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.

Hikma Pharmaceuticals USA Inc. (formerly West-Ward Pharmaceuticals) is initiating a drug recall of two (2) lots of Lorazepam Injection, USP 2mg/mL-1mL vial at the retail level. This recall is being conducted with the knowledge of the Food and Drug Administration. This recall Is being conducted due to Out of Specification for Lorazepam total related compounds observed during retain testing observed due to the elevated Related Compound-C. This recall is limited to the 2 lot numbers listed above. No other Hikma products or lots are impacted by this recall. We have received no related complaints for the subject lots to date. The services of Inmar Rx Solutions, Inc. have been enlisted to facilitate the product recall.

Item Description	Potency	Unit of sale NOC		Lot	Exp. Dates	Ship Dates
Lorazepam Injection, USP	2mg/ml	25 vials/ carton	0641-6044- 25	070086	07/2023	08/31/2020- 09/14/2020
Lorazepam Injection, USP	2mg/ml	25 vials/ carton	0641-6044- 25	070128	07/2023	10/26/2020- 12/14/2020