

INCYTE CORP  
Form 8-K  
September 28, 2005

**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: September 28, 2005**  
(Date of earliest event reported)

**INCYTE CORPORATION**

(Exact name of registrant as specified in its charter)

**Delaware**  
(State or Other  
Jurisdiction of  
Incorporation)

**0-27488**  
(Commission File  
Number)

**94-3136539**  
(I.R.S. Employer  
Identification Number)

**Experimental Station, Route  
141 & Henry Clay Road,  
Building E336  
Wilmington, DE**  
(Address of principal executive  
offices)

**19880**  
(Zip Code)

**(302) 498-6700**

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(Registrant's telephone number,  
including area code)

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:

- o Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
  - o Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
  - o Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
  - o Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))
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**ITEM 8.01 OTHER EVENTS.**

On September 28, 2005, Incyte Corporation ( Incyte ) publicly announced that it met with representatives from the Food and Drug Administration ( FDA ) on September 27, 2005, regarding the development of Reverset<sup>TM</sup>, Incyte s nucleoside reverse transcriptase inhibitor ( NRTI ) that is being developed as a therapy for treatment-experienced HIV patients.

The purpose of the meeting was to discuss the results of a recently completed Phase II trial, which were presented in July 2005 at the International AIDS Society meeting, and Incyte s plans to move Reverset into two Phase III pivotal trials. At the meeting, the FDA did not approve of Incyte s moving into Phase III studies. The agency requested that Incyte conduct another Phase II trial to provide additional data to support the efficacy and safety demonstrated in the original Phase II study with the drug.

**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.

Dated: September 28, 2005

INCYTE CORPORATION





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By:

/s/ Patricia A. Schreck









