THERAVANCE INC
Form 8-K
April 17, 2013

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: April 17, 2013
(Date of earliest event reported)

Theravance, Inc.
(Exact name of registrant as specified in its charter)
Delaware
(State or other jurisdiction
of incorporation) 000-30319
(Commission File Number) 94-3265960
(IRS Employer
Identification Number)
901 Gateway Boulevard, South San Francisco, CA
(Address of principal executive offices) 94080
(Zip Code)
650-808-6000
(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

The information contained in this Item 7.01 and in the accompanying exhibit shall not be deemed filed for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Securities Exchange Act of 1934"), or incorporated by reference in any filing under the Exchange Act or the Securities Act of 1933, as amended, except as shall be expressly set forth by specific reference in such filing.

On April 17, 2013 GlaxoSmithKline plc (GSK) and Theravance, Inc. issued a press release announcing the outcome of the meeting of the Pulmonary-Allergy Drugs Advisory Committee to the U.S. Food and Drug Administration regarding fluticasone furoate (FF) and vilanterol (VI) dry powder inhaler (proposed trade name BREO(TM) ELLIPTA(TM)) for the long-term maintenance treatment of airflow obstruction and for reducing exacerbations in patients with chronic obstructive pulmonary disease. FF/VI, an investigational once-daily inhaled corticosteroid/long-acting beta2 agonist (LABA) combination treatment, is in development under the LABA collaboration between GSK and Theravance, Inc. The press release is attached hereto as Exhibit 99.1 and incorporated herein by reference.

Item 9.01. Financial Statements and Exhibits

(d) Exhibits

99.1 Press Release dated April 17, 2013

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: April 17, 2013
THERAVANCE, INC.

By: /s/ Michael W. Aguiar
Michael W. Aguiar
Chief Financial Officer

Exhibit Index **Exhibit No.** **Description** 99.1 Press Release dated April 17, 2013