

ACORDA THERAPEUTICS INC

FORM 8-K (Current report filing)

Filed 08/19/08 for the Period Ending 08/18/08

Address	420 SAW MILL RIVER ROAD ARDSLEY, NY 10502
Telephone	914-347-4300
CIK	0001008848
Symbol	ACOR
SIC Code	2836 - Biological Products, Except Diagnostic Substances
Industry	Biotechnology & Drugs
Sector	Healthcare
Fiscal Year	12/31

**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 18, 2008**

Acorda Therapeutics, Inc.

(Exact name of registrant as specified in its charter)

Delaware
(State or other jurisdiction
of incorporation)

000-50513
(Commission
File Number)

13-3831168
(I.R.S. Employer
Identification No.)

15 Skyline Drive, Hawthorne, NY
(Address of principal executive offices)

10532
(Zip Code)

Registrant's telephone number, including area code: **(914) 347-4300**

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:

- 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 August 19, 2008, Acorda Therapeutics, Inc. (the “Registrant”) announced that the underwriter of its public offering of common stock that was announced on August 7, 2008 has exercised its entire option to purchase an additional 600,000 shares of common stock at the public offering price of \$28.50 per share. The exercise of the option increases the size of the offering to an aggregate of 4,600,000 shares of common stock.

The offering is being made pursuant to the Registrant’s shelf registration statement on Form S-3 (Registration No. 333-152826) filed with the Securities and Exchange Commission (the “Commission”) on August 6, 2008, including a related prospectus and prospectus supplement that the Registrant filed with the Commission on August 7, 2008.

The Registrant’s press release announcing that the underwriter has exercised its entire option is attached as Exhibit 99.1 to this Current Report on Form 8-K, and incorporated herein by reference.

Item 9.01. Financial Statements and Exhibits.

99.1 Press Release dated August 19, 2008

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.

Acorda Therapeutics, Inc.

August 19, 2008

By: /s/ David Lawrence

Name: David Lawrence, M.B.A.

Title: Chief Financial Officer

Exhibit Index

<u>Exhibit No.</u>	<u>Description</u>
99.1	Press Release dated August 19, 2008



Acorda Therapeutics, Inc. Announces Exercise of Option by Underwriter

HAWTHORNE, N.Y.— Aug. 19, 2008—Acorda Therapeutics, Inc. (NASDAQ: ACOR) (the “Company”) announced today that Deutsche Bank Securities Inc., as underwriter of the Company’s common stock offering that was announced on August 7, 2008, has exercised its entire option to purchase an additional 600,000 shares of common stock at the public offering price of \$28.50 per share. The exercise of the option increases the size of the offering to an aggregate of 4,600,000 shares of common stock.

Copies of the final prospectus related to the offering may be obtained by contacting Deutsche Bank Securities Inc., Deutsche Bank Securities, Prospectus Department, Harborside Financial Center, 100 Plaza One, Jersey City, New Jersey 07311-3988.

About Acorda Therapeutics

Acorda Therapeutics is a biotechnology company developing therapies for spinal cord injury, multiple sclerosis and related nervous system disorders. The Company’s marketed products include Zanaflex Capsules(R) (tizanidine hydrochloride), a short-acting drug for the management of spasticity. Acorda’s lead clinical product, Fampridine-SR, has completed two Phase 3 clinical trials to evaluate its safety and efficacy to improve walking ability in people with MS. The Company’s pipeline includes a number of products in development for the treatment, regeneration and repair of the nervous system.

Forward-Looking Statement

This press release includes forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. All statements, other than statements of historical facts, regarding management’s expectations, beliefs, goals, plans or prospects should be considered forward-looking. These statements are subject to risks and uncertainties that could cause actual results to differ materially, including delays in obtaining or failure to obtain FDA approval of Fampridine-SR, the risk of unfavorable results from future studies of Fampridine-SR, Acorda Therapeutics’ ability to successfully market and sell Fampridine-SR, if approved, and Zanaflex Capsules(R), competition, failure to protect its intellectual property or to defend against the intellectual property claims of others, the ability to obtain additional financing to support Acorda Therapeutics’ operations, and unfavorable results from its preclinical programs. These and other risks are described in greater detail in Acorda Therapeutics’ filings with the Securities and Exchange Commission. Acorda Therapeutics may not actually achieve the goals or plans described in its forward-looking statements, and investors should not place undue reliance on these statements. Acorda Therapeutics disclaims any intent or obligation to update any forward-looking statements as a result of developments occurring after the date of this press release.

CONTACT: MEDIA AND INVESTOR RELATIONS:

Acorda Therapeutics
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