
**UNITED STATES
SECURITIES AND EXCHANGE COMMISSION**
Washington, DC 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)
November 12, 2009

Cadence Pharmaceuticals, Inc.

(Exact name of registrant as specified in its charter)

Delaware
(State or other jurisdiction
of incorporation)

001-33103
(Commission File Number)

41-2142317
(IRS Employer
Identification No.)

12481 High Bluff Drive, Suite 200
San Diego, California 92130
(Address of principal executive offices, including zip code)

(858) 436-1400
(Registrant's telephone number, including area code)

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))
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Item 8.01 Other Events.

On November 13, 2009, Cadence Pharmaceuticals, Inc. (the "Company," or "Cadence") announced that the U.S. Food and Drug Administration ("FDA") has extended the Prescription Drug User Fee Act ("PDUFA") goal date for its Priority Review of the New Drug Application ("NDA") for intravenous acetaminophen by three months. The extended PDUFA goal date is February 12, 2010.

The FDA designated one of Cadence's submissions to the NDA, which contained additional clinical pharmacology data requested by the agency during the review process, as a major amendment. The FDA has the option to extend the PDUFA goal date when a sponsor submits a major amendment to an NDA within three months of the PDUFA goal date to provide time to complete the review. The FDA is not requesting any other information at this time.

