



J.P. Morgan Healthcare Conference 2026
Hangzhou Tigermed Consulting Co., Ltd.

300347.SZ / 3347.HK

San Francisco, CA, U.S.
January 2026

www.tigermedgrp.com

Disclaimer

By reading these materials, you agree to be bound by the following limitations:

The information herein has been prepared by representatives of Hangzhou Tigemed Consulting Co., Ltd. (杭州泰格医药科技股份有限公司, the “Company”) solely for your information and have not been independently verified. The information herein does not constitute any recommendation regarding any securities of the Company or any of its subsidiaries.

No representation or warranty, express or implied, is made as to, and no reliance should be placed on, the fairness, accuracy, completeness or correctness of the information, or opinions contained herein. The Company, nor any of their respective affiliates, controlling persons, directors, officers, employees, advisors or representatives shall have any responsibility or liability whatsoever (for negligence or otherwise) for any loss howsoever arising from any use of the information herein or its contents or otherwise arising in connection with the information herein. The information or opinions set out herein may be subject to updating, completion, revision, verification and amendment and such information may change materially without notice and shall only be considered current as of the date hereof. The information herein is based on the economic, regulatory, market and other conditions as in effect on the date hereof. Certain information in the materials contain information may be sourced from third parties, which has not been independently verified by the Company. It should be understood that any subsequent developments may affect the information contained herein, which the Company is not under an obligation to update, revise or affirm.

You acknowledge that you will be solely responsible for your own assessment of the market and the market position of the Company and that you will conduct your own analysis and be solely responsible for forming your own view of the potential future performance of the business of the Company and any of its subsidiaries.

Forward-Looking Statements:

The information communicated herein may contain certain “forward-looking statements”, which are not historical facts but instead include predictions about future events based on our beliefs and information currently made available to us. Although we believe that these predictions are reasonable on the date hereof, future events are inherently uncertain and these forward-looking statements may turn out to be incorrect. Forward-looking statements involve risk and uncertainty by nature because they relate to events and will depend on circumstances that will occur in the future relating to, *inter alia*, our ability to compete effectively, our ability to develop and market new service offerings, our ability to expand into new markets, the risks associated with listed subsidiaries of the Company, unforeseeable international tensions, regulatory or governmental scrutiny in certain countries, the impact of emergencies and other force majeure events. We undertake no obligations to update forward-looking statements or to adapt them to future events or developments except as required by applicable laws or listing rules. Any investment in any securities issued by the Company or its subsidiaries will also involve certain risks. There may be additional material risks that are currently not considered to be material or of which the Company and its advisors or representatives are unaware. Against the background of these uncertainties, you should not rely on these forward-looking statements.



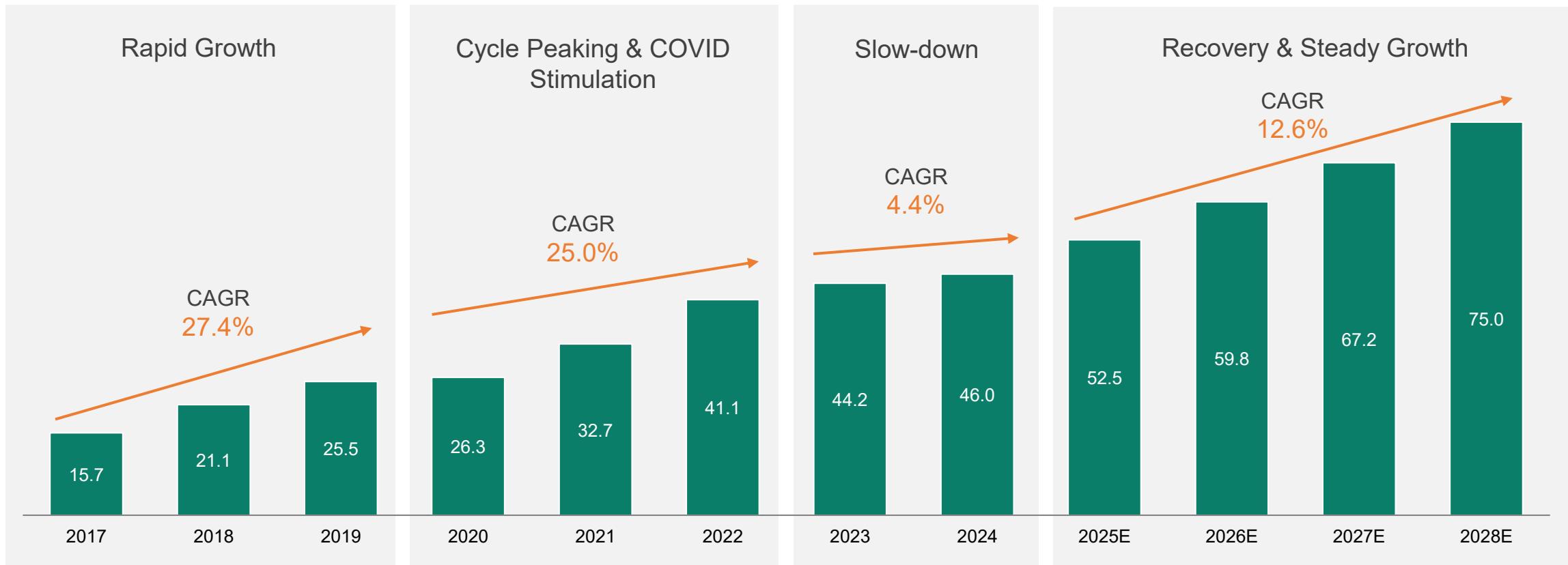
PART 1

China Clinical CRO Industry Update

China Clinical CRO Industry Expected to Recover and Repick Growth

c.50% of China clinical trials were outsourced to clinical CROs during 2019-2025 (excluding peripheral services, data from HGRAC)

China Clinical CRO Market Size *Billion RMB*



Tigermed New Bookings Outgrew the Industry during 2019-2024

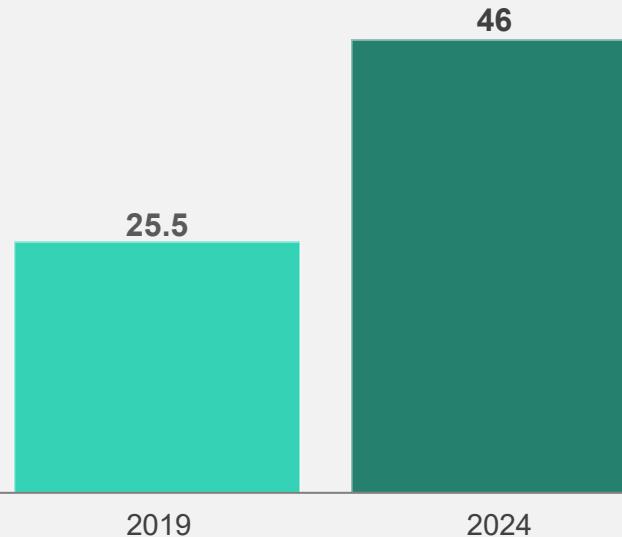
China Clinical CRO Market Size

2019-2024

Billion RMB

2019-2024 CAGR

12.5%



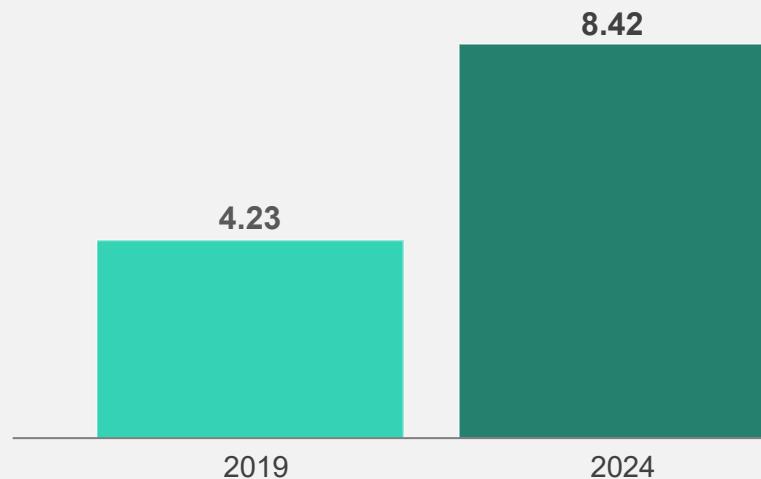
Tigermed Net New Bookings

2019-2024

Billion RMB

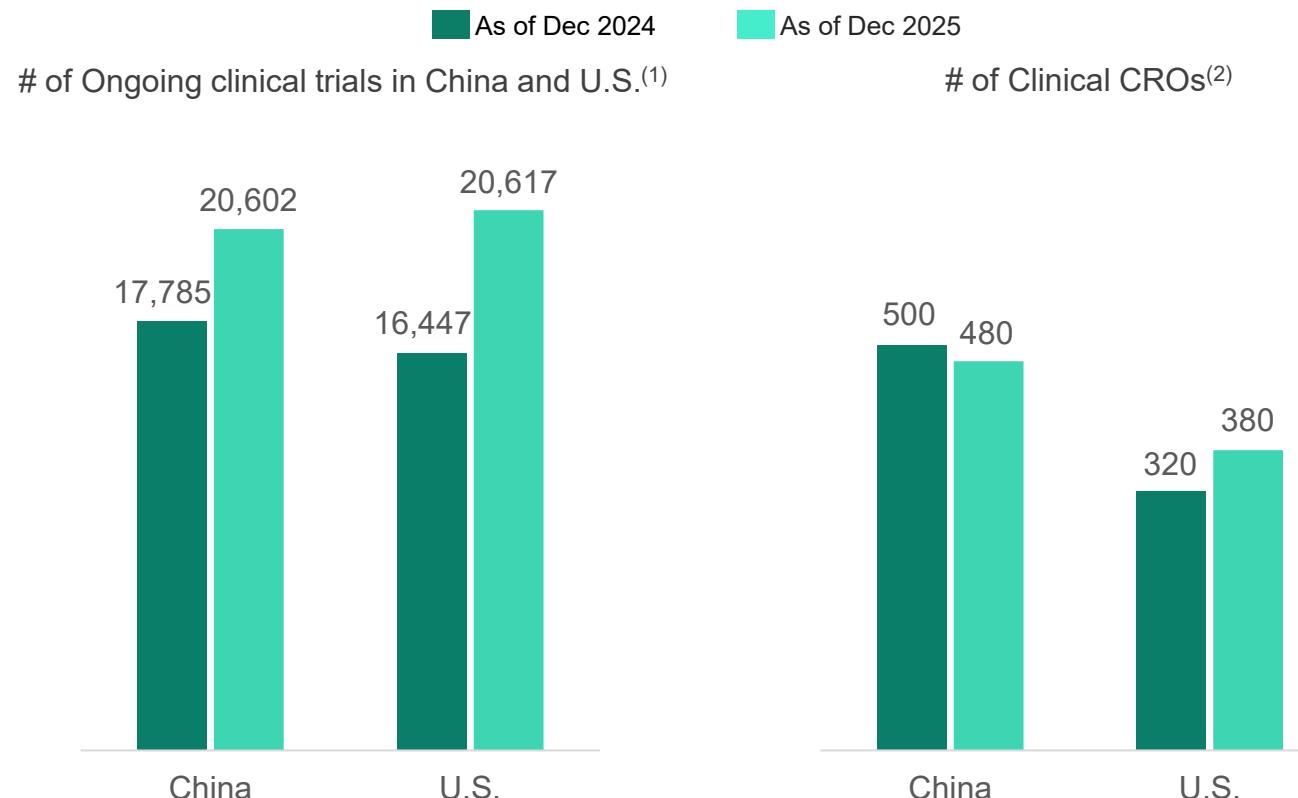
2019-2024 CAGR

14.8%

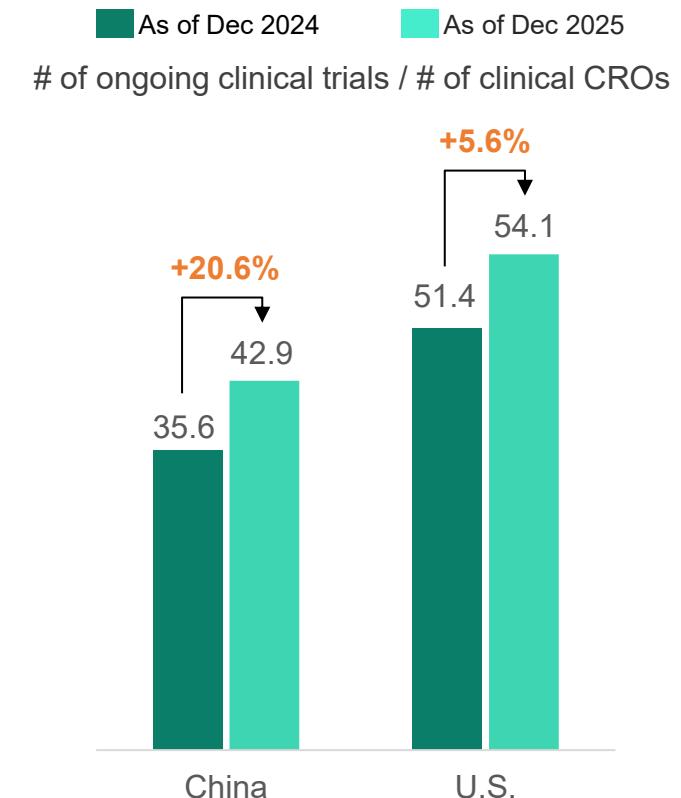


Less Overcapacity Observed Sequentially in the Industry

Supply & demand between China and the U.S. in clinical trial outsourcing



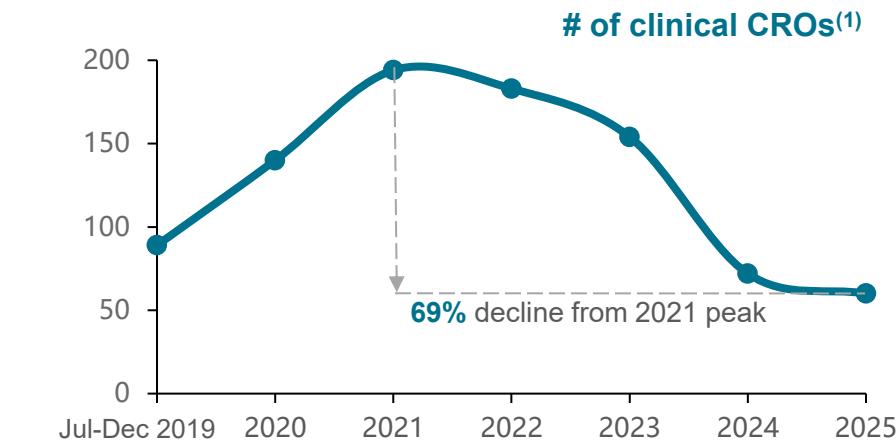
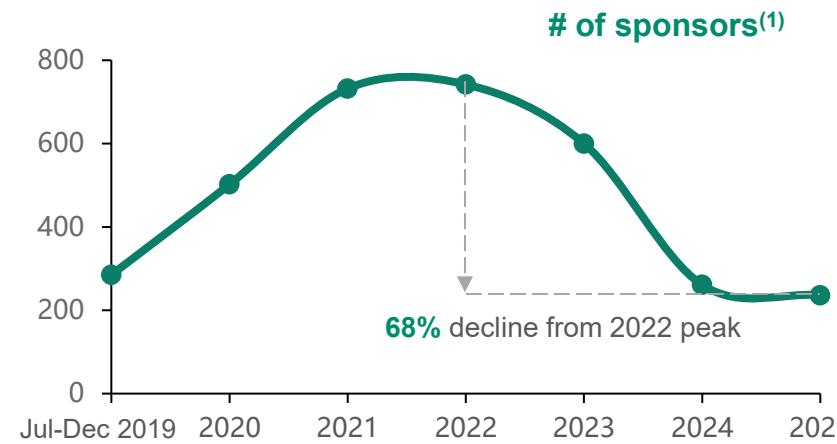
Ongoing trials available per Clinical CRO in China far behind that in the U.S.



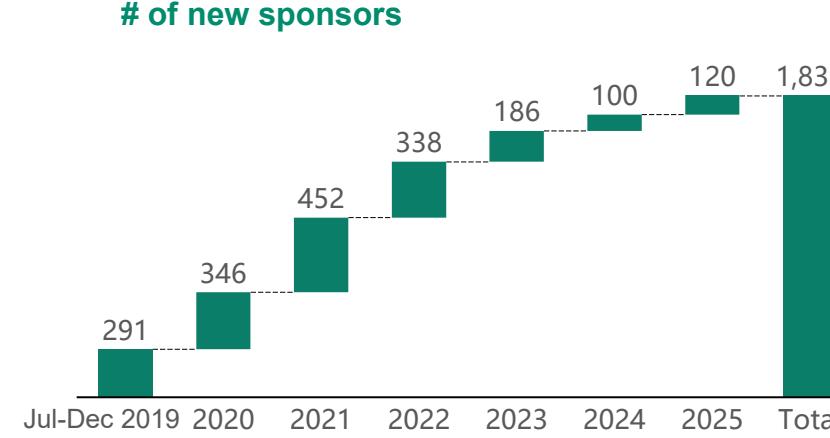
Data May Not Be Exhaustive

Supply Side Consolidation Continued in 2025

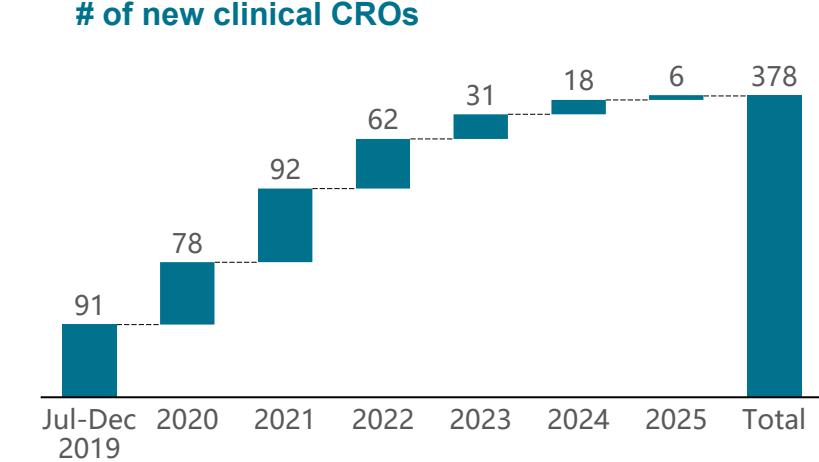
Trend comparison of number of sponsors and clinical CROs each year from HGRAC filings



Trend comparison of newly-added sponsors and clinical CROs each year from HGRAC filings



of new clinical CROs



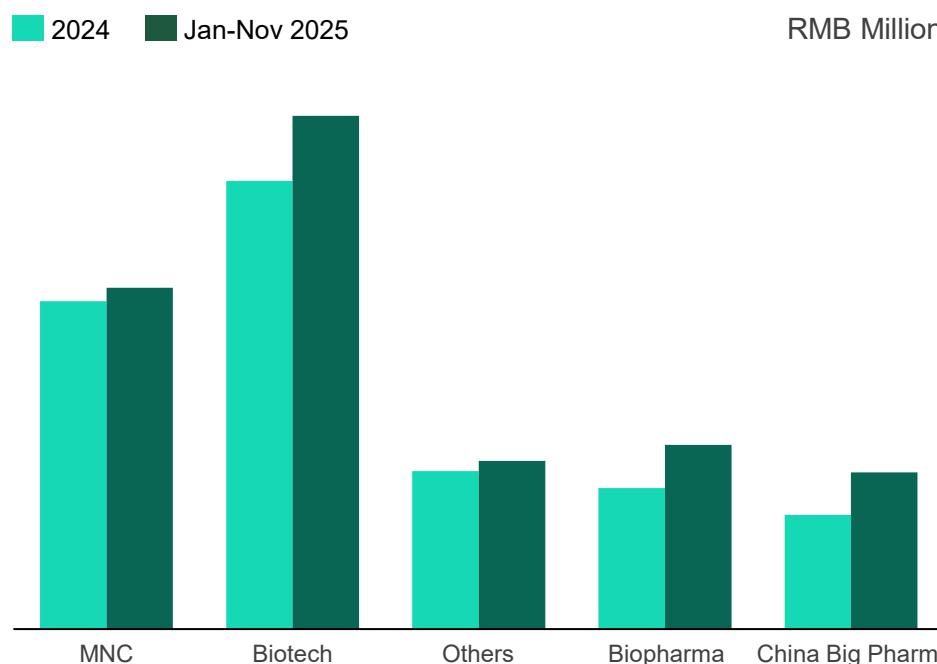


PART 2

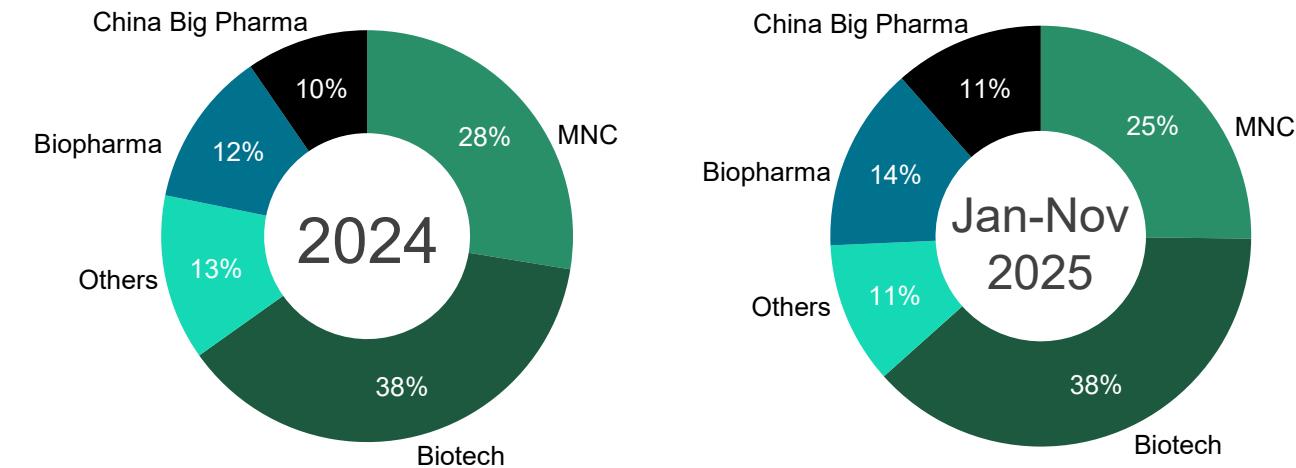
Tigermed New Booking Trend

Increasing New Bookings from MNCs and Chinese Large Pharma

New Booking Breakdown by Customer Type
by Dollar Amount 2024 vs Jan-Nov 2025



New Booking Breakdown by Customer Type
by Dollar Amount 2024 vs Jan-Nov 2025



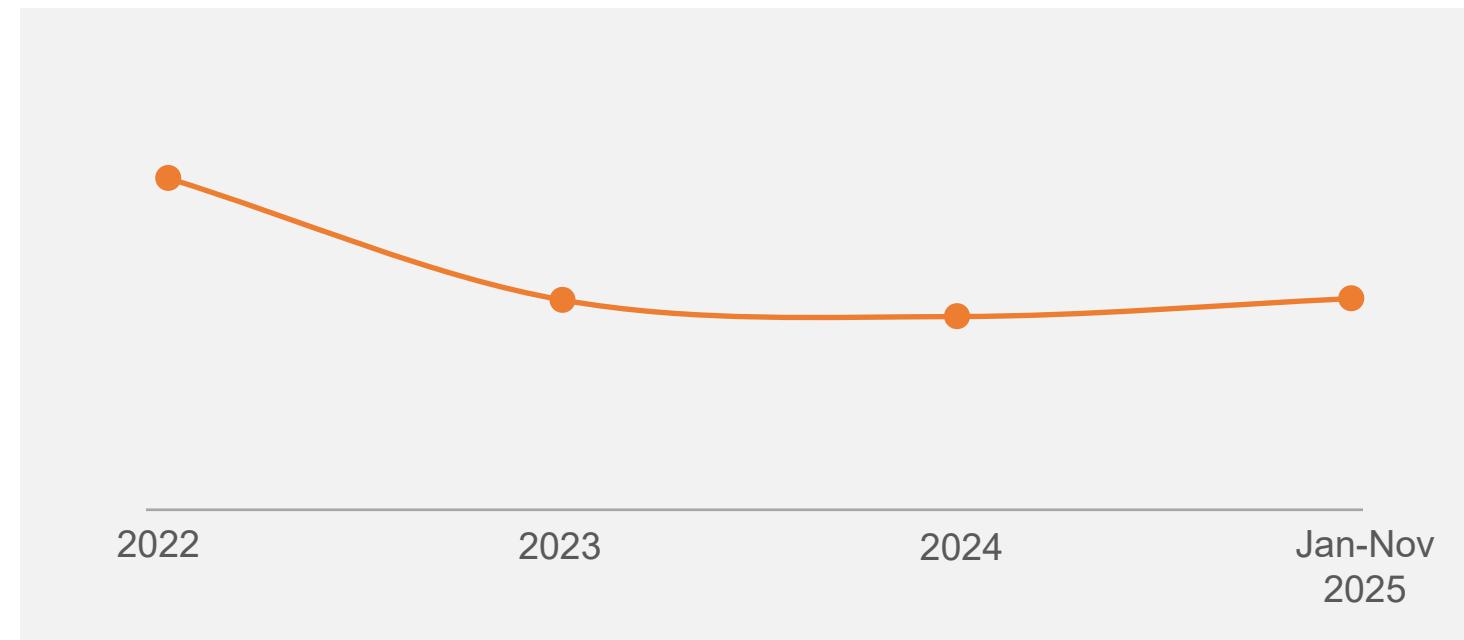
Net new booking data excluding Frontage, DreamCIS and Fantastic Bioimaging

Others include medical device companies, CROs, academic institutions, hospitals, medical/clinical centers, cosmetics and health supplement companies etc.

Domestic New Booking Price Stabilized

Domestic Phase I&II Clinical Trial New Booking Average Selling Price Trend

- ASP = total booking amount / number of trials
- Data of Tigermed new booking ASP of domestic Phase I and II clinical trial projects in 2022 - Nov 2025
- Domestic clinical trials refer to clinical trials to be conducted in China with Chinese sponsors
- Excluding Phase III clinical trials projects due to variation in size
- Data for illustrative purpose and may not be exhaustive



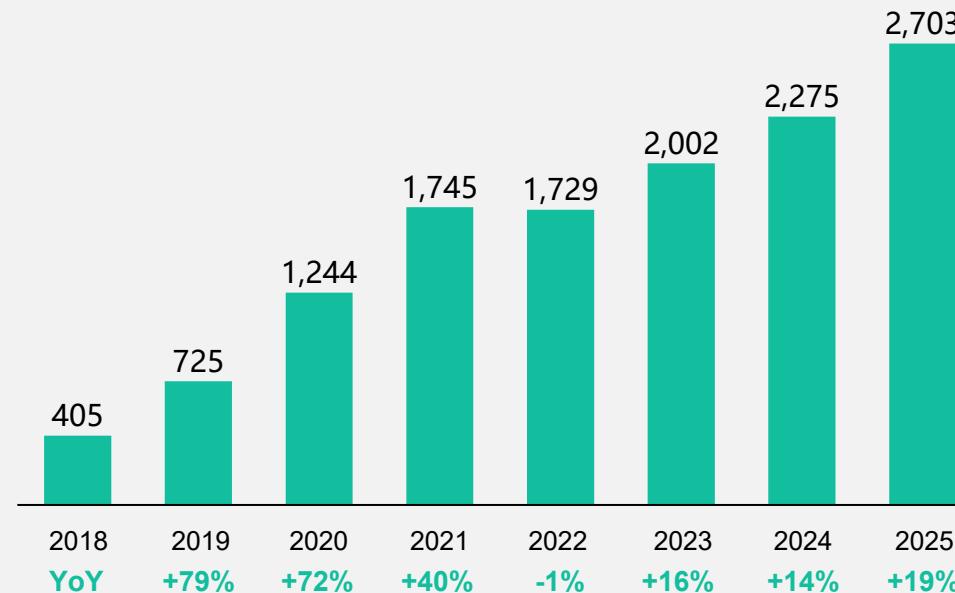


PART 3

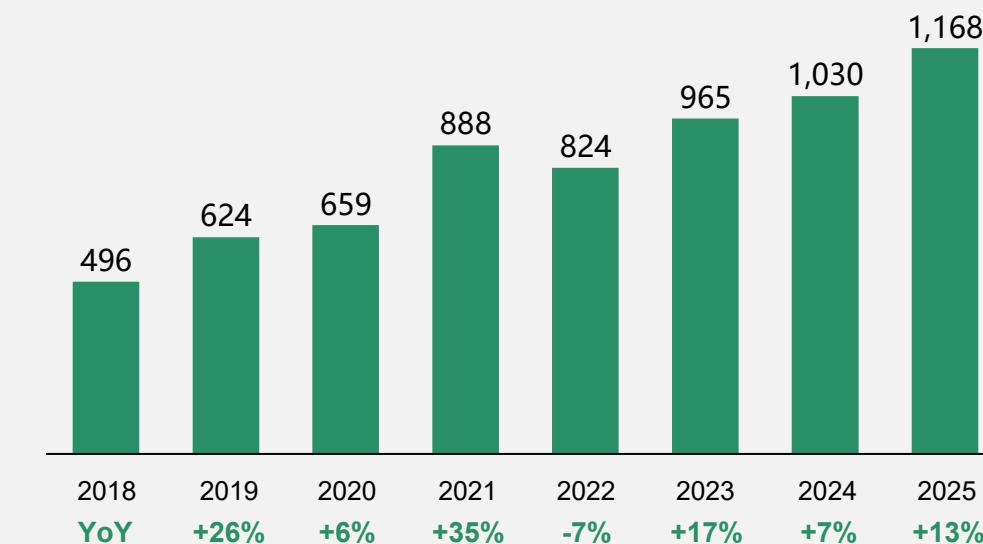
R&D Landscape – Spending and Clinical Trials

Faster Growth of INDs and First-in-Human Trials in 2025

Number of Clinical Trial IND Approvals in China
2018-2025



Number of Phase I Clinical Trials Initiated in China
2018-2025

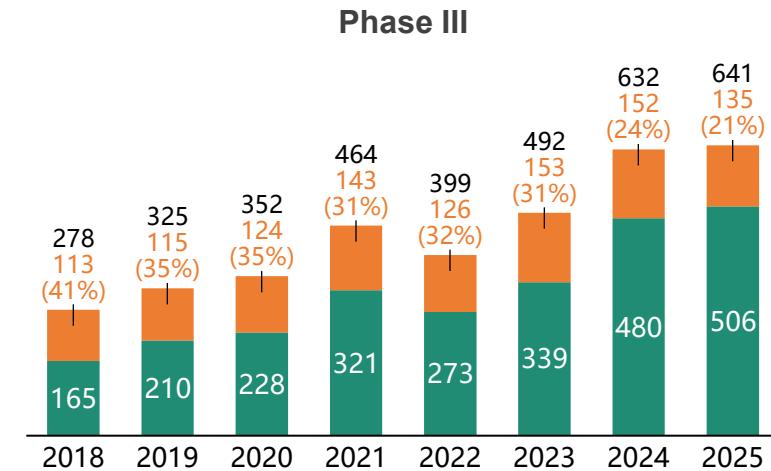
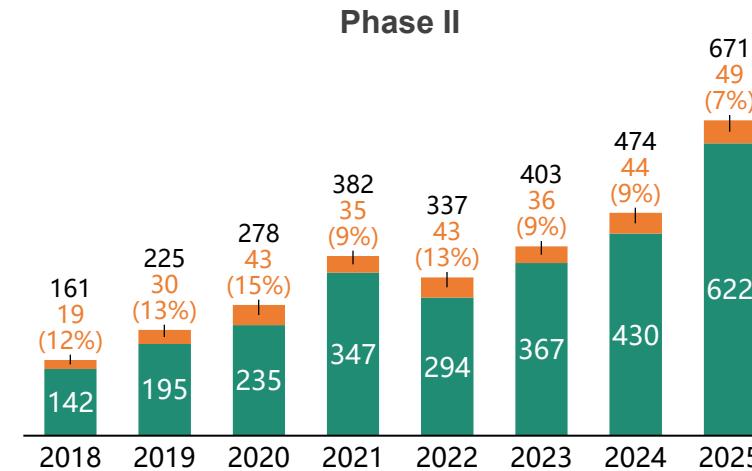


MNC is an Important Contributor of Innovative Drug Clinical Trials in China

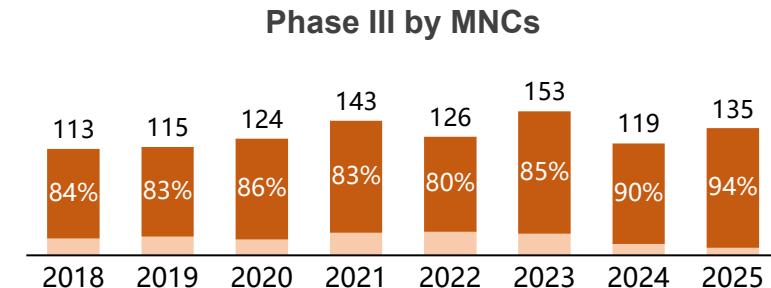
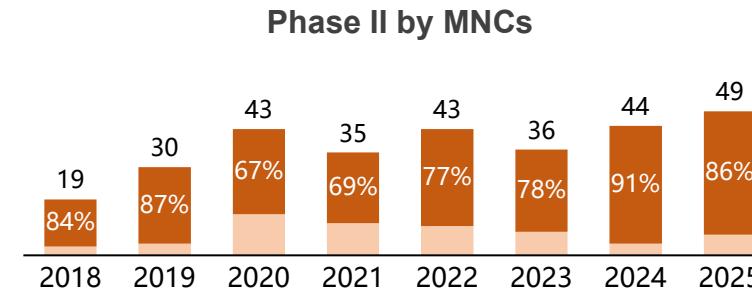
MNC is the major payer of China clinical CRO market with 25-30% of total Phase III trials by number

■ Newly started innovative drug clinical trials in China sponsored by MNCs ⁽¹⁾

■ Newly started innovative drug clinical trials in China sponsored by others



■ MRCTs (Multi-regional Clinical Trials)
■ China specific trials



Note: (1) Top 20 MNCs ranked by 2024 global revenue

MNC is Using More Local CRO Service Providers in China

Rapid development of local clinical trial industry

Improvement of service quality from local clinical CROs

Increase of clinical trial activities and peripheral services in China

- Local commercialization opportunity
- Large patient pool for clinical enrolment
- Localization of clinical trials (e.g. China-only trials and integrated evidence generation etc.)

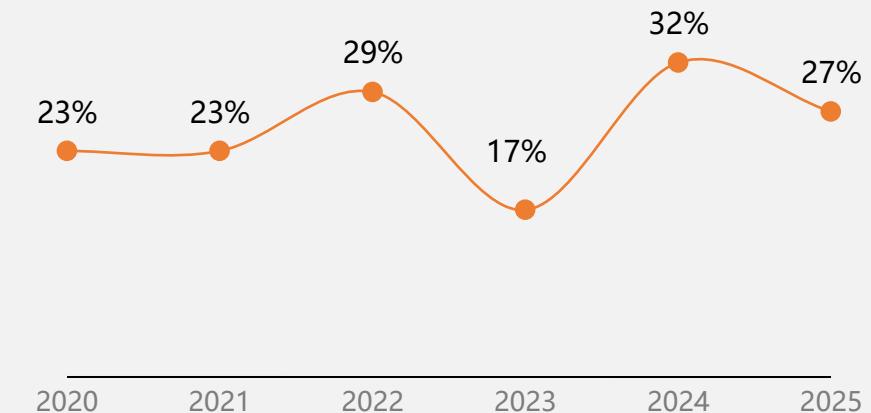
Better understanding of local regulatory environment

- Promulgation of local regulations (e.g. pharmacovigilance)

Needs for service vendor diversification and risk mitigation

Domestic CROs taking up market shares for MNC clinical operation projects in China

% — China CRO's share of MNC outsourcing projects

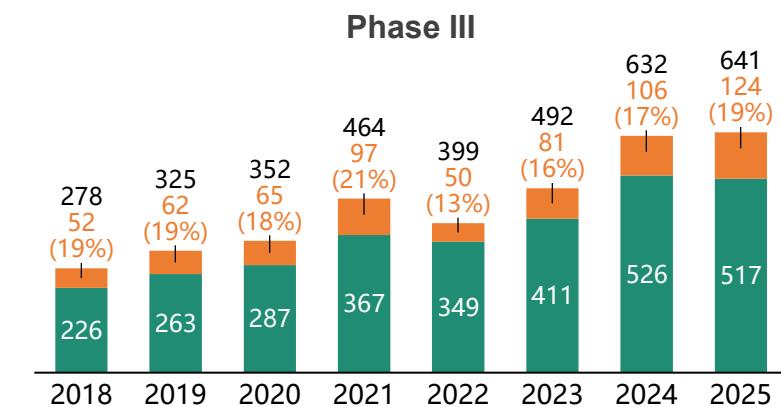
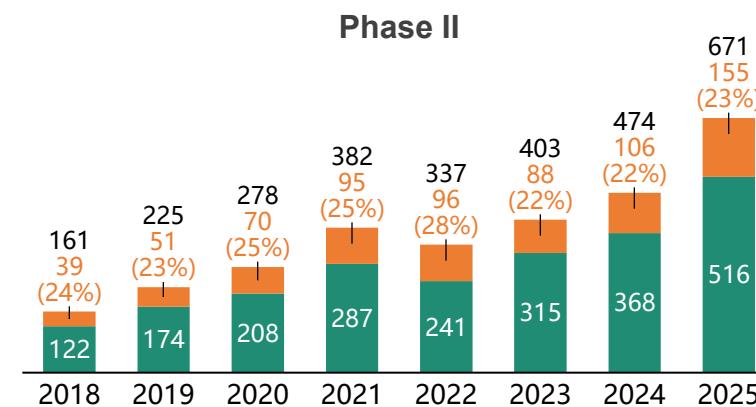


Local Pharma is Contributing More to Innovative Drug Trials in China

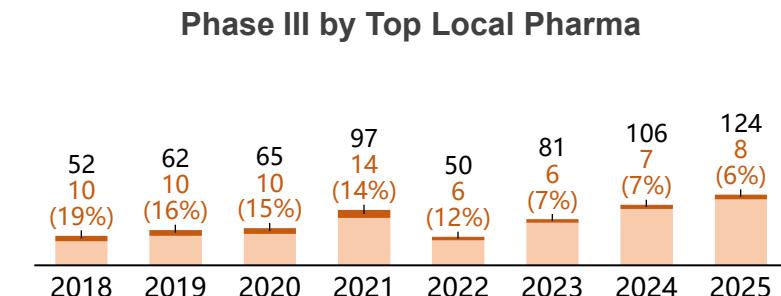
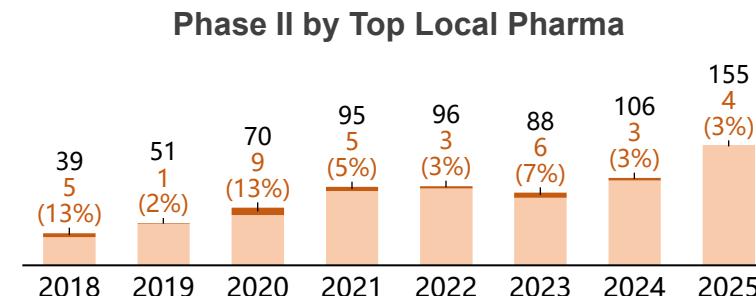
China local pharma (listed big pharma and emerging pharma) contributed 15-20% of total Phase III trials

■ Newly started innovative drug clinical trials in China sponsored by local pharma

■ Newly started innovative drug clinical trials in China sponsored by others



■ MRCTs (Multi-regional Clinical Trials)
■ China specific trials



Note:

Big pharma include Hengrui, Hansoh, Mindray, Fosun Pharma, Kelun, CSPC, Sino Biopharmaceutical, CR Pharma and Shanghai Pharma

Emerging biopharma include BeiGene, Innovent, Akeso, 3SBio, Kelun Biotech, Junshi, Remegen, SinocellTech, GenScript, ZaiLab, HutchMed, ChipScreen, Ascentage, MicroPort

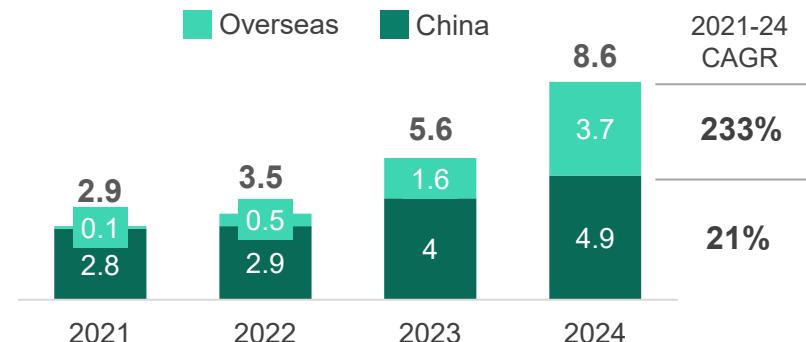
Local Pharma has Delivered and is still Growing its R&D Efforts

China-Originated Innovative Drug Sales Ramp-up⁽¹⁾

15+

China-originated innovative drugs reached
US\$100m+ sales in 2024

Sales of top 15 China-originated innovative therapies (global revenue in Billions USD)

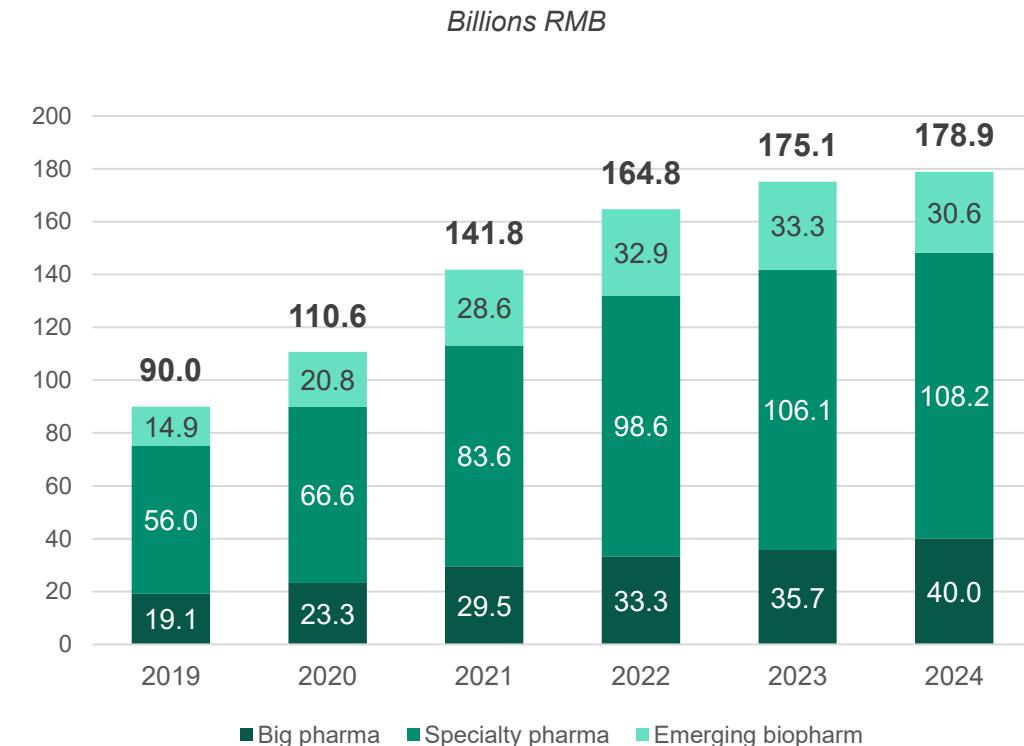


Average sales
(US\$ million)

of product
launched overseas

Note: (1) China-originated innovative drugs: China-originated assets only, not including licensed-in assets

China Local Pharma R&D Spending Continues to Grow

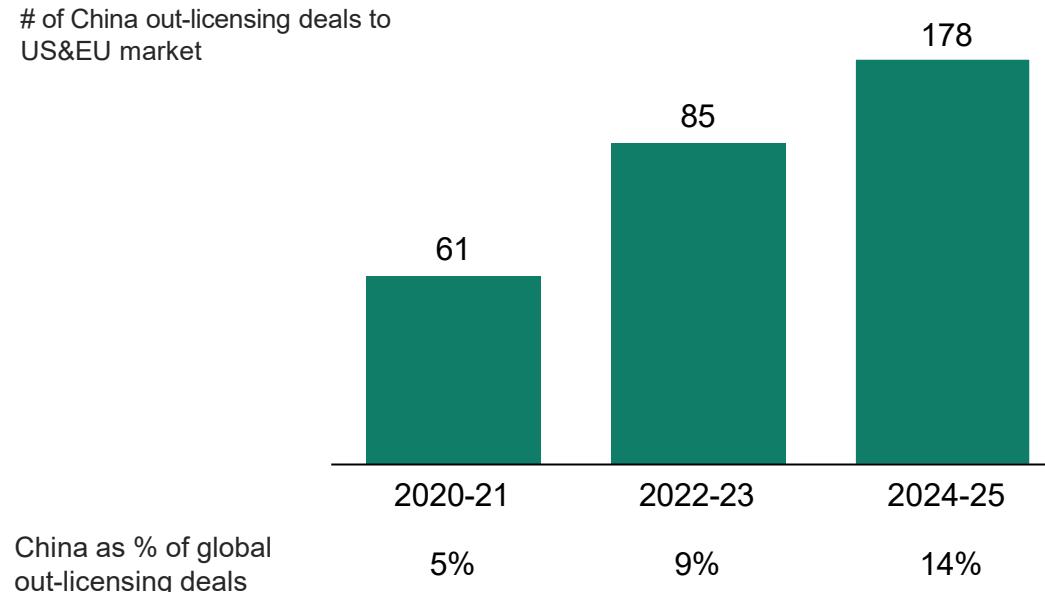


Note:

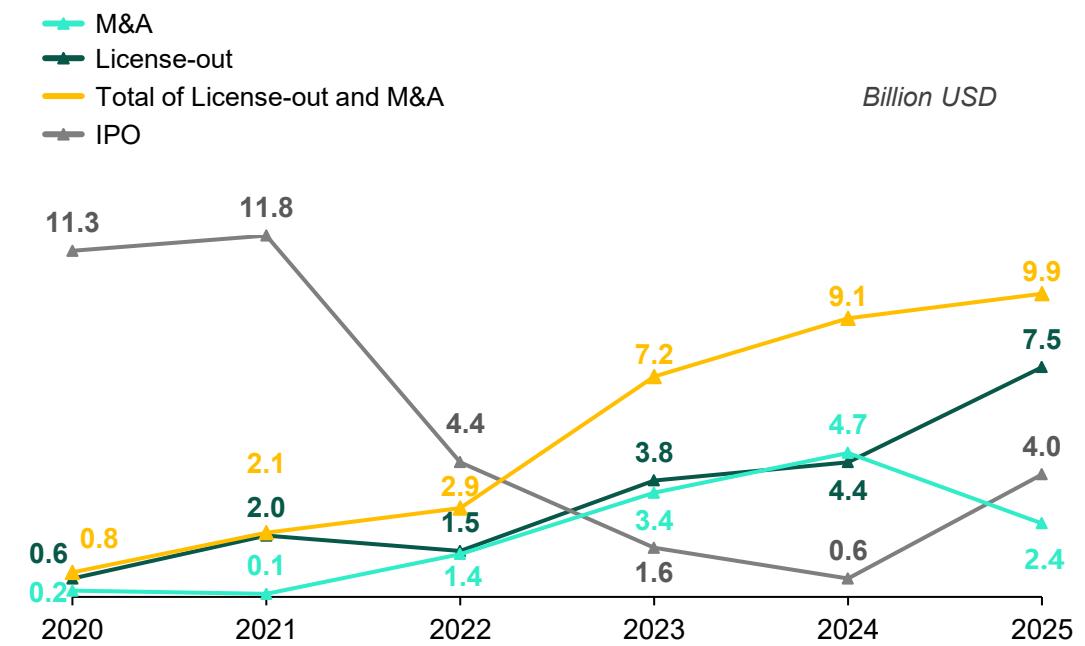
All public companies including pharmaceuticals, medtechs, vaccines and distributors
Emerging biopharma include BeiGene, Innovent, Akeso, 3SBio, Kelun Biotech, Junshi, Remegen, SinocellTech, GenScript, ZaiLab, HutchMed, ChipScreen, Ascentage, MicroPort

China Biotechs Received More Cash in 2024-2025 than 2020-2021 Mainly Because of Licensing-out and M&As Transactions

A significant increase of out-licensing deals of China-originated innovative assets to US&EU market ⁽¹⁾



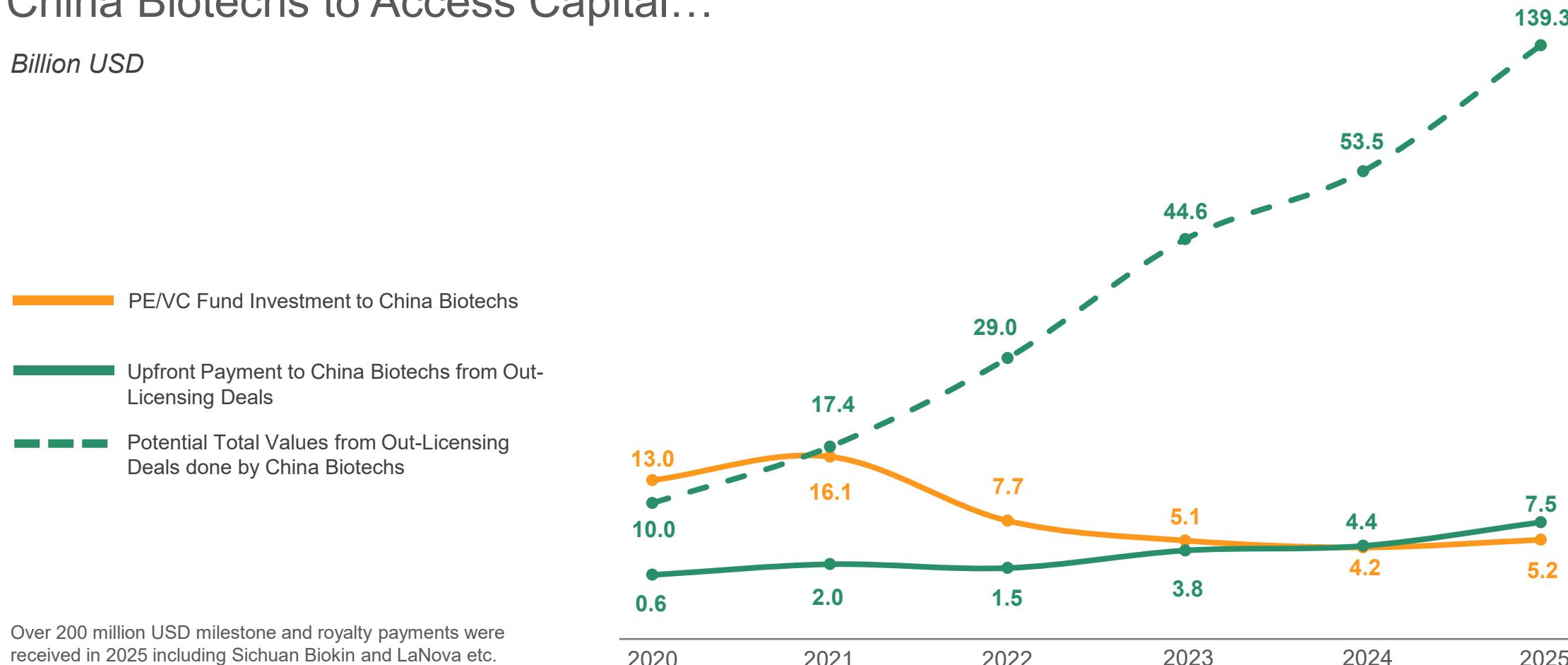
Assets-based transactions and M&As continued to be active while IPOs regaining momentum (by deal value)



Note: (1) Innovative asset deals (excluding generics and biosimilar) with China-originated licensors

Rising Upfront Payment Provides a Key Alternative for China Biotechs to Access Capital...

Billion USD



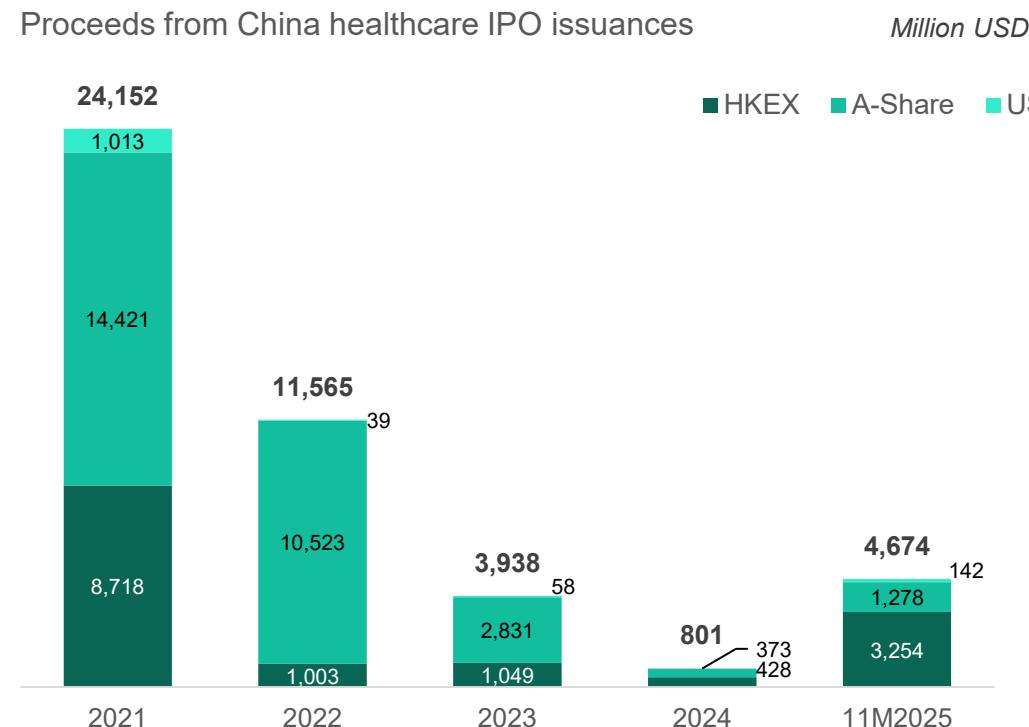
Over 200 million USD milestone and royalty payments were received in 2025 including Sichuan Biokin and LaNova etc.

May Not Be Exhaustive

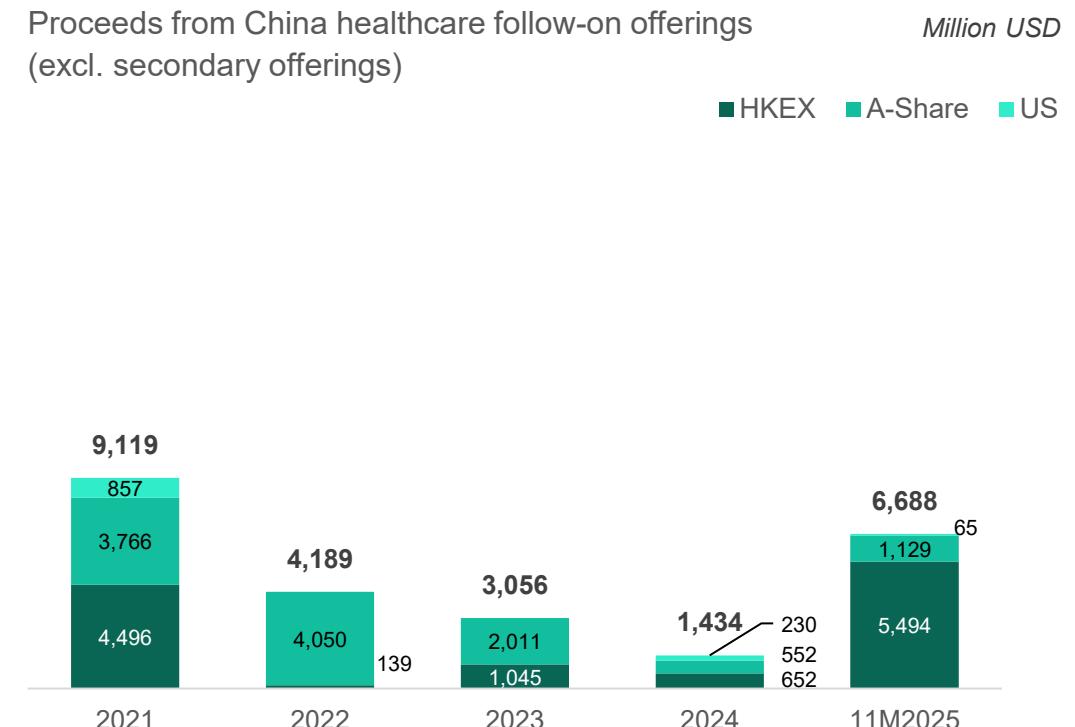
Note: China biotechs refer to China-originated biotechnology companies

...With Public Market Capital Raising Activities Revived in 2025

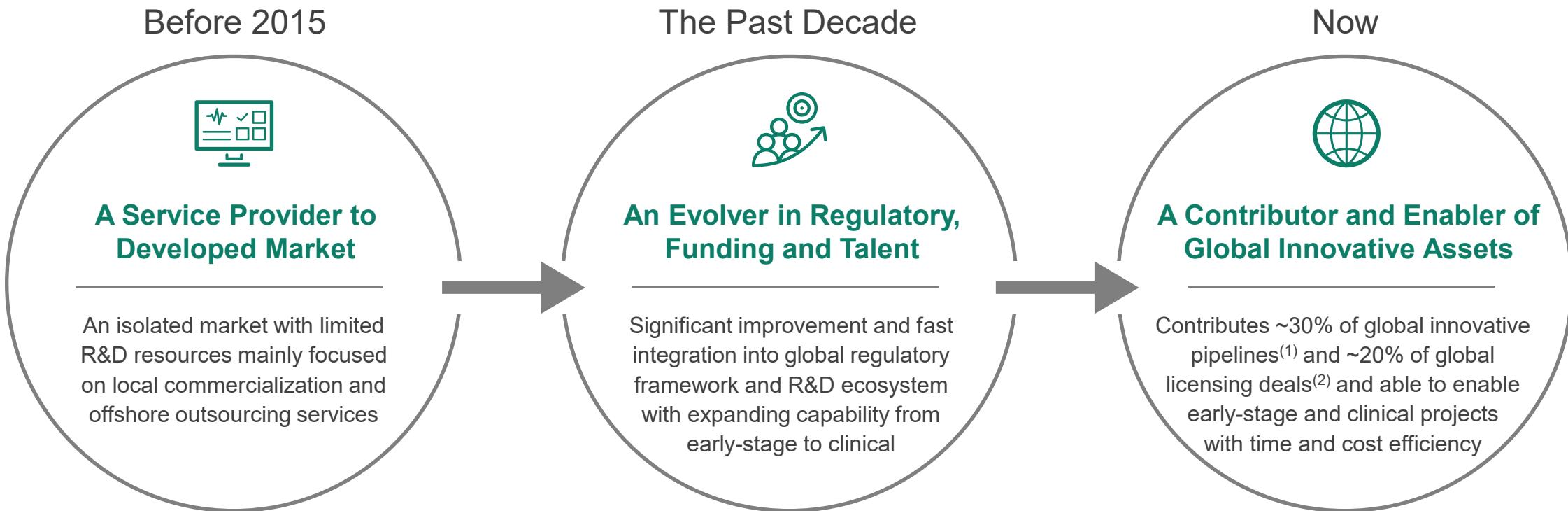
China HC Companies Raised ~20% More from IPOs in 11M2025 than 2023, Still Well Below 2021 Level Mainly Due to A-Share IPO Draught...



...While Following-on Offerings Saw Strong Rebound with HKEX FOs Well Exceeded 2021 Level and Most FO Issuers are Biotechs



China R&D Industry Plugged into Global Ecosystem...

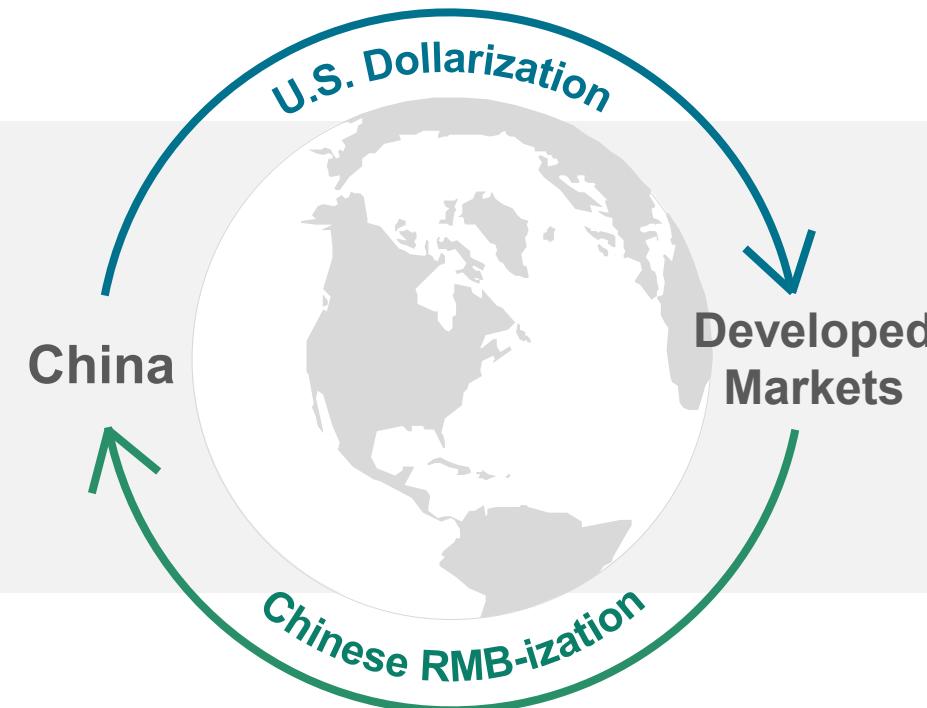


1. Including pipelines in phase 1, 2, 3, and under NDA review, excl. natural products, biosimilar and reformulation drugs

2. Innovative asset-based deals (excluding Gx and biosimilar) with licensor being China-originated companies and deal rights territory including USA/Europe market

...Followed by Value Realization

Innovative assets in China offer compelling value propositions
for US/EU companies/investors



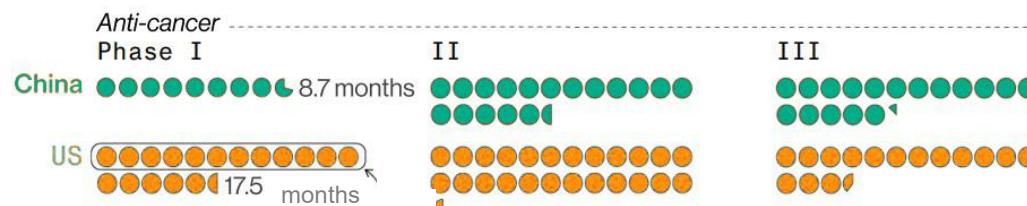
R&D services in China offer time and cost efficiency with
the same quality level

Huge Time Saving Potential by Leveraging China's R&D Resources

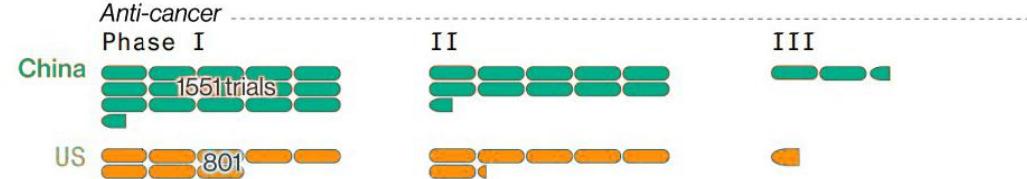
Higher Efficiency in China despite More Trials...

Data based on single-country trials conducted in 2020-2024

Median time to recruit patients for trials ○ = 1 month

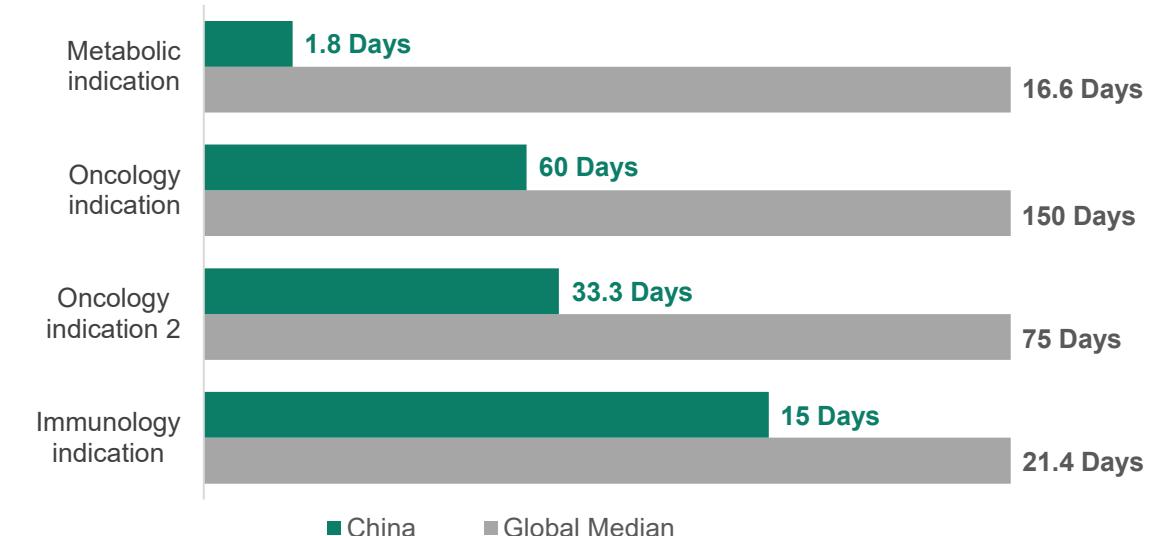


Total trial count ○ = 100 trials



...with 2-5 Times Faster in Patient Enrollement

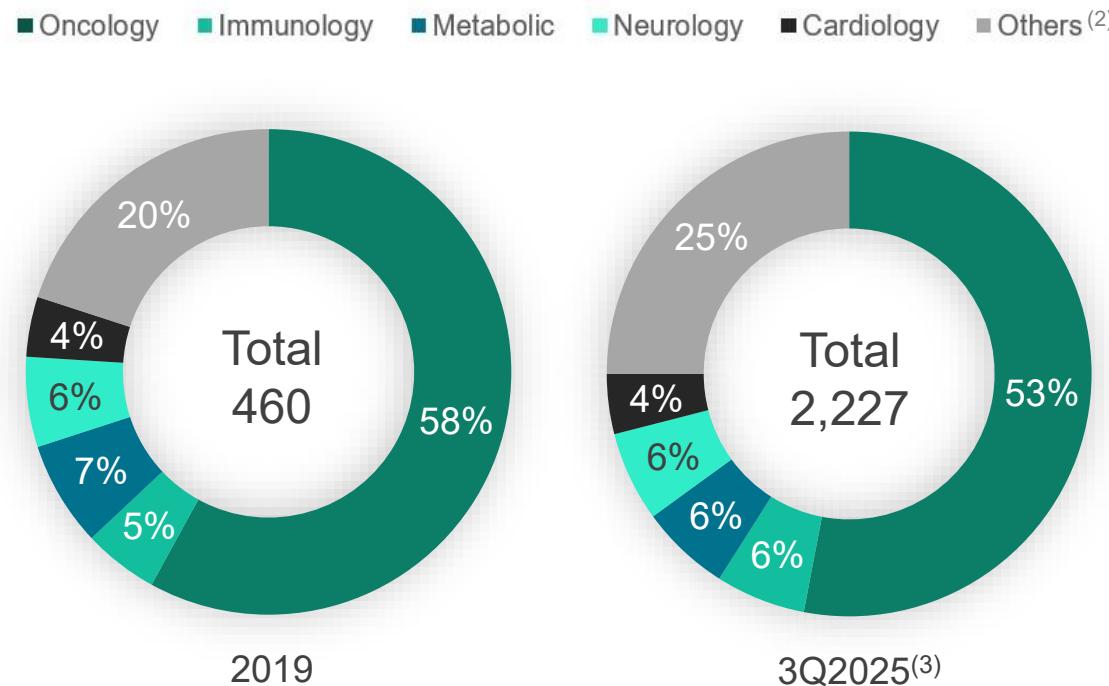
Average time to enroll a patient/site in Phase II or III trials by TAs



Illustrative And May Not Be Exhaustive

More Diversified TA Focus with Leading Position in Next-gen Modalities

China Innovative Pipelines⁽¹⁾ Breakdown by TA (total # and %)



1. Innovative pipelines from clinical phase I to pre-NDA. Excluding natural products, biosimilar, reformulation, and generic drugs

2. Including anti-infective, musculoskeletal, respiratory, dermatological, GU and blood-related etc

3. Year to date as of Sep 2025

4. CAR-T, TCR-T, Stem cell therapy and gene therapy, excluding RNA related therapies

5. siRNA, RNAi, and mRNA therapies

6. Innovative pipelines from clinical phase I-II. Excluding natural products, biosimilars, reformulations, and generic drugs

Early-stage Innovative Assets⁽⁶⁾ in Next-gen Modalities 3Q2025⁽³⁾



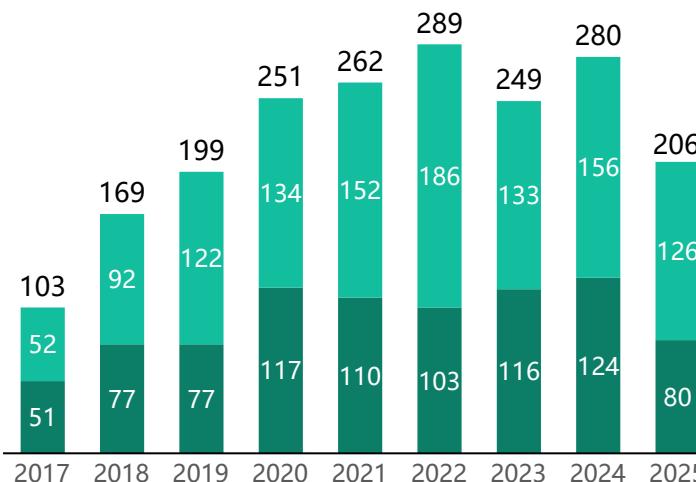
China Innovation Contribution to Global: Today and Beyond



Overseas Clinical Trials Initiated by Chinese Sponsors

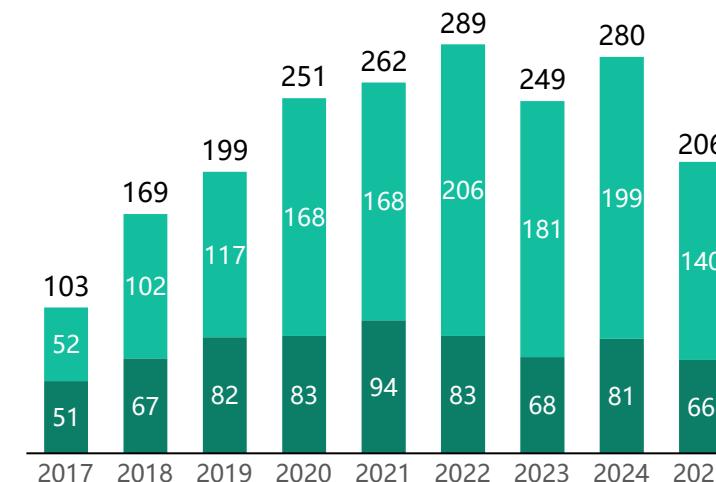
Breakdown by MRCT & Single Region 2017-2025⁽¹⁾

MRCTs Single Country



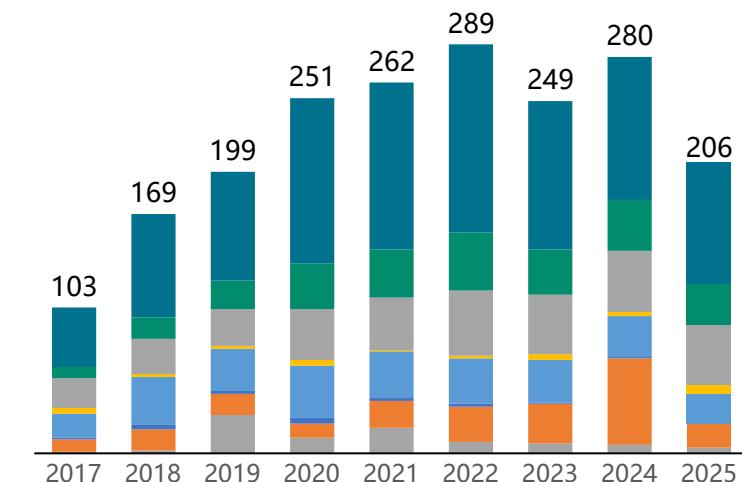
Breakdown by Pharma & Biotech 2017-2025⁽¹⁾

Pharma Biotech



Breakdown by Trial Phase 2017-2025⁽¹⁾

Phase I Phase II/III Others
Phase I/II Phase III BE
Phase II Phase IV



1. Excluding COVID-19 related clinical trials



PART 4

Tigermed Strategies



Stay Committed to Sustainability in 2025

ESG Commitment



MSCI ESG Rating

AAA

The highest rating in
the industry

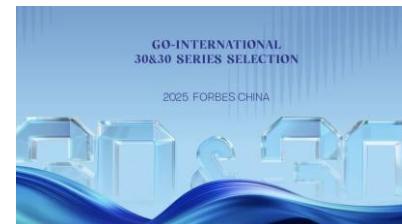


Morgan Stanley Capital International ESG
(Environment, Social, Governance) rating
received in Aug 2025

Global Recognition



Forbes China
Top 30
Go-International
Flagship Brands



Awarded by Forbes China for Go-
International 30&30 Series Selection
in Nov 2025

Overseas Expansion



Acquired
Micron
A Japanese CRO
Company



Enhance Tigermed coverage across
Japan and Asia Pacific in medical
imaging capabilities.

Domestic Leadership



Market Share

No.1

in China Clinical
CRO Market



Frost & Sullivan
Clinical Outsourcing Service Market
Report 2025

Tigermed Strategies & Outlook for 2026 and Beyond

Build a higher moat on our core clinical and related services and improve our relative competitiveness

01

Build closer ties with various stakeholders, partly through our role as an early stage industry investor, and actively participate in policy making and consultation process

03

Enhance our global service capability and quality consistence and better built our critical mass in key overseas markets

04

Monitor inorganic growth and consolidation opportunities in both China and overseas markets

06

Strategically strengthen our business relationship with MNCs and local pharma in China

02



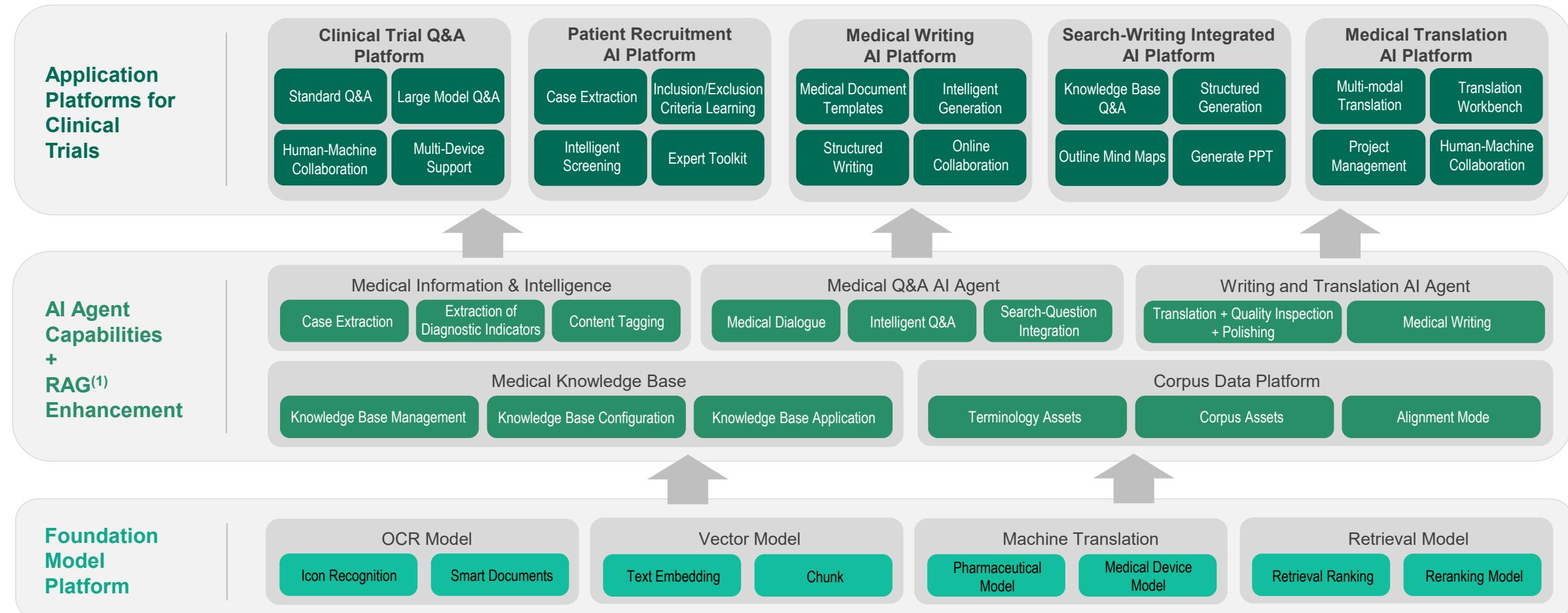
Continue to expand our emerging services and invest in new technology, AI and digital platform

05

Cope with industry cycle trough with operational resilience, profitability & cash flow focus and cost control, and reflect upon the previous cycle

07

Tigermed AI LLM for Intelligent Clinical Development



1. RAG: Retrieval-augmented Generation

Case Studies: Delivering Efficient Clinical Trials with AI Technology

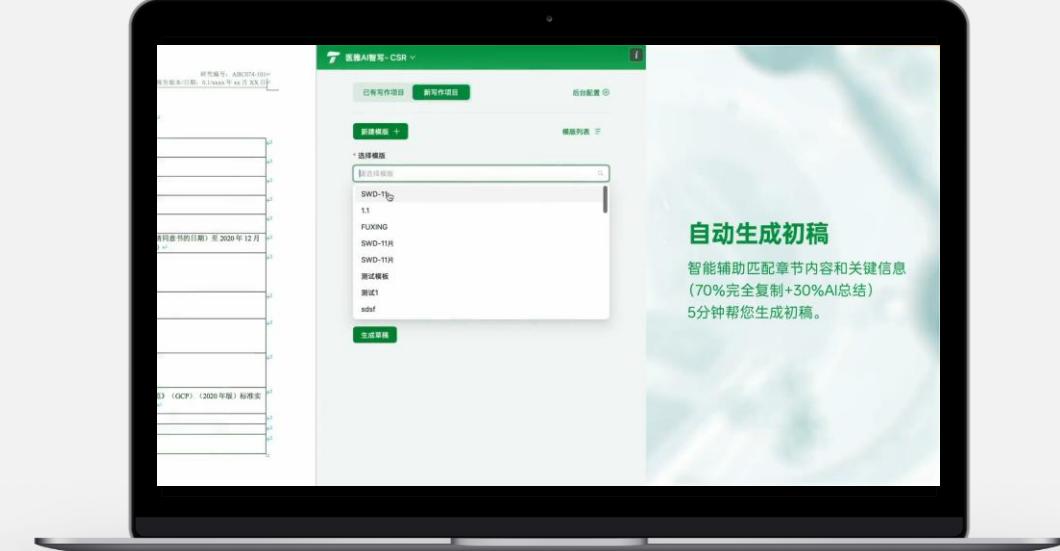
AI Translation (developed by Tigermed)

An AI Translation Platform developed by Taya (a Tigermed company) for accurate translation with multi-language and multi-model



AI Medical Writing (developed by Tigermed)

An AI Medical Writing Platform developed by Taya (a Tigermed company) to generate Clinical Study Reports (CSR) draft within 1 hour





J.P. Morgan Healthcare Conference 2026

Hangzhou Tigermed Consulting Co., Ltd.

300347.SZ / 3347.HK

Global Headquarters

F18, Building A, Shengda Science Park, 19 Jugong Road, Binjiang District, Hangzhou, China

International Headquarters

1201, Li-Ning Building, 218 Electric Rd, North Point, Hong Kong, China

www.tigermedgrp.com