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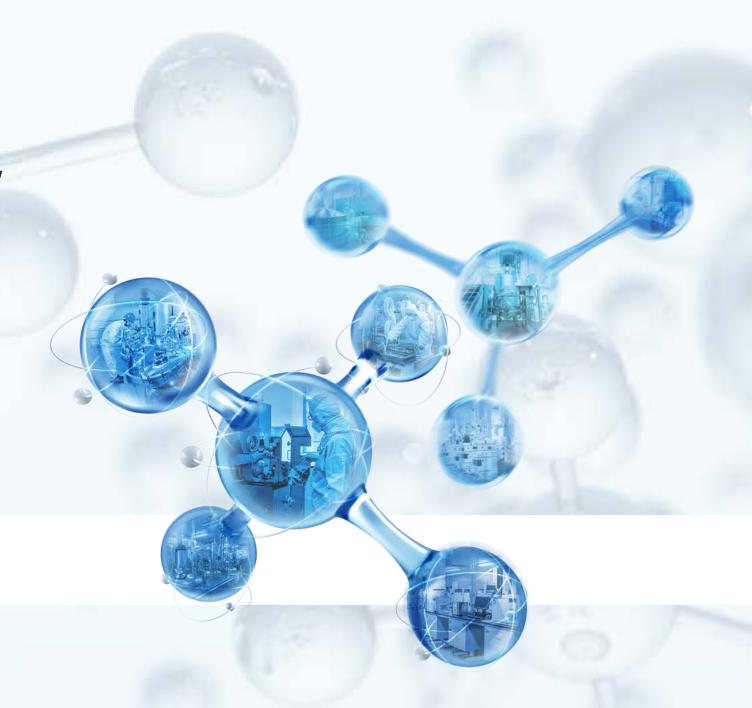
Use of Adjusted Financial Measures

We have provided adjusted net profit attributable to shareholders of the Company and adjusted net profit margin attributable to shareholders of the Company as additional financial measures, which are not required by, or presented in accordance with, the 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, non-recurring, non-cash and/or non-operating 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 the IFRS. You should not view adjusted results on a stand-alone basis or as a substitute for results under IFRS.

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2,655

Revenue (RMB mm)

(42.2%)

Excluding large orders (0.3%)

41.2%

Gross Profit Margin

(11.6pts)

@CER (12.4pts)

499

Net Profit Attributable to Shareholders of the Company (RMB mm)

970

Backlog (\$ mm)

18.8%

Net Profit Margin Attributable to Shareholders of the Company

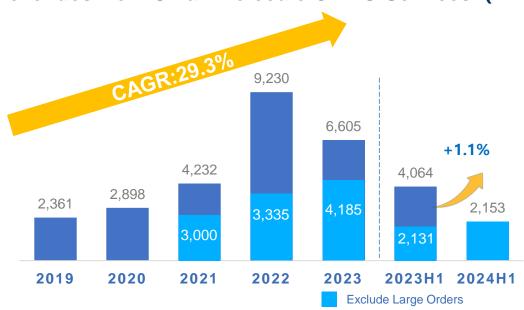
30.0%

EBITDA Margin



Dual-Engine Driven Strategy Steadily Advanced

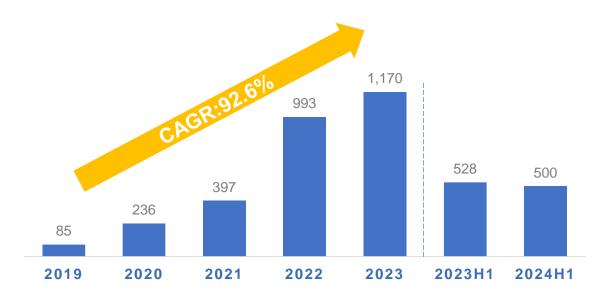
Revenues from Small Molecule CDMO Services (RMB mm)



Small Molecule CDMO Service

- Revenue reached RMB2,153 mm marking a pop¹ increase of 1.1% excluding large orders
- Delivered 353 projects with an increase of 13.9% vs. 2023H1, including:
- **61** clinical phase III projects
- 43 commercialization projects
- Commercialization projects achieved revenue of **RMB1,366 mm** with a *pop* growth of **7.1%** if excluding the effect of large orders in 2023H1

Revenues from Emerging Services (RMB mm)



Emerging Business

- Emerging business contributed RMB500 mm in revenue, experiencing a 5.3% decrease vs. 2023H1, the gross profit margin was 20.2% due to the continued downturn with domestic market and some businesses still being in a capacity ramp up phase
- Continues to focus on enhancing competitiveness and actively advancing market expansion. As of this announcement date, the estimated emerging business PPQ² reached 9, forming a sufficient reserve of commercial orders



Large Pharma and Overseas Customers Maintain Revenue Growth

RMB mm

Revenue from All Types of Customers

Large Pharma

Exclude Large Orders +10.3%

1,282

Mid-to-Small Pharma (8.4%)

1,373

Overseas Market

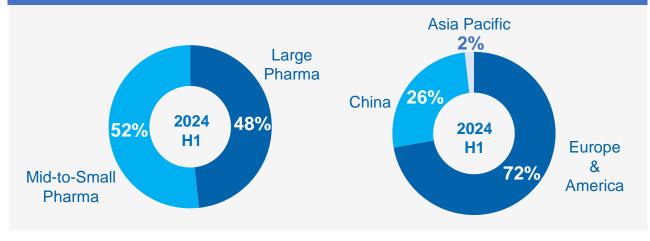
Exclude Large Orders +3.5%

1,966

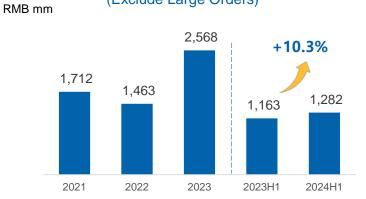
Domestic Market* (9.5%)

689

- Expanded **114** new customers during 2024H1, and achieved a *pop* growth of over **20% in new orders**
- A significant quarter-on-quarter growth in the 2nd quarter, with order growth from customers in European and American outpacing the overall order growth rate of the Company

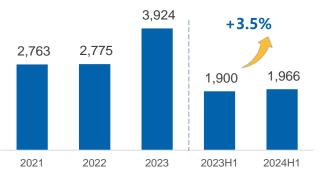


Revenue from Large Pharma (Exclude Large Orders)

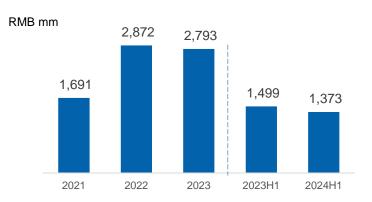


Revenue from Overseas Market

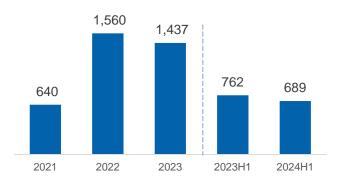
(Exclude Large Orders)



Revenue from Mid-to-Small Pharma



Revenue of Domestic Market



- European market experienced a breakthrough in revenue, with a growth of 22.1%
- U.S. Clients Revenue was RMB1,742 mm, showing a substantial pop growth of 24.8%
 vs. 2023H1 excluding large orders





Small Molecule Business Maintains Steady Growth

	2022H1	2023H1	2024H1
Pre-clinical and Early Clinical Stages	172	224	249
Phase III Clinical Stage	48	52	61
Commercial Stage	34	34	43

Despite ongoing challenges in global biopharmaceutical industry, small molecule CDMO business:

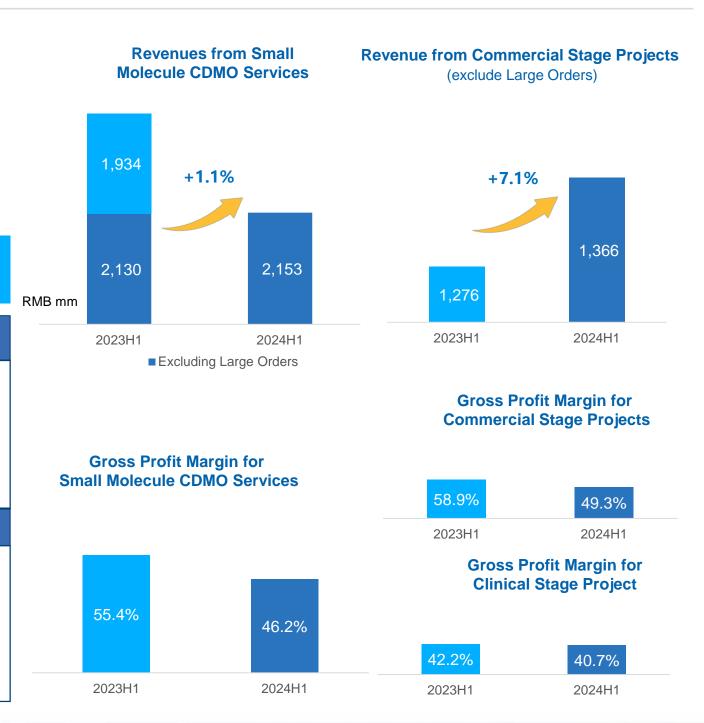
- Generated revenue of RMB2,153 mm, marking a pop increase of 1.1% excluding large orders
- Gross margin achieved 46.2% through improvement of operational efficiency and cost control

Commercialization Projects as the Backbone to Continues Revenue Growth

- Successfully progressed 43 commercialization projects with 9 projects more than 2023H1, resulting in revenue of RMB1,366 mm, and if excluding the effect of large orders, the revenue increased by 7.1%
- With a strong track record in project delivery, the Company is well positioned to foster deeper collaboration with numerous international and domestic clients in commercialization projects

Reserves of Clinical Projects Ensured for Long-term Growth

- Company had 310 clinical stage projects, including 61 clinical phase III. The recognized revenue from clinical projects reached RMB788 mm with a decrease of 7.8%
- It is expected that the number of projects reaching PPQ stage in the 2024H2 will reach 28, which established a sufficient reserve of commercial orders, providing strong support for long-term and steady performance growth





Chemical Macromolecules Business in Steady Progress

(19.3%)

Revenue

72

New Projects

+119%

Backlog

Business Progress

- 2024H1, we secured **several late stage peptide projects** from **major pharmaceutical companies** in Europe and the U.S., successfully passed **domestic GLP-1 peptide project** dynamic verification
- Continued to advance **Oligonucleotide business**, undertaking over **29** new projects, representing a *pop* **70.6**% growth, and made strides in the **commercial deployment of CpG adjuvants**
- The toxin-conjugate business is advancing rapidly, with 11 NDA projects from 8 clients steadily progressing, including 3 projects from 2 overseas clients
- **Lipid business** including cationic lipids, phospholipids, and PEG lipids, is developing rapidly. During the reporting period, **12** lipid projects were advanced in parallel, securing **4** IND stage projects from **2** MNCs¹ clients and establishing partnerships with several domestic clients

Capacity Expansion

- Company accelerated the construction of peptide commercialization production capacity to 14,250 liters of total solid-phase peptide synthesis capacity by the mid 2024 to meet the commercial production needs of domestic and international clients
- For R&D platform, we continue to advance new technology and process development. Enzyme conjugation technology has been developed and applied in the solid-phase and liquid-phase synthesis of oligonucleotides and peptides. We completed the development of substrate-related synthesis processes for RNA synthesis from scratch using enzymes and for enzyme-mediated peptide fragment conjugation. In addition, we have accumulated process development and optimization for various novel toxin conjugates



Technology Platform Enables Smooth Progress of Drug Products

(17.0%)

Revenue

No. of Projects **Completed**

Overseas Projects Completed

Business Progress

- We successfully completed 80 projects, with 150 ongoing projects in progress, encompassing 36 overseas projects. Our commercialized drug products are now ensuring stable market supply
- Underwent and passed 5 on-site inspections by drug regulatory agencies and over 20 audits by domestic and international customers, further demonstrating our robust quality management system and international standard cGMP production capacity, laying a solid foundation for providing services from clinical to commercial production
- Capability in aseptic formulation has continued to strengthen, the variety of complex formulation project types has continued to expand; rapidly undertook and delivered multiple types of LNP1 projects through the self-developed LNP technology platform
- Capacity in high-potency aseptic formulation has been established and enhanced; the number of aseptic formulation projects involving small nucleic acid and polypeptide has significantly increased

Technology Platform

- The solid dispersion technology platform continues to be solidified, successfully completing production batches for multiple late stage projects for NDA registration and PPQ, helping customers meet demands for improving bioavailability of multiple insoluble API²
- The sustained strengthening of the oral peptide delivery technology platform has successfully facilitated clinical product delivery for multiple projects, significantly enhancing the bioavailability of peptides
- Nanocrystal technology platform has been continuously refined with multiple projects steadily progressing
- New production capacity for drug product business, encompassing pre-filled syringes and pen syringes, has been initiated and is expected to commence operation by 2025, with an annual output of up to 40 mm units per single production line, providing a robust assurance for undertaking new projects



Business Scope of CRO Continues to Expand

(23.4%)

Revenue

159 New Projects

103

phase II and later stage projects

Business Progress

- Continued to implement our "One-stop Integrated Development Services" Grand Strategy, seamlessly connecting CMC, non-clinical, and clinical services to support customers in new drug research and development and undertook 19 integrated service orders and successfully obtained 3 implied China IND approvals
- Our global business continued to grow with 12 new global regulatory and clinical services orders. We initiated 2 cell therapy U.S. IND preparation for customers, contributing to 3 successful FDA submissions for customers
- Our regulatory affairs services facilitated 10 projects to obtain China IND approvals. The Company assisted 1
 phase III oncology project IDMC¹ to pass EMA¹ review
- 2024H1, Company was conducting 281 clinical research projects, including 103 phase II and later stage projects

Capability Development

- We have reinforced our strengths in oncology, immunology, infectious diseases, orthopedics, respiratory, hematology disease areas, while achieving breakthroughs in metabolism, digestion, dermatology, and urology
- We advanced further in **rare diseases**, we have been awarded projects ALS², IPAP², brain glioma, IPF², Castleman³, ATTR-CM³, PKU³ and other fields, **accumulating extensive experience and expertise**



Steady Growth in Biological Macromolecules CDMO Business



+1.9%

Revenue

100+

Orders in Hand

50%+

Proportion of ADC Projects

Business Progress

- Amassed nearly 100 orders in hand, including IND, clinical and multiple BLA stage projects, with ADC projects accounting for over 50% of the total, and successfully securing orders from several overseas customers
- Passed the **EU QP audit** and obtained the **EU QP GMP Compliance Statement** in February 2024. In 2024H1, we underwent over **10** customer audits and third-party joint audits, with no major findings

Capability Development

- Leveraging rich project experience and specialized expertise in the toxin-linker sub-field, we swiftly developed a
 one-stop ADC¹ service capability. We have established a diverse and sophisticated conjugate process
 platform, enabling us to offer a full range of CMC services, from IND to BLA for ADC
- Supported by Asymchem's renowned quality management system, we have fully established an international biopharmaceutical quality management system upon the characteristics of biopharmaceutical macromolecules



Export of New Technology Ushered Breakthroughs

~50_{mm}

Revenue

Marketing Expansion

- Undertook **9** new overseas technology output projects
- Engaged with over hundreds customers across 28 provinces and municipalities

Business Progress

- Several industrialization projects are underway, including the commissioning trial of an advanced green pesticide enterprise (3,000mt/a) scheduled in 2024H2
- Will further penetrate both domestic and overseas markets, concentrate on order securing and delivery, and consistently uphold our leadership in continuous flow technology R&D and application

Focus on ensuring order fulfillment and delivery, technological innovation, capacity building, and service capability extension

13

New Patents

21

Newly Authorized Patents

 $>30,000 \,\mathrm{m}^2$

R&D and Manufacturing
Center

>300

Team Member



Continuing to Push Forward with Our Biosynthesis Technology

Leveraging on strong R&D capabilities and over a decade of technological accumulation, we have established a mature one-stop synthetic biology service capability starting from molecular biology (recombinant expression)

Minimum enzyme evolution cycle shortened to 1 week

Carried out several commercial production projects for tonnage continuous enzymatic catalysis

Established a fully continuous platform for nonnatural amino acid synthesis

CSBT Platform

Artificial Intelligence

High Throughput Screening

Continuously Enzymatic Catalysis Technology

Cell Biosynthesis

Achieve efficient cellular synthesis of multiple biobased small molecules

Design polypeptide synthetic biology technology routes, develop high-yield strains, conduct process development, and achieve efficient production

Developed technology platform for oligopeptide, polypeptide biosynthesis

Business Progress

+92.8%

80%+

Revenue

Overseas Revenue

Touch **50+** new customers

Expanded **collaboration with multiple MNCs** to pioneer early technical pathways for enzyme engineering

Capability Development

- Built upon **integrated enzyme engineering technology platform** encompassing enzyme screening, featuring enzyme screening, development, evolution, immobilization, fermentation production and process scale-up, we have integrated cell-free protein synthesis and Al-assisted technology, significantly accelerating the speed of the enzyme development and evolution
- Developed immobilized enzyme continuous reaction technology has successfully been applied in the production of multiple ton-scale products.
 Compared to batch reactions, this technology elevates production capacity ranging from 20 to 1,000 times, significantly reducing costs, improving efficiency, and reduces three wastes
- Leveraging our established **peptide biosynthesis technology platform**, we have implemented biocatalytic synthesis technology using non-natural amino acid raw materials, microbial fermentation for peptides, and enzyme ligation technology for peptide fragments in projects, offering clear advantages in cost and yield compared to previous technologies, earning recognition from major overseas customers
- Built microbial cell factory technology platform, combined with core advantages such as enzyme evolution technology, we have developed a series of efficient strain modification technologies and HTS technologies, and possessed the capability to utilize a variety of microorganisms for product development. With these advantages above, we established the layout of diverse product pipelines and continue to advance related R&D efforts



Accelerated Globalization and Commenced the First R&D and Pilot Production in Europe



We successfully secured a **research and API pilot production** located in Sandwich, U.K., and commenced operations in Aug 2024, enhancing our global supply chain and meeting a wider range of global partners' needs



Current Capacity

- Consistently upheld a world-class capability in drug synthesis rapid route design, HTS, mature process, analysis and development as well as production and operational management
- Leveraging its advanced R&D and production facilities, a team with over 15 years of relevant experience on average, and comprehensive drug R&D technology, Asymchem will implement a comprehensive upgrade

Future Plans

- Plans are underway to expand the site include capabilities for peptides and oligonucleotides
 production, utilizing continuous flow and biocatalysis technologies to enhance sustainability
- Further advance the development of **overseas commercial capacities**, accelerate the adoption and application of new technologies abroad, and offer more partners efficient, flexible, and **high-quality one-stop CDMO solutions**

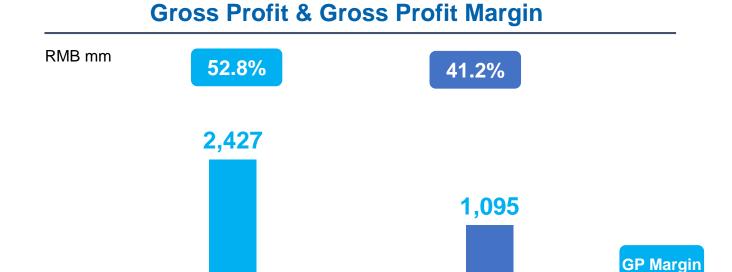




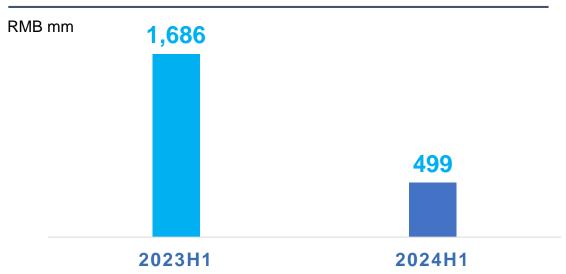
Key Financial Indicators

2024H1



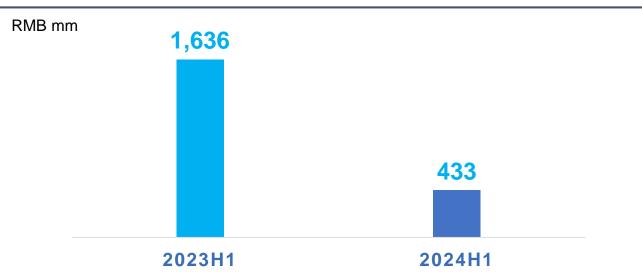


Net Profit Attributable to Shareholders of the Company



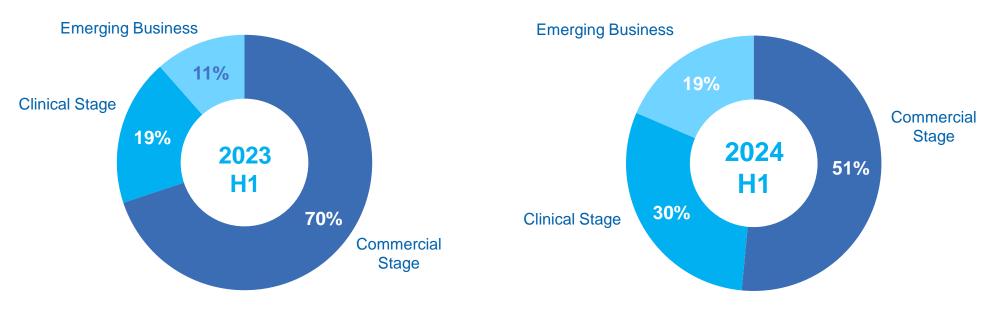
Adjusted Net Profit Attributable to Shareholders of the Company¹

2023H1

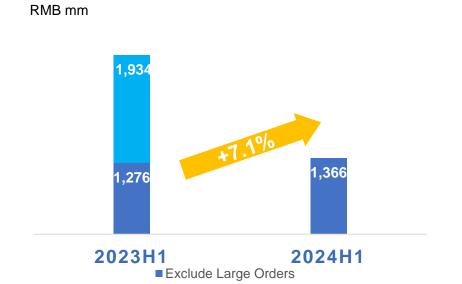




Segment Revenue



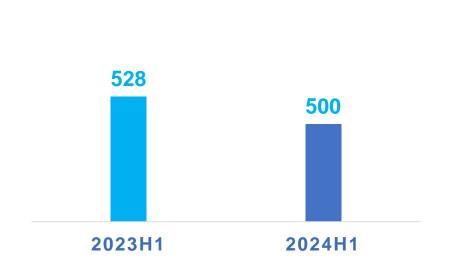




Clinical Stage CDMO Services



Emerging business

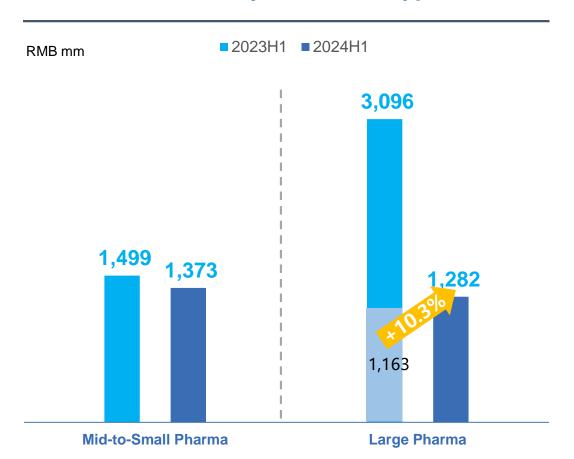




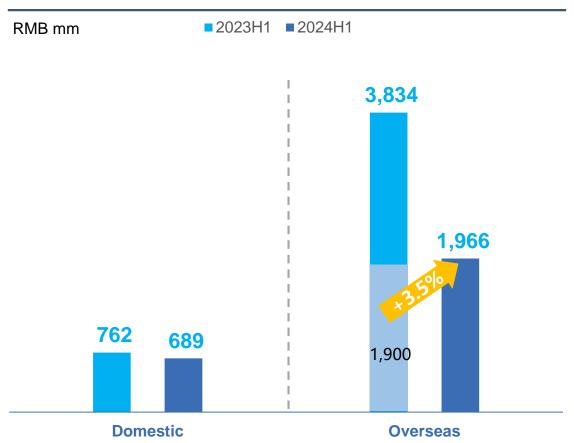
Revenue Breakdown by Customer Type

By consistently delivering high-quality services to key clients, the Company has expanded its customer base both domestically and internationally, resulting in increased market share

Revenue by Customer Type



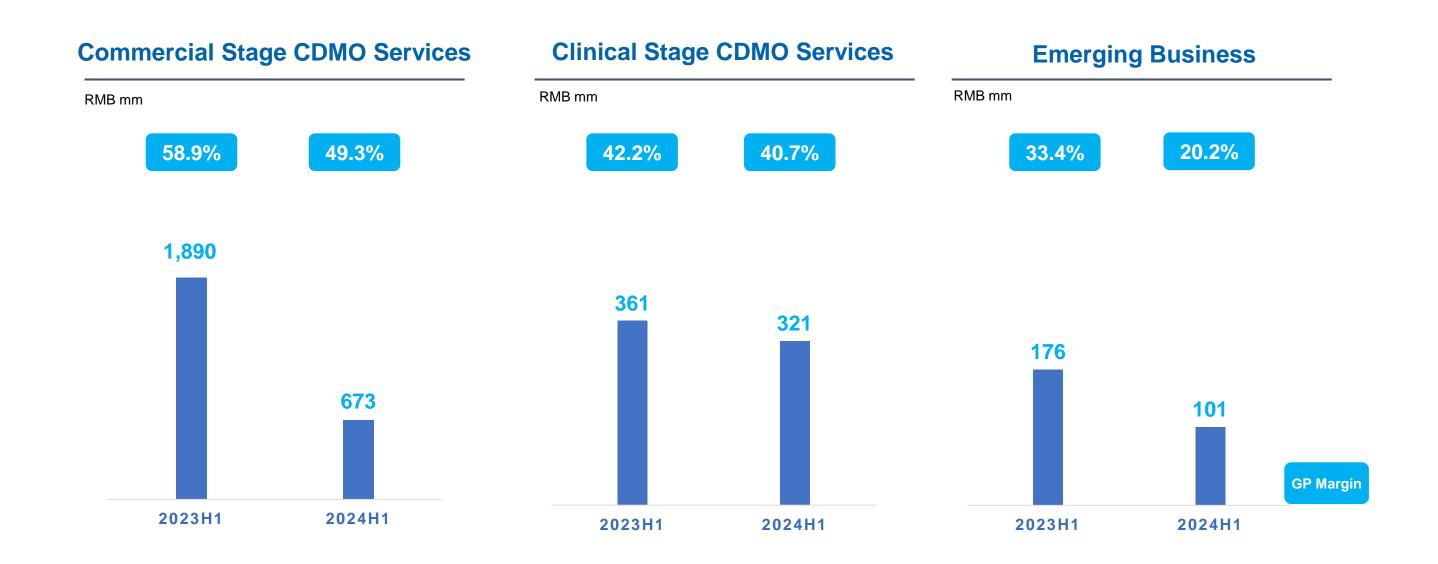
Revenue by Customer Region



Exclude Large Orders

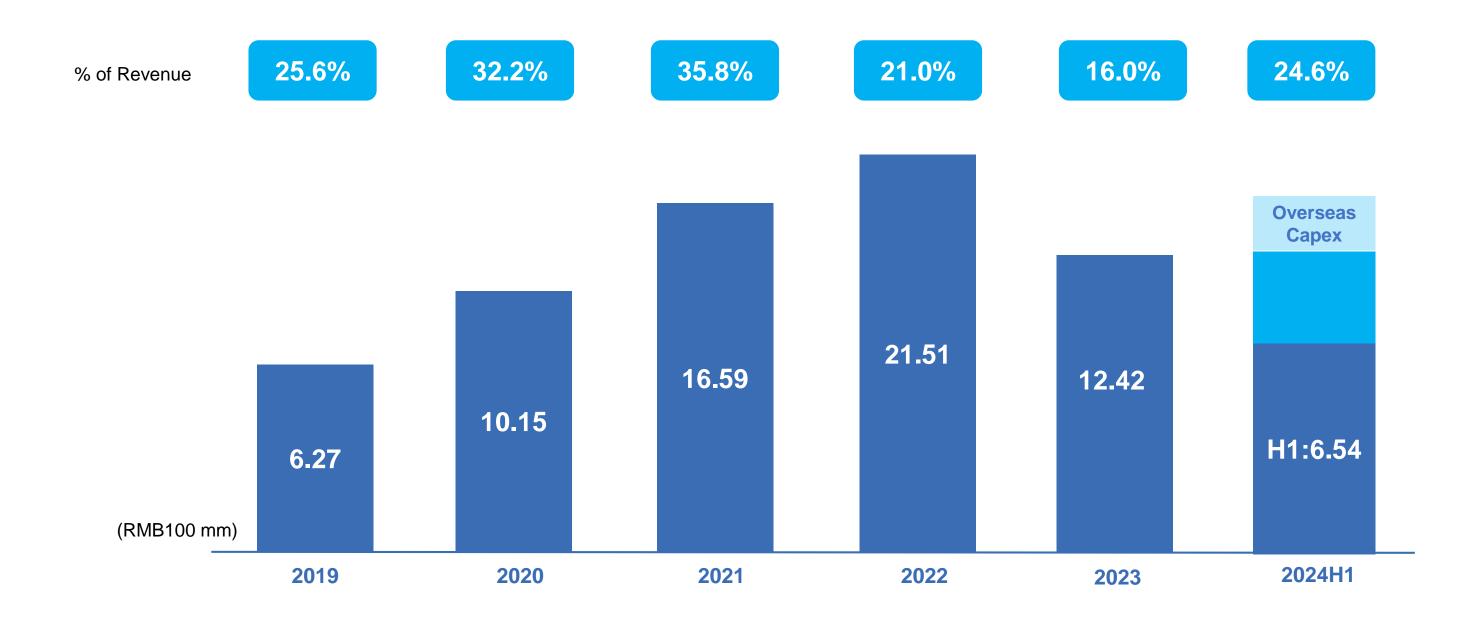


Segment Gross Profit and Gross Profit Margin







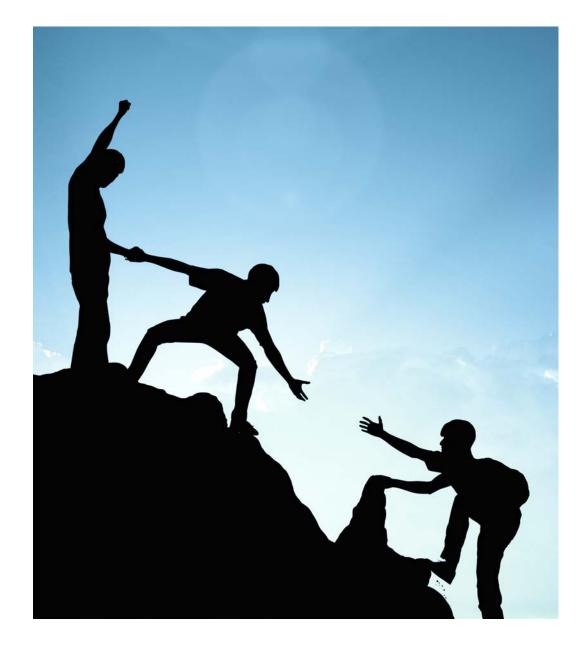




ASYMCHEM









Against the backdrop of industry restructuring, and in light of the market environment, we expected 10%+ growth in non-covid business revenue



Optimize overall profitability of the Company and ease the downward pressure on its margins



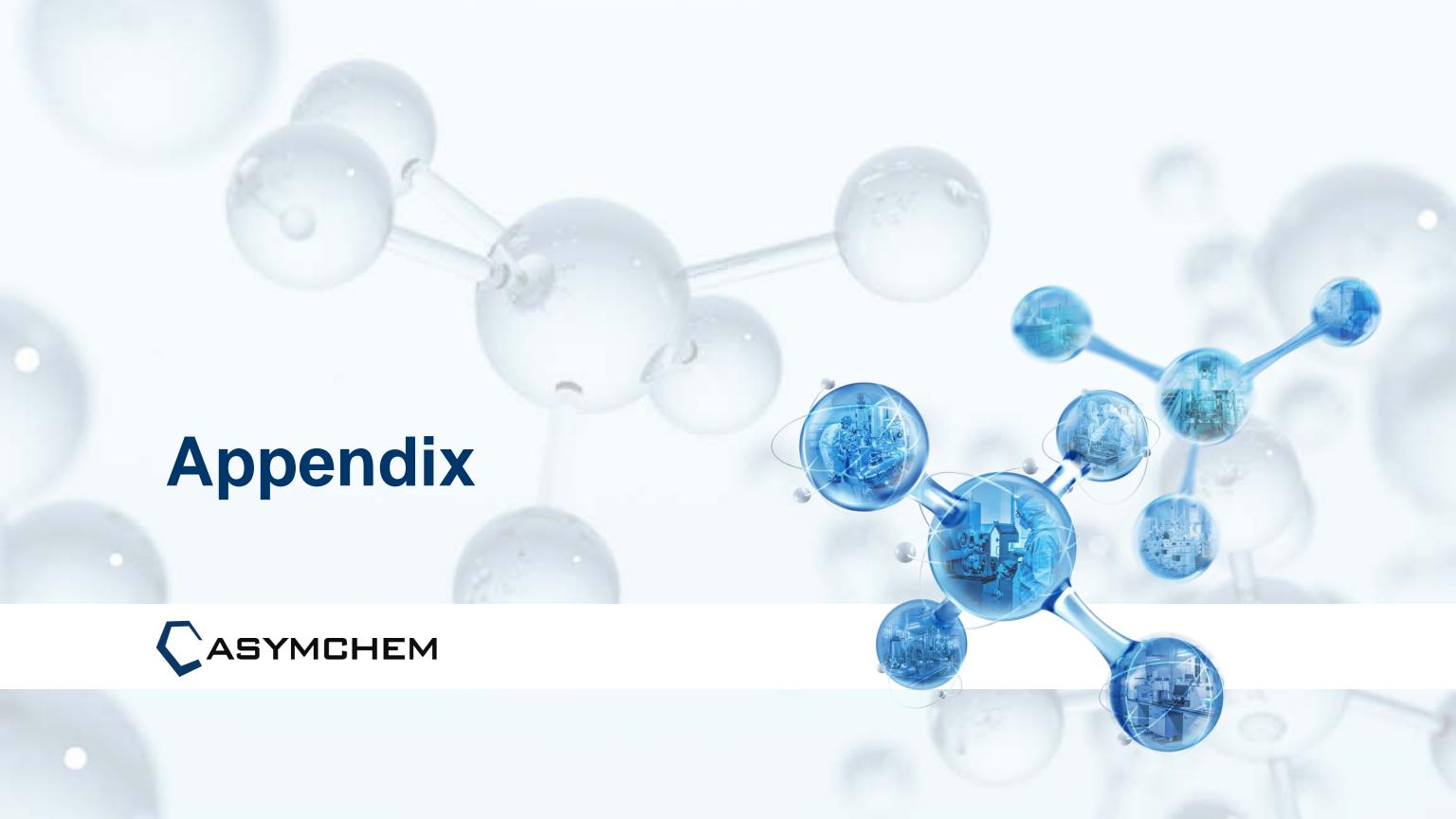
Increase market development and enhance business competitiveness, ensuring the timely delivery of vital projects in emerging businesses i.e. peptides and ADCs to lay the foundation for sustained business growth in 2025



Continuing to promote the construction of the capacity of small molecule API from pilot to commercial stage



Domestic CAPEX would be roughly in line with 2023





CONSOLIDATED STATEMENT OF PROFIT OR LOSS

	2024H1	2023H1
Revenue	2,655	4,596
Cost of sales	(1,560)	(2,169)
Gross profit	1,095	2,427
Other income and gains	258	289
Selling and distribution expenses	(102)	(82)
Administrative expenses	(376)	(351)
Research and development expenses	(329)	(323)
Losses on impairment of financial and contract assets, net	7	(16)
Other expenses	(12)	(9)
Finance costs	(3)	(3)
Share of profits/(losses) of associates	(6)	(3)
Profit before tax	532	1,928
Income tax expense	(40)	(246)
Profit for the year	492	1,682
Attributable to:		
Owners of the parent	499	1,686
Equity incentive amortization expense	34	23
Gain or loss on exchange rate fluctuations	(112)	(82)
Income tax effect	11	9
Adjusted Net Profit Attributable to Owners of the company	432	1,636



CONSOLIDATED STATEMENT OF FINANCIAL POSITION

	2024H1	2023
Non-current assets		
Property, plant and equipment	5,855	5,366
Right-of-use assets	552	526
Goodwill	146	146
Other intangible assets	50	54
Deferred tax assets	258	213
Investments in associates	533	260
Prepayments, deposits and other receivables	637	689
Financial assets at fair value through profit or loss	156	131
Equity investments at fair value through other comprehensive income	35	30
Total non-current assets	8,222	7,415
current assets		
Inventories	998	945
Trade receivables	1,483	2,011
Contract assets	98	81
Prepayments, deposits and other receivables	322	297
Tax recoverable	16	3
Financial assets at fair value through profit or loss	2,041	1,906
Cash and bank balances	5,679	7,110
Total current assets	10,637	12,352
Total assets	18,859	19,767





(continued)

	2024H1	2023
Current liabilities		
Trade payables	388	452
Other payables and accruals	1,446	1,276
Interest-bearing bank borrowings	-	12
Lease liabilities	33	29
Amounts due to related party	1	1
Tax payable	26	31
Total current liabilities	1,894	1,801
Non-current liabilities		
Deferred income	253	233
Lease liabilities	133	106
Deferred tax liabilities	126	117
Total non-current liabilities	512	456
Equity		
Share capital	369	369
Restricted Shares under share-based payment	(1,463)	(494)
Other reserves	17,523	17,605
Non-controlling interests	24	30
Total equity	16,453	17,510

