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.

On August 22, 2022, BD initiated a product notification for the BD 0.9% Sodium Chloride Injection USP in 1000mL freeflex® bag under lots 22CU10003 and 22CU10004. There have been no reports of patient harm or injury. This product is manufactured by Fresenius Medical Care (FMC), a subsidiary of Fresenius Kabi. BD is expanding the scope of this notification to include an additional lot, 22CU10002.

BD has received reports of the BD 0.9% Sodium Chloride Injection USP in 1000mL freeflex® bag Global Trade Identification Number (GTIN) scanning with an incorrect GTIN. There are no defects with respect to the actual product content, and all other contents of the labeling are correct. FMC confirmed by inspection of retained samples that both the barcode and GTIN 2D barcode on the bag label were incorrect. Upon further investigation, FMC determined that this error occurred after changeover from manufacture of the Fresenius Kabi 500mL Sodium Chloride product. As such, the barcode when scanned, may present with an error code, or tie to the 500mL Sodium Chloride product.

Product: BD 0.9% Sodium Chloride Injection USP in 1000mL freeflex® bag

Lot Number: 22CU10002

No NDC Number Provided

Expiration Date: 03/2024