

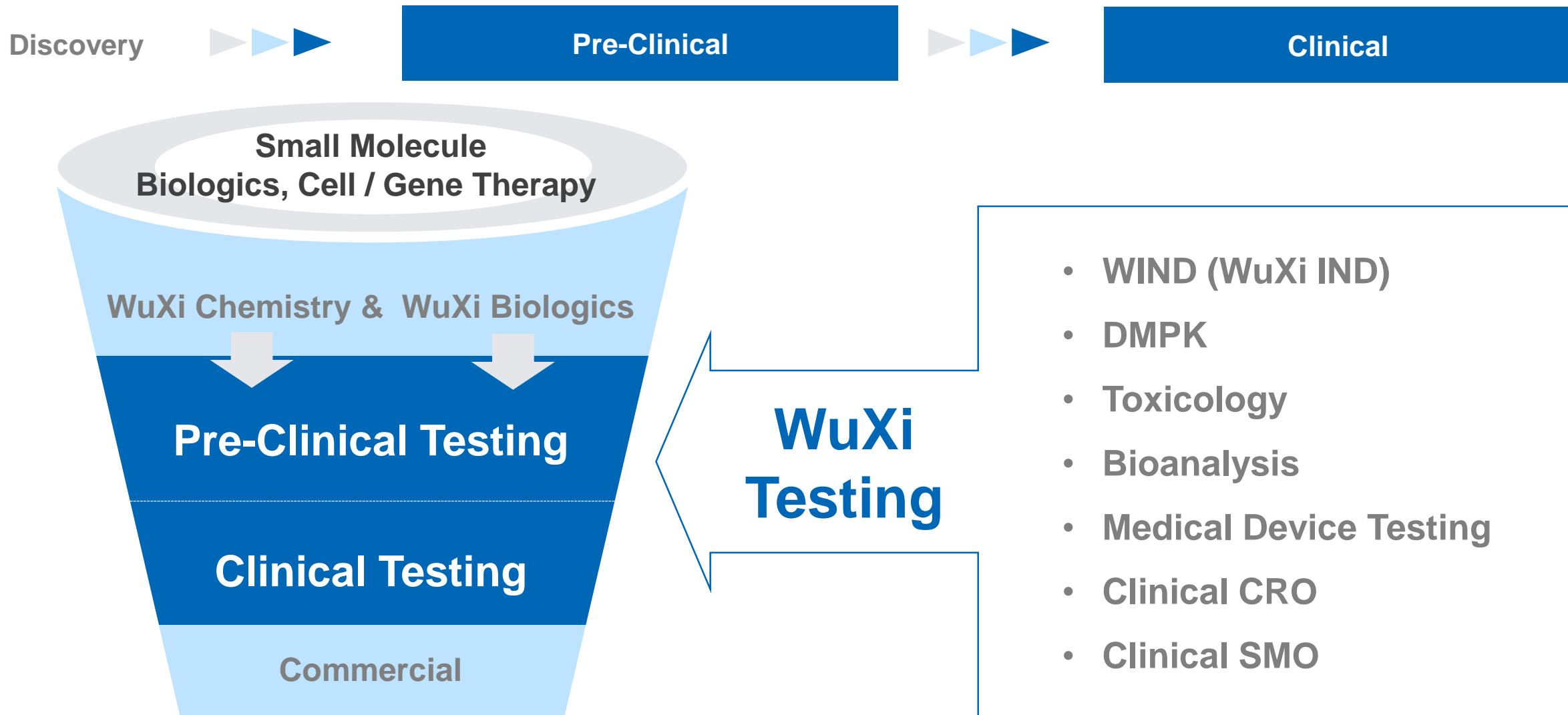


2022 WuXi AppTec Investor Day

WuXi Testing: End-to-End Testing Platform

Steve Yang, Ph.D.
Co-CEO

WuXi Testing: End-to-End Testing Platform for All Modalities



WuXi Testing: A Global Platform from Preclinical to Clinical

Employees

9,300+

Clients¹

2,200+

Projects²

8,800+

- **3,500+ Pre-clinical**
- **5,800+ Clinical**

- **2,100+ Pre-clinical**
- **800+ Clinical**

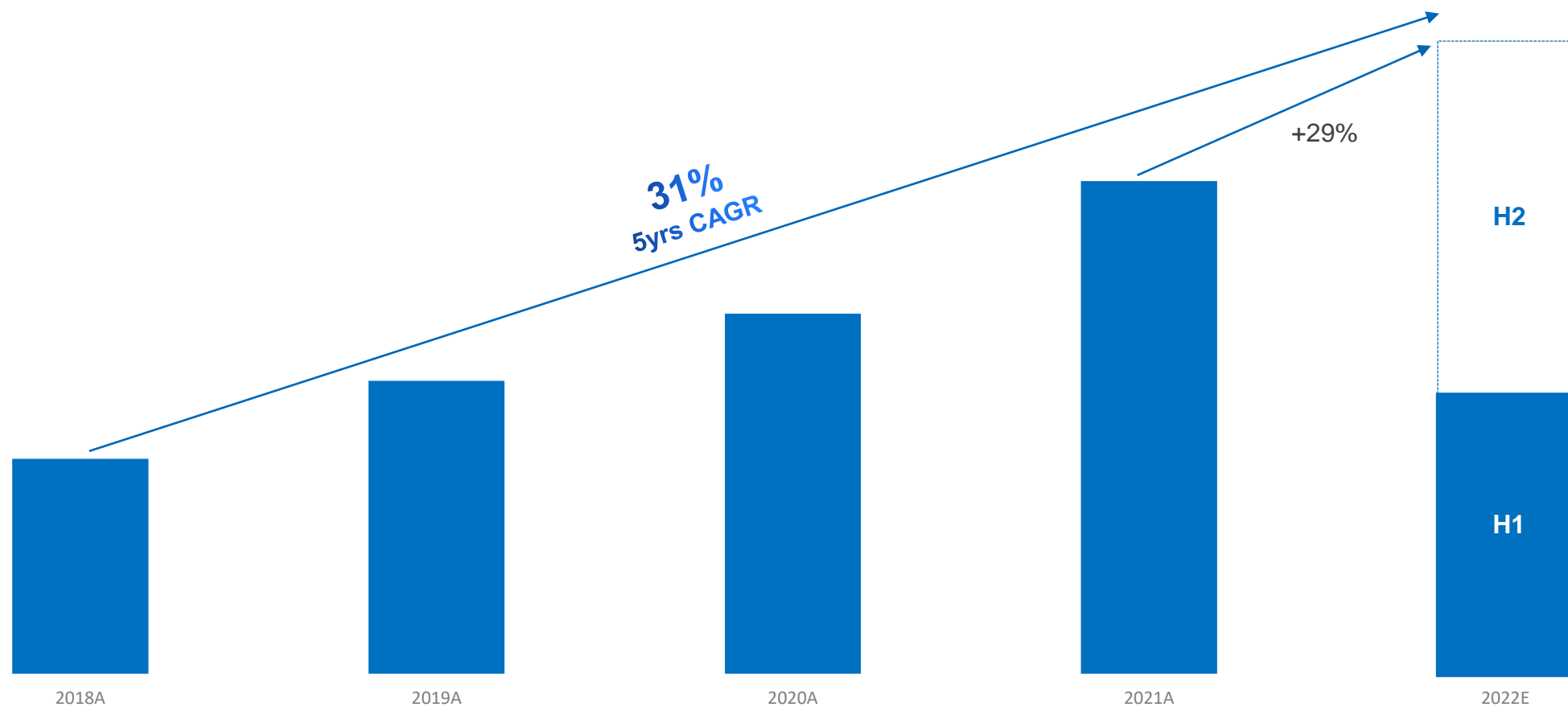
- **5,500+ Pre-clinical**
(Including 140 WIND)
- **3,300+ Clinical**

Note: 1. Active client with PO award

2. Active projects with PO award (Multiple projects may cover single molecule)

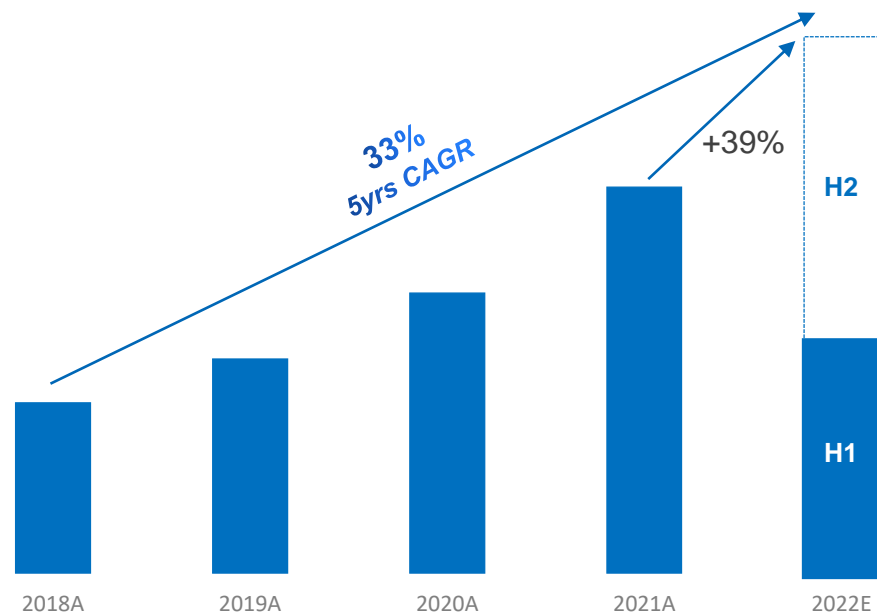
Data from: Jul 2021 to Jun 2022

WuXi Testing: Robust Track Record & Growth Forecast

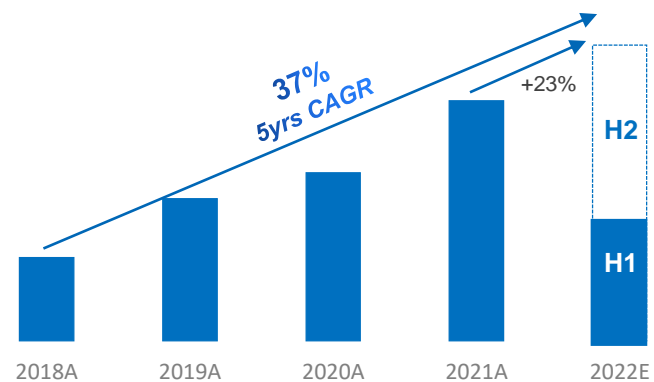


WuXi Testing: Business Performance by Division

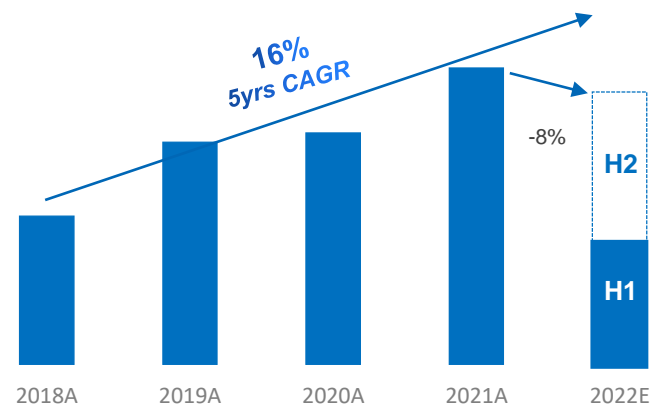
Lab Testing Business



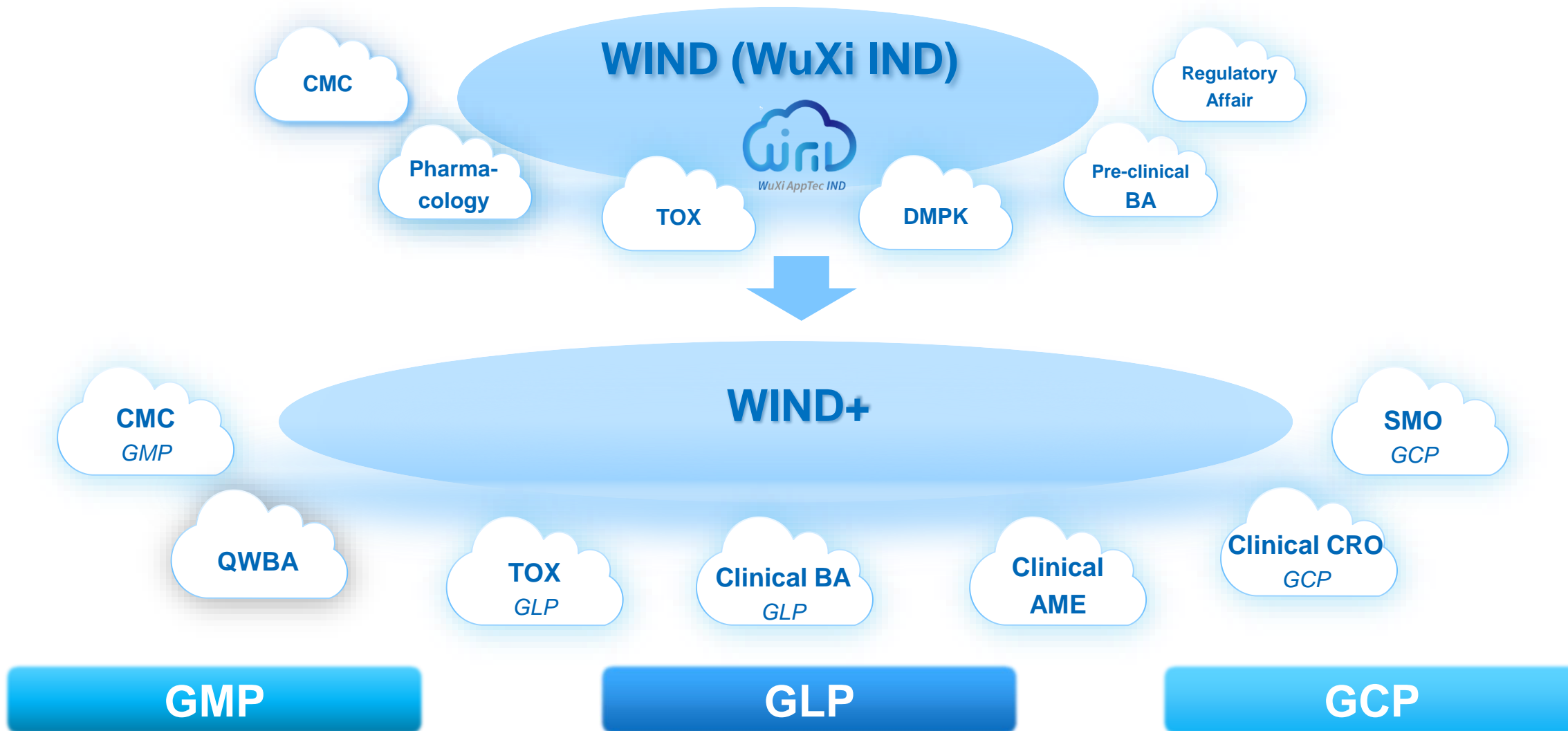
SMO Business



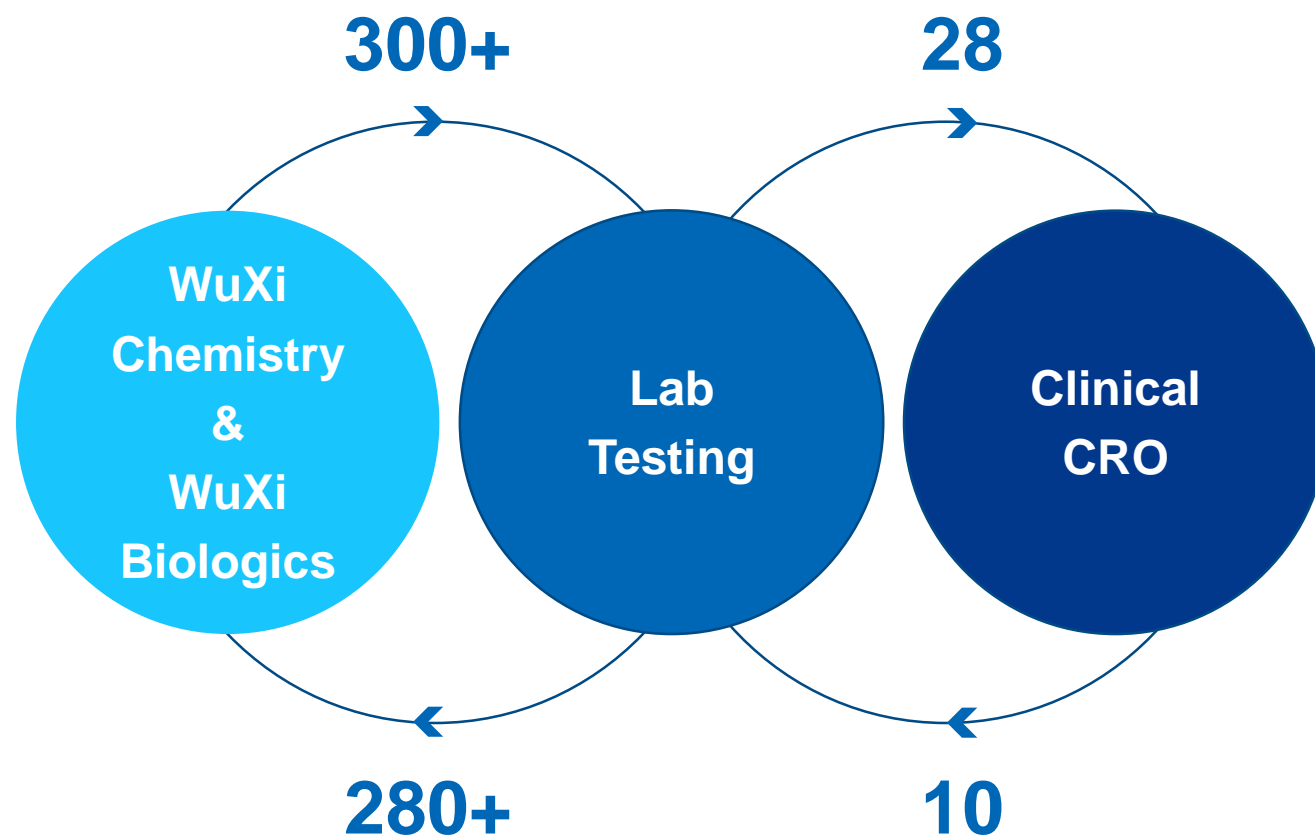
WuXi Clinical Business



Follow the Molecule: From WIND to WIND Plus

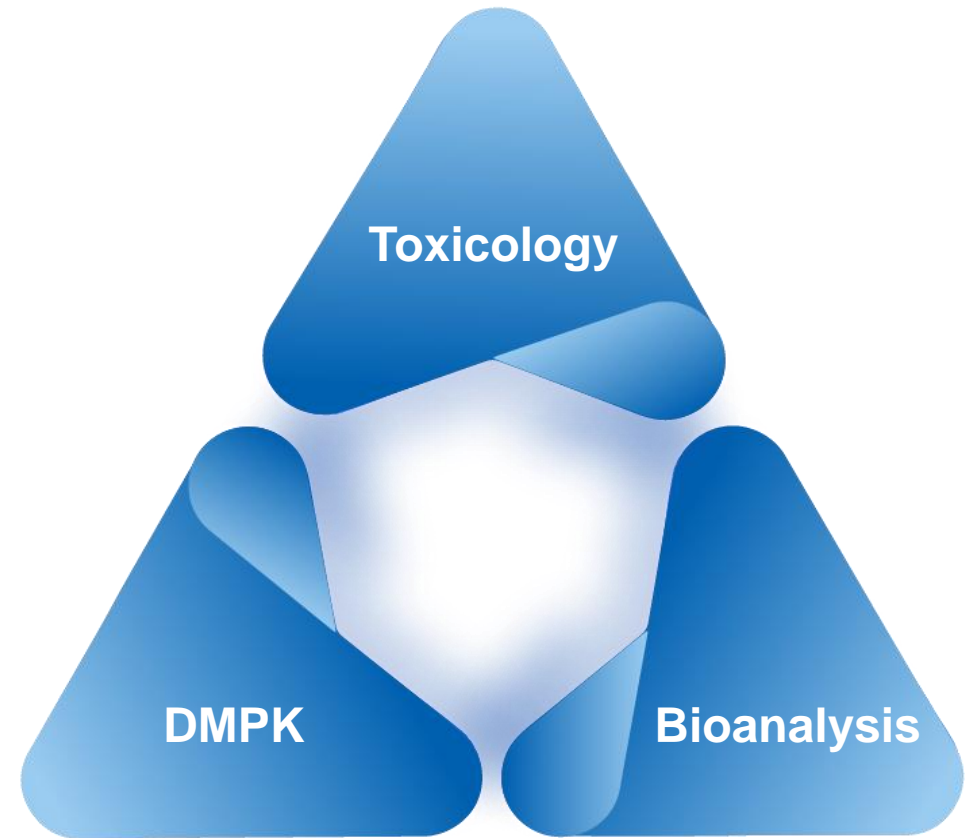
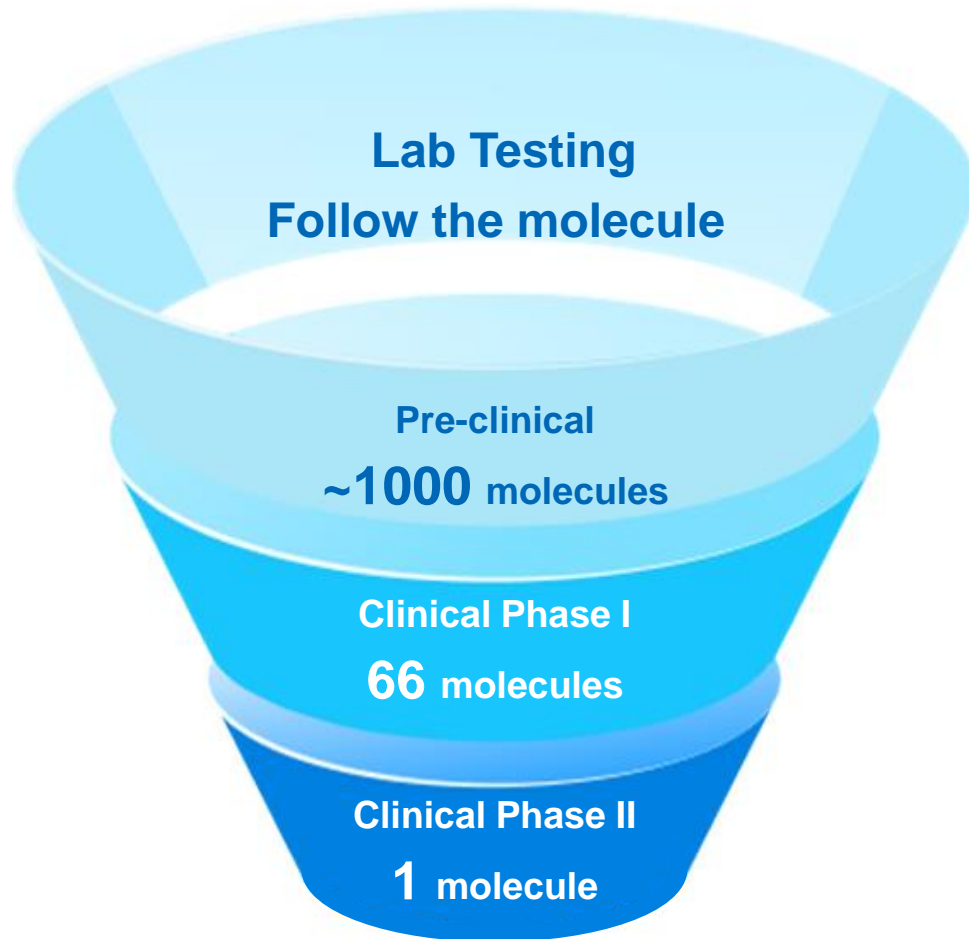


Follow the Molecule: Drive Customer and Project Conversion



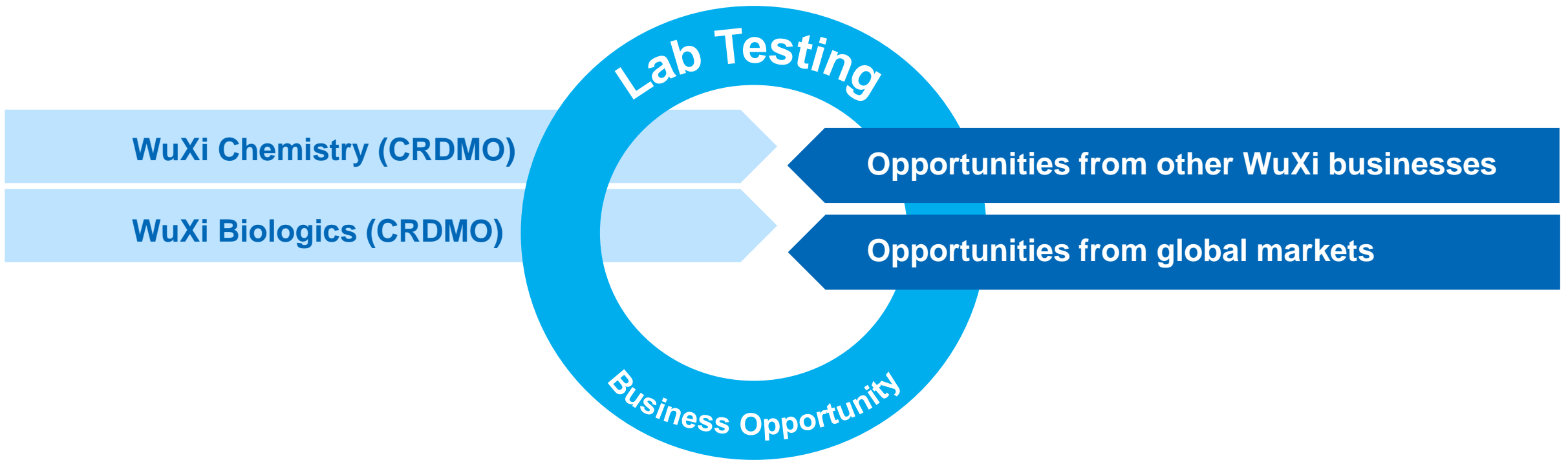
Data from: Jul 2021 to Jun 2022

Pre-Clinical to Clinical Conversion within Lab Testing



Data from: Jul 2021 to Jun 2022

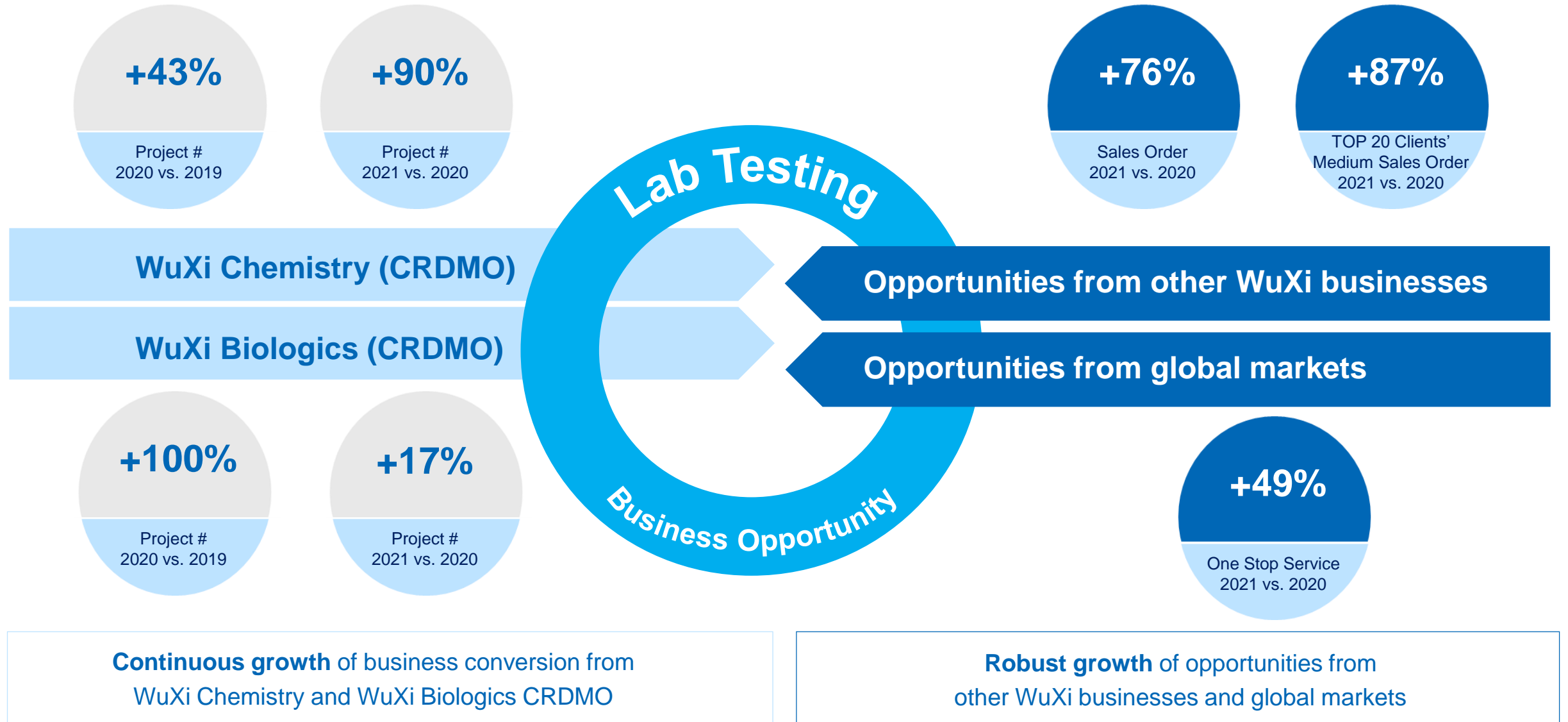
Lab Testing: Capture Opportunities from Multiple Sources



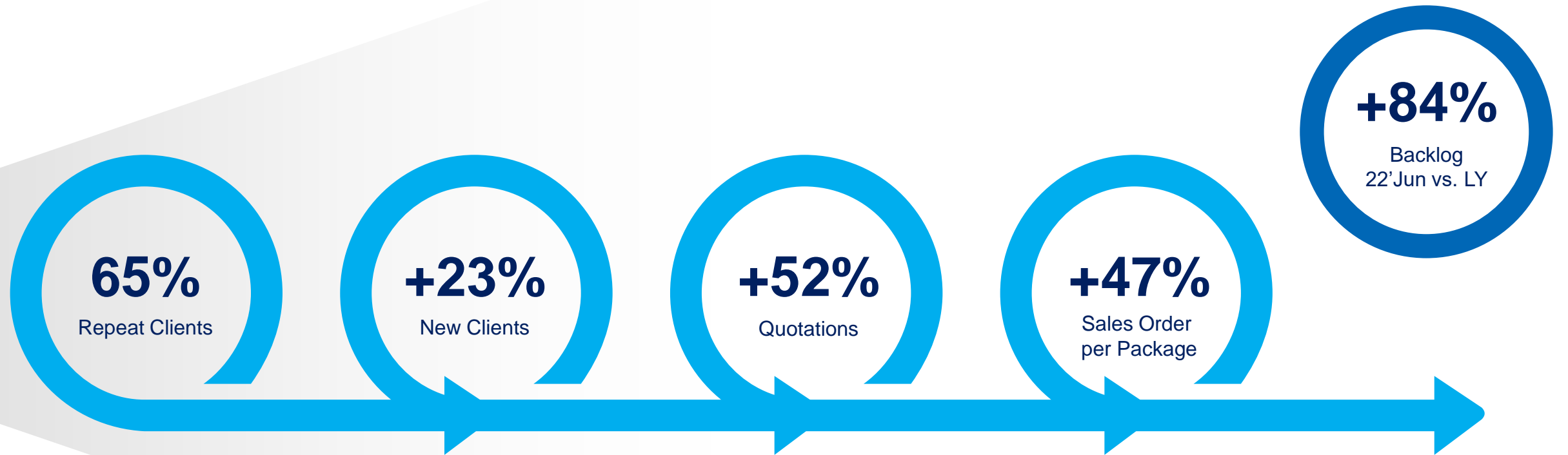
Continuous growth of business conversion from
WuXi Chemistry and WuXi Biologics CRDMO

Robust growth of opportunities from
other WuXi businesses and global markets

Lab Testing: Robust Project Inflow



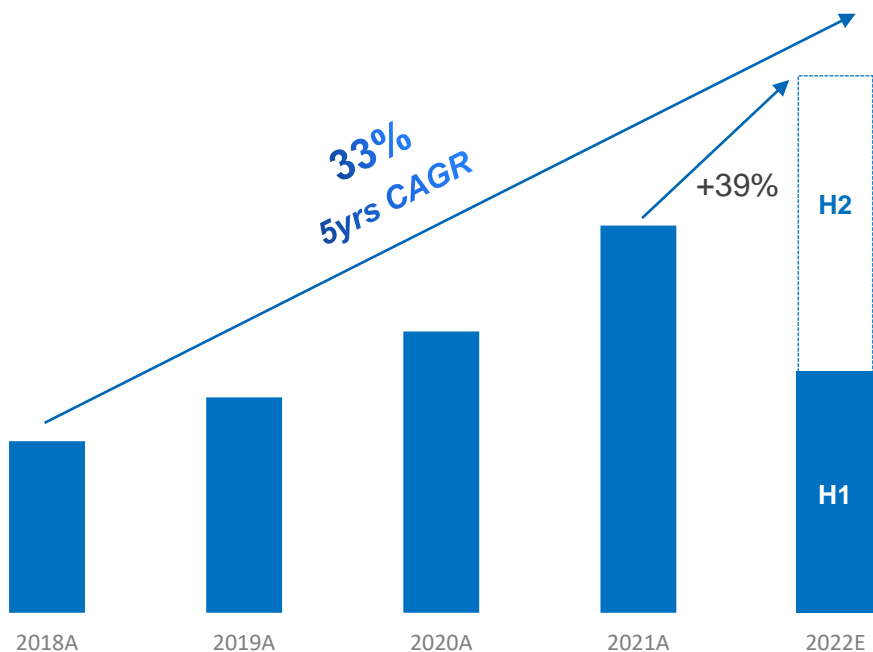
Lab Testing: Demonstrated Growth Momentum



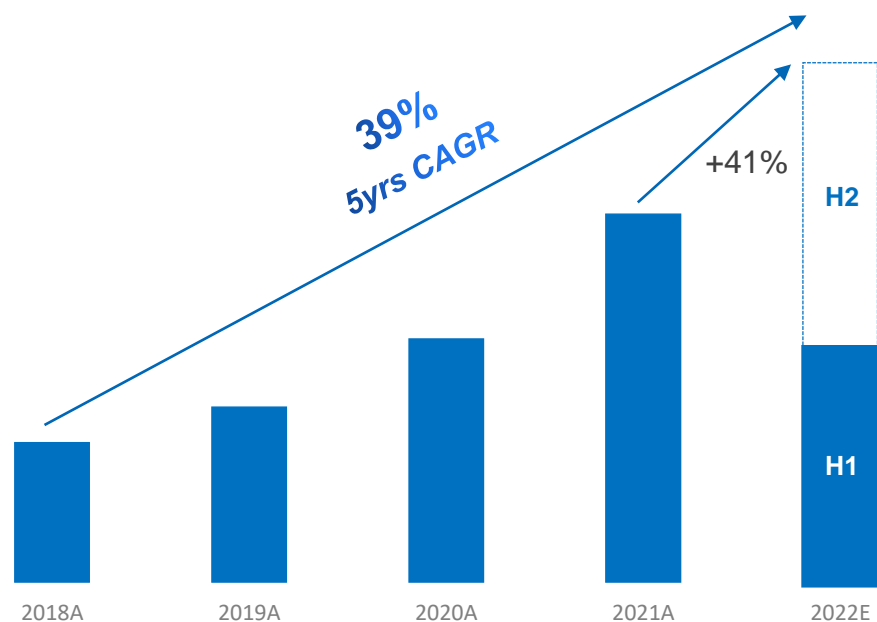
Data Compare: (Jul 2021 ~ Jun 2022) vs. (Jul 2020 ~ Jun 2021)

Lab Testing: Strong Track Record and Growth Forecast

5 Years Growth of Lab Testing Business



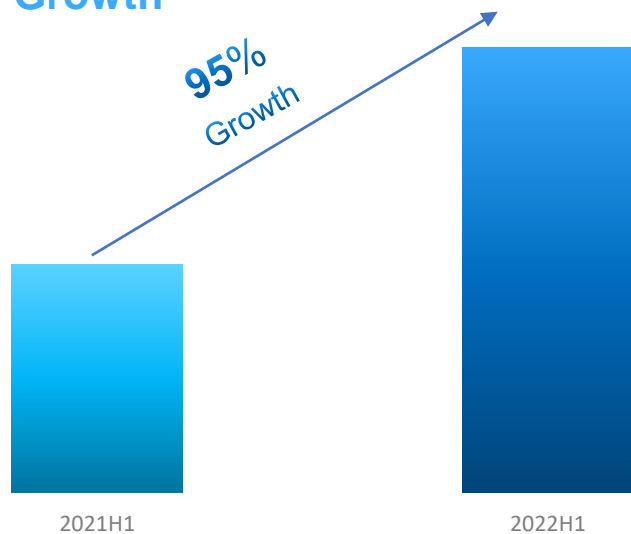
5 Years Growth of Drug R&D Business



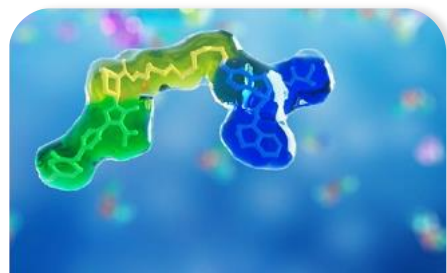
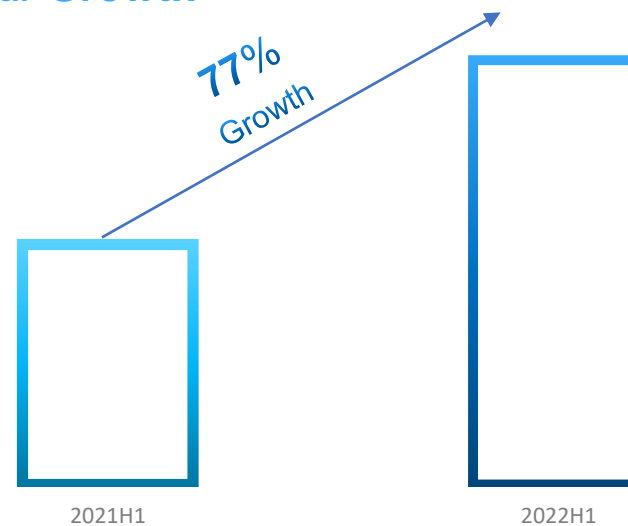
Lab Testing: Fast Growth of New Modality Business

Capability Build-up and Project Growth for New Modalities

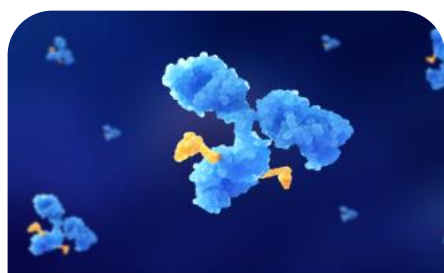
- Revenue Growth



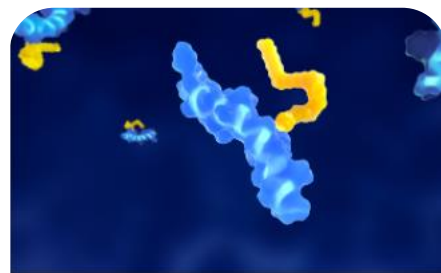
- Project# Growth



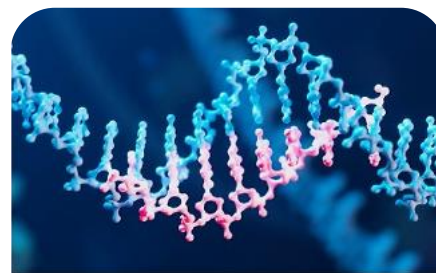
Protein Degradation



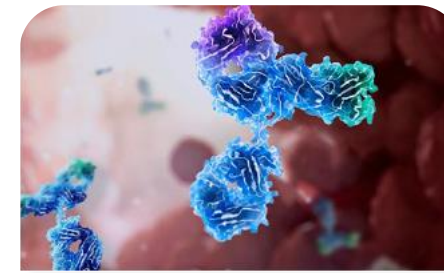
Antibody-drug Conjugate



Peptide



Oligonucleotide



Therapeutic protein

Lab Testing: Global Footprint with Increasing Capacities

China

100,000 sq.m. in use Capacity



Shanghai



Suzhou



Nanjing



Chengdu



Guangxi



Guangzhou

USA

25,000 sq.m. in use Capacity



Atlanta, GA



Cranbury, NJ



Plainsboro, NJ



St. Paul, MN



Qidong



Suzhou

Construction in Process

55,000 sq.m.

Up-coming GLP Testing & Preclinical Center: Qidong

Will Launch in 2023 Q2



GLP
Testing Lab

40,000 m²
Lab Space

GLP Certification
in 2024

Full Capacity
in 2025 Q4

Up-coming GLP Testing Lab: Suzhou

Will Launch in 2023 Q2

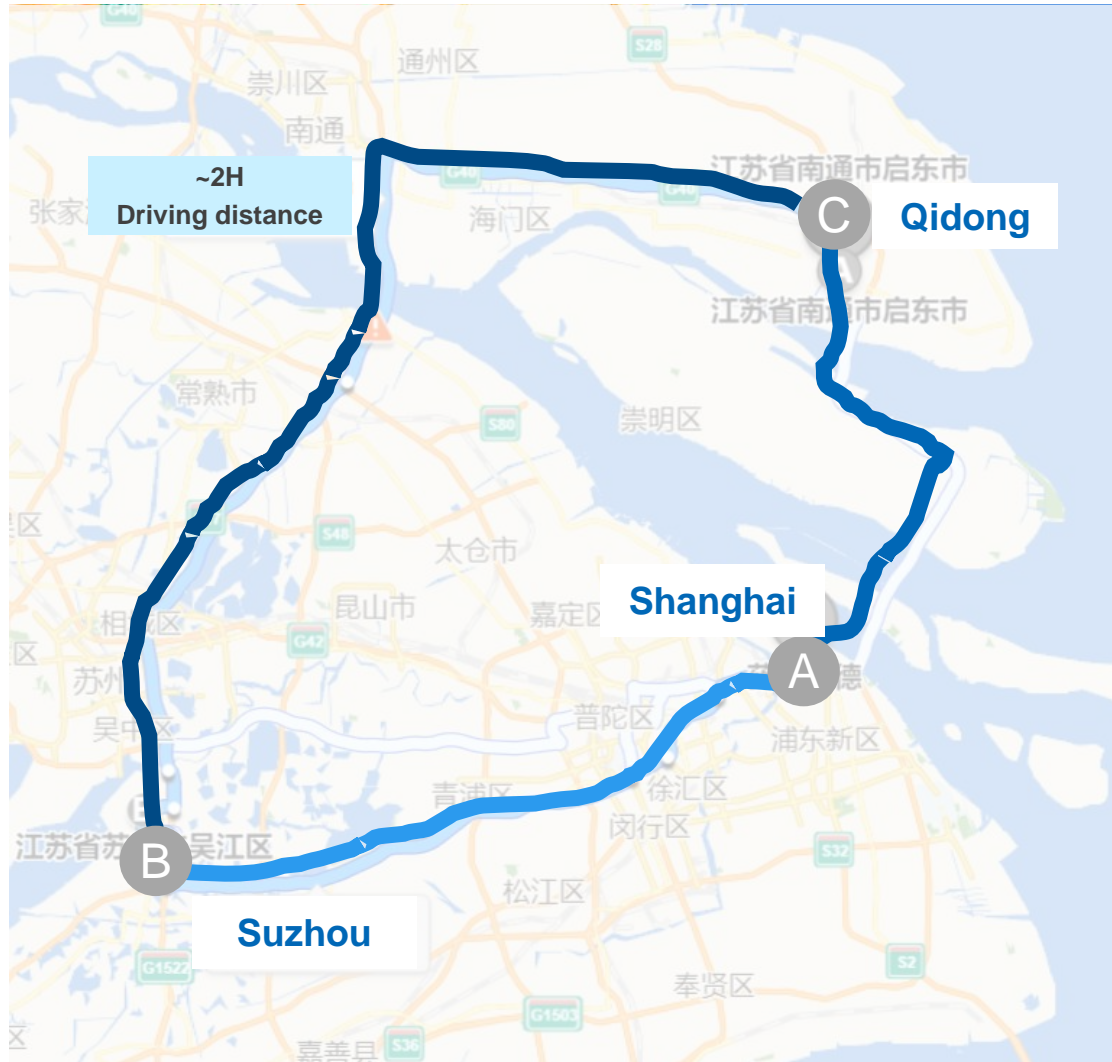


GLP
Testing Lab

15,000 m²
Lab Space

Full Capacity in
2024 Q2

Close Proximity Increases Efficiency and Resilience



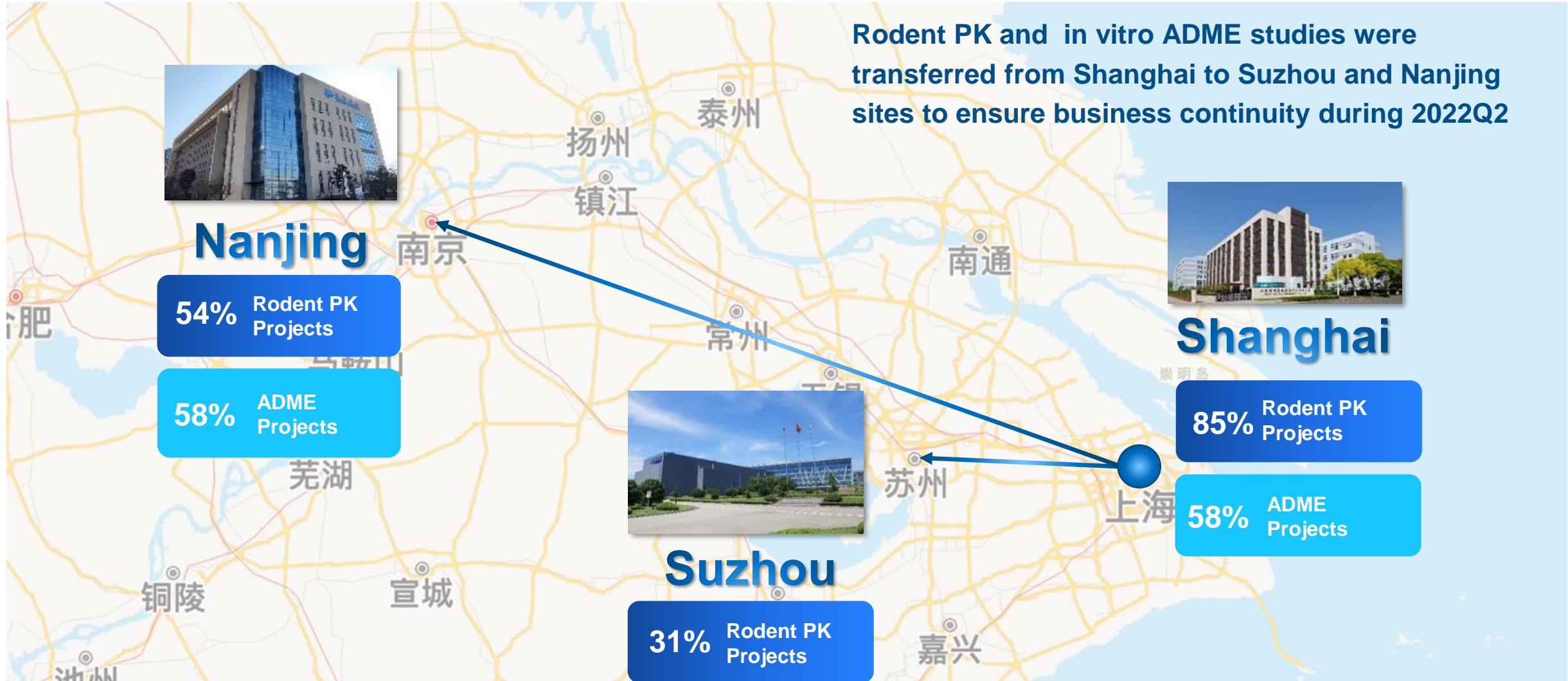
Within 2 hours driving distance

The same quality standard

The same operation system

Keep Business Sustainable and Flexible

Business Continuity Plan (BCP) Leveraging Site Network



AI-based Lab Testing: Fast, Accurate and High Quality

- **AI-based** animal room, laboratory and staff scheduling system (time to schedule a study reduced from few hours to few minutes)

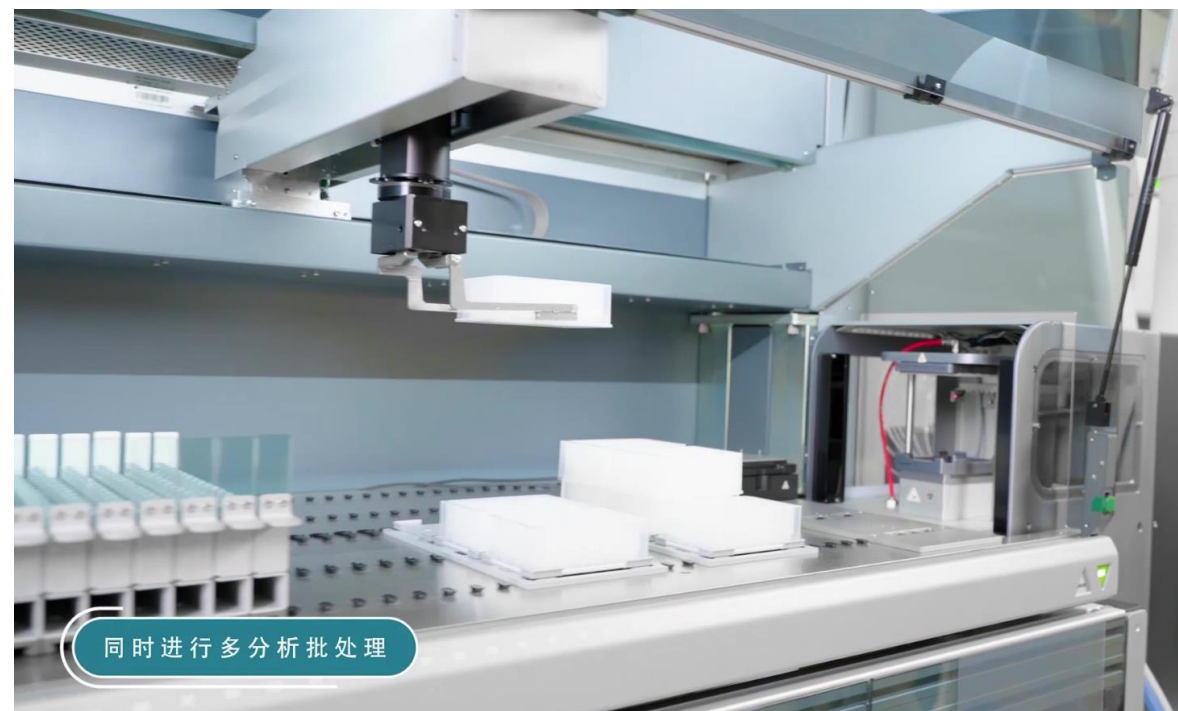
1. Intelligent Animal Facilities



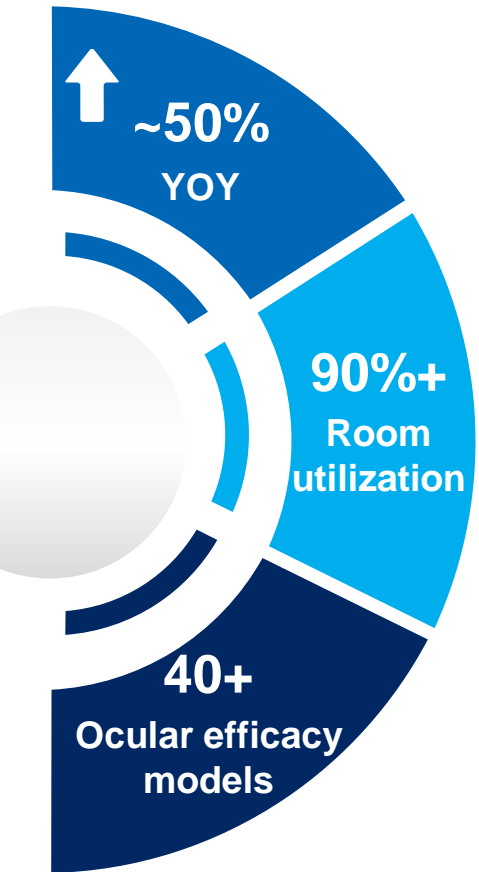
2. End-to-end Automated Labs



3. Automated Bioanalysis



Toxicology: A Growth Engine with High Quality



Comprehensive Toxicology Platform

- 56% CAGR Growth 2018-2022E
- Demand continues to be strong and outsourcing continues to accelerate in early discovery research
- Gene Therapy studies increased 400%
- A strong record of global GLP compliance

Toxicology: Substantial Backlog and Satisfied Customers

Strong Momentum Business

2,000M RMB

Backlog will convert to revenue in 2023

Molecule Conversion

13

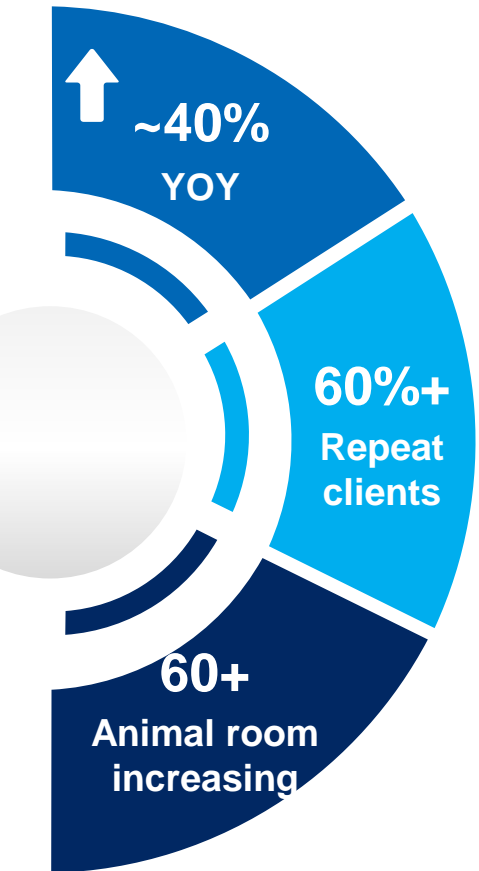
Molecules from pre-clinical to NDA long term TOX packages

Repeat customers

50%+

repeat business from existing customers

DMPK: Fast-Growing with Strong Synergy with WuXi Chemistry



Advanced end-to-end integrated platform

- Business from new modalities will maintain strong momentum
- Capacity continues to release fast in Suzhou and Qidong sites
- Long-tail customer strategy ensures a strong and growing customer base
- Strong synergy with WuXi Chemistry
- The radioactivity experimental platform has been enriched with world-class QWBA research capability

DMPK: Digital and Automation Technology Drive Efficiency

Digital Project Management

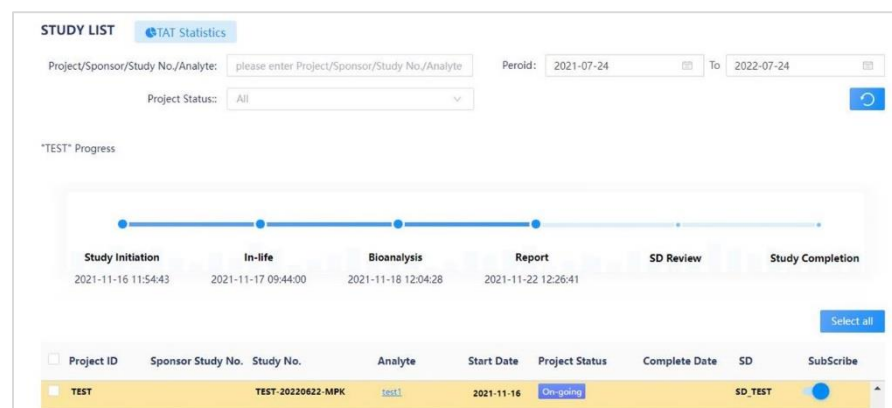
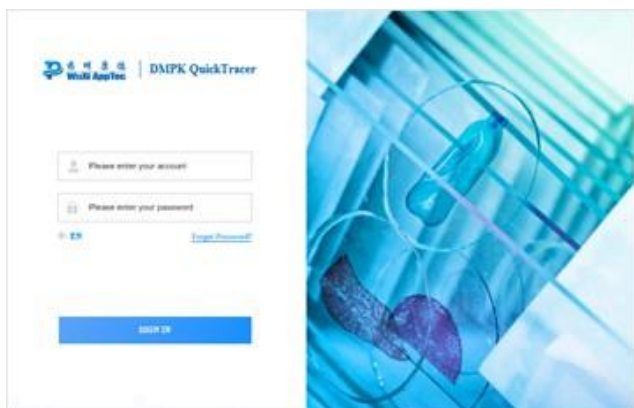
This online QuickTracer tracking system displays project progress in transparency and real-time, which ensures great customer experience and high efficient project management.

End-to-end Automated Labs

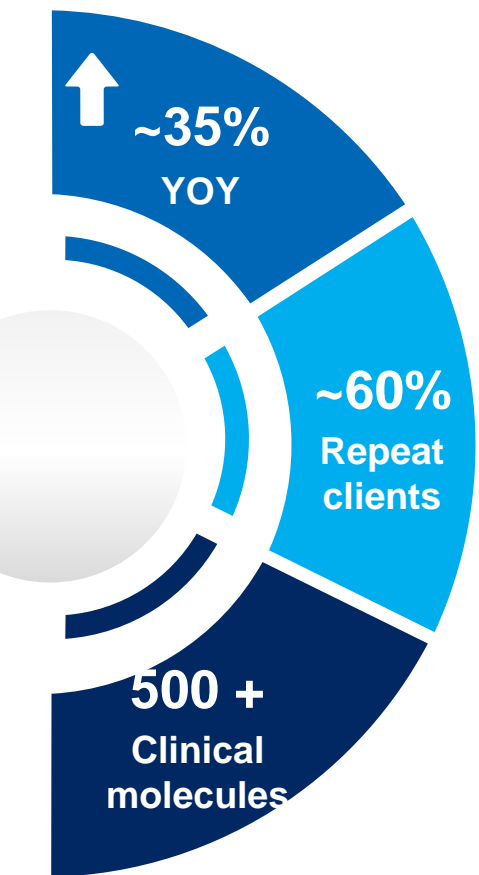
From compound management, lab testing to report writing, the highly automated DMPK labs ensure great quality and high efficiency

Intelligent Animal Facilities

The intelligent animal facilities realize the visualization of the study process and mistake prevention of experiment operation



Bioanalysis: Follow the Molecule to Drive Growth



One stop solution to cover the entire drug R&D life cycle

- Strong differentiators on bioanalytical capabilities lead to repeat business from existing customers
- “Follow the molecule” from preclinical to clinical
- “Win the molecule” through WuXi Chemistry CMC services, WuXi Biologics and XDC
- Keep strong record of global GLP compliance
- Competitive advantage of global operation to support global clinical trials

Bioanalysis: High Efficiency and High Quality

Laboratory E-Platform to Enhance Efficiency and Accuracy

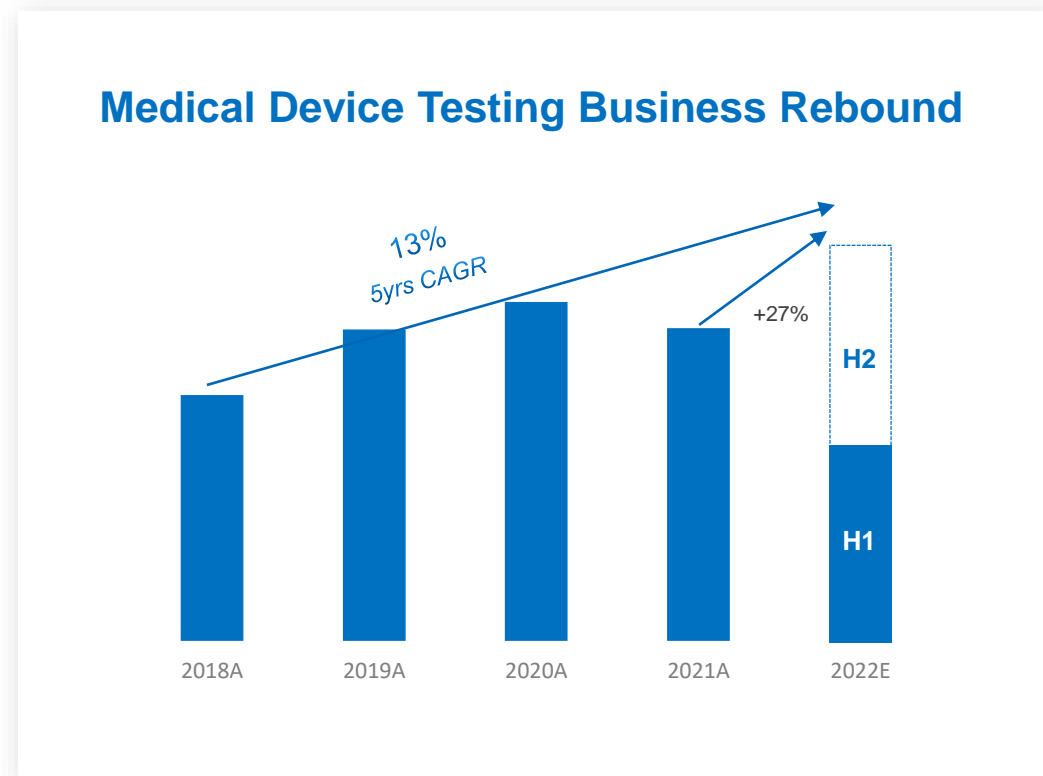
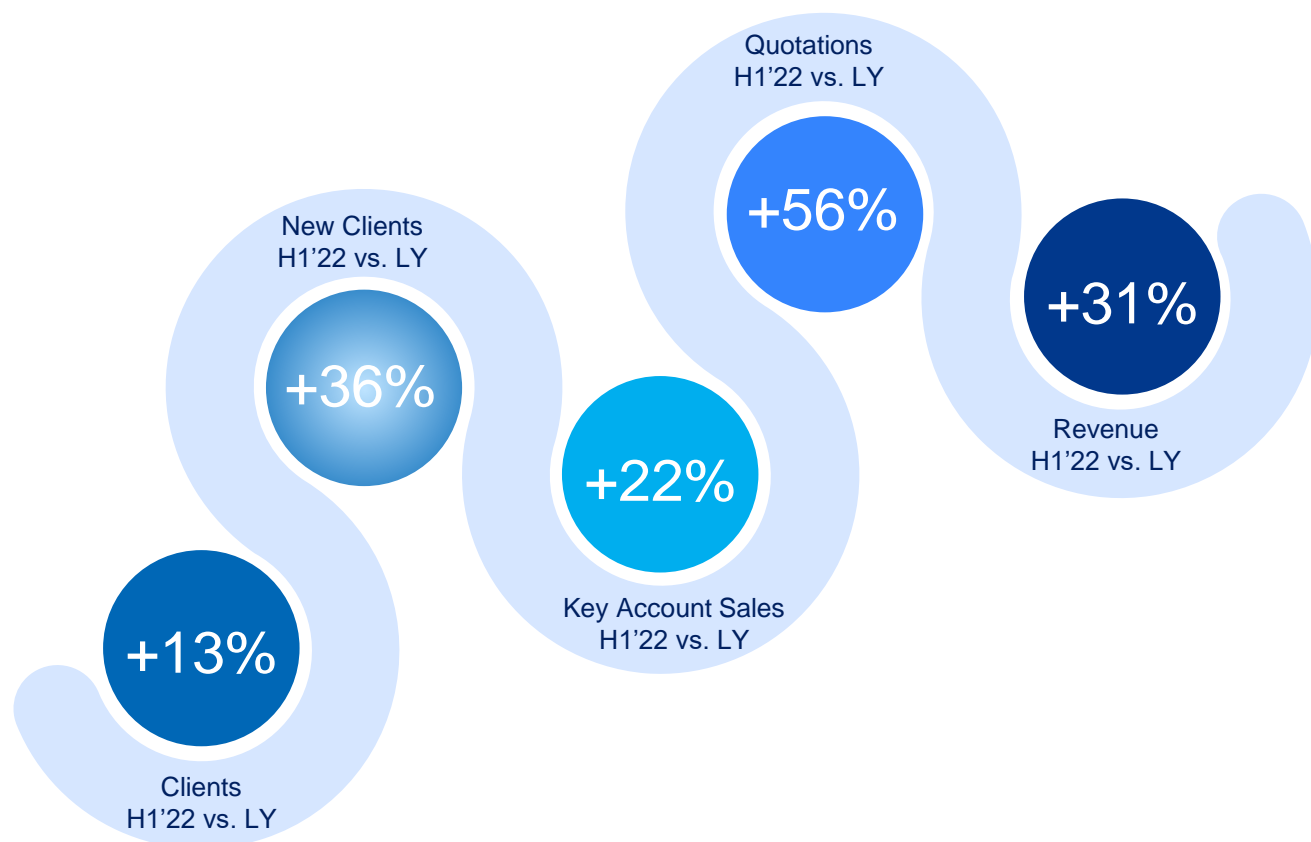


High Quality

Remote Online Audit

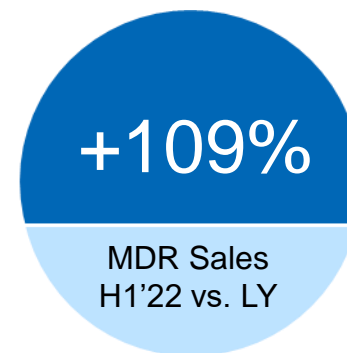
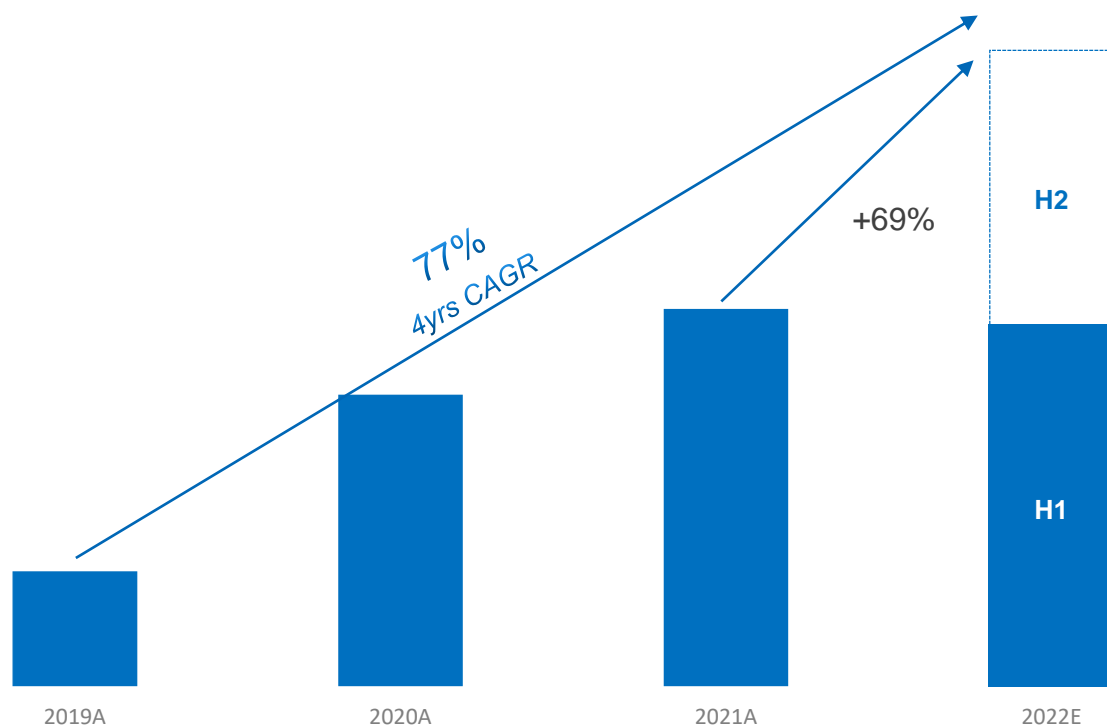
- “0 defect” passed the remote audit by US FDA in Jul 2021
- Passed OECD GLP audit and get highly acknowledged in 2022

Medical Device: Business Rebound



Medical Device: Capture MDR Opportunity

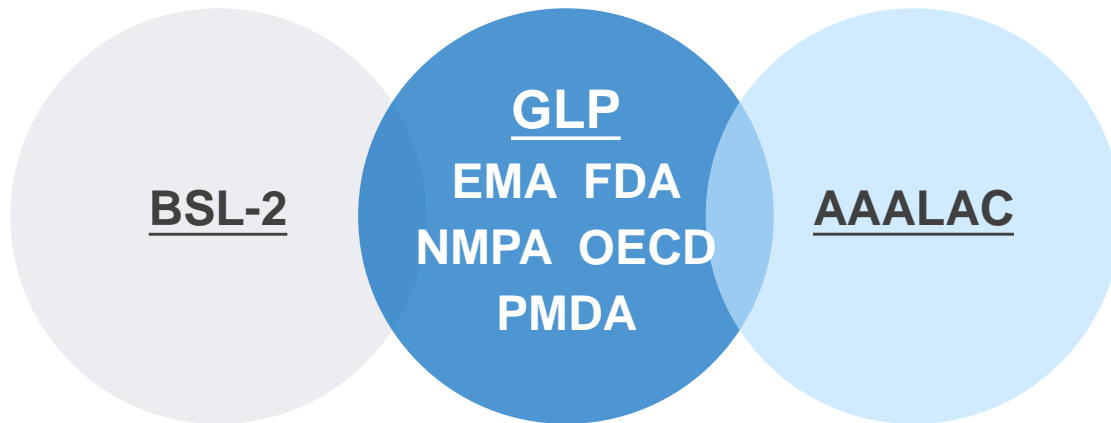
MDR Business



EU Medical Device Regulations (MDR) have driven adoption of analytical chemistry and material testing as the first step in biocompatibility risk assessment

Maintain the Highest Global Regulatory Standards

Global Regulations



China Based Facilities Certification Inspections

- **12** OECD
- **8** NMPA (CFDA) Certification
- **7** FDA
- **5** AAALAC
- **1** EMA
- **1** PMDA



WuXi Testing Growth and Synergy Opportunities

Follow the Molecule

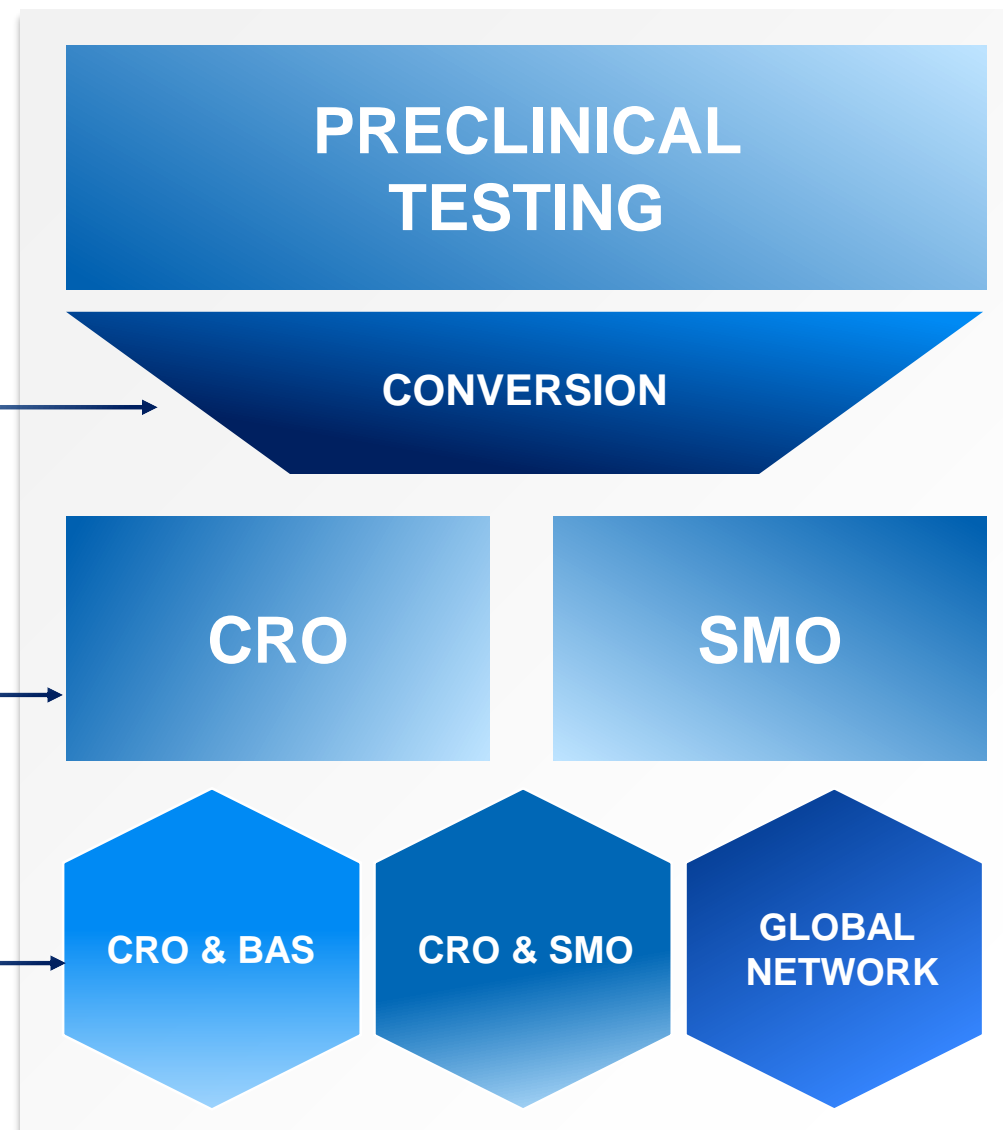
through conversion

Win the Molecule

through differentiation

Achieve Synergy

through integration



SMO: Broadest Hospital Coverage in China

~150

CITIES NATION WIDE

LOCAL CLINICAL RESEARCH TEAM

SMO: Enhanced Site Database and Management System

01. Hospital Management

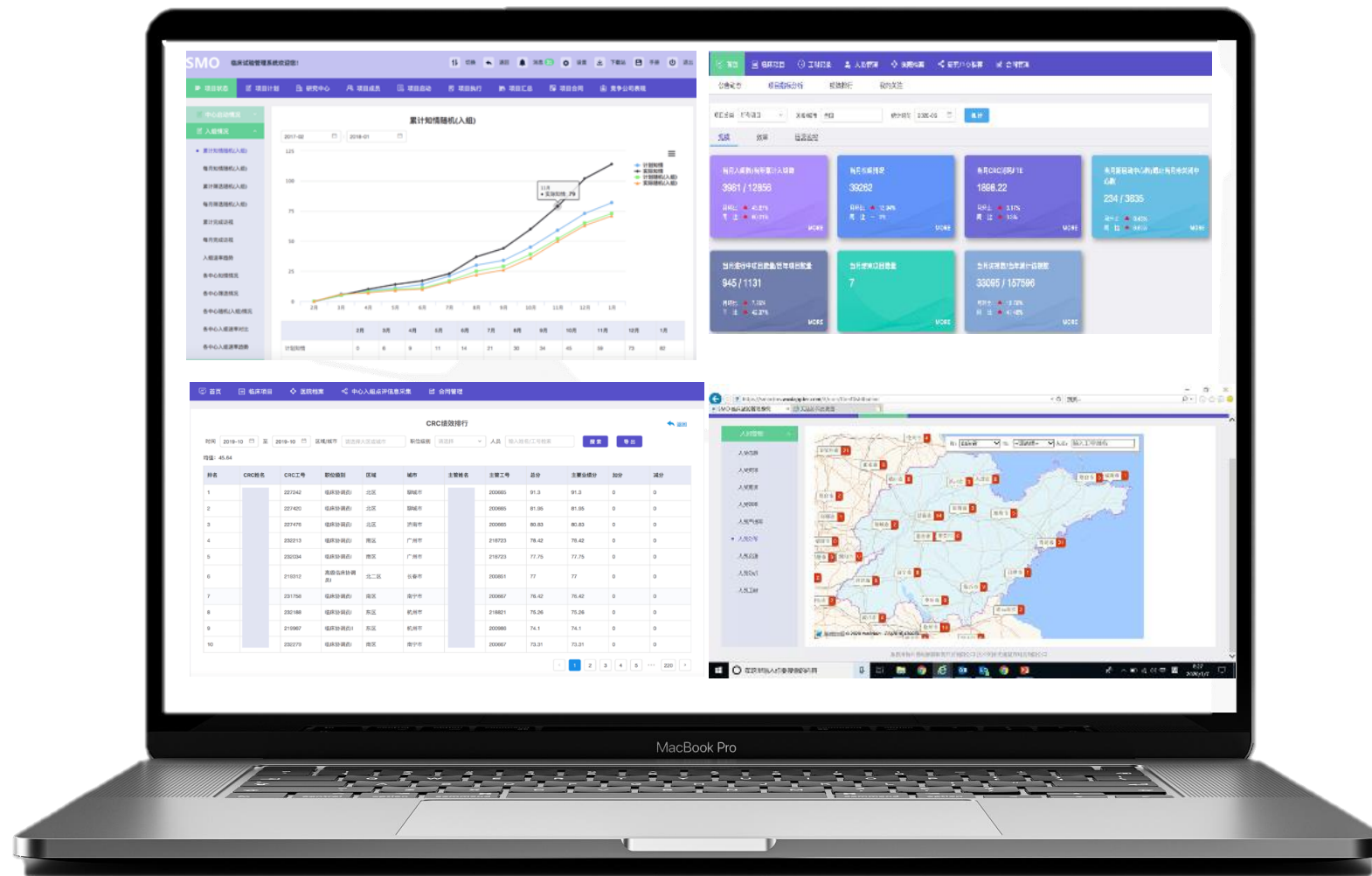
>1000 hospitals, a database of over 3,000,000 pieces of site start-up procedure items, 8000 site departments and investigators, **130,000+** patients in management.

02. Project Management

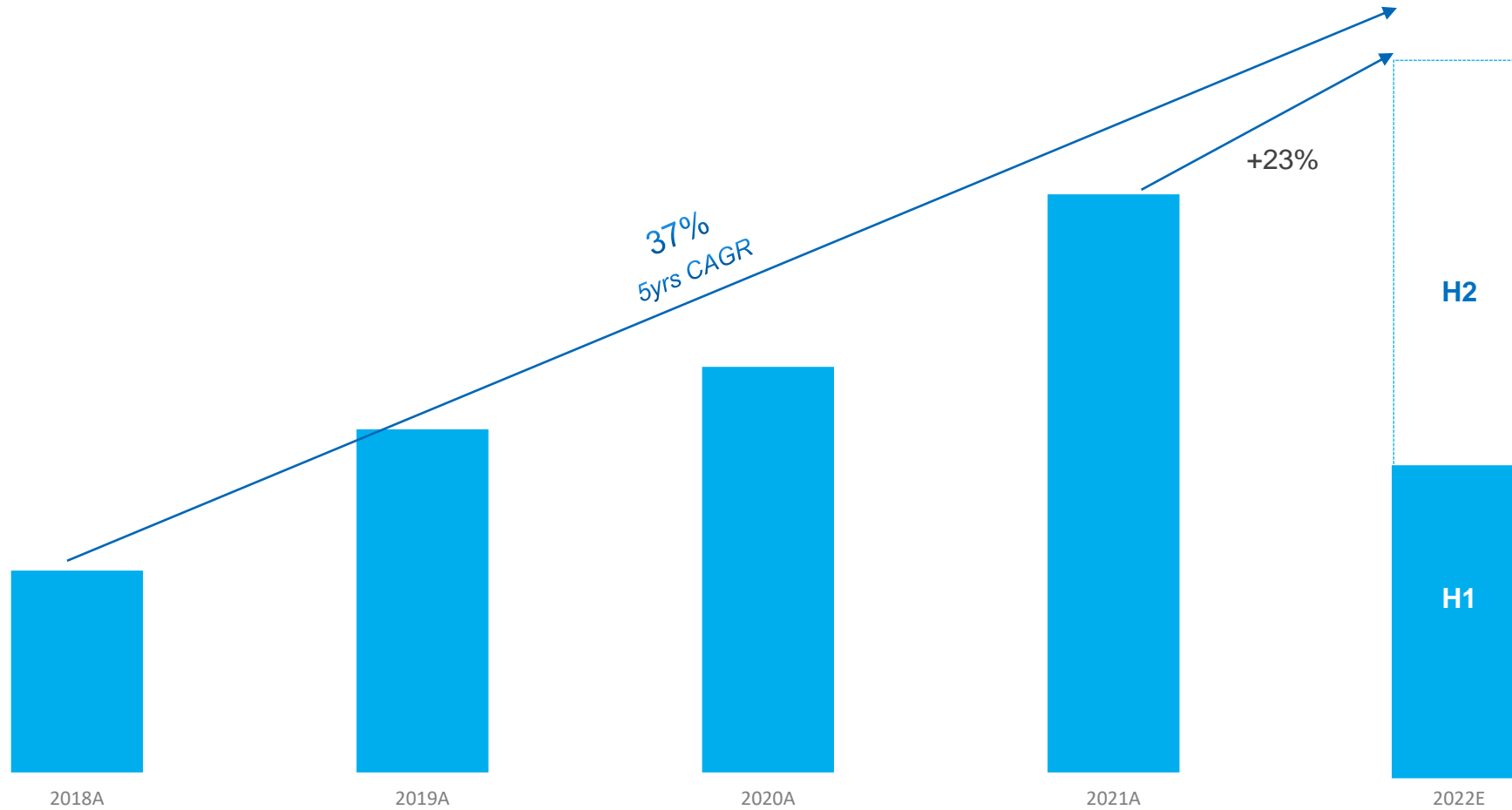
2000+ project experience, Supported 103 new drugs/medical devices approved on China / EMA / FDA market in recent 7 years

03. Personnel Management

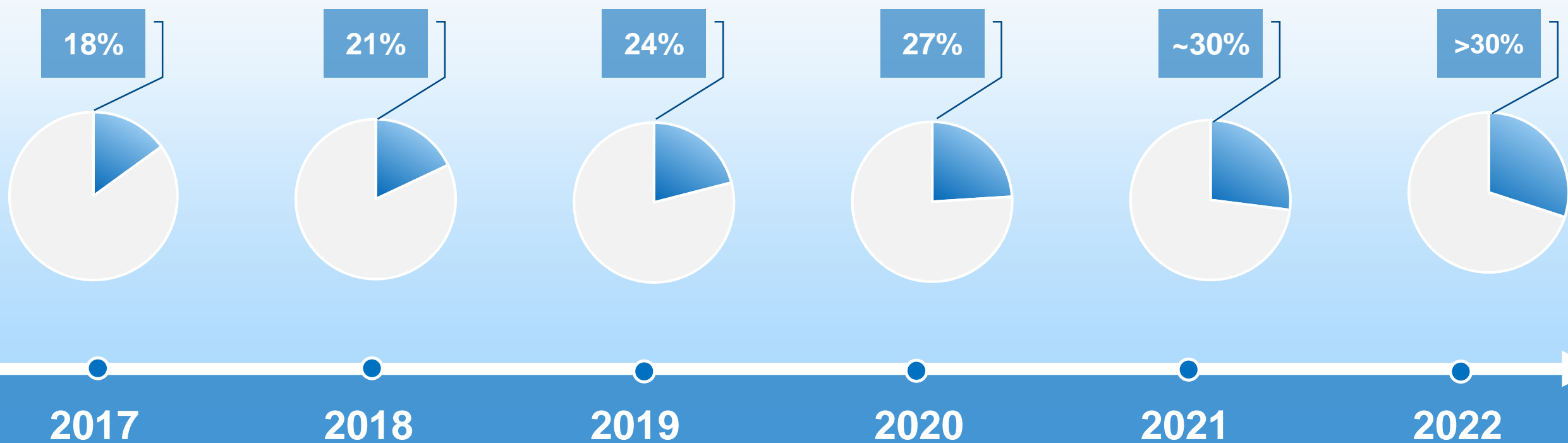
4600+ staff, efficient personnel management, resource allocation, performance appraisal



SMO Track Record and Growth Forecast

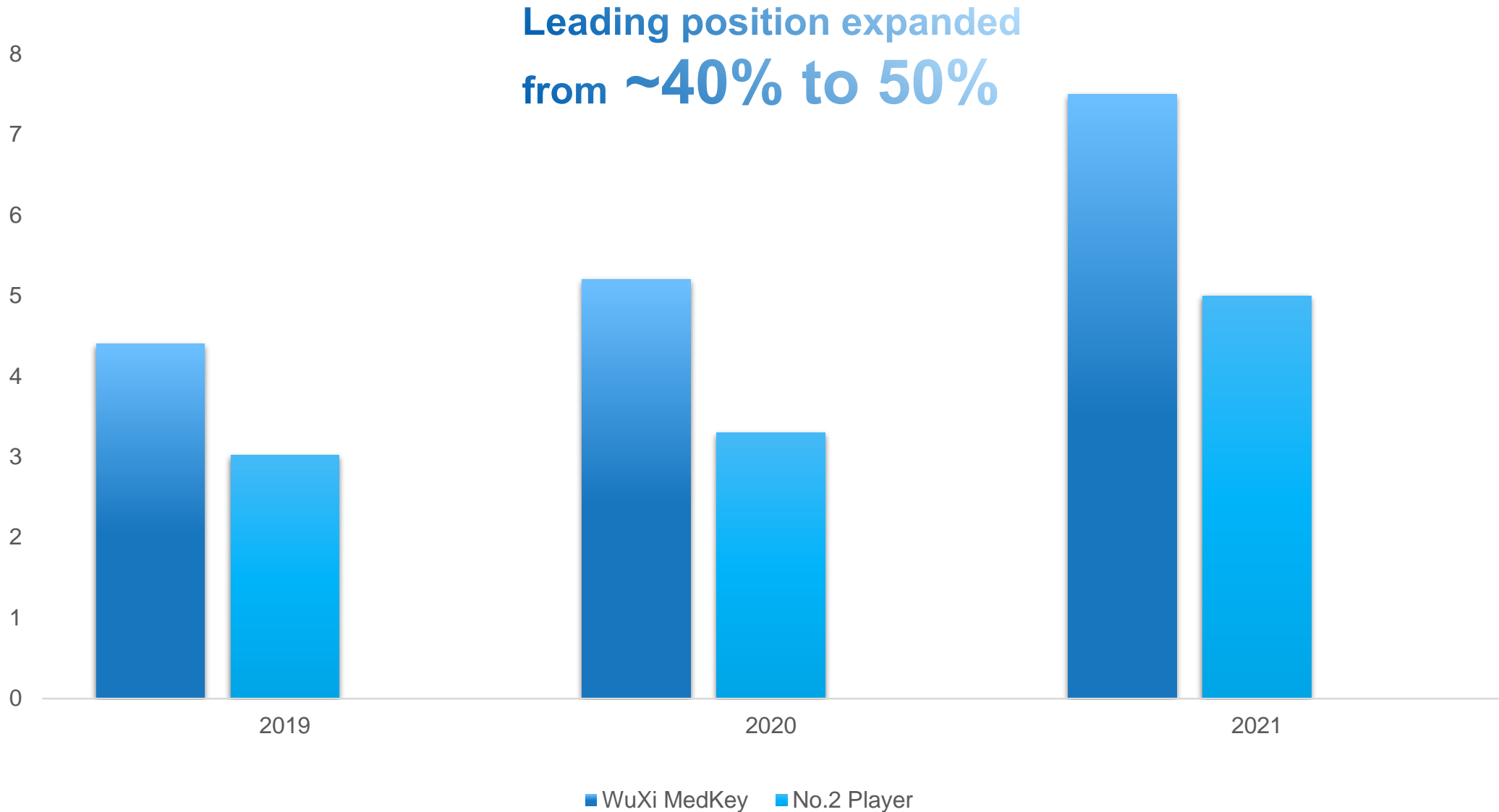


SMO Business Continues to Gain Market Share in China

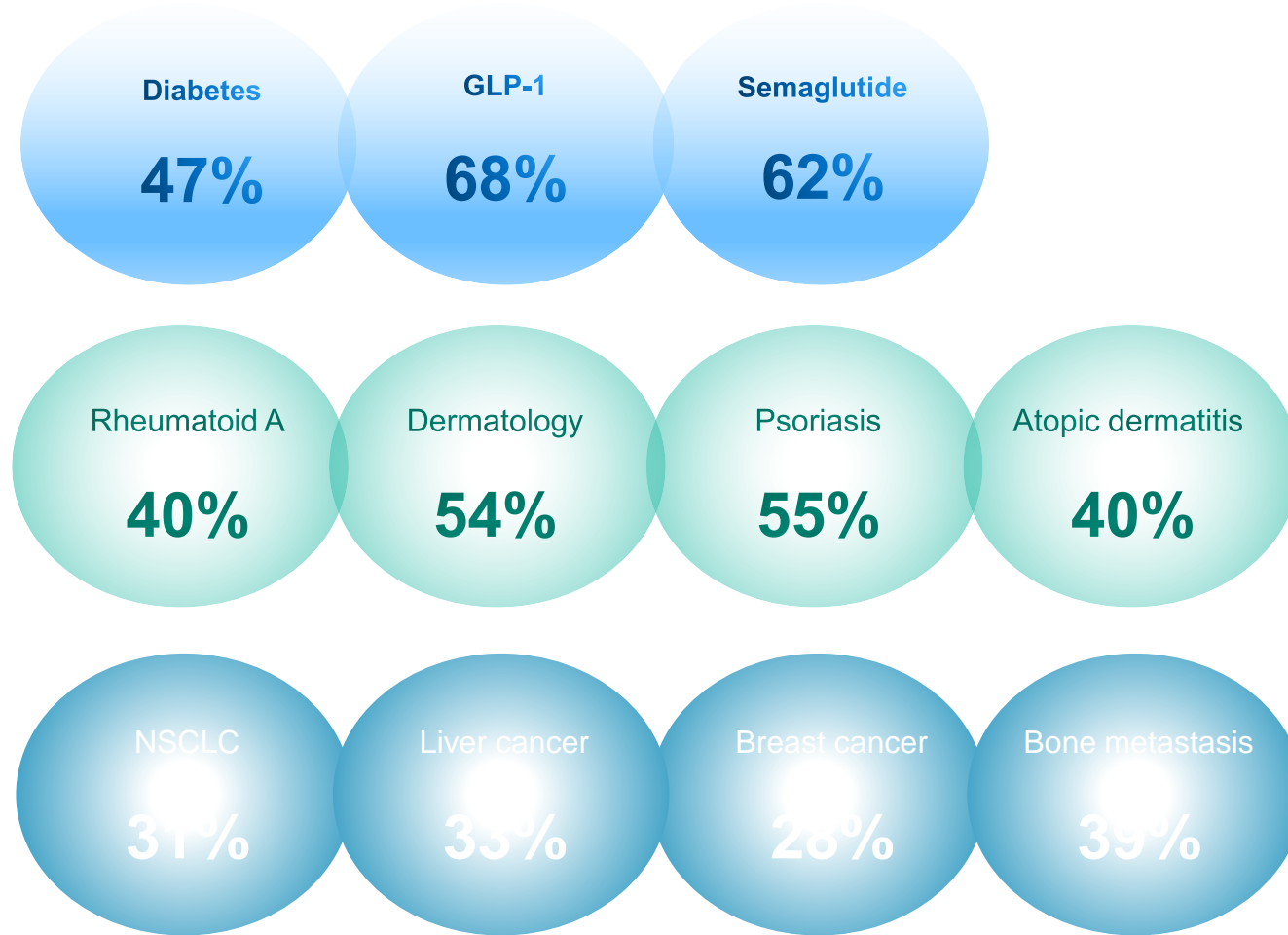


Source: internal analysis and industry reports

The Largest SMO in China with Expansion of Leadership Position



Market Leadership Across Disease Areas and Indications

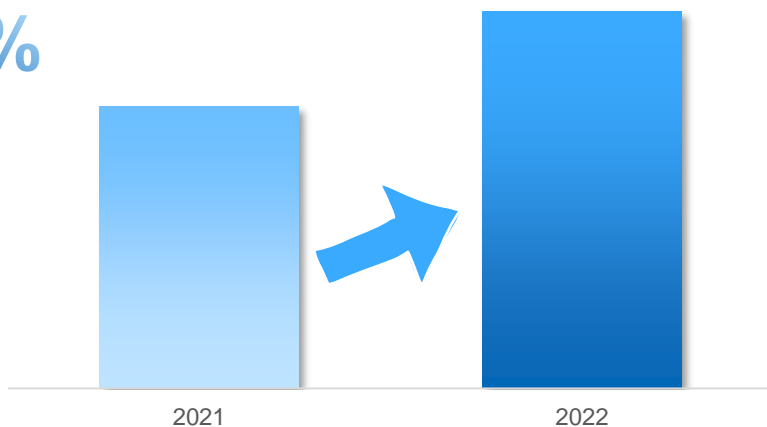


Market share as a percentage of clinical studies supported by SMO in a specific disease area / indication / drug class

SMO: Business Trend 2022H1

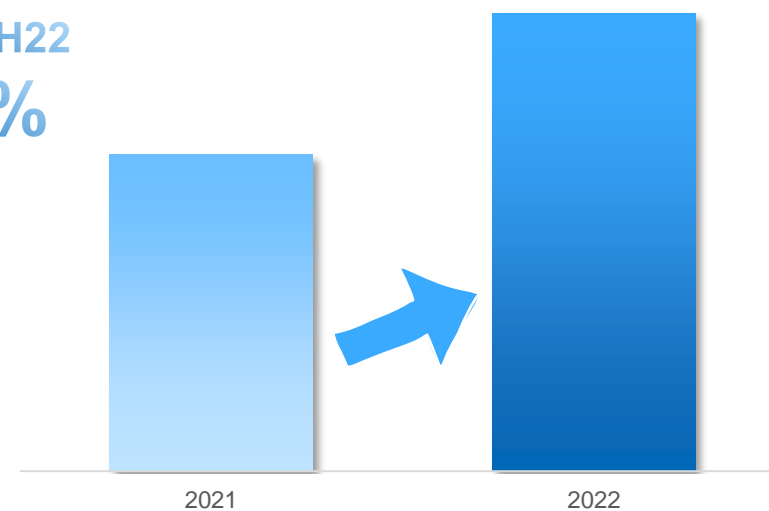
Contract value
1H21/1H22

+32%



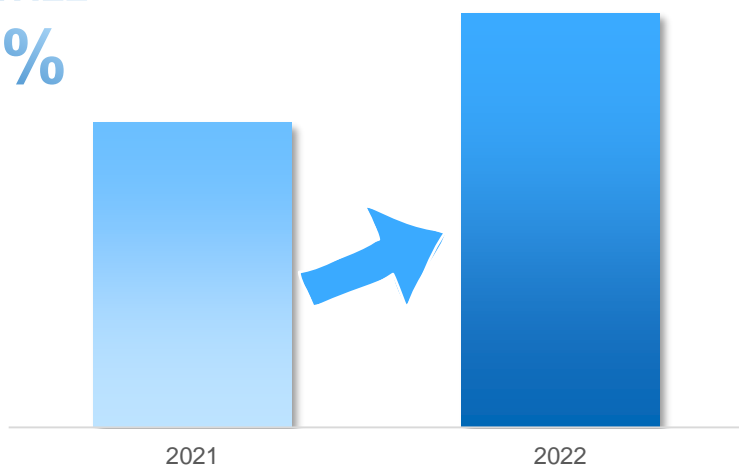
Award
1H21/1H22

+45%



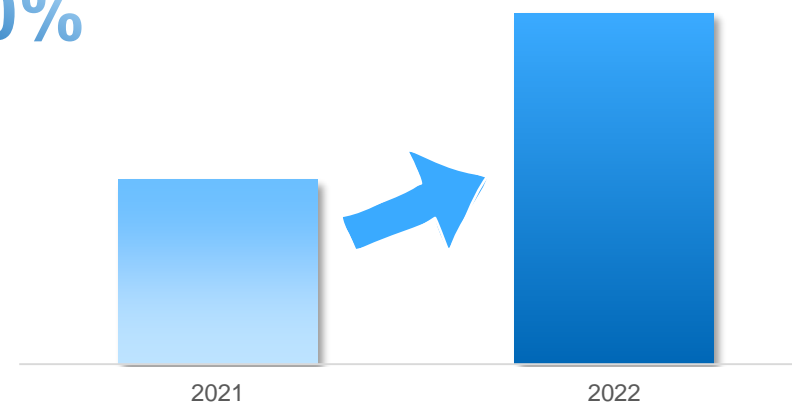
RFP
1H21/1H22

+36%



In bidding
1H21/1H22

+90%



SMO: Enable New Drug Approvals in China

16

approvals
in 2022H1

25

approvals
in 2021 whole year

- New generation GLP-1 product approved in China
- First SMA oral therapy product approved in China
- First bi-specific antibody approved in China
- First China antibody product approved in EMA
- First PD-1 product approved in China
- First CAR-T product approved in China

Strategy and Initiatives to Deliver Operational Excellence

3,066

2020 new initiated sites

5,220

2021 new initiated sites

+70%

34,888

2020 new enrolled pts.

50,593

2021 new enrolled pts.

+45%

403,036

2020 completed patient visits

560,457

2021 completed patient visits

+39%

- Establish Therapeutic Area (TA)-based Business Units to build expertise
- Focus on high productive sites and diseases
- Improve site initiation (SSU) capability
- Launch PCRC model to improve efficiency and quality
- Execute dynamic pricing strategy
- Leverage SMO clinical operation database advantage

WuXi Clinical: Global Clinical CRO Service Platform

WuXi Clinical provides Phase I to Phase IV clinical development services for products including pharmaceuticals, medical devices and IVDs

China

Australia

United States

- Shanghai, Headquarters
- Guangzhou
- Beijing
- Changsha
- Wuhan
- Xi'an
- Taipei
- Shenyang
- Chengdu

- Sydney

- Austin, Texas
- San Diego, California

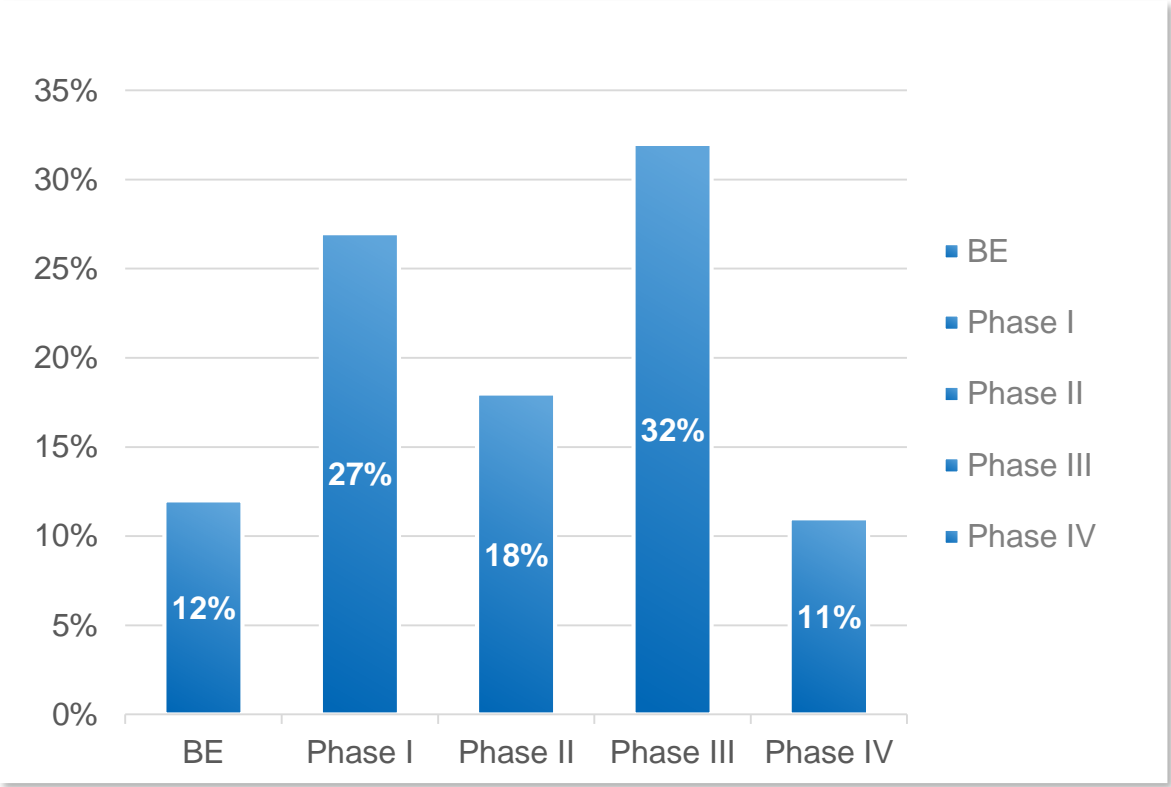
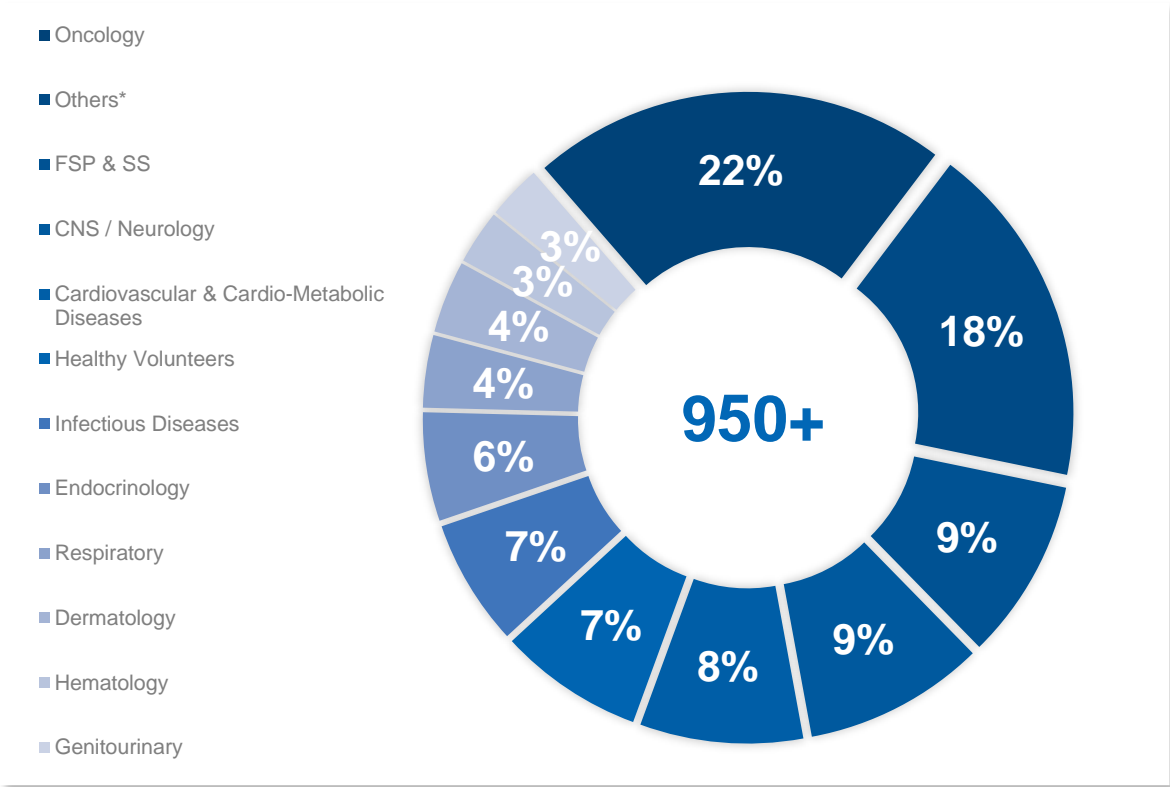
- Advantages of WuXi AppTec **Integrated** Service Platform
- Offices in **12** cities globally

- Covering **30+** major cities in China
- 830** employees globally

WuXi Clinical: Integrated CRO service in China and US

950+ projects of global clinical trials by China and US team

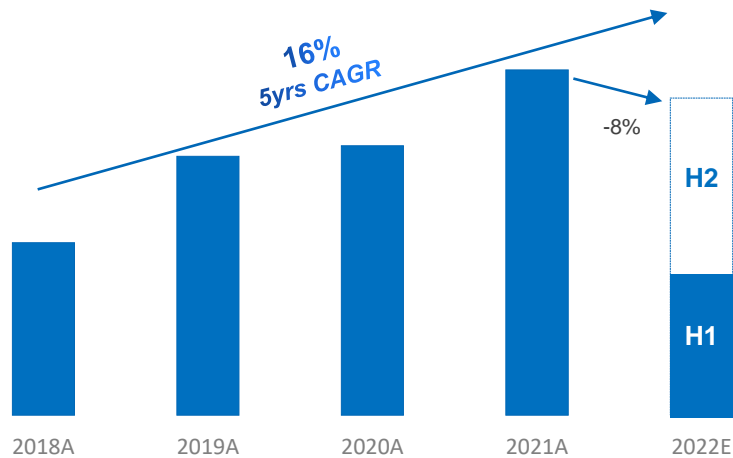
- Supported 50+ new drug applications (NDA)
- Received and passed 30+ inspections by NMPA / CFDA and US FDA in the last 6 years



* Including Immunology, Gastroenterology, Women’s Health, Hepatology, Medical Device and Musculoskeletal

WuXi Clinical: Facing Headwind with COVID Impacts

WuXi Clinical Growth Forecast



Challenges

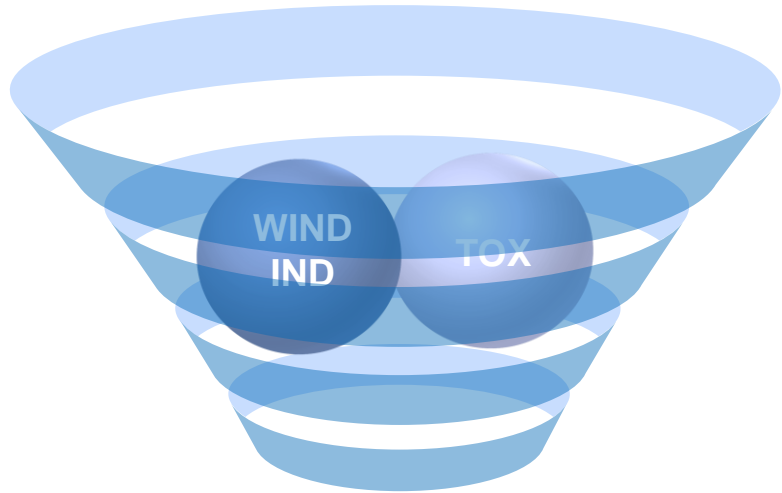
- COVID-related impacts on hospital operation and clinical trial conduct
- US BD underperforming

Solutions

- Execute Business Continuity Plan (BCP) to support ongoing clinical studies
- Engage with sponsors to catch up on study timeline after the hospitals re-open
- Dynamically manage headcount and control cost
- Pursue new opportunities with rigor
- Implement Therapeutic Area (TA) operation model to integrate global processes
- Enhance capability and delivery for global clinical studies

Leverage Preclinical Opportunities to Drive Conversion

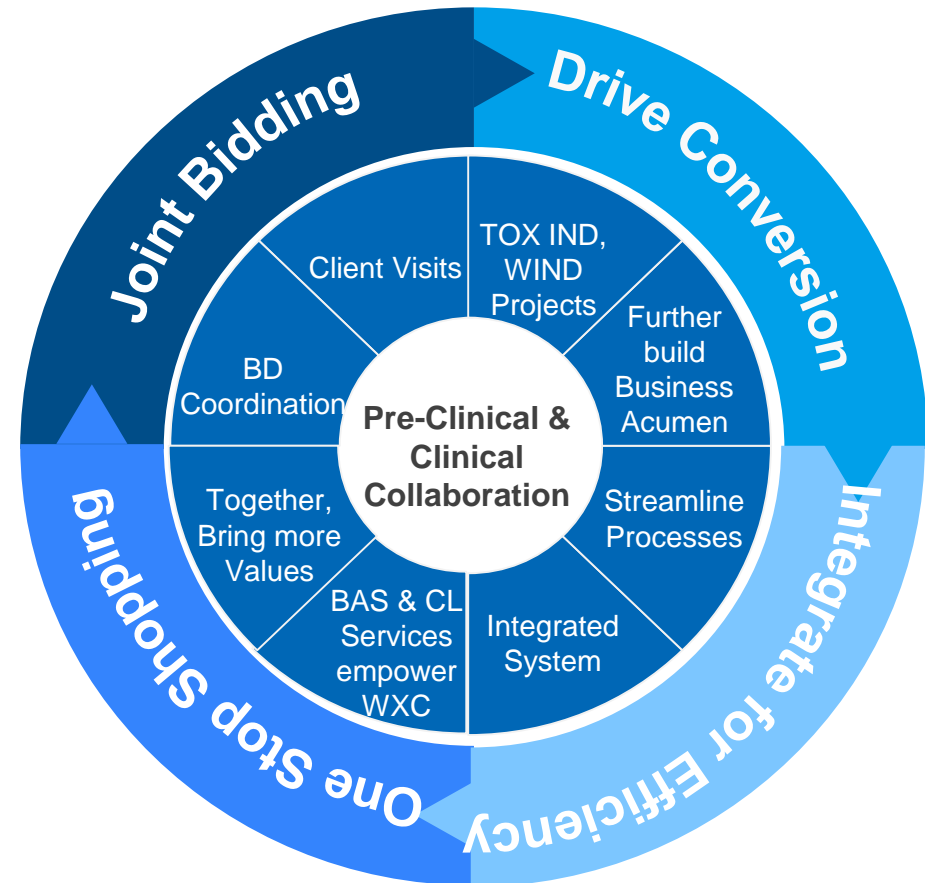
1 Drive Conversion
Leverage Preclinical Opportunities



2 Drive Conversion
Capture Clinical Wins



3 Drive Conversion
From WIND to WIND Plus

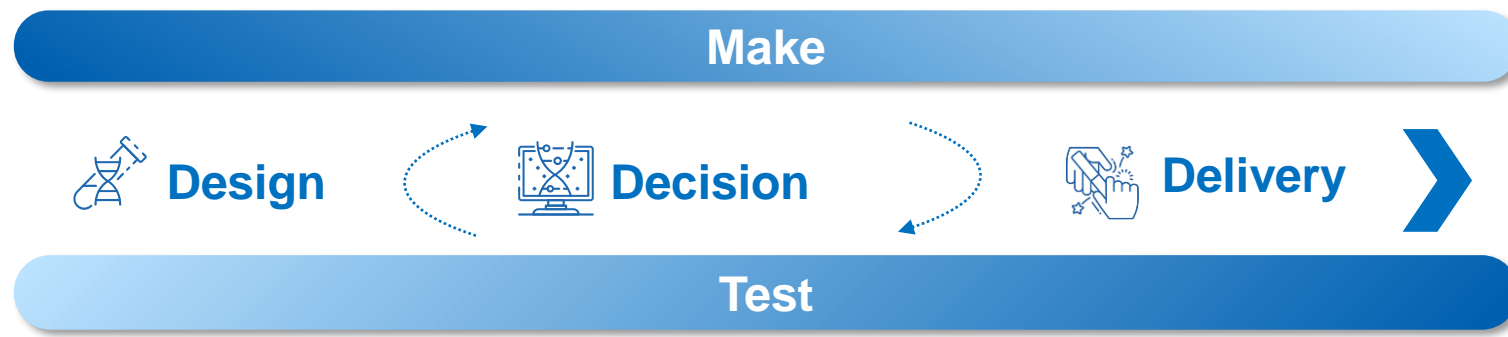


Data from: Jul 2021 to Jun 2022

Achieve Synergy in Operation between Clinical CRO and SMO

| Patient | Operation | Site |
|---|---|--|
| <ul style="list-style-type: none">• Leveraged SMO historical enrollment data for all 2022 new study enrollment projection and bidding support• Accurate projection of patient pool on over 20 co-bidding or collaborated projects in 2021, +100% vs. 2020. | <ul style="list-style-type: none">• Aligned with SMO historical site performance data of 1000+ hospitals and WXC site intelligence database to provide better site selection strategy in biddings and new win projects.• Achieved site startups speed up by about 2 weeks on average | <ul style="list-style-type: none">• Enhanced KOL alliance and built a strong KOL pool with 50+ KOLs, +30% increase from 2020• Established site partnerships in Beijing and Shanghai. More are in progress.• Conducted hundreds of remote monitoring visits |

Enable R&D Innovation through Making and Testing



WuXi Testing



WuXi Testing Key Growth Strategy

