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 notify you that Zydus Pharmaceuticals (USA) Inc. ("Zydus"), is recalling one lot of the drug product mentioned below at the **RETAIL LEVEL.** Zydus has decided to initiate a recall of one lot of Esomeprazole Magnesium for Delayed-Release Oral Suspension 40mg. The product is being recalled due a market complaint received for an incorrect NDC number where an incorrect NDC number (68382-848-93) was printed on a unit dose packet of Esomeprazole Magnesium for Delayed-Release Oral Suspension 40mg. Zydus' investigation is in progress and based on the preliminary investigation, the strength printed on the unit dose packet as well as the product inside the unit dose packet is correct; only the NDC number printed on the unit dose packet is incorrect. As patient safety remains our utmost priority, we are recalling this lot at the retail level.

PRODUCT: Esomeprazole Magnesium for Delayed Release Oral Suspension 40mg, 30 count

NDC NUMBER: 68382-849-94 (carton pack) 68382-849-93 (unit dose packet)

LOT NUMBER: M408002

EXPIRATION DATE: 05/2026

DISTRIUBUTION DATES: From 09/20/2024 to 10/31/2024