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): March 4, 2009

OPKO Health, Inc.

(Exact Name of Registrant as Specified in Charter)

Delaware (State or other jurisdiction of incorporation)

000-26648 (Commission File Number)

75-2402409 (IRS Employer Identification No.)

4400 Biscayne Blvd
Suite 1180
Miami, Florida 33137
(Address of Principal Executive Offices)

Registrant's telephone number, including area code: (305) 575-4138

(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):	
	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))

ITEM 1.01. Entry into a Material Definitive Agreement.

On March 4, 2009, Frost Gamma Investments Trust (the "Gamma Trust"), a trust controlled by Dr. Phillip Frost, the Company's Chairman and Chief Executive Officer, advanced \$3,000,000 to the Company under a Promissory Note issued by the Company to the Gamma Trust (the "Note"). The entire amount of this advance and all accrued interest thereon shall be due and payable on the earlier of May 4, 2009 or such earlier date following the closing of the previously disclosed Stock Purchase Agreement, dated February 23, 2009, between the Company and the Gamma Trust, pursuant to which the Gamma Trust agreed to make a \$20 million investment in the Company. The Note bears interest at a rate equal to 11% per annum and may be prepaid in whole or in part without penalty or premium.

ITEM 2.03. Creation of a Direct Financial Obligation or an Obligation Under an Off-Balance Sheet Arrangement of a Registrant.

The information set forth above in Item 1.01 is hereby incorporated by reference into this Item 2.03.

ITEM 8.01. Other Events.

On March 6, 2008, the Company issued a press release announcing an update on its Phase III Clinical Trial of bevasiranib. It was announced that, following the recommendation of the Independent Data Monitoring Committee, it had decided to terminate its Phase III clinical study of Bevasiranib. A copy of the press release is filed as Exhibit 99 to this report.

ITEM 9.01. Financial Statements and Exhibits.

99 Press Release

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.

OPKO Health, Inc.

By: /s/ Rao Uppaluri
Name: Rao Uppaluri

Title: Chief Financial Officer

Date March 6, 2009



OPKO Health Announces Update on Phase III Clinical Trial of Bevasiranib

MIAMI, March 6, 2009 (BUSINESS WIRE) — OPKO Health, Inc. (NYSE Alternext US:OPK)today announced that, following the recommendation of the Independent Data Monitoring Committee (IDMC), it had decided to terminate its Phase III clinical study of Bevasiranib, a first-in-class siRNA, for the treatment of wet age-related macular degeneration (wet-AMD). Although preliminary data, needing further analysis, show activity of Bevasiranib when used adjunctively with Genentech's Lucentis®, review of the data by the IDMC indicated that the trial, as structured, was unlikely to meet its primary end point. There were no systemic safety issues identified and local ocular safety was generally unremarkable. "While we are clearly disappointed with the preliminary results of this fully-enrolled study, the indications of activity are encouraging and we look forward to fully analyzing the data in the coming weeks," said Dr. Phillip Frost, Chairman and CEO of OPKO Health. "We remain committed to the continued development of our siRNA portfolio targeting Vascular Endothelial Growth Factor (VEGF), including our recently announced VEGFA165b sparing siRNA." These new proprietary siRNA's are designed to inhibit the angiogenic Vascular Endothelial Growth Factor A165 (VEGFA165 isoform) but spare the anti-angiogenic VEGFA165b isoform. VEGFA165 is known to play a critical role in diseases of the eye where the underlying cause of the problem is abnormal growth of blood vessels, such as in wet age-related macular degeneration. To the contrary, data show that VEGFA165b is an inhibitor of abnormal vessel growth.

About OPKO Health, Inc.

Miami-based OPKO is a specialty healthcare company engaged in the development, marketing, and sales of novel agents and both diagnostic and therapeutic devices for the management of ophthalmic diseases. For more information visit the company's website at www.opko.com.

This press release contains "forward-looking statements," as that term is defined under the Private Securities Litigation Reform Act of 1995 (PSLRA), which statements may be identified by words such as "expects," "plans," "projects," "will," "may," "anticipates," "believes," "should," "intends," "estimates," and other words of similar meaning, including statements regarding preliminary data, our product development efforts, our ability to significantly improve clinical outcomes in patients, and our ability to develop a preclinical pipeline of novel agents for ophthalmic diseases, as well as other non-historical statements about our expectations, beliefs or intentions regarding our business, technologies and products, financial condition, strategies or prospects. Many factors could cause our actual activities or results to differ materially from the activities and results anticipated in forward-looking statements. These factors include those described in our filings with the Securities and Exchange Commission, as well as risks inherent in funding, developing and obtaining regulatory approvals of new, commercially-viable and competitive products and treatments. In addition, forward-looking statements may also be adversely affected by general market factors, competitive product development, product availability, federal and state regulations and legislation, the regulatory process for new products and indications, manufacturing issues that may arise, patent positions and litigation, among other factors. The forward-looking statements contained in this press release speak only as of the date the statements were made, and we do not undertake any obligation to update forward-looking statements. We intend that all forward-looking statements be subject to the safe-harbor provisions of the PSLRA.

SOURCE: OPKO Health, Inc. OPKO Health, Inc., Miami Steven D. Rubin, 305 575-6015