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	NDC Number	Product Code	Batch Number	Exp. Date	Shipping Dates
Dexmedetomidine HCl in 0.9% Sodium Chloride Injection, 200 mcg / 50 mL (4 mcg / mL), 50 mL fill in a 50 mL vial	63323-671-50	671050	6132925	03/2022	04/09/2020 to 04/13/2020

Fresenius Kabi USA LLC is recalling the above batch of Dexmedetomidine HCl in 0.9% Sodium Chloride Injection, 200 mcg / 50 mL (4mcg / mL), 50 mL fill in a 50 mL vial. Fresenius Kabi is taking this action due to low level carryover of lidocaine present in the above batch. This recall is being performed to the user level.