

INSPIREMD, INC.

FORM FWP

(Free Writing Prospectus - Filing under Securities Act Rules 163/433)

Filed 02/21/17

Telephone	(888) 776-6804
CIK	0001433607
Symbol	NSPR
SIC Code	3841 - Surgical and Medical Instruments and Apparatus
Industry	Medical Equipment, Supplies & Distribution
Sector	Healthcare
Fiscal Year	12/31

Issuer Free Writing Prospectus
Filed Pursuant to Rule 433 of the
Securities Act of 1933, as amended
Registration Statement No. 333-215682



Investor Presentation

NYSE MKT: NSPR

February 2017

This presentation 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 payors 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 Quarterly Reports on Form 10-Q. 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.

This presentation highlights basic information about InspireMD, Inc. and the offering. InspireMD, Inc. has filed a registration statement on Form S-1 (Registration No. 333-215682) (including a prospectus) with the U.S. Securities and Exchange Commission (the “SEC”) for the offering to which this presentation relates. The registration statement has not yet become effective. Before you invest, you should read the prospectus in that registration statement (including, among other things, risk factors described therein) and other documents the issuer has filed with the SEC for more complete information about InspireMD, Inc. and this offering. The preliminary prospectus dated January 24, 2017, and subsequent amendments are available at the SEC website. You may get these documents for free by visiting EDGAR on the SEC website at www.sec.gov. Alternatively, InspireMD, Inc. or any underwriter or any dealer participating in the offering will arrange to send you the prospectus if you request it by contacting Dawson James Securities, Inc., Attention: Prospectus Department, 1 North Federal Highway, 5th Floor, Boca Raton, FL 33432, mmaclaren@dawsonjames.com or toll free at 866.928.0928.

The Offering



Issuer	InspireMD, Inc.
Exchange / Ticker	NYSE MKT / NSPR
Offering Type	Best Efforts
Offering Size	\$7,500,000
Security Type	<ul style="list-style-type: none">• 1 Preferred Stock – initially convertible to 4 shares of common stock• 1 5 year Series B Warrant to purchase 4 shares of common stock - exercise price 125 % premium of conversion price of the Preferred Stock• 1 6 month Series C Warrant to purchase 4 shares of common stock – exercise price equal to the conversion price of the Preferred Stock
Use of Proceeds	To further fund the expansion of our sales and marketing for CGuard™ EPS and MGuard™ EPS. If we receive sufficient proceeds from the exercise of the Series C Warrants, we plan to continue the development of and manufacturing enhancements of CGuard™ EPS and further our efforts to obtain an Investigation Device Exemption (IDE) for CGuard™ EPS. Any balance of the net proceeds will be used for general corporate purposes.
Sole Bookrunner	Dawson James Securities, Inc.

InspireMD is a commercial-stage medical device company with proprietary and innovative embolic prevention systems (EPS)/thrombus management technologies and neurovascular devices that seek to overcome the harmful consequences of conventional stenting.

COMPANY

NYSE MKT:	NSPR
Founded:	2005
Employees:	34
Headquarters:	Tel Aviv
Manufacturing Facility:	Tel Aviv

TECHNOLOGY

Proprietary MicroNet™ technology in multiple products seeking a superior solution for the treatment of complex vascular and coronary disease

PRODUCTS

Commercial:
CGuard™ Carotid EPS
MGuard™ Coronary EPS

Pipeline:
Next Gen CGuard™ - 5F
NGuard™
PVGuard™

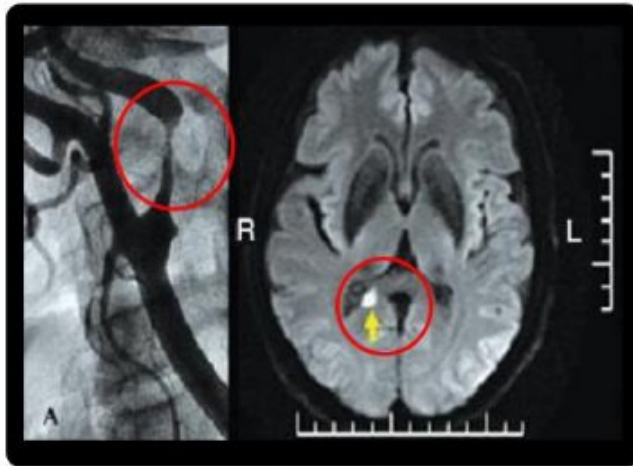
- Multi-billion dollar opportunity for MicroNet™ products for multiple vascular markets
 - Current stents do not adequately address the risk of post-procedural embolization
 - Consistent positive clinical trial results positioning CGuard™ as a potential standard-of-care in treating carotid artery disease
- Revenue growth driven by new commercialization strategy
 - Proven success with recent YOY sales growth of 67% in select markets with InspireMD managed regional distribution model
 - Transitioning from exclusive European distributor (18 countries) to established InspireMD managed regional distributor model
 - Expanding CGuard™ users to a greater number of vascular surgeons, interventional cardiologists, and interventional radiologists
- Recent leadership changes focused on sales, marketing and high value pipeline development
- Strategic collaboration outreach expanding for multiple MicroNet™ product applications
- A broad portfolio of patent-protected assets

“Plaque protrusion through stent struts occurs in up to 65% of conventional carotid stents in relation to plaque morphology/symptomatic status and stent type, providing a mechanism for post carotid artery stenting (CAS) cerebral embolization, either directly or via additional thrombus formation.”*



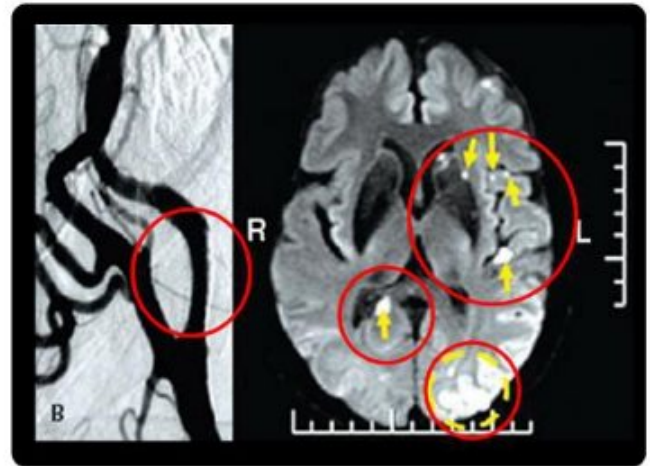
2/3 of CAS neurovascular events (stroke, TIA) are POST-procedural.**

* Musialek, et al. Eurointerventions 2016; 12 August 2016.
** Bosiers et al. Eur J Vasc Endovasc Surg Vol 33, Feb 2007.



Pre-Procedure

Pre-intervention showing 90% occlusion of the carotid artery and an MRI showing an old white matter infarction (obstruction).

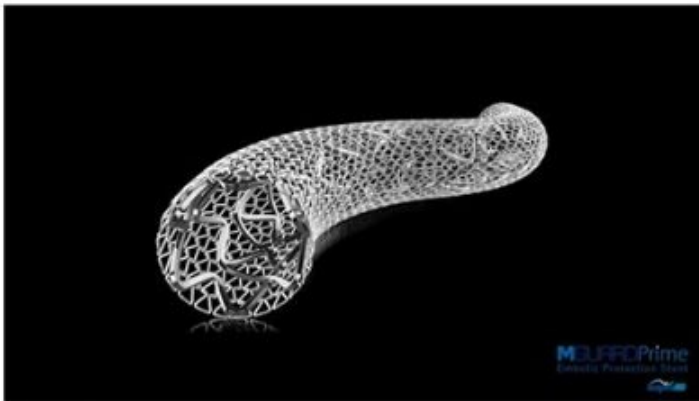
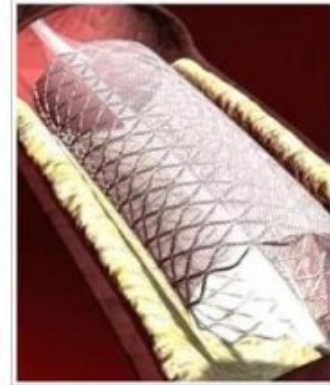


Post-Procedure

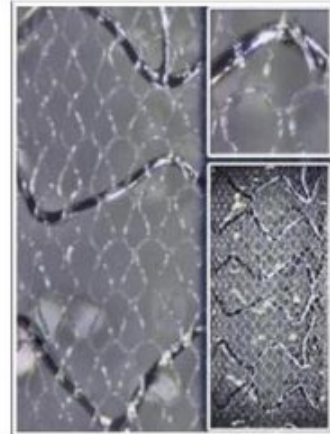
Post-intervention showing successful opening of the occluded carotid artery with conventional stenting and an MRI showing multiple micro-infarcts (obstructions) post-procedure due to liberation of embolic particles.

MicroNet™ Prevents Distal Embolization and Other Vascular Disease Challenges

- Ultrathin PET* mesh provides meaningful clinical benefit versus conventional devices
- Provides revascularization benefit
- MicroNet™ acts as a “safety net” by offering greater vessel area coverage to prevent large plaque protrusion through the scaffold into the vessel lumen
- Made of a single fiber from a biocompatible polymer, widely used in other medical implants
- Stents incorporating MicroNet™ have identical deliverability to other stents



*PET – polyethylene terephthalate



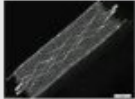



- Proprietary platform technology supported by a robust intellectual property portfolio
- Continue to strengthen and broaden patent protection globally to enable future pipeline products

PATENT RIGHTS	ISSUED	ALLOWED	PENDING
USA	6	0	12
Rest of World	16	1	15

Large & Growing Addressable Market



Embolec Prevention Products	Market Oppty	CE Mark	Focus Area
CGuard™ 	\$500M	✓	Carotid
MGuard™* 	\$1.7B	✓	Coronary AMI & SVG
NGuard™ 	\$675M	Planned Submission TBD	Neurovascular
PVGuard™ 	\$1.7B	Planned Submission TBD	Peripheral

* MGuard™ global strategy focused on drug eluting stent OEM partnership

* MGuard is a bare metal stent scaffold

- US FDA

- Pre-IDE FDA submission for CGuard™ completed
- Formal FDA meeting planned
- Planned IDE submission in H2 2017



- Next generation CGuard™ - 5 French CGuard™



- Minimally invasive devices trending smaller for broader and easier usage
- Lower profile system for cases where pre-dilatation could be problematic
- Competitive advantage in the Asia/Pacific markets
 - Smaller anatomy particularly in the female population
- Transradial delivery (delivery from the wrist vs. femoral artery) gaining favor among interventionalists

* Subject to receipt of sufficient funding

CARENET Clinical Trial (2014)

- 30 Patient Safety and Efficacy clinical trial
- **Zero major adverse cardiac or cerebral events (MACCE)** at 30 days (Comparative data 5.72%*)
- **50% fewer new ischemic lesions** with lesion volume being 10x times smaller compared to historical non-mesh carotid artery stenting data
- **All new ischemic lesions fully resolved at 30 days** except one
- 3.6% MACCE rate at 6 months (Comparative data 8.09%**)
- **Zero strokes or stroke related deaths at 12 months**

PARADIGM 101 Clinical Trial (2015 and 2016)

- 101 patient trial evaluating CGuard EPS in unselected, consecutive carotid patients (all-comers)
- **99.1% device success**
- **0% MACCE (Death/stroke/MI) @ 48 hr**
- **0% MACCE @ 30 day**
- **Zero strokes or stroke related deaths at 12 months**



"CGuard can safely be used on more than 90% of all-comer patients that have carotid artery stenosis."
P. Musialek, MD

* Trials included in analysis: ARChER pooled, ARMOUR, BEACH, CABERNET, CREATE, EMPIRE, EPIC, MAVERiC 1+2, MAVERiC International, PRIAMUS, SAPPHIRE, SECURITY, PROFi, ICSS

** Values extrapolated from event curves

Independent study conducted in 30 patients with internal carotid artery disease

Clinical results (2016)

- **100% success** in implanting the CGuard™ EPS
- No peri- or post-procedural complications
- **No deaths, major adverse events, minor or major strokes**, or new neurologic symptoms during the six months following the procedure
- All vessels treated with the CGuard™ system remained patent (open) at six months
- DW-MRI performed in 19 of 30 patients found **no new ipsilateral lesions after 30 days and after six months** compared with baseline DW-MRI studies

ENDOVASCULAR THERAPY

Clinical Results and Mechanical Properties of the Carotid CGUARD Double-Layered Embolic Prevention Stent

Christine Wissgott, MD¹, Wiltham Schmidt, PhD¹, Christoph Brandt-Wunderlich, MS¹, Peter Behrens, MS¹, and Rainer Andressen, MD²

Abstract

Purpose: To report early device outcomes with a novel double-layer stent for the internal carotid artery (ICA) and the in vitro investigation of the stent's mechanical properties. **Methods:** A prospective single-center study enrolled 30 consecutive patients (mean age 73.1±4.3 years, 31 men) with angiographic 50% or high-grade 60-70% ICA stenosis treated with the new double-layer carotid CGUARD Embolic Prevention System (EPS) stent, which has an open-cell stent design with an inner closed-cell polyethylene terephthalate layer. The average diameter of the treated artery was 8.1±1.7mm with a mean lesion length of 1.4±0.2 cm. In the laboratory, 40-mm stents were tested in vitro with respect to their radial force during expansion, the bending stiffness of the stent system and the expanded stent, as well as the collapse pressure in a sin and flexion device. The wall thickness was assessed using fluoroscopy after stent release in sin and normal vessel models. **Results:** The stent was successfully implanted in all patients. No peri- or post-procedural complications occurred; no minor or major stroke was observed in the 6-month follow-up. The bending stiffness of the expanded stent was 0.21 N/cm² and (not surprisingly) was clearly lower than that of the stent system (0.13 N/cm²). The revascularized radial force during expansion of the stent is 7.0 atm, consistent with in vivo usage, was relatively high (0.034 N/mm²), which correlates well with the collapse pressure of 6.11 bars. Residual wall adhesion was minimal and caused no impairment of the vessel after device application. **Conclusions:** Because of its structure, the novel CGUARD EPS stent is characterized by a high flexibility combined with a high radial force and very good plaque coverage. These first clinical results demonstrate a very safe implantation behavior without any stroke up to 6 months after the procedure.

Keywords: carotid artery stent, closed-cell design, double-layer stent, embolic filter, internal carotid artery, in vitro testing, mechanical behavior, radial, open-cell design, radial force, sinosis, stent.

Introduction

Several studies have demonstrated that carotid artery stenting (CAS) of the occluded internal carotid artery (ICA) is a well-tolerated and equally good option for treating intracranial carotid stenosis in comparison with carotid endarterectomy (CEA).¹⁻⁴ Although CEA is still considered the gold standard therapy of carotid stenosis,^{5,6} because of a lower risk of periprocedural and postprocedural stroke,^{7,8} the stent system is a valuable treatment option. Nevertheless, it is less invasive than CEA.⁹


Despite the fact that procedures related events can be caused by device crossing and post-procedure, 17 per-vascular stenosis has been observed on the stent design because of incomplete diffusion or rigid expansion of the stent (EPW-0081) because of the stent's structure. In previous studies with stents with a closed cell design,¹⁰

The closed-cell stent design, which is the principle for treatment of stenosis, provides the stent with a closed-cell design and thus high flexibility and conformability like an open-cell stent.¹¹ The idea of using a closed-cell stent with double radial force during expansion of the stent results with respect to its mechanical behavior based on the closed-cell design.^{12,13} This stent provides the clinical results

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"CGuard EPS is an important new treatment option for both symptomatic and asymptomatic carotid artery stenosis patients."

C. Wissgott, MD



The Iron-Guard Registry

- Physician initiated
- 12 large Italian medical centers
- 200 patients

Clinical Results

- **100% success** in implanting the CGuard EPS
- **No major adverse cerebrovascular cardiac events** at 30 days
- DW-MRI performed in 61 of 200 patients found **only 19% new lesions** between 24-72 hours
 - CARENET reported 37% new lesions in 30 patients
 - PROF1 reported 66% new lesions in 62 patients

SPECIAL ARTICLES

J CARDIOVASC SURG 2015;56:787-91

Physician-initiated prospective Italian Registry of carotid stenting with the C-Guard mesh-stent: the IRON-Guard registry. Rationale and design

C. SETACCI¹, F. SPEZIALE², G. DE DONATO³, P. SIRIGNANO⁴,
F. SETACCI¹, L. CAPOCCIA⁵, G. GALZERANO⁶, W. MANSOUR⁷
On behalf of IRON-Guard Study Group.



"The IRON-Guard Registry shows promising results in this interim analysis with a low incidence of complications and the lowest reported rate of new MRDWI lesions

F. Speziale, MD and P. Sirignano, MD

Proven success with regionally strong local distributors; YOY sales growth 67%

Representative Regional Distributors

Market	YOY Growth
Colombia	100%
Israel	151%
Italy	59%
Slovenia	95%
Chile	233%

- Replacing Penumbra with regional distributors who target all 4 clinical specialties;
 - Penumbra's focus was primarily the interventional neuroradiology market, their key customer segment
- Focus on larger markets – Germany, Italy, and Spain
- Advanced discussions with distributors in Sweden, Poland, Belgium, Netherlands and Portugal
- Successfully attracting KOLs in each of the respective markets



PD Dr. Andrej Schmidt and Dr. Sven Bräunlich from the Department of Angiology, University Hospital Leipzig, Division of Interventional Angiology, Leipzig, Germany, perform a live stent endovascular interventional procedure featuring the CGuard™ EPS Carotid Stent at the Leipzig Interventional Course (LINC) 2017



* Planning & Development Phase

CGuard™

- Engaging distribution partners in countries with current/near-term regulatory approval
- Attracting leading KOLs from around the world
- Seeking additional regulatory approvals in countries that accept CE Mark
 - Recent approval in Russia with plan to launch in first half of 2017
- Plan to file US FDA IDE in 2017*
- Plan to file CE Mark for next generation 5 French CGuard™ in second half of 2017*
- Partnership strategy targeting Asia Pacific region
 - CAS is the preferred treatment of carotid artery disease in China
 - Targeting distributors in Hong Kong, Taiwan, South Korea, Japan, and China

MGuard

- Strategy focused on formation of strategic partnerships with stent manufacturers with approved drug eluting stents

CGuard™ Approved Markets

Argentina	Ireland
Austria	Israel
Belarus	Italy
Belgium	Latvia
Chile	Lithuania
Colombia	Liechtenstein
Croatia	Luxemburg
Cyprus	Malta
Czech Republic	Norway
Denmark	Poland
Estonia	Portugal
Finland	Romania
France	Russia
Germany	Slovakia
Greece	Slovenia
Holland	Spain
Netherlands	Sweden
Hungary	Switzerland
Iceland	United Kingdom

* Subject to receipt of sufficient funding



Continued market execution and revenue growth.

* Subject to receipt of sufficient funding

Leadership



Significant track records of success

Dr. James Barry	President and CEO				
Craig Shore	CFO				
Agustin Gago	CCO				
Dr. Sol Barer	Chairman				
Isaac Blech	Vice Chairman				
Michael Berman	Director				
Paul Stuka	Director				
Dr. Campbell Rogers	Director				
Thomas Kester	Director				

NYSE MKT: NSPR

Stock Price (2/15/17):	\$2.35
Average 3 Month Volume (2/15/17):	178 K
Shares Outstanding (12/31/16):	1.5 M
Shares Outstanding Including full conversion of preferred shares and anti dilution adjustments (12/31/16):	8.7 M
Market Capitalization including full conversion of preferred shares and anti dilution adjustments (2/15/17):	\$20.4 M
Total Cash (12/31/2016):	\$7.5 M
Headquarters:	Tel Aviv, Israel
# of Employees (2/15/2017)	34



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