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 is initiating a recall 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: AHP Febuxostat Tablets, 40 mg, 30 UD

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

LOT NUMBER: 1015033

EXPIRATION DATE: 06/30/2025

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