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.

Please be advised that Breckenridge Pharmaceutical, Inc. (Breckenridge) is performing a Consumer Level Recall of Propranolol Hydrochloride Extended-Release Capsules, USP, 60mg, manufactured by Adare Pharma Solutions.

Only the lot listed below is being recalled due to potential metal contamination identified as austenitic stainless steel. The lot is being recalled because it shared the same intermediate material that was suspected to be the source of metal contamination observed for a separate lot that was not released to market.

Distribution Range is between October 26, 2026 – November 4, 2026

Use of the affected lot could potentially cause laceration of soft tissues of the oral cavity, esophagus, stomach and intestine. To date, Breckenridge is not aware of reports of adverse events that have been assessed to be related to this recall.

This recall is being initiated with the knowledge of the Food and Drug Administration and should be carried out to the Consumer Level.

PRODUCT: Propranolol Hydrochloride Extended-Release Capsules, USP, 60mg,100-count

NDC NUMBER: 51991-817-01

LOT NUMBER: PK25110035

EXPIRATION DATE: 05/2029