

Advancing Soft Tissue Reconstruction

Investor Presentation

November 2021

Forward Looking Statements

This presentation contains forward-looking statements within the meaning of The Private Securities Litigation Reform Act of 1995. All statements other than statements of historical facts contained in this document, including but not limited to statements regarding possible or assumed future results of operations, business strategies, development plans, regulatory activities, market opportunity competitive position, potential growth opportunities, and the effects of competition, are forward-looking statements. These statements involve known and unknown risks, uncertainties and other important factors that may cause TELA Bio, Inc.'s (the "Company") actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements. In some cases, you can identify forward-looking statements by terms such as "may," "will," "should," "expect," "plan," "aim," "anticipate," "could," "intend," "target," "project," "contemplate," "believe," "estimate," "predict," "potential" or "continue" or the negative of these terms or other similar expressions. The forward-looking statements in this presentation are only predictions. The Company has based these forward-looking statements largely on its current expectations and projections about future events and financial trends that it believes may affect the Company's business, financial condition and results of operations. These forward-looking statements speak only as of the date of this presentation and are subject to a number of risks, uncertainties and assumptions, some of which cannot be predicted or quantified and some of which are beyond the Company's control, including, among others: the impact to the Company's business of the ongoing COVID-19 pandemic and the development of new variants of COVID-19, such as the delta variant, including but not limited to any impact on the Company's ability to market its products, demand for the Company's products due to deferral of procedures using the Company's products or disruption in the Company's supply chain, the Company's ability to achieve or sustain profitability, the Company's ability to gain market acceptance for the Company's products and to accurately forecast and meet customer demand, the Company's ability to compete successfully, the Company's ability to enhance the Company's product offerings, development and manufacturing problems, capacity constraints or delays in production of the Company's products, maintenance of coverage and adequate reimbursement for procedures using the Company's products, product defects or failures. These and other risks and uncertainties are described more fully in the "Risk Factors" section and elsewhere in the Company's filings with the Securities and Exchange Commission and available at www.sec.gov. You should not rely on these forward-looking statements as predictions of future events. The events and circumstances reflected in the Company's forward-looking statements may not be achieved or occur, and actual results could differ materially from those projected in the forward-looking statements. Moreover, the Company operates in a dynamic industry and economy. New risk factors and uncertainties may emerge from time to time, and it is not possible for management to predict all risk factors and uncertainties that the Company may face. Except as required by applicable law, we do not plan to publicly update or revise any forward-looking statements contained herein, whether as a result of any new information, future events, changed circumstances or otherwise.



TELA BIO INC SURGICAL RECONSTRUCTION COMPANY

Redefining soft tissue reconstruction with a differentiated category of tissue reinforcement materials ~\$2B U.S Market Opportunity¹ in hernia repair, abdominal wall reconstruction and plastic and reconstructive surgery

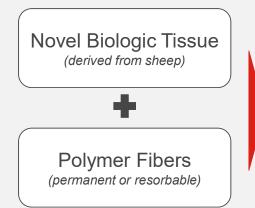
Innovative Products

- Compelling Clinical Evidence
- Products Offer Attractive Value Proposition for Hospitals

¹Management estimate. \$2B total equals \$1.5B hernia & abdominal wall reconstruction and \$0.5B plastic reconstructive surgery.

Creating Advanced Biologic Materials

Purposefully designed to address shortcomings & unmet clinical needs



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Innovative Textile Engineering

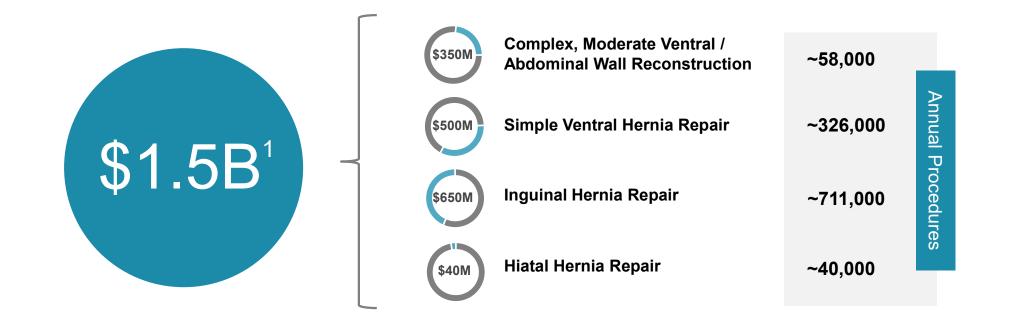


Hernia & Ab Wall Reconstruction	
Plastic Reconstruction	

Issued patents protect underlying biologic tissue and product design



OviTex: ~\$1.5 Billion Annual U.S. Hernia Market Opportunity



Source: Millennium Research Group Reports, IMS Health Data ¹Management estimate. Market size based volume weighted average selling price for OviTex



Ventral Hernia Patients Range in Complexity

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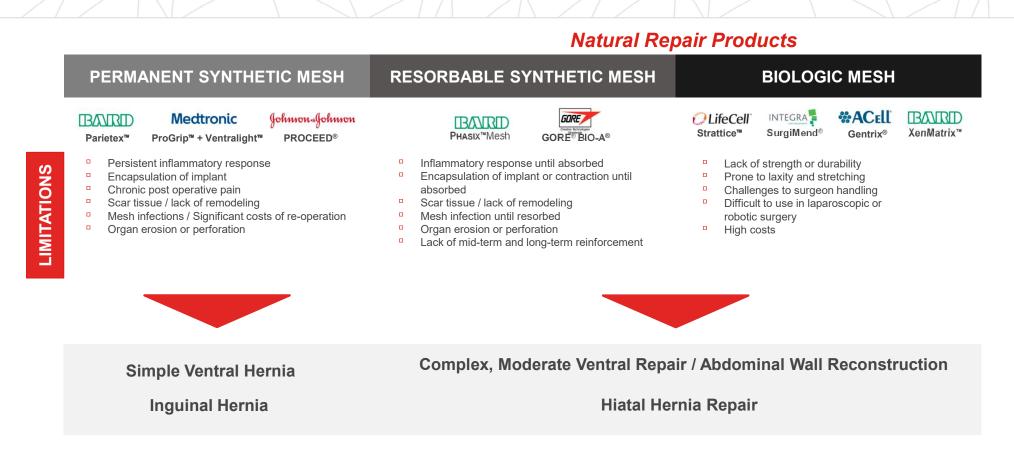
Ventral Hernia Complexity

SIMPLE	MODERATE	COMPLEX
 CDC Wound Class I (clean) Healthier patients - no co- morbidities Primary hernia repair 	 CDC Wound Class II (clean-contaminated) Patient co-morbidities (i.e., obesity, diabetes, COPD) May have prior hernia repair failure 	 CDC Wound Class III (contaminated) & IV (infected) Large defects Infected synthetic mesh removals Multiple prior hernia repair failures

Objective: provide patients the best repair the first time to prevent the simple patient from becoming the complex



Current Ventral Hernia Treatment Options: No Perfect Product



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Growing Need for Alternative to Permanent Synthetic Mesh



of surgeons agree that use of permanent synthetic mesh puts patients at long-term risk of complications¹



~15K

Hernia patients have voiced concern over use of permanent synthetic mesh in the past 12 months, according to surgeons¹

Lawsuits against permanent synthetic meshes estimated to be assembled across the U.S.²



HERNIA MESH COMPLICATIONS INCLUDE: PAIN, INFECTION RECURRENCE, ADHESION, OBSTRUCTION, & PERFORATION. THOSE AFFECTED MAY BE ELIGIBLE FOR COMPENSATION.



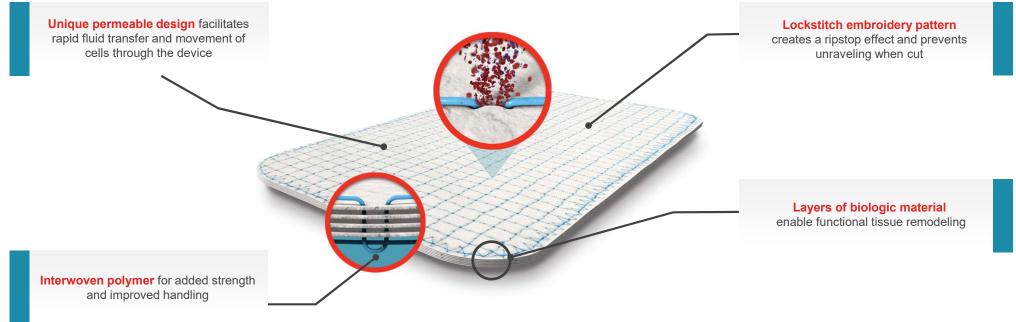
If you've been injured due to potentially defective hernia mesh, you may be entitled to financial compensation.



¹ Hernia and Abdominal Surgeries Survey (Oct 2020).
 ² www.drugwatch.com (October 2020)

OviTex Reinforced Tissue Matrix: a More Natural Hernia Repair™

An innovative reinforced tissue matrix designed to reduce stretch compared to biologic matrices and longterm complications experienced with resorbable and permanent synthetic meshes





OviTex LPR for Laparoscopic & Robotic Hernia Repair

Increase in Robotic-Assisted Hernia Repair

- Surgeons have adopted robotic-assisted techniques, primarily for inguinal & simple ventral Hernia repair, due to perceived patient and technique benefits
- Legacy biologic products are difficult to use minimally invasively (MIS) due to their thickness and handling properties



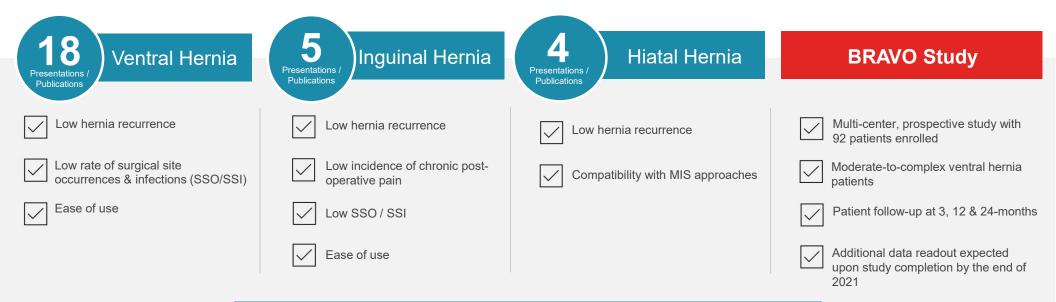
Our Solution: OviTex LPR

Tailored OviTex product designed for improved handling in MIS techniques and trocar accessibility





Compelling Clinical Evidence



OviTex supported by data from ~500 hernia patients across multiple hernia types



OviTex PRS: ~\$500 Million Annual U.S. Plastic & Reconstructive Surgery Market Opportunity

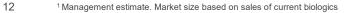


Surgeons use products to reinforce soft tissue during various reconstructive surgeries, including:

- Breast reconstruction
- Head and neck surgery
- Chest wall reconstruction
- Pelvic reconstruction
- Extremities reconstruction

Market dominated by human acellular dermal matrices (HADMs)

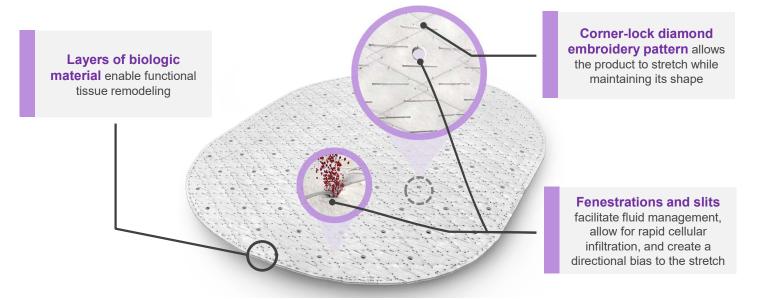
- Prone to high degree of stretch
- ^D Expensive, putting pressure on hospital systems
- ^D Often experience supply shortages, particularly when large pieces of material are required





OviTex PRS: Purposely Designed for Plastic and Reconstructive Surgery

An innovative reinforced tissue matrix designed to improve outcomes by facilitating fluid management and controlling degree and direction of stretch



Expanded commercial launch in June 2020 following limited launch initiated in 2019



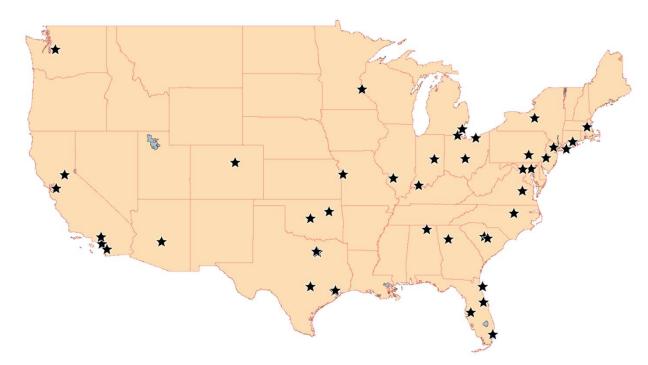
Commercial Organization

45 sales territories as of June 30, 2021

- OR-based Account Managers call on General, Plastic Recon, Colorectal & Trauma surgeons
- Carry full OviTex & OviTex PRS portfolios

6 sales regions

- Plan to scale existing regions until each region has ~8 territories
- Supported by Clinical Development and Strategic Customer Relations teams





Growth Strategy

INCREASE ADOPTION

- Promote broader awareness of OviTex & OviTex PRS products
- Employ virtual sales & marketing programs, including TELA LIVE
- Drive market awareness of risks of permanent synthetic mesh use
- Publish BRAVO clinical data

COMMERCIAL EXECUTION

- Scale direct sales force
- Drive account manager productivity
- Increase utilization within health systems under GPO contracts
- Secure additional contracts with high-potential IDNs and GPOs

MARKET EXPANSION

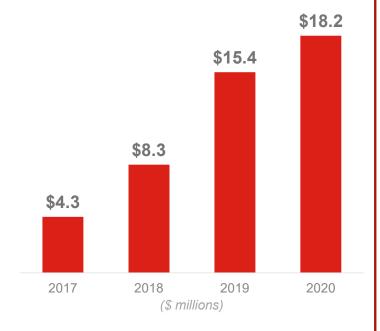
- Launch new product features and designs for OviTex and OviTex PRS
- Initiate robotic hernia post-market study
- Support investigator-led clinical studies for OviTex PRS



Delivering Revenue Growth

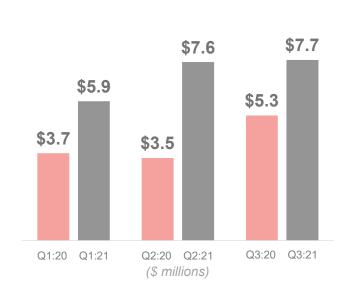
Annual Revenue

Revenue CAGR: 62%



Quarter Revenue

YTD Revenue Growth: 68%



Q3 2021 Performance

- Revenue growth of 44% year over year
- Cash and Cash equivalents (as of September 30, 2021): \$53.6M
- Robotic and MIS hernia procedures represent over 50% of all OviTex hernia procedures



Investment Highlights



Advanced reinforced tissue matrix portfolio supported by compelling clinical evidence



Focused on ~\$2.0 billion annual U.S. total addressable markets



Driving commercial adoption with targeted direct-sales approach

Recent product launches in growing markets: robotic hernia surgery + plastic and reconstructive surgery



Broad intellectual property portfolio



Established DRG-based reimbursement pathway for hernia repair



Industry leading executive team with proven track record

