



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, our ability to protect our clients' intellectual property, unforeseeable international tension, competition, the impact of emergencies and other force majeure. 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. All information provided in this presentation is as of the date of this presentation and are based on assumptions that we believe to be reasonable as of this date, and we do not undertake any obligation to update any forward-looking statement, except as required under applicable law.

Non-IFRS Financial Measures

We provide non-IFRS gross profit and non-IFRS net profit attributable to the owners of the Company, which exclude share-based compensation expenses, issuance expenses of convertible bonds, fair value gain or loss from derivative component of convertible bonds, foreign exchange-related gains or losses and amortization of intangible assets acquired in business combinations, non-financial assets impairment, etc. We also provide adjusted non-IFRS net profit attributable to the owners of the Company and earnings per share, which further exclude realized and unrealized gains or losses from our venture capital investments and joint ventures. Neither of above is 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 our core 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, non-recurring, non-cash and non-operating items that we do not consider indicative of the performance of our core business. Such non-IFRS financial measures, the management of the Company believes, is widely accepted and adopted in the industry the Company is operating in. However, the presentation of these adjusted 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.

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01 Results Overview

O2 Segment Performance

Financial Performance

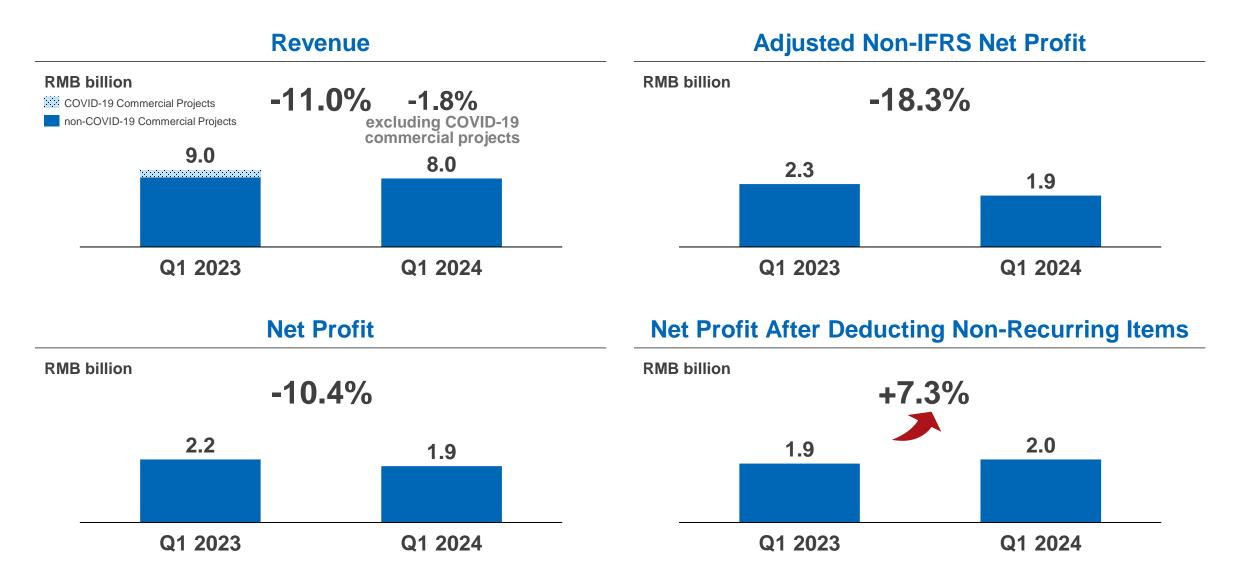
04 Business Outlook



1. Results Overview



Q1 2024 Target Achieved Despite External Challenges, Maintaining Stable Operations





Continue to Serve as a Highly-Efficient Enabler to the Industry

Strong, Loyal & Expanding Customer Base

- 300+ new customers, 6,000+ active customers
- 7.92Bn revenue from existing customers, -1.6% YoY excluding COVID-19 commercial projects
- 0.07Bn revenue from new customers

Execute Long-Tail
Strategy & Increase
Support to Large
Pharma

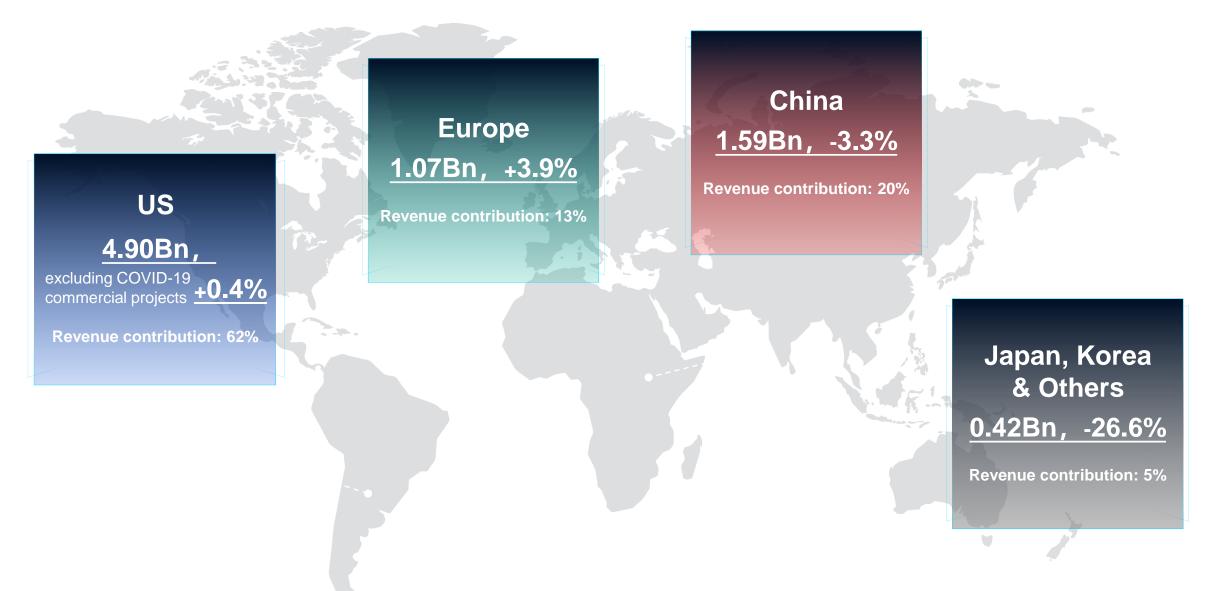
- 2.70Bn revenue from Top 20 global pharma clients, +4.2%↑ YoY excluding COVID-19 commercial projects
- 5.28Bn revenue from other customers, -4.6% YoY

Increase Customer Conversion to Sustain Growth

• 7.19Bn revenue from customers using services from multiple business units, representing over 90% of total revenue



Diversified Revenue Streams from Customers Across Regions Ensure the Stability and Resilience of the Company's Financial Performance





ESG Leading Position Further Enhanced in the Global ESG Rating Systems

Consecutively in 2023-2024: Named to the S&P Global Sustainability Yearbook, and Awarded as Industry and Regional Top-Rated Company by Sustainalytics









Consecutive "AA" ratings in 2021-2023





Consecutive "A-" rating in CDP Climate Change in 2022-2023, first "A-" rating in CDP Water Security in 2023



"Silver" medal by EcoVadis in 2023



WuXi AppTec Co., Ltd. Life Sciences Tools & Service

Sustainability Yearbook Member

S&P Global Corporate Sustainability Assessment (CSA) Score 2023











Powered by the S&P Global CSA









Unwavering Commitment to Customers' IP Protection and the Highest Quality & Compliance Standards



748¹

Quality Audits & Inspections by Global Customers, Regulatory Authorities and Independent Third Parties in 2023







83

Information Security
Audits by Global
Customers in 2023, with
O Critical Findings



24²

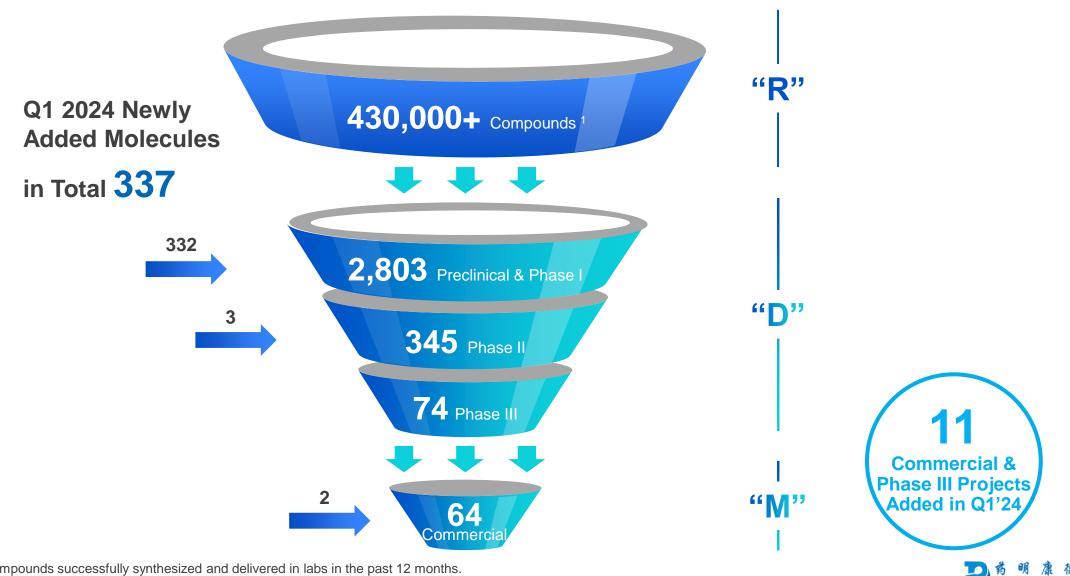
Main Operating Sites are ISO/IEC 27001 Certified



2. Segment Performance

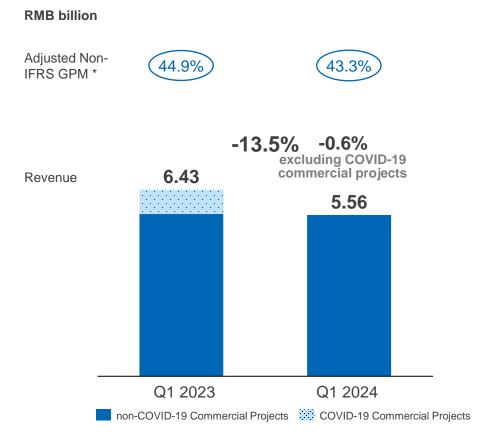


WuXi Chemistry: Growing CRDMO Pipeline Driven by "Follow the Molecule + Win the Molecule" Strategies



WuXi Chemistry: Integrated CRDMO Business Model Drives Steady Growth, with Continued Expansion in TIDES

Revenue & Gross Profit Margin



^{*} GPM @ CER: Q1 2023 44.5%, Q1 2024 40.6%

Drug Discovery (R) Generating Downstream Opportunities

- In the past 12 months, successfully synthesized and delivered
 430,000+ new compounds (grew 3% YoY)
- R to D&M conversion continued to improve efficiency, and molecules converted from R to D&M continued to grow

Continued Growth of Development and Manufacturing (D&M)

- Q1 D&M revenue¹ reached 4.00bn, excluding COVID-19 commercial projects, grew 1.2% YoY
- CDMO pipeline currently in total of 3,286 molecules (with 337 newly added in Q1); 11 commercial & Phase III projects added in Q1

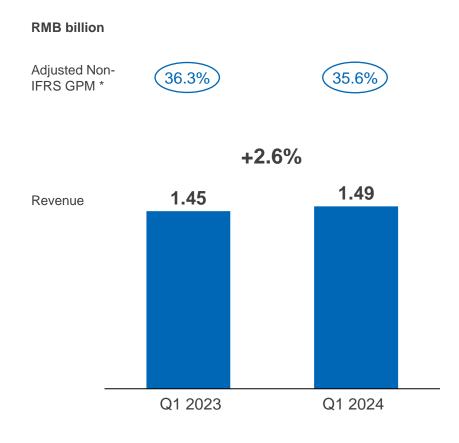
Continued Expansion of New Modalities (TIDES)

- Q1 TIDES revenue grew strongly by 43.1% YoY to 0.78bn. By end of Q1, TIDES backlog grew significantly by 110% YoY
- TIDES D&M customers grew 30% YoY to 146, and molecules grew 43% YoY to 276
- In January 2024, total reactor volume of Solid Phase Peptide Synthesizer increased to 32,000L



WuXi Testing: Drug Safety Evaluation Service & SMO Maintain Leadership Positions and Drive Steady Growth

Revenue & Gross Profit Margin



^{*} GPM @ CER: Q1 2023 36.2%, Q1 2024 34.5%

Lab Testing Services

- Q1 revenue of lab testing services down **0.6%** YoY to **1.05bn**, among which, drug safety evaluation service revenue grew **3.7%** YoY, maintaining **industry leading** position in APAC
- In Q1, Chengdu facility received NMPA¹ GLP qualification; Qidong facility received OECD² and NMPA GLP qualifications in March and April. Additional 22,000m² of facilities were GLP-qualified in Q1
- New modality business continued to develop, while new vaccine capabilities continued to improve, and market share of nucleic acids, conjugates, and mRNA further expanded

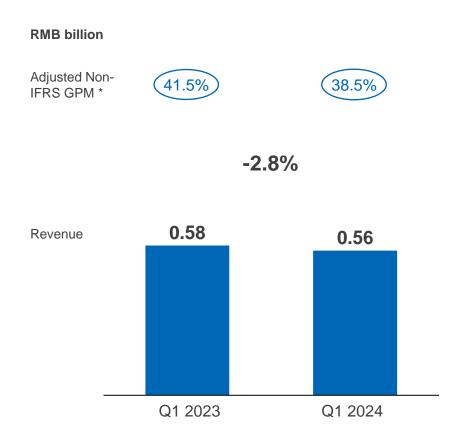
Clinical CRO & SMO

- Q1 revenue from clinical CRO and SMO grew 11.0% YoY to 0.44bn, among which, SMO revenue grew 26.4% YoY, maintaining industry leading position in China
- In Q1, clinical CRO enabled customers to obtain 7 IND approvals;
 SMO supported 10 new drug approvals
- SMO business sustains rapid growth, gaining market share in multiple therapeutic areas (lung cancer, breast cancer, dermatology, cardiovascular disease, ophthalmology, rheumatology, central nervous system, endocrinology and medical aesthetics etc.)



WuXi Biology: New Modalities Business Drives Growth; WuXi Biology Platform Continues to Generate Downstream Opportunities

Revenue & Gross Profit Margin



^{*} GPM @ CER: Q1 2023 41.3%, Q1 2024 36.6%

Discovery Biology

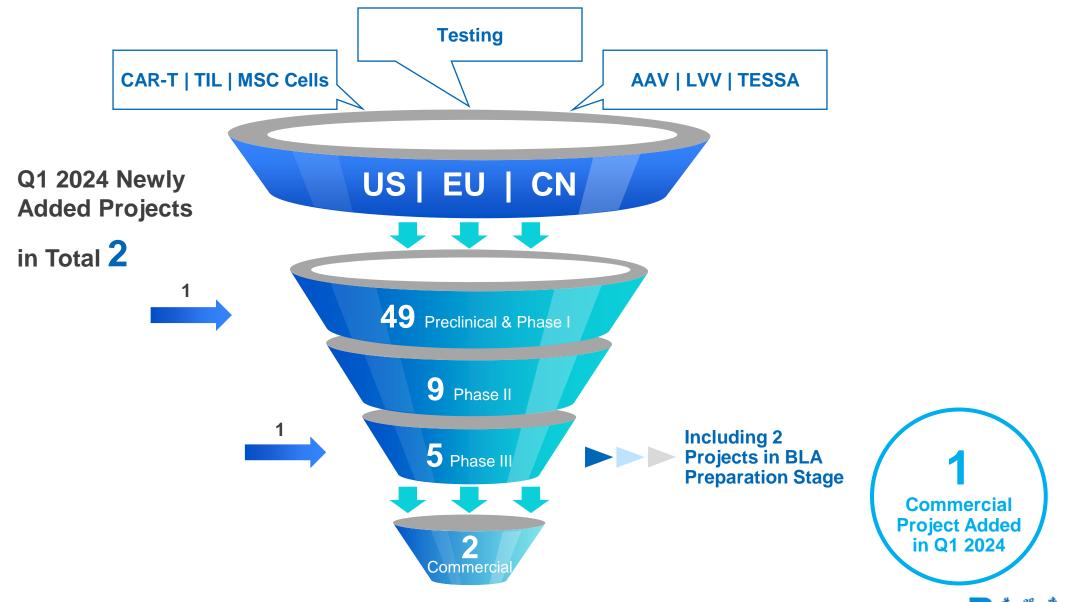
- Focused on improving capabilities related to new modalities, revenue from new modalities grew 12.2% YoY, contributing 29.2% of WuXi Biology revenue
- Number of customers and projects served by the nucleic acid
 platform continued to increase. Cumulatively provided services to
 230+ customers, and successfully delivered 1000+ projects since 2021
- Further integrated resources of *in vivo* pharmacology platform, and continued to improve platform capabilities and efficiency

Comprehensive Screening Platform

- Commenced operations at Suzhou Guoxiang research platform by end of March 2024, strengthening new capabilities of *in vitro* biology and *in vivo* pharmacology
- Continued to generate downstream opportunities and contributed
 20%+ of the Company's new customers



WuXi ATU: Advancing Pipeline on the Globally Integrated CTDMO Platform



WuXi ATU: Commercial Projects Gradually Approved, Driving Future Growth

Revenue & Gross Profit Margin

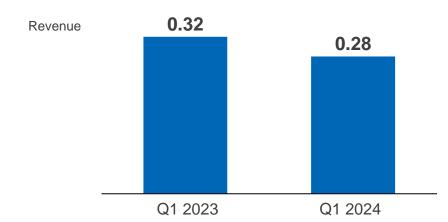


Adjusted Non-IFRS GPM *









Pipeline & Progress of Commercial Projects

- Total pipeline of 65 projects by end of Q1, among which, the world's first innovative TIL-based therapy was approved by FDA in Q1
- Preparing for BLA filing to manufacture the LVV used in a commercial CAR-T product. Completed process performance qualification (PPQ) and started post-PPQ manufacturing, expect to file preapproval submission (PAS) to FDA in H2 2024
- Expect to complete PPQ and file PAS to FDA in H2 2024 for a blockbuster commercial CAR-T product
- Q1 revenue and GPM down YoY, mainly due to: high-margin projects were completed in 2023; the new TIL product was approved in late February 2024, while manufacturing and quality staffs have been hired and came onboard in advance to prepare for commercial manufacturing, and thus costs increased accordingly; and underutilization of capacity for other projects



^{*} GPM @ CER: Q1 2023 -7.4%, Q1 2024 -34.2%

WuXi DDSU¹: Supporting Customers to Continue Advancing Pipeline Projects





NDA Approval

 By end of Q1, 3 new drugs developed for our customers obtained NMPA approvals, including 2 for COVID-19 infection treatment and 1 for tumor treatment. Continued to receive royalty income from customers for the approved new drugs. Moreover, 2 new drug candidates are in NDA review stage

Project Pipeline

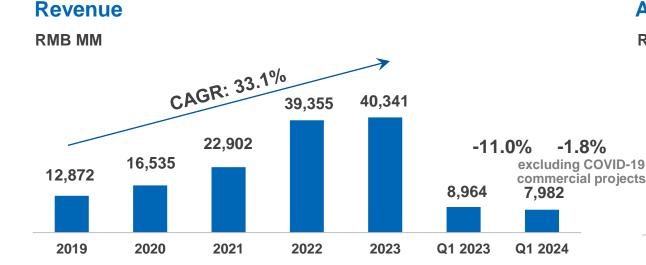
- In Q1, supported customers to file INDs for 6 drug candidates and obtained 6 Clinical Trial Approvals (CTAs)
- Cumulatively submitted 196 IND filings and obtained 175 CTAs, including 3 projects with NDA approvals, 2 projects in NDA review stage, 4 in Phase III, 35 in Phase II, and 71 in Phase I
- 17 new modality projects covering peptide/PDC, PROTAC®2 and oligo. Multiple projects have completed pre-clinical development and filed INDs, among which 6 projects have entered clinical stage
- Demand declined significantly, and thus fewer new projects were signed as expected



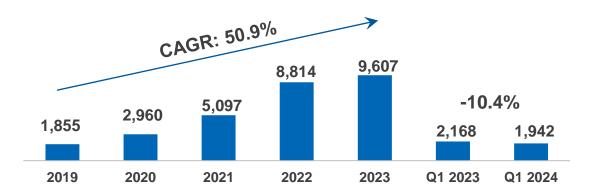
3. Financial Performance



Financial Performance



Net Profit Attributable to the Owners of the Company RMB MM



Adjusted Non-IFRS GPM **Adjusted Non-IFRS Gross Profit RMB MM** 40.0% 40.3% 38.0% 38.7% 42.0% 41.0% 38.7% CAGR: 34.6% 16,938 15,228 -16.0% 8,704 6,670 5,155 3,678 3,090

Adjusted Non-IFRS Net Profit Attributable to
the Owners of the Company

Adjusted Non-IFRS NPM

RMB MM

2022

2023

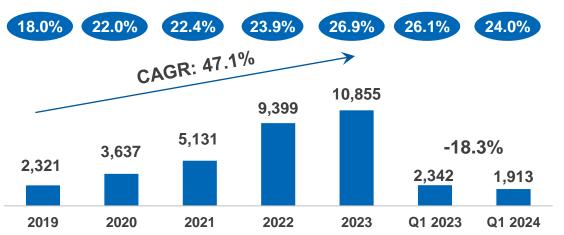
Q1 2023

Q1 2024

2019

2020

2021

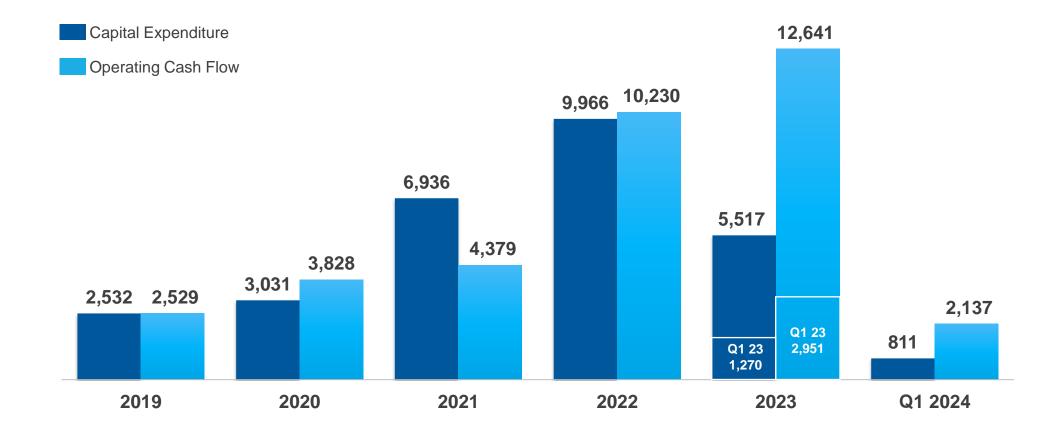




Sustained Positive FCF Driven by Business Growth & Operating Efficiency

Q1 2024 Free Cash Flow Remained Positive and Reached RMB 1.33Bn

RMB MM

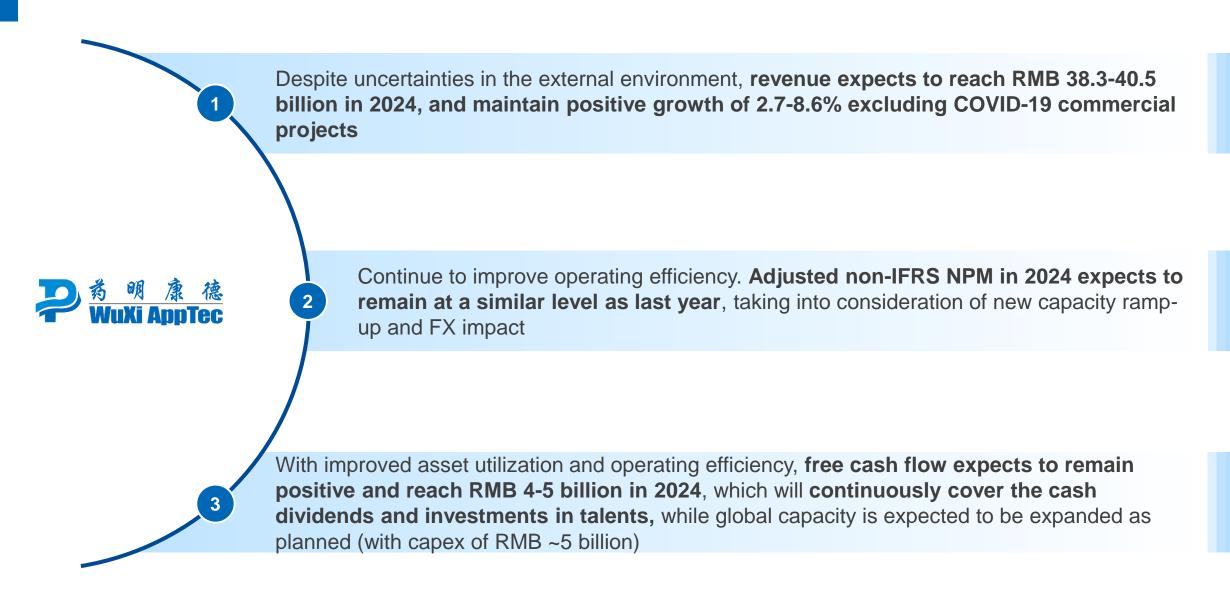




4. Business Outlook



2024 Outlook



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