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.

American Health Packaging, Inc. is initiating a drug recall to the RETAIL LEVEL for AHP Meclizine Hydrochloride Tablets, USP, 12.5 mg, 50 UD; Carton NDC#: 60687-775-65, (Individual Dose NDC#: 60687-775-11), for the lot listed below.

This recall is being initiated due to a loss on drying out-of-specification result for Meclizine Hydrochloride Tablets, USP in the American Health Packaging repackaged configuration.

Meclizine hydrochloride tablets are indicated for the treatment of vertigo associated with diseases affecting the vestibular system in adults.

The isolated loss-on-drying value of 5.3% does not present a meaningful risk to patient safety or therapeutic effectiveness, as product stability, performance, and chemical integrity remain supported by available data.

PRODUCT: AHP Meclizine Hydrochloride Tablets, USP, 12.5 mg, 50 UD

NDC NUMBER: Carton NDC#: 60687-775-65 (Individual Dose NDC#: 60687-775-11)

AHP LOT NUMBER: 1024852

EXPIRY DATE: 09/30/2026

SHIP DATES OF PRODUCT: 07/29/2025 to 09/16/2025