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.

Leading Pharma would like to notify you of a recall involving Furosemide Tablets USP, 80mg, Tablets, 80mg, 75 CC AMBER HDPE BOTTLE 100's count.

The reason for recall is the detection of Nitrosamine Drug Substance-Related Impurities (NDSRI), N-nitroso-Furosemide (NNF) above the FDA Recommended Intake Limit in Lot #H03125, Lot #H03225 and Lot #H03325. Use/consumption of this product may lead to possible potential carcinogenicity.

This product was shipped between 10/17/2025 to 02/09/2026.

PRODUCT: Furosemide Tablets USP, 80mg

NDC NUMBER: 69315-118-01

LOT NUMBERS: H03125, H03125, H03325