*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.*

This is to inform you of a product recall by **Hikma** for **Lorazepam Injection, USP 2mg/ml CIV** at the retail level**.**

This recall is being conducted due to out of specification for Lorazepam total related compounds observed during retain testing of the listed lot.

This recall is limited to the lot number listed below. **No other Hikma products or lots are impacted by this recall.**

No adverse event complaints for the subject lot to date have been received.

| Item Description | Potency | Unit of sale | NDC | Lot | Exp. Dates | Ship Dates |
|---|---|---|---|---|---|---|
| Lorazepam Injection, USP | 2mg/ml | 25 vials/ carton | 0641-6044-25 | K24118 | 10/2026 | 12/23/2024-01/13/2025 |