONATE (OTC)
 REBETOL, RIBAVIRIN
 RECARBRIO, CILASTATIN SODIUM
 RECLAST, ZOLEDRONIC ACID
 RECORLEV, LEVOKETOCONAZOLE
 RECTIV, NITROGLYCERIN
 REDITREX, METHOTREXATE
 REGLAN, METOCLOPRAMIDE HYDROCHLORIDE
 REGONOL, PYRIDOSTIGMINE BROMIDE
 RELENZA, ZANAMIVIR
 RELISTOR, METHYLNALTREXONE BROMIDE
 RELPAX, ELETRIPTAN HYDROBROMIDE
 REMERON, MIRTAZAPINE
 REMERON SOLTAB, MIRTAZAPINE
 REMIFENTANIL HYDROCHLORIDE, REMIFENTANIL HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** R **

REMODULIN, TREPROSTINIL
RENACIDIN, CITRIC ACID
RENAGEL, SEVELAMER HYDROCHLORIDE
RENOVA, TRETINOIN
RENVELA, SEVELAMER CARBONATE
REPAGLINIDE, REPAGLINIDE
RESECTISOL IN PLASTIC CONTAINER, MANNITOL
RESTASIS, CYCLOSPORINE
RESTASIS MULTIDOSE, CYCLOSPORINE
RESTORIL, TEMAZEPAM
RETEVMO, SELPERCATINIB
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, BREXPIPIRAZOLE
REYATAZ, ATAZANAVIR SULFATE
REYVOW, LASMIDITAN SUCCINATE
REZIPRES, EPHEDRINE HYDROCHLORIDE
REZUROCK, BELUMOSUDIL MESYLATE
RHINOCORT ALLERGY, BUDESONIDE (OTC)
RHOFADE, OXYMETAZOLINE HYDROCHLORIDE
RHOPRESSA, NETARSUDIL MESYLATE
RIBAVIRIN, RIBAVIRIN
RIDAURA, AURANOFIN
RIFABUTIN, RIFABUTIN
RIFADIN, RIFAMPIN
RIFAMPIN, RIFAMPIN
RILUTEK, RILUZOLE
RILUZOLE, RILUZOLE
RIMACTANE, RIFAMPIN
RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
RIMSO-50, DIMETHYL SULFOXIDE
RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
RINVOQ, UPADACITINIB
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
ROBINUL, GLYCOPYRROLATE
ROBINUL FORTE, GLYCOPYRROLATE
ROCALTROL, CALCITRIOL
ROCKLATAN, LATANOPROST
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
ROFLUMILAST, ROFLUMILAST
ROGAINE (FOR MEN), MINOXIDIL (OTC)
ROGAINE (FOR WOMEN), MINOXIDIL (OTC)
ROGAINE EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
ROMIDEPSIN, ROMIDEPSIN
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
ROSZET, EZETIMIBE

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** R **

ROWASA, MESALAMINE
ROXICODONE, OXYCODONE HYDROCHLORIDE
ROZEREM, RAMELTEON
ROZLYTREK, ENTRECTINIB
RUBRACA, RUCAPARIB CAMSYLATE
RUBY-FILL, RUBIDIUM CHLORIDE RB-82
RUFINAMIDE, RUFINAMIDE
RUKOBIA, FOSTEMSAVIR TROMETHAMINE
RUZURGI, AMIFAMPRIDINE
RYANODEX, DANTROLENE SODIUM
RYBELSUS, SEMAGLUTIDE
RYDAPT, MIDOSTAURIN
RYTARY, CARBIDOPA
RYTHMOL SR, PROPAFENONE HYDROCHLORIDE

** S **

SABRIL, VIGABATRIN
SAFYRAL, DROSPIRENONE
SALAGEN, PILOCARPINE HYDROCHLORIDE
SALONPAS, MENTHOL (OTC)
SAMSCA, TOLVAPTAN
SANCUSO, GRANISETRON
SANDIMMUNE, CYCLOSPORINE
SANDOSTATIN, OCTREOTIDE ACETATE
SANDOSTATIN LAR, OCTREOTIDE ACETATE
SAPHRIS, ASENAPINE MALEATE
SAPROPTERIN DIHYDROCHLORIDE, SAPROPTERIN DIHYDROCHLORIDE
SAVAYSA, EDOXABAN TOSYLATE
SAVELLA, MILNACIPRAN HYDROCHLORIDE
SAXENDA, LIRAGLUTIDE RECOMBINANT
SCANDONEST L, LEVONORDEFIN
SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
SCEMBLIX, ASCIMINIB HYDROCHLORIDE
SCENESSE, AFAMELANOTIDE
SCLEROSOL, TALC
SCOPOLAMINE, SCOPOLAMINE
SEASONALE, ETHINYL ESTRADIOL
SEASONIQUE, ETHINYL ESTRADIOL
SECUADO, ASENAPINE
SEGLENTIS, CELECOXIB
SEGLUOMET, ERTUGLIFLOZIN
SEIZALAM, MIDAZOLAM HYDROCHLORIDE
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
SELENIUM ACID, SELENIUM ACID
SELENIUM SULFIDE, SELENIUM SULFIDE
SELZENTRY, MARAVIROC
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
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SETLAKIN, ETHINYL ESTRADIOL
SEVELAMER CARBONATE, SEVELAMER CARBONATE
SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE
SEVOFLURANE, SEVOFLURANE
SEYSARA, SARECYCLINE HYDROCHLORIDE
SFROWASA, MESALAMINE
SIGNIFOR, PASIREOTIDE DIASPARTATE
SIGNIFOR LAR KIT, PASIREOTIDE PAMOATE

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** S **

SIKLOS, HYDROXYUREA
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILENOR, DOXEPIN HYDROCHLORIDE
 SILODOSIN, SILODOSIN
 SILVADENE, SILVER SULFADIAZINE
 SIMBRINZA, BRIMONIDINE TARTRATE
 SIMLIYA, DESOGESTREL
 SIMPESE, ETHINYL ESTRADIOL
 SIMVASTATIN, SIMVASTATIN
 SINE-AID IB, IBUPROFEN (OTC)
 SINEMET, CARBIDOPA
 SINGULAIR, MONTELUKAST SODIUM
 SINUVA, MOMETASONE FUROATE
 SIROLIMUS, SIROLIMUS
 SIRTURO, BEDAQUILINE FUMARATE
 SITAVIG, ACYCLOVIR
 SIVEXTRO, TEDIZOLID PHOSPHATE
 SKELAXIN, METAXALONE
 SKLICE, IVERMECTIN (OTC)
 SKYLA, LEVONORGESTREL
 SLYND, DROSPIRENONE
 SMOFLIPID 20%, FISH OIL
 SOAAZ, TORSEMIDE
 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 14.6%, SODIUM CHLORIDE
 SODIUM CHLORIDE 23.4%, SODIUM CHLORIDE
 SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, FERRIC OXYHYDROXIDE
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18
 SODIUM IODIDE I 123, SODIUM IODIDE I-123
 SODIUM IODIDE I 131, SODIUM IODIDE I-131
 SODIUM NITRITE, SODIUM NITRITE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 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 TETRADECYL SULFATE, SODIUM TETRADECYL SULFATE
 SODIUM THIOSULFATE, SODIUM THIOSULFATE
 SOJOURN, SEVOFLURANE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 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
 SONATA, ZALEPLON
 SOOLANTRA, IVERMECTIN
 SORBITOL 3% IN PLASTIC CONTAINER, SORBITOL
 SORBITOL 3.3% IN PLASTIC CONTAINER, SORBITOL
 SORBITOL-MANNITOL IN PLASTIC CONTAINER, MANNITOL
 SORILUX, CALCIPOTRIENE
 SORINE, SOTALOL HYDROCHLORIDE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SOTRADECOL, SODIUM TETRADECYL SULFATE

APPENDIX A - PRODUCT NAME INDEX

** S **

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
 SPRAVATO, ESKETAMINE HYDROCHLORIDE
 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
 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
 STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION
 STERITALC, TALC
 STIE-CORT, HYDROCORTISONE
 STIOLTO RESPIMAT, OLODATEROL HYDROCHLORIDE
 STIVARGA, REGORAFENIB
 STRATTERA, ATOMOXETINE HYDROCHLORIDE
 STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
 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
 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
 SULFAMYLON, MAFENIDE ACETATE
 SULFASALAZINE, SULFASALAZINE
 SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE
 SULINDAC, SULINDAC
 SUMATRIPTAN, SUMATRIPTAN
 SUMATRIPTAN, SUMATRIPTAN SUCCINATE
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SUNITINIB MALATE, SUNITINIB MALATE

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** S **

SUNOSI, SOLRIAMFETOL HYDROCHLORIDE
 SUPPRELIN LA, HISTRELIN ACETATE
 SUPRANE, DESFLURANE
 SUPRAX, CEFIXIME
 SUPREP BOWEL PREP KIT, MAGNESIUM SULFATE
 SUSTIVA, EFAVIRENZ
 SUSTOL, GRANISETRON
 SUTAB, MAGNESIUM SULFATE
 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
 SYNRIPO, OMACETAXINE MEPESUCCINATE
 SYNTHROID, LEVOTHYROXINE SODIUM **
 SYPRINE, TRIENTINE HYDROCHLORIDE

** T **

TAB-PROFEN, IBUPROFEN (OTC)
 TABRECTA, CAPMATINIB HYDROCHLORIDE
 TACLONEX, BETAMETHASONE DIPROPIONATE
 TACROLIMUS, TACROLIMUS
 TADALAFIL, TADALAFIL
 TAFINLAR, DABRAFENIB MESYLATE
 TAFLUPROST, TAFLUPROST
 TAGAMET HB, CIMETIDINE (OTC)
 TAGITOL V, BARIUM SULFATE
 TAGRISSO, OSIMERTINIB MESYLATE
 TALC, TALC
 TALICIA, AMOXICILLIN
 TALZENNA, TALAZOPARIB TOSYLATE
 TAMIFLU, OSELTAMIVIR PHOSPHATE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TARCEVA, ERLOTINIB HYDROCHLORIDE
 TARGRETIN, BEXAROTENE
 TARPEYO, BUDESONIDE
 TASCENSO ODT, FINGOLIMOD LAURYL SULFATE
 TASIGNA, NILOTINIB HYDROCHLORIDE
 TASMAR, TOLCAPONE
 TAUVID, FLORTAUCIPIR F-18
 TAVABOROLE, TAVABOROLE
 TAVALISSE, FOSTAMATINIB DISODIUM
 TAVNEOS, AVACOPAN
 TAXOTERE, DOCETAXEL
 TAYTULLA, ETHINYL ESTRADIOL
 TAZAROTENE, TAZAROTENE
 TAZICEF, CEFTAZIDIME
 TAZORAC, TAZAROTENE
 TAZTIA XT, DILTIAZEM HYDROCHLORIDE
 TAZVERIK, TAZEMETOSTAT HYDROBROMIDE

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** T **

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 SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 TECHNETIUM TC-99M MEBROFENIN, TECHNETIUM TC-99M MEBROFENIN KIT
 TECHNETIUM TC-99M MEDRONATE KIT, TECHNETIUM TC-99M MEDRONATE KIT
 TECHNETIUM TC99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT
 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
 TEMBEXA, BRINCIDOFIVIR
 TEMODAR, TEMOZOLOMIDE
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TEMSIROLIMUS, TEMSIROLIMUS
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TENORETIC 100, ATENOLOL
 TENORETIC 50, ATENOLOL
 TENORMIN, ATENOLOL
 TEPADINA, THIOTEPA
 TEPMETKO, TEPOTINIB HYDROCHLORIDE
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE (OTC)
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE 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
 TETRABENAZINE, TETRABENAZINE
 TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 TEXACORT, HYDROCORTISONE
 THALITONE, CHLORTHALIDONE
 THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
 THALOMID, THALIDOMIDE
 THAM, TROMETHAMINE
 THEO-24, 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
 THIOLA EC, TIOPRONIN
 THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
 THIOTEPA, THIOTEPA
 THIOTHIXENE, THIOTHIXENE

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** T **

THYQUIDITY, LEVOTHYROXINE SODIUM
 THYRO-TABS, LEVOTHYROXINE SODIUM **
 THYROSAFE, 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)
 TIOPRONIN, TIOPRONIN
 TIROFIBAN HYDROCHLORIDE, TIROFIBAN HYDROCHLORIDE
 TIROSINT, LEVOTHYROXINE SODIUM
 TIROSINT-SOL, LEVOTHYROXINE SODIUM
 TIS-U-SOL, MAGNESIUM SULFATE
 TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 TISSUEBLUE, BRILLIANT BLUE G
 TIVICAY, DOLUTEGRAVIR SODIUM
 TIVICAY PD, DOLUTEGRAVIR SODIUM
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOBI, TOBRAMYCIN
 TOBI PODHALER, TOBRAMYCIN
 TOBRADEX, DEXAMETHASONE
 TOBRADEX ST, DEXAMETHASONE
 TOBRAMYCIN, TOBRAMYCIN
 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
 TOFACITINIB, TOFACITINIB CITRATE
 TOFRANIL, IMIPRAMINE HYDROCHLORIDE
 TOLAK, FLUOROURACIL
 TOLCAPONE, TOLCAPONE
 TOLSURA, ITRACONAZOLE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TOLVAPTAN, TOLVAPTAN
 TOPAMAX, TOPIRAMATE
 TOPICORT, DESOXIMETASONE
 TOPIRAMATE, TOPIRAMATE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 TOPROL-XL, METOPROLOL SUCCINATE
 TOREMIFENE CITRATE, TOREMIFENE CITRATE
 TORISEL, TEMSIROLIMUS
 TORSEMIDE, TORSEMIDE
 TOSYMRA, SUMATRIPTAN
 TOVIAZ, FESOTERODINE FUMARATE
 TPN ELECTROLYTES IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 TPOXX, TECOVIRIMAT
 TRACLEER, BOSENTAN
 TRADJENTA, LINAGLIPTIN
 TRALEMENT, CUPRIC SULFATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRANDATE, LABETALOL HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** T **

TRANDOLAPRIL, TRANDOLAPRIL
 TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE, TRANDOLAPRIL
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRANSDERM SCOP, SCOPOLAMINE
 TRANXENE, CLORAZEPATE DIPOTASSIUM
 TRANYLCYPROMINE SULFATE, TRANYLCYPROMINE 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
 TREPROSTINIL, TREPROSTINIL
 TRETINOIN, TRETINOIN
 TREXALL, METHOTREXATE SODIUM
 TREXIMET, NAPROXEN SODIUM
 TREZIX, ACETAMINOPHEN
 TRI LO SPRINTC, 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
 TRI-SPRINTC, ETHINYL ESTRADIOL
 TRIACIN-C, CODEINE PHOSPHATE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIAMTERENE, TRIAMTERENE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIANEX, TRIAMCINOLONE ACETONIDE
 TRIAZOLAM, TRIAZOLAM
 TRIBENZOR, AMLODIPINE BESYLATE
 TRICOR, FENOFIBRATE
 TRIDERM, TRIAMCINOLONE ACETONIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 TRISENCE, TRIAMCINOLONE ACETONIDE
 TRIFERIC, FERRIC PYROPHOSPHATE CITRATE
 TRIFERIC AVNU, FERRIC PYROPHOSPHATE CITRATE
 TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 TRIFLURIDINE, TRIFLURIDINE
 TRIGLIDE, FENOFIBRATE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 TRIJARDY XR, EMPAGLIFLOZIN
 TRIKAFTA (COPACKAGED), ELEXACAF TOR, IVACAFTOR, TEZACAFTOR
 TRILEPTAL, OXCARBAZEPINE
 TRILIPIX, CHOLINE FENOFIBRATE
 TRIMETHOBENZAMIDE HYDROCHLORIDE, 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

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** T **

TRIZIVIR, ABACAVIR SULFATE
 TROKENDI XR, TOPIRAMATE
 TROPHAMINE, AMINO ACIDS
 TROPHAMINE 10%, AMINO ACIDS
 TROPICACYL, TROPICAMIDE
 TROPICAMIDE, TROPICAMIDE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 TRUDHESA, DIHYDROERGOTAMINE MESYLATE
 TRULANCE, PLECANATIDE
 TRUSELTIQ, INFIGRATINIB PHOSPHATE
 TRUSOPT, DORZOLAMIDE HYDROCHLORIDE
 TRUVADA, EMTRICITABINE
 TUDORZA PRESSAIR, ACLIDINIUM BROMIDE
 TUKYSA, TUCATINIB
 TURALIO, PEXIDARTINIB HYDROCHLORIDE
 TUSSICAPS, CHLORPHENIRAMINE POLISTIREX
 TUXARIN ER, CHLORPHENIRAMINE MALEATE
 TUZISTRA XR, CHLORPHENIRAMINE POLISTIREX
 TWIRLA, ETHINYL ESTRADIOL
 TWYNEO, BENZOYL PEROXIDE
 TYBLUME, ETHINYL ESTRADIOL
 TYBOST, COBICISTAT
 TYDEMY, DROSPIRENONE
 TYGACIL, TIGECYCLINE
 TYKERB, LAPATINIB DITOSYLATE
 TYLENOL, ACETAMINOPHEN (OTC)
 TYMLOS, ABALOPARATIDE
 TYRVAYA, VARENICLINE TARTRATE
 TYVASO, TREPROSTINIL
 TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE

** U **

U-CORT, HYDROCORTISONE ACETATE
 UBRELVY, UBROGEPANT
 UCERIS, BUDESONIDE
 UKONIQ, UMBRALISIB TOSYLATE
 ULORIC, FEBUXOSTAT
 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 300, IOPROMIDE
 ULTRAVIST 370, IOPROMIDE
 UNASYN, AMPICILLIN SODIUM
 UNISOM, DOXYLAMINE SUCCINATE (OTC)
 UNITHROID, LEVOTHYROXINE SODIUM **
 UPNEEQ, OXYMETAZOLINE HYDROCHLORIDE
 UPTRAVI, SELEXIPAG
 UREX, METHENAMINE HIPPURATE
 UROCIT-K, POTASSIUM CITRATE
 UROXATRAL, ALFUZOSIN HYDROCHLORIDE
 URSO 250, URSODIOL
 URSO FORTE, URSODIOL
 URSODIOL, URSODIOL
 UVADEX, METHOXSALLEN

** V **

VABOMERE, MEROPENEM
 VAGIFEM, ESTRADIOL
 VAGISTAT-1, TIOCONAZOLE (OTC)
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** v **

VALCHLOR, MECHLORETHAMINE HYDROCHLORIDE
VALCYTE, VALGANCICLOVIR HYDROCHLORIDE
VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
VALIUM, DIAZEPAM
VALNAC, BETAMETHASONE VALERATE
VALPROATE SODIUM, VALPROATE SODIUM
VALPROIC ACID, VALPROIC ACID
VALRUBICIN, VALRUBICIN
VALSARTAN, VALSARTAN
VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
VALSTAR PRESERVATIVE FREE, VALRUBICIN
VALTOCO, DIAZEPAM
VALTRES, VALACYCLOVIR HYDROCHLORIDE
VANCOCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
VANCOMYCIN, VANCOMYCIN
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
VANDAZOLE, METRONIDAZOLE
VANIQA, EFLORNITHINE HYDROCHLORIDE
VANOS, FLUOCINONIDE
VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE
VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
VARENICLINE TARTRATE, VARENICLINE TARTRATE
VARIBAR HONEY, BARIUM SULFATE
VARIBAR NECTAR, BARIUM SULFATE
VARIBAR PUDDING, BARIUM SULFATE
VARIBAR THIN HONEY, BARIUM SULFATE
VARIBAR THIN LIQUID, BARIUM SULFATE
VARITHENA, POLIDOCANOL
VARUBI, ROLAPITANT HYDROCHLORIDE
VASCEPA, ICOSAPENT ETHYL
VASERETIC, ENALAPRIL MALEATE
VASOPRESSIN, VASOPRESSIN
VASOSTRICT, VASOPRESSIN
VASOTEC, ENALAPRIL MALEATE
VAZALORE, ASPIRIN (OTC)
VAZCULEP, PHENYLEPHRINE HYDROCHLORIDE
VECTICAL, CALCITRIOL
VECURONIUM BROMIDE, VECURONIUM BROMIDE
VEKLURY, REMDESIVIR
VELCADE, BORTEZOMIB
VELETRI, EPOPROSTENOL SODIUM
VELIVET, DESOGESTREL
VELPHORO, FERRIC OXYHYDROXIDE
VELTASSA, PATIROMER SORBITE X CALCIUM
VELTIN, CLINDAMYCIN PHOSPHATE
VEMLIDY, TENOFOVIR ALAFENAMIDE FUMARATE
VENCLEXTA, VENETOCLAX
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
VENOFER, FERRIC OXYHYDROXIDE
VENTAVIS, ILOPROST
VENTOLIN HFA, ALBUTEROL SULFATE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
VERARING, ETHINYL ESTRADIOL
VERDESO, DESONIDE
VEREGEN, SINECATECHINS
VERELAN, VERAPAMIL HYDROCHLORIDE
VERELAN PM, VERAPAMIL HYDROCHLORIDE
VERKAZIA, CYCLOSPORINE
VERQUVO, VERICIGUAT
VERSACLOZ, CLOZAPINE
VERZENIO, ABEMACICLIB
VESICARE, SOLIFENACIN SUCCINATE
VESICARE LS, SOLIFENACIN SUCCINATE

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** v **

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
 VIENVA, ETHINYL ESTRADIOL
 VIGABATRIN, VIGABATRIN
 VIGADRONE, VIGABATRIN
 VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE
 VIIBRYD, VILAZODONE HYDROCHLORIDE
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 VILTEPSO, VILTOLARSEN
 VIMOVO, ESOMEPRAZOLE MAGNESIUM
 VIMPAT, LACOSAMIDE
 VINBLASTINE SULFATE, VINBLASTINE SULFATE
 VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE
 VINOELBINE TARTRATE, VINOELBINE TARTRATE
 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
 VITAMIN K1, PHYTONADIONE
 VITRAKVI, LAROTRECTINIB SULFATE
 VIVELLE-DOT, ESTRADIOL
 VIVITROL, NALTREXONE
 VIZAMYL, FLUTEMETAMOL F-18
 VIZIMPRO, DACOMITINIB
 VOCABRIA, CABOTEGRAVIR SODIUM
 VOGELXO, TESTOSTERONE
 VOLNEA, DESOGESTREL
 VOLTAREN ARTHRITIS PAIN, DICLOFENAC SODIUM (OTC)
 VORICONAZOLE, VORICONAZOLE
 VORTIOXETINE HYDROBROMIDE, VORTIOXETINE HYDROBROMIDE
 VOSEVI, SOFOSBUVIR
 VOSOL, ACETIC ACID, GLACIAL
 VOSOL HC, ACETIC ACID, GLACIAL
 VOSPIRE ER, ALBUTEROL SULFATE
 VOTRIENT, PAZOPANIB HYDROCHLORIDE
 VOXZOGO, VOSORITIDE
 VRAYLAR, CARIPRAZINE HYDROCHLORIDE
 VUIITY, PILOCARPINE HYDROCHLORIDE
 VUMERITY, DIROXIMEL FUMARATE
 VUSION, MICONAZOLE NITRATE
 VYFEMLA, ETHINYL ESTRADIOL
 VYLEESI (AUTOINJECTOR), BREMELANOTIDE ACETATE
 VYNDAMAX, TAFAMIDIS
 VYNDAQEL, TAFAMIDIS MEGLUMINE
 VYONDYS 53, GOLODIRSEN

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** V **

VYTORIN, EZETIMIBE
 VYVANSE, LISDEXAMFETAMINE DIMESYLATE
 VYXEOS, CYTARABINE
 VYZULTA, LATANOPROSTENE BUNOD

** W **

WAKIX, PITOLISANT HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM
 WEGOVY, SEMAGLUTIDE
 WELCHOL, COLESEVELAM HYDROCHLORIDE
 WELIREG, BELZUTIFAN
 WELLBUTRIN SR, BUPROPION HYDROCHLORIDE
 WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
 WERA, ETHINYL ESTRADIOL
 WINLEVI, CLASCOTERONE
 WIXELA INHUB, FLUTICASONE PROPIONATE
 WOMEN'S ROGAINE, MINOXIDIL (OTC)
 WYNZORA, BETAMETHASONE DIPROPIONATE

** X **

XACIATO, CLINDAMYCIN PHOSPHATE
 XADAGO, SAFINAMIDE MESYLATE
 XALATAN, LATANOPROST
 XALKORI, CRIZOTINIB
 XANAX, ALPRAZOLAM
 XANAX XR, ALPRAZOLAM
 XARACOLL, BUPIVACAINE HYDROCHLORIDE
 XARELTO, RIVAROXABAN
 XATMEP, METHOTREXATE SODIUM
 XCOPRI, CENOBAMATE
 KELJANZ, TOFACITINIB CITRATE
 KELJANZ XR, TOFACITINIB CITRATE
 XELODA, CAPECITABINE
 XELPROS, LATANOPROST
 XENAZINE, TETRABENAZINE
 XENICAL, ORLISTAT
 XENLETA, LEFAMULIN ACETATE
 XENON XE 133, XENON XE-133
 XEPI, OZENOXACIN
 XERAVA, ERAVACYCLINE DIHYDROCHLORIDE
 XERESE, ACYCLOVIR
 XERMELO, TELOTRISTAT ETIPRATE
 XHANCE, FLUTICASONE PROPIONATE
 XIFAXAN, RIFAXIMIN
 XIGDUO XR, DAPAGLIFLOZIN
 XIIDRA, LIFITEGRAST
 XIMINO, MINOCYCLINE HYDROCHLORIDE
 XIPERE, TRIAMCINOLONE ACETONIDE
 XOFIGO, RADIUM RA-223 DICHLORIDE
 XOFLUZA, BALOXAVIR MARBOXIL
 XOLEGEL, KETOCONAZOLE
 XOPENEX, LEVALBUTEROL HYDROCHLORIDE
 XOPENEX HFA, LEVALBUTEROL TARTRATE
 XOSPATA, GILTERITINIB FUMARATE
 XPOVIO, SELINEXOR
 XTAMPZA ER, OXYCODONE
 XTANDI, ENZALUTAMIDE
 XULANE, ETHINYL ESTRADIOL
 XURIDEN, URIDINE TRIACETATE
 XYLOCAINE, LIDOCAINE HYDROCHLORIDE
 XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
 XYOSTED (AUTOINJECTOR), TESTOSTERONE ENANTHATE
 XYREM, SODIUM OXYBATE
 XYWAV, CALCIUM OXYBATE
 XYZAL ALLERGY 24HR, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)

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** Y **

YAELA, DROSPIRENONE
YASMIN, DROSPIRENONE
YAZ, DROSPIRENONE
YONDELIS, TRABECTEDIN
YONSA, ABIRATERONE ACETATE
YUPELRI, REVEFENACIN
YUTIQ, FLUOCINOLONE ACETONIDE

** Z **

ZADITOR, KETOTIFEN FUMARATE (OTC)
ZAFIRLUKAST, ZAFIRLUKAST
ZALEPLON, ZALEPLON
ZANAFLEX, TIZANIDINE HYDROCHLORIDE
ZANOSAR, STREPTOZOCIN
ZANTAC, RANITIDINE HYDROCHLORIDE
ZARONTIN, ETHOSUXIMIDE
ZAVESCA, MIGLUSTAT
ZEGALOGUE, DASIGLUCAGON HYDROCHLORIDE
ZEGALOGUE (AUTOINJECTOR), DASIGLUCAGON HYDROCHLORIDE
ZEGERID, OMEPRAZOLE
ZEGERID OTC, OMEPRAZOLE (OTC)
ZEJULA, NIRAPARIB TOSYLATE
ZELAPAR, SELEGILINE HYDROCHLORIDE
ZELBORAF, VEMURAFENIB
ZELNORM, TEGASEROD MALEATE
ZEMBRACE SYMTOUCH, SUMATRIPTAN SUCCINATE
ZEMDRI, PLAZOMICIN SULFATE
ZEMPLAR, PARICALCITOL
ZENATANE, ISOTRETINOIN
ZEPATIER, ELBASVIR
ZEPOSIA, OZANIMOD HYDROCHLORIDE
ZEPZELCA, LURBINECTEDIN
ZERBAXA, CEFTOLOZANE SULFATE
ZERVIAE, CETIRIZINE HYDROCHLORIDE
ZESTORETIC, HYDROCHLOROTHIAZIDE
ZESTRIL, LISINAPRIL
ZETIA, EZETIMIBE
ZETONNA, CICLESONIDE
ZIAC, BISOPROLOL FUMARATE
ZIAGEN, ABACAVIR SULFATE
ZIANA, CLINDAMYCIN PHOSPHATE
ZIDOVUDINE, ZIDOVUDINE
ZILEUTON, ZILEUTON
ZILRETTA, TRIAMCINOLONE ACETONIDE
ZILXI, MINOCYCLINE HYDROCHLORIDE
ZIMHI, NALOXONE HYDROCHLORIDE
ZINACEF, CEFUROXIME SODIUM
ZINC CHLORIDE, ZINC CHLORIDE
ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE
ZINC SULFATE, ZINC SULFATE
ZINGO, LIDOCAINE HYDROCHLORIDE
ZIOPTAN, TAFLUPROST
ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
ZIPRASIDONE MESYLATE, ZIPRASIDONE MESYLATE
ZIPSOR, DICLOFENAC POTASSIUM
ZIRGAN, GANCICLOVIR
ZITHROMAX, AZITHROMYCIN
ZOCOR, SIMVASTATIN
ZOFRAN, ONDANSETRON HYDROCHLORIDE
ZOKINVY, LONAFARNIB
ZOLADEX, GOSERELIN ACETATE
ZOLEDRONIC, ZOLEDRONIC ACID
ZOLEDRONIC ACID, ZOLEDRONIC ACID
ZOLINZA, VORINOSTAT
ZOLMITRIPTAN, ZOLMITRIPTAN

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ZOLOFT, SERTRALINE HYDROCHLORIDE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
ZOLPIMIST, ZOLPIDEM TARTRATE
ZOMETA, ZOLEDRONIC ACID
ZOMIG, ZOLMITRIPTAN
ZOMIG-ZMT, ZOLMITRIPTAN
ZONALON, DOXEPIN HYDROCHLORIDE
ZONEGRAN, ZONISAMIDE
ZONISAMIDE, ZONISAMIDE
ZONTIVITY, VORAPAXAR SULFATE
ZORTRESS, EVEROLIMUS
ZORVOLEX, DICLOFENAC
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
ZULRESSO, BREXANOLONE
ZUMANDIMINE, DROSPIRENONE
ZYCLARA, IMIQUIMOD
ZYDELIG, IDELALISIB
ZYFLO, ZILEUTON
ZYFLO CR, ZILEUTON
ZYKADIA, CERITINIB
ZYLET, LOTEPREDNOL ETABONATE
ZYLOPRIM, ALLOPURINOL
ZYMAR, GATIFLOXACIN
ZYMAXID, GATIFLOXACIN
ZYNRELEF KIT, BUPIVACAINE
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 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

AADI

- * AADI BIOSCIENCE INC
FYARRO, SIROLIMUS

AAIPHARMA LLC

- * AAIPHARMA LLC
AZASAN, AZATHIOPRINE

ABBVIE

- * ABBVIE INC
ANDROGEL, TESTOSTERONE
CYCLOSPORINE, CYCLOSPORINE
DEPAKOTE ER, DIVALPROEX SODIUM
DEPAKOTE, DIVALPROEX SODIUM
GENGRAF, CYCLOSPORINE
K-TAB, POTASSIUM CHLORIDE
KALETRA, LOPINAVIR
NIMBEX PRESERVATIVE FREE, CISATRACURIUM BESYLATE
NIMBEX, CISATRACURIUM BESYLATE
NORVIR, RITONAVIR
SYNTHROID, LEVOTHYROXINE SODIUM **
TRICOR, FENOFIBRATE
TRILIPIX, CHOLINE FENOFIBRATE
ULTANE, SEVOFLURANE
ZEMPLAR, PARICALCITOL

ABBVIE ENDOCRINE INC

- * ABBVIE ENDOCRINE INC
LUPRON DEPOT, LEUPROLIDE ACETATE
LUPRON DEPOT-PED KIT, LEUPROLIDE ACETATE

ABBVIE INC

- * ABBVIE INC
DUOPA, CARBIDOPA
MAVYRET, GLECAPREVIR
NORVIR, RITONAVIR
ORIAHNN (COPACKAGED), ELAGOLIX SODIUM, ESTRADIOL, NORETHINDRONE ACETATE
ORILISSA, ELAGOLIX SODIUM
QULIPTA, ATOGEPANT
RINVOQ, UPADACITINIB
VENCLEXTA, VENETOCLAX
VUITY, PILOCARPINE HYDROCHLORIDE

ABHAI INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ABHAI INC**

DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

ABHAI LLC*** ABHAI LLC**

ATOVAQUONE, ATOVAQUONE
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
LEFLUNOMIDE, LEFLUNOMIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
URSODIOL, URSODIOL

ABON PHARMS LLC*** ABON PHARMACEUTICALS LLC**

ATOVAQUONE, ATOVAQUONE
CLOFARABINE, CLOFARABINE

ABRAXIS BIOSCIENCE*** ABRAXIS BIOSCIENCE LLC**

ABRAXANE, PACLITAXEL

ABRAXIS PHARM*** ABRAXIS PHARMACEUTICAL PRODUCTS**

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE

ACACIA*** ACACIA PHARMA LTD**

BARHEMSYS, AMISULPRIDE
BYFAVO, REMIMAZOLAM BESYLATE

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
APIXABAN, APIXABAN
ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
ARIPIPRAZOLE, ARIPIPRAZOLE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
ATROPINE SULFATE, ATROPINE SULFATE
AZACITIDINE, AZACITIDINE
BICALUTAMIDE, BICALUTAMIDE
BIVALIRUDIN, BIVALIRUDIN
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CABAZITAXEL, CABAZITAXEL
CAPECITABINE, CAPECITABINE
CARBIDOPA AND LEVODOPA, CARBIDOPA
CARBOPLATIN, CARBOPLATIN
CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
CISPLATIN, CISPLATIN
CLOFARABINE, CLOFARABINE
CLONAZEPAM, CLONAZEPAM
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
CLOZAPINE, CLOZAPINE
DALFAMPRIDINE, DALFAMPRIDINE
DAPTOMYCIN, DAPTOMYCIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACCORD HEALTHCARE INC
 DECITABINE, DECITABINE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DOCETAXEL, DOCETAXEL
 DOFETILIDE, DOFETILIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 ENTECAVIR, ENTECAVIR
 EPLERENONE, EPLERENONE
 EPTIFIBATIDE, EPTIFIBATIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ETOPOSIDE, ETOPOSIDE
 EZETIMIBE, EZETIMIBE
 FINASTERIDE, FINASTERIDE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FLUOROURACIL, FLUOROURACIL
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FULVESTRANT, FULVESTRANT
 FUROSEMIDE, FUROSEMIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLIPIZIDE, GLIPIZIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 ITRACONAZOLE, ITRACONAZOLE
 LETROZOLE, LETROZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LISINAPRIL, LISINAPRIL
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLDOPA, METHYLDOPA
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MITOMYCIN, MITOMYCIN
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 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
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 SPIRONOLACTONE, SPIRONOLACTONE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACCORD HEALTHCARE INC
 TACROLIMUS, TACROLIMUS
 TADALAFIL, TADALAFIL
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TEMSIROLIMUS, TEMSIROLIMUS
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TOPIRAMATE, TOPIRAMATE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VIGABATRIN, VIGABATRIN
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

ACCORD HLTHCARE INC

* ACCORD HEALTHCARE INC USA
 BUSULFAN, BUSULFAN
 TIGECYCLINE, TIGECYCLINE

ACELLA

* ACELLA PHARMACEUTICALS LLC
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CICLOPIROX, CICLOPIROX
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DUTASTERIDE, DUTASTERIDE
 NIFEDIPINE, NIFEDIPINE
 PHENYTOIN SODIUM, PHENYTOIN SODIUM

ACELLA PHARMS LLC

* ACELLA PHARMACEUTICALS LLC
 GABAPENTIN, GABAPENTIN

ACELRX PHARMS

* ACELRX PHARMACEUTICALS INC
 DSUVIA, SUFENTANIL CITRATE

ACERUS

* ACERUS PHARMACEUTICALS CORP
 NATESTO, TESTOSTERONE

ACI

* ACI HEALTHCARE LTD
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 LEVETIRACETAM, LEVETIRACETAM
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

ACIC PHARMS

* ACIC PHARMACEUTICALS INC
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 TRANEXAMIC ACID, TRANEXAMIC ACID

ACORDA

* ACORDA THERAPEUTICS INC
 AMPYRA, DALFAMPRIDINE
 INBRIJA, LEVODOPA

ACROTECH

* ACROTECH BIOPHARMA LLC
 BELEODAQ, BELINOSTAT
 EVOMELA, MELPHALAN HYDROCHLORIDE
 FOLOTYN, PRALATREXATE
 FUSILEV, LEVOLEUCOVORIN CALCIUM
 KHAPZORY, LEVOLEUCOVORIN

ACRUX DDS

* ACRUX DDS PTY LTD
 EFINACONAZOLE, EFINACONAZOLE

ACRUX DDS PTY

* ACRUX DDS PTY LTD
 LIDOCAINE AND PRILOCAINE, LIDOCAINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******ACS DOBFAR**

* ACS DOBFAR SPA
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
 CEFOXITIN, CEFOXITIN SODIUM
 CEFTAZIDIME, CEFTAZIDIME
 CEFTRIAXONE, CEFTRIAXONE 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
 ALBENDAZOLE, ALBENDAZOLE
 ALPRAZOLAM, ALPRAZOLAM
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 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
 DOXEPIN HYDROCHLORIDE, DOXEPIN 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
 LOVASTATIN, LOVASTATIN
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NIFEDIPINE, NIFEDIPINE
 OXAZEPAM, OXAZEPAM
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PREGABALIN, PREGABALIN
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 RANOLAZINE, RANOLAZINE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 TEMAZEPAM, TEMAZEPAM
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

* ACTAVIS ELIZABETH LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 ALPRAZOLAM, ALPRAZOLAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

ACTAVIS INC

* ACTAVIS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACTAVIS INC
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE

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)
 ISOTRETINOIN, ISOTRETINOIN
 MESALAMINE, MESALAMINE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

ACTAVIS LABS FL INC

* ACTAVIS LABORATORIES FL INC
 BUDESONIDE, BUDESONIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 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
 ENZALUTAMIDE, ENZALUTAMIDE
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 LEVETIRACETAM, LEVETIRACETAM
 METAXALONE, METAXALONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 OMEPRAZOLE, OMEPRAZOLE
 PALIPERIDONE, PALIPERIDONE
 PAROXETINE MESYLATE, PAROXETINE MESYLATE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 RAMELTEON, RAMELTEON
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TAZTIA XT, DILTIAZEM HYDROCHLORIDE
 TETRABENAZINE, TETRABENAZINE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

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
 FIORICET W/ CODEINE, ACETAMINOPHEN
 LIDOCAINE, LIDOCAINE
 PROGESTERONE, PROGESTERONE
 SCOPOLAMINE, SCOPOLAMINE
 TESTOSTERONE, TESTOSTERONE

* ACTAVIS LABORATORIES UT INC INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 PIMECROLIMUS, PIMECROLIMUS
 TESTOSTERONE, TESTOSTERONE

ACTAVIS LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * ACTAVIS LLC
 - AZACITIDINE, AZACITIDINE
 - DAPSONE, DAPSONE
 - FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 - MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
- * ACTAVIS LLC AN INDIRECT WHOLLY-OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - DOCETAXEL, DOCETAXEL
 - HYDROXOCOBALAMIN, HYDROXOCOBALAMIN
 - OXALIPLATIN, OXALIPLATIN

ACTAVIS MID ATLANTIC

- * ACTAVIS MID ATLANTIC LLC
 - ACYCLOVIR, ACYCLOVIR
 - ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 - ADAPALENE, ADAPALENE
 - BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 - CICLOPIROX, CICLOPIROX
 - CLINDAMYCIN PHOSPHATE AND TRETINOIN, CLINDAMYCIN PHOSPHATE
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - DESOXIMETASONE, DESOXIMETASONE
 - DICLOFENAC SODIUM, DICLOFENAC SODIUM
 - GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 - HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 - HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 - HYDROCORTISONE, HYDROCORTISONE
 - LEVETIRACETAM, LEVETIRACETAM
 - MESALAMINE, MESALAMINE
 - NITROFURANTOIN, NITROFURANTOIN
 - NYSTATIN, NYSTATIN
 - PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 - VALNAC, BETAMETHASONE VALERATE
- * ACTAVIS MID ATLANTIC LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - IBUPROFEN, IBUPROFEN
 - PERMETHRIN, PERMETHRIN (OTC)

ACTAVIS PHARMA

- * ACTAVIS PHARMA INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - MICONAZOLE NITRATE, MICONAZOLE NITRATE

ACTAVIS TOTOWA

- * ACTAVIS TOTOWA LLC
 - DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 - EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 - FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 - OXALIPLATIN, OXALIPLATIN
 - PACLITAXEL, PACLITAXEL
 - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 - VINORELBINE TARTRATE, VINORELBINE TARTRATE

ACTELION

- * ACTELION PHARMACEUTICALS US INC
 - 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
 - OSMOLEX ER, AMANTADINE HYDROCHLORIDE

ADAMIS PHARMS CORP

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ADAMIS PHARMACEUTICALS CORP
 SYMJEPI, EPINEPHRINE
 ZIMHI, NALOXONE HYDROCHLORIDE

ADAPTIS

* ADAPTIS PHARMA PRIVATE LTD
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM

ADARE PHARMS INC

* ADARE PHARMACEUTICALS INC
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

ADDMEDICA SAS

* ADDMEDICA SAS
 SIKLOS, HYDROXYUREA

ADHERA

* ADHERA THERAPEUTICS INC
 PRESTALIA, AMLODIPINE BESYLATE

ADIENNE SA

* ADIENNE SA
 TEPADINA, THIOTEPA

AERIE PHARMS INC

* AERIE PHARMACEUTICALS INC
 RHOPRESSA, NETARSUDIL MESYLATE
 ROCKLATAN, LATANOPROST

AET PHARMA

* AET PHARMA US INC
 POSACONAZOLE, POSACONAZOLE

AGILE

* AGILE THERAPEUTICS INC
 TWIRLA, ETHINYL ESTRADIOL

AGNITIO

* AGNITIO INC
 ETHACRYNIC ACID, ETHACRYNIC ACID
 TRIAMTERENE, TRIAMTERENE

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

AJANTA PHARMA LTD

* AJANTA PHARMA LTD
 ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 CAPTOPRIL, CAPTOPRIL
 CHLORTHALIDONE, CHLORTHALIDONE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOXEPIIN HYDROCHLORIDE, DOXEPIIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DROXIDOPA, DROXIDOPA
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 ENTACAPONE, ENTACAPONE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AJANTA PHARMA LTD**

LAMOTRIGINE, LAMOTRIGINE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 RANOLAZINE, RANOLAZINE
 RISPERIDONE, RISPERIDONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 TADALAFIL, TADALAFIL
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZOLMITRIPTAN, ZOLMITRIPTAN

AKARX INC*** AKARX INC**

DOPTELET, AVATROMBOPAG MALEATE

AKCEA THERAPS*** AKCEA THERAPEUTICS INC**

TEGSEDI, INOTERSEN SODIUM

AKORN*** AKORN OPERATING CO LLC**

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ACYCLOVIR, ACYCLOVIR
 ADENOSINE, ADENOSINE
 AK-FLUOR 10%, FLUORESCEIN SODIUM
 AK-FLUOR 25%, FLUORESCEIN SODIUM
 AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
 AKTEN, LIDOCAINE HYDROCHLORIDE
 AKTOB, TOBRAMYCIN
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALFENTA, ALFENTANIL HYDROCHLORIDE
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMICAR, AMINOCAPROIC ACID
 APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE
 ATROPINE SULFATE, ATROPINE SULFATE
 AZASITE, AZITHROMYCIN
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BAL, DIMERCAPROL
 BALANCED SALT, CALCIUM CHLORIDE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 BETIMOL, TIMOLOL
 BIMATOPROST, BIMATOPROST
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BROMFENAC SODIUM, BROMFENAC SODIUM
 CALCIPOTRIENE, CALCIPOTRIENE
 CALCITRIOL, CALCITRIOL
 CARBOPLATIN, CARBOPLATIN
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CICLOPIROX, CICLOPIROX
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 COGENTIN, BENZTROPINE MESYLATE
 CORMAX, CLOBETASOL PROPIONATE
 COSOPT PF, DORZOLAMIDE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AKORN OPERATING CO LLC
 COSOPT, DORZOLAMIDE HYDROCHLORIDE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 CYCLOPENTOLATE HYDROCHLORIDE, CYCLOPENTOLATE HYDROCHLORIDE
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 DESONIDE, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIURIL, CHLOROTHIAZIDE SODIUM
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DRONABINOL, DRONABINOL
 EMBELINE E, CLOBETASOL PROPIONATE
 EMBELINE, CLOBETASOL PROPIONATE
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 EPTIFIBATIDE, EPTIFIBATIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 FAMOTIDINE, FAMOTIDINE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
 GABAPENTIN, GABAPENTIN
 GATIFLOXACIN, GATIFLOXACIN
 GENTAK, GENTAMICIN SULFATE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 IBUPROFEN, IBUPROFEN
 IC-GREEN, INDOCYANINE GREEN
 INAPSINE, DROPERIDOL
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LACTULOSE, LACTULOSE
 LATANOPROST, LATANOPROST
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCARNITINE, LEVOCARNITINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORAZEPAM, LORAZEPAM
 LOTE Prednol Etabonate, LOTE Prednol Etabonate
 MEGESTROL ACETATE, MEGESTROL ACETATE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 MORPHINE SULFATE, MORPHINE SULFATE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AKORN OPERATING CO LLC
 NEDOCROMIL SODIUM, NEDOCROMIL SODIUM
 NEMBUTAL SODIUM, PENTOBARBITAL 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
 NYSTATIN, NYSTATIN
 OFLOXACIN, OFLOXACIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PAREMYD, HYDROXYAMPHETAMINE HYDROBROMIDE
 PARICALCITOL, PARICALCITOL
 PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PREDNISOLONE, PREDNISOLONE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
 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
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIMOLOL, TIMOLOL
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TOBRAMYCIN, TOBRAMYCIN
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIFLURIDINE, TRIFLURIDINE
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TROPICACYL, TROPICAMIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 VOSOL HC, ACETIC ACID, GLACIAL
 VOSOL, ACETIC ACID, GLACIAL
 XOPENEX, LEVALBUTEROL HYDROCHLORIDE
 ZIOPTAN, TAFLUPROST
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

ALBIREO

* ALBIREO AB
 BYLVAY, ODEVIXIBAT

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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ALCON LABORATORIES INC**

CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE
 CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE
 FLUORESCITE, FLUORESCEIN SODIUM
 ISOPTO ATROPINE, ATROPINE SULFATE
 MYDRIACYL, TROPICAMIDE
 PATADAY ONCE DAILY RELIEF, OLOPATADINE HYDROCHLORIDE (OTC)
 PATADAY TWICE DAILY RELIEF, OLOPATADINE HYDROCHLORIDE (OTC)
 SIMBRINZA, BRIMONIDINE TARTRATE

ALCON PHARMS LTD

*** ALCON PHARMACEUTICALS LTD**
 BETADINE, POVIDONE-IODINE
 ZADITOR, KETOTIFEN FUMARATE (OTC)

ALEMBIC GLOBAL

*** ALEMBIC GLOBAL HOLDING SA**
 TREPROSTINIL, TREPROSTINIL

ALEMBIC LABS

*** ALEMBIC LABS LLC**
 BENZONATATE, BENZONATATE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 ERGOCALCIFEROL, ERGOCALCIFEROL
 FENOFIBRATE, FENOFIBRATE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 LEVETIRACETAM, LEVETIRACETAM
 METRONIDAZOLE, METRONIDAZOLE

ALEMBIC LTD

*** ALEMBIC LTD**
 LITHIUM CARBONATE, LITHIUM CARBONATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE

ALEMBIC PHARMS LTD

*** ALEMBIC PHARMACEUTICALS LTD**
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ACYCLOVIR, ACYCLOVIR
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ASENAPINE MALEATE, ASENAPINE MALEATE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BIMATOPROST, BIMATOPROST
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CELECOXIB, CELECOXIB
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DEFERASIROX, DEFERASIROX
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DESVENLAFAXINE, DESVENLAFAXINE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE HCLATE, DOXYCYCLINE HCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTACAPONE, ENTACAPONE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALEMBIC PHARMACEUTICALS LTD
 FAMOTIDINE, FAMOTIDINE
 FEBUXOSTAT, FEBUXOSTAT
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 ITRACONAZOLE, ITRACONAZOLE
 LAMOTRIGINE, LAMOTRIGINE
 LEFLUNOMIDE, LEFLUNOMIDE
 LINEZOLID, LINEZOLID
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MEPROBAMATE, MEPROBAMATE
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE, METRONIDAZOLE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MODAFINIL, MODAFINIL
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PREGABALIN, PREGABALIN
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMAZEPAM, TEMAZEPAM
 TERIFLUNOMIDE, TERIFLUNOMIDE
 THEOPHYLLINE, THEOPHYLLINE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOBRAMYCIN, TOBRAMYCIN
 TRAVOPROST, TRAVOPROST
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN

ALEOR DERMACEUTICALS

* ALEOR DERMACEUTICALS LTD
 ADAPALENE, ADAPALENE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DESONIDE, DESONIDE
 LIDOCAINE, LIDOCAINE
 METRONIDAZOLE, METRONIDAZOLE
 MUPIROCIN, MUPIROCIN CALCIUM
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 TAVABOROLE, TAVABOROLE

ALEXZA PHARMS

* ALEXZA PHARMACEUTICALS INC
 ADASUVE, LOXAPINE

ALFASIGMA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALFASIGMA USA INC
ZELNORM, TEGASEROD MALEATE

ALIGNSCIENCE PHARMA

* ALIGNSCIENCE PHARMA INC
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE

ALIMERA SCIENCES INC

* ALIMERA SCIENCES INC
ILUVIEN, FLUOCINOLONE ACETONIDE

ALK ABELLO

* ALK-ABELLO INC
OTIPRIO, CIPROFLOXACIN

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
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
APREMILAST, APREMILAST
ARIPIPRAZOLE, ARIPIPRAZOLE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
AZATHIOPRINE, AZATHIOPRINE
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
CAPECITABINE, CAPECITABINE
CEFDINIR, CEFdinIR
CEFIXIME, CEFIXIME
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEPHALEXIN, CEPHALEXIN
CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
CHLORTHALIDONE, CHLORTHALIDONE
CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
CHOLESTYRAMINE, CHOLESTYRAMINE
CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
CLOBAZAM, CLOBAZAM
COLCHICINE, COLCHICINE
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
DALFAMPRIDINE, DALFAMPRIDINE
DEFERASIROX, DEFERASIROX
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
DIMETHYL FUMARATE, DIMETHYL FUMARATE
DROXIDOPA, DROXIDOPA
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
EVEROLIMUS, EVEROLIMUS
EZETIMIBE AND SIMVASTATIN, EZETIMIBE
EZETIMIBE, EZETIMIBE
FEBUXOSTAT, FEBUXOSTAT
FINASTERIDE, FINASTERIDE
FOSFOMYCIN TROMETHAMINE, FOSFOMYCIN TROMETHAMINE
GABAPENTIN, GABAPENTIN
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
IBUPROFEN AND FAMOTIDINE, FAMOTIDINE
IBUPROFEN, IBUPROFEN
ITRACONAZOLE, ITRACONAZOLE
LAMOTRIGINE, LAMOTRIGINE
LANSOPRAZOLE, LANSOPRAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALKEM LABORATORIES LTD
 LIDOCAINE, LIDOCAINE
 LINEZOLID, LINEZOLID
 MARINOL, DRONABINOL
 MESALAMINE, MESALAMINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RANOLAZINE, RANOLAZINE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 RILUZOLE, RILUZOLE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 RUFINAMIDE, RUFINAMIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SIROLIMUS, SIROLIMUS
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TACROLIMUS, TACROLIMUS
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOBRAMYCIN, TOBRAMYCIN
 TOLVAPTAN, TOLVAPTAN
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VALSARTAN, VALSARTAN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VIGABATRIN, VIGABATRIN

ALKERMES

* ALKERMES INC
 VIVITROL, NALTREXONE

ALKERMES INC

* ALKERMES INC
 ARISTADA INITIO KIT, ARIPIPRAZOLE LAUROXIL
 ARISTADA, ARIPIPRAZOLE LAUROXIL
 LYBALVI, OLANZAPINE

ALLEGIANCE HLTHCARE

* ALLEGIANCE HEALTHCARE CORP
 POVIDONE IODINE, POVIDONE-IODINE (OTC)

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
 ALOCRIL, NEDOCROMIL SODIUM
 ALPHAGAN P, BRIMONIDINE TARTRATE
 AVAGE, TAZAROTENE
 COMBIGAN, BRIMONIDINE TARTRATE
 ELESTAT, EPINASTINE HYDROCHLORIDE
 LASTACFT, ALCAFTADINE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALLERGAN INC
 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
 PRED FORTE, PREDNISOLONE ACETATE
 PRED MILD, PREDNISOLONE ACETATE
 PRED-G, GENTAMICIN SULFATE

* ALLERGAN SALES LLC
 ACTIGALL, URSODIOL
 ANDRODERM, TESTOSTERONE
 AVYCAZ, AVIBACTAM SODIUM
 BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE
 BENTYL, DICYCLOMINE HYDROCHLORIDE
 BYSTOLIC, 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, FERRIC OXYHYDROXIDE
 LEXAPRO, ESCITALOPRAM OXALATE
 LINZESS, LINACLOTIDE
 NAMENDA XR, MEMANTINE HYDROCHLORIDE
 NAMENDA, MEMANTINE HYDROCHLORIDE
 NAMZARIC, DONEPEZIL HYDROCHLORIDE
 OXYTROL FOR WOMEN, OXYBUTYNIN (OTC)
 OXYTROL, OXYBUTYNIN
 PYLERA, BISMUTH SUBCITRATE POTASSIUM
 RAPAFLO, SILODOSIN
 RECTIV, NITROGLYCERIN
 SAPHRIS, ASENAPINE MALEATE
 SAVELLA, MILNACIPRAN HYDROCHLORIDE
 TEFLARO, CEFTAROLINE FOSAMIL
 UBRELVY, UBROGEPANT
 URSO 250, URSODIOL
 URSO FORTE, URSODIOL
 VIIBRYD, VILAZODONE HYDROCHLORIDE
 VRAYLAR, CARIPRAZINE HYDROCHLORIDE

ALLERGAN HOLDINGS

* ALLERGAN HOLDINGS UNLTD CO
 VIBERZI, ELUXADOLINE

ALLERGAN INC

* ALLERGAN INC
 DURYSTA, BIMATOPROST

ALMAJECT

* ALMAJECT INC
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALMAJECT INC
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE

ALMATICA

* ALMATICA PHARMA INC
 GRALISE, GABAPENTIN

* ALMATICA PHARMA LLC
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FORFIVO XL, BUPROPION HYDROCHLORIDE
 LOREEV XR, LORAZEPAM
 MACROBID, NITROFURANTOIN
 MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE
 NAPRELAN, NAPROXEN SODIUM
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 TENORETIC 100, ATENOLOL
 TENORETIC 50, ATENOLOL
 TENORMIN, ATENOLOL
 ZESTORETIC, HYDROCHLOROTHIAZIDE
 ZESTRIL, LISINOPRIL

ALMIRALL

* ALMIRALL LLC
 ACTICLATE, DOXYCYCLINE HYCLATE
 ACZONE, DAPSONE
 ALTABAX, RETAPAMULIN
 AZELEX, AZELAIC ACID
 CORDRAN SP, FLURANDRENOLIDE
 CORDRAN, FLURANDRENOLIDE
 KLISYRI, TIRBANIBULIN
 SEYSARA, SARECYCLINE HYDROCHLORIDE
 VELTIN, CLINDAMYCIN PHOSPHATE
 VERDESO, DESONIDE
 XOLEGEL, KETOCONAZOLE

ALNYLAM PHARMS INC

* ALNYLAM PHARMACEUTICALS INC
 GIVLAARI, GIVOSIRAN SODIUM
 ONPATTRO, PATISIRAN SODIUM
 OXLUMO, LUMASIRAN SODIUM

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

ALTHERA PHARMS

* ALTHERA PHARMACEUTICALS LLC
 ROSZET, EZETIMIBE

ALVOGEN

* ALVOGEN INC
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 ATENOLOL, ATENOLOL
 BONSIITY, TERIPARATIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CARBIDOPA, CARBIDOPA
 CICLOPIROX, CICLOPIROX
 DAPSONE, DAPSONE
 DEXAMETHASONE, DEXAMETHASONE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ALVOGEN INC**

DISULFIRAM, DISULFIRAM
 DYNACIN, MINOCYCLINE HYDROCHLORIDE
 ESTRADIOL, ESTRADIOL
 ETHACRYNIC ACID, ETHACRYNIC ACID
 EXEMESTANE, EXEMESTANE
 FELBAMATE, FELBAMATE
 FLUOCINONIDE, FLUOCINONIDE
 HYDROCODONE BITARTRATE, HYDROCODONE BITARTRATE
 LIDEX, FLUOCINONIDE
 LIDEX-E, FLUOCINONIDE
 MELPHALAN, MELPHALAN
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 NEVIRAPINE, NEVIRAPINE
 NITROGLYCERIN, NITROGLYCERIN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 PYRIMETHAMINE, PYRIMETHAMINE
 RIVASTIGMINE, RIVASTIGMINE
 SPECTAZOLE, ECONAZOLE NITRATE
 THYRO-TABS, LEVOTHYROXINE SODIUM **
 TRANDATE, LABETALOL HYDROCHLORIDE
 UREX, METHENAMINE HIPPURATE

ALVOGEN PINE BROOK*** ALVOGEN PINE BROOK LLC**

HYDROCODONE BITARTRATE, HYDROCODONE BITARTRATE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE

AM REGENT*** AMERICAN REGENT INC**

ACETYLCYSTEINE, ACETYLCYSTEINE
 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
 CYANOCOBALAMIN, CYANOCOBALAMIN
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DROPERIDOL, DROPERIDOL
 ESTRADIOL VALERATE, ESTRADIOL VALERATE
 FOMEPIZOLE, FOMEPIZOLE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 INJECTAFER, FERRIC CARBOXYMALTOSIDE
 LEVOCARNITINE, LEVOCARNITINE
 METHOCARBAMOL, METHOCARBAMOL
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 MULTRYS, CUPRIC SULFATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NITROGLYCERIN, NITROGLYCERIN
 OLANZAPINE, OLANZAPINE
 SELENIOS ACID, SELENIOS ACID
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TRALEMENT, CUPRIC SULFATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VASOPRESSIN, VASOPRESSIN
 VENOFER, FERRIC OXYHYDROXIDE
 ZINC SULFATE, ZINC SULFATE

AMARIN PHARMS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMARIN PHARMACEUTICALS IRELAND LTD
VASCEPA, ICOSAPENT ETHYL

AMGEN

* AMGEN INC
SENSIPAR, CINACALCET HYDROCHLORIDE

AMGEN INC

* AMGEN INC
CORLANOR, IVABRADINE
CORLANOR, IVABRADINE HYDROCHLORIDE
LUMAKRAS, SOTORASIB
OTEZLA, APREMILAST

AMICI

* AMICI PHARMACEUTICALS LLC
CATAFLAM, DICLOFENAC POTASSIUM
METRONIDAZOLE, METRONIDAZOLE
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE

AMICUS THERAP US

* AMICUS THERAPEUTICS US LLC
GALAFOLD, MIGALASTAT HYDROCHLORIDE

AMIVAS

* AMIVAS LLC
ARTESUNATE, ARTESUNATE

AMNEAL

* AMNEAL EU LTD
AMINOCAPROIC ACID, AMINOCAPROIC ACID
APREMILAST, APREMILAST
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
BUSULFAN, BUSULFAN
CARMUSTINE, CARMUSTINE
CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
CLOFARABINE, CLOFARABINE
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
DEFERASIROX, DEFERASIROX
DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DEXAMETHASONE, DEXAMETHASONE
DIFLUPREDNATE, DIFLUPREDNATE
DIMETHYL FUMARATE, DIMETHYL FUMARATE
DOCETAXEL, DOCETAXEL
DOXERCALCIFEROL, DOXERCALCIFEROL
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
EPHEDRINE SULFATE, EPHEDRINE SULFATE
ETRAVIRINE, ETRAVIRINE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
FULVESTRANT, FULVESTRANT
GLYCOPYRROLATE, GLYCOPYRROLATE
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METYROSINE, METYROSINE
NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
OFLOXACIN, OFLOXACIN
PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
PIRFENIDONE, PIRFENIDONE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PREDNISONE, PREDNISONE
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMNEAL EU LTD
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

* AMNEAL PHARMACEUTICALS LLC
 ACTIVELLA, ESTRADIOL
 ACYCLOVIR, ACYCLOVIR
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 AZATHIOPRINE, AZATHIOPRINE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BUPRENORPHINE, BUPRENORPHINE
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 ELURYNG, ETHINYL ESTRADIOL
 ESTRADIOL, ESTRADIOL
 ETHINYL ESTRADIOL AND NORELGESTROMIN, ETHINYL ESTRADIOL
 FENOFIBRATE, FENOFIBRATE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 LIDOCAINE, LIDOCAINE
 LUBIPROSTONE, LUBIPROSTONE
 MESALAMINE, MESALAMINE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NAPROXEN, NAPROXEN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RISPERIDONE, RISPERIDONE
 RITONAVIR, RITONAVIR
 SIROLIMUS, SIROLIMUS
 SUCRALFATE, SUCRALFATE
 TAVABOROLE, TAVABOROLE
 TESTOSTERONE, TESTOSTERONE
 TIGECYCLINE, TIGECYCLINE

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
 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
 COLCHICINE, COLCHICINE
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
 DIVALPROEX SODIUM, DIVALPROEX SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMNEAL PHARMACEUTICALS
 ENTECAVIR, ENTECAVIR
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESTRADIOL, ESTRADIOL
 FELBAMATE, FELBAMATE
 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 DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN
 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
 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
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 DUTASTERIDE, DUTASTERIDE
 ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN, GUAIFENESIN (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 IRBESARTAN, IRBESARTAN
 LAMOTRIGINE, LAMOTRIGINE
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMNEAL PHARMACEUTICALS OF NEW YORK LLC
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 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
 PALIPERIDONE, PALIPERIDONE
 PARICALCITOL, PARICALCITOL
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 RIVASTIGMINE, RIVASTIGMINE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 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
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLOROTHALIDONE, CHLOROTHALIDONE
 CLOBAZAM, CLOBAZAM
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ERYTHROMYCIN, ERYTHROMYCIN
 ETHACRYNIC ACID, ETHACRYNIC ACID
 ETODOLAC, ETODOLAC
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 NADOLOL, NADOLOL
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 OXAPROZIN, OXAPROZIN
 PARICALCITOL, PARICALCITOL
 PHYTONADIONE, PHYTONADIONE
 PREGABALIN, PREGABALIN
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SILODOSIN, SILODOSIN
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 URSODIOL, URSODIOL

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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMNEAL PHARMACEUTICALS NY LLC
 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
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

* AMNEAL PHARMACEUTICALS OF NY LLC
 BEXAROTENE, BEXAROTENE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 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
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM

AMPHASTAR PHARMS INC

* AMPHASTAR PHARMACEUTICALS INC
 CORTROSYN, COSYNTROPIN
 ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
 GLUCAGON, GLUCAGON
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE

AMRING PHARMS

* AMRING PHARMACEUTICALS INC
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 LATANOPROST, LATANOPROST
 LYSTEDA, TRANEXAMIC ACID
 MESALAMINE, MESALAMINE
 NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TIMOLOL MALEATE, TIMOLOL MALEATE

AMRYT

* AMRYT PHARMACEUTICALS DAC
 JUXTAPID, LOMITAPIDE MESYLATE

AMTA

* AMTA LABS LTD
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE

ANACOR PHARMS INC

* ANACOR PHARMACEUTICALS INC
 EUCRISA, CRISABOROLE
 KERYDIN, TAVABOROLE

ANBEX

* ANBEX INC
 IOSAT, POTASSIUM IODIDE (OTC)

ANBISON LAB

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ANBISON LABORATORY CO LTD
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE

ANCHEN PHARMS

* ANCHEN PHARMACEUTICALS INC
 ALISKIREN HEMIFUMARATE, ALISKIREN HEMIFUMARATE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 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
 TRETINOIN, TRETINOIN
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

* ANCHEN PHARMACEUTICALS, INC
 ALPRAZOLAM, ALPRAZOLAM
 CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN

ANDA REPOSITORY

* ANDA REPOSITORY LLC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 FLAC, FLUOCINOLONE ACETONIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 ISONIAZID, ISONIAZID
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LEVETIRACETAM, LEVETIRACETAM
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 PRIMIDONE, PRIMIDONE

ANDOR PHARMS

* ANDOR PHARMACEUTICALS LLC
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

ANDRX LABS LLC

* ANDRX LABS LLC
 FORTAMET, METFORMIN HYDROCHLORIDE

ANI PHARMS

* ANI PHARMACEUTICALS INC
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 ARIMIDEX, ANASTROZOLE
 ATACAND HCT, CANDESARTAN CILEXETIL
 ATACAND, CANDESARTAN CILEXETIL
 BEXAROTENE, BEXAROTENE
 BRETHINE, TERBUTALINE SULFATE
 CARBIDOPA, CARBIDOPA
 CASODEX, BICALUTAMIDE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
 CORTENEMA, HYDROCORTISONE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 ETODOLAC, ETODOLAC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ANI PHARMACEUTICALS INC
 FELBAMATE, FELBAMATE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUCONAZOLE, FLUCONAZOLE
 GLIPIZIDE, GLIPIZIDE
 INDAPAMIDE, INDAPAMIDE
 INDERAL LA, PROPRANOLOL HYDROCHLORIDE
 INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
 KIONEX, SODIUM POLYSTYRENE SULFONATE
 LITHOBID, LITHIUM CARBONATE
 LUVOX, FLUVOXAMINE MALEATE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MIGLUSTAT, MIGLUSTAT
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NILUTAMIDE, NILUTAMIDE
 OXISTAT, OXICONAZOLE NITRATE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PANDEL, HYDROCORTISONE PROBUTATE
 PENICILLAMINE, PENICILLAMINE
 PINDOLOL, PINDOLOL
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PURIFIED CORTROPHIN GEL, CORTICOTROPIN
 REGLAN, METOCLOPRAMIDE HYDROCHLORIDE
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VANCOCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VEREGEN, SINECATECHINS

ANIMA

* ANIMA PHARMACEUTICALS PVT LTD
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 DEXAMETHASONE, DEXAMETHASONE

ANNORA

* ANNORA PHARMA PRIVATE LTD
 DROXIDOPA, DROXIDOPA
 LAMIVUDINE, LAMIVUDINE

ANNORA PHARMA

* ANNORA PHARMA PRIVATE LTD
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 CAPTOPRIL, CAPTOPRIL
 DEFERASIROX, DEFERASIROX
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 ITRACONAZOLE, ITRACONAZOLE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 NABUMETONE, NABUMETONE
 OXCARBAZEPINE, OXCARBAZEPINE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VIGABATRIN, VIGABATRIN
 ZAFIRLUKAST, ZAFIRLUKAST

ANTARES PHARMA INC

* ANTARES PHARMA INC
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 XYOSTED (AUTOINJECTOR), TESTOSTERONE ENANTHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******ANTIBIOTICE**

* ANTIBIOTICE SA
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 NAFCILLIN SODIUM, NAFCILLIN SODIUM

ANXIN

* ANXIN PHARMA INC
 SEVELAMER CARBONATE, SEVELAMER CARBONATE

APELLIS PHARMS

* APELLIS PHARMACEUTICALS INC
 EMPAVELI, PEGCETACOPLAN

APGDI

* ASTELLAS PHARMA GLOBAL DEVELOPMENT INC
 MYRBETRIQ GRANULES, MIRABEGRON
 MYRBETRIQ, MIRABEGRON

APIL

* ALLERGAN PHARMACEUTICALS INTERNATIONAL LTD
 ACTONEL, RISEDRONATE SODIUM
 ASACOL HD, MESALAMINE
 ATELVIA, RISEDRONATE SODIUM
 DELZICOL, MESALAMINE
 LO LOESTRIN FE, ETHINYL ESTRADIOL
 MINASTRIN 24 FE, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 TAYTULLA, ETHINYL ESTRADIOL

APNAR PHARMA LP

* APNAR PHARMA LP
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

APOTEX

* APOTEX INC
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACYCLOVIR, ACYCLOVIR
 ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATOVAQUONE, ATOVAQUONE
 AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE, AZELASTINE HYDROCHLORIDE
 BEPOTASTINE BESILATE, BEPOTASTINE BESILATE
 BUSULFAN, BUSULFAN
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CEFTAROLINE FOSAMIL, CEFTAROLINE FOSAMIL
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CIMETIDINE, CIMETIDINE (OTC)
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 CYCLOSPORINE, CYCLOSPORINE
 DASATINIB, DASATINIB
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXYCYCLINE HCLATE, DOXYCYCLINE HCLATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ETODOLAC, ETODOLAC
 EZETIMIBE, EZETIMIBE
 FAMCICLOVIR, FAMCICLOVIR
 FAMOTIDINE, FAMOTIDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* APOTEX INC**

FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FLUNISOLIDE, FLUNISOLIDE
 FLUTICASON PROPIONATE, FLUTICASON PROPIONATE (OTC)
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FULVESTRANT, FULVESTRANT
 GEMFIBROZIL, GEMFIBROZIL
 GLIPIZIDE, GLIPIZIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 ICOSAPENT ETHYL, ICOSAPENT ETHYL
 IMATINIB MESYLATE, IMATINIB MESYLATE
 LAMIVUDINE, LAMIVUDINE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OMEPRAZOLE, OMEPRAZOLE
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PENICILLAMINE, PENICILLAMINE
 PENTOXIFYLLINE, PENTOXIFYLLINE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PREGABALIN, PREGABALIN
 RAMIPRIL, RAMIPRIL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SIROLIMUS, SIROLIMUS
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TETRABENAZINE, TETRABENAZINE
 TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
 TIGECYCLINE, TIGECYCLINE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOLVAPTAN, TOLVAPTAN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRAVOPROST, TRAVOPROST
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

APOTEX CORP*** APOTEX CORP**

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

APOTEX INC*** APOTEX INC**

ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALPRAZOLAM, ALPRAZOLAM
 ARIPIPRAZOLE, ARIPIPRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 ATROPINE SULFATE, ATROPINE SULFATE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * APOTEX INC
 BENZONATATE, BENZONATATE
 BICALUTAMIDE, BICALUTAMIDE
 BIMATOPROST, BIMATOPROST
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CARBAMAZEPINE, CARBAMAZEPINE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CEFPROZIL, CEFPROZIL
 CELECOXIB, CELECOXIB
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IMIQUIMOD, IMIQUIMOD
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN, LEVOFLOXACIN
 MODAFINIL, MODAFINIL
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 OFLOXACIN, OFLOXACIN
 OLANZAPINE, OLANZAPINE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RISPERIDONE, RISPERIDONE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- * APOTEX INC ETOBICOKE SITE
 ACYCLOVIR, ACYCLOVIR
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CILOSTAZOL, CILOSTAZOL
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 ETODOLAC, ETODOLAC
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 GABAPENTIN, GABAPENTIN
 LEFLUNOMIDE, LEFLUNOMIDE
 LORATADINE, LORATADINE (OTC)
 MELOXICAM, MELOXICAM
 MIRTAZAPINE, MIRTAZAPINE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 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

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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* APOTHECON INC DIV BRISTOL MYERS SQUIBB
KENALOG-80, TRIAMCINOLONE ACETONIDE

APP PHARMS

* APP PHARMACEUTICALS LLC
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE

APPCO

* APPCO PHARMA LLC
ACETAZOLAMIDE, ACETAZOLAMIDE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
CHLORTHALIDONE, CHLORTHALIDONE
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
ELIGLUSTAT TARTRATE, ELIGLUSTAT TARTRATE
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LAMIVUDINE, LAMIVUDINE
LORATADINE, LORATADINE (OTC)
METAXALONE, METAXALONE
MODAFINIL, MODAFINIL
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
RAMELTEON, RAMELTEON
SILDENAFIL CITRATE, SILDENAFIL CITRATE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
ZOLMITRIPTAN, ZOLMITRIPTAN

APRECIA PHARMS

* APRECIA PHARMACEUTICALS LLC
SPRITAM, LEVETIRACETAM

APTAPHARMA INC

* APTAPHARMA INC
IBUPROFEN, IBUPROFEN (OTC)
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
SILDENAFIL CITRATE, SILDENAFIL CITRATE

AQUESTIVE

* AQUESTIVE THERAPEUTICS INC
SYMPAZAN, CLOBAZAM

ARBOR PHARMS LLC

* ARBOR PHARMACEUTICALS LLC
BIDIL, HYDRALAZINE HYDROCHLORIDE
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 ODT, AMPHETAMINE SULFATE
EVEKEO, AMPHETAMINE SULFATE
GLIADEL, CARMUSTINE
HORIZANT, GABAPENTIN ENACARBIL
NYMALIZE, NIMODIPINE
SKLICE, IVERMECTIN (OTC)
SOTYLIZE, SOTALOL HYDROCHLORIDE
TRIPTODUR KIT, TRIPTORELIN PAMOATE

AREVA PHARMS

* AREVA PHARMACEUTICALS INC
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
FUROSEMIDE, FUROSEMIDE
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

ARISE

* ARISE PHARMACEUTICALS LLC
IBUPROFEN, IBUPROFEN (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******ARMSTRONG PHARMS**

* ARMSTRONG PHARMACEUTICALS INC
PRIMATENE MIST, EPINEPHRINE (OTC)

ARRAY BIOPHARMA INC

* ARRAY BIOPHARMA INC
BRAFTOVI, ENCORAFENIB
MEKTOVI, BINIMETINIB

ARROW INTL

* ARROW INTERNATIONAL LTD
LENALIDOMIDE, LENALIDOMIDE

ARTHUR GRP

* ARTHUR GROUP LLC
BUSULFAN, BUSULFAN

ASCEND THERAPS US

* ASCEND THERAPEUTICS US LLC
BINOSTO, ALENDRONATE SODIUM
ESTROGEL, ESTRADIOL

ASCENT PHARMS INC

* ASCENT PHARMACEUTICALS INC
BENZONATATE, BENZONATATE
DETROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DRONABINOL, DRONABINOL
DUTASTERIDE, DUTASTERIDE
FAMOTIDINE, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE (OTC)
GABAPENTIN, GABAPENTIN
GEMFIBROZIL, GEMFIBROZIL
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
IBUPROFEN, IBUPROFEN (OTC)
LISINOPRIL, LISINOPRIL
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
POTASSIUM CITRATE, POTASSIUM CITRATE
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
TOPIRAMATE, TOPIRAMATE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

ASPEN

* ASPEN PHARMA USA INC
HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE

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
THIOGUANINE, THIOGUANINE

ASPIRO

* ASPIRO PHARMA LTD
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE

ASSERTIO

* ASSERTIO THERAPEUTICS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ASSERTIO THERAPEUTICS INC
 CAMBIA, DICLOFENAC POTASSIUM
 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 LS, SOLIFENACIN SUCCINATE
 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
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM

ASTRAZENECA

* ASTRAZENECA LP
 PULMICORT FLEXHALER, BUDESONIDE
 SYMBICORT, BUDESONIDE

* ASTRAZENECA PHARMACEUTICALS LP
 BEVESPI AEROSPHERE, FORMOTEROL FUMARATE
 BRILINTA, TICAGRELOR
 DALIRESP, ROFLUMILAST
 DUAKLIR PRESSAIR, ACLIDINIUM BROMIDE
 FASLODEX, FULVESTRANT
 IRESSA, GEFITINIB
 KOSELUGO, SELUMETINIB SULFATE
 LOKELMA, SODIUM ZIRCONIUM CYCLOSILICATE
 LYNPARZA, OLAPARIB
 NEXIUM IV, ESOMEPRAZOLE SODIUM
 NEXIUM, ESOMEPRAZOLE MAGNESIUM
 PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)
 PULMICORT RESPULES, BUDESONIDE
 SEROQUEL, QUETIAPINE FUMARATE
 TAGRISSO, OSIMERTINIB MESYLATE
 TUDORZA PRESSAIR, ACLIDINIUM BROMIDE
 ZOMIG, ZOLMITRIPTAN
 ZOMIG-ZMT, ZOLMITRIPTAN

* ASTRAZENECA UK LTD
 CALQUENCE, ACALABRUTINIB
 SEROQUEL XR, QUETIAPINE FUMARATE

ASTRAZENECA AB

* ASTRAZENECA AB
 BREZTRI AEROSPHERE, BUDESONIDE
 BYDUREON BCISE, EXENATIDE SYNTHETIC
 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)

ATHEM

* ATHEM HOLDINGS LLC
 FOLIC ACID, FOLIC ACID
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******ATHENA**

* ATHENA BIOSCIENCES LLC
 FIBRICOR, FENOFIBRIC ACID
 QDOLO, TRAMADOL HYDROCHLORIDE

ATHENEX

* ATHENEX PHARMACEUTICAL DIV
 PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM

ATHENEX INC

* ATHENEX INC
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
 ENALAPRILAT, ENALAPRILAT
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FUROSEMIDE, FUROSEMIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 VALPROATE SODIUM, VALPROATE SODIUM

ATLANTIDE

* ATLANTIDE PHARMACEUTICALS AG
 ENTACAPONE, ENTACAPONE
 LAMOTRIGINE, LAMOTRIGINE
 LISINOPRIL, LISINOPRIL
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

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
 LODOSYN, CARBIDOPA

AUCTA

* AUCTA PHARMACEUTICALS INC
 DEFERASIROX, DEFERASIROX
 VIGADRONE, VIGABATRIN

AUGUST

* AUGUST PHARMACEUTICALS LLC
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM

AURINIA

* AURINIA PHARMACEUTICALS INC
 LUPKYNIS, VOCLOSPORIN

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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* 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
 FINASTERIDE, FINASTERIDE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, 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
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 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)
 AFIRMELLE, ETHINYL ESTRADIOL
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD
 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
 ARIPIPRAZOLE, ARIPIPRAZOLE
 ARMODAFINIL, ARMODAFINIL
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATHENTIA NEXT, LEVONORGESTREL (OTC)
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 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
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BACLOFEN, BACLOFEN
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARBIDOPA, CARBIDOPA
 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)
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE (OTC)
 CHLORZOXAZONE, CHLORZOXAZONE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CLOZAPINE, CLOZAPINE
 COLCHICINE, COLCHICINE
 CYONANZ, ETHINYL ESTRADIOL
 DALFAMPRIDINE, DALFAMPRIDINE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOFETILDE, DOFETILIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DROXIDOPA, DROXIDOPA
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUTASTERIDE, DUTASTERIDE
 EFAVIRENZ, EFAVIRENZ
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ENTACAPONE, ENTACAPONE
 ENTECAVIR, ENTECAVIR
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 ESZOPICLONE, ESZOPICLONE
 EZETIMIBE, EZETIMIBE
 FAMCICLOVIR, FAMCICLOVIR
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD
 FELODIPINE, FELODIPINE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
 FINASTERIDE, FINASTERIDE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUCONAZOLE, FLUCONAZOLE
 FLUCYTOSINE, FLUCYTOSINE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GEMFIBROZIL, GEMFIBROZIL
 GLIMEPIRIDE, GLIMEPIRIDE
 GLIPIZIDE, GLIPIZIDE
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 GUAIFENESIN, GUAIFENESIN (OTC)
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 ICLEVIA, ETHINYL ESTRADIOL
 INCASSIA, NORETHINDRONE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 KALLIGA, DESOGESTREL
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LEFLUNOMIDE, LEFLUNOMIDE
 LEVETIRACETAM, LEVETIRACETAM
 LO SIMPESSE, ETHINYL ESTRADIOL
 LO-ZUMANDIMINE, DROSPIRENONE
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE, LORATADINE (OTC)
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 METRONIDAZOLE, METRONIDAZOLE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MILLI, 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
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NEVIRAPINE, NEVIRAPINE
 NEXESTA FE, ETHINYL ESTRADIOL
 NIACIN, NIACIN
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 NIFEDIPINE, NIFEDIPINE
 NYLIA 1/35, ETHINYL ESTRADIOL
 NYLIA 7/7/7, ETHINYL ESTRADIOL
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 OMEPRAZOLE, OMEPRAZOLE
 OXACILLIN SODIUM, OXACILLIN SODIUM
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PHENYTOIN SODIUM, PHENYTOIN SODIUM
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 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)
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PREDNISONE, PREDNISONE
 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
 REPAGLINIDE, REPAGLINIDE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RISPERIDONE, RISPERIDONE
 RITONAVIR, RITONAVIR
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SIMLIYA, DESOGESTREL
 SIMPESE, ETHINYL ESTRADIOL
 SPIRONOLACTONE, SPIRONOLACTONE
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 TADALAFIL, TADALAFIL
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRI-LO-MILI, ETHINYL ESTRADIOL
 TRI-MILI, ETHINYL ESTRADIOL
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VIGABATRIN, VIGABATRIN
 VORICONAZOLE, VORICONAZOLE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN
 ZONISAMIDE, ZONISAMIDE
 ZUMANDIMINE, DROSPIRENONE

* AUROBINDO PHARMA LTD INC
 ZIDOVUDINE, ZIDOVUDINE

AUROBINDO PHARMA USA

* AUROBINDO PHARMA USA INC
 BUDESONIDE, BUDESONIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA USA INC

PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
RASAGILINE MESYLATE, RASAGILINE MESYLATE**AUROLIFE PHARMA LLC**

* AUROLIFE PHARMA LLC

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
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
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE**AUSTARPHARMA**

* AUSTARPHARMA LLC

FENOFIBRATE, FENOFIBRATE
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE**AUSTARPHARMA LLC**

* AUSTARPHARMA LLC

METHOCARBAMOL, METHOCARBAMOL

AUXILIUM PHARMS INC

* AUXILIUM PHARMACEUTICALS INC

TESTOPEL, TESTOSTERONE
THEO-24, THEOPHYLLINE**AUXILIUM PHARMS LLC**

* AUXILIUM PHARMACEUTICALS LLC

EDEX, ALPROSTADIL
TESTIM, TESTOSTERONE
THEO-24, THEOPHYLLINE**AVANIR PHARMS**

* 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**AVEMA PHARMA**

* AVEMA PHARMA SOLUTIONS

IBUPROFEN, IBUPROFEN (OTC)

AVENT

* AVENT INC

PYTEST KIT, UREA, C-14
PYTEST, UREA, C-14

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******AVEO PHARMS**

* AVEO PHARMACEUTICALS INC
FOTIVDA, TIVOZANIB HYDROCHLORIDE

AVERITAS

* AVERITAS PHARMA INC
QUTENZA, CAPSAICIN

AVET

* AVET PHARMACEUTICALS INC
ACHROMYCIN V, TETRACYCLINE HYDROCHLORIDE
DOXERCALCIFEROL, DOXERCALCIFEROL

AVET LIFESCIENCES

* AVET LIFESCIENCES LTD
BICNU, CARMUSTINE
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE

AVEVA

* AVEVA DRUG DELIVERY SYSTEMS INC
BUPRENORPHINE, BUPRENORPHINE
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)

AVID RADIOPHARMS INC

* AVID RADIOPHARMACEUTICALS INC
AMYVID, FLORBETAPIR F-18
TAUVID, FLORTAUCIPIR F-18

AVION PHARMS

* AVION PHARMACEUTICALS LLC
BALCOLTRA, ETHINYL ESTRADIOL
DHIVY, CARBIDOPA
PONSTEL, MEFENAMIC ACID

AVONDALE PHARMS

* AVONDALE PHARMACEUTICALS LLC
NIACOR, NIACIN

AXAR PHARMS INC

* AXAR PHARMACEUTICALS INC
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE

AYANA PHARMA LTD

* AYANA PHARMA LTD
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE

AYTU

* AYTU BIOSCIENCE INC
KARBINAL ER, CARBINOXAMINE MALEATE
TUZISTRA XR, CHLORPHENIRAMINE POLISTIREX
ZOLPIMIST, ZOLPIDEM TARTRATE

AZURITY

* AZURITY PHARMACEUTICALS INC
EPANED, ENALAPRIL MALEATE
EPRONTIA, TOPIRAMATE
FIRVANQ KIT, VANCOMYCIN HYDROCHLORIDE
KATERZIA, AMLODIPINE BENZOATE
QBRELIS, LISINAPRIL
XATMEP, METHOTREXATE SODIUM

**** B ******B BRAUN**

* B BRAUN MEDICAL INC
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
AMINO ACIDS, AMINO ACIDS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* B BRAUN MEDICAL INC
 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
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* B BRAUN MEDICAL INC

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,
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* B BRAUN MEDICAL INC
 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
 ACETAMINOPHEN, ACETAMINOPHEN
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 CLOROTEKAL, CHLOROPROCAINE HYDROCHLORIDE
 HEPARIN SODIUM, HEPARIN SODIUM
 LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER, MEROPENEM

BAJAJ

* BAJAJ MEDICAL
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE

* BAJAJ MEDICAL LLC
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

BANNER LIFE SCIENCES

* BANNER LIFE SCIENCES LLC
 BAFIERTAM, MONOMETHYL FUMARATE

BARR

* BARR LABORATORIES INC
 AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
 ARANELLE, ETHINYL ESTRADIOL
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 BALZIVA-28, ETHINYL ESTRADIOL
 CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 DANAZOL, DANAZOL
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 DUTASTERIDE, DUTASTERIDE
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 ESTRADIOL AND NORGESTIMATE, ESTRADIOL
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 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
 LESSINA-28, ETHINYL ESTRADIOL
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 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
 PORTIA-28, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BARR LABORATORIES INC
 - SPRINTEC, ETHINYL ESTRADIOL
 - 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 INC

- * BARR LABORATORIES INC
 - ACITRETIN, ACITRETIN
 - CLOZAPINE, CLOZAPINE
 - ESTRADIOL, ESTRADIOL
 - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - OLANZAPINE, OLANZAPINE
 - OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 - TRETINOIN, TRETINOIN
 - TRI LO SPRINTEC, ETHINYL ESTRADIOL

BAUDAX

- * BAUDAX BIO INC
 - ANJESO, MELOXICAM

BAUSCH

- * BAUSCH HEALTH AMERICAS INC
 - ACANYA, BENZOYL PEROXIDE
 - ARAZLO, TAZAROTENE
 - BENZACLIN, BENZOYL PEROXIDE
 - BRYHALI, HALOBETASOL PROPIONATE
 - DUOBRII, HALOBETASOL PROPIONATE
 - EDECIN, ETHACRYNATE SODIUM
 - EDECIN, ETHACRYNIC ACID
 - EFUDEX, FLUOROURACIL
 - JUBLIA, EFINAACONAZOLE
 - LOCOID, HYDROCORTISONE BUTYRATE
 - MEPHYTON, PHYTONADIONE
 - ONEXTON, BENZOYL PEROXIDE
 - OXSORALEN-ULTRA, METHOXSALEN
 - SYPRINE, TRIENTINE HYDROCHLORIDE
 - TIMOPTIC IN OCUDOSE, TIMOLOL MALEATE
- * BAUSCH HEALTH IRELAND LTD
 - ERTACZO, SERTACONAZOLE NITRATE
 - TARGRETIN, BEXAROTENE
- * BAUSCH HEALTH US LLC
 - ALAWAY, KETOTIFEN FUMARATE (OTC)
 - ALDARA, IMIQUIMOD
 - AMMONUL, SODIUM BENZOATE
 - ANCOBON, FLUCYTOSINE
 - ALENZIN, BUPROPION HYDROBROMIDE
 - ATIVAN, LORAZEPAM
 - BRINZOLAMIDE, BRINZOLAMIDE
 - CALCIUM DISODIUM VERSENATE, EDETATE CALCIUM DISODIUM
 - CARDIZEM CD, DILTIAZEM HYDROCHLORIDE
 - CARDIZEM LA, DILTIAZEM HYDROCHLORIDE
 - CARDIZEM, DILTIAZEM HYDROCHLORIDE
 - CESAMET, NABILONE
 - CLINDAGEL, CLINDAMYCIN PHOSPHATE
 - D.H.E. 45, DIHYDROERGOTAMINE MESYLATE
 - DEMSEER, METYROSINE
 - DIASSTAT ACUDIAL, DIAZEPAM
 - DIASSTAT, DIAZEPAM
 - ELIDEL, PIMECROLIMUS
 - GRIS-PEG, GRISEOFULVIN, ULTRAMICROSIZED
 - ISORDIL, ISOSORBIDE DINITRATE
 - ISUPREL, ISOPROTERENOL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BAUSCH HEALTH US LLC

KLARON, SULFACETAMIDE SODIUM
 LIBRAX, CHLORDIAZEPOXIDE HYDROCHLORIDE
 LOCOID, HYDROCORTISONE BUTYRATE
 LOPROX, CICLOPIROX
 LUZU, LULICONAZOLE
 MESTINON, PYRIDOSTIGMINE BROMIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 METROGEL-VAGINAL, METRONIDAZOLE
 MIGRANAL, DIHYDROERGOTAMINE MESYLATE
 MINOCIN, MINOCYCLINE HYDROCHLORIDE
 SOLODYN, MINOCYCLINE HYDROCHLORIDE
 TASMAR, TOLCAPONE
 TIAZAC, DILTIAZEM HYDROCHLORIDE
 VANOS, FLUOCINONIDE
 VASERETIC, ENALAPRIL MALEATE
 VASOTEC, ENALAPRIL MALEATE
 VIRAZOLE, RIBAVIRIN
 WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
 XENAZINE, TETRABENAZINE
 XERESE, ACYCLOVIR
 ZELAPAR, SELEGILINE HYDROCHLORIDE
 ZIANA, CLINDAMYCIN PHOSPHATE
 ZOVIRAX, ACYCLOVIR
 ZYCLARA, IMIQUIMOD

BAUSCH AND LOMB

* BAUSCH AND LOMB INC

ALAWAY, KETOTIFEN FUMARATE (OTC)
 ALREX, LOTEPIREDNOL ETABONATE
 BESIVANCE, BESIFLOXACIN 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
 RETISERT, FLUOCINOLONE ACETONIDE
 SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TROPICAMIDE, TROPICAMIDE
 VYZULTA, LATANOPROSTENE BUNOD
 ZIRGAN, GANCICLOVIR
 ZYLET, LOTEPIREDNOL 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BAUSCH AND LOMB PHARMACEUTICALS INC
 NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC
 OFLOXACIN, OFLOXACIN
 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
 LOTEMAX SM, LOTEPIEDNOL ETABONATE
 LOTEMAX, LOTEPIEDNOL ETABONATE
 LUMIFY, BRIMONIDINE TARTRATE (OTC)
 XIPERE, TRIAMCINOLONE ACETONIDE

BAUSCH LOMB IRELAND

* BAUSCH AND LOMB IRELAND LTD
 FLUORESCEIN SODIUM AND BENOXINATE HYDROCHLORIDE, BENOXINATE HYDROCHLORIDE
 TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE

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
 CEFTRIAXONE IN PLASTIC CONTAINER, CEFTRIAXONE 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
 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
 CYTOXAN, CYCLOPHOSPHAMIDE
 DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BAXTER HEALTHCARE CORP

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 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 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
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BAXTER HEALTHCARE CORP

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%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 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
 PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS

BAXTER HLTHCARE CORP

* BAXTER HEALTHCARE CORP

ACETAMINOPHEN, ACETAMINOPHEN
 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
 CLINOLIPID 20%, OLIVE OIL
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOXIL (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
 EPTIFIBATIDE, EPTIFIBATIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BAXTER HEALTHCARE CORP
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FLUMAZENIL, FLUMAZENIL
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FUROSEMIDE, FUROSEMIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOFLOXACIN, LEVOFLOXACIN
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE IN 5% DEXTROSE, NOREPINEPHRINE BITARTRATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON 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
 TRANSDERM SCOP, SCOPOLAMINE
- * 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
 LAMPIT, NIFURTIMOX
 NUBEQA, DAROLUTAMIDE
 VITRAKVI, LAROTRECTINIB SULFATE

BAYER HEALTHCARE LLC

- * BAYER HEALTHCARE LLC
 CHILDREN'S CLARITIN, LORATADINE (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)
 ASTEPRO ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
 CHILDREN'S ASTEPRO ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
- * BAYER HEALTHCARE PHARMACEUTICALS INC
 ADEMPAS, RIOCIGUAT
 ANGELIQ, DROSPIRENONE
 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
 KERENDIA, FINERENONE
 KYLEENA, LEVONORGESTREL
 LEVITRA, VARDENAFIL HYDROCHLORIDE
 MENOSTAR, ESTRADIOL
 MIRENA, LEVONORGESTREL
 NATAZIA, DIENOGEST
 NEXAVAR, SORAFENIB TOSYLATE
 SAFYRAL, DROSPIRENONE
 SKYLA, LEVONORGESTREL
 STAXYN, VARDENAFIL HYDROCHLORIDE
 STIVARGA, REGORAFENIB
 ULTRAVIST (PHARMACY BULK), IOPROMIDE
 ULTRAVIST 300, IOPROMIDE
 ULTRAVIST 370, IOPROMIDE
 VITRAKVI, LAROTRECTINIB SULFATE
 XOFIGO, RADIUM RA-223 DICHLORIDE
 YASMIN, DROSPIRENONE
 YAZ, DROSPIRENONE

BAYSHORE PHARMS LLC*** BAYSHORE PHARMACEUTICALS LLC**

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 ETODOLAC, ETODOLAC
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 METOLAZONE, METOLAZONE
 PINDOLOL, PINDOLOL
 PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE

BDSI*** BIODELIVERY SCIENCES INTERNATIONAL INC**

BELBUCA, BUPRENORPHINE HYDROCHLORIDE
 ELYXYB, CELECOXIB
 SYMPROIC, NALDEMEDINE TOSYLATE

BE PHARMS*** BE PHARMACEUTICALS AG**

DAPTOMYCIN, DAPTOMYCIN
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 HEPARIN SODIUM, HEPARIN SODIUM
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE

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)

BEIGENE*** BEIGENE USA INC**

BRUKINSA, ZANUBRUTINIB

BEIJING*** BEIJING SCIECURE PHARMACEUTICAL CO LTD**

FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 NIACIN, NIACIN

BEIJING TIDE PHARM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BEIJING TIDE PHARMACEUTICAL CO LTD
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
- BEIJING YILING**
- * BEIJING YILING BIO-ENGINEERING AND TECHNOLOGY CO LTD
ANASTROZOLE, ANASTROZOLE
LETROZOLE, LETROZOLE
- BELCHER**
- * BELCHER PHARMACEUTICALS LLC
ABLYSINOL, ALCOHOL
AMINOCAPROIC ACID, AMINOCAPROIC ACID
CEFIXIME, CEFIXIME
EPINEPHRINE, EPINEPHRINE
MEFENAMIC ACID, MEFENAMIC ACID
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
TACROLIMUS, TACROLIMUS
- BELCHER PHARMS**
- * BELCHER PHARMACEUTICALS LLC
CEPHALEXIN, CEPHALEXIN
DESLORATADINE, DESLORATADINE
- BELOTECA INC**
- * BELOTECA INC
DIAZEPAM, DIAZEPAM
THIOTEPA, THIOTEPA
- BENUVIA**
- * BENUVIA THERAPEUTICS INC
SYNDROS, DRONABINOL
- BEXIMCO PHARMS USA**
- * BEXIMCO PHARMACEUTICALS USA INC
BACLOFEN, BACLOFEN
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
FLECAINIDE ACETATE, FLECAINIDE ACETATE
LOVASTATIN, LOVASTATIN
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHOCARBAMOL, METHOCARBAMOL
NADOLOL, NADOLOL
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
- BEXIMCO USA**
- * BEXIMCO PHARMACEUTICALS USA INC
CARVEDILOL, CARVEDILOL
- BIOCODEX SA**
- * BIOCODEX SA
DIACOMIT, STIRIPENTOL
- BIOCON PHARMA**
- * BIOCON PHARMA INC
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
- * BIOCON PHARMA LTD
EVEROLIMUS, EVEROLIMUS
MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
SIMVASTATIN, SIMVASTATIN
TACROLIMUS, TACROLIMUS
- BIOCRYST**
- * BIOCRYST PHARMACEUTICALS INC
ORLADEYO, BEROTRALSTAT HYDROCHLORIDE
RAPIVAB, PERAMIVIR
- BIOFRONTERA**
- * BIOFRONTERA BIOSCIENCE GMBH
AMELUZ, AMINOLEVULINIC ACID HYDROCHLORIDE
- BIOGEN IDEC**
- * BIOGEN IDEC INC
SPINRAZA, NUSINERSEN SODIUM
- BIOGEN INC**
- * BIOGEN INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******* BIOGEN INC**TECFIDERA, DIMETHYL FUMARATE
VUMERITY, DIROXIMEL FUMARATE**BIOHAVEN IRELAND***** BIOHAVEN PHARMACEUTICAL IRELAND DESIGNATED ACTIVITY CO**
NURTEC ODT, RIMEGEPANT SULFATE**BIOMARIN PHARM***** BIOMARIN PHARMACEUTICAL INC**
KUVAN, SAPROPTERIN DIHYDROCHLORIDE
VOXZOGO, VOSORITIDE**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
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
ATOVAQUONE, ATOVAQUONE
AZITHROMYCIN, AZITHROMYCIN
BENZONATATE, BENZONATATE
BEXAROTENE, BEXAROTENE
CALCITRIOL, CALCITRIOL
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
CLOBAZAM, CLOBAZAM
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
DEFERASIROX, DEFERASIROX
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
DOFETILIDE, DOFETILIDE
DROXIDOPA, DROXIDOPA
DUTASTERIDE, DUTASTERIDE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
ETHOSUXIMIDE, ETHOSUXIMIDE
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
IBUPROFEN, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
LORATADINE, LORATADINE (OTC)
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
MIDOL LIQUID GELS, IBUPROFEN (OTC)
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
NIMODIPINE, NIMODIPINE
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
PARICALCITOL, PARICALCITOL
POTASSIUM CITRATE, POTASSIUM CITRATE
PROGESTERONE, PROGESTERONE
RUFINAMIDE, RUFINAMIDE
SEVELAMER CARBONATE, SEVELAMER CARBONATE
TETRABENAZINE, TETRABENAZINE
VALPROIC ACID, VALPROIC ACID
VITAMIN D, ERGOCALCIFEROL
ZONISAMIDE, ZONISAMIDE**BIOPHARM***** BIOPHARM DISCOVERY LLC**
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
METHIMAZOLE, METHIMAZOLE
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BIOPHARM DISCOVERY LLC
VALPROIC ACID, VALPROIC ACID

BLUE EARTH

* BLUE EARTH DIAGNOSTICS LTD
AXUMIN, FLUCICLOVINE F-18

BLUEPRINT MEDICINES

* BLUEPRINT MEDICINES CORP
AYVAKIT, AVAPRITINIB

BOEHRINGER INGELHEIM

* BOEHRINGER INGELHEIM
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
TRIJARDY XR, EMPAGLIFLOZIN
VIRAMUNE XR, NEVIRAPINE
VIRAMUNE, NEVIRAPINE

BOSCOGEN

* BOSCOGEN INC
ARIPIPRAZOLE, ARIPIPRAZOLE
CAPTOPRIL, CAPTOPRIL
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
REPAGLINIDE, REPAGLINIDE
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE

BPI LABS LLC

* BPI LABS LLC
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
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
ENTERO VU 24%, 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BRACCO DIAGNOSTICS INC
 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 SMOOTHIE, 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
 VARIBAR THIN LIQUID, 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
 SUTAB, MAGNESIUM SULFATE

BRECKENRIDGE

* BRECKENRIDGE PHARMACEUTICAL INC
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ALPRAZOLAM, ALPRAZOLAM
 ASENAPINE MALEATE, ASENAPINE MALEATE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BRETILIUM TOSYLATE, BRETILIUM TOSYLATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTECAVIR, ENTECAVIR
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 EPLERENONE, EPLERENONE
 EVEROLIMUS, EVEROLIMUS
 EXEMESTANE, EXEMESTANE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 LAMIVUDINE, LAMIVUDINE
 LEVETIRACETAM, LEVETIRACETAM
 MEGESTROL ACETATE, MEGESTROL ACETATE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OMEPRAZOLE, OMEPRAZOLE
 PENICILLAMINE, PENICILLAMINE
 RIVASTIGMINE, RIVASTIGMINE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

BRECKENRIDGE PHARM

* BRECKENRIDGE PHARMACEUTICAL INC
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 TERBINAFFINE HYDROCHLORIDE, TERBINAFFINE HYDROCHLORIDE

BRIGHAM WOMENS

* BRIGHAM AND WOMENS HOSP
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

BRIGHAM WOMENS HOSP

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BRIGHAM AND WOMENS HOSP INC
AMMONIA N 13, AMMONIA N-13

BRIGHTGENE

* BRIGHTGENE BIO-MEDICAL TECHNOLOGY CO LTD
ENTECAVIR, ENTECAVIR

BRISTOL MYERS SQUIBB

* BRISTOL MYERS SQUIBB
AZACTAM, AZTREONAM
BARACLUDE, ENTECAVIR

* BRISTOL MYERS SQUIBB CO
DROXIA, HYDROXYUREA
HYDREA, HYDROXYUREA
REYATAZ, ATAZANAVIR SULFATE
SPRYCEL, DASATINIB
SUSTIVA, EFAVIRENZ

* BRISTOL MYERS SQUIBB CO PHARMACEUTICAL RESEARCH INSTITUTE
ELIQUIS, APIXABAN

BRISTOL-MYERS SQUIBB

* BRISTOL-MYERS SQUIBB CO
EVOTAZ, ATAZANAVIR SULFATE

BTCP PHARMA

* BTCP PHARMA LLC
LAZANDA, FENTANYL CITRATE
SUBSYS, FENTANYL

BTG INTL

* BTG INTERNATIONAL INC
THYROSAFE, POTASSIUM IODIDE (OTC)

BWXT ITG

* BWXT ITG CANADA INC
INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE

**** C ******CADILA**

* CADILA HEALTHCARE LTD
ACETAZOLAMIDE, ACETAZOLAMIDE
ACYCLOVIR, ACYCLOVIR
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
DESONIDE, DESONIDE
DESOXIMETASONE, DESOXIMETASONE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DUTASTERIDE, DUTASTERIDE
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
FELBAMATE, FELBAMATE
FLUOCINONIDE, FLUOCINONIDE
GEMFIBROZIL, GEMFIBROZIL
INDOMETHACIN, INDOMETHACIN
LEVOFLOXACIN, LEVOFLOXACIN
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METRONIDAZOLE, METRONIDAZOLE
MODAFINIL, MODAFINIL
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
NYSTATIN, NYSTATIN
PIROXICAM, PIROXICAM
RANOLAZINE, RANOLAZINE
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
VORICONAZOLE, VORICONAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CADILA HEALTHCARE LTD
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZONISAMIDE, ZONISAMIDE

CADILA PHARMS LTD

* CADILA PHARMACEUTICALS LTD
 ACYCLOVIR, ACYCLOVIR
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CELECOXIB, CELECOXIB
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 GEMFIBROZIL, GEMFIBROZIL
 GLYBURIDE, GLYBURIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 NATEGLINIDE, NATEGLINIDE
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 OFLOXACIN, OFLOXACIN
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 TELMISARTAN, TELMISARTAN
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

CALL INC

* CALL INC DBA ROCHESTER PHARMACEUTICALS
 ADAPALENE, ADAPALENE

CALLIDITAS

* CALLIDITAS THERAPEUTICS AB
 TARPEYO, BUDESONIDE

CAPELLON PHARMS LLC

* CAPELLON PHARMACEUTICALS LLC
 POLMON, DEXCHLORPHENIRAMINE MALEATE

CAPLIN

* CAPLIN STERILES LTD
 ARGATROBAN, ARGATROBAN
 ETOMIDATE, ETOMIDATE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

CARA THERAP

* CARA THERAPEUTICS INC
 KORSUVA, DIFELIKEFALIN ACETATE

CARDINAL HEALTH 414

* CARDINAL HEALTH 414 LLC CARDINAL HEALTH NUCLEAR PHARMACY SERVICES
 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
 TECHNETIUM TC-99M MEDRONATE KIT, TECHNETIUM TC-99M MEDRONATE KIT

CARDINAL HEALTH 418

* CARDINAL HEALTH 418 INC
 SODIUM IODIDE I 123, SODIUM IODIDE I-123

CARDINAL HLTH 414

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CARDINAL HEALTH 414 LLC
AMMONIA N 13, AMMONIA N-13

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
CILOSTAZOL, CILOSTAZOL
ENTECAVIR, ENTECAVIR
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
TENOFVIR DISOPROXIL FUMARATE, TENOFVIR DISOPROXIL FUMARATE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE

CASPER PHARMA LLC

* CASPER PHARMA LLC
ANTIVERT, MECLIZINE HYDROCHLORIDE
AQUASOL A, VITAMIN A PALMITATE
CASPORYN HC, HYDROCORTISONE
FURADANTIN, NITROFURANTOIN
LUMI-SPORYN, BACITRACIN ZINC
ROBINUL FORTE, GLYCOPYRROLATE
ROBINUL, GLYCOPYRROLATE
THALITONE, CHLORTHALIDONE
ZYLOPRIM, ALLOPURINOL

CATALENT

* CATALENT PHARMA SOLUTIONS LLC
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
VALPROIC ACID, VALPROIC ACID

CATALYST PHARMS

* CATALYST PHARMACEUTICALS INC
FIRDAPSE, AMIFAMPRIDINE PHOSPHATE

CEDIPROF INC

* CEDIPROF INC
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
LEVO-T, LEVOTHYROXINE SODIUM **

CELATOR PHARMS

* CELATOR PHARMACEUTICALS INC
VYXEOS, CYTARABINE

CELGENE

* CELGENE CORP
ISTODAX, ROMIDEPSIN
POMALYST, POMALIDOMIDE
REVLIMID, LENALIDOMIDE
THALOMID, THALIDOMIDE
VIDAZA, AZACITIDINE

CELGENE CORP

* CELGENE CORP
IDHIFA, ENASIDENIB MESYLATE
ONUREG, AZACITIDINE

CELGENE INTL

* CELGENE INTERNATIONAL II SARL
ZEPOSIA, OZANIMOD HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******CELLTRION**

* CELLTRION INC
 DEFERASIROX, DEFERASIROX
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 FAMOTIDINE, FAMOTIDINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LINEZOLID, LINEZOLID
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 RISPERIDONE, RISPERIDONE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TELMISARTAN, TELMISARTAN

CENTAUR PHARMS PVT

* CENTAUR PHARMACEUTICALS PVT LTD
 IVABRADINE HYDROCHLORIDE, IVABRADINE HYDROCHLORIDE

CEPHALON

* CEPHALON INC
 ACTIQ, FENTANYL CITRATE
 FENTORA, FENTANYL CITRATE
 GABITRIL, TIAGABINE HYDROCHLORIDE
 NUVIGIL, ARMODAFINIL
 PROVIGIL, MODAFINIL
 TREANDA, BENDAMUSTINE HYDROCHLORIDE
 TRISENOX, ARSENIC TRIOXIDE

CEROVENE INC

* CEROVENE INC
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 PYRIMETHAMINE, PYRIMETHAMINE

CHANGZHOU PHARM

* CHANGZHOU PHARMACEUTICAL FACTORY
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 PREGABALIN, PREGABALIN
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

CHARTWELL

* CHARTWELL LIFE MOLECULES LLC
 ALLOPURINOL, ALLOPURINOL
 AMOXICILLIN, AMOXICILLIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLARITHROMYCIN, CLARITHROMYCIN
 COLOCORT, HYDROCORTISONE
 FLUCONAZOLE, FLUCONAZOLE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 OXAPROZIN, OXAPROZIN
 TEMOZOLOMIDE, TEMOZOLOMIDE

* CHARTWELL PHARMA SCIENCE LLC
 SULFASALAZINE, SULFASALAZINE

* CHARTWELL SCHEDULED LLC
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN

CHARTWELL MOLECULAR

* CHARTWELL MOLECULAR HOLDINGS LLC
 CALCIUM ACETATE, CALCIUM ACETATE
 CARVEDILOL, CARVEDILOL
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 FOLIC ACID, FOLIC ACID
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IRBESARTAN, IRBESARTAN
 RAMIPRIL, RAMIPRIL
 RISPERIDONE, RISPERIDONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******CHARTWELL MOLECULES**

* CHARTWELL MOLECULES LLC
 DISULFIRAM, DISULFIRAM
 GEMFIBROZIL, GEMFIBROZIL
 NABUMETONE, NABUMETONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

CHARTWELL RX

* CHARTWELL RX SCIENCES LLC
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 CALCIUM ACETATE, CALCIUM ACETATE
 CEFPROZIL, CEFPROZIL
 CEPHALEXIN, CEPHALEXIN
 CILOSTAZOL, CILOSTAZOL
 DUVOID, BETHANECHOL CHLORIDE
 EPLERENONE, EPLERENONE
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 FULVICIN P/G 165, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 FULVICIN P/G 330, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 FULVICIN P/G, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 FULVICIN-U/F, GRISEOFULVIN, MICROCRYSTALLINE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 INDOMETHACIN, INDOMETHACIN
 LACTULOSE, LACTULOSE
 LEVETIRACETAM, LEVETIRACETAM
 LISINOPRIL, LISINOPRIL
 METHIMAZOLE, METHIMAZOLE
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 MONODOX, DOXYCYCLINE
 NABUMETONE, NABUMETONE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PREDNISOLONE, PREDNISOLONE
 REPAGLINIDE, REPAGLINIDE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 URSODIOL, URSODIOL
 VALPROIC ACID, VALPROIC ACID
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

CHARTWELL TETRA

* CHARTWELL TETRA LLC
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

CHATTEM

* CHATTEM INC
 UNISOM, DOXYLAMINE SUCCINATE (OTC)

CHATTEM SANOFI

* CHATTEM INC DBA SANOFI CONSUMER HEALTHCARE
 ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 GAVISCON, ALUMINUM HYDROXIDE (OTC)
 NICODERM CQ, NICOTINE (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
 NUVESSA, METRONIDAZOLE

CHEMOCENTRYX

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CHEMOCENTRYX INC
TAVNEOS, AVACOPAN

CHEPLAPHARM

* CHEPLAPHARM ARZNEIMITTEL GMBH
ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
KLONOPIN, CLONAZEPAM
XENICAL, ORLISTAT

CHIA TAI TIANQING

* CHIA TAI TIANQING PHARMACEUTICAL GROUP CO LTD
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
FULVESTRANT, FULVESTRANT

CHIASMA

* CHIASMA INC
MYCAPSSA, OCTREOTIDE ACETATE

CHIESI

* CHIESI USA INC
BETHKIS, TOBRAMYCIN
BRONCHITOL, MANNITOL
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
FERRIPROX, DEFERIPRONE
KENGREAL, CANGRELOR
ZYFLO CR, ZILEUTON
ZYFLO, ZILEUTON

CHILDRENS HOSP MI

* CHILDRENS HOSP MICHIGAN
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

CHIMERIX

* CHIMERIX INC
TEMBEXA, BRINCIDOFOVIR

CHINA RESOURCES

* CHINA RESOURCES SAIKE PHARMACEUTICAL CO LTD
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

CHIRHOCLIN

* CHIRHOCLIN INC
CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN

CINTEX SVCS

* CINTEX SERVICES LLC
DESONIDE, DESONIDE
FLURANDRENOLIDE, FLURANDRENOLIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

CIPHER PHARMS INC

* CIPHER PHARMACEUTICALS INC
CONZIP, TRAMADOL HYDROCHLORIDE
LIPOFEN, FENOFIBRATE

CIPLA

* CIPLA LTD
ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
ABACAVIR SULFATE, ABACAVIR SULFATE
ACYCLOVIR, ACYCLOVIR
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALENDRONATE SODIUM, ALENDRONATE SODIUM
AMBRISANTAN, AMBRISANTAN
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ANASTROZOLE, ANASTROZOLE
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
AZACITIDINE, AZACITIDINE
BUDESONIDE, BUDESONIDE
CELECOXIB, CELECOXIB
CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******* CIPLA LTD**

CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DEFERASIROX, DEFERASIROX
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIFLUPREDNATE, DIFLUPREDNATE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 EFAVIRENZ, EFAVIRENZ
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EMTRICITABINE, EMTRICITABINE
 ENTECAVIR, ENTECAVIR
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 EXEMESTANE, EXEMESTANE
 FAMCICLOVIR, FAMCICLOVIR
 FENOFIBRATE, FENOFIBRATE
 FINASTERIDE, FINASTERIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 MELOXICAM, MELOXICAM
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NEVIRAPINE, NEVIRAPINE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PREGABALIN, PREGABALIN
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 SUMATRIPTAN, SUMATRIPTAN
 TADALAFIL, TADALAFIL
 TAVABOROLE, TAVABOROLE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TESTOSTERONE, TESTOSTERONE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 ZIDOVUDINE, ZIDOVUDINE

CIPLA LTD*** CIPLA LTD**

ALBENDAZOLE, ALBENDAZOLE
 CARBOPLATIN, CARBOPLATIN
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 TOPIRAMATE, TOPIRAMATE
 ZALEPLON, ZALEPLON
 ZIDOVUDINE, ZIDOVUDINE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

CIPLA USA*** CIPLA USA INC**

ZEMDRI, PLAZOMICIN SULFATE

CISEN*** CISEN PHARMACEUTICAL CO LTD**

ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM

CLARUS*** CLARUS THERAPEUTICS INC**

JATENZO, TESTOSTERONE UNDECANOATE

CLINIGEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CLINIGEN INC
ETHYOL, AMIFOSTINE

CLINIGEN HLTHCARE

* CLINIGEN HEALTHCARE LTD
FOSCAVIR, FOSCARNET SODIUM

CLIVUNEL INC

* CLINUVEL INC
SCENESSE, AFAMELANOTIDE

CLOVIS ONCOLOGY INC

* CLOVIS ONCOLOGY INC
RUBRACA, RUCAPARIB CAMSYLATE

CMP DEV LLC

* CMP DEVELOPMENT LLC
CAROSPIR, SPIRONOLACTONE
POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC

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
TRIANEX, TRIAMCINOLONE ACETONIDE

CODY LABS INC

* CODY LABORATORIES INC A WHOLLY OWNED SUBSIDIARY OF LANNETT CO INC
NUMBRINO, COCAINE HYDROCHLORIDE

COLGATE PALMOLIVE CO

* COLGATE PALMOLIVE CO
PERIOGARD, CHLORHEXIDINE GLUCONATE

COLGATE-PALMOLIVE CO

* COLGATE-PALMOLIVE CO
PERIOGARD, CHLORHEXIDINE GLUCONATE

COLLEGIUM PHARM INC

* COLLEGIUM PHARMACEUTICAL INC
NUCYNTA ER, TAPENTADOL HYDROCHLORIDE
NUCYNTA, TAPENTADOL HYDROCHLORIDE
XTAMPZA ER, OXYCODONE

COMBE

* COMBE INC
VAGISTAT-1, TIOCONAZOLE (OTC)

COMMAVE THERAP

* COMMAVE THERAPEUTICS SA
AZSTARYS, DEXMETHYLPHENIDATE HYDROCHLORIDE

CONCORD BIOTECH LTD

* CONCORD BIOTECH LTD
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
TACROLIMUS, TACROLIMUS

CONCORDIA

* CONCORDIA PHARMACEUTICALS INC
DIBENZYLINE, PHENOXYBENZAMINE HYDROCHLORIDE
DYRENIUM, TRIAMTERENE
LANOXIN, DIGOXIN
NILANDRON, NILUTAMIDE
PANRETIN, ALITRETINOIN
PARNATE, TRANILCYPROMINE SULFATE
PLAQUENIL, HYDROXYCHLOROQUINE SULFATE
SALAGEN, PILOCARPINE HYDROCHLORIDE
UROXATRAL, ALFUZOSIN HYDROCHLORIDE
ZONEGRAN, ZONISAMIDE

CONCORDIA PHARMS INC

* CONCORDIA PHARMACEUTICALS INC
KAPVAY, CLONIDINE HYDROCHLORIDE
ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******CONTRACT PHARMACAL**

* CONTRACT PHARMACAL CORP
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 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

COREPHARMA

* COREPHARMA LLC
 LISINOPRIL, LISINOPRIL
 LOVASTATIN, LOVASTATIN
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TORSEMIDE, TORSEMIDE

COSETTE

* COSETTE PHARMACEUTICALS INC
 ACEPHEN, ACETAMINOPHEN (OTC)
 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
 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
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 INDOMETHACIN, INDOMETHACIN
 LIDOCAINE, LIDOCAINE
 METRONIDAZOLE, METRONIDAZOLE
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MIGERGOT, CAFFEINE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MYKACET, NYSTATIN
 NYSTATIN, NYSTATIN
 PROCHLORPERAZINE, PROCHLORPERAZINE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROMETHEGAN, PROMETHAZINE HYDROCHLORIDE
 TAZAROTENE, TAZAROTENE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

COVIS

* COVIS PHARMA GMBH
 ALTOPREV, LOVASTATIN
 ALVESCO, CICLESONIDE
 BETAPACE AF, SOTALOL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* COVIS PHARMA GMBH
 BETAPACE, SOTALOL HYDROCHLORIDE
 FERAHEME, FERUMOXYTOL
 LANOXIN PEDIATRIC, DIGOXIN
 LANOXIN, DIGOXIN
 MAKENA (AUTOINJECTOR), HYDROXYPROGESTERONE CAPROATE
 MAKENA PRESERVATIVE FREE, HYDROXYPROGESTERONE CAPROATE
 MAKENA, HYDROXYPROGESTERONE CAPROATE
 OMNARIS, CICLESONIDE
 PRILOSEC, OMEPRAZOLE MAGNESIUM
 RILUTEK, RILUZOLE
 SULAR, NISOLDIPINE
 ZANAFLEX, TIZANIDINE HYDROCHLORIDE
 ZETONNA, CICLESONIDE

CPPI CV

* CP PHARMACEUTICALS INTERNATIONAL CV
 SUTENT, SUNITINIB MALATE

CROSSMEDIKA SA

* CROSSMEDIKA SA
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE

CROWN LABS

* CROWN LABORATORIES INC
 ALA-CORT, HYDROCORTISONE
 TRIDERM, TRIAMCINOLONE ACETONIDE

CROWN LABS INC

* CROWN LABORATORIES INC
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN

CSPC OUYI

* CSPC OUYI PHARMACEUTICAL CO LTD
 AZITHROMYCIN, AZITHROMYCIN
 CARBAMAZEPINE, CARBAMAZEPINE
 CELECOXIB, CELECOXIB
 CONJUPRI, LEVAMLODIPINE MALEATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 GABAPENTIN, GABAPENTIN
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PALIPERIDONE, PALIPERIDONE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PREGABALIN, PREGABALIN

CSPC OUYI PHARM CO

* CSPC OUYI PHARMACEUTICAL CO LTD
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

CSPC-NBP PHARM

* CSPC-NBP PHARMACEUTICAL CO LTD
 BENZONATATE, BENZONATATE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS

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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CUMBERLAND PHARMACEUTICALS INC
 ACETADOTE, ACETYLCYSTEINE
 CALDOLOR, IBUPROFEN
 LACTULOSE, LACTULOSE
 REDITREX, METHOTREXATE
 VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE
 VIBATIV, TELAVANCIN HYDROCHLORIDE

CURIUM

* CURIUM US 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

CURRAX

* CURRAX PHARMACEUTICALS LLC
 ONZETRA XSAIL, SUMATRIPTAN SUCCINATE
 SILENOR, DOXEPIN HYDROCHLORIDE
 TREXIMET, NAPROXEN SODIUM

CUSTOPHARM INC

* CUSTOPHARM INC
 ACETAMINOPHEN, ACETAMINOPHEN
 CALCITONIN-SALMON, CALCITONIN SALMON
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
 SODIUM TETRADECYL SULFATE, SODIUM TETRADECYL SULFATE
 VALRUBICIN, VALRUBICIN

CYCLE PHARMS LTD

* CYCLE PHARMACEUTICALS LTD
 HALOPERIDOL, HALOPERIDOL
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 NITYR, NITISINONE

**** 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
 EFFIENT, PRASUGREL HYDROCHLORIDE
 EVOXAC, CEVIMELINE HYDROCHLORIDE
 SAVAYSA, EDOXABAN TOSYLATE
 TURALIO, PEXIDARTINIB HYDROCHLORIDE

DANCO LABS LLC

* DANCO LABORATORIES LLC
 MIFEPREX, MIFEPRISTONE

DARE

* DARE BIOSCIENCE INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DARE BIOSCIENCE INC
XACIATO, CLINDAMYCIN PHOSPHATE

DASH PHARMS

* DASH PHARMACEUTICALS LLC
TESTOSTERONE, TESTOSTERONE
ZILEUTON, ZILEUTON

DASTECH GENERICS

* DASTECH GENERICS LLC
NYSTATIN, NYSTATIN

DAVA PHARMS INC

* DAVA PHARMACEUTICALS INC
AMOXICILLIN, AMOXICILLIN
MORPHINE SULFATE, MORPHINE SULFATE
PROPYLTHIOURACIL, PROPYLTHIOURACIL

DAVIS AND GECK

* DAVIS AND GECK DIV AMERICAN CYANAMID CO
PRE-OP II, HEXACHLOROPHENE
PRE-OP, HEXACHLOROPHENE

DBL PHARMS

* DBL PHARMACEUTICALS INC
METHOCARBAMOL, METHOCARBAMOL

DECATUR

* DECATUR MEMORIAL HOSP
AMMONIA N 13, AMMONIA N-13
CHOLINE C-11, CHOLINE C-11
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

DECIPHERA PHARMS

* DECIPHERA PHARMACEUTICALS LLC
QINLOCK, RIPRETINIB

DENTSPLY PHARM

* DENTSPLY PHARMACEUTICAL INC
CITANEST FORTE DENTAL, EPINEPHRINE BITARTRATE
ORAQIX, LIDOCAINE

DEPROCO

* DEPROCO INC
LIGNOSPAN FORTE, EPINEPHRINE BITARTRATE
LIGNOSPAN STANDARD, EPINEPHRINE BITARTRATE
SCANDONEST L, LEVONORDEFIN
SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
SEPTOCAINE, ARTICAINE HYDROCHLORIDE

DEVA HOLDING AS

* DEVA HOLDING AS
ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
TEMOZOLOMIDE, TEMOZOLOMIDE

DEXCEL LTD

* DEXCEL LTD
DICLOFENAC SODIUM, DICLOFENAC SODIUM
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE

DEXCEL PHARMA

* DEXCEL PHARMA TECHNOLOGIES LTD
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
HEMADY, DEXAMETHASONE
LANSOPRAZOLE, LANSOPRAZOLE (OTC)
OMEPRAZOLE, OMEPRAZOLE (OTC)
PERIOCHIP, CHLORHEXIDINE GLUCONATE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
VIGABATRIN, VIGABATRIN

DIALYSIS SUPS

* DIALYSIS SUPPLIES INC
NORMOCARB HF 25, MAGNESIUM CHLORIDE
NORMOCARB HF 35, MAGNESIUM CHLORIDE

DORC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DORC INTERNATIONAL BV
 MEMBRANEBLUE, TRYPAN BLUE
 VISIONBLUE, TRYPAN BLUE

DOUGLAS PHARMS

* DOUGLAS PHARMACEUTICALS AMERICA LTD
 MYORISAN, ISOTRETINOIN

DOW PHARM

* DOW PHARMACEUTICAL SCIENCES
 ALTRENO, TRETINOIN
 ATRALIN, TRETINOIN

DR REDDYS

* DR REDDYS LABORATORIES INC
 FENOFIBRATE, FENOFIBRATE
 ICOSAPENT ETHYL, ICOSAPENT ETHYL
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 NITROGLYCERIN, NITROGLYCERIN
 PROGESTERONE, PROGESTERONE
 PROPOFOL, PROPOFOL
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TESTOSTERONE, TESTOSTERONE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

* DR REDDYS LABORATORIES LTD
 ALBENDAZOLE, ALBENDAZOLE
 AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 AZACITIDINE, AZACITIDINE
 CARFILZOMIB, CARFILZOMIB
 CARMUSTINE, CARMUSTINE
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 COLCHICINE, COLCHICINE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DAPTOMYCIN, DAPTOMYCIN
 DECITABINE, DECITABINE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FULVESTRANT, FULVESTRANT
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 LENALIDOMIDE, LENALIDOMIDE
 LIDOCAINE, LIDOCAINE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
 PARICALCITOL, PARICALCITOL
 PENICILLAMINE, PENICILLAMINE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PREGABALIN, PREGABALIN
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 SAPROPTERIN DIHYDROCHLORIDE, SAPROPTERIN DIHYDROCHLORIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TETRABENAZINE, TETRABENAZINE
 THIOTEPA, THIOTEPA
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALSARTAN, VALSARTAN
 VIGABATRIN, VIGABATRIN
 VINORELBINE TARTRATE, VINORELBINE TARTRATE

DR REDDYS LA

* DR REDDYS LABORATORIES LOUISIANA LLC
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DR REDDYS LABORATORIES LOUISIANA LLC
 LOPURIN, ALLOPURINOL
 SSD, SILVER SULFADIAZINE

DR REDDYS LABS INC

* DR REDDYS LABORATORIES INC
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 FINASTERIDE, FINASTERIDE
 FLUCONAZOLE, FLUCONAZOLE
 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)
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 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

DR REDDYS LABS LTD

* DR REDDYS LABORATORIES LIMITED
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE

* DR REDDYS LABORATORIES LTD
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BIVALIRUDIN, BIVALIRUDIN
 BORTEZOMIB, BORTEZOMIB
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CAPECITABINE, CAPECITABINE
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 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 AND DEXAMETHASONE, CIPROFLOXACIN
 CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLOFARABINE, CLOFARABINE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 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
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 ESLICARBAZEPINE ACETATE, ESLICARBAZEPINE ACETATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 ESZOPICLONE, ESZOPICLONE
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FEBUXOSTAT, FEBUXOSTAT
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DR REDDYS LABORATORIES LTD
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FINASTERIDE, FINASTERIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLIMEPIRIDE, GLIMEPIRIDE
 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)
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LAMOTRIGINE, LAMOTRIGINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LATANOPROST, LATANOPROST
 LETROZOLE, LETROZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 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 AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NATEGLINIDE, NATEGLINIDE
 NIZATIDINE, NIZATIDINE
 OFLOXACIN, OFLOXACIN
 OLANZAPINE, OLANZAPINE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 OMEPRAZOLE, OMEPRAZOLE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXAPROZIN, OXAPROZIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PHYTONADIONE, PHYTONADIONE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RAMIPRIL, RAMIPRIL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 RISPERIDONE, RISPERIDONE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SIROLIMUS, SIROLIMUS
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 TACROLIMUS, TACROLIMUS
 TADALAFIL, TADALAFIL
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VIGABATRIN, VIGABATRIN
 ZAFIRLUKAST, ZAFIRLUKAST
 ZENATANE, ISOTRETINOIN
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

DR REDDYS LABS SA

* DR REDDYS LABORATORIES SA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DR REDDYS LABORATORIES SA
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 HABITROL, NICOTINE (OTC)
 MERCAPTOPYRINE, MERCAPTOPYRINE
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 RAMELTEON, RAMELTEON
 TOBRAMYCIN, TOBRAMYCIN
 VERARING, ETHINYL ESTRADIOL

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
 VELIVET, DESOGESTREL

DUSA

* DUSA PHARMACEUTICALS INC
 LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE

DUTCH OPHTHALMIC

* DUTCH OPHTHALMIC RESEARCH CENTER INTERNATIONAL BV
 TISSUEBLUE, BRILLIANT BLUE G

REDDYS

* DOCTOR REDDYS LABORATORIES LTD
 DESLORATADINE, DESLORATADINE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE

**** E ******E5 PHARMA INC**

* E5 PHARMA INC
 DIAZOXIDE, DIAZOXIDE

EAGLE PHARMS

* EAGLE PHARMACEUTICALS INC
 BELRAPZO, BENDAMUSTINE HYDROCHLORIDE
 BENDEKA, BENDAMUSTINE HYDROCHLORIDE
 PEMFEXY, PEMETREXED
 RYANODEX, DANTROLENE SODIUM
 VASOPRESSIN, VASOPRESSIN

ECI PHARMS LLC

* ECI PHARMACEUTICALS LLC
 MONOKET, ISOSORBIDE MONONITRATE

ECOLAB

* ECOLAB INC
 CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)
 CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)

ECR PHARMA

* ECR PHARMA
 TUSSICAPS, CHLORPHENIRAMINE POLISTIREX

EDENBRIDGE PHARMS

* EDENBRIDGE PHARMACEUTICALS LLC
 ALBENDAZOLE, ALBENDAZOLE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 CARBIDOPA, CARBIDOPA
 DARTISLA ODT, GLYCOPYRROLATE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 ETODOLAC, ETODOLAC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ******* EDENBRIDGE PHARMACEUTICALS LLC**

IVERMECTIN, IVERMECTIN
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 MIGLUSTAT, MIGLUSTAT
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 TINIDAZOLE, TINIDAZOLE

EDISON THERAPS LLC

* EDISON THERAPEUTICS LLC
 METHERGINE, METHYLERGONOVINE MALEATE

EGALET

* EGALET US INC
 INDOCIN, INDOMETHACIN

EI INC

* EI INC
 THEROXIDIL, MINOXIDIL (OTC)

EIGER BIOPHARMS

* EIGER BIOPHARMACEUTICALS INC
 ZOKINVY, LONAFARNIB

EIRGEN

* EIRGEN PHARMA LTD
 RAYALDEE, CALCIFEDIOL

EISAI INC

* EISAI INC
 ARICEPT, DONEPEZIL HYDROCHLORIDE
 BANZEL, RUFINAMIDE
 DAYVIGO, LEMBOREXANT
 FYCOMPA, PERAMPANEL
 HALAVEN, ERIBULIN MESYLATE
 LENVIMA, LENVATINIB MESYLATE

ELI LILLY AND CO

* ELI LILLY AND CO
 BAQSIMI, GLUCAGON
 OLUMIANT, BARICITINIB
 PROZAC, FLUOXETINE HYDROCHLORIDE
 REYVOW, LASMIDITAN SUCCINATE
 VERZENIO, ABEMACICLIB

ELI LILLY CO

* ELI LILLY CO
 ADCIRCA, TADALAFIL
 ZYPREXA RELPREVV, OLANZAPINE PAMOATE

ELITE LABS

* ELITE LABORATORIES INC
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

ELITE LABS INC

* ELITE LABORATORIES INC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 ISRADIPINE, ISRADIPINE
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE

ELITE PHARM SOLUTION

* ELITE PHARMACEUTICAL SOLUTION INC
 NIFEDIPINE, NIFEDIPINE

ELYSIUM

* ELYSIUM PHARMACEUTICALS LTD
 CALCITRIOL, CALCITRIOL
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 TIMOLOL MALEATE, TIMOLOL MALEATE

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
 CIDOFOVIR, CIDOFOVIR
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ETOMIDATE, ETOMIDATE
 FUROSEMIDE, FUROSEMIDE
 METOCLOPRAMIDE, METOCLOPRAMIDE HYDROCHLORIDE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROPOFOL, PROPOFOL
 TRANEXAMIC ACID, TRANEXAMIC ACID

EMD SERONO INC

* EMD SERONO INC
 CETROTIDE, CETRORELIX
 MAVENCLAD, CLADRIBINE
 TEPMETKO, TEPOTINIB HYDROCHLORIDE

EMED MEDCL

* EMED MEDICAL CO LLC
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE

EMERALD INTL LTD

* EMERALD INTERNATIONAL LTD
 BACLOFEN, BACLOFEN

EMERGENT

* EMERGENT OPERATIONS IRELAND LTD
 NARCAN, NALOXONE HYDROCHLORIDE

EMMAUS MEDCL

* EMMAUS MEDICAL INC
 ENDARI, L-GLUTAMINE

ENALTEC

* ENALTEC LABS INC
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE

ENCUBE

* ENCUBE ETHICALS PRIVATE LTD
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
 KETOCONAZOLE, KETOCONAZOLE
 MUPIROCIN, MUPIROCIN CALCIUM
 TAVABOROLE, TAVABOROLE
 TESTOSTERONE, TESTOSTERONE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

ENCUBE ETHICALS

* ENCUBE ETHICALS PVT LTD
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 DESONIDE, DESONIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 PERMETHRIN, PERMETHRIN

ENDO PHARM

* ENDO PHARMACEUTICAL SOLUTIONS INC
 SUPPRELIN LA, HISTRELIN ACETATE
 VALSTAR PRESERVATIVE FREE, VALRUBICIN

ENDO PHARMS

* ENDO PHARMACEUTICALS INC
 FORTESTA, TESTOSTERONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* ENDO PHARMACEUTICALS INC
 FROVA, FROVATRIPTAN SUCCINATE
 PERCODAN, ASPIRIN

ENDO PHARMS INC

* ENDO PHARMACEUTICALS INC
 AVEED, TESTOSTERONE UNDECANOATE
 COLY-MYCIN S, COLISTIN SULFATE
 MEGACE ES, MEGESTROL ACETATE
 NASCOBAL, CYANOCOBALAMIN

EPI HLTH

* EPI HEALTH LLC
 CLODERM, CLOCORTOLONE PIVALATE
 MINOLIRA, MINOCYCLINE HYDROCHLORIDE
 RHOFAD, OXYMETAZOLINE 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
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 AZITHROMYCIN, AZITHROMYCIN
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 DEMECLOXYCLINE HYDROCHLORIDE, DEMECLOXYCLINE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE HCLATE, DOXYCYCLINE HCLATE
 FUROSEMIDE, FUROSEMIDE
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GLYBURIDE, GLYBURIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 MOLINDONE HYDROCHLORIDE, MOLINDONE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ASPIRIN, ASPIRIN
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ******EPIZYME INC**

* EPIZYME INC
TAZVERIK, TAZEMETOSTAT HYDROBROMIDE

ESPERION THERAPS INC

* ESPERION THERAPEUTICS INC
NEXLETOL, BEMPEDOIC ACID
NEXLIZET, BEMPEDOIC ACID

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 SA
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM

ETHYPHARM USA CORP

* ETHYPHARM USA CORP
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE

ETON

* ETON PHARMACEUTICALS INC
ALKINDI SPRINKLE, HYDROCORTISONE
REZIPRES, EPHEDRINE HYDROCHLORIDE

ETON PHARMS

* ETON PHARMACEUTICALS
BIORPHEN, PHENYLEPHRINE HYDROCHLORIDE

EUGIA PHARMA

* EUGIA PHARMA SPECIALITIES LTD
ACETAMINOPHEN, ACETAMINOPHEN
ACETYLCYSTEINE, ACETYLCYSTEINE
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
ADENOSINE, ADENOSINE
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
ANASTROZOLE, ANASTROZOLE
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
AZITHROMYCIN, AZITHROMYCIN
BIVALIRUDIN, BIVALIRUDIN
BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
CAFFEINE CITRATE, CAFFEINE CITRATE
CAPECITABINE, CAPECITABINE
CARBOPLATIN, CARBOPLATIN
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
CYANOCOBALAMIN, CYANOCOBALAMIN
DACTINOMYCIN, DACTINOMYCIN
DAPTOMYCIN, DAPTOMYCIN
DECITABINE, DECITABINE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
DOCETAXEL, DOCETAXEL
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
EPHEDRINE SULFATE, EPHEDRINE SULFATE
EPTIFIBATIDE, EPTIFIBATIDE
ERTAPENEM SODIUM, ERTAPENEM SODIUM
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EUGIA PHARMA SPECIALITIES LTD
 ETOMIDATE, ETOMIDATE
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FULVESTRANT, FULVESTRANT
 FUROSEMIDE, FUROSEMIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 ISOSULFAN BLUE, ISOSULFAN BLUE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 LETROZOLE, LETROZOLE
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LINEZOLID, LINEZOLID
 MEROPENEM, MEROPENEM
 METHOCARBAMOL, METHOCARBAMOL
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 PROGESTERONE, PROGESTERONE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TIGECYCLINE, TIGECYCLINE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

EUGIA PHARMA SPECLTS

* EUGIA PHARMA SPECIALITIES LTD
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE

EUROHLTH INTL SARL

* EUROHEALTH INTERNATIONAL SARL
 AZACITIDINE, AZACITIDINE
 DROPERIDOL, DROPERIDOL
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE

EVOFEM INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EVOFEM INC

PHEXXI, CITRIC ACID

EVOKE PHARMA INC

* EVOKE PHARMA INC

GIMOTI, METOCLOPRAMIDE HYDROCHLORIDE

EVUS

* EVUS HEALTH SOLUTIONS LLC

NITROMIST, NITROGLYCERIN

EXALENZ BIOSCIENCE

* EXALENZ BIOSCIENCE LTD

IDKIT:HP, CITRIC ACID

EXELA PHARMA

* EXELA PHARMA SCIENCES LLC

ACETYLCYSTEINE, ACETYLCYSTEINE

AKOVAZ, EPHEDRINE SULFATE

BLOXIVERZ, NEOSTIGMINE METHYLSULFATE

CAFFEINE CITRATE, CAFFEINE CITRATE

DILTIAZEM HYDROCHLORIDE IN DEXTROSE 5%, DILTIAZEM HYDROCHLORIDE

ELCYS, CYSTEINE HYDROCHLORIDE

ERYTHROMYCIN LACTOBIONATE, ERYTHROMYCIN LACTOBIONATE

GANZYK-RTU, GANCICLOVIR

GLYRX-PF, GLYCOPYRROLATE

MAGNESIUM SULFATE, MAGNESIUM SULFATE

NIPRIDE RTU IN SODIUM CHLORIDE 0.9%, SODIUM NITROPRUSSIDE

POTASSIUM ACETATE, POTASSIUM ACETATE

SODIUM BICARBONATE, SODIUM BICARBONATE

TRANEXAMIC ACID, TRANEXAMIC ACID

VAZCULEP, PHENYLEPHRINE HYDROCHLORIDE

VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

ZINC CHLORIDE, ZINC CHLORIDE

EXELA PHARMA SCIENCE

* EXELA PHARMA SCIENCES

CAFFEINE CITRATE, CAFFEINE CITRATE

NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE, NICARDIPINE HYDROCHLORIDE

NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE

EXELIXIS

* EXELIXIS INC

COMETRIQ, CABOZANTINIB S-MALATE

EXELIXIS INC

* EXELIXIS INC

CABOMETYX, CABOZANTINIB S-MALATE

EXELTIS USA INC

* EXELTIS USA INC

SLYND, DROSPIRENONE

TYBLUME, ETHINYL ESTRADIOL

EYEPOINT PHARMS

* EYEPOINT PHARMACEUTICALS INC

DEXYCU KIT, DEXAMETHASONE

YUTIQ, FLUOCINOLONE ACETONIDE

EYEVANCE

* EYEVANCE PHARMACEUTICALS LLC

FLAREX, FLUOROMETHOLONE ACETATE

NATACYN, NATAMYCIN

TOBRADEX ST, DEXAMETHASONE

ZERVIATE, CETIRIZINE HYDROCHLORIDE

EYWA

* EYWA PHARMA INC

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN

BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN

LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE

METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

POTASSIUM CITRATE, POTASSIUM CITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EYWA PHARMA INC
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
 VALPROIC ACID, VALPROIC ACID

EYWA PHARMA

* EYWA PHARMA PTE LTD
 ACETAZOLAMIDE, ACETAZOLAMIDE
 BACLOFEN, BACLOFEN
 URSODIOL, URSODIOL

LILLY

* ELI LILLY AND CO
 ALIMTA, PEMETREXED DISODIUM
 CIALIS, TADALAFIL
 CYMBALTA, DULOXETINE HYDROCHLORIDE
 EVISTA, RALOXIFENE HYDROCHLORIDE
 FORTEO, TERIPARATIDE
 GLUCAGON, GLUCAGON
 STRATTERA, ATOMOXETINE HYDROCHLORIDE
 SYMBYAX, FLUOXETINE HYDROCHLORIDE
 ZYPREXA ZYDIS, OLANZAPINE
 ZYPREXA, OLANZAPINE

**** F ******FDC LTD**

* FDC LTD
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 LATANOPROST, LATANOPROST
 OFLOXACIN, OFLOXACIN
 TIMOLOL MALEATE, TIMOLOL MALEATE

FDN CONSUMER

* FOUNDATION CONSUMER BRANDS LLC
 ALAVERT, LORATADINE (OTC)
 * FOUNDATION CONSUMER HEALTHCARE LLC
 PLAN B ONE-STEP, LEVONORGESTREL (OTC)

FEINSTEIN

* FEINSTEIN INSTITUTE MEDICAL RESEARCH
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 FLUORODOPA F18, FLUORODOPA F-18

FERA PHARMS

* FERA PHARMACEUTICALS LLC
 TOBRAMYCIN, TOBRAMYCIN

FERA PHARMS LLC

* FERA PHARMACEUTICALS LLC
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE

FERRER INTERNACIONAL

* FERRER INTERNACIONAL SA
 XEPI, OZENOXACIN

FERRING

* FERRING PHARMACEUTICALS INC
 ENDOMETRIN, PROGESTERONE
 FIRMAGON, DEGARELIX ACETATE

FERRING PHARMS INC

* FERRING PHARMACEUTICALS INC
 CERVIDIL, DINOPROSTONE
 CLENPIQ, CITRIC ACID
 DDAVP, DESMOPRESSIN ACETATE
 NOCDURNA, DESMOPRESSIN ACETATE

FLAMINGO PHARMS

* FLAMINGO PHARMACEUTICALS LTD
 METRONIDAZOLE, METRONIDAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FLAMINGO PHARMACEUTICALS LTD
PIROXICAM, PIROXICAM

FLEXION THERAPS INC

* FLEXION THERAPEUTICS INC
ZILRETTA, TRIAMCINOLONE ACETONIDE

FLORIDA

* FLORIDA PHARMACEUTICAL PRODUCTS LLC
LEVETIRACETAM, LEVETIRACETAM
SUMATRIPTAN, SUMATRIPTAN

FOLDRX PHARMS

* FOLDRX PHARMACEUTICALS LLC A WHOLLY OWNED SUB OF PFIZER INC
VYNDAMAX, TAFAMIDIS
VYNDAQEL, TAFAMIDIS MEGLUMINE

FORESEE PHARMS

* FORESEE PHARMACEUTICALS CO LTD
CAMCEVI KIT, LEUPROLIDE MESYLATE

FOSUN PHARMA

* FOSUN PHARMA USA INC
CARISOPRODOL, CARISOPRODOL
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
LEFLUNOMIDE, LEFLUNOMIDE

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 PRILCAINE, LIDOCAINE
METRONIDAZOLE, METRONIDAZOLE
MOMETASONE FUROATE, MOMETASONE FUROATE
MUPIROCIN, MUPIROCIN
NYSTATIN, NYSTATIN
OXISTAT, OXICONAZOLE NITRATE
PREDNICARBATE, PREDNICARBATE
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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FOUGERA PHARMACEUTICALS INC
 FLUOCINONIDE, FLUOCINONIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HYDROCORTISONE, HYDROCORTISONE
 LIDOCAINE, LIDOCAINE
 NITROGLYCERIN, NITROGLYCERIN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 TACROLIMUS, TACROLIMUS
 TERCONAZOLE, TERCONAZOLE

FRESENIUS

* FRESENIUS KABI DEUTSCHLAND GMBH
 INTRALIPID 10%, SOYBEAN OIL
 INTRALIPID 20%, SOYBEAN OIL
 INTRALIPID 30%, SOYBEAN OIL
 * FRESENIUS KABI IPSUM SRL
 NAFCILLIN SODIUM, NAFCILLIN SODIUM

FRESENIUS KABI

* FRESENIUS KABI ANTI INFECTIVES SRL
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 * FRESENIUS KABI AUSTRIA GMBH
 LACTULOSE, LACTULOSE

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
 ATROPINE SULFATE, ATROPINE SULFATE
 AZITHROMYCIN, AZITHROMYCIN
 AZTREONAM, AZTREONAM
 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 IN SODIUM CHLORIDE, CALCIUM GLUCONATE
 CALCIUM GLUCONATE, CALCIUM GLUCONATE
 CARBOPLATIN, CARBOPLATIN
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 CEFOTETAN, CEFOTETAN DISODIUM
 CHLORAMPHENICOL SODIUM SUCCINATE, CHLORAMPHENICOL SODIUM SUCCINATE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 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 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FRESENIUS KABI USA LLC
 DEXTROSE 5% AND SODIUM CHLORIDE 0.225%, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, DEXTROSE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DILAUDID, HYDROMORPHONE HYDROCHLORIDE
 DIMENHYDRINATE, DIMENHYDRINATE
 DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
 DIPRIVAN, PROPOFOL
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXY 100, DOXYCYCLINE HYCLATE
 DOXY 200, DOXYCYCLINE HYCLATE
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ETOPOSIDE, ETOPOSIDE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
 FENTANYL CITRATE, FENTANYL CITRATE
 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
 FOSCARNET SODIUM, FOSCARNET SODIUM
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FULVESTRANT, FULVESTRANT
 FUROSEMIDE, FUROSEMIDE
 GANCICLOVIR SODIUM, 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 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 IFOSFAMIDE, IFOSFAMIDE
 INDOMETHACIN, INDOMETHACIN
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 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
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FRESENIUS KABI USA LLC
 LINEZOLID, LINEZOLID
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, 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
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 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
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OMEGAVEN, FISH OIL TRIGLYCERIDES
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXACILLIN SODIUM, OXACILLIN SODIUM
 OXALIPLATIN, OXALIPLATIN
 OXYTOCIN, OXYTOCIN
 PACLITAXEL, PACLITAXEL
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PENTAM, PENTAMIDINE ISETHIONATE
 PERIKABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 POLOCAINE, MEPIVACAINE HYDROCHLORIDE
 POLOCAINE-MPF, MEPIVACAINE HYDROCHLORIDE
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC
 PROGESTERONE, PROGESTERONE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROTAMINE SULFATE, PROTAMINE SULFATE
 PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
 REMIFENTANIL HYDROCHLORIDE, REMIFENTANIL HYDROCHLORIDE
 RIFAMPIN, RIFAMPIN
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROMIDEPSIN, ROMIDEPSIN
 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
 SODIUM CHLORIDE 14.6%, SODIUM CHLORIDE
 SODIUM CHLORIDE 23.4%, SODIUM CHLORIDE
 SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FRESENIUS KABI USA LLC
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 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
 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
 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
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 OXAZEPAM, OXAZEPAM

**** G ******G1 THERAP**

* G1 THERAPEUTICS INC
 COSELA, TRILACICLIB DIHYDROCHLORIDE

GALDERMA LABS

* GALDERMA LABORATORIES INC
 CLOBEX, CLOBETASOL PROPIONATE
 EPIDUO FORTE, ADAPALENE

GALDERMA LABS LP

* GALDERMA LABORATORIES L P
 CLOBEX, CLOBETASOL PROPIONATE
 * GALDERMA LABORATORIES LP
 AKLIEF, TRIFAROTENE
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GALDERMA LABORATORIES LP
VECTICAL, CALCITRIOL

GALEN SPECIALTY

* GALEN SPECIALTY PHARMA US LLC
SYNERA, LIDOCAINE

GALT PHARMS

* GALT PHARMACEUTICALS LLC
DORAL, QUAZEPAM
ORAVIG, MICONAZOLE
ORPHENGESIC FORTE, ASPIRIN

GE HEALTHCARE

* GE HEALTHCARE
ADREVIEW, IOBENGUANE SULFATE I-123
CERETEC, TECHNETIUM TC-99M EXAMETAZIME KIT
CLARISCAN, GADOTERATE MEGLUMINE
INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE
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
VISIPAQUE 270, IODIXANOL
VISIPAQUE 320, IODIXANOL
VIZAMYL, FLUTEMETAMOL F-18

GE HLTHCARE INC

* GE HEALTHCARE INC
DATSCAN, IOFLUPANE I-123

GENBIOPRO

* GENBIOPRO INC
MIFEPRISTONE, MIFEPRISTONE

GENE YORK PHARMS

* GENE YORK PHARMACEUTICALS GROUP LLC
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

GENENTECH

* GENENTECH INC
ERIVEDGE, VISMODEGIB

GENENTECH INC

* GENENTECH INC
COTELLIC, COBIMETINIB FUMARATE
ESBRIET, PIRFENIDONE
EVRYSDI, RISDIPLAM
GAVRETO, PRALSETINIB
ROZLYTREK, ENTRECTINIB
XOFLUZA, BALOXAVIR MARBOXIL

GENERIC

* GENERICS INTERNATIONAL VENTURES ENTERPRISES LLC
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

GENEYORK PHARMS

* GENEYORK PHARMACEUTICALS GROUP LLC
PREDNISONE, PREDNISONE

GENUS

* GENUS LIFESCIENCES INC
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
HYCODAN, HOMATROPINE METHYLBROMIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
LEVETIRACETAM, LEVETIRACETAM
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GENUS LIFESCIENCES INC
 NYSTATIN, NYSTATIN

GENUS LIFESCIENCES

* GENUS LIFE SCIENCES INC
 GOPRELTO, COCAINE HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

GENZYME

* GENZYME CORP
 CLOLAR, CLOFARABINE
 MOZOBIL, PLERIXAFOR
 RENAGEL, SEVELAMER HYDROCHLORIDE
 RENVELA, SEVELAMER CARBONATE

GENZYME CORP

* GENZYME CORP
 CAPRELSA, VANDETANIB
 CERDELGA, ELIGLUSTAT TARTRATE

GILEAD

* GILEAD SCIENCES INC
 CAYSTON, AZTREONAM
 EMTRIVA, EMTRICITABINE
 HEPSERA, ADEFOVIR DIPIVOXIL
 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
 VEKLURY, REMDESIVIR
 VEMLIDY, TENOFOVIR ALAFENAMIDE FUMARATE
 VIREAD, TENOFOVIR DISOPROXIL FUMARATE
 VOSEVI, SOFOSBUVIR
 ZYDELIG, IDELALISIB

GISKIT

* GISKIT PHARMA BV
 EXEM FOAM KIT, AIR POLYMER-TYPE A

GLAND

* GLAND PHARMA LTD
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 DECITABINE, DECITABINE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 HEPARIN SODIUM, HEPARIN SODIUM
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 MEROPENEM, MEROPENEM
 MESNA, MESNA
 OXALIPLATIN, OXALIPLATIN
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

GLAND PHARMA LTD

* GLAND PHARMA LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GLAND PHARMA LTD
 ADENOSINE, ADENOSINE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BIMATOPROST, BIMATOPROST
 BROMFENAC SODIUM, BROMFENAC SODIUM
 CALCITRIOL, CALCITRIOL
 CARBOPLATIN, CARBOPLATIN
 CISPLATIN, CISPLATIN
 CLOFARABINE, CLOFARABINE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYTARABINE, CYTARABINE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOCETAXEL, DOCETAXEL
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 ERTAPENEM SODIUM, ERTAPENEM SODIUM
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ETOMIDATE, ETOMIDATE
 FLUOROURACIL, FLUOROURACIL
 FOSCARNET SODIUM, FOSCARNET SODIUM
 FUROSEMIDE, FUROSEMIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOFLOXACIN, LEVOFLOXACIN
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 METHOCARBAMOL, METHOCARBAMOL
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 MITOMYCIN, MITOMYCIN
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PACITAXEL, PACITAXEL
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TEMSIROLIMUS, TEMSIROLIMUS
 THIOTEPA, THIOTEPA
 TIROFIBAN HYDROCHLORIDE, TIROFIBAN HYDROCHLORIDE
 TOBRAMYCIN SULFATE (PHARMACY BULK), TOBRAMYCIN SULFATE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TOBRAMYCIN, TOBRAMYCIN
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ******* GLAND PHARMA LTD**

VORICONAZOLE, VORICONAZOLE
 ZIPRASIDONE MESYLATE, ZIPRASIDONE MESYLATE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID
 ZOLEDRONIC, ZOLEDRONIC ACID

GLASSHOUSE PHARMS*** GLASSHOUSE PHARMACEUTICALS LTD CANADA**

CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 HALCINONIDE, HALCINONIDE
 OXCARBAZEPINE, OXCARBAZEPINE

GLAUKOS*** GLAUKOS CORP**

PHOTREXA VISCOUS IN DEXTRAN 20%, RIBOFLAVIN 5'-PHOSPHATE SODIUM
 PHOTREXA, RIBOFLAVIN 5'-PHOSPHATE SODIUM

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
 BREO ELLIPTA, FLUTICASONE FUROATE
 FLOVENT DISKUS 100, FLUTICASONE PROPIONATE
 FLOVENT DISKUS 250, FLUTICASONE PROPIONATE
 FLOVENT DISKUS 50, FLUTICASONE PROPIONATE

GLAXOSMITHKLINE*** GLAXOSMITHKLINE**

ABREVA, DOCOSANOL (OTC)
 BECONASE AQ, BECLOMETHASONE DIPROPIONATE MONOHYDRATE
 EPIVIR-HBV, LAMIVUDINE
 IMITREX STATDOSE, SUMATRIPTAN SUCCINATE
 IMITREX, SUMATRIPTAN
 IMITREX, SUMATRIPTAN SUCCINATE
 MALARONE PEDIATRIC, ATOVAQUONE
 MALARONE, ATOVAQUONE
 NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
 NICORETTE, NICOTINE POLACRILEX (OTC)
 RELENZA, ZANAMIVIR
 VALTREX, VALACYCLOVIR HYDROCHLORIDE
 WELLBUTRIN SR, BUPROPION HYDROCHLORIDE

*** GLAXOSMITHKLINE CONSUMER HEALTHCARE HOLDINGS US LLC**

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 DUAL ACTION WITH ACETAMINOPHEN, ACETAMINOPHEN (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)
 ADVIL, IBUPROFEN SODIUM (OTC)
 AXID AR, NIZATIDINE (OTC)
 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)
 INFANT'S ADVIL, IBUPROFEN (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

- * GLAXOSMITHKLINE CONSUMER HEALTHCARE HOLDINGS US LLC
JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)
LAMISIL, TERBINAFINE HYDROCHLORIDE (OTC)
NICORETTE, NICOTINE POLACRILEX (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 LLC
ZEJULA, NIRAPARIB TOSYLATE

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, TERBINAFINE (OTC)
LAMISIL AT, TERBINAFINE HYDROCHLORIDE (OTC)
NICORETTE, NICOTINE POLACRILEX (OTC)
VOLTAREN ARTHRITIS PAIN, DICLOFENAC SODIUM (OTC)

GLAXOSMITHKLINE LLC

- * GLAXOSMITHKLINE LLC
AMERGE, NARATRIPTAN HYDROCHLORIDE
FLOLAN, EPOPROSTENOL SODIUM
LAMICTAL CD, LAMOTRIGINE
LAMICTAL ODT, LAMOTRIGINE
LAMICTAL XR, LAMOTRIGINE
LAMICTAL, LAMOTRIGINE
MEPRON, ATOVAQUONE
RYTHMOL SR, PROPAFENONE HYDROCHLORIDE

GLENMARK GENERICS

- * GLENMARK GENERICS INC USA
ADAPALENE, ADAPALENE (OTC)
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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

- * GLENMARK GENERICS LTD
 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
 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
 TERIFLUNOMIDE, TERIFLUNOMIDE
- * GLENMARK PHARMACEUTICALS INC USA
 CICLOPIROX, CICLOPIROX
 CLOTRIMAZOLE, CLOTRIMAZOLE
 MUPIROCIN, MUPIROCIN
- * GLENMARK PHARMACEUTICALS LTD
 MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
- * GLENMARK PHARMACEUTICALS SA
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ATOVAQUONE, ATOVAQUONE
 AZELAIC ACID, AZELAIC ACID
 CALCIPOTRIENE, CALCIPOTRIENE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 DESONIDE, DESONIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 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
 FULVESTRANT, FULVESTRANT
 LITHIUM CARBONATE, LITHIUM CARBONATE
 MUPIROCIN, MUPIROCIN CALCIUM
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GLENMARK PHARMACEUTICALS INC USA
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE

GLENMARK PHARMS LTD

* GLENMARK PHARMACEUTICALS LTD
ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CHLORZOXAZONE, CHLORZOXAZONE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DESONIDE, DESONIDE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIMETHYL FUMARATE, DIMETHYL FUMARATE
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
ESTRADIOL, ESTRADIOL
EZETIMIBE AND SIMVASTATIN, EZETIMIBE
EZETIMIBE, EZETIMIBE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FLUCINOLONE ACETONIDE, FLUCINOLONE ACETONIDE
FLUOCINONIDE ACETONIDE, FLUOCINOLONE ACETONIDE
FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
GABAPENTIN, GABAPENTIN
HAILEY FE 1/20, ETHINYL ESTRADIOL
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
ICATIBANT ACETATE, ICATIBANT ACETATE
INDOMETHACIN, INDOMETHACIN
LAMOTRIGINE, LAMOTRIGINE
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LEVONORGESTREL, LEVONORGESTREL (OTC)
LIDOCAINE, LIDOCAINE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
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
PIMECROLIMUS, PIMECROLIMUS
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
RANOLAZINE, RANOLAZINE
RILUZOLE, RILUZOLE
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
RUFINAMIDE, RUFINAMIDE
SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE
SIROLIMUS, SIROLIMUS
TACROLIMUS, TACROLIMUS
TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TELMISARTAN, TELMISARTAN
THEOPHYLLINE, THEOPHYLLINE
TOPIRAMATE, TOPIRAMATE
TRETINOIN, TRETINOIN
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
VORICONAZOLE, VORICONAZOLE

GLENMARK PHARMS SA

* GLENMARK PHARMACEUTICALS SA SWITZERLAND
ACYCLOVIR, ACYCLOVIR
APREPITANT, APREPITANT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

- * GLENMARK PHARMACEUTICALS SA SWITZERLAND
 - ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 - NITROGLYCERIN, NITROGLYCERIN

GLOBAL BLOOD THERAPS

- * GLOBAL BLOOD THERAPEUTICS INC
 - OXBRYTA, VOXELOTOR

GLW

- * GLW PHARMA GMBH
 - OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS

GRANULES

- * GRANULES INDIA LTD
 - ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 - ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
 - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 - GABAPENTIN, GABAPENTIN
 - GUAIFENESIN, GUAIFENESIN (OTC)
 - IBUPROFEN, IBUPROFEN
 - IBUPROFEN, IBUPROFEN (OTC)
 - LORATADINE, LORATADINE (OTC)
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - METHOCARBAMOL, METHOCARBAMOL
 - NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 - NAPROXEN, NAPROXEN
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 - ZONISAMIDE, ZONISAMIDE
- * GRANULES PHARMACEUTICALS INC
 - AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 - BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 - BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 - COLCHICINE, COLCHICINE
 - DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 - DOFETILIDE, DOFETILIDE
 - METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 - PENICILLAMINE, PENICILLAMINE
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 - PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 - RAMELTEON, RAMELTEON
 - TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 - VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 - VIGABATRIN, VIGABATRIN

GRANULES INDIA

- * GRANULES INDIA LTD
 - IBUPROFEN, IBUPROFEN (OTC)
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

GRAVITI PHARMS

- * GRAVITI PHARMACEUTICALS INC
 - GABAPENTIN, GABAPENTIN
- * GRAVITI PHARMACEUTICALS PRIVATE LTD
 - ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 - FENOFIBRATE, FENOFIBRATE
 - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 - RANOLAZINE, RANOLAZINE
 - SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE

GUARDIAN DRUG

- * GUARDIAN DRUG CO
 - GUAIFENESIN, GUAIFENESIN (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GUARDIAN DRUG CO
 IBUPROFEN, IBUPROFEN (OTC)
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE

GUERBET

* GUERBET LLC
 DOTAREM, GADOTERATE MEGLUMINE
 LIPIODOL, ETHIODIZED OIL

GW RES LTD

* GW RESEARCH LTD
 EPIDIOLEX, CANNABIDIOL

**** H ******HAEMONETICS**

* HAEMONETICS CORP
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

HAINAN POLY

* HAINAN POLY PHARM CO LTD
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

HAINAN POLY PHARM

* HAINAN POLY PHARMACEUTICAL CO LTD
 AZITHROMYCIN, AZITHROMYCIN
 BIVALIRUDIN, BIVALIRUDIN
 EPTIFIBATIDE, EPTIFIBATIDE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 LEVETIRACETAM, LEVETIRACETAM
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE

HALOCARBON PRODS

* HALOCARBON PRODUCTS CORP
 ISOFLURANE, ISOFLURANE
 SEVOFLURANE, SEVOFLURANE

HANDA

* HANDA NEUROSCIENCE LLC
 TASCENSO ODT, FINGOLIMOD LAURYL SULFATE

HANGZHOU BINJIANG

* HANGZHOU MINSHENG BINJIANG PHARMACEUTICAL CO LTD
 ALENDRONATE SODIUM, ALENDRONATE SODIUM

HANGZHOU ZHONGMEI

* HANGZHOU ZHONGMEI HUADONG PHARMACEUTICAL CO LTD
 DAPTOMYCIN, DAPTOMYCIN

HARMAN FINOCHEM

* HARMAN FINOCHEM LTD
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

HARMONY

* HARMONY BIOSCIENCES LLC
 WAKIX, PITOLISANT HYDROCHLORIDE

HBT LABS INC

* HBT LABS INC
 FULVESTRANT, FULVESTRANT

HEC PHARM

* HEC PHARM USA INC
 CLARITHROMYCIN, CLARITHROMYCIN
 LEVOFLOXACIN, LEVOFLOXACIN
 OLANZAPINE, OLANZAPINE
 PRASUGREL, PRASUGREL HYDROCHLORIDE

HEC PHARM CO LTD

* HEC PHARM CO LTD
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE

HELSINN

* HELSINN BIREX PHARMACEUTICALS LTD
 VALCHLOR, MECHLORETHAMINE HYDROCHLORIDE

HELSINN HLTHCARE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HELSINN HEALTHCARE SA
 AKYNZEO, FOSNETUPITANT CHLORIDE HYDROCHLORIDE
 AKYNZEO, NETUPITANT
 TRUSELTIQ, INFIGRATINIB PHOSPHATE

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
 BUMETANIDE, BUMETANIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DIFLUNISAL, DIFLUNISAL
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 METHIMAZOLE, METHIMAZOLE
 NADOLOL, NADOLOL
 NIFEDIPINE, NIFEDIPINE

HERITAGE PHARMA AVET

* HERITAGE PHARMA LABS INC DBA AVET PHARMACEUTICALS LABS INC
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 THEOPHYLLINE, THEOPHYLLINE

HERITAGE PHARMS INC

* HERITAGE PHARMACEUTICALS INC
 ACYCLOVIR, ACYCLOVIR
 CALCIUM ACETATE, CALCIUM ACETATE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 FELODIPINE, FELODIPINE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GLYBURIDE, GLYBURIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LEFLUNOMIDE, LEFLUNOMIDE
 METRONIDAZOLE, METRONIDAZOLE
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NIMODIPINE, NIMODIPINE
 NYSTATIN, NYSTATIN
 PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
 TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

HERON THERAPS INC

* HERON THERAPEUTICS INC
 CINVANTI, APREPITANT
 SUSTOL, GRANISETRON
 ZYNRELEF KIT, BUPIVACAINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******HETERO LABS LTD**

* HETERO LABS LTD
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE

HETERO LABS LTD III

* HETERO LABS LTD UNIT III
ABACAVIR SULFATE, ABACAVIR SULFATE
ATOVAQUONE, ATOVAQUONE
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLOBAZAM, CLOBAZAM
DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
DIMETHYL FUMARATE, DIMETHYL FUMARATE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
EFAVIRENZ, EFAVIRENZ
EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
FENOFIBRATE, FENOFIBRATE
FINASTERIDE, FINASTERIDE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
INDOMETHACIN, INDOMETHACIN
LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
LANSOPRAZOLE, LANSOPRAZOLE
LEVETIRACETAM, LEVETIRACETAM
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
LEVOCETIRIZINE HYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
LITHIUM CARBONATE, LITHIUM CARBONATE
LOPINAVIR AND RITONAVIR, LOPINAVIR
LORATADINE, LORATADINE (OTC)
METHOCARBAMOL, METHOCARBAMOL
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
NEVIRAPINE, NEVIRAPINE
OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
OMEPRAZOLE, OMEPRAZOLE
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
PREGABALIN, PREGABALIN
RITONAVIR, RITONAVIR
ROFLUMILAST, ROFLUMILAST
RUFINAMIDE, RUFINAMIDE
SIMVASTATIN, SIMVASTATIN
TADALAFIL, TADALAFIL
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
ARIPIRAZOLE, ARIPIRAZOLE
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
COLCHICINE, COLCHICINE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
ENTECAVIR, ENTECAVIR
FAMCICLOVIR, FAMCICLOVIR
FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
IRBESARTAN, IRBESARTAN
LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
LAMIVUDINE, LAMIVUDINE
LEVOFLOXACIN, LEVOFLOXACIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HETERO LABS LTD UNIT V
 LINEZOLID, LINEZOLID
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 TELMISARTAN, TELMISARTAN
 TETRABENAZINE, TETRABENAZINE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TOLVAPTAN, TOLVAPTAN
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALSARTAN, VALSARTAN

HEYL CHEMISCH

* HEYL CHEMISCH PHARMAZEUTISCHE FABRIK
 RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE (II)

HIGH TECH PHARMA

* HIGH TECHNOLOGY PHARMACAL CO INC
 VALPROIC ACID, VALPROIC ACID

HIKAL

* HIKAL LTD
 PREGABALIN, PREGABALIN

HIKMA

* HIKMA FARMACEUTICA LDA
 CEFOTAXIME, CEFOTAXIME SODIUM

* HIKMA FARMACEUTICA PORTUGAL SA
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 CEFTRIAZONE SODIUM, CEFTRIAZONE SODIUM
 DOCETAXEL, DOCETAXEL
 ESTRADIOL VALERATE, ESTRADIOL VALERATE
 ETOMIDATE, ETOMIDATE
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE

* HIKMA PHARMACEUTICALS
 AMOXICILLIN, AMOXICILLIN
 CEFACLOR, CEFACLOR
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEPHALEXIN, CEPHALEXIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 GLYBURIDE (MICRONIZED), GLYBURIDE

* HIKMA PHARMACEUTICALS INTERNATIONAL LTD
 CODEINE SULFATE, CODEINE SULFATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIGOXIN, DIGOXIN
 FUROSEMIDE, FUROSEMIDE
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 RUFINAMIDE, RUFINAMIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE

* HIKMA PHARMACEUTICALS LLC
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DOXERCALCIFEROL, DOXERCALCIFEROL
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PIROXICAM, PIROXICAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA PHARMACEUTICALS USA INC
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
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
AMRINONE LACTATE, INAMRINONE LACTATE
ATIVAN, LORAZEPAM
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
ATROPINE SULFATE, ATROPINE SULFATE
AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
BEXAROTENE, BEXAROTENE
BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
BOSENTAN, BOSENTAN
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
CAFCIT, 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
CILOSTAZOL, CILOSTAZOL
CISPLATIN, CISPLATIN
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLADRIBINE, CLADRIBINE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLOBAZAM, CLOBAZAM
CLOTRIMAZOLE, CLOTRIMAZOLE
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
CYCLOSPORINE, CYCLOSPORINE
CYTARABINE, CYTARABINE
DACARBAZINE, DACARBAZINE
DALFAMPRIDINE, DALFAMPRIDINE
DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
DEFERIPRONE, DEFERIPRONE
DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
DEXAMETHASONE INTENSOL, DEXAMETHASONE
DEXAMETHASONE, DEXAMETHASONE
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
DIAZEPAM INTENSOL, DIAZEPAM
DIAZEPAM, DIAZEPAM
DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE), DICYCLOMINE HYDROCHLORIDE
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
DIPYRIDAMOLE, DIPYRIDAMOLE
DISULFIRAM, DISULFIRAM
DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
DOPRAM, DOXAPRAM HYDROCHLORIDE
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA PHARMACEUTICALS USA INC
DOXYCYCLINE, DOXYCYCLINE HYCLATE
DROXIDOPA, DROXIDOPA
DURAMORPH PF, MORPHINE SULFATE
EPHEDRINE SULFATE, EPHEDRINE SULFATE
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
ETHACRYNIC ACID, ETHACRYNIC ACID
ETOMIDATE, ETOMIDATE
ETOPOSIDE, ETOPOSIDE
EVEROLIMUS, EVEROLIMUS
EXEMESTANE, EXEMESTANE
FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE
FEBUXOSTAT, FEBUXOSTAT
FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
FLECAINIDE ACETATE, FLECAINIDE ACETATE
FLOXURIDINE, FLOXURIDINE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
FLUCYTOSINE, FLUCYTOSINE
FLUMAZENIL, FLUMAZENIL
FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
FLUTICASONE PROPIONATE AND SALMETEROL XINAFOATE, FLUTICASONE PROPIONATE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
FUROSEMIDE, FUROSEMIDE
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
HALOPERIDOL, HALOPERIDOL LACTATE
HEPARIN SODIUM, HEPARIN SODIUM
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
ICOSAPENT ETHYL, ICOSAPENT ETHYL
IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
IFOSFAMIDE, IFOSFAMIDE
IMATINIB MESYLATE, IMATINIB MESYLATE
IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
INFUMORPH, MORPHINE SULFATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
KLOXXADO, NALOXONE HYDROCHLORIDE
LABETALOL HYDROCHLORIDE IN DEXTROSE, LABETALOL HYDROCHLORIDE
LABETALOL HYDROCHLORIDE IN SODIUM CHLORIDE, LABETALOL HYDROCHLORIDE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
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, LOSARTAN POTASSIUM
MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
MERCAPTOPYRINE, MERCAPTOPYRINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA PHARMACEUTICALS USA INC
 MESNA, MESNA
 METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 MITOMYCIN, MITOMYCIN
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NALOXONE, NALOXONE HYDROCHLORIDE
 NAPROXEN, NAPROXEN
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 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
 PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE
 PREDNISONE INTENSOL, PREDNISONE
 PREDNISONE, PREDNISONE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROPOFOL, PROPOFOL
 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
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 RUFINAMIDE, RUFINAMIDE
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUFENTANIL CITRATE, SUFENTANIL CITRATE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TETRABENAZINE, TETRABENAZINE
 TINIDAZOLE, TINIDAZOLE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TORSEMIDE, TORSEMIDE
 TRIAZOLAM, TRIAZOLAM
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VINBLASTINE SULFATE, VINBLASTINE SULFATE
 VINOELBINE TARTRATE, VINOELBINE TARTRATE
 ZALEPLON, ZALEPLON

HIKMA FARMACEUTICA

* HIKMA FARMACEUTICA (PORTUGAL) SA
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CEFOXITIN, CEFOXITIN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

- * HIKMA FARMACEUTICA (PORTUGAL) SA
 - CEFTRIAZONE, CEFTRIAZONE SODIUM
 - CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 - CIPROFLOXACIN, CIPROFLOXACIN
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - ENALAPRILAT, ENALAPRILAT
 - 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, 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
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 - OXYTOCIN, OXYTOCIN
 - TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
- * HIKMA FARMACEUTICA SA
 - ZOLEDRONIC ACID, ZOLEDRONIC ACID

HIKMA INTL PHARMS

- * HIKMA INTERNATIONAL PHARMACEUTICALS LLC
 - CAPTOPRIL, CAPTOPRIL
 - CORTISONE ACETATE, CORTISONE ACETATE
 - DIGOXIN, DIGOXIN
 - DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
 - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 - ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 - MITIGARE, COLCHICINE

HIKMA PHARM CO LTD

- * HIKMA PHARM CO LTD
 - ARGATROBAN, ARGATROBAN

HIKMA PHARMS

- * HIKMA PHARMACEUTICALS
 - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 - AMOXICILLIN, AMOXICILLIN
 - BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 - BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 - CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 - DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 - DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 - PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 - RIFAMPIN, RIFAMPIN
 - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
- * HIKMA PHARMACEUTICALS CO LTD
 - PARICALCITOL, PARICALCITOL

HILL DERMAC

- * HILL DERMACEUTICALS INC
 - DERMA-SMOOTH/FS, FLUOCINOLONE ACETONIDE
 - DERMOTIC, FLUOCINOLONE ACETONIDE

HILL DERMACEUTICALS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HILL DERMACEUTICALS INC
TOLAK, FLUOROURACIL

HISAMITSU

* HISAMITSU PHARMACEUTICAL CO INC
SECUADO, ASENAPINE

HISAMITSU PHARM CO

* HISAMITSU PHARMACEUTICAL CO INC
SALONPAS, MENTHOL (OTC)

HISUN PHARM HANGZHOU

* HISUN PHARMACEUTICAL (HANGZHOU) CO LTD
CLADRIBINE, CLADRIBINE
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IRBESARTAN, IRBESARTAN
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM
SIMVASTATIN, SIMVASTATIN

* HISUN PHARMACEUTICAL HANGZHOU CO LTD
DACTINOMYCIN, DACTINOMYCIN
DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
TICAGRELOR, TICAGRELOR

HLTHCARE

* HEALTHCARE PHARMACEUTICALS LTD
ATENOLOL, ATENOLOL
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE

HOFFMANN LA ROCHE

* HOFFMANN LA ROCHE INC
VALCYTE, VALGANCICLOVIR HYDROCHLORIDE
XELODA, CAPECITABINE
ZELBORAF, VEMURAFENIB

HOFFMANN-LA ROCHE

* HOFFMANN-LA ROCHE INC
ALECENSA, ALECTINIB HYDROCHLORIDE

HONG KONG

* HONG KONG KING FRIEND INDUSTRIAL CO LTD
BIVALIRUDIN, BIVALIRUDIN
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
DOCETAXEL, DOCETAXEL
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE

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 THERAPEUTICS IRELAND DAC
PENNSAID, DICLOFENAC SODIUM

* HORIZON THERAPEUTICS USA INC
PROCYSBI, CYSTEAMINE BITARTRATE
RAYOS, PREDNISONE

HORIZON PHARMA USA

* HORIZON PHARMA USA INC
PROCYSBI, CYSTEAMINE BITARTRATE

HORIZON THERAP

* HORIZON THERAPEUTICS LLC
BUPHENYL, SODIUM PHENYL BUTYRATE
RAVICTI, GLYCEROL PHENYL BUTYRATE

HOSPIRA

* HOSPIRA INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HOSPIRA INC

ACETYLCYSTEINE, ACETYLCYSTEINE
 ALFENTANIL, ALFENTANIL HYDROCHLORIDE
 AMIDATE, ETOMIDATE
 AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID
 AMINOPHYLLINE, AMINOPHYLLINE
 AMMONIUM CHLORIDE IN PLASTIC CONTAINER, AMMONIUM CHLORIDE
 ATROPINE SULFATE, 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, 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
 CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 CORLOPAM, FENOLDOPAM MESYLATE
 CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE
 CYTARABINE, CYTARABINE
 DACARBAZINE, DACARBAZINE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEMEROL, MEPERIDINE HYDROCHLORIDE
 DEXTROSE 25%, DEXTROSE
 DEXTROSE 5% 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
 ERYTHROCIN, ERYTHROMYCIN LACTOBIONATE
 FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
 FENTANYL CITRATE, FENTANYL CITRATE
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HOSPIRA INC

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
 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 CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 14.6%, SODIUM CHLORIDE
 SODIUM CHLORIDE 23.4%, SODIUM CHLORIDE
 SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 SUFENTANIL CITRATE, SUFENTANIL CITRATE
 TAZICEF, CEFTAZIDIME
 THAM, TROMETHAMINE
 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
 VITAMIN K1, PHYTONADIONE
 ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE

* HOSPIRA WORLDWIDE, INC

GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 NITROPRESS, SODIUM NITROPRUSSIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HOSPIRA WORLDWIDE, INC
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE

HOSPIRA INC

* HOSPIRA INC
 ADENOSINE, ADENOSINE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ARGATROBAN, ARGATROBAN
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 AZTREONAM, AZTREONAM
 BIVALIRUDIN, BIVALIRUDIN
 BUSULFAN, BUSULFAN
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 DAPTOMYCIN, DAPTOMYCIN
 DOCETAXEL, DOCETAXEL
 DOXERCALCIFEROL, DOXERCALCIFEROL
 EPINEPHRINE, EPINEPHRINE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 HEPARIN SODIUM, HEPARIN SODIUM
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 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
 MILRINONE LACTATE, MILRINONE LACTATE
 MORPHINE SULFATE, MORPHINE SULFATE
 NIPENT, PENTOSTATIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PARICALCITOL, PARICALCITOL
 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

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
 IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
 LINEZOLID, LINEZOLID
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MEROPENEM, MEROPENEM

HQ SPECIALITY PHARMA

* HQ SPECIALITY PHARMA LLC
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM

HQ SPECLT PHARMA

* HQ SPECIALTY PHARMA CORP
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******HRA PHARMA**

- * HRA PHARMA RARE DISEASES
LYSODREN, MITOTANE
METOPIRONE, METYRAPONE

HUMANWELL PURACAP

- * HUMANWELL PURACAP PHARMACEUTICAL WUHAN CO LTD
DUTASTERIDE, DUTASTERIDE
IBUPROFEN, IBUPROFEN (OTC)

HUONS

- * HUONS CO LTD
BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE

ROCHE

- * HOFFMANN LA ROCHE INC
FUZEON, ENFUVIRTIDE
TAMIFLU, OSELTAMIVIR PHOSPHATE
VALIUM, DIAZEPAM

SHUANGCHENG

- * HAINAN SHUANGCHENG PHARMACEUTICALS CO LTD
BIVALIRUDIN, BIVALIRUDIN
EPTIFIBATIDE, EPTIFIBATIDE
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
PREGABALIN, PREGABALIN

**** I ******I3 PHARMS**

- * I3 PHARMACEUTICALS LLC
CHLORZOXAZONE, CHLORZOXAZONE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
RAMELTEON, RAMELTEON
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE

IBSA INST BIO

- * IBSA INSTITUT BIOCHIMIQUE SA
LICART, DICLOFENAC EPOLAMINE

ICHNOS

- * ICHNOS SCIENCES SA
DEFERASIROX, DEFERASIROX

ICU MEDICAL INC

- * ICU MEDICAL INC
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
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
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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******* ICU MEDICAL INC**

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 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 0.45% 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 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 0.45% 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 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM
 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

IMPACT

* IMPACT BIOMEDICINES INC A WHOLLY OWNED SUB OF CELGENE CORP
 INREBIC, FEDRATINIB HYDROCHLORIDE

IMPAX

* IMPAX LABORATORIES LLC
 ADRENALICK, EPINEPHRINE
 BACLOFEN, BACLOFEN
 SEVELAMER CARBONATE, SEVELAMER CARBONATE

IMPAX LABS

* IMPAX LABORATORIES INC
 ACARBOSE, ACARBOSE
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 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
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 OMEPRAZOLE, OMEPRAZOLE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE 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
 BUDESONIDE, BUDESONIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DEXEDRINE, DEXTROAMPHETAMINE SULFATE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******* IMPAX LABORATORIES INC**

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
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RYTARY, CARBIDOPA

IMPAX PHARMS*** IMPAX PHARMACEUTICALS**

GEMFIBROZIL, GEMFIBROZIL
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE

IMPEL NEUROPHARMA*** IMPEL NEUROPHARMA**

TRUDHESA, DIHYDROERGOTAMINE MESYLATE

INCYTE CORP*** INCYTE CORP**

JAKAFI, RUXOLITINIB PHOSPHATE
 OPZELURA, RUXOLITINIB PHOSPHATE
 PEMAZYRE, PEMIGATINIB

INDCHEMIE HEALTH*** INDCHEMIE HEALTH SPECIALTIES PVT LTD**

ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE

INDICUS PHARMA*** INDICUS PHARMA LLC**

ACETAZOLAMIDE, ACETAZOLAMIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 LETROZOLE, LETROZOLE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE

INDIVIOR INC*** INDIVIOR INC**

BUPRENEX, BUPRENORPHINE HYDROCHLORIDE
 PERSERIS KIT, RISPERIDONE
 SUBLOCADE, BUPRENORPHINE
 SUBOXONE, BUPRENORPHINE HYDROCHLORIDE

INDOCO*** INDOCO REMEDIES LTD**

ALLOPURINOL, ALLOPURINOL
 APIXABAN, APIXABAN
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 FEBUXOSTAT, FEBUXOSTAT
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 OLANZAPINE, OLANZAPINE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE

INDOCO REMEDIES*** INDOCO REMEDIES LTD**

GLIMEPIRIDE, GLIMEPIRIDE

INFORLIFE*** INFORLIFE SA**

CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******* INFORLIFE SA**

LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 MIDAZOLAM IN 0.9% SODIUM CHLORIDE, MIDAZOLAM
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

INGENUS PHARMS LLC*** INGENUS PHARMACEUTICALS LLC**

ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 CABERGOLINE, CABERGOLINE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DECITABINE, DECITABINE
 DOCETAXEL, DOCETAXEL
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

INGENUS PHARMS NJ*** INGENUS PHARMACEUTICALS NJ LLC**

CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN

INNOCOLL PHARMS*** INNOCOLL PHARMACEUTICALS**

XARACOLL, BUPIVACAINE HYDROCHLORIDE

INNOGENIX*** INNOGENIX LLC**

BACLOFEN, BACLOFEN
 HALOPERIDOL, HALOPERIDOL
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 METOLAZONE, METOLAZONE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

INNOPHARMA*** INNOPHARMA LICENSING LLC A SUB OF PFIZER INC**

PROPOFOL, PROPOFOL

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
 TIROSINT-SOL, LEVOTHYROXINE SODIUM

INTAS PHARMS USA*** INTAS PHARMACEUTICALS USA**

IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE

INTELLIPHARMACEUTICS*** INTELLIPHARMACEUTICS CORP**

DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

INTERCEPT PHARMS INC*** INTERCEPT PHARMACEUTICALS INC**

OICALIVA, OBETICHOLIC ACID

INTERSECT ENT INC*** INTERSECT ENT INC**

SINUVA, MOMETASONE FUROATE

INTL ISOTOPES*** INTERNATIONAL ISOTOPES INC**

SODIUM IODIDE I 131, SODIUM IODIDE I-131

INTL MEDICATION*** INTERNATIONAL MEDICATION SYSTEM**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******* INTERNATIONAL MEDICATION SYSTEM**

LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 PHYTONADIONE, PHYTONADIONE
 PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE

INTL MEDICATION SYS*** INTERNATIONAL MEDICATION SYSTEMS LTD**

ATROPINE SULFATE, ATROPINE SULFATE
 CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 DEXTROSE 50%, DEXTROSE
 EPINEPHRINE, EPINEPHRINE
 LORAZEPAM, LORAZEPAM
 SODIUM BICARBONATE, SODIUM BICARBONATE

INTRA-CELLULAR*** INTRA-CELLULAR THERAPIES INC**

CAPLYTA, LUMATEPERONE TOSYLATE

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
 LANREOTIDE ACETATE, LANREOTIDE ACETATE
 LEVETIRACETAM, LEVETIRACETAM
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINOPRIL, LISINOPRIL
 MEPROBAMATE, MEPROBAMATE
 NABUMETONE, NABUMETONE
 NADOLOL, NADOLOL
 NAPROXEN, NAPROXEN
 OLANZAPINE, OLANZAPINE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 PENICILLAMINE, PENICILLAMINE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 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
 TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE
 VIGABATRIN, VIGABATRIN
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM
 ZOLMITRIPTAN, ZOLMITRIPTAN
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

* INVAGEN PHARMACEUTICALS INC
ZONISAMIDE, ZONISAMIDE

INVATECH

* INVATECH PHARMA SOLUTIONS LLC
DIVALPROEX SODIUM, DIVALPROEX SODIUM
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE

INVENTIA

* INVENTIA HEALTHCARE LTD
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
LANSOPRAZOLE, LANSOPRAZOLE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
PALIPERIDONE, PALIPERIDONE
TELMISARTAN, TELMISARTAN

INVENTIA HLTHCARE

* INVENTIA HEALTHCARE PRIVATE LTD
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

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
ONIVYDE, IRINOTECAN HYDROCHLORIDE

IPSEN PHARMA

* IPSEN PHARMA BIOTECH SAS
SOMATULINE DEPOT, LANREOTIDE ACETATE

IRONSHORE PHARMS

* IRONSHORE PHARMACEUTICALS AND DEVELOPMENT INC
JORNAY PM, METHYLPHENIDATE HYDROCHLORIDE

ISOLOGIC INNOVATIVE

* ISOLOGIC INNOVATIVE RADIOPHARMACEUTICALS LTD
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

* ISTITUTO BIOCHIMICO ITALIANO SPA
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PIPERACILLIN, PIPERACILLIN SODIUM

ITALFARMACO SPA

* ITALFARMACO SPA
 TIGLUTIK KIT, RILUZOLE

IVAX PHARMS

* IVAX PHARMACEUTICALS INC
 VALSARTAN, VALSARTAN

IVAX SUB TEVA PHARMS

* IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 BACLOFEN, BACLOFEN
 CABERGOLINE, CABERGOLINE
 CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CIMETIDINE, CIMETIDINE (OTC)
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLOZAPINE, CLOZAPINE
 CYCLOSPORINE, CYCLOSPORINE
 DIAZEPAM, DIAZEPAM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FAMOTIDINE, FAMOTIDINE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

**** J ******J AND J CONSUMER INC**

* JOHNSON AND JOHNSON CONSUMER INC MCNEIL CONSUMER HEALTHCARE DIV
 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)
 RHINOCORT ALLERGY, BUDESONIDE (OTC)
 SINE-AID IB, IBUPROFEN (OTC)
 SUDAFED 24 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 TYLENOL, ACETAMINOPHEN (OTC)
 ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 ZYRTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE (OTC)

JACOBUS

* JACOBUS PHARMACEUTICAL CO
 DAPSONE, DAPSONE
 PASER, AMINOSALICYLIC ACID

JACOBUS PHARM CO INC

* JACOBUS PHARMACEUTICAL CO INC
 RUZURGI, AMIFAMPRIDINE

JANSSEN BIOTECH

* JANSSEN BIOTECH INC
 BALVERSA, ERDAFITINIB
 ERLEADA, APALUTAMIDE
 ZYTIGA, ABIRATERONE ACETATE

JANSSEN PHARMS

* JANSSEN PHARMACEUTICALS INC
 CONCERTA, METHYLPHENIDATE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

* JANSSEN PHARMACEUTICALS INC
 DITROPAN XL, OXYBUTYNIN CHLORIDE
 ELMIRON, PENTOSAN POLYSULFATE SODIUM
 HALDOL, HALOPERIDOL DECANOATE
 INVEGA HAFYERA, PALIPERIDONE PALMITATE
 INVEGA SUSTENNA, PALIPERIDONE PALMITATE
 INVEGA TRINZA, PALIPERIDONE PALMITATE
 INVEGA, PALIPERIDONE
 INVOKAMET XR, CANAGLIFLOZIN
 INVOKAMET, CANAGLIFLOZIN
 INVOKANA, CANAGLIFLOZIN
 PONVORY, PONESIMOD
 RAZADYNE ER, GALANTAMINE HYDROBROMIDE
 RISPERDAL CONSTA, RISPERIDONE
 RISPERDAL, RISPERIDONE
 SPORANOX, ITRACONAZOLE
 SPRAVATO, ESKETAMINE HYDROCHLORIDE
 TOPAMAX, TOPIRAMATE
 ULTRACET, ACETAMINOPHEN
 ULTRAM, TRAMADOL HYDROCHLORIDE
 XARELTO, RIVAROXABAN

JANSSEN PRODS

* JANSSEN PRODUCTS LP
 EDURANT, RILPIVIRINE HYDROCHLORIDE
 PREZCOBIX, COBICISTAT
 PREZISTA, DARUNAVIR
 SYMTUZA, COBICISTAT
 YONDELIS, TRABECTEDIN

JANSSEN R AND D

* JANSSEN RESEARCH AND DEVELOPMENT LLC
 INTELENCE, ETRAVIRINE

JANSSEN THERAP

* JANSSEN THERAPEUTICS DIV JANSSEN PRODUCTS LP
 SIRTURO, BEDAQUILINE FUMARATE

JAZZ

* JAZZ PHARMACEUTICALS IRELAND LTD
 SUNOSI, SOLRIAMFETOL HYDROCHLORIDE
 XYWAV, CALCIUM OXYBATE
 ZEPZELCA, LURBINECTEDIN

JAZZ PHARMS

* JAZZ PHARMACEUTICALS INC
 XYREM, SODIUM OXYBATE

JAZZ PHARMS INC

* JAZZ PHARMACEUTICALS INC
 DEFITELIO, DEFIBROTIDE SODIUM

JDP

* JDP THERAPEUTICS LLC
 QUZYTIR, CETIRIZINE HYDROCHLORIDE

JIANGSU HANSOH PHARM

* JIANGSU HANSOH PHARMACEUTICAL GROUP CO LTD
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 MICAfungin SODIUM, MICAfungin SODIUM
 VINORELBINE TARTRATE, VINORELBINE TARTRATE

JIANGSU PHARMS

* JIANGSU HENGRUI PHARMACEUTICALS CO LTD
 CARMUSTINE, CARMUSTINE
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DAPTOMYCIN, DAPTOMYCIN
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOCETAXEL, DOCETAXEL
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

* JIANGSU HENGRUI PHARMACEUTICALS CO LTD
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 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)

JOURNEY

* JOURNEY MEDICAL CORP
 EURAX, CROTAMITON
 EXELDERM, SULCONAZOLE NITRATE
 QBREXZA, GLYCOPYRRONIUM TOSYLATE
 XIMINO, MINOCYCLINE HYDROCHLORIDE

JUBILANT

* JUBILANT DRAXIMAGE INC DBA JUBILANT RADIOPHARMA
 DRAX EXAMETAZIME, TECHNETIUM TC-99M EXAMETAZIME KIT
 DRAXIMAGE DTPA, TECHNETIUM TC-99M PENTETATE KIT
 DRAXIMAGE MDP-25, TECHNETIUM TC-99M MEDRONATE
 HICON, SODIUM IODIDE I-131
 RUBY-FILL, RUBIDIUM CHLORIDE RB-82
 SODIUM IODIDE I 131, SODIUM IODIDE I-131

JUBILANT CADISTA

* JUBILANT CADISTA PHARMACEUTICALS INC
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LAMOTRIGINE, LAMOTRIGINE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 PREDNISONE, PREDNISONE
 PROCOMP, PROCHLORPERAZINE MALEATE
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

JUBILANT DRAXIMAGE

* 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
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 IRBESARTAN, IRBESARTAN
 ITRACONAZOLE, ITRACONAZOLE
 OLANZAPINE, OLANZAPINE
 RISPERIDONE, RISPERIDONE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SPIRONOLACTONE, SPIRONOLACTONE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

* JUBILANT GENERICS LTD
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALSARTAN, VALSARTAN
 ZOLMITRIPTAN, ZOLMITRIPTAN

STEVENS J

* JEROME STEVENS PHARMACEUTICALS INC
 BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
 DIGOXIN, DIGOXIN
 UNITHROID, LEVOTHYROXINE SODIUM **

**** K ******GRIFFEN**

* KW GRIFFEN CO
 BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)

KADMON PHARMS LLC

* KADMON PHARMACEUTICALS LLC
 REZUROCK, BELUMOSUDIL MESYLATE

KAI PHARMS INC

* KAI PHARMACEUTICALS INC A WHOLLY OWNED SUBSIDIARY OF AMGEN INC
 PARSABIV, ETELCALCETIDE

KALA PHARMS INC

* KALA PHARMACEUTICALS INC
 EYSUVIS, LOTEPREDNOL ETABONATE
 INVELTYS, LOTEPREDNOL ETABONATE

KALEO INC

* KALEO INC
 AUVI-Q, EPINEPHRINE

KARTHA

* KARTHA PHARMACEUTICALS INC
 BACLOFEN, BACLOFEN

KARYOPHARM THERAPS

* KARYOPHARM THERAPEUTICS INC
 XPOVIO, SELINEXOR

KENTON

* KENTON CHEMICALS AND PHARMACEUTICALS CORP
 ACYCLOVIR, ACYCLOVIR
 ANASTROZOLE, ANASTROZOLE
 BICALUTAMIDE, BICALUTAMIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 RILUZOLE, RILUZOLE

KERYX BIOPHARMS

* KERYX BIOPHARMACEUTICALS INC
 AURYXIA, FERRIC CITRATE

KETTERING MEDCTR

* KETTERING MEDCTR
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

KINDEVA

* KINDEVA DRUG DELIVERY LP
 FENTANYL-100, FENTANYL
 FENTANYL-12, FENTANYL
 FENTANYL-25, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-75, FENTANYL
 PROVENTIL-HFA, ALBUTEROL SULFATE

KING PHARMS

* KING PHARMACEUTICALS INC
 SYNERCID, DALFOPRISTIN
 * KING PHARMACEUTICALS RESEARCH AND DEVELOPMENT LLC
 CYTOMEL, LIOthyronine SODIUM
 LEVOXYL, LEVOTHYROXINE SODIUM **
 * KING PHARMACEUTICALS RESEARCH AND DEVELOPMENT LLC A SUB OF PFIZER INC
 SKELAXIN, METAXALONE

KING PHARMS LLC

* KING PHARMACEUTICALS LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** K ****

* 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
 PENICILLIN G PROCAINE, PENICILLIN G PROCAINE
 SILVADENE, SILVER SULFADIAZINE

KNIGHT THERAPS

* KNIGHT THERAPEUTICS USA INC
 IMPAVIDO, MILTEFOSINE

KOWA CO

* KOWA CO LTD
 LIVALO, PITAVASTATIN CALCIUM

KOWA PHARMS

* KOWA PHARMACEUTICALS AMERICA INC
 SEGLENTIS, CELECOXIB

KRAMER

* KRAMER LABORATORIES INC
 NIZORAL ANTI-DANDRUFF, KETOCONAZOLE (OTC)

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

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
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

KVK TECH INC

* KVK TECH INC
 APADAZ, ACETAMINOPHEN
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

KYOWA KIRIN

* KYOWA KIRIN INC
 FARESTON, TOREMIFENE CITRATE
 NOURIANZ, ISTRADefylline
 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)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 JUNIOR STRENGTH IBUPROFEN, IBUPROFEN (OTC)
 LAX-LYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
 LEVONORGESTREL, LEVONORGESTREL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 MONTELUKAST SODIUM, MONTELUKAST SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******* L PERRIGO CO**

NAPROXEN, NAPROXEN
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 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
 LEVONORGESTREL, LEVONORGESTREL (OTC)

LABORATORIOS GRIFOLS

* LABORATORIOS GRIFOLS SA
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

LABORATORIOS SALVAT

* LABORATORIOS SALVAT SA
 OTOVEL, CIPROFLOXACIN HYDROCHLORIDE

LANDELA PHARM

* LANDELA PHARMACEUTICAL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE

LANNETT

* LANNETT CO INC
 ACETAZOLAMIDE, ACETAZOLAMIDE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 LANORINAL, ASPIRIN
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PRIMIDONE, PRIMIDONE
 PROBALAN, PROBENECID

LANNETT CO INC

* LANNETT CO INC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BACLOFEN, BACLOFEN
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 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)
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CODEINE SULFATE, CODEINE SULFATE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DANAZOL, DANAZOL
 DETROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXAMETHASONE, DEXAMETHASONE
 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
 HALOPERIDOL, HALOPERIDOL LACTATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LANNETT CO INC
 HYDROCORTISONE, HYDROCORTISONE
 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
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LOPINAVIR AND RITONAVIR, LOPINAVIR
 LORATADINE, LORATADINE (OTC)
 LORAZEPAM, LORAZEPAM
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METADATE CD, METHYLPHENIDATE HYDROCHLORIDE
 METAXALONE, METAXALONE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NIACIN, NIACIN
 OMEPRAZOLE, OMEPRAZOLE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 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
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 THEOPHYLLINE, THEOPHYLLINE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 URSODIOL, URSODIOL
 VALPROIC ACID, VALPROIC ACID

LANTHEUS MEDCL

* LANTHEUS MEDICAL IMAGING INC
 CARDIOLITE, TECHNETIUM TC-99M SESTAMIBI KIT
 DEFINITY RT, PERFLUTREN
 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
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LARKEN LABORATORIES INC
 ALLZITAL, ACETAMINOPHEN
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 DEXAMETHASONE, DEXAMETHASONE

LAURUS

* LAURUS LABS LTD
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EFAVIRENZ, LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

LAVIPHARM

* LAVIPHARM SA
 CATAPRES-TTS-1, CLONIDINE
 CATAPRES-TTS-2, CLONIDINE
 CATAPRES-TTS-3, CLONIDINE

LEADIANT BIOSCI INC

* LEADIANT BIOSCIENCES INC
 ABELCET, AMPHOTERICIN B
 CARNITOR SF, LEVOCARNITINE
 CARNITOR, LEVOCARNITINE
 CYSTARAN, CYSTEAMINE HYDROCHLORIDE
 MATULANE, PROCARBAZINE HYDROCHLORIDE

LEADING PHARMA LLC

* LEADING PHARMA LLC
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 FUROSEMIDE, FUROSEMIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROXYUREA, HYDROXYUREA
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LORAZEPAM, LORAZEPAM
 NIFEDIPINE, NIFEDIPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

LEO PHARMA AS

* LEO PHARMA AS
 DOVONEX, CALCIPOTRIENE
 ENSTILAR, BETAMETHASONE DIPROPIONATE
 FINACEA, AZELAIC ACID
 PROTOPIC, TACROLIMUS
 TACLONEX, BETAMETHASONE DIPROPIONATE

LG CHEM LTD

* LG CHEM LTD
 FACTIVE, GEMIFLOXACIN MESYLATE

LGM PHARMA

* LGM PHARMA SOLUTIONS LLC
 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
 CHENODIOL, CHENODIOL
 GLYCOPYRROLATE, GLYCOPYRROLATE
 MECAMYLAMINE HYDROCHLORIDE, MECAMYLAMINE HYDROCHLORIDE
 NABUMETONE, NABUMETONE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 URSODIOL, URSODIOL

LIEBEL-FLARSHEIM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LIEBEL-FLARSHEIM CO LLC
 CONRAY, IOTHALAMATE MEGLUMINE
 CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
 MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
 OPTIRAY 300, IOVERSOL
 OPTIRAY 320, IOVERSOL
 OPTIRAY 350, IOVERSOL
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

LIFE MOLECULAR

* LIFE MOLECULAR IMAGING LTD
 NEURACEQ, FLORBETABEN F-18

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)

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
 LEVETIRACETAM, LEVETIRACETAM
 * LOTUS PHARMACEUTICAL CO LTD NANTOU PLANT
 METHOTREXATE SODIUM, METHOTREXATE SODIUM

LOXO ONCOLOGY INC

* LOXO ONCOLOGY INC
 RETEVMO, SELPERCATINIB

LUITPOLD

* LUITPOLD PHARMACEUTICALS INC
 AMINOCAPROIC ACID, AMINOCAPROIC ACID

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

LUOXIN AUROVITAS

* LUOXIN AUROVITAS PHARMA CHENGDU CO LTD
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

LUPIN

* LUPIN INC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ANTARA (MICRONIZED), FENOFIBRATE
 BUDESONIDE, BUDESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 MIBELAS 24 FE, ETHINYL ESTRADIOL
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

- * LUPIN INC
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 QUINARETIC, HYDROCHLOROTHIAZIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SOLOSEC, SECNIDAZOLE
 TESTOSTERONE, TESTOSTERONE
 TOBRAMYCIN, TOBRAMYCIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
- * 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
 TRANDOLAPRIL, TRANDOLAPRIL
- LUPIN LTD**
- * 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
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 ATOVAQUONE, ATOVAQUONE
 AZITHROMYCIN, AZITHROMYCIN
 BEKYREE, DESOGESTREL
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 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
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DAYSEE, ETHINYL ESTRADIOL
 DECITABINE, DECITABINE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LUPIN LTD

DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ENSKYCE, DESOGESTREL
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESZOPICLONE, ESZOPICLONE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 FALLBACK SOLO, LEVONORGESTREL (OTC)
 FAMOTIDINE, FAMOTIDINE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FYAVOLV, ETHINYL ESTRADIOL
 GATIFLOXACIN, GATIFLOXACIN
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 JENCYCLA, NORETHINDRONE
 KAITLIB FE, ETHINYL ESTRADIOL
 KETOROLAC TROMETHAMINE AND PHENYLEPHRINE HYDROCHLORIDE, KETOROLAC TROMETHAMINE
 KURVELO, ETHINYL ESTRADIOL
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LAMOTRIGINE, LAMOTRIGINE
 LEFLUNOMIDE, LEFLUNOMIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LORAZEPAM, LORAZEPAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEFENAMIC ACID, MEFENAMIC ACID
 MELOXICAM, MELOXICAM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 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
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PENICILLAMINE, PENICILLAMINE
 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
 RANOLAZINE, RANOLAZINE
 RIFABUTIN, RIFABUTIN
 RUFINAMIDE, RUFINAMIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LUPIN LTD
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SUPRAX, CEFIXIME
 TADALAFIL, TADALAFIL
 TAVABOROLE, TAVABOROLE
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TESTOSTERONE, TESTOSTERONE
 TETRABENAZINE, TETRABENAZINE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 TYDEMY, DROSPIRENONE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VYFEMLA, ETHINYL ESTRADIOL
 ZILEUTON, ZILEUTON
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

LUPIN PHARMS

* LUPIN PHARMACEUTICALS INC
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 DESLORATADINE, DESLORATADINE
 DROXIDOPA, DROXIDOPA
 MELOXICAM, MELOXICAM
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 RIFAMPIN, RIFAMPIN
 SUPRAX, CEFIXIME
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

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

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 ****

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******MA GENERAL HOSP**

* MASSACHUSETTS GENERAL HOSP
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

MACLEODS PHARMS LTD

* MACLEODS PHARMACEUTICALS LTD
 ACYCLOVIR, ACYCLOVIR
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, 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
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 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
 LAMIVUDINE, LAMIVUDINE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE, LIDOCAINE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NEVIRAPINE, NEVIRAPINE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PYRAZINAMIDE, PYRAZINAMIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 TADALAFIL, TADALAFIL
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

MAIA PHARMS INC

* MAIA PHARMACEUTICALS INC
 ANGIOMAX RTU, BIVALIRUDIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MAIA PHARMACEUTICALS INC
 BACLOFEN, BACLOFEN
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE

MAINPOINTE

* MAINPOINTE PHARMACEUTICALS LLC
 TUXARIN ER, CHLORPHENIRAMINE MALEATE

MALLINCKRODT ARD

* MALLINCKRODT ARD INC
 ACTHAR GEL, CORTICOTROPIN

MALLINCKRODT HOSP

* MALLINCKRODT HOSP PRODUCTS IP LTD
 INOMAX, NITRIC OXIDE
 OFIRMEV, ACETAMINOPHEN
 UVADEX, METHOXSALEN

MANKIND PHARMA

* MANKIND PHARMA LTD
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 FENOFIBRATE, FENOFIBRATE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 RANOLAZINE, RANOLAZINE
 TIMOLOL MALEATE, TIMOLOL MALEATE

MARKSANS PHARMA

* MARKSANS PHARMA LTD
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 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
 IBUPROFEN, IBUPROFEN (OTC)
 LORATADINE, LORATADINE (OTC)
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NAPROXEN, NAPROXEN
 PARICALCITOL, PARICALCITOL

MARNEL PHARMS

* MARNEL PHARMACEUTICALS LLC
 ALA-SCALP, HYDROCORTISONE
 CROTAN, CROTAMITON

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
 CHLORZOXAZONE, CHLORZOXAZONE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLONIDINE, CLONIDINE
 CLOZAPINE, CLOZAPINE
 CYCLOSPORINE, CYCLOSPORINE
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIAZEPAM, DIAZEPAM
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 ERRIN, NORETHINDRONE
 ESTAZOLAM, ESTAZOLAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MAYNE PHARMA LLC
 ESTRADIOL, ESTRADIOL
 FABIOR, TAZAROTENE
 FLUOROURACIL, FLUOROURACIL
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVORA 0.15/30-28, ETHINYL ESTRADIOL
 LEXETTE, HALOBETASOL PROPIONATE
 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
 NEXTSTELLIS, DROSPIRENONE
 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
 BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DOFETILIDE, DOFETILIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NYSTATIN, NYSTATIN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 SOLTAMOX, TAMOXIFEN CITRATE

MAYNE PHARMA INTL

* MAYNE PHARMA INTERNATIONAL PTY LTD
 TOLSURA, ITRACONAZOLE

MC2

* MC2 THERAPEUTICS LTD
 WYNZORA, BETAMETHASONE DIPROPIONATE

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

MDD US

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MDD US OPERATIONS LLC

APOKYN, APOMORPHINE HYDROCHLORIDE
XADAGO, SAFINAMIDE MESYLATE**MDGH**

* MEDICINES DEVELOPMENT FOR GLOBAL HEALTH

MOXIDECTIN, MOXIDECTIN

MEDEFIL INC

* MEDEFIL INC

CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION**MEDEXUS**

* MEDEXUS PHARMA INC

RASUVO, METHOTREXATE

MEDICINES360

* MEDICINES360

LILETTA, LEVONORGESTREL

MEDICURE

* MEDICURE INTERNATIONAL INC

AGGRASTAT, TIROFIBAN HYDROCHLORIDE
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
ZYPITAMAG, PITAVASTATIN MAGNESIUM**MEDIMETRIKS PHARMS**

* 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)**MEITHEAL**

* MEITHEAL PHARMACEUTICALS INC

ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
AZACITIDINE, AZACITIDINE
BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
BUSULFAN, BUSULFAN
CARMUSTINE, CARMUSTINE
CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
CLOFARABINE, CLOFARABINE
CYTARABINE, CYTARABINE
DACTINOMYCIN, DACTINOMYCIN
DAPTOMYCIN, DAPTOMYCIN
DECITABINE, DECITABINE
DOXERCALCIFEROL, DOXERCALCIFEROL
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
ISOSULFAN BLUE, ISOSULFAN BLUE
LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
MILRINONE LACTATE, MILRINONE LACTATE
MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MEITHEAL PHARMACEUTICALS INC
NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
TIGECYCLINE, TIGECYCLINE

MELINTA

* MELINTA SUBSIDIARY CORP
BAXDELA, DELAFLOXACIN MEGLUMINE

MELINTA THERAP

* MELINTA THERAPEUTICS LLC
KIMYRSA, ORITAVANCIN DIPHOSPHATE
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
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
BELSOMRA, SUVOREXANT
DIPROLENE, BETAMETHASONE DIPROPIONATE
ELOCON, MOMETASONE FUROATE
INVANZ, ERTAPENEM SODIUM
ISENTRESS HD, RALTEGRAVIR POTASSIUM
ISENTRESS, RALTEGRAVIR POTASSIUM
JANUMET XR, METFORMIN HYDROCHLORIDE
JANUMET, METFORMIN HYDROCHLORIDE
JANUVIA, SITAGLIPTIN PHOSPHATE
LOTRISONE, BETAMETHASONE DIPROPIONATE
NOXAFIL, POSACONAZOLE
PREVYMIS, LETERMOVIR
REBETOL, RIBAVIRIN
SEGLUROMET, ERTUGLIFLOZIN
STEGLATRO, ERTUGLIFLOZIN
STEGLUJAN, ERTUGLIFLOZIN
STROMEKTOL, IVERMECTIN
TEMODAR, TEMOZOLOMIDE
VERQUVO, VERICIGUAT
ZEPATIER, ELBASVIR

* MERCK SHARP AND DOHME CORP A SUB OF MERCK AND CO INC
WELIREG, BELZUTIFAN

MERIDIAN MEDCL

* MERIDIAN MEDICAL TECHNOLOGIES INC
DUODOTE, ATROPINE

MERIDIAN MEDCL TECHN

* MERIDIAN MEDICAL TECHNOLOGIES INC
ATROPEN, ATROPINE
SEIZALAM, MIDAZOLAM HYDROCHLORIDE

MERRO PHARM USA

* MERRO PHARMACEUTICAL USA INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MERRO PHARMACEUTICAL USA INC
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

MERZ PHARMS

* MERZ PHARMACEUTICALS LLC
 CUVPOSA, GLYCOPYRROLATE

METACEL PHARMS LLC

* METACEL PHARMACEUTICALS LLC
 OZOBAX, BACLOFEN

METHAPHARM

* METHAPHARM INC
 PROVOCHOLINE, METHACHOLINE CHLORIDE

METUCHEN PHARMS

* METUCHEN PHARMACEUTICALS LLC
 STENDRA, AVANAFIL

MICRO LABS

* MICRO LABS LTD
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 BIMATOPROST, BIMATOPROST
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CELECOXIB, CELECOXIB
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 DALFAMPRIDINE, DALFAMPRIDINE
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 GLIMEPIRIDE, GLIMEPIRIDE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 LINCOMYCIN, LINCOMYCIN
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEFENAMIC ACID, MEFENAMIC ACID
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 PIROXICAM, PIROXICAM
 RANOLAZINE, RANOLAZINE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 SIMVASTATIN, SIMVASTATIN
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TAFLUPROST, TAFLUPROST
 TELMISARTAN, TELMISARTAN
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRAVOPROST, TRAVOPROST
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

MICRO LABS LTD

* MICRO LABS LTD
 NEVIRAPINE, NEVIRAPINE

MICRO LABS LTD INDIA

* MICRO LABS LTD INDIA
 ACETAZOLAMIDE, ACETAZOLAMIDE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 CROMOLYN SODIUM, CROMOLYN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MICRO LABS LTD INDIA
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN

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
 BENZONATATE, BENZONATATE
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 BUTAPAP, ACETAMINOPHEN
 CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
 CHLORZOAZONE, CHLORZOAZONE
 ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 METHAZOLAMIDE, METHAZOLAMIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

MIKART INC

* MIKART INC
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN

MILLA PHARMS

* MILLA PHARMACEUTICALS INC
 SODIUM ACETATE, SODIUM ACETATE

MILLICENT

* MILLICENT HOLDINGS LTD
 FEMRING, ESTRADIOL ACETATE
 * MILLICENT PHARMA LTD
 INTRAROSA, PRASTERONE

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

MIRUM

* MIRUM PHARMACEUTICALS INC
 LIVMARLI, MARALIXIBAT CHLORIDE

MISEMER

* MISEMER PHARMACEUTICALS INC
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 KETOPROFEN, KETOPROFEN

MISSION PHARMA

* MISSION PHARMACAL CO
 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)
 THIOLA EC, TIOPRONIN

MITSUBISHI HOLDINGS

* MITSUBISHI TANABE PHARMA HOLDINGS AMERICA INC
 EXSERVAN, RILUZOLE

MITSUBISHI TANABE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MITSUBISHI TANABE PHARMA CORP
RADICAVA, EDARAVONE

MLV

* MLV PHARMA LLC
CARISOPRODOL, CARISOPRODOL
METHOCARBAMOL, METHOCARBAMOL
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE

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
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

MSD MERCK CO

* MERCK SHARP AND DOHME CORP A SUB OF MERCK AND CO INC
DELSTRIGO, DORAVIRINE
EMEND, APREPITANT
NOXAFIL POWDERMIX KIT, POSACONAZOLE
PIFELTRO, DORAVIRINE
RECARBRIO, CILASTATIN SODIUM
SINGULAIR, MONTELUKAST SODIUM

MSN

* MSN LABORATORIES PRIVATE LTD
ABIRATERONE ACETATE, ABIRATERONE ACETATE
ALBENDAZOLE, ALBENDAZOLE
AMINOCAPROIC ACID, AMINOCAPROIC ACID
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
CAPECITABINE, CAPECITABINE
CLOFARABINE, CLOFARABINE
DECITABINE, DECITABINE
DEFERASIROX, DEFERASIROX
DIMETHYL FUMARATE, DIMETHYL FUMARATE
DOFETILIDE, DOFETILIDE
ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
FEBUXOSTAT, FEBUXOSTAT
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
PACLITAXEL, PACLITAXEL
PREGABALIN, PREGABALIN
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
SILODOSIN, SILODOSIN
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
THIOTEPA, THIOTEPA
TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE
TOREMIFENE CITRATE, TOREMIFENE CITRATE
TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******MSN PHARMS INC**

* MSN PHARMACEUTICALS INC
DROXIDOPA, DROXIDOPA

MURTY PHARMS

* MURTY PHARMACEUTICALS INC
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
DIPYRIDAMOLE, DIPYRIDAMOLE

MYLAN

* MYLAN PHARMACEUTICALS
METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

* MYLAN PHARMACEUTICALS INC
ABIRATERONE ACETATE, ABIRATERONE ACETATE
ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
ACETAMINOPHEN, ACETAMINOPHEN
ACITRETIN, ACITRETIN
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALLOPURINOL, ALLOPURINOL
ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
AMBRISENTAN, AMBRISENTAN
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
ATENOLOL, ATENOLOL
ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
AVITA, TRETINOIN
BEPOTASTINE BESILATE, BEPOTASTINE BESILATE
BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
BUDESONIDE, BUDESONIDE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
CARBIDOPA AND LEVODOPA, CARBIDOPA
CARVEDILOL, CARVEDILOL
CELECOXIB, CELECOXIB
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
CHLORTHALIDONE, CHLORTHALIDONE
CIMETIDINE, CIMETIDINE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLOBAZAM, CLOBAZAM
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CLOZAPINE, CLOZAPINE
COLCHICINE, COLCHICINE
CYSTAGON, CYSTEAMINE BITARTRATE
DENA VIR, PENCICLOVIR
DIAZEPAM, DIAZEPAM
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIMETHYL FUMARATE, DIMETHYL FUMARATE
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
ERYGEL, ERYTHROMYCIN
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
ESTRADIOL, ESTRADIOL
ETOPOSIDE, ETOPOSIDE
EVEROLIMUS, EVEROLIMUS
EVOCLIN, CLINDAMYCIN PHOSPHATE
EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
EXTINA, KETOCONAZOLE
FEBUXOSTAT, FEBUXOSTAT
FENOFIBRATE, FENOFIBRATE
FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN PHARMACEUTICALS INC
 FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 FUROSEMIDE, FUROSEMIDE
 GLATIRAMER ACETATE, GLATIRAMER ACETATE
 GLIPIZIDE, GLIPIZIDE
 HALCINONIDE, HALCINONIDE
 HALOPERIDOL, HALOPERIDOL
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IMPEKLO, CLOBETASOL PROPIONATE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LEVETIRACETAM, LEVETIRACETAM
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 LUXIQ, BETAMETHASONE VALERATE
 MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
 MENTAX, BUTENAFINE HYDROCHLORIDE
 MERCAPTOPYRINE, MERCAPTOPYRINE
 MESALAMINE, MESALAMINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHIMAZOLE, METHIMAZOLE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIRTAZAPINE, MIRTAZAPINE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NAPROXEN AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 NEVIRAPINE, NEVIRAPINE
 NISOLDIPINE, NISOLDIPINE
 OLANZAPINE, OLANZAPINE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 OLUX E, CLOBETASOL PROPIONATE
 OLUX, CLOBETASOL PROPIONATE
 ONDANSETRON, ONDANSETRON
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 PERPHENAZINE, PERPHENAZINE
 PHENYTEK, PHENYTOIN SODIUM
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREDNISONE, PREDNISONE
 PREGABALIN, PREGABALIN
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 RUFINAMIDE, RUFINAMIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SUNITINIB MALATE, SUNITINIB MALATE
 SYMFI LO, EFAVIRENZ
 TACROLIMUS, TACROLIMUS
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TETRABENAZINE, TETRABENAZINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

- * MYLAN PHARMACEUTICALS INC
 - THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
 - THIOTHIXENE, THIOTHIXENE
 - TICAGRELOR, TICAGRELOR
 - TIMOLOL MALEATE, TIMOLOL MALEATE
 - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 - TRAVOPROST, TRAVOPROST
 - TRETINOIN, TRETINOIN
 - TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 - URSODIOL, URSODIOL
 - VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 - VALSARTAN, VALSARTAN
 - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 - VUSION, MICONAZOLE NITRATE
 - WIXELA INHUB, FLUTICASONE PROPIONATE
 - ZONALON, DOXEPIN HYDROCHLORIDE
 - ZOVIRAX, ACYCLOVIR
- * MYLAN PHARMACEUTICALS INC A VIATRIS CO
 - AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
 - PREDNISONE, PREDNISONE
 - PREGABALIN, PREGABALIN
 - TELMISARTAN, TELMISARTAN
 - VALPROATE SODIUM, VALPROATE SODIUM

MYLAN ASI

- * MYLAN ASI LLC
 - ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 - ADENOSINE, ADENOSINE
 - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE

MYLAN INSTITUTIONAL

- * MYLAN INSTITUTIONAL INC
 - BUSULFAN, BUSULFAN
 - SULFAMYLOL, MAFENIDE ACETATE
- * MYLAN INSTITUTIONAL LLC
 - ALOPRIM, ALLOPURINOL SODIUM
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 - ARGATROBAN, ARGATROBAN
 - AZACITIDINE, AZACITIDINE
 - BIVALIRUDIN, BIVALIRUDIN
 - CIDOFOVIR, CIDOFOVIR
 - COSYNTROPIN, COSYNTROPIN
 - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 - 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
 - MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 - METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 - METHOCARBAMOL, METHOCARBAMOL
 - NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 - OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 - PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 - RIMSO-50, DIMETHYL SULFOXIDE
 - ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 - SOTRADECOL, SODIUM TETRADECYL SULFATE
 - THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 - TRANEXAMIC ACID, TRANEXAMIC ACID
 - ULTIVA, REMIFENTANIL HYDROCHLORIDE

MYLAN IRELAND LTD

- * MYLAN IRELAND LTD
 - ARIXTRA, FONDAPARINUX SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MYLAN IRELAND LTD**

MIACALCIN, CALCITONIN SALMON
 PRETOMANID, PRETOMANID
 YUPELRI, REVEFENACIN

MYLAN LABS LTD*** MYLAN LABORATORIES LTD**

ADENOSINE, ADENOSINE
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 BACLOFEN, BACLOFEN
 CIMDUO, LAMIVUDINE
 CLADRIBINE, CLADRIBINE
 CLOFARABINE, CLOFARABINE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 DAPTOMYCIN, DAPTOMYCIN
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOCETAXEL, DOCETAXEL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE HYCLATE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 EPTIFIBATIDE, EPTIFIBATIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 ETOMIDATE, ETOMIDATE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM, HEPARIN SODIUM
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 LAMIVUDINE, LAMIVUDINE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEVETIRACETAM, LEVETIRACETAM
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 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
 MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE
 OXALIPLATIN, OXALIPLATIN
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIFAMPIN, RIFAMPIN
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SYMFI, EFAVIRENZ
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN LABORATORIES LTD
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

MYLAN PHARMS INC

* MYLAN PHARMACEUTICALS INC
 ABACAVIR SULFATE, ABACAVIR SULFATE
 ACYCLOVIR, ACYCLOVIR
 AMNESTEEM, ISOTRETINOIN
 ARMODAFINIL, ARMODAFINIL
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AVITA, TRETINOIN
 CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 ESZOPICLONE, ESZOPICLONE
 FENOFIBRATE, FENOFIBRATE
 FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 LANSOPRAZOLE, LANSOPRAZOLE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 MAXZIDE, HYDROCHLOROTHIAZIDE
 MAXZIDE-25, HYDROCHLOROTHIAZIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 NEVIRAPINE, NEVIRAPINE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PINDOLOL, PINDOLOL
 RILUZOLE, RILUZOLE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VORICONAZOLE, VORICONAZOLE
 ZIDOVUDINE, ZIDOVUDINE

* MYLAN PHARMACEUTICALS INC.
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM

MYLAN SPCLT VIATRIS

* MYLAN SPECIALTY LP A VIATRIS CO
 DIPENTUM, OLSALAZINE SODIUM

MYLAN SPECIALITY LP

* MYLAN SPECIALTY LP
 ACCUNEB, ALBUTEROL SULFATE
 COLYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
 CORTIFOAM, HYDROCORTISONE ACETATE
 DEPEN, PENICILLAMINE
 DYMISTA, AZELASTINE HYDROCHLORIDE
 EDLUAR, ZOLPIDEM TARTRATE
 ELESTRIN, ESTRADIOL
 EPIFOAM, HYDROCORTISONE ACETATE
 EPIPEN JR., EPINEPHRINE
 EPIPEN, EPINEPHRINE
 FELBATOL, FELBAMATE
 GASTROCROM, CROMOLYN SODIUM
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* 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
SCOPOLAMINE, SCOPOLAMINE
XULANE, ETHINYL ESTRADIOL

MYOVANT SCIENCES

* MYOVANT SCIENCES GMBH
MYFEMBREE, ESTRADIOL
ORGOVYX, RELUGOLIX

**** N ******NAARI PTE LTD**

* NAARI PTE LTD
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LEVONORGESTREL, LEVONORGESTREL (OTC)
NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
NORETHINDRONE, NORETHINDRONE
NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

NABRIVA

* NABRIVA THERAPEUTICS IRELAND DAC
XENLETA, LEFAMULIN ACETATE

NAL PHARM

* NAL PHARMACEUTICAL GROUP LTD
LIDOCAINE, LIDOCAINE

NALPROPION

* NALPROPION PHARMACEUTICALS LLC
CONTRAVE, BUPROPION HYDROCHLORIDE

NAMIGEN LLC

* NAMIGEN LLC
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM

NANG KUANG PHARM CO

* NANG KUANG PHARMACEUTICAL CO LTD
ICATIBANT ACETATE, ICATIBANT ACETATE
LINEZOLID, LINEZOLID

NANJING

* NANJING SIMCERE DONGYUAN PHARMACEUTICAL CO LTD
CELECOXIB, CELECOXIB

NANJING KING-FRIEND

* NANJING KING-FRIEND BIOCHEMICAL PHARMACEUTICAL CO LTD
ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
HEPARIN SODIUM, HEPARIN SODIUM

NAPO PHARMS INC

* NAPO PHARMACEUTICALS INC
MYTESI, CROFELEMER

NATCO

* NATCO PHARMA LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NATCO PHARMA LTD
 ALPRAZOLAM, ALPRAZOLAM
 CARISOPRODOL, CARISOPRODOL
 GLYCOPYRROLATE, GLYCOPYRROLATE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE

NATCO PHARMA

* NATCO PHARMA LTD
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE

NATCO PHARMA LTD

* NATCO PHARMA LIMITED
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE

* NATCO PHARMA LTD
 ANASTROZOLE, ANASTROZOLE
 ARMODAFINIL, ARMODAFINIL
 AZACITIDINE, AZACITIDINE
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 LANSOPRAZOLE, LANSOPRAZOLE
 LANTHANUM CARBONATE, LANTHANUM CARBONATE
 LAPATINIB DITOSYLATE, LAPATINIB DITOSYLATE
 LETROZOLE, LETROZOLE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

NAVINTA LLC

* NAVINTA LLC
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 CARMUSTINE, CARMUSTINE
 FAMOTIDINE, FAMOTIDINE
 FOMEPIZOLE, FOMEPIZOLE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
 METHOCARBAMOL, METHOCARBAMOL
 PENICILLAMINE, PENICILLAMINE
 REMIFENTANIL HYDROCHLORIDE, REMIFENTANIL HYDROCHLORIDE
 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

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
 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
 BUDESONIDE, BUDESONIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NEPHRON PHARMACEUTICALS CORP

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

NESHER PHARMS

* NESHER PHARMACEUTICALS USA LLC

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
MORPHINE SULFATE, MORPHINE SULFATE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN**NEURELIS INC**

* NEURELIS INC

VALTOCO, DIAZEPAM

NEUROCRINE

* NEUROCRINE BIOSCIENCES INC

INGREZZA, VALBENZAZINE TOSYLATE
ONGENTYS, OPICAPONE**NEXTWAVE**

* NEXTWAVE PHARMACEUTICALS INC A SUB OF TRIS PHARMA INC

QUILLIVANT XR, METHYLPHENIDATE HYDROCHLORIDE

NEXTWAVE PHARMS

* NEXTWAVE PHARMACEUTICALS INC

QUILLICHEW ER, METHYLPHENIDATE HYDROCHLORIDE

NEXUS PHARMS

* NEXUS PHARMACEUTICALS INC

ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
BUSULFAN, BUSULFAN
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
EMERPHED, EPHEDRINE SULFATE
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
POTASSIUM CHLORIDE 10MEQ, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 20MEQ, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 40MEQ, POTASSIUM CHLORIDE
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE**NIAGARA PHARMS**

* NIAGARA PHARMACEUTICALS INC

PUR-WASH, PURIFIED WATER (OTC)

NIPPON SHINYAKU

* NIPPON SHINYAKU CO LTD

VILTEPSO, VILTOLARSEN

NIVAGEN PHARMS INC

* NIVAGEN PHARMACEUTICALS INC

DECITABINE, DECITABINE
GLYCOPYRROLATE, GLYCOPYRROLATE
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
TEMOZOLOMIDE, TEMOZOLOMIDE**NODEN PHARMA**

* NODEN PHARMA DAC

TEKTURNA HCT, ALISKIREN HEMIFUMARATE
TEKTURNA, ALISKIREN HEMIFUMARATE**NORTEC DEV ASSOC**

* NORTEC DEVELOPMENT ASSOC INC

MORPHINE SULFATE, MORPHINE SULFATE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE**NORTHLAND**

* NORTHLAND NUCLEAR MEDICINE LLC

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******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 REDHALER, BECLOMETHASONE DIPROPIONATE

NOSTRUM LABS INC

* NOSTRUM LABORATORIES INC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACETAZOLAMIDE, ACETAZOLAMIDE
 BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
 CALCIUM ACETATE, CALCIUM ACETATE
 CARBAMAZEPINE, CARBAMAZEPINE
 CARISOPRODOL, CARISOPRODOL
 DAPSONE, DAPSONE
 ELIXOPHYLLIN, THEOPHYLLINE
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 HYDROCODONE, HYDROCODONE BITARTRATE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 NITROFURANTOIN, NITROFURANTOIN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PINDOLOL, PINDOLOL
 PIROXICAM, PIROXICAM
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 SUCRALFATE, SUCRALFATE
 THEOPHYLLINE, THEOPHYLLINE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

NOSTRUM PHARMS LLC

* NOSTRUM PHARMACEUTICALS LLC
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

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
 ALOMIDE, LODOXAMIDE TROMETHAMINE
 ARGATROBAN, ARGATROBAN
 ARRANON, NELARABINE
 AZOPT, BRINZOLAMIDE
 BETOPTIC S, BETAXOLOL HYDROCHLORIDE
 CILOXAN, CIPROFLOXACIN HYDROCHLORIDE
 CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE
 CIPRODEX, CIPROFLOXACIN
 COARTEM, ARTEMETHER
 DESFERAL, DEFEROXAMINE MESYLATE
 DIOVAN HCT, HYDROCHLOROTHIAZIDE
 DIOVAN, VALSARTAN
 DUREZOL, DIFLUPREDNATE
 EGATEN, TRICLABENDAZOLE
 EXELON, RIVASTIGMINE
 EXFORGE HCT, AMLODIPINE BESYLATE
 EXFORGE, AMLODIPINE BESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVARTIS PHARMACEUTICALS CORP
 EXJADE, DEFERASIROX
 FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE
 GILENYA, FINGOLIMOD HYDROCHLORIDE
 GLEEVEC, IMATINIB MESYLATE
 HYCAMTIN, TOPOTECAN HYDROCHLORIDE
 ILEVRO, NEPAFENAC
 IOPIDINE, APRACLONIDINE HYDROCHLORIDE
 ISOPTO CARPINE, PILOCARPINE HYDROCHLORIDE
 JADENU SPRINKLE, DEFERASIROX
 KISQALI FEMARA CO-PACK (COPACKAGED), LETROZOLE
 KISQALI, RIBOCICLIB SUCCINATE
 LEQVIO, INCLISIRAN SODIUM
 LESCOL XL, FLUVASTATIN SODIUM
 LOTREL, AMLODIPINE BESYLATE
 MAXIDEX, DEXAMETHASONE
 MAXITROL, DEXAMETHASONE
 MAYZENT, SIPONIMOD FUMARIC ACID
 MEKINIST, TRAMETINIB DIMETHYL SULFOXIDE
 MOXEZA, MOXIFLOXACIN HYDROCHLORIDE
 MYFORTIC, MYCOPHENOLIC SODIUM
 NEORAL, CYCLOSPORINE
 NEVANAC, NEPAFENAC
 OMNIPRED, PREDNISOLONE ACETATE
 PATANASE, OLOPATADINE HYDROCHLORIDE
 PIQRAY, ALPELISIB
 PROMACTA KIT, ELTROMBOPAG OLAMINE
 PROMACTA, ELTROMBOPAG OLAMINE
 RECLAST, ZOLEDRONIC ACID
 RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE
 RITALIN, METHYLPHENIDATE HYDROCHLORIDE
 RYDAPT, MIDOSTAURIN
 SANDIMMUNE, CYCLOSPORINE
 SANDOSTATIN LAR, OCTREOTIDE ACETATE
 SANDOSTATIN, OCTREOTIDE ACETATE
 SCEMBLIX, ASCIMINIB HYDROCHLORIDE
 TAFINLAR, DABRAFENIB MESYLATE
 TASIGNA, NILOTINIB HYDROCHLORIDE
 TEGRETOL, CARBAMAZEPINE
 TEGRETOL-XR, CARBAMAZEPINE
 TOBRADEX, DEXAMETHASONE
 TOBREX, TOBRAMYCIN
 TRAVATAN Z, TRAVOPROST
 TRISENCE, TRIAMCINOLONE ACETONIDE
 TRILEPTAL, OXCARBAZEPINE
 TYKERB, LAPATINIB DITOSYLATE
 VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE
 VIVELLE-DOT, ESTRADIOL
 VOTRIENT, PAZOPANIB HYDROCHLORIDE
 XIIDRA, LIFITEGRAST
 ZOFRAN, ONDANSETRON HYDROCHLORIDE
 ZOMETA, ZOLEDRONIC ACID
 ZORTRESS, EVEROLIMUS
 ZYKADIA, CERITINIB

NOVARTIS PHARM

* NOVARTIS PHARMACEUTICAL CORP
 AFINITOR DISPERZ, EVEROLIMUS
 TABRECTA, CAPMATINIB HYDROCHLORIDE

NOVARTIS PHARMS

* NOVARTIS PHARMACEUTICALS CORP
 FEMARA, LETROZOLE

NOVARTIS PHARMS CORP

* NOVARTIS PHARMACEUTICALS CORP
 ENTRESTO, SACUBITRIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVARTIS PHARMACEUTICALS CORP
JADENU, DEFERASIROX

NOVAST LABS

* NOVAST LABORATORIES CHINA LTD
NORETHINDRONE, NORETHINDRONE

* NOVAST LABORATORIES LTD
ACETAZOLAMIDE, ACETAZOLAMIDE
CARBOPLATIN, CARBOPLATIN
CARISOPRODOL, CARISOPRODOL
CHABELINA FE, ETHINYL ESTRADIOL
CHLORTHALIDONE, CHLORTHALIDONE
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DOLISHALE, ETHINYL ESTRADIOL
ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
HER STYLE, LEVONORGESTREL (OTC)
INDOMETHACIN, INDOMETHACIN
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
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
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
LO-MALMOREDE, ETHINYL ESTRADIOL
MAFENIDE ACETATE, MAFENIDE ACETATE
MALMOREDE, ETHINYL ESTRADIOL
MELAMISA, DROSPIRENONE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
NIFEDIPINE, NIFEDIPINE
NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
NORETHINDRONE, NORETHINDRONE
OXALIPLATIN, OXALIPLATIN
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
PIMTREA, DESOGESTREL
PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE
PROBENECID AND COLCHICINE, COLCHICINE
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
QUININE SULFATE, QUININE SULFATE
SETLAKIN, ETHINYL ESTRADIOL
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TOLCAPONE, TOLCAPONE
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
TRIAZOLAM, TRIAZOLAM
YELA, DROSPIRENONE

NOVAST LABS LTD

* NOVAST LABORATORIES LTD
DASETTA 1/35, ETHINYL ESTRADIOL
DASETTA 7/7/7, ETHINYL ESTRADIOL
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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* 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
MONOBASIC SODIUM PHOSPHATE AND DIBASIC SODIUM PHOSPHATE, SODIUM PHOSPHATE, DIBASIC,
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
PEG-3350, SODIUM SULFATE, SODIUM CHLORIDE, POTASSIUM CHLORIDE, SODIUM ASCORBATE AND
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
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)

NOVEN

* NOVEN PHARMACEUTICALS INC
MINIVELLE, ESTRADIOL

NOVEN PHARMS INC

* NOVEN PHARMACEUTICALS INC
COMBIPATCH, ESTRADIOL
DAYTRANA, METHYLPHENIDATE

NOVITIUM PHARMA

* NOVITIUM PHARMA LLC
ACETAZOLAMIDE, ACETAZOLAMIDE
ACYCLOVIR, ACYCLOVIR
ALPRAZOLAM, ALPRAZOLAM
ATENOLOL AND CHLORTHALIDONE, ATENOLOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVITIUM PHARMA LLC
 BETAINE, BETAINE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 CARGLUMIC ACID, CARGLUMIC ACID
 CHLORZOXAZONE, CHLORZOXAZONE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 DAPSONE, DAPSONE
 DIAZOXIDE, DIAZOXIDE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIGOXIN, DIGOXIN
 ESTAZOLAM, ESTAZOLAM
 FAMOTIDINE, FAMOTIDINE
 FELBAMATE, FELBAMATE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 LEVOCARNITINE SF, LEVOCARNITINE
 LEVOCARNITINE, LEVOCARNITINE
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 MELOXICAM, MELOXICAM
 NAPROXEN, NAPROXEN
 NITISINONE, NITISINONE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREDNISONE, PREDNISONE
 PYRAZINAMIDE, PYRAZINAMIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RIFABUTIN, RIFABUTIN
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SIROLIMUS, SIROLIMUS
 THIOTHIXENE, THIOTHIXENE
 TRANLYCYPROMINE SULFATE, TRANLYCYPROMINE SULFATE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 VALSARTAN, VALSARTAN

NOVO

* NOVO NORDISK INC
 MACRILEN, MACIMORELIN ACETATE
 OZEMPIC, SEMAGLUTIDE
 RYBELSUS, SEMAGLUTIDE
 SAXENDA, LIRAGLUTIDE RECOMBINANT
 WEGOVY, SEMAGLUTIDE

NOVO NORDISK

* NOVO NORDISK PHARMACEUTICALS INC
 GLUCAGEN, GLUCAGON HYDROCHLORIDE

NOVO NORDISK INC

* NOVO NORDISK INC
 VAGIFEM, ESTRADIOL
 VICTOZA, LIRAGLUTIDE RECOMBINANT

NOVOCOL INC

* NOVOCOL INC
 DYCLOPRO, DYCLONINE HYDROCHLORIDE

NPS PHARMS INC

* NPS PHARMACEUTICALS INC
 GATTEX KIT, TEDUGLUTIDE RECOMBINANT

NUKEMED

* NUKEMED INC DBA SPECTRONRX
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

NUVO PHARM

* NUVO PHARMACEUTICAL INC
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NUVO PHARMACEUTICAL INC
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

NUVO PHARMS INC

* NUVO PHAMACEUTICALS INC
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
FOLIC ACID, FOLIC ACID
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
NAPROXEN, NAPROXEN
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
SULFASALAZINE, SULFASALAZINE

NXDC

* NX DEVELOPMENT CORP
GLEOLAN, AMINOLEVULINIC ACID HYDROCHLORIDE

**** O ******OCULAR THERAPEUTIX**

* OCULAR THERAPEUTIX INC
DEXTENZA, DEXAMETHASONE

OHM

* OHM CORP
IBUPROFEN, IBUPROFEN (OTC)

OHM LABS

* OHM LABORATORIES INC
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

OHM LABS INC

* OHM LABORATORIES INC
EZETIMIBE, EZETIMIBE
GUAIFENESIN, GUAIFENESIN (OTC)
VALSARTAN, VALSARTAN

OMEROS

* OMEROS CORP
OMIDRIA, KETOROLAC TROMETHAMINE

ON TARGET LABS

* ON TARGET LABORATORIES INC
CYTALUX, PAFOLACIANINE SODIUM

ONCOGEN PHARMA

* ONCOGEN PHARMA MALAYSIA SDN BHD
ABIRATERONE ACETATE, ABIRATERONE ACETATE

ONYX THERAP

* ONYX THERAPEUTICS INC A WHOLLY OWNED SUB OF AMGEN INC
KYPROLIS, CARFILZOMIB

OPTINOSE US INC

* OPTINOSE US INC
XHANCE, FLUTICASONE PROPIONATE

ORAPHARMA

* ORAPHARMA INC
ARESTIN, MINOCYCLINE HYDROCHLORIDE

ORBION PHARMS

* ORBION PHARMACEUTICALS PRIVATE LTD
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ARIPIPRAZOLE, ARIPIPRAZOLE
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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** O **

* ORBION PHARMACEUTICALS PRIVATE LTD
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN, LEVOFLOXACIN
 MODAFINIL, MODAFINIL
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZALEPLON, ZALEPLON
 ZOLMITRIPTAN, ZOLMITRIPTAN

ORCHID HLTHCARE

* ORCHID HEALTHCARE
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFDINIR, CEFDINIR
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEFPROZIL, CEFPROZIL
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CEPHALEXIN, CEPHALEXIN

OREXO US INC

* OREXO US INC
 ZUBSOLV, BUPRENORPHINE HYDROCHLORIDE

ORGANON

* ORGANON LLC
 ASMANEX HFA, MOMETASONE FUROATE
 ASMANEX TWISTHALER, MOMETASONE FUROATE
 CLARINEX-D 12 HOUR, DESLORATADINE
 DULERA, FORMOTEROL FUMARATE
 FOSAMAX PLUS D, ALENDRONATE SODIUM

* ORGANON LLC A SUB OF ORGANON AND CO
 CELESTONE SOLUSPAN, BETAMETHASONE ACETATE
 CLARINEX, DESLORATADINE
 COZAAR, LOSARTAN POTASSIUM
 HYZAAR, HYDROCHLOROTHIAZIDE
 SINEMET, CARBIDOPA
 VYTORIN, EZETIMIBE
 ZETIA, EZETIMIBE
 ZOCOR, SIMVASTATIN

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
 GANIRELIX ACETATE, GANIRELIX ACETATE
 NEXPLANON, ETONOGESTREL
 REMERON SOLTAB, MIRTAZAPINE
 REMERON, MIRTAZAPINE

ORIENT PHARMA CO LTD

* ORIENT PHARMA CO LTD
 CARISOPRODOL, CARISOPRODOL
 GLYBURIDE, GLYBURIDE
 MIGLITOL, MIGLITOL
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

ORIGIN

* ORIGIN BIOSCIENCES INC
 NULIBRY, FOSDENOPTERIN HYDROBROMIDE

ORION PHARMA

* ORION PHARMA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** O **

* ORION PHARMA

COMTAN, ENTACAPONE
 STALEVO 100, CARBIDOPA
 STALEVO 125, CARBIDOPA
 STALEVO 150, CARBIDOPA
 STALEVO 200, CARBIDOPA
 STALEVO 50, CARBIDOPA
 STALEVO 75, CARBIDOPA

OSI PHARMS

* OSI PHARMACEUTICALS LLC
 TARCEVA, ERLOTINIB HYDROCHLORIDE

OSMOTICA

* OSMOTICA KERESKEDELMI ES SZOLGALTATO KFT
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

OSMOTICA PHARM

* OSMOTICA PHARMACEUTICAL CORP
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

OSMOTICA PHARM US

* OSMOTICA PHARMACEUTICAL US LLC
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 NIFEDIPINE, NIFEDIPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

OTSUKA

* OTSUKA PHARMACEUTICAL CO LTD
 ABILIFY MYCITE KIT, ARIPIPRAZOLE
 ABILIFY, ARIPIPRAZOLE
 DACOGEN, DEGITABINE
 INQOVI, CEDAZURIDINE
 JYNARQUE, TOLVAPTAN
 REXULTI, BREXPIPRAZOLE
 SAMSCA, TOLVAPTAN

OTSUKA PHARM

* OTSUKA PHARMACEUTICAL CO LTD
 BUSULFEX, BUSULFAN

OTSUKA PHARM CO LTD

* OTSUKA PHARMACEUTICAL CO LTD
 ABILIFY MAINTENA KIT, ARIPIPRAZOLE

OTTER PHARMS

* OTTER PHARMACEUTICALS LLC
 OTREXUP, METHOTREXATE

OUTLOOK PHARMS

* OUTLOOK PHARMACEUTICALS INC
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE

OVERSEAS

* OVERSEAS PHARMACEUTICALS LTD
 LEVETIRACETAM, LEVETIRACETAM

OXFORD PHARMS

* OXFORD PHARMACEUTICALS LLC
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 BACLOFEN, BACLOFEN
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARISOPRODOL, CARISOPRODOL
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LORAZEPAM, LORAZEPAM
 METHOCARBAMOL, METHOCARBAMOL
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PRIMIDONE, PRIMIDONE
 RIMACTANE, RIFAMPIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** O ****

* OXFORD PHARMACEUTICALS LLC
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SIMVASTATIN, SIMVASTATIN
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

OYSTER POINT PHARMA

* OYSTER POINT PHARMA INC
 TYRVAYA, VARENICLINE TARTRATE

**** P ******P AND L**

* P AND L DEVELOPMENT LLC
 ADAPALENE, ADAPALENE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
 DOCOSANOL, DOCOSANOL (OTC)
 FAMOTIDINE, FAMOTIDINE (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MICONAZOLE 7, MICONAZOLE NITRATE (OTC)
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)

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

PADAGIS ISRAEL

* PADAGIS ISRAEL PHARMACEUTICALS LTD
 ACYCLOVIR, ACYCLOVIR
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 AMMONIUM LACTATE, AMMONIUM LACTATE
 AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE, AZELASTINE HYDROCHLORIDE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE AND BETHAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DESOXIMETASONE, DESOXIMETASONE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 ECONAZOLE NITRATE, ECONAZOLE NITRATE
 ESTRADIOL, ESTRADIOL
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLURANDRENOLIDE, FLURANDRENOLIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 GYNAZOLE-1, BUTOCONAZOLE NITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PADAGIS ISRAEL PHARMACEUTICALS LTD
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 IMIQUIMOD, IMIQUIMOD
 KETOCONAZOLE, KETOCONAZOLE
 MESALAMINE, MESALAMINE
 METRONIDAZOLE, METRONIDAZOLE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MUPIROCIN, MUPIROCIN
 NITROGLYCERIN, NITROGLYCERIN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 PERMETHRIN, PERMETHRIN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 SUMATRIPTAN, SUMATRIPTAN
 SUMATRIPTAN, SUMATRIPTAN SUCCINATE
 TERCONAZOLE, TERCONAZOLE
 TESTOSTERONE, TESTOSTERONE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN

PADAGIS US

* PADAGIS US LLC
 ATOVAQUONE, ATOVAQUONE
 BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BACITRACIN, BACITRACIN
 BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE, BACITRACIN ZINC
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CALCIUM ACETATE, CALCIUM ACETATE
 CENTANY, MUPIROCIN
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CICLOPIROX, CICLOPIROX
 CLINDA-DERM, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLINDESSE, CLINDAMYCIN PHOSPHATE
 CLINDETS, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 COMPRO, PROCHLORPERAZINE
 CYCLOSPORINE, CYCLOSPORINE
 DESONIDE, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 ENTOCORT EC, BUDESONIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 EVAMIST, ESTRADIOL
 FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE,
 HYDROCORTISONE, HYDROCORTISONE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 IBUPROFEN, IBUPROFEN
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 MIDAMOR, AMILORIDE HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MORPHINE SULFATE, MORPHINE SULFATE
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
 NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
 NYSTATIN, NYSTATIN
 NYSTOP, NYSTATIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PADAGIS US LLC

PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PODOFILOX, PODOFILOX
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 REPAGLINIDE, REPAGLINIDE
 SCOPOLAMINE, SCOPOLAMINE
 SELENIUM SULFIDE, SELENIUM SULFIDE
 STIE-CORT, HYDROCORTISONE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TAVABOROLE, TAVABOROLE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TRETINOIN, TRETINOIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE

PALATIN TECHNOLOGIES

* PALATIN TECHNOLOGIES INC

VYLEESI (AUTOINJECTOR), BREMELANOTIDE ACETATE

PANACEA

* PANACEA BIOTEC PHARMA 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

EVEROLIMUS, EVEROLIMUS

* PAR PHARMACEUTICAL INC

ALPRAZOLAM, ALPRAZOLAM
 AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
 CLONAZEPAM, CLONAZEPAM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROXYUREA, HYDROXYUREA
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 LAMOTRIGINE, LAMOTRIGINE
 MINOXIDIL, MINOXIDIL
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OXANDROLONE, OXANDROLONE
 PIMOZIDE, PIMOZIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 URSODIOL, URSODIOL

PAR PHARM INC

* PAR PHARMACEUTICAL INC

ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 AMBRISENTAN, AMBRISENTAN
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 BOSENTAN, BOSENTAN
 CHLORZOXAZONE, CHLORZOXAZONE
 COLCHICINE, COLCHICINE
 DEXLANSOPRAZOLE, DEXLANSOPRAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PAR PHARMACEUTICAL INC
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 ITRACONAZOLE, ITRACONAZOLE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 PENICILLAMINE, PENICILLAMINE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PRAZIQUANTEL, PRAZIQUANTEL
 SAPROPTERIN DIHYDROCHLORIDE, SAPROPTERIN DIHYDROCHLORIDE
 TOLCAPONE, TOLCAPONE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 VIGABATRIN, VIGABATRIN
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

PAR STERILE PRODUCTS

* PAR STERILE PRODUCTS LLC
 ADRENALIN, EPINEPHRINE
 ARGATROBAN, ARGATROBAN
 BREVITAL SODIUM, METHOHEXITAL SODIUM
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CALCITONIN-SALMON, CALCITONIN SALMON
 COLY-MYCIN M, COLISTIMETHATE SODIUM
 CORPHEDRA, EPHEDRINE SULFATE
 DANTRIUM, DANTROLENE SODIUM
 DELESTROGEN, ESTRADIOL VALERATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 KETALAR, KETAMINE HYDROCHLORIDE
 MICAFUNGIN, MICAFUNGIN SODIUM
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PITOCIN, OXYTOCIN
 TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
 TREPROSTINIL, TREPROSTINIL
 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
 NARDIL, PHENELZINE SULFATE
 ZARONTIN, ETHOSUXIMIDE

PARKE-DAVIS

* PARKE-DAVIS DIVISION OF PFIZER INC
 ZARONTIN, ETHOSUXIMIDE

PERRIGO

* PERRIGO CO

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PERRIGO CO
 MINOXIDIL, MINOXIDIL (OTC)
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)

* PERRIGO LLC
 DESLORATADINE, DESLORATADINE

PERRIGO NEW YORK

* PERRIGO NEW YORK INC
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 PERMETHRIN, PERMETHRIN (OTC)

PERRIGO PHARMA INTL

* PERRIGO PHARMA INTERNATIONAL DAC
 DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
 LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 LORATADINE, LORATADINE (OTC)

* PERRIGO PHARMA INTERNATIONAL DESIGNATED ACTIVITY CO
 LORATADINE, LORATADINE (OTC)
 PREVACID 24 HR, LANSOPRAZOLE (OTC)

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)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 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, IBUPROFEN (OTC)
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)

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
 PRISTIQ, DESVENLAFAXINE SUCCINATE
 RAPAMUNE, SIROLIMUS
 TORISEL, TEMSIROLIMUS
 TYGACIL, TIGECYCLINE
 VFEND, VORICONAZOLE
 XALKORI, CRIZOTINIB
 XELJANZ, TOFACITINIB CITRATE

PFIZER

* PFIZER CENTRAL RESEARCH

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** P **

* PFIZER CENTRAL RESEARCH
 DIFLUCAN, FLUCONAZOLE
 ZITHROMAX, AZITHROMYCIN

* PFIZER CHEMICALS DIV PFIZER INC
 DIFLUCAN, FLUCONAZOLE
 ZITHROMAX, AZITHROMYCIN

* PFIZER INC
 ALDACTAZIDE, HYDROCHLOROTHIAZIDE
 ALDACTONE, SPIRONOLACTONE
 AROMASIN, EXEMESTANE
 ARTHROTEC, DICLOFENAC SODIUM
 AZULFIDINE EN-TABS, SULFASALAZINE
 AZULFIDINE, SULFASALAZINE
 CALAN SR, VERAPAMIL HYDROCHLORIDE
 CAVERJECT IMPULSE, ALPROSTADIL
 CAVERJECT, ALPROSTADIL
 CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLEOCIN T, CLINDAMYCIN PHOSPHATE
 CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLEOCIN, CLINDAMYCIN PHOSPHATE
 COLESTID, COLESTIPOL HYDROCHLORIDE
 CORVERT, IBUTILIDE FUMARATE
 CYKLOKAPRON, TRANEXAMIC ACID
 CYTOTEC, MISOPROSTOL
 DAURISMO, GLASDEGIB MALEATE
 DAYPRO, OXAPROZIN
 DEPO-ESTRADIOL, ESTRADIOL CYPIONATE
 DEPO-MEDROL, METHYLPREDNISOLONE ACETATE
 DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE
 DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE
 DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 ESTRING, ESTRADIOL
 FLAGYL, METRONIDAZOLE
 FLAVORED COLESTID, COLESTIPOL HYDROCHLORIDE
 FRAGMIN, DALTEPARIN SODIUM
 GLUCOTROL XL, GLIPIZIDE
 GLUCOTROL, GLIPIZIDE
 GLYNASE, GLYBURIDE
 GLYSET, MIGLITOL
 HALCION, TRIAZOLAM
 HEMABATE, CARBOPROST TROMETHAMINE
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 IBRANCE, PALBOCICLIB
 IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE
 LINCOCIN, LINCOMYCIN HYDROCHLORIDE
 LOMOTIL, ATROPINE SULFATE
 LORBRENA, LORLATINIB
 MEDROL, METHYLPREDNISOLONE
 MERREM, MEROPENEM
 MYCOBUTIN, RIFABUTIN
 NICOTROL, NICOTINE
 NORPACE CR, DISOPYRAMIDE PHOSPHATE
 NORPACE, DISOPYRAMIDE PHOSPHATE
 OGEN 5, ESTROPIPATE
 PREPIDIL, DINOPROSTONE
 PROCARDIA, NIFEDIPINE
 PROSTIN VR PEDIATRIC, ALPROSTADIL
 PROVERA, MEDROXYPROGESTERONE ACETATE
 SONATA, ZALEPLON
 SYNAREL, NAFARELIN ACETATE
 TALZENNA, TALAZOPARIB TOSYLATE
 TESSALON, BENZONATATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

- * PFIZER INC
 - TOVIAZ, FESOTERODINE FUMARATE
 - UNASYN, AMPICILLIN SODIUM
 - VIZIMPRO, DACOMITINIB
 - XELJANZ XR, TOFACITINIB CITRATE
 - XELJANZ, TOFACITINIB CITRATE
 - ZITHROMAX, AZITHROMYCIN
 - ZYVOX, LINEZOLID
- * PFIZER LABORATORIES DIV PFIZER INC
 - 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 PRODUCTION CORP LTD
 - TIKOSYN, DOFETILIDE

PFIZER INC

- * PFIZER INC
 - CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
 - ELLENC, EPIRUBICIN HYDROCHLORIDE
 - NICOTROL, NICOTINE

PFIZER PHARMS

- * PFIZER PHARMACEUTICALS LTD
 - ACCUPRIL, QUINAPRIL HYDROCHLORIDE
 - ACCURETIC, HYDROCHLOROTHIAZIDE
 - LOPID, GEMFIBROZIL

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
 - CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 - 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
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 - PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 - PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 - PREDNISOLONE, PREDNISOLONE
 - PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 - PROMETHAZINE WITH CODEINE, CODEINE PHOSPHATE
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PHARMACEUTICAL ASSOCIATES INC
 RISPERIDONE, RISPERIDONE
 SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE
 THEOPHYLLINE, THEOPHYLLINE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 VALPROIC ACID, VALPROIC ACID
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

PHARM SOURCING

* PHARMACEUTICAL SOURCING PARTNERS INC
 MESALAMINE, MESALAMINE

PHARMA LIFE

* PHARMA LIFE SCIENCES LLC
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

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

* PHARMACIA AND UPJOHN CO LLC
 CADUET, AMLODIPINE BESYLATE
 DILANTIN, PHENYTOIN

PHARMACIA AND UPJOHN

* PHARMACIA AND UPJOHN CO
 CORTEF, HYDROCORTISONE
 EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM
 R-GENE 10, ARGININE HYDROCHLORIDE
 SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE
 SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE

PHARMACOSMOS AS

* PHARMACOSMOS AS
 MONOFERRIC, FERRIC DERISOMALTOSE

PHARMACYCLICS INC

* PHARMACYCLICS INC
 IMBRUVICA, IBRUTINIB

PHARMADAX INC

* PHARMADAX INC
 ENTECAVIR, ENTECAVIR
 LEVETIRACETAM, LEVETIRACETAM
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

PHARMALOGIC HLDGS

* PHARMALOGIC HOLDINGS CORP
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

PHARMASCIENCE INC

* PHARMASCIENCE INC
 BUSULFAN, BUSULFAN
 DECITABINE, DECITABINE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM

PHARMATHEN

* PHARMATHEN SA
 DIMETHYL FUMARATE, DIMETHYL FUMARATE

PHARMAXIS LTD

* PHARMAXIS LTD
 ARIDOL KIT, MANNITOL

PHOTOCURE ASA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PHOTOCURE ASA
 CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE

PIERRE FABRE DERMA

* PIERRE FABRE DERMATOLOGIE
 HEMANGEOL, PROPRANOLOL HYDROCHLORIDE

PIERREL

* PIERREL S.P.A.
 ORABLOC, ARTICAIN HYDROCHLORIDE

PINNACLE BIOLGS

* PINNACLE BIOLOGICS INC
 PHOTOFRIN, PORFIMER SODIUM

PIRAMAL CRITICAL

* PIRAMAL CRITICAL CARE INC
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 GABLOFEN, BACLOFEN
 GLYCOPYRROLATE, GLYCOPYRROLATE
 ISOFLURANE, ISOFLURANE
 MITIGO, MORPHINE SULFATE
 OXACILLIN SODIUM, OXACILLIN SODIUM
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 SOJOURN, SEVOFLURANE

* PIRAMAL CRITICAL CARE LTD
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM

PIRAMAL HLTHCARE UK

* PIRAMAL HEALTHCARE UK LTD
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 DEFERASIROX, DEFERASIROX
 TETRABENAZINE, TETRABENAZINE

PIRAMAL PHARMA

* PIRAMAL PHARMA LTD
 ISOFLURANE, ISOFLURANE

PLD ACQUISITIONS

* PLD ACQUISITIONS LLC DBA AVEMA PHARMA SOLUTIONS
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)

PLD ACQUISITIONS LLC

* PLD ACQUISITIONS LLC
 LORATADINE, LORATADINE (OTC)
 ZOLMITRIPTAN, ZOLMITRIPTAN

PLIVA

* PLIVA INC
 AZITHROMYCIN, AZITHROMYCIN
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 CIMETIDINE, CIMETIDINE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM

PLIVA HRVATSKA DOO

* PLIVA HRVATSKA DOO
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE

PLIVA PHARM IND

* PLIVA PHARMACEUTICAL INDUSTRY INC
 TORSEMIDE, TORSEMIDE

PLX PHARMA

* PLX PHARMA INC
 VAZALORE, ASPIRIN (OTC)

POHL BOSKAMP

* POHL BOSKAMP

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* POHL BOSKAMP

NITROLINGUAL PUMPSPRAY, NITROGLYCERIN

POLYGEN PHARMS

* POLYGEN PHARMACEUTICALS INC

AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE

POWDER PHARMS

* POWDER PHARMACEUTICALS INC

ZINGO, LIDOCAINE HYDROCHLORIDE

PRASCO

* PRASCO LLC DBA PRASCO LABORATORIES

EPLERENONE, EPLERENONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

PRASCO LABS LLC

* PRASCO LABORATORIES LLC

ESTRADIOL, ESTRADIOL

PRAXAIR DISTRIBUTION

* PRAXAIR DISTRIBUTION INC

NOXIVENT, NITRIC OXIDE

PRAXGEN

* PRAXGEN PHARMACEUTICALS LLC

FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 RANOLAZINE, RANOLAZINE

PRECISION DERMAT

* PRECISION DERMATOLOGY INC

LOCOID LIPOCREAM, HYDROCORTISONE BUTYRATE
 LOCOID, HYDROCORTISONE BUTYRATE

PRECISION DOSE INC

* PRECISION DOSE INC

PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE

PRECISION NUCLEAR

* PRECISION NUCLEAR LLC

AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

PRIMUS PHARMS

* PRIMUS PHARMACEUTICALS INC

IMPOYZ, CLOBETASOL PROPIONATE
 SERNIVO, BETAMETHASONE DIPROPIONATE

PRINSTON INC

* PRINSTON PHARMACEUTICAL INC

ACYCLOVIR, ACYCLOVIR
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 ARIPIPRAZOLE, ARIPIPRAZOLE
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CAPTOPRIL, CAPTOPRIL
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PRINSTON PHARMACEUTICAL INC
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 FUROSEMIDE, FUROSEMIDE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 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
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NEVIRAPINE, NEVIRAPINE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 PAROXETINE MESYLATE, PAROXETINE MESYLATE
 PAROXETINE, PAROXETINE HYDROCHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SILODOSIN, SILODOSIN
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMAZEPAM, TEMAZEPAM
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 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
 PYLARIFY, PIFLUFOLASTAT F-18

PROPEL PHARMA

* PROPEL PHARMA CORP
 VIGABATRIN, VIGABATRIN

PROVELL

* PROVELL PHARMACEUTICALS LLC
 EUTHYROX, LEVOTHYROXINE SODIUM **

PROVENSIS

* PROVENSIS LTD
 VARITHENA, POLIDOCANOL

PROVEPHARM SAS

* PROVEPHARM SAS
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PROVAYBLUE, METHYLENE BLUE
 TRANEXAMIC ACID, TRANEXAMIC ACID

PTC THERAP

* PTC THERAPEUTICS INC
 EMFLAZA, DEFLAZACORT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******PTS CONSULTING**

* PTS CONSULTING LLC
MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE

PULMOFLOW INC

* PULMOFLOW INC
KITABIS PAK, TOBRAMYCIN

PUMA BIOTECH

* PUMA BIOTECHNOLOGY INC
NERLYNX, NERATINIB MALEATE

PUNISKA

* PUNISKA HEALTHCARE PRIVATE LTD
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
ZOLEDRONIC ACID, ZOLEDRONIC ACID

PURACAP PHARM

* PURACAP PHARMACEUTICAL LLC
MELOXICAM, MELOXICAM

PURACAP PHARM LLC

* PURACAP PHARMACEUTICAL LLC
BENZONATATE, BENZONATATE
ERGOCALCIFEROL, ERGOCALCIFEROL
ETHOSUXIMIDE, ETHOSUXIMIDE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE

PURDUE

* PURDUE GMP CENTER LLC
SEROMYCIN, CYCLOSERINE

PURDUE PHARMA LP

* PURDUE PHARMA LP
ADHANSIA XR, METHYLPHENIDATE HYDROCHLORIDE
BUTRANS, BUPRENORPHINE
HYSINGLA ER, HYDROCODONE BITARTRATE
MS CONTIN, MORPHINE SULFATE
OXYCONTIN, OXYCODONE HYDROCHLORIDE

**** Q ******Q BIOMED**

* Q BIOMED INC
METASTRON, STRONTIUM CHLORIDE SR-89
STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE SR-89

QILU

* QILU PHARMACEUTICAL CO LTD
ABIRATERONE ACETATE, ABIRATERONE ACETATE
CEFAZOLIN SODIUM, CEFZAZOLIN SODIUM
CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
CEFTRIAZONE, CEFTRIAZONE SODIUM
EXEMESTANE, EXEMESTANE
OLANZAPINE, OLANZAPINE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
TENOFIVIR DISOPROXIL FUMARATE, TENOFIVIR DISOPROXIL FUMARATE

QILU PHARM HAINAN

* QILU PHARMACEUTICAL HAINAN CO LTD
DAPTOMYCIN, DAPTOMYCIN
DECITABINE, DECITABINE
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
OXALIPLATIN, OXALIPLATIN
PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
TADALAFIL, TADALAFIL

QINGDAO BAHEAL PHARM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Q ****

* QINGDAO BAHEAL PHARMACEUTICAL CO LTD
 BENZONATATE, BENZONATATE
 CELECOXIB, CELECOXIB
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

QOL MEDCL

* QOL MEDICAL LLC
 ETHAMOLIN, ETHANOLAMINE OLEATE

QUAGEN

* QUAGEN PHARMACEUTICALS LLC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 LIDOCAINE, LIDOCAINE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

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

RADIOMEDIX

* RADIOMEDIX INC
 DETECTNET, COPPER DOTATATE CU-64

RADIUS HEALTH INC

* RADIUS HEALTH INC
 TYMLOS, ABALOPARATIDE

RANBAXY

* RANBAXY SIGNATURE LLC
 RIOMET, METFORMIN HYDROCHLORIDE

RB HLTH

* RB HEALTH US LLC
 DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)
 MUCINEX D, GUAIFENESIN (OTC)
 MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 MUCINEX, GUAIFENESIN (OTC)

RECORDATI RARE

* RECORDATI RARE DISEASES INC
 CARBAGLU, CARGLUMIC ACID
 CHEMET, SUCCIMER
 COSMEGEN, DACTINOMYCIN
 CYSTADANE, BETAINE
 CYSTADROPS, CYSTEAMINE HYDROCHLORIDE
 DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
 ISTURISA, OSILODROSTAT PHOSPHATE
 NEOPROFEN, IBUPROFEN LYSINE
 SIGNIFOR LAR KIT, PASIREOTIDE PAMOATE
 SIGNIFOR, PASIREOTIDE DIASPARTATE
 TRANXENE, CLORAZEPATE DIPOTASSIUM

RECRO GAINESVILLE

* RECRO GAINESVILLE LLC
 VERELAN PM, VERAPAMIL HYDROCHLORIDE
 VERELAN, VERAPAMIL HYDROCHLORIDE

REDHILL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

* REDHILL BIOPHARMA INC
AEMCOLO, RIFAMYCIN SODIUM
MOVANTIK, NALOXEGOL OXALATE

* REDHILL BIOPHARMA LTD
TALICIA, AMOXICILLIN

RELIANCE LIFE

* RELIANCE LIFE SCIENCES PVT LTD
CAPECITABINE, CAPECITABINE

REMPEX

* REMPEX PHARMACEUTICALS INC A WHOLLY OWNED SUB OF MELINTA THERAPEUTICS LLC
MINOCIN, MINOCYCLINE HYDROCHLORIDE
VABOMERE, MEROPENEM

RENATA

* RENATA LTD
RISPERIDONE, RISPERIDONE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

RENEW PHARMS

* RENEW PHARMACEUTICALS LTD
INDOCYANINE GREEN, INDOCYANINE GREEN

RESILIA PHARMS

* RESILIA PHARMACEUTICALS INC
ECOZA, ECONAZOLE NITRATE

REYOUNG

* REYOUNG CORPORATION
FENOFIBRATE (MICRONIZED), FENOFIBRATE
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SILDENAFIL CITRATE, SILDENAFIL CITRATE
* REYOUNG PHARMACEUTICALS CO LTD
NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE

RHODES PHARMS

* RHODES PHARMACEUTICALS LP
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
APTENSIO XR, METHYLPHENIDATE HYDROCHLORIDE
BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DILAUDID, HYDROMORPHONE HYDROCHLORIDE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FENOFIBRATE, FENOFIBRATE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
LIDOCAINE, LIDOCAINE
MORPHINE SULFATE, MORPHINE SULFATE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
THEOPHYLLINE, THEOPHYLLINE

RHYTHM

* RHYTHM PHARMACEUTICALS INC
IMCIVREE, SETMELANOTIDE ACETATE

RICONPHARMA LLC

* RICONPHARMA LLC
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
SCOPOLAMINE, SCOPOLAMINE

RIGEL PHARMS INC

* RIGEL PHARMACEUTICALS INC
TAVALISSE, FOSTAMATINIB DISODIUM

RISING

* RISING PHARMA HOLDINGS INC
ABIRATERONE ACETATE, ABIRATERONE ACETATE
ACETIC ACID, ACETIC ACID, GLACIAL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

- * RISING PHARMA HOLDINGS INC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 AZATHIOPRINE, AZATHIOPRINE
 BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CICLOPIROX, CICLOPIROX
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DAPSONE, DAPSONE
 DESOXIMETASONE, DESOXIMETASONE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 EXEMESTANE, EXEMESTANE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCORTISONE, HYDROCORTISONE
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED), AMOXICILLIN
 LEVOCARNITINE, LEVOCARNITINE
 LEVOFLOXACIN, LEVOFLOXACIN
 METHIMAZOLE, METHIMAZOLE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 NADOLOL, NADOLOL
 NATEGLINIDE, NATEGLINIDE
 NITAZOXANIDE, NITAZOXANIDE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 PARICALCITOL, PARICALCITOL
 PERPHENAZINE, PERPHENAZINE
 PHENYTOIN, PHENYTOIN
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PREGABALIN, PREGABALIN
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RISPERIDONE, RISPERIDONE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SULFAMYLOL, MAFENIDE ACETATE
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TOREMIFENE CITRATE, TOREMIFENE CITRATE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZILEUTON, ZILEUTON
- * RISING PHARMACEUTICALS
 URSODIOL, URSODIOL

RISING PHARMA

- * RISING PHARMA HOLDING INC
 AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CAPECITABINE, CAPECITABINE
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CYTARABINE, CYTARABINE
 DIGOXIN, DIGOXIN
 FLUMAZENIL, FLUMAZENIL
 FLUNISOLIDE, FLUNISOLIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ******RK PHARMA**

* RK PHARMA INC
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 MITOMYCIN, MITOMYCIN
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 PALIPERIDONE, PALIPERIDONE

ROCHE PALO

* ROCHE PALO ALTO LLC
 CELLCEPT, MYCOPHENOLATE MOFETIL
 CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE

ROCKWELL MEDICAL INC

* ROCKWELL MEDICAL INC
 TRIFERIC AVNU, FERRIC PYROPHOSPHATE CITRATE
 TRIFERIC, FERRIC PYROPHOSPHATE CITRATE

ROMARK

* ROMARK LABORATORIES
 ALINIA, NITAZOXANIDE

ROMEG

* ROMEG THERAPEUTICS LLC
 GLOPERBA, COLCHICINE

ROXANE

* ROXANE LABORATORIES INC
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE

RUBICON

* RUBICON RESEARCH PRIVATE LTD
 ACETAZOLAMIDE, ACETAZOLAMIDE
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BACLOFEN, BACLOFEN
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARVEDILOL, CARVEDILOL
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 GABAPENTIN, GABAPENTIN
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LAMOTRIGINE, LAMOTRIGINE
 LORATADINE, LORATADINE (OTC)
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 NITROGLYCERIN, NITROGLYCERIN
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

RVL PHARMS

* RVL PHARMACEUTICALS INC
 UPNEEQ, OXYMETAZOLINE HYDROCHLORIDE

**** S ******SAGE PRODS**

* SAGE PRODUCTS INC
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

SAGE THERAP

* SAGE THERAPEUTICS INC
 ZULRESSO, BREXANOLONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******SAGENT PHARMS**

* SAGENT PHARMACEUTICALS INC
 CAFFEINE CITRATE, CAFFEINE CITRATE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 FLUMAZENIL, FLUMAZENIL
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEVETIRACETAM, LEVETIRACETAM
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXACILLIN SODIUM, OXACILLIN SODIUM
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

SAGENT PHARMS INC

* SAGENT PHARMACEUTICALS INC
 ACETYLCYSTEINE, ACETYLCYSTEINE
 AMIKACIN SULFATE, AMIKACIN SULFATE
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 DAPTOMYCIN, DAPTOMYCIN
 DECITABINE, DECITABINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 EPTIFIBATIDE, EPTIFIBATIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUOROURACIL, FLUOROURACIL
 FULVESTRANT, FULVESTRANT
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLYDO, LIDOCAINE HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 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
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OXYTOCIN, OXYTOCIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PROPOFOL, PROPOFOL
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

SALIX

* SALIX PHARMACEUTICALS INC
 FENOGLIDE, FENOFIBRATE
 PLENVU, ASCORBIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SALIX PHARMACEUTICALS INC
 RELISTOR, METHYLNALTREXONE BROMIDE
 TRULANCE, PLECANATIDE
 UCERIS, BUDESONIDE
 ZEGERID, OMEPRAZOLE

SALIX PHARMS

* SALIX PHARMACEUTICALS INC
 ANUSOL HC, HYDROCORTISONE
 DIURIL, CHLOROTHIAZIDE
 MOVIPREP, ASCORBIC ACID
 OSMOPREP, SODIUM PHOSPHATE, DIBASIC, ANHYDROUS
 RELISTOR, METHYLNALTREXONE BROMIDE
 XIFAXAN, RIFAXIMIN

SAMSON MEDCL

* SAMSON MEDICAL TECHNOLOGIES LLC
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFEPIME HYDROCHLORIDE IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
 CEFOXITIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE

SANDOZ

* SANDOZ
 DOCETAXEL, DOCETAXEL

* SANDOZ INC
 ALPRAZOLAM, ALPRAZOLAM
 AMANTADINE HYDROCHLORIDE, AMANTADINE 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
 AZITHROMYCIN, AZITHROMYCIN
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BUMETANIDE, BUMETANIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARVEDILOL, CARVEDILOL
 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
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SANDOZ INC
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 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
 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
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OMEPRAZOLE, OMEPRAZOLE
 ONDANSETRON, ONDANSETRON
 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, PIOGLITAZONE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 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
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SANDOZ INC

VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

SANDOZ CANADA INC

* SANDOZ CANADA INC

INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE
 INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID
 INFUVITE PEDIATRIC, ASCORBIC ACID

SANDOZ INC

* SANDOZ INC

ACETAMINOPHEN, ACETAMINOPHEN
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 ANECTINE, SUCCINYLCHOLINE CHLORIDE
 ANGIOMAX, BIVALIRUDIN
 ARISTOSPAN, TRIAMCINOLONE HEXACETONIDE
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 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
 CARBOPLATIN, CARBOPLATIN
 CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
 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
 CYANOCOBALAMIN, CYANOCOBALAMIN
 DECITABINE, DECITABINE
 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
 FERUMOXYTOL, FERUMOXYTOL
 FLUMAZENIL, FLUMAZENIL
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FULVESTRANT, FULVESTRANT
 GATIFLOXACIN, GATIFLOXACIN
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GLATOPA, GLATIRAMER ACETATE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 ISONIAZID, ISONIAZID
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED), AMOXICILLIN
 LATANOPROST, LATANOPROST
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 LINEZOLID, LINEZOLID
 MAXITROL, DEXAMETHASONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SANDOZ INC
 MESALAMINE, MESALAMINE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYDRIACYL, TROPICAMIDE
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 NEVIRAPINE, NEVIRAPINE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 QOLIANA, BRIMONIDINE TARTRATE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 REGONOL, PYRIDOSTIGMINE BROMIDE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILODOSIN, SILODOSIN
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TELMISARTAN, TELMISARTAN
 TERIFLUNOMIDE, TERIFLUNOMIDE
 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

SANOFI

* SANOFI AVENTIS US LLC
 FEXINIDAZOLE, FEXINIDAZOLE
 FLOMAX, TAMSULOSIN HYDROCHLORIDE
 XYZAL ALLERGY 24HR, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 * SANOFI GENZYME
 HECTOROL, DOXERCALCIFEROL
 RENVELA, SEVELAMER CARBONATE

SANOFI AVENTIS US

* SANOFI AVENTIS US INC
 JEVTANA KIT, CABAZITAXEL
 * SANOFI AVENTIS US LLC
 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
 ARAVA, LEFLUNOMIDE
 AUBAGIO, TERIFLUNOMIDE
 AVALIDE, HYDROCHLOROTHIAZIDE
 AVAPRO, IRBESARTAN
 DIABETA, GLYBURIDE
 ELOXATIN, OXALIPLATIN
 FERRLECIT, FERRIC OXYHYDROXIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SANOFI AVENTIS US LLC
 LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 LOVENOX, ENOXAPARIN SODIUM
 MULTAQ, DRONEDARONE HYDROCHLORIDE
 NASACORT ALLERGY 24 HOUR, TRIAMCINOLONE ACETONIDE (OTC)
 NASACORT AQ, TRIAMCINOLONE ACETONIDE
 PLAVIX, CLOPIDOGREL BISULFATE
 PRIFTIN, RIFAPENTINE
 PRIMAQUINE, PRIMAQUINE PHOSPHATE
 RIFADIN, RIFAMPIN
 TAXOTERE, DOCETAXEL

SANTARUS INC

* SANTARUS INC
 GLUMETZA, METFORMIN HYDROCHLORIDE

SANTEN

* SANTEN INC
 VERKAZIA, CYCLOSPORINE

SAOL THERAPS RES LTD

* SAOL THERAPEUTICS RESEARCH LTD
 LIORESAL, BACLOFEN
 LYVISPAH, BACLOFEN

SAPTALIS PHARMS

* SAPTALIS PHARMACEUTICALS LLC
 ACETIC ACID, ACETIC ACID, GLACIAL
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 ERYTHRA-DERM, ERYTHROMYCIN
 LEVOCARNITINE, LEVOCARNITINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

SAREPTA THERAPS INC

* SAREPTA THERAPEUTICS INC
 AMONDYS 45, CASIMERSSEN
 EXONDYS 51, ETEPLIRSEN
 VYONDYS 53, GOLODIRSEN

SARFE PHARMS

* SARFEZ PHARMACEUTICALS INC
 SOAANZ, TORSEMIDE

SAVIOR LIFETEC CORP

* SAVIOR LIFETEC CORP
 ERTAPENEM SODIUM, ERTAPENEM SODIUM
 MEROPENEM, MEROPENEM

SAWAI USA

* SAWAI USA INC
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM

SCHERING

* SCHERING CORP
 NOXAFIL, POSACONAZOLE

SCIARRA LABS

* SCIARRA LABORATORIES INC
 SCLEROSOL, TALC
 TALC, TALC

SCIECURE PHARMA INC

* SCIECURE PHARMA INC
 BUDESONIDE, BUDESONIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE

SCIEGEN PHARMS INC

* SCIEGEN PHARMACEUTICALS INC
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARISOPRODOL, CARISOPRODOL
 CELECOXIB, CELECOXIB

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** s ******* SCIEGEN PHARMACEUTICALS INC**

CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DROXIDOPA, DROXIDOPA
 ETHACRYNIC ACID, ETHACRYNIC ACID
 EZETIMIBE, EZETIMIBE
 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
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 METAXALONE, METAXALONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NABUMETONE, NABUMETONE
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NAPROXEN, NAPROXEN
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PREGABALIN, PREGABALIN
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 VALSARTAN, VALSARTAN

SCILEX PHARMS INC*** SCILEX PHARMACEUTICALS**

ZTLIDO, LIDOCAINE

SCINOPHARM TAIWAN*** SCINOPHARM TAIWAN LTD**

FONDAPARINUX SODIUM, FONDAPARINUX SODIUM

SCYNEXIS*** SCYNEXIS INC**

BREXAFEMME, IBREXAFUNGERP CITRATE

SEAGEN*** SEAGEN INC**

TUKYSA, TUCATINIB

SEBELA IRELAND LTD*** SEBELA IRELAND LTD**

BRISDELLE, PAROXETINE MESYLATE
 IMURAN, AZATHIOPRINE
 LOTRONEX, ALOSETRON HYDROCHLORIDE
 MICORT-HC, HYDROCORTISONE ACETATE
 MOTOFEN, ATROPINE SULFATE
 NAFTIN, NAFTIFINE HYDROCHLORIDE
 PEXEVA, PAROXETINE MESYLATE
 PRAMOSONE, HYDROCORTISONE ACETATE
 RIDAURA, AURANOFIN

SECAN PHARMS*** SECAN PHARMACEUTICALS INC**

LEVETIRACETAM, LEVETIRACETAM

SECURA*** SECURA BIO INC**

COPIKTRA, DUVELISIB

SENORES PHARMS*** SENORES PHARMACEUTICALS INC**

AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CHLORZOAZONE, CHLORZOAZONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SENORES PHARMACEUTICALS INC
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 KETOCONAZOLE, KETOCONAZOLE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE

SENTYNL THERAPS INC

* SENTYNL THERAPEUTICS INC
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE

SEPTODONT

* SEPTODONT INC
 BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE

SEPTODONT HOLDING

* SEPTODONT HOLDING SAS
 ORAVERSE, 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

SERVIER

* SERVIER PHARMACEUTICALS LLC
 TIBSOVO, IVOSIDENIB

SETON PHARM

* SETON PHARMACEUTICAL LLC
 PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE

SETON PHARMS

* SETON PHARMACEUTICALS LLC
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE

SHANDONG

* SHANDONG NEW TIME PHARMACEUTICAL CO LTD
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

SHANDONG XINHUA

* SHANDONG XINHUA PHARMACEUTICAL CO LTD
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)

SHANGHAI HENGRUI

* SHANGHAI HENGRUI PHARMACEUTICAL CO LTD
 DESFLURANE, DESFLURANE
 SEVOFLURANE, SEVOFLURANE

SHENZHEN TECHDOW

* SHENZHEN TECHDOW PHARMACEUTICAL CO LTD
 HEPARIN SODIUM, HEPARIN SODIUM

SHERTECH LABS LLC

* SHERTECH LABORATORIES LLC
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

SHIELD TX

* SHIELD TX UK LTD
 ACCRUFER, FERRIC MALTOL

SHILPA

* SHILPA MEDICARE LTD
 BUSULFAN, BUSULFAN
 CAPECITABINE, CAPECITABINE
 DOCETAXEL, DOCETAXEL
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE

SHILPA MEDICARE

* SHILPA MEDICARE LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SHILPA MEDICARE LTD
AZACITIDINE, AZACITIDINE

SHIONOGI INC

* SHIONOGI INC
FETROJA, CEFIDEROCOL SULFATE TOSYLATE
MULPLETA, LUSUTROMBOPAG

SHIRE ORPHAN THERAP

* SHIRE ORPHAN THERAPIES LLC
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
AMBRISENTAN, AMBRISENTAN
AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
ASENAPINE MALEATE, ASENAPINE MALEATE
DISULFIRAM, DISULFIRAM
DOFETILIDE, DOFETILIDE
FLUCYTOSINE, FLUCYTOSINE
GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
SODIUM PHENYL BUTYRATE, SODIUM PHENYL BUTYRATE

SINOTHERAPEUTICS INC

* SINOTHERAPEUTICS INC
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
POSACONAZOLE, POSACONAZOLE
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE

SK LIFE

* SK LIFE SCIENCE INC
XCOPRI, CENOBAMATE

SKINMEDICA

* SKINMEDICA INC
VANIQA, EFLORNITHINE HYDROCHLORIDE

SKYEPHARMA AG

* SKYEPHARMA AG
TRIGLIDE, FENOFIBRATE

SLATE

* SLATE RUN PHARMACEUTICALS LLC
CILOSTAZOL, CILOSTAZOL
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DROXIDOPA, DROXIDOPA
METHOCARBAMOL, METHOCARBAMOL
PROMETHAZINE DM, DEXTROMETHORPHAN HYDROBROMIDE

SLAYBACK PHARMA LLC

* SLAYBACK PHARMA LLC
DEOXYCHOLIC ACID, DEOXYCHOLIC ACID
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE
ICATIBANT ACETATE, ICATIBANT ACETATE
MERZEE, ETHINYL ESTRADIOL
METARAMINOL BITARTRATE, METARAMINOL BITARTRATE

SOFGEN PHARMS

* SOFGEN PHARMACEUTICALS LLC
IBUPROFEN, IBUPROFEN (OTC)
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SOFGEN PHARMACEUTICALS LLC
PROGESTERONE, PROGESTERONE

SOFIE

* SOFIE CO DBA SOFIE
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

* 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

SOL-GEL TECHNOLOGIES

* SOL-GEL TECHNOLOGIES LTD
TWYNEO, BENZOYL PEROXIDE

SOLARIS PHARMA CORP

* SOLARIS PHARMA CORP
ACYCLOVIR, ACYCLOVIR
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
METRONIDAZOLE, METRONIDAZOLE

SOMERSET

* SOMERSET PHARMACEUTICALS INC
EMSAM, SELEGILINE

* SOMERSET THERAPEUTICS LLC
CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
LATANOPROST, LATANOPROST
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

SOMERSET THERAPS LLC

* SOMERSET THERAPEUTICS LLC
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
BIMATOPROST, BIMATOPROST
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
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
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
TOBRAMYCIN, TOBRAMYCIN
TROPICAMIDE, TROPICAMIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

SOMMER PHARMS II LLC

* SOMMER PHARMACEUTICALS II LLC
TECHNETIUM TC99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT

SPECGX LLC

* SPECGX LLC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE
ANEXSIA 5/325, ACETAMINOPHEN
ANEXSIA 7.5/325, ACETAMINOPHEN
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SPECGX LLC**

FENTANYL CITRATE, FENTANYL CITRATE
 FENTANYL-100, FENTANYL
 FENTANYL-12, FENTANYL
 FENTANYL-25, FENTANYL
 FENTANYL-37, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-62, FENTANYL
 FENTANYL-75, FENTANYL
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 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

SPECTRA MDCL DEVICES

* SPECTRA MEDICAL DEVICES INC
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE

SPIL

* SUN PHARMA INDUSTRIES LTD
 AMPHOTERICIN B, AMPHOTERICIN B
 KAPSPARGO SPRINKLE, METOPROLOL SUCCINATE
 NIFEDIPINE, NIFEDIPINE
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)

SPROUT PHARMS

* SPROUT PHARMACEUTICALS INC
 ADDYI, FLIBANSERIN

SQUARE PHARMS

* SQUARE PHARMACEUTICALS LTD
 ACYCLOVIR, ACYCLOVIR
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 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, MERCAPTOPYRINE

STI PHARMA LLC

* STI PHARMA LLC
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 CARMUSTINE, CARMUSTINE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE
 THIOTEPA, THIOTEPA
 TRIACIN-C, CODEINE PHOSPHATE

STIEFEL

* STIEFEL LABORATORIES INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* STIEFEL LABORATORIES INC
DUAC, BENZOYL PEROXIDE

STRIDES PHARMA

* STRIDES PHARMA GLOBAL PTE LTD
ABACAVIR SULFATE, ABACAVIR SULFATE
ACARBOSE, ACARBOSE
ACCOLATE, ZAFIRLUKAST
ACETAZOLAMIDE, ACETAZOLAMIDE
ACYCLOVIR, ACYCLOVIR
ALBENDAZOLE, ALBENDAZOLE
ALPRAZOLAM, ALPRAZOLAM
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
BENZONATATE, BENZONATATE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
CALCITRIOL, CALCITRIOL
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
CLARITHROMYCIN, CLARITHROMYCIN
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DIAZEPAM, DIAZEPAM
DOFETILIDE, DOFETILIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DOXYCYCLINE, DOXYCYCLINE
DUTASTERIDE, DUTASTERIDE
EFAVIRENZ, EFAVIRENZ
EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
ERGOCALCIFEROL, ERGOCALCIFEROL
ETHACRYNIC ACID, ETHACRYNIC ACID
ETHOSUXIMIDE, ETHOSUXIMIDE
GABAPENTIN, GABAPENTIN
HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROCORTISONE, HYDROCORTISONE
IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
KETOCONAZOLE, KETOCONAZOLE
LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
LAMIVUDINE, LAMIVUDINE
LIDOCAINE, LIDOCAINE
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
MEGESTROL ACETATE, MEGESTROL ACETATE
MELOXICAM, MELOXICAM
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
METHOXSALEN, METHOXSALEN
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
METRONIDAZOLE, METRONIDAZOLE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
NATEGLINIDE, NATEGLINIDE
NEVIRAPINE, NEVIRAPINE
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
OLANZAPINE, OLANZAPINE
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* STRIDES PHARMA GLOBAL PTE LTD
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 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
 PREDNISONE, PREDNISONE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 ROFLUMILAST, ROFLUMILAST
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TACROLIMUS, TACROLIMUS
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TESTOSTERONE, TESTOSTERONE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 TORSEMIDE, TORSEMIDE
 TRANYLCPROMINE SULFATE, TRANYLCPROMINE SULFATE
 TRAVOPROST, TRAVOPROST
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 URSODIOL, URSODIOL
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VOSPIRE ER, ALBUTEROL SULFATE
 ZILEUTON, ZILEUTON

STRONGBRIDGE

* STRONGBRIDGE DUBLIN LTD
 RECORLEV, LEVOKETOCONAZOLE

SUCAMPO PHARMA LLC

* SUCAMPO PHARMA AMERICAS LLC
 AMITIZA, LUBIPROSTONE

SUN PHARM

* SUN PHARMACEUTICAL INDUSTRIES LTD
 ABSORICA LD, ISOTRETINOIN
 ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALPRAZOLAM, ALPRAZOLAM
 AMBRISENTAN, AMBRISENTAN
 AMIFOSTINE, AMIFOSTINE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 BOSENTAN, BOSENTAN
 BROMSITE, BROMFENAC SODIUM
 BUDESONIDE, BUDESONIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALCITRIOL, CALCITRIOL
 CAPECITABINE, CAPECITABINE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA
 CARBOPLATIN, CARBOPLATIN
 CEQUA, CYCLOSPORINE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUN PHARMACEUTICAL INDUSTRIES LTD
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DALFAMPRIDINE, DALFAMPRIDINE
 DECITABINE, DECITABINE
 DEFERASIROX, DEFERASIROX
 DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOCETAXEL, DOCETAXEL
 DOFETILIDE, DOFETILIDE
 DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
 DROXIDOPA, DROXIDOPA
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTACAPONE, ENTACAPONE
 EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 ESZOPICLONE, ESZOPICLONE
 EZALLOR SPRINKLE, ROSUVASTATIN CALCIUM
 FEBUXOSTAT, FEBUXOSTAT
 FENOFIBRATE, FENOFIBRATE
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
 FINASTERIDE, FINASTERIDE
 FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GANIRELIX ACETATE, GANIRELIX ACETATE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IMATINIB MESYLATE, IMATINIB MESYLATE
 INFUGEM, GEMCITABINE HYDROCHLORIDE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 LORATADINE, LORATADINE (OTC)
 LOTEFREDNOL ETABONATE, LOTEFREDNOL ETABONATE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 NIACIN, NIACIN
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 OPCICON ONE-STEP, LEVONORGESTREL (OTC)
 PALIPERIDONE, PALIPERIDONE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PREGABALIN, PREGABALIN
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RANOLAZINE, RANOLAZINE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** s ****

* SUN PHARMACEUTICAL INDUSTRIES LTD
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SUNITINIB MALATE, SUNITINIB MALATE
 TADALAFIL, TADALAFIL
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TETRABENAZINE, TETRABENAZINE
 TOBRAMYCIN, TOBRAMYCIN
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 WINLEVI, CLASCOTERONE
 XELPROS, LATANOPROST
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

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)
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLANZAPINE, OLANZAPINE
 ONDANSETRON, ONDANSETRON
 OXCARBAZEPINE, OXCARBAZEPINE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE

SUN PHARM INDS (IN)

* SUN PHARMACEUTICAL INDUSTRIES LTD
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ZONISAMIDE, ZONISAMIDE

SUN PHARM INDS INC

* SUN PHARMACEUTICAL INDUSTRIES INC
 ABSORICA, ISOTRETINOIN
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT
 CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT
 CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 CLONAZEPAM, CLONAZEPAM
 CLOZAPINE, CLOZAPINE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DIGOXIN, DIGOXIN
 ERGOCALCIFEROL, ERGOCALCIFEROL
 FLUMADINE, RIMANTADINE HYDROCHLORIDE
 GLIPIZIDE, GLIPIZIDE
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 HALOG, HALCINONIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 KENALOG, TRIAMCINOLONE ACETONIDE
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUN PHARMACEUTICAL INDUSTRIES INC
 ORTIKOS, BUDESONIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 REPAGLINIDE, REPAGLINIDE
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 TECHNETIUM TC-99M MEBROFENIN, TECHNETIUM TC-99M MEBROFENIN KIT
 TECHNETIUM TC99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

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
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 CARVEDILOL, CARVEDILOL
 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
 DOXYCYCLINE, DOXYCYCLINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FELODIPINE, FELODIPINE
 FENOFIBRATE, FENOFIBRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GLYCOPYRROLATE, GLYCOPYRROLATE
 LEVETIRACETAM, LEVETIRACETAM
 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
 ONDANSETRON, ONDANSETRON
 OXCARBAZEPINE, OXCARBAZEPINE
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 RILUZOLE, RILUZOLE
 RISPERIDONE, RISPERIDONE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TOPIRAMATE, TOPIRAMATE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALPROIC ACID, VALPROIC ACID

SUN PHARM INDUSTRIES

* SUN PHARMACEUTICAL INDUSTRIES INC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALLOPURINOL, ALLOPURINOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUN PHARMACEUTICAL INDUSTRIES INC
 BACTRIM DS, SULFAMETHOXAZOLE
 BACTRIM, SULFAMETHOXAZOLE
 CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
 CHLORTHALIDONE, CHLORTHALIDONE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ERGOLOID MESYLATES, ERGOLOID MESYLATES
 FELODIPINE, FELODIPINE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL, MINOXIDIL
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NYSTATIN, NYSTATIN
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PINDOLOL, PINDOLOL
 PREDNISONE, PREDNISONE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 QUALAQUIN, QUININE SULFATE
 QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
 SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SULINDAC, SULINDAC
 TEMAZEPAM, TEMAZEPAM
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 ULTRAVATE, HALOBETASOL PROPIONATE

SUN PHARMA GLOBAL

* SUN PHARMA GLOBAL FZE
 DRIZALMA SPRINKLE, DULOXETINE HYDROCHLORIDE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ODOMZO, SONIDEGIB PHOSPHATE
 YONSA, ABIRATERONE ACETATE

SUNNY

* SUNNY PHARMTECH INC
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 GLYBURIDE, GLYBURIDE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN

SUNOVION

* SUNOVION PHARMACEUTICALS INC
 BROVANA, ARFORMOTEROL TARTRATE
 XOPENEX HFA, LEVALBUTEROL TARTRATE

SUNOVION PHARMS INC

* SUNOVION PHARMACEUTICALS INC
 APTIOM, ESLICARBAZEPINE ACETATE
 KYNMOBI, APOMORPHINE HYDROCHLORIDE
 LATUDA, LURASIDONE HYDROCHLORIDE
 LUNESTA, ESZOPICLONE

SUNOVION RESP

* SUNOVION RESPIRATORY DEVELOPMENT INC
 LONHALA MAGNAIR KIT, GLYCOPYRROLATE

SUNSHINE

* SUNSHINE LAKE PHARMA CO LTD
 AZITHROMYCIN, AZITHROMYCIN
 CLARITHROMYCIN, CLARITHROMYCIN
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTACAPONE, ENTACAPONE
 FEBUXOSTAT, FEBUXOSTAT
 LINAGLIPTIN AND METFORMIN HYDROCHLORIDE, LINAGLIPTIN
 LINAGLIPTIN, LINAGLIPTIN
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUNSHINE LAKE PHARMA CO LTD
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 RANOLAZINE, RANOLAZINE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TADALAFIL, TADALAFIL

SUNSTAR AMERICAS

* SUNSTAR AMERICAS INC
 PAROEX, CHLORHEXIDINE GLUCONATE

SUPERNUS PHARMS

* SUPERNUS PHARMACEUTICALS INC
 OXTELLAR XR, OXCARBAZEPINE
 QELBREE, VILOXAZINE HYDROCHLORIDE
 TROKENDI XR, TOPIRAMATE

SUVEN PHARMS

* SUVEN PHARMACEUTICALS LTD
 CALCIUM ACETATE, CALCIUM ACETATE
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 MALATHION, MALATHION

SVC PHARMA

* SVC PHARMA LP
 DRONABINOL, DRONABINOL

SWEDISH ORPHAN

* SWEDISH ORPHAN BIOVITRUM AB PUBL
 ORFADIN, NITISINONE

SYNTHON PHARMS

* SYNTHON PHARMACEUTICALS INC
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE

**** T ******ACME LABS**

* THE ACME LABORATORIES LTD
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

GEN HOSP

* THE GENERAL HOSPITAL CORP
 AMMONIA N 13, AMMONIA N-13

METHODIST

* THE METHODIST HOSP RESEARCH INSTITUTE
 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

TAGI

* TAGI PHARMA INC
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE

TAIHO ONCOLOGY

* TAIHO ONCOLOGY INC
 LONSURF, TIPIRACIL HYDROCHLORIDE

TAKEDA PHARMS USA

* TAKEDA PHARMACEUTICALS USA INC
 ACTOPLUS MET, METFORMIN HYDROCHLORIDE
 ACTOS, PIOGLITAZONE HYDROCHLORIDE
 ADDERALL XR 10, AMPHETAMINE ASPARTATE
 ADDERALL XR 15, AMPHETAMINE ASPARTATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TAKEDA PHARMACEUTICALS USA INC
 ADDERALL XR 20, AMPHETAMINE ASPARTATE
 ADDERALL XR 25, AMPHETAMINE ASPARTATE
 ADDERALL XR 30, AMPHETAMINE ASPARTATE
 ADDERALL XR 5, AMPHETAMINE ASPARTATE
 AGRYLIN, ANAGRELIDE HYDROCHLORIDE
 ALUNBRIG, BRIGATINIB
 CARBATROL, CARBAMAZEPINE
 COLCRYS, COLCHICINE
 DEXILANT, DEXLANSOPRAZOLE
 DUETACT, GLIMEPIRIDE
 EXKIVITY, MOBOCERTINIB SUCCINATE
 FOSRENOL, LANTHANUM CARBONATE
 ICLUSIG, PONATINIB HYDROCHLORIDE
 INTUNIV, GUANFACINE HYDROCHLORIDE
 KAZANO, ALOGLIPTIN BENZOATE
 LIALDA, MESALAMINE
 LIVTENCITY, MARIBAVIR
 MOTEGRITY, PRUCALOPRIDE SUCCINATE
 MYDAYIS, AMPHETAMINE ASPARTATE
 NESINA, ALOGLIPTIN BENZOATE
 NINLARO, IXAZOMIB CITRATE
 OSENI, ALOGLIPTIN BENZOATE
 PENTASA, MESALAMINE
 PREVACID, LANSOPRAZOLE
 ROZEREM, RAMELTEON
 TRINTELLIX, VORTIOXETINE HYDROBROMIDE
 ULORIC, FEBUXOSTAT
 VELCADE, BORTEZOMIB
 VYVANSE, LISDEXAMFETAMINE DIMESYLATE

TAMARANG

* TAMARANG SA
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

TARO

* TARO PHARMACEUTICAL INDUSTRIES LTD
 ACETAZOLAMIDE, ACETAZOLAMIDE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 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)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOCORTOLONE PIVALATE, CLOCORTOLONE PIVALATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 DESONIDE, DESONIDE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ETODOLAC, ETODOLAC
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 FELBAMATE, FELBAMATE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOROURACIL, FLUOROURACIL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TARO PHARMACEUTICAL INDUSTRIES LTD
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 IMIQUIMOD, IMIQUIMOD
 IVERMECTIN, IVERMECTIN (OTC)
 KETOCONAZOLE, KETOCONAZOLE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LORATADINE, LORATADINE (OTC)
 MELOXICAM, MELOXICAM
 METRONIDAZOLE, METRONIDAZOLE
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 PHENYTOIN, PHENYTOIN
 TERIL, CARBAMAZEPINE
 WARFARIN SODIUM, WARFARIN SODIUM
- * 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
 AZELAIC ACID, AZELAIC ACID
 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
 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
 MUPIROCIN, MUPIROCIN
 MUPIROCIN, MUPIROCIN CALCIUM
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 PHENYTOIN, PHENYTOIN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TARO PHARMACEUTICALS USA INC
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE (OTC)
 TERCONAZOLE, TERCONAZOLE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)
 U-CORT, HYDROCORTISONE ACETATE

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
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 LAMOTRIGINE, LAMOTRIGINE

TARO PHARM INDS LTD

* TARO PHARMACEUTICAL INDUSTRIES LTD
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 DEFERIPRONE, DEFERIPRONE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 HYDROCORTISONE, HYDROCORTISONE
 INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TOPICORT, DESOXIMETASONE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

TARO PHARMS

* TARO PHARMACEUTICALS INC
 BUTENAFINE HYDROCHLORIDE, BUTENAFINE HYDROCHLORIDE (OTC)
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DAPSONE, DAPSONE
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 OXICONAZOLE NITRATE, OXICONAZOLE NITRATE
 PLIAGLIS, LIDOCAINE
 TAVABOROLE, TAVABOROLE
 TAZAROTENE, TAZAROTENE
 TOPICORT, DESOXIMETASONE
 TRETINOIN, TRETINOIN

TASMAN PHARMA

* TASMAN PHARMA INC
 CLOTRIMAZOLE, CLOTRIMAZOLE
 DROXIDOPA, DROXIDOPA
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 KETOCONAZOLE, KETOCONAZOLE
 VERSACLOZ, CLOZAPINE

TCG FLUENT PHARMA

* TCG FLUENT PHARMA INVESTORS LP
 FLOLIPID, SIMVASTATIN

TEIKOKU PHARMA USA

* TEIKOKU PHARMA USA INC
 LIDODERM, LIDOCAINE

TELIGENT

* TELIGENT OU
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 FORTAZ, CEFTAZIDIME
 ZANTAC, RANITIDINE HYDROCHLORIDE
 * TELIGENT PHARMA INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TELIGENT PHARMA INC
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CEFOTAN, CEFOTETAN DISODIUM
 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
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE, LIDOCAINE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 ZINACEF, CEFUROXIME SODIUM

TELI

* TELIX PHARMACEUTICALS US INC
 ILLUCCIX, GALLIUM GA-68 GOZETOTIDE

TENSHI

* TENSHI KAIZEN PVT LTD
 LORATADINE, LORATADINE (OTC)

TERSERA

* TERSERA THERAPEUTICS LLC
 ERGOMAR, ERGOTAMINE TARTRATE
 PRIALT, ZICONOTIDE ACETATE
 VARUBI, ROLAPITANT HYDROCHLORIDE
 XERMELO, TELOTRISTAT ETIPRATE
 ZOLADEX, GOSERELIN ACETATE

TETRAPHASE PHARMS

* TETRAPHASE PHARMACEUTICALS INC
 XERAVA, ERAVACYCLINE DIHYDROCHLORIDE

TEVA

* TEVA NEUROSCIENCE INC
 AZILECT, RASAGILINE MESYLATE

* TEVA PHARMACEUTICALS USA INC
 ACYCLOVIR, ACYCLOVIR
 ADIPEX-P, PHENTERMINE HYDROCHLORIDE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN PEDIATRIC, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 ATENOLOL, ATENOLOL
 AZITHROMYCIN, AZITHROMYCIN
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CALCITRIOL, CALCITRIOL
 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)
 CILOSTAZOL, CILOSTAZOL
 CIMETIDINE, CIMETIDINE
 CLARITHROMYCIN, CLARITHROMYCIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA INC
 CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
 CLONAZEPAM, CLONAZEPAM
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
 DIFLUNISAL, DIFLUNISAL
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 EPITOL, CARBAMAZEPINE
 ESZOPICLONE, ESZOPICLONE
 ETODOLAC, ETODOLAC
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FINASTERIDE, FINASTERIDE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLURBIPROFEN, FLURBIPROFEN
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 GALZIN, ZINC ACETATE
 GLYBURIDE (MICRONIZED), GLYBURIDE
 GLYBURIDE, GLYBURIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 KETOCONAZOLE, KETOCONAZOLE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LOVASTATIN, LOVASTATIN
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 MUPIROCIN, MUPIROCIN
 NAPROXEN, NAPROXEN
 NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 NEOMYCIN SULFATE, NEOMYCIN SULFATE
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 OFLOXACIN, OFLOXACIN
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXAPROZIN, OXAPROZIN
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PENICILLIN-VK, PENICILLIN V POTASSIUM
 PIROXICAM, PIROXICAM
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PRELONE, PREDNISOLONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SUCRALFATE, SUCRALFATE
 TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA BRANDED PHARMACEUTICAL PRODUCTS R AND D INC

AUSTEDO, DEUTETRABENAZINE
 CONDYLOX, PODOFILOX
 EMLA, LIDOCAINE
 LOESTRIN 21 1.5/30, ETHINYL ESTRADIOL
 LOESTRIN 21 1/20, ETHINYL ESTRADIOL
 LOESTRIN FE 1.5/30, ETHINYL ESTRADIOL
 LOESTRIN FE 1/20, ETHINYL ESTRADIOL
 LOSEASONIQUE, ETHINYL ESTRADIOL
 MICROZIDE, HYDROCHLOROTHIAZIDE
 NOR-QD, NORETHINDRONE
 PROAIR DIGIHALER, ALBUTEROL SULFATE
 PROAIR HFA, ALBUTEROL SULFATE
 PROAIR RESPICLICK, ALBUTEROL SULFATE
 PROGLYCEM, DIAZOXIDE
 QNASL, BECLOMETHASONE DIPROPIONATE
 QUARTETTE, ETHINYL ESTRADIOL
 SEASONALE, ETHINYL ESTRADIOL
 SEASONIQUE, ETHINYL ESTRADIOL
 ZIAC, BISOPROLOL FUMARATE

TEVA PARENTERAL

* TEVA PARENTERAL MEDICINES INC

LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

TEVA PHARM

* TEVA PHARMACEUTICAL INDUSTRIES LTD

AIRDUO DIGIHALER, FLUTICASONE PROPIONATE
 AIRDUO RESPICLICK, FLUTICASONE PROPIONATE
 ARMONAIR DIGIHALER, FLUTICASONE PROPIONATE

TEVA PHARMS

* TEVA PHARMACEUTICALS DEVELOPMENT INC

CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 PYRIMETHAMINE, PYRIMETHAMINE

* TEVA PHARMACEUTICALS USA

AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ANASTROZOLE, ANASTROZOLE
 BUDESONIDE, BUDESONIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFDINIR, CEFDINIR
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 FAMCICLOVIR, FAMCICLOVIR
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 IRBESARTAN, IRBESARTAN
 LANSOPRAZOLE, LANSOPRAZOLE
 LETROZOLE, LETROZOLE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LINEZOLID, LINEZOLID
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PACLITAXEL, PACLITAXEL
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUININE SULFATE, QUININE SULFATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TRANDOLAPRIL, TRANDOLAPRIL
 URSODIOL, URSODIOL
 VANDAZOLE, METRONIDAZOLE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE

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
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BEXAROTENE, BEXAROTENE
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BUDESONIDE, BUDESONIDE
 CLARAVIS, ISOTRETINOIN
 CLOZAPINE, CLOZAPINE
 COPAXONE, GLATIRAMER ACETATE
 DACARBAZINE, DACARBAZINE
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
 EPTIFIBATIDE, EPTIFIBATIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESTRADIOL, ESTRADIOL
 ETOPOSIDE, ETOPOSIDE
 FLUOROURACIL, FLUOROURACIL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 IBUPROFEN AND FAMOTIDINE, FAMOTIDINE
 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
 LOGILIA, ULIPRISTAL ACETATE
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OMEPRAZOLE, OMEPRAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PARICALCITOL, PARICALCITOL
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SUNITINIB MALATE, SUNITINIB MALATE
 TADALAFIL, TADALAFIL
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TOBRAMYCIN, TOBRAMYCIN
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 TREPROSTINIL, TREPROSTINIL
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VIGABATRIN, VIGABATRIN
 VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE
 VINOURELBINE TARTRATE, VINOURELBINE TARTRATE
 ZANOSAR, STREPTOZOCIN

* TEVA PHARMACEUTICALS USA INC
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ALYQ, TADALAFIL
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 CAPECITABINE, CAPECITABINE
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 DAPTOMYCIN, DAPTOMYCIN
 DARUNAVIR, DARUNAVIR
 DEFERASIROX, DEFERASIROX
 EFINACONAZOLE, EFINACONAZOLE
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EPINEPHRINE (AUTOINJECTOR), EPINEPHRINE
 ESTRADIOL, ESTRADIOL
 EVEROLIMUS, EVEROLIMUS
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FLUTICASONE PROPIONATE AND SALMETEROL XINAFOATE, FLUTICASONE PROPIONATE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 IVERMECTIN, IVERMECTIN
 LIOETHYRONINE SODIUM, LIOETHYRONINE SODIUM
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 MESALAMINE, MESALAMINE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METRONIDAZOLE, METRONIDAZOLE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 ONSURA, ETHINYL ESTRADIOL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 TOBRAMYCIN, TOBRAMYCIN
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

TEVA PHARMS USA INC

* TEVA PHARMACEUTICALS USA INC
 CHLORZOXAZONE, CHLORZOXAZONE
 ELIGLUSTAT TARTRATE, ELIGLUSTAT TARTRATE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 ETHINYL ESTRADIOL; ETNOGESTREL, ETHINYL ESTRADIOL
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 FULVESTRANT, FULVESTRANT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA INC
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 MIFEPRISTONE, MIFEPRISTONE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 ROMIDEPSIN, ROMIDEPSIN
 TIOPRONIN, TIOPRONIN

TG THERAPS

* TG THERAPEUTICS INC
 UKONIQ, UMBRALISIB TOSYLATE

THE FEINSTEIN INST

* THE FEINSTEIN INSTITUTE FOR MEDICAL RESEARCH
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

THEPHARMANETWORK LLC

* THEPHARMANETWORK LLC
 BENZONATATE, BENZONATATE
 ISONIAZID, ISONIAZID
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 NIMODIPINE, NIMODIPINE
 THERMAZENE, SILVER SULFADIAZINE

THERAPEUTICSMD INC

* THERAPEUTICSMD INC
 ANNOVERA, ETHINYL ESTRADIOL
 BIJUVA, ESTRADIOL
 IMVEXXY, ESTRADIOL

TIANJIN TIANYAO

* TIANJIN TIANYAO PHARMACEUTICALS CO LTD
 CELECOXIB, CELECOXIB
 METHYLPREDNISOLONE, METHYLPREDNISOLONE

TIME-CAP LABS INC

* TIME-CAP LABORATORIES INC
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE

TOLMAR

* TOLMAR INC
 AZELAIC ACID, AZELAIC ACID
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE, CALCIPOTRIENE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 KETOCONAZOLE, KETOCONAZOLE
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 METRONIDAZOLE, METRONIDAZOLE
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE

* TOLMAR INTERNATIONAL LTD
 FENSOLVI KIT, LEUPROLIDE ACETATE

TOLMAR THERAP

* TOLMAR THERAPEUTICS INC
 ELIGARD KIT, LEUPROLIDE ACETATE

TOPROL

* TOPROL ACQUISITION LLC
 TOPROL-XL, METOPROLOL SUCCINATE

TORPHARM

* TORPHARM INC
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE

TORRENT

* TORRENT PHARMA INC
 APREPITANT, APREPITANT
 ERYTHROMYCIN, ERYTHROMYCIN
 LACTULOSE, LACTULOSE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 SIROLIMUS, SIROLIMUS

* TORRENT PHARMACEUTICALS LTD
 ACYCLOVIR, ACYCLOVIR
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TORRENT PHARMACEUTICALS LTD
 ARIPIPRAZOLE, ARIPIPRAZOLE
 CELECOXIB, CELECOXIB
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GATIFLOXACIN, GATIFLOXACIN
 ITRACONAZOLE, ITRACONAZOLE
 LAMOTRIGINE, LAMOTRIGINE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 ROFLUMILAST, ROFLUMILAST
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TADALAFIL, TADALAFIL
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

TORRENT PHARMS

* TORRENT PHARMACEUTICALS LTD
 CARBAMAZEPINE, CARBAMAZEPINE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

TORRENT PHARMS LTD

* TORRENT PHARMACEUTICALS LTD
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FELODIPINE, FELODIPINE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE

TRAVERE

* TRAVERE THERAPEUTICS INC
 CHOLBAM, CHOLIC ACID

TREVENA

* TREVENA INC
 OLINVYK, OLICERIDINE

TRIPOINT

* TRIPOINT THERAPEUTICS
 ELEPSIA XR, LEVETIRACETAM

TRIS PHARMA INC

* TRIS PHARMA INC
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TRIS PHARMA INC
 DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
 DROXIDOPA, DROXIDOPA
 DYANAVEL XR 10, AMPHETAMINE
 DYANAVEL XR 15, AMPHETAMINE
 DYANAVEL XR 20, AMPHETAMINE
 DYANAVEL XR 5, AMPHETAMINE
 DYANAVEL XR, AMPHETAMINE
 GABAPENTIN, GABAPENTIN
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX, CHLORPHENIRAMINE POLISTIREX
 IBUPROFEN, IBUPROFEN (OTC)
 LEVETIRACETAM, LEVETIRACETAM
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 THEOPHYLLINE, THEOPHYLLINE

TRUSTEES UNIV PA

* TRUSTEES OF THE UNIV OF PENNSYLVANIA
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

TULEX PHARMS INC

* TULEX PHARMACEUTICALS INC
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

TWI PHARMS

* TWI PHARMACEUTICALS INC
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 NIFEDIPINE, NIFEDIPINE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 TESTOSTERONE, TESTOSTERONE

TWI PHARMS INC

* TWI PHARMACEUTICALS INC
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE

**** U ******UBI**

* UBI PHARMA INC
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE

UCB INC

* UCB INC
 BRIVIACT, BRIVARACETAM
 KEPPRA XR, LEVETIRACETAM
 KEPPRA, LEVETIRACETAM
 NAYZILAM, MIDAZOLAM
 NEUPRO, ROTIGOTINE
 VIMPAT, LACOSAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ******UCLA BIOMEDICAL**

* UCLA BIOMEDICAL CYCLOTRON
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UCSF RODIOPHARM

* UCSF RODIOPHARMACEUTICAL FACILITY
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UIHC PET IMAGING

* UNIV IOWA HOSPS AND CLINICS PET IMAGING CENTER
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 GALLIUM DOTATOC GA 68, GALLIUM DOTATOC GA-68

ULTRAGENYX PHARM INC

* ULTRAGENYX PHARMACEUTICAL INC
 DOJOLVI, TRIHEPTANOIN

UMEDICA LABS PVT LTD

* UMEDICA LABORATORIES PRIVATE LTD
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 CARBAMAZEPINE, CARBAMAZEPINE
 CELECOXIB, CELECOXIB
 CHLORTHALIDONE, CHLORTHALIDONE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TADALAFIL, TADALAFIL

UNICHEM

* UNICHEM LABORATORIES LTD
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALLOPURINOL, ALLOPURINOL
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 ATENOLOL, ATENOLOL
 BACLOFEN, BACLOFEN
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CELECOXIB, CELECOXIB
 CHLORTHALIDONE, CHLORTHALIDONE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MELOXICAM, MELOXICAM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 PIROXICAM, PIROXICAM
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 ZALEPLON, ZALEPLON
 ZONISAMIDE, ZONISAMIDE

UNICHEM LABS LTD

* UNICHEM LABORATORIES LIMITED

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

* UNICHEM LABORATORIES LIMITED
DIVALPROEX SODIUM, DIVALPROEX SODIUM

* UNICHEM LABORATORIES LTD
LAMOTRIGINE, LAMOTRIGINE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TOPIRAMATE, TOPIRAMATE

UNIMARK REMEDIES LTD

* UNIMARK REMEDIES LTD
MONTELUKAST SODIUM, MONTELUKAST SODIUM

UNIQUE PHARM

* UNIQUE PHARMACEUTICAL LABORATORIES
CARBAMAZEPINE, CARBAMAZEPINE
LORATADINE, LORATADINE (OTC)

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
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE 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
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
TINIDAZOLE, TINIDAZOLE
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE

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 CA LOS ANGELES

* UNIV CALIFORNIA LOS ANGELES
GALLIUM GA 68 PSMA-11, GALLIUM GA-68 PSMA-11

UNIV MICHIGAN

* UNIV MICHIGAN PET RADIOPHARMACEUTICAL PRODUCTION PROGRAM
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UNIV OF CA SAN FRAN

* UNIV OF CALIFORNIA SAN FRANCISCO
GALLIUM GA 68 PSMA-11, GALLIUM GA-68 PSMA-11

UNIV SOUTHERN CA

* UNIV SOUTHERN CALIFORNIA DBA USC MOLECULAR IMAGING CENTER
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 TX SW MEDCTR

* UNIV TEXAS SOUTHWESTERN MEDCTR
AMMONIA N 13, AMMONIA N-13

UNIV UTAH CYCLOTRON

* UNIV UTAH CYCLOTRON RADIOCHEMISTRY LAB
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ******UNIV WISCONSIN**

- * UNIV WISCONSIN SYSTEM
AMMONIA N 13, AMMONIA N-13

UPJOHN

- * UPJOHN MANUFACTURING IRELAND UNLTD
LIPITOR, ATORVASTATIN CALCIUM
RELPAX, ELETRIPTAN HYDROBROMIDE
- * UPJOHN US 1 LLC
CARDURA XL, DOXAZOSIN MESYLATE
CARDURA, DOXAZOSIN MESYLATE
DILANTIN, PHENYTOIN SODIUM
DILANTIN-125, PHENYTOIN
GEODON, ZIPRASIDONE HYDROCHLORIDE
GEODON, ZIPRASIDONE MESYLATE
NEURONTIN, GABAPENTIN
NITROSTAT, NITROGLYCERIN
NORVASC, AMLODIPINE BESYLATE
REVATIO, SILDENAFIL CITRATE
VIAGRA, SILDENAFIL CITRATE
ZOLOFT, SERTRALINE HYDROCHLORIDE
- * UPJOHN US 2 LLC
CELEBREX, CELECOXIB
DETROL LA, TOLTERODINE TARTRATE
DETROL, TOLTERODINE TARTRATE
EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE
INSPRA, EPLERENONE
LYRICA CR, PREGABALIN
LYRICA, PREGABALIN
XALATAN, LATANOPROST
XANAX XR, ALPRAZOLAM
XANAX, ALPRAZOLAM

UPSHER SMITH LABS

- * UPSHER SMITH LABORATORIES LLC
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
BACLOFEN, BACLOFEN
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
BEXAROTENE, BEXAROTENE
BUMETANIDE, BUMETANIDE
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CLOBAZAM, CLOBAZAM
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
ETHACRYNIC ACID, ETHACRYNIC ACID
EXEMESTANE, EXEMESTANE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
HALOPERIDOL, HALOPERIDOL
ISOTRETINOIN, ISOTRETINOIN
JANTOVEN, WARFARIN SODIUM
KLOF-CON M10, POTASSIUM CHLORIDE
KLOF-CON M15, POTASSIUM CHLORIDE
KLOF-CON M20, POTASSIUM CHLORIDE
KLOF-CON, POTASSIUM CHLORIDE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
MORPHINE SULFATE, MORPHINE SULFATE
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
NYSTATIN, NYSTATIN
ORVATEN, MIDODRINE HYDROCHLORIDE
OXANDROLONE, OXANDROLONE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

* UPSHER SMITH LABORATORIES LLC
 PACERONE, AMIODARONE HYDROCHLORIDE
 PREVALITE, CHOLESTYRAMINE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 QUDEXY XR, TOPIRAMATE
 SILODOSIN, SILODOSIN
 SORINE, SOTALOL HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TOSYMRA, SUMATRIPTAN
 VOGELXO, TESTOSTERONE
 ZEMBRACE SYMTOUCH, SUMATRIPTAN SUCCINATE

UROGEN PHARMA

* UROGEN PHARMA LTD
 JELMYTO, MITOMYCIN

UROVANT

* UROVANT SCIENCES GMBH
 GEMTESA, VIBEGRON

US ANTIBIOTICS

* US ANTIBIOTICS LLC
 AMOXIL, AMOXICILLIN
 AUGMENTIN '125', AMOXICILLIN
 AUGMENTIN '250', AMOXICILLIN
 AUGMENTIN '875', AMOXICILLIN
 AUGMENTIN ES-600, AMOXICILLIN
 LAROTID, AMOXICILLIN

USPHARMA

* USPHARMA LTD
 NITRO-DUR, NITROGLYCERIN

USPHARMA WINDLAS

* USPHARMA WINDLAS LLC
 AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 PRASUGREL, PRASUGREL HYDROCHLORIDE

USV

* USV PRIVATE LTD
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

USWM

* USWM LLC
 CORGARD, NADOLOL
 LUCEMYRA, LOFEXIDINE HYDROCHLORIDE
 REVONTO, DANTROLENE SODIUM

**** V ******VALEANT**

* VALEANT PHARMACEUTICALS INTERNATIONAL
 BONTRIL PDM, PHENDIMETRAZINE TARTRATE
 MYSOLINE, PRIMIDONE

VALEANT BERMUDA

* VALEANT INTERNATIONAL BERMUDA
 PENLAC, CICLOPIROX
 RETIN-A, TRETINOIN

VALEANT INTL

* VALEANT INTERNATIONAL BARBADOS SRL
 RETIN-A MICRO, TRETINOIN
 RETIN-A, TRETINOIN
 RETIN-A-MICRO, TRETINOIN
 * VALEANT INTERNATIONAL SRL
 BENZAMYCIN, BENZOYL PEROXIDE

VALEANT LUXEMBOURG

* VALEANT PHARMACEUTICALS LUXEMBOURG SARL
 TARGRETIN, BEXAROTENE
 VISUDYNE, VERTEPORFIN

VALEANT PHARM INTL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VALEANT PHARMACEUTICALS INTERNATIONAL
 ANDROID 25, METHYLTESTOSTERONE
 LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE

VALEANT PHARMS

* VALEANT PHARMACEUTICALS NORTH AMERICA LLC
 PENTOXIFYLLINE, PENTOXIFYLLINE

VALEANT PHARMS INTL

* VALEANT PHARMACEUTICALS INTERNATIONAL
 APRISO, MESALAMINE
 COLAZAL, BALSALAZIDE DISODIUM
 CUPRIMINE, PENICILLAMINE
 LACRISERT, HYDROXYPROPYL CELLULOSE
 TIMOPTIC, TIMOLOL MALEATE

VALEANT PHARMS LLC

* VALEANT PHARMACEUTICALS NORTH AMERICA LLC
 TIMOPTIC-XE, TIMOLOL MALEATE

VALEANT PHARMS NORTH

* VALEANT PHARMACEUTICALS NORTH AMERICA LLC
 CARAC, FLUOROURACIL
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 FENOFIBRATE, FENOFIBRATE
 NIFEDIPINE, NIFEDIPINE
 NORITATE, METRONIDAZOLE
 RENOVA, TRETINOIN
 RETIN-A, TRETINOIN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

VALIDUS PHARMS

* VALIDUS PHARMACEUTICALS LLC
 ANZEMET, DOLASETRON MESYLATE
 BUMEX, BUMETANIDE
 DRISDOL, ERGOCALCIFEROL
 EQUETRO, CARBAMAZEPINE
 LASIX, FUROSEMIDE
 LOPRESSOR HCT, HYDROCHLOROTHIAZIDE
 LOPRESSOR, METOPROLOL TARTRATE
 LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
 LOTENSIN, BENAZEPRIL HYDROCHLORIDE
 NORPRAMIN, DESIPRAMINE HYDROCHLORIDE
 PARLODEL, BROMOCRIPTINE MESYLATE
 ROCALTROL, CALCITRIOL

VALIDUS PHARMS INC

* VALIDUS PHARMACEUTICALS INC
 MARPLAN, ISOCARBOXAZID

VANDA PHARMS INC

* VANDA PHARMACEUTICALS INC
 FANAPT, ILOPERIDONE
 HETLIOZ LQ, TASIMELTEON
 HETLIOZ, TASIMELTEON

VELOXIS PHARMS INC

* VELOXIS PHARMACEUTICALS INC
 ENVARBUS XR, TACROLIMUS

VERITY

* VERITY PHARMACEUTICALS INC
 TRELSTAR, TRIPTORELIN PAMOATE

VERO BIOTECH

* VERO BIOTECH
 GENOSYL, NITRIC OXIDE

VEROSCIENCE

* VEROSCIENCE LLC
 CYCLOSET, BROMOCRIPTINE MESYLATE

VERTEX PHARMS

* VERTEX PHARMACEUTICALS INC
 KALYDECO, IVACAFTOR

VERTEX PHARMS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VERTEX PHARMACEUTICALS INC
 KALYDECO, IVACAFTOR
 ORKAMBI, IVACAFTOR
 SYMDEKO (COPACKAGED), IVACAFTOR
 TRIKAFTA (COPACKAGED), ELEXACAFTOR, IVACAFTOR, TEZACAFTOR

VERTICAL PHARMS LLC

* VERTICAL PHARMACEUTICALS LLC
 DIVIGEL, ESTRADIOL

VERU

* VERU INC
 ENTADFI, FINASTERIDE

VGYAAN

* VGYAAN PHARMACEUTICALS LLC
 CARBAMAZEPINE, CARBAMAZEPINE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 NADOLOL, NADOLOL
 URSODIOL, URSODIOL

VICURON HOLDINGS

* VICURON HOLDINGS LLC
 ERAXIS, ANIDULAFUNGIN

VIFOR FRESENIUS

* VIFOR FRESENIUS MEDICAL CARE RENAL PHARMA FRANCE
 VELPHORO, FERRIC OXYHYDROXIDE

VIFOR PHARMA

* VIFOR PHARMA INC
 VELTASSA, PATIROMER SORBITEX CALCIUM

VIIV HLTHCARE

* VIIV HEALTHCARE CO
 APRETUDE, CABOTEGRAVIR
 CABENUVA KIT, CABOTEGRAVIR
 COMBIVIR, LAMIVUDINE
 DOVATO, DOLUTEGRAVIR SODIUM
 EPIVIR, LAMIVUDINE
 EPZICOM, ABACAVIR SULFATE
 JULUCA, DOLUTEGRAVIR SODIUM
 LEXIVA, FOSAMPRENAVIR CALCIUM
 RETROVIR, ZIDOVUDINE
 RUKOBIA, FOSTEMSAVIR TROMETHAMINE
 SELZENTRY, MARAVIROC
 TIVICAY PD, DOLUTEGRAVIR SODIUM
 TIVICAY, DOLUTEGRAVIR SODIUM
 TRIUMEQ, ABACAVIR SULFATE
 TRIZIVIR, ABACAVIR SULFATE
 VOCABRIA, CABOTEGRAVIR SODIUM
 ZIAGEN, ABACAVIR SULFATE

VINTAGE

* VINTAGE PHARMACEUTICALS LLC
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 DUTASTERIDE, DUTASTERIDE

VINTAGE PHARMS

* VINTAGE PHARMACEUTICALS
 GILDAGIA, ETHINYL ESTRADIOL
 GILDESS 24 FE, ETHINYL ESTRADIOL
 KIMIDESS, DESOGESTREL
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

* VINTAGE PHARMACEUTICALS INC
 ALLOPURINOL, ALLOPURINOL
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 BACLOFEN, BACLOFEN
 IBUPROFEN, IBUPROFEN (OTC)
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 PERPHENAZINE, PERPHENAZINE
 PREDNISON, PREDNISON

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ******VINTAGE PHARMS LLC**

* VINTAGE PHARMACEUTICALS LLC
 FELODIPINE, FELODIPINE
 MYZILRA, ETHINYL ESTRADIOL
 PERCOCET, ACETAMINOPHEN
 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

VISTA PHARMS

* VISTA PHARMACEUTICALS INC
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

VISTAPHARM

* VISTAPHARM INC
 ACYCLOVIR, ACYCLOVIR
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 ARIPIPRAZOLE, ARIPIPRAZOLE
 CHLOROTHALIDONE, CHLOROTHALIDONE
 CLOBAZAM, CLOBAZAM
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 LACTULOSE, LACTULOSE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NYSTATIN, NYSTATIN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PHENYTOIN, PHENYTOIN
 THYQUIDITY, LEVOTHYROXINE SODIUM

VISUM PHARM

* VISUM PHARMACEUTICAL CO LTD
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE

VITRUVIAS THERAP

* VITRUVIAS THERAPEUTICS
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 * VITRUVIAS THERAPEUTICS LLC
 CYANOCOBALAMIN, CYANOCOBALAMIN
 LIDOCAINE, LIDOCAINE

VIVUS

* VIVUS INC
 QSYMIA, PHENTERMINE HYDROCHLORIDE

VIWIT PHARM

* VIWIT PHARMACEUTICAL CO LTD
 LEVETIRACETAM, LEVETIRACETAM
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE

VKT PHARMA

* VKT PHARMA PRIVATE LTD
 FAMOTIDINE, FAMOTIDINE
 LEVETIRACETAM, LEVETIRACETAM
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 TADALAFIL, TADALAFIL

VPI PHARMS INC

* VPI PHARMACEUTICALS INC
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM

VPNA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VALEANT PHARMACEUTICALS NORTH AMERICA
DICLOFENAC SODIUM, DICLOFENAC SODIUM

VYERA PHARMS LLC

* VYERA PHARMACEUTICALS LLC
DARAPRIM, PYRIMETHAMINE

VYNE

* VYNE PHARMACEUTICALS INC
AMZEEQ, MINOCYCLINE HYDROCHLORIDE
ZILXI, MINOCYCLINE HYDROCHLORIDE

**** 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
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE

* WATSON LABORATORIES INC
ACARBOSE, ACARBOSE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALLOPURINOL, ALLOPURINOL
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
AMOXAPINE, AMOXAPINE
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
CARISOPRODOL, CARISOPRODOL
CHLORZOXAZONE, CHLORZOXAZONE
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
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
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LAMOTRIGINE, LAMOTRIGINE
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LISINAPRIL, LISINAPRIL
LORAZEPAM, LORAZEPAM
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METRONIDAZOLE, METRONIDAZOLE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MINOXIDIL, MINOXIDIL
NABUMETONE, NABUMETONE
NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
NATEGLINIDE, NATEGLINIDE
NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
NIZATIDINE, NIZATIDINE
NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
PREDNISOLONE, PREDNISOLONE
PREDNISON, PREDNISON
PRIMIDONE, PRIMIDONE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
QUASENSE, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WATSON LABORATORIES INC
 RAMIPRIL, RAMIPRIL
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 SULFASALAZINE, SULFASALAZINE
 SULINDAC, SULINDAC
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 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
 AMBRISENTAN, AMBRISENTAN
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 BOSENTAN, BOSENTAN
 BRINZOLAMIDE, BRINZOLAMIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CELECOXIB, CELECOXIB
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 COLCHICINE, COLCHICINE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 METRONIDAZOLE, METRONIDAZOLE
 MODAFINIL, MODAFINIL
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 PENICILLAMINE, PENICILLAMINE
 PERPHENAZINE, PERPHENAZINE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PROPOFOL, PROPOFOL
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE

WATSON LABS TEVA

* WATSON LABORATORIES INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 ALVIMOPAN, ALVIMOPAN
 BICALUTAMIDE, BICALUTAMIDE
 BUPRENORPHINE, BUPRENORPHINE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 GLIPIZIDE, GLIPIZIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 ISRADIPINE, ISRADIPINE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 PROBENECID, PROBENECID
 SIMVASTATIN, SIMVASTATIN
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

WATSON PHARMS INC

* WATSON PHARMACEUTICALS INC
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE

WAYLIS THERAP

* WAYLIS THERAPEUTICS LLC
 FLUTAMIDE, FLUTAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ******WELLSTAT THERAP**

* WELLSTAT THERAPEUTICS CORP
 VISTOGARD, URIDINE TRIACETATE
 XURIDEN, URIDINE TRIACETATE

WES PHARMA INC

* WES PHARMA INC
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN

WEST WARD

* WEST WARD PHARMACEUTICAL CORP
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

WEST WARD PHARM CORP

* WEST WARD PHARMACEUTICAL CORP
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

WEST-WARD PHARMS INT

* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
 BUMETANIDE, BUMETANIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYTARABINE, CYTARABINE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DIGOXIN, DIGOXIN
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 PENTOSTATIN, PENTOSTATIN
 PHENYTOIN SODIUM, PHENYTOIN SODIUM
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, FERRIC OXYHYDROXIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 THIOTEPA, THIOTEPA

WESTMINSTER PHARMS

* WESTMINSTER PHARMACEUTICALS LLC
 EPLERENONE, EPLERENONE

WILSHIRE PHARMS INC

* WILSHIRE PHARMACEUTICALS INC
 CARISOPRODOL, CARISOPRODOL
 PERPHENAZINE, PERPHENAZINE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE

WINDER LABS LLC

* WINDER LABORATOIRES LLC
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE

WISCONSIN

* WISCONSIN MEDICAL RADIOPHARMACY LLC
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

WOCKHARDT

* WOCKHARDT LTD
 AZITHROMYCIN, AZITHROMYCIN
 CEFTRIAZONE, CEFTRIAZONE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ******* WOCKHARDT LTD**

CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 FAMOTIDINE, FAMOTIDINE (OTC)
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FUROSEMIDE, FUROSEMIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEVETIRACETAM, LEVETIRACETAM
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE

WOCKHARDT BIO AG*** WOCKHARDT BIO AG**

ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACETIC ACID, ACETIC ACID, GLACIAL
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 BROMFED-DM, BROMPHENIRAMINE MALEATE
 CARBAMAZEPINE, CARBAMAZEPINE
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 CYCLOSPORINE, CYCLOSPORINE
 DECITABINE, DECITABINE
 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
 IMATINIB MESYLATE, IMATINIB MESYLATE
 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**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WOCKHARDT LTD
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CAPTOPRIL, CAPTOPRIL
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)

WOCKHARDT USA

* WOCKHARDT USA INC
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 * WOCKHARDT USA LLC
 LANSOPRAZOLE, LANSOPRAZOLE

WOODWARD

* WOODWARD PHARMA SERVICES LLC
 ACIPHEX, RABEPRAZOLE SODIUM
 AVANDIA, ROSIGLITAZONE MALEATE
 AVODART, DUTASTERIDE
 COREG CR, CARVEDILOL PHOSPHATE
 COREG, CARVEDILOL
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 JALYN, DUTASTERIDE
 LOVAZA, OMEGA-3-ACID ETHYL ESTERS
 MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE
 MIVACURIUM CHLORIDE, MIVACURIUM CHLORIDE

WRASER PHARMS

* WRASER PHARMACEUTICALS LLC
 CETRAXAL, CIPROFLOXACIN HYDROCHLORIDE

WUSM CYCLOTRON

* WASHINGTON UNIV SCH MEDICINE CYCLOTRON FACILITY
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

WYETH PHARMS

* WYETH PHARMACEUTICALS LLC
 DUAVEE, BAZEDOXIFENE ACETATE
 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

**** X ******XELLIA PHARMS APS**

* XELLIA PHARMACEUTICALS APS
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 DAPTOMYCIN, DAPTOMYCIN
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TIGECYCLINE, TIGECYCLINE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VANCOMYCIN, VANCOMYCIN
 VORICONAZOLE, VORICONAZOLE

XERIS

* XERIS PHARMACEUTICALS INC
 GVOKE HYPOPEN, GLUCAGON
 GVOKE PFS, GLUCAGON
 KEVEYIS, DICHLORPHENAMIDE

XGEN PHARMS

* XGEN PHARMACEUTICALS DJB INC
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 AMPHOTERICIN B, AMPHOTERICIN B

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** X ****

* XGEN PHARMACEUTICALS DJB INC
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 DACTINOMYCIN, DACTINOMYCIN
 FOLIC ACID, FOLIC ACID
 IBUPROFEN LYSINE, IBUPROFEN LYSINE
 LEVETIRACETAM, LEVETIRACETAM
 LINCOMYCIN, LINCOMYCIN HYDROCHLORIDE
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
 NEOMYCIN SULFATE, NEOMYCIN SULFATE
 NYSTATIN, NYSTATIN
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TRANEXAMIC ACID, TRANEXAMIC ACID

XIAMEN LP PHARM CO

* XIAMEN LP PHARMACEUTICAL CO LTD
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

XIROMED

* XIROMED PHARMA ESPANA SL
 ACYCLOVIR, ACYCLOVIR
 ALTAVERA, ETHINYL ESTRADIOL
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 ESTARYLLA, ETHINYL ESTRADIOL
 FLUOCINONIDE, FLUOCINONIDE
 FOSFOMYCIN TROMETHAMINE, FOSFOMYCIN TROMETHAMINE
 FULVESTRANT, FULVESTRANT
 GEMMILY, ETHINYL ESTRADIOL
 GLYCOPYRROLATE, GLYCOPYRROLATE
 INTROVALE, ETHINYL ESTRADIOL
 ISIBLOOM, DESOGESTREL
 JAIMIESS, ETHINYL ESTRADIOL
 KETOCONAZOLE, KETOCONAZOLE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LORYNA, DROSPIRENONE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 OMEPRAZOLE, OMEPRAZOLE
 PROGESTERONE, PROGESTERONE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SYEDA, DROSPIRENONE
 TESTOSTERONE, TESTOSTERONE
 TRI-ESTARYLLA, ETHINYL ESTRADIOL
 TRI-LO-ESTARYLLA, ETHINYL ESTRADIOL
 VIENVA, ETHINYL ESTRADIOL
 VOLNEA, DESOGESTREL

XSPIRE PHARMA

* XSPIRE PHARMA
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 NALFON, FENOPROFEN CALCIUM
 TREZIX, ACETAMINOPHEN
 ZONTIVITY, VORAPAXAR SULFATE

* XSPIRE PHARMA LLC
 DEXAMETHASONE, DEXAMETHASONE
 FENOPROFEN CALCIUM, FENOPROFEN CALCIUM

XTTRIUM

* XTTRIUM LABORATORIES INC
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** X ****

* XTTRIUM LABORATORIES INC
EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)

XTTRIUM LABS INC

* XTTRIUM LABORATORIES INC
LACTULOSE, LACTULOSE

**** Y ******YABAO PHARM**

* YABAO PHARMACEUTICAL CO LTD BEIJING
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE

YANGLING BUCHANG

* YANGLING BUCHANG PHARMACEUTICAL CO LTD
TADALAFIL, TADALAFIL

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
FENOFIBRIC ACID, CHOLINE FENOFIBRATE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

YILING

* YILING PHARMACEUTICAL LTD
ACYCLOVIR, ACYCLOVIR
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CELECOXIB, CELECOXIB
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
FELODIPINE, FELODIPINE
LAMOTRIGINE, LAMOTRIGINE
LISINAPRIL, LISINAPRIL
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE

YOUNGTECH PHARMS INC

* YOUNGTECH PHARMACEUTICALS INC
METOPROLOL TARTRATE, METOPROLOL TARTRATE

YUNG SHIN PHARM

* YUNG SHIN PHARMACEUTICAL INDUSTRIAL CO LTD
AZITHROMYCIN, AZITHROMYCIN
CEFACLOR, CEFACLOL
CEPHALEXIN, CEPHALEXIN
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
ENTECAVIR, ENTECAVIR
FELODIPINE, FELODIPINE
MELOXICAM, MELOXICAM

**** Z ******ZAMBON SPA**

* ZAMBON SPA ITALY
MONUROL, FOSFOMYCIN TROMETHAMINE

ZEALAND PHARMA

* ZEALAND PHARMA US INC
ZEGALOGUE (AUTOINJECTOR), DASIGLUCAGON HYDROCHLORIDE
ZEGALOGUE, DASIGLUCAGON HYDROCHLORIDE

ZENNOVA

* ZENNOVA PHARMACEUTICALS CHENGDU CO LTD
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ******ZHEJIANG HISUN PHARM**

* ZHEJIANG HISUN PHARMACEUTICAL CO LTD
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL

ZHEJIANG JINGXIN

* ZHEJIANG JINGXIN PHARMACEUTICAL CO LTD
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM

ZHEJIANG JUTAI PHARM

* ZHEJIANG JUTAI PHARMACEUTICAL CO LTD
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE

ZHEJIANG YONGTAI

* ZHEJIANG YONGTAI PHARMACEUTICAL CO LTD
GABAPENTIN, GABAPENTIN
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

ZIONEXA

* ZIONEXA US CORP
CERIANNA, FLUOROESTRADIOL F-18

ZOGENIX INC

* ZOGENIX INC
FINTEPLA, FENFLURAMINE HYDROCHLORIDE

ZYDUS

* ZYDUS WORLDWIDE DMCC
AZITHROMYCIN, AZITHROMYCIN
BACLOFEN, BACLOFEN
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
LEFLUNOMIDE, LEFLUNOMIDE
LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
PHYTONADIONE, PHYTONADIONE
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
URSODIOL, URSODIOL

ZYDUS HLTHCARE

* ZYDUS HEALTHCARE USA LLC
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
LANSOPRAZOLE, LANSOPRAZOLE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

ZYDUS NOVELTECH INC

* ZYDUS NOVELTECH INC
RIVASTIGMINE, RIVASTIGMINE

ZYDUS PHARMS

* ZYDUS PHARMACEUTICALS USA INC
ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
ACETYLCYSTEINE, ACETYLCYSTEINE
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
ACYCLOVIR, ACYCLOVIR
ALBENDAZOLE, ALBENDAZOLE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALLOPURINOL, ALLOPURINOL
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMBRISENTAN, AMBRISENTAN
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
APREMILAST, APREMILAST
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ****

* ZYDUS PHARMACEUTICALS USA INC
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BOSENTAN, BOSENTAN
 BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CARBAMAZEPINE, CARBAMAZEPINE
 CARBIDOPA, CARBIDOPA
 CHLORTHALIDONE, CHLORTHALIDONE
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 COLCHICINE, COLCHICINE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DECITABINE, DECITABINE
 DEFERASIROX, DEFERASIROX
 DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIFLUNISAL, DIFLUNISAL
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DOXYCYCLINE, DOXYCYCLINE HYCLATE
 DROXIDOPA, DROXIDOPA
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 ENTECAVIR, ENTECAVIR
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 ETODOLAC, ETODOLAC
 ETOMIDATE, ETOMIDATE
 EXEMESTANE, EXEMESTANE
 EZETIMIBE, EZETIMIBE
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOCINONIDE, FLUOCINONIDE
 GLIPIZIDE, GLIPIZIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYBURIDE, GLYBURIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 INDOMETHACIN, INDOMETHACIN
 ITRACONAZOLE, ITRACONAZOLE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LAMOTRIGINE, LAMOTRIGINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LINEZOLID, LINEZOLID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ****

* ZYDUS PHARMACEUTICALS USA INC
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NADOLOL, NADOLOL
 NATEGLINIDE, NATEGLINIDE
 NELARABINE, NELARABINE
 NIFEDIPINE, NIFEDIPINE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NYSTATIN, NYSTATIN
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PERPHENAZINE, PERPHENAZINE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RAMELTEON, RAMELTEON
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SIROLIMUS, SIROLIMUS
 SPIRONOLACTONE, SPIRONOLACTONE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TADALAFIL, TADALAFIL
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOFACITINIB, TOFACITINIB CITRATE
 TOPIRAMATE, TOPIRAMATE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 VORTIOXETINE HYDROBROMIDE, VORTIOXETINE HYDROBROMIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN

ZYDUS PHARMS USA

* ZYDUS PHARMACEUTICALS USA INC
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ATENOLOL, ATENOLOL
 AZATHIOPRINE, AZATHIOPRINE
 BENZONATATE, BENZONATATE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ****

* ZYDUS PHARMACEUTICALS USA INC
 RISPERIDONE, RISPERIDONE
 SIMVASTATIN, SIMVASTATIN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM

ZYDUS PHARMS USA INC

* ZYDUS PHARMACEUTICALS USA INC
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ANASTROZOLE, ANASTROZOLE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 CARVEDILOL, CARVEDILOL
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 FINASTERIDE, FINASTERIDE
 GABAPENTIN, GABAPENTIN
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 IRBESARTAN, IRBESARTAN
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 OMEPRAZOLE, OMEPRAZOLE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN

ZYLA

* ZYLA LIFE SCIENCES US INC
 OXAYDO, OXYCODONE HYDROCHLORIDE
 SPRIX, KETOROLAC TROMETHAMINE
 ZORVOLEX, DICLOFENAC

42ND EDITION 2022 - APPROVED DRUG PRODUCTS LIST

APPENDIX C**UNIFORM TERMS*****DOSAGE FORMS***

AEROSOL, FOAM	OIL
AEROSOL, METERED	OIL/DROPS
CAPSULE	OINTMENT
CAPSULE, DELAYED REL PELLETS	OINTMENT, AUGMENTED
CAPSULE, DELAYED RELEASE	PASTE
CAPSULE, EXTENDED RELEASE	PATCH
CAPSULE, PELLETS	PELLET
CLOTH	PELLETS
CONCENTRATE	POWDER
CREAM	POWDER, EXTENDED RELEASE
CREAM, AUGMENTED	POWDER, METERED
CREAM, INSERT	RING
ELIXIR	SHAMPOO
EMULSION	SOLUTION
ENEMA	SOLUTION FOR SLUSH
FILM	SOLUTION, EXTENDED RELEASE
FILM, EXTENDED RELEASE	SOLUTION, GEL FORMING/DROPS
FOAM	SOLUTION, METERED
FOR SOLUTION	SOLUTION/DROPS
FOR SUSPENSION	SPONGE
FOR SUSPENSION, DELAYED RELEASE	SPRAY
FOR SUSPENSION, EXTENDED RELEASE	SPRAY, METERED
GAS	SUPPOSITORY
GEL	SUSPENSION
GEL, AUGMENTED	SUSPENSION, EXTENDED RELEASE
GEL, METERED	SUSPENSION, LIPOSOMAL
GRANULE	SUSPENSION/DROPS
GRANULE, DELAYED RELEASE	SWAB
GRANULES	SYRUP
GUM, CHEWING	SYSTEM
IMPLANT	TABLET
INHALANT	TABLET, CHEWABLE
INJECTABLE	TABLET, DELAYED RELEASE
INJECTABLE, LIPID COMPLEX	TABLET, EFFERVESCENT
INJECTABLE, LIPOSOMAL	TABLET, EXTENDED RELEASE
INJECTION, EXTENDED RELEASE	TABLET, EXTENDED RELEASE, CHEWABLE
INSERT	TABLET, FOR SUSPENSION
INSERT, EXTENDED RELEASE	TABLET, ORALLY DISINTEGRATING
INTRAUTERINE DEVICE	TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE
JELLY	TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE
LIQUID	TAPE
LOTION	TROCHE/LOZENGE
LOTION, AUGMENTED	
LOTION/SHAMPOO	

Note: Terms comprise currently marketed products

APPENDIX C

UNIFORM TERMS

ROUTES OF ADMINISTRATION

BUCCAL	INTRAVESICULAR
DENTAL	INTRAVITREAL
ENDOCERVICAL	IRRIGATION
ENDOTRACHEAL	IV (INFUSION)
ENTERAL	N/A
IMPLANTATION	NASAL
INHALATION	OPHTHALMIC
INJECTION	ORAL
INTERSTITIAL	ORAL-21
INTRA-ANAL	ORAL-28
INTRA-ARTERIAL	OTIC
INTRA-ARTICULAR	PERFUSION, CARDIAC
INTRACAVITARY	PERIARTICULAR
INTRACRANIAL	PERIODONTAL
INTRADERMAL	PYELOCALYCEAL
INTRAMUSCULAR	RECTAL
INTRAOCULAR	SPINAL
INTRAOSSEOUS	SUBCUTANEOUS
INTRAPERITONEAL	SUBLINGUAL
INTRAPLEURAL	TOPICAL
INTRATHECAL	TRANSDERMAL
INTRATRACHEAL	TRANSMUCOSAL
INTRAUTERINE	URETHRAL
INTRAVENOUS	VAGINAL
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
SC	SUBCUTANEOUS
SQ CM	SQUARE CENTIMETER
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 (the 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

This *Addendum* identifies:

- Drugs approved under section 505(c) of the FD&C Act that have qualified 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) of the FD&C Act
- Drugs that have qualified for Orphan Drug Exclusivity pursuant to Section 527 of the FD&C Act
- Drugs that have qualified for Pediatric Exclusivity pursuant to Section 505A of the FD&C Act
- Drugs that have qualified for Generating Antibiotics Incentives Now (GAIN) exclusivity pursuant to Section 505E of the FD&C Act
- 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) of the FD&C Act
- Generic drugs approved under section 505(j) of the FD&C Act 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 established name of the active ingredient, followed by the proprietary name (brand name or trade name) of the drug product. Active ingredient headings for multiple active ingredient, fixed-combination drug products are arranged alphabetically.

Individual descriptions of the protected use have been added to each Orphan Drug Exclusivity entry listed in the Orange Book. Such descriptions of Orphan Drug Exclusivity were included beginning with the 38th edition of the Orange Book. The ODE* code means that the timing of approval of certain follow-on applications may be subject to delay due to ODE for another drug that has the same active moiety.

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.

Patent Information

The FD&C Act requires that patent information be filed with all newly submitted Section 505(b) drug applications. In addition, patent information must be filed on Form FDA 3542 within 30 days of the date of approval of a Section 505(b) drug application.¹ FDA publishes certain information from Form FDA 3542 in the Orange Book after approval of the new drug application (NDA) or supplement.

The Orange Book includes 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 abbreviated new drug application (ANDA).²

The patents that FDA regards as covered by the statutory provisions for submission of patent information for listing in the Orange Book are:

- Patents that claim the drug for which the applicant submitted the application and are drug substance (active ingredient) patents or drug product (formulation or composition) patents; or
- Patents that claim a method of using such drug for which approval has been granted in the application.

This information, as provided by the NDA holder on Form FDA 3542, will be published as described above. 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.

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 regularly, should be consulted for the most recent patent and exclusivity information.

¹ 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).

² 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).

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ABACAVIR SULFATE; DOLUTEGRAVIR SODIUM; LAMIVUDINE - TRIUMEO</u>						
N 205551	001	8129385	Oct 05, 2027	DS DP		
		9242986	Dec 08, 2029	DS DP		
<u>ABALOPARATIDE - TYMLOS</u>						
N 208743	001	10996208	Apr 30, 2038	DP	M-270	Sep 20, 2024
		7803770	Apr 28, 2031	U-2009	NCE	Apr 28, 2022
		8148333	Nov 08, 2027	DP		
		8748382	Oct 03, 2027	U-2009		
<u>ABAMETAPIR - XEGLYZE</u>						
N 206966	001	10292389	Dec 17, 2034	DP U-2863	NCE	Jul 24, 2025
		7812163	Oct 28, 2026	DP U-2863		
		8212038	Jul 16, 2024	DP U-2863		
		9357783	Jul 16, 2024	DP		
		9839631	Jul 16, 2024	DS DP U-2863		
<u>ABEMACICLIB - VERZENIO</u>						
N 208716	001	7855211	Dec 15, 2029	DS DP U-2132	I-877	Oct 12, 2024
		7855211	Dec 15, 2029	DS DP U-2135	NCE	Sep 28, 2022
		7855211	Dec 15, 2029	DS DP U-2251	NPP	Oct 12, 2024
		7855211	Dec 15, 2029	DS DP U-3241		
		7855211	Dec 15, 2029	DS DP U-3242		
		7855211	Dec 15, 2029	DS DP U-3243		
<u>ABEMACICLIB - VERZENIO</u>						
N 208716	002	7855211	Dec 15, 2029	DS DP U-2132	I-877	Oct 12, 2024
		7855211	Dec 15, 2029	DS DP U-2135	NCE	Sep 28, 2022
		7855211	Dec 15, 2029	DS DP U-2251	NPP	Oct 12, 2024
		7855211	Dec 15, 2029	DS DP U-3241		
		7855211	Dec 15, 2029	DS DP U-3242		
		7855211	Dec 15, 2029	DS DP U-3243		
<u>ABEMACICLIB - VERZENIO</u>						
N 208716	003	7855211	Dec 15, 2029	DS DP U-2132	I-877	Oct 12, 2024
		7855211	Dec 15, 2029	DS DP U-2135	NCE	Sep 28, 2022
		7855211	Dec 15, 2029	DS DP U-2251	NPP	Oct 12, 2024
		7855211	Dec 15, 2029	DS DP U-3241		
		7855211	Dec 15, 2029	DS DP U-3242		
		7855211	Dec 15, 2029	DS DP U-3243		
<u>ABEMACICLIB - VERZENIO</u>						
N 208716	004	7855211	Dec 15, 2029	DS DP U-1981	I-877	Oct 12, 2024
		7855211	Dec 15, 2029	DS DP U-2132	NCE	Sep 28, 2022
		7855211	Dec 15, 2029	DS DP U-2135	NPP	Oct 12, 2024
		7855211	Dec 15, 2029	DS DP U-2251		
<u>ABIRATERONE ACETATE - YONSA</u>						
N 210308	001	10292990	May 20, 2034	U-2535		
		9889144	Mar 17, 2034	DP		
<u>ACALABRUTINIB - CALOQUENCE</u>						
N 210259	001	10167291	Jul 01, 2036	DP U-2145	I-817	Nov 21, 2022
		10167291	Jul 01, 2036	DP U-2666	NCE	Oct 31, 2022
		10167291	Jul 01, 2036	DP U-2667	ODE-175	Oct 31, 2024
		10167291	Jul 01, 2036	DP U-2668	ODE-274	Nov 21, 2026
		10167291	Jul 01, 2036	DP U-2669		
		10167291	Jul 01, 2036	DP U-2670		
		10167291	Jul 01, 2036	DP U-2671		
		10239883	Jul 11, 2032	U-2666		
		10239883	Jul 11, 2032	U-2668		
		10272083	Jan 21, 2035	U-2519		
		10272083	Jan 21, 2035	U-2682		
		10272083	Jan 21, 2035	U-2683		
		10272083	Jan 21, 2035	U-2684		
		10272083	Jan 21, 2035	U-2685		
		10272083	Jan 21, 2035	U-2686		
		10272083	Jan 21, 2035	U-2687		
		7459554	Nov 24, 2026	DS		
		9290504	Jul 11, 2032	DS DP		
		9758524	Jul 11, 2032	U-2145		

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<u>ACALABRUTINIB - CALQUENCE</u>						
N 210259	001	9796721	Jul 01, 2036	DS DP	U-2145	
		9796721	Jul 01, 2036	DS DP	U-2666	
		9796721	Jul 01, 2036	DS DP	U-2667	
		9796721	Jul 01, 2036	DS DP	U-2668	
		9796721	Jul 01, 2036	DS DP	U-2669	
		9796721	Jul 01, 2036	DS DP	U-2670	
		9796721	Jul 01, 2036	DS DP	U-2671	
<u>ACETAMINOPHEN - OFIRMEV</u>						
N 022450	001	10383834	Nov 13, 2028		U-2262	
		10383834	Nov 13, 2028		U-2621	
		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 - ACETAMINOPHEN</u>						
N 204767	001	8741959	Apr 19, 2030	DP		
<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		
<u>ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE - APADAZ</u>						
N 208653	002	8461137	Feb 22, 2031	DS DP		
		8748413	Jul 10, 2030	DS DP		
		8828978	Jul 01, 2030	DP		
		9132125	Jul 01, 2030	DS DP	U-2249	
		9549923	Jul 01, 2030	DS DP		
<u>ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE - APADAZ</u>						
N 208653	003	8461137	Feb 22, 2031	DS DP		
		8748413	Jul 10, 2030	DS DP		
		8828978	Jul 01, 2030	DP		
		9132125	Jul 01, 2030	DS DP	U-2249	
		9549923	Jul 01, 2030	DS DP		
<u>ACETAMINOPHEN; IBUPROFEN - ADVIL DUAL ACTION WITH ACETAMINOPHEN</u>						
N 211733	001				NP	Feb 28, 2023
<u>ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE - XARTEMIS XR</u>						
N 204031	001	7976870	Jun 01, 2027		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		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		U-1499	
<u>ACETYLCYSTEINE - ACETADOTE</u>						
N 021539	001	8148356	May 21, 2026	DP		
		8399445	Aug 24, 2025		U-1373	
		8653061	Aug 24, 2025		U-1373	
		8722738	Apr 06, 2032		U-1373	
		9327028	Jul 21, 2031		U-1839	
<u>ACETYLCYSTEINE - CETYLEV</u>						
N 207916	001	8747894	May 08, 2032	DP	U-1373	
		9427421	May 08, 2032	DP		

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<u>ACETYLCYSTEINE - CETYLEV</u>						
N 207916	001	9561204	May 08, 2032	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	U-1373		
<u>ACLIDINIUM BROMIDE - TUDORZA PRESSAIR</u>						
N 202450	001	10085974	Mar 13, 2029	DP U-2513	M-256	Mar 29, 2022
		11000517	Mar 13, 2029	DP U-2513		
		6681768	Aug 07, 2022	DP		
		8051851	Apr 22, 2027	DP		
		RE46417	Feb 10, 2025	DS DP U-2513		
<u>ACLIDINIUM BROMIDE; FORMOTEROL FUMARATE - DUAKLIR PRESSAIR</u>						
N 210595	001	10085974	Mar 13, 2029	DP U-2513	NC	Mar 29, 2022
		11000517	Mar 13, 2029	DP U-2513		
		6681768	Aug 07, 2022	DP		
		8051851	Apr 22, 2027	DP		
		RE46417	Feb 10, 2025	DS DP U-2513		
<u>ACYCLOVIR - AVACLYR</u>						
N 202408	001				ODE-235	Mar 29, 2026
<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 021753	001	7579377	Feb 23, 2025	U-818		
		7737181	Aug 29, 2024	DP		
		7834060	Mar 12, 2023	U-1078		
		7838558	Mar 12, 2023	DP		
		7868044	Mar 12, 2023	U-1078		
		8703820	Mar 12, 2023	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		
<u>ADAPALENE; BENZOYL PEROXIDE - ADAPALENE AND BENZOYL PEROXIDE</u>						
A 209148	001				PC	May 30, 2022
<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		

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<u>ADAPALENE; BENZOYL PEROXIDE - EPIDUO FORTE</u>						
N 207917	001 9814690	Dec 23, 2022	DP U-1078			
<u>AFAMELANOTIDE - SCENESSE</u>						
N 210797	001 10076555	Feb 11, 2025	U-2638		NCE	Oct 08, 2024
	8334265	Mar 11, 2029	U-2638		ODE-270	Oct 08, 2026
<u>AFATINIB DIMALEATE - GILOTRIF</u>						
N 201292	001 10004743	Jul 05, 2030	DP		ODE-115	Apr 15, 2023
	8426586	Oct 10, 2029	DS		ODE-230	Jan 12, 2025
	8545884	Dec 19, 2029	DP			
	9539258	Nov 09, 2026	U-1950			
	RE43431	Jan 13, 2026	DS			
<u>AFATINIB DIMALEATE - GILOTRIF</u>						
N 201292	002 10004743	Jul 05, 2030	DP		ODE-115	Apr 15, 2023
	8426586	Oct 10, 2029	DS		ODE-230	Jan 12, 2025
	8545884	Dec 19, 2029	DP			
	9539258	Nov 09, 2026	U-1950			
	RE43431	Jan 13, 2026	DS			
<u>AFATINIB DIMALEATE - GILOTRIF</u>						
N 201292	003 10004743	Jul 05, 2030	DP		ODE-115	Apr 15, 2023
	8426586	Oct 10, 2029	DS		ODE-230	Jan 12, 2025
	8545884	Dec 19, 2029	DP			
	9539258	Nov 09, 2026	U-1950			
	RE43431	Jan 13, 2026	DS			
<u>AIR POLYMER-TYPE A - EXEM FOAM KIT</u>						
N 212279	001 9034300	Oct 15, 2030	DP U-2663		NCE	Nov 07, 2024
	9259494	May 04, 2035	DP U-2663			
	9849199	Feb 11, 2036	DP			
<u>ALBUTEROL SULFATE - VENTOLIN HFA</u>						
N 020983	001 7500444	Feb 26, 2026	DP			
	7500444*PED	Aug 26, 2026				
	7832351	Jun 19, 2023	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			
	10561808	Jan 01, 2032	DP			
	10695512	May 18, 2031	DP			
	7105152	Sep 12, 2023	DP			
	8132712	Sep 07, 2028	DP			
	9463289	May 18, 2031	DP			
	9808587	May 18, 2031	DP			
<u>ALBUTEROL SULFATE - PROAIR RESPICLICK</u>						
N 205636	001 10022510	May 18, 2031	DP			
	10124131	May 18, 2031	DP			
	10561808	Jan 01, 2032	DP			
	10765820	May 19, 2025	DP			
	7540282	May 06, 2023	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			
<u>ALBUTEROL SULFATE - PROAIR DIGIHALER</u>						
N 205636	002 10022510	May 18, 2031	DP			
	10124131	May 18, 2031	DP			
	10561808	Jan 01, 2032	DP			
	10569034	Aug 16, 2036	DP			
	10765820	May 19, 2025	DP			
	10918816	Dec 14, 2035	DP			
	11000653	Dec 18, 2038	DP			
	11173259	Jul 06, 2040	DP			
	7540282	May 06, 2023	DP			

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<u>ALBUTEROL SULFATE - PROAIR DIGIHALER</u>						
N 205636	002	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 - COMBIVENT RESPIMAT</u>						
N 021747	001	7284474	Aug 26, 2024	DP		
		7396341	Oct 10, 2026	DP		
		7837235	Mar 13, 2028	DP		
		7896264	May 26, 2025	DP		
		8733341	Oct 16, 2030	DP		
		9027967	Mar 31, 2027	DP		
<u>ALCAFTADINE - LASTACAFT</u>						
N 022134	001	10617695	Mar 19, 2027	DP	U-3267	
		8664215	Dec 23, 2027		U-3267	
<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	10350214	Apr 24, 2035	DP	ODE-105	Dec 11, 2022
		9126931	May 29, 2031	DS	ODE-159	Nov 06, 2024
		9365514	Mar 04, 2032	DP		
		9440922	Jun 09, 2030	DP		
<u>ALENDRONATE SODIUM - BINOSTO</u>						
N 202344	001	7488496	Aug 11, 2023	DS DP		
		7964212	Mar 06, 2023	DS DP		
		9592195	Dec 05, 2031	DP		
<u>ALISKIREN HEMIFUMARATE - TEKTURNA</u>						
N 021985	001	8617595	Feb 19, 2026	DP		
		8617595*PED	Aug 19, 2026			
<u>ALISKIREN HEMIFUMARATE - TEKTURNA</u>						
N 021985	002	8617595	Feb 19, 2026	DP		
		8617595*PED	Aug 19, 2026			
<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		
<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		
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>						
N 200045	002	8183295	May 16, 2023	DP		
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>						
N 200045	003	8183295	May 16, 2023	DP		
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>						
N 200045	004	8183295	May 16, 2023	DP		

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<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>						
N 200045	005	8183295	May 16, 2023	DP		
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTRUNA HCT</u>						
N 022107	001	8618172	Jul 13, 2028	DP		
		9023893	Mar 03, 2022	DP		
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTRUNA HCT</u>						
N 022107	002	8618172	Jul 13, 2028	DP		
		9023893	Mar 03, 2022	DP		
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTRUNA HCT</u>						
N 022107	003	8618172	Jul 13, 2028	DP		
		9023893	Mar 03, 2022	DP		
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTRUNA HCT</u>						
N 022107	004	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	10183012	Nov 26, 2028		U-2104	
		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	10183012	Nov 26, 2028		U-2104	
		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	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	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	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	7807689	Jun 27, 2028	DS DP	U-1337	
		8173663	Mar 15, 2025		U-1338	
		8288539	Jun 24, 2025	DS		
		8900638	May 24, 2029	DP		

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<u>ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE - KAZANO</u>						
N 203414	002	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	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	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	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	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	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	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>ALPELISIB - PIQRAY</u>						
N 212526	001	8227462	Sep 28, 2030	DS DP	U-2539	
		8476268	Sep 10, 2029	DS DP		NCE May 24, 2024
<u>ALPELISIB - PIQRAY</u>						
N 212526	002	8227462	Sep 28, 2030	DS DP	U-2539	
		8476268	Sep 10, 2029	DS DP		NCE May 24, 2024
<u>ALPELISIB - PIQRAY</u>						
N 212526	003	8227462	Sep 28, 2030	DS DP	U-2539	
		8476268	Sep 10, 2029	DS DP		NCE May 24, 2024
<u>ALVIMOPAN - ENTEREG</u>						
N 021775	001	8946262	Feb 12, 2030		U-1655	
<u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u>						
N 208944	001	10154971	Dec 04, 2034		U-2459	
		10646456	Jun 17, 2034		U-2808	
		11065213	Aug 23, 2038	DP		
		11077073	Aug 23, 2038		U-2106	
		11077073	Aug 23, 2038		U-2224	
		11077073	Aug 23, 2038		U-3180	
		11197835	Dec 02, 2030		U-3270	
		8389578	Jan 22, 2028		U-2105	
		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	

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<u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u>						
N 208944	001	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	ODE-153 Aug 24, 2024
		10646456	Jun 17, 2034		U-2808	
		11065213	Aug 23, 2038	DP		
		11077073	Aug 23, 2038		U-2106	
		11077073	Aug 23, 2038		U-2224	
		11077073	Aug 23, 2038		U-3180	
		11197835	Dec 02, 2030		U-3270	
		8389578	Jan 22, 2028		U-2105	
		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	10213393	Feb 15, 2038		U-20	
		10213394	Feb 15, 2038		U-2497	
		10500170	Feb 15, 2038		U-20	
		10500171	Feb 15, 2038		U-2497	
		10500172	Feb 15, 2038		U-2497	
		10512617	Feb 15, 2038		U-2497	
		8252331	Mar 13, 2030	DP		
		8389578	Jan 22, 2028		U-219	
		8389578	Jan 22, 2028		U-3054	
		8574626	Nov 28, 2025	DP	U-20	
		8796337	Nov 23, 2025		U-219	
		8796337	Nov 23, 2025		U-2497	
		8796337	Nov 23, 2025		U-3054	
		8889740	Nov 23, 2025	DP		
		8895614	Nov 23, 2025	DP		
		8895615	Nov 23, 2025		U-219	
		8895615	Nov 23, 2025		U-3054	
		8895616	Nov 23, 2025		U-219	
		8895616	Nov 23, 2025		U-3054	
		8895617	Nov 23, 2025		U-219	
		8895617	Nov 23, 2025		U-3054	
		8895618	Nov 23, 2025	DP		
		8987333	Nov 23, 2025	DP		
		9072697	Nov 23, 2025		U-219	
		9072697	Nov 23, 2025		U-3054	
<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	002	10213393	Feb 15, 2038		U-20	
		10213394	Feb 15, 2038		U-2497	
		10500170	Feb 15, 2038		U-20	
		10500171	Feb 15, 2038		U-2497	
		10500172	Feb 15, 2038		U-2497	
		10512617	Feb 15, 2038		U-2497	
		8252331	Mar 13, 2030	DP		
		8389578	Jan 22, 2028		U-219	
		8389578	Jan 22, 2028		U-3054	
		8574626	Nov 28, 2025	DP	U-20	
		8796337	Nov 23, 2025		U-219	
		8796337	Nov 23, 2025		U-2497	

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<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	002	8796337				
		8889740				
		8895614				
		8895615				
		8895615				
		8895616				
		8895616				
		8895617				
		8895617				
		8895618				
		8987333				
		9072697				
		9072697				
<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	003	10213393				
		10213394				
		10500170				
		10500171				
		10500172				
		10512617				
		8252331				
		8389578				
		8389578				
		8574626				
		8796337				
		8796337				
		8796337				
		8889740				
		8895614				
		8895615				
		8895615				
		8895616				
		8895616				
		8895617				
		8895617				
		8895618				
		8987333				
		9072697				
		9072697				
<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	004	10213393				
		10213394				
		10500170				
		10500171				
		10500172				
		10512617				
		8252331				
		8389578				
		8389578				
		8574626				
		8796337				
		8796337				
		8796337				
		8889740				
		8895614				
		8895615				
		8895615				
		8895616				
		8895616				
		8895617				
		8895617				
		8895618				
		8987333				
		9072697				
		9072697				

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<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 - RUZURGI</u>						
N 209321	001				ODE-244	May 06, 2026
<u>AMIFAMPRIDINE PHOSPHATE - FIRDAPSE</u>						
N 208078	001	10793893	Apr 07, 2034	U-2956	NCE	Nov 28, 2023
		11060128	Jun 29, 2032	U-2956	ODE-223	Nov 28, 2025
<u>AMIKACIN SULFATE - ARIKAYCE KIT</u>						
N 207356	001	10251900	May 15, 2035	U-2414	ODE-214	Sep 28, 2025
		10751355	May 15, 2035	U-2414	GAIN	Sep 28, 2030
		7718189	Jun 06, 2025	DP U-2415		
		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	10357567	Jan 12, 2038	U-3163		
		11077192	Jan 12, 2038	U-3163		
		11135293	Jan 12, 2038	U-3163		
<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		
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>						
N 022325	003	6869939	May 04, 2022	DP		
		7635773	Mar 13, 2029	DP		
<u>AMISULPRIDE - BARHEMSYS</u>						
N 209510	001	10525033	Mar 10, 2031	DP	NCE	Feb 26, 2025
		9084765	Mar 10, 2031	U-1744		
		9084765	Mar 10, 2031	U-2754		
		9545426	Mar 10, 2031	U-1744		
		9545426	Mar 10, 2031	U-2754		
		9889118	Mar 10, 2031	U-1744		
		9889118	Mar 10, 2031	U-2754		
<u>AMLODIPINE BENZOATE - KATERZIA</u>						
N 211340	001	10695329	Oct 16, 2037	DP		
		10799453	Apr 11, 2039	DP		
		10894039	Oct 06, 2037	U-185		
		10894039	Oct 06, 2037	U-3		
		10952998	Oct 06, 2037	DP		
		10959991	Oct 06, 2037	U-158		
		10959991	Oct 06, 2037	U-39		
<u>AMLODIPINE BESYLATE - AMLODIPINE BESYLATE</u>						
N 022026	001	6828339	Nov 20, 2022	DS		

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<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	10350171	Jun 14, 2038	DP		
		10925835	Jun 14, 2038		U-2410	
		10945960	Jun 14, 2038	DP		
		9408837	Feb 28, 2030		U-2410	
		9662315	May 22, 2029	DP	U-2410	
<u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u>						
N 210045	002	10350171	Jun 14, 2038	DP		
		10925835	Jun 14, 2038		U-2410	
		10945960	Jun 14, 2038	DP		
		9408837	Feb 28, 2030		U-2410	
		9662315	May 22, 2029	DP	U-2410	
<u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u>						
N 210045	003	10350171	Jun 14, 2038	DP		
		10925835	Jun 14, 2038		U-2410	
		10945960	Jun 14, 2038	DP		
		9408837	Feb 28, 2030		U-2410	
		9662315	May 22, 2029	DP	U-2410	
<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>AMOXICILLIN - MOXATAG</u>						
N 050813	001	8299052	May 07, 2027		U-1304	
		8357394	Dec 08, 2026	DP		
		8778924	Dec 08, 2026	DS DP	U-897	

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<u>AMOXICILLIN; OMEPRAZOLE MAGNESIUM; RIFABUTIN - TALICIA</u>						
N 213004	001	10238606	Feb 12, 2034	DP		
		11135172	Feb 12, 2034	DP	U-2660	
		9050263	Feb 12, 2034	DP	U-2660	
		9498445	Feb 12, 2034	DP	U-2660	
		9603806	Feb 12, 2034	DP	U-2660	
<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 ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063	001	6913768	May 24, 2023	DP	U-2025	M-248 Sep 13, 2022
		8846100	Aug 24, 2029	DP		PED Mar 13, 2023
		9173857	May 12, 2026		U-2025	
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063	002	6913768	May 24, 2023	DP	U-2025	M-248 Sep 13, 2022
		8846100	Aug 24, 2029	DP		PED Mar 13, 2023
		9173857	May 12, 2026		U-2025	
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063	003	6913768	May 24, 2023	DP	U-2025	M-248 Sep 13, 2022
		8846100	Aug 24, 2029	DP		PED Mar 13, 2023
		9173857	May 12, 2026		U-2025	
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063	004	6913768	May 24, 2023	DP	U-2025	M-248 Sep 13, 2022
		8846100	Aug 24, 2029	DP		PED Mar 13, 2023
		9173857	May 12, 2026		U-2025	

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<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	001	10130580	Apr 19, 2024	DP		
		10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	002	10130580	Apr 19, 2024	DP		
		10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	003	10130580	Apr 19, 2024	DP		
		10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	004	10130580	Apr 19, 2024	DP		
		10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	005	10130580	Apr 19, 2024	DP		
		10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - 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; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR 5</u>						
N 210526	001	8337890	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		9675704	Mar 15, 2027	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR 10</u>						
N 210526	002	8337890	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		9675704	Mar 15, 2027	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR 15</u>						
N 210526	003	8337890	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		9675704	Mar 15, 2027	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR 20</u>						
N 210526	004	8337890	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		9675704	Mar 15, 2027	DP		
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360	001	10028995	Dec 18, 2034	U-2338	NCE	Dec 21, 2022
		10335451	Dec 16, 2029	U-2581		
		10493124	Dec 18, 2034	U-2679		
		10500247	Dec 16, 2029	U-2680		
		10500247	Dec 16, 2029	U-2681		
		10548943	Dec 16, 2029	U-2739		
		10548943	Dec 16, 2029	U-2740		
		11096983	Dec 18, 2034	U-3211		
		11096983	Dec 18, 2034	U-3212		
		11219662	Jan 06, 2037	U-3262		
		9220745	Dec 18, 2034	U-2217		
		9220745	Dec 18, 2034	U-2218		
		9572856	Jul 18, 2031	U-2221		
		9867863	Dec 16, 2029	U-2231		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360 002	10028995	Dec 18, 2034	U-2338		NCE	Dec 21, 2022
	10335451	Dec 16, 2029	U-2581			
	10493124	Dec 18, 2034	U-2679			
	10500247	Dec 16, 2029	U-2680			
	10500247	Dec 16, 2029	U-2681			
	10548943	Dec 16, 2029	U-2739			
	10548943	Dec 16, 2029	U-2740			
	11096983	Dec 18, 2034	U-3211			
	11096983	Dec 18, 2034	U-3212			
	11219662	Jan 06, 2037	U-3262			
	9220745	Dec 18, 2034	U-2217			
	9220745	Dec 18, 2034	U-2218			
	9572856	Nov 20, 2030	U-2221			
	9867863	Dec 16, 2029	U-2231			
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360 003	11219662	Jan 06, 2037	U-3262		NCE	Dec 21, 2022
<u>ANIDULAFUNGIN - ERAXIS</u>						
N 021632 001					NPP	Sep 22, 2023
<u>ANIDULAFUNGIN - ERAXIS</u>						
N 021632 002					NPP	Sep 22, 2023
<u>APALUTAMIDE - ERLEADA</u>						
N 210951 001	10052314	Sep 23, 2033	U-2381		I-808	Sep 17, 2022
	10052314	Sep 23, 2033	U-2382		NCE	Feb 14, 2023
	10702508	Apr 30, 2038	U-3012			
	10849888	Sep 23, 2033	U-3013			
	8445507	Sep 15, 2030	DS DP U-2237			
	8445507	Sep 15, 2030	DS DP U-2624			
	8802689	Mar 27, 2027	U-2237			
	8802689	Mar 27, 2027	U-2624			
	9388159	Mar 27, 2027	DS DP			
	9481663	Jun 04, 2033	DS			
	9884054	Sep 23, 2033	U-2237			
	9987261	Mar 27, 2027	DP			
<u>APIXABAN - ELIQUIS</u>						
N 202155 001	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	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			
	9326945	Feb 24, 2031	DP			
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875 001	10420763	Jun 11, 2030	DP U-2825		NP	May 21, 2023
	10449146	Apr 19, 2036	U-2825			
	10821074	Aug 07, 2029	DP			
	10888499	Feb 14, 2022	DP			
	10959943	Apr 19, 2036	U-2825			
	11077068	Feb 14, 2022	DP			
	8414922	Dec 16, 2031	DP U-2825			
	8603514	Apr 03, 2024	DP			
	8663687	Feb 02, 2023	DP			
	8765167	Feb 20, 2024	DP			
	8846074	Dec 16, 2031	DP U-2825			
	9044475	Jun 11, 2030	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	001	9283219	Jun 11, 2030	DP U-2825		
		9326981	Jun 11, 2030	U-2825		
		9669019	Jun 11, 2030	DP U-2825		
		9669021	Jun 11, 2030	U-2825		
		9855221	Feb 14, 2022	DP		
		9931305	Feb 14, 2022	DP		
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	002	10420763	Jun 11, 2030	DP U-2825	NP	May 21, 2023
		10449146	Apr 19, 2036	U-2825		
		10821074	Aug 07, 2029	DP		
		10888499	Feb 14, 2022	DP		
		10959943	Apr 19, 2036	U-2825		
		11077068	Feb 14, 2022	DP		
		8414922	Dec 16, 2031	DP U-2825		
		8603514	Apr 03, 2024	DP		
		8663687	Feb 02, 2023	DP		
		8765167	Feb 20, 2024	DP		
		8846074	Dec 16, 2031	DP U-2825		
		9044475	Jun 11, 2030	DP		
		9283219	Jun 11, 2030	DP U-2825		
		9326981	Jun 11, 2030	U-2825		
		9669019	Jun 11, 2030	DP U-2825		
		9669021	Jun 11, 2030	U-2825		
		9855221	Feb 14, 2022	DP		
		9931305	Feb 14, 2022	DP		
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	003	10420763	Jun 11, 2030	DP U-2825	NP	May 21, 2023
		10449146	Apr 19, 2036	U-2825		
		10821074	Aug 07, 2029	DP		
		10888499	Feb 14, 2022	DP		
		10959943	Apr 19, 2036	U-2825		
		11077068	Feb 14, 2022	DP		
		8414922	Dec 16, 2031	DP U-2825		
		8603514	Apr 03, 2024	DP		
		8663687	Feb 02, 2023	DP		
		8765167	Feb 20, 2024	DP		
		8846074	Dec 16, 2031	DP U-2825		
		9044475	Jun 11, 2030	DP		
		9283219	Jun 11, 2030	DP U-2825		
		9326981	Jun 11, 2030	U-2825		
		9669019	Jun 11, 2030	DP U-2825		
		9669021	Jun 11, 2030	U-2825		
		9855221	Feb 14, 2022	DP		
		9931305	Feb 14, 2022	DP		
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	004	10420763	Jun 11, 2030	DP U-2825	NP	May 21, 2023
		10449146	Apr 19, 2036	U-2825		
		10821074	Aug 07, 2029	DP		
		10888499	Feb 14, 2022	DP		
		10959943	Apr 19, 2036	U-2825		
		11077068	Feb 14, 2022	DP		
		8414922	Dec 16, 2031	DP U-2825		
		8603514	Apr 03, 2024	DP		
		8663687	Feb 02, 2023	DP		
		8765167	Feb 20, 2024	DP		
		8846074	Dec 16, 2031	DP U-2825		
		9044475	Jun 11, 2030	DP		
		9283219	Jun 11, 2030	DP U-2825		
		9326981	Jun 11, 2030	U-2825		
		9669019	Jun 11, 2030	DP U-2825		
		9669021	Jun 11, 2030	U-2825		
		9855221	Feb 14, 2022	DP		
		9931305	Feb 14, 2022	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	005	10420763	Jun 11, 2030	DP U-2825	NP	May 21, 2023
		10449146	Apr 19, 2036	U-2825		
		10821074	Aug 07, 2029	DP		
		10888499	Feb 14, 2022	DP		
		10959943	Apr 19, 2036	U-2825		
		11077068	Feb 14, 2022	DP		
		8414922	Dec 16, 2031	DP U-2825		
		8603514	Apr 03, 2024	DP		
		8663687	Feb 02, 2023	DP		
		8765167	Feb 20, 2024	DP		
		8846074	Dec 16, 2031	DP U-2825		
		9044475	Jun 11, 2030	DP		
		9283219	Jun 11, 2030	DP U-2825		
		9326981	Jun 11, 2030	U-2825		
		9669019	Jun 11, 2030	DP U-2825		
		9669021	Jun 11, 2030	U-2825		
		9855221	Feb 14, 2022	DP		
		9931305	Feb 14, 2022	DP		
<u>APREMILAST - OTEZLA</u>						
N 205437	001	10092541	May 29, 2034	U-2403	I-803	Jul 19, 2022
		10092541	May 29, 2034	U-2659	M-257	Apr 10, 2023
		6962940	Mar 19, 2023	U-1504	ODE-248	Jul 19, 2026
		6962940	Mar 19, 2023	U-2656		
		6962940	Mar 19, 2023	U-2657		
		6962940	Mar 19, 2023	U-2658		
		7208516	Mar 19, 2023	U-1505		
		7427638	Feb 16, 2028	DS DP		
		7659302	Mar 19, 2023	U-1505		
		7659302	Mar 19, 2023	U-1595		
		7659302	Mar 19, 2023	U-2657		
		7659302	Mar 19, 2023	U-2658		
		7893101	Dec 09, 2023	DS DP		
		8455536	Mar 19, 2023	U-1505		
		8455536	Mar 19, 2023	U-1595		
		8455536	Mar 19, 2023	U-2657		
		8455536	Mar 19, 2023	U-2658		
		8802717	Mar 19, 2023	U-1561		
		9018243	Mar 19, 2023	U-1505		
		9018243	Mar 19, 2023	U-1595		
		9018243	Mar 19, 2023	U-2656		
		9018243	Mar 19, 2023	U-2657		
		9018243	Mar 19, 2023	U-2658		
		9724330	Mar 19, 2023	U-1561		
		9724330	Mar 19, 2023	U-1595		
		9724330	Mar 19, 2023	U-2656		
		9724330	Mar 19, 2023	U-2657		
		9724330	Mar 19, 2023	U-2658		
		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	I-803	Jul 19, 2022
		10092541	May 29, 2034	U-2659	M-257	Apr 10, 2023
		6962940	Mar 19, 2023	U-1504	ODE-248	Jul 19, 2026
		6962940	Mar 19, 2023	U-2656		
		6962940	Mar 19, 2023	U-2657		
		6962940	Mar 19, 2023	U-2658		
		7208516	Mar 19, 2023	U-1505		
		7427638	Feb 16, 2028	DS DP		
		7659302	Mar 19, 2023	U-1505		
		7659302	Mar 19, 2023	U-1595		
		7659302	Mar 19, 2023	U-2657		
		7659302	Mar 19, 2023	U-2658		
		7893101	Dec 09, 2023	DS DP		
		8455536	Mar 19, 2023	U-1505		
		8455536	Mar 19, 2023	U-1595		
		8455536	Mar 19, 2023	U-2657		
		8455536	Mar 19, 2023	U-2658		

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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	8802717	Mar 19, 2023	U-1561		
		9018243	Mar 19, 2023	U-1505		
		9018243	Mar 19, 2023	U-1595		
		9018243	Mar 19, 2023	U-2656		
		9018243	Mar 19, 2023	U-2657		
		9018243	Mar 19, 2023	U-2658		
		9724330	Mar 19, 2023	U-1561		
		9724330	Mar 19, 2023	U-1595		
		9724330	Mar 19, 2023	U-2656		
		9724330	Mar 19, 2023	U-2657		
		9724330	Mar 19, 2023	U-2658		
		9872854	May 29, 2034	U-2232		
		9872854	May 29, 2034	U-2233		
<u>APREMILAST - OTEZLA</u>						
N 205437	003	10092541	May 29, 2034	U-2403	I-803	Jul 19, 2022
		10092541	May 29, 2034	U-2659	M-257	Apr 10, 2023
		6962940	Mar 19, 2023	U-1504	ODE-248	Jul 19, 2026
		6962940	Mar 19, 2023	U-2656		
		6962940	Mar 19, 2023	U-2657		
		6962940	Mar 19, 2023	U-2658		
		7208516	Mar 19, 2023	U-1505		
		7427638	Feb 16, 2028	DS DP		
		7659302	Mar 19, 2023	U-1505		
		7659302	Mar 19, 2023	U-1595		
		7659302	Mar 19, 2023	U-2657		
		7659302	Mar 19, 2023	U-2658		
		7893101	Dec 09, 2023	DS DP		
		8455536	Mar 19, 2023	U-1505		
		8455536	Mar 19, 2023	U-1595		
		8455536	Mar 19, 2023	U-2657		
		8455536	Mar 19, 2023	U-2658		
		8802717	Mar 19, 2023	U-1561		
		9018243	Mar 19, 2023	U-1505		
		9018243	Mar 19, 2023	U-1595		
		9018243	Mar 19, 2023	U-2656		
		9018243	Mar 19, 2023	U-2657		
		9018243	Mar 19, 2023	U-2658		
		9724330	Mar 19, 2023	U-1561		
		9724330	Mar 19, 2023	U-1595		
		9724330	Mar 19, 2023	U-2656		
		9724330	Mar 19, 2023	U-2657		
		9724330	Mar 19, 2023	U-2658		
		9872854	May 29, 2034	U-2232		
		9872854	May 29, 2034	U-2233		
<u>APREPITANT - EMEND</u>						
N 021549	001	8258132	Sep 26, 2027	DP U-1743		
		8258132	Sep 26, 2027	DP U-901		
<u>APREPITANT - EMEND</u>						
N 021549	002	8258132	Sep 26, 2027	DP U-1743		
		8258132	Sep 26, 2027	DP U-901		
<u>APREPITANT - EMEND</u>						
N 021549	003	8258132	Sep 26, 2027	DP U-1743		
		8258132	Sep 26, 2027	DP U-901		
<u>APREPITANT - EMEND</u>						
N 207865	001	8258132	Sep 26, 2027	DP U-1916		
<u>APREPITANT - CINVANTI</u>						
N 209296	001	10500208	Sep 18, 2035	DP		
		10624850	Sep 18, 2035	U-2161		
		10953018	Sep 18, 2035	U-2161		
		11173118	Sep 18, 2035	DP		
		9561229	Sep 18, 2035	DP U-2161		
		9808465	Sep 18, 2035	U-2161		
		9974742	Sep 18, 2035	DP		
		9974793	Sep 18, 2035	DP		

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<u>APREPITANT - CINVANTI</u>						
N 209296	001 9974794	Sep 18, 2035	DP U-2161			
<u>ARGATROBAN - ARGATROBAN IN SODIUM CHLORIDE</u>						
N 022434	001 7589106	Sep 26, 2027	DP U-1163			
	7687516	Sep 26, 2027	DP U-1164			
<u>ARIPIPIRAZOLE - ABILIFY</u>						
N 021436	001 7053092	Jan 28, 2022		U-839		
	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>ARIPIPIRAZOLE - ABILIFY</u>						
N 021436	002 7053092	Jan 28, 2022		U-839		
	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>ARIPIPIRAZOLE - ABILIFY</u>						
N 021436	003 7053092	Jan 28, 2022		U-839		
	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>ARIPIPIRAZOLE - ABILIFY</u>						
N 021436	004 7053092	Jan 28, 2022		U-839		
	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		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIPRAZOLE - ABILIFY</u>						
N 021436	005	7053092	Jan 28, 2022		U-839	
		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>ARIPIPRAZOLE - ABILIFY</u>						
N 021436	006	7053092	Jan 28, 2022		U-839	
		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>ARIPIPRAZOLE - ABILIFY</u>						
N 021713	001	6977257	Apr 24, 2022		DP	
		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>ARIPIPRAZOLE - ABILIFY</u>						
N 021729	002	7053092	Jan 28, 2022		U-839	
		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>ARIPIPRAZOLE - ABILIFY</u>						
N 021729	003	7053092	Jan 28, 2022		U-839	
		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	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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	9387182				Dec 25, 2023 U-1529
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021729	004	7053092				Jan 28, 2022 U-839
		8017615		DP		Jun 16, 2024
		8017615*PED				Dec 16, 2024
		8580796	DS			Sep 25, 2022
		8580796*PED				Mar 25, 2023
		8642600				Jan 28, 2022 U-1492
		8642600*PED				Jul 28, 2022
		8642760	DS			Sep 25, 2022
		8642760*PED				Mar 25, 2023
		9359302	DS DP			Sep 25, 2022 U-1859
		9387182				Dec 25, 2023 U-1529
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021729	005	7053092				Jan 28, 2022 U-839
		8017615		DP		Jun 16, 2024
		8017615*PED				Dec 16, 2024
		8580796	DS			Sep 25, 2022
		8580796*PED				Mar 25, 2023
		8642600				Jan 28, 2022 U-1492
		8642600*PED				Jul 28, 2022
		8642760	DS			Sep 25, 2022
		8642760*PED				Mar 25, 2023
		9359302	DS DP			Sep 25, 2022 U-1859
		9387182				Dec 25, 2023 U-1529
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021866	001	7115587				Jul 21, 2024 DP U-764
		7115587*PED				Jan 21, 2025
		7550445		DP		Jul 21, 2024
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971	001	10525057				Mar 08, 2034 U-1632
		10525057				Mar 08, 2034 U-2723
		10525057				Mar 08, 2034 U-543
		10980803				Sep 24, 2033 U-1632
		10980803				Sep 24, 2033 U-543
		11154553				Sep 24, 2033 U-1632
		11154553				Sep 24, 2033 U-3245
		11154553				Sep 24, 2033 U-814
		7807680	DP			Oct 19, 2024
		8030313				Oct 19, 2024 U-1632
		8030313				Oct 19, 2024 U-543
		8338427	DP			Mar 15, 2025 U-1633
		8338427	DP			Mar 15, 2025 U-543
		8338428	DP			Aug 06, 2023 U-1633
		8338428	DP			Aug 06, 2023 U-543
		8399469	DS			Jun 29, 2025
		8722679		DP		Oct 19, 2024
		8759351		DP		Aug 06, 2023 U-1530
		8759351		DP		Aug 06, 2023 U-1633
		8993761	DS			Sep 25, 2022
		9089567				Jan 28, 2022 U-543
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971	002	10525057				Mar 08, 2034 U-1632
		10525057				Mar 08, 2034 U-2723
		10525057				Mar 08, 2034 U-543
		10980803				Sep 24, 2033 U-1632
		10980803				Sep 24, 2033 U-543
		11154553				Sep 24, 2033 U-1632
		11154553				Sep 24, 2033 U-3245
		11154553				Sep 24, 2033 U-814
		7807680	DP			Oct 19, 2024
		8030313				Oct 19, 2024 U-1632
		8030313				Oct 19, 2024 U-543
		8338427	DP			Mar 15, 2025 U-1633
		8338427	DP			Mar 15, 2025 U-543

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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	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	003	10525057	Mar 08, 2034		U-1632	
		10525057	Mar 08, 2034		U-2723	
		10525057	Mar 08, 2034		U-543	
		10980803	Sep 24, 2033		U-1632	
		10980803	Sep 24, 2033		U-543	
		11154553	Sep 24, 2033		U-1632	
		11154553	Sep 24, 2033		U-3245	
		11154553	Sep 24, 2033		U-814	
		7807680	Oct 19, 2024	DP		
		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	10525057	Mar 08, 2034		U-1632	
		10525057	Mar 08, 2034		U-2723	
		10525057	Mar 08, 2034		U-543	
		10980803	Sep 24, 2033		U-1632	
		10980803	Sep 24, 2033		U-543	
		11154553	Sep 24, 2033		U-1632	
		11154553	Sep 24, 2033		U-3245	
		11154553	Sep 24, 2033		U-814	
		7807680	Oct 19, 2024	DP		
		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	10441194	Jul 26, 2029		DP	
		10517507	Jun 13, 2032		DP	
		7053092	Jan 28, 2022		U-1529	
		7978064	Sep 14, 2026		DP	
		8017615	Jun 16, 2024		DP	
		8114021	Jun 21, 2030		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	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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 001	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			
	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			
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 002	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	7053092	Jan 28, 2022	U-1529			
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	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			
	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			
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 003	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	7053092	Jan 28, 2022	U-1529			
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	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			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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 003	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
	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
	9941931	Nov 04, 2030	DP			
<u>ARIPIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 004	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	7053092	Jan 28, 2022				U-1529
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	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
	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
	9941931	Nov 04, 2030	DP			
<u>ARIPIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 005	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	7053092	Jan 28, 2022				U-1529
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	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

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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 005	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		
	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		
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 006	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	7053092	Jan 28, 2022		U-1529		
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	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		
	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		
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE LAUROXIL - ARISTADA</u>						
N 207533 001	10112903	Jun 24, 2030	DS	U-543		
	10226458	Mar 19, 2032		U-543		
	10238651	Mar 19, 2035		U-2402		
	10813928	Mar 19, 2035		U-2983		
	11097006	Oct 24, 2033	DP	U-764		
	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		

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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	002	10112903	Jun 24, 2030	DS	U-543	
		10226458	Mar 19, 2032		U-543	
		10238651	Mar 19, 2035		U-2402	
		10813928	Mar 19, 2035		U-2983	
		11097006	Oct 24, 2033	DP	U-764	
		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	
		10226458	Mar 19, 2032		U-543	
		10238651	Mar 19, 2035		U-2402	
		10813928	Mar 19, 2035		U-2402	
		11097006	Oct 24, 2033	DP	U-764	
		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	004	10112903	Jun 24, 2030	DS	U-543	
		10226458	Mar 19, 2032		U-543	
		10238651	Mar 19, 2035		U-2402	
		10813928	Mar 19, 2035		U-2983	
		11097006	Oct 24, 2033	DP	U-764	
		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	
		10112903	Jun 24, 2030	DS	U-543	
		10688091	Aug 17, 2035		DP	
		10849894	Aug 17, 2035		U-543	
		11154552	Aug 17, 2035		DP	
		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					

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ARSENIC TRIOXIDE - TRISENOX</u>						
N 021248	002				ODE-167	Jan 12, 2025
<u>ARTESUNATE - ARTESUNATE</u>						
N 213036	001				NCE ODE-290	May 26, 2025 May 26, 2027
<u>ASCIMINIB HYDROCHLORIDE - SCEMBLIX</u>						
N 215358	001	8829195	May 13, 2033	DS U-3249	NCE	Oct 29, 2026
<u>ASCIMINIB HYDROCHLORIDE - SCEMBLIX</u>						
N 215358	002	8829195	May 13, 2033	DS U-3249	NCE	Oct 29, 2026
<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 - PLENVU</u>						
N 209381	001	10016504	Sep 10, 2033	DP		
		10646512	Mar 25, 2032	DP		
		10780112	Mar 09, 2032	DP		
		10792306	Mar 09, 2032	DP U-2310		
		10918723	Sep 10, 2033	U-2310		
		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 - SECUADO</u>						
N 212268	001	10022445	Jul 25, 2033	DP	NP	Oct 11, 2022
		10583121	Jul 25, 2033	DP U-2763		
		10814002	Jul 25, 2033	DP U-2763		
		11123305	Jul 25, 2033	DP		
		9687474	Jul 25, 2033	DP		
<u>ASENAPINE - SECUADO</u>						
N 212268	002	10022445	Jul 25, 2033	DP	NP	Oct 11, 2022
		10583121	Jul 25, 2033	DP U-2763		
		10814002	Jul 25, 2033	DP U-2763		
		11123305	Jul 25, 2033	DP		
		9687474	Jul 25, 2033	DP		
<u>ASENAPINE - SECUADO</u>						
N 212268	003	10022445	Jul 25, 2033	DP	NP	Oct 11, 2022
		10583121	Jul 25, 2033	DP U-2763		
		10814002	Jul 25, 2033	DP U-2763		
		11123305	Jul 25, 2033	DP		
		9687474	Jul 25, 2033	DP		
<u>ASENAPINE MALEATE - SAPHRIS</u>						
N 022117	001	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	002	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		

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<u>ASENAPINE MALEATE - SAPHRIS</u>						
N 022117	002	8022228*PED	Oct 06, 2026			
<u>ASENAPINE MALEATE - SAPHRIS</u>						
N 022117	003	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	
<u>ASPIRIN; OMEPRAZOLE - YOSPRALA</u>						
N 205103	001	6926907	Feb 28, 2023	DP	U-1902	
		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	
		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; COBICISTAT - EVOTAZ</u>						
N 206353	001	10039718	Oct 06, 2032	DP		
		8148374	Sep 03, 2029	DS DP	U-1279	
<u>ATOGEFANT - OULIPTA</u>						
N 215206	001	10117836	Jan 30, 2035	DP		NCE Sep 28, 2026
		8754096	Jul 19, 2032	DS DP	U-3142	
		9499545	Nov 10, 2031	DS DP	U-3142	
		9850246	Mar 13, 2033	DS		
<u>ATOGEFANT - OULIPTA</u>						
N 215206	002	10117836	Jan 30, 2035	DP		NCE Sep 28, 2026
		8754096	Jul 19, 2032	DS DP	U-3142	
		9499545	Nov 10, 2031	DS DP	U-3142	
		9850246	Mar 13, 2033	DS		
<u>ATOGEFANT - OULIPTA</u>						
N 215206	003	10117836	Jan 30, 2035	DP		NCE Sep 28, 2026
		8754096	Jul 19, 2032	DS DP	U-3142	
		9499545	Nov 10, 2031	DS DP	U-3142	
		9850246	Mar 13, 2033	DS		
<u>ATROPINE SULFATE - ATROPINE SULFATE</u>						
A 212868	001					CGT Jan 26, 2022
<u>ATROPINE SULFATE - ATROPINE SULFATE</u>						
A 212868	002					CGT Jan 26, 2022
<u>ATROPINE SULFATE - ATROPINE SULFATE</u>						
A 212868	003					CGT Jan 26, 2022
<u>ATROPINE SULFATE - ATROPINE SULFATE</u>						
A 215624	001					CGT May 25, 2022
<u>AVACOPAN - TAVNEOS</u>						
N 214487	001	8445515	Feb 03, 2031	DS DP		NCE Oct 07, 2026
		8906938	Dec 21, 2029	DS DP		

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<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>AVAPRITINIB - AYWAKIT</u>						
N 212608	001	9200002	Oct 15, 2034	DS DP U-2726	I-863	Jun 16, 2024
		9200002	Oct 15, 2034	DS DP U-3168	I-864	Jun 16, 2024
		9944651	Oct 15, 2034	DS DP U-2726	NCE	Jan 09, 2025
		9944651	Oct 15, 2034	DS DP U-3168	ODE-356	Jun 16, 2028
		9994575	Oct 15, 2034	DS DP U-2726	ODE-366	Jan 09, 2027
		9994575	Oct 15, 2034	DS DP U-3168		
<u>AVAPRITINIB - AYWAKIT</u>						
N 212608	002	9200002	Oct 15, 2034	DS DP U-2726	I-863	Jun 16, 2024
		9200002	Oct 15, 2034	DS DP U-3168	I-864	Jun 16, 2024
		9944651	Oct 15, 2034	DS DP U-2726	NCE	Jan 09, 2025
		9944651	Oct 15, 2034	DS DP U-3168	ODE-356	Jun 16, 2028
		9994575	Oct 15, 2034	DS DP U-2726	ODE-366	Jan 09, 2027
		9994575	Oct 15, 2034	DS DP U-3168		
<u>AVAPRITINIB - AYWAKIT</u>						
N 212608	003	9200002	Oct 15, 2034	DS DP U-2726	NCE	Jan 09, 2025
		9944651	Oct 15, 2034	DS DP U-2726	ODE-356	Jun 16, 2028
		9994575	Oct 15, 2034	DS DP U-2726	ODE-366	Jan 09, 2027
<u>AVAPRITINIB - AYWAKIT</u>						
N 212608	004	9200002	Oct 15, 2034	DS DP U-3168	I-863	Jun 16, 2024
		9944651	Oct 15, 2034	DS DP U-3168	I-864	Jun 16, 2024
		9994575	Oct 15, 2034	DS DP U-3168	NCE	Jan 09, 2025
					ODE-356	Jun 16, 2028
					ODE-366	Jan 09, 2027
<u>AVAPRITINIB - AYWAKIT</u>						
N 212608	005	9200002	Oct 15, 2034	DS DP U-3168	I-863	Jun 16, 2024
		9944651	Oct 15, 2034	DS DP U-3168	I-864	Jun 16, 2024
		9994575	Oct 15, 2034	DS DP U-3168	NCE	Jan 09, 2025
					ODE-356	Jun 16, 2028
					ODE-366	Jan 09, 2027
<u>AVATROMBOPAG MALEATE - DOPTELET</u>						
N 210238	001	7638536	May 05, 2025	DS DP	I-802	Jun 26, 2022
		8338429	Jun 30, 2023	U-2577	NCE	May 21, 2023
		8765764	Jan 15, 2023	U-2314	ODE-246	Jun 26, 2026
		8765764	Jan 15, 2023	U-2578		
<u>AVIBACTAM SODIUM; CEFTAZIDIME - AVYCAZ</u>						
N 206494	001	7112592	Jan 07, 2026	DS DP U-2244	NCE	Feb 25, 2020
		7112592	Jan 07, 2026	DS DP U-2508	GAIN	Feb 25, 2025
		7112592	Jan 07, 2026	DS DP U-282		
		7612087	Nov 12, 2026	DP		
		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	10570202	Feb 03, 2035	U-2844		
		10869924	Nov 05, 2036	U-3044		
		6534524	Apr 29, 2025	DS DP		
		8791140	Dec 14, 2030	DS		

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<u>AXITINIB - INLYTA</u>						
N 202324	002	10570202	Feb 03, 2035	U-2844		
		10869924	Nov 05, 2036	U-3044		
		6534524	Apr 29, 2025	DS DP		
		8791140	Dec 14, 2030	DS		
<u>AZACITIDINE - ONUREG</u>						
N 214120	001	8846628	Jun 03, 2030	DP U-2950	NP ODE-320	Sep 01, 2023 Sep 01, 2027
<u>AZACITIDINE - ONUREG</u>						
N 214120	002	8846628	Jun 03, 2030	DP U-2950	NP ODE-320	Sep 01, 2023 Sep 01, 2027
<u>AZELAIC ACID - FINACEA</u>						
N 207071	001	10117812	Oct 18, 2027	DP U-1796		
		10322085	Oct 24, 2023	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		
<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 - ASTEPRO ALLERGY</u>						
N 213872	001	8071073	Jun 04, 2028	DP		
		8518919	Nov 22, 2025	U-3166		
		9919050	Nov 22, 2025	DP		
<u>AZELASTINE HYDROCHLORIDE - CHILDREN'S ASTEPRO ALLERGY</u>						
N 213872	002	8071073	Jun 04, 2028	DP		
		8518919	Nov 22, 2025	U-3166		
		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		
		9387249	Jul 01, 2031	U-3		

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<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		
		9387249	Jul 01, 2031		U-3	
<u>AZITHROMYCIN - ZMAX</u>						
N 050797	001	6984403	Feb 14, 2024	DP	U-282	
		7887844	Feb 14, 2024	DP		
<u>BACLOFEN - OZOBAX</u>						
N 208193	001	10610502	Aug 30, 2039		U-2779	
<u>BACLOFEN - LYVISPAH</u>						
N 215422	001	10792262	Jul 29, 2039	DP	U-3263	
<u>BACLOFEN - LYVISPAH</u>						
N 215422	002	10792262	Jul 29, 2039	DP	U-3263	
<u>BACLOFEN - LYVISPAH</u>						
N 215422	003	10792262	Jul 29, 2039	DP	U-3263	
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 210854	001	10392406	Apr 27, 2036	DS		I-811 Oct 16, 2022
		10633397	Apr 27, 2036		U-2816	NCE Oct 24, 2023
		10633397	Apr 27, 2036		U-3000	
		10759814	Aug 09, 2037	DS DP		
		8927710	May 05, 2031	DP		
		8987441	Sep 21, 2031	DS DP		
		9815835	Jun 14, 2030	DP		
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 210854	002	10392406	Apr 27, 2036	DS		I-811 Oct 16, 2022
		10633397	Apr 27, 2036		U-2816	NCE Oct 24, 2023
		10633397	Apr 27, 2036		U-3000	
		10759814	Aug 09, 2037	DS DP		
		8927710	May 05, 2031	DP		
		8987441	Sep 21, 2031	DS DP		
		9815835	Jun 14, 2030	DP		
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 210854	003	10392406	Apr 27, 2036	DS		I-811 Oct 16, 2022
		10633397	Apr 27, 2036		U-2816	NCE Oct 24, 2023
		10633397	Apr 27, 2036		U-3000	
		10759814	Aug 09, 2037	DS DP		
		8927710	May 05, 2031	DP		
		8987441	Sep 21, 2031	DS DP		
		9815835	Jun 14, 2030	DP		
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 214410	001	10392406	Apr 27, 2036	DS		NCE Oct 24, 2023
		10633397	Apr 27, 2036		U-2816	
		10633397	Apr 27, 2036		U-3000	
		10759814	Aug 09, 2037	DS DP		
		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	
<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	

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<u>BARICITINIB - OLUMIANT</u>						
N 207924	001	8158616	Jun 08, 2030	DS DP	NCE	May 31, 2023
		8420629	Mar 10, 2029	U-247		
<u>BARICITINIB - OLUMIANT</u>						
N 207924	002	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	6479535	May 06, 2024	DP U-594		
		6479535	May 06, 2024	DP U-904		
		7683051	Mar 10, 2027	DS DP U-594		
		7683051	Mar 10, 2027	DS DP U-904		
<u>BECLOMETHASONE DIPROPIONATE - QVAR 80</u>						
N 020911	001	10022509	May 18, 2031	DP		
		10022510	May 18, 2031	DP		
		10086156	May 18, 2031	DP		
		10561808	Jan 01, 2032	DP		
		10695512	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		
		10561808	Jan 01, 2032	DP		
		10695512	May 18, 2031	DP		
		9463289	May 18, 2031	DP		
		9808587	May 18, 2031	DP		
<u>BECLOMETHASONE DIPROPIONATE - QNASL</u>						
N 202813	001	10188811	Oct 21, 2031	DP		
		7780038	Jan 24, 2027	DP		
<u>BECLOMETHASONE DIPROPIONATE - QNASL</u>						
N 202813	002	10188811	Oct 21, 2031	DP		
		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		
		10561808	Jan 01, 2032	DP		
		10695512	May 18, 2031	DP		
		10792447	Jan 25, 2039	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		
		10561808	Jan 01, 2032	DP		
		10695512	May 18, 2031	DP		
		10792447	Jan 25, 2039	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	NPP	Aug 09, 2022
		8546428	Mar 19, 2029	DS DP U-1321	ODE-251	Aug 09, 2026
					ODE-307	May 27, 2027
<u>BEDAQUILINE FUMARATE - SIRTURO</u>						
N 204384	002	7498343	Dec 01, 2026	DS DP U-1321	ODE-307	May 27, 2027
		8546428	Mar 19, 2029	DS DP U-1321		

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<u>BELINOSTAT - BELEODAQ</u>						
N 206256	001 6888027	Aug 10, 2026	DS DP U-1544			
	8835501	Oct 27, 2027	DP			
<u>BELUMOSUDIL MESYLATE - REZUROCK</u>						
N 214783	001 10183931	Oct 07, 2033		U-3246	NCE	Jul 16, 2026
	10696660	Oct 07, 2033		U-3246	ODE-362	Jul 16, 2028
	8357693	Oct 30, 2029	DS DP U-3247			
	9815820	Oct 07, 2033	U-3247			
<u>BELZUTIFAN - WELIREG</u>						
N 215383	001 9908845	Sep 05, 2034	DS DP U-3201		NCE	Aug 13, 2026
	9969689	Sep 05, 2034	DS DP U-3201		ODE-364	Aug 13, 2028
<u>BEMPEDOIC ACID - NEXLETOL</u>						
N 211616	001 10118881	Dec 23, 2023		U-2747	NCE	Feb 21, 2025
	10941095	Dec 23, 2023		U-2747		
	7335799	Dec 03, 2025	DS			
	8497301	Dec 23, 2023		U-2747		
	9000041	Dec 23, 2023		U-2747		
	9624152	Dec 23, 2023		U-2748		
<u>BEMPEDOIC ACID; EZETIMIBE - NEXLIZET</u>						
N 211617	001 10118881	Dec 23, 2023		U-2746	NCE	Feb 21, 2025
	10912751	Mar 14, 2036		U-3224	NP	Feb 26, 2023
	10941095	Dec 23, 2023		U-2746		
	7335799	Dec 03, 2025	DS			
	8497301	Dec 23, 2023		U-2746		
	9000041	Dec 23, 2023		U-2746		
	9624152	Dec 23, 2023		U-2749		
<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		
	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			

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<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N 022249	003	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		
		11103483	Jan 28, 2031	DP U-1971		
		11103483	Jan 28, 2031	DP U-1972		
		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		
		11103483	Jan 28, 2031	DP U-1971		
		11103483	Jan 28, 2031	DP 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		
		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>BENOXINATE HYDROCHLORIDE; FLUORESCEIN SODIUM - FLUORESCEIN SODIUM AND BENOXINATE HYDROCHLORIDE</u>						
N 211039	001	10293047	Nov 15, 2037	DP U-2755		
		10632197	Nov 15, 2037	DP U-2755		
		10842872	Nov 15, 2037	U-3001		
<u>BENZNIDAZOLE - BENZNIDAZOLE</u>						
N 209570	001				NCE	Aug 29, 2022
					ODE-154	Aug 29, 2024
<u>BENZNIDAZOLE - BENZNIDAZOLE</u>						
N 209570	002				NCE	Aug 29, 2022
					ODE-154	Aug 29, 2024
<u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ACANYA</u>						
N 050819	001	10220049	Jun 03, 2029	DP U-916		
		10624918	Jun 03, 2029	U-916		
		8288434	Aug 05, 2029	DP U-124		
		8663699	Jun 03, 2029	U-124		

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<u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ACANYA</u>						
N 050819	001	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		
		10220049	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>BENZOYL PEROXIDE; TRETINOIN - TWYNEO</u>						
N 214902	001	10420743	Jul 12, 2038	U-3194	NC	Jul 26, 2024
		10653899	Dec 30, 2030	DP U-3194		
		11071878	Dec 30, 2030	DP		
		8617580	Feb 03, 2028	DP		
		9868103	Aug 08, 2028	DP U-3194		
<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	8784789	Jan 13, 2025	DP		
		8877168	Jul 30, 2023	DP		
<u>BEROTRALSTAT HYDROCHLORIDE - ORLADEYO</u>						
N 214094	001	10125102	Apr 07, 2035	DS U-3010	NCE	Dec 03, 2025
		10329260	Mar 09, 2035	DS	ODE-333	Dec 03, 2027
		10662160	Nov 01, 2039	DS U-3010		
		10689346	Mar 09, 2035	U-3010		
		11117867	Nov 01, 2039	DP U-3010		
<u>BEROTRALSTAT HYDROCHLORIDE - ORLADEYO</u>						
N 214094	002	10125102	Apr 07, 2035	DS U-3010	NCE	Dec 03, 2025
		10329260	Mar 09, 2035	DS	ODE-333	Dec 03, 2027
		10662160	Nov 01, 2039	DS U-3010		
		10689346	Mar 09, 2035	U-3010		
		11117867	Nov 01, 2039	DP U-3010		
<u>BESIFLOXACIN HYDROCHLORIDE - BESIVANCE</u>						
N 022308	001	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	10179137	Aug 31, 2030	DP U-1858		
		9364485	Aug 31, 2030	DP U-1858		
		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 - TACLONEX</u>						
N 022185	001				NPP	Jul 25, 2022
					PED	Jan 25, 2023
<u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE - ENSTILAR</u>						
N 207589	001	10130640	Jun 10, 2031	DP	NPP	Jul 30, 2022
		10130640*PED	Dec 10, 2031		PED	Jan 30, 2023
		10617698	Jun 10, 2031	DP		

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<u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE - ENSTILAR</u>						
N 207589	001	10660908	Jun 10, 2031	DP	U-2627	
		10682364	Jun 10, 2031	DP		
		10688108	Jun 10, 2031		U-2627	
		10716799	Jun 10, 2031	DP		
		9119781	Jun 10, 2031	DP	U-1761	
		9119781	Jun 10, 2031	DP	U-2627	
		9119781*PED	Dec 10, 2031			
		9566286	Jun 10, 2031	DP		
<u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE - WYNZORA</u>						
N 213422	001	10265265	Sep 27, 2027	DP		NDF Jul 20, 2023
<u>BETRIXABAN - BEVYXXA</u>						
N 208383	001	7598276	Nov 08, 2026	DS		NCE Jun 23, 2022
		8404724	Mar 29, 2031	DP	U-2034	
		8557852	Sep 08, 2028		U-1167	
		8557852	Sep 08, 2028		U-2030	
		8987463	Dec 28, 2030	DP		
		9555023	Nov 07, 2026		U-1502	
<u>BETRIXABAN - BEVYXXA</u>						
N 208383	002	7598276	Nov 08, 2026	DS		NCE Jun 23, 2022
		8404724	Mar 29, 2031	DP	U-2034	
		8557852	Sep 08, 2028		U-1167	
		8557852	Sep 08, 2028		U-2030	
		8987463	Dec 28, 2030	DP		
		9555023	Nov 07, 2026		U-1502	
<u>BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - BIKTARVY</u>						
N 210251	001	10385067	Jun 19, 2035		U-257	M-82 Feb 24, 2024
		10548846	Nov 08, 2036	DP		NCE Feb 07, 2023
		7390791	Apr 17, 2025	DS DP		NPP Jun 18, 2022
		7390791*PED	Oct 17, 2025			ODE-256 Jun 18, 2026
		7803788	Feb 02, 2022		U-257	
		8754065	Aug 15, 2032	DS DP	U-257	
		8754065*PED	Feb 15, 2033			
		9216996	Dec 19, 2033	DS DP		
		9296769	Aug 15, 2032	DS DP	U-257	
		9296769*PED	Feb 15, 2033			
		9708342	Jun 19, 2035	DS DP		
		9732092	Dec 19, 2033	DS DP		
<u>BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - BIKTARVY</u>						
N 210251	002	10385067	Jun 19, 2035		U-257	
		7390791	Apr 17, 2025	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	
		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	

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<u>BIMATOPROST - LATISSE</u>						
N 022369	001	8101161	May 25, 2024	U-1218		
		8263054	Jan 15, 2023	U-1277		
		8632760	Jan 15, 2023	U-1487		
		8758733	Jan 15, 2023	U-1487		
		8986715	Jan 15, 2023	U-1217		
		9216183	Jan 15, 2023	U-1487		
		9226931	Jan 15, 2023	U-1799		
<u>BIMATOPROST - DURYSTA</u>						
N 211911	001	10398707	Apr 30, 2024	U-2759	NP	Mar 04, 2023
		10441543	Dec 19, 2026	DP		
		7799336	Apr 24, 2029	DP		
		8206737	Apr 07, 2027	U-2759		
		8629185	Jul 15, 2031	DS DP		
		8673341	Feb 19, 2025	U-2759		
		9149428	Dec 19, 2026	DP		
		9492316	Oct 31, 2034	DP		
		9980974	Oct 31, 2034	U-2759		
<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		
		6713446*PED	Jul 25, 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				ODE*	Sep 05, 2024
<u>BOSENTAN - TRACLEER</u>						
N 021290	002				ODE*	Sep 05, 2024
<u>BOSENTAN - TRACLEER</u>						
N 209279	001	7959945	Dec 28, 2027	DP	ODE-161	Sep 05, 2024
		8309126	May 15, 2026	DP		
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341	001	11103497	Feb 28, 2034	U-3216	ODE-163	Dec 19, 2024
		11103497	Feb 28, 2034	U-3217		
		7417148	Dec 11, 2025	U-1283		
		7767678	Nov 23, 2026	DS DP		
		7919625	Dec 11, 2025	DP		
		RE42376	Apr 13, 2024	DS		

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<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341	002	11103497	Feb 28, 2034	U-3216	ODE-163	Dec 19, 2024
		11103497	Feb 28, 2034	U-3217		
		7417148	Dec 11, 2025	U-1283		
		7767678	Nov 23, 2026	DS DP		
		7919625	Dec 11, 2025	DP		
		RE42376	Apr 13, 2024	DS		
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341	003	11103497	Feb 28, 2034	U-3216	ODE-163	Dec 19, 2024
		11103497	Feb 28, 2034	U-3217		
		7417148	Dec 11, 2025	U-1283		
		7767678	Nov 23, 2026	DS DP		
		7919625	Dec 11, 2025	DP		
		RE42376	Apr 13, 2024	DS		
<u>BREMELANOTIDE ACETATE - VYLEESI (AUTOINJECTOR)</u>						
N 210557	001	10286034	Nov 05, 2033	U-2568	NCE	Jun 21, 2024
		6794489	Jun 28, 2025	DS DP		
		9352013	Nov 05, 2033	U-2568		
		9700592	Nov 05, 2033	U-2568		
<u>BREXANOLONE - ZULRESSO</u>						
N 211371	001	10117951	Mar 13, 2029	DP	NCE	Jun 17, 2024
		10251894	Nov 27, 2033	U-2552		
		10322139	Jan 23, 2033	DP		
		10940156	Mar 08, 2037	U-2552		
		7635773	Mar 13, 2029	DP		
		8410077	Mar 13, 2029	DP		
		9200088	Mar 13, 2029	DP		
		9750822	Mar 13, 2029	DP		
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422	001	10307419	Oct 12, 2032	DP		
		7888362	Apr 12, 2026	DS	Y	
		8349840	Apr 12, 2026	DP U-1529		
		8618109	Apr 12, 2026	U-543		
		9839637	Apr 12, 2026	DP U-1529		
		9839637	Apr 12, 2026	DP U-543		
		RE48059	Dec 23, 2028	DS		
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422	002	10307419	Oct 12, 2032	DP		
		7888362	Apr 12, 2026	DS	Y	
		8349840	Apr 12, 2026	DP U-1529		
		8618109	Apr 12, 2026	U-543		
		9839637	Apr 12, 2026	DP U-1529		
		9839637	Apr 12, 2026	DP U-543		
		RE48059	Dec 23, 2028	DS		
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422	003	10307419	Oct 12, 2032	DP		
		7888362	Apr 12, 2026	DS	Y	
		8349840	Apr 12, 2026	DP U-1529		
		8618109	Apr 12, 2026	U-543		
		9839637	Apr 12, 2026	DP U-1529		
		9839637	Apr 12, 2026	DP U-543		
		RE48059	Dec 23, 2028	DS		
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422	004	10307419	Oct 12, 2032	DP		
		7888362	Apr 12, 2026	DS	Y	
		8349840	Apr 12, 2026	DP U-1529		
		8618109	Apr 12, 2026	U-543		
		9839637	Apr 12, 2026	DP U-1529		
		9839637	Apr 12, 2026	DP U-543		
		RE48059	Dec 23, 2028	DS		
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422	005	10307419	Oct 12, 2032	DP		
		7888362	Apr 12, 2026	DS	Y	

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<u>BREXPIRAZOLE - REXULTI</u>						
N 205422	005	8349840	Apr 12, 2026	DP U-1529		
		8618109	Apr 12, 2026	U-543		
		9839637	Apr 12, 2026	DP U-1529		
		9839637	Apr 12, 2026	DP U-543		
		RE48059	Dec 23, 2028	DS		
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422	006	7888362	Apr 12, 2026	DS		Y
		8349840	Apr 12, 2026	DP U-1529		
		8618109	Apr 12, 2026	U-543		
		9839637	Apr 12, 2026	DP U-1529		
		9839637	Apr 12, 2026	DP U-543		
		RE48059	Dec 23, 2028	DS		
<u>BRIGATINIB - ALUNBRIG</u>						
N 208772	001	10385078	Nov 10, 2035	DS DP U-2837	I-847	May 22, 2023
		9012462	Jul 31, 2030	DS	NCE	Apr 28, 2022
		9273077	May 21, 2029	U-2837	ODE-142	Apr 28, 2024
		9611283	Apr 10, 2034	U-2837	ODE-300	May 22, 2027
<u>BRIGATINIB - ALUNBRIG</u>						
N 208772	002	10385078	Nov 10, 2035	DS DP U-2837	I-847	May 22, 2023
		9012462	Jul 31, 2030	DS	NCE	Apr 28, 2022
		9273077	May 21, 2029	U-2837	ODE-142	Apr 28, 2024
		9611283	Apr 10, 2034	U-2837	ODE-300	May 22, 2027
<u>BRIGATINIB - ALUNBRIG</u>						
N 208772	003	10385078	Nov 10, 2035	DS DP U-2837	I-847	May 22, 2023
		9012462	Jul 31, 2030	DS	NCE	Apr 28, 2022
		9273077	May 21, 2029	U-2837	ODE-142	Apr 28, 2024
		9611283	Apr 10, 2034	U-2837	ODE-300	May 22, 2027
<u>BRILLIANT BLUE G - TISSUEBLUE</u>						
N 209569	001				NCE	Dec 20, 2024
					ODE-282	Dec 20, 2026
<u>BRIMONIDINE TARTRATE - ALPHAGAN P</u>						
N 021262	001	9295641*PED	Jan 10, 2022			
<u>BRIMONIDINE TARTRATE - OOLIANA</u>						
N 021764	001	7265117	Aug 19, 2025	DP		
<u>BRIMONIDINE TARTRATE - ALPHAGAN P</u>						
N 021770	001	8858961	Sep 02, 2023	DP		
		8858961*PED	Mar 02, 2024			
		9295641*PED	Jan 10, 2022			
		9687443*PED	Jan 10, 2022			
<u>BRIMONIDINE TARTRATE - MIRVASO</u>						
N 204708	001	10201517	Jun 13, 2031	DP		
		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	U-2222		
		9259425	Jul 14, 2030	U-2222		
<u>BRIMONIDINE TARTRATE; BRINZOLAMIDE - SIMBRINZA</u>						
N 204251	001	9044484	Oct 30, 2030	DP		
		9421265	Jun 17, 2030	DP		

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<u>BRIMONIDINE TARTRATE; TIMOLOL MALEATE - COMBIGAN</u>						
N 021398	001	7030149	Apr 19, 2022	U-849		
		7320976	Apr 19, 2022	U-849		
		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>BRINCIDOFIVIR - TEMBEXA</u>						
N 214460	001	9303051	Aug 31, 2031	DS DP U-3165	NP ODE-354	Jun 04, 2024 Jun 04, 2028
<u>BRINCIDOFIVIR - TEMBEXA</u>						
N 214461	001	10112909	Oct 10, 2034	U-3165	NP	Jun 04, 2024
		10487061	Oct 10, 2034	DP U-3165	ODE-354	Jun 04, 2028
		8962829	Oct 10, 2034	DS DP		
		9303051	Aug 31, 2031	DS DP U-3165		
		9371344	Oct 10, 2034	DP		
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	001	10729653	Apr 09, 2030	DP	NPP	Aug 27, 2024
		6911461	Feb 21, 2026	DS DP U-2295		
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	002	10729653	Apr 09, 2030	DP	NPP	Aug 27, 2024
		6911461	Feb 21, 2026	DS DP U-2295		
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	003	10729653	Apr 09, 2030	DP	NPP	Aug 27, 2024
		6911461	Feb 21, 2026	DS DP U-2295		
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	004	10729653	Apr 09, 2030	DP	NPP	Aug 27, 2024
		6911461	Feb 21, 2026	DS DP U-2295		
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	005	10729653	Apr 09, 2030	DP	NPP	Aug 27, 2024
		6911461	Feb 21, 2026	DS DP U-2295		
<u>BRIVARACETAM - BRIVIACT</u>						
N 205837	001	6911461	Feb 21, 2026	DS DP U-1815	NPP	Aug 27, 2024
		6911461	Feb 21, 2026	DS DP U-2130		
<u>BRIVARACETAM - BRIVIACT</u>						
N 205838	001	6911461	Feb 21, 2026	DS DP U-2295	NPP	Aug 27, 2024
<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		
<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
N 020866	001	10688094	Apr 30, 2032	U-2870		
		10688094	Apr 30, 2032	U-2871		
		10688094	Apr 30, 2032	U-2872		
		10688094	Apr 30, 2032	U-2873		

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<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
N 020866 001	10688094	Apr 30, 2032	U-2874			
	10688094	Apr 30, 2032	U-2875			
	10688094	Apr 30, 2032	U-2876			
	10688094	Apr 30, 2032	U-2877			
	10688094	Apr 30, 2032	U-2878			
	10688094	Apr 30, 2032	U-2879			
	10688094	Apr 30, 2032	U-2880			
	10688094	Apr 30, 2032	U-2881			
	10688094	Apr 30, 2032	U-2882			
	10688094	Apr 30, 2032	U-2883			
	10688094	Apr 30, 2032	U-2884			
	10688094	Apr 30, 2032	U-2885			
	10688094	Apr 30, 2032	U-2886			
	10688094	Apr 30, 2032	U-2887			
	10688094	Apr 30, 2032	U-2888			
	10688155	Jun 07, 2030	U-2281			
	10688155	Jun 07, 2030	U-2890			
	10688155	Jun 07, 2030	U-2891			
	10688155	Jun 07, 2030	U-2892			
	10688155	Jun 07, 2030	U-2893			
	10688155	Jun 07, 2030	U-2894			
	10688155	Jun 07, 2030	U-2895			
	10688155	Jun 07, 2030	U-2896			
	10688155	Jun 07, 2030	U-2897			
	10688155	Jun 07, 2030	U-2898			
	10688155	Jun 07, 2030	U-2899			
	10688155	Jun 07, 2030	U-2900			
	10688155	Jun 07, 2030	U-2901			
	10688155	Jun 07, 2030	U-2902			
	10688155	Jun 07, 2030	U-2903			
	10688155	Jun 07, 2030	U-2904			
	10688155	Jun 07, 2030	U-2905			
	10688155	Jun 07, 2030	U-2906			
	10688155	Jun 07, 2030	U-2907			
	10688155	Jun 07, 2030	U-2908			
	10688155	Jun 07, 2030	U-2909			
	10688155	Jun 07, 2030	U-2910			
	10688155	Jun 07, 2030	U-2911			
	10688155	Jun 07, 2030	U-2912			
	10688155	Jun 07, 2030	U-2913			
	10688155	Jun 07, 2030	U-2914			
	10688155	Jun 07, 2030	U-2915			
	10688155	Jun 07, 2030	U-2916			
	10688155	Jun 07, 2030	U-2917			
	10688155	Jun 07, 2030	U-2918			
	10688155	Jun 07, 2030	U-2919			
	10688155	Jun 07, 2030	U-2920			
	10688155	Jun 07, 2030	U-2921			
	10688155	Jun 07, 2030	U-2922			
	10688155	Jun 07, 2030	U-2923			
	10688155	Jun 07, 2030	U-2924			
	10688155	Jun 07, 2030	U-2925			
	10688155	Jun 07, 2030	U-2926			
	10688155	Jun 07, 2030	U-2927			
	10688155	Jun 07, 2030	U-2928			
	10688155	Jun 07, 2030	U-2929			
	10688155	Jun 07, 2030	U-2930			
	10688155	Jun 07, 2030	U-2931			
	10688155	Jun 07, 2030	U-2932			
	10688155	Jun 07, 2030	U-2933			
	10688155	Jun 07, 2030	U-2934			
	10688155	Jun 07, 2030	U-2935			
	10688155	Jun 07, 2030	U-2936			
	10688155	Jun 07, 2030	U-2937			
	11000522	Apr 30, 2032	U-3119			
	11000522	Apr 30, 2032	U-3120			
	11000522	Apr 30, 2032	U-3121			
	11000522	Apr 30, 2032	U-3122			
	7888310	Jul 25, 2023	U-1433			

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<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
N 020866 001	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			
	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 - UCERIS</u>						
N 203634 001	10307375	Sep 07, 2031	DP			
	10660858	Sep 07, 2031	DP			
	8895064	Sep 07, 2031	DP			
	9132093	Sep 07, 2031	DP			
	9192581	Sep 07, 2031	DP U-1325			
<u>BUDESONIDE - ORTIKOS</u>						
N 211929 001	10172802	Sep 09, 2036	U-2554			
	9707182	Sep 09, 2036	DP U-2554			
<u>BUDESONIDE - ORTIKOS</u>						
N 211929 002	10172802	Sep 09, 2036	U-2554			
	9707182	Sep 09, 2036	DP U-2554			

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<u>BUDESONIDE - TARPEYO</u>						
N 215935	001 8491932	May 07, 2029	DP U-3269			
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
N 021929	001 10166247	Jan 29, 2023	DP U-2001			
	10166247	Jan 29, 2023	DP U-2002			
	10166247	Jan 29, 2023	DP U-2122			
	10166247*PED	Jul 29, 2023				
	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*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				
	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			
	10166247	Jan 29, 2023	DP U-2002			
	10166247	Jan 29, 2023	DP U-2122			
	10166247*PED	Jul 29, 2023				
	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*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				
	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; GLYCOPYRROLATE - BREZTRI AEROSPHERE</u>						
N 212122	001 10716753	May 28, 2030	DP U-2889		NC	Jul 23, 2023
	8324266	May 28, 2030	U-2889			
	8703806	May 28, 2030	U-2889			
	8808713	May 28, 2030	DP U-2889			
	8815258	Mar 17, 2031	U-2889			
	9415009	May 28, 2030	U-2889			
	9463161	May 28, 2030	DP U-2889			

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<u>BUPIVACAINE - EXPAREL</u>						
N 022496	001	11033495	Jan 22, 2041	DP U-3182	NPP	Mar 22, 2024
		11179336	Jan 22, 2041	DP U-3250		
<u>BUPIVACAINE - EXPAREL</u>						
N 022496	002	11033495	Jan 22, 2041	DP U-3182	NPP	Mar 22, 2024
		11179336	Jan 22, 2041	DP U-3250		
<u>BUPIVACAINE - POSIMIR</u>						
N 204803	001	8153149	Sep 15, 2025	DP	NP	Feb 01, 2024
		8153661	Sep 15, 2025	U-3074		
		8753665	Sep 15, 2025	DP U-3074		
		8846072	Sep 15, 2025	DP U-3074		
<u>BUPIVACAINE HYDROCHLORIDE - XARACOLL</u>						
N 209511	001	RE47826	May 20, 2029	U-2949	NP	Aug 28, 2023
<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988	001	10098957	Apr 20, 2035	U-3118	NP	May 12, 2024
		10213510	Apr 20, 2035	DP U-3118		
		10398686	Mar 13, 2034	DP		
		10632199	Apr 20, 2035	DP U-3118		
		10898575	Apr 20, 2035	DP U-3118		
		10980886	Apr 20, 2035	DP		
		11083730	Apr 20, 2035	DP U-3118		
		11083797	Apr 20, 2035	DP U-3118		
		9592227	Mar 13, 2034	DP U-3118		
		9694079	Apr 20, 2035	DP U-3118		
		9744163	Mar 13, 2034	DP		
		9801945	Apr 20, 2035	DP U-3118		
		9913909	Mar 13, 2034	U-3118		
<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988	002	10098957	Apr 20, 2035	U-3118	NP	May 12, 2024
		10213510	Apr 20, 2035	DP U-3118		
		10398686	Mar 13, 2034	DP		
		10632199	Apr 20, 2035	DP U-3118		
		10898575	Apr 20, 2035	DP U-3118		
		10980886	Apr 20, 2035	DP		
		11083730	Apr 20, 2035	DP U-3118		
		11083797	Apr 20, 2035	DP U-3118		
		9592227	Mar 13, 2034	DP U-3118		
		9694079	Apr 20, 2035	DP U-3118		
		9744163	Mar 13, 2034	DP		
		9801945	Apr 20, 2035	DP U-3118		
		9913909	Mar 13, 2034	U-3118		
<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988	003	10098957	Apr 20, 2035	U-3118	NP	May 12, 2024
		10213510	Apr 20, 2035	DP U-3118		
		10398686	Mar 13, 2034	DP		
		10632199	Apr 20, 2035	DP U-3118		
		10898575	Apr 20, 2035	DP U-3118		
		10980886	Apr 20, 2035	DP		
		11083730	Apr 20, 2035	DP U-3118		
		11083797	Apr 20, 2035	DP U-3118		
		9592227	Mar 13, 2034	DP U-3118		
		9694079	Apr 20, 2035	DP U-3118		
		9744163	Mar 13, 2034	DP		
		9801945	Apr 20, 2035	DP U-3118		
		9913909	Mar 13, 2034	U-3118		
<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988	004	10098957	Apr 20, 2035	U-3118	NP	May 12, 2024
		10213510	Apr 20, 2035	DP U-3118		
		10398686	Mar 13, 2034	DP		
		10632199	Apr 20, 2035	DP U-3118		
		10898575	Apr 20, 2035	DP U-3118		
		10980886	Apr 20, 2035	DP		
		11083730	Apr 20, 2035	DP U-3118		
		11083797	Apr 20, 2035	DP U-3118		

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<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988	004	9592227	Mar 13, 2034	DP	U-3118	
		9694079	Apr 20, 2035	DP	U-3118	
		9744163	Mar 13, 2034	DP		
		9801945	Apr 20, 2035	DP	U-3118	
		9913909	Mar 13, 2034		U-3118	
<u>BUPRENORPHINE - SUBLOCADE</u>						
N 209819	001	10198218	Jun 06, 2031		U-2489	
		10558394	Jun 25, 2031	DP		
		10592168	Jun 06, 2031		U-2489	
		11000520	Nov 06, 2035		U-3111	
		8921387	Jan 06, 2032	DP	U-2173	
		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 - SUBLOCADE</u>						
N 209819	002	10198218	Jun 06, 2031		U-2489	
		10558394	Jun 25, 2031	DP		
		10592168	Jun 06, 2031		U-2489	
		10646484	Jun 22, 2038		U-2489	
		11000520	Nov 06, 2035		U-3111	
		8921387	Jan 06, 2032	DP	U-2173	
		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	
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932	001	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	8147866	Jul 23, 2027	DP	U-1769	
		9655843	Jul 23, 2027	DP	U-1556	
		9901539	Dec 21, 2032		U-1556	

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<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932	003	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	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	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	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	8147866	Jul 23, 2027	DP	U-1769	
		9655843	Jul 23, 2027	DP	U-1556	
		9901539	Dec 21, 2032		U-1556	
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410	001	10285910	Oct 11, 2022	DP		
		11135216	Aug 07, 2029	DP	U-3111	
		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	10285910	Oct 11, 2022	DP		
		11135216	Aug 07, 2029	DP	U-3111	
		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	10285910	Oct 11, 2022	DP		
		11135216	Aug 07, 2029	DP	U-3111	
		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	10285910	Oct 11, 2022	DP		
		11135216	Aug 07, 2029	DP	U-3111	
		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	10946010	Sep 18, 2032	DP		
		11020388	Sep 18, 2032	DP	U-3131	
		8470361	May 22, 2030	DP	U-1425	
		8658198	Dec 03, 2027	DP	U-1494	
		8940330	Sep 18, 2032	DP		

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<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242	001	9259421	Sep 18, 2032	DP		
		9439900	Sep 18, 2032	DP		
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242	002	10874661	Sep 18, 2032	DP		
		10946010	Sep 18, 2032	DP		
		11020387	Sep 18, 2032	DP U-3131		
		11020388	Sep 18, 2032	DP U-3131		
		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	10946010	Sep 18, 2032	DP		
		11020387	Sep 18, 2032	DP U-3131		
		11020388	Sep 18, 2032	DP U-3131		
		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	10946010	Sep 18, 2032	DP		
		11020387	Sep 18, 2032	DP U-3131		
		11020388	Sep 18, 2032	DP U-3131		
		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	10946010	Sep 18, 2032	DP		
		11020387	Sep 18, 2032	DP U-3131		
		11020388	Sep 18, 2032	DP U-3131		
		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	006	10946010	Sep 18, 2032	DP		
		11020388	Sep 18, 2032	DP U-3131		
		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 - BUNAVAIL</u>						
N 205637	001	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	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	8147866	Jul 23, 2027	DP U-1521		
		8703177	Aug 20, 2032	DP		
		9522188	Apr 24, 2035	DP		
		9655843	Jul 23, 2027	DP U-2017		

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<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - BUNAVAIL</u>						
N 205637	003	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	10231964	Jul 02, 2034		U-1583	
		10307376	Nov 08, 2027		U-1585	
		10403170	Jun 05, 2033		U-1583	
		10828294	Jul 02, 2034		U-1583	
		10835527	Jul 02, 2034		U-1583	
		11033543	Jan 10, 2031		U-1583	
		11139056	Jun 05, 2033		U-1583	
		7375111	Mar 26, 2025	DP		
		7462626	Jul 20, 2024		U-1583	
		8088786	Feb 03, 2029	DP		
		8318788	Nov 08, 2027		U-1584	
		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	
		9633575	Jun 25, 2033		U-1583	
<u>CABAZITAXEL - JEVTANA KIT</u>						
N 201023	001	10583110	Oct 27, 2030		U-2753	
		10716777	Oct 27, 2030		U-2856	
		7241907	Dec 10, 2025	DS		
		7241907*PED	Jun 10, 2026			
		8927592	Oct 27, 2030		U-3200	
		8927592*PED	Apr 27, 2031			
<u>CABOTEGRAVIR SODIUM - VOCABRIA</u>						
N 212887	001	10927129	Apr 28, 2026	DS DP		
		8410103	Apr 28, 2026	DS DP	U-3061	
					NCE	Jan 21, 2026

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<u>CABOTEGRAVIR; RILPIVIRINE - CABENUVA KIT</u>						
N 212888	001	10927129	Apr 28, 2026	DS DP	NCE	Jan 21, 2026
		7125879	Apr 21, 2025	DS DP U-3059		
		8080551	Apr 11, 2023	DS DP		
		8410103	Apr 28, 2026	DS DP U-3060		
<u>CABOTEGRAVIR; RILPIVIRINE - CABENUVA KIT</u>						
N 212888	002	10927129	Apr 28, 2026	DS DP	NCE	Jan 21, 2026
		7125879	Apr 21, 2025	DS DP U-3059		
		8080551	Apr 11, 2023	DS DP		
		8410103	Apr 28, 2026	DS DP U-3060		
<u>CABOZANTINIB S-MALATE - COMETRIO</u>						
N 203756	001	11091439	Jan 15, 2030	DS		
		11091440	Jan 15, 2030	DP		
		11098015	Jan 15, 2030		U-1617	
		7579473	Aug 14, 2026	DS DP		
		8877776	Oct 08, 2030	DS DP U-1617		
		9717720	Feb 10, 2032	DP		
<u>CABOZANTINIB S-MALATE - COMETRIO</u>						
N 203756	002	11091439	Jan 15, 2030	DS		
		11091440	Jan 15, 2030	DP		
		11098015	Jan 15, 2030		U-1617	
		7579473	Aug 14, 2026	DS DP		
		8877776	Oct 08, 2030	DS DP U-1617		
		9717720	Feb 10, 2032	DP		
<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692	001	10034873	Jul 18, 2031		I-792	Jan 14, 2022
		10039757	Jul 18, 2031		I-854	Jan 22, 2024
		11091439	Jan 15, 2030	DS	I-873	Sep 17, 2024
		11091440	Jan 15, 2030	DP	ODE-227	Jan 14, 2026
		11098015	Jan 15, 2030		ODE-375	Sep 17, 2028
		11098015	Jan 15, 2030			
		11098015	Jan 15, 2030			
		11098015	Jan 15, 2030			
		11141413	Apr 17, 2037			
		7579473	Aug 14, 2026	DS DP		
		8497284	Sep 24, 2024		U-1220	
		8497284	Sep 24, 2024		U-1480	
		8497284	Sep 24, 2024		U-2488	
		8877776	Oct 08, 2030	DS DP U-3225		
		9724342	Jul 09, 2033	DP		
<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692	002	10034873	Jul 18, 2031		I-792	Jan 14, 2022
		10039757	Jul 18, 2031		I-854	Jan 22, 2024
		11091439	Jan 15, 2030	DS	I-873	Sep 17, 2024
		11091440	Jan 15, 2030	DP	ODE-227	Jan 14, 2026
		11098015	Jan 15, 2030		ODE-375	Sep 17, 2028
		11098015	Jan 15, 2030			
		11098015	Jan 15, 2030			
		11098015	Jan 15, 2030			
		11141413	Apr 17, 2037			
		7579473	Aug 14, 2026	DS DP		
		8497284	Sep 24, 2024		U-1220	
		8497284	Sep 24, 2024		U-1480	
		8497284	Sep 24, 2024		U-2488	
		8877776	Oct 08, 2030	DS DP U-3225		
		9724342	Jul 09, 2033	DP		
<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692	003	10034873	Jul 18, 2031		I-792	Jan 14, 2022
		10039757	Jul 18, 2031		I-854	Jan 22, 2024
		11091439	Jan 15, 2030	DS	I-873	Sep 17, 2024
		11091440	Jan 15, 2030	DP	ODE-227	Jan 14, 2026
		11098015	Jan 15, 2030		ODE-375	Sep 17, 2028
		11098015	Jan 15, 2030			
		11098015	Jan 15, 2030			
		11098015	Jan 15, 2030			
		11141413	Apr 17, 2037			
		7579473	Aug 14, 2026	DS DP		
		8497284	Sep 24, 2024		U-1220	
		8497284	Sep 24, 2024		U-1480	
		8497284	Sep 24, 2024		U-2488	
		8877776	Oct 08, 2030	DS DP U-3225		
		9724342	Jul 09, 2033	DP		

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<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692	003	11141413	Apr 17, 2037		U-3239	
		7579473	Aug 14, 2026	DS DP		
		8497284	Sep 24, 2024		U-1220	
		8497284	Sep 24, 2024		U-1480	
		8497284	Sep 24, 2024		U-2488	
		8877776	Oct 08, 2030	DS DP	U-3225	
		9724342	Jul 09, 2033	DP		
<u>CALCIFEDIOL - RAYALDEE</u>						
N 208010	001	10213442	Feb 02, 2027	DP		
		10300078	Mar 14, 2034	DP		
		10357502	Mar 14, 2034	DP		
		11154509	Apr 25, 2028		U-3248	
		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	May 07, 2028	DP	U-1280	NPP May 06, 2022
		8263580	May 07, 2028	DP	U-2662	NPP Nov 05, 2022
		8629128	May 26, 2026	DP	U-1280	
		8629128	May 26, 2026	DP	U-1767	
		8629128	May 26, 2026	DP	U-2662	
<u>CALCITRIOL - VECTICAL</u>						
N 022087	001					NPP Jul 17, 2023
<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 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	Jul 25, 2037	DP		
		10342813	Jul 25, 2037	DP		
<u>CALCIUM GLUCONATE - CALCIUM GLUCONATE IN SODIUM CHLORIDE</u>						
N 210906	002	10130646	Jul 25, 2037	DP		
		10342813	Jul 25, 2037	DP		
<u>CALCIUM GLUCONATE - CALCIUM GLUCONATE IN SODIUM CHLORIDE</u>						
N 210906	003	10130646	Jul 25, 2037	DP		
		10342813	Jul 25, 2037	DP		
<u>CALCIUM OXYBATE; MAGNESIUM OXYBATE; POTASSIUM OXYBATE; SODIUM OXYBATE - XYWAY</u>						
N 212690	001	10195168	Jan 11, 2033	DP	I-870	Aug 12, 2024
		10213400	Mar 15, 2033		NP	Jul 21, 2023
		10675258	Jan 11, 2033		U-2938	Jul 21, 2027
		10864181	Mar 15, 2033		U-3017	Aug 12, 2028
		8591922	Jan 11, 2033	DP		
		8731963	Dec 17, 2022		U-1110	
		8772306	Mar 15, 2033		U-1532	

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<u>CALCIUM OXYBATE; MAGNESIUM OXYBATE; POTASSIUM OXYBATE; SODIUM OXYBATE - XYWAV</u>						
N 212690	001	8772306	Mar 15, 2033	U-3198		
		8901173	Jan 11, 2033	DP		
		9050302	Mar 15, 2033	U-1532		
		9132107	Jan 11, 2033	DP		
		9486426	Mar 15, 2033	U-1532		
<u>CANAGLIFLOZIN - INVOKANA</u>						
N 204042	001	10617668	May 11, 2031	DP U-2441	I-809	Sep 27, 2022
		10617668	May 11, 2031	DP U-2632		
		10617668	May 11, 2031	DP U-2794		
		10617668	May 11, 2031	DP U-2795		
		10617668	May 11, 2031	DP U-2796		
		10617668	May 11, 2031	DP U-2797		
		10617668	May 11, 2031	DP U-2798		
		10617668	May 11, 2031	DP U-2799		
		10617668	May 11, 2031	DP U-493		
		7943582	Feb 26, 2029	DS DP U-2441		
		7943582	Feb 26, 2029	DS DP U-2632		
		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-2632		
		8222219	Apr 11, 2025	U-493		
		8513202	Dec 03, 2027	DS DP U-2441		
		8513202	Dec 03, 2027	DS DP U-2632		
		8513202	Dec 03, 2027	DS DP U-493		
<u>CANAGLIFLOZIN - INVOKANA</u>						
N 204042	002	10617668	May 11, 2031	DP U-2441	I-809	Sep 27, 2022
		10617668	May 11, 2031	DP U-2632		
		10617668	May 11, 2031	DP U-2794		
		10617668	May 11, 2031	DP U-2795		
		10617668	May 11, 2031	DP U-2796		
		10617668	May 11, 2031	DP U-2797		
		10617668	May 11, 2031	DP U-2798		
		10617668	May 11, 2031	DP U-2799		
		10617668	May 11, 2031	DP U-493		
		7943582	Feb 26, 2029	DS DP U-2441		
		7943582	Feb 26, 2029	DS DP U-2632		
		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-2632		
		8222219	Apr 11, 2025	U-493		
		8513202	Dec 03, 2027	DS DP U-2441		
		8513202	Dec 03, 2027	DS DP U-2632		
		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		
		7943582	Feb 26, 2029	DS DP U-2632		
		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-2632		
		8222219	Apr 11, 2025	U-493		
		8513202	Dec 03, 2027	DS DP U-2441		
		8513202	Dec 03, 2027	DS DP U-2632		
		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		
		7943582	Feb 26, 2029	DS DP U-2632		
		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-2632		
		8222219	Apr 11, 2025	U-493		
		8513202	Dec 03, 2027	DS DP U-2441		

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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	002	8513202	Dec 03, 2027	DS DP	U-2632	
		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	
		7943582	Feb 26, 2029	DS DP	U-2632	
		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-2632	
		8222219	Apr 11, 2025		U-493	
		8513202	Dec 03, 2027	DS DP	U-2441	
		8513202	Dec 03, 2027	DS DP	U-2632	
		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	
		7943582	Feb 26, 2029	DS DP	U-2632	
		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-2632	
		8222219	Apr 11, 2025		U-493	
		8513202	Dec 03, 2027	DS DP	U-2441	
		8513202	Dec 03, 2027	DS DP	U-2632	
		8513202	Dec 03, 2027	DS DP	U-493	
		8785403	Jul 30, 2024	DP		
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879	001	7943582	Feb 26, 2029	DS DP	U-2441	
		7943582	Feb 26, 2029	DS DP	U-2632	
		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-2632	
		8222219	Apr 11, 2025		U-493	
		8513202	Dec 03, 2027	DS DP	U-2441	
		8513202	Dec 03, 2027	DS DP	U-2632	
		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	
		7943582	Feb 26, 2029	DS DP	U-2632	
		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-2632	
		8222219	Apr 11, 2025		U-493	
		8513202	Dec 03, 2027	DS DP	U-2441	
		8513202	Dec 03, 2027	DS DP	U-2632	
		8513202	Dec 03, 2027	DS DP	U-493	
		8785403	Jul 30, 2024	DP		
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879	003	7943582	Feb 26, 2029	DS DP	U-2441	
		7943582	Feb 26, 2029	DS DP	U-2632	
		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-2632	
		8222219	Apr 11, 2025		U-493	
		8513202	Dec 03, 2027	DS DP	U-2441	
		8513202	Dec 03, 2027	DS DP	U-2632	
		8513202	Dec 03, 2027	DS DP	U-493	
		8785403	Jul 30, 2024	DP		

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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	
		7943582	Feb 26, 2029	DS DP	U-2632	
		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-2632	
		8222219	Apr 11, 2025		U-493	
		8513202	Dec 03, 2027	DS DP	U-2441	
		8513202	Dec 03, 2027	DS DP	U-2632	
		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	
		6130208	Jun 29, 2023	DP	U-1715	
		8680052	Mar 09, 2033		U-2979	
		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	NPP Jul 31, 2023
		10111840	Jun 17, 2035		U-2443	ODE-216 Sep 28, 2025
		10137095	Jun 17, 2035		U-2454	ODE-326 Jul 31, 2027
		10137095	Jun 17, 2035		U-2455	ODE-332 Jul 31, 2027
		10195159	May 07, 2022	DS		
		10603288	Jun 17, 2035		U-2780	
		10603288	Jun 17, 2035		U-2781	
		10603288	Jun 17, 2035		U-2782	
		10603288	Jun 17, 2035		U-2783	
		10709671	Jun 17, 2035		U-2862	
		10709673	Jun 17, 2035	DP		
		10709674	Jun 17, 2035		U-2780	
		10709674	Jun 17, 2035		U-2781	
		10849860	Jun 17, 2035		U-2427	
		10849860	Jun 17, 2035		U-2454	
		10918608	Oct 14, 2035		U-3071	
		10918608	Oct 14, 2035		U-3072	
		10918608	Oct 14, 2035		U-3073	
		10966939	Jun 17, 2035	DP	U-2780	
		10966939	Jun 17, 2035	DP	U-2781	
		11065209	Oct 14, 2035		U-3071	
		11096905	Oct 14, 2035	DS DP	U-2780	
		11096905	Oct 14, 2035	DS DP	U-2781	
		11154516	Jun 17, 2034		U-3235	
		11154516	Jun 17, 2034		U-3236	
		11160795	Mar 01, 2041		U-3233	
		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>CAPMATINIB HYDROCHLORIDE - TABRECTA</u>						
N 213591	001	10596178	Jul 22, 2035	DS DP		NCE May 06, 2025
		7767675	Nov 19, 2027	DS DP		ODE-291 May 06, 2027
		8420645	Jun 05, 2031	DS DP		
		8461330	Nov 19, 2027	DS DP		
		8901123	May 20, 2029		U-2813	
<u>CAPMATINIB HYDROCHLORIDE - TABRECTA</u>						
N 213591	002	10596178	Jul 22, 2035	DS DP		NCE May 06, 2025
		7767675	Nov 19, 2027	DS DP		ODE-291 May 06, 2027
		8420645	Jun 05, 2031	DS DP		
		8461330	Nov 19, 2027	DS DP		
		8901123	May 20, 2029		U-2813	

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<u>CAPMATINIB HYDROCHLORIDE - TABRECTA</u>						
N 213591	002	10596178	Jul 22, 2035	DS DP	NCE	May 06, 2025
		7767675	Nov 19, 2027	DS DP	ODE-291	May 06, 2027
		8420645	Jun 05, 2031	DS DP		
		8461330	Nov 19, 2027	DS DP		
		8901123	May 20, 2029		U-2813	
<u>CAPSAICIN - QUTENZA</u>						
N 022395	001	10034841	Sep 06, 2025	DP	I-838	Jul 17, 2023
		10463598	Sep 05, 2023	DP		
		10869827	Sep 05, 2023	DP		
		8263059	Sep 05, 2023		U-2705	
		8821920	Mar 26, 2030	DP		
		8889113	Sep 05, 2023		U-2705	
		9226903	Dec 15, 2028	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; 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		

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<u>CARBIDOPA; LEVODOPA - RYTARY</u>						
N 203312	003	7094427	May 29, 2022	DP U-1645		
		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		
		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 - 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	I-842	Aug 20, 2023
		7417042	Jul 20, 2026	DS DP		
		7491704	Apr 14, 2025		U-1260	
		7491704	Apr 14, 2025		U-2319	
		7491704	Apr 14, 2025		U-2320	
		7491704	Apr 14, 2025		U-2947	
		7737112	Dec 07, 2027	DP		
		8129346	Apr 14, 2025		U-1260	
		8129346	Apr 14, 2025		U-2319	
		8129346	Apr 14, 2025		U-2320	
		8129346	Apr 14, 2025		U-2947	
		8207125	Apr 14, 2025	DS DP		
		8207126	Apr 14, 2025	DP		
		8207127	Apr 14, 2025		U-1260	
		8207127	Apr 14, 2025		U-2319	
		8207127	Apr 14, 2025		U-2320	
		8207127	Apr 14, 2025		U-2947	
		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	I-842	Aug 20, 2023
		7417042	Jul 20, 2026	DS DP		
		7491704	Apr 14, 2025		U-1260	
		7491704	Apr 14, 2025		U-2319	
		7491704	Apr 14, 2025		U-2320	

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<u>CARFILZOMIB - KYPROLIS</u>						
N 202714 002	7491704	Apr 14, 2025	U-2947			
	7737112	Dec 07, 2027	DP			
	8129346	Apr 14, 2025	U-1260			
	8129346	Apr 14, 2025	U-2319			
	8129346	Apr 14, 2025	U-2320			
	8129346	Apr 14, 2025	U-2947			
	8207125	Apr 14, 2025	DS DP			
	8207126	Apr 14, 2025	DP			
	8207127	Apr 14, 2025	U-1260			
	8207127	Apr 14, 2025	U-2319			
	8207127	Apr 14, 2025	U-2320			
	8207127	Apr 14, 2025	U-2947			
	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		I-842	Aug 20, 2023
	7417042	Jul 20, 2026	DS DP			
	7491704	Apr 14, 2025	U-2319			
	7491704	Apr 14, 2025	U-2320			
	7491704	Apr 14, 2025	U-2947			
	7737112	Dec 07, 2027	DP			
	8129346	Apr 14, 2025	U-2319			
	8129346	Apr 14, 2025	U-2320			
	8129346	Apr 14, 2025	U-2947			
	8207125	Apr 14, 2025	DS DP			
	8207126	Apr 14, 2025	DP			
	8207127	Apr 14, 2025	U-2319			
	8207127	Apr 14, 2025	U-2320			
	8207127	Apr 14, 2025	U-2947			
	8207297	Apr 14, 2025	DS DP			
	9493582	Feb 27, 2033	DP			
	9511109	Oct 21, 2029	U-1924			
<u>CARGLUMIC ACID - CARGLUMIC ACID</u>						
A 213729 001					CGT	Jun 18, 2022
<u>CARGLUMIC ACID - CARBAGLU</u>						
N 022562 001					ODE-345	Jan 22, 2028
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 001	7737142	Sep 17, 2029	DS DP U-1750		I-798	May 24, 2022
	7737142	Sep 17, 2029	DS DP U-2543			
	7737142	Sep 17, 2029	DS DP U-2544			
	7737142	Sep 17, 2029	DS DP U-2545			
	7943621	Dec 16, 2028	DS DP			
	RE47350	Jul 16, 2029	U-1750			
	RE47350	Jul 16, 2029	U-2543			
	RE47350	Jul 16, 2029	U-2544			
	RE47350	Jul 16, 2029	U-2545			
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 002	7737142	Sep 17, 2029	DS DP U-1750		I-798	May 24, 2022
	7737142	Sep 17, 2029	DS DP U-2543			
	7737142	Sep 17, 2029	DS DP U-2544			
	7737142	Sep 17, 2029	DS DP U-2545			
	7943621	Dec 16, 2028	DS DP			
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 003	7737142	Sep 17, 2029	DS DP U-1750		I-798	May 24, 2022
	7737142	Sep 17, 2029	DS DP U-2543			
	7737142	Sep 17, 2029	DS DP U-2544			
	7943621	Dec 16, 2028	DS DP			
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 004	7737142	Sep 17, 2029	DS DP U-1750		I-798	May 24, 2022
	7737142	Sep 17, 2029	DS DP U-2543			
	7737142	Sep 17, 2029	DS DP U-2544			
	7943621	Dec 16, 2028	DS DP			

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<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370	004	7737142	Sep 17, 2029	DS DP U-1750	I-798	May 24, 2022
		7737142	Sep 17, 2029	DS DP U-2543		
		7737142	Sep 17, 2029	DS DP U-2544		
		7943621	Dec 16, 2028	DS DP		
<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>CASIMERSEN - AMONDYS 45</u>						
N 213026	001	10287586	Nov 12, 2030	DS DP	NCE	Feb 25, 2026
		10781450	Nov 12, 2030	U-3089	ODE-347	Feb 25, 2028
		8524880	Apr 02, 2026	DS DP U-3087		
		8524880	Apr 02, 2026	DS DP U-3088		
		9228187	Nov 12, 2030	DS DP		
		9447415	Jun 28, 2025	DS DP		
		9758783	Nov 12, 2030	U-3088		
		9758783	Nov 12, 2030	U-3089		
<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>CEDAZURIDINE; DECITABINE - INOOVI</u>						
N 212576	001	8268800	Aug 22, 2030	DS	U-2864	NCE
		8268800	Aug 22, 2030	DS	U-2865	ODE-316
		8268800	Aug 22, 2030	DS	U-2866	Jul 07, 2027
		8268800	Aug 22, 2030	DS	U-2867	
		8618075	Oct 16, 2028		U-2864	
		8618075	Oct 16, 2028		U-2867	
		9567363	Oct 16, 2028	DS		
<u>CEFIDEROCOL SULFATE TOSYLATE - FETROJA</u>						
N 209445	001	10004750	Sep 03, 2035	DS DP	I-844	Sep 25, 2023
		9238657	Nov 19, 2031	DS DP U-282	NCE	Nov 14, 2024
		9949982	Sep 03, 2035	DP	GAIN	Nov 14, 2029
<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	Sep 13, 2022
		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	Sep 13, 2022
		8247400	Feb 10, 2031	DP U-282		
		9629861	Sep 21, 2030	DP		

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<u>CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM - ZERBAXA</u>						
N 206829 001	10028963	Sep 07, 2032	U-2565		NCE	Dec 19, 2019
	10028963	Sep 07, 2032	U-2566		GAIN	Dec 19, 2024
	10125149	Aug 14, 2035	DP			
	10376496	Sep 09, 2034	U-2610			
	10376496	Sep 09, 2034	U-2611			
	10420841	Mar 14, 2034	U-1672			
	10420841	Mar 14, 2034	U-2631			
	10933053	Sep 09, 2034	U-3090			
	10933053	Sep 09, 2034	U-3091			
	7129232	May 15, 2028	DS DP U-1676			
	7129232	May 15, 2028	DS DP U-36			
	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			
	9724353	Sep 07, 2032	U-2565			
	9724353	Sep 07, 2032	U-2566			
	9872906	Mar 14, 2034	DP			
<u>CELECOXIB - ELYXYB</u>						
N 212157 001	10376527	May 27, 2036	DP U-2718		NP	May 05, 2023
	10722456	May 27, 2036	DP U-2718			
	10799517	May 27, 2036	DP U-2718			
	9572819	May 27, 2036	DP U-2718			
	9795620	May 27, 2036	DP U-2718			
	9949990	May 27, 2036	DP U-2718			
<u>CELECOXIB; TRAMADOL HYDROCHLORIDE - SEGLENTIS</u>						
N 213426 001	10238668	Apr 19, 2030	DS DP U-3244		NP	Oct 15, 2024
	10245276	Apr 19, 2030	DS DP			
	10548909	Apr 19, 2030	U-3244			
	8598152	Apr 19, 2030	DS DP			
	8846744	Jun 03, 2031	DP			
	9012440	Apr 19, 2030	DS DP			
<u>CENOBAMATE - XCOPRI</u>						
N 212839 001	7598279	Oct 30, 2027	DS		NCE	Mar 10, 2025
<u>CENOBAMATE - XCOPRI</u>						
N 212839 002	7598279	Oct 30, 2027	DS		NCE	Mar 10, 2025
<u>CENOBAMATE - XCOPRI</u>						
N 212839 003	7598279	Oct 30, 2027	DS		NCE	Mar 10, 2025
<u>CENOBAMATE - XCOPRI</u>						
N 212839 004	7598279	Oct 30, 2027	DS		NCE	Mar 10, 2025
<u>CENOBAMATE - XCOPRI</u>						
N 212839 005	7598279	Oct 30, 2027	DS		NCE	Mar 10, 2025
<u>CENOBAMATE - XCOPRI</u>						
N 212839 006	7598279	Oct 30, 2027	DS		NCE	Mar 10, 2025
<u>CERITINIB - ZYKADIA</u>						
N 205755 001	7893074	Apr 25, 2026	DS DP		ODE-145	May 26, 2024
	7964592	Apr 29, 2028	DS DP			
	8039479	Jun 29, 2030	DS DP			
	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			

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<u>CERITINIB - ZYKADIA</u>						
N 211225	001 7893074	Apr 25, 2026	DS DP		ODE*	May 26, 2024
	7964592	Apr 29, 2028	DS DP			
	8039479	Jun 29, 2030	DS DP			
	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		
	8829005*PED	Sep 15, 2030				
	9254286	Jul 09, 2032	DP			
	9254286*PED	Jan 09, 2033				
	9750684	Mar 15, 2030	DP			
	9993471	Mar 15, 2030		U-1680		
<u>CETIRIZINE HYDROCHLORIDE - OUZYTTIR</u>						
N 211415	001 8263581	Feb 28, 2030		U-2635	NP	Oct 04, 2022
	8314083	Feb 28, 2030		U-2634		
	8513259	Feb 11, 2030		U-2636		
	9119771	Feb 11, 2030		U-2635		
	9180090	Feb 11, 2030		U-2635		
<u>CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ZYRTEC-D 12 HOUR</u>						
N 021150	002 7014867	Jun 10, 2022	DP			
	7226614	Jun 10, 2022		U-295		
<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; 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 WITH TINT</u>						
N 020832	005 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 6729786	Mar 14, 2023	DP			
	7241065	Mar 14, 2023	DP			
	7422388	Apr 25, 2027	DP	U-1397		
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - SOLUPREP</u>						
N 208288	001 8623935	Jul 26, 2029	DP	U-1022		
<u>CHLOROPROCAINE HYDROCHLORIDE - CLOROTEKAL</u>						
N 208791	001 8969412	Sep 05, 2026	DP	U-2609		
	9504666	Dec 11, 2033	DP			
<u>CHLORPHENIRAMINE MALEATE; CODEINE PHOSPHATE - TUXARIN ER</u>						
N 206323	001 9066942	Jan 03, 2032		U-1716		
	9107921	Jan 03, 2032	DP			

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<u>CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE - ADVIL ALLERGY SINUS</u>						
N 021441	001	7863287	Feb 28, 2027	DP		
<u>CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE - CHILDREN'S ADVIL ALLERGY SINUS</u>						
N 021587	001	10238640	May 25, 2024	DP		
<u>CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX - TUZISTRA XR</u>						
N 207768	001	8062667	Mar 29, 2029	DP		
		8790700	Mar 15, 2027	DP		
<u>CHLORPROMAZINE HYDROCHLORIDE - CHLORPROMAZINE HYDROCHLORIDE</u>						
A 214542	001				CGT	Jan 29, 2022
<u>CHLORPROMAZINE HYDROCHLORIDE - CHLORPROMAZINE HYDROCHLORIDE</u>						
A 214542	002				CGT	Jan 29, 2022
<u>CHLORZOXAZONE - CHLORZOXAZONE</u>						
A 215158	001				CGT	Mar 29, 2022
<u>CHOLIC ACID - CHOLBAM</u>						
N 205750	001				ODE-91	Mar 17, 2022
<u>CHOLIC ACID - CHOLBAM</u>						
N 205750	002				ODE-91	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>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	8371292	Feb 01, 2028	U-1356		
<u>CICLESONIDE - ZETONNA</u>						
N 202129	001	8371292	Feb 01, 2028	U-1357		
<u>CILASTATIN SODIUM; IMPENEM; RELEBACTAM - RECARBRIO</u>						
N 212819	001	8487093	Nov 19, 2029	DS DP U-2586	NCE	Jul 16, 2024
		8487093	Nov 19, 2029	DS DP U-2587	GAIN	Jul 16, 2029
		8487093	Nov 19, 2029	DS DP U-2840		
<u>CINACALCET HYDROCHLORIDE - SENSIPAR</u>						
N 021688	001	7829595	Sep 22, 2026	DP U-1098		
		9375405	Sep 22, 2026	DP		
<u>CINACALCET HYDROCHLORIDE - SENSIPAR</u>						
N 021688	002	7829595	Sep 22, 2026	DP U-1098		
		9375405	Sep 22, 2026	DP		
<u>CINACALCET HYDROCHLORIDE - SENSIPAR</u>						
N 021688	003	7829595	Sep 22, 2026	DP U-1098		
		9375405	Sep 22, 2026	DP		
<u>CIPROFLOXACIN - OTIPRIO</u>						
N 207986	001	11040004	Nov 12, 2037	U-2252		
		8318817	Apr 27, 2030	U-1792		
		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; FLUOCINOLONE ACETONIDE - OTOVEL</u>						
N 208251	001	8932610	Mar 24, 2030	DP U-1578		

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<u>CIPROFLOXACIN; DEXAMETHASONE - CIPRODEX</u>						
N 021537	001	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; LACTIC ACID; POTASSIUM BITARTRATE - PHEXXI</u>						
N 208352	001	10568855	Mar 15, 2033	U-1	NP	May 22, 2023
		6706276	Mar 06, 2022	DP		
<u>CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE - PREPOPIK</u>						
N 202535	001	8450338	Oct 10, 2028	DP		
		8481083	Oct 10, 2028	DP		
<u>CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE - CLENPIO</u>						
N 209589	001	10624879	Jun 23, 2034	DP		
		11191753	Jun 23, 2034	U-3261		
		9827231	Jun 26, 2034	DP U-2162		
<u>CLADRIBINE - MAVENCLAD</u>						
N 022561	001	7713947	Oct 16, 2026	U-2520	NP	Mar 29, 2022
		7888328	Apr 11, 2024	DP U-2521		
		8377903	May 31, 2026	U-2522		
		8785415	Apr 11, 2024	DP U-2523		
<u>CLASCOTERONE - WINLEVI</u>						
N 213433	001	10159682	Aug 14, 2028	U-2942	NCE	Aug 26, 2025
		8143240	Jan 12, 2023	U-2942		
		8785427	Jul 25, 2030	DP		
		8865690	Jul 24, 2022	U-2942		
		9211295	Jul 24, 2022	DP		
		9433628	Feb 08, 2029	DP		
		9486458	Jul 24, 2028	U-2942		
<u>CLEVIDIPINE - CLEVIPREX</u>						
N 022156	001	10010537	Oct 10, 2031	DP		
		11103490	Oct 10, 2031	DP		
		8658676	Oct 10, 2031	DP		
<u>CLEVIDIPINE - CLEVIPREX</u>						
N 022156	002	10010537	Oct 10, 2031	DP		
		11103490	Oct 10, 2031	DP		
		8658676	Oct 10, 2031	DP		
<u>CLEVIDIPINE - CLEVIPREX</u>						
N 022156	003	10010537	Oct 10, 2031	DP		
		11103490	Oct 10, 2031	DP		
		8658676	Oct 10, 2031	DP		
<u>CLINDAMYCIN PHOSPHATE - CLINDESSE</u>						
N 050793	001	6899890	Apr 27, 2023	DP U-137		
		9789057	Dec 02, 2026	DP U-137		
<u>CLINDAMYCIN PHOSPHATE - EVOCLIN</u>						
N 050801	001	7141237	Feb 03, 2024	DP		
		7374747	Jan 23, 2024	DP U-921		
<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		

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<u>CLOBETASOL PROPIONATE - CLOBEX</u>						
N 021644	001 7700081	Jan 03, 2022	U-1044			
<u>CLOBETASOL PROPIONATE - OLUX E</u>						
N 022013	001 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			
	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			
	10588914	Aug 31, 2030	DP U-2771			
	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>CLONIDINE - NEXICLON XR</u>						
N 022500	001 8337890	Apr 17, 2027	DP			
	8623409	Sep 08, 2031	DP			
<u>CLONIDINE - NEXICLON XR</u>						
N 022500	002 8337890	Apr 17, 2027	DP			
	8623409	Sep 08, 2031	DP			
<u>CLOZAPINE - VERSACLOZ</u>						
N 203479	001 8057811	May 01, 2028	DP			
<u>COBICISTAT - TYBOST</u>						
N 203094	001 10039718	Oct 06, 2032	DP		NPP	Aug 22, 2022
	10039718*PED	Apr 06, 2033			ODE-260	Aug 22, 2026
	8148374	Sep 03, 2029	DS DP U-1279			
	8148374*PED	Mar 03, 2030				
<u>COBICISTAT; DARUNAVIR - PREZCOBIX</u>						
N 205395	001 10039718	Oct 06, 2032	DP			
	7700645	Dec 26, 2026	DS DP			
	7700645*PED	Jun 26, 2027				
	8148374	Sep 03, 2029	DS DP U-1279			
	8148374	Sep 03, 2029	DS DP U-2939			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>COBICISTAT; DARUNAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - SYMTUZA</u>						
N 210455	001 10039718	Oct 06, 2032	DP			
	10786518	Jul 19, 2038	U-2978			
	7390791	Apr 17, 2025	DS DP			
	7700645	Dec 26, 2026	DS DP			
	7803788	Feb 02, 2022	U-2352			
	7803788	Feb 02, 2022	U-2765			
	8148374	Sep 03, 2029	DS DP U-2353			
	8148374	Sep 03, 2029	DS DP U-2364			
	8148374	Sep 03, 2029	DS DP U-2365			
	8148374	Sep 03, 2029	DS DP U-2766			
	8148374	Sep 03, 2029	DS DP U-2767			
	8148374	Sep 03, 2029	DS DP U-2768			
	8518987	Feb 16, 2024	DS DP			
	8754065	Aug 15, 2032	DS DP U-2352			
	8754065	Aug 15, 2032	DS DP U-2765			
	9296769	Aug 15, 2032	DS DP U-2352			
	9296769	Aug 15, 2032	DS DP U-2765			
<u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - GENVOYA</u>						
N 207561	001 10039718	Oct 06, 2032	DP			
	10039718*PED	Apr 06, 2033				
	7176220	Aug 27, 2026	DS DP U-257			

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<u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - GENVOYA</u>						
N 207561	001	7176220*PED	Feb 27, 2027			
		7390791	Apr 17, 2025	DS DP		
		7390791*PED	Oct 17, 2025			
		7635704	Oct 26, 2026	DS DP U-257		
		7635704*PED	Apr 26, 2027			
		7803788	Feb 02, 2022		U-257	
		8148374	Sep 03, 2029	DS DP U-1279		
		8148374*PED	Mar 03, 2030			
		8633219	Apr 24, 2030	DP U-257		
		8633219*PED	Oct 24, 2030			
		8754065	Aug 15, 2032	DS DP U-257		
		8754065*PED	Feb 15, 2033			
		8981103	Oct 26, 2026	DS DP		
		8981103*PED	Apr 26, 2027			
		9296769	Aug 15, 2032	DS DP U-257		
		9296769*PED	Feb 15, 2033			
		9891239	Sep 03, 2029	DP U-257		
		9891239*PED	Mar 03, 2030			
<u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - STRIBILD</u>						
N 203100	001	10039718	Oct 06, 2032	DP		
		10039718*PED	Apr 06, 2033			
		7176220	Aug 27, 2026	DS DP U-257		
		7176220*PED	Feb 27, 2027			
		7635704	Oct 26, 2026	DS DP U-257		
		7635704*PED	Apr 26, 2027			
		8148374	Sep 03, 2029	DS DP U-1279		
		8592397	Jan 13, 2024	DP U-257		
		8633219	Apr 24, 2030	DP U-257		
		8633219*PED	Oct 24, 2030			
		8716264	Jan 13, 2024	DP U-257		
		8981103	Oct 26, 2026	DS DP		
		8981103*PED	Apr 26, 2027			
		9457036	Jan 13, 2024	DP U-257		
		9744181	Jan 13, 2024	DP U-257		
		9891239	Sep 03, 2029	DP U-257		
		9891239*PED	Mar 03, 2030			
<u>COBIMETINIB FUMARATE - COTELLIC</u>						
N 206192	001	10478400	Jun 29, 2036	DS DP U-1776	ODE-101	Nov 10, 2022
		10590102	Jun 30, 2036	DS DP U-1776		
		11087354	Jun 22, 2034		U-1776	
		7803839	Nov 10, 2029	DS DP		
		8362002	Oct 05, 2026		U-1776	
<u>COCAINE HYDROCHLORIDE - NUMBRINO</u>						
N 209575	001				NCE	Dec 14, 2022
					NP	Jan 10, 2023
<u>COCAINE HYDROCHLORIDE - GOPRELTO</u>						
N 209963	001	10016407	Feb 07, 2037		NCE	Dec 14, 2022
		10149843	Feb 07, 2037		U-2478	
		10149843	Feb 07, 2037		U-2479	
		10231961	Feb 07, 2037	DP		
		10413505	Feb 07, 2037		U-2479	
		10420760	Feb 07, 2037		U-2478	
		10857095	Feb 07, 2037		U-3014	
		10894012	Feb 07, 2037		U-3014	
		10933060	Feb 07, 2037		U-3014	
		10973811	Feb 07, 2037		U-2226	
		10987347	Feb 07, 2037		U-2225	
		11040032	Feb 07, 2037	DP		
		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	

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<u>COLCHICINE - COLCRYS</u>						
N 022352	001	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		
		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>COLCHICINE - GLOPERBA</u>						
N 210942	001	10226423	Dec 20, 2037	DP		
		10383820	Nov 22, 2036	DP	U-2814	
		10383821	Nov 22, 2036	DP		
		9907751	Nov 22, 2036	DP		
<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	M-232	Oct 20, 2024
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>						
N 022362	002	7229613	Apr 17, 2022	U-493	M-232	Oct 20, 2024
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>						
N 210895	001	7229613	Apr 17, 2022	U-2516		
<u>COPANLISIB DIHYDROCHLORIDE - ALIQOPA</u>						
N 209936	001	10383876	Mar 29, 2032	DS DP	NCE	Sep 14, 2022
		7511041	May 13, 2024	DS DP	ODE-155	Sep 14, 2024
		9636344	Mar 29, 2032	U-2124		
		RE46856	Oct 22, 2029	DS DP	U-2124	
<u>COPPER DOTATATE CU-64 - DETECTNET</u>						
N 213227	001	10159759	Aug 23, 2032	U-2951	NCE	Sep 03, 2025
		10383961	Aug 23, 2032	U-2951	ODE-317	Sep 03, 2027
		11160888	Aug 23, 2032	U-2951		
<u>CRISABOROLE - EUCRISA</u>						
N 207695	001	8039451	Jun 29, 2029	DS DP	NPP	Mar 23, 2023
		8039451*PED	Dec 29, 2029		PED	Sep 23, 2023
		8168614	Jan 20, 2030	U-1932		
		8168614*PED	Jul 20, 2030			
		8501712	Feb 16, 2027	U-1932		
		8501712*PED	Aug 16, 2027			
		9682092	Feb 16, 2027	U-1932		
		9682092*PED	Aug 16, 2027			
<u>CRIZOTINIB - XALKORI</u>						
N 202570	001	7230098	Aug 26, 2025	DS	I-852	Jan 14, 2024
		7825137	May 12, 2027	U-3057	ODE-111	Mar 11, 2023
		7825137	May 12, 2027	U-3058	ODE-328	Jan 14, 2028
		7858643	Oct 08, 2029	DS DP		
		8217057	Nov 06, 2029	DS DP		
		8785632	Mar 01, 2025	DS		

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<u>CRIZOTINIB - XALKORI</u>						
N 202570	002 7230098	Aug 26, 2025	DS		I-852	Jan 14, 2024
	7825137	May 12, 2027		U-3057	ODE-111	Mar 11, 2023
	7825137	May 12, 2027		U-3058	ODE-328	Jan 14, 2028
	7858643	Oct 08, 2029	DS DP			
	8217057	Nov 06, 2029	DS DP			
	8785632	Mar 01, 2025	DS			
<u>CROFELEMER - MYTESI</u>						
N 202292	001 7341744	Jun 02, 2022	DP U-1319			
	8962680	Oct 31, 2031	U-1319			
	9585868	Oct 31, 2031	U-1319			
<u>CUPRIC SULFATE; MANGANESE SULFATE; SELENIOUS ACID; ZINC SULFATE - TRALEMENT</u>						
N 209376	001				NCE	Apr 30, 2024
<u>CUPRIC SULFATE; MANGANESE SULFATE; SELENIOUS ACID; ZINC SULFATE - TRALEMENT</u>						
N 209376	002				NCE	Apr 30, 2024
<u>CUPRIC SULFATE; MANGANESE SULFATE; SELENIOUS ACID; ZINC SULFATE - MULTRY'S</u>						
N 209376	003				NCE	Apr 30, 2024
<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			
	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>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210735	001 9662342	Jun 26, 2035	DP			
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210735	002 9662342	Jun 26, 2035	DP			
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 212501	001 10993952	Feb 15, 2036	DP			
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 212501	002 10993952	Feb 15, 2036	DP			
<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			

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<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	10441630	Aug 23, 2033	DP		
		10918694	Feb 28, 2037	DP		
		8980839	Aug 23, 2033	DP	U-1483	
		9937225	Aug 23, 2033	DP	U-1483	
<u>CYCLOSPORINE - VERKAZIA</u>						
N 214965	001	7973081	Jan 27, 2026	DP	NP	Jun 23, 2023
		8298568	Nov 03, 2027	DP	ODE-358	Jun 23, 2028
		8524779	Jan 27, 2026	DP		
		9132071	Jun 02, 2029	DP		
		9220694	Jan 27, 2026	DP		
		9956289	Jan 27, 2026	DP		
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 203389	001	10143665	Aug 16, 2036		U-1399	ODE-162
		10328037	Aug 16, 2036		U-1399	ODE-97
		10548859	Aug 16, 2036		U-1399	PED
		10905662	Aug 16, 2036		U-1399	
		8026284	Sep 22, 2027		U-1399	
		8026284*PED	Mar 22, 2028			
		9173851	Jun 17, 2034	DP		
		9173851*PED	Dec 17, 2034			
		9192590	Jan 26, 2027		U-1399	
		9192590*PED	Jul 26, 2027			
		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	ODE-162
		10328037	Aug 16, 2036		U-1399	ODE-97
		10548859	Aug 16, 2036		U-1399	PED
		10905662	Aug 16, 2036		U-1399	
		8026284	Sep 22, 2027		U-1399	
		8026284*PED	Mar 22, 2028			
		9173851	Jun 17, 2034	DP		
		9173851*PED	Dec 17, 2034			
		9192590	Jan 26, 2027		U-1399	
		9192590*PED	Jul 26, 2027			
		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 213491	001	10143665	Aug 16, 2036		U-1399	ODE*
		10328037	Aug 16, 2036		U-1399	ODE*
		10548859	Aug 16, 2036		U-1399	PED
		10905662	Aug 16, 2036		U-1399	
		8026284	Sep 22, 2027		U-1399	
		8026284*PED	Mar 22, 2028			

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<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 213491	001	9173851	Jun 17, 2034	DP		
		9173851*PED	Dec 17, 2034			
		9192590	Jan 26, 2027	U-1399		
		9192590*PED	Jul 26, 2027			
		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	DP U-1399		
		9925157	Jan 26, 2027	DP U-1399		
		9925158	Jan 26, 2027	DP U-1399		
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 213491	002	10143665	Aug 16, 2036	U-1399	ODE*	Aug 14, 2022
		10328037	Aug 16, 2036	U-1399	ODE*	Dec 22, 2024
		10548859	Aug 16, 2036	U-1399	PED	Feb 14, 2023
		10905662	Aug 16, 2036	U-1399		
		8026284	Sep 22, 2027	U-1399		
		8026284*PED	Mar 22, 2028			
		9173851	Jun 17, 2034	DP		
		9173851*PED	Dec 17, 2034			
		9192590	Jan 26, 2027	U-1399		
		9192590*PED	Jul 26, 2027			
		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	DP U-1399		
		9925157	Jan 26, 2027	DP U-1399		
		9925158	Jan 26, 2027	DP U-1399		
<u>CYSTEAMINE HYDROCHLORIDE - CYSTADROPS</u>						
N 211302	001				NP	Aug 19, 2023
<u>CYSTEINE HYDROCHLORIDE - ELCYS</u>						
N 210660	001	10478453	Jan 15, 2039	DP U-2752		
		10583155	Jan 15, 2039	DP U-2752		
		10653719	Jan 15, 2039	DP		
		10905713	Jan 15, 2039	DP		
		10905714	Jan 15, 2039	DP		
		10912795	Jan 15, 2039	DP		
		10918662	Jan 15, 2039	DP		
		10933089	Jan 15, 2039	DP		
<u>CYSTEINE HYDROCHLORIDE - NOURESS</u>						
N 212535	001	10478453	Jan 15, 2039	DP U-2752		
		10493051	Mar 15, 2039	DP		
		10543186	Mar 15, 2039	U-2722		
		10583155	Jan 15, 2039	DP U-2752		
		10653719	Jan 15, 2039	DP		
		10702490	Mar 15, 2039	DP		
		10905713	Jan 15, 2039	DP		
		10905714	Jan 15, 2039	DP		
		10912795	Jan 15, 2039	DP		
		10918662	Jan 15, 2039	DP		
		10933089	Jan 15, 2039	DP		
		11045438	Mar 15, 2039	DP		
<u>CYTARABINE; DAUNORUBICIN - VYXEOS</u>						
N 209401	001	10028912	Sep 29, 2034	DP U-3149	NPP	Mar 30, 2024
		10028912	Sep 29, 2034	DP U-3150	ODE-287	Aug 03, 2024
		10166184	Oct 15, 2032	DP U-3149	ODE-350	Mar 30, 2028
		10835492	Oct 15, 2032	U-3150		
		7850990	Jan 23, 2027	DP U-3147		
		8022279	Sep 14, 2027	DP U-3147		
		8092828	Apr 01, 2029	U-3147		
		8431806	Apr 22, 2025	DP U-3147		
		8518437	Jun 07, 2026	DP		
		9271931	Jan 23, 2027	DP		

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<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 022512	001	6087380*PED			I-862	Jun 21, 2024
		7866474	DP	Y	PED	Dec 21, 2024
		7932273	DS DP			
		9034822		U-1759		
		9034822*PED				
		9925174	DP			
		9925174*PED				
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 022512	002	6087380*PED			I-862	Jun 21, 2024
		7866474	DP	Y	PED	Dec 21, 2024
		7932273	DS DP			
		9034822		U-1759		
		9034822*PED				
		9925174	DP			
		9925174*PED				
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 022512	003	6087380*PED			I-862	Jun 21, 2024
		7866474	DP	Y	PED	Dec 21, 2024
		7866474*PED				
		7932273	DS DP			
		7932273*PED				
		9034822		U-1759		
		9034822*PED				
		9925174	DP			
		9925174*PED				
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358	001	6087380*PED			NP	Jun 21, 2024
		7932273	DS DP		PED	Dec 21, 2024
		7932273*PED				
		9925174	DP			
		9925174*PED				
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358	002	6087380*PED			NP	Jun 21, 2024
		7932273	DS DP		PED	Dec 21, 2024
		7932273*PED				
		9925174	DP			
		9925174*PED				
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358	003	6087380*PED			NP	Jun 21, 2024
		7932273	DS DP		PED	Dec 21, 2024
		7932273*PED				
		9925174	DP			
		9925174*PED				
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358	004	6087380*PED			NP	Jun 21, 2024
		7932273	DS DP		PED	Dec 21, 2024
		7932273*PED				
		9925174	DP			
		9925174*PED				
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358	005	6087380*PED			NP	Jun 21, 2024
		7932273	DS DP		PED	Dec 21, 2024
		7932273*PED				
		9925174	DP			
		9925174*PED				
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358	006	6087380*PED			NP	Jun 21, 2024
		7932273	DS DP		PED	Dec 21, 2024
		7932273*PED				
		9925174	DP			
		9925174*PED				

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<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 202806 001	10869869	Aug 30, 2033		U-3185	M-246	Oct 06, 2022
	7994185	Jan 20, 2030	DS DP	U-1406	ODE-147	Jun 22, 2024
	7994185	Jan 20, 2030	DS DP	U-2031	ODE-182	Apr 30, 2025
	7994185	Jan 20, 2030	DS DP	U-2032	ODE-183	May 04, 2025
	7994185	Jan 20, 2030	DS DP	U-2296		
	8415345	Jan 20, 2030	DS DP	U-1406		
	8415345	Jan 20, 2030	DS DP	U-2031		
	8415345	Jan 20, 2030	DS DP	U-2032		
	8415345	Jan 20, 2030	DS DP	U-2296		
	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-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		
<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 202806 002	10869869	Aug 30, 2033		U-3185	M-246	Oct 06, 2022
	7994185	Jan 20, 2030	DS DP	U-1406	ODE-147	Jun 22, 2024
	7994185	Jan 20, 2030	DS DP	U-2031	ODE-182	Apr 30, 2025
	7994185	Jan 20, 2030	DS DP	U-2032	ODE-183	May 04, 2025
	7994185	Jan 20, 2030	DS DP	U-2296		
	8415345	Jan 20, 2030	DS DP	U-1406		
	8415345	Jan 20, 2030	DS DP	U-2031		
	8415345	Jan 20, 2030	DS DP	U-2032		
	8415345	Jan 20, 2030	DS DP	U-2296		
	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-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		
<u>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</u>						
N 206843 001	8329159	Jul 24, 2029	DS			
	8629171	Jun 13, 2031	DS DP	U-1724		
	8642025	Aug 11, 2027	DS DP	U-1724		
	8642025	Aug 11, 2027	DS DP	U-1725		
	8900566	Aug 08, 2027		U-1724		
	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	Jul 24, 2029	DS			
	8629171	Jun 13, 2031	DS DP	U-1724		
	8642025	Aug 11, 2027	DS DP	U-1724		
	8642025	Aug 11, 2027	DS DP	U-1725		
	8900566	Aug 08, 2027		U-1724		
	8900566	Aug 08, 2027		U-1725		
	9421192	Aug 08, 2027	DS	U-1724		
	9421192	Aug 08, 2027	DS	U-1725		

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<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	NCE May 23, 2019
		7115564	Nov 14, 2023	DP		NPP Jul 22, 2024
		7119061	Nov 14, 2023	DP		GAIN May 23, 2024
		8143212	Nov 14, 2023		U-1517	
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287	001					NPP May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287	002					NPP May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287	003					NPP May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287	004					NPP May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287	005					NPP May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287	006					NPP May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287	007					NPP May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287	008					NPP May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287	009					NPP May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287	010					NPP May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287	011					NPP May 16, 2022
<u>DANTROLENE SODIUM - RYANODEX</u>						
N 205579	001	7758890	Jul 01, 2025	DP		
		8110225	Dec 24, 2022	DP		
		8604072	Dec 24, 2022	DP		
		8685460	Feb 15, 2023		U-1546	
		9884044	Jun 13, 2022	DP	U-1546	
<u>DAPAGLIFLOZIN - FARXIGA</u>						
N 202293	001	6515117	Oct 04, 2025	DS DP	U-2139	I-834 May 05, 2023
		6515117	Oct 04, 2025	DS DP	U-493	I-841 Oct 18, 2022
		7456254	Jun 30, 2025		U-2139	I-857 Apr 30, 2024
		7851502	Aug 19, 2028		DP	M-238 Feb 22, 2022
		7919598	Dec 16, 2029	DS		
		8221786	Mar 21, 2028	DP		

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<u>DAPAGLIFLOZIN - FARXIGA</u>						
N 202293	001	8329648	Aug 18, 2026	U-2139		
		8329648	Aug 18, 2026	U-2212		
		8329648	Aug 18, 2026	U-2213		
		8361972	Mar 21, 2028	U-2139		
		8361972	Mar 21, 2028	U-493		
		8431685	Apr 13, 2025	DP U-2139		
		8461105	Apr 13, 2025	DP U-2139		
		8501698	Jun 20, 2027	DP U-493		
		8685934	May 26, 2030	U-1522		
		8716251	Mar 21, 2028	DP		
		8721615	Jan 18, 2030	DP	Y	
		8906851	Aug 18, 2026	U-2139		
		9238076	Apr 15, 2024	DP U-2139		
<u>DAPAGLIFLOZIN - FARXIGA</u>						
N 202293	002	10973836	Mar 09, 2040	U-3127	I-834	May 05, 2023
		6515117	Oct 04, 2025	DS DP U-2139	I-841	Oct 18, 2022
		6515117	Oct 04, 2025	DS DP U-493	I-857	Apr 30, 2024
		7456254	Jun 30, 2025	DP U-2139	M-238	Feb 22, 2022
		7851502	Aug 19, 2028	DP		
		7919598	Dec 16, 2029	DS		
		8221786	Mar 21, 2028	DP		
		8329648	Aug 18, 2026	U-2139		
		8329648	Aug 18, 2026	U-2212		
		8329648	Aug 18, 2026	U-2213		
		8361972	Mar 21, 2028	U-2139		
		8361972	Mar 21, 2028	U-493		
		8431685	Apr 13, 2025	DP U-2139		
		8461105	Apr 13, 2025	DP U-2139		
		8501698	Jun 20, 2027	DP U-493		
		8685934	May 26, 2030	U-1522		
		8716251	Mar 21, 2028	DP		
		8721615	Jan 18, 2030	DP	Y	
		8906851	Aug 18, 2026	U-2139		
		9238076	Apr 15, 2024	DP U-2139		
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649	001	6515117	Oct 04, 2025	DS DP U-493	I-841	Oct 18, 2022
		7919598	Dec 16, 2029	DS		
		8501698	Jun 20, 2027	DP U-493		
		8685934	May 26, 2030	U-1522		
		9616028	Nov 12, 2030	DP		
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649	002	6515117	Oct 04, 2025	DS DP U-493	I-841	Oct 18, 2022
		7919598	Dec 16, 2029	DS		
		8501698	Jun 20, 2027	DP U-493		
		8685934	May 26, 2030	U-1522		
		9616028	Nov 12, 2030	DP		
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649	003	6515117	Oct 04, 2025	DS DP U-493	I-841	Oct 18, 2022
		7919598	Dec 16, 2029	DS		
		8501698	Jun 20, 2027	DP U-493		
		8685934	May 26, 2030	U-1522		
		9616028	Nov 12, 2030	DP		
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649	004	6515117	Oct 04, 2025	DS DP U-493	I-841	Oct 18, 2022
		7919598	Dec 16, 2029	DS		
		8501698	Jun 20, 2027	DP U-493		
		8685934	May 26, 2030	U-1522		
		9616028	Nov 12, 2030	DP		
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649	005	6515117	Oct 04, 2025	DS DP U-493	I-841	Oct 18, 2022
		7919598	Dec 16, 2029	DS		
		8501698	Jun 20, 2027	DP U-493		
		8685934	May 26, 2030	U-1522		
		9616028	Nov 12, 2030	DP		

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<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649	005	6515117	Oct 04, 2025	DS DP U-493	I-841	Oct 18, 2022
		7919598	Dec 16, 2029	DS		
		8501698	Jun 20, 2027	DP U-493		
		8685934	May 26, 2030	U-1522		
		9616028	Nov 12, 2030	DP		
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874	001	6515117	Oct 04, 2025	DS DP U-493	NP	May 02, 2022
		7919598	Dec 16, 2029	DS		
		8501698	Jun 20, 2027	DP U-493		
		8628799	Jul 13, 2025	DP		
		8716251	Mar 21, 2028	DP		
		9616028	Nov 12, 2030	DP		
		RE44186	Jul 31, 2023	DS DP U-493		
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874	002	6515117	Oct 04, 2025	DS DP U-493	NP	May 02, 2022
		7919598	Dec 16, 2029	DS		
		8501698	Jun 20, 2027	DP U-493		
		8628799	Jul 13, 2025	DP		
		8716251	Mar 21, 2028	DP		
		9616028	Nov 12, 2030	DP		
		RE44186	Jul 31, 2023	DS DP U-493		
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874	003	6515117	Oct 04, 2025	DS DP U-493	NP	May 02, 2022
		7919598	Dec 16, 2029	DS		
		8501698	Jun 20, 2027	DP U-493		
		8628799	Jul 13, 2025	DP		
		8716251	Mar 21, 2028	DP		
		9616028	Nov 12, 2030	DP		
		RE44186	Jul 31, 2023	DS DP U-493		
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874	004	6515117	Oct 04, 2025	DS DP U-493	NP	May 02, 2022
		7919598	Dec 16, 2029	DS		
		8501698	Jun 20, 2027	DP U-493		
		8628799	Jul 13, 2025	DP		
		8716251	Mar 21, 2028	DP		
		9616028	Nov 12, 2030	DP		
		RE44186	Jul 31, 2023	DS DP U-493		
<u>DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE - QTERN</u>						
N 209091	001	6515117	Oct 04, 2025	DS DP U-493	I-804	May 02, 2022
		7919598	Dec 16, 2029	DS		
		8221786	Mar 21, 2028	DP		
		8361972	Mar 21, 2028	U-1976		
		8361972	Mar 21, 2028	U-1977		
		8361972	Mar 21, 2028	U-493		
		8501698	Jun 20, 2027	DP U-1976		
		8501698	Jun 20, 2027	DP U-1977		
		8501698	Jun 20, 2027	DP U-493		
		8628799	Jul 13, 2025	DP		
		8716251	Mar 21, 2028	DP		
		RE44186	Jul 31, 2023	DS DP U-493		
<u>DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE - QTERN</u>						
N 209091	002	6515117	Oct 04, 2025	DS DP U-493	NS	May 02, 2022
		7919598	Dec 16, 2029	DS		
		8221786	Mar 21, 2028	DP		
		8361972	Mar 21, 2028	U-493		
		8501698	Jun 20, 2027	DP U-493		
		8628799	Jul 13, 2025	DP		
		8716251	Mar 21, 2028	DP		
		RE44186	Jul 31, 2023	DS DP U-493		
<u>DAPSONE - ACZONE</u>						
N 207154	001	9161926	Nov 18, 2033	DP	NPP	Sep 10, 2022
		9517219	Nov 18, 2033	U-1033		

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<u>DAPTOMYCIN - CUBICIN</u>						
N 021572	002 8003673	Sep 04, 2028	U-1180			
<u>DAPTOMYCIN - CUBICIN RF</u>						
N 021572	003 9138456	Nov 23, 2030	DP			
<u>DAPTOMYCIN - DAPTOMYCIN</u>						
N 210282	001 10357535	Sep 11, 2033	DP U-3176			
	9655946	Sep 11, 2033	DP U-3175			
<u>DAPTOMYCIN - DAPTOMYCIN</u>						
N 210282	002 10357535	Sep 11, 2033	DP U-3176			
	9655946	Sep 11, 2033	DP U-3175			
<u>DAROLUTAMIDE - NUBEQA</u>						
N 212099	001 10010530	Jan 28, 2036	DS		NCE	Jul 30, 2024
	10383853	Jan 28, 2036	DS			
	10711013	Oct 27, 2030	DS DP			
	10835515	Jan 28, 2036	DP U-2605			
	11046713	Oct 27, 2030	DS			
	8975254	Oct 27, 2030	DS DP U-2605			
	9657003	Oct 27, 2030	DS DP U-2605			
<u>DARUNAVIR - PREZISTA</u>						
N 021976	001 7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976	002 7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976	003 7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976	004 7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976	005 7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976	006 7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 202895	001 7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DASABUVIR SODIUM; OMBITASVIR, PARITAPREVIR, RITONAVIR - VIEKIRA PAK (COPACKAGED)</u>						
N 206619	001 10201542	Oct 18, 2033	DP U-1753			
	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			

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<u>DASABUVIR SODIUM; OMBITASVIR, PARITAPREVIR, RITONAVIR - VIEKIRA PAK (COPACKAGED)</u>						
N 206619	001	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	
		10201541	May 17, 2032	DP		
		10201584	May 17, 2032		U-1889	
		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	7491725	Mar 28, 2026	DS DP	ODE-164	Nov 09, 2024
		7491725*PED	Sep 28, 2026		ODE-225	Dec 21, 2025
		8680103	Feb 04, 2025	DP	PED	May 09, 2025
		8680103*PED	Aug 04, 2025		PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986	002	7491725	Mar 28, 2026	DS DP	ODE-164	Nov 09, 2024
		7491725*PED	Sep 28, 2026		ODE-225	Dec 21, 2025
		8680103	Feb 04, 2025	DP	PED	May 09, 2025
		8680103*PED	Aug 04, 2025		PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986	003	7491725	Mar 28, 2026	DS DP	ODE-164	Nov 09, 2024
		7491725*PED	Sep 28, 2026		ODE-225	Dec 21, 2025
		8680103	Feb 04, 2025	DP	PED	May 09, 2025
		8680103*PED	Aug 04, 2025		PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986	004	7491725	Mar 28, 2026	DS DP	ODE-164	Nov 09, 2024
		7491725*PED	Sep 28, 2026		ODE-225	Dec 21, 2025
		8680103	Feb 04, 2025	DP	PED	May 09, 2025
		8680103*PED	Aug 04, 2025		PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986	005	7491725	Mar 28, 2026	DS DP	ODE-164	Nov 09, 2024
		7491725*PED	Sep 28, 2026		ODE-225	Dec 21, 2025
		8680103	Feb 04, 2025	DP	PED	May 09, 2025
		8680103*PED	Aug 04, 2025		PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986	006	7491725	Mar 28, 2026	DS DP	ODE-164	Nov 09, 2024
		7491725*PED	Sep 28, 2026		ODE-225	Dec 21, 2025
		8680103	Feb 04, 2025	DP	PED	May 09, 2025
		8680103*PED	Aug 04, 2025		PED	Jun 21, 2026
<u>DASIGLUCAGON HYDROCHLORIDE - ZEGALOGUE</u>						
N 214231	001	10442847	Feb 03, 2035	DS DP	NCE	Mar 22, 2026
<u>DASIGLUCAGON HYDROCHLORIDE - ZEGALOGUE (AUTOINJECTOR)</u>						
N 214231	002	10442847	Feb 03, 2035	DS DP	NCE	Mar 22, 2026

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<u>DEFERASIROX - EXJADE</u>						
N 021882	001				M-241 M-263	Jul 24, 2022 Jul 23, 2023
<u>DEFERASIROX - EXJADE</u>						
N 021882	002				M-241 M-263	Jul 24, 2022 Jul 23, 2023
<u>DEFERASIROX - EXJADE</u>						
N 021882	003				M-241 M-263	Jul 24, 2022 Jul 23, 2023
<u>DEFERASIROX - JADENU</u>						
N 206910	001 9283209	Nov 21, 2034	DS DP		M-241 M-263	Jul 24, 2022 Jul 23, 2023
<u>DEFERASIROX - JADENU</u>						
N 206910	002 9283209	Nov 21, 2034	DS DP		M-241 M-263	Jul 24, 2022 Jul 23, 2023
<u>DEFERASIROX - JADENU</u>						
N 206910	003 9283209	Nov 21, 2034	DS DP		M-241 M-263	Jul 24, 2022 Jul 23, 2023
<u>DEFERASIROX - JADENU SPRINKLE</u>						
N 207968	001				M-241 M-263	Jul 24, 2022 Jul 23, 2023
<u>DEFERASIROX - JADENU SPRINKLE</u>						
N 207968	002				M-241 M-263	Jul 24, 2022 Jul 23, 2023
<u>DEFERASIROX - JADENU SPRINKLE</u>						
N 207968	003				M-241 M-263	Jul 24, 2022 Jul 23, 2023
<u>DEFERIPRONE - FERRIPROX</u>						
N 021825	001				I-859	Apr 30, 2024
<u>DEFERIPRONE - FERRIPROX</u>						
N 021825	002				I-859	Apr 30, 2024
<u>DEFERIPRONE - FERRIPROX</u>						
N 208030	001 8703156	Oct 26, 2029	DP U-3083		I-859	Apr 30, 2024
<u>DEFERIPRONE - FERRIPROX</u>						
N 208030	002 8703156	Oct 26, 2029	DP U-3083		I-859	Apr 30, 2024
<u>DEFERIPRONE - FERRIPROX</u>						
N 212269	001 10780055 10940115 10940116	Oct 25, 2038 Oct 25, 2038 Oct 25, 2038	DP U-3083 DP U-3083 DP		I-859	Apr 30, 2024
<u>DEFIBROTIDE SODIUM - DEFITELIO</u>						
N 208114	001 11085043	Jun 22, 2032	DP		ODE-112	Mar 30, 2023
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	001				NCE ODE-130 ODE-252	Feb 09, 2022 Feb 09, 2024 Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	002				NCE ODE-130 ODE-252	Feb 09, 2022 Feb 09, 2024 Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	003				NCE ODE-130 ODE-252	Feb 09, 2022 Feb 09, 2024 Jun 07, 2026

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<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	004				NCE ODE-130 ODE-252	Feb 09, 2022 Feb 09, 2024 Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208685	001				NCE ODE-130 ODE-252	Feb 09, 2022 Feb 09, 2024 Jun 07, 2026
<u>DEGARELIX ACETATE - FIRMAGON</u>						
N 022201	001	10695398	Apr 27, 2032	U-1895		
		10729739	Feb 10, 2029	U-1978		
		10973870	Feb 10, 2029	U-1978		
		9415085	Apr 27, 2032	U-1895		
		9579359	Feb 10, 2029	U-1978		
<u>DEGARELIX ACETATE - FIRMAGON</u>						
N 022201	002	10695398	Apr 27, 2032	U-1895		
		10729739	Feb 10, 2029	U-1978		
		10973870	Feb 10, 2029	U-1978		
		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	I-815	Oct 24, 2022
		8252813	Oct 02, 2026	DP U-2028	NCE	Jun 19, 2022
		8273892	Aug 06, 2026	DS	GAIN	Jun 19, 2027
		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	I-815	Oct 24, 2022
		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		
		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>DEOXYCHOLIC ACID - KYBELLA</u>						
N 206333	001	10500214	Mar 02, 2030	DP		
		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; PSEUDOEPHEDRINE SULFATE - CLARINEX D 24 HOUR</u>						
N 021605	001	6979463	Mar 28, 2022	DP		
		7820199	Mar 28, 2022	DP		

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<u>DESMOPRESSIN ACETATE - NOCDURNA</u>						
N 022517	001	10307459	May 07, 2023	DP		
		11020448	May 21, 2029		U-2327	
		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-2326	
<u>DESMOPRESSIN ACETATE - NOCDURNA</u>						
N 022517	002	10137167	May 21, 2029		U-2327	
		10307459	May 07, 2023	DP		
		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	
		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	
		7579321	May 06, 2023		U-1980	
		9539302	Jun 15, 2030	DP		
<u>DESONIDE - VERDESO</u>						
N 021978	001	8460641	Aug 13, 2027	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	
		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	
		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	
		6673838	Mar 01, 2022	DS	U-860	
		8269040	Jul 05, 2027	DS		
<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082	001	10959996	Mar 07, 2036		U-3055	M-54 Jun 24, 2024
		10959996*PED	Sep 07, 2036			NCE Apr 03, 2022
		11179386	Mar 15, 2038	DP	U-1995	ODE-134 Apr 03, 2024
		11179386	Mar 15, 2038	DP	U-3055	PED Oct 03, 2022
		8524733	Apr 03, 2031	DS DP		PED Oct 03, 2024
		8524733*PED	Oct 03, 2031			PED Dec 24, 2024
		9233959	Sep 18, 2033	DP		
		9233959*PED	Mar 18, 2034			
		9296739	Sep 18, 2033	DP		
		9296739*PED	Mar 18, 2034			
		9550780	Sep 18, 2033	DS DP	U-1995	
		9550780	Sep 18, 2033	DS DP	U-3055	
		9550780*PED	Mar 18, 2034			

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<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082	001	9814708	Sep 18, 2033	DP		
		9814708*PED	Mar 18, 2034			
<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082	002	10959996	Mar 07, 2036	U-3055	M-54	Jun 24, 2024
		10959996*PED	Sep 07, 2036		NCE	Apr 03, 2022
		11179386	Mar 15, 2038	DP U-1995	ODE-134	Apr 03, 2024
		11179386	Mar 15, 2038	DP U-3055	PED	Oct 03, 2022
		8524733	Apr 03, 2031	DS DP	PED	Oct 03, 2024
		8524733*PED	Oct 03, 2031		PED	Dec 24, 2024
		9233959	Sep 18, 2033	DP		
		9233959*PED	Mar 18, 2034			
		9296739	Sep 18, 2033	DP		
		9296739*PED	Mar 18, 2034			
		9550780	Sep 18, 2033	DS DP U-1995		
		9550780	Sep 18, 2033	DS DP U-3055		
		9550780*PED	Mar 18, 2034			
		9814708	Sep 18, 2033	DP		
		9814708*PED	Mar 18, 2034			
<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082	003	10959996	Mar 07, 2036	U-3055	M-54	Jun 24, 2024
		10959996*PED	Sep 07, 2036		NCE	Apr 03, 2022
		11179386	Mar 15, 2038	DP U-1995	ODE-134	Apr 03, 2024
		11179386	Mar 15, 2038	DP U-3055	PED	Oct 03, 2022
		8524733	Apr 03, 2031	DS DP	PED	Oct 03, 2024
		8524733*PED	Oct 03, 2031		PED	Dec 24, 2024
		9233959	Sep 18, 2033	DP		
		9233959*PED	Mar 18, 2034			
		9296739	Sep 18, 2033	DP		
		9296739*PED	Mar 18, 2034			
		9550780	Sep 18, 2033	DS DP U-1995		
		9550780	Sep 18, 2033	DS DP U-3055		
		9550780*PED	Mar 18, 2034			
		9814708	Sep 18, 2033	DP		
		9814708*PED	Mar 18, 2034			
<u>DEXAMETHASONE - DEXAMETHASONE</u>						
A 215106	001				CGT	Apr 20, 2022
<u>DEXAMETHASONE - DEXAMETHASONE</u>						
A 215106	002				CGT	Apr 19, 2022
<u>DEXAMETHASONE - OZURDEX</u>						
N 022315	001	10076526	Jan 09, 2023	DP		
		10702539	Jan 09, 2023	U-1597		
		10702539	Jan 09, 2023	U-2868		
		10702539	Jan 09, 2023	U-985		
		6899717	Nov 01, 2023	U-1206		
		8034366	Jan 09, 2023	DP U-1204		
		8034366	Jan 09, 2023	DP U-1205		
		8034370	Jan 09, 2023	DP		
		8506987	Jan 09, 2023	U-1204		
		8506987	Jan 09, 2023	U-1205		
		9192511	Jan 09, 2023	DP		
<u>DEXAMETHASONE - DEXTENZA</u>						
N 208742	001	8409606	May 14, 2030	DP	I-800	Jun 20, 2022
		8563027	Feb 12, 2030	U-2487	I-876	Oct 07, 2024
		9254267	Sep 11, 2024	DP		
<u>DEXAMETHASONE - DEXYCU KIT</u>						
N 208912	001	10022502	Jun 22, 2034	U-2340		
		10028965	May 23, 2034	U-2340		
		10159683	May 23, 2034	DP		
		10799642	May 11, 2032	DP		
		6960346	Jul 03, 2023	DP		
		7560120	Sep 05, 2022	DP		

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<u>DEXAMETHASONE - HEMADY</u>						
N 211379	001 10537585	Dec 18, 2037	DP		ODE-271	Oct 03, 2026
<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 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				
	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			
	9233103	Mar 05, 2032		U-1805		
	9238029	Jan 17, 2026	DP			
<u>DEXLANSOPRAZOLE - DEXILANT</u>						
N 022287	002 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				
	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			
	9233103	Mar 05, 2032		U-1805		
	9238029	Jan 17, 2026	DP			
<u>DEXLANSOPRAZOLE - DEXILANT SOLUTAB</u>						
N 208056	001 6664276	Jan 30, 2023	DS DP U-950			
	6664276	Jan 30, 2023	DS DP U-951			
	6664276*PED	Jul 30, 2023				
	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				
	9011926	Feb 24, 2026	DP			
	9238029	Jan 17, 2026	DP			
	9241910	Mar 10, 2029	DP			

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<u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u>						
N 021038 002	10016396	Jan 04, 2032	DP			
	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			
	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	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			
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; SERDEXMETHYLPHENIDATE CHLORIDE - AZSTARYS</u>						
N 212994 001	10584112	Dec 09, 2037	DS DP		NCE	May 07, 2026
	10584113	Dec 09, 2037	DP			
	10759778	Dec 09, 2037	DP			
	10858341	Dec 09, 2037		U-3094		
	10954213	Dec 09, 2037		U-3094		
	9079928	Jul 27, 2032	DP			
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; SERDEXMETHYLPHENIDATE CHLORIDE - AZSTARYS</u>						
N 212994 002	10584112	Dec 09, 2037	DS DP		NCE	May 07, 2026
	10584113	Dec 09, 2037	DP			
	10759778	Dec 09, 2037	DP			
	10858341	Dec 09, 2037		U-3094		
	10954213	Dec 09, 2037		U-3094		
	9079928	Jul 27, 2032	DP			
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; SERDEXMETHYLPHENIDATE CHLORIDE - AZSTARYS</u>						
N 212994 003	10584112	Dec 09, 2037	DS DP		NCE	May 07, 2026
	10584113	Dec 09, 2037	DP			

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<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; SERDEXMETHYLPHENIDATE CHLORIDE - AZSTARYS</u>						
N 212994	003	10759778	Dec 09, 2037	DP		
		10858341	Dec 09, 2037	U-3094		
		10954213	Dec 09, 2037	U-3094		
		9079928	Jul 27, 2032	DP		
<u>DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE - NUEDEXTA</u>						
N 021879	001	7659282	Aug 13, 2026	U-1093		
		8227484	Jul 17, 2023	U-1093		
<u>DIAZEPAM - VALTOCO</u>						
N 211635	001	10265402	May 11, 2025	DP	NP	Jan 10, 2023
		8895546	Mar 27, 2029	DP	ODE-279	Jan 10, 2027
		8927497	Jul 21, 2025	DP U-2727		
		9642913	May 11, 2025	DP		
		9763876	Mar 27, 2029	DP U-2727		
<u>DIAZEPAM - VALTOCO</u>						
N 211635	002	10265402	May 11, 2025	DP	NP	Jan 10, 2023
		8895546	Mar 27, 2029	DP	ODE-279	Jan 10, 2027
		8927497	Jul 21, 2025	DP U-2727		
		9642913	May 11, 2025	DP		
		9763876	Mar 27, 2029	DP U-2727		
<u>DIAZEPAM - VALTOCO</u>						
N 211635	003	10265402	May 11, 2025	DP	NP	Jan 10, 2023
		8895546	Mar 27, 2029	DP	ODE-279	Jan 10, 2027
		8927497	Jul 21, 2025	DP U-2727		
		9642913	May 11, 2025	DP		
		9763876	Mar 27, 2029	DP U-2727		
<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		
<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	7662858	Feb 24, 2029	U-1035	NPP	May 25, 2024
		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		

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<u>DICLOFENAC SODIUM - PENNSAID</u>						
N 020947	001 8741956	Jul 10, 2029	U-1435			
<u>DICLOFENAC SODIUM - DYLOJECT</u>						
N 022396	001 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>DIFELIKEFALIN ACETATE - KORSUVA</u>						
N 214916	001 10017536	Nov 12, 2027	DS U-3204		NCE	Aug 23, 2026
	10138270	Nov 12, 2027	U-3204			
	10793596	Nov 12, 2027	DS DP U-3204			
	7402564	Nov 12, 2027	DS DP U-3204			
	7713937	Nov 12, 2027	DS DP U-3204			
	7727963	Nov 12, 2027	DS DP U-3204			
	8217007	Nov 12, 2027	U-3204			
	8236766	Nov 12, 2027	U-3204			
	8486894	Nov 12, 2027	U-3204			
	8536131	Nov 12, 2027	DS DP U-3204			
	9334305	Nov 12, 2027	U-3204			
	9359399	Nov 12, 2027	U-3204			
<u>DIHYDROERGOTAMINE MESYLATE - TRUDHESA</u>						
N 213436	001 10507295	Dec 25, 2032	DP		NP	Sep 02, 2024
	10940278	Jan 23, 2033	DP			
	11185497	Jan 04, 2039	U-3218			
	9550036	Sep 05, 2032	DP			
	9919117	Mar 17, 2033	DP U-3218			
<u>DIMETHYL FUMARATE - TECFIDERA</u>						
N 204063	001 10391160	Mar 13, 2035	U-3148		M-260	Feb 05, 2023
	10555993	Mar 13, 2035	U-3148			
	10959972	Nov 16, 2035	U-1384			
	10994003	Mar 13, 2035	U-3148			
	11007166	Nov 16, 2035	U-1384			
	11007167	Nov 16, 2035	U-1384			
	11129806	Nov 16, 2035	U-1384			
	8399514	Feb 07, 2028	U-1384			
<u>DIMETHYL FUMARATE - TECFIDERA</u>						
N 204063	002 10391160	Mar 13, 2035	U-3148		M-260	Feb 05, 2023
	10555993	Mar 13, 2035	U-3148			
	10959972	Nov 16, 2035	U-1384			
	10994003	Mar 13, 2035	U-3148			
	11007166	Nov 16, 2035	U-1384			
	11007167	Nov 16, 2035	U-1384			

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<u>DIMETHYL FUMARATE - TECFIDERA</u>						
N 204063	002	11129806	Nov 16, 2035			
		8399514	Feb 07, 2028			
<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>DIROXIMEL FUMARATE - VUMERITY</u>						
N 211855	001	10080733	Sep 20, 2033	DS DP	U-1384	
		8669281	Sep 20, 2033	DS DP		
		9090558	Sep 20, 2033		U-1384	
<u>DOCETAXEL - DOCETAXEL</u>						
N 205934	001	10842770	Aug 07, 2031	DP	U-2998	
		8940786	Sep 30, 2033	DP	U-1789	
		9308195	Sep 30, 2033	DP		
		9763880	Sep 30, 2033		U-2558	
		9763880	Sep 30, 2033		U-2559	
		9763880	Sep 30, 2033		U-2560	
		9763880	Sep 30, 2033		U-2561	
		9763880	Sep 30, 2033		U-2562	
		9763880	Sep 30, 2033		U-2563	
		9763880	Sep 30, 2033		U-2564	
<u>DOCETAXEL - DOCETAXEL</u>						
N 205934	002	10842770	Aug 07, 2031	DP	U-2998	
		8940786	Sep 30, 2033	DP	U-1789	
		9308195	Sep 30, 2033	DP		
		9763880	Sep 30, 2033		U-2558	
		9763880	Sep 30, 2033		U-2559	
		9763880	Sep 30, 2033		U-2560	
		9763880	Sep 30, 2033		U-2561	
		9763880	Sep 30, 2033		U-2562	
		9763880	Sep 30, 2033		U-2563	
		9763880	Sep 30, 2033		U-2564	
<u>DOCETAXEL - DOCETAXEL</u>						
N 205934	003	10842770	Aug 07, 2031	DP	U-2998	
		8940786	Sep 30, 2033	DP	U-1789	
		9308195	Sep 30, 2033	DP		
		9763880	Sep 30, 2033		U-2558	
		9763880	Sep 30, 2033		U-2559	
		9763880	Sep 30, 2033		U-2560	
		9763880	Sep 30, 2033		U-2561	
		9763880	Sep 30, 2033		U-2562	
		9763880	Sep 30, 2033		U-2563	
		9763880	Sep 30, 2033		U-2564	
<u>DOLUTEGRAVIR SODIUM - TIVICAY</u>						
N 204790	001	8129385	Oct 05, 2027	DS DP		
		9242986	Dec 08, 2029	DS DP		
<u>DOLUTEGRAVIR SODIUM - TIVICAY</u>						
N 204790	002	8129385	Oct 05, 2027	DS DP		
		9242986	Dec 08, 2029	DS DP		
<u>DOLUTEGRAVIR SODIUM - TIVICAY</u>						
N 204790	003	8129385	Oct 05, 2027	DS DP		
		9242986	Dec 08, 2029	DS DP		
<u>DOLUTEGRAVIR SODIUM - TIVICAY PD</u>						
N 213983	001	8129385	Oct 05, 2027	DS DP		
		9242986	Dec 08, 2029	DS DP		
<u>DOLUTEGRAVIR SODIUM; LAMIVUDINE - DOVATO</u>						
N 211994	001	8129385	Oct 05, 2027	DS DP	I-839	Aug 06, 2023
		9242986	Dec 08, 2029	DS DP	NC	Apr 08, 2022

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<u>DOLUTEGRAVIR SODIUM; RILPIVIRINE HYDROCHLORIDE - JULUCA</u>						
N 210192	001	10426780	Jan 24, 2031	DS DP	U-257	
		7125879	Apr 21, 2025	DS DP	U-257	
		8080551	Apr 11, 2023	DS DP		
		8101629	Aug 09, 2022	DP		
		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	

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<u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u>						
N 206439	003	8598233	Nov 22, 2025	DP		
		8598233*PED	May 22, 2026			
<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	I-827	Sep 19, 2022
		8486975	Oct 07, 2031	DS DP U-2630	NCE	Aug 30, 2023
<u>DORAVIRINE; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE - DELSTRIGO</u>						
N 210807	001	10603282	Nov 29, 2036	DP	I-806	Sep 19, 2022
		10842751	Nov 29, 2036	DP	NCE	Aug 30, 2023
		8486975	Oct 07, 2031	DS DP U-2395		
		8486975	Oct 07, 2031	DS DP U-2629		
<u>DOXEPIIN HYDROCHLORIDE - SILENOR</u>						
N 022036	001	10238620	May 18, 2027	U-620		
		10548871	Apr 11, 2028	U-620		
		10653660	Jul 20, 2027	U-620		
		10653662	May 18, 2027	U-620		
		11096920	Apr 11, 2028	U-620		
		11110074	Jul 20, 2027	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	10238620	May 18, 2027	U-620		
		10548871	Apr 11, 2028	U-620		
		10653660	Jul 20, 2027	U-620		
		10653662	May 18, 2027	U-620		
		11096920	Apr 11, 2028	U-620		
		11110074	Jul 20, 2027	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	U-1063		
		7211267	Apr 05, 2022	U-925		
		7232572	Apr 05, 2022	U-925		

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<u>DROSPIRENONE - SLYND</u>						
N 211367	001 11123299	Jun 28, 2031	DP			
	9603860	Jun 28, 2031	U-2553			
<u>DROSPIRENONE; ESTETROL - NEXTSTELLIS</u>						
N 214154	001 7732430	Mar 02, 2025	DP U-3152		NCE	Apr 15, 2026
<u>DROSPIRENONE; ESTRADIOL - ANGELIO</u>						
N 021355	001 8906890	Oct 22, 2031	DP			
<u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM - BEYAZ</u>						
N 022532	001 6441168	Jul 30, 2022	DS			
	7163931	Mar 03, 2022	U-1			
	8617597	Feb 08, 2030	DP			
<u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM - SAFYRAL</u>						
N 022574	001 7163931	Mar 03, 2022	U-1			
	8617597	Feb 08, 2030	DP			
<u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u>						
N 021427	001				NPP	Apr 20, 2023
<u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u>						
N 021427	002				NPP	Apr 20, 2023
<u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u>						
N 021427	004				NPP	Apr 20, 2023
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	001 10413525	Apr 13, 2037	DP			
	10959982	Apr 13, 2037	DP			
	9839626	Apr 13, 2037	DP			
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	002 10413525	Apr 13, 2037	DP			
	10959982	Apr 13, 2037	DP			
	9839626	Apr 13, 2037	DP			
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	003 10413525	Apr 13, 2037	DP			
	10959982	Apr 13, 2037	DP			
	9839626	Apr 13, 2037	DP			
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	004 10413525	Apr 13, 2037	DP			
	10959982	Apr 13, 2037	DP			
	9839626	Apr 13, 2037	DP			
<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				NCE	May 05, 2022
					ODE-144	May 05, 2024

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<u>EDARAVONE - RADICAVA</u>						
N 209176	002				NCE ODE*	May 05, 2022 May 05, 2024
<u>EDOXABAN TOSYLATE - SAVAYSA</u>						
N 206316	001	7365205	Apr 18, 2027	DS	M-243	Aug 09, 2022
		9149532	Mar 28, 2028	DP		
<u>EDOXABAN TOSYLATE - SAVAYSA</u>						
N 206316	002	7365205	Apr 18, 2027	DS	M-243	Aug 09, 2022
		9149532	Mar 28, 2028	DP		
<u>EDOXABAN TOSYLATE - SAVAYSA</u>						
N 206316	003	7365205	Apr 18, 2027	DS	M-243	Aug 09, 2022
		9149532	Mar 28, 2028	DP		
<u>EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - ATRIPLA</u>						
N 021937	001	8592397	Jan 13, 2024	DP U-1170		
		8592397	Jan 13, 2024	DP U-750		
		8598185	Apr 28, 2029	DP		
		8716264	Jan 13, 2024	DP U-257		
		9018192	Jun 13, 2026	U-1170		
		9018192	Jun 13, 2026	U-750		
		9457036	Jan 13, 2024	DP U-257		
		9545414	Jun 13, 2026	DP U-1170		
		9545414	Jun 13, 2026	DP U-750		
		9744181	Jan 13, 2024	DP U-257		
<u>EFINACONAZOLE - JUBLIA</u>						
N 203567	001	10105444	Jul 08, 2030	DP	NPP	Apr 27, 2023
		10342875	Oct 02, 2034	DP U-2720		
		10478601	Apr 25, 2035	DP U-2721		
		10512640	Jan 03, 2028	U-1969		
		10828293	Oct 02, 2034	U-2720		
		10828369	Jan 03, 2028	DP		
		10864274	Oct 02, 2034	U-2720		
		7214506	Feb 22, 2026	U-281		
		8039494	Jul 08, 2030	U-281		
		8486978	Oct 24, 2030	DP		
		9302009	Oct 24, 2030	DP		
		9566272	Jan 03, 2028	U-1969		
		9662394	Oct 02, 2034	DP		
		9861698	Jul 08, 2030	DP		
		9877955	Jan 03, 2028	U-1969		
<u>ELAGOLIX SODIUM - ORILISSA</u>						
N 210450	001	10537572	Sep 01, 2036	U-2735	NCE	Jul 23, 2023
		10682351	Sep 01, 2036	U-2850		
		7056927	Sep 10, 2024	DS DP		
		7176211	Jul 06, 2024	U-2360		
		7419983	Jul 06, 2024	DS DP U-2360		
<u>ELAGOLIX SODIUM - ORILISSA</u>						
N 210450	002	7056927	Sep 10, 2024	DS DP	NCE	Jul 23, 2023
		7176211	Jul 06, 2024	U-2360		
		7419983	Jul 06, 2024	DS DP U-2360		
<u>ELAGOLIX SODIUM, ESTRADIOL, NORETHINDRONE ACETATE; ELAGOLIX SODIUM - ORIAHNN (COPACKAGED)</u>						
N 213388	001	10881659	Mar 14, 2034	U-2842	NCE	Jul 23, 2023
		11045470	Mar 14, 2034	U-2842	NP	May 29, 2023
		7056927	Sep 10, 2024	DS DP		
		7419983	Jul 06, 2024	DS DP		
<u>ELBASVIR; GRAZOPREVRIL - ZEPATIER</u>						
N 208261	001	7973040	Jul 24, 2029	DS DP U-1813		
		8871759	May 04, 2031	DS DP U-1813		
<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 212273	001	10022352	Apr 09, 2027	DP U-2651	NCE	Feb 12, 2023
		10022352	Apr 09, 2027	DP U-3156	NCE	Oct 21, 2024
		10081621	Mar 25, 2031	DP U-2652	NPP	Jun 08, 2024

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<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 212273 001	10081621	Mar 25, 2031	DP U-3032		ODE-275	Oct 21, 2026
	10081621	Mar 25, 2031	DP U-3157		ODE-323	Dec 21, 2027
	10239867	Apr 09, 2027	DS DP U-2653		ODE-357	Jun 08, 2028
	10239867	Apr 09, 2027	DS DP U-3033			
	10239867	Apr 09, 2027	DS DP U-3158			
	10646481	Aug 13, 2029	DP			
	10758534	Oct 06, 2035	DS DP U-2645			
	10758534	Oct 06, 2035	DS DP U-3028			
	10758534	Oct 06, 2035	DS DP U-3144			
	10793547	Dec 08, 2037	DS DP U-2645			
	10793547	Dec 08, 2037	DS DP U-3028			
	10793547	Dec 08, 2037	DS DP U-3144			
	11179367	Dec 08, 2037	DP U-3253			
	7495103	May 20, 2027	DS DP			
	7645789	May 01, 2027	DS DP			
	7776905	Jun 03, 2027	DS DP			
	8324242	Aug 05, 2027	U-2645			
	8324242	Aug 05, 2027	U-3028			
	8324242	Aug 05, 2027	U-3144			
	8354427	Jul 06, 2026	U-2646			
	8354427	Jul 06, 2026	U-3029			
	8354427	Jul 06, 2026	U-3145			
	8410274	Dec 28, 2026	DP			
	8415387	Nov 12, 2027	U-2645			
	8415387	Nov 12, 2027	U-3028			
	8415387	Nov 12, 2027	U-3144			
	8598181	May 01, 2027	U-2645			
	8598181	May 01, 2027	U-3028			
	8598181	May 01, 2027	U-3144			
	8623905	May 01, 2027	DS DP			
	8629162	Jun 24, 2025	U-2648			
	8629162	Jun 24, 2025	U-3030			
	8629162	Jun 24, 2025	U-3146			
	8754224	Dec 28, 2026	DS DP			
	9012496	Jul 15, 2033	U-2649			
	9012496	Jul 15, 2033	U-3154			
	9670163	Dec 28, 2026	DP U-2650			
	9670163	Dec 28, 2026	DP U-3031			
	9670163	Dec 28, 2026	DP U-3155			
	9931334	Dec 28, 2026	DP U-2650			
	9931334	Dec 28, 2026	DP U-3031			
	9931334	Dec 28, 2026	DP U-3155			
	9974781	Apr 09, 2027	DP U-2645			
	9974781	Apr 09, 2027	DP U-3028			
	9974781	Apr 09, 2027	DP U-3144			
<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 212273 002	10022352	Apr 09, 2027	DP U-3156		NCE	Feb 12, 2023
	10081621	Mar 25, 2031	DP U-3157		NCE	Oct 21, 2024
	10239867	Apr 09, 2027	DS DP U-3158		NPP	Jun 08, 2024
	10646481	Aug 13, 2029	DP		ODE-357	Jun 08, 2028
	10758534	Oct 06, 2035	DS DP U-3144			
	10793547	Dec 08, 2037	DS DP U-3144			
	11179367	Dec 08, 2037	DP U-3253			
	7495103	May 20, 2027	DS DP			
	7645789	May 01, 2027	DS DP			
	7776905	Jun 03, 2027	DS DP			
	8324242	Aug 05, 2027	U-3144			
	8354427	Jul 06, 2026	U-3145			
	8410274	Dec 28, 2026	DP			
	8415387	Nov 12, 2027	U-3144			
	8598181	May 01, 2027	U-3144			
	8623905	May 01, 2027	DS DP			
	8629162	Jun 24, 2025	U-3146			
	8754224	Dec 28, 2026	DS DP			
	9012496	Jul 15, 2033	U-3154			
	9670163	Dec 28, 2026	DP U-3155			
	9931334	Dec 28, 2026	DP U-3155			
	9974781	Apr 09, 2027	DP U-3144			

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<u>ELIGLUSTAT TARTRATE - CERDELGA</u>						
N 205494 001	10888544	Dec 13, 2038	U-3040			
	10888544	Dec 13, 2038	U-3041			
	10888547	Jan 31, 2031	U-3042			
	10888547	Jan 31, 2031	U-3043			
	6916802	Apr 29, 2022	DS U-1571			
	7196205	Jun 26, 2026	DS			
	7253185	Apr 29, 2022	DP			
	7615573	Apr 29, 2022	U-1571			
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 001	7160870	Nov 20, 2022	DS DP U-1306		ODE-210	Nov 16, 2025
	7160870	Nov 20, 2022	DS DP U-2451			
	7160870*PED	May 20, 2023				
	7547719	Jul 13, 2025	DS DP U-1306			
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-2451			
	7547719	Jul 13, 2025	DS DP U-2452			
	7547719*PED	Jan 13, 2026				
	7795293	May 21, 2023	U-1306			
	7795293	May 21, 2023	U-2451			
	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-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-2451			
	8828430*PED	Feb 01, 2028				
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 002	7160870	Nov 20, 2022	DS DP U-1306		ODE-210	Nov 16, 2025
	7160870	Nov 20, 2022	DS DP U-2451			
	7160870*PED	May 20, 2023				
	7547719	Jul 13, 2025	DS DP U-1306			
	7547719	Jul 13, 2025	DS DP U-1575			
	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				
	7795293	May 21, 2023	U-1306			
	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-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-2451			
	8828430*PED	Feb 01, 2028				
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 003	7160870	Nov 20, 2022	DS DP U-1306		ODE-210	Nov 16, 2025
	7160870	Nov 20, 2022	DS DP U-2451			
	7160870*PED	May 20, 2023				
	7547719	Jul 13, 2025	DS DP U-1306			
	7547719	Jul 13, 2025	DS DP U-1575			

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<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 003	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				
	7795293	May 21, 2023			U-1306	
	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-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-2451			
	8828430*PED	Feb 01, 2028				
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 004	7160870	Nov 20, 2022	DS DP U-1306		ODE-210	Nov 16, 2025
	7160870	Nov 20, 2022	DS DP U-2451			
	7160870*PED	May 20, 2023				
	7547719	Jul 13, 2025	DS DP U-1306			
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-2451			
	7547719	Jul 13, 2025	DS DP U-2452			
	7547719*PED	Jan 13, 2026				
	7795293	May 21, 2023			U-1306	
	7795293	May 21, 2023			U-2451	
	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-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-2451			
	8828430*PED	Feb 01, 2028				
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 005	7160870	Nov 20, 2022	DS DP U-1306		ODE-210	Nov 16, 2025
	7160870	Nov 20, 2022	DS DP U-1575			
	7160870	Nov 20, 2022	DS DP U-1714			
	7160870	Nov 20, 2022	DS DP U-930			
	7160870*PED	May 20, 2023				
	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				
	7795293	May 21, 2023			U-1306	
	7795293	May 21, 2023			U-1575	
	7795293	May 21, 2023			U-930	
	7795293*PED	Nov 21, 2023				
<u>ELTROMBOPAG OLAMINE - PROMACTA KIT</u>						
N 207027 001	7160870	Nov 20, 2022	DS DP U-1306		ODE*	Nov 16, 2025
	7160870	Nov 20, 2022	DS DP U-1736			
	7160870*PED	May 20, 2023				
	7547719	Jul 13, 2025	DS DP U-1306			
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-1736			

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<u>ELTROMBOPAG OLAMINE - PROMACTA KIT</u>						
N 207027	001	7547719	Jul 13, 2025	DS DP	U-2452	
		7547719*PED	Jan 13, 2026			
		7795293	May 21, 2023		U-1306	
		7795293	May 21, 2023		U-1736	
		7795293*PED	Nov 24, 2023			
<u>ELTROMBOPAG OLAMINE - PROMACTA KIT</u>						
N 207027	002	7160870	Nov 20, 2022	DS DP	U-1306	ODE* Nov 16, 2025
		7160870	Nov 20, 2022	DS DP	U-1736	
		7160870*PED	May 20, 2023			
		7547719	Jul 13, 2025	DS DP	U-1306	
		7547719	Jul 13, 2025	DS DP	U-1575	
		7547719	Jul 13, 2025	DS DP	U-1736	
		7547719	Jul 13, 2025	DS DP	U-2452	
		7547719*PED	Jan 13, 2026			
		7795293	May 21, 2023		U-1306	
		7795293	May 21, 2023		U-1736	
		7795293*PED	Nov 21, 2023			
<u>ELUXADOLINE - VIBERZI</u>						
N 206940	001	10188632	Mar 14, 2033		DP	
		10213415	Mar 14, 2025	DS	U-2152	
		11007179	Mar 14, 2033		DP	
		11090291	Mar 14, 2033		DP	
		11160792	Mar 14, 2033		DP	
		7741356	May 27, 2029	DS DP		
		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	10188632	Mar 14, 2033		DP	
		10213415	Mar 14, 2025	DS	U-2152	
		11007179	Mar 14, 2033		DP	
		11090291	Mar 14, 2033		DP	
		11160792	Mar 14, 2033		DP	
		7741356	May 27, 2029	DS DP		
		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	
		7176220*PED	Feb 27, 2027			
		7635704	Oct 26, 2026	DS DP	U-257	
		7635704*PED	Apr 26, 2027			
		8981103	Oct 26, 2026	DS DP		
		8981103*PED	Apr 26, 2027			

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<u>ELVITEGRAVIR - VITEKTA</u>						
N 203093	002	7176220	Aug 27, 2026	DS DP	U-257	
		7176220*PED	Feb 27, 2027			
		7635704	Oct 26, 2026	DS DP	U-257	
		7635704*PED	Apr 26, 2027			
		8981103	Oct 26, 2026	DS DP		
		8981103*PED	Apr 26, 2027			
<u>EMPAGLIFLOZIN - JARDIANCE</u>						
N 204629	001	10258637	Apr 03, 2034		U-2290	I-869 Aug 18, 2024
		11090323	Apr 03, 2034		U-3191	
		7579449	Aug 01, 2028	DS		
		7713938	Apr 15, 2027	DS DP		
		8551957	Oct 14, 2029		U-1651	
		9949997	May 17, 2034		U-2292	
		9949997	May 17, 2034		U-3199	
		9949998	Jun 11, 2034		U-2290	
<u>EMPAGLIFLOZIN - JARDIANCE</u>						
N 204629	002	10258637	Apr 03, 2034		U-2290	
		11090323	Apr 03, 2034		U-3191	
		7579449	Aug 01, 2028	DS		
		7713938	Apr 15, 2027	DS DP		
		8551957	Oct 14, 2029		U-1651	
		9949997	May 17, 2034		U-2292	
		9949998	Jun 11, 2034		U-2290	
<u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u>						
N 206073	001	10258637	Apr 03, 2034		U-2290	M-252 Mar 30, 2023
		11033552	May 04, 2027	DP		M-258 Jul 03, 2022
		11090323	Apr 03, 2034		U-3191	
		7407955	May 02, 2025	DS DP		
		7579449	Aug 01, 2028	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	Y
		8883805	Nov 26, 2025	DP		
		9173859	May 04, 2027	DP	U-1772	Y
		9949998	Jun 11, 2034		U-2290	
<u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u>						
N 206073	002	10258637	Apr 03, 2034		U-2290	M-252 Mar 30, 2023
		11033552	May 04, 2027	DP		M-258 Jul 03, 2022
		11090323	Apr 03, 2034		U-3191	
		7407955	May 02, 2025	DS DP		
		7579449	Aug 01, 2028	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	Y
		8883805	Nov 26, 2025	DP		
		9173859	May 04, 2027	DP	U-1772	Y
		9949998	Jun 11, 2034		U-2290	
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614	001	10022379	Apr 02, 2029	DP	U-2732	
		10258637	Apr 03, 2034		U-2731	
		10406172	Jun 15, 2030	DP	U-2733	
		10596120	Mar 07, 2032	DP	U-2776	
		10596120	Mar 07, 2032	DP	U-2790	
		11090323	Apr 03, 2034		U-3192	
		7407955	May 02, 2025	DS DP		
		7579449	Aug 01, 2028	DS		
		7713938	Apr 15, 2027	DS DP		
		8119648	Aug 12, 2023		U-1652	
		8178541	Aug 12, 2023	DP	U-1652	

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<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 001	8551957	Oct 14, 2029	DP U-2730			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 002	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10406172	Jun 15, 2030	DP U-2733			
	10596120	Mar 07, 2032	DP U-2776			
	10596120	Mar 07, 2032	DP U-2790			
	11090323	Apr 03, 2034	U-3192			
	7407955	May 02, 2025	DS DP			
	7579449	Aug 01, 2028	DS			
	7713938	Apr 15, 2027	DS DP			
	8119648	Aug 12, 2023	U-1652			
	8178541	Aug 12, 2023	DP U-1652			
	8551957	Oct 14, 2029	DP U-2730			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 003	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10406172	Jun 15, 2030	DP U-2733			
	10596120	Mar 07, 2032	DP U-2776			
	10596120	Mar 07, 2032	DP U-2790			
	11090323	Apr 03, 2034	U-3192			
	7407955	May 02, 2025	DS DP			
	7579449	Aug 01, 2028	DS			
	7713938	Apr 15, 2027	DS DP			
	8119648	Aug 12, 2023	U-1652			
	8178541	Aug 12, 2023	DP U-1652			
	8551957	Oct 14, 2029	DP U-2730			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 004	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10406172	Jun 15, 2030	DP U-2733			
	10596120	Mar 07, 2032	DP U-2776			
	10596120	Mar 07, 2032	DP U-2790			
	11090323	Apr 03, 2034	U-3192			
	7407955	May 02, 2025	DS DP			
	7579449	Aug 01, 2028	DS			
	7713938	Apr 15, 2027	DS DP			
	8119648	Aug 12, 2023	U-1652			
	8178541	Aug 12, 2023	DP U-1652			
	8551957	Oct 14, 2029	DP U-2730			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111 001	10258637	Apr 03, 2034	U-2290			
	10610489	Sep 30, 2030	DP			
	11090323	Apr 03, 2034	U-3193			
	7579449	Aug 01, 2028	DS			
	7713938	Apr 15, 2027	DS DP			

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<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111	002	10258637	Apr 03, 2034			U-2290
		10610489	Sep 30, 2030	DP		
		11090323	Apr 03, 2034			U-3193
		7579449	Aug 01, 2028	DS		
		7713938	Apr 15, 2027	DS DP		
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111	003	10258637	Apr 03, 2034			U-2290
		10610489	Sep 30, 2030	DP		
		11090323	Apr 03, 2034			U-3193
		7579449	Aug 01, 2028	DS		
		7713938	Apr 15, 2027	DS DP		
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111	004	10258637	Apr 03, 2034			U-2290
		10610489	Sep 30, 2030	DP		
		11090323	Apr 03, 2034			U-3193
		7579449	Aug 01, 2028	DS		
		7713938	Apr 15, 2027	DS DP		
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	001	10258637	Apr 03, 2034			U-2290
		10596120	Mar 07, 2032	DP		U-2775
		10596120	Mar 07, 2032	DP		U-2792
		11090323	Apr 03, 2034			U-3193
		7579449	Aug 01, 2028	DS		
		7713938	Apr 15, 2027	DS DP		
		9949998	Jun 11, 2034			U-2290
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	002	10258637	Apr 03, 2034			U-2290
		10596120	Mar 07, 2032	DP		U-2775
		10596120	Mar 07, 2032	DP		U-2792
		11090323	Apr 03, 2034			U-3193
		7579449	Aug 01, 2028	DS		
		7713938	Apr 15, 2027	DS DP		
		9949998	Jun 11, 2034			U-2290
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	003	10258637	Apr 03, 2034			U-2290
		10596120	Mar 07, 2032	DP		U-2775
		10596120	Mar 07, 2032	DP		U-2792
		11090323	Apr 03, 2034			U-3193
		7579449	Aug 01, 2028	DS		
		7713938	Apr 15, 2027	DS DP		
		9949998	Jun 11, 2034			U-2290
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	004	10258637	Apr 03, 2034			U-2290
		10596120	Mar 07, 2032	DP		U-2775
		10596120	Mar 07, 2032	DP		U-2792
		11090323	Apr 03, 2034			U-3193
		7579449	Aug 01, 2028	DS		
		7713938	Apr 15, 2027	DS DP		
		9949998	Jun 11, 2034			U-2290
<u>EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR ALAFENAMIDE FUMARATE - ODEFSEY</u>						
N 208351	001	7125879	Apr 21, 2025	DS DP		U-257
		7390791	Apr 17, 2025	DS DP		
		7390791*PED	Oct 17, 2025			
		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
		8754065*PED	Feb 15, 2033			
		9296769	Aug 15, 2032	DS DP		U-257
		9296769*PED	Feb 15, 2033			

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<u>EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR DISOPROXIL FUMARATE - COMPLERA</u>						
N 202123	001	10857102	Jan 14, 2033	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	7390791	Apr 17, 2025	DS DP	I-812	Oct 03, 2022
		7390791*PED	Oct 17, 2025		ODE-284	Sep 28, 2024
		7803788	Feb 02, 2022	U-1663	ODE-285	Sep 28, 2024
		8754065	Aug 15, 2032	DS DP U-1259		
		8754065	Aug 15, 2032	DS DP U-1663		
		8754065	Aug 15, 2032	DS DP U-257		
		8754065*PED	Feb 15, 2033			
		9296769	Aug 15, 2032	DS DP U-1259		
		9296769	Aug 15, 2032	DS DP U-1663		
		9296769	Aug 15, 2032	DS DP U-257		
		9296769*PED	Feb 15, 2033			
<u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u>						
N 021752	001	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>ENALAPRIL MALEATE - ENALAPRIL MALEATE</u>						
A 212408	001				PC	Feb 13, 2022
<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		
		10772868	Mar 25, 2036	DP		
		10786482	Mar 25, 2036	DP		
		11040023	Mar 25, 2036	DP		
		11141405	Mar 25, 2036	DP		
		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	NCE	Aug 01, 2022
		10294215	Jan 07, 2033	DP U-2087	ODE-151	Aug 01, 2024
		10610125	Jun 21, 2030	U-2087		

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<u>ENASIDENIB MESYLATE - IDHIFA</u>						
N 209606 001	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		NCE	Aug 01, 2022
	10294215	Jan 07, 2033	DP U-2087		ODE-151	Aug 01, 2024
	10610125	Jun 21, 2030	U-2087			
	9512107	Jan 07, 2033	DS DP U-2087			
	9732062	Sep 16, 2034	DS			
	9738625	Aug 01, 2034	DS			
<u>ENCORAFENIB - BRAFTOVI</u>						
N 210496 001	10005761	Aug 27, 2030		U-2335	I-826	Apr 08, 2023
	8501758	Mar 04, 2031	DS DP		NCE	Jun 27, 2023
	8541575	Feb 26, 2030	DS DP U-2335		ODE-194	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	I-826	Apr 08, 2023
	10005761	Aug 27, 2030		U-2802	NCE	Jun 27, 2023
	10005761	Aug 27, 2030		U-2803	ODE-194	Jun 27, 2025
	10258622	Nov 21, 2032		U-2802		
	8501758	Mar 04, 2031	DS DP			
	8541575	Feb 26, 2030	DS DP U-2335			
	8541575	Feb 26, 2030	DS DP U-2802			
	8541575	Feb 26, 2030	DS DP U-2803			
	8946250	Jul 23, 2029	DS DP			
	9314464	Jul 04, 2031		U-2336		
	9314464	Jul 04, 2031		U-2802		
	9314464	Jul 04, 2031		U-2803		
	9387208	Nov 21, 2032	DP			
	9474754	Aug 05, 2033	DP U-2802			
	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		
	9850230	Aug 27, 2030		U-2802		
	9850230	Aug 27, 2030		U-2803		
<u>ENTRECTINIB - ROZLYTREK</u>						
N 212725 001	10231965	Feb 17, 2035		U-2617	NCE	Aug 15, 2024
	10231965	Feb 17, 2035		U-2618	ODE-265	Aug 15, 2026
	10398693	Jul 18, 2038	DP		ODE-313	Aug 15, 2026
	10561651	Feb 19, 2035		U-2745		
	10738037	May 18, 2037	DS DP U-2946			
	11091469	May 18, 2037		U-2617		
	11091469	May 18, 2037		U-2618		
	8299057	Mar 01, 2029	DS DP			
	8673893	Jul 08, 2028		U-2617		
	8673893	Jul 08, 2028		U-2618		
	9029356	Jul 08, 2028	DS DP			
	9085558	Jul 08, 2028	DP			
	9085565	May 22, 2033	DS DP			
	9255087	Jul 08, 2028		U-2617		
	9255087	Jul 08, 2028		U-2618		
	9616059	Jul 08, 2028		U-2618		
	9649306	May 22, 2033		U-2617		
	9649306	May 22, 2033		U-2618		

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<u>ENTRECTINIB - ROZLYTREK</u>						
N 212725 002	10231965	Feb 17, 2035		U-2617	NCE	Aug 15, 2024
	10231965	Feb 17, 2035		U-2618	ODE-265	Aug 15, 2026
	10398693	Jul 18, 2038	DP		ODE-313	Aug 15, 2026
	10561651	Feb 19, 2035		U-2745		
	10738037	May 18, 2037	DS DP	U-2946		
	11091469	May 18, 2037		U-2617		
	11091469	May 18, 2037		U-2618		
	8299057	Mar 01, 2029	DS DP			
	8673893	Jul 08, 2028		U-2617		
	8673893	Jul 08, 2028		U-2618		
	9029356	Jul 08, 2028	DS DP			
	9085558	Jul 08, 2028		DP		
	9085565	May 22, 2033	DS DP			
	9255087	Jul 08, 2028		U-2617		
	9255087	Jul 08, 2028		U-2618		
	9616059	Jul 08, 2028		U-2618		
	9649306	May 22, 2033		U-2617		
	9649306	May 22, 2033		U-2618		
<u>ENZALUTAMIDE - XTANDI</u>						
N 203415 001	7709517	Aug 13, 2027	DS DP		I-808	Dec 16, 2022
	8183274	Aug 24, 2026		U-1281		
	8183274	Aug 24, 2026		U-1588		
	8183274	Aug 24, 2026		U-2345		
	8183274	Aug 24, 2026		U-2708		
	9126941	May 15, 2026		U-1588		
	9126941	May 15, 2026		U-2345		
	9126941	May 15, 2026		U-2708		
<u>ENZALUTAMIDE - XTANDI</u>						
N 213674 001	7709517	Aug 13, 2027	DS DP		I-808	Dec 16, 2022
	8183274	Aug 24, 2026		U-2345		
	8183274	Aug 24, 2026		U-2708		
	9126941	May 15, 2026		U-2345		
	9126941	May 15, 2026		U-2708		
<u>ENZALUTAMIDE - XTANDI</u>						
N 213674 002	7709517	Aug 13, 2027	DS DP		I-808	Dec 16, 2022
	8183274	Aug 24, 2026		U-2345		
	8183274	Aug 24, 2026		U-2708		
	9126941	May 15, 2026		U-2345		
	9126941	May 15, 2026		U-2708		
<u>EPHEDRINE SULFATE - EMERPHED</u>						
N 213407 001	11090278	May 16, 2040		U-3183		
<u>EPHEDRINE SULFATE - EPHEDRINE SULFATE</u>						
N 213994 001	10869845	Jan 22, 2040	DP			
<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			

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<u>EPINEPHRINE - AUVI-Q</u>						
N 201739 002	10314977	Nov 23, 2024	DP			
	10335549	Apr 30, 2025	DP			
	10688244	Dec 21, 2037	DP	U-2980		
	10737028	Nov 23, 2024	DP			
	10842938	Dec 21, 2037	DP	U-2980		
	10960155	Jun 25, 2026	DP			
	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	10314977	Nov 23, 2024	DP			
	10335549	Apr 30, 2025	DP			
	10688244	Dec 21, 2037	DP	U-2980		
	10737028	Nov 23, 2024	DP			
	10842938	Dec 21, 2037	DP	U-2980		
	10960155	Jun 25, 2026	DP			
	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			
	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		

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<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		
<u>EPOPROSTENOL SODIUM - VELETRI</u>						
N 022260	001	8318802	Mar 15, 2027	DP		
		8598227	Feb 02, 2027			
<u>EPOPROSTENOL SODIUM - VELETRI</u>						
N 022260	002	8318802	Mar 15, 2027	DP		
		8598227	Feb 02, 2027			
<u>ERAVACYCLINE DIHYDROCHLORIDE - XERAVA</u>						
N 211109	001	8796245	Aug 07, 2029	U-2380	NCE	Aug 27, 2023
		8906887	Dec 28, 2030	DP	GAIN	Aug 27, 2028
<u>ERAVACYCLINE DIHYDROCHLORIDE - XERAVA</u>						
N 211109	002	8796245	Aug 07, 2029	U-2380	NCE	Aug 27, 2023
		8906887	Dec 28, 2030	DP	GAIN	Aug 27, 2028
<u>ERDAFITINIB - BALVERSA</u>						
N 212018	001	10898482	Feb 09, 2036	DP U-2518	NCE	Apr 12, 2024
		10898482	Feb 09, 2036	DP U-3065		
		10898482	Feb 09, 2036	DP U-3066		
		10898482	Feb 09, 2036	DP U-3067		
		11077106	Feb 02, 2038	U-3196		
		8895601	May 22, 2031	DS DP		
		9464071	Apr 28, 2031	U-2518		
		9902714	Mar 26, 2035	DP		
<u>ERDAFITINIB - BALVERSA</u>						
N 212018	002	10898482	Feb 09, 2036	DP U-2518	NCE	Apr 12, 2024
		10898482	Feb 09, 2036	DP U-3065		
		10898482	Feb 09, 2036	DP U-3066		
		10898482	Feb 09, 2036	DP U-3067		
		11077106	Feb 02, 2038	U-3196		
		8895601	May 22, 2031	DS DP		
		9464071	Apr 28, 2031	U-2518		
		9902714	Mar 26, 2035	DP		
<u>ERDAFITINIB - BALVERSA</u>						
N 212018	003	10898482	Feb 09, 2036	DP U-2518	NCE	Apr 12, 2024
		10898482	Feb 09, 2036	DP U-3065		
		10898482	Feb 09, 2036	DP U-3066		
		10898482	Feb 09, 2036	DP U-3067		
		11077106	Feb 02, 2038	U-3196		
		8895601	May 22, 2031	DS DP		
		9464071	Apr 28, 2031	U-2518		
		9902714	Mar 26, 2035	DP		
<u>ERIBULIN MESYLATE - HALAVEN</u>						
N 201532	001	6214865	Jul 20, 2023	DS	ODE-107	Jan 28, 2023
		RE46965	Jan 08, 2027	DP		
<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

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<u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUOMET</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 - SEGLUOMET</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 - SEGLUOMET</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 - SEGLUOMET</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
	6699871*PED	Jan 26, 2023				
	7326708	Nov 24, 2026	DS DP U-2214			
	7326708*PED	May 24, 2027				
	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
	6699871*PED	Jan 26, 2023				
	7326708	Nov 24, 2026	DS DP U-2214			
	7326708*PED	May 24, 2027				
	8080580	Jul 13, 2030	DS DP U-2214			
	9308204	Oct 21, 2030	DP			
	9439901	Oct 21, 2030	U-2214			
<u>ERYTHROMYCIN - ERYTHROMYCIN</u>						
A 211975 001					CGT	Apr 02, 2022
<u>ERYTHROMYCIN - ERYTHROMYCIN</u>						
A 211975 002					CGT	Apr 02, 2022
<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>ESKETAMINE HYDROCHLORIDE - SPRAVATO</u>						
N 211243 001	10869844	Sep 10, 2035	U-3034		I-840	Jul 31, 2023
	10869844	Sep 10, 2035	U-3035		NCE*	Mar 05, 2024
	10869844	Sep 10, 2035	U-3036			
	11173134	Sep 14, 2035	U-3257			
	8785500	Jul 09, 2031	U-2502			
	9592207	Mar 20, 2027	U-2502			
<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416 001	10675287	May 06, 2025	U-2041			
	10675287	May 06, 2025	U-2831			
	10695354	May 06, 2025	U-2501			
	10695354	May 06, 2025	U-2831			
	10702536	May 06, 2025	U-2501			
	10912781	Oct 23, 2028	DP			

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<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416	001	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	10675287	May 06, 2025		U-2041	
		10675287	May 06, 2025		U-2831	
		10695354	May 06, 2025		U-2501	
		10695354	May 06, 2025		U-2831	
		10702536	May 06, 2025		U-2501	
		10912781	Oct 23, 2028	DP		
		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	10675287	May 06, 2025		U-2041	
		10675287	May 06, 2025		U-2831	
		10695354	May 06, 2025		U-2501	
		10695354	May 06, 2025		U-2831	
		10702536	May 06, 2025		U-2501	
		10912781	Oct 23, 2028	DP		
		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	004	10675287	May 06, 2025		U-2041	
		10675287	May 06, 2025		U-2831	
		10695354	May 06, 2025		U-2501	
		10695354	May 06, 2025		U-2831	
		10702536	May 06, 2025		U-2501	
		10912781	Oct 23, 2028	DP		
		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 - ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER</u>						
N 205703	001	8829054	Mar 15, 2033	DP		
		8835505	Mar 15, 2033	DP		
<u>ESMOLOL HYDROCHLORIDE - ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER</u>						
N 205703	002	8829054	Mar 15, 2033	DP		
		8835505	Mar 15, 2033	DP		
<u>ESOMEPRAZOLE MAGNESIUM - ESOMEPRAZOLE MAGNESIUM</u>						
N 214278	001	10076494	Dec 08, 2036	DP		
		10835488	Dec 08, 2036	DP		
<u>ESOMEPRAZOLE MAGNESIUM; NAPROXEN - VIMOVO</u>						
N 022511	001	6926907	Feb 28, 2023	DP	U-1052	
		8557285	May 31, 2022	DP		
		8852636	May 31, 2022	DP	U-1052	
		8858996	May 31, 2022	DP	U-1052	

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<u>ESOMEPRAZOLE MAGNESIUM; NAPROXEN - VIMOVO</u>						
N 022511	001	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
		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		
<u>ESTRADIOL - EVAMIST</u>						
N 022014	001	6978945	Jul 31, 2022	DP		
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	001	8231906	Jul 04, 2030	DS DP		
		9730900	Jul 10, 2028			U-2086
		9833419	Jul 10, 2028	DP		
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	002	8231906	Jul 04, 2030	DS DP		
		9730900	Jul 10, 2028			U-2086
		9833419	Jul 10, 2028	DP		
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	003	8231906	Jul 04, 2030	DS DP		
		9730900	Jul 10, 2028			U-2086
		9833419	Jul 10, 2028	DP		
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	004	8231906	Jul 04, 2030	DS DP		
		9730900	Jul 10, 2028			U-2086
		9833419	Jul 10, 2028	DP		
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	005	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	10258630	Dec 20, 2033			U-2316
		10258630	Dec 20, 2033			U-2317
		10398708	Dec 20, 2033			U-2317
		10398708	Dec 20, 2033			U-2614
		10471072	Jun 18, 2033			U-2316
		10471072	Jun 18, 2033			U-2317
		10537581	Nov 21, 2032	DP		U-2316
		10537581	Nov 21, 2032	DP		U-2317
		10568891	Jun 18, 2033			U-2316
		10568891	Jun 18, 2033			U-2317
		10668082	Jun 18, 2033			U-2316
		10668082	Jun 18, 2033			U-2317
		10806697	Nov 21, 2032	DP		
		10835487	Nov 21, 2032			U-2316

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<u>ESTRADIOL - IMVEXXY</u>						
N 208564	001	10835487	Nov 21, 2032		U-2317	
		10888516	Jun 18, 2033		U-2316	
		10888516	Jun 18, 2033		U-2317	
		11065197	Jun 18, 2033	DP		
		11116717	Jun 18, 2033	DP		
		11123283	Jun 18, 2033	DP		
		9180091	Dec 20, 2033	DP	U-2316	
		9180091	Dec 20, 2033	DP	U-2317	
		9289382	Nov 21, 2032	DP		
<u>ESTRADIOL - IMVEXXY</u>						
N 208564	002	10258630	Dec 20, 2033		U-2316	
		10258630	Dec 20, 2033		U-2317	
		10398708	Dec 20, 2033		U-2317	
		10398708	Dec 20, 2033		U-2614	
		10471072	Jun 18, 2033		U-2316	
		10471072	Jun 18, 2033		U-2317	
		10537581	Nov 21, 2032	DP	U-2316	
		10537581	Nov 21, 2032	DP	U-2317	
		10568891	Jun 18, 2033		U-2316	
		10568891	Jun 18, 2033		U-2317	
		10668082	Jun 18, 2033		U-2316	
		10668082	Jun 18, 2033		U-2317	
		10806697	Nov 21, 2032	DP		
		10835487	Nov 21, 2032		U-2316	
		10835487	Nov 21, 2032		U-2317	
		10888516	Jun 18, 2033		U-2316	
		10888516	Jun 18, 2033		U-2317	
		11065197	Jun 18, 2033	DP		
		11116717	Jun 18, 2033	DP		
		11123283	Jun 18, 2033	DP		
		9180091	Dec 20, 2033	DP	U-2316	
		9180091	Dec 20, 2033	DP	U-2317	
		9289382	Nov 21, 2032	DP		
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N 021633	001	7572779	Oct 02, 2025		U-904	
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N 021633	002	7572779	Oct 02, 2025		U-904	
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N 021633	003	7572779	Oct 02, 2025		U-904	
<u>ESTRADIOL; NORETHINDRONE ACETATE; RELUGOLIX - MYFEMBREE</u>						
N 214846	001	11033551	Sep 29, 2037		U-3129	NCE
		7300935	Jan 28, 2024	DS		NP
		8058280	Jan 28, 2024	DS DP		
		9346822	Feb 17, 2024		U-3129	
<u>ESTRADIOL; PROGESTERONE - BIJUVA</u>						
N 210132	001	10052386	Nov 21, 2032	DP		
		10206932	Nov 21, 2032		U-2439	
		10639375	Nov 21, 2032	DP		
		10675288	Nov 21, 2032		U-2439	
		10806740	Nov 21, 2032	DP	U-2439	
		11033626	Nov 21, 2032	DP	U-2439	
		11103513	Nov 21, 2032		U-2439	
		11103516	Nov 21, 2032	DP		
		11110099	Nov 21, 2032	DP		
		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	

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<u>ESTRADIOL; PROGESTERONE - BIJUVA</u>						
N 210132	001	10052386	Nov 21, 2032	DP		
		10206932	Nov 21, 2032	U-2439		
		10639375	Nov 21, 2032	DP		
		10675288	Nov 21, 2032	U-2439		
		10806740	Nov 21, 2032	DP U-2439		
		11033626	Nov 21, 2032	DP U-2439		
		11103513	Nov 21, 2032	U-2439		
		11103516	Nov 21, 2032	DP		
		11110099	Nov 21, 2032	DP		
		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>ETELCALCETIDE - PARSABIV</u>						
N 208325	001	10344765	Jun 27, 2034	DP	NCE	Feb 07, 2022
		11162500	Jun 27, 2034	DP		
		8377880	Jul 29, 2030	DS DP		
		8999932	Feb 07, 2031	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	10344765	Jun 27, 2034	DP	NCE	Feb 07, 2022
		11162500	Jun 27, 2034	DP		
		8377880	Jul 29, 2030	DS DP		
		8999932	Feb 07, 2031	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	10344765	Jun 27, 2034	DP	NCE	Feb 07, 2022
		11162500	Jun 27, 2034	DP		
		8377880	Jul 29, 2030	DS DP		
		8999932	Feb 07, 2031	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	10337003	Mar 14, 2034	U-1918	ODE-122	Sep 19, 2023
		10364431	Mar 14, 2034	U-1918		
		10364431	Mar 14, 2034	U-1919		
		10781451	Jun 28, 2025	DS DP		
		9018368	Jun 28, 2025	DS DP		
		9243245	Oct 27, 2028	DS U-2097		
		9243245	Oct 27, 2028	DS U-2098		
		9506058	Mar 14, 2034	U-1918		
		9506058	Mar 14, 2034	U-1919		
		RE47751	Jun 28, 2025	U-1918		
		RE47751	Jun 28, 2025	U-2664		
		RE47751	Jun 28, 2025	U-2673		
		RE47751	Jun 28, 2025	U-2674		
		RE47769	Feb 02, 2029	DP		
		RE48468	Oct 27, 2028	U-2097		
<u>ETEPLIRSEN - EXONDYS 51</u>						
N 206488	002	10337003	Mar 14, 2034	U-1918	ODE-122	Sep 19, 2023
		10364431	Mar 14, 2034	U-1918		
		10364431	Mar 14, 2034	U-1919		
		10781451	Jun 28, 2025	DS DP		
		9018368	Jun 28, 2025	DS DP		

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<u>ETEPLIRSEN - EXONDYS 51</u>						
N 206488	002	9243245	Oct 27, 2028	DS	U-2097	
		9243245	Oct 27, 2028	DS	U-2098	
		9506058	Mar 14, 2034		U-1918	
		9506058	Mar 14, 2034		U-1919	
		RE47751	Jun 28, 2025		U-1918	
		RE47751	Jun 28, 2025		U-2664	
		RE47751	Jun 28, 2025		U-2673	
		RE47751	Jun 28, 2025		U-2674	
		RE47769	Feb 02, 2029	DP		
		RE48468	Oct 27, 2028		U-2097	
<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 - 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 - TWIRLA</u>						
N 204017	001	8246978	Aug 26, 2028	DP		NP Feb 14, 2023
		8747888	Jul 10, 2028	DP		
		9050348	Jul 10, 2028	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	7838042	Jun 01, 2027	DS	U-3251	
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - TYBLUME</u>						
N 209405	001					NDF Mar 30, 2023
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LO LOESTRIN FE</u>						
N 022501	001	7704984	Feb 02, 2029		U-1090	
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LO MINASTRIN FE</u>						
N 204654	001	7704984	Feb 02, 2029		U-1	
<u>ETHINYL ESTRADIOL; SEGESTERONE ACETATE - ANNOVERA</u>						
N 209627	001	10632066	Feb 01, 2039		U-2786	NCE Aug 10, 2023
		10632066	Feb 01, 2039		U-2787	
		10765628	Feb 01, 2039		U-2786	
		10765628	Feb 01, 2039		U-2787	
		10780047	Feb 01, 2039		U-2786	
		10780047	Feb 01, 2039		U-2787	
		10918649	Jun 21, 2039	DP		
		10925882	Jun 21, 2039	DP		
		10940157	Jun 21, 2039	DP		
<u>ETONOGESTREL - IMPLANON</u>						
N 021529	001	9757552	Jul 28, 2030	DP	U-1	
<u>ETONOGESTREL - NEXPLANON</u>						
N 021529	002	10821277	May 31, 2027	DP		
		8722037	Sep 28, 2027	DP		
		8888745	Aug 28, 2026	DP		
		9757552	Jul 28, 2030	DP	U-1	
<u>EVEROLIMUS - AFINITOR</u>						
N 022334	001	8410131	Nov 01, 2025		U-1368	ODE-108 Feb 26, 2023
		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			

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<u>EVEROLIMUS - AFINITOR</u>						
N 022334	001	9006224	Jul 01, 2028	U-1681		
<u>EVEROLIMUS - AFINITOR</u>						
N 022334	002	8410131	Nov 01, 2025	U-1368	ODE-108	Feb 26, 2023
		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	8410131	Nov 01, 2025	U-1368	ODE-108	Feb 26, 2023
		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	004	8410131	Nov 01, 2025	U-1368	ODE-108	Feb 26, 2023
		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 DISPERZ</u>						
N 203985	001	8617598	Sep 27, 2022	DP	ODE-169	Apr 10, 2025
		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	8617598	Sep 27, 2022	DP	ODE-169	Apr 10, 2025
		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	8617598	Sep 27, 2022	DP	ODE-169	Apr 10, 2025
		8617598*PED	Mar 27, 2023			
		8778962	Feb 18, 2022	U-1541		
		8778962	Feb 18, 2022	U-2280		
		8778962*PED	Aug 18, 2022			
<u>EXENATIDE SYNTHETIC - BYETTA</u>						
N 021773	001				M-232	Nov 04, 2024
<u>EXENATIDE SYNTHETIC - BYETTA</u>						
N 021773	002				M-232	Nov 04, 2024
<u>EXENATIDE SYNTHETIC - BYDUREON</u>						
N 022200	001	6515117	Oct 04, 2025	DS DP U-2588	M-240	Feb 15, 2022
		6515117*PED	Apr 04, 2026		NPP	Jul 22, 2024
		6824822	Oct 09, 2022	DP	PED	Aug 15, 2023
		7456254	Jun 30, 2025	DP U-2588	PED	Jan 22, 2025
		7456254	Jun 30, 2025	DP U-2589		
		7456254	Jun 30, 2025	DP U-2590		
		7456254	Jun 30, 2025	DP U-3188		
		7456254	Jun 30, 2025	DP U-3189		
		7456254	Jun 30, 2025	DP U-3190		
		7456254*PED	Dec 30, 2025			
		7563871	Apr 15, 2024	DP		
		7612176	Apr 13, 2025	DP U-2588		
		7612176	Apr 13, 2025	DP U-2589		

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<u>EXENATIDE SYNTHETIC - BYDUREON</u>						
N 022200 001	7612176	Apr 13, 2025	DP U-2590			
	7612176	Apr 13, 2025	DP U-3188			
	7612176	Apr 13, 2025	DP U-3189			
	7612176	Apr 13, 2025	DP U-3190			
	7612176*PED	Oct 13, 2025				
	8329648	Aug 18, 2026	U-2588			
	8329648	Aug 18, 2026	U-2589			
	8329648	Aug 18, 2026	U-2590			
	8329648	Aug 18, 2026	U-2593			
	8329648	Aug 18, 2026	U-2594			
	8329648	Aug 18, 2026	U-2595			
	8329648	Aug 18, 2026	U-2596			
	8329648	Aug 18, 2026	U-3188			
	8329648	Aug 18, 2026	U-3189			
	8329648	Aug 18, 2026	U-3190			
	8329648*PED	Feb 18, 2027				
	8361972	Mar 21, 2028	U-2588			
	8361972*PED	Sep 21, 2028				
	8431685	Apr 13, 2025	DP U-2588			
	8431685	Apr 13, 2025	DP U-2589			
	8431685	Apr 13, 2025	DP U-2590			
	8431685	Apr 13, 2025	DP U-3188			
	8431685	Apr 13, 2025	DP U-3189			
	8431685	Apr 13, 2025	DP U-3190			
	8431685*PED	Oct 13, 2025				
	8461105	Apr 13, 2025	DP U-2588			
	8461105	Apr 13, 2025	DP U-2589			
	8461105	Apr 13, 2025	DP U-2590			
	8461105	Apr 13, 2025	DP U-3188			
	8461105	Apr 13, 2025	DP U-3189			
	8461105	Apr 13, 2025	DP U-3190			
	8461105*PED	Oct 13, 2025				
	8501698	Jun 20, 2027	DP U-2588			
	8501698*PED	Dec 20, 2027				
	8906851	Aug 18, 2026	U-2588			
	8906851	Aug 18, 2026	U-2589			
	8906851	Aug 18, 2026	U-2590			
	8906851	Aug 18, 2026	U-2593			
	8906851	Aug 18, 2026	U-3188			
	8906851	Aug 18, 2026	U-3189			
	8906851	Aug 18, 2026	U-3190			
	8906851*PED	Feb 18, 2027				
	9238076	Apr 15, 2024	DP U-2588			
	9238076	Apr 15, 2024	DP U-2589			
	9238076	Apr 15, 2024	DP U-2590			
	9238076	Apr 15, 2024	DP U-2599			
	9238076	Apr 15, 2024	DP U-3188			
	9238076	Apr 15, 2024	DP U-3189			
	9238076	Apr 15, 2024	DP U-3190			
	9238076*PED	Oct 15, 2024				
	9884092	Aug 18, 2026	U-2588			
	9884092	Aug 18, 2026	U-2589			
	9884092	Aug 18, 2026	U-2590			
	9884092	Aug 18, 2026	U-2593			
	9884092	Aug 18, 2026	U-2594			
	9884092	Aug 18, 2026	U-2595			
	9884092	Aug 18, 2026	U-2596			
	9884092	Aug 18, 2026	U-3188			
	9884092	Aug 18, 2026	U-3189			
	9884092	Aug 18, 2026	U-3190			
	9884092*PED	Feb 18, 2027				
<u>EXENATIDE SYNTHETIC - BYDUREON PEN</u>						
N 022200 002	6515117	Oct 04, 2025	DS DP U-2588		M-240	Feb 15, 2022
	6515117*PED	Apr 04, 2026			NPP	Jul 22, 2024
	6824822	Oct 09, 2022	DP		PED	Aug 15, 2023
	6824822*PED	Apr 09, 2023			PED	Jan 22, 2025
	7223440*PED	Mar 03, 2022				
	7456254	Jun 30, 2025	DP U-2588			

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<u>EXENATIDE SYNTHETIC - BYDUREON PEN</u>						
N 022200 002	7456254	Jun 30, 2025	DP U-2589			
	7456254	Jun 30, 2025	DP U-2590			
	7456254	Jun 30, 2025	DP U-3188			
	7456254	Jun 30, 2025	DP U-3189			
	7456254	Jun 30, 2025	DP U-3190			
	7456254*PED	Dec 30, 2025				
	7563871	Apr 15, 2024	DP			
	7563871*PED	Oct 15, 2024				
	7612176	Apr 13, 2025	DP U-2588			
	7612176	Apr 13, 2025	DP U-2589			
	7612176	Apr 13, 2025	DP U-2590			
	7612176	Apr 13, 2025	DP U-3188			
	7612176	Apr 13, 2025	DP U-3189			
	7612176	Apr 13, 2025	DP U-3190			
	7612176*PED	Oct 13, 2025				
	8216180	Jan 12, 2028	DP			
	8216180*PED	Jul 12, 2028				
	8329648	Aug 18, 2026	U-2588			
	8329648	Aug 18, 2026	U-2589			
	8329648	Aug 18, 2026	U-2590			
	8329648	Aug 18, 2026	U-2593			
	8329648	Aug 18, 2026	U-2594			
	8329648	Aug 18, 2026	U-2595			
	8329648	Aug 18, 2026	U-2596			
	8329648	Aug 18, 2026	U-3188			
	8329648	Aug 18, 2026	U-3189			
	8329648	Aug 18, 2026	U-3190			
	8329648*PED	Feb 18, 2027				
	8361972	Mar 21, 2028	U-2588			
	8361972*PED	Sep 21, 2028				
	8431685	Apr 13, 2025	DP U-2588			
	8431685	Apr 13, 2025	DP U-2589			
	8431685	Apr 13, 2025	DP U-2590			
	8431685	Apr 13, 2025	DP U-3188			
	8431685	Apr 13, 2025	DP U-3189			
	8431685	Apr 13, 2025	DP U-3190			
	8431685*PED	Oct 13, 2025				
	8439864	Mar 25, 2028	DP			
	8439864*PED	Sep 25, 2028				
	8461105	Apr 13, 2025	DP U-2588			
	8461105	Apr 13, 2025	DP U-2589			
	8461105	Apr 13, 2025	DP U-2590			
	8461105	Apr 13, 2025	DP U-3188			
	8461105	Apr 13, 2025	DP U-3189			
	8461105	Apr 13, 2025	DP U-3190			
	8461105*PED	Oct 13, 2025				
	8501698	Jun 20, 2027	DP U-2588			
	8501698*PED	Dec 20, 2027				
	8690837	May 19, 2029	DP			
	8690837*PED	Nov 19, 2029				
	8721615	Jan 18, 2030	DP			
	8721615*PED	Jul 18, 2030				
	8758292	Nov 12, 2027	DP			
	8758292*PED	May 12, 2028				
	8827963	Feb 04, 2029	DP			
	8827963*PED	Aug 04, 2029				
	8906851	Aug 18, 2026	U-2588			
	8906851	Aug 18, 2026	U-2589			
	8906851	Aug 18, 2026	U-2590			
	8906851	Aug 18, 2026	U-2593			
	8906851	Aug 18, 2026	U-3188			
	8906851	Aug 18, 2026	U-3189			
	8906851	Aug 18, 2026	U-3190			
	8906851*PED	Feb 18, 2027				
	8998876	Jan 07, 2030	DP			
	8998876*PED	Jul 07, 2030				
	9238076	Apr 15, 2024	DP U-2588			
	9238076	Apr 15, 2024	DP U-2589			
	9238076	Apr 15, 2024	DP U-2590			

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<u>EXENATIDE SYNTHETIC - BYDUREON PEN</u>						
N 022200 002	9238076	Apr 15, 2024	DP U-2599			
	9238076	Apr 15, 2024	DP U-3188			
	9238076	Apr 15, 2024	DP U-3189			
	9238076	Apr 15, 2024	DP U-3190			
	9238076*PED	Oct 15, 2024				
	9320853	Mar 25, 2028	DP			
	9320853*PED	Sep 25, 2028				
	9884092	Aug 18, 2026	U-2588			
	9884092	Aug 18, 2026	U-2589			
	9884092	Aug 18, 2026	U-2590			
	9884092	Aug 18, 2026	U-2593			
	9884092	Aug 18, 2026	U-2594			
	9884092	Aug 18, 2026	U-2595			
	9884092	Aug 18, 2026	U-2596			
	9884092	Aug 18, 2026	U-3188			
	9884092	Aug 18, 2026	U-3189			
	9884092	Aug 18, 2026	U-3190			
	9884092*PED	Feb 18, 2027				
<u>EXENATIDE SYNTHETIC - BYDUREON BCISE</u>						
N 209210 001	6515117	Oct 04, 2025	DS DP U-2588		M-240	Feb 15, 2022
	6515117*PED	Apr 04, 2026			NPP	Jul 22, 2024
	6824822	Oct 09, 2022	DP		PED	Aug 15, 2023
	6824822*PED	Apr 09, 2023			PED	Jan 22, 2025
	7223440*PED	Mar 03, 2022				
	7456254	Jun 30, 2025	DP U-2588			
	7456254	Jun 30, 2025	DP U-2589			
	7456254	Jun 30, 2025	DP U-2590			
	7456254	Jun 30, 2025	DP U-3188			
	7456254	Jun 30, 2025	DP U-3189			
	7456254	Jun 30, 2025	DP U-3190			
	7456254*PED	Dec 30, 2025				
	7563871	Apr 15, 2024	DP			
	7563871*PED	Oct 15, 2024				
	7612176	Apr 13, 2025	DP U-2589			
	7612176	Apr 13, 2025	DP U-2590			
	7612176	Apr 13, 2025	DP U-3188			
	7612176	Apr 13, 2025	DP U-3189			
	7612176	Apr 13, 2025	DP U-3190			
	7612176*PED	Oct 13, 2025				
	8329648	Aug 18, 2026	U-2588			
	8329648	Aug 18, 2026	U-2589			
	8329648	Aug 18, 2026	U-2590			
	8329648	Aug 18, 2026	U-2593			
	8329648	Aug 18, 2026	U-2594			
	8329648	Aug 18, 2026	U-2595			
	8329648	Aug 18, 2026	U-2596			
	8329648	Aug 18, 2026	U-3188			
	8329648	Aug 18, 2026	U-3189			
	8329648	Aug 18, 2026	U-3190			
	8329648*PED	Feb 18, 2027				
	8361972	Mar 21, 2028	U-2588			
	8361972*PED	Sep 21, 2028				
	8431685	Apr 13, 2025	DP U-2588			
	8431685	Apr 13, 2025	DP U-2589			
	8431685	Apr 13, 2025	DP U-2590			
	8431685	Apr 13, 2025	DP U-2597			
	8431685	Apr 13, 2025	DP U-3188			
	8431685	Apr 13, 2025	DP U-3189			
	8431685	Apr 13, 2025	DP U-3190			
	8431685*PED	Oct 13, 2025				
	8461105	Apr 13, 2025	DP U-2588			
	8461105	Apr 13, 2025	DP U-2589			
	8461105	Apr 13, 2025	DP U-2590			
	8461105	Apr 13, 2025	DP U-2597			
	8461105	Apr 13, 2025	DP U-3188			
	8461105	Apr 13, 2025	DP U-3189			
	8461105	Apr 13, 2025	DP U-3190			
	8461105*PED	Oct 13, 2025				

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<u>EXENATIDE SYNTHETIC - BYDUREON BCISE</u>						
N 209210	001	8501698	Jun 20, 2027			U-2588
		8501698*PED	Dec 20, 2027			
		8895033	Oct 04, 2030	DP		U-2589
		8895033	Oct 04, 2030	DP		U-2590
		8895033	Oct 04, 2030	DP		U-2597
		8895033	Oct 04, 2030	DP		U-2601
		8895033	Oct 04, 2030	DP		U-2602
		8895033	Oct 04, 2030	DP		U-3188
		8895033	Oct 04, 2030	DP		U-3189
		8895033	Oct 04, 2030	DP		U-3190
		8895033*PED	Apr 04, 2031			
		8906851	Aug 18, 2026			U-2588
		8906851	Aug 18, 2026			U-2589
		8906851	Aug 18, 2026			U-2590
		8906851	Aug 18, 2026			U-2593
		8906851	Aug 18, 2026			U-2597
		8906851	Aug 18, 2026			U-3188
		8906851	Aug 18, 2026			U-3189
		8906851	Aug 18, 2026			U-3190
		8906851*PED	Feb 18, 2027			
		9238076	Apr 15, 2024	DP		U-2588
		9238076	Apr 15, 2024	DP		U-2589
		9238076	Apr 15, 2024	DP		U-2590
		9238076	Apr 15, 2024	DP		U-2597
		9238076	Apr 15, 2024	DP		U-2599
		9238076	Apr 15, 2024	DP		U-3188
		9238076	Apr 15, 2024	DP		U-3189
		9238076	Apr 15, 2024	DP		U-3190
		9238076*PED	Oct 15, 2024			
		9884092	Aug 18, 2026			U-2588
		9884092	Aug 18, 2026			U-2589
		9884092	Aug 18, 2026			U-2590
		9884092	Aug 18, 2026			U-2593
		9884092	Aug 18, 2026			U-2594
		9884092	Aug 18, 2026			U-2595
		9884092	Aug 18, 2026			U-2596
		9884092	Aug 18, 2026			U-2597
		9884092	Aug 18, 2026			U-3188
		9884092	Aug 18, 2026			U-3189
		9884092	Aug 18, 2026			U-3190
		9884092*PED	Feb 18, 2027			
<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>EZETIMIBE; ROSUVASTATIN CALCIUM - ROSZET</u>						
N 213072	001	10376470	May 01, 2033	DP		U-3095
		9763885	May 01, 2033	DP		U-3095
<u>EZETIMIBE; ROSUVASTATIN CALCIUM - ROSZET</u>						
N 213072	002	10376470	May 01, 2033	DP		U-3095
		9763885	May 01, 2033	DP		U-3095
<u>EZETIMIBE; ROSUVASTATIN CALCIUM - ROSZET</u>						
N 213072	003	10376470	May 01, 2033	DP		U-3095
		9763885	May 01, 2033	DP		U-3095
<u>EZETIMIBE; ROSUVASTATIN CALCIUM - ROSZET</u>						
N 213072	004	10376470	May 01, 2033	DP		U-3095
		9763885	May 01, 2033	DP		U-3095
<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		

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<u>FAMOTIDINE; IBUPROFEN - DUEXIS</u>						
N 022519	001 8501228	Jul 18, 2026	U-1196			
<u>FEBUXOSTAT - ULORIC</u>						
N 021856	001 7361676	Mar 08, 2024	DP			
	8372872	Sep 08, 2031	U-1346			
	9107912	Sep 08, 2031	U-1346			
<u>FEBUXOSTAT - ULORIC</u>						
N 021856	002 7361676	Mar 08, 2024	DP			
	8372872	Sep 08, 2031	U-1346			
	9107912	Sep 08, 2031	U-1346			
<u>FEDRATINIB HYDROCHLORIDE - INREBIC</u>						
N 212327	001 10391094	Jun 04, 2032	DP U-2607		NCE	Aug 16, 2024
	7528143	Dec 16, 2026	DS		ODE-259	Aug 16, 2026
	7825246	Dec 16, 2026	DS			
	8138199	Jun 30, 2028	U-2607			
<u>FENFLURAMINE HYDROCHLORIDE - FINTEPLA</u>						
N 212102	001 10452815	Jun 29, 2038	U-2859		NP	Jun 25, 2023
	10478441	May 03, 2033	U-2860		ODE-312	Jun 25, 2027
	10478442	May 03, 2033	U-2860			
	10603290	Aug 02, 2037	U-2861			
	10947183	Dec 20, 2036	DS DP			
	10950331	Sep 28, 2035	U-3098			
	11040018	Aug 02, 2037	U-2861			
	9549909	May 03, 2033	U-2858			
	9603814	May 03, 2033	U-2858			
	9603815	May 03, 2033	U-2858			
	9610260	May 03, 2033	U-2858			
<u>FENOFIBRATE - TRICOR</u>						
N 021656	001 7276249	Feb 21, 2023	DP			
	7320802	Feb 21, 2023	U-847			
<u>FENOFIBRATE - TRICOR</u>						
N 021656	002 7276249	Feb 21, 2023	DP			
	7320802	Feb 21, 2023	U-847			
<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			

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<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
		10610523	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
		10610523	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
		10610523	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
		10610523	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
		10610523	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
		10610523	Jan 25, 2027			DP
		8486972	Apr 27, 2030			DP

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<u>FENTANYL - SUBSYS</u>						
N 202788	006	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		
		10610523	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	7862832	Jun 15, 2028	DP		
		7862833	Jun 15, 2028	DP		
		8092832	Dec 30, 2024	DP		
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	002	7862832	Jun 15, 2028	DP		
		7862833	Jun 15, 2028	DP		
		8092832	Dec 30, 2024	DP		
		8119158	Dec 30, 2024	DP		
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	003	7862832	Jun 15, 2028	DP		
		7862833	Jun 15, 2028	DP		
		8092832	Dec 30, 2024	DP		
		8119158	Dec 30, 2024	DP		
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	004	7862832	Jun 15, 2028	DP		
		7862833	Jun 15, 2028	DP		
		8092832	Dec 30, 2024	DP		
		8119158	Dec 30, 2024	DP		
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	005	7862832	Jun 15, 2028	DP		
		7862833	Jun 15, 2028	DP		
		8092832	Dec 30, 2024	DP		
		8119158	Dec 30, 2024	DP		
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	001	9597288	Jul 23, 2027	DP	U-767	
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	002	9597288	Jul 23, 2027	DP	U-767	
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	003	9597288	Jul 23, 2027	DP	U-767	
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	004	9597288	Jul 23, 2027	DP	U-767	
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	005	9597288	Jul 23, 2027	DP	U-767	
<u>FENTANYL CITRATE - LAZANDA</u>						
N 022569	001	8216604	Oct 03, 2024		U-767	
		8889176	Jan 16, 2024		U-767	
		9078814	Jan 08, 2024	DP		
		9731869	Jan 26, 2032	DP		
		9814705	Jan 08, 2024	DP		

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<u>FENTANYL CITRATE - LAZANDA</u>						
N 022569	002	8216604	Oct 03, 2024	U-767		
		8889176	Jan 16, 2024	U-767		
		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	DP		
		9814705	Jan 08, 2024	DP		
<u>FENTANYL HYDROCHLORIDE - IONSY</u>						
N 021338	001	6881208	Apr 19, 2022	U-736		
		6975902	Apr 01, 2024	DP		
		8301238	Sep 30, 2031	DP		
		8428708	May 21, 2032	U-736		
		8428709	Jun 11, 2032	DP U-736		
		8781571	Mar 31, 2032	DP U-736		
		9095706	Feb 03, 2033	DP		
		9364656	Sep 30, 2031	U-736		
		9731121	Oct 17, 2031	DP		
<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565	001	11123321	Oct 20, 2023	DS DP U-2555	NPP	Nov 19, 2024
		11123321	Oct 20, 2023	DS DP U-2557		
		7612109	Feb 05, 2024	DS DP		
		7754702	Feb 15, 2028	U-1432		
		7754702	Feb 15, 2028	U-2555		
		7754702	Feb 15, 2028	U-2556		
		7754702	Feb 15, 2028	U-2557		
		8895612	Jan 08, 2027	U-1620		
		8895612	Jan 08, 2027	U-3050		
		8895612	Jan 08, 2027	U-3051		
		8895612	Jan 08, 2027	U-3115		
		8895612	Jan 08, 2027	U-3116		
		9376505	Oct 20, 2023	DS DP		
<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565	002	11123321	Oct 20, 2023	DS DP U-2555	NPP	Nov 19, 2024
		11123321	Oct 20, 2023	DS DP U-2557		
		7612109	Feb 05, 2024	DS DP		
		7754702	Feb 15, 2028	U-2555		
		7754702	Feb 15, 2028	U-2556		
		7754702	Feb 15, 2028	U-2557		
		8895612	Jan 08, 2027	U-1620		
		8895612	Jan 08, 2027	U-3050		
		8895612	Jan 08, 2027	U-3051		
		8895612	Jan 08, 2027	U-3115		
		8895612	Jan 08, 2027	U-3116		
		9376505	Oct 20, 2023	DS DP		
<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565	003	11123321	Oct 20, 2023	DS DP U-2555	NPP	Nov 19, 2024
		11123321	Oct 20, 2023	DS DP U-2557	NS	Apr 28, 2024
		7612109	Feb 05, 2024	DS DP		
		7754702	Feb 15, 2028	U-2555		
		7754702	Feb 15, 2028	U-2556		
		7754702	Feb 15, 2028	U-2557		
		8895612	Jan 08, 2027	U-1620		
		8895612	Jan 08, 2027	U-3050		
		8895612	Jan 08, 2027	U-3051		
		8895612	Jan 08, 2027	U-3115		
		8895612	Jan 08, 2027	U-3116		
		9376505	Oct 20, 2023	DS DP		
<u>FERRIC CITRATE - AURYXIA</u>						
N 205874	001	10300039	Jul 21, 2030	U-2549		
		5753706	Feb 03, 2022	DP U-1577		
		7767851	Feb 18, 2024	DS DP		
		8093423	Apr 21, 2026	U-1577		
		8299298	Feb 18, 2024	DP		

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<u>FERRIC CITRATE - AURYXIA</u>						
N 205874	001	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 DERISOMALTOSE - MONOFERRIC</u>						
N 208171	001	10414831	Mar 25, 2029	DS DP		
		8815301	Aug 14, 2029	DS DP	U-2734	
<u>FERRIC DERISOMALTOSE - MONOFERRIC</u>						
N 208171	002	10414831	Mar 25, 2029	DS DP		
		8815301	Aug 14, 2029	DS DP	U-2734	
<u>FERRIC DERISOMALTOSE - MONOFERRIC</u>						
N 208171	003	10414831	Mar 25, 2029	DS DP		
		8815301	Aug 14, 2029	DS DP	U-2734	
<u>FERRIC MALTOL - ACCRUFER</u>						
N 212320	001	10179120	Jan 06, 2035		U-2603	NCE Jul 25, 2024
		9248148	Mar 29, 2031		U-2603	
		9802973	Oct 23, 2035	DS DP	U-2603	
<u>FERRIC OXYHYDROXIDE - VELPHORO</u>						
N 205109	001	10624855	Nov 26, 2034	DP		
		10624855*PED	May 26, 2035			
		10682376	Nov 13, 2028	DP		
		10682376*PED	May 13, 2029			
		10695367	Nov 13, 2028	DP		
		10695367*PED	May 13, 2029			
		10925896	Nov 13, 2028	DP		
		10925896*PED	May 13, 2029			
		10925897	Nov 13, 2028	DP		
		10925897*PED	May 13, 2029			
		10933090	Nov 13, 2028	DP		
		10933090*PED	May 13, 2029			
		11013761	Nov 13, 2028	DP		
		11013761*PED	May 13, 2029			
		11013762	Nov 13, 2028	DP		
		11013762*PED	May 13, 2029			
		9561251	Jan 23, 2030	DP	U-1468	
		9561251*PED	Jul 23, 2030			
<u>FERRIC PYROPHOSPHATE CITRATE - TRIFERIC</u>						
N 206317	001	7816404	Apr 17, 2029	DP	U-1656	
<u>FERRIC PYROPHOSPHATE CITRATE - TRIFERIC</u>						
N 208551	001	7816404	Apr 17, 2029		U-1656	
		7857977	Sep 08, 2027		U-1656	
<u>FERRIC PYROPHOSPHATE CITRATE - TRIFERIC AVNU</u>						
N 212860	001	7816404	Apr 17, 2029	DS	U-2801	
<u>FERUMOXYTOL - FERAHEME</u>						
N 022180	001	6599498	Jun 30, 2023	DS DP		
<u>FESOTERODINE FUMARATE - TOVIAZ</u>						
N 022030	001	6858650	Jul 03, 2022	DS	U-913	I-861 Jun 17, 2024
		7807715	Jun 07, 2027	DP	U-913	PED Dec 17, 2024
		8088398	Jun 07, 2027	DP	U-913	
		8501723	Jun 07, 2027	DP		
		8501723*PED	Dec 07, 2027			

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<u>FESOTERODINE FUMARATE - TOVIAZ</u>						
N 022030	002	6858650	Jul 03, 2022	DS U-913	I-861	Jun 17, 2024
		7807715	Jun 07, 2027	DP U-913	PED	Dec 17, 2024
		8088398	Jun 07, 2027	DP U-913		
		8501723	Jun 07, 2027	DP		
		8501723*PED	Dec 07, 2027			
<u>FEXINIDAZOLE - FEXINIDAZOLE</u>						
N 214429	001				NCE	Jul 16, 2026
					ODE-359	Jul 16, 2028
<u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA ALLERGY</u>						
N 201373	001	8933097	Aug 02, 2030	DP		
<u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA HIVES</u>						
N 201373	002	8933097	Aug 02, 2030	DP		
<u>FIDAXOMICIN - DIFICID</u>						
N 201699	001	7378508	Jul 31, 2027	DS DP	NPP	Jan 24, 2023
		7863249	Jul 31, 2027	DS DP	ODE-367	Jan 24, 2027
		7906489	Mar 04, 2027		PED	Jul 24, 2023
		7906489	Mar 04, 2027			
		7906489*PED	Sep 04, 2027			
		8586551	Jul 15, 2023	DS DP		
		8586551*PED	Jan 15, 2024			
		8859510	Jul 31, 2027		U-2741	
		8859510	Jul 31, 2027		U-319	
		8859510*PED	Jan 31, 2028			
<u>FIDAXOMICIN - DIFICID</u>						
N 213138	001	7378508	Jul 31, 2027	DS DP	NP	Jan 24, 2023
		7378508*PED	Jan 31, 2028		ODE-367	Jan 24, 2027
		7863249	Jul 31, 2027	DP	PED	Jul 24, 2023
		7863249*PED	Jan 31, 2028			
		7906489	Mar 04, 2027		U-2741	
		7906489*PED	Sep 04, 2027			
		8586551	Jul 23, 2023	DS DP		
		8586551*PED	Jan 23, 2024			
		8859510	Jul 31, 2027		U-2741	
		8859510*PED	Jan 31, 2028			
		9808530	May 28, 2034	DP		
		9808530*PED	Nov 28, 2034			
<u>FINAFLOXACIN - XTORO</u>						
N 206307	001	8536167	Aug 08, 2031		U-1679	
		9119859	Jul 02, 2030		U-1679	
		9504691	Nov 21, 2033	DP	U-1679	
<u>FINERENONE - KERENDIA</u>						
N 215341	001	8436180	Apr 12, 2029	DS DP	NCE	Jul 09, 2026
<u>FINERENONE - KERENDIA</u>						
N 215341	002	8436180	Apr 12, 2029	DS DP	NCE	Jul 09, 2026
<u>FINGOLIMOD HYDROCHLORIDE - GILENYA</u>						
N 022527	001	10543179	Dec 25, 2027		U-2719	
		8324283	Mar 29, 2026	DP		
		8324283*PED	Sep 29, 2026			
		9187405	Jun 25, 2027		U-2613	
		9187405*PED	Dec 25, 2027			
<u>FINGOLIMOD HYDROCHLORIDE - GILENYA</u>						
N 022527	002	9592208	Mar 30, 2032	DP	U-2315	
		9592208*PED	Sep 30, 2032			
<u>FINGOLIMOD LAURYL SULFATE - TASCENSO ODT</u>						
N 214962	001	10555902	Jan 19, 2036		U-3268	
		10925829	Jan 19, 2036	DP		
		9925138	Jan 19, 2036	DP		

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<u>FISH OIL TRIGLYCERIDES - OMEGAVEN</u>						
N 210589	001	10350186	Nov 05, 2024	U-2585	NCE	Jul 27, 2023
		9566260	Jul 11, 2025	DP U-2366	ODE-202	Jul 27, 2025
		9629821	Jul 11, 2025	DP U-2367		
<u>FISH OIL TRIGLYCERIDES - OMEGAVEN</u>						
N 210589	002	10350186	Nov 05, 2024	U-2585	NCE	Jul 27, 2023
		9566260	Jul 11, 2025	DP U-2366	ODE-202	Jul 27, 2025
		9629821	Jul 11, 2025	DP U-2367		
<u>FLIBANSERIN - ADDYI</u>						
N 022526	001	7151103	May 09, 2023	U-1734		
		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		
<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>FLORTAUCIPIR F-18 - TAUVID</u>						
N 212123	001	8932557	May 19, 2029	DS	NCE	May 28, 2025
<u>FLORTAUCIPIR F-18 - TAUVID</u>						
N 212123	002	8932557	May 19, 2029	DS	NCE	May 28, 2025
<u>FLUCICLOVINE F-18 - AXUMIN</u>						
N 208054	001	10010632	Nov 28, 2026	DP		
		10124079	Dec 30, 2035	U-2450		
		10716868	Dec 30, 2035	U-2450		
		10933147	Dec 30, 2035	U-2450		
		10953112	Nov 28, 2026	U-1879		
		10967077	Dec 30, 2035	U-2450		
		9387266	Nov 28, 2026	U-1879		
<u>FLUDARABINE PHOSPHATE - OFORTA</u>						
N 022273	001	7148207	Dec 20, 2022	DP U-944		
<u>FLUOCINOLONE ACETONIDE - ILLUVIEN</u>						
N 201923	001	8871241	Aug 12, 2027	DP		
<u>FLUOCINOLONE ACETONIDE - YUTIQ</u>						
N 210331	001	8871241	Aug 12, 2027	DP		
<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	7220424	Jan 07, 2023	U-861		
		7794738	Sep 11, 2022	U-1084		
		8232264	Mar 09, 2023	DP		
<u>FLUORODOPA F-18 - FLUORODOPA F18</u>						
N 200655	001				NCE	Oct 10, 2024
					W	Oct 10, 2024

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUOROESTRADIOL F-18 - CERIANNA</u>						
N 212155	001				NCE	May 20, 2025
<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	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	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	002	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	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	11090294	Nov 29, 2030		I-843	Sep 09, 2023
		7439393	May 21, 2025	DS DP U-2127		
		7439393	May 21, 2025	DS DP U-2957		
		7488827	Dec 18, 2027	DS DP		
		7498440	Apr 27, 2025	DS DP		
		7776895	Sep 11, 2022	DP		
		8113199	Oct 23, 2027	DP		
		8161968	Feb 05, 2028	DP		
		8183257	Jul 27, 2025		U-2128	
		8183257	Jul 27, 2025		U-2129	
		8309572	Apr 27, 2025		U-2129	
		8511304	Jun 14, 2027	DP U-2954		
		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		
		RE44874	Mar 23, 2023	DS DP U-2955		

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - TRELEGY ELLIPTA</u>						
N 209482	002	7439393	May 21, 2025	DS DP U-2957	NS	Sep 09, 2023
		7488827	Dec 18, 2027	DS DP		
		7498440	Apr 27, 2025	DS DP		
		7776895	Sep 11, 2022	DP		
		8113199	Oct 23, 2027	DP		
		8161968	Feb 05, 2028	DP		
		8183257	Jul 27, 2025	U-2129		
		8309572	Apr 27, 2025	U-2129		
		8511304	Jun 14, 2027	DP U-2954		
		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-2955		
<u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u>						
N 204275	001	11116721	Feb 26, 2029	DP U-1401		
		11116721	Feb 26, 2029	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		
		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		
<u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u>						
N 204275	002	11116721	Feb 26, 2029	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		
		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 - FLOVENT HFA</u>						
N 021433	001	7500444	Feb 26, 2026	DP		
		7500444*PED	Aug 26, 2026			
		7832351	Jun 19, 2023	DP		
<u>FLUTICASONE PROPIONATE - FLOVENT HFA</u>						
N 021433	002	7500444	Feb 26, 2026	DP		
		7500444*PED	Aug 26, 2026			
		7832351	Jun 19, 2023	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		
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798	001	10022510	May 18, 2031	DP	NPP	Jul 09, 2024
		10022510*PED	Nov 18, 2031		PED	Jan 09, 2025
		10124131	May 18, 2031	DP		
		10124131*PED	Nov 18, 2031			
		10195375	Feb 14, 2031	DP		
		10195375*PED	Aug 14, 2031			
		10561808	Jan 01, 2032	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798	001	10561808*PED				
		Jul 01, 2032				
		10765820				
		May 19, 2025	DP			
		10765820*PED				
		Nov 19, 2025				
		7540282				
		May 06, 2023	DP			
		7540282*PED				
		Nov 06, 2023				
		8651103				
		Mar 26, 2028	DP			
		8651103*PED				
		Sep 26, 2028				
		8714149				
		Feb 25, 2032	DP			
		8714149*PED				
		Aug 25, 2032				
		8978966				
		Jan 13, 2032	DP			
		8978966*PED				
		Jul 13, 2032				
		9216260				
		Jun 28, 2031	DP			
		9216260*PED				
		Dec 28, 2031				
		9463288				
		May 19, 2025	DP			
		9463288*PED				
		Nov 19, 2025				
		9616024				
		Sep 01, 2024	DP			
		9616024*PED				
		Mar 01, 2025				
		9731087				
		May 18, 2031	DP			
		9731087*PED				
		Nov 18, 2031				
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798	002	10022510				
		May 18, 2031	DP			
		10022510*PED				
		Nov 18, 2031				
		10124131				
		May 18, 2031	DP			
		10124131*PED				
		Nov 18, 2031				
		10195375				
		Feb 14, 2031	DP			
		10195375*PED				
		Aug 14, 2031				
		10561808				
		Jan 01, 2032	DP			
		10561808*PED				
		Jul 01, 2032				
		10765820				
		May 19, 2025	DP			
		10765820*PED				
		Nov 19, 2025				
		7540282				
		May 06, 2023	DP			
		7540282*PED				
		Nov 06, 2023				
		8651103				
		Mar 26, 2028	DP			
		8651103*PED				
		Sep 26, 2028				
		8714149				
		Feb 25, 2032	DP			
		8714149*PED				
		Aug 25, 2032				
		8978966				
		Jan 13, 2032	DP			
		8978966*PED				
		Jul 13, 2032				
		9216260				
		Jun 28, 2031	DP			
		9216260*PED				
		Dec 28, 2031				
		9463288				
		May 19, 2025	DP			
		9463288*PED				
		Nov 19, 2025				
		9616024				
		Sep 01, 2024	DP			
		9616024*PED				
		Mar 01, 2025				
		9731087				
		May 18, 2031	DP			
		9731087*PED				
		Nov 18, 2031				
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798	003	10022510				
		May 18, 2031	DP			
		10022510*PED				
		Nov 18, 2031				
		10124131				
		May 18, 2031	DP			
		10124131*PED				
		Nov 18, 2031				
		10195375				
		Feb 14, 2031	DP			
		10195375*PED				
		Aug 14, 2031				
		10561808				
		Jan 01, 2032	DP			
		10561808*PED				
		Jul 01, 2032				
		10765820				
		May 19, 2025	DP			
		10765820*PED				
		Nov 19, 2025				
		7540282				
		May 06, 2023	DP			
		7540282*PED				
		Nov 06, 2023				
		8651103				
		Mar 26, 2028	DP			
		8651103*PED				
		Sep 26, 2028				
		8714149				
		Feb 25, 2032	DP			
		8714149*PED				
		Aug 25, 2032				
		8978966				
		Jan 13, 2032	DP			
		8978966*PED				
		Jul 13, 2032				
		9216260				
		Jun 28, 2031	DP			
		9216260*PED				
		Dec 28, 2031				

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<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 003	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 004	10022510	May 18, 2031	DP		NPP	Jul 09, 2024
	10022510*PED	Nov 18, 2031			PED	Jan 09, 2025
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	7540282	May 06, 2023	DP			
	7540282*PED	Nov 06, 2023				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 005	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	7540282	May 06, 2023	DP			
	7540282*PED	Nov 06, 2023				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			

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<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 005	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 006	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	7540282	May 06, 2023	DP			
	7540282*PED	Nov 06, 2023				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 007	10022510	May 18, 2031	DP		NS	Jul 09, 2024
	10022510*PED	Nov 18, 2031			PED	Jan 09, 2025
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	7540282	May 06, 2023	DP			
	7540282*PED	Nov 06, 2023				
	8651103	Mar 26, 2028	DP			

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<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798	007	8651103*PED	Sep 26, 2028			
		8714149	Feb 25, 2032	DP		
		8714149*PED	Aug 25, 2032			
		8978966	Jan 13, 2032	DP		
		8978966*PED	Jul 13, 2032			
		9216260	Jun 28, 2031	DP		
		9216260*PED	Dec 28, 2031			
		9463288	May 19, 2025	DP		
		9463288*PED	Nov 19, 2025			
		9616024	Sep 01, 2024	DP		
		9616024*PED	Mar 01, 2025			
		9731087	May 18, 2031	DP		
		9731087*PED	Nov 18, 2031			
<u>FLUTICASONE PROPIONATE - XHANCE</u>						
N 209022	001	10076614	Oct 20, 2034	DP		
		10076615	Jul 30, 2029	DP	U-2133	
		10124132	Mar 06, 2027	DP	U-2133	
		10179216	Jul 08, 2033	DP	U-2133	
		10252010	Feb 07, 2031	DP		
		10300229	Jul 07, 2035	DP	U-2133	
		10478574	Nov 04, 2033		U-2133	
		11033696	May 20, 2033	DP		
		7975690	Dec 29, 2025		U-2133	
		8327844	Oct 08, 2023		U-2133	
		8522778	May 11, 2022	DP		
		8550073	Oct 22, 2029	DP		
		8978647	Dec 06, 2030	DP		
<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		
<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		
<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		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799	001	10022510	May 18, 2031	DP	M-61	Jul 09, 2024
		10022510*PED	Nov 18, 2031		PED	Jan 09, 2025
		10124131	May 18, 2031	DP		
		10124131*PED	Nov 18, 2031			
		10195375	Feb 14, 2031	DP		
		10195375*PED	Aug 14, 2031			
		10561808	Jan 01, 2032	DP		
		10561808*PED	Jul 01, 2032			
		10765820	May 19, 2025	DP		
		10765820*PED	Nov 19, 2025			
		7540282	May 06, 2023	DP		
		7540282*PED	Nov 06, 2023			
		8651103	Mar 26, 2028	DP		
		8651103*PED	Sep 26, 2028			
		8714149	Feb 25, 2032	DP		
		8714149*PED	Aug 25, 2032			
		8978966	Jan 13, 2032	DP		
		8978966*PED	Jul 13, 2032			
		9066957	Oct 06, 2034	DP	U-645	
		9066957*PED	Apr 06, 2035			
		9216260	Jun 28, 2031	DP		
		9216260*PED	Dec 28, 2031			
		9415008	Oct 06, 2034	DP	U-645	
		9415008*PED	Apr 06, 2035			
		9463288	May 19, 2025	DP		

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<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799	001	9463288*PED	Nov 19, 2025			
		9616024	Sep 01, 2024	DP		
		9616024*PED	Mar 01, 2025			
		9731087	May 18, 2031	DP		
		9731087*PED	Nov 18, 2031			
		9987229	Sep 01, 2024	DP		
		9987229*PED	Mar 01, 2025			
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799	002	10022510	May 18, 2031	DP		
		10022510*PED	Nov 18, 2031			
		10124131	May 18, 2031	DP		
		10124131*PED	Nov 18, 2031			
		10195375	Feb 14, 2031	DP		
		10195375*PED	Aug 14, 2031			
		10561808	Jan 01, 2032	DP		
		10561808*PED	Jul 01, 2032			
		10765820	May 19, 2025	DP		
		10765820*PED	Nov 19, 2025			
		7540282	May 06, 2023	DP		
		7540282*PED	Nov 06, 2023			
		8651103	Mar 26, 2028	DP		
		8651103*PED	Sep 26, 2028			
		8714149	Feb 25, 2032	DP		
		8714149*PED	Aug 25, 2032			
		8978966	Jan 13, 2032	DP		
		8978966*PED	Jul 13, 2032			
		9066957	Oct 06, 2034	DP U-645		
		9066957*PED	Apr 06, 2035			
		9216260	Jun 28, 2031	DP		
		9216260*PED	Dec 28, 2031			
		9463288	May 19, 2025	DP		
		9463288*PED	Nov 19, 2025			
		9616024	Sep 01, 2024	DP		
		9616024*PED	Mar 01, 2025			
		9731087	May 18, 2031	DP		
		9731087*PED	Nov 18, 2031			
		9987229	Sep 01, 2024	DP		
		9987229*PED	Mar 01, 2025			
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799	003	10022510	May 18, 2031	DP		
		10022510*PED	Nov 18, 2031			
		10124131	May 18, 2031	DP		
		10124131*PED	Nov 18, 2031			
		10195375	Feb 14, 2031	DP		
		10195375*PED	Aug 14, 2031			
		10561808	Jan 01, 2032	DP		
		10561808*PED	Jul 01, 2032			
		10765820	May 19, 2025	DP		
		10765820*PED	Nov 19, 2025			
		7540282	May 06, 2023	DP		
		7540282*PED	Nov 06, 2023			
		8651103	Mar 26, 2028	DP		
		8651103*PED	Sep 26, 2028			
		8714149	Feb 25, 2032	DP		
		8714149*PED	Aug 25, 2032			
		8978966	Jan 13, 2032	DP		
		8978966*PED	Jul 13, 2032			
		9066957	Oct 06, 2034	DP U-645		
		9066957*PED	Apr 06, 2035			
		9216260	Jun 28, 2031	DP		
		9216260*PED	Dec 28, 2031			
		9463288	May 19, 2025	DP		
		9463288*PED	Nov 19, 2025			
		9616024	Sep 01, 2024	DP		
		9616024*PED	Mar 01, 2025			
		9731087	May 18, 2031	DP		
		9731087*PED	Nov 18, 2031			

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<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799 003	9987229	Sep 01, 2024	DP			
	9987229*PED	Mar 01, 2025				
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 004	10022510	May 18, 2031	DP		M-61	Jul 09, 2024
	10022510*PED	Nov 18, 2031			PED	Jan 09, 2025
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	7540282	May 06, 2023	DP			
	7540282*PED	Nov 06, 2023				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9066957	Oct 06, 2034	DP U-645			
	9066957*PED	Apr 06, 2035				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9415008	Oct 06, 2034	DP U-645			
	9415008*PED	Apr 06, 2035				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
	9987229	Sep 01, 2024	DP			
	9987229*PED	Mar 01, 2025				
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 005	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	7540282	May 06, 2023	DP			
	7540282*PED	Nov 06, 2023				
	8651103	Mar 26, 2028	DP			

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<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 005	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9066957	Oct 06, 2034	DP U-645			
	9066957*PED	Apr 06, 2035				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
	9987229	Sep 01, 2024	DP			
	9987229*PED	Mar 01, 2025				
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 006	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	7540282	May 06, 2023	DP			
	7540282*PED	Nov 06, 2023				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9066957	Oct 06, 2034	DP U-645			
	9066957*PED	Apr 06, 2035				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
	9987229	Sep 01, 2024	DP			
	9987229*PED	Mar 01, 2025				
<u>FOMEPIZOLE - ANTIZOL</u>						
N 020696 001	7553863	Jun 30, 2027	DS DP			

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<u>FORMOTEROL FUMARATE; GLYCOPYRROLATE - BEVESPI AEROSPHERE</u>						
N 208294	001	10716753	May 28, 2030	DP	U-2889	
		8324266	May 28, 2030		U-2889	
		8703806	May 28, 2030		U-2889	
		8808713	May 28, 2030	DP	U-2889	
		8815258	Mar 17, 2031		U-2889	
		9415009	May 28, 2030		U-2889	
		9463161	May 28, 2030	DP	U-2889	
<u>FORMOTEROL FUMARATE; MOMETASONE FUROATE - DULERA</u>						
N 022518	003					NS Aug 12, 2022
						PED Feb 12, 2023
<u>FOSDENOPTERIN HYDROBROMIDE - NULIBRY</u>						
N 214018	001	7504095	Jan 31, 2025	DP	U-3092	NCE Feb 26, 2026
						ODE-342 Feb 26, 2028
<u>FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u>						
N 210493	001	10208073	May 23, 2032		U-2301	NCE Apr 19, 2023
		10624911	Jun 02, 2037	DP		
		10717721	May 23, 2032	DS		
		10828297	Dec 17, 2030		U-2301	
		8426450	May 23, 2032	DS DP		
		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>FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u>						
N 210493	002	10208073	May 23, 2032		U-2301	NCE Apr 19, 2023
		10624911	Jun 02, 2037	DP		
		10717721	May 23, 2032	DS		
		10828297	Dec 17, 2030		U-2301	
		8426450	May 23, 2032	DS DP		
		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>FOSPHENYTOIN SODIUM - SESQUIENT</u>						
N 210864	001	7635773	Mar 13, 2029	DP		
		8410077	Mar 13, 2029	DP		
		9200088	Mar 13, 2029	DP		
		9493582	Feb 27, 2033	DP		
		9750822	Mar 13, 2029	DP		
<u>FOSPHENYTOIN SODIUM - SESQUIENT</u>						
N 210864	002	7635773	Mar 13, 2029	DP		
		8410077	Mar 13, 2029	DP		
		9200088	Mar 13, 2029	DP		
		9493582	Feb 27, 2033	DP		
		9750822	Mar 13, 2029	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		

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<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>FOSTEMSAVIR TROMETHAMINE - RUKOBIA</u>						
N 212950 001	7745625	Nov 19, 2027	DS		NCE	Jul 02, 2025
	8168615	Feb 25, 2025		DP		
	8461333	Feb 25, 2025	DS			
<u>FULVESTRANT - FULVESTRANT</u>						
N 210326 001	10188663	Feb 14, 2034		DP	U-2540	
	9271990	May 17, 2034		DP	U-2540	
	9833459	Feb 14, 2034		DP	U-2540	
<u>GABAPENTIN - NEURONTIN</u>						
N 021129 001	7256216	May 28, 2022		DP		
<u>GABAPENTIN - GRALISE</u>						
N 022544 001	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	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	Apr 06, 2025	DS	DP		
	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	

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<u>GABAPENTIN ENACARBIL - HORIZANT</u>						
N 022399	002	6818787	Apr 06, 2025	DS DP		
		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	001				I-801	Jul 12, 2022
<u>GADOBUTROL - GADAVIST</u>						
N 201277	002				I-801	Jul 12, 2022
<u>GADOBUTROL - GADAVIST</u>						
N 201277	003				I-801	Jul 12, 2022
<u>GADOBUTROL - GADAVIST</u>						
N 201277	004				I-801	Jul 12, 2022
<u>GADOBUTROL - GADAVIST</u>						
N 201277	005				I-801	Jul 12, 2022
<u>GADOBUTROL - GADAVIST</u>						
N 201277	006				I-801	Jul 12, 2022
<u>GALLIUM DOTATATE GA-68 - NETSPOT</u>						
N 208547	001	9375498	Aug 10, 2032	DP	ODE-120	Jun 01, 2023
<u>GALLIUM DOTATOC GA-68 - GALLIUM DOTATOC GA 68</u>						
N 210828	001				NCE W	Aug 21, 2024 Aug 21, 2024
<u>GALLIUM GA-68 PSMA-11 - GALLIUM GA 68 PSMA-11</u>						
N 212642	001				NCE W	Dec 01, 2025 Dec 01, 2025
<u>GALLIUM GA-68 PSMA-11 - GALLIUM GA 68 PSMA-11</u>						
N 212643	001				NCE W	Dec 01, 2025 Dec 01, 2025
<u>GANCICLOVIR - GANZYK-RTU</u>						
N 209347	001	9486530	Sep 02, 2034	DP		
<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		

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<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>GILTERITINIB FUMARATE - XOSPATA</u>						
N 211349	001	10786500	Jul 01, 2036	DP	NCE	Nov 28, 2023
		8969336	Jan 27, 2031	DS DP	ODE-222	Nov 28, 2025
		9487491	Jul 28, 2030	U-2456		
<u>GIVOSIRAN SODIUM - GIVLAARI</u>						
N 212194	001	10119143	Oct 03, 2034	DS DP U-2672	NCE	Nov 20, 2024
		10125364	Mar 15, 2033	DS DP U-2672	ODE-273	Nov 20, 2026
		10131907	Aug 24, 2028	DS DP U-2672		
		10273477	Mar 08, 2024	DS		
		8106022	Dec 12, 2029	DS DP U-2672		
		8546143	Jan 09, 2022	DS U-2672		
		8828956	Dec 04, 2028	DS DP U-2672		
		9133461	May 14, 2033	DS DP U-2672		
		9150605	Aug 28, 2025	DS DP		
		9631193	Mar 15, 2033	U-2672		
		9708610	Jan 01, 2024	DS DP U-2672		
		9708615	Mar 08, 2024	DS		
<u>GLASDEGIB MALEATE - DAURISMO</u>						
N 210656	001	10414748	Apr 13, 2036	DS DP	NCE	Nov 21, 2023
		11168066	Apr 13, 2036	U-3254	ODE-224	Nov 21, 2025
		8148401	Jan 30, 2031	DS DP		
		8431597	Jun 29, 2028	DP		
<u>GLASDEGIB MALEATE - DAURISMO</u>						
N 210656	002	10414748	Apr 13, 2036	DS DP	NCE	Nov 21, 2023
		11168066	Apr 13, 2036	U-3254	ODE-224	Nov 21, 2025
		8148401	Jan 30, 2031	DS DP		
		8431597	Jun 29, 2028	DP		
<u>GLECAPREVIR; PIBRENTASVIR - MAVYRET</u>						
N 209394	001	10028937	Jun 10, 2030	U-2141	D-175	Sep 26, 2022
		10028937	Jun 10, 2030	U-3237	M-259	Apr 10, 2023
		10028937*PED	Dec 10, 2030		NCE	Aug 03, 2022
		10039754	Jun 10, 2030	U-2141	NPP	Apr 30, 2022
		10039754	Jun 10, 2030	U-3237	ODE-232	Apr 30, 2026
		10039754*PED	Dec 10, 2030		ODE-233	Apr 30, 2026
		10286029	Mar 14, 2034	U-3237	ODE-372	Jun 10, 2028
		10286029*PED	Sep 14, 2034		PED	Oct 30, 2022
		8648037	Jan 19, 2032	DS DP U-2141	PED	Feb 03, 2023
		8648037	Jan 19, 2032	DS DP U-3237	PED	Mar 26, 2023
		8648037*PED	Jul 19, 2032		PED	Oct 10, 2023
		8937150	May 18, 2032	DS DP	PED	Oct 30, 2026
		8937150*PED	Nov 18, 2032		PED	Oct 30, 2026
		9321807	Jun 05, 2035	DS	PED	Dec 10, 2028
		9321807*PED	Dec 05, 2035			
		9586978	Nov 06, 2030	U-2141		
		9586978	Nov 06, 2030	U-3237		
		9586978*PED	May 06, 2031			
<u>GLECAPREVIR; PIBRENTASVIR - MAVYRET</u>						
N 215110	001	10028937	Jun 10, 2030	U-3238	D-175	Sep 26, 2022
		10028937*PED	Dec 10, 2030		M-259	Apr 10, 2023
		10039754	Jun 10, 2030	U-3238	NCE	Aug 03, 2022
		10039754*PED	Dec 10, 2030		ODE-372	Jun 10, 2028
		10286029	Mar 14, 2034	U-3238	PED	Feb 03, 2023
		10286029*PED	Sep 14, 2034		PED	Mar 26, 2023
		8648037	Jan 19, 2032	DS DP U-3238	PED	Oct 10, 2023
		8648037*PED	Jul 19, 2032		PED	Dec 10, 2028
		8937150	May 18, 2032	DS DP		
		8937150*PED	Nov 18, 2032			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>GLECAPREVIR; PIBRENTASVIR - MAVYRET</u>						
N 215110	001	9321807	Jun 05, 2035	DS DP		
		9321807*PED	Dec 05, 2035			
		9586978	Nov 06, 2030		U-3238	
		9586978*PED	May 06, 2031			
<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>GLUCAGON - BAOSIMI</u>						
N 210134	001	10213487	Feb 16, 2036	DP U-2604	NP	Jul 24, 2022
		10765602	Sep 23, 2039	DP		
		10894133	Jan 03, 2038	DP		
		6938798	Jan 03, 2022	DP		
<u>GLUCAGON - GVOKE PFS</u>						
N 212097	001	9649364	Apr 22, 2036	DP U-2742	NP	Sep 10, 2022
<u>GLUCAGON - GVOKE PFS</u>						
N 212097	002	9649364	Apr 22, 2036	DP U-2742	NP	Sep 10, 2022
<u>GLUCAGON - GVOKE HYPOPEN</u>						
N 212097	003	9649364	Apr 22, 2036	DP U-2742	NP	Sep 10, 2022
<u>GLUCAGON - GVOKE HYPOPEN</u>						
N 212097	004	9649364	Apr 22, 2036	DP U-2742	NP	Sep 10, 2022
<u>GLYCEROL PHENYLBUTYRATE - RAVICTI</u>						
N 203284	001	10045958	Sep 22, 2030	U-1816	ODE-157	Apr 28, 2024
		10045959	Sep 22, 2030	U-1816		
		10183002	Sep 22, 2030	U-1816		
		10183003	Sep 22, 2030	U-1816		
		10183004	Sep 22, 2030	U-1816		
		10183005	Sep 22, 2030	U-1816		
		10183006	Sep 22, 2030	U-1816		
		10668040	Sep 22, 2030	U-1816		
		8404215	Mar 09, 2032	U-1383		
		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 - GLYCOPYRROLATE</u>						
A 204438	001				PC	Jul 03, 2022
<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	8182838	Oct 20, 2028	DP		
		8479730	Oct 11, 2028	DP		
<u>GLYCOPYRROLATE - LONHALA MAGNAIR KIT</u>						
N 208437	001	10376661	Sep 14, 2035	DP		
		10688518	Nov 12, 2036	DP		
		10744277	Dec 07, 2036	DP U-2941		
		10940110	Feb 26, 2029	DP U-1773		
		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		

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>GLYCOPYRROLATE - LONHALA MAGNAIR KIT</u>						
N 208437	001 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		
	8182838	Oct 20, 2028	DP			
	8479730	Oct 11, 2028	DP			
<u>GLYCOPYRRONIUM TOSYLATE - OBREXZA</u>						
N 210361	001 10004717	Feb 28, 2033	DP	U-2398		
	10052267	Oct 17, 2028	DP	U-2398		
	10543192	Feb 28, 2033	DP			
	10548875	Feb 28, 2033	DS DP	U-2398		
	8618160	Dec 10, 2029	DP	U-2398		
	8859610	Feb 28, 2033	DP	U-2398		
	9259414	Feb 28, 2033		U-2398		
	9744105	Jul 18, 2030	DP	U-2398		
<u>GOLODIRSEN - VYONDYS 53</u>						
N 211970	001 10227590	Jun 28, 2025	DS DP		NCE	Dec 12, 2024
	10266827	Jun 28, 2025		U-2675	ODE-280	Dec 12, 2026
	10421966	Jun 28, 2025	DS DP			
	10968450	Jun 28, 2025	DS DP			
	10995337	Jun 28, 2025	DP	U-2675		
	9024007	Jun 28, 2025	DS DP			
	9994851	Jun 28, 2025	DS DP			
	RE47691	Jun 28, 2025	DP			
<u>GOSERELIN ACETATE - ZOLADEX</u>						
N 019726	001 7118552	Apr 13, 2022	DP			
	7220247	Apr 09, 2022	DP			
<u>GOSERELIN ACETATE - ZOLADEX</u>						
N 020578	001 7118552	Apr 13, 2022	DP			
	7220247	Apr 09, 2022	DP			
<u>GRANISETRON - SANCUSO</u>						
N 022198	001 7608282	Jan 22, 2025	DP	U-1011		
<u>GRANISETRON - SUSTOL</u>						
N 022445	001 10357570	Sep 28, 2024		U-2253		
	8252304	Sep 28, 2024	DP			
	8252305	Sep 28, 2024		U-1891		
	8715710	Sep 28, 2024	DP			
	9913910	Sep 28, 2024		U-2253		
<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>GUANFACINE HYDROCHLORIDE - INTUNIV</u>						
N 022037	001 6811794	Jul 04, 2022	DP	U-494		
	6811794*PED	Jan 04, 2023				
<u>GUANFACINE HYDROCHLORIDE - INTUNIV</u>						
N 022037	002 6811794	Jul 04, 2022	DP	U-494		
	6811794*PED	Jan 04, 2023				
<u>GUANFACINE HYDROCHLORIDE - INTUNIV</u>						
N 022037	003 6811794	Jul 04, 2022	DP	U-494		
	6811794*PED	Jan 04, 2023				
<u>GUANFACINE HYDROCHLORIDE - INTUNIV</u>						
N 022037	004 6811794	Jul 04, 2022	DP	U-494		
	6811794*PED	Jan 04, 2023				

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>HALOBETASOL PROPIONATE - ULTRAVATE</u>						
N 208183	001 8962028	Jun 19, 2033	DP U-1775		NPP	Aug 31, 2023
<u>HALOBETASOL PROPIONATE - BRYHALI</u>						
N 209355	001 10478502	Nov 02, 2031	DP U-2625			
	8809307	Nov 02, 2031	DP			
<u>HALOBETASOL PROPIONATE - LEXETTE</u>						
N 210566	001 10857159	Nov 30, 2036	DP			
	10857159*PED	May 30, 2037				
	11020407	Nov 30, 2036	DP U-3143			
<u>HALOBETASOL PROPIONATE; TAZAROTENE - DUOBRII</u>						
N 209354	001 10251895	Jun 06, 2036	DP		NP	Apr 25, 2022
	10426787	Jun 06, 2036	U-2625			
	10478502	Nov 02, 2031	DP U-2625			
	8809307	Nov 02, 2031	DP			
<u>HEXAMINOLEVULINATE HYDROCHLORIDE - CYSVIEW KIT</u>						
N 022555	001 10556010	Dec 19, 2036	U-2250			
<u>HISTRELIN ACETATE - SUPPRELIN LA</u>						
N 022058	001 8062652	Jun 16, 2026	U-1197			
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880	001 10028946	Jul 25, 2033	U-1810			
	10092559	Sep 12, 2034	U-55			
	10322120	Jul 25, 2033	DP			
	10456393	Jul 25, 2033	U-1810			
	10722511	Jul 25, 2033	U-1810			
	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			
	10322120	Jul 25, 2033	DP			
	10456393	Jul 25, 2033	U-1810			
	10722511	Jul 25, 2033	U-1810			
	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	003 10028946	Jul 25, 2033	U-1810			
	10092559	Sep 12, 2034	U-55			
	10322120	Jul 25, 2033	DP			
	10456393	Jul 25, 2033	U-1810			
	10722511	Jul 25, 2033	U-1810			
	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			

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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 003	9421200	Jul 25, 2033				
	9433619	Jul 25, 2033				
	9452163	Sep 12, 2034				
	9486451	Sep 12, 2034				
	9610286	Jul 25, 2033				
	9713611	Sep 12, 2034	DP			
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880 004	10028946	Jul 25, 2033				
	10092559	Sep 12, 2034				
	10322120	Jul 25, 2033	DP			
	10456393	Jul 25, 2033				
	10722511	Jul 25, 2033				
	9132096	Sep 12, 2034	DP			
	9265760	Jul 25, 2033				
	9326982	Jul 25, 2033				
	9333201	Jul 25, 2033				
	9339499	Jul 25, 2033				
	9421200	Jul 25, 2033				
	9433619	Jul 25, 2033				
	9452163	Sep 12, 2034				
	9486451	Sep 12, 2034				
	9610286	Jul 25, 2033				
	9713611	Sep 12, 2034	DP			
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880 005	10028946	Jul 25, 2033				
	10092559	Sep 12, 2034				
	10322120	Jul 25, 2033	DP			
	10456393	Jul 25, 2033				
	10722511	Jul 25, 2033				
	9132096	Sep 12, 2034	DP			
	9265760	Jul 25, 2033				
	9326982	Jul 25, 2033				
	9333201	Jul 25, 2033				
	9339499	Jul 25, 2033				
	9421200	Jul 25, 2033				
	9433619	Jul 25, 2033				
	9452163	Sep 12, 2034				
	9486451	Sep 12, 2034				
	9610286	Jul 25, 2033				
	9713611	Sep 12, 2034	DP			
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880 006	10028946	Jul 25, 2033				
	10092559	Sep 12, 2034				
	10322120	Jul 25, 2033	DP			
	10456393	Jul 25, 2033				
	10722511	Jul 25, 2033				
	9132096	Sep 12, 2034	DP			
	9265760	Jul 25, 2033				
	9326982	Jul 25, 2033				
	9333201	Jul 25, 2033				
	9339499	Jul 25, 2033				
	9421200	Jul 25, 2033				
	9433619	Jul 25, 2033				
	9452163	Sep 12, 2034				
	9486451	Sep 12, 2034				
	9610286	Jul 25, 2033				
	9713611	Sep 12, 2034	DP			
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627 001	10130591	Nov 20, 2023	DP			
	10369109	Jun 16, 2023	DP			
	8309060	Nov 20, 2023	DP			
	8529948	Aug 06, 2022	DP			
	8808740	Dec 21, 2031	DP			
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027				
	9095615	Aug 24, 2027	DP			

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<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	001	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	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031	DP		
		9675610	Jun 16, 2023	DP		
		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 ER</u>						
N 206627	002	10130591	Nov 20, 2023	DP	U-1819	
		10369109	Jun 16, 2023	DP		
		8309060	Nov 20, 2023	DP	U-1556	
		8529948	Aug 06, 2022	DP		
		8808740	Dec 21, 2031	DP	U-1556	
		9084816	Aug 24, 2027	DP		
		9095614	Aug 24, 2027		U-1556	
		9095615	Aug 24, 2027	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	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031	DP		
		9675610	Jun 16, 2023	DP		
		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 ER</u>						
N 206627	003	10130591	Nov 20, 2023	DP	U-1819	
		10369109	Jun 16, 2023	DP		
		8309060	Nov 20, 2023	DP	U-1556	
		8529948	Aug 06, 2022	DP		
		8808740	Dec 21, 2031	DP	U-1556	
		9084816	Aug 24, 2027	DP		
		9095614	Aug 24, 2027		U-1556	
		9095615	Aug 24, 2027	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	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031	DP		
		9675610	Jun 16, 2023	DP		
		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 ER</u>						
N 206627	004	10130591	Nov 20, 2023	DP	U-1819	
		10369109	Jun 16, 2023	DP		
		8309060	Nov 20, 2023	DP	U-1556	
		8529948	Aug 06, 2022	DP		
		8808740	Dec 21, 2031	DP	U-1556	
		9084816	Aug 24, 2027	DP		

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<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	004	9095614	Aug 24, 2027		U-1556	
		9095615	Aug 24, 2027	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	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031	DP		
		9675610	Jun 16, 2023	DP		
		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 ER</u>						
N 206627	005	10130591	Nov 20, 2023	DP	U-1819	
		10369109	Jun 16, 2023	DP		
		8309060	Nov 20, 2023	DP	U-1556	
		8529948	Aug 06, 2022	DP		
		8808740	Dec 21, 2031	DP	U-1556	
		9084816	Aug 24, 2027	DP		
		9095614	Aug 24, 2027		U-1556	
		9095615	Aug 24, 2027	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	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031	DP		
		9675610	Jun 16, 2023	DP		
		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 ER</u>						
N 206627	006	10130591	Nov 20, 2023	DP	U-1819	
		10369109	Jun 16, 2023	DP		
		8309060	Nov 20, 2023	DP	U-1556	
		8529948	Aug 06, 2022	DP		
		8808740	Dec 21, 2031	DP	U-1556	
		9084816	Aug 24, 2027	DP		
		9095614	Aug 24, 2027		U-1556	
		9095615	Aug 24, 2027	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	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031	DP		
		9675610	Jun 16, 2023	DP		
		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 ER</u>						
N 206627	007	10130591	Nov 20, 2023	DP	U-1819	
		10369109	Jun 16, 2023	DP		
		8309060	Nov 20, 2023	DP	U-1556	
		8529948	Aug 06, 2022	DP		

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<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	007	8808740	Dec 21, 2031	DP	U-1556	
		9084816	Aug 24, 2027	DP		
		9095614	Aug 24, 2027		U-1556	
		9095615	Aug 24, 2027	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	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031	DP		
		9675610	Jun 16, 2023	DP		
		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 - 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 - ALKINDI SPRINKLE</u>						
N 213876	001	9649280	May 12, 2034	DP	U-3075	
		9675559	Jan 10, 2033		U-3075	
		9717740	Nov 19, 2032		U-3075	
<u>HYDROCORTISONE - ALKINDI SPRINKLE</u>						
N 213876	002	9649280	May 12, 2034	DP	U-3075	
		9675559	Jan 10, 2033		U-3075	
		9717740	Nov 19, 2032		U-3075	
<u>HYDROCORTISONE - ALKINDI SPRINKLE</u>						
N 213876	003	9649280	May 12, 2034	DP	U-3075	
		9675559	Jan 10, 2033		U-3075	
		9717740	Nov 19, 2032		U-3075	
<u>HYDROCORTISONE - ALKINDI SPRINKLE</u>						
N 213876	004	9649280	May 12, 2034	DP	U-3075	
		9675559	Jan 10, 2033		U-3075	
		9717740	Nov 19, 2032		U-3075	
<u>HYDROCORTISONE BUTYRATE - LOCROID</u>						
N 022076	001	7378405	Dec 19, 2026	DP		
		7981877	Jan 23, 2025	DP		

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROGEN PEROXIDE - ESKATA</u>						
N 209305	001	10098910	Apr 21, 2035	DP	U-2205	
		10493103	Apr 21, 2035	DP		
		10729720	Apr 21, 2035	DP		
		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	9248229	Mar 12, 2034	DP		
		9731082	Apr 23, 2032	DP		
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID-HP</u>						
N 019034	002	9248229	Mar 12, 2034	DP		
		9731082	Apr 23, 2032	DP		
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034	003	9248229	Mar 12, 2034	DP		
		9731082	Apr 23, 2032	DP		
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034	004	9248229	Mar 12, 2034	DP		
		9731082	Apr 23, 2032	DP		
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034	005	9248229	Mar 12, 2034	DP		
		9731082	Apr 23, 2032	DP		
<u>HYDROXYPROGESTERONE CAPROATE - MAKENA (AUTOINJECTOR)</u>						
N 021945	004	10471075	May 02, 2036		U-2236	
		11154562	May 02, 2036		U-2236	
		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	
<u>HYDROXYUREA - SIKLOS</u>						
N 208843	001				ODE-177	Dec 21, 2024
<u>HYDROXYUREA - SIKLOS</u>						
N 208843	002				ODE-177	Dec 21, 2024
<u>IBANDRONATE SODIUM - BONIVA</u>						
N 021455	002	7192938	May 06, 2023		U-798	
		7410957	May 06, 2023		U-887	
		7718634	May 06, 2023		U-642	
<u>IBREXAFUNGERP CITRATE - BREXAFEMME</u>						
N 214900	001	10174074	Jan 19, 2035	DS DP		NCE Jun 01, 2026
		10370406	Jan 19, 2035		U-3159	GAIN Jun 01, 2031
		10927142	Jan 19, 2035	DS		
		8188085	Aug 28, 2030	DS DP	U-3159	
<u>IBRUTINIB - IMBRUVICA</u>						
N 205552	001	10004746	Jun 03, 2031		U-1684	M-236 Jan 25, 2022
		10004746	Jun 03, 2031		U-1946	ODE-109 Mar 04, 2023
		10004746	Jun 03, 2031		U-2241	ODE-117 May 06, 2023
		10004746	Jun 03, 2031		U-2242	ODE-128 Jan 18, 2024
		10016435	Jun 03, 2031		U-1650	ODE-152 Aug 02, 2024
		10106548	Jun 03, 2033	DS DP		ODE-86 Jan 29, 2022
		10125140	Jun 03, 2033	DS DP		
		10294231	Jun 03, 2033		DP	
		10294232	Jun 03, 2033		DP	
		10463668	Oct 24, 2034		U-2654	
		10478439	Jun 03, 2031		U-1456	
		10478439	Jun 03, 2031		U-1650	
		10478439	Jun 03, 2031		U-1684	
		10478439	Jun 03, 2031		U-1946	

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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	10478439	Jun 03, 2031		U-1947	
		10478439	Jun 03, 2031		U-2241	
		10478439	Jun 03, 2031		U-2242	
		10478439	Jun 03, 2031		U-2665	
		10653696	Jun 03, 2031		U-1456	
		10695350	Oct 24, 2034		U-2846	
		10751342	Jun 03, 2031		U-1491	
		10751342	Jun 03, 2031		U-1946	
		10751342	Jun 03, 2031		U-2943	
		10751342	Jun 03, 2031		U-2944	
		10752634	Jun 03, 2033	DP		
		10961251	Jun 03, 2033	DP		
		7514444	Dec 28, 2026	DS DP		
		8008309	Nov 13, 2027	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-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	
		9795604	Oct 24, 2034		U-2969	
		9795604	Oct 24, 2034		U-2970	
		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	M-236
		10004746	Jun 03, 2031		U-1946	Jan 25, 2022
		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		
		10294231	Jun 03, 2033	DP		
		10294232	Jun 03, 2033	DP		
		10463668	Oct 24, 2034		U-2654	
		10478439	Jun 03, 2031		U-1456	

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<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 002	10478439	Jun 03, 2031	U-1650			
	10478439	Jun 03, 2031	U-1684			
	10478439	Jun 03, 2031	U-1946			
	10478439	Jun 03, 2031	U-1947			
	10478439	Jun 03, 2031	U-2241			
	10478439	Jun 03, 2031	U-2242			
	10478439	Jun 03, 2031	U-2665			
	10653696	Jun 03, 2031	U-1456			
	10695350	Oct 24, 2034	U-2846			
	10751342	Jun 03, 2031	U-1491			
	10751342	Jun 03, 2031	U-1946			
	10751342	Jun 03, 2031	U-2943			
	10751342	Jun 03, 2031	U-2944			
	10961251	Jun 03, 2033	DP			
	7514444	Dec 28, 2026	DS DP			
	8008309	Nov 13, 2027	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			
	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-2969			
	9795604	Oct 24, 2034	U-2970			
	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		M-236	Jan 25, 2022
	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			
	10213386	Mar 03, 2036	DP			

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<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 001	10463668	Oct 24, 2034				U-2654
	10478439	Jun 03, 2031				U-1456
	10478439	Jun 03, 2031				U-1650
	10478439	Jun 03, 2031				U-1684
	10478439	Jun 03, 2031				U-1946
	10478439	Jun 03, 2031				U-1947
	10478439	Jun 03, 2031				U-2241
	10478439	Jun 03, 2031				U-2242
	10478439	Jun 03, 2031				U-2665
	10653696	Jun 03, 2031				U-1456
	10695350	Oct 24, 2034				U-2846
	10751342	Jun 03, 2031				U-1491
	10751342	Jun 03, 2031				U-1946
	10751342	Jun 03, 2031				U-2943
	10751342	Jun 03, 2031				U-2944
	10752634	Jun 03, 2033		DP		
	10828259	Mar 03, 2036		DP		
	10961251	Jun 03, 2033		DP		
	7514444	Dec 28, 2026	DS	DP		
	8008309	Nov 13, 2027	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-2969
	9795604	Oct 24, 2034				U-2970
	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

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<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 002	10004746	Jun 03, 2031		U-1684	M-236	Jan 25, 2022
	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			
	10213386	Mar 03, 2036		DP		
	10463668	Oct 24, 2034		U-2654		
	10478439	Jun 03, 2031		U-1456		
	10478439	Jun 03, 2031		U-1650		
	10478439	Jun 03, 2031		U-1684		
	10478439	Jun 03, 2031		U-1946		
	10478439	Jun 03, 2031		U-1947		
	10478439	Jun 03, 2031		U-2241		
	10478439	Jun 03, 2031		U-2242		
	10478439	Jun 03, 2031		U-2665		
	10653696	Jun 03, 2031		U-1456		
	10695350	Oct 24, 2034		U-2846		
	10751342	Jun 03, 2031		U-1491		
	10751342	Jun 03, 2031		U-1946		
	10751342	Jun 03, 2031		U-2943		
	10751342	Jun 03, 2031		U-2944		
	10828259	Mar 03, 2036		DP		
	10961251	Jun 03, 2033		DP		
	7514444	Dec 28, 2026	DS DP			
	8008309	Nov 13, 2027	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-2969		

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<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 002	9795604	Oct 24, 2034	U-2970			
	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		M-236	Jan 25, 2022
	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			
	10213386	Mar 03, 2036	DP			
	10463668	Oct 24, 2034	U-2654			
	10478439	Jun 03, 2031	U-1456			
	10478439	Jun 03, 2031	U-1650			
	10478439	Jun 03, 2031	U-1684			
	10478439	Jun 03, 2031	U-1946			
	10478439	Jun 03, 2031	U-1947			
	10478439	Jun 03, 2031	U-2241			
	10478439	Jun 03, 2031	U-2242			
	10478439	Jun 03, 2031	U-2665			
	10653696	Jun 03, 2031	U-1456			
	10695350	Oct 24, 2034	U-2846			
	10751342	Jun 03, 2031	U-1491			
	10751342	Jun 03, 2031	U-1946			
	10751342	Jun 03, 2031	U-2943			
	10751342	Jun 03, 2031	U-2944			
	10828259	Mar 03, 2036	DP			
	10961251	Jun 03, 2033	DP			
	7514444	Dec 28, 2026	DS DP			
	8008309	Nov 13, 2027	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			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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	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-2969		
		9795604	Oct 24, 2034	U-2970		
		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	004	10004746	Jun 03, 2031	U-1684	M-236	Jan 25, 2022
		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		
		10213386	Mar 03, 2036	DP		
		10463668	Oct 24, 2034	U-2654		
		10478439	Jun 03, 2031	U-1456		
		10478439	Jun 03, 2031	U-1650		
		10478439	Jun 03, 2031	U-1684		
		10478439	Jun 03, 2031	U-1946		
		10478439	Jun 03, 2031	U-1947		
		10478439	Jun 03, 2031	U-2241		
		10478439	Jun 03, 2031	U-2242		
		10478439	Jun 03, 2031	U-2665		
		10653696	Jun 03, 2031	U-1456		
		10695350	Oct 24, 2034	U-2846		
		10751342	Jun 03, 2031	U-1491		
		10751342	Jun 03, 2031	U-1946		
		10751342	Jun 03, 2031	U-2943		
		10751342	Jun 03, 2031	U-2944		
		10828259	Mar 03, 2036	DP		
		10961251	Jun 03, 2033	DP		
		7514444	Dec 28, 2026	DS DP		
		8008309	Nov 13, 2027	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		

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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	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-2969		
		9795604	Oct 24, 2034	U-2970		
		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 - CHILDREN'S ADVIL-FLAVORED</u>						
N 020589	002	10238640	May 25, 2024	DP		
<u>IBUPROFEN - CALDOLOR</u>						
N 022348	001				M-128	Nov 19, 2024
<u>IBUPROFEN - CALDOLOR</u>						
N 022348	002	8735452	Sep 30, 2029	U-981	M-128	Nov 19, 2024
		8871810	Sep 30, 2029	U-981		
		9012508	Sep 14, 2030	U-981		
		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 - CALDOLOR</u>						
N 022348	003	8735452	Sep 30, 2029	U-981	M-128	Nov 19, 2024
		8871810	Sep 30, 2029	U-981		
		9012508	Sep 14, 2030	U-981		
		9072661	Mar 16, 2032	U-2264		
		9072710	Mar 16, 2032	U-2266		
<u>IBUPROFEN LYSINE - NEOPROFEN</u>						
N 021903	001	8415337	Mar 02, 2032	DS DP		
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057	001	10010517	Apr 29, 2030	U-2690	I-819	Dec 13, 2022
		10265287	Apr 29, 2030	U-2700		
		10278935	Jun 28, 2033	U-2701		
		10278936	Jun 28, 2033	U-2702		
		10278937	Jun 28, 2033	U-2703		
		10383840	Jun 28, 2033	U-2704		
		10555924	Jun 28, 2033	U-2743		
		10555925	Jun 28, 2033	U-2744		
		10568861	Jun 28, 2033	U-2756		
		10576054	Jun 28, 2033	U-2762		
		10668042	Jun 28, 2033	U-2841		
		10786478	Jun 28, 2033	U-2959		
		10786478	Jun 28, 2033	U-2960		
		10792267	Apr 29, 2030	U-2961		
		10792270	Jun 28, 2033	U-2962		
		10842766	Apr 29, 2030	U-2997		
		10842768	Jun 15, 2030	U-2688		
		10881632	Apr 29, 2030	U-3052		
		10894028	Jun 28, 2033	U-3053		
		11000499	Jun 28, 2033	U-3126		
		11103477	Apr 29, 2030	U-3209		

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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	11116742	Jun 28, 2033	U-3221			
	11154526	Apr 29, 2030	U-3240			
	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			
	8410086	Jun 15, 2030	U-2688			
	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			
	8454994	Apr 29, 2030	U-2689			
	8455472	Jun 15, 2030	U-2690			
	8501225	Apr 29, 2030	U-1287			
	8518929	Feb 09, 2030	U-1287			
	8524698	Feb 09, 2030	U-1287			
	8546372	Feb 09, 2030	U-1287			
	8551521	Apr 29, 2030	U-1287			
	8563608	Apr 29, 2030	U-1287			
	8617593	Apr 29, 2030	U-1478			
	8617593	Apr 29, 2030	U-2691			
	8617594	Apr 29, 2030	U-1287			
	8618166	Apr 29, 2030	U-2689			
	8623406	Apr 29, 2030	U-1478			
	8623406	Apr 29, 2030	U-2692			
	8642077	Apr 29, 2030	U-2693			
	8669245	Jun 15, 2030	U-2694			
	8680144	Feb 09, 2030	U-2695			
	8691871	Apr 29, 2030	U-2689			
	8703185	Apr 29, 2030	U-2691			
	8709475	Apr 29, 2030	U-2689			
	8710041	Jun 15, 2030	U-2690			
	9198892	Sep 25, 2027	U-2706			
	9603826	Jun 28, 2033	U-2696			
	9610272	Jun 28, 2033	U-2697			
	9623001	Jun 28, 2033	U-2698			
	9693984	Jun 28, 2033	U-2697			
	9693985	Jun 28, 2033	U-2696			
	9693986	Jun 28, 2033	U-2698			
	9700537	May 31, 2027	U-2707			
	9918954	Jun 28, 2033	U-2699			
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057 002	10010517	Apr 29, 2030	U-2690		I-819	Dec 13, 2022
	10265287	Apr 29, 2030	U-2700			
	10278935	Jun 28, 2033	U-2701			
	10278936	Jun 28, 2033	U-2702			
	10278937	Jun 28, 2033	U-2703			
	10383840	Jun 28, 2033	U-2704			
	10555924	Jun 28, 2033	U-2743			
	10555925	Jun 28, 2033	U-2744			
	10568861	Jun 28, 2033	U-2756			
	10576054	Jun 28, 2033	U-2762			
	10668042	Jun 28, 2033	U-2841			
	10786478	Jun 28, 2033	U-2959			
	10786478	Jun 28, 2033	U-2960			
	10792267	Apr 29, 2030	U-2961			
	10792270	Jun 28, 2033	U-2962			
	10842766	Apr 29, 2030	U-2997			
	10842768	Jun 15, 2030	U-2688			
	10881632	Apr 29, 2030	U-3052			
	10894028	Jun 28, 2033	U-3053			

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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	002	11000499	Jun 28, 2033		U-3126	
		11103477	Apr 29, 2030		U-3209	
		11116742	Jun 28, 2033		U-3221	
		11154526	Apr 29, 2030		U-3240	
		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	
		8410086	Jun 15, 2030		U-2688	
		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	
		8454994	Apr 29, 2030		U-2689	
		8501225	Apr 29, 2030		U-1287	
		8518929	Feb 09, 2030		U-1287	
		8524698	Feb 09, 2030		U-1287	
		8546372	Feb 09, 2030		U-1287	
		8551521	Apr 29, 2030		U-1287	
		8563608	Apr 29, 2030		U-1287	
		8617593	Apr 29, 2030		U-1287	
		8617593	Apr 29, 2030		U-2691	
		8617594	Apr 29, 2030		U-1287	
		8623406	Apr 29, 2030		U-1287	
		8623406	Apr 29, 2030		U-2692	
		8642077	Apr 29, 2030		U-2693	
		8669245	Jun 15, 2030		U-2694	
		8680144	Feb 09, 2030		U-2695	
		8691871	Apr 29, 2030		U-2689	
		8703185	Apr 29, 2030		U-2691	
		8709475	Apr 29, 2030		U-2689	
		8710041	Jun 15, 2030		U-2690	
		9198892	Sep 25, 2027		U-2706	
		9603826	Jun 28, 2033		U-2696	
		9610272	Jun 28, 2033		U-2697	
		9623001	Jun 28, 2033		U-2698	
		9693984	Jun 28, 2033		U-2697	
		9693985	Jun 28, 2033		U-2696	
		9693986	Jun 28, 2033		U-2698	
		9700537	May 31, 2027		U-2707	
		9918954	Jun 28, 2033		U-2699	
<u>IDELALISIB - ZYDELIG</u>						
N 205858	001	10730879	Mar 05, 2033	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	10730879	Mar 05, 2033	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		

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<u>ILOPERIDONE - FANAPT</u>						
N 022192	001	8586610				
		Nov 02, 2027	U-1625			
		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			
		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			
		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			
		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			
		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			
		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	007	8586610				
		Nov 02, 2027	U-1625			
		8652776				
		Aug 31, 2030	U-1685			
		8999638				
		Oct 28, 2030	U-1674			
		9072742				
		Jan 16, 2031	U-1674			

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<u>ILOPERIDONE - FANAPT</u>						
N 022192	007	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 021588	001	6958335*PED	Jun 19, 2022			
<u>IMATINIB MESYLATE - GLEEVEC</u>						
N 021588	002	6958335*PED	Jun 19, 2022			
<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	10238644	Dec 11, 2029		U-68	
		10238645	Aug 18, 2029		U-1455	
		10238645	Aug 18, 2029		U-172	
		10918635	Apr 30, 2030		U-1455	
		10918635	Apr 30, 2030		U-172	
		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	
		8479730	Oct 11, 2028	DP		
<u>INDOCYANINE GREEN - SPY AGENT GREEN KIT</u>						
N 211580	001	10631746	Aug 04, 2035	DP	U-2815	
		8185176	Jun 04, 2028		U-2462	
		8406860	Apr 09, 2029		U-2463	
		8647605	Feb 11, 2029		U-2464	
		8647605	Feb 11, 2029		U-2468	
		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>INFIGRATINIB PHOSPHATE - TRUSELTIQ</u>						
N 214622	001	10278969	Dec 11, 2034	DP	NCE	May 28, 2026
		11160804	Dec 11, 2034	DP	ODE-353	May 28, 2028
		8552002	Aug 25, 2029	DS DP		
		9067896	Aug 06, 2028	DS		
<u>INFIGRATINIB PHOSPHATE - TRUSELTIQ</u>						
N 214622	002	10278969	Dec 11, 2034	DP	NCE	May 28, 2026
		11160804	Dec 11, 2034	DP	ODE-353	May 28, 2028
		8552002	Aug 25, 2029	DS DP		
		9067896	Aug 06, 2028	DS		
<u>INGENOL MEBUTATE - PICATO</u>						
N 202833	001	8278292	Jul 06, 2027	DP		
		8372827	Dec 18, 2026	DP		
		8372828	Dec 18, 2026	DP		
		8377919	Dec 18, 2026	DP		

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<u>INGENOL MEBUTATE - PICATO</u>						
N 202833	001	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	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>IOBENGUANE I-131 - AZEDRA</u>						
N 209607	001				ODE-204	Jul 30, 2025
<u>IPRATROPIUM BROMIDE - ATROVENT HFA</u>						
N 021527	001	8474447	Jan 17, 2030	DP		
<u>IRINOTECAN HYDROCHLORIDE - ONIVYDE</u>						
N 207793	001	10456360	Oct 15, 2036	DP	ODE-99	Oct 22, 2022
		10722508	May 02, 2025	DS DP		
		10980795	Jun 12, 2033	U-1848		
		10993914	Oct 15, 2036	DP		
		8147867	Aug 29, 2028	DS DP		
		8329213	Jan 06, 2027	DS DP		
		8703181	May 02, 2025	U-1434		
		8992970	May 02, 2025	DS DP		
		9339497	Jun 12, 2033	U-1848		
		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	10206879	Sep 14, 2027	DP	NCE	Mar 06, 2020
		10603280	Sep 14, 2027	DP	ODE-305	Mar 06, 2022
		6812238	Oct 31, 2025	DS	ODE-90	Mar 06, 2022
					GAIN	Mar 06, 2025
					GAIN	Mar 06, 2027
					GAIN	Mar 06, 2027
<u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u>						
N 207501	001	6812238	Oct 31, 2025	DS	NCE	Mar 06, 2020
					ODE-305	Mar 06, 2022
					ODE-90	Mar 06, 2022
					GAIN	Mar 06, 2025

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<u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u>						
N 207501	001				GAIN	Mar 06, 2027
					GAIN	Mar 06, 2027
<u>ISOPROPYL ALCOHOL - ZURAGARD</u>						
N 210872	001	10688291	Dec 20, 2034	DP U-1397	M-268	Jun 24, 2024
		8226971	May 06, 2025	DP	NP	Apr 26, 2022
		8389583	Aug 09, 2029	U-1397		
		8703828	May 23, 2028	DP		
		9011897	Feb 08, 2025	DP		
		9629368	May 23, 2028	U-1397		
		9844654	Apr 24, 2036	DP U-1397		
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913	001	9700535	Aug 04, 2035	DP		
		9750711	May 29, 2035	DP		
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913	002	9700535	Aug 04, 2035	DP		
		9750711	May 29, 2035	DP		
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913	003	9700535	Aug 04, 2035	DP		
		9750711	May 29, 2035	DP		
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913	004	9700535	Aug 04, 2035	DP		
		9750711	May 29, 2035	DP		
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913	005	9700535	Aug 04, 2035	DP		
		9750711	May 29, 2035	DP		
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913	006	9700535	Aug 04, 2035	DP		
		9750711	May 29, 2035	DP		
<u>ISTRADEFYLLINE - NOURIANZ</u>						
N 022075	001	7541363	Nov 13, 2024	DS DP	NCE	Aug 27, 2024
		7727993	Jan 28, 2023	U-2623		
		7727994	Jan 18, 2023	U-2623		
		8318201	Sep 05, 2027	DP		
<u>ISTRADEFYLLINE - NOURIANZ</u>						
N 022075	002	7541363	Nov 13, 2024	DS DP	NCE	Aug 27, 2024
		7727993	Jan 28, 2023	U-2623		
		7727994	Jan 18, 2023	U-2623		
		8318201	Sep 05, 2027	DP		
<u>ITRACONAZOLE - ONMEL</u>						
N 022484	001	8486456	Oct 03, 2028	DP U-1054		
<u>ITRACONAZOLE - TOLSURA</u>						
N 208901	001	10463740	Jun 21, 2033	DP U-2453		
		10806792	Jun 21, 2033	DP		
		8771739	Jul 25, 2023	DP		
		8921374	Jun 21, 2033	DP		
		9272046	Jun 21, 2033	DP		
		9713642	Jun 21, 2033	U-2453		
<u>IVABRADINE - CORLANOR</u>						
N 209964	001	7361649	Feb 22, 2026	DS DP U-1694	NP	Apr 22, 2022
		7361650	Feb 22, 2026	DS DP U-1694	ODE-234	Apr 22, 2026
		7867996	Dec 12, 2026	DS DP U-1694	PED	Oct 22, 2022
		7879842	Feb 22, 2026	DS DP U-1694	PED	Oct 22, 2026
<u>IVABRADINE HYDROCHLORIDE - CORLANOR</u>						
N 206143	001	7361649	Feb 22, 2026	DS DP U-1694		
		7361649*PED	Aug 22, 2026			
		7361650	Feb 22, 2026	DS DP U-1694		
		7361650*PED	Aug 22, 2026			
		7867996	Dec 12, 2026	DS DP U-1694		

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<u>IVABRADINE HYDROCHLORIDE - CORLANOR</u>						
N 206143	001	7867996*PED	Jun 12, 2027			
		7879842	Feb 22, 2026	DS DP	U-1694	
		7879842*PED	Aug 22, 2026			
<u>IVABRADINE HYDROCHLORIDE - CORLANOR</u>						
N 206143	002	7361649	Feb 22, 2026	DS DP	U-1694	
		7361649*PED	Aug 22, 2026			
		7361650	Feb 22, 2026	DS DP	U-1694	
		7361650*PED	Aug 22, 2026			
		7867996	Dec 12, 2026	DS DP	U-1694	
		7867996*PED	Jun 12, 2027			
		7879842	Feb 22, 2026	DS DP	U-1694	
		7879842*PED	Aug 22, 2026			
<u>IVACAFTOR - KALYDECO</u>						
N 203188	001	10646481	Aug 13, 2029	DP	ODE-189	Jul 31, 2024
		7495103	May 20, 2027	DS DP	ODE-190	May 17, 2024
		8324242	Aug 05, 2027		ODE-199	Aug 15, 2025
		8324242	Aug 05, 2027		U-1311	
		8324242	Aug 05, 2027		U-1906	
		8354427	Jul 06, 2026		U-1311	
		8354427	Jul 06, 2026		U-1905	
		8410274	Dec 28, 2026	DP		
		8629162	Jun 24, 2025		U-2234	
		8754224	Dec 28, 2026	DS DP		
		9670163	Dec 28, 2026	DP	U-1311	
<u>IVACAFTOR - KALYDECO</u>						
N 207925	001	10272046	Feb 27, 2033	DP	NPP	Apr 29, 2022
		10646481	Aug 13, 2029	DP	ODE-188	Mar 17, 2022
		7495103	May 20, 2027	DS DP	ODE-189	Jul 31, 2024
		8324242	Aug 05, 2027		ODE-190	May 17, 2024
		8324242	Aug 05, 2027		ODE-199	Aug 15, 2025
		8324242	Aug 05, 2027		U-2527	
		8354427	Jul 06, 2026		ODE-236	Apr 29, 2026
		8354427	Jul 06, 2026		ODE-338	Dec 21, 2027
		8354427	Jul 06, 2026		U-1311	
		8354427	Jul 06, 2026		U-1905	
		8354427	Jul 06, 2026		U-2528	
		8410274	Dec 28, 2026	DP		
		8629162	Jun 24, 2025		U-2234	
		8629162	Jun 24, 2025		U-2529	
		8754224	Dec 28, 2026	DS DP		
		8883206	Feb 27, 2033	DP		
		9670163	Dec 28, 2026	DP	U-1311	
		9670163	Dec 28, 2026	DP	U-2530	
<u>IVACAFTOR - KALYDECO</u>						
N 207925	002	10272046	Feb 27, 2033	DP	NPP	Apr 29, 2022
		10646481	Aug 13, 2029	DP	ODE-188	Mar 17, 2022
		7495103	May 20, 2027	DS DP	ODE-189	Jul 31, 2024
		8324242	Aug 05, 2027		ODE-190	May 17, 2024
		8324242	Aug 05, 2027		ODE-199	Aug 15, 2025
		8324242	Aug 05, 2027		ODE-236	Apr 29, 2026
		8354427	Jul 06, 2026		ODE-338	Dec 21, 2027
		8354427	Jul 06, 2026		U-1311	
		8354427	Jul 06, 2026		U-1906	
		8354427	Jul 06, 2026		U-2527	
		8354427	Jul 06, 2026		U-1311	
		8354427	Jul 06, 2026		U-1905	
		8354427	Jul 06, 2026		U-2528	
		8410274	Dec 28, 2026	DP		
		8629162	Jun 24, 2025		U-2234	
		8629162	Jun 24, 2025		U-2529	
		8754224	Dec 28, 2026	DS DP		
		8883206	Feb 27, 2033	DP		
		9670163	Dec 28, 2026	DP	U-1311	
		9670163	Dec 28, 2026	DP	U-2530	
<u>IVACAFTOR - KALYDECO</u>						
N 207925	003	10272046	Feb 27, 2033	DP	NPP	Apr 29, 2022
		10646481	Aug 13, 2029	DP	ODE-188	Mar 17, 2022
		7495103	May 20, 2027	DS DP	ODE-189	Jul 31, 2024
		8324242	Aug 05, 2027		ODE-190	May 17, 2024
		8324242	Aug 05, 2027		ODE-199	Aug 15, 2025
		8324242	Aug 05, 2027		ODE-236	Apr 29, 2026
		8354427	Jul 06, 2026		ODE-338	Dec 21, 2027
		8354427	Jul 06, 2026		U-1311	
		8354427	Jul 06, 2026		U-1906	
		8354427	Jul 06, 2026		U-2963	
		8354427	Jul 06, 2026		U-1311	

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<u>IVACAFTOR - KALYDECO</u>						
N 207925	003	8354427	Jul 06, 2026	U-1905		
		8354427	Jul 06, 2026	U-2964		
		8410274	Dec 28, 2026	DP		
		8629162	Jun 24, 2025	U-2234		
		8629162	Jun 24, 2025	U-2965		
		8754224	Dec 28, 2026	DS DP		
		8883206	Feb 27, 2033	DP		
		9670163	Dec 28, 2026	DP U-1311		
		9670163	Dec 28, 2026	DP U-2966		
<u>IVACAFTOR; IVACAFTOR, TEZACAFTOR - SYMDEKO (COPACKAGED)</u>						
N 210491	001	10022352	Apr 09, 2027	DP U-2343	NCE	Feb 12, 2023
		10022352	Apr 09, 2027	DP U-2573	NPP	Jun 21, 2022
		10058546	Jul 15, 2033	U-2399	ODE-173	Feb 12, 2025
		10058546	Jul 15, 2033	U-2572	ODE-247	Jun 21, 2026
		10058546	Jul 15, 2033	U-3022	ODE-335	Dec 21, 2027
		10058546	Jul 15, 2033	U-3023		
		10081621	Mar 25, 2031	DP U-2420		
		10081621	Mar 25, 2031	DP U-2571		
		10081621	Mar 25, 2031	DP U-3024		
		10081621	Mar 25, 2031	DP U-3025		
		10206877	Apr 14, 2035	DP U-2498		
		10206877	Apr 14, 2035	DP U-2570		
		10206877	Apr 14, 2035	DP U-3026		
		10206877	Apr 14, 2035	DP U-3027		
		10239867	Apr 09, 2027	DS DP U-2512		
		10239867	Apr 09, 2027	DS DP U-2569		
		10646481	Aug 13, 2029	DP		
		7495103	May 20, 2027	DS DP		
		7645789	May 01, 2027	DS DP		
		7776905	Jun 03, 2027	DS DP		
		8324242	Aug 05, 2027	U-2246		
		8354427	Jul 06, 2026	U-3021		
		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		
		9931334	Dec 28, 2026	DP U-2575		
		9974781	Apr 09, 2027	DP U-2318		
		9974781	Apr 09, 2027	DP U-2574		
<u>IVACAFTOR; IVACAFTOR, TEZACAFTOR - SYMDEKO (COPACKAGED)</u>						
N 210491	002	10022352	Apr 09, 2027	DP U-2343	NCE	Feb 12, 2023
		10022352	Apr 09, 2027	DP U-2573	NPP	Jun 21, 2022
		10058546	Jul 15, 2033	U-2399	ODE-173	Feb 12, 2025
		10058546	Jul 15, 2033	U-2572	ODE-247	Jun 21, 2026
		10058546	Jul 15, 2033	U-3022	ODE-335	Dec 21, 2027
		10058546	Jul 15, 2033	U-3023		
		10081621	Mar 25, 2031	DP U-2420		
		10081621	Mar 25, 2031	DP U-2571		
		10081621	Mar 25, 2031	DP U-3024		
		10081621	Mar 25, 2031	DP U-3025		
		10206877	Apr 14, 2035	DP U-2498		
		10206877	Apr 14, 2035	DP U-2570		
		10206877	Apr 14, 2035	DP U-3026		
		10206877	Apr 14, 2035	DP U-3027		
		10239867	Apr 09, 2027	DS DP U-2512		
		10239867	Apr 09, 2027	DS DP U-2569		
		10646481	Aug 13, 2029	DP		
		7495103	May 20, 2027	DS DP		
		7645789	May 01, 2027	DS DP		
		7776905	Jun 03, 2027	DS DP		
		8324242	Aug 05, 2027	U-2246		
		8354427	Jul 06, 2026	U-3021		

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<u>IVACAFTOR; IVACAFTOR, TEZACAFTOR - SYMDEKO (COPACKAGED)</u>						
N 210491	002	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		
		9931334	Dec 28, 2026	DP U-2575		
		9974781	Apr 09, 2027	DP U-2318		
		9974781	Apr 09, 2027	DP U-2574		
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 206038	001	10076513	Dec 04, 2028	DP U-2411	ODE-123	Sep 28, 2023
		10597384	Dec 04, 2028	DS DP U-2777	ODE-93	Jul 02, 2022
		10646481	Aug 13, 2029	DP		
		11052075	Dec 04, 2028	DP U-3181		
		7495103	May 20, 2027	DS DP		
		7973038	Nov 08, 2026	U-1973		
		8324242	Aug 05, 2027	U-1311		
		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	10597384	Dec 04, 2028	DS DP U-2777	ODE-123	Sep 28, 2023
		10646481	Aug 13, 2029	DP	ODE-93	Jul 02, 2022
		11052075	Dec 04, 2028	DP U-3181		
		7495103	May 20, 2027	DS DP		
		7973038	Nov 08, 2026	U-1973		
		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-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	10597384	Dec 04, 2028	DS DP U-2778	ODE-195	Aug 07, 2025
		10646481	Aug 13, 2029	DP		
		7495103	May 20, 2027	DS DP		
		7973038	Nov 08, 2026	U-2374		
		8324242	Aug 05, 2027	U-2374		
		8410274	Dec 28, 2026	DP		
		8507534	Sep 20, 2030	DS DP		
		8653103	Dec 04, 2028	DP		
		8716338	Sep 20, 2030	DP U-2396		

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<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 211358 001	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	10597384	Dec 04, 2028	DS DP		U-2778	ODE-195 Aug 07, 2025
	10646481	Aug 13, 2029			DP	
	7495103	May 20, 2027	DS DP			
	7973038	Nov 08, 2026			U-2374	
	8324242	Aug 05, 2027			U-2374	
	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 - SOOLANTRA</u>						
N 206255 001	10206939	Mar 13, 2034			U-1631	
	11033565	Apr 22, 2024	DP			
	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	10449184	Mar 13, 2035	DP		I-816	May 02, 2022
	10610125	Jun 21, 2030			I-875	Aug 25, 2024
	10610125	Jun 21, 2030			NCE	Jul 20, 2023
	10717764	Jan 18, 2033			ODE-203	Jul 20, 2025
	10799490	Mar 13, 2035	DP		ODE-242	May 02, 2026
	10799490	Mar 13, 2035	DP		ODE-368	Aug 25, 2028
	10980788	Jun 07, 2039			U-3112	
	10980788	Jun 07, 2039			U-3113	
	10980788	Jun 07, 2039			U-3214	
	9474779	Aug 19, 2033	DS DP		U-2350	
	9474779	Aug 19, 2033	DS DP		U-2533	
	9474779	Aug 19, 2033	DS DP		U-2534	
	9474779	Aug 19, 2033	DS DP		U-3213	
	9850277	Jan 18, 2033	DS DP		U-2350	
	9850277	Jan 18, 2033	DS DP		U-2533	
	9850277	Jan 18, 2033	DS DP		U-2534	
	9850277	Jan 18, 2033	DS DP		U-3213	
	9968595	Mar 13, 2035	DP		U-2351	
	9968595	Mar 13, 2035	DP		U-2533	
	9968595	Mar 13, 2035	DP		U-2534	
<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	

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<u>IXABEPILONE - IXEMPRA KIT</u>						
N 022065	001	7312237	Aug 21, 2024	U-965		
		RE41393	Feb 08, 2022	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		
		7312237	Aug 21, 2024	U-965		
		RE41393	Feb 08, 2022	U-961		
<u>IXAZOMIB CITRATE - NINLARO</u>						
N 208462	001	7442830	Nov 20, 2029	DS DP U-2434	ODE-103	Nov 20, 2022
		7687662	Aug 06, 2027	DS DP		
		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	Nov 20, 2029	DS DP U-2434	ODE-103	Nov 20, 2022
		7687662	Aug 06, 2027	DS DP		
		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	Nov 20, 2029	DS DP U-2434	ODE-103	Nov 20, 2022
		7687662	Aug 06, 2027	DS DP		
		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>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		
		8173707*PED	Jan 30, 2024			
		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			

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<u>KETOROLAC TROMETHAMINE; PHENYLEPHRINE HYDROCHLORIDE - OMIIDRIA</u>						
N 205388	001	9399040	Jul 30, 2023	DP		
		9399040*PED	Jan 30, 2024			
		9486406	Oct 23, 2033	DP		
		9486406*PED	Apr 23, 2034			
		9855246	Oct 23, 2033	DP		
<u>L-GLUTAMINE - ENDARI</u>						
N 208587	001				ODE-150	Jul 07, 2024
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	001	RE38551	Mar 17, 2022	DS DP U-1567	I-878	Nov 16, 2023
		RE38551	Mar 17, 2022	DS DP U-2140		
		RE38551	Mar 17, 2022	DS DP U-2989		
		RE38551	Mar 17, 2022	DS DP U-2990		
		RE38551	Mar 17, 2022	DS DP U-2999		
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	002	RE38551	Mar 17, 2022	DS DP U-1567	I-878	Nov 16, 2023
		RE38551	Mar 17, 2022	DS DP U-2140		
		RE38551	Mar 17, 2022	DS DP U-2989		
		RE38551	Mar 17, 2022	DS DP U-2990		
		RE38551	Mar 17, 2022	DS DP U-2999		
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	003	RE38551	Mar 17, 2022	DS DP U-1567	I-878	Nov 16, 2023
		RE38551	Mar 17, 2022	DS DP U-2140		
		RE38551	Mar 17, 2022	DS DP U-2989		
		RE38551	Mar 17, 2022	DS DP U-2990		
		RE38551	Mar 17, 2022	DS DP U-2999		
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	004	RE38551	Mar 17, 2022	DS DP U-1567	I-878	Nov 16, 2023
		RE38551	Mar 17, 2022	DS DP U-2140		
		RE38551	Mar 17, 2022	DS DP U-2989		
		RE38551	Mar 17, 2022	DS DP U-2990		
		RE38551	Mar 17, 2022	DS DP U-2999		
<u>LACOSAMIDE - VIMPAT</u>						
N 022254	001	RE38551	Mar 17, 2022	DS DP U-1565	I-878	Nov 16, 2023
		RE38551	Mar 17, 2022	DS DP U-1568	NPP	Nov 16, 2023
		RE38551	Mar 17, 2022	DS DP U-2989		
		RE38551	Mar 17, 2022	DS DP U-2990		
		RE38551	Mar 17, 2022	DS DP U-2999		
<u>LACOSAMIDE - VIMPAT</u>						
N 022255	001	RE38551	Mar 17, 2022	DS DP U-1567	I-878	Nov 16, 2023
		RE38551	Mar 17, 2022	DS DP U-2140		
		RE38551	Mar 17, 2022	DS DP U-2989		
		RE38551	Mar 17, 2022	DS DP U-2990		
		RE38551	Mar 17, 2022	DS DP U-2999		
<u>LACTITOL - PIZENSY</u>						
N 211281	001	10806743	May 12, 2037	U-1516	NCE	Feb 12, 2025
<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		

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<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		
<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				ODE-156	Sep 15, 2024
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>						
N 022074	002				ODE-156	Sep 15, 2024
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>						
N 022074	003				ODE-156	Sep 15, 2024
<u>LANSOPRAZOLE - LANSOPRAZOLE</u>						
N 208025	001	11077055	Apr 21, 2036	DP		
<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		
<u>LAPATINIB DITOSYLATE - TYKERB</u>						
N 022059	001	8821927	Sep 18, 2029	DS DP		

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<u>LAROTRECTINIB SULFATE - VITRAKVI</u>						
N 210861 001	10005783	Oct 21, 2029		U-2472	NCE	Nov 26, 2023
	10047097	Oct 21, 2029		U-2474	ODE-215	Nov 26, 2025
	10172861	Nov 16, 2035	DS DP		ODE-220	Nov 26, 2025
	10285993	Nov 16, 2035		U-2470	ODE-221	Nov 26, 2025
	10774085	Oct 21, 2029		U-2470		
	10799505	Aug 15, 2036	DS DP			
	10813936	Nov 16, 2035		U-2987		
	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-2475		
<u>LAROTRECTINIB SULFATE - VITRAKVI</u>						
N 210861 002	10005783	Oct 21, 2029		U-2472	NCE	Nov 26, 2023
	10047097	Oct 21, 2029		U-2474	ODE-215	Nov 26, 2025
	10172861	Nov 16, 2035	DS DP		ODE-220	Nov 26, 2025
	10285993	Nov 16, 2035		U-2470	ODE-221	Nov 26, 2025
	10774085	Oct 21, 2029		U-2470		
	10799505	Aug 15, 2036	DS DP			
	10813936	Nov 16, 2035		U-2987		
	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-2475		
<u>LAROTRECTINIB SULFATE - VITRAKVI</u>						
N 211710 001	10005783	Oct 21, 2029		U-2472	NCE	Nov 26, 2023
	10045991	Apr 04, 2037		U-2473	ODE-215	Nov 26, 2025
	10047097	Oct 21, 2029		U-2474	ODE-220	Nov 26, 2025
	10137127	Apr 04, 2037		DP	ODE-221	Nov 26, 2025
	10172861	Nov 16, 2035	DS			
	10668072	Apr 04, 2037		DP		
	10774085	Oct 21, 2029		U-2470		
	10799505	Aug 15, 2036	DS			
	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>LASMIDITAN SUCCINATE - REYVOW</u>						
N 211280 001	11053214	Dec 05, 2037	DS DP	U-1719	NCE	Jan 31, 2025
	7423050	Apr 06, 2025	DS DP	U-1719		
	8748459	Mar 27, 2023		U-1719		
<u>LASMIDITAN SUCCINATE - REYVOW</u>						
N 211280 002	11053214	Dec 05, 2037	DS DP	U-1719	NCE	Jan 31, 2025
	7423050	Apr 06, 2025	DS DP	U-1719		
	8748459	Mar 27, 2023		U-1719		
<u>LASMIDITAN SUCCINATE - REYVOW</u>						
N 211280 003	11053214	Dec 05, 2037	DS DP	U-1719	NCE	Jan 31, 2025
	7423050	Apr 06, 2025	DS DP	U-1719		
	8748459	Mar 27, 2023		U-1719		
<u>LATANOPROST - XELPROS</u>						
N 206185 001	9539262	Oct 15, 2028		U-2400		
	9629852	Sep 12, 2029		DP		
<u>LATANOPROST; NETARSUDIL DIMESYLATE - ROCKLATAN</u>						
N 208259 001	10174017	Jan 27, 2030	DS DP	U-1524	NC	Mar 12, 2022
	10532993	Jul 11, 2026		U-1524	NCE	Dec 18, 2022
	10588901	Mar 14, 2034	DS DP	U-1524		
	10654844	Jan 27, 2030	DS DP	U-1524		
	10882840	Jul 11, 2026		U-1524		

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<u>LATANOPROST; NETARSUDIL DIMESYLATE - ROCKLATAN</u>						
N 208259	001	11021456	Jul 11, 2026	U-1524		
		11028081	Jan 27, 2030	U-1524		
		11185538	Mar 14, 2034	DP		
		11197853	Mar 14, 2034	DP		
		8394826	Nov 10, 2030	DS DP U-1524		
		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		
		9993470	Mar 14, 2034	DS DP U-1524		
<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-177	Nov 15, 2022
		10039779	Jan 30, 2034	DS DP U-2370	ODE*	Aug 28, 2026
		10039779*PED	Jul 30, 2034		ODE*	Aug 28, 2026
		10456414	Sep 14, 2032	DP	ODE*	Aug 28, 2026
		7964580	Mar 26, 2029	DS DP U-1470	ODE-136	Apr 07, 2024
		7964580*PED	Sep 26, 2029		PED	Oct 07, 2024
		8088368	May 12, 2030	DS DP		
		8088368*PED	Nov 12, 2030			
		8273341	May 12, 2030	U-1470		
		8273341*PED	Nov 12, 2030			
		8334270	Mar 21, 2028	DS DP U-1470		
		8334270*PED	Sep 21, 2028			
		8580765	Mar 21, 2028	DS DP U-1470		
		8580765*PED	Sep 21, 2028			
		8618076	Dec 11, 2030	DS DP U-1470		
		8618076*PED	Jun 11, 2031			
		8633309	Mar 26, 2029	DS DP U-1470		
		8633309*PED	Sep 26, 2029			
		8735372	Mar 21, 2028	U-1470		
		8735372*PED	Sep 21, 2028			
		8822430	May 12, 2030	DS DP U-1470		
		8822430*PED	Nov 12, 2030			
		8841278	May 12, 2030	DP U-1470		
		8841278*PED	Nov 12, 2030			
		8889159	Mar 26, 2029	DP U-1470		
		8889159*PED	Sep 26, 2029			
		9085573	Mar 21, 2028	DS DP U-1470		
		9085573*PED	Sep 21, 2028			
		9284342	Sep 13, 2030	DS DP U-1470		
		9284342*PED	Mar 13, 2031			
		9393256	Sep 14, 2032	U-1470		
		9393256*PED	Mar 14, 2033			
		9511056	May 12, 2030	DP U-1470		
		9511056*PED	Nov 12, 2030			
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 205834	002	10039779	Jan 30, 2034	DS DP U-1470	D-177	Nov 15, 2022
		10039779*PED	Jul 30, 2034		ODE*	Apr 07, 2024
		10456414	Sep 14, 2032	DP	ODE*	Aug 28, 2026
		7964580	Mar 26, 2029	DS DP U-1470	ODE*	Aug 28, 2026
		7964580*PED	Sep 26, 2029		ODE*	Aug 28, 2026
		8088368	May 12, 2030	DS DP	PED	Oct 07, 2024
		8088368*PED	Nov 12, 2030			
		8273341	May 12, 2030	U-1470		
		8273341*PED	Nov 12, 2030			
		8334270	Mar 21, 2028	DS DP U-1470		
		8334270*PED	Sep 21, 2028			
		8580765	Mar 21, 2028	DS DP U-1470		
		8580765*PED	Sep 21, 2028			
		8618076	Dec 11, 2030	DS DP U-1470		
		8618076*PED	Jun 11, 2031			
		8633309	Mar 26, 2029	DS DP U-1470		

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<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 205834	002	8633309*PED	Sep 26, 2029			
		8735372	Mar 21, 2028	U-1470		
		8735372*PED	Sep 21, 2028			
		8822430	May 12, 2030	DS DP U-1470		
		8822430*PED	Nov 12, 2030			
		8841278	May 12, 2030	DP U-1470		
		8841278*PED	Nov 12, 2030			
		8889159	Mar 26, 2029	DP U-1470		
		8889159*PED	Sep 26, 2029			
		9085573	Mar 21, 2028	DS DP U-1470		
		9085573*PED	Sep 21, 2028			
		9284342	Sep 13, 2030	DS DP U-1470		
		9284342*PED	Mar 13, 2031			
		9393256	Sep 14, 2032	U-1470		
		9393256*PED	Mar 14, 2033			
		9511056	May 12, 2030	U-1470		
		9511056*PED	Nov 12, 2030			
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 212477	001	10456414	Sep 14, 2032	DP	D-177	Nov 15, 2022
		7964580	Mar 26, 2029	DS DP U-1470	ODE-262	Aug 28, 2026
		7964580*PED	Sep 26, 2029		ODE-263	Aug 28, 2026
		8088368	May 12, 2030	DS DP	ODE-264	Aug 28, 2026
		8088368*PED	Nov 12, 2030			
		8273341	May 12, 2030	U-1470		
		8273341*PED	Nov 12, 2030			
		8334270	Mar 21, 2028	DS DP U-1470		
		8334270*PED	Sep 21, 2028			
		8580765	Mar 21, 2028	DS DP U-1470		
		8580765*PED	Sep 21, 2028			
		8618076	Dec 11, 2030	DS DP U-1470		
		8618076*PED	Jun 11, 2031			
		8633309	Mar 26, 2029	DS DP U-1470		
		8633309*PED	Sep 26, 2029			
		8735372	Mar 21, 2028	U-1470		
		8735372*PED	Sep 21, 2028			
		8822430	May 12, 2030	DS DP U-1470		
		8822430*PED	Nov 12, 2030			
		8841278	May 12, 2030	DP U-1470		
		8841278*PED	Nov 12, 2030			
		8889159	Mar 26, 2029	DP U-1470		
		8889159*PED	Sep 26, 2029			
		9085573	Mar 21, 2028	DS DP U-1470		
		9085573*PED	Sep 21, 2028			
		9284342	Sep 13, 2030	DS DP U-1470		
		9284342*PED	Mar 13, 2031			
		9393256	Sep 14, 2032	U-1470		
		9393256*PED	Mar 14, 2033			
		9511056	May 12, 2030	U-1470		
		9511056*PED	Nov 12, 2030			
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 212477	002	10456414	Sep 14, 2032	DP	D-177	Nov 15, 2022
		7964580	Mar 26, 2029	DS DP U-1470	ODE-262	Aug 28, 2026
		7964580*PED	Sep 26, 2029		ODE-263	Aug 28, 2026
		8088368	May 12, 2030	DS DP	ODE-264	Aug 28, 2026
		8088368*PED	Nov 12, 2030			
		8273341	May 12, 2030	U-1470		
		8273341*PED	Nov 12, 2030			
		8334270	Mar 21, 2028	DS DP U-1470		
		8334270*PED	Sep 21, 2028			
		8580765	Mar 21, 2028	DS DP U-1470		
		8580765*PED	Sep 21, 2028			
		8618076	Dec 11, 2030	DS DP U-1470		
		8618076*PED	Jun 11, 2031			
		8633309	Mar 26, 2029	DS DP U-1470		
		8633309*PED	Sep 26, 2029			
		8735372	Mar 21, 2028	U-1470		
		8735372*PED	Sep 21, 2028			

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<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 212477	002 8822430	May 12, 2030	DS DP U-1470			
	8822430*PED	Nov 12, 2030				
	8841278	May 12, 2030	DP U-1470			
	8841278*PED	Nov 12, 2030				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9393256	Sep 14, 2032	U-1470			
	9393256*PED	Mar 14, 2033				
	9511056	May 12, 2030	U-1470			
	9511056*PED	Nov 12, 2030				
<u>LEFAMULIN ACETATE - XENLETA</u>						
N 211672	001 8071643	Jan 16, 2029	DS DP		NCE	Aug 19, 2024
	8153689	Mar 19, 2028	DS DP		GAIN	Aug 19, 2029
	9120727	May 23, 2031	DS DP			
<u>LEFAMULIN ACETATE - XENLETA</u>						
N 211673	001 8071643	Jan 16, 2029	DS DP		NCE	Aug 19, 2024
	8153689	Mar 19, 2028	DS DP		GAIN	Aug 19, 2029
<u>LEMBOREXANT - DAYVIGO</u>						
N 212028	001 10188652	Oct 21, 2035	DP U-2791		NCE	Apr 07, 2025
	8268848	Sep 20, 2031	DS DP U-2791			
<u>LEMBOREXANT - DAYVIGO</u>						
N 212028	002 10188652	Oct 21, 2035	DP U-2791		NCE	Apr 07, 2025
	8268848	Sep 20, 2031	DS DP U-2791			
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880	001 7189740	Apr 11, 2023		U-1982	I-796	May 28, 2022
	7465800	Apr 27, 2027	DS DP		I-797	May 28, 2022
	7468363	Oct 07, 2023		U-1983	ODE-131	Feb 22, 2024
	7468363	Oct 07, 2023		U-2550	ODE-241	May 28, 2026
	7468363	Oct 07, 2023		U-2551	ODE-245	May 28, 2026
	7855217	Nov 24, 2024	DS DP		ODE-88	Feb 17, 2022
	7968569	Oct 07, 2023		U-1984		
	8404717	Apr 11, 2023		U-1982		
	8492406	Oct 07, 2023		U-2550		
	8530498	May 15, 2023		U-1984		
	8648095	May 15, 2023		U-1984		
	8741929	Mar 08, 2028		U-1983		
	9056120	Apr 11, 2023		U-1982		
	9101621	May 15, 2023		U-1985		
	9101622	May 15, 2023		U-1986		
	9155730	May 15, 2023		U-2550		
	9393238	May 15, 2023		U-2550		
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880	002 7189740	Apr 11, 2023		U-1982	I-796	May 28, 2022
	7465800	Apr 27, 2027	DS DP		I-797	May 28, 2022
	7468363	Oct 07, 2023		U-1983	ODE-131	Feb 22, 2024
	7468363	Oct 07, 2023		U-2550	ODE-241	May 28, 2026
	7468363	Oct 07, 2023		U-2551	ODE-245	May 28, 2026
	7855217	Nov 24, 2024	DS DP		ODE-88	Feb 17, 2022
	7968569	Oct 07, 2023		U-1984		
	8404717	Apr 11, 2023		U-1982		
	8492406	Oct 07, 2023		U-2550		
	8530498	May 15, 2023		U-1984		
	8648095	May 15, 2023		U-1984		
	8741929	Mar 08, 2028		U-1983		
	9056120	Apr 11, 2023		U-1982		
	9101621	May 15, 2023		U-1985		
	9101622	May 15, 2023		U-1986		
	9155730	May 15, 2023		U-2550		
	9393238	May 15, 2023		U-2550		

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<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 003	7189740	Apr 11, 2023	U-1982		I-796	May 28, 2022
	7465800	Apr 27, 2027	DS DP		I-797	May 28, 2022
	7468363	Oct 07, 2023	U-1983		ODE-131	Feb 22, 2024
	7468363	Oct 07, 2023	U-2550		ODE-241	May 28, 2026
	7468363	Oct 07, 2023	U-2551		ODE-245	May 28, 2026
	7855217	Nov 24, 2024	DS DP		ODE-88	Feb 17, 2022
	7968569	Oct 07, 2023	U-1984			
	8404717	Apr 11, 2023	U-1982			
	8492406	Oct 07, 2023	U-2550			
	8530498	May 15, 2023	U-1984			
	8648095	May 15, 2023	U-1984			
	8741929	Mar 08, 2028	U-1983			
	9056120	Apr 11, 2023	U-1982			
	9101621	May 15, 2023	U-1985			
	9101622	May 15, 2023	U-1986			
	9155730	May 15, 2023	U-2550			
	9393238	May 15, 2023	U-2550			
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 004	7189740	Apr 11, 2023	U-1982		I-796	May 28, 2022
	7465800	Apr 27, 2027	DS DP		I-797	May 28, 2022
	7468363	Oct 07, 2023	U-1983		ODE-131	Feb 22, 2024
	7468363	Oct 07, 2023	U-2550		ODE-241	May 28, 2026
	7468363	Oct 07, 2023	U-2551		ODE-245	May 28, 2026
	7855217	Nov 24, 2024	DS DP		ODE-88	Feb 17, 2022
	7968569	Oct 07, 2023	U-1984			
	8404717	Apr 11, 2023	U-1982			
	8492406	Oct 07, 2023	U-2550			
	8530498	May 15, 2023	U-1984			
	8648095	May 15, 2023	U-1984			
	8741929	Mar 08, 2028	U-1983			
	9056120	Apr 11, 2023	U-1982			
	9101621	May 15, 2023	U-1985			
	9101622	May 15, 2023	U-1986			
	9155730	May 15, 2023	U-2550			
	9393238	May 15, 2023	U-2550			
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 005	7189740	Apr 11, 2023	U-1982		I-796	May 28, 2022
	7465800	Apr 27, 2027	DS DP		I-797	May 28, 2022
	7468363	Oct 07, 2023	U-1983		ODE-131	Feb 22, 2024
	7468363	Oct 07, 2023	U-2550		ODE-241	May 28, 2026
	7468363	Oct 07, 2023	U-2551		ODE-245	May 28, 2026
	7855217	Nov 24, 2024	DS DP		ODE-88	Feb 17, 2022
	7968569	Oct 07, 2023	U-1984			
	8404717	Apr 11, 2023	U-1982			
	8492406	Oct 07, 2023	U-2550			
	8530498	May 15, 2023	U-1984			
	8648095	May 15, 2023	U-1984			
	8741929	Mar 08, 2028	U-1983			
	9056120	Apr 11, 2023	U-1982			
	9101621	May 15, 2023	U-1985			
	9101622	May 15, 2023	U-1986			
	9155730	May 15, 2023	U-2550			
	9393238	May 15, 2023	U-2550			
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 006	7189740	Apr 11, 2023	U-1982		I-796	May 28, 2022
	7465800	Apr 27, 2027	DS DP		I-797	May 28, 2022
	7468363	Oct 07, 2023	U-1983		ODE-131	Feb 22, 2024
	7468363	Oct 07, 2023	U-2550		ODE-241	May 28, 2026
	7468363	Oct 07, 2023	U-2551		ODE-245	May 28, 2026
	7855217	Nov 24, 2024	DS DP		ODE-88	Feb 17, 2022
	7968569	Oct 07, 2023	U-1984			
	8404717	Apr 11, 2023	U-1982			
	8492406	Oct 07, 2023	U-2550			
	8530498	May 15, 2023	U-1984			
	8648095	May 15, 2023	U-1984			
	8741929	Mar 08, 2028	U-1983			
	9056120	Apr 11, 2023	U-1982			

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<u>LENALIDOMIDE - REVLIMID</u>						
N 021880	006	9101621	May 15, 2023	U-1985		
		9101622	May 15, 2023	U-1986		
		9155730	May 15, 2023	U-2550		
		9393238	May 15, 2023	U-2550		
<u>LENVATINIB MESYLATE - LENVIMA</u>						
N 206947	001	10259791	Aug 26, 2035	DS	I-807	Sep 17, 2022
		10407393	Aug 26, 2035	DS	I-868	Aug 10, 2024
		11186547	Aug 26, 2035	DS	M-269	Jul 21, 2024
		7253286	Oct 24, 2025	DS DP	ODE-196	Aug 15, 2025
		7612208	Sep 19, 2026	DS DP	ODE-87	Feb 13, 2022
		9006256	Jul 27, 2027	U-1695		
<u>LENVATINIB MESYLATE - LENVIMA</u>						
N 206947	002	10259791	Aug 26, 2035	DS	I-807	Sep 17, 2022
		10407393	Aug 26, 2035	DS	I-868	Aug 10, 2024
		11186547	Aug 26, 2035	DS	M-269	Jul 21, 2024
		7253286	Oct 24, 2025	DS DP	ODE-196	Aug 15, 2025
		7612208	Sep 19, 2026	DS DP	ODE-87	Feb 13, 2022
		9006256	Jul 27, 2027	U-1695		
<u>LESINURAD - ZURAMPIC</u>						
N 207988	001	10183012	Nov 26, 2028	U-2311		
		8003681	Aug 25, 2025	DS		
		8084483	Aug 17, 2029	U-1801		
		8283369	Nov 26, 2028	U-1802		
		8283369	Nov 26, 2028	U-1804		
		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	8513255	May 22, 2024	DS DP	NCE	Nov 08, 2022
		RE46791	May 22, 2024	DS DP	ODE-165	Nov 08, 2024
<u>LETERMOVIR - PREVYMIS</u>						
N 209939	002	8513255	May 22, 2024	DS DP	NCE	Nov 08, 2022
		RE46791	May 22, 2024	DS DP	ODE-165	Nov 08, 2024
<u>LETERMOVIR - PREVYMIS</u>						
N 209940	001	10603384	Feb 28, 2033	DP	NCE	Nov 08, 2022
		8513255	May 22, 2024	DS DP	ODE-165	Nov 08, 2024
		RE46791	May 22, 2024	DS DP		
<u>LETERMOVIR - PREVYMIS</u>						
N 209940	002	10603384	Feb 28, 2033	DP	NCE	Nov 08, 2022
		8513255	May 22, 2024	DS DP	ODE-165	Nov 08, 2024
		RE46791	May 22, 2024	DS DP		
<u>LETROZOLE; RIBOCICLIB SUCCINATE - KISOALI FEMARA CO-PACK (COPACKAGED)</u>						
N 209935	001	10799506	Apr 14, 2036	DP	NCE	Mar 13, 2022
		8324225	Jun 17, 2028	DS DP		
		8415355	Aug 21, 2029	DS DP		
		8685980	May 25, 2030	DS DP		
		8962630	Dec 09, 2029	U-2505		
		8962630	Dec 09, 2029	U-3264		
		9193732	Nov 09, 2031	DS DP		
		9416136	Aug 20, 2029	U-2505		
		9416136	Aug 20, 2029	U-3264		
		9868739	Nov 09, 2031	U-2505		
		9868739	Nov 09, 2031	U-3264		
<u>LEUPROLIDE ACETATE - LUPRON DEPOT</u>						
N 020517	003	8815801	Jun 28, 2022	DP		
		8921326	Feb 05, 2031	DP U-1666		

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<u>LEUPROLIDE ACETATE - ELIGARD KIT</u>						
N 021379	001 8470359	Oct 15, 2023	DS DP U-621			
<u>LEUPROLIDE ACETATE - ELIGARD KIT</u>						
N 021488	001 8470359	Oct 15, 2023	DS DP U-621			
<u>LEUPROLIDE ACETATE - ELIGARD KIT</u>						
N 021731	001 8470359	Oct 15, 2023	DS DP U-621			
<u>LEUPROLIDE ACETATE - FENSOLVI KIT</u>						
N 213150	001 8470359	Oct 15, 2023	DS DP U-2940		I-829	May 01, 2023
<u>LEUPROLIDE MESYLATE - CAMCEVI KIT</u>						
N 211488	001 10646572	Jan 16, 2027	DP		NP	May 25, 2024
	9572857	Jan 16, 2027	DP			
	9744207	Jan 16, 2027	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	Oct 31, 2027	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	Oct 31, 2027	DP			
	8535717	Feb 22, 2026	DP			
<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	001 11160786	Mar 14, 2034	DP			
	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 11160786	Mar 14, 2034	DP			
	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	

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<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	003	11160786	Mar 14, 2034	DP		
		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	11160786	Mar 14, 2034	DP		
		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>LEVOCETIRIZINE DIHYDROCHLORIDE - XYZAL ALLERGY 24HR</u>						
N 209090	001	8633194	Oct 16, 2027	DP		
<u>LEVODOPA - INBRIJA</u>						
N 209184	001	7182961	Feb 22, 2024	DP		
		7384649	Nov 20, 2022	DP		
		8404276	Mar 19, 2023		U-2484	
		8545878	Nov 16, 2032	DP		
		8586093	Mar 19, 2023		U-2484	
		8685442	Nov 16, 2032	DP		
		8945612	Nov 16, 2032	DP		
		9155699	Mar 19, 2023	DP		
		9393210	Nov 16, 2032	DP		
		RE43711	Feb 03, 2029		U-2484	
<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	Jan 11, 2026		U-839	
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	002	8481598	Mar 02, 2031		U-839	
		8865937	May 23, 2032	DS DP		
		RE43879	Jan 11, 2026		U-839	
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	003	8481598	Mar 02, 2031		U-839	
		8865937	May 23, 2032	DS DP		
		RE43879	Jan 11, 2026		U-839	
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	004	8481598	Mar 02, 2031		U-839	
		8865937	May 23, 2032	DS DP		
		RE43879	Jan 11, 2026		U-839	
<u>LEVONORGESTREL - MIRENA</u>						
N 021225	001	10561524	Sep 16, 2029		U-2948	
		10987244	Apr 01, 2031	DP		
		9615965	Sep 16, 2029	DP	U-2003	
		9668912	Apr 01, 2031	DP		
<u>LEVONORGESTREL - SKYLA</u>						
N 203159	001	10561524	Sep 16, 2029		U-2948	
		10987244	Apr 01, 2031	DP		
		7252839	Nov 13, 2023	DP		
		9615965	Sep 16, 2029	DP	U-2003	

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<u>LEVONORGESTREL - SKYLA</u>						
N 203159	001 9668912	Apr 01, 2031	DP			
<u>LEVONORGESTREL - LILETTA</u>						
N 206229	001 10028858	Mar 22, 2034	DP	U-2348		
	11090186	Oct 24, 2033		U-2348		
<u>LEVONORGESTREL - KYLEENA</u>						
N 208224	001 10561524	Sep 16, 2029			U-2948	
	10987244	Apr 01, 2031	DP			
	7252839	Nov 13, 2023	DP			
	9615965	Sep 16, 2029	DP	U-2003		
	9668912	Apr 01, 2031	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	003 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	004 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	005 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	006 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	007 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	008 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	009 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	010 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	011 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	012 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<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	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N 021301	003 6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023			U-759	

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<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 - TIROSINT</u>						
N 021924 002	7691411	Mar 14, 2024				
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924 003	7691411	Mar 14, 2024				
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924 004	7691411	Mar 14, 2024				
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924 005	7691411	Mar 14, 2024				
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924 006	7691411	Mar 14, 2024				
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924 007	7691411	Mar 14, 2024				
	7723390	Mar 14, 2024	DP			

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<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	011	7691411	Mar 14, 2024	DP		
		7723390	Mar 14, 2024	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	012	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		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	001	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	002	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	003	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	004	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	005	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	006	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	007	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	008	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		

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<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	009	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	010	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	011	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	012	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	013	11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	014	11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	015	11096913	Feb 28, 2037	DP		
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 210632	001	10398669	Dec 01, 2036	DP		
		11135190	Dec 01, 2036	DP		
		9782376	Dec 01, 2036	DP		
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 210632	002	10398669	Dec 01, 2036	DP		
		11135190	Dec 01, 2036	DP		
		9782376	Dec 01, 2036	DP		
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 210632	003	10398669	Dec 01, 2036	DP		
		11135190	Dec 01, 2036	DP		
		9782376	Dec 01, 2036	DP		
<u>LEVOTHYROXINE SODIUM - THYQUIDITY</u>						
N 214047	001	9050307	Aug 06, 2031	DP		
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 214253	001	11154498	Jul 20, 2036	DP		
<u>LIDOCAINE - ZTLIDO</u>						
N 207962	001	10765640	May 10, 2031	DP		
		10765749	May 10, 2031	DP		
		9283174	May 10, 2031	DP		
		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 - PLIAGLIS</u>						
N 021717	001	10350180	Jan 14, 2031	DP		
		10603293	Jan 14, 2031	DP		
		10751305	Jan 14, 2031	DP		
<u>LIFITEGRAST - XIIDRA</u>						
N 208073	001	10124000	Nov 05, 2024		U-1900	
		11058677	Dec 18, 2033	DP		
		7314938	Mar 10, 2025	DS	DP	
		7745460	Nov 05, 2024	DS	DP	U-1880
		7790743	Nov 05, 2024		U-1880	

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<u>LIFITEGRAST - XIIDRA</u>						
N 208073	001	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	Oct 30, 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	Oct 30, 2031	DP		
		8933030	Feb 17, 2031	DP		
		9708371	Aug 16, 2033	DP	U-1515	
<u>LINACLOTIDE - LINZESS</u>						
N 202811	003	10675325	Aug 11, 2031	DP		
		10702576	Aug 11, 2031		U-1516	
		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-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	M-252 Mar 30, 2023
		11033552	May 04, 2027		DP	M-258 Jul 03, 2022
		7407955	May 02, 2025	DS DP		
		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	Y
		8846695	Jun 04, 2030		U-1503	Y
		8853156	Mar 05, 2031		U-1642	
		8883805	Nov 26, 2025	DP		
		9486526	Aug 05, 2029		U-1915	

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<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 001	10022379	Apr 02, 2029	U-2339		M-252	Mar 30, 2023
	10973827	Apr 02, 2029	DP		M-258	Jul 03, 2022
	7407955	May 02, 2025	DS DP			
	8119648	Aug 12, 2023	U-802			
	8178541	Aug 12, 2023	DP U-775			
	8673927	May 04, 2027	U-1503	Y		
	8846695	Jun 04, 2030	U-1503			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 002	10022379	Apr 02, 2029	U-2339		M-252	Mar 30, 2023
	10973827	Apr 02, 2029	DP		M-258	Jul 03, 2022
	7407955	May 02, 2025	DS DP			
	8119648	Aug 12, 2023	U-802			
	8178541	Aug 12, 2023	DP U-775			
	8673927	May 04, 2027	U-1503	Y		
	8846695	Jun 04, 2030	U-1503			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 003	10022379	Apr 02, 2029	U-2339		M-252	Mar 30, 2023
	10973827	Apr 02, 2029	DP		M-258	Jul 03, 2022
	7407955	May 02, 2025	DS DP			
	8119648	Aug 12, 2023	U-802			
	8178541	Aug 12, 2023	DP U-775			
	8673927	May 04, 2027	U-1503	Y		
	8846695	Jun 04, 2030	U-1503	Y		
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</u>						
N 208026 001	10022379	Apr 02, 2029	U-2339		M-252	Mar 30, 2023
	7407955	May 02, 2025	DS DP		M-258	Jul 03, 2022
	8119648	Aug 12, 2023	U-802			
	8178541	Aug 12, 2023	DP U-1853			
	8673927	May 04, 2027	U-1503	Y		
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1503	Y		
	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		M-252	Mar 30, 2023
	7407955	May 02, 2025	DS DP		M-258	Jul 03, 2022
	8119648	Aug 12, 2023	U-802			
	8178541	Aug 12, 2023	DP U-1853			
	8673927	May 04, 2027	U-1503	Y		
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1503	Y		
	9415016	Apr 02, 2029	DP			
	9555001	Mar 06, 2033	DP U-1967			
	9555001	Mar 06, 2033	DP U-1968			
<u>LIRAGLUTIDE RECOMBINANT - VICTOZA</u>						
N 022341 001	6268343	Aug 22, 2022	DS DP U-968		NPP	Jun 17, 2022
	6268343*PED	Feb 22, 2023			PED	Dec 17, 2022
	7762994	May 23, 2024	DP			
	7762994*PED	Nov 23, 2024				
	8114833	Aug 13, 2025	DS DP			
	8114833*PED	Feb 13, 2026				
	8579869	Jun 30, 2023	DP			
	8579869*PED	Dec 30, 2023				

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<u>LIRAGLUTIDE RECOMBINANT - VICTOZA</u>						
N 022341 001	8846618	Jun 27, 2022	DP			
	8846618*PED	Dec 27, 2022				
	9265893	Sep 23, 2032	DP			
	9265893*PED	Mar 23, 2033				
	9968659	Jan 09, 2037		U-2313		
	9968659*PED	Jul 09, 2037				
<u>LIRAGLUTIDE RECOMBINANT - SAXENDA</u>						
N 206321 001	10220155	Jul 17, 2026	DP		NPP	Dec 04, 2023
	10220155*PED	Jan 17, 2027				
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	11097063	Jul 17, 2026	DP			
	6268343	Aug 22, 2022	DS DP	U-1255		
	6268343*PED	Feb 22, 2023				
	6899699*PED	Jul 01, 2022				
	7762994	May 23, 2024	DP			
	7762994*PED	Nov 23, 2024				
	8114833	Aug 13, 2025	DP			
	8114833*PED	Feb 13, 2026				
	8579869	Jun 30, 2023	DP			
	8579869*PED	Dec 30, 2023				
	8672898	Jan 02, 2022	DP			
	8672898*PED	Jul 02, 2022				
	8684969	Oct 20, 2025	DP			
	8684969*PED	Apr 20, 2026				
	8846618	Jun 27, 2022	DP			
	8846618*PED	Dec 27, 2022				
	8920383	Jul 17, 2026	DP			
	8920383*PED	Jan 17, 2027				
	9108002	Jan 26, 2026	DP			
	9108002*PED	Jul 26, 2026				
	9132239	Feb 01, 2032	DP			
	9132239*PED	Aug 01, 2032				
	9457154	Sep 27, 2027	DP			
	9457154*PED	Mar 27, 2028				
	9486588	Jan 02, 2022	DP			
	9486588*PED	Jul 02, 2022				
	9616180	Jan 20, 2026	DP			
	9616180*PED	Jul 20, 2026				
	9687611	Feb 27, 2027	DP			
	9687611*PED	Aug 27, 2027				
	9775953	Jul 17, 2026	DP			
	9775953*PED	Jan 17, 2027				
	9861757	Jan 20, 2026	DP			
	9861757*PED	Jul 20, 2026				
	9968659	Jan 09, 2037		U-2438		
	9968659*PED	Jul 09, 2037				
	RE46363	Aug 03, 2026	DP			
	RE46363*PED	Feb 03, 2027				
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 021977 001	7105486	Feb 24, 2023		U-727		
	7105486*PED	Aug 24, 2023				
	7223735	Feb 24, 2023	DP			
	7223735*PED	Aug 24, 2023				
	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		
	7671031*PED	Aug 24, 2023				
	7674774	Feb 24, 2023	DP	U-842		
	7674774*PED	Aug 24, 2023				
	7678770	Feb 24, 2023		U-842		
	7678770*PED	Aug 24, 2023				
	7678771	Feb 24, 2023	DP	U-842		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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 001	7678771*PED	Aug 24, 2023				
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-842			
	7687467*PED	Aug 24, 2023				
	7700561	Feb 24, 2023	DP			
	7700561*PED	Aug 24, 2023				
	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			
	7105486*PED	Aug 24, 2023				
	7223735	Feb 24, 2023	DP			
	7223735*PED	Aug 24, 2023				
	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			
	7671031*PED	Aug 24, 2023				
	7674774	Feb 24, 2023	DP U-842			
	7674774*PED	Aug 24, 2023				
	7678770	Feb 24, 2023	U-842			
	7678770*PED	Aug 24, 2023				
	7678771	Feb 24, 2023	DP U-842			
	7678771*PED	Aug 24, 2023				
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-842			
	7687467*PED	Aug 24, 2023				
	7700561	Feb 24, 2023	DP			
	7700561*PED	Aug 24, 2023				
	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 003	7105486	Feb 24, 2023	U-727			
	7105486*PED	Aug 24, 2023				
	7223735	Feb 24, 2023	DP			
	7223735*PED	Aug 24, 2023				
	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			
	7671031*PED	Aug 24, 2023				
	7674774	Feb 24, 2023	DP U-842			
	7674774*PED	Aug 24, 2023				
	7678770	Feb 24, 2023	U-842			
	7678770*PED	Aug 24, 2023				
	7678771	Feb 24, 2023	DP U-842			
	7678771*PED	Aug 24, 2023				
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-842			
	7687467*PED	Aug 24, 2023				
	7700561	Feb 24, 2023	DP			
	7700561*PED	Aug 24, 2023				
	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			
	7105486	Feb 24, 2023	U-842			
	7105486*PED	Aug 24, 2023				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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 004	7223735	Feb 24, 2023	DP			
	7223735*PED	Aug 24, 2023				
	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	
	7671031*PED	Aug 24, 2023				
	7674774	Feb 24, 2023	DP		U-842	
	7674774*PED	Aug 24, 2023				
	7678770	Feb 24, 2023			U-842	
	7678770*PED	Aug 24, 2023				
	7678771	Feb 24, 2023	DP		U-842	
	7678771*PED	Aug 24, 2023				
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP		U-842	
	7687467*PED	Aug 24, 2023				
	7700561	Feb 24, 2023	DP			
	7700561*PED	Aug 24, 2023				
	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	
	7105486*PED	Aug 24, 2023				
	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	
	7671031*PED	Aug 24, 2023				
	7674774	Feb 24, 2023	DP		U-842	
	7674774*PED	Aug 24, 2023				
	7678770	Feb 24, 2023			U-842	
	7678770*PED	Aug 24, 2023				
	7678771	Feb 24, 2023	DP		U-842	
	7678771*PED	Aug 24, 2023				
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP		U-842	
	7687467*PED	Aug 24, 2023				
	7700561	Feb 24, 2023	DP			
	7700561*PED	Aug 24, 2023				
	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	
	7105486	Feb 24, 2023			U-842	
	7105486*PED	Aug 24, 2023				
	7223735	Feb 24, 2023	DP			
	7223735*PED	Aug 24, 2023				
	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	
	7671031*PED	Aug 24, 2023				
	7674774	Feb 24, 2023	DP		U-842	
	7674774*PED	Aug 24, 2023				
	7678770	Feb 24, 2023			U-842	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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	7678770*PED	Aug 24, 2023			
		7678771	Feb 24, 2023	DP	U-842	
		7678771*PED	Aug 24, 2023			
		7687466	Feb 24, 2023	DP		
		7687467	Feb 24, 2023	DP	U-842	
		7687467*PED	Aug 24, 2023			
		7700561	Feb 24, 2023	DP		
		7700561*PED	Aug 24, 2023			
		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		
		7223735*PED	Aug 24, 2023			
		7655630	Feb 24, 2023	DS		
		7655630*PED	Aug 24, 2023			
		7659253	Feb 24, 2023	DS	DP U-727	
		7659253*PED	Aug 24, 2023			
		7659254	Feb 24, 2023		U-1034	
		7659254*PED	Aug 24, 2023			
		7662787	Feb 24, 2023	DS		
		7662787*PED	Aug 24, 2023			
		7662788	Feb 24, 2023		U-727	
		7662788*PED	Aug 24, 2023			
		7671030	Feb 24, 2023	DP	U-727	
		7671030*PED	Aug 24, 2023			
		7671031	Feb 24, 2023		U-727	
		7671031*PED	Aug 24, 2023			
		7674774	Feb 24, 2023	DP	U-842	
		7674774*PED	Aug 24, 2023			
		7678770	Feb 24, 2023		U-842	
		7678770*PED	Aug 24, 2023			
		7678771	Feb 24, 2023	DP		
		7678771*PED	Aug 24, 2023			
		7687466	Feb 24, 2023	DP		
		7687466*PED	Aug 24, 2023			
		7687467	Feb 24, 2023	DP	U-842	
		7687467*PED	Aug 24, 2023			
		7700561	Feb 24, 2023	DP		
		7700561*PED	Aug 24, 2023			
		7713936	Feb 24, 2023		U-727	
		7713936*PED	Aug 24, 2023			
		7718619	Feb 24, 2023	DP	U-842	
		7718619*PED	Aug 24, 2023			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510	001	7105486	Feb 24, 2023		U-727	
		7105486*PED	Aug 24, 2023			
		7223735	Feb 24, 2023	DP		
		7223735*PED	Aug 24, 2023			
		7655630	Feb 24, 2023	DS	DP	
		7655630*PED	Aug 24, 2023			
		7659253	Feb 24, 2023	DS	DP U-727	
		7659253*PED	Aug 24, 2023			
		7659254	Feb 24, 2023		U-727	
		7659254*PED	Aug 24, 2023			
		7662787	Feb 24, 2023	DS		
		7662787*PED	Aug 24, 2023			
		7662788	Feb 24, 2023		U-727	
		7662788*PED	Aug 24, 2023			
		7671030	Feb 24, 2023	DP	U-727	
		7671030*PED	Aug 24, 2023			
		7671031	Feb 24, 2023		U-727	
		7671031*PED	Aug 24, 2023			
		7674774	Feb 24, 2023	DP	U-727	
		7674774*PED	Aug 24, 2023			
		7678770	Feb 24, 2023		U-727	
		7678770*PED	Aug 24, 2023			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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 001	7678771	Feb 24, 2023	DP U-727			
	7678771*PED	Aug 24, 2023				
	7687466	Feb 24, 2023	DP			
	7687466*PED	Aug 24, 2023				
	7687467	Feb 24, 2023	DP U-727			
	7687467*PED	Aug 24, 2023				
	7713936	Feb 24, 2023	U-727			
	7713936*PED	Aug 24, 2023				
	7718619	Feb 24, 2023	DP U-727			
	7718619*PED	Aug 24, 2023				
	7723305	Feb 24, 2023	DP U-727			
	7723305*PED	Aug 24, 2023				
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510 002	7105486	Feb 24, 2023	U-727			
	7105486*PED	Aug 24, 2023				
	7223735	Feb 24, 2023	DP			
	7223735*PED	Aug 24, 2023				
	7655630	Feb 24, 2023	DS DP			
	7655630*PED	Aug 24, 2023				
	7659253	Feb 24, 2023	DS DP U-727			
	7659253*PED	Aug 24, 2023				
	7659254	Feb 24, 2023	U-727			
	7659254*PED	Aug 24, 2023				
	7662787	Feb 24, 2023	DS			
	7662787*PED	Aug 24, 2023				
	7662788	Feb 24, 2023	U-727			
	7662788*PED	Aug 24, 2023				
	7671030	Feb 24, 2023	DP U-727			
	7671030*PED	Aug 24, 2023				
	7671031	Feb 24, 2023	U-727			
	7671031*PED	Aug 24, 2023				
	7674774	Feb 24, 2023	DP U-727			
	7674774*PED	Aug 24, 2023				
	7678770	Feb 24, 2023	U-727			
	7678770*PED	Aug 24, 2023				
	7678771	Feb 24, 2023	DP U-727			
	7678771*PED	Aug 24, 2023				
	7687466	Feb 24, 2023	DP			
	7687466*PED	Aug 24, 2023				
	7687467	Feb 24, 2023	DP U-727			
	7687467*PED	Aug 24, 2023				
	7713936	Feb 24, 2023	U-727			
	7713936*PED	Aug 24, 2023				
	7718619	Feb 24, 2023	DP U-727			
	7718619*PED	Aug 24, 2023				
	7723305	Feb 24, 2023	DP U-727			
	7723305*PED	Aug 24, 2023				
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510 003	7105486	Feb 24, 2023	U-727			
	7105486*PED	Aug 24, 2023				
	7223735	Feb 24, 2023	DP			
	7223735*PED	Aug 24, 2023				
	7655630	Feb 24, 2023	DS DP			
	7655630*PED	Aug 24, 2023				
	7659253	Feb 24, 2023	DS DP U-727			
	7659253*PED	Aug 24, 2023				
	7659254	Feb 24, 2023	U-727			
	7659254*PED	Aug 24, 2023				
	7662787	Feb 24, 2023	DS			
	7662787*PED	Aug 24, 2023				
	7662788	Feb 24, 2023	U-727			
	7662788*PED	Aug 24, 2023				
	7671030	Feb 24, 2023	DP U-727			
	7671030*PED	Aug 24, 2023				
	7671031	Feb 24, 2023	U-727			
	7671031*PED	Aug 24, 2023				
	7674774	Feb 24, 2023	DP U-727			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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	7674774*PED	Aug 24, 2023				
	7678770	Feb 24, 2023	U-727			
	7678770*PED	Aug 24, 2023				
	7678771	Feb 24, 2023	DP U-727			
	7678771*PED	Aug 24, 2023				
	7687466	Feb 24, 2023	DP			
	7687466*PED	Aug 24, 2023				
	7687467	Feb 24, 2023	DP U-727			
	7687467*PED	Aug 24, 2023				
	7713936	Feb 24, 2023	U-727			
	7713936*PED	Aug 24, 2023				
	7718619	Feb 24, 2023	DP U-727			
	7718619*PED	Aug 24, 2023				
	7723305	Feb 24, 2023	DP U-727			
	7723305*PED	Aug 24, 2023				
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510 004	7105486	Feb 24, 2023	U-727			
	7105486*PED	Aug 24, 2023				
	7223735	Feb 24, 2023	DP			
	7223735*PED	Aug 24, 2023				
	7655630	Feb 24, 2023	DS DP			
	7655630*PED	Aug 24, 2023				
	7659253	Feb 24, 2023	DS DP U-727			
	7659253*PED	Aug 24, 2023				
	7659254	Feb 24, 2023	U-727			
	7659254*PED	Aug 24, 2023				
	7662787	Feb 24, 2023	DS			
	7662787*PED	Aug 24, 2023				
	7662788	Feb 24, 2023	U-727			
	7662788*PED	Aug 24, 2023				
	7671030	Feb 24, 2023	DP U-727			
	7671030*PED	Aug 24, 2023				
	7671031	Feb 24, 2023	U-727			
	7671031*PED	Aug 24, 2023				
	7674774	Feb 24, 2023	DP U-727			
	7674774*PED	Aug 24, 2023				
	7678770	Feb 24, 2023	U-727			
	7678770*PED	Aug 24, 2023				
	7678771	Feb 24, 2023	DP U-727			
	7678771*PED	Aug 24, 2023				
	7687466	Feb 24, 2023	DP			
	7687466*PED	Aug 24, 2023				
	7687467	Feb 24, 2023	DP U-727			
	7687467*PED	Aug 24, 2023				
	7713936	Feb 24, 2023	U-727			
	7713936*PED	Aug 24, 2023				
	7718619	Feb 24, 2023	DP U-727			
	7718619*PED	Aug 24, 2023				
	7723305	Feb 24, 2023	DP U-727			
	7723305*PED	Aug 24, 2023				
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510 005	7105486	Feb 24, 2023	U-727			
	7105486*PED	Aug 24, 2023				
	7223735	Feb 24, 2023	DP			
	7223735*PED	Aug 24, 2023				
	7655630	Feb 24, 2023	DS DP			
	7655630*PED	Aug 24, 2023				
	7659253	Feb 24, 2023	DS DP U-727			
	7659253*PED	Aug 24, 2023				
	7659254	Feb 24, 2023	U-727			
	7659254*PED	Aug 24, 2023				
	7662787	Feb 24, 2023	DS			
	7662787*PED	Aug 24, 2023				
	7662788	Feb 24, 2023	U-727			
	7662788*PED	Aug 24, 2023				
	7671030	Feb 24, 2023	DP U-727			
	7671030*PED	Aug 24, 2023				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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 005	7671031	Feb 24, 2023	U-727			
	7671031*PED	Aug 24, 2023				
	7674774	Feb 24, 2023	DP U-727			
	7674774*PED	Aug 24, 2023				
	7678770	Feb 24, 2023	U-727			
	7678770*PED	Aug 24, 2023				
	7678771	Feb 24, 2023	DP U-727			
	7678771*PED	Aug 24, 2023				
	7687466	Feb 24, 2023	DP			
	7687466*PED	Aug 24, 2023				
	7687467	Feb 24, 2023	DP U-727			
	7687467*PED	Aug 24, 2023				
	7713936	Feb 24, 2023	U-727			
	7713936*PED	Aug 24, 2023				
	7718619	Feb 24, 2023	DP U-727			
	7718619*PED	Aug 24, 2023				
	7723305	Feb 24, 2023	DP U-727			
	7723305*PED	Aug 24, 2023				
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510 006	7105486	Feb 24, 2023	U-727			
	7105486*PED	Aug 24, 2023				
	7223735	Feb 24, 2023	DP			
	7223735*PED	Aug 24, 2023				
	7655630	Feb 24, 2023	DS DP			
	7655630*PED	Aug 24, 2023				
	7659253	Feb 24, 2023	DS DP U-727			
	7659253*PED	Aug 24, 2023				
	7659254	Feb 24, 2023	U-727			
	7659254*PED	Aug 24, 2023				
	7662787	Feb 24, 2023	DS			
	7662787*PED	Aug 24, 2023				
	7662788	Feb 24, 2023	U-727			
	7662788*PED	Aug 24, 2023				
	7671030	Feb 24, 2023	DP U-727			
	7671030*PED	Aug 24, 2023				
	7671031	Feb 24, 2023	U-727			
	7671031*PED	Aug 24, 2023				
	7674774	Feb 24, 2023	DP U-727			
	7674774*PED	Aug 24, 2023				
	7678770	Feb 24, 2023	U-727			
	7678770*PED	Aug 24, 2023				
	7678771	Feb 24, 2023	DP U-727			
	7678771*PED	Aug 24, 2023				
	7687466	Feb 24, 2023	DP			
	7687466*PED	Aug 24, 2023				
	7687467	Feb 24, 2023	DP U-727			
	7687467*PED	Aug 24, 2023				
	7713936	Feb 24, 2023	U-727			
	7713936*PED	Aug 24, 2023				
	7718619	Feb 24, 2023	DP U-727			
	7718619*PED	Aug 24, 2023				
	7723305	Feb 24, 2023	DP U-727			
	7723305*PED	Aug 24, 2023				
<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			
	10265370	Nov 06, 2035	DP			
	10406199	Nov 06, 2035	U-1723			
	10406199	Nov 06, 2035	U-185			
	10406199	Nov 06, 2035	U-1864			
	10406199	Nov 06, 2035	U-1991			
	10406199	Nov 06, 2035	U-3			

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<u>LISINOPRIL - OBRELIS</u>						
N 208401	001	10406199	Nov 06, 2035	U-71		
		10406199	Nov 06, 2035	U-8		
		10940177	Nov 06, 2035	DP		
		11179434	Nov 06, 2035	DP		
		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>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		
		10555938	Mar 07, 2025	U-1316		
		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	002	10016404	Mar 07, 2025	U-1316		
		10555938	Mar 07, 2025	U-1316		
		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		
		10555938	Mar 07, 2025	U-1316		
		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		
		10555938	Mar 07, 2025	U-1316		
		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		
		10555938	Mar 07, 2025	U-1316		
		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		
		10555938	Mar 07, 2025	U-1316		

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<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	006	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>LONAFARNIB - ZOKINVY</u>						
N 213969	001	7838531	Jul 26, 2024	U-3070	NCE	Nov 20, 2025
		8828356	Oct 17, 2023	U-3070	ODE-324	Nov 20, 2027
<u>LONAFARNIB - ZOKINVY</u>						
N 213969	002	7838531	Jul 26, 2024	U-3070	NCE	Nov 20, 2025
		8828356	Oct 17, 2023	U-3070	ODE-324	Nov 20, 2027
<u>LOPERAMIDE HYDROCHLORIDE - LOPERAMIDE HYDROCHLORIDE</u>						
A 213070	001				CGT	Apr 20, 2022
<u>LORAZEPAM - LOREEV XR</u>						
N 214826	001	8999393	Jan 08, 2034	DP U-3210		
<u>LORAZEPAM - LOREEV XR</u>						
N 214826	002	8999393	Jan 08, 2034	DP U-3210		
<u>LORAZEPAM - LOREEV XR</u>						
N 214826	003	8999393	Jan 08, 2034	DP U-3210		
<u>LORLATINIB - LORBRENA</u>						
N 210868	001	10420749	Jul 27, 2036	DS DP U-2633	I-847	Mar 03, 2024
		10420749	Jul 27, 2036	DS DP U-3096	NCE	Nov 02, 2023
		11020376	Jul 27, 2036	DP	ODE-217	Nov 02, 2025
		8680111	Mar 05, 2033	DS DP	ODE-218	Nov 02, 2025
					ODE-219	Nov 02, 2025
					ODE-349	Mar 03, 2028
<u>LORLATINIB - LORBRENA</u>						
N 210868	002	10420749	Jul 27, 2036	DS DP U-2633	I-847	Mar 03, 2024
		10420749	Jul 27, 2036	DS DP U-3096	NCE	Nov 02, 2023
		11020376	Jul 27, 2036	DP	ODE-217	Nov 02, 2025
		8680111	Mar 05, 2033	DS DP	ODE-218	Nov 02, 2025
					ODE-219	Nov 02, 2025
					ODE-349	Mar 03, 2028
<u>LOTEPREDNOL ETABONATE - LOTEMAX SM</u>						
N 208219	001	10596107	Dec 23, 2036	DP U-2764	NS	Feb 22, 2022
<u>LOTEPREDNOL ETABONATE - INVELTYS</u>						
N 210565	001	10058511	May 03, 2033	DP U-2492		
		10646437	May 03, 2033	DP		
		10688045	May 03, 2033	DP		
		10864219	May 03, 2033	U-3011		
		9056057	May 03, 2033	DP U-2491		
		9393213	May 03, 2033	DP		
		9532955	May 03, 2033	U-2491		
		9737491	May 03, 2033	U-2492		
		9827191	May 03, 2033	DP U-2493		
<u>LOTEPREDNOL ETABONATE - EYSUVIS</u>						
N 210933	001	10058511	May 03, 2033	DP U-2492	NP	Oct 26, 2023
		10646436	May 03, 2033	DP		
		10688045	May 03, 2033	DP		
		10857096	May 03, 2033	U-2985		
		10940108	May 03, 2033	U-2985		
		10945948	May 03, 2033	U-2985		
		10993908	May 03, 2033	U-3117		
		9056057	May 03, 2033	DP U-2491		
		9393213	May 03, 2033	DP		
		9532955	May 03, 2033	U-2491		
		9737491	May 03, 2033	U-2492		
		9827191	May 03, 2033	DP U-2985		

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<u>LOXAPINE - ADASUVE</u>						
N 022549	001	6716416	May 20, 2022	DP		
		7458374	Aug 18, 2024	DP		
		7537009	Oct 28, 2024	DP		
		8074644	Jul 25, 2022	DP		
		8387612	Oct 23, 2026	DP		
		8991387	May 21, 2024	DP		
		9370629	May 20, 2024	DP		
<u>LUBIPROSTONE - AMITIZA</u>						
N 021908	001	6982283	Dec 04, 2022	U-1391		
		7064148	Aug 30, 2022	U-1404		
		7064148	Aug 30, 2022	U-739		
		8026393	Oct 25, 2027	DP		
		8097653	Nov 14, 2022	U-1214		
		8097653	Nov 14, 2022	U-1394		
		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	Jan 23, 2027	DP		
<u>LUBIPROSTONE - AMITIZA</u>						
N 021908	002	7064148	Aug 30, 2022	U-739		
		7064148	Aug 30, 2022	U-873		
		7795312	Sep 17, 2024	U-1085		
		8026393	Oct 25, 2027	DP		
		8338639	Jan 23, 2027	DP		
		8748481	Sep 01, 2025	U-1519		
		8779187	Jan 23, 2027	DP		
<u>LULICONAZOLE - LUZU</u>						
N 204153	001	8980931	Apr 28, 2034	DP		
		9012484	Sep 06, 2033	DS DP U-540		
		9199977	Sep 06, 2033	DS DP		
		9453006	Sep 06, 2033	DS		
<u>LUMASIRAN SODIUM - OXLUMO</u>						
N 214103	001	10131907	Aug 24, 2028	DS DP U-2995	NCE	Nov 23, 2025
		10435692	Dec 26, 2034	U-2995	ODE-339	Nov 23, 2027
		10465195	Dec 26, 2034	DS DP U-2995		
		10478500	Oct 09, 2035	DS DP U-2995		
		10487330	Dec 26, 2034	DS DP U-2995		
		10612024	Aug 14, 2035	DS DP U-2995		
		10612027	Aug 14, 2035	DS DP U-2995		
		11060093	Dec 26, 2034	DS DP U-2995		
		8106022	Dec 12, 2029	DS DP U-2995		
		8828956	Dec 04, 2028	DS DP U-2995		
		9828606	Dec 26, 2034	DS DP		
<u>LUMATEPERONE TOSYLATE - CAPLYTA</u>						
N 209500	001	10464938	Mar 12, 2028	DP	NCE	Dec 20, 2024
		10695345	Aug 30, 2039	DP U-543		
		10960009	Dec 03, 2034	U-814		
		8598119	Dec 28, 2029	U-543		
		8648077	Dec 01, 2029	DS DP		
		9199995	Mar 12, 2029	U-2713		
		9586960	Mar 12, 2029	DS DP		
		9616061	May 27, 2029	DP		
		9956227	Dec 03, 2034	U-2714		
		RE48825	Mar 12, 2029	DS DP		
		RE48839	Dec 28, 2029	U-814		
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603	001	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		

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<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603	001	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	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	
		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	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	
		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	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	
		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	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	

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<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603	005	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>LURBINECTEDIN - ZEPZELCA</u>						
N 213702	001	7763615	Dec 13, 2024	DS DP	U-2836	NCE Jun 15, 2025 ODE-304 Jun 15, 2027
<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	10596276	Jul 25, 2038	DP		NCE Jan 26, 2023
		10596278	Jul 25, 2038	DP		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	10946015	Sep 11, 2026		DP U-1445	
		7094781	Dec 05, 2025	DS DP		
		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 CHLORIDE; SODIUM SULFATE - SUTAB</u>						
N 213135	001	10143656	Aug 04, 2037	DP		NP Nov 10, 2023
		11033498	Aug 04, 2037		U-3164	
<u>MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE - SUPREP BOWEL PREP KIT</u>						
N 022372	001	6946149	Mar 07, 2023	DP	U-837	ODE-315 Aug 05, 2027
<u>MALATHION - OVIDE</u>						
N 018613	001	7560445	Feb 01, 2027	DS DP	U-986	
		7977324	Aug 14, 2026		DP	
<u>MANNITOL - BRONCHITOL</u>						
N 202049	001					NP ODE-327 Oct 30, 2023 Oct 30, 2027
<u>MARALIXIBAT CHLORIDE - LIVMARLI</u>						
N 214662	001					NCE Sep 29, 2026
<u>MARAVIROC - SELZENTRY</u>						
N 022128	001	6667314*PED	Feb 06, 2022			
		7368460	Nov 25, 2022		U-824	
		7368460*PED	May 25, 2023			
<u>MARAVIROC - SELZENTRY</u>						
N 022128	002	6667314*PED	Feb 06, 2022			
		7368460	Nov 25, 2022		U-824	
		7368460*PED	May 25, 2023			

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<u>MARAVIROC - SELZENTRY</u>						
N 022128	003	6667314*PED	Feb 06, 2022			
		7368460	Nov 25, 2022	U-824		
		7368460*PED	May 25, 2023			
<u>MARAVIROC - SELZENTRY</u>						
N 022128	004	6667314*PED	Feb 06, 2022			
		7368460	Nov 25, 2022	U-824		
		7368460*PED	May 25, 2023			
<u>MARAVIROC - SELZENTRY</u>						
N 208984	001	6667314*PED	Feb 06, 2022		NPP	Oct 30, 2023
		7368460	Nov 25, 2022	U-824	PED	Apr 30, 2024
		7368460*PED	May 25, 2023			
<u>MECHLORETHAMINE HYDROCHLORIDE - VALCHLOR</u>						
N 202317	001	7838564	Mar 07, 2026	DP		
		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>MEGESTROL ACETATE - MEGACE ES</u>						
N 021778	001	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 - 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 - ANJESO</u>						
N 210583	001	10463673	Feb 24, 2024	DP U-2750	NP	Feb 20, 2023
		10471067	Feb 24, 2024	DP U-2750		
		10709713	May 26, 2030	U-2750		
		10881663	Mar 08, 2039	U-3038		
		8512727	Dec 25, 2022	DP U-2750		
		9974746	May 26, 2030	DP		
<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 FLUFENAMIDE HYDROCHLORIDE - PEPAXTO</u>						
N 214383	001	10285946	Apr 25, 2032	DP	NCE	Feb 26, 2026
		10322182	Apr 25, 2032	DP	ODE-348	Feb 26, 2028
		10543274	Apr 25, 2032	U-3093		
		10869928	Apr 25, 2032	DP		
		6992207	Jun 25, 2022	DS DP U-3093		
<u>MELPHALAN HYDROCHLORIDE - EVOMELA</u>						
N 207155	001	10040872	Jan 30, 2034	DP	ODE-110	Mar 10, 2023
		10864183	May 28, 2030	DP		
		10940128	Jun 14, 2030	DP U-3086		
		11020363	May 28, 2030	DP		
		8410077	Mar 13, 2029	DP		
		9200088	Mar 13, 2029	DP		
		9493582	Feb 27, 2033	DP		

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<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	001	8039009	Mar 24, 2029	U-539		
		8039009*PED	Sep 24, 2029			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	002	8039009	Mar 24, 2029	U-539		
		8039009*PED	Sep 24, 2029			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	003	8039009	Mar 24, 2029	U-539		
		8039009*PED	Sep 24, 2029			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	004	8039009	Mar 24, 2029	U-539		
		8039009*PED	Sep 24, 2029			
<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>MEROPENEM; VABORBACTAM - VABOMERE</u>						
N 209776	001	10172874	Aug 08, 2031	DP	NCE	Aug 29, 2022
		10183034	Aug 08, 2031	U-2490	GAIN	Aug 29, 2027
		10561675	Aug 08, 2031	U-2490		
		11007206	Aug 08, 2031	U-3128		
		8680136	Aug 29, 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	002	8217083	Jun 06, 2028	DP		
		8436051	Jun 06, 2028	DP		
<u>MESALAMINE - LIALDA</u>						
N 022000	001				NPP	Jun 26, 2023
<u>MESALAMINE - APRISO</u>						
N 022301	001	8865688	May 01, 2030	U-1310		
<u>METAXALONE - SKELAXIN</u>						
N 013217	003	7122566	Feb 06, 2026	U-915		
<u>METFORMIN HYDROCHLORIDE - GLUMETZA</u>						
N 021748	002	7780987	Mar 23, 2025	DS DP		
		8323692	Mar 30, 2023	DP		
<u>METFORMIN HYDROCHLORIDE - RIOMET ER</u>						
N 212595	001	9962336	May 01, 2035	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	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		

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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		
		9339472	Jul 13, 2025	DP		
		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	8628799	Jul 13, 2025	DP		
		9339472	Jul 13, 2025	DP		
		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	003	8628799	Jul 13, 2025	DP		
		9339472	Jul 13, 2025	DP		
		RE44186	Jul 31, 2023	DS DP U-1097		
		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	M-187	Dec 04, 2023
		6699871*PED	Jan 26, 2023		M-244	Aug 12, 2022
		7125873	Jul 26, 2022	DP U-1036	PED	Feb 12, 2023
		7125873	Jul 26, 2022	DP U-1038	PED	Jun 04, 2024
		7125873	Jul 26, 2022	DP U-803		
		7125873*PED	Jan 26, 2023			
		7326708	Nov 24, 2026	DS DP U-802		
		7326708*PED	May 24, 2027			
		8414921	Jul 21, 2028	DP U-1036		
		8414921*PED	Jan 21, 2029			
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u>						
N 022044	002	6699871	Jul 26, 2022	DS DP U-802	M-187	Dec 04, 2023
		6699871*PED	Jan 26, 2023		M-244	Aug 12, 2022
		7125873	Jul 26, 2022	DP U-1036	PED	Feb 12, 2023
		7125873	Jul 26, 2022	DP U-1038	PED	Jun 04, 2024
		7125873	Jul 26, 2022	DP U-803		
		7125873*PED	Jan 26, 2023			
		7326708	Nov 24, 2026	DS DP U-802		
		7326708*PED	May 24, 2027			
		8414921	Jul 21, 2028	DP U-1036		
		8414921*PED	Jan 21, 2029			
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N 202270	001	6699871	Jul 26, 2022	DS DP U-1227	M-187	Dec 04, 2023
		6699871*PED	Jan 26, 2023		M-244	Aug 12, 2022
		7125873	Jul 26, 2022	DP U-1227	PED	Feb 12, 2023
		7125873*PED	Jan 26, 2023		PED	Jun 04, 2024
		7326708	Nov 24, 2026	DS DP U-1227		
		7326708*PED	May 24, 2027			

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<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N 202270 002	6699871	Jul 26, 2022	DS DP U-1227		M-187	Dec 04, 2023
	6699871*PED	Jan 26, 2023			M-244	Aug 12, 2022
	7125873	Jul 26, 2022	DP U-1227		PED	Feb 12, 2023
	7125873*PED	Jan 26, 2023			PED	Jun 04, 2024
	7326708	Nov 24, 2026	DS DP U-1227			
	7326708*PED	May 24, 2027				
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N 202270 003	6699871	Jul 26, 2022	DS DP U-1227		M-244	Aug 12, 2022
	6699871*PED	Jan 26, 2023			PED	Feb 12, 2023
	7125873	Jul 26, 2022	DP U-1227			
	7125873*PED	Jan 26, 2023				
	7326708	Nov 24, 2026	DS DP U-1227			
	7326708*PED	May 24, 2027				
<u>METHOTREXATE - OTREXUP</u>						
N 204824 001	10709844	Mar 10, 2029	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			
	8814834	May 27, 2031	DP			
	8945063	Mar 19, 2030	DP U-1442			
	9421333	Mar 19, 2030	DP U-1442			
	9533102	Jan 24, 2026	DP			
	9629959	Jan 24, 2026	DP			
	9867949	Mar 10, 2029	DP			
<u>METHOTREXATE - OTREXUP</u>						
N 204824 002	10709844	Mar 10, 2029	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			
	8814834	May 27, 2031	DP			
	8945063	Mar 19, 2030	DP U-1442			
	9421333	Mar 19, 2030	DP U-1442			
	9533102	Jan 24, 2026	DP			
	9629959	Jan 24, 2026	DP			
	9867949	Mar 10, 2029	DP			
<u>METHOTREXATE - OTREXUP</u>						
N 204824 003	10709844	Mar 10, 2029	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			
	8814834	May 27, 2031	DP			
	8945063	Mar 19, 2030	DP U-1442			
	9421333	Mar 19, 2030	DP U-1442			
	9533102	Jan 24, 2026	DP			
	9629959	Jan 24, 2026	DP			
	9867949	Mar 10, 2029	DP			
<u>METHOTREXATE - OTREXUP</u>						
N 204824 004	10709844	Mar 10, 2029	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			
	8814834	May 27, 2031	DP			
	8945063	Mar 19, 2030	DP U-1442			
	9421333	Mar 19, 2030	DP U-1442			
	9533102	Jan 24, 2026	DP			
	9629959	Jan 24, 2026	DP			
	9867949	Mar 10, 2029	DP			

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<u>METHOTREXATE - OTREXUP</u>						
N 204824	005	10709844	Mar 10, 2029	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	
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP	U-1442	
		9421333	Mar 19, 2030	DP	U-1442	
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	006	10709844	Mar 10, 2029	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	
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP	U-1442	
		9421333	Mar 19, 2030	DP	U-1442	
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	007	10709844	Mar 10, 2029	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	
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP	U-1442	
		9421333	Mar 19, 2030	DP	U-1442	
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	008	10709844	Mar 10, 2029	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	
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP	U-1442	
		9421333	Mar 19, 2030	DP	U-1442	
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - RASUVO</u>						
N 205776	001	8664231	Jun 01, 2029		U-1442	
<u>METHOTREXATE - RASUVO</u>						
N 205776	002	8664231	Jun 01, 2029		U-1442	
<u>METHOTREXATE - RASUVO</u>						
N 205776	003	8664231	Jun 01, 2029		U-1442	
<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	

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<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	10231927	Jan 02, 2033	U-1349	ODE-137	Apr 25, 2024
		10231927	Jan 02, 2033	U-1699	ODE-138	Apr 25, 2024
		10610485	Jan 02, 2033	DP		
		11116724	Jan 02, 2033	U-1349		
		11116724	Jan 02, 2033	U-1699		
		9259427	Jan 02, 2033	DP		
		9855215	Jan 02, 2033	DP		
<u>METHYLENE BLUE - PROVAYBLUE</u>						
N 204630	001				ODE-113	Apr 08, 2023
<u>METHYLENE BLUE - PROVAYBLUE</u>						
N 204630	002				ODE*	Apr 08, 2023
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>						
N 021964	001	10376584	Apr 08, 2024	DP U-1185		
		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	10376584	Apr 08, 2024	DP U-1185		
		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	10376584	Apr 08, 2024	DP U-1185		
		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	10307417	Mar 10, 2031	DP		
		10376505	Mar 10, 2031	DP		
		8420663	Sep 30, 2029	U-1185		
		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		

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<u>METHYLPHENIDATE - DAYTRANA</u>						
N 021514	001	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		
		9072680	Jun 28, 2032	DP		
		9089496	Jun 28, 2032	DP		
<u>METHYLPHENIDATE - COTEMPLA XR-ODT</u>						
N 205489	002	8840924	Jun 05, 2026	DP		
		9072680	Jun 28, 2032	DP		
		9089496	Jun 28, 2032	DP		
<u>METHYLPHENIDATE - COTEMPLA XR-ODT</u>						
N 205489	003	8840924	Jun 05, 2026	DP		
		9072680	Jun 28, 2032	DP		
		9089496	Jun 28, 2032	DP		
<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 - OUILIVANT 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	
		8956649	Feb 15, 2031	DP	U-1665	
		9040083	Feb 15, 2031	DP		
<u>METHYLPHENIDATE HYDROCHLORIDE - OUILICHEW ER</u>						
N 207960	001	10857143	Aug 14, 2033	DP	U-2993	
		11103494	Aug 14, 2033	DP		
		11103495	Aug 14, 2033	DP	U-2993	
		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 - OUILICHEW ER</u>						
N 207960	002	10857143	Aug 14, 2033	DP	U-2993	
		11103494	Aug 14, 2033	DP		
		11103495	Aug 14, 2033	DP	U-2993	
		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	

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<u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u>						
N 207960	003	10857143	Aug 14, 2033	DP	U-2993	
		11103494	Aug 14, 2033	DP		
		11103495	Aug 14, 2033	DP	U-2993	
		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	10182995	Mar 23, 2032	DP		
		10292937	Mar 23, 2032		U-2357	
		10617651	Mar 23, 2032		U-2357	
		10881618	Mar 23, 2032		U-2357	
		10905652	Mar 23, 2032	DP		
		8916588	Mar 23, 2032		U-2357	
		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	10182995	Mar 23, 2032	DP		
		10292937	Mar 23, 2032		U-2357	
		10617651	Mar 23, 2032		U-2357	
		10881618	Mar 23, 2032		U-2357	
		10905652	Mar 23, 2032	DP		
		8916588	Mar 23, 2032		U-2357	
		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	10182995	Mar 23, 2032	DP		
		10292937	Mar 23, 2032		U-2357	
		10617651	Mar 23, 2032		U-2357	
		10881618	Mar 23, 2032		U-2357	
		10905652	Mar 23, 2032	DP		
		8916588	Mar 23, 2032		U-2357	
		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	10182995	Mar 23, 2032	DP		
		10292937	Mar 23, 2032		U-2357	
		10617651	Mar 23, 2032		U-2357	
		10881618	Mar 23, 2032		U-2357	
		10905652	Mar 23, 2032	DP		
		8916588	Mar 23, 2032		U-2357	
		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	

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<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311	005	10182995	Mar 23, 2032	DP		
		10292937	Mar 23, 2032	U-2357		
		10617651	Mar 23, 2032	U-2357		
		10881618	Mar 23, 2032	U-2357		
		10905652	Mar 23, 2032	DP		
		8916588	Mar 23, 2032	U-2357		
		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 - ADHANSIA XR</u>						
N 212038	001	10111839	Oct 30, 2035	U-2357	M-82	Jun 28, 2024
		10292938	Oct 30, 2035	DP	NP	Feb 27, 2022
		10292939	Oct 30, 2035	DP U-2357		
		10449159	Oct 30, 2035	U-2357		
		10500162	Oct 30, 2035	U-2357		
		10507186	Oct 30, 2035	DP		
		10512612	Oct 30, 2035	DP		
		10512613	Oct 30, 2035	U-2357		
		10568841	Oct 30, 2035	DP U-2357		
		10688060	Oct 30, 2035	DP		
		10722473	Nov 19, 2038	U-2357		
		9974752	Oct 30, 2035	DP		
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038	002	10111839	Oct 30, 2035	U-2357	M-82	Jun 28, 2024
		10292938	Oct 30, 2035	DP	NP	Feb 27, 2022
		10292939	Oct 30, 2035	DP U-2357		
		10449159	Oct 30, 2035	U-2357		
		10500162	Oct 30, 2035	U-2357		
		10507186	Oct 30, 2035	DP		
		10512612	Oct 30, 2035	DP		
		10512613	Oct 30, 2035	U-2357		
		10568841	Oct 30, 2035	DP U-2357		
		10688060	Oct 30, 2035	DP		
		10722473	Nov 19, 2038	U-2357		
		9974752	Oct 30, 2035	DP		
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038	003	10111839	Oct 30, 2035	U-2357	M-82	Jun 28, 2024
		10292938	Oct 30, 2035	DP	NP	Feb 27, 2022
		10292939	Oct 30, 2035	DP U-2357		
		10449159	Oct 30, 2035	U-2357		
		10500162	Oct 30, 2035	U-2357		
		10507186	Oct 30, 2035	DP		
		10512612	Oct 30, 2035	DP		
		10512613	Oct 30, 2035	U-2357		
		10568841	Oct 30, 2035	DP U-2357		
		10688060	Oct 30, 2035	DP		
		10722473	Nov 19, 2038	U-2357		
		9974752	Oct 30, 2035	DP		
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038	004	10111839	Oct 30, 2035	U-2357	M-82	Jun 28, 2024
		10292938	Oct 30, 2035	DP	NP	Feb 27, 2022
		10292939	Oct 30, 2035	DP U-2357		
		10449159	Oct 30, 2035	U-2357		
		10500162	Oct 30, 2035	U-2357		
		10507186	Oct 30, 2035	DP		
		10512612	Oct 30, 2035	DP		
		10512613	Oct 30, 2035	U-2357		
		10568841	Oct 30, 2035	DP U-2357		
		10688060	Oct 30, 2035	DP		
		10722473	Nov 19, 2038	U-2357		
		9974752	Oct 30, 2035	DP		

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<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038	005	10111839	Oct 30, 2035	U-2357	M-82	Jun 28, 2024
		10292938	Oct 30, 2035	DP	NP	Feb 27, 2022
		10292939	Oct 30, 2035	DP U-2357		
		10449159	Oct 30, 2035	U-2357		
		10500162	Oct 30, 2035	U-2357		
		10507186	Oct 30, 2035	DP		
		10512612	Oct 30, 2035	DP		
		10512613	Oct 30, 2035	U-2357		
		10568841	Oct 30, 2035	DP U-2357		
		10688060	Oct 30, 2035	DP		
		10722473	Nov 19, 2038	U-2357		
		9974752	Oct 30, 2035	DP		
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038	006	10111839	Oct 30, 2035	U-2357	M-82	Jun 28, 2024
		10292938	Oct 30, 2035	DP	NP	Feb 27, 2022
		10292939	Oct 30, 2035	DP U-2357		
		10449159	Oct 30, 2035	U-2357		
		10500162	Oct 30, 2035	U-2357		
		10507186	Oct 30, 2035	DP		
		10512612	Oct 30, 2035	DP		
		10512613	Oct 30, 2035	U-2357		
		10568841	Oct 30, 2035	DP U-2357		
		10688060	Oct 30, 2035	DP		
		10722473	Nov 19, 2038	U-2357		
		9974752	Oct 30, 2035	DP		
<u>METOCLOPRAMIDE HYDROCHLORIDE - GIMOTI</u>						
N 209388	001	11020361	Dec 22, 2029	U-2843		
		8334281	May 16, 2030	DP U-2843		
<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	10238634	Jun 28, 2032	DP		
		10596155	Jun 28, 2032	DP		
		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				I-821	Dec 20, 2022
					PED	Jun 20, 2023
<u>MICAFUNGIN SODIUM - MYCAMINE</u>						
N 021506	003				I-821	Dec 20, 2022
					PED	Jun 20, 2023

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<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; PETROLATUM, WHITE; ZINC OXIDE - VUSION</u>						
N 021026	001	8147852	Mar 30, 2028	U-1426		
<u>MIDAZOLAM - NAYZILAM</u>						
N 211321	001	8217033	Jan 18, 2028	DP U-2526	NP	May 21, 2022
		8809322	Jan 18, 2028	DP	ODE-243	May 17, 2026
		9289432	Jan 18, 2028	DP U-2526		
		9687495	Jan 18, 2028	DP U-2526		
<u>MIDAZOLAM - MIDAZOLAM IN 0.9% SODIUM CHLORIDE</u>						
N 211844	001	10966990	Jun 20, 2038	DP		
<u>MIDAZOLAM - MIDAZOLAM IN 0.9% SODIUM CHLORIDE</u>						
N 211844	002	10966990	Jun 20, 2038	DP		
<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
<u>MIFEPRISTONE - KORLYM</u>						
N 202107	001	10006924	Aug 12, 2036	U-1643		
		10151763	Jan 18, 2037	U-1643		
		10166242	Apr 20, 2036	U-1643		
		10166243	Apr 20, 2036	U-1643		
		10195214	Jun 19, 2037	U-1643		
		10231983	Aug 22, 2038	U-1643		
		10314850	Aug 22, 2038	U-1643		
		10495650	Aug 12, 2036	U-1643		
		10500216	Mar 05, 2033	U-1643		
		10660904	Apr 20, 2036	U-1643		
		10780097	Aug 22, 2038	U-1643		
		10842800	Jun 19, 2037	U-1643		
		10842801	Nov 15, 2032	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
		10251873	May 30, 2038	U-2371	ODE-205	Aug 10, 2025
		10383864	May 16, 2027	U-2371		
		10406143	May 16, 2027	U-2371		
		10471053	May 30, 2038	U-2371		
		10525045	Apr 28, 2028	U-2371		
		10792278	May 30, 2038	U-2371		
		10792279	May 30, 2038	U-2371		
		10799491	May 30, 2038	U-2371		
		10806727	May 30, 2038	U-2371		
		10813921	Feb 12, 2029	U-2371		
		10849889	May 30, 2038	U-2371		
		10849890	May 30, 2038	U-2371		
		10857141	May 30, 2038	U-2371		
		10857142	May 30, 2038	U-2371		
		10874655	May 30, 2038	U-2371		
		10874656	May 30, 2038	U-2371		
		10874657	May 30, 2038	U-2371		
		10925866	Apr 28, 2028	U-2371		
		11033538	Apr 28, 2028	U-2371		
		8592362	Feb 12, 2029	U-2371		
		9000011	May 16, 2027	U-2371		
		9095584	Feb 12, 2029	U-2371		
		9480682	May 16, 2027	U-2371		

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<u>MIGALASTAT HYDROCHLORIDE - GALAFOLD</u>						
N 208623	001	9987263	May 16, 2027			U-2371
		9999618	Apr 28, 2028			U-2372
		9999618	Apr 28, 2028			U-2373
		RE48608	Feb 12, 2029			U-2371
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	001	6602911	Jan 14, 2023			U-882
		7994220	Sep 19, 2029			U-819
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	002	6602911	Jan 14, 2023			U-882
		7994220	Sep 19, 2029			U-819
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	003	6602911	Jan 14, 2023			U-882
		7994220	Sep 19, 2029			U-819
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	004	6602911	Jan 14, 2023			U-882
		7994220	Sep 19, 2029			U-819
<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
<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

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<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>MINOCYCLINE HYDROCHLORIDE - MINOLIRA</u>						
N 209269	001	11103517	Apr 07, 2036			DP
<u>MINOCYCLINE HYDROCHLORIDE - MINOLIRA</u>						
N 209269	002	11103517	Apr 07, 2036			DP
<u>MINOCYCLINE HYDROCHLORIDE - AMZEEQ</u>						
N 212379	001	10086080	Oct 01, 2030			U-2647
		10137200	Oct 01, 2030			U-2647
		10213512	Oct 01, 2030	DP		U-2647
		10265404	Oct 01, 2030	DP		
		10398641	Sep 08, 2037			U-2647
		10517882	Oct 01, 2030			U-2647
		10821187	Oct 01, 2030			U-2647
		10849847	Sep 08, 2037			U-2647
		8865139	Oct 01, 2030	DP		U-2647
		8945516	Oct 01, 2030	DP		
		8992896	Oct 01, 2030	DP		U-2647
		9675700	Oct 01, 2030	DP		U-2647
<u>MINOCYCLINE HYDROCHLORIDE - ZILXI</u>						
N 213690	001	10213512	Oct 01, 2030	DP		U-1631
		10265404	Oct 01, 2030	DP		
		10322186	Oct 01, 2030			U-1631
		10946101	Oct 01, 2030			U-1631
		8865139	Oct 01, 2030	DP		U-1631
		8945516	Oct 01, 2030	DP		
		8992896	Oct 01, 2030	DP		U-1631
		9675700	Oct 01, 2030	DP		U-1631
<u>MIPOMERSEN SODIUM - KYNAMRO</u>						
N 203568	001	7015315	Mar 21, 2023	DS		
		7101993	Sep 05, 2023	DS		

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<u>MIPOMERSEN SODIUM - KYNAMRO</u>						
N 203568	001 7511131	Jan 29, 2027	DS			
<u>MIRABEGRON - MYRBETRIO</u>						
N 202611	001 10842780	Sep 28, 2029		DP U-2996	I-855	Mar 25, 2024
	10842780*PED	Mar 28, 2030			PED	Sep 25, 2024
	6346532	Mar 27, 2022	DS DP			
	6346532*PED	Sep 27, 2022				
	7342117	Nov 04, 2023	DS			
	7982049	Nov 04, 2023		DP		
	8772315	Oct 30, 2028		U-2300		
	8772315*PED	Apr 30, 2029				
	8835474	Nov 04, 2023		U-1527		
	8835474*PED	May 04, 2024				
	RE44872	Nov 04, 2023		U-1527		
	RE44872*PED	May 04, 2024				
<u>MIRABEGRON - MYRBETRIO</u>						
N 202611	002 10842780	Sep 28, 2029		DP U-2996	I-855	Mar 25, 2024
	10842780*PED	Mar 28, 2030			PED	Sep 25, 2024
	6346532	Mar 27, 2022	DS DP			
	6346532*PED	Sep 27, 2022				
	7342117	Nov 04, 2023	DS			
	7982049	Nov 04, 2023		DP		
	8772315	Oct 30, 2028		U-2300		
	8772315*PED	Apr 30, 2029				
	8835474	Nov 04, 2023		U-1527		
	8835474*PED	May 04, 2024				
	RE44872	Nov 04, 2023		U-1527		
	RE44872*PED	May 04, 2024				
<u>MIRABEGRON - MYRBETRIO GRANULES</u>						
N 213801	001 10058536	Mar 31, 2036		DP U-3108	NP	Mar 25, 2024
	10058536*PED	Oct 01, 2036			PED	Sep 25, 2024
	6346532	Mar 27, 2022	DS			
	6346532*PED	Sep 27, 2022				
	7342117	Nov 04, 2023	DS			
	7342117*PED	May 04, 2024				
	7982049	Nov 04, 2023		DP		
	7982049*PED	May 04, 2024				
<u>MITOMYCIN - MITOSOL</u>						
N 022572	001 7806265	Feb 01, 2029		DP		
	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>MITOMYCIN - JELMYTO</u>						
N 211728	001 9040074	Jan 20, 2031		DP	NP	Apr 15, 2023
	9950069	Jan 20, 2031		DP	ODE-289	Apr 15, 2027
<u>MOBOCERTINIB SUCCINATE - EXKIVITY</u>						
N 215310	001 10227342	May 13, 2035	DS DP	U-3220	NCE	Sep 15, 2026
	9796712	May 13, 2035	DS DP		ODE-374	Sep 15, 2028
<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 - ASMANEX HFA</u>						
N 205641	003				NS	Aug 12, 2022
					PED	Feb 12, 2023
<u>MOMETASONE FUROATE - SINUVA</u>						
N 209310	001 10232152	Nov 24, 2034		DP U-2272		
	10357640	Oct 03, 2031		U-2272		
	10406332	Mar 13, 2034		DP		
	7544192	Nov 29, 2026		U-2272		

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<u>MOMETASONE FUROATE - SINUVA</u>						
N 209310	001	7662141	Mar 12, 2024			
		7713255	Mar 12, 2024			
		7951130	Mar 12, 2024			
		7951131	Mar 12, 2024			
		7951133	Mar 12, 2024			
		8025635	Jun 12, 2027	DP		
		8109918	Mar 12, 2024			
		8763222	Feb 08, 2032	DP		
		9585681	Apr 04, 2026			
<u>MONOMETHYL FUMARATE - BAFIERTAM</u>						
N 210296	001	10098863	Feb 27, 2035	DP		U-1384
		10105335	Feb 27, 2035	DP		
		10105336	Feb 27, 2035	DP		U-1384
		10105337	Feb 27, 2035	DP		
		10918615	Aug 12, 2035	DP		U-1384
		10918616	Jun 03, 2035			U-1384
		10918617	Aug 10, 2035	DP		
		10945985	Aug 14, 2035	DP		
		9326947	Feb 27, 2035	DP		
		9326965	Feb 27, 2035			U-1384
		9511043	Feb 27, 2035			U-1384
		9517209	Feb 27, 2035	DP		
		9566259	Feb 27, 2035	DP		
		9636318	Feb 27, 2035			U-1384
		9636319	Feb 27, 2035	DP		
		9814691	Feb 27, 2035			U-1384
		9814692	Feb 27, 2035			U-1384
		9820960	Feb 27, 2035	DP		
		9820961	Feb 27, 2035	DP		U-1384
<u>MONTELUKAST SODIUM - SINGULAIR</u>						
N 021409	001	8007830	Oct 24, 2022	DP		
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 022195	001				NPP	Jun 02, 2024
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 022195	002				NPP	Jun 02, 2024
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 022207	001				NPP	Jun 02, 2024
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 022207	002				NPP	Jun 02, 2024
<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
		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		

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<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	10314788	Aug 12, 2028	DP		
		7955619	Aug 12, 2028	DP		
<u>MORPHINE SULFATE - MORPHABOND ER</u>						
N 206544	002	10314788	Aug 12, 2028	DP		
		7955619	Aug 12, 2028	DP		
<u>MORPHINE SULFATE - MORPHABOND ER</u>						
N 206544	003	10314788	Aug 12, 2028	DP		
		7955619	Aug 12, 2028	DP		
<u>MORPHINE SULFATE - MORPHABOND ER</u>						
N 206544	004	10314788	Aug 12, 2028	DP		
		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		
		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		

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<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321 004	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 - MOXEZA</u>						
N 022428 001	8450311	May 29, 2029	DP			
	9114168	May 29, 2029	DP			
<u>MUPIROCIN CALCIUM - MUPIROCIN</u>						
A 213053 001					CGT	Jun 27, 2022
<u>NAFTIFINE HYDROCHLORIDE - NAFTIN</u>						
N 204286 001	10166205	Jan 31, 2033	DP			
	10166206	Jan 31, 2033	DP			
	10695303	Jan 31, 2033	DP			
	10729667	Jan 31, 2033	DP			
	8778365	Jan 31, 2033	DP			
	9161914	Jan 31, 2033	U-540			
<u>NALDEMEDINE TOSYLATE - SYMPROIC</u>						
N 208854 001	10952968	May 13, 2033	DS DP		NCE	Mar 23, 2022
	9108975	Nov 11, 2031	DS DP			
	RE46365	Jan 11, 2028	DS DP			
	RE46375	Oct 05, 2026	DS DP		U-1185	
<u>NALOXEGOL OXALATE - MOVANTIK</u>						
N 204760 001	7056500	Jun 29, 2024	DP		U-1185	
	7662365	Oct 18, 2022	DS DP			
	7786133	Sep 16, 2028	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	
	7662365	Oct 18, 2022	DS DP			
	7786133	Sep 16, 2028	DS DP			
	8067431	Dec 16, 2024			U-1185	
	8617530	Oct 18, 2022			U-1185	
	9012469	Apr 02, 2032	DS DP			

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<u>NALOXONE HYDROCHLORIDE - NALOXONE HYDROCHLORIDE</u>						
A 209522	001				PC	Jun 20, 2022
<u>NALOXONE HYDROCHLORIDE - EVZIO</u>						
N 205787	001	10143972	May 24, 2031	U-2476		
		10220158	Mar 20, 2035	DP U-2500		
		10314977	Nov 23, 2024	DP		
		10322239	Feb 28, 2031	U-1907		
		10335549	Apr 30, 2025	DP		
		10737028	Nov 23, 2024	DP		
		10960155	Jun 25, 2026	DP		
		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 (AUTOINJECTOR)</u>						
N 209862	001	10143792	May 24, 2031	U-2476		
		10220158	Mar 20, 2035	DP U-2500		
		10314977	Nov 23, 2024	DP		
		10322239	Feb 28, 2031	U-1907		
		10335549	Apr 30, 2025	DP		
		10737028	Nov 23, 2024	DP		
		10960155	Jun 25, 2026	DP		
		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		

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<u>NALOXONE HYDROCHLORIDE - EVZIO (AUTOINJECTOR)</u>						
N 209862	001	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 - KLOXXADO</u>						
N 212045	001	10722510	Aug 26, 2034	DP	U-3110	
		10973814	Aug 26, 2034	DP	U-3110	
<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	7919499	Oct 15, 2029		U-1123	
		7919499	Oct 15, 2029		U-1124	

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<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	001	7815934	Dec 12, 2027	DP		
		8685443	Jul 03, 2025		U-1508	
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	002	7815934	Dec 12, 2027	DP		
		8685443	Jul 03, 2025		U-1508	
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	003	7815934	Dec 12, 2027	DP		
		8685443	Jul 03, 2025		U-1508	
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	004	7815934	Dec 12, 2027	DP		
		8685443	Jul 03, 2025		U-1508	
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	005	7815934	Dec 12, 2027	DP		
		8685443	Jul 03, 2025		U-1508	
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	006	7815934	Dec 12, 2027	DP		
		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	
		11090280	Mar 03, 2026	DP	U-1731	
		11090280	Mar 03, 2026	DP	U-1732	
		11090280	Mar 03, 2026	DP	U-3195	
		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>NEBIVOLOL HYDROCHLORIDE; VALSARTAN - BYVALSON</u>						
N 206302	001	7803838	Aug 29, 2026	DP		
		7838552	Oct 04, 2027		U-185	
<u>NELARABINE - NELARABINE</u>						
A 215037	001				CGT	May 22, 2022
<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	
		10035788	Oct 15, 2028		U-3047	
		10035788	Oct 15, 2028		U-3097	
		7399865	Dec 29, 2030	DS DP		
		7982043	Oct 08, 2025		U-2043	
		7982043	Oct 08, 2025		U-3047	
		7982043	Oct 08, 2025		U-3097	
		8518446	Nov 20, 2030	DP	U-2043	
		8518446	Nov 20, 2030	DP	U-3047	
		8518446	Nov 20, 2030	DP	U-3097	
		8669273	Jul 18, 2031		U-3047	

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<u>NERATINIB MALEATE - NERLYNX</u>						
N 208051	001	8790708	Nov 05, 2030	DP	U-2043	
		8790708	Nov 05, 2030	DP	U-3047	
		8790708	Nov 05, 2030	DP	U-3097	
		9139558	Oct 15, 2028		U-2043	
		9139558	Oct 15, 2028		U-3047	
		9139558	Oct 15, 2028		U-3097	
		9211291	Mar 24, 2030		U-2043	
		9211291	Mar 24, 2030		U-3097	
		9265784	Aug 04, 2029		U-3047	
		9630946	Oct 15, 2028		U-2043	
		9630946	Oct 15, 2028		U-3047	
		9630946	Oct 15, 2028		U-3097	
<u>NETARSUDIL MESYLATE - RHOPRESSA</u>						
N 208254	001	10174017	Jan 27, 2030	DS DP	U-1524	NCE Dec 18, 2022
		10532993	Jul 11, 2026		U-1524	
		10588901	Mar 14, 2034	DS DP	U-1524	
		10654844	Jan 27, 2030	DS DP	U-1524	
		10882840	Jul 11, 2026		U-1524	
		11021456	Jul 11, 2026		U-1524	
		11028081	Jan 27, 2030		U-1524	
		11185538	Mar 14, 2034	DP		
		8394826	Nov 10, 2030	DS DP	U-1524	
		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	10233154	Sep 25, 2035	DS		
		10676440	Sep 25, 2035	DS DP		
		10828297	Dec 17, 2030		U-2293	
		10961195	Sep 25, 2035	DS DP		
		6297375	Mar 17, 2023	DS		
		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	10758616	Apr 18, 2027	DP		
		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	10758616	Apr 18, 2027	DP		
		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	10758616	Apr 18, 2027	DP		
		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		

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<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		
<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>NIFURTIMOX - LAMPIT</u>						
N 213464	001				NCE ODE-319	Aug 06, 2025 Aug 06, 2027
<u>NIFURTIMOX - LAMPIT</u>						
N 213464	002				NCE ODE-319	Aug 06, 2025 Aug 06, 2027
<u>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068	001	7169791	Jul 04, 2023	DS DP U-836	ODE-171	Mar 22, 2025
		7169791*PED	Jan 04, 2024		ODE-172	Mar 22, 2025
		8163904	Aug 23, 2028	DS DP	PED	Sep 22, 2025
		8163904*PED	Feb 23, 2029		PED	Sep 22, 2025
		8293756	Sep 25, 2027	DP		
		8293756*PED	Mar 25, 2028			
		8389537	Jul 18, 2026	U-1374		
		8389537*PED	Jan 18, 2027			
		8415363	Jul 18, 2026	DS DP U-1374		
		8415363	Jul 18, 2026	DS DP U-1407		
		8415363*PED	Jan 18, 2027			
		8501760	Jul 18, 2026	DP		
		8501760*PED	Jan 18, 2027			
		9061029	Apr 07, 2032	DS U-1374		
		9061029	Apr 07, 2032	DS U-3231		
		9061029*PED	Oct 07, 2032			
<u>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068	002	7169791	Jul 04, 2023	DS DP U-836	ODE-171	Mar 22, 2025
		7169791*PED	Jan 04, 2024		ODE-172	Mar 22, 2025
		8163904	Aug 23, 2028	DS DP	PED	Sep 22, 2025
		8163904*PED	Feb 23, 2029		PED	Sep 22, 2025
		8293756	Sep 25, 2027	DP		
		8293756*PED	Mar 25, 2028			
		8389537	Jul 18, 2026	U-1374		
		8389537*PED	Jan 18, 2027			
		8415363	Jul 18, 2026	DS DP U-1374		
		8415363	Jul 18, 2026	DS DP U-1407		
		8415363*PED	Jan 18, 2027			
		8501760	Jul 18, 2026	DP		
		8501760*PED	Jan 18, 2027			
		9061029	Apr 07, 2032	DS U-1374		
		9061029	Apr 07, 2032	DS U-3231		
		9061029*PED	Oct 07, 2032			
<u>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068	003	7169791	Jul 04, 2023	DS DP U-836	ODE-171	Mar 22, 2025
		7169791*PED	Jan 04, 2024		ODE-172	Mar 22, 2025
		8163904	Aug 23, 2028	DS DP	PED	Sep 22, 2025
		8163904*PED	Feb 23, 2029		PED	Sep 22, 2025
		8293756	Sep 25, 2027	DP		
		8293756*PED	Mar 25, 2028			

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<u>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068	003	8389537	Jul 18, 2026		U-1374	
		8389537*PED	Jan 18, 2027			
		8415363	Jul 18, 2026	DS DP	U-1374	
		8415363	Jul 18, 2026	DS DP	U-1407	
		8415363*PED	Jan 18, 2027			
		8501760	Jul 18, 2026		DP	
		8501760*PED	Jan 18, 2027			
		9061029	Apr 07, 2032	DS	U-1374	
		9061029	Apr 07, 2032	DS	U-3231	
		9061029*PED	Oct 07, 2032			
<u>NIMODIPINE - NYMALIZE</u>						
N 203340	002	10342787	Apr 16, 2038		DP U-2804	
		10576070	Apr 16, 2038		DP U-2804	
		7070581	Jun 23, 2023		DP	
		8517997	May 14, 2024		DP	
<u>NINTEDANIB ESYLATE - OFEV</u>						
N 205832	001	10105323	Jun 04, 2029		DP	I-805 Sep 06, 2022
		10154990	Dec 20, 2025		U-2620	I-825 Mar 09, 2023
		6762180	Oct 01, 2025	DS DP		ODE-261 Sep 06, 2026
		7119093	Feb 21, 2024	DS DP		
		9907756	Jun 07, 2029		DP	
<u>NINTEDANIB ESYLATE - OFEV</u>						
N 205832	002	10105323	Jun 04, 2029		DP	I-805 Sep 06, 2022
		10154990	Dec 20, 2025		U-2620	I-825 Mar 09, 2023
		6762180	Oct 01, 2025	DS DP		ODE-261 Sep 06, 2026
		7119093	Feb 21, 2024	DS DP		
		9907756	Jun 07, 2029		DP	
<u>NIRAPARIB TOSYLATE - ZEJULA</u>						
N 208447	001	11091459	Mar 27, 2038		DP	I-813 Oct 23, 2022
		8071579	Aug 12, 2027		U-2655	I-814 Oct 23, 2022
		8071623	Mar 27, 2031	DS DP		I-833 Apr 29, 2023
		8143241	Aug 12, 2027		U-2655	NCE Mar 27, 2022
		8436185	Apr 24, 2029	DS		ODE-133 Mar 27, 2024
		8859562	Aug 04, 2031		U-2655	ODE-277 Oct 23, 2026
						ODE-278 Oct 23, 2026
						ODE-295 Apr 29, 2027
<u>NITISINONE - ORFADIN</u>						
N 206356	001	9301932	Feb 28, 2033		DP U-1836	
<u>NITISINONE - NITYR</u>						
N 209449	001	10328029	Jan 05, 2035		DP U-1836	
<u>NITISINONE - NITYR</u>						
N 209449	002	10328029	Jan 05, 2035		DP U-1836	
<u>NITISINONE - NITYR</u>						
N 209449	003	10328029	Jan 05, 2035		DP U-1836	
<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	

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<u>NITRIC OXIDE - INOMAX</u>						
N 020845	002	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		
		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>NITRIC OXIDE - GENOSYL</u>						
N 202860	001	10124142	Aug 18, 2025	U-3037		
		10213572	Feb 12, 2036	DP		
		10737051	Oct 20, 2035	DP		
		10814092	Oct 17, 2025	U-3037		
		10926054	Aug 13, 2029	DP		
		11103669	Jun 21, 2030	DP		
		6758214	Feb 23, 2022	DP		
		7560076	Apr 21, 2027	DP		
		7618594	Oct 17, 2026	DP		
		7947227	Oct 17, 2026	U-3037		
		8057742	Jan 18, 2026	U-3037		
		8226916	Aug 18, 2025	U-3037		
		8607785	Jul 14, 2030	DP		
		8609028	Aug 18, 2025	U-3037		
		8821801	Aug 18, 2025	DP		
		8944049	Aug 13, 2029	DP		
		9522249	Aug 18, 2025	DP		
		9604028	Aug 13, 2029	U-2793		
		9701538	Jan 28, 2029	DP		
		9956373	Aug 18, 2025	U-3037		
<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	10266822	Dec 05, 2025	U-1942	ODE-127	Dec 23, 2023
		10266822	Dec 05, 2025	U-1943		
		10266822	Dec 05, 2025	U-1944		
		10436802	Sep 11, 2035	U-1941		
		10436802	Sep 11, 2035	U-1942		
		10436802	Sep 11, 2035	U-1943		
		10436802	Sep 11, 2035	U-1944		

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<u>NUSINERSEN SODIUM - SPINRAZA</u>						
N 209531	001	10436802	Sep 11, 2035	U-2093		
		10436802	Sep 11, 2035	U-2094		
		7101993	Sep 05, 2023	DS		
		7838657	Jul 11, 2027	DS		
		8110560	Dec 05, 2025	U-1942		
		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	ODE-119	May 27, 2023
		10052337	Apr 26, 2036	DP		
		10174073	Jun 17, 2033	DS		
		10751349	Apr 26, 2036	DP		
		10758549	Apr 26, 2036	U-2945		
		7138390	Nov 16, 2022	DS DP		
		8058267	Feb 21, 2022	U-1854		
		8377916	Feb 21, 2022	U-1854		
		9238673	Jun 17, 2033	DP		
		RE48286	Nov 16, 2027	DS DP		
<u>OBETICHOLIC ACID - OCALIVA</u>						
N 207999	002	10047117	Sep 06, 2033	U-1854	ODE-119	May 27, 2023
		10052337	Apr 26, 2036	DP		
		10174073	Jun 17, 2033	DS		
		10751349	Apr 26, 2036	DP		
		10758549	Apr 26, 2036	U-2945		
		7138390	Nov 16, 2022	DS DP		
		8058267	Feb 21, 2022	U-1854		
		8377916	Feb 21, 2022	U-1854		
		9238673	Jun 17, 2033	DP		
		RE48286	Nov 16, 2027	DS DP		
<u>OCTREOTIDE ACETATE - MYCAPSSA</u>						
N 208232	001	10238709	Feb 03, 2036	U-2857	NP	Jun 26, 2023
		10695397	Feb 03, 2036	U-2857		
		11052126	Feb 03, 2036	U-2857		
		11141457	Dec 28, 2040	U-3232		
		8329198	Sep 17, 2029	DP		
		8535695	Sep 17, 2029	U-2857		
		9265812	Sep 17, 2029	DP		
		9566246	Sep 17, 2029	DP		
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498	001	10011633	Nov 08, 2031	U-3186	NCE	Jul 20, 2026
		10093697	Nov 08, 2031	U-3186	ODE-363	Jul 20, 2028
		10487111	Nov 08, 2031	U-3186		
		10487111	Nov 08, 2031	U-3187		
		10975046	Jun 20, 2039	DS		
		10981952	Nov 08, 2031	U-3186		
		10981952	Nov 08, 2031	U-3187		
		7132416	Sep 05, 2022	DS DP		
		9694018	Nov 08, 2031	U-3186		
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498	002	10011633	Nov 08, 2031	U-3186	NCE	Jul 20, 2026
		10093697	Nov 08, 2031	U-3186	ODE-363	Jul 20, 2028
		10487111	Nov 08, 2031	U-3186		
		10487111	Nov 08, 2031	U-3187		
		10975046	Jun 20, 2039	DS		
		10981952	Nov 08, 2031	U-3186		
		10981952	Nov 08, 2031	U-3187		
		7132416	Sep 05, 2022	DS DP		
		9694018	Nov 08, 2031	U-3186		

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<u>ODEVIXIBAT - BYLVAY</u>						
N 215498 002	10011633	Nov 08, 2031	U-3186		NCE	Jul 20, 2026
	10093697	Nov 08, 2031	U-3186		ODE-363	Jul 20, 2028
	10487111	Nov 08, 2031	U-3186			
	10487111	Nov 08, 2031	U-3187			
	10975046	Jun 20, 2039	DS			
	10981952	Nov 08, 2031	U-3186			
	10981952	Nov 08, 2031	U-3187			
	7132416	Sep 05, 2022	DS DP			
	9694018	Nov 08, 2031	U-3186			
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498 003	10011633	Nov 08, 2031	U-3186		NCE	Jul 20, 2026
	10093697	Nov 08, 2031	U-3186		ODE-363	Jul 20, 2028
	10487111	Nov 08, 2031	U-3186			
	10487111	Nov 08, 2031	U-3187			
	10975046	Jun 20, 2039	DS			
	10981952	Nov 08, 2031	U-3186			
	10981952	Nov 08, 2031	U-3187			
	7132416	Sep 05, 2022	DS DP			
	9694018	Nov 08, 2031	U-3186			
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498 004	10011633	Nov 08, 2031	U-3186		NCE	Jul 20, 2026
	10093697	Nov 08, 2031	U-3186		ODE-363	Jul 20, 2028
	10487111	Nov 08, 2031	U-3186			
	10487111	Nov 08, 2031	U-3187			
	10975046	Jun 20, 2039	DS			
	10981952	Nov 08, 2031	U-3186			
	10981952	Nov 08, 2031	U-3187			
	7132416	Sep 05, 2022	DS DP			
	9694018	Nov 08, 2031	U-3186			
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378 001	10300054	Aug 23, 2031	DP U-3140		NCE	May 28, 2026
	10300054	Aug 23, 2031	DP U-3141			
	10716785	Aug 23, 2031	U-3136			
	10716785	Aug 23, 2031	U-3137			
	11185541	Aug 23, 2031	U-3140			
	7262298	Nov 23, 2025	DS			
	8778960	Feb 13, 2032	U-3136			
	8778960	Feb 13, 2032	U-3137			
	9119848	Aug 30, 2031	DS			
	9126977	Aug 23, 2031	DP U-3136			
	9126977	Aug 23, 2031	DP U-3137			
	9517235	Aug 23, 2031	U-3138			
	9517235	Aug 23, 2031	U-3139			
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378 002	10300054	Aug 23, 2031	DP U-3140		NCE	May 28, 2026
	10300054	Aug 23, 2031	DP U-3141			
	10716785	Aug 23, 2031	U-3136			
	10716785	Aug 23, 2031	U-3137			
	11185541	Aug 23, 2031	U-3140			
	7262298	Nov 23, 2025	DS			
	8778960	Feb 13, 2032	U-3136			
	8778960	Feb 13, 2032	U-3137			
	9119848	Aug 30, 2031	DS			
	9126977	Aug 23, 2031	DP U-3136			
	9126977	Aug 23, 2031	DP U-3137			
	9517235	Aug 23, 2031	U-3138			
	9517235	Aug 23, 2031	U-3139			
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378 003	10300054	Aug 23, 2031	DP U-3140		NCE	May 28, 2026
	10300054	Aug 23, 2031	DP U-3141			
	10716785	Aug 23, 2031	U-3136			
	10716785	Aug 23, 2031	U-3137			
	11185541	Aug 23, 2031	U-3140			
	7262298	Nov 23, 2025	DS			
	8778960	Feb 13, 2032	U-3136			

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<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378	003	8778960	Feb 13, 2032		U-3137	
		9119848	Aug 30, 2031	DS		
		9126977	Aug 23, 2031	DP	U-3136	
		9126977	Aug 23, 2031	DP	U-3137	
		9517235	Aug 23, 2031		U-3138	
		9517235	Aug 23, 2031		U-3139	
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378	004	10300054	Aug 23, 2031	DP	U-3140	NCE May 28, 2026
		10300054	Aug 23, 2031	DP	U-3141	
		10716785	Aug 23, 2031		U-3136	
		10716785	Aug 23, 2031		U-3137	
		11185541	Aug 23, 2031		U-3140	
		7262298	Nov 23, 2025	DS		
		8778960	Feb 13, 2032		U-3136	
		8778960	Feb 13, 2032		U-3137	
		9119848	Aug 30, 2031	DS		
		9126977	Aug 23, 2031	DP	U-3136	
		9126977	Aug 23, 2031	DP	U-3137	
		9517235	Aug 23, 2031		U-3138	
		9517235	Aug 23, 2031		U-3139	
<u>OLAPARIB - LYNPARZA</u>						
N 206162	001	7151102	Apr 29, 2022	DS DP		
		7449464	Oct 11, 2024	DS DP		
		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-818	Dec 27, 2022
		7449464	Oct 11, 2024	DS DP	I-831	May 08, 2023
		7981889	Oct 11, 2024	DS DP	I-832	May 19, 2023
		8071579	Aug 12, 2027		U-2101	ODE-180 Aug 17, 2024
		8071579	Aug 12, 2027		U-2103	ODE-181 Aug 17, 2024
		8071579	Aug 12, 2027		U-2480	ODE-226 Dec 19, 2025
		8071579	Aug 12, 2027		U-2481	ODE-283 Dec 27, 2026
		8071579	Aug 12, 2027		U-2482	ODE-306 May 08, 2027
		8071579	Aug 12, 2027		U-2483	
		8071579	Aug 12, 2027		U-2716	
		8071579	Aug 12, 2027		U-2819	
		8071579	Aug 12, 2027		U-2820	
		8071579	Aug 12, 2027		U-2821	
		8071579	Aug 12, 2027		U-2822	
		8071579	Aug 12, 2027		U-2823	
		8071579	Aug 12, 2027		U-2824	
		8071579	Aug 12, 2027		U-2832	
		8071579	Aug 12, 2027		U-2833	
		8143241	Aug 12, 2027		U-2101	
		8143241	Aug 12, 2027		U-2103	
		8143241	Aug 12, 2027		U-2480	
		8143241	Aug 12, 2027		U-2481	
		8143241	Aug 12, 2027		U-2482	
		8143241	Aug 12, 2027		U-2483	
		8143241	Aug 12, 2027		U-2716	
		8143241	Aug 12, 2027		U-2819	
		8143241	Aug 12, 2027		U-2820	
		8143241	Aug 12, 2027		U-2821	
		8143241	Aug 12, 2027		U-2822	
		8143241	Aug 12, 2027		U-2823	
		8143241	Aug 12, 2027		U-2824	
		8143241	Aug 12, 2027		U-2832	
		8143241	Aug 12, 2027		U-2833	
		8475842	Dec 31, 2029	DP		
		8859562	Aug 04, 2031		U-2101	
		8859562	Aug 04, 2031		U-2103	
		8859562	Aug 04, 2031		U-2480	
		8859562	Aug 04, 2031		U-2481	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OLAPARIB - LYNPARZA</u>						
N 208558	001	8859562	Aug 04, 2031	U-2482		
		8859562	Aug 04, 2031	U-2483		
		8859562	Aug 04, 2031	U-2716		
		8859562	Aug 04, 2031	U-2819		
		8859562	Aug 04, 2031	U-2820		
		8859562	Aug 04, 2031	U-2821		
		8859562	Aug 04, 2031	U-2822		
		8859562	Aug 04, 2031	U-2823		
		8859562	Aug 04, 2031	U-2824		
		8859562	Aug 04, 2031	U-2832		
		8859562	Aug 04, 2031	U-2833		
		8912187	Mar 12, 2024	U-2101		
		8912187	Mar 12, 2024	U-2480		
		8912187	Mar 12, 2024	U-2481		
		8912187	Mar 12, 2024	U-2482		
		8912187	Mar 12, 2024	U-2483		
		8912187	Mar 12, 2024	U-2819		
		8912187	Mar 12, 2024	U-2820		
		8912187	Mar 12, 2024	U-2821		
		8912187	Mar 12, 2024	U-2822		
		8912187	Mar 12, 2024	U-2823		
		8912187	Mar 12, 2024	U-2824		
		9169235	Mar 12, 2024	U-2832		
		9169235	Mar 12, 2024	U-2833		
		9566276	Mar 12, 2024	U-2716		
<u>OLAPARIB - LYNPARZA</u>						
N 208558	002	7151102	Apr 29, 2022	DS DP	I-818	Dec 27, 2022
		7449464	Oct 11, 2024	DS DP	I-831	May 08, 2023
		7981889	Oct 11, 2024	DS DP	I-832	May 19, 2023
		8071579	Aug 12, 2027	U-2101	ODE-180	Aug 17, 2024
		8071579	Aug 12, 2027	U-2103	ODE-181	Aug 17, 2024
		8071579	Aug 12, 2027	U-2480	ODE-226	Dec 19, 2025
		8071579	Aug 12, 2027	U-2481	ODE-283	Dec 27, 2026
		8071579	Aug 12, 2027	U-2482	ODE-306	May 08, 2027
		8071579	Aug 12, 2027	U-2483		
		8071579	Aug 12, 2027	U-2716		
		8071579	Aug 12, 2027	U-2819		
		8071579	Aug 12, 2027	U-2820		
		8071579	Aug 12, 2027	U-2821		
		8071579	Aug 12, 2027	U-2822		
		8071579	Aug 12, 2027	U-2823		
		8071579	Aug 12, 2027	U-2824		
		8071579	Aug 12, 2027	U-2832		
		8071579	Aug 12, 2027	U-2833		
		8143241	Aug 12, 2027	U-2101		
		8143241	Aug 12, 2027	U-2103		
		8143241	Aug 12, 2027	U-2480		
		8143241	Aug 12, 2027	U-2481		
		8143241	Aug 12, 2027	U-2482		
		8143241	Aug 12, 2027	U-2483		
		8143241	Aug 12, 2027	U-2716		
		8143241	Aug 12, 2027	U-2819		
		8143241	Aug 12, 2027	U-2820		
		8143241	Aug 12, 2027	U-2821		
		8143241	Aug 12, 2027	U-2822		
		8143241	Aug 12, 2027	U-2823		
		8143241	Aug 12, 2027	U-2824		
		8143241	Aug 12, 2027	U-2832		
		8143241	Aug 12, 2027	U-2833		
		8475842	Dec 31, 2029	DP		
		8859562	Aug 04, 2031	U-2101		
		8859562	Aug 04, 2031	U-2103		
		8859562	Aug 04, 2031	U-2480		
		8859562	Aug 04, 2031	U-2481		
		8859562	Aug 04, 2031	U-2482		
		8859562	Aug 04, 2031	U-2483		
		8859562	Aug 04, 2031	U-2716		
		8859562	Aug 04, 2031	U-2819		

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<u>OLAPARIB - LYNPARZA</u>						
N 208558	002	8859562	Aug 04, 2031	U-2820		
		8859562	Aug 04, 2031	U-2821		
		8859562	Aug 04, 2031	U-2822		
		8859562	Aug 04, 2031	U-2823		
		8859562	Aug 04, 2031	U-2824		
		8859562	Aug 04, 2031	U-2832		
		8859562	Aug 04, 2031	U-2833		
		8912187	Mar 12, 2024	U-2101		
		8912187	Mar 12, 2024	U-2480		
		8912187	Mar 12, 2024	U-2481		
		8912187	Mar 12, 2024	U-2482		
		8912187	Mar 12, 2024	U-2483		
		8912187	Mar 12, 2024	U-2819		
		8912187	Mar 12, 2024	U-2820		
		8912187	Mar 12, 2024	U-2821		
		8912187	Mar 12, 2024	U-2822		
		8912187	Mar 12, 2024	U-2823		
		8912187	Mar 12, 2024	U-2824		
		9169235	Mar 12, 2024	U-2832		
		9169235	Mar 12, 2024	U-2833		
		9566276	Mar 12, 2024	U-2716		
<u>OLICERIDINE - OLINVYK</u>						
N 210730	001	11077098	Mar 23, 2032	DS DP U-2986	NCE	Oct 30, 2025
		8835488	Mar 23, 2032	DS DP U-2986		
		9309234	Mar 23, 2032	DS DP U-2986		
		9642842	Mar 23, 2032	DP U-2986		
<u>OLICERIDINE - OLINVYK</u>						
N 210730	002	11077098	Mar 23, 2032	DS DP U-2986	NCE	Oct 30, 2025
		8835488	Mar 23, 2032	DS DP U-2986		
		9309234	Mar 23, 2032	DS DP U-2986		
		9642842	Mar 23, 2032	DP U-2986		
<u>OLICERIDINE - OLINVYK</u>						
N 210730	003	11077098	Mar 23, 2032	DS DP U-2986	NCE	Oct 30, 2025
		8835488	Mar 23, 2032	DS DP U-2986		
		9309234	Mar 23, 2032	DS DP U-2986		
		9642842	Mar 23, 2032	DP U-2986		
<u>OLODATEROL HYDROCHLORIDE - STRIVERDI RESPIMAT</u>						
N 203108	001	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	Jan 19, 2027	DS		
		7786111	Nov 10, 2023	DP		
		7837235	Mar 13, 2028	DP		
		7896264	May 26, 2025	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	7056916	Dec 07, 2023	DS DP		
		7220742	May 12, 2025	DS DP U-1703		
		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	Jan 19, 2027	DS		
		7786111	Nov 10, 2023	DP		
		7837235	Mar 13, 2028	DP		
		7837235*PED	Sep 13, 2028			
		7896264	May 26, 2025	DP		
		8034809	May 12, 2025	U-1702		
		8044046	Nov 10, 2023	U-1702		

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<u>OLODATEROL HYDROCHLORIDE; TIOTROPIUM BROMIDE - STIOLTO RESPIMAT</u>						
N 206756	001 8733341	Oct 16, 2030	DP			
	9027967	Mar 31, 2027	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 - PATADAY ONCE DAILY RELIEF</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		
<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
	10383884	Oct 31, 2037		U-2576		
	10835542	Oct 31, 2037		U-2576		
	7326696	Sep 24, 2023	DS			
	7553828	Jun 02, 2023	DS			
	8383610	Sep 23, 2030	DS			
	9265740	Mar 05, 2029		U-1569		
	9314475	Mar 18, 2031	DP			
	9724358	Mar 05, 2029		U-1569		
<u>OMADACYCLINE TOSYLATE - NUZYRA</u>						
N 209817	001 10124014	Mar 05, 2029		U-2449	NCE	Oct 02, 2023
	10383884	Oct 31, 2037		U-2576	GAIN	Oct 02, 2028
	10835542	Oct 31, 2037		U-2576		
	7326696	Sep 24, 2023	DS			
	7553828	Jun 02, 2023	DS			
	9265740	Mar 05, 2029		DP		
	9724358	Mar 05, 2029		U-1569		
<u>OMBITASVIR; PARITAPREVIR; RITONAVIR - TECHNIVIE</u>						
N 207931	001 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-CARBOXYLIC ACIDS - EPANOVA</u>						
N 205060	001 10117844	Jan 04, 2033		U-2447		
	7960370	Dec 20, 2026	DP			
	8383678	Feb 07, 2025	DP	U-1511		
	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			
	10835488	Dec 08, 2036	DP			
<u>ONDANSETRON - ZUPLENZ</u>						
N 022524	001 8580830	Nov 23, 2029	DP			
	9095577	Jul 13, 2030	DP			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ONDANSETRON - ZUPLENZ</u>						
N 022524	002	8580830				
		Nov 23, 2029	DP			
		9095577				
		Jul 13, 2030	DP			
<u>OPICAPONE - ONGENTYS</u>						
N 212489	001	10071085				
		Mar 31, 2030	DP		NCE	Apr 24, 2025
		10357468				
		May 27, 2035		U-2812		
		10583130				
		Mar 31, 2030		U-2811		
		8168793				
		Apr 02, 2029	DS DP	U-2811		
		8524746				
		Jul 14, 2029		U-2811		
		8907099				
		May 12, 2027	DS			
		9550759				
		Jul 26, 2026		U-2811		
		9550759				
		Jul 26, 2026		U-2817		
		9550759				
		Jul 26, 2026		U-2818		
		9630955				
		Dec 12, 2032	DS DP	U-2811		
		9745290				
		Oct 10, 2027	DP	U-2811		
<u>OPICAPONE - ONGENTYS</u>						
N 212489	002	10071085				
		Mar 31, 2030	DP		NCE	Apr 24, 2025
		10357468				
		May 27, 2035		U-2812		
		10583130				
		Mar 31, 2030		U-2811		
		8168793				
		Apr 02, 2029	DS DP	U-2811		
		8524746				
		Jul 14, 2029		U-2811		
		8907099				
		May 12, 2027	DS			
		9550759				
		Jul 26, 2026		U-2811		
		9550759				
		Jul 26, 2026		U-2817		
		9550759				
		Jul 26, 2026		U-2818		
		9630955				
		Dec 12, 2032	DS DP	U-2811		
		9745290				
		Oct 10, 2027	DP	U-2811		
<u>ORITAVANCIN DIPHOSPHATE - ORBACTIV</u>						
N 206334	001	8420592				
		Aug 29, 2029		U-1570	NCE	Aug 06, 2019
		9649352				
		Jul 16, 2035	DP		GAIN	Aug 06, 2024
		9682061				
		Apr 26, 2030		U-1569		
<u>ORITAVANCIN DIPHOSPHATE - KIMYRSA</u>						
N 214155	001	8420592				
		Aug 29, 2029		U-3101	NCE	Aug 06, 2019
		9649352				
		Jul 16, 2035	DS DP		NP	Mar 12, 2024
		9682061				
		Apr 26, 2030		U-3101	GAIN	Aug 06, 2024
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021087	001					
					M-251	Aug 02, 2022
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021087	002					
					M-251	Aug 02, 2022
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021087	003					
					M-251	Aug 02, 2022
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021246	001					
					M-251	Aug 02, 2022
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021246	002					
					M-251	Aug 02, 2022
<u>OSILODROSTAT PHOSPHATE - ISTURISA</u>						
N 212801	001	10143680				
		Jul 06, 2035	DP		NCE	Mar 06, 2025
		10709691				
		Oct 12, 2035		U-2770	ODE-286	Mar 06, 2027
		8314097				
		Mar 27, 2029	DS DP			
		8609862				
		Jan 13, 2031		U-2770		
		8835646				
		Aug 23, 2026	DS DP			
		9434754				
		Jan 13, 2031	DS			
<u>OSILODROSTAT PHOSPHATE - ISTURISA</u>						
N 212801	002	10143680				
		Jul 06, 2035	DP		NCE	Mar 06, 2025
		10709691				
		Oct 12, 2035		U-2770	ODE-286	Mar 06, 2027
		8314097				
		Mar 27, 2029	DS DP			
		8609862				
		Jan 13, 2031		U-2770		
		8835646				
		Aug 23, 2026	DS DP			
		9434754				
		Jan 13, 2031	DS			

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<u>OSILODROSTAT PHOSPHATE - ISTURISA</u>						
N 212801 003	10143680	Jul 06, 2035	DP		NCE	Mar 06, 2025
	10709691	Oct 12, 2035		U-2770	ODE-286	Mar 06, 2027
	8314097	Mar 27, 2029	DS DP			
	8609862	Jan 13, 2031		U-2770		
	8835646	Aug 23, 2026	DS DP			
	9434754	Jan 13, 2031	DS			
<u>OSIMERTINIB MESYLATE - TAGRISSO</u>						
N 208065 001	10183020	Jan 02, 2035	DP U-1777		I-853	Dec 18, 2023
	10183020	Jan 02, 2035	DP U-2289		ODE-102	Nov 13, 2022
	10183020	Jan 02, 2035	DP U-3016		ODE-176	Apr 18, 2025
	8946235	Aug 08, 2032	DS DP U-1777		ODE-337	Dec 18, 2027
	8946235	Aug 08, 2032	DS DP U-2289			
	8946235	Aug 08, 2032	DS DP U-3016			
	9732058	Jul 25, 2032	DS DP U-1777			
	9732058	Jul 25, 2032	DS DP U-2289			
	9732058	Jul 25, 2032	DS DP U-3016			
<u>OSIMERTINIB MESYLATE - TAGRISSO</u>						
N 208065 002	10183020	Jan 02, 2035	DP U-1777		I-853	Dec 18, 2023
	10183020	Jan 02, 2035	DP U-2289		ODE-102	Nov 13, 2022
	10183020	Jan 02, 2035	DP U-3016		ODE-176	Apr 18, 2025
	8946235	Aug 08, 2032	DS DP U-1777		ODE-337	Dec 18, 2027
	8946235	Aug 08, 2032	DS DP U-2289			
	8946235	Aug 08, 2032	DS DP U-3016			
	9732058	Jul 25, 2032	DS DP U-1777			
	9732058	Jul 25, 2032	DS DP U-2289			
	9732058	Jul 25, 2032	DS DP U-3016			
<u>OSPHEMIFENE - OSPHENA</u>						
N 203505 001	6245819	Jul 21, 2025		U-1370	I-793	Jan 25, 2022
	6245819	Jul 21, 2025		U-905		
	8236861	Aug 11, 2026		U-1369		
	8236861	Aug 11, 2026		U-1370		
	8236861	Aug 11, 2026		U-905		
	8470890	Feb 13, 2024		U-1369		
	8470890	Feb 13, 2024		U-1370		
	8470890	Feb 13, 2024		U-905		
	8642079	Jul 09, 2028	DP			
	8772353	Feb 13, 2024		U-1369		
	8772353	Feb 13, 2024		U-1370		
	8772353	Feb 13, 2024		U-905		
	9241915	Feb 13, 2024		U-1369		
	9241915	Feb 13, 2024		U-1370		
	9241915	Feb 13, 2024		U-905		
	9566252	Nov 02, 2022		U-1370		
	9855224	Feb 13, 2024		U-1369		
	9855224	Feb 13, 2024		U-1370		
	9855224	Feb 13, 2024		U-905		
<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810 001	10220042	Apr 13, 2027		U-2501		
	11166960	Apr 13, 2027	DP			
	7722898	Apr 13, 2027	DP			
	7910131	Apr 13, 2027		U-2041		
	8617600	Apr 13, 2027	DP			
	8821930	Apr 13, 2027	DP			
	9119791	Apr 13, 2027		U-2041		
	9351975	Apr 13, 2027	DP			
	9370525	Apr 13, 2027	DP			
	9855278	Apr 13, 2027	DP			
<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810 002	10220042	Apr 13, 2027		U-2501		
	11166960	Apr 13, 2027	DP			
	7722898	Apr 13, 2027	DP			
	7910131	Apr 13, 2027		U-2041		
	8617600	Apr 13, 2027	DP			
	8821930	Apr 13, 2027	DP			
	9119791	Apr 13, 2027		U-2041		

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<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810	002	9351975	Apr 13, 2027	DP		
		9370525	Apr 13, 2027	DP		
		9855278	Apr 13, 2027	DP		
<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810	003	10220042	Apr 13, 2027		U-2501	
		11166960	Apr 13, 2027	DP		
		7722898	Apr 13, 2027	DP		
		7910131	Apr 13, 2027		U-2041	
		8617600	Apr 13, 2027	DP		
		8821930	Apr 13, 2027	DP		
		9119791	Apr 13, 2027		U-2041	
		9351975	Apr 13, 2027	DP		
		9370525	Apr 13, 2027	DP		
		9855278	Apr 13, 2027	DP		
<u>OXYBUTYNIN - GELNIQUE 3%</u>						
N 202513	001	7198801	Jun 25, 2022	DP		
<u>OXYBUTYNIN CHLORIDE - GELNIQUE</u>						
N 022204	001	10449173	Nov 06, 2029	DP	U-2637	
		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	
		10188644	Sep 02, 2036	DP	U-1556	
		10525052	Jul 07, 2023	DP	U-1556	
		10525053	Jul 07, 2023	DP		
		10646485	Sep 02, 2036	DP	U-1556	
		10668060	Dec 10, 2030	DP	U-1556	
		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	002	10004729	Dec 10, 2030	DP	U-1556	
		10188644	Sep 02, 2036	DP	U-1556	
		10525052	Jul 07, 2023	DP	U-1556	
		10525053	Jul 07, 2023	DP		
		10646485	Sep 02, 2036	DP	U-1556	
		10668060	Dec 10, 2030	DP	U-1556	
		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	
		10188644	Sep 02, 2036	DP	U-1556	
		10525052	Jul 07, 2023	DP	U-1556	

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<u>OXYCODONE - XTAMPZA ER</u>						
N 208090	003	10525053	Jul 07, 2023	DP		
		10646485	Sep 02, 2036	DP U-1556		
		10668060	Dec 10, 2030	DP U-1556		
		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		
		10188644	Sep 02, 2036	DP U-1556		
		10525052	Jul 07, 2023	DP U-1556		
		10525053	Jul 07, 2023	DP		
		10646485	Sep 02, 2036	DP U-1556		
		10668060	Dec 10, 2030	DP U-1556		
		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		
		10188644	Sep 02, 2036	DP U-1556		
		10525052	Jul 07, 2023	DP U-1556		
		10525053	Jul 07, 2023	DP		
		10646485	Sep 02, 2036	DP U-1556		
		10668060	Dec 10, 2030	DP U-1556		
		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 HYDROCHLORIDE - OXYCONTIN</u>						
N 022272	001	10130591	Nov 20, 2023	DP U-1819		
		10369109	Jun 16, 2023	DP		
		10407434	Mar 30, 2025	DS		
		10675278	Nov 20, 2023	DP		
		10696684	Mar 30, 2025	DS		
		8309060	Nov 20, 2023	DP U-1556		
		8808741	Aug 24, 2027		U-1556	
		8894987	Mar 29, 2030	DP		
		8894988	Aug 24, 2027	DP		
		9073933	Mar 30, 2025	DS		

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<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272	001	9492389	Aug 24, 2027	DP		
		9492391	Aug 24, 2027	DP	U-1556	
		9492392	Aug 24, 2027	DP		
		9492393	Aug 24, 2027	DP	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	
		10369109	Jun 16, 2023	DP		
		10407434	Mar 30, 2025	DS		
		10675278	Nov 20, 2023	DP		
		10696684	Mar 30, 2025	DS		
		8309060	Nov 20, 2023	DP	U-1556	
		8808741	Aug 24, 2027	DP	U-1556	
		8894987	Mar 29, 2030	DP		
		8894988	Aug 24, 2027	DP		
		9073933	Mar 30, 2025	DS		
		9492389	Aug 24, 2027	DP		
		9492391	Aug 24, 2027	DP	U-1556	
		9492392	Aug 24, 2027	DP		
		9492393	Aug 24, 2027	DP	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	
		10369109	Jun 16, 2023	DP		
		10407434	Mar 30, 2025	DS		
		10675278	Nov 20, 2023	DP		
		10696684	Mar 30, 2025	DS		
		8309060	Nov 20, 2023	DP	U-1556	
		8808741	Aug 24, 2027	DP	U-1556	
		8894987	Mar 29, 2030	DP		
		8894988	Aug 24, 2027	DP		
		9073933	Mar 30, 2025	DS		
		9492389	Aug 24, 2027	DP		
		9492391	Aug 24, 2027	DP	U-1556	
		9492392	Aug 24, 2027	DP		
		9492393	Aug 24, 2027	DP	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	
		10369109	Jun 16, 2023	DP		
		10407434	Mar 30, 2025	DS		
		10675278	Nov 20, 2023	DP		
		10696684	Mar 30, 2025	DS		
		8309060	Nov 20, 2023	DP	U-1556	
		8808741	Aug 24, 2027	DP	U-1556	
		8894987	Mar 29, 2030	DP		
		8894988	Aug 24, 2027	DP		
		9073933	Mar 30, 2025	DS		
		9492389	Aug 24, 2027	DP		
		9492391	Aug 24, 2027	DP	U-1556	
		9492392	Aug 24, 2027	DP		
		9492393	Aug 24, 2027	DP	U-1556	
		9522919	Mar 30, 2025	DS DP		
		9675610	Jun 16, 2023	DP		
		9763933	Aug 24, 2027	DP		

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<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272	004	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	
		10369109	Jun 16, 2023	DP		
		10407434	Mar 30, 2025	DS		
		10675278	Nov 20, 2023	DP		
		10696684	Mar 30, 2025	DS		
		8309060	Nov 20, 2023	DP	U-1556	
		8808741	Aug 24, 2027		U-1556	
		8894988	Aug 24, 2027	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	
		10369109	Jun 16, 2023	DP		
		10407434	Mar 30, 2025	DS		
		10675278	Nov 20, 2023	DP		
		10696684	Mar 30, 2025	DS		
		8309060	Nov 20, 2023	DP	U-1556	
		8808741	Aug 24, 2027		U-1556	
		8894988	Aug 24, 2027	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	
		10369109	Jun 16, 2023	DP		
		10407434	Mar 30, 2025	DS		
		10675278	Nov 20, 2023	DP		
		10696684	Mar 30, 2025	DS		
		8309060	Nov 20, 2023	DP	U-1556	
		8808741	Aug 24, 2027		U-1556	
		8894988	Aug 24, 2027	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		
		8409616	Nov 26, 2023	DP		
		8637540	Nov 26, 2023	DP		
		9492443	May 26, 2024	DP		

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<u>OXYCODONE HYDROCHLORIDE - OXAYDO</u>						
N 202080	001	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 - 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	10314788	Aug 12, 2028	DP		
		7955619	Aug 12, 2028	DP		
<u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u>						
N 209777	002	10314788	Aug 12, 2028	DP		
		7955619	Aug 12, 2028	DP		
<u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u>						
N 209777	003	10314788	Aug 12, 2028	DP		
		7955619	Aug 12, 2028	DP		
<u>OXYMETAZOLINE HYDROCHLORIDE - RHOFAD</u>						
N 208552	001	10335391	Jun 11, 2035		U-2567	
		10751325	Jun 11, 2035		U-2921	
		7812049	May 02, 2028		U-1959	
		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 - UPNEEO</u>						
N 212520	001	10799481	Dec 16, 2039		U-2849	
		10814001	Dec 16, 2039	DP		
		10898573	Dec 16, 2039	DP		
		10912765	Aug 26, 2031		U-2849	
		10940138	Dec 16, 2039		U-2849	
		11103482	Dec 16, 2039	DP		
		8357714	Aug 26, 2031		U-2849	
		9867808	Aug 26, 2031		U-2849	
<u>OXYMETAZOLINE HYDROCHLORIDE; TETRACAINE HYDROCHLORIDE - KOVANAZE</u>						
N 208032	001	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	
		8309112	Feb 04, 2023	DP		
		8329216	Feb 04, 2023	DP		
		8808737	Jun 21, 2027		U-3085	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	002	7276250	Feb 04, 2023	DP	U-826	
		8309112	Feb 04, 2023	DP		
		8329216	Feb 04, 2023	DP		
		8808737	Jun 21, 2027		U-3085	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	003	7276250	Feb 04, 2023	DP	U-826	
		8309112	Feb 04, 2023	DP		
		8329216	Feb 04, 2023	DP		
		8808737	Jun 21, 2027		U-3085	
		8871779	Nov 22, 2029	DS		

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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 021610	003	7276250	Feb 04, 2023	DP	U-826	
		8309112	Feb 04, 2023	DP		
		8329216	Feb 04, 2023	DP		
		8808737	Jun 21, 2027		U-3085	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	004	7276250	Feb 04, 2023	DP	U-826	
		8309112	Feb 04, 2023	DP		
		8329216	Feb 04, 2023	DP		
		8808737	Jun 21, 2027		U-3085	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	005	7276250	Feb 04, 2023	DP	U-826	
		8309112	Feb 04, 2023	DP		
		8329216	Feb 04, 2023	DP		
		8808737	Jun 21, 2027		U-3085	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	006	7276250	Feb 04, 2023	DP	U-826	
		8309112	Feb 04, 2023	DP		
		8329216	Feb 04, 2023	DP		
		8808737	Jun 21, 2027		U-3085	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	007	7276250	Feb 04, 2023	DP	U-826	
		8309112	Feb 04, 2023	DP		
		8329216	Feb 04, 2023	DP		
		8808737	Jun 21, 2027		U-3085	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA</u>						
N 021611	001					M-14 Oct 25, 2022
<u>OXYMORPHONE HYDROCHLORIDE - OPANA</u>						
N 021611	002					M-14 Oct 25, 2022
<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		
		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		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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	003	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>OZANIMOD HYDROCHLORIDE - ZEPOSIA</u>						
N 209899	001	10239846	Nov 15, 2030		U-3132	I-860 May 27, 2024
		8481573	May 14, 2029	DS DP	U-2774	NCE Mar 25, 2025
		8796318	May 14, 2029	DS DP		
		9382217	May 14, 2029		U-2774	
<u>OZANIMOD HYDROCHLORIDE - ZEPOSIA</u>						
N 209899	002	10239846	Nov 15, 2030		U-3132	I-860 May 27, 2024
		8481573	May 14, 2029	DS DP	U-2774	NCE Mar 25, 2025
		8796318	May 14, 2029	DS DP		
		9382217	May 14, 2029		U-2774	
<u>OZANIMOD HYDROCHLORIDE - ZEPOSIA</u>						
N 209899	003	10239846	Nov 15, 2030		U-3132	I-860 May 27, 2024
		8481573	May 14, 2029	DS DP	U-2774	NCE Mar 25, 2025
		8796318	May 14, 2029	DS DP		
		9382217	May 14, 2029		U-2774	
<u>OZENOXACIN - XEPI</u>						
N 208945	001	6335447	Nov 09, 2023	DS		NCE Dec 11, 2022
		9180200	Jan 29, 2032	DP	U-805	
		9399014	Dec 15, 2029		U-805	

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<u>PACLITAXEL - ABRAXANE</u>						
N 021660 001	7758891	Feb 21, 2026	U-1434		M-14	Dec 06, 2022
	7758891*PED	Aug 21, 2026			PED	Jun 06, 2023
	7820788	Oct 27, 2024	DP U-1092			
	7820788	Oct 27, 2024	DP U-1290			
	7820788	Oct 27, 2024	DP U-1434			
	7820788*PED	Apr 27, 2025				
	7923536	Dec 09, 2023	U-1117			
	7923536	Dec 09, 2023	U-1290			
	7923536	Dec 09, 2023	U-1434			
	7923536*PED	Jun 09, 2024				
	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			
	8138229*PED	Jun 09, 2024				
	8268348	Feb 21, 2026	U-1290			
	8314156	Dec 09, 2023	U-1290			
	8314156	Dec 09, 2023	U-1434			
	8314156*PED	Jun 09, 2024				
	9101543	Feb 21, 2026	U-1434			
	9101543*PED	Aug 21, 2026				
	9393318	Mar 04, 2032	U-1290			
	9393318*PED	Sep 04, 2032				
	9511046	Jan 12, 2034	U-1434			
	9511046*PED	Jul 12, 2034				
	9597409	Mar 04, 2032	U-1290			
	9597409*PED	Sep 04, 2032				
<u>PAFOLACIANINE SODIUM - CYTALUX</u>						
N 214907 001					NCE	Nov 29, 2026
<u>PALBOCICLIB - IBRANCE</u>						
N 207103 001	10723730	Feb 08, 2034	DS DP			
	6936612	Jan 16, 2023	DS DP			
	7208489	Jan 16, 2023	DS DP	Y		
	7456168	Jan 16, 2023	U-1998			
	7456168	Jan 16, 2023	U-2515			
	RE47739	Mar 05, 2027	DS DP			
<u>PALBOCICLIB - IBRANCE</u>						
N 207103 002	10723730	Feb 08, 2034	DS DP			
	6936612	Jan 16, 2023	DS DP			
	7208489	Jan 16, 2023	DS DP	Y		
	7456168	Jan 16, 2023	U-1998			
	7456168	Jan 16, 2023	U-2515			
	RE47739	Mar 05, 2027	DS DP			
<u>PALBOCICLIB - IBRANCE</u>						
N 207103 003	10723730	Feb 08, 2034	DS DP			
	6936612	Jan 16, 2023	DS DP			
	7208489	Jan 16, 2023	DS DP	Y		
	7456168	Jan 16, 2023	U-1998			
	7456168	Jan 16, 2023	U-2515			
	RE47739	Mar 05, 2027	DS DP			
<u>PALBOCICLIB - IBRANCE</u>						
N 212436 001	10723730	Feb 08, 2034	DS DP			
	11065250	May 24, 2036	DP			
	6936612	Jan 16, 2023	DS DP			
	7456168	Jan 16, 2023	U-2515			
	RE47739	Mar 05, 2027	DS DP			
<u>PALBOCICLIB - IBRANCE</u>						
N 212436 002	10723730	Feb 08, 2034	DS DP			
	11065250	May 24, 2036	DP			
	6936612	Jan 16, 2023	DS DP			
	7456168	Jan 16, 2023	U-2515			
	RE47739	Mar 05, 2027	DS DP			

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<u>PALBOCICLIB - IBRANCE</u>						
N 212436	003	10723730	Feb 08, 2034	DS DP		
		11065250	May 24, 2036	DP		
		6936612	Jan 16, 2023	DS DP		
		7456168	Jan 16, 2023		U-2515	
		RE47739	Mar 05, 2027	DS DP		
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264	001	9439906	Jan 26, 2031		U-1901	
		9439906	Jan 26, 2031		U-2757	
		9439906	Jan 26, 2031		U-2758	
		9439906	Jan 26, 2031		U-543	
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264	002	9439906	Jan 26, 2031		U-1901	
		9439906	Jan 26, 2031		U-2757	
		9439906	Jan 26, 2031		U-2758	
		9439906	Jan 26, 2031		U-543	
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264	003	9439906	Jan 26, 2031		U-1901	
		9439906	Jan 26, 2031		U-2757	
		9439906	Jan 26, 2031		U-2758	
		9439906	Jan 26, 2031		U-543	
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264	004	9439906	Jan 26, 2031		U-1901	
		9439906	Jan 26, 2031		U-2757	
		9439906	Jan 26, 2031		U-2758	
		9439906	Jan 26, 2031		U-543	
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264	005	9439906	Jan 26, 2031		U-1901	
		9439906	Jan 26, 2031		U-2757	
		9439906	Jan 26, 2031		U-2758	
		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	
<u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u>						
N 207946	004	10143693	Apr 05, 2036		U-2457	
		10143693	Apr 05, 2036		U-2458	
<u>PALIPERIDONE PALMITATE - INVEGA HAFYERA</u>						
N 207946	005				NS	Aug 30, 2024
<u>PALIPERIDONE PALMITATE - INVEGA HAFYERA</u>						
N 207946	006				NS	Aug 30, 2024
<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		

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<u>PALONOSETRON HYDROCHLORIDE - ALOXI</u>						
N 021372	001	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>PANOBINOSTAT LACTATE - FARYDAK</u>						
N 205353	001	7989494	Jan 17, 2028	DS DP	ODE-89	Feb 23, 2022
		8883842	Jun 13, 2028	U-1669		
<u>PANOBINOSTAT LACTATE - FARYDAK</u>						
N 205353	002	7989494	Jan 17, 2028	DS DP	ODE-89	Feb 23, 2022
		8883842	Jun 13, 2028	U-1669		
<u>PANOBINOSTAT LACTATE - FARYDAK</u>						
N 205353	003	7989494	Jan 17, 2028	DS DP	ODE-89	Feb 23, 2022
		8883842	Jun 13, 2028	U-1669		
<u>PANTOPRAZOLE SODIUM - PROTONIX IV</u>						
N 020988	001	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 020819	001				ODE*	Oct 18, 2023
<u>PARICALCITOL - ZEMPLAR</u>						
N 020819	002				ODE*	Oct 18, 2023
<u>PARICALCITOL - ZEMPLAR</u>						
N 020819	003				ODE*	Oct 18, 2023
<u>PARICALCITOL - ZEMPLAR</u>						
N 021606	001				ODE-125	Oct 18, 2023
<u>PARICALCITOL - ZEMPLAR</u>						
N 021606	002				ODE-125	Oct 18, 2023

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<u>PARICALCITOL - ZEMPLAR</u>						
N 021606	003				ODE-125	Oct 18, 2023
<u>PAROXETINE HYDROCHLORIDE - PAROXETINE HYDROCHLORIDE</u>						
A 215003	001				CGT	Mar 12, 2022
<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		
<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		
		8299209	Dec 27, 2025	DS DP		
<u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u>						
N 200677	002	7473761	Dec 14, 2026	DS DP		
		8299209	Dec 27, 2025	DS DP		
<u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u>						
N 200677	003	7473761	Dec 14, 2026	DS DP		
		8299209	Dec 27, 2025	DS DP		
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	001	7473761	Dec 14, 2026	DS DP	ODE-268	Jun 29, 2025
		7759308	Oct 25, 2026	DP		
		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	ODE-268	Jun 29, 2025
		7759308	Oct 25, 2026	DP		
		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	ODE-268	Jun 29, 2025
		7759308	Oct 25, 2026	DP		
		8822637	Aug 06, 2023		U-1629	
		9351923	May 23, 2028	DP		
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	004	7473761	Dec 14, 2026	DS DP	ODE-268	Jun 29, 2025
		7759308	Oct 25, 2026	DP		
		9351923	May 23, 2028	DP		
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	005	7473761	Dec 14, 2026	DS DP	ODE-268	Jun 29, 2025
		7759308	Oct 25, 2026	DP		
		9351923	May 23, 2028	DP		
<u>PATIROMER SORBITEX CALCIUM - VELTASSA</u>						
N 205739	001	10485821	Mar 30, 2024		U-1766	
		11123363	Oct 08, 2033		U-1766	
		7556799	Feb 27, 2025		U-1766	
		8147873	Jun 20, 2028	DP		
		8216560	Mar 14, 2027		U-1766	
		8282913	May 29, 2027	DP		
		8287847	Mar 30, 2024		U-1766	

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<u>PATROMER SORBITEX CALCIUM - VELTASSA</u>						
N 205739	001	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>PATROMER SORBITEX CALCIUM - VELTASSA</u>						
N 205739	002	10485821	Mar 30, 2024		U-1766	
		11123363	Oct 08, 2033		U-1766	
		7556799	Feb 27, 2025		U-1766	
		8147873	Jun 20, 2028	DP		
		8216560	Mar 14, 2027		U-1766	
		8282913	May 29, 2027	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>PATROMER SORBITEX CALCIUM - VELTASSA</u>						
N 205739	003	10485821	Mar 30, 2024		U-1766	
		11123363	Oct 08, 2033		U-1766	
		7556799	Feb 27, 2025		U-1766	
		8147873	Jun 20, 2028	DP		
		8216560	Mar 14, 2027		U-1766	
		8282913	May 29, 2027	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	10240152	Oct 20, 2029	DS DP	U-2378	NCE Aug 10, 2023
		11079379	Aug 27, 2035	DS DP	U-2378	ODE-197 Aug 10, 2025
		11141378	Apr 15, 2029		DP	
		8058069	Apr 15, 2029		DP	
		8158601	Nov 10, 2030		DP U-2378	
		8168775	Oct 20, 2029	DS DP	U-2378	
		8334373	May 27, 2025	DS DP		
		8492359	Apr 15, 2029		DP	
		8642076	Oct 03, 2027		DP	
		8741866	Oct 20, 2029		U-2378	
		8802644	Oct 21, 2030		DP U-2378	
		8822668	Apr 15, 2029		DP U-2378	
		9234196	Oct 20, 2029		DP U-2378	
		9364435	Apr 15, 2029		DP U-2378	
		9943538	Nov 04, 2023		DP	
		9943539	Nov 04, 2023		DP	
<u>PAZOPANIB HYDROCHLORIDE - VOTRIENT</u>						
N 022465	001	7105530	Oct 19, 2023	DS DP		
<u>PAZOPANIB HYDROCHLORIDE - VOTRIENT</u>						
N 022465	002	7105530	Oct 19, 2023	DS DP		
<u>PEGCETACOPLAN - EMPAVELI</u>						
N 215014	001	10035822	Nov 15, 2033	DS		NCE May 14, 2026
		10125171	Aug 02, 2033	DS		ODE-351 May 14, 2028
		10875893	Nov 15, 2033	DS	U-3124	
		11040107	Apr 09, 2038		DP U-3172	
		11040107	Apr 09, 2038		DP U-3173	
		11040107	Apr 09, 2038		DP U-3174	
		7888323	Dec 04, 2027	DS		
		7989589	Dec 04, 2027	DS		

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<u>PEGCETACOPLAN - EMPAVELI</u>						
N 215014	001	9169307	Nov 18, 2027	DS	U-3123	
<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	
<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 - PEMFEXY</u>						
N 209472	001	7772209	May 24, 2022		U-2728	
		7772209	May 24, 2022		U-2729	
		9604990	Oct 28, 2035	DS		

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<u>PEMETREXED - PEMFEXY</u>						
N 209472	001	7772209	May 24, 2022	U-2728		
		7772209	May 24, 2022	U-2729		
		9604990	Oct 28, 2035	DS		
<u>PEMIGATINIB - PEMAZYRE</u>						
N 213736	001	10131667	Jun 12, 2033	U-2809	NCE	Apr 17, 2025
		9611267	Jan 30, 2035	DS DP	ODE-292	Apr 17, 2027
<u>PEMIGATINIB - PEMAZYRE</u>						
N 213736	002	10131667	Jun 12, 2033	U-2809	NCE	Apr 17, 2025
		9611267	Jan 30, 2035	DS DP	ODE-292	Apr 17, 2027
<u>PEMIGATINIB - PEMAZYRE</u>						
N 213736	003	10131667	Jun 12, 2033	U-2809	NCE	Apr 17, 2025
		9611267	Jan 30, 2035	DS DP	ODE-292	Apr 17, 2027
<u>PERAMIVIR - RAPIVAB</u>						
N 206426	001	10391075	Feb 12, 2027	U-2622		
		10391075	Feb 12, 2027	U-3069		
		6562861	Dec 16, 2023	DS		
		8778997	May 07, 2027	U-1627		
		8778997	May 07, 2027	U-2622		
		8778997	May 07, 2027	U-3069		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	001	6949571	Jun 08, 2022	DS DP U-106		
		6949571	Jun 08, 2022	DS DP U-2088		
		6949571	Jun 08, 2022	DS DP U-2089		
		6949571	Jun 08, 2022	DS DP U-2428		
		6949571	Jun 08, 2022	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	002	6949571	Jun 08, 2022	DS DP U-106		
		6949571	Jun 08, 2022	DS DP U-2088		
		6949571	Jun 08, 2022	DS DP U-2089		
		6949571	Jun 08, 2022	DS DP U-2428		
		6949571	Jun 08, 2022	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	003	6949571	Jun 08, 2022	DS DP U-106		
		6949571	Jun 08, 2022	DS DP U-2088		
		6949571	Jun 08, 2022	DS DP U-2089		
		6949571	Jun 08, 2022	DS DP U-2428		
		6949571	Jun 08, 2022	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	004	6949571	Jun 08, 2022	DS DP U-106		
		6949571	Jun 08, 2022	DS DP U-2088		
		6949571	Jun 08, 2022	DS DP U-2089		
		6949571	Jun 08, 2022	DS DP U-2428		
		6949571	Jun 08, 2022	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	005	6949571	Jun 08, 2022	DS DP U-106		
		6949571	Jun 08, 2022	DS DP U-2088		
		6949571	Jun 08, 2022	DS DP U-2089		
		6949571	Jun 08, 2022	DS DP U-2428		
		6949571	Jun 08, 2022	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	006	6949571	Jun 08, 2022	DS DP U-106		
		6949571	Jun 08, 2022	DS DP U-2088		
		6949571	Jun 08, 2022	DS DP U-2089		
		6949571	Jun 08, 2022	DS DP U-2428		
		6949571	Jun 08, 2022	DS DP U-2429		

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<u>PERAMPANEL - FYCOMPA</u>						
N 202834	006	8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 208277	001	6949571	Jun 08, 2022	DS DP	U-106	
		6949571	Jun 08, 2022	DS DP	U-2088	
		6949571	Jun 08, 2022	DS DP	U-2089	
		6949571	Jun 08, 2022	DS DP	U-2428	
		6949571	Jun 08, 2022	DS DP	U-2429	
		8772497	Jul 01, 2026	DS		
<u>PERFLUTREN - DEFINITY</u>						
N 021064	001	10583207	Dec 28, 2035		U-665	
		10583208	Mar 16, 2037		U-665	
		10588988	May 04, 2037		U-665	
		9789210	Mar 16, 2037		U-665	
<u>PERFLUTREN - DEFINITY RT</u>						
N 021064	002	10022460	Dec 28, 2035	DS DP		
		10583207	Dec 28, 2035		U-665	
		10583208	Mar 16, 2037		U-665	
		10588988	May 04, 2037		U-665	
		9789210	Mar 16, 2037		U-665	
<u>PEXIDARTINIB HYDROCHLORIDE - TURALIO</u>						
N 211810	001	10189833	May 05, 2036		U-2606	
		10435404	Jul 24, 2038	DP		NCE Aug 02, 2024
		10730876	May 05, 2036	DS		ODE-250 Aug 02, 2026
		10941142	Jul 24, 2038	DP		
		10961240	Jul 24, 2038		U-2606	
		7893075	Oct 13, 2028	DS		
		8404700	Nov 21, 2027	DS		
		8461169	Apr 19, 2028		U-2606	
		8722702	Nov 21, 2027	DS		
		9169250	Nov 21, 2027	DS		
		9358235	Jun 08, 2033		U-2606	
		9802932	May 05, 2036	DS		
<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	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 - QSYMIA</u>						
N 022580	002	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 - QSYMIA</u>						
N 022580	003	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	

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<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u>						
N 022580	004	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	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>PIFLUFOLASTAT F-18 - PYLARIFY</u>						
N 214793	001	10947197	Jun 09, 2037	DS DP U-3130	NCE	May 26, 2026
		8487129	Nov 07, 2027	DS DP		
		8778305	Sep 21, 2030	DS DP U-3130		
		9861713	Jul 31, 2029	DS DP U-3130		
<u>PILOCARPINE HYDROCHLORIDE - VUITY</u>						
N 214028	001	10610518	Apr 24, 2039	U-3252	NP	Oct 28, 2024
<u>PIMAVANSERIN TARTRATE - NUPLAZID</u>						
N 207318	001	10028944	Jan 15, 2024	U-1974		
		7601740	Apr 29, 2030	DS DP		
		7659285	Aug 24, 2026	U-1844		
		7732615	Jun 03, 2028	DS DP		
		7923564	Sep 26, 2025	DS DP		
		8618130	Jan 15, 2024	U-1845		
		8921393	Jan 15, 2024	U-1846		
		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		
		10517860	Mar 23, 2037	U-1974		
		10953000	Mar 23, 2037	U-1974		
		7601740	Apr 29, 2030	DS DP		
		7659285	Aug 24, 2026	U-1844		
		7732615	Jun 03, 2028	DS DP		
		7923564	Sep 26, 2025	DS DP		
		8618130	Jan 15, 2024	U-1845		
		8921393	Jan 15, 2024	U-1846		
		9566271	Jan 15, 2024	U-1974		
		9765053	Jul 27, 2022	U-1974		
<u>PIMAVANSERIN TARTRATE - NUPLAZID</u>						
N 210793	001	10028944	Jan 15, 2024	U-1974		
		10449185	Aug 27, 2038	DP		
		10646480	Aug 27, 2038	DP		
		10849891	Aug 27, 2038	DP U-1974		
		7601740	Apr 29, 2030	DS DP		
		7659285	Aug 24, 2026	U-1844		
		7732615	Jun 03, 2028	DS DP		
		7923564	Sep 26, 2025	DS DP		
		8618130	Jan 15, 2024	U-1845		
		8921393	Jan 15, 2024	U-1846		
		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		

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<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		
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u>						
N 050750	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 IN PLASTIC CONTAINER</u>						
N 050750	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 IN PLASTIC CONTAINER</u>						
N 050750	003	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		
		7635707	Apr 22, 2029	U-1609		
		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-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	10188637	Mar 28, 2037	DP		
		7566729	Apr 22, 2029	U-2077		
		7566729	Apr 22, 2029	U-2078		
		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		

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<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>PITOLISANT HYDROCHLORIDE - WAKIX</u>						
N 211150	001 8207197	Feb 25, 2029	DS DP		I-846	Oct 13, 2023
	8354430	Feb 06, 2026		U-1101	NCE	Aug 14, 2024
	8354430	Feb 06, 2026		U-1102	ODE-255	Aug 14, 2026
	8486947	Sep 26, 2029		U-1101	ODE-331	Oct 13, 2027
	8486947	Sep 26, 2029		U-1102		
<u>PITOLISANT HYDROCHLORIDE - WAKIX</u>						
N 211150	002 8207197	Feb 25, 2029	DS DP		I-846	Oct 13, 2023
	8354430	Feb 06, 2026		U-1101	NCE	Aug 14, 2024
	8354430	Feb 06, 2026		U-1102	ODE-255	Aug 14, 2026
	8486947	Sep 26, 2029		U-1101	ODE-331	Oct 13, 2027
	8486947	Sep 26, 2029		U-1102		
<u>PLAZOMICIN SULFATE - ZEMDRI</u>						
N 210303	001 8383596	Jun 02, 2031	DS	U-2328	NCE	Jun 25, 2023
	8822424	Nov 21, 2028		DP	GAIN	Jun 25, 2028
	9266919	Nov 21, 2028		U-2328		
	9688711	Nov 21, 2028	DS	U-2328		
<u>PLECANATIDE - TRULANCE</u>						
N 208745	001 10011637	Jun 05, 2034	DS		NCE	Jan 19, 2022
	11142549	Jun 05, 2034		DP		
	7041786	Jan 30, 2028	DS			
	7799897	Jun 09, 2022	DS			
	8637451	Mar 28, 2022		U-1964		
	9610321	Sep 15, 2031		U-1999		
	9610321	Sep 15, 2031		U-2230		
	9616097	Aug 20, 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 6846412	Jul 19, 2022		DP		
	7731986	Nov 17, 2024	DS DP	U-1463		
	7814943	Nov 19, 2027		DP	U-1461	
	8122917	Sep 09, 2024		DP		
	9480652	May 12, 2032		DP		
<u>POMALIDOMIDE - POMALYST</u>						
N 204026	001 10555939	May 19, 2030		DP	M-14	Nov 20, 2023
	10555939*PED	Nov 19, 2030			ODE-296	May 14, 2027
	8198262	Jun 17, 2025		U-1360	ODE-297	May 14, 2027
	8198262	Jun 17, 2025		U-2254	PED	May 20, 2024
	8198262*PED	Dec 17, 2025			PED	Nov 14, 2027
	8673939	May 15, 2023		U-1360	PED	Nov 14, 2027
	8673939	May 15, 2023		U-2254		
	8673939*PED	Nov 15, 2023				
	8735428	May 15, 2023		U-1360		
	8735428	May 15, 2023		U-2254		
	8735428*PED	Nov 15, 2023				
	8828427	Jun 21, 2031	DS DP			
	8828427*PED	Dec 21, 2031				
	9993467	May 19, 2030		DP		
	9993467*PED	Nov 19, 2030				

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<u>POMALIDOMIDE - POMALYST</u>						
N 204026 002	10555939	May 19, 2030	DP		M-14	Nov 20, 2023
	10555939*PED	Nov 19, 2030			ODE-296	May 14, 2027
	8198262	Jun 17, 2025	U-1360		ODE-297	May 14, 2027
	8198262	Jun 17, 2025	U-2254		PED	May 20, 2024
	8198262*PED	Dec 17, 2025			PED	Nov 14, 2027
	8673939	May 15, 2023	U-1360		PED	Nov 14, 2027
	8673939	May 15, 2023	U-2254			
	8673939*PED	Nov 15, 2023				
	8735428	May 15, 2023	U-1360			
	8735428	May 15, 2023	U-2254			
	8735428*PED	Nov 15, 2023				
	8828427	Jun 21, 2031	DS DP			
	8828427*PED	Dec 21, 2031				
	9993467	May 19, 2030	DP			
	9993467*PED	Nov 19, 2030				
<u>POMALIDOMIDE - POMALYST</u>						
N 204026 003	10555939	May 19, 2030	DP		M-14	Nov 20, 2023
	10555939*PED	Nov 19, 2030			ODE-296	May 14, 2027
	8198262	Jun 17, 2025	U-1360		ODE-297	May 14, 2027
	8198262	Jun 17, 2025	U-2254		PED	May 20, 2024
	8198262*PED	Dec 17, 2025			PED	Nov 14, 2027
	8673939	May 15, 2023	U-1360		PED	Nov 14, 2027
	8673939	May 15, 2023	U-2254			
	8673939*PED	Nov 15, 2023				
	8735428	May 15, 2023	U-1360			
	8735428	May 15, 2023	U-2254			
	8735428*PED	Nov 15, 2023				
	8828427	Jun 21, 2031	DS DP			
	8828427*PED	Dec 21, 2031				
	9993467	May 19, 2030	DP			
	9993467*PED	Nov 19, 2030				
<u>POMALIDOMIDE - POMALYST</u>						
N 204026 004	10555939	May 19, 2030	DP		M-14	Nov 20, 2023
	10555939*PED	Nov 19, 2030			ODE-296	May 14, 2027
	8198262	Jun 17, 2025	U-1360		ODE-297	May 14, 2027
	8198262	Jun 17, 2025	U-2254		PED	May 20, 2024
	8198262*PED	Dec 17, 2025			PED	Nov 14, 2027
	8673939	May 15, 2023	U-1360		PED	Nov 14, 2027
	8673939	May 15, 2023	U-2254			
	8673939*PED	Nov 15, 2023				
	8735428	May 15, 2023	U-1360			
	8735428	May 15, 2023	U-2254			
	8735428*PED	Nov 15, 2023				
	8828427	Jun 21, 2031	DS DP			
	8828427*PED	Dec 21, 2031				
	9993467	May 19, 2030	DP			
	9993467*PED	Nov 19, 2030				
<u>PONATINIB HYDROCHLORIDE - ICLUSIG</u>						
N 203469 001	11192895	Dec 12, 2033	U-1700		I-849	Dec 18, 2023
	11192895	Dec 12, 2033	U-1701			
	11192895	Dec 12, 2033	U-1948			
	11192897	Dec 12, 2033	DS DP U-1700			
	11192897	Dec 12, 2033	DS DP U-1701			
	11192897	Dec 12, 2033	DS DP U-1948			
	8114874	Jan 24, 2027	DS DP			
	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	11192895	Dec 12, 2033	U-1700		I-849	Dec 18, 2023
	11192895	Dec 12, 2033	U-1701			
	11192895	Dec 12, 2033	U-1948			

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<u>PONATINIB HYDROCHLORIDE - ICLUSIG</u>						
N 203469	002	11192897	Dec 12, 2033	DS DP U-1700		
		11192897	Dec 12, 2033	DS DP U-1701		
		11192897	Dec 12, 2033	DS DP U-1948		
		8114874	Jan 24, 2027	DS DP		
		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	11192895	Dec 12, 2033		I-849	Dec 18, 2023
		11192895	Dec 12, 2033		U-1701	
		11192895	Dec 12, 2033		U-1948	
		11192897	Dec 12, 2033	DS DP U-1700		
		11192897	Dec 12, 2033	DS DP U-1701		
		11192897	Dec 12, 2033	DS DP U-1948		
		8114874	Jan 24, 2027	DS DP		
		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	004	11192895	Dec 12, 2033		I-849	Dec 18, 2023
		11192895	Dec 12, 2033		U-1701	
		11192895	Dec 12, 2033		U-1948	
		11192897	Dec 12, 2033	DS DP U-1700		
		11192897	Dec 12, 2033	DS DP U-1701		
		11192897	Dec 12, 2033	DS DP U-1948		
		8114874	Jan 24, 2027	DS DP		
		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>PONESIMOD - PONVORY</u>						
N 213498	001	10220023	Dec 10, 2035		NCE	Mar 18, 2026
		8273779	Dec 17, 2025		U-2774	
		9000018	Nov 16, 2024		U-3102	
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	002	10220023	Dec 10, 2035		NCE	Mar 18, 2026
		8273779	Dec 17, 2025		U-2774	
		9000018	Nov 16, 2024		U-3102	
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	003	10220023	Dec 10, 2035		NCE	Mar 18, 2026
		8273779	Dec 17, 2025		U-2774	
		9000018	Nov 16, 2024		U-3102	
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	004	10220023	Dec 10, 2035		NCE	Mar 18, 2026
		8273779	Dec 17, 2025		U-2774	
		9000018	Nov 16, 2024		U-3102	
		9062014	May 06, 2032	DS DP U-2774		

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<u>PONESIMOD - PONVORY</u>						
N 213498	004	RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	005	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	006	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	007	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	008	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	009	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	010	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>POSACONAZOLE - NOXAFIL</u>						
N 022003	001	8263600	Apr 01, 2022	DP		
<u>POSACONAZOLE - NOXAFIL</u>						
N 205053	001				NPP ODE-355	May 31, 2024 Jun 17, 2028
<u>POSACONAZOLE - NOXAFIL</u>						
N 205596	001	10117951	Mar 13, 2029	DP	NPP	May 31, 2024
		8410077	Mar 13, 2029	DP	ODE-355	Jun 17, 2028
		9023790	Jul 04, 2031	DP U-1698		
		9023790	Jul 04, 2031	DP U-3160		
		9023790	Jul 04, 2031	DP U-3171		
		9358297	Jun 24, 2031	DP U-3160		
		9358297	Jun 24, 2031	DP U-3171		
		9493582	Feb 27, 2033	DP		
		9750822	Mar 13, 2029	DP		
<u>POSACONAZOLE - NOXAFIL POWDERMIX KIT</u>						
N 214770	001				NP	May 31, 2024
<u>POTASSIUM PHOSPHATE, DIBASIC; POTASSIUM PHOSPHATE, MONOBASIC - POTASSIUM PHOSPHATES</u>						
N 212121	001	10632150	Apr 19, 2039	DP U-2789		

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<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>PRALSETINIB - GAVRETO</u>						
N 213721	001	10030005	Nov 01, 2036	DS DP	U-2827	NCE Sep 04, 2025
		10030005	Nov 01, 2036	DS DP	U-2952	ODE-318 Sep 04, 2027
		10030005	Nov 01, 2036	DS DP	U-3002	ODE-340 Dec 01, 2027
						ODE-341 Dec 01, 2027
<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>PRASTERONE - INTRAROSA</u>						
N 208470	001	8268806	Mar 19, 2031	DP		
		8629129	Aug 07, 2028	DP		
		8957054	Aug 07, 2028		U-1922	
<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>PREDNISONE - RAYOS</u>						
N 202020	001	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	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	

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<u>PREDNISONE - RAYOS</u>						
N 202020	003	8168218		DP U-1269		
		8309124		U-1292		
		8394407		DP U-1362		
		9040085		U-1362		
		9186332		U-1362		
		9504699		U-1362		
<u>PREGABALIN - LYRICA</u>						
N 021446	001				NPP	May 23, 2022
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 021446	002				NPP	May 23, 2022
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 021446	003				NPP	May 23, 2022
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 021446	004				NPP	May 23, 2022
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 021446	005				NPP	May 23, 2022
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 021446	006				NPP	May 23, 2022
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 021446	007				NPP	May 23, 2022
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 021446	008				NPP	May 23, 2022
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 022488	001				NPP	May 23, 2022
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA CR</u>						
N 209501	001	10022447	Nov 02, 2026	U-2136		
		10022447	Nov 02, 2026	U-2137		
		10022447*PED	May 02, 2027			
		8945620	Nov 02, 2026	DP U-2136		
		8945620	Nov 02, 2026	DP U-2137		
		8945620*PED	May 02, 2027			
		9144559	Nov 02, 2026	DP		
		9144559*PED	May 02, 2027			
<u>PREGABALIN - LYRICA CR</u>						
N 209501	002	10022447	Nov 02, 2026	U-2136		
		10022447	Nov 02, 2026	U-2137		
		10022447*PED	May 02, 2027			
		8945620	Nov 02, 2026	DP U-2136		
		8945620	Nov 02, 2026	DP U-2137		
		8945620*PED	May 02, 2027			
		9144559	Nov 02, 2026	DP		
		9144559*PED	May 02, 2027			
<u>PREGABALIN - LYRICA CR</u>						
N 209501	003	10022447	Nov 02, 2026	U-2136		
		10022447	Nov 02, 2026	U-2137		
		10022447*PED	May 02, 2027			
		8945620	Nov 02, 2026	DP U-2136		
		8945620	Nov 02, 2026	DP U-2137		
		8945620*PED	May 02, 2027			

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<u>PREGABALIN - LYRICA CR</u>						
N 209501	003 9144559	Nov 02, 2026	DP			
	9144559*PED	May 02, 2027				
<u>PRETOMANID - PRETOMANID</u>						
N 212862	001				NCE	Aug 14, 2024
					ODE-253	Aug 14, 2026
					GAIN	Aug 14, 2029
<u>PROGESTERONE - MILPROSA</u>						
N 201110	001 10537584	Feb 03, 2029	DP		NP	Apr 29, 2023
	10548904	Feb 03, 2029		U-2810		
	8580293	Jan 21, 2030		U-2810		
<u>PROPOFOL - DIPRIVAN</u>						
N 019627	002 8476010	Dec 01, 2024	DS DP			
	8476010*PED	Jun 01, 2025				
<u>PROPRANOLOL HYDROCHLORIDE - HEMANGEOL</u>						
N 205410	001 8338489	Oct 16, 2028		U-1496		
	8987262	Oct 16, 2028		U-1988		
<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>PYRIMETHAMINE - PYRIMETHAMINE</u>						
A 211271	001				CGT	Apr 02, 2022
<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			
	7169780*PED	Apr 03, 2024				
	7217713	Oct 21, 2022		U-257		
	7217713*PED	Apr 21, 2023				
	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 10772888	Mar 30, 2032		U-1663		
	7169780	Oct 03, 2023	DS DP			
	7169780*PED	Apr 03, 2024				
	7217713	Oct 21, 2022		U-257		
	7217713*PED	Apr 21, 2023				
	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			
	7169780*PED	Apr 03, 2024				
	7217713	Oct 21, 2022		U-257		
	7217713*PED	Apr 21, 2023				
	7435734	Oct 21, 2022		U-257		
	7435734*PED	Apr 21, 2023				

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<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 203045	001	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		
		7169780*PED	Apr 03, 2024			
		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 205786	001	7169780	Oct 03, 2023	DS DP		
		7169780*PED	Apr 03, 2024			
		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>RASAGILINE MESYLATE - AZILECT</u>						
N 021641	001	7572834	Dec 05, 2026	DP		
		7815942	Aug 27, 2027	DS DP	U-219	
<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	
		8106183	Feb 02, 2027	DS		
		RE47301	Feb 02, 2027	DP		
<u>REGORAFENIB - STIVARGA</u>						
N 203085	001	7351834	Jun 28, 2022	DS		ODE-139
		8637553	Feb 16, 2031	DS DP		Apr 27, 2024
		8680124	Jun 02, 2030		U-1506	
		9458107	Apr 08, 2031	DP		
		9957232	Jul 09, 2032	DS		
<u>RELUGOLIX - ORGOVYX</u>						
N 214621	001	10350170	Feb 25, 2036	DP		NCE
		10449191	Sep 29, 2037		U-3020	Dec 18, 2025
		10786501	Sep 29, 2037		U-3020	
		7300935	Jan 28, 2024	DS		
		8058280	Jan 28, 2024	DS DP		
		8735401	Feb 04, 2024		U-3019	
<u>REMDESIVIR - VEKLURY</u>						
N 214787	001	10065958	Sep 16, 2031	DS		NCE
		10675296	Jul 10, 2038	DP		Oct 22, 2025
		10695361	Sep 16, 2036		U-2984	
		11007208	Sep 16, 2036		U-2984	
		8008264	Sep 06, 2029	DS DP		
		8318682	Apr 22, 2029	DS DP		
		9724360	Oct 29, 2035	DS DP		
		9949994	Oct 29, 2035	DS		
		RE46762	Apr 22, 2029	DS DP		
<u>REMDESIVIR - VEKLURY</u>						
N 214787	002	10065958	Sep 16, 2031	DS		NCE
		10675296	Jul 10, 2038	DP		Oct 22, 2025
		10695361	Sep 16, 2036		U-2984	
		11007208	Sep 16, 2036		U-2984	
		8008264	Sep 06, 2029	DS DP		
		8318682	Apr 22, 2029	DS DP		
		9724360	Oct 29, 2035	DS DP		

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<u>REMDESIVIR - VEKLURY</u>						
N 214787	002 9949994	Oct 29, 2035	DS			
	RE46762	Apr 22, 2029	DS DP			
<u>REMIMAZOLAM BESYLATE - BYFAVO</u>						
N 212295	001 10052334	Nov 07, 2031		U-2968	NCE	Oct 06, 2025
	10195210	Nov 07, 2031		U-2968		
	10342800	Nov 07, 2031		U-2968		
	10472365	Jul 10, 2027		U-2968		
	10722522	Nov 07, 2031		U-2968		
	9561236	Apr 30, 2033		U-2968		
	9737547	Nov 07, 2031		U-2968		
	9777007	Jul 10, 2027	DP			
	9827251	Nov 07, 2031		U-2968		
	9914738	Jul 10, 2027	DP			
<u>RETAPAMULIN - ALTABAX</u>						
N 022055	001 7875630	Feb 14, 2027	DS			
	8207191	Aug 30, 2024		U-805		
<u>REVEFENACIN - YUPELRI</u>						
N 210598	001 10106503	Mar 10, 2025		U-2440	NCE	Nov 09, 2023
	10343995	Mar 10, 2025		U-2440		
	10550081	Jul 14, 2030	DS			
	11008289	Jul 14, 2030		U-2440		
	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			
	8541451	Aug 25, 2031	DS			
	9765028	Jul 14, 2030	DS			
<u>RIBAVIRIN - REBETOL</u>						
N 021546	001 6790837	Apr 05, 2023	DP			
<u>RIBOCICLIB SUCCINATE - KISQALI</u>						
N 209092	001 10799506	Apr 14, 2036		DP	NCE	Mar 13, 2022
	8324225	Jun 17, 2028	DS DP			
	8415355	Aug 21, 2029	DS DP			
	8685980	May 25, 2030	DS DP			
	8962630	Dec 09, 2029		U-1981		
	8962630	Dec 09, 2029		U-2355		
	8962630	Dec 09, 2029		U-2356		
	8962630	Dec 09, 2029		U-3265		
	8962630	Dec 09, 2029		U-3266		
	9193732	Nov 09, 2031	DS DP			
	9416136	Aug 20, 2029		U-1981		
	9416136	Aug 20, 2029		U-2355		
	9416136	Aug 20, 2029		U-2356		
	9416136	Aug 20, 2029		U-3265		
	9416136	Aug 20, 2029		U-3266		
	9868739	Nov 09, 2031		U-1981		
	9868739	Nov 09, 2031		U-2355		
	9868739	Nov 09, 2031		U-2356		
	9868739	Nov 09, 2031		U-3265		
	9868739	Nov 09, 2031		U-3266		
<u>RIBOFLAVIN 5'-PHOSPHATE SODIUM - PHOTREXA</u>						
N 203324	001				ODE-116	Apr 15, 2023
					ODE-121	Jul 15, 2023
<u>RIBOFLAVIN 5'-PHOSPHATE SODIUM - PHOTREXA VISCOUS IN DEXTRAN 20%</u>						
N 203324	002				ODE-116	Apr 15, 2023
					ODE-121	Jul 15, 2023

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<u>RIFAMYCIN SODIUM - 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	10703763	Feb 27, 2026		U-1708	
		10703763	Feb 27, 2026		U-2847	
		10703763	Feb 27, 2026		U-2848	
		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 021361	002	10314828	Jul 24, 2029		U-1481	
		10335397	Jul 24, 2029		U-2579	
		10456384	Feb 26, 2029		U-2643	
		10456384	Feb 26, 2029		U-2644	
		10703763	Feb 27, 2026		U-1708	
		10703763	Feb 27, 2026		U-2847	
		10703763	Feb 27, 2026		U-2848	
		10709694	Jul 24, 2029		U-2579	
		10765667	Feb 26, 2029		U-2643	
		10765667	Feb 26, 2029		U-2644	
		7045620	Jun 19, 2024	DS		
		7612199	Jun 19, 2024	DS DP		
		7902206	Jun 19, 2024	DS DP		
		7906542	Jun 01, 2025	DS DP		
		7915275	Feb 23, 2025		U-1707	
		7915275	Feb 23, 2025		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	Jul 24, 2029		U-1481	
		9629828	Jul 24, 2029		U-1994	
<u>RILPIVIRINE HYDROCHLORIDE - EDURANT</u>						
N 202022	001	7125879	Apr 21, 2025	DS DP	U-1153	
		7125879	Apr 21, 2025	DS DP	U-1307	
		7125879	Apr 21, 2025	DS DP	U-1740	
		7125879	Apr 21, 2025	DS DP	U-3068	
		7638522	Apr 14, 2023	DP		
		8080551	Apr 11, 2023	DS DP		
		8101629	Aug 09, 2022	DP		

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<u>RILUZOLE - TIGLUTIK KIT</u>						
N 209080	001 8765150	Mar 12, 2029	DP U-2401			
<u>RILUZOLE - EXSERVAN</u>						
N 212640	001 8603514	Apr 03, 2024	DP			
	8765167	Feb 20, 2024	DP			
<u>RIMEGEPANT SULFATE - NURTEC ODT</u>						
N 212728	001 11083724	Mar 25, 2039	DP U-2718		I-865	May 27, 2024
	11083724	Mar 25, 2039	DP U-3142		NCE	Feb 27, 2025
	8314117	Feb 22, 2031	DS DP U-2718			
	8314117	Feb 22, 2031	DS DP U-3142			
	8759372	Feb 25, 2033	DS DP			
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	001 10662188	Feb 18, 2034	DS DP U-2834			
	10662188	Feb 18, 2034	DS DP U-2835			
	7173037	Dec 04, 2026	DS DP			
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	002 10662188	Feb 18, 2034	DS DP U-2834			
	10662188	Feb 18, 2034	DS DP U-2835			
	7173037	Dec 04, 2026	DS DP			
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	003 10662188	Feb 18, 2034	DS DP U-2834			
	10662188	Feb 18, 2034	DS DP U-2835			
	7173037	Dec 04, 2026	DS DP			
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	004 10662188	Feb 18, 2034	DS DP U-2834			
	10662188	Feb 18, 2034	DS DP U-2835			
	7173037	Dec 04, 2026	DS DP			
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	005 10662188	Feb 18, 2034	DS DP U-2834			
	10662188	Feb 18, 2034	DS DP U-2835			
	7173037	Dec 04, 2026	DS DP			
<u>RIPRETINIB - QINLOCK</u>						
N 213973	001 10966966	Aug 12, 2040	U-3153		NCE	May 15, 2025
	11185535	Dec 30, 2040	DP		ODE-298	May 15, 2027
	8188113	Jul 27, 2030	DS DP			
	8461179	Jun 07, 2032	DS DP			
	RE48731	Jun 07, 2032	U-3219			
<u>RISDIPLAM - EVRYSID</u>						
N 213535	001 9586955	Feb 08, 2033	DS DP		NCE	Aug 07, 2025
	9969754	May 11, 2035	DS DP U-1943		ODE-334	Aug 07, 2027
<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 - PERSERIS KIT</u>						
N 210655	001 10010612	Feb 13, 2028	DP			
	10058554	Sep 26, 2026	U-2363			
	10376590	Feb 13, 2028	U-2608			
	10406160	Jun 26, 2026	DP U-2608			
	11013809	Feb 13, 2028	DP U-3135			
	11110093	Nov 05, 2026	DP U-3135			
	9180197	Feb 13, 2028	DP			
	9186413	Feb 13, 2028	U-543			
	9597402	Sep 26, 2026	DP			

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<u>RISPERIDONE - PERSERIS KIT</u>						
N 210655	002	10010612	Feb 13, 2028	DP		
		10058554	Sep 26, 2026	U-2363		
		10376590	Feb 13, 2028	U-2608		
		10406160	Jun 26, 2026	DP U-2608		
		11013809	Feb 13, 2028	DP U-3135		
		11110093	Nov 05, 2026	DP U-3135		
		9180197	Feb 13, 2028	DP		
		9186413	Feb 13, 2028	U-543		
		9597402	Sep 26, 2026	DP		
<u>RIVAROXABAN - XARELTO</u>						
N 022406	001	7157456	Aug 28, 2024	DS DP U-1301	I-810	Oct 11, 2022
		7157456	Aug 28, 2024	DS DP U-1302	I-867	Aug 23, 2024
		7157456*PED	Feb 28, 2025		PED	Apr 11, 2023
		9415053	Nov 13, 2024	DP U-1167	PED	Feb 23, 2025
		9415053	Nov 13, 2024	DP U-2142		
		9415053	Nov 13, 2024	DP U-2640		
		9415053*PED	May 13, 2025			
		9539218	Feb 17, 2034	U-1957		
		9539218	Feb 17, 2034	U-2143		
		9539218	Feb 17, 2034	U-2641		
		9539218*PED	Aug 17, 2034			
<u>RIVAROXABAN - XARELTO</u>						
N 022406	002	7157456	Aug 28, 2024	DS DP U-1301	I-810	Oct 11, 2022
		7157456	Aug 28, 2024	DS DP U-1302	I-867	Aug 23, 2024
		7157456*PED	Feb 28, 2025		PED	Apr 11, 2023
		9415053	Nov 13, 2024	DP U-1200	PED	Feb 23, 2025
		9415053	Nov 13, 2024	DP U-1301		
		9415053	Nov 13, 2024	DP U-1302		
		9415053*PED	May 13, 2025			
		9539218	Feb 17, 2034	U-1953		
		9539218*PED	Aug 17, 2034			
<u>RIVAROXABAN - XARELTO</u>						
N 022406	003	7157456	Aug 28, 2024	DS DP U-1301	I-810	Oct 11, 2022
		7157456	Aug 28, 2024	DS DP U-1302	I-867	Aug 23, 2024
		7157456*PED	Feb 28, 2025		PED	Apr 11, 2023
		9415053	Nov 13, 2024	DP U-1200	PED	Feb 23, 2025
		9415053	Nov 13, 2024	DP U-1301		
		9415053	Nov 13, 2024	DP U-1302		
		9415053*PED	May 13, 2025			
		9539218	Feb 17, 2034	U-1953		
		9539218	Feb 17, 2034	U-1954		
		9539218	Feb 17, 2034	U-1955		
		9539218*PED	Aug 17, 2034			
<u>RIVAROXABAN - XARELTO</u>						
N 022406	004	10828310	Jan 31, 2039	U-3207	I-810	Oct 11, 2022
		10828310	Jan 31, 2039	U-3208	I-867	Aug 23, 2024
		10828310*PED	Jul 31, 2039		PED	Apr 11, 2023
		7157456	Aug 28, 2024	DS DP	PED	Feb 23, 2025
		7157456*PED	Feb 28, 2025			
		9415053	Nov 13, 2024	DP U-2435		
		9415053	Nov 13, 2024	DP U-3205		
		9415053	Nov 13, 2024	DP U-3206		
		9415053*PED	May 13, 2025			
<u>ROFLUMILAST - DALIRESP</u>						
N 022522	001	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>ROFLUMILAST - DALIRESP</u>						
N 022522	002	8431154	Feb 19, 2023	DP		
		8536206	Mar 08, 2024	U-1115		
		8604064	Mar 08, 2024	U-1115		
		8618142	Mar 08, 2024	DP		

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<u>ROFLUMILAST - DALIRESP</u>						
N 022522	002	9468598	Feb 19, 2023	DP		
<u>ROLAPITANT HYDROCHLORIDE - VARUBI</u>						
N 206500	001	7049320	Aug 19, 2028	DS DP	U-1741	
		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	Aug 19, 2028	DS DP	U-1741	
		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>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>ROSUVASTATIN CALCIUM - CRESTOR</u>						
N 021366	002	6858618*PED	Jun 17, 2022		ODE-118	May 27, 2023
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
N 021366	003	6858618*PED	Jun 17, 2022		ODE-118	May 27, 2023
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
N 021366	004	6858618*PED	Jun 17, 2022		ODE-118	May 27, 2023
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
N 021366	005				ODE-118	May 27, 2023
<u>ROSUVASTATIN CALCIUM - EZALLOR SPRINKLE</u>						
N 208647	001	10413543	Feb 12, 2036	DP		
<u>ROSUVASTATIN CALCIUM - EZALLOR SPRINKLE</u>						
N 208647	002	10413543	Feb 12, 2036	DP		
<u>ROSUVASTATIN CALCIUM - EZALLOR SPRINKLE</u>						
N 208647	003	10413543	Feb 12, 2036	DP		
<u>ROSUVASTATIN CALCIUM - EZALLOR SPRINKLE</u>						
N 208647	004	10413543	Feb 12, 2036	DP		
<u>ROTIGOTINE - NEUPRO</u>						
N 021829	001	10130589	Dec 22, 2030	DP		
		10350174	Dec 22, 2030	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	002	10130589	Dec 22, 2030	DP		
		10350174	Dec 22, 2030	DP		
		8246979	Sep 01, 2027	DP	U-1272	
		8246979	Sep 01, 2027	DP	U-1273	

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<u>ROTIGOTINE - NEUPRO</u>						
N 021829	002	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		
		10350174	Dec 22, 2030	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		
		10350174	Dec 22, 2030	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		
		10350174	Dec 22, 2030	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		
		10350174	Dec 22, 2030	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-830	May 15, 2023
		10130636	Aug 17, 2035	U-2101	ODE-126	Dec 19, 2023
		10130636	Aug 17, 2035	U-2273	ODE-168	Apr 06, 2025
		10130636	Aug 17, 2035	U-2830		
		10278974	Feb 10, 2031	DP		
		6495541	Nov 22, 2023	DS DP		
		7351701	Jul 23, 2024	U-2012		
		7351701	Jul 23, 2024	U-2273		
		7351701	Jul 23, 2024	U-2830		
		7531530	Jul 23, 2024	U-2012		
		7531530	Jul 23, 2024	U-2273		
		7531530	Jul 23, 2024	U-2830		
		8071579	Aug 12, 2027	U-2012		
		8071579	Aug 12, 2027	U-2273		
		8071579	Aug 12, 2027	U-2830		
		8143241	Aug 12, 2027	U-2012		
		8143241	Aug 12, 2027	U-2273		
		8143241	Aug 12, 2027	U-2830		
		8754072	Feb 10, 2031	DS DP		
		8859562	Aug 04, 2031	U-2012		
		8859562	Aug 04, 2031	U-2273		
		8859562	Aug 04, 2031	U-2830		
		9045487	Feb 10, 2031	DS DP		
		9861638	Feb 10, 2031	U-2012		
		9861638	Feb 10, 2031	U-2273		
		9987285	Aug 17, 2035	DP		

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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 002	10130636	Aug 17, 2035	U-2012		I-830	May 15, 2023
	10130636	Aug 17, 2035	U-2101		ODE-126	Dec 19, 2023
	10130636	Aug 17, 2035	U-2273		ODE-168	Apr 06, 2025
	10130636	Aug 17, 2035	U-2830			
	10278974	Feb 10, 2031		DP		
	6495541	Nov 22, 2023		DS DP		
	7351701	Jul 23, 2024	U-2012			
	7351701	Jul 23, 2024	U-2273			
	7351701	Jul 23, 2024	U-2830			
	7531530	Jul 23, 2024	U-2012			
	7531530	Jul 23, 2024	U-2273			
	7531530	Jul 23, 2024	U-2830			
	8071579	Aug 12, 2027	U-2012			
	8071579	Aug 12, 2027	U-2273			
	8071579	Aug 12, 2027	U-2830			
	8143241	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2273			
	8143241	Aug 12, 2027	U-2830			
	8754072	Feb 10, 2031		DS DP		
	8859562	Aug 04, 2031	U-2012			
	8859562	Aug 04, 2031	U-2273			
	8859562	Aug 04, 2031	U-2830			
	9045487	Feb 10, 2031		DS DP		
	9861638	Feb 10, 2031	U-2012			
	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-830	May 15, 2023
	10130636	Aug 17, 2035	U-2101		ODE-126	Dec 19, 2023
	10130636	Aug 17, 2035	U-2273		ODE-168	Apr 06, 2025
	10130636	Aug 17, 2035	U-2830			
	10278974	Feb 10, 2031		DP		
	6495541	Nov 22, 2023		DS DP		
	7351701	Jul 23, 2024	U-2012			
	7351701	Jul 23, 2024	U-2273			
	7351701	Jul 23, 2024	U-2830			
	7531530	Jul 23, 2024	U-2012			
	7531530	Jul 23, 2024	U-2273			
	7531530	Jul 23, 2024	U-2830			
	8071579	Aug 12, 2027	U-2012			
	8071579	Aug 12, 2027	U-2273			
	8071579	Aug 12, 2027	U-2830			
	8143241	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2273			
	8143241	Aug 12, 2027	U-2830			
	8754072	Feb 10, 2031		DS DP		
	8859562	Aug 04, 2031	U-2012			
	8859562	Aug 04, 2031	U-2273			
	8859562	Aug 04, 2031	U-2830			
	9045487	Feb 10, 2031		DS DP		
	9861638	Feb 10, 2031	U-2012			
	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				
<u>RUFINAMIDE - BANZEL</u>						
N 021911 002	6740669	Nov 14, 2022		DS DP		
	6740669*PED	May 14, 2023				
<u>RUFINAMIDE - BANZEL</u>						
N 021911 003	6740669	Nov 14, 2022		DS DP		
	6740669*PED	May 14, 2023				

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<u>RUFINAMIDE - BANZEL</u>						
N 201367	001 6740669	Nov 14, 2022	DS DP			
	6740669*PED	May 14, 2023				
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192	001 10016429	Jun 12, 2028		U-3226	I-799	May 24, 2022
	10016429	Jun 12, 2028		U-3230	I-872	Sep 22, 2024
	7598257	Dec 24, 2027	DS DP	U-3227	ODE-238	May 24, 2026
	7598257	Dec 24, 2027	DS DP	U-3228	ODE-373	Sep 22, 2028
	8415362	Dec 24, 2027	DS DP			
	8722693	Jun 12, 2028	DS DP			
	8822481	Jun 12, 2028		U-1573		
	8822481	Jun 12, 2028		U-3226		
	8822481	Jun 12, 2028		U-3227		
	8822481	Jun 12, 2028		U-3228		
	8822481	Jun 12, 2028		U-3230		
	8829013	Jun 12, 2028		U-1201		
	8829013	Jun 12, 2028		U-1622		
	8829013	Jun 12, 2028		U-3227		
	8829013	Jun 12, 2028		U-3228		
	9079912	Dec 12, 2026		U-3226		
	9079912	Dec 12, 2026		U-3227		
	9079912	Dec 12, 2026		U-3228		
	9079912	Dec 12, 2026		U-3230		
	9814722	Dec 12, 2026		U-3226		
	9814722	Dec 12, 2026		U-3230		
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192	002 10016429	Jun 12, 2028		U-3226	I-799	May 24, 2022
	10016429	Jun 12, 2028		U-3230	I-872	Sep 22, 2024
	7598257	Dec 24, 2027	DS DP	U-3227	ODE-238	May 24, 2026
	7598257	Dec 24, 2027	DS DP	U-3228	ODE-373	Sep 22, 2028
	8415362	Dec 24, 2027	DS DP			
	8722693	Jun 12, 2028	DS DP			
	8822481	Jun 12, 2028		U-1573		
	8822481	Jun 12, 2028		U-3226		
	8822481	Jun 12, 2028		U-3227		
	8822481	Jun 12, 2028		U-3228		
	8822481	Jun 12, 2028		U-3230		
	8829013	Jun 12, 2028		U-1201		
	8829013	Jun 12, 2028		U-1622		
	8829013	Jun 12, 2028		U-3227		
	8829013	Jun 12, 2028		U-3228		
	9079912	Dec 12, 2026		U-3226		
	9079912	Dec 12, 2026		U-3227		
	9079912	Dec 12, 2026		U-3228		
	9079912	Dec 12, 2026		U-3230		
	9814722	Dec 12, 2026		U-3226		
	9814722	Dec 12, 2026		U-3230		
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192	003 10016429	Jun 12, 2028		U-3226	I-799	May 24, 2022
	10016429	Jun 12, 2028		U-3230	ODE-238	May 24, 2026
	7598257	Dec 24, 2027	DS DP	U-3227	ODE-373	Sep 22, 2028
	7598257	Dec 24, 2027	DS DP	U-3228		
	8415362	Dec 24, 2027	DS DP			
	8722693	Jun 12, 2028	DS DP			
	8822481	Jun 12, 2028		U-1573		
	8822481	Jun 12, 2028		U-3226		
	8822481	Jun 12, 2028		U-3227		
	8822481	Jun 12, 2028		U-3228		
	8822481	Jun 12, 2028		U-3230		
	8829013	Jun 12, 2028		U-1201		
	8829013	Jun 12, 2028		U-1622		
	8829013	Jun 12, 2028		U-3227		
	8829013	Jun 12, 2028		U-3228		
	9079912	Dec 12, 2026		U-3226		
	9079912	Dec 12, 2026		U-3227		
	9079912	Dec 12, 2026		U-3228		
	9079912	Dec 12, 2026		U-3230		
	9814722	Dec 12, 2026		U-3226		

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<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192	003	9814722				
		Dec 12, 2026	U-3230			
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192	004	10016429			I-799	May 24, 2022
		10016429			ODE-238	May 24, 2026
		7598257	DS DP	U-3227	ODE-373	Sep 22, 2028
		7598257	DS DP	U-3228		
		8415362	DS DP			
		8722693	DS DP			
		8822481		U-1573		
		8822481		U-3226		
		8822481		U-3227		
		8822481		U-3228		
		8822481		U-3230		
		8829013		U-1201		
		8829013		U-1622		
		8829013		U-3227		
		8829013		U-3228		
		9079912		U-3226		
		9079912		U-3227		
		9079912		U-3228		
		9079912		U-3230		
		9814722		U-3226		
		9814722		U-3230		
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192	005	10016429			I-799	May 24, 2022
		10016429			ODE-238	May 24, 2026
		7598257	DS DP	U-3227	ODE-373	Sep 22, 2028
		7598257	DS DP	U-3228		
		8415362	DS DP			
		8722693	DS DP			
		8822481		U-1573		
		8822481		U-3226		
		8822481		U-3227		
		8822481		U-3228		
		8822481		U-3230		
		8829013		U-1201		
		8829013		U-1622		
		8829013		U-3227		
		8829013		U-3228		
		9079912		U-3226		
		9079912		U-3227		
		9079912		U-3228		
		9079912		U-3230		
		9814722		U-3226		
		9814722		U-3230		
<u>RUXOLITINIB PHOSPHATE - OPZELURA</u>						
N 215309	001	10610530			NP	Sep 21, 2024
		10639310				
		10758543		DP		
		10869870		U-3229		
		7598257	DS DP			
		8415362	DS DP			
		8722693	DS DP			
		8822481		U-3229		
		9079912		U-3229		
		9974790		U-3229		
<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620	001	11058667			M-82	Feb 16, 2024
		11135192			NPP	Oct 01, 2022
		7468390		DP	PED	Apr 01, 2023
		7468390*PED				
		8101659		DP		
		8101659*PED				
		8404744		DP		
		8404744*PED				
		8796331		U-1723		

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<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620	001	8796331*PED	Jul 14, 2023			
		8877938	May 27, 2027	DS DP		
		8877938*PED	Nov 27, 2027			
		9388134	Nov 08, 2026		U-1723	
		9388134*PED	May 08, 2027			
		9517226	Aug 22, 2033		U-3084	
		9937143	Aug 22, 2033		U-3084	
<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620	002	11058667	May 09, 2036		U-3170	
		11135192	Aug 22, 2033		U-3084	
		7468390	Nov 27, 2023	DP		M-82 NPP PED Feb 16, 2024 Oct 01, 2022 Apr 01, 2023
		7468390*PED	May 27, 2024			
		8101659	Jan 15, 2025	DP		
		8101659*PED	Jul 15, 2025			
		8404744	Jan 14, 2023	DP		
		8404744*PED	Jul 14, 2023			
		8796331	Jan 14, 2023		U-1723	
		8796331*PED	Jul 14, 2023			
		8877938	May 27, 2027	DS DP		
		8877938*PED	Nov 27, 2027			
		9388134	Nov 08, 2026		U-1723	
		9388134*PED	May 08, 2027			
		9517226	Aug 22, 2033		U-3084	
		9937143	Aug 22, 2033		U-3084	
<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620	003	11058667	May 09, 2036		U-3170	
		11135192	Aug 22, 2033		U-3084	
		7468390	Nov 27, 2023	DP		M-82 NPP PED Feb 16, 2024 Oct 01, 2022 Apr 01, 2023
		7468390*PED	May 27, 2024			
		8101659	Jan 15, 2025	DP		
		8101659*PED	Jul 15, 2025			
		8404744	Jan 14, 2023	DP		
		8404744*PED	Jul 14, 2023			
		8796331	Jan 14, 2023		U-1723	
		8796331*PED	Jul 14, 2023			
		8877938	May 27, 2027	DS DP		
		8877938*PED	Nov 27, 2027			
		9388134	Nov 08, 2026		U-1723	
		9388134*PED	May 08, 2027			
		9517226	Aug 22, 2033		U-3084	
		9937143	Aug 22, 2033		U-3084	
<u>SAFINAMIDE MESYLATE - XADAGO</u>						
N 207145	001	8076515	Dec 10, 2028	DS DP	U-1993	
		8278485	Jun 08, 2027	DS	U-1993	NCE Mar 21, 2022
		8283380	Mar 21, 2031		U-1993	
<u>SAFINAMIDE MESYLATE - XADAGO</u>						
N 207145	002	8076515	Dec 10, 2028	DS DP	U-1993	
		8278485	Jun 08, 2027	DS	U-1993	NCE Mar 21, 2022
		8283380	Mar 21, 2031		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			
		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	

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<u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u>						
N 022181	001	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>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	
<u>SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA</u>						
N 022350	001	7951400	Nov 30, 2028		DP	
		RE44186	Jul 31, 2023	DS DP	U-1837	
		RE44186	Jul 31, 2023	DS DP	U-995	
<u>SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA</u>						
N 022350	002	7951400	Nov 30, 2028		DP	
		RE44186	Jul 31, 2023	DS DP	U-1837	
		RE44186	Jul 31, 2023	DS DP	U-995	
<u>SECNIDAZOLE - SOLOSEC</u>						
N 209363	001	10335390	Sep 04, 2035		U-2583	I-866 Jun 30, 2024
		10682338	Sep 04, 2035		U-2583	NCE Sep 15, 2022
		10849884	Sep 04, 2035	DP	U-2583	GAIN Sep 15, 2027
		10849884	Sep 04, 2035	DP	U-3169	
		10857133	Sep 04, 2035	DP	U-2583	
		11000507	Sep 04, 2035	DP	U-2583	
		11000507	Sep 04, 2035	DP	U-3169	
		11000508	Sep 04, 2035	DP	U-2583	
		11000508	Sep 04, 2035	DP	U-3169	
		11020377	Sep 04, 2035	DP	U-2583	
		11020377	Sep 04, 2035	DP	U-3169	

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<u>SECNIDAZOLE - SOLOSEC</u>						
N 209363	001	10335390	Sep 04, 2035	U-2583	I-866	Jun 30, 2024
		10682338	Sep 04, 2035	U-2583	NCE	Sep 15, 2022
		10849884	Sep 04, 2035	DP U-2583	GAIN	Sep 15, 2027
		10849884	Sep 04, 2035	DP U-3169		
		10857133	Sep 04, 2035	DP U-2583		
		11000507	Sep 04, 2035	DP U-2583		
		11000507	Sep 04, 2035	DP U-3169		
		11000508	Sep 04, 2035	DP U-2583		
		11000508	Sep 04, 2035	DP U-3169		
		11020377	Sep 04, 2035	DP U-2583		
		11020377	Sep 04, 2035	DP U-3169		
<u>SELENIOS ACID - SELENIOS ACID</u>						
N 209379	001				NCE	Apr 30, 2024
<u>SELENIOS ACID - SELENIOS ACID</u>						
N 209379	002				NCE	Apr 30, 2024
<u>SELENIOS ACID - SELENIOS ACID</u>						
N 209379	003				NCE	Apr 30, 2024
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	001	10821108	Dec 01, 2036	DP U-2992	ODE-106	Dec 21, 2022
		10828298	Dec 01, 2036	DP U-2991		
		7205302	Oct 31, 2026	DS DP U-1797		
		8791122	Aug 01, 2030	DS DP		
		9173881	Aug 12, 2029	U-1798		
		9284280	Jun 25, 2030	U-1831		
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	002	10821108	Dec 01, 2036	DP U-2992	ODE-106	Dec 21, 2022
		10828298	Dec 01, 2036	DP U-2991		
		7205302	Oct 31, 2026	DS DP U-1797		
		8791122	Aug 01, 2030	DS DP		
		9173881	Aug 12, 2029	U-1798		
		9284280	Jun 25, 2030	U-1831		
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	003	10821108	Dec 01, 2036	DP U-2992	ODE-106	Dec 21, 2022
		10828298	Dec 01, 2036	DP U-2991		
		7205302	Oct 31, 2026	DS DP U-1797		
		8791122	Aug 01, 2030	DS DP		
		9173881	Aug 12, 2029	U-1798		
		9284280	Jun 25, 2030	U-1831		
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	004	10821108	Dec 01, 2036	DP U-2992	ODE-106	Dec 21, 2022
		10828298	Dec 01, 2036	DP U-2991		
		7205302	Oct 31, 2026	DS DP U-1797		
		8791122	Aug 01, 2030	DS DP		
		9173881	Aug 12, 2029	U-1798		
		9284280	Jun 25, 2030	U-1831		
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	005	10821108	Dec 01, 2036	DP U-2992	ODE-106	Dec 21, 2022
		10828298	Dec 01, 2036	DP U-2991		
		7205302	Oct 31, 2026	DS DP U-1797		
		8791122	Aug 01, 2030	DS DP		
		9173881	Aug 12, 2029	U-1798		
		9284280	Jun 25, 2030	U-1831		
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	006	10821108	Dec 01, 2036	DP U-2992	ODE-106	Dec 21, 2022
		10828298	Dec 01, 2036	DP U-2991		
		7205302	Oct 31, 2026	DS DP U-1797		
		8791122	Aug 01, 2030	DS DP		
		9173881	Aug 12, 2029	U-1798		
		9284280	Jun 25, 2030	U-1831		

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<u>SELEXIPAG - UPTRAVI</u>						
N 207947 007	10821108	Dec 01, 2036	DP U-2992		ODE-106	Dec 21, 2022
	10828298	Dec 01, 2036	DP U-2991			
	7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELEXIPAG - UPTRAVI</u>						
N 207947 008	10821108	Dec 01, 2036	DP U-2992		ODE-106	Dec 21, 2022
	10828298	Dec 01, 2036	DP U-2991			
	7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELEXIPAG - UPTRAVI</u>						
N 214275 001	7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELINEXOR - XPOVIO</u>						
N 212306 001	10519139	Aug 14, 2035	DS DP U-2584		I-837	Jun 22, 2023
	10519139	Aug 14, 2035	DS DP U-2855		NCE	Jul 03, 2024
	10519139	Aug 14, 2035	DS DP U-3018		ODE-257	Jul 03, 2026
	10544108	Jul 26, 2032	U-2584		ODE-310	Jun 22, 2027
	10544108	Jul 26, 2032	U-3018		ODE-346	Dec 18, 2027
	11034660	Jul 26, 2032	U-2584			
	11034660	Jul 26, 2032	U-3018			
	8999996	Sep 15, 2032	DS DP			
	9079865	Jul 26, 2032	U-2584			
	9079865	Jul 26, 2032	U-2855			
	9079865	Jul 26, 2032	U-3018			
	9714226	Jul 26, 2032	DS DP			
<u>SELINEXOR - XPOVIO</u>						
N 212306 002	10519139	Aug 14, 2035	DS DP U-2584		ODE*	Jul 03, 2026
	10519139	Aug 14, 2035	DS DP U-2855		ODE*	Jun 22, 2027
	10519139	Aug 14, 2035	DS DP U-3018		ODE*	Dec 18, 2027
	10544108	Jul 26, 2032	U-2584			
	10544108	Jul 26, 2032	U-3018			
	11034660	Jul 26, 2032	U-2584			
	11034660	Jul 26, 2032	U-3018			
	8999996	Sep 15, 2032	DS DP			
	9079865	Jul 26, 2032	U-2584			
	9079865	Jul 26, 2032	U-2855			
	9079865	Jul 26, 2032	U-3018			
	9714226	Jul 26, 2032	DS DP			
<u>SELINEXOR - XPOVIO</u>						
N 212306 003	10519139	Aug 14, 2035	DS DP U-2584		ODE*	Jul 03, 2026
	10519139	Aug 14, 2035	DS DP U-2855		ODE*	Jun 22, 2027
	10519139	Aug 14, 2035	DS DP U-3018		ODE*	Dec 18, 2027
	10544108	Jul 26, 2032	U-2584			
	10544108	Jul 26, 2032	U-3018			
	11034660	Jul 26, 2032	U-2584			
	11034660	Jul 26, 2032	U-3018			
	8999996	Sep 15, 2032	DS DP			
	9079865	Jul 26, 2032	U-2584			
	9079865	Jul 26, 2032	U-2855			
	9079865	Jul 26, 2032	U-3018			
	9714226	Jul 26, 2032	DS DP			
<u>SELINEXOR - XPOVIO</u>						
N 212306 004	10519139	Aug 14, 2035	DS DP U-2584		ODE*	Jul 03, 2026
	10519139	Aug 14, 2035	DS DP U-2855		ODE*	Jun 22, 2027
	10519139	Aug 14, 2035	DS DP U-3018		ODE*	Dec 18, 2027
	10544108	Jul 26, 2032	U-2584			
	10544108	Jul 26, 2032	U-3018			
	11034660	Jul 26, 2032	U-2584			

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<u>SELINEXOR - XPOVIO</u>						
N 212306 004	11034660	Jul 26, 2032		U-3018		
	8999996	Sep 15, 2032	DS DP			
	9079865	Jul 26, 2032		U-2584		
	9079865	Jul 26, 2032		U-2855		
	9079865	Jul 26, 2032		U-3018		
	9714226	Jul 26, 2032	DS DP			
<u>SELPERCATINIB - RETEVMO</u>						
N 213246 001	10112942	Oct 10, 2037	DS DP		NCE	May 08, 2025
	10137124	Oct 10, 2037		U-2826	ODE-301	May 08, 2027
	10137124	Oct 10, 2037		U-2827	ODE-302	May 08, 2027
	10137124	Oct 10, 2037		U-2828	ODE-303	May 08, 2027
	10172851	Oct 10, 2037		U-2826		
	10172851	Oct 10, 2037		U-2827		
	10172851	Oct 10, 2037		U-2828		
	10584124	Oct 10, 2038	DS	U-2826		
	10584124	Oct 10, 2038	DS	U-2827		
	10584124	Oct 10, 2038	DS	U-2828		
	10786489	Oct 10, 2038	DP	U-2971		
	10786489	Oct 10, 2038	DP	U-2972		
	10786489	Oct 10, 2038	DP	U-2973		
	10786489	Oct 10, 2038	DP	U-2974		
	10786489	Oct 10, 2038	DP	U-2975		
	10786489	Oct 10, 2038	DP	U-2976		
	10786489	Oct 10, 2038	DP	U-2977		
<u>SELPERCATINIB - RETEVMO</u>						
N 213246 002	10112942	Oct 10, 2037	DS DP		NCE	May 08, 2025
	10137124	Oct 10, 2037		U-2826	ODE-301	May 08, 2027
	10137124	Oct 10, 2037		U-2827	ODE-302	May 08, 2027
	10137124	Oct 10, 2037		U-2828	ODE-303	May 08, 2027
	10172851	Oct 10, 2037		U-2826		
	10172851	Oct 10, 2037		U-2827		
	10172851	Oct 10, 2037		U-2828		
	10584124	Oct 10, 2038	DS	U-2826		
	10584124	Oct 10, 2038	DS	U-2827		
	10584124	Oct 10, 2038	DS	U-2828		
	10786489	Oct 10, 2038	DP	U-2971		
	10786489	Oct 10, 2038	DP	U-2972		
	10786489	Oct 10, 2038	DP	U-2973		
	10786489	Oct 10, 2038	DP	U-2974		
	10786489	Oct 10, 2038	DP	U-2975		
	10786489	Oct 10, 2038	DP	U-2976		
	10786489	Oct 10, 2038	DP	U-2977		
<u>SELUMETINIB SULFATE - KOSELUGO</u>						
N 213756 001	7425637	Apr 11, 2024	DS		NCE	Apr 10, 2025
	8178693	Mar 13, 2023	DS DP		ODE-288	Apr 10, 2027
	9156795	Dec 12, 2026	DS DP			
	9562017	Dec 12, 2026	DS	U-2800		
<u>SELUMETINIB SULFATE - KOSELUGO</u>						
N 213756 002	7425637	Apr 11, 2024	DS		NCE	Apr 10, 2025
	8178693	Mar 13, 2023	DS DP		ODE-288	Apr 10, 2027
	9156795	Dec 12, 2026	DS DP			
	9562017	Dec 12, 2026	DS	U-2800		
<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637 001	10220155	Jul 17, 2026	DP		I-822	Jan 16, 2023
	10335462	Jun 21, 2033		U-2580	NCE	Dec 05, 2022
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	11097063	Jul 17, 2026	DP			
	6899699	Jan 02, 2022	DP			
	7762994	May 23, 2024	DP			
	8114833	Aug 13, 2025	DP			
	8129343	Dec 05, 2031	DS DP	U-2202		
	8536122	Mar 20, 2026	DS DP	U-2202		
	8579869	Jun 30, 2023	DP			
	8672898	Jan 02, 2022	DP			

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<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>SEMAGLUTIDE - OZEMPIC</u>						
N 209637	002	10220155	Jul 17, 2026	DP	I-822	Jan 16, 2023
		10335462	Jun 21, 2033	U-2580	NCE	Dec 05, 2022
		10357616	Jan 20, 2026	DP		
		10376652	Jan 20, 2026	DP		
		11097063	Jul 17, 2026	DP		
		6899699	Jan 02, 2022	DP		
		7762994	May 23, 2024	DP		
		8114833	Aug 13, 2025	DP		
		8129343	Dec 05, 2031	DS DP U-2202		
		8536122	Mar 20, 2026	DS DP U-2202		
		8579869	Jun 30, 2023	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 29, 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>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	001	10086047	Dec 16, 2031	DP	M-252	Jan 16, 2023
		10278923	May 02, 2034	U-2628	NCE	Dec 05, 2022
		10933120	Mar 15, 2033	DP	NP	Sep 20, 2022
		10960052	Dec 16, 2031	DP		
		8129343	Dec 05, 2031	DS DP U-2628		
		8536122	Mar 20, 2026	DS DP U-2628		
		9278123	Dec 16, 2031	DP U-2628		
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	002	10086047	Dec 16, 2031	DP	M-252	Jan 16, 2023
		10278923	May 02, 2034	U-2628	NCE	Dec 05, 2022
		10933120	Mar 15, 2033	DP	NP	Sep 20, 2022
		10960052	Dec 16, 2031	DP		
		8129343	Dec 05, 2031	DS DP U-2628		
		8536122	Mar 20, 2026	DS DP U-2628		
		9278123	Dec 16, 2031	DP U-2628		
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	003	10086047	Dec 16, 2031	DP	M-252	Jan 16, 2023
		10278923	May 02, 2034	U-2628	NCE	Dec 05, 2022
		10933120	Mar 15, 2033	DP	NP	Sep 20, 2022
		10960052	Dec 16, 2031	DP		
		8129343	Dec 05, 2031	DS DP U-2628		
		8536122	Mar 20, 2026	DS DP U-2628		
		9278123	Dec 16, 2031	DP U-2628		
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256	001	10888605	Aug 24, 2038	DP U-3162	NCE	Dec 05, 2022
		8129343	Dec 05, 2031	DS DP	NP	Jun 04, 2024
		8536122	Mar 20, 2026	DS DP		
		9764003	Jun 21, 2033	U-3161		

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<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256 002	10888605	Aug 24, 2038	DP U-3162		NCE	Dec 05, 2022
	8129343	Dec 05, 2031	DS DP		NP	Jun 04, 2024
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256 003	10888605	Aug 24, 2038	DP U-3162		NCE	Dec 05, 2022
	8129343	Dec 05, 2031	DS DP		NP	Jun 04, 2024
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256 004	10888605	Aug 24, 2038	DP U-3162		NCE	Dec 05, 2022
	8129343	Dec 05, 2031	DS DP		NP	Jun 04, 2024
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256 005	10888605	Aug 24, 2038	DP U-3162		NCE	Dec 05, 2022
	8129343	Dec 05, 2031	DS DP		NP	Jun 04, 2024
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SETMELANOTIDE ACETATE - IMCIVREE</u>						
N 213793 001	11129869	Jul 04, 2034	DP		NCE	Nov 25, 2025
	8039435	Oct 13, 2027	DS DP		ODE-336	Nov 25, 2027
	9458195	Oct 13, 2027	DS DP			
<u>SEVELAMER CARBONATE - RENVELA</u>						
N 022127 001	7985418	Oct 27, 2025	DP			
<u>SEVELAMER CARBONATE - RENVELA</u>						
N 022318 001	9095509	Dec 06, 2030	DP			
<u>SEVELAMER CARBONATE - RENVELA</u>						
N 022318 002	9095509	Dec 06, 2030	DP			
<u>SIMEPREVIR SODIUM - OLYSIO</u>						
N 205123 001	7671032	May 19, 2025	DS DP			
	8148399	Sep 05, 2029	DS DP U-1467			
	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	10300041	Apr 26, 2027	DP			
	9597289	Feb 23, 2030	DP			
<u>SIMVASTATIN - FLOLIPID</u>						
N 206679 002	10300041	Apr 26, 2027	DP			
	9597289	Feb 23, 2030	DP			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343 001	6699871	Jul 26, 2022	DS DP U-1188			
	6699871*PED	Jan 26, 2023				
	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			
	7125873*PED	Jan 26, 2023				
	7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				
	8168637	Jun 26, 2022	DP U-1188			

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<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343 002	6699871	Jul 26, 2022	DS DP U-1188			
	6699871*PED	Jan 26, 2023				
	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			
	7125873*PED	Jan 26, 2023				
	7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				
	8168637	Jun 26, 2022	DP U-1188			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343 003	6699871	Jul 26, 2022	DS DP U-1188			
	6699871*PED	Jan 26, 2023				
	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			
	7125873*PED	Jan 26, 2023				
	7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				
	8168637	Jun 26, 2022	DP U-1188			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343 004	6699871	Jul 26, 2022	DS DP U-1188			
	6699871*PED	Jan 26, 2023				
	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			
	7125873*PED	Jan 26, 2023				
	7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				
	8168637	Jun 26, 2022	DP U-1188			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343 005	6699871	Jul 26, 2022	DS DP U-1188			
	6699871*PED	Jan 26, 2023				
	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			
	7125873*PED	Jan 26, 2023				
	7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				
	8168637	Jun 26, 2022	DP U-1188			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343 006	6699871	Jul 26, 2022	DS DP U-1188			
	6699871*PED	Jan 26, 2023				
	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			
	7125873*PED	Jan 26, 2023				
	7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				
	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	10434059	Nov 18, 2022	DP U-172			
	7858662	Oct 02, 2026	DP U-172			
	9770406	Jul 12, 2025	DP U-172			
<u>SIPONIMOD FUMARIC ACID - MAYZENT</u>						
N 209884 001	7939519	May 19, 2024	DS DP		NCE	Mar 26, 2024
	8492441	Nov 30, 2030	U-2511			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SIPONIMOD FUMARIC ACID - MAYZENT</u>						
N 209884 001	7939519	May 19, 2024	DS DP		NCE	Mar 26, 2024
	8492441	Nov 30, 2030		U-2511		
<u>SIPONIMOD FUMARIC ACID - MAYZENT</u>						
N 209884 002	7939519	May 19, 2024	DS DP		NCE	Mar 26, 2024
	8492441	Nov 30, 2030		U-2511		
<u>SIPONIMOD FUMARIC ACID - MAYZENT</u>						
N 209884 003	7939519	May 19, 2024	DS DP		NCE	Mar 26, 2024
	8492441	Nov 30, 2030		U-2511		
<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>SIROLIMUS - FYARRO</u>						
N 213312 001	10206887	Apr 15, 2030	DP		NP	Nov 22, 2024
	10705070	Mar 05, 2036	DP			
	10973806	Jun 29, 2036		U-3258		
	8911786	Feb 14, 2029	DP	U-3259		
<u>SITAGLIPTIN PHOSPHATE - JANUVIA</u>						
N 021995 001	6699871	Jul 26, 2022	DS DP	U-774	M-187	Dec 04, 2023
	7125873	Jul 26, 2022		U-1036	M-244	Aug 12, 2022
	7125873	Jul 26, 2022		U-1037	PED	Feb 12, 2023
	7125873	Jul 26, 2022		U-1038	PED	Jun 04, 2024
	7125873	Jul 26, 2022		U-775		
	7326708	Nov 24, 2026	DS DP	U-802		
	7326708*PED	May 24, 2027				
<u>SITAGLIPTIN PHOSPHATE - JANUVIA</u>						
N 021995 002	6699871	Jul 26, 2022	DS DP	U-774	M-187	Dec 04, 2023
	7125873	Jul 26, 2022		U-1036	M-244	Aug 12, 2022
	7125873	Jul 26, 2022		U-1037	PED	Feb 12, 2023
	7125873	Jul 26, 2022		U-1038	PED	Jun 04, 2024
	7125873	Jul 26, 2022		U-775		
	7326708	Nov 24, 2026	DS DP	U-802		
	7326708*PED	May 24, 2027				
<u>SITAGLIPTIN PHOSPHATE - JANUVIA</u>						
N 021995 003	6699871	Jul 26, 2022	DS DP	U-774	M-187	Dec 04, 2023
	7125873	Jul 26, 2022		U-1036	M-244	Aug 12, 2022
	7125873	Jul 26, 2022		U-1037	PED	Feb 12, 2023
	7125873	Jul 26, 2022		U-1038	PED	Jun 04, 2024
	7125873	Jul 26, 2022		U-775		
	7326708	Nov 24, 2026	DS DP	U-802		
	7326708*PED	May 24, 2027				
<u>SODIUM ACETATE - SODIUM ACETATE</u>						
A 214805 001					CGT	Jan 03, 2022
<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			

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<u>SODIUM OXYBATE - XYREM</u>						
N 021196 001	10213400	Mar 15, 2033	U-2499		ODE-231	Oct 26, 2025
	10864181	Mar 15, 2033	U-1532		PED	Apr 26, 2026
	7668730	Jun 16, 2024	U-1110	Y		
	7668730*PED	Dec 16, 2024				
	8731963	Dec 17, 2022	U-1110			
	8731963*PED	Jun 17, 2023				
	8772306	Mar 15, 2033	U-1532			
	8772306*PED	Sep 15, 2033				
	9050302	Mar 15, 2033	U-1532			
	9050302*PED	Sep 15, 2033				
	9486426	Mar 15, 2033	U-1532			
	9486426*PED	Sep 15, 2033				
<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	10300087	Oct 14, 2035	DS U-2312		M-261	Apr 24, 2023
	10335432	Feb 10, 2032	U-2312		NCE	May 18, 2023
	10398730	Feb 10, 2032	U-2312			
	10413569	Feb 10, 2032	DS			
	10695365	Oct 22, 2033	DS			
	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			
	9913860	Oct 22, 2033	DS U-2312			
<u>SODIUM ZIRCONIUM CYCLOSILICATE - LOKELMA</u>						
N 207078 002	10300087	Oct 14, 2035	DS U-2312		M-261	Apr 24, 2023
	10398730	Feb 10, 2032	U-2312		NCE	May 18, 2023
	10413569	Feb 10, 2032	DS			
	10695365	Oct 22, 2033	DS			
	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			
	9913860	Oct 22, 2033	DS U-2312			
<u>SOFOSEBUVIR - SOVALDI</u>						
N 204671 001	7964580	Mar 26, 2029	DS DP U-1470		ODE*	Aug 28, 2026
	7964580*PED	Sep 26, 2029			ODE-135	Apr 07, 2024
	8334270	Mar 21, 2028	DS DP U-1470		PED	Oct 07, 2024
	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9549941	Mar 26, 2029	U-1958			
	9549941*PED	Sep 26, 2029				

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<u>SOFOSEBUVIR - SOVALDI</u>						
N 204671	002	7964580	Mar 26, 2029	DS DP U-1470	ODE*	Apr 07, 2024
		7964580*PED	Sep 26, 2029		ODE*	Aug 28, 2026
		8334270	Mar 21, 2028	DS DP U-1470	PED	Oct 07, 2024
		8334270*PED	Sep 21, 2028			
		8580765	Mar 21, 2028	DS DP U-1470		
		8580765*PED	Sep 21, 2028			
		8618076	Dec 11, 2030	DS DP U-1470		
		8618076*PED	Jun 11, 2031			
		8633309	Mar 26, 2029	DS DP U-1470		
		8633309*PED	Sep 26, 2029			
		8889159	Mar 26, 2029	DP U-1470		
		8889159*PED	Sep 26, 2029			
		9085573	Mar 21, 2028	DS DP U-1470		
		9085573*PED	Sep 21, 2028			
		9284342	Sep 13, 2030	DS DP U-1470		
		9284342*PED	Mar 13, 2031			
<u>SOFOSEBUVIR - SOVALDI</u>						
N 212480	001	7964580	Mar 26, 2029	DS DP U-1470	ODE-258	Aug 28, 2026
		7964580*PED	Sep 26, 2029			
		8334270	Mar 21, 2028	DS DP U-1470		
		8334270*PED	Sep 21, 2028			
		8580765	Mar 21, 2028	DS DP U-1470		
		8580765*PED	Sep 21, 2028			
		8618076	Dec 11, 2030	DS DP U-1470		
		8618076*PED	Jun 11, 2031			
		8633309	Mar 26, 2029	DS DP U-1470		
		8633309*PED	Sep 26, 2029			
		8889159	Mar 26, 2029	DP U-1470		
		8889159*PED	Sep 26, 2029			
		9085573	Mar 21, 2028	DS DP U-1470		
		9085573*PED	Sep 21, 2028			
		9284342	Sep 13, 2030	DS DP U-1470		
		9284342*PED	Mar 13, 2031			
<u>SOFOSEBUVIR - SOVALDI</u>						
N 212480	002	7964580	Mar 26, 2029	DS DP U-1470	ODE-258	Aug 28, 2026
		7964580*PED	Sep 26, 2029			
		8334270	Mar 21, 2028	DS DP U-1470		
		8334270*PED	Sep 21, 2028			
		8580765	Mar 21, 2028	DS DP U-1470		
		8580765*PED	Sep 21, 2028			
		8618076	Dec 11, 2030	DS DP U-1470		
		8618076*PED	Jun 11, 2031			
		8633309	Mar 26, 2029	DS DP U-1470		
		8633309*PED	Sep 26, 2029			
		8889159	Mar 26, 2029	DP U-1470		
		8889159*PED	Sep 26, 2029			
		9085573	Mar 21, 2028	DS DP U-1470		
		9085573*PED	Sep 21, 2028			
		9284342	Sep 13, 2030	DS DP U-1470		
		9284342*PED	Mar 13, 2031			
<u>SOFOSEBUVIR; VELPATASVIR - EPCLUSA</u>						
N 208341	001	10086011	Jan 30, 2034	U-1470	D-177	Nov 15, 2022
		10086011*PED	Jul 30, 2034		M-264	Jul 14, 2023
		11116783	Jan 30, 2034	DP U-1470	NPP	Mar 19, 2023
		11116783*PED	Jul 30, 2034		ODE-293	Mar 19, 2027
		7964580	Mar 26, 2029	DS DP U-1470	ODE-376	Jun 10, 2028
		7964580*PED	Sep 26, 2029		PED	May 15, 2023
		8334270	Mar 21, 2028	DS DP U-1470	PED	Sep 19, 2023
		8334270*PED	Sep 21, 2028		PED	Jan 14, 2024
		8575135	Nov 16, 2032	DS DP U-1470	PED	Sep 19, 2027
		8575135*PED	May 16, 2033		PED	Dec 10, 2028
		8580765	Mar 21, 2028	DS DP U-1470		
		8580765*PED	Sep 21, 2028			
		8618076	Dec 11, 2030	DS DP U-1470		
		8618076*PED	Jun 11, 2031			
		8633309	Mar 26, 2029	DS DP U-1470		
		8633309*PED	Sep 26, 2029			

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<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 208341 001	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-1470			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-1470			
	8940718*PED	May 16, 2033				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9757406	Jan 30, 2034	DP			
	9757406*PED	Jul 30, 2034				
<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 208341 002	10086011	Jan 30, 2034	U-1470		M-264	Jul 14, 2023
	10086011*PED	Jul 30, 2034			NS	Mar 19, 2023
	11116783	Jan 30, 2034	DP U-1470		ODE-293	Mar 19, 2027
	11116783*PED	Jul 30, 2034			ODE-376	Jun 10, 2028
	7964580	Mar 26, 2029	DS DP U-1470		PED	Sep 19, 2023
	7964580*PED	Sep 26, 2029			PED	Jan 14, 2024
	8334270	Mar 21, 2028	DS DP U-1470		PED	Sep 19, 2027
	8334270*PED	Sep 21, 2028			PED	Dec 10, 2028
	8575135	Nov 16, 2032	DS DP U-1470			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-1470			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-1470			
	8940718*PED	May 16, 2033				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9757406	Jan 30, 2034	DP			
	9757406*PED	Jul 30, 2034				
<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 214187 001	11116783	Jan 30, 2034	DP U-1470		ODE-376	Jun 10, 2028
	11116783*PED	Jul 30, 2034			PED	Dec 10, 2028
	7964580	Mar 26, 2029	DS DP U-1470			
	7964580*PED	Sep 26, 2029				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-1470			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-1470			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-1470			

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<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 214187 001	8940718*PED	May 16, 2033				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 214187 002	11116783	Jan 30, 2034	DP U-1470		ODE-376	Jun 10, 2028
	11116783*PED	Jul 30, 2034			PED	Dec 10, 2028
	7964580	Mar 26, 2029	DS DP U-1470			
	7964580*PED	Sep 26, 2029				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-1470			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-1470			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-1470			
	8940718*PED	May 16, 2033				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
<u>SOFOSBUVIR; VELPATASVIR; VOXILAPREVIR - VOSEVI</u>						
N 209195 001	10912814	Jun 01, 2037	DP		NCE	Jul 18, 2022
	11116783	Jan 30, 2034	DP U-2039			
	11116783	Jan 30, 2034	DP U-2040			
	11116783*PED	Jul 30, 2034				
	7964580	Mar 26, 2029	DS DP U-2039			
	7964580	Mar 26, 2029	DS DP U-2040			
	7964580*PED	Sep 26, 2029				
	8334270	Mar 21, 2028	DS DP U-2039			
	8334270	Mar 21, 2028	DS DP U-2040			
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-2039			
	8575135	Nov 16, 2032	DS DP U-2040			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-2039			
	8580765	Mar 21, 2028	DS DP U-2040			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-2039			
	8618076	Dec 11, 2030	DS DP U-2040			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-2039			
	8633309	Mar 26, 2029	DS DP U-2040			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	DS DP U-2039			
	8735372	Mar 21, 2028	DS DP U-2040			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DS DP U-2039			
	8889159	Mar 26, 2029	DS DP U-2040			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-2039			
	8921341	Nov 16, 2032	DS DP U-2040			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-2039			
	8940718	Nov 16, 2032	DS DP U-2040			
	8940718*PED	May 16, 2033				

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<u>SOFOSBUVIR; VELPATASVIR; VOXILAPREVIR - VOSEVI</u>						
N 209195	001	8957046	Mar 21, 2028	U-2039		
		8957046	Mar 21, 2028	U-2040		
		9085573	Mar 21, 2028	DS DP U-2039		
		9085573	Mar 21, 2028	DS DP U-2040		
		9085573*PED	Sep 21, 2028			
		9284342	Sep 13, 2030	DS DP U-2039		
		9284342	Sep 13, 2030	DS DP U-2040		
		9284342*PED	Mar 13, 2031			
		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 LS</u>						
N 209529	001	9918970	May 18, 2031	DP	NP PED	May 26, 2023 Nov 26, 2023
<u>SOLRIAMFETOL HYDROCHLORIDE - SUNOSI</u>						
N 211230	001	10195151	Sep 05, 2037	DP	NCE	Jun 17, 2024
		10351517	Jun 07, 2026	U-2548	ODE-254	Jun 17, 2026
		10512609	Sep 05, 2037	U-2548		
		10912754	Jun 01, 2038	U-3082		
		10940133	Mar 19, 2040	U-3099		
		10959976	Jun 01, 2038	U-3151		
		8440715	Aug 25, 2027	U-2548		
		8877806	Jun 07, 2026	U-2548		
		9604917	Jun 07, 2026	U-2548		
<u>SOLRIAMFETOL HYDROCHLORIDE - SUNOSI</u>						
N 211230	002	10195151	Sep 05, 2037	DP	NCE	Jun 17, 2024
		10351517	Jun 07, 2026	U-2548	ODE-254	Jun 17, 2026
		10512609	Sep 05, 2037	U-2548		
		10912754	Jun 01, 2038	U-3082		
		10959976	Jun 01, 2038	U-3151		
		8440715	Aug 25, 2027	U-2548		
		8877806	Jun 07, 2026	U-2548		
		9604917	Jun 07, 2026	U-2548		
<u>SONIDEGIB PHOSPHATE - ODOMZO</u>						
N 205266	001	8063043	Sep 15, 2029	DS DP		
		8178563	Jul 24, 2029	DS U-1722		
<u>SORAFENIB TOSYLATE - NEXAVAR</u>						
N 021923	001	8618141	Feb 11, 2023	U-1480		
		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 - SOTALOL HYDROCHLORIDE</u>						
N 022306	001	10512620	Aug 18, 2038	U-2769		
		10799138	Apr 05, 2039	U-3125		
<u>SOTALOL HYDROCHLORIDE - SOTYLIZE</u>						
N 205108	001	10206895	Apr 01, 2034	DP U-2096		
		10206895	Apr 01, 2034	DP U-2494		
		11013703	Apr 01, 2034	DP		
		9724297	Aug 31, 2035	DP U-2096		
<u>SOTORASIB - LUMAKRAS</u>						
N 214665	001				NCE ODE-352	May 28, 2026 May 28, 2028
<u>SPINOSAD - NATROBA</u>						
N 022408	001	6063771	Jul 25, 2023	DP U-1670	I-858	Apr 28, 2024
<u>SPIRONOLACTONE - CAROSPIR</u>						
N 209478	001	10493083	Oct 28, 2036	DP		
		10624906	Oct 28, 2036	DP		
		10660907	Oct 28, 2036	DP		

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<u>SPIRONOLACTONE - CAROSPIR</u>						
N 209478	001	10888570	Oct 28, 2036	DP		
		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>SUFENTANIL CITRATE - DSUVIA</u>						
N 209128	001	10245228	Jan 05, 2027	DP U-1351		
		10342762	Jan 05, 2027	DP		
		10507180	Jan 05, 2027	DP U-1351		
		10896751	Mar 16, 2030	DP		
		8202535	Oct 22, 2030	U-1351		
		8226978	Jan 05, 2027	DP U-1351		
		8231900	Jan 05, 2027	DP		
		8252328	Jan 05, 2027	DP		
		8252329	Jan 05, 2027	DP		
		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	RE44733	Jan 27, 2026	DS DP U-1794	M-262 NPP	Jun 09, 2023 Jun 25, 2024
<u>SUGAMMADEX SODIUM - BRIDION</u>						
N 022225	002	RE44733	Jan 27, 2026	DS DP U-1794	M-262 NPP	Jun 09, 2023 Jun 25, 2024
<u>SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSOPHERES - LUMASON</u>						
N 203684	001	10232061	Jul 06, 2038	DP	NPP	Nov 13, 2022
		10335502	Jul 06, 2038	DP		
<u>SUMATRIPTAN - TOSYMRA</u>						
N 210884	001	10603305	Jun 16, 2030	DP U-1719		
		8268791	May 09, 2026	DP		
		8440631	May 09, 2026	DP U-1719		
		9211282	Jul 19, 2031	DP U-1719		
		9283280	May 09, 2026	DP		
		9610280	Jun 16, 2030	DP U-1719		
		9974770	Jun 16, 2030	DP U-1719		

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SUMATRIPTAN - TOSYMRA</u>						
N 210884	001	10603305	Jun 16, 2030	DP U-1719		
		8268791	May 09, 2026	DP		
		8440631	May 09, 2026	DP U-1719		
		9211282	Jul 19, 2031	DP U-1719		
		9283280	May 09, 2026	DP		
		9610280	Jun 16, 2030	DP U-1719		
		9974770	Jun 16, 2030	DP U-1719		
<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 - ZECUITY</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		
		10398859	Dec 19, 2027	DP		
		10478574	Nov 04, 2033	U-2404		
		10722667	Dec 30, 2028	DP		
		7975690	Aug 18, 2025	DP U-1809		
		8047202	Jul 02, 2023	DP		
		8327844	Oct 03, 2023	U-1809		
		8550073	Oct 22, 2029	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		
		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>SUMATRIPTAN SUCCINATE - ZEMBRACE SYMTOUCH</u>						
N 208223	001	10537554	Jan 29, 2036	U-72		
<u>SUVOREXANT - BELSOMRA</u>						
N 204569	001	10098892	May 29, 2033	DP	M-253	Jan 29, 2023
		7951797	Nov 20, 2029	DS DP U-620		
<u>SUVOREXANT - BELSOMRA</u>						
N 204569	002	10098892	May 29, 2033	DP	M-253	Jan 29, 2023
		7951797	Nov 20, 2029	DS DP U-620		

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<u>SUVOREXANT - BELSOMRA</u>						
N 204569	003	10098892	May 29, 2033	DP	M-253	Jan 29, 2023
		7951797	Nov 20, 2029	DS DP U-620		
<u>SUVOREXANT - BELSOMRA</u>						
N 204569	004	10098892	May 29, 2033	DP	M-253	Jan 29, 2023
		7951797	Nov 20, 2029	DS DP U-620		
<u>TACROLIMUS - PROGRAF</u>						
N 050708	001				ODE-294	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TACROLIMUS - PROGRAF</u>						
N 050708	002				ODE-294	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TACROLIMUS - PROGRAF</u>						
N 050708	003				ODE-294	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TACROLIMUS - PROGRAF</u>						
N 050709	001				ODE-294	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TACROLIMUS - ASTAGRAF XL</u>						
N 204096	001				ODE*	May 24, 2025
<u>TACROLIMUS - ASTAGRAF XL</u>						
N 204096	002				ODE*	May 24, 2025
<u>TACROLIMUS - ASTAGRAF XL</u>						
N 204096	003				ODE*	May 24, 2025
<u>TACROLIMUS - ENVARUSUS XR</u>						
N 206406	001	10166190	May 30, 2028	DP	ODE-94	Jul 10, 2022
		10548880	Aug 30, 2024	U-2677		
		10548880	Aug 30, 2024	U-2678		
		10864199	May 30, 2028	U-2677		
		10864199	May 30, 2028	U-2678		
		11110081	May 30, 2028	U-2678		
		11123331	May 30, 2028	U-2677		
		7994214	Aug 30, 2024	DP		
		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	Aug 30, 2028	U-1752		
		8664239	Aug 30, 2028	U-2677		
		8664239	Aug 30, 2028	U-2678		
		8685998	Aug 30, 2028	DP U-1752		
		8685998	Aug 30, 2028	DP U-2677		
		8685998	Aug 30, 2028	DP U-2678		
		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	10166190	May 30, 2028	DP	ODE-94	Jul 10, 2022
		10548880	Aug 30, 2024	U-2677		
		10548880	Aug 30, 2024	U-2678		
		10864199	May 30, 2028	U-2677		
		10864199	May 30, 2028	U-2678		
		11110081	May 30, 2028	U-2678		
		11123331	May 30, 2028	U-2677		
		7994214	Aug 30, 2024	DP		
		8486993	Aug 30, 2024	DP U-1752		

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<u>TACROLIMUS - ENVARSUS XR</u>						
N 206406	002	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	Aug 30, 2028	U-1752		
		8664239	Aug 30, 2028	U-2677		
		8664239	Aug 30, 2028	U-2678		
		8685998	Aug 30, 2028	DP U-1752		
		8685998	Aug 30, 2028	DP U-2677		
		8685998	Aug 30, 2028	DP U-2678		
		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 - ENVARSUS XR</u>						
N 206406	003	10166190	May 30, 2028	DP	ODE-94	Jul 10, 2022
		10548880	Aug 30, 2024	U-2677		
		10548880	Aug 30, 2024	U-2678		
		10864199	May 30, 2028	U-2677		
		10864199	May 30, 2028	U-2678		
		11110081	May 30, 2028	U-2678		
		11123331	May 30, 2028	U-2677		
		7994214	Aug 30, 2024	DP		
		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	Aug 30, 2028	U-1752		
		8664239	Aug 30, 2028	U-2677		
		8664239	Aug 30, 2028	U-2678		
		8685998	Aug 30, 2028	DP U-1752		
		8685998	Aug 30, 2028	DP U-2677		
		8685998	Aug 30, 2028	DP U-2678		
		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 - PROGRAF</u>						
N 210115	001				ODE-269	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TACROLIMUS - PROGRAF</u>						
N 210115	002				ODE-269	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TAFAMIDIS - VYNDAMAX</u>						
N 212161	001	7214695	Apr 27, 2024	DS DP	NCE	May 03, 2024
		7214696	Dec 19, 2023	U-2524	ODE-237	May 03, 2026
		9770441	Aug 31, 2035	DS DP	U-2524	
<u>TAFAMIDIS MEGLUMINE - VYNDAQEL</u>						
N 211996	001	7214695	Apr 27, 2024	DS DP	NCE	May 03, 2024
		7214696	Dec 19, 2023	U-2524	ODE-237	May 03, 2026
		8168663	Dec 19, 2023	DS DP		
		8653119	Jan 28, 2024	U-2524		
<u>TAFENOQUINE SUCCINATE - ARAKODA</u>						
N 210607	001	10342791	Dec 02, 2035	U-2582	NCE	Jul 20, 2023
		10888558	Dec 02, 2035	U-2582		

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<u>TAFENOQUINE SUCCINATE - KRINTAFEL</u>						
N 210795	001				NCE ODE-201	Jul 20, 2023 Jul 20, 2025
<u>TAFLUPROST - ZIOPTAN</u>						
N 202514	001	10864159	May 28, 2029	DP U-778		
		5886035	Dec 18, 2022	DS DP U-778		
		9999593	May 28, 2029	DP		
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651	001	10189837	Oct 20, 2031	DS DP	NCE	Oct 16, 2023
		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	10189837	Oct 20, 2031	DS DP	NCE	Oct 16, 2023
		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	003	10189837	Oct 20, 2031	DS DP	NCE	Oct 16, 2023
		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	004	10189837	Oct 20, 2031	DS DP	NCE	Oct 16, 2023
		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
<u>TALC - STERITALC</u>						
N 205555	003				ODE-143 ODE-191	May 01, 2024 May 01, 2024
<u>TAPENTADOL HYDROCHLORIDE - NUCYNТА</u>						
N 022304	001	7994364	Jun 27, 2025	DS DP U-931		
		RE39593	Aug 05, 2022	DS DP U-931		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNТА</u>						
N 022304	002	7994364	Jun 27, 2025	DS DP U-931		
		RE39593	Aug 05, 2022	DS DP U-931		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNТА</u>						
N 022304	003	7994364	Jun 27, 2025	DS DP U-931		
		RE39593	Aug 05, 2022	DS DP U-931		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNТА ER</u>						
N 200533	001	11007156	Oct 22, 2022	DP U-1178		
		11007156	Oct 22, 2022	DP U-1276		
		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		

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<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	001	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	11007156	Oct 22, 2022		DP U-1178	
		11007156	Oct 22, 2022		DP U-1276	
		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	11007156	Oct 22, 2022		DP U-1178	
		11007156	Oct 22, 2022		DP U-1276	
		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	11007156	Oct 22, 2022		DP U-1178	
		11007156	Oct 22, 2022		DP U-1276	
		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	005	11007156	Oct 22, 2022		DP U-1178	
		11007156	Oct 22, 2022		DP U-1276	
		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		I-850 Dec 01, 2023
		10149829	Jan 25, 2033		U-2477	ODE-330 Dec 01, 2027
		10149829	Jan 25, 2033		U-3006	
		10179119	Aug 29, 2035		U-3003	
		10376487	Jul 27, 2035		U-2615	

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<u>TASIMELTEON - HETLIOZ</u>						
N 205677	001	10376487	Jul 27, 2035	U-3007		
		10449176	Jan 25, 2033	U-2149		
		10610510	Jan 25, 2033	U-2805		
		10610510	Jan 25, 2033	U-3009		
		10610511	Oct 10, 2034	U-2615		
		10610511	Oct 10, 2034	U-3007		
		10829465	Feb 12, 2035	DS DP		
		10945988	Jan 25, 2033	U-2149		
		10980770	Jan 25, 2033	U-3106		
		10980770	Jan 25, 2033	U-3107		
		11141400	Oct 10, 2034	U-2615		
		11141400	Oct 10, 2034	U-3007		
		5856529	Dec 09, 2022	DS DP	U-2149	
		5856529	Dec 09, 2022	DS DP	U-3003	
		9060995	Jan 25, 2033	U-1710		
		9539234	Jan 25, 2033	U-1934		
		9539234	Jan 25, 2033	U-3004		
		9549913	Jan 25, 2033	U-1486		
		9730910	May 17, 2034	U-2085		
		9730910	May 17, 2034	U-3005		
		9855241	Jan 25, 2033	U-2149		
		RE46604	Jan 25, 2033	U-2147		
<u>TASIMELTEON - HETLIOZ LO</u>						
N 214517	001	10071977	Feb 12, 2035	DS DP	NP	Dec 01, 2023
		10149829	Jan 25, 2033	U-3006	ODE-329	Dec 01, 2027
		10179119	Aug 29, 2035	U-3003		
		10376487	Jul 27, 2035	U-3007		
		10610510	Jan 25, 2033	U-3009		
		10610511	Oct 10, 2034	U-3007		
		10829465	Feb 12, 2035	DS DP		
		10980770	Jan 25, 2033	U-3106		
		11141400	Oct 10, 2034	U-3007		
		11202770	Dec 11, 2040	DP		
		5856529	Dec 09, 2022	DS DP	U-3003	
		9539234	Jan 25, 2033	U-3004		
		9730910	May 17, 2034	U-3005		
<u>TAZAROTENE - FABIOR</u>						
N 202428	001	10568859	Feb 24, 2030	DP	U-2760	
		10688071	Feb 24, 2030	DP	U-2760	
		8808716	Feb 24, 2030	DP		
<u>TAZAROTENE - ARAZLO</u>						
N 211882	001				NP	Dec 18, 2022
<u>TAZEMETOSTAT HYDROBROMIDE - TAZVERIK</u>						
N 211723	001	10155002	Apr 13, 2032	U-2736	I-835	Jun 18, 2023
		10245269	Apr 11, 2033	U-2737	I-836	Jun 18, 2023
		10245269	Apr 11, 2033	U-2851	NCE	Jan 23, 2025
		10245269	Apr 11, 2033	U-2854	ODE-299	Jan 23, 2027
		10369155	Oct 16, 2035	U-2736	ODE-314	Jun 18, 2027
		10369155	Oct 16, 2035	U-2852		
		10369155	Oct 16, 2035	U-2853		
		10420775	Apr 13, 2032	U-2736		
		10420775	Apr 13, 2032	U-2852		
		10420775	Apr 13, 2032	U-2853		
		10786511	Dec 19, 2035	DP		
		10821113	Apr 11, 2033	DS DP		
		11052093	Apr 13, 2032	DS DP	U-2736	
		11052093	Apr 13, 2032	DS DP	U-3179	
		8410088	Apr 13, 2032	DS DP		
		8691507	Sep 12, 2031	U-2852		
		8765732	Apr 13, 2032	U-2852		
		8765732	Apr 13, 2032	U-2853		
		8895245	Sep 12, 2031	U-2852		
		9090562	Apr 13, 2032	DS DP		
		9175331	Sep 12, 2031	U-2852		
		9333217	Sep 12, 2031	U-2852		
		9334527	Sep 12, 2031	U-2852		

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<u>TAZEMETOSTAT HYDROBROMIDE - TAZVERIK</u>						
N 211723	001	9394283	Apr 11, 2033	DS DP	U-2852	
		9394283	Apr 11, 2033	DS DP	U-2853	
		9522152	Apr 13, 2032		U-2738	
		9549931	Apr 13, 2032		U-2736	
		9549931	Apr 13, 2032		U-2852	
		9549931	Apr 13, 2032		U-2853	
		9688665	Aug 22, 2034		U-2736	
		9855275	Apr 13, 2032		U-2736	
		9872862	Apr 11, 2033		U-2738	
		9889138	Oct 16, 2035		U-2736	
		9889138	Oct 16, 2035		U-2852	
		9889138	Oct 16, 2035		U-2853	
		9949999	Sep 12, 2031		U-2851	
<u>TECHNETIUM TC-99M TETROFOSMIN KIT - MYOVUE 30ML</u>						
N 020372	002	9549999	Mar 10, 2030	DP		
<u>TECHNETIUM TC-99M TILMANOCEPT - LYMPHOSEEK KIT</u>						
N 202207	001	6409990	May 12, 2025	DS		NPP May 19, 2024
		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	10065947	Feb 03, 2030		DP	NCE Jun 20, 2019
		10442829	Feb 03, 2030		DS	NPP Jun 19, 2023
		7816379	Jun 20, 2028	DS DP	U-2507	GAIN Jun 20, 2024
		8420676	Feb 23, 2028	DS DP	U-282	
		8426389	Dec 31, 2030	DS DP	U-282	
		9624250	Feb 03, 2030	DS DP	U-2507	
		9988406	Feb 03, 2030		DP	
<u>TEDIZOLID PHOSPHATE - SIVEXTRO</u>						
N 205436	001	10065947	Feb 03, 2030		DP	NCE Jun 20, 2019
		10442829	Feb 03, 2030		DS	NPP Jun 19, 2023
		7816379	Jun 20, 2028	DS DP	U-2507	GAIN Jun 20, 2024
		8420676	Feb 23, 2028	DS DP	U-282	
		8426389	Dec 31, 2030	DS DP	U-282	
		9624250	Feb 03, 2030	DS DP	U-2507	
		9988406	Feb 03, 2030		DP	
<u>TEDUGLUTIDE RECOMBINANT - GATTEX KIT</u>						
N 203441	001	7056886	Sep 18, 2022	DP	U-1320	ODE-240 May 16, 2026
		7056886*PED	Mar 18, 2023			
		7847061	Nov 01, 2025		U-1320	
		7847061*PED	May 01, 2026			
		9060992	Nov 01, 2025		U-1320	
		9060992*PED	May 01, 2026			
		9539310	Nov 01, 2025		U-1320	
		9539310*PED	May 01, 2026			
		9545434	Nov 01, 2025		U-1320	
		9545434*PED	May 01, 2026			
		9545435	Nov 01, 2025		U-1320	
		9545435*PED	May 01, 2026			
		9555079	Nov 01, 2025		U-1320	
		9555079*PED	May 01, 2026			
		9572867	Nov 01, 2025		U-1320	
		9572867*PED	May 01, 2026			
		9592273	Nov 01, 2025		U-1320	
		9592273*PED	May 01, 2026			
		9592274	Nov 01, 2025		U-1320	
		9592274*PED	May 01, 2026			
		9968655	Nov 01, 2025		U-2308	
		9968655*PED	May 01, 2026			

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<u>TEDEGLUTIDE RECOMBINANT - GATTEX KIT</u>						
N 203441	001	9968656	Nov 01, 2025	U-2308		
		9968656*PED	May 01, 2026			
		9968658	Nov 01, 2025	U-1320		
		9968658*PED	May 01, 2026			
		9974835	Nov 01, 2025	U-1320		
		9974835*PED	May 01, 2026			
		9974837	Nov 01, 2025	U-1320		
		9974837*PED	May 01, 2026			
		9981014	Nov 01, 2025	U-1320		
		9981014*PED	May 01, 2026			
		9981016	Nov 01, 2025	U-1320		
		9981016*PED	May 01, 2026			
		9987334	Nov 01, 2025	U-1320		
		9987334*PED	May 01, 2026			
		9987335	Nov 01, 2025	U-1320		
		9987335*PED	May 01, 2026			
		9993528	Nov 01, 2025	U-1320		
		9993528*PED	May 01, 2026			
<u>TELAPREVIR - INCIVEK</u>						
N 201917	001	7820671	Feb 25, 2025	DS DP		
		8431615	May 30, 2028	U-1398		
<u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u>						
N 022110	001	6635618	Sep 11, 2023	DS DP	U-728	
		6858584	Aug 24, 2022	DP		
		7531623	Jan 01, 2027	DS		
<u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u>						
N 022110	002	6635618	Sep 11, 2023	DS DP	U-728	
		6858584	Aug 24, 2022	DP		
		7531623	Jan 01, 2027	DS		
<u>TELBIVUDINE - TYZEKA</u>						
N 022011	001	7589079	Sep 11, 2023	DS DP	U-999	
		7858594	Sep 11, 2023	DS DP	U-999	
<u>TELBIVUDINE - TYZEKA</u>						
N 022154	001	7858594	Sep 11, 2023	DS DP	U-999	
<u>TELMISARTAN - MICARDIS</u>						
N 020850	002	8003679	Oct 06, 2022	U-1176		
<u>TELOTRISTAT ETIPRATE - XERMELO</u>						
N 208794	001	7553840	Dec 11, 2027	DS	NCE	Feb 28, 2022
		7709493	Dec 11, 2027	DS	U-1979	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>TEMSTIROLIMUS - TORISEL</u>						
N 022088	001	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			
<u>TENAPANOR HYDROCHLORIDE - IBSRELA</u>						
N 211801	001	8541448	Dec 30, 2029	DS DP	NCE	Sep 12, 2024
		8969377	Dec 30, 2029	DS DP		
		9006281	May 02, 2030	U-2626		

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<u>TENAPANOR HYDROCHLORIDE - IBSRELA</u>						
N 211801	001	9408840	Dec 30, 2029	U-2626		
<u>TENOFOVIR ALAFENAMIDE FUMARATE - VEMLIDY</u>						
N 208464	001	7390791	Apr 17, 2025	DS DP	M-255	Feb 04, 2023
		7390791*PED	Oct 17, 2025		M-266	Aug 22, 2023
		7803788	Feb 02, 2022	U-999		
		8754065	Aug 15, 2032	DS DP U-999		
		8754065*PED	Feb 15, 2033			
		9296769	Aug 15, 2032	DS DP U-999		
		9296769*PED	Feb 15, 2033			
<u>TEPOTINIB HYDROCHLORIDE - TEPMETKO</u>						
N 214096	001	8329692	Oct 30, 2029	DS DP	NCE	Feb 03, 2026
		8580781	Mar 19, 2030	DS DP	ODE-325	Feb 03, 2028
		8658643	Jul 04, 2028	U-3077		
		8921357	May 30, 2028	DS DP		
		8927540	Jul 21, 2028	U-3078		
		9062029	Jul 04, 2028	DP		
		9284300	Apr 29, 2028	DP		
		9403799	Jul 04, 2028	U-3077		
<u>TERIFLUNOMIDE - AUBAGIO</u>						
N 202992	001	6794410	Sep 12, 2026	U-1285	M-61	Apr 30, 2024
		6794410*PED	Mar 12, 2027		PED	Oct 30, 2024
		8802735	Sep 14, 2030	DP		
		8802735*PED	Mar 14, 2031			
		9186346	Feb 04, 2034	U-1786		
		9186346*PED	Aug 04, 2034			
<u>TERIFLUNOMIDE - AUBAGIO</u>						
N 202992	002	6794410	Sep 12, 2026	U-1285	M-61	Apr 30, 2024
		6794410*PED	Mar 12, 2027		PED	Oct 30, 2024
		8802735	Sep 14, 2030	DP		
		8802735*PED	Mar 14, 2031			
		9186346	Feb 04, 2034	U-1786		
		9186346*PED	Aug 04, 2034			
<u>TERIPARATIDE - FORTEO</u>						
N 021318	001	7517334	Mar 25, 2025	DP		
<u>TERIPARATIDE - FORTEO</u>						
N 021318	002	7517334	Mar 25, 2025	DP		
<u>TESTOSTERONE - TESTIM</u>						
N 021454	001	7320968	Jan 18, 2025	U-843		
		7608605	Apr 21, 2023	U-1009		
		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 - ANDROGEL</u>						
N 022309	001	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		
<u>TESTOSTERONE - ANDROGEL</u>						
N 022309	002	8466136	Oct 12, 2026	DP		
		8466137	Oct 12, 2026	U-1103		
		8466138	Oct 12, 2026	U-1103		
		8486925	Oct 12, 2026	DP		

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<u>TESTOSTERONE - ANDROGEL</u>						
N 022309	002	8729057	Oct 12, 2026	DP		
		8741881	Oct 12, 2026		U-1103	
		8754070	Oct 12, 2026	DP		
		8759329	Oct 12, 2026	DP		
<u>TESTOSTERONE - ANDROGEL</u>						
N 022309	003	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		
<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	
		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	11090312	Mar 17, 2034		U-1616	
		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	10238662	Feb 19, 2035	DP	U-2418	
		10279131	Jul 31, 2031	DP		
		10357609	Aug 21, 2031	DP		
		10478560	Jan 24, 2026	DP		
		10646495	Aug 30, 2038	DP		
		10821072	Jun 04, 2033	DP	U-2418	
		10881798	Feb 11, 2034	DP		
		10905827	Aug 21, 2031	DP		
		10912782	Feb 19, 2035	DP	U-2418	
		11160751	Oct 07, 2034	DP	U-2418	
		11191908	Oct 18, 2035	DP		
		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		
		9744302	Nov 19, 2035	DP		
		9950125	Sep 04, 2036	DP	U-2418	
<u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u>						
N 209863	002	10238662	Feb 19, 2035	DP	U-2418	
		10279131	Jul 31, 2031	DP		
		10357609	Aug 21, 2031	DP		
		10478560	Jan 24, 2026	DP		
		10646495	Aug 30, 2038	DP		
		10821072	Jun 04, 2033	DP	U-2418	
		10881798	Feb 11, 2034	DP		
		10905827	Aug 21, 2031	DP		

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<u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u>						
N 209863 002	10912782	Feb 19, 2035	DP U-2418			
	11160751	Oct 07, 2034	DP U-2418			
	11191908	Oct 18, 2035	DP			
	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			
	9744302	Nov 19, 2035	DP			
	9950125	Sep 04, 2036	DP U-2418			
<u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u>						
N 209863 003	10238662	Feb 19, 2035	DP U-2418			
	10279131	Jul 31, 2031	DP			
	10357609	Aug 21, 2031	DP			
	10478560	Jan 24, 2026	DP			
	10646495	Aug 30, 2038	DP			
	10821072	Jun 04, 2033	DP U-2418			
	10881798	Feb 11, 2034	DP			
	10905827	Aug 21, 2031	DP			
	10912782	Feb 19, 2035	DP U-2418			
	11160751	Oct 07, 2034	DP U-2418			
	11191908	Oct 18, 2035	DP			
	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			
	9744302	Nov 19, 2035	DP			
	9950125	Sep 04, 2036	DP U-2418			
<u>TESTOSTERONE UNDECANOATE - AVEED</u>						
N 022219 001	7718640	Mar 14, 2027	DP			
	8338395	May 08, 2027	U-1500			
<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089 001	10543219	Apr 12, 2030	U-2506		NP	Mar 27, 2022
	10617696	Apr 12, 2030	DS DP			
	11179402	Apr 14, 2026	DS DP			
	11179403	Apr 12, 2030	U-2506			
	8241664	Mar 29, 2029	DP U-2506			
	8492369	Dec 20, 2030	DP U-2506			
	8778916	Apr 12, 2030	DP			
<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089 002	10543219	Apr 12, 2030	U-2506		NP	Mar 27, 2022
	10617696	Apr 12, 2030	DS DP			
	11179402	Apr 14, 2026	DS DP			
	11179403	Apr 12, 2030	U-2506			
	8241664	Mar 29, 2029	DP U-2506			
	8492369	Dec 20, 2030	DP U-2506			
	8778916	Apr 12, 2030	DP			
<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089 003	10543219	Apr 12, 2030	U-2506		NP	Mar 27, 2022
	10617696	Apr 12, 2030	DS DP			
	11179402	Apr 14, 2026	DS DP			
	11179403	Apr 12, 2030	U-2506			
	8241664	Mar 29, 2029	DP U-2506			
	8492369	Dec 20, 2030	DP U-2506			
	8778916	Apr 12, 2030	DP			
<u>THALIDOMIDE - THALOMID</u>						
N 020785 001	7230012	Dec 09, 2023	DP			
<u>THALIDOMIDE - THALOMID</u>						
N 020785 002	7230012	Dec 09, 2023	DP			

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<u>THALIDOMIDE - THALOMID</u>						
N 020785	003 7230012	Dec 09, 2023	DP			
<u>THIOTEPA - TEPADINA</u>						
N 208264	001				ODE-129	Jan 26, 2024
<u>THIOTEPA - TEPADINA</u>						
N 208264	002				ODE-129	Jan 26, 2024
<u>TICAGRELOR - BRILINTA</u>						
N 022433	001 10300065	Jan 27, 2036		U-2541	I-848	Nov 05, 2023
	10300065	Jan 27, 2036		U-2542	I-851	May 28, 2023
	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		
	RE46276	Oct 30, 2024	DS DP	U-2838		
	RE46276	Oct 30, 2024	DS DP	U-2839		
	RE46276	Oct 30, 2024	DS DP	U-2988		
<u>TICAGRELOR - BRILINTA</u>						
N 022433	002 10300065	Jan 27, 2036		U-2541	I-848	Nov 05, 2023
	10300065	Jan 27, 2036		U-2542	I-851	May 28, 2023
	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		
	RE46276	Oct 30, 2024	DS DP	U-2838		
	RE46276	Oct 30, 2024	DS DP	U-2839		
	RE46276	Oct 30, 2024	DS DP	U-2988		
<u>TIGECYCLINE - TYGACIL</u>						
N 021821	001 10588975	Mar 13, 2026	DP			
	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>TIOPRONIN - THIOLA</u>						
N 019569	001				ODE-267	Jun 28, 2026
<u>TIOTROPIUM BROMIDE - SPIRIVA</u>						
N 021395	001 6777423*PED	Mar 24, 2022				
	6908928*PED	Mar 24, 2022				
	7070800	Jan 22, 2022	DP	U-566		
	7070800*PED	Jul 22, 2022				
	7309707*PED	Mar 24, 2022				
	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				
	9010323	Apr 19, 2030	DP			
	RE38912*PED	Apr 11, 2022				
<u>TIOTROPIUM BROMIDE - SPIRIVA RESPIMAT</u>						
N 021936	001 7284474	Aug 26, 2024	DP			
	7284474*PED	Feb 26, 2025				
	7396341	Oct 10, 2026	DP			
	7396341*PED	Apr 10, 2027				
	7837235	Mar 13, 2028	DP			
	7837235*PED	Sep 13, 2028				
	7896264	May 26, 2025	DP			
	7896264*PED	Nov 26, 2025				
	7988001*PED	Feb 04, 2022				

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<u>TIOTROPIUM BROMIDE - SPIRIVA RESPIMAT</u>						
N 021936	001	8733341	Oct 16, 2030	DP		
		8733341*PED	Apr 16, 2031			
		9027967	Mar 31, 2027	DP		
		9027967*PED	Oct 01, 2027			
<u>TIOTROPIUM BROMIDE - SPIRIVA RESPIMAT</u>						
N 021936	002	7284474	Aug 26, 2024	DP		
		7284474*PED	Feb 26, 2025			
		7396341	Oct 10, 2026	DP		
		7396341*PED	Apr 10, 2027			
		7837235	Mar 13, 2028	DP		
		7837235*PED	Sep 13, 2028			
		7896264	May 26, 2025	DP		
		7896264*PED	Nov 26, 2025			
		7988001*PED	Feb 04, 2022			
		8733341	Oct 16, 2030	DP		
		8733341*PED	Apr 16, 2031			
		9027967	Mar 31, 2027	DP		
		9027967*PED	Oct 01, 2027			
<u>TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE - LONSURE</u>						
N 207981	001	10456399	Feb 03, 2037		U-2642	
		10457666	Jun 17, 2034	DS DP		I-794 Feb 22, 2022
		10960004	Feb 03, 2037		U-2642	ODE-229 Feb 22, 2026
		9527833	Jun 17, 2034	DS DP		
		RE46284	Sep 22, 2029		U-1751	
		RE46284	Sep 22, 2029		U-2503	
<u>TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE - LONSURE</u>						
N 207981	002	10456399	Feb 03, 2037		U-2642	
		10457666	Jun 17, 2034	DS DP		I-794 Feb 22, 2022
		10960004	Feb 03, 2037		U-2642	ODE-229 Feb 22, 2026
		9527833	Jun 17, 2034	DS DP		
		RE46284	Sep 22, 2029		U-1751	
		RE46284	Sep 22, 2029		U-2503	
<u>TIRBANIBULIN - KLISYRI</u>						
N 213189	001	10323001	Dec 28, 2027	DP		
		10617693	Mar 12, 2038		U-3015	
		10669236	Sep 07, 2038	DS DP		
		7300931	Feb 06, 2026	DS DP		
		7851470	Feb 02, 2029	DS DP	U-3015	
		8236799	Dec 28, 2025	DS DP		
		8980890	Dec 28, 2025	DS DP		
<u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N 020912	001	6770660	May 01, 2023		U-1444	
<u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N 020912	002	6770660	May 01, 2023		U-1444	
<u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N 020913	001	6770660	May 01, 2023		U-1444	
<u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N 020913	002	6770660	May 01, 2023		U-1444	
<u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N 020913	003	6770660	May 01, 2023		U-1444	
<u>TIVOZANIB HYDROCHLORIDE - FOTIVDA</u>						
N 212904	001	6821987	Apr 26, 2022	DS DP	U-3100	
		7166722	Nov 16, 2023	DS		NCE Mar 10, 2026
<u>TIVOZANIB HYDROCHLORIDE - FOTIVDA</u>						
N 212904	002	6821987	Apr 26, 2022	DS DP	U-3100	
		7166722	Nov 16, 2023	DS		NCE Mar 10, 2026

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<u>TOBRAMYCIN - TOBI PODHALER</u>						
N 201688	001	10207066	Nov 04, 2030	DP U-909		
		7368102	Dec 19, 2022	DP U-909		
		7516741	Jan 11, 2024	DP		
		7559325	Oct 27, 2025	DP		
		8069851	Sep 24, 2024	DP		
		8664187	Jun 20, 2025	U-909		
		8715623	Dec 19, 2022	DP U-909		
		8869794	Sep 12, 2028	DP U-909		
		9421166	Dec 19, 2022	DP U-909		
		RE47526	Apr 09, 2024	DP		
<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	6965027	Mar 25, 2023	DS	NPP	Sep 25, 2023
		RE41783	Dec 08, 2025	DS		
<u>TOFACITINIB CITRATE - XELJANZ</u>						
N 203214	002	6965027	Mar 25, 2023	DS		
		RE41783	Dec 08, 2025	DS		
<u>TOFACITINIB CITRATE - XELJANZ XR</u>						
N 208246	001	6965027	Mar 25, 2023	DS		
		9937181	Mar 14, 2034	DP		
		RE41783	Dec 08, 2025	DS		
<u>TOFACITINIB CITRATE - XELJANZ XR</u>						
N 208246	002	10639309	Mar 14, 2034	DP		
		6965027	Mar 25, 2023	DS		
		RE41783	Dec 08, 2025	DS		
<u>TOFACITINIB CITRATE - XELJANZ</u>						
N 213082	001	RE41783	Dec 08, 2025	DS	NP	Sep 25, 2023
<u>TOLVAPTAN - SAMSCA</u>						
N 022275	001	10905694	Apr 07, 2030	DP		
		8501730	Sep 01, 2026	DS		
<u>TOLVAPTAN - SAMSCA</u>						
N 022275	002	10905694	Apr 07, 2030	DP		
		8501730	Sep 01, 2026	DS		
<u>TOLVAPTAN - SAMSCA</u>						
N 022275	003	10905694	Apr 07, 2030	DP		
		8501730	Sep 01, 2026	DS		
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	001	10905694	Apr 07, 2030	DP	ODE-178	Apr 23, 2025
		8501730	Sep 01, 2026	DS		
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	002	10905694	Apr 07, 2030	DP	ODE-178	Apr 23, 2025
		8501730	Sep 01, 2026	DS		
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	003	10905694	Apr 07, 2030	DP	ODE-178	Apr 23, 2025
		8501730	Sep 01, 2026	DS		
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	004	10905694	Apr 07, 2030	DP	ODE-178	Apr 23, 2025
		8501730	Sep 01, 2026	DS		
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	005	10905694	Apr 07, 2030	DP	ODE-178	Apr 23, 2025
		8501730	Sep 01, 2026	DS		

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<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635	001	10314790	Nov 16, 2027	DP U-1675		
		10314790	Nov 16, 2027	DP U-1992		
		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	10314790	Nov 16, 2027	DP U-1675		
		10314790	Nov 16, 2027	DP U-1992		
		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	003	10314790	Nov 16, 2027	DP U-1675		
		10314790	Nov 16, 2027	DP U-1992		
		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	10314790	Nov 16, 2027	DP U-1675		
		10314790	Nov 16, 2027	DP U-1992		
		8298576	Apr 04, 2028	DP U-106		
		8298576	Apr 04, 2028	DP U-1992		

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<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635	004	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	10363224	Mar 19, 2033	U-766		
		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	10363224	Mar 19, 2033	U-766		
		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	10363224	Mar 19, 2033	U-766		
		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	004	10363224	Mar 19, 2033	U-766		
		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	10363224	Mar 19, 2033	U-766		
		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>TORSEMIDE - SOAANZ</u>						
N 213218	001	10154963	Oct 06, 2033	DP		
<u>TORSEMIDE - SOAANZ</u>						
N 213218	002	10154963	Oct 06, 2033	DP		
<u>TRABECTEDIN - YONDELIS</u>						
N 207953	001	8895557	Jan 07, 2028	DP	ODE-100	Oct 23, 2022
		8895557*PED	Jul 07, 2028		PED	Apr 23, 2023

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<u>TRAMADOL HYDROCHLORIDE - RYZOLT</u>						
N 021745	001 7988998	Oct 27, 2023	DP			
<u>TRAMADOL HYDROCHLORIDE - RYZOLT</u>						
N 021745	002 7988998	Oct 27, 2023	DP			
<u>TRAMADOL HYDROCHLORIDE - RYZOLT</u>						
N 021745	003 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>TRAMADOL HYDROCHLORIDE - QDOLO</u>						
N 214044	001 11103452	Sep 01, 2040	DP U-3197			
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114	001 10869869	Aug 30, 2033	U-3184		M-246	Oct 06, 2022
	7378423	May 29, 2027	DS DP		ODE-148	Jun 22, 2024
	8580304	Jan 28, 2032	DP		ODE-182	Apr 30, 2025
	8703781	Oct 15, 2030	DS DP U-1712		ODE-183	May 04, 2025
	8703781	Oct 15, 2030	DS DP U-2020			
	8703781	Oct 15, 2030	DS DP U-2037			
	8703781	Oct 15, 2030	DS DP U-2302			
	8703781	Oct 15, 2030	DS DP U-2305			
	8835443	Jun 10, 2025	U-1581			
	8835443	Jun 10, 2025	U-1582			
	8835443	Jun 10, 2025	U-2020			
	8835443	Jun 10, 2025	U-2037			
	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			
	9399021	Jan 28, 2032	DP			
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114	002 10869869	Aug 30, 2033	U-3184		M-246	Oct 06, 2022
	7378423	May 29, 2027	DS DP		ODE-148	Jun 22, 2024
	8580304	Jan 28, 2032	DP		ODE-182	Apr 30, 2025
	8703781	Oct 15, 2030	DS DP U-1712		ODE-183	May 04, 2025
	8703781	Oct 15, 2030	DS DP U-2020			
	8703781	Oct 15, 2030	DS DP U-2037			
	8703781	Oct 15, 2030	DS DP U-2302			
	8703781	Oct 15, 2030	DS DP U-2305			
	8835443	Jun 10, 2025	U-1581			
	8835443	Jun 10, 2025	U-1582			
	8835443	Jun 10, 2025	U-2020			
	8835443	Jun 10, 2025	U-2037			
	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			
	9399021	Jan 28, 2032	DP			
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114	003 10869869	Aug 30, 2033	U-3184		M-246	Oct 06, 2022
	7378423	May 29, 2027	DS DP		ODE-148	Jun 22, 2024
	8580304	Jan 28, 2032	DP		ODE-182	Apr 30, 2025
	8703781	Oct 15, 2030	DS DP U-1712		ODE-183	May 04, 2025
	8703781	Oct 15, 2030	DS DP U-2020			
	8703781	Oct 15, 2030	DS DP U-2037			
	8703781	Oct 15, 2030	DS DP U-2302			
	8703781	Oct 15, 2030	DS DP U-2305			
	8835443	Jun 10, 2025	U-1581			

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<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114	003	8835443	Jun 10, 2025			U-1582
		8835443	Jun 10, 2025			U-2020
		8835443	Jun 10, 2025			U-2037
		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		
		9399021	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	7829120	Mar 27, 2027	DP	U-796	
		8133893	Mar 13, 2029	DS	DP	
<u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u>						
N 022411	002	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		
		10695308	Dec 16, 2024		U-2845	
		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		
		10695308	Dec 16, 2024		U-2845	
		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		

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<u>TREPROSTINIL - REMODULIN</u>						
N 021272	002	9604901	Dec 15, 2028	DS		
		9713599	Dec 16, 2024		U-2036	
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	003	10076505	Dec 16, 2024	DP		
		10695308	Dec 16, 2024		U-2845	
		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		
		10695308	Dec 16, 2024		U-2845	
		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	005	10076505	Dec 16, 2024	DP		
		10695308	Dec 16, 2024		U-2845	
		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	10376525	May 14, 2027		U-1849	
		10716793	May 14, 2027		U-1849	
		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	
<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	

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<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 001	7417070	Jul 30, 2026	DS		I-820	Oct 18, 2022
	7544713	Jul 14, 2024		U-1475	ODE-272	Oct 18, 2026
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8747897	Aug 11, 2031	DP	U-2724		
	8747897	Aug 11, 2031	DP	U-2725		
	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		I-820	Oct 18, 2022
	7544713	Jul 14, 2024		U-1475	ODE-272	Oct 18, 2026
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8497393	Dec 15, 2028	DS		Y	
	8747897	Aug 11, 2031	DP	U-2724		
	8747897	Aug 11, 2031	DP	U-2725		
	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		I-820	Oct 18, 2022
	7544713	Jul 14, 2024		U-1475	ODE-272	Oct 18, 2026
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8497393	Dec 15, 2028	DS		Y	
	8747897	Aug 11, 2031	DP	U-2724		
	8747897	Aug 11, 2031	DP	U-2725		
	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		I-820	Oct 18, 2022
	7544713	Jul 14, 2024		U-1475	ODE-272	Oct 18, 2026
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8497393	Dec 15, 2028	DS		Y	
	8747897	Aug 11, 2031	DP	U-2724		
	8747897	Aug 11, 2031	DP	U-2725		
	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		I-820	Oct 18, 2022
	7544713	Jul 14, 2024		U-1475	ODE-272	Oct 18, 2026
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8747897	Aug 11, 2031	DP	U-2724		
	8747897	Aug 11, 2031	DP	U-2725		

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<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496	005 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 - ALTRENO</u>						
N 209353	001 10653656	Aug 22, 2038	DP		U-2368	
<u>TRIAMCINOLONE ACETONIDE - TRIESENC</u>						
N 022048	001 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			
	9555048	Aug 04, 2031			U-2151	
<u>TRIAMCINOLONE ACETONIDE - XIPERE</u>						
N 211950	001 8636713	May 02, 2027			U-3234	
	9636332	Nov 08, 2033			U-3234	
	9937075	May 02, 2034	DP			
<u>TRICLABENDAZOLE - EGATEN</u>						
N 208711	001					
					NCE	Feb 13, 2024
					ODE-228	Feb 13, 2026
<u>TRIFAROTENE - AKLIEF</u>						
N 211527	001 7807708	Oct 01, 2026	DS DP			
	8227507	Dec 21, 2025			U-818	
	8470871	Dec 21, 2025			U-2639	
	9084778	May 30, 2033	DP		U-134	
	9498465	May 30, 2033	DP		U-1033	
<u>TRIEPTANOIN - DOJOLVI</u>						
N 213687	001 8697748	Oct 03, 2025	DP			
	9186344	Jul 01, 2025	DP			
<u>TRILACICLIB DIHYDROCHLORIDE - COSELA</u>						
N 214200	001 10085992	Mar 14, 2034			U-3081	
	10189849	Oct 25, 2031	DS			
	10189850	Oct 25, 2031			DP	
	10927120	Oct 25, 2031			DP	
	10966984	Mar 14, 2034			U-3079	
	10966984	Mar 14, 2034			U-3080	
	11040042	Mar 14, 2034			DP	
	8598186	Oct 25, 2031	DS DP			
	8598197	Oct 25, 2031	DS DP			
	9487530	Mar 14, 2034			U-3079	
	9487530	Mar 14, 2034			U-3080	
	9957276	Oct 25, 2031	DS			
<u>TRIPTORELIN PAMOATE - TRELSTAR</u>						
N 022437	001 10166181	Jun 30, 2029	DP			
<u>TRIPTORELIN PAMOATE - TRIPTODUR KIT</u>						
N 208956	001 10166181	Jun 30, 2029	DP			
					ODE-149	Jun 29, 2024
<u>TUCATINIB - TUKYSA</u>						
N 213411	001 7452895	Nov 16, 2024	DS DP		U-2788	
	8648087	Apr 12, 2031	DS DP			
	9457093	Oct 12, 2032			DP U-2788	
	9693989	May 09, 2027			DP U-2788	
<u>TUCATINIB - TUKYSA</u>						
N 213411	002 7452895	Nov 16, 2024	DS DP		U-2788	
	8648087	Apr 12, 2031	DS DP			
	9457093	Oct 12, 2032			DP U-2788	
	9693989	May 09, 2027			DP U-2788	

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<u>UBROGEPANT - UBRELVY</u>						
N 211765	001	10117836				
		Jan 30, 2035	DP		NCE	Dec 23, 2024
		8754096	Jul 19, 2032	DS DP U-2717		
		8912210	Nov 10, 2031	DS DP		
		9499545	Nov 10, 2031	DS DP U-2718		
		9833448	Nov 10, 2031	U-2718		
<u>UBROGEPANT - UBRELVY</u>						
N 211765	002	10117836				
		Jan 30, 2035	DP		NCE	Dec 23, 2024
		8754096	Jul 19, 2032	DS DP U-2717		
		8912210	Nov 10, 2031	DS DP		
		9499545	Nov 10, 2031	DS DP U-2718		
		9833448	Nov 10, 2031	U-2718		
<u>ULIPRISTAL ACETATE - ELLA</u>						
N 022474	001	10159681				
		Apr 13, 2030		U-2510		
		10772897	Apr 13, 2030	U-2958		
		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		
		9844510	Dec 08, 2028	DP		
<u>UMBRALISIB TOSYLATE - UKONIO</u>						
N 213176	001	10072013				
		Jul 02, 2033		U-3063	NCE	Feb 05, 2026
		10072013	Jul 02, 2033	U-3064	ODE-343	Feb 05, 2028
		10414773	May 26, 2035	DS DP U-3063	ODE-344	Feb 05, 2028
		10414773	May 26, 2035	DS DP U-3064		
		10570142	Jul 02, 2033	DS DP U-3063		
		10570142	Jul 02, 2033	DS DP U-3064		
		10947244	May 26, 2035	U-3063		
		10947244	May 26, 2035	U-3064		
		9150579	Jul 02, 2033	DS DP		
		9669033	Jul 02, 2033	U-3063		
		9669033	Jul 02, 2033	U-3064		
		9969740	May 26, 2035	DS DP U-3063		
		9969740	May 26, 2035	DS DP U-3064		
<u>UMECLIDINIUM BROMIDE - INCRUSE ELLIPTA</u>						
N 205382	001	7488827				
		Dec 18, 2027	DS DP		M-245	Jun 09, 2022
		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	11090294				
		Nov 29, 2030		U-3203	M-245	Jun 09, 2022
		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		
		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		

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<u>UPADACITINIB - RINVOQ</u>						
N 211675	001	10519164	Oct 17, 2036	DP	NCE	Aug 16, 2024
		10597400	Oct 17, 2036	U-3255		
		10981923	Oct 17, 2036	DS		
		10981924	Oct 17, 2036	DP		
		11186584	Oct 17, 2036	DS		
		11198697	Oct 17, 2036	DP		
		8962629	Jan 15, 2031	DS U-3255		
		9951080	Oct 17, 2036	DS DP		
		9963459	Oct 17, 2036	DP		
		RE47221	Dec 01, 2030	DS		
<u>URIDINE TRIACETATE - VISTOGARD</u>						
N 208159	001	6258795	Jul 10, 2023	DP	ODE-104	Dec 11, 2022
		7776838	Aug 17, 2027	U-1791		
<u>URIDINE TRIACETATE - XURIDEN</u>						
N 208169	001	6258795	Jul 10, 2023	DP	ODE-98	Sep 04, 2022
<u>VALBENAZINE TOSYLATE - INGREZZA</u>						
N 209241	001	10065952	Oct 28, 2036	DS DP U-1995	NCE	Apr 11, 2022
		10844058	Oct 28, 2036	DS DP U-1995		
		10851103	Oct 28, 2036	DS DP U-1995		
		10851104	Oct 28, 2036	DS U-1995		
		10857137	Oct 10, 2037	U-1995		
		10857148	Oct 10, 2037	U-1995		
		10874648	Oct 10, 2037	U-1995		
		10874648	Oct 10, 2037	U-3046		
		10906902	Dec 22, 2036	DS DP		
		10906903	Dec 22, 2036	DS DP		
		10912771	Oct 10, 2037	U-1995		
		10912771	Oct 10, 2037	U-3076		
		10919892	Dec 22, 2036	DS DP		
		10940141	Aug 10, 2040	U-1995		
		10952997	Oct 10, 2037	U-1995		
		10993941	Oct 10, 2037	U-1995		
		11026931	Aug 14, 2039	U-1995		
		11026939	Sep 18, 2038	DP U-1995		
		11040029	Oct 10, 2037	U-1995		
		8039627	Oct 06, 2029	DS DP		
		8357697	Nov 08, 2027	U-1995		
<u>VALBENAZINE TOSYLATE - INGREZZA</u>						
N 209241	002	10065952	Oct 28, 2036	DS DP U-1995	NCE	Apr 11, 2022
		10844058	Oct 28, 2036	DS DP U-1995		
		10851103	Oct 28, 2036	DS DP U-1995		
		10851104	Oct 28, 2036	DS U-1995		
		10857137	Oct 10, 2037	U-1995		
		10857148	Oct 10, 2037	U-1995		
		10874648	Oct 10, 2037	U-1995		
		10874648	Oct 10, 2037	U-3046		
		10906902	Dec 22, 2036	DS DP		
		10906903	Dec 22, 2036	DS DP		
		10912771	Oct 10, 2037	U-1995		
		10912771	Oct 10, 2037	U-3076		
		10919892	Dec 22, 2036	DS DP		
		10940141	Aug 10, 2040	U-1995		
		10952997	Oct 10, 2037	U-1995		
		10993941	Oct 10, 2037	U-1995		
		11026931	Aug 14, 2039	U-1995		
		11026939	Sep 18, 2038	DP U-1995		
		11040029	Oct 10, 2037	U-1995		
		8039627	Oct 06, 2029	DS DP		
		8357697	Nov 08, 2027	U-1995		
<u>VALBENAZINE TOSYLATE - INGREZZA</u>						
N 209241	003	10065952	Oct 28, 2036	DS DP U-1995	NCE	Apr 11, 2022
		10844058	Oct 28, 2036	DS DP U-1995		
		10851103	Oct 28, 2036	DS DP U-1995		
		10851104	Oct 28, 2036	DS U-1995		
		10857137	Oct 10, 2037	U-1995		

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<u>VALBENZAZINE TOSYLATE - INGREZZA</u>						
N 209241	003	10857148	Oct 10, 2037		U-1995	
		10874648	Oct 10, 2037		U-1995	
		10874648	Oct 10, 2037		U-3046	
		10906902	Dec 22, 2036	DS DP		
		10906903	Dec 22, 2036	DS DP		
		10912771	Oct 10, 2037		U-1995	
		10912771	Oct 10, 2037		U-3076	
		10919892	Dec 22, 2036	DS DP		
		10940141	Aug 10, 2040		U-1995	
		10952997	Oct 10, 2037		U-1995	
		10993941	Oct 10, 2037		U-1995	
		11026931	Aug 14, 2039		U-1995	
		11026939	Sep 18, 2038	DP	U-1995	
		11040029	Oct 10, 2037		U-1995	
		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>VALSARTAN - DIOVAN</u>						
N 021283	001				NPP	Apr 19, 2024
<u>VALSARTAN - DIOVAN</u>						
N 021283	002				NPP	Apr 19, 2024
<u>VALSARTAN - DIOVAN</u>						
N 021283	004				NPP	Apr 19, 2024
<u>VANCOMYCIN - VANCOMYCIN</u>						
N 213895	001	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11000567	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - FIRVANO KIT</u>						
N 208910	001	10493028	Mar 13, 2035	DP		
		10688046	Mar 13, 2035	DP		
		10959946	Mar 13, 2035	DP		
		10959947	Mar 13, 2035	DP	U-3104	
		10959947	Mar 13, 2035	DP	U-3105	
		10959948	Mar 13, 2035	DP	U-3104	
		10959948	Mar 13, 2035	DP	U-3105	
		10959949	Mar 13, 2035	DP	U-3104	
		10959949	Mar 13, 2035	DP	U-3105	
<u>VANCOMYCIN HYDROCHLORIDE - FIRVANO KIT</u>						
N 208910	002	10493028	Mar 13, 2035	DP		
		10688046	Mar 13, 2035	DP		
		10959946	Mar 13, 2035	DP		
		10959947	Mar 13, 2035	DP	U-3104	
		10959947	Mar 13, 2035	DP	U-3105	
		10959948	Mar 13, 2035	DP	U-3104	
		10959948	Mar 13, 2035	DP	U-3105	
		10959949	Mar 13, 2035	DP	U-3104	
		10959949	Mar 13, 2035	DP	U-3105	
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	001	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	002	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		

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<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	003	10039804	Nov 06, 2035	DP U-282		
		10188697	Nov 06, 2035	DP U-282		
		10849956	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	004	10039804	Nov 06, 2035	DP U-282		
		10188697	Nov 06, 2035	DP U-282		
		10849956	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	005	10039804	Nov 06, 2035	DP U-282		
		10188697	Nov 06, 2035	DP U-282		
		10849956	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	006	10039804	Nov 06, 2035	DP U-282		
		10188697	Nov 06, 2035	DP U-282		
		10849956	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	007	10039804	Nov 06, 2035	DP U-282		
		10188697	Nov 06, 2035	DP U-282		
		10849956	Nov 06, 2035	DP		
<u>VANDETANIB - CAPRELSA</u>						
N 022405	001	8067427	Aug 08, 2028	DP		
		8642608	Feb 06, 2022		U-1490	
		RE42353	Jun 27, 2022	DS DP		
<u>VANDETANIB - CAPRELSA</u>						
N 022405	002	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		
<u>VARENICLINE TARTRATE - CHANTIX</u>						
N 021928	001	6890927	May 06, 2022	DS DP U-56	M-237	Feb 22, 2022
		6890927*PED	Nov 06, 2022		PED	Aug 22, 2022
		7265119	Aug 03, 2022	DS DP U-56		
		7265119*PED	Feb 03, 2023			
<u>VARENICLINE TARTRATE - CHANTIX</u>						
N 021928	002	6890927	May 06, 2022	DS DP U-56	M-237	Feb 22, 2022
		6890927*PED	Nov 06, 2022		PED	Aug 22, 2022
		7265119	Aug 03, 2022	DS DP U-56		
		7265119*PED	Feb 03, 2023			
<u>VARENICLINE TARTRATE - TYRVAYA</u>						
N 213978	001	10456396	Oct 19, 2035	DP U-1900	NP	Oct 15, 2024
		9504644	Oct 19, 2035	U-1900		
		9504645	Oct 19, 2035	DP		
		9532944	Oct 19, 2035	U-1900		
		9597284	Oct 19, 2035	U-1900		

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<u>VARENICLINE TARTRATE - TYRVAYA</u>						
N 213978	001	10456396	Oct 19, 2035	DP	U-1900	NP
		9504644	Oct 19, 2035		U-1900	
		9504645	Oct 19, 2035	DP		
		9532944	Oct 19, 2035		U-1900	
		9597284	Oct 19, 2035		U-1900	
<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>VASOPRESSIN - VASOSTRICT</u>						
N 204485	003	10010575	Jan 30, 2035		U-1857	
		9919026	Jan 30, 2035	DP		
		9925233	Jan 30, 2035		U-1857	
		9925234	Jan 30, 2035		U-1857	
		9962422	Jan 30, 2035		U-1857	
		9968649	Jan 30, 2035		U-1857	
		9974827	Jan 30, 2035		U-1857	
		9981006	Jan 30, 2035		U-1857	
<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485	004	10010575	Jan 30, 2035		U-1857	
		9919026	Jan 30, 2035	DP		
		9925233	Jan 30, 2035		U-1857	
		9925234	Jan 30, 2035		U-1857	
		9962422	Jan 30, 2035		U-1857	
		9968649	Jan 30, 2035		U-1857	
		9974827	Jan 30, 2035		U-1857	
		9981006	Jan 30, 2035		U-1857	
<u>VEMURAFENIB - ZELBORAF</u>						
N 202429	001	7504509	Oct 22, 2026	DS DP		ODE-158
		7863288	Jun 20, 2029	DS DP		
		8143271	Jun 21, 2026	DS DP		
		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	10730873	Nov 21, 2031	DS		I-795
		10993942	Sep 06, 2033		U-3114	M-265
		11110087	Sep 06, 2033		U-3222	ODE-114
		11110087	Sep 06, 2033		U-3223	ODE-185
		8546399	Jun 27, 2031	DS DP		ODE-211
		8722657	Jan 29, 2032	DS		ODE-239
		9174982	May 26, 2030		U-2323	
		9174982	May 26, 2030		U-2445	
		9174982	May 26, 2030		U-2446	
		9174982	May 26, 2030		U-2537	
		9539251	Sep 06, 2033		U-2538	
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573	002	10730873	Nov 21, 2031	DS		I-795
		10993942	Sep 06, 2033		U-3114	M-265
		11110087	Sep 06, 2033		U-3222	ODE-114
		11110087	Sep 06, 2033		U-3223	ODE-185
		8546399	Jun 27, 2031	DS DP		ODE-211
		8722657	Jan 29, 2032	DS		ODE-239

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<u>VENETOCLAX - VENCLEXTA</u>						
N 208573	002	9174982	May 26, 2030	U-2323		
		9174982	May 26, 2030	U-2445		
		9174982	May 26, 2030	U-2446		
		9174982	May 26, 2030	U-2537		
		9539251	Sep 06, 2033	U-2538		
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573	003	10730873	Nov 21, 2031	DS	I-795	May 15, 2022
		10993942	Sep 06, 2033	U-3114	M-265	Oct 16, 2023
		11110087	Sep 06, 2033	U-3222	ODE-114	Apr 11, 2023
		11110087	Sep 06, 2033	U-3223	ODE-185	Jun 08, 2025
		8546399	Jun 27, 2031	DS DP	ODE-211	Nov 21, 2025
		8722657	Jan 29, 2032	DS	ODE-239	May 15, 2026
		9174982	May 26, 2030	U-2323		
		9174982	May 26, 2030	U-2445		
		9174982	May 26, 2030	U-2446		
		9174982	May 26, 2030	U-2537		
		9539251	Sep 06, 2033	U-2538		
<u>VERICIGUAT - VEROUVO</u>						
N 214377	001	10736896	May 19, 2031	DS DP	NCE	Jan 19, 2026
		8420656	May 19, 2031	DS DP		
		8921377	May 19, 2031	U-3062		
		9604948	Nov 26, 2032	DS DP	U-3062	
		9993476	May 19, 2031	U-3062		
<u>VERICIGUAT - VEROUVO</u>						
N 214377	002	10736896	May 19, 2031	DS DP	NCE	Jan 19, 2026
		8420656	May 19, 2031	DS DP		
		8921377	May 19, 2031	U-3062		
		9604948	Nov 26, 2032	DS DP	U-3062	
		9993476	May 19, 2031	U-3062		
<u>VERICIGUAT - VEROUVO</u>						
N 214377	003	10736896	May 19, 2031	DS DP	NCE	Jan 19, 2026
		8420656	May 19, 2031	DS DP		
		8921377	May 19, 2031	U-3062		
		9604948	Nov 26, 2032	DS DP	U-3062	
		9993476	May 19, 2031	U-3062		
<u>VIBEGRON - GEMTESA</u>						
N 213006	001	8247415	Dec 01, 2030	DS DP	U-3045	NCE
		8653260	Apr 02, 2029	DS		Dec 23, 2025
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N 022567	001	7834020	Jun 05, 2022	DS DP	U-839	M-254
		7834020*PED	Dec 05, 2022			PED
		8193195	Jun 05, 2022		U-839	
		8193195*PED	Dec 05, 2022			
		8236804	Jun 05, 2022		U-839	
		8236804*PED	Dec 05, 2022			
		8673921	Jun 05, 2022	DS DP		
		8673921*PED	Dec 05, 2022			
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N 022567	002	7834020	Jun 05, 2022	DS DP	U-839	M-254
		7834020*PED	Dec 05, 2022			PED
		8193195	Jun 05, 2022		U-839	
		8193195*PED	Dec 05, 2022			
		8236804	Jun 05, 2022		U-839	
		8236804*PED	Dec 05, 2022			
		8673921	Jun 05, 2022	DS DP		
		8673921*PED	Dec 05, 2022			
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N 022567	003	7834020	Jun 05, 2022	DS DP	U-839	M-254
		7834020*PED	Dec 05, 2022			PED
		8193195	Jun 05, 2022		U-839	
		8193195*PED	Dec 05, 2022			
		8236804	Jun 05, 2022		U-839	

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<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N 022567	003	8236804*PED	Dec 05, 2022			
		8673921	Jun 05, 2022	DS DP		
		8673921*PED	Dec 05, 2022			
<u>VILOXAZINE HYDROCHLORIDE - OELBREE</u>						
N 211964	001	9358204	Feb 07, 2033	DP	NCE	Apr 02, 2026
		9603853	Feb 07, 2033	U-727		
		9662338	Feb 07, 2033	DP		
<u>VILOXAZINE HYDROCHLORIDE - OELBREE</u>						
N 211964	002	9358204	Feb 07, 2033	DP	NCE	Apr 02, 2026
		9603853	Feb 07, 2033	U-727		
		9662338	Feb 07, 2033	DP		
<u>VILOXAZINE HYDROCHLORIDE - OELBREE</u>						
N 211964	003	9358204	Feb 07, 2033	DP	NCE	Apr 02, 2026
		9603853	Feb 07, 2033	U-727		
		9662338	Feb 07, 2033	DP		
<u>VILTOLARSEN - VILTEPSO</u>						
N 212154	001	10870676	Aug 31, 2031	DS DP U-3039	NCE	Aug 12, 2025
		9079934	Aug 31, 2031	DS DP	ODE-280	Aug 12, 2027
<u>VISMODEGIB - ERIVEDGE</u>						
N 203388	001	7888364	Nov 11, 2028	DS DP		
		9278961	Dec 15, 2028	U-1825		
		9790183	Sep 02, 2025	U-3109		
<u>VOCLOSPORIN - LUPKYNIS</u>						
N 213716	001	10286036	Dec 07, 2037	U-3056	NCE	Jan 22, 2026
		7332472	Oct 17, 2022	DS DP U-3056		
<u>VORAPAXAR SULFATE - ZONTIVITY</u>						
N 204886	001	7304078	Apr 06, 2024	DS DP U-1512		
		7713999	May 30, 2024	DS DP U-2291		
<u>VORICONAZOLE - VFEND</u>						
N 021266	001				NPP	Jan 29, 2022
<u>VORICONAZOLE - VFEND</u>						
N 021266	002				NPP	Jan 29, 2022
<u>VORICONAZOLE - VFEND</u>						
N 021267	001				NPP	Jan 29, 2022
<u>VORICONAZOLE - VFEND</u>						
N 021630	001				NPP	Jan 29, 2022
<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	M-187	Jan 22, 2024
		8476279	Oct 02, 2022	DP U-1439	M-267	Nov 13, 2023
		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		

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<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447	002	7144884	Jun 17, 2026	DS DP U-1439	M-187	Jan 22, 2024
		8476279	Oct 02, 2022	DP U-1439	M-267	Nov 13, 2023
		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	003	7144884	Jun 17, 2026	DS DP U-1439	M-187	Jan 22, 2024
		8476279	Oct 02, 2022	DP U-1439	M-267	Nov 13, 2023
		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-187	Jan 22, 2024
		8476279	Oct 02, 2022	DP U-1439	M-267	Nov 13, 2023
		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>VOSORITIDE - VOXZOGO</u>						
N 214938	001	10646550	Aug 01, 2036	DP U-3256	NCE	Nov 19, 2026
		8198242	Jun 11, 2030	DS DP U-3256		
		9907834	Aug 01, 2036	DP		
		RE48267	May 20, 2030	U-3256		
<u>VOSORITIDE - VOXZOGO</u>						
N 214938	002	10646550	Aug 01, 2036	DP U-3256	NCE	Nov 19, 2026
		8198242	Jun 11, 2030	DS DP U-3256		
		9907834	Aug 01, 2036	DP		
		RE48267	May 20, 2030	U-3256		
<u>VOSORITIDE - VOXZOGO</u>						
N 214938	003	10646550	Aug 01, 2036	DP U-3256	NCE	Nov 19, 2026
		8198242	Jun 11, 2030	DS DP U-3256		
		9907834	Aug 01, 2036	DP		
		RE48267	May 20, 2030	U-3256		
<u>VOXELOTOR - OXBRYTA</u>						
N 213137	001	10017491	Dec 28, 2032	DP	NCE	Nov 25, 2024
		10034879	Dec 28, 2032	DS DP	ODE-281	Nov 25, 2026
		10493035	Oct 12, 2037	DP		
		10722502	Feb 06, 2035	DP		
		10806733	Dec 28, 2032	DS		
		11020382	Dec 02, 2036	U-3133		
		11020382	Dec 02, 2036	U-3134		
		9018210	Dec 28, 2032	DS DP		
		9248199	Jan 29, 2034	U-2676		
		9248199	Jan 29, 2034	U-2715		
		9447071	Feb 06, 2035	DS DP		
<u>VOXELOTOR - OXBRYTA</u>						
N 216157	001				NCE	Nov 25, 2024

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<u>ZANUBRUTINIB - BRUKINSA</u>						
N 213217	001	10570139	Apr 22, 2034	DS DP U-1745	I-871	Aug 31, 2024
		10570139	Apr 22, 2034	DS DP U-2145	I-874	Sep 14, 2024
		10570139	Apr 22, 2034	DS DP U-3063	NCE	Nov 14, 2024
		10927117	Aug 15, 2037	DS DP	ODE-276	Nov 14, 2026
		11142528	Apr 22, 2034	DS DP U-1745	ODE-370	Sep 14, 2028
		11142528	Apr 22, 2034	DS DP U-2145	ODE-371	Aug 31, 2028
		11142528	Apr 22, 2034	DS DP U-3063		
		9447106	Apr 22, 2034	DS DP U-1745		
		9447106	Apr 22, 2034	DS DP U-2145		
		9447106	Apr 22, 2034	DS DP U-3063		
<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>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>ZOLPIDEM TARTRATE - EDLUAR</u>						
N 021997	001	9265720	Feb 25, 2031	U-674		
		9597281	Apr 06, 2027	U-674		
<u>ZOLPIDEM TARTRATE - EDLUAR</u>						
N 021997	002	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		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<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			

PATENT AND EXCLUSIVITY TERMS

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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)
ODE*	FDA has not recognized Orphan-Drug Exclusivity (ODE) for this drug, but it contains the same active moiety or moieties as another drug(s) that was eligible for ODE, and also shares ODE-protected use(s) or indication(s) with that drug(s). An application seeking approval for the same active moiety or moieties, including an ANDA that cites this NDA as its basis of submission, may not be approved for such ODE-protected use(s) and indication(s)
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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

- 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/M2 OR 175MG/M2 INTRAVENOUSLY OVER THREE HOURS EVERY THREE WEEKS
- D-25 ADDITIONAL DOSAGE REGIMEN EQUAL TO HALF THE ORIGINAL DOSING REGIMEN
- 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 THIOPENTAL
- 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

- 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
- 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

- 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
- 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 ALLERGIC 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 SULFATE CO-ADMINISTERED WITH RITONAVIR FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT NAIVE PATIENTS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

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
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.

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

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
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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

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 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
D-174	MODIFICATIONS TO THE EXISTING DOSING REGIMEN TO ALLOW FOR TREATMENT INTERRUPTIONS OF UP TO 8 WEEKS FOR INTOLERABLE ADVERSE REACTIONS
D-175	EIGHT-WEEK DOSING REGIMEN FOR THE TREATMENT OF GENOTYPES 1, 2, 3, 4, 5, AND 6, CHRONIC HEPATITIS C VIRUS INFECTION IN TREATMENT-NAIVE SUBJECTS WITH COMPENSATED CIRRHOSIS BASED ON THE RESULTS FROM THE EXPEDITION-8 STUDY
D-176	IBRUTINIB IN COMBINATION WITH RITUXIMAB
D-177	INFORMATION ADDED TO THE DOSING SECTION IN REGARD TO THE TREATMENT OF CHRONIC HEPATITIS C VIRUS INFECTION IN PATIENTS WITH SEVERE RENAL IMPAIRMENT INCLUDING PATIENTS WITH END STAGE RENAL DISEASE ON DIALYSIS
D-181	DOSING REGIMEN EXTENDING THE CONTRACEPTION USE FROM 5 YEARS TO UP TO 6 YEARS
D-182	NEW DOSING REGIMEN FOR THE PREVENTION AND MANAGEMENT OF NERATINIBASSOCIATED DIARRHEA

EXCLUSIVITY INDICATION

I-1	DYSMENORRHEA
I-2	CHOLANGIOPANCREATOGRAPHY
I-3	INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY
I-4	PERIPHERAL VENOGRAPHY (PHLEBOGRAPHY)
I-5	HYSTEOSALPINGOGRAPHY
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 VENOUS 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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
- 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-59	ENDOSCOPICALLY DIAGNOSED ESOPHAGITIS, INCLUDING EROSIVE 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
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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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
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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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
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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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
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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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

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 PRILOSEC (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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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 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)

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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 ADMINISTERED 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 GLUCOCORTICOID 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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

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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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
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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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
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 \geq 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- 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 REVASCLARIZATION PROCEDURES
- 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 ARIPIPIRAZOLE 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 PAPANILLA 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- 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
- 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- 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
- 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- 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
- 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. THETAIOAOMICRON 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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
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)

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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

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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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 SIEZURES 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-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 NONSQAUMOUS 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY 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

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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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
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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

(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
- 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- 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 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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 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-790	USE OF FERRIC CITRATE FOR THE TREATMENT OF IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WITH CKD NOT ON DIALYSIS
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
I-792	TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA WHO HAVE BEEN PREVIOUSLY TREATED WITH SORAFENIB
I-793	TREATMENT OF MODERATE TO SEVERE VAGINAL DRYNESS, A SYMPTOM OF VULVAR AND VAGINAL ATROPHY, DUE TO MENOPAUSE
I-794	TREATMENT OF ADULT PATIENTS WITH METASTATIC GASTRIC OR GASTROESOPHAGEAL JUNCTION ADENOCARCINOMA PREVIOUSLY TREATED WITH AT LEAST TWO PRIOR LINES OF CHEMOTHERAPY, AND IF APPROPRIATE, HER2/NEU-TARGETED THERAPY
I-795	VENETOCLAX IN COMBINATION WITH OBINUTUZUMAB IN PREVIOUSLY UNTREATED PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LYMPHOMA
I-796	USED IN COMBINATION WITH A RITUXIMAB PRODUCT, ARE INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED FOLLICULAR LYMPHOMA (FL)
I-797	USED IN COMBINATION WITH A RITUXIMAB PRODUCT, ARE INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED MARGINAL ZONE LYMPHOMA (MZL)
I-798	TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER (BIPOLAR DEPRESSION)
I-799	TREATMENT OF STEROID-REFRACTORY ACUTE GRAFT-VERSUS-HOST DISEASE (GVHD) IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER
I-800	TREATMENT OF OCULAR INFLAMMATION FOLLOWING OPHTHALMIC SURGERY
I-801	USE IN CARDIAC MRI TO ASSESS MYOCARDIAL PERFUSION (STRESS, REST) AND LATE GADOLINIUM ENHANCEMENT IN ADULT PATIENTS WITH KNOWN OR SUSPECTED CORONARY ARTERY DISEASE (CAD)
I-802	TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC IMMUNE THROMBOCYTOPENIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-803	TREATMENT OF ADULT PATIENTS WITH ORAL ULCERS ASSOCIATED WITH BEHCETS DISEASE
I-804	EXPANDED INDICATION FOR USE AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
I-805	SLOW THE RATE OF DECLINE IN PULMONARY FUNCTION IN PATIENTS WITH SYSTEMIC SCLEROSIS-ASSOCIATED INTERSTITIAL LUNG DISEASE
I-806	EXPANDED INDICATION FOR PTS WHO ARE VIROLOGICALLY SUPPRESSED (HIV-1 RNA LESS THAN 50 COPIES/ML) ON A STABLE ARV REGIMEN WITH NO HX OF TX FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED W RESISTANCE TO DORAVIRINE, LAMIVUDINE OR TENOFOVIR DISOPROXIL FUMARATE
I-807	TREATMENT OF ADVANCED ENDOMETRIAL CARCINOMA THAT IS NOT MICROSATELLITE INSTABILITY-HIGH OR MISMATCH REPAIR DEFICIENT, WHO HAVE DISEASE PROGRESSION FOLLOWING PRIOR SYSTEMIC THERAPY AND ARE NOT CANDIDATES FOR CURATIVE SURGERY OR RADIATION
I-808	TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (MCSPC)
I-809	TO REDUCE THE RISK OF END-STAGE KIDNEY DISEASE, DOUBLING OF SERUM CREATININE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND DIABETIC NEPHROPATHY WITH ALBUMINURIA > 300 MG/DAY
I-810	PROPHYLAXIS OF VENOUS THROMBOEMBOLISM IN ACUTELY ILL MEDICAL PATIENTS AT RISK FOR THROMBOEMBOLIC COMPLICATIONS NOT AT HIGH RISK OF BLEEDING
I-811	TREATMENT OF ACUTE UNCOMPLICATED INFLUENZA IN PATIENTS 12 YEARS OF AGE OR OLDER, WHO HAVE BEEN SYMPTOMATIC FOR NO MORE THAN 48 HOURS AND ARE AT HIGH RISK OF DEVELOPING INFLUENZA-RELATED COMPLICATIONS
I-812	FOR USE IN AT RISK ADULTS AND ADOLESCENTS WEIGHING AT LEAST 35 KG FOR PRE-EXPOSURE PROPHYLAXIS TO REDUCE THE RISK OF HIV-1 INFECTION FROM SEXUAL ACQUISITION, EXCLUDING INDIVIDUALS AT RISK FROM RECEPTIVE VAGINAL SEX
I-813	TX OF ADULT PTS W/ ADV OVARIAN FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER TREATED W/ >=3 PRIOR CHEMO REGIMENS & ASSOCIATED W/ HRD DEFICIENCY POSITIVE STATUS DEFINED BY A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION
I-814	TX OF ADV OVARIAN FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER TREATED W/ >=3 PRIOR CHEMO REGIMENS & ASSOCIATED W/ HRD DEFICIENCY DEFINED BY POSITIVE STATUS GENOMIC INSTABILITY & WHO HAVE PROGRESSED >6MO AFTER RESPONSE TO LAST PLATINUM-BASED CHEMO
I-815	TREATMENT OF COMMUNITY ACQUIRED BACTERIAL PNEUMONIA (CABP) CAUSED BY DESIGNATED SUSCEPTIBLE BACTERIA IN ADULTS
I-816	TREATMENT OF ADULT PATIENTS WITH NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) WHO ARE >= 75 YEARS OLD OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
I-817	TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)
I-818	MAINTENANCE TREATMENT OF ADULT PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GBRAM METASTATIC PANCREATIC ADENOCARCINOMA WHOSE DISEASE HAS NOT PROGRESSED ON AT LEAST 16 WEEKS OF A FIRST-LINE PLATINUM-BASED CHEMOTHERAPY REGIMEN
I-819	ADJUNCT TO MAX TOLERATED STATIN TX TO REDUCE RISK OF MI, STROKE, CORONARY REVASCULARIZATION, & UNSTABLE ANGINA REQUIRING HOSPITALIZATION IN ADULTS W/ ELEVATED TG LEVELS & ESTABLISHED CV DISEASE OR DIABETES MELLITUS & 2+ RISK FACTORS FOR CV DISEASE
I-820	INDICATED FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) TO DELAY DISEASE PROGRESSION
I-821	TREATMENT OF CANDIDEMIA, ACUTE DISSEMINATED CANDIDIASIS, CANDIDA PERITONITIS AND ABSCESSSES WITHOUT MENINGOENCEPHALITIS AND/OR OCULAR DISSEMINATION IN PEDIATRIC PATIENTS YOUNGER THAN 4 MONTHS OF AGE
I-822	REDUCE THE RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS (CARDIOVASCULAR DEATH, NON-FATAL MYOCARDIAL INFARCTION OR NON-FATAL STROKE) IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE
I-823	USE IN COMBINATION WITH CAPECITABINE, FOR THE TREATMENT OF ADULT PATIENTS WITH ADVANCED OR METASTATIC HER2-POSITIVE BREAST CANCER WHO HAVE RECEIVED TWO OR MORE PRIOR ANTI-HER2 BASED REGIMENS IN THE METASTATIC SETTING
I-824	RIVAROXBAN IN COMBINATION WITH ASPIRIN, IS INDICATED TO REDUCE THE RISK OF

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- MAJOR CV EVENTS (CV DEATH, MI, AND STROKE) IN PATIENTS WITH CHRONIC CORONARY ARTERY DISEASE (CAD) OR PERIPHERAL ARTERY DISEASE (PAD)
- I-825 TREATMENT FOR CHRONIC FIBROSING INTERSTITIAL LUNG DISEASES WITH A PROGRESSIVE PHENOTYPE
- I-826 ENCORAFENIB, IN COMBINATION WITH CETUXIMAB, FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC COLORECTAL CANCER WITH A BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST, AFTER PRIOR THERAPY
- I-827 EXPANDED INDICATION FOR PATIENTS WHO ARE VIROLOGICALLY SUPPRESSED (HIV-1 RNA LESS THAN 50 COPIES/ML) ON A STABLE ARV REGIMEN WITH NO HISTORY OF TREATMENT FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED WITH RESISTANCE TO DORAVIRINE
- I-828 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- I-829 TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH CENTRAL PRECOCIOUS PUBERTY (CPP)
- I-830 TREATMENT OF ADULT PATIENTS WITH A DELETERIOUS BRCA MUTATION (GERMLINE AND/OR SOMATIC)-ASSOCIATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE BEEN TREATED WITH ANDROGEN RECEPTOR-DIRECTED THERAPY AND A TAXANE-BASED CHEMOTHERAPY
- I-831 W/BEVACIZUMAB FOR MAINTENANCE TX OF ADULTS W/ADV. EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CA IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMO & CA ASSOCIATED W/ HOMOLOGOUS RECOMBINATION DEFICIENCY POSITIVE STATUS
- I-832 TX OF ADULT PTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC HOMOLOGOUS RECOMBINATION REPAIR GENE-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE PROGRESSED FOLLOWING PRIOR TREATMENT WITH ENZALUTAMIDE OR ABIRATERONE
- I-833 MAINTENANCE TREATMENT OF ADULT PATIENTS WITH ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY
- I-834 TO REDUCE THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH HEART FAILURE (NYHA CLASS II-IV) WITH REDUCED EJECTION FRACTION
- I-835 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY (R/R) FOLLICULAR LYMPHOMA (FL) WHOSE TUMORS ARE POSITIVE FOR AN EZH2 MUTATION AS DETECTED BY AN FDA-APPROVED TEST AND WHO HAVE RECEIVED AT LEAST 2 PRIOR SYSTEMIC THERAPIES
- I-836 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY (R/R) FOLLICULAR LYMPHOMA (FL) WHO HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
- I-837 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL), NOT OTHERWISE SPECIFIED, INCLUDING DLBCL ARISING FROM FOLLICULAR LYMPHOMA, AFTER AT LEAST 2 LINES OF SYSTEMIC THERAPY
- I-838 TREATMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY (DPN) OF THE FEET
- I-839 TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPPRESSED (HIV-1 RNA LESS THAN 50 COPIES PER ML) ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO HISTORY OF TREATMENT FAILURE
- I-840 TREATMENT OF SYMPTOMS IN ADULTS WITH MAJOR DEPRESSIVE DISORDER (MDD) WITH ACUTE SUICIDAL IDEATION OR BEHAVIOR.
- I-841 TO REDUCE THE RISK OF HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE OR MULTIPLE CARDIOVASCULAR RISK FACTORS
- I-842 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY IN COMBINATION WITH DARATUMUMAB AND DEXAMETHASONE
- I-843 MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS AGED 18 YEARS AND OLDER
- I-844 INDICATED IN PATIENTS 18 YEARS OF AGE AND OLDER FOR THE TREATMENT OF HOSPITAL ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) CAUSED BY THE FOLLOWING SUSCEPTIBLE GRAM-NEGATIVE MICROORGANISMS: ACINETOBACTER BAUMANNII COMPLEX, ESCHERICHIA COLI, ENTEROBACTER CLOACAE COMPLEX, KLEBSIELLA PNEUMONIAE, PSEUDOMONAS AERUGINOSA, AND SERRATIA MARCESCENS
- I-845 TREATMENT OF ADULT PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA-APPROVED

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

TEST

- I-846 TREATMENT OF CATAPLEXY IN ADULT PATIENTS WITH NACROLEPSY
- I-847 EXPANDED INDICATION OF THE TREATMENT OF ADULT PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA-APPROVED TEST
- I-848 REDUCE THE RISK OF STROKE IN PATIENTS WITH ACUTE ISCHEMIC STROKE (NIH STROKE SCALE SCORE \leq 5) OR HIGH-RISK TRANSIENT ISCHEMIC ATTACK (TIA)
- I-849 CHRONIC PHASE (CP) CHRONIC MYELOID LEUKEMIA (CML) WITH RESISTANCE OR INTOLERANCE TO AT LEAST TWO PRIOR KINASE INHIBITORS
- I-850 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME (SMS) IN PATIENTS 16 YEARS OF AGE AND OLDER
- I-851 TO REDUCE THE RISK OF A FIRST MYOCARDIAL INFARCTION (MI) OR STROKE IN PATIENTS WITH CORONARY ARTERY DISEASE (CAD) AT HIGH RISK FOR SUCH EVENTS
- I-852 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER AND YOUNG ADULTS WITH RELAPSED OR REFRACTORY, SYSTEMIC ANAPLASTIC LARGE CELL LYMPHOMA (ALCL) THAT IS ALK-POSITIVE
- I-853 INDICATION OF OSIMERTINIB AS ADJUVANT THERAPY AFTER TUMOR RESECTION IN ADULT PATIENTS WITH 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
- I-854 FOR THE TREATMENT OF PATIENTS WITH ADVANCED RENAL CELL CARCINOMA, AS A FIRST-LINE TREATMENT IN COMBINATION WITH NIVOLUMAB
- I-855 TREATMENT OF NEUROGENIC DETRUSOR OVERACTIVITY (NDO) IN PEDIATRIC PATIENTS 3 YEARS AND OLDER AND WEIGHING 35 KILOGRAMS OR MORE
- I-856 INDICATION FOR THE TREATMENT OF PULMONARY HYPERTENSION ASSOCIATED WITH INTERSTITIAL LUNG DISEASE TO IMPROVE EXERCISE ABILITY
- I-857 TO REDUCE THE RISK OF SUSTAINED EGFR DECLINE, END-STAGE KIDNEY DISEASE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH CHRONIC KIDNEY DISEASE AT RISK OF PROGRESSION
- I-858 FOR THE TOPICAL TREATMENT OF SCABIES INFESTATIONS IN ADULT AND PEDIATRIC PATIENTS 4 YEARS OF AGE AND OLDER
- I-859 TREATMENT OF PATIENTS WITH TRANSFUSIONAL IRON OVERLOAD DUE TO SICKLE CELL DISEASE OR OTHER ANEMIAS
- I-860 FOR THE TREATMENT OF MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS IN ADULT PATIENTS
- I-861 TREATMENT OF NEUROGENIC DETRUSOR OVERACTIVITY (NDO) IN PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER AND WEIGHING GREATER THAN 25 KG
- I-862 TREATMENT OF VENOUS THROMBOEMBOLIC EVENTS (VTE) IN PEDIATRIC PATIENTS 8 TO LESS THAN 18 YEARS OF AGE WHO HAVE BEEN TREATED WITH A PARENTERAL ANTICOAGULANT FOR AT LEAST 5 DAYS AND TO REDUCE THE RISK OF RECURRENCE OF VTE IN PEDIATRIC PATIENTS 8 TO LESS THAN 18 YEARS OF AGE WHO HAVE BEEN PREVIOUSLY TREATED
- I-863 TREATMENT OF ADULT PATIENTS WITH ADVANCED SYSTEMIC MASTOCYTOSIS (ADVSM), INCLUDING PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM) AND SYSTEMIC MASTOCYTOSIS WITH AN ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN)
- I-864 TREATMENT OF ADULT PATIENTS WITH MAST CELL LEUKEMIA (MCL)
- I-865 FOR THE PREVENTIVE TREATMENT OF EPISODIC MIGRAINE IN ADULTS
- I-866 FOR THE TREATMENT OF TRICHOMONIASIS CAUSED BY TRICHOMONAS VAGINALIS IN ADULTS
- I-867 INDICATED TO REDUCE THE RISK OF MAJOR THROMBOTIC VASCULAR EVENTS (MYOCARDIAL INFARCTION, ISCHEMIC STROKE, ACUTE LIMB ISCHEMIA, AND MAJOR AMPUTATION OF VASCULAR ETIOLOGY) IN PATIENTS WITH PAD, INCLUDING PATIENTS WHO HAVE RECENTLY UNDERGONE A LOWER EXTREMITY REVASCULARIZATION PROCEDURE DUE TO SYMPTOMATIC PAD
- I-868 LENVATINIB IN COMBINATION WITH PEMBROLIZUMAB, IS INDICATED FOR THE FIRST-LINE TREATMENT OF ADULT PATIENTS WITH ADVANCED RENAL CELL CARCINOMA (RCC)
- I-869 REDUCE THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH HEART FAILURE AND REDUCED EJECTION FRACTION
- I-870 INDICATED FOR THE TREATMENT OF IDIOPATHIC HYPERSOMNIA (IH) IN ADULTS
- I-871 TREATMENT OF ADULT PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA (WM)

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-872 ADDITION OF THE INDICATION OF TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR TWO LINES OF SYSTEMIC THERAPY IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER
- I-873 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH LOCALLY ADVANCED OR METASTATIC DIFFERENTIATED THYROID CANCER (DTC) THAT HAS PROGRESSED FOLLOWING PRIOR VEGFR-TARGETED THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY OR INELIGIBLE
- I-874 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA (MZL) WHO HAVE RECEIVED AT LEAST ONE ANTI-CD20-BASED REGIMEN
- I-875 FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH AN IDH1 MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- I-876 TREATMENT OF OCULAR ITCHING ASSOCIATED WITH ALLERGIC CONJUNCTIVITIS
- I-877 INDICATION FOR THE USE OF ABEMACICLIB IN COMBINATION WITH ENDOCRINE THERAPY (TAMOXIFEN OR AN AROMATASE INHIBITOR) FOR THE ADJUVANT TREATMENT OF ADULT PATIENTS WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE, NODE-POSITIVE, EARLY CANCER (EBC) AT HIGH RISK OF RECURRENCE AND A KI-67 SCORE>20% AS DETERMINED BY AN FDA APPROVED TEST
- I-878 ADDITION OF A NEW INDICATION FOR ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- I-879 TREATMENT OF ADULT PATIENTS WITH ACTIVE ANKYLOSING SPONDYLITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS, TO THE PRESCRIBING INFORMATION
- I-880 TREATMENT OF ADULTS WITH ACTIVE PSORIATIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS

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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- 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
- 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"

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- 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
- 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- M-72 INFORMATION ABOUT USE OF INSPRA (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 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

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- 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 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

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- 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 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

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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
- 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

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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 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

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- 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.
- 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

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- 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 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 \geq 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 LOTEPIREDNOL 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
- M-236 INFORMATION ADDED TO THE PRESCRIBING INFORMATION TO INCLUDE EFFICACY AND SAFETY DATA FROM A STUDY IN PATIENTS WITH TREATMENT NAIVE CLL/SLL TREATED WITH IBRUTINIB IN COMBINATION WITH OBINUTUZUMAB OR CHLORAMBUCIL IN COMBINATION WITH OBINUTUZUMAB
- M-237 INFORMATION ADDED TO LABELING TO DESCRIBE A STUDY TO EVALUATE THE SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSATION IN ADOLESCENT SMOKERS
- M-238 INFORMATION ADDED TO THE PRESCRIBING INFORMATION TO REFLECT THAT NO DOSE ADJUSTMENT IS NEEDED FOR PATIENTS WITH AN ESTIMATED GLOMERULAR FILTRATION RATE (EGFR) OF 45 ML/MIN/1.73 M² OR GREATER AS SUPPORTED BY CLINICAL STUDY REPORT
- M-239 INFORMATION ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING REGARDING A TRIAL CONDUCTED IN TREATMENT NAIVE PEDIATRIC PATIENTS, AGES 2 YEARS TO $<$ 18 YEARS WITH TRANSFUSIONAL IRON OVERLOAD
- M-240 INFORMATION ADDED TO LABELING REGARDING A RANDOMIZED, PLACEBO-CONTROLLED CLINICAL TRIAL TO EVALUATE CARDIOVASCULAR OUTCOMES AFTER TREATMENT WITH EXENATIDE ONCE WEEKLY IN PATIENTS WITH TYPE 2 DIABETES MELLITES

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- M-241 INFORMATION ADDED TO THE LABELING FOR SAFETY & EFFICACY STUDY ENTITLED, A MULTICENTER, RANDOMIZED, DOUBLE-BLIND, PLACEBO CONTROLLED CLINICAL TRIAL OF DEFERASIROX IN PATIENTS WITH MYELODYSPLASTIC SYNDROMES (LOW/INT-1 RISK) & TRANSFUSIONAL IRON OVERLOAD
- M-242 INFORMATION ADDED TO THE LABELING REGARDING THE EFFICACY AND SAFETY OF INSULIN DEGLUDEC/LIRAGLUTIDE VS INSULIN GLARGINE IN PTS W/ TYPE 2 DIABETES INADEQUATELY CONTROLLED ON SGLT2I WITH OR WITHOUT ORAL ANTIDIABETIC THERAPIES
- M-243 INFORMATION ADDED TO LABELING FROM PROSPECTIVE, RANDOMIZED, OPEN-LABEL, BLIND EVALUATOR (PROBE) STUDY EVALUATING THE EFFICACY AND SAFETY OF LOW MOLECULAR WEIGHT HEPARIN/EDOXABAN VERSUS DALTEPARIN IN VENOUS THROMBOEMBOLISM ASSOCIATED WITH CANCER
- M-244 INFORMATION ADDED TO THE LABELING REGARDING EFFICACY AND SAFETY OF THE CONTINUATION OF SITAGLIPTIN COMPARED WITH THE WITHDRAWAL OF SITAGLIPTIN DURING INITIATION AND TITRATION OF INSULIN GLARGINE IN SUBJECTS WITH TYPE 2 DIABETES MELLITUS
- M-245 ADDITIONAL INFORMATION ADDED TO THE LABELING BASED ON SAFETY AND EFFICACY DATA FROM THE IMPACT TRIAL
- M-246 ADDITION OF STUDY BRf117277, A NON-RANDOMIZED, OPEN-LABEL, MULTI-CENTER, MULTI-COHORT TRIAL OF DABRAFENIB PLUS TRAMETINIB IN SUBJECTS WITH BRAF MUTATION-POSITIVE MELANOMA THAT HAS METASTASIZED TO THE BRAIN
- M-247 REVISIONS TO THE LABELING REGARDING CONTINUOUS SUBCUTANEOUS INSULIN INFUSION AS A CONDITION OF USE FOR INSULIN ASPART
- M-248 INFORMATION ADDED TO THE LABELING TO DESCRIBE A TRIAL EVALUATING A LOWER DOSE THAN THOSE APPROVED FOR PEDIATRIC PATIENTS 13 TO 17 YEARS OF AGE
- M-249 INFORMATION ADDED TO THE LABELING TO DESCRIBE STUDY LVM-MD-15 TO FULFILL POSTMARKETING COMMITMENT 1943-4
- M-250 REVISIONS TO THE PEDIATRIC USE SECTION TO INCLUDE AN OPEN-LABEL CLINICAL TRIAL TO FULFILL PMR 1655-1
- M-251 INFORMATION ADDED TO THE CLINICAL PHARMACOLOGY SECTION REGARDING INFLUENZA VIRUS RESISTANCE TO OSELTAMIVIR IN IMMUNOCOMPROMISED PATIENTS
- M-252 ADDITION OF INFORMATION TO CLINICAL STUDIES SECTION REGARDING CARDIOVASCULAR OUTCOME
- M-253 INFORMATION ADDED TO THE LABELING TO DESCRIBE STUDY P061, A RANDOMIZED, PLACEBO-CONTROLLED, PARALLEL GROUP, MULTI-SITE, DOUBLE-BLIND STUDY TO EVALUATE SAFETY AND EFFICACY OF SUVOREXANT FOR THE TREATMENT OF INSOMNIA IN SUBJECTS WITH ALZHEIMERS DISEASE
- M-254 INFORMATION ADDED TO THE LABELING REGARDING PEDIATRIC PATIENTS AGES 7 TO 17 YEARS OF AGE WITH MAJOR DEPRESSIVE DISORDER
- M-255 INFORMATION ADDED TO THE LABELING TO DESCRIBE STUDY GS-US-320-4018 IN VIROLOGICALLY SUPPRESSED ADULTS W/ CHRONIC HEP B INFECTION WHO SWITCHED FROM TENOFOVIR DISOPROXIL FUMARATE TO TENOFOVIR ALAFENAMIDE
- M-256 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION TO FULFILL A POST-MARKETING REQUIREMENT
- M-257 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING REGARDING THE USE OF PLAQUE PSORIASIS OF THE SCALP
- M-258 INFORMATION ADDED TO THE LABELING TO DESCRIBE CARMELINA TRIAL TO FULFILL POSTMARKETING COMMITMENT 1766-4
- M-259 INFORMATION ADDED TO THE LABELING REGARDING SAFETY AND EFFICACY IN SUBJECTS WITH HCV SUBTYPE 3B INFECTION
- M-260 INFORMATION ADDED TO THE LABELING DESCRIBING A RANDOMIZED, OPEN-LABEL STUDY THAT EXAMINED THE CONCOMITANT USE OF DIMETHYL FUMARATE AND SEVERAL NON-LIVE VACCINES IN ADULTS 27-55 YEARS OF AGE WITH RELAPSING FORMS OF MULTIPLE SCLEROSIS
- M-261 ADDITIONAL INFORMATION ADDED TO THE LABELING REGARDING THE USE IN PATIENTS ON CHRONIC HEMODIALYSIS
- M-262 REVISIONS TO THE USE IN SPECIFIC POPULATIONS SECTION OF THE PACKAGE INSERT TO INCLUDE THE RESULT OF STUDY P146 TO FULFILL THE REQUIREMENTS OF PMR 3003-4
- M-263 REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY ICL670E2419 (THETIS TRIAL) TO SUPPORT PMR 3342-2 AND 3342-3
- M-264 INFORMATION ADDED TO THE LABELING DESCRIBING A PHASE 2, MULTICENTER, OPEN-LABEL

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	STUDY TO EVALUATE THE SAFETY/EFFICACY OF SOFOSBUVIR/VELPATASVIR IN SUBJECTS WITH CHRONIC HCV INFECTION WHO HAVE RECEIVED A LIVER TRANSPLANT
M-265	REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY M15-656 (VIALE-A) AND M16-043 (VIALE-C) TO SUPPORT PMR 3545-1 AND PMR 3545-2
M-266	INFORMATION ADDED TO THE LABELING TO DESCRIBE STUDY GS-US-320-4035 IN VIROLOGICALLY SUPPRESSED ADULTS W/ CHRONIC HEP B INFECTION WHO SWITCHED FROM TENOFOVIR DISOPROXIL FUMARATE TO TENOFOVIR ALFAENAMIDE
M-267	INFORMATION ADDED TO THE LABELING REGARDING THE RESULT OF STUDY LUAA21004-402
M-268	ADDITION OF INFORMATION TO THE LABEL REGARDING A CLEAR PRODUCT PRESENTATION AND 26 ML VOLUME PRODUCTS
M-269	REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY 309/KEYNOTE-775 TO SUPPORT PMR 3696-1 AND 3700-1
M-270	INFORMATION ADDED TO CLINICAL PHARMACOLOGY SECTION

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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
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.

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- 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 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

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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, WORSENER 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.
- 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

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- 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
- 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

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- 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 THERAPY
- ODE-115 TREATMENT OF PATIENTS WITH METASTATIC, SQUAMOUS, NON-SMALL CELL LUNG CANCER PROGRESSED 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

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- 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 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

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- 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+) 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

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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) 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

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- 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 PARANGLIOMA 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 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

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- 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
- ODE-225 INDICATED FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN COMBINATION WITH CHEMOTHERAPY
- ODE-226 MAINTENANCE TREATMENT OF ADULTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC BRCA-MUTATED ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY
- ODE-227 INDICATED FOR TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC) WHO HAVE BEEN PREVIOUSLY TREATED WITH SORAFENIB
- ODE-228 INDICATED FOR THE TREATMENT OF FASCIOLIASIS IN PATIENTS 6 YEARS OF AGE AND OLDER
- ODE-229 TREATMENT OF ADULT PATIENTS WITH METASTATIC GASTRIC OR GASTROESOPHAGEAL JUNCTION ADENOCARCINOMA PREVIOUSLY TREATED WITH AT LEAST TWO PRIOR LINES OF CHEMOTHERAPY, AND IF APPROPRIATE, HER2/NEU-TARGETED THERAPY
- ODE-230 FIRST-LINE TREATMENT OF METASTATIC NONSMALL CELL LUNG CANCER WHOSE TUMORS HAVE NON-RESISTANT EPIDERMAL GROWTH FACTOR MUTATIONS OTHER THAN EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- ODE-231 INDICATED FOR THE TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN PEDIATRIC PATIENTS 7 YEARS OF AGE AND OLDER WITH NARCOLEPSY
- ODE-232 TREATMENT OF PEDIATRIC PATIENTS 12 YEARS AND OLDER OR WEIGHING AT LEAST 45 KG WITH CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1,2,3,4,5 OR 6 INFECTION WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS (CHILD-PUGH A)
- ODE-233 TREATMENT OF PEDIATRIC PATIENTS 12 YEARS AND OLDER OR WEIGHING AT LEAST 45 KG WITH HCV GENOTYPE 1 INFECTION, WHO PREVIOUSLY HAVE BEEN TREATED WITH A REGIMEN

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- CONTAINING AN HCV NS5A INHIBITOR OR AN NS3/4A PROTEASE INHIBITOR (PI), BUT NOT BOTH
- ODE-234 INDICATED FOR THE TREATMENT OF STABLE SYMPTOMATIC HEART FAILURE DUE TO DILATED CARDIOMYOPATHY (DCM) IN PEDIATRIC PATIENTS AGED 6 MONTHS AND OLDER, WHO ARE IN SINUS RHYTHM WITH AN ELEVATED HEART RATE
- ODE-235 INDICATED FOR THE TREATMENT OF ACUTE HERPETIC KERATITIS (DENDRITIC ULCERS) IN PATIENTS WITH HERPES SIMPLEX (HSV-1 AND HSV-2) VIRUS
- ODE-236 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 MONTHS TO LESS THAN 12 MONTHS WHO HAVE ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- ODE-237 TREATMENT OF THE CARDIOMYOPATHY OF WILD TYPE OR HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS (ATTR-CM) IN ADULTS TO REDUCE CARDIOVASCULAR MORTALITY AND CARDIOVASCULAR-RELATED HOSPITALIZATION
- ODE-238 TREATMENT OF STEROID-REFRACTORY ACUTE GRAFT-VERSUS-HOST DISEASE (GVHD) IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER
- ODE-239 TREATMENT OF PREVIOUSLY UNTREATED ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- ODE-240 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH SHORT BOWEL SYNDROME (SBS) WHO ARE DEPENDENT ON PARENTERAL SUPPORT
- ODE-241 INDICATED IN COMBINATION WITH A RITUXIMAB PRODUCT FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED FOLLICULAR LYMPHOMA (FL)
- ODE-242 TX OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 MUTATION AS DETECTED BY AN FDA-APPROVED TEST IN ADULT PTS WHO ARE >=75 YRS OLD OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- ODE-243 ACUTE TREATMENT OF INTERMITTENT, STEREOTYPIC EPISODES OF FREQUENT SEIZURE ACTIVITY (I.E., SEIZURE CLUSTERS, ACUTE REPETITIVE SEIZURES) THAT ARE DISTINCT FROM A PATIENT'S USUAL SEIZURE PATTERN IN PATIENTS WITH EPILEPSY 12 YEARS OF AGE AND OLDER
- ODE-244 TREATMENT OF LAMBERT-EATON MYASTHENIC SYNDROME (LEMS) IN PATIENTS 6 TO LESS THAN 17 YEARS OF AGE
- ODE-245 INDICATED IN COMBINATION WITH A RITUXIMAB PRODUCT FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED MARGINAL ZONE LYMPHOMA (MZL)
- ODE-246 TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC IMMUNE THROMBOCYTOPENIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT
- ODE-247 TX OF PTS W/ CYSTIC FIBROSIS (CF) AGE 6 TO <12 YRS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR W/ AT LEAST 1 MUTATION IN CF TRANSMEMBRANE CONDUCTANCE REGULATORY GENE RESPONSIVE TO TEZACAFTOR/IVACAFTOR BASED ON IN VITRO DATA AND/OR CLINICAL EVIDENCE
- ODE-248 TREATMENT OF ADULT PATIENTS WITH ORAL ULCERS ASSOCIATED WITH BEHCETS DISEASE
- ODE-249 TREATMENT OF PATIENTS WITH CYSTIC FIBROSIS (CF) AGE 6 YEARS TO LESS THAN 12 YEARS WHO ARE HOMOZYGOUS FOR F508DEL MUTATION OR WHO HAVE AT LEAST ONE MUTATION IN THE CF TRANSMEMBRANE CONDUCTANCE REGULATOR GENE THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR
- ODE-250 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH SYMPTOMATIC TENOSYNOVIAL GIANT CELL TUMOR (TGCT) ASSOCIATED WITH SEVERE MORBIDITY OR FUNCTIONAL LIMITATIONS AND NOT AMENABLE TO IMPROVEMENT WITH SURGERY
- ODE-251 INDICATED AS PART OF COMBINATION THERAPY IN THE TREATMENT OF PEDIATRIC PATIENTS (12 TO LESS THAN 18 YEARS OF AGE AND WEIGHING AT LEAST 30 KG) WITH PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS
- ODE-252 TREATMENT OF DUCHENE MUSCULAR DYSTROPHY IN PATIENTS 2 YEARS OF AGE TO LESS THAN 5 YEARS OF AGE
- ODE-253 INDICATED AS PART OF A COMBINATION REGIMEN WITH BEDAQUILINE AND LINEZOLID FOR THE TREATMENT OF ADULTS WITH PULMONARY EXTENSIVELY DRUG RESISTANT (XDR) OR TREATMENT-INTOLERANT OR NONRESPONSIVE MULTIDRUG-RESISTANT (MDR) TUBERCULOSIS (TB)
- ODE-254 INDICATED TO IMPROVE WAKEFULNESS IN ADULT PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY
- ODE-255 INDICATED FOR THE TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS (EDS) IN ADULT PATIENTS WITH NARCOLEPSY

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- ODE-256 FOR HIV-1 INFECTION IN PEDIATRIC PTS AT LEAST 25 KG W/ NO ANTIRETROVIRAL (ARV) TX HX OR TO REPLACE CURRENT ARV REGIMEN FOR VIROLOGICALLY-SUPPRESSED ON STABLE ARV W/ NO HX TX FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED W/ RESISTANCE TO BIC, FTC, OR TAF
- ODE-257 IN COMBO W/ DEXAMETHASONE FOR ADULTS W/ RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO RECEIVED AT LEAST 4 PRIOR THERAPIES AND REFRACTORY TO AT LEAST 2 PROTEASOME INHIBITORS, AT LEAST 2 IMMUNOMODULATORY AGENTS, AND AN ANTI-CD38 MONOCLONAL ANTIBODY
- ODE-258 FOR THE TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 2 OR 3 INFECTION IN PEDIATRIC PATIENTS BETWEEN 3 YEARS OF AGE AND 12 YEARS OF AGE OR WEIGHING 35 KG WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-259 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH INTERMEDIATE-2 OR HIGH-RISK PRIMARY OR SECONDARY (POST-POLYCYTHEMIA VERA OR POST-ESSENTIAL THROMBOCYTHEMIA) MYELOFIBROSIS (MF)
- ODE-260 INDICATED TO INCREASE SYSTEMIC EXPOSURE OF ATAZANAVIR IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS IN THE TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS WEIGHING AT LEAST 35 KG
- ODE-261 INDICATED TO SLOW THE RATE OF DECLINE IN PULMONARY FUNCTION IN PATIENTS WITH SYSTEMIC SCLEROSIS-ASSOCIATED INTERSTITIAL LUNG DISEASE (SSC-ILD)
- ODE-262 TREATMENT OF PEDIATRIC PATIENTS BETWEEN 3 YEARS OF AGE AND 12 YEARS OF AGE OR WEIGHING 35 KG WITH CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 4, 5, OR 6 INFECTION WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS
- ODE-263 TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER WITH CHRONIC HCV GENOTYPE 1 INFECTION WITH DECOMPENSATED CIRRHOSIS, FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-264 TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER WITH CHRONIC HCV GENOTYPE 1 OR 4 INFECTION WHO ARE LIVER TRANSPLANT RECIPIENTS WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS, FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-265 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS ARE ROS1-POSITIVE
- ODE-266 ADULT & PED >=12YRS OLD W/ SOLID TUMORS THAT HAVE NTRK W/O KNOWN ACQUIRED RESISTANCE MUTATION, ARE METASTATIC OR WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN SEVERE MORBIDITY & HAVE EITHER PROGRESSED FOLLOWING TX OR HAVE NO SATISFACTORY ALTERNATIVE TX
- ODE-267 INDICATED IN COMBINATION WITH HIGH FLUID INTAKE, ALKALI, AND DIET MODIFICATION, FOR THE PREVENTION OF CYSTINE STONE FORMATION IN PEDIATRIC PATIENTS 20KG TO 9 YEARS OF AGE W/SEVERE HOMOZYGOUS CYSTINURIA, WHO ARE NOT RESPONSIVE TO THESE MEASURES ALONE
- ODE-268 INDICATED FOR TREATMENT OF PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- ODE-269 PROPHYLAXIS OF ORGAN REJECTION IN PEDIATRIC PATIENTS RECEIVING ALLOGENEIC KIDNEY TRANSPLANT, LIVER TRANSPLANTS, AND HEART TRANSPLANT, IN COMBINATION WITH OTHER IMMUNOSUPPRESSANTS
- ODE-270 INDICATED TO INCREASE PAIN FREE LIGHT EXPOSURE IN ADULT PATIENTS WITH A HISTORY OF PHOTOTOXIC REACTIONS FROM ERYTHROPOIETIC PROTOPORPHYRIA (EPP)
- ODE-271 INDICATED IN COMBINATION WITH OTHER ANTI-MYELOMA PRODUCTS FOR THE TREATMENT OF ADULTS WITH MULTIPLE MYELOMA (MM)
- ODE-272 INDICATED FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) TO DELAY DISEASE PROGRESSION
- ODE-273 INDICATED FOR THE TREATMENT OF ADULTS WITH ACUTE HEPATIC PORPHYRIA (AHP)
- ODE-274 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- ODE-275 INDICATED FOR THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE
- ODE-276 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA (MCL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-277 TX OF ADULTS W/ ADV OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER TREATED W/ >=3 PRIOR CHEMO REGIMENS & CANCER ASSOCIATED W/ HRD+ STATUS DEFINED

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BY A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION

- ODE-278 TX OF ADULTS W/ ADV OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER TREATED W/ >=3 PRIOR CHEMO REGIMENS & CANCER ASSOCIATED W/ HRD+ STATUS DEFINED BY GENOMIC INSTABILITY & PROGRESSED >6 MONTHS AFTER RESPONSE TO THE LAST PLATINUM-BASED CHEMOTHERAPY
- ODE-279 INDICATED FOR THE ACUTE TX OF INTERMITTENT, STEREOTYPIC EPISODES OF FREQUENT SEIZURE ACTIVITY (I.E. SEIZURE CLUSTERS, ACUTE REPETITIVE SEIZURES) THAT ARE DISTINCT FROM A PATIENT'S USUAL SEIZURE PATTERN IN PATIENTS WITH EPILEPSY 6 YEARS OF AGE AND OLDER
- ODE-280 INDICATED FOR THE TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 53 SKIPPING
- ODE-281 INDICATED FOR THE TREATMENT OF SICKLE CELL DISEASE (SCD) IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- ODE-282 INDICATED TO SELECTIVELY STAIN THE INTERNAL LIMITING MEMBRANE (ILM)
- ODE-283 MAINTENANCE TX OF ADULTS W/ DELETERIOUS OR SUSPECTED DELETERIOUS GBRCA METASTATIC PANCREATIC ADENOCARCINOMA WHOSE DZ HAS NOT PROGRESSED ON >=16WKS OF 1ST LINE PLATINUM BASED CHEMO REGIMEN. SELECT PTS FOR THERAPY BASED ON APPROVED COMPANION DIAGNOSTIC
- ODE-284 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS, FOR THE TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS UNDER 12 YEARS OF AGE WEIGHING AT LEAST 35KG
- ODE-285 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS OTHER THAN PROTEASE INHIBITORS THAT REQUIRE A CYP3A INHIBITOR, FOR THE TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS UNDER 12 YEARS OF AGE WEIGHING AT LEAST 25KG AND LESS THAN 35KG
- ODE-286 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- ODE-287 TREATMENT OF ADULTS WITH NEWLY DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC)
- ODE-288 INDICATED FOR THE TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH NEUROFIBROMATOSIS TYPE 1 (NF1) WHO HAVE SYMPTOMATIC, INOPERABLE PLEXIFORM NEUROFIBROMAS (PN)
- ODE-289 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH LOW-GRADE UPPER TRACT UROTHELIAL CANCER (LG-UTUC)
- ODE-290 INDICATED FOR THE INITIAL TREATMENT OF SEVERE MALARIA IN ADULT AND PEDIATRIC PATIENTS TO ALWAYS BE FOLLOWED BY A COMPLETE TREATMENT COURSE OF AN APPROPRIATE ORAL ANTIMALARIAL REGIMEN
- ODE-291 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE A MUTATION THAT LEADS TO MESENCHYMAL-EPITHELIAL TRANSITION (MET) EXON 14 SKIPPING AS DETECTED BY AN FDA-APPROVED TEST
- ODE-292 INDICATED FOR THE TREATMENT OF ADULTS WITH PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FIBROBLAST GROWTH FACTOR RECEPTOR 2 (FGFR2) FUSION OR OTHER REARRANGEMENT AS DETECTED BY AN FDA-APPROVED TEST
- ODE-293 TX OF PED PTS 6 YRS OF AGE & OLDER OR WEIGHING AT LEAST 17 KG WITH CHRONIC HEPATITIS C VIRUS GENOTYPE 1, 2, 3, 4, 5, OR 6 INFECTION: WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS; OR WITH DECOMPENSATED CIRRHOSIS FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-294 PROPHYLAXIS OF ORGAN REJECTION IN PEDIATRIC PATIENTS RECEIVING ALLOGENEIC KIDNEY OR HEART TRANSPLANTS, IN COMBINATION WITH OTHER IMMUNOSUPPRESSANTS
- ODE-295 INDICATED FOR THE MAINTENANCE TREATMENT OF ADULT PATIENTS WITH ADVANCED EPITHELIAL OVARIAN CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY
- ODE-296 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH AIDS-RELATED KAPOSI SARCOMA (KS) AFTER FAILURE OF HIGHLY ACTIVE ANTIRETROVIRAL THERAPY (HAART)
- ODE-297 FOR THE TREATMENT OF KAPOSI SARCOMA (KS) IN ADULT PATIENTS WHO ARE HIV-NEGATIVE
- ODE-298 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH ADVANCED GASTROINTESTINAL STROMAL TUMOR (GIST) WHO HAVE RECEIVED PRIOR TREATMENT WITH 3 OR MORE KINASE INHIBITORS, INCLUDING IMATINIB
- ODE-299 INDICATED FOR THE TREATMENT OF ADULTS AND PEDIATRIC PATIENTS AGED 16 YEARS AND OLDER WITH METASTATIC OR LOCALLY ADVANCED EPITHELIOID SARCOMA NOT ELIGIBLE FOR

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COMPLETE RESECTION

- ODE-300 FOR THE TREATMENT OF ADULT PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)- POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA- APPROVED TEST, NOT INCLUDING PATIENTS WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- ODE-301 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC RET FUSION- POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC)
- ODE-302 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET-MUTANT MEDULLARY THYROID CANCER (MTC) WHO REQUIRE SYSTEMIC THERAPY
- ODE-303 ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY (IF RADIOACTIVE IODINE IS APPROPRIATE)
- ODE-304 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC SMALL CELL LUNG CANCER (SCLC) WITH DISEASE PROGRESSION ON OR AFTER PLATINUM-BASED CHEMOTHERAPY
- ODE-305 TREATMENT OF INVASIVE ASPERGILLOSIS
- ODE-306 W/ BEVACIZUMAB FOR MAINT TX OF ADULTS W/ ADV EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CA IN COMPLETE OR PARTIAL RESPONSE TO 1ST LINE PT BASED CHEMO & WHOSE CA IS ASSOC W/ HOMOLOGOUS RECOMB DEF + STATUS DEFINED BY GENOMIC INSTABILITY
- ODE-307 INDICATED AS PART OF COMBINATION THERAPY IN THE TREATMENT OF PEDIATRIC PATIENTS 5 YEARS AND OLDER TO LESS THAN 12 YEARS OF AGE AND WEIGHING AT LEAST 15 KG WITH PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS (MDR-TB)
- ODE-308 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) TO IMPROVE EXERCISE CAPACITY
- ODE-309 INDICATED FOR USE IN COMBINATION WITH TRASTUZUMAB AND CAPECITABINE FOR TREATMENT OF ADULT PATIENTS WITH METASTATIC HER2-POSITIVE BREAST CANCER AND BRAIN METASTASES, WHO HAVE RECEIVED ONE OR MORE PRIOR ANTI-HER2-BASED REGIMENS IN THE METASTATIC SETTING
- ODE-310 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL), NOT OTHERWISE SPECIFIED, INCLUDING DLBCL ARISING FROM FOLLICULAR LYMPHOMA, AFTER AT LEAST 2 LINES OF SYSTEMIC THERAPY
- ODE-311 INDICATED AS A SOURCE OF CALORIES AND FATTY ACIDS FOR THE TREATMENT OF PEDIATRIC AND ADULT PATIENTS WITH MOLECULARLY CONFIRMED LONG-CHAIN FATTY ACID OXIDATION DISORDERS (LC-FAOD)
- ODE-312 INDICATED FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME IN PATIENTS 2 YEARS OF AGE AND OLDER
- ODE-313 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH SOLID TUMORS THAT HAVE A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION WITHOUT A KNOWN ACQUIRED RESISTENACE MUTATION, ARE METASTATIC OR WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN SEVERE MORBIDITY, AND HAVE EITHER PROGRESSED FOLLOWING TREATMENT OR HAVE NO SATISFACTORY ALTERNATIVE THERAPY
- ODE-314 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA WHOSE TUMORS ARE POSITIVE FOR AN EZH2 MUTATION AS DETECTED BY AN FDA-APPROVED TEST AND WHO HAVE RECEIVED AT LEAST 2 PRIOR SYSTEMIC THERAPIES, AND FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA WHO HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
- ODE-315 FOR CLEANSING OF THE COLON AS A PREPARATION FOR COLONOSCOPY IN PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- ODE-316 TREATMENT OF ADULT PATIENTS WITH MYELODYSPLASTIC SYNDROMES (MDS), INCLUDING PREVIOUSLY TREATED AND UNTREATED, DE NOVO AND SECONDARY MDS WITH THE FOLLOWING FRENCH-AMERICAN-BRITISH SUBTYPES (REFRACTORY ANEMIA, REFRACTORY ANEMIA WITH RINGED SIDEROBLASTS, REFRACTORY ANEMIA WITH EXCESS BLASTS, AND CHRONIC MYELOMONOCYTIC LEUKEMIA [CMML]) AND INTERMEDIATE-1, INTERMEDIATE-2, AND HIGH-RISK INTERNATIONAL PROGNOSTIC SCORING SYSTEM GROUPS.
- ODE-317 FOR USE WITH POSITRON EMISSION TOMOGRAPHY (PET) FOR LOCALIZATION OF SOMATOSTATIN RECEPTOR POSITIVE NEUROENDOCRINE TUMORS (NETS) IN ADULT PATIENTS
- ODE-318 TREATMENT OF ADULT PATIENTS WITH METASTATIC RET FUSION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA APPROVED TEST

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- ODE-319 INDICATED IN PEDIATRIC PATIENTS (BIRTH TO LESS THAN 18 YEARS OF AGE AND WEIGHING AT LEAST 2.5 KG) FOR THE TREATMENT OF CHAGAS DISEASE (AMERICAN TRYPANOSOMIASIS) CAUSED BY TRYPANOSOMA CRUZI
- ODE-320 INDICATED FOR CONTINUED TREATMENT OF ADULT PATIENTS WITH ACUTE MYELOID LEUKEMIA WHO ACHIEVED FIRST COMPLETE REMISSION (CR) OR COMPLETE REMISSION WITH INCOMPLETE BLOOD COUNT RECOVERY (CRI) FOLLOWING INTENSIVE INDUCTION CHEMOTHERAPY AND ARE NOT ABLE TO COMPLETE INTENSIVE CURATIVE THERAPY
- ODE-321 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 4 MONTHS TO LESS THAN 6 MONTHS WHO HAVE ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- ODE-322 TREATMENT OF NARCOLEPSY
- ODE-323 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE A MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA
- ODE-324 TREATMENT OF HUTCHINSON-GILFORD PROGERIA SYNDROME (HGPS) AND PROGEROID LAMINOPATHIES
- ODE-325 TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) HARBORING MESENCHYMAL-EPITHELIAL TRANSITION (MET) EXON 14 SKIPPING ALTERATIONS
- ODE-326 TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME (LGS) OR DRAVET SYNDROME (DS) IN PATIENTS BETWEEN 1 AND 2 YEARS OF AGE
- ODE-327 ADD-ON MAINTENANCE THERAPY TO IMPROVE PULMONARY FUNCTION IN ADULT PATIENTS 18 YEARS OF AGE AND OLDER WITH CYSTIC FIBROSIS AND WHO HAVE PASSED THE BRONCHITOL TOLERANCE TEST
- ODE-328 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER AND YOUNG ADULTS WITH RELAPSED OR REFRACTORY, SYSTEMIC ANAPLASTIC LARGE CELL LYMPHOMA (ALCL) THAT IS ALK-POSITIVE
- ODE-329 FOR THE TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME (SMS) IN PEDIATRIC PATIENTS 3 TO 15 YEARS OF AGE
- ODE-330 THE TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME (SMS) IN PATIENTS 16 YEARS OF AGE AND OLDER
- ODE-331 TREATMENT OF CATAPLEXY IN ADULT PATIENTS WITH NACROLEPSY
- ODE-332 TREATMENT OF SEIZURES ASSOCIATED WITH TUBEROUS SCLEROSIS COMPLEX (TSC) IN PATIENTS 1 YEAR OF AGE AND OLDER
- ODE-333 PROPHYLAXIS TO PREVENT ATTACKS OF HEREDITARY ANGIOEDEMA (HAE) IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- ODE-334 TREATMENT OF SPINAL MUSCULAR ATROPHY (SMA) IN PATIENTS 2 MONTHS OF AGE AND OLDER
- ODE-335 FOR TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE AT LEAST ONE OF THE ADDITIONAL MUTATIONS IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE THAT HAVE BEEN IDENTIFIED AS RESPONSIVE TO TEZACAFTOR/IVACAFTOR BASED ON IN VITRO DATA AND IDENTIFIED IN THE APPROVAL ON DECEMBER 21, 2020
- ODE-336 INDICATED FOR CHRONIC WEIGHT MANAGEMENT IN ADULT AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WITH OBESITY DUE TO PROOPIOMELANOCORTIN (POMC), PROPROTEIN CONVERTASE SUBTILISIN/KEXIN TYPE 1 (PCSK1), OR LEPTIN RECEPTOR (LEPR) DEFICIENCY CONFIRMED BY GENETIC TESTING DEMONSTRATING VARIANTS IN POMC, PCSK1, OR LEPR GENES THAT ARE INTERPRETED AS PATHOGENIC, LIKELY PATHOGENIC, OR OF UNCERTAIN SIGNIFICANCE (VUS)
- ODE-337 FOR ADJUVANT THERAPY AFTER TUMOR RESECTION IN ADULT PATIENTS WITH 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-338 FOR THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 4 MONTHS AND OLDER WHO HAVE ONE OF THE ADDITIONAL MUTATIONS IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE THAT HAVE BEEN IDENTIFIED AS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON IN VITRO DATA AND IDENTIFIED IN THE APPROVAL ON DECEMBER 21, 2020
- ODE-339 TREATMENT OF PRIMARY HYPEROXALURIA TYPE 1 (PH1) TO LOWER URINARY OXALATE LEVELS IN PEDIATRIC AND ADULT PATIENTS
- ODE-340 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET-MUTANT MEDULLARY THYROID CANCER (MTC) WHO REQUIRE

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

SYSTEMIC THERAPY

- ODE-341 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY (IF RADIOACTIVE IODINE IS APPROPRIATE)
- ODE-342 INDICATED TO REDUCE THE RISK OF MORTALITY IN PATIENTS WITH MOLYBDENUM COFACTOR DEFICIENCY (MOCD) TYPE A
- ODE-343 FOR TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA (MZL) WHO HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED REGIMEN
- ODE-344 FOR TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL) WHO HAVE RECEIVED AT LEAST THREE PRIOR LINES OF SYSTEMIC THERAPY
- ODE-345 IN PEDIATRIC AND ADULT PATIENTS AS ADJUNCTIVE THERAPY TO STANDARD OF CARE FOR THE TREATMENT OF ACUTE HYPERAMMONEMIA DUE TO PROPIONIC ACIDEMIA (PA) OR METHYLMALONIC ACIDEMIA (MMA)
- ODE-346 FOR THE TREATMENT OF ADULT PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY, EXCLUDING ADULT PATIENTS COVERED BY XPOVIO'S PREVIOUS INDICATION FOR MULTIPLE MYELOMA APPROVED ON JULY 3, 2019
- ODE-347 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 45 SKIPPING
- ODE-348 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST FOUR PRIOR LINES OF THERAPY AND WHOSE DISEASE IS REFRACTORY TO AT LEAST ONE PROTEASOME INHIBITOR, ONE IMMUNOMODULATORY AGENT, AND ONE CD-38 DIRECTED MONOCLONAL ANTIBODY
- ODE-349 TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS ARE ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA-APPROVED TEST, EXCLUDING PATIENTS WHOSE DISEASE HAS PROGRESSED ON CRIZOTINIB AND AT LEAST ONE OTHER ALK INHIBITOR FOR METASTATIC DISEASE; OR ALECTINIB OR CERITINIB AS THE FIRST ALK INHIBITOR THERAPY FOR METASTATIC DISEASE
- ODE-350 TREATMENT OF NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC) IN PEDIATRIC PATIENTS AGES 1 YEAR AND OLDER
- ODE-351 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH)
- ODE-352 TREATMENT OF ADULT PATIENTS WITH KRAS G12C-MUTATED LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), AS DETERMINED BY AN FDA-APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR SYSTEMIC THERAPY
- ODE-353 FOR TREATMENT OF ADULTS WITH PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FIBROBLAST GROWTH FACTOR RECEPTOR 2 (FGFR2) FUSION OR OTHER REARRANGEMENT AS DETECTED BY AN FDA-APPROVED TEST
- ODE-354 TREATMENT OF HUMAN SMALLPOX DISEASE CAUSED BY VARIOLA VIRUS IN ADULT AND PEDIATRIC PATIENTS, INCLUDING NEONATES
- ODE-355 FOR THE TREATMENT OF INVASIVE ASPERGILLOSIS IN ADULTS AND PEDIATRIC PATIENTS 13 YEARS OF AGE AND OLDER
- ODE-356 FOR THE TREATMENT OF ADULT PATIENTS WITH ADVANCED SYSTEMIC MASTOCYTOSIS (ADVSM). ADVSM INCLUDES PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH AN ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), AND MAST CELL LEUKEMIA (MCL)
- ODE-357 FOR THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGED 6 THROUGH 11 YEARS OLD WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA
- ODE-358 FOR THE TREATMENT OF VERNAL KERATOCONJUNCTIVITIS (VKC) IN CHILDREN AND ADULTS
- ODE-359 FOR THE TREATMENT OF BOTH THE FIRST-STAGE (HEMOLYMPHATIC) AND SECOND-STAGE (MENINGOENCEPHALITIC) HUMAN AFRICAN TRY PANOSOMIASIS (HAT) DUE TO TRY PANOSOMA BRUCEI GAMBIENSE IN PATIENTS 6 YEARS OF AGE AND OLDER AND WEIGHING AT LEAST 20 KG
- ODE-360 FOR PROPHYLAXIS OF ORGAN REJECTION IN ADULT AND PEDIATRIC PATIENTS RECEIVING ALLOGENEIC LUNG TRANSPLANT
- ODE-361 INDICATED FOR THE TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN PATIENTS 7 YEARS OF AGE AND OLDER WITH NARCOLEPSY

PATENT AND EXCLUSIVITY TERMS

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ODE-362	TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER WITH CHRONIC GRAFT-VERSUS-HOST DISEASE (CHRONIC GVHD) AFTER FAILURE OF AT LEAST TWO PRIOR LINES OF SYSTEMIC THERAPY
ODE-363	TREATMENT OF PRURITUS IN PATIENTS 3 MONTHS OF AGE AND OLDER WITH PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)
ODE-364	TREATMENT OF ADULT PATIENTS WITH VON HIPPEL-LINDAU (VHL) DISEASE WHO REQUIRE THERAPY FOR ASSOCIATED RENAL CELL CARCINOMA (RCC), CENTRAL NERVOUS SYSTEM (CNS) HEMANGIOBLASTOMAS, OR PANCREATIC NEUROENDOCRINE TUMORS (PNET), NOT REQUIRING IMMEDIATE SURGERY
ODE-365	TREATMENT OF ADULT PATIENTS WITH VON HIPPEL-LINDAU (VHL) DISEASE WHO REQUIRE THERAPY FOR ASSOCIATED RENAL CELL CARCINOMA (RCC), CENTRAL NERVOUS SYSTEM (CNS) HEMANGIOBLASTOMAS, OR PANCREATIC NEUROENDOCRINE TUMORS (PNET), NOT REQUIRING IMMEDIATE SURGERY
ODE-366	INDICATED FOR THE TREATMENT OF ADULTS WITH UNRESECTABLE OR METASTATIC GIST HARBORING A PLATELET-DERIVED GROWTH FACTOR RECEPTOR ALPHA (PDGFRA) EXON 18 MUTATION, INCLUDING PDGFRA D842V MUTATIONS
ODE-367	PEDIATRIC PATIENTS AGED 6 MONTHS AND OLDER FOR THE TREATMENT OF C. DIFFICILE-ASSOCIATED DIARRHEA (CDAD)
ODE-368	TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH AN ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
ODE-369	THE TREATMENT OF IDIOPATHIC HYPERSOMNIA (IH) IN ADULTS
ODE-370	TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA (MZL) WHO HAVE RECEIVED AT LEAST ONE ANTI-CD20-BASED REGIMEN
ODE-371	TREATMENT OF ADULT PATIENTS WITH WALDENSTRM'S MACROGLOBULINEMIA (WM)
ODE-372	FOR TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE TO LESS THAN 12 YEARS OF AGE WEIGHING LESS THAN 45 KG WITH CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5 OR 6 INFECTION WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS (CHILD-PUGH A); AND TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE TO LESS THAN 12 YEARS OF AGE WEIGHING LESS THAN 45 KG WITH HCV GENOTYPE 1 INFECTION, WHO PREVIOUSLY HAVE BEEN TREATED WITH A REGIMEN CONTAINING AN HCV NS5A INHIBITOR OR AN NS3/4A PROTEASE INHIBITOR (PI), BUT NOT BOTH
ODE-373	TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR TWO LINES OF SYSTEMIC THERAPY IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER
ODE-374	TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 20 INSERTION MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, WHOSE DISEASE HAS PROGRESSED ON OR AFTER PLATINUM-BASED CHEMOTHERAPY
ODE-375	THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH LOCALLY ADVANCED OR METASTATIC DIFFERENTIATED THYROID CANCER (DTC) THAT HAS PROGRESSED FOLLOWING PRIOR VEGFR-TARGETED THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY OR INELIGIBLE
ODE-376	FOR TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE TO LESS THAN 6 YEARS OF AGE WEIGHING LESS THAN 17 KG WITH CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6 INFECTION: WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS; OR WITH DECOMPENSATED CIRRHOSIS FOR USE IN COMBINATION WITH RIBAVIRIN

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
U-6	METHOD OF PRODUCING SYMPATHOMIMETIC EFFECTS
U-7	INCREASING CARDIAC CONTRACTILITY
U-8	ACUTE MYOCARDIAL INFARCTION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

U-46 TREATMENT OF PANIC DISORDER

U-47 STIMULATION OF THE RELEASE OF GROWTH HORMONE

U-48 ANALGESIA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

U-125 TREATMENT NEUROGENERATIVE DISEASES

U-126 TREATMENT OF GASTRITIS

U-127 METHOD OF PRODUCING NEUROMUSCULAR BLOCKADE

U-128 METHOD FOR TREATMENT OF TUMORS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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-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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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....

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

PSYCHOLES, 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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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
- 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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
- 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 AMYCOPLASMA 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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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 RESONANCE 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 PERIODONTAL SCALING AND ROOT PLANNING PROCEDURES BY APPLICATION OF AN EUTECTIC MIXTURE OF LOCAL ANESTHETICS TO PERIODONTAL 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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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 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)

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 HYPONATREMIA
- 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
- 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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 SEQUESTRANT FOR LOWERING CHOLESTEROL

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 ARIPIPRAZOLE 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)

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 EROSIVE ESOPHAGITIS

U-859 EROSIVE ESOPHAGITIS, HYPERSECRETORY CONDITIONS INCLUDING ZOLLINGER-ELLISON SYNDROME, MAINTENANCE OF HEALING OF EROSIVE 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 EROSIVE 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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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, 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 TROSPIMUM 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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 PSEUDOBULBAR 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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

U-1143 REDUCTION IN RISK OF RECURRENT NONFATAL MYOCARDIAL INFARCTION BY DOSING ONCE PER DAY IN THE EVENING OR A T 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

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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, GLIMEPIRIDE & 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 OSTEODYSTROPHY, 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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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)

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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
- 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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
- 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 RYTHM 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 ARIPIPRAZOLE 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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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 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.

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 ARIPIRAZOLE IN EXTENDED RELEASE INJECTABLE SUSPENSION IN TREATING ACUTE EPISODES OF SCHIZOPHRENIA
- U-1634 TREATMENT OF BRCA MUTATED OVARIAN CANCER USING PARP INHIBITOR
- U-1635 USE OF RITONAVIR AS A POTENT CYP3A INHIBITOR TO INCREASE PLASMA DRUG CONCENTRATION OF PARITAPREVR 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 PARITAPREVR, OMBITASVIR, RITONAVIR, AND DASABUVIR WITH RIBAVIRIN.

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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
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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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
- 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 Eesomeprazole magnesium AS CLAIMED

U-1784 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN Eesomeprazole magnesium TRIHYDRATE AS CLAIMED

U-1785 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN Eesomeprazole 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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 PARITAPREXIR, 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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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-OME PRAZOLE 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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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. 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)

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 CYP1A2 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 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

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

AND TRICHOPHYTON MENTAGROPHYTES

- U-1970 TREATMENT OF ONYCHOMYCHOSIS 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 MYELODYSPLASTIC 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
- 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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
- 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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 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

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PATENT USE

- 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
- 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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
- 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 NITRIC 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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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
- 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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
- 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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
- 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 \geq 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

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

- 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
- 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

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

- 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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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 MYELOYDYSPLASIA-RELATED CHANGES (AML-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 MYELOYDYSPLASIA-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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- 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 GBRAM, 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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 LUMACAF TOR AND IVACAF TOR

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 LUMACAF TOR FORM I AND IVACAF TOR

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 LUMACAF TOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS AND LESS THAN ABOUT 30% CRYSTALLINE IVACAF TOR

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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
- 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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

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

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

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

U-2480 MAINTENANCE TREATMENT OF GBRCA- OR SBRCA-MUTATED ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY

U-2481 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY

U-2482 TREATMENT OF HR-NEGATIVE, HER-2 NEGATIVE, GBRCA-MUTATED METASTATIC BREAST CANCER, WHO HAVE BEEN TREATED WITH CHEMOTHERAPY IN THE NEOADJUVANT, ADJUVANT, OR METASTATIC SETTING

U-2483 TREATMENT OF HR-POSITIVE, HER-2 NEGATIVE, GBRCA-MUTATED METASTATIC BREAST CANCER, WHO HAVE BEEN TREATED WITH CHEMOTHERAPY IN THE NEOADJUVANT, ADJUVANT, OR METASTATIC SETTING, AND WITH ENDOCRINE THERAPY OR ARE INAPPROPRIATE FOR ENDOCRINE THERAPY

U-2484 INTERMITTENT TREATMENT OF OFF EPISODES IN PATIENTS WITH PARKINSON'S DISEASE TREATED WITH CARBIDOPA/LEVODOPA BY INHALATION OF LEVODOPA POWDER PARTICLES

U-2485 INTERMITTENT TREATMENT OF OFF EPISODES IN PATIENTS WITH PARKINSON'S DISEASE TREATED WITH CARBIDOPA/LEVODOPA BY INHALATION OF LEVODOPA POWDER PARTICLES THROUGH A SINGLE BREATH ACTIVATED STEP

U-2486 INTERMITTENT TREATMENT OF OFF EPISODES IN PATIENTS WITH PARKINSON'S DISEASE WITH A POWDER INHALER

U-2487 DEXTENZA IS APPROVED FOR THE TREATMENT OF OCULAR PAIN FOLLOWING OPHTHALMIC SURGERY

U-2488 TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC) WHO HAVE BEEN PREVIOUSLY TREATED WITH SORAFENIB

U-2489 TREATMENT OF MODERATE TO SEVERE OPIOID USE DISORDER

U-2490 TREATMENT OF COMPLICATED URINARY TRACT INFECTION (CUTI) INCLUDING PYELONEPHRITIS CAUSED BY THE FOLLOWING SUSCEPTIBLE MICROORGANISMS: ESCHERICHIA COLI, KLEBSIELLA PNEUMONIA, AND ENTEROBACTER CLOACAE SPECIES COMPLEX

U-2491 A METHOD FOR DELIVERING A COMPOSITION TO A MUCUS MEMBRANE

U-2492 A METHOD FOR DELIVERING A PHARMACEUTICAL AGENT ACROSS A MUCOSAL BARRIER

U-2493 A METHOD FOR TREATING INFLAMMATION AND/OR OTHER DISORDERS IN AN EYE OF A PATIENT

U-2494 INDICATED FOR THE TREATMENT OF VENTRICULAR ARRHYTHMIAS, SUCH AS SUSTAINED VENTRICULAR TACHYCARDIA, THAT IN THE JUDGEMENT OF THE PHYSICIAN ARE LIFE-THREATENING

U-2495 VENTRICULAR FIBRILLATION

U-2496 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY USE OF A PEN INJECTOR WITH A THREADED DRIVE SLEEVE

U-2497 TREATMENT OF DRUG-INDUCED EXTRAPYRAMIDAL REACTION IN ADULT PATIENTS

U-2498 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 10,206,877

U-2499 METHOD OF REDUCING ADVERSE EFFECTS IN PATIENTS SUFFERING FROM EXCESSIVE DAYTIME SLEEPINESS AND/OR CATAPLEXY IN NARCOLEPSY WHO ARE CONCOMITANTLY ADMINISTERED SODIUM OXYBATE AND DIVALPROEX SODIUM

U-2500 USE OF A DELIVERY DEVICE TO DELIVER A BIOEQUIVALENT DOSE OF A NALOXONE COMPOSITION VIA A NEEDLE

U-2501 TREATMENT OF PARTIAL-ONSET SEIZURES

U-2502 TREATMENT OF TREATMENT-RESISTANT DEPRESSION IN ADULT IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT

U-2503 TREATMENT OF ADULTS WITH METASTATIC GASTRIC OR GJA PREVIOUSLY TREATED WITH AT

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

LEAST TWO PRIOR LINES OF CHEMOTHERAPY THAT INCLUDED A FLUOROPYRIMIDINE, A PLATINUM, EITHER A TAXANE OR IRINOTECAN, AND IF APPROPRIATE, HER2/NEU-TARGETED THERAPY

- U-2504 TREATMENT OF HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN POSTMENOPAUSAL WOMEN IN COMBINATION WITH RIBOCICLIB AS INITIAL ENDOCRINE BASED THERAPY OR FOLLOWING DISEASE PROGRESSION ON ENDOCRINE THERAPY
- U-2505 TREATMENT OF PRE/PERIMENOPAUSAL WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-2506 METHOD OF TREATING TESTOSTERONE DEFICIENCY
- U-2507 METHOD OF TREATING ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS (ABSSSI) CAUSED BY DESIGNATED SUSCEPTIBLE BACTERIA
- U-2508 A METHOD OF TREATING BACTERIAL INFECTIONS IN COMPLICATED INTRA-ABDOMINAL INFECTION AND COMPLICATED URINARY TRACT INFECTION, INCLUDING PYELONEPHRITIS, PATIENTS COMPRISING ADMINISTERING A BACTERICIDALLY EFFECTIVE AMOUNT OF AVIBACTAM SODIUM
- U-2509 A METHOD OF TREATING A BACTERIAL INFECTION IN COMPLICATED INTRA-ABDOMINAL INFECTION (CIAI) AND COMPLICATED URINARY TRACT INFECTION (CUTI), INCLUDING PYELONEPHRITIS, PATIENTS COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AVIBACTAM SODIUM
- U-2510 A METHOD FOR CONTRACEPTION COMPRISING THE STEP OF ORAL ADMINISTRATION A DOSAGE OF 20 MG TO 30 MG OF ULIPRISTAL ACETATE TO A WOMAN WITHIN 72 HOURS AND UP TO 120 HOURS AFTER AN UNPROTECTED INTERCOURSE
- U-2511 A METHOD OF TREATING MULTIPLE SCLEROSIS BY ADMINISTERING SIPONIMOD USING A TITRATION SCHEME TO REACH A MAINTENANCE DOSE
- U-2512 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVE AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH AN EFFECTIVE AMOUNT OF TEZACAFTOR AND IVACAFTOR
- U-2513 MAINTENANCE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-2514 MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-2515 PALBOCICLIB FOR HR-POS. HER2-NEG. ADVANCED OR METASTATIC BREAST CANCER IN COMBO WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY IN POSTMENOPAUSAL WOMEN OR MEN, OR WITH FULVESTRANT IN PTS WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY
- U-2516 A METHOD FOR REDUCING SERUM GLUCOSE LEVELS IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2517 A METHOD FOR REDUCING SERUM GLUCOSE LEVELS IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2518 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA THAT HAS SUSCEPTIBLE FGFR3 OR FGFR2 GENETIC ALTERATIONS AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY
- U-2519 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY
- U-2520 TREATING MS WITH ORAL CLADRIBINE ACC. TO THE STEPS (I) INDUCTION PERIOD WITH ABOUT 1.7 MG/KG-3.5 MG/KG CLADRIBINE; (II) CLADRIBINE-FREE PERIOD OF ABOUT 8-10 MONTHS; (III) MAINTENANCE PERIOD WITH ABOUT 1.7 MG/KG CLADRIBINE; (IV) CLADRIBINE-FREE PERIOD
- U-2521 TREATMENT OF MS WITH A TABLET WITH AN ADMIXTURE OF (A) AN AMORPHOUS INCLUSION COMPLEX OF CLADRIBINE AND HYDROXYPROPYL-B-CYCLODEXTRIN AND (B) AMORPHOUS FREE CLADRIBINE AND CYCLODEXTRIN AS A NON-INCLUSION COMPLEX, CLADRIBINE/CYCLODEXTRIN 1:10-1:16 W/W
- U-2522 TREATING RRMS OR SPMS WITH ORAL CLADRIBINE: (I) 2-4 MONTHS INDUCTION WITH 1.7 MG/KG - 3.5 MG/KG CLADRIBINE; (II) CLADRIBINE-FREE PERIOD OF ABOUT 8-10 MONTHS; (III) 2-4 MONTHS MAINTENANCE WITH ABOUT 1.7 MG/KG CLADRIBINE; (IV) CLADRIBINE-FREE PERIOD
- U-2523 TREATMENT OF MS WITH AN ADMIXTURE OF (A) AN AMORPHOUS INCLUSION COMPLEX OF CLADRIBINE (2CDA) AND CYCLODEXTRIN AND (B) AMORPHOUS FREE 2CDA AND CYCLODEXTRIN AS A NON-INCLUSION COMPLEX, FORMULATED AS A SOLID ORAL FORM, W/O SIGN. AMOUNTS

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

OF CRYST. 2CDA

- U-2524 TREATMENT OF THE CARDIOMYOPATHY OF WILD TYPE OR HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS (ATTR-CM)
- U-2525 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER WITH SYMPTOMS OF URINARY FREQUENCY, URGENCY OR URGE INCONTINENCE
- U-2526 ACUTE TREATMENT OF INTERMITTENT, STEREOTYPIC EPISODES OF FREQUENT SEIZURE ACTIVITY (I.E., SEIZURE CLUSTERS, ACUTE REPETITIVE SEIZURES) THAT ARE DISTINCT FROM A PATIENT'S USUAL SEIZURE PATTERN IN PATIENTS WITH EPILEPSY 12 YEARS OF AGE AND OLDER
- U-2527 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 6 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-2528 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 6 MONTHS TO <6 YEARS WHO HAS A R117H MUTATION IN THE CFTR GENE
- U-2529 TREATMENT OF A MODERATE MILD CLINICAL PHENOTYPE OF CF USING IVACAFTOR IN A PATIENT AGE 6 MONTHS TO <6 YEARS WHO HAS ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-2530 TREATMENT OF CF IN A PATIENT AGE 6 MONTHS TO < 6 YEARS WHO HAS ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING A SOLID COMPOSITION COMPRISING AMORPHOUS (LESS THAN ABOUT 30% CRYSTALLINE) IVACAFTOR
- U-2531 TREATMENT OF CF IN A PATIENT AGE 6 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 10272046
- U-2532 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, OR 6 IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER OR WEIGHING AT LEAST 45 KG
- U-2533 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML)
- U-2534 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML)
- U-2535 USE IN COMBINATION WITH METHYLPREDNISOLONE FOR THE TREATMENT OF PATIENTS WITH PROSTATE CANCER
- U-2536 FOR TREATMENT OF STEROID-REFRACTORY ACUTE GRAFT-VERSUS-HOST DISEASE
- U-2537 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- U-2538 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LEUKEMIA (SLL) IN COMBINATION WITH A GA101 ANTIBODY SUCH AS OBINUTUZUMAB FOR ONE OR MORE DOSING PERIODS, WHEREIN THE CLL OR SLL IS A CD20-EXPRESSING CANCER
- U-2539 IN COMBINATION WITH FULVESTRANT FOR TREATMENT OF POSTMENOPAUSAL WOMEN, AND MEN, WITH HR-POSITIVE, HER-2-NEGATIVE, PIK3CA-MUTATED, ADVANCED OR METASTATIC BREAST CANCER
- U-2540 TREATMENT OF HORMONE RECEPTOR POSITIVE ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN
- U-2541 REDUCING THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION (MI), AND STROKE IN A PATIENT RECEIVING 75-100 MG ASPIRIN DAILY WITH A HISTORY OF MI BY ADMINISTERING 60 MG TICAGRELOR TWICE DAILY
- U-2542 REDUCING THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN A PATIENT RECEIVING 75-100 MG ASPIRIN DAILY AND HAVING OR WHO HAD ACUTE CORONARY SYNDROME BY ADMINISTERING 60 MG TICAGRELOR TWICE DAILY
- U-2543 TREATMENT OF SCHIZOPHRENIA WITH CARIPRAZINE
- U-2544 TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER WITH CARIPRAZINE
- U-2545 TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER (BIPOLAR DEPRESSION) WITH CARIPRAZINE
- U-2546 USE FOR THE MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
- U-2547 METHOD OF PROVIDING CONTRACEPTION IN A WOMAN HAVING A BMI OF 25 KG/M2 OR MORE WITH RESULTANT LIMITED BLEEDING EVENTS PER TREATMENT CYCLE

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- U-2548 TO IMPROVE WAKEFULNESS IN ADULT PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY OR OBSTRUCTIVE SLEEP APNEA (OSA)
- U-2549 CONTROL OF SERUM PHOSPHORUS LEVELS
- U-2550 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PREVIOUSLY TREATED FOLLICULAR LYMPHOMA IN COMBINATION WITH A RITUXIMAB PRODUCT
- U-2551 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PREVIOUSLY TREATED MARGINAL ZONE LYMPHOMA IN COMBINATION WITH A RITUXIMAB PRODUCT
- U-2552 METHOD OF TREATING POSTPARTUM DEPRESSION
- U-2553 PREVENTION OF PREGNANCY IN FEMALES OF REPRODUCTIVE AGE
- U-2554 TREATMENT OF MILD TO MODERATE ACTIVE CROHN'S DISEASE INVOLVING THE ILEUM AND/OR THE ASCENDING COLON
- U-2555 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 G OF ELEMENTAL IRON
- U-2556 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULTS WHO HAVE INTOLERANCE TO OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON ASSOCIATED WITH HEAVY UTERINE BLEEDING OR A GASTROINTESTINAL DISORDER BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE
- U-2557 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 GRAMS OF ELEMENTAL IRON
- U-2558 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED METASTATIC BREAST CANCER AFTER FAILURE OF PRIOR CHEMOTHERAPY
- U-2559 USE IN COMBINATION WITH DOXORUBICIN AND CYCLOPHOSPHAMIDE FOR ADJUVANT TREATMENT OF PATIENTS WITH OPERABLE NODE-POSITIVE BREAST CANCER
- U-2560 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER AFTER FAILURE OF PRIOR PLATINUM-BASED CHEMOTHERAPY
- U-2561 USE IN COMBINATION WITH CISPLATIN FOR TREATMENT OF UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WITHOUT PRIOR CHEMOTHERAPY TREATMENT
- U-2562 TREATMENT OF PATIENTS WITH ANDROGEN INDEPENDENT (HORMONE REFRACTORY) METASTATIC PROSTATE CANCER IN COMBINATION WITH PREDNISONE
- U-2563 TREATMENT OF ADVANCED GASTRIC ADENOCARCINOMA IN COMBINATION WITH CISPLATIN AND FLUOROURACIL IN PATIENTS THAT HAVE NOT RECEIVED PRIOR CHEMOTHERAPY
- U-2564 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED SQUAMOUS CELL CARCINOMA OF THE HEAD AND NECK IN COMBINATION WITH CISPLATIN AND FLUOROURACIL
- U-2565 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA
- U-2566 TREATMENT OF VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA
- U-2567 ONCE DAILY TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN ADULTS WITH 1% OXYMETAZOLINE HYDROCHLORIDE CREAM, WHERE THE PATIENT EXPERIENCES NO REBOUND OR WORSENING OF FACIAL ERYTHEMA POST-TREATMENT
- U-2568 TREATMENT OF HYPOACTIVE SEXUAL DESIRE DISORDER (HSDD)
- U-2569 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVE AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH AN EFFECTIVE AMOUNT OF TEZACAFTOR AND IVACAFTOR
- U-2570 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGES 6 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 10,206,877
- U-2571 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGES 6 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-2572 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 6 YEARS AND OLDER, WITH A F508DEL OR G551D CFTR GENE MUTATION AND A A455E, 2789+5G->, OR 3849+10KBC->T MUTATION, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF U.S.

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- U-2573 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 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-2574 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 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-2575 TREATING CYSTIC FIBROSIS PATIENTS AGES 6 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-2576 TREATMENT OF COMMUNITY ACQUIRED BACTERIAL PNEUMONIA
- U-2577 TREATMENT OF THROMBOCYTOPENIA IN AN ADULT PATIENT WITH CHRONIC IMMUNE THROMBOCYTOPENIA WHO HAS HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT
- U-2578 TREATMENT OF THROMBOCYTOPENIA IN AN ADULT PATIENT WITH CHRONIC LIVER DISEASE WHO IS SCHEDULED TO UNDERGO A PROCEDURE
- U-2579 REDUCTION IN A SUBJECT'S RISK OF EXPERIENCING A BREAKTHROUGH OVERT HEPATIC ENCEPHALOPATHY (HE) EPISODE
- U-2580 A METHOD OF TREATING TYPE 2 DIABETES COMPRISING ADMINISTERING SEMAGLUTIDE ONCE WEEKLY IN A AMOUNT OF 1.0 MG TO A SUBJECT IN NEED THEREOF
- U-2581 TREATING HYPOTENSION WITH ABOUT 20 NG/KG/MIN TO ABOUT 40 NG/KG/MIN ANGIOTENSIN II IN A HUMAN SUBJECT HAVING SEPTIC SHOCK
- U-2582 FOR THE ORAL PREVENTION/PROPHYLAXIS OF MALARIA IN ADULTS, COMPRISING A THREE-PHASE DOSING REGIMEN CONSISTING OF A LOADING/INITIAL DOSE, A MAINTENANCE/EXPOSURE DOSE, AND A TERMINAL/POST-EXPOSURE DOSE
- U-2583 TREATMENT OF BACTERIAL VAGINOSIS IN ADULT WOMEN
- U-2584 XPROVIO IS INDICATED IN COMBINATION WITH DEXAMETHASONE TO TREAT RELAPSED OR REFRACTORY MULTIPLE MYELOMA (REFRACTORY TO AT LEAST AN ANTI-CD38 MAB, 2 PROTEASOME INHIBITORS AND 2 IMMUNOMODULATORY AGENTS) IN ADULTS WHO RECEIVED AT LEAST 4 PRIOR THERAPIES
- U-2585 TREATMENT OF PARENTERAL NUTRITION-ASSOCIATED CHOLESTASIS IN PATIENTS UNDER THE AGE OF 12
- U-2586 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS, INCLUDING PYELONEPHRITIS (CUTI)
- U-2587 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTIONS (CIAI)
- U-2588 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH DAPAGLIFLOZIN AND METFORMIN
- U-2589 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH BASAL INSULIN OR BASAL INSULIN PLUS METFORMIN
- U-2590 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN, A SULFONYLUREA, A THIAZOLIDINEDIONE, OR COMBINATION OF ANY TWO OF THESE THERAPIES
- U-2591 LOWERING PLASMA GLUCAGON IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING EXENATIDE AS AN ADJUNCT TO DIET AND EXERCISE
- U-2592 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A SUSTAINED-RELEASE EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE
- U-2593 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION ONCE WEEKLY AS AN ADJUNCT TO DIET AND EXERCISE TO ACHIEVE A MEAN STEADY STATE PLASMA CONCENTRATION OF EXENATIDE AT LEAST 170 PG/ML
- U-2594 REDUCING FASTING PLASMA GLUCOSE IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION ONCE WEEKLY AS AN ADJUNCT TO DIET AND EXERCISE TO ACHIEVE A MEAN STEADY STATE PLASMA CONCENTRATION OF EXENATIDE AT LEAST 170 PG/ML
- U-2595 REDUCING BODY WEIGHT IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION ONCE WEEKLY AS AN ADJUNCT TO DIET AND EXERCISE TO ACHIEVE

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- A MEAN STEADY STATE PLASMA CONCENTRATION OF EXENATIDE AT LEAST 170 PG/ML
- U-2596 REDUCING HBA1C IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION ONCE WEEKLY AS AN ADJUNCT TO DIET AND EXERCISE TO ACHIEVE A MEAN STEADY STATE PLASMA CONCENTRATION OF EXENATIDE AT LEAST 170 PG/ML
- U-2597 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH DAPAGLIFLOZIN AS ADD-ON TO METFORMIN
- U-2598 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN INJECTABLE SUSTAINED RELEASE FORMULATION OF EXENATIDE AS AN ADJUNCT TO DIET AND EXERCISE
- U-2599 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE TO PROVIDE A RELEASE PROFILE HAVING A RATIO OF C-MAX TO C-AVG OF ABOUT 3 OR LESS
- U-2600 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A PRE-MIXED EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE
- U-2601 STIMULATING INSULIN RELEASE IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A PRE-MIXED EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE
- U-2602 DELAYING GASTRIC EMPTYING IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A PRE-MIXED EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE
- U-2603 METHOD OF TREATING IRON DEFICIENCY
- U-2604 TREATMENT OF SEVERE HYPOGLYCEMIA IN PATIENTS WITH DIABETES
- U-2605 TREATMENT OF PATIENTS WITH NON-METASTATIC CASTRATION RESISTANT PROSTATE CANCER
- U-2606 TREATMENT OF ADULT PATIENTS WITH SYMPTOMATIC TENOSYNOVIAL GIANT CELL TUMOR (TGCT) ASSOCIATED WITH SEVERE MORBIDITY OR FUNCTIONAL LIMITATIONS AND NOT AMENABLE TO IMPROVEMENT WITH SURGERY
- U-2607 TREATMENT OF ADULT PATIENTS WITH INTERMEDIATE-2 OR HIGH-RISK PRIMARY OR SECONDARY MYELOFIBROSIS
- U-2608 METHOD OF TREATING SCHIZOPHRENIA
- U-2609 A METHOD FOR INDUCING A REGIONAL ANAESTHESIA VIA INTRATHECAL ADMINISTRATION OF A PATENTED PRESERVATIVE FREE SOLUTION FOR INJECTION (WITH A SPECIFIC COMPOSITION, PH, OSMOLALITY AND DENSITY) CONTAINING 9-11 MG/ML CHLOROPROCAINE HCL
- U-2610 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTION IN PATIENTS WITH END-STAGE RENAL DISEASE ON HEMODIALYSIS
- U-2611 TREATMENT OF COMPLICATED URINARY TRACT INFECTION IN PATIENTS WITH END-STAGE RENAL DISEASE ON HEMODIALYSIS
- U-2612 TREATMENT OF RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN PATIENTS 10 YEARS OF AGE AND OLDER
- U-2613 TREATMENT OF RELAPSING-REMITTING SCLEROSIS (MS)
- U-2614 TREATMENT OF MODERATE TO SEVERE DYSpareunia
- U-2615 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE ADMINISTRATION OF TASIMELTEON WITH FOOD
- U-2616 TREATMENT OF ADULTS WITH MODERATELY TO SEVERELY ACTIVE RHEUMATOID ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO METHOTREXATE
- U-2617 TREATMENT OF ROS1-POSITIVE NON-SMALL CELL LUNG CANCER
- U-2618 TREATMENT OF SOLID TUMORS THAT HAVE A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION
- U-2619 TREATMENT OF ADULTS WITH COMMUNITY-ACQUIRED BACTERIAL PNEUMONIA CAUSED BY SUSCEPTIBLE MICROORGANISMS
- U-2620 USE OF NINTEDANIB FOR SLOWING THE RATE OF DECLINE IN PULMONARY FUNCTION IN PATIENTS WITH SYSTEMIC SCLEROSIS-ASSOCIATED INTERSTITIAL LUNG DISEASE (SSC-ILD)
- U-2621 MODIFIED DOSING REGIMEN FOR THE MANAGEMENT OF MILD TO MODERATE PAIN
- U-2622 TREATMENT OF ACUTE UNCOMPLICATED INFLUENZA IN PATIENTS 2 YEARS AND OLDER

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- U-2623 A METHOD OF REDUCING OFF TIME FROM L-DOPA THERAPY, COMPRISING ADMINISTERING, TO A HUMAN PATIENT WITH PARKINSON'S DISEASE, AN EFFECTIVE AMOUNT OF ISTRADÉFYLLINE, WHEREIN THE PATIENT CURRENTLY RECEIVES SAID L-DOPA THERAPY
- U-2624 TREATMENT OF METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (MCSPC)
- U-2625 TOPICAL TREATMENT OF PLAQUE PSORIASIS IN ADULTS
- U-2626 METHOD OF TREATING IRRITABLE BOWEL SYNDROME WITH CONSTIPATION BY ADMINISTERING TENAPANOR
- U-2627 TOPICAL TREATMENT OF PLAQUE PSORIASIS IN PATIENTS 12 YEARS AND OLDER
- U-2628 METHOD OF TREATING TYPE 2 DIABETES MELLITUS
- U-2629 TREATMENT OF HIV-1 INFECTION IN ADULT PATIENTS AS A REPLACEMENT THERAPY IN VIROLOGICALLY SUPPRESSED ADULTS WITH NO HISTORY OF TREATMENT FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED WITH RESISTANCE TO THE INDIVIDUAL COMPONENTS OF DELSTRIGO
- U-2630 FOR USE IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 AS A REPLACEMENT THERAPY IN VIROLOGICALLY SUPPRESSED ADULTS WITH NO HISTORY OF TREATMENT FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED WITH RESISTANCE TO DORAVIRINE
- U-2631 TREATMENT OF COMPLICATED URINARY TRACT INFECTION
- U-2632 REDUCTION OF RISK OF END STAGE KIDNEY DISEASE, DOUBLING OF SERUM CREATININE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS
- U-2633 TREATMENT OF ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER, PROGRESSED ON: CRIZOTINIB + AT LEAST 1 OTHER ALK INHIBITOR FOR METASTATIC DISEASE; OR ALECTINIB, OR CERITINIB AS FIRST ALK INHIBITOR FOR METASTATIC DISEASE.
- U-2634 METHOD OF TREATMENT IN PATIENTS WITH CONCOMITANT ANGIOEDEMA
- U-2635 TREATMENT OF ACUTE URTICARIA
- U-2636 METHOD OF INCREASING PEAK PLASMA OR ONSET OF PLASMA CONCENTRATION BY INTRAVENOUS INJECTION IN INDIVIDUALS IN NEED OF TREATMENT FOR ACUTE URTICARIA
- U-2637 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER WITH SYMPTOMS OF URINARY FREQUENCY, URGENCY, OR URGE INCONTINENCE WITH A SINGLE UNIT DOSE OF 10% OXYBUTYNIN CHLORIDE GEL
- U-2638 INCREASE PAIN-FREE LIGHT EXPOSURE IN ADULT PATIENTS WITH A HISTORY OF PHOTOTOXIC REACTIONS FROM ERYTHROPOIETIC PROTOPORPHYRIA (EPP)
- U-2639 METHOD OF ACTIVATING RARGAMMA RECEPTOR
- U-2640 PROPHYLAXIS OF VENOUS THROMBOEMBOLISM IN ACUTELY ILL MEDICAL PATIENTS AT RISK FOR THROMBOEMBOLIC COMPLICATIONS NOT AT HIGH RISK OF BLEEDING
- U-2641 PROPHYLAXIS OF VENOUS THROMBOEMBOLISM IN ACUTELY ILL MEDICAL PATIENTS AT RISK FOR THROMBOEMBOLIC COMPLICATIONS NOT AT HIGH RISK OF BLEEDING WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST 5 CONSECUTIVE DAYS
- U-2642 METHOD OF TREATING CANCER BY DETECTING A CREATININE CLEARANCE OF A PATIENT AND ADMINISTERING LONSURF
- U-2643 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS 65 YEARS OF AGE OR OLDER AND SYMPTOMS THEREOF
- U-2644 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS 65 YEARS OF AGE OR OLDER
- U-2645 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH ELEXACFTOR, TEZACFTOR, AND IVACFTOR
- U-2646 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE ONE F508DEL MUTATION AND ONE R117H MUTATION IN THE CFTR GENE WITH ELEXACFTOR, TEZACFTOR, AND IVACFTOR
- U-2647 TREATMENT OF NON-NODULAR ACNE VULGARIS
- U-2648 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH ELEXACFTOR, TEZACFTOR, AND IVACFTOR
- U-2649 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT

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LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH A COMPOSITION COMPRISING ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR; AND ANOTHER COMPOSITION COMPRISING IVACAFTOR

- U-2650 TREATMENT OF CF IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE USING A SOLID COMPOSITION COMPRISING ELEXACAFTOR, TEZACAFTOR, AMORPHOUS IVACAFTOR, AND LESS THAN ABOUT 30% CRYSTALLINE IVACAFTOR
- U-2651 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF A PHARMACEUTICAL COMPOSITION COMPRISING ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-2652 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 10081621
- U-2653 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-2654 TREATMENT OF REFRACTORY CHRONIC GRAFT-VERSUS-HOST DISEASE
- U-2655 A METHOD OF TREATMENT OF ADVANCED OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER ASSOCIATED WITH HOMOLOGOUS RECOMBINATION DEFICIENCY (HRD) POSITIVE STATUS
- U-2656 TREATMENT OF ADULT PATIENTS WITH ACTIVE PSORIATIC ARTHRITIS
- U-2657 TREATMENT OF PATIENTS WITH MODERATE TO SEVERE PLAQUE PSORIASIS WHO ARE CANDIDATES FOR PHOTOTHERAPY OR SYSTEMIC THERAPY
- U-2658 TREATMENT OF ADULT PATIENTS WITH ORAL ULCERS ASSOCIATED WITH BEHCET'S DISEASE
- U-2659 TREATMENT OF ADULT PATIENTS WITH ORAL ULCERS ASSOCIATED WITH BEHCET'S DISEASE USING A DOSAGE TITRATION SCHEDULE
- U-2660 TREATMENT OF H. PYLORI INFECTION IN ADULTS
- U-2661 CHRONIC WEIGHT MANAGEMENT IN ADULT PATIENTS USING AN EXTENDED RELEASE TABLET CONTAINING LORCARSERIN HYDROCHLORIDE HEMIHYDRATE
- U-2662 USE OF CALCIPOTRIENE FOAM FOR THE TOPICAL TREATMENT OF PLAQUE PSORIASIS IN PATIENTS AGED 4 YEARS AND OLDER
- U-2663 USE IN SONOHYSTEROSALPINOGRAPHY TO ASSESS FALLOPIAN TUBE PATENCY
- U-2664 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING BY INDUCING SKIPPING OF EXON 51
- U-2665 TREATMENT OF CHRONIC GRAFT VERSUS HOST DISEASE AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- U-2666 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA
- U-2667 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA
- U-2668 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA IN COMBINATION WITH OBINUTUZUMAB
- U-2669 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED SMALL LYMPHOCYTIC LEUKEMIA IN COMBINATION WITH OBINUTUZUMAB
- U-2670 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LEUKEMIA
- U-2671 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LEUKEMIA IN COMBINATION WITH OBINUTUZUMAB
- U-2672 TREATMENT OF ACUTE HEPATIC PORPHYRIA
- U-2673 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING BY CORRECTING A DEFECTIVE GENE FOR DYSTROPHIN
- U-2674 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING BY RESTORING OR INCREASING FUNCTIONAL DYSTROPHIN PROTEIN PRODUCTION
- U-2675 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS HAVING A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 53 SKIPPING
- U-2676 TREATMENT OF SICKLE CELL DISEASE BY ADMINISTERING VOXELOTOR, AS RECITED IN CLAIM 1

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2677 PROPHYLAXIS OF ORGAN REJECTION IN PATIENTS CONVERTED FROM TACROLIMUS IMMEDIATE-RELEASE FORMULATIONS
- U-2678 PROPHYLAXIS OF ORGAN REJECTION IN DE NOVO TRANSPLANT PATIENT
- U-2679 TREATING LOW BLOOD PRESSURE WITH ANGIOTENSIN II AT AN INITIAL RATE OF ABOUT 20 NG/KG/MIN AND TITRATING DOWN TO ACHIEVE AND/OR MAINTAIN A MAP OF ABOUT 65 MM HG OR ABOVE
- U-2680 TREATING LOW BLOOD PRESSURE WITH ANGIOTENSIN II WITH AN INITIAL RATE OF ABOUT 5 NG/KG/MIN TO ABOUT 20 NG/KG/MIN IN A SUBJECT HAVING REFRACTORY HYPOTENSION OR SEVERE HYPOTENSION
- U-2681 TREATING LOW BLOOD PRESSURE WITH ANGIOTENSIN II WITH AN INITIAL RATE OF ABOUT 5 NG/KG/MIN TO ABOUT 20 NG/KG/MIN IN A SUBJECT HAVING REFRACTORY HYPOTENSION OR SEVERE HYPOTENSION, AND TITRATING THE RATE UP
- U-2682 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100MG OF ACALABRUTINIB TWICE DAILY
- U-2683 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100MG OF ACALABRUTINIB TWICE DAILY
- U-2684 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY IN COMBINATION WITH OBINUTUZUMAB
- U-2685 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED SMALL LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY IN COMBINATION WITH OBINUTUZUMAB
- U-2686 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY
- U-2687 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LEUKEMIA IN COMBINATION WITH OBINUTUZUMAB BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY
- U-2688 USE OF VASCEPA TO LOWER TRIGLYCERIDES AND LDL-C IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (ABOUT 200 MG/DL TO LESS THAN ABOUT 500 MG/DL) AND ON STATIN THERAPY
- U-2689 USE OF VASCEPA TO TREAT MIXED DYSLIPIDEMIA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (≥ 150 MG/DL) AND ON STATIN THERAPY
- U-2690 USE OF VASCEPA TO LOWER TRIGLYCERIDES IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (ABOUT 200 MG/DL TO LESS THAN ABOUT 500 MG/DL) AND ON STATIN THERAPY
- U-2691 USE OF VASCEPA TO TREAT HYPERTRIGLYCERIDEMIA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (≥ 150 MG/DL) AND ON STATIN THERAPY
- U-2692 USE OF VASCEPA TO REDUCE TRIGLYCERIDES IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (≥ 150 MG/DL) AND ON STATIN THERAPY
- U-2693 USE OF VASCEPA TO REDUCE TRIGLYCERIDES IN A MIXED DYSLIPIDEMIA ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (≥ 150 MG/DL) AND ON STATIN THERAPY
- U-2694 USE OF VASCEPA TO LOWER TRIGLYCERIDES IN A MIXED DYSLIPIDEMIA ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (ABOUT 200 MG/DL TO LESS THAN ABOUT 500 MG/DL) AND ON STATIN THERAPY
- U-2695 USE OF VASCEPA TO TREAT MIXED HYPERTRIGLYCERIDEMIA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (≥ 150 MG/DL) AND ON STATIN THERAPY
- U-2696 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CARDIOVASCULAR DEATH, CORONARY REVASCULARIZATION, AND UNSTABLE ANGINA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS (TG ≥ 150 MG/DL TO ABOUT 500 MG/DL)
- U-2697 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CARDIOVASCULAR DEATH AND/OR UNSTABLE ANGINA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS (TG ≥ 150 MG/DL TO ABOUT 500 MG/DL)
- U-2698 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CARDIOVASCULAR DEATH AND/OR CORONARY REVASCULARIZATION IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS (TG ≥ 150 MG/DL TO ABOUT 500 MG/DL)
- U-2699 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CARDIOVASCULAR EVENT (CORONARY REVASCULARIZATION, UNSTABLE ANGINA, STROKE AND/OR MYOCARDIAL INFARCTION) IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2700 USE OF VASCEPA TO REDUCE TRIGLYCERIDES IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (ABOUT 200 MG/DL TO LESS THAN ABOUT 500 MG/DL) AND ON ROSUVASTATIN THERAPY
- U-2701 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CORONARY REVASCULARIZATION AND/OR UNSTABLE ANGINA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS (TG \geq 150 MG/DL TO ABOUT 500 MG/DL)
- U-2702 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CARDIOVASCULAR EVENT (CARDIOVASCULAR DEATH, CORONARY REVASCULARIZATION AND/OR UNSTABLE ANGINA) IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS
- U-2703 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CV EVENT (CV DEATH, CORONARY REVASCULARIZATION, UNSTABLE ANGINA, STROKE AND/OR MYOCARDIAL INFARCTION) IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS AND DIABETES MELLITUS
- U-2704 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS AND AT LEAST ONE RISK FACTOR FOR CARDIOVASCULAR DISEASE
- U-2705 METHOD OF USING CAPSAICIN IN COMBINATION WITH A GEL COMPOSITION FOR REMOVAL OF CAPSAICIN FROM A TREATMENT AREA OR UNINTENDED AREA
- U-2706 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF ONSET AND/OR RECURRENCE OF CARDIOVASCULAR EVENTS IN A PATIENT WHO HAS ESCAPED THE UNSTABLE PERIOD AFTER CARDIOVASCULAR ANGIOPLASTY
- U-2707 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE OCCURRENCE OF A CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH HYPERCHOLESTEROLEMIA
- U-2708 THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER
- U-2709 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE
- U-2710 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE
- U-2711 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE 750 MG OF ELEMENTAL IRON
- U-2712 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE 750 MG OF ELEMENTAL IRON
- U-2713 MODULATION OF 5-HYDROXYTRYPTAMINE 2 RECEPTOR ACTIVITY IN SCHIZOPHRENIA
- U-2714 TREATMENT OF SCHIZOPHRENIA WITH IMPROVEMENT IN RESIDUAL SYMPTOMS OF SCHIZOPHRENIA
- U-2715 TREATMENT OF SICKLE CELL DISEASE BY ADMINISTERING VOXELOTOR, AS RECITED IN CLAIM 2
- U-2716 MAINTENANCE TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GBRCA-MUTATED METASTATIC PANCREATIC ADENOCARCINOMA WHOSE DISEASE HAS NOT PROGRESSED ON AT LEAST 16 WEEKS OF A FIRST-LINE PLATINUM-BASED CHEMOTHERAPY REGIMEN
- U-2717 ACUTE TREATMENT OF MIGRAINE WITH HEADACHE, WITH OR WITHOUT AURA IN ADULTS
- U-2718 ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA IN ADULTS
- U-2719 TREATMENT OF RELAPSING REMITTING MULTIPLE SCLEROSIS BY DETERMINING VARICELLA ZOSTER VIRUS (VZV) STATUS AND VACCINATING PRIOR TO COMMENCING TREATMENT
- U-2720 ANTIMYCOTIC USES, SPECIFICALLY TREATMENT OF ONYCHOMYCOSIS; TOPICAL TREATMENT OF THE TOENAIL(S) DUE TO TRICHOPHYTON RUBRUM AND TRICHOPHYTON MENTAGROPHYTES
- U-2721 TOPICAL TREATMENT OF TINEA UNGUIUM BY USING AN APPLICATOR FOR APPLYING A SOLUTION FOR TREATING TINEA UNGUIUM TO AN AFFECTED PART OF A PATIENT
- U-2722 METHOD OF INTRAVENOUSLY ADMINISTERING A DILUTED CYSTEINE HYDROCHLORIDE SOLUTION TO A NEONATE IN NEED THEREOF
- U-2723 MAINTENANCE MONOTHERAPY TREATMENT OF BIPOLAR 1 DISORDER
- U-2724 A METHOD OF ORAL DELIVERY OF TREPROSTINIL COMPRISING ADMINISTERING AN ORAL OSMOTIC PHARMACEUTICAL DOSAGE FORM

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2725 A METHOD OF TREATING PULMONARY HYPERTENSION AND PULMONARY ARTERIAL HYPERTENSION BY ADMINISTERING AN ORAL OSMOTIC PHARMACEUTICAL DOSAGE FORM
- U-2726 TREATMENT OF UNRESECTABLE OR METASTATIC GASTROINTESTINAL STROMAL TUMOR (GIST) HARBORING A PLATELET-DERIVED GROWTH FACTOR RECEPTOR ALPHA (PDGFRA) EXON 18 MUTATION
- U-2727 NASAL ADMINISTRATION OF DIAZEPAM FOR TREATMENT OF INTERMITTENT, STEREOTYPIC EPISODES OF FREQUENT SEIZURE ACTIVITY IN PATIENTS 6 YEARS OF AGE AND OLDER
- U-2728 USE OF PEMETREXED WITH PRIOR AND/OR REPEATED VITAMIN B12 AND FOLIC ACID ADMINISTRATION IN PATIENTS WITH NON-SQUAMOUS NON-SMALL CELL LUNG CANCER
- U-2729 USE OF PEMETREXED WITH PRIOR AND/OR REPEATED VITAMIN B12 AND FOLIC ACID ADMINISTRATION IN PATIENTS WITH MESOTHELIOMA
- U-2730 METHOD OF TREATING TYPE 2 DIABETES MELLITUS USING A PHARMACEUTICAL COMPOSITION COMPRISING EMPAGLIFLOZIN, LINAGLIPTIN AND METFORMIN
- U-2731 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 IN COMBINATION WITH LINAGLIPTIN AND METFORMIN
- U-2732 METHOD OF TREATING TYPE 2 DIABETES USING A PHARMACEUTICAL COMPOSITION COMPRISING LINAGLIPTIN, METFORMIN, EMPAGLIFLOZIN AND A BASIC AMINO ACID
- U-2733 METHOD OF TREATING A TYPE 2 DIABETES MELLITUS PATIENT WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE THERAPY WITH METFORMIN USING A PHARMACEUTICAL COMPOSITION COMPRISING EMPAGLIFLOZIN, LINAGLIPTIN AND METFORMIN
- U-2734 METHOD OF TREATMENT OF IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON, WHO HAVE NON-HEMODIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE, BY ADMINISTERING FERRIC DERISOMALTOSE
- U-2735 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS USING 150MG ELAGOLIX WHILE CO-ADMINISTERING RIFAMPIN
- U-2736 METHOD OF TREATING EPITHELIOID SARCOMA
- U-2737 METHOD OF TREATING EPITHELIOID SARCOMA BY INHIBITING ENHANCER OF ZESTE HOMOLOG 2 (EZH2)
- U-2738 METHOD OF TREATING A LUNG METASTASIS OF EPITHELIOID SARCOMA
- U-2739 INCREASING BLOOD PRESSURE WITH AN INITIAL RATE OF ABOUT 20 NG/KG/MIN ANGIOTENSIN II IN A HUMAN SUBJECT HAVING SEPTIC SHOCK, AND TITRATING THE RATE UP.
- U-2740 INCREASING BLOOD PRESSURE WITH A RATE OF ABOUT 20 NG/KG/MIN TO ABOUT 40 NG/KG/MIN ANGIOTENSIN II IN A HUMAN SUBJECT HAVING SEPTIC SHOCK
- U-2741 TREATMENT OF CLOSTRIDIODES DIFFICILE-ASSOCIATED DIARRHEA (CDAD) IN PATIENTS FROM 6 MONTHS OF AGE AND OLDER
- U-2742 TREATMENT OF SEVERE HYPOGLYCEMIA
- U-2743 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF UNSTABLE ANGINA IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE
- U-2744 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF STROKE IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE
- U-2745 TREATMENT OF NEUROBLASTOMAS THAT HAVE A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION
- U-2746 USE OF NEXLIZET AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY TO LOWER LDL-C IN ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE
- U-2747 USE OF NEXLETOL AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY TO LOWER LDL-C IN ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE
- U-2748 USE OF NEXLETOL AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY FOR INHIBITING CHOLESTEROL SYNTHESIS TO LOWER LDL-C IN ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE
- U-2749 USE OF NEXLIZET AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY FOR INHIBITING CHOLESTEROL SYNTHESIS TO LOWER LDL-C IN ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2750 MANAGEMENT OF MODERATE-TO-SEVERE PAIN BY INTRAVENOUS INJECTION

U-2751 A TRANSDERMAL METHOD OF CONTRACEPTION

U-2752 METHOD OF USING L-CYSTEINE IN AN ADMIXTURE FOR TREATING PATIENTS NEEDING PARENTERAL NUTRITION

U-2753 INCREASING SURVIVAL IN METASTATIC CASTRATION-RESISTANT PROSTATE CANCER PATIENTS PREVIOUSLY TREATED WITH DOCETAXEL BY ADMINISTERING AS A 3 WEEK CYCLE CABAZITAXEL AFTER 5 MG DEXCHLORPHENIRAMINE, 8 MG DEXAMETHASONE, AND AN H2-AGONIST

U-2754 TREATMENT OF POST-OPERATIVE NAUSEA AND VOMITING

U-2755 OCULAR EXAMINATION, INTRAOCULAR PRESSURE MEASUREMENT, OR REMOVAL OF FOREIGN BODIES OR SUTURES, IN ADULT AND PEDIATRIC PATIENTS REQUIRING A DISCLOSING AGENT IN COMBINATION WITH A TOPICAL OPHTHALMIC ANESTHETIC

U-2756 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CARDIOVASCULAR DEATH IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE

U-2757 DOSING REGIMEN FOR THE TREATMENT OF SCHIZOPHRENIA IN ADULTS BY ADMINISTERING TWO LOADING DOSES OF PALIPERIDONE PALMITATE FOLLOWED BY MAINTENANCE DOSE(S)

U-2758 DOSING REGIMEN FOR THE TREATMENT OF SCHIZOAFFECTIVE DISORDER IN ADULTS AS A MONOTHERAPY AND AS AN ADJUNCT TO MOOD STABILIZERS OR ANTIDEPRESSANTS BY ADMINISTERING TWO LOADING DOSES OF PALIPERIDONE PALMITATE FOLLOWED BY MAINTENANCE DOSE(S)

U-2759 REDUCTION OF INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH OPEN ANGLE GLAUCOMA(OAG) OR OCULAR HYPERTENSION (OHT) WITH A BIODEGRADABLE BIMATOPROST IMPLANT

U-2760 TOPICAL TREATMENT OF ACNE VULGARIS IN PATIENTS 12 YEARS OF AGE OR OLDER

U-2761 INTRAVENOUS SOTALOL DOSING REGIMEN FOR ACHIEVING STEADY STATE CONCENTRATION (EXPOSURE) FASTER COMPARED TO THE CONVENTIONAL ORAL DOSING IN A FACILITY THAT CAN PROVIDE ELECTROCARDIOGRAPHIC MONITORING

U-2762 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A MAJOR CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH DIABETES MELLITUS AND TWO OR MORE ADDITIONAL RISK FACTORS FOR CARDIOVASCULAR DISEASE

U-2763 METHOD OF TREATING ADULTS WITH SCHIZOPHRENIA COMPRISING ADMINISTERING ASENAPINE VIA A TRANSDERMAL PATCH

U-2764 TREATMENT OF POST-OPERATIVE INFLAMMATION AND PAIN FOLLOWING OCULAR SURGERY

U-2765 TREATMENT OF HIV-1 INFECTION IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 40 KG WHO HAVE NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY OR ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS

U-2766 TX OF HIV1 INFECTION USING A COMPOSITION CONTAINING A PK ENHANCER THAT INHIBITS CY P450 MONOOXYGENASE IN ADULTS & PEDIATRIC PATIENTS AT LEAST 40KG HAVING NO PRIOR ARV TX HISTORY OR ARE VIROLOGICALLY SUPPRESSED ON A STABLE ARV REGIMEN FOR AT LEAST 6 MO

U-2767 TREATMENT OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PK ENHANCER THAT INHIBITS CY P450 MONOOXYGENASE IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 40KG WHO HAVE NO PRIOR ARV TREATMENT HISTORY

U-2768 TREATMENT OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PK ENHANCER THAT INHIBITS CY P450 MONOOXYGENASE IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 40KG WHO ARE VIROLOGICALLY SUPPRESSED ON A STABLE ARV REGIMEN FOR AT LEAST 6 MONTHS

U-2769 DOSING REGIMEN FOR INTRAVENOUS SOTALOL FOR ADMINISTRATION IN A FACILITY THAT CAN PROVIDE CONTINUOUS ELECTROCARDIOGRAPHIC MONITORING AND CARDIAC RESUSCITATION.

U-2770 CUSHING'S DISEASE

U-2771 TREATMENT OF MODERATE TO SEVERE PLAQUE PSORIASIS IN PATIENTS 18 YEARS OF AGE OR OLDER

U-2772 MAINTENANCE TREATMENT OF CHRONIC PULMONARY DISEASE (COPD)

U-2774 TREATMENT OF RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN ADULTS

U-2775 TREATMENT OF A TYPE 2 DIABETES MELLITUS PATIENT WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE METFORMIN THERAPY USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2776 TREATMENT OF A TYPE 2 DIABETES MELLITUS PATIENT WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE METFORMIN THERAPY USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN AND LINAGLIPTIN
- U-2777 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 6 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING A PHARMACEUTICAL COMPOSITION ACCORDING TO CLAIM 2 OF U.S. PATENT NO. 10,597,384, FURTHER COMPRISING IVACAFTOR
- U-2778 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 2 TO 5 YEARS OLD WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING A PHARMACEUTICAL COMPOSITION ACCORDING TO CLAIM 2 OF U.S. PATENT NO. 10,597,384, FURTHER COMPRISING IVACAFTOR
- U-2779 TREATMENT OF SPASTICITY
- U-2780 USE FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH LENNOX-GASTAUT SYNDROME
- U-2781 USE FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH DRAVET SYNDROME
- U-2782 USE FOR REDUCING CONVULSIVE SEIZURE FREQUENCY IN PATIENTS WITH LENNOX GASTAUT SYNDROME
- U-2783 USE FOR REDUCING CONVULSIVE SEIZURE FREQUENCY IN PATIENTS WITH DRAVET SYNDROME
- U-2784 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) AND WHERE THE MUTANT IDH1 HAS THE ABILITY TO CONVERT ALPHA-KETOGLUTARATE INTO 2-HYDROXYGLUTARATE (2-HG)
- U-2785 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) AND WHERE THE MUTANT IDH1 HAS THE ABILITY TO CONVERT ALPHA-KETOGLUTARATE INTO 2-HYDROXYGLUTARATE (2-HG)
- U-2786 METHOD OF PREVENTING PREGNANCY BY INSERTING A VAGINAL SYSTEM CONTAINING 103 MG OF SEGESTERONE ACETATE AND 17.4 MG ETHINYL ESTRADIOL INTO A VAGINA FOR UP TO THIRTEEN 21/7-DAY (IN/OUT) CYCLES
- U-2787 METHOD OF CONTRACEPTION BY INSERTING A VAGINAL SYSTEM FOR UP TO 13 21/7-DAY (IN/OUT) CYCLES, WHEREIN EFFICACY REQUIRES THE SYSTEM CANNOT BE OUT OF THE VAGINA FOR MORE THAN 2 CUMULATIVE HOURS IN ANY SUCH CYCLE WITHOUT USING ALTERNATIVE CONTRACEPTION
- U-2788 TREATMENT OF BREAST CANCER INCLUDING HER2 (ERBB2)-POSITIVE OR -OVEREXPRESSING BREAST CANCER
- U-2789 POTASSIUM PHOSPHATES INJECTION IS INDICATED AS A SOURCE OF PHOSPHORUS IN INTRAVENOUS FLUIDS TO CORRECT HYPOPHOSPHATEMIA IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-2790 TREATMENT OF A TREATMENT-NAIVE PATIENT WITH INADEQUATELY CONTROLLED TYPE 2 DIABETES USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN AND LINAGLIPTIN
- U-2791 TREATMENT OF ADULT PATIENTS WITH INSOMNIA, CHARACTERIZED BY DIFFICULTIES WITH SLEEP ONSET AND/OR SLEEP MAINTENANCE
- U-2792 TREATMENT OF A TREATMENT-NAIVE PATIENT WITH INADEQUATELY CONTROLLED TYPE 2 DIABETES USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN
- U-2793 A METHOD FOR DELIVERING NITRIC OXIDE TO A PATIENT WITH PULMONARY HYPERTENSION OR HYPOXIA
- U-2794 TREATMENT OF TYPE 2 DIABETES MELLITUS WITH 100 MG CANAGLIFLOZIN PER DAY
- U-2795 TREATMENT OF TYPE 2 DIABETES MELLITUS WITH 300 MG CANAGLIFLOZIN PER DAY
- U-2796 REDUCTION OF RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS WITH 100 MG CANAGLIFLOZIN PER DAY
- U-2797 REDUCTION OF RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS WITH 300 MG CANAGLIFLOZIN PER DAY
- U-2798 REDUCTION OF RISK OF END STAGE KIDNEY DISEASE, DOUBLING OF SERUM CREATININE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS WITH 100 MG CANAGLIFLOZIN PER DAY
- U-2799 REDUCTION OF RISK OF END STAGE KIDNEY DISEASE, DOUBLING OF SERUM CREATININE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS WITH 300 MG CANAGLIFLOZIN PER DAY
- U-2800 TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH NEUROFIBROMATOSIS TYPE 1 (NF1) WHO HAVE SYMPTOMATIC, INOPERABLE PLEXIFORM NEUROFIBROMAS (PN)

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2801 METHOD OF IRON ADMINISTRATION TO TREAT PATIENTS IN NEED OF IRON REPLACEMENT THERAPY
- U-2802 BRAFTOVI IS A KINASE INHIBITOR INDICATED IN COMBINATION WITH CETUXIMAB, FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC COLORECTAL CANCER (CRC) WITH A BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST, AFTER PRIOR THERAPY
- U-2803 BRAFTOVI IS A KINASE INHIBITOR 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
- U-2804 A METHOD FOR THE IMPROVEMENT OF NEUROLOGICAL OUTCOME BY REDUCING THE INCIDENCE AND SEVERITY OF ISCHEMIC DEFICITS IN ADULT PATIENTS WITH SUBARACHNOID HEMORRHAGE (SAH) FROM RUPTURED INTRACRANIAL BERRY ANEURYSMS
- U-2805 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY ADMINISTERING TASIMELTEON TO PATIENTS WITH A SMOKING HISTORY
- U-2806 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE ADMINISTRATION OF FOOD
- U-2807 TREATMENT OF MODERATE TO SEVERE MIGRAINE PAIN WITH PAIN FREE AT 2 HOURS POST ADMINISTRATION
- U-2808 TREATMENT OF DYSKINESIA, DECREASING OFF TIME, AND INCREASING ON TIME WITHOUT TROUBLESOME DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-2809 FOR THE TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FIBROBLAST GROWTH FACTOR RECEPTOR 2 (FGFR2) FUSION OR OTHER REARRANGEMENT
- U-2810 METHOD OF SUPPORTING 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-2811 METHOD OF TREATING PARKINSON'S DISEASE
- U-2812 ADJUNCTIVE TREATMENT TO LEVODOPA/CARBIDOPA IN PATIENTS WITH PARKINSON'S DISEASE
- U-2813 USE FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH A MESENCHYMAL-EPITHELIAL TRANSITION (MET) EXON 14 SKIPPING MUTATION
- U-2814 A METHOD OF PROPHYLACTIC TREATMENT OF GOUT FLARES IN ADULTS COMPRISES ADMINISTERING TO A PATIENT A LIQUID COLCHICINE ORAL SOLUTION
- U-2815 MEASURING TIME-VARYING CHANGE IN BLOOD IN A TISSUE VOLUME USING MODIFIED BEER-LAMBERT LAW IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO-AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY
- U-2816 METHOD FOR TREATING INFLUENZA
- U-2817 METHOD OF INHIBITING COMT IN THE PERIPHERY
- U-2818 METHOD OF REDUCING O-METHYLATION OF L-DOPA
- U-2819 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF ADV. EPITHELIAL OVARIAN CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION
- U-2820 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF ADV. EPITHELIAL OVARIAN CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIV WITH GENOMIC INSTABILITY
- U-2821 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF FALLOPIAN TUBE CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION
- U-2822 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF FALLOPIAN TUBE CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIV WITH GENOMIC INSTABILITY
- U-2823 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF PRIMARY PERITONEAL CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION
- U-2824 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF PRIMARY PERITONEAL CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH GENOMIC INSTABILITY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2825 TREATMENT OF "OFF" EPISODES IN PATIENTS WITH PARKINSON'S DISEASE

U-2826 TREATMENT OF ADULT PATIENTS WITH METASTATIC RET FUSION-POSITIVE NON-SMALL CELL LUNG CANCER

U-2827 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET-MUTANT MEDULLARY THYROID CANCER (MTC) WHO REQUIRE SYSTEMIC THERAPY

U-2828 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE REFRACTORY (IF RADIOACTIVE IODINE IS APPROPRIATE)

U-2829 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS

U-2830 A METHOD FOR TREATING METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC), WHEREIN THE CANCER IS ASSOCIATED WITH A DELETERIOUS BRCA-MUTATION

U-2831 TREATMENT OF PARTIAL-ONSET SEIZURES IN A PATIENT WITH REFRACTORY PARTIAL-ONSET SEIZURES

U-2832 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC HOMOLOGOUS RECOMBINATION REPAIR GENE-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER, WHICH HAS PROGRESSED FOLLOWING PRIOR TREATMENT WITH ENZALUTAMIDE OR ABIRATERONE

U-2833 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC BRCA-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER, WHICH HAS PROGRESSED FOLLOWING PRIOR TREATMENT WITH ENZALUTAMIDE OR ABIRATERONE

U-2834 TREATMENT OF ADULTS WITH PERSISTENT/RECURRENT CHRONIC THROMBOEMBOLIC PULMONARY HYPERTENSION (CTEPH), (WHO GROUP 4) AFTER SURGICAL TREATMENT, OR INOPERABLE CTEPH, TO IMPROVE EXERCISE CAPACITY AND WHO FUNCTIONAL CLASS

U-2835 TREATMENT OF ADULTS WITH PULMONARY HYPERTENSION (PAH), (WHO GROUP 1), TO IMPROVE EXERCISE CAPACITY, WHO FUNCTIONAL CLASS AND TO DELAY CLINICAL WORSENING

U-2836 TREATMENT OF ADULT PATIENTS WITH SMALL CELL LUNG CANCER (SCLC) WITH DISEASE PROGRESSION ON OR AFTER PLATINUM-BASED CHEMOTHERAPY.

U-2837 TREATMENT OF ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC)

U-2838 REDUCTION OF THE RATE OF A FIRST MYOCARDIAL INFARCTION OR STROKE IN PATIENTS WITH CORONARY ARTERY DISEASE AT HIGH RISK FOR SUCH EVENTS

U-2839 TREATMENT OF MYOCARDIAL INFARCTION OR STROKE IN PATIENTS WITH CORONARY ARTERY DISEASE AT HIGH RISK FOR SUCH EVENTS

U-2840 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP)

U-2841 USE OF VASCEPA WITH HIGH INTENSITY STATIN THERAPY TO REDUCE THE RISK OF A CV EVENT IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS AND (1) ESTABLISHED CV DISEASE, OR (2) DIABETES MELLITUS AND TWO OR MORE ADDITIONAL RISK FACTORS FOR CV DISEASE

U-2842 MANAGEMENT OF HEAVY MENSTRUAL BLEEDING ASSOCIATED WITH UTERINE LEIOMYOMAS (FIBROIDS)

U-2843 NASAL ADMINISTRATION OF METOCLOPRAMIDE FOR TREATMENT OF DIABETIC GASTROPARESIS

U-2844 IN COMBINATION WITH PEMBROLIZUMAB FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH ADVANCED RENAL CELL CARCINOMA

U-2845 A METHOD OF TREATING A HUMAN PATIENT SUFFERING FROM PULMONARY HYPERTENSION

U-2846 TREATMENT OF CHRONIC GRAFT VERSUS HOST DISEASE REFRACTORY TO SYSTEMIC THERAPY

U-2847 REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) RECURRENCE IN ADULTS

U-2848 TREATMENT OF TRAVELERS' DIARRHEA (TD) CAUSED BY NONINVASIVE STRAINS OF ESCHERIA COLI IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER

U-2849 METHOD OF TREATING BLEPHAROPTOSIS

U-2850 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS USING 150 MG ELAGOLIX WHILE CO-ADMINISTERING KETOCONAZOLE

U-2851 METHOD OF TREATING RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA POSITIVE FOR AN ENHANCER OF ZESTE HOMOLOG 2 (EZH2) MUTATION BY INHIBITING EZH2

U-2852 METHOD OF TREATING RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA POSITIVE FOR AN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- ENHANCER OF ZESTE HOMOLOG 2 (EZH2) MUTATION
- U-2853 METHOD OF TREATING RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA
- U-2854 METHOD OF TREATING RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA BY INHIBITING EZH2
- U-2855 XPOVIO IS INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL), NOT OTHERWISE SPECIFIED, INCLUDING DLBCL ARISING FROM FOLLICULAR LYMPHOMA, AFTER AT LEAST 2 LINES OF SYSTEMIC THERAPY
- U-2856 INCREASING SURVIVAL IN METASTATIC CASTRATION-RESISTANT PROSTATE CANCER PATIENTS PREVIOUSLY TREATED WITH DOCETAXEL BY ADMINISTERING 20 TO 25 MG/M2 CABAZITAXEL AFTER A PREMEDICATION REGIMEN THAT INCLUDES AN H2-ANTAGONIST
- U-2857 USE OF ORAL OCTREOTIDE FOR LONG-TERM MAINTENANCE TREATMENT IN ACROMEGALY PATIENTS WHO HAVE RESPONDED TO AND TOLERATED TREATMENT WITH OCTREOTIDE OR LANREOTIDE
- U-2858 USE IN COMBINATION WITH STIRIPENTOL, VALPROATE, AND CLOBAZAM FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-2859 USE OF CARDIAC MONITORING AND RESTRICTED DISTRIBUTION OF FENFLURAMINE TO MITIGATE RISK OF CARDIOVASCULAR TOXICITY IN THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-2860 USE IN COMBINATION WITH STIRIPENTOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-2861 USE IN COMBINATION WITH CANNABIDIOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-2862 USE FOR THE TREATMENT OF FOCAL SEIZURES IN PATIENTS WITH DRAVET SYNDROME
- U-2863 TOPICAL TREATMENT OF HEAD LICE INFESTATION IN PATIENTS 6 MONTHS OF AGE AND OLDER
- U-2864 METHOD FOR INHIBITING CYTIDINE DEAMINASE BY ADMINISTERING CEDAZURIDINE
- U-2865 TREATMENT OF MYELODYSPLASTIC SYNDROME
- U-2866 TREATMENT OF CHRONIC MYELOMONOCYTIC LEUKEMIA
- U-2867 METHOD FOR INHIBITING DEGRADATION OF A CDA SUBSTRATE BY ADMINISTERING CEDAZURIDINE
- U-2868 TREATMENT OF NON-INFECTIOUS UVEITIS AFFECTING THE POSTERIOR SEGMENT OF THE EYE
- U-2869 IV ADMINISTRATION OF CANGRELOR BEFORE PCI AND CONTINUOUS INFUSION FOR AT LEAST 2 HOURS OR THE DURATION OF THE PCI AND, DURING OR AFTER THE CONTINUOUS INFUSION, ADMINISTRATION OF A LOADING DOSE OF TICAGRELOR OR AN EQUIVALENT THERAPY (PER LABELING)
- U-2870 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 2
- U-2871 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 2 AND 3
- U-2872 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 4 AND 5
- U-2873 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 4, 5, AND 6
- U-2874 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 7 AND 8
- U-2875 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 7, 8, AND 9
- U-2876 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 10 AND 11
- U-2877 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 10, 11, AND 12

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PATENT USE

U-2878 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 13 AND 14

U-2879 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 13, 14, AND 15

U-2880 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 16 AND 17

U-2881 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 16, 17, AND 18

U-2882 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 19 AND 20

U-2883 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 19, 20, AND 21

U-2884 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 22, 23, AND 24

U-2885 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 25 AND 26

U-2886 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 25, 26, AND 27

U-2887 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 28 AND 29

U-2888 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 28, 29, AND 30

U-2889 USE FOR THE MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)

U-2890 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 AND 2 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1

U-2891 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MEYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 3 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1

U-2892 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 AND 4 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1

U-2893 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 AND 5 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1

U-2894 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 AND 6 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1

U-2895 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 AND 7 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1

U-2896 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN

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PATENT USE

SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 8 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1

U-2897 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 CLAIMS 1 AND 9

U-2898 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 CLAIMS 1 AND 10

U-2899 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 CLAIMS 1 AND 11

U-2900 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 CLAIMS 1 AND 13

U-2901 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 AND 15 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1

U-2902 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 AND 16 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1

U-2903 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 AND 17 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1

U-2904 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, 17 AND 18 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1

U-2905 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 AND 19 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1

U-2906 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 22 WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 22

U-2907 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 22 AND 23 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 22

U-2908 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 22 AND 24 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 22

U-2909 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 22 AND 25 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 22

U-2910 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 26 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26

U-2911 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 26 AND 27 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2912 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 26 AND 28 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
- U-2913 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 26 AND 29 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
- U-2914 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 26 AND 30 WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
- U-2915 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 26 AND 31 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
- U-2916 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 26 AND 32 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2917 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 26 AND 33 WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS
- U-2918 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 26 AND 34 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2919 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 26 AND 36 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2920 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 26 AND 38 AND WHEREIN THE EFFECTS ARE AS RECITED CLAIM 26
- U-2921 ONCE DAILY TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN ADULTS
- U-2922 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 26 AND 39 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
- U-2923 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 26 AND 40 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
- U-2924 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 42 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 42
- U-2925 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 42 AND 43 WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 42
- U-2926 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 44 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2927 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 44 AND 45 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44

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PATENT USE

- U-2928 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 44 AND 46 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2929 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 44 AND 47 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2930 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 44 AND 48 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2931 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 44 AND 49 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2932 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 44 AND 50 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2933 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 44 AND 51 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2934 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 44 AND 52 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2935 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 44 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 44 AND 54
- U-2936 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 44 AND 56 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2937 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 44 AND 57 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2938 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN PATIENTS 7 YEARS OF AGE AND OLDER WITH NARCOLEPSY WITH A MIXTURE OF SODIUM, POTASSIUM, MAGNESIUM, AND CALCIUM SALTS OF GHB
- U-2939 TREATMENT OF HIV INFECTION IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 40KG USING A COMPOSITION CONTAINING A PHARMACOKINETIC ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASE
- U-2940 METHOD OF TREATING PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH CENTRAL PRECOCIOUS PUBERTY
- U-2941 A METHOD OF USING AN AEROSOL DELIVERY DEVICE TO AEROSOLIZE GLYCOPYRROLATE FOR THE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-2942 METHOD OF TREATING ACNE VULGARIS WITH TOPICALLY APPLIED CORTEXOLONE 17A-PROPIONATE
- U-2943 TREATMENT OF RELAPSED OR REFRACTORY CHRONIC LYMPHOCYTIC LEUKEMIA
- U-2944 TREATMENT OF RELAPSED OR REFRACTORY SMALL LYMPHOCYTIC LYMPHOMA
- U-2945 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
- U-2946 TREATMENT OF COLORECTAL CANCER THAT HAS A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION
- U-2947 KYPROLIS IS INDICATED IN COMBINATION WITH DARATUMUMAB PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

RECEIVED ONE TO THREE LINES OF THERAPY

- U-2948 A METHOD OF POSITIONING AN INTRAUTERINE SYSTEM (IUS) BY DETERMINING A DEPTH OF THE UTERUS, HOLDING AN INSERTER HANDLE WITH ONE HAND, INSERTING THE IUS INTO THE UTERUS, AND RETRACTING A SLIDER ON THE HANDLE TO RELEASE THE IUS INTO THE UTERUS
- U-2949 A METHOD FOR INDUCING A POST-SURGICAL ANALGESIA SPARING EFFECT BY IMPLANTING AT THE SURGICAL SITE A COLLAGEN SPONGE CONTAINING BUPIVACAINE HCL WHICH PROVIDES LOCAL ANESTHESIA FOR UP TO 24 HOURS FOLLOWING IMPLANTATION
- U-2950 CONTINUED TREATMENT OF ADULTS WITH ACUTE MYELOID LEUKEMIA WHO ACHIEVED FIRST COMPLETE REMISSION (CR) OR CR WITH INCOMPLETE BLOOD COUNT RECOVERY FOLLOWING INTENSIVE INDUCTION CHEMOTHERAPY AND ARE NOT ABLE TO COMPLETE INTENSIVE CURATIVE THERAPY
- U-2951 USE OF CU-64 DOTATATE WITH POSITRON EMISSION TOMOGRAPHY (PET) FOR LOCALIZATION OF SOMATOSTATIN RECEPTOR POSITIVE NEUROENDOCRINE TUMORS (NETS) IN ADULT PATIENTS
- U-2952 TREATMENT OF ADULT PATIENTS WITH METASTATIC REARRANGED DURING TRANSFECTION (RET) FUSION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA APPROVED TEST
- U-2953 USE OF FLUTICASONE FUROATE FOR THE TREATMENT OF AN INFLAMMATORY OR ALLERGIC CONDITIONS, INCLUDING CHRONIC OBSTRUCTIVE PULMONARY DISEASE AND ASTHMA
- U-2954 METHOD OF DISPENSING A COMBINATION MEDICAMENT PRODUCT FROM CLAIMED DELIVERY DEVICE, FOR EXAMPLE FOR THE TREATMENT OF ASTHMA OR COPD
- U-2955 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA; AND ASTHMA
- U-2956 METHOD OF TREATING LAMBERT-EATON MYASTHENIC SYNDROME WITH AMIFAMPRIDINE
- U-2957 MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS 18 YRS AND OLDER, OR CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
- U-2958 A METHOD FOR CONTRACEPTION, THE METHOD COMPRISING ADMINISTERING A TABLET COMPRISING 20 MG TO 30 MG OF ULIPRISTAL ACETATE TO A WOMAN WITHIN 120 HOURS AFTER AN UNPROTECTED INTERCOURSE
- U-2959 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A THIRD AND FURTHER CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH ELEVATED TG LEVELS (≥ 150 MG/DL) AND ESTABLISHED CARDIOVASCULAR DISEASE
- U-2960 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A SECOND OR FURTHER CARDIOVASCULAR (CV) EVENT IN AN ADULT PATIENT WITH ELEVATED TG LEVELS (≥ 150 MG/DL) AND DIABETES MELLITUS AND 2 OR MORE ADDITIONAL RISK FACTORS FOR CV DISEASE
- U-2961 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF MYOCARDIAL INFARCTION, STROKE, BOTH IN AN ADULT PATIENT WITH TYPE 2 DIABETES MELLITUS
- U-2962 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CORONARY REVASCULARIZATION IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE
- U-2963 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-2964 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS A R117H MUTATION IN THE CFTR GENE
- U-2965 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF USING IVACAFTOR IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-2966 TREATMENT OF CF IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING A SOLID COMPOSITION COMPRISING AMORPHOUS (LESS THAN ABOUT 30% CRYSTALLINE) IVACAFTOR
- U-2967 TREATMENT OF CF IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 10272046
- U-2968 USE OF REMIMAZOLAM FOR INDUCTION AND MAINTENANCE OF PROCEDURAL SEDATION IN ADULTS UNDERGOING PROCEDURES LASTING 30 MINUTES OR LESS
- U-2969 TREATMENT OF ADULT PATIENTS WITH CYCLOSPORIN-RESISTANT, STEROID-DEPENDENT/REFRACTORY, OR STEROID RESISTANT CHRONIC GRAFT-VERSUS-HOST DISEASE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- U-2970 TREATMENT OF ADULT PATIENTS WITH CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- U-2971 THE TREATMENT OF ADULT PATIENTS WITH METASTATIC RET FUSION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC)
- U-2972 THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET-MUTANT MEDULLARY THYROID CANCER (MTC) WHO REQUIRE SYSTEMIC THERAPY
- U-2973 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY, WHEREIN THE CANCER IS PAPILLARY THYROID CANCER
- U-2974 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY, WHEREIN THE CANCER IS MEDULLARY THYROID CANCER
- U-2975 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY, WHEREIN THE CANCER IS DIFFERENTIATED THYROID CANCER
- U-2976 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY, WHEREIN THE CANCER IS RECURRENT THYROID CANCER
- U-2977 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE REFRACTORY DIFFERENTIATED THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND ARE RADIOACTIVE IODINE-REFRACTORY
- U-2978 TREATMENT OF HIV-1 INFECTION IN ADULT OR PEDIATRIC PATIENTS (≥ 40 KG) WITH < 50 COPIES/ML HIV-1 RNA AFTER ≥ 6 MONTHS ON PRIOR ANTIRETROVIRAL REGIMEN AND NO KNOWN DARUNAVIR OR TENOFOVIR RESISTANCE-ASSOCIATED SUBSTITUTIONS
- U-2979 METHOD COMPRISING IV ADMINISTRATION OF CANGRELOR BEFORE PCI THEN CONTINUOUS INFUSION FOR AT LEAST 2 HOURS OR THE DURATION OF PCI AND, DURING OR AFTER CONTINUOUS INFUSION, ADMINISTRATION OF A LOADING DOSE OF TICAGRELOR, OR AN EQUIVALENT METHOD
- U-2980 METHOD OF TREATING AN ALLERGIC REACTION USING AN AUTO-INJECTOR
- U-2981 A METHOD OF TREATING ACUTE MYELOGENOUS LEUKEMIA (AML) IN A SUBJECT BY ADMINISTERING A PHARMACEUTICAL COMPOSITION WHERE THE AML IS CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 AND THE AML IS NEWLY DIAGNOSED
- U-2982 A METHOD OF TREATING ACUTE MYELOGENOUS LEUKEMIA (AML) IN A SUBJECT BY ADMINISTERING A PHARMACEUTICAL COMPOSITION WHERE THE AML IS CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 AND WHERE THE AML IS RELAPSED/REFRACTORY
- U-2983 TREATMENT OF SCHIZOPHRENIA BY RAPID AND CONTINUOUS INJECTION
- U-2984 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS AND PEDIATRIC PATIENTS (AT LEAST 12 YEARS OF AGE AND 40 KG) REQUIRING HOSPITALIZATION
- U-2985 A METHOD FOR TREATING DRY EYE IN A PATIENT
- U-2986 MANAGEMENT OF ACUTE PAIN BY INTRAVENOUS INJECTION
- U-2987 METHOD OF TREATING LUNG CANCER, UNDIFFERENTIATED SARCOMA, OR COLORECTAL CANCER THAT EXHIBITS AN NTRK GENE FUSION
- U-2988 REDUCTION OF THE RISK OF STROKE IN PATIENTS WITH ACUTE ISCHEMIC STROKE OR HIGH-RISK TRANSIENT ISCHEMIC ATTACK
- U-2989 METHOD OF USE AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- U-2990 METHOD OF USE FOR TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- U-2991 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A TABLET CONTAINING SELEXIPAG
- U-2992 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A SOLID PREPARATION CONTAINING SELEXIPAG

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PATENT USE

- U-2993 A METHOD FOR TREATING A SUBJECT HAVING ADHD, SAID METHOD COMPRISING ORALLY ADMINISTERING TO SAID SUBJECT A RACEMIC METHYLPHENIDATE CHEWABLE TABLET AS CLAIMED
- U-2994 REDUCTION OF RISK OF MYOCARDIAL INFARCTION, STROKE OR CARDIOVASCULAR DEATH IN A PATIENT WITH CHRONIC CAD OR PAD BY ADMINISTERING CLINICALLY PROVEN EFFECTIVE AMOUNTS THAT ARE 2.5 MG RIVAROXABAN TWICE DAILY AND 75-100 MG ASPIRIN DAILY
- U-2995 TREATMENT OF PRIMARY HYPEROXALURIA TYPE 1 (PH1)
- U-2996 ADMINISTRATION OF AN EXTENDED RELEASE TABLET FOR THE TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY
- U-2997 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF STROKE IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDES AND ATRIAL FIBRILLATION
- U-2998 METHOD OF ADMINISTERING DOCETAXEL TO A SUBJECT COMBINING THE DOCETAXEL PRO-EMULSION FORMULATION WITH AN AQUEOUS MEDIUM TO PRODUCE DOCETAXEL EMULSION
- U-2999 METHOD OF USE OF TREATING, AS AN INITIAL LOADING DOSE FOR MONOTHERAPY OR ADJUNCTIVE THERAPY IN PARTIAL ONSET SEIZURE PATIENTS WITH EPILEPSY AGED 17 YEARS OR OLDER
- U-3000 METHOD FOR POST-EXPOSURE PROPHYLAXIS OF INFLUENZA
- U-3001 PROCEDURES IN ADULT AND PEDIATRIC PATIENTS REQUIRING A DISCLOSING AGENT IN COMBINATION WITH A TOPICAL OPHTHALMIC ANESTHETIC.
- U-3002 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY (IF RADIOACTIVE IODINE IS APPROPRIATE)
- U-3003 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY ADMINISTERING TASIMELTEON
- U-3004 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH A STRONG CYP1A2 INHIBITOR
- U-3005 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE USE OF TASIMELTEON WITH RIFAMPIN
- U-3006 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH CYP1A2 STRONG INHIBITORS
- U-3007 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE ADMINISTRATION OF TASIMELTEON WITH FOOD
- U-3008 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY ADMINISTERING TASIMELTEON TO PATIENTS WITH A SMOKING HISTORY
- U-3009 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY ADMINISTERING TASIMELTEON TO PATIENTS WITH A SMOKING HISTORY
- U-3010 PROPHYLAXIS TO PREVENT ATTACKS OF HEREDITARY ANGIOEDEMA (HAE) IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-3011 A METHOD FOR TREATING OCULAR INFLAMMATION
- U-3012 TREATMENT IN COMBINATION WITH ANDROGEN DEPRIVATION THERAPY OF NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NMCRPC) THAT IMPROVES METASTASIS FREE SURVIVAL
- U-3013 TREATMENT IN COMBINATION WITH ORCHIECTOMY OF NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NMCRPC)
- U-3014 METHOD FOR THE INDUCTION OF LOCAL ANESTHESIA OF THE MUCOUS MEMBRANES
- U-3015 TOPICAL TREATMENT OF ACTINIC KERATOSIS OF THE FACE OR SCALP
- U-3016 ADJUVANT THERAPY AFTER TUMOR RESECTION IN PATIENTS WITH NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS
- U-3017 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS AND/OR CATAPLEXY IN NARCOLEPSY PATIENTS WITH A SALT OF GAMMA-HYDROXYBUTYRATE WHEN DIVALPROEX SODIUM IS CONCOMITANTLY ADMINISTERED
- U-3018 XPOVIO IS INDICATED IN COMBINATION WITH BORTEZOMIB AND DEXAMETHASONE FOR THE

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PATENT USE

- TREATMENT OF ADULT PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-3019 TREATMENT OF ADULT PATIENTS WITH ADVANCED PROSTATE CANCER THAT IS SEX-HORMONE-DEPENDENT
- U-3020 TREATMENT OF ADULT PATIENTS WITH ADVANCED PROSTATE CANCER
- U-3021 TEZACAFTOR AND IVACAFTOR FOR THE TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE A R117H MUTATION IN THE CFTR GENE
- U-3022 TREATMENT OF CF IN PATIENTS 12 YEARS AND OLDER WHO HAVE A F508DEL OR G551D CFTR MUTATION AND A 2ND MUTATION SELECTED FROM R117H, A455E, 2789+5G->A, & 3849+10KBC->T, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF US 10058546
- U-3023 TREATMENT OF CF IN PATIENTS 6 YEARS AND OLDER WHO HAVE A F508DEL OR G551D CFTR MUTATION AND A 2ND MUTATION SELECTED FROM R117H, A455E, 2789+5G->A, AND 3849+10KBC->T, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF US 10058546
- U-3024 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 12 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR WHO HAVE AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,081,621
- U-3025 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR WHO HAVE AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,081,621
- U-3026 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 12 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,206,877
- U-3027 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,206,877
- U-3028 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-3029 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE A R117H MUTATION IN THE CFTR GENE WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-3030 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELX, TEZ, AND IVA
- U-3031 TREATMENT OF CF IN PATIENTS 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA USING A SOLID COMPOSITION COMPRISING ELX, TEZ, AMORPHOUS IVA, AND < ~30% CRYSTALLINE IVA
- U-3032 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 10081621
- U-3033 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-3034 TREATMENT OF TRD IN ADULTS BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE 2X WEEKLY FOR 4 WEEKS DURING THE INDUCTION PHASE IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3035 TREATMENT OF DEPRESSIVE SYMPTOMS IN ADULTS WITH MDD WITH ACUTE SUICIDAL IDEATION OR BEHAVIOR BY NASALLY ADMINISTERING 56MG OR 84 MG OF ESKETAMINE 2X WEEKLY FOR 4 WEEKS IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3036 TREATMENT OF TRD IN ADULTS BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE 2X WEEKLY FOR 4 WEEKS DURING THE INDUCTION PHASE FOLLOWED BY A MAINTENANCE PHASE OF 56MG OR 84 MG WEEKLY OR 1X EVERY TWO WEEKS IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3037 A METHOD OF DELIVERING NITRIC OXIDE TO A PATIENT

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- U-3038 MANAGEMENT OF MODERATE-TO-SEVERE PAIN BY INTRAVENOUS INJECTION IN PATIENTS WITH MILD RENAL IMPAIRMENT
- U-3039 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 53 SKIPPING
- U-3040 LONG-TERM TREATMENT OF ADULTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 EXTENSIVE METABOLIZERS WITH MILD HEPATIC IMPAIRMENT AND ARE CONCURRENTLY TAKING A STRONG OR MODERATE CYP3A INHIBITOR
- U-3041 LONG-TERM TREATMENT OF ADULTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 EXTENSIVE METABOLIZERS WITH MODERATE TO SEVERE RENAL IMPAIRMENT
- U-3042 LONG-TERM TREATMENT OF ADULTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 POOR METABOLIZERS WITH 84 MG ONCE DAILY OF ELIGLUSTAT (EQUIVALENT TO 100 MG OF ELIGLUSTAT TARTRATE)
- U-3043 LONG-TERM TREATMENT OF ADULTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 EXTENSIVE OR INTERMEDIATE METABOLIZERS WITH 84 MG TWICE PER DAY OF ELIGLUSTAT (EQUIVALENT TO 100 MG OF ELIGLUSTAT TARTRATE TWICE PER DAY)
- U-3044 AXITINIB IN COMBINATION WITH AVELUMAB FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH ADVANCED RENAL CELL CARCINOMA
- U-3045 TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY
- U-3046 METHOD OF ADMINISTERING VALBENAZINE WHILE AVOIDING CONCOMITANT USE OF A STRONG CYP3A4 INDUCER
- U-3047 USE IN COMBINATION WITH CAPECITABINE, FOR THE TREATMENT OF ADULT PATIENTS WITH ADVANCED OR METASTATIC HER2-POSITIVE BREAST CANCER WHO HAVE RECEIVED TWO OR MORE PRIOR ANTI-HER2 BASED REGIMENS IN THE METASTATIC SETTING
- U-3048 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE 500 TO 750 MG OF ELEMENTAL IRON
- U-3049 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE 500 TO 750 MG OF ELEMENTAL IRON
- U-3050 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING AT LEAST ABOUT 0.6G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MIN OR LESS
- U-3051 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING AT LEAST ABOUT 0.6G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MIN OR LESS
- U-3052 USE OF VASCEPA TO REDUCE TRIGLYCERIDE LEVELS IN AN ADULT PATIENT ON STATIN THERAPY AND HAVING ATRIAL FIBRILLATION AND TRIGLYCERIDE LEVELS OF GREATER THAN 500 MG/DL
- U-3053 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF MYOCARDIAL INFARCTION IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS AND ESTABLISHED CV DISEASE OR DIABETES MELLITUS AND TWO OR MORE ADDITIONAL RISK FACTORS FOR CV DISEASE
- U-3054 TREATMENT OF DRUG-INDUCED EXTRAPYRAMIDAL REACTIONS IN ADULT PATIENTS WITH PARKINSON'S DISEASE
- U-3055 A METHOD OF TREATING HUNTINGTON'S CHOREA
- U-3056 TREATMENT OF PATIENTS WITH ACTIVE LUPUS NEPHRITIS
- U-3057 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) THAT IS ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE
- U-3058 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER AND YOUNG ADULTS WITH RELAPSED OR REFRACTORY, SYSTEMIC ANAPLASTIC LARGE CELL LYMPHOMA (ALCL) THAT IS ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE
- U-3059 TREATMENT OF HIV-1 INFECTION IN ADULTS TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO HISTORY OF TREATMENT FAILURE
- U-3060 TREATMENT OF HIV INFECTION IN ADULTS
- U-3061 TREATMENT OF HIV-1 IN AN ADULT IN COMBINATION WITH RILPIVIRINE

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- U-3062 REDUCING THE RISK OF CARDIOVASCULAR DEATH AND HEART FAILURE (HF) HOSPITALIZATION FOLLOWING A HOSPITALIZATION FOR HF OR NEED FOR OUTPATIENT IV DIURETICS, IN ADULTS WITH SYMPTOMATIC CHRONIC HF AND EJECTION FRACTION LESS THAN 45%
- U-3063 RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA (MZL) WHO HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED REGIMEN
- U-3064 RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL) WHO HAVE RECEIVED AT LEAST THREE PRIOR LINES OF SYSTEMIC THERAPY
- U-3065 TREATMENT OF ADULTS WITH METASTATIC UROTHELIAL CARCINOMA THAT HAS SUSCEPTIBLE FGFR3 OR FGFR2 GENETIC ALTERATIONS AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY
- U-3066 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC, SURGICALLY UNRESECTABLE UROTHELIAL CARCINOMA THAT HAS SUSCEPTIBLE FGFR3 OR FGFR2 GENETIC ALTERATIONS AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY
- U-3067 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA THAT HAS SUSCEPTIBLE FGFR3 GENETIC ALTERATIONS AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY
- U-3068 TREATMENT IN COMBINATION WITH CABOTEGRAVIR OF HIV-1 INFECTION IN ADULTS TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO HISTORY OF TREATMENT FAILURE
- U-3069 TREATMENT OF ACUTE UNCOMPLICATED INFLUENZA IN PATIENTS 6 MONTHS AND OLDER
- U-3070 REDUCING THE RISK OF MORTALITY IN HUTCHINSON-GILFORD PROGERIA SYNDROME (HGPS)
- U-3071 USE FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH TUBEROUS SCLEROSIS COMPLEX
- U-3072 USE FOR THE TREATMENT OF GENERALIZED SEIZURES OR FOCAL SEIZURES WITH IMPAIRMENT IN PATIENTS WITH TUBEROUS SCLEROSIS COMPLEX
- U-3073 USE FOR REDUCING SEIZURE FREQUENCY IN PATIENTS WITH TUBEROUS SCLEROSIS COMPLEX
- U-3074 METHOD FOR PROVIDING SUSTAINED LOCAL ANESTHESIA FOR AT LEAST 24 HOURS
- U-3075 TREATMENT OF ADRENAL INSUFFICIENCY
- U-3076 METHOD OF TREATING TARDIVE DYSKINESIA WHILE AVOIDING CONCOMITANT USE OF A STRONG CYP3A4 INDUCER
- U-3077 TREATING A SOLID TUMOR, INCLUDING LUNG CANCER, WITH A MET ALTERATION(S), OR STABILIZING OR IMPROVING SYMPTOMS ASSOCIATED WITH HAVING A SOLID TUMOR, INCLUDING LUNG CANCER, WITH A MET ALTERATION(S), BY ADMINISTERING AN EFFECTIVE AMOUNT OF TEPOTINIB
- U-3078 TREATING A SOLID TUMOR, INCLUDING LUNG CANCER, HAVING A MET KINASE ALTERATION(S) BY ADMINISTERING AN EFFECTIVE AMOUNT OF TEPOTINIB
- U-3079 A METHOD TO DECREASE THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A PLATINUM/ETOPOSIDE-CONTAINING REGIMEN FOR EXTENSIVE-STAGE SMALL CELL LUNG CANCER
- U-3080 A METHOD TO DECREASE THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A TOPOTECAN-CONTAINING REGIMEN FOR EXTENSIVE- STAGE SMALL CELL LUNG CANCER
- U-3081 A METHOD TO DECREASE THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A CARBOPLATIN AND ETOPOSIDE-CONTAINING REGIMEN FOR EXTENSIVE-STAGE SMALL CELL LUNG CANCER
- U-3082 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH OBSTRUCTIVE SLEEP APNEA (OSA) IN AN ADULT THROUGH A DOSING REGIMEN THAT INCLUDES ORAL ADMINISTRATION OF 75 MG ONCE DAILY FOR AT LEAST 3 DAYS FOLLOWED BY 150 MG ONCE DAILY
- U-3083 METHOD OF TREATING TRANSFUSIONAL IRON OVERLOAD
- U-3084 TREATMENT OF HEART FAILURE WITH PRESERVED EJECTION FRACTION
- U-3085 DOSE MODIFICATION FOR RENAL IMPAIRMENT
- U-3086 FOR HIGH-DOSE CONDITIONING TREATMENT PRIOR TO HEMATOPOIETIC PROGENITOR (STEM) CELL TRANSPLANTATION IN PATIENTS WITH MULTIPLE MYELOMA
- U-3087 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 45 SKIPPING BY INDUCING EXON-SKIPPING OF EXON 45

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- U-3088 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 45 SKIPPING
- U-3089 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 45 SKIPPING BY RESTORING AN MRNA READING FRAME TO INDUCE DYSTROPHIN PROTEIN PRODUCTION
- U-3090 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA IN PATIENTS WITH END-STAGE RENAL DISEASE ON HEMODIALYSIS
- U-3091 TREATMENT OF VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA IN PATIENTS WITH END-STAGE RENAL DISEASE ON HEMODIALYSIS
- U-3092 METHOD OF TREATING MOLYBDENUM COFACTOR DEFICIENCY TYPE A
- U-3093 IN COMBINATION WITH DEXAMETHASONE TO TREAT RELAPSED OR REFRACTORY MULTIPLE MYELOMA (REFRACTORY TO AT LEAST 1 PROTEASOME INHIBITOR, 1 IMMUNOMODULATORY AGENT, AND 1 ANTI-CD38 MAB) IN ADULTS WHO RECEIVED AT LEAST 4 PRIOR LINES OF THERAPY
- U-3094 TREATMENT OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER (ADHD) WITH SERDEXMETHYLPHENIDATE AND DEXMETHYLPHENIDATE
- U-3095 TREATMENT OF HYPERLIPIDEMIA
- U-3096 TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS ARE ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA-APPROVED TEST
- U-3097 EXTENDED ADJUVANT TREATMENT OF ADULT PATIENTS WITH EARLY-STAGE HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-POSITIVE BREAST CANCER, TO FOLLOW ADJUVANT TRASTUZUMAB BASED THERAPY
- U-3098 TREATMENT OF REFRACTORY EPILEPSY PATIENTS WITH FENFLURAMINE THAT REDUCES THE RISK OF CARDIOVASCULAR TOXICITY BY USING CARDIAC MONITORING AND RESTRICTED DISTRIBUTION
- U-3099 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH OBSTRUCTIVE SLEEP APNEA (OSA) IN A PATIENT WITH MODERATE RENAL IMPAIRMENT
- U-3100 A METHOD OF TREATING ADULTS WITH RELAPSED OR REFRACTORY ADVANCED RENAL CELL CARCINOMA FOLLOWING TWO OR MORE PRIOR SYSTEMIC THERAPIES BY INHIBITING THE ANGIOGENESIS OF BLOOD VESSELS WITH A VASCULAR ENDOTHELIAL GROWTH FACTOR INHIBITOR
- U-3101 TREATMENT OF ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS WITH A SINGLE DOSE OF 1200MG ORITAVANCIN OR ITS SINGLE DOSE EQUIVALENT
- U-3102 REDUCTION OF CIRCULATING LYMPHOCYTES IN TREATING RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN ADULTS
- U-3103 TREATMENT OF RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN ADULTS USING A DOSE TITRATION SCHEDULE FOLLOWED BY A MAINTENANCE DOSE
- U-3104 TREATMENT OF C. DIFFICILE-ASSOCIATED DIARRHEA
- U-3105 TREATMENT OF STAPHYLOCOCCAL ENTEROCOLITIS
- U-3106 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE ADMINISTRATION OF TASIMELTEON TO SMOKERS OR TO PATIENTS BEING TREATED WITH A CYP1A2 INHIBITOR
- U-3107 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE ADMINISTRATION OF TASIMELTEON TO SMOKERS OR TO PATIENTS BEING TREATED WITH A CYP1A2 INHIBITOR
- U-3108 TREATMENT OF NEUROGENIC DETRUSOR OVERACTIVITY (NDO) IN PEDIATRIC PATIENTS AGED 3 YEARS AND OLDER BY ADMINISTRATION OF AN EXTENDED-RELEASE SUSPENSION FORMULATION OF MIRABEGRON
- U-3109 METHOD OF USING VISMODEGIB TO TREAT BASAL CELL CARCINOMA
- U-3110 USE OF NALOXONE FOR THE EMERGENCY TREATMENT OF KNOWN OR SUSPECTED OPIOID OVERDOSE, AS MANIFESTED BY RESPIRATORY AND/OR CENTRAL NERVOUS SYSTEM DEPRESSION, FOR ADULT AND PEDIATRIC PATIENTS
- U-3111 TREATING OPIOID USE DISORDER
- U-3112 TREATING NEWLY DIAGNOSED ACUTE MYELOGENOUS LEUKEMIA (AML) CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 BY ADMINISTERING A ONCE DAILY 500 MG ORAL

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DOSE TO A SUBJECT THAT HAS NOT INGESTED A HIGH-FAT MEAL

- U-3113 TREATING RELAPSED/REFRACTORY ACUTE MYELOGENOUS LEUKEMIA (AML) CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 BY ADMINISTERING A ONCE DAILY 500 MG ORAL DOSE TO A SUBJECT THAT HAS NOT INGESTED A HIGH-FAT MEAL
- U-3114 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) BY ORALLY ADMINISTERING VENETOCLAX TO AN ADULT ACCORDING TO A DOSE RAMP-UP THAT INCLUDES A DOSE OF 50 MG PER DAY FOR 1 WEEK FOLLOWED BY 100 MG PER DAY FOR 1 WEEK
- U-3115 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING ABOUT 1 G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MINUTES OR LESS
- U-3116 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING ABOUT 1 G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MINUTES OR LESS
- U-3117 ADMINISTRATION TO THE EYE OF A PATIENT FOR TREATMENT OF DRY EYE CONDITION
- U-3118 TREATMENT OF POSTSURGICAL PAIN PROVIDING ANALGESIA TO A PATIENT FOR UP TO 72 HOURS, FOR EXAMPLE, AFTER BUNIONECTOMY, OPEN INGUINAL HERNIORRHAPHY, OR TOTAL KNEE ARTHROPLASTY VIA SOFT TISSUE OR PERIARTICULAR INSTILLATION
- U-3119 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 11
- U-3120 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1, 11, AND 12
- U-3121 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 13 AND 23
- U-3122 ADJUNCT TO DIET EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 13, 23, AND 24
- U-3123 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF COMPLEMENT INHIBITOR PEGCETACOPLAN
- U-3124 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF PEGCETACOPLAN
- U-3125 USE FOR LOADING DOSE IN PATIENTS WITH SYMPTOMATIC AFIB/AFL WHO ARE CURRENTLY IN SINUS RHYTHM OR FOR THE TREATMENT OF LIFE-THREATENING VENTRICULAR TACHYCARDIA
- U-3126 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A SECOND AND FURTHER CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE
- U-3127 REDUCTION OF THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH HEART FAILURE WITH REDUCED EJECTION FRACTION AND WITHOUT TYPE II DIABETES
- U-3128 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS (CUTI) INCLUDING PYELONEPHRITIS CAUSED BY THE FOLLOWING SUSCEPTIBLE MICROORGANISMS: ESCHERICHIA COLI, KLEBSIELLA PNEUMONIA, ENTEROBACTER CLOACAE SPECIES COMPLEX WITH MEROPENEM & VABORBACTAM AS SPECIFIED
- U-3129 MANAGEMENT OF HEAVY MENSTRUAL BLEEDING ASSOCIATED WITH UTERINE LEIOMYOMAS (FIBROIDS) IN PREMENOPAUSAL WOMEN
- U-3130 METHOD OF POSITRON EMISSION TOMOGRAPHY (PET) IN MEN WITH PROSTATE CANCER
- U-3131 USE OF ZUBSOLV FOR TREATMENT OF OPIOID DEPENDENCE
- U-3132 INDICATED FOR THE TREATMENT OF MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS
- U-3133 TREATING SICKLE CELL DISEASE BY ADMINISTERING 1500 MG OF VOXELOTOR ORALLY ONCE DAILY
- U-3134 INCREASING HEMOGLOBIN TO TREAT SICKLE CELL DISEASE BY ADMINISTERING 1500 MG OF VOXELOTOR ORALLY ONCE DAILY
- U-3135 TREATING SCHIZOPHRENIA
- U-3136 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC

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- OLANZAPINE AND SAMIDORPHAN, WITH REDUCED ANTIPSYCHOTIC INDUCED WEIGHT GAIN
- U-3137 METHOD OF TREATING BIPOLAR DISORDER BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN, WITH REDUCED ANTIPSYCHOTIC INDUCED WEIGHT GAIN
- U-3138 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN, WITH REDUCTION OF THE ADVERSE METABOLIC PROFILE
- U-3139 METHOD OF TREATING BIPOLAR DISORDER BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN, WITH REDUCTION OF THE ADVERSE METABOLIC PROFILE
- U-3140 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN
- U-3141 METHOD OF TREATING BIPOLAR DISORDER BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN
- U-3142 PREVENTIVE TREATMENT OF EPISODIC MIGRAINE IN ADULTS
- U-3143 FOR THE TOPICAL TREATMENT OF PLAQUE PSORIASIS IN PATIENTS 18 YEARS OF AGE AND OLDER
- U-3144 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELEXACFTOR, TEZACFTOR, AND IVACFTOR
- U-3145 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE A R117H MUTATION IN THE CFTR GENE WITH ELEXACFTOR, TEZACFTOR, AND IVACFTOR
- U-3146 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELX, TEZ, AND IVA
- U-3147 FOR THE TREATMENT OF NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC) IN ADULTS AND PEDIATRIC PATIENTS 1 YEAR AND OLDER
- U-3148 METHOD OF TREATING RELAPSING FORMS OF MULTIPLE SCLEROSIS BEFORE AND AFTER ADMINISTERING AN INACTIVE VACCINE
- U-3149 METHOD OF RECONSTITUTING A LYOPHILIZED LIPOSOMAL COMPOSITION FOR ADMINISTERING CYTARABINE AND DAUNORUBICIN TO TREAT NEWLY-DIAGNOSED THERAPY-RELATED AML (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC) IN PATIENTS 1 YEAR AND OLDER
- U-3150 METHOD OF ADMINISTERING A RECONSTITUTED LIPOSOMAL COMPOSITION CONTAINING CYTARABINE AND DAUNORUBICIN TO TREAT NEWLY-DIAGNOSED THERAPY-RELATED AML (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC) IN PATIENTS 1 YEAR AND OLDER
- U-3151 TO IMPROVE WAKEFULNESS IN ADULT PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY OR OBSTRUCTIVE SLEEP APNEA (OSA) WITH A DOSING REGIMEN THAT INCLUDES A DOSE OF 75 MG ONCE DAILY FOR AT LEAST 3 DAYS FOLLOWED BY 150 MG ONCE DAILY
- U-3152 USE BY FEMALES OF REPRODUCTIVE POTENTIAL TO PREVENT PREGNANCY
- U-3153 TREATMENT OF ADVANCED GASTROINTESTINAL STROMAL TUMOR IN PATIENTS HAVING PROGRESSED FROM A FIRST LINE ADMINISTRATION OF IMATINIB, A SECOND LINE ADMINISTRATION OF SUNITINIB, AND A THIRD LINE ADMINISTRATION OF REGORAFENIB
- U-3154 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH A COMPOSITION COMPRISING ELEXACFTOR, TEZACFTOR, AND IVACFTOR; AND ANOTHER COMPOSITION COMPRISING IVACFTOR
- U-3155 TREATMENT OF CF IN PATIENTS 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA USING A SOLID COMPOSITION COMPRISING ELX, TEZ, AMORPHOUS IVA, AND < ~30% CRYSTALLINE IVA
- U-3156 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF A PHARMACEUTICAL COMPOSITION COMPRISING ELEXACFTOR, TEZACFTOR, AND IVACFTOR
- U-3157 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 10081621

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PATENT USE

- U-3158 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-3159 TREATMENT OF ADULT AND POST-MENARCHAL PEDIATRIC FEMALES WITH VULVOVAGINAL CANDIDIASIS (VVC)
- U-3160 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WHO ARE SEVERELY IMMUNOCOMPROMISED
- U-3161 METHOD FOR WEIGHT MANAGEMENT ACCORDING TO A DOSE ESCALATION SCHEDULE
- U-3162 METHOD FOR WEIGHT MANAGEMENT
- U-3163 TREATMENT OF ACTINIC KERATOSES OF UPPER EXTREMITIES BY PHOTODYNAMIC THERAPY
- U-3164 GASTROINTESTINAL TABLETS INDICATED FOR CLEANSING THE COLON IN PREPARATION FOR COLONOSCOPY
- U-3165 METHOD OF TREATING HUMAN SMALLPOX DISEASE
- U-3166 OTC USE: ALLERGY SYMPTOM RELIEVER; TEMPORARY RELIEF OF THESE SYMPTOMS DUE TO HAY FEVER OR OTHER UPPER RESPIRATORY ALLERGIES: NASAL CONGESTION, RUNNY NOSE, SNEEZING AND ITCHY NOSE
- U-3167 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, OR 6 IN PEDIATRIC PATIENTS 3 TO LESS THAN 12 YEARS OF AGE OR WEIGHING LESS THAN 45 KG
- U-3168 TREATMENT OF ADVANCED SYSTEMIC MASTOCYTOSIS, INCLUDING PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH AN ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), AND MAST CELL LEUKEMIA (MCL)
- U-3169 TREATMENT OF TRICHOMONIASIS IN ADULTS
- U-3170 TREATING CHRONIC HEART FAILURE WITH REDUCED EJECTION FRACTION IN PATIENTS NOT TAKING AN ACE INHIBITOR OR AN ARB OR PREVIOUSLY TAKING LOW DOSES OF THESE AGENTS, BY TITRATING UP FROM HALF THE USUALLY RECOMMENDED STARTING DOSE
- U-3171 TREATMENT OF INVASIVE ASPERGILLOSIS IN ADULTS AND PEDIATRIC PATIENTS 13 YEARS OF AGE AND OLDER
- U-3172 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF 1080 MG OF PEGCETACOPLAN
- U-3173 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF 1080 MG OF PEGCETACOPLAN TWICE WEEKLY
- U-3174 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF 1080 MG OF PEGCETACOPLAN EVERY THREE DAYS
- U-3175 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS AND STAPHYLOCOCCUS AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA) INCLUDING THOSE WITH RIGHT-SIDED INFECTIVE ENDOCARDITIS BY ADMINISTERING THE FORMULATION OF DAPTOMYCIN AS RECITED IN CLAIM 18
- U-3176 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS AND S. AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA) INCLUDING THOSE WITH RIGHT-SIDED INFECTIVE ENDOCARDITIS BY RECONSTITUTING AND ADMINISTERING THE FORMULATION AS RECITED IN CLAIM 12
- U-3177 TREATMENT OF VENOUS THROMBOTIC DISEASE
- U-3178 REDUCE THE RISK OF RECURRENCE OF VENOUS THROMBOTIC DISEASE
- U-3179 METHOD OF TREATING FOLLICULAR LYMPHOMA
- U-3180 DECREASING OFF TIME IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-3181 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 6 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE TABLET ACCORDING TO CLAIM 1 OF U.S. PATENT NO. 11,052,075, WHERE THE TABLET FURTHER COMPRISES IVACAFTOR
- U-3182 METHOD OF PROVIDING POSTSURGICAL PAIN MANAGEMENT, FOR EXAMPLE, VIA INFILTRATION FOR LOCAL ANALGESIA OR VIA INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK FOR REGIONAL ANALGESIA
- U-3183 USE OF EPHEDRINE SULFATE FOR TREATING HYPOTENSION
- U-3184 MEKINIST(R) 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

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COMPLETE RESECTION

- U-3185 TAFINLAR(R) 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-3186 METHOD OF TREATING PRURITUS IN PATIENTS 3 MONTHS OR OLDER SUFFERING FROM PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)
- U-3187 METHOD OF REDUCING SERUM BILE ACIDS IN PATIENTS 3 MONTHS OR OLDER SUFFERING FROM PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)
- U-3188 IMPROVING GLYCEMIC CONTROL IN PATIENTS 10 YEARS OF AGE AND OLDER WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A SUSTAINED-RELEASE EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE
- U-3189 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS 10 TO 17 YEARS OF AGE WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH INSULIN ALONE OR INSULIN PLUS ONE OTHER ORAL ANTIDIABETIC MEDICATION
- U-3190 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS 10 TO 17 YEARS OF AGE WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN AND/OR SULFONYLUREA
- U-3191 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT (30 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-3192 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT (EGFR < 60 ML/MIN/1.73 M²) BY INITIATION OF EMPAGLIFLOZIN, LINAGLIPTIN AND METFORMIN HCL IF EGFR ≥ 45 ML/MIN/1.73 M² AND DISCONTINUATION IF EGFR < 30 ML/MIN/1.73 M²
- U-3193 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATENT WITH RENAL IMPAIRMENT (EGFR < 60 ML/MIN/1.73 M²) BY INITIATION OF EMPAGLIFLOZIN AND METFORMIN HCL IF EGFR ≥ 45 ML/MIN/1.73 M² AND DISCONTINUATION IF EGFR < 30 ML/MIN/1.73 M²
- U-3194 TOPICAL TREATMENT OF ACNE VULGARIS IN ADULTS AND PEDIATRIC PATIENTS 9 YEARS OF AGE AND OLDER
- U-3195 U-1731 U-1732
- U-3196 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA WITH SUSCEPTIBLE FGFR3 OR FGFR2 GENETIC ALTERATIONS, AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY, WITH DOSING BASED ON SERUM PHOSPHATE LEVELS
- U-3197 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE AN OPIOID ANALGESIC AND FOR WHICH ALTERNATIVE TREATMENTS ARE INADEQUATE
- U-3198 METHOD OF TREATING PATIENTS WITH IDIOPATHIC HYPERSOMNIA WITH SODIUM OXYBATE WHEN DIVALPROEX SODIUM IS CONCOMITANTLY ADMINISTERED
- U-3199 METHOD FOR REDUCING THE RISK OF CARDIOVASCULAR DEATH PLUS HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH TYPE 2 DIABETES MELLITUS, HEART FAILURE AND REDUCED EJECTION FRACTION BY ONCE DAILY ADMINISTRATION OF EMPAGLIFLOZIN
- U-3200 INCREASING SURVIVAL IN MCRPC PATIENTS PREVIOUSLY TREATED WITH DOCETAXEL BY ADMINISTERING CABAZITAXEL IN COMBINATION WITH PREDNISONE OR PREDNISOLONE AFTER A PREMEDICATION REGIMEN THAT INCLUDES AN ANTIHISTAMINE, A CORTICOSTEROID, AND AN H₂-ANTAGONIST
- U-3201 TREATMENT OF ADULT PATIENTS WITH VON HIPPEL-LINDAU DISEASE WHO REQUIRE THERAPY FOR ASSOCIATED RENAL CELL CARCINOMA, CENTRAL NERVOUS SYSTEM HEMANGIOBLASTOMAS, OR PANCREATIC NEUROENDOCRINE TUMORS, NOT REQUIRING IMMEDIATE SURGERY
- U-3202 MAINTENANCE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) COMPRISING THE ONCE PER DAY ADMINISTRATION OF TRELEGY ELLIPTA, 100 MCG FLUTICASONE FUROATE/62.5 MCG UMECLIDINIUM/25 MCG VILANTEROL
- U-3203 MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-3204 TREATMENT OF MODERATE-TO-SEVERE PRURITUS ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD-AP) IN ADULTS UNDERGOING HEMODIALYSIS (HD)
- U-3205 REDUCTION OF RISK OF MAJOR CARDIOVASCULAR EVENTS (CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION AND STROKE) IN PATIENTS WITH CAD
- U-3206 REDUCTION OF RISK OF MAJOR THROMBOTIC VASCULAR EVENTS (MYOCARDIAL INFARCTION, ISCHEMIC STROKE, ACUTE LIMB ISCHEMIA, AND MAJOR AMPUTATION OF VASCULAR ETIOLOGY)

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IN PATIENTS WITH PAD

- U-3207 REDUCTION OF RISK OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN PATIENTS WITH CAD BY ADMINISTERING CLINICALLY PROVEN EFFECTIVE AMOUNTS THAT ARE 2.5 MG RIVAROXABAN TWICE DAILY AND 75-100 MG ASPIRIN DAILY
- U-3208 REDUCTION OF RISK OF MYOCARDIAL INFARCTION AND ISCHEMIC STROKE IN PATIENTS WITH PAD BY ADMINISTERING CLINICALLY PROVEN EFFECTIVE AMOUNTS THAT ARE 2.5 MG RIVAROXABAN TWICE DAILY AND 75-100 MG ASPIRIN DAILY
- U-3209 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK MYOCARDIAL INFARCTION IN AN ADULT PATIENT HAVING ATRIAL FIBRILLATION OR ATRIAL FLUTTER AND ELEVATED TRIGLYCERIDE LEVELS
- U-3210 ONCE DAILY TREATMENT OF ANXIETY DISORDER IN ADULTS
- U-3211 TREATING DISTRIBUTIVE SHOCK WITH ANGIOTENSIN II
- U-3212 TREATING SEPTIC SHOCK WITH ANGIOTENSIN II
- U-3213 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHEREIN THE CANCER IS PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA
- U-3214 A METHOD OF TREATING PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 BY ADMINISTERING A ONCE DAILY 500 MG ORAL DOSE TO A SUBJECT THAT HAS NOT INGESTED A HIGH-FAT MEAL
- U-3215 A METHOD OF TREATING PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA CHARACTERIZED BY AN IDH1 MUTATION
- U-3216 A METHOD FOR TREATING A BCRABL POSITIVE LEUKEMIA IN A SUBJECT THAT IS RESISTANT TO IMATINIB COMPRISING ADMINISTERING TO THE SUBJECT A THERAPEUTICALLY EFFECTIVE AMOUNT OF BOSUTINIB, WHEREIN THE SUBJECT HAS A MUTATION IN THE BCRABL PROTEIN AT 949T>C
- U-3217 A METHOD FOR TREATING A BCRABL POSITIVE LEUKEMIA IN A SUBJECT THAT IS RESISTANT TO IMATINIB COMPRISING ADMINISTERING TO THE SUBJECT A THERAPEUTICALLY EFFECTIVE AMOUNT OF BOSUTINIB, WHEREIN THE SUBJECT HAS A MUTATION IN THE BCRABL PROTEIN AT F317L
- U-3218 NASAL ADMINISTRATION OF DIHYDROERGOTAMINE MESYLATE BY METERED SPRAY FOR THE ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA
- U-3219 TREATMENT OF GASTROINTESTINAL STROMAL TUMOR
- U-3220 TREATMENT OF PATIENTS WITH NON-SMALL CELL LUNG CANCER (NSCLC) WITH EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 20 EXON INSERTION MUTATIONS WHOSE DISEASE HAS PROGRESSED ON OR AFTER PLATINUM-BASED CHEMOTHERAPY
- U-3221 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CARDIOVASCULAR EVENT IN A PATIENT WITH PRIOR PERCUTANEOUS CORONARY INTERVENTION
- U-3222 TREATMENT OF ACUTE MYELOID LEUKEMIA (AML) BY ORALLY ADMINISTERING VENETOCLAX WITH AZACITIDINE OR DECITABINE OR LOW-DOSE CYTARABINE IN ADULTS 75 YEARS OR OLDER OR HAVING CERTAIN COMORBIDITIES ACCORDING TO A DOSE RAMP-UP INCLUDING A 100 MG PER DAY DOSE
- U-3223 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) BY ORALLY ADMINISTERING VENETOCLAX TO AN ADULT ACCORDING TO A DOSE RAMP-UP INCLUDING A 100 MG PER DAY DOSE
- U-3224 A METHOD OF TREATING HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE BY DECREASING THE LEVEL OF LDL-C USING A FIXED DOSE COMBINATION OF 180 MG BEMPEDOIC ACID AND 10 MG EZETIMIBE
- U-3225 TREATMENT OF DIFFERENTIATED THYROID CANCER THAT HAS PROGRESSED FOLLOWING PRIOR VEGFR-TARGETED THERAPY
- U-3226 FOR TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR TWO LINES OF SYSTEMIC THERAPY
- U-3227 FOR TREATMENT OF INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS (MF), INCLUDING PRIMARY MF, POST-POLYCYTHEMIA VERA MF AND POST-ESSENTIAL THROMBOCYTHEMIA MF
- U-3228 FOR TREATMENT OF POLYCYTHEMIA VERA (PV) IN PATIENTS WHO HAVE HAD AN INADEQUATE RESPONSE TO OR ARE INTOLERANT OF HYDROXYUREA
- U-3229 FOR TOPICAL SHORT-TERM, NON-CONTINUOUS CHRONIC TREATMENT OF MILD TO MODERATE ATOPIC DERMATITIS IN NON-IMMUNOCOMPROMISED PATIENTS WHOSE DISEASE IS NOT ADEQUATELY CONTROLLED WITH TOPICAL PRESCRIPTION THERAPIES OR WHEN THOSE THERAPIES ARE NOT ADVISABLE

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- U-3230 FOR TREATMENT OF STEROID-REFRACTORY ACUTE GRAFT-VERSUS-HOST DISEASE (AGVHD)
- U-3231 TREATMENT OF PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH+CML) BY ADMINISTERING NILOTINIB DISPERSED IN A FRUIT PREPARATION
- U-3232 USE OF ORAL OCTREOTIDE FOR LONG-TERM MAINTENANCE TREATMENT IN ACROMEGALY PATIENTS WHILE AVOIDING CONCOMITANT ADMINISTRATION OF LEVONORGESTREL
- U-3233 USE FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH TUBEROUS SCLEROSIS COMPLEX IN PATIENTS TAKING EVEROLIMUS
- U-3234 TREATMENT OF MACULAR EDEMA ASSOCIATED WITH UVEITIS
- U-3235 USE FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-3236 USE FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME
- U-3237 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6 IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER OR WEIGHING AT LEAST 45 KG
- U-3238 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6 IN PEDIATRIC PATIENTS 3 TO LESS THAN 12 YEARS OF AGE OR WEIGHING LESS THAN 45 KG
- U-3239 TREATMENT OF ADVANCED RENAL CELL CARCINOMA (RCC) IN PATIENTS WHO HAVE RECEIVED PRIOR ANTI-ANGIOGENIC THERAPY
- U-3240 USE OF VASCEPA TO REDUCE TRIGLYCERIDE LEVELS IN AN ADULT PATIENT HAVING TRIGLYCERIDE LEVELS OF AT LEAST ABOUT 500 MG/DL, ON ANTICOAGULANT/ANTIPLATELET/THROMBOLYTIC THERAPY, AND HAVING ATRIAL FIBRILLATION AND/OR ATRIAL FLUTTER
- U-3241 IN COMBINATION WITH ENDOCRINE THERAPY (TAMOXIFEN OR AN AROMATASE INHIBITOR) FOR THE ADJUVANT TREATMENT OF ADULT PATIENTS WITH HR-POSITIVE, HER2-NEGATIVE, NODE-POSITIVE, EARLY BREAST CANCER AT HIGH RISK OF RECURRENCE AND A KI-67 SCORE $\geq 20\%$
- U-3242 IN COMBINATION WITH FULVESTRANT FOR THE TREATMENT OF ADULT PATIENTS WITH HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY
- U-3243 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN, AND MEN WITH HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-3244 A METHOD FOR TREATMENT OF PAIN IN ADULTS USING TRAMADOL HYDROCHLORIDE AND CELECOXIB
- U-3245 MAINTENANCE MONOTHERAPY TREATMENT OF BIPOLAR I DISORDER
- U-3246 FOR THE TREATMENT OF CHRONIC GRAFT VERSUS HOST DISEASE
- U-3247 FOR THE TREATMENT OF CHRONIC GRAFT VERSUS HOLD DISEASE
- U-3248 TREATING SECONDARY HYPERPARATHYROIDISM IN STAGE 3/4 CHRONIC KIDNEY DISEASE WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL WHILE AVOIDING PTH OVERSUPPRESSION
- U-3250 METHOD OF TREATING PAIN, FOR EXAMPLE, TREATING POSTSURGICAL PAIN VIA INFILTRATION FOR LOCAL ANALGESIA OR VIA INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK FOR REGIONAL ANALGESIA
- U-3251 ADMINISTRATION OF FERROUS BISGLYCINATE TABLETS
- U-3252 USE OF VUITY FOR THE TREATMENT OF PRESBYOPIA IN ADULTS
- U-3253 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOSITION ACCORDING TO AT LEAST ONE OF CLAIMS 1-9 OF US11179367
- U-3254 USE, 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
- U-3255 TREATMENT OF ADULTS WITH MODERATELY TO SEVERELY ACTIVE RHEUMATOID ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3256 USE TO INCREASE LINEAR GROWTH IN PEDIATRIC PATIENTS WITH ACHONDROPLASIA WHO ARE 5 YEARS OF AGE AND OLDER WITH OPEN EPIPHYSES
- U-3257 TREATMENT OF TRD IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE IN A MAINTENANCE PHASE WEEKLY OR 1X EVERY TWO WEEKS TO ADULTS WHO HAVE BEEN ADMINISTERED ESKETAMINE IN A INDUCTION

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PHASE FOR ABOUT 4 WEEKS

- U-3258 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC MALIGNANT PERIVASCULAR EPITHELIOID CELL TUMOR (PECOMA) WITH A DOSE BETWEEN ABOUT 56 MG/M2 AND ABOUT 100 MG/M2 ADMINISTERED ON DAYS 1 AND 8 OF A 21-DAY CYCLE
- U-3259 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC MALIGNANT PERIVASCULAR EPITHELIOID CELL TUMOR (PECOMA)
- U-3260 METHOD OF TREATING SPASTICITY
- U-3261 FOR CLEANSING OF THE COLON AS A PREPARATION FOR COLONOSCOPY
- U-3262 TREATING HYPOTENSION WITH ANGIOTENSIN II IN A PATIENT RECEIVING AN ANGIOTENSIN CONVERTING ENZYME INHIBITOR
- U-3263 METHOD FOR TREATING SPASTICITY
- U-3264 AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF ADULT PATIENTS WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-3265 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF ADULT PATIENTS WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-3266 IN COMBINATION WITH FULVESTRANT AS INITIAL ENDOCRINE-BASED THERAPY OR FOLLOWING DISEASE PROGRESSION ON ENDOCRINE THERAPY IN POSTMENOPAUSAL WOMEN OR IN MEN, FOR THE TREATMENT OF HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-3267 USE OF LASTACFT TO TEMPORARY RELIEVE ITCHY EYES DUE TO POLLEN, RAGWEED, GRASS, ANIMAL HAIR AND DANDER
- U-3268 TREATMENT OF MULTIPLE SCLEROSIS IN PEDIATRIC PATIENTS 10 YEARS OF AGE AND OLDER AND WEIGHING LESS THAN OR EQUAL TO 40 KG
- U-3269 TREATMENT OF PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) IN ADULTS AT RISK OF RAPID DISEASE PROGRESSION
- U-3270 U-2106: TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT COMCOMITANT DOPAMINERGIC MEDICATIONS
- U-3271 TREATMENT OF BIPOLAR DEPRESSION
- U-3272 AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY FOR THE TREATMENT OF ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH) OR CLINICAL ATHEROSCLEROTIC CARDIOVASCULAR DISEASE (ASCVD), BY INHIBITING EXPRESSION OF THE PCSK9 GENE
- U-3273 AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY FOR THE TREATMENT OF ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH) OR CLINICAL ATHEROSCLEROTIC CARDIOVASCULAR DISEASE (ASCVD), BY INHIBITING EXPRESSION OF THE PCSK9 GEN
- U-3274 TREATMENT OF BIPOLAR I DISORDER, BIPOLAR II DISORDER, OR BIPOLAR DEPRESSION
- U-3275 TREATMENT OF ADULTS WITH ACTIVE PSORIATIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS