

Investor Presentation August 2013

FORWARD LOOKING STATEMENT

This presentation contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. These statements include, but are not limited to, the market opportunity for and the market acceptance of the Company's products, the potential uses for the Company's tissue products, the expected growth in revenue and customer base, the availability of third-party reimbursement for the Company's products (whether by Regional Macs or private health insurers), and projected revenues. These statements are based on current information and belief, and are not guarantees of future performance. Our ability to predict results, financial or otherwise, or the actual effect of future plans or strategies is inherently uncertain and actual results may differ from those predicted depending on a variety of factors. Among the risks and uncertainties that could cause actual results to differ materially from those indicated by such forward-looking statements include that the *Company's products may not gain the anticipated acceptance in the marketplace or that acceptance* may be delayed; the effects of competition; delays or changes in reimbursement for the Company's products, delays in clinical trials or unexpected results, and the risk factors detailed from time to time in the Company's periodic Securities and Exchange Commission filings, including, without limitation, its 10-Q filing for the fiscal quarter ended June 30, 2013. By making these forward-looking statements, MiMedx Group, Inc. does not undertake to update those in any manner except as may be required by the Company's disclosure obligations in filings it makes with the Securities and Exchange Commission under the federal securities laws.



INVESTMENT HIGHLIGHTS

MiMedx (NASDAQ: MDXG) is a regenerative biomaterials company focused in delivering technologies that help the body heal itself

- Experienced & Proven management team
- ✓ Proprietary PURION Processed dHACM shown to be a "Stem Cell Magnet"
- ✓ Significant Clinical and Cost Effectiveness with Published Studies
- ✓ Strong I.P. portfolio
 - 8 Amniotic allograft patents issued and 35 pending
 - Over 100 patents issued & pending for all technologies
- ✓ In excess of 170,000 allografts distributed
- ✓ Regulatory Path Human Transplanted Tissue (HCT/P), Section 361 of the Public Health Service Act, FDA regulated under 21CFR1271
- ✓ Reimbursement of EpiFix[®] Code Q4131
 - Six of Nine Medicare Contractors are covering
- ✓ Strong placental recovery network protected by contracts



EXPERIENCED MANAGEMENT TEAM

Parker H. "Pete" Petit Chairman & CEO

Brent D. Miller Executive Vice President

Thomas J. Koob, Ph.D Chief Scientific Officer

Thornton A. Kuntz Vice President, HR & Administration William C. Taylor President & COO

Deborah L. Dean Executive Vice President

Michael W. Carlton Vice President, Global Sales

Rebeccah Brown, Ph.D. Vice President, Product Development & Quality Assurance & Regulatory Affairs Michael J. Senken Chief Financial Officer

Roberta L. McCaw General Counsel

Donald E. Fetterolf, MD Chief Medical Officer

H. Frank Burrows Vice President, Corporate Strategy





portal

HIGHMARK.



ETHICON a Johnson Johnson company



smith&nephew





ILIPS RT

RTI BIOLOGICS"

(B) SYNTHES"

SFACET TECHNOLOGIES.



TECHNOLOGY

- Transplanted amniotic membrane tissue allografts
- Placentas donated from Caesarean section deliveries
- Proprietary PURION[®] process
 - 8 Issued U.S. Patents; 33 patents pending for tissue
 - Trade secrets related to process
- Logistically superior; 5-year shelf life at room temperature storage
- Clinically effective and cost effective, Reduces time and cost to heal
 - Enhances healing
 - Reduces scar tissue
 - Reduces inflammation
 - Non-Immunogenic
- Broad patient and procedure usage to date
 - Over 170,000 grafts distributed
 - Zero tissue-related adverse events





- Chronic wounds
 - Diabetic foot ulcers (DFU's)
 - Venous stasis ulcers (VLU's)
 - Arterial ulcers
 - Pressure ulcers
- Acute/Surgical wounds
 - Burns
 - Plastic surgery
 - Scar revision
- Near term, accelerated ramp

AmnioFix

- Tendon / Ligament wrap
- Peripheral nerve wrap
- Spinal surgery– Dural Repair
- Sports Medicine
- Ortho Surgery
- Prostatectomy
- Tendinopathy & Pain Management injections
- General surgery, gynecology, etc.
- Near to long-term, moderate ramp

Soft Tissue Repair

Not simply dermis or epidermis repair

\$13+ Billion Addressable US Market



SCIENTIFIC PUBLICATION

International Wound Journal

Biological Properties of Dehydrated Human Amnion/Chorion Composite Graft: Implications for Chronic Wound Healing

Researchers from Stanford University School of Medicine and The Georgia Institute of Technology Biotechnology Complex found that:

- PURION[®] processed dHACM retains biologically active growth factors and regulatory factors that are in part responsible for its clinical effectiveness in wound healing.
- dHACM contains one or more soluble factors capable of stimulating mesenchymal stem cell migration and recruitment.
- dHACM is a multifaceted tissue graft that positively affects at least four distinct physiological processes,
 - induce regenerative cell proliferation
 - modulates inflammatory processes
 - inhibit metalloproteinase activity and matrix degradation
 - recruitment of stem cells

"Stem Cell Magnet"



EPIFIX[®] CONTAINS GROWTH FACTORS, CYTOKINES AND ENZYME INHIBITORS

• Over 70 Identified to date

Growth factors in this study

FGF	Fibroblast Growth Factor	
EGF	Epidermal Growth Factor	
GCSF	Granulocyte Colony Stimulating Factor	
PDGFAA	Platelet Derived Growth Factor alpha	
PDGFBB	Platelet Derived Growth Factor beta	
PIGF	Placental Growth Factor	
TGFβ	Transforming Growth Factor	

Inflammation regulators

Interleukins 4,8,6 and 10

Inhibitors of matrix degrading enzymes, e.g. collagenases

TIMPs 1, 2 and 4





REGULATORY

- Regulated under Section 361 of the Public Health Service Act
- Passed March 2011 FDA audit and July 2012 FDA audit with clean inspection reports and no findings
- No 510(k) or PMA required for allograft tissue if:
 - Minimally manipulated
 - Homologous use
 - Not combined with another article
 - No systemic effect and No Living cells
- Human tissue already proven safe and effective
- MiMedx processing Certified by the American Association of Tissue Banks ("AATB")
- This regulatory path is a distinct advantage for speed to market



REIMBURSEMENT

- Completed two Random Controlled Trials (RCTs)
 - Diabetic Foot Ulcers Published in International Wound Journal
 - Plantar Fasciitis Published in Foot and Ankle International
- Conducting numerous additional RCTs
- Studies currently being conducted on:
 - Diabetic foot ulcers and venous leg ulcers
 - Spine scar tissue and cranial surgeries
 - Epicondylitis (tennis & golfer's elbow)
 - Radical prostatectomies
- Multiple health plans currently reimbursing
- Medicare Code Q4131 effective Jan 1, 2013 for EpiFix[®], chronic wound care
 - Six Regional MACs have committed to paying using this code: Palmetto, Cahaba, NGS, Novitas, NHIC and WPS.
 - Total Medicare Covered lives: 32.2 million (81%)



PLACENTA DONATION

Cesarean sections

- Placenta donor history
- Serologic tests
- Recovery Network
 - 19 Hospitals currently
 - 75+ Hospitals in contract negotiation process





PROPRIETARY PURION® PROCESS

- Gentle
- Dehydrated
- Effective bio-burden reduction
- 5 year shelf life



- Specifically developed for the unique characteristics of amniotic membrane
- Minimal graft manipulation maintains structural and biological integrity

Over 170,000 Amniotic Tissue Grafts Distributed



U.S. MARKET OPPORTUNITY Annual Cases

Total Wounds Difficult to heal wounds 6 Million² 2.5 Million²

Pain Management Back/Spine Injections Epicondylitis Plantar Fasciitis

>10 million⁸ >6 million³ >1 million⁹ >17 million

Surgical (Scar reduction and wound healing)

Prostatectomies Rotator Cuff / Labrum Spine & Cranial Surgery Knees / Hips >140k⁴ >1 Million⁵ >1 Million⁶ >1 Million⁷ >3 Million pr

>3 Million procedures



U.S. MARKET OPPORTUNITY Revenue Potential

\$13+ Billion addressable US market⁸

\$2.5 Billion Chronic & Acute wounds DFUs VLUs Burn Trauma

\$7.5+ Billion Tendinopathy & Pain Mgmt Tennis & Golfer's Elbow Plantar Fasciitis Back Injections

\$3 Billion Barrier & Minimize Scar FormationSpine SurgeryOrtho SurgerySports MedicineProstatectomies

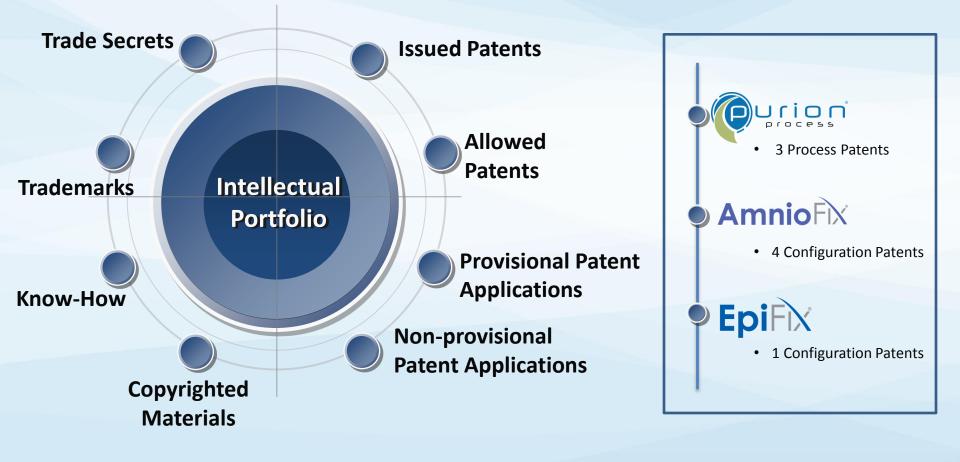
Key Market Drivers:

- Over 25 million people in US with Diabetes²
 - 1 million DFUs annually
- Chronic Wound treatment costs²
 >\$2.5 Billion annually
- Chronic wounds estimated to affect as many as 6 million patients annually²
 - DFUs, VLUs, Arterial,
 Pressure, Burns, Trauma
- Aging Population
 - Desire for active lifestyle
- Pain Management and emerging market



INTELLECTUAL PROPERTY

Patent coverage through 2028 - 2032





CLINICAL DATA

Case Study: Saving a Limb



- Wound size was 18.75 cm² Amputation already scheduled EpiFix[®] last resort
- Application of EpiFix[®] Graft with **30%** area reduction at **7 days**
- Additional 15% area reduction at Day 14
- Additional application of EpiFix[®] Graft and wound closed at day 28
- At 3 months wound remains fully closed and patient walking with custom molded shoe



CLINICAL DATA

Case Study: Healing with Reduced Scar Tissue









CLINICAL DATA Case Study: Thermal Burn

BOC



Treatment Day 1: EpiFix[®] Application



Day 6: Follow up

Day 9: Treatment & EpiFix[®] Application

CENTIMETERS



Day 16: Healed

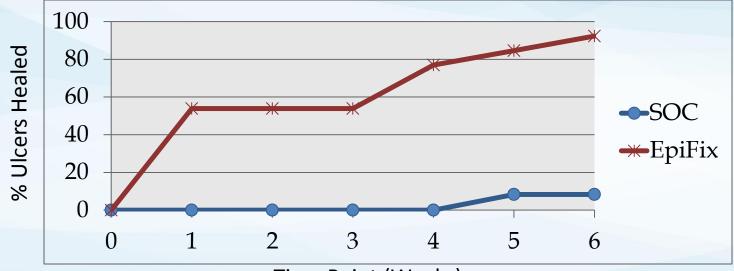
2 graft applications 16 days from initial EpiFix[®] Allograft application to healed wound



- Deep partial thickness burn to the left small finger
- Considered serious because of location
- Extremely painful

CLINICAL DATA

EpiFix® RCT on Diabetic Foot Ulcers



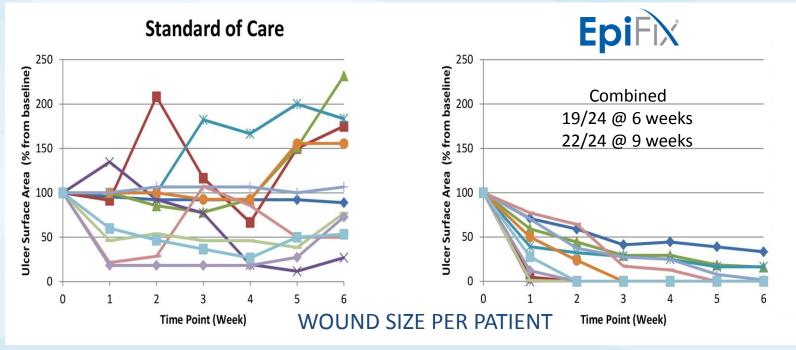
Time Point (Weeks)

Time point	Intervention Group		
	SOC (n=12)	EpiFix [®] (n=13)	ρ
4 Weeks	0 (0%)	10 (77%)	< 0.001
6 Weeks	1 (8%)	12 (92%)	<0.001



CLINICAL DATA

EpiFix® Cross-over Study on DFU's



- In the initial study period while receiving SOC only, wound sizes were inconsistent week to week.
- During the dHACM treatment period consistent reduction in wound size was noted week to week.



SALES FORCE

As of August 1



Government
AmnioFix28
28
Sales ProfessionalsCommercial Wound Care
EpiFixSales ProfessionalsEpiFix21
Sales ProfessionalsDistributors & Sales Agents
AmnioFix7
Sales Professionals



REVENUE GROWTH



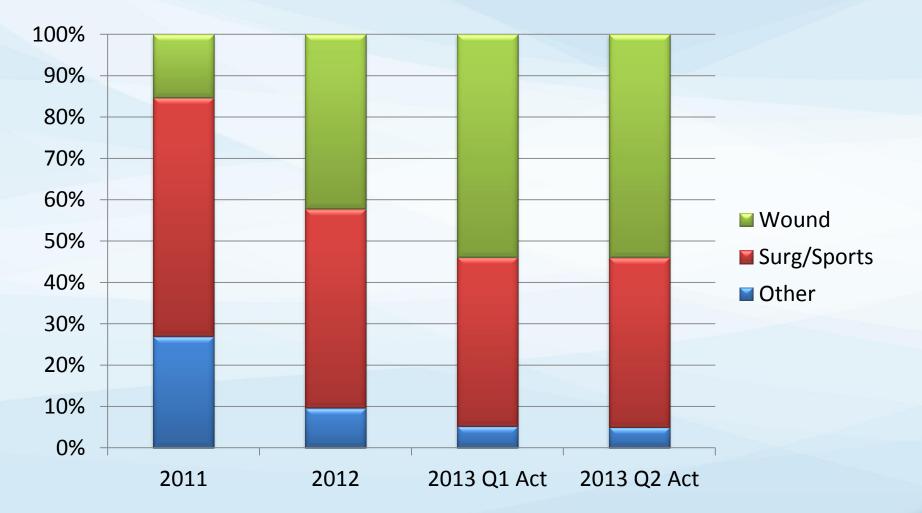


2013 REVENUE PROJECTION





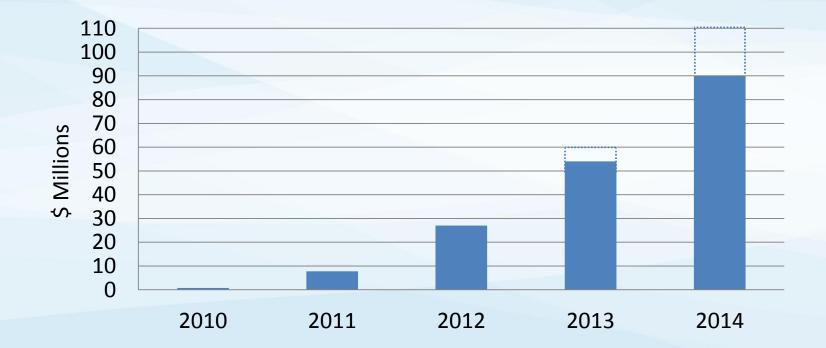
% REVENUE BY THERAPEUTIC AREA





REVENUE PROJECTIONS

329% Average Annual Growth Rate





OTHER KEY DATA

(\$ thousands)	As of June 30, 2013	As of December 31, 2012	As of December 31, 2011
EBITDA (ytd)	\$2,284	\$2,395	\$(6,314)
Current assets	21,528	18,089	6,882
Total Assets	40,283	35,183	27,096
Current Liabilities	6,141	5,071	4,732
Total Debt/ Earnout	0	9,805	12,579
Total Liabilities	7,391	15,175	15,200
Stockholders' Equity	32,892	20,007	11,897
Common Shares O/S	96,374	88,423	74,307
Employees	202	166	47



SUMMARY

- Technology platform well positioned to exploit large high growth markets, including wound care and soft tissue regeneration and stem cell like opportunities
- More predictable REGULATORY profile compared to traditional device companies due to allograft platform offerings
- EXPERIENCED management team with proven track record of success in high growth healthcare businesses
- Strong PATENT portfolio creates significant barriers to entry
- PURION[®] processed allografts demonstrate clinical and cost advantages over smaller competitors
- Near term milestones for value creation:
 - Completion of additional RCT's and scientific studies
 - Issuance of eight tissue related Patents with 33 filed
 - Expansion into additional surgical procedures with associated clinical studies
 - CMS Q code in Q1 2013 expands wound market opportunity
 - First Peer Reviewed Articles Published in Nationally Recognized Medical Journals



EpiFixattribute matrix

TREATMENT COST*	EpiFix®	Apligraf®	Dermagraft [®]
Average Applications per Patient	1-2	3-5	4-8
Average Cost to Closure	\$3,410	\$8,000	\$12,800
EASE OF USE	EpiFix®	Apligraf®	Dermagraft®
Vulnerable Viable Cells	NO	Yes	Yes
Wide Temperature Range	0-38° C	NO	NO
Frozen	NO	NO	Yes
Limited Temperature Range	NO	YES 20-23° C	YES (-80°C) ±10%
Shelf Life	5 Years	10 days	6 months
Shipping	Easy	Special (CO ₂)	Dry Ice
Multiple Sizes	YES	NO	NO

*1 EpiFix: Value Based Purchasing for Wound Care White Paper; Donald Fetterolf, MD, MBA; ² Falanga V; Margolis D; Alvarez O; Auletta M; Maggiacomo F; Altman M; Jensen J; Sabolinski M; Hardin-Young and the Human Skin Equivalent Investigators Group "Rapid healing of venous ulcers and lack of clinical rejection with an allogeneic cultured human skin equivalent." ArchDermatol. (134)3. 01-MAR-1998. pp 293-300. ³ Apligraf product promotional literature estimates of cost/reimbursements. 2010. ⁴ Apligraf product literature, "Rethink the Wound. Think Apligraf" 2010. John, T. "Human amniotic membrane transplantation: Past, present, and future." Ophthal Clin N Am. (16). 2003. pp 43-65. ⁵ Dermagraft product literature, describing treatments weekly for 8 applications.



REFERENCES

- Hansen, M.H., Anderson, J. J.: Chronic and Refractory Plantar Fasciitis Treatment with Single-Dose Dehydrated Amniotic Allograft Injection. Poster accepted and presented at the 2013 Annual Scientific Conference of the American Foot and Ankle Surgeons
- 2. BioMed GPS, LLC- Wound Biologics May 1, 2013
- 3. American Academy of Family Physicians, plus Management Estimates
- 4. Intellab, 2009
- 5. iData Research, Inc. 2012
- 6. iData Research, Inc. 2011
- 7. CDC Inpatient Data, 2011
- 8. Management Estimates
- 9. AMFAM Physician, 2011



##