# 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): June 28, 2023 (June 27, 2023)

## Atossa Therapeutics, Inc.

(Exact name of registrant as specified in its charter)

Delaware	001-35610	26-4753208
(State or other jurisdiction of incorporation)	(Commission File Number)	(I.R.S. Employer Identification No.)
107 Spring		98104
Seattle, Washington (Address of principal executive offices)		(Zip Code)
Regist	rant's telephone number, including area code: (2	206) 588-0256
Fo	<u>Not Applicable</u> ormer name or former address, if changed since	last report
☐ Soliciting material pursuant to Rule 14a-12 to ☐ Pre-commencement communications pursua ☐ Pre-commencement communications pursua	25 under the Securities Act (17 CFR 230.425) under the Exchange Act (17 CFR 240.14a-12) nt to Rule 14d-2(b) under the Exchange Act (17 nt to Rule 13e-4(c) under the Exchange Act (17 nemerging growth company as defined in Rule	' CFR 240.14d-2(b))
Emerging growth company $\Box$		
If an emerging growth company, indicate by check or revised financial accounting standards provided		extended transition period for complying with any new $\hfill\Box$
Se	ecurities registered pursuance to Section 12(b) o	of the Act:
Title of each class	Trading symbol(s)	Name of each exchange on which registered
	ATOS	The Nasdaq Capital Market

#### Item 7.01. Regulation FD Disclosure.

On June 27, 2023, Atossa Therapeutics, Inc. (the "Company") issued a press release announcing that the Board of Directors of the Company has authorized a program to repurchase common stock, par value \$0.18 per share, of the Company with an aggregate market value of up to \$10 million from time to time through December 31, 2023. The number of shares to be repurchased and the timing of the repurchases, if any, will depend on several factors, including, but not limited to, stock price, trading volume and general market conditions, along with the Company's working capital requirements and general business conditions. The repurchase authorization does not oblige the Company to acquire any particular amount of its common stock. The Board of Directors may suspend, modify, or terminate the stock repurchase program at any time without prior notice.

A copy of the press release is attached as Exhibit 99.1 to this current report and incorporated herein by reference.

The information in Items 7.01 and 9.01 of this report, including Exhibit 99.1 attached hereto, shall not be deemed to be "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or otherwise subject to the liabilities of that Section or Sections 11 and 12(a)(2) of the Securities Act of 1933, as amended. The information contained herein and in the accompanying exhibit shall not be incorporated by reference into any filing with the U.S. Securities and Exchange Commission (the "SEC") made by the Company, whether made before or after the date hereof, regardless of any general incorporation language in such filing.

## Item 9.01. Financial Statements and Exhibits.

(d) Exhibits

Exhibit No. Description

99.1 Press release, dated June 27, 2023

104 Cover page Interactive Data File (embedded within the Inline XBRL document)

#### **Forward-Looking Statements**

Forward-looking statements in this Current Report on Form 8-K and in the accompanying exhibit, which the Company undertakes no obligation to update, are subject to risks and uncertainties that may cause actual results to differ materially from the anticipated or estimated future results, including the risks and uncertainties associated with our ability to execute on our stock repurchase program, including due to changes in stock price or market or business conditions, any variation between interim and final clinical results, actions and inactions by the FDA, the outcome or timing of regulatory approvals needed by the Company including those needed to commence studies of (Z)-endoxifen, lower than anticipated rate of patient enrollment, estimated market size of drugs under development, the safety and efficacy of the Company's products, performance of clinical research organizations and investigators, obstacles resulting from proprietary rights held by others such as patent rights, whether reduction in breast density or in Ki-67 or any other result from a neoadjuvant study is an approvable endpoint for (Z)-endoxifen, whether the Company can complete acquisitions, and other risks detailed from time to time in the Company's filings with the SEC, including without limitation its periodic reports on Form 10-K and 10-Q, each as amended and supplemented from time to time.

\* \* \*

### **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.

June 28, 2023

/s/ Greg Weaver

Greg Weaver

Executive Vice President and Chief Financial Officer

## Atossa Therapeutics Announces \$10M Stock Repurchase Program

**SEATTLE, June 27, 2023 (GLOBE NEWSWIRE)** — Atossa Therapeutics, Inc. (Nasdaq: ATOS), a clinical stage biopharmaceutical company developing innovative proprietary medicines to address significant unmet needs in oncology with a focus on breast cancer, today announces that its Board of Directors has approved a share repurchase program with authorization to purchase up to \$10 million of its common stock through December 31, 2023.

"We are committed to strategically deploying capital to drive long-term value for stockholders," said Dr. Steven Quay, Atossa's President and Chief Executive Officer. "We believe that the current macroeconomic environment along with the long-term prospects of our business provide a unique buyback opportunity and this stock repurchase program reflects our confidence in the execution of our strategic priorities and the bright future of Atossa."

"Given the strength of our balance sheet and the progress we have made with our (Z)-endoxifen development program that includes multiple ongoing Phase 2 studies, we believe the current share price presents an attractive and strategic buying opportunity for our stock," said Greg Weaver, Atossa's Chief Financial Officer. "The approved share repurchase program is designed to create stockholder value over the long term. The size of the repurchase authorization is less than 10% of total cash, as reported on March 31, 2023, and should not have a material impact on our cash runway, which currently exceeds three years."

Under the stock repurchase program, Atossa expects to repurchase shares of its common stock from time to time in open market transactions, privately negotiated block transactions, or other means, including through the use of trading plans intended to qualify under Rule 10b5-1 and 10b-18 of the Securities Exchange Act of 1934, in accordance with applicable securities laws and regulations. The number of shares to be repurchased and the timing of the repurchases, if any, will depend on several factors, including, but not limited to, stock price, trading volume and general market conditions, along with the Company's working capital requirements and general business conditions. The repurchase authorization does not oblige the Company to acquire any particular amount of its common stock. The Board of Directors may suspend, modify, or terminate the stock repurchase program at any time without prior notice.

#### **About Atossa Therapeutics**

Atossa Therapeutics, Inc. is a clinical-stage biopharmaceutical company developing innovative medicines in areas of significant unmet medical need in oncology with a focus on breast cancer. For more information, please visit www.atossatherapeutics.com

#### **CONTACT:**

Eric Van Zanten VP, Investor and Public Relations 610-529-6219 eric.vanzanten@atossainc.com

#### FORWARD LOOKING STATEMENTS

Forward-looking statements in this press release, which Atossa undertakes no obligation to update, are subject to risks and uncertainties that may cause actual results to differ materially from the anticipated or estimated future results, including the risks and uncertainties associated with our ability to execute on our stock repurchase program, due to changes in stock price or market or business conditions, any variation between interim and final clinical results, actions and inactions by the FDA, the outcome or timing of regulatory approvals needed by Atossa including those needed to commence studies of (Z)-endoxifen, lower than anticipated rate of patient enrollment, estimated market size of drugs under development, the safety and efficacy of Atossa's products, performance of clinical research organizations and investigators, obstacles resulting from proprietary rights held by others such as patent rights, whether reduction in breast density or in Ki-67 or any other result from a neoadjuvant study is an approvable endpoint for (Z)-endoxifen, whether Atossa can complete acquisitions, and other risks detailed from time to time in Atossa's filings with the Securities and Exchange Commission, including without limitation its periodic reports on Form 10-K and 10-Q, each as amended and supplemented from time to time.