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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

PRESENTATION AT 2024 ANNUAL HEALTHCARE CONFERENCE

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).

The Company will attend the 42nd Annual J.P. Morgan Healthcare Conference ("Annual Healthcare Conference") on January 11, 2024 (Hong Kong time) at which it will conduct a presentation containing certain information regarding the Company's business operations (the "Presentation"). To ensure that all shareholders and potential investors of the Company have equal and timely access to such information, the Company has included in this announcement the full copy of the Presentation. Shareholders and potential investors of the Company 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.

As disclosed in the Presentation, the Company's business and financial positions have remained strong with 132 newly added projects resulting in a total of 698 projects (including 9 phase III and commercial "Win-the-Molecule" projects and 24 CMO projects) in 2023. These projects validate the customers' trust and recognition in the Company's CMO capabilities and capacity, and the increase in the number of CMO projects will further boost the Company's near term revenue. Through the "follow the molecule" strategy and enhanced capabilities to build multiple technology platforms to enable different modalities, the Company is confident that it will continue to achieve a solid growth in 2024 and maintains a positive outlook for the future.

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, January 10, 2024

As at the date of this announcement, the Board 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 High Growth

Chris Chen PhD

42nd Annual J.P. Morgan Healthcare Conference





Stock Code: 2269.HK

Forward-Looking Statements



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 EBITDA, adjusted EBITDA margin and adjusted diluted 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





Business Update

Top Global Bioprocessing Trends in 2023: Betting on the Right Trends



CMO on the Rise

The growth in contract manufacturing facilities means that biopharma manufacturers have more flexible options for their manufacturing strategies, including outsourcing their product manufacturing. Big pharma tend to outsource more, including R, D and M

Single-use technology

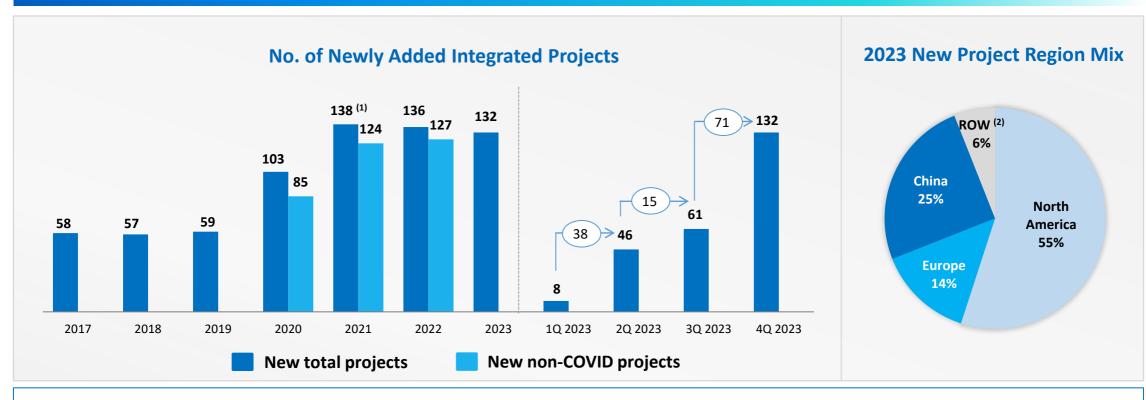
Single-use technologies
are accelerating the
replacement of stainless
steel. The bigger
deployment of singleuse technology
continues

Offshoring

Globalization of
biomanufacturing and
offshoring are growing,
especially in major
markets and Asia

New Projects Rebound from the Bottom in Q1 2023: 2023 is Now Record High of New Non-COVID Projects





- Number of new projects impacted in Q1 2023 by global biotech slowdown and rebounded since Q2 2023. Strong recovery in 2H 2023 with 2023 now highest No. of non-COVID projects, indicating more brand recognition and trust from the industry
- North America accounted for 55% of new projects and China rebounded significantly to 25%
- Strong growth in drug development with more projects inflow, raise 2024 project No. guidance from 80 to 110

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 and Deliver Additional Near-term Growth



2018

2019

2020

2021

& CMO

2022

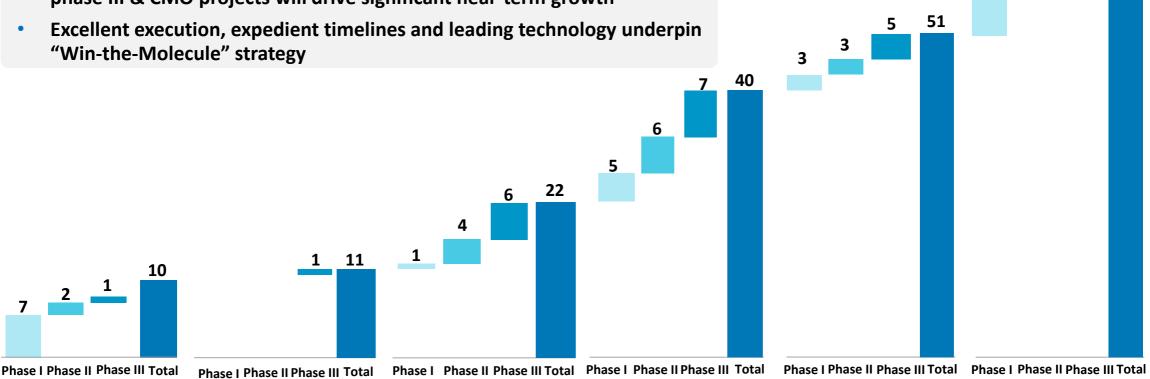
& CMO

2023

& CMO



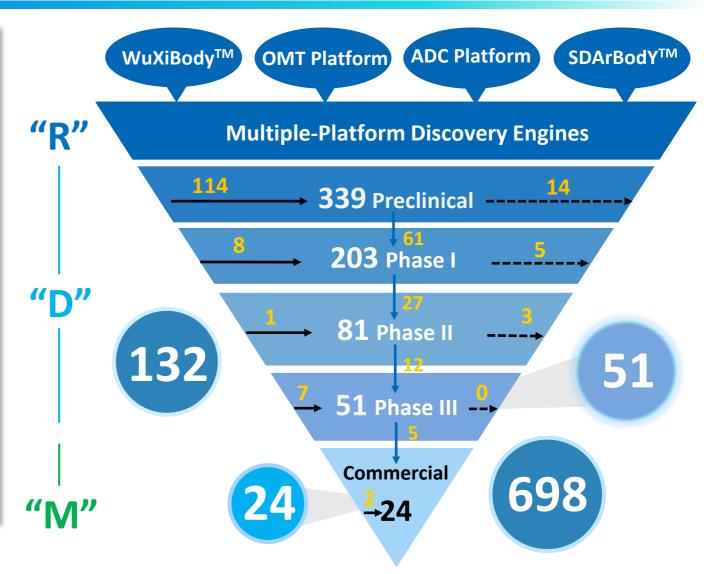
 Total 69 projects at different stages (Phase I, II and III + CMO) transferred from global CDMOs or large pharmas to WuXi Biologics since 2018: 29 phase III & CMO projects will drive significant near-term growth



New Projects Exceeded Expectations: Demonstrate Business Resilience Despite Headwinds



- Continue to gain market share for both "Follow and Win the Molecule"
- Despite headwinds, newly-added projects reached all time high excluding COVID, demonstrating the Company's resilience during the downturn cycle and ability to gain market share
- "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
- 51 Phase III projects and 24 CMO projects: poised for future manufacturing revenue growth

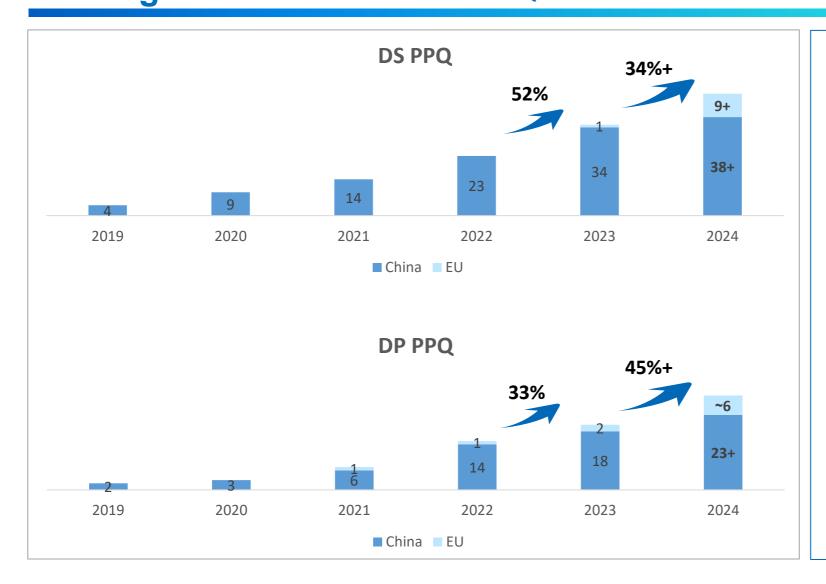


Notes:

^{1.} As of Dec 31, 2023

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

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
- 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 global competitiveness as we undergo capacity expansion 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

Explosive Growth of Commercial Projects in the Near Term



Seven manufacturing projects that could potentially generate US\$200 mm+ peak revenue per year

- Cancer bispecific A
- Cancer bispecific B
- Bispecific C
- FcRn mAb
- Autoimmune program K
- Cancer ADC Z
- Cancer ADC Y

Ten manufacturing projects that could potentially generate US\$100-200 mm peak revenue per year

- Pompe ERT
- Cancer mAb
- Cancer bispecific D
- Non-COVID Vaccine
- Global biosimilar 1
- Global biosimilar 2
- Global biosimilar 3
- Infectious disease mAb1
- TIGIT mAb
- Long-acting HGH

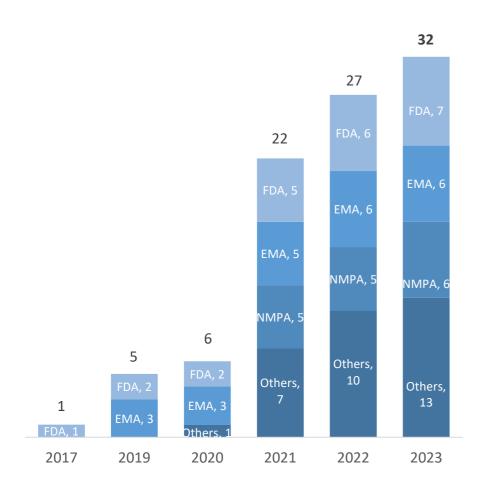
Eight manufacturing projects that could potentially generate US\$50-100 mm peak revenue per year

- Cancer bispecific E
- Cancer ADC X
- CD38 mAb
- DR5 mAb
- Global biosimilar 4
- Cancer ADC W
- Gaucher's disease ERT
- Infectious disease mAb 2
- Multiple programs and higher POS for exciting targets, signed 4 exclusive CMO deals (dual sourcing within WuXi Biologics)
- Inventory-built for biologics can start 2-3 years before approval due to complexity of manufacturing and long supply chain
- CMO revenue from these projects expected to be US\$2 bn+

QUALITY is Our Competitive Advantage







Number of License Approvals: 45 (60 by Facility) (2)

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			
2	EMA (7)	WBP XXX	WBP XXX	WBP XXX	WBP XXX	WBP XXX/ WBP XXX	WBP XXX WBP XXX		WBP XXX	
3	NMPA (5)	WBP XXX WBP XXX	WBP XXX	WBP XXX WBP XXX			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 (2)		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							
13	Saudi Arabia (1)					WBP XXX/				
14	Costa Rica (1)					WBP XXX				
15	Panama (1)									
16	Russian (1)									
17	New Zealand (1)									
18	Thailand (1)									
19	Jordan (1)									
20	Malaysia (1)									
21	HSA (1)									
#	Approvals 60	8	17	4	3	16	8	2	1	1

































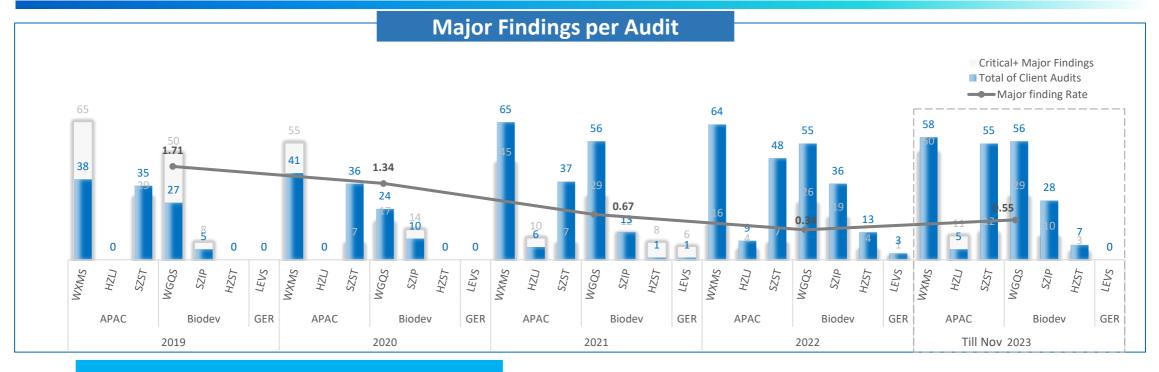






Leading Indicators of Quality/Regulatory Inspections: Continue to See Favorable Trends During Client Audits





Summary

209 Client Audits completed as of Nov 2023

0.55 (115/209) Major findings per Audit as of Nov 2023

1.71 -> 1.34 -> 0.67 -> 0.34 -> 0.55
(2019) (2020) (2021) (2022) (till Nov 2023)

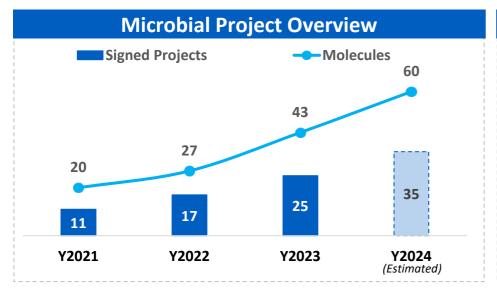
While others struggle with data integrity issues, WuXi Biologics continues to safeguard our data integrity to maintain our strong track record with customers and regulators: 0 issue with data integrity during regulatory inspections

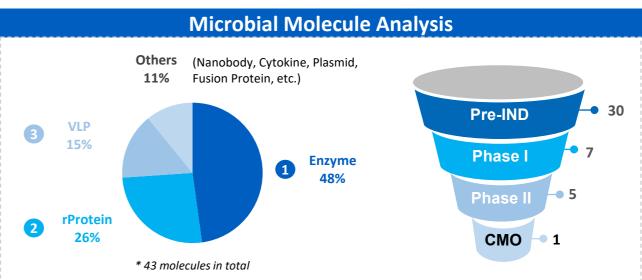
Microbial Business to be Another Growth Driver **02**

Thriving Growth in Microbial Business



End-to-end One-stop Shop from DNA to Regulatory Filling Analytical Process Strain Analytical DS GMP DP GMP Development Validation Manufacturing Validation Development Manufacturing DS Process DP Process Regulatory Formulation **QC** Release **Process Development Filling Development Development** and Stability Characterization





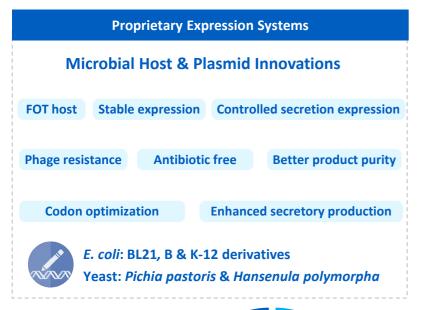
Strong "D" and Expansion for "M"

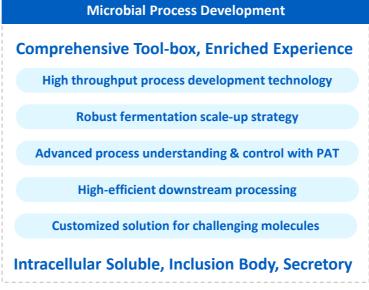




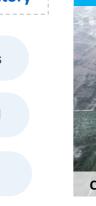
Strong development capabilities













Global **Quality System**

Technology

Depth





CMC Experience



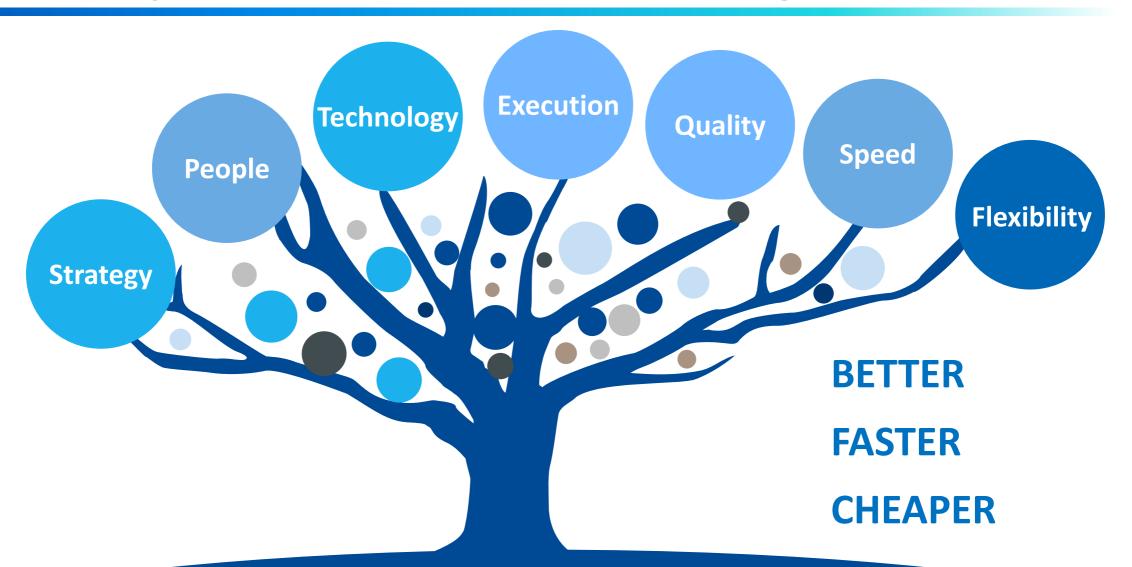




Seven Success Factors 03 also Proven on XDC

SEVEN Keys for Future Success: Sustainable High Growth





Global Leading ADC and Bioconjugate CRDMO



No.1 Globally

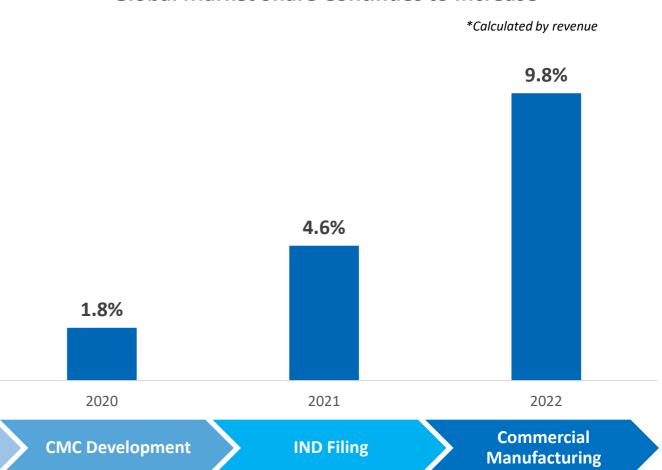
By integrated project number

No.2 Globally

By 2022 revenue

No.1 in China

By 2022 revenue



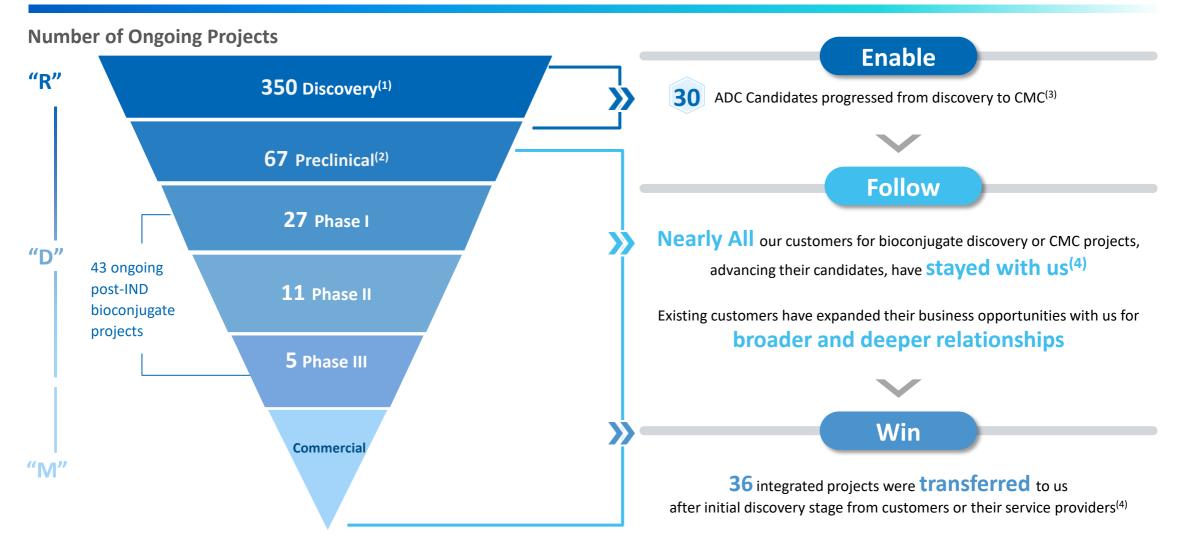
Global Market Share Continues to Increase

Antibody Discovery

Conjugation Research

"Enable-Follow-Win" Strategy Builds Robust Project Base



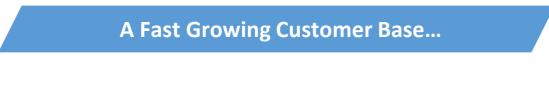


Notes

- 1. Cumulative number of discovery projects since our inception and as of Jun 30, 2023
- Number of ongoing integrated projects, as of Jun 30, 2023
- 3. Since inception in 2013 and up to Jun 30, 2023
- 4. As of Jun 30, 2023

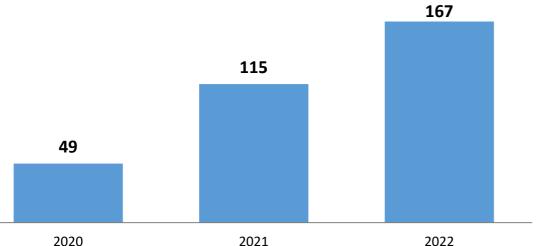
Seven Success Factors also Proven on XDC







XDC is the one and only pure-play ADC CRDMO player offering end-to-end services





Successful execution: powerful Platform Cutting Timeline by Half: From DNA to IND in < 15 months



State-of-the-art technologies underpin future growth and already gain the largest ADC/XDC pipeline globally



XDC supports >70% out-licensing deals Robust revenue growth: 1H 2023 comparable with full year of 2022

The Relations between WuXi Bio and WuXi XDC

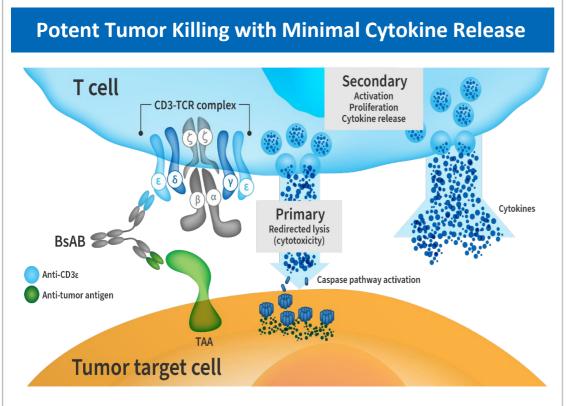
- WuXi XDC will continue to be consolidated by WuXi Biologics, and ~50% of its total earnings (as compared with 60% before the spin-off listing) will be accounted for the net profit attributable to owners of the company of WuXi Biologics
- No employee from WuXi Biologics (other than WuXi XDC) currently holds any stock option in WuXi XDC

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

04

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

WuXiUITM Performance and Strategic Applications to Biologics Commercial Manufacturing

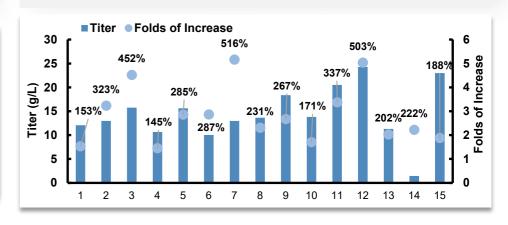




WuXiUI[™] Ultra-Intensified FB Bioprocessing Strategy

- 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 WuXiUI™ vs. TFB

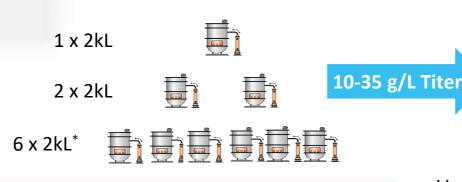




Implementation Strategy

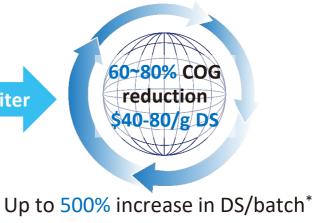


Suitable for multiple common host cell lines such as CHOK1/CHO-S/CHO-M/CHO GS



Existing facility for ease of implementation

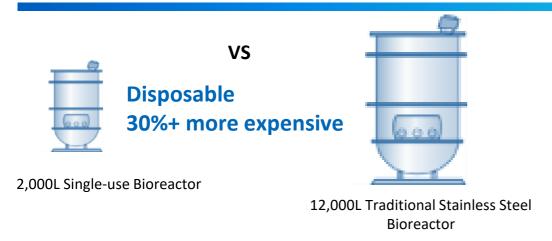
*Minor upgrade to adapt the facility

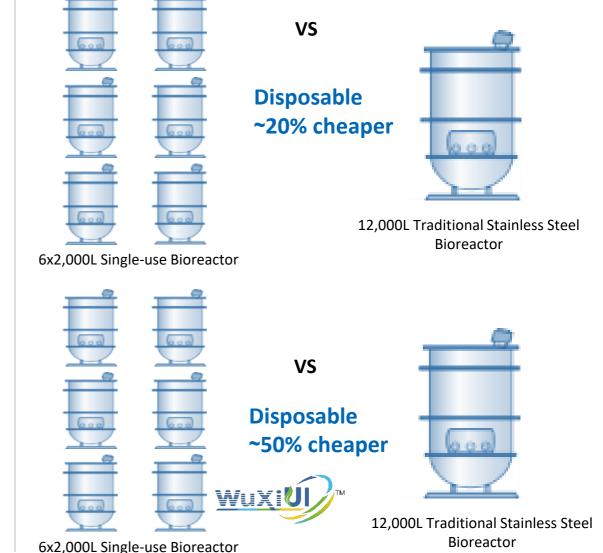


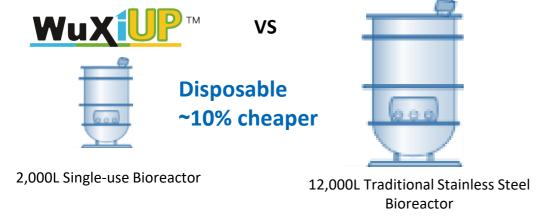
*Assuming no scale change

Disposable Bioreactors CAN Deliver Lower COGS than Traditional Stainless Steel









ESG as an Important
Component of 05
Business Strategy

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.
- Generating long-term value for stakeholders



MSCI AAA ESG Rating of 2023

- The highest rating on MSCI's AAA (Leaders) to CCC (Laggards)
- Defined as 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

ESG Performance Recognized as Industry Leader





EcoVadis Platinum Medal

- Ranked in 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







Integrated Strategy to Tackle Climate Change Advocate Energy Saving Initiatives Across Global Sites





Five Approaches



Carbon Targets



Efforts & Progress



Committed to SBTi

- Measure
- Avoid
- Reduce
- Substitute
- Offset



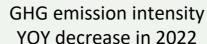
Mid-term: Reduce GHG emissions intensity by 50% by 2030

50% ↓

Net-Zero

Long-term: Net-Zero for overall operations by 2050





27% ↓

GHG emission intensity decrease since 2020



Energy Saving Across Global Sites



8,912 tCO₂e of carbon reduction



8.5 million+ kWh of electricity savings



6,550 tonnes of steam savings



200,000+ m3 of natural gas savings

Green Sites of the Future



MFG 6/7 in Ireland 100% renewable electricity supply



MFG8 in Hebei
Digital platform for energy optimization

Our Commitment to A More Sustainable Business



ISO

SBTi

Committed to SBTi in 2023

A new milestone towards net-zero

UNGC

A Signatory of UNGC in 2023

Business as a force for good



ISO 27001

Information Security Management

ISO 45001

Occupational Health and Safety

ISO 14001

Environmental Management

ISO 14064

GHG Emissions Verification

TCFD

A TCFD Supporter in 2023

Taking actions to build a more resilient financial system through climate-related disclosure

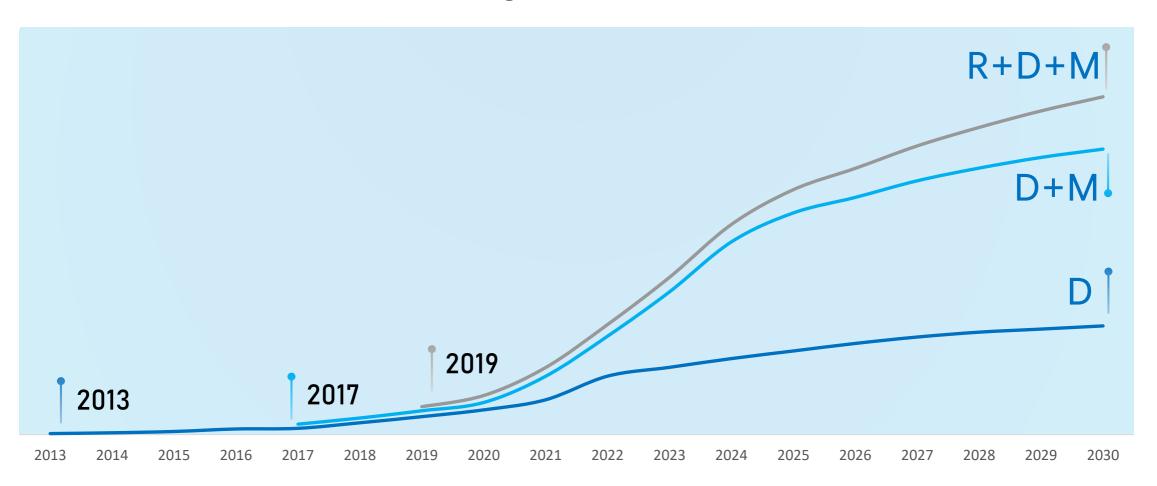
30

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



 $C \rightarrow R$

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 with 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

110 new integrated projects in 2024

Explosive growth from 1 in 2019 to 24 in 2023 to 32+ expected in 2025

Achievements

US\$6.5 bn+ milestone backlog and 50+ programs with low single digit royalties

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

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

Conversion Rate: 95%

Conversion Rate: 90%

Outlook & Summary



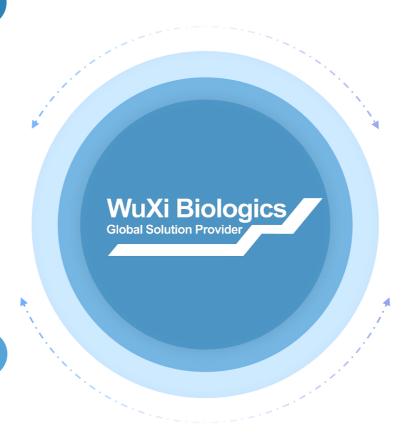
Highlights in 2023

Biotech funding slowdown and CMO revenue deferral made 2023 a challenging year, though the temporary impacts did not affect the Company's fundamentals.

Record high of 132 new projects added, 9 phase III and commercial "Win-the-Molecule" projects, 24 CMO projects. Ireland site ramp-up ahead of schedule

CMO Drives Future Growth

Remarkable growth upside expected to be driven by more CMO projects and several potential blockbusters already in pipeline



Solid Growth in 2024 and beyond

Continue to gain market share in R, D and M, especially in "D" through follow the molecule strategy. Expect a positive turnaround in Q2 2024 and maintain a positive outlook for the future

Multiple Businesses Support Additional Growth

Enhanced capabilities to build multiple technology platforms to enable different modalities and support additional growth, including ADC, bispecific, multispecific, microbial, vaccine and etc.

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

