## Edgar Filing: NOVEN PHARMACEUTICALS INC - Form 8-K

NOVEN PHARMACEUTICALS INC Form 8-K March 16, 2009

# 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 16, 2009

# Noven Pharmaceuticals, Inc.

(Exact name of registrant as specified in its charter)

Delaware	0-17254	59-2767632
(State or other jurisdiction of incorporation)	(Commission File Number)	(I.R.S. Employer Identification No.)
11960 S.W. 144th Street, Miami, Florida		33186
(Address of principal executive offices)		(Zip Code)
Registrant s telephone number, including area code:		305-253-5099
	Not Applicable	
Former name	e or former address, if changed since l	ast report
Check the appropriate box below if the Form 8-K filing he following provisions:	is intended to simultaneously satisfy	the filing obligation of the registrant under any of
<ul> <li>Written communications pursuant to Rule 425 unde</li> <li>Soliciting material pursuant to Rule 14a-12 under the</li> <li>Pre-commencement communications pursuant to Rule</li> <li>Pre-commencement communications pursuant to Rule</li> </ul>	ne Exchange Act (17 CFR 240.14a-12 ule 14d-2(b) under the Exchange Act	) (17 CFR 240.14d-2(b))

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#### <u>Top of the Form</u> Item 8.01 Other Events.

Noven Pharmaceuticals, Inc. ("Noven") today confirmed that Shire plc ("Shire"), the global licensee of Daytrana® (methylphenidate transdermal system), has announced its withdrawal of the European Marketing Authorization Application ("MAA") for Daytrana for the treatment of Attention Deficit Hyperactivity Disorder ("ADHD"). Shire stated that its decision to withdraw the MAA was based on the fact that European regulatory authorities had requested an additional clinical study for Daytrana in a European patient population, and that Shire planned to enter the European ADHD market through the previously-announced pending acquisition of an oral methylphenidate product that is already approved in Europe. Noven further confirmed that its 2009 financial guidance in its Annual Report on Form 10-K, dated March 13, 2009 did not include any revenues or other financial contribution from European sales of Daytrana.

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#### **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.

Noven Pharmaceuticals, Inc.

March 16, 2009 By: /s/ Jeff Mihm

Name: Jeff Mihm

Title: Vice President, General Counsel and Corporate

Secretary