



Investor Presentation

November 2024

helping humans heal.



Disclaimer & Cautionary Statements

This presentation includes forward-looking statements. Forward-looking statements are subject to risks and uncertainties, and the Company cautions investors against placing undue reliance on such statements. Actual results may differ materially from those set forth in the forward-looking statements. Such forward-looking statements include statements regarding:

- Future sales, sales growth, profitability and Adjusted EBITDA margins;
- Estimates of potential market size and demand for the Company's current and future products;
- Plans for expansion outside of the U.S.;
- The effectiveness of amniotic tissue as a therapy for any particular indication or condition;
- Expected spending on research and development, including to innovate and diversify our product portfolio;
- Investments in data;
- Expectations regarding the reimbursement environment for the Company's products, including Medicare Spending;
- Manner of LCD implementation;
- Expectations regarding plans to reduce customer churn and enhancing customer relationships;
- Expectations that HELIOGEN will be a meaningful contributor to our financial performance in 2025;
- The stage of development of the placental-derived products market;
- The Company's long-term strategy and goals for value creation, the status of its pipeline products, expectations for future products, and expectations for future growth and profitability





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Additional forward-looking statements may be identified by words such as "believe," "expect," "may," "plan," "potential," "will," "preliminary," and similar expressions, and are based on management's current beliefs and expectations. Forward-looking statements are subject to risks and uncertainties, and the Company cautions investors against placing undue reliance on such statements. Actual results may differ materially from those set forth in the forward-looking statements. Factors that could cause actual results to differ from expectations include:

- Future sales are uncertain and are affected by competition, access to customers, patient access to hospitals and healthcare providers, the reimbursement environment and many other factors;
- The future market for the Company's products can depend on regulatory approval of such products, which might not occur at all or when expected, and is based in part on assumptions regarding the number of patients who elect less acute and more acute treatment than the Company's products, market acceptance of the Company's products, and adequate reimbursement for such therapies;
- The process of obtaining regulatory clearances or approvals to market a biological product or medical device from the FDA or similar regulatory authorities outside of the U.S. is costly and time consuming, and such clearances or approvals may not be granted on a timely basis, or at all, and the ability to obtain the rights to market additional, suitable products depends on negotiations with third parties which may not be forthcoming; and
- The Company describes additional risks and uncertainties in the Risk Factors section of its most recent annual report and quarterly reports filed with the Securities and Exchange Commission. Any forward-looking statements speak only as of the date of this presentation and the Company assumes no obligation to update any forward-looking statement.



A Pioneer and Leader in Healing Solutions for Wound & Surgical

Our Why

Helping Humans Heal

Our Vision To be the leading global provider of healing solutions through relentless innovation to restore quality of life.



The most studied portfolio of placental-based products with **50+** clinical & scientific publications and over **300 million** payer covered lives.

Large, national placental donation network and **proprietary tissue processing**.

New product innovations leading to untapped opportunities for growth, including an increasing footprint in the Surgical market.

A key partner to healthcare professionals with industry leading support services and customer-focused approach.



>

The Unmet Need for Wound Healing Solutions Is Large and Growing





million people suffer from chronic, non-healing wounds in the U.S.¹

Favorable Demographic Trends



- Aging population
- Smoking history
- Diabetes

- Obesity
- Heart & vascular disease

Chronic Wounds
Burden Medicare
Beneficiaries



~16% of the Medicare beneficiary population is impacted by chronic wounds—and this proportion is increasing.¹

Ineffective Wound Management Leads to Poor Outcomes



It is estimated that up to 85% of amputations are avoidable with a holistic multispecialty team approach that incorporates innovative treatments and adherence to treatment parameters.²

Advances Driving Improved Outcomes for Patients



When applied following parameters for use, patients treated with EPIFIX® experienced reductions in major amputations and hospital utilization.²





Increasing Awareness of Massive Potential for Placental Tissue

The New York Times

Her Face Was Unrecognizable After an Explosion. A Placenta Restored It.

"Research has found placenta-derived grafts can reduce pain and inflammation, heal burns, prevent the formation of scartissue and adhesions around surgical sites and even restore vision. They're also gaining popularity as a treatment for the widespread issue of chronic wounds."



"...Tending to such wounds can be a matter of life and death for the millions of people with them, including 10.5 million Medicare beneficiaries as of 2022..."

"...The five-year mortality rate for people with one type, a diabetic foot ulcer, is close to 30 percent. That rate rises above 50 percent for those who require amputation."







The Patient Journey in Wound Care

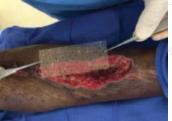
MIMEDX products are available in all settings where patients receive care...



...and are used on a range of chronic and other hard-to-heal wounds.

Acute Wounds





Mohs surgery

Burn/Trauma

Chronic Wounds





Diabetic Foot Ulcer

Venous Leg Ulcer





Limb Salvage



Dehiscence





Significant Opportunity to Drive Further Utilization in Surgical

Neuro

Mohs

Orthopedic



Cranioplasty Procedures with AMNIOFIX®

Clinical Outcomes with Conventional Methods¹



Average dissection time ~30 minutes



Frequent fibrosis reported



Fibrosis has increased complications



Clinical Outcomes with AMNIOFIX²



Average dissection time < 3 minutes

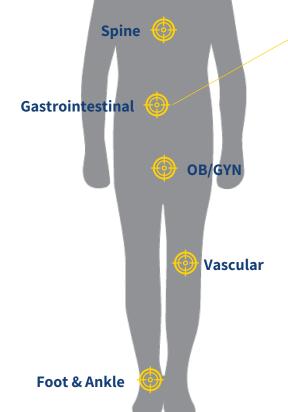


Minimal dural fibrosis in 86% of patients



No intraoperative complications

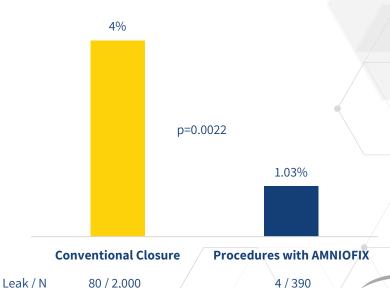






Colorectal Anastomoses Procedures with AMNIOFIX

Anastomotic Leak Rate with & without AMNIOFIX³





²⁾ Endicott L, Ehresman J, Tettelbach W, Forsyth A, Lee B. Dehydrated human amnion/chorion membrane (DHACM) use in emergent craniectomies shows minimal dural adhesions. J Wound

³⁾ F. Raymond Ortega, MD, FACS; Dennis Choat, MD, FACS, FASCRS; Emery Minnard, MD; Jeffrey Cohen, MD. The American College of Surgeons Clinical Congress, Oct 22-26, 2017, San Diego, CA.

Our Strategic Priorities

- Product Portfolio to Maximize Growth
- Continue momentum with new organic products in Wound & Surgical
- Consider additional inorganic additions to our product offering
- Drive further uptake of EPIFIX® in Japan

- Develop & Deploy
 Programs to Expand
 Footprint in Surgical
- Increase our presence in targeted surgical settings with our portfolio
- Invest in clinical data, partnering with KOLs

Enhance Customer Intimacy

Execute on initiatives to increase customer "stickiness" and reduce churn





Expanding Breadth of Skin Substitutes

Leading Human-Derived Portfolio



Best-in-Class Wound Product Portfolio



Innovative Offering for Surgical Market

Emerging Xenograft Platform



Recently announced exclusive manufacturing and supply agreement with Regenity Biosciences.

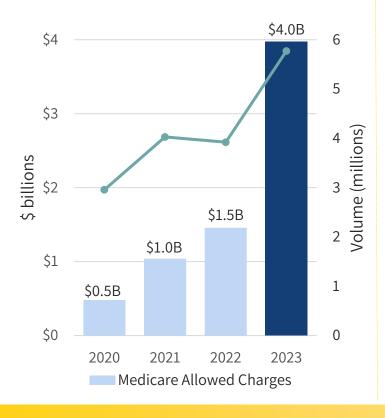
HELIOGEN builds on our goal to augment our growth through strategic portfolio expansion.

Provides MIMEDX with a bovine-derived collagen matrix particulate product that is 510(k)-cleared and indicated for the management of exudating wounds and to control minor bleeding.

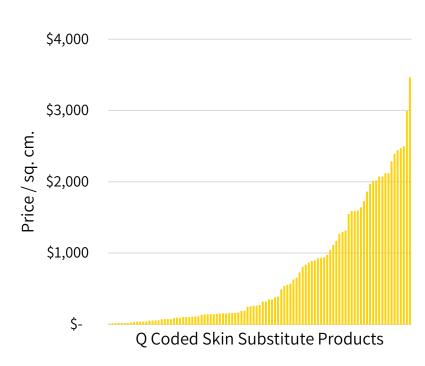


Worsening Medicare Spending Crisis Underscores Need for Overhaul

Medicare Allowed Charges¹ for skin substitutes have **exploded** since 2020...



...as have the number of skin substitutes on the Medicare ASP List², with a **significant number of products priced >\$1,000/sq.cm**...



...resulting in run rate spend of **over**\$1 billion of spend PER MONTH on products in the category

Average Price of Skin Substitutes on ASP List Has Increased More Than **Fivefold** Since Q1:22

LCDs³ proposed in April 2024 would disallow coverage of ~180 products in the category

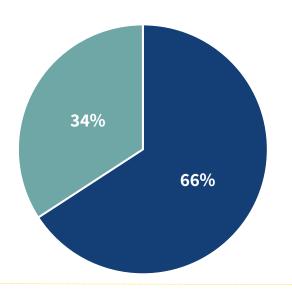
EPIFIX and EPICORD® are among the 15 covered products eligible for reimbursement in the proposal

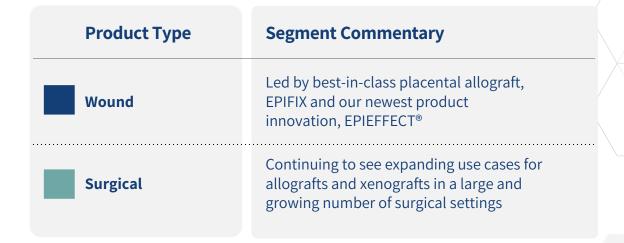
Proposed LCDs Represent Needed First Step to Curb Abuses in Private Office & Associated Care Settings

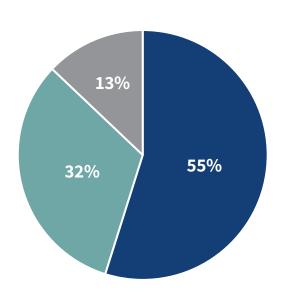


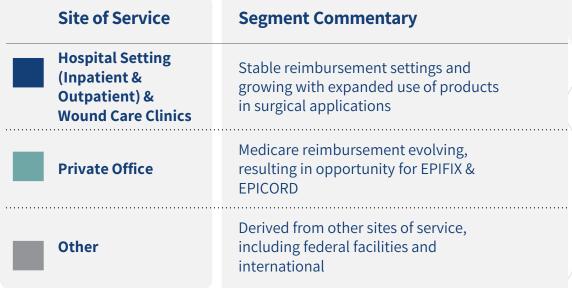


Diversified Business by Product & Across Multiple Sites of Service











Q3:24 Highlights

Net Sales \$84MM +3% year-over-year

Gross Margin 82%

GAAP Net Income \$8MM

\$18MM
22% of net sales

Cash Balance \$89MM +\$20MM vs. Q2:24



Lawmaker
Engagement to
Drive
Reimbursement
Reform

Patient Stories
Recently Featured
The New York Times





Management Team with Track Record of Success in MedTech



Joe Capper **Chief Executive** Officer



Doug Rice Chief Financial Officer



Kim Moller Chief Commercial Officer



John Harper, Ph.D. Chief Scientific Officer & SVP, R&D



Ricci Whitlow Chief Operating Officer



Butch Hulse Chief Administrative Officer & General Counsel



Kate Surdez Chief Human Resource Officer



Matt Notarianni Head of IR

Prior Roles Include:













BioTelemetry



Catalent.





































Pioneer and leader in Advanced Wound Care Expanding presence in Wound & Surgical

Committed to delivering above-market growth and profitability

helping humans heal.

