THERAVANCE INC Form 8-K September 22, 2010

	UNITED STATES	
SECURITII	ES AND EXCHANGE C	OMMISSION
	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): September 1	ber 22, 2010
	THERAVANCE, INC.	
	(Exact Name of Registrant as Specified in its Cha	rter)
elaware	000-30319	94-3265960
ner Jurisdiction of rporation)	(Commission File Number)	(I.R.S. Employer Identification Number)
	001 Cataway Bardayard	

(State or Oth Incor

> 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):		
o	Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)	
o	Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)	
o	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.

Today at the European Respiratory Society Annual Congress in Barcelona, Spain, GlaxoSmithKline plc presented two oral presentations: Phase 2 study of RELOVAIR , a once-daily combination medicine of fluticasone furoate (FF), the inhaled corticosteroid (ICS), and lanterol trifenatate (VI), the long-acting beta2 agonist (LABA) in patients with chronic obstructive pulmonary disease (COPD) and Phase 2b study of VI in patients with asthma. RELOVAIR is being developed for the treatment of patients with COPD or asthma under the LABA collaboration between GSK and Theravance, Inc. The two presentations are filed as Exhibit 99.1 and Exhibit 99.2 to this report and are incorporated herein by reference.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

Exhibit Description
Exhibit 99.1 Safety and efficacy of fluticasone furoate/vilanterol trifenatate (FF/VI) in COPD patients
Exhibit 99.2 24h duration of the novel long-acting β2 agonist vilanterol trifenatate in uncontrolled asthma

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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: September 22, 2010 By: /s/ Michael W. Aguiar

Michael W. Aguiar Chief Financial Officer

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EXHIBIT INDEX

Exhibit	Description
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Exhibit 99.2	24h duration of the novel long-acting $\beta 2$ agonist vilanterol trifenatate in uncontrolled asthma
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