PERFORMRx℠

Drug Information Update

February 2024

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NEWLY AVAILABLE GENERICS

Generic Name/ Dosage Form	Brand Name	Manufacturer	Indication
gabapentin 300 mg, 600 mg oral tablets	Gralise	Zydus Pharmaceuticals	• Management of Postherpetic Neuralgia (PHN) • Important Limitation: Gralise is not interchangeable with other gabapentin products because of differing pharmacokinetic profiles that affect the frequency of administration
mifepristone 300 mg oral tablets	Korlym	Actavis	• To control hyperglycemia secondary to hypercortisolism in adult patients with endogenous Cushing's syndrome who have type 2 diabetes mellitus or glucose intolerance and have failed surgery or are not candidates for surgery
dabigatran etexilate 110 mg oral capsules	Pradaxa	Apotex	• To reduce the risk of stroke and systemic embolism in adult patients with non-valvular atrial fibrillation • Treatment of deep venous thrombosis (DVT) and pulmonary embolism (PE) in adult patients who have been treated with a parenteral anticoagulant for 5-10 days • To reduce the risk of recurrence of DVT and PE in adult patients who have been previously treated • Prophylaxis of DVT and PE in adult patients who have undergone hip replacement surgery • Treatment of venous thromboembolic events (VTE) in pediatric patients 8 to less than 18 years of age who have been treated with a parenteral anticoagulant for at least 5 days • To reduce the risk of recurrence of VTE in pediatric patients 8 to less than 18 years of age who have been previously treated

NEW DRUG ENTITIES/STRENGTHS/COMBINATIONS

Drug Name	Generic Name	Description
Combogesic Intravenous Solution 1000-300 MG/100ML	Ibuprofen/Acetaminophen	New dosage form. Indicated for the relief of mild to moderate pain and for the management of moderate to severe pain as an adjunct to opioid analgesics in adults, where an IV route of administration is considered clinically necessary.
Rivfloza Subcutaneous Solution Prefilled Syringe 160 MG/ML, 128 MG/ML Rivfloza Subcutaneous Solution 80 MG/0.5ML	Nedosiran	New entity. A ribonucleic acid interference (RNAi) therapy indicated to lower urinary oxalate levels in children 9 years of age and older and in adults with primary hyperoxaluria type 1 (PH1) and relatively preserved kidney function (e.g. estimated glomerular filtration rate [eGFR] ≥30 mL/min/1.73 m2). Second FDA-approved treatment for PH1, after Oxlumo (lumasiran), which was approved in November 2020. Given subcutaneously once monthly, wheras Oxlumo is given once every 3 months.
Udenyca Onbody Subcutaneous Solution Prefilled Syringe 6 MG/0.6ML	Pegfilgrastim-cbqv	New dosage form. On-body injector (OBI) formulation of Udenyca. First FDA-approved biosimilar of Neulasta Onpro. Udenyca is now the only pegfilgrastim product in the U.S. that is available in three administration options: prefilled syringe, autoinjector, and OBI.
Tetracycline HCl Oral Tablet 250 MG, 500 MG	Tetracycline	New dosage form.
fentaNYL Citrate (PF) Injection Solution Prefilled Syringe 25 MCG/0.5ML	Fentanyl Citrate	New strength.
ceFAZolin Sodium Injection Solution Reconstituted 3 GM	Cefazolin	New strength.

Drug Name	Generic Name	Description
Defencath In Vitro Solution 1000-13.5 UNIT-MG/ML	Taurolidine/Heparin	New entity. Catheter lock solution (CLS) indicated to reduce catheter-related bloodstream infections in adult patients with kidney failure who are receiving chronic hemodialysis (HD) through a central venous catheter (CVC). Defencath is indicated in this limited and specific patient population.
Alvaiz Oral Tablet 9 MG, 18 mg, 36 mg, 54 mg	Eltrombopag Choline	New entity. 505(b)(2) approval. Same active ingredient as Promacta, however, it is not substitutable on a milligram per milligram basis.
Xolair Subcutaneous Solution Auto-injector 75 MG/0.5ML Xolair Subcutaneous Solution Auto-injector 150 MG/ML	Omalizumab	New dosage form. Previously only available as 75 mg/0.5 ml, 150 mg/ml subcutaneous prefilled syringes, and 150 mg subcutaneous vial for reconstitution.
Xolair Subcutaneous Solution Prefilled Syringe 300 MG/2ML Xolair Subcutaneous Solution Auto-injector 300 MG/2ML	Omalizumab	New strength and dosage form. Previously only available as 75 mg/0.5 ml, 150 mg/ml subcutaneous prefilled syringes, and 150 mg subcutaneous vial for reconstitution.
Eohilia Oral Suspension 2 MG/10ML	Budesonide	New dosage form. Indicated for 12 weeks of treatment in adult and pediatric patients 11 years of age and older with eosinophilic esophagitis (EoE). Eohilia is the first and only oral treatment approved for patients with EoE. Of note, it has not been shown to be safe and effective for the treatment of EoE for longer than 12 weeks. Eohilia joins Dupixent (dupilumab) as the second treatment option in the EoE space. Prior to the approval of Dupixent and Eohilia, management of EoE included off-label proton pump inhibitors or swallowing inhaled corticosteroids used for asthma.

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†
Zynrelef	bupivacaine/meloxicam 200 mg-6 mg/7 ml, 400 mg-12 mg/14 ml instillation vials	Heron Therapeutics	For postsurgical analgesia in adults ~~for soft tissue or periarticular instillation to produce postsurgical analgesia~~ for up to 72 hours after ~~foot and ankle, small-to-medium open abdominal, and lower extremity total joint arthroplasty surgical procedures~~: • **Soft tissue surgical procedures** • **Orthopedic surgical procedures** o **Foot and ankle procedures** o **Other orthopedic surgical procedures (e.g., total joint arthroplasty) in which direct exposure to articular cartilage is avoided**
Dupixent	dupilumab 200 mg/1.14 ml, 300 mg/2 ml subcutaneous syringe; dupilumab 200 mg/1.14 ml, 300 mg/2 ml subcutaneous pen-injector	Sanofi	• Treatment of adult and pediatric patients aged 6 months and older with moderate-to-severe atopic dermatitis whose disease is not adequately controlled with topical prescription therapies or when those therapies are not advisable • Add-on maintenance treatment of adult and pediatric patients aged 6 years and older with moderate-to-severe asthma characterized by an eosinophilic phenotype or with oral corticosteroid dependent asthma • Add-on maintenance treatment in adult patients with inadequately controlled chronic rhinosinusitis with nasal polyposis (CRSwNP) • Treatment of adult and pediatric patients aged ~~12~~ **1** year~~s~~ and older, weighing at least ~~40~~ **15** kg, with eosinophilic esophagitis (EoE)

Brand Name	Generic Name/ Dosage Form	Manufacturer	Newly Approved Indication†
			• Treatment of adult patients with prurigo nodularis (PN)
Gammagard	immune globulin (human) 1 g/10 ml, 2.5 g/25 ml, 5 g/50 ml, 10 g/100 ml, 20 g/200 ml, 30 g/300 ml injection vials	Takeda Pharmaceuticals	• Replacement therapy for primary humoral immunodeficiency (PI) in adult and pediatric patients two years of age or older • Maintenance therapy to improve muscle strength and disability in adult patients with Multifocal Motor Neuropathy (MMN) • **Therapy to improve neuromuscular disability and impairment in adult patients with Chronic Inflammatory Demyelinating Polyneuropathy (CIDP)** **Limitations of Use:** • **Safety and effectiveness of Gammagard has not been studied in immunoglobulin-naïve patients with CIDP** • **Gammagard maintenance therapy in CIDP has not been studied beyond 6 months**
Onivyde	irinotecan liposome 43 mg/10 ml intravenous vial	Ipsen Biopharmaceuticals, Inc.	• **In combination with oxaliplatin, fluorouracil and leucovorin, for the first-line treatment of adult patients with metastatic pancreatic adenocarcinoma** • In combination with fluorouracil and leucovorin, for the treatment of adult patients with metastatic pancreatic adenocarcinoma after disease progression following gemcitabine-based therapy
Tagrisso	osimertinib 40 mg, 80 mg oral tablets	AstraZeneca	• Adjuvant therapy after tumor resection in adult patients with non-small cell lung cancer (NSCLC) whose tumors have epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 L858R mutations, as detected by an FDA-approved test

Brand Name	Generic Name/ Dosage Form	Manufacturer	Newly Approved Indication†
			• First-line treatment of adult patients with metastatic NSCLC whose tumors have EGFR exon 19 deletions or exon 21 L858R mutations, as detected by an FDA-approved test • **In combination with pemetrexed and platinum-based chemotherapy, for first-line treatment of adult patients with locally advanced or metastatic NSCLC whose tumors have EGFR exon 19 deletions or exon 21 L858R mutations, as detected by an FDA-approved test** • Treatment of adult patients with metastatic EGFR T790M mutation-positive NSCLC, as detected by an FDA-approved test, whose disease has progressed on or after EGFR tyrosine kinase inhibitor (TKI) therapy
Xolair	omalizumab 75 mg/0.5 mL, 150 mg/mL, 300 mg/2 ml subcutaneous syringe; omalizumab 75 mg/0.5 mL, 150 mg/mL, 300 mg/2 mL subcutaneous auto-injector; omalizumab 150 mg subcutaneous vial	Genentech	• Moderate to severe persistent asthma in adults and pediatric patients 6 years of age and older with a positive skin test or in vitro reactivity to a perennial aeroallergen and symptoms that are inadequately controlled with inhaled corticosteroids • Chronic rhinosinusitis with nasal polyps (CRSwNP) in adult patients 18 years of age and older with inadequate response to nasal corticosteroids, as add-on maintenance treatment • **IgE-mediated food allergy in adult and pediatric patients aged 1 year and older for the reduction of allergic reactions (Type I), including anaphylaxis, that may occur with accidental exposure to one or more foods; to be used in conjunction with food allergen avoidance**

Brand Name	Generic Name/ Dosage Form	Manufacturer	Newly Approved Indication†
			• Chronic spontaneous urticaria (CSU) in adults and adolescents 12 years of age and older who remain symptomatic despite H1 antihistamine treatment

RECALLS

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Bleomycin for Injection, USP, 15 units per vial, 1 single dose glass fliptop vial, Rx Only, Distributed by: Hospira, Inc., Lake Forest, IL 68045, NDC 61703-332-18	Class I	Drugs	Lot #: BL12206A, Exp date 06/30/2024	Presence of particulate matter: glass	PFIZER
Robitussin Honey CF Max Non-Drowsy Adult (Acetaminophen 650mg, Dextromethorphan HBr 20 mg), a) 4 FL OZ (118mL) and b) 8 FL OZ (237 mL) bottles, Distributed by: GSK Consumer Healthcare, Warren, NJ 07059	Class I	Drugs	a) Lot#: T10810, Exp 10/31/2025 b) Lot#: T08730, T08731, T08732, T08733, Exp 05/31/2025; T10808, Exp 09/30/2025	Microbial Contamination of Non-Sterile Products	Haleon US Holdings LLC
Robitussin Honey CF Max Nighttime Adult (Acetaminophen 650 mg, Diphenhydramine HCl 25mg), 8 FL OZ (237 mL) bottle, Distributed by: GSK Consumer Healthcare, Warren, NJ 07059	Class I	Drugs	Lot#: T08740, T08742, Exp 06/30/2026	Microbial Contamination of Non-Sterile Products	Haleon US Holdings LLC
FentaNYL citrate PF, 1000 mcg per 100 mL 0.9% Sodium Chloride (10 mcg per mL), Item F3355, for IV use only; Leiters 13796 Compark Blvd Englewood, CO 80112; NDC 71449-072-41.	Class I	Drugs	Lot #: 2331062, Exp. Date: 02/08/2024; 2331224, Exp. Date 03/18/2024; 2331270, Exp. Date 03/28/2024.	Superpotent Drug: semi-automated IV bag filling system can malfunction and provide a double dose of drug product to IV bags.	Denver Solutions, LLC DBA Leiters Health
FentaNYL citrate PF, 2500 mcg per 250 mL 0.9% Sodium Chloride (10 mcg per mL), Item F3342, for IV use only; Leiters 13796 Compark Blvd Englewood, CO 80112; NDC 71449-072-82.	Class I	Drugs	Lot #: 2330988, Exp. Date 01/31/2024; 2331058, Exp. Date 02/18/2024; 2331150, Exp. Date 03/10/2024; 2331231, Exp. Date	Superpotent Drug: semi-automated IV bag filling system can malfunction and provide a double dose of drug product to IV bags.	Denver Solutions, LLC DBA Leiters Health

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			03/24/2024; 2331289, Exp. Date 03/30/2024.		
PHENYLephrine HCl 20 mg per 250 mL 0.9% Sodium Chloride (80 mcg per mL), Item F3360, for IV use only; Leiters 13796 Compark Blvd Englewood, CO 80112; NDC 71449-148-94.	Class I	Drugs	Lot #: 2330993, Exp. Date 02/15/2024; 2331010, Exp. Date 02/10/2024; 2331055, Exp. Date 01/18/2024; 2331113, Exp. Date 02/26/2024; 2331181, Exp. Date 03/04/2024; 2331187, Exp. Date 03/23/2024; 2331266, Exp. Date 03/31/2024; 2331343, Exp. Date 04/01/2024; 2331349, Exp. Date 04/23/2024; 2331433, Exp. Date 05/05/2024.	Superpotent Drug: semi-automated IV bag filling system can malfunction and provide a double dose of drug product to IV bags.	Denver Solutions, LLC DBA Leiters Health
PHENYLephrine HCl 40 mg per 250 mL 0.9% Sodium Chloride (160 mcg per mL), Item F3352, for IV use only; Leiters 13796 Compark Blvd Englewood, CO 80112; NDC 71449-150-82.	Class I	Drugs	Lot #: 2330939, Exp. Date 01/30/2024; 2331032, Exp. Date 02/03/2024; 2331112, Exp. Date 03/19/2024; 2331190, Exp. Date	Superpotent Drug: semi-automated IV bag filling system can malfunction and provide a double dose of drug product to IV bags.	Denver Solutions, LLC DBA Leiters Health

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			03/26/2024; 2331429, Exp. Date 04/28/2024.		
VANCOmycin HCl PF 1.25g added to 0.9% Sodium Chloride 250 mL IV bag, Item F3206, for IV use only; Leiters 13796 Compark Blvd Englewood, CO 80112; NDC 71449-028-68.	Class I	Drugs	Lot #: 2331184, Exp. Date 02/13/2024; 2331185, Exp. Date 02/10/2024; 2331189, Exp. Date 02/20/2024; 2331191, Exp. Date 02/24/2024; 2331258,Exp. Date 03/03/2024; 2331317, Exp. Date 03/15/2024.	Superpotent Drug: semi-automated IV bag filling system can malfunction and provide a double dose of drug product to IV bags.	Denver Solutions, LLC DBA Leiters Health
VANCOmycin HCl PF 1.5g added to 0.9% Sodium Chloride 250 mL IV bag, Item F3208, for IV use only; Leiters 13796 Compark Blvd Englewood, CO 80112; NDC 71449-029-68.	Class I	Drugs	Lot #: 2331140, Exp. Date 02/08/2024; 2331188, Exp. Date 02/15/2024; 2331261, Exp. Date 03/05/2024; 2331287, Exp. Date 03/14/2024.	Superpotent Drug: semi-automated IV bag filling system can malfunction and provide a double dose of drug product to IV bags.	Denver Solutions, LLC DBA Leiters Health
Zenzedi (Dextroamphetamine Sulfate) CII Tablets, USP, 30 mg, 30-count bottle, Rx only, Mfd. for: Arbor Pharmaceuticals, LLC., Atlanta, Georgia, 30328, NDC 24338-856-03	Class I	Drugs	Lot # F230169A, Exp. 06/30/2025	Labeling: Label Mix-up	Azurity Pharmaceuticals, Inc.
CVS Health brand Lubricant Eye Drops (Carboxymethylcellulose Sodium 0.5%), packaged in a) 0.5 FL OZ (15mL) bottles (Single	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Pack) (NDC 76168-702-15) and b) 0.5 FL OZ (15 mL) bottles (Twin pack) (NDC 76168-702-30), Distributed by: CVS Pharmacy Inc. One CVS Drive Woonsocket, RI 02895.					
Leader brand Lubricant Eye Drops (Carboxymethylcellulose Sodium 0.5%), packaged in a) 0.5 FL OZ (15mL) bottles (Single Pack) (NDC 70000-0090-1) and b) 0.5 FL OZ (15 mL) bottles (Twin pack) (NDC 70000-0090-2), Distributed by: Cardinal Health Dublin, Ohio 43017	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
CVS Health brand Lubricant Eye Drops (Propylene glycol 0.6%), packaged in a) 0.33 FL OZ (10mL) bottles (Single Pack) (NDC 76168-714-10) and b) 0.33 FL OZ (10 mL) bottles (Twin pack) (NDC 76168-714-20), Distributed by: CVS Pharmacy Inc. One CVS Drive Woonsocket, RI 02895.	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
Rugby brand Polyvinyl Alcohol 1.4% Lubricating Eye Drops, packaged in 0.5 FL OZ (15 mL) bottles, Distributed by: Rugby Laboratories, Livonia, MI 48152, NDC 0536-1325-94	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
CVS Health brand Lubricating Gel Drops (Polyethylene glycol 400 0.4% and Propylene glycol 0.3%), packaged in 0.33 FL OZ (10 mL) bottles, Distributed by: CVS Pharmacy Inc. One CVS Drive Woonsocket, RI 02895. NDC 76168-712-10	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
Leader brand Dry Eye Relief (Polyethylene glycol 400 0.4% and Propylene glycol 0.3%) packaged	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
in 0.33 FL OZ (10 mL) bottles, Distributed by Cardinal Health Dublin, Ohio 43017, NDC 70000-0088-1					
CVS brand Multi Action Relief Drops (Polyvinyl alcohol 0.5%, Povidone 0.6%, Tetrahydrozoline 0.05%) packaged in 0.5 FL OZ (15mL) bottles, Distributed by CVS Pharmacy Inc. One CVS Drive Woonsocket, RI 02895. NDC 76168-706-15	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
Leader brand Eye Irritation Relief (Polyvinyl alcohol 0.5%, Povidone 0.6%, Tetrahydrozoline 0.05%), packaged in 0.5 FL OZ (15 mL) bottles, Cardinal Health Dublin, Ohio 43017, NDC 70000-0087-1	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
CVS Health brand, Lubricant Gel Drops (Carboxymethylcellulose Sodium 1%) packaged in a) 0.5 FL OZ (15mL) bottles (Single Pack) (NDC 76168-704-15) and b) 0.5 FL OZ (15 mL) bottles (Twin pack) (NDC 76168-704-30), Distributed by: CVS Pharmacy Inc. One CVS Drive Woonsocket, RI 02895.	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
Leader brand Dry Eye Relief (Carboxymethylcellulose Sodium 1%), packaged in 0.5 FL OZ (15 mL) bottles, Cardinal Health Dublin, Ohio 43017, NDC 70000-0089-1	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
Target brand High Performance Lubricant Eye Drops (Polyethylene glycol 400 0.4%, Propylene glycol 0.3%) a) 0.5 FL OZ (15mL) bottles (Single Pack) (NDC 11673-522-15) and b) 0.5 FL OZ (15 mL) bottles (Twin pack) (NDC 11673-522-30),	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Distributed by: Target Corporation Minneapolis, MN 55403					
Leader brand Lubricant Eye Drops (Propylene glycol Eye Drops 0.6%), packaged in 0.33 FL OZ (10 mL) bottles, Cardinal Health Dublin, Ohio 43017, NDC 70000-0587-1	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
Velocity Pharma brand Lubricating Eye Drop (Propylene glycol Eye Drops 0.6%), packaged in 3 bottles of 0.33 FL OZ (10 mL) each, Velocity Pharma, NDC 76168-502-30	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
EQUATE brand Hydration PF Lubricant Eye Drops (Polyethylene glycol 400 0.4% and Propylene glycol 0.3%) packaged in 0.34 FL OZ (10 mL) bottles, Distributed by: Walmart Inc., Bentonville, AR 72712, NDC 79903-168-01	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
LUBRICANT EYE DROPS (Carboxymethylcellulose Sodium 0.5%), packaged 0.5 FL OZ (15mL) bottles (Twin Pack), Distributed by: Rite Aid 30 Hunter Lane, Camp Hill, PA 17011, NDC 11822-4811-5	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
CVS Health brand Mild Moderate Lubricating Eye Drops (Propylene glycol 400 0.25%), packaged in 0.5 FL OZ (15 mL) bottles, Distributed by: CVS Pharmacy Inc. One CVS Drive Woonsocket, RI 02895. NDC 76168-711-15	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
Lubricant Gel Drops (Carboxymethylcellulose sodium 1.0%), packaged in 0.5 FL OZ (15 mL) bottles, Distributed by: Rite Aid 30 Hunter Lane, Camp Hill, PA 17011, NDC 11822-4540-5	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Lubricant Gel Drops (Polyethylene glycol 400 0.4%, Propylene glycol 0.3%), packaged in 0.33 FL OZ (10 mL) bottles, Distributed by: Rite Aid 30 Hunter Lane, Camp Hill, PA 17011, NDC 11822-4540-3	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
Multi-action Relief Drops (Polyvinyl alcohol 0.5%, Providone 0.6%, Tetrahydrozoline Hydrochloride 0.05%), packaged in 0.5 FL OZ (15 mL) bottles, Distributed by: Rite Aid 30 Hunter Lane, Camp Hill, PA 17011, NDC 11822-2254-3	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
Lubricant Eye Drops (Propylene glycol 0.6%), packaged in 0.33 FL OZ (10 mL) each bottle (Twin Pack), Distributed by: Rite Aid 30 Hunter Lane, Camp Hill, PA 17011, NDC 11822-4811-3 Retail Labeler: Rite Aid	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
Rugby brand Lubricating Tears Eye Drops (Dextran 70 0.1%, Hypromellose 2910 0.3%), packaged in 0.5 FL OZ (15 mL) bottles, Distributed by: Rugby Laboratories, Livonia, MI 48152, NDC 0536-1282-94	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
Up&Up brand dry eye relief (Carboxymethylcellulose 0.5%) packaged in 0.5 FL OZ (15 mL) each bottles (Twin Pack), Distributed by: Target Corporation Minneapolis, MN 55403, NDC 76168-800-30	Class I	Drugs	All lots	Non-Sterility	Kilitch Healthcare India Limited
Methoxsalen Capsules, USP 10mg, 50-count bottle, Rx Only, Manufactured by: Strides Pharma Science Ltd. Bengaluru - 562106, India. Distributed by:Strides Pharma Inc. East Brunswick, NJ 08816, NDC 64380-752-16	Class II	Drugs	Lot #: 7253092B, Exp Date 09/30/2025	Failed Dissolution Specifications	Strides Pharma Inc.

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Fluocinolone Acetonide Oil, 0.01% (Ear Drops), package in 1 Oz. (20 mL fill volume) bottle, Rx only, Manufactured by: Glenmark Pharmaceuticals Ltd., Village: Kishanpura, Baddi Nalagarh Road, District: Solan, Himachal Pradesh - 173205, India, Manufactured for: Glenmark Pharmaceuticals Inc., USA, Mahwah, NJ 07430, NDC 68462-185-56	Class II	Drugs	Lot #: 05220346, 05220369 Exp 1/31/ 2024; 05220582, Exp 2/29/2024; 05220861 Exp 3/31/2024	Failed Excipient Specifications: OOS for assay of Isopropyl Alcohol	Glenmark Pharmaceuticals Inc., USA
Fosaprepitant for Injection 150 mg per vial, Sterile lyophilized powder for Intravenous use only after reconstitution and dilution, Single Dose Vial, Rx Only, Distributed by BE Pharmaceuticals Inc. 203 New Edition Court Cary, NC 27511, Made in India, NDC 71839-104-01.	Class II	Drugs	lot #13D012AA, Exp: 08/31/2025	Lack of Sterility Assurance: Aseptic process simulation failure.	BE PHARMACEUTICALS AG
Cinacalcet Tablets 60mg, Rx Only, 30 Tablets per bottle, Distributed by: Sun Pharmaceutical Industries, Inc., Cranbury, NJ 08512, Manufactured by: Sun Pharmaceutical Industries Limited Survey No. 259/15, Dadra-396 191 (U.T. of D & NH), India, NDC 47335-380-83.	Class II	Drugs	Lot #: DNE0702A, Exp. 06/30/2026	CGMP Deviations: Microbial contamination was reported in stagnant water in the duct of the manufacturing equipment.	SUN PHARMACEUTICAL INDUSTRIES INC
Febuxostat Tablets 40mg, Rx Only, 30 Tablets per bottle, Distributed by: Sun Pharmaceutical Industries, Inc., Cranbury, NJ 08512, Manufactured by: Sun Pharmaceutical Industries Ltd., Survey No. 259/15, Dadra-396 191 (U.T. of D & NH), INDIA, NDC 47335-721-83.	Class II	Drugs	Lot #s: DNE0866B Exp. 06/30/2025, DNE1045A, DNE1046B Exp. 08/31/2025	CGMP Deviations: Microbial contamination was reported in stagnant water in the duct of the manufacturing equipment.	SUN PHARMACEUTICAL INDUSTRIES INC

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Febuxostat Tablets 80 mg, Rx Only, 30 Tablets per bottle, Distributed by: Sun Pharmaceutical Industries, Inc., Cranbury, NJ 08512, Manufactured by: Sun Pharmaceutical Industries Ltd., Survey No. 259/15, Dadra-396 191, (U.T. of D & NH), INDIA, NDC 47335-722-83.	Class II	Drugs	Lot #s: DNE0867A Exp. 06/30/2025, DNE0894B Exp. 07/31/2025	CGMP Deviations: Microbial contamination was reported in stagnant water in the duct of the manufacturing equipment.	SUN PHARMACEUTICAL INDUSTRIES INC
Lurasidone Hydrochloride Tablets 60mg, Rx Only, 30 Tablets per bottle, Distributed by: Sun Pharmaceutical Industries, Inc., Cranbury, NJ 08512, Manufactured by: Sun Pharmaceutical Industries Ltd., Survey No. 259/15, Dadra-396 191(U.T. of D & NH), INDIA, NDC 47335-639-83.	Class II	Drugs	Lot #: DNE0620A Exp. 05/31/2025	CGMP Deviations: Microbial contamination was reported in stagnant water in the duct of the manufacturing equipment.	SUN PHARMACEUTICAL INDUSTRIES INC
Lurasidone Hydrochloride Tablets 120mg, Rx Only, 30 Tablets per bottle, Distributed by: Sun Pharmaceutical Industries, Inc., Cranbury, NJ 08512, Manufactured by: Sun Pharmaceutical Industries Ltd., Survey No. 259/15, Dadra-396 191(U.T. of D & NH), INDIA, NDC 47335-579-83.	Class II	Drugs	Lot #s: DNE0621A Exp. 11/30/2024, DNE0815A Exp. 12/31/2024	CGMP Deviations: Microbial contamination was reported in stagnant water in the duct of the manufacturing equipment.	SUN PHARMACEUTICAL INDUSTRIES INC
Mesalamine Delayed-Release Tablets, USP 1.2 g per tablet, Rx Only, 120 Tablets per bottle, Once Daily, Distributed by: Sun Pharmaceutical Industries, Inc., Cranbury, NJ 08512, Manufactured by: Sun Pharmaceutical Industries Limited Survey No. 259/15, Dadra-396 191 (U.T. of D & NH), INDIA, NDC 63304-175-13.	Class II	Drugs	Lot #s: DNE0875A Exp. 01/31/2025; DNE0876A, DNE0877A, DNE1080A, DNE1081A Exp. 02/28/2025; DNE1147A, DNE1148A Exp. 03/31/2025.	CGMP Deviations: Microbial contamination was reported in stagnant water in the duct of the manufacturing equipment.	SUN PHARMACEUTICAL INDUSTRIES INC

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Niacin Extended-Release Tablets, USP 1000mg, Rx Only, 90 Tablets per bottle, Distributed by: Sun Pharmaceutical Industries, Inc., Cranbury, NJ 08512, Manufactured by: Sun Pharmaceutical Industries Limited, Survey No. 259/15, Dadra-396 191(U.T. of D & NH), INDIA, NDC 47335-613-81.	Class II	Drugs	Lot #: DNE0788A Exp. 07/31/2025	CGMP Deviations: Microbial contamination was reported in stagnant water in the duct of the manufacturing equipment.	SUN PHARMACEUTICAL INDUSTRIES INC
Niacin Extended-Release Tablets, USP 500mg, Rx Only, 90 Tablets per bottle, Distributed by: Sun Pharmaceutical Industries, Inc., Cranbury, NJ 08512, Manufactured by: Sun Pharmaceutical Industries Limited, Survey No. 259/15, Dadra-396 191(U.T. of D & NH), INDIA, NDC 47335-539-81.	Class II	Drugs	Lot #s: DNE0771A Exp. 06/30/2025; DNE0857A, DNE0959A Exp. 07/31/2025.	CGMP Deviations: Microbial contamination was reported in stagnant water in the duct of the manufacturing equipment.	SUN PHARMACEUTICAL INDUSTRIES INC
Zolpidem Tartrate Extended-Release Tablets, USP 6.25 mg, Rx Only, 100 Tablets per bottle, Distributed by: Sun Pharmaceutical Industries, Inc., Cranbury, NJ 08512, Manufactured by: Sun Pharmaceutical Industries Limited, Survey No. 259/15, Dadra-396 191, (U.T. of D & NH), INDIA, NDC 47335-307-88.	Class II	Drugs	Lot #: DNE0892A Exp. 07/31/2026	CGMP Deviations: Microbial contamination was reported in stagnant water in the duct of the manufacturing equipment.	SUN PHARMACEUTICAL INDUSTRIES INC
Zolpidem Tartrate Extended-Release Tablets, USP 12.5mg, Rx Only, 100 Tablets per bottle, Distributed by: Sun Pharmaceutical Industries, Inc., Cranbury, NJ 08512, Manufactured by: Sun Pharmaceutical Industries Limited, Survey No. 259/15, Dadra-396 191, (U.T. of D & NH), INDIA, NDC 47335-308-88.	Class II	Drugs	Lot #: DNE0893A Expires 07/31/2026	CGMP Deviations: Microbial contamination was reported in stagnant water in the duct of the manufacturing equipment.	SUN PHARMACEUTICAL INDUSTRIES INC

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Fexofenadine Hydrochloride Tablets, USP 180mg, Antihistamine, Allergy, 24 hour, a) 30 Tablets per bottle, NDC 51660-998-30; Distributed by: Ohm Laboratories Inc., New Brunswick, NJ 08901, Made in India, b) 45 Tablets per bottle, NDC 51316-800-45; Distributed by: CVS Pharmacy, Inc., Woonsocket, RI 02895, Made in India, c) 150 Tablets per bottle, NDC 51660-998-55, Distributed by: Ohm Laboratories Inc., New Brunswick, NJ 08901, Made in India.	Class II	Drugs	Lot #s: a) DNE0792A Exp. 06/31/2025; DNE1027A Exp. 08/31/2025. b) DNE0793A Exp. 06/31/2025. c) DNE0789A, DNE0790A, DNE0791A Exp. 06/2025, DNE1026A Exp. 08/31/025.	CGMP Deviations: Microbial contamination was reported in stagnant water in the duct of the manufacturing equipment.	SUN PHARMACEUTICAL INDUSTRIES INC
Cefixime for Oral Suspension 100mg/5mL, 50 mL bottle, Rx Only, Manufactured for: Lupin Pharmaceuticals, Inc., Baltimore, Maryland 21202, Manufactured by:Lupin Limited, Mandideep 462 046 INDIA, NDC 68180-405-01	Class II	Drugs	Lot #: F304833, Exp 06/2025	Failed Impurities/Degradation Specifications	Lupin Pharmaceuticals Inc.
Budesonide Extended-Release Tablets 9mg, 30-count bottle, Rx Only, Manufactured by: Actavis Laboratories FL, Inc., Fort Lauderdale, FL 33314 USA, Distributed by: Actavis Pharma, Inc., Parsippany, NJ 07054 USA, NDC 0591-2510-30	Class II	Drugs	Lot # 100047273; Exp. 07/2025	Failed Dissolution Specifications	Teva Pharmaceuticals USA, Inc
Rifampin Capsules USP 150mg, 30-count bottle, Rx Only, Manufactured for: Lupin Pharmaceuticals, Inc. Baltimore, Maryland 21202 United States, Manufactured Lupin Limited, Aurangabad 431 210 INDIA, NDC 68180-658-06	Class II	Drugs	A200816 exp 1/2024 A201248 exp 3/2024	Failed Impurities/Degradation Specifications	Lupin Pharmaceuticals Inc.

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Rifampin Capsules USP 300mg, 30-count bottle, Rx Only, Manufactured for: Lupin Pharmaceuticals, Inc. Baltimore, Maryland 21202 United States, Manufactured Lupin Limited, Aurangabad 431 210 INDIA, NDC 68180-659-06	Class II	Drugs	A200817 exp 1/2024	Failed Impurities/Degradation Specifications	Lupin Pharmaceuticals Inc.
OralProCare medicated lip treatment, Net Wt 9.9 g (0.35 oz) tube, Ethyl alcohol 6.0% Antibacterial, Manufactured by: Den-Mat Holdings, LLC, Lompoc, CA, 93436, USA, NDC 59883-500-01, UPC: 3 59883 00000 4.	Class II	Drugs	Lot #s: 2319500023, Exp: 01- 12-2024; 2330400002, Exp: 11-02-2024.	CGMP Deviations: products may not conform to the labeled specifications.	Den-Mat Holdings, LLC
Hydrogen Peroxide Oral Rinse, Significantly Reduces Bacteria, Fresh Mint Flavor, Alcohol Free, a)16 fl. oz. (473 mL) bottle, NDC 59883-202-16, UPC 3 59883 00009 7); b) 64 fl. oz. (1.89 L) bottle, NDC 59883-202-64; c) 128 fl. oz. (1 gal) 3.78L bottle, NDC 59883-202-28, UPC 3 59883 00007 3), Manufactured by: Den-Mat Holdings, LLC, Lompoc, CA 93436.	Class II	Drugs	Lot #: a) 2214500032, 2216700041, 2301900118, 2312600002, 2204800002, Exp. 02/05/2024; 2306100034, 2322000078, 2322200021 02/22/2024; 2318000026, Exp 02/24/2024; 2323000070, 2324400005, 2323300016, Exp 08/29/2025; 2324800079, Exp 08/30/2025; 2325800008, Exp	CGMP Deviations: products may not conform to the labeled specifications.	Den-Mat Holdings, LLC

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			09/18/2025; 2328600011, 2328600032, 10/10/2025; 2324800123, Exp 10/11/2025; 2329700005, 2329700090, Exp 10/19/2025; 2330400003, Exp 10/25/2025; 2334100101, Exp 11/13/2025; 2335300019, 2335400038, Exp 12/19/2025. b) 2212200014, Exp 02/22/2024; 2323300017, Exp 08/29/2025; 2324800124, Exp 09/18/2025. c) 2211100001, Exp 02/24/2024; 2330500009, Exp 10/25/2025; 2333200014, Exp12/19/2025.		

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Rifampin Capsules USP, 150 mg, 30 Capsules (3 x 10) unit doses per carton, Rx Only, Distributed by: American Health Packaging, Columbus, Ohio 43217. NDC Carton: 60687-575-21; NDC Unit Dose: 60687-575-11	Class II	Drugs	Lot #: 1008111, Exp. Date 01/31/2024	Failed Impurities/Degradation Specification.	Amerisource Health Services LLC
Phenoxybenzamine Hydrochloride Capsules, USP 10mg, 100-count bottle, Rx only, Manufactured by: Amneal Pharmaceuticals Pvt. Ltd, Oral Solid Dosage Unit, Ahmedabad 382213, India, Distributed by: Amneal Pharmaceuticals LLC, Bridgewater, NJ 08807, NDC 60219-1502-01	Class II	Drugs	Lot # AM221153, Exp. date 06/30/2024; AM230497, Exp. date 02/29/2025	Failed Impurities/Degradation Specifications: Out of specification for unknown impurity.	Amneal Pharmaceuticals of New York, LLC
Medline Remedy Clinical TREAT Antifungal Cream, 2% Miconazole Nitrate, Paraben Free, 4 FL OZ (118 mL) tube, Manufactured for Medline Industries, LP Three Lakes Drive, Northfield, IL 60093. NDC: 53329-147-44	Class II	Drugs	Lot #s: 08926; 08926A Exp. 8/2025; 08928 Exp. 9/2025	Labeling: Not Elsewhere Classified; Product labeling contains the claim of 'Paraben Free' while the product does in fact contain parabens as part of the formulation/ ingredient list.	MEDLINE INDUSTRIES, LP - Northfield
Omeza Lidocaine Lavage pain relief oil 10*2mL VIALS	Class II	Drugs	Lots: 28622 2-1, Exp. 04/24/2024 286222-3, Exp. 1/25/2024 or 04/25/2024	CGMP Deviations	OMEZA LLC
Omeza Skin Protectant, Skin Protectant Gel 10*2mL Vials	Class II	Drugs	Lot: 19123 4-1, Exp. 05/11/2024	CGMP Deviations	OMEZA LLC
Lansoprazole Delayed-Release Capsules USP, 15 mg, 14-count bottle, Manufactured by: Natco Pharma Limited Kothur- 509 228, India,	Class II	Drugs	Lot # 411987 Exp: 05/2025	CGMP Deviations: Inadequate induction sealing on bottles,	NATCO Pharma Limited

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Distributed by: Rising Pharma Holdings, Inc. East Brunswick, NJ. 08816. NDC 16571-742-41				capsules were observed with bubbles on band seal, capsules with holes and spheres sticking to capsules. Also coding details were missing on one bottle.	
Febuxostat Tablets, 40 mg, 30 Tablets (3 x 10) per carton, Rx only, Distributed by: American Health Packaging, Columbus, Ohio 43217. NDC Carton: 60687-538-21, NDC Unit dose 60687-538-11	Class II	Drugs	Lot # 1015033, exp. 06/30/2025; 1016409, exp. 08/31/2025	CGMP Deviations	Amerisource Health Services LLC
HydrALAZINE Hydrochloride Tablets, USP 10mg, packaged in 100-count (10x10 blister cards), Lot T04680, Rx only, Manufactured for Heritage Pharmaceuticals, Inc. Eatontown, NJ, Distributed by Major Pharmaceuticals Livonia MI. NDC 0904-6440-61	Class II	Drugs	Lot #: T04680, Exp. Date 6/2024	an out of specification result obtained during routine stability testing for Impurities. There is a remote possibility that use of this product could cause a medically reversible or transient adverse health consequences.	The Harvard Drug Group LLC dba Major Pharmaceuticals and Rugby Laboratories
Vasopressin 2 Units/2 mL in 0.9% Sodium Chloride Sterile Syringe for Injection, Concentration: 1 Unit/mL, 2 mL Syringe, Rx Only, 719 Kasota Ave SE, Minneapolis, MN, Compounded Drug Not for Resale. Office Use Only, NDC 71139-0190-1.	Class II	Drugs	Lot #s: 20230929VAS-2, Exp. 02/29/2024; 20231004VAS-2, Exp. 04/01/2024; Lot 20231010VAS-2,Exp. 04/07/2024; Lot	Sub-potent drug: failure to maintain potency through the duration of the labeled expiration/beyond-use date.	IntegraDose Compounding Services LLC

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			20231013VAS-2, Exp. 04/10/2024		
Benzonatate Capsules, USP, 100 mg, 100 Capsules (10 capsules x 10 unit dose cards), Rx only, Distributed by: American Health Packaging, Columbus, Ohio 43217. Carton NDC 68084-214-01; Individual Dose NDC 68084-214-11	Class III	Drugs	Lot # 1014208, Exp Mar/31/2025	Superpotent drug: Assay results were slightly above specification at the time zero point.	Amerisource Health Services LLC
Glimepiride Tablets, USP 1mg, 100-count bottles, Rx Only, Mfd. By: Dr. Reddy's Laboratories Limited, Srikakulam-532 409 INDIA, NDC: 55111-320-01	Class III	Drugs	Lot: T2303622; Exp. 06/2026 Lot: T2303626; Exp. 06/2026 Lot: T2303627; Exp. 06/2026 Lot: T2303628; Exp. 06/2026 Lot: T2303629; Exp. 06/2026	Misprint on tablet	Dr. Reddy's Laboratories, Inc.
Glimepiride Tablets, USP 1mg, 500-count bottles, Rx Only, Mfd. By: Dr. Reddy's Laboratories Limited, Srikakulam-532 409 INDIA, NDC: 55111-320-05	Class III	Drugs	Lot: T2303609; Exp. 06/2026 Lot: T2303610; Exp. 06/2026	Misprint on tablet	Dr. Reddy's Laboratories, Inc.
HydrALAZINE Hydrochloride Tablets, USP, 10 mg, 100-count (10 x 10) per unit dose carton, Rx Only, Distributed by: American Health Packaging, Columbus, Ohio 43217. NDC Carton: 68084-447-01; NDC Unit Dose: 68084-447-11	Class III	Drugs	Lot #: 1012275, Exp 02/28/2025	Failed Impurities/Degradation Specifications: Out of Specification (OOS) result in the repackaged product for Related Compounds (Impurities)	Amerisource Health Services LLC

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
				at the 6-month time point.	
Nortrel (norethindrone and ethinyl estradiol tablets USP) 0.5/35, packaged in cartons, each carton contains 3 blister cards, each card contains 28 tablets, Rx only, TEVA PHARMACEUTICALS USA, INC., North Wales, PA 19454, NDC 0555-9008-67	Class III	Drugs	Lot #: 100042978, Exp 7/31/2024	Discoloration: discolored tablets (shades of blue) mixed in with the white inert remainder tablets.	Teva Pharmaceuticals USA, Inc
Nortrel 7/7/7 (norethindrone and ethinyl estradiol tablets USP- triphasic regimen), packaged in cartons, each carton contains 6 blister cards, each card contains 28 tablets, Rx only, TEVA PHARMACEUTICALS USA, INC., North Wales, PA 19454, NDC 0555-9012-58	Class III	Drugs	Lot #: 100040731, Exp 7/31/2024	Discoloration: discolored tablets (shades of blue) mixed in with the white inert remainder tablets.	Teva Pharmaceuticals USA, Inc

*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

Albuterol Sulfate Solution
Alprostadil Suppository
Amifostine Injection
Amino Acid Injection
Amoxapine Tablet
Amoxicillin Powder, For Suspension
Amphetamine Aspartate Monohydrate, Amphetamine Sulfate, Dextroamphetamine Saccharate, Dextroamphetamine Sulfate Tablet
Atropa Belladonna, Opium Suppository
Atropine Sulfate Injection
Azacitidine Injection
Bazedoxifene Acetate, Estrogens, Conjugated Tablet, Film Coated
Bumetanide Injection
Bupivacaine Hydrochloride Injection
Bupivacaine Hydrochloride, Epinephrine Bitartrate Injection, Solution
Capecitabine Tablet
Carboplatin Injection
Cefixime Capsule
Cefotaxime Sodium Injection
Cefotetan Disodium Injection
Chloroprocaine Hydrochloride Injection
Cisplatin Injection
Clindaymycin Phosphate Injection
Clonazepam Tablet
Collagenase Clostridium Histolyticum Ointment
Cromolyn Sodium Concentrate
Cyclopentolate Hydrochloride Ophthalmic Solution
Cyclopentolate Hydrochloride, Phenylephrine Hydrochloride Ophthalmic Solution
Cytarabine Injection, Solution
Dacarbazine Injection
Desmopressin Acetate Spray
Dexamethasone Sodium Phosphate Injection
Dexmedetomidine Hydrochloride Injection
Dextrose Monohydrate Injection
Dextrose Monohydrate, Lidocaine Hydrochloride Anhydrous Injection
Diazepam Gel
Difluprednate Emulsion
Digoxin Injection

Diltiazem Hydrochloride Injection
Disopyramide Phosphate Capsule
Dobutamine Hydrochloride Injection
Dopamine Hydrochloride Injection
Dulaglutide Injection
Echothiophate Iodide Ophthalmic Solution
Enalaprilat Injection
Epinephrine Bitartrate, Lidocaine Hydrochloride Injection
Epinephrine Injection
Erythromycin Ointment
Etomidate Injection
Fentanyl Citrate Injection
Fluconazole Injection
Fludarabine Phosphate Injection
Flurazepam Hydrochloride Capsule
Furosemide Injection
Gentamicin Sulfate Injection
Heparin Sodium Injection
Hydrocortisone Sodium Succinate Injection
Hydromorphone Hydrochloride Injection
Hydroxypropyl Cellulose (1600000 Wamw) Insert
Indigotindisulfonate Sodium Injection
Isoniazid Tablet
Ketamine Hydrochloride Injection
Ketorolac Tromethamine Injection
Leucovorin Calcium Injection
Lidocaine Hydrochloride Injection
Lidocaine Hydrochloride Solution
Liraglutide Injection
Lisdexamfetamine Dimesylate Capsule
Lisdexamfetamine Dimesylate Tablet, Chewable
Lorazepam Injection
Methamphetamine Hydrochloride Tablet
Methotrexate Sodium Injection
Methotrexate Sodium Tablet
Methylphenidate Hydrochloride Tablet, Extended Release
Methylprednisolone Acetate Injection
Metronidazole Injection
Midazolam Hydrochloride Injection
Morphine Sulfate Injection
Neomycin Sulfate Tablet
Nitroglycerin Injection
Nizatidine Capsule
Oxybutynin Chloride Syrup
Parathyroid Hormone Injection
Penicillin G Benzathine Injection

Potassium Acetate Injection
Potassium Chloride Injection
Promethazine Hydrochloride Injection
Propranolol Hydrochloride Injection
Quinapril Hydrochloride Tablet
Quinapril/Hydrochlorothiazide Tablet
Remifentanil Hydrochloride Injection
Rifampin Capsule
Rifampin Injection
Rifapentine Tablet, Film Coated
Riluzole Oral Suspension
Rocuronium Bromide Injection
Ropivacaine Hydrochloride Injection
Semaglutide Injection
Sodium Acetate Injection
Sodium Bicarbonate Injection
Sodium Chloride 0.9% Injection
Sodium Chloride 0.9% Irrigation
Sodium Chloride 14.6% Injection
Sodium Chloride 23.4% Injection
Sodium Phosphate, Dibasic, Anhydrous, Sodium Phosphate, Monobasic, Monohydrate Injection, Solution
Sodium Pyrophosphate Injection
Somatropin Injection
Sterile Water Injection
Sterile Water Irrigant
Streptozocin Powder, For Solution
Sucralfate Tablet
Sufentanil Citrate Injection
Sulfasalazine Tablet
Technetium TC-99M Pyrophosphate Kit Injection
Tirzepatide Injection
Triamcinolone Acetonide Injection
Triamcinolone Hexacetonide Injection
Trimethobenzamide Hydrochloride Capsule
Valproate Sodium Injection
Vecuronium Bromide Injection
Vinblastine Sulfate Injection