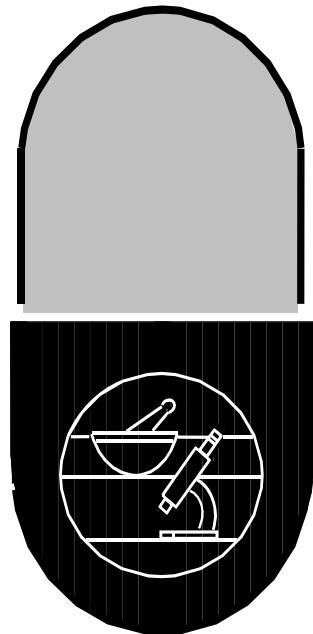


**CUMULATIVE
SUPPLEMENT 7
July 2012**



**APPROVED
DRUG PRODUCTS**

**WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS**

32nd EDITION

**Department of Health and Human Services
Food and Drug Administration
Center for Drug Evaluation and Research
Office of Generic Drugs**

2012

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

**APPROVED DRUG PRODUCTS
with
THERAPEUTIC EQUIVALENCE EVALUATIONS**

32nd EDITION

Cumulative Supplement 7

July 2012

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 Levothyroxine Sodium.....	v
1.5 Availability of the Edition	vi
1.6 Report of Counts for the Prescription Drug Product List	vii
1.7 Cumulative Supplement Legend	viii
 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 <i>in vivo</i> 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**

32nd EDITION

**CUMULATIVE SUPPLEMENT 7
July 2012**

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, 30th 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are for exportation, are for military use, have been discontinued from marketing or that have had their approvals withdrawn for other than safety or efficacy reasons.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 32nd Edition List will then be added to the "Discontinued Drug Product List" appearing in the 33rd 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.
- All product changes received and processed as of the date of publication.
 - o Refer to CS Section 1.8 Cumulative Supplement Legend for types of changes
 - o 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 Staff received a letter November 7 that the product has been discontinued from manufacturing and marketing. The Orange Book 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 Staff 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.

Every effort is made to ensure the Cumulative Supplement is current and accurate. Applicant holders are requested to inform the FDA Orange Book Staff (OBS) of any changes or corrections. The OBS can be contacted by email at drugproducts@fda.hhs.gov. Send Changes by FAX: 240-276-8974; mail to:

FDA/CDER Orange Book Staff
Office of Generic Drugs, HFD-610
7620 Standish Place
Rockville, MD 20855-2773

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.

FORMER APPLICANT NAME
(FORMER ABBREVIATED NAME)

ALCON UNIVERSAL LTD
(ALCON UNIVERSAL)

OVATION PHARMACEUTICALS INC
(OVATION PHARMS)

NEW APPLICANT NAME
(NEW ABBREVIATED NAME)

ALCON PHARMACEUTICALS LTD
(ALCON PHARMS LTD)

OAK PHARMACEUTICALS INC
(OAK PHARMS)

1.4 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) and Levo-T (Alara NDA 21342) 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 (Merck KGAA 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 (Merck KGAA ANDA 76752) tablets have been determined to be therapeutically equivalent to corresponding strengths of Levoxyl (King Pharms NDA 021301) tablets.

Levothyroxine Sodium (Mylan ANDA 76187) tablets have been determined to be therapeutically equivalent to corresponding strengths of Levothroid (Lloyd NDA 021116) tablets.

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

Trade Name	Applicant	Potency	TE Code	Appl No	Product No
UNITHROID	STEVENS J	0.025MG	AB1	21210	001
LEVOTHYROXINE SODIUM	MYLAN	0.025MG	AB1	76187	001
LEVOXYL	KING PHARMS	0.025MG	AB1	21301	001
SYNTHROID	ABBOTT	0.025MG	AB1	21402	001
LEVO-T	ALARA PHARM	0.025MG	AB1	21342	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	MERCK KGAA	0.025MG	AB2	76752	001
LEVOXYL	KUNG PHARMS	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	MERCK KGAA	0.025MG	AB3	76752	001
LEVOTHROID	LLOYD	0.025MG	AB4	21116	001
LEVOTHYROXINE SODIUM	MYLAN	0.025MG	AB4	76187	001

1.5 AVAILABILITY OF THE EDITION

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

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

The downloaded Annual Edition and Cumulative Supplements are also available in a paper version (Approved Drug Products with Therapeutic Equivalence Evaluations, ADP) from the U.S. Government Printing Office: <http://bookstore.gpo.gov>; toll free 866-512-1800.

There are historical lists of Orange Book cumulative supplement product monthly changes at

<http://www.fda.gov/Drugs/InformationOnDrugs/ucm086229.htm>. There are ASCII text files of the Orange Book drug product, patent, and exclusivity data at <http://www.fda.gov/Drugs/InformationOnDrugs/ucm129689.htm>. The drug product text files are provided in eobzip.zip format. The files are updated concurrently with the monthly cumulative supplements. The annual Orange Book Edition Appendices A, B, and C in PDF format 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>.

1.6 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 2011) 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

CATEGORIES COUNTED	DEC 2011	MAR 2012	JUN 2012	SEPT 2012	DEC 2012
DRUG PRODUCTS LISTED	14480	14711	14834		
SINGLE SOURCE	2451 (16.9%)	2446 (16.6%)	2428 (16.4%)		
MULTISOURCE	11953 (82.5%)	12187 (82.8%)	12328 (83.1%)		
THERAPEUTICALLY EQUIVALENT	11792 (81.4%)	12037 (81.8%)	12187 (82.2%)		
NOT THERAPEUTICALLY EQUIVALENT	161 (1.1%)	150 (1.0%)	141 (1.0%)		
EXCEPTIONS ¹	76 (0.5%)	78 (0.5%)	78 (0.5%)		
NEW MOLECULAR ENTITIES					
APPROVED	6	8	7		
NUMBER OF APPLICANTS	810	826	826		

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

1.7 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 application number preceded by "N" is a New Drug Application (NDA or innovator). The application number preceded by an "A" is an Abbreviated New Drug Application (ANDA or generic). 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 proceeded 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 - 32ND EDITION

RX DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

1-1

ABACAVIR SULFATE

TABLET; ORAL

ABACAVIR SULFATE

AB	MYLAN PHARMS INC	EQ 300MG BASE	A091294 001 Jun 18, 2012 Jun NEWA
	ZIAGEN		
AB	+ VIIV HLTHCARE	EQ 300MG BASE	N020977 001 Dec 17, 1998 Jun CFTG

ACARBOSE

TABLET; ORAL

ACARBOSE

AB	EMCURE PHARMS LTD	25MG	A202271 001 Feb 07, 2012 Jan NEWA
AB		50MG	A202271 002 Feb 07, 2012 Jan NEWA
AB		100MG	A202271 003 Feb 07, 2012 Jan NEWA

ACETAMINOPHEN; BUTALBITAL

TABLET; ORAL

BUTAPAP

+ MIKART

325MG;50MG

A089987 001 Oct 26, 1992 Jun CRLD

PHRENILIN

@ VALEANT

325MG;50MG

A087811 001 Jun 19, 1985 Jun DISC

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE

>A>	NEXGEN PHARMA INC	300MG;50MG;40MG;30MG	A076560 002 Jul 19, 2012 Jul NEWA
-----	-------------------	----------------------	-----------------------------------

ACETAMINOPHEN; CODEINE PHOSPHATE

SOLUTION; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

AA	+ PHARM ASSOC	120MG/5ML;12MG/5ML	A087508 001 Jan CRLD
	SUSPENSION; ORAL		
	CAPITAL AND CODEINE		
+ VALEANT	120MG/5ML;12MG/5ML	A086024 001 Jan CTEC	
+ VALEANT PHARMS LLC	120MG/5ML;12MG/5ML	A086024 001 Jun CAHN	

ACETAMINOPHEN; HYDROCODONE BITARTRATE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

@ NESHER PHARMS 500MG/15ML;7.5MG/15ML

A040366 001 Jan 23, 2002 Jun DISC

AA	VISTAPHARM	325MG/15ML;7.5MG/15ML
----	------------	-----------------------

A200343 001 Jan 25, 2012 Jan NEWA

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

AA	AUROLIFE PHARMA LLC	325MG;5MG
AA		325MG;7.5MG
AA		325MG;10MG

A201013 001 Apr 11, 2012 Mar NEWA

A201013 002 Apr 11, 2012 Mar NEWA

A201013 003 Apr 11, 2012 Mar NEWA

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE AND ACETAMINOPHEN

AA	ACTAVIS TOTOWA	325MG;7.5MG
AA		325MG;10MG
>A>	AA ALVOGEN INC	325MG;7.5MG
>A>	AA	325MG;10MG

A040800 001 Apr 03, 2012 Mar NEWA

A040800 002 Apr 03, 2012 Mar NEWA

A202677 001 Jul 26, 2012 Jul NEWA

A202677 002 Jul 26, 2012 Jul NEWA

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

AB ALVOGEN INC 325MG;37.5MG A202076 001 Mar 30, 2012 Mar NEWA

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

AP SAGENT STRIDES EQ 500MG BASE/VIAL A200880 001 May 09, 2012 Apr NEWA

AP + X GEN PHARMS EQ 500MG BASE/VIAL A040784 001 Dec 10, 2008 Apr CRLD

DIAMOX

@ TEVA WOMENS EQ 500MG BASE/VIAL N009388 001 Dec 05, 1990 Apr DISC

ACETYLCYSTEINE

SOLUTION; INHALATION, ORAL

ACETYLCYSTEINE

AN INNOPHARMA INC 20% A203853 001 Jun 21, 2012 Jun NEWA

>A> ACLIDINIUM BROMIDE

>A> POWDER, METERED; INHALATION

>A> TUDORZA PRESSAIR

>A> + FOREST LABS INC 0.375MG/INH N202450 001 Jul 23, 2012 Jul NEWA

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

@ HERITAGE PHARMS INC 200MG

A074889 001 Oct 31, 1997 May CAHN

TABLET; ORAL

ACYCLOVIR

@ HERITAGE PHARMS INC 400MG

A074891 001 Oct 31, 1997 May CAHN

@ 800MG

A074891 002 Oct 31, 1997 May CAHN

ADAPALENE

CREAM; TOPICAL

ADAPALENE

AB FOUGERA PHARMS 0.1% A090824 001 Jun 30, 2010 Jan CAHN

GEL; TOPICAL

ADAPALENE

AB TOLMAR 0.3% A200298 001 Jun 14, 2012 Jun NEWA

DIFFERIN

AB + GALDERMA LABS LP 0.3% N021753 001 Jun 19, 2007 Jun CFTG

LOTION; TOPICAL

DIFFERIN

+ GALDERMA LABS LP 0.1%

N022502 001 Mar 17, 2010 Apr CAHN

ADENOSINE

INJECTABLE; INJECTION

ADENOSINE

@ TEVA PARENTERAL 3MG/ML

A078676 001 Jul 31, 2008 Mar DISC

@ 3MG/ML

A076564 001 Jun 16, 2004 Jan DISC

ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

SOLUTION; INHALATION

ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE

@ SANDOZ INC EQ 0.083% BASE;0.017%

A076867 001 Dec 21, 2006 Apr DISC

ALCLOMETASONE DIPROPIONATE

CREAM; TOPICAL
 ALCLOMETASONE DIPROPIONATE
 AB FOUGERA PHARMS 0.05% A076973 001 Jul 12, 2005 Jan CAHN
 OINTMENT; TOPICAL
 ALCLOMETASONE DIPROPIONATE
 AB FOUGERA PHARMS 0.05% A076884 001 Jul 18, 2005 Jan CAHN

ALENDRONATE SODIUM

TABLET, EFFERVESCENT; ORAL
 BINOSTO
 + MISSION PHARMA EQ 70MG BASE N202344 001 Mar 12, 2012 Mar NEWA

ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL
 ALFUZOSIN HYDROCHLORIDE
 AB INVAGEN PHARMS 10MG A090284 001 Jan 17, 2012 Jan NEWA
 >A> AB WOCKHARDT LTD 10MG A090221 001 Aug 10, 2012 Jul NEWA

ALGLUCERASE

INJECTABLE; INJECTION
 CEREDASE
 @ GENZYME 80 UNITS/ML N020057 003 Apr 05, 1991 Mar DISC

ALPRAZOLAM

TABLET, EXTENDED RELEASE; ORAL
 ALPRAZOLAM
 @ SANDOZ INC 0.5MG A077777 001 Jun 30, 2006 Mar DISC
 @ 1MG A077777 002 Jun 30, 2006 Mar DISC
 @ 2MG A077777 003 Jun 30, 2006 Mar DISC
 @ 3MG A077777 004 Jun 30, 2006 Mar DISC
 >D> AB VINTAGE 0.5MG A078442 001 Oct 15, 2007 Jul DISC
 >D> AB 1MG A078442 002 Oct 15, 2007 Jul DISC
 >D> AB 2MG A078442 003 Oct 15, 2007 Jul DISC
 >D> AB 3MG A078442 004 Oct 15, 2007 Jul DISC
 >A> @ VINTAGE PHARMS 0.5MG A078442 001 Oct 15, 2007 Jul DISC
 >A> @ 1MG A078442 002 Oct 15, 2007 Jul DISC
 >A> @ 2MG A078442 003 Oct 15, 2007 Jul DISC
 >A> @ 3MG A078442 004 Oct 15, 2007 Jul DISC

TABLET, ORALLY DISINTEGRATING; ORAL
 NIRAVAM

AB UCB INC	0.25MG	N021726 001 Jan 19, 2005 Jan CAHN
AB	0.5MG	N021726 002 Jan 19, 2005 Jan CAHN
AB +	1MG	N021726 003 Jan 19, 2005 Jan CAHN
AB	2MG	N021726 004 Jan 19, 2005 Jan CAHN

ALVIMOPAN

CAPSULE; ORAL
 ENTEREG
 + CUBIST PHARMS 12MG N021775 001 May 20, 2008 Feb CAHN

AMCINONIDE

CREAM; TOPICAL
 AMCINONIDE
 AB + FOUGERA PHARMS 0.1% A076065 001 May 15, 2003 Jan CAHN

LOTION; TOPICAL

AMCINONIDE

+ FOUGERA PHARMS 0.1% A076329 001 Nov 06, 2002 Jan CAHN

OINTMENT; TOPICAL

AMCINONIDE

AB + FOUGERA PHARMS 0.1% A076096 001 Nov 19, 2002 Jan CAHN

AMIKACIN SULFATEINJECTABLE; INJECTION

AMIKACIN SULFATE

@ HOSPIRA

EQ 50MG BASE/ML

A063263 001 Nov 30, 1994 Jun DISC

AMINO ACIDSINJECTABLE; INJECTION

AMINO ACIDS

B BRAUN 15%(300GM/2000ML)

A091112 002 Apr 13, 2012 Mar NEWA

15%(150GM/1000ML)

A091112 001 Apr 13, 2012 Mar NEWA

CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER

BAXTER HLTHCARE 15% (15GM/100ML)

A020512 001 Aug 30, 1996 Jun CAHN

NOVAMINE 15%

@ HOSPIRA INC

15% (75GM/500ML)

N017957 004 Nov 28, 1986 Mar CPOT

AMINOCAPROIC ACIDINJECTABLE; INJECTION

AMICAR

@ CLOVER PHARMS

250MG/ML

N015229 002

Mar CAHN

SYRUP; ORAL

AMICAR

AA + CLOVER PHARMS 1.25GM/5ML

N015230 002

Mar CAHN

TABLET; ORAL

AMICAR

AB CLOVER PHARMS 500MG

N015197 001

Mar CAHN

+ 1GM

N015197 002 Jun 24, 2004 Mar CAHN

AMIODARONE HYDROCHLORIDEINJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

@ BEDFORD 50MG/ML

A076018 001 Oct 15, 2002 Jun DISC

@ BEDFORD LABS 50MG/ML

A076299 001 Oct 24, 2002 Jun DISC

@ TEVA PARENTERAL 50MG/ML

A076163 001 Sep 05, 2003 Jan DISC

TABLET; ORAL

AMIODARONE HYDROCHLORIDE

AB MURTY PHARMS 200MG

A077069 001 Apr 08, 2005 May CAHN

AB 400MG

A077069 002 Apr 08, 2005 May CAHN

AMITRIPTYLINE HYDROCHLORIDETABLET; ORAL

AMITRIPTYLINE HYDROCHLORIDE

AB CARACO 10MG

A040816 002 Jun 27, 2008 Apr NEWA

AB 50MG

A040816 003 Jun 27, 2008 Apr NEWA

AB 75MG

A040816 004 Jun 27, 2008 Apr NEWA

AB 100MG

A040816 005 Jun 27, 2008 Apr NEWA

AB 150MG

A040816 006 Jun 27, 2008 Apr NEWA

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TABLET; ORAL

CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE

MYLAN PHARMS INC	EQ 12.5MG BASE;5MG	A071297 002	Dec 10, 1986	Feb	CTEC
+	EQ 25MG BASE;10MG	A071297 001	Dec 10, 1986	Feb	CRLD
LIMBITROL					
@ VALEANT PHARM INTL	EQ 12.5MG BASE;5MG	N016949 001		Feb	DISC
LIMBITROL DS					
@ VALEANT PHARM INTL	EQ 25MG BASE;10MG	N016949 002		Feb	DISC

AMLODIPINE BESYLATE

TABLET; ORAL

AMLODIPINE BESYLATE

AB	MACLEODS PHARMS LTD	EQ 5MG BASE	A201380 001	Apr 13, 2012	Mar	NEWA
AB		EQ 10MG BASE	A201380 002	Apr 13, 2012	Mar	NEWA

AMOXICILLIN

TABLET, FOR SUSPENSION; ORAL

DISPERMOX

@ RANBAXY LABS LTD 600MG

A065159 001 Dec 04, 2003 Apr DISC

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

>D>	AB	LEK PHARMS	200MG/5ML;EQ 28.5MG BASE/5ML	A065098 001	Dec 16, 2002	Jul	CAHN
>D>	AB		400MG/5ML;EQ 57MG BASE/5ML	A065098 002	Dec 16, 2002	Jul	CAHN
>D>	AB	LEK PHARMS DD	600MG/5ML;EQ 42.9MG BASE/5ML	A065358 001	Aug 13, 2007	Jul	CAHN
>A>	AB	SANDOZ INC	200MG/5ML;EQ 28.5MG BASE/5ML	A065098 001	Dec 16, 2002	Jul	CAHN
>A>	AB		400MG/5ML;EQ 57MG BASE/5ML	A065098 002	Dec 16, 2002	Jul	CAHN
>A>	AB		600MG/5ML;EQ 42.9MG BASE/5ML	A065358 001	Aug 13, 2007	Jul	CAHN

TABLET; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

AB	AUROBINDO PHARMA LTD	250MG;EQ 125MG BASE	A091569 001	Jan 20, 2012	Jan	NEWA	
AB		500MG;EQ 125MG BASE	A091569 002	Jan 20, 2012	Jan	NEWA	
AB		875MG;EQ 125MG BASE	A091568 001	Jan 20, 2012	Jan	NEWA	
>D>	AB	LEK PHARMS	500MG;EQ 125MG BASE	A065117 001	Nov 27, 2002	Jul	CAHN
>D>	AB		875MG;EQ 125MG BASE	A065093 001	Nov 21, 2002	Jul	CAHN
>A>	AB	SANDOZ INC	500MG;EQ 125MG BASE	A065117 001	Nov 27, 2002	Jul	CAHN
>A>	AB		875MG;EQ 125MG BASE	A065093 001	Nov 21, 2002	Jul	CAHN

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

ADDERALL XR 10

AB	SHIRE	2.5MG;2.5MG;2.5MG;2.5MG	N021303 001	Oct 11, 2001	Jun	CFTG
AB	ADDERALL XR 15					
AB	SHIRE	3.75MG;3.75MG;3.75MG;3.75MG	N021303 006	May 22, 2002	Jun	CFTG
AB	ADDERALL XR 20					
AB	SHIRE	5MG;5MG;5MG;5MG	N021303 002	Oct 11, 2001	Jun	CFTG
AB	ADDERALL XR 25					
AB	SHIRE	6.25MG;6.25MG;6.25MG;6.25MG	N021303 004	May 22, 2002	Jun	CFTG
AB	ADDERALL XR 30					
AB	+ SHIRE	7.5MG;7.5MG;7.5MG;7.5MG	N021303 003	Oct 11, 2001	Jun	CFTG
AB	ADDERALL XR 5					
AB	SHIRE	1.25MG;1.25MG;1.25MG;1.25MG	N021303 005	May 22, 2002	Jun	CFTG

CAPSULE, EXTENDED RELEASE; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB	ACTAVIS ELIZABETH	1.25MG;1.25MG;1.25MG;1.25MG	A077302 001	Jun 22, 2012	Jun	NEWA
AB		2.5MG;2.5MG;2.5MG;2.5MG	A077302 002	Jun 22, 2012	Jun	NEWA
AB		3.75MG;3.75MG;3.75MG;3.75MG	A077302 003	Jun 22, 2012	Jun	NEWA
AB		5MG;5MG;5MG;5MG	A077302 004	Jun 22, 2012	Jun	NEWA
AB		6.25MG;6.25MG;6.25MG;6.25MG	A077302 005	Jun 22, 2012	Jun	NEWA
AB		7.5MG;7.5MG;7.5MG;7.5MG	A077302 006	Jun 22, 2012	Jun	NEWA

AMPHETAMINE SULFATE

TABLET; ORAL

AMPHETAMINE SULFATE

>A>	INDEPENDENCE PHARMS	5MG	A200166 001	Aug 09, 2012	Jul	NEWA
>A>	+	10MG	A200166 002	Aug 09, 2012	Jul	NEWA

AMPHOTERICIN B

INJECTABLE, LIPID COMPLEX; INJECTION

AMPHOTEC

+	ALKOPHARMA USA	50MG/VIAL
+		100MG/VIAL

N050729 001	Nov 22, 1996	May	CAHN
N050729 002	Nov 22, 1996	May	CAHN

ANAGRELIDE HYDROCHLORIDE

CAPSULE; ORAL

AGRYLIN

AB	SHIRE LLC	EQ 0.5MG BASE	N020333 001	Mar 14, 1997	Jan	CAHN
	ANAGRELIDE HYDROCHLORIDE					
@	SANDOZ INC	EQ 0.5MG BASE	A076683 001	Apr 18, 2005	Mar	DISC

@ EQ 1MG BASE

A076683 002 Apr 18, 2005 Mar DISC

ANASTROZOLE

TABLET; ORAL

ANASTROZOLE

AB	APOTEX INC	1MG	A200654 001	May 11, 2012	Apr	NEWA
AB	IMPAX LABS INC	1MG	A091242 001	May 31, 2012	May	NEWA

APROTININ

INJECTABLE; INJECTION

TRASYLOL

@ BAYER HLTHCARE 10,000KIU/ML

N020304 001 Dec 29, 1993 May DISC

ARGATROBAN

INJECTABLE; INJECTION

ACOVA

AP	+ PFIZER	250MG/2.5ML (100MG/ML)	N020883 001	Jun 30, 2000	Jan	CTNA
AP	ARGATROBAN					

AP HIKMA PHARM CO LTD 250MG/2.5ML (100MG/ML)

N203049 001 Jan 05, 2012 Jan NEWA

ARMODAFINIL

TABLET; ORAL

ARMODAFINIL

AB	MYLAN PHARMS INC	50MG	A200043 001	Jun 01, 2012	May	NEWA
AB		150MG	A200043 002	Jun 01, 2012	May	NEWA
AB		250MG	A200043 003	Jun 01, 2012	May	NEWA
	NUVIGIL					
AB	CEPHALON	50MG	N021875 001	Jun 15, 2007	May	CFTG
AB		150MG	N021875 003	Jun 15, 2007	May	CFTG

TABLET; ORAL

NUVIGIL

AB + CEPHALON 250MG N021875 004 Jun 15, 2007 May CFTG

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

FOR SOLUTION; ORAL

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

AA NOVEL LABS INC 4.7GM;100GM;1.015GM;5.9MG;2.691GM A090145 001 Jan 25, 2012 Jan NEWA ;7.5GM

MOVIPREP

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

ASENAPINE MALEATE

TABLET; SUBLINGUAL

SAPHRIS

>A> ORGANON SUB MERCK EQ 5MG BASE N022117 001 Aug 13, 2009 Jul CAHN

>A> + EQ 10MG BASE N022117 002 Aug 13, 2009 Jul CAHN

>D> ORGANON USA INC EQ 5MG BASE N022117 001 Aug 13, 2009 Jul CAHN

>D> + EQ 10MG BASE N022117 002 Aug 13, 2009 Jul CAHN

ASPIRIN; CARISOPRODOL

TABLET; ORAL

CARISOPRODOL AND ASPIRIN

AB PROSAM LABS 325MG;200MG A040252 001 Dec 10, 1997 Feb CAHN

ASPIRIN; CARISOPRODOL; CODEINE PHOSPHATE

TABLET; ORAL

CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE

AB PROSAM LABS 325MG;200MG;16MG A040283 001 Dec 29, 1998 Feb CAHN

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

ATOMOXETINE HYDROCHLORIDE

@ ZYDUS PHARMS USA INC 18MG A079017 001 Sep 17, 2010 Jun DISC

@ 25MG A079017 002 Sep 17, 2010 Jun DISC

@ 40MG A079017 003 Sep 17, 2010 Jun DISC

@ 60MG A079017 004 Sep 17, 2010 Jun DISC

@ 80MG A079017 005 Sep 17, 2010 Jun DISC

@ 100MG A079017 006 Sep 17, 2010 Jun DISC

STRATTERA

LILLY 10MG N021411 002 Nov 26, 2002 Jun CTEC

18MG N021411 003 Nov 26, 2002 Jun CTEC

25MG N021411 004 Nov 26, 2002 Jun CTEC

40MG N021411 005 Nov 26, 2002 Jun CTEC

+ 60MG N021411 006 Nov 26, 2002 Jun CTEC

80MG N021411 007 Feb 14, 2005 Jun CTEC

100MG N021411 008 Feb 14, 2005 Jun CTEC

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

AB APOTEX INC EQ 10MG BASE A090548 001 May 29, 2012 May NEWA

AB EQ 20MG BASE A090548 002 May 29, 2012 May NEWA

AB EQ 40MG BASE A090548 003 May 29, 2012 May NEWA

TABLET; ORALATORVASTATIN CALCIUM

AB	APOTEX INC	EQ 80MG BASE	A090548 004	May 29, 2012	May	NEWA
>A>	AB DR REDDYS LABS LTD	EQ 10MG BASE	A091650 001	Jul 17, 2012	Jul	NEWA
>A>	AB	EQ 20MG BASE	A091650 002	Jul 17, 2012	Jul	NEWA
>A>	AB	EQ 40MG BASE	A091650 003	Jul 17, 2012	Jul	NEWA
>A>	AB	EQ 80MG BASE	A202357 001	Jul 17, 2012	Jul	NEWA
AB	MYLAN PHARMS INC	EQ 10MG BASE	A091226 001	May 29, 2012	May	NEWA
AB		EQ 20MG BASE	A091226 002	May 29, 2012	May	NEWA
AB		EQ 40MG BASE	A091226 003	May 29, 2012	May	NEWA
AB		EQ 80MG BASE	A091226 004	May 29, 2012	May	NEWA
AB	SANDOZ INC	EQ 10MG BASE	A077575 001	May 29, 2012	May	NEWA
AB		EQ 20MG BASE	A077575 002	May 29, 2012	May	NEWA
AB		EQ 40MG BASE	A077575 003	May 29, 2012	May	NEWA
AB		EQ 80MG BASE	A077575 004	May 29, 2012	May	NEWA
AB	TEVA PHARMS	EQ 10MG BASE	A078773 001	May 29, 2012	May	NEWA
AB		EQ 20MG BASE	A078773 002	May 29, 2012	May	NEWA
AB		EQ 40MG BASE	A078773 003	May 29, 2012	May	NEWA
AB		EQ 80MG BASE	A078773 004	May 29, 2012	May	NEWA

ATRACURIUM BESYLATEINJECTABLE; INJECTIONATRACURIUM BESYLATE

AP	+ BEDFORD	10MG/ML	A074901 001	Jul 18, 1997	Feb	CTEC
AP	SAGENT PHARMS	10MG/ML	A091489 001	Feb 17, 2012	Feb	NEWA
	ATRACURIUM BESYLATE PRESERVATIVE FREE					
AP	+ BEDFORD	10MG/ML	A074900 001	Jul 18, 1997	Feb	CTEC
AP	SAGENT PHARMS	10MG/ML	A091488 001	Feb 17, 2012	Feb	NEWA

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDESOLUTION; ORALDIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

AA	+ ROXANE	0.025MG/5ML;2.5MG/5ML	A087708 001	May 03, 1982	Apr	CRLD
	LOMOTIL @ GD SEARLE LLC	0.025MG/5ML;2.5MG/5ML	N012699 001		Apr	DISC

AVANAFILTABLET; ORALSTENDRAVIVUS

	50MG	N202276 001	Apr 27, 2012	Apr	NEWA
	100MG	N202276 002	Apr 27, 2012	Apr	NEWA
+	200MG	N202276 003	Apr 27, 2012	Apr	NEWA

AXITINIBTABLET; ORALINLYTAPFIZER

	1MG	N202324 001	Jan 27, 2012	Jan	NEWA
+	5MG	N202324 002	Jan 27, 2012	Jan	NEWA

AZELAIC ACIDGEL; TOPICALFINACEA

+	BAYER HLTHCARE	15%	N021470 001	Dec 24, 2002	Jun	CAHN
---	----------------	-----	-------------	--------------	-----	------

AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC AZELASTINE HYDROCHLORIDE AT ALCON PHARMA	0.05%	A202305 001 May 31, 2012 May NEWA
SPRAY, METERED; NASAL AZELASTINE HYDROCHLORIDE AB SUN PHARMA GLOBAL	EQ 0.125MG BASE/SPRAY	A090423 001 May 23, 2012 May NEWA

AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE

SPRAY, METERED; NASAL DYMISTA + MEDA PHARMS	EQ 0.125MG BASE/SPRAY; 0.05MG/SPRAY	N202236 001 May 01, 2012 May NEWA
---	--	-----------------------------------

AZILSARTAN KAMEDOXOMIL

TABLET; ORAL EDARBI TAKEDA PHARMS USA	EQ 40MG MEDOXOMIL	N200796 001 Feb 25, 2011 Apr CAHN
+ +	EQ 80MG MEDOXOMIL	N200796 002 Feb 25, 2011 Apr CAHN

AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE

TABLET; ORAL EDARBYCLOL TAKEDA PHARMS USA	EQ 40MG MEDOXOMIL; 12.5MG EQ 40MG MEDOXOMIL; 25MG	N202331 001 Dec 20, 2011 Apr CAHN
+ +		N202331 002 Dec 20, 2011 Apr CAHN

AZITHROMYCIN

INJECTABLE; INJECTION AZITHROMYCIN >D> AP PLIVA HRVATSKA DOO	EQ 500MG BASE/VIAL	A065265 001 Jan 18, 2007 Jul DISC
>A> @	EQ 500MG BASE/VIAL	A065265 001 Jan 18, 2007 Jul DISC
@ TEVA PARENTERAL	EQ 500MG BASE/VIAL	N050809 001 Dec 19, 2006 Jun DISC
@	EQ 2.5GM BASE/VIAL	N050809 002 Dec 19, 2006 Jun DISC

BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE >A> AT AKORN	400 UNITS/GM; 1%; EQ 3.5MG BASE/GM; 10,000 UNITS/GM	A065213 001 Jul 25, 2012 Jul NEWA
>D> + BAUSCH AND LOMB	400 UNITS/GM; 1%; EQ 3.5MG BASE/GM; 10,000 UNITS/GM	A064068 001 Oct 30, 1995 Jul CTEC
>A> AT +	400 UNITS/GM; 1%; EQ 3.5MG BASE/GM; 10,000 UNITS/GM	A064068 001 Oct 30, 1995 Jul CTEC

BACLOFEN

INJECTABLE; INTRATHECAL GABLOFEN CNS THERAPS INC	1MG/ML	N022462 004 Jun 22, 2012 Jun NEWA
TABLET; ORAL BACLOFEN AB PROSAM LABS	10MG	A077089 001 Oct 31, 2007 Feb CAHN
	20MG	A077088 001 Oct 31, 2007 Feb CAHN
TABLET, ORALLY DISINTEGRATING; ORAL KEMSTRO @ UCB INC	10MG	N021589 001 Oct 30, 2003 Jan CAHN
@	20MG	N021589 002 Oct 30, 2003 Jan CAHN

BALSALAZIDE DISODIUM

TABLET; ORAL
GIAZO
+ SALIX PHARMS 1.1GM N022205 001 Feb 03, 2012 Feb NEWA

BECLOMETHASONE DIPROPIONATE

AEROSOL, METERED; NASAL
QNASL
+ TEVA BRANDED PHARM 0.08MG/ACTIVATION N202813 001 Mar 23, 2012 Mar NEWA

BENZONATATE

CAPSULE; ORAL
BENZONATATE
@ NESHER PHARMS 100MG A040795 001 Oct 31, 2007 Jun DISC
@ 200MG A040795 002 Oct 31, 2007 Jun DISC

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL
BENZACLIN
BT + VALEANT INTL 5%;EQ 1% BASE N050756 002 Apr 20, 2007 Apr CAHN
AB 5%;EQ 1% BASE N050756 001 Dec 21, 2000 Apr CAHN
CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE
AB PERRIGO ISRAEL 5%;1.2% A090979 001 Jun 26, 2012 Jun NEWA
DUAC
AB + STIEFEL 5%;1.2% N050741 001 Aug 26, 2002 Jun CFTG

BENZOYL PEROXIDE; ERYTHROMYCIN

GEL; TOPICAL
BENZAMYCIN
AB + VALEANT INTL 5%;3% N050557 001 Oct 26, 1984 Jan CAHN
BENZAMYCIN PAK
+ VALEANT INTL 5%;3% N050769 001 Nov 27, 2000 Jun CAHN

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL
BENZPHETAMINE HYDROCHLORIDE
AA EMCURE PHARMS LTD 50MG A202061 001 Jan 27, 2012 Jan NEWA

BENZTROPINE MESYLATE

INJECTABLE; INJECTION
BENZTROPINE MESYLATE
AP APP PHARMS LLC 1MG/ML A090233 001 Jul 28, 2009 May CAHN
COGENTIN
AP + OAK PHARMS AKORN 1MG/ML N012015 001 Feb CAHN
TABLET; ORAL
BENZTROPINE MESYLATE
@ LANNETT HOLDINGS INC 0.5MG A088877 001 Apr 11, 1985 Jan DISC
@ 1MG A088894 001 Apr 11, 1985 Jan DISC
@ 2MG A088895 001 Apr 11, 1985 Jan DISC
AA PROSAM LABS 0.5MG A040699 001 Feb 14, 2008 Feb CMFD
AA 1MG A040705 001 Feb 14, 2008 Feb CMFD
AA 2MG A040706 001 Feb 14, 2008 Feb CMFD
AA + USL PHARMA 0.5MG A040103 001 Dec 12, 1996 Jan CRLD
AA + 1MG A040103 002 Dec 12, 1996 Jan CRLD
AA + 2MG A040103 003 Dec 12, 1996 Jan CRLD

BETAMETHASONE

SYRUP; ORAL
 CELESTONE
 @ SCHERING 0.6MG/5ML N014215 002 May DISC

BETAMETHASONE DIPROPIONATE

CREAM; TOPICAL
 BETAMETHASONE DIPROPIONATE
 AB + FOUGERA PHARMS EQ 0.05% BASE N019137 001 Jun 26, 1984 Jan CAHN
 CREAM, AUGMENTED; TOPICAL
 BETAMETHASONE DIPROPIONATE
 AB FOUGERA PHARMS EQ 0.05% BASE A076215 001 Dec 09, 2003 Jan CAHN
 GEL, AUGMENTED; TOPICAL
 BETAMETHASONE DIPROPIONATE
 AB + FOUGERA PHARMS EQ 0.05% BASE A075276 001 May 13, 2003 Jan CAHN
 LOTION, AUGMENTED; TOPICAL
 BETAMETHASONE DIPROPIONATE
 AB FOUGERA PHARMS EQ 0.05% BASE A077111 001 May 21, 2007 Jan CAHN
 OINTMENT, AUGMENTED; TOPICAL
 BETAMETHASONE DIPROPIONATE
 AB FOUGERA PHARMS EQ 0.05% BASE A075373 001 Jun 22, 1999 Jan CAHN

BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

CREAM; TOPICAL
 CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE
 AB FOUGERA PHARMS EQ 0.05% BASE;1% A075502 001 Jun 05, 2001 Jan CAHN
 LOTION; TOPICAL
 CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE
 AB FOUGERA PHARMS EQ 0.05% BASE;1% A076516 001 Jun 16, 2005 Jan CAHN
 LOTRISONE
 AB + SCHERING CORP EQ 0.05% BASE;1% N020010 001 Dec 08, 2000 Feb CAHN

BETAMETHASONE VALERATE

AEROSOL, FOAM; TOPICAL
 LUXIQ
 + STIEFEL EQ 0.12% BASE N020934 001 Feb 28, 1999 Jan CAHN

BISOPROLOL FUMARATE

TABLET; ORAL
 ZEBETA
 AB TEVA WOMENS 5MG N019982 002 Jul 31, 1992 Jan CAHN
 AB + 10MG N019982 001 Jul 31, 1992 Jan CAHN

BOCEPREVIR

CAPSULE; ORAL
 VICTRELIS
 + MERCK SHARP DOHME 200MG N202258 001 May 13, 2011 Mar CAHN

BORTEZOMIB

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS
 VELCADE
 + MILLENNIUM PHARMS 3.5MG/VIAL N021602 001 May 13, 2003 Jan CDFR

BOSENTAN

TABLET; ORAL
 TRACLEER
 ACTELION PHARMS LTD 62.5MG N021290 001 Nov 20, 2001 May CAHN
 + 125MG N021290 002 Nov 20, 2001 May CAHN

BUDESONIDE

POWDER, METERED; INHALATION
 PULMICORT
 @ ASTRAZENECA 0.16MG/INH N020441 002 Jun 24, 1997 Jun DISC
 SUSPENSION; INHALATION
 BUDESONIDE
 >A> AN WATSON LABS INC 0.25MG/2ML A078404 001 Jul 31, 2012 Jul NEWA
 >A> AN 0.5MG/2ML A078404 002 Jul 31, 2012 Jul NEWA

BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE

AEROSOL, METERED; INHALATION
 SYMBICORT
 + ASTRAZENECA 0.08MG/INH;0.0045MG/INH N021929 001 Jul 21, 2006 Jan CDFR
 + 0.16MG/INH;0.0045MG/INH N021929 002 Jul 21, 2006 Jan CDFR

BUPIVACAINE

INJECTABLE, LIPOSOMAL; INJECTION
 EXPAREL
 >D> + PACIRA PHARMS INC 133MG/10ML (13.3MG/ML) N022496 001 Oct 28, 2011 Jul DISC
 >A> @ 133MG/10ML (13.3MG/ML) N022496 001 Oct 28, 2011 Jul DISC

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

FILM; SUBLINGUAL
 SUBOXONE
 RECKITT BENCKISER EQ 2MG BASE;EQ 0.5MG BASE N022410 001 Aug 30, 2010 Apr CAIN
 + EQ 8MG BASE;EQ 2MG BASE N022410 002 Aug 30, 2010 Apr CAIN
 TABLET; SUBLINGUAL
 SUBOXONE
 RECKITT BENCKISER EQ 2MG BASE;EQ 0.5MG BASE N020733 001 Oct 08, 2002 Apr CPOT
 + EQ 8MG BASE;EQ 2MG BASE N020733 002 Oct 08, 2002 Apr CPOT

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL
 FORFIVO XL
 + EDGEMONT PHARMS LLC 450MG N022497 001 Nov 10, 2011 Mar CAHN

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL
 BUSPIRONE HYDROCHLORIDE
 @ APOTEX 5MG A075521 001 Apr 05, 2002 Mar DISC
 @ 10MG A075521 002 Apr 05, 2002 Mar DISC
 @ 15MG A075521 003 Apr 05, 2002 Mar DISC
 @ EGIS 5MG A075119 001 Mar 14, 2002 Mar DISC
 @ 10MG A075119 002 Mar 14, 2002 Mar DISC
 @ 15MG A075119 003 Jan 23, 2003 Mar DISC
 @ NESHER PHARMS 5MG A075572 001 Feb 27, 2002 Jun DISC
 @ 10MG A075572 002 Feb 27, 2002 Jun DISC
 @ 15MG A075572 003 Feb 27, 2002 Jun DISC
 @ PROSAM LABS 5MG A075388 001 May 09, 2002 Mar DISC

TABLET; ORAL

BUSPIRONE HYDROCHLORIDE

AB	PROSAM LABS	5MG	A075388 001	May 09, 2002	Feb	CMFD
	@	10MG	A075388 002	May 09, 2002	Mar	DISC
AB		10MG	A075388 002	May 09, 2002	Feb	CMFD
	@	15MG	A075388 003	May 09, 2002	Mar	DISC
AB		15MG	A075388 003	May 09, 2002	Feb	CMFD
AB		30MG	A078302 001	Dec 17, 2007	Feb	CMFD

BUTOCONAZOLE NITRATE

CREAM; VAGINAL

BUTOCONAZOLE NITRATE

@ PERRIGO ISRAEL 2%

A200923 001 May 18, 2012 Apr NEWA

CAFFEINE CITRATE

SOLUTION; INTRAVENOUS

CAF'CIT

AP	+	BEDFORD LABS	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N020793 001	Sep 21, 1999	Jan	CAHN
----	---	--------------	---------------------------------------	-------------	--------------	-----	------

SOLUTION; ORAL

CAF'CIT

AA	+	BEDFORD LABS	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N020793 002	Apr 12, 2000	Jan	CAHN
----	---	--------------	---------------------------------------	-------------	--------------	-----	------

CALCIPOURIENE

CREAM; TOPICAL

CALCIPOURIENE

AB		TOLMAR	0.005%	A200935 001	May 30, 2012	May	NEWA
----	--	--------	--------	-------------	--------------	-----	------

DOVONEX

AB	+	LEO PHARMA AS	0.005%	N020554 001	Jul 22, 1996	May	CFTG
----	---	---------------	--------	-------------	--------------	-----	------

SOLUTION; TOPICAL

CALCIPOURIENE

AT		FOUGERA PHARMS	0.005%	A078305 001	May 06, 2008	Jan	CAHN
----	--	----------------	--------	-------------	--------------	-----	------

CALCIUM ACETATE

CAPSULE; ORAL

CALCIUM ACETATE

AB		PADDOCK LLC	EQ 169MG CALCIUM	A091312 001	Jun 01, 2012	May	NEWA
----	--	-------------	------------------	-------------	--------------	-----	------

CARBIDOPA; ENTACAPONE; LEVODOPA

TABLET; ORAL

CARBIDOPA, LEVODOPA AND ENTACAPONE

AB		SUN PHARMA GLOBAL	25MG;200MG;100MG	A079085 001	May 10, 2012	Apr	NEWA
----	--	-------------------	------------------	-------------	--------------	-----	------

AB			37.5MG;200MG;150MG	A079085 002	May 10, 2012	Apr	NEWA
----	--	--	--------------------	-------------	--------------	-----	------

STALEVO 100

AB		ORION PHARMA	25MG;200MG;100MG	N021485 002	Jun 11, 2003	Apr	CFTG
----	--	--------------	------------------	-------------	--------------	-----	------

STALEVO 150

AB		ORION PHARMA	37.5MG;200MG;150MG	N021485 003	Jun 11, 2003	Apr	CFTG
----	--	--------------	--------------------	-------------	--------------	-----	------

CARBIDOPA; LEVODOPA

TABLET, ORALLY DISINTEGRATING; ORAL

PARCOPA

AB		UCB INC	10MG;100MG	A076699 001	Aug 27, 2004	Jan	CAHN
----	--	---------	------------	-------------	--------------	-----	------

AB			25MG;100MG	A076699 002	Aug 27, 2004	Jan	CAHN
----	--	--	------------	-------------	--------------	-----	------

AB	+		25MG;250MG	A076699 003	Aug 27, 2004	Jan	CAHN
----	---	--	------------	-------------	--------------	-----	------

CARBOPLATIN

INJECTABLE; INJECTION

CARBOPLATIN

AP	CIPLA LTD	50MG/VIAL	A077383 001	Jan 27, 2006	Mar	CAHN
AP		150MG/VIAL	A077383 002	Jan 27, 2006	Mar	CAHN
AP		450MG/VIAL	A077383 003	Jan 27, 2006	Mar	CAHN
AP	ONCO THERAPIES LTD	50MG/VIAL	A091510 001	May 29, 2012	May	NEWA
AP		150MG/VIAL	A091510 002	May 29, 2012	May	NEWA
AP		450MG/VIAL	A091510 003	May 29, 2012	May	NEWA

PARAPLATIN

@ CORDEN PHARMA	50MG/VIAL	N019880 001	Mar 03, 1989	Feb	CAHN
@	150MG/VIAL	N019880 002	Mar 03, 1989	Feb	CAHN
@	450MG/VIAL	N019880 003	Mar 03, 1989	Feb	CAHN

INJECTABLE; IV (INFUSION)

CARBOPLATIN

AP	ACTAVIS TOTOWA	50MG/5ML (10MG/ML)	A078732 001	Feb 06, 2012	Jan	NEWA
AP		150MG/15ML (10MG/ML)	A078732 002	Feb 06, 2012	Jan	NEWA
AP		450MG/45ML (10MG/ML)	A078732 003	Feb 06, 2012	Jan	NEWA
AP		600MG/60ML (10MG/ML)	A078732 004	Feb 06, 2012	Jan	NEWA
AP	CIPLA LTD	50MG/5ML (10MG/ML)	A077861 001	Jan 18, 2007	Mar	CAHN
AP		150MG/15ML (10MG/ML)	A077861 002	Jan 18, 2007	Mar	CAHN
AP		450MG/45ML (10MG/ML)	A077861 003	Jan 18, 2007	Mar	CAHN
AP		600MG/60ML (10MG/ML)	A077861 004	Jan 18, 2007	Mar	CAHN

>A>

CARFILZOMIB

>A>

POWDER; INTRAVENOUS

>A>

KYPROLIS

>A>

+ ONYX PHARMS

60MG/VIAL

N202714 001 Jul 20, 2012 Jul NEWA

CARISOPRODOL

TABLET; ORAL

CARISOPRODOL

AA	PROSAM LABS	350MG	A040188 001	Mar 07, 1997	Feb	CAHN
----	-------------	-------	-------------	--------------	-----	------

CEFACLOR

CAPSULE; ORAL

CEFACLOR

@ RANBAXY

EQ 250MG BASE

A064156 001 Aug 28, 1997 Apr DISC

@

EQ 500MG BASE

A064156 002 Aug 28, 1997 Apr DISC

FOR SUSPENSION; ORAL

CEFACLOR

@ RANBAXY

EQ 125MG BASE/5ML

A064166 001 Oct 02, 1997 Apr DISC

AB

+

EQ 125MG BASE/5ML

A064166 001 Oct 02, 1997 Feb CTEC

@

EQ 187MG BASE/5ML

A064165 001 Oct 02, 1997 Apr DISC

AB

+

EQ 187MG BASE/5ML

A064165 001 Oct 02, 1997 Feb CTEC

@

EQ 250MG BASE/5ML

A064164 001 Oct 02, 1997 Apr DISC

AB

+

EQ 250MG BASE/5ML

A064164 001 Oct 02, 1997 Feb CTEC

@

EQ 375MG BASE/5ML

A064155 001 Oct 02, 1997 Apr DISC

AB

+

EQ 375MG BASE/5ML

A064155 001 Oct 02, 1997 Feb CTEC

AB

YUNG SHIN PHARM

EQ 125MG BASE/5ML

A065412 001 Feb 17, 2012 Feb NEWA

AB

EQ 187MG BASE/5ML

A065412 002 Feb 17, 2012 Feb NEWA

AB

EQ 250MG BASE/5ML

A065412 003 Feb 17, 2012 Feb NEWA

AB

EQ 375MG BASE/5ML

A065412 004 Feb 17, 2012 Feb NEWA

TABLET, CHEWABLE; ORAL

RANICLOR

@ RANBAXY LABS LTD	EQ 125MG BASE	A065092 001	Dec 22, 2003	Apr	DISC
@	EQ 187MG BASE	A065092 002	Dec 22, 2003	Apr	DISC
@	EQ 250MG BASE	A065092 003	Dec 22, 2003	Apr	DISC
@	EQ 375MG BASE	A065092 004	Dec 22, 2003	Apr	DISC

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE; ORAL

CEFADROXIL

@ RANBAXY LABS LTD	EQ 500MG BASE	A065015 001	Jun 22, 1999	Apr	DISC
--------------------	---------------	-------------	--------------	-----	------

TABLET; ORAL

CEFADROXIL

@ RANBAXY	EQ 1GM BASE	A065018 001	Apr 23, 1999	Apr	DISC
-----------	-------------	-------------	--------------	-----	------

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

CEFAZOLIN AND DEXTROSE

B BRAUN	EQ 2GM BASE/VIAL	N050779 003	Jan 13, 2012	May	NEWA
---------	------------------	-------------	--------------	-----	------

CEFDITOREN PIVOXIL

TABLET; ORAL

SPECTRACEF

VANSEN PHARMA	200MG	N021222 001	Aug 29, 2001	Mar	CAHN
+	400MG	N021222 002	Jul 21, 2008	Mar	CAHN

CEFEPIMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

CEFEPIMIDE HYDROCHLORIDE

>A> AP	HOSPIRA INC	EQ 1GM BASE/VIAL	A202268 001	Jul 30, 2012	Jul	NEWA
>A> AP		EQ 2GM BASE/VIAL	A202268 002	Jul 30, 2012	Jul	NEWA

CEFIXIME

CAPSULE; ORAL

SUPRAX

+ LUPIN LTD	400MG	N203195 001	Jun 01, 2012	Jun	NEWA
-------------	-------	-------------	--------------	-----	------

CEFOTAXIME SODIUM

INJECTABLE; INJECTION

CLAFORAN

AP + SANOFI AVENTIS US	EQ 1GM BASE/VIAL	A062659 001	Jan 13, 1987	Mar	CRLD
AP +	EQ 2GM BASE/VIAL	A062659 002	Jan 13, 1987	Mar	CRLD

>D> CEFOXITIN

>D> INJECTABLE; INJECTION

>D> CEFOXITIN

>D> AP	ANTIBIOTICOS BRASIL	EQ 1GM BASE/VIAL	A065467 001	Aug 31, 2011	Jul	CAIN
>D> AP		EQ 2GM BASE/VIAL	A065467 002	Aug 31, 2011	Jul	CAIN
>D> AP		EQ 10GM BASE/VIAL	A065464 001	Aug 31, 2011	Jul	CAIN

>A> CEFOXITIN SODIUM

>A> INJECTABLE; INJECTION

>A> CEFOXITIN

>A> AP	ANTIBIOTICOS BRASIL	EQ 1GM BASE/VIAL	A065467 001	Aug 31, 2011	Jul	CAIN
>A> AP		EQ 2GM BASE/VIAL	A065467 002	Aug 31, 2011	Jul	CAIN
>A> AP		EQ 10GM BASE/VIAL	A065464 001	Aug 31, 2011	Jul	CAIN

CEPHALEXIN

FOR SUSPENSION; ORAL

KEFLEX

>D>	@ LEX PHARMS	EQ 100MG BASE/ML	N050406 003	Jul	CAHN
>D>	@	EQ 125MG BASE/5ML	N050406 001	Jul	CAHN
>D>	@	EQ 250MG BASE/5ML	N050406 002	Jul	CAHN
>A>	@ SHIONOGI INC	EQ 100MG BASE/ML	N050406 003	Jul	CAHN
>A>	@	EQ 125MG BASE/5ML	N050406 001	Jul	CAHN
>A>	@	EQ 250MG BASE/5ML	N050406 002	Jul	CAHN

CETIRIZINE HYDROCHLORIDE

SYRUP; ORAL

CETIRIZINE HYDROCHLORIDE

>D> AA	RANBAXY	5MG/5ML	A077472 001	Jun 18, 2008	Jul	DISC
>A>	@ RANBAXY LABS LTD	5MG/5ML	A077472 001	Jun 18, 2008	Jul	DISC
AA	SILARX	5MG/5ML	A078876 001	May 11, 2012	Apr	NEWA

CHLORAMBUCIL

TABLET; ORAL

LEUKERAN

+ ASPEN GLOBAL INC 2MG

N010669 002 May CAHN

CHLORPHENIRAMINE POLISTIREX; HYDROCODONE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

HYDROCODONE POLISTIREX AND CHLORPHENIRAMNE POLISTIREX

AB	CORNERSTONE THERAP	EQ 8MG MALEATE/5ML;EQ 10MG BITARTRATE/5ML	A091671 001	Jun 29, 2012	Jun	NEWA
----	--------------------	---	-------------	--------------	-----	------

CICLESONIDE

AEROSOL, METERED; INHALATION

ALVESCO

NYCOMED GMBH 0.08MG/INH

N021658 002 Jan 10, 2008 May CAHN

+ 0.16MG/INH

N021658 003 Jan 10, 2008 May CAHN

AEROSOL, METERED; NASAL

ZETONNA

+ NYCOMED GMBH 0.037MG/INH

N202129 001 Jan 20, 2012 Jan NEWA

SPRAY, METERED; NASAL

OMNARIS

+ NYCOMED GMBH 0.05MG/INH

N022004 001 Oct 20, 2006 May CAHN

CICLOPIROX

CREAM; TOPICAL

CICLOPIROX

AB	FOUGERA PHARMS	0.77%	A076435 001	Dec 29, 2004	Jan	CAHN
----	----------------	-------	-------------	--------------	-----	------

GEL; TOPICAL

CICLOPIROX

AB	FOUGERA PHARMS	0.77%	A077896 001	Jun 10, 2008	Jan	CAHN
----	----------------	-------	-------------	--------------	-----	------

AB	GLENMARK GENERICS	0.77%	A091595 001	Feb 29, 2012	Feb	NEWA
----	-------------------	-------	-------------	--------------	-----	------

SHAMPOO; TOPICAL

CICLOPIROX

AT	FOUGERA PHARMS	1%	A090146 001	May 25, 2010	Jan	CAHN
----	----------------	----	-------------	--------------	-----	------

SOLUTION; TOPICAL

CICLOPIROX

AT	MYLAN PHARMS INC	8%	A078567 001	Sep 18, 2007	Feb	CAHN
----	------------------	----	-------------	--------------	-----	------

SUSPENSION; TOPICAL

CICLOPIROX

AB FOUGERA PHARMS 0.77% A076422 001 Aug 06, 2004 Jan CAHN

CIDOFOVIR

INJECTABLE; INJECTION

CIDOFOVIR

>A>	AP EMCURE PHARMS LTD	EQ 75MG BASE/ML	A202501 001 Jul 26, 2012 Jul NEWA
	AP MYLAN LLC	EQ 75MG BASE/ML	A201276 001 Jun 27, 2012 Jun NEWA
	VISTIDE		
	AP + GILEAD SCIENCES INC	EQ 75MG BASE/ML	N020638 001 Jun 26, 1996 Jun CFTG

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

@ HOSPIRA	EQ 300MG BASE/2ML	A074344 001 Jan 31, 1995 Jun DISC
@	EQ 300MG BASE/2ML	A074345 001 Jan 31, 1995 Jun DISC

CIPROFLOXACIN

INJECTABLE; INJECTION

CIPROFLOXACIN

@ TEVA PARENTERAL	200MG/20ML (10MG/ML)	A077782 001 Aug 28, 2006 Mar DISC
@	400MG/40ML (10MG/ML)	A077782 002 Aug 28, 2006 Jan DISC
CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER		
@ BAXTER HLTHCARE	200MG/100ML	A077888 001 Mar 18, 2008 Jun DISC
@	400MG/200ML	A077888 002 Mar 18, 2008 Jun DISC

CIPROFLOXACIN HYDROCHLORIDE

TABLET; ORAL

CIPROFLOXACIN HYDROCHLORIDE

@ PLIVA	EQ 100MG BASE	A076426 001 Jun 15, 2005 Jan DISC
@	EQ 250MG BASE	A076426 002 Jun 15, 2005 Jan DISC
@	EQ 500MG BASE	A076426 003 Jun 15, 2005 Jan DISC
@	EQ 750MG BASE	A076426 004 Jun 15, 2005 Jan DISC

CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CIPROFLOXACIN EXTENDED RELEASE

AB DR REDDYS LABS LTD 212.6MG;EQ 287.5MG BASE A077701 002 Oct 31, 2007 Apr NEWA

CISATRACURIUM BESYLATE

INJECTABLE; INJECTION

CISATRACURIUM BESYLATE

AP SANDOZ INC	EQ 2MG BASE/ML	A200159 001 Feb 03, 2012 Jan NEWA
CISATRACURIUM BESYLATE PRESERVATIVE FREE		
AP SANDOZ INC	EQ 2MG BASE/ML	A200154 001 Feb 03, 2012 Jan NEWA
AP	EQ 10MG BASE/ML	A200154 002 Feb 03, 2012 Jan NEWA
NIMBEX		
AP + ABBOTT	EQ 2MG BASE/ML	N020551 001 Dec 15, 1995 Jan CFTG
NIMBEX PRESERVATIVE FREE		
AP + ABBOTT	EQ 2MG BASE/ML	N020551 003 Dec 15, 1995 Jan CFTG
AP +	EQ 10MG BASE/ML	N020551 002 Dec 15, 1995 Jan CFTG

CISPLATIN

INJECTABLE; INJECTION

CISPLATIN

AP	ONCO THERAPIES LTD @ TEVA PARENTERAL	1MG/ML 1MG/ML	A091062 001 Apr 18, 2012 Apr NEWA
			A074814 001 May 16, 2000 Mar DISC

>A> CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE

>A> FOR SOLUTION; ORAL

>A> PREPOPIK

>A>	+ FERRING PHARMS AS	12GM/PACKET;3.5GM/PACKET;10MG/PAC KET	N202535 001 Jul 16, 2012 Jul NEWA
-----	---------------------	---------------------------------------	-----------------------------------

CLARITHROMYCIN

TABLET; ORAL

CLARITHROMYCIN

>A> AB	AUROBINDO	250MG	A065489 001 Jul 25, 2012 Jul NEWA
>A> AB		500MG	A065489 002 Jul 25, 2012 Jul NEWA

CLINDAMYCIN PHOSPHATE

CREAM; VAGINAL

CLINDAMYCIN PHOSPHATE

AB	FOUGERA PHARMS	EQ 2% BASE	A065139 001 Dec 27, 2004 Jan CAHN
----	----------------	------------	-----------------------------------

GEL; TOPICAL

CLINDAMYCIN PHOSPHATE

AB	FOUGERA PHARMS	EQ 1% BASE	A064160 001 Jan 28, 2000 Jan CAHN
----	----------------	------------	-----------------------------------

INJECTABLE; INJECTION

CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER

AP	+ PHARMACIA AND UPJOHN	EQ 6MG BASE/ML	N050639 001 Aug 30, 1989 May CFTG
----	------------------------	----------------	-----------------------------------

AP	+	EQ 12MG BASE/ML	N050639 002 Aug 30, 1989 May CFTG
----	---	-----------------	-----------------------------------

AP	+	EQ 18MG BASE/ML	N050639 003 Apr 10, 1991 May CFTG
----	---	-----------------	-----------------------------------

CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER

AP	SANDOZ INC	EQ 6MG BASE/ML	A201692 001 May 31, 2012 May NEWA
----	------------	----------------	-----------------------------------

AP		EQ 12MG BASE/ML	A201692 002 May 31, 2012 May NEWA
----	--	-----------------	-----------------------------------

AP		EQ 18MG BASE/ML	A201692 003 May 31, 2012 May NEWA
----	--	-----------------	-----------------------------------

LOTION; TOPICAL

CLINDAMYCIN PHOSPHATE

AB	FOUGERA PHARMS	EQ 1% BASE	A065067 001 Jan 31, 2002 Jan CAHN
----	----------------	------------	-----------------------------------

SOLUTION; TOPICAL

CLINDAMYCIN PHOSPHATE

AT	FOUGERA PHARMS	EQ 1% BASE	A065254 001 Feb 14, 2006 Jan CAHN
----	----------------	------------	-----------------------------------

CLINDAMYCIN PHOSPHATE; TRETINOIN

GEL; TOPICAL

VELTIN

BX	+ STIEFEL GSK	1.2%;0.025%	N050803 001 Jul 16, 2010 May CRLD
----	---------------	-------------	-----------------------------------

CLOBAZAM

TABLET; ORAL

ONFI

LUNDBECK LLC

		5MG	N202067 001 Oct 21, 2011 Mar CAHN
--	--	-----	-----------------------------------

		10MG	N202067 002 Oct 21, 2011 Mar CAHN
--	--	------	-----------------------------------

	+	20MG	N202067 003 Oct 21, 2011 Mar CAHN
--	---	------	-----------------------------------

CLOBETASOL PROPIONATE

AEROSOL, FOAM; TOPICAL
CLOBETASOL PROPIONATE

>D>	AB	PERRIGO ISRAEL	0.05%	A077763	001	Mar 10, 2008	Jul	CTEC
>A>	AB1		0.05%	A077763	001	Mar 10, 2008	Jul	CTEC
>A>	AB2		0.05%	A201402	001	Aug 14, 2012	Jul	NEWA

OLUX

>D>	AB	STIEFEL LABS INC	0.05%	N021142	001	May 26, 2000	Jul	CTEC
>A>	AB1	+	0.05%	N021142	001	May 26, 2000	Jul	CTEC

OLUX E

>D>	AB	STIEFEL LABS INC	0.05%	N022013	001	Jan 12, 2007	Jul	CFTG
>A>	AB2	+	0.05%	N022013	001	Jan 12, 2007	Jul	CFTG

CREAM; TOPICAL
CLOBETASOL PROPIONATE

@ G AND W LABS INC	0.05%	A074139	001	Aug 03, 1994	Jun	CAHN
--------------------	-------	---------	-----	--------------	-----	------

CLOBETASOL PROPIONATE (EMOLLIENT)

AB2	FOUGERA PHARMS	0.05%	A075430	001	May 26, 1999	Jan	CAHN
-----	----------------	-------	---------	-----	--------------	-----	------

TEMOVATE

AB1	+ FOUGERA PHARMS	0.05%	N019322	001	Dec 27, 1985	Jan	CAHN
-----	------------------	-------	---------	-----	--------------	-----	------

GEL; TOPICAL
CLOBETASOL PROPIONATE

AB	FOUGERA PHARMS	0.05%	A075368	001	Feb 15, 2000	Jan	CAHN
----	----------------	-------	---------	-----	--------------	-----	------

TEMOVATE

AB	+ FOUGERA PHARMS	0.05%	N020337	001	Apr 29, 1994	Jan	CAHN
----	------------------	-------	---------	-----	--------------	-----	------

LOTION; TOPICAL
CLOBETASOL PROPIONATE

AB	TARO	0.05%	A200302	001	Jul 02, 2012	Jun	NEWA
----	------	-------	---------	-----	--------------	-----	------

OINTMENT; TOPICAL
CLOBETASOL PROPIONATE

AB	FOUGERA PHARMS	0.05%	A074407	001	Feb 23, 1996	Jan	CAHN
----	----------------	-------	---------	-----	--------------	-----	------

TEMOVATE

AB	+ FOUGERA PHARMS	0.05%	N019323	001	Dec 27, 1985	Jan	CAHN
----	------------------	-------	---------	-----	--------------	-----	------

SHAMPOO; TOPICAL
CLOBETASOL PROPIONATE

>A>	AB	PERRIGO ISRAEL	0.05%	A090974	001	Aug 09, 2012	Jul	NEWA
-----	----	----------------	-------	---------	-----	--------------	-----	------

SOLUTION; TOPICAL
CLOBETASOL PROPIONATE

AT	FOUGERA PHARMS	0.05%	A075391	001	Feb 08, 1999	Jan	CAHN
@ G AND W LABS INC	0.05%	A074331	001	Dec 15, 1995	Jun	CAHN	

TEMOVATE

AT	+ FOUGERA PHARMS	0.05%	N019966	001	Feb 22, 1990	Jan	CAHN
----	------------------	-------	---------	-----	--------------	-----	------

CLONIDINE HYDROCHLORIDE

INJECTABLE; INJECTION
DURACLON

AP	MYLAN INSTITUTIONAL	1 MG/10 ML (0.1 MG/ML)	N020615	001	Oct 02, 1996	Feb	CAHN
AP	+	5 MG/10 ML (0.5 MG/ML)	N020615	002	Apr 27, 1999	Feb	CAHN

TABLET, EXTENDED RELEASE; ORAL
KAPVAY

SHIONOGI INC	0.2MG	N022331	004	Sep 28, 2010	May	CMFD
--------------	-------	---------	-----	--------------	-----	------

CLOPIDOGREL BISULFATE

TABLET; ORAL
CLOPIDOGREL BISULFATE

AB	APOTEX INC	EQ 75MG BASE	A076274	001	May 17, 2012	Apr	NEWA
----	------------	--------------	---------	-----	--------------	-----	------

TABLET; ORALCLOPIDOGREL BISULFATE

AB	AUROBINDO PHARMA LTD	EQ 75MG BASE	A090540 001	May 17, 2012	Apr	NEWA	
AB	DR REDDYS LABS INC	EQ 75MG BASE	A076273 001	Jan 14, 2008	Mar	CMFD	
AB	DR REDDYS LABS LTD	EQ 300MG BASE	A091023 001	May 17, 2012	Apr	NEWA	
AB	GATE PHARMS	EQ 300MG BASE	A091216 001	May 17, 2012	Apr	NEWA	
AB	MYLAN PHARMS INC	EQ 75MG BASE	A077665 001	May 17, 2012	Apr	NEWA	
AB		EQ 300MG BASE	A077665 002	May 17, 2012	Apr	NEWA	
AB	ROXANE	EQ 75MG BASE	A078004 001	May 17, 2012	Apr	NEWA	
AB	SUN PHARMA GLOBAL	EQ 75MG BASE	A090494 001	May 17, 2012	Apr	NEWA	
AB	TEVA	EQ 75MG BASE	A076999 001	May 17, 2012	Apr	NEWA	
AB	TEVA PHARMS	EQ 300MG BASE	A090625 001	May 17, 2012	Apr	NEWA	
AB	TORRENT PHARMS LTD	EQ 75MG BASE	A090844 001	May 17, 2012	Apr	NEWA	
>A>	AB	WOCKHARDT LTD	EQ 75MG BASE	A202266 001	Aug 14, 2012	Jul	NEWA
		PLAVIX					
AB	SANOFI AVENTIS US	EQ 75MG BASE	N020839 001	Nov 17, 1997	Mar	CTEC	
AB	+	EQ 300MG BASE	N020839 002	Sep 20, 2007	Apr	CTEC	

CLORAZEPATE DIPOTASSIUMCAPSULE; ORALTRANXENE

@	LUNDBECK LLC	3.75MG	N017105 001	Mar	CAHN
@		7.5MG	N017105 002	Mar	CAHN
@		15MG	N017105 003	Mar	CAHN

TABLET; ORALTRANXENE

AB	LUNDBECK LLC	3.75MG	N017105 006	Mar	CAHN
AB		7.5MG	N017105 007	Mar	CAHN
AB	+	15MG	N017105 008	Mar	CAHN
	TRANXENE SD				
@	LUNDBECK LLC	11.25MG	N017105 005	Mar	CAHN
@		22.5MG	N017105 004	Mar	CAHN

CLOTRIMAZOLECREAM; TOPICALCLOTRIMAZOLE

AB	FOUGERA PHARMS	1%	A078338 001	Sep 02, 2008	Jan	CAHN
----	----------------	----	-------------	--------------	-----	------

CLOZAPINETABLET, ORALLY DISINTEGRATING; ORALFAZACLO ODT

JAZZ	12.5MG	N021590 004	May 30, 2007	Apr	CAHN
	25MG	N021590 001	Feb 10, 2004	Apr	CAHN
@	50MG	N021590 003	Jun 03, 2005	Apr	CAHN
+	100MG	N021590 002	Feb 10, 2004	Apr	CAHN
	150MG	N021590 005	Jul 09, 2010	Apr	CAHN
	200MG	N021590 006	Jul 09, 2010	Apr	CAHN
JAZZ PHARMS III	12.5MG	N021590 004	May 30, 2007	Jun	CAHN
	25MG	N021590 001	Feb 10, 2004	Jun	CAHN
@	50MG	N021590 003	Jun 03, 2005	Jun	CAHN
+	100MG	N021590 002	Feb 10, 2004	Jun	CAHN
	150MG	N021590 005	Jul 09, 2010	Jun	CAHN
	200MG	N021590 006	Jul 09, 2010	Jun	CAHN

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE

AA TRIS PHARMA INC 10MG/5ML; 6.25MG/5ML A200386 001 Jun 29, 2012 Jun NEWA

COPPER

INTRAUTERINE DEVICE; INTRAUTERINE

PARAGARD T 380A

+ TEVA WOMENS 309MG/COPPER N018680 001 Nov 15, 1984 May CAHN

COSYNTROPIN

INJECTABLE; INJECTION

COSYNTROPIN

AP INNOPHARMA INC 0.25MG/VIAL A202147 001 Jun 29, 2012 Jun NEWA

CROMOLYN SODIUM

AEROSOL, METERED; INHALATION

INTAL

@ KING PFIZER 0.8MG/INH N018887 001 Dec 05, 1985 Jun DISC

CONCENTRATE; ORAL

GASTROCROM

AA + JAZZ PHARMS COMMERCIAL 100MG/5ML N020479 001 Feb 29, 1996 Mar CAHN

AA + JAZZ PHARMS INTL 100MG/5ML N020479 001 Feb 29, 1996 Jun CAHN

CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

>D> AB MYLAN 10MG A073144 001 May 30, 1991 Jul CRLD

>A> AB + MYLAN PHARMS INC 10MG A073144 001 May 30, 1991 Jul CRLD

AB PROSAM LABS 5MG A077291 001 Feb 03, 2006 Feb CAHN

AB 10MG A077209 001 Oct 04, 2005 Feb CAHN

FLEXERIL

>D> AB + JANSSEN R AND D 10MG N017821 002 Jul DISC

>D> AB JANSSEN RES AND DEV 5MG N017821 001 Jul DISC

>A> @ 5MG N017821 001 Jul DISC

>A> @ 10MG N017821 002 Jul DISC

CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION

CYTOXAN

AP + BAXTER HLTHCARE 500MG/VIAL N012142 003 Apr CRLD

AP + 1GM/VIAL N012142 004 Aug 30, 1982 Apr CRLD

AP + 2GM/VIAL N012142 005 Aug 30, 1982 Apr CRLD

CYCLOSPORINE

CAPSULE; ORAL

CYCLOSPORINE

AB1 WATSON LABS INC 25MG A065044 002 Dec 20, 2000 May CAHN

AB1 100MG A065044 001 Dec 20, 2000 May CAHN

CYTARABINE

INJECTABLE; INJECTION

CYTARABINE

AP ONCO THERAPIES LTD 100MG/ML A201784 001 Jan 30, 2012 Jan NEWA

DACTINOMYCIN

INJECTABLE; INJECTION

COSMEGEN

AP + LUNDBECK LLC 0.5MG/VIAL N050682 001 Mar CAHN

DANTROLENE SODIUM

CAPSULE; ORAL

DANTROLENE SODIUM

>D>	@ ACTAVIS TOTOWA	25MG	A076686 001 Oct 24, 2005 Jul CAHN
>D>	@	50MG	A076686 002 Oct 24, 2005 Jul CAHN
>D>	@	100MG	A076686 003 Oct 24, 2005 Jul CAHN
>A>	@ MIKAH PHARMA	25MG	A076686 001 Oct 24, 2005 Jul CAHN
>A>	@	50MG	A076686 002 Oct 24, 2005 Jul CAHN
>A>	@	100MG	A076686 003 Oct 24, 2005 Jul CAHN

DAPIPRAZOLE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

>A>	DAPIPRAZOLE HYDROCHLORIDE		
>A>	@ FERA PHARMS	0.5%	N019849 001 Dec 31, 1990 Jul CTNA
>D>	REV-EYES		
>D>	@ FERA PHARMS	0.5%	N019849 001 Dec 31, 1990 Jul CTNA

DARUNAVIR ETHANOLATE

SUSPENSION; ORAL

PREZISTA

+ JANSSEN PRODS EQ 100MG BASE/ML N202895 001 Dec 16, 2011 Feb CAHN

TABLET; ORAL

PREZISTA

JANSSEN PRODS	EQ 75MG BASE	N021976 004 Dec 18, 2008 Feb CAHN
	EQ 150MG BASE	N021976 005 Dec 18, 2008 Feb CAHN
@	EQ 300MG BASE	N021976 001 Jun 23, 2006 Feb CAHN
	EQ 400MG BASE	N021976 003 Oct 21, 2008 Feb CAHN
+	EQ 600MG BASE	N021976 002 Feb 25, 2008 Feb CAHN

DEMECLOCYCLINE HYDROCHLORIDE

TABLET; ORAL

DECLOMYCIN

>D>	COREPHARMA	150MG	N050261 002 Jul DISC
>A>	@	150MG	N050261 002 Jul DISC
>D>	AB +	300MG	N050261 003 Jul DISC
>A>	@	300MG	N050261 003 Jul DISC
	DEMECLOCYCLINE HYDROCHLORIDE		
>D>	AB AMNEAL PHARM	300MG	A065425 002 Feb 27, 2008 Jul CRLD
>A>	AB +	300MG	A065425 002 Feb 27, 2008 Jul CRLD

DESFLURANE

LIQUID; INHALATION

SUPRANE

>A>	+ BAXTER HLTHCARE	99.9%	N020118 001 Sep 18, 1992 Jul CAHN
>D>	+ BAXTER HLTHCARE CORP	99.9%	N020118 001 Sep 18, 1992 Jul CAHN

DESLORATADINE

TABLET; ORAL

DESLORATADINE

AB BELCHER PHARMS 5MG A078355 001 Apr 19, 2012 Apr NEWA

TABLET; ORAL

DESLORATADINE

AB MYLAN PHARMS INC 5MG A078351 001 Feb 10, 2012 Jan NEWA

DESMOPRESSIN ACETATE

SOLUTION; NASAL

DDAVP

AB + SANOFI AVENTIS US 0.01% N017922 001 Mar CFTG

DESMOPRESSIN ACETATE

AB SUN PHARM INDS 0.01% A077212 001 Apr 12, 2012 Mar NEWA

DESOGESTREL; ETHINYLL ESTRADIOL

TABLET; ORAL-28

MIRCETTE

AB + TEVA BRANDED PHARM 0.15MG,N/A;0.02MG,0.01MG N020713 001 Apr 22, 1998 Jun CAHN

VIORELE

AB GLENMARK GENERICS .15MG,N/A; 0.02MG,0.01MG A091346 001 Apr 02, 2012 Mar NEWA

DESONIDE

LOTION; TOPICAL

DESONIDE

AB FOUGERA PHARMS 0.05% A075860 001 Mar 19, 2002 Jan CAHN

OINTMENT; TOPICAL

DESONIDE

AB FOUGERA PHARMS 0.05% A075751 001 Mar 12, 2001 Jan CAHN

DESOXIMETASONE

CREAM; TOPICAL

DESOXIMETASONE

AB FOUGERA PHARMS 0.25% A078369 001 Jun 29, 2010 Jan CAHN

TOPICORT

+ TARO 0.05%

A073210 001 Nov 30, 1990 Apr CTNA

+ 0.25%

A073193 001 Nov 30, 1990 Apr CTNA

@ TARO PHARMS NORTH 0.25%

N017856 001 Apr DISC

TOPICORT LP

@ TARO PHARMS NORTH 0.05%

N018309 001 Apr DISC

GEL; TOPICAL

TOPICORT

AB + TARO 0.05% A074904 001 Jul 14, 1998 Apr CTNA

@ TARO PHARMS NORTH 0.05%

N018586 001 Mar 29, 1982 Apr DISC

OINTMENT; TOPICAL

DESOXIMETASONE

AB + TARO 0.25% A074286 001 Jun 07, 1996 Mar CRLD

TOPICORT

+ TARO 0.25%

A074286 001 Jun 07, 1996 Apr CTNA

@ TARO PHARMS NORTH 0.25%

N018763 001 Sep 30, 1983 Mar DISC

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

DEXAMETHASONE SODIUM PHOSPHATE

AP + APP PHARMS LLC EQ 10MG PHOSPHATE/ML A040572 001 Apr 22, 2005 Feb CRLD

DEXRAZOXANE HYDROCHLORIDE

INJECTABLE; INJECTION

TOTECT

+ APRICUS PHARMS EQ 500MG BASE/VIAL

N022025 001 Sep 06, 2007 Jun CAHN

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

AA TRIS PHARMA INC 15MG/5ML; 6.25MG/5ML A091687 001 Jun 28, 2012 Jun NEWA

DIAZEPAM

INJECTABLE; INJECTION

DIAZEPAM

>D>	@ BAXTER HLTHCARE	5MG/ML	A071309 001 Jul 17, 1987 Jul CAHN
>D>	@	5MG/ML	A070313 001 Dec 16, 1985 Jul CAHN
>D>	@	5MG/ML	A070311 001 Dec 16, 1985 Jul CAHN
>D>	@	5MG/ML	A071308 001 Jul 17, 1987 Jul CAHN
>D>	@	5MG/ML	A070312 001 Dec 16, 1985 Jul CAHN
>D>	@	5MG/ML	A071310 001 Jul 17, 1987 Jul CAHN
>A>	@ HIKMA MAPLE	5MG/ML	A070312 001 Dec 16, 1985 Jul CAHN
>A>	@	5MG/ML	A070311 001 Dec 16, 1985 Jul CAHN
>A>	@	5MG/ML	A070313 001 Dec 16, 1985 Jul CAHN
>A>	@	5MG/ML	A071308 001 Jul 17, 1987 Jul CAHN
>A>	@	5MG/ML	A071309 001 Jul 17, 1987 Jul CAHN
>A>	@	5MG/ML	A071310 001 Jul 17, 1987 Jul CAHN
	@ HOSPIRA	5MG/ML	A071584 001 Oct 13, 1987 Apr DISC

DICLOFENAC POTASSIUM

CAPSULE; ORAL

ZIPSOR

+ DEPOMED INC 25MG N022202 001 Jun 16, 2009 Jun CAHN

DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE; ORAL

ARTHROTEC

AB	GD SEARLE LLC	50MG; 0.2MG	N020607 001 Dec 24, 1997 Jun CFTG
AB	+	75MG; 0.2MG	N020607 002 Dec 24, 1997 Jun CFTG
DICLOFENAC SODIUM AND MISOPROSTOL			
AB	WATSON LABS INC	50MG; 0.2MG	A201089 001 Jul 09, 2012 Jun NEWA
AB		75MG; 0.2MG	A201089 002 Jul 09, 2012 Jun NEWA

DIDANOSINE

TABLET, CHEWABLE; ORAL

DIDANOSINE

>A>	AUROBINDO	100MG	A077275 001 Aug 14, 2012 Jul NEWA
>A>		150MG	A077275 002 Aug 14, 2012 Jul NEWA
>A>		200MG	A077275 003 Aug 14, 2012 Jul NEWA

DIFLORASONE DIACETATE

CREAM; TOPICAL

DIFLORASONE DIACETATE

BX	+ FOUGERA PHARMS	0.05%	A076263 001 Dec 20, 2002 Jan CAHN
AB1	+	0.05%	A075187 001 Mar 30, 1998 Jan CAHN
OINTMENT; TOPICAL			
DIFLORASONE DIACETATE			
AB	FOUGERA PHARMS	0.05%	A075374 001 Apr 27, 1999 Jan CAHN

DIFLUNISAL

TABLET; ORAL

DIFLUNISAL

AB	EMCURE PHARMS USA	500MG	A202845 001 Mar 08, 2012 Feb NEWA
AB	+ TEVA	500MG	A073673 001 Jul 31, 1992 Feb CTEC

DIGOXIN

INJECTABLE; INJECTION

DIGOXIN

>D>	AP	BAXTER HLTHCARE	0.25MG/ML	A083391 001 Jul CAHN
>A>	AP	HIKMA MAPLE	0.25MG/ML	A083391 001 Jul CAHN
	@ HOSPIRA		0.25MG/ML	A040093 001 May 16, 1996 Jun DISC
		TABLET; ORAL		
		DIGOXIN		
>D>		@ ACTAVIS TOTOWA	0.125MG	A040282 001 Dec 23, 1999 Jul CAHN
>D>		@	0.25MG	A040282 002 Dec 23, 1999 Jul CAHN
>A>		@ MYLAN PHARMS INC	0.125MG	A040282 001 Dec 23, 1999 Jul CAHN
>A>		@	0.25MG	A040282 002 Dec 23, 1999 Jul CAHN

DILTIAZEM HYDROCHLORIDE

INJECTABLE; INJECTION

DILTIAZEM HYDROCHLORIDE

>D>	AP	BAXTER HLTHCARE	5MG/ML	A078538 001 Dec 17, 2008 Jul CAHN
>A>	AP	HIKMA FARMACEUTICA	5MG/ML	A202651 001 Aug 09, 2012 Jul NEWA
>A>	AP	HIKMA MAPLE	5MG/ML	A078538 001 Dec 17, 2008 Jul CAHN
	@ TEVA PARENTERAL		5MG/ML	A074894 001 Aug 26, 1997 Jan DISC

DINOPROSTONEINSERT, EXTENDED RELEASE; VAGINAL
CERVIDIL

>D>	+	CONTROLLED THERAP	10MG	N020411 001 Mar 30, 1995 Jul CAHN
>A>	+	FERRING CONTROLLED	10MG	N020411 001 Mar 30, 1995 Jul CAHN

DIPHENHYDRAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

BENADRYL

@ MCNEIL CONS 50MG/ML

N006146 002 Mar DISC

BENADRYL PRESERVATIVE FREE

@ MCNEIL CONS 50MG/ML

N009486 001 Mar DISC

DIPHENHYDRAMINE HYDROCHLORIDE

>D>	AP	+	BAXTER HLTHCARE	50MG/ML	A080817 002 Jul CAHN
>D>		@		50MG/ML	A083183 001 Jul CAHN
	AP	+		50MG/ML	A080817 002 Apr CRLD
>A>	AP	+	HIKMA MAPLE	50MG/ML	A080817 002 Jul CAHN
>A>		@		50MG/ML	A083183 001 Jul CAHN

DIPYRIDAMOLE

INJECTABLE; INJECTION

DIPYRIDAMOLE

>D>	AP	BAXTER HLTHCARE	5MG/ML	A074521 001 Oct 18, 1996 Jul CAHN
>A>	AP	HIKMA MAPLE	5MG/ML	A074521 001 Oct 18, 1996 Jul CAHN
	@ TEVA PARENTERAL		5MG/ML	A074952 001 Nov 26, 1997 Jan DISC

TABLET; ORAL

DIPYRIDAMOLE

AB	PROSAM LABS	25MG	A040542 001 Apr 21, 2006 Feb CAHN
----	-------------	------	-----------------------------------

TABLET; ORAL

DIPYRIDAMOLE

AB	PROSAM LABS	50MG	A040542 002	Apr 21, 2006	Feb	CAHN
AB		75MG	A040542 003	Apr 21, 2006	Feb	CAHN

DISOPYRAMIDE PHOSPHATE

CAPSULE, EXTENDED RELEASE; ORAL

DISOPYRAMIDE PHOSPHATE

@ NESHER PHARMS	EQ 150MG BASE	A071200 001	Dec 15, 1987	Jun	DISC
-----------------	---------------	-------------	--------------	-----	------

DIVALPROEX SODIUM

TABLET, EXTENDED RELEASE; ORAL

DIVALPROEX SODIUM

AB	DR REDDYS LABS LTD	EQ 250MG VALPROIC ACID	A090161 001	Mar 15, 2012	Feb	NEWA
AB	REDDYS	EQ 500MG VALPROIC ACID	A090070 001	Mar 12, 2012	Feb	NEWA

DOCETAXEL

INJECTABLE; INJECTION

DOCETAXEL

AP	ACCORD HLTHCARE	20MG/ML (20MG/ML)	N201195 003	Apr 20, 2012	Apr	NEWA
AP		20MG/0.5ML (40MG/ML)	N201195 001	Jun 08, 2011	Jan	CTEC
AP		80MG/4ML (20MG/ML)	N201195 004	Apr 20, 2012	Apr	NEWA
AP		80MG/2ML (40MG/ML)	N201195 002	Jun 08, 2011	Jan	CTEC
AP		160MG/8ML (20MG/ML)	N201195 005	Apr 20, 2012	Apr	NEWA
AP	APOTEX INC	20MG/0.5ML (40MG/ML)	N022312 001	Jan 11, 2012	Jan	NEWA
AP		80MG/2ML (40MG/ML)	N022312 002	Jan 11, 2012	Jan	NEWA
TAXOTERE						
AP	+ SANOFI AVENTIS US	20MG/ML (20MG/ML)	N020449 003	Aug 03, 2010	Apr	CTEC
AP	+	80MG/4ML (20MG/ML)	N020449 004	Aug 02, 2010	Apr	CTEC

DONEPEZIL HYDROCHLORIDE

TABLET; ORAL

DONEPEZIL HYDROCHLORIDE

>A>	AB	INDICUS PHARMA	5MG	A201634 001	Jun 13, 2012	Jul	NEWA
>A>	AB		10MG	A201634 002	Jun 13, 2012	Jul	NEWA
AB		PRINSTON INC	5MG	A200292 001	May 31, 2011	Jun	CAHN
AB			10MG	A200292 002	May 31, 2011	Jun	CAHN

DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

COSOPT PF

+ MERCK SHARP DOHME	EQ 2% BASE;EQ 0.5% BASE	N202667 001	Feb 01, 2012	Feb	NEWA
---------------------	-------------------------	-------------	--------------	-----	------

DOXAZOSIN MESYLATE

TABLET; ORAL

DOXAZOSIN MESYLATE

@ NESHER PHARMS	EQ 1MG BASE	A075609 001	Oct 18, 2000	Feb	DISC
@	EQ 2MG BASE	A075609 002	Oct 18, 2000	Feb	DISC
@	EQ 4MG BASE	A075609 003	Oct 18, 2000	Feb	DISC
@	EQ 8MG BASE	A075609 004	Oct 18, 2000	Feb	DISC

DOXEPIN HYDROCHLORIDE

CREAM; TOPICAL

ZONALON

+ FOUGERA PHARMS	5%	N020126 001	Apr 01, 1994	Jan	CAHN
------------------	----	-------------	--------------	-----	------

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

DOXORUBICIN HYDROCHLORIDE

AP	ONCO THERAPIES LTD	2MG/ML	A200901 001	Feb 14, 2012	Jan	NEWA
>A>	AP	SANDOZ INC	A200146 001	Jul 18, 2012	Jul	NEWA
AP	SUN PHARM IND	2MG/ML	A091418 001	Feb 15, 2012	Jan	NEWA
INJECTABLE, LIPOSOMAL; INJECTION						
DOXIL						
+ JANSSEN R AND D		20MG/10ML (2MG/ML)	N050718 001	Nov 17, 1995	Jan	CAHN
+ JANSSEN R AND D		50MG/25ML (2MG/ML)	N050718 002	Jun 13, 2000	Jan	CAHN

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

AB	MYLAN PHARMS INC	EQ 150MG BASE	A202778 001	Jun 08, 2012	May	NEWA
	@ SANDOZ INC	EQ 50MG BASE	A065032 001	Jun 30, 2000	Mar	DISC
	@	EQ 100MG BASE	A065032 002	Jun 30, 2000	Mar	DISC
MONODOX						
>A> AB	AQUA PHARMS	EQ 50MG BASE	N050641 002	Feb 10, 1992	Jul	CAHN
>A> AB		EQ 75MG BASE	N050641 003	Oct 18, 2006	Jul	CAHN
>A> AB +		EQ 100MG BASE	N050641 001	Dec 29, 1989	Jul	CAHN
>D> AB	WATSON PHARMS	EQ 50MG BASE	N050641 002	Feb 10, 1992	Jul	CAHN
>D> AB		EQ 75MG BASE	N050641 003	Oct 18, 2006	Jul	CAHN
>D> AB +		EQ 100MG BASE	N050641 001	Dec 29, 1989	Jul	CAHN
TABLET; ORAL						
DOXYCYCLINE						
	@ SANDOZ INC	EQ 50MG BASE	A065353 001	Nov 27, 2006	Mar	DISC
	@	EQ 75MG BASE	A065353 002	Nov 27, 2006	Mar	DISC
	@	EQ 100MG BASE	A065353 003	Nov 27, 2006	Mar	DISC

DOXYCYCLINE HYCLATE

INJECTABLE; INJECTION

DOXYCYCLINE

>D>	@ BAXTER HLTHCARE	EQ 100MG BASE/VIAL	A062450 001	Oct 27, 1983	Jul	CAHN
>D>	@	EQ 200MG BASE/VIAL	A062450 002	Oct 27, 1983	Jul	CAHN
>A>	@ HIKMA MAPLE	EQ 100MG BASE/VIAL	A062450 001	Oct 27, 1983	Jul	CAHN
>A>	@	EQ 200MG BASE/VIAL	A062450 002	Oct 27, 1983	Jul	CAHN
DOXYCYCLINE HYCLATE						
>D>	@ BAXTER HLTHCARE	EQ 100MG BASE/VIAL	A062992 001	Feb 16, 1989	Jul	CAHN
>D>	@	EQ 200MG BASE/VIAL	A062992 002	Feb 16, 1989	Jul	CAHN
>A>	@ HIKMA MAPLE	EQ 100MG BASE/VIAL	A062992 001	Feb 16, 1989	Jul	CAHN
>A>	@	EQ 200MG BASE/VIAL	A062992 002	Feb 16, 1989	Jul	CAHN

TABLET; ORAL

DOXYCYCLINE HYCLATE

AB	LARKEN LABS	EQ 20MG BASE	A065287 001	Feb 28, 2006	Apr	CMFD
TABLET, DELAYED RELEASE; ORAL						
DORYX						
AB +	MAYNE PHARMA	EQ 150MG BASE	N050795 003	Jun 20, 2008	Jan	CTEC
DOXYCYCLINE HYCLATE						
AB	MYLAN PHARMS INC	EQ 150MG BASE	A091052 001	Feb 08, 2012	Jan	NEWA

DROSPIRENONE; ESTRADIOL

TABLET; ORAL

ANGELIQ

BAYER HLTHCARE

0.25MG;0.5MG

N021355 001 Feb 29, 2012 Feb NEWA

ECONAZOLE NITRATE

CREAM; TOPICAL
ECONAZOLE NITRATE
AB + FOUGERA PHARMS 1% A076075 001 Nov 26, 2002 Jan CAHN

EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL
COMPLERA
+ GILEAD SCIENCES INC 200MG;EQ 25MG BASE;300MG N202123 001 Aug 10, 2011 Apr CAIN

ENALAPRIL MALEATE

TABLET; ORAL
ENALAPRIL MALEATE
@ SANDOZ INC 2.5MG A075621 001 Aug 22, 2000 Feb DISC
@ 2.5MG A075496 001 Aug 22, 2000 Jan DISC
@ 5MG A075621 002 Aug 22, 2000 Feb DISC
@ 5MG A075496 002 Aug 22, 2000 Jan DISC
@ 10MG A075621 003 Aug 22, 2000 Feb DISC
@ 10MG A075459 001 Aug 22, 2000 Jan DISC
@ 20MG A075621 004 Aug 22, 2000 Feb DISC
@ 20MG A075459 002 Aug 22, 2000 Jan DISC

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL
ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE
@ SANDOZ INC 5MG;12.5MG A076116 001 Sep 19, 2001 Mar DISC
@ 10MG;25MG A076116 002 Sep 19, 2001 Mar DISC

ENFLURANE

LIQUID; INHALATION
ETHRANE
>A> AN + BAXTER HLTHCARE 99.9% N017087 001 Jul CAHN
>D> AN + BAXTER HLTHCARE CORP 99.9% N017087 001 Jul CAHN

ENTACAPONE

TABLET; ORAL
COMTAN
>D> + ORION PHARMA 200MG N020796 001 Oct 19, 1999 Jul CFTG
>A> AB + 200MG N020796 001 Oct 19, 1999 Jul CFTG
+ 200MG N020796 001 Oct 19, 1999 May CAHN
>A> ENTACAPONE
>A> AB SUN PHARMA GLOBAL 200MG A090690 001 Jul 16, 2012 Jul NEWA
>A> AB WOCKHARDT LTD 200MG A078941 001 Aug 16, 2012 Jul NEWA

EPINASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC
EPINASTINE HYDROCHLORIDE
AT LUITPOLD 0.05% A090951 001 Oct 31, 2011 Mar CAHN

EPINEPHRINE

INJECTABLE; IM-SC
ADRENAClick
BX + COREPHARMA EQ 0.15MG /DELIVERY N020800 003 Nov 25, 2009 Mar CAHN
BX + EQ 0.3MG /DELIVERY N020800 004 Nov 25, 2009 Mar CAHN

INJECTABLE; IM-SC

TWINJECT 0.15

BX	+ COREPHARMA	EQ 0.15MG /DELIVERY	N020800 002 May 28, 2004 Mar CAHN
	TWINJECT 0.3		
BX	+ COREPHARMA	EQ 0.3MG /DELIVERY	N020800 001 May 30, 2003 Mar CAHN

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

OCTOCOCAINE

AP	+ SEPTODONT	0.01MG/ML; 2%	A084048 001	May CRLD
	XYLOCAINE DENTAL WITH EPINEPHRINE			
	@ DENTSPLY PHARM	0.01MG/ML; 2%	N021381 001	May DISC
	@	0.02MG/ML; 2%	N021381 002	May DISC

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

EPIRUBICIN HYDROCHLORIDE

AP	CIPLA LTD	50MG/25ML (2MG/ML)	A065361 001 Oct 22, 2007 Mar CAHN
AP		200MG/100ML (2MG/ML)	A065361 002 Oct 22, 2007 Mar CAHN
AP	HISUN PHARM HANGZHOU	50MG/25ML (2MG/ML)	A090075 001 Mar 25, 2010 Mar CAHN
AP		200MG/100ML (2MG/ML)	A090075 002 Mar 25, 2010 Mar CAHN
AP	ONCO THERAPIES LTD	50MG/25ML (2MG/ML)	A091599 001 Mar 12, 2012 Mar NEWA
AP		200MG/100ML (2MG/ML)	A091599 002 Mar 12, 2012 Mar NEWA

EPOPROSTENOL SODIUM

INJECTABLE; INJECTION

VELETRI

	ACTELION PHARMS LTD	EQ 0.5MG BASE/VIAL	N022260 002 Jun 28, 2012 Jun NEWA
+		EQ 1.5MG BASE/VIAL	N022260 001 Jun 27, 2008 May CAHN

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS; ORAL

ERYC

AB	WARNER CHILCOTT LLC	250MG	A062338 001	Jun CAHN
	GEL; TOPICAL			
	E-GLADES			
>D>	AT COREPHARMA	2%	A065009 001 Mar 18, 2002 Jul DISC	
>A>	@	2%	A065009 001 Mar 18, 2002 Jul DISC	
	ERYTHROMYCIN			
AT	FOUGERA PHARMS	2%	A064184 001 Sep 30, 1997 Jan CAHN	
	SOLUTION; TOPICAL			
	C-SOLVE-2			
AT	FOUGERA PHARMS	2%	A062468 001 Jul 03, 1985 Jan CMFD	
	SWAB; TOPICAL			
	ERYCETTE			
	@ JOHNSON AND JOHNSON	2%	N050594 001 Feb 15, 1985 Apr CAHN	
	ERYTHROMYCIN			
AT	+ FOUGERA PHARMS	2%	A065320 001 Jul 25, 2006 Jan CAHN	

ESCITALOPRAM OXALATE

SOLUTION; ORAL

ESCITALOPRAM OXALATE

AA	AMNEAL PHARMS	EQ 5MG BASE/5ML	A202227 001 Mar 14, 2012 Feb NEWA
AA	AUROBINDO PHARMA LTD	EQ 5MG BASE/5ML	A079062 001 Apr 02, 2012 Mar NEWA
AA	HETERO LABS LTD III	EQ 5MG BASE/5ML	A202221 001 Jun 12, 2012 May NEWA
AA	TARO	EQ 5MG BASE/5ML	A079121 001 May 03, 2012 Apr NEWA

ESTROGENS, CONJUGATED SYNTHETIC A

TABLET; ORAL

CENESTIN

TEVA BRANDED PHARM	0.3MG 0.45MG 0.625MG 0.9MG +	N020992 001 Jun 21, 2002 Jun CAHN N020992 005 Feb 05, 2004 Jun CAHN N020992 002 Mar 24, 1999 Jun CAHN N020992 003 Mar 24, 1999 Jun CAHN N020992 004 Mar 13, 2000 Jun CAHN
--------------------	--	---

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL

LEVONORGESTREL AND ETHINYL ESTRADIOL

+ WATSON LABS	0.02MG;0.09MG	A079218 001 Jun 06, 2011 May CRLD
---------------	---------------	-----------------------------------

LOSEASONIQUE

AB TEVA BRANDED PHARM	0.02MG,0.01MG;0.1MG,N/A	N022262 001 Oct 24, 2008 Jun CAHN
LYBREL @ WYETH PHARMS INC	0.02MG;0.09MG	N021864 001 May 22, 2007 May DISC
PREVEN EMERGENCY CONTRACEPTIVE KIT @ TEVA BRANDED PHARM	0.05MG;0.25MG	N020946 001 Sep 01, 1998 Jun CAHN
SEASONALE		

>D> AB + DURAMED RES	0.03MG;0.15MG	N021544 001 Sep 05, 2003 Jul CAHN
----------------------	---------------	-----------------------------------

>A> AB + TEVA BRANDED PHARM	0.03MG;0.15MG	N021544 001 Sep 05, 2003 Jul CAHN
-----------------------------	---------------	-----------------------------------

SEASONIQUE

AB + TEVA BRANDED PHARM	0.03MG,0.01MG;0.15MG,N/A	N021840 001 May 25, 2006 Jun CAHN
-------------------------	--------------------------	-----------------------------------

TABLET; ORAL-21

NORDETTE-21

@ TEVA BRANDED PHARM 0.03MG;0.15MG

N018668 001 May 10, 1982 Jun CAHN

TABLET; ORAL-28

FALMINA

AB1 NOVAST LABS LTD	0.02MG;0.1MG	A090721 001 Mar 28, 2012 Mar NEWA
---------------------	--------------	-----------------------------------

MARLISSA

AB GLENMARK GENERICS	0.03MG;0.15MG	A091452 001 Feb 29, 2012 Feb NEWA
----------------------	---------------	-----------------------------------

MYZILRA

AB VINTAGE PHARMS	0.03MG,0.04MG,0.03MG;0.05MG,0.075 MG,0.125MG	A077502 001 Nov 23, 2011 May CDFR
-------------------	--	-----------------------------------

NORDETTE-28

AB + TEVA BRANDED PHARM	0.03MG;0.15MG	N018782 001 Jul 21, 1982 Jun CAHN
-------------------------	---------------	-----------------------------------

ETHINYL ESTRADIOL; NORELGESTROMIN

FILM, EXTENDED RELEASE; TRANSDERMAL

ORTHO EVRA

+ JANSSEN PHARMS	0.75MG;6MG	N021180 001 Nov 20, 2001 Jun CPOT
------------------	------------	-----------------------------------

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-28

ALYACEN 1/35

AB GLENMARK GENERICS	0.035MG;1MG	A091634 001 Jan 19, 2012 Jan NEWA
----------------------	-------------	-----------------------------------

ALYACEN 7/7/7

AB GLENMARK GENERICS	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A091636 001 Jan 19, 2012 Jan NEWA
----------------------	--	-----------------------------------

OVCON-35

AB + WARNER CHILCOTT LLC	0.035MG;0.4MG	N017716 001 Jun CAHN
--------------------------	---------------	----------------------

OVCON-50

+ WARNER CHILCOTT LLC	0.05MG;1MG	N017576 001 Jun CAHN
-----------------------	------------	----------------------

WERA

AB NOVAST LABS LTD	0.035MG;0.5MG	A091204 001 Mar 27, 2012 Mar NEWA
--------------------	---------------	-----------------------------------

TABLET, CHEWABLE; ORAL

FEMCON FE

AB + WARNER CHILCOTT LLC 0.035MG;0.4MG N021490 001 Nov 14, 2003 Jun CAHN

ETHINYLY ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL

FEMHRT

AB + WARNER CHILCOTT LLC 0.0025MG;0.5MG N021065 001 Jan 14, 2005 Jun CAHN
N021065 002 Oct 15, 1999 Jun CAHN

LO LOESTRIN FE

AB + WARNER CHILCOTT LLC 0.01MG,0.01MG;1MG,N/A N022501 001 Oct 21, 2010 Jun CAHN

TABLET; ORAL-21

ESTROSTEP 21

AB @ WARNER CHILCOTT LLC 0.02MG,0.03MG,0.035MG;1MG,1MG,1MG N020130 001 Oct 09, 1996 Jun CAHN
LOESTRIN 21 1.5/30

AB WARNER CHILCOTT LLC 0.03MG;1.5MG N017875 001 Jun CAHN

LOESTRIN 21 1/20

AB WARNER CHILCOTT LLC 0.02MG;1MG N017876 001 Jun CAHN

TABLET; ORAL-28

ESTROSTEP FE

AB + WARNER CHILCOTT LLC 0.02MG,0.03MG,0.035MG;1MG,1MG,1MG N020130 002 Oct 09, 1996 Jun CAHN
LOESTRIN FE 1.5/30

AB + WARNER CHILCOTT LLC 0.03MG;1.5MG N017355 001 Jun CAHN

ETHINYLY ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

MONO-LINYAH

AB NOVAST LABS LTD 0.035MG;0.25MG A090523 001 May 23, 2012 May NEWA

NORGESTIMATE AND ETHINYLY ESTRADIOL

AB GLENMARK GENERICS 0.035MG;0.25MG A200538 001 Apr 05, 2012 Mar NEWA

AB LUPIN PHARMS 0.025MG,0.025MG,0.025MG;0.18MG,0. A200541 001 Jun 25, 2012 Jun NEWA
215MG,0.25MG

TRI-LINYAH

AB NOVAST LABS LTD 0.035MG,0.035MG,0.035MG;0.18MG,0. A090524 001 May 30, 2012 May NEWA
215MG,0.25MGETHINYLY ESTRADIOL; NORGESTREL

TABLET; ORAL-28

ELINEST

AB NOVAST LABS LTD 0.03MG;0.3MG A091105 001 Mar 28, 2012 Mar NEWA

ETODOLAC

TABLET; ORAL

ETODOLAC

AB PROSAM LABS 400MG A074819 001 Feb 28, 1997 Feb CMFD

AB 500MG A074819 002 Apr 28, 1998 Feb CMFD

ETOMIDATE

INJECTABLE; INJECTION

ETOMIDATE

AP MYLAN INSTITUTIONAL 2MG/ML A091297 001 Jun 20, 2012 Jun NEWA

ETOPOSIDE

CAPSULE; ORAL

VEPESID

AB @ CORDEN PHARMA 50MG N019557 001 Dec 30, 1986 Feb CAHN

CAPSULE; ORAL

VEPESID

@ CORDEN PHARMA

100MG

N019557 002 Dec 30, 1986 Feb CAHN

INJECTABLE; INJECTION

ETOPOSIDE

AP + APP PHARMS LLC

20MG/ML

A074983 001 Sep 30, 1998 Mar CRLD

AP BEDFORD

20MG/ML

A074290 001 Jul 17, 1995 Mar CRLD

VEPESID

@ CORDEN PHARMA

20MG/ML

N018768 001 Nov 10, 1983 Feb CAHN

ETRAVIRINETABLET; ORAL

INTELENCE

JANSSEN R AND D

25MG

N022187 003 Mar 26, 2012 Mar NEWA

100MG

N022187 001 Jan 18, 2008 Feb CAHN

+

200MG

N022187 002 Dec 22, 2010 Feb CAHN

EVEROLIMUSTABLET; ORAL

AFINITOR

NOVARTIS

7.5MG

N022334 004 Mar 30, 2012 Apr NEWA

EXENATIDE SYNTHETICFOR SUSPENSION, EXTENDED RELEASE; SUBCUTANEOUS

BYDUREON

+ AMYLIN

2MG/VIAL

N022200 001 Jan 27, 2012 Jan NEWA

EZETIMIBETABLET; ORAL

ZETIA

+ MSD INTL GMBH

10MG

N021445 001 Oct 25, 2002 Feb CAHN

+ MSP SINGAPORE

10MG

N021445 001 Oct 25, 2002 Jan CAHN

EZETIMIBE; SIMVASTATINTABLET; ORAL

VYTORIN

MSD INTL

10MG;10MG

N021687 001 Jul 23, 2004 Jan CAHN

10MG;20MG

N021687 002 Jul 23, 2004 Jan CAHN

10MG;40MG

N021687 003 Jul 23, 2004 Jan CAHN

+

10MG;80MG

N021687 004 Jul 23, 2004 Jan CAHN

EZOGABINETABLET; ORAL

POTIGA

GLAXOSMITHKLINE

50MG

N022345 001 Jun 10, 2011 May CAHN

200MG

N022345 002 Jun 10, 2011 May CAHN

300MG

N022345 003 Jun 10, 2011 May CAHN

+

400MG

N022345 004 Jun 10, 2011 May CAHN

FAMCICLOVIRTABLET; ORAL

FAMCICLOVIR

AB MACLEODS PHARMS LTD

125MG

A201022 001 Jan 12, 2012 Jan NEWA

AB

250MG

A201022 002 Jan 12, 2012 Jan NEWA

AB

500MG

A201022 003 Jan 12, 2012 Jan NEWA

FAMOTIDINE

INJECTABLE; INJECTION

FAMOTIDINE

>D>	AP	+	BAXTER HLTHCARE	10MG/ML	A075488	001	Apr 16, 2001	Jul	CAHN
>D>	AP	+		10MG/ML	A075799	001	Apr 30, 2002	Jul	CAHN
>A>	AP	+	HIKMA MAPLE	10MG/ML	A075488	001	Apr 16, 2001	Jul	CAHN
>A>	AP	+		10MG/ML	A075799	001	Apr 30, 2002	Jul	CAHN
			FAMOTIDINE PRESERVATIVE FREE						
>D>	AP	+	BAXTER HLTHCARE	10MG/ML	A075789	001	Apr 30, 2002	Jul	CAHN
>D>	AP	+		10MG/ML	A075486	001	Apr 16, 2001	Jul	CAHN
>A>	AP	+	HIKMA MAPLE	10MG/ML	A075486	001	Apr 16, 2001	Jul	CAHN
>A>	AP	+		10MG/ML	A075789	001	Apr 30, 2002	Jul	CAHN
			TABLET; ORAL						
			PEPCID						
AB			MARATHON PHARMS	20MG	N019462	001	Oct 15, 1986	Jan	CAHN
AB		+		40MG	N019462	002	Oct 15, 1986	Jan	CAHN
			TABLET, ORALLY DISINTEGRATING; ORAL						
			FLUXID						
	@	UCB INC		20MG	N021712	001	Sep 24, 2004	Jan	CAHN
	@			40MG	N021712	002	Sep 24, 2004	Jan	CAHN

FEBUXOSTAT

TABLET; ORAL

ULORIC

	TAKEDA PHARMS USA	40MG	N021856	001	Feb 13, 2009	Apr	CAHN
	+	80MG	N021856	002	Feb 13, 2009	Apr	CAHN

FELODIPINE

TABLET, EXTENDED RELEASE; ORAL

FELODIPINE

>A>	AB	WOCKHARDT LTD	2.5MG	A091484	001	Aug 15, 2012	Jul	NEWA
>A>	AB		5MG	A091484	002	Aug 15, 2012	Jul	NEWA
>A>	AB		10MG	A091484	003	Aug 15, 2012	Jul	NEWA

FENOFIBRATE

CAPSULE; ORAL

ANTARA (MICRONIZED)

AB	LUPIN ATLANTIS	43MG	N021695	001	Nov 30, 2004	Feb	CFTG
AB	+	130MG	N021695	003	Nov 30, 2004	Feb	CFTG
FENOFIBRATE (MICRONIZED)							
AB	DR REDDYS LABS SA	43MG	A090859	001	Mar 01, 2012	Feb	NEWA
AB		130MG	A090859	002	Mar 01, 2012	Feb	NEWA

TABLET; ORAL

FENOFIBRATE

AB	IMPAK LABS	160MG	A076509	002	Mar 26, 2008	May	CRLD
AB	+	160MG	A076509	002	Mar 26, 2008	Mar	CRLD
AB	TEVA	54MG	A076433	001	May 13, 2005	May	CMFD
	@	54MG	A076433	001	May 13, 2005	Mar	DISC
AB	+	160MG	A076433	002	May 13, 2005	May	CMFD
	@	160MG	A076433	002	May 13, 2005	Mar	DISC
AB	VALEANT INTL	48MG	A090715	001	Apr 05, 2012	Mar	NEWA
AB		145MG	A090715	002	Apr 05, 2012	Mar	NEWA

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL							
FENTANYL-100							
AB	AVEVA	100MCG/HR	A077449	004	Oct 20, 2008	Jun	CAHN
FENTANYL-25							
AB	AVEVA	25MCG/HR	A077449	001	Oct 20, 2008	Jun	CAHN
FENTANYL-50							
AB	AVEVA	50MCG/HR	A077449	002	Oct 20, 2008	Jun	CAHN
FENTANYL-75							
AB	AVEVA	75MCG/HR	A077449	003	Oct 20, 2008	Jun	CAHN
SPRAY; SUBLINGUAL							
SUBSYS							
INSYS THERAP							
		0.1MG	N202788	001	Jan 04, 2012	Mar	CPOT
		0.1MCG	N202788	001	Jan 04, 2012	Jan	NEWA
		0.2MG	N202788	002	Jan 04, 2012	Mar	CPOT
		0.2MCG	N202788	002	Jan 04, 2012	Jan	NEWA
+		0.4MG	N202788	003	Jan 04, 2012	Mar	CPOT
+		0.4MCG	N202788	003	Jan 04, 2012	Jan	NEWA
		0.6MG	N202788	004	Jan 04, 2012	Mar	CPOT
		0.6MCG	N202788	004	Jan 04, 2012	Jan	NEWA
		0.8MG	N202788	005	Jan 04, 2012	Mar	CPOT
		0.8MCG	N202788	005	Jan 04, 2012	Jan	NEWA

FENTANYL CITRATE

SPRAY, METERED; NASAL							
LAZANDA							
	ARCHIMEDES	EQ 0.1MG BASE	N022569	001	Jun 30, 2011	Jan	CPOT
+		EQ 0.4MG BASE	N022569	002	Jun 30, 2011	Jan	CPOT
TROCHE/LOZENGE; TRANSMUCOSAL							
FENTANYL CITRATE							
AB	PAR PHARM	EQ 0.2MG BASE	A077312	001	Oct 30, 2009	Feb	CAHN
AB		EQ 0.4MG BASE	A077312	002	Oct 30, 2009	Feb	CAHN
AB		EQ 0.6MG BASE	A077312	003	Oct 30, 2009	Feb	CAHN
AB		EQ 0.8MG BASE	A077312	004	Oct 30, 2009	Feb	CAHN
AB		EQ 1.2MG BASE	A077312	005	Oct 30, 2009	Feb	CAHN
AB		EQ 1.6MG BASE	A077312	006	Oct 30, 2009	Feb	CAHN

FERUMOXIDES

INJECTABLE; INJECTION							
FERIDEX I.V.							
	@ AMAG PHARMS INC	EQ 11.2MG IRON/ML	N020416	001	Aug 30, 1996	Jun	DISC

FEXOFENADINE HYDROCHLORIDE

SUSPENSION; ORAL								
ALLEGRA								
>D>	+	SANOFI AVENTIS US	30MG/5ML	N021963	001	Oct 16, 2006	Jul	CFTG
>A>	AB	+	30MG/5ML	N021963	001	Oct 16, 2006	Jul	CFTG
FEXOFENADINE HYDROCHLORIDE								
>A>	AB	ACTAVIS MID ATLANTIC	30MG/5ML	A201311	001	Jul 25, 2012	Jul	NEWA

FLORBETAPIR F-18

SOLUTION; INTRAVENOUS							
AMYVID							
+	AVID RADIOPHARMS INC	10-50ML (13.5-51MCI/ML)	N202008	003	Apr 06, 2012	Jun	CPOT
+		10-50ML (13.5-51CM/ML)	N202008	003	Apr 06, 2012	Apr	NEWA

SOLUTION; INTRAVENOUS

AMYVID

+ AVID RADIOPHARMS INC 10-30ML (13.5-51MCI/ML)
+ 10ML (13.5-51MCI/ML)

N202008 002 Apr 06, 2012 Apr NEWA
N202008 001 Apr 06, 2012 Apr NEWA

FLUCONAZOLE

INJECTABLE; INJECTION

FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER

AP HIKMA FARMACEUTICA 200MG/100ML (2MG/ML)
AP 400MG/200ML (2MG/ML)

A078764 001 Jan 30, 2012 Jan NEWA
A078764 002 Jan 30, 2012 Jan NEWA

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP HIKMA FARMACEUTICA 200MG/100ML (2MG/ML)
AP 400MG/200ML (2MG/ML)

A078698 001 Jan 30, 2012 Jan NEWA
A078698 002 Jan 30, 2012 Jan NEWA

TABLET; ORAL

FLUCONAZOLE

>D>	AB	AMNEAL PHARM	50MG	
>D>	AB		100MG	
>D>	AB		150MG	
>D>	AB		200MG	
>A>	AB	HARRIS PHARM	50MG	
>A>	AB		100MG	
>A>	AB		150MG	
>A>	AB		200MG	
		@ PLIVA	50MG	
		@	100MG	
		@	150MG	
		@	200MG	
		@ RANBAXY LABS LTD	50MG	
		@	100MG	
		@	150MG	
		@	200MG	

A078423 001	Mar 07, 2011	Jul	CAHN
A078423 002	Mar 07, 2011	Jul	CAHN
A078423 003	Mar 07, 2011	Jul	CAHN
A078423 004	Mar 07, 2011	Jul	CAHN
A078423 001	Mar 07, 2011	Jul	CAHN
A078423 002	Mar 07, 2011	Jul	CAHN
A078423 003	Mar 07, 2011	Jul	CAHN
A078423 004	Mar 07, 2011	Jul	CAHN
A076424 001	Jul 29, 2004	Feb	DISC
A076424 002	Jul 29, 2004	Feb	DISC
A076424 003	Jul 29, 2004	Feb	DISC
A076424 004	Jul 29, 2004	Feb	DISC
A076386 001	Jul 29, 2004	Apr	DISC
A076386 002	Jul 29, 2004	Apr	DISC
A076386 003	Jul 29, 2004	Apr	DISC
A076386 004	Jul 29, 2004	Apr	DISC

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARA

@ GENZYME CORP 50MG/VIAL

N020038 001 Apr 18, 1991 Mar DISC

AP + HOSPIRA 50MG/VIAL

A077790 001 Apr 06, 2007 Mar CRLD

TABLET; ORAL

OFORTA

@ SANOFI AVENTIS US 10MG

N022273 001 Dec 18, 2008 May DISC

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

>D>	AP	BAXTER HLTHCARE	0.5MG/5ML (0.1MG/ML)	
>D>	AP		1MG/10ML (0.1MG/ML)	
>A>	AP	HIKMA MAPLE	0.5MG/5ML (0.1MG/ML)	
>A>	AP		1MG/10ML (0.1MG/ML)	
		@ TEVA PARENTERAL	0.5MG/5ML (0.1MG/ML)	
		@	1MG/10ML (0.1MG/ML)	

A076787 002	Oct 12, 2004	Jul	CAHN
A076787 001	Oct 12, 2004	Jul	CAHN
A076787 002	Oct 12, 2004	Jul	CAHN
A076787 001	Oct 12, 2004	Jul	CAHN
A076589 002	Oct 12, 2004	Jan	DISC
A076589 001	Oct 12, 2004	Jan	DISC

FLUNISOLIDE

AEROSOL, METERED; INHALATION

AEROSPAN HFA

>D> + ACTON PHARMS EQ 78MCG BASE/INH

N021247 001 Jan 27, 2006 Jul CPOT

AEROSOL, METERED; INHALATION

AEROSPAH HFA

>A> + ACTON PHARMS EQ 78MG BASE/INH N021247 001 Jan 27, 2006 Jul CPOT

FLUOCINOLONE ACETONIDE

OIL/DROPS; OTIC

FLUOCINOLONE ACETONIDE

AT IDENTI PHARMS INC 0.01% A091306 001 Oct 17, 2011 Jan CPOT

FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE EMULSIFIED BASE

AB2 FOUGERA PHARMS 0.05% A076586 001 Jun 23, 2004 Jan CAHN
@ G AND W LABS INC 0.05% A074204 001 Jun 13, 1995 May CAHN

OINTMENT; TOPICAL

FLUOCINONIDE

AB FOUGERA PHARMS 0.05% A074905 001 Aug 26, 1997 Jan CAHN

FLUOROURACIL

CREAM; TOPICAL

CARAC

>D> + SANOFI AVENTIS US 0.5% N020985 001 Oct 27, 2000 Jul CAHN

>A> + VALEANT INTL 0.5% N020985 001 Oct 27, 2000 Jul CAHN

FLUOROPLEX

+ AQUA PHARMS 1% N016988 001 Jan CAHN

INJECTABLE; INJECTION

FLUOROURACIL

>A> AP ONCO THERAPIES LTD 500MG/10ML (50MG/ML) A202668 001 Jul 17, 2012 Jul NEWA

>A> AP 1GM/20ML (50MG/ML) A202668 002 Jul 17, 2012 Jul NEWA

>A> AP 2.5GM/50ML (50MG/ML) A202669 001 Jul 17, 2012 Jul NEWA

>A> AP 5GM/100ML (50MG/ML) A202669 002 Jul 17, 2012 Jul NEWA

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL

PROZAC

AB1 ELI LILLY AND CO EQ 10MG BASE N018936 006 Dec 23, 1992 Apr CAHN

AB1 EQ 20MG BASE N018936 001 Dec 29, 1987 Apr CAHN

AB + EQ 40MG BASE N018936 003 Jun 15, 1999 Apr CAHN

@ EQ 60MG BASE N018936 004 Jun 15, 1999 Apr CAHN

SARAFEM

AB2 ELI LILLY AND CO EQ 10MG BASE N018936 007 Jul 06, 2000 Apr CAHN

AB2 + EQ 20MG BASE N018936 008 Jul 06, 2000 Apr CAHN

TABLET; ORAL

SARAFEM

WARNER CHILCOTT LLC EQ 10MG BASE N021860 001 May 19, 2006 Jun CAHN

EQ 15MG BASE N021860 002 May 19, 2006 Jun CAHN

+ EQ 20MG BASE N021860 003 May 19, 2006 Jun CAHN

FLUOXETINE HYDROCHLORIDE; OLANZAPINE

CAPSULE; ORAL

OLANZAPINE AND FLUOXETINE HYDROCHLORIDE

AB TEVA PHARMS EQ 25MG BASE;EQ 6MG BASE A077528 001 Jun 19, 2012 Jun NEWA

AB EQ 25MG BASE;EQ 12MG BASE A077528 002 Jun 19, 2012 Jun NEWA

AB EQ 50MG BASE;EQ 6MG BASE A077528 003 Jun 19, 2012 Jun NEWA

AB EQ 50MG BASE;EQ 12MG BASE A077528 004 Jun 19, 2012 Jun NEWA

CAPSULE; ORAL

SYMBYAX

AB	LILLY	EQ 25MG BASE;EQ 6MG BASE	N021520 002	Dec 24, 2003	Jun	CFTG
AB		EQ 25MG BASE;EQ 12MG BASE	N021520 004	Dec 24, 2003	Jun	CFTG
AB	+	EQ 50MG BASE;EQ 6MG BASE	N021520 003	Dec 24, 2003	Jun	CFTG
AB		EQ 50MG BASE;EQ 12MG BASE	N021520 005	Dec 24, 2003	Jun	CFTG

FLURANDRENOLIDE

CREAM; TOPICAL

CORDRAN SP

>A>	AQUA PHARMS	0.025%	N012806 003	Jul	CAHN
>A>		0.05%	N012806 002	Jul	CAHN
>D>	WATSON PHARMS	0.025%	N012806 003	Jul	CAHN
>D>		0.05%	N012806 002	Jul	CAHN
		0.05%	N012806 002	Apr	CMFD

LOTION; TOPICAL

CORDRAN

>A>	+ AQUA PHARMS	0.05%	N013790 001	Jul	CAHN
>D>	+ WATSON LABS	0.05%	N013790 001	Jul	CAHN

OINTMENT; TOPICAL

CORDRAN

>A>	@ AQUA PHARMS	0.025%	N012806 004	Jul	CAHN
>A>	@	0.05%	N012806 001	Jul	CAHN
>D>	@ WATSON PHARMS	0.025%	N012806 004	Jul	CAHN
>D>	@	0.05%	N012806 001	Jul	CAHN

FLUTAMIDE

CAPSULE; ORAL

FLUTAMIDE

AB	+ IVAX SUB TEVA PHARMS	125MG	A075780 001	Sep 19, 2001	Mar	CRLD
AB	SANDOZ INC	125MG	A075818 001	Sep 18, 2001	Mar	CRLD

FLUTICASONE PROPIONATE

CREAM; TOPICAL

CUTIVATE

AB	+ FOUGERA PHARMS	0.05%	N019958 001	Dec 18, 1990	Jan	CAHN
----	------------------	-------	-------------	--------------	-----	------

FLUTICASONE PROPIONATE

AB	FOUGERA PHARMS	0.05%	A076451 001	May 14, 2004	Jan	CAHN
	@ NESHER PHARMS	0.05%	A076865 001	Sep 10, 2004	Jun	DISC

OINTMENT; TOPICAL

FLUTICASONE PROPIONATE

AB	FOUGERA PHARMS	0.005%	A076300 001	May 14, 2004	Jan	CAHN
----	----------------	--------	-------------	--------------	-----	------

FLUVASTATIN SODIUM

CAPSULE; ORAL

FLUVASTATIN SODIUM

AB	MYLAN PHARMS INC	EQ 20MG BASE	A090595 001	Apr 11, 2012	Mar	NEWA
AB		EQ 40MG BASE	A090595 002	Apr 11, 2012	Mar	NEWA
AB	TEVA PHARMS	EQ 20MG BASE	A078407 001	Jun 12, 2012	May	NEWA
AB		EQ 40MG BASE	A078407 002	Jun 12, 2012	May	NEWA

LESCOL

AB	NOVARTIS	EQ 20MG BASE	N020261 001	Dec 31, 1993	Mar	CFTG
AB	+	EQ 40MG BASE	N020261 002	Dec 31, 1993	Mar	CFTG

FLUVOXAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

LUVOX CR

JAZZ PHARMS 100MG
+ 150MGN022033 001 Feb 28, 2008 Jun CAHN
N022033 002 Feb 28, 2008 Jun CAHNFORMOTEROL FUMARATE; MOMETASONE FUROATE

AEROSOL, METERED; INHALATION

DULERA

+ MERCK SHARP DOHME 0.005MG/INH;0.1MG/INH
+ 0.005MG/INH;0.2MG/INHN022518 001 Jun 22, 2010 Jun CAHN
N022518 002 Jun 22, 2010 Jun CAHNFOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

@ RANBAXY LABS LTD 10MG

A076580 001 Apr 23, 2004 Apr DISC

@ 20MG

A076580 002 Apr 23, 2004 Apr DISC

@ 40MG

A076580 003 Apr 23, 2004 Apr DISC

MONOPRIL

@ BRISTOL MYERS SQUIBB 10MG

N019915 002 May 16, 1991 Mar DISC

@ 20MG

N019915 003 May 16, 1991 Mar DISC

@ 40MG

N019915 004 Mar 28, 1995 Mar DISC

FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

FOSPHENYTOIN SODIUM

>D> AP BAXTER HLTHCARE EQ 50MG PHENYTOIN NA/ML
>A> AP HIKMA MAPLE EQ 50MG PHENYTOIN NA/ML
@ TEVA PARENTERAL EQ 50MG PHENYTOIN NA/MLA077989 001 Aug 06, 2007 Jul CAHN
A077989 001 Aug 06, 2007 Jul CAHN
A076886 001 Aug 06, 2007 Mar DISCFOSPROPOFOL DISODIUM

SOLUTION; INTRAVENOUS

LUSEDRA

>D> + EISAI INC 1050MG/30ML (35MG/ML)
>A> @ 1050MG/30ML (35MG/ML)N022244 001 Dec 12, 2008 Jul DISC
N022244 001 Dec 12, 2008 Jul DISCFUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

@ HOSPIRA 10MG/ML

A070578 001 Jul 08, 1987 Jun DISC

GABAPENTIN

SOLUTION; ORAL

GABAPENTIN

AA ACELLA PHARMS LLC 250MG/5ML
AA AMNEAL PHARMS 250MG/5ML
AA KIEL 250MG/5MLA076403 001 May 01, 2012 Jun CAHN
A202024 001 Mar 23, 2012 Mar NEWA
A076403 001 May 01, 2012 Apr NEWAGANCICLOVIR

CAPSULE; ORAL

GANCICLOVIR

@ RANBAXY LABS LTD 250MG
@ 500MGA076457 001 Jun 27, 2003 Apr DISC
A076457 002 Jun 27, 2003 Apr DISC

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

+ HOSPIRA INC	200MG/5.26ML (38MG/ML)	N200795 001 Aug 04, 2011 Apr CTNA
+	1GM/26.3ML (38MG/ML)	N200795 002 Aug 04, 2011 Apr CTNA
+	2GM/52.6ML (38MG/ML)	N200795 003 Aug 04, 2011 Apr CTNA

GLIMEPIRIDE

TABLET; ORAL

GLIMEPIRIDE

AB AUROBINDO PHARMA LTD	1MG	A202759 001 Jun 29, 2012 Jun NEWA
AB	2MG	A202759 002 Jun 29, 2012 Jun NEWA
AB	4MG	A202759 003 Jun 29, 2012 Jun NEWA
AB MICRO LABS USA	1MG	A091220 001 Jun 29, 2012 Jun NEWA
AB	2MG	A091220 002 Jun 29, 2012 Jun NEWA
	3MG	A091220 003 Jun 29, 2012 Jun NEWA
AB	4MG	A091220 004 Jun 29, 2012 Jun NEWA
	6MG	A091220 005 Jun 29, 2012 Jun NEWA
AB	8MG	A091220 006 Jun 29, 2012 Jun NEWA
@ RANBAXY LABS LTD	1MG	A076875 001 Oct 06, 2005 Apr DISC
@	2MG	A076875 002 Oct 06, 2005 Apr DISC
@	4MG	A076875 003 Oct 06, 2005 Apr DISC
@	8MG	A076875 004 Oct 06, 2005 Apr DISC

GLYCOPYRROLATE

TABLET; ORAL

GLYCOPYRROLATE

AA NEXGEN PHARMA	1.5MG	A091522 001 Mar 12, 2012 Feb NEWA
------------------	-------	-----------------------------------

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE

AP CIPLA LTD	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)	A078262 001 Dec 31, 2007 Mar CAHN
AP	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A078258 001 Jun 30, 2008 Mar CAHN
AP	EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	A078258 002 Jun 30, 2008 Mar CAHN
	GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE	
AP + APP PHARMS LLC	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A078096 001 Jun 30, 2008 Apr CRLD
@ TEVA PARENTERAL	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A077165 001 Dec 31, 2007 Apr DISC

GRISEOFULVIN, MICROCRYSTALLINE

SUSPENSION; ORAL

GRIFULVIN V

AB + VALEANT PHARM NORTH	125MG/5ML	A062483 001 Jan 26, 1984 Feb CAHN
--------------------------	-----------	-----------------------------------

TABLET; ORAL

GRIFULVIN V

>D- @ ORTHO JANSSEN	125MG	A062279 001 Jul CAHN
>D- @	250MG	A062279 002 Jul CAHN
>D- +	500MG	A062279 003 Jul CAHN
>A- @ VALEANT INTL	125MG	A062279 001 Jul CAHN
>A- @	250MG	A062279 002 Jul CAHN
>A- +	500MG	A062279 003 Jul CAHN

HALOBETASOL PROPIONATE

CREAM; TOPICAL
 HALOBETASOL PROPIONATE

AB	FOUGERA PHARMS	0.05%	A077001 001 Dec 16, 2004 Jan CAHN
OINTMENT; TOPICAL			
HALOBETASOL PROPIONATE			
AB	FOUGERA PHARMS	0.05%	A076903 001 Dec 16, 2004 Jan CAHN
	@ VENUS PHARMS	0.05%	A077109 001 Jun 14, 2005 Jun CAHN

HEPARIN SODIUM

INJECTABLE; INJECTION
 HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

>D>	AP	B BRAUN	200 UNITS/100ML	N019953 001 Jul 20, 1992 Jul CRLD
>A>	AP	+	200 UNITS/100ML	N019953 001 Jul 20, 1992 Jul CRLD
HEPARIN SODIUM 10,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER				
@ BAXTER HLTHCARE 2,000 UNITS/100ML N018814 002 Jul 09, 1985 Mar DISC				
HEPARIN SODIUM 20,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER				
@ BAXTER HLTHCARE 4,000 UNITS/100ML N018814 001 Oct 31, 1983 Mar DISC				
HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER				
>D>	AP	B BRAUN	4,000 UNITS/100ML	N019952 001 Jul 20, 1992 Jul CRLD
>A>	AP	+	4,000 UNITS/100ML	N019952 001 Jul 20, 1992 Jul CRLD
HEPARIN SODIUM 25,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER				
@ BAXTER HLTHCARE 5,000 UNITS/100ML N018814 003 Jul 09, 1985 Mar DISC				
@ 10,000 UNITS/100ML N018814 004 Jul 02, 1987 Mar DISC				
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER				
>D>	AP	B BRAUN	5,000 UNITS/100ML	N019952 004 Jul 20, 1992 Jul CRLD
>A>	AP	+	5,000 UNITS/100ML	N019952 004 Jul 20, 1992 Jul CRLD
>D>	AP		10,000 UNITS/100ML	N019952 005 Jul 20, 1992 Jul CRLD
>A>	AP	+	10,000 UNITS/100ML	N019952 005 Jul 20, 1992 Jul CRLD

HEXAMINOLEVULINATE HYDROCHLORIDE

FOR SOLUTION; INTRAVESICAL
 CYSVIEW KIT
 + PHOTOCURE ASA 100MG/VIAL N022555 001 May 28, 2010 May CAHN

HYALURONIDASE

INJECTABLE; INJECTION
 HYDASE
 @ AKORN INC 150 UNITS/ML N021716 001 Oct 25, 2005 Jun DISC

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL
 HYDROCHLOROTHIAZIDE
 AB LANNETT HOLDINGS INC 12.5MG A091662 001 Jan 27, 2012 Jan NEWA

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL
 AVALIDE
 AB + SANOFI AVENTIS 12.5MG;150MG N020758 002 Sep 30, 1997 Mar CFTG
 AB + 12.5MG;300MG N020758 003 Aug 31, 1998 Mar CFTG
 AB + 12.5MG;300MG N020758 003 Aug 31, 1998 Feb CRLD
 IRBESARTAN AND HYDROCHLOROTHIAZIDE
 AB TEVA 12.5MG;150MG A077369 001 Mar 30, 2012 Mar NEWA
 AB 12.5MG;300MG A077369 002 Mar 30, 2012 Mar NEWA
 AB @ 25MG;300MG A077369 003 Mar 30, 2012 Apr DISC

TABLET; ORAL

IRBESARTAN AND HYDROCHLOROTHIAZIDE
TEVA 25MG;300MG

A077369 003 Mar 30, 2012 Mar NEWA

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE

AB	ALEMBIC LTD	12.5MG;50MG	A091617 001	Feb 17, 2012	Feb	NEWA
AB		12.5MG;100MG	A091617 002	Feb 17, 2012	Feb	NEWA
AB		25MG;100MG	A091617 003	Feb 17, 2012	Feb	NEWA
>A>	AB MACLEODS PHARMS LTD	12.5MG;50MG	A202289 001	Aug 09, 2012	Jul	NEWA
>A>	AB	12.5MG;100MG	A202289 002	Aug 09, 2012	Jul	NEWA
>A>	AB	25MG;100MG	A202289 003	Aug 09, 2012	Jul	NEWA

HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

DUTOPROL

	ASTRAZENECA	12.5MG;EQ 25MG TARTRATE	N021956 001	Aug 28, 2006	Jan	CMFD
		12.5MG;EQ 50MG TARTRATE	N021956 002	Aug 28, 2006	Jan	CMFD
+		12.5MG;EQ 100MG TARTRATE	N021956 003	Aug 28, 2006	Apr	CRLD
		12.5MG;EQ 100MG TARTRATE	N021956 003	Aug 28, 2006	Jan	CMFD

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE

AB	MYLAN	50MG;100MG	A076792 003	Aug 20, 2004	Jan	CTEC
AB	SUN PHARM IND	25MG;50MG	A090654 001	Jan 19, 2012	Jan	NEWA
AB		25MG;100MG	A090654 002	Jan 19, 2012	Jan	NEWA
AB		50MG;100MG	A090654 003	Jan 19, 2012	Jan	NEWA

HYDROCHLOROTHIAZIDE; MOXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

@ PADDICK LLC	12.5MG;7.5MG	A090096 001	Sep 25, 2008	Jan	DISC
@	12.5MG;15MG	A090096 002	Sep 25, 2008	Jan	DISC
@	25MG;15MG	A090096 003	Sep 25, 2008	Jan	DISC

HYDROCORTISONE

CREAM; TOPICAL

NUTRACORT

@ DOW PHARM	0.5%	A080442 002		Jun	CAHN
@	1%	A080442 003		Jun	CAHN

LOTION; TOPICAL

CETACORT

@ DOW PHARM	0.5%	A080426 002		Jun	CAHN
@	1%	A080426 001		Jun	CAHN

HYDROCORTISONE

AT	+ FOUGERA PHARMS	2.5%	A040351 001	Jul 25, 2000	Jan	CAHN
	NUTRACORT					

@ DOW PHARM	0.5%	A080443 002		Jun	CAHN
	1%	A080443 003		Jun	CAHN

AT		2.5%	A087644 001	Aug 24, 1982	Jun	CAHN

OINTMENT; TOPICAL

HYDROCORTISONE

AT	+ FOUGERA PHARMS	1%	A080692 001		Jan	CAHN

HYDROCORTISONE ACETATE; UREA

CREAM; TOPICAL
CARMOL HC
AT FOUGERA PHARMS 1%;10% A080505 001 Jan CAHN

HYDROCORTISONE BUTYRATE

CREAM; TOPICAL
LOCOID
AB + PRECISION DERMAT 0.1% N018514 001 Mar 31, 1982 Jun CAHN
LOCOID LIPOCREAM
+ PRECISION DERMAT 0.1% N020769 001 Sep 08, 1997 Jun CAHN
LOTION; TOPICAL
LOCOID
+ PRECISION DERMAT 0.1% N022076 001 May 18, 2007 May CAHN
OINTMENT; TOPICAL
LOCOID
AB + PRECISION DERMAT 0.1% N018652 001 Oct 29, 1982 May CAHN
SOLUTION; TOPICAL
LOCOID
AT + PRECISION DERMAT 0.1% N019116 001 Feb 25, 1987 Jun CAHN

HYDROCORTISONE VALERATE

CREAM; TOPICAL
HYDROCORTISONE VALERATE
AB + TARO 0.2% A075042 001 Aug 25, 1998 Mar CRLD
WESTCORT
@ RANBAXY 0.2% N017950 001 Mar DISC
OINTMENT; TOPICAL
HYDROCORTISONE VALERATE
AB FOUGERA PHARMS 0.2% A075085 001 Jul 31, 2001 Jan CAHN

HYDROFLUMETHIAZIDE

TABLET; ORAL
SALURON
AB + SHIRE LLC 50MG N011949 001 Jan CAHN

HYDROMORPHONE HYDROCHLORIDE

TABLET; ORAL
HYDROMORPHONE HYDROCHLORIDE
@ NESHER PHARMS 2MG A077311 001 Nov 09, 2005 Jun DISC
@ 4MG A077311 002 Nov 09, 2005 Jun DISC
@ 8MG A077311 003 Nov 09, 2005 Jun DISC

HYDROXYZINE HYDROCHLORIDE

SYRUP; ORAL
HYDROXYZINE HYDROCHLORIDE
@ STI PHARMA LLC 10MG/5ML A086880 001 Feb CAHN

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE
>D @ ACTAVIS TOTOWA 10MG A040600 001 Dec 28, 2004 Jul CAHN
>D @ 25MG A040602 001 Dec 28, 2004 Jul CAHN
>D @ 50MG A040604 001 Dec 28, 2004 Jul CAHN
>A @ MIKAH PHARMA 10MG A040600 001 Dec 28, 2004 Jul CAHN
>A @ 25MG A040602 001 Dec 28, 2004 Jul CAHN
>A @ 50MG A040604 001 Dec 28, 2004 Jul CAHN

HYDROXYZINE PAMOATE

>D>	SUSPENSION; ORAL					
>D>	VISTARIL					
>D>	+ PFIZER	EQ 25MG HCL/5ML	N011795 001	Jul	DISC	
>A>	@	EQ 25MG HCL/5ML	N011795 001	Jul	DISC	

IBANDRONATE SODIUM

TABLET; ORAL						
BONIVA						
AB	+ HOFFMANN LA ROCHE	EQ 150MG BASE	N021455 002	Mar 24, 2005	Mar	CFTG
IBANDRONATE SODIUM						
AB	APOTEX INC	EQ 150MG BASE	A078948 001	Mar 19, 2012	Mar	NEWA
AB	DR REDDYS LABS LTD	EQ 150MG BASE	A078997 001	Apr 30, 2012	Apr	NEWA
>A>	AB MUTUAL PHARM CO INC	EQ 150MG BASE	A078996 001	Aug 15, 2012	Jul	NEWA
AB	MYLAN PHARMS INC	EQ 150MG BASE	A078995 001	Mar 19, 2012	Mar	NEWA
AB	ORCHID HLTHCARE	EQ 150MG BASE	A078998 001	Mar 19, 2012	Mar	NEWA
AB	WATSON LABS INC	EQ 150MG BASE	A079003 001	Mar 20, 2012	Mar	NEWA

IBUPROFEN

TABLET; ORAL						
IBUPROFEN						
	@ NORTHLSTAR HLTHCARE	400MG	A078132 001	Sep 10, 2007	Jun	DISC
	@	600MG	A078132 002	Sep 10, 2007	Jun	DISC
	@	800MG	A078132 003	Sep 10, 2007	Jun	DISC
	@ OHM LABS	400MG	A070818 001	Dec 26, 1985	Jun	DISC

IBUPROFEN LYSINE

INJECTABLE; INTRAVENOUS						
NEOPROFEN						
+ LUNDBECK LLC	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	N021903 001	Apr 13, 2006	Mar	CAHN	

>A>	<u>ICOSAPENT ETHYL</u>					
>A>	CAPSULE; ORAL					
>A>	VASCEPA					
>A>	+ AMARIN PHARMA INC	1GM	N202057 001	Jul 26, 2012	Jul	NEWA

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION						
IDARUBICIN HYDROCHLORIDE						
@ TEVA PARENTERAL	5MG/VIAL	A065037 003	May 01, 2002	Mar	DISC	
@	10MG/VIAL	A065037 002	May 01, 2002	Mar	DISC	
@	20MG/VIAL	A065037 001	May 01, 2002	Mar	DISC	

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL						
IMIPRAMINE HYDROCHLORIDE						
AB	PROSAM LABS	10MG	A040753 001	Feb 28, 2008	Feb	CMFD
AB		25MG	A040752 001	Feb 28, 2008	Feb	CMFD
AB		50MG	A040751 001	Feb 28, 2008	Feb	CMFD

IMIQUIMOD

CREAM; TOPICAL						
IMIQUIMOD						
AB	APOTEX INC	5%	A091308 001	Apr 06, 2012	Mar	NEWA

CREAM; TOPICAL

IMIQUIMOD

AB	FOUGERA PHARMS	5%	A078548 001 Feb 25, 2010 Jan CAHN
AB	GLENMARK GENERICS	5%	A201994 001 Mar 06, 2012 Feb NEWA

INDOMETHACIN SODIUM

INJECTABLE; INJECTION

INDOCIN

AP	+	LUNDBECK LLC	EQ 1MG BASE/VIAL	N018878 001 Jan 30, 1985 Mar CAHN
----	---	--------------	------------------	-----------------------------------

INGENOL MEBUTATE

GEL; TOPICAL

PICATO

LEO PHARMA AS	0.015%	N202833 001 Jan 23, 2012 Jan NEWA
+	0.05%	N202833 002 Jan 23, 2012 Jan NEWA

INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT

INJECTABLE; INJECTION

HUMALOG MIX 50/50 PEN

@ LILLY	50 UNITS/ML;50 UNITS/ML	N021018 003 Dec 22, 1999 Jun DISC
---------	-------------------------	-----------------------------------

HUMALOG MIX 75/25 PEN

@ LILLY	75 UNITS/ML;25 UNITS/ML	N021017 003 Dec 22, 1999 Jun DISC
---------	-------------------------	-----------------------------------

INSULIN LISPRO RECOMBINANT

INJECTABLE; INJECTION

HUMALOG PEN

@ LILLY	100 UNITS/ML	N020563 002 Aug 06, 1998 Jun DISC
---------	--------------	-----------------------------------

IRBESARTAN

TABLET; ORAL

AVAPRO

AB	SANOFI AVENTIS US	75MG	N020757 001 Sep 30, 1997 Mar CFTG
AB		150MG	N020757 002 Sep 30, 1997 Mar CFTG
AB	+	300MG	N020757 003 Sep 30, 1997 Mar CFTG
	IRBESARTAN		
AB	TEVA PHARMS	75MG	A077159 001 Mar 30, 2012 Mar NEWA
AB		150MG	A077159 002 Mar 30, 2012 Mar NEWA
AB		300MG	A077159 003 Mar 30, 2012 Mar NEWA

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

AP	CIPILA LTD	40MG/2ML (20MG/ML)	A077219 001 Feb 20, 2008 Mar CAHN
AP		100MG/5ML (20MG/ML)	A077219 002 Feb 20, 2008 Mar CAHN
AP	EMCURE PHARMS LTD	40MG/2ML (20MG/ML)	A200771 001 Feb 14, 2012 Jan NEWA
AP		100MG/5ML (20MG/ML)	A200771 002 Feb 14, 2012 Jan NEWA
AP	HISUN PHARM HANGZHOU	40MG/2ML (20MG/ML)	A090016 001 Jan 28, 2009 Mar CAHN
AP		100MG/5ML (20MG/ML)	A090016 002 Jan 28, 2009 Mar CAHN
AP	TEVA PARENTERAL	40MG/2ML (20MG/ML)	A090101 002 Feb 27, 2008 Jun NEWA
AP		100MG/5ML (20MG/ML)	A090101 003 Feb 27, 2008 Jun NEWA
AP		500MG/25ML (20MG/ML)	A090101 001 Nov 26, 2008 Jun CMFD
	@	500MG/25ML (20MG/ML)	A090101 001 Nov 26, 2008 Mar DISC

ISOFLURANE

LIQUID; INHALATION
FORANE

>A>	AN	+	BAXTER HLTHCARE	99.9%	N017624 001	Jul	CAHN
>D>	AN	+	BAXTER HLTHCARE CORP	99.9%	N017624 001	Jul	CAHN

ISOSORBIDE MONONITRATE

TABLET, EXTENDED RELEASE; ORAL
ISOSORBIDE MONONITRATE

AB	ALKERMES GAINESVILLE	60MG	A075041 001	Sep 22, 1998	Jun	CAHN
	@ SKYEPHARMA AG	60MG	A075166 001	Oct 07, 1999	Jan	DISC

ISOTRETINOIN

CAPSULE; ORAL
ABSORICA

CIPHER	10MG	N021951 001	May 25, 2012	May	NEWA
	20MG	N021951 002	May 25, 2012	May	NEWA
	30MG	N021951 003	May 25, 2012	May	NEWA
	40MG	N021951 004	May 25, 2012	May	NEWA

MYORISAN

AB	DOUGLAS PHARMS	10MG	A076485 001	Jan 19, 2012	Jan	NEWA
AB		20MG	A076485 002	Jan 19, 2012	Jan	NEWA
AB		40MG	A076485 003	Jan 19, 2012	Jan	NEWA

ITRACONAZOLE

CAPSULE; ORAL
ITRACONAZOLE

>A>	AB	MYLAN PHARMS INC	100MG	A200463 001	Jul 20, 2012	Jul	NEWA
-----	----	------------------	-------	-------------	--------------	-----	------

IVACAFTOR

TABLET; ORAL
KALYDECO

+	VERTEX PHARMS	150MG	N203188 001	Jan 31, 2012	Jan	NEWA
---	---------------	-------	-------------	--------------	-----	------

IVERMECTIN

LOTION; TOPICAL
SKLICE

+	SANOFI PASTEUR	0.5%	N202736 001	Feb 07, 2012	Feb	NEWA
---	----------------	------	-------------	--------------	-----	------

KETOCONAZOLE

CREAM; TOPICAL
KETOCONAZOLE

AB	FOUGERA PHARMS	2%	A076294 001	Apr 28, 2004	Jan	CAHN
----	----------------	----	-------------	--------------	-----	------

KETOPROFEN

CAPSULE; ORAL
KETOPROFEN

>A>		DORADO PHARMA	25MG	A074014 001	Jan 29, 1993	Jul	CAHN
>A>	AB		50MG	A074014 002	Jan 29, 1993	Jul	CAHN
>A>	AB		75MG	A074014 003	Jan 29, 1993	Jul	CAHN
>D>		HERITAGE PHARMS INC	25MG	A074014 001	Jan 29, 1993	Jul	CAHN
>D>	AB		50MG	A074014 002	Jan 29, 1993	Jul	CAHN
>D>	AB		75MG	A074014 003	Jan 29, 1993	Jul	CAHN

CAPSULE, EXTENDED RELEASE; ORAL
KETOPROFEN

AB ALKERMES GAINESVILLE 200MG A074879 001 Dec 10, 1997 Jun CAHN

LABETALOL HYDROCHLORIDE

INJECTABLE; INJECTION
LABETALOL HYDROCHLORIDE

AP GLAND PHARMA LTD 5MG/ML A090699 001 Apr 03, 2012 Mar NEWA
TABLET; ORAL
LABETALOL HYDROCHLORIDE
AB PAR FORM 100MG A200908 001 Jul 10, 2012 Jun NEWA
AB 200MG A200908 002 Jul 10, 2012 Jun NEWA
AB 300MG A200908 003 Jul 10, 2012 Jun NEWA

LACTULOSE

FOR SOLUTION; ORAL
LACTULOSE

+ CUMBERLAND PHARMS 10GM/PACKET A074712 001 Dec 10, 1997 Jun CAHN
+ 20GM/PACKET A074712 002 Dec 10, 1997 Jun CAHN

SOLUTION; ORAL
LACTULOSE

AA FRESENIUS KABI 10GM/15ML A090503 001 Jan 25, 2012 Jan NEWA
SOLUTION; ORAL, RECTAL
LACTULOSE
AA FRESENIUS KABI 10GM/15ML A090502 001 Jan 25, 2012 Jan NEWA

LAMIVUDINE; ZIDOVUDINE

TABLET; ORAL
LAMIVUDINE AND ZIDOVUDINE

AB AUROBINDO PHARMA LTD 150MG;300MG A202418 001 May 15, 2012 Apr NEWA
AB LUPIN LTD 150MG;300MG A090246 001 May 15, 2012 Apr NEWA

LAMOTRIGINE

TABLET; ORAL
LAMOTRIGINE

AB GLENMARK GENERICS 25MG A090169 001 May 12, 2012 Apr NEWA
AB 100MG A090169 002 May 12, 2012 Apr NEWA
AB 150MG A090169 003 May 12, 2012 Apr NEWA
AB 200MG A090169 004 May 12, 2012 Apr NEWA

LANREOTIDE ACETATE

INJECTABLE; SUBCUTANEOUS
SOMATULINE DEPOT

+ IPSSEN INC EQ 60MG BASE N022074 001 Aug 30, 2007 Apr CAHN
+ EQ 90MG BASE N022074 002 Aug 30, 2007 Apr CAHN
+ EQ 120MG BASE N022074 003 Aug 30, 2007 Apr CAHN
+ IPSSEN PHARMA EQ 60MG BASE N022074 001 Aug 30, 2007 May CAHN
+ EQ 90MG BASE N022074 002 Aug 30, 2007 May CAHN
+ EQ 120MG BASE N022074 003 Aug 30, 2007 May CAHN

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL
PREVACID

AB TAKEDA PHARMS USA 15MG N020406 001 May 10, 1995 Mar CAHN
AB + 30MG N020406 002 May 10, 1995 Mar CAHN

TABLET, DELAYED RELEASE, ORALLY DISINTEGRATING; ORAL

LANSOPRAZOLE

@ TEVA PHARMS 15MG

A078730 001 Oct 15, 2010 Mar DISC

@ 30MG

A078730 002 Oct 15, 2010 Mar DISC

PREVACID

TAKEDA PHARMS USA 15MG

N021428 001 Aug 30, 2002 Mar CTEC

+ 30MG

N021428 002 Aug 30, 2002 Mar CTEC

LANTHANUM CARBONATE

TABLET, CHEWABLE; ORAL

FOSRENOL

@ SHIRE LLC EQ 250MG BASE

N021468 001 Oct 26, 2004 Feb CAHN

EQ 500MG BASE

N021468 002 Oct 26, 2004 Feb CAHN

EQ 750MG BASE

N021468 003 Nov 23, 2005 Feb CAHN

+ EQ 1GM BASE

N021468 004 Nov 23, 2005 Feb CAHN

LETROZOLE

TABLET; ORAL

LETROZOLE

AB APOTEX INC 2.5MG

A091303 001 Apr 19, 2012 Apr NEWA

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

@ TEVA PARENTERAL

EQ 50MG BASE/VIAL

A081278 001 Sep 28, 1993 Mar DISC

LEUPROLIDE ACETATE

INJECTABLE; INJECTION

LUPRON DEPOT

+ ABBOTT ENDOCRINE 3.75MG/VIAL

N020011 001 Oct 22, 1990 May CAHN

+ 7.5MG/VIAL

N019732 001 Jan 26, 1989 May CAHN

+ 11.25MG/VIAL

N020708 001 Mar 07, 1997 May CAHN

LEVETIRACETAM

INJECTABLE; IV (INFUSION)

LEVETIRACETAM

AP APP PHARMS LLC 500MG/5ML (100MG/ML)

A090813 001 May 26, 2010 May CAHN

AP HOSPIRA INC 500MG/ML (100MG/ML)

A202869 001 Apr 06, 2012 Mar NEWA

>D> AP INNOPHARMA LLC 500MG/5ML (100MG/ML)

A091485 001 Aug 05, 2011 Jul CAHN

AP PHARMAFORCE 500MG/5ML (100MG/ML)

A202143 001 Jan 31, 2012 Jan NEWA

>A> AP X GEN PHARMS 500MG/5ML (100MG/ML)

A091485 001 Aug 05, 2011 Jul CAHN

SOLUTION; ORAL

LEVETIRACETAM

AA HI-TECH PHARMACAL 100MG/ML

A090601 001 Feb 28, 2012 Feb NEWA

AA VINTAGE PHARMS 100MG/ML

A090079 001 Apr 11, 2012 Mar NEWA

TABLET; ORAL

LEVETIRACETAM

>D> AB BIOKEY 500MG

A090906 001 Nov 05, 2010 Jul CAHN

>A> AB LOTUS PHARM CO LTD 500MG

A090906 001 Nov 05, 2010 Jul CAHN

AB WATSON LABS INC 250MG

A078797 002 Jan 15, 2009 Apr NEWA

AB 500MG

A078797 003 Jan 15, 2009 Apr NEWA

AB 750MG

A078797 004 Jan 15, 2009 Apr NEWA

TABLET, EXTENDED RELEASE; ORAL

LEVETIRACETAM

AB BOCA PHARMA 500MG

A201464 001 May 25, 2012 May NEWA

AB 750MG

A201464 002 May 25, 2012 May NEWA

TABLET, EXTENDED RELEASE; ORAL
LEVETIRACETAM

AB	TORRENT PHARMS LTD	500MG	A091338 001 May 29, 2012 May NEWA
AB		750MG	A091338 002 May 29, 2012 May NEWA

LEVOCETIRIZINE DIHYDROCHLORIDE

TABLET; ORAL
LEVOCETIRIZINE DIHYDROCHLORIDE

AB	HETERO LABS LTD III	5MG	A091264 001 Jun 29, 2012 Jun NEWA
----	---------------------	-----	-----------------------------------

LEVOFLOXACIN

TABLET; ORAL
LEVOFLOXACIN

AB	CIPLA LTD	250MG	A076890 001 Mar 30, 2012 Mar NEWA
AB		500MG	A076890 002 Mar 30, 2012 Mar NEWA
AB		750MG	A076890 003 Mar 30, 2012 Mar NEWA
AB	MACLEODS PHARMS LTD	250MG	A200839 001 Mar 22, 2012 Mar NEWA
AB		500MG	A200839 002 Mar 22, 2012 Mar NEWA
AB		750MG	A200839 003 Mar 22, 2012 Mar NEWA
AB	ORCHID HLTHCARE	250MG	A202200 001 Jan 30, 2012 Jan NEWA
AB		500MG	A202200 002 Jan 30, 2012 Jan NEWA
AB		750MG	A202200 003 Jan 30, 2012 Jan NEWA

LEVOLEUCOVORIN CALCIUM

SOLUTION; IV (INFUSION)
FUSILEV
@ SPECTRUM PHARMS EQ 250MG BASE/25ML (EQ 10MG
BASE/ML) N020140 003 Apr 29, 2011 Jun DISC

LEVONORGESTREL

TABLET; ORAL
LEVONORGESTREL

AB	WATSON LABS INC	1.5MG	A200670 001 Jul 12, 2012 Jun NEWA
	PLAN B		
	@ TEVA BRANDED PHARM	0.75MG	N021045 001 Jul 28, 1999 May CAHN
AB	+	0.75MG	N021045 002 Aug 24, 2006 May CAHN
	PLAN B ONE-STEP		
AB	+	DURAMED	1.5MG N021998 001 Jul 10, 2009 Jun CFTG

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION
XYLOCAINE DENTAL
@ DENTSPLY PHARM 2% N021380 001 May DISC

LIDOCAINE; PRILOCaine

CREAM; TOPICAL
LIDOCAINE AND PRILOCaine
AB FOUGERA PHARMS 2.5%;2.5% A076453 001 Aug 18, 2003 Jan CAHN

LIDOCAINE; TETRACAINe

CREAM; TOPICAL
PLIAGLIS
+ GALDERMA LABS LP 7%;7% N021717 001 Jun 29, 2006 Jun CTNA

LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

JENTADUETO

BOEHRINGER INGELHEIM	2.5MG;500MG
	2.5MG;850MG
+	2.5MG;1GM

N201281 001	Jan 30, 2012	Jan	NEWA
N201281 002	Jan 30, 2012	Jan	NEWA
N201281 003	Jan 30, 2012	Jan	NEWA

LINEZOLID

INJECTABLE; INJECTION

LINEZOLID

AP	TEVA PHARMS	200MG/100ML
	ZYVOX	
AP	+ PHARMACIA AND UPJOHN	200MG/100ML

A200222 001	Jun 22, 2012	Jun	NEWA
N021131 001	Apr 18, 2000	Jun	CFTG

LITHIUM CARBONATE

TABLET, EXTENDED RELEASE; ORAL

LITHIUM CARBONATE

AB	MYLAN PHARMS INC	300MG
>A>	AB	450MG

A202288 001	Jun 29, 2012	Jun	NEWA
A202219 001	Aug 08, 2012	Jul	NEWA

LOMUSTINE

CAPSULE; ORAL

CEENU

BRISTOL MYERS SQUIBB	10MG
@	10MG
	40MG
@	40MG
+	100MG
@	100MG

N017588 001	Jun	CMFD
N017588 001	May	DISC
N017588 002	Jun	CMFD
N017588 002	May	DISC
N017588 003	Jun	CMFD
N017588 003	May	DISC

LORAZEPAM

CONCENTRATE; ORAL

LORAZEPAM

AA	HI-TECH PHARMA CO	2MG/ML
	INJECTABLE; INJECTION	
	LORAZEPAM	
@	BEDFORD	2MG/ML
@		4MG/ML
@	HOSPIRA	2MG/ML

A200169 001	Jan 30, 2012	Jan	NEWA
A077076 001	Jul 13, 2005	Jun	DISC
A077076 002	Jul 13, 2005	Jun	DISC
A074300 001	Apr 12, 1994	Jun	DISC

LORCASERIN HYDROCHLORIDE

TABLET; ORAL

BELVIQ

>D>	+ ARENA PHARMS INC	10MG
	+	10MG
>A>	+ EISAI INC	10MG

N022529 001	Jun 27, 2012	Jul	CAHN
N022529 001	Jun 27, 2012	Jun	NEWA
N022529 001	Jun 27, 2012	Jul	CAHN

LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM

AB	MACLEODS PHARMS LTD	25MG
AB		50MG
AB		100MG

A202230 001	May 30, 2012	May	NEWA
A202230 002	May 30, 2012	May	NEWA
A202230 003	May 30, 2012	May	NEWA

LOXAPINE SUCCINATE

CAPSULE; ORAL							
LOXAPINE SUCCINATE							
>D>	@ ACTAVIS TOTOWA	EQ 5MG BASE	A076868	001	Aug 04, 2005	Jul	CAHN
>D>	@	EQ 10MG BASE	A076868	002	Aug 04, 2005	Jul	CAHN
>D>	@	EQ 25MG BASE	A076868	003	Aug 04, 2005	Jul	CAHN
>D>	@	EQ 50MG BASE	A076868	004	Aug 04, 2005	Jul	CAHN
>A>	@ MIKAH PHARMA	EQ 5MG BASE	A076868	001	Aug 04, 2005	Jul	CAHN
>A>	@	EQ 10MG BASE	A076868	002	Aug 04, 2005	Jul	CAHN
>A>	@	EQ 25MG BASE	A076868	003	Aug 04, 2005	Jul	CAHN
>A>	@	EQ 50MG BASE	A076868	004	Aug 04, 2005	Jul	CAHN

LUCINACTANT

SUSPENSION; INTRATRACHEAL							
SURFAXIN							
+ DISCOVERY LABS	8.5ML	N021746	001	Mar 06, 2012	Apr	NEWA	

LURASIDONE HYDROCHLORIDE

TABLET; ORAL							
LATUDA							
SUNOVION PHARMS INC	20MG	N200603	003	Dec 07, 2011	Apr	NEWA	
+ +	40MG	N200603	001	Oct 28, 2010	Apr	CRLD	
	80MG	N200603	002	Oct 28, 2010	Apr	CRLD	
	120MG	N200603	004	Apr 26, 2012	Apr	NEWA	

LUTROPIN ALFA

INJECTABLE; SUBCUTANEOUS							
LUVERIS							
@ EMD SERONO	75 IU/VIAL	N021322	001	Oct 08, 2004	Jun	DISC	

MAFENIDE ACETATE

CREAM; TOPICAL							
SULFAMYLYON							
+ MYLAN LLC	EQ 85MG BASE/GM	N016763	001		Jun	CAHN	
FOR SOLUTION; TOPICAL							
SULFAMYLYON							
+ MYLAN LLC	5%	N019832	003	Jun 05, 1998	Jun	CAHN	

MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE

TABLET; ORAL							
MAGNESIUM HYDROXIDE AND OMEPRAZOLE AND SODIUM BICARBONATE							
@ SANTARUS	343MG;20MG;750MG	N022456	001	Dec 04, 2009	Jun	DISC	
@	343MG;40MG;750MG	N022456	002	Dec 04, 2009	Jun	DISC	

MAGNESIUM SULFATE

INJECTABLE; INJECTION							
MAGNESIUM SULFATE IN PLASTIC CONTAINER							
HOSPIRA	20GM/500ML (40MG/ML)	N020309	004	Jan 18, 1995	Jan	NEWA	
	40GM/1000ML(40MG/ML)	N020309	005	Jan 18, 1995	Jan	NEWA	

MALATHION

LOTION; TOPICAL							
MALATHION							
AT MYLAN PHARMS INC	0.5%	A078743	001	Mar 06, 2009	Feb	CAHN	
AT SUVEN LIFE	0.5%	A091559	001	May 23, 2012	May	NEWA	

MANNITOL

POWDER; INHALATION
 ARIDOL KIT
 + PHARMAXIS LTD N/A,5MG,10MG,20MG,40MG N022368 001 Oct 05, 2010 May CAHN

MECASERMIN RECOMBINANT

INJECTABLE; SUBCUTANEOUS
 INCRELEX
 + IPSEN INC 40MG/4ML (10MG/ML) N021839 001 Aug 30, 2005 May CAHN

MECLIZINE HYDROCHLORIDE

TABLET; ORAL
 MECLIZINE HYDROCHLORIDE
 AA EPIC PHARMA LLC 12.5MG A200294 001 Apr 13, 2012 Mar NEWA
 AA 25MG A200294 002 Apr 13, 2012 Mar NEWA

MEDROXYPROGESTERONE ACETATE

INJECTABLE; INJECTION
 MEDROXYPROGESTERONE ACETATE
 @ TEVA PARENTERAL 150MG/ML A076552 001 Oct 27, 2004 Apr DISC

MEMANTINE HYDROCHLORIDE

TABLET; ORAL
 MEMANTINE HYDROCHLORIDE
 @ ORCHID HLTHCARE 5MG A090044 001 Mar 12, 2012 Mar DISC
 AB 5MG A090044 001 Mar 12, 2012 Feb NEWA
 @ 10MG A090044 002 Mar 12, 2012 Mar DISC
 AB 10MG A090044 002 Mar 12, 2012 Feb NEWA
 NAMENDA
 FOREST LABS 5MG N021487 001 Oct 16, 2003 Mar CTEC
 AB 5MG N021487 001 Oct 16, 2003 Feb CTEC
 + 10MG N021487 002 Oct 16, 2003 Mar CTEC
 AB + 10MG N021487 002 Oct 16, 2003 Feb CTEC

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION
 POLOCAINE
 @ DENTSPLY PHARM 3% A088653 001 Aug 21, 1984 May DISC

MEQUINOL; TRETINOIN

SOLUTION; TOPICAL
 SOLAGE
 @ AQUA PHARMS 2%;0.01% N020922 001 Dec 10, 1999 Jun DISC

MESALAMINE

SUPPOSITORY; RECTAL
 CANASA
 >A> @ APTALIS PHARMA US 500MG N021252 001 Jan 05, 2001 Jul CAHN
 >A> 1GM N021252 002 Nov 05, 2004 Jul CAHN
 >D> @ AXCAN 500MG N021252 001 Jan 05, 2001 Jul CAHN
 >D> + 1GM N021252 002 Nov 05, 2004 Jul CAHN
 ROWASA
 @ MEDA PHARMS 500MG N019919 001 Dec 18, 1990 Jan CAHN

MESNA

INJECTABLE; INTRAVENOUS

MESNA

AP MYLAN INSTITUTIONAL 100MG/ML

A076488 001 Mar 08, 2012 Feb NEWA

METFORMIN HYDROCHLORIDE

TABLET; ORAL

METFORMIN HYDROCHLORIDE

AB MARKSANS PHARMA 500MG

A090888 001 Mar 12, 2012 Feb NEWA

AB 850MG

A090888 002 Mar 12, 2012 Feb NEWA

AB 1GM

A090888 003 Mar 12, 2012 Feb NEWA

TABLET, EXTENDED RELEASE; ORAL

METFORMIN HYDROCHLORIDE

>A> AB1 AUROBINDO PHARMA LTD 500MG

A079118 001 Jul 20, 2012 Jul NEWA

>A> AB 750MG

A079118 002 Jul 20, 2012 Jul NEWA

AB1 INVENTIA HLTHCARE 500MG

A201991 001 Jan 18, 2012 Jan NEWA

>A> AB2 MYLAN PHARMS INC 500MG

A200690 001 Aug 01, 2012 Jul NEWA

>A> AB 1GM

A200690 002 Aug 01, 2012 Jul NEWA

@ RANBAXY LABS LTD 500MG

A076413 001 Jun 18, 2004 Apr DISC

@ 750MG

A077211 001 Jun 29, 2005 Apr DISC

METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

KOMBIGLYZE XR

BRISTOL MYERS SQUIBB 500MG;EQ 5MG BASE

N200678 001 Nov 05, 2010 Apr CAIN

1GM;EQ 2.5MG BASE

N200678 003 Nov 05, 2010 Apr CAIN

+ 1GM;EQ 5MG BASE

N200678 002 Nov 05, 2010 Apr CAIN

METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE

TABLET; ORAL

JANUMET

MERCK SHARP DOHME 500MG;EQ 50MG BASE

N022044 001 Mar 30, 2007 May CAHN

+ 1GM;EQ 50MG BASE

N022044 002 Mar 30, 2007 May CAHN

TABLET, EXTENDED RELEASE; ORAL

JANUMET XR

MERCK SHARP DOHME 500MG;EQ 50MG BASE

N202270 001 Feb 02, 2012 Feb NEWA

1GM;EQ 50MG BASE

N202270 002 Feb 02, 2012 Feb NEWA

+ 1GM;EQ 100MG BASE

N202270 003 Feb 02, 2012 Feb NEWA

METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

AB BOCA PHARMA 5MG

A202068 001 Mar 07, 2012 Feb NEWA

AB 10MG

A202068 002 Mar 07, 2012 Feb NEWA

METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL

AA SOLCO HLTHCARE 500MG

A086989 001

Mar CMFD

AA 750MG

A086988 001

Mar CMFD

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE PRESERVATIVE FREE

AP APP PHARMS LLC EQ 1GM BASE/VIAL

A040266 001 Feb 26, 1999 Jan CMFD

INJECTABLE; INJECTION

METHOTREXATE PRESERVATIVE FREE

AP	PHARMACHEMIE BV	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	A200171 001 Feb 27, 2012 Feb NEWA
METHOTREXATE SODIUM			
AP	+ BEDFORD	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	A089341 001 Sep 16, 1986 Feb CTEC
METHOTREXATE SODIUM PRESERVATIVE FREE			
AP	+ MYLAN INSTITUTIONAL	EQ 50MG BASE/2ML (EQ 25MG BASE/ML)	A040767 001 Apr 30, 2007 Feb CAHN
AP	ONCO THERAPIES LTD	EQ 50MG BASE/2ML (EQ 25MG BASE/ML)	A201529 001 Mar 29, 2012 Mar NEWA
AP		EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	A201529 002 Mar 29, 2012 Mar NEWA
AP		EQ 200MG BASE/8ML (EQ 25MG BASE/ML)	A201529 003 Mar 29, 2012 Mar NEWA
AP		EQ 250MG BASE/10ML (EQ 25MG BASE/ML)	A201529 004 Mar 29, 2012 Mar NEWA
AP		EQ 1GM BASE/40ML (EQ 25MG BASE/ML)	A201530 001 Mar 29, 2012 Mar NEWA

METHYCLOTHIAZIDE

TABLET; ORAL

ENDURON

@ ABBOTT LABS PHARM	2.5MG	N012524 001 Mar DISC
@	5MG	N012524 004 Mar DISC
METHYCLOTHIAZIDE		
AB + MYLAN PHARMS INC	5MG	A087672 001 Aug 17, 1982 Mar CRLD

METHYLDOPATE HYDROCHLORIDE

INJECTABLE; INJECTION

METHYLDOPATE HYDROCHLORIDE

@ TEVA PARENTERAL 50MG/ML

A072974 001 Nov 22, 1991 Mar DISC

METHYLNALTREXONE BROMIDE

INJECTABLE; SUBCUTANEOUS

RELISTOR

SALIX PHARMS	8MG/0.4ML	N021964 002 Sep 27, 2010 May CAHN
+	12MG/0.6ML (12MG/0.6ML)	N021964 003 Apr 24, 2008 Jun NEWA
SOLUTION; SUBCUTANEOUS		
RELISTOR		
+ SALIX PHARMS	12MG/0.6ML (12MG/0.6ML)	N021964 001 Apr 24, 2008 May CAHN

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

METADATE CD

>D> BX	UCB INC	10MG	N021259 003 May 27, 2003 Jul CFTG
>A> AB		10MG	N021259 003 May 27, 2003 Jul CFTG
>D> BX		20MG	N021259 001 Apr 03, 2001 Jul CFTG
>A> AB2		20MG	N021259 001 Apr 03, 2001 Jul CFTG
>D> BX		30MG	N021259 002 Jun 19, 2003 Jul CFTG
>A> AB2		30MG	N021259 002 Jun 19, 2003 Jul CFTG
>D> BX		40MG	N021259 004 Feb 19, 2006 Jul CFTG
>A> AB2		40MG	N021259 004 Feb 19, 2006 Jul CFTG
>D>		50MG	N021259 005 Feb 19, 2006 Jul CFTG
>A> AB		50MG	N021259 005 Feb 19, 2006 Jul CFTG
>D> +		60MG	N021259 006 Feb 19, 2006 Jul CFTG
>A> AB +		60MG	N021259 006 Feb 19, 2006 Jul CFTG

CAPSULE, EXTENDED RELEASE; ORAL
METHYLPHENIDATE HYDROCHLORIDE

>D>	AB	ACTAVIS	20MG	A078458 001	Dec 01, 2011	Jul	CTEC
>A>	AB1		20MG	A078458 001	Dec 01, 2011	Jul	CTEC
>D>	AB		30MG	A078458 002	Dec 01, 2011	Jul	CTEC
>A>	AB1		30MG	A078458 002	Dec 01, 2011	Jul	CTEC
>D>	AB		40MG	A078458 003	Dec 01, 2011	Jul	CTEC
>A>	AB1		40MG	A078458 003	Dec 01, 2011	Jul	CTEC
>D>	AB	BARR LABS INC	20MG	A079031 001	Jul 13, 2012	Jul	CTEC
>A>	AB1		20MG	A079031 001	Jul 13, 2012	Jul	CTEC
		AB	20MG	A079031 001	Jul 13, 2012	Jun	NEWA
>D>	AB		30MG	A079031 002	Jul 13, 2012	Jul	CTEC
>A>	AB1		30MG	A079031 002	Jul 13, 2012	Jul	CTEC
		AB	30MG	A079031 002	Jul 13, 2012	Jun	NEWA
>D>	AB		40MG	A079031 003	Jul 13, 2012	Jul	CTEC
>A>	AB1		40MG	A079031 003	Jul 13, 2012	Jul	CTEC
		AB	40MG	A079031 003	Jul 13, 2012	Jun	NEWA
>A>	AB	TEVA PHARMS	10MG	A077707 001	Jul 19, 2012	Jul	NEWA
>A>	AB2		20MG	A077707 002	Jul 19, 2012	Jul	NEWA
>A>	AB2		30MG	A077707 003	Jul 19, 2012	Jul	NEWA
>A>	AB2		40MG	A078873 001	Jul 19, 2012	Jul	NEWA
>A>	AB		50MG	A078873 002	Jul 19, 2012	Jul	NEWA
>A>	AB		60MG	A078873 003	Jul 19, 2012	Jul	NEWA

RITALIN LA

>D>	AB	NOVARTIS	20MG	N021284 001	Jun 05, 2002	Jul	CTEC
>A>	AB1		20MG	N021284 001	Jun 05, 2002	Jul	CTEC
>D>	AB		30MG	N021284 002	Jun 05, 2002	Jul	CTEC
>A>	AB1		30MG	N021284 002	Jun 05, 2002	Jul	CTEC
>D>	AB	+	40MG	N021284 003	Jun 05, 2002	Jul	CTEC
>A>	AB1	+	40MG	N021284 003	Jun 05, 2002	Jul	CTEC

TABLET; ORAL

METHYLPHENIDATE HYDROCHLORIDE

AB	SUN PHARM INDS INC	5MG	A090710 001	Mar 15, 2012	Feb	NEWA
AB		10MG	A090710 002	Mar 15, 2012	Feb	NEWA
AB		20MG	A090710 003	Mar 15, 2012	Feb	NEWA

TABLET, EXTENDED RELEASE; ORAL

>D>		METADATE ER					
>D>	AB	UCB INC	10MG	A040306 001	Oct 20, 1999	Jul	DISC
>A>		@	10MG	A040306 001	Oct 20, 1999	Jul	DISC

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE SODIUM SUCCINATE

@	BEDFORD LABS	EQ 40MG BASE/VIAL	A040662 001	Feb 21, 2007	Jun	DISC
@		EQ 125MG BASE/VIAL	A040641 002	Feb 21, 2007	Jun	DISC
@		EQ 500MG BASE/VIAL	A040641 003	Feb 21, 2007	Jun	DISC
@		EQ 500MG BASE/VIAL	A040709 001	Feb 21, 2007	Jun	DISC
@		EQ 1GM BASE/VIAL	A040641 004	Feb 21, 2007	Jun	DISC
@		EQ 1GM BASE/VIAL	A040709 002	Feb 21, 2007	Jun	DISC
@	TEVA PARENTERAL	EQ 125MG BASE/VIAL	A081266 001	Nov 30, 1992	Mar	DISC

METOCLOPRAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

METOCLOPRAMIDE HYDROCHLORIDE

@	TEVA PARENTERAL	EQ 5MG BASE/ML	A073135 001	Nov 27, 1991	Jan	DISC
---	-----------------	----------------	-------------	--------------	-----	------

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL						
MINOCIN						
AB PRECISION DERMAT	EQ 50MG BASE	N050649 001	May 31, 1990	May	CAHN	
@	EQ 75MG BASE	N050649 003	Feb 12, 2001	May	CAHN	
AB	EQ 100MG BASE	N050649 002	May 31, 1990	May	CAHN	
>A> CAPSULE, EXTENDED RELEASE; ORAL						
>A> XIMINO						
>A> RANBAXY LABS LTD	EQ 45MG BASE	N201922 001	Jul 11, 2012	Jul	NEWA	
>A>	EQ 67.5MG BASE	N201922 002	Jul 11, 2012	Jul	NEWA	
>A>	EQ 90MG BASE	N201922 003	Jul 11, 2012	Jul	NEWA	
>A>	EQ 112.5MG BASE	N201922 004	Jul 11, 2012	Jul	NEWA	
>A> +	EQ 135MG BASE	N201922 005	Jul 11, 2012	Jul	NEWA	
INJECTABLE; INJECTION						
MINOCIN						
+ PRECISION DERMAT	EQ 100MG BASE/VIAL	N050444 001		May	CAHN	
SUSPENSION; ORAL						
MINOCIN						
@ PRECISION DERMAT	EQ 50MG BASE/5ML	N050445 001		May	CAHN	

MIRABEGRON

TABLET, EXTENDED RELEASE; ORAL						
MYRBETRIQ						
APGDI	25MG	N202611 001	Jun 28, 2012	Jun	NEWA	
+	50MG	N202611 002	Jun 28, 2012	Jun	NEWA	

MISOPROSTOL

TABLET; ORAL						
MISOPROSTOL						
>A> AB NOVEL LABS INC	0.1MG	A091667 001	Jul 25, 2012	Jul	NEWA	
>A> AB	0.2MG	A091667 002	Jul 25, 2012	Jul	NEWA	

MITOMYCIN

FOR SOLUTION; TOPICAL						
MITOSOL						
+ MOBIUS THERAP	0.2MG/VIAL	N022572 001	Feb 07, 2012	Feb	NEWA	

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION						
MITOXANTRONE HYDROCHLORIDE						
@ FRESENIUS KABI ONCOL	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	A078606 001	May 14, 2008	Jun	DISC	
@	EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)	A078606 002	May 14, 2008	Jun	DISC	
@	EQ 30MG BASE/15ML (EQ 2MG BASE/ML)	A078606 003	May 14, 2008	Jun	DISC	

MIVACURIUM CHLORIDE

INJECTABLE; INJECTION						
MIVACURIUM CHLORIDE						
@ STRIDES ARCOLAB LTD	EQ 2MG BASE/ML	A078562 001	Apr 30, 2009	Jun	CAHN	

MODAFINIL

TABLET; ORAL						
MODAFINIL						
AB MYLAN PHARMS INC	100MG	A076594 001	Jul 16, 2012	Jun	NEWA	

TABLET; ORAL

MODAFINIL

AB	MYLAN PHARMS INC	200MG	A076594 002 Jul 16, 2012 Jun NEWA
	PROVIGIL		
AB	CEPHALON	100MG	N020717 001 Dec 24, 1998 Jun CFTG
AB	+	200MG	N020717 002 Dec 24, 1998 Jun CFTG

MOMETASONE FUROATE

CREAM; TOPICAL

ELOCON

AB	+	MERCK SHARP DOHME	0.1%	N019625 001 May 06, 1987 Jun CAHN	
		MOMETASONE FUROATE			
AB		FOUGERA PHARMS	0.1%	A076171 001 Apr 08, 2005 Jan CAHN	
		LOTION; TOPICAL			
		ELOCON			
>A>	AB	+	MERCK SHARP DOHME	0.1%	N019796 001 Mar 30, 1989 Jul CAHN
>D>	AB	+	SCHERING	0.1%	N019796 001 Mar 30, 1989 Jul CAHN
		MOMETASONE FUROATE			
AB		FOUGERA PHARMS	0.1%	A075919 001 Nov 29, 2007 Jan CAHN	
		OINTMENT; TOPICAL			
		ELOCON			
AB	+	MERCK SHARP DOHME	0.1%	N019543 001 Apr 30, 1987 May CAHN	
		MOMETASONE FUROATE			
AB		FOUGERA PHARMS	0.1%	A077061 001 Mar 28, 2005 Jan CAHN	

MONTELUKAST SODIUM

GRANULE; ORAL

MONTELUKAST SODIUM

>A>		TEVA PHARMS	EQ 4MG BASE/PACKET	A090955 001 Aug 03, 2012 Jul NEWA
		SINGULAIR		
>D>	+	MERCK	EQ 4MG BASE/PACKET	N021409 001 Jul 26, 2002 Jul CFTG
>A>	AB	+	EQ 4MG BASE/PACKET	N021409 001 Jul 26, 2002 Jul CFTG

TABLET; ORAL

MONTELUKAST SODIUM

>A>	AB	APOTEX CORP	EQ 10MG BASE	A201294 001 Aug 03, 2012 Jul NEWA
>A>	AB	AUROBINDO PHARMA LTD	EQ 10MG BASE	A202468 001 Aug 03, 2012 Jul NEWA
>A>	AB	DR REDDYS LABS LTD	EQ 10MG BASE	A201582 001 Aug 06, 2012 Jul NEWA
>A>	AB	ENDO PHARMS	EQ 10MG BASE	A091576 001 Aug 03, 2012 Jul NEWA
>A>	AB	GLENMARK GENERICS	EQ 10MG BASE	A090926 001 Aug 03, 2012 Jul NEWA
>A>	AB	KUDCO IRELAND	EQ 10MG BASE	A201522 001 Aug 03, 2012 Jul NEWA
>A>	AB	MYLAN PHARMS INC	EQ 10MG BASE	A079103 001 Aug 03, 2012 Jul NEWA
>A>	AB	ROXANE	EQ 10MG BASE	A090655 001 Aug 03, 2012 Jul NEWA
>A>	AB	SANDOZ INC	EQ 10MG BASE	A200889 001 Aug 03, 2012 Jul NEWA
>A>	AB	TEVA PHARMS	EQ 10MG BASE	A078605 001 Aug 03, 2012 Jul NEWA
>A>	AB	TORRENT PHARMS LTD	EQ 10MG BASE	A201515 001 Aug 03, 2012 Jul NEWA

SINGULAIR

>D>	+	MERCK	EQ 10MG BASE	N020829 002 Feb 20, 1998 Jul CFTG
>A>	AB	+	EQ 10MG BASE	N020829 002 Feb 20, 1998 Jul CFTG

TABLET, CHEWABLE; ORAL

MONTELUKAST SODIUM

>A>	AB	APOTEX INC	EQ 4MG BASE	A201508 001 Aug 03, 2012 Jul NEWA
>A>	AB		EQ 5MG BASE	A201508 002 Aug 03, 2012 Jul NEWA
>A>	AB	AUROBINDO PHARMA LTD	EQ 4MG BASE	A202096 001 Aug 03, 2012 Jul NEWA
>A>	AB		EQ 5MG BASE	A202096 002 Aug 03, 2012 Jul NEWA
>A>	AB	DR REDDYS LABS LTD	EQ 4MG BASE	A201581 001 Aug 06, 2012 Jul NEWA
>A>	AB		EQ 5MG BASE	A201581 002 Aug 06, 2012 Jul NEWA

TABLET, CHEWABLE; ORAL

>A>	MONTELUKAST SODIUM											
>A> AB	ENDO PHARMS	EQ 4MG BASE	A091588	001	Aug 03,	2012	Jul	NEWA				
>A> AB		EQ 5MG BASE	A091588	002	Aug 03,	2012	Jul	NEWA				
>A> AB	KUDCO IRELAND	EQ 4MG BASE	A200405	001	Aug 03,	2012	Jul	NEWA				
>A> AB		EQ 5MG BASE	A200405	002	Aug 03,	2012	Jul	NEWA				
>A> AB	MYLAN PHARMS INC	EQ 4MG BASE	A079142	001	Aug 03,	2012	Jul	NEWA				
>A> AB		EQ 5MG BASE	A079142	002	Aug 03,	2012	Jul	NEWA				
>A> AB	ROXANE	EQ 4MG BASE	A091128	001	Aug 03,	2012	Jul	NEWA				
>A> AB		EQ 5MG BASE	A091128	002	Aug 03,	2012	Jul	NEWA				
>A> AB	SANDOZ INC	EQ 4MG BASE	A091414	001	Aug 03,	2012	Jul	NEWA				
>A> AB		EQ 5MG BASE	A091414	002	Aug 03,	2012	Jul	NEWA				
>A> AB	TEVA PHARMS	EQ 4MG BASE	A078723	001	Aug 03,	2012	Jul	NEWA				
>A> AB		EQ 5MG BASE	A078723	002	Aug 03,	2012	Jul	NEWA				
>A> AB	TORRENT PHARMS LTD	EQ 4MG BASE	A090984	001	Aug 03,	2012	Jul	NEWA				
>A> AB		EQ 5MG BASE	A090984	002	Aug 03,	2012	Jul	NEWA				
<u>SINGULAIR</u>												
>D>	MERCK	EQ 4MG BASE	N020830	002	Mar 03,	2000	Jul	CFTG				
>A> AB		EQ 4MG BASE	N020830	002	Mar 03,	2000	Jul	CFTG				
>D>	+	EQ 5MG BASE	N020830	001	Feb 20,	1998	Jul	CFTG				
>A> AB	+	EQ 5MG BASE	N020830	001	Feb 20,	1998	Jul	CFTG				

MORPHINE SULFATECAPSULE, EXTENDED RELEASE; ORALKADIAN

>A>	ACTAVIS ELIZABETH	40MG	N020616	009	Jul 09,	2012	Jul	NEWA				
>A>		70MG	N020616	010	Jul 09,	2012	Jul	NEWA				
>A>		130MG	N020616	011	Jul 09,	2012	Jul	NEWA				
>A>		150MG	N020616	012	Jul 09,	2012	Jul	NEWA				

INJECTABLE; INJECTIONDURAMORPH PF

AP	+	HIKMA MAPLE	0.5MG/ML	N018565	001	Sep 18,	1984	Jun	CAHN			
AP	+		1MG/ML	N018565	002	Sep 18,	1984	Jun	CAHN			
<u>INFUMORPH</u>												
	+	HIKMA MAPLE	10MG/ML	N018565	003	Jul 19,	1991	Jun	CAHN			
	+		25MG/ML	N018565	004	Jul 19,	1991	Jun	CAHN			

INJECTABLE, LIPOSOMAL; EPIDURALDEPODUR

@	PACIRA PHARMS INC	10MG/ML (10MG/ML)	N021671	001	May 18,	2004	May	CAHN				
@		15MG/1.5ML (10MG/ML)	N021671	002	May 18,	2004	May	CAHN				
@		20MG/2ML (10MG/ML)	N021671	003	May 18,	2004	May	CAHN				

SOLUTION; ORALMORPHINE SULFATE

>A> AA	PADDOCK LLC	100MG/5ML	A201574	001	Aug 06,	2012	Jul	NEWA				
--------	-------------	-----------	---------	-----	---------	------	-----	------	--	--	--	--

TABLET, EXTENDED RELEASE; ORALMORPHINE SULFATE

AB	RANBAXY LABS LTD	15MG	A078761	001	May 11,	2012	Apr	NEWA				
AB		30MG	A078761	002	May 11,	2012	Apr	NEWA				
AB		60MG	A078761	003	May 11,	2012	Apr	NEWA				
AB		100MG	A078761	004	May 11,	2012	Apr	NEWA				
AB		200MG	A078761	005	May 11,	2012	Apr	NEWA				

ORAMORPH SR

@	XANODYNE PHARMS INC	15MG	N019977	004	Nov 23,	1994	May	DISC				
@		30MG	N019977	001	Aug 15,	1991	May	DISC				
@		60MG	N019977	002	Aug 15,	1991	May	DISC				
@		100MG	N019977	003	Aug 15,	1991	May	DISC				

MUPIROCIN

OINTMENT; TOPICAL

MUPIROCIN

AB FOUGERA PHARMS 2% A065192 001 Nov 30, 2005 Jan CAHN

NABUMETONE

TABLET; ORAL

NABUMETONE

AB PROSAM LABS 500MG A079093 001 Feb 27, 2009 Feb CMFD
AB 750MG A079093 002 Feb 27, 2009 Feb CMFDNAFTIFINE HYDROCHLORIDE

CREAM; TOPICAL

NAFTIN

+ MERZ PHARMS 2% N019599 002 Jan 13, 2012 Jan NEWA

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

@ HIKMA MAPLE 0.4MG/ML A070298 001 Sep 24, 1986 Jun CAHN
@ 0.4MG/ML A070299 001 Sep 24, 1986 Jun CAHN
@ 0.4MG/ML A070496 001 Sep 24, 1986 Jun CAHN

NALOXONE HYDROCHLORIDE

@ HIKMA MAPLE 0.02MG/ML A071272 001 May 24, 1988 Jun CAHN
@ 1MG/ML A071273 001 May 24, 1988 Jun CAHN
@ 1MG/ML A071274 001 May 24, 1988 Jun CAHN
@ 1MG/ML A071287 001 May 24, 1988 Jun CAHNNALTREXONE HYDROCHLORIDE

TABLET; ORAL

NALTREXONE HYDROCHLORIDE

AB SUN PHARMA GLOBAL 50MG A090356 001 Feb 24, 2012 Feb NEWA

NAPROXEN SODIUM

TABLET; ORAL

ANAPROX

AB HOFFMANN LA ROCHE EQ 250MG BASE N018164 001 May CAHN

ANAPROX DS

AB + HOFFMANN LA ROCHE EQ 500MG BASE N018164 003 Sep 30, 1987 May CAHN

NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE

TABLET; ORAL

TREXIMET

+ SMITHKLINE BEECHAM 500MG;EQ 85MG BASE N021926 001 Apr 15, 2008 May CAHN

NARATRIPTAN HYDROCHLORIDE

TABLET; ORAL

NARATRIPTAN

AB MYLAN PHARMS INC EQ 1MG BASE A202431 001 May 31, 2012 May NEWA

AB EQ 2.5MG BASE A202431 002 May 31, 2012 May NEWA

AB ORCHID HLTHCARE EQ 1MG BASE A091441 001 Apr 30, 2012 Apr NEWA

AB EQ 2.5MG BASE A091441 002 Apr 30, 2012 Apr NEWA

NATEGLINIDE

TABLET; ORAL

NATEGLINIDE

>D>	AB	TEVA PHARMS	60MG	A077467	001	Sep 09, 2009	Jul	DISC
>A>		@	60MG	A077467	001	Sep 09, 2009	Jul	DISC
>D>	AB		120MG	A077467	002	Sep 09, 2009	Jul	DISC
>A>		@	120MG	A077467	002	Sep 09, 2009	Jul	DISC

NEVIRAPINE

SUSPENSION; ORAL

NEVIRAPINE

AA AUROBINDO 50MG/5ML A077702 001 May 22, 2012 May NEWA
 AA + BOEHRINGER INGELHEIM 50MG/5ML N020933 001 Sep 11, 1998 May CFTG

TABLET; ORAL

NEVIRAPINE

AB	APOTEX INC	200MG	A203021	001	May 22, 2012	May	NEWA
AB	AUROBINDO	200MG	A077521	001	May 22, 2012	May	NEWA
AB	CIPLA	200MG	A077956	001	May 22, 2012	May	NEWA
AB	HETERO LABS LTD III	200MG	A078584	001	May 22, 2012	May	NEWA
AB	MICRO LABS LTD	200MG	A203080	001	May 22, 2012	May	NEWA
AB	MYLAN LABS	200MG	A078864	001	May 22, 2012	May	NEWA
AB	MYLAN PHARMS INC	200MG	A202523	001	May 22, 2012	May	NEWA
AB	PRINSTON INC	200MG	A078644	001	May 22, 2012	May	NEWA
AB	SCIEGEN PHARMS INC	200MG	A203176	001	May 22, 2012	May	NEWA
AB	STRIDES	200MG	A078195	001	May 22, 2012	May	NEWA
VIRAMUNE							
AB	+ BOEHRINGER INGELHEIM	200MG	N020636	001	Jun 21, 1996	May	CFTG

NIFEDIPINE

TABLET, EXTENDED RELEASE; ORAL

NIFEDIPINE

AB1 ACTAVIS 90MG A077899 003 May 25, 2012 May NEWA

NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACRODANTIN

>A>	AB	ALMATICCA	25MG	N016620 003	Jul	CAHN
>A>	AB		50MG	N016620 001	Jul	CAHN
>A>	AB	+	100MG	N016620 002	Jul	CAHN
>D>	AB	ALVOGEN	25MG	N016620 003	Jul	CAHN
>D>	AB		50MG	N016620 001	Jul	CAHN
>D>	AB	+	100MG	N016620 002	Jul	CAHN

NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACROBID

>A>	AB	+	ALMATICA	75MG; 25MG	N020064	001	Dec 24, 1991	Jul	CAHN
>D>	AB	+	ALVOGEN	75MG; 25MG	N020064	001	Dec 24, 1991	Jul	CAHN
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)									
@ RANBAXY LABS LTD 75MG; 25MG					A076951	001	Mar 30, 2005	Apr	DISC

NITROGLYCERIN

FILM, EXTENDED RELEASE; TRANSDERMAL

MINITRAN

>D>	AB1	GRACEWAY	0.1MG/HR	A089771	001	Aug 30,	1996	Jul	CAHN
>D>	AB1		0.2MG/HR	A089772	001	Aug 30,	1996	Jul	CAHN
>D>	AB1		0.4MG/HR	A089773	001	Aug 30,	1996	Jul	CAHN
>D>	AB1		0.6MG/HR	A089774	001	Aug 30,	1996	Jul	CAHN
>A>	AB1	MEDICIS	0.1MG/HR	A089771	001	Aug 30,	1996	Jul	CAHN
>A>	AB1		0.2MG/HR	A089772	001	Aug 30,	1996	Jul	CAHN
>A>	AB1		0.4MG/HR	A089773	001	Aug 30,	1996	Jul	CAHN
>A>	AB1		0.6MG/HR	A089774	001	Aug 30,	1996	Jul	CAHN
OINTMENT; INTRA-ANAL									
RECTIV									
+ APTALIS PHARMA			0.4%						
				N021359	001	Jun 21,	2011	Jan	CAHN

NIZATIDINE

CAPSULE; ORAL

AXID

@ SMITHKLINE BEECHAM	150MG	N019508	001	Apr 12,	1988	Mar	DISC
@	300MG	N019508	002	Apr 12,	1988	Mar	DISC

NIZATIDINE

@ MYLAN PHARMS INC	150MG	A075934	001	Jul 09,	2002	Apr	DISC
AB +	300MG	A075806	002	Jul 05,	2002	Apr	CRLD

@	300MG	A075934	002	Jul 09,	2002	Apr	DISC
---	-------	---------	-----	---------	------	-----	------

NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

NOREPINEPHRINE BITARTRATE

AP CLARIS LIFESCIENCES	EQ 1MG BASE/ML	A040859	001	Mar 27,	2012	May	CAHN
AP SINTETICA	EQ 1MG BASE/ML	A040859	001	Mar 27,	2012	Mar	NEWA

NORETHINDRONE ACETATE

TABLET; ORAL

NORETHIDRONE ACETATE

>A> AB AMNEAL PHARMS	5MG	A200275	001	Jul 30,	2012	Jul	NEWA
----------------------	-----	---------	-----	---------	------	-----	------

NYSTATIN

CREAM; TOPICAL

NYSTATIN

AT FOUGERA PHARMS	100,000 UNITS/GM	A062129	001				Jan	CAHN
OINTMENT; TOPICAL								

NYSTATIN

AT + FOUGERA PHARMS	100,000 UNITS/GM	A062124	002	Sep 23,	1982	Jan	CAHN
---------------------	------------------	---------	-----	---------	------	-----	------

POWDER; TOPICAL

NYSTATIN

@ NESHER PHARMS	100,000 UNITS/GM	A065321	001	Aug 18,	2006	Jun	DISC
-----------------	------------------	---------	-----	---------	------	-----	------

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYKACET

@ G AND W LABS INC	100,000 UNITS/GM;0.1%	A062367	001	May 28,	1985	Jun	CAHN
--------------------	-----------------------	---------	-----	---------	------	-----	------

OINTMENT; TOPICAL

MYKACET

@ G AND W LABS INC	100,000 UNITS/GM;0.1%	A062733	001	Mar 09,	1987	Jun	CAHN
--------------------	-----------------------	---------	-----	---------	------	-----	------

OFLOXACIN

TABLET; ORAL

OFLOXACIN

@ RANBAXY LABS LTD	200MG
@	300MG
@	400MG

A076220 001	Sep 02, 2003	Apr	DISC
A076220 002	Sep 02, 2003	Apr	DISC
A076220 003	Sep 02, 2003	Apr	DISC

OLANZAPINE

INJECTABLE; INTRAMUSCULAR

OLANZAPINE

AP PHARMAFORCE	10MG/VIAL
AP SANDOZ INC	10MG/VIAL

A201741 001	Mar 20, 2012	Mar	NEWA
A201588 001	Oct 24, 2011	Mar	CAHN

TABLET; ORAL

OLANZAPINE

AB APOTEX INC	2.5MG
AB	5MG
AB	7.5MG
AB	10MG
AB	15MG
AB	20MG
AB AUROBINDO PHARMA LTD	2.5MG
AB	5MG
AB	7.5MG
AB	10MG
AB	15MG
AB	20MG
AB DR REDDYS LABS LTD	2.5MG
AB	5MG
AB	7.5MG
AB	10MG
AB	15MG
AB MYLAN	2.5MG
AB	5MG
AB	7.5MG
AB	10MG
AB	15MG
AB	20MG
AB ORCHID HLTHCARE	2.5MG
AB	5MG
AB	7.5MG
AB	10MG
AB	15MG
AB	20MG
AB SUN PHARM INDs	2.5MG
AB	5MG
AB	7.5MG
AB	10MG
AB	15MG
AB	20MG
AB TORRENT PHARMS LTD	2.5MG
AB	5MG
AB	7.5MG
AB	10MG
AB	15MG
AB	20MG

A090798 001	Apr 23, 2012	Apr	NEWA
A090798 002	Apr 23, 2012	Apr	NEWA
A090798 003	Apr 23, 2012	Apr	NEWA
A090798 004	Apr 23, 2012	Apr	NEWA
A090798 005	Apr 23, 2012	Apr	NEWA
A090798 006	Apr 23, 2012	Apr	NEWA
A202050 001	Apr 23, 2012	Apr	NEWA
A202050 002	Apr 23, 2012	Apr	NEWA
A202050 003	Apr 23, 2012	Apr	NEWA
A202050 004	Apr 23, 2012	Apr	NEWA
A202050 005	Apr 23, 2012	Apr	NEWA
A202050 006	Apr 23, 2012	Apr	NEWA
A076255 001	Apr 23, 2012	Apr	NEWA
A076255 002	Apr 23, 2012	Apr	NEWA
A076255 003	Apr 23, 2012	Apr	NEWA
A076255 004	Apr 23, 2012	Apr	NEWA
A076133 001	Apr 23, 2012	Apr	NEWA
A076866 001	Apr 23, 2012	Apr	NEWA
A076866 002	Apr 23, 2012	Apr	NEWA
A076866 003	Apr 23, 2012	Apr	NEWA
A076866 004	Apr 23, 2012	Apr	NEWA
A076866 005	Apr 23, 2012	Apr	NEWA
A076866 006	Apr 23, 2012	Apr	NEWA
A202287 001	Apr 23, 2012	Apr	NEWA
A202287 002	Apr 23, 2012	Apr	NEWA
A202287 003	Apr 23, 2012	Apr	NEWA
A202287 004	Apr 23, 2012	Apr	NEWA
A202287 005	Apr 23, 2012	Apr	NEWA
A202287 006	Apr 23, 2012	Apr	NEWA
A091038 001	Apr 23, 2012	Apr	NEWA
A091038 002	Apr 23, 2012	Apr	NEWA
A091038 003	Apr 23, 2012	Apr	NEWA
A091038 004	Apr 23, 2012	Apr	NEWA
A091038 005	Apr 23, 2012	Apr	NEWA
A091038 006	Apr 23, 2012	Apr	NEWA
A091434 001	Apr 23, 2012	Apr	NEWA
A091434 002	Apr 23, 2012	Apr	NEWA
A091434 003	Apr 23, 2012	Apr	NEWA
A091434 004	Apr 23, 2012	Apr	NEWA
A091434 005	Apr 23, 2012	Apr	NEWA
A091434 006	Apr 23, 2012	Apr	NEWA

TABLET, ORALLY DISINTEGRATING; ORAL

OLANZAPINE

AB	BARR LABS INC	5MG	A077243 001	Jan 30, 2012	Jan	NEWA
AB		10MG	A077243 002	Jan 30, 2012	Jan	NEWA
AB		15MG	A077243 003	Jan 30, 2012	Jan	NEWA
AB		20MG	A077243 004	Jan 30, 2012	Jan	NEWA
AB	SUN PHARM IND	5MG	A090881 001	Feb 28, 2012	Feb	NEWA
AB		10MG	A090881 002	Feb 28, 2012	Feb	NEWA
AB		15MG	A090881 003	Feb 28, 2012	Feb	NEWA
AB		20MG	A090881 004	Feb 28, 2012	Feb	NEWA

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL

OMEPRAZOLE

AB	DR REDDYS LABS LTD	10MG	A078490 002	Mar 16, 2009	Apr	NEWA
AB		20MG	A078490 003	Mar 16, 2009	Apr	NEWA

ONDANSETRON

TABLET, ORALLY DISINTEGRATING; ORAL

ONDANSETRON

@	NESHER PHARMS	4MG	A077717 001	Jun 25, 2007	Jun	DISC
@		8MG	A077717 002	Jun 25, 2007	Jun	DISC

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ONDANSETRON HYDROCHLORIDE

AP	GLAND PHARMA LTD	EQ 2MG BASE/ML	A090648 001	Jun 15, 2012	Jun	NEWA
AP	HIKMA MAPLE	EQ 2MG BASE/ML	A077365 001	Dec 26, 2006	Jun	CAHN
	ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE					
AP	HIKMA MAPLE	EQ 2MG BASE/ML	A077541 001	Dec 26, 2006	Jun	CAHN

ORPHENADRINE CITRATE

TABLET, EXTENDED RELEASE; ORAL

ORPHENADRINE CITRATE

>A>	AB	CNTY LINE PHARMS	100MG	A040249 001	Jan 29, 1999	Jul	CAHN
>A>	AB	INVAGEN PHARMS	100MG	A091158 001	Jul 27, 2012	Jul	NEWA
>D>	AB	KIEL	100MG	A040249 001	Jan 29, 1999	Jul	CAHN

OSELTAMIVIR PHOSPHATE

FOR SUSPENSION; ORAL

TAMIFLU

+	ROCHE	EQ 6MG BASE/ML	N021246 002	Mar 21, 2011	Jan	CRLD
---	-------	----------------	-------------	--------------	-----	------

OXACILLIN SODIUM

INJECTABLE; INJECTION

OXACILLIN SODIUM

AP	SAGENT PHARMS	EQ 1GM BASE/VIAL	A091246 001	Mar 30, 2012	Mar	NEWA
AP		EQ 2GM BASE/VIAL	A091246 002	Mar 30, 2012	Mar	NEWA
AP		EQ 10GM BASE/VIAL	A091245 001	Mar 30, 2012	Mar	NEWA

OXALIPLATIN

INJECTABLE; IV (INFUSION)

OXALIPLATIN

>A>	AP	ACTAVIS TOTOWA	50MG/VIAL	A078803 001	Aug 08, 2012	Jul	NEWA
>A>	AP		100MG/VIAL	A078803 002	Aug 08, 2012	Jul	NEWA
>A>	AP	ONCO THERAPIES LTD	50MG/VIAL	A200979 001	Aug 08, 2012	Jul	NEWA

INJECTABLE; IV (INFUSION)OXALIPLATIN

>A>	AP	ONCO THERAPIES LTD	50MG/10ML (5MG/ML)	A091358 001 Aug 07, 2012 Jul NEWA
>A>	AP		100MG/20ML (5MG/ML)	A091358 002 Aug 07, 2012 Jul NEWA
>A>	AP		100MG/VIAL	A200979 002 Aug 08, 2012 Jul NEWA
	AP +	TEVA PHARMS	50MG/10ML (5MG/ML)	N022160 001 Aug 07, 2009 Mar CRLD
	AP +		100MG/20ML (5MG/ML)	N022160 002 Aug 07, 2009 Mar CRLD

OXICONAZOLE NITRATECREAM; TOPICALOXISTAT

+ FOUGERA PHARMS	EQ 1% BASE
------------------	------------

N019828 001 Dec 30, 1988 Jan CAHN

OXTRIPHYLLINETABLET, EXTENDED RELEASE; ORALCHOLEDYL SA

+ WARNER CHILCOTT LLC	400MG	A087863 001 May 24, 1983 Jun CAHN
+	600MG	A086742 001 Jun CAHN

OXYBUTYNINGEL, METERED; TRANSDERMALANTUROL

+ ARROW INTL	3%	N202513 001 Dec 07, 2011 Apr CAHN
+ WATSON LABS INC	3%	N202513 001 Dec 07, 2011 Feb CAHN

OXYBUTYNIN CHLORIDETABLET; ORALDITROPAN

@ JANSSEN PHARMS	5MG	N017577 001 Mar DISC
OXYBUTYNIN CHLORIDE		

AB + VINTAGE PHARMS	5MG	A075079 001 Oct 31, 1997 Mar CRLD
TABLET, EXTENDED RELEASE; ORAL		
OXYBUTYNIN CHLORIDE		
AB MYLAN PHARMS INC	15MG	A076644 002 May 10, 2007 Mar NEWA

OXYCODONE HYDROCHLORIDECAPSULE; ORAL

>A> OXYCODONE HYDROCHLORIDE		
>A> AB COASTAL PHARMS	5MG	A203107 001 Jul 26, 2012 Jul NEWA
>D> + LEHIGH VALLEY	5MG	N200534 001 Oct 20, 2010 Jul CFTG
>A> AB +	5MG	N200534 001 Oct 20, 2010 Jul CFTG

SOLUTION; ORALOXYCODONE HYDROCHLORIDE

>D> + LEHIGH VALLEY	100MG/5ML	N200535 001 Oct 20, 2010 Jul CFTG
>A> AA +	100MG/5ML	N200535 001 Oct 20, 2010 Jul CFTG
	5MG/5ML	N201194 001 Jan 12, 2012 Jan NEWA
>A> AA	100MG/5ML	A202537 001 Jul 30, 2012 Jul NEWA

TABLET; ORALOXYCODONE HYDROCHLORIDE

AB MALLINCKRODT INC	5MG	A076758 003 Mar 19, 2007 Apr NEWA
---------------------	-----	-----------------------------------

OXYMORPHONE HYDROCHLORIDETABLET, EXTENDED RELEASE; ORALOPANA ER

@ ENDO PHARMS	5MG	N021610 001 Jun 22, 2006 May DISC
@	10MG	N021610 002 Jun 22, 2006 May DISC

TABLET, EXTENDED RELEASE; ORAL

OPANA ER

@ ENDO PHARMS	20MG	N021610 003	Jun 22, 2006	May	DISC
@	30MG	N021610 007	Feb 29, 2008	May	DISC
@	40MG	N021610 004	Jun 22, 2006	May	DISC

OXYMORPHONE HYDROCHLORIDE

IMPAX LABS	5MG	A079087 001	Jun 14, 2010	May	CTEC
	10MG	A079087 003	Jun 14, 2010	May	CTEC
	20MG	A079087 005	Jun 14, 2010	May	CTEC
	30MG	A079087 006	Jul 22, 2010	May	CTEC
	40MG	A079087 007	Jun 14, 2010	May	CTEC

OXYTOCIN

INJECTABLE; INJECTION

OXYTOCIN

@ TEVA PARENTERAL	10USP UNITS/ML (10USP UNITS/ML)	A077453 001	Jan 24, 2008	Jan	DISC
@	100USP UNITS/10ML (10USP UNITS/ML)	A077453 002	Jan 24, 2008	Mar	DISC

PACLITAXEL

INJECTABLE; INJECTION

TAXOL

@ CORDEN PHARMA	6MG/ML	N020262 001	Dec 29, 1992	Mar	CAHN
-----------------	--------	-------------	--------------	-----	------

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

AREDIA

@ NOVARTIS	30MG/VIAL	N020036 001	Oct 31, 1991	Mar	DISC
------------	-----------	-------------	--------------	-----	------

PANCRELIPASE (AMYLASE;LIPASE;PROTEASE)

CAPSULE, DELAYED RELEASE; ORAL

CREON

ABBOTT LABS	15,000USP UNITS;3,000USP UNITS;9,500USP UNITS	N020725 004	Jul 12, 2011	Jun	NEWA
	30,000USP UNITS;6,000USP UNITS;19,000USP UNITS	N020725 001	Apr 30, 2009	Feb	CAHN
	60,000USP UNITS;12,000USP UNITS;38,000USP UNITS	N020725 002	Apr 30, 2009	Feb	CAHN
+	120,000USP UNITS;24,000USP UNITS;76,000USP UNITS	N020725 003	Apr 30, 2009	Feb	CAHN

PERTZYE

DIGESTIVE CARE INC	30,250USP/UNITS;8,000USP/UNITS;28,750USP/UNITS	N022175 001	May 17, 2012	May	NEWA
+	60,500USP/UNITS;16,000USP/UNITS;5,7,500USP/UNITS	N022175 002	May 17, 2012	May	NEWA

ULTRESA

APTALIS PHARMA US	27,600USP UNITS;13,800USP UNITS;27,600USP UNITS	N022222 001	Mar 01, 2012	Mar	NEWA
	41,400USP UNITS;20,700USP UNITS;41,400USP UNITS	N022222 002	Mar 01, 2012	Mar	NEWA
+	46,000USP UNITS;23,000USP UNITS;46,000USP UNITS	N022222 003	Mar 01, 2012	Mar	NEWA

ZENPEP

APTALIS PHARMA US	16,000USP UNITS;3,000USP UNITS;10,000USP UNITS	N022210 005	Jun 15, 2011	Apr	NEWA
	109,000USP UNITS;20,000USP UNITS;68,000USP UNITS	N022210 004	Aug 27, 2009	Apr	CRLD
+	136,000USP UNITS;25,000USP UNITS;85,000USP UNITS	N022210 006	Jul 13, 2011	Apr	NEWA

TABLET; ORAL

VIOKACE

APITALIS PHARMA US	39,150USP UNITS;10,440USP UNITS;39,150USP UNITS	N022542 001 Mar 01, 2012 Mar NEWA
+	78,300USP UNITS;20,880USP UNITS;78,300USP UNITS	N022542 002 Mar 01, 2012 Mar NEWA

PANTOPRAZOLE SODIUM

TABLET, DELAYED RELEASE; ORAL

PANTOPRAZOLE SODIUM

AB APOTEX INC	EQ 20MG BASE	A090807 001 May 02, 2012 Apr NEWA
AB	EQ 40MG BASE	A090807 002 May 02, 2012 Apr NEWA
AB MACLEODS PHARMS LTD	EQ 20MG BASE	A200821 001 Feb 16, 2012 Jan NEWA
AB	EQ 40MG BASE	A200821 002 Feb 16, 2012 Jan NEWA
AB RANBAXY LABS LTD	EQ 20MG BASE	A200794 001 May 02, 2012 Apr NEWA
AB	EQ 40MG BASE	A200794 002 May 02, 2012 Apr NEWA

PAROXETINE HYDROCHLORIDE

TABLET; ORAL

PAROXETINE HYDROCHLORIDE

AB ACTAVIS ELIZABETH	EQ 10MG BASE	A076968 001 Jun 21, 2010 Feb CMFD
AB	EQ 20MG BASE	A076968 002 Jun 21, 2010 Feb CMFD
AB	EQ 30MG BASE	A076968 003 Jun 21, 2010 Feb CMFD
AB	EQ 40MG BASE	A076968 004 Jun 21, 2010 Feb CMFD

PEGAPTANIB SODIUM

INJECTABLE; INTRAVITREAL

MACUGEN

+ VALEANT PHARMS LLC	EQ 0.3MG ACID/0.09ML	N021756 001 Dec 17, 2004 May CAHN
----------------------	----------------------	-----------------------------------

PEGINESATIDE ACETATE

SOLUTION; INTRAVENOUS, SUBCUTANEOUS

OMONTYS

+ AFFYMAX	EQ 10MG BASE/ML (EQ 10MG BASE/ML)	N202799 007 Mar 27, 2012 Mar NEWA
+	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	N202799 008 Mar 27, 2012 Mar NEWA
OMONTYS PRESERVATIVE FREE		
+ AFFYMAX	EQ 1MG BASE/0.5ML (EQ 1MG BASE/0.5ML)	N202799 001 Mar 27, 2012 Mar NEWA
+	EQ 2MG BASE/0.5ML (EQ 2MG BASE/0.5ML)	N202799 002 Mar 27, 2012 Mar NEWA
+	EQ 3MG BASE/0.5ML (EQ 3MG BASE/0.5ML)	N202799 003 Mar 27, 2012 Mar NEWA
+	EQ 4MG BASE/0.5ML (EQ 4MG BASE/0.5ML)	N202799 004 Mar 27, 2012 Mar NEWA
+	EQ 5MG BASE/0.5ML (EQ 5MG BASE/0.5ML)	N202799 005 Mar 27, 2012 Mar NEWA
+	EQ 6MG BASE/0.5ML (EQ 6MG BASE/0.5ML)	N202799 006 Mar 27, 2012 Mar NEWA

PENTOBARBITAL SODIUM

INJECTABLE; INJECTION

NEMBUTAL SODIUM

+ OAK PHARMS	50MG/ML	A083246 001 Feb CAHN
--------------	---------	----------------------

PERINDOPRIL ERBUMINE

TABLET; ORAL

ACEON

AB XOMA	2MG	N020184 001 Dec 30, 1993 Jan CAHN
---------	-----	-----------------------------------

TABLET; ORAL

ACEON

AB	XOMA	4MG	N020184 002 Dec 30, 1993 Jan CAHN
AB	+	8MG	N020184 003 Dec 30, 1993 Jan CAHN

PERMETHRIN

CREAM; TOPICAL

ELIMITE

AB	+	RENAISSANCE PHARMA	5%	N019855 001 Aug 25, 1989 May CAHN
----	---	--------------------	----	-----------------------------------

PHENOXYBENZAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIBENZYLINE

AB	+	WELLSPRING PHARM	10MG	N008708 001 Jun CFTG
AB			PHENOXYBENZAMINE HYDROCHLORIDE	
AB		ROXANE	10MG	A201050 001 Jul 16, 2012 Jun NEWA

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

PHENTERMINE HYDROCHLORIDE

>D>	@ ACTAVIS TOTOWA	15MG	A040460 001 Jan 14, 2003 Jul CAHN
>D>	@	30MG	A040227 001 Jun 18, 1997 Jul CAHN
>D>	@	30MG	A040448 001 Jan 22, 2003 Jul CAHN
>D>	@	37.5MG	A040228 001 Jun 19, 1997 Jul CAHN
AA	LANNETT	15MG	A087022 002 Jan 20, 2012 Feb NEWA
>A>	@ MIKAH PHARMA	15MG	A040460 001 Jan 14, 2003 Jul CAHN
>A>	@	30MG	A040448 001 Jan 22, 2003 Jul CAHN
>A>	@	30MG	A040227 001 Jun 18, 1997 Jul CAHN
>A>	@	37.5MG	A040228 001 Jun 19, 1997 Jul CAHN

TABLET; ORAL

PHENTERMINE HYDROCHLORIDE

>D>	@ ACTAVIS TOTOWA	37.5MG	A040190 001 May 30, 1997 Jul CAHN
>A>	@ MIKAH PHARMA	37.5MG	A040190 001 May 30, 1997 Jul CAHN
	@ SANDOZ INC	30MG	A088605 001 Sep 28, 1987 Feb DISC

TABLET, ORALLY DISINTEGRATING; ORAL

SUPRENZA

CITIUS PHARMS

	30MG	N202088 002 Jun 13, 2011 Apr CRLD
+	37.5MG	N202088 003 Mar 27, 2012 Apr CRLD
	37.5MG	N202088 003 Mar 27, 2012 Mar NEWA

>A> PHENTERMINE HYDROCHLORIDE; TOPIRAMATE

CAPSULE, EXTENDED RELEASE; ORAL

QSYMIA

VIVUS

EQ 3.75MG BASE;23MG

N022580 001 Jul 17, 2012 Jul NEWA

EQ 7.5MG BASE;46MG

N022580 002 Jul 17, 2012 Jul NEWA

EQ 11.25MG BASE;69MG

N022580 003 Jul 17, 2012 Jul NEWA

>A> + EQ 15MG BASE;92MG

N022580 004 Jul 17, 2012 Jul NEWA

PHENTERMINE RESIN COMPLEX

CAPSULE, EXTENDED RELEASE; ORAL

PHENTERMINE RESIN COMPLEX

+	LANNETT HOLDINGS INC	EQ 15MG BASE	A040872 001 Jul 28, 2011 Jan CRLD
+		EQ 30MG BASE	A040872 002 Jul 28, 2011 Jan CRLD

PHENYTOIN SODIUM

INJECTABLE; INJECTION

PHENYTOIN SODIUM

AP + HIKMA MAPLE 50MG/ML A084307 001 Jun CAHN

PHYTONADIONE

INJECTABLE; INJECTION

PHYTONADIONE

BP + INTL MEDICATION 1MG/0.5ML A083722 001 Feb CRLD
TABLET; ORAL
MEPHYTON
+ VALEANT PHARMS 5MG N010104 003 Jun CAHNPINDOLOL

TABLET; ORAL

PINDOLOL

@ MYLAN PHARMS INC 5MG A074013 001 Sep 24, 1992 May CAHN
@ 10MG A074018 001 Sep 24, 1992 May CAHNPIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

ACTOS

TAKEDA PHARMS USA EQ 15MG BASE N021073 001 Jul 15, 1999 Mar CAHN
EQ 30MG BASE N021073 002 Jul 15, 1999 Mar CAHN
+ EQ 45MG BASE N021073 003 Jul 15, 1999 Mar CAHNPOTASSIUM CHLORIDE

TABLET, EXTENDED RELEASE; ORAL

K+10

@ FUTURE PAK 10MEQ A070999 001 Oct 22, 1987 Mar DISC

KAON CL-10

@ SAVAGE LABS 10MEQ N017046 002 Mar DISC

KLOR-CON M20

AB + UPSHER SMITH LABS 20MEQ A074726 001 Nov 20, 1998 Mar CRLD
KLOTRIX
@ APOTHECON 10MEQ N017850 001 Mar DISC
POTASSIUM CHLORIDE
@ NESHER PHARMS 20MEQ A076044 001 Apr 05, 2002 Mar DISC
@ SCHERING 10MEQ N019439 002 Jun 13, 1986 Mar DISC
@ 20MEQ N019439 001 Jun 13, 1986 Mar DISCPOTASSIUM CITRATE

FOR SOLUTION; ORAL

POTASSIUM CITRATE

@ NOVA K 10MEQ/PACKET N019647 002 Oct 13, 1988 Feb CAHN
@ 20MEQ/PACKET N019647 001 Oct 13, 1988 Feb CAHNPRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

AB APOTEX INC 0.125MG A090151 001 Apr 30, 2012 Apr NEWA
AB 0.25MG A090151 002 Apr 30, 2012 Apr NEWA
AB 0.5MG A090151 003 Apr 30, 2012 Apr NEWA
AB 0.75MG A090151 006 Apr 30, 2012 Apr NEWA
AB 1MG A090151 004 Apr 30, 2012 Apr NEWA

TABLET; ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

AB APOTEX INC 1.5MG A090151 005 Apr 30, 2012 Apr NEWA

PRAVASTATIN SODIUM

TABLET; ORAL

PRAVASTATIN SODIUM

@ PLIVA HRVATSKA DOO	10MG	A077730 001	Nov 21, 2006	Jun	DISC
@	20MG	A077730 002	Nov 21, 2006	Jun	DISC
@	30MG	A077730 003	Nov 21, 2006	Jun	DISC
@	40MG	A077730 005	Nov 21, 2006	Jun	DISC
@ RANBAXY LABS LTD	10MG	A076445 001	Apr 23, 2007	Apr	DISC
@	20MG	A076445 002	Apr 23, 2007	Apr	DISC
@	40MG	A076445 003	Apr 23, 2007	Apr	DISC
@	80MG	A076445 004	Apr 23, 2007	Apr	DISC

PREDNICARBATE

CREAM; TOPICAL

PREDNICARBATE

AB FOUGERA PHARMS 0.1% A077287 001 Sep 19, 2006 Jan CAHN

OINTMENT; TOPICAL

PREDNICARBATE

AB FOUGERA PHARMS 0.1% A077236 001 Mar 09, 2007 Jan CAHN

PREDNISOLONE

SYRUP; ORAL

PREDNISOLONE

>D> AA HI TECH PHARMA	15MG/5ML	A040401 001	Feb 27, 2003	Jul	CRLD
>A> AA + HI TECH PHARMA CO	15MG/5ML	A040401 001	Feb 27, 2003	Jul	CRLD
@ NESHER PHARMS	5MG/5ML	A040423 001	Oct 22, 2001	Jun	DISC
@	15MG/5ML	A040364 001	Apr 10, 2002	Jun	DISC

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION; ORAL

PREDNISOLONE SODIUM PHOSPHATE

>A> + MISSION PHARMA	EQ 25MG BASE/5ML	A091396 001	Sep 13, 2010	Jul	CAHN
@ NESHER PHARMS	EQ 5MG BASE/5ML	A076982 001	May 24, 2005	Jun	DISC
>D> + PHARM ASSOC	EQ 25MG BASE/5ML	A091396 001	Sep 13, 2010	Jul	CAHN

PREDNISONE

>A> TABLET, DELAYED RELEASE; ORAL					
>A> RAYOS					
>A> HORIZON PHARMA	1MG	N202020 001	Jul 26, 2012	Jul	NEWA
>A>	2MG	N202020 002	Jul 26, 2012	Jul	NEWA
>A> +	5MG	N202020 003	Jul 26, 2012	Jul	NEWA

PREGABALIN

CAPSULE; ORAL

LYRICA

AB PF PRISM	25MG	N021446 001	Dec 30, 2004	Jun	CFTG
	25MG	N021446 001	Dec 30, 2004	Jan	CAHN
AB	50MG	N021446 002	Dec 30, 2004	Jun	CFTG
	50MG	N021446 002	Dec 30, 2004	Jan	CAHN
AB	75MG	N021446 003	Dec 30, 2004	Jun	CFTG
	75MG	N021446 003	Dec 30, 2004	Jan	CAHN
	100MG	N021446 004	Dec 30, 2004	Jan	CAHN

CAPSULE; ORAL

LYRICA

AB		PF PRISM	150MG	N021446	005	Dec 30,	2004	Jun	CFTG
			150MG	N021446	005	Dec 30,	2004	Jan	CAHN
AB			200MG	N021446	006	Dec 30,	2004	Jun	CFTG
			200MG	N021446	006	Dec 30,	2004	Jan	CAHN
AB			225MG	N021446	007	Dec 30,	2004	Jun	CFTG
			225MG	N021446	007	Dec 30,	2004	Jan	CAHN
AB	+		300MG	N021446	008	Dec 30,	2004	Jun	CFTG
	+		300MG	N021446	008	Dec 30,	2004	Jan	CAHN
PREGABALIN									
AB	LUPIN LTD		25MG	A091040	001	Jul 03,	2012	Jun	NEWA
AB			50MG	A091040	002	Jul 03,	2012	Jun	NEWA
AB			75MG	A091040	003	Jul 03,	2012	Jun	NEWA
AB			100MG	A091040	004	Jul 03,	2012	Jun	NEWA
AB			150MG	A091040	005	Jul 03,	2012	Jun	NEWA
AB			200MG	A091040	006	Jul 03,	2012	Jun	NEWA
AB			225MG	A091040	007	Jul 03,	2012	Jun	NEWA
AB			300MG	A091040	008	Jul 03,	2012	Jun	NEWA
AB	TEVA PHARMS		25MG	A091219	001	Jul 03,	2012	Jun	NEWA
AB			50MG	A091219	002	Jul 03,	2012	Jun	NEWA
AB			75MG	A091224	001	Jul 03,	2012	Jun	NEWA
AB			100MG	A091224	002	Jul 03,	2012	Jun	NEWA
AB			150MG	A091224	003	Jul 03,	2012	Jun	NEWA
AB			200MG	A091224	004	Jul 03,	2012	Jun	NEWA
AB			225MG	A091224	005	Jul 03,	2012	Jun	NEWA
AB			300MG	A091224	006	Jul 03,	2012	Jun	NEWA
AB	WATSON LABS INC		25MG	A091221	001	Jul 03,	2012	Jun	NEWA
AB			50MG	A091221	002	Jul 03,	2012	Jun	NEWA
AB			75MG	A091221	003	Jul 03,	2012	Jun	NEWA
AB			100MG	A091221	004	Jul 03,	2012	Jun	NEWA
AB			150MG	A091221	005	Jul 03,	2012	Jun	NEWA
AB			200MG	A091221	006	Jul 03,	2012	Jun	NEWA
AB			225MG	A091221	007	Jul 03,	2012	Jun	NEWA
AB			300MG	A091221	008	Jul 03,	2012	Jun	NEWA

SOLUTION; ORAL

LYRICA

+ PF PRISM 20MG/ML N022488 001 Jan 04, 2010 Jan CAHN

PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST PLAIN DENTAL

@ DENTSPLY PHARM 4% N021382 001 May DISC
PRILOCAINE HYDROCHLORIDE
SEPTODONT INC 4% A070235 001 Sep 29 2010 May CRBL

Procainamide Hydrochloride

INJECTABLE: INJECTION

BROCAINAMIDE HYDROCHLORIDE

@ HIKMA MAPLE 100MG/ML A089029 001 Apr 17, 1986 Jun CAHN
@ 500MG/ML A089030 001 Apr 17, 1986 Jun CAHN

PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

@ HIKMA MAPLE EQ 5MG BASE/ML A089523 001 May 03, 1988 Jun CAHN

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

AP + HIKMA MAPLE EQ 5MG BASE/ML A089903 001 Aug 29, 1989 Jun CAHN

PROGESTERONE

CAPSULE; ORAL

PROGESTERONE

AB TEVA PHARMS 100MG A202121 001 Feb 29, 2012 Feb NEWA

AB 200MG A202121 002 Feb 29, 2012 Feb NEWA

PROMETRIUM

AB ABBOTT LABS 100MG N019781 001 May 14, 1998 Feb CFTG

AB + 200MG N019781 002 Oct 15, 1999 Feb CFTG

PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETHAZINE HYDROCHLORIDE

AA TRIS PHARMA INC 6.25MG/5ML A091675 001 Jun 28, 2012 Jun NEWA

PROPAFENONE HYDROCHLORIDE

TABLET; ORAL

PROPAFENONE HYDROCHLORIDE

@ NESHER PHARMS 150MG A076193 001 Feb 07, 2002 Jun DISC

@ 225MG A076193 002 Feb 07, 2002 Jun DISC

@ 300MG A076193 003 Feb 07, 2002 Jun DISC

PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

VIVACTIL

>D> @ ODYSSEY PHARMS 5MG N016012 001 Jul CAHN

>D> @ 10MG N016012 002 Jul CAHN

>A> @ TEVA WOMENS R AND D 5MG N016012 001 Jul CAHN

>A> @ 10MG N016012 002 Jul CAHN

PYRIDOSTIGMINE BROMIDE

TABLET; ORAL

MESTINON

AB + VALEANT PHARMS LLC 60MG N009829 002 Jun CAHN

TABLET, EXTENDED RELEASE; ORAL

MESTINON

+ VALEANT PHARMS LLC 180MG N011665 001 Jun CAHN

QUETIAPINE FUMARATE

TABLET; ORAL

QUETIAPINE FUMARATE

AB ACCORD HLTHCARE INC EQ 25MG BASE A202152 001 Mar 27, 2012 Mar NEWA

AB EQ 50MG BASE A202152 002 Mar 27, 2012 Mar NEWA

AB EQ 100MG BASE A202152 003 Mar 27, 2012 Mar NEWA

AB EQ 200MG BASE A202152 004 Mar 27, 2012 Mar NEWA

AB EQ 300MG BASE A202152 005 Mar 27, 2012 Mar NEWA

AB EQ 400MG BASE A202152 006 Mar 27, 2012 Mar NEWA

AB APOTEX INC EQ 25MG BASE A090960 001 Mar 27, 2012 Mar NEWA

AB EQ 50MG BASE A090960 002 Mar 27, 2012 Mar NEWA

AB EQ 100MG BASE A090960 003 Mar 27, 2012 Mar NEWA

AB EQ 200MG BASE A090960 004 Mar 27, 2012 Mar NEWA

AB EQ 300MG BASE A090960 005 Mar 27, 2012 Mar NEWA

AB EQ 400MG BASE A090960 006 Mar 27, 2012 Mar NEWA

TABLET; ORAL

QUETIAPINE FUMARATE

AB	AUROBINDO PHARMA LTD	EQ 25MG BASE	A091388 001	Mar 27, 2012	Mar	NEWA
AB		EQ 50MG BASE	A091388 002	Mar 27, 2012	Mar	NEWA
AB		EQ 100MG BASE	A091388 003	Mar 27, 2012	Mar	NEWA
AB		EQ 150MG BASE	A091388 004	Mar 27, 2012	Mar	NEWA
AB		EQ 200MG BASE	A091388 005	Mar 27, 2012	Mar	NEWA
AB		EQ 300MG BASE	A091388 006	Mar 27, 2012	Mar	NEWA
AB		EQ 400MG BASE	A091388 007	Mar 27, 2012	Mar	NEWA
AB	DR REDDYS LABS LTD	EQ 25MG BASE	A077380 001	Mar 27, 2012	Mar	NEWA
AB		EQ 50MG BASE	A077380 002	Mar 27, 2012	Mar	NEWA
AB		EQ 100MG BASE	A077380 003	Mar 27, 2012	Mar	NEWA
AB		EQ 150MG BASE	A077380 004	Mar 27, 2012	Mar	NEWA
AB		EQ 200MG BASE	A077380 005	Mar 27, 2012	Mar	NEWA
AB		EQ 300MG BASE	A077380 006	Mar 27, 2012	Mar	NEWA
AB		EQ 400MG BASE	A077380 007	Mar 27, 2012	Mar	NEWA
AB	LUPIN LTD	EQ 25MG BASE	A201109 001	Mar 27, 2012	Mar	NEWA
AB		EQ 50MG BASE	A201109 002	Mar 27, 2012	Mar	NEWA
AB		EQ 100MG BASE	A201109 003	Mar 27, 2012	Mar	NEWA
AB		EQ 200MG BASE	A201109 004	Mar 27, 2012	Mar	NEWA
AB		EQ 300MG BASE	A201109 005	Mar 27, 2012	Mar	NEWA
AB		EQ 400MG BASE	A201109 006	Mar 27, 2012	Mar	NEWA
AB	MYLAN PHARMS INC	EQ 25MG BASE	A090323 001	Mar 27, 2012	Mar	NEWA
AB	ROXANE	EQ 25MG BASE	A090120 001	Mar 27, 2012	Mar	NEWA
AB		EQ 50MG BASE	A090749 001	Mar 27, 2012	Mar	NEWA
AB		EQ 100MG BASE	A090749 002	Mar 27, 2012	Mar	NEWA
AB		EQ 200MG BASE	A090749 003	Mar 27, 2012	Mar	NEWA
AB		EQ 300MG BASE	A090749 004	Mar 27, 2012	Mar	NEWA
AB		EQ 400MG BASE	A090749 005	Mar 27, 2012	Mar	NEWA
AB	SUN PHARMA GLOBAL	EQ 25MG BASE	A201190 001	Mar 27, 2012	Mar	NEWA
AB		EQ 50MG BASE	A201190 002	Mar 27, 2012	Mar	NEWA
AB		EQ 100MG BASE	A201190 003	Mar 27, 2012	Mar	NEWA
AB		EQ 200MG BASE	A201190 004	Mar 27, 2012	Mar	NEWA
AB		EQ 300MG BASE	A201190 005	Mar 27, 2012	Mar	NEWA
AB		EQ 400MG BASE	A201190 006	Mar 27, 2012	Mar	NEWA
AB	TEVA PHARMS	EQ 25MG BASE	A077745 001	Mar 27, 2012	Mar	NEWA
AB		EQ 50MG BASE	A077745 002	Mar 27, 2012	Mar	NEWA
AB		EQ 100MG BASE	A077745 003	Mar 27, 2012	Mar	NEWA
AB		EQ 150MG BASE	A077745 004	Mar 27, 2012	Mar	NEWA
AB		EQ 200MG BASE	A077745 005	Mar 27, 2012	Mar	NEWA
AB		EQ 300MG BASE	A077745 006	Mar 27, 2012	Mar	NEWA
AB		EQ 400MG BASE	A077745 007	Mar 27, 2012	Mar	NEWA
AB	TORRENT PHARMS LTD	EQ 25MG BASE	A200363 001	Mar 27, 2012	Mar	NEWA
AB		EQ 50MG BASE	A200363 002	Mar 27, 2012	Mar	NEWA
AB		EQ 100MG BASE	A200363 003	Mar 27, 2012	Mar	NEWA
AB		EQ 200MG BASE	A200363 004	Mar 27, 2012	Mar	NEWA
AB		EQ 300MG BASE	A200363 005	Mar 27, 2012	Mar	NEWA
AB		EQ 400MG BASE	A200363 006	Mar 27, 2012	Mar	NEWA
	SEROQUEL					
AB	+ ASTRAZENECA	EQ 25MG BASE	N020639 001	Sep 26, 1997	Mar	CFTG
AB		EQ 50MG BASE	N020639 007	Oct 04, 2005	Mar	CFTG
AB		EQ 100MG BASE	N020639 002	Sep 26, 1997	Mar	CFTG
AB		EQ 200MG BASE	N020639 003	Sep 26, 1997	Mar	CFTG
AB	+	EQ 300MG BASE	N020639 005	Jul 26, 2000	Mar	CFTG
AB		EQ 400MG BASE	N020639 006	Oct 04, 2005	Mar	CFTG

RAMELTEON

TABLET; ORAL
ROZEREM

>D>	+ TAKEDA GLOBAL	8MG	N021782 001 Jul 22, 2005 Jul CAHN
>A>	+ TAKEDA PHARMS USA	8MG	N021782 001 Jul 22, 2005 Jul CAHN

RAMIPRIL

CAPSULE; ORAL
RAMIPRIL

@ RANBAXY LABS LTD	5MG	A078849 001 Mar 06, 2009 Apr DISC
@	10MG	A078849 002 Mar 06, 2009 Apr DISC

TABLET; ORAL
ALTACE

@ KING PFIZER	1.25MG	N022021 001 Feb 27, 2007 Jan DISC
@	2.5MG	N022021 002 Feb 27, 2007 Jan DISC
@	5MG	N022021 003 Feb 27, 2007 Jan DISC
@	10MG	N022021 004 Feb 27, 2007 Jan DISC

RANITIDINE HYDROCHLORIDE

INJECTABLE; INJECTION
ZANTAC

AP + COVIS PHARMA	EQ 25MG BASE/ML	N019090 001 Oct 19, 1984 Mar CAHN
ZANTAC IN PLASTIC CONTAINER		
+ COVIS PHARMA	EQ 1MG BASE/ML	N019593 002 Sep 27, 1991 Mar CAHN
@	EQ 50MG BASE/100ML	N019593 001 Dec 17, 1986 Mar CAHN

RIFAMPIN

CAPSULE; ORAL
RIMACTANE

AB PROSAM LABS	300MG	N050429 001 Feb CMFD
----------------	-------	----------------------

RILPIVIRINE HYDROCHLORIDE

TABLET; ORAL
EDURANT

+ JANSSEN PRODS	EQ 25MG BASE	N202022 001 May 20, 2011 Feb CAHN
-----------------	--------------	-----------------------------------

RISEDRONATE SODIUM

TABLET; ORAL
ACTONEL

AB WARNER CHILCOTT LLC	5MG	N020835 002 Apr 14, 2000 Jun CAHN
AB	30MG	N020835 001 Mar 27, 1998 Jun CAHN
AB +	35MG	N020835 003 May 25, 2002 Jun CAHN
@	75MG	N020835 004 Apr 16, 2007 Jun CAHN
+	150MG	N020835 005 Apr 22, 2008 Jun CAHN

TABLET, DELAYED RELEASE; ORAL
ATELVIA

+ WARNER CHILCOTT LLC	35MG	N022560 001 Oct 08, 2010 Jun CAHN
-----------------------	------	-----------------------------------

RISPERIDONE

TABLET; ORAL
RISPERIDONE

AB PROSAM LABS	0.25MG	A078071 001 Jun 17, 2009 Feb CMFD
AB	0.5MG	A078071 002 Jun 17, 2009 Feb CMFD
AB	1MG	A078071 003 Jun 17, 2009 Feb CMFD
AB	2MG	A078071 004 Jun 17, 2009 Feb CMFD

TABLET; ORAL

RISPERIDONE

AB	PROSAM LABS	3MG	A078071 005	Jun 17, 2009	Feb	CMFD
AB		4MG	A078071 006	Jun 17, 2009	Feb	CMFD

TABLET, ORALLY DISINTEGRATING; ORAL

RISPERIDONE

AB	TEVA	0.5MG	A076908 001	Mar 12, 2012	Feb	NEWA
AB		1MG	A076908 002	Mar 12, 2012	Feb	NEWA
AB		2MG	A076908 003	Mar 12, 2012	Feb	NEWA

RIVASTIGMINE TARTRATE

CAPSULE; ORAL

RIVASTIGMINE TARTRATE

AB	ALEMBIC PHARMS LTD	EQ 1.5MG BASE	A091689 001	Jun 12, 2012	May	NEWA
AB		EQ 3MG BASE	A091689 002	Jun 12, 2012	May	NEWA
AB		EQ 4.5MG BASE	A091689 003	Jun 12, 2012	May	NEWA
AB		EQ 6MG BASE	A091689 004	Jun 12, 2012	May	NEWA

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

ROPINIROLE HYDROCHLORIDE

AB	APOTEX	EQ 0.25MG BASE	A079165 001	Feb 07, 2012	Jan	NEWA
AB		EQ 0.5MG BASE	A079165 002	Feb 07, 2012	Jan	NEWA
AB		EQ 1MG BASE	A079165 003	Feb 07, 2012	Jan	NEWA
AB		EQ 2MG BASE	A079165 004	Feb 07, 2012	Jan	NEWA
AB		EQ 3MG BASE	A079165 005	Feb 07, 2012	Jan	NEWA
AB		EQ 4MG BASE	A079165 006	Feb 07, 2012	Jan	NEWA
AB		EQ 5MG BASE	A079165 007	Feb 07, 2012	Jan	NEWA

TABLET, EXTENDED RELEASE; ORAL

REQUIP XL

AB	+ SMITHKLINE BEECHAM	EQ 2MG BASE	N022008 001	Jun 13, 2008	Apr	CFTG
AB		EQ 4MG BASE	N022008 003	Jun 13, 2008	Apr	CFTG
AB		EQ 6MG BASE	N022008 006	Apr 10, 2009	Apr	CFTG
AB		EQ 8MG BASE	N022008 004	Jun 13, 2008	Apr	CFTG
AB		EQ 12MG BASE	N022008 005	Oct 31, 2008	Apr	CFTG

ROPINIROLE HYDROCHLORIDE

AB	ACTAVIS	EQ 2MG BASE	A090869 001	May 17, 2012	Apr	NEWA
AB		EQ 4MG BASE	A090869 002	May 17, 2012	Apr	NEWA
AB		EQ 6MG BASE	A090869 003	May 17, 2012	Apr	NEWA
AB		EQ 8MG BASE	A090869 004	May 17, 2012	Apr	NEWA
AB		EQ 12MG BASE	A090869 005	May 17, 2012	Apr	NEWA
AB	DR REDDYS LABS LTD	EQ 2MG BASE	A201576 001	Jun 06, 2012	May	NEWA
AB		EQ 4MG BASE	A201576 002	Jun 06, 2012	May	NEWA
AB		EQ 6MG BASE	A201576 003	Jun 06, 2012	May	NEWA
AB		EQ 8MG BASE	A201576 004	Jun 06, 2012	May	NEWA
AB		EQ 12MG BASE	A201576 005	Jun 06, 2012	May	NEWA
AB	SANDOZ INC	EQ 2MG BASE	A201047 001	Jun 06, 2012	May	NEWA
AB		EQ 4MG BASE	A201047 003	Jun 06, 2012	May	NEWA
AB		EQ 6MG BASE	A201047 004	Jun 06, 2012	May	NEWA
AB		EQ 8MG BASE	A201047 005	Jun 06, 2012	May	NEWA
AB		EQ 12MG BASE	A201047 006	Jun 06, 2012	May	NEWA
AB	WATSON LABS INC	EQ 2MG BASE	A200431 001	Jun 06, 2012	May	NEWA
AB		EQ 4MG BASE	A200431 002	Jun 06, 2012	May	NEWA
AB		EQ 6MG BASE	A200431 003	Jun 06, 2012	May	NEWA
AB		EQ 8MG BASE	A200431 004	Jun 06, 2012	May	NEWA
AB		EQ 12MG BASE	A200431 005	Jun 06, 2012	May	NEWA

ROTIGOTINE

FILM, EXTENDED RELEASE; TRANSDERMAL

NEUPRO

UCB INC

1MG/24HR

N021829 004 Apr 02, 2012 Apr NEWA

2MG/24HR

N021829 001 May 09, 2007 Apr CMFD

3MG/24HR

N021829 005 Apr 02, 2012 Apr NEWA

4MG/24HR

N021829 002 May 09, 2007 Apr CMFD

6MG/24HR

N021829 003 May 09, 2007 Apr CMFD

+

8MG/24HR

N021829 006 Apr 02, 2012 Apr NEWA

SAMARIUM SM-153 LEXIDRONAM PENTASODIUM

INJECTABLE; INJECTION

QUADRAMET

>D>	+	EUSA PHARMA USA	50mCi/ML	N020570 001 Mar 28, 1997 Jul CAHN
>A>	+	JAZZ EUSA PHARMA	50mCi/ML	N020570 001 Mar 28, 1997 Jul CAHN

SERTACONAZOLE NITRATE

CREAM; TOPICAL

ERTACZO

+ VALEANT INTL

2%

N021385 001 Dec 10, 2003 Jan CAHN

SERTRALINE HYDROCHLORIDE

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

AB	PROSAM LABS	EQ 25MG BASE	A078175 001 Jul 21, 2010 Feb CMFD
AB		EQ 50MG BASE	A078175 002 Jul 21, 2010 Feb CMFD
AB		EQ 100MG BASE	A078175 003 Jul 21, 2010 Feb CMFD

SILVER SULFADIAZINE

CREAM; TOPICAL

THERMAZENE

AB	THEPHARMANETWORK LLC	1%	N018810 001 Dec 23, 1985 Jun CAHN
----	----------------------	----	-----------------------------------

SIMVASTATIN

TABLET; ORAL

SIMVASTATIN

AB	PROSAM LABS	5MG	A078735 001 Aug 30, 2010 Feb CMFD
AB		10MG	A078735 002 Aug 30, 2010 Feb CMFD
AB		20MG	A078735 003 Aug 30, 2010 Feb CMFD
AB		40MG	A078735 004 Aug 30, 2010 Feb CMFD
AB		80MG	A078735 005 Aug 30, 2010 Feb CMFD

SITAGLIPTIN PHOSPHATE

TABLET; ORAL

JANUVIA

MERCK SHARP DOHME

EQ 25MG BASE

N021995 001 Oct 16, 2006 May CAHN

EQ 50MG BASE

N021995 002 Oct 16, 2006 May CAHN

+

EQ 100MG BASE

N021995 003 Oct 16, 2006 May CAHN

SODIUM BICARBONATE; TARTARIC ACID

GRANULE, EFFERVESCENT; ORAL

BAROS

@ MALLINCKRODT INC

460MG/GM;420MG/GM

N018509 001 Aug 07, 1985 Jan CAHN

SODIUM CHLORIDE

INJECTABLE; INJECTION
SODIUM CHLORIDE 0.9%
AP HIKMA (MAPLE) 9MG/ML A201850 001 Jan 20, 2012 Jan NEWA
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
MEDEFIL 9MG/ML N202832 001 Jan 06, 2012 Jan NEWA

SODIUM CHROMATE CR-51

INJECTABLE; INJECTION
CHROMITOPE SODIUM
@ BRACCO 200uCi/ML N013993 001 Jun DISC

SODIUM FERRIC GLUCONATE COMPLEX

INJECTABLE; INJECTION
SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE
AB HIKMA PHARMS 62.5MG/5ML A078215 001 Mar 31, 2011 May CAHN

SODIUM IODIDE I-123

CAPSULE; ORAL
SODIUM IODIDE I 123
@ GE HEALTHCARE 100uCi N017630 001 May DISC
SOLUTION; ORAL
SODIUM IODIDE I 123
>D> + GE HEALTHCARE 2mCi/ML N017630 002 Jul DISC
>A> @ 2mCi/ML N017630 002 Jul DISC

SODIUM IODIDE I-131

CAPSULE; ORAL
SODIUM IODIDE I-131
JUBILANT DRAXIMAGE 9-100mCi N021305 006 May 19, 2005 Feb CAHN
@ 2-200mCi N021305 004 Nov 18, 2004 Jan DISC
SOLUTION; ORAL
HICON
@ JUBILANT DRAXIMAGE 1-500mCi/0.5ML N021305 003 Jan 24, 2003 Feb DISC
@ 1-250mCi/0.25ML N021305 002 Jan 24, 2003 Feb DISC
@ 1-1000mCi/ML N021305 005 Apr 04, 2006 Feb DISC
+ 250-1000mCi N021305 007 Dec 05, 2011 Jan NEWA

SODIUM NITRITE

SOLUTION; INTRAVENOUS
SODIUM NITRITE
+ HOPE PHARMS 300MG/10ML (30MG/ML) N203922 001 Feb 14, 2012 Feb NEWA

SODIUM OXYBATE

SOLUTION; ORAL
XYREM
+ JAZZ PHARMS 500MG/ML N021196 001 Jul 17, 2002 Jun CAHN

SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL
SODIUM POLYSTYRENE SULFONATE
@ CITRUSPHRMA 454GM/BOT A040909 001 Dec 03, 2008 Jun DISC

SODIUM THIOSULFATE

SOLUTION; INTRAVENOUS
 SODIUM THIOSULFATE
 + HOPE PHARMS 12.5GM/50ML (250MG/ML) N203923 001 Feb 14, 2012 Feb NEWA

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION

>D>	NORDITROPIN							
>D> BX	NOVO NORDISK INC	5MG/1.5ML		N021148	001	Jun 20, 2000	Jul	DISC
>A>	@	5MG/1.5ML		N021148	001	Jun 20, 2000	Jul	DISC
>D> BX		10MG/1.5ML		N021148	002	Jun 20, 2000	Jul	DISC
>A>	@	10MG/1.5ML		N021148	002	Jun 20, 2000	Jul	DISC
>D>	+	15MG/1.5ML		N021148	003	Jun 20, 2000	Jul	DISC
>A>	@	15MG/1.5ML		N021148	003	Jun 20, 2000	Jul	DISC

SOTALOL HYDROCHLORIDETABLET; ORAL
 SOTALOL HYDROCHLORIDE

AB2	EPIC PHARMA INC	80MG	A077070	001	Nov 04, 2005	Jun	CAHN
AB2		120MG	A077070	002	Nov 04, 2005	Jun	CAHN
AB2		160MG	A077070	003	Nov 04, 2005	Jun	CAHN

SPINOSAD

SUSPENSION; TOPICAL

NATROBA							
+ PARAPRO LLC		0.9%	N022408	001	Jan 18, 2011	May	CAHN

SUCCIMERCAPSULE; ORAL
 CHEMET
 + LUNDBECK LLC

100MG	N019998	002	Jan 30, 1991	Mar	CAHN
-------	---------	-----	--------------	-----	------

SULFACETAMIDE SODIUMLOTION; TOPICAL
 SULFACETAMIDE SODIUM

AB	FOUGERA PHARMS	10%	A077015	001	Nov 17, 2006	Jan	CAHN
----	----------------	-----	---------	-----	--------------	-----	------

SULFANILAMIDECREAM; VAGINAL
 AVC

+ JAZZ PHARMS COMMERCIAL	15%	N006530	003	Jan 27, 1987	Mar	CAHN
+ JAZZ PHARMS II	15%	N006530	003	Jan 27, 1987	Jul	CAHN
>D> + JAZZ PHARMS III	15%	N006530	003	Jan 27, 1987	Jul	CAHN
+ @	15%	N006530	003	Jan 27, 1987	Jun	CAHN

SUPPOSITORY; VAGINAL

AVC							
@ JAZZ PHARMS COMMERCIAL	1.05GM	N006530	004	Jan 27, 1987	Mar	CAHN	
@ JAZZ PHARMS II	1.05GM	N006530	004	Jan 27, 1987	Jul	CAHN	
>D> @ JAZZ PHARMS III	1.05GM	N006530	004	Jan 27, 1987	Jul	CAHN	
@	1.05GM	N006530	004	Jan 27, 1987	Jun	CAHN	

SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS
ALSUMA

BX	MERIDIAN MEDCL	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N022377 001 Jun 29, 2010 Jun CRLD
AP	SUMATRIPTAN SUCCINATE INJECTALIA	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A090310 001 Aug 11, 2010 Mar CAHN
BX	+ ZOGENIX INC	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N022239 001 Jul 15, 2009 Jun CTEC
AB	TABLET; ORAL SUMATRIPTAN SUCCINATE APOTEX INC	EQ 25MG BASE	A200263 001 Jun 19, 2012 Jun NEWA
AB		EQ 50MG BASE	A200263 002 Jun 19, 2012 Jun NEWA
AB		EQ 100MG BASE	A200263 003 Jun 19, 2012 Jun NEWA

TAFLUPROST

SOLUTION/DROPS; OPHTHALMIC
ZIOPTAN

+ MERCK SHARP DOHME	0.0015%	N202514 001 Feb 10, 2012 Feb NEWA
---------------------	---------	-----------------------------------

TALIGLUCERASE ALFA

POWDER; IV (INFUSION)
ELELYSO

+ PFIZER	200 UNITS/VIAL	N022458 001 May 01, 2012 May NEWA
----------	----------------	-----------------------------------

TAMOXIFEN CITRATE

TABLET; ORAL
TAMOXIFEN CITRATE

@ AEGIS PHARMS	EQ 10MG BASE	A076398 001 Mar 31, 2003 Jun DISC
@	EQ 20MG BASE	A076398 002 Mar 31, 2003 Jun DISC

TAZAROTENE

AEROSOL, FOAM; TOPICAL
FABIOR

+ STIEFEL LABS INC	0.1%	N202428 001 May 11, 2012 May NEWA
--------------------	------	-----------------------------------

TECHNETIUM TC-99M MEDRONATE

INJECTABLE; INJECTION
DRAXIMAGE MDP-10

@ JUBILANT DRAXIMAGE	N/A	N018035 001 May CAHN
DRAXIMAGE MDP-25		
+ JUBILANT DRAXIMAGE	N/A	N018035 002 Feb 27, 2004 May CAHN

TEMAZEPAM

CAPSULE; ORAL
TEMAZEPAM

AB	NOVEL LABS INC	7.5MG	A071457 002 Jun 22, 2012 Jun NEWA
AB		22.5MG	A071457 003 Jun 22, 2012 Jun NEWA

TEMOZOLOMIDE

CAPSULE; ORAL
TEMODAR

AB	MERCK SHARP DOHME	5MG	N021029 001 Aug 11, 1999 Jun CAHN
AB		20MG	N021029 002 Aug 11, 1999 Jun CAHN

CAPSULE; ORAL

TEMODAR

AB	MERCK SHARP DOHME	100MG	N021029 003 Aug 11, 1999 Jun CAHN
AB		140MG	N021029 005 Oct 19, 2006 Jun CAHN
AB		180MG	N021029 006 Oct 19, 2006 Jun CAHN
AB	+	250MG	N021029 004 Aug 11, 1999 Jun CAHN

POWDER; INTRAVENOUS

TEMODAR

+ MERCK SHARP DOHME	100MG/VIAL	N022277 001 Feb 27, 2009 Jun CAHN
---------------------	------------	-----------------------------------

TENOFOVIR DISOPROXIL FUMARATE

POWDER; ORAL

VIREAD

+ GILEAD SCIENCES INC	40MG/SCOOPFUL	N022577 001 Jan 18, 2012 Jan NEWA
-----------------------	---------------	-----------------------------------

TABLET; ORAL

VIREAD

GILEAD SCIENCES INC	150MG	N021356 002 Jan 18, 2012 Apr NEWA
	200MG	N021356 003 Jan 18, 2012 Apr NEWA
	250MG	N021356 004 Jan 18, 2012 Apr NEWA

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

TERAZOSIN HYDROCHLORIDE

@ RANBAXY LABS LTD	EQ 1MG BASE	A076021 001 Aug 22, 2002 Apr DISC
@	EQ 2MG BASE	A076021 002 Aug 22, 2002 Apr DISC
@	EQ 5MG BASE	A076021 003 Aug 22, 2002 Apr DISC
@	EQ 10MG BASE	A076021 004 Aug 22, 2002 Apr DISC

TERBUTALINE SULFATE

INJECTABLE; INJECTION

TERBUTALINE SULFATE

@ TEVA PARENTERAL	1MG/ML	A076853 001 Jul 20, 2004 Jan DISC
-------------------	--------	-----------------------------------

TERCONAZOLE

CREAM; VAGINAL

TERCONAZOLE

AB FOUGERA PHARMS	0.4%	A076712 001 Feb 18, 2005 Jan CAHN
-------------------	------	-----------------------------------

SUPPOSITORY; VAGINAL

TERCONAZOLE

AB FOUGERA PHARMS	80MG	A076850 001 Jul 12, 2006 Jan CAHN
-------------------	------	-----------------------------------

TESAMORELIN ACETATE

POWDER; SUBCUTANEOUS

EGRIFTA

@ EMD SERONO	EQ 1MG BASE/VIAL	N022505 001 Nov 10, 2010 Jun DISC
+	EQ 1MG BASE/VIAL	N022505 001 Nov 10, 2010 May CAHN
+	EQ 2MG BASE/VIAL	N022505 002 Nov 29, 2011 Jun NEWA

TESTOSTERONE

GEL; TRANSDERMAL

ANDROGEL

BX + ABBOTT LABS	1% (50MG/5GM PACKET)	N021015 002 Feb 28, 2000 Feb CPOT
	1% (25MG/2.5GM PACKET)	N021015 001 Feb 28, 2000 Feb CPOT

TESTIM

BX + AUXILIUM PHARMS	1% (50MG/5GM PACKET)	N021454 001 Oct 31, 2002 Feb CPOT
----------------------	----------------------	-----------------------------------

GEL; TRANSDERMAL
TESTOSTERONE
TEVA PHARMS 25MG/2.5GM PACKET N202763 001 Feb 14, 2012 Feb NEWA
50MG/5GM PACKET N202763 002 Feb 14, 2012 Feb NEWA

GEL, METERED; TRANSDERMAL
ANDROGEL
+ ABBOTT LABS 1% (1.25GM/ACTUATION) N021015 003 Sep 26, 2003 Jun CPOT

PELLET; IMPLANTATION
TESTOPEL
+ ACTIENT PHARMS 75MG A080911 001 May CAHN

TESTOSTERONE CYPIONATE
INJECTABLE; INJECTION
TESTOSTERONE CYPIONATE
AO HIKMA FARMACEUTICA 200MG/ML A091244 001 May 01, 2012 Apr NEWA
AO MYLAN INSTITUTIONAL 200MG/ML A040652 001 Dec 11, 2006 Feb CAHN

TESTOSTERONE ENANTHATE
INJECTABLE; INJECTION
TESTOSTERONE ENANTHATE
AO MYLAN INSTITUTIONAL 200MG/ML A040647 001 Oct 05, 2009 Feb CAHN

TETRABENAZINE
TABLET; ORAL
XENAZINE
VALEANT PHARMS 12.5MG N021894 001 Aug 15, 2008 Apr CAHN
+ 25MG N021894 002 Aug 15, 2008 Apr CAHN

TETRAHYDROZOLINE HYDROCHLORIDE
SOLUTION; NASAL
TYZINE
+ FOUGERA PHARMS 0.05% A086576 002 Jan CAHN
0.1% A086576 001 Jan CAHN

SPRAY; NASAL
TYZINE
+ FOUGERA PHARMS 0.1% A086576 003 Jan CAHN

THEOPHYLLINE
CAPSULE, EXTENDED RELEASE; ORAL
THEO-24
UCB INC 100MG A087942 001 Aug 22, 1983 May CTEC
200MG A087943 001 Aug 22, 1983 May CTEC
300MG A087944 001 Aug 22, 1983 May CTEC
+ 400MG A081034 001 Feb 28, 1992 May CRLD

THEOPHYLLINE
@ INWOOD LABS 100MG A040052 001 Feb 14, 1994 May DISC
@ 125MG A040052 002 Feb 14, 1994 May DISC
@ 200MG A040052 003 Feb 14, 1994 May DISC
@ 300MG A040052 004 Feb 14, 1994 May DISC

INJECTABLE; INJECTION
THEOPHYLLINE AND DEXTROSE 5% IN PLASTIC CONTAINER
@ BAXTER HLTHCARE 4MG/ML N018649 007 Jul 26, 1982 Mar DISC
@ 40MG/100ML N018649 001 Jul 26, 1982 Mar DISC
@ 80MG/100ML N018649 002 Jul 26, 1982 Mar DISC
@ 160MG/100ML N018649 003 Jul 26, 1982 Mar DISC
@ 200MG/100ML N018649 004 Jul 26, 1982 Mar DISC

INJECTABLE; INJECTION

THEOPHYLLINE AND DEXTROSE 5% IN PLASTIC CONTAINER

@ BAXTER HLTHCARE	320MG/100ML
@	400MG/100ML

N018649 006	Nov 13, 1985	Mar	DISC
N018649 005	Jul 26, 1982	Mar	DISC

SOLUTION; ORAL

THEOPHYLLINE

AA + SILARX	80MG/15ML
AA TRIS PHARMA INC	80MG/15ML

A091156 001	Apr 13, 2011	Jun	CTEC
A091586 001	Jun 15, 2012	Jun	NEWA

TABLET; ORAL

THEOLAIR

>D> + GRACEWAY	125MG
>D> +	250MG
>A> + MEDICIS	125MG
>A> +	250MG

A086399 001	Jul	CAHN
A086399 002	Jul	CAHN
A086399 001	Jul	CAHN
A086399 002	Jul	CAHN

THIAMINE HYDROCHLORIDEINJECTABLE; INJECTION

THIAMINE HYDROCHLORIDE

AP MYLAN INSTITUTIONAL	100MG/ML
------------------------	----------

A091623 001	Jun 25, 2012	Jun	NEWA
-------------	--------------	-----	------

TIAGABINE HYDROCHLORIDETABLET; ORAL

GABITRIL

@ CEPHALON	6MG
@	8MG
@	10MG

N020646 006	Nov 29, 2005	Jan	DISC
N020646 007	Nov 29, 2005	Jan	DISC
N020646 008	Nov 29, 2005	Jan	DISC

TINIDAZOLETABLET; ORAL

TINDAMAX

AB MISSION PHARMA	250MG
AB +	500MG
TINIDAZOLE	
AB NOVEL LABS INC	250MG
AB	500MG
AB ROXANE	250MG
AB	500MG

N021618 001	May 17, 2004	Apr	CFTG
N021618 002	May 17, 2004	Apr	CFTG
A202044 001	Apr 30, 2012	Apr	NEWA
A202044 002	Apr 30, 2012	Apr	NEWA
A201172 001	Apr 30, 2012	Apr	NEWA
A201172 002	Apr 30, 2012	Apr	NEWA

TINZAPARIN SODIUMINJECTABLE; INJECTION

INNOHEP

@ LEO PHARMA AS	20,000 IU/ML
-----------------	--------------

N020484 001	Jul 14, 2000	Jun	DISC
-------------	--------------	-----	------

TIZANIDINE HYDROCHLORIDECAPSULE; ORAL

TIZANIDINE HYDROCHLORIDE

AB APOTEX INC	EQ 2MG BASE
AB	EQ 4MG BASE
AB	EQ 6MG BASE

A078868 001	Feb 03, 2012	Jan	NEWA
A078868 002	Feb 03, 2012	Jan	NEWA
A078868 003	Feb 03, 2012	Jan	NEWA

ZANAFLEX

AB ACORDA	EQ 2MG BASE
AB	EQ 4MG BASE
AB +	EQ 6MG BASE

N021447 001	Aug 29, 2002	Jan	CFTG
N021447 002	Aug 29, 2002	Jan	CFTG
N021447 003	Aug 29, 2002	Jan	CFTG

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE

AB PROSAM LABS	EQ 2MG BASE
----------------	-------------

A076281 001	Oct 20, 2003	Feb	CMFD
-------------	--------------	-----	------

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE

AB	PROSAM LABS	EQ 4MG BASE	A076281 002 Oct 20, 2003 Feb CMFD
AB	SANDOZ INC	EQ 2MG BASE	A076280 001 Nov 26, 2002 Feb NEWA

TOLBUTAMIDE

TABLET; ORAL

TOLBUTAMIDE

+ MYLAN PHARMS INC	500MG	A086445 001 Mar CRLD
@ WATSON LABS	500MG	A086109 001 Mar DISC
@	500MG	A087318 001 Mar DISC

TOLCAPONE

TABLET; ORAL

TASMAR

+ VALEANT PHARMS LLC	100MG	N020697 001 Jan 29, 1998 May CAHN
@	200MG	N020697 002 Jan 29, 1998 May CAHN

TOPOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

TOPOTECAN HYDROCHLORIDE

>A> AP	INNOPHARMA INC	EQ 4MG BASE/VIAL	A201166 001 Aug 08, 2012 Jul NEWA
--------	----------------	------------------	-----------------------------------

TRAMADOL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TRAMADOL HYDROCHLORIDE

AB2	ACTAVIS	100MG	A091609 001 Jun 27, 2012 Jun NEWA
AB2		200MG	A091609 002 Jun 27, 2012 Jun NEWA
AB2		300MG	A091609 003 Jun 27, 2012 Jun NEWA
AB2	ANCHEN PHARMS	100MG	A200491 001 Jun 27, 2012 Jun NEWA
AB2		200MG	A200491 002 Jun 27, 2012 Jun NEWA
AB2		300MG	A200491 003 Jun 27, 2012 Jun NEWA

TABLET, ORALLY DISINTEGRATING; ORAL

RYBIX ODT

+ SHIONOGI INC	50MG	N021693 001 May 05, 2005 Mar CTNA
----------------	------	-----------------------------------

TRANEXAMIC ACID

INJECTABLE; INJECTION

TRANEXAMIC ACID

AP	APP PHARMS LLC	100MG/ML	A091596 001 Mar 02, 2012 May CAHN
AP	NEXUS PHARMS	100MG/ML	A091596 001 Mar 02, 2012 Feb NEWA

TRANYLCYPROMINE SULFATE

TABLET; ORAL

PARNATE

AB	+ COVIS PHARMA	EQ 10MG BASE	N012342 003 Aug 16, 1985 Jan CAHN
----	----------------	--------------	-----------------------------------

TRETINOIN

CREAM; TOPICAL

AVITA

AB	MYLAN PHARMS INC	0.025%	N020404 003 Jan 14, 1997 Jun CAHN
----	------------------	--------	-----------------------------------

RENOVA

+ VALEANT INTL	0.02%	N021108 001 Aug 31, 2000 Jan CAHN
----------------	-------	-----------------------------------

AB2 +		0.05%	N019963 001 Dec 29, 1995 Jan CAHN
-------	--	-------	-----------------------------------

RETIN-A

AB + VALEANT INTL	0.025%	N019049 001 Sep 16, 1988 Jan CAHN
-------------------	--------	-----------------------------------

CREAM; TOPICAL

RETIN-A

AB1	+	VALEANT INTL	0.05%	N017522 001	Jan	CAHN
AB	+		0.1%	N017340 001	Jan	CAHN
TRETINOIN						
AB		PRECISION DERMAT	0.025%	A075264 001	Dec 24, 1998	May CAHN
	+		0.0375%	A090098 001	Mar 22, 2010	May CAHN
AB1			0.05%	A075265 001	Dec 24, 1998	May CAHN
AB			0.1%	A075213 001	Dec 24, 1998	May CAHN
AB2		SUNEVA MEDCL	0.05%	A076498 001	Sep 15, 2005	May CAHN

GEL; TOPICAL

RETIN-A

AB	+	VALEANT INTL	0.01%	N017955 001	Jan	CAHN
AB	+		0.025%	N017579 002	Jan	CAHN
TRETINOIN						
AB		PRECISION DERMAT	0.01%	A075589 001	Jun 11, 2002	May CAHN
AB			0.025%	A075529 001	Feb 22, 2000	May CAHN

SOLUTION; TOPICAL

RETIN-A

AT	+	VALEANT INTL	0.05%	N016921 001	Jan	CAHN
SWAB; TOPICAL						
RETIN-A						
	@	VALEANT INTL	0.05%	N016921 002	Jan	CAHN

TRIAMCINOLONE ACETONIDE

AEROSOL, METERED; INHALATION

AZMACORT

@ ABBOTT		0.1MG/INH	N018117 001	Apr 23, 1982	May DISC
----------	--	-----------	-------------	--------------	----------

CREAM; TOPICAL

TRIAMCINOLONE ACETONIDE

AT		FOUGERA PHARMS	0.025%	A085692 001	Jan	CAHN
AT			0.1%	A085692 003	Jan	CAHN
AT	+		0.5%	A085692 002	Jan	CAHN

LOTION; TOPICAL

TRIAMCINOLONE ACETONIDE

AT		FOUGERA PHARMS	0.025%	A040467 001	Apr 21, 2003	Jan CAHN
AT			0.1%	A040467 002	Apr 21, 2003	Jan CAHN

OINTMENT; TOPICAL

TRIAMCINOLONE ACETONIDE

AT		FOUGERA PHARMS	0.025%	A085691 001	Jan	CAHN
AT			0.1%	A085691 003	Jan	CAHN
AT	+		0.5%	A085691 002	Jan	CAHN

SPRAY, METERED; NASAL

TRIAMCINOLONE ACETONIDE

AB		TEVA PHARMS	0.055MG/SPRAY	A078104 001	Jul 30, 2009	Mar CAHN
----	--	-------------	---------------	-------------	--------------	----------

TRIMETHOBENZAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

TRIMETHOBENZAMIDE HYDROCHLORIDE

@ HOSPIRA		100MG/ML	A088804 001	Apr 03, 1987	Jun DISC
-----------	--	----------	-------------	--------------	----------

VALACYCLOVIR HYDROCHLORIDE

TABLET; ORAL

VALACYCLOVIR HYDROCHLORIDE

AB		JUBILANT ORGANOSYS	EQ 500MG BASE	A201506 001	Apr 03, 2012	Mar NEWA
AB			EQ 1GM BASE	A201506 002	Apr 03, 2012	Mar NEWA

VALGANCICLOVIR HYDROCHLORIDE

FOR SOLUTION; ORAL
 VALCYTE
 + HOFFMANN LA ROCHE 50MG/ML N022257 001 Aug 28, 2009 May CAHN
 TABLET; ORAL
 VALCYTE
 + HOFFMANN LA ROCHE EQ 450MG BASE N021304 001 Mar 29, 2001 May CAHN

VANCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL
 VANCOCIN HYDROCHLORIDE
 AB VIROPHARMA EQ 125MG BASE N050606 001 Apr 15, 1986 Mar CFTG
 AB + EQ 250MG BASE N050606 002 Apr 15, 1986 Mar CFTG
 VANCOMYCIN HYDROCHLORIDE
 AB AKORN EQ 125MG BASE A065478 001 Apr 09, 2012 Mar NEWA
 AB EQ 250MG BASE A065478 002 Apr 09, 2012 Mar NEWA
 AB APP PHARMS LLC EQ 125MG BASE A065453 001 Jun 18, 2012 Jun NEWA
 AB EQ 250MG BASE A065453 002 Jun 18, 2012 Jun NEWA
 AB STRIDES ARCOLAB LTD EQ 125MG BASE A065490 001 Apr 09, 2012 Mar NEWA
 AB EQ 250MG BASE A065490 002 Apr 09, 2012 Mar NEWA
 AB WATSON LABS EQ 125MG BASE A065510 001 Apr 09, 2012 Mar NEWA
 AB EQ 250MG BASE A065510 002 Apr 09, 2012 Mar NEWA
 INJECTABLE; INJECTION
 VANCOMYCIN HYDROCHLORIDE
 >A> AP SAGENT PHARMS EQ 5GM BASE/VIAL A200837 001 Aug 10, 2012 Jul NEWA
 >A> AP SANDOZ INC EQ 5GM BASE/VIAL A201048 001 Aug 10, 2012 Jul NEWA
 >A> AP EQ 10GM BASE/VIAL A201048 002 Aug 10, 2012 Jul NEWA

VANDETANIB

TABLET; ORAL
 CAPRELSA
 IPR PHARMS INC 100MG N022405 001 Apr 06, 2011 Feb CTNA
 + 300MG N022405 002 Apr 06, 2011 Feb CTNA

VARDENAFIL HYDROCHLORIDE

TABLET; ORAL
 LEVITRA
 AB BAYER HLTHCARE 2.5MG N021400 003 Aug 19, 2003 Apr CFTG
 AB 5MG N021400 001 Aug 19, 2003 Apr CFTG
 AB 10MG N021400 002 Aug 19, 2003 Apr CFTG
 AB + 20MG N021400 004 Aug 19, 2003 Apr CFTG
 VARDENAFIL HYDROCHLORIDE
 AB TEVA PHARMS 2.5MG A091347 001 May 03, 2012 Apr NEWA
 AB 5MG A091347 002 May 03, 2012 Apr NEWA
 AB 10MG A091347 003 May 03, 2012 Apr NEWA
 AB 20MG A091347 004 May 03, 2012 Apr NEWA

VECURONIUM BROMIDE

INJECTABLE; INJECTION
 VECURONIUM BROMIDE
 AP BEDFORD 10MG/VIAL A075549 001 Jun 13, 2000 Jun CRLD
 20MG/VIAL A075549 002 Jun 13, 2000 Jun CRLD
 AP + SUN PHARMA GLOBAL 10MG/VIAL A079001 001 Jun 17, 2009 Jun CRLD
 20MG/VIAL A079001 002 Jun 17, 2009 Jun CRLD

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL
 VENLAFAXINE HYDROCHLORIDE

AB	ANCHEN PHARMS	EQ 37.5MG BASE	A078087 001	Mar 16, 2012	Mar	NEWA
AB		EQ 75MG BASE	A078087 002	Mar 16, 2012	Mar	NEWA
AB		EQ 150MG BASE	A078087 003	Mar 16, 2012	Mar	NEWA

VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL
 CALAN SR

AB	+	PFIZER	120MG	N019152 003	Mar 06, 1991	Feb	CTNA
AB	+		180MG	N019152 002	Dec 15, 1989	Feb	CTNA
AB	+		240MG	N019152 001	Dec 16, 1986	Feb	CTNA

VERAPAMIL HYDROCHLORIDE

AB	APOTEX CORP	120MG	A200878 001	Apr 20, 2012	Apr	NEWA
AB		180MG	A200878 002	Apr 20, 2012	Apr	NEWA
AB		240MG	A200878 003	Apr 20, 2012	Apr	NEWA

VIGABATRIN

FOR SOLUTION; ORAL
 SABRIL

+ LUNDBECK LLC	500MG/PACKET	N022006 001	Aug 21, 2009	Apr	CAHN
----------------	--------------	-------------	--------------	-----	------

TABLET; ORAL
 SABRIL

+ LUNDBECK LLC	500MG	N020427 001	Aug 21, 2009	Apr	CAHN
----------------	-------	-------------	--------------	-----	------

VILAZODONE HYDROCHLORIDE

TABLET; ORAL
 VIIBRYD

+ FOREST LABS INC	10MG	N022567 001	Jan 21, 2011	Apr	CRLD
	40MG	N022567 003	Jan 21, 2011	Apr	CRLD

VISMODEGIB

CAPSULE; ORAL
 ERIVEDGE

+ GENENTECH	150MG	N203388 001	Jan 30, 2012	Jan	NEWA
-------------	-------	-------------	--------------	-----	------

VORICONAZOLE

INJECTABLE; IV (INFUSION)
 VFEND

AP	+	PFIZER	200MG/VIAL	N021267 001	May 24, 2002	May	CFTG
----	---	--------	------------	-------------	--------------	-----	------

VORICONAZOLE

AP	SANDOZ INC	200MG/VIAL	A090862 001	May 30, 2012	May	NEWA
----	------------	------------	-------------	--------------	-----	------

TABLET; ORAL
 VORICONAZOLE

AB	TEVA PHARMS	50MG	A091658 001	Apr 06, 2012	Mar	NEWA
AB		200MG	A091658 002	Apr 06, 2012	Mar	NEWA

ZICONOTIDE

INJECTABLE; INTRATHECAL
 PRIALT

@ JAZZ PHARMS COMMERCIAL	200MCG/2ML (100MCG/ML)	N021060 003	Dec 28, 2004	Mar	CAHN
@ JAZZ PHARMS INTL	200MCG/2ML (100MCG/ML)	N021060 003	Dec 28, 2004	Jun	CAHN

ZICONOTIDE ACETATE

INJECTABLE; INTRATHECAL

PRIALT

+ JAZZ PHARMS COMMERCIAL	100MCG/1ML (100MCG/ML)	N021060 002	Dec 28, 2004	Mar	CAHN
+	500MCG/20ML (25MCG/ML)	N021060 001	Dec 28, 2004	Mar	CAHN
+	500MCG/5ML (100MCG/ML)	N021060 004	Dec 28, 2004	Mar	CAHN
+ JAZZ PHARMS INTL	100MCG/1ML (100MCG/ML)	N021060 002	Dec 28, 2004	Jun	CAHN
+	500MCG/20ML (25MCG/ML)	N021060 001	Dec 28, 2004	Jun	CAHN
+	500MCG/5ML (100MCG/ML)	N021060 004	Dec 28, 2004	Jun	CAHN

ZIDOVUDINE

TABLET; ORAL

ZIDOVUDINE

@ RANBAXY LABS LTD

300MG

A077327 001 Sep 19, 2005 Apr DISC

ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

GEODON

AB + PFIZER	EQ 20MG BASE	N020825 001	Feb 05, 2001	Feb	CFTG
AB	EQ 40MG BASE	N020825 002	Feb 05, 2001	Feb	CFTG
AB	EQ 60MG BASE	N020825 003	Feb 05, 2001	Feb	CFTG
AB	EQ 80MG BASE	N020825 004	Feb 05, 2001	Feb	CFTG

ZIPRASIDONE HYDROCHLORIDE

AB APOTEX CORP	EQ 20MG BASE	A077561 001	Mar 02, 2012	Feb	NEWA
AB	EQ 40MG BASE	A077561 002	Mar 02, 2012	Feb	NEWA
AB	EQ 60MG BASE	A077561 003	Mar 02, 2012	Feb	NEWA
AB	EQ 80MG BASE	A077561 004	Mar 02, 2012	Feb	NEWA
AB DR REDDYS LABS INC	EQ 20MG BASE	A077565 001	Mar 02, 2012	Feb	NEWA
AB	EQ 40MG BASE	A077565 002	Mar 02, 2012	Feb	NEWA
AB	EQ 60MG BASE	A077565 003	Mar 02, 2012	Feb	NEWA
AB	EQ 80MG BASE	A077565 004	Mar 02, 2012	Feb	NEWA
AB LUPIN PHARMS	EQ 20MG BASE	A077560 001	Mar 02, 2012	Feb	NEWA
AB	EQ 40MG BASE	A077560 002	Mar 02, 2012	Feb	NEWA
AB	EQ 60MG BASE	A077560 003	Mar 02, 2012	Feb	NEWA
AB	EQ 80MG BASE	A077560 004	Mar 02, 2012	Feb	NEWA
AB SANDOZ INC	EQ 20MG BASE	A077562 001	Jun 01, 2012	May	NEWA
AB	EQ 40MG BASE	A077562 002	Jun 01, 2012	May	NEWA
AB	EQ 60MG BASE	A077562 003	Jun 01, 2012	May	NEWA
AB	EQ 80MG BASE	A077562 004	Jun 01, 2012	May	NEWA

ZOLPIDEM TARTRATE

TABLET; ORAL

ZOLPIDEM TARTRATE

>A> AB CIPLA LTD	5MG	A077388 001	Jul 30, 2012	Jul	NEWA
>A> AB	10MG	A077388 002	Jul 30, 2012	Jul	NEWA

OTC DRUG PRODUCT LIST - 32ND EDITION

OTC DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

2-1

CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE

TABLET, CHEWABLE; ORAL
 PEPCID COMPLETE
 + MCNEIL CONS 800MG;10MG;165MG N020958 001 Oct 16, 2000 Mar CAHN

CETIRIZINE HYDROCHLORIDE

SYRUP; ORAL
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY
 >D> RANBAXY 5MG/5ML A090183 002 Apr 24, 2008 Jul DISC
 >A> @ RANBAXY LABS LTD 5MG/5ML A090183 002 Apr 24, 2008 Jul DISC
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF
 >D> RANBAXY 5MG/5ML A090183 001 Apr 24, 2008 Jul DISC
 >A> @ RANBAXY LABS LTD 5MG/5ML A090183 001 Apr 24, 2008 Jul DISC

CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL

SWAB; TOPICAL
 CHLORASCRUB MAXI SWABSTICK
 + PROF DSPLS 3.15%;70% (5.1ML) N021524 003 Jun 03, 2005 Mar CAHN
 CHLORASCRUB SWAB
 + PROF DSPLS 3.15%;70% (1ML) N021524 001 Jun 03, 2005 Mar CAHN
 CHLORASCRUB SWABSTICK
 + PROF DSPLS 3.15%;70% (1.6ML) N021524 002 Jun 03, 2005 Mar CAHN

DEXTROMETHORPHAN POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL
 DEXTROMETHORPHAN POLISTIREX
 TRIS PHARMA INC EQ 30MG HBR/5ML A091135 001 May 25, 2012 May NEWA

DIPHENHYDRAMINE HYDROCHLORIDE; IBUPROFEN

CAPSULE; ORAL
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE
 STRIDES ARCOLAB LTD 25MG;EQ 200MG FREE ACID AND POTASSIUM SALT A200888 001 Mar 05, 2012 Feb NEWA

FAMOTIDINE

TABLET, CHEWABLE; ORAL
 PEPCID AC
 @ MCNEIL CONS 10MG N020801 001 Sep 24, 1998 Mar CAHN
 + 20MG N020801 002 Dec 17, 2007 Mar CAHN
 TABLET; ORAL
 PEPCID AC
 MCNEIL CONS 10MG N020325 001 Apr 28, 1995 Mar CAHN
 + 20MG N020325 002 Sep 23, 2003 Mar CAHN
 PEPCID AC (GELTAB)
 MCNEIL CONS 10MG N020902 001 Aug 05, 1999 Mar CAHN

FEXOFENADINE HYDROCHLORIDE

TABLET; ORAL
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY
 SUN PHARM INDs 30MG A091567 002 Feb 06, 2012 Jan NEWA
 WOCKHARDT LTD 30MG A079112 002 Feb 08, 2012 Jan NEWA
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES
 SUN PHARM INDs 30MG A091567 001 Feb 06, 2012 Jan NEWA

TABLET; ORAL

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES		
WOCKHARDT LTD	30MG	A079112 001 Feb 08, 2012 Jan NEWA
FEXOFENADINE HYDROCHLORIDE ALLERGY		
SUN PHARM INDs	60MG	A091567 004 Feb 06, 2012 Jan NEWA
	180MG	A091567 006 Feb 06, 2012 Jan NEWA
WOCKHARDT LTD	60MG	A079112 004 Feb 08, 2012 Jan NEWA
	180MG	A079112 006 Feb 08, 2012 Jan NEWA
FEXOFENADINE HYDROCHLORIDE HIVES		
SUN PHARM INDs	60MG	A091567 003 Feb 06, 2012 Jan NEWA
	180MG	A091567 005 Feb 06, 2012 Jan NEWA
WOCKHARDT LTD	60MG	A079112 003 Feb 08, 2012 Jan NEWA
	180MG	A079112 005 Feb 08, 2012 Jan NEWA

IBUPROFENCAPSULE; ORAL

IBUPROFEN		
ACCUCAPS INDs	EQ 200MG FREE ACID AND POTASSIUM SALT	A077338 001 Jul 10, 2009 Jan CAHN

IBUPROFEN SODIUMTABLET; ORAL

ADVIL		
+ PFIZER CONS HLTHCARE	EQ 200MG BASE	N201803 001 Jun 12, 2012 Jun NEWA

LANSOPRAZOLECAPSULE, DELAYED REL PELLETS; ORAL

LANSOPRAZOLE		
DR REDDYS LABS LTD	15MG	A202194 001 May 18, 2012 Apr NEWA
PERRIGO R AND D	15MG	A202319 001 May 18, 2012 Apr NEWA
WOCKHARDT LTD	15MG	A202727 001 May 18, 2012 Apr NEWA
PREVACID 24 HR		
+ NOVARTIS	15MG	N022327 001 May 18, 2009 Apr CFTG

LEVONORGESTRELTABLET; ORAL

LEVONORGESTREL		
WATSON LABS INC	1.5MG	A200670 001 Jul 12, 2012 Jun NEWA
PLAN B		
+ TEVA BRANDED PHARM	0.75MG	N021045 002 Aug 24, 2006 May CAHN
PLAN B ONE-STEP		
+ DURAMED	1.5MG	N021998 001 Jul 10, 2009 Jun CFTG

LORATADINESYRUP; ORAL

LORATADINE		
>D> RANBAXY	1MG/ML	A076529 001 Aug 20, 2004 Jul DISC
>A> @ RANBAXY LABS LTD	1MG/ML	A076529 001 Aug 20, 2004 Jul DISC

NICOTINEFILM, EXTENDED RELEASE; TRANSDERMAL

NICOTINE		
AVEVA	7MG/24HR	A074612 002 Jul 28, 2003 Apr NEWA
	14MG/24HR	A074612 003 Oct 20, 1997 Apr NEWA

RANITIDINE HYDROCHLORIDE

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

APOTEX INC	EQ 150MG BASE	A200172 001 May 31, 2012 May NEWA
SHASUN CHEMS	EQ 75MG BASE	A201745 001 Feb 29, 2012 Feb NEWA

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

CUMULATIVE SUPPLEMENT NUMBER 07 JULY 2012

NO JULY 2012 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 JULY 2012 ADDITIONS

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ACETYLCYSTEINE - ACETADOTE</u>						
N021539 001	8148356	May 21, 2026	DP			
<u>ACLIDINIUM BROMIDE - TUDORZA PRESSAIR</u>						
N0202450 001				>A> NCE		Jul 23, 2017
<u>ADAPALENE; BENZOYL PEROXIDE - EPIDUO</u>						
N022320 001	8105618	Dec 23, 2022	U-1078			
	8129362	Jul 18, 2027	U-1078			
<u>ALBUTEROL SULFATE - PROAIR HFA</u>						
N021457 001	6446627	Dec 18, 2017	DP			
	8132712	Sep 07, 2028	DP			
<u>ALENDRONATE SODIUM - BINOSTO</u>						
N0202344 001	7488496	Aug 11, 2023	DS DP			
	7964212	Mar 06, 2023	DS DP			
<u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALTURNA</u>						
N022217 001	8168616	Jul 05, 2025	DP			
<u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALTURNA</u>						
N022217 002	8168616	Jul 05, 2025	DP			
<u>ALVIMOPAN - ENTEREG</u>						
N021775 001	8112290	Jul 31, 2030	U-1225			
<u>AMBRISENTAN - LETAIRIS</u>						
N022081 001	RE42462	Jul 29, 2018	DS			
<u>AMBRISENTAN - LETAIRIS</u>						
N022081 002	RE42462	Jul 29, 2018	DS			
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N022314 001	8101599	May 16, 2023	DP			
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N022314 002	8101599	May 16, 2023	DP			
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N022314 003	8101599	May 16, 2023	DP			
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N022314 004	8101599	May 16, 2023	DP			
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N022314 005	8101599	May 16, 2023	DP			
<u>ARFORMOTEROL TARTRATE - BROVANA</u>						
N021912 001	6589508	Apr 03, 2013	U-793			
	6667344	Jun 22, 2021	DP			
	6814953	Jun 22, 2021	U-793			
	8110706	Nov 09, 2021	DP			
<u>ATOVAQUONE; PROGUANIL HYDROCHLORIDE - ATOVAQUONE AND PROGUANIL HYDROCHLORIDE</u>						
A091211 001				PC		Mar 13, 2012
<u>AVANAFIL - STENDRA</u>						
N202276 001	6656935	Sep 13, 2020	DS DP U-155	NCE		Apr 27, 2017

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AVANAFIL - STENDRA</u>						
N202276 002	6656935	Sep 13, 2020	DS DP U-155		NCE	Apr 27, 2017
<u>AVANAFIL - STENDRA</u>						
N202276 003	6656935	Sep 13, 2020	DS DP U-155		NCE	Apr 27, 2017
<u>AXITINIB - INLYTA</u>						
N202324 001	6534524	Jun 30, 2020	DS DP		NCE	Jan 27, 2017
	7141581	Jun 30, 2020	U-1220			
<u>AXITINIB - INLYTA</u>						
N202324 002	6534524	Jun 30, 2020	DS DP		NCE	Jan 27, 2017
	7141581	Jun 30, 2020	U-1220			
<u>AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE - DYMISTA</u>						
N202236 001	8163723	Aug 29, 2023	U-81		NC	May 01, 2015
	8163723	Aug 29, 2023	U-77			
	8163723	Aug 29, 2023	U-707			
	8163723	Aug 29, 2023	U-644			
	8168620	Feb 24, 2026	DP			
<u>AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE - EDARBYCLOR</u>						
N202331 001	5583141	Dec 10, 2013	DS DP U-3			
	5736555	Jun 25, 2012	DS DP U-3			
	7157584	May 22, 2025	DS			
	7572920	Jan 07, 2025	DP U-3			
<u>AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE - EDARBYCLOR</u>						
N202331 002	5583141	Dec 10, 2013	DS DP U-3			
	5736555	Jun 25, 2012	DS DP U-3			
	7157584	May 22, 2025	DS			
	7572920	Jan 07, 2025	DP U-3			
<u>BALSALAZIDE DISODIUM - GIAZO</u>						
N022205 001	6197341	Mar 13, 2018	DP U-1229		NDF	Feb 03, 2015
	7452872	Aug 24, 2026	U-1229			
	7625884	Aug 24, 2026	U-1229			
<u>BECLOMETHASONE DIPROPIONATE - QNASL</u>						
N202813 001	5605674	Feb 25, 2014	DP		NP	Mar 23, 2015
	5683677	Nov 04, 2014	DP			
	5776432	Jul 07, 2015	DP			
	7780038	Jan 24, 2027	DP			
<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N022249 001					NCE	Mar 20, 2013
					ODE	Mar 20, 2015
					PED	Sep 20, 2015
					PED	Sep 20, 2013
<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N022249 002					NCE	Mar 20, 2013
					ODE	Mar 20, 2015
					PED	Sep 20, 2015
					PED	Sep 20, 2013

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BIMATOPROST - LATISSE</u>						
N022369 001	8101161	May 25, 2024		U-1218		
	8101161	May 25, 2024		U-1217		
<u>BIVALIRUDIN - ANGIOMAX</u>						
N020873 001	5196404	Dec 15, 2014	DS DP	U-1232		
	5196404	Dec 15, 2014	DS DP	U-1040		
	5196404*PED	Jun 15, 2015				
<u>BOCEPREVIR - VICTRELIS</u>						
N202258 001	8119602	Mar 17, 2027		U-1233		
	RE43298	Feb 22, 2022	DS DP	U-1128		
<u>BORTEZOMIB - VELCADE</u>						
N021602 001					NR	Jan 23, 2015
<u>BRIMONIDINE TARTRATE; TIMOLOL MALEATE - COMBIGAN</u>						
N021398 001	8133890	Apr 19, 2022		U-1235		
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
N021929 001	8143239	Jan 29, 2023	DP	U-1073		
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
N021929 002	8143239	Jan 29, 2023	DP	U-1073		
<u>BUPIVACAINE - EXPAREL</u>						
N022496 001	6132766	Nov 16, 2013	DP			
	8182835	Sep 18, 2018	DP	U-1246		
<u>BUPIVACAINE - EXPAREL</u>						
N022496 002	6132766	Nov 16, 2013	DP			
	8182835	Sep 18, 2018	DP	U-1246		
<u>CARFILZOMIB - KYPROLIS</u>						
N202714 001	>A> 7232818	Apr 14, 2025	DS DP		>A> NCE	Jul 20, 2017
	>A> 7417042	Jun 07, 2026	DS DP			
	>A> 7491704	Apr 14, 2025		U-1260		
	>A> 7737112	Dec 07, 2027	DP			
	>A> 8129346	Dec 25, 2026		U-1260		
	>A> 8207125	Apr 14, 2025	DS DP			
	>A> 8207126	Apr 14, 2025	DP			
	>A> 8207127	Apr 14, 2025		U-1260		
	>A> 8207297	Apr 14, 2025	DS DP			
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N022012 001	6022562	Oct 17, 2015	DP		Y	
	8101209	Sep 11, 2025	DP			
	8101209*PED	Mar 11, 2026				
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N022012 002	6022562	Oct 17, 2015	DP		Y	
	8101209	Sep 11, 2025	DP			
	8101209*PED	Mar 11, 2026				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N022012 003	6022562	Oct 17, 2015	DP	Y		
	8101209	Sep 11, 2025	DP			
	8101209*PED	Mar 11, 2026				
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N022012 004	6022562	Oct 17, 2015	DP	Y		
	8101209	Sep 11, 2025	DP			
	8101209*PED	Mar 11, 2026				
<u>CHLORHEXIDINE GLUCONATE - CHLORHEXIDINE GLUCONATE</u>						
N021669 001	7717889	Feb 27, 2025	DP U-1022			
<u>CICLESONIDE - ZETONNA</u>						
N202129 001	5482934	Oct 24, 2017	DS DP U-1002		NP	Jan 20, 2015
	5605674	Feb 25, 2014	DP			
	5683677	Nov 04, 2014	DP			
	5775321	Jul 07, 2015	DP			
	6006745	Dec 28, 2016	DP			
	6036942	Apr 30, 2013	DP			
	6120752	May 13, 2018	DP			
	6264923	May 13, 2018	DP			
<u>CICLOPIROX - LOPROX</u>						
N021159 001	>A> 8227490	Sep 16, 2017	U-1256			
<u>CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE - PREPOPIK</u>						
N202535 001				>A> NP		Jul 16, 2015
<u>CLEVIDIPINE BUTYRATE - CLEVIPREX</u>						
N022156 001	5856346	Jan 05, 2021	DS DP U-893			
<u>CLEVIDIPINE BUTYRATE - CLEVIPREX</u>						
N022156 002	5856346	Jan 05, 2021	DS DP U-893			
<u>CLOBETASOL PROPIONATE - CLOBETASOL PROPIONATE</u>						
A078223 001				PC		Jun 30, 2012
<u>CLOBETASOL PROPIONATE - CLOBETASOL PROPIONATE</u>						
A078854 001				PC		Jun 30, 2012
<u>CLOPIDOGREL BISULFATE - CLOPIDOGREL BISULFATE</u>						
A077665 002				PC		Nov 13, 2012
<u>CLOPIDOGREL BISULFATE - CLOPIDOGREL BISULFATE</u>						
A091023 001				PC		Nov 13, 2012
<u>COLCHICINE - COLCRYS</u>						
N022352 001	7964648	Oct 06, 2028	U-1161			
	8097655	Oct 06, 2028	U-1020			
<u>CRIZOTINIB - XALKORI</u>						
N202570 001	>A> 8217057	Nov 06, 2029	DS DP			
<u>CRIZOTINIB - XALKORI</u>						
N202570 002	>A> 8217057	Nov 06, 2029	DS DP			
<u>DAPTOMYCIN - CUBICIN</u>						
N021572 002	8129342	Nov 28, 2020	DS DP			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DARUNAVIR ETHANOLATE - PREZISTA</u>						
N021976 001	5843946*PED	Jun 01, 2016				
	6037157	Jun 26, 2016		U-935		
	6037157	Jun 26, 2016		U-1209		
	6037157*PED	Dec 26, 2016				
	6335460	Aug 25, 2012	DS DP	U-935		
	6335460	Aug 25, 2012	DS DP	U-903		
	6335460	Aug 25, 2012	DS DP	U-744		
	6335460	Aug 25, 2012	DS DP	U-1209		
	6335460*PED	Feb 25, 2013				
	6703403	Jun 26, 2016		U-935		
	6703403	Jun 26, 2016		U-1209		
	6703403*PED	Dec 26, 2016				
	7470506	Jun 23, 2019		U-935		
	7470506	Jun 23, 2019		U-1209		
	7470506*PED	Dec 23, 2019				
<u>DARUNAVIR ETHANOLATE - PREZISTA</u>						
N021976 002	5843946*PED	Jun 01, 2016				
	6037157	Jun 26, 2016		U-935		
	6037157	Jun 26, 2016		U-1209		
	6037157*PED	Dec 26, 2016				
	6335460	Aug 25, 2012	DS DP	U-935		
	6335460	Aug 25, 2012	DS DP	U-903		
	6335460	Aug 25, 2012	DS DP	U-744		
	6335460	Aug 25, 2012	DS DP	U-1209		
	6335460*PED	Feb 25, 2013				
	6703403	Jun 26, 2016		U-935		
	6703403	Jun 26, 2016		U-1209		
	6703403*PED	Dec 26, 2016				
	7470506	Jun 23, 2019		U-935		
	7470506	Jun 23, 2019		U-1209		
	7470506*PED	Dec 23, 2019				
<u>DARUNAVIR ETHANOLATE - PREZISTA</u>						
N021976 003	5843946*PED	Jun 01, 2016				
	6037157	Jun 26, 2016		U-1209		
	6037157	Jun 26, 2016		U-935		
	6037157*PED	Dec 26, 2016				
	6335460	Aug 25, 2012	DS DP	U-935		
	6335460	Aug 25, 2012	DS DP	U-903		
	6335460	Aug 25, 2012	DS DP	U-744		
	6335460	Aug 25, 2012	DS DP	U-1209		
	6335460*PED	Feb 25, 2013				
	6703403	Jun 26, 2016		U-935		
	6703403	Jun 26, 2016		U-1209		
	6703403*PED	Dec 26, 2016				
	7470506	Jun 23, 2019		U-935		
	7470506	Jun 23, 2019		U-1209		
	7470506*PED	Dec 23, 2019				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DARUNAVIR ETHANOLATE - PREZISTA</u>						
N021976 004	5843946*PED	Jun 01, 2016				
	6037157	Jun 26, 2016		U-935		
	6037157	Jun 26, 2016		U-1209		
	6037157*PED	Dec 26, 2016				
	6335460	Aug 25, 2012	DS DP	U-935		
	6335460	Aug 25, 2012	DS DP	U-903		
	6335460	Aug 25, 2012	DS DP	U-744		
	6335460	Aug 25, 2012	DS DP	U-1209		
	6335460*PED	Feb 25, 2013				
	6703403	Jun 26, 2016		U-935		
	6703403	Jun 26, 2016		U-1209		
	6703403*PED	Dec 26, 2016				
	7470506	Jun 23, 2019		U-935		
	7470506	Jun 23, 2019		U-1209		
	7470506*PED	Dec 23, 2019				
<u>DARUNAVIR ETHANOLATE - PREZISTA</u>						
N021976 005	5843946*PED	Jun 01, 2016				
	6037157	Jun 26, 2016		U-935		
	6037157	Jun 26, 2016		U-1209		
	6037157*PED	Dec 26, 2016				
	6335460	Aug 25, 2012	DS DP	U-935		
	6335460	Aug 25, 2012	DS DP	U-903		
	6335460	Aug 25, 2012	DS DP	U-744		
	6335460	Aug 25, 2012	DS DP	U-1209		
	6335460*PED	Feb 25, 2013				
	6703403	Jun 26, 2016		U-935		
	6703403	Jun 26, 2016		U-1209		
	6703403*PED	Dec 26, 2016				
	7470506	Jun 23, 2019		U-935		
	7470506	Jun 23, 2019		U-1209		
	7470506*PED	Dec 23, 2019				
<u>DARUNAVIR ETHANOLATE - PREZISTA</u>						
N202895 001	5843946	Dec 01, 2015	DP	U-1209		
	5843946*PED	Jun 01, 2016				
	6037157	Jun 26, 2016		U-1209		
	6037157*PED	Dec 26, 2016				
	6248775	Aug 13, 2014	DS			
	6248775*PED	Feb 13, 2015				
	6335460	Aug 25, 2012	DS DP	U-1209		
	6335460*PED	Feb 25, 2013				
	6703403	Jun 26, 2016		U-1209		
	6703403*PED	Dec 26, 2016				
	7470506	Jun 23, 2019		U-1209		
	7470506*PED	Dec 23, 2019				
	7700645	Dec 26, 2026	DS DP			
	7700645*PED	Jun 26, 2027				
	RE42889	Oct 19, 2016	DP			
	RE42889*PED	Apr 19, 2017				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DEGARELIX ACETATE - FIRMAGON</u>						
N022201 001	5925730	May 18, 2021	DS DP U-943			
<u>DEGARELIX ACETATE - FIRMAGON</u>						
N022201 002	5925730	May 18, 2021	DS DP U-943			
<u>DESLORATADINE; PSEUDOEPHEDRINE SULFATE - CLARINEX-D 12 HOUR</u>						
N021313 001	8187630	Dec 19, 2020	DP U-1017			
<u>DEXAMETHASONE - OZURDEX</u>						
N022315 001	8088407	Oct 20, 2020	DP U-1205			
<u>DEXLANSOPRAZOLE - DEXILANT</u>						
N022287 001	8105626	Sep 27, 2026	DP			
	8105626*PED	Mar 27, 2027				
	8173158	Mar 17, 2030	U-951			
	8173158	Mar 17, 2030	U-950			
	8173158	Mar 17, 2030	U-949			
	8173158*PED	Sep 17, 2030				
<u>DEXLANSOPRAZOLE - DEXILANT</u>						
N022287 002	8105626	Sep 27, 2026	DP			
	8105626*PED	Mar 27, 2027				
	8173158	Mar 17, 2030	U-951			
	8173158	Mar 17, 2030	U-950			
	8173158	Mar 17, 2030	U-949			
	8173158*PED	Sep 17, 2030				
<u>DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE - NUEDEXTA</u>						
N021879 001	>A> 8227484	Jul 17, 2023	U-1093			
<u>DICLOFENAC POTASSIUM - CAMBIA</u>						
N022165 001	8097651	Jun 16, 2026	DS DP U-436			
<u>DICLOFENAC POTASSIUM - ZIPSOR</u>						
N022202 001	8110606	Feb 24, 2029	U-980			
<u>DICLOFENAC SODIUM - PENNSAID</u>						
N020947 001	8217078	Jul 10, 2029	U-1248			
<u>DIENOGEST; ESTRADIOL VALERATE - NATAZIA</u>						
N022252 001	8153616	Jan 30, 2028	U-1240	I-648	Mar 14, 2015	
<u>DIFLUPREDNATE - DUREZOL</u>						
N022212 001				I-653 ODE	Jun 13, 2015 Jun 13, 2019	
<u>DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE - COSOPT PF</u>						
N202667 001				NP	Feb 01, 2015	
<u>DOXYCYCLINE - ORACEA</u>						
N050805 001	8206740	Dec 24, 2025	DP U-925			
<u>DRONEDARONE HYDROCHLORIDE - MULTAQ</u>						
N022425 001	>A> 5223510	Jul 26, 2012	DS DP U-992			
	>A> 5223510	Jul 26, 2012	DS DP U-1261			
<u>DROSPIRENONE; ESTRADIOL - ANGELIQ</u>						
N021355 001				NS	Mar 01, 2015	

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u>						
N021427 001	5023269	Jun 11, 2013	DS DP U-882		I-632	Nov 04, 2013
	5023269	Jun 11, 2013	DS DP U-839		I-617	Nov 19, 2012
	5023269	Jun 11, 2013	DS DP U-799		PED	May 04, 2014
	5023269	Jun 11, 2013	DS DP U-797		PED	May 19, 2013
	5023269	Jun 11, 2013	DS DP U-796			
	5023269	Jun 11, 2013	DS DP U-795			
	5023269	Jun 11, 2013	DS DP U-605			
	5023269	Jun 11, 2013	DS DP U-398			
	5023269	Jun 11, 2013	DS DP U-1094			
	5023269*PED	Dec 11, 2013				
	5508276	Jul 18, 2014	DP			
	5508276*PED	Jan 18, 2015				
	6596756	Sep 10, 2019	U-882			
	6596756*PED	Mar 10, 2020				
<u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u>						
N021427 002	5023269	Jun 11, 2013	DS DP U-882		I-632	Nov 04, 2013
	5023269	Jun 11, 2013	DS DP U-839		I-617	Nov 19, 2012
	5023269	Jun 11, 2013	DS DP U-799		PED	May 04, 2014
	5023269	Jun 11, 2013	DS DP U-797		PED	May 19, 2013
	5023269	Jun 11, 2013	DS DP U-796			
	5023269	Jun 11, 2013	DS DP U-795			
	5023269	Jun 11, 2013	DS DP U-605			
	5023269	Jun 11, 2013	DS DP U-398			
	5023269	Jun 11, 2013	DS DP U-1094			
	5023269*PED	Dec 11, 2013				
	5508276	Jul 18, 2014	DP			
	5508276*PED	Jan 18, 2015				
	6596756	Sep 10, 2019	U-882			
	6596756*PED	Mar 10, 2020				
<u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u>						
N021427 004	5023269	Jun 11, 2013	DS DP U-882		I-632	Nov 04, 2013
	5023269	Jun 11, 2013	DS DP U-839		I-617	Nov 19, 2012
	5023269	Jun 11, 2013	DS DP U-799		PED	May 04, 2014
	5023269	Jun 11, 2013	DS DP U-797		PED	May 19, 2013
	5023269	Jun 11, 2013	DS DP U-796			
	5023269	Jun 11, 2013	DS DP U-795			
	5023269	Jun 11, 2013	DS DP U-605			
	5023269	Jun 11, 2013	DS DP U-398			
	5023269	Jun 11, 2013	DS DP U-1094			
	5023269*PED	Dec 11, 2013				
	5508276	Jul 18, 2014	DP			
	5508276*PED	Jan 18, 2015				
	6596756	Sep 10, 2019	U-882			
	6596756*PED	Mar 10, 2020				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - ATRIPLA</u>						
N021937 001	5663169	Sep 02, 2014	U-750			
	5663169	Sep 02, 2014	U-1170			
	5811423	Aug 07, 2012	U-750			
	5811423	Aug 07, 2012	U-1170			
	5922695	Jul 25, 2017	DS	U-750		
	5922695	Jul 25, 2017	DS	U-1170		
	5935946	Jul 25, 2017	DS DP	U-750		
	5935946	Jul 25, 2017	DS DP	U-1170		
	5977089	Jul 25, 2017	DS DP	U-750		
	5977089	Jul 25, 2017	DS DP	U-1170		
	6043230	Jul 25, 2017		U-750		
	6043230	Jul 25, 2017		U-1170		
	6642245	Nov 04, 2020		U-750		
	6642245	Nov 04, 2020		U-1170		
<u>EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR DISOPROXIL FUMARATE - COMPLERA</u>						
N202123 001	8080551	Apr 11, 2023	DS DP		NCE	May 20, 2016
	8101629	Aug 09, 2022	DP			
<u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u>						
N021752 001	>A> 5922695	Jul 25, 2017	DS	U-1259		
	>A> 5922695	Jul 25, 2017	DS	U-248		
	>A> 5922695	Jul 25, 2017	DS	U-541		
	>A> 5922695	Jul 25, 2017	DS	U-1170		
	>A> 5935946	Jul 25, 2017	DS DP	U-541		
	>A> 5935946	Jul 25, 2017	DS DP	U-1170		
	>A> 5935946	Jul 25, 2017	DS DP	U-1259		
	>A> 5935946	Jul 25, 2017	DS DP	U-248		
	>A> 5977089	Jul 25, 2017	DS DP	U-1259		
	>A> 5977089	Jul 25, 2017	DS DP	U-248		
	>A> 5977089	Jul 25, 2017	DS DP	U-541		
	>A> 5977089	Jul 25, 2017	DS DP	U-1170		
	>A> 6043230	Jul 25, 2017	DP	U-541		
	>A> 6043230	Jul 25, 2017	DP	U-1170		
	>A> 6043230	Jul 25, 2017	DP	U-1259		
	>A> 6043230	Jul 25, 2017	DP	U-248		
<u>EPROSARTAN MESYLATE - EPROSARTAN MESYLATE</u>						
A202012 001					PC	Jun 17, 2012
<u>EPROSARTAN MESYLATE - EPROSARTAN MESYLATE</u>						
A202012 002					PC	Jun 17, 2012
<u>ERIBULIN MESYLATE - HALAVEN</u>						
N201532 001	8097648	Jan 22, 2021		U-1096		
<u>ESCITALOPRAM OXALATE - ESCITALOPRAM OXALATE</u>						
A076765 001					PC	Sep 10, 2012
<u>ESCITALOPRAM OXALATE - ESCITALOPRAM OXALATE</u>						
A076765 002					PC	Sep 10, 2012
<u>ESCITALOPRAM OXALATE - ESCITALOPRAM OXALATE</u>						
A076765 003					PC	Sep 10, 2012

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>						
N021957 001	5877192	May 27, 2014	U-773			
	5877192	May 27, 2014	U-729			
	5877192	May 27, 2014	U-1207			
	5877192*PED	Nov 27, 2014				
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>						
N021957 002	5877192	May 27, 2014	U-773			
	5877192	May 27, 2014	U-729			
	5877192	May 27, 2014	U-1207			
	5877192*PED	Nov 27, 2014				
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>						
N021957 003	>A> 5690960	Nov 25, 2014	DP U-1207			
	>A> 5690960*PED	May 25, 2015				
	>A> 5714504	Feb 03, 2015	DP U-1207			
	>A> 5714504*PED	Aug 03, 2015				
	>A> 5877192	May 27, 2014	U-1207			
	>A> 5877192*PED	Nov 27, 2014				
	>A> 5900424	May 04, 2016	DS U-1207			
	>A> 5900424*PED	Nov 04, 2016				
	>A> 6369085	May 25, 2018	DS DP U-1207			
	>A> 6369085*PED	Nov 25, 2018				
	>A> 6428810	Nov 03, 2019	DP U-1207			
	>A> 6428810*PED	May 03, 2020				
	>A> 6875872	May 27, 2014	DS			
	>A> 6875872*PED	Nov 27, 2014				
	>A> 7411070	May 25, 2018	DS			
	>A> 7411070*PED	Nov 25, 2018				
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>						
N021957 004	>A> 5690960	Nov 25, 2014	DP U-1207			
	>A> 5690960*PED	May 25, 2015				
	>A> 5714504	Feb 03, 2015	DP U-1207			
	>A> 5714504*PED	Aug 03, 2015				
	>A> 5877192	May 27, 2014	U-1207			
	>A> 5877192*PED	Nov 27, 2014				
	>A> 5900424	May 04, 2016	DS U-1207			
	>A> 5900424*PED	Nov 04, 2016				
	>A> 6369085	May 25, 2018	DS DP U-1207			
	>A> 6369085*PED	Nov 25, 2018				
	>A> 6428810	Nov 03, 2019	DP U-1207			
	>A> 6428810*PED	May 03, 2020				
	>A> 6875872	May 27, 2014	DS			
	>A> 6875872*PED	Nov 27, 2014				
	>A> 7411070	May 25, 2018	DS			
	>A> 7411070*PED	Nov 25, 2018				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ESZOPICLONE - LUNESTA</u>						
N021476 001	6319926	Jan 16, 2012		U-620		
	6319926*PED	Jul 16, 2012				
	6444673	Feb 14, 2014	DS DP			
	6444673*PED	Aug 14, 2014				
	6864257	Aug 30, 2012		U-629		
	6864257*PED	Mar 02, 2013				
	7381724	Jan 16, 2012	DS DP	U-629		
	7381724*PED	Jul 16, 2012				
<u>ESZOPICLONE - LUNESTA</u>						
N021476 002	6319926	Jan 16, 2012		U-620		
	6319926*PED	Jul 16, 2012				
	6444673	Feb 14, 2014	DS DP			
	6444673*PED	Aug 14, 2014				
	6864257	Aug 30, 2012		U-629		
	6864257*PED	Mar 02, 2013				
	7381724	Jan 16, 2012	DS DP	U-629		
	7381724*PED	Jul 16, 2012				
<u>ESZOPICLONE - LUNESTA</u>						
N021476 003	6319926	Jan 16, 2012		U-620		
	6319926*PED	Jul 16, 2012				
	6444673	Feb 14, 2014	DS DP			
	6444673*PED	Aug 14, 2014				
	6864257	Aug 30, 2012		U-629		
	6864257*PED	Mar 02, 2013				
	7381724	Jan 16, 2012	DS DP	U-629		
	7381724*PED	Jul 16, 2012				
<u>ETONOGESTREL - NEXPLANON</u>						
N021529 002					NP	May 31, 2014
<u>ETRAVIRINE - INTELENCE</u>						
N022187 001	6878717	Nov 05, 2019		U-256		
	6878717	Nov 05, 2019		U-1237		
	6878717	Nov 05, 2019		U-1016		
	7037917	Dec 13, 2020	DS DP	U-256		
	7037917	Dec 13, 2020	DS DP	U-1237		
	7037917	Dec 13, 2020	DS DP	U-1016		
<u>ETRAVIRINE - INTELENCE</u>						
N022187 002	6878717	Nov 05, 2019		U-256		
	6878717	Nov 05, 2019		U-1237		
	6878717	Nov 05, 2019		U-1016		
	7037917	Dec 13, 2020	DS DP	U-256		
	7037917	Dec 13, 2020	DS DP	U-1237		
	7037917	Dec 13, 2020	DS DP	U-1016		

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ETRAVIRINE - INTELENCE</u>						
N022187 003	6878717	Nov 05, 2019	U-256		NPP	Mar 26, 2015
	6878717	Nov 05, 2019	U-1237		NCE	Jan 18, 2013
	6878717	Nov 05, 2019	U-1016			
	7037917	Dec 13, 2020	DS DP U-1237			
	7887845	Mar 25, 2019	DP			
	8003789	Nov 01, 2019	DS DP			
<u>EVEROLIMUS - AFINITOR</u>						
N022334 001	5665772	Sep 09, 2019	DS DP	>A>	I-655	Jul 20, 2015
	5665772*PED	Mar 09, 2020			I-650	Apr 26, 2015
	6004973	Jul 12, 2016	DP		I-638	May 05, 2014
	6004973*PED	Jan 12, 2017			I-630	Oct 29, 2013
	7297703	Dec 06, 2019	DP		NCE	Mar 30, 2014
	7297703*PED	Jun 06, 2020			ODE	Apr 26, 2019
					ODE	May 05, 2018
					ODE	Oct 29, 2017
					PED	Oct 26, 2019
					PED	Nov 05, 2018
					PED	Apr 29, 2018
					PED	Oct 26, 2015
					PED	Nov 05, 2014
					PED	Sep 30, 2014
					PED	Apr 29, 2014
<u>EVEROLIMUS - AFINITOR</u>						
N022334 002	5665772	Sep 09, 2019	DS DP	>A>	I-655	Jul 20, 2015
	5665772*PED	Mar 09, 2020			I-650	Apr 26, 2015
	6004973	Jul 12, 2016	DP		I-638	May 05, 2014
	6004973*PED	Jan 12, 2017			I-630	Oct 29, 2013
	7297703	Dec 06, 2019	DP		NCE	Mar 30, 2014
	7297703*PED	Jun 06, 2020			ODE	Apr 26, 2019
					ODE	May 05, 2018
					ODE	Oct 29, 2017
					PED	Oct 26, 2019
					PED	Nov 05, 2018
					PED	Apr 29, 2018
					PED	Oct 26, 2015
					PED	Nov 05, 2014
					PED	Sep 30, 2014
					PED	Apr 29, 2014
<u>EVEROLIMUS - AFINITOR</u>						
N022334 003	5665772	Sep 09, 2019	DS DP	>A>	I-655	Jul 20, 2015
	5665772*PED	Mar 09, 2020			I-650	Apr 26, 2015
	6004973	Jul 12, 2016	DP		I-638	May 05, 2014
	6004973*PED	Jan 12, 2017			I-630	Oct 29, 2013
	7297703	Dec 06, 2019	DP		NCE	Mar 30, 2014
	7297703*PED	Jun 06, 2020			ODE	Apr 26, 2019
					ODE	May 05, 2018
					ODE	Oct 29, 2017
					PED	Oct 26, 2019
					PED	Nov 05, 2018
					PED	Apr 29, 2018
					PED	Oct 26, 2015
					PED	Nov 05, 2014
					PED	Sep 30, 2014
					PED	Apr 29, 2014

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
EVEROLIMUS - AFINITOR							
N022334 004	5665772	Sep	09, 2019	DS DP	>A>	I-655	Jul 20, 2015
	5665772*PED	Mar	09, 2020			I-650	Apr 26, 2015
	6004973	Jul	12, 2016	DP		I-638	May 05, 2014
	6004973*PED	Jan	12, 2017			I-630	Oct 29, 2013
	7297703	Dec	06, 2019	DP		NCE	Mar 30, 2014
	7297703*PED	Jun	06, 2020			ODE	Apr 26, 2019
						ODE	May 05, 2018
						ODE	Oct 29, 2017
						PED	Oct 26, 2019
						PED	Nov 05, 2018
						PED	Apr 29, 2018
						PED	Oct 26, 2015
						PED	Nov 05, 2014
						PED	Sep 30, 2014
						PED	Apr 29, 2014
EVEROLIMUS - ZORTRESS							
N021560 001	5665772	Sep	09, 2019	DS DP U-1049		NP	Apr 20, 2013
	5665772*PED	Mar	09, 2020			NCE	Mar 30, 2014
	6004973	Jul	12, 2016	DP U-1049		PED	Sep 30, 2014
	6004973*PED	Jan	12, 2017			PED	Oct 20, 2013
	6239124	Aug	11, 2017	U-1049			
	6239124*PED	Feb	11, 2018				
	6440990	Sep	24, 2013	DP U-1049			
	6440990*PED	Mar	24, 2014				
	6455518	Jul	29, 2017	U-1049			
	6455518*PED	Jan	29, 2018				
EVEROLIMUS - ZORTRESS							
N021560 002	5665772	Sep	09, 2019	DS DP U-1049		NP	Apr 20, 2013
	5665772*PED	Mar	09, 2020			NCE	Mar 30, 2014
	6004973	Jul	12, 2016	DP U-1049		PED	Sep 30, 2014
	6004973*PED	Jan	12, 2017			PED	Oct 20, 2013
	6239124	Aug	11, 2017	U-1049			
	6239124*PED	Feb	11, 2018				
	6440990	Sep	24, 2013	DP U-1049			
	6440990*PED	Mar	24, 2014				
	6455518	Jul	29, 2017	U-1049			
	6455518*PED	Jan	29, 2018				
EVEROLIMUS - ZORTRESS							
N021560 003	5665772	Sep	09, 2019	DS DP U-1049		NP	Apr 20, 2013
	5665772*PED	Mar	09, 2020			NCE	Mar 30, 2014
	6004973	Jul	12, 2016	DP U-1049		PED	Sep 30, 2014
	6004973*PED	Jan	12, 2017			PED	Oct 20, 2013
	6239124	Aug	11, 2017	U-1049			
	6239124*PED	Feb	11, 2018				
	6440990	Sep	24, 2013	DP U-1049			
	6440990*PED	Mar	24, 2014				
	6455518	Jul	29, 2017	U-1049			
	6455518*PED	Jan	29, 2018				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>EXENATIDE SYNTHETIC - BYDUREON</u>						
N022200 001	5424286	Dec 01, 2016	U-1108		NP	Jan 27, 2015
	6479065	Aug 10, 2020	DP			
	6495164	May 25, 2020	DP			
	6667061	May 25, 2020	DP			
	6824822	Oct 09, 2022	DP			
	6858576	Jan 06, 2017	U-656			
	6872700	Jan 14, 2020	U-654			
	6956026	Jan 07, 2018	U-687			
	7223440	Aug 31, 2021	DP			
	7456254	Jun 30, 2025	DP U-1223			
	7563871	Apr 15, 2024	DP			
	7612176	Apr 13, 2025	DP U-1223			
	7741269	Jan 07, 2018	U-1224			
<u>EXENATIDE SYNTHETIC - BYETTA</u>						
N021773 001				M-113	Oct 19, 2014	
				M-111	Oct 19, 2014	
<u>EXENATIDE SYNTHETIC - BYETTA</u>						
N021773 002				M-113	Oct 19, 2014	
				M-111	Oct 19, 2014	
<u>EZETIMIBE - ZETIA</u>						
N021445 001				M-109	Jan 24, 2015	
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>						
N021687 001				M-109	Jan 24, 2015	
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>						
N021687 002				M-109	Jan 24, 2015	
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>						
N021687 003				M-109	Jan 24, 2015	
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>						
N021687 004				M-109	Jan 24, 2015	
<u>FAMCICLOVIR - FAMVIR</u>						
N020363 001				M-112	Feb 09, 2015	
<u>FAMCICLOVIR - FAMVIR</u>						
N020363 002				M-112	Feb 09, 2015	
<u>FAMCICLOVIR - FAMVIR</u>						
N020363 003				M-112	Feb 09, 2015	
<u>FENOFIBRATE - FENOGLIDE</u>						
N022118 001	8124125	Oct 01, 2024	DP U-1234			
<u>FENOFIBRATE - FENOGLIDE</u>						
N022118 002	8124125	Oct 01, 2024	DP U-1234			
<u>FENTANYL - SUBSYS</u>						
N202788 001				NP	Jan 04, 2015	
<u>FENTANYL - SUBSYS</u>						
N202788 002				NP	Jan 04, 2015	
<u>FENTANYL - SUBSYS</u>						
N202788 003				NP	Jan 04, 2015	

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FENTANYL - SUBSYS</u>						
	N202788 004				NP	Jan 04, 2015
<u>FENTANYL - SUBSYS</u>						
	N202788 005				NP	Jan 04, 2015
<u>FENTANYL CITRATE - FENTORA</u>						
N021947 001	8092832	Dec 30, 2024	DP			
<u>FENTANYL CITRATE - FENTORA</u>						
N021947 002	8092832	Dec 30, 2024	DP			
	8119158	Dec 30, 2024	DP			
<u>FENTANYL CITRATE - FENTORA</u>						
N021947 003	8092832	Dec 30, 2024	DP			
	8119158	Dec 30, 2024	DP			
<u>FENTANYL CITRATE - FENTORA</u>						
N021947 004	8092832	Dec 30, 2024	DP			
	8119158	Dec 30, 2024	DP			
<u>FENTANYL CITRATE - FENTORA</u>						
N021947 005	8092832	Dec 30, 2024	DP			
	8119158	Dec 30, 2024	DP			
<u>FENTANYL CITRATE - LAZANDA</u>						
N022569 001	>A> 8216604	Oct 03, 2024		U-767		
<u>FENTANYL CITRATE - LAZANDA</u>						
N022569 002	>A> 8216604	Oct 03, 2024		U-767		
<u>FENTANYL CITRATE - ONSOLIS</u>						
N022266 001	7579019	Jan 22, 2020		U-767		
<u>FENTANYL CITRATE - ONSOLIS</u>						
N022266 002	7579019	Jan 22, 2020		U-767		
<u>FENTANYL CITRATE - ONSOLIS</u>						
N022266 003	7579019	Jan 22, 2020		U-767		
<u>FENTANYL CITRATE - ONSOLIS</u>						
N022266 004	7579019	Jan 22, 2020		U-767		
<u>FENTANYL CITRATE - ONSOLIS</u>						
N022266 005	7579019	Jan 22, 2020		U-767		
<u>FESOTERODINE FUMARATE - TOVIAZ</u>						
N022030 001	8088398	Jun 07, 2027	DP U-913			
<u>FESOTERODINE FUMARATE - TOVIAZ</u>						
N022030 002	8088398	Jun 07, 2027	DP U-913			
<u>FINGOLIMOD - GILENYA</u>						
N022527 001	6004565	Sep 23, 2017		U-1086		
<u>FLORBETAPIR F-18 - AMYVID</u>						
N202008 001	7687052	Apr 30, 2027	DS DP		NCE	Apr 06, 2017
<u>FLORBETAPIR F-18 - AMYVID</u>						
N202008 002	7687052	Apr 30, 2027	DS DP		NCE	Apr 06, 2017

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLORBETAPIR F-18 - AMYVID</u>						
N020208 003	7687052	Apr 30, 2027	DS DP		NCE	Apr 06, 2017
<u>FLUNISOLIDE - AEROSPIN HFA</u>						
N021247 001	5776433	Jul 07, 2015	DP			
	5980867	Jul 06, 2018	DP			
<u>FLUOCINONIDE - VANOS</u>						
N021758 001	>A> 8232264	Mar 09, 2023	DP			
<u>FOSAMPRENAVIR CALCIUM - LEXIVA</u>						
N021548 001					NPP PED	Apr 27, 2015 Oct 27, 2015
<u>FOSAMPRENAVIR CALCIUM - LEXIVA</u>						
N022116 001					NPP PED	Apr 27, 2015 Oct 27, 2015
<u>FOSAPREPITANT DIMEGLUMINE - EMEND</u>						
N022023 002	5512570	Mar 04, 2014	U-850			
	5538982	Jul 23, 2013	U-850			
	5716942	Feb 10, 2015	U-850			
	7214692	Sep 18, 2012	U-850			
<u>FOSPROPOFOL DISODIUM - LUSEDRA</u>						
N022244 001	6204257	Jul 01, 2022	DS DP U-945			
<u>GABAPENTIN - GRALISE</u>						
N022544 001	8192756	Oct 25, 2022	DP U-1114			
<u>GABAPENTIN - GRALISE</u>						
N022544 002	8192756	Oct 25, 2022	DP U-1114			
<u>GABAPENTIN ENACARBIL - HORIZANT</u>						
N022399 001	8048917	Nov 06, 2022	DS DP U-1247		I-652	Jun 06, 2015
	8114909	Apr 11, 2026	U-1231			
<u>GABAPENTIN ENACARBIL - HORIZANT</u>						
N022399 002	8048917	Nov 06, 2022	DS DP U-1247		I-652	Jun 06, 2015
	8114909	Apr 11, 2026	U-1231			
<u>GADOBENATE DIMEGLUMINE - MULTIHANCE</u>						
N021357 001					I-654	Jul 06, 2015
<u>GADOBENATE DIMEGLUMINE - MULTIHANCE</u>						
N021357 002					I-654	Jul 06, 2015
<u>GADOBENATE DIMEGLUMINE - MULTIHANCE</u>						
N021357 003					I-654	Jul 06, 2015
<u>GADOBENATE DIMEGLUMINE - MULTIHANCE</u>						
N021357 004					I-654	Jul 06, 2015
<u>GADOBENATE DIMEGLUMINE - MULTIHANCE MULTIPACK</u>						
N021358 001					I-654	Jul 06, 2015
<u>GADOBENATE DIMEGLUMINE - MULTIHANCE MULTIPACK</u>						
N021358 002					I-654	Jul 06, 2015
<u>GADOFOSVESET TRISODIUM - ABLAVAR</u>						
N021711 001	6676929	May 04, 2020	DP			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>GADOFOSVESET TRISODIUM - ABLAVAR</u>						
N021711 002	6676929	May 04, 2020	DP			
<u>GADOXETATE DISODIUM - EOVIST</u>						
N022090 001	5798092	Aug 25, 2015	DS DP			
	6039931	Nov 13, 2021	U-1239			
<u>GLYCOPYRROLATE - CUVPOSA</u>						
N022571 001					ODE	Jul 28, 2017
<u>HYDROCHLOROTHIAZIDE; IRBESARTAN - IRBESARTAN AND HYDROCHLOROTHIAZIDE</u>						
A077369 001					PC	Sep 26, 2012
<u>HYDROCHLOROTHIAZIDE; IRBESARTAN - IRBESARTAN AND HYDROCHLOROTHIAZIDE</u>						
A077369 002					PC	Sep 26, 2012
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N202057 001	>A> 8188146	Jan 27, 2020	DS DP			
<u>IMATINIB MESYLATE - GLEEVEC</u>						
N021588 001					ODE	Dec 19, 2015
<u>IMATINIB MESYLATE - GLEEVEC</u>						
N021588 002					ODE	Dec 19, 2015
<u>IMIQUIMOD - ZYCLARA</u>						
N022483 001	>A> 8236816	Dec 11, 2029	U-68			
<u>IMIQUIMOD - ZYCLARA</u>						
N022483 002	>A> 8222270	Dec 11, 2029	U-68			
<u>INGENOL MEBUTATE - PICATO</u>						
N202833 001	6432452	Aug 19, 2018	DS U-68		NCE	Jan 23, 2017
	6844013	Aug 19, 2018	DS DP U-1221			
	7410656	Aug 19, 2018	DS U-1222			
<u>INGENOL MEBUTATE - PICATO</u>						
N202833 002	6432452	Aug 19, 2018	DS U-68		NCE	Jan 23, 2017
	6844013	Aug 19, 2018	DS DP U-1221			
	7410656	Aug 19, 2018	DS U-1222			
<u>INSULIN DETEMIR RECOMBINANT - LEVEMIR</u>						
N021536 001	>A> 5750497	Jun 16, 2019	DS DP U-668		M-117 M-115	May 18, 2015 Apr 06, 2015
<u>IOFLUPANE I-123 - DATSCAN</u>						
N022454 001	5310912	Feb 25, 2013	DS			
<u>IRBESARTAN - IRBESARTAN</u>						
A077159 001					PC	Sep 26, 2012
<u>IRBESARTAN - IRBESARTAN</u>						
A077159 002					PC	Sep 26, 2012
<u>IRBESARTAN - IRBESARTAN</u>						
A077159 003					PC	Sep 26, 2012
<u>ISOTRETINOIN - ABSORICA</u>						
N021951 001	7435427	Sep 21, 2021	DP		NP	May 25, 2015
<u>ISOTRETINOIN - ABSORICA</u>						
N021951 002	7435427	Sep 21, 2021	DP		NP	May 25, 2015

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ISOTRETINOIN - ABSORICA</u>						
N021951 003	7435427	Sep 21, 2021	DP		NP	May 25, 2015
<u>ISOTRETINOIN - ABSORICA</u>						
N021951 004	7435427	Sep 21, 2021	DP		NP	May 25, 2015
<u>IVACAFTOR - KALYDECO</u>						
N203188 001	7495103	May 20, 2027	DS DP		NCE ODE	Jan 31, 2017 Jan 31, 2019
<u>IVERMECTIN - SKLICE</u>						
N202736 001	6103248	May 22, 2018	DP		NP	Feb 07, 2015
<u>IXABEPILONE - IXEMPRA KIT</u>						
N022065 001	RE41911	Sep 28, 2020	DS DP U-961			
	RE41911*PED	Mar 28, 2021				
<u>IXABEPILONE - IXEMPRA KIT</u>						
N022065 002	RE41911	Sep 28, 2020	DS DP U-961			
	RE41911*PED	Mar 28, 2021				
<u>KETOCONAZOLE - EXTINA</u>						
N021738 001	8026238	Oct 19, 2018	DP U-1213			
<u>KETOROLAC TROMETHAMINE - ACULAR LS</u>						
N021528 001	>A> 8008338*PED	Nov 24, 2027				
	>A> 8207215	May 28, 2024	U-1251			
	>A> 8207215*PED	Nov 28, 2024				
<u>LAMIVUDINE; ZIDOVUDINE - LAMIVUDINE AND ZIDOVUDINE</u>						
A079081 001					PC	May 15, 2012
<u>LENALIDOMIDE - REVLIMID</u>						
N021880 001	5635517	Oct 04, 2019	DS	U-1211		
	6045501	Aug 28, 2018		U-1210		
	6281230	Jul 24, 2016		U-1212		
	6315720	Oct 23, 2020		U-1210		
	6555554	Jul 24, 2016	DP	U-1211		
	6561976	Aug 28, 2018		U-1210		
	6561977	Oct 23, 2020		U-1210		
	6755784	Oct 23, 2020		U-1210		
	6908432	Aug 28, 2018		U-1210		
	7189740	Apr 11, 2023		U-1215		
	7465800	Apr 27, 2027	DS DP			
	7968569	Oct 07, 2023		U-1216		
	8204763	Aug 28, 2018		U-1249		

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LENALIDOMIDE - REVLIMID</u>						
N021880 002	5635517	Oct 04, 2019	DS	U-1211		
	6045501	Aug 28, 2018		U-1210		
	6281230	Jul 24, 2016		U-1212		
	6315720	Oct 23, 2020		U-1210		
	6555554	Jul 24, 2016	DP	U-1211		
	6561976	Aug 28, 2018		U-1210		
	6561977	Oct 23, 2020		U-1210		
	6755784	Oct 23, 2020		U-1210		
	6908432	Aug 28, 2018		U-1210		
	7189740	Apr 11, 2023		U-1215		
	7465800	Apr 27, 2027	DS DP			
	7968569	Oct 07, 2023		U-1216		
	8204763	Aug 28, 2018		U-1249		
<u>LENALIDOMIDE - REVLIMID</u>						
N021880 003	5635517	Oct 04, 2019	DS	U-1211		
	6045501	Aug 28, 2018		U-1210		
	6281230	Jul 24, 2016		U-1212		
	6315720	Oct 23, 2020		U-1210		
	6555554	Jul 24, 2016	DP	U-1211		
	6561976	Aug 28, 2018		U-1210		
	6561977	Oct 23, 2020		U-1210		
	6755784	Oct 23, 2020		U-1210		
	6908432	Aug 28, 2018		U-1210		
	7189740	Apr 11, 2023		U-1215		
	7465800	Apr 27, 2027	DS DP			
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023		U-1216		
	8204763	Aug 28, 2018		U-1249		
<u>LENALIDOMIDE - REVLIMID</u>						
N021880 004	5635517	Oct 04, 2019	DS	U-1211		
	6045501	Aug 28, 2018		U-1210		
	6281230	Jul 24, 2016		U-1212		
	6315720	Oct 23, 2020		U-1210		
	6555554	Jul 24, 2016	DP	U-1211		
	6561976	Aug 28, 2018		U-1210		
	6561977	Oct 23, 2020		U-1210		
	6755784	Oct 23, 2020		U-1210		
	6908432	Aug 28, 2018		U-1210		
	7189740	Apr 11, 2023		U-1215		
	7465800	Apr 27, 2027	DS DP			
	7968569	Oct 07, 2023		U-1216		
	8204763	Aug 28, 2018		U-1249		

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LENALIDOMIDE - REVLIMID</u>						
N021880 005	5635517	Oct 04, 2019	DS	U-1211		
	6045501	Aug 28, 2018		U-1210		
	6281230	Jul 24, 2016		U-1212		
	6315720	Oct 23, 2020		U-1210		
	6555554	Jul 24, 2016	DP	U-1211		
	6561976	Aug 28, 2018		U-1210		
	6561977	Oct 23, 2020		U-1210		
	6755784	Oct 23, 2020		U-1210		
	6908432	Aug 28, 2018		U-1210		
	7119106	Jul 24, 2016	DP			
	7189740	Apr 11, 2023		U-1215		
	7465800	Apr 27, 2027	DS DP			
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023		U-1216		
	8204763	Aug 28, 2018		U-1249		
<u>LEVETIRACETAM - KEPPRA</u>						
N021035 001				>A> NPP		Dec 16, 2014
<u>LEVETIRACETAM - KEPPRA</u>						
N021035 002				>A> NPP		Dec 16, 2014
<u>LEVETIRACETAM - KEPPRA</u>						
N021035 003				>A> NPP		Dec 16, 2014
<u>LEVETIRACETAM - KEPPRA</u>						
N021035 004				>A> NPP		Dec 16, 2014
<u>LEVETIRACETAM - KEPPRA</u>						
N021505 001				>A> NPP		Dec 16, 2014
<u>LIDOCAINE; TETRACAIN - SYNERA</u>						
N021623 001	>A> 6465709	Jul 07, 2020	DP			
<u>LINAGLIPTIN - TRADJENTA</u>						
N201280 001	8119648	Aug 12, 2023		U-774		
	8178541	Aug 12, 2023		U-775		
	8178541	Aug 12, 2023		U-1245		
	8178541	Aug 12, 2023		U-1244		
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N201281 001	6303661	Apr 24, 2017		U-802	NCE	May 02, 2016
	6890898	Feb 02, 2019		U-1039	NC	Jan 30, 2015
	7078381	Feb 02, 2019		U-1039		
	7407955	Aug 12, 2023	DS DP			
	7459428	Feb 02, 2019		U-1039		
	8119648	Aug 12, 2023		U-802		
	8178541	Aug 12, 2023	DP	U-775		

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LINAGLIPITIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N201281 002	6303661	Apr 24, 2017	U-802		NCE	May 02, 2016
	6890898	Feb 02, 2019	U-1039		NC	Jan 30, 2015
	7078381	Feb 02, 2019	U-1039			
	7407955	Aug 12, 2023	DS DP			
	7459428	Feb 02, 2019	U-1039			
	8119648	Aug 12, 2023	U-802			
	8178541	Aug 12, 2023	DP U-775			
<u>LINAGLIPITIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N201281 003	6303661	Apr 24, 2017	U-802		NCE	May 02, 2016
	6890898	Feb 02, 2019	U-1039		NC	Jan 30, 2015
	7078381	Feb 02, 2019	U-1039			
	7407955	Aug 12, 2023	DS DP			
	7459428	Feb 02, 2019	U-1039			
	8119648	Aug 12, 2023	U-802			
	8178541	Aug 12, 2023	DP U-775			
<u>LIRAGLUTIDE RECOMBINANT - VICTOZA</u>						
N022341 001	8114833	Aug 13, 2025	DP		M-115	Apr 06, 2015
<u>LISDEXAMFETAMINE Dimesylate - VYVANSE</u>						
N021977 001	7662788	Feb 24, 2023	U-727		I-645	Jan 31, 2015
	7713936	Feb 24, 2023	U-727			
<u>LISDEXAMFETAMINE Dimesylate - VYVANSE</u>						
N021977 002	7662788	Feb 24, 2023	U-727		I-645	Jan 31, 2015
	7713936	Feb 24, 2023	U-727			
<u>LISDEXAMFETAMINE Dimesylate - VYVANSE</u>						
N021977 003	7662788	Feb 24, 2023	U-727		I-645	Jan 31, 2015
	7713936	Feb 24, 2023	U-727			
<u>LISDEXAMFETAMINE Dimesylate - VYVANSE</u>						
N021977 004	7662788	Feb 24, 2023	U-727		I-645	Jan 31, 2015
	7671030	Feb 24, 2023	DP U-727			
	7674774	Mar 18, 2023	DP U-842			
	7678771	Mar 25, 2023	DP U-842			
	7687467	Apr 08, 2023	DP U-842			
	7713936	Feb 24, 2023	U-727			
	7718619	Feb 24, 2023	DP U-842			
	7723305	Feb 24, 2023	DP U-842			
<u>LISDEXAMFETAMINE Dimesylate - VYVANSE</u>						
N021977 005	7662788	Feb 24, 2023	U-727		I-645	Jan 31, 2015
	7713936	Feb 24, 2023	U-727			
<u>LISDEXAMFETAMINE Dimesylate - VYVANSE</u>						
N021977 006	7662788	Feb 24, 2023	U-727		I-645	Jan 31, 2015
	7713936	Feb 24, 2023	U-727			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LORCASERIN HYDROCHLORIDE - BELVIO</u>						
N022529 001	>A> 6953787	Apr 10, 2023	DS DP U-1253		NCE	Jun 27, 2017
	>A> 6953787	Apr 10, 2023	DS DP U-1254			
	>A> 6953787	Apr 10, 2023	DS DP U-1255			
	>A> 6953787	Apr 10, 2023	DS DP U-1252			
	>A> 7514422	Apr 10, 2023		U-1255		
	>A> 7514422	Apr 10, 2023		U-1252		
	>A> 7514422	Apr 10, 2023		U-1253		
	>A> 7514422	Apr 10, 2023		U-1254		
	>A> 7977329	Apr 10, 2023	DS DP	U-1253		
	>A> 7977329	Apr 10, 2023	DS DP	U-1254		
	>A> 7977329	Apr 10, 2023	DS DP	U-1255		
	>A> 7977329	Apr 10, 2023	DS DP	U-1252		
	>A> 8168624	Apr 18, 2029	DS DP			
	>A> 8207158	Apr 10, 2023		U-1255		
	>A> 8207158	Apr 10, 2023		U-1252		
	>A> 8207158	Apr 10, 2023		U-1253		
	>A> 8207158	Apr 10, 2023		U-1254		
<u>LUBIPROSTONE - AMITIZA</u>						
N021908 001	8097649	Oct 16, 2020	DP			
	8097653	Nov 14, 2022		U-1214		
	8114890	Sep 05, 2020	DP			
<u>LUBIPROSTONE - AMITIZA</u>						
N021908 002	8097649	Oct 16, 2020	DP			
	8114890	Sep 05, 2020	DP			
<u>LUCINACTANT - SURFAXIN</u>						
N021746 001	5407914	Nov 17, 2012	DS DP	U-1242		
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N200603 001				D-134	Apr 26, 2015	
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N200603 002				D-134	Apr 26, 2015	
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N200603 003				D-134	Apr 26, 2015	
				NCE	Oct 28, 2015	
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N200603 004				D-134	Apr 26, 2015	
				NCE	Oct 28, 2015	
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET</u>						
N021842 001				M-116	May 17, 2015	
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET</u>						
N021842 002				M-116	May 17, 2015	

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N202270 001	6303661	Apr 24, 2017	U-1227			
	6340475	Sep 19, 2016	DP			
	6635280	Sep 19, 2016	DP			
	6699871	Jul 26, 2022	DS DP U-1227			
	6890898	Feb 02, 2019	U-1228			
	7078381	Feb 02, 2019	U-1227			
	7125873	Jul 26, 2022	DP U-1227			
	7326708	Apr 11, 2026	DS DP U-1227			
	7459428	Feb 02, 2019	U-1227			
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N202270 002	6303661	Apr 24, 2017	U-1227			
	6340475	Sep 19, 2016	DP			
	6635280	Sep 19, 2016	DP			
	6699871	Jul 26, 2022	DS DP U-1227			
	6890898	Feb 02, 2019	U-1228			
	7078381	Feb 02, 2019	U-1227			
	7125873	Jul 26, 2022	DP U-1227			
	7326708	Apr 11, 2026	DS DP U-1227			
	7459428	Feb 02, 2019	U-1227			
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N202270 003	6303661	Apr 24, 2017	U-1227			
	6340475	Sep 19, 2016	DP			
	6635280	Sep 19, 2016	DP			
	6699871	Jul 26, 2022	DS DP U-1227			
	6890898	Feb 02, 2019	U-1228			
	7078381	Feb 02, 2019	U-1227			
	7125873	Jul 26, 2022	DP U-1227			
	7326708	Apr 11, 2026	DS DP U-1227			
	7459428	Feb 02, 2019	U-1227			
<u>METHYLPHENIDATE HYDROCHLORIDE - CONCERTA</u>						
N021121 001	8163798	Jul 31, 2017	DP			
	8163798*PED	Jan 31, 2018				
<u>METHYLPHENIDATE HYDROCHLORIDE - CONCERTA</u>						
N021121 002	8163798	Jul 31, 2017	DP			
	8163798*PED	Jan 31, 2018				
<u>METHYLPHENIDATE HYDROCHLORIDE - CONCERTA</u>						
N021121 003	8163798	Jul 31, 2017	DP			
	8163798*PED	Jan 31, 2018				
<u>METHYLPHENIDATE HYDROCHLORIDE - CONCERTA</u>						
N021121 004	8163798	Jul 31, 2017	DP			
	8163798*PED	Jan 31, 2018				
<u>METHYLPHENIDATE HYDROCHLORIDE - METHYLPHENIDATE HYDROCHLORIDE</u>						
A078458 001				PC		Jul 01, 2012
<u>METHYLPHENIDATE HYDROCHLORIDE - METHYLPHENIDATE HYDROCHLORIDE</u>						
A078458 002				PC		Jul 01, 2012

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>METHYLPHENIDATE HYDROCHLORIDE - METHYLPHENIDATE HYDROCHLORIDE</u>						
A078458 003					PC	Jul 01, 2012
<u>MICONAZOLE - ORAVIG</u>						
N022404 001	7651698	Sep 11, 2022	U-1051			
<u>MIFEPRISTONE - KORLYM</u>						
N202107 001					NP	Feb 17, 2015
<u>MINOCYCLINE HYDROCHLORIDE - ARRESTIN</u>						
N050781 001	6682348	Mar 29, 2022	DP			
<u>MIRABEGRON - MYRBETRIQ</u>						
N202611 001	>A> 6346532	Oct 15, 2018	DS DP		NCE	Jun 28, 2017
	>A> 6562375	Aug 01, 2020	DP			
	>A> 6699503	Sep 10, 2013	DP			
	>A> 7342117	Nov 04, 2023	DS			
	>A> 7750029	Dec 18, 2023	U-913			
	>A> 7982049	Nov 04, 2023	DP			
<u>MIRABEGRON - MYRBETRIQ</u>						
N202611 002	>A> 6346532	Oct 15, 2018	DS DP		NCE	Jun 28, 2017
	>A> 6562375	Aug 01, 2020	DP			
	>A> 6699503	Sep 10, 2013	DP			
	>A> 7342117	Nov 04, 2023	DS			
	>A> 7750029	Dec 18, 2023	U-913			
	>A> 7982049	Nov 04, 2023	DP			
<u>MITOMYCIN - MITOSOL</u>						
N022572 001	7806265	Feb 01, 2029	DP		ODE	Feb 07, 2019
	8186511	Jul 19, 2026	DP			
<u>MOMETASONE FUROATE - ASMANEX TWISTHALER</u>						
N021067 001	8173172	Mar 17, 2018	DP			
	8173172*PED	Sep 17, 2018				
<u>MOMETASONE FUROATE - ASMANEX TWISTHALER</u>						
N021067 002	8173172	Mar 17, 2018	DP			
	8173172*PED	Sep 17, 2018				
<u>MONTELUKAST SODIUM - SINGULAIR</u>						
N020829 002					NPP	Mar 26, 2015
<u>MONTELUKAST SODIUM - SINGULAIR</u>						
N020830 001					NPP	Mar 26, 2015
<u>MONTELUKAST SODIUM - SINGULAIR</u>						
N020830 002					NPP	Mar 26, 2015
<u>MONTELUKAST SODIUM - SINGULAIR</u>						
N021409 001					NPP	Mar 26, 2015
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N022321 001	8158156	Jun 19, 2027	U-1241			
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N022321 002	8158156	Jun 19, 2027	U-1241			
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N022321 003	8158156	Jun 19, 2027	U-1241			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N022321 004	8158156	Jun 19, 2027	U-1241			
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N022321 005	8158156	Jun 19, 2027	U-1241			
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N022321 006	8158156	Jun 19, 2027	U-1241			
<u>NAFTIFINE HYDROCHLORIDE - NAFTIN</u>						
N019599 002				NS		Jan 13, 2015
<u>NICOTINE - NICODERM CQ</u>						
N020165 004	8075911	May 22, 2021	DP			
<u>NICOTINE - NICODERM CQ</u>						
N020165 005	8075911	May 22, 2021	DP			
<u>NICOTINE - NICODERM CQ</u>						
N020165 006	8075911	May 22, 2021	DP			
<u>NILOTINIB HYDROCHLORIDE MONOHYDRATE - TASIGNA</u>						
N022068 001	8163904	Aug 23, 2028	DS DP			
<u>NILOTINIB HYDROCHLORIDE MONOHYDRATE - TASIGNA</u>						
N022068 002	8163904	Aug 23, 2028	DS DP			
<u>NITRIC OXIDE - INOMAX</u>						
N020845 002	5558083	Nov 22, 2013	DP U-1226			
	5558083*PED	May 22, 2014				
	5732693	Dec 13, 2016	DP U-1230			
	5732693*PED	Jun 13, 2017				
	5752504	Dec 13, 2016	DP U-1230			
	5752504*PED	Jun 13, 2017				
<u>NITRIC OXIDE - INOMAX</u>						
N020845 003	5558083	Nov 22, 2013	DP U-1226			
	5558083*PED	May 22, 2014				
	5732693	Dec 13, 2016	DP U-1230			
	5732693*PED	Jun 13, 2017				
	5752504	Dec 13, 2016	DP U-1230			
	5752504*PED	Jun 13, 2017				
<u>NITROGLYCERIN - NITROLINGUAL PUMPSPRAY</u>						
N018705 002	7872049	Mar 14, 2028	DP U-39			
<u>OXCARBAZEPINE - TRILEPTAL</u>						
N021285 001	8119148	Dec 19, 2020	DP U-724			
	8119148*PED	Jun 19, 2021				
<u>OXYBUTYNIN - ANTUROL</u>						
N202513 001	7029694	Apr 26, 2020	DP U-318			
	7179483	Apr 26, 2020	U-318			
	7198801	Jun 25, 2022	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N022272 001	8114383	Oct 10, 2024	DP			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N022272 002	8114383	Oct 10, 2024	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N022272 003	8114383	Oct 10, 2024	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N022272 004	8114383	Oct 10, 2024	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N022272 005	8114383	Oct 10, 2024	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N022272 006	8114383	Oct 10, 2024	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N022272 007	8114383	Oct 10, 2024	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 001	7851482	Jul 10, 2029	DS			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 002	7851482	Jul 10, 2029	DS			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 003	7851482	Jul 10, 2029	DS			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 004	7851482	Jul 10, 2029	DS			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 005	7851482	Jul 10, 2029	DS			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 006	7851482	Jul 10, 2029	DS			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 007	7851482	Jul 10, 2029	DS			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
<u>PACLITAXEL - ABRAXANE</u>						
N021660 001	8138229	Dec 09, 2023	DP U-1092			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u>						
	N020725 004			I-625 M-93 NCE	Apr 30, 2013 Jul 29, 2019 Apr 30, 2014	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - PERTZYE</u>						
	N022175 001			NCE	May 17, 2017	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - PERTZYE</u>						
	N022175 002			NCE	May 17, 2017	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ULTRESA</u>						
	N022222 001			NCE	Mar 01, 2017	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ULTRESA</u>						
	N022222 002			NCE	Mar 01, 2017	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ULTRESA</u>						
	N022222 003			NCE	Mar 01, 2017	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - VIOKACE</u>						
	N022542 001			NCE	Mar 01, 2017	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - VIOKACE</u>						
	N022542 002			NCE	Mar 01, 2017	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
	N022210 005			NCE	Aug 27, 2014	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
	N022210 006			NCE	Aug 27, 2014	
<u>PAZOPANIB HYDROCHLORIDE - VOTRIENT</u>						
N022465 001	8114885	Dec 19, 2021	DS DP	I-649 ODE	Apr 26, 2015 Apr 26, 2019	
<u>PAZOPANIB HYDROCHLORIDE - VOTRIENT</u>						
N022465 002	8114885	Dec 19, 2021	DS DP	I-649 ODE	Apr 26, 2015 Apr 26, 2019	
<u>PEGINESATIDE ACETATE - OMONTYS</u>						
N202799 007	7084245	May 12, 2024	DS DP U-1238			
	7414105	May 12, 2024	DS DP U-1238			
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026	U-1238			
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026	U-1238			
<u>PEGINESATIDE ACETATE - OMONTYS</u>						
N202799 008	7084245	May 12, 2024	DS DP U-1238			
	7414105	May 12, 2024	DS DP U-1238			
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026	U-1238			
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026	U-1238			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N202799 001	7084245	May 12, 2024	DS DP U-1238		NCE	Mar 27, 2017
	7414105	May 12, 2024	DS DP U-1238			
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026		U-1238		
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026		U-1238		
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N202799 002	7084245	May 12, 2024	DS DP U-1238		NCE	Mar 27, 2017
	7414105	May 12, 2024	DS DP U-1238			
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026		U-1238		
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026		U-1238		
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N202799 003	7084245	May 12, 2024	DS DP U-1238		NCE	Mar 27, 2017
	7414105	May 12, 2024	DS DP U-1238			
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026		U-1238		
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026		U-1238		
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N202799 004	7084245	May 12, 2024	DS DP U-1238		NCE	Mar 27, 2017
	7414105	May 12, 2024	DS DP U-1238			
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026		U-1238		
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026		U-1238		
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N202799 005	7084245	May 12, 2024	DS DP U-1238		NCE	Mar 27, 2017
	7414105	May 12, 2024	DS DP U-1238			
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026		U-1238		
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026		U-1238		
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N202799 006	7084245	May 12, 2024	DS DP U-1238		NCE	Mar 27, 2017
	7414105	May 12, 2024	DS DP U-1238			
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026		U-1238		
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026		U-1238		
<u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u>						
N202088 003	6149938	Jul 23, 2018		DP U-1243		
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u>						
N022580 001					>A> NC	Jul 17, 2015
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u>						
N022580 002					>A> NC	Jul 17, 2015

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u>						
	N022580 003			>A> NC		Jul 17, 2015
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u>						
	N022580 004			>A> NC		Jul 17, 2015
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u>						
N050684 001	8133883	Apr 14, 2023	DP U-282			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u>						
N050684 002	8133883	Apr 14, 2023	DP U-282			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u>						
N050684 003	8133883	Apr 14, 2023	DP U-282			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u>						
N050684 004	8133883	Apr 14, 2023	DP U-282			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u>						
N050750 001	8133883	Apr 14, 2023	DP U-282			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u>						
N050750 002	8133883	Apr 14, 2023	DP U-282			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u>						
N050750 003	8133883	Apr 14, 2023	DP U-282			
<u>PREDNISOLONE ACETATE - FLO-PRED</u>						
N022067 001	7799331	Oct 11, 2028	DP U-139			
	7799331	Oct 11, 2028	DP U-1068			
<u>PREDNISOLONE ACETATE - FLO-PRED</u>						
N022067 002	7799331	Oct 11, 2028	DP U-139			
	7799331	Oct 11, 2028	DP U-1068			
<u>PREGABALIN - LYRICA</u>						
N021446 001	>A> 6001876	Dec 30, 2018	U-819	Y	I-651	Jun 20, 2015
	>A> 6001876	Dec 30, 2018	U-55	Y		
	>A> RE41920	Dec 30, 2018	U-1250			
<u>PREGABALIN - LYRICA</u>						
N021446 002	>A> 6001876	Dec 30, 2018	U-819	Y	I-651	Jun 20, 2015
	>A> 6001876	Dec 30, 2018	U-55	Y		
	>A> RE41920	Dec 30, 2018	U-1250			
<u>PREGABALIN - LYRICA</u>						
N021446 003	>A> 6001876	Dec 30, 2018	U-819	Y	I-651	Jun 20, 2015
	>A> 6001876	Dec 30, 2018	U-55	Y		
	>A> RE41920	Dec 30, 2018	U-1250			
<u>PREGABALIN - LYRICA</u>						
N021446 004	>A> 6001876	Dec 30, 2018	U-819	Y	I-651	Jun 20, 2015
	>A> 6001876	Dec 30, 2018	U-55	Y		
	>A> RE41920	Dec 30, 2018	U-1250			
<u>PREGABALIN - LYRICA</u>						
N021446 005	>A> 6001876	Dec 30, 2018	U-819	Y	I-651	Jun 20, 2015
	>A> 6001876	Dec 30, 2018	U-55	Y		
	>A> RE41920	Dec 30, 2018	U-1250			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PREGABALIN - LYRICA</u>						
N021446 006	>A> 6001876	Dec 30, 2018	U-819	Y	I-651	Jun 20, 2015
	>A> 6001876	Dec 30, 2018	U-55	Y		
	>A> RE41920	Dec 30, 2018	U-1250			
<u>PREGABALIN - LYRICA</u>						
N021446 007	>A> 6001876	Dec 30, 2018	U-819	Y	I-651	Jun 20, 2015
	>A> 6001876	Dec 30, 2018	U-55	Y		
	>A> RE41920	Dec 30, 2018	U-1250			
<u>PREGABALIN - LYRICA</u>						
N021446 008	>A> 6001876	Dec 30, 2018	U-819	Y	I-651	Jun 20, 2015
	>A> 6001876	Dec 30, 2018	U-55	Y		
	>A> RE41920	Dec 30, 2018	U-1250			
<u>PREGABALIN - LYRICA</u>						
N022488 001	>A> 6001876	Dec 30, 2018	U-819	Y		
	>A> 6001876	Dec 30, 2018	U-55	Y		
	>A> RE41920	Dec 30, 2018	U-1250			
<u>PROPRANOLOL HYDROCHLORIDE - INNOPRAN XL</u>						
N021438 001	6500454	Oct 04, 2021	DP			
<u>PROPRANOLOL HYDROCHLORIDE - INNOPRAN XL</u>						
N021438 002	6500454	Oct 04, 2021	DP			
<u>Raltegravir Potassium - ISENTRESS</u>						
N022145 001					M-114	Mar 28, 2015
<u>Raltegravir Potassium - ISENTRESS</u>						
N203045 001	7169780	Oct 03, 2023	DS DP		M-114	Mar 28, 2015
	7217713	Oct 21, 2022	U-257			
	7435734	Oct 21, 2022	U-257			
	7754731	Mar 11, 2029	DS DP U-257			
<u>Raltegravir Potassium - ISENTRESS</u>						
N203045 002	7169780	Oct 03, 2023	DS DP		M-114	Mar 28, 2015
	7217713	Oct 21, 2022	U-257			
	7435734	Oct 21, 2022	U-257			
	7754731	Mar 11, 2029	DS DP U-257			
<u>Regadenoson - LEXISCAN</u>						
N022161 001	8106029	Jun 22, 2019	U-1042			
	8106183	Feb 02, 2027	DS			
	8133879	Jun 22, 2019	DP			
	8183226	Jun 22, 2019	U-116			
<u>Retapamulin - ALTABAX</u>						
N022055 001	RE43390	Apr 12, 2021	DS DP U-805			
<u>Rifaximin - XIFAXAN</u>						
N021361 001	8158644	Jun 19, 2024	DP			
	8158781	Jun 19, 2024	DS			
	8193196	Sep 02, 2027	DS DP			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RIFAXIMIN - XIFAXAN</u>						
N022554 001	8158644	Jun 19, 2024	DP			
	8158781	Jun 19, 2024	DS			
	8193196	Sep 02, 2027	DS DP			
<u>RILPIVIRINE HYDROCHLORIDE - EDURANT</u>						
N202022 001	8080551	Apr 11, 2023	DS DP			
	8101629	Aug 09, 2022	DP			
<u>RIZATRIPTAN BENZOATE - MAXALT</u>						
N020864 001	5602162	Jan 28, 2012		Y		
	5602162*PED	Jul 28, 2012				
<u>RIZATRIPTAN BENZOATE - MAXALT</u>						
N020864 002	5602162	Jan 28, 2012		Y		
	5602162*PED	Jul 28, 2012				
<u>RIZATRIPTAN BENZOATE - MAXALT-MLT</u>						
N020865 001	5602162	Jan 28, 2012	U-240	Y		
	5602162*PED	Jul 28, 2012				
<u>RIZATRIPTAN BENZOATE - MAXALT-MLT</u>						
N020865 002	5602162	Jan 28, 2012	U-240	Y		
	5602162*PED	Jul 28, 2012				
<u>ROMIDEPSIN - ISTODAX</u>						
N022393 001	>A> 4977138	Aug 22, 2013	DS DP			
<u>ROTIGOTINE - NEUPRO</u>						
N021829 001				I-647	Apr 02, 2015	
				I-646	Apr 02, 2015	
<u>ROTIGOTINE - NEUPRO</u>						
N021829 002				I-647	Apr 02, 2015	
				I-646	Apr 02, 2015	
<u>ROTIGOTINE - NEUPRO</u>						
N021829 003				I-647	Apr 02, 2015	
				I-646	Apr 02, 2015	
<u>ROTIGOTINE - NEUPRO</u>						
N021829 004	6699498	Nov 27, 2020	DP	I-647	Apr 02, 2015	
	6884434	Mar 30, 2021	DP	I-646	Apr 02, 2015	
	7413747	Mar 18, 2019	DP			
<u>ROTIGOTINE - NEUPRO</u>						
N021829 005	6699498	Nov 27, 2020	DP	I-647	Apr 02, 2015	
	6884434	Mar 30, 2021	DP	I-646	Apr 02, 2015	
	7413747	Mar 18, 2019	DP			
<u>ROTIGOTINE - NEUPRO</u>						
N021829 006	6699498	Nov 27, 2020	DP	I-647	Apr 02, 2015	
	6884434	Mar 30, 2021	DP	I-646	Apr 02, 2015	
	7413747	Mar 18, 2019	DP			
<u>RUFINAMIDE - BANZEL</u>						
N201367 001	>A> 6740669	Nov 14, 2022	DS DP			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SILDENAFIL CITRATE - REVATIO</u>						
N021845 001	5250534	Mar 27, 2012	DS DP		I-598	May 07, 2012
	5250534*PED	Sep 27, 2012			PED	Nov 07, 2012
<u>SILDENAFIL CITRATE - REVATIO</u>						
N022473 001	5250534	Mar 27, 2012	DS DP		NDF	Nov 20, 2012
	5250534*PED	Sep 27, 2012			PED	May 20, 2013
<u>SILDENAFIL CITRATE - VIAGRA</u>						
N020895 001	5250534	Mar 27, 2012				
	5250534*PED	Sep 27, 2012				
	6469012	Oct 22, 2019		U-155		
	6469012*PED	Apr 22, 2020				
<u>SILDENAFIL CITRATE - VIAGRA</u>						
N020895 002	5250534	Mar 27, 2012				
	5250534*PED	Sep 27, 2012				
	6469012	Oct 22, 2019	U-155			
	6469012*PED	Apr 22, 2020				
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N202343 001	8168637	Jun 26, 2022	DP U-1188			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N202343 002	8168637	Jun 26, 2022	DP U-1188			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N202343 003	8168637	Jun 26, 2022	DP U-1188			
<u>SODIUM NITRITE - SODIUM NITRITE</u>						
N203922 001					ODE	Jan 14, 2018
<u>SODIUM THIOSULFATE - SODIUM THIOSULFATE</u>						
N203923 001					ODE	Jan 14, 2018
<u>SUMATRIPTAN SUCCINATE - SUMAVEL DOSEPRO</u>						
N022239 001	8118771	Aug 10, 2023	DP			
<u>TAFLUPROST - ZIOPTAN</u>						
N202514 001	5886035	Dec 18, 2017	DS DP U-778		NCE	Feb 10, 2017
<u>TALIGLUCERASE ALFA - ELELYSO</u>						
N022458 001					NCE	May 01, 2017
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N200533 001	8114383	Oct 10, 2024	DP			
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N200533 002	8114383	Oct 10, 2024	DP			
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N200533 003	8114383	Oct 10, 2024	DP			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N200533 004	8114383	Oct 10, 2024	DP			
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N200533 005	8114383	Oct 10, 2024	DP			
<u>TAZAROTENE - FABIOR</u>						
N202428 001					NDF	May 11, 2015
<u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u>						
N022110 001	8101575	May 01, 2021	DP			
	8158580	May 01, 2021	DP			
<u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u>						
N022110 002	8101575	May 01, 2021	DP			
	8158580	May 01, 2021	DP			
<u>TELBIVUDINE - TYZEKA</u>						
N022154 001	7858594	Sep 11, 2023	DS DP U-999			
<u>TEMSIROLIMUS - TORISEL</u>						
N022088 001	5362718	Apr 18, 2014	DS DP		M-92	Jul 09, 2013
	5362718*PED	Oct 18, 2014			M-91	Apr 26, 2013
	8026276	Jan 20, 2026	DP		ODE	May 30, 2014
	8026276*PED	Jul 20, 2026			PED	Nov 30, 2014
					PED	Jan 09, 2014
					PED	Oct 26, 2013
<u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u>						
N021356 001					NPP	Jan 18, 2015
					PED	Jul 18, 2015
<u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u>						
N021356 002	5922695	Jul 25, 2017	DS U-250		M-95	Oct 01, 2013
	5922695	Jul 25, 2017	DS U-256		NPP	Jan 18, 2015
	5922695	Jul 25, 2017	DS U-999		NPP	Mar 24, 2013
	5922695	Jul 25, 2017	DS U-248		ODE	Mar 24, 2017
	5922695*PED	Jan 25, 2018			PED	Sep 24, 2017
	5935946	Jul 25, 2017	DS DP U-999		PED	Jul 18, 2015
	5935946	Jul 25, 2017	DS DP U-248		PED	Apr 01, 2014
	5935946	Jul 25, 2017	DS DP U-250		PED	Sep 24, 2013
	5935946	Jul 25, 2017	DS DP U-256			
	5935946*PED	Jan 25, 2018				
	5977089	Jul 25, 2017	DS DP U-250			
	5977089	Jul 25, 2017	DS DP U-256			
	5977089	Jul 25, 2017	DS DP U-999			
	5977089	Jul 25, 2017	DS DP U-248			
	5977089*PED	Jan 25, 2018				
	6043230	Jul 25, 2017	U-248			
	6043230	Jul 25, 2017	U-250			
	6043230	Jul 25, 2017	U-256			
	6043230	Jul 25, 2017	U-999			
	6043230*PED	Jan 25, 2018				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u>						
N021356 003	5922695	Jul 25, 2017	DS U-250		M-95	Oct 01, 2013
	5922695	Jul 25, 2017	DS U-256		NPP	Jan 18, 2015
	5922695	Jul 25, 2017	DS U-999		NPP	Mar 24, 2013
	5922695	Jul 25, 2017	DS U-248		ODE	Mar 24, 2017
	5922695*PED	Jan 25, 2018			PED	Sep 24, 2017
	5935946	Jul 25, 2017	DS DP U-999		PED	Jul 18, 2015
	5935946	Jul 25, 2017	DS DP U-248		PED	Apr 01, 2014
	5935946	Jul 25, 2017	DS DP U-250		PED	Sep 24, 2013
	5935946	Jul 25, 2017	DS DP U-256			
	5935946*PED	Jan 25, 2018				
	5977089	Jul 25, 2017	DS DP U-250			
	5977089	Jul 25, 2017	DS DP U-256			
	5977089	Jul 25, 2017	DS DP U-999			
	5977089	Jul 25, 2017	DS DP U-248			
	5977089*PED	Jan 25, 2018				
	6043230	Jul 25, 2017	U-248			
	6043230	Jul 25, 2017	U-250			
	6043230	Jul 25, 2017	U-256			
	6043230	Jul 25, 2017	U-999			
	6043230*PED	Jan 25, 2018				
<u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u>						
N021356 004	5922695	Jul 25, 2017	DS U-250		M-95	Oct 01, 2013
	5922695	Jul 25, 2017	DS U-256		NPP	Jan 18, 2015
	5922695	Jul 25, 2017	DS U-999		NPP	Mar 24, 2013
	5922695	Jul 25, 2017	DS U-248		ODE	Mar 24, 2017
	5922695*PED	Jan 25, 2018			PED	Sep 24, 2017
	5935946	Jul 25, 2017	DS DP U-999		PED	Jul 18, 2015
	5935946	Jul 25, 2017	DS DP U-248		PED	Apr 01, 2014
	5935946	Jul 25, 2017	DS DP U-250		PED	Sep 24, 2013
	5935946	Jul 25, 2017	DS DP U-256			
	5935946*PED	Jan 25, 2018				
	5977089	Jul 25, 2017	DS DP U-250			
	5977089	Jul 25, 2017	DS DP U-256			
	5977089	Jul 25, 2017	DS DP U-999			
	5977089	Jul 25, 2017	DS DP U-248			
	5977089*PED	Jan 25, 2018				
	6043230	Jul 25, 2017	U-248			
	6043230	Jul 25, 2017	U-250			
	6043230	Jul 25, 2017	U-256			
	6043230	Jul 25, 2017	U-999			
	6043230*PED	Jan 25, 2018				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u>						
N022577 001	5922695	Jul 25, 2017	DS U-250		M-95	Oct 01, 2013
	5922695	Jul 25, 2017	DS U-256		NPP	Mar 24, 2013
	5922695	Jul 25, 2017	DS U-999		NDF	Jan 18, 2015
	5922695	Jul 25, 2017	DS U-248		ODE	Mar 24, 2017
	5922695*PED	Jan 25, 2018			PED	Sep 24, 2017
	5935946	Jul 25, 2017	DP U-999	Y	PED	Jul 18, 2015
	5935946	Jul 25, 2017	DP U-248	Y	PED	Apr 01, 2014
	5935946	Jul 25, 2017	DP U-250	Y		
	5935946	Jul 25, 2017	DP U-256	Y		
	5935946*PED	Jan 25, 2018				
	5977089	Jul 25, 2017	DS DP U-250			
	5977089	Jul 25, 2017	DS DP U-256			
	5977089	Jul 25, 2017	DS DP U-999			
	5977089	Jul 25, 2017	DS DP U-248			
	5977089*PED	Jan 25, 2018				
	6043230	Jul 25, 2017	U-248			
	6043230	Jul 25, 2017	U-250			
	6043230	Jul 25, 2017	U-256			
	6043230	Jul 25, 2017	U-999			
	6043230*PED	Jan 25, 2018				
<u>TESAMORELIN ACETATE - EGRIFTA</u>						
N022505 002					NCE	Nov 10, 2015
<u>TESTOSTERONE - TESTIM</u>						
N021454 001	8178518	Apr 21, 2023	DP			
<u>TESTOSTERONE - TESTOSTERONE</u>						
N202763 001					NP	Feb 14, 2015
<u>THALIDOMIDE - THALOMID</u>						
N020785 001	8143283	Mar 01, 2013	U-1236			
	8204763	Aug 28, 2018	U-1249			
<u>THALIDOMIDE - THALOMID</u>						
N020785 002	8143283	Mar 01, 2013	U-1236			
	8204763	Aug 28, 2018	U-1249			
<u>THALIDOMIDE - THALOMID</u>						
N020785 003	8143283	Mar 01, 2013	U-1236			
	8204763	Aug 28, 2018	U-1249			
<u>THALIDOMIDE - THALOMID</u>						
N020785 004	8143283	Mar 01, 2013	U-1236			
	8204763	Aug 28, 2018	U-1249			
<u>TIZANIDINE HYDROCHLORIDE - ZANAFLEX</u>						
N021447 001	>A> 6455557	Nov 28, 2021		Y		
<u>TIZANIDINE HYDROCHLORIDE - ZANAFLEX</u>						
N021447 002	>A> 6455557	Nov 28, 2021		Y		
<u>TIZANIDINE HYDROCHLORIDE - ZANAFLEX</u>						
N021447 003	>A> 6455557	Nov 28, 2021		Y		

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TOPIRAMATE - TOPAMAX</u>						
N020505 001					NPP	Jul 15, 2014
<u>TOPIRAMATE - TOPAMAX</u>						
N020505 002					NPP	Jul 15, 2014
<u>TOPIRAMATE - TOPAMAX</u>						
N020505 004					NPP	Jul 15, 2014
<u>TOPIRAMATE - TOPAMAX</u>						
N020505 005					NPP	Jul 15, 2014
<u>TOPIRAMATE - TOPAMAX</u>						
N020844 001					NPP	Jul 15, 2014
<u>TOPIRAMATE - TOPAMAX</u>						
N020844 002					NPP	Jul 15, 2014
<u>TOPOTECAN HYDROCHLORIDE - HYCAMTIN</u>						
N020981 001	8158645	Dec 10, 2024	DP			
<u>TOPOTECAN HYDROCHLORIDE - HYCAMTIN</u>						
N020981 002	8158645	Dec 10, 2024	DP			
<u>TRAMADOL HYDROCHLORIDE - TRAMADOL HYDROCHLORIDE</u>						
A091607 001					PC	Jun 27, 2012
<u>TRAMADOL HYDROCHLORIDE - TRAMADOL HYDROCHLORIDE</u>						
A091607 002					PC	Jun 27, 2012
<u>TRAMADOL HYDROCHLORIDE - TRAMADOL HYDROCHLORIDE</u>						
A091607 003					PC	Jun 27, 2012
<u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u>						
N022411 001	8133893	Mar 13, 2029	DS DP			
<u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u>						
N022411 002	8133893	Mar 13, 2029	DS DP			
<u>TRIAMCINOLONE ACETONIDE - TRIESENCE</u>						
N022048 001	8128960	Dec 17, 2029	DP			
>A> 8211880		Mar 10, 2029	U-1258			
>A> 8211880		Mar 10, 2029	U-1257			
<u>TRYPAN BLUE - MEMBRANEBLUE</u>						
N022278 001					>A> ODE	Feb 20, 2016
<u>VEMURAFENIB - ZELBORAF</u>						
N202429 001	8143271	Jun 21, 2026	DS DP			
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N022567 001	8193195	Jun 05, 2022	U-839			
>A> 8236804		Jun 05, 2022	U-839			
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N022567 002	8193195	Jun 05, 2022	U-839			
>A> 8236804		Jun 05, 2022	U-839			
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N022567 003	8193195	Jun 05, 2022	U-839			
>A> 8236804		Jun 05, 2022	U-839			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2012

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VISMODEGIB - ERIVEDGE</u>						
	N203388 001				NCE	Jan 30, 2017
<u>VORINOSTAT - ZOLINZA</u>						
N021991 001	8093295	May 16, 2026	DP			
	8101663	Mar 04, 2023	U-892			
<u>ZIPRASIDONE HYDROCHLORIDE - ZIPRASIDONE HYDROCHLORIDE</u>						
	A077560 001				PC	Aug 29, 2012
<u>ZIPRASIDONE HYDROCHLORIDE - ZIPRASIDONE HYDROCHLORIDE</u>						
	A077560 002				PC	Aug 29, 2012
<u>ZIPRASIDONE HYDROCHLORIDE - ZIPRASIDONE HYDROCHLORIDE</u>						
	A077560 003				PC	Aug 29, 2012
<u>ZIPRASIDONE HYDROCHLORIDE - ZIPRASIDONE HYDROCHLORIDE</u>						
	A077560 004				PC	Aug 29, 2012

Footnote:

1. 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).
2. 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.

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, 31st 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/patternsall.cfm>

The current complete list of patent terms is available at
<http://www.accessdata.fda.gov/scripts/cder/ob/docs/excltermsall.cfm>