ORASURE TECHNOLOGIES INC Form 8-K July 03, 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): July 3, 2012

OraSure Technologies, Inc.

(Exact Name of Registrant as Specified in Charter)

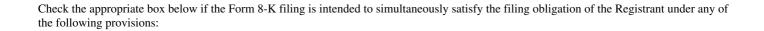
Delaware (State or Other Jurisdiction

001-16537 (Commission 36-4370966 (I.R.S. Employer

of Incorporation) File Number) Identification No.)

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)



- " 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 8.01 Other Events.

On July 3, 2012, OraSure Technologies, Inc. (the Company) issued a press release announcing the receipt of U.S. Food and Drug Administration (FDA) approval of the Company s OraQhibk-Home HIV Test. A copy of the press release is attached as Exhibit 99.1 to this Form 8-K and is incorporated herein by reference.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

Exhibit	
Number	Description
99.1	Press Release, dated July 3, 2012, announcing the receipt of FDA approval of the OraOuick® In-Home HIV Test.

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: July 3, 2012 By: /s/ Jack E. Jerrett

Jack E. Jerrett

Senior Vice President, General Counsel and Secretary

Index to Exhibits

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99.1	Press Release, dated July 3, 2012, announcing the receipt of FDA approval of the OraQuick® In-Home HIV Test.