

**CUMULATIVE
SUPPLEMENT 5
JAN'88-MAY'88**

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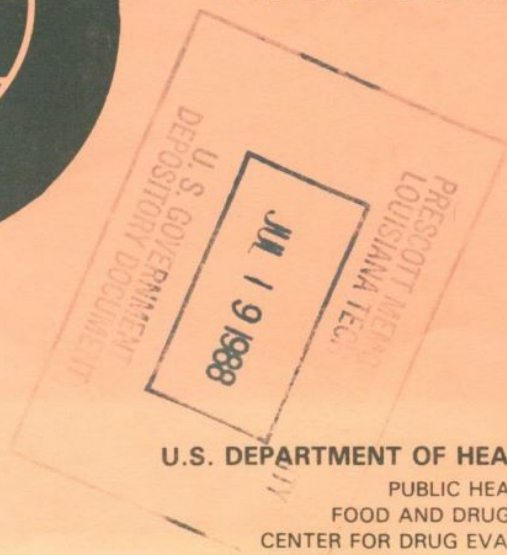
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APPROVED DRUG PRODUCTS

**WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS**

8TH EDITION



U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
PUBLIC HEALTH SERVICE
FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH
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Prepared By
Division of Drug Information Resources
Office of Management
Center for Drug Evaluation and Research, FDA

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8TH EDITION

CUMULATIVE SUPPLEMENT 5

MAY 1988

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8th EDITION
CUMULATIVE SUPPLEMENT 5
MAY 1988

1.0 INTRODUCTION

1.1 HOW TO USE THE CUMULATIVE SUPPLEMENT

This Cumulative Supplement is one of a series of monthly updates to the Approved Drug Products with Therapeutic Equivalence Evaluations, 8th Edition (the List). The List is composed of four parts: approved prescription drug products with therapeutic equivalence evaluations, over-the-counter (OTC) drug products that require approved applications as a condition of marketing, drug products in the Division of Blood and Blood Products approved under Section 505 of the Act, and products discontinued from marketing or products which have had their approval withdrawn for other than safety or efficacy reasons.

The Cumulative Supplement provides, among other things, information on newly approved drugs and, if necessary, revised therapeutic equivalence evaluations and updated patent and exclusivity data. The Addendum contains appropriate drug patent and exclusivity information required of the Agency by the "Drug Price Competition and Patent Term Restoration Act of 1984" for the Prescription, OTC, and Drug Products in the Division of Blood and Blood Products Approved Under Section 505 of the Act lists.

The Patent and Exclusivity Lists are arranged in alphabetical order by active ingredient name. For those products with multiple active ingredients, only the first active ingredient (in alphabetical sort) will appear. In addition, the trade name will be displayed to the right of the active ingredient name for each product, along with the application number and product number (FDA's internal file number). All patents with their expiration dates are displayed for each application number. 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 in the Patent and Exclusivity Information Addendum for an explanation of all codes and abbreviations.

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 place an asterisk (*) 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. [Strength(s) which already exist in the List will not be repeated for context.] The effective date for the approved drug product (the earliest date a product may be marketed) appears, when appropriate, to the left of the approval date.

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.)

Additions new to the Prescription Drug Product List, OTC Drug Product List, List of Drug Products in the Division of Blood and Blood Products Approved Under Section 505 of the Act and the Patent and Exclusivity Data are indicated by the symbol >ADD> to the left of the line on which new information exists. The >ADD> symbol is then dropped in subsequent Cumulative Supplements for that item. A newly approved product is also identified by a lozenge (■) to the right of its strength which remains throughout all Cumulative Supplements for this edition.

Deletions new to the Prescription Drug Product List, OTC Drug Product List, List of Drug Products in the Division of Blood and Blood Products Approved Under Section 505 of the Act and the Patent and Exclusivity Data are indicated by the symbol >DLT> (DELETE) to the left of the line containing overstruck print. The >DLT> symbol is dropped in subsequent Cumulative Supplements for that item. The overstruck print will remain in the Prescription Drug Product List and OTC Drug Product Lists in all Cumulative Supplements for this edition. However, the overstruck print in the List of Drug Products in the Division of Blood and Blood Products Approved Under Section 505 of the Act and the Patent and Exclusivity Data will be dropped in subsequent Cumulative Supplements.

Products discontinued from marketing or products which have had their approval withdrawn for other than safety or effectiveness reasons, will be flagged in this Cumulative Supplement with the "⦿" symbol to designate their non-marketed status. All products having a "⦿" symbol in the 12th Cumulative Supplement of the 8th Edition List will then be added to the "Discontinued Drug Product List" appearing in the 9th Edition.

1.2 PRODUCTS REQUIRING REVISED LABELING FOR FULL APPROVAL

Drug products in this category (1) initially received approval only on the basis of safety before effectiveness studies were required, or (2) were conditionally approved under the temporary exemption that allowed these products to be marketed while effectiveness studies were being conducted. Listed below are those drugs which are now required to revise their labeling and provide additional information necessary for full approval on the basis of requirements listed in the Federal Register. As approval is granted by the Agency for a specific product, based on additional information submitted by the applicant, the product will be included in the appropriate Drug Product List.

<u>Products</u>	<u>Federal Register Reference</u>
Nitroglycerin (capsule, controlled release;oral)	SEP 7, 1984 (49 FR 35428)
Nitroglycerin (ointment;topical)	SEP 3, 1986 (51 FR 31371)
Nitroglycerin (tablet, controlled release;oral)	SEP 7, 1984 (49 FR 35428)
Nitroglycerin (tablet, controlled release;buccal)	JUL 5, 1985 (50 FR 27688)
Tranlylcypromine Sulfate	MAR 22, 1984 (49 FR 10708)

1.3 APPLICANT (NAME) CHANGES

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

1.4 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 sections 505 and 507 of the Federal Food, Drug, and Cosmetic Act. Excluded are approved drug products marketed by distributors; those marketed solely abroad; and those now regarded as medical devices, biologics or foods.

The baseline column (Dec 1987) 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.

COUNTS CUMULATIVE BY QUARTER¹

CATEGORIES COUNTED	DEC 1987	MAR 1988	JUN 1988	SEP 1988
DRUG PRODUCTS LISTED	9709	9528*		
SINGLE SOURCE	2096 (21.6%)	1997 (21.0%)		
MULTISOURCE	7613 (78.4%)	7531 (79.0%)		
THERAPEUTICALLY EQUIVALENT	6691 (68.9%)	6660 (69.9%)		
NOT THERAPEUTICALLY EQUIVALENT	848 (8.7%)	770 (8.1%)		
EXCEPTIONS ²	74 (0.8%)	101 (1.0)		
NEW MOLECULAR ENTITIES APPROVED	--	1		
NUMBER OF APPLICANTS	349	361		

*This number is inclusive of products discontinued since December 1987, and any products approved or discontinued through March 1988.

(1) Cumulative counts are calculated from January 1, 1988 to, and including, the month indicated.

(2) Amino acid-containing products of varying composition (see Introduction, page 1-8 of the List).

PRESCRIPTION DRUG PRODUCT LIST

8TH EDITION

CUMULATIVE SUPPLEMENT NUMBER 5 / JAN'88 - MAY'88

1

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE
MIKART

> ADD >
> ADD >

N89451 001
MAY 23, 1988

TABLET; ORAL

TYLENOL W/ CODEINE NO. 4
/AA/ /MCNEIL/LABS/ /300MG; 60MG/
AA MCNEIL PHARM 300MG; 60MG

/N85055/004/
N85055 004

BUTALBITAL, APAP, AND CAFFEINE

AB HALSEY DRUG 325MG; 50MG; 40MG

N89536 001
FEB 16, 1988

ACETAMINOPHEN; CODEINE PHOSPHATE

ELIXIR; ORAL

MYAPAP AND CODEINE

MY K LABS

/MYAPAP W/ CODEINE/
/MYK/LABS/

> ADD >
> ADD >
> DLT >
> DLT >

120MG/5ML; 12MG/5ML

/140MG/5ML; 12MG/5ML/
/140MG/5ML; 12MG/5ML/
/140MG/5ML; 12MG/5ML/
120MG/5ML; 12MG/5ML

N87006 001

/N87006/001/
/N85057/001/
N85057 001

N89360 001
MAR 02, 1988

N89696 001
APR 21, 1988

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

/BOOTS/LABS/

300MG; 30MG

> DLT >
> DLT >

AA MUTUAL PHARM

300MG; 15MG

N89671 001

/DEC/10/1982/
FEB 10, 1988

AA

300MG; 30MG

N89672 001

FEB 10, 1988

AA

300MG; 60MG

N89673 001

FEB 10, 1988

AA PHARMAFAIR

300MG; 30MG

N87762 001

DEC 10, 1982

TYLENOL W/ CODEINE

/MCNEIL/LABS/

325MG; 7.5MG

325MG; 15MG

/N85056/001/
/N85056/002/
/N85056/003/
/N85056/004/
N85056 001
N85056 002
N85056 003
N85056 004

/N85056/001/
/N85056/002/
/N85056/003/
/N85056/004/
N85056 001
N85056 002
N85056 003
N85056 004

N89619 001
JAN 13, 1988

N09388 001

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

QUAD PHARMS

500MG/VIAL

N89619 001
JAN 13, 1988

N09388 001

DIAMOX

AP LEDERLE LABS

500MG/VIAL

N09388 001

ALBUTEROL SULFATE

CAPSULE; INHALATION

VENTOLIN ROTACAPS

GLAXO

EQ 0.2MG BASE

N19489 001
MAY 04, 1988

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIL

AMITRIL
/PARKE/DAVIS/
/AB/

1999/51269N/NO9215 001

100MG/	100MG/
150MG/	150MG/
AB/	AB/
AB/	AB/

WARNER CHILCOTT

<u>AB</u>	<u>10MG</u>
<u>AB</u>	<u>25MG</u>
<u>AB</u>	<u>50MG</u>

	<u>AB</u>	<u>AB</u>	<u>75MG</u>	<u>50MG</u>	<u>25MG</u>
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2					
3					
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100					

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TRIPTYLINE H

10

/N18792/001
 JAN/17, 1984/
 N18792 001

AB

N72277 001
MAY 09, 1988
N72278 001
MAY 09, 1988
N70477 001
JAN 12, 1988
N70478 001
JAN 12, 1988

10

ICILLIN

1986297/061/
AUG/19./1983/
1986298/061/

AD
CLONTEL CHERB

AB 500MGH

POWDER FOR RECONSTITUTION; ORAL

AB
250MG/5ML A

1700/0740N
N07320 002

NO7320 002 / NO7336/6621

N62885 001
MAR 08, 1988
N62885 002
MAR 08, 1988

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AB AMPICILLIN
CLONWEL CHEMS

EQ 250MG BASEM

N62883 001
FEB 25, 1988

EQ 500MG BASEM

N62882 001
FEB 25, 1988

POLYICILLIN

AB BRSTL MYRS IND

EQ 250MG BASEM

N62888 001
MAR 04, 1988

EQ 500MG BASEM

N62888 002
MAR 04, 1988

ASPIRIN; CAFFEINE; PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL
COMPOUND 65

/AA/ /BANMAX/PHARMS/

/389MG;32.4MG;65MG/

389MG;32.4MG;65MG

/N64553/001/
/AUG 17, 1983/
N84553 002
AUG 17, 1983

ASPIRIN; HYDROCODONE BITARTRATE

TABLET; ORAL
AZDONE

CENTRAL PHARMS

500MG;5MG

N89420 001
JAN 25, 1988

BACLOFEN

TABLET; ORAL
BACLOFEN

PHARM BASICS

10MG

N71260 001
MAY 06, 1988

20MG

N71261 001
MAY 06, 1988

10MG

N71901 001
APR 13, 1988

20MG

N71902 001
APR 13, 1988

LIQRESAL

GEIGY PHARMS

10MG

N17851 001

LIQRESAL DS

GEIGY PHARMS

20MG

N17851 003
JAN 20, 1982

BENZTHIAZIDE

TABLET; ORAL

BENZTHIAZIDE

/BP/ /PRIVATE/FMLTNS/
a PRIVATE FMLTNS

/50MG/
50MG

/N63206/001/
N83206 001

BENZTROPINE MESYLATE

TABLET; ORAL

BENZTROPINE MESYLATE

PAR PHARM

0.5MG

N88877 001
APR 11, 1985

1MG

N88894 001
APR 11, 1985

2MG

N88895 001
APR 11, 1985

/0.5MG/

/1MG/

/2MG/

/0.5MG/

/1MG/

/2MG/

QUANTUM PHARMS

0.5MG

N88514 001
JAN 31, 1984

1MG

N88510 001
JAN 31, 1984

2MG

N88511 001
JAN 31, 1984

/0.5MG/

/1MG/

/2MG/

/0.5MG/

/1MG/

/2MG/

COGENTIN

MS&D

> ADD > AA

> ADD > AA

> ADD > AA

> DLT > /BP/

> DLT > /BP/

> DLT > /BP/

> DLT > /BP/

> DLT > /BP/

0.5MG

1MG

2MG

/0.5MG/

/1MG/

/2MG/

N09193 004
N09193 003

N09193 002
/N09193/004/

/N09193/003/

/N09193/002/

EQ 0.1% BASEM

N72041 001
JAN 06, 1988

BETAMETHASONE VALERATE

LOTION; TOPICAL

BETAMETHASONE VALERATE

AB COPLEY PHARM

EQ 0.1% BASE

N71883 001
APR 22, 1988BETHANECHOL CHLORIDE

INJECTABLE; INJECTION

BETHANECHOL CHLORIDE

AP QUAD PHARMS

5MG/ML

N89815 001
APR 12, 1988URECHOLINE

AP MSD

5MG/ML

N06536 001

BRETYLIUM IOSYLATE

INJECTABLE; INJECTION

BRETYLIUM IOSYLATE

AP QUAD PHARMS

50MG/ML

N71181 001
FEB 16, 1988BROMPHENIRAMINE MALEATE

TABLET; ORAL

BROMPHENIRAMINE MALEATE/BA/ /BARR/LABS/
3 BARR LABS4MG/
4MG/N84468/001/
N84468 001CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER

KENDALL MCGAM

10MG/100ML; 2.5GM/100ML; 15MG/100ML;
300MG/100ML;
160MG/100ML

N19634 001

DEXTROSE 4% IN MODIFIED LACTATED RINGER'S IN PLASTIC CONTAINER

KENDALL MCGAM

4MG/100ML; 4GM/100ML; 6MG/100ML;
120MG/100ML; 62MG/100MLN19634 002
FEB 24, 1988CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER

AP KENDALL MCGAM

20MG/100ML; 5GM/100ML; 30MG/100ML;
600MG/100ML;
310MG/100MLN19634 003
FEB 24, 1988CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

LACTATED RINGER'S IN PLASTIC CONTAINER

AP KENDALL MCGAM

20MG/100ML; 30MG/100ML; 600MG/100ML;
310MG/100MLN19632 001
FEB 29, 1988CARBAMAZEPINE

TABLET, CHEMABLE; ORAL

CARBAMAZEPINE

AB WARNER CHILCOTT

100MG

N71940 001
FEB 01, 1988TEGRETOL

AB GEIGY PHARMS

100MG

N18281 001

CEFACTOR

POWDER FOR RECONSTITUTION; ORAL

CECLOR

LILLY

EQ 187MG BASE/5ML

N62206 003

EQ 375MG BASE/5ML

APR 20, 1988
N62206 004
APR 20, 1988CEFAZOLIN SODIUM

INJECTABLE; INJECTION

ANCEF

AP SK&F LABS

EQ 5GM BASE/VIAL

N50461 004

CEFAZOLIN SODIUMINJECTABLE; INJECTION
CEFAZOLIN SODIUM
ELKINS SINN

AP	EQ 250MG BASE/VIALM	N62807 001	AB	N62850 001
		JAN 12, 1988		APR 22, 1988
AP	EQ 500MG BASE/VIALM	N62807 002	AB	N62851 001
		JAN 12, 1988		APR 22, 1988
AP	EQ 1GM BASE/VIALM	N62807 003	AB	N62813 001
		JAN 12, 1988		FEB 25, 1988
AP	EQ 5GM BASE/VIALM	N62807 004	AB	N62813 002
		JAN 12, 1988		FEB 25, 1988
AP	EQ 10GM BASE/VIALM	N62807 005		
		JAN 12, 1988		
AP	EQ 20GM BASE/VIALM	N62807 006		
		JAN 12, 1988		

CEFOTETAN DISODIUMINJECTABLE; INJECTION
CEFOTAN
STUART PHARMS

EQ 10GM BASE/VIALM	N50588 003
	APR 25, 1988

CEPHALEXINCAPSULE; ORAL
CEPHALEXIN
JEROME STEVENS

AB	EQ 250MG BASEM	N62870 001		
		MAR 17, 1988		
AB	EQ 500MG BASEM	N62869 001		
		MAR 17, 1988		
AB	EQ 250MG BASEM	N62821 001		
		FEB 05, 1988		
AB	EQ 500MG BASEM	N62823 001		
		FEB 05, 1988		

POWDER FOR RECONSTITUTION; ORAL

CEPHALEXIN
TAG PHARMS

> ADD >	AB	EQ 125MG BASE/5MLM	N62873 001	
> ADD >			MAY 23, 1988	
> ADD >	AB	EQ 250MG BASE/5MLM	N62867 001	
			APR 15, 1988	

CEPHALEXIN HYDROCHLORIDETABLET; ORAL
KEFTAB
LILLY

> ADD >		EQ 333MG BASEM	N50614 003	
> ADD >			MAY 16, 1988	

CEPHRADINECAPSULE; ORAL
CEPHRADINE
BARR LABS

AB	250MGM	N62850 001	
		APR 22, 1988	
AB	500MGM	N62851 001	
		APR 22, 1988	
AB	250MGM	N62813 001	
		FEB 25, 1988	
AB	500MGM	N62813 002	
		FEB 25, 1988	

POWDER FOR RECONSTITUTION; ORAL

> ADD >	AB	125MG/5MLM	N62858 001	
> ADD >			MAY 19, 1988	
> ADD >	AB	250MG/5MLM	N62859 001	
> ADD >			MAY 19, 1988	

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HCL
/PUREPAC/PHARM/

/AB/	/5MG/	/N85155/001/
/AB/	/10MG/	/N84939/002/
/AB/	/25MG/	/N85144/001/
		N85155 001
		N84939 002
		N85144 001

PUREPAC PHARM

LYGEN

/BANMAX/PHARMS/

/AB/	/5MG/	/N85167/002/
/AB/	/10MG/	/N85009/001/
/AB/	/25MG/	/N85168/001/
		N85107 002
		N85009 001
		N85108 001

BANMAX PHARMS

CHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CHLOROTHIAZIDE W/ RESERPINE

> DLT >	/BP/	/25MG; 0.125MG/
> ADD >		/BOLAR PHARM

/N84853/001/
N84853 001

CHLORPHENIRAMINE MALEATE

TABLET; ORAL

CHLORPHENIRAMINE MALEATE
/AA/ /BARR/LABS/ 4MG/4MG
BARR LABS

CLINDAMYCIN PHOSPHATE

INJECTABLE; INJECTION

CLINDAMYCIN PHOSPHATE
> ADD > AP
> ADD >
/N80700/001/
N80700 001

N62905 001
MAY 09, 1988

EQ 150MG BASE/MLM

CHLORPROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHLORPROMAZINE HCL
AP MARSAM PHARMS 25MG/MLM

N62953 001
APR 21, 1988

EQ 150MG BASE/MLM

N62889 001
APR 25, 1988

EQ 150MG BASE/MLM

N62877 001
MAR 15, 1988

EQ 150MG BASE/MLM

N62819 001
MAR 15, 1988

EQ 150MG BASE/MLM

N62852 001
MAR 17, 1988

EQ 150MG BASE/MLM

CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE
BARR LABS 500MG
> ADD > AA
> ADD >
CORD LABS 250MG
> ADD > AA
> ADD >
LEMMON 500MG
> ADD > AA
> ADD >
PARAFON FORTE DSC
MCNEIL PHARM 500MG
> ADD > AA
> ADD >

N89895 001
MAY 04, 1988
N89852 001
MAY 04, 1988
N89853 001
MAY 04, 1988
N89859 001
MAY 04, 1988

CLOFIBRATE

CAPSULE; ORAL

CLOFIBRATE
> ADD > AB
> ADD >
CORD LABS 500MG

N72191 001
MAY 02, 1988

500MG

CLONIDINE HYDROCHLORIDE

TABLET; ORAL

CLONIDINE HCL
AB LEDERLE LABS

0.1MG

N71783 001
APR 05, 1988

0.2MG

N71784 001
APR 05, 1988

0.3MG

N71785 001
APR 05, 1988

0.3MG

N71785 001
APR 05, 1988

CLORAZEPATE DIPOTASSIUM

CAPSULE; ORAL

CLORAZEPATE DIPOTASSIUM
AB CHELSEA LABS 3.75MG

3.75MG

N71878 001
MAR 15, 1988

7.5MG

N71879 001
MAR 15, 1988

15MG

N71860 001
MAR 15, 1988

15MG

N71860 001
MAR 15, 1988

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

CLEOCIN HCL
UPJOHN EQ 300MG BASEM

N50162 003
APR 14, 1988

EQ 300MG BASEM

CLORAZEPATE DIPOTASSIUMCAPSULE; ORALCLORAZEPATE DIPOTASSIUMPUREPAC PHARM 3.75MGAB AB N71924 001
APR 25, 1988AB AB N71925 001
APR 25, 1988AB AB N71926 001
APR 25, 1988AB AB N71774 001
MAR 01, 1988AB AB N71775 001
MAR 01, 1988AB AB N71776 001
MAR 01, 1988TABLET; ORALCLORAZEPATE DIPOTASSIUMWARNER CHILCOTT 3.75MGAB AB N71828 001
MAR 03, 1988AB AB N71829 001
MAR 03, 1988AB AB N71830 001
MAR 03, 1988AB AB N71852 001
FEB 09, 1988AB AB N71853 001
FEB 09, 1988AB AB N71854 001
FEB 09, 1988GEN-XENE

ALRA LABS

AB AB N71787 001
APR 26, 1988AB AB N71788 001
APR 26, 1988AB AB N71789 001
APR 26, 1988CLOXACILLIN SODIUMPOWDER FOR RECONSTITUTION; ORALCLOXACILLIN SODIUMAA AB EQ 125MG BASE/5ML
/EQ 125MG BASE/5ML/AA AB BIOCRRAFT LABS
N62268 001
/N62268/001/AA AB EQ 125MG BASE/5ML
/EQ 125MG BASE/5ML/AA AB TEGOPEN
N50192 001AA AB BRISTOL LABS
N61453 001AA AB EQ 125MG BASE/5ML
/EQ 125MG BASE/5ML/AA AB EQ 125MG BASE/5ML
/EQ 125MG BASE/5ML/COLCHICINE; PROBENECIDTABLET; ORALPROBENECID AND COLCHICINE

/BP/ /BEECHAM LABS/ /0.5MG;500MG/

/N84321/001/

/BP/ /BEECHAM LABS/ /0.5MG;500MG/

/BP/ /LEDERLE LABS/ /0.5MG;500MG/

/BP/ /LEDERLE LABS/ /0.5MG;500MG/

CYCLOBENZAPRINE HYDROCHLORIDETABLET; ORALCYCLOBENZAPRINE HCLAB AB DANBURY PHARMA 10MG

MAY 03, 1989 : FEB 29, 1988

FLEXERILAB AB MS&D 10MG

N17821 002

DACARBAZINEINJECTABLE; INJECTIONDACARBAZINE

QUAD PHARMS

500MG/VIAL

N71563 001
MAY 06, 1988DESIPRAMINE HYDROCHLORIDETABLET; ORALDESIPRAMINE HCL

CORD LABS

AB AB 10MGN72099 001
MAY 24, 1988AB AB 25MGN72100 001
MAY 24, 1988AB AB 50MGN72101 001
MAY 24, 1988AB AB 10MGN72167 001
FEB 03, 1988AB AB 150MGN72254 001
FEB 03, 1988MORPRAMIN

MERRELL DOW

AB AB 10MGN14399 007
FEB 11, 1982AB AB 150MG

N14399 006

DEXAMETHASONE

TABLET; ORAL

DEXAMETHASONE

/BP/ 0.25MG /BARR LABS/ /N84013/001/
 /BP/ 0.25MG /BARR LABS/ /N84764/001/
 /BP/ 0.5MG /BARR LABS/ /N84084/001/
 /BP/ 0.5MG /BARR LABS/ /N84081/001/
 /BP/ 0.75MG /BARR LABS/ /N84765/001/
 /BP/ 0.75MG /BARR LABS/ /N84086/001/
 /BP/ 1.5MG /BARR LABS/ /N84763/001/
 3 BARR LABS
 3 0.25MG
 3 0.5MG
 3 0.75MG
 3 0.75MG
 3 1.5MG
 3 1.5MG

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 10% IN PLASTIC CONTAINER

AP KENDALL MCGAM 10GM/100MLM /N19626 004/
 FEB 02, 1988
 DEXTROSE 2.5% IN PLASTIC CONTAINER
 KENDALL MCGAM 2.5GM/100MLM /N19626 001/
 FEB 02, 1988

DEXTROSE 5% IN PLASTIC CONTAINER

AP KENDALL MCGAM 5GM/100MLM /N19626 002/
 FEB 02, 1988
 DEXTROSE 7.7% IN PLASTIC CONTAINER
 KENDALL MCGAM 7.7GM/100MLM /N19626 003/
 FEB 02, 1988

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% AND SODIUM CHLORIDE 0.3% W/ POTASSIUM CHLORIDE 0.07% IN PLASTIC CONTAINER

/AP/ 5GM/100ML; 5MG/100ML; 110MG/100MLM /N19630 001/
 FEB 17, 1988
 DEXTROSE 5% AND SODIUM CHLORIDE 0.3% W/ POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER
 /AP/ 5GM/100ML; 5MG/100ML; 330MG/100MLM /N19630 013/
 FEB 17, 1988

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

/DEX/ 5% AND SODIUM CHLORIDE 0.3% W/ POTASSIUM CHLORIDE 0.07% IN PLASTIC CONTAINER /N19630 001/
 FEB 17, 1988
 DEXTROSE 5% AND SODIUM CHLORIDE 0.3% W/ POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER /N19630 013/
 FEB 17, 1988

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER
 KENDALL MCGAM 10GM/100ML; 37MG/100ML; 200MG/100MLM /N19630 031/
 FEB 17, 1988
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER
 KENDALL MCGAM 10GM/100ML; 37MG/100ML; 450MG/100MLM /N19630 037/
 FEB 17, 1988
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
 KENDALL MCGAM 10GM/100ML; 37MG/100ML; 900MG/100MLM /N19630 043/
 FEB 17, 1988
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER
 KENDALL MCGAM 5GM/100ML; 37MG/100ML; 110MG/100MLM /N19630 001/
 FEB 17, 1988
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER
 KENDALL MCGAM 5GM/100ML; 37MG/100ML; 200MG/100MLM /N19630 007/
 FEB 17, 1988
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER
 KENDALL MCGAM 5GM/100ML; 37MG/100ML; 330MG/100MLM /N19630 013/
 FEB 17, 1988

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER
KENDALL MCGAM
5GM/100ML; 37MG/100ML;
450MG/100MLM
N19630 019
FEB 17, 1988

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
KENDALL MCGAM
5GM/100ML; 37MG/100ML;
900MG/100MLM
N19630 025
FEB 17, 1988

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER
KENDALL MCGAM
10GM/100ML; 75MG/100ML;
200MG/100MLM
N19630 032
FEB 17, 1988

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER
KENDALL MCGAM
10GM/100ML; 75MG/100ML;
450MG/100MLM
N19630 038
FEB 17, 1988

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
KENDALL MCGAM
10GM/100ML; 75MG/100ML;
900MG/100MLM
N19630 044
FEB 17, 1988

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER
KENDALL MCGAM
5GM/100ML; 75MG/100ML;
200MG/100MLM
N19630 008
FEB 17, 1988

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER
KENDALL MCGAM
5GM/100ML; 75MG/100ML;
330MG/100MLM
N19630 014
FEB 17, 1988

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER
KENDALL MCGAM
5GM/100ML; 75MG/100ML;
450MG/100MLM
N19630 020
FEB 17, 1988

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
KENDALL MCGAM
5GM/100ML; 75MG/100ML;
900MG/100MLM
N19630 026
FEB 17, 1988

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER
KENDALL MCGAM
5GM/100ML; 75MG/100ML;
110MG/100MLM
N19630 002
FEB 17, 1988

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER
KENDALL MCGAM
10GM/100ML; 110MG/100ML;
200MG/100MLM
N19630 033
FEB 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER
KENDALL MCGAM
10GM/100ML; 110MG/100ML;
450MG/100MLM
N19630 039
FEB 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
KENDALL MCGAM
10GM/100ML; 110MG/100ML;
900MG/100MLM
N19630 045
FEB 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER
KENDALL MCGAM
5GM/100ML; 110MG/100ML;
110MG/100MLM
N19630 003
FEB 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER
KENDALL MCGAM
5GM/100ML; 110MG/100ML;
200MG/100MLM
N19630 009
FEB 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER
KENDALL MCGAM
5GM/100ML; 110MG/100ML;
330MG/100MLM
N19630 015
FEB 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER
KENDALL MCGAM
5GM/100ML; 110MG/100ML;
450MG/100MLM
N19630 021
FEB 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
KENDALL MCGAM
5GM/100ML; 110MG/100ML;
900MG/100MLM
N19630 027
FEB 17, 1988

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER
KENDALL MCGAM
10GM/100ML; 150MG/100ML;
200MG/100MLM
N19630 034
FEB 17, 1988

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER
KENDALL MCGAW

10GM/100ML;150MG/100ML;
450MG/100ML

N19630 040
FEB 17, 1988

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
KENDALL MCGAW

10GM/100ML;150MG/100ML;
900MG/100ML

N19630 046
FEB 17, 1988

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM

CHLORIDE 0.2% IN PLASTIC CONTAINER

KENDALL MCGAW
5GM/100ML;150MG/100ML;
200MG/100ML

N19630 010
FEB 17, 1988

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM

CHLORIDE 0.33% IN PLASTIC CONTAINER

KENDALL MCGAW
5GM/100ML;150MG/100ML;
330MG/100ML

N19630 016
FEB 17, 1988

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM

CHLORIDE 0.45% IN PLASTIC CONTAINER

KENDALL MCGAW
5GM/100ML;150MG/100ML;
450MG/100ML

N19630 022
FEB 17, 1988

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM

CHLORIDE 0.9% IN PLASTIC CONTAINER

KENDALL MCGAW
5GM/100ML;150MG/100ML;
900MG/100ML

N19630 028
FEB 17, 1988

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER
KENDALL MCGAW

5GM/100ML;150MG/100ML;
110MG/100ML

N19630 004
FEB 17, 1988

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER
KENDALL MCGAW

10GM/100ML;220MG/100ML;
200MG/100ML

N19630 035
FEB 17, 1988

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER
KENDALL MCGAW

10GM/100ML;220MG/100ML;
450MG/100ML

N19630 041
FEB 17, 1988

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
KENDALL MCGAW

10GM/100ML;220MG/100ML;
900MG/100ML

N19630 047
FEB 17, 1988

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER
KENDALL MCGAW

5GM/100ML;220MG/100ML;
110MG/100ML

N19630 005
FEB 17, 1988

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER
KENDALL MCGAW

5GM/100ML;220MG/100ML;
200MG/100ML

N19630 011
FEB 17, 1988

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER
KENDALL MCGAW

5GM/100ML;220MG/100ML;
330MG/100ML

N19630 017
FEB 17, 1988

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER
KENDALL MCGAW

5GM/100ML;220MG/100ML;
450MG/100ML

N19630 023
FEB 17, 1988

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
KENDALL MCGAW

5GM/100ML;220MG/100ML;
900MG/100ML

N19630 029
FEB 17, 1988

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER
KENDALL MCGAW

10GM/100ML;300MG/100ML;
200MG/100ML

N19630 036
FEB 17, 1988

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER
KENDALL MCGAW

10GM/100ML;300MG/100ML;
450MG/100ML

N19630 042
FEB 17, 1988

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
KENDALL MCGAW

10GM/100ML;300MG/100ML;
900MG/100ML

N19630 048
FEB 17, 1988

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE

0.2% IN PLASTIC CONTAINER

KENDALL MCGAW
5GM/100ML;300MG/100ML;
200MG/100ML

N19630 012
FEB 17, 1988

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE

0.33% IN PLASTIC CONTAINER

KENDALL MCGAW
5GM/100ML;300MG/100ML;
330MG/100ML

N19630 018
FEB 17, 1988

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE

0.9% IN PLASTIC CONTAINER

KENDALL MCGAW
5GM/100ML;300MG/100ML;
330MG/100ML

N19630 018
FEB 17, 1988

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

AP KENDALL MCGAM 5GM/100ML; 300MG/100ML; 450MG/100MLM N19630 024 FEB 17, 1988

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP KENDALL MCGAM 5GM/100ML; 300MG/100ML; 900MG/100MLM N19630 030 FEB 17, 1988

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

KENDALL MCGAM 5GM/100ML; 300MG/100ML; 110MG/100MLM N19630 006 FEB 17, 1988

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

AP ABBOTT LABS 5GM/100ML; 14.9MG/100ML; 450MG/100MLM N18362 009 JUL 05, 1983

AP 5GM/100ML; 14.9MG/100ML; 450MG/100MLM N18362 005 MAR 28, 1988

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP ABBOTT LABS 5GM/100ML; 14.9MG/100ML; 900MG/100MLM N19691 002 MAR 24, 1988

AP 5GM/100ML; 14.9MG/100ML; 900MG/100MLM N19691 004 MAR 24, 1988

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER

ABBOTT LABS 5GM/100ML; 74.5MG/100ML; 225MG/100MLM N18365 002 JUL 05, 1983

5GM/100ML; 14.9MG/100ML; 225MG/100MLM N18365 006 MAR 28, 1988

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

ABBOTT LABS 5GM/100ML; 74.5MG/100ML; 300MG/100ML N18876 001 JAN 17, 1986

5GM/100ML; 14.9MG/100ML; 300MG/100MLM N18876 006 MAR 28, 1988

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP TRAVENOL LABS 5GM/100ML; 75MG/100ML; 900MG/100MLM N19308 004 APR 05, 1985

POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

AP ABBOTT LABS 5GM/100ML; 224MG/100ML; 450MG/100MLM N18362 006 MAR 28, 1988

POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP ABBOTT LABS 5GM/100ML; 224MG/100ML; 900MG/100MLM N19691 006 MAR 24, 1988

POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER

ABBOTT LABS 5GM/100ML; 224MG/100ML; 225MG/100MLM N18365 008 MAR 28, 1988

POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

ABBOTT LABS 5GM/100ML; 224MG/100ML; 300MG/100MLM N18876 007 MAR 28, 1988

POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

AP ABBOTT LABS 5GM/100ML; 14.9MG/100ML; 450MG/100MLM N18362 010 JUL 05, 1983

5GM/100ML; 298MG/100ML; 450MG/100MLM N18362 007 MAR 28, 1988

POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP ABBOTT LABS 5GM/100ML; 14.9MG/100ML; 900MG/100MLM N19691 005 MAR 24, 1988

5GM/100ML; 298MG/100ML; 900MG/100MLM N19691 008 MAR 24, 1988

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER
ABBOTT LABS

5GM/100ML;14.9MG/100ML;
225MG/100ML N18365 001
5GM/100ML;298MG/100ML;
225MG/100ML N18365 009

MAR 28, 1988

POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER
ABBOTT LABS

5GM/100ML;14.9MG/100ML;
300MG/100ML

N18876 002
JAN 17, 1986

5GM/100ML;298MG/100ML;
300MG/100ML

N18876 008
MAR 28, 1988

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER
ABBOTT LABS

5GM/100ML;224MG/100ML;
450MG/100ML

N18362 002

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
ABBOTT LABS

5GM/100ML;224MG/100ML;
900MG/100ML

N19691 007
MAR 24, 1988

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER
ABBOTT LABS

5GM/100ML;224MG/100ML;
225MG/100ML

N18365 003
JUL 05, 1983

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER
ABBOTT LABS

5GM/100ML;224MG/100ML;
300MG/100ML

N18876 003
JAN 17, 1986

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
TRAVENOL LABS

5GM/100ML;224MG/100ML;
900MG/100ML

N19308 006
APR 05, 1985

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER
ABBOTT LABS

5GM/100ML;298MG/100ML;
450MG/100ML

N18362 003

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
ABBOTT LABS

5GM/100ML;298MG/100ML;
900MG/100ML

N19691 009
MAR 24, 1988

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER
ABBOTT LABS

5GM/100ML;298MG/100ML;
225MG/100ML

N18365 004
JUL 05, 1983

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER
ABBOTT LABS

5GM/100ML;298MG/100ML;
300MG/100ML

N18876 004
MAR 28, 1988

POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER
ABBOTT LABS

5GM/100ML;74.5MG/100ML;
450MG/100ML

N18362 008
MAR 28, 1988

5GM/100ML;14.9MG/100ML;
450MG/100ML

N18362 004
MAR 28, 1988

POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
ABBOTT LABS

5GM/100ML;74.5MG/100ML;
900MG/100ML

N19691 001
MAR 24, 1988

5GM/100ML;14.9MG/100ML;
900MG/100ML

N19691 003
MAR 24, 1988

POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER

5GM/100ML;74.5MG/100ML;
225MG/100ML

N18365 005
MAR 28, 1988

5GM/100ML;14.9MG/100ML;
225MG/100ML

N18365 007
MAR 28, 1988

POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER
ABBOTT LABS

5GM/100ML;74.5MG/100ML;
300MG/100ML

N18876 005
MAR 28, 1988

5GM/100ML;14.9MG/100ML;
300MG/100ML

N18876 009
MAR 28, 1988

DEXTROSE; SODIUM CHLORIDEINJECTABLE; INJECTION

DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

KENDALL MCGAW 10GM/100ML; 110MG/100ML N19631 011
FEB 24, 1988

DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

KENDALL MCGAW 10GM/100ML; 200MG/100ML N19631 012
FEB 24, 1988

DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER

KENDALL MCGAW 10GM/100ML; 330MG/100ML N19631 013
FEB 24, 1988

DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

KENDALL MCGAW 10GM/100ML; 450MG/100ML N19631 014
FEB 24, 1988

DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

KENDALL MCGAW 10GM/100ML; 900MG/100ML N19631 015
FEB 24, 1988

DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

KENDALL MCGAW 2.5GM/100ML; 110MG/100ML N19631 001
FEB 24, 1988

DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

KENDALL MCGAW 2.5GM/100ML; 200MG/100ML N19631 002
FEB 24, 1988

DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER

KENDALL MCGAW 2.5GM/100ML; 330MG/100ML N19631 003
FEB 24, 1988

DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

KENDALL MCGAW 2.5GM/100ML; 450MG/100ML N19631 004
FEB 24, 1988

DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

KENDALL MCGAW 2.5GM/100ML; 900MG/100ML N19631 005
FEB 24, 1988

DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

KENDALL MCGAW 5GM/100ML; 110MG/100ML N19631 006
FEB 24, 1988

DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

KENDALL MCGAW 5GM/100ML; 200MG/100ML N19631 007
FEB 24, 1988

DEXTROSE; SODIUM CHLORIDEINJECTABLE; INJECTION

DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER

AP KENDALL MCGAW 5GM/100ML; 330MG/100ML N19631 008
FEB 24, 1988

DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

AP KENDALL MCGAW 5GM/100ML; 450MG/100ML N19631 009
FEB 24, 1988

DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP KENDALL MCGAW 5GM/100ML; 900MG/100ML N19631 010
FEB 24, 1988

DIAZEPAMTABLET; ORALQ-PAM

AB QUANTUM PHARMCS

2MGM

N72431 001
APR 29, 1988

AB 5MGM

N72432 001
APR 29, 1988

AB 10MGM

N72433 001
APR 29, 1988

DIAZOXIDEINJECTABLE; INJECTIONDIAZOXIDE

AP QUAD PHARMS

15MG/ML

N71908 001
JAN 26, 1988

DOPAMINE HYDROCHLORIDEINJECTABLE; INJECTION

DOPAMINE HCL IN DEXTROSE 5% IN PLASTIC CONTAINER

AP BAXTER

80MG/100ML

N19615 001
MAR 27, 1987

AP 160MG/100ML

160MG/100ML

N19615 002
MAR 27, 1987

AP 320MG/100ML

320MG/100ML

N19615 003
MAR 27, 1987

AP 640MG/100ML

640MG/100ML

N19615 004
MAR 27, 1987

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOPAMINE HCL IN DEXTROSE 5% IN PLASTIC CONTAINER

AB / TRAVENCO LABS / 30MG/100ML / N16515/001 / MAR/27, 1987
 AB / 160MG/100ML / N16515/002 / MAR/27, 1987
 AB / 320MG/100ML / N16515/003 / MAR/27, 1987
 AB / 640MG/100ML / N16515/004 / MAR/27, 1987

DOXYCYCLINE HYCLATE

INJECTABLE; INJECTION

DOXYCYCLINE

AP BEN VENUE LABS / EQ 100MG BASE/VIAL / N62569 001 / MAR 09, 1988
 AP / EQ 200MG BASE/VIAL / N62569 002 / MAR 09, 1988

DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL

AP ABBOTT LABS / 2.5MG/ML / N71981 001 / FEB 29, 1988
 AP DUPONT CRI CARE / 2.5MG/ML / N71645 001 / APR 07, 1988

DOXEPIH HYDROCHLORIDE

CAPSULE; ORAL

ADAPIN

AB PENNWALT / EQ 150MG BASE / N16987 007 / APR 13, 1987

DOXEPIH HCL

BARR LABS

AB / EQ 25MG BASE / N71502 001 / FEB 18, 1988
 AB / EQ 50MG BASE / N71653 001 / FEB 18, 1988
 AB / EQ 75MG BASE / N71654 001 / FEB 18, 1988
 AB / EQ 100MG BASE / N71521 001 / FEB 18, 1988

CHELSEA LABS

AB / EQ 75MG BASE / N71763 001 / FEB 09, 1988
 AB / EQ 150MG BASE / N71764 001 / FEB 09, 1988

LEDERLE LABS

AB / EQ 10MG BASE / N71685 001 / JAN 05, 1988
 AB / EQ 25MG BASE / N71686 001 / JAN 05, 1988
 AB / EQ 50MG BASE / N71673 001 / JAN 05, 1988

AB / EQ 75MG BASE / N71674 001 / JAN 05, 1988
 AB / EQ 100MG BASE / N71675 001 / JAN 05, 1988
 AB / EQ 150MG BASE / N71676 001 / JAN 05, 1988

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

DOXYCYCLINE HYCLATE

VITARINE

AB / EQ 50MG BASE / N62780 001 / APR 12, 1988

DROPERIDOL; FENTANYL CITRATE

INJECTABLE; INJECTION

FENTANYL CITRATE AND DROPERIDOL

> ADD > ABBOTT LABS / 2.5MG/ML / N71982 001 / MAY 04, 1988
 > ADD > EQ 0.05MG BASE/ML

INNOVAR

JANSSEN PHARMA

> ADD > AP / 2.5MG/ML / N16049 001
 > ADD > EQ 0.05MG BASE/ML

EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

REVERSOL

ORGANON

> ADD > AP / 10MG/ML / N89624 001 / MAY 13, 1988
 > ADD >

ENALAPRILAT

INJECTABLE; INJECTION

VASOTEC

MS&D RES LABS

> ADD > 1.25MG/ML / N19309 001 / FEB 09, 1988

EPINEPHRINE; LIDOCAINE HYDROCHLORIDEERYTHROMYCIN ETHYLSUCCINATE; SULFISOXAZOLE ACETYLINJECTABLE; INJECTIONOCTOCAINE

> DLT > /AB/ /NOVOCOL/CHEN/ /0.01MG/ML; 2Z/

> ADD > AP /NOVOCOL PHARM/ 0.01MG/ML; 2Z/

> DLT > /AB/ /NOVOCOL/CHEN/ /0.02MG/ML; 2Z/

> ADD > AP /NOVOCOL PHARM/ 0.02MG/ML; 2Z/

GRANULE; ORALERYTHROMYCIN ETHYLSUCCINATE AND SULFISOXAZOLE ACETYL

> ADD > /N84048/001/

> ADD > AB BARR LABS EQ 200MG BASE/5 ML;

> ADD > N84048 001 EQ 600MG BASE/5MLM

> ADD > /N84048/002/

> ADD > N84048 002

> ADD > AB

> ADD > PEDIAZOLE EQ 200MG BASE/5ML;

> ADD > ROSS LABS EQ 600MG BASE/5MLM

> ADD > N50529 001

ERGOLOID MESYLATESTABLET; ORAL

> DLT > /AB/ /ERGOLOID MESYLATES/

> DLT > /AB/ /CHELSEA LABS/

> DLT > /AB/ /CHELSEA LABS/

GERMAL

> ADD > AB

> ADD > AB

> ADD > AB

1MG

/N88207/001/

/MAR/22/1984/

MAR 22, 1984

TABLET; SUBLINGUALCIRCANOL

> DLT > /AA/ /RIKER LABS/

> DLT > /AA/ /RIKER LABS/

> ADD > AA

> ADD > AA

> ADD > AA

/0.5MG/

1MG

/N84868/001/

/N85809/001/

> DLT > /AA/ /ERGOLOID MESYLATES/

> DLT > /AA/ /CHELSEA LABS/

> DLT > /AA/ /CHELSEA LABS/

GERMAL

> ADD > AA

> ADD > AA

> ADD > AA

0.5MG

1MG

/N86189/001/

/N86188/001/

ERYTHROMYCINSOLUTION; TOPICALETS-2Z

> ADD > AA

> ADD > AA

> ADD > AA

2ZM

N62687 001

FEB 05, 1988

ERYTHROMYCIN STEARATETABLET; ORALBRISTAMYCIN

> DLT > /AB/ /BRISTOL LABS/

> DLT > /AB/ /BRISTOL LABS/

> ADD > AB

> ADD > AB

> ADD > AB

/EQ 250MG BASE/

/EQ 250MG BASE/

/EQ 250MG BASE/

/N61304/001/

/N61304/001/

/N61887 001

ERYPAR

> DLT > /AB/ /PARKE DAVIS/

> DLT > /AB/ /PARKE DAVIS/

> ADD > AB

> ADD > AB

> ADD > AB

/EQ 250MG BASE/

/EQ 250MG BASE/

/EQ 250MG BASE/

/N62032/001/

/N62032/002/

/N62032 001

ERYTHROCIN STEARATE

> DLT > /AB/ /ABBOTT LABS/

> DLT > /AB/ /ABBOTT LABS/

> ADD > AB

> ADD > AB

> ADD > AB

/EQ 125MG BASE/

/EQ 125MG BASE/

/EQ 125MG BASE/

/N60359/002/

/N60359 002

/N60359 002

ERYTHROMYCIN STEARATE

> DLT > /AB/ /LEDERLE LABS/

> DLT > /AB/ /LEDERLE LABS/

> ADD > AB

> ADD > AB

> ADD > AB

/EQ 500MG BASE/

/EQ 500MG BASE/

/EQ 500MG BASE/

/N62089/001/

/N62089/002/

/N62089 001

PFIZER-E

> DLT > /AB/ /PFIZER LABS/

> DLT > /AB/ /PFIZER LABS/

> ADD > AB

> ADD > AB

> ADD > AB

/EQ 500MG BASE/

/EQ 500MG BASE/

/EQ 500MG BASE/

/N61791/002/

/N61791 002

/N61791 002

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21

NORETHIN 1/35E-21

AB

SEARLE PHARMS

0.035MG; 1MG

N71480 001
APR 12, 1988

NORETHINDRONE AND ETHINYL ESTRADIOL

AB

WATSON LABS

0.035MG; 0.5MG AND 1MG

N71043 001
APR 01, 1988

ORTHO-NOVUM 10/11-21

AB

ORTHO PHARM

0.035MG; 0.5MG AND 1MG

N18354 001
JAN 11, 1982

TABLET; ORAL-28

NORETHIN 1/35E-28

AB

SEARLE PHARMS

0.035MG; 1MG

N71481 001
APR 12, 1988

NORETHINDRONE AND ETHINYL ESTRADIOL

AB

WATSON LABS

0.035MG; 0.5MG AND 1MG

N71044 001
APR 01, 1988

ORTHO-NOVUM 10/11-28

AB

ORTHO PHARM

0.035MG; 0.5MG AND 1MG

N18354 002
JAN 11, 1982

FENOPROFEN CALCIUM

CAPSULE; ORAL

FENOPROFEN CALCIUM

QUANTUM PHARMS

AB

EQ 200MG BASE

N72214 001
AUG 17, 1988 : APR 14, 1988

AB

QUANTUM PHARMS

EQ 300MG BASE

N71738 001
AUG 17, 1988 : APR 14, 1988

AB

HALFON

EQ 300MG BASE

N17604 002

AB

DISTA PRODS

EQ 200MG BASE

N17604 003

TABLET; ORAL

FENOPROFEN CALCIUM

LEDERLE LABS

AB

EQ 600MG BASE

N72326 001
AUG 17, 1988 : APR 20, 1988

> ADD > AB

PUREPAC PHARM

EQ 600MG BASE

N72274 001
AUG 17, 1988 : MAY 02, 1988

> ADD > AB

QUANTUM PHARMS

EQ 600MG BASE

N72194 001
AUG 17, 1988 : APR 14, 1988

AB

HALFON

EQ 600MG BASE

N17710 001

FLUOCINOLONE ACETONIDE

OIL; TOPICAL

DERMA-SMOOTH/FS
HILL DERM

0.01%

N19452 001
FEB 03, 1988

FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE

AB

CLAY PARK LABS

0.05%

N71790 001
JUL 13, 1988 : APR 25, 1988

FLUOROURACIL

INJECTABLE; INJECTION

FLUOROURACIL

AP

BEN VENUE LABS

50MG/ML

N89508 001
JAN 26, 1988

FLUPHENAZINE HYDROCHLORIDE

TABLET; ORAL

PERMITIL

BP

SCHERING

1.5MG

N12034/004/

BP

SCHERING

2.5MG

N12034/004/

BP

SCHERING

5MG

N12034/005/

BP

SCHERING

10MG

N12034/006/

FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL

FLURAZEPAM HCL

AB

HALSEY DRUG

15MG

N71808 001
JAN 07, 1988

AB

HALSEY DRUG

30MG

N71809 001
JAN 07, 1988

FOLIC ACID

TABLET; ORAL

/AA/ FOLIC ACID
/BARR/LABS/

/1MG/

3 BARR LABS

/N89177/001/
/JAN/88/1985/
N89177 001
JAN 08, 1986

1MG

AA

CONCENTRATE; ORAL
HALOPERIDOL INTENSOL
ROXANE LABS

EQ 2MG BASE/MLM

N72045 001
APR 12, 1988FUROSEMIDE

INJECTABLE; INJECTION

/AB/ FUROSEMIDE
/PARKE/DAVIS/

/10MG/ML/

AP WARNER CHILCOTT

/N18420/001/
/FEB/88/1982/
N18420 001
FEB 26, 1982

10MG/ML

TABLET; ORAL

/AB/ FUROSEMIDE
BARR LABS

80MG

/AB/ DANBURY PHARMA

80MG

N70100 001
JAN 26, 1988
N71594 001
FEB 09, 1988HALOPERIDOL

TABLET; ORAL

HALDOL SOLUTAB
/3/MCNEIL/LABS/
3 MCNEIL PHARM/1MG/
1MG/N17079/001/
N17079 001

/AB/ BARR LABS

5MG

/AB/ BARR LABS

10MG

/AB/ BARR LABS

20MG

/AB/ CORD LABS

10MG

/AB/ CORD LABS

20MG

N71212 001
JAN 07, 1988
N71173 001
JAN 07, 1988
N71177 001
JAN 07, 1988
N71210 001
MAR 11, 1988
N71211 001
MAR 11, 1988

INJECTABLE; INJECTION

HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER/AB/ HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER
/ABBOTT/LABS/

/10,000 UNITS/100ML/

3 ABBOTT LABS

10,000 UNITS/100ML

/N19339/001/
/MAR/87/1985/
N19339 003
MAR 27, 1985HEPARIN SODIUM 1000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER/AB/ HEPARIN SODIUM 1000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
/BAXTER/

200 UNITS/100ML

N18609 001
APR 28, 1982/AB/ HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER
/TRAVENOL/LABS/

/200 UNITS/100ML/

/N18609/001/
/MAR/87/1985/
/APR/88/1982/HEPARIN SODIUM 2000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER/AB/ HEPARIN SODIUM 2000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
/ABBOTT/LABS/

/5,000 UNITS/100ML/

/N19339/001/
/MAR/87/1985/
N19339 001
MAR 27, 1985HEPARIN SODIUM 2000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER/AB/ HEPARIN SODIUM 2000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
/BAXTER/

200 UNITS/100ML

N18609 002
APR 28, 1982HEPARIN SODIUM 2000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER/AB/ HEPARIN SODIUM 2000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
/TRAVENOL/LABS/

/200 UNITS/100ML/

/N18609/001/
/APR/88/1982/HALOPERIDOL LACTATE

CONCENTRATE; ORAL

/AA/ HALDOL
/MCNEIL/LABS/
MCNEIL PHARM/EQ 2MG BASE/ML/
EQ 2MG BASE/ML/N15922/001/
N15922 001

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

METHYLDOPA AND HYDROCHLOROTHIAZIDE

AB NOVOPHARM 15MG;250MG

AB 25MG;250MG

AB 30MG;500MG

AB 50MG;500MG

/AB/ /PUREPAC/PHARM/ 50MG;500MG/

a PUREPAC PHARM 50MG;500MG

AB ZENITH LABS 15MG;250MG

AB 25MG;250MG

AB 30MG;500MG

AB 50MG;500MG

N71819 001

APR 08, 1988

N71820 001

APR 08, 1988

N71821 001

APR 08, 1988

N71822 001

APR 08, 1988

/N70689/001/

/APR/24, 1988/

N70689 001

APR 24, 1986

N71458 001

MAR 08, 1988

N71459 001

MAR 08, 1988

N71460 001

MAR 08, 1988

N71461 001

MAR 08, 1988

CAPSULE; ORAL

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

AB VITARINE 25MG;50MG

N71737 001

FEB 12, 1988

TABLET; ORAL

MAXZIDE-25

> ADD >

> ADD >

> ADD >

25MG;37.5MG

N19129 003

MAY 13, 1988

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

AB DANBURY PHARMA 50MG;75MG

N71969 001

JAN 15, 1988

> ADD >

> ADD >

> ADD >

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

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50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

50MG;75MG

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HCL AND HYDROCHLOROTHIAZIDE

AB SIDMAK LABS 25MG;40MG

AB 25MG;80MG

AB 25MG;40MG

AB 25MG;80MG

N72042 001

MAR 14, 1988

N72043 001

MAR 14, 1988

N71771 001

JAN 26, 1988

N71772 001

JAN 26, 1988

CREAM; TOPICAL

HYDROCORTISONE

AT NASKA PHARMA 1/2

N89706 001

MAR 10, 1988

N89682 001

MAR 10, 1988

N87795 001

MAY 03, 1983

N87427 001

APR 04, 1988

/N87795/001/

/MAY/03, 1983/

/N87795/001/

/MAY/03, 1983/

/N87795/001/

/MAY/03, 1983/

/N87795/001/

/MAY/03, 1983/

/N87795/001/

/MAY/03, 1983/

/N87795/001/

/MAY/03, 1983/

/N87795/001/

/MAY/03, 1983/

/N87795/001/

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/MAY/03, 1983/

/N87795/001/

/MAY/03, 1983/

/N87795/001/

/MAY/03, 1983/

/N87795/001/

/MAY/03, 1983/

/N87795/001/

/MAY/03, 1983/

/N87795/001/

/MAY/03, 1983/

/N87795/001/

LOTION; TOPICAL

BETA-HC

AT BETA DERM 1/2

N89495 001

JAN 25, 1988

HYDROCORTISONE

AT NASKA PHARMA 1/2

N89705 001

APR 25, 1988

OINTMENT; TOPICAL

HC (HYDROCORTISONE)

> DLT > /AT/

> DLT > /AT/

> ADD > /AT/

> ADD > /AT/

> ADD > /AT/

> ADD > /AT/

> ADD > /AT/

> ADD > /AT/

> ADD > /AT/

> ADD > /AT/

> ADD > /AT/

> ADD > /AT/

> ADD > /AT/

> ADD > /AT/

/0.52/

/1.2/

/1.2/

/1.2/

/1.2/

/1.2/

/1.2/

/1.2/

/1.2/

/1.2/

/1.2/

/1.2/

/1.2/

/1.2/

/N80481/001/

/N80481/001/

/N80481/001/

/N80481/001/

/N80481/001/

/N80481/001/

/N80481/001/

/N80481/001/

/N80481/001/

/N80481/001/

/N80481/001/

/N80481/001/

/N80481/001/

/N80481/001/

HYDROCORTISONE

OINTMENT; TOPICAL

AI HYDROCORTISONE
NASKA PHARMA

1/2M

N89704 001
MAR 10, 1988

> ADD > AB
> ADD > AB
> ADD > AB
> ADD > AB
> ADD > AB

10MG

N89366 001
MAY 02, 1988
N89117 001
MAY 02, 1988
N89396 001
MAY 02, 1988

TABLET; ORAL

/BP/ HYDROCORTISONE
BARR/LABS/
BARR LABS

1/20MG/
20MG

N83999/001/
N83999 001

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SUSPENSION/DROPS; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

> ADD >
> ADD > AI
> ADD >
> ADD >

STERIS LABS
1/2; EQ 3.5MG BASE/ML;
10,000 UNITS/MLM

N62874 001
MAY 11, 1988

AB

IBUPROFEN

800MG

N72137 001
FEB 05, 1988
N72064 001
JAN 14, 1988

AB

INVAMED

400MG

AB

600MG

N72065 001
JAN 14, 1988
N71938 001
JAN 14, 1988

HYDROCORTISONE ACETATE; PRAMOXINE HYDROCHLORIDE

AEROSOL; TOPICAL

EPIFOAM

REED & CARRICK

1/2; 1/2;
HYDROCORTISONE ACETATE 1/2 AND PRAMOXINE HCL 1/2

N86457 001
N89440 001
MAY 17, 1988

AB

MEDICOPHARMA

400MG

N71644 001
FEB 01, 1988
N71964 001
FEB 01, 1988

AB

PUREPAC PHARM

800MG

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE HCL

/AP/ HYDROXYZINE HCL
ALTANA

1/25MG/ML

N87273/001/
APR 20, 1982

/AP/

1/50MG/ML

N87273/002/
APR 20, 1982

a ALTANA

25MG/ML

N87273 001

a

50MG/ML

N87273 002

APR 20, 1982

SYRUP; ORAL

HYDROXYZINE HCL

NASKA PHARMA

10MG/5MLM

N88785 001
FEB 03, 1988

AB

INDOMETHACIN

25MG

N71342 001
APR 18, 1988

AB

NOVOPHARM

50MG

N71343 001
APR 18, 1988

HYDROXYZINE HYDROCHLORIDE

TABLET; ORAL

HYDROXYZINE HCL

HALSEY DRUG

10MG

N89366 001
MAY 02, 1988

25MG

N89117 001
MAY 02, 1988

50MG

N89396 001
MAY 02, 1988

IBUPROFEN

TABLET; ORAL

IBUPROFEN

800MG

N72137 001
FEB 05, 1988

AB

HALSEY DRUG

400MG

N72064 001
JAN 14, 1988

AB

INVAMED

600MG

N72065 001
JAN 14, 1988

AB

MEDICOPHARMA

800MG

N71938 001
JAN 14, 1988

AB

PUREPAC PHARM

400MG

N71644 001
FEB 01, 1988

AB

PUREPAC PHARM

800MG

N71964 001
FEB 01, 1988

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

PRAMINE

AB/ PRAMINE
BANMAX/PHARMS/
AB/

10MG

N83827/001/
APR 18, 1988

AB

BANMAX PHARMS

25MG

N83827/002/
APR 18, 1988

AB

BANMAX PHARMS

50MG

N83827/003/
APR 18, 1988

a

BANMAX PHARMS

10MG

N83827/001
APR 18, 1988

a

BANMAX PHARMS

25MG

N83827/002
APR 18, 1988

INDOMETHACIN

CAPSULE; ORAL

INDOMETHACIN

25MG

N71342 001
APR 18, 1988

AB

NOVOPHARM

50MG

N71343 001
APR 18, 1988

AB

NOVOPHARM

50MG

N71343 001
APR 18, 1988

IOHEXOL

INJECTABLE; INJECTION

OMIPAGUE 180

/AP/ 12/STERLING/DRUGS/

/18.82/

AP STERLING DRUG-

38.82

/N18956/001/
DEC 26, 1985

AP
KETAMINE HCL
QUAD PHARMS
EQ 10MG BASE/ML
AP
EQ 50MG BASE/ML
AP
EQ 100MG BASE/ML

N71949 001
APR 11, 1988
N71950 001
APR 11, 1988
N71951 001
APR 11, 1988

ISOSORBIDE DINITRATE

TABLET; ORAL

ISOSORBIDE DINITRATE

BARR LABS

30MG

AB

CORD LABS

5MG

AB

10MG

20MG

AB

DANBURY PHARMA

5MG

AB

10MG

30MG

AB

PAR PHARM

2.5MG

AB

5MG

10MG

AB

2.5MG

5MG

TABLET; SUBLINGUAL

ISOSORBIDE DINITRATE

CORD LABS

2.5MG

AB

DANBURY PHARMA

5MG

AB

10MG

2.5MG

AB

5MG

KETAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

KETALAR

PARKE DAVIS

EQ 10MG BASE/ML

EQ 50MG BASE/ML

EQ 100MG BASE/ML

N16812 001
N16812 002
N16812 003

LEVONORDEFIN; MEPIVACAIN HYDROCHLORIDE

INJECTABLE; INJECTION

ISOCATINE HCL W/ LEVONORDEFIN

/NOVOCOL/CHEN/

NOVOCOL PHARM

POLOCATINE W/ LEVONORDEFIN

ASTRA PHARM PRODS

0.05MG/ML; 2Z

/N84697/001/
N84697 001

N89517 001
APR 14, 1988

LACTULOSE

SYRUP; ORAL

LACTULOSE

3 KALI DUPHAR

/AA/ /ROXANE/LABS/

10GM/15ML

10GM/15ML

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

BEN VENUE LABS

EQ 100MG BASE/VIAL

EQ 100MG BASE/VIAL

EQ 100MG BASE/VIAL

N89717 001
MAR 28, 1988
N08107 004
MAY 23, 1988
N89636 001
DEC 24, 1987

LEVODOPA

CAPSULE; ORAL

BENDOPA

/ICN/PHARMS/

100MG

250MG

500MG

/N16948/001/
N16948 001
N16948 002
N16948 003
N16948 001
N16948 002

100MG

250MG

500MG

100MG

250MG

500MG

MEFENAMIC ACID

CAPSULE; ORAL

MEFENAMIC ACID

VITARINE

250MG

N72179 001
APR 21, 1988

/AB/

ENOVID-E
/SEARLE/
a SEARLE

/0.1MG; 2.5MG/
0.1MG; 2.5MG

/N10976; 006/
N10976 006

PONSTEL

PARKE DAVIS PR

250MG

N15034 003

METAPROTERENOL SULFATE

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ISOGAINE HCL

/NOVOCOL/CHEN/

NOVOCOL PHARM

/3 1/2/
3 1/2

/N80925; 001/
N80925 001

METAPROTERENOL SULFATE
PHARM BASICS

10MG
20MG

N15874 002
N15874 001

MEPROBAMATE

TABLET; ORAL

MEPROBAMATE

/MALLARD/

a MALLARD

/500MG/
400MG

/N15072; 002/
N15072 002

METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL

/BARR/LABS/

a BARR LABS

/500MG/
500MG

/N84488; 001/
N84488 001

a PHARM BASICS

200MG

N87825 001

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE

LEDERLE LABS

EQ 1GM BASE/VIAL

N11719 009
APR 07, 1988

MESTRANOL; NORETHINDRONE

TABLET; ORAL-21

NORETHIN 1/50M-21

SEARLE PHARMS

0.05MG; 1MG

N71539 001
APR 12, 1988

METHOXSALEN

CAPSULE; ORAL

OXSORALEN

/ELDER/PHARMS/

8-HOP

ELDER PHARMS

/10MG/
10MG

/N80948; 001/
N80948 001

TABLET; ORAL-28

NORETHIN 1/50M-28

SEARLE PHARMS

0.05MG; 1MG

N71540 001
APR 12, 1988

MESTRANOL; NORETHYNODREL

TABLET; ORAL-20

ENOVID

/SEARLE/

a SEARLE

/0.075MG; 5MG/
0.075MG; 5MG

/N10976; 004/
N10976 004

TABLET; ORAL

METHYLDOPA

CORD LABS

125MG

N71700 001
MAR 02, 1988

METHYLDOPA

TABLET; ORAL

METHYLDOPA

HALSEY DRUG

AB 125MG
AB 250MG
AB 500MG

N71751 001
MAR 28, 1988
N71752 001
MAR 28, 1988
N71753 001
MAR 28, 1988

INJECTABLE; INJECTION

MORPHINE SULFATE PF

ABBOTT LABS

> ADD > AP
> ADD > AP
> ADD > AP
> ADD > AP

N71849 001
MAY 11, 1988
N71850 001
MAY 11, 1988

0.5MG/ML

1MG/ML

TABLET, CONTROLLED RELEASE; ORAL

MS CONTIN

PURDUE FRDRK

60MG

N19516 002
APR 08, 1988

METHYLPREDNISOLONE

TABLET; ORAL

MEDROL

UP JOHN

AB 24MG
AB 32MG

N11153 005
N11153 006

METHYLPREDNISOLONE

PAR PHARM

AB 16MG
AB 24MG
AB 32MG

N89207 001
APR 25, 1988
N89208 001
APR 25, 1988
N89209 001
APR 25, 1988

NAFTIFINE HYDROCHLORIDE

CREAM; TOPICAL

NAFTIN

HERBERT LABS

1/4

N19599 001
FEB 29, 1988

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE HCL

ELKINS SINN

0.02MG/ML

> ADD > AP
> ADD > AP
> ADD > AP
> ADD > AP
> ADD > AP
> ADD > AP
> ADD > AP

N71272 001
MAY 24, 1988
N71273 001
MAY 24, 1988
N71274 001
MAY 24, 1988
N71287 001
MAY 24, 1988
N72076 001
MAR 24, 1988
N72115 001
APR 27, 1988

1MG/ML

1MG/ML

1MG/ML

1MG/ML

1MG/ML

1MG/ML

INTL MEDTN SYS

AP

AP

NITROGLYCERIN

INJECTABLE; INJECTION

NITROGLYCERIN

LUITPOLD PHARMS

5MG/ML

N71492 001
MAY 24, 1988
N72034 001
MAY 24, 1988

5MG/ML

> ADD > AP
> ADD > AP
> ADD > AP
> ADD > AP

N62336 003
MAR 10, 1988

METOCLOPRAMIDE HYDROCHLORIDE

TABLET; ORAL

METOCLOPRAMIDE HCL

SIDMAK LABS

EQ 10MG BASE

N71250 001
FEB 03, 1988

MEZLOCILLIN SODIUM MONOHYDRATE

INJECTABLE; INJECTION

MEZLIN

MILES PHARM

EQ 20GM BASE/VIAL

N50549 005
MAR 02, 1988

EQ 20GM BASE/VIAL

N62372 004
MAR 02, 1988

MITIOMYCIN

INJECTABLE; INJECTION

MUTAMYCIN

BRISTOL MYERS

40MG/VIAL

NIZATIDINE

CAPSULE; ORAL
AXID
LILLY

150MG
300MG

N19508 001
APR 12, 1988
N19508 002
APR 12, 1988

CAPSULE; ORAL
OXAZEPAM
CORD LABS

10MG

N71813 001
APR 19, 1988

15MG

N71756 001
APR 19, 1988

30MG

N71814 001
APR 19, 1988

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL
AVENTYL HCL
LILLY

EQ 10MG BASE
EQ 25MG BASE
/EQ/10MG/BASE/
/EQ/25MG/BASE/

N14684 001
N14684 002
/N14684/001/
/N14684/002/

BD
BD
/BP/
/BP/
PAMELOR
SANDOZ PHARMS

EQ 10MG BASE
EQ 25MG BASE
/EQ/10MG/BASE/
/EQ/25MG/BASE/

N18013 001
N18013 002
/N18013/001/
/N18013/002/

NYSTATIN

SUSPENSION; ORAL
NYSTATIN
THAMES PHARMA

100,000 UNITS/ML

N62876 001
FEB 29, 1988

OXAZEPAM

CAPSULE; ORAL
OXAZEPAM
AM THERPTCS

10MG

N71955 001
MAR 03, 1988

15MG

N71956 001
MAR 03, 1988

30MG

N71957 001
MAR 03, 1988

CHELSEA LABS

10MG

N71661 001
MAR 02, 1988

15MG

N71662 001
MAR 02, 1988

30MG

N71663 001
MAR 02, 1988

OXYBUTYRIN CHLORIDE

TABLET; ORAL
DITROPAN
MARION LABS

5MG

N17577 001

OXYBUTYRIN CHLORIDE

PHARM BASICS

5MG

N70746 001
MAR 10, 1988

SERAX
WYETH

10MG

N15539 002

15MG

N15539 004

30MG

N15539 006

10MG

N15539/002/

15MG

N15539/004/

30MG

N15539/006/

ZAXOPAM

QUANTUM PHARMS

10MG

N70650 001

15MG

N70640 001

30MG

N70641 001

PENICILLIN V POTASSIUM

TABLET; ORAL

AB/ AB/	PENICILLIN V POTASSIUM/ PUREPAC PHARM/	/EQ 450MG BASE/ /EQ 500MG BASE/
	3 PUREPAC PHARM	EQ 250MG BASE
	3	EQ 500MG BASE

/N61571/002/
 /N61571/003/
 N61571 002
 N61571 003

IMG/MLM

SYRUP; ORAL
TRILAFON
/SCHERING/
a SCHERING

PHENDIMETRAZINE TARTRATE

TABLET; ORAL
PHENMETRAZINE TARTRATE
/BARR/LABS/

BARR LABS

POWDER FOR RECONSTITUTION; ORAL

N61740/002

AB/	PENICILLIN G POTASSIUM /PUREPAC/ PHARM/	2 PUREPAC PHARM
AB/		2

/N61588/002/
 /N61588/003/
 N61588 002
 N61588 003

POWDER FOR RECONSTITUTION; ORAL

N61758/002
N61758 002

CAPSULE, CONTROLLED RELEASE: ORAL

AB	<u>IONATHIN-30</u>	<u>EQ 30MG BASE</u>
	PENNWALT	
	<u>PHENTERMINE RESIN 30</u>	<u>EQ 30MG BASE</u>
AB	QUANTUM PHARMS	

N11613 002
N89120 001
FEB 04, 1988

PIPERACETAZINE

TABLET; ORAL

GUIDE

/DOW/PHARM/ 10MG
a DOW PHARMS
/DOW/PHARM/ 25MG
a DOW PHARMS

/N13615/001/
N13615 001
/N13615/002/
N13615 002

AB SUPERPHARM

5MG

N88865 001
OCT 25, 1984
N88866 001
OCT 25, 1984
N88867 001
OCT 25, 1984

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM
BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE, ANHYDROUS

POWDER FOR RECONSTITUTION; ORAL

COLOHILITE

DYNAPHARM

227.1GM/PACKET; 2.82GM/PACKET;
6.36GM/PACKET; 5.53GM/PACKET;
21.5GM/PACKET

N71320 001
APR 20, 1988

PROCAINAMIDE HYDROCHLORIDE

COLYTE

REED & CARNICK

227.1GM/PACKET; 2.82GM/PACKET;
6.36GM/PACKET; 5.53GM/PACKET;
21.5GM/PACKET

N18983 004
OCT 26, 1984

INJECTABLE; INJECTION

PROCAINAMIDE HCL

WARNER CHILCOTT

100MG/ML

N89528 001
MAY 03, 1988
N89529 001
MAY 03, 1988

POTASSIUM CHLORIDE

INJECTABLE; INJECTION
POTASSIUM CHLORIDE

STERIS LABS

2MEQ/ML

N89163 001
MAR 10, 1988

PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

ELKINS SINN

EQ 5MG BASE/ML

N89523 001
MAY 03, 1988
N89637 001
FEB 01, 1988
N89638 001
FEB 01, 1988
N89703 001
APR 07, 1988

PREDNISOLONE

TABLET; ORAL

PREDNISOLONE

a BARR LABS

/N84426/002/
N84426 002

5MG

PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROMETHAZINE HCL

MARSAM PHARMS

25MG/ML

N89463 001
MAY 02, 1988
N89477 001
MAY 02, 1988

PROMETHAZINE HYDROCHLORIDE

TABLET; ORAL

PROMETHAZINE HCL

/BP/ /BARR/LABS/ /1.5MG/ /N84555/001/

/BP/ /BARR/LABS/ /2.5MG/ /N84554/001/

/BP/ /BARR/LABS/ /5MG/ /N84557/001/

2 BARR LABS /12.5MG/ N84555 001

2 /25MG/ N84554 001

2 /50MG/ N84557 001

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

PROPOXYPHENE HCL

/AA/ /BANMAX/PHARMS/ /65MG/ /N83184/001/

/AA/ /BANMAX/PHARMS/ /65MG/ N83184 001

/AA/ /BARR/LABS/ /65MG/ /N83186/001/

2 BARR LABS /65MG/ N83186 001

PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HCL

AB SIDMAK LABS /10MG/ N71972 001

AB /20MG/ APR 06, 1988

AB /40MG/ N71973 001

AB /60MG/ APR 06, 1988

AB /80MG/ N71974 001

AB /90MG/ APR 06, 1988

AB /100MG/ N71975 001

AB /120MG/ APR 06, 1988

AB /140MG/ N71976 001

AB /160MG/ APR 06, 1988

AB /180MG/ N71977 001

AB /200MG/ APR 06, 1988

PROTAMINE SULFATE

INJECTABLE; INJECTION

PROTAMINE SULFATE

/AP/ /UPJOHN/ /50MG/VIAL/ /N87413/001/

/AP/ /UPJOHN/ /250MG/VIAL/ /N87413/001/

2 UPJOHN /50MG/VIAL/ N87413 001

2 /250MG/VIAL/ N87413 002

AUG 02, 1984

QUINESTROL

TABLET; ORAL

ESTROVIS

> DLT > /N84555/001/

> ADD > /N84554/001/

> DLT > /N84557/001/

> ADD > /N84555/001/

> ADD > /N84554/001/

> ADD > /N84557/001/

RAUMOLFIA SERPENTINA

TABLET; ORAL

RAUVAL

/BP/ /VALE/CHEN/ /50MG/ /N84168/002/

/BP/ /VALE/CHEN/ /100MG/ /N84168/004/

/BP/ /VALE/CHEN/ /100MG/ N84168 002

/BP/ /VALE/CHEN/ /100MG/ N84168 004

RESERPINE

TABLET; ORAL

RESERPINE

/BP/ /BARR/LABS/ /0.25MG/ /N80721/002/

/BP/ /BARR/LABS/ /0.25MG/ N80721 002

SECOBARBITAL SODIUM

CAPSULE; ORAL

SODIUM SECOBARBITAL

/AA/ /BARR/LABS/ /100MG/ /N84225/001/

/AA/ /BARR/LABS/ /100MG/ N84225 001

SODIUM CHLORIDE

INJECTABLE; INJECTION

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP KENDALL MCGAW /900MG/100MLM /N19635/002/

AP KENDALL MCGAW /900MG/100MLM MAR 09, 1988

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

AP KENDALL MCGAW /450MG/100MLM /N19635/001/

AP KENDALL MCGAW /450MG/100MLM MAR 09, 1988

SODIUM CHLORIDE 3% IN PLASTIC CONTAINER

AP KENDALL MCGAW /3GM/100MLM /N19635/003/

AP KENDALL MCGAW /3GM/100MLM MAR 09, 1988

SODIUM CHLORIDE 5% IN PLASTIC CONTAINER

AP KENDALL MCGAW /5GM/100MLM /N19022/001/

AP KENDALL MCGAW /5GM/100MLM NOV 01, 1983

AP KENDALL MCGAW /5GM/100MLM /N19635/004/

AP KENDALL MCGAW /5GM/100MLM MAR 09, 1988

SODIUM CHLORIDEINJECTABLE; INJECTION

AP SODIUM CHLORIDE 5% IN PLASTIC CONTAINER
TRAVENOL LABS 5GM/100ML

N19022 002
 NOV 01, 1983

AB SULINDAC
DANBURY PHARMA

150MG
200MG

N71891 001
 APR 03, 1990 : MAR 03, 1988
 N71795 001
 APR 03, 1990 : MAR 03, 1988

SULFAMETHOXAZOLETABLET; ORAL

/AB/ SULFAMETHOXAZOLE
BARR/LABS

500MG

3 BARR LABS

500MG

/N87189/001/
 /JUL/25/1983/
 N87189 001
 JUL 25, 1983

TEMASEPAM

CAPSULE; ORAL
TEMASEPAM
CORD LABS

15MG
30MG

N71427 001
 JAN 12, 1988
 N71428 001
 JAN 12, 1988

SULFAMETHOXAZOLE; TRIMETHOPRIMSUSPENSION; ORAL

TRIMETH/SULFA
NASKA PHARMA

> ADD >
 > ADD >
 > ADD >
 > ADD >
 > ADD >
 > ADD >

200MG/5ML; 40MG/5ML

N72289 001
 MAY 23, 1988
 N72398 001
 MAY 23, 1988
 N72399 001
 MAY 23, 1988

> ADD >
 > ADD >
 > ADD >
 > ADD >

TERCONAZOLE

SUPPOSITORY; VAGINAL
TERAZOL 3
ORTHO PHARM

80MG

N19641 001
 MAY 24, 1988

SULFISOXAZOLETABLET; ORAL

/AB/ SULFISOXAZOLE
BARR/LABS
 3 BARR LABS

500MG
500MG

/N84031/001/
 N84031 001

> ADD >
 > ADD >

AA THEOPHYLLINE
NASKA PHARMA

80MG/15ML

N89223 001
 MAY 27, 1988

SULINDACTABLET; ORAL

CLINORIL
MS&D

150MG
200MG

N17911 001
 N17911 002

AB THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER
ABBOTT LABS
320MG/100ML

320MG/100ML

N18649 006
 NOV 13, 1985
 N19211 006
 JAN 20, 1988

AB SULINDAC
AM THERPTCS

150MG
200MG

N72171 001
 APR 03, 1990 : MAY 23, 1988
 N72172 001
 APR 03, 1990 : MAY 23, 1988

BC THEOLAIR-SR
RIKER LABS
250MG

N86363 002
 JUL 16, 1987
 /N86363/002/
 /JUL/16/1987/

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

VANCOCIN HCL

LILLY

EQ 1GM BASE/VIAL

N60180 002

MAR 21, 1986

EQ 1GM BASE/VIAL

N62476 002

MAR 21, 1986

EQ 1GM BASE/VIAL

N62716 002

MAR 13, 1987

EQ 1GM BASE/VIAL

N62812 002

NOV 17, 1987

EQ 10GM BASE/VIAL

N62812 003

NOV 17, 1987

VANCOLED

LEDERLE LABS

EQ 1GM BASE/VIAL

N62682 002

MAR 30, 1988

EQ 10GM BASE/VIAL

N62682 005

MAY 11, 1988

EQ 2GM BASE/VIAL

N62682 003

MAY 11, 1988

EQ 5GM BASE/VIAL

N62682 004

MAY 11, 1988

VERAPAMIL HYDROCHLORIDE

TABLET; ORAL

CALAN

/SEARLE/

/160MG/

AB SEARLE PHARMS

40MG

N18817 003

FEB 23, 1988

160MG

N18817 004

FEB 23, 1988

ISOPTIN

KNOLL PHARM

40MG

N18593 003

NOV 23, 1987

VERAPAMIL HCL

CORD LABS

80MG

N71423 001

MAY 25, 1988

120MG

N71424 001

MAY 25, 1988

80MG

N71880 001

APR 05, 1988

120MG

N71881 001

APR 05, 1988

80MG

N71488 001

JAN 13, 1988

120MG

N71489 001

JAN 13, 1988

VINCRISTINE SULFATE

INJECTABLE; INJECTION

VINCRISTINE SULFATE

BULL LABS

1MG/VIAL

N71559 001

APR 11, 1988

2MG/VIAL

N71560 001

APR 11, 1988

5MG/VIAL

N71561 001

APR 11, 1988

1MG/VIAL

N71222 001

MAR 07, 1988

2MG/VIAL

N71223 001

MAR 07, 1988

5MG/VIAL

N71937 001

MAR 07, 1988

VINCRISTINE SULFATE PFS

BULL LABS

1MG/ML

N71484 001

APR 19, 1988

WATER FOR INJECTION, STERILE

LIQUID; N/A

STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

100/ML

N18632 002

APR 19, 1988

100/ML

N19633 001

FEB 29, 1988

XYLOSE

POWDER; ORAL

XYLOSE

/LYNE LABS/

/25GM/BOT/

N18856 001

MAR 26, 1987

2 LYNE LABS

25GM/BOT

MAR 26, 1987

ACETAMINOPHEN

SUPPOSITORY; RECTAL
TYLENOL
MCNEIL CONSUMER
/MCNEIL LABS/
/120MG/
/650MG/
/120MG/
/650MG/

N17756 002
N17756 001
/N17756/002/
/N17756/001/

INSULIN ZINC SUSP EXTENDED BIOSYNTHETIC HUMAN

INJECTABLE; INJECTION

HUMULIN U
/LILLY/

/40 UNITS/ML/
40 UNITS/ML

/N19571/001/
/JUN/10, 1987/
N19571 001
JUN 10, 1987

IBUPROFEN

TABLET; ORAL
IBUPROFEN

DANBURY PHARMA

200MG

N71905 001

INTERPHARM

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INVAMED

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MEDICOPHARMA

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MYLAN PHARMS

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ZENITH LABS

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LOPERAMIDE HYDROCHLORIDE

SOLUTION; ORAL

IMODIUM A-D

MCNEIL CONSUMER

1MG/5ML

N19487 001

NONOXYNOL-9

SPONGE; VAGINAL

TODAY

/V.I./

/164/

/N18683/001/
/APR/01, 1983/
N18683 001
APR 01, 1983

WHITEHALL LABS

1GM

INSULIN SEMISYNTHETIC PURIFIED HUMAN; INSULIN SUSP ISOPHANE
SEMISYNTHETIC PURIFIED HUMAN

INJECTABLE; INJECTION
MIXTARD HUMAN 70/30
NORDISK USA

30 UNITS/ML;
70 UNITS/ML

N19585 001
MAR 11, 1988

DRUG PRODUCTS IN THE DIVISION OF BLOOD AND BLOOD PRODUCTS / CUMULATIVE SUPPLEMENT NUMBER 5 / JAN '88 - MAY '88
APPROVED UNDER SECTION 505 OF THE ACT LIST

DEXTRAN 70, 6% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

MACRODEX(R)

PHARMACIA INC

6GM/100ML; 0.9GM/100ML

N 06826

BIOPHARMACEUTIC GUIDANCE AVAILABILITY

THE FOLLOWING IS A LIST OF GUIDANCES AVAILABLE FOR IN VIVO BIOEQUIVALENCE STUDIES AND IN VITRO DISSOLUTION TESTING AVAILABLE FROM THE DIVISION OF BIOEQUIVALENCE, HFN-250, ROOM 17B-06, 5600 FISHERS LANE, ROCKVILLE, MD 20857. COMMENTS AND SUGGESTIONS CONCERNING THESE GUIDANCES ARE ENCOURAGED AND SHOULD BE SENT TO THE DIVISION OF BIOEQUIVALENCE.

REFER BACK TO THE APPROVED DRUG PRODUCTS WITH THERAPEUTIC EQUIVALENCE EVALUATIONS, 8TH EDITION FOR A FULL LISTING OF BIOPHARMACEUTIC GUIDANCE AVAILABILITY DATA. ONLY NEW DATA WILL BE ADDED TO THE CUMULATIVE SUPPLEMENT.

NAME OF DRUG (DOSAGE FORM)	DATE	REVISED DATE
CARBAMAZEPINE (TABLET)	JAN 01, 1988	
CLINDAMYCIN (CAPSULE)	MAY 31, 1988	
CYCLOBENZAPRINE HYDROCHLORIDE (TABLET)	JAN 25, 1988	
DOXYCYCLINE HYCLATE (CAPSULE AND TABLET)	APR 11, 1988	
FENOPROFEN (CAPSULE AND TABLET)	AUG 27, 1987	FEB 03, 1988
INDOMETHACIN (CAPSULE)	JAN 27, 1988	
MESTRANOL; NORETHYNODREL (TABLET)	MAY 13, 1988	
METAPROTERENOL SULFATE (TABLET)	MAR 18, 1988	
NORETHINDRONE; ETHINYL ESTRADIOL (TABLET)	MAR 18, 1988	

ANDA SUITABILITY PETITIONS

THE FOLLOWING ARE TWO LISTS OF PETITIONS FILED UNDER SECTION 505(j)(2)(C) OF THE ACT WHERE THE AGENCY HAS DETERMINED THAT THE REFERENCED PRODUCT: (1) IS SUITABLE FOR SUBMISSION AS AN ANDA (PETITIONS APPROVED) OR (2) IS NOT SUITABLE FOR SUBMISSION AS AN ANDA (PETITIONS DENIED). THE DETERMINATION THAT AN ANDA WILL BE APPROVED IS NOT MADE UNTIL THE ANDA ITSELF IS SUBMITTED AND REVIEWED BY THE AGENCY. A COPY OF EACH PETITION IS LISTED BY DOCKET NUMBER ON PUBLIC DISPLAY IN FDA'S DOCKETS MANAGEMENT BRANCH, HFA-305, ROOM 4-62, 5600 FISHERS LANE, ROCKVILLE, MD 20857.

REFER BACK TO THE APPROVED DRUG PRODUCTS WITH THERAPEUTIC EQUIVALENCE EVALUATIONS, 8TH EDITION FOR A FULL LISTING OF ANDA SUITABILITY PETITIONS DATA. ONLY NEW DATA WILL BE ADDED TO THE CUMULATIVE SUPPLEMENT.

PETITIONS APPROVED

DRUG NAME DOSAGE FORM; ROUTE	STRENGTH (CONTAINER SIZE)	DOCKET NUMBER	PETITIONER	REASON FOR PETITION	STATUS
ASPIRIN; HYDROCODONE BITARTRATE TABLET; ORAL	325MG 5MG	87 P-0376/CP	ANABOLIC	NEW STRENGTH	APPROVED FEB 12, 1988
ASPIRIN; HYDROCODONE BITARTRATE TABLET; ORAL	650MG 5MG	87 P-0376/ CP0002	ANABOLIC	NEW STRENGTH	APPROVED FEB 12, 1988
CHLORZOXAZONE CAPSULE; ORAL	500MG	82 N-0032/ CP0006	MIKART	NEW DOSAGE FORM	APPROVED JAN 13, 1988

ANDA SUITABILITY PETITIONS

PETITIONS APPROVED

DRUG NAME DOSAGE FORM; ROUTE	STRENGTH (CONTAINER SIZE)	DOCKET NUMBER	PETITIONER	REASON FOR PETITION	STATUS
CISPLATIN INJECTABLE; INJECTION	1MG/ML (10ML/VIAL) (50ML/VIAL) (100ML/VIAL)	87 P-0421/CP	BULL LABS	NEW DOSAGE FORM NEW STRENGTH	APPROVED FEB 29, 1988
CISPLATIN INJECTABLE; INJECTION	1MG/ML (20ML/VIAL)	88 P-0010/CP	LYPHOMED	NEW DOSAGE FORM NEW STRENGTH	APPROVED APR 01, 1988
FLUOROURACIL INJECTABLE; INJECTION	50MG/ML (5ML/VIAL)	88 P-0052/CP	BEN VENUE LABS	NEW STRENGTH	APPROVED MAR 21, 1988
HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE SOFT GELATIN CAPSULE; ORAL	1.5MG 5MG	88 P-0061/CP	KLEINFELD, KAPLAN AND BECKER	NEW DOSAGE FORM	APPROVED MAY 12, 1988
HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE SOLUTION; ORAL	25MG/5ML 40MG/5ML	87 P-0399/CP	BURDITT, BOWLES, RADZIUS AND RUBERRY	NEW DOSAGE FORM	APPROVED FEB 16, 1988

ANDA SUITABILITY PETITIONS

PETITIONS APPROVED

DRUG NAME DOSAGE FORM; ROUTE	STRENGTH (CONTAINER SIZE)	DOCKET NUMBER	PETITIONER	REASON FOR PETITION	STATUS
HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE SOLUTION; ORAL	25MG/5ML 80MG/5ML	87 P-0399/CP	BURDITT, BOWLES, RADZIUS AND RUBERRY	NEW DOSAGE FORM	APPROVED FEB 16, 1988
HYDROCHLOROTHIAZIDE; TRIAMTERENE TABLET; ORAL	25MG 50MG	87 P-0335/CP	PAR PHARM	NEW DOSAGE FORM	APPROVED FEB 26, 1988
MEPERIDINE HYDROCHLORIDE INJECTABLE; INJECTION	10MG/ML (50ML/CONTAINER)	88 P-0008/CP	LYPHOMED	NEW STRENGTH	APPROVED APR 01, 1988
NIFEDIPINE CAPSULE; ORAL	10MG	88 P-0072/CP	MARTEC PHARMS	NEW DOSAGE FORM	APPROVED MAY 11, 1988
NIFEDIPINE CAPSULE; ORAL	20MG	88 P-0072/ CP0002	MARTEC PHARMS	NEW DOSAGE FORM	APPROVED MAY 11, 1988

ANDA SUITABILITY PETITIONS

PETITIONS APPROVED

DRUG NAME DOSAGE FORM; ROUTE	STRENGTH (CONTAINER SIZE)	DOCKET NUMBER	PETITIONER	REASON FOR PETITION	STATUS
PHENYTOIN SODIUM INJECTABLE; INJECTION	100MG/VIAL	87 P-0367/CP	LYPHOMED	NEW DOSAGE FORM	APPROVED FEB 16, 1988
PHENYTOIN SODIUM INJECTABLE; INJECTION	250MG/VIAL	87 P-0367/CP	LYPHOMED	NEW DOSAGE FORM	APPROVED FEB 16, 1988
THEOPHYLLINE CAPSULE, CONTROLLED RELEASE; ORAL	450MG	88 P-0119/CP	RIKER LABS	NEW STRENGTH	APPROVED MAY 11, 1988
THIOTEPA (WITH DILUENT) INJECTABLE; INJECTION	15MG/VIAL	87 P-0382/CP	LYPHOMED	NEW DOSAGE FORM	APPROVED MAY 12, 1988
VERAPAMIL HYDROCHLORIDE CAPSULE, CONTROLLED RELEASE; ORAL	120MG 240MG	87 P-0233/CP	SEARLE	NEW DOSAGE FORM NEW STRENGTH	APPROVED FEB 26, 1988

ANDA SUITABILITY PETITIONS

PETITIONS DENIED

DRUG NAME DOSAGE FORM; ROUTE	STRENGTH (CONTAINER SIZE)	DOCKET NUMBER	PETITIONER	REASON FOR PETITION	STATUS
BENZOYL METRONIDAZOLE SUSPENSION; ORAL	200MG/5ML	85 P-0258/CP	APKON LABS	NEW ESTER NEW INGREDIENT	DENIED MAR 19, 1986
BROMPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PHENYLPROPANOLAMINE HYDROCHLORIDE SYRUP; ORAL	2MG/5ML 2.5MG/5ML 12.5MG/5ML	85 P-0237/CP	MIKART	NEW COMBINATION	DENIED MAY 11, 1988
BROMDIPHENHYDRAMINE HYDROCHLORIDE; HYDROCODONE BITARTRATE SOLUTION; ORAL	12.5MG/5ML 2.5MG/5ML	85 P-0255/CP	MIKART	NEW COMBINATION	DENIED MAY 11, 1988
BROMDIPHENHYDRAMINE HYDROCHLORIDE; HYDROCODONE BITARTRATE SYRUP; ORAL	12.5MG/5ML 2.5MG/5ML	85 P-0255/CP	MIKART	NEW COMBINATION	DENIED MAY 11, 1988
CYCLOPHOSPHAMIDE INJECTABLE; INJECTION	100MG/ML (1ML/VIAL) (2ML/VIAL)	87 P-0283/CP	LYPHOMED	NEW DOSAGE FORM NEW STRENGTH	DENIED JAN 21, 1988
CYCLOPHOSPHAMIDE INJECTABLE; INJECTION	500MG/ML (1ML/VIAL) (2ML/VIAL) (4ML/VIAL)	87 P-0283/CP	LYPHOMED	NEW DOSAGE FORM NEW STRENGTH	DENIED JAN 21, 1988

ANDA SUITABILITY PETITIONS

PETITIONS APPROVED

DRUG NAME DOSAGE FORM; ROUTE	STRENGTH (CONTAINER SIZE)	DOCKET NUMBER	PETITIONER	REASON FOR PETITION	STATUS
PHENYTOIN SODIUM INJECTABLE; INJECTION	100MG/VIAL	87 P-0367/CP	LYPHOMED	NEW DOSAGE FORM	APPROVED FEB 16, 1988
PHENYTOIN SODIUM INJECTABLE; INJECTION	250MG/VIAL	87 P-0367/CP	LYPHOMED	NEW DOSAGE FORM	APPROVED FEB 16, 1988
THEOPHYLLINE CAPSULE, CONTROLLED RELEASE; ORAL	450MG	88 P-0119/CP	RIKER LABS	NEW STRENGTH	APPROVED MAY 11, 1988
THIOTEPA (WITH DILUENT) INJECTABLE; INJECTION	15MG/VIAL	87 P-0382/CP	LYPHOMED	NEW DOSAGE FORM	APPROVED MAY 12, 1988
VERAPAMIL HYDROCHLORIDE CAPSULE, CONTROLLED RELEASE; ORAL	120MG 240MG	87 P-0233/CP	SEARLE	NEW DOSAGE FORM NEW STRENGTH	APPROVED FEB 26, 1988

ANDA SUITABILITY PETITIONS

PETITIONS DENIED

DRUG NAME DOSAGE FORM; ROUTE	STRENGTH (CONTAINER SIZE)	DOCKET NUMBER	PETITIONER	REASON FOR PETITION	STATUS
BENZOYL METRONIDAZOLE SUSPENSION; ORAL	200MG/5ML	85 P-0258/CP	APKON LABS	NEW ESTER NEW INGREDIENT	DENIED MAR 19, 1986
BROMPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PHENYLPROPANOLAMINE HYDROCHLORIDE SYRUP; ORAL	2MG/5ML 2.5MG/5ML 12.5MG/5ML	85 P-0237/CP	MIKART	NEW COMBINATION	DENIED MAY 11, 1988
BROMDIPHENHYDRAMINE HYDROCHLORIDE; HYDROCODONE BITARTRATE SOLUTION; ORAL	12.5MG/5ML 2.5MG/5ML	85 P-0255/CP	MIKART	NEW COMBINATION	DENIED MAY 11, 1988
BROMDIPHENHYDRAMINE HYDROCHLORIDE; HYDROCODONE BITARTRATE SYRUP; ORAL	12.5MG/5ML 2.5MG/5ML	85 P-0255/CP	MIKART	NEW COMBINATION	DENIED MAY 11, 1988
CYCLOPHOSPHAMIDE INJECTABLE; INJECTION	100MG/ML (1ML/VIAL) (2ML/VIAL)	87 P-0283/CP	LYPHOMED	NEW DOSAGE FORM NEW STRENGTH	DENIED JAN 21, 1988
CYCLOPHOSPHAMIDE INJECTABLE; INJECTION	500MG/ML (1ML/VIAL) (2ML/VIAL) (4ML/VIAL)	87 P-0283/CP	LYPHOMED	NEW DOSAGE FORM NEW STRENGTH	DENIED JAN 21, 1988

ANDA SUITABILITY PETITIONS

PETITIONS APPROVED

DRUG NAME DOSAGE FORM; ROUTE	STRENGTH (CONTAINER SIZE)	DOCKET NUMBER	PETITIONER	REASON FOR PETITION	STATUS
DIPHENHYDRAMINE HYDROCHLORIDE CAPSULE, SUSTAINED RELEASE; ORAL	75MG	87 P-0355/CP	PARKE DAVIS	NEW DOSAGE FORM NEW STRENGTH	DENIED MAY 11, 1988
HYDROCODONE BITARTRATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE SYRUP; ORAL	1.66MG/5ML 5MG/5ML 6.25MG/5ML	85 P-0389/CP	UAD LABS	NEW COMBINATION	DENIED MAY 11, 1988
HYDROCODONE BITARTRATE; PROMETHAZINE HYDROCHLORIDE SYRUP; ORAL	2.5MG/5ML 6.25MG/5ML	85 P-0256/CP	MIKART	NEW COMBINATION	DENIED MAY 11, 1988
METOCLOPRAMIDE HYDROCHLORIDE INJECTABLE; INJECTION	1MG/ML (50ML/VIAL)	87 P-0090/CP	INTL MEDTN SYS	NEW STRENGTH	DENIED FEB 08, 1988

ANDA SUITABILITY PETITIONS

PETITIONS APPROVED

DRUG NAME DOSAGE FORM; ROUTE	STRENGTH (CONTAINER SIZE)	DOCKET NUMBER	PETITIONER	REASON FOR PETITION	STATUS
METOCLOPRAMIDE HYDROCHLORIDE INJECTABLE; INJECTION	1MG/ML (75ML/VIAL)	87 P-0090/CP	INTL MEDTN SYS	NEW STRENGTH	DENIED FEB 08, 1988
METOCLOPRAMIDE HYDROCHLORIDE INJECTABLE; INJECTION	1MG/ML (100ML/VIAL)	87 P-0090/CP	INTL MEDTN SYS	NEW STRENGTH	DENIED FEB 08, 1988

EXCLUSIVITY TERMS

DUE TO SPACE LIMITATIONS IN THE EXCLUSIVITY COLUMN, ABBREVIATIONS AND REFERENCES HAVE BEEN DEVELOPED. REFER BACK TO THE APPROVED DRUG PRODUCTS WITH THERAPEUTIC EQUIVALENCE EVALUATIONS, 8TH EDITION FOR A FULL LISTING OF EXCLUSIVITY TERMS (ABBREVIATIONS, NEW DOSING SCHEDULE, NEW INDICATIONS AND PATENT USE CODES). ONLY NEW CODES WILL BE ADDED TO THE CUMULATIVE SUPPLEMENT.

REFERENCES

NEW DOSING SCHEDULE

D-12	BEDTIME DOSING OF 800MG FOR TREATMENT
D-12	BEDTIME DOSING OF 800MG FOR TREATMENT OF ACTIVE DUODENAL ULCER
D-13	INCREASED MAXIMUM DAILY DOSAGE RECOMMENDATION
D-14	BEDTIME DOSING OF 800MG FOR TREATMENT OF ACTIVE BENIGN GASTRIC ULCER

NEW INDICATION

I-72	PHOTOPHERESIS IN THE PALLIATIVE TREATMENT OF SKIN MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PERSONS NOT RESPONSIVE TO OTHER TREATMENT
I-73	FOLLICULAR STIMULATION IN VITRO FERTILIZATION

EXCLUSIVITY TERMS

REFERENCES

PATENT USE CODE

U-26	METHOD OF TREATING ANIMALS SUFFERING FROM AN APPETITE DISORDER
U-27	METHOD OF BLOCKING THE UPTAKE OF MONOAMINES BY BRAIN NEURONS IN ANIMALS
U-28	METHOD FOR IMPROVING MEMORY IN MAMMALS
U-29	METHOD FOR TREATING AMNESIA
U-30	METHOD OF POTENTIATING CODEINE ANALGESIA IN MAMMALS
U-31	USE IN LUNG SCANNING PROCEDURES
U-32	TREATMENT OF VENTRICULAR AND SUPRAVENTRICULAR ARRHYTHMIAS
U-33	METHOD FOR INHIBITING GASTRIC SECRETION IN MAMMALS

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA

46

APPL/PROD	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
>ADD> >ADD>	19489 001 ALBUTEROL SULFATE; VENTOLIN ROTACAPS	3705233	DEC 05, 1989		NDF	MAY 04, 1991
	17920 002 CIMETIDINE; TAGAMET	3644353	FEB 22, 1989		D-14	MAR 31, 1991
	17920 003 CIMETIDINE; TAGAMET	4024271	MAY 17, 1994		D-12	APR 30, 1989
	17920 004 CIMETIDINE; TAGAMET	3950333	APR 13, 1993		D-14	MAR 31, 1991
	17920 005 CIMETIDINE; TAGAMET	4024271	MAY 17, 1994		D-12	APR 30, 1989
	17924 001 CIMETIDINE HYDROCHLORIDE; TAGAMET	3950333	APR 13, 1993		D-14	MAR 31, 1991
	19309 001 ENALAPRILAT; VASOTEC	4024271	MAY 17, 1994		D-12	APR 30, 1989
	18981 002 ENCAINIDE HYDROCHLORIDE; ENKAID	3950333	APR 13, 1993		D-14	MAR 31, 1991
	18981 003 ENCAINIDE HYDROCHLORIDE; ENKAID	4024271	MAY 17, 1994		D-12	APR 30, 1989
	18981 004 ENCAINIDE HYDROCHLORIDE; ENKAID	3950333	APR 13, 1993		D-14	MAR 31, 1991
	19452 001 FLUOCINOLONE ACETONIDE; DERMA-SMOOTHIE/FS	4024271	MAY 17, 1994		D-12	APR 30, 1989
	18936 001 FLUOXETINE HYDROCHLORIDE; PROZAC	4024271	MAY 17, 1994		NDF	FEB 09, 1991
		RE30811	FEB 22, 2000	U-32		
		RE30811	DEC 20, 1996	U-32		
		RE30811	DEC 20, 1996	U-32		
		4683235	JUL 28, 2004	U-30		
		4647591	MAR 03, 2004	U-28		
		4647591	MAR 03, 2004	U-29		
		4626549	DEC 02, 2003	U-26		
		4626549	DEC 02, 2003	U-27		
		4194009	APR 19, 1994			
		3793457	FEB 19, 1993			
		3655663	APR 11, 1989			
		4021481	MAY 03, 1994			
		3681500	AUG 01, 1991			
		4374829	FEB 22, 2000			
		4374829	FEB 22, 2000			
		4374829	FEB 22, 2000			
		3714159	JAN 30, 1990			
		4087545	MAY 02, 1997	U-7		
		4282251	AUG 04, 1998			
		4382090	MAY 03, 2000	U-33		
		4375547	MAR 01, 2000			
		4382090	MAY 03, 2000	U-33		
		4375547	MAR 01, 2000			
		4021481	MAY 03, 1994			
		3681500	AUG 01, 1991			
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		4382090	MAY 03, 2000	U-33		
		4375547	MAR 01, 2000			
		4382090	MAY 03, 2000	U-33		
		4375547	MAR 01, 2000			
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		4021481	MAY 03, 1994			
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		3714159	JAN 30, 1990			
		4087545	MAY 02, 1997	U-7		
		4282251	AUG 04, 1998			
		4382090	MAY 03, 2000	U-33		
		4375547	MAR 01, 2000			
		4382090	MAY 03, 2000	U-33		
		4375547	MAR 01, 2000			
		4021481	MAY 03, 1994			
		3681500	AUG 01, 1991			
		4374829	FEB 22, 2000			
		4374829	FEB 22, 2000			
		4374829	FEB 22, 2000			
		3714159	JAN 30, 1990			
		4087545	MAY 02, 1997	U-7		
		4282251	AUG 04, 1998			
		4382090	MAY 03, 2000	U-33		
		4375547	MAR 01, 2000			
		4382090	MAY 03, 2000	U-33		
		4375547	MAR 01, 2000			
		4021481	MAY 03, 1994			
		3681500	AUG 01, 1991			
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		4087545	MAY 02, 1997	U-7		
		4282251	AUG 04, 1998			
		4382090	MAY 03, 2000	U-33		
		4375547	MAR 01, 2000			
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		4375547	MAR 01, 2000			
		4021481	MAY 03, 1994			
		3681500	AUG 01, 1991			
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		4087545	MAY 02, 1997	U-7		
		4282251	AUG 04, 1998			
		4382090	MAY 03, 2000	U-33		
		4375547	MAR 01, 2000			
		4382090	MAY 03, 2000	U-33		
		4375547	MAR 01, 2000			
		4021481	MAY 03, 1994			
		3681500	AUG 01, 1991			
		4374829	FEB 22, 2000			
		4374829	FEB 22, 2000			
		4374829	FEB 22, 2000			
		3714159	JAN 30, 1990			
		4087545	MAY 02, 1997	U-7		
		4282251	AUG 04, 1998			
		4382090	MAY 03, 2000	U-33		
		4375547	MAR 01, 2000			
		4382090	MAY 03, 2000	U-33		
		4375547	MAR 01, 2000			
		4021481	MAY 03, 1994			

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA

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APPL/PROD	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
19009 001	PIRBUTEROL ACETATE; EXIREL	4175128	NOV 20, 1996			
		3786160	JAN 15, 1993			
		3700681	OCT 24, 1989		NCE	DEC 30, 1991
17881 001	TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT; TECHNETIUM TC 99M	3863004	MAR 18, 1992	U-31		
19057 001	TERAZOSIN HYDROCHLORIDE; HYTRIN	4251532	JAN 28, 1992	U-5	NCE	AUG 07, 1992
		4112097	FEB 17, 1998	U-5		
		4026894	SEP 05, 1995			
19057 002	TERAZOSIN HYDROCHLORIDE; HYTRIN	4026894	MAY 31, 1994	U-5	NCE	AUG 07, 1992
		4251532	FEB 17, 1998	U-5		
		4112097	SEP 05, 1995			
19057 003	TERAZOSIN HYDROCHLORIDE; HYTRIN	4026894	MAY 31, 1994	U-5	NCE	AUG 07, 1992
		4251532	FEB 17, 1998	U-5		
		4112097	SEP 05, 1995			
19057 004	TERAZOSIN HYDROCHLORIDE; HYTRIN	4026894	MAY 31, 1994	U-5	NCE	AUG 07, 1992
		4251532	FEB 17, 1998	U-5		
		4112097	SEP 05, 1995			
19641 001	TERCONAZOLE; TERAZOL 3	4026894	MAY 31, 1994	U-5	NCE	AUG 07, 1992
		4358449	NOV 09, 1999			
18207 003	TRAZODONE HYDROCHLORIDE; DESYREL	4258027	MAR 24, 1998		NCE	DEC 31, 1992
		4215104	JUL 29, 1997	NDF		MAY 24, 1991
19415 002	UROFOLLITROPIN; METRODIN				I-73	MAR 01, 1991
18817 003	VERAPAMIL HYDROCHLORIDE; CALAN				I-50	DEC 16, 1989
					I-51	DEC 16, 1989
18817 004	VERAPAMIL HYDROCHLORIDE; CALAN				I-50	DEC 16, 1989
					I-51	DEC 16, 1989



SUBSCRIPTION FORM
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- ☐ Check/money order enclosed for \$ _____
(Make check or money order payable to Superintendent of Documents)

AUTHORIZING
SIGNATURE:

DATE:

DESCRIPTION	QUANTITY	UNIT PRICE	TOTAL PRICE
The 8th Edition is published in March 1988. Subscription includes the Approved Drug Products publication and monthly Cumulative Supplements.			
DOMESTIC (Stock No. 917-001-00000-6)		@ \$ 79.00	\$
FOREIGN (Stock No. 917-001-00000-6)		@ \$ 98.75	\$
ENTER TOTAL			\$