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CRYOLIFE INC Form 8-K April 21, 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

	SECURI	TIES EXCHANGE ACT OF 1934	
	Date of Report (Da	te of earliest event reported): Apr	il 16, 2008
	(Exact name	CRYOLIFE, INC. of registrant as specified in its cha	rter)
	Florida —	1-13165	59-2417093
	(State or Other Jurisdiction of Incorporation)	(Commission File Number)	(IRS Employer Identification No.)
	(Address of	ulevard, N.W., Kennesaw, Georgia principal executive office) (zip co e number, including area code: (77	de)
	(Former name or f	former address, if changed since la	est report)
	neck the appropriate box below if the Forgistrant under any of the following pro-		
O	Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)		
0	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))

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Section 1 Registrant's Business and Operations

Item 1.01 Entry into a Material Definitive Agreement

On April 16, 2008, CryoLife, Inc. (the "Company" or "CryoLife") entered into an exclusive distribution agreement with Medafor, Inc. ("Medafor"). Pursuant to the agreement, CryoLife will distribute an absorbable blood clotting agent manufactured and developed by Medafor. A copy of the press release announcing the agreement is attached as Exhibit 99.1 hereto and incorporated by reference hereto.

It is anticipated that CryoLife will begin distribution of the absorbable blood clotting agent, marketed by CryoLife as Hemostase MPH, on May 1, 2008. The agreement has a three-year term with a three-year right of renewal by CryoLife if CryoLife meets the annual minimum purchase requirements. The agreement gives CryoLife the exclusive distribution rights for Hemostase MPH in specified territories for all applications in cardiac and vascular surgery, and with respect to countries other than the United States, China and Japan, for general surgery applications, excluding orthopedic and ear, nose, and throat surgeries.

Commencing May 1, 2008, CryoLife's distribution territory under the distribution agreement includes the United States (excluding approximately 41 hospitals where Medafor will retain distribution rights until no later than December 31, 2008 and Department of Defense facilities), the United Kingdom, Germany and Canada. Subject to the exceptions noted above, the territory will expand to include the entire world except Japan and China beginning in January 1, 2009. Department of Veterans Affairs hospitals are not considered Department of Defense facilities as part of the agreement.

If CryoLife fails to purchase certain specified annual amounts under the agreement, Medafor may terminate the distribution agreement, with no further obligation owed by CryoLife. If CryoLife terminates the distribution agreement, other than for certain breaches by Medafor, then CryoLife is required to reimburse Medafor for Medafor's cost of up to three months of Hemostase MPH inventory maintained by Medafor or purchase such inventory from Medafor.

Section 9 Financial Statements and Exhibits.

Item 9.01 Financial Statements and Exhibits.

(a) Financial Statements.

Not applicable.

(b) Pro Forma Financial Information.

Not applicable.

(c) Shell Company Transactions.

Not applicable.

(d) Exhibits.

99.1 Press Release dated April 17, 2008.

2

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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: April 21, 2008 By:/s/ D.A. Lee

Name: D. Ashley Lee

Title: Executive Vice President, Chief

Operating Officer and Chief

Financial Officer

3