THERAVANCE INC
Form 8-K
July 07, 2014

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: July 07, 2014
(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)
951 Gateway Boulevard, South San Francisco, CA
(Address of principal executive offices) 94080
(Zip Code)
650-238-9600
(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 (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 8.01. Other Events

On July 7, 2014, GlaxoSmithKline plc (GSK) and Theravance, Inc. issued a press release announcing that the Japanese Ministry of Health, Labour and Welfare (MHLW) has approved ANORO(R) ELLIPTA(R) (umeclidinium/vilanterol) for the relief of various symptoms due to airway obstruction with chronic obstructive pulmonary diseases (chronic bronchitis, pulmonary emphysema) (in the case where concurrent use of long-acting inhaled muscarinic antagonist and long-acting inhaled beta2 agonist is required). ANORO(R) is a once-daily combination treatment comprised of two bronchodilators, umeclidinium (UMEC), a long-acting muscarinic antagonist (LAMA), and vilanterol (VI), a long-acting beta2 agonist (LABA), in a single inhaler, the ELLIPTA(R). The approved dose in Japan is UMEC/VI 62.5/25mcg delivered once daily. Under the terms of the 2002 LABA collaboration agreement, Theravance is obligated to make a milestone payment of $10 million (USD) to GSK following MHLW approval of UMEC/VI in Japan. UMEC/VI has been developed under the LABA collaboration agreement between Glaxo Group Limited and Theravance, Inc. A copy of the press release is filed as Exhibit 99.1 to this report and is incorporated herein by reference.

Item 9.01. Financial Statements and Exhibits

(d) Exhibits

99.1 Press Release dated July 07, 2014

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: July 07, 2014
THERAVANCE, INC.

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

Exhibit Index **Exhibit No.** **Description** 99.1 Press Release dated July 07, 2014