

Adamis – Recall of Symjepi® (epinephrine) injection

- On March 21, 2022, [Adamis announced](#) a voluntary consumer-level recall of some lots of [Symjepi \(epinephrine\)](#) pre-filled single dose syringes due to the potential clogging of the needle preventing the dispensing of epinephrine.
- US WorldMeds exclusively markets and distributes Symjepi in the U.S., under license from Adamis.

Product Description	NDC#	Lot# (Expiration Date)
Symjepi (epinephrine) 0.15 mg/0.3 mL injection	78670-131-02	21101Y (11/30/2022)
Symjepi (epinephrine) 0.3 mg/0.3 mL injection	78670-130-02	21041W (8/31/2022); 21081W (11/30/2022); 21102W (2/28/2023)

- Symjepi is indicated in the emergency treatment of allergic reactions (Type I) including anaphylaxis to stinging insects (e.g., order Hymenoptera, which include bees, wasps, hornets, yellow jackets, and fire ants) and biting insects (e.g., triatoma, mosquitoes), allergen immunotherapy, foods, drugs, diagnostic testing substances (e.g., radiocontrast media) and other allergens, as well as idiopathic anaphylaxis or exercise-induced anaphylaxis.
- If a person is experiencing an allergic reaction and/or anaphylaxis and is unable to access life-saving epinephrine due to the syringe malfunction, it can lead to life threatening consequences including death.
- Although not confirmed to be related to the recall, there have been two different customer complaints on three syringes, regarding difficulty in dispensing the product, to date. However, neither US WorldMeds nor Adamis has received, or is aware of, any adverse events related to this recall.
- Anyone with an existing inventory of the recalled product should stop use, distribution and quarantine the product immediately. Recalled product may be returned or discarded.
- Patients should contact their physician or health care provider if they have experienced any problems that may be related to taking or using the recalled Symjepi.
- Contact US WorldMeds by phone at **1-888-900-8796** or by email at medinfo@usworldmeds.com for more information about the recall.