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): **October 10, 2012**

Corcept Therapeutics Incorporated

(Exact name of registrant as specified in its charter)

000-50679
(Commission File Number)

Delaware	000-50679	77-0487658
(State or other jurisdiction of incorporation)	(Commission File Number)	(I.R.S. Employer Identification No.)

149 Commonwealth Drive
Menlo Park, CA 94025
(Address of principal executive offices, with zip code)

(650) 327-3270
(Registrant's telephone number, including area code)

(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):

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 5.02 Departure of Directors or Certain Officers; Election of Directors; Appointment of Certain Officers; Compensatory Arrangements of Certain Officers

On October 10, 2012, the Board of Directors, or Board, of Corcept Therapeutics Incorporated, or the Company, elected Daniel M. Bradbury to serve as a director of the Company until the next annual meeting. Mr. Bradbury was also appointed to the Compensation Committee of the Board. Mr. Bradbury is an independent director under the criteria established by the Nasdaq Stock Market, and qualifies as a "non-employee director" for purposes of Rule 16b-3 under the Securities Exchange Act of 1934, as amended, and as an "outside director" for purposes of Section 162(m) of the Internal Revenue Code of 1986, as amended. A copy of the press release announcing Mr. Bradbury's appointment to the Board is attached hereto as Exhibit 99.1.

In connection with Mr. Bradbury's appointment as a director and member of the Compensation Committee, he will receive cash compensation in the amount of $15,000 per year in accordance with the Company's standard practices for non-employee director compensation. In addition, in accordance with Company practice for option grants to a new director, on October 10, 2012, Mr. Bradbury was granted an option to purchase 70,000 shares of the Company's common stock, at an exercise price of $2.70 per share, the closing price of the Company's common stock on the Nasdaq Stock Market on the date of grant. This option will vest over a 4-year period with 25% vesting on the first annual anniversary of the date of grant and the remainder vesting at the rate of 2.08334% on each monthly anniversary thereafter, until fully vested, subject to Mr. Bradbury's continued service. The option will expire 10 years from date of grant.

In connection with Mr. Bradbury's appointment, Mr. Bradbury and the Company also entered into an Indemnification Agreement in the same form as has previously been entered into with the Company's other current directors. The Indemnification Agreement generally requires the Company to indemnify Mr. Bradbury against liabilities incurred in the performance of his duties to the Company to the maximum extent permitted by applicable law. The foregoing description is qualified in its entirety by reference to the Company's standard form of Indemnification Agreement, which was filed as Exhibit 10.7 to its Quarterly Report on Form 10-Q filed on November 14, 2007.

Item 9.01. Financial Statements and Exhibits

(d) Exhibits.

99.1 Press Release of Corcept Therapeutics Incorporated dated October 15, 2012

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.

CORCEPT THERAPEUTICS INCORPORATED

Date: October 16, 2012 By: /s/ G. Charles Robb
G. Charles Robb
Chief Financial Officer

EXHIBIT INDEX

Exhibit Number	Description
99.1	Press Release of Corcept Therapeutics Incorporated dated October 15, 2012

Exhibit 99.1

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CORCEPT THERAPEUTICS APPOINTS DANIEL M. BRADBURY TO BOARD OF DIRECTORS

MENLO PARK, Calif., (October 15, 2012) -- Corcept Therapeutics Incorporated (NASDAQ: CORT), a pharmaceutical company engaged in the discovery, development and commercialization of drugs for the treatment of severe metabolic and psychiatric disorders, announced today that Daniel M. Bradbury has been appointed to the Company's Board of Directors.

"We are pleased that Dan has joined our Board," said Joseph K. Belanoff, M.D., Corcept's Chief Executive Officer. "His long experience with every aspect of the biotech industry, from drug discovery through approval and product launch, will be invaluable as we commercialize Korlym™ for Cushing's syndrome, continue our Phase 3 trial of mifepristone for the treatment of the psychotic features of psychotic depression and pursue development of our next-generation GR-II antagonists."

Mr. Bradbury's career in the biotechnology and pharmaceutical industry spans 30 years. He served as Chief Executive Officer of Amylin Pharmaceuticals, Inc. from March 2007 until its acquisition by Bristol-Myers Squibb Company in August 2012. From June 2006 until August 2012 he was a member of Amylin's board of directors and served on its Finance and Risk Management Committee. Mr. Bradbury also served as Amylin's President (2006-2007), Chief Operating Officer (2003-2006) and Executive Vice President (2000-2003). From 1994-2003 he held a variety of sales and marketing positions at the company. Before joining Amylin, Mr. Bradbury worked in marketing and sales roles for ten years at SmithKline Beecham Pharmaceuticals. He also serves on the board of directors of Illumina, Inc., Geron Corporation and Castle Biosciences, Inc., the board of trustees of the Keck Graduate Institute, the Investor Growth Capital Advisory Board, and the BioMed Ventures Advisory Committee.

Mr. Bradbury currently serves on the University of California San Diego's Rady School of Management's Advisory Council, the RAND Health Board of Advisors and the University of Miami's Innovation Corporate Advisory Council. He received a Bachelor of Pharmacy from Nottingham University and a Diploma in Management Studies from Harrow and Ealing Colleges of Higher Education in the United Kingdom.

"This is an exciting time for Corcept," said Mr. Bradbury. "Its platform of cortisol receptor antagonists, of which I believe Korlym for endogenous Cushing's syndrome is just the first, could generate treatments for many severe illnesses." He added, "I look forward to helping bring those medicines to patients."

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 and is generated by tumors that produce cortisol or ACTH. Cushing's syndrome is an orphan indication that 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 annually 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 Korlym™ (mifepristone) 300 mg Tablets

Korlym blocks the glucocorticoid receptor type II (GR-II) to which cortisol normally binds, thereby inhibiting the effects of excess cortisol in Cushing's syndrome patients. In April 2012, Corcept made Korlym available as a once-daily oral treatment of hyperglycemia secondary to endogenous Cushing's syndrome in adult patients with glucose intolerance or diabetes mellitus type 2 who have failed surgery or are not candidates for surgery. Korlym is the first and only FDA-approved treatment for that illness and the FDA has designated it as an Orphan Drug for that indication. Orphan Drug designation is a special status designed to encourage the development of medicines for rare diseases and conditions. Because Korlym is an Orphan Drug, Corcept will have marketing exclusivity for the approved indication in the United States until February 2019.

About Psychotic Depression

Psychotic depression is a serious psychiatric disorder that affects approximately three million people annually in the United States. It is more prevalent than either schizophrenia or bipolar I disorder. The disorder is characterized by severe depression accompanied by delusions, hallucinations or both. People with psychotic depression are approximately 70 times more likely to commit suicide than the general population and often require lengthy and expensive hospital stays. There is no FDA-approved treatment for psychotic depression.

About Corcept Therapeutics Incorporated

Corcept is a pharmaceutical company engaged in the discovery, development and commercialization of drugs for the treatment of severe metabolic and psychiatric disorders. Korlym, a first generation GR-II antagonist, is the company's first FDA-approved medication. The company owns extensive intellectual property covering the use of GR-II antagonists, including mifepristone, in the treatment of a wide variety of metabolic and psychiatric disorders. It also holds composition of matter patents for its selective GR-II antagonists.

Statements made in this news release, other than statements of historical fact, are forward-looking statements within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934. Forward-looking statements are subject to a number of known and unknown risks and uncertainties that might cause results to differ materially from those expressed or implied by such statements. For example, there can be no assurances regarding the amount of Corcept's revenues from Korlym or any other source, Korlym's acceptance by physicians and patients, the reimbursement decisions of government or private insurances, the FDA's response to any of the company's future submissions, the effects of rapid technological change and competition, the protections afforded by Corcept's intellectual property rights, or the cost, pace and success of Corcept's commercialization of Korlym for Cushing's syndrome or the company's product development efforts. These and other risks are set forth in the company's SEC filings, all of which are available from the company's website (**http://www.corcept.com**) or from the SEC's website (**http://www.sec.gov**). Corcept disclaims any duty to update any forward-looking statement made in this news release.