Judi Rx Drug Recall Report

JULY 2026

Welcome to the **Judi 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 Judi Rx, demonstrate our commitment to providing clients and partners the tools and resources they desire.

WHO WE ARE

Judi 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. Judi 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 **7/15/2026****

Privacy Statement:

Judi Rx, Inc., on behalf of itself and its subsidiaries (including but not limited to Judi Health, LLC and Amino, LLC) ("we," "our," or "us") is committed to ensuring that your privacy is protected. This privacy policy describes the types of information we may collect from you or that you may provide to us and our practices for collecting, using, maintaining, protecting, and disclosing that information. This policy applies to information collected or provided through any services offered on or through judihealth.com such as the member web portal, our care navigation platform, and our mobile application accessible at the Google Play Store and iOS App Store under the name Judi Rx | Judi Health (collectively, our "Site").

RECALL DATE	RECALL TYPE	DRUG RECALLED	MANUFACTURER	NDC(S) IMPACTED	IMPACTED LOT(S)	REASON FOR RECALL
6/24/2026	Class 2	Alyacen 7/7/7 0.5/0.75/1-35 Mg-Mcg Tabs	Glenmark Pharmaceuticals Inc., USA	68462-0556-29	20240411, EXP 6/30/2026; 20250252, EXP 3/31/2027	Out-of-specification results for total impurities.
6/24/2026	Class 2	Aripiprazole 30 Mg Tabs	Ajanta Pharma USA Inc	27241-0056-03	PA00805, EXP 01/31/2029	A product mix-up; a bottle containing Voriconazole Tablets 50 mg was labelled and distributed as Aripiprazole Tablets USP 30 mg.

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6/24/2026	Class 2	Chlorthalidone 25 Mg Tabs	Inventia Healthcare Limited	64980-0599-01, 64980-0599-10	100 TABLET BOTTLE] BATCH RISA24001; [1000 TABLET BOTTLE] BATCH RISB24002; EXP 04/2027	Failure to meet dissolution specifications, which may result in the tablets not dissolving as intended.
7/01/2026	Class 2	Corlanor 5 Mg Tabs	Amgen, Inc.	55513-0800-99, 55513-0800-60	A) 1138901, EXP 08/31/2026; 1149846, EXP 04/30/2027 B) #1138201, 1138202, 08/31/2026, 1138900, 1141143, EXP 08/31/2026; 1140280, 1140281, 1142063, EXP 10/31/2026; 1142062, 1142942, 1142943, EXP 11/30/2026; 1144081, 1144104, EXP 12/31/2026; 1146373, 1146374, EXP 01/31/2027; 1147491, 1147492, 1149843, EXP 03/31/2027; 1151113, 1151890, 1153401, EXP 07/31/2027; 1151114, 1151115, EXP 09/30/2027; 1152412, 1153399, 1153400, EXP 06/30/2027; 1155834, 1155835, EXP 04/30/2027; 1155836, EXP 09/30/2027; 1158480, 1158481, 1158482, EXP 10/31/2027; 1160354, 1160355, EXP 03/31/2028; 1161593, 1161594, 1161595, EXP 04/30/2028; 1164097, 1164098, 1164099, EXP 06/30/2028; 1165078, 1165079, EXP 07/31/2028; 1165080, EXP 02/29/2028; 1168005, 1168006, 1168007, EXP 09/30/2028; 1169288, 1169289, EXP 10/31/2028; 1172885, EXP 12/31/2028	The presence of a foreign substance.

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7/01/2026	Class 2	Corlanor 7.5 Mg Tabs	Amgen, Inc.	55513-0810-60	1138203, 1142065, EXP 07/31/2026; 1145151, 1145152, 1145153, EXP 2/28/2027; 1148908, 1148909, EXP 05/31/2027; 1162845, 1162846, EXP 11/30/2027; 1166471, 1166472, 1166473, EXP 05/31/2028; 1170615, EXP 08/31/2028	The presence of a foreign substance.
7/01/2026	Class 2	Fenofibrate Micronized 200 Mg Caps	Ajanta Pharma USA Inc	27241-0120-04	PA02216; EXP 12/2029	Deviations from Current Good Manufacturing Practices (CGMP).
7/01/2026	Class 2	Lacosamide 100 Mg Tabs	Annora Pharma Private Limited	31722-0813-60	A253999, A254000, EXP 09/30/2027	The presence of foreign tablets. Complaint received about a possible mix-up of Selexipag 1000 mcg tablets in a bottle of Lacosamide Tablets USP, 100mg.
7/01/2026	Class 2	Perampanel 6 Mg Tabs	SUN PHARMACEUTICAL INDUSTRIES INC	51672-4206-06	AE01763, EXP 9/30/2027	A label mix-up, 10mg Perampanel tablet was found in a Perampanel bottle labeled Perampanel 6 mg tablets.
7/08/2026	Class 2	Amphetamine-Dextroamphet Er 5 Mg Cp24	Lannett Company Inc.	00527-0790-37	25285401A, EXP 04/2027	Deviations from the Current Good Manufacturing Practices (CGMP), N-Nitroso-Cinacalcet being higher than the acceptable limit.
7/08/2026	Class 2	Levothyroxine Sodium 300 Mcg Tabs	ACCORD HEALTHCARE, INC.	16729-0458-15	D2401522, EXP 06/2026	The product being subpotent (not strong enough).

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7/08/2026	Class 2	Prednisolone Acetate 1 % Susp	Lupin Pharmaceuticals Inc.	70748-0332-02, 70748-0332-03	A) HA00937, HA00939, HA00941, HA00971, H00973, HA00975, EXPIRES: 7/31/2026; HA00987, HA00989, HA00991, HA01010, HA01012, HA01014, HA01016, HA01048, EXP 8/31/2026; HA01120, HA01137, HA01129, HA01138, HA01133, EXP 9/30/2026; HA01148, HA01196, HA01198, HA01200, HA01206, HA01208, HA01210, HA01212, HA01214, HA01216, HA01218, HA01220, EXP 10/31/2026; HA01236, HA01238, HA01240, HA01244, HA01246, HA01248, HA01261, HA01263, HA01265, HA01267, HA01278, HA01280, HA01282, HA01284, HA01288, HA01290, HA01292, HA01298, EXP 11/30/2026; HB00002, HB00004, HB00006, HB00017, HB00019, HB00021, HB00023, HB00025, HB00031, HB00033, HB00035, HB00037, HB00039, HB00041, HB00043, EXP 12/31/2026; HB00049, HB00051, HB00076, HB00078, HB00080, HB00082, HB00084, EXP 1/31/2027; HB00128, HB00130, HB00132, HB00134, HB00157, HB00159, HB00161, HB00196, HB00198, HB00200, EXP 2/28/2027; HB00224, HB00226, HB00228, HB00230, HB00240, HB00242, EXP 3/31/2027; HB00416, HB00417, HB00419, HB00439, HB00441, HB00443, HB00463, HB00465, EXP 5/31/2027; HB00561, EXP 6/30/2027; HB00569, HB00657, HB00659, HB00663, HB00671, HB00673, HB00675, HB00677, HB00679, EXP 10/31/2027;	The presence of a foreign substance.

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7/08/2026	Class 2	Prednisolone Acetate 1 % Susp	Lupin Pharmaceuticals Inc.	70748-0332-02, 70748-0332-03	HB00747, HB00749, HB00751, HB00753, EXP 11/30/2027; HB00761, HB00763, HB00765, HB00783, HB00785, HC00002, HC00004, HC00006, HC00018, HC00020, HC00028, EXP 12/31/2027; HC00030, HC00032, EXP 1/31/2028; HC00056, HC00058, HC00062, HC00064, HC00073, HC00075, HC00079, HC00081, HC00083, EXP 2/29/2028; HC00091, EXP 3/31/2028; B) HA00945, HA00947, HA00949, EXP 7/31/2026; HA01160, HA01162, HA01164, EXP 10/31/2026; HA01234, HA01300, HA01302, EXP 11/30/2026; HB00053, HB00055, HB00057, EXP 1/31/2027; HB00107, HB00109, HB00111, HB00151, HB00153, HB00155, EXP 2/28/2027; HB00386, HB00388, HB00390, HB00445, HB00447, EXP 5/31/2027; HB00549, HB00551, HB00553, HB00555, EXP 6/30/2027; HB00700, HB00702, EXP 10/31/2027; HB00771, EXP 11/30/2027; HC00008, HC00010, EXP 12/31/2027; HC00034, HC00038, HC00054, HC00060, EXP 1/31/2028;	The presence of a foreign substance.

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7/08/2026	Class 2	Prednisolone Acetate 1 % Susp	Lupin Pharmaceuticals Inc.	70748-0332-02, 70748-0332-03	C) HA00951, HA00953, HA00955, EXP 7/31/2026; HA01188, HA01190, HA01192, EXP 10/31/2026; HA01304, EXP 11/30/2026; HB00113, HB00115, EXP 1/31/2027; HB00166, HB00168, EXP 2/28/2027; HB00392, HB00394, HB00396, EXP 5/31/2027; HB00545, HB00547, EXP 6/30/2027; HB00704, EXP 10/31/2027; HB00773, EXP 11/30/2027; HC00012, HC00014, EXP 12/31/2027; HC00066, HC00068, EXP 2/29/2028	The presence of a foreign substance.
7/08/2026	Class 2	Synjardy Xr 12.5-1000 Mg Tb24	Boehringer Ingelheim Pharmaceuticals, Inc.	00597-0300-45	3244413; EXP 12/31/2028	Deviations from the Current Good Manufacturing Practices (CGMP). An incorrect detergent was used and the required swab sampling was not performed on a production hopper that had previously been used for a different product.

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	Alyacen 7/7/7 0.5/0.75/1-35 Mg-Mcg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=220817
Class 2	Aripiprazole 30 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=220708
Class 2	Chlorthalidone 25 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=220914
Class 2	Corlanor 5 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=220747
Class 2	Corlanor 7.5 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=221134
Class 2	Fenofibrate Micronized 200 Mg Caps	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=220905
Class 2	Lacosamide 100 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=221039
Class 2	Perampanel 6 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=221159
Class 2	Amphetamine-Dextroamphet Er 5 Mg Cp24	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=221274
Class 2	Cinacalcet Hcl 30 Mg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=221386
Class 2	Levothyroxine Sodium 300 Mcg Tabs	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=221025
Class 2	Prednisolone Acetate 1 % Susp	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=220940
Class 2	Synjardy Xr 12.5-1000 Mg Tb24	https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=221116