The Board of Pharmacy has received notice of the following product recall. The Board strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.

URGENT: DRUG RECALL: ***Pravastatin Sodium Tablets, USP 80 mg container pack (Tablets) (Bottle pack, 90's) NDC: 68462-198-90***

This is to inform you of a voluntary recall to ***Retail level*** *involving the following drug product:* ***Pravastatin Sodium Tablets, USP 80 mg***

NDC	Product Name	Batch Number	Expiry Date
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17211525	Jul-24
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17211535	Jul-24
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17211549	Jul-24
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17211787	Aug-24
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17211801	Aug-24
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17212041	Sep-24
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17212088	Oct-24
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17212106	Oct-24
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17212346	Nov-24
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17212345	Nov-24
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17220053	Dec-24
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17220054	Dec-24
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17220055	Dec-24
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17220309	Jan-25
68462-198-90	PRAVASTATIN SODIUM TABLETS, USP 80MG 90'S	17220310	Jan-25

Recall of these batches have been initiated as an abundance of caution against the Out of Specification results for the test of dissolution (By UV) for the batches 17210413 & 17210903, reported at shelf life time point in the long term stability (commercial batches). Batches 17210413 & 17210903 are not part of recall as these have been expired.

Please see details of product batches listed in above table and refer enclosed product labels for ease in identifying the product. Glenmark Pharmaceuticals Inc. initiated shipment of this product (from 15 batches) from September 14, 2021.

Examine your inventory and if you have any inventory available pertains to batch specified in above table then quarantine it immediately.