

THERAVANCE INC  
Form 8-K  
March 14, 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: March 14, 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)**  
**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 (see General Instruction A.2. below):

- 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))

---

**Item 8.01. Other Events**

On March 14, 2014, GlaxoSmithKline plc (GSK) and Theravance, Inc. issued a press release announcing positive results from three Phase 3 studies. Two studies comparing the efficacy and safety of the combination anticholinergic/long-acting beta2-adrenergic agonist, ANORO(TM) ELLIPTA(TM) (umeclidinium/vilanterol, 'UMEC/VI') with inhaled corticosteroid/long-acting beta2-adrenergic agonist combination, ADVAIR(R) DISKUS(R) (fluticasone propionate/salmeterol 'FSC 250/50') and the third comparing the efficacy and safety of ANORO(TM) ELLIPTA(TM) with SERETIDE(R) DISKUS(R) 'FSC 500/50' in patients with chronic obstructive pulmonary disease (COPD) and no history of moderate to severe COPD exacerbations in the last year. 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 March 14, 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: March 14, 2014  
**THERAVANCE, INC.**

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

---

**Exhibit Index** **Exhibit No. Description** 99.1 Press Release dated March 14, 2014