PERFORMRx℠

Drug Information Update

July 2023

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NEWLY AVAILABLE GENERICS

Generic Name/ Dosage Form	Brand Name	Manufacturer	Indication
Meclizine 50 mg oral tablet	Antivert	Rising Pharmaceuticals	• Prevention and treatment of nausea, vomiting, or dizziness associated with motion sickness • Treatment of vertigo associated with diseases affecting the vestibular system
Plerixafor 24 mg/1.2 ml subcutaneous vial	Mozobil	Fresenius Kabi	• To be used in combination with granulocyte-colony stimulating factor (G-CSF) to mobilize hematopoietic stem cells (HSCs) to the peripheral blood for collection and subsequent autologous transplantation in patients with non-Hodgkin's lymphoma or multiple myeloma

NEW DRUG ENTITIES/STRENGTHS/COMBINATIONS

Drug Name	Generic Name	Description
Zejula 100, 200, 300 mg oral tablet	niraparib tosylate	New tablet formulation (caps only available 100 mg). No new indications or dosing.
Hulio 40 mg/0.8 ml subcutaneous auto-injector	adalimumab-fkjp	New drug launch, biosimilar adalimumab.
Hulio 20 mg/0.4 ml, 40 mg/0.8 ml subcutaneous prefilled syringe	adalimumab-fkjp	New drug launch, biosimilar adalimumab.
Adalimumab-fkjp 20 mg/0.4 ml, 40 mg/0.8 ml subcutaneous prefilled syringe	adalimumab-fkjp	New drug launch, biosimilar adalimumab. Unbranded Hulio with lower pricing.
Idacio 40 mg/0.8 ml subcutaneous auto-injector	adalimumab-aacf	New drug launch, biosimilar adalimumab.
Idacio 40 mg/0.8 ml subcutaneous prefilled syringe	adalimumab-aacf	New drug launch, biosimilar adalimumab.
Rezzayo 200 mg intravenous solution	rezafungin acetate	Once-weekly IV treatment of candidemia and invasive candidiasis in adults with limited or no alternative treatment options.
Cyltezo 40 mg/0.8 ml subcutaneous prefilled syringe	adalimumab-adbm	Humira biosimilar. Only available in low concentration strengths.
Cyltezo 10 mg/0.2 ml, 20 mg/0.4 ml subcutaneous prefilled syringe	adalimumab-adbm	Humira biosimilar. Only available in low concentration strengths.
Rystiggo 280 mg/2 ml subcutaneous Solution	rozanolixizumab-noli	Indicated for the treatment of generalized myasthenia gravis (gMG) in adult patients who are anti-acetylcholine receptor (AChR) or anti-muscle-specific tyrosine kinase (MuSK) antibody positive.
Suflave 178.7 g oral solution	peg 3350/sodium sulfate/potassium chloride/magnesium/sodium chloride	New bowel prep treatment option that is touted to be better tasting than other pre-colonoscopy treatments. Meant to improve palatability for patients with a taste that's similar to a lemon-lime sports drink.
Elevidys intravenous kit	delandistrogene moxeparvovec-rokl	First gene therapy for the treatment of ambulatory pediatric patients 4 through 5 years of age with Duchenne muscular dystrophy (DMD) with a confirmed mutation in the DMD gene.

Drug Name	Generic Name	Description
		Indication is approved under accelerated approval based on expression of micro-dystrophin observed in patients treated with Elevidys.
Adalimumab-adaz 40 mg/0.4 ml subcutaneous auto-injector	adalimumab-adaz	Humira biosimilar. Unbranded version of Hyrimoz. High concentration strength.
Adalimumab-adaz 40 mg/0.4 ml subcutaneous prefilled syringe	adalimumab-adaz	Humira biosimilar. Unbranded version of Hyrimoz. High concentration strength.
Hyrimoz 40 mg/0.4 ml, 80 mg/0.8 ml subcutaneous auto-injector	adalimumab-adaz	Humira biosimilar. High concentration strength.
Hyrimoz 10 mg/0.1 ml, 20 mg/0.2 ml, 40 mg/0.4 ml, 80 mg/0.8 ml subcutaneous prefilled syringe	adalimumab-adaz	Humira biosimilar. High concentration strength.
Yuflyma 40 mg/0.4 ml subcutaneous auto-injector	adalimumab-aaty	Humira biosimilar. Only available in high concentration strength.
Yusimry 40 mg/0.8 ml subcutaneous auto-injector	adalimumab-aqvh	Humira biosimilar. Only available in low concentration strength.
Hadlima 40 mg/0.4 ml, 40 mg/0.8 ml subcutaneous prefilled syringe	adalimumab-bwwd	Humira biosimilar. Available in both high and low concentration strengths.
Hadlima PushTouch 40 mg/0.4 ml, 40 mg/0.8 ml subcutaneous auto-injector	adalimumab-bwwd	Humira biosimilar. Available in both high and low concentration strengths.
Litfulo 50 mg oral capsule	ritlecitinib	New JAK inhibitor indicated for the treatment of individuals 12 years of age and older with severe alopecia areata. Joins Olumiant as the only FDA-approved treatments for alopciea areata, however, unlike Olumiant this can be used adolescents (12+).
Austedo XR 6 mg/12 mg/24 mg extended release oral tablet pack	deutetrabenazine	New titration pack for Austedo XR
Abrysvo 120 mcg/0.5 ml intramuscular solution	respiratory syncytial virus vaccine	New RSV vaccine from Pfizer for the prevention of lower respiratory tract disease (LRTD) caused by respiratory syncytial virus RSV in individuals 60 years and older. Will compete with Arexvy.
Adstiladrin 300,000,000,000 vp/ml intravesical suspension	adstiladrin intravesical	Novel gene therapy for the treatment of adult patients with high-risk, Bacillus Calmette-Guérin (BCG)-unresponsive non-muscle invasive bladder cancer (NMIBC) with carcinoma in situ (CIS)

Drug Name	Generic Name	Description
		with or without papillary tumors. It is administered every 3 months and target's the patient's own bladder wall cells to enhance natural defenses against cancer.
Arexvy 120 mcg/0.5m intramuscular suspension	respiratory syncytial virus vaccine, adjuvanted	New RSV vaccine from GSK for the prevention of lower respiratory tract disease (LRTD) caused by respiratory syncytial virus (RSV) in individuals 60 years of age and older. Will compete with Abrysvo.
Roctavian 20,000,000,000,000 vg/ml intravenous suspension	valoctocogene roxaparvovec-rvox	First gene therapy for the treatment of adults with severe hemophilia A (congenital factor VIII (FVIII) deficiency with FVIII activity < 1 IU/dL) without antibodies to adeno-associated virus serotype 5 (AAV5) detected by an FDA-approved test.
Brenzavvy oral tablet 20 mg	bexagliflozin	New SGLT2 inhbitor for type 2 diabetes.

NEW INDICATIONS (EXISTING DRUGS)

†**Bolded** items reflect newly approved indication; ~~strikethrough~~ of removed indication/age limit/etc.

Brand Name	Generic Name/ Dosage Form	Manufacturer	Newly Approved Indication†
Leqembi	lecanemab-irmb 200 mg/2 mL, 500 mg/5 ml intravenous vials	Eisai	Treatment of Alzheimer's disease. Treatment with Leqembi should be initiated in patients with mild cognitive impairment or mild dementia stage of disease, the population in which treatment was initiated in clinical trials. ~~There are no safety or effectiveness data on initiating treatment at earlier or later stages of the disease than were studied.~~ ~~This indication is approved under accelerated approval based on reduction in amyloid beta plaques observed in patients treated with LEQEMBI. Continued approval for this indication may be contingent upon verification of clinical benefit in a confirmatory trial~~.
Liletta	levonorgestrel 52 mg intrauterine system	AbbVie	• Prevention of pregnancy for up to 8 years • **Treatment of heavy menstrual bleeding for up to 5 years in patients who choose intrauterine contraception as their method of contraception**
Leqvio	inclisiran 284 mg/1.5 ml subcutaneous syringe	Novartis	Adjunct to diet and ~~maximally tolerated~~ statin therapy for the treatment of adults with **primary hyperlipidemia, including** heterozygous familial hypercholesterolemia (HeFH) ~~or clinical atherosclerotic cardiovascular disease (ASCVD), who require additional lowering of~~ **to reduce** low-density lipoprotein cholesterol (LDL-C)

RECALLS

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Artificial Tears (Carboxymethylcellulose Sodium) Lubricant Eye Drops, 10 MG in 1 ml, 1/2 fl oz (15 ml) bottle, Distributed by. EzriCare, LLC, Lakewood, NJ, NDC 79503-0101-15.	Class I	Drugs	Lot #: PCMI005, Exp. date AUG-2024; PCMJ001, PCMJ002, PCMJ004, PCMJ005, PCMJ006, PCMJ008, PCMJ009, PCMJ010, PCMJ011, PCMJ012, PCMJ013, PCMJ014, PCMJ015, PCMJ016, Exp. date MAR-2025	Non-Sterility: FDA analysis found unopened products to have bacterial contamination.	Global Pharma Healthcare Private Limited
Delsam Pharma's ARTIFICIAL TEARS (Carboxymethylcellulose Sodium) Lubricant Eye Drops, 10 MG in 1 ml, 1%, 1/2 fl oz (15 ml) bottle, Distributed By: Delsam Pharma Llc, Bronx, New York 10467, NDC 72570 121 15.	Class I	Drugs	Lot #: PCMH001, PCMH002, PCMH005, PCMH007, Exp. date NOV-2023	Non-Sterility: FDA analysis found unopened products to have bacterial contamination.	Global Pharma Healthcare Private Limited
Albuterol Sulfate Inhalation Aerosol, 90 mcg, 200 Metered Inhalation, net content 6.7 g canister packaged in a box, Rx only, Manufactured by: Cipla Ltd, Indore SE Z, Pithampur, India, Manufactured for: Cipla USA, Inc., 10 Independence Boulevard, Suite 300, Warren, NJ, 07059, NDC 69097-142-60	Class I	Drugs	Lot # IB20045, IB20055, IB20056, IB20057, IB20059, IB20072, Exp Nov. 30, 2023	Defective container: empty inhaler and leakage observed through the inhaler valve due to partially missing bottom seat (gasket).	Cipla USA, Inc.
Amphotericin B Liposome for Injection, 50mg/vial, Rx only, Distributed by Sun Pharmaceutical Industries, Inc. Cranbury, NJ 08512, Manufactured by Sun Pharmaceutical Medicare Limited, Baska	Class II	Drugs	Lot # BAD0089A, Exp 01/2025; BAD0330A, Exp 07/2025	Subpotent drug	SUN PHARMACEUTICAL INDUSTRIES INC

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Ujeti Road, Ujeti, Halol-389350, Gujarat, India, NDC 62756-0233-01					
Kanjo Arnica Cream, 2 oz. gel & cream NDC# 80551-201-02 Mfg. For: Acutens,Inc. Sheridan, WY 82801	Class II	Drugs	Lot # 2446, exp. date Mar-24	cGMP deviations	Homeocare Laboratories, Inc.
Arnica Cream, 2 oz. NDC# 61727-114-02 Mfg. For: Brazmedics, LLC New York, NY 10021	Class II	Drugs	Lot # 2446, exp. Mar-24	cGMP deviations	Homeocare Laboratories, Inc.
Bebelyn Colics, 0.50 fl. oz. distributed by Pharmadel LLC New Castle, DE 19720 NDC# 55758-036-01	Class II	Drugs	Lot # 2414, exp. date Sep-24	cGMP deviations	Homeocare Laboratories, Inc.
Babelyn Diarrhea, 0.50 fl. oz. distributed by Pharadel LLC NDC# 55758-035-01	Class II	Drugs	Lot # 2413, exp. date Sep-24	cGMP deviations	Homeocare Laboratories, Inc.
DoloEar, 0.50 fl. oz. distributed by Pharadel LLC NDC# 55758-035-15	Class II	Drugs	Lot # 2423, exp. date Nov-24; 2445, exp. date Apr-25	cGMP deviations	Homeocare Laboratories, Inc.
SnoreStop Naso Spray Bulk, liquid distributed by Green Pharma NDC# 61152-199-99	Class II	Drugs	Lot # 2436, exp. date: unknown	cGMP deviations	Homeocare Laboratories, Inc.
StellaLife Peppermint Dental Rinse, 16 oz. bottles NDC# 69685-103-16	Class II	Drugs	Lot # 2357B,2333, exp. date Sep-2; 2367C, 2367D, exp. date Nov-23; 2367E, 2367F, exp. date Jan-24; 2395A, 2406, exp. date Feb-2; 2395B, exp. date Mar-24; 2395C, exp. date Apr-24	cGMP deviations	Homeocare Laboratories, Inc.

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
StellaLife Coconut Dental Rinse, 16 oz. bottles NDC# 69685-113-16	Class II	Drugs	Lot # 2396, exp. date Apr-24	cGMP deviations	Homeocare Laboratories, Inc.
StellaLife Dental Spray, 1 fl. oz. NDC# 69685-101-02	Class II	Drugs	Lot # 2335, exp. date Sep-23; 2364, exp. date Nov-23; 2408, exp. date Feb-24	cGMP deviations	Homeocare Laboratories, Inc.
StellaLife Dental Gel, 1 fl. oz. NDC# 69685-102-02	Class II	Drugs	Lot #2334, exp. date Sep-23; 2363, exp. date Nov-23; 2365, exp. date Jan-24	cGMP deviations	Homeocare Laboratories, Inc.
StellaLife Dental Gel, 1 fl. oz. NDC# 69685-112-01	Class II	Drugs	Lot #2407, exp. date Feb-24; 2409A, exp. date Apr-24; 2409B, exp. date May-24	cGMP deviations	Homeocare Laboratories, Inc.
Endometrin (progesterone) Vaginal Insert 100mg, packaged in a carton of 21 vaginal inserts with 21 disposable vaginal applicators, Manufactured for: Ferring Pharmaceuticals Inc., Parsippany, NJ	Class II	Drugs	AA200A, AA201A, Exp. 12/2025	cGMP deviations: potential for Microbial Contamination of Non-Sterile Products	Ferring Pharmaceuticals Inc
Tramadol Hydrochloride Tablets, USP 50 mg, Unit Dose 100-Count Tablets Rx only, Manufactured by: Sun Pharmaceutical Industries Ltd. Survey No. 259/15, Dadra-396 191, (U.T. of D & NH), India. Distributed by: Sun Pharmaceutical Industries, Inc. Cranbury, NJ 08512, Distributed by: Major Pharmaceuticals Livonia, MI 48152 USA, NDC 0904-7179-61	Class II	Drugs	Lot #: M04343, Exp. 04/2024	Packaging defect: blister packaging inadequately sealed.	The Harvard Drug Group
Losartan Potassium Tablets, USP, 25 mg, 1000 film coated tablets per bottle, Rx Only, Manufactured by: Vivimed Life Sciences	Class II	Drugs	Lot#: 7901903A, exp. date 04/2024	Presence of Foreign Substance: Presence of a small piece of blue plastic embedded in the tablet.	Strides Pharma Inc.

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Private Limited, Plot No. 101, 102, 107 & 108, SIDCO Pharmaceutical Complex, Alathur, Kanchipuram- 603 110, Tamilnadu, India, NDC 64380-933-08					
Tizanidine Tablets, USP, 4 mg, 100 Tablets (10 tablets x 10 unit dose blister packs) per carton, Rx Only, Packaged and Distributed by: American Health Packaging, Columbus, Ohio 43217, NDC 68084-645-01 (carton), barcode (01) 003 68084 645 11 2.	Class II	Drugs	Lot 1004835, Exp 7/31/2023	Failed Dissolution Specifications: this repackaged product was recalled by the manufacturer, Dr. Reddy's Laboratories, Inc., due to out of specification results for dissolution.	Amerisource Health Services LLC
Tizanidine Hydrochloride Tablet 4mg, packaged in a) 20 count-bottles (NDC 68788-7781-2), b)30-count bottles (NDC: 68788-7781-3), c) 60-count bottles, (NDC: 68788-7781-6), d) 90-count bottles (NDC: 68788-7781-9), e) 120-count bottles (NDC: 68788-7781-8), Rx only, Mfg: Dr. Reddy's Laboratories Limited.	Class II	Drugs	Lot#: a) H1621S, Exp: 12/31/2023; b) H2321C, Exp: 12/31/2023; c) H0421B, Exp: 12/31/2023; d) H1721E, H1921T, H3121M, Exp: 12/31/2023, e) H2021G, Exp: 12/31/2023	Failed Stability Specifications	Preferred Pharmaceuticals, Inc.
Artificial Tears (Carboxymethylcellulose Sodium) Lubricant Eye Drops, 10 MG in 1 ml, 1/2 fl oz (15 ml) bottle, Distributed by. EzriCare, LLC, Lakewood, NJ, NDC 79503-0101-15.	Class II	Drugs	Lot #: PCMH009, PCMH010, PCMH011, PCMH012, PCMH013, PCMH014, PCMH015, PCMH016, exp. date Nov-2023; PCMI001, PCMI002, PCMI003, PCMI004, PCMI006, PCMI 07, PCMI 008, Exp. date AUG-2024; PCMJ003, PCMJ007, Exp. date MAR-2025	CGMP Deviations: All other lots of eye drops are being recalled due to CGMP Deviations because they were manufactured in the same facility under the same conditions as the lots found to be contaminated.	Global Pharma Healthcare Private Limited

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Delsam Pharma's ARTIFICIAL TEARS (Carboxymethylcellulose Sodium) Lubricant Eye Drops, 10 MG in 1 ml, 1%, 1/2 fl oz (15 ml) bottle, Distributed By: Delsam Pharma Llc, Bronx, New York 10467, NDC 72570 121 15.	Class II	Drugs	Lot #: PCMH003, PCMH004, PCMH006, PCMH008, Exp. date NOV-2023	CGMP Deviations: All other lots of eye drops are being recalled due to CGMP Deviations because they were manufactured in the same facility under the same conditions as the lots found to be contaminated.	Global Pharma Healthcare Private Limited
Allergy, Fexofenadine Hydrochloride Tablets, 180 mg, packaged in a) 30-count cartons (NDC 72288-847-39), and b)150-count cartons (NDC 72288-847-47 and 72288-847-37), Distributed By: Amazon.com Services LLC., 410 Terry Avenue N., Seattle, WA 98109.	Class II	Drugs	Lot # a) 2GR0329, Exp. date 04/24 Lot # b) 2DR0472, Exp. date 02/23; 2MR0417, Exp. date 07/24	Failed Impurities/Degradation Specifications	L. Perrigo Company
Non-Drowsy Allergy Relief, Fexofenadine Hydrochloride Tablets, 180 mg, 150-count cartons, Distributed By: BJ's Wholesale Club, 25 Research Drive, Westborough, MA 01581. NDC 68391-847-47	Class II	Drugs	Lot #:2DV1863, 2HV2698, Exp. date 12/23; 2GV1583, Exp. date 02/24; 2GV1950, 2HV2697, Exp. Date 01/24.	Failed Impurities/Degradation Specifications	L. Perrigo Company
Allergy Relief, Fexofenadine Hydrochloride Tablets, 180 mg, 15-count cartons, Distributed By: Adusa Distribution, LLC, Salisbury, NC 28147. NDC 72476-847-22	Class II	Drugs	Lot #: 2JE2185, Exp. date 01/24	Failed Impurities/Degradation Specifications	L. Perrigo Company
Allergy Relief, Fexofenadine Hydrochloride Tablets, 180 mg, 30-count cartons, Distributed By: CVS Pharmacy, Inc., One CVS Drive, Woonsocket, RI 02895. NDC 69842-0914-39	Class II	Drugs	Lot #: 2DV1925, Exp. date 06/23	Failed Impurities/Degradation Specifications	L. Perrigo Company
aller-ease, Fexofenadine Hydrochloride Tablets, 180 mg, 30-count cartons, Packaged	Class II	Drugs	Lot #: 2ER0285,Exp. date 01/24	Failed Impurities/Degradation Specifications	L. Perrigo Company

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
For: Your Military Exchanges, By: Perrigo Company, Allegan, MI USA 49010. NDC 55301-847-39					
Allergy ALLER-FEX, Fexofenadine Hydrochloride Tablets, 180 mg, 180-count cartons, Packaged by: Perrigo, 515 Eastern Ave., Allegan, MI 49010 USA. NDC 63981-847-48	Class II	Drugs	Lot #: 2DV1487, 2DV1870, 2DV1871, 2DV1873, 2EV1676, 2GV2132, 2HV2679, Exp. date 12/23; 2DV2000, exp. date 11/23; 2LV1573, 2MV1314, Exp. date 04/24; 2FV1764, 2FV1765 2FV1766, 2GV1579, 2GV1937, 2GV1941 , 2GV1942, 2GV2157, 2HV1886, 2HV1997, 2HV2019, 2HV2047, Exp. datev02/24; 2EV1666, 2EV1667, 2EV1668, 2EV1670, 2EV1671, 2EV1672, 2EV1674, 2EV1677, 2EV1678, 2HV2017, Exp. date 01/24.	Failed Impurities/Degradation Specifications	L. Perrigo Company
allergy relief, Fexofenadine Hydrochloride Tablets, 180 mg, packaged in a)15-count cartons (NDC 79481-0847-0), b)30-count cartons (NDC 79481-0847-1), and c) 45-count cartons (NDC 79481-0847-2), Distributed by: MEIJER DISTRIBUTION, INC., GRAND RAPIDS, MI 49544.	Class II	Drugs	Lot #: a) 2JE1882, Exp. date 01/24 b) 2FV1918, 2ER0411, Exp. date 01/24; 2GV1902, Exp. date 04/24; 2CR0652, 2DR0465, Exp. date 12/23; 2GR0329, Exp. date 04/24 c)2CR0653, 2DR0466, Exp. date 12/23;	Failed Impurities/Degradation Specifications	L. Perrigo Company

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			2ER0287, 2ER0412, Exp. date 01/24		
Fexofenadine Hydrochloride Tablets, 180 mg, 100-count cartons, Distributed by: Perrigo, Allegan, MI 49010. NDC 45802-847-78	Class II	Drugs	Lot #: 2DR0351, Exp. date 12/23	Failed Impurities/Degradation Specifications	L. Perrigo Company
Fexofenadine Hydrochloride Tablets, 180 mg, packaged in a) 30-count cartons (NDC 0113-0847-39), and b) 45-count cartons (NDC 0113-0847-95), Distributed by: Perrigo, Allegan, MI 49010.	Class II	Drugs	Lot #: a) 2DV1869, 2EV1613, 2EV1614, Exp. date 12/23; 2EV1820, 2FV1943, Exp. date 01/24; 2GV1893, Exp. date 04/24. Lot #: b) 2CR0653, 2DR0466, Exp. date 12/23, 2ER2087, Exp. date 01/24	Failed Impurities/Degradation Specifications	L. Perrigo Company
allergyrelief, Fexofenadine Hydrochloride Tablets, 180 mg, packaged in a)30-count cartons (NDC 56062-847-39); and b) 45-count cartons (NDC 56062-847-95), Distributed by: Publix Super Markets, Inc., 3300 Publix Corporate Parkway, Lakeland, FL 33811.	Class II	Drugs	Lot #: a) 2CR0652, 2DR0464, 2DR0465,Exp. Date 12/23; 2ER0285, Exp. Date 01/24. b) 2DR0466, Exp. Date 12/23; 2ER0412, Exp. Date 01/24; 2GR0330, Exp. Date 04/24	Failed Impurities/Degradation Specifications	L. Perrigo Company
allergy relief, Fexofenadine Hydrochloride Tablets, 180 mg, packaged in a)15-count cartons (NDC 11673-617-22), b)30-count cartons (NDC 11673-617-39), c)70-count cartons (NDC 11673-617-01), and d)150-count cartons (11673-617-47), Distributed by: Target Corporation, Minneapolis, MN 55403.	Class II	Drugs	Lot #: a) 2HE2032, 2JE1882, 2JE2185 Exp. Date 01/24; b) 2DR0464, 2ER0410, Exp. Date 12/23; 2ER0286, 2ER0411, Exp. Date 01/24 c) 2DR0467, 2DR0468, 2DR0469, 2ER0288, Exp. Date 12/23 d) 2ER0414, Exp. Date	Failed Impurities/Degradation Specifications	L. Perrigo Company

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			01/24, 2GR0333, Exp. Date 02/24		
Picnic, Fexofenadine Hydrochloride Tablets, 180 mg, Antihistamine, 90-count cartons, Distributed by: Thirty Madison, Inc., New York, NY 10001. NDC 45 tablets: 80159-112-03	Class II	Drugs	Lot#: 2DR0471, Exp. Date 12/23	Failed Impurities/Degradation Specifications	L. Perrigo Company
Allergy Relief, Fexofenadine Hydrochloride Tablets, 180 mg, packaged in a) 30-count cartons (NDC 36800-691-39), b)45-count cartons (NDC 36800-691-95), and 90-count cartons (NDC 36800-691-75), Distributed by: Topco Associates LLC, Elk Grove Village, IL 60007.	Class II	Drugs	Lot #: a) 2FV1948, 2ER0285, 2ER0411, Exp. Date 01/24 b)2CR0653, Exp. Date 12/23 c)2GR0331, Exp. Date 04/24	Failed Impurities/Degradation Specifications	L. Perrigo Company
Original Eye Drops; Redness Reliever; (Tetrahydrozoline HCl), 0.05%, 0.5 FL OZ (15mL); distributed by a) Original Formula Eye Drops, DG health, DISTRIBUTED BY OLD EAST MAIN CO, 100 MISSION RIDGE, GOODLETTSVILLE, TN 37072, UPC 0 95072 00556 5; b) Publix, DISTRIBUTED BY PUBLIX SUPER MARKETS, INC, 3300 PUBLIX CORPORATE PARKWAY, LAKELAND, FL 33811, UPC 0 41415 01076 5; c) sterile eye drops, Original Formula, sunmark, Distributed by McKesson, 6555 State Highway 161, Irving, TX 75039, UPC 0 10939 16733 0, NDC 49348-037-29; d) TopCare health, DISTRIBUTED BY TOPCO ASSOCIATES LLC, ELKGROVE VILLAGE, IL 60007, UPC 0	Class II	Drugs	Lot #: a) RG21F01, Exp: 6/30/2023; b) RG21F01, Exp: 6/30/2023; c) RG21F01, Exp: 6/30/2023; d) RG21F01, RG21F02, Exp 6/30/2023; e) RG21F02, RG21F03, Exp 6/30/2023; f) RG21F02, Exp 6/30/2023; g) RG21F02, Exp 6/30/2023; h) RG21F02, Exp 6/30/2023; i) RG21F02, Exp 6/30/2023; j) RG21F02, Exp 6/30/2023; k) RG21F02, Exp 6/30/2023; l) RG21F02, Exp	CGMP Deviations: good manufacturing deficiencies related to a lack of documentation of the fill line.	K.C. Pharmaceuticals, Inc

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
36800 03639 0; e) Eye Drops, Original Formula, GoodSense, Distributed By: Geiss, Destin & Dunn, Inc., Peachtree City, GA 30269, UPC 1 80410 00015 6, NDC 50804-141-01; f) sterile eye drops, Circle K, Product manufactured for: Lil' Drug Store Products, Inc., 9300 Earhart Lane SW, Cedar Rapids, IA 52404 Proudly distributed by Circle K Stores Inc., UPC 1 94283 65185 8; g) Sterile Eye Drops, Regular Formula, Lil Drug Store, Product manufactured for: Lil' Drug Store Products, Inc., 9300 Earhart Lane SW, Cedar Rapids, IA 52404; UPC 3 66715 68324 3; h) Tetrahydrozoline Ophthalmic Solution, Rugby, Distributed by: Rugby Laboratories, 17177 N Laurel Park Drive, Suite 233, Livonia, MI 48152, UPC 3 05361 21794 5, NDC 0536-1217-94, i) Leader, DISTRIBUTED BY CARDINAL HEALTH, DUBLIN, OHIO 43017, UPC 0 96295 13645 6, NDC 70000-0454-1; j) REDNESS RELIEF EYE DROPS, CAREone, Distributed by: FOODHOLD USA, LLC, LANDOVER, MD 20785; UPC 0 41520 86531 1, NDC 41520-431-05; k) Eye Drops, Original Formula, Good Neighbor Pharmacy, Distributed By AmerisourceBergen, 1300 Morris Drive, Chesterbrook, PA 19087, UPC 0 87701 14975 7, NDC 24385-075-05; l) Eye Drops, ORIGINAL FORMULA, Walgreens, DISTRIBUTED BY: WALGREEN CO., 200			6/30/2023; m) RG21F03, Exp 6/30/2023; n) RG21F03, Exp 6/30/2023; o) RG21F03, Exp 6/30/2023; p) RG21F03, Exp 6/30/2023		

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
WILMOT RD., DEERFIELD, IL 60015, UPC 3 11917 20076 7; m) CVS Health, Distributed by: CVS Pharmacy, Inc., One CVS Drive, Woonsocket, RI 02895, UPC 0 50428 36131 3; n) H-E-B, MADE WITH PRIDE & CARE FOR H-E-B, SAN ANTONIO, TX 78204, UPC 0 41220 43747 4; o) redness relief, Original Redness Reliever Eye Drops, meijer, DIST. BY MEIJER DISTRIBUTION INC., GRAND RAPIDS, MI 49544, UPC 0 41250 82916 4, NDC 41250-814-01; p) Eye Drops, ORIGINAL, Best Choice, PROUDLY DISTRIBUTED BY: VALU MERCHANDISERS, CO., 5000 KANSAS AVE, KANSAS CITY, KS 66106, UPC 0 70038 47011 3					
Dry Eye Relief Lubricant Eye Drops, (Glycerin 0.2%, Hypromellose 0.2 %, Polyethylene glycol 400 1%), 0.5 FL OZ (15mL) bottle, packaged in a) equate, DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716, 2-0.5 FL OZ (15 mL) Bottles, 1 FL OZ (30 mL) TOTAL, UPC 6 81131 36701 1, NDC 49035-280-02; b) DG health, DISTRIBUTED BY OLD EAST MAIN CO., 100 MISSION RIDGE, GOODLETTSVILLE, TN 37072, UPC 0 95072 02656 0; c) sunmark, Distributed By McKesson, 6555 State Highway 161, Irving, TX 75039, UPC 0 10939 62144 3, NDC 49348-037-29; d) TopCare health, DISTRIBUTED BY TOPCO ASSOCIATES LLC,	Class II	Drugs	Lot #: a) LT21FO2, LT21F03, Exp 6/2023; b) LT21FO2, LT21F03, Exp 6/2023; c) LT21FO2, LT21F03, Exp 6/2023; d) LT21FO2, LT21F03, Exp 6/2023; e) LT21F02, Exp 6/2023; f) LT21F02, Exp 6/2023; g) LT21F03, Exp 6/2023; h) LT21F03, Exp 6/2023; i) LT21F03, Exp 6/2023	CGMP Deviations: good manufacturing deficiencies related to a lack of documentation of the fill line.	K.C. Pharmaceuticals, Inc

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
ELK GROVE VILLAGE, IL 60007, UPC 0 36800 36100 3; e) HealthMart, Distributed by McKesson, 6555 State Highway 161, Irving, TX 75039, UPC 0 52569 13715 4; f) exchange select Artificial Tears, Manufactured for your Military Exchanges by: KC Pharmaceuticals, Inc., Pomona, CA 91768, UPC 6 14299 05620 6; g) meijer, DIST. BY MEIJER DISTRIBUTION, INC., GRAND RAPIDS, MI 49544, UPC 7 13733 29692 2, NDC 41250-718-01; h) H.E.B, MADE WITH PRIDE AND CARE FOR H.E.B, SAN ANTONIO, TX 78204, UPC 0 41220 43741 2; i) GeriCare, Distributed by: Gericare Pharmaceuticals Corp., 1650 63rd St., Brooklyn, NY 11204, UPC 3 57896 18405 6, NDC 57896-181-05					
Big Machine Distillery Hand Sanitizer Non-Sterile Solution Alcohol Antiseptic 80% Topical Solution, Produced and Distributed by Big Machine Distillery 1800 Abernathy Road Lynnville, TN 38472 a) 50 mL UPC 8 59105 00452 5; b) 100 mL UPC 8 59105 00453 2; c) 375 mL; d) 1/2 gallon; e) 1 gallon; f) 5 gallon; g) 55 gallon; i) Indianapolis Motor Speedway 50 mL UPC 8 59105 00452 5; j) Indianapolis Motor Speedway 375 mL UPC 8 59105 00454 9; k) Indianapolis Motor Speedway 1 gallon UPC 8 59105 00451 7; l) The Contributor 50 mL; m) Tony Kanaan Last Lap 100 mL; n) Tony	Class II	Drugs	All lots remaining within expiry.	CGMP Deviation: impurities exceed allowable limits.	Tenn South Distillery LLC dba Big Machine Distillery

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Kanaan Last Lap 375 mL; o) Trexis Insurance 50 mL; p) Team Penske 50 mL; q) Middle Tennessee State University Lightning 50 mL.					
Sunitinib Malate Capsules, 12.5 mg, 28-count bottles, Rx only, Manufactured By: Teva Pharmaceuticals USA, Inc., Parsippany, NJ 07054, NDC 0093-8199-28	Class II	Drugs	Lot # 100037220, Exp 10/2024	Failed Moisture Limits: Water (moisture) content above the approved product specifications.	Teva Pharmaceuticals USA Inc
Atropine Sulfate Injection, USP 8 mg per 20 mL (0.4 mg per mL), 20 mL Multiple Dose Vials, Rx only, Manufactured for: Accord Healthcare, Inc. Durham, NC 27703 Manufactured by: Intas Pharmaceuticals Limited Ahmedabad-382 210, India, NDC 16729-512-43.	Class II	Drugs	Lot #: M2210154 Exp. date 06/2025; M2212575 Exp. date 08/2025	Presence of Particulate Matter: Particulate matter identified as fiber.	Accord Healthcare, Inc.
Bivalirudin for Injection 250 mg, 10 Single-Dose Vials, Rx Only, Manufactured for: Accord Healthcare, Inc. Durham, NC 27703 Manufactured by: Intas Pharmaceuticals Limited Ahmedabad-382 210, India, NDC 16729-275-67.	Class II	Drugs	Lot #: M2212070 Exp. date 08/2024	Presence of Particulate Matter: Particulate matter identified as fiber.	Accord Healthcare, Inc.
0.9% Sodium Chloride Injection USP, 1000 mL Excel Plus Container, Rx only, B. Braun Medical Inc. Bethlehem, PA 18018-3524 USA, NDC 0264-5802-00	Class II	Drugs	Lot#: 0061858305, 0061858306 Exp 3/31/2025	Lack of assurance of sterility: bags have the potential to leak.	B. Braun Medical Inc.
Tiagabine Hydrochloride Tablets, 2 mg, 30-count bottle, Distributed by Sun Pharmaceutical Industries. Inc. Cranbury NJ 08512, Manufactured by: Sun Pharmaceutical Industries Limited Halol-	Class II	Drugs	Lot HAC3339A, Expires 07/2023	Failed Impurities: Out of Specification (OOS) result observed during Related Substances testing	SUN PHARMACEUTICAL INDUSTRIES INC

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Baroda Highway, Halol-389 350, Gujarat, India. NDC#: 62756-200-83					
Tizanidine Tablets USP, 4 mg, 1000-Count bottle, Rx Only, Manufactured by Dr. Reddy's Laboratories Limited Srikakulam - 532 409 India. NDC 55111-180-10	Class II	Drugs	Lot: T2100585, T2100586, T2100587, Exp 12/2023	Failed dissolution specification: Out of specification results observed in 24-month long term stability testing.	Dr Reddy's Laboratories Limited
Famotidine Tablets, USP, 20mg, 200-count bottle within a carton, Distributed by: Glenmark Therapeutics Inc., USA, Mahwah, NJ 07430, Made in India, NDC 72657-113-20.	Class II	Drugs	FA2022001B, Exp 03/2025	Labeling: Label Error on Declared Strength: some cartons labeled and containing 20 mg may have an external label placed on the side of the carton indicating strength as 10 mg.	Glenmark Therapeutics, Inc.
Sodium Bicarbonate in 5% Dextrose Injection, 150 mEq per 1,000 mL (12.6 mg per mL), 1,000 mL Single Dose bag, packaged in 1000 mL x 6 units per case, SterRx, 141 Idaho Ave., Plattsburgh, NY 12903, NDC 70324-326-01.	Class II	Drugs	Lot #: 489344, Exp 25-Jul-23; 489352, Exp 26-Jul-23; 489361, Exp 27-Jul-23; 489387, 491647, Exp 28-Jul-23; 492181, Exp 29-Jul-23; 492199, 492201, Exp 2-Aug-23; 492210, 492228, Exp 3-Aug-23; 492236, 492244, Exp 4-Aug-23; 492261, Exp 5-Aug-23; 492279, 492287, Exp 8-Aug-23; 492295, 492308, Exp 9-Aug-23; 492957, 492965, Exp 19-Sep-23; 492973, 492981, Exp 20-Sep-23; 494194, 494207, Exp 21-Sep-23;494215, 494223, Exp	Lack of Assurance of Sterility	SterRx, LLC

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			22-Sep-23; 494231, 495200, Exp 23-Sep-23; 495218, 495453, Exp 26-Sep-23; 495470, 495488, Exp 27-Sep-23; 495496, 495509, Exp 28-Sep-23; 495517, 495525, Exp 29-Sep-23; 495533, Exp 30-Sep-23; 495699, 495701, Exp 3-Oct-23; 495710, 495728, Exp 4-Oct-23; 496106, 497491, Exp 15-Nov-23; 497504, 497512, Exp 16-Nov-23; 497521, 497539, Exp 22-Nov-23; 497547, 497555, Exp 23-Nov-23; 497563, Exp 24-Nov-23; 497571, Exp 29-Nov-23; 497598, Exp 30-Nov-23; 497619, 498363, Exp 1-Dec-23; 498371, 498380, 498398, Exp 5-Dec-23; 498401, Exp 6-Dec-23; 498419, 498427, Exp 7-Dec-23; 498435, Exp 8-Dec-23; 498443, 498451, Exp 12-Dec-23; 498460, Exp 13-Dec-23; 498478, Exp 14-Dec-23; 498507, Exp 24-Jan-24;		

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			498515, 498523, Exp 25-Jan-24; 498531, 498540, Exp 26-Jan-24; 498558, Exp 27-Jan-24; 498574, Exp 30-Jan-24; 498582, 499665, Exp 31-Jan-24; 500355, 500363, Exp 1-Feb-24; 500371, 500380, Exp 2-Feb-24; 500398, 500401, Exp 3-Feb-24; 500419, 500427, Exp 6-Feb-24; 500435, 500443, Exp 7-Feb-24; 500451, 500460, Exp 8-Feb-24; 500478, Exp 9-Feb-24		
Norepinephrine 4 mg per 250 mL (16 mcg per mL) in 0.9% Sodium Chloride, 250 mL Single Dose bag, packaged in 250 mL x 12 units per case, Rx only, SterRx, 141 Idaho Ave., Plattsburgh, NY 12903, NDC 70324-552-01.	Class II	Drugs	Lot #: 490062, Exp 22-Jul-23; 490151, Exp 28-Jul-23; 490732, Exp 6-Aug-23; 491591, Exp 17-Aug-23; 491604, Exp 18-Aug-23; 494418, Exp 16-Nov-23; 494426, Exp 17-Nov-23; 494434, Exp 23-Nov-23; 496755, Exp 25-Nov-23; 497424, Exp 3-Dec-23; 497432, Exp 4-Dec-23; 497441, 497459, Exp 7-Dec-23; 498005, Exp 10-Dec-23; 499075, Exp 22-Jan-24; 499083, 499091,	Lack of Assurance of Sterility	SterRx, LLC

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			Exp 25-Jan-24; 499391, Exp 26-Jan-24; 501964, Exp 23-Mar-24; 501972, Exp 28-Mar-24; 501981, Exp 30-Mar-24; 501999, Exp 6-Apr-24; 502001, Exp 7-Apr-24		
Norepinephrine 8 mg per 250 mL (32 mcg per mL) in 0.9% Sodium Chloride, 250 mL Single Dose bag, packaged in 250 mL x 12 units per case, Rx only, SterRx, 141 Idaho Ave., Plattsburgh, NY 12903, NDC 70324-577-01.	Class II	Drugs	Lot #: 490071, Exp 30-Jul-23; 490089, Exp 4-Aug-23; 490118, 490126, Exp 5-Aug-23; 490169, Exp 10-Aug-23; 491276, Exp 11-Aug-23; 491284, Exp 12-Aug-23; 491612, Exp 18-Aug-23; 497467, Exp 8-Dec-23; 497475, Exp 9-Dec-23; 499104, Exp 18-Jan-24; 499112, Exp 19-Jan-24; 499121, 499278, Exp 20-Jan-24; 499358, Exp 21-Jan-24; 502019, Exp 24-Mar-24; 502027, Exp 29-Mar-24; 502035, Exp 5-Apr-24; 503513, Exp 18-Apr-24; 504217, Exp 20-Apr-24	Lack of Assurance of Sterility	SterRx, LLC
Norepinephrine 16 mg per 250 mL (64 mcg per mL) in 0.9% Sodium Chloride, 250 mL Single Dose bag, packaged in 250 mL x 12 units per carton, Rx only, SterRx, 141 Idaho	Class II	Drugs	Lot #: 490097, Exp 3-Aug-23; 490142, Exp 10-Aug-23; 491292, Exp 13-Aug-23; 494442, Exp 26-Nov-	Lack of Assurance of Sterility	SterRx, LLC

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Ave., Plattsburgh, NY 12903, NDC 70324-602-01.			23; 496763, Exp 30-Nov-23; 496771, Exp 1-Dec-23; 497408, Exp 2-Dec-23; 497416, Exp 3-Dec-23; 499438, Exp 28-Jan-24; 501032, Exp 8-Feb-24; 501059, Exp 15-Feb-24; 502043, Exp 28-Mar-24; 502051, Exp 31-Mar-24; 503505, Exp 4-Apr-24; 504031, Exp 7-Apr-24; 504250, Exp 19-Apr-24		
Norepinephrine 32 mg per 250 mL (128 mcg per mL) in 0.9% Sodium Chloride, 250 mL Single Dose bag, packaged in 250 mL x 12 units per case, Rx only, SterRx, 141 Idaho Ave., Plattsburgh, NY 12903, NDC 70324-702-01.	Class II	Drugs	Lot #: 490100, Exp 27-Jul-23	Lack of Assurance of Sterility	SterRx, LLC
Fingolimod Capsules, 0.5 mg, 30-count bottle, Rx Only, Manufactured by: Alkem Laboratories Ltd., INDIA; Distributed by: Ascend Laboratories, LLC, Parsippany, NJ 07054, NDC 67877-476-30.	Class II	Drugs	Lot 22122841, Exp August 2025	Failed Dissolution Specifications	Ascend Laboratories, LLC
SUCCINYLcholine Chloride 200 mg/10mL (20 mg/mL), 10 mL BD Syringe, Rx Only, Advanced Compounding Solutions, 4 Constitution Way Ste L Woburn, MA 01801-1042; NDC: 71546-083-10	Class II	Drugs	Lot # 20230524-23C29D, Exp 21SEP2023	CGMP Violations- that spaces adjacent to the production area may have been compromised at the time of production.	New England Life Care, Inc. dba Advanced Compounding Solutions
PHENYLephrine HCl 10mg added to 0.9% Sodium Chloride 250mL, 250 mL IV Bag @60	Class II	Drugs	Lot # 20230524-6FBB77, Exp 21OCT2023	CGMP Violations- that spaces adjacent to the production area	New England Life Care, Inc.

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Total volume), RX only, Advanced Compounding Solutions, 4 Constitution Way Ste L Woburn, MA 01801-1042, NDC: 71546-450-25;				may have been compromised at the time of production.	dba Advanced Compounding Solutions
ROcuronium Bromide 50 mg/5 mL (10 mg/mL), 5mL Syringe, RX Only, Advanced Compounding Solutions, 4 Constitution Way Ste L Woburn, MA 01801-1042, NDC: 71546-090-05	Class II	Drugs	Lot # 20230524-530F73, Exp 21OCT2023	CGMP Violations- that spaces adjacent to the production area may have been compromised at the time of production.	New England Life Care, Inc. dba Advanced Compounding Solutions
Vancomycin HCl 1.5 g added to 0.9% Sodium Chloride 500 mL, 500 mL IV Bag (515 total volume), Rx Only, Advanced Compounding Solutions, 4 Constitution Way Ste L Woburn, MA 01801-1042, NDC 71546-310-50;	Class II	Drugs	Lot # 20230524-0113AB, Exp 22AUG2023	CGMP Violations- that spaces adjacent to the production area may have been compromised at the time of production.	New England Life Care, Inc. dba Advanced Compounding Solutions
PANCREAZE (pancrelipase) Delayed-Release Capsules, 100-count bottles, Rx only, Rx only, Manufactured by VIVUS LLC, Campbell, CA 95008, UPC: N3 62541-401-10 5, NDC 62541-401-10,	Class III	Drugs	Lot #: 102101, Exp: 31 July 2024	Failed Stability Specifications	Vivus, Inc.
Amlodipine Besylate Tablets, USP 10 mg, 1000-count bottles, Rx only, Manufactured for: Lupin Pharmaceuticals, Inc. Baltimore, Maryland 21202 U.S, Manufactured by: Lupin Limited, Aurangabad 431 210 India, NDC 68180-721-03	Class III	Drugs	Lot #: A102887, Exp. 6/2023	Subpotent Drug: Out-of-Specification test results observed in assay test at 21-month long term stability study.	Lupin Pharmaceuticals Inc.
Lotepredrednol Etabonate Ophthalmic Suspension, 5 mg/mL (0.5%), packaged as one bottle in a carton in a) 10 mL bottle (NDC# 62756-232-55) and b) 15 mL bottle (NDC # 62756-232-56), Rx only, Distributed	Class III	Drugs	Lot#: a) BAC0334A, Exp 06/2023; BAC0532A, Exp 11/2023 BAD0407A, Exp 08/2024 BAD0425A, Exp 08/2024; b) BAC0335A,	Superpotent Drug: Out of Specification (OOS) results observed for unit dose content.	SUN PHARMACEUTICAL INDUSTRIES INC

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
by: Sun Pharmaceutical Ind., Inc., NJ 08512, Manufactured by: Sun Pharmaceutical Medicare Ltd., India.			Exp 06/2023, BAC0533A, Exp 10/2023, BAD0148A, Exp 03/2024, BAD0320A, Exp 07/2024		

*Please refer to FDA website for further information at: http://www.fda.gov/Safety/Recalls

CURRENT DRUG SHORTAGES

Below is the list of drugs listed by the FDA as currently in shortage. Please refer to the FDA website for more information at: https://www.accessdata.fda.gov/scripts/drugshortages/default.cfm

Generic Name or Active Ingredient

0.9% Sodium Chloride Irrigation
Albuterol Sulfate Inhalational Solution
Alprostadil (Muse) Suppository
Amifostine Injection
Amino Acids
Amoxapine Tablets
Amoxicillin Oral Powder for Suspension
Amphetamine; Dextroamphetamine Tablets
Atropine Sulfate Injection
Azacitidine for Injection
Azithromycin (Azasite) Ophthalmic Solution 1%
Bacteriostatic 0.9% Sodium Chloride Injection
Bacteriostatic Water for Injection
Belatacept (Nulojix) Lyophilized Powder for Injection
Belladonna and Opium Suppositories
Bumetanide Injection
Bupivacaine Hydrochloride and Epinephrine Injection
Bupivacaine Hydrochloride Injection
Calcium Gluconate Injection
Capecitabine Tablets
Carboplatin Injection
Cefixime Oral Capsules
Cefotaxime Sodium Injection
Cefotetan Disodium Injection
Chloramphenicol Sodium Succinate Injection
Chloroprocaine Hydrochloride Injection
Chlorothiazide Oral Suspension
Cisplatin Injection
Clindamycin Phosphate Injection
Clonazepam Tablets
Collagenase Ointment
Conivaptan Hydrochloride (Vaprisol) in 5% Dextrose Plastic Container
Conjugated Estrogens/Bazedoxifene (DUAVEE) Tablet, Film Coated
Cyclopentolate Ophthalmic Solution
Cytarabine Injection
Dacarbazine Injection
Desmopressin Acetate Nasal Spray
Dexamethasone Sodium Phosphate Injection

Dexmedetomidine Injection
Dextrose 10% Injection
Dextrose 25% Injection
Dextrose 5% Injection
Dextrose 50% Injection
Diazepam Rectal Gel
Diflunisal Tablets
Difluprednate Ophthalmic Emulsion
Digoxin Injection
Diltiazem Hydrochloride Injection
Dimercaprol (Bal in Oil) Injection
Disopyramide Phosphate (Norpace) Capsules
Dobutamine Hydrochloride Injection
Dopamine Hydrochloride Injection
Dulaglutide (Trulicity) Injection
Echothiophate Iodide (Phospholine Iodide) Ophthalmic Solution
Edetate Calcium Disodium Injection
Enalaprilat Injection
Epinephrine Injection, 0.1 mg/mL
Erythromycin Ophthalmic Ointment
Etomidate Injection
Fentanyl Citrate (Sublimaze) Injection
Fludarabine Phosphate Injection
Fluorescein Injection
Flurazepam Hydrochloride Capsules
Furosemide Injection
Gentamicin Sulfate Injection
Guanfacine Hydrochloride Tablets
Heparin Sodium and Sodium Chloride 0.9% Injection
Hydrocortisone Sodium Succinate Injection
Hydromorphone Hydrochloride Injection
Hydroxypropyl (Lacrisert) Cellulose Ophthalmic Insert
Ibutilide Fumarate Injection
Indigotindisulfonate Sodium Injection
Isoniazid Injection
Isoniazid Tablets
IV Fat Emulsion
Ketamine Injection
Ketorolac Tromethamine Injection
Leucovorin Calcium Injection
Lidocaine Hydrochloride (Viscous) Oral Topical Solution
Lidocaine Hydrochloride (Xylocaine) and Dextrose Injection Solution-Premix Bags
Lidocaine Hydrochloride (Xylocaine) Injection
Lidocaine Hydrochloride (Xylocaine) Injection with Epinephrine
Liraglutide Injection
Lisdexamfetamine Dimesylate Capsules

Lorazepam Injection
Lutetium Lu 177 Vipivotide Tetraxetan (Pluvicto) Injection
Mannitol Injection
Mepivacaine Hydrochloride Injection
Methamphetamine Hydrochloride Tablets
Methotrexate Injection
Methotrexate Tablets
Methyldopa Tablets
Methylphenidate Hydrochloride Extended Release Tablets
Methylprednisolone Acetate Injection
Metronidazole Injection
Midazolam Injection
Morphine Sulfate Injection
Multi-Vitamin Infusion (Adult and Pediatric)
Neomycin Sulfate Tablets
Nizatidine Capsules
Oxybutynin Chloride Syrup
Oxytocin Injection
Palifermin (Kepivance) Lyophilized Powder for Injection
Pantoprazole Sodium for Injection
Parathyroid Hormone Injection
Penicillin G Benzathine Injectable Suspension
Physostigmine Salicylate Injection
Potassium Acetate Injection
Potassium Chloride Concentrate Injection
Quinapril and Hydrochlorothiazide Tablets
Quinapril Hydrochloride Tablets
Remifentanil Injection
Rifampin Capsules
Rifampin Injection
Rifapentine Tablets
Rocuronium Bromide Injection
Ropivacaine Hydrochloride Injection
Semaglutide (Ozempic) Injection
Semaglutide (Wegovy) Injection
Sodium Acetate Injection
Sodium Bicarbonate Injection
Sodium Chloride 0.9% Injection Bags
Sodium Chloride 14.6% Injection
Sodium Chloride 23.4% Injection
Sodium Chloride Injection USP, 0.9% Vials and Syringes
Sodium Phosphates Injection
Somatropin Injection
Sterile Water for Injection
Sterile Water for Irrigation
Streptozocin (Zanosar) Sterile Powder

Sucralfate Tablets
Sufentanil Citrate Injection
Sulfasalazine Tablets
Technetium TC-99M Mebrofenin Injection
Tirzepatide Injection
Triamcinolone Acetonide Injectable Suspension
Triamcinolone Hexacetonide Injectable suspension
Trimethobenzamide Hydrochloride Capsules
Valproate Sodium Injection
Vecuronium Bromide for Injection