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: November 04, 2014 (Date of earliest event reported)

Corcept Therapeutics

(Exact name of registrant as specified in its charter)

DE (State or other jurisdiction of incorporation)

000-50679 (Commission File Number) **77-0487658** (IRS Employer Identification Number)

149 Commonwealth, Menlo Park CA (Address of principal executive offices)

94025 (Zip Code)

(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 2.02. Results of Operations and Financial Condition

Item 7.01. Regulation FD Disclosure

On November 4, 2014, Corcept Therapeutics Incorporated (the "Company") issued a press release announcing its financial results for the quarter ended September 30, 2014 and updating its revenue guidance for the year ending December 31, 2014. The press release is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

The information in this Item 2.02 and Item 7.01 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 or Sections 11 and 12(a)(2) of the Securities Act of 1933, as amended. The information in this Item 2.02 and the information contained in the press release attached as Exhibit 99.1 is not incorporated by reference into any filing with the U.S. Securities and Exchange Commission made by the Company, whether made before or after the date hereof, regardless of any general incorporation language in the filing unless specifically stated so therein.

Item 9.01. Financial Statements and Exhibits

(d) Exhibits

99.1 Press Release of Corcept Therapeutics dated November 04, 2014

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: November 10, 2014

CORCEPT THERAPEUTICS

By: <u>/s/ G. Charles Robb</u>
G. Charles Robb
Chief Financial Officer

Exhibit Index

Exhibit No.

Description

99.1

Press Release of Corcept Therapeutics dated November 04, 2014

Corcept Therapeutics Announces Third Quarter 2014 Financial Results and Provides Corporate Update

Revenue Increases 177 Percent From the Comparable Period in Prior Year and 24 Percent From Prior Quarter on Continued Uptake of Korlym(R) for the Treatment of Cushing's Syndrome; GAAP Net Loss Decreases to \$0.06 per Share; Non-GAAP Net Loss Decreases to \$0.04 per Share; Company Tightens 2014 Revenue Guidance Range to \$25-27 Million; Phase 1/2 Trial of Korlym for Treatment of Triple-Negative Breast Cancer to Generate Results in 2015; First Patients Enrolled in Phase 1 Clinical Trial of Next-Generation Selective GR Antagonist

MENLO PARK, CA -- (Marketwired - November 04, 2014) - Corcept Therapeutics Incorporated (NASDAQ: CORT), a pharmaceutical company engaged in the discovery, development and commercialization of drugs for the treatment of severe metabolic, oncologic and psychiatric disorders, today reported its financial results for the quarter ended September 30, 2014. The company also provided an update on its clinical programs and revised its 2014 revenue guidance.

Third Quarter Financial Results and 2014 Revenue Guidance

- Corcept recognized \$7.3 million in net revenue for the third quarter of 2014 compared to \$5.9 million in the second quarter, an increase of 24 percent. The company's net loss in the third quarter on a GAAP basis was \$6.0 million, or \$0.06 per share, compared to a net loss of \$7.6 million, or \$0.07 per share, in the second quarter.
- The company's net loss on a GAAP basis included significant non-cash items of \$2.1 million in the third quarter and \$2.2 million in the second quarter. Excluding these items, the company's net loss on a non-GAAP basis was \$3.9 million, or \$0.04 per share, for the third quarter of 2014 and \$5.4 million or \$0.05 per share, for the second quarter.
- As of September 30, 2014, the company held cash and cash equivalents of \$26.8 million.
- The company revised the range of its 2014 revenue guidance to \$25-27 million.

"Our Cushing's syndrome commercial business continues to grow and now generates cash to fund an increasing portion of our research and development activities," said Joseph K. Belanoff, M.D., Corcept's Chief Executive Officer. "While there is a group of endocrinologists who have seen Korlym's benefits and have become repeat prescribers, there are many physicians who have not yet written their first prescription. These physicians represent the potential for better treatment for more patients and significant future revenues."

Clinical Pipeline Progress

"We expect to have identified tolerable doses of Korlym and Halaven® (eribulin) in our triple-negative breast cancer study by yearend, when the study's efficacy phase is scheduled to begin. In September, we began dosing patients in our Phase 1 trial of CORT 125134. We will advance additional selective glucocorticoid receptor (GR) antagonists into the clinic next year. GR antagonism is an exciting area of research that will generate interesting data in 2015, both from our own studies and from those being undertaken by our academic collaborators."

Financial Results

For the third quarter of 2014, Corcept recognized net product revenue of \$7.3 million. The company reported a net loss of \$6.0 million, or \$0.06 per share, for the third quarter of 2014 compared to a net loss of \$10.9 million, or \$0.11 per share, for the same period in 2013.

The net loss on a GAAP basis for the third quarter of 2014 and for the third quarter of 2013 included non-cash stock-based compensation expenses of \$1.2 million and \$1.3 million, respectively. The company also recognized non-cash interest expense related to its capped royalty financing transaction of \$895,000 in the third quarter of 2014 and \$1.1 million in the same period in 2013.

After adjusting for these items, the company's net loss on a non-GAAP basis was \$3.9 million, or \$0.04 per share, for the third quarter of 2014, compared to \$8.5 million, or \$0.08 per share, for the third quarter of 2013. A reconciliation of GAAP net loss to non-GAAP net loss is included below.

Operating expenses were \$12.4 million for the third quarter of 2014 and also for the corresponding period in 2013.

- Selling, general and administrative expenses in the third quarter were \$9.1 million compared to \$7.2 million for the comparable period in 2013.
- Research and development expenses in the third quarter of 2014 were \$3.0 million, compared to \$5.2 million for the third quarter of 2013.

Corcept's cash balance as of September 30, 2014 was \$26.8 million, compared to \$54.9 million at December 31, 2013. Net cash used in the company's operating activities for the third quarter of 2014 was \$5.9 million as compared to \$9.0 million for the second quarter of 2014 and \$11.1 million for the first quarter of 2014.

Conference Call

Corcept will hold a conference call on November 4, 2014, at 5:00 p.m. Eastern Time (2:00 p.m. Pacific Time) to discuss this announcement. To participate, dial 1-888-771-4371 from the United States or +1-847-585-4405 internationally approximately ten minutes before the start of the call. The passcode is 38270277.

A replay will be available through November 18, 2014 at 1-888-843-7419 from the United States and +1-630-652-3042 internationally. The passcode is 38270277.

About Korlym®

Korlym competitively blocks the glucocorticoid receptor type II (GR), one of the two receptors to which cortisol normally binds, thereby inhibiting the effects of excess cortisol in patients with Cushing's syndrome. 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 was the first FDA-approved treatment for that illness and the FDA has designated it as an Orphan Drug for that indication.

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 20-50. An estimated 10-15 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 Triple-Negative Breast Cancer

Triple-negative breast cancer is a form of the disease in which the three receptors that fuel most breast cancer growth -- estrogen, progesterone and the HER-2/neu gene -- are not present. Because the tumor cells lack the necessary receptors, treatments that target estrogen, progesterone and HER-2 receptors are ineffective. In 2013, approximately 40,000 women were diagnosed with triple-negative breast cancer. There is no FDA-approved treatment and neither a targeted treatment nor an approved standard chemotherapy regimen for relapsed triple-negative breast cancer patients exists.

About CORT 125134

CORT 125134 is a potent, competitive antagonist at the GR-II receptor, but does not have affinity for the progesterone, estrogen, AR androgen or GR-I (mineralocorticoid) receptors. The company has begun a Phase 1 study of the safety and tolerability of CORT 125134, one of its next-generation selective GR-II antagonists.

About Corcept Therapeutics Incorporated

Corcept is a pharmaceutical company engaged in the discovery, development and commercialization of drugs for the treatment of severe metabolic, oncologic and psychiatric disorders. Korlym, a first generation competitive GR antagonist, is the company's first FDA-approved medication. The company has a Phase 1 trial of mifepristone for the treatment of triple-negative breast cancer, a Phase 1 trial of CORT 125134, one of its proprietary, selective GR-II antagonists, and a portfolio of other selective GR antagonists that competitively block the effects of cortisol but not progesterone. It owns extensive intellectual property covering the use of GR antagonists, including mifepristone, in the treatment of a wide variety of metabolic, oncologic and psychiatric disorders. It also holds composition of matter patents for its selective GR antagonists.

Non-GAAP Measures of Net Loss

To supplement Corcept's financial results presented on a GAAP basis, the company uses non-GAAP measures of net loss that exclude significant non-cash expenses related to stock-based compensation expense and the recognition of interest expense under the company's capped royalty financing transaction. We believe that this non-GAAP measure of net loss helps investors better evaluate the company's past financial performance and potential future results. Non-GAAP measures should not be considered in isolation or as a substitute for comparable GAAP accounting and investors should read them in conjunction with the company's financial statements prepared in accordance with GAAP. The non-GAAP measure of net loss we use may be different from, and not directly comparable to, similarly titled measures used by other companies.

Forward-Looking Statements

Statements made in this news release, other than statements of historical fact, are forward-looking statements. These forward-looking statements, including statements regarding anticipated future revenues, the timing of clinical trials and clinical trial results, clinical pipeline progress and the advancement of additional compounds, are subject to known and unknown risks and uncertainties that might cause actual results to differ materially from those expressed or implied by such statements, including the pace of Korlym's acceptance by physicians and patients, the reimbursement decisions of government or private insurers, the pace of

enrollment in or the outcome of the company's Phase 1/2 study of mifepristone in the treatment of triple-negative breast cancer and the Phase 1 study of its next-generation selective GR antagonist, CORT 125134, the effects of rapid technological change and competition, the protections afforded by Korlym's Orphan Drug designation or by Corcept's other intellectual property rights, or the cost, pace and success of Corcept's other 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 intention or duty to update any forward-looking statement made in this news release.

Halaven® is a registered trademark used by Eisai Inc. under license from Eisai R&D Management Co., Ltd.

CORCEPT THERAPEUTICS INCORPORATED CONDENSED BALANCE SHEETS (in thousands)

	September 30, 2014 (Unaudited)		Dec	cember 31, 2013
			(Note)	
ASSETS: Cash and cash equivalents Trade receivables, net Inventory Other assets	\$	26,780 2,746 5,510 2,123		54,877 1,428 5,546 1,226
Total assets	\$ ====	37,159 =======	\$ ====	63,077
LIABILITIES AND STOCKHOLDERS' EQUITY: Accounts payable Deferred revenue Long-term obligation Other liabilities Stockholders' equity	\$	1,960 50 34,651 1,815 (1,317)		2,381 25 35,065 4,589 21,017
Total liabilities and stockholders' equity	\$ =====	37,159 ======	\$ ====	63,077

Note: Derived from audited financial statements at that date.

CORCEPT THERAPEUTICS INCORPORATED CONDENSED STATEMENTS OF OPERATIONS (in thousands, except per share amounts)

(Unaudited)

	Three Months Ended September 30,							
		2014		2013	-	2014		2013
Revenues: Product sales, net	\$	7,282	\$	2,634	\$	17,537	\$	6,242
Operating expenses: Cost of sales Research and development Selling, general and administrative		3,047		5,155		624 14,583 26,872		13,903
Total operating expenses					-	42,079		
Loss from operations		(5,103)		(9,740)		(24,542)		(31,466)
Interest and other expense		(903)		(1,166)	_	(2,946)		(3,421)
Net loss and comprehensive loss	\$ ==	(6,006) =====	\$	(10,906) ======	\$	(27,488) ======	\$	(34,887) ======
Basic and diluted net loss per share	\$	(0.06)	\$	(0.11)	\$	(0.27)	\$	(0.35)

Shares used in computing basis					
Shares used in computing basic and diluted net loss per share	101,134	99,814	100,880	99,814	
	=======	=======	=======	=======	

CORCEPT THERAPEUTICS INCORPORATED RECONCILIATION OF GAAP TO NON-GAAP NET LOSS (in thousands, except per share amounts)

(Unaudited)

	Septem		Nine Months Ended September 30,		
	2014	2013	2014	2013	
GAAP net loss	\$ (6,006)	\$ (10,906)	\$ (27,488)	\$ (34,887)	
Significant non-cash expenses: Stock-based compensation Research and development Selling, general and administrative		161 1,148	3,300	3,417	
Total stock-based compensation		1,309		3,883	
Accretion of interest expense related to long-term obligation Non-GAAP net loss	895	1,133 \$ (8,464)	2,874	3,340	
6.4 2000		=======			
GAAP basic and diluted net loss per share	, ,	\$ (0.11) =======	` ,	,	
Non-GAAP basic and diluted net loss per share as adjusted for significant non-cash expenses		\$ (0.08) =====			
Shares used in computing basic and diluted net loss per share	•	99,814	•	99,814	

CONTACT:

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