Sanara MedTech Overview

December 2024



Disclaimers



This presentation contains forward-looking statements that discuss expectations as to future trends, plans, events, results of operations or financial condition, or state other information relating to Sanara MedTech Inc. (the "Company," "Sanara," "we," "our" or "us"). All statements other than statements of historical fact contained herein are forward-looking statements. These statements may be identified by terms such as "aims," "anticipates," "believes," "contemplates," "could," "estimates," "expect," "forecast," "guidance," "intend," "may," "plan," "possible," "potential," "predicts," "preliminary," "projects," "seeks," "should," "targets," "will," or "would," or the negatives of these terms, variations of these terms or other similar expressions. These forward-looking statements include, among others, statements concerning the development and launch of new products, the timing of commercialization of our products, the regulatory approval process and expansion of the Company's business in telehealth and wound care. These items involve risks, contingencies and uncertainties such as our ability to build out our executive team, our ability to identify and effectively utilize the net proceeds of the CRG term loan to support the Company's growth initiatives, the development and process for obtaining regulatory approval for new products, the extent of product demand, market and customer acceptance, the effect of economic conditions, competition, pricing, the ability to consummate and integrate acquisitions, and other risks, contingencies and uncertainties detailed in the Company's filings with the Securities and Exchange Commission ("SEC"), including the Company's most recently filed Annual Report on Form 10-K and the Company's Quarterly Reports on Form 10-Q as well as other documents the Company files with the SEC. Investors and security holders are urged to read these documents free of charge on the SEC's website at http://www.sec.gov. Forward-looking statement, whether as a result of new information, future eve

This presentation contains statistical and market data that we obtained from industry publications, reports generated by third parties, third-party studies and public filings. Although we believe that the publications, reports, studies and filings are reliable as of the date of this presentation, we have not independently verified such statistical or market data.

The trademarks and service marks included herein are the property of the owners thereof and are used for reference purposes only. Such use should not be construed as an endorsement of such products.

This presentation shall not constitute an offer to sell or the solicitation of an offer to buy securities, nor shall there be any sales of securities, in any state or jurisdiction in which such an offer, solicitation, or sale would be unlawful prior to registration or qualification under the securities laws of any such state or jurisdiction.

CAUTION: This presentation concerns certain products that are under clinical investigation and which have not yet been cleared for marketing by the U.S. Food and Drug Administration. These products are currently limited by federal law to investigational use, and no representation is made as to the safety or effectiveness of these products for the purposes for which they are being investigated.

Sanara MedTech at a Glance



A medical technology company focused on developing and commercializing transformative technologies to improve clinical outcomes and reduce healthcare expenditures in the surgical, chronic wound and skincare markets.

Ticker (Nasdaq)	SMTI
Market Cap	\$317.7M ⁽¹⁾
Avg. Daily Volume (TTM)	~20,240 ⁽²⁾
Business Segments	Sanara SurgicalTissue Health Plus
Net Revenue (TTM)	\$78.1M ⁽³⁾
Net Loss (TTM)	\$8.5M ⁽³⁾
Adj. EBITDA (TTM)	\$2.2M ^{(3) (4)}
Cash Balance	\$16.3M ⁽³⁾

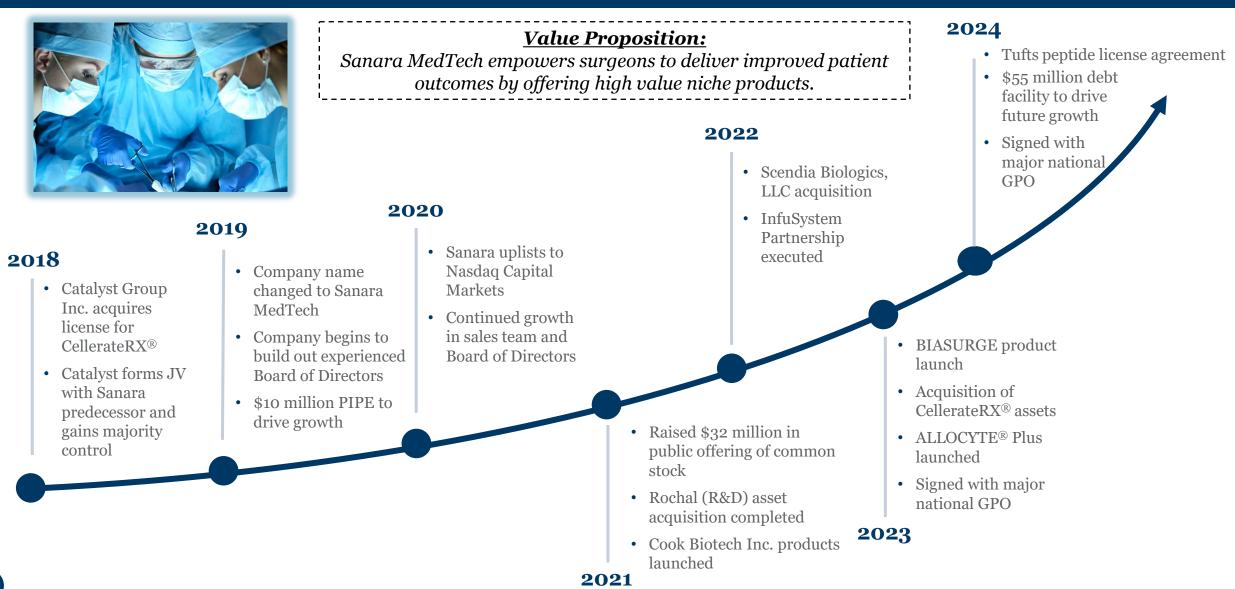
(1) Market cap. based on common shares outstanding of 8.7M as of 11/8/24 x share price of \$36.34 as of market close on 11/29/24

- (2) As of market close on 11/29/24
- (3) As of 9/30/2024

(4) Adjusted EBITDA is a non-GAAP financial measure. See the discussion and reconciliation in the appendix for additional information

The Sanara MedTech Story Continuous Innovation and Expansion





Board of Directors Strategic Leadership from Experienced Professionals





Ron Nixon

CEO & Executive Chairman

- Experience: Founder, The Catalyst Group, Inc.
- Education: B.S. in Mechanical Engineering from the University of Texas at Austin



Ann Beal Salamone, M.S.

Director

- Experience: Rochal Industries, EDC, National Academy of Engineering, TAMEST
- Education: B.S. from Western Kentucky University and M.S. from Rice University



Bob DeSutter

Director

- **Experience:** Piper Sandler
- Education: B.B.A. from the University of Minnesota and M.B.A from the University of Virginia Graduate School of Business



Sara N. Ortwein

Director

- Experience: ExxonMobil, XTO Energy, National Academy of Engineering, TAMEST
- Education: B.S. in Civil Engineering from the University of Texas at Austin



Roszell Mack III

Director

- Experience: Mack & Co., LLC, Ascend Venture Group, LLC, Goldman Sachs
- Education: B.A. in Engineering Sciences from Yale University and M.B.A. from Harvard Business School



Eric D. Tanzberger Director

- **Experience:** Service Corporation International, Coopers and Lybrand LLP
- Education: B.B.A. in Business Administration from the University of Notre Dame



Eric Major

Director

- Experience: HIGHRIDGE Medical, BarTrack, Inc., Stryker Spine, K2M, American OsteoMedix
- Education: B.S. from James Madison University



Keith Myers

Director

- Experience:
 - LHC Group, Co-Founder, Chairman & CEO
 - Optum, Senior Advisor
 - · Partnership for Quality Home Healthcare

🖸 CATALYST 🚔 Rochal PIPER SANDLER ExconMobil







Sanara Surgical Segment

















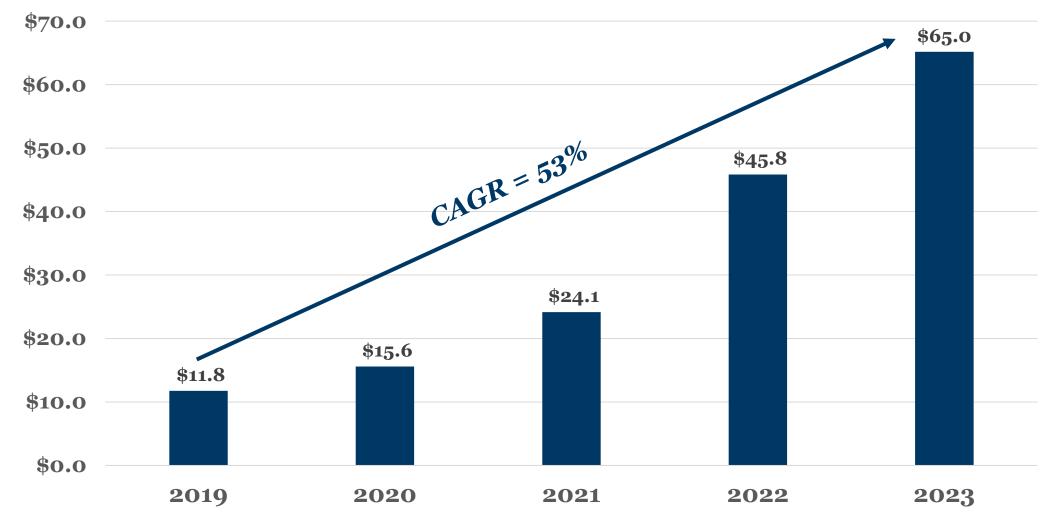




Sanara Surgical Segment: Demonstrated Track Record of Strong, Sustained Revenue Growth



Net revenues in millions \$



* For the periods presented, represents net revenues on a consolidated basis, as the Company managed its business on the basis of one operating and reportable segment prior to Q2 2024. All net revenues for the periods presented would have been attributable to the Surgical Segment had net revenues been presented on a segmented basis.

7



Empowering surgeons to deliver improved patient outcomes by offering high value niche products.

Sales Overview

- Products contracted or approved to be sold in >4,000 hospitals⁽¹⁾
- Products sold in >1,200 hospitals across 34 states and the District of Columbia⁽²⁾
- Net Revenue (TTM): $78.1M^{(1)}$, +25% year-over-year

Key Differentiators Driving Future Growth

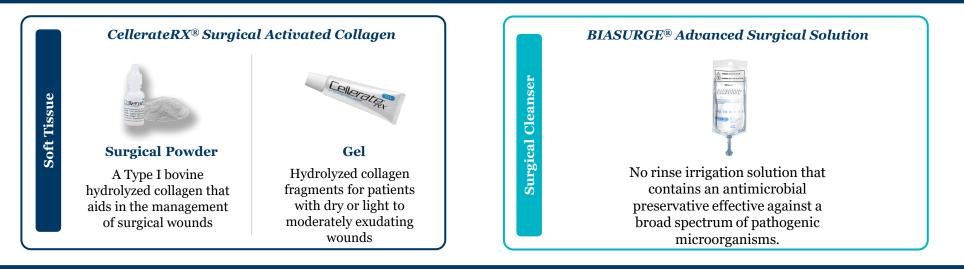
- Highly accomplished sales team
- Multiple selling opportunities per case
- Significant greenfield hospital opportunities
- In-house experienced R&D team driving new product development



Sanara Surgical Segment: Superior Product Offering with Significant Opportunity Runway



Platform Products



Ancillary Products

Fusion

Bone

FORTIFY TRG ® Tissue Repair Graft Multi-layered 100% small intestine submucosa (SIS) sheet indicated for implantation to reinforce soft tissue. Example: Applied by surgeons to reinforce Achilles tendon after rupture repair surgery FORTIFY FLOWABLE ® ECM Porcine small intestinal submucosa (SIS) ECM cryo-fractured to micronized ECM for cell migration and capillary growth Example: Applied by surgeons to aid in wound healing after amputation of the foot TEXAGEN ® Amniotic Membrane Allograft

Multilayer amniotic membrane that may be used as a soft tissue barrier and wound covering in numerous applications.

Example: Applied by surgeons as a wound covering after deep debridement

BiFORM® Bioactive Moldable Matrix

Bioactive and osteoconductive bone graft that can be hydrated and used as a strip or molded into a putty to fill a bone defect.

ACTIGEN[™] Verified Inductive Bone Matrix

Naturally derived, differentiated allograft matrix with robust handling properties.

ALLOCYTE[®] Plus Advanced Cellular Bone Matrix

Next generation cellular bone matrix (CBM) containing bonederived progenitor cells and conformable bone fibers.

Example: All three products above used for support of bone fusion (cervical spine fusion after trauma) depending on surgeon's preference

Soft Tissue

CellerateRX[®] Surgical Proven, Safe, and Effective

<u>Overview</u>

- CellerateRX[®] Surgical Powder is a medical hydrolysate of Type I bovine collagen and contains no additives
- Indicated for the management of acute and chronic wounds including:
 - Surgical wounds
 - Traumatic wounds
 - Partial and full-thickness wounds
 - First and second-degree burns
- Used to treat surgical site infections in multiple specialties primarily ortho and spine today





CellerateRX® Surgical Studies Supports Wound Management To Reduce Surgical Site Infection

Hydrolyzed Collagen Significantly Reduces Surgical Site Infections (Nowrouzi and Awad 2023)⁽¹⁾

Objective: Evaluate the effect of using type 1 hydrolyzed collagen in reducing incidence of surgical site infections associated with elective surgeries

Design: A retrospective analysis of 5,335 patients with similar patient profiles who underwent various surgeries (plastic, vascular, general, oncology, orthopedic, neuro, CVCT, and gynecology) were matched in a 3:1 (non-CellerateRX:CellerateRX) cohort and assessed for SSI rates.

59% reduction in Surgical Site Infections of clean contaminated cases with CellerateRX[®] Surgical Clinically meaningful difference in in SSI rate with CellerateRX[®] Surgical, 1.54% vs. 3.36%







Existing Acute Surgical Wound Growth

- The majority of CellerateRX[®] Surgical cases have been in the Ortho and Spine specialties.
- Team is working to further penetrate the >1,200 hospitals where our products are currently sold...
- ...while also expanding into the contracted hospitals that we are not selling in today.
 - Recently executed a contract with a national GPO giving SMTI sales team access to 1,000+ hospitals.
 - Products are contracted or approved to be sold in >4,000 hospitals⁽¹⁾

Expansion into Additional Acute Surgical Specialties

• Implementing a surgical strategy to further penetrate additional specialties including trauma, vascular, and general surgery.

Scale Recently Launched BIASURGE® Advanced Surgical Solution

• Focused on penetrating existing hospitals where CellerateRX® Surgical is currently sold.

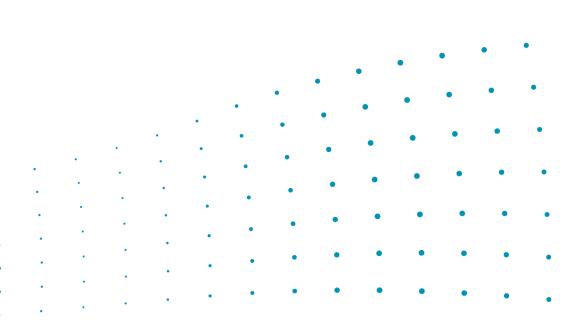
Inorganic Growth Opportunities

- Management anticipates synergistic potential transactions over time as a key growth driver that will complement strong organic growth.
- Pursuing multiple surgical M&A and partnership opportunities to complement its existing portfolio.



Tissue Health Plus Segment







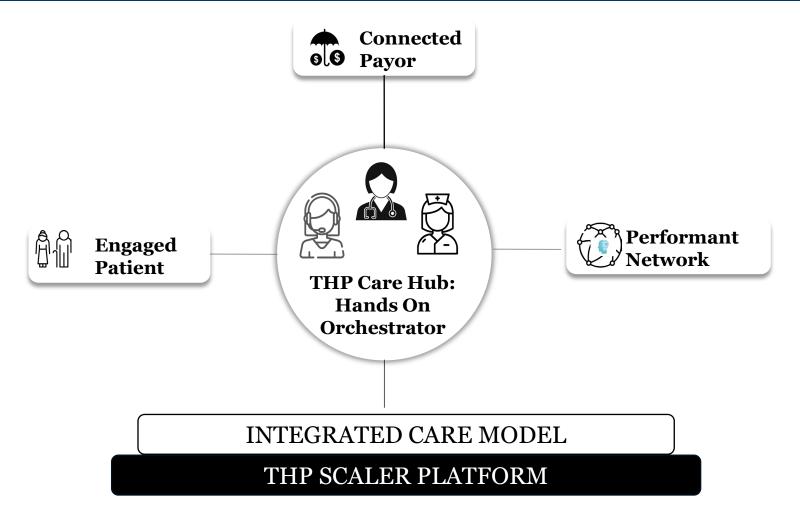
Significant Opportunity to Reduce Cost of Care in Non-Surgical Wound Care:

- Represents ~\$69B⁽¹⁾ of annual expenditure in U.S. hospitals, the majority of which is preventable
- Wound healing rates remain low
- 15% of individuals over 65 suffer from chronic non-healing wounds⁽¹⁾
- 5-year mortality rate for diabetic foot ulcers is over 30%⁽²⁾

Overview of Tissue Health Plus ("THP"):

- A first of its kind value-based wound care program targeting payors
- THP is designed to...
 - Coordinate delivery of wound care for patients under management
 - Integrate science and evidence-based medicine protocols to standardize wound prevention and treatment
- **Objective**: enable payors to divest wound care spend risk, reduce wound related hospitalizations and improve patient quality of life
- (1) THP Analysis + Carter MJ, DaVanzo J, Haught R, et al. Chronic wound prevalence and the associated cost of treatment in Medicare beneficiaries: Changes between 2014 and 2019. J Med Econ 2023;26(1): 894–901; doi: 10.1080/13696998.2023.2232256
- (2) Sheets, AR, Hwang, C and IM Herman 2016. Developing Smart Point-of-Care Diagnostic Tools for "Next-Generation" Wound Care In Translating Regenerative Medicine to the Clinic. http://dx.doi.org/10.1016/B978-0-12-800548-4.00017-6. Copyright © 2016 Elsevier Inc. IM Herman. 2016 Translating Regenerative Medicine to the Clinic Placeholder for footnote with source

Planned Tissue Health Plus ("THP") Model



Sanara MedTech

Commercial launch expected in mid-2025

Q3 2024 Financial Highlights



Sanara MedTech Evidence Based Healing

Q3 2024 Financial Highlights (Unaudited)



Total Company

- Net revenue of \$21.7M (+35% year-over-year)
 - Twelfth consecutive record net revenue quarter
 - Highest net revenue quarter in the Company's history
- Net loss of \$2.9M in Q3 and \$8.2M YTD
- Adjusted EBITDA⁽¹⁾ of \$0.8M in Q3 and \$1.7M YTD
- Cash balance: $16.3M^{(2)}$
- Additional borrowing capacity: $24.5M^{(2)}$

- Sanara Surgical Segment
 - Segment net loss of \$0.2M in Q3 and \$2.9M YTD
 - Segment EBITDA⁽¹⁾ of \$2.6M in Q3 and \$5.1M YTD
- Tissue Health Plus Segment
 - Segment net loss of \$2.7M in Q3 and \$5.3M YTD
 - Segment EBITDA⁽¹⁾ of (\$1.7M) in Q3 and (\$3.4M) YTD
 - Continued investment to build this strategy through the expected commercial launch in mid-2025
 - Continued investment through mid-2025 estimated at approximately \$5 to \$10 million
 - Pursuing like-minded partners to invest in the execution of this strategy

Thank You



• • • • • •

•

· · ·

Appendix



• • • •

•

• • • •



• Revenue

- For the three months ended September 30, 2024, Sanara generated net revenue of \$21.7 million compared to net revenue of \$16.0 million for the three months ended September 30, 2023, a 35% increase from the prior year period.
- The higher net revenue for the three months ended September 30, 2024 was primarily due to increased sales of soft tissue repair products, including CellerateRX[®] Surgical, as a result of our increased market penetration, geographic expansion, and our continuing strategy to expand our independent distribution network in both new and existing U.S. markets.
- SG&A
 - SG&A expenses for the three months ended September 30, 2024 were \$19.0 million compared to SG&A expenses of \$13.9 million for the three months ended September 30, 2023.
 - The higher SG&A expenses for the three months ended September 30, 2024 were primarily due to the buildout of our THP platform and infrastructure which increased by approximately \$1.2 million compared to the prior year period. In addition, higher direct sales and marketing expenses increased by approximately \$3.7 million compared to the prior year period due largely to higher sales commissions as a result of higher revenues.

• R&D Expenses

- R&D expenses for the three months ended September 30, 2024 were \$1.4 million compared to R&D expenses of \$1.0 million for the three months ended September 30, 2023.
- R&D expenses included \$0.6 million and \$0.8 million attributable to our THP segment for the three months ended September 30, 2024 and 2023, respectively.
- The higher R&D expenses for the three months ended September 30, 2024 compared to the three months ended September 30, 2023 were primarily due to new projects associated with CellerateRX[®] Surgical.

Q3 2024 Financial Summary (Continued)



• Depreciation, Amortization and Interest Expense

- Depreciation and amortization expenses for the three months ended September 30, 2024 were \$1.1 million compared to \$1.0 million for the same period in 2023. The higher depreciation and amortization expenses in 2024 were due to amortization of intangible assets acquired from Applied Nutritionals in August 2023.
- Interest expense was \$0.9 million for the three months ended September 30, 2024 compared to \$0.2 million during Q3 2023. The higher interest expense in 2024 was related to our new term loan with CRG.
- Net Loss
 - Sanara had a net loss of \$2.9 million for the three months ended September 30, 2024, compared to a net loss of \$1.1 million for the three months ended September 30, 2023.
 - The net loss included \$2.7 million and \$1.7 million attributable to our THP segment for the three months ended September 30, 2024 and 2023, respectively.
 - The higher net loss for the three months ended September 30, 2024 was primarily due to higher SG&A costs related to the buildout of our THP platform and infrastructure, higher interest expense related to our new term loan with CRG, an increase in expense due to change in fair value of earnout liabilities and higher amortization of our acquired intangible assets as discussed above, partially offset by higher gross profit.
- Cash Balances at End of Quarter (in millions)
 - \$6.2 (Q3-23), \$5.1 (Q4-23), \$2.8 (Q1-24), \$6.2 (Q2-24), \$16.3 (Q3-24)



- Segment Reporting Overview
 - Break out of Tissue Health Plus and Sanara Surgical demonstrates our strategic rationale of the acute/post acute comprehensive strategy investments.

Three Months Ended September 30, 2024

	<u>Surgical</u>	THP	<u>Total</u>
Net Revenue	\$21.7	-	\$21.7
Net loss	\$(0.2)	\$(2.7)	\$(2.9)
Segment EBITDA / Adjusted EBITDA (consolidated) ⁽¹⁾	\$2.6	\$(1.7)	\$0.8

Three Months Ended September 30, 2023

	<u>Surgical</u>	<u>THP</u>	<u>Total</u>
Net Revenue	\$16.0	-	\$16.0
Net income (loss)	\$0.6	\$(1.7)	\$(1.1)
Segment EBITDA / Adjusted EBITDA (consolidated) ⁽¹⁾	\$1.4	\$(1.1)	\$0.3

Nine Months Ended September 30, 2024

	<u>Surgical</u>	<u>THP</u>	<u>Total</u>
Net Revenue	\$60.4	-	\$60.4
Net loss	\$(2.9)	\$(5.3)	\$(8.2)
Segment EBITDA / Adjusted EBITDA (consolidated) ⁽¹⁾	\$5.1	\$(3.4)	\$1.7

Nine Months Ended September 30, 2023

	<u>Surgical</u>	<u>THP</u>	<u>Total</u>
Net Revenue	\$47.3	-	\$47.3
Net income (loss)	\$1.2	\$(5.4)	\$(4.2)
Segment EBITDA / Adjusted EBITDA (consolidated) ⁽¹⁾	\$3.8	\$(4.1)	\$(0.3)



Use of Non-GAAP Financial Measures

To supplement the Company's financial information presented in accordance with generally accepted accounting principles in the United States ("GAAP"), we present certain non-GAAP financial measures in this presentation, including Adjusted EBITDA and Segment EBITDA. The Company's management uses these non-GAAP financial measures, both internally and externally, to assess and communicate the financial performance of the Company. The Company defines Adjusted EBITDA as net income (loss) excluding interest expense/income, provision/benefit for income taxes, depreciation and amortization, non-cash share-based compensation expense, change in fair value of earnout liabilities, share of losses from equity method investment, executive separation costs, legal and diligence expenses related to acquisitions and gains/losses from the disposal of property and equipment, as each is applicable to the periods presented. The Company believes Adjusted EBITDA and Segment EBITDA are useful to investors because they facilitate comparisons of its core business operations across periods on a consistent basis. Accordingly, the Company adjusts for items such as change in fair value of earnout liabilities when calculating Adjusted EBITDA and Segment EBITDA because the Company believes that they are not related to the Company's core business operations. Segment EBITDA is calculated in the same manner as Adjusted EBITDA but is presented on a segment basis.

The Company's non-GAAP financial measures are not in accordance with, nor an alternative for, measures conforming to GAAP and may be different from non-GAAP financial measures used by other companies. In addition, these non-GAAP financial measures are not based on any comprehensive set of accounting rules or principles. The Company continues to provide all information required by GAAP, but it believes that evaluating its ongoing operating results may not be as useful if an investor or other user is limited to reviewing only GAAP financial measures. The Company does not, nor does it suggest that investors should, consider these non-GAAP financial measures in isolation from, or as a substitute for, financial information prepared in accordance with GAAP. Material limitations associated with the use of such measures include that they do not reflect all costs included in operating expenses and may not be comparable with similarly named financial measures of other companies. Furthermore, these non-GAAP financial measures are based on subjective determinations of management regarding the nature and classification of events and circumstances. The Company presents these non-GAAP financial measures to provide investors with information to evaluate the Company's operating results in a manner similar to how management evaluates business performance. To compensate for any limitations in such non-GAAP financial measures, management believes that it is useful in understanding and analyzing the results of the business to review both GAAP information and the related non-GAAP financial measures. Whenever the Company uses a non-GAAP financial measure, it provides a reconciliation of the non-GAAP financial measure to the most directly comparable GAAP financial measure. Investors are encouraged to review and consider these reconciliations.

Segment EBITDA is reported to the chief operating decision maker for purposes of making decisions about allocating resources to the segments and assessing their performance.

Reconciliation of GAAP to Non-GAAP Financial Measures (Consolidated)



	Three Months Ended September 30,						
	-	2024				2023	
	Sanar	a			Sanara		
	Surgic	al <u>THP</u>	<u> </u>	tal	Surgical	THP	Total
Net Income (Loss)	\$ (180,4	88) \$(2,702,	564) \$(2,88	33,052)	\$ 567,235	5 \$(1,662,184)	\$(1,094,949)
Adjustments:							
Interest expense	927,5			27,577	188,294		188,294
Depreciation and amortization Noncash share-based	n 696,8	406,	966 1,10)3,854	590,563	3 407,111	997,674
compensation	1,003,5	99 21,	831 1,02	25,430	813,600	6 43,920	857,526
Change in fair value of earno liabilities	ut	- 147,	000 1/	47,000	(758,783	3) 77,030	(681,753)
Share of losses from equity		- 14/,	000 1-	17,000	(750,70.) 11,050	(001,755)
method investment	31,4	48	_ 3	31,448			-
Executive separation costs	59,6			59,685			-
Acquisition costs	24,8			30,019			-
Segment EBITDA (on a segme					•		
basis) / Adjusted EBITDA							
(consolidated)	\$2,563,5	21 \$(1,721,	560) \$ 84	41,961	\$1,400,910	5 \$(1,134,124)	\$ 266,792
			Nine N	Month	s Ended		
				otembe			
		2024	•		,	2023	
	Sanara			<u> </u>	Sanara		
	Surgical	THP	Total	Sı	urgical	ТНР	Total
Net Income (Loss)	\$(2,872,286)	\$(5,339,011)	\$(8,211,29	97) \$ 1	,181,296 \$	(5,358,754) \$	(4,177,458)
Adjustments:							
Interest expense Depreciation and	1,839,259	-	1,839,25	59	188,300	-	188,300
amortization	2,093,797	1,220,984	3,314,78	81 1	,359,180	1,221,063	2,580,243
Noncash share-based compensation	2,803,536	108,031	2,911,56	57 2	,423,335	158,828	2,582,163
Change in fair value of earnout liabilities	(14,451)	82,000	67,54	49 (1	,385,914)	(108,996)	(1,494,910)
Share of losses from equity							
method investment	31,448	-	31,44	48	-	-	-
Executive separation costs (1)	964,466	-	964,46	56	-	-	-
Acquisition costs	249,901	577,892	827,79	93			-
Segment EBITDA (on a							

segment basis) / Adjusted EBITDA (consolidated)

§ 5,095,670 **§**(3,350,104) **§** 1,745,566 **§** 3,766,196 **§**(4,087,858) **§** (321,662)

(1)- Includes \$328,795 of share-based compensation related to executive separation costs for the nine months ended September 30, 2024.