

INSPIREMD, INC.

FORM 8-K (Current report filing)

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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 15, 2015

InspireMD, Inc.

(Exact name of registrant as specified in its charter)

	Delaware	001-35731	26-2123838
	(State or other	(Commission File Number)	(IRS Employer
	jurisdiction		Identification No.)
	of incorporation)		
	321 Columbus Aven	ue	
Boston, Massachusetts (Address of principal executive		tts	02116
		ve offices)	(Zip Code)
	Registrant	's telephone number, including area code: (857) 45	3-6553
	(Forme	r name or former address, if changed since last rep	ort)
any	Check the appropriate box below if the For	rm 8-K filing is intended to simultaneously satisfy	the filing obligation of the registrant under
	Written communications pursuant to Rule 42	25 under the Securities Act (17 CFR 230.425)	
	Soliciting material pursuant to Rule 14a-12 u	under the Exchange Act (17 CFR 240.14a-12)	
	Pre-commencement communications pursua	nt to Rule 14d-2(b) under the Exchange Act (17 Cl	FR 240.14d-2(b))
	Pre-commencement communications pursua	nt to Rule 13e-4 (c) under the Exchange Act (17 C	FR 240.13e-4(c))

Item 2.02 Results of Operations and Financial Condition.

On April 15, 2015, InspireMD, Inc. (the "Company") issued a press release pre-announcing its revenues for the fiscal quarter ended March 31, 2015. A copy of this press release is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

In accordance with General Instruction B.2 of Form 8-K, the information in this Current Report on Form 8-K with respect to the Company's revenues for the fiscal quarter ended March 31, 2015, including the announcement of the Company's revenues for the fiscal quarter ended March 31, 2015 in Exhibit 99.1 hereto, that is furnished pursuant to this Item 2.02 shall not be deemed to be "filed" for the purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise subject to the liabilities of that section, and shall not be incorporated by reference into any registration statement or other document filed under the Securities Act of 1933, as amended, or the Exchange Act, except as shall be expressly set forth by specific reference in such filing.

Item 8.01 Other Events.

On April 15, 2015, the Company issued a press release announcing the completion of its sales restructuring program which began on January 4, 2015.

A copy of the press release is attached as Exhibit 99.1 to this Current Report on Form 8-K and is incorporated by reference herein.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

Exhibit Number		Description	
99.1	Press release dated April 15, 2015	•	

SIGNATURES

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

INSPIREMD, INC.

Date: April 15, 2015 By: /s/ Craig Shore

Name: Craig Shore Title: Chief Financial Officer



InspireMD Announces Completion of Sales Restructuring Program

BOSTON, MA – April 15, 2015 – <u>InspireMD, Inc.</u> (NYSE MKT: NSPR) ("InspireMD" or the "Company"), a leader in stent embolic protection systems ("EPS"), today announced the completion of its sales restructuring program which began on January 4, 2015. The restructuring was implemented to conserve capital and to align the organization to a new commercial strategy built on third party distribution partners. Reflecting some impact from the transition, as well as the trend towards drug eluting stent use in STEMI patients discussed during our fourth quarter 2014 earnings call last month, total revenues for the first quarter 2015 were approximately \$500,000, down sequentially from \$850,000 in the fourth quarter 2014.

Alan Milinazzo, CEO of InspireMD, commented, "2015 is a transition year for us, as we pivot our strategy to exploit the immediate carotid and emerging neurovascular opportunities while being more selective with our sales resource allocation and shift to a distributor based commercial model."

Mr. Milinazzo, concluded, "With our sales restructuring program now behind us, our cash projections remain on track and we look forward to improving our sales performance, mindful of less linear growth in the coming quarters, as we intend to lead our commercial efforts with our carotid platform. We began shipping our new CGuard RX product during the quarter and while quantities were limited due to short term supply constraints, we expect improvements beginning in the second quarter."

InspireMD will report full first quarter 2015 results and host its investor earnings conference call on May 11, 2015.

About InspireMD, Inc.

InspireMD seeks to utilize its proprietary MGuardTM with MicroNetTM technology to make its products the industry standard for embolic protection and to provide a superior solution to the key clinical issues of current stenting in patients with a high risk of distal embolization, no reflow and major adverse cardiac events.

InspireMD intends to pursue applications of this MicroNet technology in coronary, carotid (CGuardTM) and peripheral artery procedures. InspireMD's common stock is quoted on the NYSE MKT under the ticker symbol NSPR.



Forward-looking Statements

This press release contains "forward-looking statements." Such statements may be preceded by the words "intends," "may," "will," "plans," "expects," "anticipates," "projects," "predicts," "estimates," "aims," "believes," "hopes," "potential" or similar words. Forward-looking statements are not guarantees of future performance, are based on certain assumptions and are subject to various known and unknown risks and uncertainties, many of which are beyond the Company's control, and cannot be predicted or quantified and consequently, actual results may differ materially from those expressed or implied by such forward-looking statements. Such risks and uncertainties include, without limitation, risks and uncertainties associated with (i) market acceptance of our existing and new products, (ii) negative clinical trial results or lengthy product delays in key markets, (iii) an inability to secure regulatory approvals for the sale of our products, (iv) intense competition in the medical device industry from much larger, multinational companies, (v) product liability claims, (vi) product malfunctions, (vii) our limited manufacturing capabilities and reliance on subcontractors for assistance, (viii) insufficient or inadequate reimbursement by governmental and other third party payers for our products, (ix) our efforts to successfully obtain and maintain intellectual property protection covering our products, which may not be successful, (x) legislative or regulatory reform of the healthcare system in both the U.S. and foreign jurisdictions, (xi) our reliance on single suppliers for certain product components, (xii) the fact that we will need to raise additional capital to meet our business requirements in the future and that such capital raising may be costly, dilutive or difficult to obtain and (xiii) the fact that we conduct business in multiple foreign jurisdictions, exposing us to foreign currency exchange rate fluctuations, logistical and communications challenges, burdens and costs of compliance with foreign laws and political and economic instability in each jurisdiction. More detailed information about the Company and the risk factors that may affect the realization of forward looking statements is set forth in the Company's filings with the Securities and Exchange Commission (SEC), including the Company's Annual Report on Form 10-K and its Ouarterly Reports on Form 10-O. Investors and security holders are urged to read these documents free of charge on the SEC's web site at http://www.sec.gov. The Company assumes no obligation to publicly update or revise its forward-looking statements as a result of new information, future events or otherwise.

Investor Contacts:

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