

ORASURE TECHNOLOGIES INC
Form 8-K
April 04, 2012

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): April 4, 2012

OraSure Technologies, Inc.

(Exact Name of Registrant as Specified in Charter)

Delaware
(State or Other Jurisdiction

of Incorporation)

001-16537
(Commission

File Number)

36-4370966
(I.R.S. Employer

Identification No.)

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220 East First Street

Bethlehem, Pennsylvania
(Address of Principal Executive Offices)

Registrant's telephone number, including area code: 610-882-1820

18015-1360
(Zip 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:

- .. 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 7.01 Regulation FD Disclosure.

On April 4, 2012, OraSure Technologies, Inc. (the Company) issued a press release announcing that the U.S. Food and Drug Administration's Blood Products Advisory Committee will consider the Company's application for the approval of its OraQuick® Rapid HIV-1/2 test for sale in the U.S. consumer or over-the-counter market at a meeting scheduled for May 15, 2012. A copy of the press release is attached as Exhibit 99 to this Form 8-K and is incorporated herein by reference.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

Exhibit Number	Description
99	Press Release, dated April 4, 2012, announcing that the U.S. Food and Drug Administration's Blood Products Advisory Committee will consider the Company's application for the approval of its OraQuick® Rapid HIV-1/2 test for sale in the U.S. consumer or over-the-counter market at a meeting scheduled for May 15, 2012.

Signatures

Pursuant to the requirements of the Securities and Exchange Act of 1934, the Registrant has duly caused this report to be signed on its behalf by the undersigned, hereunto duly authorized.

ORASURE TECHNOLOGIES, INC.

Date: April 4, 2012

By: */s/ Jack E. Jerrett*
Jack E. Jerrett
Senior Vice President, General Counsel and Secretary

Index to Exhibits

**Exhibit
No.**

Description

99	Press Release, dated April 4, 2012, announcing that the U.S. Food and Drug Administration's Blood Products Advisory Committee will consider the Company's application for the approval of its OraQuick [®] Rapid HIV-1/2 test for sale in the U.S. consumer or over-the-counter market at a meeting scheduled for May 15, 2012.
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