

Drug Recall

DATE OF RECALL: December 27, 2023

DRUG NAME: Various Injectable Products

RECALLING FIRM: Pfizer

REASON FOR RECALL: This recall was issued due to the potential for presence of glass particulate matter, identified during product inspection.

FDA LINK: N/A

OTHER DETAILS: N/A

RECALLED PRODUCT:

Product Description	NDC Number	Lot Number	Expiration Date
4.2% Sodium Bicarbonate Injection, USP ABBOJECT® Glass Syringe 5 mEq/10 mL (0.5 mEq/mL) Carton	00409-5534-24	GX1542	1/1/2025
4.2% Sodium Bicarbonate Injection, USP ABBOJECT® Glass Syringe 5 mEq/10 mL (0.5 mEq/mL) Case	00409-5534-14	GX1542	1/1/2025
8.4% Sodium Bicarbonate Injection, USP Lifeshield® ABBOJECT® Glass Syringe 50 mEq/50 mL (1 mEq/mL) Carton	00409-6637-24	HA7295	3/1/2025
8.4% Sodium Bicarbonate Injection, USP Lifeshield® ABBOJECT® Glass Syringe 50 mEq/50 mL (1 mEq/mL) Case	00409-6637-14	HA7295	3/1/2025
Atropine Sulfate Injection, USP LifeShield® ABBOJECT® Glass Syringe 1 mg/10 mL (0.1 mg/mL) Carton	00409-4911-11	GY2496	2/1/2023
Atropine Sulfate Injection, USP LifeShield® ABBOJECT® Glass Syringe 1 mg/10 mL (0.1 mg/mL) Case	00409-4911-34	GY2496	2/1/2023