

Acorda Therapeutics to Host Conference Call to Discuss First Quarter 2014 Financial Results on May 6, 2014

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ARDSLEY, N.Y.--(BUSINESS WIRE)-- Acorda Therapeutics, Inc. (Nasdaq:ACOR) announced that it will host a conference call and webcast to report its first quarter 2014 financial results on Tuesday, May 6 at 8:30 a.m. ET.

To participate in the conference call, please dial 800-706-7745 (domestic) or 617-614-3472 (international) and reference the access code 66119945. The presentation will be available via live webcast on the Investor section of www.acorda.com. Please log in approximately 5 minutes before the scheduled time of the presentation to ensure a timely connection.

A replay of the call will be available from 12:30 a.m. ET on May 6, 2014 until midnight on June 3, 2014. To access the replay, please dial 888-286-8010 (domestic) or 617-801-6888 (international) and reference the access code 83511569. The archived webcast will be available in the Investor Relations section of the Acorda website at www.acorda.com.

About Acorda Therapeutics

Founded in 1995, **Acorda Therapeutics** is a biotechnology company focused on developing therapies that restore function and improve the lives of people with neurological conditions.

Acorda markets three FDA-approved therapies including: **AMPYRA® (dalfampridine)** Extended Release Tablets, 10 mg, a treatment to improve walking in patients with multiple sclerosis (MS); **ZANAFLEX CAPSULES® (tizanidine hydrochloride)** and Zanaflex tablets, a short-acting drug for the management of spasticity; and Qutenza® (capsaicin) 8% Patch, for the management of neuropathic pain associated with postherpetic neuralgia. AMPYRA is marketed outside the United States as FAMPYRA® (prolonged-release fampridine tablets) by Biogen Idec under a

licensing agreement from Acorda.

Acorda has one of the leading pipelines in the industry of novel neurological therapies. The Company is currently developing six clinical-stage therapies and one preclinical stage therapy that address a range of disorders including post-stroke deficits, epilepsy, stroke, peripheral nerve damage, spinal cord injury, neuropathic pain, and heart failure. For more information, please visit the Company's website at: www.acorda.com.

Forward-Looking Statements

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 our ability to successfully market and sell Ampyra in the U.S.; third party payers (including governmental agencies) may not reimburse for the use of Ampyra or our other products at acceptable rates or at all and may impose restrictive prior authorization requirements that limit or block prescriptions; the risk of unfavorable results from future studies of Ampyra or from our other research and development programs, including Diazepam Nasal Spray or any other acquired or in-licensed programs; we may not be able to complete development of, obtain regulatory approval for, or successfully market Diazepam Nasal Spray or other products under development; the occurrence of adverse safety events with our products; delays in obtaining or failure to obtain regulatory approval of or to successfully market Fampyra outside of the U.S. and our dependence on our collaboration partner Biogen Idec in connection therewith; competition, including the impact of generic competition on Zanaflex Capsules revenues; failure to protect our intellectual property, to defend against the intellectual property claims of others or to obtain third party intellectual property licenses needed for the commercialization of our products; failure to comply with regulatory requirements could result in adverse action by regulatory agencies; and the ability to obtain additional financing to support our operations. These and other risks are described in greater detail in Acorda Therapeutics' filings with the Securities & Exchange Commission. Acorda may not actually achieve the goals or plans described in its forward-looking statements, and investors should not place undue reliance on these statements. Forward-looking statements made in this release are made only as of the date hereof, and Acorda disclaims any intent or obligation to update any forward-looking statements as a result of developments occurring after the date of this release.

Source: Acorda Therapeutics, Inc.

Acorda Therapeutics

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