

Drug Recall

DATE OF RECALL: August 04, 2026

DRUG NAME: Morphine Sulfate Injection

RECALLING FIRM: Fresenius Kabi

REASON FOR RECALL: Fresenius Kabi announced a voluntary, consumer level recall of one lot of morphine sulfate injection, Simplist® 2 mg/1 mL due to a label mix-up. The MicroVault labeled as morphine 2 mg/1 mL, may contain a prefilled syringe of Dilaudid® (hydromorphone) 0.5 mg/0.5 mL.

FDA LINK: [Fresenius Kabi Issues Voluntary Nationwide Recall of One Lot of Morphine Sulfate Injection, USP in a Simplist® Prefilled Syringe Due to Labeling Mix-up](#)

OTHER DETAILS: N/A

RECALLED PRODUCT:

Product Description	NDC Number	Lot Number	Expiration Date
Morphine Sulfate Injection USP, Simplist® 2 mg / 1 mL, 1 mL fill in a single dose 1 mL Simplist® prefilled syringe	76045-004-01	6402820	12/2028