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): June 15, 2018

SCPHARMACEUTICALS INC.

(Exact Name of Registrant as Specified in its Charter)

	Delaware **(State or other jurisdiction of incorporation or organization)**	
001-38293 **(Commission File Number)**		**46-5184075** **(I.R.S. Employer Identification No.)**
2400 District Avenue, Suite 310 **Burlington, Massachusetts** **(Address of principal executive offices)**		**01803** **(Zip Code)**

Registrant's telephone number, including area code: (617) 517-0730

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 (*see* General Instruction A.2. below):

☐ 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))

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 ☒

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. ☒

Item 8.01 Other Events.

As previously disclosed, on June 11, 2018, scPharmaceuticals Inc. (the "Company") received a complete response letter ("CRL") from the U.S. Food and Drug Administration (the "FDA") regarding its New Drug Application (the "NDA") under section 505(b)(2) of the Federal Food, Drug, and Cosmetic Act for Furoscix™ Infusor (furosemide), 80 mg/10 mL, drug-device combination product. On June 15, 2018 management implemented a restructuring plan to reduce operating costs and better align its workforce with the needs of its business following receipt of the CRL. Under this restructuring plan, the Company reduced its workforce by approximately 36%, to 27 employees. The Company currently estimates that it will record a charge in the second quarter of 2018 of approximately $529,000, consisting of severance, benefits and outplacement services.

By filing this information, the Company makes no admission as to the materiality of any information in this report. The information contained in this report is intended to be considered in the context of the Company's filings with the U.S. Securities and Exchange Commission (the "Commission") and other public announcements that the Company makes, by press release or otherwise, from time to time. The Company undertakes no duty or obligation to publicly update or revise the information contained in this report, although it may do so from time to time as its management believes is appropriate. Any such updating may be made through the filing of other reports or documents with the Commission, through press releases or through other public disclosure.

This report contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995 and other federal securities laws. Any statements contained herein which do not describe historical facts, including but not limited to statements regarding the effects of the Company's restructuring plan, reduction in force charges, the potential cost saving resulting from these Company's restructuring and reduction in force, and the timing of the FDA review process and the Company's expectations with regard to its NDA for Furoscix are forward-looking statements which involve risks and uncertainties that could cause actual results to differ materially from those discussed in such forward-looking statements. Such risks and uncertainties include, without limitation, risks associated with whether the Company will be able to address the deficiencies raised in the CRL and the receipt of regulatory approval for Furoscix, as well as other risks set forth under the caption "Risk Factors" in the Company's Annual Report on Form 10-K for the year ended December 31, 2017 and its subsequent public filings with the Commission. The Company cautions investors not to place undue reliance on any forward-looking statements, which speak only as of the date they are made. Except as required by law, the Company disclaims any obligation to publicly update or revise any such statements to reflect any change in expectations or in events, conditions or circumstances on which any such statements may be based, or that may affect the likelihood that actual results will differ from those set forth in the forward-looking statements.

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.

SCPHARMACEUTICALS INC.

Date: June 15, 2018

By: /s/ John H. Tucker
Name: John H. Tucker
Title: President, Chief Executive Officer and Principal Executive Officer