

Enforcement Report - Week of September 23, 2020

Class I Drugs Event

Event ID:
86105

Status:
Ongoing

Recall Initiation Date:
07/23/2020

Center Classification Date:
09/11/2020

Recalling Firm:

Real Clean Distribuciones SA de CV
Recursos Petroleros Fracc Industrial La Loma 3; Tlalnepantla De Baz
Tlalnepantla De Baz Mexico

Distribution Pattern:
Distributed Nationwide in the USA

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:

BORN BASIC., ANTI-BAC HAND SANITIZER, 70% Alcohol, (Ethyl Alcohol 70%), 16.9 FL OZ (500 mL) bottle, UPC 8 40038 21456 3, Made in Mexico; Distributed by Scent Theory Products, LLC, New York, NY 10018.

Product Quantity:

Reason for Recall:

Chemical Contamination: Born Basic Anti Bac was found to be below the label claim for ethanol content and contained methanol. Other products were recalled because they were manufactured in the same facility as the product found to contain methanol.

Recall Number:

D-1583-2020

Code Information:

Lots: 1033420, 1133420, 1233420, 2233420, 2333420, 2433420, 2533420

Class I Drugs Event

Event ID:
86127

Status:
Ongoing

Recall Initiation Date:
07/27/2020

Center Classification Date:
09/11/2020

Recalling Firm:

Broncolin, S.A. de C.V.
Calle Sur 16 No. 353 Col. Agricola Oriental, Iztacalco
Ciudad De Mexico Mexico

Distribution Pattern:
Distributed Nationwide in the USA

Product Type:
Drugs

Date Terminated:

Voluntary / Mandated:
Voluntary: Firm initiated

Initial Firm Notification of Consignee or Public:
Press Release

Associated Products

Product Description:

HERBACIL, Antiseptic Hand Sanitizer, 70% Alcohol, (Alcohol) 70%, packaged as a) 4.22 fl. oz. (125ml) bottle, UPC 7 14706 91368 6; b) 8.4 fl. oz. (250 ml) bottle, UPC 7 14706 91367 9; c) 33.8 fl. oz. (1 Liter) bottle, UPC 7 14706 91365 5; , Made in Mexico by: Broncolin, S.A. de C.V. Sur

16 No. 353, Col. Agricola Oriental, Iztacalco, C.P. 08500, Ciudad de Mexico, Mexico. Importer/Distributor: INBC TRADING LLC 4404 Merle Drive, Austin, TX 78745.

Product Quantity:

26,972 bottle

Reason for Recall:

Chemical Contamination: HERBACIL, Antiseptic Hand Sanitizer, 70% Alcohol, was found to be below the label claim for ethanol content and contained methanol.

Recall Number:

D-1578-2020

Code Information:

Lot# L 201117, Exp 05/2022

Class I Drugs Event

Event ID:

86268

Status:

Ongoing

Recall Initiation Date:

08/17/2020

Center Classification Date:

09/13/2020

Recalling Firm:

Maison Terre, LLC
226 Melrose Cir
North Little Rock AR United States

Distribution Pattern:

Nationwide.

Product Type:

Drugs

Date Terminated:**Voluntary / Mandated:**

Voluntary: Firm initiated

Initial Firm Notification of Consignee or Public:

Press Release

Associated Products

Product Description:

Goldenseal Root Powder Organic, 1 oz. clear plastic bag, Maison Terre Natural Products, www.maisonterre.net

Product Quantity:

670 bags

Reason for Recall:

Microbial Contamination of Non-Sterile Products: FDA laboratory analysis found product to be contaminated with various microorganisms including Enterobacter cloacae, Cronobacter sakazakii, Cronobacter dublinensis, among others.

Recall Number:

D-1584-2020

Code Information:

All lots distributed 01/25/2015 to 08/04/2020

Class II Drugs Event

Event ID:

86105

Status:

Ongoing

Recall Initiation Date:

07/23/2020

Center Classification Date:

09/11/2020

Recalling Firm:

Real Clean Distribuciones SA de CV

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

Recursos Petroleros Fracc Industrial La Loma 3; Tlalnepantla De Baz
Tlalnepantla De Baz Mexico

Distribution Pattern:

Distributed Nationwide in the USA

Associated Products**Product Description:**

scent theory KEEP IT CLEAN, pure clean,(Ethyl Alcohol 70%), 16.9 FL OZ/500 mL, Distributed by Scent Theory Products, LLC New York, NY 10018, Made in Mexico, UPC 8 40038 21434 1

Product Quantity:

504,000 pieces

Reason for Recall:

CGMP Deviations: Other products were recalled because they were manufactured in the same facility as the product found to contain methanol.

Recall Number:

D-1579-2020

Code Information:

Lots: 0133220, 0233220, 0333220, 0433220, 0533220, 0633220, 0733220, 0833220, 0933220, 1033220 1133220, 1233220, 1333220, 1433220, 1533220

Product Description:

scent theory KEEP CLEAN Moisturizing Hand Sanitizer, 70% alcohol (Ethyl Alcohol 70%), 16.9 FL OZ/500 mL bottle, Distributed by Scent Theory Products, LLC New York, NY 10018. Made in Mexico, UPC 8 40038 21434 1

Product Quantity:

917,280 bottles

Reason for Recall:

CGMP Deviations: Born Basic Anti Bac was found to be below the label claim for ethanol content and contained methanol. Other products were recalled because they were manufactured in the same facility as the product found to contain methanol.

Recall Number:

D-1580-2020

Code Information:

Lots: 1633220, 1733220, 1833220, 1933220, 2033220, 2133220, 2233220, 2333220, 2433220, 2533220, 2633220, 2733220, 2833220, 2933220, 3033220, 3133220, 3233220, 3333220, 3433220, 3533220, 3633220, 3733220, 3833220, 3933220, 4033220, 4133220

Product Description:

BORN BASIC., ANTI-BAC HAND SANITIZER, 70% Alcohol, (Ethyl Alcohol 70%), packaged as a) 9.5 FL OZ (280 mL) bottle, UPC 8 40038 21462 4; b) 16.9 FL OZ (500 mL) bottle, UPC 8 40038 21456 3; and c) 34 FL OZ (1L) bottle, UPC 8 40038 21447 1; Made in Mexico; Distributed by Scent Theory Products, LLC, New York, NY 10018.

Product Quantity:

975,720 bottles

Reason for Recall:

CGMP Deviations: Born Basic Anti Bac was found to be below the label claim for ethanol content and contained methanol. Other products were recalled because they were manufactured in the same facility as the product found to contain methanol.

Recall Number:

D-1581-2020

Code Information:

Lots: a) 0133720, 0233720 b) 0833420, 0933420, 1333420, 1433420, 1533420, 1633420, 1733420, 1833420, 1933420, 2033420, 2133420, 2633420, 2733420, 2833420, 2933420 c) 0133420, 0233420, 0333420, 0433420, 0533420, 0633420

Product Description:

LUX EOI Hand Sanitizing Gel, (Ethyl Alcohol 70%), (16.9 FL OZ) 500mL bottle, Distributed by: Procurement Services, LLC Nashville, TN 37215, Product of Mexico. NDC 74882-007-02

Product Quantity:

90,288 bottles

Reason for Recall:

CGMP Deviations: Born Basic Anti Bac was found to be below the label claim for ethanol content and contained methanol. Other products were recalled because they were manufactured in the same facility as the product found to contain methanol.

Recall Number:

D-1582-2020

Code Information:

Lots: 0133920, 0233920, 0333920

Class II Drugs Event**Event ID:**

86127

Status:

Ongoing

Recall Initiation Date:

07/27/2020

Center Classification Date:

09/11/2020

Recalling Firm:

Broncolin, S.A. de C.V.
Calle Sur 16 No. 353 Col. Agricola Oriental, Iztacalco
Ciudad De Mexico Mexico

Distribution Pattern:

Distributed Nationwide in the USA

Product Type:

Drugs

Date Terminated:**Voluntary / Mandated:**

Voluntary: Firm initiated

Initial Firm Notification of Consignee or Public:

Press Release

Associated Products**Product Description:**

HERBACIL, Antiseptic Hand Sanitizer, 70% Alcohol, (Alcohol) 70%, packaged as a) 4.22 fl. oz. (125ml) bottle, UPC 7 14706 91368 6; b) 8.4 fl. oz. (250 ml) bottle, UPC 714706 91367 9; c) 16.9 fl. oz. (500 ml) bottle, UPC 7 14706 91366 2; and d) 33.8 fl. oz. (1 Liter) bottle, UPC 7 14706 91365 5; Made in Mexico by: Broncolin, S.A. de C.V. Sur 16 No. 353, Col. Agricola Oriental, Iztacalco, C.P. 08500, Ciudad de Mexico, Mexico. Importer/Distributor: INBC TRADING LLC 4404 Merle Drive, Austin, TX 78745

Product Quantity:

482,416 bottles

Reason for Recall:

CGMP Deviations: products were recalled because they were manufactured in the same facility as the product found to contain methanol.

Recall Number:

D-1577-2020

Code Information:

Lots: 201111, 201113, 201115 Exp. 04/30/2022; 201116, 201118, 201119, 201120, 201121, Exp. 05/31/2022,

Class III Drugs Event**Event ID:**

86288

Status:

Ongoing

Recall Initiation Date:

05/05/2020

Center Classification Date:

09/14/2020

Recalling Firm:

Exela Pharma Sciences LLC
1245 Blowing Rock Blvd
Lenoir NC United States

Distribution Pattern:

Nationwide and Puerto Rico

Product Type:

Drugs

Date Terminated:**Voluntary / Mandated:**

Voluntary: Firm initiated

Initial Firm Notification of Consignee or Public:

Letter

Associated Products**Product Description:**

Potassium Acetate Injection, USP 40 mEq/20 mL (2 mEq/mL) Single Dose Vials, Rx only, Manufactured By: Exela Pharma Sciences, LLC. Lenoir, NC NDC 51754-2001-4

Product Quantity:

23,775 vials

Reason for Recall:

Short Fill

Recall Number:

D-1585-2020

Code Information:

Lot #P0000471, Catalog #2020TS002, Exp Date: 04/2021

Not Yet Classified Drugs Event

Event ID:

86343

Product Type:

Drugs

Status:

Ongoing

Date Terminated:**Recall Initiation Date:**

08/25/2020

Voluntary / Mandated:

Voluntary: Firm initiated

Center Classification Date:**Initial Firm Notification of Consignee or Public:**

E-Mail

Recalling Firm:

RLC Labs Inc.
1850 E Riverview Dr
Phoenix AZ United States

Distribution Pattern:

Nationwide in the United States

Associated Products

Product Description:

Nature-Throid, Thyroid USP [liothyronine (T3) 9 mcg and levothyroxine (T4) 38 mcg], 1 Grain (65 mg) Tablets, packaged in a) 30-count bottles, NDC 64727-3300-4; b) 60-count bottles, NDC 64727-3300-5; c) 90-count bottles, NDC 64727-3300-6; d) 100-count bottles, NDC 64727-3300-1; and e) 1,000-count bottles, NDC 64727-3300-2; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034.

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

Nature-Throid, Thyroid USP [liothyronine (T3) 6.75 mcg and levothyroxine (T4) 28.5 mcg], 3/4 Grain (48.75 mg) Tablets, packaged in a) 30-count bottles, NDC 64727-3302-4; b) 60-count bottles, NDC 64727-3302-5; c) 90-count bottles, NDC 64727-3302-6; d) 100-count bottles, NDC 64727-3302-1; and e) 1,000-count bottles, NDC 64727-3302-2; Rx Only, Manufactured by: RLC Labs Phoenix, AZ 85034.

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

Nature-Throid, Thyroid USP [liothyronine (T3) 4.5 mcg and levothyroxine (T4) 19 mcg], 1/2 Grain (32.5 mg) Tablets, packaged in a) 30-count bottles, NDC 64727-3299-4; b) 60-count bottles, NDC 64727-3299-5; c) 90-count bottles, NDC 64727-3299-6; d) 100-count bottles, NDC 64727-3299-1; and e) 1000-count bottles, NDC 64727-3299-2; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034.

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

Nature-Throid, Thyroid USP [liothyronine (T3) 11.25 mcg and levothyroxine (T4) 47.5 mcg], 1.25 Grain (81.25 mg) Tablets, packaged in a) 30-count bottles, NDC 64727-3303-4; b) 60-count bottles, NDC 64727-3303-5; c) 90-count bottles, NDC 64727-3303-6; d) 100-count bottles, NDC 64727-3303-1; e) 1,000-count bottles, NDC 64727-3303-2; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034.

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

Nature-Throid, Thyroid USP [liothyronine (T3) 13.5 mcg and levothyroxine (T4) 57 mcg], 1.5 Grain (97.5 mg) Tablets, packaged in a) 30-count bottles, NDC 64727-3305-4; b) 60-count bottles, NDC 64727-3305-5; c) 90-count bottles, NDC 64727-3305-6; d) 100-count bottles, NDC 64727-3305-1; and e) 1,000-count bottles, NDC 64727-3305-2; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034.

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

Nature-Throid, Thyroid USP [liothyronine (T3) 15.75 mcg and levothyroxine (T4) 66.5 mcg], 1.75 Grain (113.75 mg) Tablets, packaged in a) 30-count bottles, NDC 64727-3307-4; b) 60-count bottles, NDC 64727-3307-5; c) 90-count bottles, NDC 64727-3307-6; d) 100-count bottles, NDC 64727-3307-1; and e) 1,000-count bottles, NDC 64727-3307-2; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034.

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

Nature-Throid, Thyroid USP [liothyronine (T3) 18 mcg and levothyroxine (T4) 76 mcg], 2 Grain (130 mg) Tablets, packaged in a) 30-count bottles, NDC 64727-3308-4; b) 60-count bottles, NDC 64727-3308-5; c) 90-count bottles, NDC 64727-3308-6; d) 100-count bottles, NDC 64727-3308-1; and e) 1,000-count bottles, NDC 64727-3308-2; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034.

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

Nature-Throid, Thyroid USP [liothyronine (T3) 20.25 mcg and levothyroxine (T4) 85.5 mcg], 2.25 Grain (146.25 mg) Tablets, packaged in a) 30-count bottles, NDC 64727-3309-4; b) 60-count bottles, NDC 64727-3309-5; c) 90-count bottles, NDC 64727-3309-6; d) 100-count bottles, NDC 64727-3309-1; and e) 1,000-count bottles, NDC 64727-3309-2; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034.

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

Nature-Throid, Thyroid USP [liothyronine (T3) 22.5 mcg and levothyroxine (T4) 95 mcg], 2.5 Grain (162.5 mg) Tablets, packaged in a) 30-count bottles, NDC 64727-3310-4; b) 60-count bottles, NDC 64727-3310-5; c) 90-count bottles, NDC 64727-3310-6; d) 100-count bottles, NDC 64727-3310-1; and e) 1,000-count bottles, NDC 64727-3310-2; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034.

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

Nature-Throid, Thyroid USP [liothyronine (T3) 27 mcg and levothyroxine (T4) 114 mcg], 3 Grain (195 mg) Tablets, packaged in a) 30-count bottles, NDC 64727-3312-4; b) 60-count bottles, NDC 64727-3312-5; c) 90-count bottles, NDC 64727-3312-6; d) 100-count bottles, NDC 64727-3312-1; and e) 1,000-count bottles, NDC 64727-3312-2; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034.

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

WP Thyroid, Westhroid Pure, Thyroid USP, [liothyronine (T3) 2.25 mcg and levothyroxine (T4) 9.5 mcg], 1/4 Grain (16.25 mg) Tablets, packaged in a) 30-count bottles, NDC 64727-5450-4; b) 60-count bottles, NDC 64727-5450-5; c) 90-count bottles, NDC 64727-5450-6; d) 100-count bottles, NDC 64727-5450-1; and e) 1,000-count bottles, NDC 64727-5450-2; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034.

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

WP Thyroid, Westhroid Pure, 1/2 Grain (32.5 mg) Thyroid USP packaged in a) 1,000 Tablets, NDC 64727-5550-2 ; b) 30 Tablets, NDC 64727-5550-4; c) 60 Tablets, NDC 64727-5550-5; d) 90 Tablets, NDC 64727-5550-6; e) 100 Tablets, NDC 64727-5550-1; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

WP Thyroid, Westhroid Pure, 3/4 Grain (48.75 mg) Thyroid USP, packaged in a) 1,000 Tablets, NDC 64727-5650-2 ; b) 30 Tablets, NDC 64727-5650-4; c) 60 Tablets, NDC 64727-5650-5; d) 90 Tablets, NDC 64727-5650-6; e) 100 Tablets, NDC 64727-5650-1; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

WP Thyroid, Westhroid Pure, 1 Grain (65 mg) Thyroid USP, packaged in a) 1,000 Tablets, NDC 64727-5750-2 ; b) 30 Tablets, NDC 64727-

5750-4; c) 60 Tablets, NDC 64727-5750-5; d) 90 Tablets, NDC 64727-5750-6; e) 100 Tablets, NDC 64727-5750-1; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

WP Thyroid, Westhroid Pure, 1.25 Grain (81.25 mg) Thyroid USP packaged in a) 1,000 Tablets, NDC 64727-6050-2 ; b) 30 Tablets, NDC 64727-6050-4; c) 60 Tablets, NDC 64727-6050-5; d) 90 Tablets, NDC 64727-6050-6; e) 100 Tablets, NDC 64727-6050-1; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

WP Thyroid, Westhroid Pure, 1.5 Grain (97.5 mg) Thyroid USP packaged in a) 1,000 Tablets, NDC 64727-5850-2 ; b) 30 Tablets, NDC 64727-5850-4; c) 60 Tablets, NDC 64727-5850-5; d) 90 Tablets, NDC 64727-5850-6; e) 100 Tablets, NDC 64727-5850-1; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

WP Thyroid, Westhroid Pure, 1.75 Grain (113.75 mg) Thyroid USP, packaged in a) 1,000 Tablets, NDC 64727-6150-2 ; b) 30 Tablets, NDC 64727-6150-4; c) 60 Tablets, NDC 64727-6150-5; d) 90 Tablets, NDC 64727-6150-6; e) 100 Tablets, NDC 64727-6150-1; Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

WP Thyroid, Westhroid Pure, 2 Grain (130 mg) Thyroid USP packaged in a) 1,000 Tablets, NDC 64727-5950-2 ; b) 30 Tablets, NDC 64727-5950-4; c) 60 Tablets, NDC 64727-5950-5; d) 90 Tablets, NDC 64727-5950-6; e) 100 Tablets, NDC 64727-5950-1; Rx Only, Manufactured by: RLC Labs Phoenix, AZ 85034

Product Quantity:**Reason for Recall:**

CGMP Deviations: manufactured under the same conditions as those found to be subpotent.

Recall Number:**Code Information:**

All lots with expiry between 10/2020-07/2023

Product Description:

WP Thyroid, Westhroid Pure, 1/2 Grain (32.5 mg) Thyroid USP [Liothyronine (T3) 4.5 mcg, Levothyroxine (T4) 19 mcg], packaged in a) 30-count bottles (NDC 64727-5550-4), b) 90-count bottles (NDC 64727-5550-6), c) 100-count bottles (NDC 64727-5550-1), Rx only, Manufactured by: RLC Labs Phoenix, AZ 85024

Product Quantity:

Reason for Recall:

Subpotent Drug: FDA analysis found product to be subpotent

Recall Number:**Code Information:**

Lot #: 119142, Exp. Date 01/2022

Product Description:

Nature Throid, 1/2 Grain (32.5 mg) Thyroid USP [Liothyronine (T3) 4.5 mcg, Levothyroxine (T4) 19 mcg], packaged in a) 90-count bottles (NDC 64727-3299-6), b) 100-count bottles (NDC 64727-3299-1), Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034

Product Quantity:**Reason for Recall:**

Subpotent Drug: FDA analysis found product to be subpotent

Recall Number:**Code Information:**

Lot #: 099186, Exp. Date 11/2022

Product Description:

Nature Throid, 3/4 Grain (48.75 mg) Thyroid USP [Liothyronine (T3) 6.75 mcg, Levothyroxine (T4) 28.5 mcg], packaged in a) 100-count bottles (NDC 64727-3302-1) and b) 1000-count bottles (NDC 64727-3302-2), Rx only, Manufactured by: RLC Labs Phoenix, AZ 85034

Product Quantity:**Reason for Recall:**

Subpotent Drug: FDA analysis found product to be subpotent

Recall Number:**Code Information:**

Lot #: 109242, Exp. Date 11/2022

Product Description:

Nature Throid, 1.25 Grain (81.25 mg) Thyroid USP [Liothyronine (T3) 11.25 mcg, Levothyroxine (T4) 47.5 mcg], 100-count bottles, Rx only, Manufactured by: RLC Labs Phoenix, AZ 85024, NDC 64727-3303-1

Product Quantity:**Reason for Recall:**

Subpotent Drug: FDA analysis found product to be subpotent

Recall Number:**Code Information:**

Lot #: 109032, Exp. Date 10/2022

Product Description:

Nature Throid, 2 Grain (130 mg) Thyroid USP [Liothyronine (T3) 18 mcg, Levothyroxine (T4) 76 mcg], packaged in a) 30-count bottles (NDC 64727-3308-4) and b) 100-count bottles (NDC 64727-3308-1), Rx only, Manufactured by: RLC Labs Phoenix, AZ 85024

Product Quantity:**Reason for Recall:**

Subpotent Drug: FDA analysis found product to be subpotent

Recall Number:**Code Information:**

Lot #: 109264, Exp. Date 10/2022

Product Description:

Nature Throid, 1.5 Grain (97.5 mg) Thyroid USP [Liothyronine (T3) 13.5 mcg, Levothyroxine (T4) 57 mcg], packaged in a) 30-count bottles (NDC 64727-3305-4), b) 60-count bottles (NDC 64727-3305-5), c) 90-count bottles (NDC 64727-3305-6), and 100-count bottles (NDC 64727-3305-1), Rx only, Manufactured by: RLC Labs Phoenix, AZ 85024, NDC 64727-3305-1

Product Quantity:**Reason for Recall:**

Subpotent Drug: FDA analysis found product to be subpotent

Recall Number:**Code Information:**

Lot #: 109286, Exp. Date 10/2022

