

ALKERMES INC
Form 8-K
May 27, 2005

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**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): May 27, 2005

ALKERMES, INC.

(Exact Name of Registrant as Specified in its Charter)

PENNSYLVANIA
(State or Other Jurisdiction of
Incorporation)

1-14131
(Commission
File Number)

23-2472830
(I.R.S. Employer
Identification No.)

88 Sidney Street
Cambridge, Massachusetts
(Address of principal executive offices)

02139
(Zip Code)

Registrant's telephone number, including area code: **(617) 494-0171**

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 8.01 Other Events

SIGNATURE

EX-99.1 Alkermes, Inc.

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Item 8.01 Other Events

On May 27, 2005, Alkermes, Inc. announced that the New Drug Application (NDA) for Vivitrol[®] (naltrexone long-acting injection) has been accepted for review by the United States Food and Drug Administration (FDA) and has been granted a Priority Review designation. A copy of the press release is attached hereto as Exhibit 99.1.

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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.

ALKERMES, INC.

Date: May 27, 2005

By: /s/ James M. Frates

James M. Frates
Vice President, Chief Financial Officer and
Treasurer

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EXHIBIT INDEX

- 99.1 Press Release issued by the Company on May 27, 2005 announcing priority review granted for Vivitrex[®] (naltrexone long-acting injection) NDA submission