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 Febuxostat Tablets, 40 mg, 30 UD; Carton NDC#: 60687-538-21, (Individual Dose NDC: 60687-538-11), for the lot listed below. This recall is being initiated in support of the recall by the manufacturer (Sun Pharmaceutical Industries, Inc.) dated January 16, 2024, which included lots that were repackaged by American Health Packaging. Sun stated that "This recall has been initiated in response to microbial contamination in stagnant water found in the duct of the manufacturing equipment." Febuxostat tablets are xanthine oxidase (XO) inhibitor indicated for the chronic management of hyperuricemia in adult patients with gout who have an inadequate response to a maximally titrated dose of allopurinol, who are intolerant to allopurinol, or for whom treatment with allopurinol is not advisable.

PRODUCT NAME: AHP Febuxostat Tablets, 40 mg, 30 UD

NDC NUMBER: Carton NDC#: 60687-538-21(Individual Dose NDC: 60687-538-11)

AHP LOT NUMBER: 1015033

EXPIRATION DATE: 06/30/2025

PRODUCT SHIP DATES: 10/11/2023 to 01/22/2024