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: May 03, 2010
(Date of earliest event reported)

Corcept Therapeutics Incorporated
(Exact name of registrant as specified in its charter)

DE	**000-50679**	**77-0487658**
(State or other jurisdiction of incorporation)	(Commission File Number)	(IRS Employer Identification Number)

149 Commonwealth Drive, Menlo Park, CA	**94025**
(Address of principal executive offices)	(Zip Code)

650-327-3270
(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:

o Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
o Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
o Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
o Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Item 7.01. Regulation FD Disclosure

On May 3, 2010, we issued a press release updating the status of enrollment in our Phase 3 trial in Cushing's Syndrome and discussing our recent financing. The press release is attached hereto as Exhibit 99.1 and incorporated by reference herein.

This information and the information contained in the press release attached as Exhibit 99.1 shall not be deemed "filed" for purposes of Section 18 of the Exchange Act of 1934, as amended, or otherwise subject to the liabilities of that Section. This information and the information contained in the press release attached as Exhibit 99.1 is not incorporated by reference into any filings of Corcept Therapeutics Incorporated made under the Securities Act of 1933, as amended, whether made before or after the date of this current report on Form 8-K, regardless of any general incorporation language in the filing unless specifically stated so therein.

Item 8.01. Other Events

On May 3, 2010, we announced that we are nearing completion of enrollment in our 50-patient open-label Phase 3 trial of CORLUX in patients with Cushing's Syndrome. Forty-five patients have received treatment with CORLUX and we have numerous patients in the screening phase. The FDA has indicated that this single 50-patient open-label Phase 3 study of CORLUX may provide a reasonable basis for the submission of an NDA for Cushing's Syndrome. We expect to submit an NDA for Cushing's Syndrome by year end.

As previously disclosed, on April 21, 2010, we completed a financing that raised gross proceeds of approximately $7.7 million from the exercise of existing warrants and the private placement of new warrants. With the additional funds from this financing, we now believe we have sufficient capital to support our operations into the second quarter of 2011.

Statements made in this current report on Form 8-K, other than statements of historical fact, are forward-looking statements, including, for example, statements relating to our clinical development and research programs, the timing of the introduction of CORLUX and future product candidates, estimates of the timing of enrollment or completion of our clinical trials and the anticipated results of those trials, and the timing of submission of the NDA, if submitted at all, the ability to create value from CORLUX or other future product candidates and our estimates regarding our capital requirements, spending plans and needs for additional financing. Forward-looking statements are subject to a number of known and unknown risks and uncertainties that might cause actual results to differ materially from those expressed or implied by such statements. For example, we cannot assure you that the cost, rate of spending, completion or success of clinical trials will be favorable, that our financial projections will be accurate or that we will pursue further activities with respect to the development of CORLUX, CORT 108297, or any of our other selective GR-II antagonists. These and other risk factors are set forth in our annual report on Form 10-K for the fiscal year ended December 31, 2009 and subsequent SEC filings. We disclaim any intention or duty to update any forward-looking statement made in this current report on Form 8-K.

Item 9.01. Financial Statements and Exhibits

(d) Exhibits

99.1 Press Release of Corcept Therapeutics Incorporated dated May 03, 2010

SIGNATURE

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.

Dated: May 05, 2010

CORCEPT THERAPEUTICS INCORPORATED

By: /s/ Caroline M. Loewy
Caroline M. Loewy
Chief Financial Officer

Exhibit Index

Exhibit No.	Description
99.1	Press Release of Corcept Therapeutics Incorporated dated May 03, 2010

Corcept Therapeutics Updates Cushing's Syndrome Development and Recent Financing

MENLO PARK, CA -- (Marketwire - May 03, 2010) - Corcept Therapeutics Incorporated (NASDAQ: CORT), a pharmaceutical company engaged in the discovery and development of drugs for the treatment of severe metabolic and psychiatric disorders, today updated the status of enrollment in its Phase 3 trial in Cushing's Syndrome and discussed its recent financing.

"Forty-five of fifty patients have been dosed in our Phase 3 study of CORLUX® for the treatment of Cushing's Syndrome, and the remaining patients are now being screened to enable completion of enrollment in the coming weeks," said Joseph Belanoff, M.D., Chief Executive Officer of Corcept. "We continue to expect to announce results of this study in the fourth quarter of 2010, and we remain on track to submit our New Drug Application (NDA) for the use of CORLUX in Cushing's Syndrome by year end."

Cushing's Syndrome Development Update

We are nearing completion of enrollment in our 50-patient open-label Phase 3 trial of CORLUX in patients with Cushing's Syndrome, which is being conducted at 20 leading medical facilities throughout the United States. The pace of enrollment has been difficult to predict given the unprecedented nature of the trial. However, with 45 patients having received treatment with CORLUX, and numerous patients in the screening phase, we are confident that the end of the enrollment phase of the trial will be quite soon.

The FDA has indicated that this single 50-patient open-label Phase 3 study of CORLUX may provide a reasonable basis for the submission of an NDA for Cushing's Syndrome. In the study, each patient's dose is titrated to clinical benefit by their clinical investigator and the endpoints are measured at the end of 24 weeks. We expect to announce results of this study in the fourth quarter of 2010. All other portions of the NDA are complete or will be completed in time to submit an NDA by year end.

Warrant Transaction

We recently completed a financing which raised gross proceeds of $7.7 million from the exercise of existing warrants and the private placement of new warrants. Investors who received warrants in Corcept's October 2009 private placement agreed to exercise those cash warrants for $1.66 per share. For each warrant exercised, the investor was given the opportunity to purchase a new warrant for $0.125 per share with an exercise price equivalent to the market price at the close of the transaction, $2.96 per share.

With the additional funds from this financing, we now have sufficient capital to support our operations into the second quarter of 2011. Participants in this transaction included existing investors Longitude Capital, Sutter Hill Ventures, Federated Kauffman Funds, and members of our Board of Directors.

About Cushing's Syndrome

Endogenous Cushing's Syndrome is caused by prolonged exposure of the body's tissues to high levels of the hormone cortisol due to either cortisol or ACTH production by tumors. Cushing's Syndrome is an orphan indication which most commonly affects adults aged 20 to 50. An estimated 10 to 15 of every one million people are newly diagnosed with this syndrome each year, resulting in over 3,000 new patients in the United States. An estimated 20,000 patients in the United States have Cushing's Syndrome. Symptoms vary, but most people have one or more of the following manifestations: high blood sugar, diabetes, high blood pressure, upper body obesity, rounded face, increased fat around the neck, thinning arms and legs, severe fatigue and weak muscles. Irritability, anxiety, cognitive disturbances and depression are also common. Cushing's Syndrome can affect every organ system in the body and can be lethal if not treated effectively.

About CORLUX

Corcept's first-generation compound, CORLUX, also known as mifepristone, directly blocks the cortisol (GR-II) receptor and the progesterone (PR) receptor. Intellectual property protection is in place to protect important methods of use for CORLUX. Corcept retains worldwide rights to its intellectual property related to CORLUX.

About Corcept Therapeutics Incorporated

Corcept is a pharmaceutical company engaged in the discovery and development of drugs for the treatment of severe metabolic and psychiatric disorders. The company has two ongoing Phase 3 programs: CORLUX for the treatment of Cushing's Syndrome, and CORLUX for the treatment of the psychotic features of psychotic depression. Corcept also has a Phase 1 program for CORT 108297. Corcept has developed an extensive intellectual property portfolio that covers the use of GR-II antagonists in the treatment of a wide variety of psychiatric and metabolic disorders, including the prevention of weight gain caused by the use of antipsychotic medication, as well as composition of matter patents for our selective GR-II antagonists.

Statements made in this news release, other than statements of historical fact, are forward-looking statements, including, for example, statements relating to Corcept's clinical development and research programs, the timing of the introduction of CORLUX and future product candidates, including CORT 108297, estimates of the timing of enrollment or completion of our clinical trials and the anticipated results of those trials, and the timing of submission of the NDA, if submitted at all, the ability to create value from CORLUX or other future product candidates and our estimates regarding our capital requirements, spending plans and needs for additional financing. Forward-looking statements are subject to a number of known and unknown risks and uncertainties that might cause actual results to differ materially from those expressed or implied by such statements. For example, there can be no assurances with respect to the cost, rate of spending, completion or success of clinical trials; financial projections may not be

accurate; there can be no assurances that Corcept will pursue further activities with respect to the development of CORLUX, CORT 108297, or any of its other selective GR-II antagonists. These and other risk factors are set forth in the Company's SEC filings, all of which are available from our website (www.corcept.com) or from the SEC's website (www.sec.gov). We disclaim any intention or duty to update any forward-looking statement made in this news release.

CONTACT:
Caroline Loewy
Chief Financial Officer
Corcept Therapeutics
650-688-8783
cloewy@corcept.com
www.corcept.com