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 Carbamazepine Extended-Release Tablets USP, 400 mg, 30 UD; Carton NDC#: 60687-594-21, (Individual Dose NDC: 60687-594-11).

This recall is being initiated due to an Out of Specification (OOS) result in the repackaged product for dissolution. Carbamazepine Extended-Release Tablets USP are indicated for use as an anticonvulsant drug and in the treatment of the pain associated with true trigeminal neuralgia. There is a potential that the patient may not receive the active ingredient in the expected time frame. No market complaints have been received for this batch at this time. Out of an abundance of caution, American Health Packaging is initiating a voluntary recall of the finished product batch.

PRODUCT: AHP Carbamazepine Extended-Release Tablets USP 400 mg, 30 UD

NDC NUMBER: Carton NDC#: 60687-594-21 (Individual Dose NDC: 60687-594-11)

LOT NUMBER: 1024078

EXPIRATION DATE: 08/31/2026

SHIP DATES OF PRODUCT: 06/05/2025 to 08/29/2025