# 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): September 18, 2024

# THERAVANCE BIOPHARMA, INC.

(Exact Name of Registrant as Specified in its Charter)

Cayman Islands (State or Other Jurisdiction of Incorporation) 001-36033

(Commission File Number)

98-1226628 (I.R.S. Employer Identification Number)

c/o Theravance Biopharma US, Inc. 901 Gateway Boulevard South San Francisco, CA 94080 (650) 808-6000

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

following provisions (see General Instruction A.2. below):		e filling obligation of the registrant under any of the
Written communications pursuant to Rule 425 under to Soliciting material pursuant to Rule 14a-12 under the Pre-commencement communications pursuant to Rule Pre-commencement communications pursuant to Rule Pre-commencement communications pursuant to Rule	Exchange Act (17 CFR 240.14a-12) e 14d-2(b) under the Exchange Act (17 CF	\$ 77
Securities registered pursuant to Section 12(b) of the Act:		
Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Ordinary Share \$0.00001 Par Value	ТВРН	NASDAQ Global Market
Indicate by check mark whether the registrant is an emer chapter) or Rule 12b-2 of the Securities Exchange Act of 1		405 of the Securities Act of 1933 (§ 230.405 of this
		Emerging growth company $\Box$
If an emerging growth company, indicate by check mark is or revised financial accounting standards provided pursuant	•	

#### Item 8.01 Other Events

On September 18, 2024, Theravance Biopharma R&D IP, LLC, Theravance Biopharma US, Inc. and Theravance Biopharma Ireland Limited, subsidiaries of the Registrant Theravance Biopharma, Inc. (together, "Theravance"), and Mylan Ireland Limited and Mylan Specialty L.P. (together, "Mylan"), entered into a Settlement Agreement (the "Settlement Agreement") with Qilu Pharmaceutical Co., Ltd. and Qilu Pharma Inc. (together "Qilu") relating to Theravance and Mylan's YUPELRI® (revefenacin) inhalation solution. A Theravance entity owns and Mylan is the exclusive sub-licensee of United States Patent Nos. 8,541,451; 9,765,028; 10,550,081; 11,008,289; 11,484,531; 11,691,948; and 11,858,898 (the "Patents-in-Suit").

The Settlement Agreement resolves ongoing patent litigation brought by Theravance and Mylan against Qilu pursuant to the Hatch-Waxman Act based on Qilu's filing of an abbreviated new drug application ("ANDA") seeking approval to market a generic version of YUPELRI® (revefenacin) inhalation solution prior to expiration of the Patents-in-Suit.

Under the Settlement Agreement, Theravance and Mylan granted Qilu a royalty-free, non-exclusive, non-sublicensable, non-transferable license to manufacture and market Qilu's generic version of YUPELRI® (revefenacin) inhalation solution in the United States on or after the Licensed Launch Date of April 23, 2039, subject to certain exceptions as is customary in these type of agreements.

As required by law, the settlement is subject to review by the U.S. Department of Justice and the Federal Trade Commission.

The patent litigation previously disclosed by the Company remains pending against three other ANDA filers: Cipla Limited; Eugia Pharma Specialities Ltd.; and Mankind Pharma Ltd.; along with certain affiliates.

### **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 BIOPHARMA, INC.

Date: September 23, 2024 By: /s/ Brett Grimaud

Brett Grimaud General Counsel