

NEKTAR THERAPEUTICS
Form 8-K
November 04, 2009

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 4, 2009

NEKTAR THERAPEUTICS
(Exact Name of Registrant as Specified in Charter)

Delaware
(State or Other Jurisdiction
of Incorporation)

0-24006
(Commission
File Number)

94-3134940
(IRS Employer
Identification No.)

201 Industrial Road
San Carlos, California 94070
(Address of Principal Executive Offices and Zip Code)

Registrant's telephone number, including area code: (650) 631-3100

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))
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Item 2.02 Results of Operations and Financial Condition

On November 4, 2009, Nektar Therapeutics issued a press release (the “Press Release”) announcing its financial results for the third quarter ended September 30, 2009. A copy of the Press Release is furnished herewith as Exhibit 99.1.

On October 30, 2009, the company announced that management would hold a conference call on November 4, 2009 to review its financial results for the third quarter ended September 30, 2009 and provide an update on the company’s business. On this conference call, management expects to make certain forward-looking statements regarding certain pre-clinical and clinical development results and progress for certain of the company’s proprietary drug development programs, the value of the company’s pegylation and advanced polymer chemistry technology platform, potential future revenues that may be realized in the future under certain of the company’s collaboration agreements, and management’s financial guidance for 2009. These forward-looking statements involve substantial risks and uncertainties including but not limited to: (i) the company’s proprietary drug candidates, including NKTR-118, NKTR-102 and NKTR-105, are in early to mid-stage clinical development and the risk of failure remains high and can unexpectedly occur at any stage prior to regulatory approval due to lack of efficacy, safety issues or other factors; (ii) the amount and timing of future payments that may become payable to the company under the agreement with AstraZeneca for NKTR-118 and NKTR-119 is subject to a number of development, regulatory and commercial risks such as the risk of failure to obtain regulatory approval for NKTR-118 and/or NKTR-119 based on safety, efficacy or other issues, the risk of a lack of government or private insurance reimbursement limiting commercial potential, the risk of competition from alternative competing therapies, and other important risks and uncertainties described or referenced herein; (iii) the timing or success of the commencement or end of clinical trials and commercial launch of new drugs may be delayed or unsuccessful due to regulatory delays, clinical trial design, slower than anticipated patient enrollment, drug manufacturing challenges, changing standards of care, clinical outcomes, or delay or failure in obtaining regulatory approval in one or more important markets; (iv) scientific discovery of new medical breakthroughs is an inherently uncertain process and the future success of the application of the company’s technology platforms to potential new drug candidates is therefore very uncertain and unpredictable; (v) clinical trials are long, expensive and uncertain processes and the risk of failure of any drug candidate that is in clinical development and prior to regulatory approval remains high and can occur at any stage due to efficacy, safety or other factors; (vi) management’s financial projections for the company’s 2009 annual revenue, cash used in operations and year-end cash position are subject to the significant risk of unplanned revenue short-falls and unplanned expenses, which could adversely affect the company’s actual 2009 annual financial results and end of year cash position; (vii) the company’s patent applications for its proprietary or partner product candidates may not issue, patents that have issued may not be enforceable, or intellectual property licenses from third parties may be required in the future; (viii) the outcome of any existing or future intellectual property or other litigation related to the company’s proprietary product candidates; (ix) the market sizes and revenue potential of the company’s proprietary and partnered product programs are based on management’s current estimates only and actual market sizes may differ materially; (x) the overall market size for the partnered product programs and revenue and profit contribution potential to the company will depend upon successful sales and marketing efforts by our partners, competition from competing therapies (if any), government and private insurance reimbursement, changing standards of care, commercial product profile and product pricing; (xi) if the company is unable to establish and maintain collaboration partnerships on attractive commercial terms, our business, results of operations and financial condition could suffer; and (xii) certain other important risks and uncertainties set forth in the company’s Annual Report on Form 10-K filed with the Securities and Exchange Commission on March 6, 2009, the company’s most recent Quarterly Report on Form 10-Q for the quarter ended June 30, 2009, and the company’s Quarterly Report on Form 10-Q for the quarter ended September 30, 2009 expected to be filed by the company on or about November 5, 2009. Actual results could differ materially from the forward-looking statements made by management during the conference call and in the press release attached as Exhibit 99.1 hereto. The company undertakes no obligation to update forward-looking statements, whether as a result of new information, future events or otherwise.

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The information in this report, including the exhibit hereto, is being furnished and shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or otherwise subject to the liabilities of that Section or Sections 11 and 12(a)(2) of the Securities Act of 1933, as amended. The information contained herein and in the accompanying exhibit shall not be incorporated by reference into any filing with the Securities and Exchange Commission made by Nektar Therapeutics, whether made before or after the date hereof, regardless of any general incorporation language in such filing.

Item 9.01 Financial Statements and Exhibits

Exhibit	Description
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No.	
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99.1	Press release titled “Nektar Therapeutics Reports Third Quarter 2009 Financial Results” issued on November 4, 2009.
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SIGNATURES

Pursuant to the requirement of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned thereunto duly authorized.

By: /s/ Gil M. Labrucherie
Gil M. Labrucherie
General Counsel and Secretary

Date: November 4, 2009
