

ORASURE TECHNOLOGIES INC  
Form 8-K  
June 25, 2010

**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, 2010**

**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.)

Edgar Filing: ORASURE TECHNOLOGIES INC - Form 8-K

**220 East First Street**

**Bethlehem, Pennsylvania**  
(Address of Principal Executive Offices)

**18015-1360**  
(Zip Code)

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

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 June 25, 2010, 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 OraQuick<sup>®</sup> HCV Rapid Antibody Test for use with venous whole blood samples. 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**

<b>Exhibit Number</b>	<b>Description</b>
99	Press Release, dated June 25, 2010, announcing the receipt of FDA approval of the Company's OraQuick <sup>®</sup> HCV Rapid Antibody Test for use with venous whole blood samples.

**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: June 25, 2010

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

**Index to Exhibits**

<b>Exhibit No.</b>	<b>Description</b>
99	Press Release, dated June 25, 2010, announcing the receipt of FDA approval of the Company's OraQuic® HCV Rapid Antibody Test for use with venous whole blood samples.