Enforcement Report - Week of May 31, 2023

Class II Drugs Event

Event ID:
92058

Product Type:
Drugs

Status:
Ongoing

Date Terminated:

Recall Initiation Date:
03/17/2023

Voluntary / Mandated:
Voluntary: Firm initiated

Center Classification Date:
05/19/2023

Initial Firm Notification of Consignee or Public:
Letter

Recalling Firm:
Amerisource Health Services LLC
2550 John Glenn Ave Ste A
Columbus OH United States

Distribution Pattern:
Nationwide USA

Associated Products

Product Description:
Glimepiride Tablets, USP, 1 mg, RX, Packaged as a)100-count bottle, NDC# 68001-177-00; b) 500-count bottle, NDC# 68001-177-03; Manufactured by: Intas Pharmaceuticals Limited, Pharmez, Ahmedabad-382 213. INDIA. For BluePoint Laboratories

Product Quantity:
63,335 bottles

Reason for Recall:
CGMP Deviations: recalling drug products following an FDA inspection.

Recall Number:
D-0757-2023

Code Information:
Batches a) P2002616, EXP 04/30/2023; P2006509, EXP 11/30/2023; P2103572, EXP 04/30/2024; P2106811, EXP 09/30/2024; R2200578, EXP 04/30/2025 b)P2100095, EXP 11/30/2023; P2100624, EXP 01/31/2024; P2101780, EXP 02/29/2024; P2107383, EXP 09/30/2024; P2201505, EXP 02/28/2025; R2201109, EXP 06/30/2025

Product Description:
Glimepiride Tablets, USP, 2 mg, RX, Packaged as a) 100-count bottle, NDC# 68001-178-00; b) 500-count bottle; NDC# 68001-178-03 Manufactured by: Intas Pharmaceuticals Limited, Pharmez, Ahmedabad-382 213, INDIA. For BluePoint Laboratories

Product Quantity:
105,361 bottles

Reason for Recall:
CGMP Deviations: recalling drug products following an FDA inspection.

Recall Number:
D-0758-2023

Code Information:
Batches a)P2003493, EXP 05/31/2023; P2100120, EXP 11/30/2023; P2100683, EXP 01/31/2024; P2106002, EXP 07/31/2024; R2200148, EXP 12/31/2024; R2201125, EXP 06/30/2025 b) P2003403, EXP 05/31/2023; b) P2005800, EXP 09/30/2023; P2101156, EXP 01/31/2024; P2105401, EXP 07/31/2024; R2200083, EXP 12/31/2024; R2201004, EXP 07/31/2025

Product Description:
Glimepiride Tablets, USP, 4 mg, RX, Packaged as a) 100-count bottle, NDC# 68001-179-00; b) 500-count bottle, NDC# 68001-179-03, Manufactured by: Intas Pharmaceuticals Limited, Pharmez, Ahmedabad-382 213. INDIA. For BluePoint Laboratories

Product Quantity:

129,849 bottles

Reason for Recall:
CGMP Deviations: recalling drug products following an FDA inspection.

Recall Number:
D-0759-2023

Code Information:
Batches a) P2003403, EXP 05/31/2023; P2006593, EXP 11/30/2023; P2101152, EXP 01/31/2024; P2105014, EXP 06/30/2024; R2101440, EXP 09/30/2024; P2200774, EXP 01/31/2025; R2200664, EXP 04/30/2025 b) P2100705, EXP 01/31/2024; P2104672, EXP 06/30/2024; R2101435, EXP 09/30/2024; R2200102, EXP 12/31/2024; R2200577, EXP 04/30/2025; P2205870, EXP 08/31/2025 [500 count] Lot, expiry: P2100121, exp 11/30/2023; P2100705, exp 01/31/2024; P2104672, exp 06/30/2024; R2101435, exp 09/30/2024; R2200102, exp 12/31/2024; R2200577, exp R2200577;P2205870, exp 08/31/2025

Class II Drugs Event

Event ID:
92253

Product Type:
Drugs

Status:
Ongoing

Date Terminated:

Recall Initiation Date:
04/28/2023

Voluntary / Mandated:
Voluntary: Firm initiated

Center Classification Date:
05/19/2023

Initial Firm Notification of Consignee or Public:
E-Mail

Recalling Firm:
Central Admixture Pharmacy Services, Inc.
6580 Snowdrift Rd Ste 100
Allentown PA United States

Distribution Pattern:
Nationwide in the USA.

Associated Products

Product Description:
Microplegia (MSA/MSG 0.92 Molar) packaged in 125 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0001-1.

Product Quantity:
307 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0648-2023

Code Information:
Lot # 37-894942, Exp 05/01/2023; 37-896485, Exp 05/07/2023; 37-896793, Exp 05/08/2023; 37-898247, Exp 05/14/2023; 37-899149, Exp 05/19/2023; 37-900610, Exp 05/25/2023

Product Description:
Cardioplegia Solution, Warm Induction 4:1 High Potassium (40 mEq) packaged in 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0002-1.

Product Quantity:
215 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0649-2023

Code Information:
Lot # 37-893802, Exp 04/28/2023; 37-894943, Exp 05/01/2023; 37-896487, Exp 05/07/2023; 37-898248, Exp 05/14/2023; 37-899171, Exp 05/19/2023

Product Description:
Cardioplegia Solution, Reperfusate No Potassium, packaged in 238.75 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0005-1.

Product Quantity:
255 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0650-2023

Code Information:
Lot # 37-893790, Exp 04/28/2023; 37-896488, Exp 05/07/2023; 37-898250, Exp 05/14/2023; 37-899235, Exp 05/19/2023; 37-900609, Exp 05/25/2023

Product Description:
Cardioplegia Solution, Reperfusate No Potassium, packaged in 477.5 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0006-1.

Product Quantity:
151 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0651-2023

Code Information:
Lot # 37-895691, Exp 05/05/2023; 37-896792, Exp 05/08/2023; 37-899170, Exp 05/19/2023

Product Description:
Cardioplegia Solution, Reperfusate 4:1 low potassium, 7.5 mEq K, packaged in 238.75 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0007-1.

Product Quantity:
140 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0652-2023

Code Information:
Lot # 37-896489, Exp 05/07/2023; 37-897805, Exp 05/12/2023; 37-900959, Exp 05/26/2023

Product Description:
Cardioplegia Solution, Reperfusate 4:1 low potassium, 15 mEq K, packaged in 477.5 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0008-1.

Product Quantity:
338 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0653-2023

Code Information:
Lot # 37-893847, Exp 04/28/2023; 37-895646, Exp 05/05/2023; 37-897804, Exp 05/12/2023; 37-899236, Exp 05/19/2023

Product Description:
Cardioplegia Solution, Reperfusate 4:1 low potassium/low tromethamine, 15 mEq K, packaged in 500 mL per bag, Rx only, Central Admixture

Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0009-1.

Product Quantity:
254 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0654-2023

Code Information:
Lot # 37-893848, Exp 04/28/2023; 37-895681, Exp 05/05/2023; 37-899250, Exp 05/19/2023; 37-901355, Exp 05/27/2023

Product Description:
Cardioplegia Solution, Warm Induction 4:1 HIGH POTASSIUM/low tromethamine, 40 mEq K, packaged in 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0011-1.

Product Quantity:
12 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0655-2023

Code Information:
Lot # 37-897075, Exp 05/11/2023

Product Description:
Microplegia Solution, MSA/MSG 0.92 Molar with CP2D, packaged in 120 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0012-2.

Product Quantity:
144 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0656-2023

Code Information:
Lot # 37-893787, Exp 04/28/2023; 37-895645, Exp 05/05/2023; 37-897071, Exp 05/11/2023; 37-897802, Exp 05/12/2023; 37-899136, Exp 05/19/2023

Product Description:
Transplant Solution (Plasma-Lyte A), packaged in 165 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0014-2.

Product Quantity:
24 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0657-2023

Code Information:
Lot # 37-897801, Exp 05/12/2023

Product Description:
Cardioplegia Solution, Induction 4:1, High Potassium, 60 mEq K, packaged in 830 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0100-1.

Product Quantity:
267 bags

Reason for Recall:

Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0658-2023

Code Information:
Lot # 37-894304, Exp 04/29/2023; 37-894946, Exp 05/01/2023; 37-896482, Exp 05/07/2023; 37-897933, Exp 05/13/2023; 37-900461, Exp 05/22/2023

Product Description:
Cardioplegia Solution, Induction 4:1, HIGH POTASSIUM/low tromethamine, 36 mEq K, packaged in 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-0101-1.

Product Quantity:
350 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0659-2023

Code Information:
Lot # 37-895647, Exp 05/05/2023; 37-898310, Exp 05/14/2023; 37-899622, Exp 05/20/2023; 37-900951, Exp 05/26/2023

Product Description:
Cardioplegia Solution, Induction 8:1 High Potassium, 108 mEq K, packaged in 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0102-1.

Product Quantity:
544 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0660-2023

Code Information:
Lot # 37-894305, Exp 04/29/2023; 37-896153, Exp 05/06/2023; 37-897073, Exp 05/11/2023; 37-897925, 37-897941, Exp 05/13/2023; 37-899607, 37-899608, Exp 05/20/2023; 37-900320, Exp 05/22/2023

Product Description:
Cardioplegia Solution, Maintenance 4:1 low potassium, 20 mEq K, packaged in 810 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0103-1.

Product Quantity:
269 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0661-2023

Code Information:
Lot # 37-893803, Exp 04/28/2023; 37-896483, Exp 05/07/2023; 37-897939, Exp 05/13/2023; 37-900952, Exp 05/26/2023

Product Description:
Cardioplegia Solution, Maintenance 4:1 low potassium/low tromethamine, 36 mEq K, packaged in 1000 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0104-1.

Product Quantity:
476 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0662-2023

Code Information:
Lot # 37-893850, Exp 04/28/2023; 37-894947, Exp 05/01/2023; 37-895695, Exp 05/05/2023; 37-896484, Exp 05/07/2023; 37-898314, Exp 05/14/2023; 37-898811, Exp 05/18/2023; 37-900616, Exp 05/25/2023

Product Description:
Cardioplegia Solution, Maintenance 8:1 low potassium, 24 mEq K, packaged in 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0105-1.

Product Quantity:
308 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0663-2023

Code Information:
Lot # 37-893846, Exp 04/28/2023; 37-895704, Exp 05/05/2023; 37-897079, Exp 05/11/2023; 37-897927, Exp 05/13/2023; 37-898814, Exp 05/18/2023; 37-900617, Exp 05/25/2023

Product Description:
Cardioplegia Solution, Induction 4:1 High Potassium, 30 mEq K, packaged in 415 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0106-1.

Product Quantity:
84 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0664-2023

Code Information:
Lot # 37-894307, Exp 04/29/2023; 37-896158, Exp 05/06/2023; 37-898281, Exp 05/14/2023; 37-900321, Exp 05/22/2023

Product Description:
Cardioplegia Solution, Induction 8:1 High Potassium/low dextrose, 100 mEq K, packaged in 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0107-1.

Product Quantity:
48 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0665-2023

Code Information:
Lot # 37-897926, Exp 05/13/2023; 37-900464, Exp 05/22/2023

Product Description:
Cardioplegia Solution, Induction 4:1 Plasma-Lyte/Tromethamine, High Potassium, packaged in 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0111-1.

Product Quantity:
613 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0666-2023

Code Information:
Lot # 37-894299, Exp 04/29/2023; 37-897051, Exp 05/11/2023; 37-897368, Exp 05/12/2023; 37-898794, Exp 05/18/2023; 37-899614, Exp 05/20/2023; 37-900608, Exp 05/25/2023

Product Description:
Cardioplegia Solution, Maintenance 4:1 Plasma-Lyte/Tromethamine, low potassium, packaged in 1000 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0112-1.

Product Quantity:
476 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0667-2023

Code Information:
Lot # 37-894300, Exp 04/29/2023; 37-896114, Exp 05/06/2023; 37-896417, Exp 05/07/2023; 37-897388, Exp 05/12/2023; 37-898798, Exp 05/18/2023; 37-899618, Exp 05/20/2023

Product Description:
Cardioplegia Solution, del Nido Formula, packaged in 1,052.8 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0202-1.

Product Quantity:
12,888 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0668-2023

Code Information:
Lot # 37-890033, 37-890035, Exp 04/28/2023; 37-890518, 37-890527, 37-890536, 37-890538, Exp 04/29/2023; 37-890844, 37-890845, Exp 04/30/2023; 37-891159, 37-891161, 37-891187, 37-891190, 37-891212, 37-891213, 37-891222, 37-891223, Exp 05/01/2023; 37-891432, 37-891433, 37-891434, 37-891435, 37-891440, 37-891442, 37-891445, 37-891446, 37-891447, Exp 05/02/2023; 37-891746, 37-891754, 37-891755, Exp 05/05/2023; 37-892005, 37-892007, Exp 05/06/2023; 37-892502, 37-892503, 37-892505, 37-892517, 37-892518, 37-892519, 37-892538, Exp 05/07/2023; 37-892871, 37-892873, 37-892874, 37-892879, 37-892880, 37-892886, 37-892887, 37-892888, 37-892899, Exp 05/08/2023; 37-893129, 37-893131, 37-893158, 37-893160, 37-893162, 37-893163, 37-893164, Exp 05/09/2023; 37-893681, 37-893689, 37-893698, 37-893760, 37-893761, 37-893762, Exp 05/13/2023; 37-894275, 37-894276, 37-894277, 37-894301, Exp 05/14/2023; 37-894651, 37-894654, 37-894658, 37-894660, 37-894662, 37-894663, Exp 05/15/2023; 37-894948, 37-894967, 37-894970, Exp 05/16/2023; 37-895276, 37-895278, 37-895281, 37-895284, 37-895285, 37-895286, 37-895289, 37-895297, Exp 05/19/2023; 37-895650, 37-895652, Exp 05/20/2023; 37-896051, 37-896052, 37-896054, 37-896055, 37-896057, 37-896058, 37-896455, Exp 05/21/2023; 37-896418, 37-896419, 37-896420, 37-896438, 37-896453, 37-896454, 37-896855, Exp 05/22/2023; 37-896814, 37-896816, 37-896818, 37-896823, 37-896824, 37-896841, 37-896843, 37-896942, Exp 05/23/2023; 37-897417, 37-897429, Exp 05/27/2023; 37-897856, 37-897857, 37-897858, 37-897880, 37-897881, 37-897882, 37-898264, Exp 05/28/2023; 37-898180, 37-898222, 37-898225, 37-898226, 37-898227, 37-898228, Exp 05/29/2023; 37-898487, 37-898488, 37-898490, Exp 05/30/2023; 37-898781, 37-898782, 37-898784, 37-898787, 37-898788, Exp 06/02/2023; 37-899102, 37-899103, 37-899104, 37-899105, Exp 06/03/2023; 37-899590, 37-899598, 37-899599, 37-900116, Exp 06/04/2023; 3700000900051, 37-900052, 37-900054, 37-900055, Exp 06/05/2023; 37-900306, 37-900309, 37-900310, 37-900311, 37-900313, 37-900314, 37-900315, 37-900316, Exp 06/06/2023

Product Description:
Cardioplegia Solution, Modified St Thomas Solution, low potassium, HIGH SODIUM BICARBONATE, 62 mEq K, packaged in 1000 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0203-1.

Product Quantity:
182 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0669-2023

Code Information:
Lot # 37-894704, Exp 04/30/2023; 37-896126, Exp 05/06/2023; 37-898276, Exp 05/14/2023; 37-899141, Exp 05/19/2023

Product Description:
Cardioplegia Solution, Modified St Thomas Solution, HIGH POTASSIUM, HIGH SODIUM BICARBONATE, 106 mEq K, packaged in 1000 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0204-1.

Product Quantity:
241 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0670-2023

Code Information:
Lot # 37-894705, Exp 04/30/2023; 37-896127, Exp 05/06/2023; 37-898278, Exp 05/14/2023; 37-899148, Exp 05/19/2023; 37-900619, Exp 05/25/2023

Product Description:
Cardioplegia Solution, Maintenance 4:1 in Ringer's, low potassium, 12 mEq K, packaged in 504.8 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0205-1.

Product Quantity:
280 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0671-2023

Code Information:
Lot # 37-894325, Exp 04/29/2023; 37-895211, Exp 05/04/2023; 37-899265, Exp 05/19/2023

Product Description:
Cardioplegia Solution, Induction 4:1 in Ringer's, HIGH POTASSIUM, 48 mEq K, packaged in 522.8 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0206-1.

Product Quantity:
333 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0672-2023

Code Information:
Lot # 37-894326, Exp 04/29/2023; 37-895212, Exp 05/04/2023; 37-896794, Exp 05/08/2023; 37-897286, Exp 05/12/2023; 37-899267, Exp 05/19/2023

Product Description:
Cardioplegia Solution, Modified St Thomas Formula, HIGH POTASSIUM, 122 mEq K, packaged in 1000 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0209-1.

Product Quantity:
329 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0673-2023

Code Information:
Lot # 37-893789, Exp 04/28/2023; 37-894318, Exp 04/29/2023; 37-895648, Exp 05/05/2023; 37-897301, Exp 05/12/2023; 37-898815, Exp 05/18/2023

Product Description:
Cardioplegia Solution, Modified St Thomas Formula, low potassium, 70 mEq K, packaged in 1000 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0210-1.

Product Quantity:
208 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0674-2023

Code Information:
Lot # 37-894320, Exp 04/29/2023; 37-894324, Exp 04/29/2023; 37-897297, Exp 05/12/2023; 37-898509, Exp 05/15/2023

Product Description:
Cardioplegia Solution, Maintenance 4:1 Plasmalyte, low potassium, low K, packaged in 1047 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0211-1.

Product Quantity:
226 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0675-2023

Code Information:
Lot # 37-894309, 37-894321, Exp 04/29/2023; 37-896159, Exp 05/06/2023; 37-898510, Exp 05/15/2023; 37-900324, Exp 05/22/2023

Product Description:
Cardioplegia Solution, Induction 4:1 Plasmalyte, HIGH POTASSIUM, HIGH K, packaged in 542 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0212-1.

Product Quantity:
52 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0676-2023

Code Information:
Lot # 37-895217, Exp 05/04/2023; 37-898803, Exp 05/18/2023; 37-900614, Exp 05/25/2023

Product Description:
Microplegia Solution, HIGH POTASSIUM (100 mEq), packaged in 200 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0213-1.

Product Quantity:
79 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0677-2023

Code Information:
Lot # 37-895226, Exp 05/04/2023; 37-900615, Exp 05/25/2023

Product Description:
Cardioplegia Solution, Induction 8:1 non-enriched, HIGH POTASSIUM, 70 mEq K, packaged in 300 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0214-1.

Product Quantity:
363 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0678-2023

Code Information:
Lot # 37-894310, Exp 04/29/2023; 37-895259, Exp 05/04/2023; 37-898303, Exp 05/14/2023; 37-901367, Exp 05/27/2023

Product Description:
Cardioplegia Solution, Maintenance 8:1 non-enriched, low potassium, 24 mEq K, packaged in 300 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0215-1.

Product Quantity:
325 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0679-2023

Code Information:
Lot # 37-895694, 37-896050, Exp 05/05/2023; 37-898304, Exp 05/14/2023; 37-899234, Exp 05/19/2023; 37-901375, Exp 05/27/2023

Product Description:
Cardioplegia Solution, LEESBURG CARDIOPLEGIA, packaged in 1030.2 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0218-1.

Product Quantity:
79 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0680-2023

Code Information:
Lot # 37-895256, Exp 05/04/2023; 37-898816, Exp 05/18/2023

Product Description:
Modified del Nido Microplegia, packaged in 40 mL per syringe, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0219-1.

Product Quantity:
690 syringes

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0681-2023

Code Information:
Lot # 37-893821, Exp 04/28/2023; 37-894813, 37-894815, Exp 04/30/2023; 37-895489, 37-895490, Exp 05/05/2023; 37-897810, 37-897811, Exp 05/13/2023; 37-899585, Exp 05/20/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 2%/Dextrose 10%, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0400-1.

Product Quantity:
80 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0682-2023

Code Information:
Lot # 37-895214, Exp 05/04/2023; 37-897950, Exp 05/13/2023; 37-899603, Exp 05/20/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 3%/Dextrose 10% with CALCIUM, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0405-1.

Product Quantity:

558 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0683-2023

Code Information:
Lot # 37-893806, Exp 04/28/2023; 37-894701, Exp 04/30/2023; 37-895626, Exp 05/05/2023; 37-897277, Exp 05/12/2023; 37-899175, Exp 05/19/2023; 37-900971, Exp 05/26/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 3.5%/Dextrose 10%, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0406-1.

Product Quantity:
71 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0684-2023

Code Information:
Lot # 37-894328, Exp 04/29/2023; 37-897792, Exp 05/12/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 3.5%/Dextrose 10% with CALCIUM, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0407-1.

Product Quantity:
138 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0685-2023

Code Information:
Lot # 37-894709, Exp 04/30/2023; 37-896128, Exp 05/06/2023; 37-897894, Exp 05/13/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 4%/Dextrose 10%, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0408-1.

Product Quantity:
167 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0686-2023

Code Information:
Lot # 37-896130, Exp 05/06/2023; 37-897957, Exp 05/13/2023; 37-899605, Exp 05/20/2023; 37-901357, Exp 05/27/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 3%/Dextrose 5% with CALCIUM, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0413-1.

Product Quantity:
258 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:

D-0687-2023

Code Information:
Lot # 37-894313, Exp 04/29/2023; 37-895257, Exp 05/04/2023; 37-897360, Exp 05/12/2023; 37-897890, Exp 05/13/2023; 37-900964, Exp 05/26/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 2%/Dextrose 10% with CALCIUM and HEPARIN, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0416-1.

Product Quantity:
139 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0688-2023

Code Information:
Lot # 37-897800, Exp 05/12/2023; 37-897895, Exp 05/13/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 2.5%/Dextrose 10% with CALCIUM and HEPARIN, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0418-1.

Product Quantity:
92 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0689-2023

Code Information:
Lot # 37-894334, Exp 04/29/2023; 37-898299, Exp 05/14/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 3%/Dextrose 5% with CALCIUM and HEPARIN, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0419-1.

Product Quantity:
418 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0690-2023

Code Information:
Lot # 37-893810, Exp 04/28/2023; 37-894698, Exp 04/30/2023; 37-896144, Exp 05/06/2023; 37-897896, Exp 05/13/2023; 37-899172, Exp 05/19/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 3%/Dextrose 10% with CALCIUM and HEPARIN, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0420-1.

Product Quantity:
877 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0691-2023

Code Information:
Lot # 37-894314, Exp 04/29/2023; 37-894690, 37-894699, Exp 04/30/2023; 37-895627, Exp 05/05/2023; 37-896494, Exp 05/07/2023; 37-897082, Exp 05/11/2023; 37-897955, Exp 05/13/2023; 37-899254, Exp 05/19/2023; 37-900057, 37-900060, Exp 05/21/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 3.5%/Dextrose 10% with CALCIUM and HEPARIN, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0421-1.

Product Quantity:
427 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0692-2023

Code Information:
Lot # 37-893804, Exp 04/28/2023; 37-894713, Exp 04/30/2023; 37-896134, Exp 05/06/2023; 37-896497, Exp 05/07/2023; 37-897310, Exp 05/12/2023; 37-899623, Exp 05/20/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 4%/Dextrose 10% with CALCIUM and HEPARIN, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0422-1.

Product Quantity:
747 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0693-2023

Code Information:
Lot # 37-893819, Exp 04/28/2023; 37-894702, 37-894710, Exp 04/30/2023; 37-895215, Exp 05/04/2023; 37-897282, Exp 05/12/2023; 37-897921, Exp 05/13/2023; 37-898309, Exp 05/14/2023; 37-899199, Exp 05/19/2023; 37-900058, Exp 05/21/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 6%/Dextrose 10% with CALCIUM and HEPARIN, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0423-1.

Product Quantity:
128 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0694-2023

Code Information:
Lot # 37-896152, Exp 05/06/2023; 37-897923, Exp 05/13/2023; 37-899173, Exp 05/19/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 2%/Dextrose 10% with low calcium and HEPARIN, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0425-1.

Product Quantity:
135 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0695-2023

Code Information:
Lot # 37-894712, Exp 04/30/2023; 37-897924, Exp 05/13/2023; 37-899606, Exp 05/20/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 3%/Dextrose 5% with low calcium and HEPARIN, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0427-1.

Product Quantity:
259 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0696-2023

Code Information:
Lot # 37-894338, Exp 04/29/2023; 37-896505, Exp 05/07/2023; 37-898301, Exp 05/14/2023; 37-898305, Exp 05/14/2023; 37-900061, Exp 05/21/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 3%/Dextrose 10% with low calcium and HEPARIN, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0428-1.

Product Quantity:
485 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0697-2023

Code Information:
Lot # 37-893805, Exp 04/28/2023; 37-894711, Exp 04/30/2023; 37-895258, Exp 05/04/2023; 37-896501, Exp 05/07/2023; 37-898160, Exp 05/13/2023; 37-900064, Exp 05/21/2023; 37-900620, Exp 05/25/2023

Product Description:
Neonatal PN Starter Bag, Amino Acids (Trophamine) 3.5%/Dextrose 10% with low calcium and HEPARIN, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100 Allentown, PA 18106, NDC 71285-0429-1.

Product Quantity:
328 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0698-2023

Code Information:
Lot # 37-894695, Exp 04/30/2023; 37-895693, Exp 05/05/2023; 37-900065, Exp 05/21/2023; 37-900622, Exp 05/25/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 6%/Dextrose 10% with low calcium and HEPARIN, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0430-1.

Product Quantity:
60 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0699-2023

Code Information:
Lot # 37-896506, Exp 05/07/2023; 37-898306, Exp 05/14/2023; 37-900062, Exp 05/21/2023

Product Description:
Neonatal TPN Starter Bag, Amino Acids (Trophamine) 4.5%/Dextrose 10% with HEPARIN, packaged in 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., Lehigh Valley, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-0432-1.

Product Quantity:
175 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0700-2023

Code Information:
Lot # 37-894962, Exp 05/01/2023; 37-898285, Exp 05/14/2023

Product Description:
EPINEPHrine added to dextrose 5%, 2 mg/250 mL* (8 mcg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6030-1.

Product Quantity:
3338 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0701-2023

Code Information:
Lot # 37-884448, Exp 05/04/2023; 37-887386, Exp 05/16/2023; 37-888271, Exp 05/21/2023; 37-891189, Exp 05/31/2023; 37-893456, Exp 06/11/2023; 37-894656, Exp 06/14/2023; 37-898333, Exp 06/28/2023; 37-898559, Exp 06/29/2023

Product Description:
oxyTOCIN 20 units added to dextrose 5%/Lactated Ringer's 1,000 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6031-1.

Product Quantity:
7595 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0702-2023

Code Information:
Lot # 37-883206, 37-883207, 37-883210, 37-883213, 37-883216, 37-883217, 37-883217, Exp 04/30/2023; 37-884447, 37-884450, Exp 05/04/2023; 37-886269, Exp 05/11/2023; 37-887516, Exp 05/16/2023; 37-888306, 37-888308, 37-888310, 37-888311, Exp 05/21/2023; 37-890097, 37-890099, 37-890101, 37-890108, Exp 05/28/2023; 37-892636, 37-892637, 37-892639, 37-892640, Exp 06/06/2023; 37-894451, 37-894452, 37-894476, 37-894482, Exp 06/13/2023; 37-895311, 37-895312, 37-895313, Exp 06/18/2023; 37-895804, Exp 06/19/2023; 37-896607, Exp 06/21/2023

Product Description:
oxyTOCIN 30 units added to dextrose 5%/Lactated Ringer's 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6032-1.

Product Quantity:
2624 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0703-2023

Code Information:
Lot # 37-883948, Exp 05/02/2023; 37-886037, 37-886040, Exp 05/10/2023; 37-889593, Exp 05/24/2023; 37-891270, Exp 05/31/2023; 37-893223, Exp 06/08/2023; 37-896230, Exp 06/20/2023; 37-898520, Exp 06/29/2023

Product Description:
oxyTOCIN 10 units added to Lactated Ringer's 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 72185-6036-1.

Product Quantity:
2989 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:

D-0704-2023

Code Information:
Lot # 37-891283, Exp 05/06/2023; 37-892501, Exp 05/12/2023; 37-894413, Exp 05/19/2023; 37-896228, Exp 05/26/2023; 37-896815, 37-896820, 37-896826, Exp 05/28/2023; 37-900126, Exp 06/10/2023

Product Description:
oxyTOCIN 15 units added to Lactated Ringer's 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6037-1.

Product Quantity:
560 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0705-2023

Code Information:
Lot # 37-891282, Exp 05/06/2023; 37-900124, Exp 06/10/2023

Product Description:
oxyTOCIN 20 units added to Lactated Ringer's 1,000 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6038-1.

Product Quantity:
14362 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0706-2023

Code Information:
Lot # 37-889158, 37-889187, 37-889189, Exp 04/28/2023; 37-889494, 37-889507, 37-889522, 37-889525, 37-889527, 37-889608, Exp 04/29/2023; 37-889906, 37-889912, 37-889913, Exp 04/30/2023; 37-890110, 37-890121, Exp 05/03/2023; 37-891160, 37-891162, 37-891186, 37-891188, 37-891199, 37-891211, Exp 05/06/2023; 37-891564, 37-891565, 37-891567, 37-891569, 37-891570, Exp 05/07/2023; 37-892591, Exp 05/12/2023; 37-892893, 37-892897, 37-892900, 37-892902, 37-892903, 37-892904, 37-892909, 37-892910, Exp 05/13/2023; 37-894715, 37-894720, 37-894749, 37-894754, 37-894762, 37-894763, 37-894765, Exp 05/20/2023; 37-894991, 37-894992, 37-894993, Exp 05/21/2023; 37-896187, 37-896188, 37-896190, 37-896191, 37-896199, 05/26/2023; 37-896521, 37-896522, 37-896523, Exp 05/27/2023; 37-896883, 37-896884, Exp 05/28/2023; 37-900067, 37-900068, Exp 06/10/2023

Product Description:
oxyTOCIN 30 units added to Lactated Ringer's 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6039-1.

Product Quantity:
21871 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0707-2023

Code Information:
Lot # 37-889853, 37-889854, 37-889856, Exp 04/30/2023; 37-890087, 37-890088, 37-890095, 37-890098, 37-890100, 37-890107, 37-890109, Exp 05/03/2023; 37-890876, 37-890878, 37-890885, 37-890891, Exp 05/05/2023; 37-891572, Exp 05/07/2023; 37-892504, 37-892510, 37-892512, 37-892520, 37-892560, 37-892575, 37-892585, 37-892587, Exp 05/12/2023; 37-892988, 37-892989, Exp 05/13/2023; 37-894323, 37-894356, 37-894366, 37-894385, 37-894386, 37-894402, 37-894409, 37-894410, Exp 05/19/2023; 37-894778, Exp 05/20/2023; 37-894998, 37-894999, 37-895000, 37-895001, 37-895002, Exp 05/21/2023; 37-896186, 37-896189, 37-896192, 37-896194, 37-896200, 37-896202, 37-896227, 37-896231, 37-896240, 37-896241, Exp 05/26/2023; 37-896844, 37-896851, 37-896853, 37-896856, Exp 05/28/2023; 37-897827, 37-897836, 37-897837, 37-897853, 37-897854, 37-897928, Exp 06/02/2023; 37-900117, 37-900122, 37-900123, Exp 06/10/2023

Product Description:
oxyTOCIN 15 units added to 0.9% sodium chloride 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 72185-6042-1.

Product Quantity:
3865 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0708-2023

Code Information:
Lot # 37-883926, Exp 05/02/2023; 37-886032, Exp 05/10/2023; 37-887681, Exp 05/17/2023; 37-888317, Exp 05/21/2023; 37-890906, Exp 05/30/2023; 37-892146, Exp 06/05/2023; 37-892507, Exp 06/06/2023; 37-894483, Exp 06/13/2023; 37-896595, Exp 06/21/2023; 37-898568, Exp 06/29/2023

Product Description:
oxyTOCIN 20 units added to 0.9% sodium chloride 1,000 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 72185-6043-1.

Product Quantity:
5544 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0709-2023

Code Information:
Lot # 37-883938, Exp 05/02/2023; 37-884910, 37-884915, Exp 05/07/2023; 37-886041, Exp 05/10/2023; 37-887515, 37-887520, Exp 05/16/2023; 37-888318, Exp 05/21/2023; 37-888847, Exp 05/22/2023; 37-890535, Exp 05/29/2023; 37-890957, Exp 05/30/2023; 37-891566, 37-891568, Exp 06/01/2023; 37-892156, Exp 06/05/2023; 37-893165, 37-893166, Exp 06/08/2023; 37-893994, 37-893995, Exp 06/12/2023; 37-895771, 37-895773, 37-895775, Exp 06/19/2023; 37-896614, Exp 06/21/2023; 37-900377, Exp 07/06/2023

Product Description:
oxyTOCIN 30 units added to 0.9% sodium chloride 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6044-1.

Product Quantity:
104807 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0710-2023

Code Information:
Lot # 37-883041, 37-883043, 37-883044, 37-883045, 37-883047, 37-883059, 37-883060, 37-883067, 37-883071, 37-883072, 37-883131, 37-883185, Exp 04/28/2023; 37-883492, 37-883499, 37-883501, 37-883503, 37-883514, 37-883525, 37-883526, 37-883539, 37-883540, 37-883548, 37-883549, 37-883550, 37-883551, Exp 05/01/2023; 37-884229, 37-884230, 37-884245, 37-884249, 37-884252, 37-884256, 37-884260, 37-884261, 37-884262, 37-884266, 37-884270, Exp 05/03/2023; 37-884452, 37-884453, 37-884457, 37-884461, 37-884464, 37-884466, 37-884468, 37-884470, 37-884472, 37-884473, Exp 05/04/2023; 37-884793, 37-884812, 37-884817, 37-884821, 37-884822, 37-884830, 37-884833, 37-884874, 37-884881, 37-884886, 37-884895, 37-884905, 37-884907, 37-884916, 37-884917, Exp 05/05/2023; 37-884807, 37-884941, 37-884942, 37-884943, 37-884944, 37-884945, 37-884946, 37-884954, 37-884955, 37-884956, Exp 05/07/2023; 37-885187, 37-885217, 37-885244, 37-885247, 37-885261, 37-885268, 37-885279, 37-885282, 37-885291, 37-885293, Exp 05/08/2023; 37-885584, 37-885598, 37-885608, Exp 05/09/2023; 37-886056, 37-886057, 37-886060, 37-886081, 37-886082, 37-886088, 37-886089, Exp 05/10/2023; 37-886251, 37-886256, 37-886257, 37-886259, 37-886263, 37-886264, Exp 05/11/2023; 37-886499, 37-886503, 37-886510, 37-886527, 37-886530, 37-886532, 37-886534, 37-886538, 37-886541, 37-886542, 37-886543, Exp 05/14/2023; 37-887304, 37-887306, 37-887307, 37-887308, 37-887309, 37-887310, 37-887311, 37-887312, Exp 05/16/2023; 37-887672, 37-887693, 37-887718, 37-887724, 37-887728, 37-887753, Exp 05/17/2023; 37-888081, 37-888082, 37-888084, 37-888086, Exp 05/18/2023; 37-888304, 37-888305, 37-888307, 37-888309, 37-888312, 37-888313, 37-888315, 37-888316, Exp 05/21/2023; 37-888695, 37-888705, 37-888722, 37-888742, 37-888764, Exp 05/22/2023; 37-889048, 37-889049, 37-889061, 37-889065, 37-889118, 37-889133, 37-889139, Exp 05/23/2023; 37-889530, 37-889534, 37-889541, 37-889561, 37-889562, 37-889563, 37-889571, 37-889573, 37-889703, Exp 05/24/2023; 37-889733, 37-889747, 37-889793, 37-889803, 37-889828, 37-889829, 37-889839, 37-889841, 37-889847, 37-889851, Exp 05/25/2023; 37-890413, 37-890419, 37-890427, 37-890430, 37-890435, 37-890443, 37-890444, 37-890473, Exp 05/29/2023; 37-890958, Exp 05/30/2023; 37-891217, 37-891221, 37-891224, 37-891229, 37-891231, 37-891233, 37-891248, Exp 05/31/2023; 37-891475, 37-891479, 37-891481, 37-891484, 37-891486, 37-891488, 37-891531, 37-891549, 37-891550, 37-891562, Exp 06/01/2023; 37-892164, 37-892183, 37-892195, Exp 06/05/2023; 37-892878, 37-892883, 37-892885, 37-892891, 37-892892, 37-892895, 37-892896, Exp 06/07/2023; 37-893167, 37-893168, 37-893200, 37-893203, 37-893205, 37-893206, 37-893219, Exp 06/08/2023; 37-893457, 37-893462, 37-893468, Exp 06/11/2023; 37-

893798, 37-893799, 37-893808, 37-893809, 37-893814, 37-893844, Exp 06/12/2023; 37-894655, 37-894661, 37-894664, 37-894666, 37-894686, 37-894703, Exp 06/14/2023; 37-895029, 37-895030, 37-895036, 37-895037, 37-895038, 37-895048, 37-895056, 37-895063, 37-895068, 37-895072, 37-895073, 37-895074, 37-895075, Exp 06/15/2023; 37-895260, 37-895271, 37-895273, 37-895274, Exp 06/18/2023; 37-895623, 37-895624, 37-895644, 37-895653, 37-895661, 37-895676, 37-895677, 37-895678, 37-895679, 37-895774, 37-895776, Exp 06/19/2023; 37-896536, 37-896540, 37-896541, 37-896542, 37-896543, 37-896547, 37-896566, 37-896585, 37-896587, 37-896596, 37-896597, 37-896598, 37-896599, 37-896615, Exp 06/21/2023; 37-896864, 37-896868, 37-896870, 37-896877, 37-896879, 37-896882, 37-896882, 37-896885, 37-896899, 37-896901, 37-896903, 37-896914, 37-896915, 37-896916, Exp 06/22/2023; 37-897454, 37-897500, Exp 06/26/2023; 37-898273, Exp 06/28/2023; 37-898517, Exp 06/29/2023

Product Description:
oxyTOCIN 60 units added to 0.9% sodium chloride 1,000 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6045-1.

Product Quantity:
1332 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0711-2023

Code Information:
Lot # 37-883073, Exp 04/28/2023; 37-887776, Exp 05/17/2023; 37-889209, Exp 05/23/2023; 37-892588, Exp 06/06/2023; 37-894449, Exp 06/13/2023; 37-898552, Exp 06/29/2023

Product Description:
dilTIAZem added to dextrose 5%, 125 mg/125 mL* (1 mg/mL), 125 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 72185-6054-1.

Product Quantity:
1230 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0712-2023

Code Information:
Lot # 37-896365, 37-896367, 37-896370, Exp 05/31/2023; 37-896857, 37-896858, 37-896859, Exp 06/02/2023

Product Description:
dilTIAZem added to 0.9% sodium chloride, 125 mg/125 mL* (1 mg/mL), 125 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6055-1.

Product Quantity:
3040 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0713-2023

Code Information:
Lot # 37-896374, 37-896375, Exp 05/31/2023; 37-896876, 37-896878, Exp 06/02/2023; 37-897077, 37-897078, Exp 06/05/2023; 37-900074, 37-900075, 37-900076, 37-900111, 37-900112, 37-900115, Exp 06/15/2023; 37-900328, Exp 06/16/2023

Product Description:
norepinephrine 4 mg added to dextrose 5% 250 mL*, 16 mcg/mL, 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6056-1.

Product Quantity:
2041 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:

D-0714-2023

Code Information:
Lot # 37-884760, Exp 05/05/2023; 37-885601, Exp 05/09/2023; 37-887668, Exp 05/17/2023; 37-889594, Exp 05/24/2023; 37-892622, Exp 06/06/2023; 37-894316, Exp 06/13/2023

Product Description:
PHENYLephrine added to 0.9% sodium chloride, 10 mg/250 mL* (40 mcg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6058-1.

Product Quantity:
6407 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0715-2023

Code Information:
Lot # 37-884272, Exp 05/03/2023; 37-887513, Exp 05/16/2023; 37-888083, Exp 05/18/2023; 37-889123, 37-889423, Exp 05/23/2023; 37-891563, Exp 06/01/2023; 37-892577, 37-892582, Exp 06/06/2023; 37-893248, 37-893257, Exp 06/08/2023; 37-894436, Exp 06/13/2023; 37-895805, Exp 06/19/2023; 37-898324, Exp 06/28/2023

Product Description:
potassium phosphate 15 mmole added to 0.9% sodium chloride 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6060-1.

Product Quantity:
9432 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0716-2023

Code Information:
Lot # 37-883769, Exp 05/02/2023; 37-884449, Exp 05/04/2023; 37-885585, Exp 05/09/2023; 37-885922, Exp 05/10/2023; 37-886526, Exp 05/14/2023; 37-887387, Exp 05/16/2023; 37-887659, 37-887662, Exp 05/17/2023; 37-889064, Exp 05/23/2023; 37-889735, 37-889745, Exp 05/25/2023; 37-890036, 37-890055, Exp 05/28/2023; 37-890934, Exp 05/30/2023; 37-892890, Exp 06/07/2023; 37-893813, Exp 06/12/2023; 37-894317, Exp 06/13/2023; 37-895275, Exp 06/18/2023; 37-896822, 37-896852, Exp 06/22/2023; 37-897826, 37-897839, Exp 06/27/2023

Product Description:
vancomycin added to 0.9% sodium chloride, 1 g/250 mL* (4 mg/mL), 25 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6064-1.

Product Quantity:
4390 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0717-2023

Code Information:
Lot # 37-885351, Exp 05/08/2023; 37-885979, Exp 05/10/2023; 37-887721, Exp 05/17/2023; 37-889212, Exp 05/23/2023; 37-889607, Exp 05/24/2023; 37-891247, Exp 05/31/2023; 37-892937, Exp 06/07/2023; 37-894025, Exp 06/12/2023; 37-894748, Exp 06/14/2023; 37-895282, Exp 06/18/2023; 37-896173, Exp 06/20/2023

Product Description:
vancomycin added to 0.9% sodium chloride, 750 mg/250 mL* (3 mg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6071-1.

Product Quantity:
3075 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0718-2023

Code Information:
Lot # 37-884161, Exp 05/03/2023; 37-885995, 37-885997, Exp 05/10/2023; 37-887729, Exp 05/17/2023; 37-889595, Exp 05/24/2023; 37-891272, Exp 05/31/2023; 37-892162, Exp 06/05/2023; 37-894716, Exp 06/14/2023

Product Description:
vancomycin added to dextrose 5%, 1.25 g/250 mL* (5 mg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6073-1.

Product Quantity:
13772 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0719-2023

Code Information:
Lot # 37-883198, 37-883202, 37-883203, 37-883204, Exp 04/30/2023; 37-883799, Exp 05/02/2023; 37-884159, Exp 05/03/2023; 37-884888, 37-884892, 37-884899, 37-884901, Exp 05/07/2023; 37-886836, 37-886855, 37-886861, Exp 05/15/2023; 37-887381, Exp 05/16/2023; 37-887664, Exp 05/17/2023; 37-888256, 37-888259, 37-888267, Exp 05/21/2023; 37-888624, 37-888625, Exp 05/22/2023; 37-890038, 37-890041, 37-890042, Exp 05/28/2023; 37-890331, 37-890336, 37-890350, Exp 05/29/2023; 37-891163, Exp 05/31/2023; 37-892008, 37-892012, 37-892015, Exp 06/05/2023; 37-892882, 37-892884, Exp 06/07/2023

Product Description:
vancomycin added to 0.9% sodium chloride, 1.25 g/250 mL* (5 mg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6074-1.

Product Quantity:
27780 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0720-2023

Code Information:
Lot # 37-883407, 37-883436, Exp 05/01/2023; 37-883796, 37-883797, 37-883800, 37-883802, 37-883806, Exp 05/02/2023; 37-884162, 37-884163, Exp 05/03/2023; 37-885332, 37-885336, 37-885337, Exp 05/08/2023; 37-885749, 37-885753, 37-885755, Exp 05/09/2023; 37-886008, 37-886027, 37-886031, 37-886035, Exp 05/10/2023; 37-886273, 37-886274, 37-886275, Exp 05/11/2023; 37-887448, 37-887457, Exp 05/16/2023; 37-887765, 37-887772, 37-887775, Exp 05/17/2023; 37-888062, 37-888072, Exp 05/18/2023; 37-888811, 37-888817, Exp 05/22/2023; 37-889193, 37-889208, 37-889210, Exp 05/23/2023; 37-889450, 37-889451, 37-889454, Exp 05/24/2023; 37-890940, 37-890941, 37-890952, Exp 05/30/2023; 37-891137, 37-891138, 37-891139, Exp 05/31/2023; 37-892633, 37-892634, 37-892635, Exp 06/06/2023; 37-892934, 37-892961, 37-892964, 37-892965, Exp 06/07/2023; 37-893524, Exp 06/11/2023; 37-893993, 37-893996, Exp 06/12/2023; 37-894465, 37-894466, 37-894467, 37-894468, Exp 06/13/2023; 37-894706, 37-894714, Exp 06/14/2023; 37-895680, 37-895682, Exp 06/19/2023; 37-896123, Exp 06/20/2023; 37-896564, 37-896579, 37-896586, Exp 06/21/2023

Product Description:
vancomycin added to dextrose 5%, 1.5 g/250 mL* (6 mg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6075-1.

Product Quantity:
10282 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0721-2023

Code Information:
Lot # 37-883196, 37-883197, 37-883205, Exp 04/30/2023; 37-884882, 37-884890, 37-884898, Exp 05/07/2023; 37-886840, 37-886858, 37-886863, Exp 05/15/2023; 37-887365, Exp 05/16/2023; 37-887663, Exp 05/17/2023; 37-888254, 37-888257, 37-888265, 37-888303, Exp 05/21/2023; 37-889453, Exp 05/24/2023; 37-890037, 37-890039, 37-890043, Exp 05/28/2023; 37-890330, 37-890333, 37-890344, Exp 05/29/2023; 37-892009, 37-

892011, 37-892016, Exp 06/05/2023; 37-892881, Exp 06/07/2023; 37-893448, Exp 06/11/2023; 37-895221, Exp 06/18/2023; 37-898530, Exp 06/29/2023

Product Description:
vancomycin added to 0.9% sodium chloride, 1.5 g/250 mL* (6 mg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6076-1.

Product Quantity:
9037 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0722-2023

Code Information:
Lot # 37-883390, 37-883435, Exp 05/01/2023; 37-884157, Exp 05/03/2023; 37-885716, 37-886036, 37-886039, Exp 05/09/2023; 37-887499, 37-887500, Exp 05/16/2023; 37-887778, Exp 05/17/2023; 37-888105, Exp 05/18/2023; 37-889216, Exp 05/23/2023; 37-890923, 37-890928, 37-890936, Exp 05/30/2023; 37-891271, Exp 05/31/2023; 37-892649, Exp 06/06/2023; 37-892962, Exp 06/07/2023; 37-893966, 37-893967, 37-893969, 37-893998, Exp 06/12/2023; 37-896160, 37-896162, 37-896171, Exp 06/20/2023

Product Description:
vancomycin added to 0.9% sodium chloride, 2 g/500 mL* (4 mg/mL), 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6078-1.

Product Quantity:
7632 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0723-2023

Code Information:
Lot # 37-884151, 37-884153, Exp 05/03/2023; 37-885670, 37-885869, 37-885900, Exp 05/09/2023; 37-886271, Exp 05/11/2023; 37-886857, 37-886862, 37-886865, 37-886923, Exp 05/15/2023; 37-889190, 37-889191, Exp 05/23/2023; 37-889596, Exp 05/24/2023; 37-890482, 37-890487, 37-890488, Exp 05/29/2023; 37-891230, Exp 05/31/2023; 37-892589, 37-892590, Exp 06/06/2023; 37-893779, 37-893780, 37-893781, Exp 06/12/2023; 37-894665, 37-894667, Exp 06/14/2023; 37-895608, 37-895625, Exp 06/19/2023; 37-896519, Exp 06/21/2023

Product Description:
PHENYLephrine added to 0.9% sodium chloride, 40 mg/250 mL* (160 mcg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6092-1.

Product Quantity:
14205 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0724-2023

Code Information:
Lot # 37-883910, Exp 05/02/2023; 37-884221, Exp 05/03/2023; 37-884756, Exp 05/05/2023; 37-885358, Exp 05/08/2023; 37-885683, Exp 05/09/2023; 37-886267, 37-886268, Exp 05/11/2023; 37-886539, Exp 05/14/2023; 37-887458, 37-887459, Exp 05/16/2023; 37-888080, Exp 05/18/2023; 37-889192, Exp 05/23/2023; 37-889575, 37-889578, Exp 05/24/2023; 37-889852, 37-889855, Exp 05/25/2023; 37-890120, 37-890126, Exp 05/28/2023; 37-891485, 37-891491, Exp 06/01/2023; 37-892901, 37-892966, Exp 06/07/2023; 37-893236, 37-893237, Exp 06/08/2023; 37-893997, Exp 06/12/2023; 37-894777, Exp 06/14/2023; 37-895317, 37-895321, 37-895323, Exp 06/18/2023; 37-896201, Exp 06/20/2023; 37-896923, Exp 06/22/2023; 37-898558, Exp 06/29/2023

Product Description:
vancomycin added to 0.9% sodium chloride, 1.5 g/500 mL* (3 mg/mL), 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-6176-1.

Product Quantity:
11338 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0725-2023

Code Information:
Lot # 37-883372, 37-883387, Exp 05/01/2023; 37-883790, Exp 05/02/2023; 37-884160, Exp 05/03/2023; 37-885320, 37-885321, 37-885324, Exp 05/08/2023; 37-885921, 37-885928, 37-885930, Exp 05/10/2023; 37-886270, Exp 05/11/2023; 37-887511, Exp 05/16/2023; 37-887666, 37-887670, Exp 05/17/2023; 37-888115, Exp 05/18/2023; 37-888841, Exp 05/22/2023; 37-889457, 37-889462, Exp 05/24/2023; 37-892638, 37-892641, 37-892648, Exp 06/06/2023; 37-892907, Exp 06/07/2023; 37-894437, 37-894443, 37-894450, Exp 06/13/2023; 37-895707, 37-895718, 37-895723, Exp 06/19/2023; 37-896532, 37-896535, Exp 06/21/2023; 37-898527, Exp 06/29/2023

Product Description:
heparin added to 0.9% sodium chloride, 7,500 units/1,000 mL* (7.5 units/mL), 1000 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-7009-1.

Product Quantity:
3055 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0726-2023

Code Information:
Lot # 37-888668, 37-888685, 37-888701, Exp 05/02/2023; 37-890550, 37-890556, Exp 05/09/2023; 37-892201, 37-892216, Exp 05/16/2023; 37-893866, 37-893917, 37-893918, Exp 05/23/2023; 37-895705, 37-895717, Exp 05/30/2023

Product Description:
PHENYLephrine added to 0.9% sodium chloride, 25 mg/250 mL* (100 mcg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-7011-1.

Product Quantity:
5304 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0727-2023

Code Information:
Lot # 37-883924, Exp 05/02/2023; 37-884918, Exp 05/07/2023; 37-885677, Exp 05/09/2023; 37-887318, Exp 05/16/2023; 37-889211, Exp 05/23/2023; 37-889848, Exp 05/25/2023; 37-891150, Exp 05/31/2023; 37-892482, Exp 06/06/2023; 37-894484, Exp 06/13/2023; 37-896458, 37-896480, Exp 06/21/2023; 37-897967, Exp 06/27/2023; 37-898551, Exp 06/29/2023

Product Description:
potassium phosphate 30 mmole added to 0.9% sodium chloride 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-7016-1.

Product Quantity:
2146 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0728-2023

Code Information:
Lot # 37-883219, Exp 04/30/2023; 37-885926, Exp 05/10/2023; 37-887382, Exp 05/16/2023; 37-889455, Exp 05/24/2023; 37-890060, Exp 05/28/2023; 37-893444, Exp 06/11/2023; 37-894987, Exp 06/15/2023

Product Description:
EPINEPHrine added to dextrose 5%, 4 mg/250 mL* (16 mcg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-7018-1.

Product Quantity:

5627 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0729-2023

Code Information:
Lot # 37-883218, Exp 04/30/2023; 37-885235, Exp 05/08/2023; 37-885923, Exp 05/10/2023; 37-889050, Exp 05/23/2023; 37-890053, 37-890054, Exp 05/28/2023; 37-891439, 37-891441, Exp 06/01/2023; 37-892500, Exp 06/06/2023; 37-894308, Exp 06/13/2023; 37-896125, Exp 06/20/2023; 37-896926, Exp 06/22/2023; 37-900396, Exp 07/06/2023

Product Description:
EPINEPHrine added to dextrose 5%, 8 mg/250 mL* (32 mcg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-7019-1.

Product Quantity:
2542 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0730-2023

Code Information:
Lot # 37-884156, Exp 05/03/2023; 37-885207, Exp 05/08/2023; 37-885929, Exp 05/10/2023; 37-886496, Exp 05/14/2023; 37-886817, Exp 05/15/2023; 37-887420, Exp 05/16/2023; 37-893786, Exp 06/12/2023; 37-894303, Exp 06/13/2023; 37-897893, Exp 06/27/2023; 37-900073, Exp 07/05/2023

Product Description:
heparin added to 0.9% sodium chloride, 4,000 units/1,000 mL* (4 units/mL), 1,000 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-7022-1.

Product Quantity:
11117 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0731-2023

Code Information:
Lot # 37-888633, 37-888637, 37-888641, 37-888658, 37-888666, 37-888839, Exp 05/02/2023; 37-890332, 37-890383, 37-890394, 37-890424, 37-890428, 37-890442, Exp 05/09/2023; 37-892010, 37-892021, 37-892046, 37-892061, 37-892116, 37-892120, 37-892150, 37-892153, Exp 05/16/2023; 37-893467, 37-893469, 37-893470, 37-893471, 37-893508, 37-893512, 37-893519, Exp 05/22/2023; 37-893925, 37-893937, 37-893939, 37-893940, 37-893958, 37-893968, Exp 05/23/2023; 37-895218, 37-895220, 37-895224, 37-895231, 37-895236, 37-895305, 37-895306, Exp 05/29/2023; 37-895754, 37-895756, 37-895757, Exp 05/30/2023

Product Description:
heparin added to 0.9% sodium chloride, 5,000 units/500 mL* (10 units/mL), 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-7023-1.

Product Quantity:
5207 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0732-2023

Code Information:
Lot # 37-888736, 37-888769, Exp 05/02/2023; 37-890465, 37-890483, Exp 05/09/2023; 37-892088, 37-892117, 37-892119, 37-892145, Exp 05/16/2023; 37-893759, 37-893765, Exp 05/23/2023; 37-895742, 37-895751, Exp 05/30/2023

Product Description:
PHENYLephrine added to 0.9% sodium chloride, 20 mg/250 mL* (80 mcg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services,

Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-7025-1.

Product Quantity:
13953 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0733-2023

Code Information:
Lot # 37-883074, Exp 04/28/2023; 37-883237, 37-883238, Exp 04/30/2023; 37-883940, Exp 05/02/2023; 37-884271, Exp 05/03/2023; 37-884749, 37-884751, Exp 05/05/2023; 37-885338, Exp 05/08/2023; 37-885756, Exp 05/09/2023; 37-886272, Exp 05/11/2023; 37-887512, Exp 05/16/2023; 37-888027, 37-888030, 37-888040, 37-888052, 37-888063, Exp 05/18/2023; 37-888314, Exp 05/21/2023; 37-890489, 37-890497, 37-890534, Exp 05/29/2023; 37-890933, Exp 05/30/2023; 37-891152, Exp 05/31/2023; 37-892985, 37-892986, 37-892995, Exp 06/07/2023; 37-893511, 37-893517, Exp 06/11/2023; 37-896481, Exp 06/21/2023; 37-897828, Exp 06/27/2023; 37-898523, Exp 06/29/2023

Product Description:
NORepinephrine added to 0.9% sodium chloride, 16 mg/250 mL* (64 mcg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-7036-1.

Product Quantity:
6341 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0734-2023

Code Information:
Lot # 37-882998, 37-883005, 37-883145, Exp 04/28/2023; 37-883144, 37-883220, 37-883221, 37-883222, Exp 04/30/2023; 37-883925, Exp 05/02/2023; 37-884180, 37-884181, Exp 05/03/2023; 37-884505, 37-884506, 37-884507, 37-884508, 37-884509, Exp 05/04/2023; 37-884834, 37-884869, 37-884870, Exp 05/07/2023; 37-886058, Exp 05/10/2023; 37-886536, 37-886537, Exp 05/14/2023; 37-887062, 37-887389, Exp 05/15/2023; 37-887784, 37-887787, Exp 05/17/2023; 37-888029, 37-888048, Exp 05/18/2023; 37-888261, 37-888263, 37-888269, 37-888282, 37-888292, 37-888294, 37-888301, 37-888302, Exp 05/21/2023; 37-890058, 37-890064, 37-890074, 37-890076, Exp 05/28/2023; 37-891747, 37-891748, 37-891750, 37-891751, 37-891753, Exp 06/04/2023; 37-893530, 37-893531, 37-893532, Exp 06/11/2023; 37-894760, 37-894761, Exp 06/14/2023; 37-894988, 37-894990, Exp 06/15/2023; 37-896260, Exp 06/20/2023; 37-896621, Exp 06/21/2023; 37-898518, 37-898519, Exp 06/29/2023; 37-898789, 37-898790, 37-898791, 37-898792, 37-898793, Exp 07/02/2023; 37-899718, Exp 07/04/2023; 37-900591, 37-900596, Exp 07/09/2023; 37-901149, Exp 07/10/2023

Product Description:
phenylephrine 50 mg added to 0.9% sodium chloride 250 mL*, 200 mcg/mL*, 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-7039-1.

Product Quantity:
1735 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0735-2023

Code Information:
Lot # 37-887056, 37-887057, 37-887059, Exp 05/15/2023; 37-893807, Exp 06/12/2023; 37-896880, Exp 06/22/2023

Product Description:
vancomycin added to 0.9% sodium chloride, 1.75 g/500 mL* (3.5 mg/mL), 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-7060-1.

Product Quantity:
8252 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:

D-0736-2023

Code Information:
Lot # 37-884152, 37-884158, Exp 05/03/2023; 37-885300, 37-885302, 37-885307, Exp 05/08/2023; 37-887418, 37-887431, 37-887432, Exp 05/16/2023; 37-888850, 37-888852, 37-888853, Exp 05/22/2023; 37-889188, Exp 05/23/2023; 37-890892, 37-890894, 37-890897, Exp 05/30/2023; 37-891232, Exp 05/31/2023; 37-892014, 37-892017, Exp 06/05/2023; 37-894411, 37-894412, 37-894414, Exp 06/13/2023; 37-894659, Exp 06/14/2023; 37-896078, 37-896089, 37-896097, Exp 06/20/2023; 37-897369, Exp 06/26/2023

Product Description:
diphenhydrAMINE 25 mg added to 0.9% sodium chloride 50 mL in 100 mL Partial Additive Bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-7089-1.

Product Quantity:
5627 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0737-2023

Code Information:
Lot # 37-884145, 37-884154, Exp 05/03/2023; 37-885931, Exp 05/10/2023; 37-888293, Exp 05/21/2023; 37-889057, 37-889063, Exp 05/23/2023; 37-891437, Exp 06/01/2023; 37-893455, Exp 06/11/2023; 37-893788, Exp 06/12/2023; 37-894302, Exp 06/13/2023; 37-894652, Exp 06/14/2023; 37-896817, Exp 06/22/2023; 37-897263, Exp 06/26/2023

Product Description:
diphenhydrAMINE 50 mg added to 0.9% sodium chloride 50 mL in 100 mL Partial Additive Bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-7090-1.

Product Quantity:
3999 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0738-2023

Code Information:
Lot # 37-884155, Exp 05/03/2023; 37-885992, Exp 05/10/2023; 37-887348, Exp 05/16/2023; 37-887660, Exp 05/17/2023; 37-889052, Exp 05/23/2023; 37-891155, 37-891156, Exp 05/31/2023; 37-891443, Exp 06/01/2023; 37-892475, Exp 06/06/2023

Product Description:
heparin added to 0.9% sodium chloride, 2,500 units/250 mL* (10 units/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-8000-1.

Product Quantity:
1300 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0739-2023

Code Information:
Lot # 37-888810, Exp 05/02/2023; 37-892217, Exp 05/16/2023; 37-895759, Exp 05/30/2023; 37-899174, Exp 06/13/2023

Product Description:
oxyTOCIN 40 units added to 0.9% sodium chloride 1,000 bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-8069-1.

Product Quantity:
2186 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:

D-0740-2023

Code Information:
Lot # 37-886265, Exp 05/11/2023; 37-886266, Exp 05/11/2023; 37-889903, Exp 05/25/2023; 37-890551, Exp 05/29/2023; 37-892521, 37-892620, Exp 06/06/2023; 37-896246, 37-896247, Exp 06/20/2023; 37-898553, Exp 06/29/2023

Product Description:
tromethamine 0.3 Molar, 50 mL syringe, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-8086-2.

Product Quantity:
225 syringes

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0741-2023

Code Information:
Lot # 37-884580, Exp 05/04/2023; 37-887061, Exp 05/15/2023; 37-892981, Exp 06/07/2023

Product Description:
EPINEPHrine added to 0.9% sodium chloride, 4 mg/250 mL* (16 mcg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-8093-1.

Product Quantity:
8171 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0742-2023

Code Information:
Lot # 37-882988, Exp 04/28/2023; 37-883194, 37-883208, Exp 04/30/2023; 37-884746, Exp 05/05/2023; 37-885215, Exp 05/08/2023; 37-885927, Exp 05/10/2023; 37-887044, 37-887051, Exp 05/15/2023; 37-887667, Exp 05/17/2023; 37-888297, 37-888298, Exp 05/21/2023; 37-890886, Exp 05/30/2023; 37-891438, Exp 06/01/2023; 37-892497, Exp 06/06/2023; 37-893159, Exp 06/08/2023; 37-893843, Exp 06/12/2023; 37-894649, Exp 06/14/2023; 37-896928, Exp 06/22/2023; 37-899304, Exp 07/03/2023

Product Description:
NORepinephrine added to dextrose 5%, 16 mg/250 mL* (64 mcg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-8095-1.

Product Quantity:
2890 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0743-2023

Code Information:
Lot # 37-883007, 37-883015, 37-883017, Exp 04/28/2023; 37-883236, Exp 04/30/2023; 37-884187, 37-884189, 37-884190, Exp 05/03/2023; 37-884567, 37-884568, 37-884569, Exp 05/04/2023; 37-887789, 37-887790, Exp 05/17/2023; 37-889142, 37-889143, 37-889144, 37-889149, Exp 05/23/2023; 37-889606, Exp 05/24/2023; 37-890519, 37-890528, Exp 05/29/2023; 37-890946, 37-890951, Exp 05/30/2023; 37-891281, 37-891284, 37-891292, 37-891294, Exp 05/31/2023; 37-891756, 37-891757, Exp 06/04/2023; 37-897959, 37-897961, Exp 06/27/2023; 37-898795, Exp 07/02/2023

Product Description:
NORepinephrine added to 0.9% sodium chloride, 8 mg/250 mL* (32 mcg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-8096-1.

Product Quantity:
6414 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0744-2023

Code Information:
Lot # 37-882989, Exp 04/28/2023; 37-883827, 37-883829, 37-883830, 37-883833, 37-883858, Exp 05/02/2023; 37-884758, Exp 05/05/2023; 37-885357, Exp 05/08/2023; 37-885748, 37-885751, Exp 05/09/2023; 37-887409, 37-887410, Exp 05/16/2023; 37-888863, 37-888867, Exp 05/22/2023; 37-889226, Exp 05/23/2023; 37-889576, 37-889577, Exp 05/24/2023; 37-889804, 37-889817, 37-889830, Exp 05/25/2023; 37-891280, Exp 05/31/2023; 37-891758, 37-891759, Exp 06/04/2023; 37-894311, 37-894315, 37-894319, Exp 06/13/2023; 37-895304, Exp 06/18/2023; 37-896544, 37-896545, 37-896546, 37-896552, Exp 06/21/2023; 37-898514, 37-898515, 37-898516, Exp 06/29/2023; 37-899305, Exp 07/03/2023; 37-900362, Exp 07/06/2023

Product Description:
NORepinephrine added to 0.9% sodium chloride, 4 mg/250 mL* (16 mcg/mL), 250 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-8097-1.

Product Quantity:
4994 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0745-2023

Code Information:
Lot # 37-884510, Exp 05/04/2023; 37-885671, Exp 05/09/2023; 37-886285, Exp 05/11/2023; 37-888322, Exp 05/21/2023; 37-888851, Exp 05/22/2023; 37-889840, Exp 05/25/2023; 37-890130, Exp 05/28/2023; 37-890966, Exp 05/30/2023; 37-893529, Exp 06/11/2023; 37-894766, Exp 06/14/2023; 37-895300, 37-895301, Exp 06/18/2023; 37-898521, Exp 06/29/2023; 37-900338, Exp 07/06/2023

Product Description:
heparin added to 0.9% sodium chloride, 2,500 units/500 mL* (5 units/mL), 500 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-8100-1.

Product Quantity:
2992 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0746-2023

Code Information:
Lot # 37-888833, Exp 05/02/2023; 37-890537, Exp 05/09/2023; 37-892161, 37-892163, Exp 05/16/2023; 37-893979, 37-893992, Exp 05/23/2023; 37-895755, 37-895758, Exp 05/30/2023

Product Description:
vasopressin 20 units added to 0.9% sodium chloride 100 mL*, 0.2 units/mL*, 100 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-9000-1.

Product Quantity:
1858 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0747-2023

Code Information:
Lot # 37-886252, 37-886254, Exp 05/11/2023; 37-889741, 37-889746, Exp 05/25/2023; 37-891444, Exp 06/01/2023; 37-893155, 37-893156, Exp 06/08/2023; 37-896812, Exp 06/22/2023

Product Description:
vasopressin 40 units added to 0.9% sodium chloride 100 mL*, 0.4 units/mL*, 100 mL per bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-9001-1.

Product Quantity:
100 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0748-2023

Code Information:
Lot # 37-889738, 37-889742, Exp 05/25/2023

Product Description:
vasopressin 50 units added to 0.9% sodium chloride 50 mL*, 1 unit/mL*, 50 mL in 100 mL Partial Additive Bag, Rx only, Central Admixture Pharmacy Services, Inc., 6580 Snowdrift Rd., Ste 100, Allentown, PA 18106, NDC 71285-9002-1.

Product Quantity:
4259 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0749-2023

Code Information:
Lot # 37-883209, 37-883211, 37-883212, 37-883214, 37-883215, Exp 04/30/2023; 37-883776, Exp 05/02/2023; 37-884451, Exp 05/04/2023; 37-884805, 37-884806, 37-884808, 37-884810, 37-884811, Exp 05/07/2023; 37-886498, 37-886506, 37-886507, 37-886509, 37-886511, 37-886512, Exp 05/14/2023; 37-887319, 37-887321, Exp 05/16/2023; 37-889572, 37-889574, Exp 05/24/2023; 37-890881, 37-890883, 37-890884, 37-890887, Exp 05/30/2023; 37-892003, 37-892004, 37-892006, Exp 06/05/2023; 37-893157, Exp 06/08/2023; 37-893460, 37-893461, 37-893463, Exp 06/11/2023; 37-894619, 37-894648, 37-894650, Exp 06/14/2023; 37-895213, 37-895216, 37-895219, 37-895223, 37-895225, Exp 06/18/2023; 37-896813, Exp 06/22/2023; 37-897069, 37-897074, Exp 06/25/2023

Product Description:
MSA 7.84% MSG 8.56% (0.92M) Comp. Sol. 1000 ml bag, Rx only, Central Admixture Pharmacy Services, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-8029-1, code 7128580291.

Product Quantity:
653 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0750-2023

Code Information:
Lot # 37-882955, Exp 04/29/2023; 37-884753, 37-884755, Exp 05/06/2023; 37-887536, 37-887537, Exp 05/17/2023; 37-888532, Exp 05/23/2023; 37-892474, Exp 06/07/2023; 37-896761, Exp 06/23/2023; 37-897261, Exp 06/27/2023

Product Description:
Sodium Phosphates Injection 4 mEq/3 mMol/mL, 500 ml bag, Rx only, Central Admixture Pharmacy Services, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-8077-1, code 7128580771.

Product Quantity:
3165 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0751-2023

Code Information:
Lot # 37-883239, 37-883240, Exp 05/01/2023; 37-884419, 37-884423, Exp 05/04/2023; 37-889009, Exp 05/23/2023; 37-890261, 37-890262, Exp 05/29/2023; 37-891250, Exp 05/31/2023; 37-891412, Exp 06/01/2023; 37-895488, Exp 06/19/2023; 37-896456, Exp 06/21/2023; 37-898159, Exp 06/28/2023

Product Description:
Potassium Acetate Injection, 2 mEq/mL, 500 ml bag, Rx only, Central Admixture Pharmacy Services, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-8078-1, code 7128580781.

Product Quantity:

3326 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0752-2023

Code Information:
Lot # 37-884108, 37-884109, Exp 05/03/2023; 37-884957, Exp 05/07/2023; 37-885047, 37-885048, Exp 05/08/2023; 37-885919, Exp 05/10/2023; 37-889426, 37-889432, Exp 05/24/2023; 37-891933, Exp 06/05/2023; 37-894274, Exp 06/13/2023; 37-896048, 37-896049, Exp 06/20/2023; 37-897798, Exp 06/27/2023

Product Description:
Lidocaine 2% HCl Inj, 500mL bag, Rx only, Central Admixture Pharmacy Services, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-8091-1, code 7128580911.

Product Quantity:
1516 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0753-2023

Code Information:
Lot # 37-870983, Exp 06/10/2023; 37-879653, 37-879654, Exp 07/16/2023; 37-883186, Exp 07/29/2023; 37-888533, Exp 08/20/2023; 37-893416, Exp 09/09/2023

Product Description:
HyperLyte CR Injection, 500 mL bag, Rx only, Central Admixture Pharmacy Services, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-8094-1, code 7128580941.

Product Quantity:
554 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0754-2023

Code Information:
Lot # 37-874236, Exp 06/20/2023; 37-878694, Exp 07/10/2023

Product Description:
HyperLyte CR Injection, 250 mL bag, Rx only, Central Admixture Pharmacy Services, 6580 Snowdrift Road, Allentown, PA 18106, NDC 71285-8094-2, code 7128580942.

Product Quantity:
1673 bags

Reason for Recall:
Lack of Assurance of Sterility: after an FDA inspection called into question the sterility of the products intended to be sterile.

Recall Number:
D-0755-2023

Code Information:
Lot # 37-885553, 37-885554, Exp 05/09/2023; 37-887923, 37-887924, Exp 05/18/2023; 37-890778, 37-890779, Exp 05/30/2023; 37-892847, Exp 06/07/2023; 37-895049, Exp 06/15/2023; 37-898480, 37-898481, Exp 06/29/2023

Class II Drugs Event

Event ID:
92289

Product Type:
Drugs

Status:

Date Terminated:

Ongoing

Recall Initiation Date:
05/09/2023

Voluntary / Mandated:
Voluntary: Firm initiated

Center Classification Date:
05/23/2023

Initial Firm Notification of Consignee or Public:
Letter

Recalling Firm:
SUN PHARMACEUTICAL INDUSTRIES INC
2 Independence Way
Princeton NJ United States

Distribution Pattern:
Nationwide in the U.S.A

Associated Products

Product Description:
buPROPrion Hydrochloride Extended-Release Tablets, USP (SR) 150 mg, 60 Tablets, Rx Only, Distributed by: Sun Pharmaceutical Industries, Inc., Cranbury, NJ, Manufactured by: Sun Pharmaceutical Industries Limited, Gujrat, India, NDC 47335-737-86

Product Quantity:
5,344 Bottles

Reason for Recall:
Failed Dissolution Specifications; during stability testing

Recall Number:
D-0761-2023

Code Information:
Lots: HAC2240A, Exp 05/2023; HAC3162A, Exp 07/2023

Class II Drugs Event

Event ID:
92329

Product Type:
Drugs

Status:
Ongoing

Date Terminated:

Recall Initiation Date:
05/11/2023

Voluntary / Mandated:
Voluntary: Firm initiated

Center Classification Date:
05/25/2023

Initial Firm Notification of Consignee or Public:
Letter

Recalling Firm:
Revive Rx LLC dba Revive Rx Pharmacy
3831 Golf Dr Ste A
Houston TX United States

Distribution Pattern:
Nationwide in the US

Associated Products

Product Description:
Tirzepatide 10 mg/0.5 mL Sterile Solution-2 mL Vial, Rx Only, For Sub-Q Use Only, Compounded Drug Product By: Revive Rx, 3831 Golf Dr A, Houston, TX 77018, NDC: 99000-9278-64.

Product Quantity:
45 Vials

Reason for Recall:
Sub-potent Drug

Recall Number:
D-0771-2023

Code Information:
Lot: 1643397 BUD: 10/16/2023

Class II Drugs Event

Event ID:
92347

Product Type:
Drugs

Status:
Ongoing

Date Terminated:

Recall Initiation Date:
05/16/2023

Voluntary / Mandated:
Voluntary: Firm initiated

Center Classification Date:
05/22/2023

Initial Firm Notification of Consignee or Public:
E-Mail

Recalling Firm:
Alpha Aromatics
294 Alpha Dr
Pittsburgh PA United States

Distribution Pattern:
Product was distributed to one account in NC.

Associated Products

Product Description:
FOAMING HAND SANITIZER, fresh mint scent, 62% alcohol, 1 Gallon / 3.78 Liters bottle, 290 Alpha Drive RIDC industrial Park, Pittsburgh, PA 15238,

Product Quantity:
544 1 gallon bottles

Reason for Recall:
CGMP Deviations: FDA analysis found product to contain acetaldehyde and acetal above specification limits.

Recall Number:
D-0760-2023

Code Information:
Lot: 2020-013211 and 2020-013629, No EXP date on label.

Class II Drugs Event

Event ID:
92376

Product Type:
Drugs

Status:
Ongoing

Date Terminated:

Recall Initiation Date:
05/19/2023

Voluntary / Mandated:
Voluntary: Firm initiated

Center Classification Date:
05/23/2023

Initial Firm Notification of Consignee or Public:
Letter

Recalling Firm:
Sagent Pharmaceuticals Inc
1901 N Roselle Rd Ste 450
Schaumburg IL United States

Distribution Pattern:
Nationwide USA

Associated Products

Product Description:
Nafcillin for Injection, USP, 1 gram per vial, Rx only, Mfd. for SAGENT Pharmaceuticals, Schaumburg, IL 60195. NDC: 25021-139-10

Product Quantity:
2,249,980 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0762-2023

Code Information:
Lot: NFG101, EXP: 1/31/2024; Lot: NFG102, EXP: 9/30/2024; Lot: NFG201, 5/31/2025

Product Description:
Nafcillin for Injection, USP, 2 gram per vial, Rx only, Mfd. for SAGENT Pharmaceuticals, Schaumburg, IL 60195. NDC: 25021-140-10

Product Quantity:
2,249,980 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0763-2023

Code Information:
Lots: NFL101, NFL102, EXP: 1/31/2024; Lots: NFL103, NFL104, EXP: 5/31/2024; Lot: NFL201, EXP: 03/31/2025; Lot: NFL202, EXP: 5/31/2025;

Product Description:
Nafcillin for Injection, USP, 10 gram per Pharmacy Bulk Package, Rx only, Mfd. for SAGENT Pharmaceuticals, Schaumburg, IL 60195. NDC: 25021-141-99

Product Quantity:
370,200 bottles

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0764-2023

Code Information:
Lot: NFT101, EXP: 5/31/2024; Lots: NFT201, NFT202, EXP: 2/28/2025; Lots: NFT203, NFT204, EXP: 5/31/2025

Class II Drugs Event

Event ID:
92380

Product Type:
Drugs

Status:
Ongoing

Date Terminated:

Recall Initiation Date:
05/22/2023

Voluntary / Mandated:
Voluntary: Firm initiated

Center Classification Date:
05/24/2023

Initial Firm Notification of Consignee or Public:
Letter

Recalling Firm:
Aurobindo Pharma USA Inc.
279 Princeton Hightstown Rd
East Windsor NJ United States

Distribution Pattern:
The recalled product was distributed to one Retail Distributor (Walgreens) who further distributed Nationwide in the USA.

Associated Products

Product Description:
Pain Reliever, Acetaminophen USP Caplets, 500 mg, 225-count bottles packaged in a cardboard carton, Distributed by: Walgreen Co., 200 Wilmot Rd, Deerfield, IL 60015; NDC 0363-9947-35.

Product Quantity:
87,360 bottles

Reason for Recall:
Failed Impurities/Degradation Specifications: firm's investigation due to customer complaints for discoloration found that the product was out of specification for an impurity.

Recall Number:
D-0765-2023

Code Information:
Lot: P2200101, P2200178, Exp. date 11/2023; P2200230, Exp. date 12/2023

Class III Drugs Event

Event ID:
92133

Product Type:
Drugs

Status:
Ongoing

Date Terminated:

Recall Initiation Date:
04/27/2023

Voluntary / Mandated:
Voluntary: Firm initiated

Center Classification Date:
05/24/2023

Initial Firm Notification of Consignee or Public:
Letter

Recalling Firm:
Teva Pharmaceuticals USA Inc
400 Interpace Pkwy Bldg A
Parsippany NJ United States

Distribution Pattern:
US Nationwide

Associated Products

Product Description:
Fentanyl Buccal Tablets CII, 100mcg, packaged in cartons of 28 Buccal Tablets (4 tablets x 7 cards), Rx only, Distributed by: Mayne Pharma, Greenville, NC 27834, NDC 51862-634-28

Product Quantity:
438

Reason for Recall:
Labeling: Incorrect or Missing Package Insert

Recall Number:
D-0766-2023

Code Information:
Lot# 42617828, Exp 06/2023; 100020465, Exp 01/2024

Product Description:
Fentanyl Buccal Tablets CII, 200mcg, packaged in cartons of 28 Buccal Tablets (4 tablets x 7 cards), Rx only, Distributed by: Mayne Pharma, Greenville, NC 27834, NDC 51862-635-28

Product Quantity:
397

Reason for Recall:
Labeling: Incorrect or Missing Package Insert

Recall Number:
D-0767-2023

Code Information:
Lot #: 100020528, Exp 09/2024; 100026699, Exp 11/2024

Product Description:
Fentanyl Buccal Tablets CII, 400mcg, packaged in cartons of 28 Buccal Tablets (4 tablets x 7 cards), Rx only, Distributed by: Mayne Pharma, Greenville, NC 27834, NDC 51862-636-28

Product Quantity:
1272

Reason for Recall:
Labeling: Incorrect or Missing Package Insert

Recall Number:
D-0768-2023

Code Information:
Lot#: 100020351, Exp 11/2024; 100020522, Exp 09/2024; 100026700, Exp 11/2024

Product Description:
Fentanyl Buccal Tablets CII, 600mcg, packaged in cartons of 28 Buccal Tablets (4 tablets x 7 cards), Rx only, Distributed by: Mayne Pharma, Greenville, NC 27834, NDC 51862-637-28

Product Quantity:
1908

Reason for Recall:
Labeling: Incorrect or Missing Package Insert

Recall Number:
D-0769-2023

Code Information:
Lot#: 42617831, Exp 06/2023; 42619585, Exp 11/2023; 100029649, Exp 11/2024

Product Description:
Fentanyl Buccal Tablets CII, 800mcg, packaged in cartons of 28 Buccal Tablets (4 tablets x 7 cards), Rx only, Distributed by: Mayne Pharma, Greenville, NC 27834, NDC 51862-638-28

Product Quantity:
3032

Reason for Recall:
Labeling: Incorrect or Missing Package Insert

Recall Number:
D-0770-2023

Code Information:
Lot#: 42617832, Exp 06/2023; 42619530, Exp 08/2023; 100020532, Exp 11/2024

Class III Drugs Event

Event ID:
92249

Product Type:
Drugs

Status:
Ongoing

Date Terminated:

Recall Initiation Date:
05/08/2023

Voluntary / Mandated:
Voluntary: Firm initiated

Center Classification Date:
05/19/2023

Initial Firm Notification of Consignee or Public:
Letter

Recalling Firm:
Amneal Pharmaceuticals of New York, LLC

50 Horseblock Rd
Brookhaven NY United States

Distribution Pattern:
Nationwide in the USA and PR

Associated Products

Product Description:
Methylprednisolone Acetate Injectable Suspension, USP 80 mg/mL, For Intramuscular, Intrasynovial and Soft Tissue Injection Only, Not for Intravenous Use, 1 mL Single Dose Vial, Manufactured by: Amneal Pharmaceuticals Pvt. Ltd. Parenteral Unit, Ahmedabad 382213, India, Amneal Pharamceuticals LLC, Bridgewater, NJ 08807, NDC 70121-1574-01.

Product Quantity:
69,239 vials

Reason for Recall:
Labeling: Not Elsewhere Classified; A typographical error was observed in the NDC number on the preprinted Individual Folding Cartons (secondary packaging only). Incorrect NDC Number 70121-1573-1; Correct NDC Number 70121-1574-1.

Recall Number:
D-0756-2023

Code Information:
Lot Numbers: AP220481, Exp 09/2024; AP220536A, Exp 10/2024