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.

BE Pharmaceuticals is recalling Fosaprepitant for Injection, 150 mg per vial, NDC # 71839- 104-01. This recall is being carried out due to an aseptic process simulation failure. BE Pharmaceuticals has not received any reports of adverse events on the subject lot. Injection of potentially contaminated product may result in non-gastrointestinal tract infections which are serious, life threatening or may lead to fatal outcome based on the immune status of the patient.

PRODUCT NAME: Fosaprepitant for Injection, 150 mg per vial (1x24s)

NDC NUMBER: 71839- 104-01

LOT NUMBER: 13D012AA

EXPIRATION DATE: 08/31/2025

DISTRIBUTION DATES: October 25, 2023 to November 08, 2023