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)

June 25, 2007

VIVUS, INC.

(Exact name of registrant as specified in its charter)

Delaware	**000-23490**	**94-3136179**
(State or other jurisdiction of incorporation)	(Commission File Number)	(IRS Employer Identification No.)

1172 CASTRO STREET
MOUNTAIN VIEW, CA 94040
(Address of principal executive offices, including zip code)

(650) 934-5200
(Registrant's telephone number, including area code)

N/A
(Former name or former address, if changed since last report)

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))

Item 8.01. Other Events

On June 25, 2007, VIVUS, Inc. issued a press release titled "VIVUS Announces Abstract Published at American Diabetes Association Scientific Sessions." A copy of the press release is attached hereto as Exhibit 99.1.

The information in this Form 8-K and the exhibit attached hereto shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or incorporated by reference into any of the Registrant's filings under the Securities Act of 1933, as amended, or the Exchange Act, except as shall be expressly set forth by specific reference in any such filing.

Item 9.01. Financial Statements and Exhibits

(d) Exhibits.

Exhibit No.	Description
99.1	Press Release dated June 25, 2007

SIGNATURES

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.

VIVUS, INC.

By: /s/ Lee B. Perry
Lee B. Perry
Vice President and Chief Accounting Officer

Date: **June 25, 2007**

EXHIBIT INDEX

Exhibit No.	Description
99.1	Press Release dated June 25, 2007

CONTACT:

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Chief Financial Officer
650-934-5200

The Trout Group
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FOR IMMEDIATE RELEASE

VIVUS Announces Abstract Published at American Diabetes Association Scientific Sessions
A Retrospective Review of Data from Obese Diabetic Patients Treated in Private Clinic Practice with Topiramate plus Phentermine

MOUNTAIN VIEW, Calif., June 25, 2007 VIVUS, Inc. (NASDAQ: VVUS), a pharmaceutical company dedicated to the development and commercialization of novel therapeutic products, announced that Abstract #2209-PO, *Evaluation of Glycemic Parameters in Obese Diabetic Patients Treated with Phentermine and Topiramate: Outcomes in a Private Practice Setting* has been published in the American Diabetes Association 2007 Scientific Sessions Abstract book. The abstract is available on the American Diabetes Association website.

The purpose of analyzing these data compiled from Dr. Najarian's private medical practice was to evaluate the glycemic effects of topiramate plus phentermine in type 2 diabetic patients with metabolic co-morbidities. The abstract was based on a series of consecutive obese type 2 diabetic patients that had both baseline and on-treatment HbA1c measurements. The results showed that topiramate plus phentermine reduced HbA1c levels and the need for other anti-diabetic medications. Metabolic co-morbidities were also reduced.

About Dr. Najarian

Thomas Najarian, M.D., a Board Certified Specialist in Internal Medicine, is a graduate of MIT (SB, SM in Mechanical Engineering) and Harvard Medical School. He is a former faculty member of the Harvard Medical School. Dr. Najarian was the former Medical Director and Vice-President of Medical Affairs at Interneuron Pharmaceuticals (now Indevus Pharmaceuticals). He is the author of many scientific publications as well as the holder of several U.S. and international patents. Dr. Najarian has 25 years experience in treating obesity and discovered the proprietary combination of topiramate plus phentermine for the treatment of obesity and related conditions. In 2001, VIVUS in-licensed Dr. Najarian's invention and intellectual property associated therewith in exchange for certain milestone payments, stock options and future royalty payments based on product sales. In May of 2006, Dr. Najarian was hired by VIVUS as part-time Principal Scientist. VIVUS is currently developing Qnexa, which is a proprietary pharmaceutical treatment that is a combination of low doses of phentermine and topiramate that is being investigated for the treatment of obesity. The abstract discussed by Dr. Najarian and published in the American Diabetes Association 2007 Scientific Sessions abstract book was not based on Qnexa. VIVUS provided funding for Dr. Najarian's analysis.

About VIVUS

VIVUS, Inc. is a pharmaceutical company dedicated to the development and commercialization of novel therapeutic products. The current portfolio includes investigational products addressing obesity and sexual health. VIVUS has three products that are positioned to enter Phase 3 clinical trials, and one product currently under NDA review by the FDA. The pipeline includes: Qnexa™, for which a Phase 2 study has been completed for the treatment of obesity; Testosterone MDTS®, for which a Phase 2 study has been completed for the treatment of Hypoactive Sexual Desire Disorder (HSDD); EvaMist™, for which a Phase 3 study has been completed and an NDA submitted for the treatment of menopausal symptoms, and on May 15, 2007, the EvaMist assets were transferred to KV Pharmaceutical Company; and avanafil, for which a Phase 2 study has been completed for the treatment of erectile dysfunction (ED). MUSE® is approved and currently on the market for the treatment of ED. For more information on clinical trials and products, please visit the company's web site at www.vivus.com.

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Certain statements in this press release are forward-looking within the meaning of the Private Securities Litigation Reform Act of 1995. These statements may be identified by the use of forward-looking words such as "anticipate," "believe," "forecast," "estimated" and "intend," among others. These forward-looking statements are based on VIVUS' current expectations and actual results could differ materially. There are a number of factors that could cause actual events to differ materially from those indicated by such forward-looking statements. These factors include, but are not limited to, substantial competition; uncertainties of patent protection and litigation; uncertainties of government or third party payer reimbursement; reliance on sole source suppliers; limited sales and marketing efforts and dependence upon third parties; risks related to the development of innovative products; and risks related to failure to obtain FDA clearances or approvals and noncompliance with FDA regulations. As with any pharmaceutical under development, there are significant risks in the development, regulatory approval and commercialization of new products. There are no guarantees that the EvaMist NDA submission will be approved in a timely basis, or at all. There are no guarantees that future clinical studies discussed in this press release will be completed or successful or that any product will receive regulatory approval for any indication or prove to be commercially successful. VIVUS does not undertake an obligation to update or revise any forward-looking statement. Investors should read the risk factors set forth in VIVUS' Form 10- K for the year ended December 31, 2006 and periodic reports filed with the Securities and Exchange Commission.
