

THERAVANCE INC  
Form 8-K  
January 09, 2013

**UNITED STATES**

**SECURITIES AND EXCHANGE COMMISSION**

Washington, D.C. 20549

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**FORM 8-K**

**CURRENT REPORT**

Pursuant to Section 13 or 15(d) of the  
Securities Exchange Act of 1934

**Date of Report: January 09, 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:

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
- 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))

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**Item 8.01. Other Events**

On January 9, 2013, GlaxoSmithKline plc (GSK) and Theravance, Inc. (the "Company") issued a press release announcing that a Marketing Authorization Application (MAA) for UMEC/VI with the proposed proprietary name ANORO(TM) has been submitted to the European Medicines Agency (EMA), for the maintenance bronchodilator treatment to relieve symptoms in adult patients with chronic obstructive pulmonary disease. UMEC/VI is a combination of two investigational bronchodilator molecules - GSK573719 or umeclidinium bromide (UMEC), a long-acting muscarinic antagonist and vilanterol (VI), a long-acting beta2 agonist (LABA), administered once-daily using the ELLIPTA(TM) inhaler. UMEC/VI is currently in development under the LABA collaboration between GSK and the Company. 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 of Theravance, Inc. dated January 09, 2013

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**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: January 09, 2013

**THERAVANCE, INC.**

By: /s/ Michael W. Aguiar

Michael W. Aguiar

*Chief Financial Officer*

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**Exhibit Index** **Exhibit No. Description** 99.1 Press Release of Theravance, Inc. dated January 09, 2013