

THERAVANCE INC  
Form 8-K  
April 06, 2009

**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: April 06, 2009**  
**(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 7.01. Regulation FD Disclosure**

The information contained in this Item 7.01 and in the accompanying exhibit shall not be deemed filed for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or incorporated by reference in any filing under the Exchange Act or the Securities Act of 1933, as amended, except as shall be expressly set forth by specific reference in such filing.

On April 6, 2009 Theravance, Inc. announced that the U.S. Food and Drug Administration ("FDA") accepted for review its New Drug Application for telavancin, a novel, bactericidal, once-daily injectable investigational antibiotic, for the treatment of nosocomial pneumonia (also known as hospital-acquired pneumonia, or HAP) caused by Gram-positive bacteria such as methicillin-resistant Staphylococcus aureus (MRSA). This filing of the New Drug Application with the FDA triggers a \$10 million milestone payment from Theravance's partner Astellas Pharma Inc. A copy of the press release is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

**Item 9.01. Financial Statements and Exhibits**

**(d) Exhibits**

99.1 Press Release of Theravance, Inc. dated April 06, 2009

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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: April 06, 2009  
**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 April 06, 2009