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 1934

Date of Report (date of earliest event reported): September 2, 2009

MIMEDX GROUP, INC.

(Exact name of registrant as specified in charter)

Florida

000-52491

90-0300868

(State or other jurisdiction of incorporation)

(Commission File Number)

(IRS Employer Identification No.)

811 Livingston Court, Suite B, Marietta, GA 30067

(Address of principal executive offices, including zip code)

(678) 384-6720

(Registrant's telephone number, including area code)

1234 Airport Road, Suite 105 Destin, Florida 32541

(Former address)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of registrant under any of the following provisions:

- 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 2 — Financial Information

Item 2.02 Results of Operations and Financial Condition.

The following information is intended to be furnished under item 2.02 of Form 8-K, "Results of Operations and Financial Condition." This information shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or incorporated by reference in any filing under the Securities Act of 1933, as amended (the "Securities Act"), or the Exchange Act, except as shall be expressly set forth by specific reference in such a filing.

The Company is planning to give a presentation to prospective investors. The presentation discusses financial projections as well as strategic and tactical goals. The presentation is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

Section 7 — Regulation FD

Item 7.01 Regulation FD Disclosure

The information furnished under Item 2.02 of this Form 8-K is hereby incorporated by reference into this Item 7.01.

Section 9 — Financial Statements and Exhibits

Item 9.01 Financial Statements and Exhibits

(d) Exhibits

Exhibit	Description
99.1	Presentation to Prospective Investors

SIGNATURES

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

Dated: September 2, 2009 MIMEDX GROUP, INC.

By: /s/ Michael J. Culumber

Michael J. Culumber, Acting Chief Financial

Officer

EXHIBIT INDEX

Exhibit	Description	
99.1	Presentation to Prospective Investors	



INNOVATIONS IN BIOMATERIALS









COMPANY PRESENTATION CONFIDENTIAL



FORWARD-LOOKING STATEMENT

This document contains forward-looking statements within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934. Such statements may be identified by their use of terms or phrases such as "expects," "estimates," "projects," "believes," "anticipates," "plans," "intends," and similar terms and phrases. Forward-looking statements are based upon the current beliefs and expectations of our management and are inherently subject to risks and uncertainties, some of which cannot be predicted or quantified, which could cause future events and actual results to differ materially from those set forth in, contemplated by, or underlying the forward-looking statements. For example, the Company (i) may be unable to continue its efforts on particular products due to future laboratory results, (ii) may be unable to obtain needed additional capital on acceptable terms, if at all, or meet its financial forecasts, (iii) may be unable to maintain, retain and develop its intellectual property, (iv) must address regulatory, competitive and marketing challenges in the course of developing its products. These and other risks are described in our reports filed with the Securities and Exchange Commission, including our Form 10-K filed June 15, 2009, as subsequently amended. We disclaim any obligation to update or revise any forward-looking statements to reflect actual results or changes in the factors affecting the forward-looking information.

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INNOVATIONS IN BIOMATERIALS

MiMedx Group, Inc., (OTCBB: MDXG) is an integrated developer, manufacturer, and marketer of patent protected biomaterial-based products.

- •In 2009, MiMedx Group expects to transition from a pre-revenue, developmentfocused concern to an operating company focused on sales growth and profitability
- •MiMedx Group holds exclusive rights to 8 issued and 33 pending U.S. and foreign patent applications
- •MiMedx Group went public through a reverse merger with an inactive entity in February 2008
- •MiMedx Group has recently restructured the management team with the addition of experienced and proven healthcare executives

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Board of Directors

Parker H. "Pete" Petit, Chairman

Management Team

Pete Petit President & CEO
Bill Taylor COO (acting)
To Be Determined EVP Sales

Mike Culumber Chief Financial Officer
Roberta McCaw General Counsel
Matt Miller EVP Business Dev.



SpineMedica

Collagen-Based Biomaterials Durable Hydrogel Biomaterials

Thomas Koob, PhD, CSO Based in Tampa, FL Rebeccah Brown, PhD, COO Based in Atlanta, GA

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BOARD OF DIRECTORS

Parker H. (Pete) Petit — Chairman

- Former Chairman and CEO of Matria Healthcare which was sold to Inverness Medical Innovations in May, 2008
- Matria Healthcare was a former subsidiary of Healthdyne, which Mr. Petit founded in 1970
- Served as Chairman and CEO of Healthdyne and some of its publicly traded subsidiaries after Healthdyne became a publicly traded company in 1981
- · For further information, see www.thepetitgroup.com

Steve Gorlin

- Founder of MiMedx Group, Inc.
- Founder of several biotechnology/pharmaceutical companies, including Hycor Biomedical, Theragenics (NYSE:TGX), CyrRx (Nasdaq:CYTR), Medicis Pharmaceutical (NYSE:MRX), EntreMed (Nasdaq:ENMD), Surgi-Vision, DARA BioSciences, (Nasdaq:DARA), SpineMedica and Medivation (Nasdaq: MDVN)

Charles Koob

- Successful M&A, Anti-trust, Corporate attorney at Simpson Thatcher and Bartlett
- Experience includes the defense of Appleton Papers, representation of Virgin Atlantic Airways, and the representation of a special committee of Archer Daniels Midland in a federal grand jury investigation of price-fixing

Larry Papasan

- Venture Partner at MB Venture Partners
- Former President of Smith & Nephew Orthopedics
- Chairman at BioMimetic Therapeutics (Nasdaq:BMTI), Board Member at Memphis Biotech Group and Plough Foundation
- Chairman at Lebonheur Children's Hospital Foundation, Pres. of the Board at the Biblical Resource Center & Museum

Kurt Eichle

- Serves as an EVP of LCOR Incorporated, a real estate investment and development company, in charge of operations in the metropolitan New York region
- Formerly in the Real Estate Debt and Equity Finance Group of Merrill Lynch, Hubbard
- · Member of the Board of Governors, Real Estate Board of NY

Kreamer Rooke

- · Initial investor in MiMedx
- · Advised on the acquisition of SpineMedica Corp.
- Partner at the Gorlin Companies
- Former Investment Banking Analyst in the Global Healthcare Group of Collins Stewart, Inc.
- Graduate of the University of Pennsylvania: B.A. in Economics

Joseph G. Bleser (under consideration to fill a vacancy)

•Experienced Healthcare Exec, CFO, Board member, Audit Chair

•Over 20 years as CFO of Public Healthcare & Technology companies

Companies include: HBO, Healthdyne Information Enterprises,
Allegiant Physician Svs, Healthcare.com, Transcend Services
BOD: Transcend Services (NASDAQ: TRCR), Matria Healthcare
Licensed CPA with 10 yrs public accounting for international firm

Terry Dewberry (under consideration to fill a vacancy)

- ·Experienced Healthcare Exec, Board member, Audit Chair
- ·Previously Director of Respironics, and Matria Healthcare
- •Currently on Board of DrTango, a private multicultural
- communications and health management company
- Previously, Director, Vice Chairman, President and COO, and CFO of Healthdyne, 1981 1996
- •Experience buy & sell side Corp M&A, transactions up to \$5 billion

COMPANY PRESENTATION CONFIDENTIAL



CONSOLIDATED PRO FORMA INCOME STATEMENTS

		Cale	ndar Year	
	2009	2010	2011	2012
Net Revenue	105	8,876	21,281	36,000
Cost of Sales	1,648	3,272	5,345	7,884
Gross Profit	(1,543)	5,604	15,936	28,116
Total Operating Expenses	5,765	8,288	12,564	19,097
EBITDA	(7,308)	(2,684)	3,372	9,019
Depreciation & Amortization	1,152	1,384	1,691	1,800
Operating Margin	(8,460)	(4,068)	1,681	7,219
Interest Income, Net	(69)	(88)	(92)	(100)
Other Income (Expense)	565			
Pretax Profit	(7,964)	(4,156)	1,589	7,119
Income Taxes		-		
Net Earnings	(7,964)	(4,156)	1,589	7,119
Weighted Average Shares (Fully Diluted)	46,887	56,540	61,040	61,040
EPS	\$ (0.17)	\$ (0.07)	\$ 0.03	\$ 0.12



CONSOLIDATED PRO FORMA INCOME STATEMENTS

	2009 Plan					2010 Plan				2011 Plan					
	Q1	Q2	Q3	Q4	YTD	Q1	Q2	Q3	Q4	YTD	Q1	Q2	Q3	Q4	YTD
MiMedx															
Revenues					-		480	925	1,875	3,280	2,427	2,843	3,225	3,584	12,079
Gross Profit	(258)	(206)	(95)	(150)	(709)	(190)	146	595	1,451	2,002	1,927	2,147	2,466	2,741	9,281
Operating Profit	(816)	(763)	(975)	(1,011)	(3,565)	(892)	(650)	(364)	241	(1,665)	492	565	721	896	2,674
SpineMedica															
Revenues				105	105	632	1,340	1,719	1,905	5,596	2,084	2,216	2,364	2,540	9,204
Gross Profit	(197)	(144)	(207)	(129)	(677)	308	911	1,220	1,378	3,817	1,535	1,645	1,770	1,920	6,870
Operating Profit	(676)	(620)	(698)	(752)	(2,746)	(406)	(55)	157	403	99	364	321	455	586	1,726
MiMedx Group Overhea	d														
Revenues			-	-		-	-	-		-	-	-			
Gross Profit		(52)	(52)	(52)	(156)	(53)	(54)	(54)	(54)	(215)	(54)	(54)	(54)	(54)	(216
Operating Profit	(541)	(675)	(467)	(467)	(2,150)	(720)	(648)	(588)	(545)	(2,501)	(623)	(629)	(737)	(730)	(2,719
Consolidated MiMedx															
Revenues				105	105	632	1,820	2,644	3,780	8,876	4,511	5,059	5,589	6,124	21,283
Gross Profit	(455)	(402)	(354)	(331)	(1,542)	65	1,003	1,761	2,775	5,604	3,408	3,738	4,182	4,607	15,935
Operating Profit	(2,033)	(2,058)	(2,140)	(2,230)	(8,461)	(2,018)	(1,353)	(795)	99	(4,067)	233	257	439	752	1,681
Interest Expense	1	(20)	(25)	(25)	(69)	(22)	(22)	(22)	(22)	(88)	(23)	(23)	(23)	(23)	(92
Other Income (Expense)		565	-	1-21	565	-	-	-	-	-	-	-	-	-	-
Net Income	(2,032)	(1,513)	(2,165)	(2,255)	(7,965)	(2,040)	(1,375)	(817)	77	(4,155)	210	234	416	729	1,589

COMPANY PRESENTATION CONFIDENTIAL



STRATEGIC and TACTICAL GOALS

STRATEGIC

•Create a fast growth and very profitable biomaterials company from the MiMedx Group's diverse intellectual property

TACTICAL

- •Exploit intellectual property through internal product development as well as strategic partnerships
- Develop an independent sales representative national group as the company's primary product distribution outlet
- •Carefully select a minimum number of larger orthopedic device manufacturers as joint product development partners who could also serve as distribution outlets
- •Over time, add direct sales persons to supplement the sales rep groups
- •Keep management structure, controls and processes maturing ahead of the revenue growth profile
- •Maintain excellent Wall Street rapport with analysts and brokerage houses

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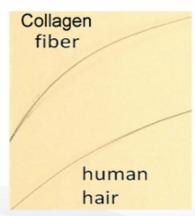


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Collagen Technology with Exceptional Mechanical Performance and Biocompatibility for Connective Tissue Repair





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MIMEDX COLLAGEN: A BIOMATERIAL SCAFFOLD

MiMedx's collagen-based biomaterial products act as platforms or scaffolds on which a patient's tissue grows during the healing process in order to achieve *healthy* tissue reconstruction without significant scarring.

While the patient's own cells are fabricating their own natural structure, MiMedx implants provide structural integrity, promote better and faster healing and are eventually absorbed, being replaced by newly formed, competent, healthy tissue rather than scar tissue.

Our collagen fiber is intended to facilitate the repair by:

- Providing mechanical integrity during the healing process
- •Biodegrading and being absorbed by the surrounding tissue without insult or future surgical intervention

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PRODUCT CHARACTERISTICS

- Collagen is cross-linked using a proprietary and patented process
- Although intended only to augment soft tissue during repair, our collagen fibers are stronger human tendons and ligaments
- Approximately same stiffness as human tendons and ligaments
- · Forms a scaffold when implanted which allows for cell in-growth
- Biological—entirely organic materials, no synthetics
- Biodegradable—products are absorbed by the body during and after healing occurs
- · Manufactured as continuous fibers

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PRODUCT POTENTIAL USES

- · Orthopedic soft tissue augmentation and repair—tendons and ligaments
 - o Repair for shoulder & rotator cuff, hand & elbow, knee, hip
 - Repair for foot and ankle—posterior tibial, peroneal, extensor and lateral tendons
- · General surgery & wound care
 - Diabetic foot ulcers, plantar soft tissue augmentation, tendons and ligaments of the foot
- · Gynecological repair and organ support
 - o Urethral slings, vaginal prolapse repair
- Neurological surgery
 - Nerve wraps
- Plastic and reconstructive surgery
 - Soft tissue support, cranial maxillofacial reconstruction

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PRODUCT OPPORTUNITIES

Orthopedics and Connective Tissue Reconstruction

- · Tendon and ligament augmentation during repair
- · Sutures for tying tissue
- · Tubes for anchoring to bone

Wound Repair

- · Patches for wound healing which immediately covers and protects against infection
- · May act as a scaffold for full repair and healing

Other Opportunities

- Slings or supports for gynecological uses
- · Patches for hernia repair and other general soft tissue repair

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PRODUCT COMPETITION

Tendon Protectors

 TenoGlide™ by Integra Life Sciences and TenoMend™ by Collagen Matrix: collagen wraps for protection of repaired tendons

Tendon Augmentation or Repair Devices

- OrthAdapt® by Synovis/Pegasus: a collagen soft tissue scaffold in the form of a patch for reinforcement of rotator cuff and Achilles tendon repair
- GraftJacket® by Wright Medical: a collagen patch used for connective tissue repair and wound management
- FiberTape™ and FiberWire® by Arthrex: synthetic materials for reattaching all severed tendons and ligaments

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ADVANTAGES OVER THE COMPETITON

Potential advantages over TenoGlide and TenoMend:

- · Our products will also supply protection to the repair site
- Our products will go beyond simple repair protection by augmenting the repair

Potential advantages over GraftJacket, OrthAdapt, and FiberWire:

- Scalability: we can target fiber geometries to match the anatomy, GraftJacket and OrthAdapt are only offered as a patch
- Biologic: FiberWire is made of polyester, a synthetic material that can cause inflammation and is a permanent implant
- Stiffness: we can match native tissue stiffness, independent of implant size
- Strength: depending on size, 5x to 10x stronger than GraftJacket or OrthAdapt
- Biocompatible: novel cross-linking technology has no associated inflammatory response while degrading

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PROPRIETARY TECHNOLOGY

Patent No. 6,565,960—A broad process patent protects the technology

<u>Our collagen-fiber cross-linking patent is strong and broad: no competitor has a collagen fiber as strong, stiff, expandable, scalable, biocompatible and biomimetic, from a biological source.</u>

All patents relating to specific products, such as an Achilles tendon augmentation device, are available for the company to file, at which time we will have protection for the *product's* entire patent-lifespan.

Patents Issued: 2 Patents Pending: 10

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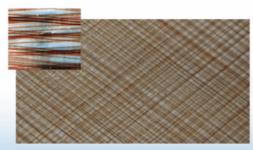
PRODUCT CONFIGURATIONS



Cables



Tubes



Patches

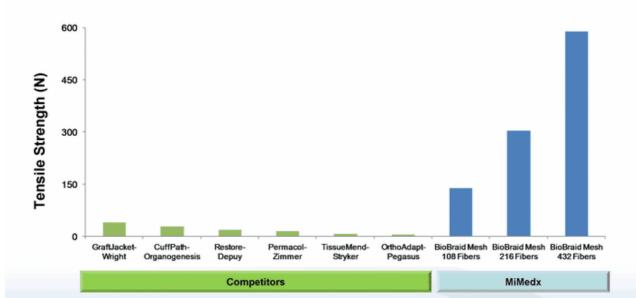


Knits

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SUPERIOR STRENGTH VERSUS COMPETITORS

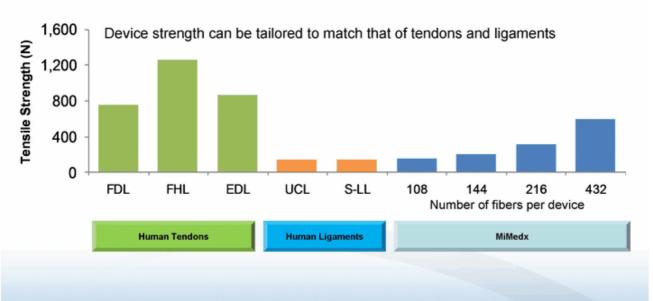


Individual values taken from published & Company literature

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STRENGTH COMPARISON TO NATIVE TISSUE

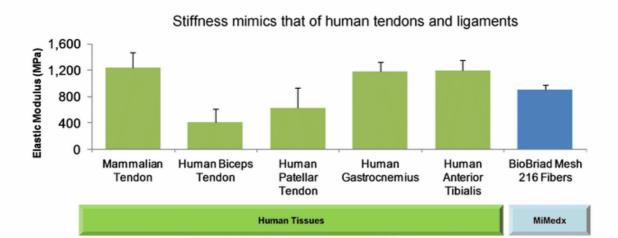


Data from published & Company literature

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STIFFNESS COMPARISON TO NATIVE TISSUE



Data from published & company literature

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MIMEDX PHYSICIAN ADVISORY BOARD

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- •The Andrews Institute; Gulf Breeze, FL

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•The Andrews Institute; Gulf Breeze, Florida

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•Kerlan Jobe Orthopedic Center; Los Angeles, CA

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·Beacon Orthopedics; Cincinnati, OH

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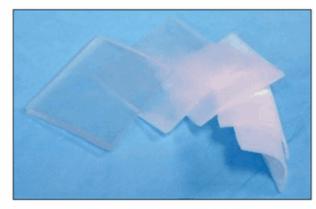
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DURABLE, BIOCOMPATIBLE HYDROGEL TECHNOLOGY

- Durable hydrogel products have numerous opportunities with initial product focus in sheet applications
- Sheets will have broad market opportunities
- First product to the market is the Paradis Vaso Shield™

The Paradis Vaso Shield™ is FDA cleared for use as a cover for vessels following anterior vertebral surgery.



Paradìs Vaso Shield™

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PRODUCT CHARACTERISTICS

- · Silicone-like material with properties that can be tailored
- · A durable hydrogel compound made of polyvinyl alcohol
- · Prohibits cellular attachment or in-growth
- · Patented and proprietary manufacturing processes
- Biocompatible
- Durable

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PRODUCT POTENTIAL USES

- · Orthopedic Spine Repair
 - FDA cleared as a cover for vessel following anterior vertebral surgery
- Possible cervical and thoracic anterior and full spine posterior vertebral surgery
- · Future Product Uses
 - o Cardiovascular and open heart surgery
 - General surgery
 - o Hernia repair
 - Plastic and reconstructive surgery
 - Obstetrics & gynecological surgery

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PRODUCT OPPORTUNITIES

Orthopedics

- · Currently FDA cleared as a vessel cover following anterior vertebral surgery
- · Providing a plane of dissection for surgeon accessibility for revision surgery
 - o Expedites the surgical process
 - o Reduces risk to patient during revision surgery

Foot and Ankle

- · Cushion device
- · Wound covering

Other Opportunities

· May possibly act as an anti-adhesion patch

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PRODUCT COMPETITION

Preclude® Vessel Guard by W.L. Gore

- · Only direct competitor
- · Similar FDA indication to Vaso Shield
- · Product is opaque
- Made of ePTFE (expanded-PolyTetraFloroEthylene)

Seprafilm® by Genzyme

- · FDA indication as an anti-adhesion barrier in open abdominal surgery
- · Sometimes is used off-label for anterior vertebral surgery
- · Resorbs over time

Interceed® by Ethicon/J&J

- · FDA indication for gynecological use
- Acts as a barrier following surgery
- · Sometimes is used off-label for anterior vertebral surgery
- · Resorbs over time

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ADVANTAGES OVER THE COMPETITON

Potential advantages of Vaso Shield over Preclude Vessel Guard:

- Translucent
 - Surgeon can see through implant
 - o Makes suturing easier
 - Makes implanting faster
 - Makes using safer
- · MRI-clearly visible on MRI, Preclude is not
- · Thickness—easier to handle and does not fold over on itself
- · Thickness—easier to feel edges and find during revision surgery

Potential advantages over Interceed and Seprafilm:

- The Vaso Shield is FDA cleared for use as a cover for vessels following anterior vertebral surgery
- Vaso Shield provides permanent and predictable protection
- · Interceed and Seprafilm resorbe

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PROPRIETARY TECHNOLOGY



Hydrogel devices have been used in the knee and foot for several years

Durable Viscoelastic Polymer

- · Own and Licensed IP for products and manufacturing
 - 6 Issued, 23 pending patents securing technology—US & Foreign
 - Material texture is similar to silicone without associated negative properties, such as silicone wear debris, biocompatibility issues
- · Mimics properties of human tissue
- · Biocompatible & highly durable
- · Economical to produce
- Wide variety of proprietary manufacturing methods
- · Easily integrated with other materials
- · Proven pre-clinical & clinical performance
 - Knee Cartilage (EU) & Nerve Cuff (US)

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VASO SHIELD

FDA Cleared 510(k) as a vessel cover following anterior vertebral surgery

- Visible on MRI
- ·Sutured into place





Vaso Shield being implanted



First implant procedure

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