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): July 31, 2023

Eagle Pharmaceuticals, Inc.

(Exact name of registrant as specified in its charter)

Delaware

001-36306 (Commission File Number) **20-8179278** (IRS Employer Identification No.)

(State or other jurisdiction of incorporation)

50 Tice Boulevard, Suite 315 Woodcliff Lake, NJ (Address of principal executive offices)

07677 (Zip Code)

Registrant's telephone number, including area code: (201) 326-5300

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligations of the registrant under any of the following provisions:

□ 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 Act:

Title of each class	Trading Symbol	Name of each exchange on which registered
Common Stock (par value \$0.001 per share)	EGRX	The Nasdaq Stock Market LLC

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (17 CFR §230.405) or Rule 12b-2 of the Securities Exchange Act of 1934 (17 CFR §240.12b-2).

Emerging growth company \Box

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 7.01 Regulation FD Disclosure.

On July 31, 2023, Eagle Pharmaceuticals, Inc., or the Company, issued a press release announcing updates to certain of its financial guidance for fiscal year 2023. A copy of the press release is furnished as Exhibit 99.1 to this Current Report on Form 8-K and is incorporated herein by reference.

The information furnished under this Item 7.01, including Exhibit 99.1, shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or subject to the liabilities of that section. The information shall not be deemed incorporated by reference into any other filing with the Securities and Exchange Commission made by the Company, regardless of any general incorporation language in such filing.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

Exhibit No.	Description
<u>99.1</u>	Press Release of the Company, dated July 31, 2023.
104	Cover Page Interactive Data File (embedded within the Inline XBRL document).

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.

Dated: July 31, 2023

EAGLE PHARMACEUTICALS, INC.

By: /s/ Scott Tarriff

Scott Tarriff Chief Executive Officer



For Immediate Release

Eagle Pharmaceuticals Raises 2023 Adjusted non-GAAP EPS¹ Guidance

- -- FY 2023 Adjusted non-GAAP earnings per share now expected to be \$4.40 \$4.70¹ --
- -- FY 2023 Adjusted non-GAAP EBITDA expected to be \$78.0 \$84.0 million¹ --
- -- FY 2023 Adjusted non-GAAP R&D expense is reiterated at \$41.0-\$45.0 million¹ --
- -- FY 2023 Adjusted non-GAAP SG&A expense is reiterated at \$86.0-\$90.0 million¹ --

WOODCLIFF LAKE, N.J. — July 31, 2023 — Eagle Pharmaceuticals, Inc. (Nasdaq: EGRX) ("Eagle" or the "Company") today announced that it has raised its Adjusted non-GAAP earnings per share and Adjusted non-GAAP EBITDA guidance for the 2023 fiscal year. Full-year 2023 Adjusted non-GAAP earnings per share is now expected to range from \$4.40 to \$4.70, up from the previously disclosed estimated range of \$4.20 to \$4.53. Full-year 2023 Adjusted non-GAAP EBITDA is now expected to range from \$78.0 to \$84.0 million, up from the previously disclosed estimated range of \$74.0 to \$80.0 million. Full-year 2023 Adjusted non-GAAP research and development and Adjusted non-GAAP SG&A continue to be expected to range from \$41.0-\$45.0 million and \$86.0-\$90.0 million, respectively.

"Due to the strength and momentum of our business, we are raising our full year Adjusted non-GAAP EPS and Adjusted non-GAAP EBITDA guidance. Our previously disclosed Adjusted non-GAAP research and development and Adjusted non-GAAP SG&A expense ranges remain the same as we continue to invest in our robust pipeline of products. We look forward to sharing additional details during our second quarter earnings call on August 8th," stated Scott Tarriff, President and Chief Executive Officer of Eagle Pharmaceuticals.

About Eagle Pharmaceuticals, Inc.

Eagle is a fully integrated pharmaceutical company with research and development, clinical, manufacturing and commercial expertise. Eagle is committed to developing innovative medicines that result in meaningful improvements in patients' lives. Eagle's commercialized products include PEMFEXY®, RYANODEX®, BENDEKA®, BELRAPZO®, TREAKISYM® (Japan), and BYFAVO® and BARHEMSYS® through its wholly owned subsidiary Acacia Pharma Inc. Eagle's oncology and CNS/metabolic critical care pipeline includes product candidates with the potential to address underserved therapeutic areas across multiple disease states. Additional information is available on Eagle's website at <u>www.eagleus.com</u>.

¹ Adjusted non-GAAP EBITDA, adjusted non-GAAP earnings per share, adjusted non-GAAP R&D expense and adjusted non-GAAP SG&A expense are non-GAAP financial measures. For descriptions of these non-GAAP financial measures, please see below.

Forward-Looking Statements

This press release contains "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995, as amended, and other securities law. Forward-looking statements are statements that are not historical facts. Words and phrases such as "anticipated," "forward," "will," "would," "could," "may," "remain," "potential," "prepare," "expected," "believe," "plan," "near future," "belief," "guidance," and similar expressions are intended to identify forward-looking statements. These statements include, but are not limited to, statements regarding the Company's expectations for the design and timing of the planned Phase 2 study, including with respect to enrollment and site selection and the timing thereof; statements regarding the potential of CAL02 to be a medical breakthrough and offer unique or meaningful therapeutic benefits to seriously ill patients, potentially improving the treatment regimen for patients with severe community-acquired pneumonia, shortening the duration of illness and improving patient outcomes; statements regarding potential regulatory exclusivity, CAL02's potential eligibility for fast track and breakthrough therapy designations and the potential for a CAL02 new drug application for the treatment of SCABP to qualify for priority review; statements regarding the Company's expectation to strengthen the patent portfolio for CAL02; and the potential of the Company's pipeline and product candidates to address underserved therapeutic areas across multiple disease states. All of such statements are subject to certain risks and uncertainties, many of which are difficult to predict and generally beyond the Company's control, that could cause actual results to differ materially from those expressed in, or implied or projected by, the forward-looking information and statements. Such risks and uncertainties include, but are not limited to: the impacts of the ongoing COVID-19 pandemic, including interruptions or other adverse effects on clinical trials and delays in regulatory review or further disruption or delay of any pending or future litigation; delay in or failure to obtain regulatory approval of the Company's product candidates and successful compliance with FDA, European Medicines Agency and other governmental regulations applicable to product approvals; the outcome of litigation involving any of its products or that may have an impact on any of its products; the strength and enforceability of the Company's intellectual property rights or the rights of third parties; the risks inherent in drug development and in conducting clinical trials; and those risks and uncertainties identified in the "Risk Factors" sections of the Company's Annual Report on Form 10-K for the year ended December 31, 2022, filed with the Securities and Exchange Commission (the "SEC") on March 23, 2023, the Company's Quarterly Reports on Form 10-Q for the quarters ended March 31, 2023, filed with the SEC on May 9, 2023, and its other subsequent filings with the SEC. Readers are cautioned not to place undue reliance on these forward-looking statements. All forward-looking statements contained in this press release speak only as of the date on which they were made. Except to the extent required by law, the Company undertakes no obligation to update such statements to reflect events that occur or circumstances that exist after the date on which they were made.

Non-GAAP Financial Performance Measures

This press release contains guidance as to adjusted non-GAAP EBITDA, adjusted non-GAAP earnings per share, adjusted non-GAAP R&D expense and adjusted non-GAAP SG&A expense. The Company believes these measures provide investors and management with supplemental information relating to operating performance and trends that facilitate comparisons between periods and with respect to projected information.

Adjusted non-GAAP net income and related earnings per share information excludes amortization expense, stock-based compensation expense, depreciation expense, severance expense, non-cash interest expense, fair value adjustments on equity investment, fair value adjustments related to derivative instruments, foreign currency exchange gain or loss, amortization of inventory step-up and the tax effect of these adjustments.

Adjusted non-GAAP EBITDA excludes interest expense net of interest income, income tax provision, depreciation and amortization expense, stock-based compensation expense, fair value adjustments on equity investment, fair value adjustments related to derivative instruments, foreign currency exchange gain or loss, and severance expense.

Adjusted non-GAAP R&D expense excludes stock-based compensation expense, depreciation expense and severance expense.

Adjusted non-GAAP SG&A expense excludes stock-based compensation expense, depreciation expense, and severance expense, .

The Company believes the use of non-GAAP financial measures helps indicate underlying trends in the Company's business and are important in comparing current results with prior period results and understanding projected operating performance. Non-GAAP financial measures provide the Company and its investors with an indication of the Company's baseline performance before items that are considered by the Company not to be reflective of the Company's ongoing results.

Investors should note that reconciliations of the forward-looking or projected non-GAAP financial measures included in this press release to their most comparable GAAP financial measures cannot be provided because the Company cannot do so without unreasonable efforts due to the unavailability of information needed to calculate the reconciling items and the variability, complexity, and limited visibility of comparable GAAP measures, and the reconciling items that would be excluded from the non-GAAP financial measures in the future. Likewise, the Company is unable to provide projected GAAP financial measures. GAAP projections and reconciliations of the components of projected adjusted non-GAAP EBITDA, adjusted non-GAAP R&D expenses, and adjusted non-GAAP earning per share to their most comparable GAAP financial measures are not provided because the quantification of projected GAAP R&D expenses, and adjusted non-GAAP earnings per share and the reconciling items between projected GAAP to adjusted non-GAAP EBITDA, adjusted non-GAAP earnings per share cannot be reasonably calculated or predicted at this time without unreasonable efforts. For example, with respect to GAAP net income and R&D Expense, the Company is not able to calculate the favorable or unfavorable expenses related to the fair value adjustments on equity investments and derivative instruments primarily due to nature of these transactions. Such unavailable information could be significant such that actual GAAP net income, R&D expenses, and earnings per share.

These non-GAAP financial measures should be considered in addition to, but not as a substitute for, the information prepared in accordance with U.S. GAAP. In addition, from time to time in the future there may be other items that the company may exclude for purposes of its non-GAAP financial measures; and the Company has ceased, and may in the future cease, to exclude items that it has historically excluded for purposes of its non-GAAP financial measures. For example, commencing in 2023, the Company no longer excludes expense of acquired in-process research & development from the Company's adjusted non-GAAP net income or adjusted non-GAAP EBITDA, their line item components, and non-GAAP earnings per share. Likewise, the Company may determine to modify the nature of its adjustments to arrive at its non-GAAP financial measures. The Company strongly encourages investors to review its consolidated financial statements and publicly-filed reports in their entirety and cautions investors that the non-GAAP financial measures used by other companies, even when similar terms are used to identify such measures.

Investor Relations for Eagle Pharmaceuticals, Inc.:

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