

ADVENTRX PHARMACEUTICALS INC

Form 8-K

November 07, 2006



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**Item 7.01. Regulation FD Disclosure.**

Evan M. Levine, Chief Executive Officer of Adventrx Pharmaceuticals, Inc. ( Adventrx ), will present the information reflected in the slides attached as Exhibit 99.1 to this Current Report on Form 8-K (this Report ) on November 7, 2006 at the CIBC World Markets 17th Annual Healthcare Conference at The Waldorf-Astoria Hotel in New York City. In addition, Mr. Levine will present the information reflected in certain selected slides attached as Exhibit 99.1 to this Report on November 7, 2006 at the Rodman and Renshaw 8th Annual Healthcare Conference at the Palace Hotel in New York City.

The information in this Report, including the slides attached hereto as Exhibit 99.1, is being furnished pursuant to this Item 7.01 and shall not be deemed filed for purposes of Section 18 of the Securities Exchange Act of 1934 (the Exchange Act ) or otherwise subject to the liabilities of that section, and it shall not be deemed incorporated by reference in any filing under the Securities Act of 1933 or under the Exchange Act, whether made before or after the date hereof, except as expressly set forth by specific reference in such filing to this Report.

By filing this Report and furnishing this information, Adventrx makes no admission as to the materiality of any information in this Report. The information contained in the slides is summary information that is intended to be considered in the context of Adventrx filings with the Securities and Exchange Commission (the SEC ) and other public announcements that Adventrx makes, by press release or otherwise, from time to time. Adventrx undertakes no duty or obligation to publicly update or revise the information contained in this Report, although it may do so from time to time as its management believes is appropriate. Any such updating may be made through the filing of other reports or documents with the SEC, through press releases or through other public disclosure.

*Adventrx cautions you that information included in the slides attached hereto as Exhibit 99.1 that are not a description of historical facts are forward-looking statements that involve risks, uncertainties, assumptions and other factors that, if they do not materialize or prove to be accurate, could cause Adventrx's results to differ materially from historical results or those expressed or implied by such forward-looking statements. Such forward-looking statements are made based on management's current expectations and beliefs and should not be regarded as a statement or representation by Adventrx that any of its plans, including its anticipated milestones, will be achieved on time or at all. The potential risks and uncertainties that could cause actual results to differ materially include, but are not limited to: the risk that Adventrx will be unable to raise sufficient capital to fund the projects necessary to meet its anticipated or stated goals and milestones; the potential to attract a strategic partner and the terms of any related transaction; the ability to timely enroll subjects in Adventrx's current and anticipated clinical trials; the results of pending clinical trials for CoFactor® or Adventrx's other product candidates; the potential for CoFactor® and Adventrx's other product candidates to receive regulatory approval for one or more indications on a timely basis or at all, and the uncertain process of seeking regulatory approval; other difficulties or delays in developing, testing, manufacturing and marketing of and obtaining regulatory approval for CoFactor® or Adventrx's other product candidates; the market potential for fluoropyrimidine biomodulators and other target markets, and Adventrx's ability to compete in those markets; unexpected adverse side effects or inadequate therapeutic efficacy of CoFactor® or Adventrx's other*

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*products that could delay or prevent regulatory approval or commercialization, or that could result in recalls or product liability claims; the risk that preclinical and clinical results are not indicative of the success of subsequent clinical trials and that products will not perform as preclinical and clinical data suggests or as otherwise anticipated; the potential for regulatory authorities to require additional preclinical work or other clinical requirements to support regulatory filings; the scope and validity of patent protection for CoFactor® and Adventrx's other product candidates; and other risks and uncertainties more fully described in Adventrx's press releases and periodic filings with the Securities and Exchange Commission. Adventrx's public filings with the Securities and Exchange Commission are available at <http://www.sec.gov>.*

*You are cautioned not to place undue reliance on these forward-looking statements, which speak only as of the date when made. All forward-looking statements are qualified in their entirety by this cautionary statement and Adventrx assumes no obligation to revise or update any forward-looking statement, including any information included in the slides attached hereto as Exhibit 99.1, to reflect events or circumstances arising after the date on which it was made. This caution is made under the safe harbor provisions of Section 21E of the Securities Exchange Act of 1934.*

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

(d) Exhibits.

The list of exhibits called for by this Item is incorporated by reference to the Index to Exhibits filed with this report.

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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.

**ADVENTRX Pharmaceuticals, Inc.**

Dated: November 7, 2006

By: /s/ Evan M. Levine

Name: Evan M. Levine

Title: Chief Executive Officer

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99.1 Presentation Slides dated November 7, 2006.