UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 8-K

CURRENT REPORT

PURSUANT TO SECTION 13 OR 15(d) OF THE
SECURITIES EXCHANGE ACT OF 1934

Date of report (Date of earliest event reported): March 21, 2022

ADAMIS PHARMACEUTICALS CORPORATION

(Exact Name of Registrant as Specified in Charter)

Delaware	**0-26372**	**82-0429727**
(State or other jurisdiction of incorporation)	(Commission File Number)	(IRS Employer Identification No.)

11682 El Camino Real, Suite 300 **San Diego, CA** (Address of Principal Executive Offices)	**92130** (Zip Code)

Registrant's telephone number, including area code: **(858) 997-2400**

(Former name or Former Address, if Changed Since Last Report.)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions (see General Instruction A.2. below):

☐ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
☐ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
☐ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
☐ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Securities registered pursuant to Section 12(b) of the Exchange Act:

Title of each class	**Trading Symbol(s)**	**Name of each exchange on which registered**
Common Stock	ADMP	NASDAQ Capital Market

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company ☐

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☐

Item 8.01 Other Events.

On March 21, 2022, Adamis Pharmaceuticals Corporation (the "Company") issued a press release announcing that it is voluntarily recalling certain lots of SYMJEPI (epinephrine) Injection 0.15 mg (0.15 mg/0.3 mL) and 0.3 mg (0.3 mg/0.3 mL) Pre-Filled Single-Dose Syringes to the consumer level. The two batches identified in the table contained in the press release are being recalled due to the potential clogging of the needle preventing the dispensing of epinephrine. US WorldMeds ("USWM") exclusively markets and distributes SYMJEPI in the United States, under a license from the Company, the NDA holder. USWM will handle the entire recall process for the Company, with Company oversight. SYMJEPI is manufactured and tested for the Company by Catalent Belgium S.A. As of the date of this Report, neither USWM nor the Company has received, or is aware of, any adverse events related to this recall. The recall is being conducted with the knowledge of the U.S. Food and Drug Administration.

A copy of the Company's press release is filed with this Report as Exhibit 99.1 and is incorporated by reference into this Item 8.01.

Forward Looking Statements

This Current Report on Form 8-K, and the press release filed as an exhibit with this Report, contain forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995, that are subject to risks, uncertainties and other factors, including risks and uncertainties related to this voluntary recall and any future recalls of SYMJEPI and any potential adverse events related to the recall(s). These risks, uncertainties and other factors could cause actual results to differ materially from those referred to in the forward-looking statements. You should not place undue reliance on any forward-looking statements. Further, any forward-looking statement speaks only as of the date on which it is made, and except as may be required by applicable law, we undertake no obligation to update or release publicly the results of any revisions to these forward-looking statements or to reflect events or circumstances arising after the date of this Report. Certain of these risks and additional risks, uncertainties, and other factors are described in greater detail in the Company's filings from time to time with the Securities and Exchange Commission, including its annual report on Form 10-K for the year ended December 31, 2020 and subsequent filings with the SEC, which Adamis strongly urges you to read and consider, all of which are available free of charge on the SEC's web site at http://www.sec.gov.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

99.1	Press release dated March 21, 2022.

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the Registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

ADAMIS PHARMACEUTICALS CORPORATION

Dated: March 22, 2022

By: /s/ David C. Benedicto
Name: David C. Benedicto
Title: Chief Financial Officer

Exhibit 99.1

Adamis Pharmaceuticals Corporation Issues Nationwide Voluntary Recall of SYMJEPI® (epinephrine) Injection for Potential Manufacturing Defect

San Diego, March 21, 2022 – Adamis Pharmaceuticals Corporation (Nasdaq: ADMP) is voluntarily recalling certain lots of SYMJEPI (epinephrine) Injection 0.15 mg (0.15 mg/0.3 mL) and 0.3 mg (0.3 mg/0.3 mL) Pre-Filled Single-Dose Syringes to the consumer level. The batches in the table below are being recalled due to the potential clogging of the needle preventing the dispensing of epinephrine. US WorldMeds (USWM) exclusively markets and distributes SYMJEPI in the United States, under license from Adamis, the NDA holder. USWM will handle the entire recall process for Adamis, with Adamis oversight. SYMJEPI is manufactured and tested for Adamis by Catalent Belgium S.A.

Risk Statement:
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 Pharmaceuticals has received, or is aware of, any adverse events related to this recall.

The recall encompasses all of the following batches, within expiry:

Product	Strength	NDC	Lot	Expiration
SYMJEPI (epinephrine) Injection	0.15 mg/0.3 mL	78670-131-02	21101Y	11/30/2022
	0.3 mg/0.3 mL	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.

The products are packaged in 2-count Pre-Filled Single-Dose Syringes per carton and were distributed nationwide in the USA and directly to customers and/or medical facilities. The products can be identified by the label containing the US WorldMeds name and logo pictured on the cartons in the links below.

US WorldMeds is notifying its customers by email, FDA alerts, and direct outreach. Consumers and institutions that have products that are subject to this recall should stop using the products immediately and may either return or discard the recalled lots. Consumers with questions regarding this recall can call (888) 900-8796 or e-mail questions at medinfo@usworldmeds.com Monday-Friday from 8:00 am to 4:00 pm ET.

Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's MedWatch Adverse Event Reporting program either online, by regular mail or by fax.

- Complete and submit the report Online
- Regular Mail or Fax: Download form or call 1-800-332-1088 to request a reporting form, then complete and return to the address on the pre-addressed form, or submit by fax to 1-800-FDA-0178

This recall is being conducted with the knowledge of the U.S. Food and Drug Administration.

Adamis Contacts:

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