

## **Drug Recall**

DATE OF RECALL: March 23, 2022

**DRUG NAME:** SYMJEPI (epinephrine) Injection

**RECALLING FIRM:** Adamis Pharmaceuticals Corporation

**REASON FOR RECALL:** This recall was issued due to the potential clogging of the needle preventing the

dispensing of epinephrine.

**FDA LINK:** <a href="https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/adamis-pharmaceuticals-corporation-issues-nationwide-voluntary-recall-symjepir-epinephrine-injection">https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/adamis-pharmaceuticals-corporation-issues-nationwide-voluntary-recall-symjepir-epinephrine-injection</a>

**OTHER DETAILS:** N/A

## **RECALLED PRODUCT:**

Product Description	NDC Number	Affected Lots	<b>Expiration Date</b>
SYMJEPI (epinephrine) Injection 0.15 mg/0.3 mL	78670-0131-02	21101Y	11/30/2022
SYMJEPI (epinephrine) Injection 0.3 mg/0.3 mL	78670-0130-02	21041W	08/31/2022
		21081W	11/30/2022
		21102W	02/28/2023