PERFORMRx℠

Drug Information Update

August 2022

TABLE OF CONTENTS

NEWLY AVAILABLE GENERICS

Generic Name/ Dosage Form	Brand Name	Manufacturer	Indication
Barium sulfate 2% oral suspension	Readi-cat 2	Bracco Diagnostics	For use in computed tomography (CT) of the abdomen to delineate the gastrointestinal (GI) tract in adult and pediatric patients

NEW DRUG ENTITIES/STRENGTHS/COMBINATIONS

Drug Name	Generic Name	Description	Comments
ammonia N-13 3.75 mCi to 37.5 mCi/mL intravenous syringe	ammonia n-13	New syringe version of diagnostic agent used in PET imaging of the heart. It was previously only available as a vial.	New Entity
Injectafer 100 mg iron/2 mL intravenous solution	ferric carboxymaltose	New smaller package size of 100 mg/2mL to accommodate younger patients. Was previously only available as 750 mg/15 mL. Indication was expanded to pediatric patients aged 1 year and older, in 11/2021, for patients who have an intolerance or unsatisfactory response to oral iron or non-dialysis dependent CKD.	New package size
Tascenso ODT 0.25 mg disintegrating tablet	fingolimod lauryl sulfate	New ODT version of fingolimod having the same indication of relapsing forms of MS in patients 10 years and older weighing less than or equal to 40 kg.	New entity
Caplyta 10.5 mg capsule	lumateperone tosylate	New lower strength capsules likely to be used in patients with moderate to severe hepatic impairment. It was previously only available as a 42 mg capsule. Caplyta was previously only indicated in schizophrenia but has a new indication for bipolar I and II as of 12/2021.	New dosage strength
Caplyta 21 mg capsule	lumateperone tosylate	New lower strength capsules. It was previously only available as a 42 mg capsule. Caplyta was previously only indicated in schizophrenia but has a new indication for bipolar I and II as of 12/2021.	New dosage strength
carmustine 50 mg intravenous solution	carmustine	New strengths of carmustine. Was previously only available as 100 mg vials.	New strength
carmustine 300 mg intravenous solution	carmustine	New strengths of carmustine. Was previously only available as 100 mg vials.	New strength

Drug Name	Generic Name	Description	Comments
Flumist Quad 2022-2023 10exp6.5-7.5 FF unit/0.2 mL nasal spray syringe	Flu vacc qv live 2022(2-49yrs)	Influenza vaccine	2022-2023 vaccine
Qwo 1.84 mg subcutaneous solution	collagenase C. Histolyt-aaes	New cosmetic drug used to treat moderate to severe cellulite in the buttocks of adult women.	New entity
Qwo 0.92 mg subcutaneous solution	collagenase C. Histolyt-aaes	New cosmetic drug used to treat moderate to severe cellulite in the buttocks of adult women.	New entity
quetiapine 150 mg tablet	quetiapine fumarate	New strength of quetiapine having the same indications (Schizophrenia, PTSD, Bipolar disorder, GAD, etc).	New Strength
Zoryve 0.3% topical cream	roflumilast	First topical PDE-4 inhibitor approved for plaque psoriasis, including intertriginous areas, in patients 12 years and older.	New Route, Dosage Form and Strength
Entadfi 5 mg-5 mg capsule	finasteride/tadalafil	Combination of finasteride and tadalafil indicated to initiate treatment of the signs and symptoms of BPH in men with an enlarged prostate for up to 26 weeks.	New Combination
Calquence (maleate) 100 mg tablet	Acalabrutinib maleate	New tablet dosage form of acalabrutinib, a medication used to treat various types of non-Hodgkin lymphoma, previously only in capsule form	New Dosage Form
fluorodopa F-18 0.42 mCi/mL to 8.33 mCi/mL intravenous solution	Fluorodopa f-18	New radiopharmaceutical injection is indicated for use in positron emission tomography (PET) to visualize dopaminergic nerve terminals in the striatum for the evaluation of adult patients with suspected Parkinsonian syndromes (PS)	New Entity
fluorodopa F-18 1 mCi/mL to 40 mCi/mL intravenous solution	Fluorodopa f-18	New radiopharmaceutical injection is indicated for use in positron emission tomography (PET) to visualize dopaminergic nerve terminals in the striatum for the evaluation of adult patients with suspected Parkinsonian syndromes (PS)	New Entity

Drug Name	Generic Name	Description	Comments
pirfenidone 534 mg tablet	Pirfenidone	New strength for pirfenidone, a drug indicated in adults for the treatment of mild to moderate idiopathic pulmonary fibrosis; was previously only available as 267 mg, 801 mg tablets	New Strength
Xaciato 2 % vaginal gel	Clindamycin phosphate	New dosage form of xaciato, approved for treatment of bacterial vaginosis; previously oral clindamycin or vaginal ovules only available dosage forms	New Dosage Form

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†
Benlysta	Belimumab 200 mg/mL IV/SQ autoinjector, syringe; 120, 400 mg IV/SQ vial	GlaxoSmithKline	B-lymphocyte stimulator (BLyS)-specific inhibitor indicated for the treatment of: • patients aged 5 years and older with active, autoantibody-positive systemic lupus erythematosus (SLE) who are receiving standard therapy • ~~adult~~ **patients aged 5 years and older with active lupus nephritis who are receiving standard therapy**
Stelara	Ustekinumab injection, 45 mg/0.5 mL and 130 mg/26 mL; 45 mg/0.5 mL and 90 mg/mL prefilled syringes	Janssen	Human interleukin-12 and -23 antagonist indicated for the treatment of: Adult patients with: • moderate to severe plaque psoriasis (Ps) who are candidates for phototherapy or systemic therapy • active psoriatic arthritis (PsA) • moderately to severely active Crohn's disease (CD) • moderately to severely active ulcerative colitis Pediatric patients 6 years and older with: • moderate to severe plaque psoriasis, who are candidates for phototherapy or systemic therapy • **active psoriatic arthritis (PsA)**
Myfembree	relugolix 40 mg, estradiol 1 mg, and norethindrone 0.5 mg oral tablets	Myovant sciences	Myfembree is a combination of relugolix, a gonadotropin-releasing hormone (GnRH) receptor antagonist, estradiol, an estrogen, and norethindrone acetate, a progestin, indicated in premenopausal women for the: • management of heavy menstrual bleeding associated with uterine leiomyomas (fibroids) • **management of moderate to severe pain associated with endometriosis**
Nubeqa	Darolutamide 300 mg oral tablet	Bayer healthcare	Nubeqa is an androgen receptor inhibitor indicated for the treatment of adult patients with: • non-metastatic castration-resistant prostate cancer (nmCRPC) • **metastatic hormone-sensitive prostate cancer (mHSPC) in combination with docetaxel**

Brand Name	Generic Name/ Dosage Form	Manufacturer	Newly Approved Indication†
Xofluza	baloxavir marboxil 40, 80 mg oral tablet	Genentech	An influenza virus polymerase acidic (PA) endonuclease inhibitor indicated for: • Treatment of acute uncomplicated influenza in patients who have been symptomatic for no more than 48 hours and who are: o otherwise healthy adults and pediatric patients ~~12~~ **5** years of age and older, OR o adults and pediatric patients 12 years of age and older who are at high-risk of developing influenza-related complications. • Post-exposure prophylaxis of influenza in patients 5 years of age and older following contact with an individual who has influenza
Enhertu	fam-trastuzumab deruxtecan-nxki 100 mg vial	Daiichi Sankyo	HER2-directed antibody and topoisomerase inhibitor conjugate indicated for the treatment of: • adult patients with unresectable or metastatic HER2-positive breast cancer who have received a prior anti-HER2-based regimen either: o in the metastatic setting, or o in the neoadjuvant or adjuvant setting and have developed disease recurrence during or within six months of completing therapy • **adult patients with unresectable or metastatic HER2-low (IHC 1+ or IHC 2+/ISH-) breast cancer who have received a prior chemotherapy in the metastatic setting or developed disease recurrence during or within 6 months of completing adjuvant chemotherapy.** • **adult patients with unresectable or metastatic non-small cell lung cancer (NSCLC) whose tumors have activating HER2 (ERBB2) mutations, as detected by an FDA-approved test, and who have received a prior systemic therapy** o **This indication is approved under accelerated approval based on objective response rate and duration of response. Continued approval for this indication may be contingent upon verification and description of clinical benefit in a confirmatory trial** • adult patients with locally advanced or metastatic HER2-positive gastric or gastroesophageal junction adenocarcinoma who have received a prior trastuzumab-based regimen

Brand Name	Generic Name/ Dosage Form	Manufacturer	Newly Approved Indication†
Mirena	levonorgestrel-releasing intrauterine system	Bayer Healthcare	Mirena is a progestin-containing intrauterine system (IUS) indicated for: Prevention of pregnancy for up to ~~7~~ **8** years • Treatment of heavy menstrual bleeding for women who choose to use intrauterine contraception as their method of contraception for up to 5 years

RECALLS

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Nifedipine WSP 0.2% Ointment, 60 gram tubes, Rx only, Valor Compounding Pharmacy, 2461 Shattuck Ave., Berkeley, CA 94704	Class I	D-1286-2022	Class I	Drugs	Lot #: 06162022@30, BUD 12/13/2022
Insulin Glargine (Insulin glargine-yfgn) Injection, 100 units/mL (U-100), 3 mL prefilled pens (NDC 49502-394-71), packaged in cartons of 5 prefilled pens (NDC 49502-394-75), Rx only, Manufactured by: Mylan Pharmaceuticals Inc., Morgantown, WV 15317, Manufactured for: Mylan Specialty L.P., Morgantown, WV	Class I	Drugs	Lot #: BF21002895, Exp. Date Aug 2023	Labeling: Missing label: Label missing from some prefilled pens.	Mylan Pharmaceuticals Inc
Propofol Injectable Emulsion, 1 g/100 mL (10 mg/mL), packaged in 100 mL per glass fliptop vial (NDC 0409-4699-54) further packaged in a tray of 10 vials (NDC Carton: 0409-4699-24), Rx only, Distributed by Hospira, Inc., Lake Forest, IL 60045 USA	Class I	Drugs	Lot #: DX9067, Exp 5/1/2023	Presence of particulate matter: particulate identified as a beetle.	Pfizer Inc.
SUSTANGO (Pendenadril Tytrate Blend) Capsules, 400 mg, 10-count blisters per carton, Formulated by: Male FX Labs, Bangor, ME, ASIN X0024468I9.	Class I	Drugs	Lot DAP272109, Exp: 4/1/2026	Marketed Without an Approved NDA/ANDA: Analytical testing showed the presence of tadalafil.	Ultra Supplement LLC
SANGTER Energy Supplement Capsules, 3000mg x 7 grain, 7-count blister pack within a carton, Distributed by: Distributor RFR, LLC, (800) 519-0204 Miami 33172 FL, USA; UPC 0 705632 523285	Class I	Drugs	Lot: 48656, Exp. 01/2025	Marketed Without An Approved NDA/ANDA: FDA analysis found the product to be tainted with sildenafil.	Distributor RFR, LLC.

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
M Hand Sanitizer Ethyl Alcohol Antiseptic 80%v/v, Topical Solution 128 oz/3,785 mL, Made in Mexico by: Grupo Plast-Y-Kosas S.A. de C.V. Puebla 105 Col. Rodriguez Reynosa, Tam, Mexico C.P. 88630, NDC Code: 77797-001-01, Distributed by: Medek, LLC 315 E. Business Hwy 83 Alamo, TX 78516, NDC Code: 75432-001-02.	Class I	Drugs	All lots	Chemical Contamination and Subpotent Drug: FDA analysis found product to contain methanol and below label claim for ethanol.	MEDEK LLC
Divalproex Sodium Delayed-Release Tablets, USP 500mg, Rx Only, 100 Tablets, Distributed by: Sun Pharmaceutical Industries, Inc., Cranbury, NJ 08512, Manufactured by: Sun Pharmaceutical Industries, Ltd., Halol-Baroda Highway, Halol- 389 350, Gujarat, India, NDC 62756-798-88.	Class II	Drugs	Lot: HAC1312A, EXP. 05/2024	Failed Dissolution Specifications: Failure occurred during routine stability testing of dissolution test.	SUN PHARMACEUTICAL INDUSTRIES INC
Testosterone Gel 1% (25mg testosterone/2.5g of gel) 2.5 g per unit dose, Rx Only, 30 unit-dose packets per box. Manufactured by: Actavis Laboratories UT, Inc., Salt Lake City, US 84108, USA, Distributed by: Actavis Pharma, Inc., Parsippany, NJ 07054, USA, Sachet NDC 0591-3216-17, Carton NDC 0591-3216-30	Class II	Drugs	Lot: 1403180, EXP. 10/2022	Superpotent Drug: Out of specification assay result was obtained during stability testing.	Teva Pharmaceuticals USA Inc
Lansoprazole Delayed-Release Orally Disintegrating Tablets, 15 mg, 100-count blisters per carton, 10 Packs of 10 Tablets Each, Rx Only, Distributed by: Dr. Reddy's Laboratories, Inc., Princeton, NJ 08540,	Class II	Drugs	Lot T2100514, Exp 01/2023	FAILED DISSOLUTION SPECIFICATIONS	Dr. Reddy's Laboratories, Inc.

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Made in India, blister barcode 4359856079, NDC 43598-560-78					
Lansoprazole Delayed-Release Orally Disintegrating Tablets, 35 mg, 100-count blisters per carton, 10 Packs of 10 Tablets Each, Rx Only, Distributed by: Dr. Reddy's Laboratories, Inc., Princeton, NJ 08540, Made in India, blister barcode 4359856179, NDC 43598-561-78	Class II	Drugs	Lot T2100515, Exp 01/2023	FAILED DISSOLUTION SPECIFICATIONS	Dr. Reddy's Laboratories, Inc.
Irbesartan Tablets, USP, 150mg, 90- count bottles, Rx only, Manufactured by: Jubilant Generics Ltd. Roorkee- 247661, India, Marketed by: Jubilant Cadista Pharmaceuticals Inc. Salisbury, MD 21801, USA, NDC 59746-448-90	Class II	Drugs	Lot #: IB220023A, exp. date 08/2022	Failed dissolution specifications.	Jubilant Cadista Pharmaceuticals, Inc.
Irbesartan Tablets, USP, 75 mg, 90- count bottle, Rx only, Manufactured by: Jubilant Generics Ltd. Roorkee- 247661, India, Marketed by: Jubilant Cadista Pharmaceuticals Inc. Salisbury, MD 21801, USA, NDC 59746-447-90	Class II	Drugs	Lot #: IB120012A, IB120013A, IB120014A Exp. date 08/2022	Failed Dissolution Specifications.	Jubilant Cadista Pharmaceuticals, Inc.
Naftifine Hydrochloride Gel, USP 1%, packaged in cartons with: a) Net Wt. 90g tube (NDC 0115-1510-48), b) Net Wt. 40g tube (NDC 0115-1510-63), c) Net Wt. 60g tube (NDC 0115-1510-58), Rx only, Manufactured by: Tolmar, Inc., Fort Collins, CO, 80526, Distributed by: Amneal Pharmaceuticals LLC, Bridgewater, NJ 08807	Class II	Drugs	Lot #: a) 12070A, Exp 5/2023; 11801A, Exp 9/2022; b) 12386A, Exp 8/2023; 11800A, Exp 9/2022; c) 11940A, Exp 12/2022	Failed Impurities/ degradation specifications: Out-of-Specification test results obtained for Unspecified Impurity Relative Retention Time	Tolmar, Inc.

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Matzim LA (Diltiazem Hydrochloride) Extended-Release Tablets, 180 mg, 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 52544-691-30.	Class II	Drugs	Lot 1411593A, Exp 09/22	Failed Dissolution Specifications: below specification limits for dissolution.	Teva Pharmaceuticals USA Inc
Matzim LA (Diltiazem Hydrochloride) Extended-Release Tablets, 240 mg, 30-count bottles, Rx Only, Manufactured by: Actavis Laboratories FL, Inc., Fort Lauderdale, FL USA 33314, USA, Distributed by: Actavis Pharma Inc., Parsippany, NJ 07054 USA; NDC 58544-692-30.	Class II	Drugs	Lot 1411596A, Exp 09/22	Failed Dissolution Specifications: below specification limits for dissolution.	Teva Pharmaceuticals USA Inc
Telmisartan and Hydrochlorothiazide Tablets USP, 80 mg/25 mg, 30 Tablets, Rx Only, Manufactured by: Glenmark Pharmaceuticals Ltd., Plot no 2, Phase-2, Pharma Zone, SEZ Pithampur, District Dhar, Madhya Pradesh 454775, India, Manufactured for: Glenmark Pharmaceuticals Inc, USA, Mahwah, NJ 07430, NDC 68462-842-13.	Class II	Drugs	Lots 17210935 & 17210936., Exp Date 05/2023 Lot 17211206, Exp Date 06/2023 Lots 17211652, 17211655 & 17211658, Exp Date 08/2023	Packaging : Blister package issues.	Glenmark Pharmaceuticals Inc., USA
Telmisartan and Hydrochlorothiazide Tablets USP, 80 mg/12.5 mg, 30 Tablets, Rx Only, Manufactured by: Glenmark Pharmaceuticals Ltd., Plot no 2, Phase-2, Pharma Zone, SEZ Pithampur, District Dhar, Madhya Pradesh 454775, India, Manufactured for: Glenmark	Class II	Drugs	Lots 17210929 & 17210930, Exp Date 05/2023; Lot 17211203, Exp Date 06/2023 & Lots 17211643, 17211646 &	Packaging : Blister package issues.	Glenmark Pharmaceuticals Inc., USA

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Pharmaceuticals Inc,m USA, Mahwah, NJ 07430, NDC 68462-841-13.			17211649, Exp Date 08/2023		
Triple Antibiotic Ointment, Bacitracin zinc, Neomycin sulfate, Polymixin B sulfate, First Aid Antibiotic, Triple Antibiotic Ointment, 144 packets per box, Net wt. per packet 0.5 g, Honeywell Safety Products, NDC 0498-0750-36.	Class II	Drugs	Part# 231209G	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
Bisacodyl Suppositories, Fast Acting Stimulant Laxative, 100 suppositories per box, 10 mg each, Health Star, NDC 57896-443-01.	Class II	Drugs	Part# 444-01-HST	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
Naphcon A eye drops, Naphazoline HCl 0.025% and Pheniramine Maleate 0.3%, Redness Reliever and Antihistamine Eye Drops, Sterile, 15 mL (0.5 FL OZ) bottle per box, Alcon, a Novartis company, NDC 0065-0085-15.	Class II	Drugs	Part# 0065008515	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
Systane, Lubricant Eye Drops, Polyethylene Glycol 400 0.4% Lubricant, Propylene Glycol 0.3% Lubricant, Original, Long Lasting Dry Eye Relief, Sterile, a) 15 mL (0.5 FL OZ) NDC 0065-0429-15, b) 30 mL (1 FL OZ) NDC 0065-0429-30, bottle per box, Alcon Surgical Inc.	Class II	Drugs	Part# a) 0065042915, b) 0065042930	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
Eye-stream, eye wash solution, sterile, 4 FL OZ (118 mL) bottle per box, Alcon, NDC 0065-0530-04.	Class II	Drugs	Part# 0065053004	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Systane Balance, Lubricant Eye Drops, Propylene Glycol 0.6% lubricant, Restorative Formula, Sterile, 10 mL (1/3 FL OZ) bottle per box, Alcon, NDC 0065-1433-02.	Class II	Drugs	Part# 0065143302	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
Systane Zaditor, ketotifen fumarate ophthalmic solution 0.035%, Antihistamine eye drops, Eye Itch Relief, up to 12 Hours, Sterile, 30 day supply, 5mL (0.17 FL OZ) bottle per box, Alcon, NDC 0065-4011-05.	Class II	Drugs	Part# 0065401105	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
Debrox, Carbamide Peroxide, Earwax Removal Aid, 0.5 FL OZ (15 mL) bottle per box), MedTech Products Inc., NDC 63029-321-01.	Class II	Drugs	Part# 04203710478	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
Miralax (Polyethylene Glycol 3350), Powder for Solution, Osmotic Laxative, 30 Once-Daily Doses, Net WT 17.9 OZ (510 g) bottle, Bayer Healthcare Pharmaceuticals, NDC 11523-7234-4.	Class II	Drugs	Part# 11523723404	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
GenTeal Tears, Lubricant Eye Ointment, Night-Time Ointment, Sterile, 3.5 gm (0.12 FL OZ) per box, Alcon, NDC 0065-0518-01.	Class II	Drugs	Part# 30065051801	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
Pataday, Once Daily Relief, Olopatadine hydrochloride ophthalmic solution 0.2%, Antihistamine, Eye Allergy Itch Relief, Once Daily, Sterile, 2.5 mL (0.085 FL OZ) bottle per box, Alcon, NDC 0065-8150-01.	Class II	Drugs	Part# 00065815001	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
A&D Original Ointment, Diaper Rash Ointment & Skin Protectant, 16 oz. Jar, Bayer Healthcare Pharmaceutica, NDC 11523-0096-3.	Class II	Drugs	NA	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
Dakin's Solution, a) sodium hypochlorite 0.125%, quarter strength, NDC 0436-0672-16, b) sodium hypochlorite 0.25%, half strength, NDC 0436-0936-16, c) sodium hypochlorite 0.5% full strength, NDC 0436-0946-16, Antimicrobial, 473 mL (16 fl oz) bottle, Century Pharmaceuticals, Inc.	Class II	Drugs	NA	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
Asthmanefrin Racephinephrine Inhalation Solution Bronchodilator, For temporary relief of mild symptoms of intermittent asthma, Preservative Free, Sterile, For Oral Inhalation Only, 30 vials per box, Nephron Pharmaceuticals Corporation, NDC 0487-2784-01.	Class II	Drugs	NA	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
Racepinephrine Inhalation Solution, USP 2.25%, Bronchodilator, For Oral Inhalation Only, Sulfite Free, Preservative Free, 30 x 0.5 mL Sterile Unit-of-Use Vials, each in a foil pouch, per carton, Nephron Pharmaceuticals Corporation, NDC 0487-5901-99.	Class II	Drugs	NA	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
Sterile Alcohol Prep Pads, Sterile, Latex Free, 100 large pads per box, manufactured for: Dynarex Corporation, NY 10962, NDC 67777-121-16.	Class II	Drugs	Part# 1116	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Alcohol Swabsticks, Antiseptic, 50 4" saturated individual packets per box, Manufactured for: Dynarex Corporation, NY 10962, Made in Mexico, NDC 67777-300-01 (current NDC) NDC# 67777-120-10 (discontinued)	Class II	Drugs	Part# 1203	CGMP Deviations: products were stored outside the drug label specifications.	Mckesson Medical-Surgical Inc. Corporate Office
Lorazepam Injection, USP, 2mg/mL, 1 mL vial, 25 vials per carton, RX Only, Manufactured by West-Ward Eatontown, NJ 07724, Carton NDC# 0641-6044-25, Vial NDC# 0641-6044-01	Class II	Drugs	Lot # 060064, Exp. 06/2023, 070084, Exp. 07/2023, 070126, Exp. 07/2023, 080091, Exp. 08/2023, 080060, Exp. 08/2023	Failed Impurities/Degradation Specifications: Out-of-specification results observed for total related compounds during testing of retain samples.	Hikma Pharmaceuticals USA Inc.
Ativan Injection (lorazepam injection, USP), 2mg/mL, 1 mL vial, 25 vials per carton, RX Only, Manufactured by West-Ward Eatontown, NJ 07724, Carton NDC# 0641-6001-25, Vial NDC# 0641-6001-01	Class II	Drugs	Lot # 060064Z, Exp. 06/2023	Failed Impurities/Degradation Specifications: Out-of-specification results observed for total related compounds during testing of retain samples.	Hikma Pharmaceuticals USA Inc.
Lorazepam Injection, USP, 2mg/mL, 1 mL vial, 25 vial per carton, Rx Only, Novaplus, Manufactured by Hikma Berkeley Heights, NJ 07922. Carton NDC# 0641-6048-25, Vial NDC# 0641-6048-01	Class II	Drugs	Lot # 070088, exp. date 07/2023	Failed Impurities/Degradation Specifications: Out-of-specification results observed for total related compounds during testing of retain samples.	Hikma Pharmaceuticals USA Inc.
Lorazepam Injection, USP, 4mg/mL, 1 mL vial, 25 vial per carton, Rx Only, Novaplus, Manufactured by Hikma Berkeley Heights, NJ 07922. Carton NDC# NDC# 0641-6045-25, Vial NDC# 0641-6045-01	Class II	Drugs	Lot # 070096, exp. date 07/2023	Failed Impurities/Degradation Specifications: Out-of-specification results observed for total related compounds during testing of retain samples.	Hikma Pharmaceuticals USA Inc.
Prednisone Tablets USP, 20 mg, 100-count bottle, Rx Only, Manufactured by: Strides Pharma Science Ltd., Bengaluru - 562106,	Class II	Drugs	Lot #: 7248988B, Exp 9/2023	Presence of foreign tablet: 2.5 mg tablet in a 20 mg bottle of Prednisone Tablets	Strides Pharma Inc.

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
India, Distributed by: Strides Pharma Inc., East Brunswick, NJ 08816, NDC 64380-785-06.					
FentaNYL Citrate in 0.9% Sodium Chloride 1 mg per 100 mL (10 mcg per mL) IV bags, Rx only, SterRx, 141 Idaho Ave., Plattsburgh, NY, 12903, NDC 70324-327-01.	Class II	Drugs	Lot #: CHI, Exp. Date Dec 22, 2022	Lack of Assurance of Sterility	SterRx, LLC
fentaNYL Citrate in 0.9% Sodium Chloride, 2.5 mg per 250 mL, (10 mcg per mL) IV bags, Rx Only, SterRx, 141 Idaho Avenue, Plattsburgh, NY 12903, NDC 70324-327-02.	Class II	Drugs	Lot #: CHL, Exp. Date Dec 29, 2022	Lack of Assurance of Sterility	SterRx, LLC
Difluprednate Ophthalmic Emulsion 0.05%, For Ophthalmic Use Only, Sterile, 5 mL bottles, Manufactured by Cipla Ltd., India, Manufactured for: Cipla USA, Inc., NJ 07059, NDC 69097-341-35.	Class II	Drugs	Lot # DEG3LC2, Exp 05/2023	Lack of Assurance of Sterility: Complaints received of defective container closure.	CIPLA
Rifampin Capsules, USP, 150 mg, 30 count HDPE bottles, Rx Only, Manufactured for Lupin Pharmaceuticals, Inc., Baltimore, MD, 21202 NDC: 68180-658-06	Class II	Drugs	Lot #A200170, exp. date December 2023	CGMP Deviations:OOS result was observed in 1-Methyl-4-Nitroso Piperazine (MNP) impurity.	Lupin Pharmaceuticals Inc.
Fulvestrant Injection 250mg/5mL (50 mg/mL), Contains 2 Single-Dose Prefilled Syringes, Rx Only, Product of India, Manufactured by: Cadila Healthcare Limited, Ahmadabad, India, Distributed by: Zydus Pharmaceuticals (USA) Inc.m Pennington, NJ 08534, NDC 70710-1688-8.	Class II	Drugs	Lot # B200076; Exp 31 JAN 2024	Failed Impurities/Degradation Specifications	Zydus Pharmaceuticals (USA) Inc
Avant Foaming Hand Sanitizer, ethanol 62%, a) 18 fl. oz. bottle (530 mL), b) 1000 mL (33.9 fl. oz.) pouch, c) 55 gallon drum, Fragrance	Class II	Drugs	[Product number], lot code, expiry: a) [46016] Lot 722770, exp 07/22 b)	CGMP Deviations: product manufactured using deionized water from a system lacking	Aire-Master of America Inc

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Free, Manufactured for B4 Brands, Lisbon, Iowa 62253			[46017] Lot 722995-exp 07/22, 727370-exp 07/23, 727898-exp 09/23, 728005-exp 08/23, 729522-exp12/23, 730412-exp 03/24, 730861-exp 04/24 c) [46138] Lot 725188-exp 12/22	appropriate microbial control post deionization.	
Avant Foaming Hand Sanitizer Ophardt, Fragrance Free, 1000 mL (33.9 fl. oz.) per plastic carton, Manufactured for B4 Brands, Lisbon, Iowa 52253	Class II	Drugs	Product Number 46076 Lot 722896-exp 07/22, 722996-exp 07/22	CGMP Deviations: product manufactured using deionized water from a system lacking appropriate microbial control post deionization.	Aire-Master of America Inc
Stage 2-Ophardt Foaming Hand Sanitizer, Fragrance-Free, 1000 mL (33.9 fl. oz.) per bottle, Manufactured for 2XL Corporation, 2 Gateway Ct, Ste A, Bolingbrook, IL 60440	Class II	Drugs	Product Number 46101, Lots 722712-exp 07/22, 724755-exp 11/22, 725054-exp 12/22	CGMP Deviations: product manufactured using deionized water from a system lacking appropriate microbial control post deionization.	Aire-Master of America Inc
Protect U Guard Foaming Hand Sanitizer Ophardt, Fragrance Free, 1000 mL (33.9 fl. oz.) per carton, Manufactured for Protect U Guard, Tampa 33606.	Class II	Drugs	Product Number 46112, Lots 722782 -exp 08/22	CGMP Deviations: product manufactured using deionized water from a system lacking appropriate microbial control post deionization.	Aire-Master of America Inc
Protect U Guard Foaming Hand Sanitizer, Fragrance Free, 18 fl/oz. (530 mL) per bottle, Manufactured for Protect U Guard, Tampa, FL 33606	Class II	Drugs	Product Number 46111, Lot 722781-exp 09/22	CGMP Deviations: product manufactured using deionized water from a system lacking appropriate microbial control post deionization.	Aire-Master of America Inc

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Common Sense Fragrance Free Hand Sanitizer, 250 Gallon Tote, Microbe Solutions, LLC, 344-5 Route 9, Suite 237, Lanoka Harbor, NJ 08734	Class II	Drugs	Product Number 46137, Lot 724640-exp 11/22, 729955-exp 01/24	CGMP Deviations: product manufactured using deionized water from a system lacking appropriate microbial control post deionization.	Aire-Master of America Inc
PrednisoLONE Oral Solution USP, 15 mg per 5 mL, 240 ml bottle, Rx Only, HI-TECH PHARMACAL CO., INC., Amityville, NY 11701. NDC: 50383-042-24	Class II	Drugs	Lot# 379804, Exp. 8/31/2023	Defective Container: Product has incomplete induction seals.	Akorn, Inc
Acetaminophen Injection, 10 mg/mL, 1,000 mg/100 mL, 100 mL VIAFLO container bag, Single Dose Container, For Intravenous Use Only, Rx Only, Baxter Healthcare Corporation, Deerfield, IL 60015 USA; NDC 36000-306-60	Class II	Drugs	Lots: 20K19G64T1, Exp 10/31/2022; 21K23G65, 21K25G65, 21K26G65, 21K29G67, Exp 10/31/2023; 21L10G65, 21L13G66, 21L14G66, 21L15G65, Exp 11/30/2023	Temperature Abuse: Product distributed in refrigerated trucks with labels attached to pallets indicating "Refrigerate Upon Arrival", however product is labeled to be stored in a controlled room temperature environment.	Baxter Healthcare Corporation
Ketorolac Tromethamine Injection, USP 60 mg/2 mL (30 mg/mL), packaged in 2 mL single dose vials, Rx Only, Nephron Pharmaceutical Corporation 4500 12th Street Extension West Columbia, SV 29172, NDC 0487-6232-01	Class II	Drugs	Lot#: 023011 Exp 8/31/2022	cGMP Deviations: deviations leading to potential cross-contamination.	Nephron Sc Inc
PF-Neostigmine Methylsulfate Injection, USP, 3 mg/3 mL (1 mg/mL), One 3 mL Unit-Dose Vial, packaged in 30 x 3 mL Sterile Unit-Dose Vials per carton, Rx Only, Nephron 503B Outsourcing Facility, 4500 12th Street Extension, West Columbia, SC 29172; NDC 69374-932-33	Class II	Drugs	Lot: NE1057A, Exp. 10/23/2022	CGMP Deviations: Potential for cross-contamination due to product carryover during manufacturing.	Nephron Sterile Compounding Center LLC

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Trisodium Citrate 0.5% Solution, (0.5%/4L), contains Per Liter: Sodium 140 mmol/L, Chloride 86 mmol/L, Citrate 18 mmol/L, 4000 mL IV bag, packaged in 1 x 1 IV bag per carton, Rx Only, Nephron 503B Outsourcing Facility, 4500 12th Street Extension, West Columbia, SC 29172; NDC 69374-910-04	Class II	Drugs	Lots: TC2007D, Exp. 8/26/2022; TC2010A, Exp. 9/25/2022	CGMP Deviations: Potential for cross-contamination due to product carryover during manufacturing.	Nephron Sterile Compounding Center LLC
PF-0.125% Bupivacaine HCl Injection, USP, 625 mg/500 mL (1.25 mg/mL), 500 mL bag, packaged in 10 x 1 IV Bag per case, Rx Only, Nephron 503B Outsourcing Facility, 4500 12th Street Extension, West Columbia, SC 29172; NDC 69374-970-05	Class II	Drugs	Lots: BH2003A, Exp. 8/19/2022; BH2011A, Exp. 11/9/2022	CGMP Deviations: Potential for cross-contamination due to product carryover during manufacturing.	Nephron Sterile Compounding Center LLC
PF-Labetalol HCl Injection, USP, 20 mg/4 mL (5 mg/mL), One 4 mL Unit-Dose Vial, packaged in 30 x 4 mL Sterile Unit-dose Vials per carton, Rx Only, Nephron 503B Outsourcing Facility, 4500 12th Street Extension, West Columbia, SC 29172; NDC 69374-946-34	Class II	Drugs	Lot: LB2005A, Exp 3/2/2023	CGMP Deviations: Potential for cross-contamination due to product carryover during manufacturing.	Nephron Sterile Compounding Center LLC
Juice Beauty, The Organic Solution, SPF 8, Joyful Lip Moisturizer, Hydratant pour les Levres, NET WT 0.15 oz (4.25 g) per tube, Active Ingredient: Zinc oxide 4%, Manufactured for Juice Beauty, Inc., 709 Fifth Ave., San Rafael, California 94901-3566	Class III	Drugs	Lot#: 90121-20321 Exp: 11/2023	Failed Dissolution Specifications	Eco Lips, Inc
Juice Beauty, The Organic Solution, SPF 8, Naturally Clear Lip Moisturizer, Hydratant pour les Levres, NET WT 0.15 oz (4.25 g) per tube, Active Ingredient: Zinc oxide 4%,	Class III	Drugs	Lot #: 90122-21015, Exp: 01/24; Lot#: 90122-20356, Exp: 12/23	Failed Dissolution Specifications	Eco Lips, Inc

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
Manufactured for Juice Beauty, Inc., 709 Fifth Ave., San Rafael, California 94901-3566					
Crest 3D White Fluoride Anticavity Toothpaste, Advanced Triple Whitening, 0.243% sodium fluoride, Net Wt. 5.6 oz (158 g) a) Individual carton, UPC 0 37000 598534; b) 5-count Bundle, UPC 037000171867, Distributed. By Procter & Gamble, Cincinnati, OH 45202.	Class III	Drugs	Lot: a) 13191707B4, Exp. 10/31/2023; b)13511707Y1, Exp. 10/31/2023.	Labeling: Missing label: the product tube was missing a label and contained a different formulation.	The Procter & Gamble Company
Tacrolimus Ointment, 0.1%, For Dermatological Use Only, Not for Ophthalmic Use, Rx Only, a) 30 g tube, NDC 68462-534-35, b) 60 g tube, NDC 68462-534-65, c) 100 g tube, NDC 68462-534-94, Manufactured for: Glenmark Pharmaceuticals Inc., USA, Mahwah, NJ 07430, Product of India.	Class III	Drugs	Lot #s: a) 15200058, 15200066, 15200067, Exp 06/2022; 15200075, Exp 07/2022; 15200094, Exp 08/2022; 15200119, 15200120, 15200122, Exp 09/2022; 15200131, 15200132, Exp 10/2022; 15200133, 15200134, 15200136, 15200139, Exp 11/2022; 15210033, 15210035, 15210036, 15210038 Exp 01/2023; 15210092, 15210094, 15210097, 15210099, Exp 03/2023; 15210100, 15210101, 15210104, Exp 04/2023; 15210122, Exp 05/2023; 15210148, 15210149, 15210153, 15210155, 15210157,	Defective Container: Tube split from side seam	Glenmark Pharmaceuticals Inc., USA

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			15210158, Exp 06/2023; 15210173, 15210174, 15210176, 15210177, 15210178, 15210179, Exp 07/2023; 15210184, 15210186, Exp 08/2023; 15210214, Exp 09/2023; 15210225, 15210226, 15210228, 15210230, 15210231, Exp 10/2023; 15220001, 15220002, Exp 12/2023. b) 15200092, Exp 08/2022; 15200114, 15200115, Exp 09/2022; 15200123, 15200124, Exp 10/2022; 15200142, 15200144, Exp 11/2022; 15210014, 15210015, 15210021, Exp 12/2022; 15210031, Exp 01/2023; 15210055, Exp 02/2023; 15210073, 15210075, Exp 03/2023; 15210117, 15210118, Exp 04/2023; 15210144, 15210147, 15210160, 15210162, Exp 06/2023; 15210171, Exp 07/2023; 15210187, 15210190, 15210191, 15210193, Exp 08/2023;		

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			15210212, Exp 09/2023; 15210244, 15210247, Exp 11/2023; 15220007, Exp 12/2023; 15220019, 15220028, Exp 01/2024; 15220032, 15220034, 15220053, Exp 02/2024; 15220062, 15220064, 15220072, Exp 03/2024. c) 15200065, Exp 06/2022; 15200108, 15200109, Exp 09/2022; 15200125, Exp 10/2022; 15200148, 15200149, 15200150, Exp 11/2022; 15210007, 15210009, 15210012, 15210018, Exp 12/2022; 15210049, Exp 02/2023; 15210076, 15210077, 15210089, Exp 03/2023; 15210112, 15210114, Exp 04/2023; 15210124, 15210125, 15210127, Exp 05/2023; 15210169, 15210170, Exp 07/2023; 15210196, 15210199, Exp 08/2023; 15210204, 15210206, 15210207, 15210210, Exp 09/2023; 15210248, Exp 11/2023;		

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			15220004, 15220013, 15220014, 15220015, Exp 12/2023; 15220025, 15220026, Exp 01/2024.		
Azacitidine for Injection 100mg/vial Lyophilized Powder, Rx Only, Mfd. in Romania By: Sindan Pharma SRL For BluePoint Laboratories, NDC 68001-313-56	Class III	Drugs	Lot: FE22001A, Exp 01/2024	Subpotent Drug - Out of specification (OOS) result obtained during monitoring stability study for Assay. Results below specification.	Teva Pharmaceuticals USA Inc
Lamotrigine Tablets, USP 100 mg, 1000-count bottles, Rx Only, Manufactured for: AvKARE, Inc. Pulaski, TN 38478, NDC 42291-367-10, UPC 3 42291 36710 4	Class III	Drugs	Lot #: 42581 Exp. 12/2024; 42484 Exp. 11/2024; 41204 Exp. 05/2024; 38723 Exp. 02/2023; 37623 Exp. 10/2022	Labeling: Label Error on Declared Strength	AVKARE Inc.
Azacitidine for Injection, 100 mg Lyophilized Powder, Single-Dose Vials, Rx Only, Manufactured by: Intas Pharmaceuticals Limited, Pharmez, Ahmedabad-382 213 India. Manufactured for BluePoint Laboratories. NDC 68001-0313-56	Class III	Drugs	Lot #: FE22001A, Exp. Date 01/2024	Subpotent Drug	Amerisource Health Services LLC
Acetaminophen 325 mg tablets, packaged in a) 100-count bottle (NDC 71399-8024-01); b) 1000-count bottle (NDC 71399-8024-02), Akorn Pharma Manufactured for: Akorn Pharma, Inc., Fairfield, NJ	Class III	Drugs	Lot #: AXA2001, AXA2002, AXA2003, AXA2004, AXA2005, AXA2006, AXA2007, AXA2008, AXA2009, AXA2010, AXA2011, AXA2012, AXA2013, AXA2014, Exp Feb-23; AKK2021, AKK30421, AKK40421, AKK50421, AKK60421,	Failed Tablet/Capsule Specifications: Imprint "AP 325" is missing from the tablet.	Akron Pharma, Inc.

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			AKK70421, AKK80421, AKK90421, Exp Mar-24; AKL10421, AKL20421, AKL10521, AKL20521, AKL30521, AKL40521, AKL50521, AKL60521, AKL70521, AKL80521, AKL90521, Exp Apr-24; AKM10521, Exp Apr-24; AKA10621, AKA20621, AKA30621, AKA40621, AKA50621, AKA60621, AKA70621, AKA80621, AKA90621, Exp May-24; AKB10621, Exp May-24		
Acetaminophen 500 mg tablet, Extra Strength, packaged in a) 100-count bottle (NDC 71399-8022-01), b) 1000-count bottle (NDC 71399-8022-02), Akorn Pharma Manufactured for: Akorn Pharma, Inc., Fairfield, NJ	Class III	Drugs	Lot#: AXA2014, Exp Feb-23; AXB2001, Exp Nov-22; AXB2002, AXB2003, AXB2004, AXB2005, AXB2006, AXB2007, AXB2008, AXB2009, AXB2010, AXB2011, Exp Dec-22; AXB2012, AXB2013, AXB2014, AXB2015, AXB2016, AXB2017, AXB2018, AXB2019, AXB2020, AXB2021, AXB2022, AXB2023, AXB2024,	Failed Tablet/Capsule Specifications: Imprint "AP 325" is missing from the tablet.	Akron Pharma, Inc.

Product Description	Classification	Product Type	Code Info	Reason for recall	Recalling Firm
			AXB2025, AXB2026, AXB2027, Exp Feb-23		
Zynrelef (bupivacaine and meloxicam), 400 mg bupivacaine and 12 mg meloxicam single dose application, packaged in a kit, Rx only, Manufactured for Heron Therapeutics, Inc., San Diego, CA, NDC 47426-301-02	Class III	Drugs	Lot #: 01126739, Exp 7/31/2023	Defective Delivery System: An incorrect 10 mL (12 mL) Luer (slip) syringe packaged in one lot of Zynrelef 400 mg/12 mg kit	HERON THERAPEUTICS, 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

Amifostine Injection
Amino Acids
Amoxapine 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 Disodium Versenate Injection
Calcium Gluconate Injection
Cefazolin Injection
Cefixime Oral Capsules
Cefotaxime Sodium Injection
Cefotetan Disodium Injection
Chlordiazepoxide Hydrochloride Capsules
Chloroprocaine Hydrochloride Injection
Conivaptan Hydrochloride (Vaprisol) in 5% Dextrose Plastic Container
Continuous Renal Replacement Therapy (CRRT) Solutions
Cortisone Acetate Tablets
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
Digoxin Injection
Diltiazem Hydrochloride Injection

Disopyramide Phosphate (Norpace) Capsules
Dobutamine Hydrochloride Injection
Dopamine Hydrochloride Injection
Echothiophate Iodide (Phospholine Iodide) Ophthalmic Solution
Enalaprilat Injection
Epinephrine Injection, 0.1 mg/mL
Epinephrine Injection, Auto-Injector
Erythromycin Ophthalmic Ointment
Fentanyl Citrate (Sublimaze) Injection
Floxuridine for Injection
Fludarabine Phosphate Injection
Fluorescein Injection
Flurazepam Hydrochloride Capsules
Fluvoxamine ER Capsules
Furosemide Injection
Gentamicin Sulfate Injection
Guanfacine Hydrochloride Tablets
Heparin Sodium and Sodium Chloride 0.9% Injection
Hydromorphone Hydrochloride Injection
Hydroxypropyl (Lacrisert) Cellulose Ophthalmic Insert
Ibutilide Fumarate Injection
Indigotindisulfonate Sodium Injection
Iodixanol Injection
Iohexol Injection
Iomeprol injection
Iopromide (Ultravist) Injection
Isoniazid Injection
Ketamine Injection
Ketoprofen Capsules
Ketorolac Tromethamine Injection
Leucovorin Calcium Lyophilized Powder for Injection
Leuprolide Acetate Injection
Lidocaine Hydrochloride (Xylocaine) and Dextrose Injection Solution-Premix Bags
Lidocaine Hydrochloride (Xylocaine) Injection
Lidocaine Hydrochloride (Xylocaine) Injection with Epinephrine
Lipid Injection
Lithium Oral Solution
Lorazepam Injection
Mannitol Injection
Mepivacaine Hydrochloride Injection
Methyldopa Tablets
Methylprednisolone Acetate Injection
Metronidazole Injection
Midazolam Injection
Morphine Sulfate Injection
Multi-Vitamin Infusion (Adult and Pediatric)

Nizatidine Capsules
Paclitaxel Injection (protein-bound particles)
Pantoprazole Sodium for Injection
Parathyroid Hormone (Natpara) Injection
Pentostatin Injection
Physostigmine Salicylate Injection
Potassium Acetate Injection
Potassium Chloride Concentrate Injection
Promethazine (Phenergan) Injection
Propofol Injectable Emulsion
Protamine Sulfate Injection
Remifentanil Injection
Rifampin Capsules
Rifampin Injection
Rifapentine Tablets
Ropivacaine Hydrochloride Injection
Semaglutide (Ozempic) Injection
Semaglutide (WEGOVY®) Injection
Sincalide (Kinevac) Lyophilized Powder for 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
Sterile Water for Injection
Streptozocin (Zanosar) Sterile Powder
Sufentanil Citrate Injection
Sulfasalazine Tablets
Technetium TC-99M Mebrofenin Injection
Technetium Tc99m Succimer Injection (DMSA)
Teprotumumab-trbw
Thiothixene Capsules
Triamcinolone Acetonide Injectable Suspension
Triamcinolone Hexacetonide Injectable suspension
Trimethobenzamide Hydrochloride Capsules
Valproate Sodium Injection
Vandetanib Tablets
Vecuronium Bromide for Injection