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ALKERMES INC Form 8-K May 27, 2005

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	1-14131	23-2472830
(State or Other Jurisdiction of	(Commission	(I.R.S. Employer
Incorporation)	File Number)	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):

- 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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TABLE OF CONTENTS

Item 8.01 Other Events
SIGNATURE
EX-99.1 Alkermes, Inc.

Item 8.01 Other Events

On May 27, 2005, Alkermes, Inc. announced that the New Drug Application (NDA) for VivitRexnaltrexone 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.

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

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