

CURRENT REPORT
Pursuant to Section 13 OR 15(d) of The Securities Exchange Act of 1934

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☐

Item 7.01 Regulation FD Disclosure.

On June 26, 2026, Cumberland Pharmaceuticals Inc. (the “Company” or “Cumberland”) presented data from its Phase 2 FIGHT DMD trial evaluating ifetroban, a novel oral therapy for Duchenne muscular dystrophy (DMD) heart disease, at the annual Parent Project Muscular Dystrophy conference in Orlando, Florida. The presentation included a slide indicating that long-term safety and efficacy data from the ongoing open-label extension was expected in August 2026. The Company is providing this update to note that such data is not yet available and, accordingly, the Company no longer expects to present this data in August 2026. The Company intends to present this information at a future date when it becomes available.

Forward-Looking Statements

This Form 8-K contains forward-looking statements, which are subject to certain risks and reflect Cumberland’s current views on future events based on what it believes are reasonable assumptions. No assurance can be given that these events will occur. Forward-looking statements include, among other things, statements regarding the Company’s intent, belief or expectations, and can be identified by the use of terminology such as “may,” “will,” “expect,” “believe,” “intend,” “plan,” “estimate,” “goal,” “should,” “seek,” “anticipate,” “look forward” and other comparable terms or the negative thereof. As with any business, all phases of Cumberland’s operations are subject to factors outside of its control, and any one or combination of these factors could materially affect Cumberland’s operation results. These factors include risks and uncertainties related to the strategic transaction, risks related to the Company’s ability to develop its pipeline of new product candidates, the risks inherent in clinical development and regulatory review, macroeconomic conditions, including changes in interest rates, inflation, tariffs, competition and other events beyond the Company’s control as more fully discussed in its most recent quarterly report on Form 10-Q as filed with the U.S. Securities and Exchange Commission (“SEC”), as well as the Company’s other filings with the SEC from time to time. There can be no assurance that results anticipated by the Company will be realized or that they will have the expected effects. Readers are cautioned not to place undue reliance on forward-looking statements, which speak only as of the date hereof. The Company does not undertake any obligation to publicly revise these statements to reflect events after the date hereof.

SIGNATURES

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.

Cumberland Pharmaceuticals Inc.

Dated: August 28, 2026

By:

/s/ John Hamm

John Hamm

Chief Financial Officer