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): December 13, 2019

Eagle Pharmaceuticals, Inc.

(Exact name of registrant as specified in its charter)

Delaware(State or other jurisdiction of incorporation)

001-36306 (Commission File Number)

20-8179278 (IRS Employer Identification No.)

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 provisions:	is intended to simultaneously satisfy the	e filing obligations of the registrant under any of the following
\square Written communications pursuant to Rule 425 u	under the Securities Act (17 CFR 230.42)	5)
\square Soliciting material pursuant to Rule 14a-12 und	ler the Exchange Act (17 CFR 240.14a-1	2)
\square Pre-commencement communications pursuant t	to Rule 14d-2(b) under the Exchange Act	t (17 CFR 240.14d-2(b))
\square Pre-commencement communications pursuant t	to Rule 13e-4(c) under the Exchange Act	t (17 CFR 240.13e-4(c))
Securities registered pursuant to Section 12(b) of the Ad	et:	
Title of each class Common Stock (par value \$0.001 per share)	Trading Symbol EGRX	Name of each exchange on which registered The Nasdaq Global Market
ndicate by check mark whether the registrant is an emeRule 12b-2 of the Securities Exchange Act of 1934 (17) Emerging growth company \Box		le 405 of the Securities Act of 1933 (17 CFR §230.405) or
f an emerging growth company, indicate by check marl evised financial accounting standards provided pursuar	<u> </u>	he extended transition period for complying with any new or \Box

Item 8.01 Other Events.

On December 13, 2019, Eagle Pharmaceuticals, Inc., or the Company, issued a press release announcing that on December 13, 2019, or the Effective Date, the Company entered into a settlement agreement with Eli Lilly and Company relating to the Company's product PEMFEXYTM (pemetrexed for injection), a branded alternative to ALIMTA®. The settlement agreement provides for a release of all claims by the parties and allows for an initial entry of PEMFEXY into the market (equivalent to approximately a three week supply of current ALIMTA utilization) on February 1, 2022 and a subsequent uncapped entry on April 1, 2022. The Company intends to file the information required by Item 1.01 of Form 8-K in a separate Current Report on Form 8-K within four business days of the Effective Date.

A copy of the full text of the press release referenced above is filed as Exhibit 99.1 to this Current Report on Form 8-K and is incorporated herein by reference.

Item 9.01	Financial Statements and Exhibits.
(d)	Exhibits
Exhibit No.	Description
<u>99.1</u>	Press Release dated December 13, 2019

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.

Eagle Pharmaceuticals, Inc.

Dated: December 16, 2019

By: /s/ Scott Tarriff

Scott Tarriff

Chief Executive Officer



For Immediate Release

Eagle Pharmaceuticals Reaches Settlement Agreement with Eli Lilly for PEMFEXY (pemetrexed for injection)

WOODCLIFF LAKE, N.J. — December 13, 2019 — Eagle Pharmaceuticals, Inc. (Nasdaq: EGRX) ("Eagle" or the "Company") today announced that it has reached a settlement agreement with Eli Lilly and Company (NYSE: LLY) ("Lilly") related to the Company's novel product, PEMFEXYTM (pemetrexed for injection), a branded alternative to ALIMTA[®]. The agreement provides for a release of all claims by the parties and allows for an initial entry of PEMFEXYTM into the market (equivalent to approximately a three week supply of current ALIMTA[®] utilization) on February 1, 2022 and a subsequent uncapped entry on April 1, 2022.

About Eagle Pharmaceuticals, Inc.

Eagle is a specialty pharmaceutical company focused on developing and commercializing innovative and differentiated injectable products that address the shortcomings, as identified by physicians, pharmacists and other stakeholders, of existing commercially successful injectable products. Additional information is available on the Company's website at www.eagleus.com.

Forward-Looking Statements

This press release contains forward-looking information within the meaning of the Private Securities Litigation Reform Act of 1995, as amended, and other securities laws. Forward-looking statements are statements that are not historical facts. Words and phrases such as "anticipated," "forward," "will," "would," "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 future events including, but not limited to: the outcome of the review by the U.S. Department of Justice and the Federal Trade Commission of the settlement agreement; anticipated timing of final approval of the PEMFEXY NDA by the U.S. Food and Drug Administration, if at all; and the timing of Eagle's PEMFEXY launch. All of such statements are subject to certain risks and uncertainties, many of which are difficult to predict and generally beyond Eagle'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. Readers are cautioned not to place undue reliance on these forward-looking statements that speak only as of the date hereof, and the Company does not undertake any obligation to revise and disseminate forward-looking statements to reflect events or circumstances after the date hereof, or to reflect the occurrence of or non-occurrence of any events.

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