

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.

Product Name: Fosaprepitant for Injection, 150 mg / vial in a 10 mL vial

NDC Number: 63323-972-10

Product Code: 972010

Batch Number	Expiration Date	First Ship Date	Last Ship Date
6122760	08/2021	09/30/2019	12/30/2019
6122761	08/2021	12/30/2019	04/06/2020
6122762	09/2021	12/02/2019	06/16/2020
6123883	03/2022	05/20/2020	06/22/2020

Fresenius Kabi USA LLC is recalling the above-mentioned batches of Fosaprepitant for Injection, 150 mg / vial in a 10 mL vial to the user level. Fresenius Kabi has decided to take this action due to the carton label and product insert incorrectly stating the quantity of the excipient edetate disodium (EDTA) as 5.4 mg / vial, rather than the actual amount of 18.8 mg / vial. The investigation reveals that this issue is limited to the product batches indicated above.