Capital Rx Drug Recall Report

FEBRUARY 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 **2/18/2026****

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
1/21/2026	Class 1	Freestyle Libre 3 Plus Sensor Misc	Abbott Diabetes Care, Inc.	50090-7248-00, 57599-0844-00	Confirm impacted lot numbers by visiting www.FreeStyleCheck.com	The sensor potentially providing incorrect low glucose results
1/21/2026	Class 1	Freestyle Libre 3 Sensor Misc	Abbott Diabetes Care, Inc.	50090-6385-00, 57599-0818-00	Confirm impacted lot numbers by visiting www.FreeStyleCheck.com	The sensor potentially providing incorrect low glucose results

RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
1/21/2026	Class 2	Levothyroxine Sodium 150 Mcg Tabs	Alvogen, Inc	47781-0662-10	MHA21825, EXP 12/31/2027	A product mix up. A single bottle of this lot was reported by a pharmacy to contain 88 mcg tablets instead of 150 mcg tablets
1/28/2026	Class 2	Furosemide 40 Mg Tabs	Graviti Pharmaceuticals Private Limited	64980-0563-10	FUB125042G; EXP 05/13/2027	The presence of a foreign substance
1/28/2026	Class 2	Icosapent Ethyl 1 Gm Caps	Zydus Pharmaceuticals (USA) Inc	70710-1592-07	S2520304, S2520333, EXP 2/28/2027; S2540186, EXP 4/30/2027	The product being subpotent (not strong enough). This happened because some of the capsules leaked, which caused the medicine inside to break down. If someone uses this product, it might not work the way it should. It could also cause more stomach-related side effects for some people
2/4/2026	Class 2	Methylprednisolone 4 Mg Tbpk	Greenstone LLC	59762-4440-02	LG7675, EXP 11/2026	A labeling issue. The foil on the blister pack was put on the wrong way, so the dosing directions printed on it are not correct
2/11/2026	Class 2	Bisoprolol-HCTZ 2.5-6.25 Mg Tabs	Unichem Pharmaceuticals USA Inc.	29300-0187-01	GBHL24005A, EXP 09/2026	Deviations from the Current Good Manufacturing Practices (CGMP), not meeting the N-Nitroso Bisoprolol impurity specification limits
2/11/2026	Class 2	Trazodone Hcl 50 Mg Tabs	Granules Pharmaceuticals Inc.	70010-0231-01	BATCH # 6160008A, EXP 12/31/2026	The presence of tablets/capsules from another product found in Trazadone 50mg package

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 1	Freestyle Libre 3 Plus Sensor Misc	https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/abbott-initiates-medical-device-correction-certain-freestyle-librer-3-and-freestyle-libre-3-plus
Class 1	Freestyle Libre 3 Sensor Misc	https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/abbott-initiates-medical-device-correction-certain-freestyle-librer-3-and-freestyle-libre-3-plus
Class 2	Levothyroxine Sodium 150 Mcg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=217920
Class 2	Furosemide 40 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=218176
Class 2	Icosapent Ethyl 1 Gm Caps	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=217936
Class 2	Methylprednisolone 4 Mg Tbpk	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=218100
Class 2	Bisoprolol-Hydrochlorothiazide 2.5-6.25 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=218244
Class 2	Trazodone Hcl 50 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=218354