

Exchange Act. o

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Item 2.02. Results of Operations and Financial Condition.

The following information and Exhibit 99.1 attached hereto shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, (the “Exchange Act”) or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, (the “Securities Act”) except as expressly set forth by specific reference in such filing. On February 14, 2018, AMAG Pharmaceuticals, Inc. (“the Company”) issued a press release, described below, including an update to the 2017 Makena revenues. A copy of the Company’s press release is furnished herewith as Exhibit 99.1.

Item 7.01 Regulation FD.

On February 14, 2018, the Company issued a press release announcing that the U.S. Food and Drug Administration has approved the Makena[®] subcutaneous auto-injector drug-device combination product as a ready-to-administer treatment to reduce the risk of preterm birth in women who are pregnant with one baby and who spontaneously delivered one preterm baby in the past.

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	<u>Press release, dated February 14, 2018</u>

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 14, 2018