

THERAVANCE INC  
Form 8-K  
March 20, 2013

**UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION**

Washington, DC 20549

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**FORM 8-K**

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

Date of Report (Date of earliest event Reported): **March 20, 2013**

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**THERAVANCE, INC.**

(Exact Name of Registrant as Specified in its Charter)

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**Delaware**

(State or Other Jurisdiction of  
Incorporation)

**000-30319**

(Commission File Number)

**94-3265960**

(I.R.S. Employer Identification Number)

**901 Gateway Boulevard  
South San Francisco, California 94080  
(650) 808-6000**

## Edgar Filing: THERAVANCE INC - Form 8-K

(Addresses, including zip code, and telephone numbers, including area code, of principal executive offices)

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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))
  
  - 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 March 20, 2013, the Office of the Federal Register published an advanced display of the Federal Register Notice that on April 17, 2013 the U.S. Food and Drug Administration's Pulmonary-Allergy Drugs Advisory Committee will discuss the new drug application (NDA) 204275, for fluticasone furoate and vilanterol dry powder inhaler (proposed trade name BREO ELLIPTA<sup>®</sup>), sponsored by GlaxoSmithKline plc (GSK), for the long-term maintenance treatment of airflow obstruction and for reducing exacerbations in patients with chronic obstructive pulmonary disease. Fluticasone furoate and vilanterol, 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.

**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.

**THERAVANCE, INC.**

Date: March 20, 2013

By:

*/s/ Michael W. Aguiar*

**Michael W. Aguiar**  
**Chief Financial Officer**