

MANNKIND CORP  
Form 8-K  
April 02, 2014

**UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION**

**Washington, D.C. 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 1, 2014**

**MannKind Corporation**

(Exact name of registrant as specified in its charter)

**Delaware**  
(State or other jurisdiction  
of incorporation)

**000-50865**  
(Commission File Number)

**13-3607736**  
(IRS Employer Identification No.)

**28903 North Avenue Paine Valencia, California**  
(Address of principal executive offices)

**91355**  
(Zip Code)

Registrant's telephone number, including area code: **(661) 775-5300**

**N/A**  
(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))
- 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 1, 2014, we announced that the U.S. Food and Drug Administration's (FDA) Endocrinologic and Metabolic Drugs Advisory Committee (EMDA) voted 13 to 1 to recommend that AFREZZA<sup>®</sup> (insulin human [rDNA origin]) Inhalation Powder be granted marketing approval by the FDA to improve glycemic control in adults with type 1 diabetes and voted 14 to 0 to recommend that AFREZZA be granted marketing approval by the FDA to improve glycemic control in adults with type 2 diabetes.

Although the EMDA provides recommendations to the FDA, the FDA makes the final decision with respect to approval of a drug. The FDA has assigned a Prescription Drug User Fee Act (PDUFA) action date of April 15, 2014 for its review of our New Drug Application (NDA) for AFREZZA<sup>®</sup>.

A copy of the press release is attached as Exhibit 99.1 to this current report.

**Item 9.01. Financial Statements and Exhibits.**

(d) Exhibits. The following exhibits are filed herewith:

99.1 Press Release of MannKind Corporation dated April 1, 2014, announcing FDA Advisory Committee's recommendation to approve MannKind's Investigational Drug to Treat Diabetes

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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.

**MannKind Corporation**

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(Registrant)

**/s/ DAVID THOMSON, PH.D., J.D.**

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**April 1, 2014**

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(Date)

David Thomson, Ph.D., J.D.

*Corporate Vice President, General Counsel and Secretary*