AMAG PHARMACEUTICALS INC.
Form 8-K
February 05, 2018

UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
WASHINGTON, DC 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): February 5, 2018

AMAG PHARMACEUTICALS, INC.
(Exact name of registrant as specified in its charter)

Delaware
(State or other jurisdiction of incorporation)

001-10865	04-2742593
(Commission File Number)	(IRS Employer Identification No.)

1100 Winter Street Waltham, Massachusetts	02451
(Address of principal executive offices)	(Zip Code)

(617) 498-3300
(Registrant’s telephone number, including area code)

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:

o Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

o Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

o Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

o Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

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

Emerging growth company o

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. o

Item 8.01. Other Events.

On February 5, 2018, AMAG Pharmaceuticals, Inc. issued a press release announcing that the U.S. Food and Drug Administration has approved its application to broaden the existing label for Feraheme® (ferumoxytol injection) beyond the current chronic kidney disease indication to include all eligible adult iron deficiency anemia patients who have intolerance to oral iron or have had unsatisfactory response to oral iron.

A copy of the press release is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

Item 9.01 Financial Statements and Exhibits

(d) Exhibits

Exhibit No.	Description
99.1	Press release, dated February 5, 2018

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 thereunto duly authorized.

AMAG
PHARMACEUTICALS, INC.
By: /s/ Joseph D. Vittiglio
Joseph D. Vittiglio
Executive Vice President,
General Counsel, Quality &
Corporate Secretary

Date: February 5, 2018