

ACHILLION PHARMACEUTICALS INC  
Form 8-K  
July 01, 2013

**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 28, 2013**

**Achillion Pharmaceuticals, Inc.**

(Exact name of registrant as specified in its charter)

**Delaware**  
(State or other jurisdiction  
of incorporation)

**001-33095**  
(Commission  
File Number)

**52-2113479**  
(IRS Employer  
Identification No.)

**300 George Street**  
**New Haven, CT**  
(Address of principal executive offices)

**Registrant's telephone number, including area code: (203) 624-7000**

**06511**  
(Zip Code)

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N/A

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

Check the appropriate box if the Form 8-K 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**

Achillion Pharmaceuticals, Inc. (the Company) announced today that the Company received notice from the U.S. Food and Drug Administration (the FDA) that a clinical hold has been placed on sovalprevir after elevations in liver enzymes associated with significantly higher than anticipated exposures to atazanavir and sovalprevir were noted in a Phase 1 healthy subject drug-drug interaction study evaluating the effects of concomitant administration of sovalprevir with ritonavir-boosted atazanavir. The FDA has allowed continued enrollment and treatment of patients in the Phase 2 -007 clinical trial evaluating 12-weeks of sovalprevir in combination with ACH-3102 and ribavirin for patients with treatment-naïve genotype 1 hepatitis C viral infection.

The full text of the press release issued in connection with this announcement is attached as Exhibit 99.1 to this Current report on Form 8-K.

**Item 9.01. Financial Statements and Exhibits**

(d) Exhibits

The following exhibit relating to Item 8.01 shall be deemed to be furnished, and not filed:

99.1 Press Release dated July 1, 2013

SIGNATURE

Pursuant to the requirements of the Securities Exchange Act of 1934, as amended, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

ACHILLION PHARMACEUTICALS, INC.

Date: July 1, 2013

By: /s/ Mary Kay Fenton  
Mary Kay Fenton

Chief Financial Officer

Exhibit Index

99.1 Press Release dated July 1, 2013