

HORIZON PHARMA, INC.  
Form 8-K  
August 28, 2013

**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): August 26, 2013**

**Horizon Pharma, Inc.**

**(Exact name of registrant as specified in its charter)**

**Delaware**  
**(State of incorporation)**

**001-35238**  
**(Commission File No.)**

**27-2179987**  
**(IRS Employer Identification No.)**

**520 Lake Cook Road, Suite 520, Deerfield, Illinois**

**60015**

Edgar Filing: HORIZON PHARMA, INC. - Form 8-K

(Address of principal executive offices)

(Zip Code)

Registrant's telephone number, including area code: (224) 383-3000

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))

**Item 8.01 Other Events.**

On August 26, 2013, we filed suit in the United States District Court for the District of New Jersey against Watson Laboratories, Inc. Florida, Actavis Pharma, Inc., Andrx Corp., and Actavis, Inc. (collectively, WLF ). The lawsuit alleges that WLF has infringed U.S. Patent Nos. 6,488,960, 6,677,326, 8,168,218, 8,309,124, and 8,394,407 by filing an Abbreviated New Drug Application ( ANDA ) seeking approval from the U.S. Food and Drug Administration ( FDA ) to market generic versions of RAYO<sup>®</sup>s containing 1 mg, 2 mg, and 5 mg of prednisone prior to the expiration of the patents. The subject patents are listed in the FDA s Approved Drug Products with Therapeutic Equivalence Evaluations, commonly known as the Orange Book. Our commencement of the patent infringement lawsuit stays, or bars, FDA approval of WLF s ANDA for 30 months or until an earlier district court decision that the subject patents are not infringed or invalid.

**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.

Date: August 28, 2013

Horizon Pharma, Inc.

By: /s/ Robert J. De Vaere  
Robert J. De Vaere  
Executive Vice President and Chief Financial Officer