

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): November 2, 2009**

**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 (*see* General Instruction A.2. below):

- 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))
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## **Item 2.02 Results of Operations and Financial Condition**

On November 2, 2009, MannKind Corporation issued a press release announcing its financial results for the third quarter of 2009. A copy of the press release is attached as Exhibit 99.1 to this Current Report and is incorporated herein by reference.

The information in this Current Report is being furnished and shall not be deemed “filed” for the purposes of Section 18 of the Securities and Exchange Act of 1934, as amended, or otherwise subject to the liabilities of that Section. The information in this Current Report shall not be incorporated by reference into any registration statement or other document pursuant to the Securities Act of 1933, as amended.

### **Item 9.01 Financial Statements and Exhibits**

(c) Exhibits. The following exhibit is furnished herewith:

99.1 Press Release of MannKind Corporation dated November 2, 2009, reporting MannKind’s financial results for the third quarter of 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.

**MANKIND CORPORATION**

By: /s/ MATTHEW J. PFEFFER

Name: Matthew J. Pfeffer

Title: Corporate Vice President and  
Chief Financial Officer

Dated: November 2, 2009

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## EXHIBIT INDEX

Number	Description
99.1	Press Release of MannKind Corporation dated November 2, 2009, reporting MannKind's financial results for the third quarter of 2009.

**MannKind Corporation Reports Third Quarter Financial Results****- Conference Call Today at 5:00 p.m. EST -**

VALENCIA, Calif.--(BUSINESS WIRE)--November 2, 2009--**MannKind Corporation (Nasdaq:MNKD)** today reported financial results for the third quarter ended September 30, 2009.

For the third quarter of 2009, total operating expenses were \$42.8 million, compared to \$69.1 million for the third quarter of 2008. Research and development (R&D) expenses were \$30.5 million for the third quarter of 2009 compared to \$55.6 million for the same quarter in 2008, a decrease of \$25.2 million. This 45% decrease was primarily due to reduced costs associated with the clinical development of AFRESA® as the Company completed its pivotal AFRESA trials in 2008, as well as decreases in clinical supplies costs. General and administrative (G&A) expenses decreased by \$1.2 million or 9% to \$12.3 million for the third quarter of 2009 compared to \$13.4 million in the third quarter of 2008.

For the first nine months of 2009, operating expenses totaled \$154.0 million, compared to \$224.0 million in the first nine months of 2008. R&D expenses for the first nine months were \$113.2 million, compared to \$181.7 million in 2008, a decrease of \$68.4 million. The 38% decrease in R&D expenses for the first nine months was primarily due to decreased costs associated with the clinical development of AFRESA as the Company completed its pivotal AFRESA trials in 2008, as well as decreases in cost of clinical supplies. G&A expenses decreased by \$1.6 million or 4% to \$40.7 million for the first nine months of 2009 as compared to \$42.4 million in the same period in 2008. The decrease in G&A expenses for the first nine months of 2009 was primarily due to the nonrecurrence of costs associated with the purchase of patents during the first quarter of 2008, which was partially offset in 2009 by increased professional fees related to the insulin acquisition transaction with Pfizer Inc. during the second quarter of 2009 and partnership discussions during the third quarter of 2009.

The net loss applicable to common stockholders for the third quarter of 2009 was \$45.6 million, or \$0.42 per share, based on 108.8 million weighted average shares outstanding. This compares to a net loss applicable to common stockholders of \$68.5 million, or \$0.67 per share, based on 101.6 million weighted average shares outstanding for the third quarter of 2008.

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The net loss applicable to common stockholders for the first nine months of 2009 was \$160.6 million, or \$1.54 per share based on 104.4 million weighted average shares outstanding, compared with a net loss applicable to common stockholders of \$219.7 million, or \$2.17 per share based on 101.5 million weighted average shares outstanding, for the first nine months of 2008.

Cash, cash equivalents and marketable securities were \$56.6 million at September 30, 2009 and \$46.5 million at December 31, 2008. Currently, the Company has \$200.0 million of available borrowings under the loan agreement with an entity controlled by the Company's principal stockholder.

"While we continue to be fully engaged in the task of navigating our NDA through the FDA review process, we are continuing to run additional studies of AFRESA. The results to date are preliminary, but we have nonetheless observed some interesting and exciting findings," said Alfred Mann, Chairman and Chief Executive Officer. "The more we study our product in different settings and under different conditions, the more we can appreciate how it has the potential to change the way diabetes is treated. In addition, as we get closer to our January 2010 PFUFA date for the first generation of AFRESA, we are increasing the pace of development of the next generation inhaler in order to provide soon an even better therapeutic option for the millions of patients with diabetes."

### **Conference Call**

MannKind management will host a conference call to discuss these results today at 5:00 p.m. Eastern Time. To participate in the call please dial (888) 677-5721 or (210) 839-8507. To listen to the call via the Internet please visit <http://www.mannkindcorp.com>. The web site replay will be available for 14 days. A telephone replay will be accessible for approximately 14 days following completion of the call by dialing (800) 393-9645 or (203) 369-3721 and entering conference number 7091682.

Presenting from the Company will be:

- Chairman and Chief Executive Officer Alfred Mann
  - President and Chief Operating Officer Hakan Edstrom
  - Corporate Vice President and Chief Financial Officer Matthew Pfeffer
  - Corporate Vice President and Chief Scientific Officer Peter Richardson
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## **About MannKind Corporation**

MannKind Corporation (Nasdaq:MNKD) focuses on the discovery, development and commercialization of therapeutic products for patients with diseases such as diabetes and cancer. Its pipeline includes AFRESA<sup>®</sup>, MKC253, MKC1106-PP, and MKC1106-MT. MannKind has submitted an NDA to the FDA requesting approval of AFRESA for the treatment of adults with type 1 or type 2 diabetes for the control of hyperglycemia. Its other programs are currently in Phase 1 clinical trials. MannKind maintains a website at <http://www.mannkindcorp.com> to which MannKind regularly posts copies of its press releases as well as additional information about MannKind. Interested persons can subscribe on the MannKind website to e-mail alerts that are sent automatically when MannKind issues press releases, files its reports with the Securities and Exchange Commission or posts certain other information to the website.

## **Forward-Looking Statements**

This press release contains forward-looking statements, including statements related to the potential of the Company's products, including AFRESA, and the results of clinical trials, that involve risks and uncertainties. Words such as "believes," "anticipates," "plans," "expects," "intend," "will," "goal," "potential" and similar expressions are intended to identify forward-looking statements. These forward-looking statements are based upon the Company's current expectations. Actual results and the timing of events could differ materially from those anticipated in such forward-looking statements as a result of these risks and uncertainties, which include, without limitation, risks inherent in the progress, timing and results of clinical trials, difficulties or delays in seeking or obtaining regulatory approval, intellectual property matters and other risks detailed in MannKind's filings with the Securities and Exchange Commission, including the Annual Report on Form 10-K for the year ended December 31, 2008 and periodic reports on Form 10-Q and Form 8-K. You are cautioned not to place undue reliance on these forward-looking statements, which speak only as of the date of this press release. All forward-looking statements are qualified in their entirety by this cautionary statement, and MannKind undertakes no obligation to revise or update any forward-looking statements to reflect events or circumstances after the date of this press release.

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**MannKind Corporation**  
**(A Development Stage Company)**  
**Condensed Consolidated Statements of Operations**  
**(Unaudited)**

(In thousands, except per share amounts)

	<b>Three months ended</b>		<b>Nine months ended</b>		<b>Cumulative period from</b>
	<b>September 30,</b>		<b>September 30,</b>		<b>February 14, 1991</b>
	<b>2009</b>	<b>2008</b>	<b>2009</b>	<b>2008</b>	<b>(date of inception)</b>
	<b>2009</b>	<b>2008</b>	<b>2009</b>	<b>2008</b>	<b>to September 30,</b>
	<b>\$</b>	<b>\$</b>	<b>\$</b>	<b>\$</b>	<b>2009</b>
Revenue	\$ —	\$ —	\$ —	\$ 20	\$ 2,988
Operating expenses:					
Research and development	30,494	55,645	113,232	181,665	1,110,714
General and administrative	12,273	13,435	40,727	42,365	286,569
In-process research and development costs	—	—	—	—	19,726
Goodwill impairment	—	—	—	—	151,428
Total operating expenses	<u>42,767</u>	<u>69,080</u>	<u>153,959</u>	<u>224,030</u>	<u>1,568,437</u>
Loss from operations	(42,767)	(69,080)	(153,959)	(224,010)	(1,565,449)
Other income (expense)	149	(7)	503	(7)	(1,440)
Interest expense on note payable to principal stockholder	(1,816)	—	(3,806)	—	(5,329)
Interest expense on senior convertible notes	(1,130)	(124)	(3,376)	(585)	(9,333)
Interest income	9	715	67	4,858	36,928
Loss before provision for income taxes	(45,555)	(68,496)	(160,571)	(219,744)	(1,544,623)
Income taxes	—	—	—	—	(26)
Net loss	(45,555)	(68,496)	(160,571)	(219,744)	(1,544,649)
Deemed dividend related to beneficial conversion feature of convertible preferred stock	—	—	—	—	(22,260)
Accretion on redeemable preferred stock	—	—	—	—	(952)
Net loss applicable to common stockholders	<u>\$ (45,555)</u>	<u>\$ (68,496)</u>	<u>\$ (160,571)</u>	<u>\$ (219,744)</u>	<u>\$ (1,567,861)</u>
Net loss per share applicable to common stockholders — basic and diluted	<u>\$ (0.42)</u>	<u>\$ (0.67)</u>	<u>\$ (1.54)</u>	<u>\$ (2.17)</u>	
Shares used to compute basic and diluted net loss per share applicable to common stockholders	<u>108,779</u>	<u>101,647</u>	<u>104,402</u>	<u>101,495</u>	



**MannKind Corporation**  
**(A Development Stage Company)**  
**Condensed Consolidated Balance Sheet**  
**(Unaudited)**  
(in thousands)

	<b>September 30, 2009</b>	<b>December 31, 2008</b>
<b>Assets</b>		
Current assets:		
Cash and cash equivalents	\$ 53,918	\$ 27,648
Marketable securities	2,649	18,844
State research and development credit exchange receivable — current	1,500	1,500
Prepaid expenses and other current assets	5,247	5,983
Total current assets	63,314	53,975
Property and equipment — net	224,057	226,436
State research and development credit exchange receivable — net of current portion	700	1,500
Other assets	584	548
Total	\$ 288,655	\$ 282,459
<b>Liabilities and Stockholders' Equity (Deficit)</b>		
Current liabilities		
Senior convertible notes	\$ 28,425	\$ 53,472
Note payable to principal stockholder	112,635	112,253
Stockholders' equity (deficit)	150,000	30,000
Total	(2,405)	86,734
Total	\$ 288,655	\$ 282,459

**CONTACT:**

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