A Public Entity

Inland Empire Health Plan

PHARMACY TIMES

BY IEHP PHARMACEUTICAL SERVICES DEPARTMENT

May 5, 2021

URGENT: Recalls, Market Withdrawals, & Safety Alerts

Dear IEHP Providers,

According to the FDA recall guidance, Title 21 Code of Federal Regulations Part 7 (21 CFR part 7), recalled products must be promptly removed or corrected. In an effort to promote health and wellness of our members, please review your records and notify members who may have been impacted by these recalls and market withdrawals.

Product Name	Product Code	Lot # and Exp. Date	Classification	Recalling Firm
Acetaminophen Extra Strength 500 mg Tablets	NDC: 50090-5350-00	323206, 323207, 323208, 323209, 323210, 323211, 323212, 323213, 323214, 323215, 323216, 323218, 323219, 323220, 323222, 323223, 323224, 323238, 335353, 335354, 335355, 335356, 335358, 335359, 335360, 335361, 335362, 335363, 335364, 335365, 335366, 33536, 335368, 335369, 335370, 335371, 335372, 335373, 335374, 335375, 335376, 335377, 335395, 352116 Exp07/31/2022 or 08/31/2022	MW	A-S Medication Solutions, LLM
Acyclovir Sodium Injection 1000mg/20mL (50mg/mL) Vial	NDC: 68382-0049-01; 68382-0049-10	L000155 Exp. 12/2021; L000156 Exp. 1/2022	Class I	Zydus Pharmaceuticals (USA) Inc

Acyclovir Sodium Injection 500mg/10mL (50mg/mL) Vial	NDC: 68382-0048-01; 68382-0048-10	L000126, L000127 Exp. 12/31/2021	Class I	Zydus Pharmaceuticals (USA) Inc
ZOMA-Jet 5 Demonstration Kit, Needle-free delivery device for use with Zomacton (somatropin) for injection 5mg vial	UPC: 3 55566 18031 5	201601320031 201702320207 201731220039 201817120044 201912720086 201912320044	Class II	Ferring Pharmaceuticals Inc
ZOMA-Jet 10 Demonstration Kit, Needle-free delivery device for use with Zomacton (somatropin) for injection 10 mg vial	UPC: 3 55566 19031 4	201827020015 201826020065 201835320058 201835320009 201834920003 201835220007 201901820009 201901420200 201902120010 201901120019 201901420005 201901020180 201901520015 201900720001 201900920004 201904620003 201902120011 201904420007 201904520015 201904920002 201905020111 201904620004 201905720011 201907320003 201907320030 201907820007 201907420017 201910820009 201915620014 201910920016 201910620017 201910820020 201911320048 201911320095 201913720148 201912920021	Class II	Ferring Pharmaceuticals Inc

Telmisartan 20 mg Tablets	NDC: 62332-0087-30	1905005661 Exp. 03/2022	Class I	Alembic Pharmaceuticals Limited
Neomycin Sulfate 500 mg Tablets	NDC: 39822-0310-05	CFMBX Exp. 9/2022	Class II	X-Gen Pharmaceuticals Inc.
Ganirelix Acetate 250 mcg/0.5 mL Injection	NDC: 55566-1000-01	JKU1212A, JKU1503A, JKU1504A, JKU1505A, JKU1506A Exp. 03/2021; JKU3313A & JKU3314A Exp. 08/2021	Class II	SUN PHARMACEUTICAL INDUSTRIES INC
Mometasone Furoate 0.1% Topical Solution (Lotion)	NDC: 00713-0701-85; 00713-0701-53	30 mL: 1014611. 1014612 Exp. 12/2022 60 mL: 1014593, 1014594, 1014595 Exp. 10/2022	Class II	Cosette Pharmaceuticals, Inc.
Guanfacine 2 mg Extended-Release Tablets	NDC: 60505-3928-01	RX1662, RX1663, RX1664 Exp. 11/2022	Class II	Apotex Corp.
Cefprozil 250mg/5mL for Oral Suspension, 50 mL (when mixed) bottle	NDC: 68180-0402-01; 68180-0402-02; 68180-0402-03	F801122, F801123, F801124 Exp. 06/2021	Class II	Lupin Pharmaceuticals Inc.
Riomet (metformin hydrochloride oral solution) 500 mg/5 mL Cherry Flavor, 16 fl. oz.	NDC: 10631-0206-02	J190386A, X190354A Exp. 03/2021; J190393A Exp. 05/2021; A200035A Exp. 06/2021; B200064A Exp. 08/2021; H200236A Exp. 01/2022	Class II	SUN PHARMACEUTICAL INDUSTRIES INC
Itraconazole 100 mg Capsules	NDC: 59746-0282-30	IT119008B Exp. 05/2021; IT120001A, IT120002A Exp. 12/2021	Class II	Jubilant Cadista Pharmaceuticals, Inc.
Betadine (Povidone-Iodine) 5%, 0.5mL per syringe Single Use Syringe Rx only, For Topical Ophthalmic Use (Do Not Inject) Sterile Ophthalmic Solution, Preservative Free	NDC Not Provided	02-2021-16@4 BUD 5/31/2021	Class II	Edge Pharma, LLC

Minivelle (estradiol transdermal system) Delivers 0.075 mg/day, 8 patches/box	NDC: 68968-6675-08	88584 Exp. 03/2022	Class II	Noven Pharmaceuticals Inc
Estradiol Transdermal System Delivers 0.0375 mg/day, 8 Systems/box	NDC: 68968-3437-08	88321 Exp. 02/2022	Class II	Noven Pharmaceuticals Inc
NP Thyroid 15 mg Tablets	NDC: 42192-0327-01; 42192-0327-07	M327L19-1 Exp. 04/30/2021; M327H19-3A Exp. 07/31/2021; M327D20-1, M327D20-3 Exp. 03/31/2022; M327D20-1 Exp. 03/31/2022	MW	Acella Pharmaceuticals, LLC
NP Thyroid 30 mg Tablets	NDC: 42192-0329-01; 42192-0329-07	M329D20-1, M329D20-2, M329D20-3 Exp. 03/31/2022; M329D20-2 Exp. 03/31/2022	MW	Acella Pharmaceuticals, LLC
NP Thyroid 60 mg Tablets	NDC: 42192-0330-01	M330J19-2A, M330J19-4A, M330J19-5A, M330J19-6A, M330J19-7A, M330J19-9A Exp. 08/31/2021; M330K19-10, M330K19-1A, M330K19-9 Exp. 09/30/2021; M330D20-1, M330D20-2 Exp. 03/31/2022;	MW	Acella Pharmaceuticals, LLC
NP Thyroid 90 mg Tablets	NDC: 42192-0331-01	M331J19-10A, M331J19-11, M331J19-2A, M331J19-6A Exp. 08/31/2021; M331K19-1, M331K19-2, M331K19-6 Exp. 09/30/2021	MW	Acella Pharmaceuticals, LLC

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NP Thyroid 120 mg Tablets	NDC: 42192-0328-01; 42192-0328-07	M328H19-2B, M328J19-11, M328J19-2A, M328J19-3A, M328J19-4A, M328J19-5A, M328J19-6A, M328J19-7A Exp. 08/31/2021; M328K19-2, M328K19-4A Exp. 9/30/2021; M328J19-9B Exp. 08/31/2021	MW	Acella Pharmaceuticals, LLC

I = Class I Recall, II = Class II Recall, MW = Market Withdrawal

Additional information can be found at:

1. FDA Recalls, Market Withdrawals, & Safety Alerts: https://www.fda.gov/Safety/Recalls/default.htm
2. FDA Enforcement Report: http://www.fda.gov/Safety/Recalls/EnforcementReports/default.htm
3. IEHP Safety Resources: https://ww3.iehp.org/en/providers/pharmaceutical-services/clinical-information/safety-resources/

If you have any questions or comments regarding this recall, please call IEHP Pharmaceutical Services Department at 909-890-2049, 8am – 5pm (PST), Monday through Friday.

Sincerely,

IEHP Pharmaceutical Services