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

August 11, 2020

Date of Report (Date of earliest event reported)

ETON PHARMACEUTICALS, INC.

(Exact name of registrant as specified in its charter)

Delaware (State of incorporation) 001-38738 (Commission File Number) 37-1858472 (I.R.S. Employer Identification Number)

21925 W. Field Parkway, Suite 235 Deer Park, Illinois 60010-7208 (Address of principal executive offices) (Zip code)

(847) 787-7361 (Registrant's telephone number, including area code)

Common Stock, par value \$0.001 per share	ETON	NASDAQ Global Market
Title of each class	Trading symbol(s)	Name of each exchange on which registered
Securitie	s registered pursuant to Section 12	(b) of the Act:
] Pre-commencement communications pursuant to Rule	e 13e-4(c) under the Exchange Act	(17 CFR 240.13e-4(c))
] Pre-commencement communications pursuant to Rule	e 14d-2(b) under the Exchange Act	(17 CFR 240.14d-2(b))
Soliciting material pursuant to Rule 14a-12 under the	Exchange Act (17 CFR 240.14a-12	2)
Written communications pursuant to Rule 425 under to	the Securities Act (17 CFR 230.425	5)
Check the appropriate box below if the Form 8-K filing following provisions:	g is intended to simultaneously s	atisfy the filing obligation of the registrant under any of the

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 [X]

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. [X]

Item 8.01 Other Events.

On August 11, 2020, Eton Pharmaceuticals, Inc. issued a press release reporting that its partner has not yet received a communication from the U.S. Food and Drug Administration (FDA) regarding its decision on the review of EM-100. A copy of the press release is attached as Exhibit 99.1 to this Current Report on Form 8-K.

Item 9.01 Financial Statements and Exhibits

Exhibit 99.1 Press Release dated August 11, 2020

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.

Date: August 11, 2020 By: /s/ W. Wilson Troutman

W. Wilson Troutman Chief Financial Officer and Secretary (Principal Financial Officer)

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Eton Pharmaceuticals Provides Update on the FDA Review of EM-100

DEER PARK, Ill., Aug 11, 2020 (GLOBE NEWSWIRE) — Eton Pharmaceuticals, Inc (Nasdaq: ETON), a specialty pharmaceutical company focused on developing and commercializing innovative drug products, reported that its partner has not yet received a communication from the U.S. Food and Drug Administration (FDA) regarding its decision on the review of EM-100. EM-100's Generic Drug User Fee Act (GDUFA) target action date was August 10th. The company is not aware of any information requests outstanding and expects the FDA to communicate a decision in the coming days.

About Eton Pharmaceuticals

Eton Pharmaceuticals, Inc. is a specialty pharmaceutical company focused on developing, acquiring, and commercializing innovative products. Eton is primarily focused on hospital injectable and pediatric rare disease products. The company's first commercial product, Biorphen, is the only FDA approved ready-to-use formulation of phenylephrine injection and was launched in December 2019. The company's lead pediatric product is the orphan drug Alkindi® Sprinkle, which is currently under review with the FDA. The company has an additional seven products under development, including five that are under review with the FDA.

Forward-Looking Statements

Statements contained in this press release regarding matters that are not historical facts are "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995, including statements associated with the expected ability of Eton to undertake certain activities and accomplish certain goals and objectives. These statements include but are not limited to statements regarding Eton's business strategy, Eton's plans to develop and commercialize its product candidates, the safety and efficacy of Eton's product candidates, Eton's plans and expected timing with respect to regulatory filings and approvals, and the size and growth potential of the markets for Eton's product candidates. Because such statements are subject to risks and uncertainties, actual results may differ materially from those expressed or implied by such forward-looking statements. Words such as "believes," "anticipates," "plans," "expects," "intends," "will," "goal," "potential" and similar expressions are intended to identify forward-looking statements. These forward-looking statements are based upon Eton's current expectations and involve assumptions that may never materialize or may prove to be incorrect. Actual results and the timing of events could differ materially from those anticipated in such forward-looking statements as a result of various risks and uncertainties, which include, without limitation, risks associated with the process of discovering, developing and commercializing drugs that are safe and effective for use as human therapeutics, and in the endeavor of building a business around such drugs. These and other risks concerning Eton's development programs and financial position are described in additional detail in Eton's filings with the Securities and Exchange Commission. All forward-looking statements contained in this press release speak only as of the date on which they were made. Eton undertakes no obligation to update such statements to reflect events that occur or circumstances that exist

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