

NYSE MKT: NSPR

August 2016

InspireMD



Pioneering fully integrated embolic prevention systems and other advanced medical technology for vascular procedures

NYSE MKT: NSPR

Stock Price (8/5/16):	\$0.19
52 Week Range:	\$0.17-\$2.5
Average Volume:	1.1 M
Shares Outstanding (8/5/16):	29.6 M
Shares Outstanding Including Future Pref. Stock Conv. (8/5/16):	90.3 M
Market Capitalization (8/5/16):	\$17.2 M
Analyst Coverage:	H.C. Wainwright: Yi Chen Empire Asset Management: Cathy Reese
Total Cash (6/30/16):	\$0.9 M
US Headquarters:	Boston, MA
International Headquarters:	Tel Aviv, Israel
# of Employees (8/5/2016):	32

Investment Highlights



Focused on commercial execution of CGuard™ EPS and development of pipeline products

- 2016 focus on revenue growth driven by a broader EU launch of CGuard
 - Strategic distribution partnership with Penumbra (NYSE:PEN)
 - Significant growth in Italy over the last 3 quarters serving as a leading indicator
 - Positive clinical trial results using CGuard in a broad patient population, including high risk patients
- Advancing into the growing neurovascular and peripheral vascular markets
 - 2017E CE Mark Submission for **NGuard**[™] flow diverter for treatment of cerebral vascular aneurysms enabling EU commercialization post approval
- A broad portfolio of assets supported by aggressive pursuit of intellectual property protection
- Well positioned for strategic collaboration on multiple MicroNet[™] product applications
- Continued financial discipline in line with development and growth initiatives

The Problem



Embolization can lead to catastrophic health events

THE WALL STREET JOURNAL. $\equiv | u.s.$



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Stents Boost Stroke Recovery, Study Finds

Using Devices to Pull Clots From Brain Arteries Can Help Patients

By THOMAS M. BURTON

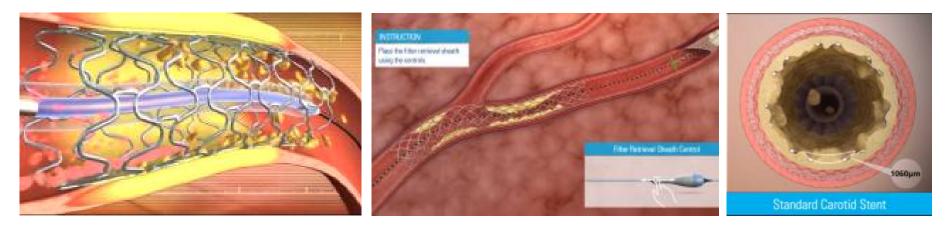
Using a device to extract blood clots from brain arteries can significantly improve patients' ability to rebound from a stroke, according to a landmark study published Wednesday.







"Plaque protrusion through stent struts occurs in up to 65.5% 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.**

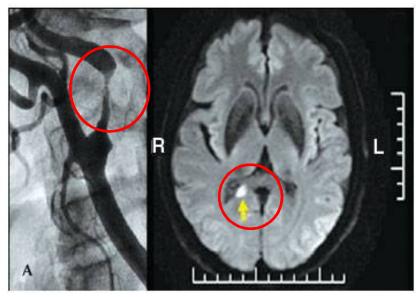
* Musialek, et.al. Eurointerventions 2016;12 published online ahead of print May 2016

** Bosiers et al. Eur J Vasc Endovasc Surg Vol 33, Feb 2007

The Consequences

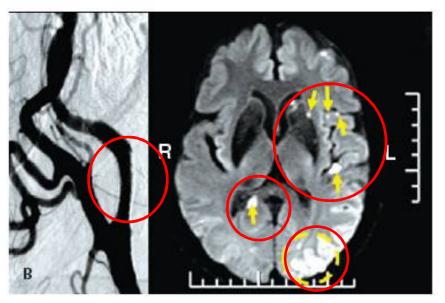


Range from neurological deficit to stroke to death



Pre-Procedure

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



Post-Procedure

B. 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.

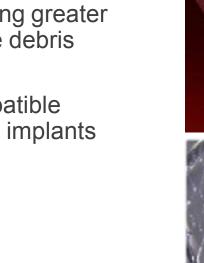
MicroNet[™]

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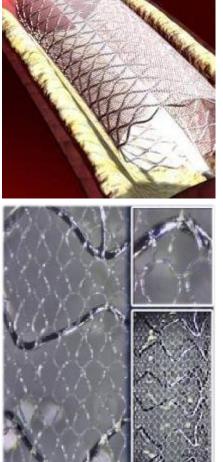
Proprietary technology for preventing distal embolization and other vascular disease challenges

Ultrathin PET mesh provides meaningful clinical benefit* to conventional devices

- Provides revascularization benefit •
- MicroNet acts as "safety net" by offering greater vessel area coverage to prevent large debris flow through the scaffold
- Made of a single fiber from a biocompatible polymer, widely used in other medical implants



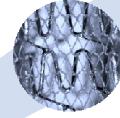




Large Addressable Market



Expanding the MicroNet[™] Platform



CGuard™

- ✓ \$500M Market
- ✓ CE Mark Cleared
- ✓ FDA IDE draft protocol
- synopsis
- ✓ Carotid

NGuard

- ✓ \$125M Flow Diversion Market
- ✓ \$550M Aneurysm Market
- ✓ 2017E CE Mark Planned Submission for Flow Diverter
- ✓ Neurovascular

PVGuard

- ✓ \$1.7B Market
- ✓ 2018E CE Mark Planned
- Submission
- ✓ Peripheral



MGuard™* ✓ \$1.7B AMI Market ✓ CE Mark Cleared

✓ Coronary AMI, SVG

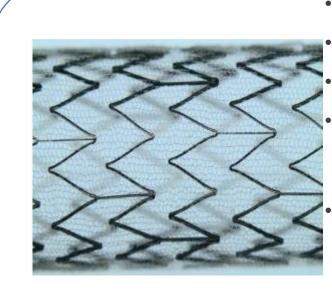
Carotid Solution: Mesh Covered Technology



An emerging market opportunity

CGuard[™] Embolic Prevention System(EPS)

Combines stent and embolic protection in a single device



- CE marked
- Self-expanding nitinol stent
- Global market valued at \$500M*
- Positive CARENET data released 9/14, 1/15 and 5/16 documenting the safety and patency of the CGuard EPS
- Positive all-comer data from PARADIGM trials presented in May 2016 at EuroPCR documenting the safety and benefits of the CGuard EPS
- Ongoing launch in Europe, Latin America, South America, & other regions

Positive CGuard[™] Clinical Experience InspireMD



CARENET Clinical Trial

- 30 Patient Safety and Efficacy clinical trial •
- Zero major adverse cardiac or cerebral events (MACCE) at 30 days (Comparative data 5.72%*) ۰
- 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 full resolved at 30 days except one •
- 3.6% MACCE rate at 6 months (Comparative data 8.09%**) ۰
- Zero strokes or stroke related deaths at 12 months

PARADIGM 101 Clinical Trial

101 patient trial evaluating CGuard EPS in unselected, consecutive carotid patients (all-comers)

- 99.1% device success •
- 0% MACCE (Death/stroke/MI) @ 48 hr ٠
- 0% MACCE @ 30 day as determined by independent neurological and angiographic evaluation ٠



"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

Carotid Market Opportunity



"Game Changing" Minimally Invasive Solution

- Current standard of care: Surgery
 - Carotid Endarterectomy (CEA)
- The risk of post-procedural cerebral events has been related to [conventional] carotid stents.¹
 - Higher risks of stroke at 10 years appear to be attributable to the peri-procedural differences in risk.²
 - Mesh-covered carotid stents may lower the rates of peri-procedural stroke.²
- CGuard[™] clinical studies have demonstrated superior safety
 - CARENET
 - PARADIGM
 - PARADIGM 101
- Immediate EU commercial opportunity (non-US)
 - EU pursued via new strategic partner Penumbra
 - Europe, Latin America and other regions are covered by experienced distributors
 - U.S. development and clinical plan to follow



"The most important theme during [EuroPCR 2016] was carotid artery stenting....[The double layered mesh stents] will resolve the main problem of carotid artery stents which was late embolic events." A. Cremonesi

¹ Musialek, EuroIntervention 2016:12 online publish ahead of print May 2016

² Brott, T. Long-Term Results of Stenting versus Endarterectomy for Carotid-Artery Stenosis, New England Journal of Medicine, March 17, 2016

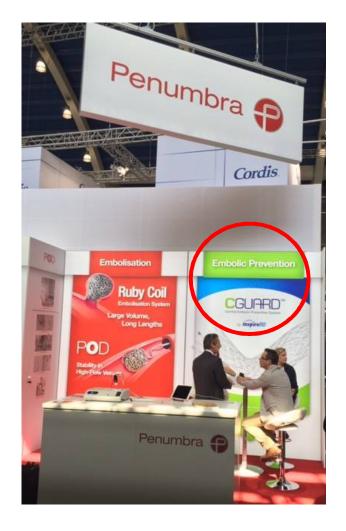
Strategic Distribution Partnership



Broad European commercialization support from a growing neurovascular leader



- Distribution agreement with Penumbra
- 18 European markets with opportunity to expand
- Comprehensive neurovascular product portfolio
- CGuard is a synergistic product offering
- Growing direct sales force throughout Europe
- Establishing a direct sales force focused on peripheral vascular

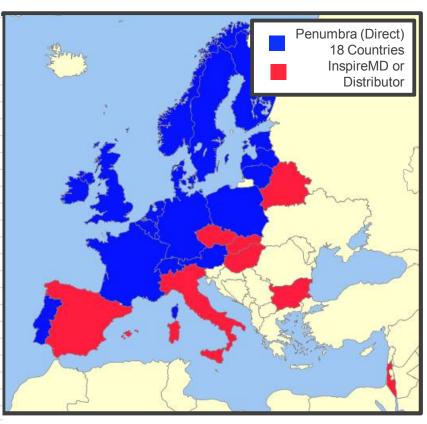


CGuard Distribution (EU)



	Sales Channel	Q115	Q215	Q315	Q415	Q116	Q216E	Q316 E
UK	Р			OLT		HOC	IND	
France	Ρ			OLT	HOC		IND	
Belgium	Ρ			OLT	HOC		IND	
Netherlands	Ρ			OLT		HOC	IND	
Luxemburg	Ρ			OLT		HOC	IND	
Norway	Ρ			OLT			HOC	IND
Sweden	Ρ			OLT			HOC	IND
Denmark	Ρ			OLT			HOC	IND
Finland	Ρ			OLT			HOC	IND
Germany	Ρ					OLT	HOC	IND
Austria	Ρ					OLT	HOC	IND
Switzerland	Ρ					OLT	HOC	IND
Portugal	Ρ			OLT	HOC	IND		
Poland	Ρ			OLT	HOC	IND		
Latvia	Ρ			OLT		HOC	IND	
Lithuania	Ρ			OLT		HOC	IND	
Estonia	Ρ			OLT		HOC	IND	
Italy	СК	OLT	HOC	IND				
Israel	IN	OLT	HOC	IND				
NOTE: (P) Penur	nbra, CK	(Crossmed/Ka	ster), IZ (Izasa),	, IN (InspireMD/I	Distributor)			
	-			-				
_	OLT	Online		HOC	•Hands On Training •Case			endent

Observation

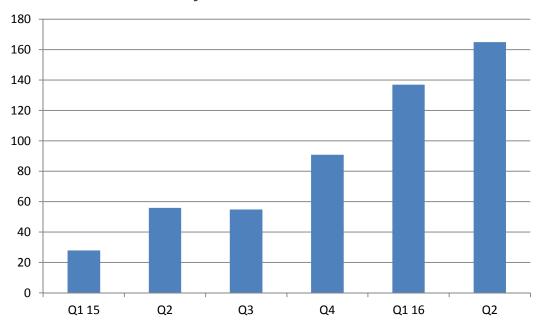


CGuard[™] Country Case Study



Italy – A leading indicator of CGuard Growth

- CGuard covered by 2 distributors
- Initial success drove 29 Italian carotid interventionalists to initiate the IRON-Guard* registry last year



Italy CGuard Revenue

Robust Pipeline



Expanding Indications with MicroNet[™] Commercial CE Mark Cleared 12-month MASTER I, II MGuard™ Coronary Commercial EPS · CE Mark Cleared CARENET Trial **Exploring Market** RGuard™ CGuard™ PARADIGM all-comer trials **Opportunity Renal*** • FDA IDE draft protocol Carotid synopsis Penumbra Strategic MicroNet™ Partnership **Platform** Technology **Exploring Market** 2018E development towards **Opportunities: PVGuard**[™] NGuard™ **CE Mark Planned** Flow Diverter **Peripheral* Neurovascular*** Intra-Cranial Stent Submission 2017E CE Mark Planned Submission MGuard™ for Flow Diverter Drug Eluting* Strategic partner opportunity

Neurovascular Aneurysms

Flow Diversion

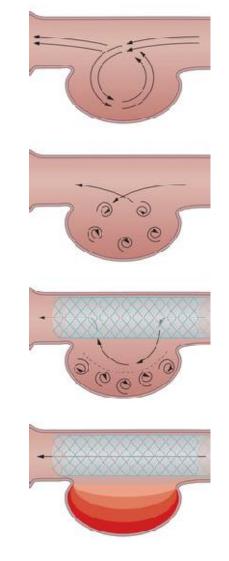
Objective

• Seal the aneurysm and prevent rupture

Current device therapies

- Coils to pack the aneurysm
- Flow diverters
 - Highly flexible, dense metal "tube"
 - Placed in main artery to seal off aneurysm and cause aneurysm thrombosis
 - Precise delivery required to avoid blocking other vessels





Neurovascular Market Opportunity

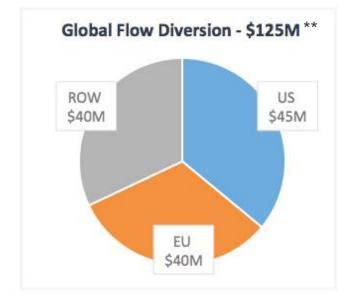


Innovation leads growth

Flow Diversion For Unruptured Brain Aneurysms

Next Generation Technology

- Aneurysm Therapy (all types): \$550M*
- Flow diverters are estimated to be 25% of the aneurysm market
- Neurovascular products: estimated 15% CAGR from 2010-2016

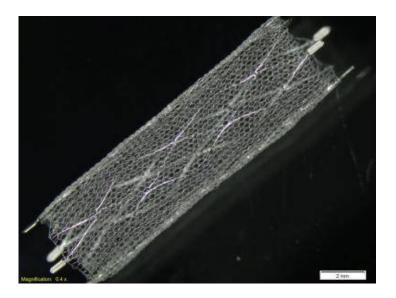


2014 Competitive Landscape: Relatively Fewer Players					
Product	Company	Approval			
Pipeline	Medtronic/Covidien	CE Mark 2014 FDA 2011			
Surpass	Stryker	CE Mark 2010			
Silk	Balt Extrusion	CE Mark 2008			

* 2013 MRG Neuro Report, 2010 Ev3 Revenue Data ** 2014 projection based on 2013 actuals

InspireMD Flow Diverter Advantage





- Low profile, flexible, open cell scaffold = Easy to delivery
- Low metal ratio = Potential for reduced anti-thrombosis medication
- Re-accessible through MicroNet[™] = Allows for further treatment, if needed which is impossible with current flow diverters
- Can be placed in side branches and bifurcations = Will not block blood flow into major side vessels, which is impossible with current technology
- Published success with MicroNet in coronary and carotid aneurysms



PATENT RIGHTS	Issued	Allowed	Pending
US	4	0	11
Rest of World (ROW)	16	0	13

Continue to strengthen and broaden patent protection globally. Progress over the last year imparts important rights on existing products and technologies and will enable future pipeline products.

Leadership



Significant Track Records of Success

EXECUTIVE TEAM

Dr. James Barry, President and CEO

- Boston Scientific
- Howmedica Division of Pfizer

Craig Shore, CFO

- Pfizer
- General Electric

David Blossom, VP Global Marketing & Strategy

- Boston Scientific
- Covidien

BOARD OF DIRECTORS

- Dr. Sol Barer, Chairman
 - Former Chairman and CEO, Celgene

Isaac Blech, Vice Chairman

• Private financier in the life science industries

Dr. James Barry, President and CEO

- SVP Corporate Technology Development at Boston Scientific
- Howmedica Division of Pfizer

Michael Berman

- Pres. Boston Scientific/Scimed
- Founder, Velocimed
- Director Lutonix

Paul Stuka

- Founder, Osiris
- Fidelity Management and Research

Dr. Campbell Rogers

- CMO, Heartflow
- CSO, Cordis/JNJ
- Associate Professor, Harvard School of Medicine

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