

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 5/20/2026****

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RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
4/22/2026	Class 2	Cinacalcet Hcl 30 Mg Tabs	Cipla USA, Inc.	69097-0410-02	EXP 4PB0526, 4PB0527, 4PB0528, EXP SEPTEMBER-2026; 5PB0173, EXP JANUARY-2027	Deviations from the Current Good Manufacturing Practices (CGMP), N-Nitroso-Cinacalcet being higher than the acceptable limit.
4/22/2026	Class 2	Cinacalcet Hcl 60 Mg Tabs	Cipla USA, Inc.	69097-0411-02	EXP 4PB0215, 4PB0216, EXP MARCH 2026; 4PB0515, 4PB0516, EXP SEPTEMBER-2026; 4PB0517, EXP SEPTEMBER-2026; 5PB0167, EXP JANUARY-2027	Deviations from the Current Good Manufacturing Practices (CGMP), N-Nitroso-Cinacalcet being higher than the acceptable limit.

RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
4/22/2026	Class 2	Cinacalcet Hcl 90 Mg Tabs	Cipla USA, Inc.	69097-0412-02	EXP 4PB0224, EXP MARCH-2026; 4PB0505, 4PB0506, 4PB0507, EXP SEPTEMBER-2026; 5PB0564, EXP APRIL-2027	Deviations from the Current Good Manufacturing Practices (CGMP), N-Nitroso-Cinacalcet being higher than the acceptable limit.
4/22/2026	Class 2	Fluocinonide 0.05 % Crea	Sun Pharmaceutical Industries Inc.	A) 51672-1386-01 B) 51672-1386-02 C) 51672-1386-03	A) AD76252, EXP 6/30/2026 B) AD76251, EXP 6/30/2026 C) AD76250, EXP 6/30/2026	During testing, the product was found to be thicker than expected and it did not meet quality standards.
4/22/2026	Class 1	Omnipod 5 Dexg7g6 Intro Gen 5 Kit	Insulet Corporation	85083-0000-01	MODEL/CATALOG NUMBER: PT-001662 ASM 5PK POD STRL OPS G6G7. UDI-DI: 10385083000527. CODES: PH1U02252541, PH1U03282511, PH1U03282522, PH1U03312511, PH1U03312521, PH1U04012511, PH1U04012521, PH1U05052511, PH1U08162531, PH1U08182531, PH1U08182541, PH1U09242511, PH1U09242521, PH1U09242531, PH1U09252521, PH1U09252531, PH1U10152541, PH1U10162531, PH1U10162541, PH1U10172531, PH1U10172541, PH1U10182531, PH1U10182541, PH1U10202511, PH1U10202521, PH1U10202531, PH1U10202541, PH1U10212531, PH1U10212541, PH1U10222531, PH1U10222541, PH1U10232531, PH1U10232541, PH1U10242521, PH1U10242531, PH1U10242541.	Pods potentially having a small tear in the internal tubing that delivers insulin. This may leak inside the Pod instead of being fully infused into the body as intended.

RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
4/22/2026	Class 1	Omnipod 5 Dexg7g6 Pods Gen 5 Misc	Insulet Corporation	85083-0000-21, 08508-3000-75	MODEL/CATALOG NUMBER: PT-001662 ASM 5PK POD STRL OPS G6G7. UDI-DI: 10385083000527. CODES: PH1U02252541, PH1U03282511, PH1U03282522, PH1U03312511, PH1U03312521, PH1U04012511, PH1U04012521, PH1U05052511, PH1U08162531, PH1U08182531, PH1U08182541, PH1U09242511, PH1U09242521, PH1U09242531, PH1U09252521, PH1U09252531, PH1U10152541, PH1U10162531, PH1U10162541, PH1U10172531, PH1U10172541, PH1U10182531, PH1U10182541, PH1U10202511, PH1U10202521, PH1U10202531, PH1U10202541, PH1U10212531, PH1U10212541, PH1U10222531, PH1U10222541, PH1U10232531, PH1U10232541, PH1U10242521, PH1U10242531, PH1U10242541.	Pods potentially having a small tear in the internal tubing that delivers insulin. This may leak inside the Pod instead of being fully infused into the body as intended.

RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
4/22/2026	Class 1	Omnipod 5 G7 Intro (Gen 5) Kit	Insulet Corporation	08508-3000-50	MODEL/CATALOG NUMBER: PT-001662 ASM 5PK POD STRL OPS G6G7. UDI-DI: 10385083000527. CODES: PH1U02252541, PH1U03282511, PH1U03282522, PH1U03312511, PH1U03312521, PH1U04012511, PH1U04012521, PH1U05052511, PH1U08162531, PH1U08182531, PH1U08182541, PH1U09242511, PH1U09242521, PH1U09242531, PH1U09252521, PH1U09252531, PH1U10152541, PH1U10162531, PH1U10162541, PH1U10172531, PH1U10172541, PH1U10182531, PH1U10182541, PH1U10202511, PH1U10202521, PH1U10202531, PH1U10202541, PH1U10212531, PH1U10212541, PH1U10222531, PH1U10222541, PH1U10232531, PH1U10232541, PH1U10242521, PH1U10242531, PH1U10242541.	Pods potentially having a small tear in the internal tubing that delivers insulin. This may leak inside the Pod instead of being fully infused into the body as intended.

RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
4/22/2026	Class 1	Omnipod 5 G7 Pods (Gen 5) Misc	Insulet Corporation	08508-3000-53	MODEL/CATALOG NUMBER: PT-001662 ASM 5PK POD STRL OPS G6G7. UDI-DI: 10385083000527. CODES: PH1U02252541, PH1U03282511, PH1U03282522, PH1U03312511, PH1U03312521, PH1U04012511, PH1U04012521, PH1U05052511, PH1U08162531, PH1U08182531, PH1U08182541, PH1U09242511, PH1U09242521, PH1U09242531, PH1U09252521, PH1U09252531, PH1U10152541, PH1U10162531, PH1U10162541, PH1U10172531, PH1U10172541, PH1U10182531, PH1U10182541, PH1U10202511, PH1U10202521, PH1U10202531, PH1U10202541, PH1U10212531, PH1U10212541, PH1U10222531, PH1U10222541, PH1U10232531, PH1U10232541, PH1U10242521, PH1U10242531, PH1U10242541.	Pods potentially having a small tear in the internal tubing that delivers insulin. This may leak inside the Pod instead of being fully infused into the body as intended.

RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
4/22/2026	Class 1	Omnipod 5 Libre2 Plus G6 Pods Misc	Insulet Corporation	08508-3000-42	MODEL/CATALOG NUMBER: PT-001662 ASM 5PK POD STRL OPS G6G7. UDI-DI: 10385083000527. CODES: PH1U02252541, PH1U03282511, PH1U03282522, PH1U03312511, PH1U03312521, PH1U04012511, PH1U04012521, PH1U05052511, PH1U08162531, PH1U08182531, PH1U08182541, PH1U09242511, PH1U09242521, PH1U09242531, PH1U09252521, PH1U09252531, PH1U10152541, PH1U10162531, PH1U10162541, PH1U10172531, PH1U10172541, PH1U10182531, PH1U10182541, PH1U10202511, PH1U10202521, PH1U10202531, PH1U10202541, PH1U10212531, PH1U10212541, PH1U10222531, PH1U10222541, PH1U10232531, PH1U10232541, PH1U10242521, PH1U10242531, PH1U10242541.	Inside the Pod instead of being fully infused into the body as intended.
4/29/2026	Class 2	Furosemide 80 Mg Tabs	Leading Pharma, LLC	69315-0118-01	H03125, H03225, H03325, EXP 2027/08	Deviations from the Current Good Manufacturing Practices (CGMP). Presence of N-nitroso-Furosemide (NNF) above the recommended limit.
4/29/2026	Class 2	Pantoprazole Sodium 40 Mg Tbec	Hetero Labs Limited (Unit V)	31722-0713-10	FD253967 EXP 06/26/2027	Discoloration. The firm received five (5) complaints stating that, "Tablets discolored darker than normal and have lighter-colored spots."

RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
5/5/2026	Class 1	Gnp True Metrix Glucose Meter W/Device Kit	Trividia Health, Inc.	56151-1494-01, 08528-1474-01, 87701-0427-39, 87701-0426-25, 11917-0173-89	All TRUE METRIX, TRUE METRIX AIR, TRUE METRIX GO, and TRUE METRIX PRO, branded Blood Glucose Meters distributed in the United States.	The system labeling (user manual and online labeling) did not provide adequate directions for users to act on the E-5 error code. The error code description and recommended actions for the E-5 error code in the labeling do not: (a) clearly emphasize that an E-5 error code could represent a very high blood glucose level, and (b) prominently convey the appropriate immediate actions, such as contacting a healthcare professional. This could lead to a delay in treatment if the user does not seek medical attention immediately when receiving an E-5 error code and experiencing symptoms of high glucose.
5/5/2026	Class 1	True Metrix Air Glucose Meter W/Device Kit	Trividia Health, Inc.	56151-1494-01, 08528-1474-01, 87701-0427-39, 87701-0426-25, 11917-0173-89	All TRUE METRIX, TRUE METRIX AIR, TRUE METRIX GO, and TRUE METRIX PRO, branded Blood Glucose Meters distributed in the United States.	The system labeling (user manual and online labeling) did not provide adequate directions for users to act on the E-5 error code. The error code description and recommended actions for the E-5 error code in the labeling do not: (a) clearly emphasize that an E-5 error code could represent a very high blood glucose level, and (b) prominently convey the appropriate immediate actions, such as contacting a healthcare professional. This could lead to a delay in treatment if the user does not seek medical attention immediately when receiving an E-5 error code and experiencing symptoms of high glucose.

RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
5/5/2026	Class 1	True Metrix Go Glucose Meter W/Device Kit	Trividia Health, Inc.	56151-1494-01, 08528-1474-01, 87701-0427-39, 87701-0426-25, 11917-0173-89	All TRUE METRIX, TRUE METRIX AIR, TRUE METRIX GO, and TRUE METRIX PRO, branded Blood Glucose Meters distributed in the United States.	The system labeling (user manual and online labeling) did not provide adequate directions for users to act on the E-5 error code. The error code description and recommended actions for the E-5 error code in the labeling do not: (a) clearly emphasize that an E-5 error code could represent a very high blood glucose level, and (b) prominently convey the appropriate immediate actions, such as contacting a healthcare professional. This could lead to a delay in treatment if the user does not seek medical attention immediately when receiving an E-5 error code and experiencing symptoms of high glucose.
5/5/2026	Class 1	True Metrix Meter W/Device Kit	Trividia Health, Inc.	56151-1494-01, 08528-1474-01, 87701-0427-39, 87701-0426-25, 11917-0173-89	All TRUE METRIX, TRUE METRIX AIR, TRUE METRIX GO, and TRUE METRIX PRO, branded Blood Glucose Meters distributed in the United States.	The system labeling (user manual and online labeling) did not provide adequate directions for users to act on the E-5 error code. The error code description and recommended actions for the E-5 error code in the labeling do not: (a) clearly emphasize that an E-5 error code could represent a very high blood glucose level, and (b) prominently convey the appropriate immediate actions, such as contacting a healthcare professional. This could lead to a delay in treatment if the user does not seek medical attention immediately when receiving an E-5 error code and experiencing symptoms of high glucose.
5/6/2026	Class 2	Gentle Severe 0.3 % Gel	Alcon Research LLC	00065-8064-01	9T20, 9T50, 9T59, 1U30, 1U48, EXP 4/30/2026; 4V15, EXP 8/31/2026; 7V61, EXP 12/31/2026; 1W47, 1W49, 1X14, EXP 3/31/2027	A lack of assurance of sterility. Due to FDA inspection observations that it believes may impact product quality.

RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
5/6/2026	Class 2	Systane Night 0.3 % Gel	Alcon Research LLC	00065-0474-01	9T21, EXP 4/30/2026; 1U63, 2U47, EXP 5/31/2026; 6V00, 6V12, 8V54, EXP 1/31/2027; 9V55, 8V58, 9V39, 9V97, EXP 2/28/2027; 1W39, 1W40, EXP 3/31/2027; 1X76, EXP 4/30/2027	A lack of assurance of sterility. Due to FDA inspection observations that it believes may impact product quality.
5/13/2026	Class 2	Duloxetine Hcl 20 Mg Cpep	Ajanta Pharma Ltd.	27241-0097-06	PA10734, EXP JUN 2026	Deviations from the Current Good Manufacturing Practices (CGMP), an impurity called N-nitroso-duloxetine being higher than the acceptable limit of 0.83 ppm, identified at the 12-month and 18-month long-term stability intervals.
5/13/2026	Class 2	Duloxetine Hcl 30 Mg Cpep	Ajanta Pharma Ltd.	27241-0098-09, 27241-0098-03, 27241-0098-10	A) PA10774, EXP MAY 2026; B) PA10794, PA12174, EXP JUN 2026; C) PA10804, EXP JUN 2026	Deviations from the Current Good Manufacturing Practices (CGMP), an impurity called N-nitroso-duloxetine being higher than the acceptable limit of 0.83 ppm, identified at the 12-month and 18-month long-term stability intervals.
5/13/2026	Class 2	Duloxetine Hcl 60 Mg Cpep	A) Ajanta Pharma Ltd. B) Breckenridge Pharmaceutical, Inc.	A) 27241-0099-03 B) 51991-0748-90	A) PA07434, EXP MAY 2026 B) 241069C, EXP 05/31/2027	Deviations from the Current Good Manufacturing Practices (CGMP), an impurity called N-nitroso-duloxetine being higher than the acceptable limit of 0.83 ppm, identified at the 12-month and 18-month long-term stability intervals.
5/13/2026	Class 2	Enalapril Maleate 20 Mg Tabs	JB Chemicals and Pharmaceuticals Ltd	64980-0688-10	GEH25023, EXP 6/30/2027	Failing to meet impurity and degradation standards.
5/13/2026	Class 2	Naproxen 125 Mg/5ml Susp	Acella Pharmaceuticals, LLC	42192-0619-16	23F02, EXP 05/2026; 25A37, EXP 01/2028	Chemical contamination, the presence of lead and lithium above specification.
5/13/2026	Class 2	Octreotide Acetate 30 Mg Kit	Teva Pharmaceuticals USA, Inc	00480-9262-08, 00480-9260-01, 00480-9262-08, 00480-9263-21	45011002, EXP 03/31/2027	A lack of assurance of sterility. Due to FDA inspection observations that it believes may impact product quality.

RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
5/20/2026	Class 2	Metoprolol Succinate Er 25 Mg Tb24	Ascend Laboratories, LLC	67877-0590-01	25140859; EXP JAN 2027	Failing to meet dissolution specifications, which may result in the tablets not dissolving properly.

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	Cinacalcet Hcl 30 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219408
Class 2	Cinacalcet Hcl 60 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219409
Class 2	Cinacalcet Hcl 90 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219410
Class 2	Fluocinonide 0.05 % Crea	A) https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219416 B) https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219417 C) https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219418
Class 1	Omnipod 5 Dextg7g6 Intro Gen 5 Kit	https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/insulet-initiates-voluntary-medical-device-correction-certain-omnipodr-5-pods-us#recall-announcement
Class 1	Omnipod 5 Dextg7g6 Pods Gen 5 Misc	https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/insulet-initiates-voluntary-medical-device-correction-certain-omnipodr-5-pods-us#recall-announcement
Class 1	Omnipod 5 G7 Intro (Gen 5) Kit	https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/insulet-initiates-voluntary-medical-device-correction-certain-omnipodr-5-pods-us#recall-announcement
Class 1	Omnipod 5 G7 Pods (Gen 5) Misc	https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/insulet-initiates-voluntary-medical-device-correction-certain-omnipodr-5-pods-us#recall-announcement
Class 1	Omnipod 5 Libre2 Plus G6 Pods Misc	https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/insulet-initiates-voluntary-medical-device-correction-certain-omnipodr-5-pods-us#recall-announcement
Class 2	Furosemide 80 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219264
Class 2	Pantoprazole Sodium 40 Mg Tbec	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219405
Class 1	Gnp True Metrix Glucose Meter W/Device Kit	https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/trividia-health-correction-true-metrix-blood-glucose-monitoring-systems
Class 1	True Metrix Air Glucose Meter W/Device Kit	https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/trividia-health-correction-true-metrix-blood-glucose-monitoring-systems
Class 1	True Metrix Go Glucose Meter W/Device Kit	https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/trividia-health-correction-true-metrix-blood-glucose-monitoring-systems

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	True Metrix Meter W/Device Kit	https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/trividia-health-correction-true-metrix-blood-glucose-monitoring-systems
Class 2	Genteal Severe 0.3 % Gel	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219875
Class 2	Systane Night 0.3 % Gel	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219874
Class 2	Duloxetine Hcl 20 Mg Cpep	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=220018
Class 2	Duloxetine Hcl 30 Mg Cpep	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219822
Class 2	Duloxetine Hcl 60 Mg Cpep	A) https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219823 B) https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219850
Class 2	Enalapril Maleate 20 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219842
Class 2	Naproxen 125 Mg/5ml Susp	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219776
Class 2	Octreotide Acetate 30 Mg Kit	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219871
Class 2	Metoprolol Succinate Er 25 Mg Tb24	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=219887