

Enforcement Report - Week of March 11, 2026

Class I Drugs Event

Event ID:

98258

Status:

Ongoing

Recall Initiation Date:

10/20/2025

Center Classification Date:

03/13/2026

Recalling Firm:

ANTHONY TRINH, 123Herbals LLC
7214 Berne St
Rosemead, CA 91770-3844
United States

Distribution Pattern:

USA nationwide

Product Type:

Drugs

Date Terminated:

N/A

Voluntary / Mandated:

Voluntary: Firm initiated

Initial Firm Notification of Consignee or Public:

Press Release

Associated Products

Product Description:

SILINTAN 25/pills, packaged in a 25-count bottle, Shanghai Chinese Medical Works, Shanghai, China

Product Quantity:

N/A

Reason for Recall:

Marketed Without an Approved NDA/ANDA: FDA analysis revealed the presence of undeclared meloxicam.

Recall Number:

D-0388-2026

Code Information:

All lots within expiry

Class II Drugs Event

Event ID:

98291

Status:

Ongoing

Recall Initiation Date:

01/20/2026

Center Classification Date:

03/02/2026

Recalling Firm:

ACME UNITED CORPORATION
1 Waterview Dr Ste 200
Shelton, CT 06484-4368
United States

Distribution Pattern:

Nationwide within the United States

Product Type:

Drugs

Date Terminated:

N/A

Voluntary / Mandated:

Voluntary: Firm initiated

Initial Firm Notification of Consignee or Public:

Letter

Associated Products

Product Description:

First Aid Only BZK Antiseptic Towelettes (Benzalkonium Chloride 0.13%) 100 Wipes, NDC 0924-7116-04

Product Quantity:

212,607,900 containers

Reason for Recall:

CGMP Deviations

Recall Number:

D-0358-2026

Code Information:

Lot #: MN26222, MN34722, MN25922, Exp. Date Nov 2027; MN00523, MN00523, Exp. Date Dec 2027.

Product Description:

First Aid Only BZK Antiseptic Towelettes (Benzalkonium Chloride 0.13%), 50 Wipes, NDC 0924-7116-03.

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0359-2026

Code Information:

Lot #: MN34722, MN34722, Exp. Date Nov 2027; MN34722B, Exp. Date Aug 2027.

Product Description:

Dukal BZK Antiseptic Towelette (Benzalkonium Chloride 0.13%) 1000 Packs per case, Manufactured For: Dukal, LLC, Ronkonkoma, NY 11779, Made in USA, NDC 65517-0004-1

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0360-2026

Code Information:

Lot #: MN31621, Exp. Date Oct 2026; MN14522, MN15522, Exp. Date Apr 2027; MN28122, MN31522, Exp. Date Sept 2027; MN04223, MN06023, Exp. Date Feb 2028; MN17223, Exp. Date May 2028; MN25123, Exp Date Aug 2028; MN33123, MN33523, Exp. Date Nov 2028.

Product Description:

First Aid Only BZK Antiseptic Towelette (Benzalkonium Chloride 0.13%), 5x7, bulk, Acme United Corporation, 2280 Tanner Road, Rocky Mount, NC, 27801 Made in USA, NDC 0924-7116-00.

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0361-2026

Code Information:

Lot #: MN12121, MN13521, Exp. Date Apr 2026; MN15221, MN16721, MN16921, MN17021, MN17021B, Exp. Date May 2026; MN18521, MN18621, MN18621B, Exp. Date Jun 2026; MN24721, MN25021, Exp. Date Aug 2026; MN01123, Exp. Date Dec 2027; MN27421, MN27721, Exp. Date Sep 2026; MN31321, MN31421, Exp. Date Oct 2026; MN34321, MN34521, MN34721, Exp. Date Nov 2026; MN02022, MN03022, Exp. Date Dec 2026; MN03522, MN04622, Exp. Date Jan 2027; MN06022, MN06122, Exp. Date Feb 2027; MN12322, MN12422, MN14422, Exp. Date Apr 2027; MN15322, Exp. May 2027; MN22022, Exp. Date Jul 2027; MN25222, MN25322, MN26422, Exp. Date Aug 2027; MN27822, Exp. Date Sep 2027; MN15322, Exp. Date Nov 2027; MN06123, Exp. Date Aug 2027; MN06423, MN06523, Exp. Date Feb 2028; MN09423, MN09523, MN09723, MN10223 Exp Date Mar 2028; MN12623, MN13323, Exp. Date Apr 2028; MN15322, MN16523, Exp. Date May 2028; MN18623, MN19323, MN19423, MN20223, Exp. Date Jun 2028; MN24323, MN24423 Exp. Date Jul 2028; MN34523, Exp. Date Nov 2028; 301424, 301524, 301624, 301724, 303124, Exp. Date Dec 2028.

Product Description:

1) BZK Antiseptic Towelettes (Benzalkonium Chloride 0.13%) 5"x7" Piedmont Sanitizing Hand Wipes, 1,000 ea Bulk, Made in USA for: Food Service Resources, Flowery Branch, GA 30542, NDC 0924-7130-00. 2) BZK Antiseptic Towelettes (Benzalkonium Chloride 0.13%) 5" x 7" Touchpoint, 1000 ea Bulk, Made in USA for : Food Service Resources, Flowery Branch, GA 30542. NDC 0924-7130-00. 3) BZK Antiseptic Towelettes (Benzalkonium Chloride 0.13%) 5" x 7" Wellstar Cobb Hand Wipes, 1000 ea Bulk, Made in the USA for Food Service Resources, Flowery Branch, GA 30542, NDC 0924-7130-00. 4) BZK Antiseptic Towelettes (Benzalkonium Chloride 0.13%) 5" x 7" FSR Sanitizing Hand Wipes, 1000 ea Bulk, Made in USA for Food Service Resources, Flowery Branch, GA 30542, NDC 0924-7130-00. Inner Packaging - Instant Hand Sanitizing Wipes 5) BZK Antiseptic Towelettes (Benzalkonium Chloride 0.13%) 5" x7" Children's 1000 ea Bulk, Made in USA for: Food Service Resources, Flowery Branch, GA 30542 NDC 0924-713-00. Inner packaging -Children's Healthcare of Atlanta

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0362-2026

Code Information:

1) Lot #: MN22222, Exp. Date Jul 2027; MN05523, Exp. Date Jan 2028; MN32023, Exp. Date Oct 2028; MN15523, Exp. Date May 2028. 2) Lots MN11422B, MN11422, Exp. Date Mar 2027; MN03823, Exp. Date Jan 2028; MN09223 Exp. Date Mar 2028; MN32123, Exp. Date Oct 2028; MN20323, Exp. Date Jun 2028; MN30822 Exp. Date Oct 2027; MN304024, Exp. Date Dec 2028; MN13023, Exp. Date Apr 2028; MN12222 Exp. Date Apr 2027-Apr; MN16023, Exp. Date May 2028; MN27423, Exp. Date Aug 2028. 3) Lot #: MN304024, Exp. Date Dec 2028; MN13023, Exp. Date Apr 2028; MN12222, Exp. Date Apr 2027; MN16023, Exp. Date May 2028; MN27423, Exp. Date Aug2028. 4) Lot#: MN04423, Exp. Date Jan 2028; MN10022, Exp. Date Jul 2027; MN30722, Exp. Date Oct 2027; MN03523, Exp. Date Jan 2028; MN09323, Exp. Date Mar 2028; MN20223, Exp. Date July 2028; MN32223, Exp. Date Oct 2028 5) Lots MN12122 2027-Apr, MN27523 2028-Aug & MN30922 2027-Oct.

Product Description:

Max Packaging Antibacterial Towelette (Benzalkonium Chloride 0.13%), 1 towelette, 4x7, Max Packaging, Made in the USA, NDC 59647-750-01

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0363-2026

Code Information:

Lot #: MN24523, Exp. Date July 2028; MN27123, Exp. Date Aug 2028; MN28223, Exp. Date Sep 2028.

Product Description:

Savannah Antibacterial Towelettes (Benzalkonium Chloride 0.13%), 1 towelette, 4 x 7, R&R Enterprise, NDC 59647-749-02

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0364-2026

Code Information:

Lot #: MN24623, Exp. Date Jul 2028; MN27023, Exp. Date Aug 2028; MN32423, Exp. Oct 2028; MN300724, Exp. Date Dec 2028.

Product Description:

Medi-First Antiseptic Wipes (BZK)(Benzalkonium Chloride 0.13%), 100 wipes per box, Manufactured for Medique Products, Fort Meyers, FL, 33967, NDC 47682-122-33.

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0365-2026

Code Information:

Lot #: MN15922, Exp. Date Apr 2027; MN31022, Exp. Date Oct 2027; MN27223, Exp. Date 2028-Aug.

Product Description:

Med Nap BZK Antiseptic Towelettes (Benzalkonium Chloride 0.13%), 100 Towelettes, Manufactured by: Acme United Corporation, 1 Waterview Dr, Shelton, CT 06481, NDC 0924-0246-01.

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0366-2026

Code Information:

Lot #: MN09322, Exp. Date Mar 2027; MN15222, Exp. Date May 2027; MN21622, Exp. Date Jul 2027; MN31822, Exp. Date Oct 2027; MN06223, MN06323, Exp. Date Feb 2028; MN12523, Exp. Date Apr 2028; MN16123, Exp. Date May 2028; MN20523, MN20623, MN20723, Exp. Date Jun 2028; MN23223, Exp. Date Jul 2028; MN31123, Exp. Date Oct 2028; MN300524, MN300624, MN302624, MN302724, MN303324, Exp. Date Dec 2028.

Product Description:

BZK Antiseptic Towelettes (Benzalkonium Chloride 0.13%), 5" x7", 1000 ea Bulk, Med-Nap, Brooksville, FL 34601, Made in USA, NDC 0924-0246-02.

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0367-2026

Code Information:

Lot # MN23323, Exp. Date Jul 2028; MN31323, Exp. Date Oct 2028; MN21622, Exp. Date Jul 2027; MN13123, Exp. Date Apr 2028.

Product Description:

Wipes Plus, Hand Sanitizing Alcohol Free BZK Wipes (Benzalkonium Chloride 0.13%), Manufactured for: CFS Brands DBA WipesPlus, 4711 E. Heiner Rd. Oklahoma City, OK 73331, NDC 67151-333-01.

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0368-2026

Code Information:

Lot #: MN16022, Exp. Date Aug 2027; MN33922, MN31222, Exp. Date Apr 2027; MN16522, Exp. Date Oct 2027; MN17022, Exp. Date Apr 2027; MN03323, Exp. Date Jan 2028; MN34622, Exp. Date Oct 2027.

Product Description:

Dynarex Obstetrical Towelettes Cleanser (Benzalkonium Chloride 0.13%), 100 packets, 5" x 7", Manufactured for Dynarex Corporation, 10 Glenshaw Street, Orangeburg, NY 10962, USA Made in USA, NDC 67777-244-02

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0369-2026

Code Information:

Lot #: 52347-R, 52348-R, 52349-R, 52350-R, 52351-R, 52352-R, 53518, 53519, Exp. Date Mar 2027.

Product Description:

Cleansing Towelette (Benzalkonium Chloride 0.13%), 5"x 7", 1,000 case/Bulk, Med-Nap, Brooksville, FL 34601, Made in USA, NDC 0924-0243-00.

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0370-2026

Code Information:

Lot #: MN15923, Exp. Date May 2028; MN25822, Exp. Date Aug 2027.

Product Description:

Med-Nap Cleansing Towelettes (Benzalkonium Chloride 0.13%), 100 Towelettes, Manufactured by: Acme United Corporation, 1 Waterview Dr., Stelton, CT 06484, NDC 0924-0243-01.

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0371-2026

Code Information:

Lot #: 52347, 52348, 52349, 52350, 52351, Exp. Date Mar 2027.

Product Description:

WP WipesPlus Hand Sanitizing Wipes (Benzalkonium Chloride 0.13%), 5x7, 100 CT, Progressive Products, 211 S. Ridge Street, Rye Brook, NY 10573. NDC 67151-727-01.

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0372-2026

Code Information:

Lot #: MN27922, MN21722, MN21822, Exp. Date Jul 2027; MN27422, MN27522, Exp. Date Sep 2027; MN25422, Exp. Date Aug 2027; MN27922, MN27922, Exp. Date Sep 2027.

Product Description:

Hand Sanitizing Wipes (Benzalkonium Chloride 0.13%), 1000 per case, Acme United Corporation, 2280 Tanner Road, Rocky Mount, NC 27801, NDC 0924-0301-00

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0373-2026

Code Information:

Lot #: MN24922, Exp. Date Aug 2027; MN31122, Exp. Date Oct 2027.

Product Description:

Green Guard Antiseptic Wipes (Benzalkonium Chloride 0.13%), 25 wipes per box, Distributed by Green Guard, St. Louis, MO 63045, NDC 47682-056-73

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0374-2026

Code Information:

Lot #: MN15922, Exp. Date Apr 2027; MN18223, Exp. Date May 2028; MN10123, Exp. Date Mar 2028; MN27223, Exp. Date Aug 2028; MN01023, Exp. Date Dec 2027.

Product Description:

Dynarex, BZK Antiseptic Towelettes (Benzalkonium Chloride 0.13%), 100 packets, 5x7, Manufactured for Dynarex Corporation, 10 Glenshaw Street, Orangeburg, NY 10962,USA, NDC 67777-245-01

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0375-2026

Code Information:

Lot #: 52347-R, 52348-R, 52349-R, 52350, 52351-R, 52352, Exp. Date Mar 2027; 53518, Exp. Date Jun 2027.

Product Description:

Dynarex, BZK Antiseptic Towelettes (Benzalkonium Chloride 0.13%), 5x7, 1000 packets, 5x7, Manufactured for Dynarex Corporation, 10 Glenshaw Street, Orangeburg, NY 10962,USA, NDC 67777-245-04.

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0376-2026

Code Information:

Lot #: 53518, 53519, Exp. Date Jun 2027.

Product Description:

Children's Healthcare of Atlanta, Instant Hand Sanitizing Wipe(Benzalkonium Chloride 0.13%), 1000 per case, Made in USA for Food Service Resources, Flowery Branch, GA 30542, NDC 0924-7130-00

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0377-2026

Code Information:

Lot #: MN12122, Exp. Date Apr 2027; MN27523, Exp. Date Aug 2028; MN30922, Exp. Date Oct 2027.

Product Description:

Antimicrobial Alcohol Hand Wipe, Isopropyl Alcohol 70%, Manufactured for : Custom Packaging Co., Inc, Louisville, KY 4023, NDC 0924-251-00.

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations

Recall Number:

D-0378-2026

Code Information:

Lot #: MN15423 , MN16422, Exp. Date Apr 2027-Apr; MN26322, MN26322, Exp. Date Aug 2027; MN03423, MN03423, Exp. Date Jan 2028; MN09722, Exp. Date Mar 2027; MN31722, Exp. Date Oct 2027; MN15722, MN13722 Exp. Date Apr 2027; MN31722, Exp. Date Oct 2027; MN09722, Exp. Date Mar 2027.

Class II Drugs Event**Event ID:**

98348

Product Type:

Drugs

Status:

Ongoing

Date Terminated:

N/A

Recall Initiation Date:

01/29/2026

Voluntary / Mandated:

Voluntary: Firm initiated

Center Classification Date:

02/27/2026

Initial Firm Notification of Consignee or Public:

Letter

Recalling Firm:

Teva Pharmaceuticals USA, Inc
400 Interpace Pkwy Bldg A
Parsippany, NJ 07054-1120
United States

Distribution Pattern:

Nationwide

Associated Products**Product Description:**

Metoprolol Succinate Extended-Release Tablets, 25 mg, 100 tablets bottles (NDC: 45963-709-11) and 1,000 tablets bottles (NDC: 45963-709-96), Rx Only, Teva Pharmaceuticals, Parsippany, NJ 07054

Product Quantity:

18,780 bottles

Reason for Recall:

Failed Dissolution Specifications

Recall Number:

D-0354-2026

Code Information:

Lot#: a) 0715J251, Exp.: 12/2026; 0716J251, Exp.: 01/2027; 0717J251, Exp.: 01/2027; b) 0715J252, Exp.: 12/2026; 0716J252, Exp.: 01/2027, 0717J252, Exp.: 01/2027.

Product Description:

Metoprolol Succinate Extended-Release Tablets, 50 mg, 100 tablets bottles (NDC: 45963-676-11) and 1,000 tablets bottles (NDC: 45963-676-96), Rx Only, Teva Pharmaceuticals, Parsippany, NJ 07054

Product Quantity:

67,043 bottles

Reason for Recall:

Failed Dissolution Specifications

Recall Number:

D-0355-2026

Code Information:

Lot#: a) 0486G251, 0487G251, 0488G251, Exp.: 01/2027; b) 0486G252, 0487G253, 0488G252, Exp.: 01/2027.

Product Description:

Metoprolol Succinate Extended-Release Tablets, Rx Only, 100 mg, 100 tablets bottles (NDC: 45963-677-11) and 1,000 tablets bottles (NDC: 45963-677-96), Teva Pharmaceuticals, Parsippany, NJ 07054.

Product Quantity:

7,448 bottles

Reason for Recall:

Failed Dissolution Specifications

Recall Number:

D-0356-2026

Code Information:

Lot#: a) 0718J251, 0729J251, 0730J251, Exp.: 12/2026; b) 0718J252, 0729J252, 0730J252, Exp.: 12/2026.

Product Description:

Metoprolol Succinate Extended-Release Tablets, 200 mg, 100 tablets bottles, Rx Only, Teva Pharmaceuticals, Parsippany, NJ 07054, NDC: 45963-678-11

Product Quantity:

19,056 bottles

Reason for Recall:

Failed Dissolution Specifications

Recall Number:

D-0357-2026

Code Information:

Lot#: a) 0686H251, 0687H251, 10688H251, Exp.: 02/2027. .

Class II Drugs Event

Event ID:

98438

Product Type:

Drugs

Status:

Ongoing

Date Terminated:

N/A

Recall Initiation Date:

02/13/2026

Voluntary / Mandated:

Voluntary: Firm initiated

Center Classification Date:

03/02/2026

Initial Firm Notification of Consignee or Public:

E-Mail

Recalling Firm:

Harbin Jixianglong Biotech Co., Ltd.
 North Of Baoan Road, East Of Changqing Street Hulan District
 Haerbin
 China

Distribution Pattern:

Nationwide within the United States

Associated Products

Product Description:

Semaglutide, For Rx compounding use only, packaged in a) 1g, NDC 84385-106-01; b) 5g, NDC 84385-106-02; c) 10g, NDC 84385-106-06; d) 25g, NDC 84385-106-03; d) 50g, NDC 84385-106-04 Manufacturer: Harbin Jixianglong Biotech Co., Ltd., North of Baoan Road, East of Changqing Street, Limin Development Zone, Harbin, China

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations This recall has been initiated due to failing to complete process validation and bacterial endotoxin method validation before distribution

Recall Number:

D-0379-2026

Code Information:

Lot #: CP-030-20250711, Exp. Date Jul. 24th, 2027

Product Description:

Semaglutide, For Rx compounding use only, packaged in a) 1g, NDC 84385-106-01; b) 5g, NDC 84385-106-02; c) 10g, NDC 84385-106-06; d) 25g, NDC 84385-106-03; d) 50g, NDC 84385-106-04; e) 100g, NDC 84385-106-05 Manufacturer: Harbin Jixianglong Biotech Co., Ltd., North of Baoan Road, East of Changqing Street, Limin Development Zone, Harbin, China.

Product Quantity:

N/A

Reason for Recall:

CGMP Deviations This recall has been initiated due to failing to complete process validation and bacterial endotoxin method validation before distribution

Recall Number:

D-0380-2026

Code Information:

Batch # CP-030-20250911, Mfg Date: Sep. 25th, 2025, Retest Date: Sep. 24th, 2027.

Class II Drugs Event

Event ID:

98456

Product Type:

Drugs

Status:

Ongoing

Date Terminated:

N/A

Recall Initiation Date:

02/17/2026

Voluntary / Mandated:

Voluntary: Firm initiated

Center Classification Date:

03/03/2026

Initial Firm Notification of Consignee or Public:

Letter

Recalling Firm:

The Harvard Drug Group LLC
7000 Cardinal Pl
Dublin, OH 43017-1091
United States

Distribution Pattern:

Nationwide US.

Associated Products

Product Description:

Midodrine Hydrochloride Tablets, USP, 5 mg, 50 Tablets (5 x 10 blister packs), Rx only, Packaged and Distributed by: Major Pharmaceuticals, Indianapolis, IN 46268 USA, NDC 0904-6818-06.

Product Quantity:

8892 cartons

Reason for Recall:

Defective container; inadequately sealed blister packaging.

Recall Number:

D-0383-2026

Code Information:

Lot # N02640, Exp Date: 08/2027

Class II Drugs Event

Event ID:

98470

Product Type:

Drugs

Status:
Ongoing

Date Terminated:
N/A

Recall Initiation Date:
01/26/2026

Voluntary / Mandated:
Voluntary: Firm initiated

Center Classification Date:
03/13/2026

Initial Firm Notification of Consignee or Public:
Letter

Recalling Firm:
Radnostix
4137 Commerce Cir
Idaho Falls, ID 83401-1205
United States

Distribution Pattern:
Nationwide in the USA and Puerto Rico

Associated Products

Product Description:
Sodium Iodide (I-131) Solution, Therapeutic Oral, packaged as a) 1 ml V-vial, NDC 69208-003-15; Order 49317, 49323, 49325, 49329, 49333, 49339, 49341, 49355, 49364, 49380, 49392. (b) 2 ml V-Vial, NDC 69208-003-25; Order 49353, 49457, 49477. (c) 3 ml V-vial, NDC 69208-003-35; Order 49454, 49406. Rx Only, Manufactured by: International Isotopes Inc., Idaho Falls, ID.

Product Quantity:
16 units

Reason for Recall:
Presence of Particulate Matter: Due to production issues

Recall Number:
D-0389-2026

Code Information:
Batch I012626R-01, Exp 02/09/2026

Class II Drugs Event

Event ID:
98494

Product Type:
Drugs

Status:
Ongoing

Date Terminated:
N/A

Recall Initiation Date:
02/18/2026

Voluntary / Mandated:
Voluntary: Firm initiated

Center Classification Date:
03/05/2026

Initial Firm Notification of Consignee or Public:
E-Mail

Recalling Firm:
HTO Nevada Inc. dba Kirkman
6400 Rosewood St
Lake Oswego, OR 97035-5392
United States

Distribution Pattern:
Nationwide within the U.S

Associated Products

Product Description:
FIVE-STAR, VASOCAINE Spray, (Lidocaine HCl 4%, Racepinephrine HCl 0.01%), 4oz bottles, Distributed By: Dermal Source, Portland, OR 97232, NDC 80069-016.

Product Quantity:
27,380 bottles.

Reason for Recall:

Stability Data Does Not Support Expiry Date.

Recall Number:

D-0385-2026

Code Information:

Lot #: 839CP-0016, Exp. Date 03/31/2026; 839CP-0017, Exp. Date 06/30/2026; 839CP-0018, Exp. Date 08/31/2026; 839CP-0019, Exp. Date 10/31/2026; 839CP-0020, Exp. Date 03/31/2027; 839CP-0021, Exp. Date 07/31/2027; 839CP-0022, Exp. Date 11/30/2027; 839CP-0023, Exp. Date 02/29/2028; 839CP-0024, Exp. Date 07/31/2028.

Product Description:

MAXIMUM, ZONE 1, 4% Lidocaine Cream, 1/2 oz bottles, Distributed By: Dermal Source, Portland, OR 97232, NDC 80069-014.

Product Quantity:

178 bottles

Reason for Recall:

Stability Data Does Not Support Expiry Date.

Recall Number:

D-0386-2026

Code Information:

Lot #: 807CP-0003, Exp. Date 07/31/2026.

Product Description:

MAXIMUM, ZONE 2, 4% Lidocaine Cream, (Lidocaine 4% and Epinephrine 0.01%), 1 oz bottles, Distributed By: Dermal Source, Portland, OR 97232, NDC 80069-015.

Product Quantity:

243 bottles

Reason for Recall:

Stability Data Does Not Support Expiry Date.

Recall Number:

D-0387-2026

Code Information:

Lot #: 795CP-0005, Exp. Date 06/30/2026

Class II Drugs Event

Event ID:

98523

Status:

Ongoing

Recall Initiation Date:

03/03/2026

Center Classification Date:

03/04/2026

Recalling Firm:

Rising Pharma Holding, Inc.
2 Tower Center Blvd Ste 1401
East Brunswick, NJ 08816-1149
United States

Distribution Pattern:

Nationwide within the U.S

Product Type:

Drugs

Date Terminated:

N/A

Voluntary / Mandated:

Voluntary: Firm initiated

Initial Firm Notification of Consignee or Public:

Letter

Associated Products

Product Description:

Product label: Temozolomide Capsules, 5mg, packaged in 5-capsule bottles, Rx only, Manufactured for: Rising Pharmaceuticals, Inc., Allendale, NJ

07401, Country of origin: Taiwan, NDC 16571-816-51.

Product Quantity:

1200 bottles

Reason for Recall:

Failed Impurities/Degradation Specifications: An out-of-specification result observed during 9th-month long term stability testing

Recall Number:

D-0384-2026

Code Information:

Lot #: 1TM0524003A, Exp. Date 09/2026.

Class III Drugs Event

Event ID:

98452

Product Type:

Drugs

Status:

Ongoing

Date Terminated:

N/A

Recall Initiation Date:

02/18/2026

Voluntary / Mandated:

Voluntary: Firm initiated

Center Classification Date:

03/03/2026

Initial Firm Notification of Consignee or Public:

Letter

Recalling Firm:

Cipla USA, Inc.
10 Independence Blvd
Warren, NJ 07059-2730
United States

Distribution Pattern:

U.S.A. Nationwide

Associated Products

Product Description:

Nilotinib Capsules, 150 mg per capsule, packaged in cartons, Rx only, Manufactured by: Cipla Ltd., Verna, Goa, India, Manufactured for: Cipla USA, Inc., 10 Independence Boulevard, Suite 300, Warren, NJ 07059, Outer carton: 112 capsules (4 individual packs containing 28 capsules each) NDC 69097-031-74; Inner carton: 28 capsules (4 blisters of 7 capsules) NDC 69097-031-56; Foil blister: NDC 69097-031-17

Product Quantity:

271 cartons - Ex: 84 outer cartons x 4 inner cartons x 4 blisters x 7 capsules

Reason for Recall:

Failed Tablet/Capsule Specifications: Observed OOS results at 6-months long-term stability condition for Description test and Appearance by Visual Inspection test.

Recall Number:

D-0381-2026

Code Information:

Lot #: 5GJ0220, 5GJ0221, 5GJ0222, Exp 04/30/2027

Product Description:

Nilotinib Capsules, 200 mg per capsule, packaged in cartons, Rx only, Manufactured for: Cipla USA, Inc., 10 Independence Boulevard, Suite 300, Warren, NJ 07059, Outer carton: 112 capsules (4 individual packs containing 28 capsules each), NDC 69097-032-74; Inner carton: 28 capsules (4 blisters of 7 capsules), NDC 69097-032-56; Foil blister: NDC 69097-032-17

Product Quantity:

164 cartons

Reason for Recall:

Failed Tablet/Capsule Specifications: Observed OOS results at 6-months long-term stability condition for Description test and Appearance by Visual Inspection test.

Recall Number:

D-0382-2026

Code Information:

Lot #: 5GJ0223, Exp 04/30/2027

Not Yet Classified Drugs Event

Event ID:

97796

Status:

Ongoing

Recall Initiation Date:

10/08/2025

Center Classification Date:

N/A

Recalling Firm:

Mohamed Hagar
4952 Highway 169 N
Brooklyn Ctr, MN 55428-4026
United States

Distribution Pattern:

U.S. Nationwide

Product Type:

Drugs

Date Terminated:

N/A

Voluntary / Mandated:

Voluntary: Firm initiated

Initial Firm Notification of Consignee or Public:

E-Mail

Associated Products

Product Description:

Mojo Max Fusion XXX, 500 mg, one capsule per blister card, 20 count box, Distributed by: Max Nutrition Inc. UPC 6 78945 36675 0

Product Quantity:

125 boxes

Reason for Recall:

Marketed Without an Approved NDA/ANDA-FDA analysis revealed undeclared sildenafil (13.0 mg/capsule) and tadalafil (13.7 mg/capsule)

Recall Number:

N/A

Code Information:

Mj25 600M2, exp 12/31/2027

Not Yet Classified Drugs Event

Event ID:

98256

Status:

Ongoing

Recall Initiation Date:

12/15/2025

Center Classification Date:

N/A

Recalling Firm:

StuffbyNainax
2830 Lake Rd Apt 1102
Huntsville, TX 77340-5662
United States

Distribution Pattern:

Product was distributed to 4 customers in the US.

Product Type:

Drugs

Date Terminated:

N/A

Voluntary / Mandated:

Voluntary: Firm initiated

Initial Firm Notification of Consignee or Public:

Press Release

Associated Products

Product Description:

MR. 7 SUPER 700000 capsules, 1 capsule blister card, Distributed by mR. 7

Product Quantity:

4

Reason for Recall:

Marketed Without an Approved NDA/ANDA: FDA analysis revealed the presence of undeclared sildenafil and tadalafil

Recall Number:

N/A

Code Information:

All codes; Exp 12/31/2029

Not Yet Classified Drugs Event

Event ID:

98477

Product Type:

Drugs

Status:

Ongoing

Date Terminated:

N/A

Recall Initiation Date:

02/24/2026

Voluntary / Mandated:

Voluntary: Firm initiated

Center Classification Date:

N/A

Initial Firm Notification of Consignee or Public:

Letter

Recalling Firm:

MYLAN PHARMACEUTICALS INC
1311 Pineview Dr
Morgantown, WV 26505-3276
United States

Distribution Pattern:

Nationwide within the United States

Associated Products

Product Description:

Isotretinoin Capsules, USP 40 mg, packaged in 3 x 10-Count Prescription Packs, Rx Only, Manufactured for: Mylan Pharmaceuticals Inc., Morgantown, WV, 26505 USA, Made in France, NDC 0378-6614-93.

Product Quantity:

34850 blister packs

Reason for Recall:

Failed Dissolution Specifications

Recall Number:

N/A

Code Information:

Lot #: 3116110, 3116130, Exp. Date 07/2026

Not Yet Classified Drugs Event

Event ID:

98496

Product Type:

Drugs

Status:

Ongoing

Date Terminated:

N/A

Recall Initiation Date:
Voluntary / Mandated:

02/26/2026

Voluntary: Firm initiated

Center Classification Date:

N/A

Initial Firm Notification of Consignee or Public:

Letter

Recalling Firm:

New Life Pharma LLC
265 Livingston St
Northvale, NJ 07647-1901
United States

Distribution Pattern:

OH

Associated Products

<p>Product Description: Semaglutide Inj, 2mg x 10, Sterile Multi-Dose Vial, Rx only, Nomida, 25055 W Valley Pkwy, Suite 106, Olathe, KS 66061, NDC 84223-001-07.</p> <p>Product Quantity: 820 vials</p> <p>Reason for Recall: Lack of Assurance of Sterility</p> <p>Recall Number: N/A</p> <p>Code Information: Lot 241113, Exp Date 11/30/2026 & Lot 250103, Exp Date 1/31/2027</p>

<p>Product Description: Semaglutide Inj., 2 mg x 5, Sterile Multi-Dose Vial, Rx only, Nomida, 25055 W Valley Pkwy, Suite 106, Olathe, KS 66061, NDC 84223-001-06</p> <p>Product Quantity: 2 vials</p> <p>Reason for Recall: Lack of Assurance of Sterility</p> <p>Recall Number: N/A</p> <p>Code Information: Lot 240903, exp 9/30/2026</p>
--

<p>Product Description: Tirzepatide Inj, 10mg, Sterile Multi-Dose Vial, Rx only, Nomida, 25055 W Valley Pkwy, Suite 106, Olathe, KS 66061, NDC 84223-002-03.</p> <p>Product Quantity: 23 vials</p> <p>Reason for Recall: Lack of Assurance of Sterility</p> <p>Recall Number: N/A</p> <p>Code Information: Lot 240709, exp 7/31/2026</p>

<p>Product Description: Tirzepatide Inj, 15mg x 4, Sterile Multi-Dose Vial, Rx only, Nomida, 25055 W Valley Pkwy, Suite 106, Olathe, KS 66061, NDC 84223-002-04.</p> <p>Product Quantity: 45 vials</p> <p>Reason for Recall: Lack of Assurance of Sterility</p> <p>Recall Number:</p>

N/A

Code Information:

Lot 240709, exp 7/31/2026