

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): March 2, 2017

SCYNEXIS, Inc.

(Exact name of registrant as specified in its charter)

Delaware
(State or other jurisdiction
of incorporation)

001-36365
(Commission
File Number)

56-2181648
(IRS Employer
Identification No.)

101 Hudson Street, Suite 3610
Jersey City, New Jersey 07302-6548
(Address of principal executive offices, including zip code)

(201)-884-5485
(Registrant's telephone number, including area code)

N/A
(Former name or former address, if changed since last report)

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:

- 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))
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Item 8.01 Other Events.

On March 2, 2017, SCYNEXIS, Inc. announced that the U.S. Food and Drug Administration (the "FDA") informed SCYNEXIS to hold the initiation of any new clinical studies with SCYNEXIS' intravenous ("IV") formulation of SCY-078 until the FDA completes a review of all available pre-clinical and clinical data of the IV formulation of SCY-078. Ongoing and future clinical development using the oral formulation of SCY-078 are unaffected by this regulatory action. A meeting with the FDA to discuss these data and to agree on subsequent clinical studies with the IV formulation of SCY-078 is scheduled for the second quarter of 2017.

Item 9.01 Financial Statements and Exhibits

Exhibit No. Description

99.1 Press release of SCYNEXIS, Inc., dated March 2, 2017

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: March 2, 2017

SCYNEXIS, Inc.
By: /s/ Eric Francois
Eric Francois
Chief Financial Officer

EXHIBIT INDEX

Exhibit No. Description

99.1 Press release of SCYNEXIS, Inc., dated March 2, 2017



SCYNEXIS delays initiation of new clinical studies using the IV formulation of SCY-078 at FDA's request

Ongoing and future clinical development using the oral formulation of SCY-078 are unaffected

JERSEY CITY, N.J., March 2, 2017 (GLOBE NEWSWIRE) – Drug development company SCYNEXIS, Inc. (Nasdaq: SCYX) today announced that the United States Food and Drug Administration (FDA) has informed the Company to hold the initiation of any new clinical studies with the intravenous (IV) formulation of SCY-078 until the FDA completes a review of all available pre-clinical and clinical data of the IV formulation of SCY-078. Ongoing and future trials using the oral formulation of SCY-078 are unaffected by this regulatory action. A meeting with the FDA to discuss these data and to agree on subsequent clinical studies with the IV formulation of SCY-078 is scheduled for the second quarter of 2017.

The clinical hold decision was issued by the FDA following a review of three mild-to-moderate thrombotic events in healthy volunteers receiving the IV formulation of SCY-078 at the highest doses and highest concentrations in a Phase 1 study. The potential contribution of the IV formulation of SCY-078 to these events cannot be ruled out even though rates of thrombotic events due to intravenous catheters reported in the literature are comparable to those observed in the Phase 1 study.

SCYNEXIS is working closely with the FDA to review the data supporting the use of the IV formulation and dose regimen of SCY-078 selected by the Company for its upcoming clinical trials.

About SCYNEXIS

SCYNEXIS is a drug development company committed to the development and commercialization of novel anti-infectives to address significant unmet therapeutic needs. We are developing our lead product candidate, SCY-078, as an oral and IV drug for the treatment of several fungal infections, including serious and life-threatening invasive fungal infections. For more information, visit www.scynexis.com.

Forward Looking Statement

Statements contained in this press release, including but not limited to the statement, regarding the timing of the next meeting with the FDA to discuss the data and agree on subsequent studies, are "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995. Because such statements are subject to risks and uncertainties, actual results may differ materially from those expressed or implied by such forward-looking statements. These risks and uncertainties include, but are not limited, to: risks inherent in SCYNEXIS' ability to successfully develop SCY-078, including SCYNEXIS' ability to resolve the FDA's concerns to lift the clinical hold and obtain FDA approval for SCY-078; the expected costs of studies and when they might begin or be concluded; and SCYNEXIS' reliance on third parties to conduct SCYNEXIS' clinical studies. These and other risks are described more fully in SCYNEXIS' filings with the Securities and Exchange Commission, including without limitation, its most recent Annual Report on Form 10-K under the caption "Risk Factors" and other documents

subsequently filed with or furnished to 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. SCYNEXIS undertakes no obligation to update such statements to reflect events that occur or circumstances that exist after the date on which they were made.

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