CRYOLIFE INC
Form 8-K
February 12, 2008

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): February 7, 2008

CRYOLIFE, INC.
(Exact name of registrant as specified in its charter)

Florida	1-13165	59-2417093
(State or Other Jurisdiction of Incorporation)	(Commission File Number)	(IRS Employer Identification No.)

1655 Roberts Boulevard, N.W., Kennesaw, Georgia 30144
(Address of principal executive office) (zip code)

Registrant's telephone number, including area code: (770) 419-3355

(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))

Section 8 Other Events

Item 8.01 Other Events.

On February 7, 2008, CryoLife, Inc. (“CryoLife” or the “Company”) issued a press release announcing that the Food and Drug Administration (“FDA”) notified CryoLife that it has received 510(k) clearance from the FDA for its CryoValve SG pulmonary heart valve processed with the Company’s proprietary SynerGraft technology. CryoLife hereby incorporates by reference herein the information set forth in its Press Release dated February 7, 2008, a copy of which is attached hereto as Exhibit 99.1

Except for the historical information contained in this report, the statements made by CryoLife are forward-looking statements that involve risks and uncertainties. All such statements are subject to the safe harbor created by the Private Securities Litigation Reform Act of 1995. These statements include those regarding anticipated effectiveness, benefits and indications for use of CryoValve SG as well as the timing of use of the SynerGraft technology. These future events may not occur as and when expected, if at all, and, together with the Company's business, are subject to various risks and uncertainties. These risks and uncertainties include that CryoValve SG may not perform as well as expected or provide all of the benefits anticipated, the Company may not be able to begin processing the majority of its pulmonary valves by the anticipated time, nor may the first shipments of CryoValve SG occur as expected, and other risk factors detailed in CryoLife's Securities and Exchange Commission filings, including CryoLife's Form 10-K filing for the year ended December 31, 2006, its most recent Form 10-Q, and the Company's other SEC filings. The Company does not undertake to update its forward-looking statements.

Section 9 Financial Statements and Exhibits

Item 9.01(c) Exhibits.

(a) Financial Statements.
Not applicable.

(b) Pro Forma Financial Information.
Not applicable.

(c) Shell Company Transactions.
Not applicable.

(d) Exhibits.

Exhibit Number	Description
99.1	Press release dated February 7, 2008

SIGNATURES

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

CRYOLIFE, INC.

Date: February 11, 2008

By: /s/ D. A. Lee
Name: D. Ashley Lee
Title: Executive Vice President, Chief Operating Officer and Chief Financial Officer