Hong Kong Exchanges and Clearing Limited and The Stock Exchange of Hong Kong Limited take no responsibility for the contents of this announcement, make no representation as to its accuracy or completeness and expressly disclaim any liability whatsoever for any loss howsoever arising from or in reliance upon the whole or any part of the contents of this announcement.



WUXI BIOLOGICS (CAYMAN) INC.

藥明生物技術有限公司*

(Incorporated in the Cayman Islands with limited liability)
(Stock Code: 2269)

INSIDE INFORMATION 2023 ANNUAL RESULTS PRESENTATION

This announcement is made by WuXi Biologics (Cayman) Inc. (the "Company") pursuant to Rule 13.09 of the Rules Governing the Listing of Securities on The Stock Exchange of Hong Kong Limited (the "Listing Rules") and the Inside Information Provisions (as defined in the Listing Rules) under Part XIVA of the Securities and Futures Ordinance (Chapter 571 of the Laws of Hong Kong).

In order to enable shareholders and investors to have a deeper and more comprehensive understanding of its 2023 annual results and business operations, the Company will convene conference calls at 9:30 a.m. and 8:30 p.m. (Hong Kong time) on March 27, 2024, at which it will conduct a presentation regarding the Company's financial results and business operations (the "**Presentation**"). Investors may attend the conference calls at the scheduled time at the following links:

Conference call in Chinese at 9:30 a.m. (Hong Kong time):

If you are within the mainland of China:

https://morganstanley.cnwebcasts.cn/starthere.jsp?ei=1660437&tp_key=f2bee92149

If you are outside the mainland of China:

https://morganstanley.webcasts.com/starthere.jsp?ei=1660437&tp_key=f2bee92149

Conference call in English at 8:30 p.m. (Hong Kong time):

https://jefferies.zoom.us/webinar/register/9117095533914/WN_Ey-wgpL5Qt2pcTHGEZ85UQ

Please use the links provided above to complete the online registration process in advance of the conference calls.

Further, to ensure that all shareholders and investors have equal and timely access to such information, the Company has included in this announcement the full copy of the Presentation. Shareholders and investors are reminded that the Presentation may contain forward-looking statements, which are, by their nature, subject to risks and uncertainties, and any estimate and future proposals stated in the Presentation are based on certain assumptions and estimates and on management's judgements in light of currently available information only.

Shareholders and potential investors of the Company are advised not to place undue reliance on the information contained in the Presentation and should exercise caution when dealing in the securities of the Company.

By order of the Board
WuXi Biologics (Cayman) Inc.
Dr. Ge Li
Chairman

Hong Kong, March 26, 2024

As at the date of this announcement, the board of directors of the Company comprises Dr. Zhisheng Chen and Dr. Weichang Zhou as executive Directors; Dr. Ge Li, Dr. Yibing Wu and Mr. Yanling Cao as non-executive Directors; and Mr. William Robert Keller, Mr. Kenneth Walton Hitchner III, Mr. Jackson Peter Tai and Dr. Jue Chen as independent non-executive Directors.

* For identification purpose only

Global Premier CRDMO: Enabling Global Partners and Delivering Sustainable Growth

2023 Annual Results
March 2024





Stock Code: 2269.HK





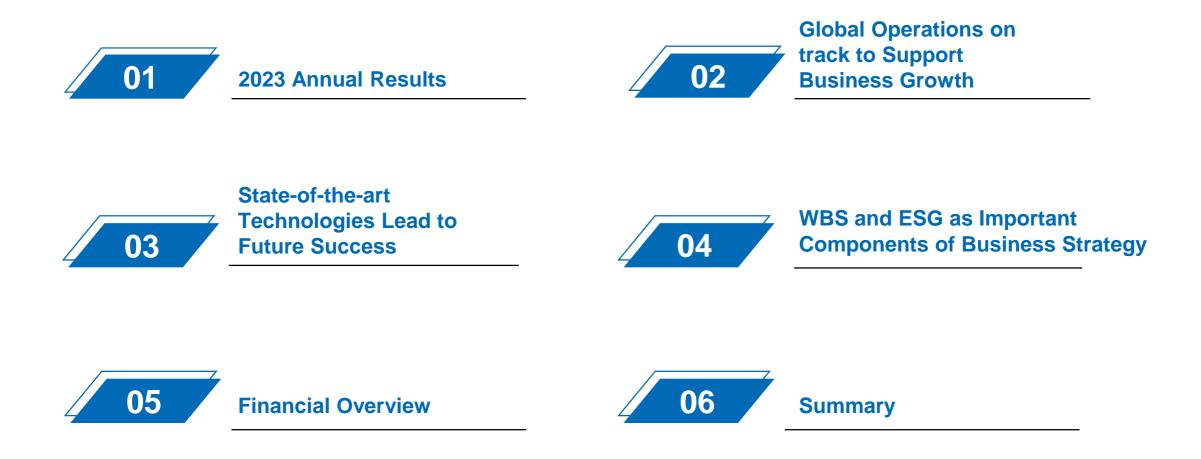
This presentation may contain certain "forward-looking statements" which are not historical facts, but instead are predictions about future events based on our beliefs as well as assumptions made by and information currently available to our management. Although we believe that our predictions are reasonable, future events are inherently uncertain and our forward-looking statements may turn out to be incorrect. Our forward-looking statements are subject to risks relating to, among other things, the ability of our service offerings to compete effectively, our ability to meet timelines for the expansion of our service offerings, and our ability to protect our clients' intellectual property. Our forward-looking statements in this presentation speak only as of the date on which they are made, and we assume no obligation to update any forward-looking statements except as required by applicable law or listing rules. Accordingly, you are strongly cautioned that reliance on any forward-looking statements involves known and unknown risks and uncertainties. All forward-looking statements contained herein are qualified by reference to the cautionary statements set forth in this section.

Use of Adjusted Financial Measures (Non-IFRS Measures)

We have provided adjusted net profit, adjusted net profit margin, adjusted gross profit, adjusted gross profit margin, adjusted EBITDA, adjusted EBITDA margin and adjusted basic earnings per share for the corresponding periods, which excludes the share-based compensation expenses, listing expenses, gains or losses from equity investments and foreign exchange gains or losses, and are not required by, or presented in accordance with, IFRS. We believe that the adjusted financial measures used in this presentation are useful for understanding and assessing underlying business performance and operating trends, and we believe that management and investors may benefit from referring to these adjusted financial measures in assessing our financial performance by eliminating the impact of certain unusual and non-recurring items that we do not consider indicative of the performance of our business. However, the presentation of these non-IFRS financial measures is not intended to be considered in isolation or as a substitute for the financial information prepared and presented in accordance with IFRS. You should not view adjusted results on a stand-alone basis or as a substitute for results under IFRS, or as being comparable to results reported or forecasted by other companies.

CONTENTS





2023 Annual Results



588 ^{18.7}% **698**

Integrated Projects YoY

37.7%

Non-COVID Revenue Growth (YoY)

132

New Projects Added

17^{41.2%} 24

Commercial Projects YoY

20.59

Total Backlog (US\$ Bn)

12,740/4,432

Employees / Development Scientists



15.27^{11.6%} **17.03**

Revenue (RMB Bn) YoY

5.05 ^{-2.0%} **4.95**

Adj Net Profit (RMB Bn) YoY

45.3%

Adj Gross Profit Margin

29.1%

Adj Net Profit Margin

41.1%

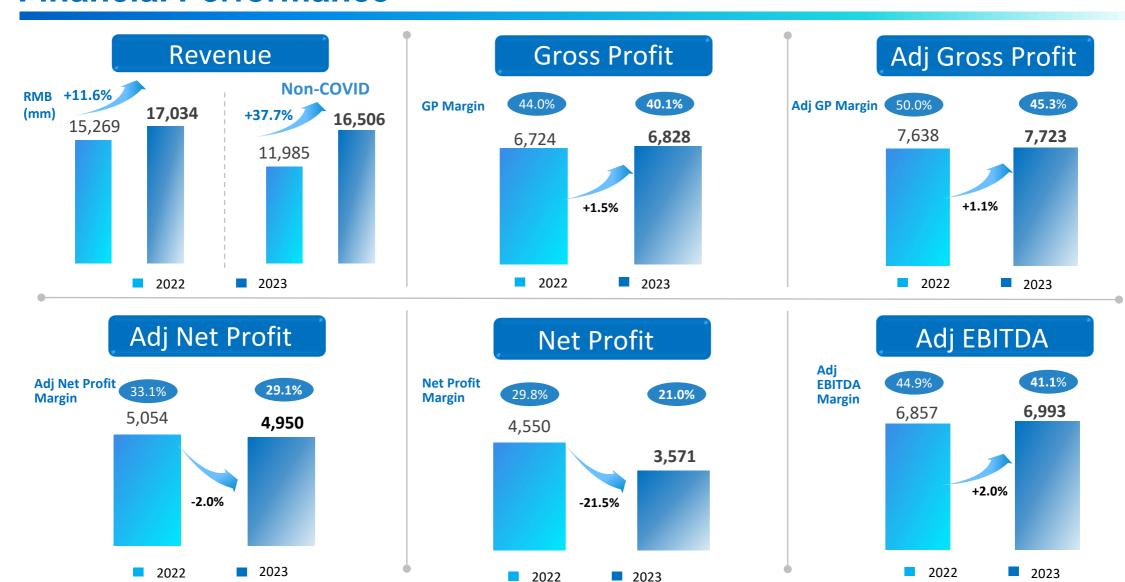
Adj EBITDA Margin

1.13

Adj. Basic EPS (RMB)

Financial Performance





Key Financials



AVAILABLE FUNDS

- Available funds approx. RMB11.2 bn as of Dec 31, 2023
- Total Liability to Equity Ratio 28.5%, expect to have sufficient funds for capacity increase

CAPEX

- 2023 CAPEX approx. RMB4.1 bn, mainly for capacity increase in Europe and U.S.
- 2024 CAPEX Plan: approx. RMB5 bn

LOAN

- Approx. RMB2.1 bn borrowings as of Dec 31, 2023
- Available bank credit facilities of around RMB6 bn

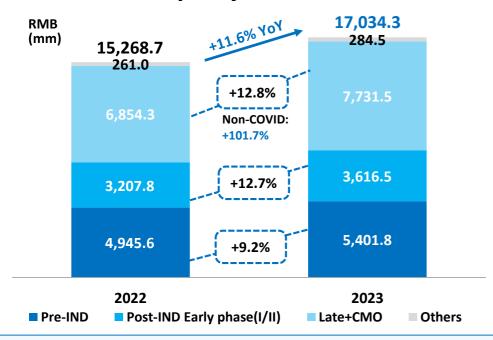
CASH FLOW

- Positive Free Cash Flow of RMB0.6 bn in 2023
- Continue to target free cash flow positive in 2024

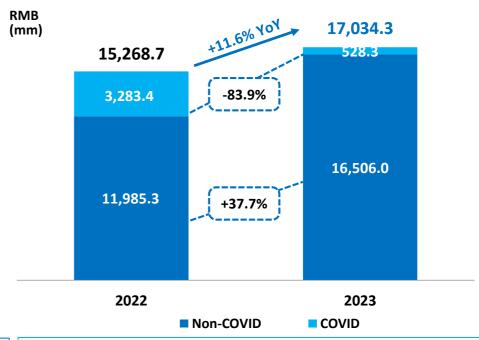
Non-COVID Projects Continued to Accelerate and Delivered Steady Revenue Growth







COVID VS Non-COVID

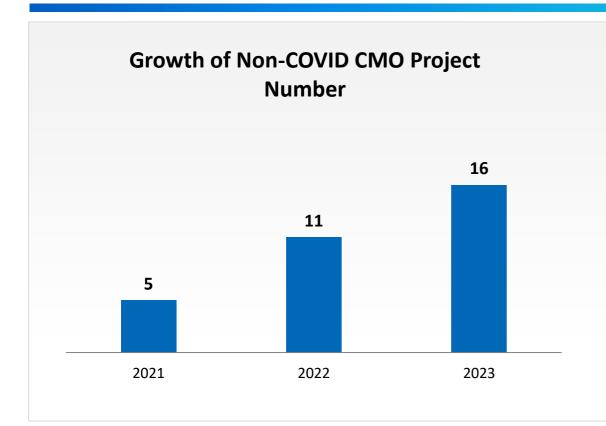


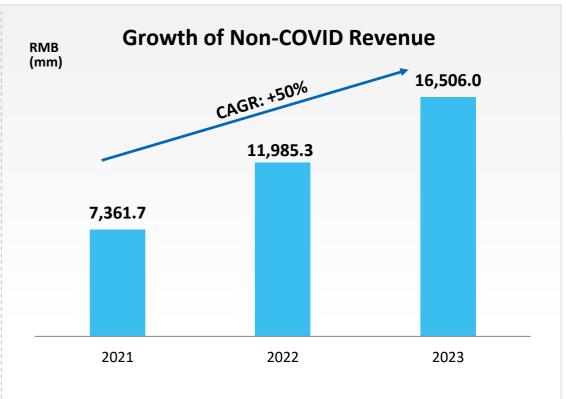
- Due to biotech funding constraints, pre-IND revenue increased by 9.2%. With more projects signed in 4Q, revenue from pre-IND is expected to regain its growth momentum
- Early-phase revenue grew by 12.7% YoY, contributed by the strong execution of "Follow and Win the Molecule" strategies. The inflow of our early-phase projects will continue to feed the late-phase and commercial manufacturing growth
- Late-phase and CMO revenue achieved steady growth despite of the declining revenue from COVID, non-COVID revenue from late phase and CMO stage grew 101.7% YoY

- Non-COVID projects delivered 37.7% YoY revenue growth in 2023, , despite of the diminishing COVID sales
- With COVID transition behind us, our non-COVID technology platforms and speed of execution will drive our growth in the long-run



Non-COVID Projects Driving Growth with Sustained Momentum



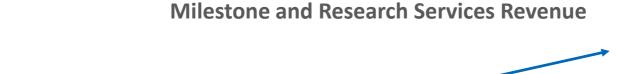


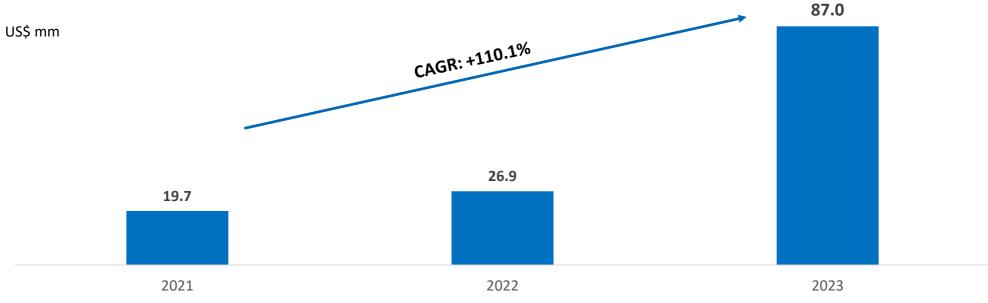
- CMO growth and the Non-COVID growth are the sustainable pillars for the Company's growth
- The successful implementation of "Follow and Win the Molecule" strategies will enhance the growing momentum of CMO projects and boost the future growth

R in CRDMO



Accelerated Growth in Milestone and Research Services Revenue

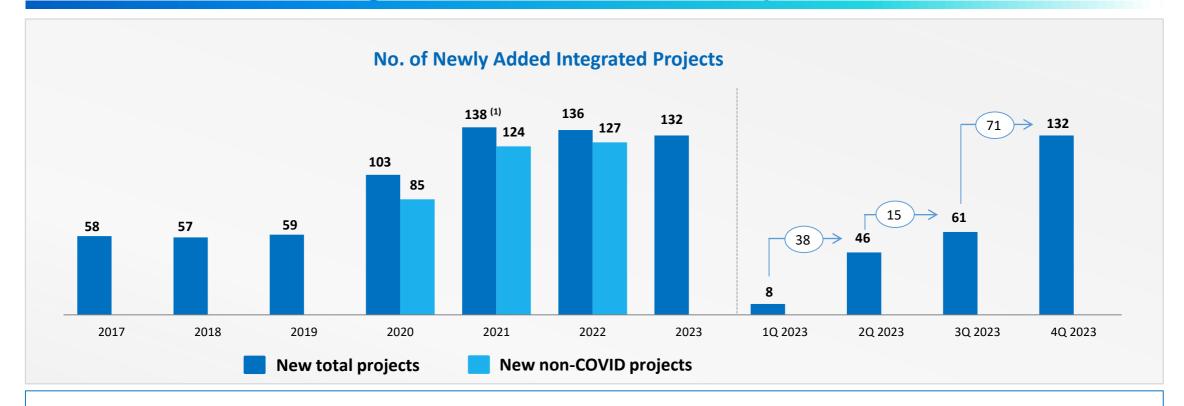




- Benefitted from the comprehensive technology platforms, milestone and research services revenue achieved 110.1% CAGR growth from 2021 to 2023, demonstrating the accelerated growth in R and solidifying growth potential
- The Company will continue to enhance its capabilities to expand service business, enable more discovery projects, thus optimizing the spectrum of services offered to the global biologics industry and bringing more business opportunities

New Projects Rebound from the Bottom in Q1 2023: 2023 is One of the Highest Levels of New Projects





- Number of new projects impacted in Q1 2023 by global biotech slowdown and rebounded since Q2 2023. Strong recovery in 2H 2023 and 2023 is one of the highest No. of new projects, indicating more brand recognition and trust from the industry
- Strong growth in drug development with more projects inflow

Notes

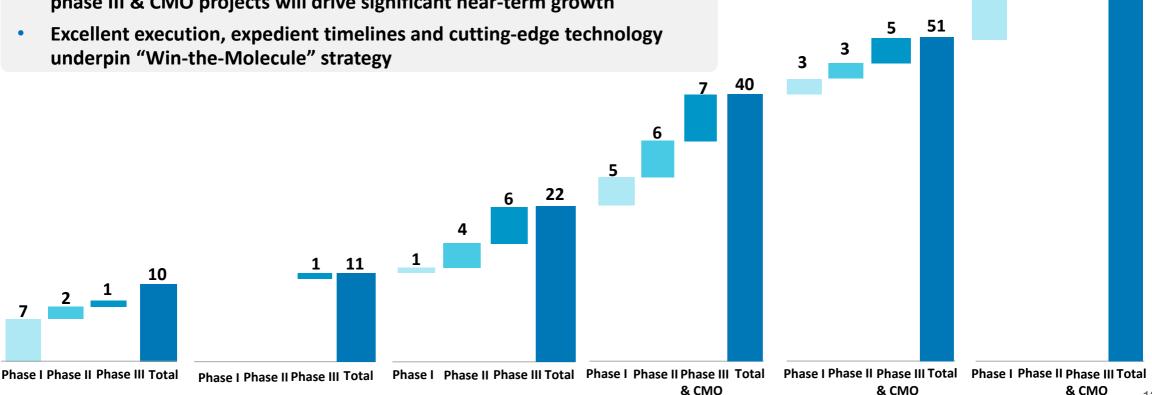
- 1. Newly-added integrated project number in 2021 has excluded the projects from CMAB acquisition
- 2. The rest market primarily includes Singapore, Japan, South Korea, and Australia

"Win-the-Molecule" Strategy: Another Driver to Expand **Pipeline**





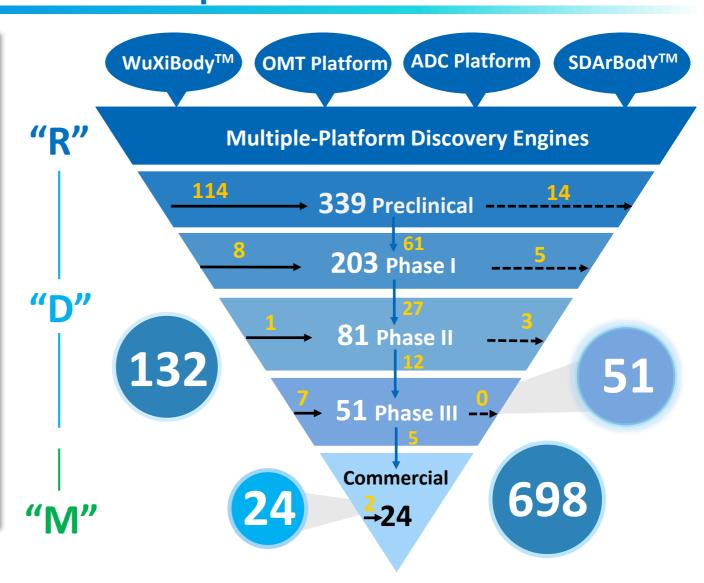
- 9 phase III and Commercial "Win-the-Molecule" projects: record high, significantly higher than our original expectation of 5
- Total 69 projects at different stages (Phase I, II and III + CMO) transferred from global CDMOs or big pharmas to WuXi Biologics since 2018: 29 phase III & CMO projects will drive significant near-term growth
- **Excellent execution, expedient timelines and cutting-edge technology** underpin "Win-the-Molecule" strategy



New Projects Exceeded Expectations: Demonstrate Business Resilience Despite Headwinds



- Continue to gain growth opportunities for both "Follow and Win the Molecule"
- Despite headwinds, newly-added projects reached one of the highest levels since inception, demonstrating the Company's resilience during the downturn cycle and ability to maintain the business growth
- "Win-the-Molecule" strategy continued to excel: 18 external projects transferred into the pipeline as of Dec 31, 2023, including 7 phase III projects and 2 CMO for blockbuster products
- Milestone and research services revenue exceeded US\$87 million, indicating the accelerated growth in R services
- 51 Phase III projects and 24 CMO projects: poised for future manufacturing revenue growth



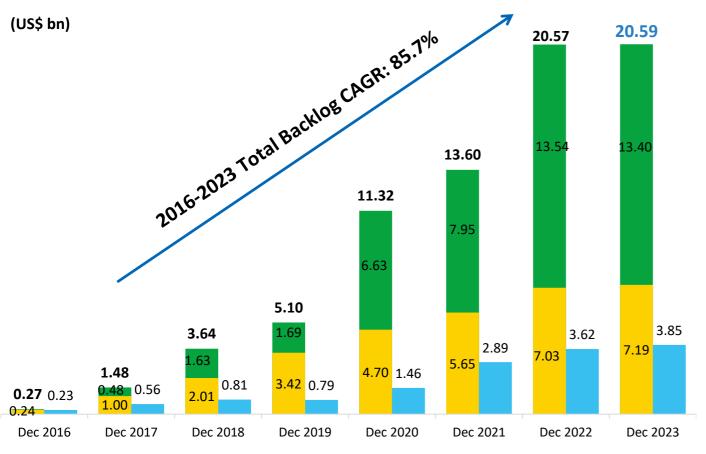
Notes:

As of Dec 31, 2023

Backlog Underpins Future Performance



- **Service Backlog**
- **Upcoming Potential Milestone Fees** (1)
- **Backlog within 3 Years**



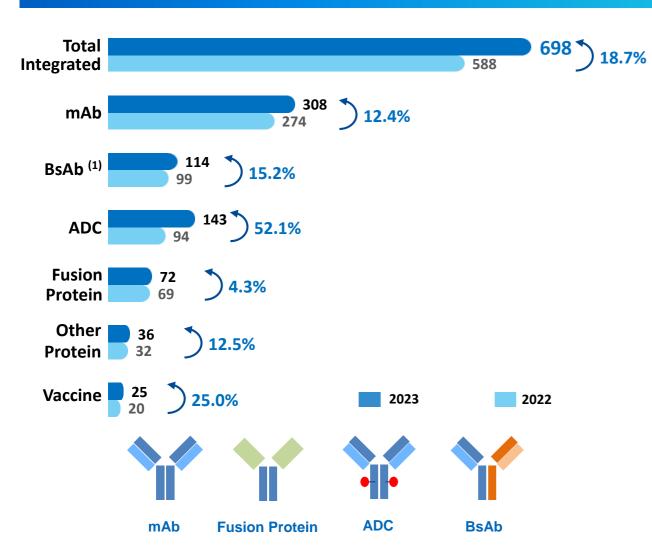
- As of Dec 31, 2023, total backlog reached US\$20.6 bn. US\$13.4 bn service backlog as of Dec 31, 2023, the slight decrease is partially attributable to the COVID cancellation
- **Upcoming potential milestone backlog** reached a record high of US\$7.2 bn, research service capability got validated again to benefit from technology enabling platforms and unique CRDMO business model
- As of Dec 31, 2023, backlog within 3 years approximated US\$3.9 bn, providing high visibility of short-term growth
- **Excellent execution of pre-clinical projects** and increased capacity would shorten the order to fulfillment cycle
- Strong backlog does not indicate lack of capacity for new projects. Additional projects can be initiated within 4 weeks

Results may not foot due to rounding

Upcoming milestone revenue may take longer to receive at the various development stages as it depends on the success rate and progress of the projects

Rich Pipeline across All Biologics Modalities





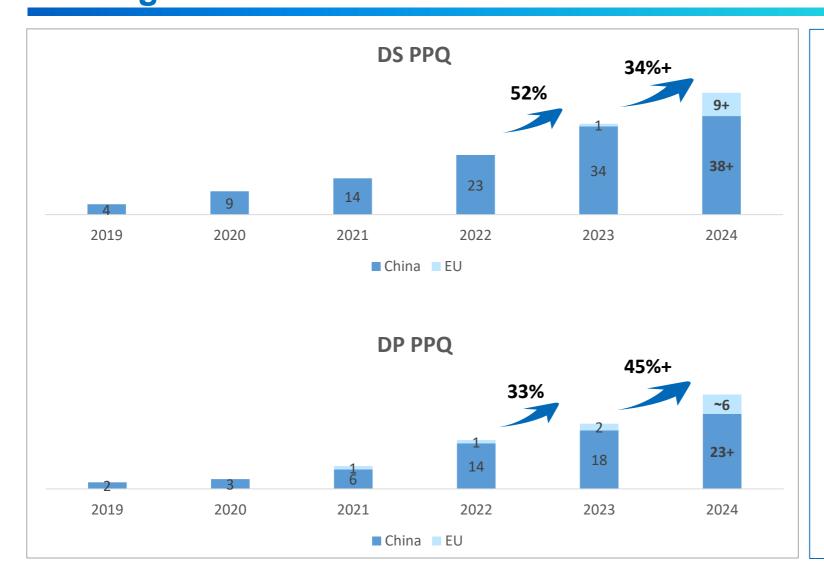
- 259 First-in-class programs
- 25 vaccine projects, including 19 non-COVID vaccines
- 114 bispecific projects covering different formats, several in phase III and commercial stage with upside potential
- 143 Antibody Drug Conjugates (ADC) projects with 52.1% YoY growth driven by increasing industry demands with ~35% global ADC outsourcing market CAGR growth between 2018 and 2022 and ~28% CAGR growth between 2022 and 2030
- 19 CNS (Central Nervous System) programs from domestic and global companies with exciting potential, the fastest one already in phase III
- One of the largest portfolios of complex biologics, consisting of mAbs, bispecifics, multispecifics, ADCs, fusion proteins and vaccines, etc.

Notes

^{1.} As of Dec 31, 2023, compared with projects number as of Dec 31, 2022

CMO Continues with Strong Momentum as Shown by Leading Indicators: No. of PPQs





- Robust growth in drug substance (DS) and drug product (DP) PPQs, in line with our CMO project increase: will add more in 2024
- Growing number of PPQs
 executed in 2023 and
 scheduled for 2024, indicating
 the solid growth for future
 CMO
- PPQ success of 97%+: one of the best performers in the industry, premier quality demonstrated

Sustained Growth with Diverse Engines





North America 20.2% Non-COVID Growth





30.2% of Revenue

47.4% of Revenue

18.3% of Revenue

4.1% of Revenue

- Europe: the market with the fastest growth rate. Non-COVID revenue grew by 172.4% YoY and EU market accounted for 30.2% of our total revenue in 2023, increased from 16.7% in 2022. Extended more collaboration with MNCs and witnessed more revenue contribution from them. Expect continued strong performance from EU market
- North America: Non-COVID revenue grew by 20.2% YoY. Riding on unwavering execution and established track record, WuXi Bio continues to enable the clients in North America
- China: -1.2% non-COVID revenue growth attributable to the impact from biotech funding constraints. Collaborated more with high-quality clients, especially those who licensed out the products to global MNCs. Signed more projects in Q4 and still maintain positive outlook for Chinese market in the long run
- Rest of the World: continued to enable customers, saw massive potential in this region especially in Southeast Asia and explored more collaborative opportunities

Bispecifics to be Another Growth Driver − WuXiBodyTM

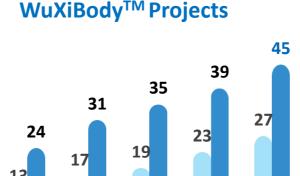


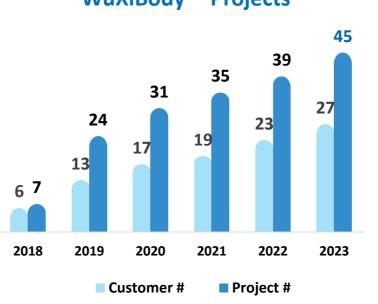


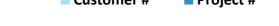
Empower to discover best or first-in-class molecules

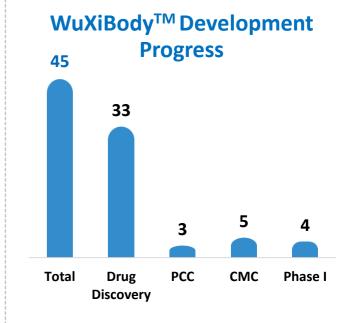












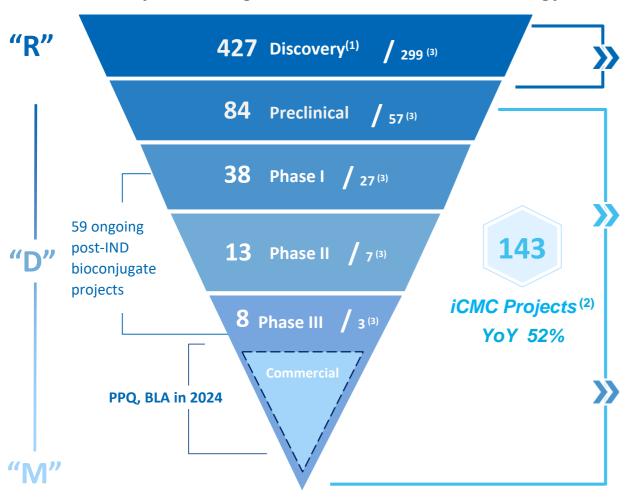
- WuXiBodyTM continues to gain worldwide recognition, with 45 projects as of Dec 31, 2023
- 4 projects at Phase I, 5 projects at CMC, and 3 projects at PCC, demonstrating state-of-the-art technology of WuXiBody™
- 2 WuXiBody[™] projects are expected to get IND approval in 2024

18

WuXi XDC's Powerful Platform and Business Model to Secure Project Wins and Fuel The Growth



Number of Projects Through "Enable - Follow - Win" Strategy



"Research": Enable discovery to PCC

36 projects advanced from discovery to iCMC stage ⁽¹⁾

"Development": Fastest route to IND

Capacity to support 40 INDs each year

Seamless transition between early & late stage PD

50 new integrated projects signed in 2023

21 Phase II & Phase III projects

"Manufacturing": Commercialization

5 PPQ projects with more potential BLA submissions

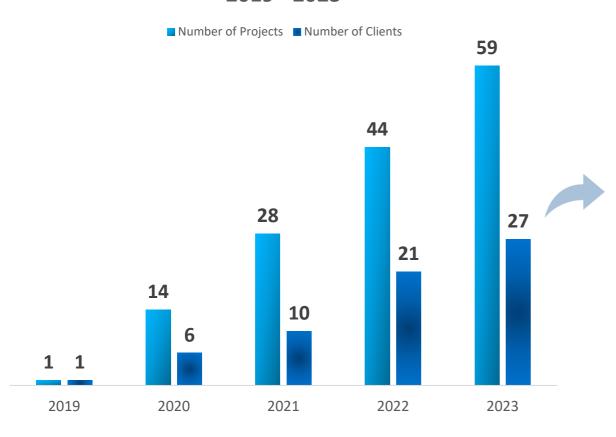
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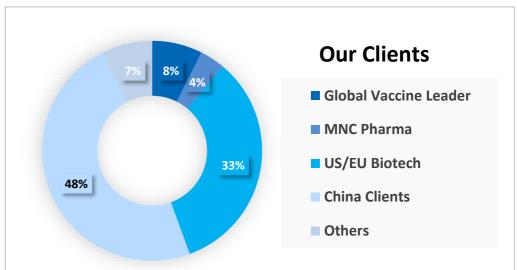
- Cumulative number since our inception and as of Dec 31, 2023
- 2. Number of ongoing integrated CMC projects, as of Dec 31, 2023
 - The small-sized figures account for the number of projects as at December 31, 2022, save for the number of projects at discovery stage which is cumulative from the Group's inception up until December 31, 2022

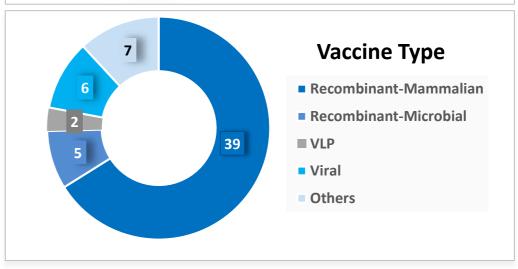
Progress of WuXi Vaccines Serving 27 Clients on 59 Projects



Cumulative Number of Projects and Clients 2019 - 2023





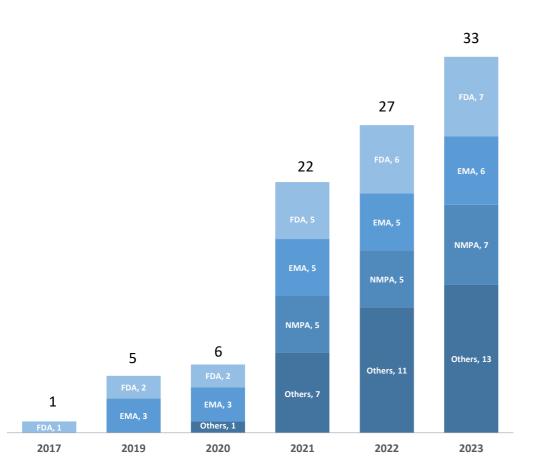


1. As of Dec 31, 2023

QUALITY is Our Competitive Advantage



Number of Regulatory Inspections Completed: 33(1)



Number of License Approvals: 48 (64 by Facility)

Ag	ency Facility	MFG1	MFG2F	MFG2P	MFG4	MFG5	DP1	DP4	MFG3(CB)	MFG3
1	FDA (6)	WBP XXX WBP XXX WBP XXX	WBP XXX	WBP XXX		WBP XXX/ WBP XXX	WBP XXX WBP XXX WBP XXX			
2	EMA (7)	WBP XXX WBP XXX	WBP XXX	WBP XXX	WBP XXX	WBP XXX/ WBP XXX	WBP XXX		WBP XXX	
3	NMPA (6)	WBP XXX WBP XXX		WBP XXX(2) WBP XXX			WBP XXX WBP XXX(2) WBP XXX	WBP XXX WBP XXX		WBP XXX
4	ANVISA (3)		WBP XXX WBP XXX		WBP XXX					
5	WHO (1)				WBP XXX					
6	TGA (2)		WBP XXX WBP XXX							
7	Hong Kong (1)		WBP XXX							
8	MHRA (4)	WBP XXX WBP XXX	WBP XXX	WBP XXX			WBP XXX			
9	PMDA (2)		WBP XXX							
10	Switzerland (2)		WBP XXX							
11	United Arab Emirates (2)		WBP XXX							
12	Canada (3)		WBP XXX WBP XXX			MDD MM/				
13	Saudi Arabia (1)					WBP XXX/ WBP XXX				
14	Costa Rica (1)					(14)				
15	Panama (1)									
16 17	Russian (1)									
18	New Zealand (1) Thailand (1)									
19	Jordan (1)									
20	Malaysia (1)									
21	HSA (1)									
#	Approvals 64	10	15	6	3	± 16	10	2	1	NDD A





























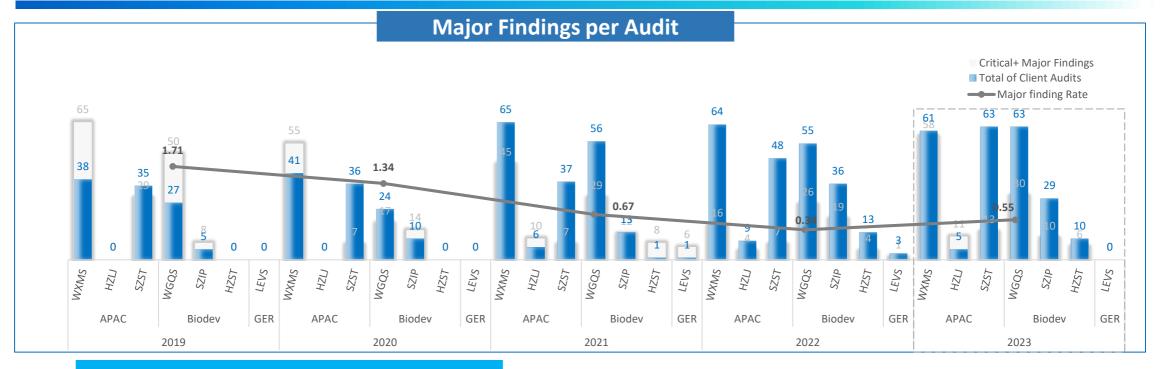






Leading Indicators of Quality/Regulatory Inspections: Continue to See Favorable Trends During Client Audits





Summary

While others struggle with data integrity issues, WuXi Biologics continues to safeguard our data integrity to maintain our strong track record with customers and regulators: 0 issue with data integrity during regulatory inspections

Global Operations on track to Support 02
Business Growth

Robust Global Network to Enable Partners: Multiple Nodes with Geographic Diversity



Global CRDMO: 3 R centers + 8 D centers + 9 M centers

R: Shanghai WGQ, Shanghai FX, Boston

D: Shanghai WGQ, Wuxi, Shanghai FX, Chengdu, Hangzhou, Suzhou, Cranbury NJ and Singapore

M: Wuxi, Hebei, Chengdu, Hangzhou, Wuppertal, Leverkusen, Dundalk, Worcester MA and Singapore



Groundbreaking on CRDMO Center in Singapore





The Company has broken ground on its new 13.5-hectare CRDMO center in Singapore. The center will provide integrated biologics research, development and manufacturing services



The CRDMO site will add 120,000L manufacturing capacity to WuXi Biologics' global network



The site is expected to employ 1,500 people when fully operational





New Chapter for Global Operations





- MFG6/7 GMP released in Q4 2022, and received first GMP Certificate from Ireland
- 7 CMO contracts signed and almost fully booked in 2025
- 3 ENG run completed with 100% success rate, completed first 16,000L manufacturing run, 3 PPQ batches in progress and 5 PPQ expected in 2024

MFG6/7 in Dundalk, Ireland



- MFG19 will further increase to 12x2,000L
- DP7 (liquid/lyo commercial facility, Germany/EMA certified): with an annual capacity of approximately ten million doses, is being upgraded to include a second variable filling line in 2025



- MFG18, the first clinical manufacturing facility in U.S., started GMP operations in August, 2022
- Produced 10+ GMP batches and achieved 100% success rate
- GMP slots for 2024 are 70% booked



- In response to global clients' increasing demands, MFG11 will increase its capacity from original planned 24,000L to 36,000L
- Anticipates being operational in 2025 with GMP release in 2026



- A 48,000L commercial DS manufacturing facility GMP released in Oct, 2022
- Completed 6 batches with 100% success rate in 2023
- Expect 1st PPQ completed in 2024

MFG19/DP7 in Germany

MFG18 in New Jersey, U.S.

MFG11 in Massachusetts, U.S.

MFG8 in Hebei, China

Ireland Ramp-up Ahead of Schedule: Expect to Break Even in 2024







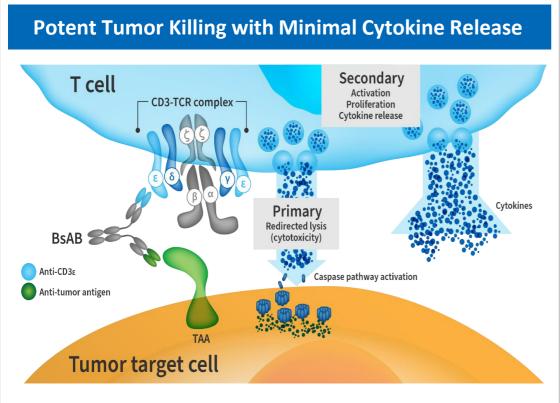
- Ireland site completed construction with record speed and received the 2023 Facility of the Year Award (FOYA) in the Operations category from ISPE
- GMP runs at both facilities are proceeding as expected: the first PPQ batch was successfully completed at MFG6 and successfully completed first 16,000L manufacturing run at MFG7
- Significant commercial manufacturing demands from 2024 onwards, almost fully booked in 2025
- Expect to break even in 2024, one year earlier than the original target. This is a strong testament of our continued business momentum as we undergo capacity increase outside of China
- 70% of the pipeline are from "Win-the-Molecule" execution, most of which are commercialized blockbuster products on the market
- Fast ramp-up in Ireland can be replicated to other global sites

State-of-the-art
Technologies Lead to
Future Success

03

WuXi Bio's Proprietary Clinical Stage Anti-CD3 mAb Empowers Discovery of Best-in-Class T Cell Engagers (TCEs)





Hypothesis – Adjusting CD3 Affinity & Binding Kinetics to Disassociate Tumor Cytotoxicity from Cytokine Release

WuXi CD3 Platform Highlights

- 3 TCEs in clinical stage (Ph1 clinical trials)
- Cynomolgus monkey cross-reactive CD3 binder
- Unique epitope with intermediate affinity
- Fast-on & fast-off binding kinetics
- Low cytokine release with potent tumor killing
- Several partnered with leading pharmaceutical companies

Full Kinetic Affinity to Human CD3εδ	ka (1/Ms)	kd (1/s)	K _D (nM)
CD3xCD20 (Regeneron)	2.21E+04	7.39E-05	<mark>3.4</mark>
CD3xDLL3 (Amgen)	4.60E+05	2.22E-03	<mark>4.8</mark>
CD3xDLL3 (Harpoon)	6.01E+06	4.03E-03	<mark>6.7</mark>
CD3xDLL3 (BI)	5.28E+05	<mark>6.23E-03</mark>	12
CD3xPSMA (Amgen)	1.45E+05	2.25E-03	16
CD3xBCMA (Celgene/Engmab)	8.64E+04	3.12E-03	36
CD3xGPC3 (Chugai)	3.98E+04	1.69E-03	43
CD3xCEACAM5 (Roche/Chugai)	7.94E+04	3.95E-03	50
CD3xFolR1(Roche)	2.25E+05	2.00E-03	89
WuXi Bio Lead	4.61E+05	1.86E-02	40

WuXiUI[™] Performance and Strategic Applications to Biologics Commercial Manufacturing

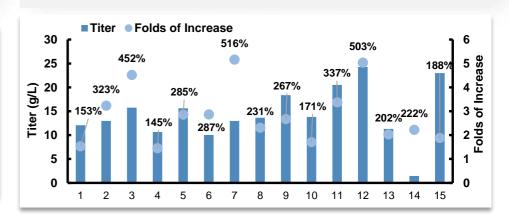




WuXiUI™ Ultra-Intensified FB Bioprocessing Strategy

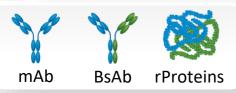
- 3~6 folds of TFB productivity with WuXiUI™
- 10~35 g/L upstream titer for popular CHO cell lines
- Implementing ESG concept in process design

Productivity Comparison WuXiUITM vs. TFB

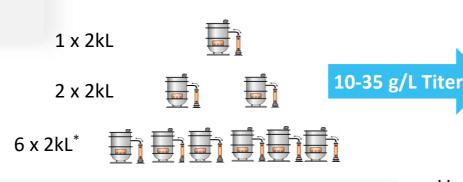




Implementation Strategy

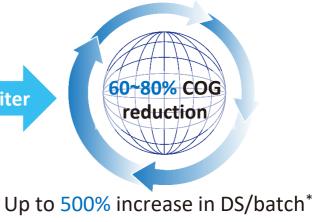


Suitable for multiple common host cell lines such as CHOK1/CHO-S/CHO-M/CHO GS



Existing facility for ease of implementation

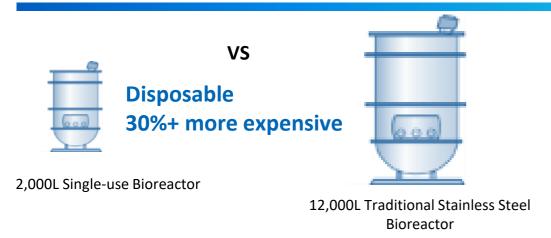
*Minor upgrade to adapt the facility

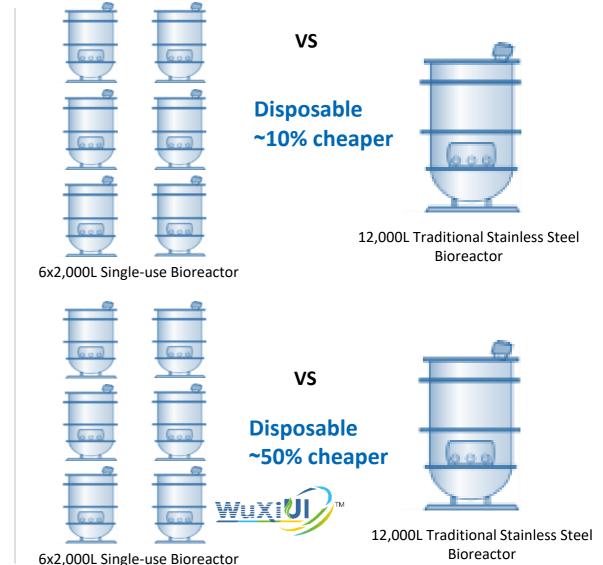


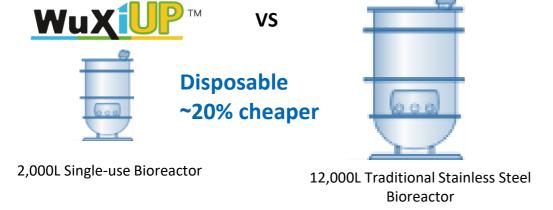
*Assuming no scale change

Disposable Bioreactors CAN Deliver Lower COGS than Traditional Stainless Steel









Global MNC Applies Single-use Technology at its Ireland Site Another Demonstration for This Manufacturing Technology



Eli Lilly picks ABEC for single-use Ireland facility

by Millie Nelson Thursday, July 13, 2023 5:33 am

ABEC will deliver multiple Customer Single Run (CSR) single-use systems to Eli Lilly's manufacturing facility in Limerick, Ireland.

Eli Lilly announced plans to build a €400+ million (\$433 million) greenfield site in Limerick, Ireland <u>last year</u>. The firm then magnified the investment to around \$1 billion to further support its biologics ambition in the country. in March.

Now, Lilly has selected ABEC, a solutions and services for biotech manufacturing to support Basis of Design (BOD) engineering of the plant's upstream and downstream processes.

ABEC will implement its single-use technology and its large-scale CSR bioreactors will be "at the heart of the facility," used for Lilly's high-density culture processes.



WuXi Bio's Track Record

WuXi Bio delivered ~4 tons of COVID-19 neutralizing antibodies with single-use technology, demonstrating that such technology is comparable to stainless steel in large-scale manufacturing



Big Pharma's Choice

One big pharma plans to expand its facility in Ireland with single-use technology, showcasing the strength and trend of such manufacturing technology



Single-Use in Capacity Expansion*

Growing trend: 44% single-use technology is applied in new capacity; 36% single-use technology is applied in CMO capacity

* Source: Morgan Stanley report

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Thriving Growth in Microbial Business



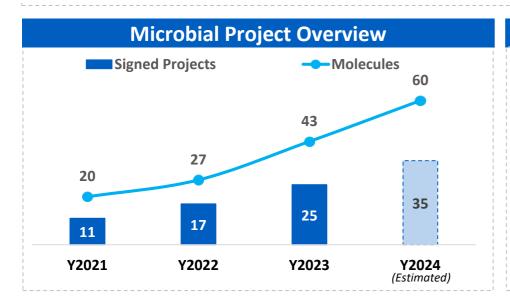
Regulatory

Filling

Strain Development Development

QC Release

and Stability

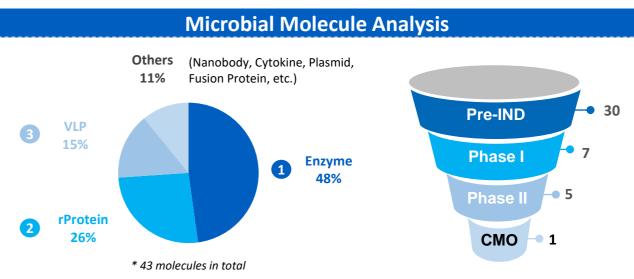


Formulation

Development

DP Process

Development



Process

Characterization

DS Process

Development

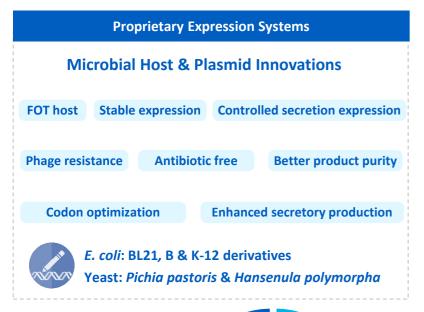
Strong "D" and Expansion for "M" in Microbial Business

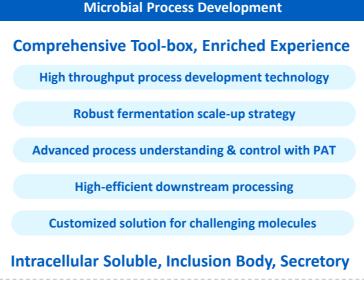




Strong development capabilities









Plan for Large-scale Microbial Production





CMC

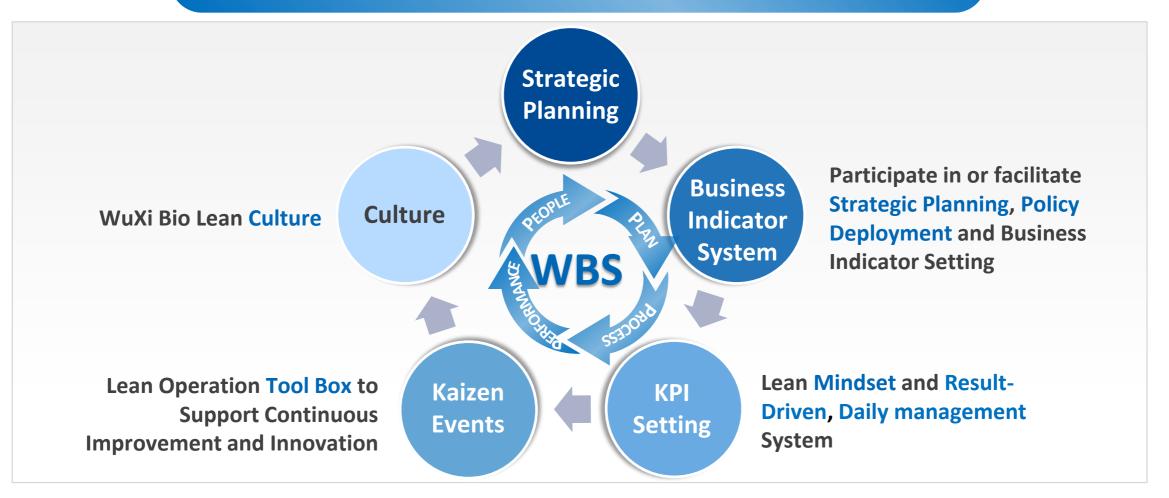
WBS and ESG as Important Components 04 of Business Strategy

WuXi Biologics Business System (WBS) launched in 2021



WuXi Biologics' Lean Operation and Management System

HIGHER QUALITY, FASTER DELIVERY, BETTER VALUE



WBS Achievements in 2023





376 kaizen projects are completed in 2023

Improvements





- Reduction of QC assay failure rate, ↓ 28%
- On-time completion rate of Deviation increasing 6%



Material Cost Saving

- Materials cost saving for one project in MFG, ↓ 13.7mm
- Cost saving for non-GMP materials in one site, ↓ 5.9mm
- Material cost saving in PD, ↓
 5.2mm



Labor Efficiency Improvement

- Working hours reduction for MFG downstream, ↓
 16,000+ hr/yr
- Re-validation strategy
 optimization for SZ site , ↓

 7,000+ hr/yr



ESG

- Low-energy model establishment for HBST facility, ↓ 5.8mm/yr
- Standardize operation of membrane pack, ↓ 3.2mm/yr

ESG Performance Recognized as Industry Leader





Dow Jones Sustainability World Index
Dow Jones Sustainability Emerging Markets Index

- Ranked No.1 in its sector and identified as a global sustainability leader
- Recognition in talent development, innovation, quality, environment and climate change, etc.
- Inclusion into Global Sustainability Yearbook, generating long-term value for stakeholders



MSCI AAA ESG Rating of 2023

- The highest rating on MSCI's AAA (Leaders) to CCC (Laggards)
- A company leading its industry in managing the most significant ESG opportunities
- MSCI is a major ESG data and rating provider, covering 10,000+ companies

lote:

ESG Performance Recognized as Industry Leader





EcoVadis Platinum Medal

- Ranked top 1% of 100,000+ companies
- Outstanding performance across all areas:
 Environment, Labor & Human Rights, Ethics,
 and Sustainable Procurement
- A trusted partner by global clients for its strong sustainability commitment

This is really fantastic news!
So excited to hear WuXi Bio is
doing an excellent job with
EcoVadis."

- Feedback from one of Top Pharma clients



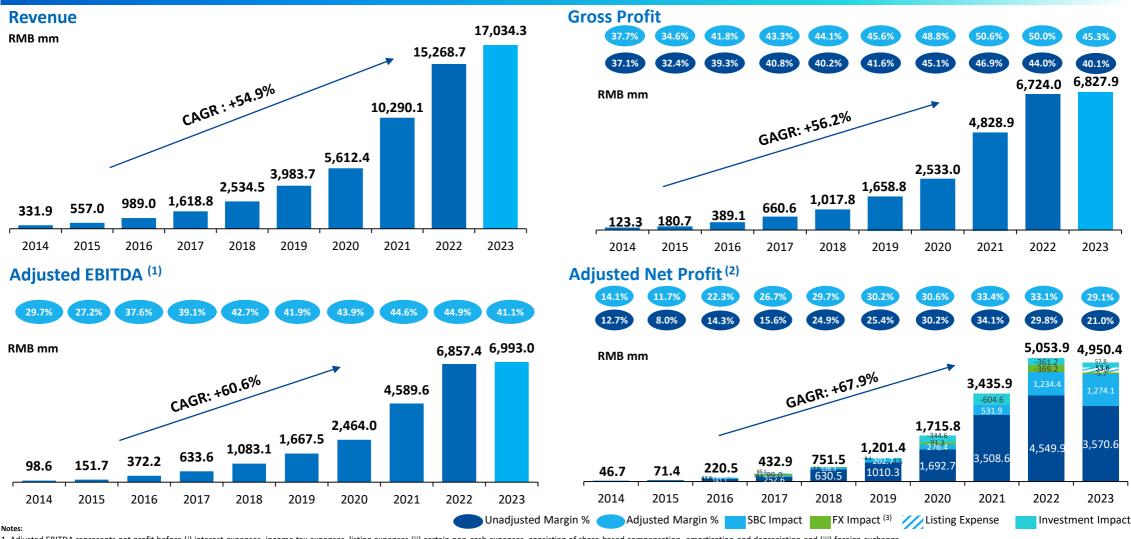




Financial Overview 05

Solid Financial Performance Achieved





^{1.} Adjusted EBITDA represents net profit before (i) interest expenses, income tax expenses, listing expenses (ii) certain non-cash expenses, consisting of share-based compensation, amortization and depreciation and (iii) foreign exchange gains/losses and (iv) fair value gains/losses on investment portfolios

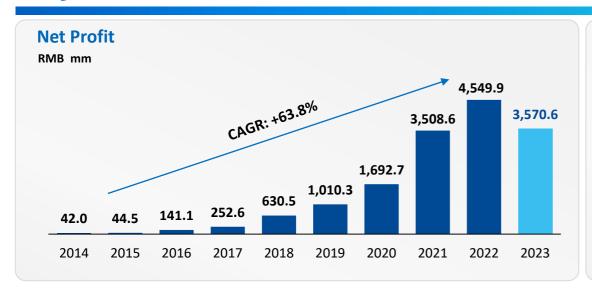
^{2.} Adjusted net profit excludes the share-based compensation expenses, fair value gains/losses on investment portfolios, foreign exchange gains/losses and listing expenses

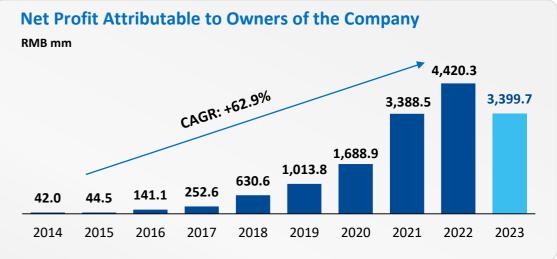
^{3.} Refers to foreign exchange gains/losses

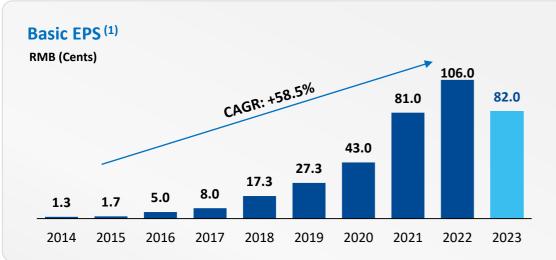
^{4.} Adjusted EBITDA and adjusted net profit of 2019 have been restated to further exclude the fair value gains/losses on the Group's investment portfolios

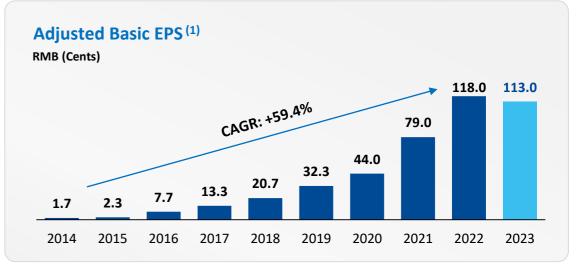
Key Profit Metrics











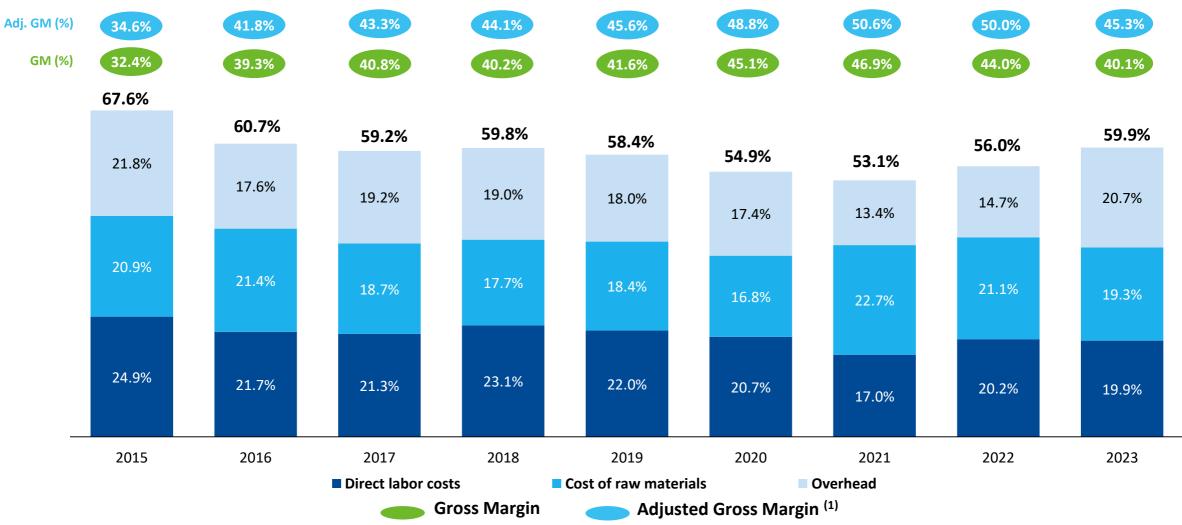
Note:

1. The authorized and issued shares of the Company were subdivided on the basis that every one (1) issued share is subdivided into three (3) subdivided shares (the "Share Subdivision"), which became effective on November 16, 2020. Basic and diluted earnings per share were stated after taking into account the effect of the Share Subdivision. Comparative figures have also been restated on the assumption that the Share Subdivision had been effective in the prior year

Gross Profit and Cost of Sales

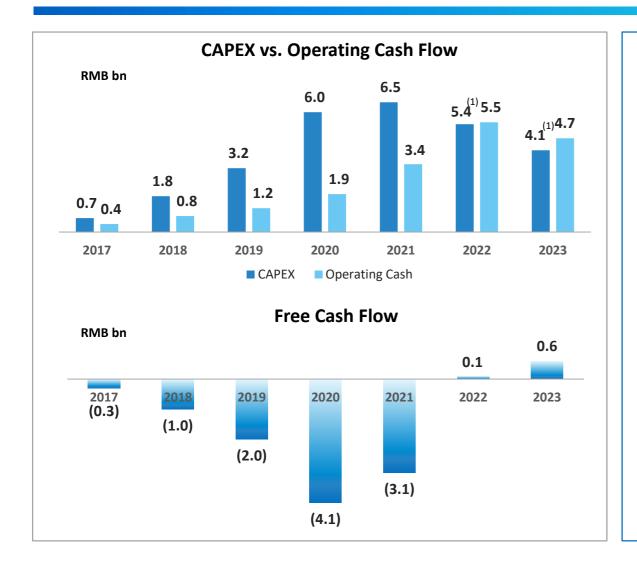


Cost of Services as % of Revenue



Free Cash Flow Positive since 2022





- Net operating cash flow recorded ~51%
 CAGR growth from 2017 to 2023
- Free cash flow turned positive in 2022 and continued to improve in 2023: critical milestone for company growth
- Target at continued free cash flow positive in
 2024 and beyond
- Solid cash flow management to support global capacity and capabilities enhancement

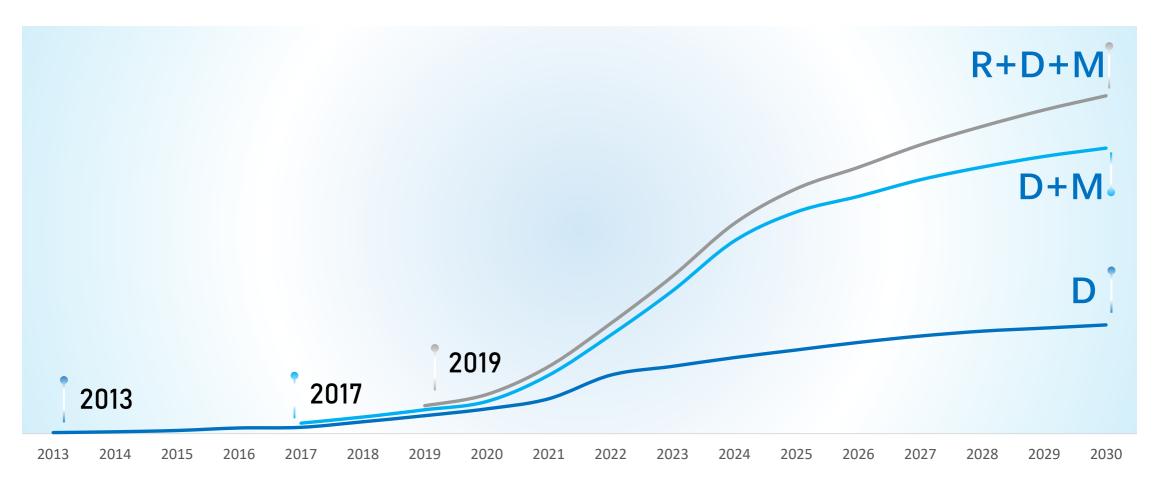
1. CAPEX excluded of asset acquisitions via M&A deals

06 **Summary**

CRDMO: Three Growth Curves Drive Sustainable Long-term Growth



Three Long-Term Growth Curves



Proven CRDMO Business Model Continues to Deliver Sustainable Growth



Contract Services

- **Target Selection**
- Reagent/Protein Generation & Assay Development
- **Antibody Generation**
- Complex Biologics Engineering (e.g., ADCs, bsAb)
- Lead ID & Optimization
- **Developability Assessment**
- Characterization: (e.g., PK, PD, Efficacy, & Exploratory Tox
- **PCC Selection**
- **Regulatory Support**

- Cell Line Engineering
- **Assay Development**
- Process Development
- **Drug Product Development**
- DS & DP Scale Up
- Cell Banking & Characterization
- Pilot Scale Manufacturing
- Viral Clearance
- Late-stage Development & PC/PV
- **IND & BLA Filing Support**

DS GMP Manufacturing

- **DP GMP Manufacturing**
- QC Lot Release & Stability
- Global Dual Source Support

One partner expertise in

all areas

Requisites

State-of-the-art technology platforms enhance drug discovery capabilities

Best-in-class technology platforms, maximum scalability, speed and execution

Large CAPEX, validated quality and unwavering execution

Current State

- One of the best mAb platforms demonstrated by Arcus deal in 2018
- One of the best bispecific platforms demonstrated by GSK collaboration

132 new integrated projects in 2023

Robust growth of non-COVID CMO from 1 in 2019 to 16 in 2023

Achievements

US\$7.2 bn milestone backlog plus 55+ programs with low single digit royalties

- ~US\$1 bn revenue per year from new projects only
- ~700 programs

- Strong growth potential benefiting from feed from R&D
- "Win-the-Molecule" expedites M

Conversion Rate: 95%

Conversion Rate: 90%

2023 Highlights: Significant Progress in R, D and M, Best in Company Record





Most important reason for the 2023 slow-down is 84% decrease of COVID revenue.

Overall non-COVID revenue grew 37.7% YoY in 2023. R&D is still core to the Company with high profitability and customer stickiness



R: record high revenue, 110% 3-year CAGR growth for milestone and research services revenue, blockbuster deals signed with global pharma



D: We saw significantly fewer projects in Q1 2023 due to biotech funding challenges, but great recovery to full year record high of 132 new projects added. New technology platform progressing well: WuXiUITM and Microbial. Continue to gain market share despite all challenges



M: non-COVID revenue from late phase and CMO stage grew 101.7% YoY. Ireland site ramp-up ahead of schedule

2024 Outlook





R&D: Continue to expect strong growth. Strong momentum from global biotech recovery in Q4 2023 continued in Q1 2024. 25 new projects signed as of Mar 22, 2024 vs 8 in Q1 2023



M: "Follow the Molecule" projects track well. Several potential blockbuster projects move to PPQ stages which will generate significant revenue in 2025 and beyond. "Win-the-Molecule" projects will be potentially more challenging due to geopolitical concerns. As a result, we will continue to invest in Germany, US and Singapore for commercial MFG capacities



Due to uncertainties in the external environment, 2024 revenue is expected to achieve 5%-10% growth, non-COVID revenue is expected to grow 8%-14%



Business fundamentals remain strong







2023	2022	Change	
17,034.3	15,268.7	11.6%	
(10,206.4)	(8,544.6)		
6,827.9	6,724.0	1.5%	
416.7	305.5		
(320.0)	(258.5)		
36.5	766.5		
(294.0)	(162.9)		
(1,495.4)	(1,269.6)		
(53.6)	-		
(785.8)	(682.8)		
(158.5)	(64.4)		
4,173.8	5,357.8	-22.1%	
(603.2)	(807.9)		
3,570.6	4,549.9	-21.5%	
0.82	1.06		
1.13	1.18		
	17,034.3 (10,206.4) 6,827.9 416.7 (320.0) 36.5 (294.0) (1,495.4) (53.6) (785.8) (158.5) 4,173.8 (603.2) 3,570.6 0.82	17,034.3 15,268.7 (10,206.4) (8,544.6) 6,827.9 6,724.0 416.7 305.5 (320.0) (258.5) 36.5 766.5 (294.0) (162.9) (1,495.4) (1,269.6) (53.6) - (785.8) (682.8) (158.5) (64.4) 4,173.8 5,357.8 (603.2) (807.9) 3,570.6 4,549.9 0.82 1.06	

Notes:
1. Results may not foot due to rounding

Reconciliation for Adjusted Net Profit and Adjusted EBITDA



(RMB million)	2023	2022	Change
Adjusted Net Profit Reconciliation			
Net Profit	3,570.6	4,549.9	
Share-based Compensation Expense	1,274.1	1,234.4	
Foreign Exchange Gain	(5.7)	(369.2)	
Losses (Gains) from Equity Investments	57.8	(361.2)	
Listing Expenses	53.6	-	
Adjusted Net Profit	4,950.4	5,053.9	-2.0%

Adjusted EBITDA Reconciliation			
EBITDA	5,613.2	6,353.4	
Share-based Compensation Expense	1,274.1	1,234.4	
Foreign Exchange Gain	(5.7)	(369.2)	
Losses (Gains) from Equity Investments	57.8	(361.2)	
Listing Expenses	53.6	-	
Adjusted EBITDA	6,993.0	6,857.4	2.0%

Notes:

1. Results may not foot due to rounding

WuXi Bio Vision

"Every drug can be made and every disease can be treated" by building an open-access platform with the most comprehensive capabilities and technologies in the global biologics industry

