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 Glipizide Extended-Release Tablets, 2.5 mg, 30 UD; Carton NDC#: 60687-480-21, (Individual Dose NDC: 60687-480-11), for the lot listed below. This recall is being initiated due to an out-of-specification for dissolution at the initial stability time point of the product repackaged configuration. Glipizide extended-release tablets are indicated as an adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus.

PRODUCT: AHP Glipizide Extended-Release Tablets,2.5 mg, 30 UD

NDC NUMBER: Carton NDC#: 60687-480-21 (Individual Dose NDC: 60687-480-11)

LOT NUMBER: 1012910

EXPIRATION DATE: 04/30/2025

SHIP DATES OF PRODUCTS: 06/05/2023 to 09/05/2023