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 05, 2015
(Date of earliest event reported)

Corcept Therapeutics
(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, 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 5, 2015, Corcept Therapeutics Incorporated (the "Company") issued a press release announcing its financial results for the quarter ended September 30, 2015. 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 Item 7.01 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 05, 2015

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: November 05, 2015

CORCEPT THERAPEUTICS

By: /s/ G. Charles Robb
G. Charles Robb
Chief Financial Officer

Exhibit Index

Exhibit No.	**Description**
99.1	Press Release of Corcept Therapeutics dated November 05, 2015

Corcept Therapeutics Announces Third Quarter 2015 Results and Provides Corporate Update

MENLO PARK, CA -- (Marketwired - November 05, 2015) - Corcept Therapeutics Incorporated (NASDAQ: CORT)

- **GAAP net loss in the third quarter narrows to $0.6 million, from $6.0 million in the third quarter of 2014**
- **Non-GAAP net income in the third quarter increases to $1.6 million, compared to a non-GAAP net loss of $3.9 million in the third quarter of 2014**
- **Third quarter revenue grows to $13.3 million, compared to $7.3 million in the third quarter of 2014, an 82 percent increase; the company reiterates its 2015 revenue guidance of $49 - $53 million**
- **Preliminary efficacy results of the company's Phase 1/2 trial of Korlym® to treat triple-negative breast cancer will be presented at the San Antonio Breast Cancer Symposium on December 10, 2015**
- **Phase 2 trials of next-generation selective cortisol modulator CORT125134, in Cushing's syndrome and an oncology indication, planned to begin in the first quarter of 2016**

Corcept Therapeutics Incorporated (NASDAQ: CORT), a pharmaceutical company engaged in the discovery, development and commercialization of drugs that treat severe metabolic, oncologic and psychiatric disorders by modulating the effects of cortisol, today reported its financial results for the quarter ended September 30, 2015 and provided a corporate update.

Corcept recorded net revenue of $13.3 million in the third quarter of 2015, compared to $7.3 million for the same period in 2014, an increase of 82 percent. The company reiterated its 2015 revenue guidance of $49 - $53 million.

Corcept reported a net loss on a GAAP basis of $0.6 million for the third quarter of 2015, compared to a GAAP net loss of $6.0 million for the same period in 2014. Excluding non-cash expenses, Corcept generated non-GAAP net income in the third quarter of $1.6 million, compared to a non-GAAP net loss of $3.9 million in the third quarter of 2014. A reconciliation of GAAP to non-GAAP net operating results is set forth below.

At September 30, 2015, Corcept held cash and cash equivalents of $36.5 million, compared to $37.0 million at the end of the prior quarter. This change in cash reflects the repayment of $1.8 million in principal due under the company's capped royalty financing agreement (Royalty Financing). The company entered into the Royalty Financing in 2012 to fund the commercialization of Korlym® and expects to make its final payment in 2017.

Based on its current plans, the company expects to reach cash flow breakeven, including payments of principal due under the Royalty Financing, without needing to raise additional funds.

"Our Cushing's syndrome franchise continues to grow as more physicians appreciate that using Korlym to modulate the effect of excess cortisol can greatly improve their patients' health," said Joseph K. Belanoff, M.D., Corcept's Chief Executive Officer. "As we have said before, our efficient cost structure and the revenue growth in our Cushing's syndrome business allows us to build our clinical infrastructure, further develop Korlym and advance our next-generation compounds."

Korlym for the Treatment of Triple-Negative Breast Cancer (TNBC)

In December 2015, at the San Antonio Breast Cancer Symposium, Corcept will present preliminary efficacy results of its Phase 1/2 open-label trial of Korlym in combination with eribulin (Halaven®) to treat patients with metastatic TNBC.

CORT125134 Phase 2 Trials

CORT125134 is the lead compound in Corcept's portfolio of proprietary next-generation cortisol modulators. It was well-tolerated in its Phase 1 trial and showed that it shares Korlym's ability to potently modulate activity at the glucocorticoid receptor (GR), the essential quality in treating Cushing's syndrome. In addition, when administered with a chemotherapeutic agent, CORT125134 slows tumor growth significantly in mouse models of TNBC and castration-resistant prostate cancer. In vitro, it similarly slows the growth of ovarian cancer tumor cells; in vivo testing in mouse models of ovarian cancer is in progress.

In the first quarter of 2016, the company plans to begin two Phase 2 clinical trials with CORT125134. One trial will be for the treatment of patients with Cushing's syndrome. The other will administer CORT125134 with a companion chemotherapeutic or hormonal agent to treat patients with a solid tumor cancer.

"It's an exciting time to join Corcept," said Robert S. Fishman, M.D., Corcept's Chief Medical Officer. "We are exploring extending Korlym's label to cover oncologic indications, our lead next-generation cortisol modulator, CORT125134, is about to enter multiple Phase 2 trials and we're advancing additional selective cortisol modulating compounds toward the clinic."

Financial Discussion

Corcept recorded a GAAP net loss of $0.6 million in the third quarter of 2015, compared to $6.0 million in the third quarter of 2014, including non-cash stock-based compensation and accreted interest expense generated from the Royalty Financing of $2.2 million and $2.1 million in the third quarter of 2015 and 2014, respectively. Excluding these non-cash items, Corcept generated net income on a non-GAAP basis of $1.6 million in the third quarter of 2015, compared to a non-GAAP net loss of $3.9 million in the third quarter of 2014.

Operating expenses for the third quarter were $13.2 million, compared to $12.4 million for the third quarter of 2014. The increase was primarily due to increased staffing costs and additional spending on the Phase 1/2 trial of Korlym for the treatment of TNBC

and the development of next-generation cortisol modulators.

In the third quarter of 2015, Corcept made a payment of $2.5 million under the Royalty Financing, of which $0.7 million represented accreted interest and $1.8 million reduced outstanding principal. In the same period of 2014, Corcept paid $1.3 million, of which $0.9 million represented accreted interest and $0.4 million reduced outstanding principal. Corcept expects to make its final payment under the Royalty Financing in 2017.

Corcept's cash balance at September 30, 2015 was $36.5 million, compared to $37.0 million at June 30, 2015 and $24.2 million at December 31, 2014.

Conference Call

Corcept will hold a conference call on November 5, 2015, at 5:00 p.m. Eastern Time (2:00 p.m. Pacific Time), to discuss this announcement. To participate, dial 1-888-771-4371 in the United States or 1-847-585-4405 internationally approximately ten minutes before the start of the call. The passcode is 41022099. A replay will be available through November 19, 2015 at 1-888-843-7419 in the United States and 1-630-652-3042 internationally. The passcode will be 41022099.

About Korlym

Korlym modulates the effects of excess cortisol in patients with Cushing's syndrome. Since 2012, Corcept has marketed Korlym 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 Korlym 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, half of whom are cured by surgery. 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 HER-2 -- are not present. Because the tumor cells lack these receptors, treatments that target estrogen, progesterone and HER-2 are ineffective. Approximately 40,000 women are diagnosed with triple-negative breast cancer each year. It is estimated that more than 75 percent of these women's tumor cells express GR. 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 CORT125134

CORT125134 is one of Corcept's next-generation selective cortisol modulators. It is a potent, competitive modulator of GR, but does not have affinity for the progesterone, estrogen, androgen or mineralocorticoid receptors. The company has completed a Phase 1 study of the safety and tolerability of CORT125134 and plans to begin Phase 2 trials of the compound as a treatment for Cushing's syndrome and an oncology indication early in 2016.

About Corcept Therapeutics Incorporated

Corcept discovers, develops and commercializes drugs that treat severe metabolic, oncologic and psychiatric disorders by modulating the effects of cortisol. Korlym is the company's first FDA-approved medication. The company is conducting a Phase 1/2 trial of Korlym for the treatment of triple-negative breast cancer and is planning Phase 2 trials of CORT125134, one of its next-generation selective cortisol modulators, for the treatment of Cushing's syndrome and an oncology indication. The company has developed a proprietary portfolio of other compounds that modulate the effects of cortisol, but not progesterone. It holds composition of matter patents on these molecules. Corcept also owns or has exclusively licensed extensive intellectual property covering the use of cortisol modulators, including Korlym, in the treatment of a wide variety of serious disorders.

Non-GAAP Measures of Net Income / (Loss)

To supplement Corcept's financial results presented on a GAAP basis, the company uses non-GAAP measures of net income and net loss that exclude non-cash expenses related to stock-based compensation expense and the accretion of interest expense under the Royalty Financing. The company believes that these non-GAAP measures help 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 measures of net income / (loss) the company uses may be different from, and not directly comparable to, similarly titled measures used by other companies.

Forward-Looking Statements

Statements made in this press release, other than statements of historical fact, are forward-looking statements that 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. Forward-looking statements include statements regarding anticipated revenues and scheduled debt payments for 2015 and beyond, the company's expectation to reach cash flow breakeven, growth of the company's Cushing syndrome franchise, the timing of clinical trials and clinical trial results and the company's presentation of preliminary efficacy results at the San Antonio Breast Cancer Symposium, the pace of Korlym's acceptance by physicians and patients, the pace of enrollment in or the outcome of the company's clinical trials and the advancement of its next-generation selective cortisol modulators, 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, and 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, which are available at the company's website (www.corcept.com) or from the SEC's website (www.sec.gov). Corcept disclaims any intention or duty to update any forward-looking statement made in this press 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, 2015	December 31, 2014
	(Unaudited)	(Note)
ASSETS:		
Cash and cash equivalents	$ 36,450	$ 24,248
Trade receivables	5,945	3,334
Inventory	4,594	5,297
Other assets	1,388	1,751
Total assets	$ 48,377	$ 34,630
LIABILITIES AND STOCKHOLDERS' EQUITY:		
Accounts payable	$ 1,740	$ 1,886
Capped royalty financing obligation	29,640	33,887
Other liabilities	3,520	2,245
Stockholders' equity (deficit)	13,477	(3,388)
Total liabilities and stockholders' equity	$ 48,377	$ 34,630

Note: Derived from audited financial statements at that date.

CORCEPT THERAPEUTICS INCORPORATED
CONDENSED STATEMENTS OF COMPREHENSIVE INCOME / (LOSS)
(in thousands, except per share amounts)

(Unaudited)

	Three Months Ended September 30,		Nine Months Ended September 30,	
	2015	2014	2015	2014
Revenues:				
Product sales, net	$ 13,261	$ 7,282	$ 35,319	$ 17,537
Operating expenses:				
Cost of sales	256	235	997	624
Research and development	3,612	3,047	11,330	14,583
Selling, general and administrative	9,291	9,103	28,086	26,872
Total operating expenses	13,159	12,385	40,413	42,079
Income / (Loss) from operations	102	(5,103)	(5,094)	(24,542)
Interest and other expense	(703)	(903)	(2,273)	(2,946)

Net income / (loss) and comprehensive loss	$ (601)	$ (6,006)	$ (7,367)	$ (27,488)
Basic and diluted net income / (loss) per share	$ (0.01)	$ (0.06)	$ (0.07)	$ (0.27)
Shares used in computing basic and diluted net income / (loss) per share	108,461	101,134	106,104	100,880

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

(Unaudited)

	Three Months Ended September 30,		Nine Months Ended September 30,	
	2015	2014	2015	2014
GAAP net income / (loss)	$ (601)	$ (6,006)	$ (7,367)	$ (27,488)
Non-cash expenses:				
Stock-based compensation				
Research and development	196	183	579	514
Selling, general and administrative	1,346	1,027	3,941	3,300
Total stock-based compensation	1,542	1,210	4,520	3,814
Accretion of interest expense related to Capped royalty financing obligation	698	895	2,196	2,874
Non-GAAP net income / (loss)	$ 1,639	$ (3,901)	$ (651)	$ (20,800)
GAAP basic and diluted net income / (loss) per share	$ (0.01)	$ (0.06)	$ (0.07)	$ (0.27)
Non-GAAP basic and diluted net income / (loss) per share excluding non-cash expenses	$ 0.02	$ (0.04)	$ (0.01)	$ (0.21)
Shares used in computing basic and diluted net income / (loss) per share	108,461	101,134	106,104	100,880

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