

THERAVANCE INC  
Form 8-K  
May 10, 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: May 10, 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))

---

**Item 8.01. Other Events**

On May 10, 2013, GlaxoSmithKline plc (GSK) and Theravance, Inc. issued a press release announcing that the U.S. Food and Drug Administration has approved BREO(TM) ELLIPTA(TM) as an inhaled long-term, once-daily maintenance treatment of airflow obstruction in patients with chronic obstructive pulmonary disease (COPD), including chronic bronchitis and/or emphysema. It is also indicated to reduce exacerbations of COPD in patients with a history of exacerbations. BREO(TM) ELLIPTA(TM) is not indicated for the relief of acute bronchospasm or the treatment of asthma. BREO(TM) ELLIPTA(TM), a combination of the inhaled corticosteroid, fluticasone furoate "FF", and the long-acting beta2 agonist (LABA), vilanterol "VI" (FF/VI 100/25 mcg), is in development under the LABA collaboration between GSK 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 May 10, 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: May 10, 2013  
**THERAVANCE, INC.**

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

---

**Exhibit Index** **Exhibit No. Description** 99.1 Press Release dated May 10, 2013