

Drug Recall List

Last Updated: July 2022

Drug	Recall Details	Contact	Date	Drug Recall Class*
Morphine Sulfate Extended-Release 60mg 63629108901	Labeling: Label Mix-up	If you have questions about this recall, Bryant Ranch Prepack, Inc, 1-877-885-0882	July 2022	Class I
Morphine Sulfate Extended-Release 30mg 63629108801	Labeling: Label Mix-up	If you have questions about this recall, Bryant Ranch Prepack, Inc, 1-877-885-0882	July 2022	Class I
Losartan Potassium 06438093305 06438093308 06438093405 06438093408 06438093505 06438093508	CGMP Deviations	If you have questions about this recall, Strides Inc.1-609-773-5000	June 2022	Class II
Losartan HCTZ 70518257800 68180021709 70518256400 68180021609 33342005010	CGMP Deviations	If you have questions about this recall, RemedyRepack Inc., 1-866-845-3791	June 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

70518323100				
Losartan Potassium & HCT 68788775809 68788775803	CGMP Deviations	If you have questions about this recall, Preferred Pharmaceuticals, Inc., 1-714-777-3729	June 2022	Class II
Vitamin D3 73198007530	Lack of assurance of sterility	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	June 2022	Class II
Testosterone Cypionate 73198005410 73198005505	Lack of assurance of sterility	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	June 2022	Class II
Daytrana (methylphenidate transdermal system) 06896855523	Defective Delivery Syst	If you have questions about this recall, Noven Pharmaceuticals Inc, 1-305-253-5099	June 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

EPINEPHrine 69374-544-10 69374-925-10	Lack of Assurance of Sterility	If you have questions about this recall, Nephron Sterile Compounding Center LLC, 1-800-443-4313	June 2022	Class II
Zonisamide 06191977590	CGMP Deviations	If you have questions about this recall, Direct Rx, 1-678-619-5510	June 2022	Class II
Losartan Pot/HCTZ 07218929090 07218929790 07218928990 07218916730 07218916790	CGMP Deviations	If you have questions about this recall, Direct Rx, 1-678-619-5510	June 2022	Class II
Trulicity (dulaglutide) 00002143480 00002143380	TEMPERATURE ABUSE	If you have questions about this recall, Eli Lilly & Company, 1-800-545-5979	June 2022	Class II
Humalog KwikPen 00002879959 00002751659	TEMPERATURE ABUSE	If you have questions about this recall, Eli Lilly & Company, 1-800-545-5979	June 2022	Class II
Xanax XR (alprazolam) extended release 00009006807	Failed Dissolution Specifications	If you have questions about this recall, Viatris Inc, 1-800.796.9526	May 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Accupril (Quinapril HCl Tablets) 00071053023 00071053223 00071053523	CGMP Deviations	If you have questions about this recall, Pfizer Inc., 1-800-438-1985	May 2022	Class II
GaviLyte -C (Polyethylene Glycol 3350, 240 g) 04338606019	Failed Stability Specification	If you have questions about this recall, Lupin Pharmaceuticals Inc. 1-410-576-2000 ext 3	May 2022	Class II
Trulicity (dulaglutide) 00002143480 00002143380	TEMPERATURE ABUSE	If you have questions about this recall, Eli Lilly & Company, 1-800-545-5979	May 2022	Class II
Esomeprazole Magnesium Delayed-Release 04229200916 04229201016	Failed Impurities/Degradation Specifications	If you have questions about this recall, call Mylan 1-888-406-9305	May 2022	Class II
Pantoprazole Sodium 01366809690	CGMP deviation	If you have questions about this recall, Torrent Pharma Inc, 1-888-280-2040	May 2022	Class II
Lidocaine 00591207072 00591207030	cGMP Deviations	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	May 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Vancomycin 70004092444	CGMP Deviations	If you have questions about this recall, SCA Pharmaceuticals 1-877-550-5059	May 2022	Class II
Fentanyl 70004023132	CGMP Deviations	If you have questions about this recall, SCA Pharmaceuticals 1-877-550-5059	May 2022	Class II
Norepinephrine 7000407840	CGMP Deviations	If you have questions about this recall, SCA Pharmaceuticals 1-877-550-5059	May 2022	Class II
Losartan Potassium 7051832821 7051832820	CGMP Deviations	If you have questions about this recall, RemedyRepack Inc., 1-866-845-3791	May 2022	Class II
Erythromycin Topical 06373905368	CGMP Deviations	If you have questions about this recall, call McKesson Drug Company, 1-330-487-0740.	May 2022	Class II
Betamethasone Dipropionate 06373999665	CGMP Deviations	If you have questions about this recall, call McKesson Drug	May 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

		Company, 1-330-487-0740.		
Lidocaine Prilocaine 06373905466	CGMP Deviations	If you have questions about this recall, call McKesson Drug Company, 1-330-487-0740.	May 2022	Class II
Lidocaine Hydrochloride 06373997764	CGMP Deviations	If you have questions about this recall, call McKesson Drug Company, 1-330-487-0740.	May 2022	Class II
Halobetasol Propionate 06373997764	CGMP Deviations	If you have questions about this recall, call McKesson Drug Company, 1-330-487-0740.	May 2022	Class II
Losartan Potassium & Hydrochlorothiazide 03334205007 03334205010 03334205044 03334205207 03334205210	CGMP Deviations	If you have questions about this recall, Macleods Pharma Usa Inc, 1-888-943-3210	May 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

03334205244 03334205107 03334205110 03334205144				
Losartan Potassium 03334204410 03334204444 03334204507 03334204510 03334204544 03334204607 03334204610 03334204644	CGMP Deviations	If you have questions about this recall, Macleods Pharma Usa Inc, 1-888-943-3210	May 2022	Class II
Losartan Potassium 06818037603 06818037609 06818037703 06818037709 06818037803 06818037809 06818037803 06818037809 06818021706 06818021709 06818021606 06818021609	CGMP Deviations	If you have questions about this recall, Lupin Pharmaceuticals Inc. 1-410-576-2000 ext 3	May 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Zonisamide 06846212801 06846212901 06846213001 06846213005	cGMP deviations	If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115	May 2022	Class II
Lansoprazole 04359856178	Failed Dissolution Specifications	If you have questions about this recall, Dr. Reddy's Laboratories, Inc., 1-888-375-3784	May 2022	Class II
Hydroxocobalamin 73198008030	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
Ultratest, Testosterone Cypionate 73198005810	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
Testosterone Cypionate 73198005405 73198005510 73198005505	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Formula F2, Papaverine 73198000210	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
T-50, Papaverine 73198002210	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
PGE-3, Alprostadil 73198003010	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
PGE-2, Alprostadil 73198002910	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
PGE-1, Alprostadil 73198-0028-10.	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

FA, Papaverine 73198000610	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
BIMIX-3, Papaverine 73198002710	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
RE-2, Papaverine 73198001610	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
RE-1, Papaverine 73198001510 73198001503	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

QM-4, Papaverine 73198002010	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
QM-3, Papaverine 73198001910	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
ST-2, Papaverine 73198001210	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
ST-1, Papaverine 73198001110	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
SB-6, Papaverine 73198002510	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

SB-5, Papaverine 73198002410	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
SB-4, Papaverine 73198002310	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
T-101, Papaverine 73198001410 73198001405	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
T-106, Papaverine 73198001310 73198001305	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

NB-243, Papaverine 731980009-0	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
Formula F9, Papaverine 73198000410	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
T-105, Papaverine 73198000510 73198000505 73198000503	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
QM-2 Papaverine 073198001810	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
Betamethasone Dipropionate 06373999665	CGMP Deviations	If you have questions about this recall, call McKesson Drug Company, 1-330-487-0740.	May 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

AT-6, Papaverine 73198004010	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
AT-1, Papaverine 073198003910	CGMP Deviations	If you have questions about this recall, call Olympia Compounding Pharmacy, 1-407-673-2222.	May 2022	Class II
alprazolam XR 05976200681	Viatis Inc	If you have questions about this recall, call Viatis Inc, 1-724-514-1800	May 2022	Class II
Insulin Glargine (insulin glargine-yfgn) 04950239380	Labeling: Missing label on the vial	If you have questions about this recall, call Mylan 1-888-406-9305	April 2022	Class II
Travoprost 00378965132	Subpotent Drug and Failed Impurities/Degradation Specifications	If you have questions about this recall, call Mylan 1-888-406-9305	April 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

TETRACAINE 1% Tetracaine HCl 04249443710	Lack of Assurance of Sterility	If you have questions about this recall, Vitae Enim Vitae Scientific, Inc, Inc, 1-303-236-3010	April 2022	Class II
Sucralfate Oral 06933914817 06933914819	Labeling: Label Mix-Up	If you have questions about this recall, Vitae Enim Vitae Scientific, Inc, Inc, 1-303-236-3010	April 2022	Class II
Rifampin 06745744560		If you have questions about this recall, call Mylan 1-888-406-9305	April 2022	Class II
quinapril HCl/hydrochlorothiazide 05976252259 05976202201 05976202231	CGMP Deviations	If you have questions about this recall, Pfizer Inc., 1-800-438-1985	April 2022	Class II
PHENOBARBITAL Sodium 04249441525 04249441503 04249441625 04249441603	Lack of Assurance of Sterility	If you have questions about this recall, Vitae Enim Vitae Scientific, Inc, Inc, 1-303-236-3010	April 2022	Class II
PAPAVERINE HYDROCHLORIDE 072516-02425 007251602410	Lack of Assurance of Sterility	If you have questions about this recall, Vitae Enim Vitae Scientific, Inc, Inc, 1-303-236-3010	April 2022	Class II
Orphenadrine Citrate Extended-Release 00185002201	CGMP Deviations	If you have questions about this recall, Sandoz, Inc, 1-609-627-8500	April 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Lansoprazole Delayed-Release 068788639009	Out of specification	If you have questions about this recall, Preferred Pharmaceuticals, Inc., 1-714-777-3729	April 2022	Class II
Janumet 00006057502 00006057503	Presence of Foreign Substance	If you have questions about this recall, MERCK SHARP & DOHME CORP, 1-(908) 423-1000	April 2022	Class II
Glycopyrrolate 01310701401	Presence of Foreign Substance	If you have questions about this recall, Aurolife Pharma, LLC, 1-732-839-9400 option 2	April 2022	Class II
Econazole Nitrate 068788740603	CGMP Deviations	If you have questions about this recall, Preferred Pharmaceuticals, Inc., 1-714-777-3729	April 2022	Class II
Diclofenac Sodium 68788791801	CGMP Deviations	If you have questions about this recall, Preferred Pharmaceuticals, Inc., 1-714-777-3729	April 2022	Class II
Clobetasol Propionate 068788776801	CGMP Deviations	If you have questions about this recall, Preferred Pharmaceuticals, Inc., 1-714-777-3729	April 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Accuretic (quinapril HCl/hydrochlorothiazide) 00071311223 00071022223 00071022023 00071022323	CGMP Deviations	If you have questions about this recall, Pfizer Inc., 1-800-438-1985	April 2022	Class II
Accuretic 00071521223	CGMP Deviations	If you have questions about this recall, Pfizer Inc., 1-800-438-1985	April 2022	Class II
IDArubicin Hydrochloride 00703415411	Presence of Particulate Matte	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	April 2022	Class I
SYMJEPI 07867013002 07867013102	Defective Delivery Syst	If you have questions about this recall, Adamis Pharmaceuticals Corporation, 1-858-997-2400	April 2022	Class I
Triamcinolone Acetonide 05256505630 05256505680 05256505626 05256504815 05256501059 05256501415	cGMP deviations	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

05256501480 05256501426				
Nystatin and Triamcinolone Acetonide 052565004215 005256504230 05256504260	cGMP deviations	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II
Lidocaine 05256512215 05256512230 05256512207 05256500814 00536128128 05038334135	cGMP deviations	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II
Hydrocortisone Butyrate 05256508702 05256508704 05186215904 05256508702 05256508704	cGMP deviations	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II
Halobetasol Propionate 07051203350 05256507315 05256507351 06373999867	cGMP deviations	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Gentamicin Sulfate 05256508515 05256508530 05256509015 05256509030 07051203630	cGMP deviations	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II
Fluocinonide 05256505415 05256505460 05256505430 05256507911 05256502520 05256502559	cGMP deviations	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II
Econazole Nitrate 05256502215 05256502285 05256502230	cGMP deviations	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II
Diflorasone Diacetate 07051203160 05256506360	cGMP deviations	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II
Diclofenac Sodium 06191967505	Defective Container	If you have questions about this recall, Direct Rx, 1-678-619-5510	March 2022	Class II
Diclofenac Sodium 05256500205 07051202505	cGMP deviations	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Desoximetasone 05256504599 05256504560 07051203710 05256503099 05256503060 05256503015	cGMP deviations	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II
Desonide 05256503815 05256503860	cGMP deviations	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II
Clobetasol Propionate 05256509415 05256509430 05256509445 05256505502 05256505504 05256503915 05256503930 05256503945 05256503960 05256508215 05256508230 05256508260 07051202860	cGMP deviation	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II
Betamethasone Dipropionate 05256502329 05256502359	cGMP deviation	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	March 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Alprazolam 06042950418	CGMP Deviation	If you have questions about this recall, Golden State Medical Supply Inc., 1-805-477-9866 Ext 4.	March 2022	Class II
hydrALAZINE 00904644061	Failed Impurities/Degradation Specifications	If you have questions about this recall, The Harvard Drug Group, 1-800-875-0123	March 2022	Class II
Alprazolam C-IV 06191983660 72189005860	CGMP Deviations	If you have questions about this recall, Direct Rx, 1-678-619-5510	March 2022	Class II
PALIPERIDONE EXTENDED-RELEASE 00904693761	Failed Dissolution Specifications	If you have questions about this recall, The Harvard Drug Group, 1-800-875-0123.	March 2022	Class II
Oxycodone Hydrochloride 00360687406 06068-40677 06068740640	Impurity failure at 0-time of the repackaged lot	If you have questions about this recall, American Health Packaging, 1-614-492-8177.	March 2022	Class II
Methylphenidate Hydrochloride 6498022101	Failed Tablet Specifications	If you have questions about this recall, RISING PHARMACEUTICALS, 1-866-562-4597	March 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Diazepam 00527176836	Failed Impurities/Degradation Specifications	If you have questions about this recall, Lannett Company, Inc. 1-215-333-9000	February 2022	Class II
Morphine Sulfate 73177010504	Labeling; label mix-up	If you have questions about this recall, STAQ Pharma, Inc., 1-833-397-0106.	February 2022	Class I
Hydromorphone HCL 73177010405	Labeling; label mix-up	If you have questions about this recall, STAQ Pharma, Inc., 1-833-397-0106.	February 2022	Class I
Doxylamine Succinate and Pyridoxine Hydrochloride 0591-2132-01	Failed Dissolution Specifications	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	February 2022	Class II
Tretinoin 00555080802	Failed Dissolution Specifications	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	February 2022	Class II
Proctofoam 00037682210	cGMP deficiencies	If you have questions about this recall, call Mylan 1-888-406-9305	February 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Metformin Hydrochloride 07257803601	CGMP Deviations	If you have questions about this recall, VIONA PHARMACEUTICALS INC, 1-888-304-5022	February 2022	Class II
Proctofoam HC 00037682210	cGMP deficiencies	If you have questions about this recall, call Mylan 1-888-406-9305	February 2022	Class II
Diazepam 00527176836	Failed Impurities/Degradation Specifications	If you have questions about this recall, Lannett Company, Inc. 1-215-333-9000.	February 2022	Class II
Trypan Blue 05446134801	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Vancomycin HCl 05446134801	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Tetracaine HCl 05446119503	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see *FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls* for additional information.

Promethazine HCl Topical 05446134101	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Vitamin K Oral 05446113203	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Lidocaine HCl/Phenylephrine HCl 05446104503	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Phenol, Topical 05446121103	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Profound-PE Dental Gel 05446040810	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Profound-PE Dental Gel11 05446101810	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Profound Dental Gel 05446079010 05446040710	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Lidocaine HCl / Oxymetazoline HCl 05446125601	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
LET Topical Gel 05446060701	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
LT Topical 05446164701	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Dibutyl Squaric Acid 05446104703 05446115603	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Dexamethasone sodium phosphate 05446062201	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Cantharidin PLUS 05446097003	CGMP Deviations	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Cantharidin Gel 05446057203	CGMP Deviations	If you have questions about this recall,	February 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

		Edge Pharma, LLC. 1-802-992-1178.		
Vancomycin HCl 05446073601	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Betadine (povidone-iodine) 05446168001	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Phenylephrine HCl/Tropicamide/Cyclopentolate HCl/ Ketorolac 05446085903	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Phenylephrine HCl/Tropicamide/Cyclopentolate HCl/Ketorolac 05446099301	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Phenylephrine HCl/Tropicamide 05446081501	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Phenylephrine HCl/Lidocaine 05446111801	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Phenylephrine HCl 05446127001 05446154505 05446154410 05446165201	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
MVASI 05446166113	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Neostigmine 05446154905	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Moxifloxacin 05446105001	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Mitomycin-C 05446141601 05446100901 05446101101	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Methotrexate 05446150505	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Methacholine 05446160005 05446124601 05446124701 05446124901	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Lidocaine HCL / Bupivacaine HCL 05446154818	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Gemcitabine 05446156650	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Epinephrine/Lidocaine 05446086301	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Edetate Disodium (EDTA) 00544614271 05446142810	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Dexamethasone sodium phosphate 05446084801	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Cefuroxime 05446100301	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Ceftazidime 05446073301	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Lidocaine HCl/Epinephrine 05446126801	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Lidocaine HCl 05446085010	Lack of Assurance of Sterility	If you have questions about this recall, Edge Pharma, LLC. 1-802-992-1178.	February 2022	Class II
Polymyxin B 05515023410	Presence of Particulate Matter	If you have questions about this recall, AuroMedics Pharma LLC, 1-732-839-9400 ext 2.	February 2022	Class II
Moxifloxacin 06586284003	Failed impurities/degradation specifications	If you have questions about this recall, Aurobindo Pharma, 1-866-850-2876	February 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Pioglitazone 05723722105	Superpotent and Failed Tablet/Capsule Specifications	If you have questions about this recall, Aurobindo Pharma, 1-866- 850-2876	February 2022	Class II
Metoprolol Succinate Extended- Release 06787759001	Failed Dissolution Specifications	If you have questions about this recall, Ascend Laboratories LLC, 1-201- 476-1977.	February 2022	Class II
Pyrazinamide 06725366010	cGMP Deviations	If you have questions about this recall, ANI Pharmaceuticals, Inc., 1- 800-308-6755.	February 2022	Class II
Alprazolam 06725390110 06725390150 06725390111 06725390210 06725390250 06725390211 06725390310 06725390350	cGMP Deviations	If you have questions about this recall, ANI Pharmaceuticals, Inc., 1- 800-308-6755.	February 2022	Class II
Metoprolol Succinate Extended- Release 06800150103	Failed Dissolution Specifications	If you have questions about this recall, American Health Packaging, 1-614- 492-8177.	February 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Lexette (halobetasol propionate) 05186260450	CGMP Deviation	If you have questions about this recall, Mayne Pharma Inc, 844 825 8500	January 2022	Class II
Clobazam 07778905804	Failed Stability Specifications	If you have questions about this recall, Ingenus Pharmaceuticals LLC, 877-530-1633.	January 2022	Class II
Clobetasol Propionate 05074230450 05074230401	CGMP Deviations	If you have questions about this recall, Ingenus Pharmaceuticals LLC, 1-407-354-5740	January 2022	Class II
Brinzolamide 00591212779 00591212712	Defective Container	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	January 2022	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

8.4% Sodium Bicarbonate 05175450011	Lack of Assurance of Sterility	If you have questions about this recall, Exela Pharma Sciences LLC, 1-888-451-4321	January 2022	Class II
Metformin 70518292000	CGMP Deviations	If you have questions about this recall, RemedyRepack Inc., 1-866-845-3791	January 2022	Class II
Norepinephrine Bitartrate 00703115303	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	January 2022	Class II
MethylPREDNISolone Acetate 01671409001	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	January 2022	Class II
Carbamazepine 01366826801	Failed Dissolution Specifications	If you have questions about this recall, Torrent Pharma Inc, 1-888-280-2040.	January 2022	Class II
Clobetasol Propionate 05167212593	Microbial Contamination of Non-Sterile Products	If you have questions about this recall, Taro Pharmaceuticals U.S.A., Inc., 1-800-544-1449 ext 6066.	January 2022	Class I

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Pregabalin 04733568788	Failed Tablet/Capsule Specifications	If you have questions about this recall, SUN PHARMACEUTICAL INDUSTRIES INC, 1-800-818-4555	January 2022	Class II
Lidocaine Hydrochloride 06373999764	Superpotent Drug	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	January 2022	Class II
Lidocaine Hydrochloride 06373999764	Superpotent Drug	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	January 2022	Class I
Betamethasone Dipropionate 05256502329	Failed Stability Specifications	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	January 2022	Class II
Penicillin V Potassium 00093412573 00093412574	Subpotent	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	December 2021	Class II
Fexofenadine Hydrochloride 00904697940	Failed Impurities/Degradation Specifications	If you have questions about this recall, The Harvard Drug Group, 1-800-875-0123.	December 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Clindamycin and Benzoyl Peroxide Gel 00781726368	Superpotent Drug	If you have questions about this recall, TOLMAR, Inc., 1-970-212-4500	December 2021	Class II
Lidocaine Hydrochloride 05256500950	Superpotent Drug	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	December 2021	Class I
Lidocaine Hydrochloride 06373999764 05256500950	CGMP Deviations	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	December 2021	Class II
Cefixime 70518274902	Failed Impurities/Degradation Specifications	If you have questions about this recall, RemedyRepack Inc., 1-866-845-3791	December 2021	Class II
Diclofenac Sodium 68788770701	Defective container	If you have questions about this recall, Preferred Pharmaceuticals, Inc., 1-714-777-3729	December 2021	Class II
Hydrocodone Bitartrate and Acetaminophen 03172299701	Product Mix-up	If you have questions about this recall, Ascent Pharmaceuticals, Inc., 1-631-851-0550	December 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Cefixime 06787758450	Failed impurities/degradation specifications	If you have questions about this recall, Ascend Laboratories, LLC, 1-201-476-1977	December 2021	Class II
Ascorbic Acid	Lack of Assurance of Sterility	If you have questions about this recall, ASP Cares, 1-210-615-7400	December 2021	Class II
Diclofenac Sodium 05266500205 05266502505	Defective Container	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	December 2021	Class II
Floclonolone Acetonide 06516270486 06516270386	Subpotent Drug	If you have questions about this recall, Amneal Pharmaceuticals., 1-833-582-0812.	December 2021	Class II
Cefixime 06787758450	Failed Impurities/Degradation Specifications	If you have questions about this recall, Ascend Laboratories LLC, 1-201-476-1977	November 2021	Class II
Tadalafil 01671407501 01671407701	Incorrect Product Formulation	If you have questions about this recall, SUN PHARMACEUTICAL INDUSTRIES INC, 1-800-818-4555	November 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Ezetimibe and Simvastatin 05140719305	Failed Excipient Specification	If you have questions about this recall, Dr. Reddy's Laboratories, Inc., 1-888-375-3784	November 2021	Class II
Ezetimibe and Simvastatin 05140719290 05140719205	cGMP Deviations	If you have questions about this recall, Dr. Reddy's Laboratories, Inc., 1-888-375-3784	November 2021	Class II
Methocarbamol 07133517952 07133517954 07133517957	Labeling: Label Error on Declared Strength	If you have questions about this recall, 1-877-885-0882.	November 2021	Class I
Irbesartan and Hydrochlorothiazide 06818041306 06818041309 06818041406 06818041409	CGMP Deviations	If you have questions about this recall, Lupin Pharmaceuticals Inc. 1-410-576-2000 ext 3	November 2021	Class II
Irbesartan 06818041006 06818041009 06818041106 06818041109 06818041206 06818041209	CGMP Deviations	If you have questions about this recall, Lupin Pharmaceuticals Inc. 1-410-576-2000 ext 3	November 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Ezetimibe and Simvastatin 04359874290 04359874210 04359874430 04359874490 04359874530 04359874590 04359874505 04359874330 04359874390 04359874305	Failed Excipient Specifications	If you have questions about this recall, Dr. Reddy's Laboratories, Inc., 1-888-375-3784	November 2021	Class II
Rocuronium Bromide 66794022841	Labeling: Label Lacks Warning or Rx Legend	If you have questions about this recall, Piramal Critical Care, Inc. 1-800-414-1901	October 2021	Class II
Imipramine Pamoate	Out of specification result observed in a dissolution test	If you have questions about this recall, Lupin Pharmaceuticals Inc. 1-410-576-2000 ext 3	October 2021	Class II
Omeprazole Delayed-Release 07000002321	CGMP Deviations	If you have questions about this recall, Dr. Reddy's Laboratories, Inc., 1-888-375-3784	October 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

AirDuo Digihaler 232/14 (fluticasone propionate 113 mcg and salmeterol 14 mcg) 05931013606	Subpotent drug	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	October 2021	Class II
AirDuo Digihaler 113/14 (fluticasone propionate 113 mcg and salmeterol 14 mcg) 05931012906	Subpotent drug	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	October 2021	Class II
AirDuo Digihaler 55/14 (fluticasone propionate 55 mcg and salmeterol 14 mcg) 05931011106	Subpotent drug	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	October 2021	Class II
AirDuo Digihaler 232/14 (fluticasone propionate 232 mcg and salmeterol 14 mcg) 05931053008	Subpotent drug	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	October 2021	Class II
AirDuo Digihaler 113/14 (fluticasone propionate 113 mcg and salmeterol 14 mcg) 05931052008	Subpotent drug	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	October 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

AirDuo Digihaler 113/14 (fluticasone propionate 113 mcg and salmeterol 14 mcg) 05931052008	Subpotent drug	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	October 2021	Class II
GlipiZIDE Extended-Release 06808429521 06808429511	Failed Dissolution Specifications: results were above specification.	If you have questions about this recall, American Health Packaging, 1-614-492-8177	October 2021	Class II
Glucagon Emergency Kit 00002803101	Subpotent Drug	If you have questions about this recall, Eli Lilly & Company, 1-800-545-5979	October 2021	Class I
Betaxolol Ophthalmic 01747870510 01747870511	Microbial Contamination of Sterile Products	If you have questions about this recall, Akorn, Inc., 1-855-526- 4827	October 2021	Class II
Artesunate 07360700101 07360700111 07360700101	Lack of Assurance of Sterility	If you have questions about this recall, AMIVAS (US), LLC, 1-855-526-4827	October 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

07360700102 07360700110				
Meclizine HCl Tablets, 25 mg 01657175201	Labeling: Incorrect Instructions	If you have questions about this recall, RISING PHARMACEUTICALS, 1-866-562-4597	October 2021	Class II
Firvanq (vancomycin hydrochloride for oral solution) 06562820605	Product Mix-up	If you have questions about this recall, Azurity Pharmaceuticals, Inc., 1-800-461-7449	October 2021	Class I
Morphine Sulfate 06332345201	Defective container	If you have questions about this recall, Fresenius Kabi USA, 1-800-551-7176.	October 2021	Class II
Valproic Acid 06068726256 06068726242	CGMP Deviations	If you have questions about this recall, American Health Packaging, 1-800-707-4621.	October 2021	Class II
Lyrica CR (pregabalin) 00071102901	Failed Dissolution Specifications	If you have questions about this recall, Pfizer Inc., 1-800-438-1985.	October 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Zonisamide 61919077530	CGMP deviations	If you have questions about this recall, Direct Rx, 1-678-619-5510	September 2021	Class II
Metoprolol Tartrate 06586206499	Presence of Foreign Substance	If you have questions about this recall, Aurobindo Pharma USA Inc., 1-866-850-2876	September 2021	Class II
Entacapone 00904682204	Failed Dissolution Specifications	If you have questions about this recall, The Harvard Drug Group, 1-800-875-0123.	September 2021	Class II
Valproic Acid 06043262116	CGMP Deviations	If you have questions about this recall, Morton Grove Pharmaceuticals, Inc., 1-888-721-7115	September 2021	Class II
Promethazine With Codeine 06043260616	CGMP Deviations	If you have questions about this recall, Morton Grove Pharmaceuticals, Inc., 1-888-721-7115	September 2021	Class II
Promethazine Syrup 06043260816	CGMP Deviations	If you have questions about this recall, Morton Grove Pharmaceuticals, Inc., 1-888-721-7115	September 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Arformoterol Tartrate 06846283365 06846283335	Lack of Assurance of Sterility	If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115	September 2021	Class II
Zonisamide 06846212901 06846213001 06846213005	CGMP Deviations	If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115	September 2021	Class II
Chlorzoxazone 06846272401 06846272501	CGMP Deviations	If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115	September 2021	Class II
Naproxen Sodium 06846217801 06846217901 06846217905	CGMP Deviations	If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115	September 2021	Class II
Fulvestrant 06846231732	Lack of Assurance of Sterility	If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115	September 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Carbamazepine 00904617261	Failed Dissolution Specifications	If you have questions about this recall, The Harvard Drug Group, 1-800-875-0123.	September 2021	Class II
Betadine Solution Swabstick 06761815301 06761815303	Subpotent Drug	If you have questions about this recall, AVRIO HEALTH L.P, 1-888-726-7535	September 2021	Class II
Chantix (varenicline) 00069047103 00069046856 00069046956 00069046903	CGMP Deviations	If you have questions about this recall, Pfizer Inc., 1-800-438-1985.	September 2021	Class II
Oxycodone Hydrochloride 04285800201	Presence of Foreign Tablets/Capsules; A single foreign tablet Hydrochlorothiazide/Lisinopril 25/20 was found in one bottle	If you have questions about this recall, Akorn, Inc, Rhodes Pharmaceuticals, L.P., 1-401-262-9400, Prompt 2	September 2021	Class II
Betamethasone Dipropionate 06174848030	Failed impurities/degradation specification: Out of Specification for an unknown impurity observed in topical product.	If you have questions about this recall, Akorn, Inc, 1-800-932-5676, Prompt 2	September 2021	Class II
Naproxen Sodium 06220776236	CGMP Deviations	If you have questions about this recall, Granules USA, Inc., 1877-770-3183	September 2021	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Clopidogrel 03334206015	Presence of foreign matter	If you have questions about this recall, Macleods Pharma Usa Inc, 1-888-943-3210	September 2021	II
Cyclobenzaprine Hydrochloride 00591333001 07019901401 05723726601	CGMP Deviations	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	September 2021	II
FLUDARABINE PHOSPHATE FOR INJECTION 02420123701	Lack of Assurance of Sterility: the manufacturing firm had microbial recoveries during environmental monitoring in aseptic areas of manufacturing.	If you have questions about this recall, Custopharm, Inc., 1760-683-0901	September 2021	II
Erythromycin Topical Solution 05256502759	Defective container: possibility for lack of seal integrity.	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	September 2021	II
Atovaquone Oral Suspension 01070222321	Temperature abuse: the firm received customer complaints of unusual grittiness in the product.	If you have questions about this recall, KVK-Tech, Inc., 1-215-579-1842	September 2020	II
Carvedilol 06838209505 70518182601	Presence of Foreign Tablets/Capsules; report of two Paroxetine tablets were found in the bottle	If you have questions about this recall, RemedyRepack Inc., 1-866-845-3791	September 2021	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/Drugs/DrugSafetyandAvailability/DrugRecalls) for additional information.

Adenosine 01671418001	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
Leucovorin Calcium 00703514001 00703514591	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
Octreotide Acetate 00703331101 00703330101	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
Methylprednisolone Acetate 00703005101 00703006301	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
Alprostadil 00703150101	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
Metoclopramide 00703450201	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
Adenosine Injection 00703877601	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Norepinephrine Bitartrate 00703115301	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
Epoprostenol Sodium 00703199501	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
Leucovorin Calcium 00703514501	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
Octreotide Acetate 00703333301	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
Vecuronium Bromide 00703291401	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
Idarubicin Hydrochloride 00703415611	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
Amikacin Sulfate 00703904001	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Haloperidol Decanoate 00703713101 00703713301	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
Methylprednisolone Acetate 00703003101 00703004301 00703004501	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
DAUNOrubicin Hydrochloride 00703523313	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	August 2021	II
GaviLyte™ 04338606019	Failed Stability Specification; Out of specification for Osmolarity	If you have questions about this recall, Lupin Pharmaceuticals Inc. 1-410-576-2000 ext 3	August 2021	II
Combipatch 06896805148 06896805258	Failed Stability Specifications; out of specification for shear.	If you have questions about this recall, Noven Pharmaceuticals Inc, 1-305-253-5099	August 2021	II
Tizanidine HCl 05511118015	Failed Tablet/Capsule Specification: Some tablets are shaved	If you have questions about this recall, Dr. Reddy's Laboratories, Inc., 1-888-375-3784	August 2021	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Triamcinolone Acetonide 05256501480	Correct Labeled Product Mispack	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	August 2021	II
Econazole Nitrate 05256501480	Correct Labeled Product Mispack	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	August 2021	II
Ethosuximide 00121067016	Voluntary: Firm initiated	If you have questions about this recall, PAI Holdings, LLC. dba Pharmaceutical Associates Inc1-864-277-7282 ext 0	August 2021	II
Cimetidine Hydrochloride 00121064908	Voluntary: Firm initiated	If you have questions about this recall, PAI Holdings, LLC. dba Pharmaceutical Associates Inc1-864-277-7282 ext 0	August 2021	II
Nystatin Oral 00121086816 00121086802	Voluntary: Firm initiated	If you have questions about this recall, PAI Holdings, LLC. dba Pharmaceutical Associates Inc1-864-277-7282 ext 0	August 2021	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Venlafaxine 01671465701	Voluntary: Firm initiated		August 2021	II
Sulfamethoxazole and Trimethoprim 06586242005	Voluntary: Firm initiated	If you have questions about this recall, Aurobindo Pharma USA Inc., 1-866-850-2876	August 2021	II
Zyprexa Intramuscular, Olanzapine 00002759701	Voluntary: Firm initiated	If you have questions about this recall, Eli Lilly & Company. 1-317-276-2000	August 2021	II
Chantix 00069046856 00069047103 00069046956	CGMP Deviations	If you have questions about this recall, Pfizer Inc., 1-800-438-1985.	August 2021	II
Estriol 04614430001	cGMP Deviations	If you have questions about this recall, API Solutions Inc., 1-855-878-1489	August 2021	II
NIFEdipine EXTENDED-RELEASE 00904708061 00904708006	Failed Dissolution Specification	If you have questions about this recall, Ingenus Pharmaceutical. 1-877-748-1970	July 2021	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Solifenacin Succinate 06909726102	CGMP Deviations	If you have questions about this recall, CIPLA, 1-866-604-3268.	July 2021	II
Xylocaine-MPF with Epinephrine 06332348737 06332348707	Low out of specification results for epinephrine assay	If you have questions about this recall, Genentech Inc, 1-888-835-2555.	July 2021	II
Xolair (omalizumab) 05024221501	Failed Stability Specifications	If you have questions about this recall, Fresenius Kabi USA, 1-800-551-7176.	July 2021	II
Buprenorphine and Naloxone Sublingual Film 04778135503	Subpotent drug	If you have questions about this recall, Alvogen, Inc. 1-866-770-3024	July 2021	II
Topotecan Injection 00703471471	Presence of Particulate Matter	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	July 2021	I
Metformin Hydrochloride Extended-Release 07257803601	CGMP Deviations	If you have questions about this recall, VIONA PHARMACEUTICALS INC, 1-888-304-5022	June 2021	Class II
DermOtic Oil (fluocinolone acetonide oil) 68791-103-20	Presence of Foreign Substance	If you have questions about this recall, Hill Dermaceuticals, Inc., 1-407-323-1998	June 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Diflorasone Diacetate Ointment 05256506315 05256506330	Presence of Foreign Substance	If you have questions about this recall, Teligent Pharma, 1-856-697-1441	June 2021	Class II
BusPIRone Hydrochloride 06958409310	Presence of Foreign Tablets/Capsules	If you have questions about this recall, Oxford Pharmaceuticals, LLC, 1-205-434-9649	June 2021	Class II
Micardis 0597004137	Subpotent Drug	If you have questions about this recall, Boehringer Ingelheim Pharmaceuticals, Inc., 1-908-927-1400	June 2021	Class II
ChloroPrep With Tint 2% w/v chlorhexidine gluconate 005436540011	Microbial Contamination of Non-Sterile Product	If you have questions about this recall, CareFusion 213, LLC, 1-201-847-6800.	May 2021	Class I
ChloroPrep One-Step 2% w/v chlorhexidine gluconate 005436540001	Microbial Contamination of Non-Sterile Product	If you have questions about this recall, CareFusion 213, LLC, 1-201-847-6800.	May 2021	Class I
BD ChloroPrep Hi-Lite Orange 2 05436540033	Non-sterility	If you have questions about this recall, CareFusion 213, LLC, 1-201-847-6800.	May 2021	Class I

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

BD Chloraprep Clear 05436540032	Non-sterility	If you have questions about this recall, CareFusion 213, LLC, 1-201-847-6800.	May 2021	Class I
Estradiol Transdermal System 06896834378	Defective Delivery System	If you have questions about this recall, Noven Pharmaceuticals Inc, 1-305-253-5099	April 2021	Class II
Minivelle 06896866758	Defective Delivery System	If you have questions about this recall, Noven Pharmaceuticals Inc, 1-305-253-5099	April 2021	Class II
Itraconazole 05974628230	Failed Dissolution Specifications	If you have questions about this recall, Jubilant Cadista Pharmaceuticals, Inc., 1-410- 860-2836	April 2021	Class II
Riomet (metformin hydrochloride oral solution) 01063120602	Microbial Contamination of Non-Sterile Product	If you have questions about this recall, SUN PHARMACEUTICAL INDUSTRIES INC, 1-800-818-4555	April 2021	Class II
Cefprozil 06818040201 06818040202 06818040203	Superpotent Drug	If you have questions about this recall, Lupin Pharmaceuticals Inc., 1-866-587-4617	April 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/Drugs/DrugSafetyandAvailability/DrugRecalls) for additional information.

Guanfacine Extended-Release 06050539281 36050539281	Cross Contamination	If you have questions about this recall, Apotex Corp., 1 866.390-4411.	April 2021	Class II
Mometasone Furoate 00713070185 00713070153	CGMP Deviatons	If you have questions about this recall, Cosette Pharmaceuticals, Inc., 1 866.390-4411.	April 2021	Class II
Neomycin Sulfate 03982203105	Failed Stability	If you have questions about this recall, X-Gen Pharmaceuticals Inc., 1-866-390-4411.	April 2021	Class II
Telmisartan 06233208730	Labeling: Label-mixup	If you have questions about this recall, Alembic Pharmaceuticals Limited, 1-908-393-9604	April 2021	Class I
Progesterone Capsules 04359835001	Failed Dissolution Specifications	If you have questions about this recall, Dr. Reddy's Laboratories, Inc., 1-888-375-3784	March 2021	Class II
Metoclopramide Injection 0703450204	Chemical contamination	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	March 2021	Class II
Famotidine Tablets 06586286099	Presence of foreign tablets/capsules	If you have questions about this recall, Aurobindo Pharma USA Inc., 1-866-850-2876	March 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/Drugs/DrugSafetyandAvailability/DrugRecalls) for additional information.

Phenylephrine HCl 02502131501	Lack of Assurance of Sterility	If you have questions about this recall, Sagent Pharmaceuticals Inc, 1-866-625-1618	March 2021	Class II
Gabapentin 05038331107	Failed Impurities/Degradation Specifications	If you have questions about this recall, Akorn, Inc, 1-800-932-5676, Prompt 2	March 2021	Class II
Nortriptyline HCL 06191985330	Failed Impurities/Degradation Specifications	If you have questions about this recall, Direct Rx, 1-678-619-5510	March 2021	Class II
Spironolactone Tablets 63629106701	Labeling: Label Mix-Up	If you have questions about this recall, BRP Pharmaceuticals, 1-877-885-0882	March 2021	Class II
Daytrana (methylphenidate transdermal system) 06896855523 06896855533 06896855543 06896855553	Defective Delivery System: Out of specification for mechanical peel	If you have questions about this recall, Noven Pharmaceuticals Inc, 1-305-253-5099	March 2021	Class II
Omeprazole Delayed Release Capsules 05199164310	Failed Impurities/Degradation Specifications	If you have questions about this recall, Breckenridge Pharmaceutical, Inc, 1-860-828-8140	March 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Omeprazole Delayed Release Capsules 05140712910	Failed Impurities/Degradation Specifications	If you have questions about this recall, Golden State Medical Supply Inc., 1-805-477-9866	March 2021	Class II
Fludeoxyglucose F 18 07631833450	Lack of Assurance of Sterility	If you have questions about this recall, Massachusetts General Hospital PET Center, 1-617-726-2000	March 2021	Class II
Irinotecan Hydrochloride 05992371402	CGMP Deviations	If you have questions about this recall, Areva Pharmaceuticals Inc, 1-855-853-4760	March 2021	Class II
Imatinib Mesylate 04359834490 00435984431	Failed Dissolution Specifications	If you have questions about this recall, Dr. Reddy's Laboratories, Inc., 1-888-375-3784	March 2021	Class II
Epoprostenol Sodium 00703199501	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	March 2021	Class II
Vecuronium Bromide 00703291401 00703291403	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	March 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Toposar 00703565701	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800	March 2021	Class II
Metoclopramide Injection 00703450201 00703450204	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800.	March 2021	Class II
Leucovorin Calcium 00703514001 00703514501	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800.	March 2021	Class II
MethyIPREDNISolone Acetate 00703003101 00703005101 00703005104 00703004501 00703004301 00703006301	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800.	March 2021	Class II
Epoprostenol Sodium 00703198501	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800.	March 2021	Class II
Sterile Diluent for Epoprostenol Sodium 0703925801	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800.	March 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Desmopressin Acetate 00703505101 00703505103	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800.	March 2021	Class II
Dacarbazine 00703507501 00703507503	Lack of Assurance of Sterility	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800.	March 2021	Class II
Cisatracurium Besylate 07128871206	Labeling: Label mix-up	If you have questions about this recall, Meitheal Pharmaceuticals Inc, 1-224-443-4617.	March 2021	Class I
BD Chloraprep Hi-Lite Orange 2% w/v chlorhexidine gluconate (CHG) and 70% v/v isopropyl alcohol (IPA), Sterile Solution, 0.01 fl. oz 05436540033	Non-sterility	If you have questions about this recall, CareFusion 213, LLC, 1-201-847-6800.	March 2021	Class I
Chloraprep One-Step 2% w/v chlorhexidine gluconate (CHG) and 70% v/v isopropyl alcohol (IPA) Non-Sterile Solution -Clear, 0.10 fl. Oz 05436540001	Non-sterility	If you have questions about this recall, CareFusion 213, LLC, 1-201-847-6800.	March 2021	Class I
Chloraprep With Tint 2% w/v chlorhexidine gluconate (CHG) and 70% v/v isopropyl alcohol (IPA) Non-Sterile Solution - Hi-Lite Orange, 0.10 fl. Oz	Microbial Contamination			Class I

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

05436540011				
BD Chloraprep Clear, 2% w/v chlorhexidine gluconate (CHG) and 70% v/v isopropyl alcohol (IPA) Sterile Solution, 0.10 fl 05436540032	Non-sterility	If you have questions about this recall, CareFusion 213, LLC, 1-201-847-6800.	March 2021	Class I
Meclizine HCl Tablets 05253612901 05253613301	Failed Dissolution Specifications	If you have questions about this recall, Wilshire Pharmaceuticals, Inc, Ltd, 1-877-495-6856.	February 2021	Class II
Cephalexin for Oral Suspension 06787754568	CGMP Deviations	If you have questions about this recall, Alkem Laboratories, Ltd, 1-636-343-5664.	February 2021	Class II
Metformin Hydrochloride Extended-Release Tablets 02903305601	CGMP Deviations	If you have questions about this recall, Nostrum Laboratories Inc, 1-816-841-4636.	February 2021	Class II
Nortriptyline HCl Capsules 05167240011	CGMP Deviations	If you have questions about this recall, Taro Pharmaceuticals U.S.A., Inc., 1-800-544-1449 ext 6066.	February 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Acetaminophen Injection 05515030701	Discoloration and failed pH specifications	If you have questions about this recall, AuroMedics Pharma LLC, 1-732-839-9400 ext 2.	February 2021	Class II
Ketorolac Tromethamine Injection 06332316201	Presence of Particulate Matter	If you have questions about this recall, Fresenius Kabi USA, 1-800-551-7176.	February 2021	Class I
Nitrofurantoin Capsules 06808444601 06808444611	Failed Dissolution Specifications	If you have questions about this recall, American Health Packaging, 1-800-707-4621.	January 2021	Class II
Chlorhexidine Gluconate 06809402861	cGMP Deviations	If you have questions about this recall, Precision Dose Inc., 1-800-397-9228.	January 2021	Class II
Metformin Hydrochloride 02903305601	cGMP Deviations	If you have questions about this recall, Nostrum Laboratories Inc, 1-816-308-4941.	January 2021	Class II
Paroex (Chlorhexidine Gluconate) 05237602102 05237602104	cGMP Deviations	If you have questions about this recall, Sunstar Americas, Inc., 1-800-528-8527	January 2021	Class II
Levetiracetam 05038324116	Defective container	If you have questions about this recall, Akorn, Inc, 1-800-932-5676, Prompt 2	January 2021	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Cephalexin 06787754488 06787754468	Failed Impurity/Degradation Specifications	If you have questions about this recall, Ascend Laboratories LLC, 1-877-272-7901.	January 2021	Class II
Esomeprazole Magnesium 06909752734 06909752834 06909752934	Cross- contamination with other products	If you have questions about this recall, CIPLA, 1-866-604-3268.	January 2021	Class II
Cephalexin 06787754568 06909752734	Failed Impurity/Degradation Specifications	If you have questions about this recall, Ascend Laboratories LLC, 1-877-272-7901.	January 2021	Class II
Auryxia (ferric citrate) 05992263101	CGMP Deviations	If you have questions about this recall, Akebia Therapeutics dba Keryx Biopharmaceuticals, Inc, 1-617-871-2098.	December 2020	Class II
Hydroxyzine Hydrochloride 06043215004 06043215016	Failed Impurities/Degradation Specification	If you have questions about this recall, Morton Grove Pharmaceuticals, Inc., 1-847-967-5600, Prompt 4.	December 2020	Class II
Vumerity (diroximel fumarate) 06440602001	Failed dissolution specifications	If you have questions about this recall, Biogen MA Inc., 1-844-477-4672.	December 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Lansoprazole Delayed-Release 06838277177	Failed Dissolution Specifications	If you have questions about this recall, Zydus Pharmaceuticals (USA) Inc, 1-877-993-8779, Prompt 2.	December 2020	Class II
Lansoprazole Delayed-Release 06838277277	Failed Dissolution Specifications	If you have questions about this recall, Zydus Pharmaceuticals (USA) Inc, 1-877-993-8779, Prompt 2.	December 2020	Class II
Sildenafil 04229174801	Product mix-up	If you have questions about this recall, AVKARE Inc., 1-855-361-3993.	December 2020	Class II
TraZODONE Hydrochloride 04229183410	Product mix-up	If you have questions about this recall, Shilpa Medicare Limited, 1-732-637-1971.	December 2020	Class II
Imatinib Mesylate 07248520330	GMP Deviations	If you have questions about this recall, Shilpa Medicare Limited, 1-732-637-1971.	December 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Imatinib Mesylate 07248520290d	GMP Deviations	If you have questions about this recall, Shilpa Medicare Limited, 1-732-637-1971.	December 2020	Class II
Anagrelide 01366846201	Failed Dissolution Specifications	If you have questions about this recall, Torrent Pharma Inc, 1-888-280-2040.	December 2020	Class I
(Lidocaine HCL 2%) Topical Anesthetic Hydrogel 06697710703	Microbial Contamination of Non-Sterile Drug Product	If you have questions about this recall, MPM Medical LLC, 1-800-232-5512.	December 2020	Class I
Chlorhexidine Gluconate 007016602715	Microbial contamination of non-sterile products	If you have questions about this recall, Lohxa LLC, 1-800-641-5564.	December 2020	Class I
Tizanidine 06787761415	Failed Dissolution Specifications	If you have questions about this recall, Ascend Laboratories LLC, 1-877-272-7901.	December 2020	Class I
Aripiprazole 00904651204	Failed Dissolution Specifications	If you have questions about this recall, The Harvard Drug Group, 1-800-875-0123.	December 2020	Class II
ARIPIPRAZOLE 06042944930	FAILED DISSOLUTION SPECIFICATIONS	If you have questions about this recall, Golden	December 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

		State Medical Supply Inc., 1-805-477-9866.		
Paroex (chlorhexidine gluconate) 05237602102 05237602104	Microbial Contamination of Non-sterile Product	If you have questions about this recall, Sunstar Americas, Inc., 1-609-751-9600.	December 2020	Class I
Levetiracetam 07193006352	Presence of foreign tablet	If you have questions about this recall, Jubilant EYWA PHARMA INC, 1-609-751-9600.	December 2020	Class II
clomiPRAMINE Hydrochloride 05974671130	Failed Tablet	If you have questions about this recall, Jubilant Cadista Pharmaceuticals, Inc., 1-800-308-3985.	December 2020	Class II
Lansoprazole Delayed-Release Orally Disintegrating Tablets 06838277177	Failed Dissolution Specifications	If you have questions about this recall, Zydus Pharmaceuticals, 1-877-993-8779.	November 2020	Class II
Lansoprazole Delayed-Release Orally Disintegrating Tablets 06838277277	Failed Dissolution Specifications	If you have questions about this recall, Zydus Pharmaceuticals, 1-877-993-8779.	November 2020	Class II
Mesalamine Delayed-Release Tablets 00591224522	Failed Dissolution Specifications	If you have questions about this recall, Teva Pharmaceuticals USA, 1-888-838-2872 ext 5	November 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/Drugs/DrugSafetyandAvailability/DrugRecalls) for additional information.

metformin HCL ER 750 mg 04306390230 04306390260	CGMP Deviations	If you have questions about this recall, PD-Rx Pharmaceuticals, Inc., 1-800-299-7379	November 2020	Class II
metformin HCL ER 500 mg tablets 07278900930 07278900960 07278900993 07278900990	CGMP Deviations	If you have questions about this recall, PD-Rx Pharmaceuticals, Inc., 1-800-299-7379	November 2020	Class II
Metformin Hydrochloride Extended-Release Tablets 02903305501	CGMP Deviations	If you have questions about this recall, Nostrum Laboratories Inc, 1-816-841-4636.	November 2020	Class II
Metformin Hydrochloride Extended-Release Tablets 02903305601	CGMP Deviations	If you have questions about this recall, Nostrum Laboratories Inc, 1-816-841-4636.	November 2020	Class II
Chlorhexidine Gluconate 20% 05329600120	Discoloration	If you have questions about this recall, Medichem S.A., 1-201-420-1800	November 2020	Class II
Metformin Hydrochloride Extended Release Tablets 04948362401	CGMP Deviations	If you have questions about this recall, Medichem S.A., 631-753-9090; ext. 160	November 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Metformin Hydrochloride - Extended-Release Tablets 04948362309 04948362301 04948362350 04948362310 04948362350	CGMP Deviations	If you have questions about this recall, Marksans Pharma Limited, 631-753-9090; ext. 160	November 2020	Class II
Metformin HCL E/R 500 mg 07218906490	CGMP Deviations	If you have questions about this recall, Direct Rx, 1-678-619-5510	November 2020	Class II
Metformin Hydrochloride Extended-Release Tablets, USP 500 mg 07093430930 07093430960 07093430990 07093430998	CGMP Deviations	If you have questions about this recall, Denton Pharma, Inc., 1-800-722- 0772.	November 2020	II
Metformin Hydrochloride Extended-Release Tablets 07093433430	CGMP Deviations	If you have questions about this recall, Denton Pharma, Inc., 1-800-722- 0772.	November 2020	II
Mesalamine Delayed-Release Tablets 04229156412	Failed Dissolution Specifications	If you have questions about this recall, AVKARE Inc., 1-931-292-6222.	November 2020	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Metformin Hydrochloride Extended Release 750 mg 70518248000	CGMP Deviations	If you have questions about this recall, RemedyRepack Inc., 1- 866-845-3791	November 2020	II
Hydrocortisone butyrate Cream, 0.1% 06868227015	SUBPOTENT DRUG	If you have questions about this recall, Boehringer Ingelheim Pharmaceuticals, Inc., 1- 908-927-1400	November 2020	II
Catapres (clonidine hydrochloride, USP) 0.3 mg 00597001101	An extraneous peak was observed for dissolution testing.	If you have questions about this recall, Boehringer Ingelheim Pharmaceuticals, Inc., 1- 800-243-0127	November 2020	II
Catapres (clonidine hydrochloride, USP) 0.2 mg 00597000701	An extraneous peak was observed for dissolution testing.	If you have questions about this recall, Boehringer Ingelheim Pharmaceuticals, Inc., 1- 800-243-0127	November 2020	II
Catapres (clonidine hydrochloride, USP) 0.1 mg 0597000601	An extraneous peak was observed for dissolution testing.	If you have questions about this recall, Boehringer Ingelheim Pharmaceuticals, Inc., 1- 800-243-0127	November 2020	II
NP Thyroid 120, Thyroid Tablets, USP 2 grain (120 mg) 04219232701	Subpotent Drug	If you have questions about this recall, Acella Pharmaceuticals, LLC 1- 800-541-4802 ext 1,	October 2020	I

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

NP Thyroid 120, Thyroid Tablets, USP 2 grain (120 mg) 04219232801	Subpotent Drug	If you have questions about this recall, Acella Pharmaceuticals, LLC 1-800-541-4802 ext 1,	October 2020	I
Losartan Pot/HCTZ 50/12.5 mg 06191904090	CGMP Deviations	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	October 2020	II
RIOMET ER (metformin hydrochloride for extended-release oral suspension) 01063101917	CGMP Deviations	If you have questions about this recall, Sun Pharmaceutical Industries, Inc., 1-800-818-4555	October 2020	II
Eye Itch Relief, Ketotifen Fumarate Ophthalmic Solution 0.035% 05977992001	CGMP Deviations	If you have questions about this recall, Akorn, Inc, 1-800-932-5676.	October 2020	II
Potassium Chloride Extended-Release Tablets 06438086006	Failed Dissolution Specifications	If you have questions about this recall, Strides Inc. 1-609-773-5000.	October 2020	II
Nature-Throid, 1.5 Grain, (97.5 mg), (Thyroid U.S.P. 1.5 gr. (97.5mg)/Liothyronine (T3) 13.5mcg/Levothyroxine (T4) 57mcg 68788760509	Subpotent Drug	If you have questions about this recall, Preferred Pharmaceuticals, Inc. 1-714-777-3729.	September 2020	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Nature-Throid, 1 Grain, 65 mg (Thyroid U.S.P. 1 gr. (65mg)/Liothyronine (T3) 9mcg/Levothyroxine (T4) 38mcg 68788760409 68788760401	Subpotent Drug	If you have questions about this recall, Preferred Pharmaceuticals, Inc. 1-714-777-3729.	September 2020	II
Nature-Throid, 3/4 Grain (48.75 mg) Thyroid U.S.P. 3/4 gr. (48.75 mg)/Liothyronine (T3) 6.75mcg/Levothyroxine (T4) 28.5mcg 68788686009	Subpotent Drug	If you have questions about this recall, Preferred Pharmaceuticals, Inc. 1-714-777-3729.	September 2020	II
Nature-Throid, 1/2 Grain, 32.5 mg (Thyroid U.S.P. 1/2 gr. (32.5 mg)/Liothyronine (T3) 4.5mcg/Levothyroxine (T4) 19mch 68788928301	Subpotent Drug	If you have questions about this recall, Preferred Pharmaceuticals, Inc. 1-714-777-3729.	September 2020	II
WP Thyroid, Westthroid Pure, 2 Grain (130 mg) Thyroid USP 06472759502 06472759504 06472759505 06472759506 06472759501	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II
WP Thyroid, Westthroid Pure, 1.75 Grain (113.75 mg) Thyroid USP 06472761502	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

06472761504 06472761505 06472761506 06472761501				
WP Thyroid, Westhroid Pure, 1.5 Grain (97.5 mg) Thyroid USP 06472758502 06472758504 06472758505 06472758506 06472758501	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II
WP Thyroid, Westhroid Pure, 1.25 Grain (81.25 mg) Thyroid USP 06472760502 06472760504 06472760505 06472760506 06472760501	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II
WP Thyroid, Westhroid Pure, 1 Grain (65 mg) Thyroid USP 06472757502 06472757504 06472757505 06472757506 06472757501	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

WP Thyroid, Westhroid Pure, 1/2 Grain (32.5 mg) Thyroid USP 06472755502 06472755504 06472755505 06472755506 06472755501	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II
WP Thyroid, Westhroid Pure, Thyroid USP, [lithothyronine (T3) 2.25 mcg and levothyroxine (T4) 9.5 mcg 06472754504 06472754505 06472754506 06472754501 06472754502	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II
Nature-Throid, Thyroid USP [lithothyronine (T3) 27 mcg and levothyroxine (T4) 114 mcg 06472733124 06472733125 06472733126 06472733121 06472733122	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II
Nature-Throid, Thyroid USP [lithothyronine (T3) 22.5 mcg and levothyroxine (T4) 95 mcg 06472733104 06472733105	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

06472733106 06472733101 06472733102				
Nature-Throid, Thyroid USP [lithyronine (T3) 20.25 mcg and levothyroxine (T4) 85.5 mcg 06472733094 06472733095 06472733096 06472733091 06472733092	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II
Nature-Throid, Thyroid USP [lithyronine (T3) 18 mcg and levothyroxine (T4) 76 mcg 06472733084 06472733085 06472733086 06472733081 06472733082	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II
Nature-Throid, Thyroid USP [lithyronine (T3) 15.75 mcg and levothyroxine (T4) 66.5 mcg 06472733074 06472733075 06472733076 06472733071 06472733072	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Nature-Throid, Thyroid USP [liothyronine (T3) 13.5 mcg and levothyroxine (T4) 57 mcg 06472733054 06472733055 06472733056 00472733051 06472733052	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II
Nature-Throid, Thyroid USP [liothyronine (T3) 11.25 mcg and levothyroxine (T4) 47.5 mcg], 1.25 Grain (81.25 mg) 06472733034 06472733035 06472733036 06472733031 06472733032	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II
Nature-Throid, Thyroid USP [liothyronine (T3) 4.5 mcg and levothyroxine (T4) 19 mcg 06472732994 06472732995 06472732996 06472732991 06472732992	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Nature-Throid, Thyroid USP [liothyronine (T3) 6.75 mcg 06472733024 06472733025 06472733026 06472733021 06472733022	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II
Nature-Throid, Thyroid USP [liothyronine (T3) 9 mcg and levothyroxine (T4) 38 mcg 06472733004 06472733005 06472733006 06472733001 06472733002	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	II
Nature Throid, 1.5 Grain (97.5 mg) Thyroid USP [Liothyronine (T3) 13.5 mcg, Levothyroxine (T4) 57 mcg 06472733054 06472733055 06472733056 06472733051 06472733051	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	I
Nature Throid, 2 Grain (130 mg) Thyroid USP [Liothyronine (T3) 18 mcg, Levothyroxine (T4) 76 mcg 06472733084	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	I

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

06472733081				
Nature Throid, 1.25 Grain (81.25 mg) Thyroid USP [Liothyronine (T3) 11.25 mcg, Levothyroxine (T4) 47.5 mcg 06472733031	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	I
Nature Throid, 3/4 Grain (48.75 mg) Thyroid USP [Liothyronine (T3) 6.75 mcg, Levothyroxine (T4) 28.5 mcg 06472733021 06472733022	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	I
Nature Throid, 1/2 Grain (32.5 mg) Thyroid USP [Liothyronine (T3) 4.5 mcg, Levothyroxine (T4) 19 mcg 06472732996 06472732991	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	I
WP Thyroid, Westthroid Pure, 1/2 Grain (32.5 mg) Thyroid USP [Liothyronine (T3) 4.5 mcg, Levothyroxine (T4) 19 mcg 06472755504 06472755506 06472755501	Subpotent Drug	If you have questions about this recall, RLC Labs Inc., 1-877-797-7997	September 2020	I

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Sulfamethoxazole and Trimethoprim Tablets	Presence of Foreign Substance: product complaints were received by the firm for the presence of metal wire in the tablet(s).	If you have questions about this recall, Aurobindo Pharma USA Inc., 1-866-850-2876	September 2020	II
BusPIRone Hydrochloride Tablets	Failed Impurity /Degradation Specifications	If you have questions about this recall, Par Pharmaceutical Inc., 1-845-573-5500	September 2020	II
Buprenorphine HCl Injection	Sub-potent Drug: Out-of-Specification assay results found at 3-month stability testing.	If you have questions about this recall, West-Ward Pharmaceuticals , 1-877-845-0689.	September 2020	II
Metformin Hydrochloride Extended-Release Tablets 07638512810	CGMP Deviations	If you have questions about this recall, BAYSHORE PHARMACEUTICALS, 1-877-372-6093	September 2020	II
Metformin Hydrochloride Extended-Release Tablets 07638512901	CGMP Deviations	If you have questions about this recall, BAYSHORE PHARMACEUTICALS, 1-877-372-6093	September 2020	II
Cephalexin for Oral Suspension 00093417773	Labeling: Label Error on Declared Strength	If you have questions about this recall, Teva Pharmaceuticals USA, 1-888-838-2872 ext 5	September 2020	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/Drugs/DrugSafetyandAvailability/DrugRecalls) for additional information.

Lidocaine Patch 5% 00591352530	Labeling: Incorrect or Missing Lot and/or Exp date on the individual transdermal pouches but not in the carton	If you have questions about this recall, Teva Pharmaceuticals USA, 1-888-838-2872 ext 5	August 2020	II
Mibelas 24 Fe 06818091113	Failed Impurities/Degradation Specifications: Out of specification result observed in related substance test	If you have questions about this recall, Lupin Pharmaceuticals Inc. 1-410-576-2000 ext 3	August 2020	II
Nystatin Oral Suspension 00121061016	Subpotent drug: Out of specification for assay at the 15-month test interval	If you have questions about this recall, PAI Holdings, LLC. dba Pharmaceutical Associates Inc 1-864-277-7282 ext 0	August 2020	II
Ear Pain MD Pain Relief Drops 07242900708	cGMP Deviations.	If you have questions about this recall, Eosera, Inc. 1-844-732-7929	August 2020	II
Ear Pain MD Pain Relief Drops 07242900722	cGMP Deviations.	If you have questions about this recall, Eosera, Inc. 1-844-732-7929	August 2020	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Heparin Sodium 70004065046	Subpotent Drug: Out-of-Specification potency results at the 30-day stability timepoint.	If you have questions about this recall, SCA Pharmaceuticals 1-877-550-5059	August 2020	II
BD Chloraprep Clear 5436540032213	Non-Sterility: Product is being recalled due to presence of <i>Aspergillus Penicilloides</i>	If you have questions about this recall, CareFusion LLC 1-201-847-6800.	August 2020	I
Chloraprep With Tint 05436540011	Non-Sterility: Product is being recalled due to presence of <i>Aspergillus Penicilloides</i>	If you have questions about this recall, CareFusion LLC 1-201-847-6800.	August 2020	I
Chloraprep One-Step 05436540001	Non-Sterility: Product is being recalled due to presence of <i>Aspergillus Penicilloides</i>	If you have questions about this recall, CareFusion LLC 1-201-847-6800.	August 2020	I

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

DDAVP Nasal 05556625000	Superpotent Drug	If you have questions about this recall, Ferring Pharmaceuticals Inc1-888-337-7464 ext2	August 2020	I
Desmopressin Acetate 06991850105	Superpotent Drug	If you have questions about this recall, Ferring Pharmaceuticals Inc1-888-337-7464 ext2	August 2020	I
STIMATE (desmopressin acetate) 00053687100	Superpotent Drug	If you have questions about this recall, Ferring Pharmaceuticals Inc1-888-337-7464 ext2	August 2020	I
Hydrochlorothiazide 16729018317	CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches	If you have questions about this recall, Calvin Scott & Company, Inc.1-800-545-6545.	August 2020	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Hydrochlorothiazide (HCTZ) 29300012810	CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches	If you have questions about this recall, Calvin Scott & Company, Inc.1-800-545-6545.	August 2020	II
Hydrochlorothiazide 16729018417	CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches	If you have questions about this recall, Calvin Scott & Company, Inc.1-800-545-6545.	August 2020	II
Thyroid 1 Grain 64727330002	CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches	If you have questions about this recall, Calvin Scott & Company, Inc.1-800-545-6545.	August 2020	II
Thyroid 1 Gr 64727330001	CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches	If you have questions about this recall, Calvin Scott & Company, Inc.1-800-545-6545.	August 2020	II
Thyroid Neutral 64727330802	CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches	If you have questions about this recall, Calvin Scott & Company, Inc.1-800-545-6545.	August 2020	II
Topiramate 47335070713	CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches	If you have questions about this recall, Calvin Scott & Company, Inc.1-800-545-6545.	August 2020	II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Phendimetrazine 00702007710	CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches	If you have questions about this recall, Calvin Scott & Company, Inc.1-800-545-6545.	August 2020	II
Phendimetrazine 69543041011	CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches	If you have questions about this recall, Calvin Scott & Company, Inc.1-800-545-6545.	August 2020	II
Phendimetrazine 69543040911	CGMP Deviations: Light sensitive drug products repackaged in transparent/partially transparent pouches	If you have questions about this recall, Calvin Scott & Company, Inc.1-800-545-6545.	August 2020	Class II
Prednisone 00527293137	Labeling: Label Mix Up: bottle labeled to contain Prednisone Tablets	If you have questions about this recall, Lannett Company, Inc.1-215-333-9000	August 2020	Class II
Elitek (rasburicase) 00024515175	Failed Stability Specifications: Out of Specification result for enzyme activity levels noted during routine stability testing	If you have questions about this recall, .Sanofi-Aventis U.S. LLC1-800-981-2491 ext 2	August 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/Drugs/DrugSafetyandAvailability/DrugRecalls) for additional information.

Mibelas 24 Fe 06818091111	Failed Impurities/Degradation Specifications: Out of specification result observed in related substance test	If you have questions about this recall, Lupin Pharmaceuticals Inc. 1- 410-576-2000 ext 3	August 2019	Class II
Daptomycin 06745781350	Presence of Particulate Matter	If you have questions about this recall, Mylan Institutional LLC, 1-800- 796-9526 ext 1.	August 2020	Class II
Fentanyl Citrate 00409909412	Lack of Assurance of Sterility	If you have questions about this recall, Pfizer Inc., 1-800-438-1985.	August 2020	Class II
Lisinopril 06800133408	Presence of Foreign Tablets/Capsules	If you have questions about this recall, Lupin Pharmaceuticals Inc., 1- 866-587-4617 ext 2.	August 2020	Class II
Cefdinir 06818072320	Superpotent Drug	If you have questions about this recall, Lupin Pharmaceuticals Inc., 1- 866-587-4617 ext 2.	July 2020	Class II
Metformin HCl Extended Release Tablets	CGMP Deviation	If you have questions about this recall, Preferred Pharmaceuticals, Inc, 1- 714-777-3729.	July 2020	Class II
Metformin Hydrochloride Extended-Release 00904579461	CGMP Deviation	If you have questions about this recall, The Time-Cap, Labs Inc, 1- 631-753-9090 ext 160	July 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Clozapine Tablets 06586284605	Presence of foreign tablet	If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876	July 2020	Class II
Metformin Hydrochloride Extended-release Tablets 06818033607	CGMP Deviations	If you have questions about this recall, Lupin Pharmaceuticals Inc., 1-866-587-4617.	July 2020	Class II
Aripiprazole Tablets 06233209730	Labeling: Label mix up	If you have questions about this recall, Alembic Pharmaceuticals Limited, 1-908-393-9604.	July 2020	Class II
Nystatin Cream 00316022130 00316022115	Subpotent Drug	If you have questions about this recall, Crown Laboratories, Inc., 1-800-877-8869.	July 2020	Class II
metFORMIN HCL ER 07278900930 07278900960 07278900990 07278900993 49483062301	CGMP Deviations	If you have questions about this recall, Amneal Pharmaceuticals., 1-833-582-0812.	July 2020	Class II
Metformin Hydrochloride Extended-Release Tablets 04948362301	CGMP Deviations	If you have questions about this recall, Time-Cap Labs, 1-631-753-9090 ext 160	July 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/Drugs/DrugSafetyandAvailability/DrugRecalls) for additional information.

Metformin Hydrochloride Extended - Release Tablets 06203757101 06203757110 06203757701 06203757710	CGMP Deviations	If you have questions about this recall, Teva Pharmaceuticals USA, 1- 800-545-8800.	July 2020	Class II
Metformin Hydrochloride Extended-Release Tablets 53746017801 06516217809 06516217850 06516217810 53746017901 06516217910 00537460178 53746017805 53746017810 65162017811 00537460179 65162017901	CGMP Deviations	If you have questions about this recall, Amneal Pharmaceuticals, 1-833- 582-0812	July 2020	Class II
Heparin Sodium 70004065046	Subpotent Drug	If you have questions about this recall, SCA Pharmaceuticals, 1-877- 550-5059.	June 2020	Class II
Irinotecan HCL Injection 00143958301	Defective Container	If you have questions about this recall, West- Ward Pharmaceutical Corp, 1-877-845-0689.	June 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Irinotecan HCL Injection 00143970101	Defective Container	If you have questions about this recall, West-Ward Pharmaceutical Corp, 1-877-845-0689.	June 2020	Class II
metFORMIN HCL ER 04306342830 04306342860 04306342890 04306342898 04306342893 53746017805	CGMP Deviations	If you have questions about this recall, Amneal Pharmaceuticals, 1-833-582-0812	June 2020	Class II
Metformin Hydrochloride Extended-Release Tablets 05026853115 04229161090 04229161018 04229161036 04229161010	CGMP Deviations	If you have questions about this recall, AVKARE Inc., 1-931-292-6222.	June 2020	Class II
Metformin Hydrochloride Extended-Release Tablets 04229161190 04229161118 04229161150	CGMP Deviations	If you have questions about this recall, AVKARE Inc., 1-931-292-6222.	June 2020	Class II
Metformin Hydrochloride Extended-Release Tablets 06050502601	CGMP Deviations	If you have questions about this recall, Apotex Inc., 1-800-268-4623.	June 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Gaviscon Liquid Antacid Extra Strength, Cool Mint 00135009541	Labeling: Label lacks warning	If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings Inc., 1-800-743-4014	June 2020	Class II
Gaviscon Extra Strength Liquid Antacid Extra Strength Cherry 00135057441	Labeling: Label lacks warning	If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings Inc., 1-800-743-4014	June 2020	Class II
Gaviscon Regular Strength Liquid Antacid Cool Mint 00135009441	Labeling: Label lacks warning	If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings Inc., 1-800-743-4014	June 2020	Class II
Gaviscon Regular Strength Liquid Antacid Cool Mint 00135009442	Labeling: Label lacks warning	If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings Inc., 1-800-743-4014	June 2020	Class II
oxyTOCIN 30 Units 070092106807	Subpotent drug	If you have questions about this recall, QuVa Pharma, Inc., 1-888-339-0874.	June 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Lisinopril 06818051303	Product Mix Up	If you have questions about this recall, Lupin Pharmaceuticals Inc., 1-866-587-4617.	June 2020	Class II
Unasyn (ampicillin sodium/sulbactam) 00049001381 00049001383	Presence of Particulate Matter	If you have questions about this recall, Pfizer Inc., 1-800-879-3477.	June 2020	Class II
Doxycycline Hyclate 06258469311	Failed dissolution specification	If you have questions about this recall, American Health Packaging, 1-614-492-8177.	June 2020	Class II
Ketorolac Tromethamine 06332316200 06332316203	Presence of Particulate Matter	If you have questions about this recall, Fresenius Kabi USA, LLC, 1-847-550-2300.	June 2020	Class II
Doxycycline Hyclate 00904043004 00904043006	Failed Dissolution Specification	If you have questions about this recall, The Harvard Drug Group, 1-732-542-1191.	June 2020	Class II
Dextroamphetamine Saccharate, Amphetamine Aspartate, Dextroamphetamine Sulfate and Amphetamine Sulfate 00555097102 00555097302 00555077702	Some bottles may contain mixed strengths of the product.	If you have questions about this recall, Teva Pharmaceuticals USA, 1-800-545-8800.	June 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Estriol USP Micronized 71092997702	Failed impurities/ degradation specifications	If you have questions about this recall, TRIOVA PHARMACEUTICALS LLC, 1-539-777-0720.	June 2020	Class II
NP Thyroid 90 04219233101	Superpotent Drug	If you have questions about this recall, Acella Pharmaceuticals, LLC 1- 800-541-4802.	June 2020	Class I
NP Thyroid 60 04219233001	Superpotent Drug	If you have questions about this recall, Acella Pharmaceuticals, LLC 1- 800-541-4802.	June 2020	Class I
NP Thyroid 30 04219232901	Superpotent Drug	If you have questions about this recall, Acella Pharmaceuticals, LLC 1- 800-541-4802.	June 2020	Class I
Doxycycline Hyclate tablets 05528986606 05528986610 05528986614 05528986620 05528986628 05528986630 05528986660 05528986698 05528986671 05528986687 05528986674	Failed dissolution specifications	If you have questions about this recall, PD-Rx Pharmaceuticals, Inc., 1- 800-299-7379.	June 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Doxycycline Hyclate Tablets 00143211250 00143211205	Failed dissolution specification	If you have questions about this recall, West-Ward Columbus Inc, 1-877-845-0689.	June 2020	Class II
Lactated Ringers 0409795309	Presence of Particulate Matter	If you have questions about this recall, Hospira Inc., 1-877-946-7747.	May 2020	Class II
Aloprim (allopurinol sodium) 06745718750	Discoloration	If you have questions about this recall, Mylan Pharmaceuticals Inc., 1-551-233-2700.	May 2020	Class II
Infuvite PEDIatric Pharmacy 05464356470	Defective container	If you have questions about this recall, Sandoz, Inc, 1-609-627-8500	May 2020	Class II
Epinephrine Injection 00093598627	CGMP Deviations	If you have questions about this recall, Teva Pharmaceuticals USA, 1-888-838-2872.	May 2020	Class II
Nizatidine 06084630115	CGMP Deviations	If you have questions about this recall, Amneal Pharmaceuticals of New York, 1-631-952-0214	May 2020	Class II
Ceftazidime 00264314511	Failed Stability Specifications	If you have questions about this recall, Braun Medical Inc., 1-949-660-2000	May 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/Drugs/DrugSafetyandAvailability/DrugRecalls) for additional information.

Lisinopril Tablets 68180098103	Product Mix-Up	If you have questions about this recall, Lupin Pharmaceuticals Inc., 1-866-587-4617	April 2020	Class II
Cefixime for Oral Suspension 06818040501	Subpotent Drug	If you have questions about this recall, Lupin Pharmaceuticals Inc., 1-866-587-4617	April 2020	Class II
Daytrana 06896855553	Defective Delivery System	If you have questions about this recall, Noven Therapeutics, LLC, 1-305-253-5099.	April 2020	Class II
Daytrana 06896855543	Defective Delivery System	If you have questions about this recall, Noven Therapeutics, LLC, 1-305-253-5099.	April 2020	Class II
Daytrana 06896855523	Defective Delivery System	If you have questions about this recall, Noven Therapeutics, LLC, 1-305-253-5099.	April 2020	Class II
Acetaminophen and Codeine Phosphate 01310705801	CGMP Deviations	If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876.	April 2020	Class II
Acetaminophen and Codeine Phosphate 01310705999	CGMP Deviations	If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876.	April 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Acetaminophen and Codeine Phosphate 01310706001	CGMP Deviations	If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876.	April 2020	Class II
Gabapentin Capsules 06586219899	CGMP Deviations	If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876.	April 2020	Class II
Levetiracetam Tablet 06586224708	CGMP Deviations	If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876.	April 2020	Class II
Simvastatin 06586205390	CGMP Deviations	If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876.	April 2020	Class II
Mirtazapine 01310703134 06586219899	CGMP Deviations	If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876.	April 2020	Class II
Phentermine Hydrochloride 06586205390 06586219899	CGMP Deviations	If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876.	April 2020	Class II
Oxycodone and Acetaminophen 01310704601 6586219899	CGMP Deviations	If you have questions about this recall, Aurobindo Pharma USA, Inc, 1-866-850-2876.	April 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Losartan Potassium 06042931630 06042931690 06042931610	CGMP Deviations	If you have questions about this recall, Golden State Medical Supply Inc., 1-805-477-9866 Ext 4.	April 2020	Class II
Estradiol 06327599008 06327599005 06327599004	CGMP Deviations	If you have questions about this recall, B & B Pharmaceuticals, Inc., 1-800-499-3100.	April 2020	Class II
Acetaminophen and Codeine Phosphate 300/30mg 52959000310 52959000312 52959000314 52959000315 52959000316 52959000320 52959000330 52959000360	CGMP Deviations	If you have questions about this recall, H.J. Harkins Co, 1-805-929-1333	April 2020	Class II
succinylcholine Chloride 07101934104	lack of sterility assurance.	If you have questions about this recall, PharMEDium Services, 1-800-523-7749 Option 1	April 2020	Class II
rocuronium Bromide 07101932110 07101932105	lack of sterility assurance.	If you have questions about this recall, PharMEDium Services, 1-800-523-7749 Option 1	April 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Phytonadione Injectable Emulsion 04359840511	Defective Container	If you have questions about this recall, Dr. Reddy's Laboratories Inc., 1-888-375-3784	April 2020	Class I
nICARdipine HCl 07101920601	lack of sterility assurance.	If you have questions about this recall, PharMEDium Services, 1-800-523-7749 Option 1	April 2020	Class II
Losartan Potassium tablets 00591374500 00591374600 00591374700	CGMP Deviations	If you have questions Teva Pharmaceuticals USA, 1-888-483-8279	April 2020	Class II
Losartan Potassium tablets 02315564503 02315564509 02315564510 02315564509 02315564603 02315564609 02315564610	CGMP Deviations	If you have questions Avet Pharmaceuticals, Inc., 1-800-967-5952 Option 1	April 2020	Class II
Lisinopril Tablets 06818098201	Presence of Foreign Tablet/ Capsule	If you have questions about this recall, Lupin Pharmaceuticals Inc., 1-800-399-2561	April 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

HYDROmorphine in 0.9% Sodium Chloride HCl 06155316644	lack of sterility assurance.	If you have questions about this recall, PharMEDium Services, 1-800-523-7749 Option 1	April 2020	Class II
Glycopyrrolate Tablets 49884006501	lack of sterility assurance.	If you have questions about this recall, Par Pharmaceutical Inc., 1-845-573-5500	April 2020	Class II
fentaNYL Citrate Injection 06155330633 06155330328	lack of sterility assurance.	If you have questions about this recall, PharMEDium Services, LLC, 1-800-523-7749 Option 1	April 2020	Class II
fentaNYL Citrate 06155311350 06155367244	lack of sterility assurance.	If you have questions about this recall, PharMEDium Services, LLC, 1-800-523-7749 Option 1	April 2020	Class II
ePHEDrine Sulfate 07103000312 07103000309 07103000302 07103000109	lack of sterility assurance.	If you have questions about this recall, PharMEDium Services, LLC, 1-800-523-7749 Option 1	April 2020	Class II
C-*Vancomycin Opthal 14 mg drops 67457034001	Lack of Assurance of Sterility	If you have questions about this recall, Medical Center Pharmacy, Inc., 1-800-523-7749	April 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

C-*Gentamicin/Bacitracin Bladder Irrigation in N.S., 250 mL 63323001002	Lack of Assurance of Sterility	If you have questions about this recall, Medical Center Pharmacy, Inc., 1-800-523-7749	April 2020	Class II
C-*Albumin Eye Drop 10% S 44206025105	Lack of Assurance of Sterility	If you have questions about this recall, Medical Center Pharmacy, Inc., 1-800-523-7749	April 2020	Class II
*Morphine 2 mg/mL Cassette 00409113403	Lack of Assurance of Sterility	If you have questions about this recall, Medical Center Pharmacy, Inc., 1-800-523-7749	April 2020	Class II
*Mitomycin 0.04% Ophth DR eye drops 067457051805	Lack of Assurance of Sterility	If you have questions about this recall, Medical Center Pharmacy, Inc., 1-800-523-7749	April 2020	Class II
Nystatin Oral Suspension 00 21081016	SubPotent Drug: Low out-of-specification results for assay testing.	If you have questions about this recall, PAI Holdings, LLC. dba Pharmaceutical Associates Inc, 1-864-277-7282.	April 2020	Class II
Theophylline (Anhydrous) Extended-Release 02903300101	CGMP Deviations: poor manufacturing practices resulted in Labeling	If you have questions about this recall, Jubilant Draximage Inc, 1-800-361-2356.	April 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

DRAXIMAGE DTPA (KIT FOR THE PREPARATION OF TECHNETIUM TC 99M PENTETATE INJECTION) 06517428805 06717528830	Failed Stability Specifications	If you have questions about this recall, Nostrum Laboratories Inc, 1-816-841-4636.	April 2020	Class II
Advil Liqui-Gel Mini 160+20+20 CT 00573171559	Labeling: Lacks Warning or Rx Legend- Certain lots does not include required safety warning information in the Drug Facts Panel.	If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings, +44 20 8047 5000.	April 2020	Class II
Ibuprofen 50 mg per 1.25 mL Oral Suspension Advil Infant Concentrated Drops White Grape 00573019175 00573019150	Labeling: Lacks Warning or Rx Legend- Certain lots does not include required safety warning information in the Drug Facts Panel.	If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings, +44 20 8047 5000.	April 2020	Class II
Ibuprofen 200 mg liquid filled capsules Advil Liqui-Gel Minis 00573176989 00573176995	Labeling: Lacks Warning or Rx Legend- Certain lots does not include required safety warning information in the Drug Facts Panel.	If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings, +44 20 8047 5000.	April 2020	Class II
Advil Allergy & Congestion Relief 00573019610	Labeling: Lacks Warning or Rx Legend- Certain lots does not include required safety warning information in the Drug Facts Panel.	If you have questions about this recall, Glaxosmithkline Consumer Healthcare	April 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

		Holdings, +44 20 8047 5000.		
Phenylephrine 10 mg tablets. 00573019620	Labeling: Lacks Warning or Rx Legend- Certain lots does not include required safety warning information in the Drug Facts Panel.	If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings, +44 20 8047 5000.	April 2020	Class II
Ibuprofen 200 mg Chlorpheniramine Maleate 4mg 00573019610	Labeling: Lacks Warning or Rx Legend- Certain lots does not include required safety warning information in the Drug Facts Panel.	If you have questions about this recall, Glaxosmithkline Consumer Healthcare Holdings, +44 20 8047 5000.	April 2020	Class II
Doxycycline 06330461501 06330461650	CGMP Deviations: During manufacturing Solifenacin Succinate Tablets might convert to Solifenacin Tartrate Tablets.	If you have questions about this recall, Sun Pharmaceutical Industries, Inc., 1-800-818-4555.	March 2020	Class II
Solifenacin Succinate 05199189333 05199189390	CGMP Deviations: During manufacturing Solifenacin Succinate Tablets might convert to Solifenacin Tartrate Tablets.	If you have questions about this recall, Breckenridge Pharmaceutical, Inc, 1-860-828 - 8140.	March 2020	Class II
Atorvastatin Calcium 06330482905	Presence of foreign substance: Foreign matter has been identified as latex glove in one lot of Atorvastatin Calcium Tablets USP 40 mg.	If you have questions about this recall, Sun Pharmaceutical Industries, Inc., 1-800-818-4555.	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/Drugs/DrugSafetyandAvailability/DrugRecalls) for additional information.

Pantoprazole Sodium Delayed-Release	CGMP Deviations: Presence of dark brown discoloration on edges of tablets	If you have questions about this recall, Jubilant Cadista Pharmaceuticals, Inc., 1-410- 860-2836.	March 2020	Class II
Sotalol HCL 00378512301	Presence of particulate matter. presence of metal particles.	If you have questions about this recall, Mylan Pharmaceuticals Inc., 1-551-233-2700.	March 2020	Class II
Daytrana (methylphenidate transdermal system) 06896855523 06896855533 06896855543 06896855553	Defective Delivery System: Out of specification for mechanical peel and shear.	If you have questions about this recall, Noven Therapeutics, LLC, 1 - 800-796-9526.	March 2020	Class II
Ranitidine 00113785282 05977954082 03780850782 02113011682 00363085282	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II
Ranitidine 05977995001	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II
Ranitidine 00113785201 06839185256 05977954001 05591001101	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

06398185256 03014260056 04125085201 01182208524 00363085201				
Ranitidine 04903580075 01167395075	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II
Ranitidine 04903560875 01167302375	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II
Ranitidine 00113795009 05977995009 04903580009 04612253309 03014289109 03680095009 01167395009 00363095009	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II
Ranitidine 04152039209 05977954009 05531985209 04612253209 03780850709 04934810954	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

0 4125089109 01182208523 03701285209 05059485209 04934810954 01167385209 00363085209				
Ranitidine 05591085271 04612222471 00113085271 06925604171 03014260071 00904671651 04125085271 05606209971 01182208522 02113011671 03680085271	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II
Ranitidine 01167395058	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II
Ranitidine 04125095002 01182209500 03701295062 02113056862	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

03680095062 00363095002				
Ranitidine 04699485262 04152039202 05977954002 05591085202 04116385262 04903560802 05530185202 05531985202 04612222462 00113085262 06925604162 04934810904 03014260002 00904671624 04125085202 05606209902 01182208525 01020285262 04119085262 03701285262 02113011602 05059485202 06201102821 03680085202	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

01167302302 00363085262				
Ranitidine 00113085251 00363085251	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II
Ranitidine 06984229306	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II
Ranitidine 03780887647 04903587647	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II
Ranitidine 03780887647 04903587647	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II
Ranitidine 00113787627 07000003752 01167387627 00363187627	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Ranitidine 06201102832 09046715524 00125025272 04934813612	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II
Ranitidine 06984229365 04116393165 05531987665 00113087665 03780887665 06925687665 06201102831 06201102831 00904671546 02113011865 04934813644 01167387665 00363187665	CGMP Deviation; Possible contamination with impurity N-nitrosodimethylamine.	If you have questions about this recall, Perrigo Company PLC, 1-800-719-9260	March 2020	Class II
Lisinopril/HCTZ 68180051902 70518038203	Presence of Foreign Tablets/Capsules	If you have questions about this recall, RemedyRepack Inc., 1-866-845-3791	March 2020	Class II
Calcium Chloride Injection 07128302253	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Cholecalciferol (Vitamin D3) Injection, VITAMIN D3 [P] 1,000 IU/ML INJECTABLE 07128302103	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Zinc Chloride Injection, ZINC CHLORIDE 10MG/ML INJECTABLE 07128302243	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Cholecalciferol (Vitamin D3) Injection, VITAMIN D3 [P] 100,000 IU/ML INJECTABLE 07128302513	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
ElELYso (taliglucerase alfa) for injection 069010601	Lack of Assurance of Sterility	If you have questions about this recall, Pfizer Labs, 1-877-225-9750	March 2020	Class II
Mesalamine Delayed-Release Tablets 591224522	Failed Dissolution Specifications: Low out of specification dissolution result observed during stability testing.	If you have questions about this recall, Teva Pharmaceuticals USA, Inc, 1-877-685-8222	March 2020	Class II
Triamcinolone Diacetate Injectable Suspension, TRIAMCINOLONE DIACETATE (PF) [2ML] CMC 40 MG/ML INJ SUSP, For IM, IA7 0128306282	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Triamcinolone Diacetate Injectable Suspension, TRIAMCINOLONE DIACETATE [10ML] CMC 80 MG/ML INJ SUSP, For IM, IA 07128306341	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Triamcinolone Diacetate Injectable Suspension, TRIAMCINOLONE Diacetate [10ML] CMC 10MG/ML INJ SUSP, For IM, IA 7128306251	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Triamcinolone Acetonide (PF) Injectable Suspension, TRIAMCINOLONE ACETONIDE (PF) [2ML] 40MG/ML INJ SUSP, For IM, IA7128306352	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Triamcinolone Acetonide (PF) Injectable Suspension, TRIAMCINOLONE ACETONIDE (PF) [10ML] 50MG/ML INJ SUSP, For IM, IA7128306331	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Triamcinolone Acetonide/Bupivacaine Hydrochloride Injectable Suspension, TRIAMCINOLONE ACET/BUPIVACAINE HCL	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

[10ML] 40MG/5MG/ML INJ SUSP, For IM, IA 07128306321				
Testosterone Cypionate Injection, TESTOSTERONE CYP IN GRAPESEED OIL [1ML] 200 MG/ML INJECTABLE 07128305302	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Triamcinolone Acetonide (PF) Injectable Suspension, TRIAMCINOLONE ACETONIDE (PF) [10ML] 50MG/ML INJ SUSP, For IM, IA 07128306331	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Triamcinolone Acetonide (PF) Injectable Suspension, TRIAMCINOLONE ACETONIDE (PF) [2ML] 40MG/ML INJ SUSP, For IM, IA 07128306352	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Triamcinolone Diacetate Injectable Suspension, TRIAMCINOLONE Diacetate [10ML] CMC 10MG/ML INJ SUSP, For IM, IA	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

071283062511				
Testosterone Cypionate/Progesterone Injection, TESTOSTERONE CYP/PROGESTERONE [2ML] 200MG/2.5MG/ML INJECTABLE 7128305312	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Testosterone Cypionate Injection, TESTOSTERONE CYP IN GRAPESEED OIL [10ML] 200 MG/ML INJECTABLE 07128305301	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Selenium Injection, SELENIUM 200MCG/ML INJECTABLE 07128302273	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Super MIC Injection, SUPER MIC* INJECTABLE 07128302343	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Nicotinamide Adenine Dinucleotide (PF) Injection, NICOTINAMIDE ADENINE DINUCLEOTIDE (PF) 20 MG/ML INJECTABLE	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

07128303361				
Iohexol (PF) Injection, OMNIPAQUE INJECTION [5ML] 300MG I/ML INJECTABLE 07128314135	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Pyridoxine Hydrochloride Injection, PYRIDOXINE HCL 100 MG/ML INJECTABLE 07128302163	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Nicotinamide Adenine Dinucleotide (PF) Injection, NICOTINAMIDE ADENINE DINUCLEOTIDE (PF) 50 MG/ML INJECTABLE 07128303061	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
MIC-PLUS Injection, Vitamin Complex, MIC-COMBO* 25MG/50MG/50MG/1MG/20MG/ 5MG/ML INJECTABLE 07128302313	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Mitomycin-C (PF) Irrigation Solution, MITOMYCIN-C (PF) 0.5 MG/ML PF SYRINGE 07128308128	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
MIC-PLEX Injection, Vitamin Complex, MIC-COMBO* 25MG/50MG/50MG/1MG/20MG/5MG/ML INJECTABLE7128302333	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
MIC-B12 Injection, MIC-B12 25MG/50MG/50MG/1MG/ML INJECTABLE7128302303	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Methylprednisolone Acetate Injectable Suspension, METHYLPREDNISOLONE ACETATE [10ML] CMC 50 MG/ML INJ SUSP, For IM, IA 07128306361	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Methylprednisolone Acetate Injectable Suspension, METHYLPREDNISOLONE ACETATE [10ML] CMC 100 MG/ML INJ SUSP 07128306191	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Methylprednisolone Acetate/Bupivacaine Hydrochloride Injectable Suspension 07128306241	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Methylprednisolone Acetate Injectable Suspension, METHYLPREDNISOLONE ACETATE (PF) CMC [2ML] 80 MG/ML INJ SUSP, For IM, IA 07128306392	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Methylprednisolone Acetate/Bupivacaine Hydrochloride Injectable Suspension, METHYLPRED ACETATE/BUPIV [10ML] CMC 40MG/5MG/ML INJ SUSP, For IM, IA 07128306231	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Methylcobalamin Injection, METHYLCOBALAMIN [CD] 10MG/100MG/ML INJECTABLE, For IV, IM 07128303383	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Methylcobalamin Injection, METHYLCOBALAMIN 1 MG/ML INJECTABLE 07128303313	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Lysine Hydrochloride Injection, LYSINE HCL 100MG/ML INJECTABLE 07128303423	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Levocarnitine Injection, LEVOCARNITINE 500 MG/ML INJECTABLE 07128392603	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Lidocaine HCL (PF) Injection, LIDOCAINE HCL 4% (PF) 40 MG/ML INJECTABLE 07128310115	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Polyoxyl Lauryl Ether (Polidocanol) Injection, LAURETH-9 (POLIDOCANOL) 5% INJECTABLE 0712830603	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Hydroxyprogesterone Caproate Injection, HYDROXYPROGESTERONE CAPROATE [4ML] 250 MG/ML INJECTABLE	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

07128305325				
Arginine Hydrochloride Injection, L-ARGININE HCL 100MG/ML INJECTABLE 07128303413	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Human Chorionic Gonadotropin (hCG) Injection, HCG [10ML] 1000 IU/ML INJECTABLE 07128305331	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Glycine Injection, GLYCINE USP 50MG/ML INJECTABLE 07128303443	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Glycerin (PF) Injection, GLYCERIN 99% INJECTABLE 07128304641	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Glutathione Injection, GLUTATHIONE 200MG/ML INJECTABLE 07128312313	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Hydroxocobalamin Injection, HYDROXOCOBALAMIN 5MG/ML INJECTABLE 07128303393	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Glutathione Injection, GLUTATHIONE 200MG/ML INJECTABLE 07128312313	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Dexpanthenol Injection, DEXPANTHENOL 250 MG/ML INJECTABLE 07128302143	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Dexamethasone (LA) Injectable Suspension, DEXAMETHASONE LA [10ML] 16MG INJ SUSP 07128306301	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Dexamethasone (LA) Injectable Suspension, DEXAMETHASONE LA [10ML] 16MG INJ SUSP 07128306301	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Deoxycholic Acid Sodium Injection, DEOXYCHOLIC ACID SODIUM 1.67% INJECTABLE 07128304623	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
Cyanocobalamin Injection 07128303303	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

		Pharmaceuticals, Inc, 1-877-685-8222		
Cyanocobalamin/Folinic Acid Injection, CYANOCOBALAMIN : FOLINIC ACID 2000 MCG/ML: 500MCG/ML INJECTABLE 07128303333	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Injection, COENZYME Q-10 20MG/ML OIL INJ SOLN 07128304111	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Biotin (Vitamin H) Injectable Suspension, BIOTIN 10 MG/ML INJ SUSP, For IM 07128302203	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
BETAMETHASONE ACETATE/BETAMETHASONE (PF) Injectable Suspension, Betamethasone Acetate/Betamethasone (PF) CMC [5ML] 7MG/ML INJ SUSP, For IM, IA 07128306215	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

BETAMETHASONE ACETATE/BETAMETHASONE (PF) Injectable Suspension, Betamethasone Acetate/Betamethasone (PF) CMC [10ML] 7MG/ML INJ SUSP, For IM, IA 07128306211	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
BETAMETHASONE ACETATE/BETAMETHASONE (PF) Injectable Suspension, Betamethasone Acetate/Betamethasone (PF) CMC [2ML] 6MG/ML INJ SUSP, For IM, 07128306202	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
B-COMPLEX 110 INJECTABLE 07128302123	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
AscorbiX (Buffered C) Injection, Ascorbix (50ML) 500MG/ML Injectable0 7128302265	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II
AscorbiX (Buffered C) Injection, Ascorbix (30ML) 500MG/ML Injectable 07128302263	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1- 877-685-8222	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Alprostadil/Papaverine Hydrochloride/Phentolamine Mesylate Injection 07128305462	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Alprostadil/Papaverine Hydrochloride/Phentolamine Mesylate Injection 07128305452	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Alprostadil (prostaglandin E1) 80 MCG/ML Injectable 07128305432	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Alprostadil (prostaglandin E1) 150 MCG/ML Injectable 027128305382	Patent ductus arteriosus	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Alprostadil/Papaverine Hydrochloride/Phentolamine Mesylate/Atropine Sulfate Injection 07128305502	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Alprostadil/Papaverine Hydrochloride/Phentolamine Mesylate/Atropine Sulfate Injection 07128305492	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Alprostadil/Papaverine Hydrochloride/Phentolamine Mesylate/Atropine Sulfate Injection 07128305472	Lack of Assurance of Sterility	If you have questions about this recall, Fusion IV Pharmaceuticals, Inc, 1-877-685-8222	March 2020	Class II
Phenytoin Oral Suspension 05167240691	Resuspension Problems	If you have questions about this recall, Taro Pharmaceuticals U.S.A, 1-866-923-4914.	March 2020	Class II
Glycopyrrolate Tabs 00615817039	Failed Impurities	If you have questions about this recall, NCS Healthcare of Kentucky Inc, 1-270-651-6188	February 2020	Class II
Desmopressin Acetate Tablets 06808460421	GMP Deviations	If you have questions about this recall, American Health Packaging, 1-800-967-5952.	February 2020	Class II
Desmopressin Acetate Tablets 06808460621	GMP Deviations	If you have questions about this recall, American Health Packaging, 1-800-967-5952.	February 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Lamotrigine 05167241311	Cross contamination	If you have questions about this recall, Taro Pharmaceuticals U.S.A, 1-866-923-4914.	February 2020	Class II
Methylphenidate hydrochloride Extended-Release 06203772501	Product bottle may be absent of desiccant.	If you have questions about this recall, Teva Pharmaceuticals, 1-888-838-2872.	February 2020	Class II
Fentanyl Citrate Inj 0409909412	Defective container	If you have questions about this recall, Pfizer Inc, 1-800-879-3477.	February 2020	Class II
Ethacrynate Sodium for Injection 06838224601	cGMP Deviations	If you have questions about this recall, Zydus Pharmaceuticals, 1-877-993-8779.	February 2020	Class II
Caduet (amlodipine besylate/atorvastatin calcium) 00069218030	Defective container	If you have questions about this recall, Pfizer Inc, 1-800-879-3477.	February 2020	Class II
Hydrocortisone and Acetic Acid Otic 05038390110	Subpotent Drug	If you have questions about this recall, TECH PHARMACAL CO., INC, 1-800-932-5676.	February 2020	Class II
Ranitidine Hydrochloride 04306384414, 04306384430, 04306384490, 04306384401	cGMP Deviations	If you have questions about this recall, PD-Rx Pharmaceuticals, Inc., 1-405-942-3040.	February 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see *FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls* for additional information.

Olmesartan Medoxomil Tablets 06787744690	cGMP Deviations	If you have questions about this recall, Ascend Laboratories LLC, 1-201-476-1977.	February 2020	Class II
Atorvastatin Calcium Tablets 07037702711	Presence of Foreign Tablets	If you have questions about this recall, Graviti Pharmaceuticals Private Limited, 040 6815 5555.	February 2020	Class II
Walgreens Acne Cleansing Bar, Benzoyl Peroxide 10% 00363013711	Presence of Foreign Substance	If you have questions about this recall, Shandex Personal Care Manufacturing Inc., 1-613-267-1881	February 2020	Class II
Nystatin Oral Suspension 00121081016	Subpotent	If you have questions about this recall, Pharmaceutical Associates Inc, 1-864-277-7282.	February 2020	Class II
Ranitidine Tablets 70518171400	CGMP Deviation	If you have questions about this recall, RemedyRepack Inc, 1-866-845-3791.	January 2020	Class II
Ranitidine Tablets 06878873883, 06878873889, 06878873881, 06878863821, 06878863823, 06878863829,	Trace amounts of an impurity	If you have questions about this recall, Preferred Pharmaceuticals, Inc, 1-714-777-3729.	January 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

06878863820				
Nizatidine Capsules 00378515091, 00 378530093	Trace amounts of an impurity	If you have questions about this recall, Mylan Pharmaceuticals Inc, 1-304-599-2595.	January 2020	Class II
Ranitidine Tablets 52959050207 52959050214 52959050220 52959050230 52959050260	Trace amounts of an impurity	If you have questions about this recall, H.J. Harkins Company, Inc, 1-800-841-5554.	January 2020	Class II
Ranitidine Tablets 07093401704, 07093401720, 07093401724, 07093401730, 07093401790, 07093428715, 07093428790	Trace amounts of an impurity	If you have questions about this recall, Denton Pharma, Inc., 1-800-722-0772.	January 2020	Class II
Netspot 06948800140	Defective container	If you have questions about this recall, Advanced Accelerator Applications USA, Inc., 1-862-263-0820.	January 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Estriol USP Micronized 62991215906, 62991215903, 62991215905, 62991215902	Subpotent	If you have questions about this recall, Letco Medical LLC, 1-800-239-5288.	January 2020	Class II
Dutasteride 5026828213	cGMP Deviations	If you have questions about this recall, AvKARE, Inc, 1-931-292-6222	January 2020	Class II
Testosterone Cypionate 06275601740,06275601540, 06275601640	cGMP Deviations	If you have questions about this recall, Sun Pharmaceutical Industries, Inc., 1-609-495-2800.	January 2020	Class II
Sumatriptan Succinate 06275652169, 06275652188	Failed Impurities/Degradation Specifications; out-of-specification results obtained for related substance.	If you have questions about this recall, Sun Pharmaceutical Industries, Inc., 1-609-495-2800.	January 2020	Class II
Ranitidine 06255969060, 06255969005	Impurities	If you have questions about this recall, Appco Pharma LLC, 1-732-253-7735.	January 2020	Class II
Ranitidine 03780819601,03780819604	Impurities	If you have questions about this recall, AAA Pharmaceutical, Inc, 1-609-288-6060.	January 2020	Class II
Blisovi Fe 1.5/30 06818086611	Failed tablet	If you have questions about this recall, Lupin Pharmaceuticals Inc, 1-410-576-2000.	January 2020	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Estriol 07642004936,07642004937, 07642004938	Presence of impurities	If you have questions about this recall, Asclemed USA Inc, 1-310-320-0100.	January 2020	Class II
Ranitidine 06846224860,68486224860, 06846224801, 68486224860, 06846224930, 68486224860, 06846224901, 68486224860, 06846224920, 68486224860, 06846224805, 68486224860	Presence of impurities	If you have questions about this recall, Glenmark Pharmaceuticals Inc, 1-888-721-7115	January 2020	Class II
Ranitidine 06220777332	Presence of impurities	If you have questions about this recall, Granules India Limited, 1-877-770-3183.	January 2020	Class II
Myorisan® 0 6174830213	Unit dose mispackaging	If you have questions about this recall, Akorn, Inc, 1-800-932-5676.	January 2020	Class II
Ranitidine 06068726069	Presence of impurities	If you have questions about this recall, American Health Packaging, 1-800-707-4621.	January 2020	Class II
Amantadine Hydrochloride 5974669901	Presence of foreign substance	If you have questions about this recall, Jubilant Cadista Pharmaceuticals, Inc. 1-410-860-8500	December 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

25% Dextrose® 00409177510	Labeling error	If you have questions about this recall, Pfizer Inc1-877-225-9750	December 2019	Class II
Vancomycin Hydrochloride 05515020420	Product discoloration	If you have questions about this recall, AuroMedics Pharma LLC, 888-238-7880.	December 2019	Class II
Lidocaine Hcl 70004072309	Presence of foreign substance	If you have questions about this recall, SCA Pharmaceuticals, LLC, 1-877-550-5059.	December 2019	Class II
Zantac® 06671597362, 06671597363,06671597368, 00597012201 00597012208, 00597012213 00597012237, 00597012240 00597012254, 00597012261 00597012281, 00597012296	Presence of impurities	If you have questions about this recall, Sanofi-Aventis U.S. LLC, 1- 800-633-1610	December 2019	Class II
Regular Strength Zantac® 05026922225, 06775115101, 06775115201, 06775115202, 06815125840	Presence of impurities	If you have questions about this recall, Sanofi-Aventis U.S. LLC, 1- 800-633-1610	December 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Maximum Strength Zantac® 00597012101, 00597012106, 00597012108, 00597012109, 00597012111, 00597012124, 00597012138, 00597012150, 00597012164, 00597012166, 00597012168, 0059701218, 00597012180, 00597012182, 00597012185, 00597012190, 0059701294	Presence of impurities	If you have questions about this recall, Sanofi- Aventis U.S. LLC, 1- 800- 633-1610	December 2019	Class II
Regular Strength Zantac® 04116703000, 04116703001, 04116703003, 04116703005, 04116703006, 04116703007, 04116703008,00526922025	Presence of impurities	If you have questions about this recall, Sanofi- Aventis U.S. LLC, 1- 800- 633-1610	December 2019	Class II
Cool Mint Tablets Maximum Strength Zantac® 00597012006,00597012008, 00597012009,00597012024, 00597012038,00597012050, 00597012076, 00597012078 00597012080,00597012082, 00597012087, 04116703201, 04116703202, 04116703203 04116703204,04116703205 04116703206, 04116703207	Presence of impurities	If you have questions about this recall, Sanofi- Aventis U.S. LLC, 1- 800- 633-1610	December 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Ibuprofen 068788726801	Presence of foreign substance	If you have questions about this recall, Preferred Pharmaceuticals Inc 1-714-777-3729	December 2019	Class II
Lidocaine hcl 05515016505	Presence of foreign substance	If you have questions about this recall, AuroMedics Pharma, 1-888-238-7880	December 2019	Class II
Ranitidine 04229172460, 04229172530,	Presence of impurities	If you have questions about this recall, AvKARE, Inc, 1-931-292-6222	December 2019	Class II
Ranitidine 06516225306, 06516225310, 06516225318, 06516225350, 06516225311, 06516225430, 06516225410, 06516225425, 06516266490, 05374625310, 05374625402	Presence of impurities	If you have questions about this recall, Amneal Pharmaceuticals, Inc., 1-908-947-3120	December 2019	Class II
Ranitidine 068788707803, 068788707806	Presence of impurities	If you have questions about this recall, Preferred Pharmaceuticals Inc 1-714-777-3729	December 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Ranitidine 05591009279	Presence of impurities	If you have questions about this recall, Dolgencorp LLC, Inc, 1-615-855-4000	December 2019	Class II
Ranitidine 05965114460, 059651145-30, 06586243174, 05965114405,	Presence of impurities	If you have questions about this recall, Aurobindo Pharma, 1-866-850-2876	December 2019	Class II
Walgreens Sodium Chloride Ophthalmic Solution 00363019313	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Walgreens Sodium Chloride Ophthalmic Ointment 00363750050	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Walgreen's Lubricant Eye Ointment, Mineral Oil 00363019150	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Equate Support Harmony Lubricant Eye Drops 04903514510	Lack of Assurance of Sterility	If you have questions about this recall, Altaire	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

		Pharmaceuticals, Inc, 1-631-722-5988 Ext 16		
Equate Support Advanced 04903588549	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Equate Support Advanced Lubricating Eye Drops 04903588254	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Equate Support Advanced Lubricant Gel Drops 0903588252	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Equate Sterile Lubricant Styel Ointment 04903587550	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Equate Restore Tears Lubricant Eye Drops Twin Pack 04903518949	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Equate Restore PM Nighttime Lubricant Eye Ointment 04903519150	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Equate Night & Day Restore Tears Lubricant Eye Pack 04903588359	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Equate Eye Allergy Relief Drops 4903587413	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Equate Eye Allergy Relief Drops 04903588713	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Equate Comfort Gel Lubricant Eye Gel 04903519749	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Puralube Petrolatum Ophthalmic Ointment® 00574402535	Lack of Assurance of Sterility	If you have questions about this recall, Altaire	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

		Pharmaceuticals, Inc, 1-631-722-5988 Ext 16		
Polycin (bacitracin zinc and polymyxin B sulfate) Ophthalmic Ointment® 00574402135	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Tetacaine (Tetracaine Hydrochloride) Ophthalmic Solution® 05479950215	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
CVS Health Preservative Free Lubricant Eye Drops® 50428302958	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
TRP Styel Relief® 01731201413	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
TRP Pink Eye Relief® 01731201315	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

TRP Blur Relief 01731200211	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Perrigo Sulfacetamide Sodium Ophthalmic Ointment® 00574419035	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Perrigo Sterile Neo-Polycin HC (neomycin and polymixin B sulfates, bacitracin zinc and hydrocortisone acetate) Ophthalmic® 00574414435	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Perrigo Neo-Polycin neomycin and polymixin B sulfates and bacitracin zinc Ophthalmic Ointment® 0574425035	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Perrigo Neomycin and Polymixin B Sulfates and Dexamethasone Ophthalmic® 0574416035	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Perrigo Bacitracin Ophthalmic Ointment® 0574402235	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Ofloxacin Ophthalmic Solution 05939014005	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
OCuSOFT Tetravisc Tetracaine HCl 0.5% Sterile Anesthetic® 05479950505	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
OCuSOFT Tetravisc Tetracaine HCl 0.5% Sterile Anesthetic 05479950501®	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
OCuSOFT Tetravisc Forte Tetracaine HCl 0.5 % Sterile Anesthetic® 05479950405	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
OCuSOFT Tetravisc Forte (Tetracaine HCl) 0.5% Sterile Anesthetic®	Lack of Assurance of Sterility	If you have questions about this recall, Altaire	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

05479950401		Pharmaceuticals, Inc, 1-631-722-5988 Ext 16		
OCuSOFT Tears Again Sterile Lubricant Eye Drops® 05479990430	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Ocusoft Tears Again Lubricant Eye Drops® 05479990415	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
OCuSOFT Homatropine Hydrobromide Ophthalmic Solution® 05479943105	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Ocusoft Goniosoft Hypromellose 2.5% Ophthalmic Demulcent Solution® 05479950315	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
OCuSOFT Flucaine Proparacaine Hydrochloride and Fluorescein Sodium Ophthalmic Solution® 05479950721	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

OCuSOFT Eye Wash Sterile Isotonic Buffered Solution® 05479956501	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
OCuSOFT Eye Wash Sterile Isotonic 05479956559	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Natural Ophthalmics Women's Tear Stimulation Dry Eye Drops® 06877010315	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Natural Ophthalmics Tear Stimulation Forte Dry Eye Drops® 06877010415	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Natural Ophthalmics Ortho-K Thin Eye Drops® 06877014415	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Natural Ophthalmics Ortho-K Thick Comfort Gel 06877014315	Lack of Assurance of Sterility	If you have questions about this recall, Altaire	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

		Pharmaceuticals, Inc, 1-631-722-5988 Ext 16		
Natural Ophthalmics Cataract Eye Drops® 06877013015	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Natural Ophthalmics Allergy Desensitization Eye Drops® 06877012015	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Nano Tears XP Clear Emollient Lubricant Gel Drops® 05939014351	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Nano Tears TF Clear Emollient Lubricant Gel Drops® 05939014249	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Nano Tears TF Clear Emollient Lubricant Gel Drops® 05939014156	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Lubricant Eye Drops Moisturizing Twin Pack® 00363018549	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
iSolutions NanoTears HA Preservative Free Multi - Dose Lubricant Gel Drops® 05939020810	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
iSolutions Nano Tears XP Clear Emollient Lubricant Gel Drops® 05939014310	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
iSolutions Nano Tears TF Clear Emollient Lubricant Gel Drops® 05939014213	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
iSolutions Nano Tears TF Clear Emollient Lubricant Gel Drops® 05939014152	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
iSolutions Nano Tears MXP Forte Clear Emollient Lubricant Gel Drops®	Lack of Assurance of Sterility	If you have questions about this recall, Altaire	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

05939014756		Pharmaceuticals, Inc, 1-631-722-5988 Ext 16		
iSolutions Nano Tears MXP Forte Clear Emollient Lubricant Gel Drops® 05939014410	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
iSolutions Nano Tears MO Clear Emollient Lubricant Drops® 05939014510	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
iSolutions ActivEyes Nighttime Lubricant Eye Ointment® 05939019050	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
FreshKote Lubricant Eye Drops® 01582110115	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Altaire Sterile Eye Wash® 05939017513, 05939017535, 05939017518	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Altaire Homatropaire Homatropine Hyrdobromide Ophthalmic Solution® 05939019205	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1- 631-722-5988 Ext 16	November 2019	Class II
Altaire Goniotaire Hypromellose 2.5% Opthlamic Demulcent Solution® 05939018213	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1- 631-722-5988 Ext 16	November 2019	Class II
Altaire Fluorescein Sodium with Proparacaine Hydrochloride Ophthalmic Solution 05939020505®	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1- 631-722-5988 Ext 16	November 2019	Class II
Altaire Diclofenac Sodium Ophthalmic Solution® 5939014905	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1- 631-722-5988 Ext 16	November 2019	Class II
Altaire Diclofenac Sodium Ophthalmic Solution® 05939014902	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1- 631-722-5988 Ext 16	November 2019	Class II
Altaire Ciprofloxacin Ophthalmic Solution® 05939021710	Lack of Assurance of Sterility	If you have questions about this recall, Altaire	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

		Pharmaceuticals, Inc, 1-631-722-5988 Ext 16		
Altaire Ciprofloxacin Ophthalmic Solution® 05939021702	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Altaire Ciprofloxacin HCl Ophthalmic Solution® 05939021705	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Altacaine (Tetracaine Hydrochloride) Ophthalmic Solution ® 05939018113	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
ActivEyes Sterile Altalube Ointment® 05939019850	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
ActivEyes Preservative Free Multi-Dose Lubricant Gel Drops® 05939014852	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

ActivEyes Preservative Free Multi-Dose Lubricant Drops® 05939014652	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
ActivEyes Sterile Altalube Ointment® 05939019850	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
ActivEyes Preservative Free Multi-Dose Lubricant Gel Drops® 05939014852	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
ActivEyes Preservative Free Multi-Dose Lubricant Drops® 05939014652	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
ActivEyes Lubricant Eye Ointment® 05939018950	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
ActivEyes Altachlore Solution 05939018313®	Lack of Assurance of Sterility	If you have questions about this recall, Altaire	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

		Pharmaceuticals, Inc, 1-631-722-5988 Ext 16		
ActivEyes Altachlore Sodium Chloride Hypertonicity Ophthalmic Ointment® 05939018450	Lack of Assurance of Sterility	If you have questions about this recall, Altaire Pharmaceuticals, Inc, 1-631-722-5988 Ext 16	November 2019	Class II
Ranitidine 05140709705	impurity detected	If you have questions about this recall, Golden State Medical Supply Inc. 1-805-477-9866	November 2019	Class II
Novitium Pharma Ranitidine® 07095400120 07095400140 07095400210 07095400240	impurity detected	If you have questions about this recall, Novitium Pharma LLC. 1-855-204-1431	November 2019	Class II
Rite Aid Pharmacy Maximum Strength Ranitidine® 01182260522 01182260518	Possible contamination	If you have questions about this recall, Apotex Inc. 1-800-706-5575	November 2019	Class II
Equate Maximum Strength Ranitidine® 04903511706	Possible contamination	If you have questions about this recall, Apotex Inc. 1-800-706-5575	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Rite Aid Pharmacy Maximum Strength Ranitidine® 01182261074	Possible contamination	If you have questions about this recall, Apotex Inc. 1-800-706-5575	November 2019	Class II
Equate Maximum Strength Ranitidine® 04903510007 04903510000	Possible contamination	If you have questions about this recall, Apotex Inc. 1-800-706-5575	November 2019	Class II
Walgreens Regular Strength Wal-Zan 75 Ranitidine® 00363102903	Possible contamination	If you have questions about this recall, Apotex Inc. 1-800-706-5575	November 2019	Class II
Walgreens Maximum Strength Wal-Zan 150 Ranitidine® 00363103007 00363103006	Possible contamination	If you have questions about this recall, Apotex Inc. 1-800-706-5575	November 2019	Class II
Alprazolam tablets 00378400305	Presence of foreign substance	If you have questions about this recall, Mylan Pharmaceuticals. 1-304-599-2595	November 2019	Class II
Ranitidine tablets 00363001061 ,00363001062 ,00363001001, 04903540461, 04903540413,04903540465, 03014250534, 03014250550, 00150062076, 06984287130, 06984287180,06984287137,	Impurity detected in product.	If you have questions about this recall, Dr. Reddy's Laboratories, Inc. 1-309-375-9900	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

03014213130, 06386848230, 06386848260, 04359880862 04359880865, 07171320302 07171320305, 05789671524, 01167384940, 05511113160, 05511140434, 00363013130 00363013180, 06386848024 06386848050, 05789671705				
Ranitidine capsules 05511112960, 05511112905, 05511113030, 05511113001	Impurity detected in product.	If you have questions about this recall, Dr. Reddy's Laboratories, Inc. 1-309-375-9900	November 2019	Class II
Atorvastatin calcium 06050525808	Presence of foreign substance	If you have questions about this recall, Apotex, Inc. 1-800-706-5575	November 2019	Class II
Gatifloxacin Ophthalmic Solution® 06131467225	Labeling error	If you have questions about this recall, Sandoz, Inc. 1-609-627-8500	November 2019	Class II
Neomycin and Polymyxin B Sulfates and Dexamethasone Ophthalmic Suspension 06131463006	Incorrect or missing package insert	If you have questions about this recall, Sandoz, Inc. 1-609-627-8500	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Bimatoprost Ophthalmic Solution 00781620675	Incorrect or missing package insert	If you have questions about this recall, Sandoz, Inc. 1-609-627-8500	November 2019	Class II
Testosterone Cypionate Injection 05253662501	Labeling error	If you have questions about this recall, Arbor Pharmaceuticals. 1-866-284-8792	November 2019	Class II
Ranitidine tablets 05591009279	Empty bottles	If you have questions about this recall, AuroMedeics Pharma LLC.	November 2019	Class II
Ranitidine hydrochloride capsules 04229173550,04229173650	Impurities	If you have questions about this recall, AvKare, Inc. 1-931-292-6222	November 2019	Class II
Estradiol vaginal inserts 06846271188	Impurities	If you have questions about this recall, Glenmark Pharmaceuticals. 1-888-721-7115	November 2019	Class II
Prasugrel tablets® 00378518593	Failed dissolution specification	If you have questions about this recall, Mylan Pharmaceuticals. 1-304-599-2595	November 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Ibuprofen suspension 05167213858,05167221308, 51672138509,05167221301, 6709132104	Presence of foreign substance	If you have questions about this recall, Taro Pharmaceuticals. 1-866-923-4914	November 2019	Class II
Leucovorin Calcium 05074246450	Presence of particulate matter	If you have questions about this recall, Ingenus Pharmaceutical. 1-877-748-1970	October 2019	Class II
Pioglitazone Hydrochloride 03334205407	Superpotent	If you have questions about this recall, Macleods Pharmaceuticals Ltd, 91-22-6676-2800	October 2019	Class II
Dextroamphetamine Sacharate, Amphetamine Aspartate, Dextroamphetamine Sulfate and Amphetamine Sulfate 01310707301	Superpotent Drug: Amphetamine Mixed Salts 20mg have been found to be out of specification for weight and thickness.	If you have questions about this recall, Aurobindo Pharma USA Inc. 1-732-839-9400	October 2019	Class II
Fentanyl Citrate® 49452003206	Potential contamination	If you have questions about this recall, Spectrum Laboratory Products. 1-800-772-8786	October 2019	Class II
Ascorbic Acid 07159150050	Labeling issue	If you have questions about this recall, Atlas	October 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

		Pharmaceutical. 1-410-860-8500		
Pantoprazole Sodium Delayed Release 05974628490	Discoloration on edges of tablets	If you have questions about this recall, Jubilant Cadista Pharmaceuticals, Inc. 1-410-860-8500	October 2019	Class II
Vivitrol® 65757030001	Mislabeled	If you have questions about this recall, Alkermes, Inc. 1-800-848-4876	September 2019	Class II
Relpax® 00049234005, 00049234045	Microbial contamination	If you have questions about this recall, Pfizer Inc 1-877-225-9750	September 2019	Class II
Hydrocortisone cream with aloe, Hydrocortisone 00363062003, 00363062004	Potential microbial contamination	If you have questions about this recall, US Pharmaceuticals Inc. 1-888-337-7464	August 2019	Class II
Lisinopril and hydrochlorothiazide tablets 68180051801, 68180051802	Brownish/blackish stains on the tablets	If you have questions about this recall, Lupin Limited 1-866-587-4617	August 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Nitrofurantoin monohydrate/macrocrystals capsules 47781030301	Failed dissolution specifications	If you have questions about this recall, Alvogen, Inc. 1-866-770-3024	August 2019	Class II
Macrobid Urinary Tract Antibacterial 52427028501	Failed dissolution specifications	If you have questions about this recall, Alvogen, Inc. 1-866-770-3024	August 2019	Class II
Aspirin and extended-release dipyridamole capsules 60687030532	Failed impurities/degradation specifications	If you have questions about this recall, American Health Packaging 1-614-492-8177	August 2019	Class II
Doxycycline hyclate 62584069321	Failed dissolution specifications	If you have questions about this recall, American Health Packaging	August 2019	Class II
Neomycin 3.5 mg/g / polymyxin b10000 usp units/g / dexamethasone 1 mg/g ophthalmic ointment 52959040701	Insufficient quality	If you have questions about this recall, H.J. Harkins Company 1-805-929-1333	July 2019	Class II
Diphenhydramine 49035033002, 49035033096, 49035033045	Contamination	If you have questions about this recall, LNK International 1-631-543-3787	July 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Norethindrone and ethinyl estradiol 68180090313	Failed Impurities	If you have questions about this recall, Lupin Pharmaceuticals 1-800- 399-2561	July 2019	Class II
Milrinone lactate 00409277602, 00409277623	Bags have the potential to leak	If you have questions about this recall, Pfizer Inc. 1-800-879-3477	July 2019	Class II
Fluorouracil injection 63323011710, 63323011719, 63323011728, 63323011761, 63323011761, 63323011769, 63323011718, 63323011720, 63323011751, 63323011759, 63323011768	Glass Particles	If you have questions about this recall, Fresenius Kabi 1-888-386-1300	June 2019	Class I
Heparin Sodium® 00264958720	Not potent	If you have questions about this recall, Braun Medical Inc 1-800-523- 9676	June 2019	Class II
Estradiol vaginal inserts 68462071171, 68462071188	Defective delivery	If you have questions about this recall, Glenmark Pharmaceuticals Inc 1-201-684-8000	June 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Zyflo Cr® 10122090212, 10122090220	Failed dissolution	If you have questions about this recall, Chiesi USA, 1-919-678-6611	June 2019	Class II
Pramipexole dihydrochloride 68084079325, 68084097425	Possible cross contamination	If you have questions about this recall, American Health Packaging, 1-800-707-4621	May 2019	Class II
Zileuton® 66993048532	Failed dissolution	If you have questions about this recall, Chiesi USA, 1-919-678-6611	May 2019	Class II
Testosterone 69699170230, 69699170210	Lack of sterility	If you have questions about this recall, Pharm D. Solutions 1-713-790-1693	May 2019	Class II
Hydrocodone bitartrate and homatropine methylbromide oral solution 59741026216, 13668057710	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Risperidone 23155031751	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/Drugs/DrugSafetyandAvailability/DrugRecalls) for additional information.

Cetirizine hydrochloride 23155029251, 23155029252, 13668059607, 13668059611	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912- 9561	May 2019	Class II
Phenylephrine hydrochloride 00536138912, 00536138935, 00904768822	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912- 9561	May 2019	Class II
Bisacodyl 00536135501, 00536135512	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912- 9561	May 2019	Class II
Hydrocortisone acetate suppositories 59741030101, 59741030124, 59741030112	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912- 9561	May 2019	Class II
Memantine hydrochloride 39328055112 13668057309	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912- 9561	May 2019	Class II
Lactulose solution	Contamination with burkholderia	If you have questions about this recall, Torrent	May 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see *FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls* for additional information.

13668058008, 13668058012, 13668057408, 13668057412, 13668057410		Pharma Inc 1-800-912-9561		
Pseudoephedrine Hydrochloride 00536185085, 00536185097	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Diphenhydramine hydrochloride 00536077085, 00536077097	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Guaifenesin & dextromethorphan hydrobromide 00536097085, 00536097097	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Guaifenesin 00536082585, 00536082597	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Acetaminophen 00536012285, 00536012297	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Phenobarbital 16571033016	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Guaifenesin, Codeine Phosphate & Pseudoephedrine Hydrochloride 16571030116	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Guaifenesin AC 16571030216	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Acetic Acid 64980042415	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Hyoscyamine sulfate 39328004715, 39328004816	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Cyproheptadine hydrochloride 39328004416	Contamination with burkholderia	If you have questions about this recall, Torrent	May 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

		Pharma Inc 1-800-912-9561		
Bisacodyl 00904505812, 00904505860	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Pedia Relief 00904505020	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Robafen dm cough sugar free clear 00904630620	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Oxymetazoline Hydrochloride 00904571130, 00904571135	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Pseudoephedrine hydrochloride 00536185085, 00536185097	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Banofen 904517416	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Robafen DM 00904005300, 00904005316 00904005309	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Robafen AC 00904647916	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Risperidone 13668058906	Contamination with burkholderia	If you have questions about this recall, Torrent Pharma Inc 1-800-912-9561	May 2019	Class II
Pecgen dmx 52083063016	Incorrect labeling instructions	If you have questions about this recall, Novis PR 1-800-727-6711	May 2019	Class I
Cefdinir for oral suspension 68180072310, 68180072320	Metal piece identified in the product bottle	If you have questions about this recall, Lupin	May 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

		Pharmaceuticals Inc 1-410-576-2000		
Articaine Hydrochloride and Epinephrine Bitartrate® 66312060116	Mislabeling	If you have questions about this recall, Septodont Inc 1-717-286-0100	May 2019	Class II
Losartan Potassium tablets 23155064503, 23155064509, 23155064510, 23155064603, 23155064610, 23155064609, 23155064403, 23155064410, 23155064409	Presence of an impurity	If you have questions about this recall, Heritage Pharmaceuticals Inc 1-866-901-DRUG (3784)	May 2019	Class II
Amitiza® 64764008060, 64764024010, 64764024060	24 mcg capsules were found in a 8 mcg	If you have questions about this recall, Mallinckrodt Pharmaceuticals 1-314-654-2000	May 2019	Class II
Losartan Potassium tablets 60429031610, 60429031690, 60429031630, 60429031810, 60429031890, 60429031830	Presence of an impurity	If you have questions about this recall, Golden State Medical Supply Inc 1-800-284-8633 ext. 215	May 2019	Class II
Divalproex sodium 00904636345, 00904636361	Exposed above 50% relative humidity levels	If you have questions about this recall, Major	April 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

		Pharmaceuticals 1-800-875-0123 option 5		
Losartan Potassium tab 68788004809	Presence of an impurity	If you have questions about this recall, Preferred Pharmaceuticals Inc 1-714-777-3729	April 2019	Class II
Carvedilol tablets 65841061605	Label Mix-up	If you have questions about this recall, Zydus Pharmaceuticals 1-877-993-8779	April 2019	Class II
Ketorolac tromethamine 25021070101, 25021070102	Lack of sterility	If you have questions about this recall, Sagent Pharmaceuticals 1-866-625-1618	April 2019	Class II
Acyclovir tablets 68382079101, 68382079106, 68382079116, 68382079105, 68382079110, 68382079130	Label mix-up	If you have questions about this recall, Zydus Pharmaceuticals 1-877-993-8779	April 2019	Class II
Fentanyl transdermal system, 12 mcg/h 47781042347	Label mix-up	If you have questions about this recall, Alvogen 1-866-770-3024	April 2019	Class I

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Losartan Potassium/hctz 70518156000	Presence of an impurity	If you have questions about this recall, Remedy Repack 1-866-845-3791	April 2019	Class II
Losartan Potassium and hydrochlorothiazide tablets 13668011610, 13668011674, 13668011630, 13668011690, 13668011710, 13668011774, 13668011730, 13668011790, 13668011810, 13668011874, 13668011830, 13668011890	Presence of an impurity	If you have questions about this recall, Torrent Pharmaceuticals 1-800-912-9561	April 2019	Class II
Losartan Potassium 68645049454	Presence of an impurity	If you have questions about this recall, Legacy Pharmaceutical Packaging LLC 1-314-813-1555	April 2019	Class II
Losartan Potassium 50268051715, 50268051615	Presence of an impurity	If you have questions about this recall, Avkare Inc 1-931-292-6222	April 2019	Class !!
Losartan Potassium 13668011310, 13668011390, 13668011374, 13668040990, 13668040930, 13668040910, 13668040974, 13668011510, 13668011574, 13668011530, 13668011590	Presence of an impurity	If you have questions about this recall, Torrent Pharmaceuticals 1-800-912-9561	April 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Gavilyte- N 43386-0050-19	Labeling incorrectly classified	If you have questions about this recall, Lupin Pharmaceuticals 1-800-399-2561	March 2019	Class II
Combigan 00023921103, 00023921110, 00023921105, 00023921115	Trace amounts of an impurity	If you have questions about this recall, Ecolab 1-800-433-8871	March 2019	Class II
Pravastatin Sodium 68462-0196-05, 68462-0196-90	Foreign tablet in bottle	If you have questions about this recall, Glenmark Pharmaceuticals, Inc. 201-684-8000	March 2019	Class II
Losartan 70518-0588-01	Trace amounts of impurity	If you have questions about this recall, Remedy repack Inc 1-886-845-3791	March 2019	Class II
Phenobarbital 70166-0536-02	The label contains the incorrect expiration date	If you have questions about this recall, Lohxa LLC 1-800-641-5564	March 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see [FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls](https://www.fda.gov/drugs/drug-safety-and-availability/drug-recalls) for additional information.

Losartan Potassium 31722-0702-30	Impurity found	If you have questions about this recall, Pharma Pac 1-805-929-1333	March 2019	Class II
Volumex 50914-7720-08	Lack of assurance	If you have questions about this recall, Iso-Tex Diagnostics 1-800-477-4539	March 2019	Class II
Hydrocortisone and Acetic Acid 50383-0301-10	Sub Potent Drug	If you have questions about this recall, call Akon Inc. 1-800-477-4539	March 2019	Class II
Losartan Potassium 68645-0578-54	Trace amounts of impurity	If you have questions about this recall, Call Legacy 1-877-538-8443	February 2019	Class II
Losartan Potassium/Hydrochlorothiazide 13668-0118-10, 13668-0118-74, 13668-0118-30, 13668-0118-90, 13668-0117-10, 13668-0117-74, 13668-0117-30, 13668-0117-90, 13668-0116-10, 13668-0116-74, 13668-0116-30, 13668-0116-90	Presence of Impurity	If you have questions about this recall, Call Torrent Pharmaceuticals 1-800-912-9561	February 2019	Class II
Losartan Potassium 13668-0113-10, 13668-0113-90, 13668-0113-74, 13668-0409-10,	Presence of an impurity	If you have questions about this recall, Call	February 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

13668-0409-74, 13668-0409-30, 13668-0409-90, 13668-0115-10, 13668-0115-74, 13668-0115-30, 13668-0115-90		Torrent Pharmaceuticals 1-800-912-9561		
Valsartan 52343-0122-30, 52343-0123-90, 52343-0124-90, 52343-0125-90	Presence of an impurity	If you have questions about this recall, Call Preferred Pharmaceuticals 1-855-544-9419	February 2019	Class II
Losartan Potassium 68788-6882-03, 68788-6882-09	Presence of an impurity	If you have questions about this recall, Call Preferred Pharmaceuticals 1-855-544-9419	February 2019	Class II
Valsartan 60687-0139-01	Trace amounts of an impurity	If you have questions about this recall, call Aurobindo 1-800-912- 9572	February 2019	Class II
Losartan Potassium 31722-0702-05, 31722-0702-30, 31722-0702-90, 31722-0702-10, 31722-0702-60, 50268-0517-15, 50268-0513-15, 50268-0514-15, 68645-0577-54	Trace amounts of an impurity	If you have questions about this recall, call Aurobindo 1-800-912- 9572	February 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

Alprazolam 51079-0788-20	Failed impurities/degradation	If you have questions about this recall, call Mylan 1-888-406-9305	January 2019	Class II
Ceftriaxone for Injection 68180-0611-01	Presence of Particulate Matter: Product complaints received of grey flecks, identified as shredded rubber particulate matter from the stopper observed in reconstituted vials	If you have questions about this recall, Lupin Pharmaceuticals Inc. 111 S Calvert St Fl 21ST Baltimore, MD 21202-6174	January 2019	Class I
Vecuronium Bromide for Injection 10 mg 47335-0931-44	Presence of Particulate Matter: Foreign matter identified as glass detected in Vecuronium Bromide for Injection.	If you have questions about this recall, Sun Pharmaceutical Industries, Inc. 270 Prospect Plains Rd Cranbury, NJ 08512-3605	January 2019	Class I
OZURDEX 0023-3348-07	Deviations: A silicone particulate was noted in Ozurdex	If you have questions about this recall, Allergan, PLC. 5 Giralda Farms Madison, NJ 07940-1027 (714) 246-4500	January 2019	Class II
Estradiol Vaginal Inserts 68462-711-71 68462-711-88	Defective Delivery System: Customer complaints of malfunctioning plunger of the applicator	If you have questions about this recall, Glenmark Pharmaceuticals Inc., USA	January 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

		750 Corporate Dr Mahwah, NJ 07430-2009		
Dianeal Low Calcium 00941-0424-52 Lot #: Y281477 Expiration: 02/2020	Lack of Assurance of Sterility: Confirmed customer complaints for leaks on the tubing	If you have questions about this recall, Baxter Healthcare Corporation 1 Baxter Pkwy Deerfield, IL 60015-4625	January 2019	Class II
Cefdinir for Oral Suspension USP 125mg/5mL 68180-0722-10 68180-0722-20	Deviations: Product complaints received indicating reconstituted suspension was observed to be thick	If you have questions about this recall, Lupin Pharmaceuticals Inc. 111 S Calvert St Fl 21ST Baltimore, MD 21202- 6174 866-587-4617	January 2019	Class II
Cidofovir Injection 375mg 23255-0216-31	Lack of Assurance of Sterility: complaints received about dried powder on the outside of bottle	If you have questions about this recall, Heritage Pharmaceuticals, Inc. 1 Tower Center Blvd Ste 1700 East Brunswick, NJ 08816-1145 732) 429-1000	January 2019	Class II
Cephalexin for Oral Suspension USP 68180-0124-01	Deviation; manufacturing batch record could not be located	If you have questions about this recall, Lupin Pharmaceuticals Inc.	January 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see *FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls* for additional information.

68180-0124-02		111 S Calvert St Fl 21ST Baltimore, MD 21202 866-587-4617		
Valsartan tablets USP 320 mg 65862-0573-05 65862-0573-90	Deviations: FDA lab confirmed presence an impurity, N-nitrosodimethylamine (NDEA) contained in the API used to manufacture the product above the interim acceptable daily intake level of 0.083 parts per million.	If you have questions about this recall, Aurobindo Pharma USA Inc. 279 Princeton Hightstown Rd East Windsor, NJ 08520-1401 732-839-9400	January 2019	Class II
Valsartan and Hydrochlorothiazide tablets 65862-0549-10 65862-0549-90 65862-0549-99	Deviations: FDA lab confirmed presence an impurity, N-nitrosodimethylamine (NDEA) contained in the API used to manufacture the product above the interim acceptable daily intake level of 0.083 parts per million	If you have questions about this recall, Aurobindo Pharma USA Inc. 279 Princeton Hightstown Rd East Windsor, NJ 08520 732- 839-9400	January 2019	Class II
Amlodipine and Valsartan Tablets USP 10 mg/320 mg	Deviations: FDA lab confirmed presence an impurity, N-	If you have questions about this recall,	January 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

65862-0740-03 65862-0740-30 65862-0740-90	nitrosodimethylamine (NDEA) contained in the API used to manufacture the product above the interim acceptable daily intake level of 0.083 parts per million.	Aurobindo Pharma USA Inc. 279 Princeton Hightstown Rd East Windsor, NJ 08520-1401 <u>732 839 9400</u>		
EEMT (esterified estrogens and methyltestosterone) 15310-0010-01	The combination of esterified estrogens and methyltestosterone is used to treat symptoms of menopause such as hot flashes, and vaginal dryness, burning, and irritation.	If you have questions about this recall, Syntho Pharmaceuticals, Inc. 230 Sherwood Ave Farmingdale, NY 11735-1718 631-755-9898	January 2019	Class II
Olmesartan Medoxomil and Hydrochlorothiazide Tablets, 40 mg/25 mg 00093761756 00093761798	Failed dissolution specifications	If you have questions about this recall, Teva Pharmaceuticals USA 1090 Horsham Rd North Wales, PA 19454-1505 888-TEVA-USA (888-838-2872)	January 2019	Class II

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see *FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls* for additional information.

<p>Fluocinolone Acetonide Topical Solution, USP, 0.01 %, 60 mL bottle</p> <p>Lot #: S700214, Exp Apr-19; S700447, Exp Jun-19; S700787, Exp Oct-19; S701057, Exp Nov-19; S800107, Exp Feb-20; S800266, Exp Mar-20; S800524, Exp May-20; S800791 Kul Exp Jul-20</p>	<p>Failed Impurities/Degradation Specifications: Expansion of October 2018 recall due to elevated out of specification results for total impurities that have been chemically identified as oxidative degradation products of the fluocinolone active pharmaceutical ingredient</p>	<p>If you have questions about this recall, Contact: LUPIN SOMERSET 400 Campus Dr Somerset, NJ 08873-1145</p>	<p>January 2019</p>	<p>Class II</p>
<p>Ezetimibe and Simvastatin Tablets 10mg/80 mg, 1000-count bottles</p> <p>Lot #: 43E021 and 43E023, Exp. 01/2020</p>	<p>Presence of Foreign Substance: Product complaint of black speckles observed on tablets.</p>	<p>If you have questions about this recall, Contact: Dr. Reddy's Laboratories, Inc. 107 College Rd E Princeton, NJ 08540-6623</p>	<p>January 2019</p>	<p>Class II</p>
<p>Losartan potassium and hydrochlorothiazide 1000mg/25mg tablets</p> <p>ONLY Lot: JB8912</p>	<p>This product is being recalled due to the presence of an impurity, N-nitrosodiethylamine. NDEA occurs naturally in manufacturing processes and could cause cancer based on laboratory test results. No adverse effects related to this recall have been reported to date.</p>	<p>If you have questions about this recall, Contact Sandoz Inc. at 1-800-525-8747 Monday-Friday 8:30 AM – 5:00 PM (EST) or email usdrugsafety.operations@novartis.com.</p>	<p>November 2018</p>	<p>Class I</p>

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see *FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls* for additional information.

Montelukast Sodium 10Mg	This medicine is made by Camber Pharmaceuticals, Inc. This medicine is being recalled because sealed bottles labeled as Montelukast sodium tablets, 10 mg, 30-count bottle from Camber were found to instead contain 90 tablets of losartan potassium tablets, 50 mg. This could result in a life-threatening condition.	If you have questions about this recall, call Camber Pharmaceuticals Med Line at 1-866-495-1995	September 2018	Class I
Diphenoxylate Hydrochloride and Atropine Sulfate tablets, 2.5 mg/0.025 mg (Lomotil).	This medication is made by Greenstone LLC, a wholly owned subsidiary of Pfizer, Inc. This medication is being recalled due to super potency. The use of the impacted super potent product when used as labeled has a low probability of being associated with adverse events of limited severity such as lethargy, skin flush, and drowsiness. Serious adverse events such as coma and respiratory depression are improbable.	If you have questions about this recall, please contact Pfizer Customer Support at 800-879-3477.	January 2018	Class I

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.

EpiPen and EpiPen Jr Auto-Injector	This medication is made by Mylan. This medication is being recalled due to failure to activate the device or increased force needed to activate due to a potential defect in a supplier component. The potential defect could make the device difficult to activate in an emergency. This could result in a life-threatening condition.	If you have questions about this recall, please contact your physician, pharmacy or Mylan Customer Relations at 800-796-9526 or customer.service@mylan.com .	May 2017	Class I

* Drug Recall Class

Class 1 Recall: Reasonable probability that using the drug will cause serious adverse health consequences or death.

Class 2 Recall: Using the drug may cause temporary or medical reversible adverse health consequences, the probability of serious adverse health consequences is remote.

Class 3 Recall: Using the drug is not likely to cause adverse health consequences.

NOTE: This is not a complete list of all recalls. Please see FDA.gov >Drugs>Drug Safety and Availability> Drug Recalls for additional information.