Enforcement Report - Week of January 11, 2017

Class II Drugs Event

Event ID:
75289

Product Type:
Drugs

Status:
Ongoing

Date Terminated:

Recall Initiation Date:
09/21/2016

Voluntary / Mandated:
Voluntary: Firm Initiated

Center Classification Date:
01/04/2017

Initial Firm Notification of Consignee or Public:
Press Release

Recalling Firm:
Wells Pharmacy Network LLC
1210 SW 33rd Ave
Ocala FL United States

Distribution Pattern:
Nationwide

Associated Products

Product Description:
Testosterone Cypionate in Grapeseed Oil Injectable, 200 mg/mL, a) 1 mL b) 3 mL c) 5 mL and d) 10 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 694 vials b) 389 vials c) 695 vials d) 5193 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0327-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot codes: 03282016@16, 03302016@4, 04042016@10, 04262016@3, 05162016@2, 06032016@2, 07062016@10, 08262016@2 b) Known lot code: 04012016@4, 04182016@6, 04282016@7, 05022016@4, 06142016@19, 08102016@22. c) Known lot code: 04042016@7, 05022016@9, 05272016@2, 06272016@13, 07292016@8, 08192016@10 d) Known lot code: 03242016@12, 04082016@6, 04142016@2, 04192016@5, 04272016@4, 05092016@4, 05232016@17, 06122016@6, 06082016@2, 06142016@148, 06162016@9, 06172016@13, 06232016@78, 06282016@31, 07122016@20, 08012016@25, 08032016@12, 08042016@13.

Product Description:
ULTRA-TEST (CYP 80%/PROP 20%) Injectable, 250 mg/mL 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
157 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0328-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known Lot No. 03282016@22, 05162016@5, 07252016@16. Known lot codes: 03282016@22, 05162016@5, 07262016@16

Product Description:
Testosterone Enanthate in Grapeseed Oil Injectable, 200 mg/mL, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
1,250 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0329-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot codes: 03282016@3, 04132016@1, 05262016@3, 06242016@97, 07282016@1

Product Description:
QUAD3 (PAP/PHEN/ALPRO/ATRO) 30 mg/3 mg/30 mcg/0.1mL Injectable, a) 2 mL and 5 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 1,312 vials and b) 3,354 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0330-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 03292016@12, 0422016@10, 04222016@10, 05052016@33, 05122016@11, 05132016@19, 05242016@2, 06212016@17, 07262016@36, 09072016@21, 09142015@17 b) Known lot code: 03242016@9, 04112016@10, 04212016@19, 05042016@20, 05132016@21, 05202016@71, 06152016@7, 07122016@21, 08032016@17, 09092016@35.

Product Description:
Phenylephrine HCl Injectable, 1 mg/mL (0.1%), 5 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
1,968 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0331-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot codes: 03292016@3, 04202016@4, 05162016@6, 06102016@9, 07202016@44, 08032016@38

Product Description:
MIC Combo - Standard 15 mg/50 mg/100 mg/mL Injectable, 30 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
308 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0332-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 03292016@5, 04122016@76, 04202016@5, 05182016@2, 06102016@2, 06282016@71, 09092016@1

Product Description:
Nicotinamide Adenine Dinucleotide (NADH) - Lyophilized (P.F.) 100 mg Injectable, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
Unknown

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0333-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 03302016@13, 06302016@95, 08082016@58.

Product Description:
ULTRA-TEST in Grapeseed Oil (CYP 90%/PROP 10%) Injectable, 200 mg/mL, 8 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
714 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0334-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 03302016@5, 04202016@2, 05172016@5, 06082016@44, 06102016@6, 08042016@14

Product Description:
ULTRA-TEST (CYP 80%/PROP 20%) Injectable, 200 mg/mL, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
325 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0335-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04152016@2, 05052016@2, 05192016@7, 07142016@28, 09122016@23

Product Description:
Chorionic Gonadotropin, Lyophilized (HCG) 11,000 Unit Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
10 kits

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0336-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 04052016@16, 04072016@26, 04142016@65, 05172016@105, 06062016@137, 06152016@28, 06232016@61, 07272016@56, 08192016@108, 09062016@127 Known vial lot code: 03252016@6, 03312016@1, 04042016@30, 05052016@134, 05092016@57, 06072016@32, 06172016@120, 07222016@26, 07222016@26, 08122016@3, 08252016@3.

Product Description:
Alprostadil 20 mcg/mL Injectable, a) 5 mL and b) 10 mL vials, Compounded by Wells Pharmacy Network LLC, Ocala FL

Product Quantity:
a) 24 vials, b) 91 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0337-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 03312016@6 b) Known lot code: 04222016@8, 15202016@85, 06302016@4, 08012016@32

Product Description:
Testosterone Cypionate in Sesame Oil Injectable, 210 mg/mL, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
21 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0338-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 03312016@9

Product Description:
Alprostadil 40 mcg/mL Injectable, 10 mL vials, Compounded by Wells Pharmacy Network LLC, Ocala FL

Product Quantity:
37 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0339-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 05092016@3, 07192016@8

Product Description:
Arginine HCl 100 mg/mL Injectable,10 mL, Compounded by Wells Pharmacy Network, LLC, Ocala, FL

Product Quantity:
351 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0340-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 05122016@3, 07072016@15, 08302016@65

Product Description:
Arginine/Citrulline/Ornit+Lido 100/100/100/10 mg/mL Injectable, 10 mL vials, Compounded by Wells Pharmacy Network, LLC, Ocala, FL

Product Quantity:
30 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0341-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 06032016@7, 09092016@5

Product Description:
B-complex + Vit C 20 mg/mL Injectable, 30 mL vial, Compounded by Wells Pharmacy Network, LLC, Ocala, FL

Product Quantity:
180 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0342-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 03252016@2, 04292016@11, 06122016@9, 07212016@11

Product Description:
Bimix 30 mg/1 mg/mL Injectable, a) 5 mL and b) 10 mL vial, Compounded by Wells Pharmacy Network, LLC, Ocala, FL

Product Quantity:
a) 80 vials, b) 23 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0343-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 04012016@7, 04252016@17, 05262016@13, 06142016@39, 08052016@13 b) Known lot code: 05242016@11, 09072016@18

Product Description:
Carnitine-L 250 mg/mL Injectable, 30 mL vials, Rx only, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
44 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0344-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 07222016@2, 08172016@1

Product Description:
Ceftazidime, Lyophilized, Ophthalmic 2.25% Kit, Rx only, Compounded by Wells Pharmacy, Ocala , FL

Product Quantity:
46 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0345-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 06062016@118. Vial lot code 05272016@47

Product Description:
Chorionic Gonadotropin (New York State), 11,000 unit Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
18 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0346-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 06072016@44, 07292016@92, 08092016@80, 09132016@33. Known vial lot code: 04042016@30, 07222016@26, 040
42016@30, 08252016@3

Product Description:
Chorionic Gonadotropin, Lyophilized, 3,500 unit Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
1753 kits (number of vials per kit vary)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0347-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 04212016@46, 05232016@215, 05232016@216, 07052016@89, 08042016@103, 09132016@116. Known vial lot code:
04132016@91, 05162016@136, 06242016@96, 07282016@51, 09092016@20

Product Description:
Chorionic Gonadotropin, Lyophilized, 5,000 unit Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
5277 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0348-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 04142016@75, 04252016@52, 05042016@143, 05042016@158, 06092016@169, 06242016@158, 08022016@154, 08
112016@56, 08252016@82, 09082016@41. Known vial lot code: 04062016@7, 04182016@49, 04292016@80, 04282016@43, 06012016@44, 06152016@57, 07132016@68, 08042016@35, 08192016@45, 08242016@33

Product Description:
Chorionic Gonadotropin, Lyophilized, 6,000 unit Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
2,339 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0349-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code:04212016@49, 05262016@154, 06092016@181, 08082016@118, 08252016@89. Known vial lot code: 04142016@34, 0
5202016@55, 06032016@42, 08012016@41, 08192016@47.

Product Description:
Chorionic Gonadotropin, Lyophilized, 7,500 unit Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
1,638 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0350-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code:04052016@14, 05162016@126, 06092016@173, 06202016@137, 08042016@97, 09132016@117. Known vial lot code:
03242016@31, 04262016@25, 06032016@44, 06202016@137, 07252016@56, 09092016@17.

Product Description:
Chorionic Gonadotropin Lyophilized, 20,000 unit Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
908 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0351-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code:04212016@42, 06092016@164, 08122016@96. Known vial lot code: 04132016@96, 05242016@1, 08082016@50.

Product Description:
Chorionic Gonadotropin+B12(C), Lyophilized, 10,000 unit + 40 mcg Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
724 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0352-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 05042016@162, 06062016@122, 08192016@110. Known vial lot code: 04252016@168, 05252016@117, 08152016@4
0.

Product Description:
Dexamethasone Sodium Phosphate 200 mg/mL, Injectable, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
15 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0353-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 07082016@25.

Product Description:
Dexamethasone Sodium Phosphate 24 mg/mL (2.4%), Injectable, 2 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
16 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0354-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 08082016@45.

Product Description:
DMSO (Dimethyl Sulfoxide) with Lidocaine 50%/0.5% mL Solution, 50 mL bottle, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
438 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0355-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04132016@69, 04272016@9, 05142016@13, 06162016@129, 07052016@12, 07292016@2.

Product Description:
Fluorescein/Indocyanine, Lyophilized (P.F.), 400 mg/15 mg, 1 mL, Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
203 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0356-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 06082016@133. Known vial lot code: 05202016@4.

Product Description:
Forskolin/PAPAV/PHEN/PGE1, 100 mcg/30 mcg/3 mcg/60 mcg/mL, 5 mL vial, Injectable, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
22 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0357-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 05272016@18.

Product Description:
Glutamine/ORNI/Arginine/LYS/CITRU+LIDO 75 mg/75 mg/150 mg/50 mg/100 mg/10 mg/mL Injectable, 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
147 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0358-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 06232016@14, 07212016@2, 08012016@35, 08192016@11.

Product Description:
Glutamine/Arginine/Carnitine 25 mg/100 mg/200 mg/mL Injectable, 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
137 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0359-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 07012016@10, 08032016@42, 08182016@11.

Product Description:
Glutamine/ORNI/ARGIN/LYSINE/LIDO Injectable, 75 mg/75 mg/150 mg/150 mg/10 mg/mL, 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
199 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0360-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 06292016@9, 07252016@39, 08102016@24.

Product Description:
Glutathione 200 mg/mL Injectable, 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
148 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0361-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 07252016@46, 08112016@19.

Product Description:
Indocyanine Green (P.F.), 15 mg Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
131 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0362-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 08232016@42. Known vial lot code: 08172016@45.

Product Description:
Inositol/Choline/B-Complex+Leucine+CARN+CHROM+LIDO (LipoLean), Injectable, 30 mL vial, Compounded by Wells Pharmacy. Ocala, FL

Product Quantity:
433 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0363-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04132016@32, 05062016@4, 05312016@5, 06132016@93, 06282016@36, 08172016@5.

Product Description:
Leucine/Isoleucine/Valine 10 mg/10 mg/5 mg/5 mg/mL Injectable, 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
189 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0364-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04062016@3, 04252016@4, 05172016@4, 06162016@6, 07212016@8.

Product Description:
Methylcobalamin 1,000 mcg/mL Injectable, a) 10 mL vial and b) 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 499 vials b) 1214 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0365-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 04062016@5, 05172016@1, 06162016@2, 07202016@47, 08222016@6 b) Known lot code: 04052016@1, 05032016@1, 05182016@1, 05262016@32, 06162016@13, 06232016@74, 07012016@18, 08092016@6, 09062016@33

Product Description:
Methylcobalamin 5,000 mcg/mL Injectable, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
265 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0366-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04152016@3, 05192016@1, 06132016@23, 07132016@10, 08112016@22, 09062016@32.

Product Description:
Methylcobalamin Injectable (Dr. Hall Kit) - 1,000 mcg/mL, 30 mL vial Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
7 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0367-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 04292016@93, 07282016@26, 09082016@37. Known vial lot code: 04052016@1, 05262016@32, 08092016@6.

Product Description:
MIC/B12(M) 15 mg/50 mg/100 mg/1 mg/mL Injectable, 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
633 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0368-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04222016@4, 05112016@24, 05162016@14, 06082016@8, 06222016@42, 07272016@8

Product Description:
MIC/B12/CAR/CHR/LYS/DMG 25 mg/50 mg/50 mg/0.5 mg/50 mg/25 mcg/25 mg/25 mg/mL Injectable, 30 mL vial, Injectable, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
106 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0369-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 06292016@10, 08092016@19, 08232016@5.

Product Description:
MIC/B-COMP/(M)/CHROM/CARN, 20 mg/40 mg/50 mg/25 mcg/25 mg/mL Injectable, 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
933 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0370-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04082016@4, 05102016@1, 05172016@6, 06032016@3, 06092016@17, 06212016@42, 07012016@32, 07272016@16, 08172016@13, 4252016@21.

Product Description:
MIC/BCOMP+CAR+ARG+CHR+LID 12.5 mg/25 mg/25 mg/25 mg/12.5 mg/0.2 mg/20 mg/mL Injectable, a) 10 mL and b) 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
511 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0371-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: a) 04192016@7, 05132016@7, 07052016@19, 08102016@9 and b) 04262016@4, 05132016@9, 05192016@9, 05252016@25, 08122016@1.

Product Description:
MIC/B-COMP+CARN+LIDO, 15 mg/30 mg/30 mg/85 mg/20 mg/mL Injectable, 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
1716 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0372-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 03302016@9, 04072016@15, 05022016@14, 05052016@5, 05162016@16, 06012016@11, 06092016@30, 06152016@15, 06212016@2, 07152016@8, 08042016@21.

Product Description:
MIC/B-COMP+LIDO 20 mg/40 mg/50 mg/10 mg/mL Injectable, 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
4 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0373-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 07132016@5.

Product Description:
MIC/B-COMPLEX 15 mg/50 mg/100 mg/mL Injectable, 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
984 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0374-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04052016@5, 04082016@3, 04182016@37, 05032016@2, 05182016@3, 05232016@22, 06072016@18, 06282016@45, 07252016@52, 08262016@8.

Product Description:
MIC/Chromium/B12(M) Injectable, 25 mg/50 mg/50 mg/25 mcg/1 mg/mL, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
648 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0375-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 03302016@8, 04152016@5, 05032016@15, 05262016@24, 06142016@1, 07202016@55.

Product Description:
MIC+B12(M)+Carnitine 12.5 mg/50 mg/100 mg/1 mg/50 mg/mL Injectable, 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
159

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0376-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 03252016@5, 04192016@8, 05242016@15, 06222016@28, 08102016@2.

Product Description:
Mitomycin Solution Injectable, Sterile 0.02%, 200 mcg/mL, 1 mL syringe, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
122 syringes

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0377-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04142016@107, 05242016@128

Product Description:
Mitomycin Solution Injectable, Sterile 0.04%, 400 mcg/mL, 1 mL syringe, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
41 syringes

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0378-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 05022016@160

Product Description:
Mitomycin 20 mg Injectable, Lyophilized (Buffered), 1 mL vial, Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
315 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0379-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 05122016@84, 07012016@107. Known vial lot code: 04212016@149, 06082016@7.

Product Description:
Mitomycin 40 mg Injectable, Lyophilized (Buffered), 1 mL vial, Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
2,336 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0380-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 04112016@201, 04202016@33, 05182016@46, 05232016@227, 06072016@55, 06092016@88, 06142016@60, 07012016@112, 07132016@142, 07292016@121, 07292016@129. Known vial lot code: 04062016@106, 04182016@10, 04262016@56, 05022016@189, 05102016@54, 05192016@84, 05242016@135, 06232016@70, 07012016@112, 07072016@94.

Product Description:
NADH/Carnitine/Taurine/B12(M) Injectable, 100 mg/150 mg/150 mg/10 mg, Lyophilized Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
219 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0381-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 04072016@29, 08122016@103. Known vial lot code: 03302016@13, 03252016@16, 08082016@58, 08082016@67.

Product Description:
Nandrolone Decanote (H), 200 mg/mL Injectable, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
526 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0382-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04112016@16, 04282016@3, 05202016@6, 06132016@4, 07052016@1, 07272016@7.

Product Description:
QUAD1(PAP/PHEN/ALPROS/ATRO) 0.9 mg/0.2 mg/20 mcg/0.01 mg/mL Injectable, a) 2 mL and b) 5 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 1,165 vials b) 2,217 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0383-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 04122016@17, 04142016@14, 05112016@23, 06072016@17, 06232016@13, 07122016@41, 08052016@12 b) Known lot code: 03242016@8, 04142016@15, 05032016@11, 05262016@12, 06242016@11, 07122016@42, 08052016@11.

Product Description:
QUAD2 (PAP/PHEN/ALPROS/ATRO) 9 mg/1 mg/10 mg/0.1 mg/mL Injectable a) 2 mL and b) 5 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 191 vials and b) 794 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0384-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 04222016@8, 06142016@42, 08152016@17 b) Known lot code: 03312016@8, 05182016@15, 08092016@18

Product Description:
QUAD4 (PAP/PHEN/ALPROS/ATRO) 30 mg/3 mg/60 mcg/0.2 mg/mL Injectable, a) 2 mL and b) 5 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
114 vials and b) 1,017 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0385-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 04072016@8, 06272016@22, 09082016@29 b) Known lot code: 04142016@16, 04272016@8, 06102016@13, 07062016@29, 08052016@1.

Product Description:
Sermorelin Acetate 15 mg Injectable Kit., Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
568 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0386-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 05182016@173, 07052016@148, 09062016@114. Known vial lot code: 04252016@165, 06102016@56, 08262016@42.

Product Description:
Sermorelin Acetate 28 mg Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
330 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0387-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 05182016@51, 05262016@146, 06212016@76, 08052016@74. Known vial lot code: 04252016@163, 05042016@87, 07192016@31.

Product Description:
Sermorelin Acetate 9 mg Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
629 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0388-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 04212016@55, 06222016@106. Known vial lot code: 04142016@31, 06012016@50.

Product Description:
Sermorelin Acetate/GHRP (2) & (6) 9 mg/3.15 mg/3.15 mg Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
3,089 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0389-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 05122016@180, 06132016@155, 06232016@147, 07062016@156, 07272016@68, 08232016@24, 09062016@74. Kn own vial lot code: 04212016@110, 052412016@76, 05312016@40, 06102016@52, 07152016@56, 08192016@20, 08102016@56.

Product Description:
Sermorelin Acetate/GHRP (2) & (6) 9 mg/9 mg/9 mg Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
1,319 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0390-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 04072016@19, 05272016@55, 06102016@62, 09062016@75. Known vial lot code: 03302016@11, 050412016@98, 05 172016@10, 08252016@18.

Product Description:
Sermorelin Acetate/GHRP (2) 9 mg/9 mg Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
158 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0391-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 05182016@215, 09062016@92. Known vial lot code: 04252016@158, 082612016@45.

Product Description:
Sermorelin Acetate/GHRP (2) Kit 6 mg/4.5 mg Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
176 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0392-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 04062016@170, 05272016@67, 09132016@98. Known vial lot code: 03302016@12, 050412016@84, 08222016@27.

Product Description:
Sermorelin Acetate/GHRP (2)/Theanine 15 mg/5.4 mg/75 mg Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
1,255 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0393-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 05162016@37, 06172016@161, 06242016@135, 08112016@121. Known vial lot code: 04202016@8, 05252016@33, 0 5312016@47, 08042016@39.

Product Description:
Sermorelin Acetate/GHRP (2) 3 mg/4.5 mg Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
3 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0394-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 05272016@100. Known vial lot code: 05042016@94.

Product Description:
Sermorelin Acetate/GHRP (2) 9 mg/3 mg Injectable Kit, Compounded by Wells Pharmacy, Ocala FL

Product Quantity:
2,264 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0395-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 04152016@149, 05182016@176, 06172016@142, 07272016@83, 08232016@32, 09122016@138. Known vial lot code: 04072016@91, 04262016@27, 05272016@41, 07192016@29, 08192016@18, 09062016@44.

Product Description:
Sermorelin Acetate/GHRP (6) 3 mg/3 mg Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
106 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0396-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 05272016@73. Known vial lot code: 05042016@91.

Product Description:
Sermorelin Acetate/GHRP (6) 6 mg/3 mg Injectable Kit, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
1,232 kits (number of vials per kit vary by prescription)

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0397-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known kit lot code: 04192016@24, 05172016@229, 06032016@136, 06222016@121, 06222016@148, 09132016@102. Known vial lot code: 03242016@27, 04202016@7, 05162016@141, 06032016@47, 05312016@53, 08222016@54.

Product Description:
Sodium Bicarbonate, 8.4% Injectable, 30 mL MultiDose vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
67 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0398-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04062016@4, 05232016@120, 06142016@9, 09122016@9.

Product Description:
Testosterone Cypionate / Zinc Sulf in Grapeseed Oil Injectable, 200 mg/200 mcg/mL, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
266 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0399-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04012016@2, 05052016@4, 06022016@13, 06102016@35, 07052016@6.

Product Description:
Testosterone Cypionate in Grapeseed Oil Injectable, 100 mg/mL, a) 1mL and b) 5 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 303 vials and b) 413 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0400-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 04212016@5, 05102016@123, 05232016@21, 07132016@18 b) Known lot code: 04142016@7, 05042016@6, 05252016@2, 06222016@30, 07132016@25, 07282016@5.

Product Description:
Testosterone Cypionate in Grapeseed Oil Injectable, 20 mg/mL, 5 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
146 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0401-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04142016@5, 04292016@5, 05022016@6, 07062016@7.

Product Description:
Testosterone Cypionate in Sesame Oil Injectable 200 mg/mL, a) 1 mL and b) 10 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 798 vials and b) 3,191 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0402-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 03202016@10, 04072016@85, 04182016@4, 05062016@5, 05262016@1, 06132016@1, 08112016@4 b) Known lot code: 03282016@6, 04122016@2, 04182016@33, 05092016@6, 05192016@3, 06022016@5, 06072016@3, 06172016@3, 06292016@2, 07112016@5, 07262016@29, 07292016@12.

Product Description:
Testosterone Cypionate/Anastrozole Injectable, 200 mg/1 mg/mL, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
203 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0403-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 03242016@4, 04122016@1, 05102016@10, 06092016@4, 07062016@12, 08152016@3.

Product Description:
Testosterone Propionate Injectable, 100 mg/mL, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
469 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0404-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04042016@2, 04282016@1, 05262016@2, 06272016@8, 08162016@12.

Product Description:
Titan Up (INOS/CHOL/LEU/CAR/CHRO/LID) Injectable 25 mg/25 mg/1.5 mg/25 mg/0.025 mg/10 mg/mL, 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
433 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0405-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 03252016@4, 04012016@9, 04042016@9, 04132016@14, 05062016@3, 06022016@7, 06152016@22, 06292016@25, 07282016@12.

Product Description:
TitanMax-MIC/12/CA/CH/L/D Injectable 25 mg/50 mg/50/0.5 mg/50 mg/25 mcg/25 mg/25 mg/mL, 30 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
18 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0406-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 07212016@15.

Product Description:
Tri 1 (PAP/PHEN/ATROPINE) NR Injectable, 3.6 mg/0.4 mg/0.04 mg/mL, a) 2 mL and b) 5 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 257 vials and b) 460 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0407-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 04072016@10, 05102016@18, 06232016@12, 08182016@26 b) Known lot code: 04122016@19, 05042016@18, 06142016@43, 07122016@40, 07262016@47.

Product Description:
Tri 2 (PAP/PHEN/ATROPINE) NR Injectable, 30 mg/3 mg/0.3 mg/mL, a) 2 mL and b) 5 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 238 vials and b) 607 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0408-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 05042016@19, 06142016@28, 07262016@34, 09072016@22 b) Known lot code: 04112016@8, 05232016@2, 06212016@21, 07012016@22, 09082016@31.

Product Description:
Trimix with Atropine Injectable 30 mg/3 mg/150 mcg/0.2 mg/mL, a) 2 mL b) 5 mL and 10 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 6 vials, b) 19 vials and c) 41 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0409-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 04042016@22 b) Known lot code: 07192016@3 c) Known lot code: 04112016@13.

Product Description:
Trimix with Atropine Injectable 30 mg/3 mg/300 mcg/0.2 mg/mL, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
3 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0410-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 07192016@8.

Product Description:
Trimix with Atropine Injectable 30 mg/3 mg/60 mcg/0.2 mg/mL, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
1 vial

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0411-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04112016@6.

Product Description:
Trimix Injectable 30 mg/2 mg/40 mcg, 5 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
42 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0412-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code:03292016@79, 05262016@15, 07082016@9, 09082016@25.

Product Description:
Trimix Injectable 30 mg/2 mg/20 mcg, a) 5 mL and b) 10 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 56 vials and b) 48 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0413-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code:05182016@17, 07222016@16, 08182016@27 b) Known lot code:03312016@7, 05192016@23, 08032016@34.

Product Description:
Trimix Injectable 17.65 mg/0.59 mg/5.9 mcg/mL a) 2 mL, b) 5 mL and c) 10 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 2 vials b) 3 vials and c) 7 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0414-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 07192016@25 b) Known lot code 06202016@157 c) Known lot code: 05042016@16.

Product Description:
Trimix Injectable 30 mg/1 mg/10 mcg/mL a) 2 mL and b) 5 mL and c) 10 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 9 vials and b) 78 vials, c) 210 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0415-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 07222016@15 b) Known lot code: 04122016@17, 05182016@16, 09082016@28 c) Known lot code: 04072016@13, 05192016@22, 06132016@36, 08032016@35.

Product Description:
Trimix Injectable 30 mg/1 mg/25 mcg/mL, 5 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
7 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0416-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 07192016@26.

Product Description:
Trimix Injectable 30 mg/3 mg/100 mcg/mL, a) 5 mL and 10 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 1 vial and b) 12 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0417-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 07192016@7 b) Known lot code: 04122016@18.

Product Description:
Trimix Injectable 30 mg/3 mg/60 mcg/mL, a) 2 mL, b) 5 mL and c) 10 mL vials, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
a) 2 vials,, b) 9 vials and 6 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0418-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. a) Known lot code: 07192016@27 b) Knonw lot code: 06202016@162 c) Known lot code: 07262016@49.

Product Description:
Tri-Test 200 (CEP 100-75-25) Injectable, 200 mg/mL, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
550 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0419-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04122016@4, 05042016@3, 06012016@8, 06212016@5, 07252016@22.

Product Description:
Vitamin D3 in Sesame Oil Injectable 100,000 IU/mL, 5 mL, vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
237 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0420-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04112016@23, 05132016@3, 06062016@7, 06272016@17.

Product Description:
Vitamin D3 in Sesame Oil Injectable 50,000 IU/mL, 5 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
335 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0421-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04202016@1, 05052016@6, 06062016@1, 08012016@34.

Product Description:
Testosterone Cypionate in Grapeseed Oil Injectable, (NY STATE), 200 mg/mL, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
4 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0422-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04082016@6, 06012016@6, 06142016@148

Product Description:
Testosterone Cypionate in Sesame Oil Injectable, (NY STATE), 200 mg/mL, 10 mL vial, Compounded by Wells Pharmacy, Ocala, FL

Product Quantity:
3 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0423-2017

Code Information:
All lot codes of product prepared between 02/22/2016 and 09/14/2016, within expiry. Known lot code: 04182016@33, 06022016@5

Class II Drugs Event

Event ID:
75763

Status:
Ongoing

Recall Initiation Date:
11/17/2016

Center Classification Date:
01/04/2017

Recalling Firm:
Tri-Coast Pharmacy
14125 US Highway 1
Juno Beach FL United States

Distribution Pattern:
Nationwide

Product Type:
Drugs

Date Terminated:

Voluntary / Mandated:
Voluntary: Firm Initiated

Initial Firm Notification of Consignee or Public:
Two or more of the following: Email, Fax, Letter, Press Release, Telephone, Visit

Associated Products

Product Description:
Amino Energy Cocktail (30 ML), (Methionine (L) 12.5 mg/Inositol 25 mg/Choline Chloride 25 mg/Methylcobalamin 300 mcg/Carnitine (L) 125 mg/Thiamine HCL 50 mg/Riboflavin-5-Phosphate Sod 5 mg/Niacinamide 20 mg/Pyridoxine HCL 2 mg/Folic Acid 2 mg/Ascorbic Acid 50 mg/Chromium Picolinate 200 mcg/Glutamine (L) 30 mg/Lidocaine HCL 10 mg/mL). Compounded by Tri-Coast Pharmacy.

Product Quantity:
59 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0246-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 06012016A (11/28/2016), 07282016A (12/31/2016).

Product Description:
Amino Acid Recover Eaze (50 ML), Phenylaline (L) 8 mg/ml + Tyrosine (L) 0.45 mg/ml + L-Taurine 0.7 mg/ml. Compounded by Tri-Coast Pharmacy.

Product Quantity:
105 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0247-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 06072016C (11/15/2016), 08162016E (09/08/2016).

Product Description:
Ascorbic Acid 500 mg/mL (50 ML). Compounded by Tri-Coast Pharmacy.

Product Quantity:
201 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0248-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 07192016WW (01/15/2017), 07152016G (01/11/2017), 08292016A (02/25/2017), 10262016B (02/28/2017).

Product Description:
BCAA (30 ML) (Isoleucine (L) 15 mg + Leucine (L) 10 mg+ Valine 40 mg+ Lidocaine HCL 10 mg/mL). Compounded by Tri-Coast Pharmacy.

Product Quantity:
265 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0249-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 07222016A (01/18/2017), 08162016C (02/12/2017), 08042016A (01/31/2017), 11012016I (04/30/2017), 1017 2016B (02/28/2017).

Product Description:
B-Complex (30 ML), (Thiamine HCL 100 mg/Riboflavin-5-Phosphate 2 mg/Pyridoxine HCL 2 mg/Niacinamide 100 mg/Methylcobalamin 300 mcg/ml). Compounded by Tri-Coast Pharmacy.

Product Quantity:
67 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0250-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 07262016E (12/31/2016), 10262016E (02/28/2017).

Product Description:
BiMix Phentolamine/Papaverine 0.06 mg/1.8 mg/mL (1 ML). Compounded by Tri-Coast Pharmacy.

Product Quantity:
28 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0251-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 06092016B (0724/2016), 07212016C (09/04/2016), 09162016A (10/31/2016), 11012016A (12/16/2016).

Product Description:
BiMix Phentolamine/Papaverine 1 mg/30 mg/mL (1 ML). Compounded by Tri-Coast Pharmacy

Product Quantity:
5 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0252-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 07222016H (09/05/2016)

Product Description:
BiMix Phentolamine/Papaverine 2 mg/30 mg/mL (1 ML). Compounded by Tri-Coast Pharmacy

Product Quantity:
2 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0253-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 09302016D (11/14/2016).

Product Description:
Calcium Gluconate 10% Injection (30 ML). Compounded by Tri-Coast Pharmacy

Product Quantity:
12 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0254-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 08172016A (02/13/2017).

Product Description:
Calm Me (Aminobutyric Acid (GABA) 50 mg/Magnesium Chloride 50 mg/Taurine 50 mg/Theanine (L) 50 mg/Tryptophan (L) 10 mg/Lidocaine HCL 10 mg/ml), (30 ML). Compounded by Tri-Coast Pharmacy.

Product Quantity:
10 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0255-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 09212016A (01/31/2017), 08032016C (01/31/2017).

Product Description:
Dexpanthenol 250 mg/mL (30 ML). Compounded by Tri-Coast Pharmacy

Product Quantity:
5 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0256-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 08162016B (02/12/2017).

Product Description:
GAC, L-Glutamine/L-Arginine/L-Carnitine 25 mg/100 mg/200 mg/mL (30 mL). Tri-Coast Pharmacy.

Product Quantity:
114 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0257-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 06172016A (12/14/2016), 08092016B (02/05/2014), 09262016A (03/25/2017), 10132016E (02/28/2017).

Product Description:
Glucosamine 200 mg/mL (30 ML). Compounded by Tri-Coast Pharmacy

Product Quantity:
30 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0258-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 10192016D (10/22/2016).

Product Description:
Glutamine 25 mg/mL (30 ML). Compounded by Tri-Coast Pharmacy

Product Quantity:
12 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0259-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp date): 08032016A (01/30/2017).

Product Description:
Glutathione 200 mg/mL (30 ML). Compounded by Tri-Coast Pharmacy

Product Quantity:
367 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0260-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 06082016A (09/06/2016), 07052016C (10/03/2016), 08082016A (11/06/2016), 09132016D (12/12/2016), 1026 2016A (01/24/2016).

Product Description:
HCG 5,000 IU Vial. Compounded by Tri-Coast Pharmacy.

Product Quantity:
4,884 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0261-2017

Code Information:
Lot codes: 06222016D, 06062016C, 07192016C07062016A, 08182016B, 08112016A, 08012016A, 09282016A, 09122016D, 10182016C, 11042016B, 11012016F; Exp. 11/16

Product Description:
HCG/Hydroxycobalamin 5,000 IU/5 mg Vial. Compounded by Tri-Coast Pharmacy

Product Quantity:
1,304 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0262-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 06232016A (12/20/2016), 07192016B (01/15/2017), 08262016A (01/31/2017), 08092016C (01/31/2017), 0919 2016C (01/31/2017), 11042016C (11/04/2016), 10252016A (02/28/2017)

Product Description:
HCG 11,000 IU Vial. Compounded by Tri-Coast Pharmacy

Product Quantity:
5,229 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0263-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 06152016A (12/12/2016), 07252016D (01/21/2017), 07112016C (01/07/2017), 08312016B (02/27/2017), 0826 2016B (02/22/2017), 08092016D (02/05/2017), 08052016A (02/01/2017), 09282016B (02/28/2017), 09222016D (02/28/2017), 09072016A (02/28/2017), 10252016J (02/28/2017), 10182016D (02/28/2017), 10102016H (02/2 8/2017), 11042016A (02/28/2017), 11012016E (02/28/2017)

Product Description:
Infuvite IV (Ascorbic Acid 200 mg+ Vitamin A Acetate 330 IU+Vitamin D3 200 IU+Thiamine HCL 6 mg+Riboflavin-5-Phosphate Sod 3.6 mg+Pyridoxine HCL 6 mg+Niacinamide 40 mg+Dexpanthenol 15 mg+Vitamin 40E Succinate 10 IU+Phytonadione (K1) 150 mcg/5 mL). 30 ML. Compounded by Tri-Coast Pharmacy.

Product Quantity:
161 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0264-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date, if known): 07062016D (10/30/2016), 08312016A, 08232016C, 08222016C, 08102016E, 08102016D, 08012016 E, 09262016C, 09132016C

Product Description:
IV Cocktail #2 (Ascorbic Acid 200 mg+Vitamin A Acetate 330 IU+Vitamin D3 200 IU+Thiamine HCL 6 mg+Riboflavin-5-Phosphate Sod 3.6 mg+Pyridoxine HCL 6 mg+Niacinamide 40 mg+Dexpanthenol 15 mg+Vitamin 40E Succinate 10 IU+Phytonadione (K1) 150 mcg/5 mL), (30 ML). Compounded by Tri-Coast Pharmacy

Product Quantity:
132 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0265-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date, if known): 05252016D (05/28/2016), 05202016A (05/23/2016), 05182016C (05/21/2016), 05092016C (05/12/2 016), 05022016C (05/05/2016), 06132016D (06/16/2016), 06072016A (06/10/2016), 06062016A (06/09/2016), 07282106E (07/31/2016), 07252016G (07/28/2016), 09122016A (09/15/2016), 09192016B (09/22/2016)

Product Description:
L-Arginine HCL 100 mg/mL, (30 ML). Compounded by Tri-Coast Pharmacy

Product Quantity:
41 vials

Reason for Recall:
Lack of Assurance of Sterility

Recall Number:
D-0266-2017

Code Information:
Recall includes all sterile products compounded between 05/17/2016 and 11/17/2016. Known lot codes (Exp Date): 07282016H (01/24/2017), 09282016E (03/27/2017)

Product Description:
L-Carnitine 250 mg/mL, (30 ML). Compounded by Tri-Coast Pharmacy

Product Quantity:
14 vials