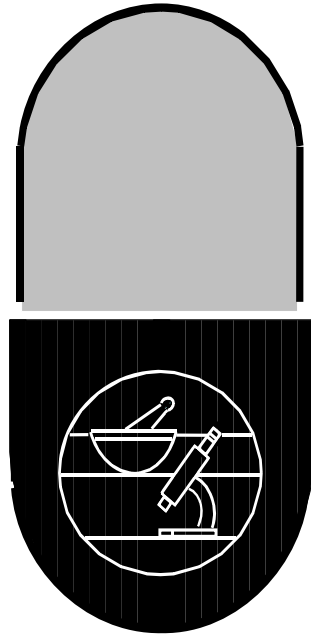


**CUMULATIVE
SUPPLEMENT 09
September 2006**



**APPROVED
DRUG PRODUCTS**

**WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS**

26th EDITION

Department of Health and Human Services

**Food and Drug Administration
Center for Drug Evaluation and Research
Office of Generic Drugs**

2006

Prepared By
Office of Generic Drugs
Center for Drug Evaluation and Research
Food and Drug Administration

**APPROVED DRUG PRODUCTS
with
THERAPEUTIC EQUIVALENCE EVALUATIONS**

26th EDITION

Cumulative Supplement 09

September 2006

CONTENTS

| | <i>PAGE</i> |
|--|-------------|
| 1.0 INTRODUCTION | iii |
| 1.1 How to use the Cumulative Supplement | iii |
| 1.2 Cumulative Supplement Content | iv |
| 1.3 Applicant Name Changes..... | v |
| 1.4 Availability of the Edition | vi |
| 1.5 Report of Counts for the Prescription Drug Product List | vii |
| 1.6 Zocor (simvastatin) patent relisting | viii |
| 1.7 Levothyroxine Sodium..... | viii |
| 1.8 Cumulative Supplement Legend | x |
| | |
| DRUG PRODUCT LISTS | |
| Prescription Drug Product List | 1-1 |
| OTC Drug Product List | 2-1 |
| Drug Products with Approval under Section 505 of the Act | |
| Administered by the Center for Biologics Evaluation and Research List..... | 3-1 |
| Orphan Product Designations and Approvals List | 4-1 |
| Drug Products Which Must Demonstrate in vivo Bioavailability | |
| Only if Product Fails to Achieve Adequate Dissolution | 5-1 |
| | |
| PATENT AND EXCLUSIVITY INFORMATION ADDENDUM | |
| A. Patent and Exclusivity Lists | A-1 |
| B. Patent and Exclusivity Terms | B-1 |

APPROVED DRUG PRODUCTS
with
THERAPEUTIC EQUIVALENCE EVALUATIONS

26th EDITION

CUMULATIVE SUPPLEMENT 09
September 2006

1.0 INTRODUCTION

1.1 HOW TO USE THE CUMULATIVE SUPPLEMENT

This Cumulative Supplement is one of a series of monthly updates to the Approved Drug Products with Therapeutic Equivalence Evaluations, 25th Edition (the List). The List is composed of four parts: approved prescription drug products with therapeutic equivalence evaluations; over-the-counter (OTC) drug products that require approved applications as a condition of marketing; drug products with approval under Section 505 of the Act administered by the Center for Biologics Evaluation and Research; and products that have never been marketed, are for exportation, are for military use, have been discontinued from marketing or that have had their approvals withdrawn for other than safety or efficacy reasons.

The Cumulative Supplement provides, among other things, information on newly approved drugs and, if necessary, revised therapeutic equivalence evaluations and updated patent and exclusivity data. The Addendum contains appropriate drug patent and exclusivity information required of the Agency by the "Drug Price Competition and Patent Term Restoration Act of 1984" for the Prescription, OTC, and Drug Products with Approval under Section 505 of the Act Administered by the Center for Biologics Evaluation and Research Lists.

Because all parts of the publication are subject to changes, additions, or deletions, the List must be used in conjunction with the most current Cumulative Supplement. Users may wish to mark to the left of the ingredient(s) in the List to indicate that changes to that entry appear in the Cumulative Supplement. Drug product information is provided in each Cumulative Supplement for completeness to assist in locating the proper place in the List for the revision.

The presence of any therapeutic equivalence code indicates that the drug product is multisource; the deletion of a therapeutic equivalence code indicates that the drug product has become single source. (An infrequent exception exists when a therapeutic equivalence code is revised. In that case the deletion of the therapeutic equivalence code is followed immediately by the addition of the revised one.)

Products that have never been marketed, are for exportation, are for military use, or have been discontinued from marketing or that have had their approvals withdrawn for other than safety or efficacy reasons, will be flagged in this Cumulative Supplement with the "@" symbol to designate their non-marketed status. All products having a "@" symbol in the 12th Cumulative Supplement of the 25th Edition List will then be added to the "Discontinued Drug Product List" appearing in the 26th Edition. The current edition

Section 2. How To Use The Drug Product Lists describes the layout and usage of the List.

New additions to the Prescription Drug Product List and OTC Drug Product List are indicated by the symbol >A>. The Patent and Exclusivity List new additions are indicated by the symbol >A> to the left of Patent Number or Exclusivity Code. The >A> symbol is then dropped in subsequent Cumulative Supplements for that item.

New deletions to the Prescription Drug Product List and OTC Drug Product List are indicated by the symbol >D> (DELETE) to the left of the line. The information line with the >D> symbol is dropped in subsequent Cumulative Supplements for that item.

The Patent and Exclusivity List is arranged in alphabetical order by active ingredient name(s) and trade name. The trade name will follow the active ingredient name separated by a dash symbol. Also shown is the application number and product number (FDA's internal file number) for reference purposes. All patents with their expiration dates are displayed for each application number. Drug substance and drug product patents are indicated as such with DS or DP in the Patent codes column. Use patents are indicated with the symbol "U" followed by a number representing a specific use. Exclusivity information for a specific drug is indicated by an abbreviation followed by the date upon which the exclusivity expires. Refer to the Exclusivity Terms, Section B, in the Patent and Exclusivity Information Addendum for an explanation of all codes and abbreviations. Refer to Section 1.3 for internet access to the most current list of Patent and Exclusivity terms.

1.2 CUMULATIVE SUPPLEMENT CONTENT

Since February 2005, we have been providing daily Electronic Orange Book (EOB) product information for new generic drug approvals. Daily generic updates provide the consumer with the current list of approved generic products which is important for substitution purposes. Previously, a first-time-generic product approved early in the month would not be published in the Cumulative Supplement (CS) for several weeks.

The CS monthly update publish goal is by the end of the following month's second work week (e.g., November's supplement will be updated by the end of the second full work week in December).

Currently, the monthly PDF CS includes:

- Generic product ANDA (Abbreviated New Drug Approval) approvals as of the date of publication.
- We also include all product changes received and processed as of the date of publication.
 - Refer to CS Section 1.8 Cumulative Supplement Legend for types of changes
 - Discontinued products will be processed as of the date of publication. There will be circumstances where a product is discontinued in one month, however, it will be reported in a different month's CS. For example, the Orange Book received a letter November 7 that the product has been discontinued from manufacturing and marketing. The Orange Books subsequently publishes the October CS on November 14. The product will show in the October CS that it is discontinued even though the date of discontinuance is the day that the Orange Book receives notification (November 7).

- New Drug Application (NDA) approvals (20,000 and 50,000 series) appear in the CS month they were approved.
- Patent information, also updated daily in the EOB, is current to the date of publication.
- Exclusivity information is updated monthly and current to the date of publication.

1.3 APPLICANT NAME CHANGES

It is not practical to identify in the Cumulative Supplement each and every product involved when an applicant transfers its entire line of approved drug products to another applicant, or when an applicant changes its name. Therefore, the cumulation of these transfers and name changes will be identified in this section only. Where only partial lines of approved products are transferred between applicants, each approved product involved will appear as an applicant name change entry in the Cumulative Supplement.

It is also not practical to identify each and every product involved when an applicant name is changed to meet internal publication standards (e.g., MSD or Zenith [Former Abbreviated Names] are changed, respectively, to Merck Sharp Dohme or Zenith Labs [New Abbreviated Names]). When this occurs, each product involved (either currently in the Cumulative Supplement or in the following year's edition) will reflect the new abbreviated name. Consequently, it will not appear as an applicant name change entry in the Cumulative Supplement nor will the cumulation of these name changes appear in this section. The Electronic Orange Book Query, updated monthly, will contain the most current applicant holder name.

| <u>FORMER APPLICANT NAME</u> <u>(FORMER ABBREVIATED NAME)</u> | <u>NEW APPLICANT NAME</u> <u>(NEW ABBREVIATED NAME)</u> |
|--|--|
| AMERICAN PHARMACEUTICAL PARTNERS INC (AM PHARM PARTNERS) | ABRAXIS PHARMACEUTICAL PRODUCTS (ABRAXIS PHARM) |
| AMERICAN PHARMACEUTICAL CO INC SUB BURR CORP (AM PHARM) | ABRAXIS PHARMACEUTICAL PRODUCTS (ABRAXIS PHARM) |
| AMIDE PHARMACEUTICAL INC (AMIDE PHARM) | ACTAVIS TOTOWA LLC (ACTAVIS TOTOWA) |
| APOTEX CORP (APOTEX) | APOTEX INC ETOBICOKE SITE (APOTEX INC) |
| APOTEX CORP (APOTEX) | APOTEX INC RICHMOND HILL (APOTEX INC) |
| APOTEX INC (APOTEX) | APOTEX INC ETOBICOKE SITE (APOTEX INC) |
| APOTEX INC (APOTEX) | APOTEX INC RICHMOND HILL (APOTEX INC) |
| APOTEX INC (APOTEX INC) | APOTEX INC ETOBICOKE SITE (APOTEX INC) |
| APOTEX INC (APOTEX INC) | APOTEX INC RICHMOND HILL (APOTEX INC) |
| AVENTIS PHARMACEUTICALS INC (AVENTIS) | SANOFI AVENTIS US LLC (SANOFI AVENTIS US) |
| AVENTIS PHARMACEUTICAL PRODUCTS INC (AVENTIS PHARMS) | SANOFI AVENTIS US LLC (SANOFI AVENTIS US) |
| CLAY PARK LABORATORIES INC (CLAY PARK) | PERRIGO NEW YORK INC (PERRIGO NEW YORK) |

| | |
|--|---|
| CLAY PARK LABS INC (CLAY PARK) | PERRIGO NEW YORK INC (PERRIGO NEW YORK) |
| DERMIK LABORATORIES INC (DERMIK LABS) | SANOFI AVENTIS US LLC (SANOFI AVENTIS US) |
| DERMIK LABORATORIES INC SUB RORER (DERMIK LABS) | SANOFI AVENTIS US LLC (SANOFI AVENTIS US) |
| FIRST HORIZON PHARMACEUTICAL COMPANY (FIRST HORIZON) | SCIELE PHARMA INC (SCIELE PHARMA INC) |
| LOREX PHARMACEUTICALS (LOREX) | SANOFI AVENTIS US LLC (SANOFI AVENTIS US) |
| MARTEC PHARMACEUTICALS (MARTEC) | MARTEC USA LLC (MARTEC USA LLC) |
| MARTEC SCIENTIFIC INC (MARTEC) | MARTEC USA LLC (MARTEC USA LLC) |
| MCNEIL CONSUMER AND SPECIALTY PHARMACEUTICALS DIV MCNEIL PCC IN (MCNEIL CONS SPECLT) | MCNEIL PEDIATRICS (MCNEIL PED) |
| ORPHAN MEDICAL INC (ORPHAN MEDCL) | JAZZ PHARMACEUTICALS (JAZZ) |
| PHARMACEUTICAL FORMULATIONS INC (PHARM FORM) | LEINER HEALTH PRODUCTS INC (LEINER HLTH PRODS) |
| PHARMA TEK INC (PHARMA TEK) | X GEN PHARMACEUTICALS INC (X GEN) |
| PRIVATE FORMULATIONS INC (PRIVATE FMLTNS) | LEINER HEALTH PRODUCTS INC (LEINER HLTH PRODS) |
| SANKYO PHARMA INC (SANKYO) | DAIICHI SANKYO INC (DAIICHI SANKYO) |
| SANOFI AVENTIS US INC (SANOFI AVENTIS US) | SANOFI AVENTIS US LLC (SANOFI AVENTIS US) |
| SANOFI-AVENTIS US INC (SANOFI AVENTIS) | SANOFI AVENTIS US LLC (SANOFI AVENTIS US) |
| SANOFI INC (SANOFI) | SANOFI AVENTIS US LLC (SANOFI AVENTIS US) |
| SANOFI SYNTHELABO INC (SANOFI SYNTHELABO) | SANOFI AVENTIS US LLC (SANOFI AVENTIS US) |
| SANOFI SYNTHELABO RESEARCH DIV SANOFI SYNTHELABO INC (SANOFI SYN RES) | SANOFI AVENTIS US LLC (SANOFI AVENTIS US) |
| STERIS LABORATORIES INC (STERIS) | WATSON LABORATORIES INC (WATSON) |
| TRIGEN LABORATORIES INC (TRIGEN) | JUBILANT PHARMACEUTICALS INC (JUBILANT PHARMS) |
| UCB PHARMA INC (UCB PHARMA) | UCB INC (UCB INC) |

1.4 AVAILABILITY OF THE EDITION

Commencing with the 25th edition, the Annual Edition and monthly Cumulative Supplements will not be available in a published paper version.

Since 1997, the Electronic Orange Book (EOB <http://www.fda.gov/cder/ob/default.htm>), has been available on the internet and has become the updated-every-month Orange Book.

The 25th edition and current monthly supplement are available in an electronic downloadable Portable Document Format (PDF) at the EOB home page by clicking on the Annual Edition. The PDF annual and cumulative supplements will duplicate previous paper versions. Over time, there will be an archive for the annuals and each year's December Cumulative Supplement.

The Electronic Orange Book Query (EOB) is at <http://www.fda.gov/cder/ob/default.htm>. The Query provides searching of the approved drug list by active ingredient, proprietary name, applicant

holder, applicant number or patent number. Product search categories are: prescription, over-the-counter, discontinued drugs. There are links to patent and exclusivity information that may be applicable to each product. Currently, In addition to monthly updates, in the public interest, the EOB is updated on a daily basis with new generic product approval information and new patent information. Current month updates are accomplished by the third week of the following month.

There are historical lists of Orange Book cumulative supplement product monthly changes at <http://www.fda.gov/cder/rxotcdpl/pdplarchive.htm>

There are ASCII text files of the Orange Book drug product, patent, and exclusivity data at <http://www.fda.gov/cder/orange/obreadme.htm>. The drug product text files are zipped into eobzip.exe. The files are updated concurrently with the monthly cumulative supplements. Appendix A and Appendix B text files of the annual Orange Book Edition are updated quarterly.

Effective August 18, 2003, patent submissions for publication in the Orange Book and Docket *95S-0117 need to be submitted on form FDA-3542 which may be downloaded from the FDA Forms List, <http://www.fda.gov/opacom/morechoices/fdaforms/default.html>.

The current listing of the Orphan Product Designations and Approvals is available at <http://www.fda.gov/orphan/designat/list.htm>.

1.5 REPORT OF COUNTS FOR THE PRESCRIPTION DRUG PRODUCT LIST

DESCRIPTION OF REPORT

This report provides summary counts derived from the product information in the Prescription Drug Product List and the current Cumulative Supplement. Products included in the counts are domestically marketed drug products approved for both safety and effectiveness under section 505 of the Federal Food, Drug, and Cosmetic Act. Excluded are approved drug products marketed by distributors; those marketed solely abroad; and those now regarded as medical devices, biologics or foods.

The baseline column (Dec 2004) refers to the products in the Prescription Drug Product List. For each three-month period, a column of quarterly data is added which incorporates counts of product activity from the previous quarter(s) with those in the baseline count.

DEFINITIONS

Drug Product

For this report, a drug product is the representation in the Prescription Drug Product List of an active moiety (molecular entity and its salts, esters and derivatives) either as a single ingredient or as a combination product provided in a specific dosage form and strength for a given route of administration with approval for marketing by a firm under a particular generic or trade name.

New Molecular Entity

A new molecular entity is considered an active moiety that has not previously been approved (either as the parent compound or as a salt, ester or derivative of the parent compound) in the United States for use in a drug product either as a single ingredient or as part of a

combination.

REPORT OF COUNTS FOR THE PRESCRIPTION DRUG PRODUCT LIST
COUNTS CUMULATIVE BY QUARTER

| <u>CATEGORIES COUNTED</u> | <u>DEC 2005</u> | <u>MAR 2006</u> | <u>JUN 2006</u> | <u>SEPT 2006</u> |
|------------------------------------|------------------|-----------------|-----------------|------------------|
| DRUG PRODUCTS LISTED | 11368 | 11487 | 11636 | |
| SINGLE SOURCE | 2428 (21.4%) | 2461 (21.4%) | 2447 (21.0%) | |
| MULTISOURCE | 8851 (77.9%) | 8937 (77.8%) | 9000 (78.2%) | |
| THERAPEUTICALLY EQUIVALENT | 8642 (76.04%) | 8730 (76.0%) | 8900 (76.5%) | |
| NOT THERAPEUTICALLY EQUIVALENT | 209 (1.8%) | 207 (1.8%) | 200 (1.7%) | |
| EXCEPTIONS ¹ | 89 (0.8%) | 89 (0.8%) | 89 (0.8%) | |
| NEW MOLECULAR ENTITIES APPROVED | 11 | 6 | 8 | |
| NUMBER OF APPLICANTS | 628 | 629 | 642 | |

¹Amino acid-containing products of varying composition (see Introduction, page xx of the List).

1.6 ZOCOR (SIMVASTATIN) PATENT RELISTING

U.S. Patent Nos. RE 36481 and RE 36520 are being relisted for Zocor (NDA 19-766) pursuant to the decision and related order in *Ranbaxy Labs. v. Leavitt*, No. 05-1838 (D.D.C. April 30, 2006). The '481 and '520 patents will remain listed in *Approved Drug Products with Therapeutic Equivalence Evaluations* until any applicable periods of exclusivity pursuant to section 505(j)(5)(B)(iv) of the Federal Food, Drug, and Cosmetic Act have been triggered and run, unless the agency's appeal of the decision to the U.S. Court of Appeals for the District of Columbia is decided in the agency's favor before the exclusivity periods have expired. While the patents remain listed, any new or pending ANDA referencing Zocor must contain patent certifications to these patents. For additional information on this matter, please refer to Docket Nos. 2005P-0008 and 2005P-0046.

1.7 LEVOTHYROXINE SODIUM

The Description of Special Situations, Levothyroxine Sodium, published in the 26th Annual Edition of the Orange Book, has been modified in the Cumulative Supplement to include information on a supplemental approval for Genpharm ANDA 76752 approved in 2006. The full discussion as published in the 26th Annual Edition is repeated in the Cumulative Supplement and includes recent approval information on Levothyroxine Sodium.

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

Levothyroxine Sodium (Mylan ANDA 76187) tablets have been determined to be therapeutically equivalent to corresponding strengths of Unithroid (Jerome Stevens NDA 021210) tablets.

Levo-T (Alara NDA 021342), Levothyroxine Sodium (Mylan ANDA 76187), Unithroid (Jerome Stevens NDA 021210) and Levothyroxine Sodium (Genpharm ANDA 76752) tablets have been determined to be therapeutically equivalent to corresponding strengths of Synthroid (Abbott NDA 021402) tablets.

Levo-T (Alara NDA 021342), Unithroid (Jerome Stevens NDA 021210), Levothyroxine Sodium (Mylan ANDA 076187) and Levothyroxine Sodium (Genpharm ANDA 76752) tablets have been determined to be therapeutically equivalent to corresponding strengths of Levoxyl (King/Jones Pharma NDA 021301) tablets.

Novothyrox (Genpharm NDA 021292) requires further investigation and review to establish therapeutic equivalence to corresponding strengths of any other Levothyroxine Sodium drug products and is rated BX.

Thyro-Tabs (Lloyd NDA 021116) requires further investigation and review to establish therapeutic equivalence to corresponding strengths of any other Levothyroxine Sodium drug products and is rated BX.

Levolet (Vintage NDA 021137) requires further investigation and review to establish therapeutic equivalence to corresponding strengths of any other Levothyroxine Sodium drug products and is rated BX.

The chart outlines TE codes for all 0.025mg products with other products being similar. Therapeutic equivalence has been established between products that have the same AB+number TE code. More than one TE code may apply to some products. One common TE code indicates therapeutic equivalence between products.

| Trade Name | Applicant | Potency | TE Code | Appl No | Product No |
|-----------------------|--------------|---------|---------|---------|------------|
| UNITHROID | STEVENS J | 0.025MG | AB1 | 21210 | 001 |
| LEVOTHYROXINE SODIUM | MYLAN | 0.025MG | AB1 | 76187 | 001 |
| LEVOXYL | JONES PHARMA | 0.025MG | AB1 | 21301 | 001 |
| SYNTHROID | ABBOTT | 0.025MG | AB1 | 21402 | 001 |
| | | | | | |
| SYNTHROID | ABBOTT | 0.025MG | AB2 | 21402 | 001 |
| LEVOTHYROXINE SODIUM | MYLAN | 0.025MG | AB2 | 76187 | 001 |
| LEVO-T | ALARA PHARM | 0.025MG | AB2 | 21342 | 001 |
| UNITHROID | STEVENS J | 0.025MG | AB2 | 21210 | 001 |
| LEVOTHYROXINE SODIUM | GENPHARM | 0.025MG | AB2 | 76752 | 001 |
| | | | | | |
| LEVOXYL | JONES PHARMA | 0.025MG | AB3 | 21301 | 001 |
| LEVO-T | ALARA PHARM | 0.025MG | AB3 | 21342 | 001 |
| UNITHROID | STEVENS J | 0.025MG | AB3 | 21210 | 001 |
| LEVOTHYROXINE SODIUM | MYLAN | 0.025MG | AB3 | 76187 | 001 |
| LEVOTHYROXINE SODIUM* | GENPHARM | 0.025MG | AB3 | 76752 | 001 |
| | | | | | |
| NOVOTHYROX | GENPHARM | 0.025MG | BX | 21292 | 001 |
| | | | | | |
| THYRO-TABS | LLOYD | 0.025MG | BX | 21116 | 001 |
| | | | | | |
| LEVOLET | VINTAGE | 0.025MG | BX | 21137 | 001 |

| Trade Name | Applicant | Potency | TE Code | Appl No | Product No |
|------------|-----------|---------|---------|---------|------------|
| | PHARMS | | | | |

*Revised September 2006

1.8 CUMULATIVE SUPPLEMENT LEGEND

The List is sorted by Ingredient(s) and, within each grouping, by the Dosage Form; Route and then by trade name.

The individual product record contains the Therapeutic Equivalence Code, Reference Listed Drug symbol, applicant holder, strength(s), New Drug Application number, product number, and approval date. The last two columns describe the action. The Action Month is the CS month the action occurred. The OB Action is the type of change that has occurred.

New ingredient(s), new dosage form; route(s), new trade names, and new product additions are preceded by >A> during the action month. The change month is the current CS month; the change code for new approvals is NEWA. Following months will display the same information without the >A>.

Changes to currently listed products will list two records. The deleted product record will be preceded by >D>. The product record change addition being made will be preceded by >A>. Following months will display only the >A> record without the >A>. All changes that occur to the product through the Annual year will be listed. The change month and change code will document the change.

The change code and description:

| | |
|------|---|
| NEWA | New drug product approval usually in the supplement month. |
| CAHN | Applicant holder firm name has changed. |
| CAIN | Change. There has been a change in the Ingredient(s) name. All products will be deleted under the old name and all products will be added under the changed ingredient(s) name. |
| CDFR | Change. Dosage Form; Route of Administration. |
| CFTG | Change. A first time generic for the innovator product. A TE Code is added. |
| CMFD | Change. The product is moved from the Discontinued Section due to a change in marketing status. |
| CMS1 | Change. Miscellaneous addition to list. |
| CMS2 | Change. Miscellaneous deletion from list. |
| CPOT | Change. Potency amount/unit. |
| CRLD | Change. Reference Listed Drug. |
| CTEC | Change. Therapeutic Equivalence Code. |
| CTNA | Change. Trade Name. |
| DISC | Discontinued. The Rx or OTC listed product is not being marketed and will be moved to the discontinued section in the next edition. |

PRESCRIPTION DRUG PRODUCT LIST - 26TH EDITION

RX DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 9 - September 2006

1-1

| | | | | | | | | | |
|-----|---|---|--------------------|--------|-----|--------------|-----|------|--|
| >D> | <u>ABARELIX</u> | | | | | | | | |
| >D> | INJECTABLE; INTRAMUSCULAR | | | | | | | | |
| >D> | PLENAXIS | | | | | | | | |
| >D> | + | PRAECIS | 100MG/VIAL | N21320 | 001 | Nov 25, 2003 | Sep | DISC | |
| >A> | | @ | 100MG/VIAL | N21320 | 001 | Nov 25, 2003 | Sep | DISC | |
| | <u>ACEBUTOLOL HYDROCHLORIDE</u> | | | | | | | | |
| | CAPSULE; ORAL | | | | | | | | |
| | SECTRAL | | | | | | | | |
| AB | | DR REDDYS LABS INC | EQ 200MG BASE | N18917 | 001 | Dec 28, 1984 | May | CAHN | |
| AB | + | | EQ 400MG BASE | N18917 | 003 | Dec 28, 1984 | May | CAHN | |
| | <u>ACETAMINOPHEN; BUTALBITAL</u> | | | | | | | | |
| | TABLET; ORAL | | | | | | | | |
| | BUTAPAP | | | | | | | | |
| AB | + | MIKART | 650MG;50MG | N89988 | 001 | Oct 26, 1992 | Jan | CRLD | |
| | | SEDAPAP | | | | | | | |
| | | @ MAYRAND | 650MG;50MG | N88944 | 001 | Oct 17, 1985 | Jan | DISC | |
| | <u>ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE</u> | | | | | | | | |
| | TABLET; ORAL | | | | | | | | |
| | ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE | | | | | | | | |
| >D> | + | MIKART | 712.8MG;60MG;32MG | N40316 | 001 | Apr 28, 1999 | Sep | CFTG | |
| >A> | AB | + | 712.8MG;60MG;32MG | N40316 | 001 | Apr 28, 1999 | Sep | CFTG | |
| | | ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITATRATE | | | | | | | |
| >A> | AB | WEST WARD | 712.8MG;60MG;32MG | N40637 | 001 | Sep 22, 2006 | Sep | NEWA | |
| | <u>ACETAMINOPHEN; CODEINE PHOSPHATE</u> | | | | | | | | |
| | SOLUTION; ORAL | | | | | | | | |
| | ACETAMINOPHEN AND CODEINE PHOSPHATE | | | | | | | | |
| AA | + | ACTAVIS MID ATLANTIC | 120MG/5ML;12MG/5ML | N85861 | 001 | | Jul | CAHN | |
| | | @ CLONMEL | 120MG/5ML;12MG/5ML | N40098 | 001 | Sep 20, 1996 | Jan | DISC | |
| | SUSPENSION; ORAL | | | | | | | | |
| | CAPITAL AND CODEINE | | | | | | | | |
| AA | | ACTAVIS MID ATLANTIC | 120MG/5ML;12MG/5ML | N85883 | 001 | | Jul | CAHN | |
| | TABLET; ORAL | | | | | | | | |
| | ACETAMINOPHEN AND CODEINE PHOSPHATE | | | | | | | | |
| AA | + | TEVA | 300MG;60MG | N88629 | 001 | Mar 06, 1985 | Apr | CRLD | |
| | | ACETAMINOPHEN W/ CODEINE NO. 3 | | | | | | | |
| | | @ ROXANE | 300MG;30MG | N84656 | 001 | | Jul | DISC | |
| | <u>ACETAMINOPHEN; HYDROCODONE BITARTRATE</u> | | | | | | | | |
| | TABLET; ORAL | | | | | | | | |
| | HYDROCODONE BITARTRATE AND ACETAMINOPHEN | | | | | | | | |
| | | @ ENDO PHARMS | 500MG;7.5MG | N40280 | 001 | Sep 30, 1998 | Feb | DISC | |
| | | @ | 650MG;7.5MG | N40280 | 002 | Sep 30, 1998 | Feb | DISC | |
| | | @ | 650MG;10MG | N40280 | 003 | Sep 30, 1998 | Feb | DISC | |
| | | @ | 750MG;7.5MG | N40281 | 002 | Sep 30, 1998 | Feb | DISC | |
| AA | | INTERPHARM | 325MG;5MG | N40736 | 001 | Aug 25, 2006 | Aug | NEWA | |
| AA | | | 325MG;10MG | N40746 | 001 | Aug 25, 2006 | Aug | NEWA | |
| AA | | | 500MG;5MG | N40729 | 001 | Aug 25, 2006 | Aug | NEWA | |
| AA | | | 500MG;7.5MG | N40748 | 001 | Aug 25, 2006 | Aug | NEWA | |
| AA | | | 650MG;7.5MG | N40754 | 001 | Aug 25, 2006 | Aug | NEWA | |

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

| | | | | | | | | |
|----|----------------|-------------|-------------|--------|--------------|--------------|------|------|
| AA | INTERPHARM | 650MG;10MG | N40757 | 001 | Aug 25, 2006 | Aug | NEWA | |
| AA | | 750MG;7.5MG | N40769 | 001 | Aug 28, 2006 | Aug | NEWA | |
| | MIKART | 300MG;5MG | N40658 | 001 | Jan 19, 2006 | Jan | NEWA | |
| | + | 300MG;7.5MG | N40556 | 002 | Mar 24, 2006 | Mar | NEWA | |
| AA | VINTAGE PHARMS | 325MG;5MG | N40655 | 001 | Jan 19, 2006 | Jan | NEWA | |
| AA | | 325MG;7.5MG | N40656 | 001 | Jan 19, 2006 | Jan | NEWA | |
| | HY-PHEN | | | | | | | |
| | @ ASCHER | 500MG;5MG | N87677 | 001 | May 03, 1982 | Mar | DISC | |
| | ZYDONE | | | | | | | |
| | + | ENDO PHARMS | 400MG;5MG | N40288 | 001 | Nov 27, 1998 | May | CTNA |
| | + | | 400MG;7.5MG | N40288 | 002 | Nov 27, 1998 | May | CTNA |
| | + | | 400MG;10MG | N40288 | 003 | Nov 27, 1998 | May | CTNA |

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

SOLUTION; ORAL

>A> OXYCODONE AND ACETAMINOPHEN

| | | | | | | | | |
|-----|----|--------------|-------------------|--------|-----|--------------|-----|------|
| >A> | AA | MALLINCKRODT | 325MG/5ML;5MG/5ML | N40680 | 001 | Sep 29, 2006 | Sep | NEWA |
| | | ROXICET | | | | | | |
| >D> | + | ROXANE | 325MG/5ML;5MG/5ML | N89351 | 001 | Dec 03, 1986 | Sep | CFTG |
| >A> | AA | + | 325MG/5ML;5MG/5ML | N89351 | 001 | Dec 03, 1986 | Sep | CFTG |

TABLET; ORAL

OXYCODONE AND ACETAMINOPHEN

| | | | | | | | | |
|--|---|--------|-------------|--------|-----|--------------|-----|------|
| | + | MIKART | 400MG;2.5MG | N40679 | 001 | May 16, 2006 | May | NEWA |
| | + | | 400MG;5MG | N40687 | 001 | Apr 27, 2006 | Apr | NEWA |
| | + | | 400MG;7.5MG | N40698 | 001 | Apr 27, 2006 | Apr | NEWA |
| | + | | 400MG;10MG | N40692 | 001 | Apr 27, 2006 | Apr | NEWA |
| | | | 500MG;10MG | N40676 | 001 | Apr 19, 2006 | Apr | NEWA |

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET; ORAL

PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN

| | | | | | | | |
|----|-------------------|-------------|--------|-----|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 650MG;100MG | N70910 | 001 | Jan 02, 1987 | Jun | CAHN |
| | CORNERSTONE | 325MG;100MG | N76743 | 001 | May 07, 2004 | Jul | CAHN |
| AB | | 500MG;100MG | N76750 | 001 | Jun 28, 2004 | Jul | CAHN |

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

| | | | | | | | |
|----|------|--------------|--------|-----|--------------|-----|------|
| AB | BARR | 325MG;37.5MG | N76914 | 001 | Jul 26, 2006 | Jul | NEWA |
|----|------|--------------|--------|-----|--------------|-----|------|

ACETAZOLAMIDE

TABLET; ORAL

ACETAZOLAMIDE

| | | | | | | | | |
|----|---|-----------------------|-------|--------|-----|--------------|-----|------|
| AB | + | TARO | 250MG | N40195 | 002 | May 28, 1997 | Mar | CRLD |
| | | DIAMOX | | | | | | |
| | | @ DURAMED PHARMS BARR | 125MG | N08943 | 001 | | Mar | DISC |
| | | @ | 250MG | N08943 | 002 | | Mar | DISC |

ACETIC ACID, GLACIAL

SOLUTION/DROPS; OTIC

ACETASOL

| | | | | | | | |
|----|----------------------|----|--------|-----|--|-----|------|
| AT | ACTAVIS MID ATLANTIC | 2% | N87146 | 001 | | Jul | CAHN |
|----|----------------------|----|--------|-----|--|-----|------|

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS; OTIC

ACETASOL HC

| | | | | | | |
|----|--------------------------------|-------|------------|--------------|-----|------|
| AT | ACTAVIS MID ATLANTIC | 2%;1% | N87143 001 | Jan 13, 1982 | Jul | CAHN |
| | HYDROCORTISONE AND ACETIC ACID | | | | | |
| AT | VINTAGE | 2%;1% | N40609 001 | Feb 06, 2006 | Jan | NEWA |

ACETYLCHOLINE CHLORIDE

FOR SOLUTION; OPHTHALMIC

| | | | | | | |
|-----|------------|-----------|------------|--|-----|------|
| >D> | MIOCHOL | | | | | |
| >D> | + NOVARTIS | 20MG/VIAL | N16211 001 | | Sep | DISC |
| >A> | @ | 20MG/VIAL | N16211 001 | | Sep | DISC |

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

| | | | | | | |
|----|-------------------|-------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 200MG | N74906 001 | Aug 26, 1997 | Jun | CAHN |
| AB | CLONMEL HLTHCARE | 200MG | N74833 001 | Apr 22, 1997 | May | CAHN |

SUSPENSION; ORAL

ACYCLOVIR

| | | | | | | |
|----|----------------------|-----------|------------|--------------|-----|------|
| AB | ACTAVIS MID ATLANTIC | 200MG/5ML | N74738 001 | Apr 28, 1997 | Jul | CAHN |
|----|----------------------|-----------|------------|--------------|-----|------|

TABLET; ORAL

ACYCLOVIR

| | | | | | | |
|----|-------------------|-------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 400MG | N74870 001 | Jun 05, 1997 | Jun | CAHN |
| AB | | 800MG | N74870 002 | Jun 05, 1997 | Jun | CAHN |
| AB | CLONMEL HLTHCARE | 400MG | N74946 001 | Nov 19, 1997 | May | CAHN |
| AB | | 800MG | N74946 002 | Nov 19, 1997 | May | CAHN |

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR SODIUM

| | | | | | | |
|--|-----------|--------------------|------------|--------------|-----|------|
| | @ HOSPIRA | EQ 500MG BASE/VIAL | N74663 001 | Apr 22, 1997 | Jun | DISC |
| | @ | EQ 1GM BASE/VIAL | N74663 002 | Apr 22, 1997 | Jun | DISC |

ALBUTEROL

AEROSOL, METERED; INHALATION

ALBUTEROL

| | | | | | | |
|--|------------|------------|------------|--------------|-----|------|
| | @ GENPHARM | 0.09MG/INH | N73045 001 | Aug 19, 1997 | Feb | DISC |
| | @ PLIVA | 0.09MG/INH | N74072 001 | Aug 01, 1996 | Feb | DISC |

ALBUTEROL SULFATE

SOLUTION; INHALATION

ALBUTEROL SULFATE

| | | | | | | |
|----|----------------------|----------------|------------|--------------|-----|------|
| AN | ACTAVIS MID ATLANTIC | EQ 0.083% BASE | N73533 001 | Sep 26, 1995 | Jul | CAHN |
| AN | RXELITE | EQ 0.083% BASE | N77569 001 | Apr 04, 2006 | Mar | NEWA |

SYRUP; ORAL

ALBUTEROL SULFATE

| | | | | | | |
|----|----------------------|-----------------|------------|--------------|-----|------|
| AA | ACTAVIS MID ATLANTIC | EQ 2MG BASE/5ML | N74454 001 | Sep 25, 1995 | Jul | CAHN |
| AA | | EQ 2MG BASE/5ML | N75262 001 | Mar 30, 1999 | Jul | CAHN |

TABLET, EXTENDED RELEASE; ORAL

VOSPIRE ER

| | | | | | | |
|-----|-----------------|-------------|------------|--------------|-----|------|
| >A> | DAVA PHARMS INC | EQ 4MG BASE | N76130 002 | Sep 26, 2002 | Sep | CAHN |
| >A> | + | EQ 8MG BASE | N76130 003 | Sep 26, 2002 | Sep | CAHN |
| >D> | ODYSSEY PHARMS | EQ 4MG BASE | N76130 002 | Sep 26, 2002 | Sep | CAHN |

TABLET, EXTENDED RELEASE; ORAL

VOSPIRE ER

| | | | | | | | |
|-----|---|----------------|-------------|------------|--------------|-----|------|
| >D> | + | ODYSSEY PHARMS | EQ 8MG BASE | N76130 003 | Sep 26, 2002 | Sep | CAHN |
|-----|---|----------------|-------------|------------|--------------|-----|------|

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

INJECTABLE; INJECTION

INFUVITE ADULT

| | | | | | | |
|---|--------|--|------------|--------------|-----|------|
| + | SANDOZ | 2 IU/ML;40MG/ML;12UGM/ML;40 IU/ML;1UGM/ML;3MG/ML;120UGM/ML;8M G/ML;1.2MG/ML;0.72MG/ML;1.2MG/ML; 660 IU/ML;0.03MG/ML | N21163 001 | May 18, 2000 | Jan | CAHN |
|---|--------|--|------------|--------------|-----|------|

INJECTABLE; IV (INFUSION)

INFUVITE ADULT

| | | | | | | |
|---|--------|---|------------|--------------|-----|------|
| + | SANDOZ | 2 IU/ML;40MG/ML;12UGM/ML;40 IU/ML;1UGM/ML;3MG/ML;120UGM/ML;8M G/ML;1.2MG/ML;0.72MG/ML;1.2MG/ML; 660 IU/ML;30UGM/ML | N21559 001 | Jun 16, 2003 | Jan | CAHN |
|---|--------|---|------------|--------------|-----|------|

ALPRAZOLAM

TABLET; ORAL

ALPRAZOLAM

| | | | | | | |
|----|-------------------|--------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 0.25MG | N74342 001 | Oct 31, 1993 | Jun | CAHN |
| AB | | 0.5MG | N74342 002 | Oct 31, 1993 | Jun | CAHN |
| AB | | 1MG | N74342 003 | Oct 31, 1993 | Jun | CAHN |
| AB | | 2MG | N74342 004 | Oct 31, 1993 | Jun | CAHN |

TABLET, EXTENDED RELEASE; ORAL

ALPRAZOLAM

| | | | | | | |
|----|--------|-------|------------|--------------|-----|------|
| AB | BARR | 0.5MG | N77725 001 | Jul 31, 2006 | Jul | NEWA |
| AB | | 1MG | N77725 002 | Jul 31, 2006 | Jul | NEWA |
| AB | | 2MG | N77725 004 | Jul 31, 2006 | Aug | NEWA |
| AB | | 3MG | N77725 003 | Jul 31, 2006 | Jul | NEWA |
| AB | MYLAN | 0.5MG | N77391 002 | Jan 26, 2006 | Jan | NEWA |
| AB | | 1MG | N77391 003 | Jan 26, 2006 | Jan | NEWA |
| AB | | 2MG | N77391 004 | Jan 26, 2006 | Jan | NEWA |
| AB | | 3MG | N77391 001 | Jan 26, 2006 | Jan | NEWA |
| AB | SANDOZ | 0.5MG | N77777 001 | Jun 30, 2006 | Jun | NEWA |
| AB | | 1MG | N77777 002 | Jun 30, 2006 | Jun | NEWA |
| AB | | 2MG | N77777 003 | Jun 30, 2006 | Jun | NEWA |
| AB | | 3MG | N77777 004 | Jun 30, 2006 | Jun | NEWA |

XANAX XR

| | | | | | | |
|----|----------------------|-------|------------|--------------|-----|------|
| AB | PHARMACIA AND UPJOHN | 0.5MG | N21434 001 | Jan 17, 2003 | Jan | CFTG |
| AB | | 1MG | N21434 002 | Jan 17, 2003 | Jan | CFTG |
| AB | | 2MG | N21434 003 | Jan 17, 2003 | Jan | CFTG |
| AB | + | 3MG | N21434 004 | Jan 17, 2003 | Jan | CFTG |

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL

AMANTADINE HYDROCHLORIDE

| | | | | | | |
|----|-------------|-------|------------|--------------|-----|------|
| AB | AMIDE PHARM | 100MG | N77659 001 | Feb 23, 2006 | Feb | NEWA |
|----|-------------|-------|------------|--------------|-----|------|

SYRUP; ORAL

AMANTADINE HYDROCHLORIDE

| | | | | | | | |
|----|---|----------------------|----------|------------|--------------|-----|------|
| AA | + | ACTAVIS MID ATLANTIC | 50MG/5ML | N72655 001 | Oct 30, 1990 | Jul | CAHN |
|----|---|----------------------|----------|------------|--------------|-----|------|

AMCINONIDE

CREAM; TOPICAL

AMCINONIDE

| | | | | | | | |
|-----|----|-----------|----------|------------|--------------|-----|------|
| >D> | AB | ALTANA | 0.1% | N76065 001 | May 15, 2003 | Sep | CRLD |
| >A> | AB | + | 0.1% | N76065 001 | May 15, 2003 | Sep | CRLD |
| >D> | | CYCLOCORT | | | | | |
| >D> | AB | + | ASTELLAS | 0.1% | N18116 002 | Sep | DISC |
| >A> | | @ | 0.1% | N18116 002 | | Sep | DISC |

LOTION; TOPICAL

AMCINONIDE

| | | | | | | | | |
|-----|----|-----------|----------|------------|--------------|--------------|------|------|
| >D> | AB | ALTANA | 0.1% | N76329 001 | Nov 06, 2002 | Sep | CRLD | |
| >A> | | + | 0.1% | N76329 001 | Nov 06, 2002 | Sep | CRLD | |
| >D> | | CYCLOCORT | | | | | | |
| >D> | AB | + | ASTELLAS | 0.1% | N19729 001 | Jun 13, 1988 | Sep | DISC |
| >A> | | @ | 0.1% | N19729 001 | Jun 13, 1988 | Sep | DISC | |

OINTMENT; TOPICAL

AMCINONIDE

| | | | | | | | |
|-----|----|-----------|----------|------------|--------------|-----|------|
| >D> | AB | ALTANA | 0.1% | N76096 001 | Nov 19, 2002 | Sep | CRLD |
| >A> | AB | + | 0.1% | N76096 001 | Nov 19, 2002 | Sep | CRLD |
| >D> | | CYCLOCORT | | | | | |
| >D> | AB | + | ASTELLAS | 0.1% | N18498 001 | Sep | DISC |
| >A> | | @ | 0.1% | N18498 001 | | Sep | DISC |

AMILORIDE HYDROCHLORIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE

| | | | | | | | |
|--|---|-----------|-----|------------|--------------|-----|------|
| | + | PAR PHARM | 5MG | N70346 001 | Jan 22, 1986 | Jun | CRLD |
| | | MIDAMOR | | | | | |
| | | @ MERCK | 5MG | N18200 001 | | Jun | DISC |

AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

| | | | | | | | |
|----|---|----------------|-----------------------|------------|--------------|-----|------|
| AB | + | MYLAN | EQ 5MG ANHYDROUS;50MG | N73209 001 | Oct 31, 1991 | Jun | CRLD |
| | | MODURETIC 5-50 | | | | | |
| | | @ MERCK | EQ 5MG ANHYDROUS;50MG | N18201 001 | | Jun | DISC |

AMINOPHYLLINE

SOLUTION; ORAL

AMINOPHYLLINE DYE FREE

| | | | | | | | |
|----|---|----------------------|-----------|------------|--------------|-----|------|
| AA | + | ACTAVIS MID ATLANTIC | 105MG/5ML | N87727 001 | Apr 16, 1982 | Jul | CAHN |
|----|---|----------------------|-----------|------------|--------------|-----|------|

AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

| | | | | | | | |
|----|---|-------|---------|------------|--------------|-----|------|
| AP | + | AKORN | 50MG/ML | N76232 001 | Jul 05, 2006 | Jun | NEWA |
|----|---|-------|---------|------------|--------------|-----|------|

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE; ORAL

LOTREL

NOVARTIS

| | | | | | | | |
|--|---|--|-------------------|------------|--------------|-----|------|
| | | | EQ 5MG BASE;40MG | N20364 007 | Apr 11, 2006 | Apr | NEWA |
| | | | EQ 10MG BASE;20MG | N20364 005 | Jun 20, 2002 | Apr | CRLD |
| | + | | EQ 10MG BASE;40MG | N20364 006 | Apr 11, 2006 | Apr | NEWA |

AMMONIUM LACTATE

CREAM; TOPICAL

AMMONIUM LACTATE

| | | | | | | |
|----|---------|-------------|------------|--------------|-----|------|
| AB | PADDOCK | EQ 12% BASE | N76829 001 | Feb 07, 2006 | Jan | NEWA |
|----|---------|-------------|------------|--------------|-----|------|

AMOXICILLIN

CAPSULE; ORAL

AMOXICILLIN

| | | | | | | |
|----|----------------|-------|------------|--|-----|------|
| AB | AM ANTIBIOTICS | 250MG | N62058 001 | | Jan | CAHN |
| AB | | 500MG | N62058 002 | | Jan | CAHN |

FOR SUSPENSION; ORAL

AMOXICILLIN

| | | | | | | |
|----|----------------|-----------|------------|--------------|-----|------|
| AB | AM ANTIBIOTICS | 125MG/5ML | N62059 001 | | Jan | CAHN |
| AB | | 250MG/5ML | N62059 002 | | Jan | CAHN |
| AB | HIKMA | 125MG/5ML | N65322 002 | Jun 19, 2006 | Jun | NEWA |
| AB | | 200MG/5ML | N65325 002 | Jun 19, 2006 | Jun | NEWA |
| AB | | 250MG/5ML | N65322 001 | Jun 19, 2006 | Jun | NEWA |
| AB | | 400MG/5ML | N65325 001 | Jun 19, 2006 | Jun | NEWA |

TABLET; ORAL

AMOXICILLIN

| | | | | | | |
|----|-------|-------|------------|--------------|-----|------|
| AB | HIKMA | 875MG | N65255 001 | Mar 29, 2006 | Mar | NEWA |
|----|-------|-------|------------|--------------|-----|------|

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

| | | | | | | |
|----|-------------|-----------------------------|------------|--------------|-----|------|
| AB | TEVA PHARMS | 1.25MG;1.25MG;1.25MG;1.25MG | N40472 001 | Sep 30, 2003 | May | CAHN |
| AB | | 2.5MG;2.5MG;2.5MG;2.5MG | N40472 002 | Sep 30, 2003 | May | CAHN |
| AB | | 5MG;5MG;5MG;5MG | N40472 003 | Sep 30, 2003 | May | CAHN |
| AB | | 7.5MG;7.5MG;7.5MG;7.5MG | N40472 004 | Sep 30, 2003 | May | CAHN |

AMPHOTERICIN B

INJECTABLE; INJECTION

AMPHOTERICIN B

| | | | | | | |
|----|--------------|-----------|------------|--------------|-----|------|
| AP | X GEN PHARMS | 50MG/VIAL | N63206 001 | Apr 29, 1992 | Jun | CAHN |
|----|--------------|-----------|------------|--------------|-----|------|

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

| | | | | | | |
|----|--------|--|------------|--------------|-----|------|
| AP | SANDOZ | EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL | N65241 001 | Jul 25, 2006 | Jul | NEWA |
| AP | | EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL | N65310 001 | Jul 25, 2006 | Jul | NEWA |
| AP | | EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL | N65241 002 | Jul 25, 2006 | Jul | NEWA |
| AP | | EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL | N65310 002 | Jul 25, 2006 | Jul | NEWA |
| AP | | EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL | N65240 001 | Jul 25, 2006 | Jul | NEWA |

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMPICILLIN TRIHYDRATE

@ AM ANTIBIOTICS

@

| | | | | | |
|--|---------------|------------|--|-----|------|
| | EQ 250MG BASE | N61602 001 | | Jan | CAHN |
| | EQ 500MG BASE | N61602 002 | | Jan | CAHN |

FOR SUSPENSION; ORAL

AMPICILLIN TRIHYDRATE

@ AM ANTIBIOTICS

| | | | | | |
|--|-------------------|------------|--|-----|------|
| | EQ 125MG BASE/5ML | N61601 001 | | Jan | CAHN |
|--|-------------------|------------|--|-----|------|

FOR SUSPENSION; ORAL
 AMPICILLIN TRIHYDRATE
 @ AM ANTIBIOTICS

EQ 250MG BASE/5ML

N61601 002

Jan CAHN

ANAGRELIDE HYDROCHLORIDE

CAPSULE; ORAL
 ANAGRELIDE HYDROCHLORIDE

AB ALPHAPHARM

EQ 0.5MG BASE

N77613 001 Jun 27, 2006 Jun NEWA

AB

EQ 1MG BASE

N77613 002 Jun 27, 2006 Jun NEWA

ANIDULAFUNGIN

INJECTABLE; IV (INFUSION)
 ERAXIS

+ VICURON

50MG/VIAL

N21632 001 Feb 17, 2006 Feb NEWA

ANISINDIONE

TABLET; ORAL
 MIRADON

@ SCHERING

50MG

N10909 003

Jan DISC

APREPITANT

CAPSULE; ORAL
 EMEND

MERCK

40MG

N21549 003 Jun 30, 2006 Jun NEWA

ARIPIRAZOLE

>A> INJECTABLE; INTRAMUSCULAR

>A> ABILIFY

>A> + OTSUKA 9.75MG/1.3ML (7.5MG/ML)

N21866 001 Sep 20, 2006 Sep NEWA

TABLET; ORAL

ABILIFY

OTSUKA

2MG

N21436 006 Nov 15, 2002 Jul CMFD

>D> 10MG

N21436 001 Nov 15, 2002 Sep CRLD

>A> + 10MG

N21436 001 Nov 15, 2002 Sep CRLD

TABLET, ORALLY DISINTEGRATING; ORAL

ABILIFY

OTSUKA

10MG

N21729 002 Jun 07, 2006 Jun NEWA

15MG

N21729 003 Jun 07, 2006 Jun NEWA

20MG

N21729 004 Jun 07, 2006 Jun NEWA

+ 30MG

N21729 005 Jun 07, 2006 Jun NEWA

ARTICAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE; INJECTION
 SEPTOCAINE

DEPROCO

4%;EQ 0.005MG BASE/ML

N22010 001 Mar 30, 2006 Mar NEWA

+ 4%; EQ 0.017MG BASE/1.7ML (4%; EQ 0.01MG BASE/ML)

N20971 001 Apr 03, 2000 Mar CPOT

ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION
 SEPTOCAINE

+ DEPROCO

4%;EQ 0.0085MG BASE/1.7ML(4%; EQ 0.005MG BASE/ML)

N22010 001 Mar 30, 2006 Apr CAIN

+ 4%; EQ 0.017MG BASE/1.7ML (4%; EQ 0.01MG BASE/ML)

N20971 001 Apr 03, 2000 Apr CAIN

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

INJECTABLE; IV (INFUSION)

INFUVITE PEDIATRIC

| | | | | | | | |
|--|---|--------|---|------------|--------------|-----|------|
| | + | SANDOZ | 80MG/VIAL;0.02MG/VIAL;400 IU/VIAL;0.001MG/VIAL;5MG/VIAL;0.1 4MG/VIAL;17MG/VIAL;1MG/VIAL;1.4MG /VIAL;1.2MG/VIAL;7 IU/VIAL;2,300 IU/VIAL;0.2MG/VIAL | N21265 001 | Feb 21, 2001 | Jan | CAHN |
| | | | INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE) | | | | |
| | + | SANDOZ | 80MG/VIAL;0.02MG/VIAL;400 IU/VIAL;0.001MG/VIAL;5MG/VIAL;0.1 4MG/VIAL;17MG/VIAL;1MG/VIAL;1.4MG /VIAL;1.2MG/VIAL;7 IU/VIAL;2,300 IU/VIAL;0.2MG/VIAL | N21646 001 | Jan 29, 2004 | Jan | CAHN |

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

FOR SOLUTION; ORAL

MOVIPREP

| | | | | | | | |
|-----|---|--------------|---|------------|--------------|-----|------|
| >D> | + | NORGINE B V | 4.7GM;100GM;1.015GM;5.9GM;2.691GM ;7.5GM | N21881 001 | Aug 02, 2006 | Sep | CAHN |
| | + | | 4.7GM;100GM;1.015GM;5.9GM;2.691GM ;7.5GM | N21881 001 | Aug 02, 2006 | Aug | NEWA |
| >A> | + | SALIX PHARMS | 4.7GM;100GM;1.015GM;5.9GM;2.691GM ;7.5GM | N21881 001 | Aug 02, 2006 | Sep | CAHN |

ASPIRIN; BUTALBITAL; CAFFEINE

TABLET; ORAL

ASPIRIN AND CAFFEINE W/ BUTALBITAL

| | | | | | | | |
|----|--|-------------------|-----------------|------------|--------------|-----|------|
| AB | | ACTAVIS ELIZABETH | 325MG;50MG;40MG | N86710 002 | Aug 23, 1983 | Jun | CAHN |
|----|--|-------------------|-----------------|------------|--------------|-----|------|

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

| | | | | | | | |
|----|---|---------------|----------------------|------------|--------------|-----|------|
| | | @ ENDO PHARMS | 325MG;50MG;40MG;30MG | N75351 001 | Mar 05, 1999 | Feb | DISC |
| AB | + | WATSON PHARMS | 325MG;50MG;40MG;30MG | N19429 003 | Oct 26, 1990 | Apr | CTNA |

ASPIRIN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

ASPIRIN AND OXYCODONE

| | | | | | | | |
|--|---|-------------|----------------|------------|--------------|-----|------|
| | + | ENDO PHARMS | 325MG;4.8355MG | N07337 007 | Aug 05, 2005 | Jun | NEWA |
|--|---|-------------|----------------|------------|--------------|-----|------|

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET; ORAL

PERCODAN-DEMI

| | | | | | | | |
|--|--|---------------|---------------------|------------|--|-----|------|
| | | @ ENDO PHARMS | 325MG;2.25MG;0.19MG | N07337 005 | | Feb | DISC |
|--|--|---------------|---------------------|------------|--|-----|------|

ATENOLOL

TABLET; ORAL

ATENOLOL

| | | | | | | | |
|----|--|-------------------|-------|------------|--------------|-----|------|
| AB | | UNIQUE PHARM LABS | 25MG | N77443 001 | Sep 13, 2006 | Aug | NEWA |
| AB | | | 50MG | N77443 002 | Sep 13, 2006 | Aug | NEWA |
| AB | | | 100MG | N77443 003 | Sep 13, 2006 | Aug | NEWA |

ATOVAQUONE

TABLET; ORAL

MEPRON

@ GLAXOSMITHKLINE

250MG

N20259 001 Nov 25, 1992 May DISC

>A>

ATROPINE; PRALIDOXIME CHLORIDE

>A>

INJECTABLE; INTRAMUSCULAR

>A>

DUODOTE

>A>

+ MERIDIAN MEDCL

2.1MG/0.7ML;600MG/2ML

N21983 001 Sep 28, 2006 Sep NEWA

AZITHROMYCIN

FOR SUSPENSION; ORAL

AZITHROMYCIN

AB PLIVA

EQ 100MG BASE/5ML

N65246 002 Jul 05, 2006 Jun NEWA

AB

EQ 200MG BASE/5ML

N65246 001 Jul 05, 2006 Jun NEWA

>A>

AB

SANDOZ

EQ 100MG BASE/5ML

N65297 001 Sep 18, 2006 Sep NEWA

>A>

AB

ZITHROMAX

EQ 200MG BASE/5ML

N65297 002 Sep 18, 2006 Sep NEWA

PFIZER

EQ 100MG BASE/5ML

N50710 001 Oct 19, 1995 Jun CFTG

AB

+

ZITHROMAX

EQ 200MG BASE/5ML

N50710 002 Oct 19, 1995 Jun CFTG

BACAMPICILLIN HYDROCHLORIDE

FOR SUSPENSION; ORAL

SPECTROBID

@ PFIZER

125MG/5ML

N50556 001 Mar 23, 1982 Feb DISC

TABLET; ORAL

SPECTROBID

@ PFIZER

400MG

N50520 001 Feb DISC

BACLOFEN

TABLET; ORAL

BACLOFEN

AB CARACO

10MG

N77984 001 Aug 14, 2006 Aug NEWA

AB

BACLOFEN

20MG

N77862 002 Aug 14, 2006 Aug NEWA

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

AB APOTEX INC

5MG

N77128 001 Mar 08, 2006 Feb NEWA

AB

BENAZEPRIL

10MG

N77128 002 Mar 08, 2006 Feb NEWA

AB

BENAZEPRIL

20MG

N77128 003 Mar 08, 2006 Feb NEWA

AB

BENAZEPRIL

40MG

N77128 004 Mar 08, 2006 Feb NEWA

AB

BIOKEY

5MG

N76820 001 Feb 03, 2006 Jan NEWA

AB

BIOKEY

10MG

N76820 002 Feb 03, 2006 Jan NEWA

AB

BIOKEY

20MG

N76820 003 Feb 03, 2006 Jan NEWA

AB

BIOKEY

40MG

N76820 004 Feb 03, 2006 Jan NEWA

BENDROFLUMETHIAZIDE

TABLET; ORAL

NATURETIN-5

@ APOTHECON

5MG

N12164 002 Jun DISC

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL

BENZPHETAMINE HYDROCHLORIDE

| | | | | | | |
|----|------------------------|------|------------|--------------|-----|------|
| AA | PADDOCK | 50MG | N40578 001 | Apr 17, 2006 | Apr | NEWA |
| AA | DIDREX | | | | | |
| AA | + PHARMACIA AND UPJOHN | 50MG | N12427 002 | | Apr | CFTG |

BENZQUINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

EMETE-CON

| | | | | | | |
|--|----------|-------------------|------------|--|-----|------|
| | @ PFIZER | EQ 50MG BASE/VIAL | N16820 001 | | Mar | DISC |
|--|----------|-------------------|------------|--|-----|------|

BETAINE, ANHYDROUS

FOR SOLUTION; ORAL

CYSTADANE

| | | | | | | |
|--|--------|--------------|------------|--------------|-----|------|
| | + JAZZ | 1GM/SCOOPFUL | N20576 001 | Oct 25, 1996 | Feb | CAHN |
|--|--------|--------------|------------|--------------|-----|------|

BETAMETHASONE DIPROPIONATE

CREAM; TOPICAL

ALPHATREX

| | | | | | | |
|--|---------------|---------------|------------|--------------|-----|------|
| | @ SAVAGE LABS | EQ 0.05% BASE | N19138 001 | Jun 26, 1984 | Jun | DISC |
|--|---------------|---------------|------------|--------------|-----|------|

BETAMETHASONE DIPROPIONATE

| | | | | | | |
|----|----------------------|---------------|------------|--------------|-----|------|
| AB | ACTAVIS MID ATLANTIC | EQ 0.05% BASE | N70885 001 | Feb 03, 1987 | Jun | CAHN |
|----|----------------------|---------------|------------|--------------|-----|------|

| | | | | | | |
|----|-----------|---------------|------------|--------------|-----|------|
| AB | + FOUGERA | EQ 0.05% BASE | N19137 001 | Jun 26, 1984 | Jun | CRLD |
|----|-----------|---------------|------------|--------------|-----|------|

GEL, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

| | | | | | | |
|----|----------|---------------|------------|--------------|-----|------|
| AB | + ALTANA | EQ 0.05% BASE | N75276 001 | May 13, 2003 | Aug | CRLD |
|----|----------|---------------|------------|--------------|-----|------|

DIPROLENE

| | | | | | | |
|--|------------|---------------|------------|--------------|-----|------|
| | @ SCHERING | EQ 0.05% BASE | N19408 002 | Nov 22, 1991 | Aug | DISC |
|--|------------|---------------|------------|--------------|-----|------|

LOTION; TOPICAL

ALPHATREX

| | | | | | | |
|--|---------------|---------------|------------|--------------|-----|------|
| | @ SAVAGE LABS | EQ 0.05% BASE | N70273 001 | Aug 12, 1985 | Jun | DISC |
|--|---------------|---------------|------------|--------------|-----|------|

BETAMETHASONE DIPROPIONATE

| | | | | | | |
|----|----------------------|---------------|------------|--------------|-----|------|
| AB | ACTAVIS MID ATLANTIC | EQ 0.05% BASE | N70281 001 | Jul 31, 1985 | Jul | CAHN |
|----|----------------------|---------------|------------|--------------|-----|------|

OINTMENT; TOPICAL

BETAMETHASONE DIPROPIONATE

| | | | | | | |
|----|----------------------|---------------|------------|--------------|-----|------|
| AB | ACTAVIS MID ATLANTIC | EQ 0.05% BASE | N71012 001 | Feb 03, 1987 | Jun | CAHN |
|----|----------------------|---------------|------------|--------------|-----|------|

OINTMENT, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

| | | | | | | |
|----|----------------------|---------------|------------|--------------|-----|------|
| AB | ACTAVIS MID ATLANTIC | EQ 0.05% BASE | N74304 001 | Aug 31, 1995 | Jun | CAHN |
|----|----------------------|---------------|------------|--------------|-----|------|

BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE

OINTMENT; TOPICAL

TACLONEX

| | | | | | | |
|--|-------------------|---------------|------------|--------------|-----|------|
| | + LEO PHARM PRODS | 0.064%;0.005% | N21852 001 | Jan 09, 2006 | Jan | NEWA |
|--|-------------------|---------------|------------|--------------|-----|------|

BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

CREAM; TOPICAL

CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE

| | | | | | | |
|----|----------------------|------------------|------------|--------------|-----|------|
| AB | ACTAVIS MID ATLANTIC | EQ 0.05% BASE;1% | N76002 001 | Aug 02, 2002 | Jun | CAHN |
|----|----------------------|------------------|------------|--------------|-----|------|

BETAMETHASONE VALERATE

CREAM; TOPICAL

VALNAC

| | | | | | | | |
|----|----------------------|--------------|--------|-----|--------------|-----|------|
| AB | ACTAVIS MID ATLANTIC | EQ 0.1% BASE | N70050 | 001 | Oct 10, 1984 | Jun | CAHN |
|----|----------------------|--------------|--------|-----|--------------|-----|------|

LOTION; TOPICAL

BETAMETHASONE VALERATE

| | | | | | | | |
|----|----------------------|--------------|--------|-----|--------------|-----|------|
| AB | ACTAVIS MID ATLANTIC | EQ 0.1% BASE | N70052 | 001 | Jul 31, 1985 | Jul | CAHN |
|----|----------------------|--------------|--------|-----|--------------|-----|------|

OINTMENT; TOPICAL

BETAMETHASONE VALERATE

| | | | | | | | |
|----|----------------------|--------------|--------|-----|--------------|-----|------|
| AB | ACTAVIS MID ATLANTIC | EQ 0.1% BASE | N70051 | 001 | Oct 10, 1984 | Jun | CAHN |
|----|----------------------|--------------|--------|-----|--------------|-----|------|

>A> BISKALCITRATE; METRONIDAZOLE; TETRACYCLINE

>A> CAPSULE; ORAL

>A> PYLERA

| | | | | | | | | |
|-----|---|-------------------|-------------------|--------|-----|--------------|-----|------|
| >A> | + | AXCAN SCANDIPHARM | 140MG;125MG;125MG | N50786 | 001 | Sep 28, 2006 | Sep | NEWA |
|-----|---|-------------------|-------------------|--------|-----|--------------|-----|------|

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

| | | | | | | | |
|----|-------------------|--------------|--------|-----|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 2.5MG;6.25MG | N75672 | 001 | Sep 25, 2000 | Jun | CAHN |
|----|-------------------|--------------|--------|-----|--------------|-----|------|

| | | | | | | | |
|----|--|------------|--------|-----|--------------|-----|------|
| AB | | 5MG;6.25MG | N75672 | 002 | Sep 25, 2000 | Jun | CAHN |
|----|--|------------|--------|-----|--------------|-----|------|

| | | | | | | | |
|----|--|-------------|--------|-----|--------------|-----|------|
| AB | | 10MG;6.25MG | N75672 | 003 | Sep 25, 2000 | Jun | CAHN |
|----|--|-------------|--------|-----|--------------|-----|------|

BLEOMYCIN SULFATE

INJECTABLE; INJECTION

BLEOMYCIN SULFATE

| | | | | | | | |
|---|--------------|-----------------------|--------|-----|--------------|-----|------|
| @ | SICOR PHARMS | EQ 15 UNITS BASE/VIAL | N64084 | 001 | Jun 01, 1996 | Jun | DISC |
|---|--------------|-----------------------|--------|-----|--------------|-----|------|

| | | | | | | | |
|---|--|-----------------------|--------|-----|--------------|-----|------|
| @ | | EQ 30 UNITS BASE/VIAL | N64084 | 002 | Jun 01, 1996 | Jun | DISC |
|---|--|-----------------------|--------|-----|--------------|-----|------|

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

ALPHAGAN P

| | | | | | | | | |
|----|---|----------|-------|--------|-----|--------------|-----|------|
| AT | + | ALLERGAN | 0.15% | N21262 | 001 | Mar 16, 2001 | May | CTEC |
|----|---|----------|-------|--------|-----|--------------|-----|------|

BRIMONIDINE TARTRATE

| | | | | | | | | |
|----|--|-------|------|--------|-----|--------------|-----|------|
| AT | | AKORN | 0.2% | N76439 | 001 | Mar 14, 2006 | Feb | NEWA |
|----|--|-------|------|--------|-----|--------------|-----|------|

| | | | | | | | | |
|----|--|-----------|-------|--------|-----|--------------|-----|------|
| AT | | ALCON RES | 0.15% | N21764 | 001 | May 22, 2006 | May | NEWA |
|----|--|-----------|-------|--------|-----|--------------|-----|------|

BRINZOLAMIDE

SUSPENSION/DROPS; OPHTHALMIC

AZOPT

| | | | | | | | |
|---|-------|----|--------|-----|--------------|-----|------|
| + | ALCON | 1% | N20816 | 001 | Apr 01, 1998 | Feb | CAHN |
|---|-------|----|--------|-----|--------------|-----|------|

BROMODIPHENHYDRAMINE HYDROCHLORIDE; CODEINE PHOSPHATE

SYRUP; ORAL

AMBENYL

| | | | | | | | |
|---|-------------|---------------------|--------|-----|--------------|-----|------|
| @ | FOREST LABS | 12.5MG/5ML;10MG/5ML | N09319 | 006 | Jan 10, 1984 | Aug | DISC |
|---|-------------|---------------------|--------|-----|--------------|-----|------|

MYBANIL

| | | | | | | | |
|---|--------------|---------------------|--------|-----|--------------|-----|------|
| + | MORTON GROVE | 12.5MG/5ML;10MG/5ML | N88626 | 001 | Oct 12, 1984 | Aug | CRLD |
|---|--------------|---------------------|--------|-----|--------------|-----|------|

BUDESONIDE

POWDER, METERED; INHALATION

BUDESONIDE

| | | | | | | | |
|--|-------------|------------|--------|-----|--------------|-----|------|
| | ASTRAZENECA | 0.08MG/INH | N21949 | 001 | Jul 12, 2006 | Jul | NEWA |
|--|-------------|------------|--------|-----|--------------|-----|------|

| | | | | | | | |
|---|--|------------|--------|-----|--------------|-----|------|
| + | | 0.16MG/INH | N21949 | 002 | Jul 12, 2006 | Jul | NEWA |
|---|--|------------|--------|-----|--------------|-----|------|

SPRAY, METERED; NASAL

RHINOCORT

| | | | | | | | |
|--|---|-------------|-------------|------------|--------------|-----|------|
| | + | ASTRAZENECA | 0.032MG/INH | N20746 001 | Oct 01, 1999 | Mar | CRLD |
| | | @ | 0.064MG/INH | N20746 002 | Oct 01, 1999 | Mar | DISC |

BUDESONIDE; FORMOTEROL FUMARATE

SPRAY, METERED; INHALATION

SYMBICORT

| | | | | | | | |
|--|---|-------------|-----------------------------|------------|--------------|-----|------|
| | | ASTRAZENECA | 0.08MG/INH;EQ 0.045MG BASE | N21929 001 | Jul 21, 2006 | Jul | NEWA |
| | + | | 0.016MG/INH;EQ 0.045MG BASE | N21929 002 | Jul 21, 2006 | Jul | NEWA |

BUMETANIDE

INJECTABLE; INJECTION

BUMETANIDE

| | | | | | | | |
|----|---|---------|-----------|------------|--------------|-----|------|
| AP | + | BEDFORD | 0.25MG/ML | N74441 001 | Jan 27, 1995 | Feb | CRLD |
| | | BUMEX | | | | | |
| | | @ ROCHE | 0.25MG/ML | N18226 001 | Feb 28, 1983 | Feb | DISC |

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE

| | | | | | | | |
|-----|----|---|-------------------------|------------|--------------|-----|------|
| | + | HOSPIRA | 0.5%;EQ 0.009MG BASE/ML | N22046 001 | Jul 13, 1983 | Aug | CTNA |
| >A> | | BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE | | | | | |
| >A> | AP | SEPTODONT | 0.5%;0.0091MG/ML | N77250 001 | Sep 27, 2006 | Sep | NEWA |
| | | BUPIVACAINE HYDROCHLORIDE W/EPINEPHRINE | | | | | |
| | + | HOSPIRA | 0.5%;EQ 0.009MG BASE/ML | N22046 001 | Jul 13, 1983 | Apr | NEWA |

BUPROPION HYDROCHLORIDE

TABLET; ORAL

BUPROPION HYDROCHLORIDE

| | | | | | | | |
|----|--|------------|-------|------------|--------------|-----|------|
| AB | | APOTEX INC | 75MG | N76143 001 | Jan 17, 2006 | Jan | NEWA |
| AB | | | 100MG | N76143 002 | Jan 17, 2006 | Jan | NEWA |

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPIRONE HYDROCHLORIDE

| | | | | | | | |
|----|--|----------------|------|------------|--------------|-----|------|
| AB | | ACTAVIS TOTOWA | 5MG | N75388 001 | May 09, 2002 | Aug | CAHN |
| AB | | | 10MG | N75388 002 | May 09, 2002 | Aug | CAHN |
| AB | | | 15MG | N75388 003 | May 09, 2002 | Aug | CAHN |

BUSULFAN

INJECTABLE; INJECTION

BUSULFEX

| | | | | | | | |
|--|---|-------------------|--------|------------|--------------|-----|------|
| | + | PDL BIOPHARMA INC | 6MG/ML | N20954 001 | Feb 04, 1999 | Jan | CAHN |
|--|---|-------------------|--------|------------|--------------|-----|------|

CAFFEINE CITRATE

SOLUTION; INTRAVENOUS

CAF CIT

| | | | | | | | | |
|-----|----|---|--------------|------------------------------------|------------|--------------|-----|------|
| >A> | AP | + | MEAD JOHNSON | EQ 30MG BASE/3ML (EQ 10MG BASE/ML) | N20793 001 | Sep 21, 1999 | Sep | CFTG |
|-----|----|---|--------------|------------------------------------|------------|--------------|-----|------|

| | | | | | | | | |
|-----|----|------------------|-------------|------------------------------------|------------|--------------|-----|------|
| >A> | | CAFFEINE CITRATE | | | | | | |
| >A> | AP | | PHARMAFORCE | EQ 30MG BASE/3ML (EQ 10MG BASE/ML) | N77233 001 | Sep 21, 2006 | Sep | NEWA |

SOLUTION; INTRAVENOUS, ORAL

CAF CIT

| | | | | | | | | |
|-----|--|---|--------------|------------------------------------|------------|--------------|-----|------|
| >D> | | + | MEAD JOHNSON | EQ 30MG BASE/3ML (EQ 10MG BASE/ML) | N20793 001 | Sep 21, 1999 | Sep | CFTG |
|-----|--|---|--------------|------------------------------------|------------|--------------|-----|------|

SOLUTION; ORAL

CAFCIT

| | | | | | | | |
|-----|----|------------------|---------------------------------------|------------|--------------|-----|------|
| >D> | + | MEAD JOHNSON | EQ 30MG BASE/3ML (EQ 10MG BASE/ML) | N20793 002 | Apr 12, 2000 | Sep | CFTG |
| >A> | AA | + | EQ 30MG BASE/3ML (EQ 10MG BASE/ML) | N20793 002 | Apr 12, 2000 | Sep | CFTG |
| >A> | | CAFFEINE CITRATE | | | | | |
| >A> | AA | PHARMAFORCE | EQ 30MG BASE/3ML (EQ 10MG BASE/ML) | N77304 001 | Sep 21, 2006 | Sep | NEWA |

CAFFEINE; ERGOTAMINE TARTRATE

SUPPOSITORY; RECTAL

CAFERGOT

| | | | | | | | |
|--|---|--------------|-----------|------------|--------------|-----|------|
| | @ | NOVARTIS | 100MG;2MG | N09000 002 | | Feb | DISC |
| | | MIGERGOT | | | | | |
| | + | G AND W LABS | 100MG;2MG | N86557 001 | Oct 04, 1983 | Feb | CRLD |

CALCIPOTRIENE

CREAM; TOPICAL

DOVONEX

| | | | | | | | |
|--|---|-----------|--------|------------|--------------|-----|------|
| | + | LEO PHARM | 0.005% | N20554 001 | Jul 22, 1996 | Feb | CAHN |
|--|---|-----------|--------|------------|--------------|-----|------|

OINTMENT; TOPICAL

DOVONEX

| | | | | | | | |
|--|---|-----------|--------|------------|--------------|-----|------|
| | + | LEO PHARM | 0.005% | N20273 001 | Dec 29, 1993 | Feb | CAHN |
|--|---|-----------|--------|------------|--------------|-----|------|

SOLUTION; TOPICAL

DOVONEX

| | | | | | | | |
|--|---|-----------|--------|------------|--------------|-----|------|
| | + | LEO PHARM | 0.005% | N20611 001 | Mar 03, 1997 | Feb | CAHN |
|--|---|-----------|--------|------------|--------------|-----|------|

CALCITONIN SALMON RECOMBINANT

SPRAY, METERED; NASAL

FORTICAL

| | | | | | | | |
|--|---|--------------|--------------|------------|--------------|-----|------|
| | + | UPSHER SMITH | 200 IU/SPRAY | N21406 001 | Aug 12, 2005 | Jun | CAHN |
|--|---|--------------|--------------|------------|--------------|-----|------|

CALCITONIN, SALMON

INJECTABLE; INJECTION

MIACALCIN

| | | | | | | | |
|--|---|----------|-----------|------------|--------------|-----|------|
| | + | NOVARTIS | 200 IU/ML | N17808 002 | Mar 29, 1991 | Jan | CTEC |
|--|---|----------|-----------|------------|--------------|-----|------|

CALCITRIOL

CAPSULE; ORAL

CALCITRIOL

| | | | | | | | |
|----|--|--------|---------|------------|--------------|-----|------|
| AB | | ROXANE | 0.25UGM | N76917 001 | Mar 27, 2006 | Mar | NEWA |
|----|--|--------|---------|------------|--------------|-----|------|

INJECTABLE; INJECTION

CALCITRIOL

| | | | | | | | |
|----|--|--------------|------------|------------|--------------|-----|------|
| AP | | GENIX THERAP | 0.001MG/ML | N77102 001 | Feb 08, 2006 | Jan | NEWA |
|----|--|--------------|------------|------------|--------------|-----|------|

CALCIUM ACETATE

TABLET; ORAL

PHOSLO

@ NABI

| | | | | | | |
|--|--|------------------|------------|--------------|-----|------|
| | | EQ 169MG CALCIUM | N19976 001 | Dec 10, 1990 | Jun | DISC |
|--|--|------------------|------------|--------------|-----|------|

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

SOLUTION; IRRIGATION

| | | | | | | | |
|-----|----|------------------------|---|------------|--------------|-----|------|
| >A> | | BALANCED SALT SOLUTION | | | | | |
| >A> | AT | AKORN | 0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9M G/ML;6.4MG/ML;1.7MG/ML | N75503 001 | Sep 27, 2006 | Sep | NEWA |

SOLUTION; IRRIGATION

BSS

| | | | | | | | |
|-----|----|-------|-----------------------------------|------------|--------------|-----|------|
| >D> | + | ALCON | 0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9M | N20742 001 | Dec 10, 1997 | Sep | CFTG |
| | | | G/ML;6.4MG/ML;1.7MG/ML | | | | |
| >A> | AT | + | 0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9M | N20742 001 | Dec 10, 1997 | Sep | CFTG |
| | | | G/ML;6.4MG/ML;1.7MG/ML | | | | |

CANDESARTAN CILEXETIL

TABLET; ORAL

ATACAND

ASTRAZENECA

8MG

N20838 002 Jun 04, 1998 May CRLD

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

@ CLONMEL HLTHCARE

12.5MG

N74423 001 Feb 13, 1996 Jan DISC

@

25MG

N74423 002 Feb 13, 1996 Jan DISC

@

50MG

N74423 003 Feb 13, 1996 Jan DISC

@

100MG

N74423 004 Feb 13, 1996 Jan DISC

@ ENDO LABS

12.5MG

N74418 001 Feb 13, 1996 Feb DISC

@

25MG

N74418 002 Feb 13, 1996 Feb DISC

@

50MG

N74418 003 Feb 13, 1996 Feb DISC

@

100MG

N74418 004 Feb 13, 1996 Feb DISC

CAPTOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CAPTOPRIL AND HYDROCHLOROTHIAZIDE

@ ENDO LABS

25MG;15MG

N74788 001 Dec 29, 1997 Feb DISC

@

25MG;25MG

N74788 002 Dec 29, 1997 Feb DISC

@

50MG;15MG

N74788 004 Dec 29, 1997 Feb DISC

@

50MG;25MG

N74788 003 Dec 29, 1997 Feb DISC

CARBAMAZEPINE

TABLET, CHEWABLE; ORAL

CARBAMAZEPINE

AB

JUBILANT PHARMS

100MG

N71940 001 Feb 01, 1988 Jul CAHN

CARBIDOPA; LEVODOPA

TABLET; ORAL

CARBIDOPA AND LEVODOPA

AB

ACTAVIS ELIZABETH

10MG;100MG

N74260 001 Sep 03, 1993 Jun CAHN

AB

ACTAVIS ELIZABETH

25MG;100MG

N74260 002 Sep 03, 1993 Jun CAHN

AB

ACTAVIS ELIZABETH

25MG;250MG

N74260 003 Sep 03, 1993 Jun CAHN

CARBOPLATIN

INJECTABLE; INJECTION

CARBOPLATIN

AP

WATSON LABS

50MG/VIAL

N77383 001 Jan 27, 2006 Jan NEWA

AP

WATSON LABS

150MG/VIAL

N77383 002 Jan 27, 2006 Jan NEWA

AP

WATSON LABS

450MG/VIAL

N77383 003 Jan 27, 2006 Jan NEWA

INJECTABLE; IV (INFUSION)

CARBOPLATIN

@ AM PHARM

EQ 50MG/5ML (10MG/ML)

N77247 001 Oct 21, 2004 Feb DISC

AP

@

EQ 50MG/5ML (10MG/ML)

N77266 001 Feb 15, 2006 Jan NEWA

@

EQ 150MG/15ML (10MG/ML)

N77247 002 Oct 21, 2004 Feb DISC

AP

@

EQ 150MG/15ML (10MG/ML)

N77266 002 Feb 15, 2006 Jan NEWA

INJECTABLE; IV (INFUSION)

CARBOPLATIN

| | | | | | | |
|-----|--------------|-------------------------|-------------------------|--------------|--------------|----------|
| AP | AM PHARM | EQ 450MG/45ML (10MG/ML) | N77266 003 | Feb 15, 2006 | Jan | NEWA |
| AP | | EQ 600MG/60ML (10MG/ML) | N77266 004 | Feb 15, 2006 | Jan | NEWA |
| AP | BEDFORD LABS | EQ 600MG/60ML (10MG/ML) | N77244 004 | Jan 20, 2006 | Jan | NEWA |
| >A> | AP | DABUR ONCOLOGY PLC | EQ 150MG/15ML (10MG/ML) | N77432 002 | Sep 29, 2006 | Sep NEWA |
| >A> | AP | | EQ 450MG/45ML (10MG/ML) | N77432 003 | Sep 29, 2006 | Sep NEWA |
| >A> | AP | | EQ 50MG/5ML (10MG/ML) | N77432 001 | Sep 29, 2006 | Sep NEWA |

CASPOFUNGIN ACETATE

INJECTABLE; IV (INFUSION)

CANCIDAS

| | | | | | | |
|---|-------|-----------|------------|--------------|-----|------|
| + | MERCK | 50MG/VIAL | N21227 001 | Jan 26, 2001 | May | CAHN |
| + | | 70MG/VIAL | N21227 002 | Jan 26, 2001 | May | CAHN |

CEFACTOR

TABLET, EXTENDED RELEASE; ORAL

CEFACTOR

| | | | | | | |
|----|---|-----------|---------------|------------|--------------|----------|
| AB | + | PAR PHARM | EQ 500MG BASE | N65057 001 | Jan 05, 2001 | May CAHN |
|----|---|-----------|---------------|------------|--------------|----------|

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE; ORAL

CEFADROXIL

| | | | | | | |
|-----|----|-----------------|---------------|------------|--------------|----------|
| AB | + | IVAX PHARMS | EQ 500MG BASE | N62766 001 | Mar 03, 1987 | Mar CRLD |
| >A> | AB | ORCHID HLTHCARE | EQ 500MG BASE | N65309 001 | Sep 18, 2006 | Sep NEWA |
| AB | | SANDOZ | EQ 500MG BASE | N62291 001 | | Aug CAHN |
| AB | | TEVA PHARMS | EQ 500MG BASE | N65282 001 | Jan 20, 2006 | Jan NEWA |
| AB | | WESTWARD | EQ 500MG BASE | N65311 001 | Feb 07, 2006 | Jan NEWA |

DURICEF

@ WARNER CHILCOTT

EQ 500MG BASE N50512 001 Jan DISC

FOR SUSPENSION; ORAL

CEFADROXIL

| | | | | | | |
|----|-------------|-------------------|------------|--------------|-----|------|
| + | RANBAXY | EQ 125MG BASE/5ML | N65115 001 | Mar 26, 2003 | Jul | CRLD |
| | | EQ 125MG BASE/5ML | N65115 001 | Mar 26, 2003 | Feb | CTEC |
| AB | TEVA PHARMS | EQ 250MG BASE/5ML | N65278 001 | Jan 20, 2006 | Jan | NEWA |
| AB | | EQ 500MG BASE/5ML | N65278 002 | Jan 20, 2006 | Jan | NEWA |

DURICEF

@ WARNER CHILCOTT

EQ 125MG BASE/5ML N50527 002 Feb DISC

TABLET; ORAL

CEFADROXIL

| | | | | | | |
|-----|----|-------------------|-------------|------------|--------------|----------|
| AB | | HIKMA | EQ 1GM BASE | N65260 001 | Mar 30, 2006 | Mar NEWA |
| AB | + | IVAX PHARMS | EQ 1GM BASE | N62774 001 | Apr 08, 1987 | Feb CRLD |
| >A> | AB | ORCHID HLTHCARE | EQ 1GM BASE | N65301 001 | Sep 18, 2006 | Sep NEWA |
| | | DURICEF | | | | |
| | | @ WARNER CHILCOTT | EQ 1GM BASE | N50528 001 | | Jan DISC |

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

CEFAZOLIN AND DEXTROSE

@ B BRAUN

EQ 500MG BASE/VIAL N50779 001 Jul 27, 2000 May DISC

CEFDINIR

CAPSULE; ORAL

CEFDINIR

| | | | | | | |
|----|--|-------|-------|------------|--------------|----------|
| AB | | LUPIN | 300MG | N65264 001 | May 19, 2006 | May NEWA |
|----|--|-------|-------|------------|--------------|----------|

| | | | | | | |
|---|---|-------------------|--------------------|--------|-----|-----------------------|
| CAPSULE; ORAL | | | | | | |
| OMNICEF | | | | | | |
| AB | + | ABBOTT | 300MG | N50739 | 001 | Dec 04, 1997 May CFTG |
| FOR SUSPENSION; ORAL | | | | | | |
| CEFDIRINR | | | | | | |
| AB | | LUPIN | 125MG/5ML | N65259 | 001 | May 31, 2006 Jul CAHN |
| AB | | LUPIN (USA) | 125MG/5ML | N65259 | 001 | May 31, 2006 May NEWA |
| OMNICEF | | | | | | |
| AB | | ABBOTT | 125MG/5ML | N50749 | 001 | Dec 04, 1997 May CFTG |
| <u>CEFOTAXIME SODIUM</u> | | | | | | |
| INJECTABLE; INJECTION | | | | | | |
| CEFOTAXIME | | | | | | |
| AP | | WOCKHARDT | EQ 1GM BASE/VIAL | N65197 | 001 | Aug 29, 2006 Aug NEWA |
| CEFOTAXIME AND DEXTROSE 2.4% IN PLASTIC CONTAINER | | | | | | |
| | | @ B BRAUN | EQ 2GM BASE | N50792 | 001 | Jul 29, 2004 May DISC |
| CEFOTAXIME AND DEXTROSE 3.9% IN PLASTIC CONTAINER | | | | | | |
| | | @ B BRAUN | EQ 1GM BASE | N50792 | 002 | Jul 29, 2004 May DISC |
| CEFOTAXIME SODIUM | | | | | | |
| AP | | ORCHID HLTHCARE | EQ 500MG BASE/VIAL | N65290 | 001 | Aug 11, 2006 Aug NEWA |
| AP | | | EQ 1GM BASE/VIAL | N65290 | 002 | Aug 11, 2006 Aug NEWA |
| AP | | | EQ 1GM BASE/VIAL | N65293 | 001 | Aug 10, 2006 Aug NEWA |
| AP | | | EQ 2GM BASE/VIAL | N65290 | 003 | Aug 11, 2006 Aug NEWA |
| AP | | | EQ 2GM BASE/VIAL | N65293 | 002 | Aug 10, 2006 Aug NEWA |
| AP | | | EQ 10GM BASE/VIAL | N65292 | 001 | Aug 10, 2006 Aug NEWA |
| <u>CEFOTETAN DISODIUM</u> | | | | | | |
| INJECTABLE; INJECTION | | | | | | |
| CEFOTAN | | | | | | |
| | | @ ASTRAZENECA | EQ 1GM BASE/VIAL | N63293 | 001 | Apr 29, 1993 Jun DISC |
| | | @ | EQ 2GM BASE/VIAL | N63293 | 002 | Apr 29, 1993 Jun DISC |
| <u>CEFOXITIN SODIUM</u> | | | | | | |
| INJECTABLE; INJECTION | | | | | | |
| CEFOXITIN | | | | | | |
| AP | | ORCHID HLTHCARE | EQ 1GM BASE/VIAL | N65313 | 001 | Jan 23, 2006 Jan NEWA |
| AP | | | EQ 2GM BASE/VIAL | N65313 | 002 | Jan 23, 2006 Jan NEWA |
| AP | | | EQ 10GM BASE/VIAL | N65312 | 001 | Feb 13, 2006 Jan NEWA |
| CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER | | | | | | |
| AP | | B BRAUN | EQ 1GM BASE/VIAL | N65214 | 001 | Mar 10, 2006 Feb NEWA |
| AP | | | EQ 2GM BASE/VIAL | N65214 | 002 | Mar 10, 2006 Feb NEWA |
| <u>CEFPROZIL</u> | | | | | | |
| FOR SUSPENSION; ORAL | | | | | | |
| CEFPROZIL | | | | | | |
| AB | | RANBAXY | 125MG/5ML | N65202 | 001 | Jun 30, 2006 Jun NEWA |
| AB | | | 250MG/5ML | N65202 | 002 | Jun 30, 2006 Jun NEWA |
| <u>CEFTAZIDIME (ARGININE FORMULATION)</u> | | | | | | |
| INJECTABLE; INJECTION | | | | | | |
| CEPTAZ | | | | | | |
| | | @ GLAXOSMITHKLINE | 1GM/VIAL | N50646 | 002 | Sep 27, 1990 May DISC |
| | | @ | 2GM/VIAL | N50646 | 003 | Sep 27, 1990 May DISC |
| | | @ | 10GM/VIAL | N50646 | 004 | Sep 27, 1990 May DISC |

CEFTAZIDIME SODIUM

INJECTABLE; INJECTION

CEFTAZIDIME SODIUM IN PLASTIC CONTAINER

| | | | | | | |
|---|-----------------|-----------------|------------|--------------|-----|------|
| @ | BAXTER HLTHCARE | EQ 10MG BASE/ML | N63221 001 | Apr 29, 1993 | Aug | DISC |
| @ | | EQ 20MG BASE/ML | N63221 002 | Apr 29, 1993 | Aug | DISC |
| @ | | EQ 40MG BASE/ML | N63221 003 | Apr 29, 1993 | Aug | DISC |

FORTAZ IN PLASTIC CONTAINER

| | | | | | | |
|---|-----------------|-----------------|------------|--------------|-----|------|
| + | GLAXOSMITHKLINE | EQ 20MG BASE/ML | N50634 002 | Apr 28, 1989 | Aug | CTEC |
| + | | EQ 40MG BASE/ML | N50634 003 | Apr 28, 1989 | Aug | CTEC |

CEFTRIAZONE SODIUM

INJECTABLE; IM-IV

CEFTRIAZONE

| | | | | | | |
|----|-------------------|--------------------|------------|--------------|-----|------|
| AP | AM PHARM PARTNERS | EQ 250MG BASE/VIAL | N65245 001 | Feb 15, 2006 | Jan | NEWA |
| AP | | EQ 500MG BASE/VIAL | N65245 002 | Feb 15, 2006 | Jan | NEWA |
| AP | | EQ 1GM BASE/VIAL | N65245 003 | Feb 15, 2006 | Jan | NEWA |
| AP | | EQ 2GM BASE/VIAL | N65245 004 | Feb 15, 2006 | Jan | NEWA |

CEFTRIAZONE SODIUM

| | | | | | | |
|----|------|------------------|------------|--------------|-----|------|
| AP | TEVA | EQ 1GM BASE/VIAL | N65262 001 | Jun 29, 2006 | Jun | NEWA |
| AP | | EQ 2GM BASE/VIAL | N65262 002 | Jun 29, 2006 | Jun | NEWA |

INJECTABLE; INJECTION

CEFTRIAZONE

| | | | | | | |
|----|-----------|-------------------|------------|--------------|-----|------|
| AP | AM PHARM | EQ 10GM BASE/VIAL | N65252 001 | Feb 15, 2006 | Jan | NEWA |
| AP | LUPIN | EQ 10GM BASE/VIAL | N65263 001 | Sep 12, 2006 | Aug | NEWA |
| AP | WOCKHARDT | EQ 1GM BASE/VIAL | N65180 001 | May 12, 2006 | May | NEWA |

CEFTRIAZONE SODIUM

| | | | | | | |
|----|------|-------------------|------------|--------------|-----|------|
| AP | TEVA | EQ 10GM BASE/VIAL | N65274 001 | May 01, 2006 | Apr | NEWA |
|----|------|-------------------|------------|--------------|-----|------|

CEFUROXIME AXETIL

TABLET; ORAL

CEFUROXIME AXETIL

| | | | | | | |
|----|----------------------|---------------|------------|--------------|-----|------|
| AB | AUROBINDO PHARMA LTD | EQ 125MG BASE | N65308 001 | Mar 29, 2006 | Mar | NEWA |
| AB | | EQ 250MG BASE | N65308 002 | Mar 29, 2006 | Mar | NEWA |
| AB | | EQ 500MG BASE | N65308 003 | Mar 29, 2006 | Mar | NEWA |

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

| | | | | | | |
|----|-------|---------------|------------|--------------|-----|------|
| AB | HIKMA | EQ 250MG BASE | N65215 001 | Jan 24, 2006 | Jan | NEWA |
| AB | | EQ 500MG BASE | N65215 002 | Jan 24, 2006 | Jan | NEWA |

KEFLEX

| | | | | | | |
|----|----------------|---------------|------------|--------------|-----|------|
| AB | ADVANCIS PHARM | EQ 333MG BASE | N50405 004 | May 12, 2006 | May | NEWA |
| AB | | EQ 500MG BASE | N50405 003 | | May | CRLD |
| + | | EQ 750MG BASE | N50405 005 | May 12, 2006 | May | NEWA |

FOR SUSPENSION; ORAL

CEPHALEXIN

| | | | | | | |
|----|-----------------|-------------------|------------|--------------|-----|------|
| AB | ORCHID HLTHCARE | EQ 125MG BASE/5ML | N65326 001 | Jul 10, 2006 | Jun | NEWA |
| AB | | EQ 250MG BASE/5ML | N65326 002 | Jul 10, 2006 | Jun | NEWA |

CEPHRADINE

CAPSULE; ORAL

ANSPOR

| | | | | | | |
|---|-----------------|-------|------------|--|-----|------|
| @ | GLAXOSMITHKLINE | 250MG | N61859 001 | | Mar | DISC |
| @ | | 500MG | N61859 002 | | Mar | DISC |

CETYL ALCOHOL; COLFOSCERIL PALMITATE; TYLOXAPOL

FOR SUSPENSION; INTRATRACHEAL

EXOSURF NEONATAL

| | | | | | | | |
|---|-----------------|-------------------------------|--------|-----|--------------|-----|------|
| @ | GLAXOSMITHKLINE | 12MG/VIAL;108MG/VIAL;8MG/VIAL | N20044 | 001 | Aug 02, 1990 | May | DISC |
|---|-----------------|-------------------------------|--------|-----|--------------|-----|------|

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL

CHLORHEXIDINE GLUCONATE

| | | | | | | | |
|----|----------------------|-------|--------|-----|--------------|-----|------|
| AT | ACTAVIS MID ATLANTIC | 0.12% | N74291 | 001 | Dec 28, 1995 | Jul | CAHN |
|----|----------------------|-------|--------|-----|--------------|-----|------|

CHLOROPROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

NESACAINE

| | | | | | | | | |
|----|---|--------------------|----|--------|-----|--------------|-----|------|
| | + | ABRAXIS BIOSCIENCE | 1% | N09435 | 001 | | Jul | CAHN |
| AP | | | 2% | N09435 | 002 | | Jul | CAHN |
| | | NESACAINE-MPF | | | | | | |
| AP | + | ABRAXIS BIOSCIENCE | 2% | N09435 | 006 | May 02, 1996 | Jul | CAHN |
| | | @ | 2% | N09435 | 003 | | Jul | CAHN |
| | | @ | 3% | N09435 | 004 | | Jul | CAHN |
| AP | + | | 3% | N09435 | 007 | May 02, 1996 | Jul | CAHN |

CHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

DIUPRES-250

| | | | | | | | |
|---|-------|---------------|--------|-----|--------------|-----|------|
| @ | MERCK | 250MG;0.125MG | N11635 | 003 | Aug 26, 1987 | Jun | DISC |
|---|-------|---------------|--------|-----|--------------|-----|------|

DIUPRES-500

| | | | | | | | |
|---|-------|---------------|--------|-----|--------------|-----|------|
| @ | MERCK | 500MG;0.125MG | N11635 | 006 | Aug 26, 1987 | Jun | DISC |
|---|-------|---------------|--------|-----|--------------|-----|------|

CHLORPROMAZINE

SUPPOSITORY; RECTAL

THORAZINE

| | | | | | | | |
|---|-----------------|------|--------|-----|--|-----|------|
| @ | GLAXOSMITHKLINE | 25MG | N09149 | 024 | | May | DISC |
|---|-----------------|------|--------|-----|--|-----|------|

| | | | | | | | |
|---|--|-------|--------|-----|--|-----|------|
| @ | | 100MG | N09149 | 033 | | May | DISC |
|---|--|-------|--------|-----|--|-----|------|

CHLORPROMAZINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

THORAZINE

| | | | | | | | |
|---|-----------------|-------|--------|-----|--|-----|------|
| @ | GLAXOSMITHKLINE | 200MG | N11120 | 019 | | May | DISC |
|---|-----------------|-------|--------|-----|--|-----|------|

| | | | | | | | |
|---|--|-------|--------|-----|--|-----|------|
| @ | | 300MG | N11120 | 020 | | May | DISC |
|---|--|-------|--------|-----|--|-----|------|

CONCENTRATE; ORAL

CHLORPROMAZINE HYDROCHLORIDE

| | | | | | | | |
|----|----------------------|----------|--------|-----|--|-----|------|
| AA | ACTAVIS MID ATLANTIC | 100MG/ML | N86863 | 001 | | Jul | CAHN |
|----|----------------------|----------|--------|-----|--|-----|------|

CICLOPIROX

CREAM; TOPICAL

CICLOPIROX

| | | | | | | | |
|----|---------|-------|--------|-----|--------------|-----|------|
| AB | PERRIGO | 0.77% | N77364 | 001 | Mar 03, 2006 | Mar | CAHN |
|----|---------|-------|--------|-----|--------------|-----|------|

| | | | | | | | |
|----|------------------|-------|--------|-----|--------------|-----|------|
| AB | PERRIGO NEW YORK | 0.77% | N77364 | 001 | Mar 03, 2006 | Feb | NEWA |
|----|------------------|-------|--------|-----|--------------|-----|------|

CILASTATIN SODIUM; IMIPENEM

INJECTABLE; INJECTION

PRIMAXIN

| | | | | | | | |
|---|-------|-------------------------------|--------|-----|--------------|-----|------|
| @ | MERCK | EQ 750MG BASE/VIAL;750MG/VIAL | N50630 | 002 | Dec 14, 1990 | Jun | DISC |
|---|-------|-------------------------------|--------|-----|--------------|-----|------|

CILOSTAZOL

TABLET; ORAL

CILOSTAZOL

| | | | | | | |
|----|--------------|-------|------------|--------------|-----|------|
| AB | MUTUAL PHARM | 50MG | N77208 002 | Mar 29, 2006 | Mar | NEWA |
| AB | | 100MG | N77208 001 | Mar 29, 2006 | Mar | NEWA |
| AB | MYLAN | 50MG | N77323 002 | Apr 20, 2006 | Apr | NEWA |
| AB | | 100MG | N77323 001 | Apr 20, 2006 | Apr | NEWA |

CIMETIDINE

TABLET; ORAL

CIMETIDINE

@ ENDO PHARMS

200MG

N74281 001 May 17, 1994 Feb DISC

@

300MG

N74281 002 May 17, 1994 Feb DISC

@

400MG

N74281 003 May 17, 1994 Feb DISC

@

800MG

N74329 001 May 17, 1994 Feb DISC

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

@ ENDO PHARMS

EQ 300MG BASE/2ML

N74005 001 Aug 31, 1994 Feb DISC

SOLUTION; ORAL

CIMETIDINE HYDROCHLORIDE

AA + ACTAVIS MID ATLANTIC

EQ 300MG BASE/5ML

N74176 001 Jun 01, 1994 Jul CAHN

@ ENDO PHARMS

EQ 300MG BASE/5ML

N74251 001 Dec 22, 1994 Feb DISC

CIPROFLOXACIN

INJECTABLE; INJECTION

CIPRO

AP + BAYER PHARMS

400MG/40ML (10MG/ML)

N19847 001 Dec 26, 1990 Aug CFTG

AP +

200MG/20ML (10MG/ML)

N19847 002 Dec 26, 1990 Aug CFTG

@

1200MG/120ML (10MG/ML)

N19847 003 Dec 26, 1990 Aug DISC

CIPROFLOXACIN

AP ABRAXIS PHARM

400MG/40ML (10MG/ML)

N76484 002 Aug 28, 2006 Aug NEWA

AP

200MG/20ML (10MG/ML)

N76484 001 Aug 28, 2006 Aug NEWA

AP BEDFORD LABS

400MG/40ML (10MG/ML)

N76992 002 Aug 28, 2006 Aug NEWA

AP

200MG/20ML (10MG/ML)

N76992 001 Aug 28, 2006 Aug NEWA

1200MG/120ML (10MG/ML)

N76993 001 Aug 28, 2006 Aug NEWA

AP HOSPIRA

400MG/40ML (10MG/ML)

N77245 002 Aug 28, 2006 Aug NEWA

AP

200MG/20ML (10MG/ML)

N77245 001 Aug 28, 2006 Aug NEWA

AP SICOR PHARMS

200MG/20ML (10MG/ML)

N77782 001 Aug 28, 2006 Aug NEWA

AP

400MG/40ML (10MG/ML)

N77782 002 Aug 28, 2006 Aug NEWA

CITALOPRAM HYDROBROMIDE

SOLUTION; ORAL

CITALOPRAM HYDROBROMIDE

AA AUROBINDO PHARMA LTD

EQ 10MG BASE/5ML

N77812 001 Aug 28, 2006 Aug NEWA

AA

SILARX

EQ 10MG BASE/5ML

N77629 001 Jun 15, 2006 May NEWA

TABLET; ORAL

CITALOPRAM HYDROBROMIDE

AB ACTAVIS ELIZABETH

EQ 10MG BASE

N77033 001 Oct 28, 2004 Jun CAHN

AB

EQ 20MG BASE

N77033 002 Oct 28, 2004 Jun CAHN

AB

EQ 40MG BASE

N77033 003 Oct 28, 2004 Jun CAHN

>A>

AB INVAGEN PHARMS

EQ 10MG BASE

N77534 001 Oct 03, 2006 Sep NEWA

>A>

AB

EQ 20MG BASE

N77534 002 Oct 03, 2006 Sep NEWA

TABLET; ORAL

CITALOPRAM HYDROBROMIDE

| | | | | | | | |
|-----|----|----------------|--------------|------------|--------------|-----|------|
| >A> | AB | INVAGEN PHARMS | EQ 40MG BASE | N77534 003 | Oct 03, 2006 | Sep | NEWA |
| | AB | MUTUAL PHARM | EQ 10MG BASE | N77052 001 | Jul 03, 2006 | Jun | NEWA |
| | AB | | EQ 20MG BASE | N77052 002 | Jul 03, 2006 | Jun | NEWA |
| | AB | | EQ 40MG BASE | N77052 003 | Jul 03, 2006 | Jun | NEWA |
| | AB | TARO | EQ 10MG BASE | N77278 001 | Mar 22, 2006 | Mar | NEWA |
| | AB | | EQ 20MG BASE | N77278 002 | Mar 22, 2006 | Mar | NEWA |
| | AB | | EQ 40MG BASE | N77278 003 | Mar 22, 2006 | Mar | NEWA |
| | AB | TEVA PHARMS | EQ 10MG BASE | N77213 001 | Mar 31, 2006 | Mar | NEWA |
| | AB | | EQ 20MG BASE | N77213 002 | Mar 31, 2006 | Mar | NEWA |
| | AB | | EQ 40MG BASE | N77213 003 | Mar 31, 2006 | Mar | NEWA |

CITRIC ACID; MAGNESIUM OXIDE; SODIUM CARBONATE

SOLUTION; IRRIGATION

IRRIGATING SOLUTION G IN PLASTIC CONTAINER

| | | | | | | | |
|---|--|---------------------------------|--------------------------------------|------------|--------------|-----|------|
| | | @ BAXTER HLTHCARE | 3.24GM/100ML;380MG/100ML;430MG/100ML | N18519 001 | Jun 22, 1982 | Jun | DISC |
| | | UROLOGIC G IN PLASTIC CONTAINER | | | | | |
| + | | HOSPIRA | 3.24GM/100ML;380MG/100ML;430MG/100ML | N18904 001 | May 27, 1983 | Jun | CTEC |

CITRIC ACID; UREA, C-13

FOR SOLUTION, TABLET, FOR SOLUTION; ORAL

IDKIT:HP

| | | | | | | | |
|-----|--|----------------------|------------------|------------|--------------|-----|------|
| >A> | | @ BREATHID 2006 | N/A,4GM;75MG,N/A | N21314 001 | Dec 17, 2002 | Sep | CAHN |
| >D> | | @ ORIDION MEDCL 1987 | N/A,4GM;75MG,N/A | N21314 001 | Dec 17, 2002 | Sep | CAHN |

CLARITHROMYCIN

TABLET; ORAL

CLARITHROMYCIN

| | | | | | | | |
|---|----|--------------------------------|-------|------------|--------------|-----|------|
| | AB | WOCKHARDT | 250MG | N65266 001 | May 31, 2006 | May | NEWA |
| | AB | | 500MG | N65266 002 | May 31, 2006 | May | NEWA |
| | | TABLET, EXTENDED RELEASE; ORAL | | | | | |
| | | CLARITHROMYCIN | | | | | |
| + | | RANBAXY | 1GM | N65210 001 | Jan 26, 2005 | Apr | CRLD |

CLEMASTINE FUMARATE

SYRUP; ORAL

CLEMASTINE FUMARATE

| | | | | | | | |
|----|--|----------------------|-------------------|------------|--------------|-----|------|
| AA | | ACTAVIS MID ATLANTIC | EQ 0.5MG BASE/5ML | N74075 001 | Oct 31, 1993 | Jul | CAHN |
|----|--|----------------------|-------------------|------------|--------------|-----|------|

CLINDAMYCIN PHOSPHATE

SOLUTION; TOPICAL

CLINDAMYCIN PHOSPHATE

| | | | | | | | |
|----|--|----------------------|------------|------------|--------------|-----|------|
| AT | | ACTAVIS MID ATLANTIC | EQ 1% BASE | N62811 001 | Sep 01, 1988 | Jul | CAHN |
| AT | | ALTANA | EQ 1% BASE | N65254 001 | Feb 14, 2006 | Jan | NEWA |

CLOBETASOL PROPIONATE

CREAM; TOPICAL

CLOBETASOL PROPIONATE

| | | | | | | | |
|-----|--|----------------------|-------|------------|--------------|-----|------|
| AB1 | | ACTAVIS MID ATLANTIC | 0.05% | N74139 001 | Aug 03, 1994 | Jun | CAHN |
|-----|--|----------------------|-------|------------|--------------|-----|------|

OINTMENT; TOPICAL

CLOBETASOL PROPIONATE

| | | | | | | | |
|----|--|----------------------|-------|------------|--------------|-----|------|
| AB | | ACTAVIS MID ATLANTIC | 0.05% | N74128 001 | Aug 03, 1994 | Jun | CAHN |
|----|--|----------------------|-------|------------|--------------|-----|------|

SOLUTION; TOPICAL

CLOBETASOL PROPIONATE

| | | | | | | | |
|----|----------------------|-------|--------|-----|--------------|-----|------|
| AT | ACTAVIS MID ATLANTIC | 0.05% | N74331 | 001 | Dec 15, 1995 | Jul | CAHN |
| | @ ALTANA | 0.05% | N75391 | 001 | Feb 08, 1999 | May | DISC |

SPRAY; TOPICAL

CLOBEX

| | | | | | | | |
|---|------------------|-------|--------|-----|--------------|-----|------|
| + | GALDERMA LABS LP | 0.05% | N21835 | 001 | Oct 27, 2005 | Feb | CAHN |
|---|------------------|-------|--------|-----|--------------|-----|------|

CLOFAZIMINE

CAPSULE; ORAL

LAMPRENE

| | | | | | | | |
|---|----------|------|--------|-----|--------------|-----|------|
| + | NOVARTIS | 50MG | N19500 | 002 | Dec 15, 1986 | Jun | CMFD |
|---|----------|------|--------|-----|--------------|-----|------|

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

| | | | | | | | |
|----|-------------------|-------|--------|-----|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 0.5MG | N74869 | 001 | Oct 31, 1996 | Jun | CAHN |
| AB | | 1MG | N74869 | 002 | Oct 31, 1996 | Jun | CAHN |
| AB | | 2MG | N74869 | 003 | Oct 31, 1996 | Jun | CAHN |
| AB | VINTAGE PHARMS | 0.5MG | N77856 | 001 | Jun 28, 2006 | Jun | NEWA |
| AB | | 1MG | N77856 | 002 | Jun 28, 2006 | Jun | NEWA |
| AB | | 2MG | N77856 | 003 | Jun 28, 2006 | Jun | NEWA |

CLONIDINE HYDROCHLORIDE

TABLET; ORAL

CLONIDINE HYDROCHLORIDE

| | | | | | | | |
|----|-------------------|-------|--------|-----|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 0.1MG | N70974 | 001 | Dec 16, 1986 | Jun | CAHN |
| AB | | 0.2MG | N70975 | 001 | Dec 16, 1986 | Jun | CAHN |
| AB | | 0.3MG | N70976 | 001 | Dec 16, 1986 | Jun | CAHN |

CLOPIDOGREL BISULFATE

TABLET; ORAL

CLOPIDOGREL BISULFATE

| | | | | | | | | |
|----|--------|-------------------|--------------|--------|--------------|--------------|------|------|
| AB | APOTEX | EQ 75MG BASE | N76274 | 001 | Jan 20, 2006 | Jan | NEWA | |
| AB | + | SANOFI SYNTHELABO | EQ 75MG BASE | N20839 | 001 | Nov 17, 1997 | Jan | CFTG |

CLOZAPINE

TABLET, ORALLY DISINTEGRATING; ORAL

FAZACLO ODT

| | | | | | | | |
|---|---------------|-------|--------|-----|--------------|-----|------|
| | AVANIR PHARMS | 25MG | N21590 | 001 | Feb 10, 2004 | Jul | CAHN |
| | | 50MG | N21590 | 003 | Jun 03, 2005 | Jul | CAHN |
| + | | 100MG | N21590 | 002 | Feb 10, 2004 | Jul | CAHN |

COBALT CHLORIDE, CO-57; CYANOCOBALAMIN; CYANOCOBALAMIN, CO-57; INTRINSIC FACTOR

N/A; N/A

RUBRATOPE-57 KIT

| | | | | | | | |
|---|--------|-----------------|--------|-----|--|-----|------|
| @ | BRACCO | N/A;N/A;N/A;N/A | N16089 | 001 | | Jun | DISC |
|---|--------|-----------------|--------|-----|--|-----|------|

CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETH VC W/ CODEINE

| | | | | | | | |
|---|----------------------|-----------------------------|--------|-----|--------------|-----|------|
| + | ACTAVIS MID ATLANTIC | 10MG/5ML;5MG/5ML;6.25MG/5ML | N88764 | 001 | Oct 31, 1984 | Jul | CAHN |
| + | ALPHARMA US PHARMS | 10MG/5ML;5MG/5ML;6.25MG/5ML | N88764 | 001 | Oct 31, 1984 | Jan | CTEC |

SYRUP; ORAL

PROMETHAZINE VC W/ CODEINE

| | | | | | |
|----------------|-----------------------------|------------|--------------|-----|------|
| @ MORTON GROVE | 10MG/5ML;5MG/5ML;6.25MG/5ML | N88896 001 | Jan 04, 1985 | Jan | DISC |
|----------------|-----------------------------|------------|--------------|-----|------|

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETH W/ CODEINE

| | | | | | | |
|------|----------------------|---------------------|------------|--------------|-----|------|
| AA + | ACTAVIS MID ATLANTIC | 10MG/5ML;6.25MG/5ML | N88763 001 | Oct 31, 1984 | Jul | CAHN |
|------|----------------------|---------------------|------------|--------------|-----|------|

PROMETHAZINE WITH CODEINE SYRUP

| | | | | | | |
|----|---------|---------------------|------------|--------------|-----|------|
| AA | VINTAGE | 10MG/5ML;6.25MG/5ML | N40650 001 | Jan 31, 2006 | Jan | NEWA |
|----|---------|---------------------|------------|--------------|-----|------|

CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP; ORAL

TRIACIN-C

| | | | | | | |
|----|----------------------|------------------------------|------------|--------------|-----|------|
| AA | ACTAVIS MID ATLANTIC | 10MG/5ML;30MG/5ML;1.25MG/5ML | N88704 001 | Mar 22, 1985 | Jul | CAHN |
|----|----------------------|------------------------------|------------|--------------|-----|------|

COLESTIPOL HYDROCHLORIDE

GRANULE; ORAL

COLESTID

| | | | | | | |
|----|----------------------|--------------|------------|--------------|-----|------|
| AB | PHARMACIA AND UPJOHN | 5GM/SCOOPFUL | N17563 003 | Sep 22, 1995 | Apr | CFTG |
|----|----------------------|--------------|------------|--------------|-----|------|

| | | | | | | |
|------|--|------------|------------|--------------|-----|------|
| AB + | | 5GM/PACKET | N17563 004 | Sep 22, 1995 | Apr | CFTG |
|------|--|------------|------------|--------------|-----|------|

COLESTIPOL HYDROCHLORIDE

| | | | | | | |
|----|------------|--------------|------------|--------------|-----|------|
| AB | IMPAX LABS | 5GM/SCOOPFUL | N77277 001 | May 02, 2006 | Apr | NEWA |
|----|------------|--------------|------------|--------------|-----|------|

| | | | | | | |
|----|--|------------|------------|--------------|-----|------|
| AB | | 5GM/PACKET | N77277 002 | May 02, 2006 | Apr | NEWA |
|----|--|------------|------------|--------------|-----|------|

FLAVORED COLESTID

PHARMACIA AND UPJOHN 5GM/PACKET

| | | | |
|------------|--|-----|------|
| N17563 001 | | Apr | CFTG |
|------------|--|-----|------|

CROMOLYN SODIUM

CONCENTRATE; ORAL

GASTROCROM

| | | | | | |
|---------------|-----------|------------|--------------|-----|------|
| + AZUR PHARMA | 100MG/5ML | N20479 001 | Feb 29, 1996 | May | CDFR |
|---------------|-----------|------------|--------------|-----|------|

SOLUTION; INHALATION

CROMOLYN SODIUM

| | | | | | | |
|----|----------------------|---------|------------|--------------|-----|------|
| AN | ACTAVIS MID ATLANTIC | 10MG/ML | N75067 001 | Jul 19, 1999 | Jul | CAHN |
|----|----------------------|---------|------------|--------------|-----|------|

SOLUTION, CONCENTRATE; ORAL

GASTROCROM

| | | | | | |
|---------------|-----------|------------|--------------|-----|------|
| + AZUR PHARMA | 100MG/5ML | N20479 001 | Feb 29, 1996 | Feb | CAHN |
|---------------|-----------|------------|--------------|-----|------|

CYANOCOBALAMIN

GEL, METERED; NASAL

NASCOBAL

| | | | | | |
|-------------|-----------|------------|--------------|-----|------|
| @ QOL MEDCL | 0.5MG/INH | N19722 001 | Nov 05, 1996 | Mar | DISC |
|-------------|-----------|------------|--------------|-----|------|

CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

| | | | | | | |
|----|-----------------|-----|------------|--------------|-----|------|
| AB | JUBILANT PHARMS | 5MG | N77563 001 | Apr 19, 2006 | Apr | NEWA |
|----|-----------------|-----|------------|--------------|-----|------|

| | | | | | | |
|----|--|------|------------|--------------|-----|------|
| AB | | 10MG | N77563 002 | Apr 19, 2006 | Apr | NEWA |
|----|--|------|------------|--------------|-----|------|

CYCLOBENZAPRINE HYDROCHLORIDE

| | | | | | | |
|----|-------------|-----|------------|--------------|-----|------|
| AB | AMIDE PHARM | 5MG | N77291 001 | Feb 03, 2006 | Jan | NEWA |
|----|-------------|-----|------------|--------------|-----|------|

| | | | | | | |
|----|--------------|-----|------------|--------------|-----|------|
| AB | MUTUAL PHARM | 5MG | N73541 002 | Apr 06, 2006 | Mar | NEWA |
|----|--------------|-----|------------|--------------|-----|------|

| | | | | | | |
|----|-------|-----|------------|--------------|-----|------|
| AB | MYLAN | 5MG | N73144 002 | Feb 03, 2006 | Jan | NEWA |
|----|-------|-----|------------|--------------|-----|------|

| | | | | | | |
|----|--------|-----|------------|--------------|-----|------|
| AB | SANDOZ | 5MG | N72854 002 | Feb 03, 2006 | Jan | NEWA |
|----|--------|-----|------------|--------------|-----|------|

| | | | | | | |
|----|-------------|-----|------------|--------------|-----|------|
| AB | WATSON LABS | 5MG | N71611 002 | Feb 03, 2006 | Jan | NEWA |
|----|-------------|-----|------------|--------------|-----|------|

| | | | | | | |
|--|--|-------|------------|--------------|-----|------|
| | | 7.5MG | N71611 003 | Feb 03, 2006 | Jan | NEWA |
|--|--|-------|------------|--------------|-----|------|

| | | | | | | | | |
|----|-------------------------------------|-----------------------|---------------|------------|------------|--------------|-----|------|
| | TABLET; ORAL | | | | | | | |
| | FLEXERIL | | | | | | | |
| AB | MCNEIL CONS SPECLT | 5MG | | N17821 001 | | | Jan | CFTG |
| | <u>CYPROHEPTADINE HYDROCHLORIDE</u> | | | | | | | |
| | SYRUP; ORAL | | | | | | | |
| | CYPROHEPTADINE HYDROCHLORIDE | | | | | | | |
| AA | + | ACTAVIS MID ATLANTIC | 2MG/5ML | | N86833 001 | | Jul | CAHN |
| AA | + | ALPHARMA US PHARMS | 2MG/5ML | | N86833 001 | | Jun | CTEC |
| AA | | LYNE | 2MG/5ML | | N40668 001 | Jun 28, 2006 | Jun | NEWA |
| | TABLET; ORAL | | | | | | | |
| | CYPROHEPTADINE HYDROCHLORIDE | | | | | | | |
| | | @ TG UNITED LABS | 4MG | | N88212 001 | May 26, 1983 | May | CAHN |
| | CYPROHEPTADINE HYDROCHLORIDE | | | | | | | |
| AA | | STASON PHARMS | 4MG | | N40644 001 | May 30, 2006 | May | NEWA |
| | <u>DAPTOMYCIN</u> | | | | | | | |
| | INJECTABLE; IV (INFUSION) | | | | | | | |
| | CUBICIN | | | | | | | |
| | | @ CUBIST | 250MG/VIAL | | N21572 001 | Sep 12, 2003 | Jun | DISC |
| | <u>DARUNAVIR ETHANOLATE</u> | | | | | | | |
| | TABLET; ORAL | | | | | | | |
| | PREZISTA | | | | | | | |
| | + | TIBOTEC | EQ 300MG BASE | | N21976 001 | Jun 23, 2006 | Jun | NEWA |
| | <u>DASATINIB</u> | | | | | | | |
| | TABLET; ORAL | | | | | | | |
| | SPRYCEL | | | | | | | |
| | | BRISTOL MYERS SQUIBB | 20MG | | N21986 001 | Jun 28, 2006 | Jun | NEWA |
| | | | 50MG | | N21986 002 | Jun 28, 2006 | Jun | NEWA |
| | + | | 70MG | | N21986 003 | Jun 28, 2006 | Jun | NEWA |
| | <u>DECITABINE</u> | | | | | | | |
| | INJECTABLE; INTRAVENOUS | | | | | | | |
| | DACOGEN | | | | | | | |
| | + | MGI PHARMA INC | 50MG/VIAL | | N21790 001 | May 02, 2006 | May | NEWA |
| | <u>DEFEROXAMINE MESYLATE</u> | | | | | | | |
| | INJECTABLE; INJECTION | | | | | | | |
| | DEFEROXAMINE MESYLATE | | | | | | | |
| AP | | SICOR PHARMS | 500MG/VIAL | | N76806 001 | Mar 31, 2006 | Mar | NEWA |
| AP | | | 2GM/VIAL | | N76806 002 | Mar 31, 2006 | Mar | NEWA |
| | <u>DEMECLOCYCLINE HYDROCHLORIDE</u> | | | | | | | |
| | TABLET; ORAL | | | | | | | |
| | DECLOMYCIN | | | | | | | |
| | | @ GLADES PHARMS LLC | 75MG | | N50261 001 | | Mar | CAHN |
| AB | | | 150MG | | N50261 002 | | Mar | CAHN |
| AB | + | | 300MG | | N50261 003 | | Mar | CAHN |
| | | @ PROTEIN DESIGN LABS | 75MG | | N50261 001 | | Feb | CAHN |
| AB | | | 150MG | | N50261 002 | | Feb | CAHN |
| AB | + | | 300MG | | N50261 003 | | Feb | CAHN |

DESLORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

CLARINEX-D 12 HOUR

| | | | | | | |
|---|----------|-------------|------------|--------------|-----|------|
| + | SCHERING | 2.5MG;120MG | N21313 001 | Feb 01, 2006 | Feb | NEWA |
|---|----------|-------------|------------|--------------|-----|------|

DESMOPRESSIN ACETATE

INJECTABLE; INJECTION

DESMOPRESSIN ACETATE

| | | | | | | |
|---|---------|------------|------------|--------------|-----|------|
| @ | BEDFORD | 0.004MG/ML | N74575 001 | Feb 18, 2000 | Jan | DISC |
|---|---------|------------|------------|--------------|-----|------|

DESMOPRESSIN ACETATE PRESERVATIVE FREE

| | | | | | | |
|---|---------|------------|------------|--------------|-----|------|
| @ | BEDFORD | 0.004MG/ML | N74574 001 | Feb 18, 2000 | Jan | DISC |
|---|---------|------------|------------|--------------|-----|------|

TABLET; ORAL

DESMOPRESSIN ACETATE

| | | | | | | |
|----|--------|-------|------------|--------------|-----|------|
| AB | APOTEX | 0.1MG | N77414 001 | Mar 07, 2006 | Feb | NEWA |
|----|--------|-------|------------|--------------|-----|------|

| | | | | | | |
|----|--|-------|------------|--------------|-----|------|
| AB | | 0.2MG | N77414 002 | Mar 07, 2006 | Feb | NEWA |
|----|--|-------|------------|--------------|-----|------|

| | | | | | | |
|----|-------------|-------|------------|--------------|-----|------|
| AB | TEVA PHARMS | 0.1MG | N77122 001 | Jan 25, 2006 | Jan | NEWA |
|----|-------------|-------|------------|--------------|-----|------|

| | | | | | | |
|----|--|-------|------------|--------------|-----|------|
| AB | | 0.2MG | N77122 002 | Jan 25, 2006 | Jan | NEWA |
|----|--|-------|------------|--------------|-----|------|

DESOGESTREL; ETHINYL ESTRADIOL

TABLET; ORAL-28

MIRCETTE

| | | | | | | | |
|----|---|---------|--------------------------|------------|--------------|-----|------|
| AB | + | DURAMED | 0.15MG,N/A;0.02MG,0.01MG | N20713 001 | Apr 22, 1998 | Feb | CAHN |
|----|---|---------|--------------------------|------------|--------------|-----|------|

| | | | | | | | |
|----|---|--|--------------------------|------------|--------------|-----|------|
| AB | + | | 0.15MG,N/A;0.02MG,0.01MG | N20713 001 | Apr 22, 1998 | Feb | CAHN |
|----|---|--|--------------------------|------------|--------------|-----|------|

DESONIDE

>A> AEROSOL, FOAM; TOPICAL

>A> VERDESO

| | | | | | | | |
|-----|---|-----------|-------|------------|--------------|-----|------|
| >A> | + | CONNETICS | 0.05% | N21978 001 | Sep 19, 2006 | Sep | NEWA |
|-----|---|-----------|-------|------------|--------------|-----|------|

DEXAMETHASONE

ELIXIR; ORAL

DEXAMETHASONE

| | | | | | | | |
|----|---|----------------------|-----------|------------|--|-----|------|
| AA | + | ACTAVIS MID ATLANTIC | 0.5MG/5ML | N84754 001 | | Jul | CAHN |
|----|---|----------------------|-----------|------------|--|-----|------|

| | | | | | | | |
|----|---|--------------------|-----------|------------|--|-----|------|
| AA | + | ALPHARMA US PHARMS | 0.5MG/5ML | N84754 001 | | Jun | CRLD |
|----|---|--------------------|-----------|------------|--|-----|------|

HEXADROL

| | | | | | | |
|---|-----------------|-----------|------------|--|-----|------|
| @ | ORGANON USA INC | 0.5MG/5ML | N12674 001 | | Jun | DISC |
|---|-----------------|-----------|------------|--|-----|------|

MYMETHASONE

| | | | | | | | |
|----|---|--------------|-----------|------------|--------------|-----|------|
| AA | + | MORTON GROVE | 0.5MG/5ML | N88254 001 | Jul 27, 1983 | Jun | CRLD |
|----|---|--------------|-----------|------------|--------------|-----|------|

TABLET; ORAL

DECADRON

| | | | | | | |
|---|-------|-------|------------|--|-----|------|
| @ | MERCK | 0.5MG | N11664 001 | | Aug | DISC |
|---|-------|-------|------------|--|-----|------|

| | | | | | | |
|---|--|--------|------------|--|-----|------|
| @ | | 0.75MG | N11664 002 | | Aug | DISC |
|---|--|--------|------------|--|-----|------|

DEXAMETHASONE

| | | | | | | |
|----|--------|-------|------------|--|-----|------|
| BP | ROXANE | 0.5MG | N84611 001 | | Aug | CTEC |
|----|--------|-------|------------|--|-----|------|

| | | | | | | |
|----|--|--------|------------|--|-----|------|
| BP | | 0.75MG | N84613 001 | | Aug | CTEC |
|----|--|--------|------------|--|-----|------|

HEXADROL

| | | | | | | |
|---|-----------------|-----|------------|--|-----|------|
| @ | ORGANON USA INC | 4MG | N12675 010 | | Jun | DISC |
|---|-----------------|-----|------------|--|-----|------|

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

DEXAMETHASONE

| | | | | | | |
|----|-----------------|----------------------|------------|--------------|-----|------|
| AP | BAXTER HLTHCARE | EQ 10MG PHOSPHATE/ML | N87702 001 | Sep 07, 1982 | Mar | CAHN |
|----|-----------------|----------------------|------------|--------------|-----|------|

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL
 FOCALIN XR
 NOVARTIS 15MG

N21802 004 Aug 01, 2006 Aug NEWA

DEXTROAMPHETAMINE SULFATE

TABLET; ORAL
 DEXTROAMPHETAMINE SULFATE
 @ ENDO PHARMS 5MG

N40299 001 May 13, 1999 Feb DISC

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL
 PROMETH W/ DEXTROMETHORPHAN
 AA + ACTAVIS MID ATLANTIC 15MG/5ML;6.25MG/5ML
 PROMETHAZINE DM
 AA VINTAGE 15MG/5ML;6.25MG/5ML

N88762 001 Oct 31, 1984 Jul CAHN

N40649 001 Feb 14, 2006 Jan NEWA

DIAZEPAM

TABLET; ORAL
 DIAZEPAM
 AB ACTAVIS ELIZABETH 2MG
 AB 5MG
 AB 10MG
 AB VINTAGE PHARMS 2MG
 AB 5MG
 AB 10MG

N70781 001 Mar 19, 1986 Jun CAHN

N70706 001 Mar 19, 1986 Jun CAHN

N70707 001 Mar 19, 1986 Jun CAHN

N77749 001 Mar 31, 2006 Mar NEWA

N77749 002 Mar 31, 2006 Mar NEWA

N77749 003 Mar 31, 2006 Mar NEWA

DICLOFENAC SODIUM

TABLET, DELAYED RELEASE; ORAL
 DICLOFENAC SODIUM
 AB ACTAVIS ELIZABETH 50MG
 AB 75MG
 AB SANDOZ 75MG
 VOLTAREN
 AB + NOVARTIS 75MG
 TABLET, EXTENDED RELEASE; ORAL
 DICLOFENAC SODIUM
 AB ACTAVIS ELIZABETH 100MG

N74514 001 Mar 26, 1996 Jun CAHN

N74514 002 Mar 26, 1996 Jun CAHN

N74394 001 Nov 30, 1995 May CRLD

N19201 003 Jul 28, 1988 May CMFD

N75910 001 Jan 07, 2002 Jun CAHN

DIETHYLPROPION HYDROCHLORIDE

TABLET; ORAL
 DIETHYLPROPION HYDROCHLORIDE
 @ TG UNITED LABS 25MG
 @ 25MG

N88267 001 Aug 25, 1983 May CAHN

N88268 001 Aug 25, 1983 May CAHN

DIFLORASONE DIACETATE

CREAM; TOPICAL
 DIFLORASONE DIACETATE
 BX + ALTANA 0.05%
 FLORONE
 @ PHARMACIA AND UPJOHN 0.05%
 FLORONE E
 @ PHARMACIA AND UPJOHN 0.05%

N76263 001 Dec 20, 2002 Jan CRLD

N17741 001 Jan DISC

N19259 001 Aug 28, 1985 Jan DISC

OINTMENT; TOPICAL

| | | | | | | | | |
|----|------------------------|-------|--------|-----|--------------|-----|------|--|
| | DIFLORASONE DIACETATE | | | | | | | |
| AB | + TARO | 0.05% | N75331 | 001 | May 14, 1999 | Jan | CRLD | |
| | FLORONE | | | | | | | |
| | @ PHARMACIA AND UPJOHN | 0.05% | N17994 | 001 | | Jan | DISC | |
| | PSORCON | | | | | | | |
| | @ PHARMACIA AND UPJOHN | 0.05% | N19260 | 001 | Aug 28, 1985 | Jan | DISC | |

DIFLUNISAL

TABLET; ORAL

| | | | | | | | | |
|----|------------|-------|--------|-----|--------------|-----|------|--|
| | DIFLUNISAL | | | | | | | |
| AB | + TEVA | 500MG | N73673 | 001 | Jul 31, 1992 | Jun | CRLD | |
| | DOLOBID | | | | | | | |
| | @ MERCK | 250MG | N18445 | 001 | Apr 19, 1982 | Jun | DISC | |
| | @ | 500MG | N18445 | 002 | Apr 19, 1982 | Jun | DISC | |

DIGOXIN

INJECTABLE; INJECTION

| | | | | | | | | |
|----|---------|-----------|--------|-----|--------------|-----|------|--|
| | DIGOXIN | | | | | | | |
| AP | SANDOZ | 0.25MG/ML | N40481 | 001 | Aug 21, 2003 | Jan | CAHN | |

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

| | | | | | | | | |
|-----|-------------------------|-------|--------|-----|--------------|-----|------|--|
| | DILTIAZEM HYDROCHLORIDE | | | | | | | |
| AB3 | ACTAVIS ELIZABETH | 120MG | N74984 | 001 | Dec 20, 1999 | Jun | CAHN | |
| AB3 | | 180MG | N74984 | 002 | Dec 20, 1999 | Jun | CAHN | |
| AB3 | | 240MG | N74984 | 003 | Dec 20, 1999 | Jun | CAHN | |
| AB3 | | 300MG | N74984 | 004 | Dec 20, 1999 | Jun | CAHN | |
| AB4 | KV PHARM | 120MG | N76563 | 002 | Sep 12, 2006 | Aug | NEWA | |
| AB4 | | 180MG | N76563 | 003 | Sep 12, 2006 | Aug | NEWA | |
| AB4 | | 240MG | N76563 | 004 | Sep 12, 2006 | Aug | NEWA | |
| AB4 | | 300MG | N76563 | 005 | Sep 12, 2006 | Aug | NEWA | |
| AB4 | | 360MG | N76563 | 006 | Sep 12, 2006 | Aug | NEWA | |
| AB4 | | 420MG | N76563 | 001 | Sep 12, 2006 | Aug | NEWA | |
| | DILTZAC | | | | | | | |
| AB4 | APOTEX INC | 120MG | N76395 | 001 | Feb 01, 2006 | Jan | NEWA | |
| AB4 | | 180MG | N76395 | 002 | Feb 01, 2006 | Jan | NEWA | |
| AB4 | | 240MG | N76395 | 003 | Feb 01, 2006 | Jan | NEWA | |
| AB4 | | 300MG | N76395 | 004 | Feb 01, 2006 | Jan | NEWA | |
| AB4 | | 360MG | N76395 | 005 | Feb 01, 2006 | Jan | NEWA | |
| | TIAZAC | | | | | | | |
| AB4 | + BIOVAIL | 420MG | N20401 | 006 | Oct 16, 1998 | Aug | CFTG | |

DIMETHYL SULFOXIDE

SOLUTION; INTRAVESICAL

| | | | | | | | | |
|----|-------------------|-----|--------|-----|--|-----|------|--|
| | RIMSO-50 | | | | | | | |
| AT | + BIONICHE PHARMA | 50% | N17788 | 001 | | Jul | CAHN | |

DIPYRIDAMOLE

TABLET; ORAL

| | | | | | | | | |
|----|-------------------|------|--------|-----|--------------|-----|------|--|
| | DIPYRIDAMOLE | | | | | | | |
| AB | AMIDE PHARM | 25MG | N40542 | 001 | Apr 21, 2006 | Apr | NEWA | |
| AB | | 50MG | N40542 | 002 | Apr 21, 2006 | Apr | NEWA | |
| AB | | 75MG | N40542 | 003 | Apr 21, 2006 | Apr | NEWA | |
| | @ GLENMARK PHARMA | 25MG | N88999 | 001 | Feb 05, 1991 | Jul | CAHN | |

TABLET; ORAL

| | | | | | | | | | |
|----|-----------------|------|--------|-----|--------------|-----|------|--|--|
| | DIPYRIDAMOLE | | | | | | | | |
| AB | GLENMARK PHARMA | 50MG | N89000 | 001 | Feb 05, 1991 | Jul | CAHN | | |
| | @ | 75MG | N89001 | 001 | Feb 05, 1991 | Jul | CAHN | | |

DISOPYRAMIDE PHOSPHATE

CAPSULE; ORAL

DISOPYRAMIDE PHOSPHATE

| | | | | | | | | | |
|--|---------------|---------------|--------|-----|--------------|-----|------|--|--|
| | @ IVAX PHARMS | EQ 100MG BASE | N70186 | 001 | Nov 18, 1985 | Jan | DISC | | |
| | @ | EQ 150MG BASE | N70187 | 001 | Nov 18, 1985 | Jan | DISC | | |
| | @ SANDOZ | EQ 100MG BASE | N70470 | 001 | Dec 10, 1985 | Jan | DISC | | |
| | @ | EQ 150MG BASE | N70471 | 001 | Dec 10, 1985 | Jan | DISC | | |

DOLASETRON MESYLATE

INJECTABLE; INJECTION

ANZEMET

| | | | | | | | | | |
|---|-------------------|--|--------|-----|--------------|-----|------|--|--|
| + | SANOFI AVENTIS US | EQ 12.5MG BASE/0.625ML (EQ 20MG BASE/ML) | N20624 | 002 | Sep 11, 1997 | Mar | CAIN | | |
| + | | EQ 100MG BASE/5ML (EQ 20MG BASE/ML) | N20624 | 001 | Sep 11, 1997 | Mar | CAIN | | |
| + | | EQ 500MG BASE/25ML (EQ 20MG BASE/ML) | N20624 | 003 | Dec 11, 2001 | Mar | CAIN | | |

TABLET; ORAL

ANZEMET

| | | | | | | | | | |
|---|-------------------|---------------|--------|-----|--------------|-----|------|--|--|
| | SANOFI AVENTIS US | EQ 50MG BASE | N20623 | 001 | Sep 11, 1997 | Mar | CAIN | | |
| + | | EQ 100MG BASE | N20623 | 002 | Sep 11, 1997 | Mar | CAIN | | |

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOPAMINE HYDROCHLORIDE

| | | | | | | | | | |
|--|-------------------|---------|--------|-----|--------------|-----|------|--|--|
| | @ HOSPIRA | 40MG/ML | N74403 | 001 | May 23, 1996 | Apr | DISC | | |
| | @ INTL MEDICATION | 40MG/ML | N18014 | 001 | | Apr | DISC | | |
| | @ SICOR PHARMS | 40MG/ML | N72999 | 001 | Oct 23, 1991 | Apr | DISC | | |
| | @ | 80MG/ML | N73000 | 001 | Oct 23, 1991 | Apr | DISC | | |

DOXAZOSIN MESYLATE

TABLET; ORAL

DOXAZOSIN MESYLATE

| | | | | | | | | | |
|----|-------------------|-------------|--------|-----|--------------|-----|------|--|--|
| AB | ACTAVIS ELIZABETH | EQ 1MG BASE | N75574 | 001 | Oct 18, 2000 | Jun | CAHN | | |
| AB | | EQ 2MG BASE | N75574 | 002 | Oct 18, 2000 | Jun | CAHN | | |
| AB | | EQ 4MG BASE | N75574 | 003 | Oct 18, 2000 | Jun | CAHN | | |
| AB | | EQ 8MG BASE | N75574 | 004 | Oct 18, 2000 | Jun | CAHN | | |

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

| | | | | | | | | | |
|--|-------------|---------------|--------|-----|--------------|-----|------|--|--|
| | @ PAR PHARM | EQ 75MG BASE | N65055 | 004 | Apr 18, 2005 | Mar | DISC | | |
| | @ | EQ 150MG BASE | N65055 | 003 | Jul 15, 2005 | Mar | DISC | | |
| | RANBAXY | EQ 75MG BASE | N65053 | 003 | Sep 10, 2003 | Mar | CTEC | | |

CAPSULE, DELAYED RELEASE; ORAL

ORACEA

| | | | | | | | | | |
|---|-------------------|------|--------|-----|--------------|-----|------|--|--|
| + | COLLAGENEX PHARMS | 40MG | N50805 | 001 | May 26, 2006 | May | NEWA | | |
|---|-------------------|------|--------|-----|--------------|-----|------|--|--|

TABLET; ORAL

DOXYCYCLINE

| | | | | | | | | | |
|----|-----------|--------------|--------|-----|--------------|-----|------|--|--|
| AB | PAR PHARM | EQ 75MG BASE | N65070 | 003 | Dec 30, 2002 | May | CFTG | | |
| | | EQ 75MG BASE | N65070 | 003 | Dec 30, 2002 | Mar | CMFD | | |
| AB | RANBAXY | EQ 50MG BASE | N65356 | 001 | May 31, 2006 | May | NEWA | | |

TABLET; ORAL

DOXYCYCLINE

| | | | | | | |
|----|---------|---------------|------------|--------------|-----|------|
| AB | RANBAXY | EQ 75MG BASE | N65356 002 | May 31, 2006 | May | NEWA |
| AB | | EQ 100MG BASE | N65356 003 | May 31, 2006 | May | NEWA |

DOXYCYCLINE HYCLATE

CAPSULE, COATED PELLETS; ORAL

DORYX

| | | | | | | |
|-----|----------------------|---------------|------------|--------------|-----|------|
| >D> | @ FH FAULDING CO LTD | EQ 75MG BASE | N50582 002 | Aug 13, 2001 | Sep | CAHN |
| >D> | @ | EQ 100MG BASE | N50582 001 | Jul 22, 1985 | Sep | CAHN |

CAPSULE, DELAYED RELEASE; ORAL

DORYX

| | | | | | | |
|-----|----------------------|---------------|------------|--------------|-----|------|
| >D> | @ FH FAULDING CO LTD | EQ 75MG BASE | N50582 002 | Aug 13, 2001 | Sep | CAHN |
| | @ | EQ 75MG BASE | N50582 002 | Aug 13, 2001 | Apr | DISC |
| >D> | @ | EQ 100MG BASE | N50582 001 | Jul 22, 1985 | Sep | CAHN |
| | @ | EQ 100MG BASE | N50582 001 | Jul 22, 1985 | Apr | DISC |
| >A> | @ MAYNE PHARMA INTL | EQ 75MG BASE | N50582 002 | Aug 13, 2001 | Sep | CAHN |
| >A> | @ | EQ 100MG BASE | N50582 001 | Jul 22, 1985 | Sep | CAHN |
| | @ WARNER CHILCOTT | EQ 100MG BASE | N62653 001 | Oct 30, 1985 | Apr | DISC |

DOXYCYCLINE HYCLATE

SANDOZ

| | | | | | | |
|--|--|--------------|------------|--------------|-----|------|
| | | EQ 75MG BASE | N65281 001 | Dec 21, 2005 | Apr | CTEC |
|--|--|--------------|------------|--------------|-----|------|

| | | | | | | |
|---|--|---------------|------------|--------------|-----|------|
| + | | EQ 100MG BASE | N65281 002 | Dec 21, 2005 | Apr | CRLD |
|---|--|---------------|------------|--------------|-----|------|

TABLET; ORAL

DOXYCYCLINE HYCLATE

| | | | | | | |
|----|-----------|--------------|------------|--------------|-----|------|
| AB | PAR PHARM | EQ 20MG BASE | N65287 001 | Feb 28, 2006 | Feb | NEWA |
|----|-----------|--------------|------------|--------------|-----|------|

DROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL

YAZ

| | | | | | | |
|---|-------------|------------|------------|--------------|-----|------|
| + | BERLEX | 3MG;0.02MG | N21676 001 | Mar 16, 2006 | Jun | CAHN |
| + | BERLEX LABS | 3MG;0.02MG | N21676 001 | Mar 16, 2006 | Mar | NEWA |

DYPHYLLINE

TABLET; ORAL

DILOR

| | | | | | | |
|--|---------------|-------|------------|--|-----|------|
| | @ SAVAGE LABS | 200MG | N84514 001 | | Jun | DISC |
|--|---------------|-------|------------|--|-----|------|

DILOR-400

| | | | | | | |
|--|---------------|-------|------------|--|-----|------|
| | @ SAVAGE LABS | 400MG | N84751 001 | | Jun | DISC |
|--|---------------|-------|------------|--|-----|------|

LUFYLLIN

MEDPOINTE PHARM HLC

| | | | | | | |
|--|--|-------|------------|--|-----|------|
| | | 200MG | N84566 001 | | Jun | CTEC |
|--|--|-------|------------|--|-----|------|

| | | | | | | |
|---|--|-------|------------|--|-----|------|
| + | | 400MG | N84566 002 | | Jun | CRLD |
|---|--|-------|------------|--|-----|------|

EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

ATRIPLA

| | | | | | | |
|---|--------|-------------------|------------|--------------|-----|------|
| + | GILEAD | 600MG;200MG;300MG | N21937 001 | Jul 12, 2006 | Jul | NEWA |
|---|--------|-------------------|------------|--------------|-----|------|

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

XYLOCAINE W/ EPINEPHRINE

| | | | | | | | |
|----|---|--------------------|-----------------|------------|--------------|-----|------|
| AP | + | ABRAXIS BIOSCIENCE | 0.005MG/ML;0.5% | N06488 012 | | Jul | CAHN |
| AP | + | | 0.005MG/ML;1% | N06488 018 | Nov 13, 1986 | Jul | CAHN |
| AP | + | | 0.005MG/ML;1.5% | N06488 017 | | Jul | CAHN |
| AP | + | | 0.005MG/ML;2% | N06488 019 | Nov 13, 1986 | Jul | CAHN |
| AP | + | | 0.01MG/ML;1% | N06488 004 | | Jul | CAHN |

INJECTABLE; INJECTION

XYLOCAINE W/ EPINEPHRINE

| | | | | | | |
|---|--------------------|--------------|------------|--|-----|------|
| @ | ABRAXIS BIOSCIENCE | 0.01MG/ML;2% | N06488 003 | | Jul | CAHN |
| @ | | 0.02MG/ML;2% | N06488 005 | | Jul | CAHN |
| + | DENTSPLY PHARM | 0.02MG/ML;2% | N21381 002 | | Mar | CRLD |

PATCH; IONTOPHORESIS, TOPICAL

LIDOSITE TOPICAL SYSTEM KIT

| | | | | | | |
|---|---------|--------------------------|------------|--------------|-----|------|
| + | VYTERIS | 1.05MG/PATCH;100MG/PATCH | N21504 001 | May 06, 2004 | May | CDFR |
|---|---------|--------------------------|------------|--------------|-----|------|

SOLUTION; IONTOPHORESIS, TOPICAL

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

| | | | | | | |
|---|------|--------------|------------|--------------|-----|------|
| + | EMPI | 0.01MG/ML;2% | N21486 001 | Oct 26, 2004 | May | CDFR |
|---|------|--------------|------------|--------------|-----|------|

EPIRUBICIN HYDROCHLORIDE

>A> INJECTABLE; IV (INFUSION)

>A> EPIRUBICIN HYDROCHLORIDE

| | | | | | | | |
|-----|---|------------------|------------|------------|--------------|-----|------|
| >A> | + | MAYNE PHARMA USA | 50MG/VIAL | N50807 001 | Sep 15, 2006 | Sep | NEWA |
| >A> | + | | 200MG/VIAL | N50807 002 | Sep 15, 2006 | Sep | NEWA |

ERYTHROMYCIN

SOLUTION; TOPICAL

A/T/S

| | | | | | | |
|----|-------------------|----|------------|--------------|-----|------|
| AT | TARO PHARMS NORTH | 2% | N62405 001 | Nov 18, 1982 | Feb | CAHN |
|----|-------------------|----|------------|--------------|-----|------|

SWAB; TOPICAL

ERYTHROMYCIN

| | | | | | | |
|----|--------|----|------------|--------------|-----|------|
| AT | ALTANA | 2% | N65320 001 | Jul 25, 2006 | Jul | NEWA |
|----|--------|----|------------|--------------|-----|------|

ERYTHROMYCIN ETHYLSUCCINATE

TABLET, CHEWABLE; ORAL

E.E.S.

| | | | | | | |
|---|--------|---------------|------------|--|-----|------|
| @ | ABBOTT | EQ 200MG BASE | N50297 002 | | Aug | DISC |
|---|--------|---------------|------------|--|-----|------|

ERYPED

| | | | | | | |
|---|--------|---------------|------------|--------------|-----|------|
| @ | ABBOTT | EQ 200MG BASE | N50297 003 | Jul 05, 1988 | Aug | DISC |
|---|--------|---------------|------------|--------------|-----|------|

ESCITALOPRAM OXALATE

TABLET; ORAL

ESCITALOPRAM OXALATE

| | | | | | | |
|----|-------------|------|------------|--------------|-----|------|
| AB | IVAX PHARMS | 5MG | N76765 001 | May 22, 2006 | May | NEWA |
| AB | | 10MG | N76765 002 | May 22, 2006 | May | NEWA |
| AB | | 20MG | N76765 003 | May 22, 2006 | May | NEWA |

LEXAPRO

| | | | | | | |
|----|-------------|------|------------|--------------|-----|------|
| AB | FOREST LABS | 5MG | N21323 001 | Aug 14, 2002 | May | CFTG |
| AB | | 10MG | N21323 002 | Aug 14, 2002 | May | CFTG |
| AB | + | 20MG | N21323 003 | Aug 14, 2002 | May | CFTG |

ESTAZOLAM

TABLET; ORAL

ESTAZOLAM

| | | | | | | |
|----|-----------|-----|------------|--------------|-----|------|
| AB | PAR PHARM | 1MG | N74826 001 | Jul 03, 1997 | Apr | CAHN |
| AB | | 2MG | N74826 002 | Jul 03, 1997 | Apr | CAHN |

ESTRADIOL

FILM, EXTENDED RELEASE; TRANSDERMAL

CLIMARA

| | | | | | | |
|-----|--------|---------------|------------|--------------|-----|------|
| AB2 | BERLEX | 0.025MG/24HR | N20375 004 | Mar 05, 1999 | Aug | CRLD |
| AB | | 0.0375MG/24HR | N20375 005 | May 27, 2003 | Aug | CRLD |

FILM, EXTENDED RELEASE; TRANSDERMAL

CLIMARA

| | | | | | | | |
|-----|---|--------|---------------|------------|--------------|-----|------|
| AB | + | BERLEX | 0.0375MG/24HR | N20375 005 | May 27, 2003 | Jul | CFTG |
| AB2 | | | 0.05MG/24HR | N20375 001 | Dec 22, 1994 | Aug | CRLD |
| AB | | | 0.06MG/24HR | N20375 006 | May 27, 2003 | Aug | CRLD |
| AB | + | | 0.06MG/24HR | N20375 006 | May 27, 2003 | Jul | CFTG |
| AB2 | | | 0.075MG/24HR | N20375 003 | Mar 23, 1998 | Aug | CRLD |

ESTRADERM

| | | | | | | | |
|----|--|----------|-------------|------------|--------------|-----|------|
| BX | | NOVARTIS | 0.05MG/24HR | N19081 002 | Sep 10, 1986 | Aug | CRLD |
|----|--|----------|-------------|------------|--------------|-----|------|

ESTRADIOL

| | | | | | | | |
|----|--|--------------------|---------------|------------|--------------|-----|------|
| AB | | MYLAN TECHNOLOGIES | 0.0375MG/24HR | N75182 004 | Jul 20, 2006 | Jul | NEWA |
| AB | | | 0.06MG/24HR | N75182 005 | Jul 20, 2006 | Jul | NEWA |

MENOSTAR

| | | | | | | | |
|---|--|--------|--------------|------------|--------------|-----|------|
| + | | BERLEX | 0.014MG/24HR | N21674 001 | Jun 08, 2004 | Jun | CAHN |
|---|--|--------|--------------|------------|--------------|-----|------|

VIVELLE-DOT

| | | | | | | | |
|-----|--|----------|---------------|------------|--------------|-----|------|
| BX | | NOVARTIS | 0.025MG/24HR | N20538 009 | May 03, 2002 | Aug | CRLD |
| BX | | | 0.0375MG/24HR | N20538 005 | Jan 08, 1999 | Aug | CRLD |
| AB1 | | | 0.05MG/24HR | N20538 006 | Jan 08, 1999 | Aug | CRLD |
| BX | | | 0.075MG/24HR | N20538 007 | Jan 08, 1999 | Aug | CRLD |

GEL; TOPICAL

ESTROGEL

| | | | | | | | |
|---|--|--------|-------|------------|--------------|-----|------|
| @ | | ASCEND | 0.06% | N21166 001 | Feb 09, 2004 | Jan | CAHN |
|---|--|--------|-------|------------|--------------|-----|------|

GEL, METERED; TOPICAL

ESTROGEL

| | | | | | | | |
|---|--|--------|-------|------------|--------------|-----|------|
| + | | ASCEND | 0.06% | N21166 002 | Feb 09, 2004 | Jan | CAHN |
|---|--|--------|-------|------------|--------------|-----|------|

TABLET; ORAL

ESTRADIOL

| | | | | | | | |
|---|--|---------------|-------|------------|--------------|-----|------|
| @ | | RADIUS PHARMS | 0.5MG | N40275 001 | Dec 29, 1998 | Aug | CAHN |
| @ | | | 1MG | N40275 002 | Dec 29, 1998 | Aug | CAHN |
| @ | | | 2MG | N40275 003 | Dec 29, 1998 | Aug | CAHN |

ESTRADIOL HEMIHYDRATE

EMULSION; TOPICAL

ESTRASORB

| | | | | | | | |
|---|--|---------------|-------|------------|--------------|-----|------|
| + | | ESPRIT PHARMA | 0.25% | N21371 001 | Oct 09, 2003 | Feb | CAHN |
|---|--|---------------|-------|------------|--------------|-----|------|

ESTROGENS, CONJUGATED SYNTHETIC B

TABLET; ORAL

ENJUVIA

| | | | | | | | |
|---|--|---------|---------|------------|--------------|-----|------|
| | | DURAMED | 0.3MG | N21443 001 | Dec 20, 2004 | Mar | CMFD |
| | | | 0.45MG | N21443 002 | Dec 20, 2004 | Mar | CMFD |
| | | | 0.625MG | N21443 003 | May 10, 2004 | Mar | CMFD |
| + | | | 1.25MG | N21443 004 | May 10, 2004 | Mar | CMFD |

ETHACRYNATE SODIUM

INJECTABLE; INJECTION

EDECIN

| | | | | | | | |
|---|--|------|-------------------|------------|--|-----|------|
| + | | ATON | EQ 50MG BASE/VIAL | N16093 001 | | Jul | CAHN |
|---|--|------|-------------------|------------|--|-----|------|

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL

QUASENSE

| | | | | | | | |
|----|---|-------------|---------------|------------|--------------|-----|------|
| AB | | WATSON LABS | 0.03MG;0.15MG | N77101 001 | Sep 06, 2006 | Aug | NEWA |
| AB | + | DURAMED | 0.03MG;0.15MG | N21544 001 | Sep 05, 2003 | Aug | CFTG |

| | | | | | | | | | |
|---|--------------------|---|--------|-----|--------------|-----|------|--|--|
| TABLET; ORAL | | | | | | | | | |
| SEASONIQUE | | | | | | | | | |
| | + DURAMED RES | 0.03MG,0.01MG;0.15MG,N/A | N21840 | 001 | May 25, 2006 | May | NEWA | | |
| TABLET; ORAL-28 | | | | | | | | | |
| LEVONORGESTREL AND ETHINYL ESTRADIOL | | | | | | | | | |
| AB2 | WATSON LABS | 0.02MG;0.1MG | N77681 | 001 | May 31, 2006 | May | NEWA | | |
| <u>ETHINYL ESTRADIOL; NORETHINDRONE</u> | | | | | | | | | |
| TABLET; ORAL-21 | | | | | | | | | |
| BALZIVA-21 | | | | | | | | | |
| | BARR | 0.035MG;0.4MG | N76198 | 001 | Apr 22, 2004 | Mar | CRLD | | |
| TABLET; ORAL-28 | | | | | | | | | |
| OVCON-35 | | | | | | | | | |
| AB | + WARNER CHILCOTT | 0.035MG;0.4MG | N17716 | 001 | | Mar | CRLD | | |
| <u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE</u> | | | | | | | | | |
| TABLET; ORAL | | | | | | | | | |
| LOESTRIN 24 FE | | | | | | | | | |
| | + WARNER CHILCOTT | 0.02MG;1MG | N21871 | 001 | Feb 17, 2006 | Feb | NEWA | | |
| <u>ETHINYL ESTRADIOL; NORGESTIMATE</u> | | | | | | | | | |
| TABLET; ORAL-28 | | | | | | | | | |
| NORGESTIMATE AND ETHINYL ESTRADIOL | | | | | | | | | |
| AB | WATSON LABS | 0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG | N76626 | 001 | Aug 17, 2006 | Aug | NEWA | | |
| AB | | 0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG | N76626 | 001 | Aug 17, 2006 | Aug | NEWA | | |
| AB | | 0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG | N76626 | 001 | Aug 17, 2006 | Aug | NEWA | | |
| AB | | 0.035MG;0.25MG | N76627 | 001 | Aug 17, 2006 | Aug | NEWA | | |
| <u>ETODOLAC</u> | | | | | | | | | |
| CAPSULE; ORAL | | | | | | | | | |
| ETODOLAC | | | | | | | | | |
| | @ ENDO PHARMS | 200MG | N74842 | 001 | Jul 17, 1997 | Feb | DISC | | |
| | @ | 300MG | N74842 | 002 | Jul 17, 1997 | Feb | DISC | | |
| AB | + TARO | 300MG | N75078 | 002 | Apr 30, 1998 | Mar | CRLD | | |
| LODINE | | | | | | | | | |
| | @ WYETH PHARMS INC | 300MG | N18922 | 003 | Jan 31, 1991 | Mar | DISC | | |
| TABLET; ORAL | | | | | | | | | |
| ETODOLAC | | | | | | | | | |
| AB | ACTAVIS ELIZABETH | 400MG | N74819 | 001 | Feb 28, 1997 | Jun | CAHN | | |
| | @ ENDO PHARMS | 400MG | N74841 | 001 | Jun 27, 1997 | Feb | DISC | | |
| <u>ETONOGESTREL</u> | | | | | | | | | |
| IMPLANT; IMPLANTATION | | | | | | | | | |
| IMPLANON | | | | | | | | | |
| | + ORGANON USA INC | 68MG/IMPLANT | N21529 | 001 | Jul 17, 2006 | Jul | NEWA | | |
| <u>ETOPOSIDE</u> | | | | | | | | | |
| INJECTABLE; INJECTION | | | | | | | | | |
| TOPOSAR | | | | | | | | | |
| | @ SICOR PHARMS | 20MG/ML | N74166 | 001 | Feb 27, 1995 | Jun | DISC | | |

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

| | | | | | | |
|----|-------------------|------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 20MG | N75650 001 | Sep 14, 2001 | Jun | CAHN |
| AB | | 40MG | N75650 002 | Sep 14, 2001 | Jun | CAHN |

FENOFIBRATE

CAPSULE; ORAL

ANTARA (MICRONIZED)

OSCIENT

| | | | | | | |
|--|---|-------|------------|--------------|-----|------|
| | | 43MG | N21695 001 | Nov 30, 2004 | Aug | CAHN |
| | @ | 87MG | N21695 002 | Nov 30, 2004 | Aug | CAHN |
| | + | 130MG | N21695 003 | Nov 30, 2004 | Aug | CAHN |

LIPOFEN

CIPHER

| | | | | | | |
|--|---|-------|------------|--------------|-----|------|
| | | 50MG | N21612 001 | Jan 11, 2006 | Jan | NEWA |
| | | 100MG | N21612 002 | Jan 11, 2006 | Jan | NEWA |
| | + | 150MG | N21612 003 | Jan 11, 2006 | Jan | NEWA |

TABLET; ORAL

FENOFIBRATE

| | | | | | | | |
|----|---|----------|-------|------------|--------------|-----|------|
| AB | + | TEVA | 160MG | N76433 002 | May 13, 2005 | Jan | CRLD |
| | | TRICOR | | | | | |
| | | @ ABBOTT | 54MG | N21203 001 | Sep 04, 2001 | Jan | DISC |
| | | @ | 160MG | N21203 003 | Sep 04, 2001 | Jan | DISC |

FENOLDOPAM MESYLATE

INJECTABLE; INJECTION

FENOLDOPAM MESYLATE

| | | | | | | | |
|----|--|--------|-----------------|------------|--------------|-----|------|
| AP | | SANDOZ | EQ 10MG BASE/ML | N77155 001 | Feb 15, 2005 | Jan | CAHN |
|----|--|--------|-----------------|------------|--------------|-----|------|

FENOPROFEN CALCIUM

TABLET; ORAL

FENOPROFEN CALCIUM

| | | | | | | | |
|----|--|--------------------|---------------|------------|--------------|-----|------|
| AB | | ACTAVIS ELIZABETH | EQ 600MG BASE | N72274 001 | May 02, 1988 | Jun | CAHN |
| | | @ CLONMEL HLTHCARE | EQ 600MG BASE | N72326 001 | Aug 17, 1988 | Jan | DISC |

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL

FENTANYL

| | | | | | | | |
|----|--|----------------|-----------|------------|--------------|-----|------|
| AB | | LAVIPHARM LABS | 25UGM/HR | N77051 001 | Aug 04, 2006 | Jul | NEWA |
| AB | | | 50UGM/HR | N77051 002 | Aug 04, 2006 | Jul | NEWA |
| AB | | | 75UGM/HR | N77051 003 | Aug 04, 2006 | Jul | NEWA |
| AB | | | 100UGM/HR | N77051 004 | Aug 04, 2006 | Jul | NEWA |

FENTANYL CITRATE

>A> TABLET; BUCCAL

>A> FENTORA

| | | | | | | | |
|-----|---|----------|---------------|------------|--------------|-----|------|
| >A> | | CEPHALON | EQ 0.1MG BASE | N21947 001 | Sep 25, 2006 | Sep | NEWA |
| >A> | | | EQ 0.2MG BASE | N21947 002 | Sep 25, 2006 | Sep | NEWA |
| >A> | + | | EQ 0.4MG BASE | N21947 003 | Sep 25, 2006 | Sep | NEWA |
| >A> | | | EQ 0.6MG BASE | N21947 004 | Sep 25, 2006 | Sep | NEWA |
| >A> | | | EQ 0.8MG BASE | N21947 005 | Sep 25, 2006 | Sep | NEWA |

TROCHE/LOZENGE; ORAL

FENTANYL

@ CEPHALON

@

| | | | | | | |
|--|--|---------------|------------|--------------|-----|------|
| | | EQ 0.1MG BASE | N20195 007 | Oct 30, 1995 | Jul | CAHN |
| | | EQ 0.2MG BASE | N20195 001 | Oct 04, 1993 | Jul | CAHN |

TROCHE/LOZENGE; ORAL

FENTANYL

| | | | | | | |
|---|----------|---------------|------------|--------------|-----|------|
| @ | CEPHALON | EQ 0.3MG BASE | N20195 002 | Oct 04, 1993 | Jul | CAHN |
| @ | | EQ 0.4MG BASE | N20195 003 | Oct 04, 1993 | Jul | CAHN |

TROCHE/LOZENGE; TRANSMUCOSAL

ACTIQ (SUGAR-FREE)

| | | | | | | |
|---|----------|---------------|------------|--------------|-----|------|
| | CEPHALON | EQ 0.2MG BASE | N20747 001 | Nov 04, 1998 | Mar | CTNA |
| + | | EQ 0.4MG BASE | N20747 002 | Nov 04, 1998 | Mar | CTNA |
| | | EQ 0.6MG BASE | N20747 003 | Nov 04, 1998 | Mar | CTNA |
| | | EQ 0.8MG BASE | N20747 004 | Nov 04, 1998 | Mar | CTNA |
| | | EQ 1.2MG BASE | N20747 005 | Nov 04, 1998 | Mar | CTNA |
| | | EQ 1.6MG BASE | N20747 006 | Nov 04, 1998 | Mar | CTNA |

FEXOFENADINE HYDROCHLORIDE

TABLET; ORAL

FEXOFENADINE HYDROCHLORIDE

| | | | | | | |
|----|--------------------|-------|------------|--------------|-----|------|
| AB | DR REDDYS LABS LTD | 30MG | N76502 001 | Apr 11, 2006 | Mar | NEWA |
| AB | | 60MG | N76502 002 | Apr 11, 2006 | Mar | NEWA |
| AB | | 180MG | N76502 003 | Apr 11, 2006 | Mar | NEWA |

FINASTERIDE

TABLET; ORAL

FINASTERIDE

| | | | | | | |
|----|--------------------|-----|------------|--------------|-----|------|
| AB | DR REDDYS LABS INC | 1MG | N76436 001 | Jul 28, 2006 | Jul | NEWA |
| AB | IVAX PHARMS | 5MG | N76340 001 | Jun 19, 2006 | Jun | NEWA |
| AB | PROPECIA | | | | | |
| AB | + MERCK | 1MG | N20788 001 | Dec 19, 1997 | Jul | CFTG |
| AB | PROSCAR | | | | | |
| AB | + MERCK | 5MG | N20180 001 | Jun 19, 1992 | Jun | CFTG |

FLUCONAZOLE

TABLET; ORAL

FLUCONAZOLE

| | | | | | | |
|----|-----------------|-------|------------|--------------|-----|------|
| AB | GLENMARK PHARMA | 50MG | N77253 001 | Jan 25, 2006 | Jan | NEWA |
| AB | | 100MG | N77253 002 | Jan 25, 2006 | Jan | NEWA |
| AB | | 150MG | N77253 003 | Jan 25, 2006 | Jan | NEWA |
| AB | | 200MG | N77253 004 | Jan 25, 2006 | Jan | NEWA |

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

| | | | | | | |
|----|--------|----------------------|------------|--------------|-----|------|
| AP | SANDOZ | 1MG/10ML (0.1MG/ML) | N77071 002 | May 03, 2005 | Jan | CAHN |
| AP | | 0.5MG/5ML (0.1MG/ML) | N77071 001 | May 03, 2005 | Jan | CAHN |

FLUNISOLIDE

AEROSOL, METERED; INHALATION

AEROSPAN HFA

| | | | | | | |
|---|-------------|-------------------|------------|--------------|-----|------|
| + | FOREST LABS | EQ 78UGM BASE/INH | N21247 001 | Jan 27, 2006 | Jan | NEWA |
|---|-------------|-------------------|------------|--------------|-----|------|

SPRAY, METERED; NASAL

FLUNISOLIDE

| | | | | | | |
|----|-------------------|---------------|------------|--------------|-----|------|
| AB | + BAUSCH AND LOMB | 0.025MG/SPRAY | N74805 001 | Feb 20, 2002 | Jul | CTEC |
| AB | QPHARMA | 0.025MG/SPRAY | N77704 001 | Aug 03, 2006 | Jul | NEWA |

FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE

| | | | | | | | |
|-----|------------------------------|-------|--------|-----|--------------|-----|------|
| AB1 | ACTAVIS MID ATLANTIC | 0.05% | N73085 | 001 | Feb 14, 1992 | Jun | CAHN |
| | FLUOCINONIDE EMULSIFIED BASE | | | | | | |
| AB2 | ACTAVIS MID ATLANTIC | 0.05% | N74204 | 001 | Jun 13, 1995 | Jun | CAHN |
| | SOLUTION; TOPICAL | | | | | | |
| | FLUOCINONIDE | | | | | | |
| AT | ACTAVIS MID ATLANTIC | 0.05% | N71535 | 001 | Dec 02, 1988 | Jul | CAHN |

FLUORESCEIN SODIUM

INJECTABLE; INTRAVENOUS

FLUORESCITE

| | | | | | | | |
|---|-----------|--------------------------------------|--------|-----|--------------|-----|------|
| + | ALCON | EQ 500MG BASE/5ML (EQ 100MG BASE/ML) | N21980 | 001 | Mar 28, 2006 | May | CAHN |
| + | ALCON RES | EQ 500MG BASE/5ML (EQ 100MG BASE/ML) | N21980 | 001 | Mar 28, 2006 | Mar | NEWA |

FLUOROURACIL

INJECTABLE; INJECTION

ADRUCIL

| | | | | | | | |
|----|----------------|---------|--------|-----|--------------|-----|------|
| | @ SICOR PHARMS | 50MG/ML | N40023 | 001 | Oct 18, 1991 | Jun | DISC |
| AP | + | 50MG/ML | N40023 | 001 | Oct 18, 1991 | Mar | CRLD |
| | @ | 50MG/ML | N81225 | 001 | Aug 28, 1991 | Jun | DISC |

FLUOROURACIL

| | | | | | | | | |
|----|---------------|-------------------|---------|--------|--------------|--------------|------|------|
| AP | + | AM PHARM PARTNERS | 50MG/ML | N40278 | 001 | Sep 30, 1998 | Mar | CRLD |
| AP | + | | 50MG/ML | N40279 | 001 | Sep 30, 1998 | Mar | CRLD |
| AP | + | | 50MG/ML | N40291 | 001 | Mar 24, 1999 | Mar | CRLD |
| AP | + | | 50MG/ML | N40379 | 001 | Nov 15, 2000 | Mar | CRLD |
| | @ BEDFORD | 50MG/ML | N89508 | 001 | Jan 26, 1988 | Apr | DISC | |
| AP | + | | 50MG/ML | N89508 | 001 | Jan 26, 1988 | Mar | CRLD |
| AP | + | SICOR PHARMS | 50MG/ML | N40333 | 001 | Jan 27, 2000 | Mar | CRLD |
| AP | + | | 50MG/ML | N40334 | 001 | Feb 25, 2000 | Mar | CRLD |
| AP | + | STERIS | 50MG/ML | N87792 | 001 | Oct 13, 1982 | Mar | CRLD |
| | @ WATSON LABS | 50MG/ML | N87792 | 001 | Oct 13, 1982 | Apr | DISC | |

FLUOXETINE HYDROCHLORIDE

SOLUTION; ORAL

FLUOXETINE

| | | | | | | | |
|----|--------------------------|------------------|--------|-----|--------------|-----|------|
| AA | ACTAVIS MID ATLANTIC | EQ 20MG BASE/5ML | N75690 | 001 | Jan 31, 2002 | Jul | CAHN |
| | TABLET; ORAL | | | | | | |
| | FLUOXETINE HYDROCHLORIDE | | | | | | |
| + | IVAX PHARMS | EQ 40MG BASE | N75865 | 003 | Aug 30, 2004 | Aug | CRLD |
| | SARAFEM | | | | | | |
| | WARNER CHILCOTT | EQ 10MG BASE | N21860 | 001 | May 19, 2006 | May | NEWA |
| | | EQ 15MG BASE | N21860 | 002 | May 19, 2006 | May | NEWA |
| + | | EQ 20MG BASE | N21860 | 003 | May 19, 2006 | May | NEWA |

FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

FLUPHENAZINE HYDROCHLORIDE

| | | | | | | | |
|---|---------------|--------|--------|-----|--------------|-----|------|
| + | PHARM ASSOC | 5MG/ML | N74725 | 001 | Sep 16, 1996 | Aug | CRLD |
| | @ TEVA PHARMS | 5MG/ML | N73058 | 001 | Aug 30, 1991 | Aug | DISC |
| | PROLIXIN | | | | | | |
| | @ APOTHECON | 5MG/ML | N70533 | 001 | Nov 07, 1985 | Jul | DISC |

ELIXIR; ORAL

FLUPHENAZINE HYDROCHLORIDE

| | | | | | | |
|---|---------------|-----------|------------|--------------|-----|------|
| + | PHARM ASSOC | 2.5MG/5ML | N40146 001 | Aug 21, 1996 | Apr | CRLD |
| | @ TEVA PHARMS | 2.5MG/5ML | N81310 001 | Apr 29, 1993 | Apr | DISC |
| | PROLIXIN | | | | | |
| | @ APOTHECON | 2.5MG/5ML | N12145 003 | | Apr | DISC |

FLURBIPROFEN SODIUM

SOLUTION/DROPS; OPHTHALMIC

FLURBIPROFEN SODIUM

| | | | | | | |
|----|-----------------|-------|------------|--------------|-----|------|
| AT | BAUSCH AND LOMB | 0.03% | N74447 001 | Jan 04, 1995 | Aug | CAHN |
|----|-----------------|-------|------------|--------------|-----|------|

FLUTAMIDE

CAPSULE; ORAL

EULEXIN

| | | | | | | |
|--|------------|-------|------------|--------------|-----|------|
| | @ SCHERING | 125MG | N18554 001 | Jan 27, 1989 | May | DISC |
|--|------------|-------|------------|--------------|-----|------|

FLUTAMIDE

| | | | | | | |
|----|-----------|-------|------------|--------------|-----|------|
| AB | PAR PHARM | 125MG | N75298 001 | Sep 18, 2001 | Apr | CAHN |
| AB | + SANDOZ | 125MG | N75818 001 | Sep 18, 2001 | May | CRLD |

FLUTICASONE PROPIONATE

CREAM; TOPICAL

FLUTICASONE PROPIONATE

| | | | | | | |
|----|--------------|-------|------------|--------------|-----|------|
| AB | G AND W LABS | 0.05% | N77055 001 | Jun 30, 2006 | Jun | NEWA |
|----|--------------|-------|------------|--------------|-----|------|

OINTMENT; TOPICAL

FLUTICASONE PROPIONATE

| | | | | | | |
|----|--------------|--------|------------|--------------|-----|------|
| AB | G AND W LABS | 0.005% | N77168 001 | Mar 03, 2006 | Feb | NEWA |
|----|--------------|--------|------------|--------------|-----|------|

SPRAY, METERED; NASAL

FLONASE

| | | | | | | |
|----|-------------------|--------------|------------|--------------|-----|------|
| AB | + GLAXOSMITHKLINE | 0.05MG/SPRAY | N20121 001 | Oct 19, 1994 | Feb | CFTG |
|----|-------------------|--------------|------------|--------------|-----|------|

FLUTICASONE PROPIONATE

| | | | | | | |
|----|--------|--------------|------------|--------------|-----|------|
| AB | ROXANE | 0.05MG/SPRAY | N76504 001 | Feb 22, 2006 | Feb | NEWA |
|----|--------|--------------|------------|--------------|-----|------|

FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE

AEROSOL, METERED; INHALATION

ADVAIR HFA

| | | | | | | |
|---|-----------------|---------------------------------|------------|--------------|-----|------|
| + | GLAXOSMITHKLINE | 0.045MG/INH;EQ 0.021MG BASE/INH | N21254 001 | Jun 08, 2006 | Jun | NEWA |
| + | | 0.115MG/INH;EQ 0.021MG BASE/INH | N21254 002 | Jun 08, 2006 | Jun | NEWA |
| + | | 0.23MG/INH;EQ 0.021MG BASE/INH | N21254 003 | Jun 08, 2006 | Jun | NEWA |

FLUVOXAMINE MALEATE

TABLET; ORAL

FLUVOXAMINE MALEATE

| | | | | | | |
|----|-------------------|-------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 25MG | N75901 001 | Dec 28, 2000 | Jun | CAHN |
| AB | | 50MG | N75901 002 | Dec 28, 2000 | Jun | CAHN |
| AB | | 100MG | N75901 003 | Dec 28, 2000 | Jun | CAHN |
| AB | CARACO | 25MG | N75900 001 | Feb 23, 2006 | Feb | NEWA |
| AB | | 50MG | N75900 002 | Feb 23, 2006 | Feb | NEWA |
| AB | | 100MG | N75900 003 | Feb 23, 2006 | Feb | NEWA |

FOLLITROPIN ALFA/BETA

INJECTABLE; SUBCUTANEOUS

FOLLISTIM AQ

| | | | | | | |
|---|-------------------|---------------|------------|--------------|-----|------|
| | @ ORGANON USA INC | 150 IU/0.18ML | N21211 003 | Feb 11, 2004 | Jul | DISC |
| + | | 300 IU/0.36ML | N21211 001 | Mar 23, 2004 | Jul | CMFD |

INJECTABLE; SUBCUTANEOUS

FOLLISTIM AQ

| | | | | | |
|-------------------|---------------|------------|--------------|-----|------|
| @ ORGANON USA INC | 300 IU/0.36ML | N21211 001 | Mar 23, 2004 | Jun | DISC |
|-------------------|---------------|------------|--------------|-----|------|

FOSCARNET SODIUM

INJECTABLE; INJECTION

FOSCARNET SODIUM

| | | | | | | |
|----|---------|-------------|------------|--------------|-----|------|
| AP | HOSPIRA | 2.4GM/100ML | N77174 001 | May 31, 2005 | Feb | CAHN |
|----|---------|-------------|------------|--------------|-----|------|

FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

| | | | | | | |
|----|--------|------|------------|--------------|-----|------|
| AB | COBALT | 10MG | N77531 001 | Aug 31, 2006 | Aug | NEWA |
| AB | | 20MG | N77531 002 | Aug 31, 2006 | Aug | NEWA |
| AB | | 40MG | N77531 003 | Aug 31, 2006 | Aug | NEWA |

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

| | | | | | | |
|----|----------|-------------|------------|--------------|-----|------|
| AB | GENPHARM | 10MG;12.5MG | N77705 001 | Aug 14, 2006 | Aug | NEWA |
| AB | | 20MG;12.5MG | N77705 002 | Aug 14, 2006 | Aug | NEWA |
| AB | TEVA | 10MG;12.5MG | N76945 001 | Jul 05, 2006 | Jun | NEWA |
| AB | | 20MG;12.5MG | N76945 002 | Jul 05, 2006 | Jun | NEWA |

FUROSEMIDE

TABLET; ORAL

FUROSEMIDE

| | | | | | | | |
|-----|----|----------|------|------------|--------------|-----|------|
| >A> | AB | OHM LABS | 20MG | N78010 001 | Sep 18, 2006 | Sep | NEWA |
| >A> | AB | | 40MG | N78010 002 | Sep 18, 2006 | Sep | NEWA |
| >A> | AB | | 80MG | N78010 003 | Sep 18, 2006 | Sep | NEWA |

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

| | | | | | | |
|----|--------------------|-------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 100MG | N75350 001 | Sep 12, 2003 | Jun | CAHN |
| AB | | 300MG | N75350 002 | Sep 12, 2003 | Jun | CAHN |
| AB | | 400MG | N75350 003 | Sep 12, 2003 | Jun | CAHN |
| AB | SANDOZ | 100MG | N75428 001 | Jan 24, 2006 | Jan | NEWA |
| AB | | 300MG | N75428 002 | Jan 24, 2006 | Jan | NEWA |
| AB | | 400MG | N75428 003 | Jan 24, 2006 | Jan | NEWA |
| AB | SUN PHARM INDS LTD | 100MG | N77242 001 | Aug 24, 2006 | Aug | NEWA |
| AB | | 300MG | N77242 002 | Aug 24, 2006 | Aug | NEWA |
| AB | | 400MG | N77242 003 | Aug 24, 2006 | Aug | NEWA |

TABLET; ORAL

GABAPENTIN

| | | | | | | | |
|-----|-------------------|-----------------|------------|--------------|--------------|------|------|
| AB | ACTAVIS ELIZABETH | 600MG | N75694 001 | Oct 21, 2004 | Jun | CAHN | |
| AB | | 800MG | N75694 002 | Oct 21, 2004 | Jun | CAHN | |
| >A> | AB | APOTEX INC | 100MG | N77894 001 | Oct 10, 2006 | Sep | NEWA |
| >A> | AB | | 300MG | N77894 002 | Oct 10, 2006 | Sep | NEWA |
| >A> | AB | | 400MG | N77894 003 | Oct 10, 2006 | Sep | NEWA |
| | AB | | 600MG | N77661 004 | Sep 13, 2006 | Aug | NEWA |
| | AB | | 800MG | N77661 005 | Sep 13, 2006 | Aug | NEWA |
| | AB | GLENMARK PHARMS | 600MG | N77662 001 | Aug 18, 2006 | Aug | NEWA |
| | AB | | 800MG | N77662 002 | Aug 18, 2006 | Aug | NEWA |
| >D> | | IVAX PHARMS | 100MG | N76017 001 | Apr 28, 2004 | Sep | CFTG |

TABLET; ORAL

GABAPENTIN

| | | | | | | | |
|-----|----|--------------------|-------|------------|--------------|-----|------|
| >A> | AB | IVAX PHARMS | 100MG | N76017 001 | Apr 28, 2004 | Sep | CFTG |
| >D> | | | 300MG | N76017 002 | Apr 28, 2004 | Sep | CFTG |
| >A> | AB | | 300MG | N76017 002 | Apr 28, 2004 | Sep | CFTG |
| >D> | | | 400MG | N76017 003 | Apr 28, 2004 | Sep | CFTG |
| >A> | AB | | 400MG | N76017 003 | Apr 28, 2004 | Sep | CFTG |
| | AB | SANDOZ | 600MG | N76120 001 | Jan 27, 2006 | Jan | NEWA |
| | AB | | 600MG | N76877 001 | Jul 06, 2006 | Jun | NEWA |
| | AB | | 800MG | N76120 002 | Jan 27, 2006 | Jan | NEWA |
| | AB | | 800MG | N76877 002 | Jul 06, 2006 | Jun | NEWA |
| | AB | SUN PHARM INDS LTD | 600MG | N77525 001 | Aug 24, 2006 | Aug | NEWA |
| | AB | | 800MG | N77525 002 | Aug 24, 2006 | Aug | NEWA |

GADOVERSETAMIDE

INJECTABLE; INJECTION

OPTIMARK

| | | | | | | |
|---|--------------|----------------------------|------------|--------------|-----|------|
| + | MALLINCKRODT | 1654.5MG/5ML (330.9MG/ML) | N20937 001 | Dec 08, 1999 | Jan | CPOT |
| + | | 3309MG/10ML (330.9MG/ML) | N20937 002 | Dec 08, 1999 | Jan | NEWA |
| + | | 4963.5MG/15ML (330.9MG/ML) | N20937 003 | Dec 08, 1999 | Jan | NEWA |
| + | | 6618MG/20ML (330.9MG/ML) | N20937 004 | Dec 08, 1999 | Jan | NEWA |
| + | | 16.545GM/50ML (330.9MG/ML) | N20975 001 | Dec 08, 1999 | Jan | CPOT |

OPTIMARK IN PLASTIC CONTAINER

| | | | | | | |
|---|--------------|----------------------------|------------|--------------|-----|------|
| + | MALLINCKRODT | 1654.5MG/5ML (330.9MG/ML) | N20976 001 | Dec 08, 1999 | Jan | CPOT |
| + | | 3309MG/10ML (330.9MG/ML) | N20976 002 | Dec 08, 1999 | Jan | NEWA |
| + | | 4963.5MG/15ML (330.9MG/ML) | N20976 003 | Dec 08, 1999 | Jan | NEWA |
| + | | 6618MG/20ML (330.9MG/ML) | N20976 004 | Dec 08, 1999 | Jan | NEWA |

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE; ORAL

RAZADYNE ER

| | | | | | | |
|---|---------|--------------|------------|--------------|-----|------|
| + | JANSSEN | EQ 8MG BASE | N21615 001 | Apr 01, 2005 | May | CMS2 |
| | | EQ 16MG BASE | N21615 002 | Apr 01, 2005 | May | CMS2 |
| | | EQ 24MG BASE | N21615 003 | Apr 01, 2005 | May | CMS2 |

GANCICLOVIR

CAPSULE; ORAL

CYTOVENE

| | | | | | | |
|---|------------|-------|------------|--------------|-----|------|
| @ | ROCHE PALO | 250MG | N20460 001 | Dec 22, 1994 | Jun | DISC |
| @ | | 500MG | N20460 002 | Dec 12, 1997 | Jun | DISC |

GANCICLOVIR

| | | | | | | |
|---|---------|-------|------------|--------------|-----|------|
| | RANBAXY | 250MG | N76457 001 | Jun 27, 2003 | Jun | CTEC |
| + | | 500MG | N76457 002 | Jun 27, 2003 | Jun | CRLD |

GATIFLOXACIN

INJECTABLE; INJECTION

TEQUIN

| | | | | | | |
|---|----------------------|---------------------|------------|--------------|-----|------|
| @ | BRISTOL MYERS SQUIBB | 400MG/40ML(10MG/ML) | N21062 004 | Dec 17, 1999 | May | DISC |
|---|----------------------|---------------------|------------|--------------|-----|------|

TEQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | | |
|---|----------------------|---------------------|------------|--------------|-----|------|
| @ | BRISTOL MYERS SQUIBB | 200MG/100ML(2MG/ML) | N21062 001 | Dec 17, 1999 | May | DISC |
| @ | | 400MG/200ML(2MG/ML) | N21062 002 | Dec 17, 1999 | May | DISC |

SUSPENSION; ORAL

TEQUIN

| | | | | | | |
|---|----------------------|-----------|------------|--------------|-----|------|
| @ | BRISTOL MYERS SQUIBB | 200MG/5ML | N21678 001 | Aug 27, 2004 | Jun | DISC |
|---|----------------------|-----------|------------|--------------|-----|------|

TABLET; ORAL

TEQUIN

@ BRISTOL MYERS SQUIBB 200MG

N21061 001 Dec 17, 1999 May DISC

@ 400MG

N21061 002 Dec 17, 1999 May DISC

GEMFIBROZIL

TABLET; ORAL

GEMFIBROZIL

AB INVAGEN PHARMS 600MG

N77836 001 Jul 27, 2006 Jul NEWA

GENTAMICIN SULFATE

SOLUTION/DROPS; OPHTHALMIC

GENTACIDIN

@ NOVARTIS EQ 0.3% BASE

N62480 001 Mar 30, 1984 Jun DISC

GLIMEPIRIDE

TABLET; ORAL

GLIMEPIRIDE

AB COBALT 1MG

N77280 001 Feb 03, 2006 Jan NEWA

AB 2MG

N77280 002 Feb 03, 2006 Jan NEWA

AB 4MG

N77280 003 Feb 03, 2006 Jan NEWA

AB GENPHARM 1MG

N77486 001 Feb 10, 2006 Jan NEWA

AB 2MG

N77486 002 Feb 10, 2006 Jan NEWA

AB 4MG

N77486 003 Feb 10, 2006 Jan NEWA

GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

DUETACT

TAKEDA GLOBAL 2MG;30MG

N21925 001 Jul 28, 2006 Jul NEWA

>D> 4MG;30MG

N21925 002 Jul 28, 2006 Sep CRLD

>A> + 4MG;30MG

N21925 002 Jul 28, 2006 Sep CRLD

4MG;30MG

N21925 002 Jul 28, 2006 Jul NEWA

GLIPIZIDE

TABLET; ORAL

GLIPIZIDE

AB CARACO 5MG

N77820 001 Jul 11, 2006 Jun NEWA

AB 10MG

N77820 002 Jul 11, 2006 Jun NEWA

@ ENDO PHARMS 5MG

N74378 001 Nov 28, 1994 Feb DISC

@ 10MG

N74378 002 Nov 28, 1994 Feb DISC

TABLET, EXTENDED RELEASE; ORAL

GLIPIZIDE

AB WATSON LABS 2.5MG

N76467 003 Mar 27, 2006 Mar NEWA

GLUCOTROL XL

AB PFIZER 2.5MG

N20329 003 Aug 10, 1999 Mar CFTG

GLYBURIDE

TABLET; ORAL

DIABETA

BX + SANOFI AVENTIS US 5MG

N17532 003 May 01, 1984 Feb CRLD

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLYBURIDE AND METFORMIN HYDROCHLORIDE

AB ACTAVIS ELIZABETH 1.25MG;250MG

N76716 001 Jun 28, 2005 Jun CAHN

TABLET; ORAL

GLYBURIDE AND METFORMIN HYDROCHLORIDE

| | | | | | | |
|----|-------------------|-------------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 2.5MG;500MG | N76716 002 | Jun 28, 2005 | Jun | CAHN |
| AB | | 5MG;500MG | N76716 003 | Jun 28, 2005 | Jun | CAHN |

GLYCOPYRROLATE

TABLET; ORAL

GLYCOPYRROLATE

| | | | | | | |
|---------------|---------------------|-----|------------|--------------|-----|------|
| AA | KALI LABS | 1MG | N40653 001 | Aug 31, 2006 | Aug | NEWA |
| AA | | 2MG | N40653 002 | Aug 31, 2006 | Aug | NEWA |
| ROBINUL | | | | | | |
| AA | + SCIELE PHARMA INC | 1MG | N12827 001 | | Jun | CAHN |
| ROBINUL FORTE | | | | | | |
| AA | + SCIELE PHARMA INC | 2MG | N12827 002 | | Jun | CAHN |

GUANFACINE HYDROCHLORIDE

TABLET; ORAL

TENEX

| | | | | | | |
|----|--------------------|-------------|------------|--------------|-----|------|
| AB | DR REDDYS LABS INC | EQ 1MG BASE | N19032 001 | Oct 27, 1986 | May | CAHN |
| AB | + @ | EQ 2MG BASE | N19032 002 | Nov 07, 1988 | May | CAHN |
| | | EQ 3MG BASE | N19032 003 | Nov 07, 1988 | May | CAHN |

HALOBETASOL PROPIONATE

CREAM; TOPICAL

HALOBETASOL PROPIONATE

| | | | | | | |
|----|----------------|-------|------------|--------------|-----|------|
| AB | PERRIGO ISRAEL | 0.05% | N77123 001 | Dec 16, 2004 | Apr | CAHN |
|----|----------------|-------|------------|--------------|-----|------|

OINTMENT; TOPICAL

HALOBETASOL PROPIONATE

| | | | | | | |
|----|----------------------|-------|------------|--------------|-----|------|
| AB | ACTAVIS MID ATLANTIC | 0.05% | N77109 001 | Jun 14, 2005 | Jun | CAHN |
| AB | G AND W LABS | 0.05% | N77721 001 | Sep 07, 2006 | Aug | NEWA |
| AB | PERRIGO | 0.05% | N76872 001 | Dec 16, 2004 | Mar | CAHN |

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALOPERIDOL DECANOATE

| | | | | | | |
|----|--------|------------------|------------|--------------|-----|------|
| AO | SANDOZ | EQ 50MG BASE/ML | N76463 001 | Jun 24, 2005 | Jan | CAHN |
| AO | | EQ 100MG BASE/ML | N76463 002 | Jun 24, 2005 | Jan | CAHN |

HALOPERIDOL LACTATE

INJECTABLE; INJECTION

HALOPERIDOL

| | | | | | | |
|----|--------|----------------|------------|--------------|-----|------|
| AP | SANDOZ | EQ 5MG BASE/ML | N76464 001 | Sep 29, 2004 | Jan | CAHN |
|----|--------|----------------|------------|--------------|-----|------|

SOLUTION; ORAL

HALOPERIDOL LACTATE

| | | | | | | |
|--|----------------------|----------------|------------|--------------|-----|------|
| | ACTAVIS MID ATLANTIC | EQ 1MG BASE/ML | N74536 001 | Nov 02, 1995 | Jul | CAHN |
|--|----------------------|----------------|------------|--------------|-----|------|

HALOTHANE

LIQUID; INHALATION

HALOTHANE

| | | | | | | |
|--|-----------|--------|------------|--|-----|------|
| | @ HOSPIRA | 99.99% | N83254 001 | | Aug | DISC |
|--|-----------|--------|------------|--|-----|------|

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYDROCODONE COMPOUND

| | | | | | | |
|----|----------------------|-------------------|------------|--------------|-----|------|
| AA | ACTAVIS MID ATLANTIC | 1.5MG/5ML;5MG/5ML | N88017 001 | Jul 05, 1983 | Jul | CAHN |
|----|----------------------|-------------------|------------|--------------|-----|------|

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDRALAZINE HYDROCHLORIDE

| | | | | | |
|----------------|---------|------------|--------------|-----|------|
| @ SICOR PHARMS | 20MG/ML | N40373 001 | Feb 23, 2000 | Aug | DISC |
|----------------|---------|------------|--------------|-----|------|

TABLET; ORAL

APRESOLINE

| | | | | | |
|------------|------|------------|--|-----|------|
| @ NOVARTIS | 10MG | N08303 004 | | Feb | DISC |
|------------|------|------------|--|-----|------|

| | | | | | |
|---|------|------------|--|-----|------|
| @ | 25MG | N08303 001 | | Feb | DISC |
|---|------|------------|--|-----|------|

| | | | | | |
|---|------|------------|--|-----|------|
| @ | 50MG | N08303 002 | | Feb | DISC |
|---|------|------------|--|-----|------|

| | | | | | |
|---|-------|------------|--|-----|------|
| @ | 100MG | N08303 005 | | Feb | DISC |
|---|-------|------------|--|-----|------|

HYDRALAZINE HYDROCHLORIDE

| | | | | | | |
|------|-------|------|------------|--------------|-----|------|
| AA + | PLIVA | 10MG | N89097 001 | Dec 18, 1985 | Feb | CRLD |
|------|-------|------|------------|--------------|-----|------|

| | | | | | | |
|------|--|------|------------|--------------|-----|------|
| AA + | | 25MG | N88467 001 | May 01, 1984 | Feb | CRLD |
|------|--|------|------------|--------------|-----|------|

| | | | | | | |
|------|--|------|------------|--------------|-----|------|
| AA + | | 50MG | N88468 001 | May 01, 1984 | Feb | CRLD |
|------|--|------|------------|--------------|-----|------|

| | | | | | | |
|------|--|-------|------------|--------------|-----|------|
| AA + | | 100MG | N89098 001 | Dec 18, 1985 | Feb | CRLD |
|------|--|-------|------------|--------------|-----|------|

| | | | | | |
|-----------------|------|------------|--|-----|------|
| @ RADIUS PHARMS | 25MG | N86243 001 | | Aug | CAHN |
|-----------------|------|------------|--|-----|------|

| | | | | | |
|---|------|------------|--|-----|------|
| @ | 50MG | N86242 002 | | Aug | CAHN |
|---|------|------------|--|-----|------|

| | | | | | |
|------------------|------|------------|--------------|-----|------|
| @ TG UNITED LABS | 10MG | N88846 001 | Feb 26, 1985 | May | CAHN |
|------------------|------|------------|--------------|-----|------|

| | | | | | |
|---|------|------------|--------------|-----|------|
| @ | 25MG | N88847 001 | Feb 26, 1985 | May | CAHN |
|---|------|------------|--------------|-----|------|

| | | | | | |
|---|------|------------|--------------|-----|------|
| @ | 50MG | N88848 001 | Feb 26, 1985 | May | CAHN |
|---|------|------------|--------------|-----|------|

| | | | | | |
|---|-------|------------|--------------|-----|------|
| @ | 100MG | N88849 001 | Feb 26, 1985 | May | CAHN |
|---|-------|------------|--------------|-----|------|

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CAM-AP-ES

| | | | | | |
|------------------|-----------------|------------|--|-----|------|
| @ TG UNITED LABS | 25MG;15MG;0.1MG | N84897 001 | | May | CAHN |
|------------------|-----------------|------------|--|-----|------|

HYDROCHLOROTHIAZIDE

TABLET; ORAL

HYDROCHLOROTHIAZIDE

| | | | | | | |
|----|-------------------|------|------------|--|-----|------|
| AB | ACTAVIS ELIZABETH | 25MG | N85054 002 | | Jun | CAHN |
|----|-------------------|------|------------|--|-----|------|

| | | | | | | |
|----|--|------|------------|--|-----|------|
| AB | | 50MG | N85208 001 | | Jun | CAHN |
|----|--|------|------------|--|-----|------|

| | | | | | |
|------------------|------|------------|--|-----|------|
| @ TG UNITED LABS | 25MG | N85683 001 | | May | CAHN |
|------------------|------|------------|--|-----|------|

| | | | | | |
|---|------|------------|--|-----|------|
| @ | 50MG | N83965 001 | | May | CAHN |
|---|------|------------|--|-----|------|

| | | | | | | |
|----|-----------|------|------------|--------------|-----|------|
| AB | WEST WARD | 25MG | N84878 002 | Jul 12, 2006 | Jun | NEWA |
|----|-----------|------|------------|--------------|-----|------|

HYDROCHLOROTHIAZIDE; LISINAPRIL

TABLET; ORAL

LISINAPRIL AND HYDROCHLOROTHIAZIDE

| | | | | | | |
|----|-------------------|-------------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 12.5MG;10MG | N76230 001 | Jul 01, 2002 | Jun | CAHN |
|----|-------------------|-------------|------------|--------------|-----|------|

| | | | | | | |
|----|--|-------------|------------|--------------|-----|------|
| AB | | 12.5MG;20MG | N76230 002 | Jul 01, 2002 | Jun | CAHN |
|----|--|-------------|------------|--------------|-----|------|

| | | | | | | |
|----|--|-----------|------------|--------------|-----|------|
| AB | | 25MG;20MG | N76230 003 | Jul 01, 2002 | Jun | CAHN |
|----|--|-----------|------------|--------------|-----|------|

| | | | | | | |
|----|-----------|-------------|------------|--------------|-----|------|
| AB | AUROBINDO | 12.5MG;10MG | N77606 001 | Mar 14, 2006 | Feb | NEWA |
|----|-----------|-------------|------------|--------------|-----|------|

| | | | | | | |
|----|--|-------------|------------|--------------|-----|------|
| AB | | 12.5MG;20MG | N77606 002 | Mar 14, 2006 | Feb | NEWA |
|----|--|-------------|------------|--------------|-----|------|

| | | | | | | |
|----|--|-----------|------------|--------------|-----|------|
| AB | | 25MG;20MG | N77606 003 | Mar 14, 2006 | Feb | NEWA |
|----|--|-----------|------------|--------------|-----|------|

| | | | | | | | |
|-----|----|-------|-------------|------------|--------------|-----|------|
| >A> | AB | LUPIN | 12.5MG;10MG | N77912 001 | Sep 27, 2006 | Sep | NEWA |
|-----|----|-------|-------------|------------|--------------|-----|------|

| | | | | | | | |
|-----|----|--|-------------|------------|--------------|-----|------|
| >A> | AB | | 12.5MG;20MG | N77912 002 | Sep 27, 2006 | Sep | NEWA |
|-----|----|--|-------------|------------|--------------|-----|------|

| | | | | | | | |
|-----|----|--|-----------|------------|--------------|-----|------|
| >A> | AB | | 25MG;20MG | N77912 003 | Sep 27, 2006 | Sep | NEWA |
|-----|----|--|-----------|------------|--------------|-----|------|

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

ALDORIL 15

| | | | | | |
|---------|-----------|------------|--|-----|------|
| @ MERCK | 15MG;25MG | N13402 001 | | Jun | DISC |
|---------|-----------|------------|--|-----|------|

TABLET; ORAL

| | | | | | | | | | |
|------------------------------------|------------|--|--------|-----|--------------|-----|------|------|--|
| ALDORIL 25 | | | | | | | | | |
| @ MERCK | 25MG;250MG | | N13402 | 002 | | | Jun | DISC | |
| ALDORIL D30 | | | | | | | | | |
| @ MERCK | 30MG;500MG | | N13402 | 003 | | | Jun | DISC | |
| ALDORIL D50 | | | | | | | | | |
| @ MERCK | 50MG;500MG | | N13402 | 004 | | | Jun | DISC | |
| METHYLDOPA AND HYDROCHLOROTHIAZIDE | | | | | | | | | |
| MYLAN | 15MG;250MG | | N70264 | 001 | Jan 23, 1986 | Jun | CTEC | | |
| + | 25MG;250MG | | N70265 | 001 | Jan 23, 1986 | Jun | CRLD | | |
| @ PAR PHARM | 15MG;250MG | | N70616 | 001 | Feb 02, 1987 | Jun | DISC | | |
| @ | 25MG;250MG | | N70612 | 001 | Feb 02, 1987 | Jun | DISC | | |
| @ | 30MG;500MG | | N70613 | 001 | Feb 02, 1987 | Jun | DISC | | |
| @ | 50MG;500MG | | N70614 | 001 | Feb 02, 1987 | Jun | DISC | | |
| @ SANDOZ | 15MG;250MG | | N70182 | 001 | Jan 15, 1986 | Jun | DISC | | |
| @ | 25MG;250MG | | N70183 | 001 | Jan 15, 1986 | Jun | DISC | | |
| @ | 30MG;500MG | | N70543 | 001 | Jan 15, 1986 | Jun | DISC | | |
| @ | 50MG;500MG | | N70544 | 001 | Jan 15, 1986 | Jun | DISC | | |
| @ WATSON LABS | 15MG;250MG | | N70958 | 001 | Feb 06, 1989 | Jun | DISC | | |
| @ | 25MG;250MG | | N70959 | 001 | Jan 19, 1989 | Jun | DISC | | |
| @ | 30MG;500MG | | N71069 | 001 | Jan 19, 1989 | Jun | DISC | | |
| @ | 50MG;500MG | | N70960 | 001 | Feb 06, 1989 | Jun | DISC | | |

HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

| | | | | | | | | | |
|-------------|--------------------------|--|--------|-----|--------------|-----|------|--|--|
| DUTOPROL | | | | | | | | | |
| ASTRAZENECA | 12.5MG;EQ 25MG TARTRATE | | N21956 | 001 | Aug 28, 2006 | Aug | NEWA | | |
| | 12.5MG;EQ 50MG TARTRATE | | N21956 | 002 | Aug 28, 2006 | Aug | NEWA | | |
| + | 12.5MG;EQ 100MG TARTRATE | | N21956 | 003 | Aug 28, 2006 | Aug | NEWA | | |

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

| | | | | | | | | | |
|---|-----------|--|--------|-----|--------------|-----|------|--|--|
| PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE | | | | | | | | | |
| AB ACTAVIS ELIZABETH | 25MG;40MG | | N70851 | 001 | May 15, 1986 | Jun | CAHN | | |
| AB | 25MG;80MG | | N70852 | 001 | May 15, 1986 | Jun | CAHN | | |

HYDROCHLOROTHIAZIDE; TIMOLOL MALEATE

TABLET; ORAL

| | | | | | | | | | |
|----------------|-----------|--|--------|-----|--|-----|------|--|--|
| TIMOLIDE 10-25 | | | | | | | | | |
| @ MERCK | 25MG;10MG | | N18061 | 001 | | Jun | DISC | | |

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

| | | | | | | | | | |
|------------|--------------|--|--------|-----|--------------|-----|------|--|--|
| DIOVAN HCT | | | | | | | | | |
| NOVARTIS | 12.5MG;320MG | | N20818 | 004 | Apr 28, 2006 | Apr | NEWA | | |
| | 25MG;160MG | | N20818 | 003 | Jan 17, 2002 | Apr | CRLD | | |
| + | 25MG;320MG | | N20818 | 005 | Apr 28, 2006 | Apr | NEWA | | |

HYDROCORTISONE

CREAM; TOPICAL

| | | | | | | | | | |
|-------------------------|------|--|--------|-----|--------------|-----|------|--|--|
| HYDROCORTISONE | | | | | | | | | |
| AT ACTAVIS MID ATLANTIC | 1% | | N87795 | 001 | May 03, 1983 | Jun | CAHN | | |
| AT | 2.5% | | N89682 | 001 | Mar 10, 1988 | Jun | CAHN | | |

| | | | | | | | | |
|----|--|-------------------------------|--------|-----|--------------|-----|------|--|
| | OINTMENT; TOPICAL | | | | | | | |
| | HYDROCORTISONE | | | | | | | |
| AT | ACTAVIS MID ATLANTIC | 1% | N87796 | 001 | Oct 13, 1982 | Jun | CAHN | |
| | POWDER; FOR RX COMPOUNDING | | | | | | | |
| | HYDRO-RX | | | | | | | |
| | + X GEN PHARMS | 100% | N85982 | 001 | | Jun | CTNA | |
| | TABLET; ORAL | | | | | | | |
| | CORTEF | | | | | | | |
| | @ PHARMACIA AND UPJOHN | 10MG | N08697 | 001 | | Jun | CTEC | |
| | HYDROCORTONE | | | | | | | |
| | @ MERCK | 10MG | N08506 | 007 | | Jun | DISC | |
| | @ | 20MG | N08506 | 011 | | Jun | DISC | |
| | <u>HYDROCORTISONE SODIUM SUCCINATE</u> | | | | | | | |
| | INJECTABLE; INJECTION | | | | | | | |
| | A-HYDROCORT | | | | | | | |
| AP | HOSPIRA | EQ 100MG BASE/VIAL | N40666 | 001 | Apr 06, 2006 | Mar | NEWA | |
| | <u>HYDROCORTISONE; NEOMYCIN; POLYMYXIN B SULFATE</u> | | | | | | | |
| | SUSPENSION/DROPS; OTIC | | | | | | | |
| | NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE | | | | | | | |
| AT | PHARMAFORCE | 1%;EQ 3.5MG BASE;10,000 UNITS | N65219 | 001 | May 01, 2006 | Apr | NEWA | |
| | <u>HYDROXYZINE HYDROCHLORIDE</u> | | | | | | | |
| | SYRUP; ORAL | | | | | | | |
| | ATARAX | | | | | | | |
| | @ ROERIG | 10MG/5ML | N10485 | 001 | | Jun | DISC | |
| | HYDROXYZINE HYDROCHLORIDE | | | | | | | |
| AA | + ACTAVIS MID ATLANTIC | 10MG/5ML | N86880 | 001 | | Jul | CAHN | |
| AA | + ALPHARMA US PHARMS | 10MG/5ML | N86880 | 001 | | Jun | CRLD | |
| AA | + HI TECH PHARMA | 10MG/5ML | N40010 | 001 | Oct 28, 1994 | Jun | CRLD | |
| AA | + MORTON GROVE | 10MG/5ML | N87294 | 001 | Apr 12, 1982 | Jun | CRLD | |
| AA | + VINTAGE PHARMS | 10MG/5ML | N40391 | 001 | Apr 10, 2002 | Jun | CRLD | |
| | <u>HYDROXYZINE PAMOATE</u> | | | | | | | |
| | CAPSULE; ORAL | | | | | | | |
| | HYDROXYZINE PAMOATE | | | | | | | |
| | BARR | EQ 100MG HCL | N88488 | 001 | Jun 15, 1984 | Mar | CTEC | |
| | VISTARIL | | | | | | | |
| | @ PFIZER | EQ 100MG HCL | N11459 | 006 | | Mar | DISC | |
| | <u>IBANDRONATE SODIUM</u> | | | | | | | |
| | INJECTABLE; INTRAVENOUS | | | | | | | |
| | BONIVA | | | | | | | |
| | + ROCHE | EQ 3MG BASE/3ML | N21858 | 001 | Jan 06, 2006 | Jan | NEWA | |
| | <u>IBUPROFEN</u> | | | | | | | |
| | SUSPENSION; ORAL | | | | | | | |
| | IBUPROFEN | | | | | | | |
| AB | ACTAVIS MID ATLANTIC | 100MG/5ML | N74978 | 001 | Mar 25, 1998 | Jul | CAHN | |
| | TABLET; ORAL | | | | | | | |
| | IBU | | | | | | | |
| | @ BASF | 400MG | N18197 | 001 | | Jun | DISC | |
| | @ | 400MG | N70083 | 001 | Feb 22, 1985 | Jun | DISC | |
| | @ | 600MG | N70099 | 001 | Mar 29, 1985 | Jun | DISC | |

TABLET; ORAL

IBU

| | | | | | |
|--------|-------|------------|--------------|-----|------|
| @ BASF | 800MG | N70745 001 | Jul 23, 1986 | Jun | DISC |
|--------|-------|------------|--------------|-----|------|

IBUPROFEN LYSINE

INJECTABLE; INTRAVENOUS

NEOPROFEN

| | | | | | |
|---------------|------------------------------------|------------|--------------|-----|------|
| + FARMACON IL | EQ 20MG BASE/2ML (EQ 10MG BASE/ML) | N21903 001 | Apr 13, 2006 | Apr | NEWA |
|---------------|------------------------------------|------------|--------------|-----|------|

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

| | | | | | | |
|----|-------------------|--------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 1.25MG | N74722 001 | Jun 17, 1996 | Jun | CAHN |
| AB | | 2.5MG | N74722 002 | Jun 17, 1996 | Jun | CAHN |

INDIUM IN-111 OXYQUINOLINE

INJECTABLE; INJECTION

INDIUM IN-111 OXYQUINOLINE

| | | | | | |
|-----------------|---------|------------|--------------|-----|------|
| + GE HEALTHCARE | 1mCi/ML | N19044 001 | Dec 24, 1985 | Aug | CRLD |
|-----------------|---------|------------|--------------|-----|------|

INDIUM IN-111 PENTETATE DISODIUM

INJECTABLE; INTRATHECAL

MPI INDIUM DTPA IN 111

| | | | | | |
|-----------------|---------|------------|--------------|-----|------|
| + GE HEALTHCARE | 1mCi/ML | N17707 001 | Feb 18, 1982 | Aug | CRLD |
|-----------------|---------|------------|--------------|-----|------|

INDIUM IN-111 PENTETREOTIDE KIT

INJECTABLE; INJECTION

OCTREOSCAN

| | | | | | |
|----------------|---------|------------|--------------|-----|------|
| + MALLINCKRODT | 3mCi/ML | N20314 001 | Jun 02, 1994 | Aug | CRLD |
|----------------|---------|------------|--------------|-----|------|

INDOMETHACIN

CAPSULE; ORAL

INDOCIN

| | | | | | |
|---------|------|------------|--|-----|------|
| @ MERCK | 25MG | N16059 001 | | Jun | DISC |
|---------|------|------------|--|-----|------|

| | | | | | |
|---|------|------------|--|-----|------|
| @ | 50MG | N16059 002 | | Jun | DISC |
|---|------|------------|--|-----|------|

INDO-LEMMON

| | | | | | |
|--------|------|------------|--------------|-----|------|
| @ TEVA | 25MG | N70266 001 | Nov 07, 1985 | Jun | DISC |
|--------|------|------------|--------------|-----|------|

| | | | | | |
|---|------|------------|--------------|-----|------|
| @ | 50MG | N70267 001 | Nov 07, 1985 | Jun | DISC |
|---|------|------------|--------------|-----|------|

INDOMETHACIN

| | | | | | |
|--------------------|------|------------|--------------|-----|------|
| @ CLONMEL HLTHCARE | 25MG | N18851 001 | May 18, 1984 | Jun | DISC |
|--------------------|------|------------|--------------|-----|------|

| | | | | | |
|---|------|------------|--------------|-----|------|
| @ | 50MG | N18851 002 | May 18, 1984 | Jun | DISC |
|---|------|------------|--------------|-----|------|

| | | | | | |
|---------------|------|------------|--------------|-----|------|
| @ IVAX PHARMS | 25MG | N70719 001 | Feb 12, 1986 | Jun | DISC |
|---------------|------|------------|--------------|-----|------|

| | | | | | |
|---|------|------------|--------------|-----|------|
| @ | 50MG | N70756 001 | Feb 12, 1986 | Jun | DISC |
|---|------|------------|--------------|-----|------|

| | | | | | |
|----------------|------|------------|--------------|-----|------|
| @ MUTUAL PHARM | 25MG | N70899 001 | Feb 09, 1987 | Jun | DISC |
|----------------|------|------------|--------------|-----|------|

| | | | | | |
|---|------|------------|--------------|-----|------|
| @ | 50MG | N70900 001 | Feb 09, 1987 | Jun | DISC |
|---|------|------------|--------------|-----|------|

| | | | | | |
|-------|------|------------|--------------|-----|------|
| MYLAN | 25MG | N18858 001 | Apr 20, 1984 | Jun | CTEC |
|-------|------|------------|--------------|-----|------|

| | | | | | | |
|----|---|------|------------|--------------|-----|------|
| AB | + | 50MG | N18858 002 | Apr 20, 1984 | Jun | CRLD |
|----|---|------|------------|--------------|-----|------|

| | | | | | |
|-------------|------|------------|--------------|-----|------|
| @ PAR PHARM | 25MG | N18829 002 | Aug 06, 1984 | Jun | DISC |
|-------------|------|------------|--------------|-----|------|

| | | | | | |
|---|------|------------|--------------|-----|------|
| @ | 50MG | N18829 001 | Aug 06, 1984 | Jun | DISC |
|---|------|------------|--------------|-----|------|

| | | | | | |
|---|------|------------|--------------|-----|------|
| @ | 50MG | N70651 001 | Mar 05, 1986 | Jun | DISC |
|---|------|------------|--------------|-----|------|

| | | | | | |
|---------|------|------------|--------------|-----|------|
| @ PLIVA | 25MG | N71148 001 | Mar 18, 1987 | Jun | DISC |
|---------|------|------------|--------------|-----|------|

| | | | | | |
|---|------|------------|--------------|-----|------|
| @ | 50MG | N71149 001 | Mar 18, 1987 | Jun | DISC |
|---|------|------------|--------------|-----|------|

| | | | | | |
|-----------------|------|------------|--------------|-----|------|
| @ RADIUS PHARMS | 25MG | N18851 001 | May 18, 1984 | Aug | CAHN |
|-----------------|------|------------|--------------|-----|------|

| | | | | | |
|---|------|------------|--------------|-----|------|
| @ | 50MG | N18851 002 | May 18, 1984 | Aug | CAHN |
|---|------|------------|--------------|-----|------|

| | | | | | |
|----------|------|------------|--------------|-----|------|
| @ SANDOZ | 25MG | N70673 001 | Apr 29, 1987 | Jun | DISC |
|----------|------|------------|--------------|-----|------|

CAPSULE; ORAL

INDOMETHACIN

| | | | | | |
|----------|------|------------|--------------|-----|------|
| @ SANDOZ | 50MG | N70674 001 | Apr 29, 1987 | Jun | DISC |
| @ TEVA | 25MG | N71342 001 | Apr 18, 1988 | Jun | DISC |
| @ | 50MG | N71343 001 | Apr 18, 1988 | Jun | DISC |

CAPSULE, EXTENDED RELEASE; ORAL

INDOCIN SR

| | | | | | |
|---------------|------|------------|--------------|-----|------|
| + SANDOZ | 75MG | N74464 001 | May 28, 1998 | Jun | CTEC |
| @ INWOOD LABS | 75MG | N72410 001 | Mar 15, 1989 | Jun | DISC |

SUPPOSITORY; RECTAL

INDOMETHACIN

| | | | | | |
|----------------|------|------------|--------------|-----|------|
| + G AND W LABS | 50MG | N73314 001 | Aug 31, 1992 | Apr | CTNA |
|----------------|------|------------|--------------|-----|------|

INSULIN GLULISINE RECOMBINANT

INJECTABLE; SUBCUTANEOUS

APIDRA

| | | | | | |
|---------------------|--------------------------------|------------|--------------|-----|------|
| + SANOFI AVENTIS US | 1000 UNITS/10ML (100 UNITS/ML) | N21629 001 | Apr 16, 2004 | Mar | CMFD |
| + | 300 UNITS/3ML (100 UNITS/ML) | N21629 002 | Dec 20, 2005 | Mar | CMFD |

INSULIN PURIFIED PORK

INJECTABLE; INJECTION

ILETIN II

| | | | | | |
|---------|--------------|------------|--|-----|------|
| @ LILLY | 500 UNITS/ML | N18344 002 | | Jun | DISC |
|---------|--------------|------------|--|-----|------|

INSULIN RECOMBINANT HUMAN

POWDER; INHALATION

EXUBERA

| | | | | | |
|--------|---------|------------|--------------|-----|------|
| PFIZER | 1MG/INH | N21868 001 | Jan 27, 2006 | Jan | NEWA |
| + | 3MG/INH | N21868 002 | Jan 27, 2006 | Jan | NEWA |

IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION

ATROVENT

| | | | | | |
|------------------------|-------------|------------|--------------|-----|------|
| @ BOEHRINGER INGELHEIM | 0.018MG/INH | N19085 001 | Dec 29, 1986 | May | DISC |
|------------------------|-------------|------------|--------------|-----|------|

SOLUTION; INHALATION

ATROVENT

| | | | | | |
|------------------------|-------|------------|--------------|-----|------|
| @ BOEHRINGER INGELHEIM | 0.02% | N20228 001 | Sep 29, 1993 | May | DISC |
|------------------------|-------|------------|--------------|-----|------|

IPRATROPIUM BROMIDE

| | | | | | | |
|----|----------------------|-------|------------|--------------|-----|------|
| AN | ACTAVIS MID ATLANTIC | 0.02% | N75111 001 | Apr 22, 1999 | Jul | CAHN |
| AN | + DEY | 0.02% | N74755 001 | Jan 10, 1997 | May | CRLD |
| | @ ROXANE | 0.02% | N75867 001 | Jul 22, 2002 | May | DISC |

ISONIAZID

INJECTABLE; INJECTION

ISONIAZID

| | | | | | | |
|----|--------|----------|------------|--------------|-----|------|
| AP | SANDOZ | 100MG/ML | N40648 001 | Jul 05, 2005 | Jan | CAHN |
|----|--------|----------|------------|--------------|-----|------|

ISOSORBIDE MONONITRATE

TABLET; ORAL

ISMO

| | | | | | | |
|----|------------------------|------|------------|--------------|-----|------|
| AB | DR REDDYS LABS INC | 20MG | N19091 001 | Dec 30, 1991 | May | CAHN |
| | ISOSORBIDE MONONITRATE | | | | | |
| AB | ACTAVIS ELIZABETH | 10MG | N75037 002 | Oct 30, 1998 | Jun | CAHN |
| AB | | 20MG | N75037 001 | Oct 30, 1998 | Jun | CAHN |

TABLET, EXTENDED RELEASE; ORAL

ISOSORBIDE MONONITRATE

| | | | | | | |
|----|-------------------|------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 30MG | N75306 001 | Dec 31, 1998 | Jun | CAHN |
| AB | | 60MG | N75306 002 | Dec 31, 1998 | Jun | CAHN |
| AB | WEST WARD | 30MG | N76813 002 | Mar 30, 2006 | Mar | NEWA |

ISOTRETINOIN

CAPSULE; ORAL

CLARAVIS

| | | | | | | |
|----|---------|------|------------|--------------|-----|------|
| AB | BARR | 30MG | N76135 003 | May 11, 2006 | May | NEWA |
| AB | SOTRET | | | | | |
| AB | RANBAXY | 30MG | N76503 001 | Jun 20, 2003 | May | CTEC |

ISRADIPINE

CAPSULE; ORAL

DYNACIRC

@ RELIANT PHARMS

2.5MG

N19546 001 Dec 20, 1990 May DISC

ISRADIPINE

| | | | | | | |
|----|-----------------|-------|------------|--------------|-----|------|
| AB | + ABRIKA PHARMS | 5MG | N77317 002 | Jan 05, 2006 | Jun | CRLD |
| AB | | 5MG | N77317 002 | Jan 05, 2006 | Apr | CTEC |
| AB | AMIDE PHARM | 2.5MG | N77169 001 | Apr 24, 2006 | Apr | NEWA |
| AB | | 5MG | N77169 002 | Apr 24, 2006 | Apr | NEWA |

IVERMECTIN

TABLET; ORAL

STROMEKTOL

+ MERCK

3MG

N50742 002 Oct 08, 1998 Jun CRLD

@

6MG

N50742 001 Nov 22, 1996 Jun DISC

KETOCONAZOLE

GEL; TOPICAL

XOLEGEL

+ BARRIER THERAP

2%

N21946 001 Jul 28, 2006 Jul NEWA

KETOPROFEN

CAPSULE; ORAL

KETOPROFEN

| | | | | | | |
|----|---------------|------|------------|--------------|-----|------|
| AB | RADIUS PHARMS | 25MG | N74014 001 | Jan 29, 1993 | Jul | CAHN |
| AB | | 50MG | N74014 002 | Jan 29, 1993 | Jul | CAHN |
| AB | | 75MG | N74014 003 | Jan 29, 1993 | Jul | CAHN |

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

| | | | | | | |
|----|--------|---------|------------|--------------|-----|------|
| AP | SANDOZ | 15MG/ML | N76271 001 | Oct 06, 2004 | Jan | CAHN |
| AP | | 30MG/ML | N76271 002 | Oct 06, 2004 | Jan | CAHN |

KETOTIFEN FUMARATE

SOLUTION/DROPS; OPHTHALMIC

KETOTIFEN FUMARATE

| | | | | | | |
|----|------------|----------------|------------|--------------|-----|------|
| AT | APOTEX | EQ 0.025% BASE | N77354 001 | May 09, 2006 | Apr | NEWA |
| AT | ZADITOR | | | | | |
| AT | + NOVARTIS | EQ 0.025% BASE | N21066 001 | Jul 02, 1999 | Apr | CFTG |

LACTULOSE

SOLUTION; ORAL

CONSTULOSE

AA + ACTAVIS MID ATLANTIC 10GM/15ML N70288 001 Aug 15, 1988 Jul CAHN

SOLUTION; ORAL, RECTAL

ENULOSE

AA + ACTAVIS MID ATLANTIC 10GM/15ML N71548 001 Aug 15, 1988 Jul CAHN

LAMOTRIGINE

TABLET; ORAL

LAMICTAL

AB + GLAXOSMITHKLINE 25MG N20241 005 Dec 27, 1994 Aug CFTG

AB 100MG N20241 001 Dec 27, 1994 Aug CFTG

AB 150MG N20241 002 Dec 27, 1994 Aug CFTG

AB 200MG N20241 003 Dec 27, 1994 Aug CFTG

LAMOTRIGINE

AB TEVA 25MG N76388 001 Aug 30, 2006 Aug NEWA

AB 100MG N76388 002 Aug 30, 2006 Aug NEWA

AB 150MG N76388 003 Aug 30, 2006 Aug NEWA

AB 200MG N76388 004 Aug 30, 2006 Aug NEWA

TABLET, CHEWABLE; ORAL

LAMICTAL CD

AB GLAXOSMITHKLINE 5MG N20764 001 Aug 24, 1998 Jun CFTG

AB + 25MG N20764 002 Aug 24, 1998 Jun CFTG

LAMOTRIGINE

AB TEVA 5MG N76420 001 Jun 21, 2006 Jun NEWA

AB 25MG N76420 002 Jun 21, 2006 Jun NEWA

LANSOPRAZOLE; NAPROXEN

CAPSULE, DELAYED REL PELLETS, TABLET; ORAL

PREVACID NAPRAPAC 250 (COPACKAGED)

TAP PHARM 15MG,N/A;N/A,250MG N21507 002 Nov 14, 2003 Feb CTNA

PREVACID NAPRAPAC 375 (COPACKAGED)

TAP PHARM 15MG,N/A;N/A,375MG N21507 003 Nov 14, 2003 Feb CTNA

PREVACID NAPRAPAC 500 (COPACKAGED)

+ TAP PHARM 15MG,N/A;N/A,500MG N21507 004 Nov 14, 2003 Feb CTNA

LENALIDOMIDE

CAPSULE; ORAL

REVLIMID

CELGENE 10MG N21880 002 Dec 27, 2005 Jul CRLD

15MG N21880 003 Jun 29, 2006 Jul NEWA

+ 25MG N21880 004 Jun 29, 2006 Jul NEWA

LEVETIRACETAM

INJECTABLE; IV (INFUSION)

KEPPRA

+ UCB INC 500MG/5ML (100MG/ML) N21872 001 Jul 31, 2006 Jul NEWA

TABLET; ORAL

KEPPRA

UCB INC 750MG N21035 003 Nov 30, 1999 Mar CRLD

+ 1GM N21035 004 Jan 06, 2006 Mar NEWA

LEVOBETAXOLOL HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

BETAXON

@ ALCON

EQ 0.5% BASE

N21114 001 Feb 23, 2000 Feb CAHN

LEVOCABASTINE HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

LIVOSTIN

@ NOVARTIS

EQ 0.05% BASE

N20219 001 Nov 10, 1993 May DISC

LEVONORGESTREL

TABLET; ORAL

PLAN B

+ DURAMED

0.75MG

N21045 002 Aug 24, 2006 Aug NEWA

LEVOTHYROXINE SODIUM**

**Refer to Preface Section 1.7 Levothyroxine Sodium for amplifying information

TABLET; ORAL

LEVOTHYROXINE SODIUM

| | | | | | | | |
|-----|------|----------|---------|------------|--------------|-----|------|
| >D> | AB2 | GENPHARM | 0.025MG | N76752 001 | Jun 16, 2005 | Sep | CTEC |
| >A> | AB2, | | 0.025MG | N76752 001 | Jun 16, 2005 | Sep | CTEC |
| | AB3 | | | | | | |
| >D> | AB2 | | 0.05MG | N76752 002 | Jun 16, 2005 | Sep | CTEC |
| >A> | AB2, | | 0.05MG | N76752 002 | Jun 16, 2005 | Sep | CTEC |
| | AB3 | | | | | | |
| >D> | AB2 | | 0.075MG | N76752 003 | Jun 16, 2005 | Sep | CTEC |
| >A> | AB2, | | 0.075MG | N76752 003 | Jun 16, 2005 | Sep | CTEC |
| | AB3 | | | | | | |
| >D> | AB2 | | 0.088MG | N76752 004 | Jun 16, 2005 | Sep | CTEC |
| >A> | AB2, | | 0.088MG | N76752 004 | Jun 16, 2005 | Sep | CTEC |
| | AB3 | | | | | | |
| >D> | AB2 | | 0.1MG | N76752 005 | Jun 16, 2005 | Sep | CTEC |
| >A> | AB2, | | 0.1MG | N76752 005 | Jun 16, 2005 | Sep | CTEC |
| | AB3 | | | | | | |
| >D> | AB2 | | 0.112MG | N76752 006 | Jun 16, 2005 | Sep | CTEC |
| >A> | AB2, | | 0.112MG | N76752 006 | Jun 16, 2005 | Sep | CTEC |
| | AB3 | | | | | | |
| >D> | AB2 | | 0.125MG | N76752 007 | Jun 16, 2005 | Sep | CTEC |
| >A> | AB2, | | 0.125MG | N76752 007 | Jun 16, 2005 | Sep | CTEC |
| | AB3 | | | | | | |
| >D> | AB2 | | 0.15MG | N76752 008 | Jun 16, 2005 | Sep | CTEC |
| >A> | AB2, | | 0.15MG | N76752 008 | Jun 16, 2005 | Sep | CTEC |
| | AB3 | | | | | | |
| >D> | AB2 | | 0.175MG | N76752 009 | Jun 16, 2005 | Sep | CTEC |
| >A> | AB2, | | 0.175MG | N76752 009 | Jun 16, 2005 | Sep | CTEC |
| | AB3 | | | | | | |
| >D> | AB2 | | 0.2MG | N76752 010 | Jun 16, 2005 | Sep | CTEC |
| >A> | AB2, | | 0.2MG | N76752 010 | Jun 16, 2005 | Sep | CTEC |
| | AB3 | | | | | | |
| >D> | AB2 | | 0.3MG | N76752 011 | Jun 16, 2005 | Sep | CTEC |
| >A> | AB2, | | 0.3MG | N76752 011 | Jun 16, 2005 | Sep | CTEC |
| | AB3 | | | | | | |

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

XYLOCAINE

AP + ABRAXIS BIOSCIENCE

0.5%

N06488 008

Jul CAHN

AP +

1%

N06488 007

Jul CAHN

AP +

1.5%

N06488 010

Jul CAHN

INJECTABLE; INJECTION

| | | | | | | | | |
|----|---|--------------------------------|-----|--|--------|-----|--------------|----------|
| | | XYLOCAINE | | | | | | |
| | | @ ABRAXIS BIOSCIENCE | 2% | | N06488 | 002 | | Jul CAHN |
| | | XYLOCAINE 4% PRESERVATIVE FREE | | | | | | |
| AP | + | ABRAXIS BIOSCIENCE | 4% | | N10417 | 001 | | Jul CAHN |
| | | XYLOCAINE PRESERVATIVE FREE | | | | | | |
| AP | | ABRAXIS BIOSCIENCE | 1% | | N16801 | 005 | Jan 19, 1988 | Jul CAHN |
| AP | + | | 2% | | N16801 | 001 | | Jul CAHN |
| AP | | | 4% | | N16801 | 002 | | Jul CAHN |
| AP | + | | 10% | | N16801 | 003 | | Jul CAHN |
| AP | + | | 20% | | N16801 | 004 | | Jul CAHN |

INJECTABLE; SPINAL

| | | | | | | | | |
|--|--|---------------------------------|------|--|--------|-----|--|----------|
| | | XYLOCAINE 1.5% W/ DEXTROSE 7.5% | | | | | | |
| | | @ ABRAXIS BIOSCIENCE | 1.5% | | N16297 | 001 | | Jul CAHN |

JELLY; TOPICAL

| | | | | | | | | |
|----|---|--------------------|----|--|--------|-----|--|----------|
| | | ANESTACON | | | | | | |
| AT | + | POLYMEDICA | 2% | | N80429 | 001 | | Jan CDFR |
| | | XYLOCAINE | | | | | | |
| AT | + | ABRAXIS BIOSCIENCE | 2% | | N08816 | 001 | | Jul CAHN |

SOLUTION; ORAL

| | | | | | | | | |
|----|---|---------------------------------|----|--|--------|-----|--|----------|
| | | LIDOCAINE HYDROCHLORIDE VISCOUS | | | | | | |
| AT | | ACTAVIS MID ATLANTIC | 2% | | N86578 | 001 | | Jul CAHN |
| | | XYLOCAINE VISCOUS | | | | | | |
| AT | + | ABRAXIS BIOSCIENCE | 2% | | N09470 | 001 | | Jul CAHN |

SOLUTION; TOPICAL

| | | | | | | | | |
|----|---|--------------------------------|----|--|--------|-----|--|----------|
| | | XYLOCAINE 4% PRESERVATIVE FREE | | | | | | |
| AT | + | ABRAXIS BIOSCIENCE | 4% | | N10417 | 002 | | Jul CAHN |

LIDOCAINE; PRILOCAINE

CREAM; TOPICAL

| | | | | | | | | |
|----|---|--------------------|-----------|--|--------|-----|--------------|----------|
| | | EMLA | | | | | | |
| AB | + | ABRAXIS BIOSCIENCE | 2.5%;2.5% | | N19941 | 001 | Dec 30, 1992 | Jul CAHN |

LIDOCAINE; TETRACAINE

CREAM; TOPICAL

| | | | | | | | | |
|--|---|--------------------------|-------|--|--------|-----|--------------|----------|
| | | LIDOCAINE AND TETRACAINE | | | | | | |
| | + | ZARS | 7%;7% | | N21717 | 001 | Jun 29, 2006 | Jun NEWA |

PATCH; TOPICAL

| | | | | | | | | |
|--|---|-------------|-----------|--|--------|-----|--------------|----------|
| | | SYNERA | | | | | | |
| | + | ENDO PHARMS | 70MG;70MG | | N21623 | 001 | Jun 23, 2005 | Feb CAHN |

LINDANE

LOTION; TOPICAL

| | | | | | | | | |
|----|---|----------------------|----|--|--------|-----|--|----------|
| | | LINDANE | | | | | | |
| AT | + | ACTAVIS MID ATLANTIC | 1% | | N87313 | 001 | | Jul CAHN |

SHAMPOO; TOPICAL

| | | | | | | | | |
|----|---|----------------------|----|--|--------|-----|--|----------|
| | | LINDANE | | | | | | |
| AT | + | ACTAVIS MID ATLANTIC | 1% | | N87266 | 001 | | Jul CAHN |

LIOTHYRONINE SODIUM

INJECTABLE; INJECTION

LIOTHYRONINE SODIUM

| | | | | | | | | |
|-----|----|--------------|-------------------|--|--------|-----|--------------|----------|
| >D> | AP | PHARMAFORCE | EQ 0.01MG BASE/ML | | N76923 | 001 | Aug 17, 2005 | Sep CAHN |
| >A> | AP | X GEN PHARMS | EQ 0.01MG BASE/ML | | N76923 | 001 | Aug 17, 2005 | Sep CAHN |

LISINOPRIL

TABLET; ORAL

LISINOPRIL

| | | | | | | |
|----|-------------------|-------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 2.5MG | N76180 001 | Jul 01, 2002 | Jun | CAHN |
| AB | | 5MG | N76180 002 | Jul 01, 2002 | Jun | CAHN |
| AB | | 10MG | N76180 003 | Jul 01, 2002 | Jun | CAHN |
| AB | | 20MG | N76164 001 | Jul 01, 2002 | Jun | CAHN |
| AB | | 30MG | N76164 002 | Jul 01, 2002 | Jun | CAHN |
| AB | | 40MG | N76164 003 | Jul 01, 2002 | Jun | CAHN |
| AB | AUROBINDO | 2.5MG | N77622 001 | Feb 22, 2006 | Feb | NEWA |
| AB | | 5MG | N77622 002 | Feb 22, 2006 | Feb | NEWA |
| AB | | 10MG | N77622 003 | Feb 22, 2006 | Feb | NEWA |
| AB | | 20MG | N77622 004 | Feb 22, 2006 | Feb | NEWA |
| AB | | 30MG | N77622 005 | Feb 22, 2006 | Feb | NEWA |
| AB | | 40MG | N77622 006 | Feb 22, 2006 | Feb | NEWA |
| | PRINIVIL | | | | | |
| | @ MERCK | 2.5MG | N19558 006 | Jan 28, 1994 | Jun | DISC |

LITHIUM CARBONATE

TABLET; ORAL

LITHIUM CARBONATE

| | | | | | | |
|----|--------------------------------|-------|------------|--------------|-----|------|
| | @ PFIZER | 300MG | N16834 001 | | Aug | DISC |
| + | ROXANE | 300MG | N18558 001 | Jan 29, 1982 | Aug | CRLD |
| | TABLET, EXTENDED RELEASE; ORAL | | | | | |
| | ESKALITH CR | | | | | |
| | @ GLAXOSMITHKLINE | 450MG | N18152 001 | Mar 29, 1982 | Jul | DISC |
| | LITHIUM CARBONATE | | | | | |
| | @ BARR | 450MG | N76366 001 | Aug 21, 2003 | Jul | DISC |
| AB | + ROXANE | 450MG | N76691 001 | Jan 05, 2004 | Jul | CRLD |

LORAZEPAM

TABLET; ORAL

LORAZEPAM

| | | | | | | |
|----|-------------------|-------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 0.5MG | N71403 001 | Apr 21, 1987 | Jun | CAHN |
| AB | | 1MG | N71404 001 | Apr 21, 1987 | Jun | CAHN |
| AB | | 2MG | N71141 001 | Apr 21, 1987 | Jun | CAHN |
| AB | MYLAN | 0.5MG | N77657 001 | Mar 16, 2006 | Mar | NEWA |
| AB | | 1MG | N77657 002 | Mar 16, 2006 | Mar | NEWA |
| AB | | 2MG | N77657 003 | Mar 16, 2006 | Mar | NEWA |
| AB | VINTAGE PHARMS | 0.5MG | N77754 001 | May 10, 2006 | Apr | NEWA |
| AB | | 1MG | N77754 002 | May 10, 2006 | Apr | NEWA |
| AB | | 2MG | N77754 003 | May 10, 2006 | Apr | NEWA |

LOVASTATIN

TABLET; ORAL

LOVASTATIN

| | | | | | | |
|----|-------------------|------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 10MG | N75828 001 | Dec 17, 2001 | Jun | CAHN |
| AB | | 20MG | N75828 002 | Dec 17, 2001 | Jun | CAHN |
| AB | | 40MG | N75828 003 | Dec 17, 2001 | Jun | CAHN |
| AB | MUTUAL PHARM | 10MG | N77520 001 | Apr 14, 2006 | Apr | NEWA |
| AB | | 20MG | N77520 002 | Apr 14, 2006 | Apr | NEWA |
| AB | | 40MG | N77520 003 | Apr 14, 2006 | Apr | NEWA |
| | MEVACOR | | | | | |
| | @ MERCK | 10MG | N19643 002 | Mar 28, 1991 | Aug | DISC |

LOVASTATIN; NIACIN

TABLET, EXTENDED RELEASE; ORAL
ADVICOR

| | | | | | | |
|---|----------|------------|------------|--------------|-----|------|
| + | KOS LIFE | 20MG;750MG | N21249 002 | Dec 17, 2001 | Feb | CMFD |
| + | | 40MG;1GM | N21249 004 | Apr 27, 2006 | Jul | NEWA |

LUBIPROSTONE

CAPSULE; ORAL
AMITIZA

| | | | | | | |
|---|----------------|-------|------------|--------------|-----|------|
| + | SUCAMPO PHARMS | 24UGM | N21908 001 | Jan 31, 2006 | Jan | NEWA |
|---|----------------|-------|------------|--------------|-----|------|

MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

SOLUTION; INJECTION
NORMOCARB HF 25

| | | | | | | |
|---|---------------|---------------------------------------|------------|--------------|-----|------|
| + | DIALYSIS SUPS | 0.21GM/100ML;2.8GM/100ML;9.07GM/100ML | N21910 001 | Jul 26, 2006 | Jul | NEWA |
|---|---------------|---------------------------------------|------------|--------------|-----|------|

NORMOCARB HF 35

| | | | | | | |
|---|---------------|---------------------------------------|------------|--------------|-----|------|
| + | DIALYSIS SUPS | 0.21GM/100ML;3.97GM/100ML;8.3GM/100ML | N21910 002 | Jul 26, 2006 | Jul | NEWA |
|---|---------------|---------------------------------------|------------|--------------|-----|------|

MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE

TABLET, CHEWABLE; ORAL
ZEGERID

| | | | | | | |
|---|----------|------------------|------------|--------------|-----|------|
| | SANTARUS | 700MG;20MG;600MG | N21850 001 | Mar 24, 2006 | Mar | NEWA |
| + | | 700MG;40MG;600MG | N21850 002 | Mar 24, 2006 | Mar | NEWA |

MEDROXYPROGESTERONE ACETATE

INJECTABLE; SUBCUTANEOUS
DEPO-SUBQ PROVERA 104

| | | | | | | |
|---|----------------------|--------------|------------|--------------|-----|------|
| + | PHARMACIA AND UPJOHN | 104MG/0.65ML | N21583 001 | Dec 17, 2004 | Jan | CAHN |
|---|----------------------|--------------|------------|--------------|-----|------|

MEGESTROL ACETATE

SUSPENSION; ORAL
MEGESTROL ACETATE

| | | | | | | |
|----|--------|---------|------------|--------------|-----|------|
| AB | APOTEX | 40MG/ML | N77404 001 | Feb 16, 2006 | Jan | NEWA |
|----|--------|---------|------------|--------------|-----|------|

MELOXICAM

TABLET; ORAL
MELOXICAM

| | | | | | | |
|-----|--------------------|-------|------------|--------------|-----|------|
| AB | ACTAVIS TOTOWA | 7.5MG | N77938 001 | Jul 19, 2006 | Jul | NEWA |
| AB | | 15MG | N77938 002 | Jul 19, 2006 | Jul | NEWA |
| AB | APOTEX INC | 7.5MG | N77882 001 | Jul 20, 2006 | Jul | NEWA |
| AB | | 15MG | N77882 002 | Jul 20, 2006 | Jul | NEWA |
| >A> | AUROBINDO PHARMA | 7.5MG | N78008 001 | Oct 02, 2006 | Sep | NEWA |
| >A> | | 15MG | N78008 002 | Oct 02, 2006 | Sep | NEWA |
| AB | BRECKENRIDGE PHARM | 7.5MG | N77920 001 | Jul 19, 2006 | Jul | NEWA |
| AB | | 15MG | N77920 002 | Jul 19, 2006 | Jul | NEWA |
| AB | CARACO | 7.5MG | N77937 001 | Jul 19, 2006 | Jul | NEWA |
| AB | | 15MG | N77937 002 | Jul 19, 2006 | Jul | NEWA |
| AB | COREPHARMA | 7.5MG | N77930 001 | Jul 19, 2006 | Jul | NEWA |
| AB | | 15MG | N77930 002 | Jul 19, 2006 | Jul | NEWA |
| AB | DR REDDYS LABS INC | 7.5MG | N77931 001 | Jul 25, 2006 | Jul | NEWA |
| AB | | 15MG | N77931 002 | Jul 25, 2006 | Jul | NEWA |
| AB | GENPHARM | 7.5MG | N77934 001 | Jul 20, 2006 | Jul | NEWA |
| AB | | 15MG | N77934 002 | Jul 20, 2006 | Jul | NEWA |

TABLET; ORAL

MELOXICAM

| | | | | | | |
|-------|----------------------|-------|------------|--------------|-----|------|
| AB | GLENMARK PHARMS LTD | 7.5MG | N77932 001 | Jul 19, 2006 | Jul | NEWA |
| AB | | 15MG | N77932 002 | Jul 19, 2006 | Jul | NEWA |
| AB | LUPIN PHARMS | 7.5MG | N77944 001 | Jul 19, 2006 | Jul | NEWA |
| AB | | 15MG | N77944 002 | Jul 19, 2006 | Jul | NEWA |
| AB | MUTUAL PHARM | 7.5MG | N77935 001 | Jul 19, 2006 | Jul | NEWA |
| AB | | 15MG | N77935 002 | Jul 19, 2006 | Jul | NEWA |
| AB | MYLAN | 7.5MG | N77923 001 | Jul 19, 2006 | Jul | NEWA |
| AB | | 15MG | N77923 002 | Jul 19, 2006 | Jul | NEWA |
| AB | PAR PHARM | 7.5MG | N77933 001 | Jul 19, 2006 | Jul | NEWA |
| AB | | 15MG | N77933 002 | Jul 19, 2006 | Jul | NEWA |
| AB | ROXANE | 7.5MG | N77925 001 | Jul 19, 2006 | Jul | NEWA |
| AB | | 15MG | N77925 002 | Jul 19, 2006 | Jul | NEWA |
| AB | TEVA PHARMS | 7.5MG | N77936 001 | Jul 19, 2006 | Jul | NEWA |
| AB | | 15MG | N77936 002 | Jul 19, 2006 | Jul | NEWA |
| AB | WATSON LABS | 7.5MG | N77929 001 | Jul 19, 2006 | Jul | NEWA |
| AB | | 15MG | N77929 002 | Jul 19, 2006 | Jul | NEWA |
| AB | ZYDUS PHARMS USA | 7.5MG | N77921 001 | Jul 19, 2006 | Jul | NEWA |
| AB | | 15MG | N77921 002 | Jul 19, 2006 | Jul | NEWA |
| MOBIC | | | | | | |
| AB | BOEHRINGER INGELHEIM | 7.5MG | N20938 001 | Apr 13, 2000 | Jul | CFTG |
| AB | + | 15MG | N20938 002 | Aug 23, 2000 | Jul | CFTG |

MEPROBAMATE

TABLET; ORAL

MEPROBAMATE

| | | | | | | |
|---------|-----------------------|-------|------------|--|-----|------|
| | @ ROXANE | 600MG | N84332 001 | | Jan | DISC |
| | @ SANDOZ | 200MG | N14547 002 | | Jan | DISC |
| | @ | 400MG | N14547 001 | | Jan | DISC |
| | @ | 400MG | N80655 001 | | Jan | DISC |
| | @ SCHERER LABS | 400MG | N83343 001 | | Jan | DISC |
| | @ TABLICAPS | 400MG | N83494 001 | | Jan | DISC |
| AA | + WATSON LABS | 200MG | N83304 001 | | Jan | CRLD |
| | @ | 200MG | N85720 001 | | Jan | DISC |
| | + | 400MG | N83308 001 | | Jan | CRLD |
| | @ | 400MG | N85721 001 | | Jan | DISC |
| MILTOWN | | | | | | |
| | @ MEDPOINTE PHARM HLC | 200MG | N09698 004 | | Jan | DISC |
| | @ | 400MG | N09698 002 | | Jan | DISC |
| TRANMEP | | | | | | |
| | @ SOLVAY | 400MG | N16249 001 | | Jan | DISC |

MESALAMINE

ENEMA; RECTAL

ROWASA

| | | | | | | |
|---------------------|---------------------|----------|------------|--------------|-----|------|
| AB | + ALAVEN PHARM | 4GM/60ML | N19618 001 | Dec 24, 1987 | Apr | CAHN |
| SUPPOSITORY; RECTAL | | | | | | |
| CANASA | | | | | | |
| | @ AXCAN SCANDIPHARM | 500MG | N21252 001 | Jan 05, 2001 | May | DISC |
| ROWASA | | | | | | |
| | @ ALAVEN PHARM | 500MG | N19919 001 | Dec 18, 1990 | Jul | CAHN |

METARAMINOL BITARTRATE

INJECTABLE; INJECTION

ARAMINE

| | | | | | | |
|---|---------------|-----------------|------------|--------------|-----|------|
| @ | MERCK | EQ 10MG BASE/ML | N09509 002 | Dec 22, 1987 | Jun | DISC |
| + | ABRAXIS PHARM | EQ 10MG BASE/ML | N80722 001 | | Jun | CRLD |

METFORMIN HYDROCHLORIDE

TABLET; ORAL

METFORMIN HYDROCHLORIDE

| | | | | | | |
|----|--------------------|-------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 500MG | N76033 001 | Jan 24, 2002 | Jun | CAHN |
| AB | | 850MG | N76033 002 | Jan 24, 2002 | Jun | CAHN |
| AB | | 1GM | N76033 003 | Jan 24, 2002 | Jun | CAHN |
| AB | AMNEAL PHARM | 500MG | N77853 001 | Jul 28, 2006 | Jul | NEWA |
| AB | | 850MG | N77853 002 | Jul 28, 2006 | Jul | NEWA |
| AB | | 1GM | N77853 003 | Jul 28, 2006 | Jul | NEWA |
| AB | DR REDDYS LABS INC | 500MG | N77787 001 | Aug 23, 2006 | Aug | NEWA |
| AB | | 850MG | N77787 002 | Aug 23, 2006 | Aug | NEWA |
| AB | | 1GM | N77787 003 | Aug 23, 2006 | Aug | NEWA |
| AB | INTERPHARM | 500MG | N77880 001 | Jun 05, 2006 | May | NEWA |
| AB | | 850MG | N77880 002 | Jun 05, 2006 | May | NEWA |
| AB | | 1GM | N77880 003 | Jun 05, 2006 | May | NEWA |

TABLET, EXTENDED RELEASE; ORAL

GLUMETZA

| | | | | | | |
|----|-------------|-------|------------|--------------|-----|------|
| BX | DEPOMED INC | 500MG | N21748 001 | Jun 03, 2005 | Jan | CAHN |
| BX | + | 1GM | N21748 002 | Jun 03, 2005 | Aug | CRLD |
| BX | | 1GM | N21748 002 | Jun 03, 2005 | Jan | CAHN |

METFORMIN HYDROCHLORIDE

| | | | | | | |
|----|---------------------|-------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 500MG | N76450 001 | Oct 01, 2004 | Jun | CAHN |
| AB | | 750MG | N76878 001 | Apr 13, 2005 | Jun | CAHN |
| AB | NOSTRUM | 500MG | N76756 001 | Jul 26, 2006 | Jul | NEWA |
| AB | SUN PHARM INDS (IN) | 500MG | N77336 001 | Feb 09, 2006 | Jan | NEWA |
| AB | | 750MG | N77336 002 | Feb 09, 2006 | Jan | NEWA |

METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

| | | | | | | |
|----|--------------|------|------------|--------------|-----|------|
| + | CEDAR PHARMS | 20MG | N40547 004 | Feb 18, 2005 | May | CRLD |
| AB | JONES PHARMA | 5MG | N40320 001 | Mar 31, 2000 | May | CTNA |
| AB | | 10MG | N40320 002 | Mar 31, 2000 | May | CTNA |

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE

| | | | | | | | |
|----|---|---------------|------------------------------|------------|--------------|-----|------|
| AP | + | ABRAXIS PHARM | EQ 50MG BASE/2ML (25MG/ML) | N40263 001 | Feb 26, 1999 | Jun | CMFD |
| AP | + | | EQ 250MG BASE/10ML (25MG/ML) | N40263 002 | Feb 26, 1999 | Jul | CRLD |
| AP | | | EQ 250MG BASE/10ML (25MG/ML) | N40263 002 | Feb 26, 1999 | Jun | NEWA |

METHOTREXATE PRESERVATIVE FREE

| | | | | | | |
|---|------------------|-----------------------------|------------|--------------|-----|------|
| + | BEDFORD | EQ 1GM BASE/VIAL | N40632 001 | Aug 12, 2005 | Apr | CAIN |
| + | MAYNE PHARMA USA | EQ 1GM BASE/40ML (25MG/ML) | N11719 012 | Apr 13, 2005 | Jun | CPOT |
| + | | EQ 1GM BASE/40ML (25 MG/ML) | N11719 012 | Apr 13, 2005 | Apr | CPOT |

METHOTREXATE SODIUM

| | | | | | | | |
|----|---|---------|-----------------------------|------------|--------------|-----|------|
| AP | + | BEDFORD | EQ 50MG BASE/2ML (25MG/ML) | N89340 001 | Sep 16, 1986 | Apr | CPOT |
| + | | | EQ 100MG BASE/4ML (25MG/ML) | N89341 001 | Sep 16, 1986 | Apr | CTEC |

INJECTABLE; INJECTION

METHOTREXATE SODIUM

| | | | | | | |
|---|---------|-------------------------------|------------|--------------|-----|------|
| + | BEDFORD | EQ 200MG BASE/8ML (25MG/ML) | N89342 001 | Sep 16, 1986 | Apr | CTEC |
| + | | EQ 250MG BASE/10ML (25 MG/ML) | N89343 001 | Sep 16, 1986 | Apr | CTEC |

TABLET; ORAL

METHOTREXATE SODIUM

| | | | | | | | |
|----|---|--------------|---------------|------------|--|-----|------|
| AB | + | STADA PHARMS | EQ 2.5MG BASE | N08085 002 | | Aug | CAHN |
|----|---|--------------|---------------|------------|--|-----|------|

METHYLPHENIDATE

FILM, EXTENDED RELEASE; TRANSDERMAL

DAYTRANA

| | | | | | | |
|---|-------|---------------------|------------|--------------|-----|------|
| + | SHIRE | 10MG/9HR (1.1MG/HR) | N21514 001 | Apr 06, 2006 | Apr | NEWA |
| + | | 15MG/9HR (1.6MG/HR) | N21514 002 | Apr 06, 2006 | Apr | NEWA |
| + | | 20MG/9HR (2.2MG/HR) | N21514 003 | Apr 06, 2006 | Apr | NEWA |
| + | | 30MG/9HR (3.3MG/HR) | N21514 004 | Apr 06, 2006 | Apr | NEWA |

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

METADATE CD

| | | | | | | | |
|----|---|---------|------|------------|--------------|-----|------|
| BX | | UCB INC | 40MG | N21259 004 | Feb 19, 2006 | Feb | NEWA |
| | | | 50MG | N21259 005 | Feb 19, 2006 | Feb | NEWA |
| | + | | 60MG | N21259 006 | Feb 19, 2006 | Feb | NEWA |

RITALIN LA

| | | | | | | | |
|----|---|----------|------|------------|--------------|-----|------|
| BX | + | NOVARTIS | 40MG | N21284 003 | Jun 05, 2002 | Feb | CTEC |
|----|---|----------|------|------------|--------------|-----|------|

TABLET; ORAL

METHYLPHENIDATE HYDROCHLORIDE

| | | | | | | | |
|----|--|-------------------|------|------------|--------------|-----|------|
| AB | | ACTAVIS ELIZABETH | 5MG | N40321 001 | Feb 05, 2002 | Jun | CAHN |
| AB | | | 10MG | N40321 002 | Feb 05, 2002 | Jun | CAHN |
| AB | | | 20MG | N40321 003 | Feb 05, 2002 | Jun | CAHN |

TABLET, EXTENDED RELEASE; ORAL

METHYLPHENIDATE HYDROCHLORIDE

| | | | | | | | |
|----|--|-------------------|------|------------|--------------|-----|------|
| AB | | ACTAVIS ELIZABETH | 20MG | N75450 001 | Dec 21, 2001 | Jun | CAHN |
|----|--|-------------------|------|------------|--------------|-----|------|

METOCLOPRAMIDE HYDROCHLORIDE

SOLUTION; ORAL

METOCLOPRAMIDE

| | | | | | | | |
|----|---|--------------|-----------------|------------|--------------|-----|------|
| AA | + | JVL | EQ 5MG BASE/5ML | N74703 001 | Oct 31, 1997 | Aug | CRLD |
| | | @ VISTAPHARM | EQ 5MG BASE/5ML | N75051 001 | Jan 26, 2001 | Aug | DISC |

METOCLOPRAMIDE HYDROCHLORIDE

| | | | | | | | |
|----|--|----------------------|-----------------|------------|--------------|-----|------|
| AA | | ACTAVIS MID ATLANTIC | EQ 5MG BASE/5ML | N71340 001 | Aug 18, 1988 | Jul | CAHN |
| | | @ ROXANE | EQ 5MG BASE/5ML | N72038 001 | Dec 05, 1988 | Aug | DISC |
| | | @ TEVA | EQ 5MG BASE/5ML | N70819 001 | Jul 10, 1987 | Aug | DISC |
| | | @ | EQ 5MG BASE/5ML | N71315 001 | Jun 30, 1993 | Aug | DISC |

TABLET; ORAL

METOCLOPRAMIDE

| | | | | | | | |
|----|--|----------------|--------------|------------|--------------|-----|------|
| AB | | VINTAGE PHARMS | EQ 5MG BASE | N77878 001 | Aug 28, 2006 | Aug | NEWA |
| AB | | | EQ 10MG BASE | N77878 002 | Aug 28, 2006 | Aug | NEWA |

METOCLOPRAMIDE HYDROCHLORIDE

| | | | | | | | |
|----|--|-------------------|--------------|------------|--------------|-----|------|
| AB | | ACTAVIS ELIZABETH | EQ 10MG BASE | N70581 001 | Oct 17, 1985 | Jun | CAHN |
| AB | | MUTUAL PHARM | EQ 5MG BASE | N71536 002 | Jan 16, 1997 | Apr | CMFD |
| AB | | | EQ 10MG BASE | N71536 001 | Apr 28, 1993 | Apr | CMFD |

TABLET, ORALLY DISINTEGRATING; ORAL

REGLAN ODT

@ SCHWARZ PHARMA

@

| | | | | | | | |
|--|--|--|--------------|------------|--------------|-----|------|
| | | | EQ 5MG BASE | N21793 001 | Jun 10, 2005 | May | DISC |
| | | | EQ 10MG BASE | N21793 002 | Jun 10, 2005 | May | DISC |

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

METOPROLOL SUCCINATE

| | | | | | | |
|----|-------------|------------------|------------|--------------|-----|------|
| AB | SANDOZ | EQ 25MG TARTRATE | N76969 001 | Jul 31, 2006 | Jul | NEWA |
| | TOPROL-XL | | | | | |
| AB | ASTRAZENECA | EQ 25MG TARTRATE | N19962 004 | Feb 05, 2001 | Jul | CFTG |

METRONIDAZOLE

GEL; TOPICAL

METROGEL

| | | | | | | |
|----|--------------------|-------|------------|--------------|-----|------|
| AB | + GALDERMA LABS LP | 0.75% | N19737 001 | Nov 22, 1988 | May | CFTG |
| | METRONIDAZOLE | | | | | |
| AB | ALTANA | 0.75% | N77018 001 | Jun 06, 2006 | May | NEWA |
| AB | QLT USA | 0.75% | N77547 001 | Jul 13, 2006 | Jun | NEWA |
| AB | TARO | 0.75% | N77819 001 | Jul 18, 2006 | Jul | NEWA |

LOTION; TOPICAL

METROLOTION

| | | | | | | |
|----|--------------------|-------|------------|--------------|-----|------|
| AB | + GALDERMA LABS LP | 0.75% | N20901 001 | Nov 24, 1998 | May | CFTG |
| | METRONIDAZOLE | | | | | |
| AB | ALTANA | 0.75% | N77197 001 | May 24, 2006 | May | NEWA |

MICAFUNGIN SODIUM

INJECTABLE; IV (INFUSION)

MYCAMINE

ASTELLAS

100MG/VIAL

N21506 003 Jun 27, 2006 Jun NEWA

MICONAZOLE NITRATE

SUPPOSITORY; VAGINAL

MICONAZOLE NITRATE

| | | | | | | |
|----|----------------------|-------|------------|--------------|-----|------|
| AB | ACTAVIS MID ATLANTIC | 200MG | N73508 001 | Nov 19, 1993 | Jun | CAHN |
| | MONISTAT 3 | | | | | |
| AB | + PERSONAL PRODS | 200MG | N18888 001 | Aug 15, 1984 | Aug | CAHN |

MICONAZOLE NITRATE; PETROLATUM, WHITE; ZINC OXIDE

OINTMENT; TOPICAL

VUSION

+ BARRIER

0.25%;81.35%;15%

N21026 001 Feb 16, 2006 Feb NEWA

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

| | | | | | | |
|----|-----------|----------------|------------|--------------|-----|------|
| AP | + HOSPIRA | EQ 5MG BASE/ML | N75293 002 | Jun 20, 2000 | Mar | CRLD |
|----|-----------|----------------|------------|--------------|-----|------|

MIDODRINE HYDROCHLORIDE

TABLET; ORAL

MIDODRINE HYDROCHLORIDE

| | | | | | | |
|----|------------|-------|------------|--------------|-----|------|
| AB | APOTEX INC | 2.5MG | N77746 001 | Sep 12, 2006 | Aug | NEWA |
| AB | | 5MG | N77746 002 | Sep 12, 2006 | Aug | NEWA |
| AB | | 10MG | N77746 003 | Sep 12, 2006 | Aug | NEWA |

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

MINOCIN

| | | | | | | |
|----|--------------|--------------|------------|--------------|-----|------|
| AB | TRIAx PHARMS | EQ 50MG BASE | N50649 001 | May 31, 1990 | Feb | CAHN |
|----|--------------|--------------|------------|--------------|-----|------|

CAPSULE; ORAL

MINOCIN

| | | | | | | |
|----|----------------|---------------|------------|--------------|-----|------|
| | @ TRIAX PHARMS | EQ 75MG BASE | N50649 003 | Feb 12, 2001 | Feb | CAHN |
| AB | + | EQ 100MG BASE | N50649 002 | May 31, 1990 | Feb | CAHN |

TABLET, EXTENDED RELEASE; ORAL

SOLODYN

| | | | | | | |
|--|---------|---------------|------------|--------------|-----|------|
| | MEDICIS | EQ 45MG BASE | N50808 001 | May 08, 2006 | May | NEWA |
| | | EQ 90MG BASE | N50808 002 | May 08, 2006 | May | NEWA |
| | + | EQ 135MG BASE | N50808 003 | May 08, 2006 | May | NEWA |

MIRTAZAPINE

TABLET; ORAL

MIRTAZAPINE

| | | | | | | |
|----|-------------------|------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 15MG | N76308 001 | Jun 20, 2003 | Jun | CAHN |
| AB | | 30MG | N76308 002 | Jun 20, 2003 | Jun | CAHN |
| AB | | 45MG | N76308 003 | Jun 20, 2003 | Jun | CAHN |

TABLET, ORALLY DISINTEGRATING; ORAL

MIRTAZAPINE

| | | | | | | |
|----|----------------------|------|------------|--------------|-----|------|
| AB | AUROBINDO PHARMA LTD | 45MG | N77376 004 | Feb 28, 2006 | Feb | NEWA |
| AB | BARR | 45MG | N76307 003 | Feb 28, 2006 | Feb | NEWA |

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

MITOXANTRONE

| | | | | | | | |
|----|------------------|------------------------------|------------------------------|--------------|--------------|------|------|
| AP | AM PHARM | EQ 20MG BASE/10ML (2MG/ML) | N77496 001 | Apr 11, 2006 | Mar | NEWA | |
| AP | | EQ 25MG BASE/12.5ML (2MG/ML) | N77496 002 | Apr 11, 2006 | Mar | NEWA | |
| AP | | EQ 30MG BASE/15ML (2MG/ML) | N77496 003 | Apr 11, 2006 | Mar | NEWA | |
| AP | BEDFORD | EQ 20MG BASE/10ML (2MG/ML) | N76611 001 | Apr 11, 2006 | Mar | NEWA | |
| AP | | EQ 25MG BASE/12.5ML (2MG/ML) | N76611 002 | Apr 11, 2006 | Mar | NEWA | |
| AP | | EQ 30MG BASE/15ML (2MG/ML) | N76611 003 | Apr 11, 2006 | Mar | NEWA | |
| AP | MAYNE PHARMA USA | EQ 20MG BASE/10ML (2MG/ML) | N76871 001 | Apr 11, 2006 | Mar | NEWA | |
| AP | | EQ 25MG BASE/12.5ML (2MG/ML) | N76871 002 | Apr 11, 2006 | Mar | NEWA | |
| AP | | EQ 30MG BASE/15ML (2MG/ML) | N76871 003 | Apr 11, 2006 | Mar | NEWA | |
| AP | SICOR PHARMS | EQ 20MG BASE/10ML (2MG/ML) | N77356 001 | Apr 11, 2006 | Mar | NEWA | |
| AP | | EQ 25MG BASE/12.5ML (2MG/ML) | N77356 002 | Apr 11, 2006 | Mar | NEWA | |
| AP | | EQ 30MG BASE/15ML (2MG/ML) | N77356 003 | Apr 11, 2006 | Mar | NEWA | |
| | NOVANTRONE | | | | | | |
| AP | + | SERONO INC | EQ 20MG BASE/10ML (2MG/ML) | N19297 001 | Dec 23, 1987 | Mar | CFTG |
| AP | + | | EQ 25MG BASE/12.5ML (2MG/ML) | N19297 002 | Dec 23, 1987 | Mar | CFTG |
| AP | + | | EQ 30MG BASE/15ML (2MG/ML) | N19297 003 | Dec 23, 1987 | Mar | CFTG |

>D> MIVACURIUM CHLORIDE

>D> INJECTABLE; INJECTION

>D> MIVACRON

| | | | | | | | |
|-----|---|-------------------------|----------------------|------------|--------------|-----|------|
| >D> | + | ABBOTT | EQ 2MG BASE/ML | N20098 001 | Jan 22, 1992 | Sep | DISC |
| >A> | | @ | EQ 2MG BASE/ML | N20098 001 | Jan 22, 1992 | Sep | DISC |
| >D> | | MIVACRON IN DEXTROSE 5% | IN PLASTIC CONTAINER | | | | |
| >D> | + | ABBOTT | EQ 50MG BASE/100ML | N20098 003 | Jan 22, 1992 | Sep | DISC |
| >A> | | @ | EQ 50MG BASE/100ML | N20098 003 | Jan 22, 1992 | Sep | DISC |

MOMETASONE FUROATE

CREAM; TOPICAL

MOMETASONE FUROATE

| | | | | | | |
|----|--------------|------|------------|--------------|-----|------|
| AB | G AND W LABS | 0.1% | N77447 001 | May 22, 2006 | May | NEWA |
|----|--------------|------|------------|--------------|-----|------|

LOTION; TOPICAL

MOMETASONE FUROATE

| | | | | | | |
|----|---------|------|------------|--------------|-----|------|
| AB | PERRIGO | 0.1% | N77180 001 | Apr 06, 2005 | Mar | CAHN |
| AB | TARO | 0.1% | N76788 001 | Mar 15, 2006 | Feb | NEWA |

OINTMENT; TOPICAL

MOMETASONE FUROATE

| | | | | | | |
|----|--------------|------|------------|--------------|-----|------|
| AB | G AND W LABS | 0.1% | N77401 001 | Jun 20, 2006 | Jun | NEWA |
|----|--------------|------|------------|--------------|-----|------|

MORPHINE SULFATE

INJECTABLE; INJECTION

MORPHINE SULFATE

| | | | | | | |
|--|---------|--------|------------|--------------|-----|------|
| | HOSPIRA | 5MG/ML | N19916 002 | Mar 30, 2006 | Mar | NEWA |
|--|---------|--------|------------|--------------|-----|------|

NABILONE

CAPSULE; ORAL

CESAMET

| | | | | | | |
|---|---------|-----|------------|--------------|-----|------|
| + | VALEANT | 1MG | N18677 001 | Dec 26, 1985 | May | CMFD |
|---|---------|-----|------------|--------------|-----|------|

NABUMETONE

TABLET; ORAL

NABUMETONE

| | | | | | | |
|----|----------------------|-------|------------|--------------|-----|------|
| | @ COPLEY PHARM | 750MG | N75179 001 | Jun 06, 2000 | Jun | DISC |
| AB | PAR PHARM | 500MG | N76009 001 | Jan 24, 2003 | Apr | CAHN |
| AB | | 750MG | N76009 002 | Jan 24, 2003 | Apr | CAHN |
| AB | + TEVA | 750MG | N75189 002 | Sep 24, 2001 | Jul | CRLD |
| | RELAFEN | | | | | |
| | @ SMITHKLINE BEECHAM | 500MG | N19583 001 | Dec 24, 1991 | Jul | DISC |
| | @ | 750MG | N19583 002 | Dec 24, 1991 | Jul | DISC |

NADOLOL

TABLET; ORAL

CORGARD

| | | | | | | |
|----|-------------|-------|------------|--------------|-----|------|
| AB | KING PHARMS | 20MG | N18063 005 | Oct 28, 1986 | Jun | CAHN |
| AB | | 40MG | N18063 001 | | Jun | CAHN |
| AB | | 80MG | N18063 002 | | Jun | CAHN |
| AB | | 120MG | N18063 003 | | Jun | CAHN |
| AB | + | 160MG | N18063 004 | | Jun | CAHN |

NAFCILLIN SODIUM

INJECTABLE; INJECTION

NAFCILLIN SODIUM

| | | | | | | |
|----|----------|-------------------|------------|--------------|-----|------|
| AP | + SANDOZ | EQ 1GM BASE/VIAL | N62527 002 | Aug 02, 1984 | Apr | CRLD |
| AP | + | EQ 1GM BASE/VIAL | N62732 001 | Dec 23, 1986 | Apr | CRLD |
| AP | + | EQ 2GM BASE/VIAL | N62527 003 | Aug 02, 1984 | Apr | CRLD |
| AP | + | EQ 2GM BASE/VIAL | N62732 002 | Dec 23, 1986 | Apr | CRLD |
| AP | + | EQ 10GM BASE/VIAL | N62527 004 | Aug 02, 1984 | Apr | CRLD |

NAFTIFINE HYDROCHLORIDE

CREAM; TOPICAL

NAFTIN

| | | | | | | |
|-----|-------------|----|------------|--------------|-----|------|
| >D> | MERZ PHARMS | 1% | N19599 001 | Feb 29, 1988 | Sep | CRLD |
| >A> | + | 1% | N19599 001 | Feb 29, 1988 | Sep | CRLD |

NALTREXONE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR
VIVITROL

+ ALKERMES 380MG/VIAL N21897 001 Apr 13, 2006 Apr NEWA

NAPROXEN

TABLET, DELAYED RELEASE; ORAL

NAPROXEN

AB ACTAVIS ELIZABETH 375MG N74936 001 Feb 24, 1998 Jun CAHN
AB 500MG N74936 002 Feb 24, 1998 Jun CAHN

NEOMYCIN SULFATE

TABLET; ORAL

NEOMYCIN SULFATE

@ BRISTOL MYERS SQUIBB 500MG N60365 001 Jul CPOT
@ LANNETT 500MG N60607 001 Jul CPOT
@ LILLY 500MG N60385 001 Jul CPOT
@ ROXANE 500MG N62173 001 Jul CPOT
@ SANDOZ 500MG N61586 001 Jul CPOT
AB + TEVA 500MG N60304 001 Jul CFTG
AB X GEN PHARMS 500MG N65220 001 Jul 28, 2006 Jul NEWA

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION; IRRIGATION

NEOMYCIN AND POLYMYXIN B SULFATE

AT WATSON LABS EQ 40MG BASE/ML;200,000 UNITS/ML N62664 001 Apr 08, 1986 Jul CMFD
AT X GEN PHARMS EQ 40MG BASE/ML;200,000 UNITS/ML N65106 001 Jan 31, 2006 Jun CTNA
AT EQ 800MG BASE/20ML;4,000,000 UNITS/20ML N65108 001 Jan 31, 2006 Jun CTNA
NEOSPORIN AND POLYMYXIN B SULFATE
AT X GEN PHARMS EQ 40MG BASE/ML;200,000 UNITS/ML N65106 001 Jan 31, 2006 Jan NEWA
AT EQ 800MG BASE/20ML;4,000,000 UNITS/20ML (EQ 40MG BASE/ML;200,000 UNITS/ML) N65108 001 Jan 31, 2006 Jan NEWA
NEOSPORIN G.U. IRRIGANT
AT + MONARCH PHARMS EQ 40MG BASE/ML;200,000 UNITS/ML N60707 001 Jan CTEC
AT + EQ 800MG BASE/20ML;4,000,000 UNITS/20ML (EQ 40MG BASE/ML;200,000 UNITS/ML) N60707 002 Jan NEWA

NIACIN

TABLET, EXTENDED RELEASE; ORAL

>D> NIACIN
>D> AB BARR 500MG N76378 001 Apr 26, 2005 Sep DISC
>A> @ 500MG N76378 001 Apr 26, 2005 Sep DISC
>D> AB 750MG N76378 002 Apr 26, 2005 Sep DISC
>A> @ 750MG N76378 002 Apr 26, 2005 Sep DISC
>D> AB 1GM N76250 001 Apr 14, 2005 Sep DISC
>A> @ 1GM N76250 001 Apr 14, 2005 Sep DISC
NIASPAN
>D> AB + KOS LIFE 500MG N20381 002 Jul 28, 1997 Sep CTEC
>A> + 500MG N20381 002 Jul 28, 1997 Sep CTEC
>D> AB + 750MG N20381 003 Jul 28, 1997 Sep CTEC
>A> + 750MG N20381 003 Jul 28, 1997 Sep CTEC
>D> AB + 1GM N20381 004 Jul 28, 1997 Sep CTEC
>A> + 1GM N20381 004 Jul 28, 1997 Sep CTEC

NICARDIPINE HYDROCHLORIDE

CAPSULE; ORAL

NICARDIPINE HYDROCHLORIDE

| | | | | | | |
|----|------|------|------------|--------------|-----|------|
| AB | BARR | 20MG | N74439 001 | Dec 10, 1996 | Apr | CAHN |
| AB | | 30MG | N74439 002 | Dec 10, 1996 | Apr | CAHN |

INJECTABLE; INJECTION

CARDENE

| | | | | | | |
|---|-------------------|----------|------------|--------------|-----|------|
| + | PDL BIOPHARMA INC | 2.5MG/ML | N19734 001 | Jan 30, 1992 | Jan | CAHN |
|---|-------------------|----------|------------|--------------|-----|------|

NIFEDIPINE

CAPSULE; ORAL

NIFEDIPINE

| | | | | | | |
|----|-------------------|------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 10MG | N72579 001 | Jan 08, 1991 | Jun | CAHN |
| AB | | 20MG | N72556 001 | Sep 20, 1990 | Jun | CAHN |

TABLET, EXTENDED RELEASE; ORAL

AFEDITAB CR

| | | | | | | |
|-----|-------------|------|------------|--------------|-----|------|
| AB1 | WATSON LABS | 30MG | N75128 001 | Mar 10, 2000 | Jan | CAHN |
| AB1 | | 60MG | N75659 001 | Oct 26, 2001 | Jan | CAHN |

NISOLDIPINE

TABLET, EXTENDED RELEASE; ORAL

SULAR

| | | | | | | |
|---|-------------------|------|------------|--------------|-----|------|
| + | SCIELE PHARMA INC | 10MG | N20356 001 | Feb 02, 1995 | Jun | CAHN |
| | | 20MG | N20356 002 | Feb 02, 1995 | Jun | CAHN |
| + | | 30MG | N20356 003 | Feb 02, 1995 | Jun | CAHN |
| + | | 40MG | N20356 004 | Feb 02, 1995 | Jun | CAHN |

NITROGLYCERIN

AEROSOL; SUBLINGUAL

NITROLINGUAL

| | | | | | | |
|---|--------------|-------------|------------|--------------|-----|------|
| @ | POHL BOSKAMP | 0.4MG/SPRAY | N18705 001 | Oct 31, 1985 | Jun | CAHN |
|---|--------------|-------------|------------|--------------|-----|------|

SPRAY, METERED; SUBLINGUAL

NITROLINGUAL PUMPSPRAY

| | | | | | | |
|---|--------------|-------------|------------|--------------|-----|------|
| + | POHL BOSKAMP | 0.4MG/SPRAY | N18705 002 | Jan 10, 1997 | Jun | CAHN |
|---|--------------|-------------|------------|--------------|-----|------|

NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

NOREPINEPHRINE BITARTRATE

| | | | | | | |
|----|---------------|----------------|------------|--------------|-----|------|
| AP | METRICS PHARM | EQ 1MG BASE/ML | N40522 001 | Sep 30, 2004 | May | CAHN |
|----|---------------|----------------|------------|--------------|-----|------|

NORTRIPTYLINE HYDROCHLORIDE

SOLUTION; ORAL

NORTRIPTYLINE HYDROCHLORIDE

| | | | | | | |
|----|------|------------------|------------|--------------|-----|------|
| AA | TARO | EQ 10MG BASE/5ML | N77965 001 | Jun 20, 2006 | Jun | NEWA |
|----|------|------------------|------------|--------------|-----|------|

NYSTATIN

CREAM; TOPICAL

NYSTATIN

| | | | | | | |
|----|----------------------|------------------|------------|--------------|-----|------|
| AT | ACTAVIS MID ATLANTIC | 100,000 UNITS/GM | N62949 001 | Jun 13, 1988 | Jun | CAHN |
| | @ TARO | 100,000 UNITS/GM | N62457 001 | Jul 28, 1983 | May | DISC |
| AT | VINTAGE | 100,000 UNITS/GM | N65315 001 | May 31, 2006 | May | NEWA |

OINTMENT; TOPICAL

NYSTATIN

| | | | | | | |
|----|----------------------|------------------|------------|--------------|-----|------|
| AT | ACTAVIS MID ATLANTIC | 100,000 UNITS/GM | N62840 001 | Nov 13, 1987 | Jun | CAHN |
|----|----------------------|------------------|------------|--------------|-----|------|

POWDER; ORAL

NILSTAT

| | | | | | | |
|-----|--------------------|------|------------|--------------|-----|------|
| >D> | @ CLONMEL HLTHCARE | 100% | N50576 001 | Dec 22, 1983 | Sep | CAHN |
| >A> | @ STADA PHARMS | 100% | N50576 001 | Dec 22, 1983 | Sep | CAHN |

POWDER; TOPICAL

NYSTATIN

| | | | | | | |
|----|----------|------------------|------------|--------------|-----|------|
| AT | KV PHARM | 100,000 UNITS/GM | N65321 001 | Aug 18, 2006 | Aug | NEWA |
|----|----------|------------------|------------|--------------|-----|------|

SUSPENSION; ORAL

NYSTATIN

| | | | | | | |
|----|----------------------|------------------|------------|--------------|-----|------|
| AA | ACTAVIS MID ATLANTIC | 100,000 UNITS/ML | N62349 001 | Jul 14, 1982 | Jul | CAHN |
| AA | TARO | 100,000 UNITS/ML | N62876 001 | Feb 29, 1988 | May | CMFD |

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYKACET

| | | | | | | |
|----|----------------------|-----------------------|------------|--------------|-----|------|
| AT | ACTAVIS MID ATLANTIC | 100,000 UNITS/GM;0.1% | N62367 001 | May 28, 1985 | Jun | CAHN |
|----|----------------------|-----------------------|------------|--------------|-----|------|

OINTMENT; TOPICAL

MYKACET

| | | | | | | |
|----|----------------------|-----------------------|------------|--------------|-----|------|
| AT | ACTAVIS MID ATLANTIC | 100,000 UNITS/GM;0.1% | N62733 001 | Mar 09, 1987 | Jun | CAHN |
|----|----------------------|-----------------------|------------|--------------|-----|------|

OCTREOTIDE ACETATE

INJECTABLE; INJECTION

OCTREOTIDE ACETATE

| | | | | | | |
|----|----------|--|------------|--------------|-----|------|
| AP | AM PHARM | EQ 0.2MG BASE/ML | N77450 001 | Feb 10, 2006 | Jan | NEWA |
| AP | | EQ 1MG BASE/ML | N77450 002 | Feb 10, 2006 | Jan | NEWA |
| | | OCTREOTIDE ACETATE (PRESERVATIVE FREE) | | | | |
| AP | AM PHARM | EQ 0.05MG BASE/ML | N77457 001 | Feb 10, 2006 | Jan | NEWA |
| AP | | EQ 0.1MG BASE/ML | N77457 002 | Feb 10, 2006 | Jan | NEWA |
| AP | | EQ 0.5MG BASE/ML | N77457 003 | Feb 10, 2006 | Jan | NEWA |

OFLOXACIN

SOLUTION/DROPS; OTIC

OFLOXACIN

| | | | | | | |
|----|------------|------|------------|--------------|-----|------|
| AT | APOTEX INC | 0.3% | N76527 001 | Nov 18, 2005 | Aug | CAHN |
|----|------------|------|------------|--------------|-----|------|

TABLET; ORAL

OFLOXACIN

| | | | | | | |
|----|--------------------|-------|------------|--------------|-----|------|
| AB | DR REDDYS LABS LTD | 200MG | N77098 001 | Feb 10, 2006 | Jan | NEWA |
| AB | | 300MG | N77098 002 | Feb 10, 2006 | Jan | NEWA |
| AB | | 400MG | N77098 003 | Feb 10, 2006 | Jan | NEWA |

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE; ORAL

ZEGERID

| | | | | | | |
|---|----------|------------|------------|--------------|-----|------|
| | SANTARUS | 20MG;1.1GM | N21849 001 | Feb 27, 2006 | Feb | NEWA |
| + | | 40MG;1.1GM | N21849 002 | Feb 27, 2006 | Feb | NEWA |

FOR SUSPENSION; ORAL

ZEGERID

| | | | | | | |
|---|----------|---------------------------|------------|--------------|-----|------|
| | SANTARUS | 20MG/PACKET;1.68GM/PACKET | N21636 001 | Jun 15, 2004 | Feb | CAIN |
| + | | 40MG/PACKET;1.68GM/PACKET | N21706 001 | Dec 21, 2004 | Feb | CAIN |

ORPHENADRINE CITRATE

INJECTABLE; INJECTION

ORPHENADRINE CITRATE

| | | | | | | |
|----|-------|---------|------------|--------------|-----|------|
| AP | AKORN | 30MG/ML | N40484 001 | May 24, 2006 | May | NEWA |
|----|-------|---------|------------|--------------|-----|------|

OXALIPLATIN

INJECTABLE; IV (INFUSION)

ELOXATIN

@ SANOFI AVENTIS US 50MG/VIAL
 @ 100MG/VIAL

N21492 001 Aug 09, 2002 Jun DISC
 N21492 002 Aug 09, 2002 Jun DISC

OXAPROZIN

TABLET; ORAL

OXAPROZIN

AB ACTAVIS ELIZABETH 600MG

N75843 001 Oct 03, 2001 Jun CAHN

OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

AB ACTAVIS ELIZABETH 10MG
 AB 15MG
 AB 30MG

N72251 001 Apr 14, 1988 Jun CAHN
 N72252 001 Apr 14, 1988 Jun CAHN
 N72253 001 Apr 14, 1988 Jun CAHN

OXYCODONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

OXYCONTIN

>A> PURDUE PHARMA LP 15MG
 >A> 30MG
 >A> 60MG

N20553 006 Sep 18, 2006 Sep NEWA
 N20553 007 Sep 18, 2006 Sep NEWA
 N20553 008 Sep 18, 2006 Sep NEWA

OXYMETHOLONE

TABLET; ORAL

ANADROL-50

+ ALAVEN PHARM 50MG

N16848 001 Apr CAHN

OXYMORPHONE HYDROCHLORIDE

TABLET; ORAL

OPANA

ENDO PHARMS 5MG
 + 10MG

N21611 001 Jun 22, 2006 Jun NEWA
 N21611 002 Jun 22, 2006 Jun NEWA

TABLET, EXTENDED RELEASE; ORAL

OPANA ER

ENDO PHARMS 5MG
 10MG
 20MG
 + 40MG

N21610 001 Jun 22, 2006 Jun NEWA
 N21610 002 Jun 22, 2006 Jun NEWA
 N21610 003 Jun 22, 2006 Jun NEWA
 N21610 004 Jun 22, 2006 Jun NEWA

PACLITAXEL

FOR SUSPENSION; IV (INFUSION)

ABRAXANE

+ ABRAXIS BIOSCIENCE 100MG/VIAL

N21660 001 Jan 07, 2005 Apr CAHN

PARICALCITOL

INJECTABLE; INJECTION

ZEMPLAR

+ ABBOTT 0.002MG/ML

N20819 002 Feb 01, 2000 Aug CRLD

PAROXETINE MESYLATE

TABLET; ORAL

PEXEVA

| | | | | | |
|------------|--------------|------------|--------------|-----|------|
| JDS PHARMS | EQ 10MG BASE | N21299 001 | Jul 03, 2003 | Apr | CAHN |
| | EQ 20MG BASE | N21299 002 | Jul 03, 2003 | Apr | CAHN |
| | EQ 30MG BASE | N21299 003 | Jul 03, 2003 | Apr | CAHN |
| + | EQ 40MG BASE | N21299 004 | Jul 03, 2003 | Apr | CAHN |

PEGAPTANIB SODIUM

INJECTABLE; INTRAVITREAL

MACUGEN

| | | | | | | |
|---|-------------|----------------------|------------|--------------|-----|------|
| + | OSI EYETECH | EQ 0.3MG ACID/0.09ML | N21756 001 | Dec 17, 2004 | May | CAHN |
|---|-------------|----------------------|------------|--------------|-----|------|

PENICILLAMINE

CAPSULE; ORAL

CUPRIMINE

| | | | | | | |
|---|-------|-------|------------|--|-----|------|
| @ | ATON | 125MG | N19853 002 | | Aug | CAHN |
| + | | 250MG | N19853 001 | | Aug | CAHN |
| @ | MERCK | 125MG | N19853 002 | | Jun | DISC |

PENICILLIN V POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN V POTASSIUM

| | | | | | | |
|----|----------------|-------------------|------------|--|-----|------|
| AA | AM ANTIBIOTICS | EQ 125MG BASE/5ML | N61529 001 | | Jan | CAHN |
| AA | | EQ 250MG BASE/5ML | N61529 002 | | Jan | CAHN |

TABLET; ORAL

PENICILLIN V POTASSIUM

| | | | | | | |
|---|----------------|---------------|------------|--|-----|------|
| @ | AM ANTIBIOTICS | EQ 250MG BASE | N61528 001 | | Jan | CAHN |
| @ | | EQ 500MG BASE | N61528 002 | | Jan | CAHN |

PENTOSTATIN

INJECTABLE; INJECTION

NIPENT

| | | | | | | |
|---|------------------|-----------|------------|--------------|-----|------|
| + | MAYNE PHARMA USA | 10MG/VIAL | N20122 001 | Oct 11, 1991 | Aug | CAHN |
|---|------------------|-----------|------------|--------------|-----|------|

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL

PENTOXIFYLLINE

| | | | | | | |
|----|-------------------|-------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 400MG | N74878 001 | Jul 09, 1997 | Jun | CAHN |
| AB | RADIUS PHARMS | 400MG | N74877 001 | Jul 08, 1997 | Jul | CAHN |

PERGOLIDE MESYLATE

TABLET; ORAL

PERGOLIDE MESYLATE

| | | | | | | |
|----|-----------|----------------|------------|--------------|-----|------|
| AB | PAR PHARM | EQ 0.05MG BASE | N76061 001 | Nov 27, 2002 | Apr | CAHN |
| AB | | EQ 0.25MG BASE | N76061 002 | Nov 27, 2002 | Apr | CAHN |
| AB | | EQ 1MG BASE | N76061 003 | Nov 27, 2002 | Apr | CAHN |
| | PERMAX | | | | | |
| AB | VALEANT | EQ 0.05MG BASE | N19385 001 | Dec 30, 1988 | Jan | CRLD |
| AB | + | EQ 0.25MG BASE | N19385 002 | Dec 30, 1988 | Jan | CRLD |

PERMETHRIN

CREAM; TOPICAL

PERMETHRIN

| | | | | | | | |
|----|----------------------|----|--------|-----|--------------|-----|------|
| AB | ACTAVIS MID ATLANTIC | 5% | N74806 | 001 | Jan 23, 1998 | Jun | CAHN |
|----|----------------------|----|--------|-----|--------------|-----|------|

PHENDIMETRAZINE TARTRATE

TABLET; ORAL

CAM-METRAZINE

| | | | | | | | |
|--|------------------|------|--------|-----|--|-----|------|
| | @ TG UNITED LABS | 35MG | N83922 | 001 | | May | CAHN |
| | @ | 35MG | N85318 | 001 | | May | CAHN |
| | @ | 35MG | N85320 | 001 | | May | CAHN |
| | @ | 35MG | N85321 | 001 | | May | CAHN |

PHENDIMETRAZINE TARTRATE

| | | | | | | | |
|--|------------------|------|--------|-----|--------------|-----|------|
| | @ TG UNITED LABS | 35MG | N85761 | 001 | | May | CAHN |
| | @ | 35MG | N85941 | 001 | Jun 27, 1983 | May | CAHN |

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

PHENTERMINE HYDROCHLORIDE

| | | | | | | | |
|--|------------------|---------|--------|-----|--------------|-----|------|
| | @ TG UNITED LABS | 18.75MG | N88576 | 001 | May 23, 1984 | May | CAHN |
| | @ | 30MG | N85417 | 001 | | May | CAHN |
| | @ | 30MG | N86732 | 002 | | May | CAHN |
| | @ | 30MG | N87215 | 001 | | May | CAHN |
| | @ | 37.5MG | N87915 | 001 | Dec 22, 1983 | May | CAHN |
| | @ | 37.5MG | N87918 | 001 | Dec 22, 1983 | May | CAHN |
| | @ | 37.5MG | N87930 | 001 | Oct 14, 1983 | May | CAHN |
| | @ | 37.5MG | N88610 | 001 | Jun 04, 1984 | May | CAHN |
| | @ | 37.5MG | N88611 | 001 | Jun 04, 1984 | May | CAHN |
| | @ | 37.5MG | N88625 | 001 | Aug 23, 1984 | May | CAHN |

TABLET; ORAL

PHENTERMINE HYDROCHLORIDE

| | | | | | | | |
|----|-------------------|--------|--------|-----|--------------|-----|------|
| AA | ACTAVIS ELIZABETH | 37.5MG | N40276 | 001 | Nov 25, 1998 | Jun | CAHN |
| | @ TG UNITED LABS | 8MG | N83923 | 001 | | May | CAHN |
| | @ | 8MG | N85319 | 001 | | May | CAHN |
| | @ | 37.5MG | N87805 | 001 | Dec 06, 1982 | May | CAHN |
| | @ | 37.5MG | N88596 | 001 | Apr 04, 1984 | May | CAHN |

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETH VC PLAIN

| | | | | | | | |
|---|-----------------------|---------------------|--------|-----|--------------|-----|------|
| + | ACTAVIS MID ATLANTIC | 5MG/5ML; 6.25MG/5ML | N88761 | 001 | Nov 08, 1984 | Jul | CAHN |
| + | ALPHARMA US PHARMS | 5MG/5ML; 6.25MG/5ML | N88761 | 001 | Nov 08, 1984 | Jan | CTEC |
| | PROMETHAZINE VC PLAIN | | | | | | |
| | @ MORTON GROVE | 5MG/5ML; 6.25MG/5ML | N88897 | 001 | Jan 04, 1985 | Jan | DISC |

PHENYTOIN

SUSPENSION; ORAL

PHENYTOIN

| | | | | | | | |
|----|----------------------|-----------|--------|-----|--------------|-----|------|
| AB | ACTAVIS MID ATLANTIC | 125MG/5ML | N89892 | 001 | Sep 25, 1992 | Jul | CAHN |
|----|----------------------|-----------|--------|-----|--------------|-----|------|

PHENYTOIN SODIUM

CAPSULE; ORAL

EXTENDED PHENYTOIN SODIUM

| | | | | | | | |
|----|------|----------------|--------|-----|--------------|-----|------|
| AB | TARO | 100MG EXTENDED | N40684 | 001 | Sep 05, 2006 | Aug | NEWA |
|----|------|----------------|--------|-----|--------------|-----|------|

INJECTABLE; INJECTION

PHENYTOIN SODIUM

| | | | | | | |
|----|--------------------|---------|------------|--------------|-----|------|
| AP | HIKMA FARMACEUTICA | 50MG/ML | N40573 001 | Sep 13, 2006 | Aug | NEWA |
|----|--------------------|---------|------------|--------------|-----|------|

PHYTONADIONE

INJECTABLE; INJECTION

AQUAMEPHYTON

@ MERCK

1MG/0.5ML

N12223 002

Feb DISC

@

10MG/ML

N12223 001

Feb DISC

VITAMIN K1

| | | | | | | | |
|----|---|---------|-----------|------------|--------------|-----|------|
| BP | + | HOSPIRA | 1MG/0.5ML | N87954 001 | Jul 25, 1983 | Feb | CRLD |
|----|---|---------|-----------|------------|--------------|-----|------|

| | | | | | | | |
|--|---|--|---------|------------|--------------|-----|------|
| | + | | 10MG/ML | N87955 001 | Jul 25, 1983 | Feb | CRLD |
|--|---|--|---------|------------|--------------|-----|------|

PILOCARPINE HYDROCHLORIDE

TABLET; ORAL

PILOCARPINE HYDROCHLORIDE

| | | | | | | |
|----|------------|-----|------------|--------------|-----|------|
| AB | IMPAX LABS | 5MG | N77248 001 | Mar 31, 2006 | Mar | NEWA |
|----|------------|-----|------------|--------------|-----|------|

| | | | | | | |
|----|--|-------|------------|--------------|-----|------|
| AB | | 7.5MG | N77248 002 | Mar 31, 2006 | Mar | NEWA |
|----|--|-------|------------|--------------|-----|------|

SALAGEN

| | | | | | | | |
|----|---|----------------|-------|------------|--------------|-----|------|
| AB | + | MGI PHARMA INC | 7.5MG | N20237 002 | Apr 18, 2003 | Mar | CFTG |
|----|---|----------------|-------|------------|--------------|-----|------|

POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL

POLYETHYLENE GLYCOL 3350

| | | | | | | |
|----|----------------|---------------|------------|--------------|-----|------|
| AA | COASTAL PHARMS | 17GM/SCOOPFUL | N77893 001 | May 26, 2006 | May | NEWA |
|----|----------------|---------------|------------|--------------|-----|------|

| | | | | | | |
|----|-----------|---------------|------------|--------------|-----|------|
| AA | KALI LABS | 17GM/SCOOPFUL | N77736 001 | May 26, 2006 | May | NEWA |
|----|-----------|---------------|------------|--------------|-----|------|

| | | | | | | |
|----|-------------|---------------|------------|--------------|-----|------|
| AA | TEVA PHARMS | 17GM/SCOOPFUL | N77445 001 | May 04, 2006 | Apr | NEWA |
|----|-------------|---------------|------------|--------------|-----|------|

| | | | | | | | |
|-----|----|------------|---------------|------------|--------------|-----|------|
| >A> | AA | YVR THERAP | 17GM/SCOOPFUL | N77706 001 | Sep 27, 2006 | Sep | NEWA |
|-----|----|------------|---------------|------------|--------------|-----|------|

>A> POSACONAZOLE

>A> SUSPENSION; ORAL

>A> NOXAFIL

| | | | | | | | |
|-----|---|----------|---------|------------|--------------|-----|------|
| >A> | + | SCHERING | 40MG/ML | N22003 001 | Sep 15, 2006 | Sep | NEWA |
|-----|---|----------|---------|------------|--------------|-----|------|

POTASSIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER

| | | | | | | | |
|----|---|-----------------|-----------|------------|--------------|-----|------|
| AP | + | BAXTER HLTHCARE | 14.9MG/ML | N19904 001 | Dec 26, 1989 | Jun | CTEC |
|----|---|-----------------|-----------|------------|--------------|-----|------|

| | | | | | | | |
|----|---|--|-------------|------------|--------------|-----|------|
| AP | + | | 746MG/100ML | N19904 005 | Dec 17, 1990 | Jun | CTEC |
|----|---|--|-------------|------------|--------------|-----|------|

POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER

| | | | | | | | |
|----|---|-----------------|--------------|------------|--------------|-----|------|
| AP | + | BAXTER HLTHCARE | 1.49GM/100ML | N19904 006 | Dec 17, 1990 | Jun | CTEC |
|----|---|-----------------|--------------|------------|--------------|-----|------|

TABLET, EXTENDED RELEASE; ORAL

KLOR-CON

| | | | | | | |
|----|--------------|------|------------|--------------|-----|------|
| AB | UPSHER SMITH | 8MEQ | N19123 001 | Apr 17, 1986 | Jan | CRLD |
|----|--------------|------|------------|--------------|-----|------|

POTASSIUM CHLORIDE

| | | | | | | | |
|----|---|--------------|------|------------|--------------|-----|------|
| AB | + | COPELY PHARM | 8MEQ | N70618 001 | Sep 09, 1987 | Jan | CRLD |
|----|---|--------------|------|------------|--------------|-----|------|

POTASSIUM CITRATE

TABLET, EXTENDED RELEASE; ORAL

POTASSIUM CITRATE

| | | | | | | |
|----|------------|------|------------|--------------|-----|------|
| AB | COREPHARMA | 5MEQ | N77440 001 | Jun 09, 2006 | May | NEWA |
|----|------------|------|------------|--------------|-----|------|

| | | | | | | |
|----|--|-------|------------|--------------|-----|------|
| AB | | 10MEQ | N77440 002 | Jun 09, 2006 | May | NEWA |
|----|--|-------|------------|--------------|-----|------|

UROCIT-K

| | | | | | | |
|----|----------------|------|------------|--------------|-----|------|
| AB | MISSION PHARMA | 5MEQ | N19071 001 | Aug 30, 1985 | May | CFTG |
|----|----------------|------|------------|--------------|-----|------|

| | | | | | | | |
|----|---|--|-------|------------|--------------|-----|------|
| AB | + | | 10MEQ | N19071 002 | Aug 31, 1992 | May | CFTG |
|----|---|--|-------|------------|--------------|-----|------|

PRAVASTATIN SODIUM

TABLET; ORAL

PRAVACHOL

| | | | | | | |
|--------------------|----------------------|------|------------|--------------|-----|------|
| AB | BRISTOL MYERS SQUIBB | 10MG | N19898 002 | Oct 31, 1991 | Apr | CFTG |
| AB | | 20MG | N19898 003 | Oct 31, 1991 | Apr | CFTG |
| AB | | 40MG | N19898 004 | Mar 22, 1993 | Apr | CFTG |
| PRAVASTATIN SODIUM | | | | | | |
| AB | TEVA | 10MG | N76056 001 | Apr 24, 2006 | Apr | NEWA |
| AB | | 20MG | N76056 002 | Apr 24, 2006 | Apr | NEWA |
| AB | | 40MG | N76056 003 | Apr 24, 2006 | Apr | NEWA |

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

PRAZOSIN HYDROCHLORIDE

@ CLONMEL HLTHCARE

EQ 1MG BASE

N72705 001 May 16, 1989 Jan DISC

@

EQ 5MG BASE

N72707 001 May 16, 1989 Jan DISC

PREDNICARBATE

CREAM; TOPICAL

DERMATOP E EMOLLIENT

| | | | | | | | |
|-----|----|-------------------|------|------------|--------------|-----|------|
| >D> | + | SANOFI AVENTIS US | 0.1% | N20279 001 | Oct 29, 1993 | Sep | CFTG |
| >A> | AB | + | 0.1% | N20279 001 | Oct 29, 1993 | Sep | CFTG |
| >A> | | PREDNICARBATE | | | | | |
| >A> | AB | ALTANA | 0.1% | N77287 001 | Sep 19, 2006 | Sep | NEWA |

PREDNISOLONE ACETATE

INJECTABLE; INJECTION

PREDNISOLONE ACETATE

@ STERIS

25MG/ML

N83398 001

Mar DISC

@

50MG/ML

N83764 001

Mar DISC

SUSPENSION/DROPS; OPHTHALMIC

OMNIPRED

| | | | | | | |
|----|-------|----|------------|--|-----|------|
| AB | ALCON | 1% | N17469 001 | | May | CTNA |
|----|-------|----|------------|--|-----|------|

PREDNISOLONE SODIUM PHOSPHATE

TABLET, ORALLY DISINTEGRATING; ORAL

ORAPRED ODT

BIOMARIN PHARM

EQ 10MG BASE

N21959 001 Jun 01, 2006 Jun NEWA

EQ 15MG BASE

N21959 002 Jun 01, 2006 Jun NEWA

+

EQ 30MG BASE

N21959 003 Jun 01, 2006 Jun NEWA

PRIMIDONE

TABLET; ORAL

PRIMIDONE

| | | | | | | |
|----|-----------|-------|------------|--------------|-----|------|
| AB | WEST WARD | 50MG | N40667 001 | Jul 27, 2006 | Jul | NEWA |
| AB | | 250MG | N40667 002 | Jul 27, 2006 | Jul | NEWA |

PROCHLORPERAZINE

SUPPOSITORY; RECTAL

COMPAZINE

@ GLAXOSMITHKLINE

2.5MG

N11127 003

May DISC

@

5MG

N11127 001

May DISC

PROCHLORPERAZINE EDISYLATE

SYRUP; ORAL

COMPAZINE

| | | | | | |
|-------------------|-----------------|------------|--|-----|------|
| @ GLAXOSMITHKLINE | EQ 5MG BASE/5ML | N11188 001 | | May | DISC |
|-------------------|-----------------|------------|--|-----|------|

PROCHLORPERAZINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

COMPAZINE

| | | | | | |
|-------------------|--------------|------------|--------------|-----|------|
| @ GLAXOSMITHKLINE | EQ 10MG BASE | N21019 001 | Oct 06, 1999 | May | DISC |
|-------------------|--------------|------------|--------------|-----|------|

PROGESTERONE

GEL; VAGINAL

CRINONE

| | | | | | |
|---------------|----|------------|--------------|-----|------|
| COLUMBIA LABS | 4% | N20701 001 | Jul 31, 1997 | May | CAHN |
|---------------|----|------------|--------------|-----|------|

| | | | | | |
|---|----|------------|--------------|-----|------|
| + | 8% | N20701 002 | Jul 31, 1997 | May | CAHN |
|---|----|------------|--------------|-----|------|

PROMETHAZINE HYDROCHLORIDE

SUPPOSITORY; RECTAL

PHENERGAN

| | | | | | |
|--------------------|--------|------------|--|-----|------|
| @ WYETH PHARMS INC | 12.5MG | N10926 002 | | Mar | DISC |
|--------------------|--------|------------|--|-----|------|

| | | | | | |
|---|------|------------|--|-----|------|
| @ | 25MG | N10926 001 | | Mar | DISC |
|---|------|------------|--|-----|------|

PROMETHAZINE HYDROCHLORIDE

| | | | | | | | |
|----|---|--------------|------|------------|--------------|-----|------|
| AB | + | G AND W LABS | 25MG | N40428 001 | Feb 05, 2002 | Mar | CRLD |
|----|---|--------------|------|------------|--------------|-----|------|

PROMETHEGAN

| | | | | | | |
|---|--------------|------|------------|--------------|-----|------|
| + | G AND W LABS | 50MG | N87165 001 | Aug 14, 1987 | Jan | CRLD |
|---|--------------|------|------------|--------------|-----|------|

SYRUP; ORAL

PROMETH PLAIN

| | | | | | |
|------------------------|------------|------------|--|-----|------|
| @ ACTAVIS MID ATLANTIC | 6.25MG/5ML | N85953 001 | | Jul | CAHN |
|------------------------|------------|------------|--|-----|------|

PROMETHAZINE HYDROCHLORIDE

| | | | | | | |
|----|---------|------------|------------|--------------|-----|------|
| AA | VINTAGE | 6.25MG/5ML | N40643 001 | Apr 26, 2006 | Apr | NEWA |
|----|---------|------------|------------|--------------|-----|------|

TABLET; ORAL

PROMETHAZINE HYDROCHLORIDE

| | | | | | | |
|----|--------------|------|------------|--------------|-----|------|
| AB | KVK-TECH INC | 25MG | N40712 001 | Jul 31, 2006 | Jul | NEWA |
|----|--------------|------|------------|--------------|-----|------|

| | | | | | | |
|----|--|------|------------|--------------|-----|------|
| AB | | 50MG | N40713 001 | Jul 31, 2006 | Jul | NEWA |
|----|--|------|------------|--------------|-----|------|

| | | | | | | |
|----|----------------|--------|------------|--------------|-----|------|
| AB | VINTAGE PHARMS | 12.5MG | N40622 001 | Jul 18, 2006 | Jul | NEWA |
|----|----------------|--------|------------|--------------|-----|------|

| | | | | | | |
|----|--|------|------------|--------------|-----|------|
| AB | | 25MG | N40622 002 | Jul 18, 2006 | Jul | NEWA |
|----|--|------|------------|--------------|-----|------|

| | | | | | | |
|----|--|------|------------|--------------|-----|------|
| AB | | 50MG | N40622 003 | Jul 18, 2006 | Jul | NEWA |
|----|--|------|------------|--------------|-----|------|

| | | | | | | |
|----|--|------|------------|--------------|-----|------|
| AB | | 50MG | N40622 003 | Jul 18, 2006 | Jun | NEWA |
|----|--|------|------------|--------------|-----|------|

| | | | | | | |
|----|------------------|--------|------------|--------------|-----|------|
| AB | ZYDUS PHARMS USA | 12.5MG | N40596 001 | Nov 18, 2005 | Jul | CTEC |
|----|------------------|--------|------------|--------------|-----|------|

PROPOFOL

INJECTABLE; INJECTION

DIPRIVAN

| | | | | | | | |
|----|---|--------------------|---------|------------|--------------|-----|------|
| AB | + | ABRAXIS BIOSCIENCE | 10MG/ML | N19627 002 | Jun 11, 1996 | Jul | CAHN |
|----|---|--------------------|---------|------------|--------------|-----|------|

| | | | | | |
|---|---------|------------|--------------|-----|------|
| @ | 10MG/ML | N19627 001 | Oct 02, 1989 | Jul | CAHN |
|---|---------|------------|--------------|-----|------|

PROPOFOL

| | | | | | | |
|----|---------|---------|------------|--------------|-----|------|
| AB | HOSPIRA | 10MG/ML | N77908 001 | Mar 17, 2006 | Mar | NEWA |
|----|---------|---------|------------|--------------|-----|------|

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DOLENE

| | | | | | |
|-----------------|------|------------|--|-----|------|
| @ RADIUS PHARMS | 65MG | N80530 001 | | Aug | CAHN |
|-----------------|------|------------|--|-----|------|

PROPOXYPHENE HYDROCHLORIDE

| | | | | | | |
|----|-----------|------|------------|--|-----|------|
| AA | PAR PHARM | 65MG | N80269 001 | | Mar | CAHN |
|----|-----------|------|------------|--|-----|------|

PROPRANOLOL HYDROCHLORIDE

INJECTABLE; INJECTION
PROPRANOLOL HYDROCHLORIDE

| | | | | | | |
|----|--------|--------|------------|--------------|-----|------|
| AP | SANDOZ | 1MG/ML | N76400 001 | Feb 26, 2003 | Jan | CAHN |
|----|--------|--------|------------|--------------|-----|------|

PROPYLTHIOURACIL

TABLET; ORAL
PROPYLTHIOURACIL

| | | | | | | |
|-----|-----------------------|------|------------|--|-----|------|
| BD | ACTAVIS ELIZABETH | 50MG | N80172 001 | | Jun | CAHN |
| >D> | BD + CLONMEL HLTHCARE | 50MG | N06188 001 | | Sep | CAHN |
| >A> | BD + STADA PHARMS | 50MG | N06188 001 | | Sep | CAHN |

PYRIDOSTIGMINE BROMIDE

INJECTABLE; INJECTION
REGONOL

| | | | | | | |
|----|--------|--------|------------|--|-----|------|
| AP | SANDOZ | 5MG/ML | N17398 001 | | Jan | CAHN |
|----|--------|--------|------------|--|-----|------|

QUAZEPAM

TABLET; ORAL
DORAL
@ QUESTCOR PHARMS
+
15MG

| | | | |
|------------|--------------|-----|------|
| N18708 003 | Feb 26, 1987 | May | CAHN |
| N18708 001 | Dec 27, 1985 | May | CAHN |

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL
QUINAPRIL

| | | | | | | |
|----|-------|--------------|------------|--------------|-----|------|
| AB | LUPIN | EQ 5MG BASE | N77690 001 | Jun 20, 2006 | Jun | NEWA |
| AB | | EQ 10MG BASE | N77690 002 | Jun 20, 2006 | Jun | NEWA |
| AB | | EQ 20MG BASE | N77690 003 | Jun 20, 2006 | Jun | NEWA |
| AB | | EQ 40MG BASE | N77690 004 | Jun 20, 2006 | Jun | NEWA |

QUINAPRIL HYDROCHLORIDE

| | | | | | | |
|----|----------|--------------|------------|--------------|-----|------|
| AB | TORPHARM | EQ 5MG BASE | N76240 001 | Jan 26, 2006 | Jan | NEWA |
| AB | | EQ 10MG BASE | N76240 002 | Jan 26, 2006 | Jan | NEWA |
| AB | | EQ 20MG BASE | N76240 003 | Jan 26, 2006 | Jan | NEWA |
| AB | | EQ 40MG BASE | N76240 004 | Jan 26, 2006 | Jan | NEWA |

QUINIDINE GLUCONATE

TABLET, EXTENDED RELEASE; ORAL
QUINIDINE GLUCONATE

| | | | | | | |
|----|----------------|-------|------------|--------------|-----|------|
| BX | + MUTUAL PHARM | 324MG | N89338 001 | Feb 11, 1987 | Jan | CTEC |
| BX | WATSON LABS | 324MG | N87810 001 | Sep 29, 1982 | Jan | CMFD |

QUINIDINE SULFATE

TABLET; ORAL
QUINIDINE SULFATE
@ CLONMEL HLTHCARE
@ LANNETT
@ MUTUAL PHARM

| | | | | | | |
|----|--------------|-------|------------|--------------|-----|------|
| | | 200MG | N87011 001 | | Jan | DISC |
| | | 200MG | N83743 001 | | Jan | DISC |
| | | 100MG | N81029 001 | Apr 14, 1989 | Jan | DISC |
| AB | | 300MG | N81031 001 | Apr 14, 1989 | Jul | CMFD |
| | @ PHARM FORM | 200MG | N83808 001 | | Jan | DISC |
| | @ SANDOZ | 200MG | N84631 001 | | Jan | DISC |
| | @ | 200MG | N84914 001 | | Jan | DISC |
| AB | | 200MG | N88072 002 | | Jan | NEWA |
| | @ | 300MG | N89839 001 | Sep 29, 1988 | Jan | DISC |

TABLET; ORAL

QUINIDINE SULFATE

| | | | | | |
|----|-------------|-------|------------|-----|------|
| AB | WATSON LABS | 200MG | N83288 001 | Jul | CMFD |
| | @ | 200MG | N83288 001 | Jan | DISC |
| | @ | 200MG | N85140 002 | Jan | DISC |

QUININE SULFATE

CAPSULE; ORAL

QUININE SULFATE

| | | | | | | | |
|-----|---|-------------------|-------|------------|--------------|-----|------|
| >A> | + | AR HOLDING CO INC | 324MG | N21799 001 | Aug 12, 2005 | Sep | CAHN |
| >D> | + | MUTUAL PHARM | 324MG | N21799 001 | Aug 12, 2005 | Sep | CAHN |

RANITIDINE

INJECTABLE; INJECTION

RANITIDINE

| | | | | | | |
|----|---------|-----------------|------------|--------------|-----|------|
| AP | BEDFORD | EQ 25MG BASE/ML | N77458 001 | Feb 16, 2006 | Feb | NEWA |
|----|---------|-----------------|------------|--------------|-----|------|

RANOLAZINE

TABLET, EXTENDED RELEASE; ORAL

RANEXA

| | | | | | | | |
|--|---|-----------|-------|------------|--------------|-----|------|
| | + | CV THERAP | 500MG | N21526 002 | Jan 27, 2006 | Jan | NEWA |
|--|---|-----------|-------|------------|--------------|-----|------|

RASAGILINE MESYLATE

TABLET; ORAL

AZILECT

| | | | | | | | |
|--|---|------|---------------|------------|--------------|-----|------|
| | + | TEVA | EQ 0.5MG BASE | N21641 001 | May 16, 2006 | May | NEWA |
| | | | EQ 1MG BASE | N21641 002 | May 16, 2006 | May | NEWA |

RIBAVIRIN

TABLET; ORAL

RIBAVIRIN

| | | | | | | | |
|-----|----|--------|-------|------------|--------------|-----|------|
| >A> | AB | SANDOZ | 200MG | N77743 001 | Oct 03, 2006 | Sep | NEWA |
|-----|----|--------|-------|------------|--------------|-----|------|

RISPERIDONE

TABLET, ORALLY DISINTEGRATING; ORAL

RISPERDAL

| | | | | | | | |
|--|--|----------------|-----|------------|--------------|-----|------|
| | | JANSSEN PHARMA | 3MG | N21444 004 | Dec 23, 2004 | Mar | CMFD |
| | | | 4MG | N21444 005 | Dec 23, 2004 | Mar | CMFD |

ROPIVACAINE HYDROCHLORIDE MONOHYDRATE

INJECTABLE; INJECTION

NAROPIN

| | | | | | | | |
|--|---|--------------------|----------|------------|--------------|-----|------|
| | | ABRAXIS BIOSCIENCE | 2MG/ML | N20533 001 | Sep 24, 1996 | Jul | CAHN |
| | | | 5MG/ML | N20533 003 | Sep 24, 1996 | Jul | CAHN |
| | | | 7.5MG/ML | N20533 004 | Sep 24, 1996 | Jul | CAHN |
| | + | | 10MG/ML | N20533 005 | Sep 24, 1996 | Jul | CAHN |

SECRETIN SYNTHETIC PORCINE

FOR SOLUTION; INTRAVENOUS

SECFLO

| | | | | | | | |
|--|---|------------|------------|------------|--------------|-----|------|
| | + | CHIRHOCLIN | 16UGM/VIAL | N21136 001 | Apr 04, 2002 | May | CTNA |
|--|---|------------|------------|------------|--------------|-----|------|

SELEGILINE

FILM, EXTENDED RELEASE; TRANSDERMAL
EMSAM

| | | | | | | |
|---|----------|-----------|------------|--------------|-----|------|
| | SOMERSET | 6MG/24HR | N21336 001 | Feb 27, 2006 | Mar | CAIN |
| | | 9MG/24HR | N21336 002 | Feb 27, 2006 | Mar | CAIN |
| + | | 12MG/24HR | N21336 003 | Feb 27, 2006 | Mar | CAIN |

SELEGILINE HYDROCHLORIDE

CAPSULE; ORAL

| | | | | | | |
|-----|--------------------------|---------------|-----|------------|--------------|----------|
| >D> | SELEGILINE HYDROCHLORIDE | | | | | |
| >D> | AB | AAIPHARMA LLC | 5MG | N75145 001 | Sep 15, 2003 | Sep DISC |
| >A> | | @ | 5MG | N75145 001 | Sep 15, 2003 | Sep DISC |

FILM, EXTENDED RELEASE; TRANSDERMAL
EMSAM

| | | | | | | |
|---|----------|-----------|------------|--------------|-----|------|
| | SOMERSET | 6MG/24HR | N21336 001 | Feb 27, 2006 | Feb | NEWA |
| | | 9MG/24HR | N21336 002 | Feb 27, 2006 | Feb | NEWA |
| + | | 12MG/24HR | N21336 003 | Feb 27, 2006 | Feb | NEWA |

TABLET, ORALLY DISINTEGRATING; ORAL
ZELAPAR

| | | | | | | |
|---|--------------------|--------|------------|--------------|-----|------|
| + | VALEANT PHARM INTL | 1.25MG | N21479 001 | Jun 14, 2006 | Jun | NEWA |
|---|--------------------|--------|------------|--------------|-----|------|

SELENIUM SULFIDE

LOTION/SHAMPOO; TOPICAL

SELENIUM SULFIDE

| | | | | | | |
|----|----------------------|------|------------|--|-----|------|
| AT | ACTAVIS MID ATLANTIC | 2.5% | N84394 001 | | Jul | CAHN |
|----|----------------------|------|------------|--|-----|------|

SERTRALINE HYDROCHLORIDE

CONCENTRATE; ORAL

SERTRALINE HYDROCHLORIDE

| | | | | | | |
|----|--------|-----------------|------------|--------------|-----|------|
| AB | ROXANE | EQ 20MG BASE/ML | N76934 001 | Jun 30, 2006 | Jun | NEWA |
|----|--------|-----------------|------------|--------------|-----|------|

ZOLOFT

| | | | | | | |
|----|---|--------|-----------------|------------|--------------|----------|
| AB | + | PFIZER | EQ 20MG BASE/ML | N20990 001 | Dec 07, 1999 | Jun CFTG |
|----|---|--------|-----------------|------------|--------------|----------|

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

| | | | | | | |
|----|-------------|---------------|------------|--------------|-----|------|
| AB | IVAX PHARMS | EQ 25MG BASE | N75719 003 | Jun 30, 2006 | Jun | NEWA |
| AB | | EQ 50MG BASE | N75719 001 | Jun 30, 2006 | Jun | NEWA |
| AB | | EQ 100MG BASE | N75719 002 | Jun 30, 2006 | Jun | NEWA |
| AB | TEVA | EQ 25MG BASE | N76465 001 | Aug 11, 2006 | Aug | NEWA |
| AB | | EQ 50MG BASE | N76465 002 | Aug 11, 2006 | Aug | NEWA |
| AB | | EQ 100MG BASE | N76465 003 | Aug 11, 2006 | Aug | NEWA |

ZOLOFT

| | | | | | | |
|----|--------|---------------|------------|--------------|-----|------|
| AB | PFIZER | EQ 25MG BASE | N19839 005 | Mar 06, 1996 | Jun | CFTG |
| AB | | EQ 50MG BASE | N19839 001 | Dec 30, 1991 | Jun | CFTG |
| AB | + | EQ 100MG BASE | N19839 002 | Dec 30, 1991 | Jun | CFTG |

SIMVASTATIN

TABLET; ORAL

SIMVASTATIN

| | | | | | | |
|----|-------------|------|------------|--------------|-----|------|
| AB | IVAX PHARMS | 5MG | N76052 001 | Jun 23, 2006 | Jun | NEWA |
| AB | | 10MG | N76052 002 | Jun 23, 2006 | Jun | NEWA |
| AB | | 20MG | N76052 003 | Jun 23, 2006 | Jun | NEWA |
| AB | | 40MG | N76052 004 | Jun 23, 2006 | Jun | NEWA |
| AB | RANBAXY | 80MG | N76285 005 | Jun 23, 2006 | Jun | NEWA |

TABLET; ORAL

ZOCOR

| | | | | | | | |
|----|---|-------|------|------------|--------------|-----|------|
| AB | | MERCK | 5MG | N19766 001 | Dec 23, 1991 | Jun | CFTG |
| AB | | | 10MG | N19766 002 | Dec 23, 1991 | Jun | CFTG |
| AB | | | 20MG | N19766 003 | Dec 23, 1991 | Jun | CFTG |
| AB | | | 40MG | N19766 004 | Dec 23, 1991 | Jun | CFTG |
| AB | + | | 80MG | N19766 005 | Jul 10, 1998 | Jun | CFTG |

SODIUM CHLORIDE

INJECTABLE; INJECTION

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | | | | |
|----|---|---------------------|------------------------|------------|--------------|-----|------|
| AP | + | B BRAUN | 900MG/100ML | N17464 001 | | May | CRLD |
| AP | + | | 900MG/100ML | N19635 002 | Mar 09, 1988 | May | CRLD |
| AP | + | BAXTER HLTHCARE | 900MG/100ML | N16677 001 | | May | CRLD |
| AP | + | | 900MG/100ML | N20178 001 | Dec 07, 1992 | May | CRLD |
| AP | + | HOSPIRA | 900MG/100ML | N16366 001 | | May | CRLD |
| AP | + | | 900MG/100ML | N19465 001 | Jul 15, 1985 | May | CRLD |
| AP | + | | 900MG/100ML | N19480 001 | Sep 17, 1985 | May | CRLD |
| | + | MALLINCKRODT | 45MG/50ML (9MG/ML) | N21569 001 | Jul 27, 2006 | Aug | CDFR |
| | | | 112.5MG/125ML (9MG/ML) | N21569 002 | Jul 27, 2006 | Aug | CDFR |
| AP | | TARO PHARMS IRELAND | 9MG/ML | N77407 001 | Aug 11, 2006 | Aug | NEWA |

INJECTABLE; INTRAVASCULAR

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | | | | |
|--|---|--------------|--------------------------|------------|--------------|-----|------|
| | + | MALLINCKRODT | 45MG/50ML (0.9MG/ML) | N21569 001 | Jul 27, 2006 | Jul | NEWA |
| | | | 112.5MG/125ML (0.9MG/ML) | N21569 002 | Jul 27, 2006 | Jul | NEWA |

SODIUM IODIDE, I-123

CAPSULE; ORAL

SODIUM IODIDE I 123

| | | | | | | | |
|-----|----|---------------|--------|------------|--------------|-----|------|
| >D> | AA | GE HEALTHCARE | 100uCi | N17630 001 | | Sep | CRLD |
| >A> | AA | + | 100uCi | N17630 001 | | Sep | CRLD |
| >D> | AA | SYNCOR PHARMS | 100uCi | N18671 001 | May 27, 1982 | Sep | CRLD |
| >A> | AA | + | 100uCi | N18671 001 | May 27, 1982 | Sep | CRLD |
| >D> | AA | | 200uCi | N18671 002 | May 27, 1982 | Sep | CRLD |
| >A> | AA | + | 200uCi | N18671 002 | May 27, 1982 | Sep | CRLD |

SOLUTION; ORAL

SODIUM IODIDE I 123

| | | | | | | | |
|-----|--|---------------|---------|------------|--|-----|------|
| >D> | | GE HEALTHCARE | 2mCi/ML | N17630 002 | | Sep | CRLD |
| >A> | | + | 2mCi/ML | N17630 002 | | Sep | CRLD |

SODIUM IODIDE, I-131

CAPSULE; ORAL

SODIUM IODIDE I 131

| | | | | | | | |
|--|--|-----------|--------|------------|--------------|-----|------|
| | | DRAXIMAGE | 100mCi | N21305 004 | Nov 18, 2004 | Jun | NEWA |
|--|--|-----------|--------|------------|--------------|-----|------|

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL

OSMOPREP

| | | | | | | | |
|--|---|--------------|-----------------|------------|--------------|-----|------|
| | + | SALIX PHARMS | 0.398GM;1.102GM | N21892 001 | Mar 16, 2006 | Mar | NEWA |
|--|---|--------------|-----------------|------------|--------------|-----|------|

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION

GENOTROPIN

| | | | | | | | |
|----|---|----------------------|------------|------------|--------------|-----|------|
| BX | + | PHARMACIA AND UPJOHN | 5.8MG/VIAL | N20280 006 | Aug 24, 1995 | May | CTEC |
|----|---|----------------------|------------|------------|--------------|-----|------|

INJECTABLE; INJECTION

| GENOTROPIN PRESERVATIVE FREE | | | | | | | | | |
|------------------------------|----------------------|-------------|--------|-----|--------------|-----|------|--|--|
| BX | PHARMACIA AND UPJOHN | 1.5MG/VIAL | N20280 | 004 | Aug 24, 1995 | May | CTEC | | |
| NUTROPIN DEPOT | | | | | | | | | |
| | @ GENENTECH | 13.5MG/VIAL | N21075 | 001 | Dec 22, 1999 | Jul | DISC | | |
| | @ | 18MG/VIAL | N21075 | 002 | Dec 22, 1999 | Jul | DISC | | |
| | @ | 22.5MG/VIAL | N21075 | 003 | Dec 22, 1999 | Jul | DISC | | |
| OMNITROPE | | | | | | | | | |
| BX | SANDOZ | 1.5MG/VIAL | N21426 | 002 | May 30, 2006 | May | NEWA | | |
| BX | | 5.8MG/VIAL | N21426 | 001 | May 30, 2006 | May | NEWA | | |

SPIRONOLACTONE

TABLET; ORAL

| SPIRONOLACTONE | | | | | | | | | |
|----------------|-------------------|-------|--------|-----|--------------|-----|------|--|--|
| AB | ACTAVIS ELIZABETH | 25MG | N40353 | 003 | Mar 15, 2006 | Jun | CAHN | | |
| AB | | 50MG | N40353 | 001 | Jul 29, 1999 | Jun | CAHN | | |
| AB | | 100MG | N40353 | 002 | Jul 29, 1999 | Jun | CAHN | | |
| AB | PUREPAC PHARM | 25MG | N40353 | 003 | Mar 15, 2006 | Feb | NEWA | | |
| AB | VINTAGE | 25MG | N40750 | 001 | Aug 29, 2006 | Aug | NEWA | | |
| AB | | 50MG | N40750 | 002 | Aug 29, 2006 | Aug | NEWA | | |
| AB | | 100MG | N40750 | 003 | Aug 29, 2006 | Aug | NEWA | | |

SUCCINYLCHOLINE CHLORIDE

INJECTABLE; INJECTION

| ANECTINE | | | | | | | | | |
|--------------------------|-------------------|------------|--------|-----|--|-----|------|--|--|
| AP | + SANDOZ | 20MG/ML | N08453 | 002 | | Jan | CAHN | | |
| | @ | 50MG/ML | N08453 | 003 | | Jan | CAHN | | |
| | @ | 500MG/VIAL | N08453 | 001 | | Jan | CAHN | | |
| | @ | 1GM/VIAL | N08453 | 004 | | Jan | CAHN | | |
| SUCCINYLCHOLINE CHLORIDE | | | | | | | | | |
| | @ ORGANON USA INC | 20MG/ML | N80997 | 001 | | Jun | DISC | | |

SULCONAZOLE NITRATE

SOLUTION; TOPICAL

| EXELDERM | | | | | | | | | |
|----------|-------------------|----|--------|-----|--------------|-----|------|--|--|
| | + WESTWOOD SQUIBB | 1% | N18738 | 001 | Aug 30, 1985 | Jun | CMFD | | |

SULFACETAMIDE SODIUM

SOLUTION/DROPS; OPHTHALMIC

| SULFACETAMIDE SODIUM | | | | | | | | | |
|----------------------|-------|-----|--------|-----|--------------|-----|------|--|--|
| AT | ALCON | 30% | N89068 | 001 | May 05, 1987 | Apr | CAHN | | |

SULFAMETHOXAZOLE; TRIMETHOPRIM

SUSPENSION; ORAL

| BACTRIM PEDIATRIC | | | | | | | | | |
|---------------------|----------------------|---------------------|--------|-----|--------------|-----|------|--|--|
| AB | MUTUAL PHARM | 200MG/5ML; 40MG/5ML | N17560 | 002 | | Apr | CMFD | | |
| SULFATRIM | | | | | | | | | |
| AB | ACTAVIS MID ATLANTIC | 200MG/5ML; 40MG/5ML | N18615 | 002 | Jan 07, 1983 | Jul | CAHN | | |
| SULFATRIM PEDIATRIC | | | | | | | | | |
| AB | ACTAVIS MID ATLANTIC | 200MG/5ML; 40MG/5ML | N18615 | 001 | Jan 07, 1983 | Jul | CAHN | | |

SULFASALAZINE

TABLET; ORAL

| SULFASALAZINE | | | | | | | | | |
|---------------|-----------------|-------|--------|-----|--|-----|------|--|--|
| | @ RADIUS PHARMS | 500MG | N80197 | 001 | | Aug | CAHN | | |

SULINDAC

TABLET; ORAL

CLINORIL

@ MERCK 150MG N17911 001 Jun DISC

SULINDAC

@ RADIUS PHARMS 150MG N73261 001 Sep 06, 1991 Jul CAHN

@ 200MG N73262 001 Sep 06, 1991 Jul CAHN

SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS

IMITREX

+ GLAXOSMITHKLINE EQ 6MG BASE/0.5ML (12MG/ML) N20080 001 Dec 28, 1992 Feb CDFR

IMITREX STATDOSE

+ GLAXOSMITHKLINE EQ 4MG BASE/0.5ML (8MG/ML) N20080 002 Feb 01, 2006 Feb NEWA

+ EQ 6MG BASE/0.5ML (12MG/ML) N20080 003 Dec 23, 1996 Feb NEWA

SUNITINIB MALATE

CAPSULE; ORAL

SUTENT

CPPI CV

EQ 12.5MG BASE N21938 001 Jan 26, 2006 Jul CAHN

EQ 25MG BASE N21938 002 Jan 26, 2006 Jul CAHN

+ EQ 50MG BASE N21938 003 Jan 26, 2006 Jul CAHN

PFIZER

EQ 12.5MG BASE N21938 001 Jan 26, 2006 Jun CPOT

12.5MG N21938 001 Jan 26, 2006 Jan NEWA

EQ 25MG BASE N21938 002 Jan 26, 2006 Jun CPOT

25MG N21938 002 Jan 26, 2006 Jan NEWA

+ EQ 50MG BASE N21938 003 Jan 26, 2006 Jun CPOT

+ 50MG N21938 003 Jan 26, 2006 Jan NEWA

TAMOXIFEN CITRATE

SOLUTION; ORAL

SOLTAMOX

+ SAVIENT PHARMA EQ 10MG BASE/5ML N21807 001 Oct 29, 2005 Jun CAHN

+ SAVIENT PHARMS EQ 10MG BASE/5ML N21807 001 Oct 29, 2005 Jul CAHN

TECHNETIUM TC-99M APCITIDE

INJECTABLE; INJECTION

ACUTECT

@ CIS BIO INTL SA N/A N20887 001 Sep 14, 1998 May DISC

N/A N20887 001 Sep 14, 1998 Apr CAHN

TECHNETIUM TC-99M DEPREOTIDE

INJECTABLE; INJECTION

NEO TECT KIT

@ CIS BIO INTL SA N/A N21012 001 Aug 03, 1999 May DISC

+ N/A N21012 001 Aug 03, 1999 Apr CAHN

TECHNETIUM TC-99M MEBROFENIN KIT

INJECTABLE; INJECTION

CHOLETEC

+ BRACCO N/A N18963 001 Jan 21, 1987 Aug CRLD

TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE; INJECTION

MYOVIEW 30ML

@ GE HEALTHCARE

N/A

N20372 002 Jul 07, 2005 May DISC

TEMAZEPAM

CAPSULE; ORAL

TEMAZEPAM

AB ACTAVIS ELIZABETH

15MG

N71638 001 Aug 07, 1987 Jun CAHN

AB 30MG

N71620 001 Aug 07, 1987 Jun CAHN

TERBUTALINE SULFATE

INJECTABLE; INJECTION

BRETHINE

@ AAIPHARMA LLC

1MG/ML

N18571 001 Aug DISC

TERBUTALINE SULFATE

AP + BEDFORD

1MG/ML

N76770 001 Apr 23, 2004 Aug CRLD

TERCONAZOLE

CREAM; VAGINAL

TERAZOL 3

>D> @ ORTHO MCNEIL PHARM 0.8%

N19964 001 Feb 21, 1991 Sep CMFD

>A> AB + 0.8%

N19964 001 Feb 21, 1991 Sep CMFD

@ 0.8%

N19964 001 Feb 21, 1991 Jul DISC

TERAZOL 7

>D> @ ORTHO MCNEIL PHARM 0.4%

N19579 001 Dec 31, 1987 Sep CMFD

>A> AB + 0.4%

N19579 001 Dec 31, 1987 Sep CMFD

@ 0.4%

N19579 001 Dec 31, 1987 Jul DISC

TERCONAZOLE

>D> AB + TARO 0.4%

N76043 001 Jan 19, 2005 Sep CRLD

>A> AB 0.4%

N76043 001 Jan 19, 2005 Sep CRLD

AB + 0.4%

N76043 001 Jan 19, 2005 Jul CRLD

>D> BX + 0.8%

N75953 001 Apr 06, 2004 Sep CRLD

>A> AB 0.8%

N75953 001 Apr 06, 2004 Sep CRLD

BX + 0.8%

N75953 001 Apr 06, 2004 Jul CRLD

SUPPOSITORY; VAGINAL

TERAZOL 3

AB + ORTHO MCNEIL PHARM 80MG

N19641 001 May 24, 1988 Mar CFTG

TERCONAZOLE

AB ALTANA 80MG

N76850 001 Jul 12, 2006 Jun NEWA

AB PERRIGO NEW YORK 80MG

N77149 001 Mar 17, 2006 Mar NEWA

TESTOSTERONE

GEL; TRANSDERMAL

ANDROGEL

AB + UNIMED PHARMS 1%

N21015 001 Feb 28, 2000 Jan CTEC

TESTOSTERONE

AB WATSON LABS 1%

N76737 001 Jan 27, 2006 Jan NEWA

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

TESTOSTERONE CYPIONATE

AO SANDOZ 100MG/ML

N40615 001 Aug 10, 2006 Aug NEWA

AO 200MG/ML

N40615 002 Aug 10, 2006 Aug NEWA

TESTOSTERONE ENANTHATE

| | | | | | |
|----|-----------------------|----------|------------|--------------|----------|
| | INJECTABLE; INJECTION | | | | |
| | DELATESTRYL | | | | |
| | @ INDEVUS PHARMS | 200MG/ML | N09165 001 | Jan | CAHN |
| AO | + | 200MG/ML | N09165 003 | Jan | CAHN |
| | TESTOSTERONE ETHANATE | | | | |
| AO | PADDOCK | 200MG/ML | N40575 001 | Jun 14, 2006 | May NEWA |

TETRACYCLINE HYDROCHLORIDE

| | | | | | |
|--|-------------------|-----------|------------|-----|------|
| | CAPSULE; ORAL | | | | |
| | ACHROMYCIN V | | | | |
| | @ RADIUS PHARMS | 250MG | N50278 003 | May | CAHN |
| | @ | 500MG | N50278 001 | May | CAHN |
| | @ SCIREG INTL INC | 250MG | N50278 003 | Apr | CAHN |
| | @ | 500MG | N50278 001 | Apr | CAHN |
| | SUSPENSION; ORAL | | | | |
| | SUMYCIN | | | | |
| | + PAR PHARM | 125MG/5ML | N60400 001 | Aug | CAHN |

THALLOUS CHLORIDE, TL-201

| | | | | | |
|----|--------------------------|---------|------------|--------------|----------|
| | INJECTABLE; INJECTION | | | | |
| | THALLOUS CHLORIDE TL 201 | | | | |
| AP | TRACE RADIOCHEMICALS | 1mCi/ML | N75569 001 | Nov 21, 2001 | Feb CAHN |

THEOPHYLLINE

| | | | | | |
|----|--------------------------------|-----------|------------|--------------|----------|
| | ELIXIR; ORAL | | | | |
| | THEOPHYLLINE | | | | |
| AA | ACTAVIS MID ATLANTIC | 80MG/15ML | N85863 001 | Jul | CAHN |
| | SOLUTION; ORAL | | | | |
| | THEOLAIR | | | | |
| | @ 3M | 80MG/15ML | N86107 001 | Aug | DISC |
| | THEOPHYLLINE | | | | |
| | + ROXANE | 80MG/15ML | N87449 001 | Sep 15, 1983 | Aug CRLD |
| | TABLET, EXTENDED RELEASE; ORAL | | | | |
| | THEOPHYLLINE | | | | |
| AB | NOSTRUM | 400MG | N40595 001 | Apr 21, 2006 | Apr NEWA |
| AB | | 600MG | N40560 002 | Apr 21, 2006 | Apr NEWA |
| | T-PHYL | | | | |
| | @ PURDUE FREDERICK | 200MG | N88253 001 | Aug 17, 1983 | Jun DISC |
| | UNIPHYL | | | | |
| AB | + PURDUE FREDERICK | 400MG | N87571 001 | Sep 01, 1982 | Apr CTEC |
| AB | + | 600MG | N40086 001 | Apr 15, 1996 | Apr CTEC |

THIABENDAZOLE

| | | | | | |
|--|------------------|-----------|------------|-----|------|
| | SUSPENSION; ORAL | | | | |
| | MINTEZOL | | | | |
| | @ MERCK | 500MG/5ML | N16097 001 | Jun | DISC |

THIORIDAZINE HYDROCHLORIDE

| | | | | | |
|----|----------------------------|----------|------------|--------------|----------|
| | CONCENTRATE; ORAL | | | | |
| | THIORIDAZINE HYDROCHLORIDE | | | | |
| AA | ACTAVIS MID ATLANTIC | 100MG/ML | N88229 001 | Aug 23, 1983 | Jul CAHN |

TICARCILLIN DISODIUM

INJECTABLE; INJECTION

TICAR

| | | | | | |
|---|-----------------|-------------------|------------|--------------|----------|
| @ | GLAXOSMITHKLINE | EQ 1GM BASE/VIAL | N50497 001 | May | DISC |
| @ | | EQ 3GM BASE/VIAL | N50497 002 | May | DISC |
| @ | | EQ 20GM BASE/VIAL | N50497 004 | May | DISC |
| @ | | EQ 30GM BASE/VIAL | N50497 005 | Apr 04, 1984 | May DISC |

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLOPIDINE HYDROCHLORIDE

| | | | | | |
|----|-------------------|-------|------------|--------------|----------|
| AB | ACTAVIS ELIZABETH | 250MG | N75253 001 | Aug 20, 1999 | Jun CAHN |
|----|-------------------|-------|------------|--------------|----------|

TIMOLOL MALEATE

TABLET; ORAL

BLOCADREN

| | | | | | |
|---|-------|------|------------|-----|------|
| @ | MERCK | 5MG | N18017 001 | Jun | DISC |
| @ | | 10MG | N18017 002 | Jun | DISC |
| @ | | 20MG | N18017 004 | Jun | DISC |

TIMOLOL MALEATE

| | | | | | | |
|----|---|-------|------|------------|--------------|----------|
| AB | + | MYLAN | 20MG | N72668 001 | Jun 08, 1990 | Jun CRLD |
|----|---|-------|------|------------|--------------|----------|

TINIDAZOLE

TABLET; ORAL

TINDAMAX

| | | | | | |
|---|----------------|-------|------------|--------------|----------|
| | MISSION PHARMA | 250MG | N21618 001 | May 17, 2004 | Jan CAHN |
| + | | 500MG | N21618 002 | May 17, 2004 | Jan CAHN |

TIROFIBAN HYDROCHLORIDE

INJECTABLE; INJECTION

AGGRASTAT

| | | | | | |
|---|----------|-------------------|------------|--------------|----------|
| + | MEDICURE | EQ 0.05MG BASE/ML | N20913 001 | May 14, 1998 | Aug CAHN |
| + | | EQ 0.25MG BASE/ML | N20912 001 | May 14, 1998 | Aug CAHN |

TIZANIDINE HYDROCHLORIDE

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE

| | | | | | |
|----|-------------------|-------------|------------|--------------|----------|
| AB | ACTAVIS ELIZABETH | EQ 2MG BASE | N76283 001 | Jul 12, 2002 | Jun CAHN |
| AB | | EQ 4MG BASE | N76283 002 | Jul 12, 2002 | Jun CAHN |

TOBRAMYCIN

SOLUTION; INHALATION

TOBI

| | | | | | |
|---|-----------------|-----------|------------|--------------|----------|
| + | NOVARTIS PHARMS | 300MG/5ML | N50753 001 | Dec 22, 1997 | Jul CAHN |
|---|-----------------|-----------|------------|--------------|----------|

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

TOBRAMYCIN

| | | | | | | |
|----|---|---------------|-----------------|------------|--------------|----------|
| AP | + | ABRAXIS PHARM | EQ 40MG BASE/ML | N65122 002 | Nov 29, 2002 | Jul CRLD |
|----|---|---------------|-----------------|------------|--------------|----------|

TOLMETIN SODIUM

CAPSULE; ORAL

TOLMETIN SODIUM

| | | | | | |
|----|-------------------|---------------|------------|--------------|----------|
| AB | ACTAVIS ELIZABETH | EQ 400MG BASE | N73308 001 | Jan 24, 1992 | Jun CAHN |
|----|-------------------|---------------|------------|--------------|----------|

TABLET; ORAL

TOLMETIN SODIUM

| | | | | | | |
|----|-------------------|---------------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | EQ 600MG BASE | N73527 001 | Jun 30, 1992 | Jun | CAHN |
|----|-------------------|---------------|------------|--------------|-----|------|

TOPIRAMATE

TABLET; ORAL

TOPAMAX

| | | | | | | | |
|----|---|--------------------|-------|------------|--------------|-----|------|
| AB | + | ORTHO MCNEIL PHARM | 25MG | N20505 004 | Dec 24, 1996 | Aug | CFTG |
| AB | | | 100MG | N20505 001 | Dec 24, 1996 | Aug | CFTG |
| AB | | | 200MG | N20505 002 | Dec 24, 1996 | Aug | CFTG |

TOPIRAMATE

| | | | | | | |
|----|-------|-------|------------|--------------|-----|------|
| AB | MYLAN | 25MG | N76314 001 | Sep 11, 2006 | Aug | NEWA |
| AB | | 100MG | N76314 002 | Sep 11, 2006 | Aug | NEWA |
| AB | | 200MG | N76314 003 | Sep 11, 2006 | Aug | NEWA |

TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE

| | | | | | | |
|----|-------------------|------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 50MG | N75960 001 | Jun 19, 2002 | Jun | CAHN |
| | @ IVAX PHARMS | 50MG | N75963 001 | Jul 03, 2002 | Jan | DISC |

TRANLYCYPROMINE SULFATE

TABLET; ORAL

PARNATE

| | | | | | | | |
|----|---|-----------------|--------------|------------|--------------|-----|------|
| AB | + | GLAXOSMITHKLINE | EQ 10MG BASE | N12342 003 | Aug 16, 1985 | Jun | CFTG |
|----|---|-----------------|--------------|------------|--------------|-----|------|

TRANLYCYPROMINE SULFATE

| | | | | | | |
|----|-----------|--------------|------------|--------------|-----|------|
| AB | KALI LABS | EQ 10MG BASE | N40640 001 | Jun 29, 2006 | Jun | NEWA |
|----|-----------|--------------|------------|--------------|-----|------|

TRAVOPROST

SOLUTION/DROPS; OPHTHALMIC

| | | | | | | | |
|-----|------------|-------|--------|------------|--------------|-----|------|
| >A> | TRAVATAN Z | | | | | | |
| >A> | + | ALCON | 0.004% | N21994 001 | Sep 21, 2006 | Sep | NEWA |

TRAZODONE HYDROCHLORIDE

TABLET; ORAL

DESYREL

@ APOTHECON

@

@

@

@

| | | | | | | | |
|-----|-------------------|-------|------------|--------------|--------------|------|------|
| | | 50MG | N18207 001 | | Aug | DISC | |
| | | 100MG | N18207 002 | | Aug | DISC | |
| | | 150MG | N18207 003 | Mar 25, 1985 | Aug | DISC | |
| | | 300MG | N18207 004 | Nov 07, 1988 | Aug | DISC | |
| AB | ACTAVIS ELIZABETH | 50MG | N71636 001 | Apr 18, 1988 | Jun | CAHN | |
| AB | | 100MG | N71514 001 | Apr 18, 1988 | Jun | CAHN | |
| >D> | AB | BARR | 300MG | N71196 003 | Apr 26, 1999 | Sep | CRLD |
| >A> | AB | + | 300MG | N71196 003 | Apr 26, 1999 | Sep | CRLD |

TRIAMCINOLONE

TABLET; ORAL

ARISTOCORT

| | | | | | | | |
|-----|----|---|----------|-----|------------|-----|------|
| >D> | BP | + | ASTELLAS | 4MG | N11161 007 | Sep | DISC |
| >A> | | | @ | 4MG | N11161 007 | Sep | DISC |

KENACORT

| | | | | | | |
|-----|----|----------------------|-----|------------|-----|------|
| >D> | BP | BRISTOL MYERS SQUIBB | 4MG | N11283 006 | Sep | CRLD |
| >A> | | + | 4MG | N11283 006 | Sep | CRLD |

TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

| | | | | | | | | | |
|-----|----|--------------|--------|--------|-----|--------------|-----|------|--|
| >D> | | ARISTOCORT A | | | | | | | |
| >D> | AT | ASTELLAS | 0.025% | N88818 | 001 | Oct 16, 1984 | Sep | DISC | |
| >A> | | @ | 0.025% | N88818 | 001 | Oct 16, 1984 | Sep | DISC | |
| >D> | AT | | 0.1% | N88819 | 001 | Oct 16, 1984 | Sep | DISC | |
| >A> | | @ | 0.1% | N88819 | 001 | Oct 16, 1984 | Sep | DISC | |
| >D> | AT | | 0.5% | N88820 | 001 | Oct 16, 1984 | Sep | DISC | |
| >A> | | @ | 0.5% | N88820 | 001 | Oct 16, 1984 | Sep | DISC | |

TRIAMCINOLONE ACETONIDE

| | | | | | | | | | |
|----|--|----------------------|--------|--------|-----|--------------|-----|------|--|
| AT | | ACTAVIS MID ATLANTIC | 0.1% | N87798 | 001 | Jun 04, 1982 | Jun | CAHN | |
| AT | | VINTAGE | 0.025% | N40671 | 001 | Jun 09, 2006 | May | NEWA | |
| AT | | | 0.1% | N40671 | 002 | Jun 09, 2006 | May | NEWA | |

OINTMENT; TOPICAL

TRIAMCINOLONE ACETONIDE

| | | | | | | | | | |
|----|--|----------------------|------|--------|-----|--------------|-----|------|--|
| AT | | ACTAVIS MID ATLANTIC | 0.1% | N87799 | 001 | Jun 07, 1982 | Jun | CAHN | |
|----|--|----------------------|------|--------|-----|--------------|-----|------|--|

TRIAMCINOLONE DIACETATE

INJECTABLE; INJECTION

ARISTOCORT

| | | | | | | | | | |
|---|--------|---------|--|--------|-----|--|-----|------|--|
| @ | SANDOZ | 25MG/ML | | N11685 | 003 | | Jan | CAHN | |
| @ | | 40MG/ML | | N12802 | 001 | | Jan | CAHN | |

TRIAMCINOLONE HEXACETONIDE

INJECTABLE; INJECTION

ARISTOSPAN

| | | | | | | | | | |
|---|--------|---------|--|--------|-----|--|-----|------|--|
| + | SANDOZ | 5MG/ML | | N16466 | 001 | | Jan | CAHN | |
| + | | 20MG/ML | | N16466 | 002 | | Jan | CAHN | |

TRICHLORMETHIAZIDE

TABLET; ORAL

TRICHLORMETHIAZIDE

| | | | | | | | | | |
|---|----------------|-----|--|--------|-----|--|-----|------|--|
| @ | TG UNITED LABS | 4MG | | N85568 | 001 | | May | CAHN | |
|---|----------------|-----|--|--------|-----|--|-----|------|--|

TRIFLUOPERAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

STELAZINE

| | | | | | | | | | |
|---|-----------------|----------------|--|--------|-----|--|-----|------|--|
| @ | GLAXOSMITHKLINE | EQ 2MG BASE/ML | | N11552 | 005 | | May | DISC | |
|---|-----------------|----------------|--|--------|-----|--|-----|------|--|

TRIMIPRAMINE MALEATE

CAPSULE; ORAL

SURMONTIL

| | | | | | | | | | |
|----|----------------|---------------|--|--------|-----|--------------|-----|------|--|
| AB | ODYSSEY PHARMS | EQ 25MG BASE | | N16792 | 001 | | Jul | CFTG | |
| AB | | EQ 50MG BASE | | N16792 | 002 | | Jul | CFTG | |
| AB | + | EQ 100MG BASE | | N16792 | 003 | Sep 15, 1982 | Jul | CFTG | |

TRIMIPRAMINE MALEATE

| | | | | | | | | | |
|----|----------------|---------------|--|--------|-----|--------------|-----|------|--|
| AB | ACTAVIS TOTOWA | EQ 25MG BASE | | N77361 | 001 | Aug 02, 2006 | Jul | NEWA | |
| AB | | EQ 50MG BASE | | N77361 | 002 | Aug 02, 2006 | Jul | NEWA | |
| AB | | EQ 100MG BASE | | N77361 | 003 | Aug 02, 2006 | Jul | NEWA | |

TRIPLENNAMINE HYDROCHLORIDE

TABLET; ORAL

PBZ

| | | | | | | | | | |
|---|----------|------|--|--------|-----|--|-----|------|--|
| @ | NOVARTIS | 50MG | | N05914 | 002 | | Jan | DISC | |
|---|----------|------|--|--------|-----|--|-----|------|--|

TROLEANDOMYCIN

CAPSULE; ORAL

TAO

@ PFIZER

EQ 250MG BASE

N50336 002

Mar DISC

UNOPROSTONE ISOPROPYL

SOLUTION/DROPS; OPHTHALMIC

RESCULA

@ R TECH UENO LTD

0.15%

N21214 001 Aug 03, 2000 Jun DISC

+

0.15%

N21214 001 Aug 03, 2000 Feb CAHN

UREA

INJECTABLE; INJECTION

UREAPHIL

@ HOSPIRA

40GM/VIAL

N12154 001

Jun DISC

UROKINASE

INJECTABLE; INJECTION

ABBOKINASE

@ IMARX THERAP

5,000 IU/VIAL

N21846 003

Apr CAHN

@

9,000 IU/VIAL

N21846 002

Apr CAHN

+

250,000 IU/VIAL

N21846 001

Apr CAHN

URSODIOL

CAPSULE; ORAL

URSODIOL

AB

COREPHARMA

300MG

N77895 001 Jul 27, 2006 Jul NEWA

VALRUBICIN

SOLUTION; INTRAVESICAL

VALSTAR PRESERVATIVE FREE

+ VALERA

40MG/ML

N20892 001 Sep 25, 1998 Jul CAHN

VARENICLINE TARTRATE

TABLET; ORAL

CHANTIX

>D>

CPPI CV

EQ 0.5MG BASE

N21928 001 May 10, 2006 Sep CAHN

EQ 0.5MG BASE

N21928 001 May 10, 2006 Jul CAHN

>D>

+

EQ 1MG BASE

N21928 002 May 10, 2006 Sep CAHN

+

EQ 1MG BASE

N21928 002 May 10, 2006 Jul CAHN

PFIZER

EQ 0.5MG BASE

N21928 001 May 10, 2006 May NEWA

+

EQ 1MG BASE

N21928 002 May 10, 2006 May NEWA

>A>

PFIZER INC

EQ 0.5MG BASE

N21928 001 May 10, 2006 Sep CAHN

>A>

+

EQ 1MG BASE

N21928 002 May 10, 2006 Sep CAHN

VECURONIUM BROMIDE

INJECTABLE; INJECTION

VECURONIUM BROMIDE

AP

+

BEDFORD

20MG/VIAL

N75549 002 Jun 13, 2000 Jan CRLD

VENLAFAXINE HYDROCHLORIDE

TABLET; ORAL

EFFEXOR

AB

WYETH PHARMS INC

EQ 25MG BASE

N20151 002 Dec 28, 1993 Jul CFTG

TABLET; ORAL

EFFEXOR

| | | | | | | |
|----|------------------|----------------|------------|--------------|-----|------|
| AB | WYETH PHARMS INC | EQ 37.5MG BASE | N20151 006 | Dec 28, 1993 | Jul | CFTG |
| AB | + | EQ 50MG BASE | N20151 003 | Dec 28, 1993 | Jul | CFTG |
| AB | | EQ 75MG BASE | N20151 004 | Dec 28, 1993 | Jul | CFTG |
| AB | | EQ 100MG BASE | N20151 005 | Dec 28, 1993 | Jul | CFTG |

VENLAFAXINE HYDROCHLORIDE

| | | | | | | |
|----|------|----------------|------------|--------------|-----|------|
| AB | TEVA | EQ 25MG BASE | N76690 001 | Aug 03, 2006 | Jul | CTNA |
| AB | | EQ 37.5MG BASE | N76690 002 | Aug 03, 2006 | Jul | NEWA |
| AB | | EQ 50MG BASE | N76690 003 | Aug 03, 2006 | Jul | NEWA |
| AB | | EQ 75MG BASE | N76690 004 | Aug 03, 2006 | Jul | CTNA |
| AB | | EQ 100MG BASE | N76690 005 | Aug 03, 2006 | Jul | NEWA |

VERAPAMIL HYDROCHLORIDE

TABLET; ORAL

VERAPAMIL HYDROCHLORIDE

| | | | | | | |
|----|-------------------|-------|------------|--------------|-----|------|
| AB | ACTAVIS ELIZABETH | 80MG | N71019 001 | Sep 24, 1986 | Jun | CAHN |
| AB | | 120MG | N70468 001 | Sep 24, 1986 | Jun | CAHN |
| | @ RADIUS PHARMS | 80MG | N71880 001 | Apr 05, 1988 | Jul | CAHN |
| | @ | 120MG | N71881 001 | Apr 05, 1988 | Jul | CAHN |

WARFARIN SODIUM

TABLET; ORAL

WARFARIN SODIUM

| | | | | | | |
|----|------------------|-------|------------|--------------|-----|------|
| AB | PLIVA | 1MG | N40616 009 | Jul 05, 2006 | Jun | NEWA |
| AB | | 2MG | N40616 001 | Jul 05, 2006 | Jun | NEWA |
| AB | | 2.5MG | N40616 002 | Jul 05, 2006 | Jun | NEWA |
| AB | | 3MG | N40616 003 | Jul 05, 2006 | Jun | NEWA |
| AB | | 4MG | N40616 004 | Jul 05, 2006 | Jun | NEWA |
| AB | | 5MG | N40616 005 | Jul 05, 2006 | Jun | NEWA |
| AB | | 6MG | N40616 006 | Jul 05, 2006 | Jun | NEWA |
| AB | | 7.5MG | N40616 007 | Jul 05, 2006 | Jun | NEWA |
| AB | | 10MG | N40616 008 | Jul 05, 2006 | Jun | NEWA |
| AB | ZYDUS PHARMS USA | 1MG | N40663 001 | May 30, 2006 | May | NEWA |
| AB | | 2MG | N40663 002 | May 30, 2006 | May | NEWA |
| AB | | 2.5MG | N40663 003 | May 30, 2006 | May | NEWA |
| AB | | 3MG | N40663 004 | May 30, 2006 | May | NEWA |
| AB | | 4MG | N40663 005 | May 30, 2006 | May | NEWA |
| AB | | 5MG | N40663 006 | May 30, 2006 | May | NEWA |
| AB | | 6MG | N40663 007 | May 30, 2006 | May | NEWA |
| AB | | 7.5MG | N40663 008 | May 30, 2006 | May | NEWA |
| AB | | 10MG | N40663 009 | May 30, 2006 | May | NEWA |

WATER FOR INJECTION, STERILE

LIQUID; N/A

STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

| | | | | | | |
|----|---------------------|------|------------|--------------|-----|------|
| AP | TARO PHARMS IRELAND | 100% | N77393 001 | Aug 11, 2006 | Aug | NEWA |
|----|---------------------|------|------------|--------------|-----|------|

ZIDOVUDINE

CAPSULE; ORAL

RETROVIR

| | | | | | | | |
|----|---|----------------------|-------|------------|--------------|-----|------|
| AB | + | GLAXOSMITHKLINE | 100MG | N19655 001 | Mar 19, 1987 | Mar | CFTG |
| | | ZIDOVUDINE | | | | | |
| AB | | AUROBINDO PHARMA LTD | 100MG | N78128 001 | Mar 27, 2006 | Mar | NEWA |

ZIPRASIDONE HYDROCHLORIDE

SUSPENSION; ORAL

GEODON

| | | | | | | | |
|---|------------|-----------------|--------|-----|--------------|-----|------|
| + | PFIZER INC | EQ 10MG BASE/ML | N21483 | 001 | Mar 29, 2006 | Mar | NEWA |
|---|------------|-----------------|--------|-----|--------------|-----|------|

ZONISAMIDE

CAPSULE; ORAL

ZONISAMIDE

| | | | | | | | |
|-----|---------------------|-------|--------|-----|--------------|-----|------|
| AB | BANNER PHARMACAPS | 25MG | N77813 | 001 | Aug 16, 2006 | Aug | NEWA |
| AB | | 50MG | N77813 | 002 | Aug 16, 2006 | Aug | NEWA |
| AB | | 100MG | N77813 | 003 | Aug 16, 2006 | Aug | NEWA |
| >A> | DR REDDYS LABS LTD | 25MG | N77891 | 001 | Sep 29, 2006 | Sep | NEWA |
| >A> | | 50MG | N77891 | 002 | Sep 29, 2006 | Sep | NEWA |
| AB | GLENMARK PHARMS | 25MG | N77651 | 001 | Jan 30, 2006 | Jan | NEWA |
| AB | | 50MG | N77651 | 002 | Jan 30, 2006 | Jan | NEWA |
| AB | | 100MG | N77651 | 003 | Jan 30, 2006 | Jan | NEWA |
| AB | INVAGEN PHARMS | 25MG | N77869 | 001 | May 31, 2006 | May | NEWA |
| AB | | 50MG | N77869 | 002 | May 31, 2006 | May | NEWA |
| AB | | 100MG | N77869 | 003 | May 31, 2006 | May | NEWA |
| AB | SUN PHARM INDS (IN) | 25MG | N77634 | 001 | Mar 17, 2006 | Mar | NEWA |
| AB | | 50MG | N77634 | 002 | Mar 17, 2006 | Mar | NEWA |
| AB | | 100MG | N77634 | 003 | Mar 17, 2006 | Mar | NEWA |
| AB | WATSON LABS | 25MG | N77650 | 001 | Apr 20, 2006 | Apr | NEWA |
| AB | | 50MG | N77650 | 002 | Apr 20, 2006 | Apr | NEWA |
| AB | | 100MG | N77650 | 003 | Apr 20, 2006 | Apr | NEWA |
| AB | WOCKHARDT | 25MG | N77636 | 003 | Jul 27, 2006 | Jul | NEWA |
| AB | | 50MG | N77636 | 002 | Jul 27, 2006 | Jul | NEWA |

OTC DRUG PRODUCT LIST - 26TH EDITION

OTC DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 9 - September 2006

2-1

ACETAMINOPHEN

SUPPOSITORY; RECTAL

ACETAMINOPHEN

| | | | | | |
|--------------------------------|-------------|------------|--------------|--------------|----------|
| ACTAVIS MID ATLANTIC | 120MG | N18337 003 | Sep 12, 1983 | Jun | CAHN |
| | 325MG | N18337 002 | | Jun | CAHN |
| + | 650MG | N18337 001 | | Jun | CAHN |
| INFANTS' FEVERALL | | | | | |
| ACTAVIS MID ATLANTIC | 80MG | N18337 004 | Aug 26, 1992 | Jun | CAHN |
| TYLENOL | | | | | |
| @ MCNEIL CONS | 120MG | N17756 002 | | Jun | CAHN |
| @ | 650MG | N17756 001 | | Jun | CAHN |
| TABLET, EXTENDED RELEASE; ORAL | | | | | |
| TYLENOL (CAPLET) | | | | | |
| + | MCNEIL CONS | 650MG | N19872 001 | Jun 08, 1994 | Jun CAHN |
| TYLENOL (GELTAB) | | | | | |
| + | MCNEIL CONS | 650MG | N19872 002 | Jan 11, 2001 | Jun CAHN |

AVOBENZONE; ECAMSULE; OCTOCRYLENE

CREAM; TOPICAL

ANTHELIOS SX

| | | | | | |
|---|------------|-----------|------------|--------------|----------|
| + | LOREAL USA | 2%;2%;10% | N21502 001 | Jul 21, 2006 | Jul NEWA |
|---|------------|-----------|------------|--------------|----------|

CHLORHEXIDINE GLUCONATE

CLOTH; TOPICAL

HALO

| | | | | | |
|---|------------|----|------------|--------------|----------|
| + | SAGE PRODS | 2% | N21669 001 | Apr 25, 2005 | Aug CTNA |
|---|------------|----|------------|--------------|----------|

SPONGE; TOPICAL

CHLORHEXIDINE GLUCONATE

| | | | | | |
|--|------------------|----|------------|--------------|----------|
| | BECTON DICKINSON | 4% | N72525 001 | Oct 24, 1989 | Aug CAHN |
| | @ KENDALL IL | 4% | N19490 001 | Mar 27, 1987 | Aug DISC |

CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL

SPONGE; TOPICAL

CHLORAPREP WITH TINT

| | | | | | |
|---|---------------|-----------------|------------|--------------|----------|
| + | MEDI FLEX INC | 2%;70% (10.5ML) | N20832 005 | Apr 03, 2006 | Apr NEWA |
|---|---------------|-----------------|------------|--------------|----------|

CLOTRIMAZOLE

CREAM; VAGINAL

CLOTRIMAZOLE

| | | | | | |
|--|----------------------|----|------------|--------------|----------|
| | ACTAVIS MID ATLANTIC | 1% | N74165 001 | Jul 16, 1993 | Jun CAHN |
|--|----------------------|----|------------|--------------|----------|

CROMOLYN SODIUM

SPRAY, METERED; NASAL

CROMOLYN SODIUM

| | | | | | |
|-----------|----------------------|-------------|------------|--------------|----------|
| | ACTAVIS MID ATLANTIC | 5.2MG/SPRAY | N74800 001 | Jul 26, 2001 | Jul CAHN |
| | ALPHARMA US PHARMS | 5.2MG/SPRAY | N74800 001 | Jul 26, 2001 | Jan CPOT |
| + | BAUSCH AND LOMB | 5.2MG/SPRAY | N75702 001 | Jul 03, 2001 | Jan CRLD |
| NASALCROM | | | | | |
| | @ PHARMACIA UPJOHN | 5.2MG/SPRAY | N20463 001 | Jan 03, 1997 | Jan DISC |

DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL

MUCINEX DM

| | | | | | |
|-------------------|------------|------------|--------------|-----|------|
| ADAMS RESP THERAP | 30MG;600MG | N21620 002 | Apr 29, 2004 | May | CAHN |
| + | 60MG;1.2GM | N21620 001 | Apr 29, 2004 | May | CAHN |

DEXTROMETHORPHAN POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

DELSYM

| | | | | | |
|-------------------|-----------------|------------|--------------|-----|------|
| ADAMS RESP THERAP | EQ 30MG HBR/5ML | N18658 001 | Oct 08, 1982 | Jun | CAHN |
|-------------------|-----------------|------------|--------------|-----|------|

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

| | | | | | | |
|-----|--------------------|------|------------|--------------|-----|------|
| >A> | DR REDDYS LABS LTD | 20MG | N77367 001 | Sep 25, 2006 | Sep | NEWA |
| >A> | PERRIGO | 20MG | N77351 001 | Sep 25, 2006 | Sep | NEWA |

GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL

HUMABID

| | | | | | |
|-------------------|-------|------------|--------------|-----|------|
| ADAMS RESP THERAP | 1.2GM | N21282 002 | Dec 18, 2002 | Aug | CTNA |
|-------------------|-------|------------|--------------|-----|------|

MUCINEX

| | | | | | |
|-------------------|-------|------------|--------------|-----|------|
| ADAMS RESP THERAP | 600MG | N21282 001 | Jul 12, 2002 | May | CAHN |
| + | 1.2GM | N21282 002 | Dec 18, 2002 | May | CAHN |

GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

MUCINEX D

| | | | | | |
|-------------------|-------------|------------|--------------|-----|------|
| ADAMS RESP THERAP | 600MG;60MG | N21585 001 | Jun 22, 2004 | May | CAHN |
| + | 1.2GM;120MG | N21585 002 | Jun 22, 2004 | May | CAHN |

IBUPROFEN

SUSPENSION/DROPS; ORAL

CHILDREN'S MOTRIN

| | | | | | |
|-------------|---------|------------|--------------|-----|------|
| MCNEIL CONS | 40MG/ML | N20603 001 | Jun 10, 1996 | Jun | CAHN |
|-------------|---------|------------|--------------|-----|------|

SUSPENSION; ORAL

CHILDREN'S ELIXSURE

| | | | | | |
|------------------|-----------|------------|--------------|-----|------|
| ALTERNA-TCHP LLC | 100MG/5ML | N21604 001 | Jan 07, 2004 | Jul | CAHN |
|------------------|-----------|------------|--------------|-----|------|

IBUPROFEN

| | | | | | |
|----------------------|-----------|------------|--------------|-----|------|
| ACTAVIS MID ATLANTIC | 100MG/5ML | N74916 001 | Apr 30, 1999 | Jul | CAHN |
|----------------------|-----------|------------|--------------|-----|------|

TABLET, CHEWABLE; ORAL

CHILDREN'S MOTRIN

| | | | | | |
|-------------|------|------------|--------------|-----|------|
| MCNEIL CONS | 50MG | N20601 001 | Nov 15, 1996 | Jun | CAHN |
|-------------|------|------------|--------------|-----|------|

JUNIOR STRENGTH MOTRIN

| | | | | | |
|-------------|-------|------------|--------------|-----|------|
| MCNEIL CONS | 100MG | N20601 003 | Nov 15, 1996 | Jun | CAHN |
|-------------|-------|------------|--------------|-----|------|

TABLET; ORAL

JUNIOR STRENGTH MOTRIN

| | | | | | |
|-------------|-------|------------|--------------|-----|------|
| MCNEIL CONS | 100MG | N20602 001 | Jun 10, 1996 | Jun | CAHN |
|-------------|-------|------------|--------------|-----|------|

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

SUSPENSION; ORAL

CHILDREN'S MOTRIN COLD

| | | | | | |
|-------------|--------------------|------------|--------------|-----|------|
| MCNEIL CONS | 100MG/5ML;15MG/5ML | N21128 001 | Aug 01, 2000 | Jun | CAHN |
|-------------|--------------------|------------|--------------|-----|------|

TABLET; ORAL

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

DR REDDYS LABS LTD 200MG;30MG N77628 001 Aug 14, 2006 Aug NEWA

SINE-AID IB

MCNEIL CONS 200MG;30MG N19899 001 Dec 31, 1992 Jun CAHN

INSULIN PURIFIED PORK

INJECTABLE; INJECTION

REGULAR ILETIN II (PORK)

@ LILLY 100 UNITS/ML N18344 001 Jun DISC

INSULIN SUSP ISOPHANE BEEF/PORK

INJECTABLE; INJECTION

NPH ILETIN I (BEEF-PORK)

@ LILLY 100 UNITS/ML N17936 002 Jun DISC

INSULIN SUSP ISOPHANE PURIFIED PORK

INJECTABLE; INJECTION

NPH ILETIN II (PORK)

@ LILLY 100 UNITS/ML N18345 001 Jun DISC

INSULIN ZINC SUSP EXTENDED RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN U

@ LILLY 100 UNITS/ML N19571 002 Jun 10, 1987 Jun DISC

INSULIN ZINC SUSP PURIFIED PORK

INJECTABLE; INJECTION

LENTE ILETIN II (PORK)

@ LILLY 100 UNITS/ML N18347 001 Jun DISC

>A> IODINE POVACRYLEX; ISOPROPYL ALCOHOL

>A> SPONGE; TOPICAL

>A> DURAPREP

>A> 3M EQ 0.7% IODINE;74% (26ML) N21586 002 Sep 29, 2006 Sep NEWA

>A> + EQ 0.7% IODINE; 74% (6ML) N21586 001 Sep 29, 2006 Sep NEWA

KETOCONAZOLE

SHAMPOO; TOPICAL

NIZORAL A-D

+ MCNEIL CONS 1% N20310 001 Oct 10, 1997 Jun CAHN

KETOPROFEN

TABLET; ORAL

ACTRON

@ BAYER 12.5MG N20499 001 Oct 06, 1995 Feb DISC

ORUDIS KT

@ WYETH CONS 12.5MG N20429 001 Oct 06, 1995 Feb DISC

LEVONORGESTREL

TABLET; ORAL

PLAN B

+ DURAMED 0.75MG N21045 002 Aug 24, 2006 Aug NEWA

LOPERAMIDE HYDROCHLORIDE

SOLUTION; ORAL

IMODIUM A-D

| | | | | | |
|---------------|---------|------------|--------------|-----|------|
| + MCNEIL CONS | 1MG/5ML | N19487 001 | Mar 01, 1988 | Jun | CAHN |
|---------------|---------|------------|--------------|-----|------|

SUSPENSION; ORAL

IMODIUM A-D

| | | | | | |
|----------|-----------|------------|--------------|-----|------|
| + MCNEIL | 1MG/7.5ML | N19487 002 | Jul 08, 2004 | Mar | CDFR |
|----------|-----------|------------|--------------|-----|------|

| | | | | | |
|---------------|-----------|------------|--------------|-----|------|
| + MCNEIL CONS | 1MG/7.5ML | N19487 002 | Jul 08, 2004 | Jun | CAHN |
|---------------|-----------|------------|--------------|-----|------|

TABLET; ORAL

IMODIUM A-D

| | | | | | |
|---------------|-----|------------|--------------|-----|------|
| + MCNEIL CONS | 2MG | N19860 001 | Nov 22, 1989 | Jun | CAHN |
|---------------|-----|------------|--------------|-----|------|

LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET; ORAL

IMODIUM ADVANCED

| | | | | | |
|---------------|-----------|------------|--------------|-----|------|
| + MCNEIL CONS | 2MG;125MG | N21140 001 | Nov 30, 2000 | Jun | CAHN |
|---------------|-----------|------------|--------------|-----|------|

LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE

| | | | | | |
|---------|-----------|------------|--------------|-----|------|
| RANBAXY | 2MG;125MG | N77500 001 | Sep 06, 2006 | Aug | NEWA |
|---------|-----------|------------|--------------|-----|------|

LORATADINE

SYRUP; ORAL

LORATADINE

SILARX 1MG/ML

| | | | |
|------------|--------------|-----|------|
| N77421 001 | Jun 29, 2006 | Jun | NEWA |
|------------|--------------|-----|------|

TABLET, CHEWABLE; ORAL

CHILDREN'S CLARITIN

| | | | | | |
|-------------------|-----|------------|--------------|-----|------|
| + SCHERING PLOUGH | 5MG | N21891 001 | Aug 23, 2006 | Aug | NEWA |
|-------------------|-----|------------|--------------|-----|------|

TABLET; ORAL

LORATADINE

APOTEX 10MG

| | | | |
|------------|--------------|-----|------|
| N76471 001 | Feb 14, 2006 | Jan | NEWA |
|------------|--------------|-----|------|

MICONAZOLE NITRATE

CREAM, SUPPOSITORY; TOPICAL, VAGINAL

M-ZOLE 3 COMBINATION PACK

ACTAVIS MID ATLANTIC 2%,200MG

| | | | |
|------------|--------------|-----|------|
| N74926 001 | Apr 16, 1999 | Jun | CAHN |
|------------|--------------|-----|------|

2%,200MG

| | | | |
|------------|--------------|-----|------|
| N74926 001 | Apr 16, 1999 | Jun | CAHN |
|------------|--------------|-----|------|

2%,200MG

| | | | |
|------------|--------------|-----|------|
| N74926 001 | Apr 16, 1999 | Jun | CAHN |
|------------|--------------|-----|------|

2%,200MG

| | | | |
|------------|--------------|-----|------|
| N74926 001 | Apr 16, 1999 | Jun | CAHN |
|------------|--------------|-----|------|

2%,200MG

| | | | |
|------------|--------------|-----|------|
| N74926 001 | Apr 16, 1999 | Jun | CAHN |
|------------|--------------|-----|------|

2%,200MG

| | | | |
|------------|--------------|-----|------|
| N74926 001 | Apr 16, 1999 | Jun | CAHN |
|------------|--------------|-----|------|

2%,200MG

| | | | |
|------------|--------------|-----|------|
| N74926 001 | Apr 16, 1999 | Jun | CAHN |
|------------|--------------|-----|------|

2%,200MG

| | | | |
|------------|--------------|-----|------|
| N74926 001 | Apr 16, 1999 | Jun | CAHN |
|------------|--------------|-----|------|

M-ZOLE 7 DUAL PACK

ACTAVIS MID ATLANTIC 2%,100MG

| | | | |
|------------|--------------|-----|------|
| N74586 001 | Jul 17, 1997 | Jun | CAHN |
|------------|--------------|-----|------|

CREAM; VAGINAL

MICONAZOLE 7

ACTAVIS MID ATLANTIC 2%

| | | | |
|------------|--------------|-----|------|
| N74164 001 | Mar 29, 1996 | Jun | CAHN |
|------------|--------------|-----|------|

SUPPOSITORY; VAGINAL

MICONAZOLE NITRATE

ACTAVIS MID ATLANTIC 100MG

| | | | |
|------------|--------------|-----|------|
| N73507 001 | Nov 19, 1993 | Jun | CAHN |
|------------|--------------|-----|------|

MINOXIDIL

AEROSOL, FOAM; TOPICAL

MEN'S ROGAINE

| | | | | | |
|------------------------|----|------------|--------------|-----|------|
| + PHARMACIA AND UPJOHN | 5% | N21812 001 | Jan 20, 2006 | Jan | NEWA |
|------------------------|----|------------|--------------|-----|------|

SOLUTION; TOPICAL

MINOXIDIL (FOR MEN)

ACTAVIS MID ATLANTIC 2%

N74588 001 Apr 05, 1996 Jul CAHN

MINOXIDIL EXTRA STRENGTH (FOR MEN)

ACTAVIS MID ATLANTIC 5%

N75518 001 Nov 17, 2000 Jul CAHN

NAPROXEN SODIUM

CAPSULE; ORAL

NAPROXEN SODIUM

| | | | | | | |
|-----|---------------------|---------------|------------|--------------|-----|------|
| >D> | @ BANNER PHARMACAPS | EQ 200MG BASE | N21920 001 | Feb 17, 2006 | Sep | CMFD |
| >D> | @ | EQ 200MG BASE | N21920 001 | Feb 17, 2006 | Sep | CMFD |
| >A> | | EQ 200MG BASE | N21920 001 | Feb 17, 2006 | Sep | CMFD |
| | @ | EQ 200MG BASE | N21920 001 | Feb 17, 2006 | Jul | DISC |
| | + | EQ 200MG BASE | N21920 001 | Feb 17, 2006 | Feb | NEWA |

NAPROXEN SODIUM; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE

| | | | | | | |
|-----|--------------------|---------------------|------------|--------------|-----|------|
| >A> | DR REDDYS LABS INC | EQ 220MG BASE;120MG | N77381 001 | Sep 27, 2006 | Sep | NEWA |
|-----|--------------------|---------------------|------------|--------------|-----|------|

NICOTINE POLACRILEX

GUM, CHEWING; BUCCAL

NICORETTE

GLAXOSMITHKLINE

EQ 2MG BASE

N18612 003 Dec 23, 1998 Mar CRLD

EQ 2MG BASE

N18612 004 Sep 25, 2000 Mar CRLD

EQ 4MG BASE

N20066 003 Dec 23, 1998 Mar CRLD

EQ 4MG BASE

N20066 004 Sep 25, 2000 Mar CRLD

NICORETTE (MINT)

GLAXOSMITHKLINE

EQ 2MG BASE

N18612 003 Dec 23, 1998 Apr CTNA

EQ 4MG BASE

N20066 003 Dec 23, 1998 Apr CTNA

NICOTINE POLACRILEX

PERRIGO

EQ 2MG BASE

N76776 001 Sep 16, 2004 Apr CTNA

EQ 2MG BASE

N76777 001 Sep 16, 2004 Apr CTNA

EQ 4MG BASE

N76778 001 Sep 16, 2004 Apr CTNA

EQ 4MG BASE

N76779 001 Sep 16, 2004 Apr CTNA

WATSON LABS

EQ 2MG BASE

N76569 001 Jul 29, 2004 Apr CTNA

EQ 4MG BASE

N76568 002 Jul 29, 2004 Apr CTNA

TROCHE/LOZENGE; ORAL

NICOTINE POLACRILEX

PERRIGO R AND D

EQ 2MG BASE

N77007 001 Jan 31, 2006 Jan NEWA

EQ 4MG BASE

N77007 002 Jan 31, 2006 Jan NEWA

PERMETHRIN

LOTION; TOPICAL

PERMETHRIN

ACTAVIS MID ATLANTIC 1%

N75014 001 Mar 28, 2000 Jun CAHN

POVIDONE-IODINE

SOLUTION; TOPICAL

E-Z PREP

@ CLINIPAD

10%

N19382 001 Jul 25, 1989 Aug DISC

SPONGE; TOPICAL

E-Z PREP

@ CLINIPAD

5%

N19382 002 Jul 25, 1989 Aug DISC

E-Z PREP 220

@ CLINIPAD

5%

N19382 003 Jul 25, 1989 Aug DISC

RANITIDINE HYDROCHLORIDE

TABLET; ORAL

RANITIDINE

WOCKHARDT

EQ 75MG BASE

N76760 001 Feb 24, 2006 Feb NEWA

TERBINAFINE

GEL; TOPICAL

LAMISIL AT

+ NOVARTIS

1%

N21958 001 Jul 24, 2006 Jul NEWA

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

CUMULATIVE SUPPLEMENT NUMBER 09 SEPTEMBER 2006

NO SEPTEMBER 2006 APPROVALS

ORPHAN PRODUCT DESIGNATIONS AND APPROVALS LIST

The list of List of Orphan Designations and Approvals is available at:

<http://www.fda.gov/orphan/designat/list.htm>

**DRUG PRODUCTS WHICH MUST DEMONSTRATE *IN VIVO* BIOAVAILABILITY
ONLY IF PRODUCT FAILS TO ACHIEVE ADEQUATE DISSOLUTION**

NO SEPTEMBER 2006 ADDITIONS

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---|-----------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>ABACAVIR SULFATE; LAMIVUDINE - EPZICOM</u> | | | | | |
| 021652 001 | 5034394 | Dec 18, 2011 | DS DP | D-40 | Aug 02, 2007 |
| | 5034394*PED | Jun 18, 2012 | | | |
| | 5047407 | Nov 17, 2009 | DS DP | U-257 | |
| | 5047407*PED | May 17, 2010 | | | |
| | 5089500 | Jun 26, 2009 | | U-257 | |
| | 5089500*PED | Dec 26, 2009 | | | |
| | 5905082 | May 18, 2016 | DS DP | | |
| | 5905082*PED | Nov 18, 2016 | | | |
| | 6294540 | May 14, 2018 | DS DP | U-257 | |
| | 6294540*PED | Nov 14, 2018 | | | |
| <u>ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE - TRIZIVIR</u> | | | | | |
| 021205 001 | >A> 5034394 | Dec 18, 2011 | DS DP | | |
| | >A> 5034394*PED | Jun 18, 2012 | | | |
| <u>ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE - ULTRACET</u> | | | | | |
| 021123 001 | RE39221 | Aug 09, 2011 | DS DP | U-55 | |
| <u>ALBUTEROL SULFATE - PROAIR HFA</u> | | | | | |
| 021457 001 | | | | I-235 | Feb 03, 2009 |
| <u>ALBUTEROL SULFATE - VENTOLIN HFA</u> | | | | | |
| 020983 001 | 6131566 | Apr 14, 2015 | DP | U-716 | |
| | 6131566 | Apr 14, 2015 | DP | U-589 | |
| | 6532955 | Apr 14, 2015 | DP | U-716 | |
| | 6532955 | Apr 14, 2015 | DP | U-590 | |
| <u>ALENDRONATE SODIUM; CHOLECALCIFEROL - FOSAMAX PLUS D</u> | | | | | |
| 021762 001 | | | | NC | Apr 07, 2008 |
| <u>ALFUZOSIN HYDROCHLORIDE - UROXATRAL</u> | | | | | |
| 021287 001 | 4661491 | May 27, 2007 | | U-706 | |
| <u>ALITRETINOIN - PANRETIN</u> | | | | | |
| 020886 001 | 7056954 | Aug 02, 2012 | DP | | |
| <u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u> | | | | | |
| 020364 006 | 4879303 | Mar 25, 2007 | DS DP | NS | Apr 11, 2009 |
| | 6162802 | Dec 19, 2017 | DS DP | U-185 | |
| <u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u> | | | | | |
| 020364 007 | 4879303 | Mar 25, 2007 | DS DP | NS | Apr 11, 2009 |
| | 6162802 | Dec 19, 2017 | DS DP | U-185 | |
| <u>ANIDULAFUNGIN - ERAXIS</u> | | | | | |
| 021632 001 | 5965525 | Oct 12, 2016 | DS DP | U-540 | NCE |
| | 6384013 | Mar 19, 2012 | DS | | |
| | 6743777 | Mar 19, 2012 | DP | U-540 | |
| | 6960564 | Apr 12, 2021 | DP | U-540 | |
| <u>APREPITANT - EMEND</u> | | | | | |
| 021549 001 | 5145684 | Jan 25, 2011 | DP | | |
| | 5719147 | Jun 29, 2012 | DS DP | | |
| | 6096742 | Jul 01, 2018 | DS DP | U-745 | |
| | 6235735 | Jun 29, 2012 | | U-746 | |
| | 6235735 | Jun 29, 2012 | | U-747 | |
| <u>APREPITANT - EMEND</u> | | | | | |
| 021549 002 | 5145684 | Jan 25, 2011 | DP | | |
| | 5719147 | Jun 29, 2012 | DS DP | | |
| | 6096742 | Jul 01, 2018 | DS DP | U-745 | |
| | 6235735 | Jun 29, 2012 | | U-747 | |
| | 6235735 | Jun 29, 2012 | | U-746 | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|-----------------------------------|--|
| <u>APREPITANT - EMEND</u> | | | | | |
| 021549 003 | 5145684 | Jan 25, 2011 | DP | I-498 | Jun 30, 2009 |
| | 5719147 | Jun 29, 2012 | DS DP | NS | Jun 30, 2009 |
| | 6096742 | Jul 01, 2018 | DS DP U-745 | NCE | Mar 26, 2008 |
| | 6235735 | Jun 29, 2012 | U-747 | | |
| | 6235735 | Jun 29, 2012 | U-746 | | |
| <u>ARIPIPRAZOLE - ABILIFY</u> | | | | | |
| 021436 001 | | | | I-488 | Mar 01, 2008 |
| <u>ARIPIPRAZOLE - ABILIFY</u> | | | | | |
| 021436 002 | | | | I-488 | Mar 01, 2008 |
| <u>ARIPIPRAZOLE - ABILIFY</u> | | | | | |
| 021436 003 | | | | I-488 | Mar 01, 2008 |
| <u>ARIPIPRAZOLE - ABILIFY</u> | | | | | |
| 021436 004 | | | | I-488 | Mar 01, 2008 |
| <u>ARIPIPRAZOLE - ABILIFY</u> | | | | | |
| 021436 005 | | | | I-488 | Mar 01, 2008 |
| <u>ARIPIPRAZOLE - ABILIFY</u> | | | | | |
| 021436 006 | | | | I-488 | Mar 01, 2008 |
| <u>ARIPIPRAZOLE - ABILIFY</u> | | | | | |
| 021713 001 | 6977257 | Apr 24, 2022 | DS DP | I-488 | Mar 01, 2008 |
| <u>ARIPIPRAZOLE - ABILIFY</u> | | | | | |
| 021729 002 | | | | >A> I-488 >A> I-437 >A> NCE | Mar 01, 2008 Sep 29, 2007 Nov 15, 2007 |
| <u>ARIPIPRAZOLE - ABILIFY</u> | | | | | |
| 021729 003 | | | | >A> I-488 >A> I-437 >A> NCE | Mar 01, 2008 Sep 29, 2007 Nov 15, 2007 |
| <u>ARIPIPRAZOLE - ABILIFY</u> | | | | | |
| 021729 004 | | | | >A> I-488 >A> I-437 >A> NCE | Mar 01, 2008 Sep 29, 2007 Nov 15, 2007 |
| <u>ARIPIPRAZOLE - ABILIFY</u> | | | | | |
| 021729 005 | | | | >A> I-488 >A> I-437 >A> NCE | Mar 01, 2008 Sep 29, 2007 Nov 15, 2007 |
| <u>ARIPIPRAZOLE - ABILIFY</u> | | | | | |
| 021866 001 | | | | >A> NDF >A> NCE | Sep 20, 2009 Nov 15, 2007 |
| <u>ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE - SEPTOCAINE</u> | | | | | |
| 022010 001 | | | | NP | Mar 30, 2009 |
| <u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - MOVIPREP</u> | | | | | |
| 021881 001 | | | | NP | Aug 02, 2009 |
| <u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u> | | | | | |
| 021411 001 | 5658590 | Nov 26, 2016 | | U-494 | |
| | 5658590*PED | May 26, 2017 | | U-494 | |
| <u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u> | | | | | |
| 021411 002 | 5658590 | Nov 26, 2016 | | U-494 | |
| | 5658590*PED | May 26, 2017 | | U-494 | |
| <u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u> | | | | | |
| 021411 003 | 5658590 | Nov 26, 2016 | | U-494 | |
| | 5658590*PED | May 26, 2017 | | U-494 | |
| <u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u> | | | | | |
| 021411 004 | 5658590 | Nov 26, 2016 | | U-494 | |
| | 5658590*PED | May 26, 2017 | | U-494 | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u> | | | | | |
| 021411 005 | 5658590 | Nov 26, 2016 | U-494 | | |
| | 5658590*PED | May 26, 2017 | U-494 | | |
| <u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u> | | | | | |
| 021411 006 | 5658590 | Nov 26, 2016 | U-494 | | |
| | 5658590*PED | May 26, 2017 | U-494 | | |
| <u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u> | | | | | |
| 021411 007 | 5658590 | Nov 26, 2016 | U-494 | | |
| | 5658590*PED | May 26, 2017 | U-494 | | |
| <u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u> | | | | | |
| 021411 008 | 5658590 | Nov 26, 2016 | U-494 | | |
| | 5658590*PED | May 26, 2017 | U-494 | | |
| <u>AVOBENZONE; ECAMSULE; OCTOCRYLENE - ANTHELIOS SX</u> | | | | | |
| 021502 001 | 4585597 | Jun 16, 2007 | DS DP | U-752 | NC Jul 21, 2009 |
| | 5587150 | Dec 24, 2013 | DP | U-752 | |
| <u>AZELASTINE HYDROCHLORIDE - ASTELIN</u> | | | | | |
| 020114 001 | | | | | D-102 Feb 17, 2009 |
| <u>BALSALAZIDE DISODIUM - COLAZAL</u> | | | | | |
| 020610 001 | 4412992*PED | Jan 08, 2007 | | | |
| <u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE - TACLONEX</u> | | | | | |
| 021852 001 | 4866048 | Dec 29, 2007 | DS DP | U-88 | NC Jan 09, 2009 |
| | 4866048 | Dec 29, 2007 | DS DP | U-193 | |
| | 5763426 | Jun 09, 2015 | DS DP | | |
| | 6753013 | Jan 27, 2020 | DP | U-88 | |
| | 6753013 | Jan 27, 2020 | DP | U-193 | |
| <u>BETAMETHASONE VALERATE - LUXIQ</u> | | | | | |
| 020934 001 | 7078058 | Mar 01, 2016 | DS DP | | |
| <u>BETAXOLOL HYDROCHLORIDE - BETOPTIC S</u> | | | | | |
| 019845 001 | 4911920 | Mar 27, 2007 | | | |
| | 4911920*PED | Sep 27, 2007 | | | |
| <u>BIMATOPROST - LUMIGAN</u> | | | | | |
| 021275 001 | 5688819 | Aug 19, 2014 | | U-446 | |
| | 6403649 | Sep 21, 2012 | DS DP | U-446 | |
| <u>BIVALIRUDIN - ANGIOMAX</u> | | | | | |
| 020873 001 | | | | | I-486 Nov 30, 2008 |
| <u>BORTEZOMIB - VELCADE</u> | | | | | |
| 021602 001 | | | | | ODE Mar 25, 2012 |
| <u>BRIMONIDINE TARTRATE - ALPHAGAN P</u> | | | | | |
| 021770 001 | 5424078 | Jun 13, 2012 | DP | | |
| | 5424078*PED | Dec 13, 2012 | | | |
| | 6562873 | Jul 10, 2021 | DP | | |
| | 6562873*PED | Jan 10, 2022 | | | |
| | 6627210 | Jul 18, 2021 | DP | | |
| | 6627210*PED | Jan 18, 2022 | | | |
| | 6641834 | Jul 28, 2021 | DP | | |
| | 6641834*PED | Jan 28, 2022 | | | |
| | 6673337 | Jul 26, 2021 | DP | | |
| | 6673337*PED | Jan 26, 2022 | | | |
| <u>BRINZOLAMIDE - AZOPT</u> | | | | | |
| 020816 001 | 5240923 | Aug 31, 2010 | | U-224 | |
| | 5240923*PED | Mar 01, 2011 | | | |
| | 5378703 | Apr 01, 2012 | | U-224 | |
| | 5378703*PED | Oct 01, 2012 | | | |
| | 5461081 | Oct 24, 2012 | | U-225 | |
| | 5461081*PED | Apr 24, 2013 | | | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-----------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>BROMFENAC SODIUM - XIBROM</u> | | | | | |
| 021664 | 001 | | | I-485 | Jan 27, 2009 |
| <u>BUDESONIDE - BUDESONIDE</u> | | | | | |
| 021949 | 001 | | | NP | Jul 12, 2009 |
| <u>BUDESONIDE - BUDESONIDE</u> | | | | | |
| 021949 | 002 | | | NP | Jul 12, 2009 |
| <u>BUDESONIDE - PULMICORT RESPULES</u> | | | | | |
| 020929 | 001 | 6598603*PED | Jun 23, 2019 | | |
| | | 6899099 | Dec 23, 2018 | U-751 | |
| | | 6899099*PED | Jun 23, 2019 | | |
| <u>BUDESONIDE - PULMICORT RESPULES</u> | | | | | |
| 020929 | 002 | 6899099 | Dec 23, 2018 | U-751 | |
| | | 6899099*PED | Jun 23, 2019 | | |
| <u>BUDESONIDE - RHINOCORT</u> | | | | | |
| 020746 | 001 | 6986904 | Apr 29, 2017 | DP U-699 | |
| <u>BUDESONIDE - RHINOCORT</u> | | | | | |
| 020746 | 002 | 6986904 | Apr 29, 2017 | DP U-699 | |
| <u>BUDESONIDE; FORMOTEROL FUMARATE - SYMBICORT</u> | | | | | |
| 021929 | 001 | 5674860 | Oct 07, 2014 | DP U-754 | NC Jul 21, 2009 |
| | | 5972919 | Dec 17, 2012 | DP U-754 | |
| | | 6123924 | Sep 26, 2017 | DP | |
| | | 6641800 | Sep 26, 2017 | DP | |
| <u>BUDESONIDE; FORMOTEROL FUMARATE - SYMBICORT</u> | | | | | |
| 021929 | 002 | 5674860 | Oct 07, 2014 | DP U-754 | NC Jul 21, 2009 |
| | | 5972919 | Dec 17, 2012 | DP U-754 | |
| | | 6123924 | Sep 26, 2017 | DP | |
| | | 6641800 | Sep 26, 2017 | DP | |
| <u>BUPROPION HYDROCHLORIDE - WELLBUTRIN XL</u> | | | | | |
| 021515 | 001 | | | I-497 | Jun 12, 2009 |
| <u>BUPROPION HYDROCHLORIDE - WELLBUTRIN XL</u> | | | | | |
| 021515 | 002 | | | I-497 | Jun 12, 2009 |
| <u>CALCIUM CARBONATE; RISEDRONATE SODIUM - ACTONEL WITH CALCIUM (COPACKAGED)</u> | | | | | |
| 021823 | 001 | | | M-52 | Jan 24, 2009 |
| <u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 100</u> | | | | | |
| 021485 | 002 | 4963590 | Nov 27, 2007 | DP U-219 | |
| | | 5112861 | May 12, 2009 | U-219 | |
| | | 5135950 | Oct 31, 2010 | DS DP U-219 | |
| | | 6500867 | Jun 29, 2020 | DP U-219 | |
| <u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 150</u> | | | | | |
| 021485 | 003 | 4963590 | Nov 27, 2007 | DP U-219 | |
| | | 5112861 | May 12, 2009 | U-219 | |
| | | 5135950 | Oct 31, 2010 | DS DP U-219 | |
| | | 6500867 | Jun 29, 2020 | DP U-219 | |
| <u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 50</u> | | | | | |
| 021485 | 001 | 4963590 | Nov 27, 2007 | DP U-219 | |
| | | 5112861 | May 12, 2009 | U-219 | |
| | | 5135950 | Oct 31, 2010 | DS DP U-219 | |
| | | 6500867 | Jun 29, 2020 | DP U-219 | |
| <u>CARVEDILOL - COREG</u> | | | | | |
| 020297 | 001 | | | M-56 | Aug 28, 2009 |
| <u>CARVEDILOL - COREG</u> | | | | | |
| 020297 | 002 | | | M-56 | Aug 28, 2009 |
| <u>CARVEDILOL - COREG</u> | | | | | |
| 020297 | 003 | | | M-56 | Aug 28, 2009 |
| <u>CARVEDILOL - COREG</u> | | | | | |
| 020297 | 004 | | | M-56 | Aug 28, 2009 |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>CEFDITOREN PIVOXIL - SPECTRACEF</u> | | | | | |
| 021222 001 | 4839350 | Apr 14, 2009 | DS DP | | |
| <u>CELECOXIB - CELEBREX</u> | | | | | |
| 020998 001 | 5466823 | Nov 30, 2013 | DS | I-466 | Jul 29, 2008 |
| | 5466823*PED | May 30, 2014 | | PED | Jan 29, 2009 |
| | 5563165 | Nov 30, 2013 | DP | | |
| | 5563165*PED | May 30, 2014 | | | |
| | 5760068 | Jun 02, 2015 | | U-672 | |
| | 5760068 | Jun 02, 2015 | | U-299 | |
| | 5760068*PED | Dec 02, 2015 | | | |
| | 5972986 | Oct 14, 2017 | | U-299 | |
| | 5972986*PED | Apr 14, 2018 | | | |
| <u>CELECOXIB - CELEBREX</u> | | | | | |
| 020998 002 | 5466823 | Nov 30, 2013 | DS | I-466 | Jul 29, 2008 |
| | 5466823*PED | May 30, 2014 | | PED | Jan 29, 2009 |
| | 5563165 | Nov 30, 2013 | DP | | |
| | 5563165*PED | May 30, 2014 | | | |
| | 5760068 | Jun 02, 2015 | | U-672 | |
| | 5760068 | Jun 02, 2015 | | U-299 | |
| | 5760068*PED | Dec 02, 2015 | | | |
| | 5972986 | Oct 14, 2017 | | U-299 | |
| | 5972986*PED | Apr 14, 2018 | | | |
| <u>CELECOXIB - CELEBREX</u> | | | | | |
| 020998 003 | 5466823 | Nov 30, 2013 | DS | I-466 | Jul 29, 2008 |
| | 5466823*PED | May 30, 2014 | | PED | Jan 29, 2009 |
| | 5563165 | Nov 30, 2013 | DP | | |
| | 5563165*PED | May 30, 2014 | | | |
| | 5760068 | Jun 02, 2015 | | U-672 | |
| | 5760068 | Jun 02, 2015 | | U-299 | |
| | 5760068*PED | Dec 02, 2015 | | | |
| | 5972986 | Oct 14, 2017 | | U-299 | |
| | 5972986*PED | Apr 14, 2018 | | | |
| <u>CETIRIZINE HYDROCHLORIDE - ZYRTEC</u> | | | | | |
| 019835 001 | 4525358 | Jun 25, 2007 | DS DP | U-565 | |
| | 4525358*PED | Dec 25, 2007 | | | |
| <u>CETIRIZINE HYDROCHLORIDE - ZYRTEC</u> | | | | | |
| 019835 002 | 4525358 | Jun 25, 2007 | DS DP | U-565 | |
| | 4525358*PED | Dec 25, 2007 | | | |
| <u>CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ZYRTEC-D 12 HOUR</u> | | | | | |
| 021150 001 | 7014867 | Jun 10, 2022 | | DP | |
| <u>CHLORHEXIDINE GLUCONATE - HALO</u> | | | | | |
| 021669 001 | 7066916 | Feb 17, 2024 | | U-737 | |
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u> | | | | | |
| 020832 004 | 5690958 | Sep 30, 2016 | | DP | |
| | 6536975 | Nov 10, 2020 | | DP | |
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP FREPP</u> | | | | | |
| 020832 003 | 5538353 | Aug 25, 2015 | | DP | |
| | 5690958 | Sep 30, 2016 | | DP | |
| | 5752363 | Apr 22, 2017 | | DP | |
| | 5772346 | Apr 22, 2017 | | DP | |
| | D386849 | Nov 25, 2011 | | DP | |
| | D396911 | Aug 11, 2012 | | DP | |
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP SINGLE SWABSTICK</u> | | | | | |
| 021555 002 | 5690958 | Sep 30, 2016 | | DP | |
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u> | | | | | |
| 020832 002 | 6991393 | Mar 14, 2023 | | DP | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u> | | | | | |
| 020832 005 | 5690958 | Sep 30, 2016 | DP | | |
| | 6536975 | Nov 10, 2020 | DP | | |
| | 6729786 | Mar 14, 2023 | DP | | |
| | 6991393 | Jan 31, 2024 | DP | | |
| <u>CICLOPIROX - LOPROX</u> | | | | | |
| 020519 001 | 7018656 | Sep 05, 2018 | DP | | |
| | 7026337 | Apr 02, 2018 | | U-714 | |
| <u>CIPROFLOXACIN - CIPRO</u> | | | | | |
| 019847 002 | | | | >A> I-421 | Mar 25, 2007 |
| | | | | >A> PED | Sep 25, 2007 |
| <u>CIPROFLOXACIN - CIPRO</u> | | | | | |
| 019847 003 | | | | >A> I-421 | Mar 25, 2007 |
| | | | | >A> PED | Sep 25, 2007 |
| <u>CLOBETASOL PROPIONATE - CLOBEX</u> | | | | | |
| 021835 001 | 5972920 | Feb 12, 2018 | DP | NDF | Oct 27, 2008 |
| | 5990100 | Mar 24, 2018 | DP | U-742 | |
| <u>CLOPIDOGREL BISULFATE - CLOPIDOGREL BISULFATE</u> | | | | | |
| 076274 001 | | | | >A> PC | Feb 04, 2007 |
| <u>CLOPIDOGREL BISULFATE - PLAVIX</u> | | | | | |
| 020839 001 | | | | I-502 | Aug 17, 2009 |
| <u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u> | | | | | |
| 021176 001 | >A> 5693675 | Dec 02, 2014 | DS | | |
| | >A> 5917007 | Apr 29, 2014 | DS | U-323 | |
| | >A> 5919832 | Jun 10, 2014 | DS | | |
| | >A> 6066678 | Jun 10, 2014 | DS | U-323 | |
| | >A> 6433026 | Jun 10, 2014 | DS | | |
| | >A> 6784254 | Jun 10, 2014 | DS | DP | |
| | >A> 7101960 | Apr 29, 2014 | DS | DP | U-757 |
| <u>CONIVAPTAN HYDROCHLORIDE - VAPRISOL</u> | | | | | |
| 021697 001 | 5723606 | Mar 03, 2015 | DS | DP | U-698 |
| <u>DAPSONE - ACZONE</u> | | | | | |
| 021794 001 | 5863560 | Sep 11, 2016 | DP | | |
| | 6620435 | Sep 11, 2016 | DP | | |
| <u>DAPTOMYCIN - CUBICIN</u> | | | | | |
| 021572 001 | 6468967 | Sep 24, 2019 | | U-282 | |
| | RE39071 | Jun 15, 2016 | DS | DP | U-728 |
| <u>DAPTOMYCIN - CUBICIN</u> | | | | | |
| 021572 002 | 6468967 | Sep 24, 2019 | | U-282 | |
| | RE39071 | Jun 15, 2016 | DS | DP | U-728 |
| <u>DARUNAVIR ETHANOLATE - PREZISTA</u> | | | | | |
| 021976 001 | 5843946 | Dec 01, 2015 | DP | U-744 | NCE Jun 23, 2011 |
| | 6248775 | Aug 25, 2012 | DS | | |
| | 6335460 | Aug 25, 2012 | DS | DP | U-744 |
| <u>DASATINIB - SPRYCEL</u> | | | | | |
| 021986 001 | 6596746 | Apr 13, 2020 | DS | DP | U-748 NCE Jun 28, 2011 |
| <u>DASATINIB - SPRYCEL</u> | | | | | |
| 021986 002 | 6596746 | Apr 13, 2020 | DS | DP | U-748 NCE Jun 28, 2011 |
| <u>DASATINIB - SPRYCEL</u> | | | | | |
| 021986 003 | 6596746 | Apr 13, 2020 | DS | DP | U-748 NCE Jun 28, 2011 |
| <u>DECITABINE - DACOGEN</u> | | | | | |
| 021790 001 | | | | NCE | May 02, 2011 |
| <u>DEFERASIROX - EXJADE</u> | | | | | |
| 021882 001 | 6465504 | Jun 24, 2017 | DS | DP | |
| | 6596750 | Jun 24, 2017 | DS | | U-735 |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-----------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>DEFERASIROX - EXJADE</u> | | | | | |
| 021882 002 | 6465504 | Jun 24, 2017 | DS DP | | |
| | 6596750 | Jun 24, 2017 | DS | U-735 | |
| <u>DEFERASIROX - EXJADE</u> | | | | | |
| 021882 003 | 6465504 | Jun 24, 2017 | DS DP | | |
| | 6596750 | Jun 24, 2017 | DS | U-735 | |
| <u>DESFLURANE - SUPRANE</u> | | | | | |
| 020118 001 | >A> 4762856 | Feb 02, 2007 | | U-67 | |
| | >A> 4762856*PED | Aug 02, 2007 | | | |
| | >A> 5617906 | Apr 08, 2014 | DP | | |
| | >A> 5617906*PED | Oct 08, 2014 | | | |
| <u>DESLORATADINE - CLARINEX</u> | | | | | |
| 021165 001 | 4659716 | Mar 31, 2007 | DP | U-725 | |
| | 4659716 | Mar 31, 2007 | DP | U-427 | |
| | 4659716*PED | Oct 01, 2007 | | U-427 | |
| <u>DESLORATADINE - CLARINEX</u> | | | | | |
| 021300 001 | 4659716 | Mar 31, 2007 | DP | U-725 | |
| | 4659716 | Mar 31, 2007 | DP | U-611 | |
| | 4659716*PED | Oct 01, 2007 | | | |
| <u>DESLORATADINE - CLARINEX</u> | | | | | |
| 021312 001 | 4659716 | Mar 31, 2007 | DP | U-725 | |
| | 4659716 | Mar 31, 2007 | DP | U-427 | |
| | 4659716*PED | Oct 01, 2007 | | U-427 | |
| | 5178878 | Jan 12, 2010 | DP | | |
| | 5607697 | Jun 07, 2015 | DP | | |
| <u>DESLORATADINE - CLARINEX</u> | | | | | |
| 021312 002 | 4659716 | Mar 31, 2007 | DP | U-725 | |
| | 4659716 | Mar 31, 2007 | DP | U-427 | |
| | 4659716*PED | Oct 01, 2007 | DP | | |
| | 5178878 | Jan 12, 2010 | DP | | |
| | 5607697 | Jun 07, 2015 | DP | | |
| <u>DESLORATADINE; PSEUDOEPHEDRINE SULFATE - CLARINEX D 24 HOUR</u> | | | | | |
| 021605 001 | 4659716 | Mar 31, 2007 | DP | U-726 | |
| | 4659716 | Mar 31, 2007 | DP | U-644 | |
| | 4659716*PED | Oct 01, 2007 | | | |
| | 6979463 | Mar 28, 2022 | DP | | |
| <u>DESLORATADINE; PSEUDOEPHEDRINE SULFATE - CLARINEX-D 12 HOUR</u> | | | | | |
| 021313 001 | 4659716 | Mar 31, 2007 | DP | U-707 | >A> NP |
| | 4659716*PED | Oct 01, 2007 | | | NCE |
| | 6100274 | Jul 07, 2019 | DP | | NC |
| | 6100274*PED | Jan 07, 2020 | | | PED |
| | 6709676 | Feb 18, 2021 | DP | U-707 | |
| <u>DESMOPRESSIN ACETATE - DDAVP</u> | | | | | |
| 019955 001 | 7022340 | Apr 30, 2023 | DP | | |
| <u>DESMOPRESSIN ACETATE - DDAVP</u> | | | | | |
| 019955 002 | 7022340 | Apr 30, 2023 | DP | | |
| <u>DESONIDE - VERDESO</u> | | | | | |
| 021978 001 | | | | >A> NDF | Sep 19, 2009 |
| <u>DEXMEDETOMIDINE - PRECEDEX</u> | | | | | |
| 021038 001 | >A> 4910214 | Jul 15, 2013 | DS DP | U-421 | |
| <u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u> | | | | | |
| 021802 004 | | | | >A> NDF | May 26, 2008 |
| <u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u> | | | | | |
| 021392 001 | >A> 7108866 | Dec 17, 2019 | DP | U-107 | |
| <u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u> | | | | | |
| 021392 002 | >A> 7108866 | Dec 17, 2019 | DP | U-107 | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u> | | | | | |
| 021392 003 | >A> 7108866 | Dec 17, 2019 | DP U-107 | | |
| <u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u> | | | | | |
| 021392 004 | >A> 7108866 | Dec 17, 2019 | DP U-107 | | |
| <u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u> | | | | | |
| 021392 005 | >A> 7108866 | Dec 17, 2019 | DP U-107 | | |
| <u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u> | | | | | |
| 021392 006 | >A> 7108866 | Dec 17, 2019 | DP U-107 | | |
| <u>DOCETAXEL - TAXOTERE</u> | | | | | |
| 020449 001 | 5750561 | Jul 03, 2012 | DP | I-490 | Mar 22, 2009 |
| <u>DONEPEZIL HYDROCHLORIDE - ARICEPT ODT</u> | | | | | |
| 021720 001 | 4895841 | Nov 25, 2010 | DS DP | U-713 | |
| <u>DONEPEZIL HYDROCHLORIDE - ARICEPT ODT</u> | | | | | |
| 021720 002 | 4895841 | Nov 25, 2010 | DS DP | U-713 | |
| <u>DROSPIRENONE; ETHINYL ESTRADIOL - YAZ</u> | | | | | |
| 021676 001 | 5569652 | Oct 29, 2013 | | U-1 | NP |
| | 5798338 | Jul 10, 2015 | DP | | |
| | 6787531 | Aug 31, 2020 | DP | | |
| | 6933395 | Aug 11, 2017 | DP | | |
| | 6958326 | Dec 20, 2021 | DP | | |
| | RE37564 | Jun 30, 2014 | DP | | |
| | RE37838 | Jun 30, 2014 | DP | | |
| | RE38253 | Jun 30, 2014 | DP | | |
| <u>EFAVIRENZ - SUSTIVA</u> | | | | | |
| 020972 001 | 5519021 | May 21, 2013 | DS DP | | |
| | 5663169 | Sep 02, 2014 | | U-257 | |
| | 5811423 | Aug 07, 2012 | DS DP | U-256 | |
| | 6238695 | Apr 06, 2019 | DP | | |
| <u>EFAVIRENZ - SUSTIVA</u> | | | | | |
| 020972 002 | 5519021 | May 21, 2013 | DS DP | | |
| | 5663169 | Sep 02, 2014 | | U-257 | |
| | 5811423 | Aug 07, 2012 | DS DP | U-256 | |
| | 6238695 | Apr 06, 2019 | DP | | |
| <u>EFAVIRENZ - SUSTIVA</u> | | | | | |
| 020972 003 | 5519021 | May 21, 2013 | DS DP | | |
| | 5663169 | Sep 02, 2014 | | U-257 | |
| | 5811423 | Aug 07, 2012 | DS DP | U-256 | |
| | 6238695 | Apr 06, 2019 | DP | | |
| <u>EFAVIRENZ - SUSTIVA</u> | | | | | |
| 021360 002 | 5519021 | May 21, 2013 | DS DP | | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|----|------------------------|--------------------------------|
| <u>EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - ATRIPLA</u> | | | | | | |
| 021937 001 | 5210085 | May 11, 2010 | | | U-750 | |
| | 5210085*PED | Nov 11, 2010 | | | | |
| | 5519021 | May 21, 2013 | DS | DP | | |
| | 5663169 | Sep 02, 2014 | | | U-750 | |
| | 5811423 | Aug 07, 2012 | | | U-750 | |
| | 5814639 | Sep 29, 2015 | DS | DP | | |
| | 5814639*PED | Mar 29, 2016 | | | | |
| | 5914331 | Sep 29, 2015 | DS | | | |
| | 5914331*PED | Mar 29, 2016 | | | | |
| | 5922695 | Jul 25, 2017 | DS | | U-750 | |
| | 5935946 | Jul 25, 2017 | DS | DP | U-750 | |
| | 5977089 | Jul 25, 2017 | DS | DP | U-750 | |
| | 6043230 | Jul 25, 2017 | | | U-750 | |
| | 6238695 | Apr 06, 2019 | | DP | | |
| | 6555133 | Apr 06, 2019 | | | U-750 | |
| | 6639071 | Nov 13, 2021 | DS | | | |
| | 6642245 | Nov 04, 2020 | | | U-750 | |
| | 6642245*PED | May 04, 2021 | | | | |
| | 6703396 | Mar 09, 2021 | DS | DP | | |
| | 6703396*PED | Sep 09, 2021 | | | | |
| | 6939964 | Jan 20, 2018 | DS | | | |
| <u>EMTRICITABINE - EMTRIVA</u> | | | | | | |
| 021500 001 | 5210085 | May 11, 2010 | | | NCE | Jul 02, 2008 |
| | 5210085*PED | Nov 11, 2010 | | | PED | Jan 02, 2009 |
| | 5814639 | Sep 29, 2015 | | | | |
| | 5814639*PED | Mar 29, 2016 | | | | |
| | 5914331 | Sep 29, 2015 | | | | |
| | 5914331*PED | Mar 29, 2016 | | | | |
| | 6642245 | Nov 04, 2020 | | | U-541 | |
| | 6642245*PED | May 04, 2021 | | | | |
| | 6703396 | Mar 09, 2021 | DS | DP | | |
| | 6703396*PED | Sep 09, 2021 | | | | |
| <u>EMTRICITABINE - EMTRIVA</u> | | | | | | |
| 021896 001 | 5210085 | May 11, 2010 | | | U-257 | |
| | 5210085*PED | Nov 11, 2010 | | | NDF | Sep 27, 2008 |
| | 5814639 | Sep 29, 2015 | DS | DP | NCE | Jul 02, 2008 |
| | 5814639*PED | Mar 29, 2016 | | | PED | Mar 27, 2009 |
| | 5914331 | Sep 29, 2015 | DS | | PED | Jan 02, 2009 |
| | 5914331*PED | Mar 29, 2016 | | | | |
| | 6642245 | Nov 04, 2020 | | | U-257 | |
| | 6642245*PED | May 04, 2021 | | | | |
| | 6703396 | Mar 09, 2021 | DS | DP | | |
| | 6703396*PED | Sep 09, 2021 | | | | |
| <u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u> | | | | | | |
| 021752 001 | 5210085 | May 11, 2010 | | | U-248 | |
| | 5210085 | May 11, 2010 | | | U-541 | |
| | 5210085*PED | Nov 11, 2010 | | | NCE | Jul 02, 2008 |
| | 5814639 | Sep 29, 2015 | DS | DP | PED | Jan 02, 2009 |
| | 5814639*PED | Mar 29, 2016 | | | | |
| | 5914331 | Sep 29, 2015 | DS | DP | U-248 | |
| | 5914331*PED | Mar 29, 2016 | | | | |
| | 6642245 | Nov 04, 2020 | | | U-248 | |
| | 6642245 | Nov 04, 2020 | | | U-541 | |
| | 6642245*PED | May 04, 2021 | | | | |
| | 6703396 | Mar 09, 2021 | DS | DP | | |
| | 6703396*PED | Sep 09, 2021 | | | | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>ENTACAPONE - COMTAN</u> | | | | | |
| 020796 001 | 4963590 | Nov 27, 2007 | DP U-219 | | |
| | 5112861 | May 12, 2009 | U-219 | | |
| | 5135950 | Oct 31, 2010 | DS DP U-219 | | |
| | 6599530 | Sep 14, 2018 | DP U-219 | | |
| <u>EPLERENONE - INSPRA</u> | | | | | |
| 021437 001 | 4559332 | Apr 09, 2007 | DS DP U-537 | | |
| <u>EPLERENONE - INSPRA</u> | | | | | |
| 021437 002 | 4559332 | Apr 09, 2007 | DS DP U-537 | | |
| <u>EPLERENONE - INSPRA</u> | | | | | |
| 021437 003 | 4559332 | Apr 09, 2007 | DS DP U-537 | | |
| <u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u> | | | | | |
| 021153 001 | 4738974 | Apr 19, 2007 | DS DP U-635 | I-484 | Nov 24, 2007 |
| | 4738974 | Apr 19, 2007 | DS DP U-373 | NPP | Apr 28, 2009 |
| | 4738974*PED | Oct 19, 2007 | U-373 | | |
| | 4786505 | Apr 20, 2007 | DP U-373 | | |
| | 4786505 | Apr 20, 2007 | DP U-729 | | |
| | 4853230 | Apr 20, 2007 | DP U-729 | | |
| | 4853230 | Apr 20, 2007 | DP U-373 | | |
| | 5690960 | Nov 25, 2014 | DP U-729 | | |
| | 5690960 | Nov 25, 2014 | DP U-373 | | |
| | 5714504 | Feb 03, 2015 | DP U-729 | | |
| | 5714504 | Feb 03, 2015 | DP U-373 | | |
| | 5877192 | May 27, 2014 | DP U-373 | | |
| | 5877192 | May 27, 2014 | DP U-729 | | |
| | 5900424 | May 04, 2016 | DS U-729 | | |
| | 5900424 | May 04, 2016 | DS U-373 | | |
| | 6369085 | May 25, 2018 | DS DP U-729 | | |
| | 6428810 | Nov 03, 2019 | DP U-469 | | |
| | 6428810 | Nov 03, 2019 | DP U-729 | | |
| <u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u> | | | | | |
| 021153 002 | 4738974 | Apr 19, 2007 | DS DP U-635 | I-484 | Nov 24, 2007 |
| | 4738974 | Apr 19, 2007 | DS DP U-373 | NPP | Apr 28, 2009 |
| | 4738974*PED | Oct 19, 2007 | U-373 | | |
| | 4786505 | Apr 20, 2007 | DP U-373 | | |
| | 4786505 | Apr 20, 2007 | DP U-729 | | |
| | 4853230 | Apr 20, 2007 | DP U-729 | | |
| | 4853230 | Apr 20, 2007 | DP U-373 | | |
| | 5690960 | Nov 25, 2014 | DP U-729 | | |
| | 5690960 | Nov 25, 2014 | DP U-373 | | |
| | 5714504 | Feb 03, 2015 | DP U-729 | | |
| | 5714504 | Feb 03, 2015 | DP U-373 | | |
| | 5877192 | May 27, 2014 | DP U-373 | | |
| | 5877192 | May 27, 2014 | DP U-729 | | |
| | 5900424 | May 04, 2016 | DS U-729 | | |
| | 5900424 | May 04, 2016 | DS U-373 | | |
| | 6369085 | May 25, 2018 | DS DP U-729 | | |
| | 6428810 | Nov 03, 2019 | DP U-469 | | |
| | 6428810 | Nov 03, 2019 | DP U-729 | | |
| <u>ESTRADIOL - ESTRADIOL</u> | | | | | |
| 075182 004 | | | | PC | Feb 06, 2007 |
| <u>ESTRADIOL - ESTRADIOL</u> | | | | | |
| 075182 005 | | | | PC | Feb 06, 2007 |
| <u>ETHINYL ESTRADIOL; LEVONORGESTREL - SEASONIQUE</u> | | | | | |
| 021840 001 | | | | NP | May 25, 2009 |
| <u>ETHINYL ESTRADIOL; NORELGESTROMIN - ORTHO EVRA</u> | | | | | |
| 021180 001 | 5876746 | Nov 20, 2015 | DP U-514 | | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>ETHINYL ESTRADIOL; NORETHINDRONE - OVCON-35</u> | | | | | |
| 021490 001 | 6667050 | Apr 06, 2019 | DP U-1 | | |
| <u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LOESTRIN 24 FE</u> | | | | | |
| 021871 001 | 5552394 | Jul 22, 2014 | U-1 | NP | Feb 17, 2009 |
| <u>ETONOGESTREL - IMPLANON</u> | | | | | |
| 021529 001 | 4957119 | Aug 05, 2008 | DP | NDF | Jul 17, 2009 |
| | 5150718 | Sep 29, 2009 | U-749 | | |
| <u>EXEMESTANE - AROMASIN</u> | | | | | |
| 020753 001 | | | | I-495 | Oct 05, 2008 |
| <u>EZETIMIBE - ZETIA</u> | | | | | |
| 021445 001 | 7030106 | Jan 25, 2022 | DP | I-493 | May 23, 2009 |
| | >A> RE37721 | Oct 25, 2016 | DS DP | U-473 | |
| <u>EZETIMIBE; SIMVASTATIN - VYTORIN</u> | | | | | |
| 021687 001 | >A> RE37721 | Oct 25, 2016 | DS DP | U-473 | |
| <u>EZETIMIBE; SIMVASTATIN - VYTORIN</u> | | | | | |
| 021687 002 | >A> RE37721 | Oct 25, 2016 | DS DP | U-473 | |
| <u>EZETIMIBE; SIMVASTATIN - VYTORIN</u> | | | | | |
| 021687 003 | >A> RE37721 | Oct 25, 2016 | DS DP | U-473 | |
| <u>EZETIMIBE; SIMVASTATIN - VYTORIN</u> | | | | | |
| 021687 004 | >A> RE37721 | Oct 25, 2016 | DS DP | U-473 | |
| <u>FAMCICLOVIR - FAMVIR</u> | | | | | |
| 020363 001 | | | | D-103 I-501 | Jul 28, 2009 Jul 28, 2009 |
| <u>FAMCICLOVIR - FAMVIR</u> | | | | | |
| 020363 002 | | | | D-103 I-501 | Jul 28, 2009 Jul 28, 2009 |
| <u>FAMCICLOVIR - FAMVIR</u> | | | | | |
| 020363 003 | | | | D-103 I-501 | Jul 28, 2009 Jul 28, 2009 |
| <u>FENOFIBRATE - ANTARA (MICRONIZED)</u> | | | | | |
| 021695 001 | 7101574 | Aug 20, 2020 | DS DP | M-47 | Oct 21, 2008 |
| <u>FENOFIBRATE - ANTARA (MICRONIZED)</u> | | | | | |
| 021695 003 | 7101574 | Aug 20, 2020 | DS DP | M-47 | Oct 21, 2008 |
| <u>FENOFIBRATE - FENOFIBRATE</u> | | | | | |
| 076433 001 | | | | PC | May 22, 2006 |
| <u>FENOFIBRATE - FENOFIBRATE</u> | | | | | |
| 076433 002 | | | | PC | May 22, 2006 |
| <u>FENOFIBRATE - LIPOFEN</u> | | | | | |
| 021612 001 | 5545628 | Jan 10, 2015 | | U-701 | |
| <u>FENOFIBRATE - LIPOFEN</u> | | | | | |
| 021612 002 | 5545628 | Jan 10, 2015 | | U-701 | |
| <u>FENOFIBRATE - LIPOFEN</u> | | | | | |
| 021612 003 | 5545628 | Jan 10, 2015 | | U-701 | |
| <u>FENOFIBRATE - TRICOR</u> | | | | | |
| 021656 001 | 7037529 | Jan 09, 2018 | | DP | |
| | 7041319 | Jan 09, 2018 | | DP | |
| <u>FENOFIBRATE - TRICOR</u> | | | | | |
| 021656 002 | 7037529 | Jan 09, 2018 | | DP | |
| | 7041319 | Jan 09, 2018 | | DP | |
| <u>FENTANYL CITRATE - FENTORA</u> | | | | | |
| 021947 001 | | | | >A> NDF | Sep 25, 2009 |
| <u>FENTANYL CITRATE - FENTORA</u> | | | | | |
| 021947 002 | | | | >A> NDF | Sep 25, 2009 |
| <u>FENTANYL CITRATE - FENTORA</u> | | | | | |
| 021947 003 | | | | >A> NDF | Sep 25, 2009 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>FENTANYL CITRATE - FENTORA</u> | | | | | |
| 021947 004 | | | | >A> NDF | Sep 25, 2009 |
| <u>FENTANYL CITRATE - FENTORA</u> | | | | | |
| 021947 005 | | | | >A> NDF | Sep 25, 2009 |
| <u>FENTANYL HYDROCHLORIDE - IONSYS</u> | | | | | |
| 021338 001 | 5232438 | Oct 03, 2008 | DP U-736 | NDF | May 22, 2009 |
| | 5445606 | Dec 11, 2011 | DP | | |
| | 5697896 | Dec 16, 2014 | DP | | |
| | 5843014 | Dec 01, 2015 | DP | | |
| | 6169920 | Jan 02, 2018 | DP | | |
| | 6171294 | Jun 05, 2015 | | U-736 | |
| | 6181963 | Nov 02, 2019 | DP | | |
| | 6195582 | Jan 28, 2019 | DP | U-736 | |
| | 6216033 | Jun 05, 2015 | DP | | |
| | 6317629 | Jun 02, 2012 | DP | | |
| | 6425892 | Jun 05, 2015 | | U-736 | |
| | 6842640 | Jun 02, 2015 | DP | | |
| | 6881208 | Apr 19, 2022 | | U-736 | |
| | 6975902 | Apr 01, 2024 | DP | | |
| | 7018370 | Jun 05, 2015 | | U-736 | |
| | 7027859 | Sep 26, 2014 | DP | | |
| <u>FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ALLEGRA D 24 HOUR</u> | | | | | |
| 021704 001 | RE39069 | May 29, 2018 | DP | | |
| <u>FINASTERIDE - FINASTERIDE</u> | | | | | |
| 076340 001 | | | | PC | Dec 16, 2006 |
| <u>FLUNISOLIDE - AEROSPAN HFA</u> | | | | | |
| 021247 001 | | | | NP | Jan 27, 2009 |
| <u>FLUOCINOLONE ACETONIDE - RETISERT</u> | | | | | |
| 021737 001 | 6217895 | Mar 22, 2019 | DP U-708 | | |
| | 6548078 | Mar 22, 2019 | DP U-708 | | |
| <u>FLUOCINONIDE - VANOS</u> | | | | | |
| 021758 001 | | | | I-487 | Mar 02, 2009 |
| <u>FLUOXETINE HYDROCHLORIDE - PROZAC WEEKLY</u> | | | | | |
| 021235 001 | RE39030 | May 29, 2017 | DP U-397 | | |
| | RE39030 | May 29, 2017 | DP U-396 | | |
| <u>FLUTICASONE PROPIONATE - FLOVENT HFA</u> | | | | | |
| 021433 001 | 5658549 | Sep 19, 2014 | DP U-710 | NPP | Feb 28, 2009 |
| | 5674472 | Oct 07, 2014 | DP | | |
| | 6251368 | Dec 04, 2012 | DP | | |
| <u>FLUTICASONE PROPIONATE - FLOVENT HFA</u> | | | | | |
| 021433 002 | 5658549 | Sep 19, 2014 | DP U-710 | NPP | Feb 28, 2009 |
| | 5674472 | Oct 07, 2014 | DP | | |
| | 6251368 | Dec 04, 2012 | DP | | |
| <u>FLUTICASONE PROPIONATE - FLOVENT HFA</u> | | | | | |
| 021433 003 | 5658549 | Sep 19, 2014 | DP U-710 | NPP | Feb 28, 2009 |
| | 5674472 | Oct 07, 2014 | DP | | |
| | 6251368 | Dec 04, 2012 | DP | | |
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 100/50</u> | | | | | |
| 021077 001 | 4992474 | Feb 12, 2008 | | U-211 | |
| | 4992474*PED | Aug 12, 2008 | | U-211 | |
| | 5126375 | Feb 12, 2008 | | | |
| | 5126375*PED | Aug 12, 2008 | | | |
| | 5225445 | Feb 12, 2008 | | U-211 | |
| | 5225445*PED | Aug 12, 2008 | | U-211 | |
| | 6536427 | Mar 01, 2011 | DP | | |
| | 6536427*PED | Sep 01, 2011 | | | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 250/50</u> | | | | | |
| 021077 002 | 4992474 | Feb 12, 2008 | | U-211 | |
| | 4992474*PED | Aug 12, 2008 | | U-211 | |
| | 5126375 | Feb 12, 2008 | | | |
| | 5126375*PED | Aug 12, 2008 | | | |
| | 5225445 | Feb 12, 2008 | | U-211 | |
| | 5225445*PED | Aug 12, 2008 | | U-211 | |
| | 6536427 | Mar 01, 2011 | DP | | |
| | 6536427*PED | Sep 01, 2011 | | | |
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 500/50</u> | | | | | |
| 021077 003 | 4992474 | Feb 12, 2008 | | U-211 | |
| | 4992474*PED | Aug 12, 2008 | | U-211 | |
| | 5126375 | Feb 12, 2008 | | | |
| | 5126375*PED | Aug 12, 2008 | | | |
| | 5225445 | Feb 12, 2008 | | U-211 | |
| | 5225445*PED | Aug 12, 2008 | | U-211 | |
| | 6536427 | Mar 01, 2011 | DP | | |
| | 6536427*PED | Sep 01, 2011 | | | |
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u> | | | | | |
| 021254 001 | 4992474 | Feb 12, 2008 | DS DP | U-738 | NP Jun 08, 2009 |
| | 4992474*PED | Aug 12, 2008 | | | |
| | 5126375 | Feb 12, 2008 | DS DP | | |
| | 5126375*PED | Aug 12, 2008 | | | |
| | 5225445 | Feb 12, 2008 | | U-738 | |
| | 5225445*PED | Aug 12, 2008 | | | |
| | 5270305 | Sep 07, 2010 | | U-738 | |
| | 5658549 | Aug 19, 2014 | DP | U-738 | |
| | 5674472 | Oct 07, 2014 | DP | | |
| | 6143277 | Apr 14, 2015 | DP | U-738 | |
| | 6251368 | Dec 04, 2012 | DP | | |
| | 6253762 | Apr 14, 2015 | DP | U-738 | |
| | 6315173 | Dec 23, 2017 | DP | | |
| | 6510969 | Dec 23, 2017 | DP | | |
| | 6524555 | Apr 14, 2015 | DP | | |
| | 6546928 | Apr 14, 2015 | DP | | |
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u> | | | | | |
| 021254 002 | 4992474 | Feb 12, 2008 | DS DP | U-738 | NP Jun 08, 2009 |
| | 4992474*PED | Aug 12, 2008 | | | |
| | 5126375 | Feb 12, 2008 | DS DP | | |
| | 5126375*PED | Aug 12, 2008 | | | |
| | 5225445 | Feb 12, 2008 | | U-738 | |
| | 5225445*PED | Aug 12, 2008 | | | |
| | 5270305 | Sep 07, 2010 | | U-738 | |
| | 5658549 | Aug 19, 2014 | DP | U-738 | |
| | 5674472 | Oct 07, 2014 | DP | | |
| | 6143277 | Apr 14, 2015 | DP | U-738 | |
| | 6251368 | Dec 04, 2012 | DP | | |
| | 6253762 | Apr 14, 2015 | DP | U-738 | |
| | 6315173 | Dec 23, 2017 | DP | | |
| | 6510969 | Dec 23, 2017 | DP | | |
| | 6524555 | Apr 14, 2015 | DP | | |
| | 6546928 | Apr 14, 2015 | DP | | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u> | | | | | |
| 021254 003 | 4992474 | Feb 12, 2008 | DS DP | U-738 | NP |
| | 4992474*PED | Aug 12, 2008 | | | |
| | 5126375 | Feb 12, 2008 | DS DP | | |
| | 5126375*PED | Aug 12, 2008 | | | |
| | 5225445 | Feb 12, 2008 | | U-738 | |
| | 5225445*PED | Aug 12, 2008 | | | |
| | 5270305 | Sep 07, 2010 | | U-738 | |
| | 5658549 | Aug 19, 2014 | DP | U-738 | |
| | 5674472 | Oct 07, 2014 | DP | | |
| | 6143277 | Apr 14, 2015 | DP | U-738 | |
| | 6251368 | Dec 04, 2012 | DP | | |
| | 6253762 | Apr 14, 2015 | DP | U-738 | |
| | 6315173 | Dec 23, 2017 | DP | | |
| | 6510969 | Dec 23, 2017 | DP | | |
| | 6524555 | Apr 14, 2015 | DP | | |
| | 6546928 | Apr 14, 2015 | DP | | |
| <u>FROVATRIPTAN SUCCINATE - FROVA</u> | | | | | |
| 021006 001 | 5464864 | Nov 07, 2015 | | U-436 | |
| <u>FULVESTRANT - FASLODEX</u> | | | | | |
| 021344 001 | >A> 4659516 | Dec 11, 2007 | DS DP | U-596 | |
| <u>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</u> | | | | | |
| 021615 001 | | | | NDF | Apr 01, 2008 |
| <u>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</u> | | | | | |
| 021615 002 | | | | NDF | Apr 01, 2008 |
| <u>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</u> | | | | | |
| 021615 003 | | | | NDF | Apr 01, 2008 |
| <u>GANIRELIX ACETATE - GANIRELIX ACETATE INJECTION</u> | | | | | |
| 021057 001 | 4801577 | Feb 05, 2012 | DS DP | | |
| <u>GEMCITABINE HYDROCHLORIDE - GEMZAR</u> | | | | | |
| 020509 001 | | | | I-499 | Jul 14, 2009 |
| <u>GEMCITABINE HYDROCHLORIDE - GEMZAR</u> | | | | | |
| 020509 002 | | | | I-499 | Jul 14, 2009 |
| <u>GEMIFLOXACIN MESYLATE - FACTIVE</u> | | | | | |
| 021158 001 | 5776944 | Apr 04, 2017 | DS DP | | |
| <u>GLIMEPIRIDE - AMARYL</u> | | | | | |
| 020496 001 | | | | M-54 PED | Nov 28, 2008 May 28, 2009 |
| <u>GLIMEPIRIDE - AMARYL</u> | | | | | |
| 020496 002 | | | | M-54 PED | Nov 28, 2008 May 28, 2009 |
| <u>GLIMEPIRIDE - AMARYL</u> | | | | | |
| 020496 003 | | | | M-54 PED | Nov 28, 2008 May 28, 2009 |
| <u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u> | | | | | |
| 021925 001 | 4687777 | Jan 17, 2011 | DS | | |
| | 6150383 | Jun 19, 2016 | | U-753 | |
| | 6211205 | Jun 19, 2016 | | U-753 | |
| | 6303640 | Aug 09, 2016 | | U-753 | |
| | 6329404 | Jun 19, 2016 | DP | U-753 | |
| <u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u> | | | | | |
| 021925 002 | 4687777 | Jan 17, 2011 | DS | | |
| | 6150383 | Jun 19, 2016 | | U-753 | |
| | 6211205 | Jun 19, 2016 | | U-753 | |
| | 6303640 | Aug 09, 2016 | | U-753 | |
| | 6329404 | Jun 19, 2016 | DP | U-753 | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u> | | | | | |
| 021700 001 | 5002953 | Sep 17, 2011 | DS DP | U-690 | |
| | 5002953*PED | Mar 17, 2012 | | | |
| | 5741803 | Apr 21, 2015 | DS DP | U-690 | |
| | 5741803*PED | Oct 21, 2015 | | | |
| <u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u> | | | | | |
| 021700 002 | 5002953 | Sep 17, 2011 | DS DP | U-690 | |
| | 5002953*PED | Mar 17, 2012 | | | |
| | 5741803 | Apr 21, 2015 | DS DP | U-690 | |
| | 5741803*PED | Oct 21, 2015 | | | |
| <u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u> | | | | | |
| 021700 003 | 5002953 | Sep 17, 2011 | DS DP | U-690 | |
| | 5002953*PED | Mar 17, 2012 | | | |
| | 5741803 | Apr 21, 2015 | DS DP | U-690 | |
| | 5741803*PED | Oct 21, 2015 | | | |
| <u>HYALURONIDASE RECOMBINANT HUMAN - HYLENEX RECOMBINANT</u> | | | | | |
| 021859 001 | | | | NCE | Dec 02, 2010 |
| <u>HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE - DUTOPROL</u> | | | | | |
| 021956 001 | | | | NC | Aug 28, 2009 |
| <u>HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE - DUTOPROL</u> | | | | | |
| 021956 002 | | | | NC | Aug 28, 2009 |
| <u>HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE - DUTOPROL</u> | | | | | |
| 021956 003 | | | | NC | Aug 28, 2009 |
| <u>HYDROCHLOROTHIAZIDE; VALSARTAN - DIOVAN HCT</u> | | | | | |
| 020818 004 | | | | NS | Apr 28, 2009 |
| <u>HYDROCHLOROTHIAZIDE; VALSARTAN - DIOVAN HCT</u> | | | | | |
| 020818 005 | | | | NS | Apr 28, 2009 |
| <u>IBANDRONATE SODIUM - BONIVA</u> | | | | | |
| 021455 001 | 4927814 | Jul 09, 2007 | DS DP | U-642 | |
| | 6143326 | Apr 21, 2017 | | U-642 | |
| <u>IBANDRONATE SODIUM - BONIVA</u> | | | | | |
| 021858 001 | 4927814 | Jul 09, 2007 | DS DP | U-700 | NDF Jan 06, 2009 |
| | 5662918 | Sep 02, 2014 | DP | | NCE May 16, 2008 |
| <u>IBUPROFEN LYSINE - NEOPROFEN</u> | | | | | |
| 021903 001 | | | | NE | Apr 13, 2009 |
| | | | | ODE | Apr 13, 2013 |
| <u>IMATINIB MESYLATE - GLEEVEC</u> | | | | | |
| 021335 001 | 5521184 | Jan 04, 2015 | DS DP | I-392 | May 20, 2006 |
| | 5521184*PED | Jul 04, 2015 | | I-376 | Dec 20, 2005 |
| | 6894051 | May 23, 2019 | DS DP | U-649 | NCE May 10, 2006 |
| | 6894051*PED | Nov 23, 2019 | | | ODE Feb 01, 2009 |
| | 6958335 | Dec 19, 2021 | DS DP | | ODE May 10, 2008 |
| | 6958335*PED | Jun 19, 2022 | | | PED Aug 01, 2009 |
| | | | | | PED Nov 10, 2008 |
| | | | | | PED Nov 20, 2006 |
| | | | | | PED Nov 10, 2006 |
| | | | | | PED Jun 20, 2006 |
| <u>IMATINIB MESYLATE - GLEEVEC</u> | | | | | |
| 021335 002 | 5521184 | Jan 04, 2015 | | I-392 | May 20, 2006 |
| | 5521184*PED | Jul 04, 2015 | | I-376 | Dec 20, 2005 |
| | 6894051 | May 23, 2019 | DS DP | U-649 | NCE May 10, 2006 |
| | 6894051*PED | Nov 23, 2019 | | | ODE Feb 01, 2009 |
| | 6958335 | Dec 19, 2021 | DS DP | | ODE May 10, 2008 |
| | 6958335*PED | Jun 19, 2022 | | | PED Aug 01, 2009 |
| | | | | | PED Nov 10, 2008 |
| | | | | | PED Nov 20, 2006 |
| | | | | | PED Nov 10, 2006 |
| | | | | | PED Nov 10, 2006 |
| | | | | | PED Jun 20, 2006 |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>IMATINIB MESYLATE - GLEEVEC</u> | | | | | |
| 021588 001 | 5521184 | Jan 04, 2015 | DS DP | I-392 | May 20, 2006 |
| | 5521184*PED | Jul 04, 2015 | | I-376 | Dec 20, 2005 |
| | 6894051 | May 23, 2019 | DS DP U-649 | NCE | May 10, 2006 |
| | 6894051*PED | Nov 23, 2019 | | ODE | Feb 01, 2009 |
| | 6958335 | Dec 19, 2021 | DS DP | ODE | May 10, 2008 |
| | 6958335*PED | Jun 19, 2022 | | PED | Aug 01, 2009 |
| | | | | PED | Nov 10, 2008 |
| | | | | PED | Nov 20, 2006 |
| | | | | PED | Nov 10, 2006 |
| | | | | PED | Jun 20, 2006 |
| <u>IMATINIB MESYLATE - GLEEVEC</u> | | | | | |
| 021588 002 | 5521184 | Jan 04, 2015 | | I-392 | May 20, 2006 |
| | 5521184*PED | Jul 04, 2015 | | I-376 | Dec 20, 2005 |
| | 6894051 | May 23, 2019 | DS DP U-649 | NCE | May 10, 2006 |
| | 6894051*PED | Nov 23, 2019 | | ODE | Feb 01, 2009 |
| | 6958335 | Dec 19, 2021 | DS DP | ODE | May 10, 2008 |
| | 6958335*PED | Jun 19, 2022 | | PED | Aug 01, 2009 |
| | | | | PED | Nov 10, 2008 |
| | | | | PED | Nov 20, 2006 |
| | | | | PED | Nov 10, 2006 |
| | | | | PED | Jun 20, 2006 |
| <u>INSULIN DETEMIR RECOMBINANT - LEVEMIR</u> | | | | | |
| 021536 001 | | | | I-489 | Oct 19, 2008 |
| <u>INSULIN RECOMBINANT HUMAN - EXUBERA</u> | | | | | |
| 021868 001 | 5740794 | Apr 21, 2015 | DP | NP | Jan 27, 2009 |
| | 5997848 | Mar 07, 2014 | | U-704 | |
| | 6051256 | Mar 07, 2014 | DP | | |
| | 6257233 | May 14, 2019 | | U-704 | |
| | 6423344 | Mar 07, 2014 | DP | | |
| | 6543448 | Sep 21, 2014 | DP | | |
| | 6546929 | May 14, 2019 | | U-704 | |
| | 6582728 | Jun 24, 2020 | DP | | |
| | 6592904 | Mar 07, 2014 | DP | | |
| | 6685967 | Sep 11, 2018 | DP | | |
| | 6737045 | Mar 07, 2014 | | U-704 | |
| | RE37872 | Feb 12, 2010 | DP | | |
| | RE38385 | Feb 12, 2010 | DP | | |
| <u>INSULIN RECOMBINANT HUMAN - EXUBERA</u> | | | | | |
| 021868 002 | 5740794 | Apr 21, 2015 | DP | NP | Jan 27, 2009 |
| | 5997848 | Mar 07, 2014 | | U-704 | |
| | 6051256 | Mar 07, 2014 | DP | | |
| | 6257233 | May 14, 2019 | | U-704 | |
| | 6423344 | Mar 07, 2014 | DP | | |
| | 6543448 | Sep 21, 2014 | DP | | |
| | 6546929 | May 14, 2019 | | U-704 | |
| | 6582728 | Jun 24, 2020 | DP | | |
| | 6592904 | Mar 07, 2014 | DP | | |
| | 6685967 | Sep 11, 2018 | DP | | |
| | 6737045 | Mar 07, 2014 | | U-704 | |
| | RE37872 | Feb 12, 2010 | DP | | |
| | RE38385 | Feb 12, 2010 | DP | | |
| <u>IPRATROPIUM BROMIDE - ATROVENT HFA</u> | | | | | |
| 021527 001 | 6983743 | May 26, 2020 | DP | | |
| <u>LAMOTRIGINE - LAMOTRIGINE</u> | | | | | |
| 076420 001 | | | | >A> PC | Feb 25, 2007 |
| <u>LAMOTRIGINE - LAMOTRIGINE</u> | | | | | |
| 076420 002 | | | | PC | Dec 25, 2006 |
| <u>LANSOPRAZOLE - PREVACID</u> | | | | | |
| 020406 001 | 6749864 | Feb 13, 2007 | DP | | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>LANSOPRAZOLE - PREVACID</u> | | | | | |
| 020406 002 | 6749864 | Feb 13, 2007 | DP | | |
| <u>LANSOPRAZOLE - PREVACID</u> | | | | | |
| 021281 001 | 6749864 | Feb 13, 2007 | DP | | |
| <u>LANSOPRAZOLE - PREVACID</u> | | | | | |
| 021281 002 | 6749864 | Feb 13, 2007 | DP | | |
| <u>LANSOPRAZOLE - PREVACID</u> | | | | | |
| 021428 001 | 6749864 | Feb 13, 2007 | DP | | |
| <u>LANSOPRAZOLE - PREVACID</u> | | | | | |
| 021428 002 | 6749864 | Feb 13, 2007 | DP | | |
| <u>LANTHANUM CARBONATE - FOSRENOL</u> | | | | | |
| 021468 003 | 5968976 | Mar 19, 2016 | DP | U-613 | |
| <u>LANTHANUM CARBONATE - FOSRENOL</u> | | | | | |
| 021468 004 | 5968976 | Mar 19, 2016 | DP | U-613 | |
| <u>LENALIDOMIDE - REVLIMID</u> | | | | | |
| 021880 001 | 5635517 | Jul 24, 2016 | DS | | Dec 27, 2012 |
| | 6045501 | Aug 28, 2018 | | U-694 | |
| | 6315720 | Oct 23, 2020 | | U-694 | |
| | 6555554 | Jul 24, 2016 | DP | | |
| | 6561976 | Aug 28, 2018 | | U-694 | |
| | 6561977 | Oct 23, 2020 | | U-694 | |
| | 6755784 | Oct 23, 2020 | | U-694 | |
| | 6908432 | Aug 28, 2018 | | U-694 | |
| <u>LENALIDOMIDE - REVLIMID</u> | | | | | |
| 021880 002 | 5635517 | Jul 24, 2016 | DS | | Dec 27, 2012 |
| | 6045501 | Aug 28, 2018 | | U-694 | |
| | 6315720 | Oct 23, 2020 | | U-694 | |
| | 6555554 | Jul 24, 2016 | DP | | |
| | 6561976 | Aug 28, 2018 | | U-694 | |
| | 6561977 | Oct 23, 2020 | | U-694 | |
| | 6755784 | Oct 23, 2020 | | U-694 | |
| | 6908432 | Aug 28, 2018 | | U-694 | |
| <u>LENALIDOMIDE - REVLIMID</u> | | | | | |
| 021880 003 | 5635517 | Jul 24, 2016 | DS | | Jun 29, 2009 |
| | 6045501 | Aug 28, 2018 | | U-694 | Dec 27, 2010 |
| | 6315720 | Oct 23, 2020 | | U-694 | |
| | 6555554 | Jul 24, 2016 | DP | | |
| | 6561976 | Aug 28, 2018 | | U-694 | |
| | 6561977 | Oct 23, 2020 | | U-694 | |
| | 6755784 | Oct 23, 2020 | | U-694 | |
| | 6908432 | Aug 28, 2018 | | U-694 | |
| <u>LENALIDOMIDE - REVLIMID</u> | | | | | |
| 021880 004 | 5635517 | Jul 24, 2016 | DS | | Jun 29, 2009 |
| | 6045501 | Aug 28, 2018 | | U-694 | Dec 27, 2010 |
| | 6315720 | Oct 23, 2020 | | U-694 | |
| | 6555554 | Jul 24, 2016 | DP | | |
| | 6561976 | Aug 28, 2018 | | U-694 | |
| | 6561977 | Oct 23, 2020 | | U-694 | |
| | 6755784 | Oct 23, 2020 | | U-694 | |
| | 6908432 | Aug 28, 2018 | | U-694 | |
| <u>LEVALBUTEROL TARTRATE - XOPENEX HFA</u> | | | | | |
| 021730 001 | >A> 5362755 | Mar 25, 2013 | | U-636 | |
| <u>LEVETIRACETAM - KEPPRA</u> | | | | | |
| 021035 004 | 4943639 | Jul 14, 2008 | DS | | Jun 21, 2008 |
| <u>LEVETIRACETAM - KEPPRA</u> | | | | | |
| 021872 001 | 4943639 | Jul 14, 2008 | DS | | Jul 31, 2009 |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>LEVOBETAXOLOL HYDROCHLORIDE - BETAXON</u> | | | | | |
| 021114 001 | 4911920 | Mar 27, 2007 | | U-369 | |
| | 4911920*PED | Sep 27, 2007 | | | |
| | 5540918 | Jul 30, 2013 | | | |
| | 5540918*PED | Jan 30, 2014 | | | |
| <u>LEVONORGESTREL - PLAN B</u> | | | | | |
| 021045 002 | | | | NP | Aug 24, 2009 |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | |
| 021301 001 | 7067148 | Feb 15, 2022 | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | |
| 021301 002 | 7067148 | Feb 15, 2022 | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | |
| 021301 003 | 7067148 | Feb 15, 2022 | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | |
| 021301 004 | 7067148 | Feb 15, 2022 | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | |
| 021301 005 | 7067148 | Feb 15, 2022 | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | |
| 021301 006 | 7067148 | Feb 15, 2022 | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | |
| 021301 007 | 7067148 | Feb 15, 2022 | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | |
| 021301 008 | 7067148 | Feb 15, 2022 | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | |
| 021301 009 | 7067148 | Feb 15, 2022 | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | |
| 021301 010 | 7067148 | Feb 15, 2022 | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | |
| 021301 011 | 7067148 | Feb 15, 2022 | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | |
| 021301 012 | 7067148 | Feb 15, 2022 | DP | | |
| <u>LIDOCAINE; TETRACAINE - LIDOCAINE AND TETRACAINE</u> | | | | | |
| 021717 001 | 5919479 | Jul 28, 2015 | DP | NP | Jun 29, 2009 |
| | 6528086 | Sep 28, 2019 | DP | | |
| <u>LIDOCAINE; TETRACAINE - SYNERA</u> | | | | | |
| 021623 001 | | | | NC | Jun 23, 2008 |
| <u>LOPINAVIR; RITONAVIR - KALETRA</u> | | | | | |
| 021906 001 | 5541206 | Jul 30, 2013 | DS DP | U-688 | |
| | 5541206*PED | Jan 30, 2014 | | | |
| | 5635523 | Jun 03, 2014 | | U-688 | |
| | 5635523*PED | Dec 03, 2014 | | | |
| | 5648497 | Jul 15, 2014 | DS DP | | |
| | 5648497*PED | Jan 15, 2015 | | | |
| | 5674882 | Oct 07, 2014 | | U-688 | |
| | 5674882*PED | Apr 07, 2015 | | | |
| | 5846987 | Dec 29, 2012 | | U-688 | |
| | 5846987*PED | Jun 29, 2013 | | | |
| | 5886036 | Dec 29, 2012 | DP | | |
| | 5886036*PED | Jun 29, 2013 | | | |
| | 6037157 | Jun 26, 2016 | | U-688 | |
| | 6037157*PED | Dec 26, 2016 | | | |
| | 6703403 | Jun 26, 2016 | | U-688 | |
| | 6703403*PED | Dec 26, 2016 | | | |
| <u>LOVASTATIN; NIACIN - ADVICOR</u> | | | | | |
| 021249 001 | 7011848 | Sep 20, 2013 | | U-712 | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-----------|---------------------------|-----------------|----------------------------|--|
| <u>LOVASTATIN; NIACIN - ADVICOR</u> | | | | | |
| 021249 002 | 7011848 | Sep 20, 2013 | | U-712 | |
| <u>LOVASTATIN; NIACIN - ADVICOR</u> | | | | | |
| 021249 003 | 7011848 | Sep 20, 2013 | | U-712 | |
| <u>LUBIPROSTONE - AMITIZA</u> | | | | | |
| 021908 001 | 5284858 | Feb 08, 2011 | DS DP | NCE | Jan 31, 2011 |
| | 5317032 | May 31, 2011 | DS DP | U-717 | |
| | 6414016 | Sep 05, 2020 | DS DP | U-717 | |
| | 6583174 | Oct 16, 2020 | DS DP | | |
| | 7064148 | Aug 30, 2022 | DS DP | U-739 | |
| <u>MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u> | | | | | |
| 021850 001 | 6489346 | Jul 16, 2016 | DS DP | U-623 | |
| | 6489346 | Jul 16, 2016 | DS DP | U-588 | |
| | 6645988 | Jul 16, 2016 | DS DP | U-623 | |
| | 6645988 | Jul 16, 2016 | DS DP | U-588 | |
| | 6699885 | Jul 16, 2016 | | U-623 | |
| | 6699885 | Jul 16, 2016 | | U-588 | |
| <u>MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u> | | | | | |
| 021850 002 | 6489346 | Jul 16, 2016 | DS DP | U-623 | |
| | 6489346 | Jul 16, 2016 | DS DP | U-588 | |
| | 6645988 | Jul 16, 2016 | DS DP | U-623 | |
| | 6645988 | Jul 16, 2016 | DS DP | U-588 | |
| | 6699885 | Jul 16, 2016 | | U-623 | |
| | 6699885 | Jul 16, 2016 | | U-588 | |
| <u>MECASERMIN RINFABATE RECOMBINANT - IPLEX</u> | | | | | |
| 021884 001 | 5200509 | Apr 06, 2010 | DS | | |
| | 5681818 | Oct 28, 2014 | | U-697 | |
| <u>MEGESTROL ACETATE - MEGACE ES</u> | | | | | |
| 021778 001 | 7101576 | Apr 22, 2024 | | U-755 | |
| <u>MELOXICAM - MOBIC</u> | | | | | |
| 020938 001 | | | | ODE PED | Aug 11, 2012 Feb 11, 2013 |
| <u>MELOXICAM - MOBIC</u> | | | | | |
| 020938 002 | | | | ODE PED | Aug 11, 2012 Feb 11, 2013 |
| <u>MELOXICAM - MOBIC</u> | | | | | |
| 021530 001 | | | | I-469 ODE PED PED | Aug 11, 2008 Aug 11, 2012 Feb 11, 2013 Feb 11, 2009 |
| <u>MEQUINOL; TRETINOIN - SOLAGE</u> | | | | | |
| 020922 001 | 5194247 | Dec 10, 2013 | DP | U-294 | |
| <u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u> | | | | | |
| 021410 001 | 5002953 | Sep 17, 2011 | DS DP | U-690 | I-494 |
| | 5002953 | Sep 17, 2011 | DS DP | U-734 | May 19, 2009 |
| | 5002953 | Sep 17, 2011 | DS DP | U-691 | |
| | 5002953 | Sep 17, 2011 | DS DP | U-493 | |
| | 5741803 | Apr 21, 2015 | DS DP | U-734 | |
| | 5741803 | Apr 21, 2015 | DS DP | U-493 | |
| <u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u> | | | | | |
| 021410 002 | 5002953 | Sep 17, 2011 | DS DP | U-690 | I-494 |
| | 5002953 | Sep 17, 2011 | DS DP | U-734 | May 19, 2009 |
| | 5002953 | Sep 17, 2011 | DS DP | U-691 | |
| | 5002953 | Sep 17, 2011 | DS DP | U-493 | |
| | 5741803 | Apr 21, 2015 | DS DP | U-734 | |
| | 5741803 | Apr 21, 2015 | DS DP | U-493 | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u> | | | | | |
| 021410 003 | 5002953 | Sep 17, 2011 | DS DP | U-691 | I-494 May 19, 2009 |
| | 5002953 | Sep 17, 2011 | DS DP | U-734 | |
| | 5002953 | Sep 17, 2011 | DS DP | U-690 | |
| | 5002953 | Sep 17, 2011 | DS DP | U-493 | |
| | 5741803 | Apr 21, 2015 | DS DP | U-734 | |
| | 5741803 | Apr 21, 2015 | DS DP | U-493 | |
| <u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u> | | | | | |
| 021410 004 | 5002953 | Sep 17, 2011 | DS DP | U-691 | I-494 May 19, 2009 |
| | 5002953 | Sep 17, 2011 | DS DP | U-734 | |
| | 5002953 | Sep 17, 2011 | DS DP | U-690 | |
| | 5002953 | Sep 17, 2011 | DS DP | U-493 | |
| | 5741803 | Apr 21, 2015 | DS DP | U-734 | |
| | 5741803 | Apr 21, 2015 | DS DP | U-493 | |
| <u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u> | | | | | |
| 021410 005 | 5002953 | Sep 17, 2011 | DS DP | U-690 | I-494 May 19, 2009 |
| | 5002953 | Sep 17, 2011 | DS DP | U-734 | |
| | 5002953 | Sep 17, 2011 | DS DP | U-691 | |
| | 5002953 | Sep 17, 2011 | DS DP | U-493 | |
| | 5741803 | Apr 21, 2015 | DS DP | U-734 | |
| | 5741803 | Apr 21, 2015 | DS DP | U-493 | |
| | 5741803*PED | Oct 21, 2015 | | | |
| <u>METHYLPHENIDATE - DAYTRANA</u> | | | | | |
| 021514 001 | 5958446 | Dec 12, 2012 | DP | | NDF Apr 06, 2009 |
| | 6210705 | Sep 30, 2018 | DP | U-727 | |
| | 6348211 | Sep 30, 2018 | DP | U-727 | |
| <u>METHYLPHENIDATE - DAYTRANA</u> | | | | | |
| 021514 002 | 5958446 | Dec 12, 2012 | DP | | NDF Apr 06, 2009 |
| | 6210705 | Sep 30, 2018 | DP | U-727 | |
| | 6348211 | Sep 30, 2018 | DP | U-727 | |
| <u>METHYLPHENIDATE - DAYTRANA</u> | | | | | |
| 021514 003 | 5958446 | Dec 12, 2012 | DP | | NDF Apr 06, 2009 |
| | 6210705 | Sep 30, 2018 | DP | U-727 | |
| | 6348211 | Sep 30, 2018 | DP | U-727 | |
| <u>METHYLPHENIDATE - DAYTRANA</u> | | | | | |
| 021514 004 | 5958446 | Dec 12, 2012 | DP | | NDF Apr 06, 2009 |
| | 6210705 | Sep 30, 2018 | DP | U-727 | |
| | 6348211 | Sep 30, 2018 | DP | U-727 | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u> | | | | | |
| 021259 001 | 6344215 | Oct 27, 2020 | DP | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u> | | | | | |
| 021259 002 | 6344215 | Oct 27, 2020 | DP | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u> | | | | | |
| 021259 003 | 6344215 | Oct 27, 2020 | DP | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u> | | | | | |
| 021259 004 | 6344215 | Oct 27, 2020 | DP | | |
| <u>METOPROLOL SUCCINATE - TOPROL-XL</u> | | | | | |
| 019962 001 | 4927640 | May 22, 2007 | DP | | D-95 Feb 15, 2008 |
| | 4927640*PED | Nov 22, 2007 | | | PED Aug 15, 2008 |
| | 4957745 | Sep 18, 2007 | DP | U-107 | |
| | 4957745*PED | Mar 18, 2008 | | | |
| | 5001161 | Sep 18, 2007 | DP | | |
| | 5001161*PED | Mar 18, 2008 | | | |
| | 5081154 | Sep 18, 2007 | DS | | |
| | 5081154*PED | Mar 18, 2008 | | | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>METOPROLOL SUCCINATE - TOPROL-XL</u> | | | | | |
| 019962 002 | 4927640 | May 22, 2007 | DP | D-95 | Feb 15, 2008 |
| | 4927640*PED | Nov 22, 2007 | | PED | Aug 15, 2008 |
| | 4957745 | Sep 18, 2007 | DP | U-107 | |
| | 4957745*PED | Mar 18, 2008 | | | |
| | 5001161 | Sep 18, 2007 | DP | | |
| | 5001161*PED | Mar 18, 2008 | | | |
| | 5081154 | Sep 18, 2007 | DS | | |
| | 5081154*PED | Mar 18, 2008 | | | |
| <u>METOPROLOL SUCCINATE - TOPROL-XL</u> | | | | | |
| 019962 003 | 4927640 | May 22, 2007 | DP | D-95 | Feb 15, 2008 |
| | 4927640*PED | Nov 22, 2007 | | PED | Aug 15, 2008 |
| | 4957745 | Sep 18, 2007 | DP | U-107 | |
| | 4957745*PED | Mar 18, 2008 | | | |
| | 5001161 | Sep 18, 2007 | DP | | |
| | 5001161*PED | Mar 18, 2008 | | | |
| | 5081154 | Sep 18, 2007 | DS | | |
| | 5081154*PED | Mar 18, 2008 | | | |
| <u>METOPROLOL SUCCINATE - TOPROL-XL</u> | | | | | |
| 019962 004 | 4927640 | May 22, 2007 | DP | D-95 | Feb 15, 2008 |
| | 4927640*PED | Nov 22, 2007 | | PED | Aug 15, 2008 |
| | 4957745 | Sep 18, 2007 | DP | U-107 | |
| | 4957745*PED | Mar 18, 2008 | | | |
| | 5001161 | Sep 18, 2007 | DP | U-107 | |
| | 5001161*PED | Mar 18, 2008 | | | |
| | 5081154 | Sep 18, 2007 | DS | U-107 | |
| | 5081154*PED | Mar 18, 2008 | | | |
| <u>METRONIDAZOLE - METROGEL</u> | | | | | |
| 021789 001 | 6881726 | Feb 21, 2022 | DP | U-743 | |
| <u>MICONAZOLE NITRATE; PETROLATUM, WHITE; ZINC OXIDE - VUSION</u> | | | | | |
| 021026 001 | 4911932 | Mar 27, 2007 | DP | U-718 | NP Feb 16, 2009 |
| <u>MINOXIDIL - MEN'S ROGAINE</u> | | | | | |
| 021812 001 | 6946120 | Apr 20, 2019 | DP | U-702 | NDF Jan 20, 2009 |
| <u>MODAFINIL - PROVIGIL</u> | | | | | |
| 020717 001 | 4927855 | May 22, 2007 | | U-255 | I-449 Jan 23, 2007 |
| | 4927855*PED | Nov 22, 2007 | | | ODE Dec 24, 2005 |
| | RE37516 | Oct 06, 2014 | | U-255 | PED Jul 23, 2007 |
| | RE37516*PED | Apr 06, 2015 | | | PED Jun 24, 2006 |
| <u>MODAFINIL - PROVIGIL</u> | | | | | |
| 020717 002 | 4927855 | May 22, 2007 | | U-255 | I-449 Jan 23, 2007 |
| | 4927855*PED | Nov 22, 2007 | | | ODE Dec 24, 2005 |
| | RE37516 | Oct 06, 2014 | | U-255 | PED Jul 23, 2007 |
| | RE37516*PED | Apr 06, 2015 | | | PED Jun 24, 2006 |
| <u>MORPHINE SULFATE - KADIAN</u> | | | | | |
| 020616 004 | 5378474 | Mar 23, 2010 | | | |
| <u>MORPHINE SULFATE - KADIAN</u> | | | | | |
| 020616 005 | 5202128 | Apr 13, 2010 | | | |
| | 5378474 | Mar 23, 2010 | | | |
| <u>MOXIFLOXACIN HYDROCHLORIDE - AVELOX</u> | | | | | |
| 021085 001 | 4990517 | Dec 08, 2011 | DS DP | U-298 | |
| | 6610327 | Oct 29, 2019 | DP | U-298 | |
| <u>MOXIFLOXACIN HYDROCHLORIDE - AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER</u> | | | | | |
| 021277 001 | 4990517 | Dec 08, 2011 | DS DP | U-298 | |
| | 6548079 | Jul 25, 2020 | DP | U-298 | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>MOXIFLOXACIN HYDROCHLORIDE - VIGAMOX</u> | | | | | |
| 021598 001 | 4990517 | Dec 08, 2011 | DS DP | U-709 | |
| | 4990517*PED | Jun 08, 2012 | | | |
| <u>NALTREXONE - VIVITROL</u> | | | | | |
| 021897 001 | 5792477 | May 02, 2017 | DP | NDF | Apr 13, 2009 |
| | 5916598 | May 02, 2017 | DP | | |
| | 6110503 | May 02, 2017 | DP | | |
| | 6194006 | Dec 30, 2018 | DP | | |
| | 6264987 | May 19, 2020 | DP | | |
| | 6331317 | Nov 12, 2019 | DP | | |
| | 6379703 | Dec 30, 2018 | DP | | |
| | 6379704 | May 19, 2020 | DP | | |
| | 6395304 | Nov 12, 2019 | DP | | |
| | 6403114 | May 02, 2017 | DP | | |
| | 6495164 | May 25, 2020 | DP | | |
| | 6495166 | Nov 12, 2019 | DP | | |
| | 6534092 | May 19, 2020 | DP | | |
| | 6537586 | Nov 12, 2019 | DP | | |
| | 6596316 | Dec 30, 2018 | DP | | |
| | 6667061 | May 25, 2020 | DP | | |
| | 6713090 | Nov 12, 2019 | DP | | |
| | 6939033 | Nov 12, 2019 | DP | | |
| <u>NELARABINE - ARRANON</u> | | | | | |
| 021877 001 | 5747472 | Feb 20, 2013 | | U-696 | |
| | 5747472 | Feb 20, 2013 | | U-695 | |
| | 5747472 | Feb 20, 2013 | | U-689 | |
| | 5821236 | Feb 20, 2013 | | U-695 | |
| <u>NIACIN - NIASPAN</u> | | | | | |
| 020381 001 | 7011848 | Sep 20, 2013 | | U-712 | |
| <u>NIACIN - NIASPAN</u> | | | | | |
| 020381 002 | 7011848 | Sep 20, 2013 | | U-712 | |
| <u>NIACIN - NIASPAN</u> | | | | | |
| 020381 003 | 7011848 | Sep 20, 2013 | | U-712 | |
| <u>NIACIN - NIASPAN</u> | | | | | |
| 020381 004 | 7011848 | Sep 20, 2013 | | U-712 | |
| <u>NIACIN - NIASPAN TITRATION STARTER PACK</u> | | | | | |
| 020381 005 | 7011848 | Sep 20, 2013 | | U-712 | |
| <u>NICOTINE POLACRILEX - NICOTINE POLACRILEX</u> | | | | | |
| 077007 001 | | | | PC | Aug 21, 2006 |
| <u>NICOTINE POLACRILEX - NICOTINE POLACRILEX</u> | | | | | |
| 077007 002 | | | | PC | Aug 21, 2006 |
| <u>OCTREOTIDE ACETATE - SANDOSTATIN</u> | | | | | |
| 019667 001 | 5753618 | May 19, 2015 | | | |
| | 5753618*PED | Nov 19, 2015 | | | |
| <u>OCTREOTIDE ACETATE - SANDOSTATIN</u> | | | | | |
| 019667 002 | 5753618 | May 19, 2015 | | | |
| | 5753618*PED | Nov 19, 2015 | | | |
| <u>OCTREOTIDE ACETATE - SANDOSTATIN</u> | | | | | |
| 019667 003 | 5753618 | May 19, 2015 | | | |
| | 5753618*PED | Nov 19, 2015 | | | |
| <u>OCTREOTIDE ACETATE - SANDOSTATIN</u> | | | | | |
| 019667 004 | 5753618 | May 19, 2015 | | | |
| | 5753618*PED | Nov 19, 2015 | | | |
| <u>OCTREOTIDE ACETATE - SANDOSTATIN</u> | | | | | |
| 019667 005 | 5753618 | May 19, 2015 | | | |
| | 5753618*PED | Nov 19, 2015 | | | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---|-----------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>OCTREOTIDE ACETATE - SANDOSTATIN LAR</u> | | | | | |
| 021008 001 | 5538739 | Jul 23, 2013 | DP | M-55 | May 10, 2009 |
| | 5538739*PED | Jan 23, 2014 | | ODE | Nov 25, 2005 |
| | 5639480 | Jun 17, 2014 | DP | PED | Nov 10, 2009 |
| | 5639480*PED | Dec 17, 2014 | | PED | May 25, 2006 |
| | 5688530 | Nov 18, 2014 | | U-268 | |
| | 5688530*PED | May 18, 2015 | | | |
| | 5922338 | Jul 13, 2016 | DP | | |
| | 5922338*PED | Jan 13, 2017 | | | |
| | 5922682 | Jul 13, 2016 | DP | | |
| | 5922682*PED | Jan 13, 2017 | | | |
| <u>OCTREOTIDE ACETATE - SANDOSTATIN LAR</u> | | | | | |
| 021008 002 | 5538739 | Jul 23, 2013 | | M-55 | May 10, 2009 |
| | 5538739*PED | Jan 23, 2014 | | ODE | Nov 25, 2005 |
| | 5639480 | Jun 17, 2014 | DP | PED | Nov 10, 2009 |
| | 5639480*PED | Dec 17, 2014 | | PED | May 25, 2006 |
| | 5688530 | Nov 18, 2014 | | U-268 | |
| | 5688530*PED | May 18, 2015 | | | |
| | 5922338 | Jul 13, 2016 | DP | | |
| | 5922338*PED | Jan 13, 2017 | | | |
| | 5922682 | Jul 13, 2016 | DP | | |
| | 5922682*PED | Jan 13, 2017 | | | |
| <u>OCTREOTIDE ACETATE - SANDOSTATIN LAR</u> | | | | | |
| 021008 003 | 5538739 | Jul 23, 2013 | | M-55 | May 25, 2009 |
| | 5538739*PED | Jan 23, 2014 | | ODE | Nov 25, 2005 |
| | 5639480 | Jun 17, 2014 | DP | PED | Nov 25, 2009 |
| | 5639480*PED | Dec 17, 2014 | | PED | May 25, 2006 |
| | 5688530 | Nov 18, 2014 | | U-268 | |
| | 5688530*PED | May 18, 2015 | | | |
| | 5922338 | Jul 13, 2016 | DP | | |
| | 5922338*PED | Jan 13, 2017 | | | |
| | 5922682 | Jul 13, 2016 | DP | | |
| | 5922682*PED | Jan 13, 2017 | | | |
| <u>OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u> | | | | | |
| 021849 001 | 6489346 | Jul 16, 2016 | DS DP | U-588 | |
| | 6645988 | Jul 16, 2016 | DS DP | | |
| | 6699885 | Jul 16, 2016 | | U-588 | |
| <u>OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u> | | | | | |
| 021849 002 | 6489346 | Jul 16, 2016 | DS DP | U-623 | |
| | 6489346 | Jul 16, 2016 | DS DP | U-588 | |
| | 6645988 | Jul 16, 2016 | DS DP | | |
| | 6699885 | Jul 16, 2016 | | U-623 | |
| | 6699885 | Jul 16, 2016 | | U-588 | |
| <u>OXALIPLATIN - ELOXATIN</u> | | | | | |
| 021492 001 | >A> 5290961 | Jan 12, 2013 | | >A> I-441 | Nov 04, 2007 |
| | >A> 5290961*PED | Jul 12, 2013 | | >A> I-425 | Jan 09, 2007 |
| | >A> 5338874 | Apr 07, 2013 | DS | >A> NCE | Aug 09, 2007 |
| | >A> 5338874*PED | Oct 07, 2013 | | >A> PED | May 04, 2008 |
| | >A> 5420319 | Aug 09, 2016 | DS | >A> PED | Feb 09, 2008 |
| | >A> 5420319*PED | Aug 09, 2016 | | >A> PED | Jul 09, 2007 |
| <u>OXALIPLATIN - ELOXATIN</u> | | | | | |
| 021492 002 | >A> 5290961 | Jan 12, 2013 | | >A> I-441 | Nov 04, 2007 |
| | >A> 5290961*PED | Jul 12, 2013 | | >A> I-425 | Jan 09, 2007 |
| | >A> 5338874 | Apr 07, 2013 | DS | >A> NCE | Aug 09, 2007 |
| | >A> 5338874*PED | Oct 07, 2013 | | >A> PED | May 04, 2008 |
| | >A> 5420319 | Aug 09, 2016 | DS | >A> PED | Feb 09, 2008 |
| | >A> 5420319*PED | Aug 09, 2016 | | >A> PED | Jul 09, 2007 |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-----------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>OXALIPLATIN - ELOXATIN</u> | | | | | |
| 021759 001 | >A> 5290961 | Jan 12, 2013 | DS | >A> I-441 | Nov 04, 2007 |
| | >A> 5290961*PED | Jul 12, 2013 | | >A> NCE | Aug 09, 2007 |
| | >A> 5338874 | Apr 07, 2013 | DS | >A> PED | May 04, 2008 |
| | >A> 5338874*PED | Oct 07, 2013 | | >A> PED | Feb 09, 2008 |
| | >A> 5420319 | Aug 08, 2016 | DS | | |
| | >A> 5420319*PED | Feb 08, 2017 | | | |
| | >A> 5716988 | Aug 07, 2015 | DP | | |
| | >A> 5716988*PED | Feb 07, 2016 | | | |
| <u>OXALIPLATIN - ELOXATIN</u> | | | | | |
| 021759 002 | >A> 5290961 | Jan 12, 2013 | DS | >A> I-441 | Nov 04, 2007 |
| | >A> 5290961*PED | Jul 12, 2013 | | >A> NCE | Aug 09, 2007 |
| | >A> 5338874 | Apr 07, 2013 | DS | >A> PED | May 04, 2008 |
| | >A> 5338874*PED | Oct 07, 2013 | | >A> PED | Feb 09, 2008 |
| | >A> 5420319 | Aug 08, 2016 | DS | | |
| | >A> 5420319*PED | Feb 08, 2017 | | | |
| | >A> 5716988 | Aug 07, 2015 | DP | | |
| | >A> 5716988*PED | Feb 07, 2016 | | | |
| <u>OXCARBAZEPINE - TRILEPTAL</u> | | | | | |
| 021014 001 | 7037525 | Feb 12, 2018 | | U-724 | |
| <u>OXCARBAZEPINE - TRILEPTAL</u> | | | | | |
| 021014 002 | 7037525 | Feb 12, 2018 | | U-724 | |
| <u>OXCARBAZEPINE - TRILEPTAL</u> | | | | | |
| 021014 003 | 7037525 | Feb 12, 2018 | | U-724 | |
| <u>OXCARBAZEPINE - TRILEPTAL</u> | | | | | |
| 021285 001 | 7037525 | Feb 12, 2018 | | U-724 | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA</u> | | | | | |
| 021611 001 | | | | NDF | Jun 22, 2009 |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA</u> | | | | | |
| 021611 002 | | | | NDF | Jun 22, 2009 |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | |
| 021610 001 | 5128143 | Sep 19, 2008 | DP | NDF | Jun 22, 2009 |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | |
| 021610 002 | 5128143 | Sep 19, 2008 | DP | NDF | Jun 22, 2009 |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | |
| 021610 003 | 5128143 | Sep 19, 2008 | DP | NDF | Jun 22, 2009 |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | |
| 021610 004 | 5128143 | Sep 19, 2008 | DP | NDF | Jun 22, 2009 |
| <u>PAROXETINE HYDROCHLORIDE - PAXIL</u> | | | | | |
| 020031 004 | 6133289 | May 19, 2015 | | U-358 | |
| <u>PAROXETINE HYDROCHLORIDE - PAXIL</u> | | | | | |
| 020031 005 | 6133289 | May 19, 2015 | | U-358 | |
| <u>POSACONAZOLE - NOXAFIL</u> | | | | | |
| 022003 001 | | | | >A> NCE | Sep 15, 2011 |
| <u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u> | | | | | |
| 020667 001 | 4843086 | Jun 27, 2006 | | U-231 | |
| <u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u> | | | | | |
| 020667 002 | 4843086 | Jun 27, 2006 | | U-231 | |
| <u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u> | | | | | |
| 020667 003 | 4843086 | Jun 27, 2006 | | U-231 | |
| <u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u> | | | | | |
| 020667 004 | 4843086 | Jun 27, 2006 | | U-231 | |
| <u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u> | | | | | |
| 020667 005 | 4843086 | Jun 27, 2006 | | U-231 | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u> | | | | | |
| 020667 006 | 4843086 | Jun 27, 2006 | | U-231 | |
| <u>PRAVASTATIN SODIUM - PRAVASTATIN SODIUM</u> | | | | | |
| 076056 001 | | | | PC | Oct 21, 2006 |
| <u>PRAVASTATIN SODIUM - PRAVASTATIN SODIUM</u> | | | | | |
| 076056 002 | | | | PC | Oct 21, 2006 |
| <u>PRAVASTATIN SODIUM - PRAVASTATIN SODIUM</u> | | | | | |
| 076056 003 | | | | PC | Oct 21, 2006 |
| <u>QUETIAPINE FUMARATE - SEROQUEL</u> | | | | | |
| 020639 006 | 4879288 | Sep 26, 2011 | DS DP | U-550 | |
| <u>QUETIAPINE FUMARATE - SEROQUEL</u> | | | | | |
| 020639 007 | 4879288 | Sep 26, 2011 | DS DP | U-550 | |
| <u>RALOXIFENE HYDROCHLORIDE - EVISTA</u> | | | | | |
| 020815 001 | RE38968 | Jul 28, 2012 | | U-662 | |
| | RE38968 | Jul 28, 2012 | | U-657 | |
| | RE39049 | Jul 28, 2012 | | U-662 | |
| | RE39049 | Jul 28, 2012 | | U-657 | |
| | RE39050 | Mar 02, 2014 | | U-662 | |
| | RE39050 | Mar 02, 2014 | | U-657 | |
| <u>RAMIPRIL - ALTACE</u> | | | | | |
| 019901 001 | 5061722 | Oct 19, 2008 | | | |
| <u>RANOLAZINE - RANEXA</u> | | | | | |
| 021526 002 | 4567264 | May 18, 2007 | DS | | Jan 27, 2011 |
| | 6303607 | May 27, 2019 | | U-705 | |
| | 6369062 | May 27, 2019 | | DP | |
| | 6479496 | May 27, 2019 | | U-705 | |
| | 6503911 | May 27, 2019 | | DP | |
| | 6525057 | May 27, 2019 | | U-705 | |
| | 6562826 | May 27, 2019 | | U-705 | |
| | 6617328 | May 27, 2019 | | DP | |
| | 6620814 | May 27, 2019 | | U-705 | |
| | 6852724 | May 27, 2019 | | U-705 | |
| | 6864258 | May 27, 2019 | | U-705 | |
| <u>RASAGILINE MESYLATE - AZILECT</u> | | | | | |
| 021641 001 | 5387612 | Feb 07, 2012 | | U-219 | May 16, 2011 |
| | 5453446 | Feb 07, 2012 | | U-219 | |
| | 5457133 | Feb 07, 2012 | DS DP | | |
| | 5532415 | Jul 02, 2013 | DS | | |
| | 5786390 | Feb 07, 2012 | | DP | |
| | 6126968 | Sep 18, 2016 | | DP | |
| <u>RASAGILINE MESYLATE - AZILECT</u> | | | | | |
| 021641 002 | 5387612 | Feb 07, 2012 | | U-219 | May 16, 2011 |
| | 5453446 | Feb 07, 2012 | | U-219 | |
| | 5457133 | Feb 07, 2012 | DS DP | | |
| | 5532415 | Jul 02, 2013 | DS | | |
| | 5786390 | Feb 07, 2012 | | DP | |
| | 6126968 | Sep 18, 2016 | | DP | |
| <u>RIFAXIMIN - XIFAXAN</u> | | | | | |
| 021361 001 | 7045620 | May 22, 2024 | DS | | |
| <u>RISEDRONATE SODIUM - ACTONEL</u> | | | | | |
| 020835 001 | | | | M-52 | Jan 24, 2009 |
| <u>RISEDRONATE SODIUM - ACTONEL</u> | | | | | |
| 020835 002 | | | | M-52 | Jan 24, 2009 |
| <u>RISEDRONATE SODIUM - ACTONEL</u> | | | | | |
| 020835 003 | >A> 5583122 | Dec 10, 2013 | DS DP | U-756 | Aug 11, 2009 |
| | >A> 5583122 | Dec 10, 2013 | DS DP | U-222 | Jan 24, 2009 |
| | >A> 6096342 | Nov 22, 2011 | | DP | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>RISPERIDONE - RISPERDAL</u> | | | | | |
| 020588 001 | RE39181 | Jul 11, 2014 | DP | | |
| <u>RIVASTIGMINE TARTRATE - EXELON</u> | | | | | |
| 020823 003 | 4948807 | Aug 14, 2012 | DS | U-322 | |
| <u>RIVASTIGMINE TARTRATE - EXELON</u> | | | | | |
| 020823 004 | 4948807 | Aug 14, 2012 | DS | U-322 | |
| <u>RIVASTIGMINE TARTRATE - EXELON</u> | | | | | |
| 020823 005 | 4948807 | Aug 14, 2012 | DS | U-322 | |
| <u>RIVASTIGMINE TARTRATE - EXELON</u> | | | | | |
| 020823 006 | 4948807 | Aug 14, 2012 | DS | U-322 | |
| <u>RIVASTIGMINE TARTRATE - EXELON</u> | | | | | |
| 021025 001 | 4948807 | Aug 14, 2012 | DS | U-322 | |
| <u>ROCURONIUM BROMIDE - ZEMURON</u> | | | | | |
| 020214 003 | 4894369 | Apr 13, 2008 | | | |
| <u>SALMETEROL XINAFOATE - SEREVENT</u> | | | | | |
| 020236 001 | 4992474 | Feb 12, 2008 | | | |
| | 4992474*PED | Aug 12, 2008 | | | |
| | 5126375 | Feb 12, 2008 | | | |
| | 5126375*PED | Aug 12, 2008 | | | |
| | 5225445 | Feb 12, 2008 | | U-182 | |
| | 5225445*PED | Aug 12, 2008 | | | |
| <u>SALMETEROL XINAFOATE - SEREVENT</u> | | | | | |
| 020692 001 | 4992474 | Feb 12, 2008 | | | |
| | 4992474*PED | Aug 12, 2008 | | | |
| | 5126375 | Feb 12, 2008 | | | |
| | 5126375*PED | Aug 12, 2008 | | | |
| | 5225445 | Feb 12, 2008 | | U-211 | |
| | 5225445*PED | Aug 12, 2008 | | | |
| <u>SELEGILINE - EMSAM</u> | | | | | |
| 021336 001 | 7070808 | May 10, 2018 | DS DP | NDF | Feb 27, 2009 |
| | RE34579 | Aug 18, 2007 | DS DP | U-711 | |
| <u>SELEGILINE - EMSAM</u> | | | | | |
| 021336 002 | RE34579 | Aug 18, 2007 | DS DP | U-711 | NDF Feb 27, 2009 |
| <u>SELEGILINE - EMSAM</u> | | | | | |
| 021336 003 | RE34579 | Aug 18, 2007 | DS DP | U-711 | NDF Feb 27, 2009 |
| <u>SELEGILINE HYDROCHLORIDE - ZELAPAR</u> | | | | | |
| 021479 001 | 5648093 | Jul 15, 2014 | DP | | |
| | 6423342 | Mar 01, 2016 | DP | | |
| <u>SERTRALINE HYDROCHLORIDE - SERTRALINE HYDROCHLORIDE</u> | | | | | |
| 075719 001 | | | | PC | Feb 06, 2007 |
| <u>SERTRALINE HYDROCHLORIDE - SERTRALINE HYDROCHLORIDE</u> | | | | | |
| 075719 002 | | | | PC | Feb 06, 2007 |
| <u>SERTRALINE HYDROCHLORIDE - SERTRALINE HYDROCHLORIDE</u> | | | | | |
| 075719 003 | | | | PC | Feb 06, 2007 |
| <u>SERTRALINE HYDROCHLORIDE - SERTRALINE HYDROCHLORIDE</u> | | | | | |
| 076934 001 | | | | PC | Feb 03, 2007 |
| <u>SERTRALINE HYDROCHLORIDE - ZOLOFT</u> | | | | | |
| 020990 001 | 6727283 | Oct 11, 2019 | DP | U-580 | |
| | 6727283*PED | Apr 11, 2020 | | | |
| | 7067555 | Nov 10, 2019 | DP | | |
| | 7067555*PED | May 10, 2020 | | | |
| <u>SEVELAMER HYDROCHLORIDE - RENAGEL</u> | | | | | |
| 021179 001 | 7014846 | Aug 11, 2013 | DP | U-246 | |
| <u>SEVELAMER HYDROCHLORIDE - RENAGEL</u> | | | | | |
| 021179 002 | 7014846 | Aug 11, 2013 | DP | U-246 | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-----------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>SIMVASTATIN - SIMVASTATIN</u> | | | | | |
| 076052 001 | | | | PC | Dec 20, 2006 |
| <u>SIMVASTATIN - SIMVASTATIN</u> | | | | | |
| 076052 002 | | | | PC | Dec 20, 2006 |
| <u>SIMVASTATIN - SIMVASTATIN</u> | | | | | |
| 076052 003 | | | | PC | Dec 20, 2006 |
| <u>SIMVASTATIN - SIMVASTATIN</u> | | | | | |
| 076052 004 | | | | PC | Dec 20, 2006 |
| <u>SIMVASTATIN - SIMVASTATIN</u> | | | | | |
| 076285 005 | | | | PC | Dec 20, 2006 |
| <u>SIMVASTATIN - ZOCOR</u> | | | | | |
| 019766 001 | RE36481 *** | Jul 10, 2007 | | U-300 | |
| | RE36481*PED | Jan 10, 2008 | | U-300 | |
| | RE36520 *** | May 26, 2009 | | U-300 | |
| | RE36520*PED | Nov 26, 2009 | | U-300 | |
| <u>SIMVASTATIN - ZOCOR</u> | | | | | |
| 019766 002 | RE36481 *** | Jul 10, 2007 | | U-300 | |
| | RE36481*PED | Jan 10, 2008 | | U-300 | |
| | RE36520 *** | May 26, 2009 | | U-300 | |
| | RE36520*PED | Nov 26, 2009 | | U-300 | |
| <u>SIMVASTATIN - ZOCOR</u> | | | | | |
| 019766 003 | RE36481 *** | Jul 10, 2007 | | U-300 | |
| | RE36481*PED | Jan 10, 2008 | | U-300 | |
| | RE36520 *** | May 26, 2009 | | U-300 | |
| | RE36520*PED | Nov 26, 2009 | | U-300 | |
| <u>SIMVASTATIN - ZOCOR</u> | | | | | |
| 019766 004 | RE36481 *** | Jul 10, 2007 | | U-300 | |
| | RE36481*PED | Jan 10, 2008 | | U-300 | |
| | RE36520 *** | May 26, 2009 | | U-300 | |
| | RE36520*PED | Nov 26, 2009 | | U-300 | |
| <u>SIMVASTATIN - ZOCOR</u> | | | | | |
| 019766 005 | RE36481 *** | Jul 10, 2007 | | U-300 | |
| | RE36481*PED | Jan 10, 2008 | | U-300 | |
| | RE36520 *** | May 26, 2009 | | U-300 | |
| | RE36520*PED | Nov 26, 2009 | | U-300 | |
| <u>SIROLIMUS - RAPAMUNE</u> | | | | | |
| 021083 001 | >A> 5100899 | Jul 07, 2013 | | U-290 | |
| | >A> 5100899*PED | Jan 07, 2014 | | | |
| <u>SIROLIMUS - RAPAMUNE</u> | | | | | |
| 021110 001 | >A> 5100899 | Jul 07, 2013 | | U-290 | |
| | >A> 5100899*PED | Jan 07, 2014 | | | |
| <u>SIROLIMUS - RAPAMUNE</u> | | | | | |
| 021110 002 | >A> 5100899 | Jul 07, 2013 | | U-290 | |
| | >A> 5100899*PED | Jan 07, 2014 | | | |
| <u>SIROLIMUS - RAPAMUNE</u> | | | | | |
| 021110 003 | >A> 5100899 | Jul 07, 2013 | | U-290 | |
| | >A> 5100899*PED | Jan 07, 2014 | | | |
| <u>SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE - OSMOPREP</u> | | | | | |
| 021892 001 | 5616346 | May 18, 2013 | DP | U-715 NP | Mar 16, 2009 |
| <u>SOMATROPIN RECOMBINANT - GENOTROPIN</u> | | | | | |
| 020280 006 | 4968299 | Jun 28, 2008 | DP | I-496 | Apr 27, 2009 |
| <u>SOMATROPIN RECOMBINANT - GENOTROPIN</u> | | | | | |
| 020280 007 | 4968299 | Jun 28, 2008 | DP | I-496 | Apr 27, 2009 |
| <u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u> | | | | | |
| 020280 001 | 5435076 | Apr 16, 2013 | DP | I-496 | Apr 27, 2009 |
| | 5716338 | Feb 10, 2015 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-------------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u> | | | | | |
| 020280 002 | 5435076 | Apr 16, 2013 | DP | I-496 | Apr 27, 2009 |
| | 5716338 | Feb 10, 2015 | DP | | |
| <u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u> | | | | | |
| 020280 003 | 5435076 | Apr 16, 2013 | DP | I-496 | Apr 27, 2009 |
| | 5716338 | Feb 10, 2015 | DP | | |
| <u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u> | | | | | |
| 020280 004 | | | | I-496 | Apr 27, 2009 |
| <u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u> | | | | | |
| 020280 005 | 5435076 | Apr 16, 2013 | DP | I-496 | Apr 27, 2009 |
| | 5716338 | Feb 10, 2015 | DP | | |
| <u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u> | | | | | |
| 020280 008 | 5435076 | Apr 16, 2013 | DP | I-496 | Apr 27, 2009 |
| | 5716338 | Feb 10, 2015 | DP | | |
| <u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u> | | | | | |
| 020280 009 | 5435076 | Apr 16, 2013 | DP | I-496 | Apr 27, 2009 |
| | 5716338 | Feb 10, 2015 | DP | | |
| <u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u> | | | | | |
| 020280 010 | 5435076 | Apr 16, 2013 | DP | I-496 | Apr 27, 2009 |
| | 5716338 | Feb 10, 2015 | DP | | |
| <u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u> | | | | | |
| 020280 011 | 5435076 | Apr 16, 2013 | DP | I-496 | Apr 27, 2009 |
| | 5716338 | Feb 10, 2015 | DP | | |
| <u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u> | | | | | |
| 020280 012 | 5435076 | Apr 16, 2013 | DP | I-496 | Apr 27, 2009 |
| | 5716338 | Feb 10, 2015 | DP | | |
| <u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u> | | | | | |
| 020280 013 | 5435076 | Apr 16, 2013 | DP | I-496 | Apr 27, 2009 |
| | 5716338 | Feb 10, 2015 | DP | | |
| <u>SOMATROPIN RECOMBINANT - OMNITROPE</u> | | | | | |
| 021426 001 | | | | NP | May 30, 2009 |
| <u>SOMATROPIN RECOMBINANT - OMNITROPE</u> | | | | | |
| 021426 002 | | | | NP | May 30, 2009 |
| <u>SORAFENIB TOSYLATE - NEXAVAR</u> | | | | | |
| 021923 001 | | | | ODE | Dec 20, 2012 |
| <u>SUMATRIPTAN SUCCINATE - IMITREX STATDOSE</u> | | | | | |
| 020080 003 | 4816470 | Dec 28, 2006 | | U-72 | |
| | 4816470*PED | Jun 28, 2007 | | | |
| | 5037845 | Aug 06, 2008 | | U-72 | |
| | 5037845*PED | Feb 06, 2009 | | | |
| <u>SUNITINIB MALATE - SUTENT</u> | | | | | |
| 021938 001 | 6573293 | Feb 15, 2021 | DS DP | U-703 | Jan 26, 2011 |
| <u>SUNITINIB MALATE - SUTENT</u> | | | | | |
| 021938 002 | 6573293 | Feb 15, 2021 | DS DP | U-703 | Jan 26, 2011 |
| <u>SUNITINIB MALATE - SUTENT</u> | | | | | |
| 021938 003 | 6573293 | Feb 15, 2021 | DS DP | U-703 | Jan 26, 2011 |
| <u>TACROLIMUS - PROGRAF</u> | | | | | |
| 050708 001 | | | | ODE | Mar 29, 2013 |
| <u>TACROLIMUS - PROGRAF</u> | | | | | |
| 050708 002 | | | | ODE | Mar 29, 2013 |
| <u>TACROLIMUS - PROGRAF</u> | | | | | |
| 050708 003 | | | | ODE | Mar 29, 2013 |
| <u>TACROLIMUS - PROGRAF</u> | | | | | |
| 050709 001 | | | | ODE | Mar 29, 2013 |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-----------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>TERIPARATIDE RECOMBINANT HUMAN - FORTEO</u> | | | | | |
| 021318 001 | 6977077 | Aug 19, 2019 | | U-597 | |
| <u>THALIDOMIDE - THALOMID</u> | | | | | |
| 020785 001 | 5629327 | Mar 01, 2013 | | U-731 | |
| | 6045501 | Aug 28, 2018 | | U-731 | |
| | 6045501 | Aug 28, 2018 | | U-371 | |
| | 6235756 | Mar 01, 2013 | | U-731 | |
| | 6315720 | Oct 23, 2020 | | U-442 | |
| | 6315720 | Oct 23, 2020 | | U-731 | |
| | 6561976 | Aug 28, 2018 | | U-731 | |
| | 6561976 | Aug 28, 2018 | | U-371 | |
| | 6561977 | Oct 23, 2020 | | U-371 | |
| | 6561977 | Oct 23, 2020 | | U-731 | |
| | 6755784 | Sep 23, 2020 | | U-731 | |
| | 6755784 | Sep 23, 2020 | | U-371 | |
| | 6869399 | Oct 23, 2020 | | U-371 | |
| | 6869399 | Oct 23, 2020 | | U-732 | |
| | 6869399 | Oct 23, 2020 | | U-733 | |
| | 6869399 | Oct 23, 2020 | | U-731 | |
| | 6908432 | Aug 28, 2018 | | U-371 | |
| | 6908432 | Aug 28, 2018 | | U-731 | |
| <u>THALIDOMIDE - THALOMID</u> | | | | | |
| 020785 002 | 5629327 | Mar 01, 2013 | | U-731 | |
| | 6045501 | Aug 28, 2018 | | U-731 | |
| | 6045501 | Aug 28, 2018 | | U-371 | |
| | 6235756 | Mar 01, 2013 | | U-731 | |
| | 6315720 | Oct 23, 2020 | | U-442 | |
| | 6315720 | Oct 23, 2020 | | U-731 | |
| | 6561976 | Aug 28, 2018 | | U-731 | |
| | 6561976 | Aug 28, 2018 | | U-371 | |
| | 6561977 | Oct 23, 2020 | | U-371 | |
| | 6561977 | Oct 23, 2020 | | U-731 | |
| | 6755784 | Sep 23, 2020 | | U-731 | |
| | 6755784 | Sep 23, 2020 | | U-371 | |
| | 6869399 | Oct 23, 2020 | | U-371 | |
| | 6869399 | Oct 23, 2020 | | U-733 | |
| | 6869399 | Oct 23, 2020 | | U-732 | |
| | 6869399 | Oct 23, 2020 | | U-731 | |
| | 6908432 | Aug 28, 2018 | | U-371 | |
| | 6908432 | Aug 28, 2018 | | U-731 | |
| <u>THALIDOMIDE - THALOMID</u> | | | | | |
| 020785 003 | 5629327 | Mar 01, 2013 | | U-731 | |
| | 6045501 | Aug 28, 2018 | | U-731 | |
| | 6045501 | Aug 28, 2018 | | U-371 | |
| | 6235756 | Mar 01, 2013 | | U-731 | |
| | 6315720 | Oct 23, 2020 | | U-442 | |
| | 6315720 | Oct 23, 2020 | | U-731 | |
| | 6561976 | Aug 28, 2018 | | U-731 | |
| | 6561976 | Aug 28, 2018 | | U-371 | |
| | 6561977 | Oct 23, 2020 | | U-371 | |
| | 6561977 | Oct 23, 2020 | | U-731 | |
| | 6755784 | Sep 23, 2020 | | U-731 | |
| | 6755784 | Sep 23, 2020 | | U-371 | |
| | 6869399 | Oct 23, 2020 | | U-371 | |
| | 6869399 | Oct 23, 2020 | | U-732 | |
| | 6869399 | Oct 23, 2020 | | U-733 | |
| | 6869399 | Oct 23, 2020 | | U-731 | |
| | 6908432 | Aug 28, 2018 | | U-371 | |
| | 6908432 | Aug 28, 2018 | | U-731 | |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--|-----------|---------------------------|-----------------|-------|------------------------|--------------------------------|
| <u>THYROTROPIN ALFA - THYROGEN</u> | | | | | | |
| 020898 001 | | | | | M-53 | Jan 23, 2009 |
| <u>TIOTROPIUM BROMIDE MONOHYDRATE - SPIRIVA</u> | | | | | | |
| 021395 001 | 7070800 | Jan 22, 2022 | DP | U-566 | | |
| | RE38912 | Oct 11, 2021 | DP | | | |
| <u>TOPIRAMATE - TOPAMAX</u> | | | | | | |
| 020505 001 | 7018983 | Oct 13, 2015 | | U-723 | | |
| <u>TOPIRAMATE - TOPAMAX</u> | | | | | | |
| 020505 002 | 7018983 | Oct 13, 2015 | | U-723 | | |
| <u>TOPIRAMATE - TOPAMAX</u> | | | | | | |
| 020505 003 | 7018983 | Oct 13, 2015 | | U-723 | | |
| <u>TOPIRAMATE - TOPAMAX</u> | | | | | | |
| 020505 004 | 7018983 | Oct 13, 2015 | | U-723 | | |
| <u>TOPIRAMATE - TOPAMAX</u> | | | | | | |
| 020505 005 | 7018983 | Oct 13, 2015 | | U-723 | | |
| <u>TOPIRAMATE - TOPAMAX</u> | | | | | | |
| 020505 006 | 7018983 | Oct 13, 2015 | | U-723 | | |
| <u>TOPIRAMATE - TOPAMAX SPRINKLE</u> | | | | | | |
| 020844 001 | 7018983 | Oct 13, 2015 | | U-723 | | |
| <u>TOPIRAMATE - TOPAMAX SPRINKLE</u> | | | | | | |
| 020844 002 | 7018983 | Oct 13, 2015 | | U-723 | | |
| <u>TOPIRAMATE - TOPAMAX SPRINKLE</u> | | | | | | |
| 020844 003 | 7018983 | Oct 13, 2015 | | U-723 | | |
| <u>TOPOTECAN HYDROCHLORIDE - Hycamtin</u> | | | | | | |
| 020671 001 | 5004758 | May 28, 2010 | DS DP | U-741 | | |
| <u>TREPROSTINIL SODIUM - REMODULIN</u> | | | | | | |
| 021272 001 | 5153222 | Oct 06, 2014 | | U-455 | | |
| <u>TREPROSTINIL SODIUM - REMODULIN</u> | | | | | | |
| 021272 002 | 5153222 | Oct 06, 2014 | | U-455 | | |
| <u>TREPROSTINIL SODIUM - REMODULIN</u> | | | | | | |
| 021272 003 | 5153222 | Oct 06, 2014 | | U-455 | | |
| <u>TREPROSTINIL SODIUM - REMODULIN</u> | | | | | | |
| 021272 004 | 5153222 | Oct 06, 2014 | | U-455 | | |
| <u>URSODIOL - URSO FORTE</u> | | | | | | |
| 020675 002 | 4859660 | Nov 19, 2007 | | U-740 | | |
| <u>VARENICLINE TARTRATE - CHANTIX</u> | | | | | | |
| 021928 001 | 6410550 | Nov 13, 2018 | DS DP | U-56 | NCE | May 10, 2011 |
| | 6890927 | May 06, 2022 | DS DP | U-56 | | |
| <u>VARENICLINE TARTRATE - CHANTIX</u> | | | | | | |
| 021928 002 | 6410550 | Nov 13, 2018 | DS DP | U-56 | NCE | May 10, 2011 |
| | 6890927 | May 06, 2022 | DS DP | U-56 | | |
| <u>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u> | | | | | | |
| 076690 001 | | | | | PC | Jan 30, 2007 |
| <u>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u> | | | | | | |
| 076690 002 | | | | | PC | Jan 30, 2007 |
| <u>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u> | | | | | | |
| 076690 003 | | | | | PC | Jan 30, 2007 |
| <u>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u> | | | | | | |
| 076690 004 | | | | | PC | Jan 30, 2007 |
| <u>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u> | | | | | | |
| 076690 005 | | | | | PC | Jan 30, 2007 |

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---|-----------|---------------------------|-----------------|------------------------|--------------------------------|
| <u>ZANAMIVIR - RELENZA</u> | | | | | |
| 021036 001 | 5648379 | Jul 15, 2014 | U-722 | I-491 | Mar 29, 2009 |
| | 5648379 | Jul 15, 2014 | U-721 | | |
| | 5648379 | Jul 15, 2014 | U-274 | | |
| | 6294572 | Dec 15, 2014 | DS DP | | |
| <u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u> | | | | | |
| 020825 001 | 4831031 | Mar 02, 2012 | DS DP U-720 | I-492 | Aug 19, 2007 |
| <u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u> | | | | | |
| 020825 002 | 4831031 | Mar 02, 2012 | DS DP U-720 | I-492 | Aug 19, 2007 |
| <u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u> | | | | | |
| 020825 003 | 4831031 | Mar 02, 2012 | DS DP U-720 | I-492 | Aug 19, 2007 |
| <u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u> | | | | | |
| 020825 004 | 4831031 | Mar 02, 2012 | DS DP U-720 | I-492 | Aug 19, 2007 |
| <u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u> | | | | | |
| 021483 001 | 4831031 | Mar 02, 2012 | DS DP U-720 | I-492 | Aug 19, 2007 |
| | 5312925 | Sep 01, 2012 | DS DP U-720 | | |
| | 6150366 | May 27, 2019 | DP U-719 | | |
| | 6245766 | Dec 18, 2018 | U-601 | | |
| <u>ZIPRASIDONE MESYLATE - GEODON</u> | | | | | |
| 020919 001 | 4831031 | Mar 02, 2012 | DS DP U-720 | | |
| <u>ZOLEDRONIC ACID - ZOMETA</u> | | | | | |
| 021223 001 | 4939130 | Sep 02, 2012 | DS DP U-53 | | |
| <u>ZOLEDRONIC ACID - ZOMETA</u> | | | | | |
| 021223 002 | 4939130 | Sep 02, 2012 | DS DP U-53 | | |

Footnotes:

- Patents are published upon receipt by the Orange Book Staff and may not reflect the official receipt date as described in 21 CFR 314.53(d)(5).
- Patents submitted on FDA Form 3542 and listed after August 18, 2003 will have one to three patent codes indicating specific patent claims as submitted by the sponsor:
DS = Drug Substance claim
DP = Drug Product claim
U and number = Method of Use claim (may be multiple). Specific Method of use claims are listed at <http://www.fda.gov/cder/orange/patex.htm>
- Patents listed prior to August 18, 2003 are flagged with method of use claims only as applicable and submitted by the sponsor. They may not be flagged with respect to other claims which may apply.
- *PED and PED represent pediatric exclusivity. Patents with pediatric exclusivity granted after August 18, 2003 will be indicated with *PED as was done prior to August 18, 2003. Patents with *PED added after August 18, 2003 will not contain any information relative to the patent itself other than the *PED extension. Information related specifically to the patent will be conveyed on the original patent only.
- *** U.S. Patent Nos. RE 36481 and RE 36520 are being relisted for Zocor (NDA 19-766) pursuant to the decision and related order in Ranbaxy Labs. v. Leavitt, No. 05-1838 (D.D.C. April 30, 2006). The '481 and '520 patents will remain listed in Approved Drug Products with Therapeutic Equivalence Evaluations until any applicable periods of exclusivity pursuant to section 505(j)(5)(B)(iv) of the Federal Food, Drug, and Cosmetic Act have been triggered and run, unless the agency's appeal of the decision to the U.S. Court of Appeals for the District of Columbia is decided in the agency's favor before the exclusivity periods have expired. While the patents remain listed, any new or pending ANDA referencing Zocor must contain patent certifications to these patents. For additional information on this matter, please refer to Docket Nos. 2005P-0008 and 2005P-0046.

PATENT AND EXCLUSIVITY TERMS

Due to space limitations in the patent and exclusivity columns, abbreviations and references have been developed. Refer to the APPROVED DRUG PRODUCTS WITH THERAPEUTIC EQUIVALENCE EVALUATIONS, 25th Edition for a full listing of patent and exclusivity terms (Abbreviations, Dosing Schedule, Indications, and Patent Use Codes).

The current complete list of patent terms is available at <http://www.accessdata.fda.gov/scripts/cder/ob/docs/pattermsall.cfm>

The current complete list of exclusivity terms is available at <http://www.accessdata.fda.gov/scripts/cder/ob/docs/excltermsall.cfm>