



INTERIM RESULTS 2025

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Qyuns at a Glance - Powering The Boom of China's Autoimmune Sector with Comprehensive Capabilities

A clinical-stage biotech company exclusively focused on biologic therapies for autoimmune and allergic diseases



Exclusive Focus

Focus on **autoimmune and allergic disease therapies** since our inception



Full Coverage

Cover **four major disease areas** in the fields of Skin, Rheumatic, Respiratory and Digestive diseases



Huge Market

Global market size of **US\$187.5 billion** in 2022 of autoimmune and allergic diseases, the **second-largest therapeutic area** globally



Comprehensive Pipeline

1 product approved
9 drug candidates
21 IND approvals
Multiple ongoing clinical trials



Advanced Development Status

QX001S: 1st Ustekinumab biosimilar (SAILEXIN) in China (BLA approved)
QX002N: for AS (Phase III)
QX005N: for AD, PN (Phase III)
QX004N: for Ps (Phase III)



In-house Manufacturing

Established **commercial-scale cGMP-standard**
4 x 2,000L single-use bioreactors
~300 kg annual capacity



Strategic Partnership

Practical commercialization model to partner with **Huadong Medicine, Joincare and Hansoh Pharma.**
Reached an agreement with **Lilly Asia Ventures** and other funds for global NewCo's cooperation



Management Team

Experienced and diverse management team led by a successful serial entrepreneur and industry veteran

Exclusive Focus on Autoimmune and Allergic Disease Therapies, Covering Four Major Disease Areas And Key Therapeutic Pathways

mAbs bring commercial certainty; bsAbs boost efficacy and patient compliance



Seasoned Management Team with Extensive Industry Experience And Successful Entrepreneurial Track Records



Jiwan Qiu
Executive Director, Chairman, General Manager

30 years of extensive R&D experience in biotechnology industry with depth knowledge and understandings of innovation

Various entrepreneurial achievements, founded and led several antibody-focused biotech companies

Previously founded Jiangsu T-mab, developed 4 therapeutic biologic drugs, including LA-GCSF, anti-VEGF mAb, Denosumab biosimilar, GLP-1 analogue

Genetics and genetic engineering, **Fudan University**



Jianwei Li, Ph.D.
Deputy General Manager



Yiliang Wu
Executive Director,
General Manager of Cellularforce



Weidong Lin
Executive Director,
Deputy General Manager



Shenglong Wu
Deputy General Manager



Xiao Liu, Ph.D.
Deputy General Manager



Overview

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Innovative R&D Capability :
Dual Engines Linking Past And Forging Future

02

Strategic Partnership Capability:
Future Commercialization Guarantee

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Self-Sustaining Operations:
Stronger Financial Performance Insurance

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Financial Results

01

Innovative R&D Capability :

Dual Engines Linking Past And Forging Future

2025H1 Highlights



2025H1 sales performance of SAILEXIN is encouraging

- On Oct 29, 2024, SAILEXIN, China's first ustekinumab biosimilar, was approved with rapid nationwide sales launch;
- As of June 30, 2025, Qyuns has shipped over **60,000** units to Zhongmei Huadong.



Early stage bsAbs gradually form an innovative pipeline matrix

- As of the interim results announcement date, Qyuns has disclosed **4** innovative bsAbs, **3** of which are scheduled for IND filing within this year.



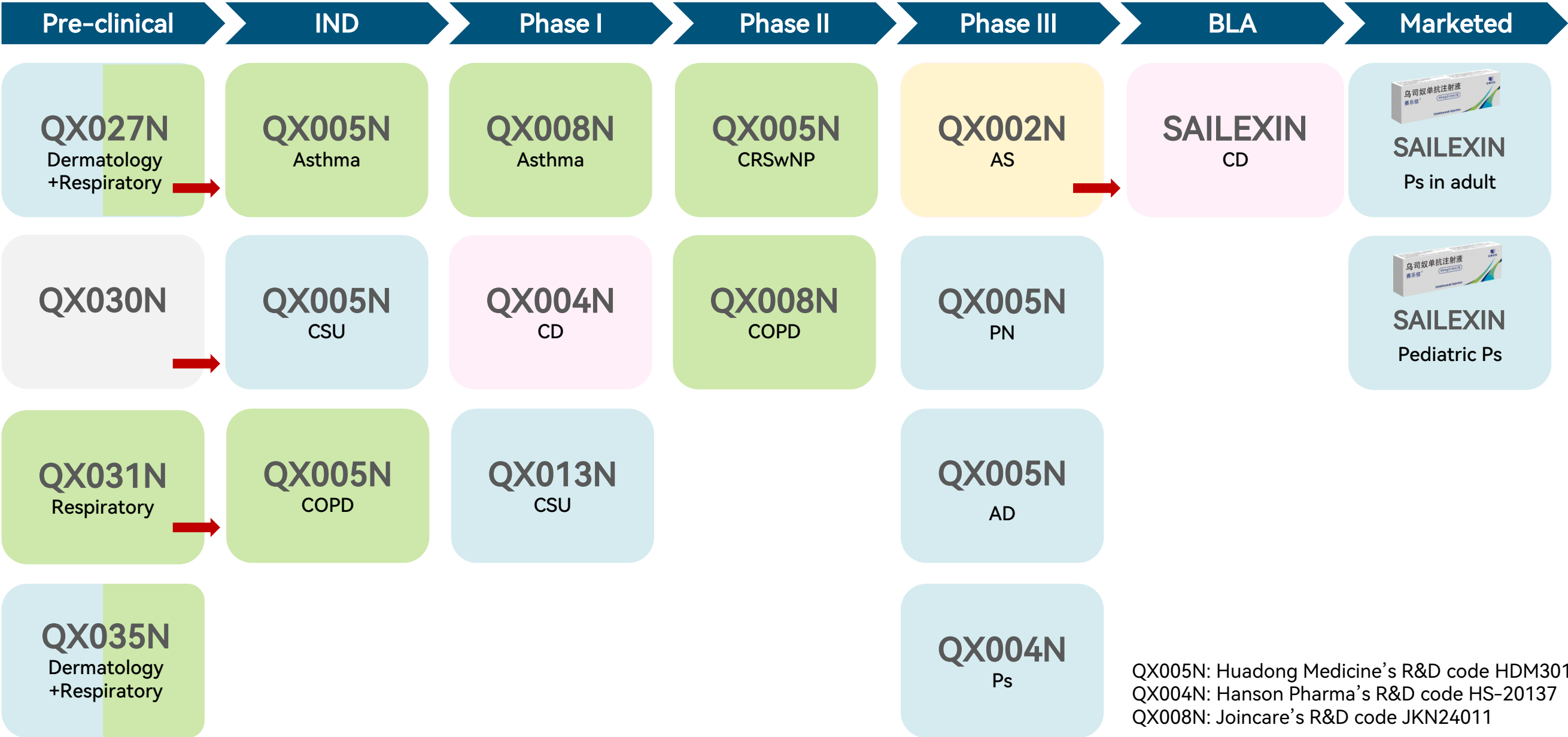
First step for globalization: first NewCo deal, USD 555 mil in total

- In April 2025, Qyuns and Caldera have entered into an out-license agreement granting Caldera an exclusive right to develop and commercialize QX030N globally.

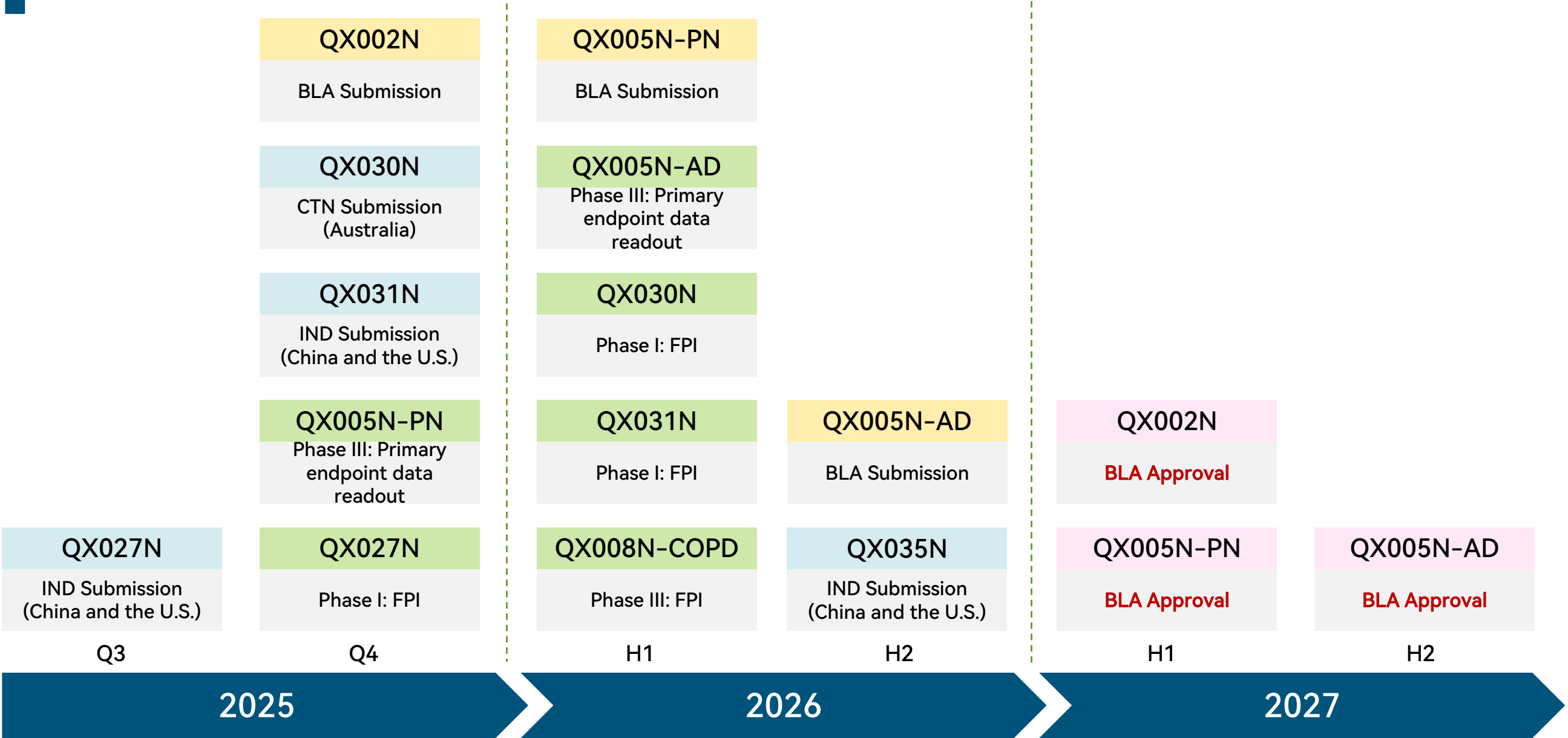
Comprehensive and Synergized Pipeline

Drug			Target	Indication	Preclinical	IND Approval	Phase I	Phase II	Phase III	BLA Approval	Partners
	QX001S SAILEXIN	IL-12/ IL-23p40	Ps								华东医药 HUADONG MEDICINE
			CD								
	QX005N	IL-4Ra	PN								华东医药 HUADONG MEDICINE
			AD								
			CRSwNP								
			CSU								
			Asthma								
			COPD								
	QX002N	IL-17A	AS								
	QX004N	IL-23p19	Ps								翰森制药 HANSON PHARMA
			CD								
	QX008N	TSLP	Asthma								健康元 Joincare
			COPD								
	QX013N	c-kit	CSU								
	QX027N	BsAb	Respiratory+Dermatology								
	QX030N	BsAb	Undisclosed								Caldera THERAPEUTICS
	QX031N	BsAb	Respiratory								
	QX035N	BsAb	Respiratory+Dermatology								

Bridging Past & Future: MAbs for Strong Delivery; BsAbs for Innovative Portfolio



Expected Progress of Certain Products in The Next Three Years



IND/CTN Submission

Clinical Milestones

BLA Submission

BLA Approval

QX005N: Huadong Medicine’s R&D code HDM3016
 QX004N: Hanson Pharma’s R&D code HS-20137
 QX008N: Joincare’s R&D code JKN24011

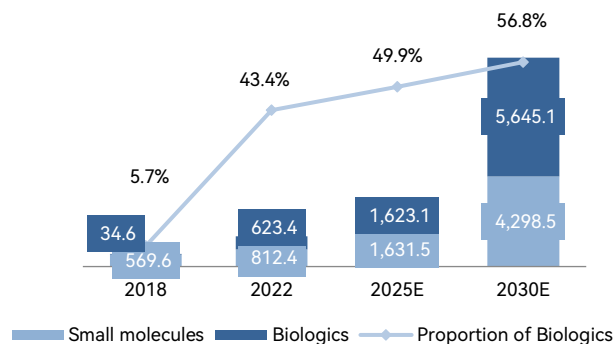
SAILEXIN – 1st Ustekinumab Biosimilar Approved in China & Expected Blockbuster

Market Opportunities and Competition

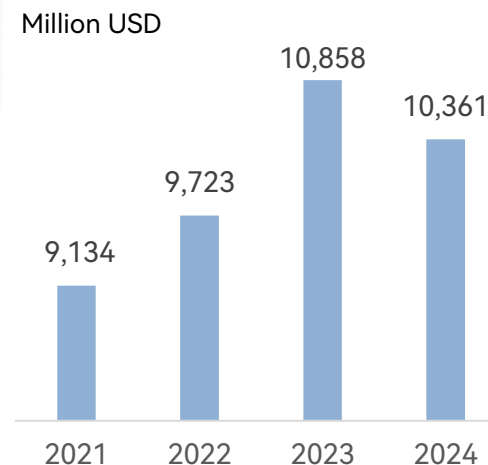
Market Opportunities

In 2023, the number of psoriasis patients in China is estimated to be 7.2 million, among whom 60% have moderate-to-severe psoriasis

Ps drug market in China (million USD)



Ustekinumab Global Sales



Cooperation with Huadong Medicine



- In 2020, Zhongmei Huadong was granted the rights for joint development and exclusive commercialization of QX001S in **mainland China**
- We have received the upfront payment and milestone payments from Zhongmei Huadong, amounting to a total of RMB **50 million**
- After offsetting the attributable loss from the commercialization of QX001S, the parties will share the cumulative pre-tax profits from QX001S **on a 50:50 basis**
- The strategic partnership with Zhongmei Huadong will ensure **more efficient commercialization** of QX001S

Source: NMPA, CDE, Frost & Sullivan, prospectus of Dermavon

Competitive Advantages

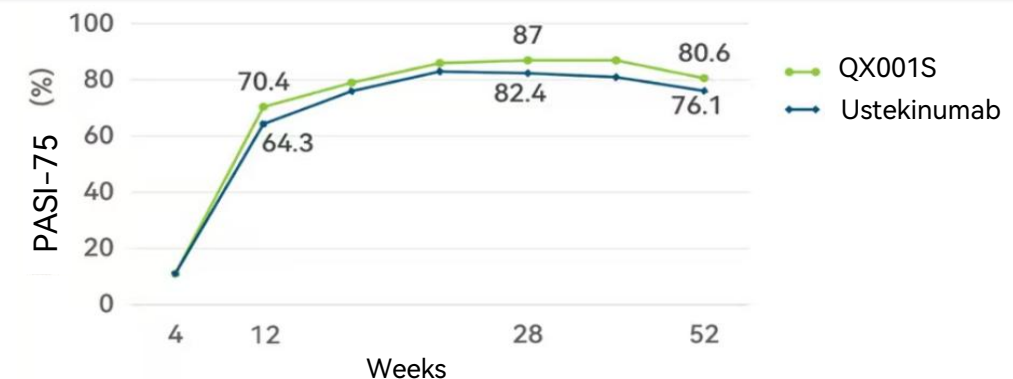
The first marketed Ustekinumab Biosimilar in China (SAILEXIN)

- ✓ The commercial collaboration with Huadong Medicine has ensured rapid commercialization. As of 2025H1, over **1,200** hospitals have prescribed the product, and Qyuns has shipped more than **60,000 units** by 2025H1.
- ✓ Expect **better accessibility** than Stelara® (annual cost ~RMB16,063 for maintenance treatment) given Qyuns' manufacturing capability

Ustekinumab: 10+ years clinical evidences on safety and efficacy

- ✓ **Higher drug survival rate: more effective** than TNF-α and IL-17 mAbs in long term
- ✓ **Convenient treatment regimen:** Q12W (QX001S) vs. Q4W (IL-17A mAb)
- ✓ In Phase III clinical trial for Ps, **QX001S demonstrated clinical equivalence to Ustekinumab** in terms of efficacy, safety, immunogenicity and PK profile

Clinical Development and Commercialization Process



Oct 2024

Feb 2025

Mar 2025

BLA approval for Adult Ps

BLA accepted for CD

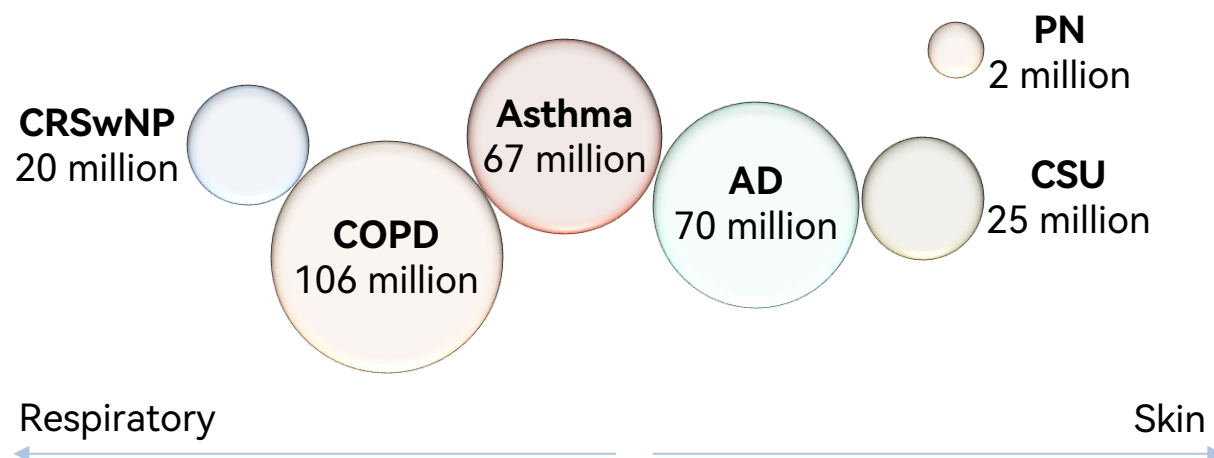
BLA approval for pediatric Ps

QX005N – One of the Only Two IL-4Ra mAbs with **BT**D in China

QX005N

- IL-4Ra controls the signaling of both IL-4 and IL-13, which is critical in the initiation of type 2 inflammation
- QX005N is designed to inhibit IL-4Ra, a well-validated, broad-acting target for a wide range of indications

2022 Prevalence of Covered Indications in China

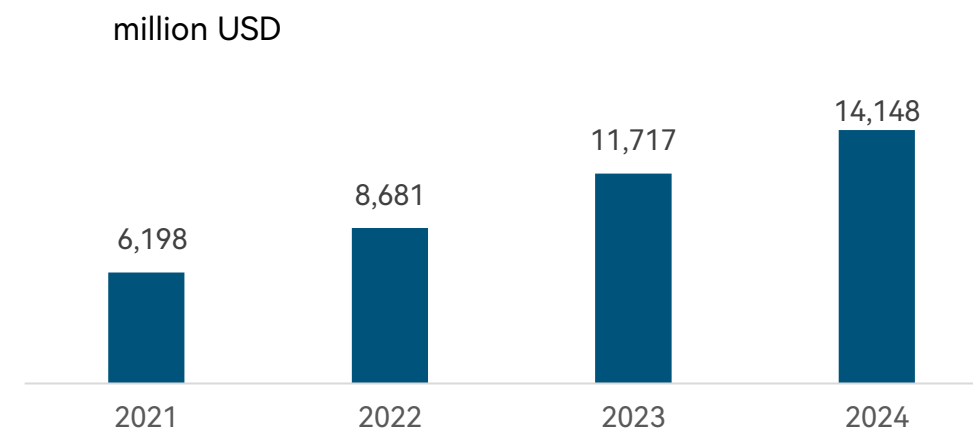


Source: Frost & Sullivan, Sanofi annual reports



- In July 2024, the **exclusive joint development rights, exclusive commercialization options, and MAH transfer priority** were granted to Zhongmei Huadong in the designated area
- Zhongmei Huadong will **bear 50% of the Ph III clinical costs** for the cooperative indications

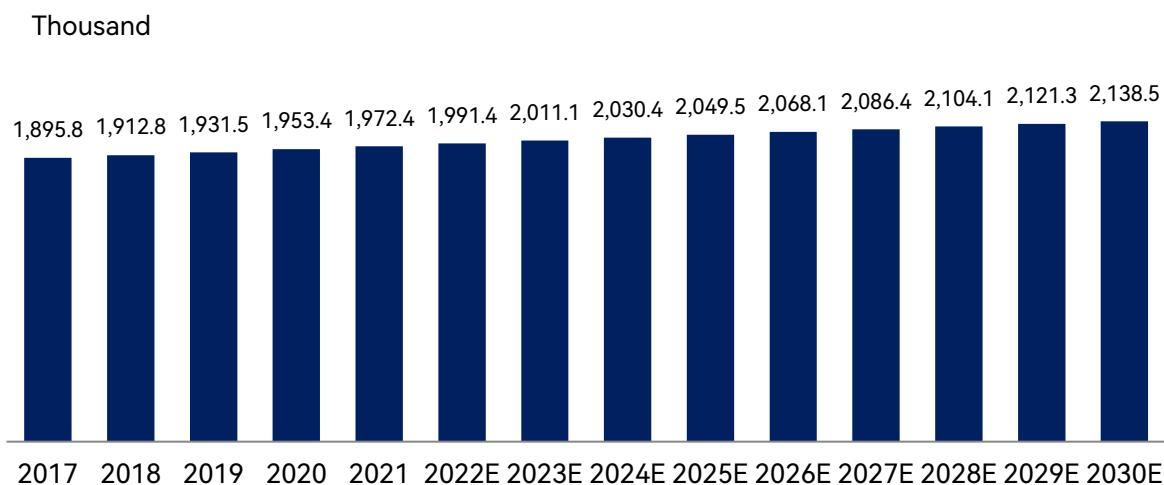
Dupixent® Global Sales, 2021 – 2024



QX005N – 1st Biologic Drug Candidate for PN Developed by Chinese Domestic Company

Market Opportunities and Competition

Prevalence of PN in China, 2017-2030E



- Commonly associated with other skin diseases or underlying medical conditions that affect multiple body systems
- Biologic drugs have become a guideline treatment option for PN

Marketed Targeted Biologics for PN in China

Brand Name	INN	Company	Target	NMPA Approval Time
Dupixent	Dupilumab	Sanofi	IL-4Rα	2023

Unmet Medical Needs

Limitation of current treatment

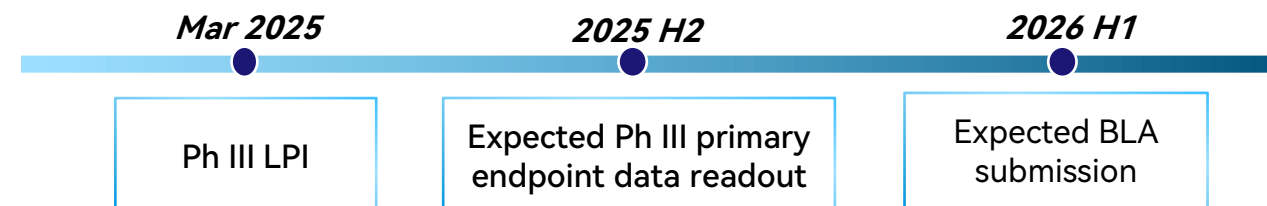
- Development of the PN drug market in China is still at an early stage
- Current treatment (topical steroids and topical anesthetics): only for **limited duration** due to **side effects**
- Lack of biologic drug: **Dupixent®** is the only treatment approved by FDA and by NMPA for PN
- Dupixent® is the only biologic drug approved for PN in China

Lack of effective treatment

Insufficient study

- **2024.1 QX005N BTD received for PN**
- In the Phase II trial, the proportions of patients achieving ≥ 4 -point improvement in WI-NRS score from baseline at Week 16 were **76.7% (300mg), 83.3% (450mg), and 76.7% (600mg) respectively, all demonstrating statistically significant superiority over the placebo group (30%; $P < 0.0001$)**

Clinical Development



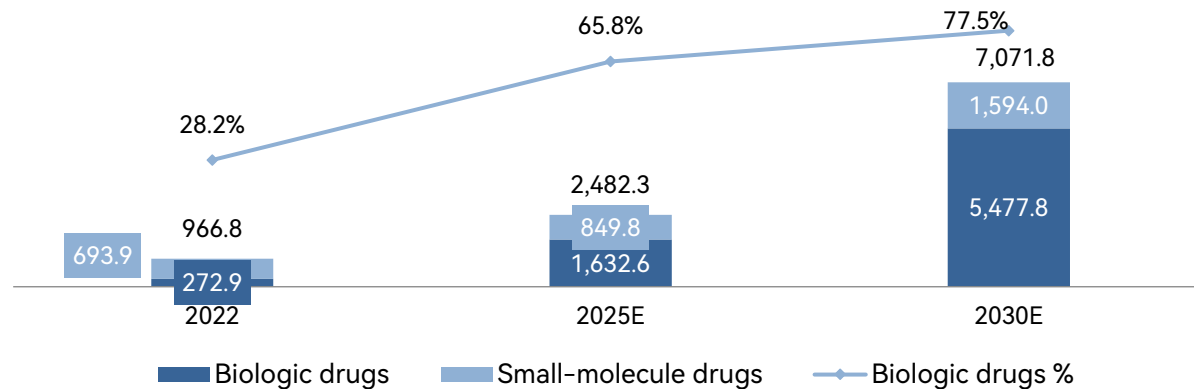
QX005N – Treatment for Moderate-to-Severe AD

Market Opportunities and Competition

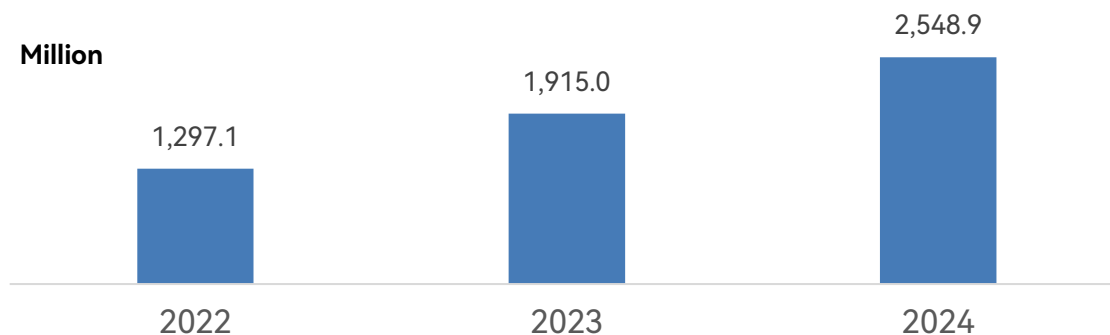
Market Opportunities

AD patients in China reached 70.3 million in 2022 and is anticipated to reach 78.5 million in 2030, 30% of which are suffered from moderate-to-severe AD.

AD drug market in China (million USD)



Sales Volume of Dupilumab in China



Source: Company data, Frost & Sullivan, Menet

Unmet Medical Needs

- **Systemic immunosuppressants** face significant safety concerns and inadequate efficacy in the long-term treatment of moderate-to-severe AD
- **Children above 6 months** demand absolutely safe and effective treatment of AD

Clinical Development and Commercialization

Commercialization

- > Key products in the dermatology portfolio
- > It is expected to quickly realize its value through Huadong Medicine's channels

Phase III

- > Adolescent (12-17 yrs old) and adult patients are enrolled simultaneously
- > Targeting a broader population

Phase I~II

- > Good tolerability and safety, with significant efficacy

2025 Q3

2026 H1

2026 H2

Phase III LPI

Expected Ph III primary
endpoint data readout

Expected BLA
submission

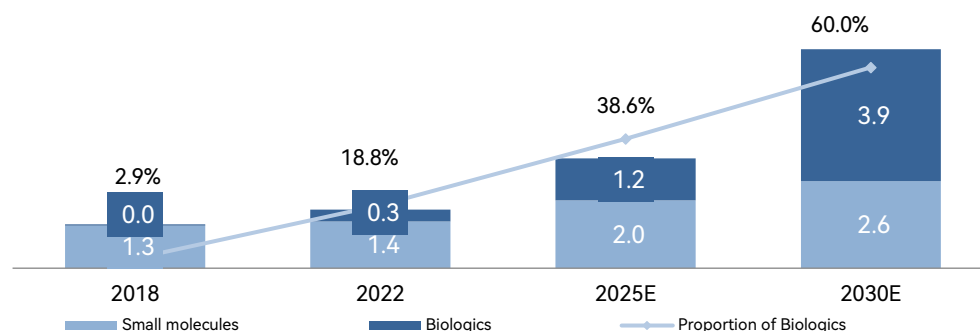
QX002N – IL-17A mAb for AS

Market Opportunities and Competition

Market Opportunities

The AS patient population in China reached 3.9 million in 2022, mainly younger adults

AS drug market in China (billion USD)



Unmet Medical Needs

- **40%** of AS patients intolerant to / inadequate disease control with **anti-TNF therapies**
- Guided as 2L standalone treatment for AS (the same designation as TNF inhibitors) for AS patients **with high disease activity after receiving first-line traditional treatments**

Competitive Advantages

★ IL-17A vs. TNF

- ✓ Guided as 2L standalone treatment for AS, IL-17A inhibitors have shown **clear clinical benefit** in patients who are intolerant to or fail to achieve adequate disease with TNF-α inhibitors
- ✓ IL-17A inhibitors are **more targeted** and with generally **fewer warnings and precautions**

★ Cost-effective in-house commercialization

Clinical Development

Phase III

> Ph III primary endpoint data read-out in Feb 2025

Data

- > **ASAS40 40.4% vs 18.9%** (QX002N vs Placebo)
- > **ASAS20 65.2% vs 41.3%** (QX002N vs Placebo)
- > Good safety

Feb 2025

2025 H2

Ph III primary endpoint
data read-out

Expected BLA
submission

QX004N – IL-23p19 mAb for Psoriasis and Crohn's Disease

IL-23p19 is currently a better target for psoriasis
The treatment effect of psoriasis will be further improved,
and it is conducive to long-term management

Strategic cooperation with Hansoh Pharma



- On 24 April, 2024, Qyuns and Hansoh Pharma reached a strategic cooperation
- Under the terms of the agreement, **Hansoh Pharma has paid an upfront payment of RMB 75.0 million**, and shall pay potential milestone payments of no more than RMB 1,032.0 million upon the achievement of development, regulatory and sales-based commercialization milestones, plus tiered royalties on future product sales
- **Recently, Qyuns has received the Ph III milestone payment and other payments for Ps from Hansoh Pharma in accordance with the license agreement, totaling RMB 58M**

Same target competitor

The global sales of Skyrizi®, the same target competitor, were \$1.59 billion in 2020, \$2.94 billion in 2021, \$5.17 billion in 2022, and \$7.76 billion in 2023, with an **average annual growth rate of 70.3%**.
At the end of 2024, its global sales reached **\$11.7 billion with the annual growth rate of 50.9%**.

Clinical Development

Ps

- > Dec 2024: Ph I data published in JAMA Dermatology
- > Mar 2025: Hansoh presented Ph II data as a late-breaking oral presentation at the 2025 AAD Annual Meeting
- > **92.3% PASI 75 and 76.9% PASI 90** response rates at 16 weeks (200mg dose)

CD

- > 2024.5 Ph Ia completed
- > Led by Hansoh subsequently

May 2024

Aug 2024

2025 Q2

CD Ph Ia completed

Ph II primary endpoint
data read-out for Ps

Ph III initiated

QX008N – TSLP mAb for Respiratory Diseases

- TSLP is a cytokine expressed by the airway epithelium and sits at the top of multiple inflammatory cascades
- TSLP monoclonal antibody is the only biologic drug that is independent of eosinophilic levels and reduces the exacerbation of severe asthma in a broad population

Same target competitor

The global sales of Tezspire®, a competitor with the same target, were \$174 million in 2022 and **\$653 million** in 2023, with an annual increase of **275.3%**. At the end of 2024, its global sales reached **\$1.219 billion with the annual growth rate of 86.7%**.

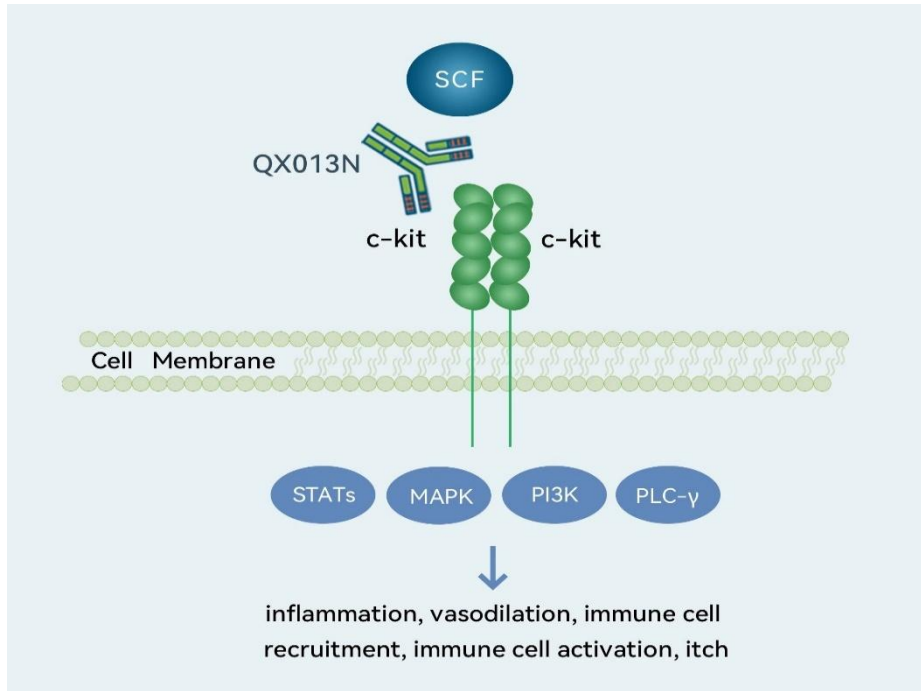
Tezspire® recently received **breakthrough therapy designation** from the FDA as an additional maintenance therapy for patients with moderate-to-severe COPD.

Strategic Cooperation with Joincare



- In January 2024, we entered into a technology transfer agreement with Joincare to grant Joincare an exclusive license to develop, manufacture and commercialize QX008N in China, Hong Kong and Macau. Joincare will be responsible for the BLA application and will be the MAH of QX008N in the aforementioned area, once approved. We retain the exclusive rights to develop, manufacture and commercialize QX008N outside China, Hong Kong and Macau
- Joincare has completed Phase II patient enrollment for COPD in China as the leading player in this field, and is expected to initiate Ph III trial in 2026H1

QX013N – First Biologic Drug Candidate Targeting c-kit in China



QX013N is a humanized IgG1 mAb independently developed by Qyuns and is the first biologic drug candidate targeting c-kit in China. Used for the treatment of chronic spontaneous urticaria (CSU).

Mast cells are a key driver of the inflammatory response, yet current therapies focus on blocking mast cells and releasing medium, not the source.

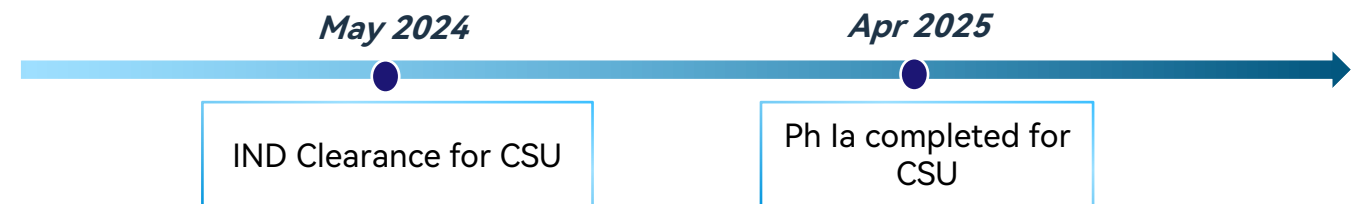
Direct inhibition of mast cell activation and degranulation is an innovative therapy for the treatment of a variety of allergic diseases.

Potential future indications include atopic dermatitis, asthma, allergic rhinitis, etc.

Indication Introduction

CSU is a common chronic inflammatory skin disease characterized by the spontaneous appearance of wind masses and/or angioedema on the skin, lasting more than 6 weeks. The disease is prone to relapse and is often accompanied by a persistent itching or burning sensation, seriously affecting patients' quality of life and physical and mental health. Public information shows that 5.1 million patients in China are receiving treatment and 2 million are not under control.

Clinical Development



Innovative BsAbs Portfolio: Delivering Superior Solutions for Chronic Disease Management and Marking a New Era of Company Pipeline



QX027N

- Focusing on respiratory and skin diseases
- Planned IND filings in China and US in Q3 2025



QX031N

- Focusing on respiratory diseases
- Planned IND filings in China and US in Q4 2025



QX030N



- Undisclosed indication area
- Planned CTN submission in Australia in Q4 2025



QX035N

- Focusing on respiratory and skin diseases
- Planned IND filings in China and US in Q4 2026

Dermatology

- For AD, immediate relief of itching and skin lesions, reduction of recurrence risk, and significant prolongation of recurrence interval
- For CSU, immediate symptom relief to meet the needs of refractory patients

Respiratory

- Strengthen disease-modifying therapy
- Delay, prevent and even reverse disease progression
- Maintain long-term efficacy and achieve the goal of clinical cure

IBD

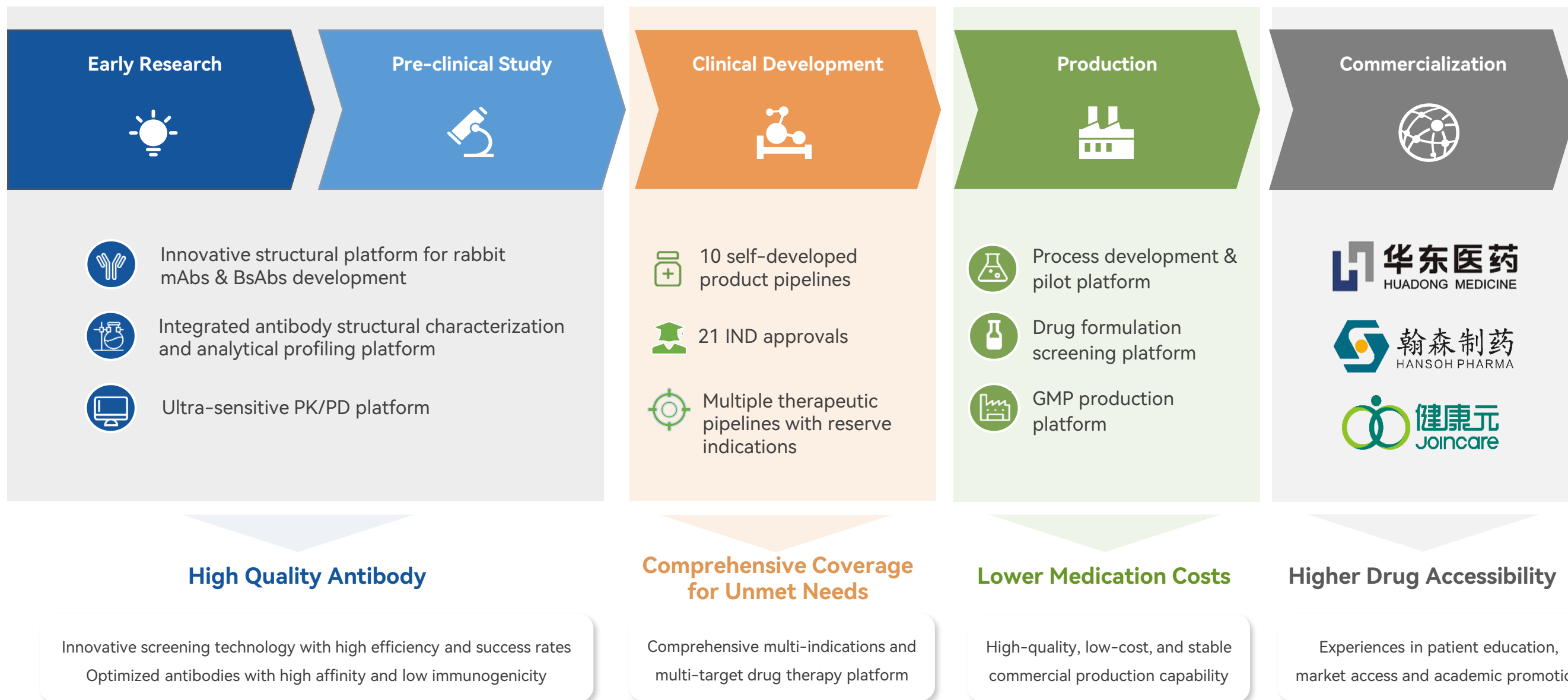
- Improve clinical and endoscopic remission rates
- Meet the needs of alternative treatment for patients who have received biologic therapies

Overall goal: Improve drug efficacy, optimize dosing intervals, enhance patient compliance, and reduce medication costs

02

Strategic Partnership Capability: Future Commercialization Guarantee

R&D-Production-Commercialization Integrated Strategic Alliance



Strategic Cooperation: Verify Pipeline Value, Generate Cash Income, Accelerate Clinical Progress, Enhance Commercial Certainty



QX001S: strategic cooperation in Aug 2020
QX005N: strategic cooperation in Jul 2024



QX004N: strategic cooperation in Apr 2024



QX008N: strategic cooperation in Jan 2024



QX030N: strategic cooperation in Apr 2025

QYuns
荃信生物

03

Self-Sustaining Operations: Stronger Financial Performance Insurance

Cellularforce: Commercial Scale Production Capacity Supporting CDMO Business Transformation

One of only a few Chinese biotech companies that are focused on autoimmune and allergic diseases have established **commercial-scale in-house manufacturing capability**

Manufacturing facility

- **4 x 2,000L** single-use bioreactors
- Approximately **300 kg annual capacity of therapeutic antibodies**



cGMP-standard manufacturing facility with excelled CMC capability and quality management will secure **high quality, cost controllable and reliable supply of products** for clinical study and commercialization

CMC capability and quality management

- **Strong CMC capability** to improve production efficiency
- **Excelled quality management** through QA and QC



Strong Operational Cash-generating Capability Delivering Sustained Capital Reserves



Income from BD
RMB 180M +



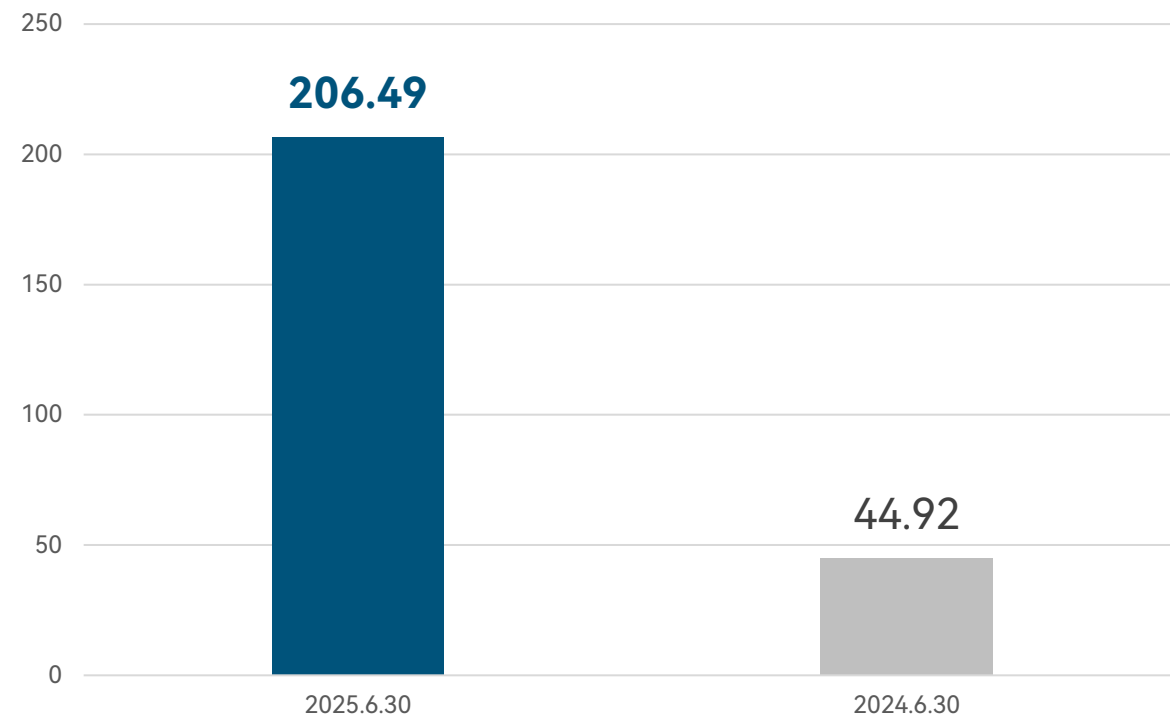
Income from CDMO & R&D services
RMB 22M +

04

Financial Results

Significant Operating Income Increase further Strengthened Capital Reserves

Mil RMB



Total Revenue in 2024 H1 and 2025 H1

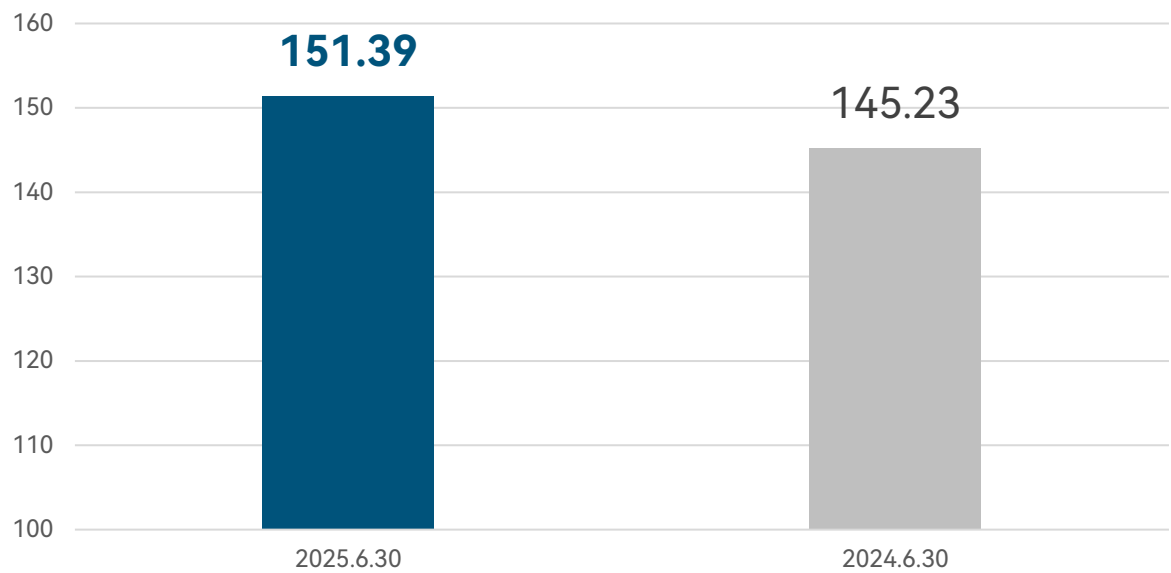
In 2025 H1, Qyuns recorded a total revenue of **RMB 206.49 million, representing 359.7% increase period-to-period.**

In 2025 H1, company's revenue mainly derived from upfront and milestone payments from licensing agreements, provision of R&D services, and CDMO services and supply of SAILEXIN.

(million RMB)	2025.6.30	2024.6.30
Revenue	206.49	44.92

Effective Expense Management Improves Operational Efficiency

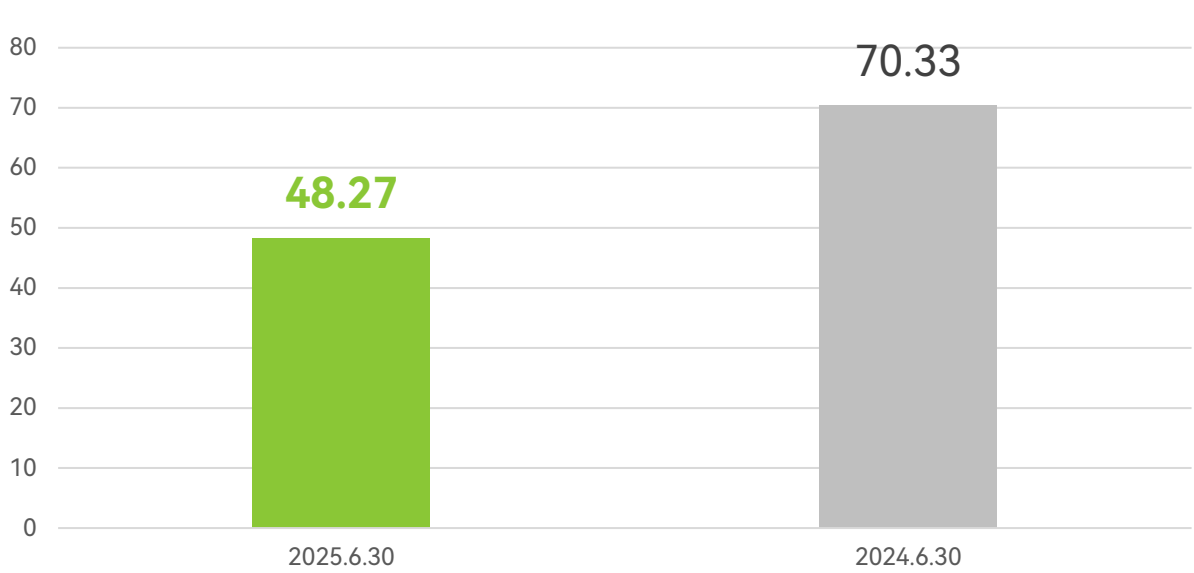
Mil RMB R&D Expenses of 2024 H1 – 2025 H1



In 2025 H1, Qyuns R&D expenses amounted to **RMB 151.39 million**, a net effect of decrease of RMB 5.87 million in amortization of equity-settled share-based payment and increase of **RMB 12.04 million** in R&D expenses primarily attributable to the increase in clinical trial costs resulting from advancement of clinical trials.

(Mil RMB)	2025.6.30	2024.6.30
R&D Expenses	151.39	145.23

Mil RMB Administrative Expenses of 2024 H1 – 2025 H1



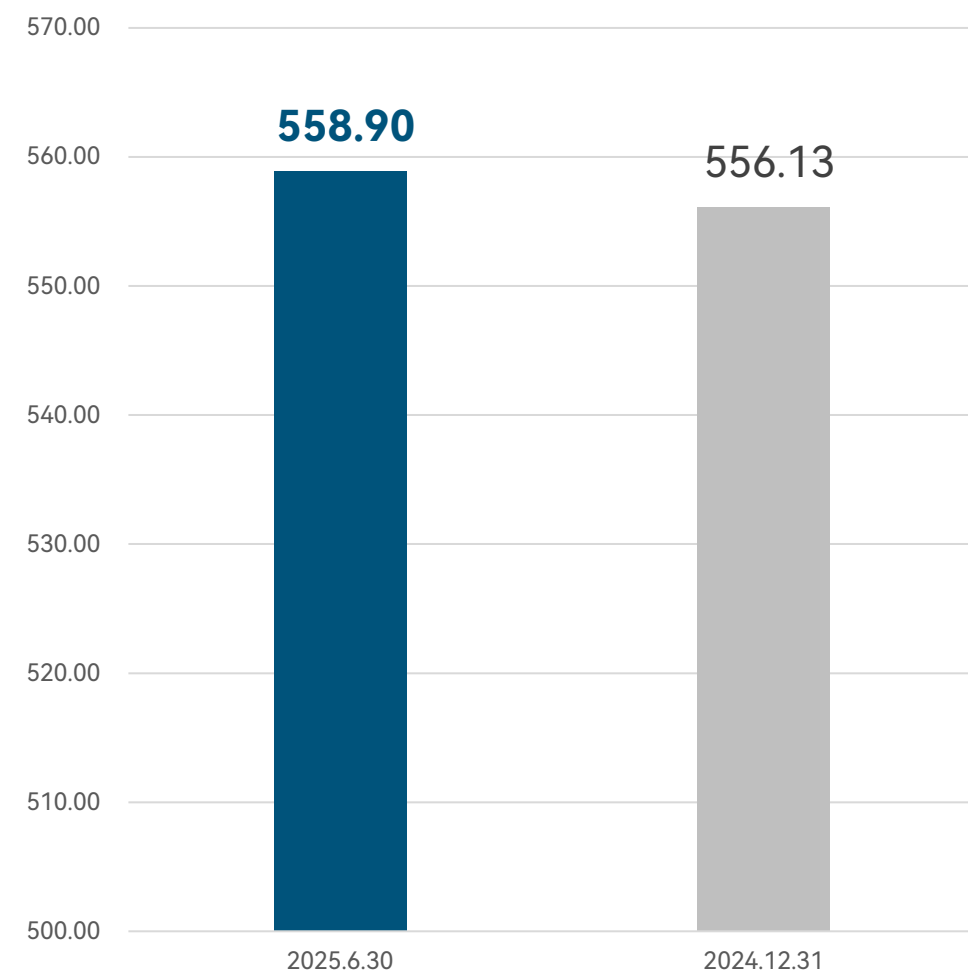
In 2025 H1, Qyuns incurred administrative expenses of **RMB 48.27 million**, **31.36% lower** than RMB 70.33 million in the prior-year period, primarily attributable to a decrease in equity-settled share-based payment expenses.

(Mil RMB)	2025.6.30	2024.6.30
Administrative Expenses	48.27	70.33

Cash Reserves Remain Stable with Optimized Loan Structure

Cash and cash equivalents, time deposits, and financial assets at fair value through profit or loss

Mil RMB



Stable cash reserves: As of June 30, 2025, Cash and cash equivalents, time deposits, and financial assets at fair value through profit or loss amounted to **RMB 558.90 million**, an increase of **RMB 2.77 million** compared to **RMB 556.13 million** as of December 31, 2024, **indicating stable cash reserves**.

Sufficient credit lines and optimized loan structure: As of June 30, 2025, the unutilized credit facility for working capital use available to the Company amounted to RMB180.73 million; as of June 30, 2025, the balance of working capital loan with terms of 2 to 3 years accounted for 74.9% of the total working capital loan balance (December 31, 2024: 39.1%).

(Mil RMB)	2025.6.30	2024.12.31
Cash and cash equivalents, time deposits, and financial assets at fair value through profit or loss	558.90	556.13
Total non-current assets	453.09	367.15
Total current assets	679.06	616.73
Total non-current liabilities	465.28	332.67
Total current liabilities	451.02	430.16
Net current assets	228.02	186.56
Total equity	215.83	221.05



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