

Capital Rx Drug Recall Report

JANUARY 2026



Welcome to the **Capital Rx Drug Recall Report**. This report is designed to keep you up to date on the latest FDA Class 1 and Class 2 recalled drugs and market withdrawals that impact our members. It is one of the many ways we, at Capital Rx, demonstrate our commitment to providing clients and partners the tools and resources they desire.

WHO WE ARE

Capital Rx is a full-service pharmacy benefit manager (PBM) and pharmacy benefit administrator (PBA), advancing our nation's electronic healthcare infrastructure to improve drug price visibility and patient outcomes. As a Certified B Corp™, Capital Rx is executing its mission through the deployment of JUDI®, the company's cloud-native enterprise health platform, and a Single-Ledger Model™, which increases visibility and reduces variability in drug prices. JUDI connects every aspect of the pharmacy ecosystem in one efficient, scalable platform, servicing over 2.4 million members for Medicare, Medicaid, and commercial plans. Together with our clients, we are reimagining the administration of pharmacy benefits and rebuilding trust in healthcare. ****The drug recall report is subject to change: information in this report is current as of 1/14/2026****

Privacy Statement:

This privacy policy describes the types of information we may collect from you or that you may provide when you visit the website cap-rx.com and our practices for collecting, using, maintaining, protecting, and disclosing that information. Capital Rx, Inc. ("we," "our," or "us") is committed to ensuring that your privacy is protected. This policy applies to information we may collect through cap-rx.com, including any services offered on or through cap-rx.com such as the prescription benefit member web portal, and our mobile application accessible at the Google Play Store and iOS App Store under the name Capital Rx (collectively, our "Site").

RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
12/24/2025	Class 2	Ciclopirox 1 % Sham	Sun Pharma / Taro Pharmaceuticals	51672-1351-08	AD37059, AD37060, AD37061, AD37062, AD37065, AD37066, AD37067, AD37068, EXP 1/31/2026	Failing to meet impurity and degradation standards.
01/07/2026	Class 2	Ondansetron 4 Mg Tbdp	Glenmark Pharmaceuticals Inc., USA	68462-0157-13	19251311, EXP APRIL 2027	A defective container. Preferred Pharmaceuticals received a letter from the manufacturer Glenmark, that the blister packs are not fully sealed and tablets falling out. Preferred Pharmaceuticals purchased the finished product and repackaged the product for sale.

RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
01/07/2026	Class 2	Wegovy 0.5 Mg/0.5ml Soaj	Novo Nordisk Inc.	00169-4505-14	RZFHD52, RZFW93; EXP 10/31/2026	Hair found in a prefilled syringe.
01/07/2026	Class 2	Wegovy 1 Mg/0.5ml Soaj	Novo Nordisk Inc.	00169-4501-14	RZFYK06, RZFYA53; EXP 3/31/2027	Hair found in a prefilled syringe.
01/14/2026	Class 2	Oxycodone-Acetaminophen 10-325 Mg Tabs	SpecGx, LLC	00406-0523-01	0523J23904, EXP 05/2027; 0523J24426, 0523J24427, EXP 06/2027	A possibility for the imprint to be missing on tablets.
01/14/2026	Class 2	Oxycodone-Acetaminophen 7.5-325 Mg Tabs	SpecGx, LLC	00406-0522-01	0522J23493, EXP 03/2027	A possibility for the imprint to be missing on tablets.

How do I find out more information about the recall? View the FDA website URL for more information.

RECALL TYPE	DRUG RECALLED	FDA NOTIFICATION URL
Class 2	Ciclopirox 1 % Sham	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=217622
Class 2	Ondansetron 4 Mg Tbdp	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=217948
Class 2	Wegovy 0.5 Mg/0.5ml Soaj	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=217653
Class 2	Wegovy 1 Mg/0.5ml Soaj	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=217889
Class 2	Oxycodone-Acetaminophen 10-325 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=217695
Class 2	Oxycodone-Acetaminophen 7.5-325 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=217751