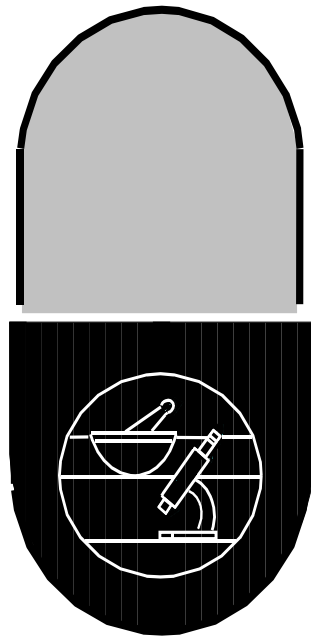


**CUMULATIVE  
SUPPLEMENT 2  
FEBRUARY 2020**



**APPROVED  
DRUG PRODUCTS**

**WITH  
THERAPEUTIC EQUIVALENCE EVALUATIONS**

**40th EDITION**

**Department of Health and Human Services**

**Food and Drug Administration  
Office of Medical Products and Tobacco  
Center for Drug Evaluation and Research  
Office of Generic Drugs  
Office of Generic Drug Policy**

**2020**

Prepared By  
Food and Drug Administration  
Office of Medical Products and Tobacco  
Center for Drug Evaluation and Research  
Office of Generic Drugs  
Office of Generic Drug Policy

**APPROVED DRUG PRODUCTS  
with  
THERAPEUTIC EQUIVALENCE EVALUATIONS**

**40<sup>th</sup> EDITION**

**Cumulative Supplement 2  
February 2020**

**CONTENTS**

*PAGE*

**Contents**

1.0	INTRODUCTION.....	v
1.1	HOW TO USE THE CUMULATIVE SUPPLEMENT .....	vi
1.2	CUMULATIVE SUPPLEMENT CONTENT .....	vi
1.3	APPLICANT NAME CHANGES .....	vii
1.4	LEVOTHYROXINE SODIUM .....	viii
1.5	AVAILABILITY OF THE EDITION.....	ix
1.6	REPORT OF COUNTS FOR THE PRESCRIPTION DRUG PRODUCT LIST.....	x
1.7	CUMULATIVE SUPPLEMENT LEGEND.....	xi
<b>DRUG PRODUCT LISTS</b>		
	Prescription Drug Product List .....	1-1
	OTC Drug Product List .....	2-1
	Drug Products with Approval under Section 505 of the Act Administered by the Center for Biologics Evaluation and Research List .....	3-1
	Orphan Product Designations and Approvals List .....	4-1
	Drug Products Which Must Demonstrate in vivo Bioavailability Only if Product Fails to Achieve Adequate Dissolution .....	5-1
<b>PATENT AND EXCLUSIVITY INFORMATION ADDENDUM</b>		
A.	Patent and Exclusivity Lists .....	A-1
B.	Patent and Exclusivity Terms .....	B-1

**APPROVED DRUG PRODUCTS  
with  
THERAPEUTIC EQUIVALENCE EVALUATIONS**

**40<sup>th</sup> EDITION**

**CUMULATIVE SUPPLEMENT 2  
FEBRUARY 2020**

**1.0 INTRODUCTION**

This Cumulative Supplement is one of a series of monthly updates to the *Approved Drug Products with Therapeutic Equivalence Evaluations* (the List, commonly known as the Orange Book). The List is composed of four parts: approved prescription drug products with therapeutic equivalence evaluations; approved over-the-counter (OTC) drug products for those drugs that may not be marketed without NDAs or ANDAs because they are not covered under existing OTC monographs; drug products with approval under Section 505 of the Federal Food, Drug, and Cosmetic Act (FD&C Act) administered by the Center for Biologics Evaluation and Research; and approved products that have never been marketed, are for exportation, are for military use, have been discontinued from marketing and we have not determined that they were withdrawn for safety or effectiveness reasons, 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, Discontinued Drug Product, 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, have been discontinued from marketing and we have not determined that they were withdrawn for safety or effectiveness reasons 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 this Edition List will then be added to the "Discontinued Drug Product List" appearing in the next Edition. The current Annual Edition Section 2., How To Use The Drug Product Lists, describes the layout and usage of the List.

## 1.1 HOW TO USE THE CUMULATIVE SUPPLEMENT

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.
  - Refer to CS Section 1.7 Cumulative Supplement Legend for types of changes
- New Drug Application (NDA) approvals 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 is current as of the date of publication.

Every effort is made to ensure the Cumulative Supplement is 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 [orangebook@fda.hhs.gov](mailto:orangebook@fda.hhs.gov).

### 1.3 APPLICANT NAME CHANGES

It is not practical to identify in the Cumulative Supplement each 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)

NEW APPLICANT NAME  
(NEW ABBREVIATED NAME)

AMNEAL PHARMACEUTICALS CO GMBH  
(AMNEAL PHARMS CO)

AMNEAL EU LTD  
(AMNEAL EU)

ANDA REPOSITORY LLC  
(ANDA REPOSITORY)

PHARMACEUTICS INTERNATIONAL INC  
(PII)

## 1.4 LEVOTHYROXINE SODIUM<sup>1</sup>

Because there are multiple reference listed drugs for levothyroxine sodium tablets and some reference listed drugs' sponsors have conducted studies to establish their drugs' therapeutic equivalence to other reference listed drugs, FDA has determined that its usual practice of assigning two or three character therapeutic equivalence codes may be potentially confusing and inadequate for these drug products. Looking at the Orange Book listing alone for a product identified as a reference listed drug or reference standard, it may be difficult to determine to which therapeutic equivalence code the reference listed drugs and/or reference standard designation corresponds. For example, Unithroid 0.3 mg strength has been assigned the therapeutic equivalence codes AB1, AB2, and AB3 and it is identified as the reference listed drug and reference standard, but it is unclear that the reference listed drug and reference standard designations are associated with the AB1 therapeutic equivalence code.

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

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

TE Code	Proprietary Name	Applicant	Strength	Appl No	RLD	RS
AB1	UNITHROID	STEVENS J	0.3MG	N021210	RLD	RS
AB2	SYNTHROID	ABBVIE	0.3MG	N021402	RLD	RS
AB3	LEVOXYL	KING PHARMS	0.2MG	N021301	RLD	RS
AB4	THYRO-TABS	ALVOGEN GROUP	0.3MG	N021116	RLD	-

<sup>1</sup> In previous editions of the Orange Book, FDA provided a chart outlining therapeutic equivalence codes for all .025 mg levothyroxine sodium drug products in the Active Section of the Orange Book. FDA has decided, for ease of review, to revise the chart to identify the NDAs for the reference listed drugs for each therapeutic equivalence code (i.e., AB1, AB2, AB3, and AB4), and their corresponding reference standards, which are identified in 0.2 and 0.3 mg strengths.

<sup>2</sup> Please consult the Active Section of the Orange Book for information on other strengths.

		HOLDINGS 4 LLC				
AB4	LEVOTHYROXINE SODIUM <sup>3</sup>	MYLAN	0.3MG	A076187	-	RS

## 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,  
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,

---

<sup>3</sup> Lloyd's Thyro-Tabs tablets (NDA 021116) (previously known as Levothroid) previously was listed in the Discontinued Drug Product List section of the Orange Book. It is the RLD for therapeutic equivalents identified with the AB4 code. During this time, Mylan's levothyroxine product (ANDA 076187) was selected as the reference standard for ANDA applicants to use to establish bioequivalence to Thyro-Tabs. It remains the reference standard for ANDA applicants to use to establish bioequivalence to Thyro-Tabs. If an ANDA that uses Mylan's levothyroxine product as its reference standard is approved, the ANDA will receive an AB4 rating. The ANDA applicant also may obtain an AB rating for its product to the other reference listed drugs (i.e., Unithroid, Synthroid, and Levoxyl) by submitting supplements that demonstrate that the generic product is bioequivalent to these other reference listed drugs and satisfies all other therapeutic equivalence criteria with respect to these reference listed drugs. See Letter from Janet Woodcock, M.D., Director, Center for Drug Evaluation and Research, FDA to Teri Nataline, Principal Consultant, Lachman Consultant Services, Inc., Docket No. FDA-2015-P-0403 (May 27, 2016).



<https://www.fda.gov/about-fda/reports-manuals-forms/forms>.

The current listing of the Orphan Product Designations and Approvals is available at <https://www.fda.gov/industry/developing-products-rare-diseases-conditions/designating-orphan-product-drugs-and-biological-products>.

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

### DEFINITIONS

#### Drug Product

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

#### New Molecular Entity

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

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

<u>CATEGORIES</u> <u>COUNTED</u>	<u>DEC</u> <u>2019</u>	<u>MAR</u> <u>2020</u>	<u>JUN</u> <u>2020</u>	<u>SEP</u> <u>2020</u>	<u>DEC</u> <u>2020</u>
DRUG PRODUCTS LISTED SINGLE SOURCE	20567	2743 (13.3%)			
MULTISOURCE	17824 (86.7%)				

THERAPEUTICALLY EQUIVALENT	17740 (86.3%)
NOT THERAPEUTICALLY EQUIVALENT	84 (0.4%)
EXCEPTIONS <sup>4</sup>	54 (0.3%)
NEW MOLECULAR ENTITIES APPROVED	17
NUMBER OF APPLICANTS	1162

## 1.7 CUMULATIVE SUPPLEMENT LEGEND

The List is sorted by Ingredient(s) and, within each grouping, by the Dosage Form; Route of Administration and then by trade name (or established name of the active ingredient, if no trade name exists).

The individual product record contains the Therapeutic Equivalence Code, Reference Listed Drug symbol, Reference Standard 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, new route(s) of administration, new trade names, and new product additions are preceded by >A> during the action month. The change month is the current CS month; the change code for new approvals is NEWA. Following months will display the same information without the >A>.

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

The change code and description:

NEWA	New drug product approval
NFTG	New first-time generic approval
CAHN <sup>5</sup>	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

<sup>4</sup> Amino acid containing products of varying composition (see Introduction, page xx of the List).

<sup>5</sup> The Cumulative Supplement (CS) currently displays a condensed 20 character collapsed applicant holder firm name and the Electronic Orange Book (EOB) query may display up to a 250-character full applicant holder firm name. An applicant holder firm name change usually changes both the collapsed name and long name. On occasion, only the long name is changed resulting in the CS displaying only the collapsed name for the >D> and >A> action. The new firm long name will display in the EOB query.

CFTG Change. A TE Code is added when a first time generic for an innovator is approved.

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

CHRS Change. Reference Standard

CTEC Change. Therapeutic Equivalence Code

CTNA Change. Trade Name

DISC Discontinued. The Rx or OTC listed product is not being marketed and will appear in the discontinued section in the next edition.

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

>D>	AA	HIKMA PHARMS	325MG; 50MG; 40MG	A089718	001	Jun 12, 1995	Feb	DISC
>A>		@	325MG; 50MG; 40MG	A089718	001	Jun 12, 1995	Feb	DISC

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

>D>	AA	ACTAVIS LABS FL INC	300MG; 5MG	A206470	001	Jun 02, 2016	Feb	DISC
>A>		@	300MG; 5MG	A206470	001	Jun 02, 2016	Feb	DISC
>D>	AA		300MG; 7.5MG	A206470	002	Jun 02, 2016	Feb	DISC
>A>		@	300MG; 7.5MG	A206470	002	Jun 02, 2016	Feb	DISC
>D>	AA		300MG; 10MG	A206470	003	Jun 02, 2016	Feb	DISC
>A>		@	300MG; 10MG	A206470	003	Jun 02, 2016	Feb	DISC
>D>	AA	XIROMED	325MG; 5MG	A211690	001	Feb 07, 2020	Feb	CAHN
>A>	AA		325MG; 5MG	A211690	001	Feb 07, 2020	Feb	CAHN
	AA		325MG; 5MG	A211690	001	Feb 07, 2020	Jan	NEWA
>D>	AA		325MG; 7.5MG	A211690	002	Feb 07, 2020	Feb	CAHN
>A>	AA		325MG; 7.5MG	A211690	002	Feb 07, 2020	Feb	CAHN
	AA		325MG; 7.5MG	A211690	002	Feb 07, 2020	Jan	NEWA
>D>	AA		325MG; 10MG	A211690	003	Feb 07, 2020	Feb	CAHN
>A>	AA		325MG; 10MG	A211690	003	Feb 07, 2020	Feb	CAHN
	AA		325MG; 10MG	A211690	003	Feb 07, 2020	Jan	NEWA

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

SOLUTION; ORAL

OXYCODONE AND ACETAMINOPHEN

>D>		@ MIKART INC	300MG/5ML; 10MG/5ML	A202142	001	Nov 27, 2018	Feb	CMFD
>A>			300MG/5ML; 10MG/5ML	A202142	001	Nov 27, 2018	Feb	CMFD

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

>D>	AB	APOTEX	325MG; 37.5MG	A078778	001	Apr 07, 2014	Feb	DISC
>A>		@	325MG; 37.5MG	A078778	001	Apr 07, 2014	Feb	DISC

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

>D>	AP	! X GEN PHARMS	EQ 500MG BASE/VIAL	A040784	001	Dec 10, 2008	Feb	CAHN
>A>	AP	! XGEN PHARMS	EQ 500MG BASE/VIAL	A040784	001	Dec 10, 2008	Feb	CAHN

ACYCLOVIR

CREAM; TOPICAL

ACYCLOVIR

AB	!	PERRIGO UK FINCO	5%	A208702	001	Feb 04, 2019	Jan	CHRS
----	---	------------------	----	---------	-----	--------------	-----	------

AB	+	BAUSCH	5%	N021478	001	Dec 30, 2002	Jan	CHRS
----	---	--------	----	---------	-----	--------------	-----	------

OINTMENT; TOPICAL

ACYCLOVIR

AB		XIROMED	5%	A201501	001	Jan 29, 2020	Jan	NEWA
----	--	---------	----	---------	-----	--------------	-----	------

TABLET; ORAL

ACYCLOVIR

AB	!	HETERO LABS LTD V	800MG	A203834	002	Oct 29, 2013	Jan	CHRS
----	---	-------------------	-------	---------	-----	--------------	-----	------

ZOVIRAX

+ @ MYLAN

+ @

400MG	N020089	001	Apr 30, 1991	Jan	DISC
800MG	N020089	002	Apr 30, 1991	Jan	DISC

ALBUTEROL SULFATE

AEROSOL, METERED; INHALATION

ALBUTEROL SULFATE

>A>	AB	PERRIGO PHARMS CO	EQ 0.09MG BASE/INH	A203760	001	Feb 24, 2020	Feb	NFTG
-----	----	-------------------	--------------------	---------	-----	--------------	-----	------

PROAIR HFA

>D>	BX	+! TEVA BRANDED PHARM	EQ 0.09MG BASE/INH	N021457	001	Oct 29, 2004	Feb	CFTG
-----	----	-----------------------	--------------------	---------	-----	--------------	-----	------

>A>	AB	+!	EQ 0.09MG BASE/INH	N021457	001	Oct 29, 2004	Feb	CFTG
-----	----	----	--------------------	---------	-----	--------------	-----	------

SOLUTION; INHALATION

ALBUTEROL SULFATE

>D>	AN	! BAUSCH AND LOMB	EQ 0.5% BASE	A075050	001	Jun 18, 1998	Feb	DISC
-----	----	-------------------	--------------	---------	-----	--------------	-----	------

SOLUTION; INHALATION  
ALBUTEROL SULFATE

>A>	@		EQ 0.5% BASE	A075050	001	Jun 18, 1998	Feb DISC
>D>	AN	NEPHRON	EQ 0.5% BASE	A075664	001	Jun 26, 2001	Feb CHRS
>A>	AN	!	EQ 0.5% BASE	A075664	001	Jun 26, 2001	Feb CHRS

ALPRAZOLAM

TABLET; ORAL  
ALPRAZOLAM

>D>	AB	DAVA INTL INC	0.25MG	A074174	001	Oct 19, 1993	Feb DISC
>A>	@		0.25MG	A074174	001	Oct 19, 1993	Feb DISC
>D>	AB		0.5MG	A074174	002	Oct 19, 1993	Feb DISC
>A>	@		0.5MG	A074174	002	Oct 19, 1993	Feb DISC
>D>	AB		1MG	A074174	003	Oct 19, 1993	Feb DISC
>A>	@		1MG	A074174	003	Oct 19, 1993	Feb DISC
>D>	AB		2MG	A074174	004	Oct 19, 1993	Feb DISC
>A>	@		2MG	A074174	004	Oct 19, 1993	Feb DISC

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL  
AMANTADINE HYDROCHLORIDE

>A>	AB	INVATECH	100MG	A210129	001	Mar 02, 2020	Feb NEWA
>A>	AB	INVATECH	100MG	A210215	001	Mar 10, 2020	Feb NEWA

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

EMULSION; INTRAVENOUS  
PERIKABIVEN IN PLASTIC CONTAINER

+!	FRESENIUS KABI USA	2.4%;20MG/100ML;6.8GM/100ML;68MG/100ML;124MG/100ML;170MG/100ML;105MG/100ML;3.5GM/100ML (2400ML)	N200656	003	Aug 25, 2014	Jan CHRS
----	--------------------	---	---------	-----	--------------	----------

AMINOCAPROIC ACID

SYRUP; ORAL  
AMINOCAPROIC ACID

>A>	AA	VISTAPHARM	1.25GM/5ML	A212814	001	Feb 26, 2020	Feb NEWA
-----	----	------------	------------	---------	-----	--------------	----------

AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION  
AMIODARONE HYDROCHLORIDE

>D>	AP	! HOSPIRA	50MG/ML	A075955	001	Oct 18, 2002	Feb DISC
>A>	@		50MG/ML	A075955	001	Oct 18, 2002	Feb DISC

>A> AMISULPRIDE

SOLUTION; INTRAVENOUS  
BARHEMSYS

>A>	+!	ACACIA PHARMA LTD	5MG/2ML (2.5MG/ML)	N209510	001	Feb 26, 2020	Feb NEWA
-----	----	-------------------	--------------------	---------	-----	--------------	----------

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL  
AMITRIPTYLINE HYDROCHLORIDE

AB	SANDOZ	10MG	A085968	004		Jan CMS1
AB	!	25MG	A085968	002		Jan CMS1
AB		75MG	A085968	006		Jan CMS1
AB		100MG	A085968	003		Jan CMS1
AB		150MG	A085968	005		Jan CMS1

AMLODIPINE BESYLATE

TABLET; ORAL  
AMLODIPINE BESYLATE

>D>	AB	HIKMA	EQ 2.5MG BASE	A077262	001	Jul 09, 2007	Feb DISC
>A>	@		EQ 2.5MG BASE	A077262	001	Jul 09, 2007	Feb DISC
>D>	AB		EQ 5MG BASE	A077262	002	Jul 09, 2007	Feb DISC
>A>	@		EQ 5MG BASE	A077262	002	Jul 09, 2007	Feb DISC
>D>	AB		EQ 10MG BASE	A077262	003	Jul 09, 2007	Feb DISC
>A>	@		EQ 10MG BASE	A077262	003	Jul 09, 2007	Feb DISC

AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM

TABLET;ORAL

AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM

>D>	@	ZYDUS PHARMS	EQ 2.5MG BASE;EQ 10MG BASE	A207762	001	Jan 11, 2019	Feb CMFD
>A>	AB		EQ 2.5MG BASE;EQ 10MG BASE	A207762	001	Jan 11, 2019	Feb CMFD
>D>	@		EQ 2.5MG BASE;EQ 20MG BASE	A207762	002	Jan 11, 2019	Feb CMFD
>A>	AB		EQ 2.5MG BASE;EQ 20MG BASE	A207762	002	Jan 11, 2019	Feb CMFD
>D>	@		EQ 2.5MG BASE;EQ 40MG BASE	A207762	003	Jan 11, 2019	Feb CMFD
>A>	AB		EQ 2.5MG BASE;EQ 40MG BASE	A207762	003	Jan 11, 2019	Feb CMFD
>D>	@		EQ 5MG BASE;EQ 10MG BASE	A207762	004	Jan 11, 2019	Feb CMFD
>A>	AB		EQ 5MG BASE;EQ 10MG BASE	A207762	004	Jan 11, 2019	Feb CMFD
>D>	@		EQ 5MG BASE;EQ 20MG BASE	A207762	005	Jan 11, 2019	Feb CMFD
>A>	AB		EQ 5MG BASE;EQ 20MG BASE	A207762	005	Jan 11, 2019	Feb CMFD
>D>	@		EQ 5MG BASE;EQ 40MG BASE	A207762	006	Jan 11, 2019	Feb CMFD
>A>	AB		EQ 5MG BASE;EQ 40MG BASE	A207762	006	Jan 11, 2019	Feb CMFD
>D>	@		EQ 5MG BASE;EQ 80MG BASE	A207762	007	Jan 11, 2019	Feb CMFD
>A>	AB		EQ 5MG BASE;EQ 80MG BASE	A207762	007	Jan 11, 2019	Feb CMFD
>D>	@		EQ 10MG BASE;EQ 10MG BASE	A207762	008	Jan 11, 2019	Feb CMFD
>A>	AB		EQ 10MG BASE;EQ 10MG BASE	A207762	008	Jan 11, 2019	Feb CMFD
>D>	@		EQ 10MG BASE;EQ 20MG BASE	A207762	009	Jan 11, 2019	Feb CMFD
>A>	AB		EQ 10MG BASE;EQ 20MG BASE	A207762	009	Jan 11, 2019	Feb CMFD
>D>	@		EQ 10MG BASE;EQ 40MG BASE	A207762	010	Jan 11, 2019	Feb CMFD
>A>	AB		EQ 10MG BASE;EQ 40MG BASE	A207762	010	Jan 11, 2019	Feb CMFD
>D>	@		EQ 10MG BASE;EQ 80MG BASE	A207762	011	Jan 11, 2019	Feb CMFD
>A>	AB		EQ 10MG BASE;EQ 80MG BASE	A207762	011	Jan 11, 2019	Feb CMFD

AMOXAPINE

TABLET;ORAL

AMOXAPINE

@	WATSON PHARMS TEVA	25MG	A072418	001	Aug 01, 1989	Jan CMS1
@		50MG	A072419	001	Aug 01, 1989	Jan CMS1

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB	ANI PHARMS INC	1.25MG;1.25MG;1.25MG;1.25MG	A205401	001	Jan 22, 2019	Jan CAHN
AB		2.5MG;2.5MG;2.5MG;2.5MG	A205401	002	Jan 22, 2019	Jan CAHN
AB		3.75MG;3.75MG;3.75MG;3.75MG	A205401	003	Jan 22, 2019	Jan CAHN
AB		5MG;5MG;5MG;5MG	A205401	004	Jan 22, 2019	Jan CAHN
AB		6.25MG;6.25MG;6.25MG;6.25MG	A205401	005	Jan 22, 2019	Jan CAHN
AB		7.5MG;7.5MG;7.5MG;7.5MG	A205401	006	Jan 22, 2019	Jan CAHN
	@ NESHER PHARMS	1.25MG;1.25MG;1.25MG;1.25MG	A210080	001	Jul 17, 2019	Jan DISC
	@	2.5MG;2.5MG;2.5MG;2.5MG	A210080	002	Jul 17, 2019	Jan DISC
	@	3.75MG;3.75MG;3.75MG;3.75MG	A210080	003	Jul 17, 2019	Jan DISC
	@	5MG;5MG;5MG;5MG	A210080	004	Jul 17, 2019	Jan DISC
	@	6.25MG;6.25MG;6.25MG;6.25MG	A210080	005	Jul 17, 2019	Jan DISC
	@	7.5MG;7.5MG;7.5MG;7.5MG	A210080	006	Jul 17, 2019	Jan DISC

AMPHETAMINE SULFATE

TABLET;ORAL

AMPHETAMINE SULFATE

AA	CEROVENE INC	5MG	A212582	001	Feb 04, 2020	Jan NEWA	
AA		10MG	A212582	002	Feb 04, 2020	Jan NEWA	
>A>	AA	PRINSTON INC	5MG	A211861	001	Mar 11, 2020	Feb NEWA
>A>	AA		10MG	A211861	002	Mar 11, 2020	Feb NEWA

AMPHOTERICIN B

INJECTABLE;INJECTION

AMPHOTERICIN B

>D>	!	X GEN PHARMS	50MG/VIAL	A063206	001	Apr 29, 1992	Feb CAHN
>A>	!	XGEN PHARMS	50MG/VIAL	A063206	001	Apr 29, 1992	Feb CAHN

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE;INJECTION

AMPICILLIN AND SULBACTAM

@	PHARM ASSOC	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A065316	001	Jun 29, 2007	Jan CAHN
@		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A065316	002	Jun 29, 2007	Jan CAHN

ANASTROZOLE

TABLET; ORAL

ANASTROZOLE

>D>	AB	APOTEX INC	1MG	A200654	001	May 11, 2012	Feb	DISC
>A>		@	1MG	A200654	001	May 11, 2012	Feb	DISC
		@ MYLAN	1MG	A091051	001	Jun 28, 2010	Jan	DISC

APIXABAN

TABLET; ORAL

APIXABAN

		@ MYLAN	2.5MG	A210128	001	Dec 23, 2019	Jan	DISC
		@	5MG	A210128	002	Dec 23, 2019	Jan	DISC

ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

SEPTOCAINE

>D>	AP	+!	DEPROCO	4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML)	N020971	001	Apr 03, 2000	Feb	CTEC
>A>		+!		4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML)	N020971	001	Apr 03, 2000	Feb	CTEC
>D>			ULTACAN FORTE						
>D>	AP		HANSAMED INC	4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML)	A201750	001	Jul 11, 2017	Feb	DISC
>A>		@		4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML)	A201750	001	Jul 11, 2017	Feb	DISC

ASENAPINE MALEATE

TABLET; SUBLINGUAL

SAPHRIS

		+	ALLERGAN	EQ 2.5MG BASE	N022117	003	Mar 12, 2015	Jan	CMS1
--	--	---	----------	---------------	---------	-----	--------------	-----	------

ATAZANAVIR SULFATE

CAPSULE; ORAL

ATAZANAVIR SULFATE

>D>	AB		MYLAN	EQ 150MG BASE	A208177	001	Sep 24, 2018	Feb	DISC
>A>		@		EQ 150MG BASE	A208177	001	Sep 24, 2018	Feb	DISC
>D>	AB			EQ 200MG BASE	A208177	002	Sep 24, 2018	Feb	DISC
>A>		@		EQ 200MG BASE	A208177	002	Sep 24, 2018	Feb	DISC
>D>	AB			EQ 300MG BASE	A208177	003	Sep 24, 2018	Feb	DISC
>A>		@		EQ 300MG BASE	A208177	003	Sep 24, 2018	Feb	DISC

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

AB			LUPIN LTD	EQ 10MG BASE	A204991	001	Mar 06, 2019	Jan	CMFD
AB				EQ 20MG BASE	A204991	002	Mar 06, 2019	Jan	CMFD
AB				EQ 40MG BASE	A204991	003	Mar 06, 2019	Jan	CMFD
AB				EQ 80MG BASE	A204991	004	Mar 06, 2019	Jan	CMFD

ATROPINE SULFATE

SOLUTION/DROPS; OPHTHALMIC

ISOPTO ATROPINE

>D>			ALCON LABS INC	1%	N208151	001	Dec 01, 2016	Feb	CHRS
>A>		!		1%	N208151	001	Dec 01, 2016	Feb	CHRS
>A>		!+		1%	N208151	001	Dec 01, 2016	Feb	CRLD

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

TABLET; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

>A>	AA		LEADING PHARMA LLC	0.025MG; 2.5MG	A213413	001	Feb 20, 2020	Feb	NEWA
-----	----	--	--------------------	----------------	---------	-----	--------------	-----	------

AVAPRITINIB

TABLET; ORAL

AYVAKIT

		+	BLUEPRINT MEDICINES	100MG	N212608	001	Jan 09, 2020	Jan	NEWA
		+		200MG	N212608	002	Jan 09, 2020	Jan	NEWA
		+!		300MG	N212608	003	Jan 09, 2020	Jan	NEWA

AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC  
AZELASTINE HYDROCHLORIDE

>A> AT GLAND PHARMA LTD 0.05% A210092 001 Feb 25, 2020 Feb NEWA

AZITHROMYCIN

TABLET;ORAL  
AZITHROMYCIN

AB ALEMBIC PHARMS LTD EQ 250MG BASE A211791 001 Jan 28, 2020 Jan NEWA  
 AB EQ 500MG BASE A211792 001 Jan 28, 2020 Jan NEWA  
 AB EQ 600MG BASE A211793 001 Jan 27, 2020 Jan NEWA  
 @ MYLAN EQ 600MG BASE A065360 001 Jan 08, 2007 Jan DISC

BACLOFEN

TABLET;ORAL  
BACLOFEN  
@ MYLAN

20MG A077121 002 Jul 29, 2005 Jan DISC

>A> BEMPEDOIC ACID

>A> TABLET;ORAL  
>A> NEXLETOL

>A> +! ESPERION THERAPS INC 180MG N211616 001 Feb 21, 2020 Feb NEWA

>A> BEMPEDOIC ACID; EZETIMIBE

>A> TABLET;ORAL  
>A> NEXLIZET

>A> +! ESPERION THERAPS INC 180MG;10MG N211617 001 Feb 26, 2020 Feb NEWA

BENAZEPRIL HYDROCHLORIDE

TABLET;ORAL  
BENAZEPRIL HYDROCHLORIDE

>D> AB APOTEX 5MG A077128 001 Mar 08, 2006 Feb CAHN  
 >D> AB 10MG A077128 002 Mar 08, 2006 Feb CAHN  
 >D> AB 20MG A077128 003 Mar 08, 2006 Feb CAHN  
 >D> AB 40MG A077128 004 Mar 08, 2006 Feb CAHN  
 >A> AB COREPHARMA 5MG A077128 001 Mar 08, 2006 Feb CAHN  
 >A> AB 10MG A077128 002 Mar 08, 2006 Feb CAHN  
 >A> AB 20MG A077128 003 Mar 08, 2006 Feb CAHN  
 >A> AB 40MG A077128 004 Mar 08, 2006 Feb CAHN

BEPOTASTINE BESILATE

SOLUTION/DROPS;OPHTHALMIC  
BEPOTASTINE BESILATE

>D> MYLAN 1.5% A206220 001 Mar 18, 2019 Feb DISC  
 >A> @ 1.5% A206220 001 Mar 18, 2019 Feb DISC  
 >D> AT +! BAUSCH AND LOMB INC 1.5% N022288 001 Sep 08, 2009 Feb CTEC  
 >A> +! 1.5% N022288 001 Sep 08, 2009 Feb CTEC

BETAMETHASONE DIPROPIONATE

CREAM, AUGMENTED;TOPICAL  
BETAMETHASONE DIPROPIONATE

>D> AB GLENMARK GENERICS EQ 0.05% BASE A078930 001 Sep 23, 2008 Feb CHRS  
 >A> AB ! EQ 0.05% BASE A078930 001 Sep 23, 2008 Feb CHRS  
 >D> DIPROLENE AF  
 >D> AB +! MERCK SHARP DOHME EQ 0.05% BASE N019555 001 Apr 27, 1987 Feb DISC  
 >A> + @ EQ 0.05% BASE N019555 001 Apr 27, 1987 Feb DISC

BETHANECHOL CHLORIDE

TABLET;ORAL  
BETHANECHOL CHLORIDE

AA ! AMNEAL PHARM 5MG A040855 001 Nov 21, 2007 Jan CHRS  
 AA ! 10MG A040855 002 Nov 21, 2007 Jan CHRS  
 AA ! 25MG A040855 003 Nov 21, 2007 Jan CHRS  
 @ IMPAX LABS 25MG A040721 003 Nov 01, 2006 Jan CMS1  
 URECHOLINE  
 @ ODYSSEY PHARMS 5MG A089095 001 Dec 19, 1985 Jan DISC  
 @ 10MG A088440 001 May 29, 1984 Jan DISC  
 @ 25MG A088441 001 May 29, 1984 Jan DISC



BEXAROTENE

CAPSULE;ORAL  
 BEXAROTENE  
 AB ANI PHARMS INC 75MG A209861 001 May 08, 2018 Jan CAHN

BICALUTAMIDE

TABLET;ORAL  
 BICALUTAMIDE  
 @ MYLAN 50MG A079185 001 Jul 06, 2009 Jan DISC

BIMATOPROST

SOLUTION/DROPS;TOPICAL  
 BIMATOPROST  
 AT ALEMBIC PHARMS LTD 0.03% A210515 001 Jan 21, 2020 Jan NEWA

BIVALIRUDIN

INJECTABLE;INTRAVENOUS  
 BIVALIRUDIN  
 >D> AP APOTEX 250MG/VIAL A204876 001 Jul 06, 2017 Feb DISC  
 >A> @ 250MG/VIAL A204876 001 Jul 06, 2017 Feb DISC

BOSENTAN

TABLET;ORAL  
 BOSENTAN  
 AB ALEMBIC PHARMS LTD 62.5MG A211461 001 Jan 23, 2020 Jan NEWA  
 AB 125MG A211461 002 Jan 23, 2020 Jan NEWA  
 @ MYLAN 62.5MG A205173 001 Jan 15, 2020 Jan DISC  
 @ 125MG A205173 002 Jan 15, 2020 Jan DISC

BUDESONIDE

AEROSOL, FOAM;RECTAL  
 UCERIS  
 >A> +! SALIX 2MG/ACTUATION N205613 001 Oct 07, 2014 Feb CAHN  
 >D> +! VALEANT PHARMS INTL 2MG/ACTUATION N205613 001 Oct 07, 2014 Feb CAHN

BUMETANIDE

TABLET;ORAL  
 BUMETANIDE  
 AB RISING 0.5MG A212019 001 Dec 12, 2019 Jan CAHN  
 AB 1MG A212019 002 Dec 12, 2019 Jan CAHN  
 AB 2MG A212019 003 Dec 12, 2019 Jan CAHN

BUPROPION HYDROCHLORIDE

TABLET;ORAL  
 BUPROPION HYDROCHLORIDE  
 AB CADILA PHARMS LTD 75MG A208606 001 Jan 16, 2020 Jan NEWA  
 AB 100MG A208606 002 Jan 16, 2020 Jan NEWA  
 TABLET, EXTENDED RELEASE;ORAL  
 BUPROPION HYDROCHLORIDE  
 >D> AB1 JUBILANT GENERICS 100MG A202774 001 Oct 11, 2013 Feb DISC  
 >A> @ 100MG A202774 001 Oct 11, 2013 Feb DISC  
 >D> AB1 150MG A202774 002 Oct 11, 2013 Feb DISC  
 >A> @ 150MG A202774 002 Oct 11, 2013 Feb DISC  
 >D> AB2 150MG A202775 001 Oct 11, 2013 Feb DISC  
 >A> @ 150MG A202775 001 Oct 11, 2013 Feb DISC  
 >D> AB3 150MG A207459 001 Jun 30, 2017 Feb DISC  
 >A> @ 150MG A207459 001 Jun 30, 2017 Feb DISC  
 >D> AB1 200MG A202774 003 Oct 11, 2013 Feb DISC  
 >A> @ 200MG A202774 003 Oct 11, 2013 Feb DISC  
 >D> AB3 300MG A207459 002 Jun 30, 2017 Feb DISC  
 >A> @ 300MG A207459 002 Jun 30, 2017 Feb DISC

BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL  
 CONTRAVE  
 +! NALPROPION 90MG;8MG N200063 001 Sep 10, 2014 Jan CAHN

BUSULFAN

INJECTABLE; INJECTION  
BUSULFAN

>D>	AP	MYLAN LABS LTD	6MG/ML	A205184	001	Jul 13, 2018	Feb DISC
>A>		@	6MG/ML	A205184	001	Jul 13, 2018	Feb DISC

CABERGOLINE

TABLET; ORAL  
CABERGOLINE  
@ MYLAN

0.5MG

A202947 001 Dec 02, 2013 Jan DISC

CALCITRIOL

INJECTABLE; INJECTION  
CALCITRIOL

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

A078066 001 Jan 29, 2008 Jan CTEC  
A211030 001 Feb 03, 2020 Jan NEWA

CALCIUM ACETATE

CAPSULE; ORAL  
CALCIUM ACETATE

>A>	AB	@ LOTUS PHARM CO LTD	667MG	A203298	001	Jul 26, 2016	Jan DISC
		SUVEN LIFE	667MG	A211038	001	Feb 21, 2020	Feb NEWA

CAPTOPRIL

TABLET; ORAL  
CAPTOPRIL

>D>		@ APOTEX	12.5MG	A074737	001	Oct 28, 1998	Feb CAHN
>D>		@	25MG	A074737	002	Oct 28, 1998	Feb CAHN
>D>		@	50MG	A074737	003	Oct 28, 1998	Feb CAHN
>D>		@	100MG	A074737	004	Oct 28, 1998	Feb CAHN
>A>		@ COREPHARMA	12.5MG	A074737	001	Oct 28, 1998	Feb CAHN
>A>		@	25MG	A074737	002	Oct 28, 1998	Feb CAHN
>A>		@	50MG	A074737	003	Oct 28, 1998	Feb CAHN
>A>		@	100MG	A074737	004	Oct 28, 1998	Feb CAHN

CARBIDOPA

TABLET; ORAL  
CARBIDOPA

AB		ANI PHARMS INC	25MG	A203261	001	Mar 10, 2014	Jan CAHN
----	--	----------------	------	---------	-----	--------------	----------

CARBIDOPA; LEVODOPA

TABLET, EXTENDED RELEASE; ORAL  
SINEMET CR

+ @ MERCK SHARP DOHME 25MG; 100MG

N019856 002 Dec 24, 1992 Jan DISC

CARISOPRODOL

TABLET; ORAL  
CARISOPRODOL

>D>	AA	HIKMA INTL PHARMS	350MG	A040124	001	Jan 24, 1996	Feb DISC
>A>		@	350MG	A040124	001	Jan 24, 1996	Feb DISC

CEFIXIME

FOR SUSPENSION; ORAL  
SUPRAX

@ LUPIN PHARMS 100MG/5ML

A065129 001 Feb 23, 2004 Jan DISC

TABLET; ORAL  
SUPRAX

@ LUPIN PHARMS 400MG

A065130 001 Feb 12, 2004 Jan DISC

CEFTRIAZONE SODIUM

INJECTABLE; INJECTION  
CEFTRIAZONE

@ HOSPIRA INC EQ 10GM BASE/VIAL

A065232 001 Aug 02, 2005 Jan DISC

CELECOXIB

CAPSULE; ORAL  
CELECOXIB

>A>	AB	TIANJIN TIANYAO	50MG	A207872	001	Feb 25, 2020	Feb NEWA
>A>	AB		100MG	A207872	002	Feb 25, 2020	Feb NEWA
>A>	AB		200MG	A207872	003	Feb 25, 2020	Feb NEWA
>A>	AB		400MG	A207872	004	Feb 25, 2020	Feb NEWA

CAPSULE;ORAL

CELECOXIB

AB	UMEDICA LABS PVT LTD	50MG	A210628	001	Nov 27, 2019	Jan CDFR
AB		100MG	A210628	002	Nov 27, 2019	Jan CDFR
AB		200MG	A210628	003	Nov 27, 2019	Jan CDFR
AB		400MG	A210628	004	Nov 27, 2019	Jan CDFR
>A>	YILING PHARM LTD	50MG	A211412	001	Mar 06, 2020	Feb NEWA
>A>		100MG	A211412	002	Mar 06, 2020	Feb NEWA
>A>		200MG	A211412	003	Mar 06, 2020	Feb NEWA
>A>		400MG	A211412	004	Mar 06, 2020	Feb NEWA

CHLOROQUINE PHOSPHATE

TABLET;ORAL

CHLOROQUINE PHOSPHATE

>D>	!	HIKMA PHARMS	EQ 150MG BASE	A083082	001	Feb DISC
>A>	@		EQ 150MG BASE	A083082	001	Feb DISC
>D>	AA	NATCO PHARMA LTD	EQ 150MG BASE	A091621	001	Jan 21, 2011 Feb CHRS
>A>	AA	!	EQ 150MG BASE	A091621	001	Jan 21, 2011 Feb CHRS

CHLORPROMAZINE HYDROCHLORIDE

TABLET;ORAL

CHLORPROMAZINE HYDROCHLORIDE

AB	ZYDUS	10MG	A213368	001	Jan 17, 2020	Jan NEWA
AB		25MG	A213368	002	Jan 17, 2020	Jan NEWA
AB		50MG	A213368	003	Jan 17, 2020	Jan NEWA
AB		100MG	A213368	004	Jan 17, 2020	Jan NEWA
AB		200MG	A213368	005	Jan 17, 2020	Jan NEWA

CHLORTHALIDONE

TABLET;ORAL

CHLORTHALIDONE

AB	ALKEM LABS LTD	25MG	A213412	001	Feb 11, 2020	Jan NEWA
AB		50MG	A213412	002	Feb 11, 2020	Jan NEWA
>D>	AB	ATHEM	25MG	A211063	001	Feb 26, 2019 Feb CAHN
>D>	AB		50MG	A211063	002	Feb 26, 2019 Feb CAHN
>A>	AB	VISTAPHARM	25MG	A211063	001	Feb 26, 2019 Feb CAHN
>A>	AB		50MG	A211063	002	Feb 26, 2019 Feb CAHN

CHOLESTYRAMINE

POWDER;ORAL

LOCHOLEST

>A>	@	ALLIED	EQ 4GM RESIN/PACKET	A074561	001	Aug 15, 1996 Feb CAHN
>A>	@		EQ 4GM RESIN/SCOOPFUL	A074561	002	Aug 15, 1996 Feb CAHN
>D>	@	INVATECH	EQ 4GM RESIN/PACKET	A074561	001	Aug 15, 1996 Feb CAHN
>D>	@		EQ 4GM RESIN/PACKET	A074561	001	Aug 15, 1996 Jan CAHN
>D>	@		EQ 4GM RESIN/SCOOPFUL	A074561	002	Aug 15, 1996 Feb CAHN
>D>	@		EQ 4GM RESIN/SCOOPFUL	A074561	002	Aug 15, 1996 Jan CAHN
		LOCHOLEST LIGHT				
>A>	@	ALLIED	EQ 4GM RESIN/PACKET	A074562	001	Aug 15, 1996 Feb CAHN
>A>	@		EQ 4GM RESIN/SCOOPFUL	A074562	002	Aug 15, 1996 Feb CAHN
>D>	@	INVATECH	EQ 4GM RESIN/PACKET	A074562	001	Aug 15, 1996 Feb CAHN
>D>	@		EQ 4GM RESIN/SCOOPFUL	A074562	002	Aug 15, 1996 Feb CAHN

CINACALCET HYDROCHLORIDE

TABLET;ORAL

CINACALCET HYDROCHLORIDE

@ LUPIN LTD

@

@

EQ 30MG BASE	A210548	001	Jun 28, 2019	Jan DISC
EQ 60MG BASE	A210548	002	Jun 28, 2019	Jan DISC
EQ 90MG BASE	A210548	003	Jun 28, 2019	Jan DISC

CISATRACURIUM BESYLATE

INJECTABLE;INJECTION

CISATRACURIUM BESYLATE

AP	HOSPIRA INC	EQ 2MG BASE/ML	A203238	001	Mar 30, 2018	Jan CMFD
----	-------------	----------------	---------	-----	--------------	----------

CLINDAMYCIN PHOSPHATE

AEROSOL, FOAM;TOPICAL

CLINDAMYCIN PHOSPHATE

>A>	AT	TARO PHARM INDS LTD	1%	A210004	001	Mar 11, 2020	Feb NEWA
-----	----	---------------------	----	---------	-----	--------------	----------

CLOBAZAMSUSPENSION;ORAL  
CLOBAZAM

>A>	AB	HETERO LABS LTD III	2.5MG/ML	A209796	001	Feb 24, 2020	Feb NEWA
	AB	TEVA PHARMS USA	2.5MG/ML	A211032	001	Jan 31, 2020	Jan NEWA

CLOBETASOL PROPIONATE

CREAM;TOPICAL

CLOBETASOL PROPIONATE

AB1		ALEOR DERMACEUTICALS	0.05%	A213291	001	Jan 27, 2020	Jan NEWA
-----	--	----------------------	-------	---------	-----	--------------	----------

SPRAY;TOPICAL

CLOBETASOL PROPIONATE

@ ALEOR DERMACEUTICALS 0.05%

A211191	001	Oct 02, 2019	Jan DISC
---------	-----	--------------	----------

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE;ORAL

CLOMIPRAMINE HYDROCHLORIDE

AB		LEADING PHARMA LLC	25MG	A211364	001	Feb 07, 2020	Jan NEWA
AB			50MG	A211364	002	Feb 07, 2020	Jan NEWA
AB			75MG	A211364	003	Feb 07, 2020	Jan NEWA
AB		MANKIND PHARMA	25MG	A211767	001	Apr 08, 2019	Jan CMFD
AB			50MG	A211767	002	Apr 08, 2019	Jan CMFD
AB			75MG	A211767	003	Apr 08, 2019	Jan CMFD
>D>	AB	TEVA	25MG	A074958	001	Aug 26, 1997	Feb DISC
>A>		@	25MG	A074958	001	Aug 26, 1997	Feb DISC
>D>	AB		50MG	A074958	002	Aug 26, 1997	Feb DISC
>A>		@	50MG	A074958	002	Aug 26, 1997	Feb DISC
>D>	AB		75MG	A074958	003	Aug 26, 1997	Feb DISC
>A>		@	75MG	A074958	003	Aug 26, 1997	Feb DISC

CLONIDINE HYDROCHLORIDE

INJECTABLE;INJECTION

CLONIDINE HYDROCHLORIDE

>D>	AP	X-GEN PHARMS INC	1MG/10ML (0.1MG/ML)	A203167	001	Oct 29, 2013	Feb CAHN
>D>	AP		5MG/10ML (0.5MG/ML)	A203167	002	Oct 29, 2013	Feb CAHN
>A>	AP	XGEN PHARMS	1MG/10ML (0.1MG/ML)	A203167	001	Oct 29, 2013	Feb CAHN
>A>	AP		5MG/10ML (0.5MG/ML)	A203167	002	Oct 29, 2013	Feb CAHN

TABLET, EXTENDED RELEASE;ORAL

CLONIDINE HYDROCHLORIDE

>D>		@ UPSHER SMITH LABS	0.1MG	A211433	001	Oct 12, 2018	Feb CMFD
>A>	AB1		0.1MG	A211433	001	Oct 12, 2018	Feb CMFD

CLORAZEPATE DIPOTASSIUM

CAPSULE;ORAL

CLORAZEPATE DIPOTASSIUM

>D>		@ AUROLIFE PHARMA LLC	3.75MG	A072112	002	Aug 11, 2017	Feb CMS1
>A>		@	3.75MG	A072112	002	Aug 26, 1988	Feb CMS1
>D>		@	7.5MG	A072112	003	Aug 11, 2017	Feb CMS1
>A>		@	7.5MG	A072112	003	Aug 26, 1988	Feb CMS1

COCAINE HYDROCHLORIDE

SOLUTION;NASAL

NUMBRINO

CODY LABS INC 4%

N209575	001	Jan 10, 2020	Jan NEWA
---------	-----	--------------	----------

COLCHICINE

TABLET;ORAL

COLCHICINE

AB		AMNEAL PHARMS	0.6MG	A204711	001	Sep 28, 2016	Jan CMFD
AB		GRANULES PHARMS	0.6MG	A210425	001	Feb 05, 2020	Jan NEWA

COLISTIMETHATE SODIUM

INJECTABLE;INJECTION

COLISTIMETHATE SODIUM

>D>	AP	X GEN PHARMS	EQ 150MG BASE/VIAL	A064216	001	Feb 26, 1999	Feb CAHN
>A>	AP	XGEN PHARMS	EQ 150MG BASE/VIAL	A064216	001	Feb 26, 1999	Feb CAHN

CYCLOPHOSPHAMIDECAPSULE;ORAL  
CYCLOPHOSPHAMIDE

AB	ANI PHARMS INC	25MG	A207014	001	Mar 19, 2018	Jan CAHN
AB		50MG	A207014	002	Mar 19, 2018	Jan CAHN

CYSTEAMINE BITARTRATEGRANULE;ORAL  
PROCYSBI

>A>	+	HORIZON PHARMA USA	EQ 75MG BASE/PACKET	N213491	001	Feb 14, 2020	Feb NEWA
>A>	+	!	EQ 300MG BASE/PACKET	N213491	002	Feb 14, 2020	Feb NEWA

DABIGATRAN ETEXILATE MESYLATECAPSULE;ORAL  
DABIGATRAN ETEXILATE MESYLATE

>A>	AB	ALKEM LABS LTD	EQ 75MG BASE	A208040	001	Mar 11, 2020	Feb NFTG
>A>	AB		EQ 150MG BASE	A208040	002	Mar 11, 2020	Feb NFTG
		PRADAXA					
>D>	+	BOEHRINGER INGELHEIM	EQ 75MG BASE	N022512	001	Oct 19, 2010	Feb CFTG
>A>	AB	+	EQ 75MG BASE	N022512	001	Oct 19, 2010	Feb CFTG
>D>	+	!	EQ 150MG BASE	N022512	002	Oct 19, 2010	Feb CFTG
>A>	AB	+	EQ 150MG BASE	N022512	002	Oct 19, 2010	Feb CFTG

DACTINOMYCININJECTABLE;INJECTION  
DACTINOMYCIN

>D>	AP	X-GEN PHARMS INC	0.5MG/VIAL	A203999	001	May 20, 2019	Feb CAHN
>A>	AP	XGEN PHARMS	0.5MG/VIAL	A203999	001	May 20, 2019	Feb CAHN

DAPSONEGEL;TOPICAL  
DAPSONE

>A>	AB	TARO PHARMS	7.5%	A210191	001	Jun 26, 2019	Feb NEWA
-----	----	-------------	------	---------	-----	--------------	----------

DARIFENACIN HYDROBROMIDETABLET, EXTENDED RELEASE;ORAL  
DARIFENACIN HYDROBROMIDE

>D>	AB	ANCHEN PHARMS	EQ 7.5MG BASE	A091190	001	Mar 13, 2015	Feb DISC
>A>	@		EQ 7.5MG BASE	A091190	001	Mar 13, 2015	Feb DISC
>D>	AB		EQ 15MG BASE	A091190	002	Mar 13, 2015	Feb DISC
>A>	@		EQ 15MG BASE	A091190	002	Mar 13, 2015	Feb DISC

DEFERASIROXTABLET;ORAL  
DEFERASIROX

>A>	AB	AMNEAL	90MG	A210727	001	Dec 27, 2019	Feb CAHN
>A>	AB		360MG	A210727	002	Dec 27, 2019	Feb CAHN
>D>	AB	AMNEAL PHARMS CO	90MG	A210727	001	Dec 27, 2019	Feb CAHN
>D>	AB		360MG	A210727	002	Dec 27, 2019	Feb CAHN
	AB	CIPLA	90MG	A211852	001	Feb 11, 2020	Jan NEWA
	AB		360MG	A211852	002	Feb 11, 2020	Jan NEWA

DESIRUDIN RECOMBINANTINJECTABLE;SUBCUTANEOUS  
IPRIVASK

	+	@ BAUSCH	15MG/VIAL	N021271	001	Apr 04, 2003	Jan CAHN
--	---	----------	-----------	---------	-----	--------------	----------

DESMOPRESSIN ACETATEINJECTABLE;INJECTION  
DESMOPRESSIN ACETATE

>A>	AP	UBI	0.004MG/ML	A210218	001	Feb 14, 2020	Feb NEWA
-----	----	-----	------------	---------	-----	--------------	----------

DESOXIMETASONECREAM;TOPICAL  
DESOXIMETASONE

>A>	AB	COSETTE	0.25%	A209595	001	Mar 04, 2020	Feb NEWA
-----	----	---------	-------	---------	-----	--------------	----------

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE;ORAL  
DESVENLAFAXINE SUCCINATE

>D>	AB	MYLAN	EQ 50MG BASE	A204095	001	Jun 29, 2015	Feb DISC
>A>		@	EQ 50MG BASE	A204095	001	Jun 29, 2015	Feb DISC

DEXAMETHASONE; TOBRAMYCIN

SUSPENSION/DROPS;OPHTHALMIC  
TOBRADEX ST

>A>	+	EYEVANCE	0.05%;0.3%	N050818	001	Feb 13, 2009	Feb CAHN
>D>	+	NOVARTIS	0.05%;0.3%	N050818	001	Feb 13, 2009	Feb CAHN

DEXMEDETOMIDINE HYDROCHLORIDE

INJECTABLE;INJECTION  
DEXMEDETOMIDINE

AP		JIANGSU HENGRUI MED	EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)	A209065	001	Sep 19, 2017	Jan CTNA
AP		HIKMA	EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)	A206407	001	Jan 30, 2020	Jan NEWA
AP			EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)	A206407	002	Jan 30, 2020	Jan NEWA

DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL  
DEXTROAMPHETAMINE SULFATE

	@	PII	5MG	A205077	001	Jun 21, 2019	Jan DISC
	@		10MG	A205077	002	Jun 21, 2019	Jan DISC
	@		15MG	A205077	003	Jun 21, 2019	Jan DISC

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL  
PROMETHAZINE DM

>A>		@ SLATE	15MG/5ML;6.25MG/5ML	A040649	001	Feb 14, 2006	Feb CAHN
>D>		@ VINTAGE	15MG/5ML;6.25MG/5ML	A040649	001	Feb 14, 2006	Feb CAHN

DIAZEPAM

SPRAY;NASAL  
VALTOCO

	+	NEURELIS INC	5MG/SPRAY	N211635	001	Jan 10, 2020	Jan NEWA
	+		7.5MG/SPRAY	N211635	002	Jan 10, 2020	Jan NEWA
	+		10MG/SPRAY	N211635	003	Jan 10, 2020	Jan NEWA

DICLOFENAC EPOLAMINE

>D>			PATCH;TOPICAL				
>D>			FLECTOR				
>D>	+	INST BIOCHEM	1.3%	N021234	001	Jan 31, 2007	Feb CDFR
>A>			SYSTEM;TOPICAL				
>A>			FLECTOR				
>A>	+	INST BIOCHEM	1.3%	N021234	001	Jan 31, 2007	Feb CDFR

DICLOFENAC POTASSIUM

TABLET;ORAL  
CATAFLAM

	+	@ NOVARTIS	50MG	N020142	002	Nov 24, 1993	Jan CRLD
--	---	------------	------	---------	-----	--------------	----------

DICLOFENAC SODIUM

GEL;TOPICAL  
DICLOFENAC SODIUM

AB		ENCUBE	1%	A210986	001	Jan 27, 2020	Jan NEWA
----	--	--------	----	---------	-----	--------------	----------

DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE;ORAL  
DICLOFENAC SODIUM AND MISOPROSTOL

>A>	AB	YUNG SHIN PHARM	50MG;0.2MG	A205143	001	Feb 19, 2020	Feb NEWA
>A>	AB		75MG;0.2MG	A205143	002	Feb 19, 2020	Feb NEWA

DIHYDROERGOTAMINE MESYLATE

SPRAY, METERED;NASAL

>A>		DIHYDROERGOTAMINE MESYLATE						
>A>	AB	CUSTOPHARM INC	0.5MG/SPRAY	A211393	001	Feb 28, 2020	Feb	NFTG
		MIGRANAL						
>D>		+! BAUSCH	0.5MG/SPRAY	N020148	001	Dec 08, 1997	Feb	CFTG
>A>	AB	+!	0.5MG/SPRAY	N020148	001	Dec 08, 1997	Feb	CFTG

DISULFIRAM

TABLET;ORAL  
DISULFIRAM

>D>	AB	HIKMA	250MG	A202652	001	Feb 05, 2014	Feb	DISC
>A>		@	250MG	A202652	001	Feb 05, 2014	Feb	DISC
>D>	AB		500MG	A202652	002	Feb 05, 2014	Feb	DISC
>A>		@	500MG	A202652	002	Feb 05, 2014	Feb	DISC

DIVALPROEX SODIUM

CAPSULE, DELAYED REL PELLETS;ORAL  
DIVALPROEX SODIUM

>A>	AB	AJANTA PHARMA LTD	EQ 125MG VALPROIC ACID	A213181	001	Mar 02, 2020	Feb	NEWA
		@ MYLAN	EQ 125MG VALPROIC ACID	A090407	001	Mar 28, 2011	Jan	DISC
		TABLET, DELAYED RELEASE;ORAL						
		DIVALPROEX SODIUM						
		@ MYLAN	EQ 125MG VALPROIC ACID	A090062	001	Mar 17, 2009	Jan	DISC
		@	EQ 250MG VALPROIC ACID	A090062	002	Mar 17, 2009	Jan	DISC
		@	EQ 500MG VALPROIC ACID	A090062	003	Mar 17, 2009	Jan	DISC

DOCETAXEL

INJECTABLE;INJECTION  
DOCETAXEL

AP		SHILPA MEDICARE LTD	20MG/ML (20MG/ML)	A210327	001	May 16, 2019	Jan	CMFD
AP			80MG/4ML (20MG/ML)	A210327	002	May 16, 2019	Jan	CMFD
AP			160MG/8ML (20MG/ML)	A210327	003	May 16, 2019	Jan	CMFD

DOFETILIDE

CAPSULE;ORAL  
DOFETILIDE

AB		MSN	0.125MG	A213220	001	Jan 29, 2020	Jan	NEWA
AB			0.25MG	A213220	002	Jan 29, 2020	Jan	NEWA
AB			0.5MG	A213220	003	Jan 29, 2020	Jan	NEWA

DOXAZOSIN MESYLATE

TABLET;ORAL  
DOXAZOSIN MESYLATE

		@ ANI PHARMS INC	EQ 1MG BASE	A075432	001	Oct 18, 2000	Jan	DISC
		@	EQ 2MG BASE	A075432	002	Oct 18, 2000	Jan	DISC
		@	EQ 4MG BASE	A075432	003	Oct 18, 2000	Jan	DISC
		@	EQ 8MG BASE	A075432	004	Oct 18, 2000	Jan	DISC

DOXERCALCIFEROL

INJECTABLE;INJECTION  
DOXERCALCIFEROL

>A>	AP	MEITHEAL	4MCG/2ML (2MCG/ML)	A211670	001	Feb 07, 2020	Feb	CAHN
>D>	AP	NANJING KING-FRIEND	4MCG/2ML (2MCG/ML)	A211670	001	Feb 07, 2020	Feb	CAHN
	AP		4MCG/2ML (2MCG/ML)	A211670	001	Feb 07, 2020	Jan	NEWA

DOXYCYCLINE

CAPSULE;ORAL  
DOXYCYCLINE

		@ MYLAN PHARMS INC	EQ 150MG BASE	A202778	001	Jun 08, 2012	Jan	DISC
--	--	--------------------	---------------	---------	-----	--------------	-----	------

DOXYCYCLINE HYCLATE

TABLET;ORAL  
DOXYCYCLINE HYCLATE

>A>	AB	ALEMBIC PHARMS LTD	EQ 20MG BASE	A210537	001	Mar 03, 2020	Feb	NEWA
		TABLET, DELAYED RELEASE;ORAL						
		DOXYCYCLINE HYCLATE						
>A>	AB	PRINSTON INC	EQ 50MG BASE	A207494	003	Feb 19, 2019	Feb	NEWA

DRONABINOL

		CAPSULE;ORAL							
		DRONABINOL							
AB		ASCENT PHARMS INC	2.5MG	A207421	001	Feb 07, 2020	Jan	NEWA	
AB			5MG	A207421	002	Feb 07, 2020	Jan	NEWA	
AB			10MG	A207421	003	Feb 07, 2020	Jan	NEWA	
		MARINOL							
AB	+	ALKEM LABS LTD	2.5MG	N018651	001	May 31, 1985	Jan	CAHN	
AB	+	!	5MG	N018651	002	May 31, 1985	Jan	CAHN	
AB	+		10MG	N018651	003	May 31, 1985	Jan	CAHN	
		SOLUTION;ORAL							
		SYNDROS							
>A>	+	BENUVIA	5MG/ML	N205525	001	Mar 23, 2017	Feb	CAHN	
>D>	+	INSYS DEV CO INC	5MG/ML	N205525	001	Mar 23, 2017	Feb	CAHN	

DROSPIRENONE; ETHINYL ESTRADIOL

		TABLET;ORAL							
		DROSPIRENONE AND ETHINYL ESTRADIOL							
AB		ALLIED	3MG;0.02MG	A203291	001	Jul 18, 2017	Jan	CAHN	
		TABLET;ORAL-28							
		DROSPIRENONE AND ETHINYL ESTRADIOL							
AB		HETERO LABS LTD	3MG;0.03MG	A213034	001	Jan 24, 2020	Jan	NEWA	

DUTASTERIDE

		CAPSULE;ORAL							
		DUTASTERIDE							
>D>	AB	@	ACTAVIS LABS FL INC	0.5MG	A202808	001	Nov 20, 2015	Jan	DISC
>A>	AB		APOTEX INC	0.5MG	A204292	001	Nov 24, 2015	Feb	DISC
>A>		@		0.5MG	A204292	001	Nov 24, 2015	Feb	DISC
>D>	AB		HIKMA	0.5MG	A202204	001	Nov 23, 2015	Feb	DISC
>A>		@		0.5MG	A202204	001	Nov 23, 2015	Feb	DISC

EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE

		TABLET, EXTENDED RELEASE;ORAL							
		TRIJARDY XR							
	+	BOEHRINGER INGELHEIM	5MG;2.5MG;1GM	N212614	001	Jan 27, 2020	Jan	NEWA	
	+		10MG;5MG;1GM	N212614	002	Jan 27, 2020	Jan	NEWA	
	+		12.5MG;2.5MG;1GM	N212614	003	Jan 27, 2020	Jan	NEWA	
	+	!	25MG;5MG;1GM	N212614	004	Jan 27, 2020	Jan	NEWA	

EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

		TABLET;ORAL							
		EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE							
>D>	AB		MYLAN	200MG;300MG	A206436	001	Apr 09, 2018	Feb	DISC
>A>		@		200MG;300MG	A206436	001	Apr 09, 2018	Feb	DISC
>A>	AB		ZYDUS PHARMS	200MG;300MG	A212689	001	Feb 28, 2020	Feb	NEWA

ENTECAVIR

		TABLET;ORAL							
		ENTECAVIR							
	@	MYLAN	0.5MG	A206226	001	Mar 26, 2019	Jan	DISC	
	@		1MG	A206226	002	Mar 26, 2019	Jan	DISC	

EPLERENONE

		TABLET;ORAL							
		EPLERENONE							
>D>	AB		APOTEX	25MG	A078482	001	Jul 30, 2008	Feb	CAHN
>D>	AB			50MG	A078482	002	Jul 30, 2008	Feb	CAHN
>A>	AB		COREPHARMA	25MG	A078482	001	Jul 30, 2008	Feb	CAHN
>A>	AB			50MG	A078482	002	Jul 30, 2008	Feb	CAHN

ESCITALOPRAM OXALATE

		SOLUTION;ORAL							
		ESCITALOPRAM OXALATE							
AA		ANTRIM PHARMS LLC	EQ 5MG BASE/5ML	A203967	001	May 26, 2015	Jan	CAHN	



ESOMEPRAZOLE MAGNESIUM; NAPROXEN

TABLET, DELAYED RELEASE;ORAL  
 NAPROXEN AND ESOMEPRAZOLE MAGNESIUM

>A>	AB	DR REDDYS LABS LTD	EQ 20MG BASE;375MG	A204206	001	Feb 18, 2020	Feb NEWA
>A>	AB		EQ 20MG BASE;500MG	A204206	002	Feb 18, 2020	Feb NEWA
		VIMOVO					
>D>	+	HORIZON	EQ 20MG BASE;375MG	N022511	002	Apr 30, 2010	Feb CTEC
>A>	AB	+	EQ 20MG BASE;375MG	N022511	002	Apr 30, 2010	Feb CTEC
>D>		+	EQ 20MG BASE;500MG	N022511	001	Apr 30, 2010	Feb CTEC
>A>	AB	+	EQ 20MG BASE;500MG	N022511	001	Apr 30, 2010	Feb CTEC

ESTRADIOL; NORETHINDRONE ACETATE

TABLET;ORAL  
 ESTRADIOL AND NORETHINDRONE ACETATE  
 @ TEVA PHARMS USA 0.5MG;0.1MG

A200747	001	Mar 08, 2012	Jan DISC
---------	-----	--------------	----------

ESTROPIPATE

TABLET;ORAL  
 ESTROPIPATE  
 @ MYLAN 0.75MG  
 @ 1.5MG

A040359	001	Aug 26, 1999	Jan DISC
A040359	002	Aug 26, 1999	Jan DISC

ESZOPICLONE

TABLET;ORAL  
 ESZOPICLONE

>D>	AB	HIKMA	1MG	A091153	001	Apr 15, 2014	Feb DISC
>A>		@	1MG	A091153	001	Apr 15, 2014	Feb DISC
>D>	AB		2MG	A091153	002	Apr 15, 2014	Feb DISC
>A>		@	2MG	A091153	002	Apr 15, 2014	Feb DISC
>D>	AB		3MG	A091153	003	Apr 15, 2014	Feb DISC
>A>		@	3MG	A091153	003	Apr 15, 2014	Feb DISC

ETHACRYNIC ACID

TABLET;ORAL  
 ETHACRYNIC ACID  
 UPSHER SMITH LABS 25MG

>A>	AB			A212417	001	Feb 19, 2020	Feb NEWA
-----	----	--	--	---------	-----	--------------	----------

ETHINYL ESTRADIOL; LEVONORGESTREL

>A> SYSTEM;TRANSDERMAL  
 >A> TWIRLA  
 >A> +! AGILE 0.03MG/24HR;0.12MG/24HR

N204017	001	Feb 14, 2020	Feb NEWA
---------	-----	--------------	----------

TABLET;ORAL  
 BALCOLTRA  
 +! AVION PHARMS 0.02MG;0.1MG  
 + 0.02MG;0.1MG

N208612	001	Jan 09, 2018	Jan CHRS
N208612	001	Jan 09, 2018	Jan CRLD

TABLET;ORAL-28  
 LEVONORGESTREL AND ETHINYL ESTRADIOL

>D>	AB	MYLAN LABS LTD	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A202970	001	Mar 23, 2018	Feb DISC
>A>		@	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A202970	001	Mar 23, 2018	Feb DISC

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET;ORAL  
 LOESTRIN 24 FE  
 AB + TEVA BRANDED PHARM 0.02MG;1MG

N021871	001	Feb 17, 2006	Jan CAHN
---------	-----	--------------	----------

TABLET;ORAL-21  
 LOESTRIN 21 1.5/30  
 AB + TEVA BRANDED PHARM 0.03MG;1.5MG  
 LOESTRIN 21 1/20

N017875	001		Jan CAHN
---------	-----	--	----------

AB + TEVA BRANDED PHARM 0.02MG;1MG  
 LOESTRIN FE 1.5/30  
 AB +! TEVA BRANDED PHARM 0.03MG;1.5MG

N017876	001		Jan CAHN
N017355	001		Jan CAHN

TABLET;ORAL-28  
 LOESTRIN FE 1/20  
 AB + TEVA BRANDED PHARM 0.02MG;1MG  
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

>D>	AB	MYLAN LABS LTD	0.02MG,0.03MG,0.035MG;1MG,1MG,1MG	A205069	001	Jun 22, 2018	Feb DISC
>A>		@	0.02MG,0.03MG,0.035MG;1MG,1MG,1MG	A205069	001	Jun 22, 2018	Feb DISC

ETODOLAC

TABLET; ORAL

ETODOLAC

>D>	AB	TEVA	400MG	A075009	001	Nov 26, 1997	Feb DISC
>A>		@	400MG	A075009	001	Nov 26, 1997	Feb DISC
>D>	AB		500MG	A075009	002	Dec 28, 1999	Feb DISC
>A>		@	500MG	A075009	002	Dec 28, 1999	Feb DISC

ETOMIDATE

INJECTABLE; INJECTION

ETOMIDATE

>D>	AP	MYLAN LABS LTD	2MG/ML	A078289	001	Jan 02, 2009	Feb DISC
>A>		@	2MG/ML	A078289	001	Jan 02, 2009	Feb DISC

EXEMESTANE

TABLET; ORAL

EXEMESTANE

@ MAYNE PHARMA INC 25MG

A208764 001 Aug 08, 2019 Jan DISC

EZETIMIBE

TABLET; ORAL

EZETIMIBE

>D>	AB	MYLAN	10MG	A201790	001	Apr 26, 2019	Feb DISC
>A>		@	10MG	A201790	001	Apr 26, 2019	Feb DISC

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

AB	!	AUROBINDO PHARMA LTD	40MG	A206530	002	Dec 22, 2015	Jan CHRS
>D>	AB	MYLAN	20MG	A075704	001	Apr 16, 2001	Feb DISC
>A>		@	20MG	A075704	001	Apr 16, 2001	Feb DISC
		PEPCID					
	+	@ VALEANT PHARMS NORTH	20MG	N019462	001	Oct 15, 1986	Jan DISC
	+	@	40MG	N019462	002	Oct 15, 1986	Jan DISC

FENOFIBRATE

CAPSULE; ORAL

FENOFIBRATE (MICRONIZED)

AB		ANI PHARMS INC	67MG	A209504	001	Apr 30, 2018	Jan CAHN
AB			134MG	A209504	002	Apr 30, 2018	Jan CAHN
AB			200MG	A209504	003	Apr 30, 2018	Jan CAHN

TABLET; ORAL

FENOFIBRATE

AB		ALEMBIC PHARMS LTD	54MG	A213252	001	Jan 17, 2020	Jan NEWA
AB			160MG	A213252	002	Jan 17, 2020	Jan NEWA
AB		APPCO	54MG	A210670	001	Sep 06, 2019	Jan CAHN
AB			160MG	A210670	002	Sep 06, 2019	Jan CAHN
AB		GRAVITI PHARMS	54MG	A210606	001	Aug 17, 2018	Jan CMFD
AB			160MG	A210606	002	Aug 17, 2018	Jan CMFD

FENTANYL CITRATE

INJECTABLE; INJECTION

FENTANYL CITRATE PRESERVATIVE FREE

>A>	AP	! HIKMA	EQ 0.05MG BASE/ML	N019101	001	Jul 11, 1984	Feb CAHN
>D>	AP	! WEST-WARD PHARMS INT	EQ 0.05MG BASE/ML	N019101	001	Jul 11, 1984	Feb CAHN

FERRIC DERISOMALTOSE

SOLUTION; INTRAVENOUS

MONOFERRIC

+	!	PHARMACOSMOS AS	100MG/ML (100MG/ML)	N208171	001	Jan 16, 2020	Jan NEWA
+	!		500MG/5ML (100MG/ML)	N208171	002	Jan 16, 2020	Jan NEWA
+	!		1GM/10ML (100MG/ML)	N208171	003	Jan 16, 2020	Jan NEWA

FESOTERODINE FUMARATE

TABLET, EXTENDED RELEASE; ORAL

FESOTERODINE FUMARATE

>D>	AB	DR REDDYS LABS LTD	4MG	A204975	001	Aug 13, 2019	Feb DISC
>A>		@	4MG	A204975	001	Aug 13, 2019	Feb DISC
>D>	AB		8MG	A204975	002	Aug 13, 2019	Feb DISC
>A>		@	8MG	A204975	002	Aug 13, 2019	Feb DISC

FIDAXOMICIN

FOR SUSPENSION;ORAL  
DIFICID

+! CUBIST PHARMS LLC 40MG/ML N213138 001 Jan 24, 2020 Jan NEWA

FLUCONAZOLE

FOR SUSPENSION;ORAL  
FLUCONAZOLE

>D> AB HIKMA 50MG/5ML A076246 001 Jul 29, 2004 Feb DISC  
>A> @ 50MG/5ML A076246 001 Jul 29, 2004 Feb DISC  
>D> AB 200MG/5ML A076246 002 Jul 29, 2004 Feb DISC  
>A> @ 200MG/5ML A076246 002 Jul 29, 2004 Feb DISC

TABLET;ORAL  
FLUCONAZOLE

@ MYLAN 50MG A076351 001 Jul 29, 2004 Jan DISC  
@ 100MG A076351 002 Jul 29, 2004 Jan DISC  
@ 150MG A076351 003 Jul 29, 2004 Jan DISC  
@ 200MG A076351 004 Jul 29, 2004 Jan DISC

FLUDEOXYGLUCOSE F-18

INJECTABLE;INTRAVENOUS  
FLUDEOXYGLUCOSE F18

AP UNIV TX SW MEDCTR 20-200mCi/ML A210265 001 Feb 06, 2020 Jan NEWA

FLUOCINONIDE

CREAM;TOPICAL  
FLUOCINONIDE

>A> AB CADILA 0.1% A208989 001 Feb 10, 2020 Feb CAHN  
>D> AB ZYDUS PHARMS 0.1% A208989 001 Feb 10, 2020 Feb CAHN  
AB 0.1% A208989 001 Feb 10, 2020 Jan NEWA

FOLIC ACID

INJECTABLE;INJECTION  
FOLIC ACID

>D> AP X-GEN PHARMS INC 5MG/ML A202522 001 Nov 06, 2019 Feb CAHN  
AP 5MG/ML A202522 001 Nov 06, 2019 Jan CAHN  
>A> AP XGEN PHARMS 5MG/ML A202522 001 Nov 06, 2019 Feb CAHN

FULVESTRANT

INJECTABLE;INTRAMUSCULAR  
FULVESTRANT

AO CHIA TAI TIANQING 50MG/ML A211422 001 Feb 07, 2020 Jan NEWA

FUROSEMIDE

TABLET;ORAL  
FUROSEMIDE

@ AVET 20MG N018413 001 Nov 30, 1983 Jan DISC  
@ 40MG N018413 002 Nov 30, 1983 Jan DISC

GABAPENTIN

CAPSULE;ORAL  
GABAPENTIN

AB CSPC OUYI 100MG A075477 001 Mar 23, 2005 Jan CMFD  
AB 300MG A075477 002 Mar 23, 2005 Jan CMFD  
AB 400MG A075477 003 Mar 23, 2005 Jan CMFD

TABLET;ORAL  
GRALISE

BX +! ALMATICA 300MG N022544 001 Jan 28, 2011 Jan CAHN  
BX +! 600MG N022544 002 Jan 28, 2011 Jan CAHN

GALANTAMINE HYDROBROMIDE

TABLET;ORAL  
GALANTAMINE HYDROBROMIDE

>D> AB HIKMA EQ 4MG BASE A077608 001 Feb 11, 2009 Feb DISC  
>A> @ EQ 4MG BASE A077608 001 Feb 11, 2009 Feb DISC  
>D> AB EQ 8MG BASE A077608 002 Feb 11, 2009 Feb DISC  
>A> @ EQ 8MG BASE A077608 002 Feb 11, 2009 Feb DISC  
>D> AB EQ 12MG BASE A077608 003 Feb 11, 2009 Feb DISC  
>A> @ EQ 12MG BASE A077608 003 Feb 11, 2009 Feb DISC

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION  
GEMCITABINE HYDROCHLORIDE

>D>	AP	APOTEX	200MG/5.26ML (38MG/ML)	A206776	001	May 23, 2017	Feb	DISC
>A>		@	200MG/5.26ML (38MG/ML)	A206776	001	May 23, 2017	Feb	DISC
>D>	AP		1GM/26.3ML (38MG/ML)	A206776	002	May 23, 2017	Feb	DISC
>A>		@	1GM/26.3ML (38MG/ML)	A206776	002	May 23, 2017	Feb	DISC
>D>	AP		2GM/52.6ML (38MG/ML)	A206776	003	May 23, 2017	Feb	DISC
>A>		@	2GM/52.6ML (38MG/ML)	A206776	003	May 23, 2017	Feb	DISC
	AP	SHILPA MEDICARE LTD	EQ 200MG BASE/VIAL	A207575	001	Feb 22, 2019	Jan	CMFD
	AP		EQ 1GM BASE/VIAL	A207575	002	Feb 22, 2019	Jan	CMFD

GEMFIBROZIL

TABLET; ORAL  
GEMFIBROZIL

>A>	AB	CADILA	600MG	A204189	001	Aug 28, 2018	Feb	CAHN
>D>	AB	TEVA	600MG	A074256	001	Oct 31, 1993	Feb	DISC
>A>		@	600MG	A074256	001	Oct 31, 1993	Feb	DISC
>D>	AB	ZYDUS PHARMS	600MG	A204189	001	Aug 28, 2018	Feb	CAHN

GLYBURIDE

TABLET; ORAL  
GLYBURIDE (MICRONIZED)

		@ MYLAN	1.5MG	A074792	001	Jun 26, 1998	Jan	DISC
		@	3MG	A074792	002	Jun 26, 1998	Jan	DISC
		@	6MG	A074792	003	Aug 17, 1999	Jan	DISC

GLYCOPYRROLATE

INJECTABLE; INJECTION  
GLYCOPYRROLATE

>A>	AP	RICONPHARMA LLC	0.2MG/ML	A210083	001	Feb 21, 2020	Feb	NEWA
-----	----	-----------------	----------	---------	-----	--------------	-----	------

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION  
GRANISETRON HYDROCHLORIDE

>A>	AP	YUNG SHIN PHARM	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)	A202647	001	Mar 06, 2020	Feb	NEWA
-----	----	-----------------	-------------------------------------	---------	-----	--------------	-----	------

TABLET; ORAL

		@ MYLAN	EQ 1MG BASE	A078725	001	Jan 30, 2008	Jan	DISC
--	--	---------	-------------	---------	-----	--------------	-----	------

HEPARIN SODIUM

INJECTABLE; INJECTION  
HEPARIN SODIUM

>A>	AP	NANJING KING-FRIEND	20,000 UNITS/ML	A211004	001	Feb 24, 2020	Feb	NEWA
-----	----	---------------------	-----------------	---------	-----	--------------	-----	------

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

TABLET; ORAL  
HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE

AA	!	NOVEL LABS INC	1.5MG;5MG	A091528	001	Apr 20, 2011	Jan	CHRS
		TUSSIGON						
		@ KING PHARMS	1.5MG;5MG	A088508	001	Jul 30, 1985	Jan	DISC

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION  
HYDRALAZINE HYDROCHLORIDE

>D>	AP	X-GEN PHARMS INC	20MG/ML	A203110	001	Jun 29, 2015	Feb	CAHN
>A>	AP	XGEN PHARMS	20MG/ML	A203110	001	Jun 29, 2015	Feb	CAHN

TABLET; ORAL

		@ STRIDES PHARMA	10MG	A200770	004	Jun 25, 2019	Jan	DISC
--	--	------------------	------	---------	-----	--------------	-----	------

HYDROCHLOROTHIAZIDE

TABLET; ORAL  
HYDROCHLOROTHIAZIDE

		@ MYLAN PHARMS INC	25MG	A040735	002	Jan 23, 2007	Jan	DISC
		@	50MG	A040735	003	Jan 23, 2007	Jan	DISC

HYDROCHLOROTHIAZIDE; LISINAPRIL

## TABLET; ORAL

## LISINAPRIL AND HYDROCHLOROTHIAZIDE

>D>	@	APOTEX	12.5MG;10MG	A076674	001	Oct 05, 2004	Feb CAHN
>D>	@		12.5MG;20MG	A076674	002	Oct 05, 2004	Feb CAHN
>D>	@		25MG;20MG	A076674	003	Oct 05, 2004	Feb CAHN
>A>	@	COREPHARMA	12.5MG;10MG	A076674	001	Oct 05, 2004	Feb CAHN
>A>	@		12.5MG;20MG	A076674	002	Oct 05, 2004	Feb CAHN
>A>	@		25MG;20MG	A076674	003	Oct 05, 2004	Feb CAHN

HYDROCHLOROTHIAZIDE; TRIAMTERENE

## CAPSULE; ORAL

## TRIAMTERENE AND HYDROCHLOROTHIAZIDE

>A>	AB	CADILA	25MG;37.5MG	A208358	001	Feb 11, 2019	Feb CAHN
>D>	AB	ZYDUS PHARMS	25MG;37.5MG	A208358	001	Feb 11, 2019	Feb CAHN

HYDROCODONE BITARTRATE

## CAPSULE, EXTENDED RELEASE; ORAL

## HYDROCODONE BITARTRATE

AB		ALVOGEN	10MG	A206986	001	Jan 21, 2020	Jan NFTG
AB			15MG	A206986	002	Jan 21, 2020	Jan NFTG
AB			20MG	A206986	003	Jan 21, 2020	Jan NFTG
AB			30MG	A206986	004	Jan 21, 2020	Jan NFTG
AB			40MG	A206986	005	Jan 21, 2020	Jan NFTG
AB			50MG	A206986	006	Jan 21, 2020	Jan NFTG
		ZOHYDRO ER					
AB	+	PERSION	10MG	N202880	001	Oct 25, 2013	Jan CFTG
AB	+		15MG	N202880	002	Oct 25, 2013	Jan CFTG
AB	+		20MG	N202880	003	Oct 25, 2013	Jan CFTG
AB	+		30MG	N202880	004	Oct 25, 2013	Jan CFTG
AB	+		40MG	N202880	005	Oct 25, 2013	Jan CFTG
AB	+		50MG	N202880	006	Oct 25, 2013	Jan CFTG

HYDROCORTISONE

## TABLET; ORAL

## HYDROCORTISONE

>D>	AB	PII	5MG	A207029	001	Apr 27, 2017	Feb CAHN
>D>	AB		10MG	A207029	002	Apr 27, 2017	Feb CAHN
>D>	AB		20MG	A207029	003	Apr 27, 2017	Feb CAHN
>A>	AB	STRIDES PHARMA	5MG	A207029	001	Apr 27, 2017	Feb CAHN
>A>	AB		10MG	A207029	002	Apr 27, 2017	Feb CAHN
>A>	AB		20MG	A207029	003	Apr 27, 2017	Feb CAHN

HYDROCORTISONE VALERATE

## OINTMENT; TOPICAL

## HYDROCORTISONE VALERATE

>A>	AB	COSETTE	0.2%	A211764	001	Mar 04, 2020	Feb NEWA
-----	----	---------	------	---------	-----	--------------	----------

HYDROMORPHONE HYDROCHLORIDE

## INJECTABLE; INJECTION

## DILAUDID

+	!	FRESENIUS KABI USA	0.2MG/ML	N019034	006	Jan 16, 2020	Jan NEWA
---	---	--------------------	----------	---------	-----	--------------	----------

## TABLET; ORAL

## HYDROMORPHONE HYDROCHLORIDE

>D>	AB	ELITE LABS	8MG	A076723	001	Oct 18, 2005	Feb CAHN
>A>	AB	NOSTRUM LABS INC	8MG	A076723	001	Oct 18, 2005	Feb CAHN

HYDROXYPROGESTERONE CAPROATE

## SOLUTION; INTRAMUSCULAR

## HYDROXYPROGESTERONE CAPROATE

>D>	AP1	AM REGENT	1250MG/5ML (250MG/ML)	A210724	001	Aug 09, 2019	Feb DISC
>A>		@	1250MG/5ML (250MG/ML)	A210724	001	Aug 09, 2019	Feb DISC
	AP1		1250MG/5ML (250MG/ML)	A210724	001	Aug 09, 2019	Jan CMS1

IBUPROFEN LYSINE

## INJECTABLE; INTRAVENOUS

## IBUPROFEN LYSINE

>D>	AP	X-GEN PHARMS INC	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	A202402	001	Mar 30, 2016	Feb CAHN
>A>	AP	XGEN PHARMS	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	A202402	001	Mar 30, 2016	Feb CAHN

ICATIBANT ACETATE

INJECTABLE;SUBCUTANEOUS  
ICATIBANT ACETATE

>A> AP JIANGSU HANSOH PHARM EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A211021 001 Mar 09, 2020 Feb NEWA

INDAPAMIDE

TABLET;ORAL  
INDAPAMIDE

AB ANI PHARMS INC 1.25MG A075201 001 Dec 04, 1998 Jan CAHN  
AB 2.5MG A075201 002 Dec 04, 1998 Jan CAHN

INDOMETHACIN

CAPSULE;ORAL  
INDOMETHACIN

AB HERITAGE 25MG N018851 001 May 18, 1984 Jan CAHN  
AB 50MG N018851 002 May 18, 1984 Jan CAHN

INSULIN LISPRO RECOMBINANT

INJECTABLE;INJECTION  
HUMALOG KWIKPEN

+! LILLY 200UNITS/ML N020563 004 Jan 06, 2017 Jan CMS1

IPRATROPIUM BROMIDE

SPRAY, METERED;NASAL  
IPRATROPIUM BROMIDE

AB APOTEX INC 0.021MG/SPRAY A076156 001 Apr 18, 2003 Jan CMFD

IRBESARTAN

TABLET;ORAL  
IRBESARTAN

>D> AB HIKMA 75MG A090201 001 Oct 15, 2012 Feb DISC  
>A> @ 75MG A090201 001 Oct 15, 2012 Feb DISC  
>D> AB 150MG A090201 002 Oct 15, 2012 Feb DISC  
>A> @ 150MG A090201 002 Oct 15, 2012 Feb DISC  
>D> AB 300MG A090201 003 Oct 15, 2012 Feb DISC  
>A> @ 300MG A090201 003 Oct 15, 2012 Feb DISC

ISOSORBIDE MONONITRATE

TABLET, EXTENDED RELEASE;ORAL  
ISOSORBIDE MONONITRATE

>D> AB HIKMA INTL PHARMS 30MG A076813 002 Mar 30, 2006 Feb DISC  
>A> @ 30MG A076813 002 Mar 30, 2006 Feb DISC  
>D> AB 60MG A076813 001 Jan 07, 2005 Feb DISC  
>A> @ 60MG A076813 001 Jan 07, 2005 Feb DISC

ITRACONAZOLE

SOLUTION;ORAL  
ITRACONAZOLE

>D> AA APOTEX 10MG/ML A208481 001 Aug 02, 2019 Feb DISC  
>A> @ 10MG/ML A208481 001 Aug 02, 2019 Feb DISC

KETOROLAC TROMETHAMINE

TABLET;ORAL  
KETOROLAC TROMETHAMINE

>D> AB ! MYLAN 10MG A074761 001 May 16, 1997 Feb CHRS  
>A> AB 10MG A074761 001 May 16, 1997 Feb CHRS  
>D> AB TEVA 10MG A074754 001 May 16, 1997 Feb CHRS  
>A> AB ! 10MG A074754 001 May 16, 1997 Feb CHRS

LABETALOL HYDROCHLORIDE

INJECTABLE;INJECTION  
NORMODYNE

+ @ SCHERING 5MG/ML N018686 001 Aug 01, 1984 Jan CRLD

TABLET;ORAL  
LABETALOL HYDROCHLORIDE

>D> @ HERITAGE PHARMA 100MG A074787 001 Aug 03, 1998 Feb CMFD  
>A> AB 100MG A074787 001 Aug 03, 1998 Feb CMFD  
>D> @ 200MG A074787 002 Aug 03, 1998 Feb CMFD  
>A> AB 200MG A074787 002 Aug 03, 1998 Feb CMFD

>A>	<u>LACTITOL</u>						
>A>	FOR SOLUTION;ORAL						
>A>	PIZENSY						
>A>	+!	BRAINTREE LABS	10GM	N211281	001	Feb 12, 2020	Feb NEWA
>A>	+		10GM/PACKET	N211281	001	Feb 12, 2020	Feb NEWA
	<u>LASMIDITAN SUCCINATE</u>						
	TABLET;ORAL						
	REYVOW						
	+	ELI LILLY AND CO	50MG	N211280	001	Jan 31, 2020	Jan NEWA
	+!		100MG	N211280	002	Jan 31, 2020	Jan NEWA
	<u>LEFLUNOMIDE</u>						
	TABLET;ORAL						
	LEFLUNOMIDE						
AB		LUPIN LTD	10MG	A211863	001	Feb 04, 2020	Jan NEWA
AB			20MG	A211863	002	Feb 04, 2020	Jan NEWA
	<u>LEVETIRACETAM</u>						
	INJECTABLE;INTRAVENOUS						
	LEVETIRACETAM						
>D>	AP	X GEN PHARMS	500MG/5ML (100MG/ML)	A091485	001	Aug 05, 2011	Feb CAHN
>A>	AP	XGEN PHARMS	500MG/5ML (100MG/ML)	A091485	001	Aug 05, 2011	Feb CAHN
	<u>LEVOCETIRIZINE DIHYDROCHLORIDE</u>						
	SOLUTION;ORAL						
	LEVOCETIRIZINE DIHYDROCHLORIDE						
>D>	AA	APOTEX	2.5MG/5ML	A202915	001	Aug 21, 2014	Feb DISC
>A>		@	2.5MG/5ML	A202915	001	Aug 21, 2014	Feb DISC
	TABLET;ORAL						
	LEVOCETIRIZINE DIHYDROCHLORIDE						
>D>	AB	APOTEX	5MG	A203027	001	Feb 13, 2015	Feb DISC
>A>		@	5MG	A203027	001	Feb 13, 2015	Feb DISC
	<u>LEVOTHYROXINE SODIUM **</u>						
	**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium						
	TABLET;ORAL						
	THYRO-TABS						
AB2,	+	ALVOGEN	0.3MG	N021116	009	Oct 24, 2002	Jan CHRS
AB4							
	<u>LINCOMYCIN HYDROCHLORIDE</u>						
	INJECTABLE;INJECTION						
	LINCOMYCIN						
>D>	AP	X-GEN PHARMS INC	EQ 300MG BASE/ML	A201746	001	Jun 04, 2015	Feb CAHN
>A>	AP	XGEN PHARMS	EQ 300MG BASE/ML	A201746	001	Jun 04, 2015	Feb CAHN
	<u>LIOTHYRONINE SODIUM</u>						
	INJECTABLE;INJECTION						
	LIOTHYRONINE SODIUM						
>D>	AP	X GEN PHARMS	EQ 0.01MG BASE/ML	A076923	001	Aug 17, 2005	Feb CAHN
>A>	AP	XGEN PHARMS	EQ 0.01MG BASE/ML	A076923	001	Aug 17, 2005	Feb CAHN
	TABLET;ORAL						
	LIOTHYRONINE SODIUM						
AB		SUN PHARM	EQ 0.005MG BASE	A091382	001	Apr 20, 2016	Jan CMFD
AB			EQ 0.025MG BASE	A091382	002	Apr 20, 2016	Jan CMFD
AB			EQ 0.05MG BASE	A091382	003	Apr 20, 2016	Jan CMFD
	<u>LISINAPRIL</u>						
	TABLET;ORAL						
	LISINAPRIL						
>D>	AB	APOTEX INC	2.5MG	A076102	001	Sep 30, 2002	Feb CAHN
>D>	AB		5MG	A076102	002	Sep 30, 2002	Feb CAHN
>D>	AB		10MG	A076102	003	Sep 30, 2002	Feb CAHN
>D>	AB		20MG	A076102	004	Sep 30, 2002	Feb CAHN
>D>	AB		30MG	A076102	005	Sep 30, 2002	Feb CAHN
>D>	AB		40MG	A076102	006	Sep 30, 2002	Feb CAHN
>A>	AB	COREPHARMA	2.5MG	A076102	001	Sep 30, 2002	Feb CAHN
>A>	AB		5MG	A076102	002	Sep 30, 2002	Feb CAHN
>A>	AB		10MG	A076102	003	Sep 30, 2002	Feb CAHN
>A>	AB		20MG	A076102	004	Sep 30, 2002	Feb CAHN

TABLET;ORAL							
LISINOPRIL							
>A>	AB	30MG	A076102	005	Sep 30, 2002	Feb	CAHN
>A>	AB	40MG	A076102	006	Sep 30, 2002	Feb	CAHN
<u>LITHIUM CARBONATE</u>							
CAPSULE;ORAL							
LITHIUM CARBONATE							
AB	+ HIKMA	150MG	N017812	002	Jan 28, 1987	Jan	CAHN
AB	+	300MG	N017812	001		Jan	CAHN
AB	+!	600MG	N017812	003	Jan 28, 1987	Jan	CAHN
	@ MYLAN	150MG	A076243	002	Feb 24, 2003	Jan	DISC
	@	300MG	A076243	001	Jun 27, 2002	Jan	DISC
<u>LITHIUM CITRATE</u>							
SYRUP;ORAL							
LITHIUM CITRATE							
AA	+! HIKMA	EQ 300MG CARBONATE/5ML	N018421	001		Jan	CAHN
<u>LORCASERIN HYDROCHLORIDE</u>							
TABLET;ORAL							
BELVIQ							
>D>	+! EISAI INC	10MG	N022529	001	Jun 27, 2012	Feb	DISC
>A>	+ @	10MG	N022529	001	Jun 27, 2012	Feb	DISC
TABLET, EXTENDED RELEASE;ORAL							
BELVIQ XR							
>D>	+! EISAI INC	20MG	N208524	001	Jul 15, 2016	Feb	DISC
>A>	+ @	20MG	N208524	001	Jul 15, 2016	Feb	DISC
<u>LOSARTAN POTASSIUM</u>							
TABLET;ORAL							
LOSARTAN POTASSIUM							
>D>	AB HIKMA	25MG	A077459	001	Oct 06, 2010	Feb	DISC
>A>	@	25MG	A077459	001	Oct 06, 2010	Feb	DISC
>D>	AB	50MG	A077459	002	Oct 06, 2010	Feb	DISC
>A>	@	50MG	A077459	002	Oct 06, 2010	Feb	DISC
>D>	AB	100MG	A077459	003	Oct 06, 2010	Feb	DISC
>A>	@	100MG	A077459	003	Oct 06, 2010	Feb	DISC
<u>LOVASTATIN</u>							
TABLET;ORAL							
LOVASTATIN							
>D>	AB APOTEX INC	10MG	A077748	001	Feb 28, 2007	Feb	CAHN
>D>	AB	20MG	A077748	002	Feb 28, 2007	Feb	CAHN
>D>	AB	40MG	A077748	003	Feb 28, 2007	Feb	CAHN
>A>	AB COREPHARMA	10MG	A077748	001	Feb 28, 2007	Feb	CAHN
>A>	AB	20MG	A077748	002	Feb 28, 2007	Feb	CAHN
>A>	AB	40MG	A077748	003	Feb 28, 2007	Feb	CAHN
<u>MAZINDOL</u>							
TABLET;ORAL							
SANOREX							
	+ @ NOVARTIS	1MG	N017247	001		Jan	CAHN
	+ @	2MG	N017247	002		Jan	CAHN
<u>MECLIZINE HYDROCHLORIDE</u>							
TABLET;ORAL							
MECLIZINE HYDROCHLORIDE							
	@ MYLAN PHARMS INC	12.5MG	A202640	001	Sep 17, 2012	Jan	DISC
	@	25MG	A202640	002	Sep 17, 2012	Jan	DISC
<u>MEFLOQUINE HYDROCHLORIDE</u>							
TABLET;ORAL							
MEFLOQUINE HYDROCHLORIDE							
>D>	AB ! BARR	250MG	A076392	001	Dec 29, 2003	Feb	CTEC
>A>	!	250MG	A076392	001	Dec 29, 2003	Feb	CTEC
>D>	AB HIKMA	250MG	A076523	001	Oct 01, 2004	Feb	DISC
>A>	@	250MG	A076523	001	Oct 01, 2004	Feb	DISC



MEGESTROL ACETATE

SUSPENSION;ORAL  
MEGESTROL ACETATE

>D>	AB	!	HIKMA	40MG/ML	A 075997	001	Feb 15, 2002	Feb	DISC
>A>		@		40MG/ML	A 075997	001	Feb 15, 2002	Feb	DISC
>D>	AB		PAR PHARM	40MG/ML	A 075671	001	Jul 25, 2001	Feb	CHRS
>A>	AB	!		40MG/ML	A 075671	001	Jul 25, 2001	Feb	CHRS

MELOXICAM

SOLUTION;INTRAVENOUS  
ANJESO

>A>		+!	BAUDAX	30MG/ML (30MG/ML)	N210583	001	Feb 20, 2020	Feb	NEWA
-----	--	----	--------	-------------------	---------	-----	--------------	-----	------

MELPHALAN HYDROCHLORIDE

INJECTABLE;INJECTION  
MELPHALAN HYDROCHLORIDE

>A>	AP		INGENUS PHARMS LLC	EQ 50MG BASE/VIAL	A 210947	001	Feb 18, 2020	Feb	NEWA
>D>	AP		PAR STERILE PRODUCTS	EQ 50MG BASE/VIAL	A 204773	001	Aug 22, 2016	Feb	DISC
>A>		@		EQ 50MG BASE/VIAL	A 204773	001	Aug 22, 2016	Feb	DISC
>A>		@	TWI PHARMS	EQ 50MG BASE/VIAL	A 211463	001	Sep 13, 2019	Jan	DISC

POWDER;INTRAVENOUS  
EVOMELA

>D>		+!	ACROTECH	EQ 50MG BASE/VIAL	N 207155	001	Mar 10, 2016	Feb	CFTG
>A>	AP	+!		EQ 50MG BASE/VIAL	N 207155	001	Mar 10, 2016	Feb	CFTG
>A>			MELPHALAN HYDROCHLORIDE						
>A>	AP		ACTAVIS LLC	EQ 50MG BASE/VIAL	A 209323	001	Mar 06, 2020	Feb	NFTG

MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL  
MEMANTINE HYDROCHLORIDE

>A>	AB		ANI PHARMS INC	7MG	A 205365	001	Feb 28, 2020	Feb	NEWA
>A>	AB			14MG	A 205365	002	Feb 28, 2020	Feb	NEWA
>A>	AB			21MG	A 205365	003	Feb 28, 2020	Feb	NEWA
>A>	AB			28MG	A 205365	004	Feb 28, 2020	Feb	NEWA

TABLET;ORAL

MEMANTINE HYDROCHLORIDE

>A>	AB		HIKMA PHARMS	5MG	A 208173	001	Feb 28, 2020	Feb	NEWA
>A>	AB			10MG	A 208173	002	Feb 28, 2020	Feb	NEWA

MEPERIDINE HYDROCHLORIDE

TABLET;ORAL

MEPERIDINE HYDROCHLORIDE

>D>	AA		EPIC PHARMA	50MG	A 040331	001	May 28, 1999	Feb	CHRS
>A>	AA	!		50MG	A 040331	001	May 28, 1999	Feb	CHRS
>D>	AA			100MG	A 040331	002	May 28, 1999	Feb	CHRS
>A>	AA	!		100MG	A 040331	002	May 28, 1999	Feb	CHRS
>D>	AA	!	HIKMA	50MG	A 040110	001	Mar 12, 1997	Feb	DISC
>A>		@		50MG	A 040110	001	Mar 12, 1997	Feb	DISC
>D>	AA	!		100MG	A 040110	002	Mar 12, 1997	Feb	DISC
>A>		@		100MG	A 040110	002	Mar 12, 1997	Feb	DISC

MESALAMINE

SUPPOSITORY;RECTAL  
MESALAMINE

AB			ZYDUS PHARMS	1GM	A 208953	001	Feb 12, 2020	Jan	NEWA
----	--	--	--------------	-----	----------	-----	--------------	-----	------

METFORMIN HYDROCHLORIDE

SOLUTION;ORAL

METFORMIN HYDROCHLORIDE

>A>	AB		SAPTALIS PHARMS	500MG/5ML	A 211309	001	Mar 03, 2020	Feb	NFTG
>D>		+!	RANBAXY	500MG/5ML	N 021591	001	Sep 11, 2003	Feb	CFTG
>A>	AB	+!		500MG/5ML	N 021591	001	Sep 11, 2003	Feb	CFTG
>A>		+!		500MG/5ML	N 021591	001	Sep 11, 2003	Jan	CAHN

TABLET;ORAL

METFORMIN HYDROCHLORIDE

		@	MYLAN	500MG	A 075976	001	Jan 24, 2002	Jan	DISC
		@		850MG	A 075976	002	Jan 24, 2002	Jan	DISC
		@		1GM	A 075976	003	Jan 24, 2002	Jan	DISC

METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL

>D>	AA	HIKMA INTL PHARMS	500MG	A085159	001		Feb	DISC
>A>		@	500MG	A085159	001		Feb	DISC
>D>	AA		750MG	A085123	001		Feb	DISC
>A>		@	750MG	A085123	001		Feb	DISC

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE SODIUM PRESERVATIVE FREE

		@ MYLAN LABS LTD	EQ 50MG BASE/2ML (EQ 25MG BASE/ML)	A201529	001	Mar 29, 2012	Jan	DISC
		@	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	A201529	002	Mar 29, 2012	Jan	DISC
		@	EQ 200MG BASE/8ML (EQ 25MG BASE/ML)	A201529	003	Mar 29, 2012	Jan	DISC
		PHARMACHEMIE BV	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	A040843	003	Feb 27, 2012	Jan	CTEC

TABLET; ORAL

METHOTREXATE SODIUM

AB		ACCORD HLTHCARE	EQ 2.5MG BASE	A213343	001	Jan 24, 2020	Jan	NEWA
AB		EUGIA PHARMA	EQ 2.5MG BASE	A210454	001	Jan 30, 2020	Jan	NEWA

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

METOPROLOL SUCCINATE

		@ ACTAVIS LABS FL INC	EQ 25MG TARTRATE	A076862	002	Aug 03, 2009	Jan	DISC
		@	EQ 100MG TARTRATE	A077298	001	Apr 15, 2010	Jan	DISC
		@	EQ 200MG TARTRATE	A077298	002	Apr 15, 2010	Jan	DISC

METOPROLOL TARTRATE

TABLET; ORAL

METOPROLOL TARTRATE

>D>	AB	HERITAGE PHARMA	50MG	A074141	001	Jan 31, 1995	Feb	DISC
>A>		@	50MG	A074141	001	Jan 31, 1995	Feb	DISC
>D>	AB		100MG	A074141	002	Jan 31, 1995	Feb	DISC
>A>		@	100MG	A074141	002	Jan 31, 1995	Feb	DISC
AB		YOUNGTECH PHARMS INC	25MG	A208955	001	Feb 05, 2020	Jan	NEWA
AB			50MG	A208955	002	Feb 05, 2020	Jan	NEWA
AB			100MG	A208955	003	Feb 05, 2020	Jan	NEWA

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE

		@ MYLAN ASI	EQ 1MG BASE/ML	A090315	001	Nov 29, 2010	Jan	DISC
		@	EQ 5MG BASE/ML	A090315	002	Nov 29, 2010	Jan	DISC

MINOCYCLINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

XIMINO

>A>		JOURNEY	EQ 45MG BASE	N201922	001	Jul 11, 2012	Feb	CAHN
>A>		@	EQ 67.5MG BASE	N201922	002	Jul 11, 2012	Feb	CAHN
>A>			EQ 90MG BASE	N201922	003	Jul 11, 2012	Feb	CAHN
>A>		@	EQ 112.5MG BASE	N201922	004	Jul 11, 2012	Feb	CAHN
>A>			EQ 135MG BASE	N201922	005	Jul 11, 2012	Feb	CAHN
>D>		SUN PHARM INDS INC	EQ 45MG BASE	N201922	001	Jul 11, 2012	Feb	CAHN
>D>		@	EQ 67.5MG BASE	N201922	002	Jul 11, 2012	Feb	CAHN
>D>			EQ 90MG BASE	N201922	003	Jul 11, 2012	Feb	CAHN
>D>		@	EQ 112.5MG BASE	N201922	004	Jul 11, 2012	Feb	CAHN
>D>			EQ 135MG BASE	N201922	005	Jul 11, 2012	Feb	CAHN

MIRABEGRON

TABLET, EXTENDED RELEASE; ORAL

MIRABEGRON

		@ SAWAI USA	25MG	A209446	001	Dec 27, 2019	Jan	DISC
		MYRBETRIQ						
		+! APGDI	25MG	N202611	001	Jun 28, 2012	Jan	CTEC

MONTELUKAST SODIUM

		GRANULE; ORAL							
		MONTELUKAST SODIUM							
>A>	AB	AUROBINDO PHARMA LTD	EQ 4MG BASE/PACKET	A213471	001	Feb 18, 2020	Feb	NEWA	
		TABLET; ORAL							
		MONTELUKAST SODIUM							
>D>	AB	HIKMA	EQ 10MG BASE	A090655	001	Aug 03, 2012	Feb	DISC	
>A>		@	EQ 10MG BASE	A090655	001	Aug 03, 2012	Feb	DISC	
		TABLET, CHEWABLE; ORAL							
		MONTELUKAST SODIUM							
		@ JUBILANT GENERICS	EQ 4MG BASE	A203795	001	Feb 27, 2015	Jan	DISC	
		@	EQ 5MG BASE	A203795	002	Feb 27, 2015	Jan	DISC	

MORPHINE SULFATE

		INJECTABLE; INJECTION							
		DURAMORPH PF							
>A>	AP	+!	HIKMA	0.5MG/ML	N018565	001	Sep 18, 1984	Feb	CAHN
>A>	AP	+!		1MG/ML	N018565	002	Sep 18, 1984	Feb	CAHN
>D>	AP	+!	WEST-WARD PHARMS INT	0.5MG/ML	N018565	001	Sep 18, 1984	Feb	CAHN
>D>	AP	+!		1MG/ML	N018565	002	Sep 18, 1984	Feb	CAHN
		INFUMORPH							
>A>	AP	+!	HIKMA	10MG/ML	N018565	003	Jul 19, 1991	Feb	CAHN
>A>	AP	+!		25MG/ML	N018565	004	Jul 19, 1991	Feb	CAHN
>D>	AP	+!	WEST-WARD PHARMS INT	10MG/ML	N018565	003	Jul 19, 1991	Feb	CAHN
>D>	AP	+!		25MG/ML	N018565	004	Jul 19, 1991	Feb	CAHN
		MORPHINE SULFATE							
>D>	AP		EUROHLTH INTL SARL	4MG/ML	A205758	001	May 21, 2015	Feb	CAHN
>D>	AP			8MG/ML	A205758	002	May 21, 2015	Feb	CAHN
>D>	AP			10MG/ML	A205758	003	May 21, 2015	Feb	CAHN
>A>	AP		HIKMA	4MG/ML	A205758	001	May 21, 2015	Feb	CAHN
>A>	AP			8MG/ML	A205758	002	May 21, 2015	Feb	CAHN
>A>	AP			10MG/ML	A205758	003	May 21, 2015	Feb	CAHN

NALBUPHINE HYDROCHLORIDE

		INJECTABLE; INJECTION							
		NALBUPHINE HYDROCHLORIDE							
>D>	AP		MYLAN LABS LTD	10MG/ML	A206506	001	Feb 06, 2019	Feb	DISC
>A>		@		10MG/ML	A206506	001	Feb 06, 2019	Feb	DISC
>D>	AP			10MG/ML	A207595	001	Jan 11, 2019	Feb	DISC
>A>		@		10MG/ML	A207595	001	Jan 11, 2019	Feb	DISC
>D>	AP			20MG/ML	A206506	002	Feb 06, 2019	Feb	DISC
>A>		@		20MG/ML	A206506	002	Feb 06, 2019	Feb	DISC
>D>	AP			20MG/ML	A207595	002	Jan 11, 2019	Feb	DISC
>A>		@		20MG/ML	A207595	002	Jan 11, 2019	Feb	DISC

NALOXONE HYDROCHLORIDE

		INJECTABLE; INJECTION							
		NALOXONE HYDROCHLORIDE							
AP		PAR STERILE PRODUCTS	0.4MG/ML	A211286	001	Jan 17, 2020	Jan	NEWA	

NAPROXEN

		TABLET; ORAL							
		NAPROXEN							
		@ MYLAN	250MG	A074121	001	Dec 21, 1993	Jan	DISC	
		@	375MG	A074121	002	Dec 21, 1993	Jan	DISC	
		@	500MG	A074121	003	Dec 21, 1993	Jan	DISC	
>A>	AB	SCIEGEN PHARMS INC	250MG	A212517	001	Feb 21, 2020	Feb	NEWA	
>A>	AB		375MG	A212517	002	Feb 21, 2020	Feb	NEWA	
>A>	AB		500MG	A212517	003	Feb 21, 2020	Feb	NEWA	

NEOMYCIN SULFATE

		TABLET; ORAL							
		NEOMYCIN SULFATE							
>D>	AA	X GEN PHARMS	500MG	A065220	001	Jul 28, 2006	Feb	CAHN	
>A>	AA	XGEN PHARMS	500MG	A065220	001	Jul 28, 2006	Feb	CAHN	

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION;IRRIGATION

NEOMYCIN AND POLYMYXIN B SULFATE

>D>	AT	X GEN PHARMS	EQ 40MG BASE/ML;200,000 UNITS/ML	A 065106	001	Jan 31, 2006	Feb CAHN
>D>	AT		EQ 40MG BASE/ML;200,000 UNITS/ML	A 065108	001	Jan 31, 2006	Feb CAHN
>A>	AT	XGEN PHARMS	EQ 40MG BASE/ML;200,000 UNITS/ML	A 065106	001	Jan 31, 2006	Feb CAHN
>A>	AT		EQ 40MG BASE/ML;200,000 UNITS/ML	A 065108	001	Jan 31, 2006	Feb CAHN

>D> NETARSUDIL DIMESYLATE

>D> SOLUTION/DROPS;OPHTHALMIC  
>D> RHOPRESSA

>D>	+	AERIE PHARMS INC	EQ 0.02% BASE	N 208254	001	Dec 18, 2017	Feb CAIN
-----	---	------------------	---------------	----------	-----	--------------	----------

>A> NETARSUDIL MESYLATE

>A> SOLUTION/DROPS;OPHTHALMIC  
>A> RHOPRESSA

>A>	+	AERIE PHARMS INC	EQ 0.02% BASE	N 208254	001	Dec 18, 2017	Feb CAIN
-----	---	------------------	---------------	----------	-----	--------------	----------

NIACIN

TABLET, EXTENDED RELEASE;ORAL

NIACIN

@	MYLAN	500MG	A 203742	001	Feb 22, 2019	Jan DISC
@		750MG	A 203742	002	Feb 22, 2019	Jan DISC
@		1GM	A 203742	003	Feb 22, 2019	Jan DISC

NIRAPARIB TOSYLATE

CAPSULE;ORAL

ZEJULA

+	GLAXOSMITHKLINE	EQ 100MG BASE	N 208447	001	Mar 27, 2017	Jan CAHN
---	-----------------	---------------	----------	-----	--------------	----------

NORETHINDRONE

TABLET;ORAL-28

NOR-QD

AB1	+	TEVA BRANDED PHARM	0.35MG	N 017060	001		Jan CAHN
-----	---	--------------------	--------	----------	-----	--	----------

NYSTATIN

POWDER;TOPICAL

NYSTATIN

>D>	AT	X GEN PHARMS	100,000 UNITS/GM	A 065175	001	Dec 17, 2004	Feb CAHN
>A>	AT	XGEN PHARMS	100,000 UNITS/GM	A 065175	001	Dec 17, 2004	Feb CAHN

OCTREOTIDE ACETATE

INJECTABLE;INJECTION

OCTREOTIDE ACETATE

>D>	AP	HERITAGE PHARMS INC	EQ 0.05MG BASE/ML	A 204669	001	Dec 27, 2018	Feb DISC
>A>		@	EQ 0.05MG BASE/ML	A 204669	001	Dec 27, 2018	Feb DISC
>D>	AP		EQ 0.1MG BASE/ML	A 204669	002	Dec 27, 2018	Feb DISC
>A>		@	EQ 0.1MG BASE/ML	A 204669	002	Dec 27, 2018	Feb DISC
>D>	AP		EQ 0.5MG BASE/ML	A 204669	003	Dec 27, 2018	Feb DISC
>A>		@	EQ 0.5MG BASE/ML	A 204669	003	Dec 27, 2018	Feb DISC

SOLUTION;SUBCUTANEOUS

BYNFEZIA PEN

+	SUN PHARM	EQ 2.5MG BASE/ML (EQ 2.5MG BASE/ML)	N 213224	001	Jan 28, 2020	Jan NEWA
---	-----------	-------------------------------------	----------	-----	--------------	----------

OLANZAPINE

TABLET;ORAL

OLANZAPINE

@	TEVA PHARMS	2.5MG	A 076000	001	Oct 24, 2011	Jan DISC
@		5MG	A 076000	002	Oct 24, 2011	Jan DISC
@		7.5MG	A 076000	003	Oct 24, 2011	Jan DISC
@		10MG	A 076000	004	Oct 24, 2011	Jan DISC
@		15MG	A 076000	005	Oct 24, 2011	Jan DISC

OLMESARTAN MEDOXOMIL

TABLET;ORAL

OLMESARTAN MEDOXOMIL

>D>	AB	JUBILANT GENERICS	5MG	A 205482	001	Apr 24, 2017	Feb DISC
>A>		@	5MG	A 205482	001	Apr 24, 2017	Feb DISC
>D>	AB		20MG	A 205482	002	Apr 24, 2017	Feb DISC
>A>		@	20MG	A 205482	002	Apr 24, 2017	Feb DISC

## TABLET;ORAL

## OLMESARTAN MEDOXOMIL

>D>	AB		40MG	A205482	003	Apr 24, 2017	Feb DISC
>A>	@		40MG	A205482	003	Apr 24, 2017	Feb DISC

OLOPATADINE HYDROCHLORIDE

## SOLUTION/DROPS;OPHTHALMIC

## OLOPATADINE HYDROCHLORIDE

>A>	AT	WATSON LABS INC	EQ 0.7% BASE	A208637	001	Feb 19, 2020	Feb NFTG
		PAZEO					
>D>		+! NOVARTIS	EQ 0.7% BASE	N206276	001	Jan 30, 2015	Feb CFTG
>A>	AT	+!	EQ 0.7% BASE	N206276	001	Jan 30, 2015	Feb CFTG

## SPRAY, METERED;NASAL

## OLOPATADINE HYDROCHLORIDE

AB		AMNEAL PHARMS LLC	0.665MG/SPRAY	A210901	001	Jan 28, 2020	Jan NEWA
----	--	-------------------	---------------	---------	-----	--------------	----------

OMEGA-3-ACID ETHYL ESTERS

## CAPSULE;ORAL

## OMEGA-3-ACID ETHYL ESTERS

>D>		@ SOFGEN PHARMS	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A211355	001	Jul 10, 2019	Feb CMFD
>A>	AB		1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A211355	001	Jul 10, 2019	Feb CMFD

OMEPRAZOLE

## CAPSULE, DELAYED REL PELLETS;ORAL

## OMEPRAZOLE

>D>	AB	MYLAN	10MG	A205070	001	Jun 29, 2018	Feb DISC
>A>		@	10MG	A205070	001	Jun 29, 2018	Feb DISC
		@	20MG	A075876	002	May 29, 2003	Jan DISC
>D>	AB		20MG	A205070	002	Jun 29, 2018	Feb DISC
>A>		@	20MG	A205070	002	Jun 29, 2018	Feb DISC
>D>	AB		40MG	A205070	003	Jun 29, 2018	Feb DISC
>A>		@	40MG	A205070	003	Jun 29, 2018	Feb DISC

ONDANSETRON HYDROCHLORIDE

## TABLET;ORAL

## ONDANSETRON HYDROCHLORIDE

		@ MYLAN	EQ 4MG BASE	A076930	001	Jun 25, 2007	Jan DISC
		@	EQ 8MG BASE	A076930	002	Jun 25, 2007	Jan DISC

OSELTAMIVIR PHOSPHATE

## CAPSULE;ORAL

## OSELTAMIVIR PHOSPHATE

>A>	AB	SUNSHINE LAKE	EQ 30MG BASE	A212739	001	Mar 04, 2020	Feb NEWA
>A>	AB		EQ 45MG BASE	A212739	002	Mar 04, 2020	Feb NEWA
>A>	AB		EQ 75MG BASE	A212739	003	Mar 04, 2020	Feb NEWA

OXCARBAZEPINE

## TABLET;ORAL

## OXCARBAZEPINE

>D>	AB	HIKMA	150MG	A077795	001	Oct 09, 2007	Feb DISC
>A>		@	150MG	A077795	001	Oct 09, 2007	Feb DISC
>D>	AB		300MG	A077795	002	Oct 09, 2007	Feb DISC
>A>		@	300MG	A077795	002	Oct 09, 2007	Feb DISC
>D>	AB		600MG	A077795	003	Oct 09, 2007	Feb DISC
>A>		@	600MG	A077795	003	Oct 09, 2007	Feb DISC

OXYBUTYNIN CHLORIDE

## TABLET, EXTENDED RELEASE;ORAL

## OXYBUTYNIN CHLORIDE

		@ MYLAN PHARMS INC	10MG	A076644	001	Nov 09, 2006	Jan DISC
		@	15MG	A076644	002	May 10, 2007	Jan DISC

OXYCODONE HYDROCHLORIDE

## SOLUTION;ORAL

## OXYCODONE HYDROCHLORIDE

		@ ANI PHARMS INC	100MG/5ML	A203447	001	Aug 30, 2017	Jan DISC
AA		AUROLIFE PHARMA LLC	5MG/5ML	A212429	001	Jan 27, 2020	Jan NEWA
AA			100MG/5ML	A212429	002	Jan 27, 2020	Jan NEWA
>D>	AA	HIKMA	100MG/5ML	A203208	001	Jul 12, 2013	Feb DISC

SOLUTION;ORAL

OXYCODONE HYDROCHLORIDE

>A> @ 100MG/5ML A203208 001 Jul 12, 2013 Feb DISC

PAROXETINE HYDROCHLORIDE

TABLET;ORAL

PAROXETINE HYDROCHLORIDE

@ JUBILANT GENERICS

EQ 10MG BASE A205528 001 Nov 27, 2015 Jan DISC

@ EQ 20MG BASE A205528 002 Nov 27, 2015 Jan DISC

@ EQ 30MG BASE A205528 003 Nov 27, 2015 Jan DISC

@ EQ 40MG BASE A205528 004 Nov 27, 2015 Jan DISC

PASIREOTIDE DIASPARTATE

SOLUTION;SUBCUTANEOUS

SIGNIFOR

>D> + NOVARTIS EQ 0.3MG BASE/ML (EQ 0.3MG BASE/ML) N200677 001 Dec 14, 2012 Feb CAHN

>D> + EQ 0.6MG BASE/ML (EQ 0.6MG BASE/ML) N200677 002 Dec 14, 2012 Feb CAHN

>D> +! EQ 0.9MG BASE/ML (EQ 0.9MG BASE/ML) N200677 003 Dec 14, 2012 Feb CAHN

>A> + RECORDATI RARE EQ 0.3MG BASE/ML (EQ 0.3MG BASE/ML) N200677 001 Dec 14, 2012 Feb CAHN

>A> + EQ 0.6MG BASE/ML (EQ 0.6MG BASE/ML) N200677 002 Dec 14, 2012 Feb CAHN

>A> +! EQ 0.9MG BASE/ML (EQ 0.9MG BASE/ML) N200677 003 Dec 14, 2012 Feb CAHN

PASIREOTIDE PAMOATE

FOR SUSPENSION;INTRAMUSCULAR

SIGNIFOR LAR KIT

>D> + NOVARTIS EQ 10MG BASE/VIAL N203255 004 Jun 29, 2018 Feb CAHN

>D> + EQ 20MG BASE/VIAL N203255 001 Dec 15, 2014 Feb CAHN

>D> + EQ 30MG BASE/VIAL N203255 005 Jun 29, 2018 Feb CAHN

>D> + EQ 40MG BASE/VIAL N203255 002 Dec 15, 2014 Feb CAHN

>D> +! EQ 60MG BASE/VIAL N203255 003 Dec 15, 2014 Feb CAHN

>A> + RECORDATI RARE EQ 10MG BASE/VIAL N203255 004 Jun 29, 2018 Feb CAHN

>A> + EQ 20MG BASE/VIAL N203255 001 Dec 15, 2014 Feb CAHN

>A> + EQ 30MG BASE/VIAL N203255 005 Jun 29, 2018 Feb CAHN

>A> + EQ 40MG BASE/VIAL N203255 002 Dec 15, 2014 Feb CAHN

>A> +! EQ 60MG BASE/VIAL N203255 003 Dec 15, 2014 Feb CAHN

PAZOPANIB HYDROCHLORIDE

TABLET;ORAL

VOTRIENT

+ @ NOVARTIS EQ 400MG BASE N022465 002 Oct 19, 2009 Jan CRLD

PEGVISOMANT

INJECTABLE;SUBCUTANEOUS

SOMAVERT

+! PHARMACIA 10MG/VIAL N021106 001 Mar 25, 2003 Jan CAHN

+! 15MG/VIAL N021106 002 Mar 25, 2003 Jan CAHN

+! 20MG/VIAL N021106 003 Mar 25, 2003 Jan CAHN

+! 25MG/VIAL N021106 004 Jul 31, 2014 Jan CAHN

+! 30MG/VIAL N021106 005 Jul 31, 2014 Jan CAHN

PEMETREXED

>A> SOLUTION;INTRAVENOUS

>A> PEMFEXY

>A> +! EAGLE PHARMS 500MG/20ML (25MG/ML) N209472 001 Feb 08, 2020 Feb NEWA

PENICILLAMINE

CAPSULE;ORAL

PENICILLAMINE

AB ANI PHARMS INC 250MG A209921 001 May 07, 2019 Jan CAHN

TABLET;ORAL

PENICILLAMINE

>A> AB TEVA PHARMS USA 250MG A211497 001 Feb 13, 2020 Feb NEWA

PERMETHRIN

CREAM; TOPICAL

&gt;D&gt; ELIMITE

>D> AB	+!	MYLAN	5%	N019855	001	Aug 25, 1989	Feb	DISC
>A>	+ @		5%	N019855	001	Aug 25, 1989	Feb	DISC

PERMETHRIN

>D> AB		PERRIGO ISRAEL	5%	A076369	001	Apr 21, 2003	Feb	CHRS
>A> AB	!		5%	A076369	001	Apr 21, 2003	Feb	CHRS

PHENYLEPHRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

PHENYLEPHRINE HYDROCHLORIDE

AP1		RICONPHARMA LLC	10MG/ML (10MG/ML)	A209967	001	Jan 16, 2020	Jan	NEWA
AP1			50MG/5ML (10MG/ML)	A209967	002	Jan 16, 2020	Jan	NEWA
AP1			100MG/10ML (10MG/ML)	A209967	003	Jan 16, 2020	Jan	NEWA

PILOCARPINE HYDROCHLORIDE

TABLET; ORAL

PILOCARPINE HYDROCHLORIDE

>D> AB		AUROBINDO PHARMA LTD	5MG	A212377	001	Aug 13, 2019	Feb	DISC
>A>	@		5MG	A212377	001	Aug 13, 2019	Feb	DISC
>D> AB			7.5MG	A212377	002	Aug 13, 2019	Feb	DISC
>A>	@		7.5MG	A212377	002	Aug 13, 2019	Feb	DISC

PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

PIOGLITAZONE HYDROCHLORIDE

@	MYLAN PHARMS INC	EQ 15MG BASE	A076801	001	Aug 17, 2012	Jan	DISC
@		EQ 30MG BASE	A076801	002	Aug 17, 2012	Jan	DISC
@		EQ 45MG BASE	A076801	003	Aug 17, 2012	Jan	DISC

PODOFILOX

SOLUTION; TOPICAL

CONDYLOX

AT	+!	TEVA BRANDED PHARM	0.5%	N019795	001	Dec 13, 1990	Jan	CAHN
----	----	--------------------	------	---------	-----	--------------	-----	------

POLYMYXIN B SULFATE

INJECTABLE; INJECTION

AEROSPORIN

@	GLAXOSMITHKLINE	EQ 500,000 UNITS BASE/VIAL	A062036	001		Jan	CMS1
---	-----------------	----------------------------	---------	-----	--	-----	------

POLYMYXIN B SULFATE

>D> AP		X GEN PHARMS	EQ 500,000 UNITS BASE/VIAL	A063000	001	Sep 30, 1994	Feb	CAHN
>A> AP		XGEN PHARMS	EQ 500,000 UNITS BASE/VIAL	A063000	001	Sep 30, 1994	Feb	CAHN

POTASSIUM CHLORIDE

FOR SUSPENSION, EXTENDED RELEASE; ORAL

MICRO-K LS

>D>	@	KV PHARM	20MEQ/PACKET	N019561	003	Aug 26, 1988	Feb	CRLD
>A>	+ @		20MEQ/PACKET	N019561	003	Aug 26, 1988	Feb	CRLD

TABLET, EXTENDED RELEASE; ORAL

POTASSIUM CHLORIDE

>A> AB2		GRANULES PHARMS	8MEQ	A211797	001	Mar 04, 2020	Feb	NEWA
>A> AB2			10MEQ	A211797	002	Mar 04, 2020	Feb	NEWA
>D>	@	PII	8MEQ	A206881	001	Jan 22, 2019	Feb	CAHN
>D>	@		10MEQ	A206881	002	Jan 22, 2019	Feb	CAHN
>D>	@		10MEQ	A210097	001	Jun 17, 2019	Feb	CAHN
>D>	@		20MEQ	A210098	001	Apr 26, 2019	Feb	CAHN
>A>	@	STRIDES PHARMA	8MEQ	A206881	001	Jan 22, 2019	Feb	CAHN
>A>	@		10MEQ	A206881	002	Jan 22, 2019	Feb	CAHN
>A>	@		10MEQ	A210097	001	Jun 17, 2019	Feb	CAHN
>A>	@		20MEQ	A210098	001	Apr 26, 2019	Feb	CAHN

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

>D> AB		APOTEX INC	0.125MG	A090151	001	Apr 30, 2012	Feb	CAHN
>D> AB			0.25MG	A090151	002	Apr 30, 2012	Feb	CAHN
>D> AB			0.5MG	A090151	003	Apr 30, 2012	Feb	CAHN
>D> AB			0.75MG	A090151	006	Apr 30, 2012	Feb	CAHN
>D> AB			1MG	A090151	004	Apr 30, 2012	Feb	CAHN
>D> AB			1.5MG	A090151	005	Apr 30, 2012	Feb	CAHN

TABLET;ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

	@ MYLAN	0.125MG	A077854	001	Oct 08, 2010	Jan	DISC
	@	0.25MG	A077854	002	Oct 08, 2010	Jan	DISC
	@	0.5MG	A077854	003	Oct 08, 2010	Jan	DISC
	@	0.75MG	A090764	001	Apr 09, 2010	Jan	DISC
	@	1MG	A077854	004	Oct 08, 2010	Jan	DISC
	@	1.5MG	A077854	005	Oct 08, 2010	Jan	DISC
>A>	AB ZENNOVA	0.125MG	A090151	001	Apr 30, 2012	Feb	CAHN
>A>	AB	0.25MG	A090151	002	Apr 30, 2012	Feb	CAHN
>A>	AB	0.5MG	A090151	003	Apr 30, 2012	Feb	CAHN
>A>	AB	0.75MG	A090151	006	Apr 30, 2012	Feb	CAHN
>A>	AB	1MG	A090151	004	Apr 30, 2012	Feb	CAHN
>A>	AB	1.5MG	A090151	005	Apr 30, 2012	Feb	CAHN

PREDNICARBATE

OINTMENT;TOPICAL

DERMATOP

>D>	AB +!	VALEANT PHARMS NORTH	0.1%	N019568	001	Sep 23, 1991	Feb	DISC
>A>		+ @	0.1%	N019568	001	Sep 23, 1991	Feb	DISC
		PREDNICARBATE						
>D>	AB	FOUGERA PHARMS	0.1%	A077236	001	Mar 09, 2007	Feb	CHRS
>A>		!	0.1%	A077236	001	Mar 09, 2007	Feb	CTEC
>A>	AB	!	0.1%	A077236	001	Mar 09, 2007	Feb	CHRS

PREDNISOLONE ACETATE

SUSPENSION/DROPS;OPHTHALMIC

OMNIPRED

AB	+!	NOVARTIS	1%	N017469	001		Jan	CHRS
AB	+		1%	N017469	001		Jan	CRLD

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION;ORAL

PREDNISOLONE SODIUM PHOSPHATE

>A>		PHARM ASSOC	EQ 30MG BASE/5ML	A204962	001	Mar 11, 2020	Feb	NEWA
-----	--	-------------	------------------	---------	-----	--------------	-----	------

PREGABALIN

CAPSULE;ORAL

PREGABALIN

>D>	AB	MYLAN	25MG	A091228	001	Sep 20, 2019	Feb	DISC
>A>		@	25MG	A091228	001	Sep 20, 2019	Feb	DISC
>D>	AB		50MG	A091228	002	Sep 20, 2019	Feb	DISC
>A>		@	50MG	A091228	002	Sep 20, 2019	Feb	DISC
>D>	AB		75MG	A091228	003	Sep 20, 2019	Feb	DISC
>A>		@	75MG	A091228	003	Sep 20, 2019	Feb	DISC
>D>	AB		100MG	A091228	004	Sep 20, 2019	Feb	DISC
>A>		@	100MG	A091228	004	Sep 20, 2019	Feb	DISC
>D>	AB		150MG	A091228	005	Sep 20, 2019	Feb	DISC
>A>		@	150MG	A091228	005	Sep 20, 2019	Feb	DISC
>D>	AB		200MG	A091228	006	Sep 20, 2019	Feb	DISC
>A>		@	200MG	A091228	006	Sep 20, 2019	Feb	DISC
>D>	AB		225MG	A091228	007	Sep 20, 2019	Feb	DISC
>A>		@	225MG	A091228	007	Sep 20, 2019	Feb	DISC
>D>	AB		300MG	A091228	008	Sep 20, 2019	Feb	DISC
>A>		@	300MG	A091228	008	Sep 20, 2019	Feb	DISC

PROMETHAZINE HYDROCHLORIDE

INJECTABLE;INJECTION

PROMETHAZINE HYDROCHLORIDE

>D>	AP	X-GEN PHARMS	25MG/ML	A040737	001	Apr 24, 2008	Feb	CAHN
>D>	AP		50MG/ML	A040737	002	Apr 24, 2008	Feb	CAHN
>A>	AP	XGEN PHARMS	25MG/ML	A040737	001	Apr 24, 2008	Feb	CAHN
>A>	AP		50MG/ML	A040737	002	Apr 24, 2008	Feb	CAHN

PYRAZINAMIDE

TABLET;ORAL

PYRAZINAMIDE

>D>	AB	AKORN	500MG	A081319	001	Jun 30, 1992	Feb	CTEC
>A>		!	500MG	A081319	001	Jun 30, 1992	Feb	CHRS
>A>			500MG	A081319	001	Jun 30, 1992	Feb	CTEC
>D>	AB	!	DAVA PHARMS INC	500MG	A080157	001	Feb	DISC
>A>		@		500MG	A080157	001	Feb	DISC



PYRIMETHAMINE

TABLET;ORAL  
DARAPRIM

>D>	+	VYERA PHARMS LLC	25MG	N008578	001		Feb	CFTG
>A>	AB	+	25MG	N008578	001		Feb	CFTG
>A>		PYRIMETHAMINE						
>A>	AB	CEROVENE INC	25MG	A207127	001	Feb 28, 2020	Feb	NFTG

QUETIAPINE FUMARATE

TABLET;ORAL  
QUETIAPINE FUMARATE

>D>	AB	APOTEX INC	EQ 25MG BASE	A090960	001	Mar 27, 2012	Feb	CAHN
>D>	AB		EQ 50MG BASE	A090960	002	Mar 27, 2012	Feb	CAHN
>D>	AB		EQ 100MG BASE	A090960	003	Mar 27, 2012	Feb	CAHN
>D>	AB		EQ 200MG BASE	A090960	004	Mar 27, 2012	Feb	CAHN
>D>	AB		EQ 300MG BASE	A090960	005	Mar 27, 2012	Feb	CAHN
>D>	AB		EQ 400MG BASE	A090960	006	Mar 27, 2012	Feb	CAHN
>A>	AB	ZENNOVA	EQ 25MG BASE	A090960	001	Mar 27, 2012	Feb	CAHN
>A>	AB		EQ 50MG BASE	A090960	002	Mar 27, 2012	Feb	CAHN
>A>	AB		EQ 100MG BASE	A090960	003	Mar 27, 2012	Feb	CAHN
>A>	AB		EQ 200MG BASE	A090960	004	Mar 27, 2012	Feb	CAHN
>A>	AB		EQ 300MG BASE	A090960	005	Mar 27, 2012	Feb	CAHN
>A>	AB		EQ 400MG BASE	A090960	006	Mar 27, 2012	Feb	CAHN

RANITIDINE HYDROCHLORIDE

TABLET;ORAL  
RANITIDINE HYDROCHLORIDE

		@ AMNEAL PHARMS NY	EQ 150MG BASE	A077824	001	Oct 13, 2006	Jan	DISC
		@	EQ 300MG BASE	A077824	002	Oct 13, 2006	Jan	DISC
		@ STRIDES PHARMA	EQ 150MG BASE	A210010	001	Aug 01, 2018	Jan	DISC
AB	!	@	EQ 300MG BASE	A205512	002	Aug 22, 2016	Jan	CHRS
		@	EQ 300MG BASE	A210010	002	Aug 01, 2018	Jan	DISC

RANOLAZINE

TABLET, EXTENDED RELEASE;ORAL  
RANOLAZINE

AB		ANI PHARMS INC	500MG	A210482	001	Oct 29, 2019	Jan	CAHN
AB			1GM	A210482	002	Oct 29, 2019	Jan	CAHN
AB		MANKIND PHARMA	500MG	A212284	001	Feb 12, 2020	Jan	NEWA
AB			1GM	A212284	002	Feb 12, 2020	Jan	NEWA
>A>	AB	MICRO LABS	500MG	A211745	001	Feb 27, 2020	Feb	NEWA
>A>	AB		1GM	A211745	002	Feb 27, 2020	Feb	NEWA

RIBAVIRIN

CAPSULE;ORAL  
RIBASPHERE

>D>	AB	KADMON PHARMS LLC	200MG	A076203	001	Apr 06, 2004	Feb	DISC
>A>		@	200MG	A076203	001	Apr 06, 2004	Feb	DISC

TABLET;ORAL  
RIBAVIRIN

>D>	AB	BEXIMCO PHARMS USA	200MG	A202546	001	Aug 12, 2014	Feb	DISC
>A>		@	200MG	A202546	001	Aug 12, 2014	Feb	DISC
>D>	AB		400MG	A202546	002	Aug 12, 2014	Feb	DISC
>A>		@	400MG	A202546	002	Aug 12, 2014	Feb	DISC
>D>	AB		500MG	A202546	003	Aug 12, 2014	Feb	DISC
>A>		@	500MG	A202546	003	Aug 12, 2014	Feb	DISC
>D>	AB		600MG	A202546	004	Aug 12, 2014	Feb	DISC
>A>		@	600MG	A202546	004	Aug 12, 2014	Feb	DISC
>D>	AB	KADMON PHARMS LLC	200MG	A077456	001	Dec 05, 2005	Feb	DISC
>A>		@	200MG	A077456	001	Dec 05, 2005	Feb	DISC
>D>	AB		400MG	A077456	002	Dec 05, 2005	Feb	DISC
>A>		@	400MG	A077456	002	Dec 05, 2005	Feb	DISC
>D>	AB	!	600MG	A077456	003	Dec 05, 2005	Feb	DISC
>A>		@	600MG	A077456	003	Dec 05, 2005	Feb	DISC
>D>	AB	ZYDUS PHARMS USA	400MG	A077094	002	Mar 16, 2007	Feb	DISC
>A>		@	400MG	A077094	002	Mar 16, 2007	Feb	DISC
>D>	AB		500MG	A077094	004	Apr 18, 2008	Feb	DISC
>A>		@	500MG	A077094	004	Apr 18, 2008	Feb	DISC
>D>	AB		600MG	A077094	003	Mar 16, 2007	Feb	DISC
>A>		@	600MG	A077094	003	Mar 16, 2007	Feb	DISC

>A> RIMEGEPANT SULFATE  
 >A> TABLET, ORALLY DISINTEGRATING;ORAL  
 >A> NURTEC ODT  
 >A> +! BIOHAVEN PHARM EQ 75MG BASE N212728 001 Feb 27, 2020 Feb NEWA

RITODRINE HYDROCHLORIDE  
 INJECTABLE; INJECTION  
 YUTOPAR  
 @ ASTRAZENECA 15MG/ML N018580 002 Sep 27, 1984 Jan CMS1

RIZATRIPTAN BENZOATE  
 TABLET, ORALLY DISINTEGRATING;ORAL  
 RIZATRIPTAN BENZOATE  
 >D> AB MYLAN PHARMS INC EQ 5MG BASE A078173 001 Dec 31, 2012 Feb DISC  
 >A> @ EQ 5MG BASE A078173 001 Dec 31, 2012 Feb DISC  
 >D> AB EQ 10MG BASE A078173 002 Dec 31, 2012 Feb DISC  
 >A> @ EQ 10MG BASE A078173 002 Dec 31, 2012 Feb DISC

ROPINIROLE HYDROCHLORIDE  
 TABLET;ORAL  
 ROPINIROLE HYDROCHLORIDE  
 >D> AB HIKMA EQ 0.25MG BASE A077852 001 May 05, 2008 Feb DISC  
 >A> @ EQ 0.25MG BASE A077852 001 May 05, 2008 Feb DISC  
 >D> AB EQ 0.5MG BASE A077852 002 May 05, 2008 Feb DISC  
 >A> @ EQ 0.5MG BASE A077852 002 May 05, 2008 Feb DISC  
 >D> AB EQ 1MG BASE A077852 003 May 05, 2008 Feb DISC  
 >A> @ EQ 1MG BASE A077852 003 May 05, 2008 Feb DISC  
 >D> AB EQ 2MG BASE A077852 004 May 05, 2008 Feb DISC  
 >A> @ EQ 2MG BASE A077852 004 May 05, 2008 Feb DISC  
 >D> AB EQ 3MG BASE A077852 005 May 05, 2008 Feb DISC  
 >A> @ EQ 3MG BASE A077852 005 May 05, 2008 Feb DISC  
 >D> AB EQ 4MG BASE A077852 006 May 05, 2008 Feb DISC  
 >A> @ EQ 4MG BASE A077852 006 May 05, 2008 Feb DISC  
 >D> AB EQ 5MG BASE A077852 007 May 19, 2008 Feb DISC  
 >A> @ EQ 5MG BASE A077852 007 May 19, 2008 Feb DISC

SELEGILINE HYDROCHLORIDE  
 CAPSULE;ORAL  
 SELEGILINE HYDROCHLORIDE  
 >D> AB DAVA PHARMS INC 5MG A075352 001 Nov 30, 1998 Feb DISC  
 >A> @ 5MG A075352 001 Nov 30, 1998 Feb DISC

SELEXIPAG  
 TABLET;ORAL  
 UPTRAVI  
 +! ACTELION PHARMS LTD 0.4MG N207947 002 Dec 21, 2015 Jan CHRS  
 + 1.6MG N207947 008 Dec 21, 2015 Jan CHRS

SEMAGLUTIDE  
 SOLUTION;SUBCUTANEOUS  
 OZEMPIC  
 +! NOVO 4MG/3ML (1.34MG/ML) N209637 002 Apr 09, 2019 Jan NEWA

SILDENAFIL CITRATE  
 TABLET;ORAL  
 SILDENAFIL CITRATE  
 >A> AB APPCO 25MG A207178 001 Mar 02, 2020 Feb NEWA  
 >A> AB 50MG A207178 002 Mar 02, 2020 Feb NEWA  
 >A> AB 100MG A207178 003 Mar 02, 2020 Feb NEWA

SILODOSIN  
 CAPSULE;ORAL  
 SILODOSIN  
 >A> AB HETERO LABS LTD V 4MG A204793 001 Feb 14, 2020 Feb NEWA  
 >A> AB 8MG A204793 002 Feb 14, 2020 Feb NEWA

SODIUM IODIDE I-123

CAPSULE;ORAL

SODIUM IODIDE I 123

>A>	AA	CURIUM	100uCi	A 071909	001	Feb 28, 1989	Feb CAHN
>A>	AA		200uCi	A 071910	001	Feb 28, 1989	Feb CAHN
>D>	AA	MALLINKRODT NUCLEAR	100uCi	A 071909	001	Feb 28, 1989	Feb CAHN
>D>	AA		200uCi	A 071910	001	Feb 28, 1989	Feb CAHN

SODIUM IODIDE I-131

SOLUTION;ORAL

HICON

AA	+!	JUBILANT DRAXIMAGE	250-1000mCi	N021305	007	Dec 05, 2011	Jan CFTG
		SODIUM IODIDE I 131					
AA		INTL ISOTOPES	250-1000mCi	A209166	001	Feb 05, 2020	Jan NFTG

SODIUM PHENYL BUTYRATE

TABLET;ORAL

SODIUM PHENYL BUTYRATE

@ ALVOGEN

500MG

A 090910 001 Nov 18, 2011 Jan DISC

SODIUM POLYSTYRENE SULFONATE

SUSPENSION;ORAL, RECTAL

SODIUM POLYSTYRENE SULFONATE

>D>	AA	HIKMA	15GM/60ML	A 089049	001	Nov 17, 1986	Feb DISC
>A>		@	15GM/60ML	A 089049	001	Nov 17, 1986	Feb DISC

SOLRIAMFETOL HYDROCHLORIDE

TABLET;ORAL

SUNOSI

+	JAZZ	EQ 75MG BASE	N211230	001	Jun 17, 2019	Jan CPOT
+		75MG	N211230	001	Jun 17, 2019	Jan CAIN
+		EQ 150MG BASE	N211230	002	Jun 17, 2019	Jan CPOT
+		150MG	N211230	002	Jun 17, 2019	Jan CAIN

SOMATROPIN

INJECTABLE;INJECTION

ACCRETROPIN

@ EMERGENT

5MG/ML (5MG/ML)

N 021538 001 Jan 23, 2008 Jan CAIN

HUMATROPE

@ LILLY

2MG/VIAL

N 019640 001 Jun 23, 1987 Jan CAIN

BX	+!		5MG/VIAL	N 019640	004	Mar 08, 1987	Jan CAIN
BX	+		6MG/VIAL	N 019640	005	Feb 04, 1999	Jan CAIN
	+!		12MG/VIAL	N 019640	006	Feb 04, 1999	Jan CAIN
	+!		24MG/VIAL	N 019640	007	Feb 04, 1999	Jan CAIN

STREPTOMYCIN SULFATE

INJECTABLE;INJECTION

STREPTOMYCIN SULFATE

>D>	!	X GEN PHARMS	EQ 1GM BASE/VIAL	A 064210	001	Jun 30, 1998	Feb CAHN
>A>	!	XGEN PHARMS	EQ 1GM BASE/VIAL	A 064210	001	Jun 30, 1998	Feb CAHN

SUMATRIPTAN SUCCINATE

TABLET;ORAL

SUMATRIPTAN SUCCINATE

>D>	AB	APOTEX INC	EQ 25MG BASE	A 200263	001	Jun 19, 2012	Feb CAHN
>D>	AB		EQ 50MG BASE	A 200263	002	Jun 19, 2012	Feb CAHN
>D>	AB		EQ 100MG BASE	A 200263	003	Jun 19, 2012	Feb CAHN
>A>	AB	COREPHARMA	EQ 25MG BASE	A 200263	001	Jun 19, 2012	Feb CAHN
>A>	AB		EQ 50MG BASE	A 200263	002	Jun 19, 2012	Feb CAHN
>A>	AB		EQ 100MG BASE	A 200263	003	Jun 19, 2012	Feb CAHN

TAPENTADOL HYDROCHLORIDE

SOLUTION;ORAL

NUCYNTA

>D>	+	@ ASSERTIO	EQ 20MG BASE/ML	N 203794	001	Oct 15, 2012	Feb CAHN
>A>	+	@ COLLEGIUM PHARM INC	EQ 20MG BASE/ML	N 203794	001	Oct 15, 2012	Feb CAHN

TABLET;ORAL

NUCYNTA

>A>	+	COLLEGIUM PHARM INC	EQ 50MG BASE	N 022304	001	Nov 20, 2008	Feb CAHN
>A>	+		EQ 75MG BASE	N 022304	002	Nov 20, 2008	Feb CAHN
>A>	+!		EQ 100MG BASE	N 022304	003	Nov 20, 2008	Feb CAHN

## TABLET;ORAL

## NUCYNTA

>D>	+	DEPO NF	EQ 50MG BASE	N022304	001	Nov 20, 2008	Feb CAHN
>D>	+		EQ 75MG BASE	N022304	002	Nov 20, 2008	Feb CAHN
>D>	+!		EQ 100MG BASE	N022304	003	Nov 20, 2008	Feb CAHN

## TABLET, EXTENDED RELEASE;ORAL

## NUCYNTA ER

>A>	+	COLLEGIUM PHARM INC	EQ 50MG BASE	N200533	001	Aug 25, 2011	Feb CAHN
>A>	+		EQ 100MG BASE	N200533	002	Aug 25, 2011	Feb CAHN
>A>	+		EQ 150MG BASE	N200533	003	Aug 25, 2011	Feb CAHN
>A>	+		EQ 200MG BASE	N200533	004	Aug 25, 2011	Feb CAHN
>A>	+!		EQ 250MG BASE	N200533	005	Aug 25, 2011	Feb CAHN
>D>	+	DEPO NF	EQ 50MG BASE	N200533	001	Aug 25, 2011	Feb CAHN
>D>	+		EQ 100MG BASE	N200533	002	Aug 25, 2011	Feb CAHN
>D>	+		EQ 150MG BASE	N200533	003	Aug 25, 2011	Feb CAHN
>D>	+		EQ 200MG BASE	N200533	004	Aug 25, 2011	Feb CAHN
>D>	+!		EQ 250MG BASE	N200533	005	Aug 25, 2011	Feb CAHN

TAZEMETOSTAT HYDROBROMIDE

## TABLET;ORAL

## TAZVERIK

+!	EPIZYME INC	EQ 200MG BASE	N211723	001	Jan 23, 2020	Jan NEWA
----	-------------	---------------	---------	-----	--------------	----------

TECHNETIUM TC-99M MERTIATIDE KIT

## INJECTABLE;INJECTION

## TECHNETIUM TC99M MERTIATIDE KIT

AP	SOMMER PHARMS II LLC	N/A	A206489	001	Feb 06, 2020	Jan NEWA
----	----------------------	-----	---------	-----	--------------	----------

TECHNETIUM TC-99M SESTAMIBI KIT

## INJECTABLE;INJECTION

## TECHNETIUM TC 99M SESTAMIBI

>A>	AP	CURIUM	N/A	A078098	001	Sep 22, 2008	Feb CAHN
>D>	AP	MALLINKRODT NUCLEAR	N/A	A078098	001	Sep 22, 2008	Feb CAHN

TELMISARTAN

## TABLET;ORAL

## TELMISARTAN

@	JUBILANT GENERICS	20MG	A204164	001	Aug 22, 2016	Jan DISC
---	-------------------	------	---------	-----	--------------	----------

@		40MG	A204164	002	Aug 22, 2016	Jan DISC
---	--	------	---------	-----	--------------	----------

@		80MG	A204164	003	Aug 22, 2016	Jan DISC
---	--	------	---------	-----	--------------	----------

TENOFOVIR DISOPROXIL FUMARATE

## TABLET;ORAL

## TENOFOVIR DISOPROXIL FUMARATE

>D>	AB	MYLAN	150MG	A206569	001	Nov 27, 2018	Feb DISC
>A>	@		150MG	A206569	001	Nov 27, 2018	Feb DISC
>D>	AB		200MG	A206569	002	Nov 27, 2018	Feb DISC
>A>	@		200MG	A206569	002	Nov 27, 2018	Feb DISC
>D>	AB		250MG	A206569	003	Nov 27, 2018	Feb DISC
>A>	@		250MG	A206569	003	Nov 27, 2018	Feb DISC
>D>	AB		300MG	A206569	004	Nov 27, 2018	Feb DISC
>A>	@		300MG	A206569	004	Nov 27, 2018	Feb DISC

TERAZOSIN HYDROCHLORIDE

## CAPSULE;ORAL

## TERAZOSIN HYDROCHLORIDE

>D>	AB	HERITAGE PHARMA	EQ 1MG BASE	A075614	002	Jan 30, 2001	Feb DISC
>A>	@		EQ 1MG BASE	A075614	002	Jan 30, 2001	Feb DISC
>D>	AB		EQ 2MG BASE	A075614	001	Jan 30, 2001	Feb DISC
>A>	@		EQ 2MG BASE	A075614	001	Jan 30, 2001	Feb DISC
>D>	AB		EQ 5MG BASE	A075614	003	Jan 30, 2001	Feb DISC
>A>	@		EQ 5MG BASE	A075614	003	Jan 30, 2001	Feb DISC
>D>	AB		EQ 10MG BASE	A075614	004	Jan 30, 2001	Feb DISC
>A>	@		EQ 10MG BASE	A075614	004	Jan 30, 2001	Feb DISC

TERIFLUNOMIDE

## TABLET;ORAL

## TERIFLUNOMIDE

>A>	AB	MYLAN	7MG	A209702	001	Feb 28, 2020	Feb NEWA
>A>	AB		14MG	A209702	002	Feb 28, 2020	Feb NEWA

TERIPARATIDE

INJECTABLE; SUBCUTANEOUS  
FORTEO

+!	LILLY	0.6MG/2.4ML (0.25MG/ML)	N021318	002	Jun 25, 2008	Jan	CAIN
@		0.75MG/3ML (0.25MG/ML)	N021318	001	Nov 26, 2002	Jan	CAIN

SOLUTION; SUBCUTANEOUS  
BONSITY

	ALVOGEN	0.62MG/2.48ML (0.25MG/ML)	N211939	001	Oct 04, 2019	Jan	CAIN
+!	LILLY	0.6MG/2.4ML (0.25MG/ML)	N021318	002	Jun 25, 2008	Jan	CDFR
@		0.75MG/3ML (0.25MG/ML)	N021318	001	Nov 26, 2002	Jan	CDFR

TETRABENAZINE

TABLET; ORAL

	TETRABENAZINE						
AB	PIRAMAL HLTHCARE UK	12.5MG	A213316	001	Jan 22, 2020	Jan	NEWA
AB		25MG	A213316	002	Jan 22, 2020	Jan	NEWA

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

TETRACYCLINE HYDROCHLORIDE

>A>	AB	STRIDES PHARMA	250MG	A212635	001	Mar 03, 2020	Feb	NEWA
>A>	AB		500MG	A212635	002	Mar 03, 2020	Feb	NEWA

THALLOUS CHLORIDE TL-201

INJECTABLE; INTRAVENOUS

THALLOUS CHLORIDE TL 201

>A>	@	CURIUM	2mCi/ML	A077698	001	Nov 09, 2006	Feb	CAHN
>D>	@	MALLINKRODT NUCLEAR	2mCi/ML	A077698	001	Nov 09, 2006	Feb	CAHN

THIOTEPA

POWDER; INTRAVENOUS

TEPADINA

>D>	+!	ADIENNE SA	15MG/VIAL	N208264	001	Jan 26, 2017	Feb	CFTG
>A>	AP	+!	15MG/VIAL	N208264	001	Jan 26, 2017	Feb	CFTG
>D>	+!		100MG/VIAL	N208264	002	Jan 26, 2017	Feb	CFTG
>A>	AP	+!	100MG/VIAL	N208264	002	Jan 26, 2017	Feb	CFTG
>A>		THIOTEPA						
>A>	AP	MSN	15MG/VIAL	A213049	001	Mar 04, 2020	Feb	NFTG
>A>	AP		100MG/VIAL	A213049	002	Mar 04, 2020	Feb	NFTG

TOBRAMYCIN

SOLUTION; INHALATION

TOBRAMYCIN

>D>	AN	MYLAN	300MG/5ML	A209554	001	Oct 13, 2017	Feb	DISC
>A>	@		300MG/5ML	A209554	001	Oct 13, 2017	Feb	DISC

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

TOBRAMYCIN SULFATE

>D>	AP	!	X GEN PHARMS	EQ 1.2GM BASE/VIAL	A065013	001	Aug 17, 2001	Feb	CAHN
>A>	AP	!	XGEN PHARMS	EQ 1.2GM BASE/VIAL	A065013	001	Aug 17, 2001	Feb	CAHN

TOLTERODINE TARTRATE

TABLET; ORAL

TOLTERODINE TARTRATE

	@	MYLAN PHARMS INC	1MG	A202641	001	Nov 27, 2012	Jan	DISC
	@		2MG	A202641	002	Nov 27, 2012	Jan	DISC
AB		UNIQUE PHARM LABS	1MG	A204721	001	Jan 24, 2020	Jan	NEWA
AB			2MG	A204721	002	Jan 24, 2020	Jan	NEWA

TORSEMIDE

TABLET; ORAL

TORSEMIDE

>D>	AB	APOTEX INC	5MG	A076894	001	May 31, 2005	Feb	CAHN
>D>	AB		10MG	A076894	002	May 31, 2005	Feb	CAHN
>D>	AB		20MG	A076894	003	May 31, 2005	Feb	CAHN
>D>	AB		100MG	A076894	004	May 31, 2005	Feb	CAHN
>A>	AB	COREPHARMA	5MG	A076894	001	May 31, 2005	Feb	CAHN
>A>	AB		10MG	A076894	002	May 31, 2005	Feb	CAHN
>A>	AB		20MG	A076894	003	May 31, 2005	Feb	CAHN

	TABLET;ORAL						
	TORSEMIDE						
>A>	AB	100MG	A076894	004	May 31, 2005	Feb	CAHN
	<u>TRANEXAMIC ACID</u>						
	INJECTABLE;INJECTION						
	TRANEXAMIC ACID						
	@	CAPLIN	100MG/ML	A212360	001	Jul 17, 2019	Jan DISC
>D>	AP	X-GEN PHARMS INC	100MG/ML	A201580	001	Jun 14, 2013	Feb CAHN
>A>	AP	XGEN PHARMS	100MG/ML	A201580	001	Jun 14, 2013	Feb CAHN
	TABLET;ORAL						
	TRANEXAMIC ACID						
	@	ANI PHARMS INC	650MG	A203256	001	Jul 25, 2016	Jan CAHN
	<u>TRAZODONE HYDROCHLORIDE</u>						
	TABLET;ORAL						
	TRAZODONE HYDROCHLORIDE						
AB	AUROLIFE PHARMA LLC	50MG	A204852	001	Feb 05, 2020	Jan	NEWA
AB		100MG	A204852	002	Feb 05, 2020	Jan	NEWA
AB		150MG	A204852	003	Feb 05, 2020	Jan	NEWA
AB		300MG	A204852	004	Feb 05, 2020	Jan	NEWA
	<u>TRIAZOLAM</u>						
	TABLET;ORAL						
	TRIAZOLAM						
	@	MYLAN PHARMS INC	0.125MG	A074031	001	Mar 25, 1994	Jan DISC
	@		0.25MG	A074031	002	Mar 25, 1994	Jan DISC
	<u>TRIENTINE HYDROCHLORIDE</u>						
	CAPSULE;ORAL						
	TRIENTINE HYDROCHLORIDE						
>A>	AB	RISING	250MG	A212238	001	Feb 20, 2020	Feb NEWA
	<u>URSODIOL</u>						
	CAPSULE;ORAL						
	URSODIOL						
AB	RISING	300MG	A213200	001	Feb 12, 2020	Jan	NEWA
	<u>VALACYCLOVIR HYDROCHLORIDE</u>						
	TABLET;ORAL						
	VALACYCLOVIR HYDROCHLORIDE						
	@	TEVA PHARMS	EQ 500MG BASE	A077655	001	May 24, 2010	Jan DISC
	@		EQ 1GM BASE	A077655	002	May 24, 2010	Jan DISC
	<u>VALGANCICLOVIR HYDROCHLORIDE</u>						
	FOR SOLUTION;ORAL						
	VALGANCICLOVIR HYDROCHLORIDE						
AB	GRANULES PHARMS	50MG/ML	A213306	001	Jan 31, 2020	Jan	NEWA
	<u>VANCOMYCIN HYDROCHLORIDE</u>						
	INJECTABLE;INJECTION						
	VANCOMYCIN HYDROCHLORIDE						
AP	PHARM ASSOC	EQ 500MG BASE/VIAL	A065401	001	Jun 30, 2008	Jan	CAHN
AP		EQ 1GM BASE/VIAL	A065401	002	Jun 30, 2008	Jan	CAHN
	<u>VARDENAFIL HYDROCHLORIDE</u>						
	TABLET, ORALLY DISINTEGRATING;ORAL						
	VARDENAFIL HYDROCHLORIDE						
>A>	AB	MACLEODS PHARMS LTD	10MG	A205988	001	Mar 10, 2020	Feb NEWA
	<u>VINCRISTINE SULFATE</u>						
	INJECTABLE;INJECTION						
	VINCRISTINE SULFATE PFS						
>D>	!	HOSPIRA	1MG/ML	A071484	001	Apr 19, 1988	Feb CTEC
>A>	AP	!	1MG/ML	A071484	001	Apr 19, 1988	Feb CTEC
>D>	@	TEVA PHARMS USA	1MG/ML	A075493	001	Sep 01, 1999	Feb CMFD
>A>	AP		1MG/ML	A075493	001	Sep 01, 1999	Feb CMFD

ZOLEDRONIC ACIDINJECTABLE; INTRAVENOUS  
ZOLEDRONIC ACID

>A>	AP	INGENUS PHARMS LLC	EQ 4MG BASE/5ML	A208968	001	Feb 19, 2020	Feb NEWA
-----	----	--------------------	-----------------	---------	-----	--------------	----------

ZOLMITRIPTANTABLET, ORALLY DISINTEGRATING; ORAL  
ZOLMITRIPTAN

>D>	AB	MYLAN	2.5MG	A202855	001	Sep 20, 2019	Feb DISC
>A>		@	2.5MG	A202855	001	Sep 20, 2019	Feb DISC
>D>	AB		5MG	A202855	002	Sep 20, 2019	Feb DISC
>A>		@	5MG	A202855	002	Sep 20, 2019	Feb DISC

>A>	<u>ACETAMINOPHEN; IBUPROFEN</u>						
>A>	TABLET;ORAL						
>A>	ADVIL DUAL ACTION WITH ACETAMINOPHEN						
>A>	+! PFIZER INC	250MG;125MG		N211733	001	Feb 28, 2020	Feb NEWA
	<u>FAMOTIDINE</u>						
	TABLET;ORAL						
	FAMOTIDINE						
	DR REDDYS LABS LTD	10MG		A077367	002	Aug 17, 2001	Jan CMS1
	@ MYLAN	10MG		A075674	001	Dec 21, 2001	Jan DISC
	<u>FEXOFENADINE HYDROCHLORIDE</u>						
	TABLET;ORAL						
	FEXOFENADINE HYDROCHLORIDE						
>A>	L PERRIGO CO	60MG		A212971	001	Feb 24, 2020	Feb NEWA
>A>		180MG		A212971	002	Feb 24, 2020	Feb NEWA
	<u>LEVOCETIRIZINE DIHYDROCHLORIDE</u>						
	TABLET;ORAL						
	LEVOCETIRIZINE DIHYDROCHLORIDE						
>D>	@ PERRIGO R AND D	5MG		A211983	001	Mar 28, 2019	Feb CMFD
>A>		5MG		A211983	001	Mar 28, 2019	Feb CMFD
	<u>LEVONORGESTREL</u>						
	TABLET;ORAL						
	LEVONORGESTREL						
>D>	APOTEX	1.5MG		A205329	001	Sep 18, 2018	Feb CAHN
		1.5MG		A205329	001	Sep 18, 2018	Jan CMFD
	LABORATOIRE HRA	1.5MG		A204044	001	Jul 03, 2018	Jan CAHN
>A>	XIROMED	1.5MG		A205329	001	Sep 18, 2018	Feb CAHN
	<u>LORATADINE</u>						
	TABLET;ORAL						
	LORATADINE						
	@ MYLAN	10MG		A075790	001	Nov 07, 2008	Jan DISC
	@	10MG		A078447	001	Aug 12, 2011	Jan DISC
	<u>NICOTINE POLACRILEX</u>						
	TROCHE/LOZENGE;ORAL						
	NICOTINE POLACRILEX						
>A>	DR REDDYS LABS SA	EQ 2MG BASE		A212983	001	Feb 21, 2020	Feb NEWA
>A>		EQ 4MG BASE		A212983	002	Feb 21, 2020	Feb NEWA
	<u>POLYETHYLENE GLYCOL 3350</u>						
	FOR SOLUTION;ORAL						
	GLYCOLAX						
	LANNETT CO INC	17GM/PACKET		A090600	001	Oct 06, 2009	Jan CMFD
		17GM/SCOOPFUL		A090600	002	Oct 06, 2009	Jan CMFD



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

**CUMULATIVE SUPPLEMENT NUMBER 02 FEBRUARY 2020**

NO FEBRUARY 2020 APPROVALS

## **ORPHAN PRODUCT DESIGNATIONS AND APPROVALS LIST**

The list of Orphan Designations and Approvals is available at:

[https://www.fda.gov/industry/developing-products-rare-diseases-conditions/designating-orphan-product-drugs-and-biological-products.](https://www.fda.gov/industry/developing-products-rare-diseases-conditions/designating-orphan-product-drugs-and-biological-products)

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

NO FEBRUARY 2020 ADDITIONS

PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ACETAMINOPHEN; BUTALBITAL - BUTALBITAL AND ACETAMINOPHEN</u>						
A 213115	001				CGT	Jul 28, 2020
<u>AMISULPRIDE - BARHEMSYS</u>						
N 209510	001				>A> NCE	Feb 26, 2025
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360	001	>A> 10548943	Dec 16, 2029	U-2739		
		>A> 10548943	Dec 16, 2029	U-2740		
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360	002	>A> 10548943	Dec 16, 2029	U-2739		
		>A> 10548943	Dec 16, 2029	U-2740		
<u>ARIPIPRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971	001	10525057	Mar 08, 2034	U-1632		
		10525057	Mar 08, 2034	U-2723		
		10525057	Mar 08, 2034	U-543		
<u>ARIPIPRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971	002	10525057	Mar 08, 2034	U-1632		
		10525057	Mar 08, 2034	U-2723		
		10525057	Mar 08, 2034	U-543		
<u>ARIPIPRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971	003	10525057	Mar 08, 2034	U-1632		
		10525057	Mar 08, 2034	U-2723		
		10525057	Mar 08, 2034	U-543		
<u>ARIPIPRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971	004	10525057	Mar 08, 2034	U-1632		
		10525057	Mar 08, 2034	U-2723		
		10525057	Mar 08, 2034	U-543		
<u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202	001	10517507	Jun 13, 2032	DP		
<u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202	002	10517507	Jun 13, 2032	DP		
<u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202	003	10517507	Jun 13, 2032	DP		
<u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202	004	10517507	Jun 13, 2032	DP		
<u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202	005	10517507	Jun 13, 2032	DP		
<u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202	006	10517507	Jun 13, 2032	DP		
<u>AVAPRITINIB - AYPVAKIT</u>						
N 212608	001	9200002	Oct 15, 2034	DS DP U-2726	NCE	Jan 09, 2025
		9944651	Oct 15, 2034	DS DP U-2726		
		9994575	Oct 15, 2034	DS DP U-2726		
<u>AVAPRITINIB - AYPVAKIT</u>						
N 212608	002	9200002	Oct 15, 2034	DS DP U-2726	NCE	Jan 09, 2025
		9944651	Oct 15, 2034	DS DP U-2726		
		9994575	Oct 15, 2034	DS DP U-2726		

## PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>AVAPRITINIB - AYVAKIT</u>						
N 212608 002	9200002	Oct 15, 2034	DS DP U-2726		NCE	Jan 09, 2025
	9944651	Oct 15, 2034	DS DP U-2726			
	9994575	Oct 15, 2034	DS DP U-2726			
<u>AVAPRITINIB - AYVAKIT</u>						
N 212608 003	9200002	Oct 15, 2034	DS DP U-2726		NCE	Jan 09, 2025
	9944651	Oct 15, 2034	DS DP U-2726			
	9994575	Oct 15, 2034	DS DP U-2726			
<u>AVIBACTAM SODIUM; CEFTAZIDIME - AVYCAZ</u>						
N 206494 001	7112592	Jan 07, 2026	DS DP U-2244			
	7112592	Jan 07, 2026	DS DP U-2508			
	7112592	Jan 07, 2026	DS DP U-282			
<u>AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE - AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE</u>						
A 207712 001					>A> PC	Aug 29, 2020
<u>BEMPEDOIC ACID - NEXLETO</u>						
N 211616 001	>A> 10118881	Dec 23, 2023	U-2747		>A> NCE	Feb 21, 2025
	>A> 7335799	Dec 03, 2025	DS			
	>A> 8497301	Dec 23, 2023	U-2747			
	>A> 9000041	Dec 23, 2023	U-2747			
	>A> 9624152	Dec 23, 2023	U-2748			
<u>BEMPEDOIC ACID; EZETIMIBE - NEXLIZET</u>						
N 211617 001	>A> 10118881	Dec 23, 2023	U-2746		>A> NCE	Feb 21, 2025
	>A> 7335799	Dec 03, 2025	DS		>A> NP	Feb 26, 2023
	>A> 8497301	Dec 23, 2023	U-2746			
	>A> 9000041	Dec 23, 2023	U-2746			
	>A> 9624152	Dec 23, 2023	U-2749			
<u>BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - BIKTARVY</u>						
N 210251 001	>A> 10548846	Nov 08, 2036	DP			
<u>BRILLIANT BLUE G - TISSUEBLUE</u>						
N 209569 001					NCE ODE-282	Dec 20, 2024 Dec 20, 2026
<u>BUPRENORPHINE - SUBLOCADE</u>						
N 209819 001	>A> 10558394	Jun 25, 2031	DP			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 001	>A> 8513202	Dec 03, 2027	DS DP U-2441			
	>A> 8513202	Dec 03, 2027	DS DP U-2632			
	>A> 8513202	Dec 03, 2027	DS DP U-493			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 002	>A> 8513202	Dec 03, 2027	DS DP U-2441			
	>A> 8513202	Dec 03, 2027	DS DP U-2632			
	>A> 8513202	Dec 03, 2027	DS DP U-493			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 003	>A> 8513202	Dec 03, 2027	DS DP U-2441			
	>A> 8513202	Dec 03, 2027	DS DP U-2632			
	>A> 8513202	Dec 03, 2027	DS DP U-493			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 004	>A> 8513202	Dec 03, 2027	DS DP U-2441			
	>A> 8513202	Dec 03, 2027	DS DP U-2632			
	>A> 8513202	Dec 03, 2027	DS DP U-493			

## PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 001	>A> 7943582	Feb 26, 2029	DS DP U-2441			
	>A> 7943582	Feb 26, 2029	DS DP U-2632			
	>A> 7943582	Feb 26, 2029	DS DP U-493			
	>A> 8222219	Apr 11, 2025	U-2441			
	>A> 8222219	Apr 11, 2025	U-2632			
	>A> 8222219	Apr 11, 2025	U-493			
	>A> 8513202	Dec 03, 2027	DS DP U-2441			
	>A> 8513202	Dec 03, 2027	DS DP U-2632			
	>A> 8513202	Dec 03, 2027	DS DP U-493			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 002	>A> 7943582	Feb 26, 2029	DS DP U-2441			
	>A> 7943582	Feb 26, 2029	DS DP U-2632			
	>A> 7943582	Feb 26, 2029	DS DP U-493			
	>A> 8222219	Apr 11, 2025	U-2441			
	>A> 8222219	Apr 11, 2025	U-2632			
	>A> 8222219	Apr 11, 2025	U-493			
	>A> 8513202	Dec 03, 2027	DS DP U-2441			
	>A> 8513202	Dec 03, 2027	DS DP U-2632			
	>A> 8513202	Dec 03, 2027	DS DP U-493			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 003	>A> 7943582	Feb 26, 2029	DS DP U-2441			
	>A> 7943582	Feb 26, 2029	DS DP U-2632			
	>A> 7943582	Feb 26, 2029	DS DP U-493			
	>A> 8222219	Apr 11, 2025	U-2441			
	>A> 8222219	Apr 11, 2025	U-2632			
	>A> 8222219	Apr 11, 2025	U-493			
	>A> 8513202	Dec 03, 2027	DS DP U-2441			
	>A> 8513202	Dec 03, 2027	DS DP U-2632			
	>A> 8513202	Dec 03, 2027	DS DP U-493			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 004	>A> 7943582	Feb 26, 2029	DS DP U-2441			
	>A> 7943582	Feb 26, 2029	DS DP U-2632			
	>A> 7943582	Feb 26, 2029	DS DP U-493			
	>A> 8222219	Apr 11, 2025	U-2441			
	>A> 8222219	Apr 11, 2025	U-2632			
	>A> 8222219	Apr 11, 2025	U-493			
	>A> 8513202	Dec 03, 2027	DS DP U-2441			
	>A> 8513202	Dec 03, 2027	DS DP U-2632			
	>A> 8513202	Dec 03, 2027	DS DP U-493			
<u>CRISABOROLE - EUCRISA</u>						
N 207695 001	>A> 8039451	Jun 11, 2026	DS DP		>A> NCE	Dec 14, 2021
	>A> 8039451*PED	Dec 11, 2026			>A> PED	Jun 14, 2022
	>A> 8168614	Jan 20, 2030	U-1932			
	>A> 8168614*PED	Jul 20, 2030				
	>A> 8501712	Feb 16, 2027	U-1932			
	>A> 8501712*PED	Aug 16, 2027				
	>A> 9682092	Feb 16, 2027	U-1932			
	>A> 9682092*PED	Aug 16, 2027				
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 203389 001	>A> 10548859	Aug 16, 2036	U-1399			
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 203389 002	>A> 10548859	Aug 16, 2036	U-1399			
<u>CYSTEINE HYDROCHLORIDE - ELCYS</u>						
N 210660 001	>A> 10478453	Jan 15, 2039	DP U-2752			
	>A> 10583155	Jan 15, 2039	DP U-2752			

## PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CYSTEINE HYDROCHLORIDE - NOURESS</u>						
N 212535 001	10493051	Mar 15, 2039	DP			
	10543186	Mar 15, 2039	U-2722			
<u>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</u>						
N 206843 001	8329159	Jul 24, 2029	DS			
<u>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</u>						
N 206843 002	8329159	Jul 24, 2029	DS			
<u>DEFERIPRONE - FERRIPROX</u>						
N 021825 002	7049328	Jun 28, 2021	U-735			
<u>DEOXYCHOLIC ACID - KYBELLA</u>						
N 206333 001	10500214	Mar 02, 2030	DP			
<u>DEXAMETHASONE - HEMADY</u>						
N 211379 001	10537585	Dec 18, 2037	DP			
<u>DIAZEPAM - VALTOCO</u>						
N 211635 001	10265402	May 11, 2025	DP		NP	Jan 10, 2023
	8895546	Mar 27, 2029	DP		ODE-279	Jan 10, 2027
	8927497	Jul 21, 2025	DP U-2727			
	9642913	May 11, 2025	DP			
	9763876	Mar 27, 2029	DP U-2727			
<u>DIAZEPAM - VALTOCO</u>						
N 211635 002	10265402	May 11, 2025	DP		NP	Jan 10, 2023
	8895546	Mar 27, 2029	DP		ODE-279	Jan 10, 2027
	8927497	Jul 21, 2025	DP U-2727			
	9642913	May 11, 2025	DP			
	9763876	Mar 27, 2029	DP U-2727			
<u>DIAZEPAM - VALTOCO</u>						
N 211635 003	10265402	May 11, 2025	DP		NP	Jan 10, 2023
	8895546	Mar 27, 2029	DP		ODE-279	Jan 10, 2027
	8927497	Jul 21, 2025	DP U-2727			
	9642913	May 11, 2025	DP			
	9763876	Mar 27, 2029	DP U-2727			
<u>DOXEPIN HYDROCHLORIDE - SILENOR</u>						
N 022036 001 >A>	10548871	Apr 11, 2028	U-620			
<u>DOXEPIN HYDROCHLORIDE - SILENOR</u>						
N 022036 002 >A>	10548871	Apr 11, 2028	U-620			
<u>EFINACONAZOLE - JUBLIA</u>						
N 203567 001	10342875	Oct 02, 2034	DP U-2720			
	10478601	Apr 25, 2035	DP U-2721			
<u>ELAGOLIX SODIUM - ORILISSA</u>						
N 210450 001 >A>	10537572	Sep 01, 2036	U-2735			
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 001 >A>	10022379	Apr 02, 2029	DP U-2732			
	>A> 10258637	Apr 03, 2034	U-2731			
	>A> 10406172	Jun 15, 2030	DP U-2733			
	>A> 6488962	Jun 20, 2020	DP			
	>A> 7407955	May 02, 2025	DS DP			
	>A> 7579449	Nov 05, 2025	DS			
	>A> 7713938	Apr 15, 2027	DS DP			
	>A> 8119648	Aug 12, 2023	U-1652			
	>A> 8178541	Aug 12, 2023	DP U-1652			
	>A> 8551957	Oct 14, 2029	DP U-2730			

## PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 001	>A> 8673927	May 04, 2027	U-1652			
	>A> 8883805	Nov 26, 2025	DP			
	>A> 9155705	May 21, 2030	DP			
	>A> 9173859	May 04, 2027	DP U-1652			
	>A> 9173859	May 04, 2027	DP U-2730			
	>A> 9415016	Apr 02, 2029	DP			
	>A> 9949998	Jun 11, 2034	U-2731			
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 002	>A> 10022379	Apr 02, 2029	DP U-2732			
	>A> 10258637	Apr 03, 2034	U-2731			
	>A> 10406172	Jun 15, 2030	DP U-2733			
	>A> 6488962	Jun 20, 2020	DP			
	>A> 7407955	May 02, 2025	DS DP			
	>A> 7579449	Nov 05, 2025	DS			
	>A> 7713938	Apr 15, 2027	DS DP			
	>A> 8119648	Aug 12, 2023	U-1652			
	>A> 8178541	Aug 12, 2023	DP U-1652			
	>A> 8551957	Oct 14, 2029	DP U-2730			
	>A> 8673927	May 04, 2027	U-1652			
	>A> 8883805	Nov 26, 2025	DP			
	>A> 9155705	May 21, 2030	DP			
	>A> 9173859	May 04, 2027	DP U-1652			
	>A> 9173859	May 04, 2027	DP U-2730			
	>A> 9415016	Apr 02, 2029	DP			
	>A> 9949998	Jun 11, 2034	U-2731			
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 003	>A> 10022379	Apr 02, 2029	DP U-2732			
	>A> 10258637	Apr 03, 2034	U-2731			
	>A> 10406172	Jun 15, 2030	DP U-2733			
	>A> 6488962	Jun 20, 2020	DP			
	>A> 7407955	May 02, 2025	DS DP			
	>A> 7579449	Nov 05, 2025	DS			
	>A> 7713938	Apr 15, 2027	DS DP			
	>A> 8119648	Aug 12, 2023	U-1652			
	>A> 8178541	Aug 12, 2023	DP U-1652			
	>A> 8551957	Oct 14, 2029	DP U-2730			
	>A> 8673927	May 04, 2027	U-1652			
	>A> 8883805	Nov 26, 2025	DP			
	>A> 9155705	May 21, 2030	DP			
	>A> 9173859	May 04, 2027	DP U-1652			
	>A> 9173859	May 04, 2027	DP U-2730			
	>A> 9415016	Apr 02, 2029	DP			
	>A> 9949998	Jun 11, 2034	U-2731			
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 004	>A> 10022379	Apr 02, 2029	DP U-2732			
	>A> 10258637	Apr 03, 2034	U-2731			
	>A> 10406172	Jun 15, 2030	DP U-2733			
	>A> 6488962	Jun 20, 2020	DP			
	>A> 7407955	May 02, 2025	DS DP			
	>A> 7579449	Nov 05, 2025	DS			
	>A> 7713938	Apr 15, 2027	DS DP			
	>A> 8119648	Aug 12, 2023	U-1652			
	>A> 8178541	Aug 12, 2023	DP U-1652			
	>A> 8551957	Oct 14, 2029	DP U-2730			
	>A> 8673927	May 04, 2027	U-1652			
	>A> 8883805	Nov 26, 2025	DP			
	>A> 9155705	May 21, 2030	DP			
	>A> 9173859	May 04, 2027	DP U-1652			
	>A> 9173859	May 04, 2027	DP U-2730			
	>A> 9415016	Apr 02, 2029	DP			
	>A> 9949998	Jun 11, 2034	U-2731			



## PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - DESCOVY</u>						
N 208215	001				NPP ODE-284 ODE-285	Sep 25, 2020 Sep 28, 2024 Sep 28, 2024
<u>ENZALUTAMIDE - XTANDI</u>						
N 203415	001	8183274	Aug 24, 2026	U-1281		
		8183274	Aug 24, 2026	U-1588		
		8183274	Aug 24, 2026	U-2345		
		8183274	Aug 24, 2026	U-2708		
		9126941	May 15, 2026	U-1588		
		9126941	May 15, 2026	U-2345		
		9126941	May 15, 2026	U-2708		
<u>EPINEPHRINE - ADRENALIN</u>						
N 020800	003	10166334	Jan 21, 2025	DP		
<u>EPINEPHRINE - ADRENALIN</u>						
N 020800	004	10166334	Jan 21, 2025	DP		
<u>ESTRADIOL - IMVEXXY</u>						
N 208564	001	>A> 10537581	Nov 21, 2032	DP U-2316		
		>A> 10537581	Nov 21, 2032	DP U-2317		
<u>ESTRADIOL - IMVEXXY</u>						
N 208564	002	>A> 10537581	Nov 21, 2032	DP U-2316		
		>A> 10537581	Nov 21, 2032	DP U-2317		
<u>ETEPLIRSEN - EXONDYS 51</u>						
N 206488	001	10533174	May 04, 2021	DP		
<u>ETEPLIRSEN - EXONDYS 51</u>						
N 206488	002	10533174	May 04, 2021	DP		
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - TWIRLA</u>						
N 204017	001	>A> 7045145	Mar 14, 2021	DP U-2751	>A> NP	Feb 14, 2023
		>A> 7384650	Mar 14, 2021	DP		
		>A> 8221784	Mar 14, 2021	DP		
		>A> 8221785	Mar 14, 2021	DP		
		>A> 8246978	Aug 26, 2028	DP		
		>A> 8747888	Jul 10, 2028	DP		
		>A> 8883196	Nov 22, 2020	DP U-2751		
		>A> 9050348	Jul 10, 2028	DP		
<u>EXENATIDE SYNTHETIC - BYDUREON</u>						
N 022200	001	>A> 6414126	Oct 04, 2020	DS DP U-2588		
		>A> 6515117	Oct 04, 2020	DS DP U-2588		
		>A> 6872700	Jan 14, 2020	U-2588		
		>A> 6872700	Jan 14, 2020	U-2589		
		>A> 6872700	Jan 14, 2020	U-2590		
		>A> 6872700	Jan 14, 2020	U-2591		
		>A> 6936590	Oct 04, 2020	U-2588		
		>A> 7456254	Jun 30, 2025	DP U-2588		
		>A> 7456254	Jun 30, 2025	DP U-2589		
		>A> 7456254	Jun 30, 2025	DP U-2590		
		>A> 7456254	Jun 30, 2025	DP U-2592		
		>A> 7612176	Apr 13, 2025	DP U-2588		
		>A> 7612176	Apr 13, 2025	DP U-2589		
		>A> 7612176	Apr 13, 2025	DP U-2590		
		>A> 7612176	Apr 13, 2025	DP U-2592		
		>A> 8329648	Aug 18, 2026	U-2588		
		>A> 8329648	Aug 18, 2026	U-2589		
		>A> 8329648	Aug 18, 2026	U-2590		
		>A> 8329648	Aug 18, 2026	U-2593		
		>A> 8329648	Aug 18, 2026	U-2594		
		>A> 8329648	Aug 18, 2026	U-2595		

## PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EXENATIDE SYNTHETIC - BYDUREON</u>						
N 022200 001	>A> 8329648	Aug 18, 2026	U-2596			
	>A> 8361972	Mar 21, 2028	U-2588			
	>A> 8431685	Apr 13, 2025	DP U-2588			
	>A> 8431685	Apr 13, 2025	DP U-2589			
	>A> 8431685	Apr 13, 2025	DP U-2590			
	>A> 8431685	Apr 13, 2025	DP U-2598			
	>A> 8461105	Apr 13, 2025	DP U-2588			
	>A> 8461105	Apr 13, 2025	DP U-2589			
	>A> 8461105	Apr 13, 2025	DP U-2590			
	>A> 8461105	Apr 13, 2025	DP U-2598			
	>A> 8501698	Jun 20, 2027	DP U-2588			
	>A> 8906851	Aug 18, 2026	U-2588			
	>A> 8906851	Aug 18, 2026	U-2589			
	>A> 8906851	Aug 18, 2026	U-2590			
	>A> 8906851	Aug 18, 2026	U-2593			
	>A> 9198925	Oct 04, 2020	U-2588			
	>A> 9238076	Apr 15, 2024	DP U-2588			
	>A> 9238076	Apr 15, 2024	DP U-2589			
	>A> 9238076	Apr 15, 2024	DP U-2590			
	>A> 9238076	Apr 15, 2024	DP U-2599			
	>A> 9884092	Aug 18, 2026	U-2588			
	>A> 9884092	Aug 18, 2026	U-2589			
	>A> 9884092	Aug 18, 2026	U-2590			
	>A> 9884092	Aug 18, 2026	U-2593			
	>A> 9884092	Aug 18, 2026	U-2594			
	>A> 9884092	Aug 18, 2026	U-2595			
	>A> 9884092	Aug 18, 2026	U-2596			
<u>EXENATIDE SYNTHETIC - BYDUREON PEN</u>						
N 022200 002	>A> 6414126	Oct 04, 2020	DS DP U-2588			
	>A> 6515117	Oct 04, 2020	DS DP U-2588			
	>A> 6872700	Jan 14, 2020	U-2588			
	>A> 6872700	Jan 14, 2020	U-2589			
	>A> 6872700	Jan 14, 2020	U-2590			
	>A> 6872700	Jan 14, 2020	U-2591			
	>A> 6936590	Oct 04, 2020	U-2588			
	>A> 7456254	Jun 30, 2025	DP U-2588			
	>A> 7456254	Jun 30, 2025	DP U-2589			
	>A> 7456254	Jun 30, 2025	DP U-2590			
	>A> 7456254	Jun 30, 2025	DP U-2592			
	>A> 7612176	Apr 13, 2025	DP U-2588			
	>A> 7612176	Apr 13, 2025	DP U-2589			
	>A> 7612176	Apr 13, 2025	DP U-2590			
	>A> 7612176	Apr 13, 2025	DP U-2592			
	>A> 8329648	Aug 18, 2026	U-2588			
	>A> 8329648	Aug 18, 2026	U-2589			
	>A> 8329648	Aug 18, 2026	U-2590			
	>A> 8329648	Aug 18, 2026	U-2593			
	>A> 8329648	Aug 18, 2026	U-2594			
	>A> 8329648	Aug 18, 2026	U-2595			
	>A> 8329648	Aug 18, 2026	U-2596			
	>A> 8361972	Mar 21, 2028	U-2588			
	>A> 8431685	Apr 13, 2025	DP U-2588			
	>A> 8431685	Apr 13, 2025	DP U-2589			
	>A> 8431685	Apr 13, 2025	DP U-2590			
	>A> 8431685	Apr 13, 2025	DP U-2598			
	>A> 8461105	Apr 13, 2025	DP U-2588			
	>A> 8461105	Apr 13, 2025	DP U-2589			
	>A> 8461105	Apr 13, 2025	DP U-2590			
	>A> 8461105	Apr 13, 2025	DP U-2598			
	>A> 8501698	Jun 20, 2027	DP U-2588			
	>A> 8906851	Aug 18, 2026	U-2588			
	>A> 8906851	Aug 18, 2026	U-2589			
	>A> 8906851	Aug 18, 2026	U-2590			
	>A> 8906851	Aug 18, 2026	U-2593			
	>A> 9198925	Oct 04, 2020	U-2588			
	>A> 9238076	Apr 15, 2024	DP U-2588			
	>A> 9238076	Apr 15, 2024	DP U-2589			

## PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EXENATIDE SYNTHETIC - BYDUREON PEN</u>						
N 022200 002	>A> 9238076	Apr 15, 2024	DP U-2590			
	>A> 9238076	Apr 15, 2024	DP U-2599			
	>A> 9884092	Aug 18, 2026	U-2588			
	>A> 9884092	Aug 18, 2026	U-2589			
	>A> 9884092	Aug 18, 2026	U-2590			
	>A> 9884092	Aug 18, 2026	U-2593			
	>A> 9884092	Aug 18, 2026	U-2594			
	>A> 9884092	Aug 18, 2026	U-2595			
	>A> 9884092	Aug 18, 2026	U-2596			
<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565 001	10519252	Oct 20, 2023	DS U-2709			
	10519252	Oct 20, 2023	DS U-2710			
	10519252	Oct 20, 2023	DS U-2711			
	10519252	Oct 20, 2023	DS U-2712			
<u>FERRIC CITRATE - AURYXIA</u>						
N 205874 001	5753706	Feb 03, 2021	DP U-1577			
<u>FERRIC DERISOMALTOSE - MONOFERRIC</u>						
N 208171 001	>A> 10414831	Mar 25, 2029	DS DP			
	>A> 8815301	Aug 14, 2029	DS DP U-2734			
<u>FIDAXOMICIN - DIFICID</u>						
N 201699 001	>A> 7906489	Mar 04, 2027	U-2741		NPP	Jan 24, 2023
	>A> 7906489	Mar 04, 2027	U-319		PED	Jul 24, 2023
	>A> 7906489*PED	Sep 04, 2027				
	>A> 8859510	Jul 31, 2027	U-2741			
	>A> 8859510	Jul 31, 2027	U-319			
	>A> 8859510*PED	Jan 31, 2028				
<u>FIDAXOMICIN - DIFICID</u>						
N 213138 001	>A> 7378508	Jul 31, 2027	DS DP		NP	Jan 24, 2023
	>A> 7378508*PED	Jan 31, 2028			PED	Jul 24, 2023
	>A> 7863249	Jul 31, 2027	DP			
	>A> 7863249*PED	Jan 31, 2028				
	>A> 7906489	Mar 04, 2027	U-2741			
	>A> 7906489*PED	Sep 04, 2027				
	>A> 8586551	Jul 23, 2023	DS DP			
	>A> 8586551*PED	Jan 23, 2024				
	>A> 8859510	Jul 31, 2027	U-2741			
	>A> 8859510*PED	Jan 31, 2028				
	>A> 9808530	May 28, 2034	DP			
	>A> 9808530*PED	Nov 28, 2034				
<u>FINGOLIMOD HYDROCHLORIDE - GILENYA</u>						
N 022527 001	10543179	Dec 25, 2027	U-2719			
<u>GLUCAGON - GVOKE PFS</u>						
N 212097 001	>A> 9649364	Apr 22, 2036	DP U-2742			
<u>GLUCAGON - GVOKE PFS</u>						
N 212097 002	>A> 9649364	Apr 22, 2036	DP U-2742			
<u>GLUCAGON - GVOKE HYOPEN</u>						
N 212097 003	>A> 9649364	Apr 22, 2036	DP U-2742			
<u>GLUCAGON - GVOKE HYOPEN</u>						
N 212097 004	>A> 9649364	Apr 22, 2036	DP U-2742			
<u>GLYCOPYRRONIUM TOSYLATE - OBREXZA</u>						
N 210361 001	>A> 10543192	Feb 28, 2033	DP			
	>A> 10548875	Feb 28, 2033	DS DP U-2398			

## PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>GOLODIRSEN - VYONDYS 53</u>						
N 211970 001	10533174	May 04, 2021	DP		ODE-280	Dec 12, 2026
<u>HEXAMINOLEVULINATE HYDROCHLORIDE - CYSVIEW KIT</u>						
N 022555 001	>A> 10556010	Dec 19, 2036	U-2250			
<u>HYDROGEN PEROXIDE - ESKATA</u>						
N 209305 001	10493103	Apr 21, 2035	DP			
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057 001	>A> 10555924	Jun 28, 2033	U-2743			
	>A> 10555925	Jun 28, 2033	U-2744			
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057 002	>A> 10555924	Jun 28, 2033	U-2743			
	>A> 10555925	Jun 28, 2033	U-2744			
<u>LACTITOL - PIZENSY</u>						
N 211281 001					>A> NCE	Feb 12, 2025
<u>LASMIDITAN SUCCINATE - REYVOW</u>						
N 211280 001					NCE	Jan 31, 2025
<u>LASMIDITAN SUCCINATE - REYVOW</u>						
N 211280 002					NCE	Jan 31, 2025
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 001	>A> 10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 002	>A> 10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 003	>A> 10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 004	>A> 10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 005	>A> 10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 006	>A> 10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 007	>A> 10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 008	>A> 10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 009	>A> 10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 010	>A> 10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 011	>A> 10537538	Feb 28, 2037	DP			

## PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 012	>A> 10537538	Feb 28, 2037	DP			
<u>LUMATEPERONE TOSYLATE - CAPLYTA</u>						
N 209500 001	10464938	Mar 12, 2028	DP		NCE	Dec 20, 2024
	7183282	Jun 15, 2020	DS DP			
	8598119	Dec 28, 2029		U-543		
	8648077	Dec 01, 2029	DS DP			
	9199995	Mar 12, 2029		U-2713		
	9586960	Mar 12, 2029	DS DP			
	9616061	May 27, 2029	DP			
	9956227	Dec 03, 2034		U-2714		
	RE39680	Jun 15, 2020	DS DP	U-543		
<u>MELOXICAM - ANJESO</u>						
N 210583 001	>A> 10463673	Feb 24, 2024	DP	U-2750	>A> NP	Feb 20, 2023
	>A> 10471067	Feb 24, 2024	DP	U-2750		
	>A> 8512727	Dec 25, 2022	DP	U-2750		
	>A> 9974746	May 26, 2030	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 001	>A> 10568841	Oct 30, 2035	DP	U-2357		
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 002	>A> 10568841	Oct 30, 2035	DP	U-2357		
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 003	>A> 10568841	Oct 30, 2035	DP	U-2357		
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 004	>A> 10568841	Oct 30, 2035	DP	U-2357		
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 005	>A> 10568841	Oct 30, 2035	DP	U-2357		
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 006	>A> 10568841	Oct 30, 2035	DP	U-2357		
<u>MICAFUNGIN SODIUM - MYCAMINE</u>						
N 021506 002					I-821 PED	Dec 20, 2022 Jun 20, 2023
<u>MICAFUNGIN SODIUM - MYCAMINE</u>						
N 021506 003					I-821 PED	Dec 20, 2022 Jun 20, 2023
<u>MIGALASTAT HYDROCHLORIDE - GALAFOLD</u>						
N 208623 001	10525045	Apr 28, 2028	U-2371			
<u>MINOCYCLINE HYDROCHLORIDE - AMZEEQ</u>						
N 212379 001	10517882	Oct 01, 2030	U-2647			
<u>NERATINIB MALEATE - NERLYNX</u>						
N 208051 001					>A> I-823	Feb 25, 2023
<u>NETUPITANT; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u>						
N 205718 001	>A> 6297375	Mar 17, 2023	DS			
<u>OCTREOTIDE ACETATE - BYNFEZIA PEN</u>						
N 213224 001	10342850	May 15, 2038	DP			

## PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>OLAPARIB - LYNPARZA</u>						
N 208558 001	8143241	Aug 12, 2027	U-2101		ODE-283	Dec 27, 2026
	8143241	Aug 12, 2027	U-2103			
	8143241	Aug 12, 2027	U-2480			
	8143241	Aug 12, 2027	U-2481			
	8143241	Aug 12, 2027	U-2482			
	8143241	Aug 12, 2027	U-2483			
	8143241	Aug 12, 2027	U-2716			
	8859562	Aug 04, 2031	U-2101			
	8859562	Aug 04, 2031	U-2480			
	8859562	Aug 04, 2031	U-2481			
	8859562	Aug 04, 2031	U-2482			
	8859562	Aug 04, 2031	U-2483			
	8859562	Aug 04, 2031	U-2716			
	9566276	Mar 12, 2024	U-2716			
<u>OLAPARIB - LYNPARZA</u>						
N 208558 002	8143241	Aug 12, 2027	U-2101		ODE-283	Dec 27, 2026
	8143241	Aug 12, 2027	U-2103			
	8143241	Aug 12, 2027	U-2480			
	8143241	Aug 12, 2027	U-2481			
	8143241	Aug 12, 2027	U-2482			
	8143241	Aug 12, 2027	U-2483			
	8143241	Aug 12, 2027	U-2716			
	8859562	Aug 04, 2031	U-2101			
	8859562	Aug 04, 2031	U-2480			
	8859562	Aug 04, 2031	U-2481			
	8859562	Aug 04, 2031	U-2482			
	8859562	Aug 04, 2031	U-2483			
	8859562	Aug 04, 2031	U-2716			
	9566276	Mar 12, 2024	U-2716			
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021087 001					M-251	Aug 02, 2022
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021087 002					M-251	Aug 02, 2022
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021087 003					M-251	Aug 02, 2022
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021246 001					M-251	Aug 02, 2022
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021246 002					M-251	Aug 02, 2022
<u>PEMETREXED - PEMFEXY</u>						
N 209472 001	7772209	May 24, 2022	U-2728			
	7772209	May 24, 2022	U-2729			
<u>POMALIDOMIDE - POMALYST</u>						
N 204026 001	>A> 10555939	May 19, 2030	DP			
<u>POMALIDOMIDE - POMALYST</u>						
N 204026 002	>A> 10555939	May 19, 2030	DP			
<u>POMALIDOMIDE - POMALYST</u>						
N 204026 003	>A> 10555939	May 19, 2030	DP			
<u>POMALIDOMIDE - POMALYST</u>						
N 204026 004	>A> 10555939	May 19, 2030	DP			

## PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>REVEFENACIN - YUPELRI</u>						
N 210598 001	10550081	Jul 14, 2030	DS			
<u>RIMEGEPANT SULFATE - NURTEC ODT</u>						
N 212728 001					>A> NCE	Feb 27, 2025
<u>RIVAROXABAN - XARELTO</u>						
N 022406 004					>A> I-824	Oct 11, 2021
<u>SELINEXOR - XPROVIO</u>						
N 212306 001	10519139	Aug 14, 2035	DS DP U-2584			
	>A> 10544108	Jul 26, 2032	U-2584			
<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637 001					I-822	Jan 16, 2023
<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637 002					I-822	Jan 16, 2023
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051 001					M-252	Jan 16, 2023
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051 002					M-252	Jan 16, 2023
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051 003					M-252	Jan 16, 2023
<u>SOLRIAMFETOL HYDROCHLORIDE - SUNOSI</u>						
N 211230 001	10512609	Sep 05, 2037	U-2548			
<u>SOLRIAMFETOL HYDROCHLORIDE - SUNOSI</u>						
N 211230 002	10512609	Sep 05, 2037	U-2548			
<u>SUMATRIPTAN SUCCINATE - ZEMBRACE SYMTOUCH</u>						
N 208223 001	10537554	Jan 29, 2036	U-72			
<u>SUVOREXANT - BELSOMRA</u>						
N 204569 001					M-253	Jan 29, 2023
<u>SUVOREXANT - BELSOMRA</u>						
N 204569 002					M-253	Jan 29, 2023
<u>SUVOREXANT - BELSOMRA</u>						
N 204569 003					M-253	Jan 29, 2023
<u>SUVOREXANT - BELSOMRA</u>						
N 204569 004					M-253	Jan 29, 2023
<u>TAZEMETOSTAT HYDROBROMIDE - TAZVERIK</u>						
N 211723 001	>A> 10155002	Apr 13, 2032	U-2736		NCE	Jan 23, 2025
	>A> 10245269	Apr 11, 2033	U-2737			
	>A> 10369155	Oct 16, 2035	U-2736			
	>A> 10420775	Apr 13, 2032	U-2736			
	>A> 8410088	Apr 13, 2032	DS DP			
	>A> 9090562	Apr 13, 2032	DS DP			
	>A> 9394283	Apr 11, 2033	DS DP			
	>A> 9522152	Apr 13, 2032	U-2738			
	>A> 9549931	Apr 13, 2032	U-2736			
	>A> 9688665	Aug 22, 2034	U-2736			
	>A> 9855275	Apr 13, 2032	U-2736			
	>A> 9872862	Apr 11, 2033	U-2738			
	>A> 9889138	Oct 16, 2035	U-2736			

## PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TAZEMETOSTAT HYDROBROMIDE - TAZVERIK</u>						
N 211723 001	>A> 10155002	Apr 13, 2032	U-2736		NCE	Jan 23, 2025
	>A> 10245269	Apr 11, 2033	U-2737			
	>A> 10369155	Oct 16, 2035	U-2736			
	>A> 10420775	Apr 13, 2032	U-2736			
	>A> 8410088	Apr 13, 2032	DS DP			
	>A> 9090562	Apr 13, 2032	DS DP			
	>A> 9394283	Apr 11, 2033	DS DP			
	>A> 9522152	Apr 13, 2032	U-2738			
	>A> 9549931	Apr 13, 2032	U-2736			
	>A> 9688665	Aug 22, 2034	U-2736			
	>A> 9855275	Apr 13, 2032	U-2736			
	>A> 9872862	Apr 11, 2033	U-2738			
	>A> 9889138	Oct 16, 2035	U-2736			
<u>TEDIZOLID PHOSPHATE - SIVEXTRO</u>						
N 205435 001	>A> 7816379	Apr 20, 2026	DS DP U-2507			
	>A> 7816379	Apr 20, 2026	DS DP U-282			
<u>TEDIZOLID PHOSPHATE - SIVEXTRO</u>						
N 205436 001	>A> 7816379	Apr 20, 2026	DS DP U-2507			
	>A> 7816379	Apr 20, 2026	DS DP U-282			
<u>TENOFOVIR ALAFENAMIDE FUMARATE - VEMLIDY</u>						
N 208464 001					>A> M-255	Feb 04, 2023
<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089 001	>A> 10543219	Apr 12, 2030	U-2506			
<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089 002	>A> 10543219	Apr 12, 2030	U-2506			
<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089 003	>A> 10543219	Apr 12, 2030	U-2506			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 001	8747897	Aug 11, 2031	DP U-2724			
	8747897	Aug 11, 2031	DP U-2725			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 002	8747897	Aug 11, 2031	DP U-2724			
	8747897	Aug 11, 2031	DP U-2725			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 003	8747897	Aug 11, 2031	DP U-2724			
	8747897	Aug 11, 2031	DP U-2725			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 004	8747897	Aug 11, 2031	DP U-2724			
	8747897	Aug 11, 2031	DP U-2725			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 005	8747897	Aug 11, 2031	DP U-2724			
	8747897	Aug 11, 2031	DP U-2725			
<u>TRIPTORELIN PAMOATE - TRELSTAR</u>						
N 022437 001	>A> 10166181	Jun 30, 2029	DP			
<u>TRIPTORELIN PAMOATE - TRIPTODUR KIT</u>						
N 208956 001	10166181	Jun 30, 2029	DP			



## PATENT AND EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 02 - February 2020

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>UBROGEPANT - UBRELVY</u>						
N 211765 001	10117836	Jan 30, 2035	DP			
	8754096	Jul 19, 2032	DS DP U-2717			
	8912210	Nov 10, 2031	DS DP			
	9499545	Nov 10, 2031	DS DP U-2718			
	9833448	Nov 10, 2031	U-2718			
<u>UBROGEPANT - UBRELVY</u>						
N 211765 002	10117836	Jan 30, 2035	DP			
	8754096	Jul 19, 2032	DS DP U-2717			
	8912210	Nov 10, 2031	DS DP			
	9499545	Nov 10, 2031	DS DP U-2718			
	9833448	Nov 10, 2031	U-2718			
<u>UPADACITINIB - RINVOO</u>						
N 211675 001	10519164	Oct 17, 2036	DP			
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573 001	>A> 8722657	Jan 29, 2032	DS			
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573 002	>A> 8722657	Jan 29, 2032	DS			
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573 003	>A> 8722657	Jan 29, 2032	DS			
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N 022567 001	7834020	Jun 05, 2022	DS DP U-839		>A> M-254	Jan 31, 2023
	7834020*PED	Dec 05, 2022			>A> PED	Jul 31, 2023
	8193195	Jun 05, 2022	U-839			
	8193195*PED	Dec 05, 2022				
	8236804	Jun 05, 2022	U-839			
	8236804*PED	Dec 05, 2022				
	8673921	Jun 05, 2022	DS DP			
	8673921*PED	Dec 05, 2022				
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N 022567 002	7834020	Jun 05, 2022	DS DP U-839		>A> M-254	Jan 31, 2023
	7834020*PED	Dec 05, 2022			>A> PED	Jul 31, 2023
	8193195	Jun 05, 2022	U-839			
	8193195*PED	Dec 05, 2022				
	8236804	Jun 05, 2022	U-839			
	8236804*PED	Dec 05, 2022				
	8673921	Jun 05, 2022	DS DP			
	8673921*PED	Dec 05, 2022				
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N 022567 003	7834020	Jun 05, 2022	DS DP U-839		>A> M-254	Jan 31, 2023
	7834020*PED	Dec 05, 2022			>A> PED	Jul 31, 2023
	8193195	Jun 05, 2022	U-839			
	8193195*PED	Dec 05, 2022				
	8236804	Jun 05, 2022	U-839			
	8236804*PED	Dec 05, 2022				
	8673921	Jun 05, 2022	DS DP			
	8673921*PED	Dec 05, 2022				
<u>VOXELOTOR - OXBRYTA</u>						
N 213137 001	10017491	Dec 28, 2032	DP		ODE-281	Nov 25, 2026
	10034879	Dec 28, 2032	DS DP			
	10493035	Oct 12, 2037	DP			
	9018210	Dec 28, 2032	DS DP			
	9248199	Jan 29, 2034	U-2676			
	9248199	Jan 29, 2034	U-2715			
	9447071	Feb 06, 2035	DS DP			

## 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, 40<sup>th</sup> 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/results\\_patent.cfm](http://www.accessdata.fda.gov/scripts/cder/ob/results_patent.cfm)

The current complete list of exclusivity terms is available at [http://www.accessdata.fda.gov/scripts/cder/ob/results\\_exclusivity.cfm](http://www.accessdata.fda.gov/scripts/cder/ob/results_exclusivity.cfm)