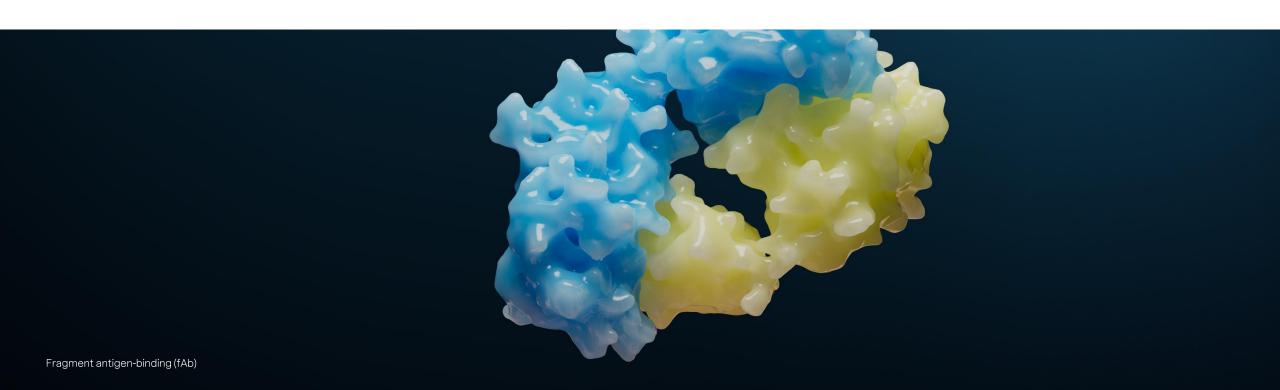


Lonza – Capturing Growth and Value in the CDMO Market

Philippe Deecke, Chief Financial Officer

9 January 2024



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Lonza today and our path forward

How we win

Concluding remarks

Q&A



A Leading Strategic Partner to the Healthcare Industry

Attractive Business Delivering Strong KPIs and Financials

2022 Figures

>1,025 Molecules^{1,2}

~800 CDMO customers

Large Ongoing Growth Projects³

~70% of **CDMO** sales in commercial

6.2_{bn} Sales in CHF

+16% Sales CAGR 2019 – 2022 in CER⁴

CORE4 EBITDA margin

23% Average Capex as % Sales 2019 - 2022

¹ Large molecules including mammalian, microbial, bioconjugates and cell and gene therapy products (pers. medicines included for pre-clinical and clinical molecules only)

² Small molecules including active pharmaceutical ingredients (API), Highly Potent API (HPAPI), dosage form and delivery systems and particle engineering

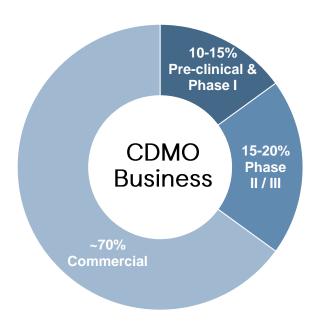
³ Projects started since 2018

⁴ CORE results and Constant Exchange Rates (CER) are non-IFRS measures. For Lonza's definition of CORE results, also refer to the Alternative Performance Measures Brochure published in conjunction with the Lonza Half-Year Report 2023

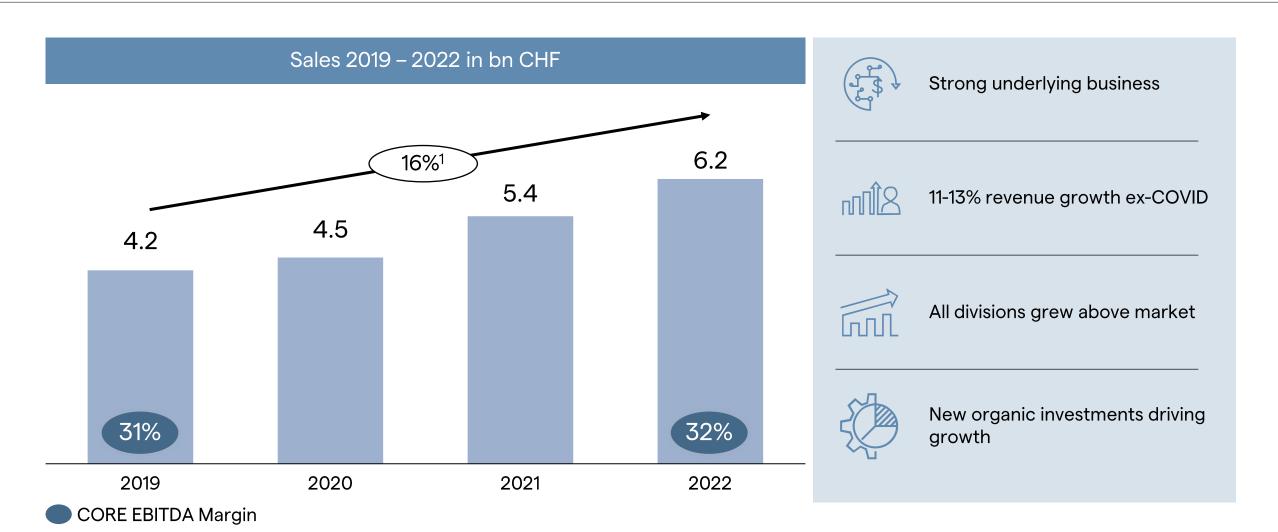
Unique End-to-End Offering Across Multiple Modalities Creating Competitive Advantage



Majority of the Business From Commercial Products



Lonza's Journey of Delivering Strong Topline Growth



Our Strategic Priorities



INNOVATION

- Anticipating technological trends
- **Building IP** and protected capabilities
- Investing in data analytics and Al



COLLABORATION

- Long-term partnership with customers
- Creating joint customized solutions



SERVICE

- Best-in-class quality track record
- Customer satisfaction and retention



PERFORMANCE

- Continuous process improvements
- Accelerate path to drug commercialization



VALUE CREATION

- Deliver on Lonza's guidance
- Disciplined capital allocation
- Delivering for shareholders. stakeholders and society



ESG

- Provide safe workplaces and foster employee participation
- ESG targets in remuneration policy since 2022
- Committed to SBTi

Lonza is Uniquely Positioned to Capture Growth and Value in CDMO Market

Global leader and pure-play CDMO

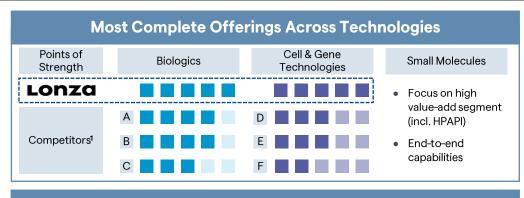
End-to-end offering across all divisions

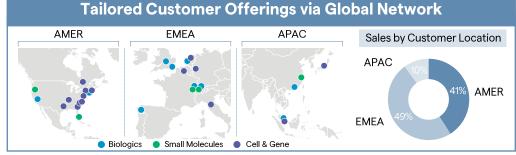
Strong and long-standing customer relationships

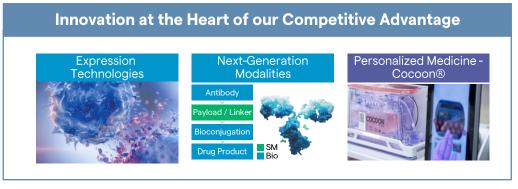
Global, western-focused manufacturing network

Strong project execution capabilities

Leader in innovation and industry pioneer







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Dynamic End-Markets with Attractive Growth Outlook

Strong end market growth

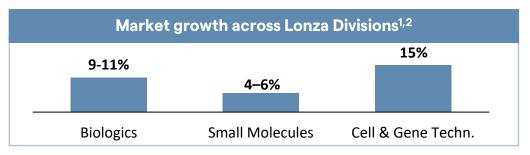
Continued growth in CDMO share of business

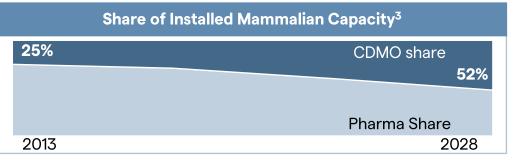
Further tightening of biologics capacity utilization

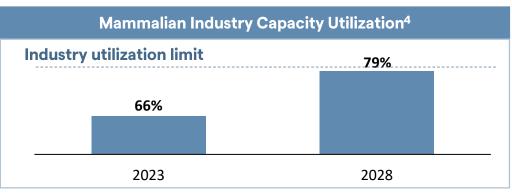
Record number of small molecules and cell & gene molecules in development

Market competes on quality, regulatory expertise above price

High risks for new market entrants (capital intensity, technological capabilities, regulatory and quality expertise)







¹²⁰²³⁻²⁰²⁸ CAGR for Biologics and Small Molecules in USD bn, 2024-2028 CAGR for Cell & Gene Technologies in CHF bn

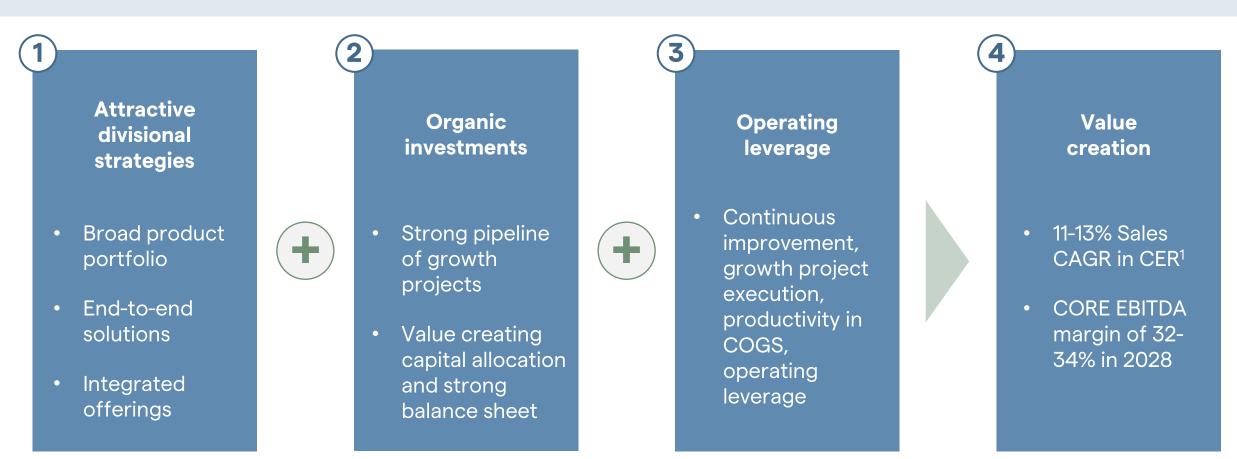
² Source: Lonza internal analysis, EvaluatePharma, Citeline

³ Source: Lonza internal analysis; publicly announced capital expansions (2023)

⁴Source: Lonza internal analysis, IQVIA, EvaluatePharma, Citeline, publicly announced capacity expansions (2023)

Lonza Delivers Value Through Growth Investments and Operational Excellence

The CDMO market remains highly attractive with strong demand from end-markets Lonza is serving



)Biologics

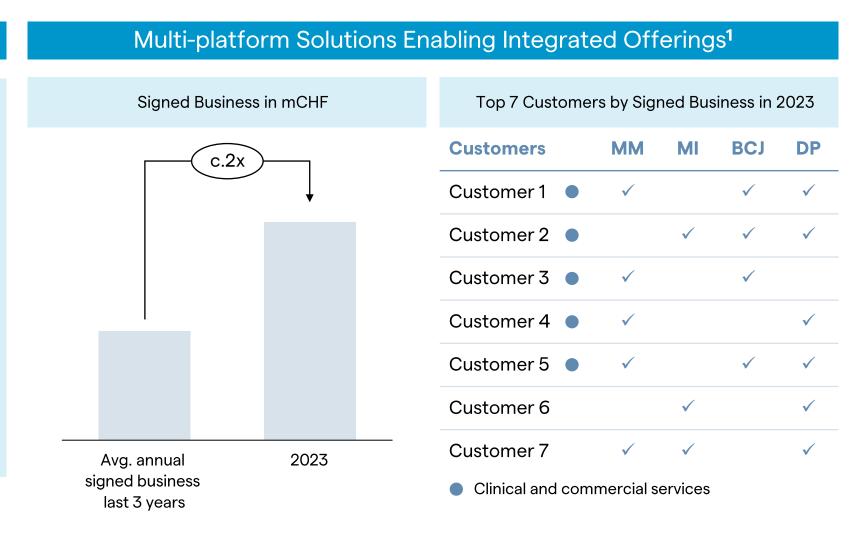
Delivering a Complete CDMO Offering Across the Lifecycle

Strategic Priorities

Full lifecycle management

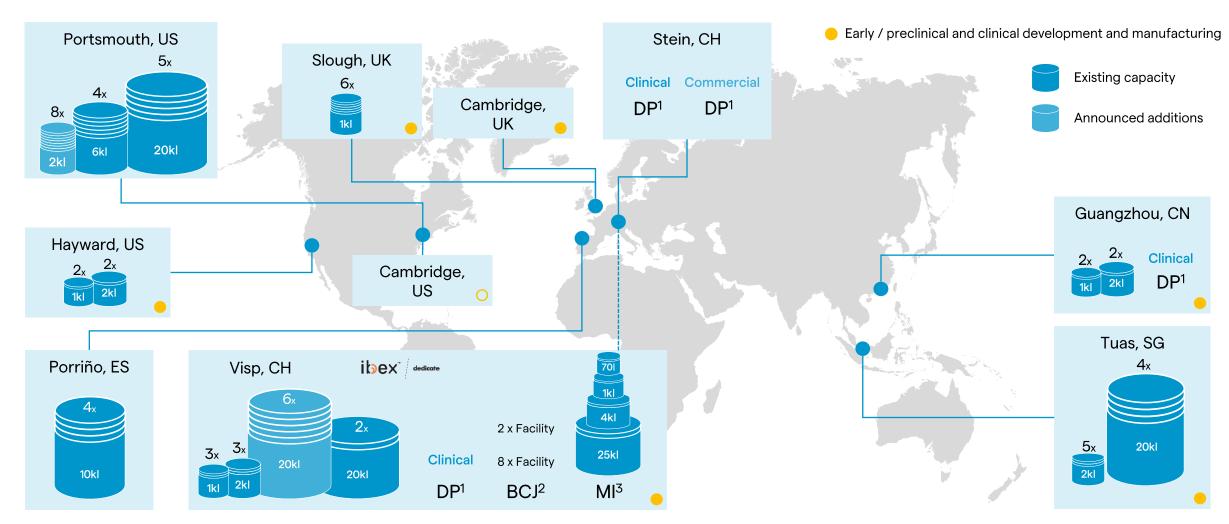
End-to-end offering

Sustainable global manufacturing, site specialization



(1)Biologics

Tailored Customer Offerings on a Global Scale



1)Small Molecules

Meeting our Customers' Complex Small Molecules Needs

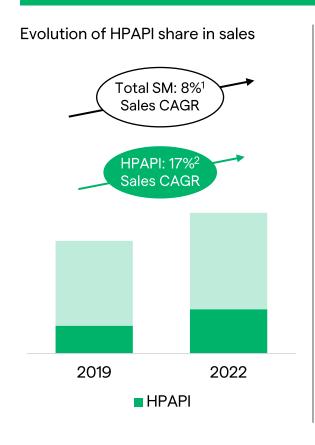
Strategic Priorities

Be a strong development partner driven by science

Be present throughout the product lifecycle

Make the customer experience great

Leading Capabilities in Highly Potent API (HPAPI)





HPAPI Drug Substance manufacturing Clinical to commercial scale-up Large pharma – long-term contract

Operational since 2021



Dedicated ADC production suite

Clinical to commercial scale-up

Large pharma – long-term contract

Operational in 2024

Integrated supply chain with **Biologics**

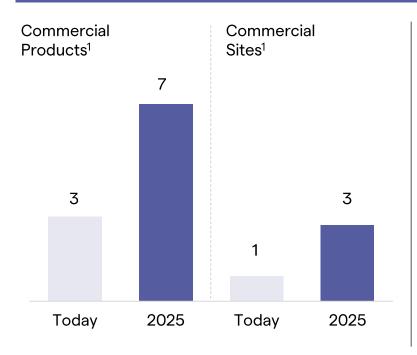
Cell & Gene

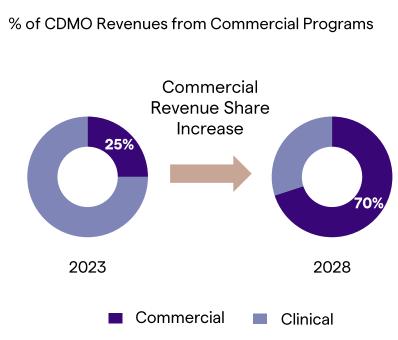
Driving Commercialization of Cell & Gene Therapies

Strategic Priorities

- Manufacturing expertise
- Commercialization track record
- Innovation
- Partnerships with frontrunners

Increasing Focus on Commercial Portfolio in CGT





Three commercial products already in portfolio and four more expected in the next 18 months

(1)Capsules & Health Ingredients

Leading with a High-Value and Innovative Capsule and Service Offering

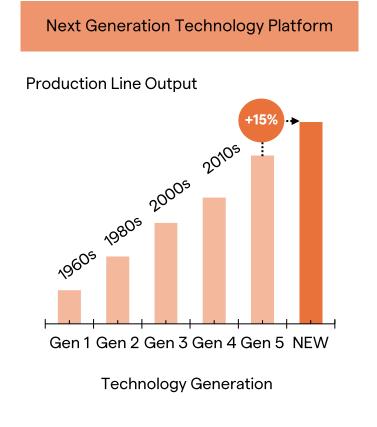
Strategic Priorities

Growth above the market

Process innovation and network optimization

Product and service innovation

Process and Product Innovation to Extend Leadership

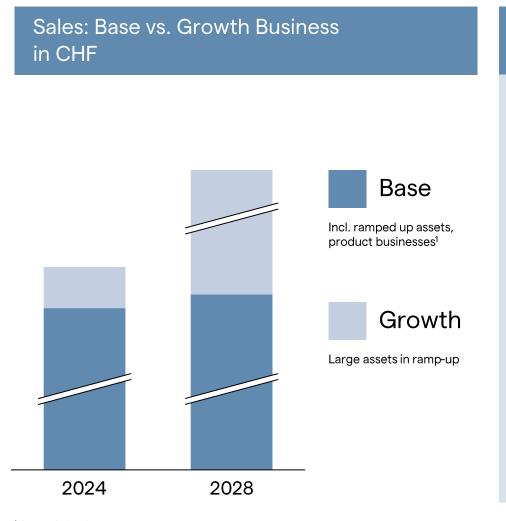




1 Ahead of Market Sales Growth and Margin Improvements Across Divisions

Division	Sales CAGR 2024 – 2028 (CER)	CORE EBITDA margin 2024 – 2028
Biologics	Mid-teens	>35%
Small Molecules	Mid-to-high single-digit	>30%
Cell & Gene	Mid-teens	>25%
Capsules & Health Ingredients	Low-to-mid single-digit	>30%

(2)Large and Stable Base Complemented by Dynamic Growth Portfolio



Base Business

Stable, growing and resilient business

High asset utilization

High revenue visibility

Strong cash generation

Strong margin

Growth

New assets

Fast growth

Low margin during ramp-up

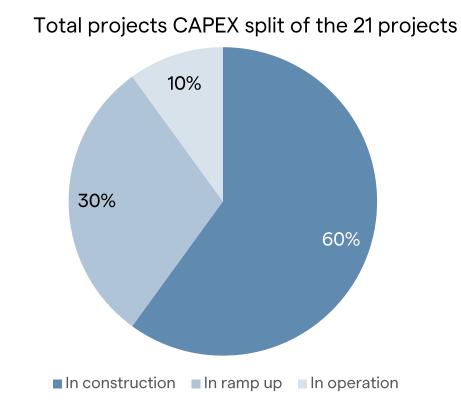
Margin accretive at peak

De-risked through contracting

(2) Growth Project Portfolio at a Glance



CAPEX above CHF 50m



Modalities

CHF 1.8bn Cumulative sales 2018 - 2023E

90% invested in commercial / mixed assets

75% of Capex invested in growth 2019-2022

15% IRR threshold

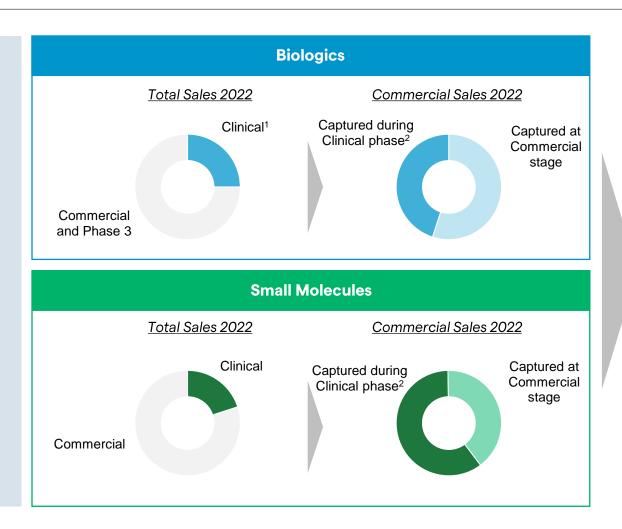
30% **ROIC** threshold (at peak)

(2) Trusted Partner for Scale-up and Commercialization

Acquiring molecules early in the lifecycle - key to fuel future commercial pipeline

Targeted investments in earlystage capacity and capabilities

Proven track record of successful commercialization

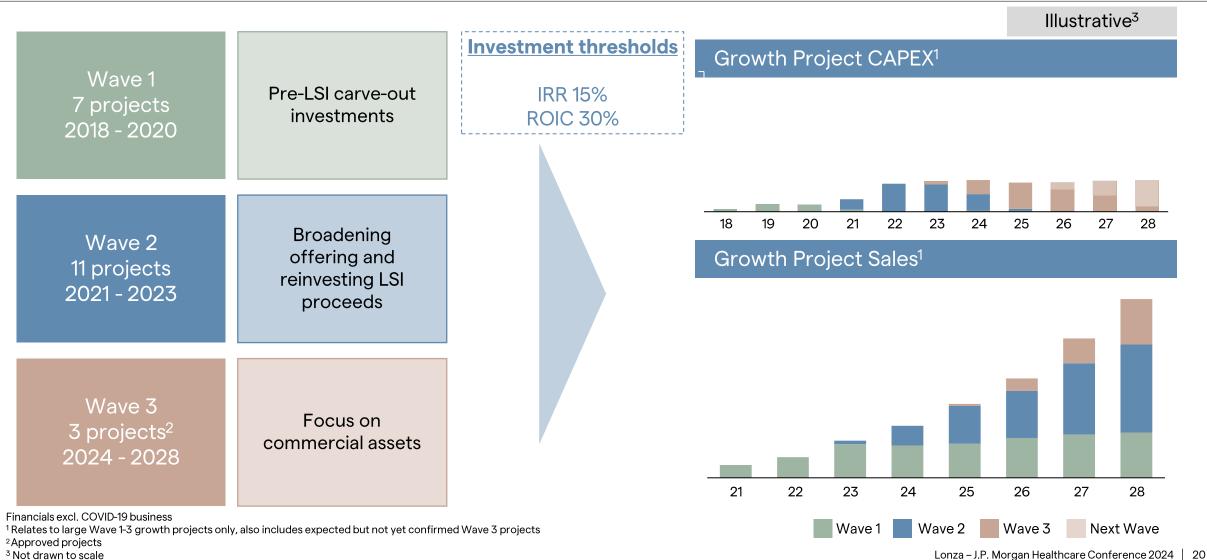


c.50% of commercial sales captured in Phase II or earlier

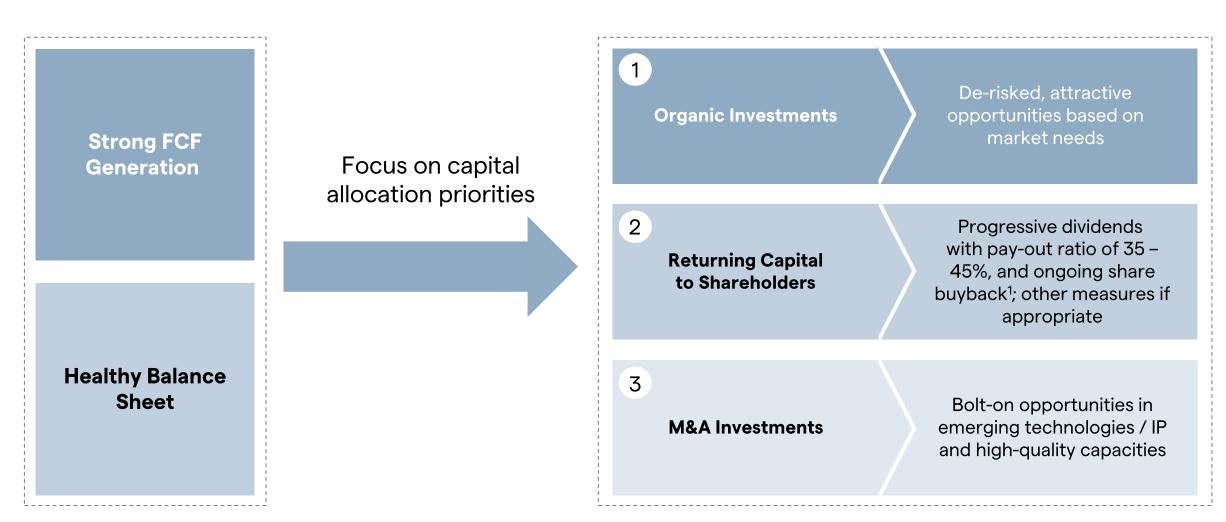
¹Defined for Biologics as programs up to and including phase 2

² Sales % generated from molecules captured in pre-clinical, clinical Phase 1 and 2

(2)Investment Program Securing Industry Leadership



(2)Our Capital Allocation Strategy Prioritizes Organic Growth, Shareholder Return and Bolt-On M&A



¹ Note: In April 2023 Lonza launched share buyback program of up to CHF 2bn over a maximum period of 2 years, as of end of November 2023 Lonza repurchased shares of CHF 890m corresponding to 2.8% of Lonza's share capital

(4) Reiterating our Group Mid-Term Guidance

Sales growth driven by new Biologics assets

Margin improvement from growth projects, Cell & Gene maturing, operational excellence

CAPEX expected to decrease to mid-to-high teens as % of sales by 2028

Strong balance sheet with commitment to investment grade rating of BBB+

11 – 13% Sales CAGR in CER (2024 - 2028)

32 – 34% CORE EBITDA margin in 2028

Double-digit ROIC in 2028

1.5 - 2.0xNet Debt / CORE EBITDA

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Capturing Growth and Value in the CDMO Market

Organic investments in growing modalities to capture value in attractive market

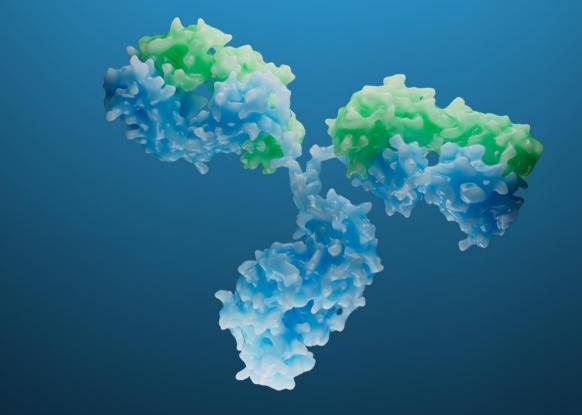
Broad range of technologies and capabilities to meet complex customer needs

Best-in-class and industry reference in technical and regulatory expertise

Solid strategies and world-class teams in place to maintain leadership role in industry

Confident in delivering on our guidance

Lonza



Q&A

Additional Information and Disclaimer

Lonza Group Ltd has its headquarters in Basel, Switzerland, and is listed on the SIX Swiss Exchange. It has a secondary listing on the Singapore Exchange Securities Trading Limited ("SGX-ST"). Lonza Group Ltd is not subject to the SGX-ST's continuing listing requirements but remains subject to Rules 217 and 751 of the SGX-ST Listing Manual.

Forward-looking statements contained herein are qualified in their entirety as there are certain factors that could cause results to differ materially from those anticipated. Any statements contained herein that are not statements of historical fact (including statements containing the words "outlook," "guidance," "believes," "plans," "anticipates," "expects," "estimates" and similar expressions) should be considered to be forward-looking statements. Investors are cautioned that all forward-looking statements involve risks and uncertainty.

There are a number of important factors that could cause actual results or events to differ materially from those indicated by such forwardlooking statements, including the timing and strength of new product offerings; pricing strategies of competitors; the company's ability to continue to receive adequate products from its vendors on acceptable terms, or at all, and to continue to obtain sufficient financing to meet its liquidity needs;

difficulty to maintain relationships with employees, customers and other business partners; and changes in the political, social and regulatory framework in which the company operates, or in economic or technological trends or conditions, including currency fluctuations, inflation and consumer confidence, on a global, regional or national basis.

In particular, the assumptions underlying the section "Looking to the Future" herein may not prove to be correct. The statements in the section "Looking to the Future" constitute forward-looking statements and are not guarantees of future financial performance.

Lonza's actual results of operations could deviate materially from those set forth in the section "Looking to the Future" as a result of the factors described above or other factors. Investors should not place undue reliance on the statements in the section "Looking to the Future". Except as otherwise required by law, Lonza disclaims any intention or obligation to update any forward-looking statements as a result of developments occurring after this presentation was published.