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.

Alembic is initiating a Recall to the RETAIL LEVEL for One Lot of Celecoxib 200 mg. This recall has been initiated due to a Product Mix-up. Alembic received a complaint stating one pill of Tadalafil 5mg table was found in a bottle of Celecoxib 200mg capsules.

Health Hazard: If accidentally administered, chances of developing of serious adverse event associated with single dose of Tadalafil 5 mg are unlikely. This recall is being carried out with the knowledge of the US Food and Drug Administration. The below marketed product is subject to the recall. This recall is to the Retail Level Only.

PRODUCT: Celecoxib 200mg 50ct

NDC NUMBER: 50268-169-15

LOT NUMBER: 47881

INITIAL SHIP DATE: 03/24/2025

EXPIRATION DATE: 05/2026