Drug Recall

DATE OF RECALL: September 02, 2020

DRUG NAME: Dexmedetomidine HCl in 0.9% Sodium Chloride Injection

RECALLING FIRM: Fresenius Kabi USA, LLC

REASON FOR RECALL: This recall was issued due to the possibility of a trace amount of lidocaine present in two lots and our investigation indicates that this issue is limited to two product lots.

FDA LINK: https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/fresenius-kabi-issues-voluntary-nationwide-recall-two-lots-dexmedetomidine-hydrochloride-injection

OTHER DETAILS: See FDA link.

RECALLED PRODUCT:

Product Description	NDC Number	Affected Lots	Expiration Date
Dexmedetomidine HCl in 0.9% Sodium Chloride Injection, 200 mcg/50 mL, 50 mL Dose Bottle, Carton	63323-0671-50	6121853 6122207	05/2021 06/2021
Dexmedetomidine HCl in 0.9% Sodium Chloride Injection, 200 mcg/50 mL, 50 mL Dose Bottle	63323-0671-05	6121853 6122207	05/2021 06/2021