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): **June 10, 2013**

Supernus Pharmaceuticals, Inc.

(Exact name of registrant as specified in its charter)

Delaware
(State or other jurisdiction of Incorporation)

0-50440	**20-2590184**
(Commission File Number)	(IRS Employer Identification No.)
1550 East Gude Drive, Rockville MD	**20850**
(Address of principal executive offices)	(Zip Code)

Registrant's telephone number, including area code: **(301) 838-2500**

Not Applicable
(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 (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))

Item 8.01 Other Events

On June 10, 2013, Supernus issued a press release announcing the receipt of a tentative approval letter from the Food & Drug Administration (the "FDA") for Trokendi XR™, a novel once-daily extended release formulation of topiramate. The letter states that the FDA completed its review of the Trokendi XR™ New Drug Application as amended in December 2012 and the product is tentatively approved for use as recommended in the submitted and agreed-upon labeling. A copy of the press release is furnished as Exhibit 99.1 hereto and is incorporated herein by reference.

Item 9.01 Financial Statements and Exhibits

(d) Exhibits

The following document is furnished as an Exhibit pursuant to Item 8.01 hereof:

Exhibit 99.1 — Press Release Dated June 10, 2013.

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.

SUPERNUS PHARMACEUTICALS, INC.

DATED: June 12, 2013

By: /s/ Gregory S. Patrick
Gregory S. Patrick
Vice-President and Chief Financial Officer

EXHIBIT INDEX

Number	Description	
99.1	Press Release dated June 10, 2013.	Attached

Exhibit 99.1

FOR IMMEDIATE RELEASE

Supernus Receives Tentative Approval for Trokendi XR™

Rockville, MD, June 10, 2013 — Supernus Pharmaceuticals, Inc. (Nasdaq:SUPN), a specialty pharmaceutical company, received a tentative approval letter from the Food & Drug Administration (the "FDA") for Trokendi XR™, a novel once-daily extended release formulation of topiramate (formerly known as SPN-538). The letter states that the FDA completed its review of the Trokendi XR™ New Drug Application (the "NDA") as amended in December 2012 and the product is tentatively approved for use as recommended in the submitted and agreed-upon labeling.

"As expected and as previously communicated, most recently during our May 13, 2013 quarterly earnings call, since this approval was granted before the June 22, 2013 date of expiration of the Topamax® data exclusivity, this approval came in the form of a tentative approval. We will now submit the "Request for Final Approval" letter to the FDA based on which we expect to receive Final Approval and then launch our product, as anticipated, in the third quarter of 2013," said Jack Khattar, Chief Executive Officer, President and Director of Supernus.

About Supernus Pharmaceuticals, Inc.

Supernus Pharmaceuticals, Inc. is a specialty pharmaceutical company focused on developing and commercializing products for the treatment of central nervous system, or CNS, diseases. The Company has one marketed product for epilepsy, Oxtellar XR™ (extended release oxcarbazepine), and one tentatively approved product for epilepsy, Trokendi XR™ (extended release topiramate). The Company is also developing several product candidates in psychiatry to address large market opportunities in ADHD including ADHD patients with impulsive aggression. These product candidates include SPN-810 for impulsive aggression in ADHD and SPN-812 for ADHD.

Forward Looking Statements

This press release contains forward-looking statements regarding the timing of the availability of Trokendi XR™ to the market. Actual results may differ materially from those in these forward-looking statements as a result of various factors, including, but not limited to, risks regarding the Company's ability to obtain final approval for its products, commercialize its products successfully, whether physicians will prescribe and patients will use its products, once available, and competition in their respective markets. For a further description of these and other risks facing the company, please see the risk factors described in the Company's Annual Report on Form 10-K that was filed with the United States Securities and Exchange Commission on March 15, 2013 and under the caption "Risk Factors" and the updates to these risk factors in the Company's quarterly report form 10-Q that was filed with the commission on May 15, 2013. Forward-looking statements speak only as of the date of this press release, and the company undertakes no obligation to update or revise these statements, except as may be required by law.

CONTACTS:
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