

Important Information Regarding Forward-Looking Statements

This presentation includes 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. These statements, among other things, relate to the Company's growth drivers and expected levels of our organic growth; the impact of our investment in development and commercial initiatives; financial guidance, including timing of revenues and EBITDA; our ability to manage costs and to achieve our financial goals; our ability to operate under lending covenants; our ability to maintain sufficient liquidity to operate the business; our ability to pay our debt under our credit agreement and to maintain relationships with CDMO commercial partners and develop additional commercial and development partnerships. The words "anticipate", "believe", "could", "goal, "objective", "estimate", "upcoming", "expect", "intend", "may", "might", "plan", "predict", "project", "will". "should", "can have", likely and similar terms and phrases may be used to identify forward-looking statements in this presentation. The forward-looking statements in this presentation are only predictions.

Our operations involve risks and uncertainties, many of which are outside our control, and any one of which, or a combination of which, could materially affect our results of operations and whether the forward-looking statements ultimately prove to be correct. Factors that could cause the company's actual outcomes to differ materially from those expressed in or underlying these forward-looking statements include, but are not limited to, unstable market and macroeconomic conditions, including any adverse impact on the customer ordering patterns or inventory rebalancing or disruption in raw materials or supply chain; demand for the company's services, which depends in part on customers' research and development funding, their clinical plans and the market success of their products; customers' changing inventory requirements and manufacturing plans; customers and prospective customers decisions to move forward with the company's manufacturing services; the average profitability, or mix, of the products the company manufactures; the company's ability to enhance existing or introduce new services in a timely manner; fluctuations in the costs, availability, and suitability of the components of the products the company manufactures, including active pharmaceutical ingredients, excipients, purchased components and raw materials, or the company's customers facing increasing or new competition; the Company's ability to successfully enact its business strategies, including with respect to installation, capacity generation and its ability to attract demand for its services; the Company's ability to remain current with its reports with the Securities and Exchange Commission (the "SEC"); the Company's ability to collect on customers' receivable balances; the extent to which health epidemics and other outbreaks of communicable diseases could disrupt our operations; and other risks and uncertainties discussed in our filings with the Securities and Exchange Commission at www.sec.gov. These forward-looking statements are based o

Any historical or projected financial information contained in this presentation are not intended to be indicative of future financial results. The events and circumstances reflected in these forward-looking statements, may not be achieved or occur, and actual results could differ materially from those projected in the forward-looking statements. Undue reliance should not be placed on the forward-looking statements. Moreover, we operate in a dynamic industry and economy. New risk factors could emerge from time to time, and it is not possible for our management to predict all uncertainties that the Company may face.





Non-GAAP Financial Measures

This presentation contains non-GAAP financial information including Adjusted EBITDA. The Company has included a reconciliation of Adjusted EBITDA to Net (loss) income, the most directly comparable financial measure calculated in accordance with GAAP. We define Adjusted EBITDA Net (loss) income as determined under GAAP excluding (i) interest expense, net of interest income, (ii) provision for income tax expense (benefit), (iii) depreciation and amortization, (iv) stock-based compensation, (v) change in fair value derivatives, (vi) financing fees (non-interest), (vii) reorganization costs, (viii) restructuring costs, (ix) franchise tax equivalent to income tax, (x) contract cancellations, (xi) stockholder activist settlement costs, and (xii) start-up costs.

The Company has disclosed these non-GAAP financial measures to supplement its consolidated financial statements presented in accordance with GAAP. These non-GAAP financial measures exclude/include certain items that are included in the Company's results reported in accordance with GAAP. Management believes these non-GAAP financial measures provide useful additional information to investors about trends in the Company's operations and are useful for period-over-period comparisons. These non-GAAP financial measures should not be considered in isolation or as a substitute for the comparable GAAP measures. In addition, these non-GAAP financial measures may not be the same as similar measures provided by other companies due to the potential differences in methods of calculation and items being excluded/included. These non-GAAP financial measures should be read in conjunction with the Company's consolidated financial statements presented in accordance with GAAP.





Investor Day Key Takeaways

CDMO industry leader in complex injectables with broad capabilities that support large and growing markets

Actionable strategy to drive strong growth of a 12%+ revenue CAGR and adj. EBITDA margins of 25%+ in mid-term

A proven and experienced team dedicated to execution and creating value for customers, employees, and shareholders

Current projected capacity to support ~ \$300 million in annual revenue





About Lifecore - Our Journey

Market Overview

Growing Our Business

Sustaining Objectives

Question & Answer





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Market Overview

Growing Our Business

Sustaining Objectives

Question & Answer



Our Journey: Transformation to Standalone CDMO













Our Journey: Transformation to Standalone CDMO

NOW:



Best-in-class technical capabilities

Strengthened financial position

Doubled revenue-generating capacity

Enhanced business development resources & strategy

Nasdaq / regulatory compliance

Leadership transition complete





Experienced Management Team with Proven Ability to Execute

Paul Josephs
President &
Chief Executive Officer



Joined: 2024 30+ years experience

- President & Chief Executive Officer at Woodstock Sterile Solutions
- Head of CDMO-Global Business Development at Viatris (formerly Mylan)

Ryan Lake Chief Financial Officer



Joined: 2024 24+ years experience

- Extensive senior financial and strategic life sciences leadership experience
- Chief Financial Officer of Societal CDMO, Recro Pharma, Baudax Bio, Aspire Bariatrics, DSM Biomedical, Kensey Nash

Thomas Guldager
VP of Operations



Joined: 2024 20+ years experience

 Senior executive, manufacturing and site leader at Xellia Pharmaceuticals

Jackie Klecker

EVP Quality and Development Services



Joined: 2001 30+ years experience

 Served in various roles at Lifecore surrounding Quality Assurance and Regulatory Affairs

Darren Hieber

SVP of Corporate Development & Partnerships



Joined: 2021 20+ years experience

 VP of Business Development, Drug Product at Catalent

Brikkelle Thompson

SVP of Human Resources



Joined: 2024 24+ years experience

- Head of Human Resources
 the Americas at Teleflex
- VP of Human Resources at Nonin Medical





Lifecore at a Glance

Fully integrated CDMO offering development and fill/finish of sterile injectable pharmaceuticals

450 Employees

Driven Culture

Inclusive, Performance-

Corporate
Headquarters

Milmar Minneapolis

Chaska O hester O Austin O Chaska O Chas

Projected Revenues* (FY2025E)

\$126.5M - \$130M

Projected Adj. EBITDA* (FY2025E)

\$19M - \$21M

- Founded in 1965
- Leader in Sodium Hyaluronate (HA)
- Global Regulatory Capabilities







Campus Overview

248,000 Sqft State-of-the-art facilities, within 2 square miles

Employees

Site 1 - HQ (Lyman Blvd.)

150,000 sqft



Site 2 (Lakeview Drive) 78,000 sqft



Site 3 (Shelby Court) 20,000 sqft



Manufacturing Operations

- Sodium hyaluronate manufacturing (fermentation)
- Drug and medical device formulation and filling
- · Secondary packaging
- · Microbiology and analytical quality control laboratories
- Warehousing: 6,400 sqft CRT; 1,500 sqft cooler
- Distribution

Contract Development

Pilot laboratory

Manufacturing Operations

- Final packaging
- Warehousing: 16,400 sqft CRT; 4,000 sqft cooler
- Distribution
- Quality control laboratory
- Particulate lab

Contract Development

Analytical development laboratory

Manufacturing Operations

- · Receipt, inspection, & warehousing of raw materials and components
- 10,000 ft² CRT; 1,795 sqft cooler







Our Commitment







Shareholders

Deliver a strong financial track record and generate sustainable value over time

Customers

Be a value-added partner that provides the "hands" and the "brain" and delivers highquality solutions to our customers development and manufacturing needs

Employees

Be a fantastic, inclusive, performance-driven organization that lives its core values daily







About Lifecore - Our Journey

Market Overview

Growing Our Business

Sustaining Objectives

Question & Answer



We Serve Large and Growing Markets with Strong Tailwinds

Global CDMO

\$120B

Market1

+8% CAGR

Hyaluronic Acid

\$9.8B

Market²

+7% CAGR

Global Injectable CDMO

\$10B

Market¹

+10% CAGR

GLP-1

\$47B

Market³

Expected to Increase 10X

Biosecure Act 50%+

of annual US drug approvals are injectables⁴



^{1.} Jefferies September 2024 PBOA - 8th Annual Meeting Uncovering Life Sciences Investment Trends / J. Miller October 2024 - Outsourcing Includes drug product (finished dose form), drug substance (active pharmaceutical ingredients (API)



^{2.} Global Market Insights March 2024 – Hyaluronic Acid Market Size & Share – Trends Reports, 2024-2032

^{3.} Markets and Markets July 2024- GLP-1 Analogues Market Size, Share & Trends 2032

^{4.} William Blair Equity Research August 2024 - Percent of FDA Approvals for 2023 and YTD as of July 31, 2024



About Lifecore - Our Journey

Market Overview

Growing Our Business

Sustaining Objectives

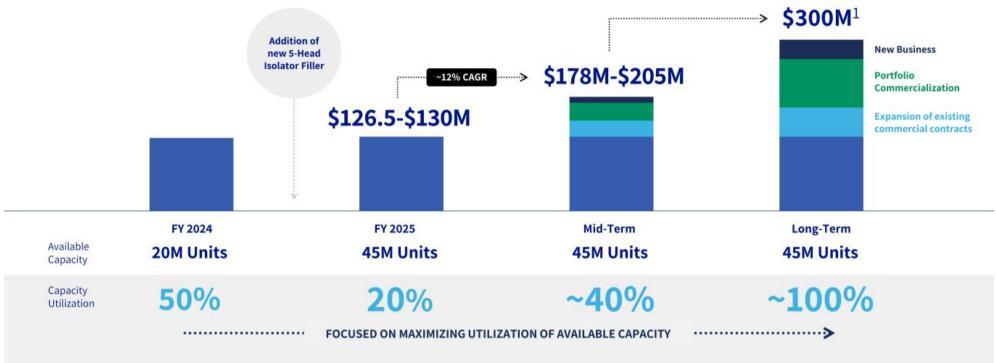
Question & Answer





Mid-Term and Long-Term Revenue Outlook

Revenue growth driven by maximization of existing customer base, portfolio commercialization, and new business





For illustrative purposes only, timing, estimates, assumptions and the actual growth of revenue and capacity utilization may vary significantly, and we may not be able to achieve our anticipated financial goals. The information provided is illustrative only; the growth cycle may not be achieved and there is continued uncertainty relating to any guidance contained herein. There can be no assurance that such results will occur or that such results may be materially different from actual results.

1. Based on estimates derived from internal testing and historical capacity data. There can be no assurance that such results will occur or that such results will be materially different from actual results.



Expanding Existing Customer Relationships

Know our customers

Establish trust and reliability

Establish Lifecore as a partner-ofchoice for the future CDMO needs of existing customers

Anticipate customers' growing needs Efficient onboarding of new programs Consistent engagement

Focus on commercial excellence

Maintain/increase margin profile

Lifecore prides itself on building long-term relationships, with multiple customer relationships ranging from

20 yrs to nearly 40 yrs¹





Fill & Finish: Pathway to Doubling Commercial Demand





- Significant inflection point expected from minimum volumes beginning in 2027
- Potential upside to contractual minimums



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HA Fermentation: Strong & Steady Demand

Lifecore manufactures >20 commercially approved HA injectable products

LIFECORE'S PREMIUM SODIUM **HYALURONATE:**

More than 150 million¹

doses sold worldwide

Proven Applications Worldwide:

- Ophthalmology
- Orthopedics
- Drug delivery
- Biomaterials

- Aesthetics
- Oncology
- Pain management





Strong, Diverse Pipeline

Total Pipeline Represents

\$100M - \$200M1

in Incremental Commercial Revenue Potential



- Impactful commercial revenue potential over the mid-term
- Strong development project pipeline: vials, syringes, cartridges
- Diversification across broad customer base



^{1.} Assumes full realization of management's estimates for annual commercial revenue potential from pipeline projects at peak sales. Information presented is not risk and probability adjusted and the actual revenue realization may vary significantly. This does not assume new customer additions or attrition. There can be no assurance that such results will occur or that such results will be materially different from actual results.

Projects are defined as individual drugs or devices for which Lifecore provides manufacturing services; as of 09/24





Late-Stage Development Portfolio: Impactful Revenue Potential¹

Customer	Product Type	2025	2026	2027	2028
Specialty Pharma	Med Device		0		
Large Pharma	Med Device			0	
Specialty Pharma	NDA*			0	
Specialty Pharma	Med Device			0	
Specialty Pharma	NDA			0	
Specialty Pharma	NDA				0
Specialty Pharma	Med Device				0
Specialty Pharma	Med Device		Estimated Annual Revenue Potential ¹ \$10MM + \$5MM - \$10MM \$ < \$5MM	ı О < \$5ММ	0
Specialty Pharma	NDA		7.5	0-	
Specialty Pharma	Med Device				0



^{*}Large Pharma company retains commercial rights to product

1. Assumes full realization of management's estimates for annual commercial revenue potential from pipeline projects as of Sept. 2024 at peak sales (not risk-adjusted). Information presented is not risk and probability-adjusted, and the actual revenue realization may vary significantly. There can be no assurance that such results will occur and that such results will be materially different from actual results.

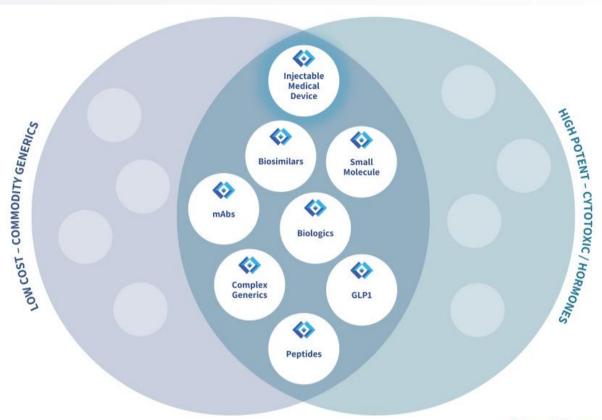






Strategically Expand Target Markets

- Expanding beyond high-viscosity legacy
- Attractive therapeutic areas
- NCEs in Phase 2, Phase 3
- Unique, injectable delivery systems
- Ophthalmic and orthopedic medical devices
- Commercial site transfers









Expanded Targets Lead to Growing Pipeline

Prospective Opportunities

In process of being qualified - Inform & educate on Lifecore capabilities -

Active Opportunities

Within our capabilities with an identified close date

- Strong, diverse and growing universe of 50+ potential future business opportunities¹
- · Mix of both large and specialty pharma
- Subset of opportunities are HA-related, representing a broadening of our pipeline
- Significant number of late-stage development or commercial site transfer programs







New Technology Opens Door to New Business

State-of-the-Art, 5-Head Isolator Filler

- Full isolator technology, state-of-the-art containment
- Significantly expanded available capacity
- Broad capability: vials, syringes & cartridges
- Strengthens compliance





Based on estimates derived from internal testing and historical capacity data. There can be no assurance that such results will occur or that such results will be materially different from actual results.







Expanding Business Development & Brand Awareness

- Team approach to business development
- Adding proven talent
- Expanded visibility at industry events
- Enhanced marketing strategy: SEO, content, social media









About Lifecore - Our Journey

Market Overview

Growing Our Business

Sustaining Objectives

Question & Answer





Reduced Operational Expenses

Working to drive operational efficiency and make OpEx a progressively smaller percentage of revenue:

- Cost-conscious practices
- Optimize key processes
- Eliminate low-value activities
- Leverage technology
- Culture of continuous improvement

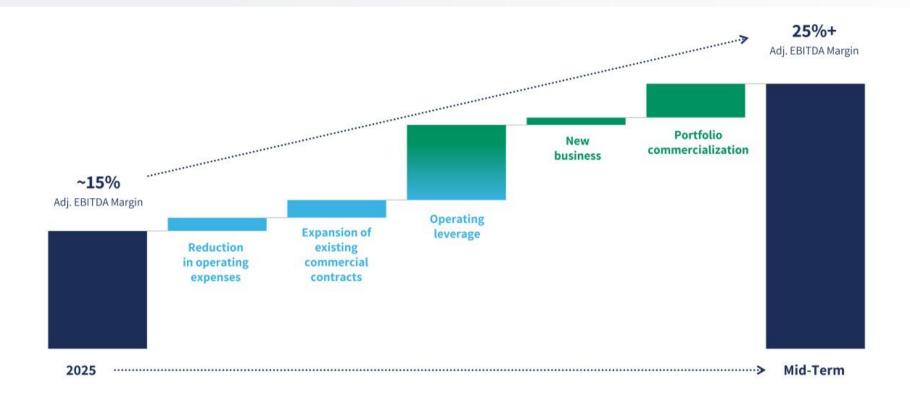


Build long-term efficiencies aligned with growth that leads to improved profitability





Efficiency and Revenue Growth Drive Margin Improvement





For illustrative purposes only, timing, estimates, assumptions and the actual growth of adjusted EBITDA may vary significantly; we may not be able to manage our costs and achieve our anticipated financial goals. The information provided is illustrative only, the growth cycle may not be achieved and there is continued uncertainty relating to any guidance contained herein. There can be no assurance that such results will occur or that such results may be materially different from actual results.



Performance-Driven Culture

Commitment to a high-performance culture driven by exceptional talent

Performance Driven

Metrics / KPI driven Linked to strategic goals Talent Management

Define "A" players Local / National recruitment Incentivize and Recognize

Aligned with shareholder value enhancement Manage, Monitor and Adjust

Consistent review and evaluation of talent





Commitment to Quality

Lifecore provides the highest quality pharmaceutical products and services to our customers and their patients.







40+ Years of Strong Track Record with Global Regulatory Bodies







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Reconciliation of Non-GAAP Financial Measures

	Fiscal Year Ending	Fiscal Year Ended May 26, 2024	
(in thousands)	May 25, 2025		
	(estimate)		
Net Loss (GAAP)	\$ (25,900) — \$ (23,900)	\$ 9,331	
Interest expense, net	20,900	18,090	
Provision for income tax (benefit) expense	7 7.	183	
Depreciation and amortization on property, plant, and equipment	8,600	7,954	
Stock-based compensation	9,700	6,201	
Change in fair value of debt derivatives	(4,800)	(39,500)	
Financing fees (non-interest)	400	3,513	
Reorganization costs (a)	7,100	9,796	
Restructuring costs (a)	1,300	1,656	
Franchise tax equivalent to income tax	200	272	
Contract cancellation and other costs	_	567	
Stockholder activist settlement (a)	1,500	459	
Start-up costs	<u> </u>	1,684	
Adjusted EBITDA	\$ 19,000 — \$ 21,000	\$ 20,206	

To supplement the company's financial results determined by U.S. generally accepted accounting principles ("GAAP"), the company has disclosed in the table below the following non-GAAP information about Adjusted EBITDA. $^{\rm 1}$

Adjusted EBITDA is net (loss) income as determined under GAAP excluding (i) interest expense, net of interest income, (ii) provision for income tax expense (benefit), (iii) depreciation and amortization, (iv) stock-based compensation, (v) change in fair value derivatives, (vi) financing fees (non-interest), (vii) reorganization costs, (viii) restructuring costs, (ix) franchise tax equivalent to income tax, (x) contract cancellations, (xi) stockholder activist settlement costs, and (xii) start-up costs.

The company believes that non-GAAP financial measures, such as Adjusted EBITDA, are helpful in understanding its business as it is useful to investors in allowing for greater transparency of supplementation information used by management. Adjusted EBITDA, is used by investors, as well as management in assessing the company's performance. Non-GAAP financial measures should be considered in addition to, but not as substitute for, reported GAAP results. Further, non-GAAP financial measures, even if similarly titled, may not be calculated in the same manner by all companies, and therefore should not be compared.

(a) We previously estimated restructuring, reorganization, stockholder activist settlement costs to be in a range of \$5.5 to \$6.5 million, which we now estimate will be approximately \$9.9 million.

Reorganization costs include costs not expected to be incurred on a normalized basis associated with Lifecore becoming a stand-alone entity, divestitures, litigation related with former owners of acquired businesses, restatements of financial statements and change in auditors.

Restructuring costs are related to board approved actions consisting primarily of employee severance, lease cost of exited facilities, and costs associated with divested businesses.



See disclaimers and important information on Slides 2 and 3





About Lifecore - Our Journey

Market Overview

Growing Our Business

Sustaining Objectives

Question & Answer



