

THERAVANCE INC  
Form 8-K  
April 22, 2014

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UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION  
Washington, DC 20549

FORM 8-K

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

Date of Report (Date of earliest event Reported): April 22, 2014

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 (I.R.S. Employer Identification Number)
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901 Gateway Boulevard  
South San Francisco, California 94080  
(650) 808-6000

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

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))



Item 8.01 Other Events.

On April 22, 2014, GlaxoSmithKline plc and Theravance, Inc. issued a press release announcing that the Therapeutic Goods Administration (TGA) has approved BREO™ ELLIPTA® (fluticasone furoate/vilanterol trifenate) for the treatment of patients with asthma or chronic obstructive pulmonary disease (COPD) in Australia. BREO™ ELLIPTA® is a combination of the inhaled corticosteroid, fluticasone furoate (FF), and the long-acting beta2-agonist (LABA), vilanterol (VI). Two strengths of BREO™ ELLIPTA® have been licensed for the treatment of asthma (100/25 mcg and 200/25 mcg) and one strength has been licensed for the treatment of COPD (100/25 mcg). Both strengths will be administered once-daily using the new ELLIPTA® dry powder inhaler. FF/VI has been developed under the 2002 LABA collaboration between Glaxo Group Limited and Theravance, Inc. 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.

Exhibit	Description
<u>Exhibit 99.1</u>	Press Release dated April 22, 2014

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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.

THERAVANCE, INC.

Date: April 22, 2014

By: /s/ Michael W. Aguiar

Michael W. Aguiar  
Chief Financial Officer

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EXHIBIT INDEX

Exhibit No.	Description
<u>99.1</u>	Press Release dated April 22, 2014

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