

2024 1Q Results

May 2024



China's Leading Innovative Pharmaceutical Enterprise

R&D Capabilities

- 8 R&D platforms
- 5 R&D centres located in China & the U.S.
- ~ 2,000 R&D professionals
- ~ 300 R&D projects (~130 innovative projects)

Commercialization Capabilities

- **10,000+** professional sales personnel
- Covered 35,000+ medical institutions across the country, of which 2,900+ Class 3 hospitals (more than 90%), 7,000+ Class 2 hospitals (more than 70%), 26,000+ other terminals and 350,000+ drug stores
- Products exported to 114 countries/regions in 6 continents, including the U.S. and Europe; marketing centers established in the U.S., Germany and Brazil



Manufacturing Capabilities

- **10+** pharmaceutical production bases
- Nano formulation: 27 production lines built with production capacity of 20M doses/year; 2 production lines under construction with production capacity of 2M doses/year
- Biologics: fermentation capacity of 40,000L
- Chemical drugs: production capacity of OSD~30B tablets/year, production capacity of injection ~3B doses/year
- mRNA vaccines: GMP-compliant production plant has been built
- siRNA: 2 pilot scale production lines has been built; commercial scale production line is under construction

2024 1Q Highlights

R&D

1 new drug approvals:

• Mingfule(AIS): first approved in China in similar products, which is the second indication

18 IND approvals :

China 7 for the first indication 9 for additional indications

Results

- Revenue increased by 11.5% to RMB 8.98 B
- Underlying profit attributable to shareholders* (see page 6) increased by 11.6% to RMB 17.2 B

North America

2 for the first indication -JMT106 injection(GPC3/INF) -SYH2039 tablet (MAT2A inhibitor)



Part 01

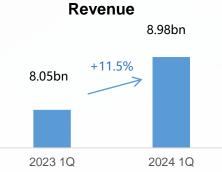
Financial Highlights



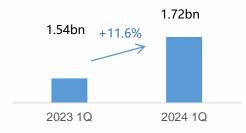
Financial Highlights

Unit: RMB' M

| | 2024 1Q | 2023 1Q | Change |
|---|---------|---------|--------|
| Revenue | 8,983 | 8,053 | +11.5% |
| Gross profit | 6,495 | 5,519 | +17.7% |
| Gross profit margin | 72.3% | 68.5% | +3.8% |
| R&D expenses | 1,169 | 1,008 | +16.0% |
| Underlying profit attributable to shareholders* | 1,724 | 1,545 | +11.6% |
| Reported profit attributable to shareholders | 1,613 | 1,429 | +12.9% |
| Basic earnings per share (RMB cents) | | | |
| Based on underlying profit attributable to shareholders | 14.55 | 12.96 | +12.3% |
| Based on reported profit attributable to shareholders | 13.61 | 11.99 | +13.5% |

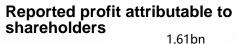


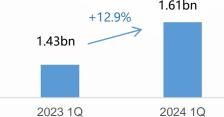
Underlying profit attributable to shareholders



*Note:

Underlying profit attributable to shareholders, a non-HKFRS measure, represents profit excluding fair-value changes on financial assets measured at FVTPL and employee share-based compensation expense.







Revenue by product category Unit: RMB' M Finished drug revenue

Unit: RMB 'M

License fee income

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| | 2024 1Q | 2023 1Q | Change | | | 2024 1Q | 2023 1Q | Change |
|-------------------------------|------------|---------|------------|---|------------------------|---------|---------|--------|
| Finished drugs | 7,561 | 6,422 | +17.7% | | Nervous system | 2,707 | 2,125 | +27.4% |
| | 486 | 551 | -11.8% | , | Oncology | 1,611 | 1,444 | +11.6% |
| Bulk vitamin C | | | 11.070 | | Anti-infective | 1,350 | 1,230 | +9.8% |
| Bulk antibiotics | 449 | 466 | -3.6% | | Cardiovascular | 720 | 589 | +22.3% |
| Functional Food and Others | 486 | 615 | -21.0% | | Respiratory system | 467 | 498 | -6.2% |
| | 100S Veals | | elpresente | | Digestion & metabolism | 314 | 196 | +60.0% |
| | | | | | Others | 392 | 304 | +28.5% |

Note: certain percentage changes of financial figures contained in this material are calculated based on the corresponding financial figures in RMB for two periods/years, rounded to the nearest thousand. Therefore, the percentage changes listed in certain tables may differ from those calculated based on the financial figures in RMB for two periods/years, which are presented in million.



Unit: RMB' M

| | 2024 1Q | 2023 1Q | Change | 2024 1Q OPM | 2023 1Q OPM | Change |
|-------------------------------|---------|---------|---------|----------------|----------------|--------|
| Finished drugs | 1,885 | 1,621 | +16.3% | 24.9% | 25.2% | -0.3% |
| Bulk vitamin C | 13 | 31 | -60.1% | 2.6% | 5.7% | -3.1% |
| Bulk antibiotics | 106 | 26 | +301.7% | 23.6% | 5.7% | +17.9% |
| Functional Food and Others | 121 | 144 | -15.7% | 24.9% | 23.4% | +1.5% |

Note: certain percentage changes of financial figures contained in this material are calculated based on the corresponding financial figures in RMB for two periods/years, rounded to the nearest thousand. Therefore, the percentage changes listed in certain tables may differ from those calculated based on the financial figures in RMB for two periods/years, which are presented in million.

Part 02

Business Review



Finished Drugs Overview by Therapeutic Areas

| | Nervous system | Major products include NBP (butylphthalide soft capsules, butylphthalide and sodium chloride injection), Shuanling (pentoxifylline extended-release tablets, pentoxifylline injection), Oulaining (oxiracetam capsules, oxiracetam for injection), Enxi (pramipexole dihydrochloride tablets), Enliwei (lacosamide injection, lacosamide tablets), and Oushuan (paliperidone extended-release tablets) |
|--|------------------------|--|
| Digestion & metabolism Others 4.2% 5.1% | Oncology | Major products include Duomeisu (doxorubicin hydrochloride liposome injection), Jinyouli (PEG- rhG-CSF injection), Keaili (paclitaxel for injection (albuminbound)) Duoenda (mitoxantrone hydrochloride liposome injection), Duoenyi (irinotecan hydrochloride liposome injection), Jinlitai (Narlumosbart injection), Copiktra (duvelisib capsules) and Geruite (lenvatinib mesilate capsules) |
| Respiratory Nervou system 6.2% System Cardi- Total : | Anti-infective | Major products include Anfulike (amphotericin B cholesteryl sulfate complex for injection), Shuluoke (meropenem for injection), Nuomoling (amoxicillin capsules), Xianqu (Ceftriaxone Sodium for Injection), Xianwu (Cefazolin Sodium for Injection), Zhongnuo Lixin(Cefuroxime Sodium For Injection), and Weihong (azithromycin tablets/capsules/enteric-coated tablets, azithromycin for injection) |
| vascular 9.5% (exclude License fee income) | Cardio- vascular | Major products include Xuanning (levamlodipine maleate tablets and dispersible tablets), Mingfule (recombinant human TNK tissue-type plasminogen activator for injection), Encun (clopidogrel bisulfate tablets), Daxinning (dronedarone hydrochloride tablets), Abikang (Aspirin enteric-coated tablets), Yishuning (nifedipine controlled-release tablets), and Meiluolin (Ticagrelor tablets) |
| Anti-infective 17.9% Oncology | Respiratory system | Major products include Yiluoda (nintedanib capsules), Qixin (oseltamivir phosphate capsules), Qixiao (arbidol hydrochloride tablets), Nuoyian (montelukast sodium tablets/chewable tablets), Zhongnuo Like (ambroxol hydrochloride oral solution) and Zhongnuoping (ambroxol hydrochloride extended-release tablets) |
| 21.3% | Digestion & metabolism | Major products include Debixin (omeprazole enteric capsules/tablets/injections), Linmeixin (glimepiride dispersible tablets), Shuanglexin (metformin hydrochloride tablets/extendedrelease tablets), Xinweiping (acarbose tablets), and Obeituo (Esomeprazole magnesium enteric-coated capsules) |
| 10 | Others | Major products include Oubida (Apgumilast tablets), Gujie (Tofacitib citrate sustained release tablets), Gubang (alendronate sodium tablets/enteric tablets), Xianpai (omeprazole sodium for injection) and Qimaite (tramadol hydrochloride tablets) |

Key Products Overview

NBP

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Butylphthalide soft capsules and injections

- 1st Class 1 new drug of cardiocerebrovascular field in China
- Price cut after negotiation improves affordability and accessibility, benefiting more patients
- Significant growth in OTC and E- channels
- New indication vascular dementia (VaD) under clinical trails

Mingfule

Recombinant human TNK tissue-type plasminogen activator for injection

- For thrombolytic therapy in patients with acute ischemic stroke
- For thrombolysis in patients with acute myocardial infarction within 6h
- Preferred thrombolytic drug recommended by authoritative guidelines such as "Chinese Expert Consensus on Pre-hospital Thrombolysis", "STEMI Rational Drug Use Guide", "2023 SIGN Clinical Management Guide", and "Chinese Guidelines for the Diagnosis and Treatment of Acute Ischemic Stroke"

Ouyuexin

Desvenlafaxine succinate extendedrelease tablets

- The third-generation antidepressant, the only two-channel drug that does not require dose-titration, easy to use with good safety
- "The second batch of encourage imitation drug catalogue" variety, China's first imitation, exclusive listing
- Approved in 2023, was included in the NRDL the same year, benefiting more depressed patients

Xuanning

Levamlodipine maleate tablets and dispersible tablets

- The first Chinese innovative antihypertensive drug fully approved by the U.S. FDA
- Has served 50 million hypertensive patients
 in China
- Recommended by authoritative guidelines such as "China Hypertension Prevention Guide" and "Guidelines for Rational Drug Use of Hypertension"

Anfulike

Amphotericin B cholesteryl sulfate complex for injection

- Exclusive product, obtained marketing approval in March 2021; was included in the NRDL in December 2021
- Have covered approx.1,600 hospitals
- Form of discoid compound, has a unique drug uptake and release mechanism, significantly decrease nephrotoxicity and increase dosage

Key Products Overview

Jinyouli

PEG-rhG-CSF

- 1st long-acting white blood cell booster drug in China
- Expanding coverage in major municipal hospitals and county-level markets
- Was included in the centralised procurement of the Guangdong Alliance, which consists of 11 provinces, and improving access to the drug would accelerate wider clinical use

Duomeisu

Doxorubicin Hydrochloride liposome injection

- Largest market share in China
- The first player who passed consistency evaluation

Duoenda

Mitoxantrone hydrochloride liposome injection

- Obtained marketing approval in January 2022, exclusive new preparation worldwide authorized by multinational patents, more potent and safer than mitoxantrone
- Was included in the NRDL through negotiation Dec. 2023, benefiting more patients
- Various clinical trails in solid tumors undergoing, blockbuster potential

Duoenyi

Irinotecan hydrochloride liposome injection

- First generic drug in domestic market
- In combination with 5-fluorouracil (5-FU) and leucovorin (LV), for the treatment of patients with metastatic pancreatic cancer after disease progression following gemcitabinebased therapy
- Recommended by domestic and foreign authoritative guidelines (NCCN/CSCO/CACA)

Jinlitai

Narlumosbart for injection

- The first IgG4 subtype fully human monoclonal antibody against RANKL obtaining marketing approval in the world
- Compared with denosumab, the Product has significant enhancement in uniformity and quality controllability, with favorable efficacy and safety profile
- New indications of tumor bone metastasis and osteoporosis are under development

Bulk Product Business, Functional Food and Others

Bulk vitamin C

- Major products: vitamin C, vitamin C - sodium, vitamin C calcium and granular vitamin C
- Sales of vitamin C products decreased by 11.8% to RMB486 million due to weakened demand

Bulk antibiotics

- Major products: 7-ACA (intermediate), cefazolin sodium, penicillin potassium, penicillin sodium, azithromycin, and ertapenem sodium
- Sales of antibiotic products remained stable, which decreased slightly by 3.6% to RMB449 million

Functional food and others

- Revenue of the functional food and other business decreased, mainly affected by the decrease in price of caffeine products
- The overall market share of caffeine products has exceeded 60%

Part 03 R&D Capability









R&D Centre

- 5 R&D centres located in China & the U.S.
- R&D expenses in 2024
 1Q: RMB1.17 B

Technology Platform

- 8 national science & technology qualifications
- 2 state key labs
- 8 R&D technology platforms

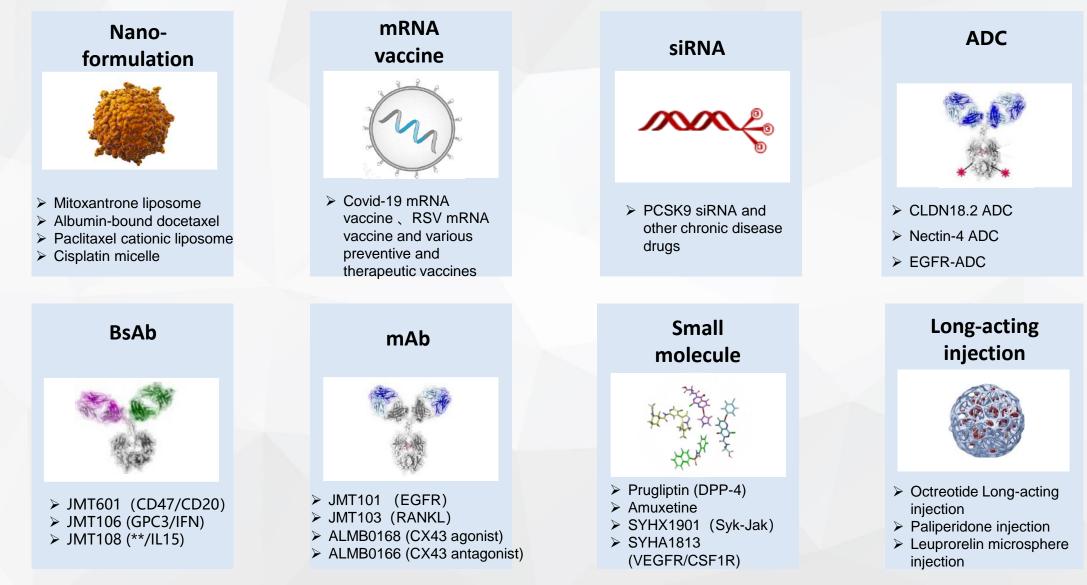
Project under Development & IPs

- Approx.300 projects under development (approx. 130 innovative drug projects)
- 1856 IPs applications
- 921 IPs authorised

Science Projects & Government Support

- 87 national projects
- RMB890M government
 grant support
- 8 national prizes

Innovative R&D Platforms



Note: only shows the representative products on each platform

Nano-formulation Platform

| Nano-formulation development and manufacturing platform | Novel drug carrier design | Novel drug delivery technology | Novel preparation method | Novel Industrialized production technology |
|---|--|---|---|---|
| • | Invented Albumin nanoemulsion Developed new cationic materials and new delivery system | Invented ammonium salt gradient method of sulfobutylether-β-cyclodextrin and 5-sulfosalicylate Cholesterol PEGylation modification method and post single layer PEGylation | Invented single-phase solution lyophilization technology, O/W type Emulsification technology, crossflow mixing technology, continuous flow reaction technology, etc. Invented bottom up nanocrystal preparation technology, enabling continuous production | Invented continuous flow technology, employing linear amplifier, overcome barriers to industrialized production Illustrated that all nano drugs are able to be prepared by permutation and combination of four key processes |
| Nano-formulation assessment system | rticle characterisation method | PK determination method | Mature animal screening models | Particle characterisation technique guided in vivo PK、PD、TOX evaluation |
| | Developed nano- formulation assessment technology for lipsome, albumin nanoparticles, emulsion, micelles, etc. | Established multiple PK determination methods for nano drugs including lipsome, albumin nano- particle, micelles etc. | Established multiple animal disease model for efficacy assessment Established animal models for evaluating ABC phenomenon、CARPA response and HFS, enabling quick screening | Illustrated influence of drug release rate of lipsome, mode of administration and animal model on ABC phenomenon Detailed study of CARPA and HFS laid the foundations for rational design of nanoparticles |

The largest R&D and industrialization base for Nano- Formulation

mRNA Vaccine platform

Advantages of antigen design

- Mutation prediction platform
- The combination of bioinformatics and structural biology to obtain effective epitopes
- Superior immunogenicity from sitespecific mutation of antigen

No observed SAE in clinical trials

Base modification mitigates innate

launched products

immunogenicity

Excellent safety profile

• Excipients proven to be low toxicity by

Formulation ensures long-term stability



- Base modification, UTR screening, codon optimization and structural elements inclusion
- Structural energy optimization to enhance antigen expression

5 Streamlined CMC Strategy

- One-step API manufacturing process
- API purification process: up to 99%
 purity
- Highly scalable LNP manufacturing process
- Short turnaround time: ~2 days

6 Highly expandable platform

Manufacturing capabilities

Manufacturing capabilities of CSPC

Manufacturing capacity reaches to 1.5

Top tier LNP R&D platform

billion doses per year

- Each individual component can be continuously upgraded
- Expansion from linear mRNA to circRNA; from liver-target delivery to extrahepatic delivery
- From preventive to therapeutic application; from vaccine to CGT



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siRNA Platform



- Rational sequence design based on bioinformatics and experienced scientists
- Comprehensive in vitro and in vivo PK/PD characterization

4 Excellent safety profile

- Build off-target risk assessment platform
- Chemical modification to mitigate immunogenicity
- Long-term stability



- Build strong oligonucleotides CMC platform based on QbD strategy
- Develop liquid synthesis technology

5 Nucleotides building blocks

- Develop novel building blocks
- Develop Galnac molecule with inhouse IP
- Scalable building blocks
 manufacturing technology

3

Manufacturing capabilities

- Manufacturing capabilities of CSPC
- Two GMP-compliant production lines have been built

6 Highly expandable platform

- Each individual component can be continuously upgraded
- Integrated manufacturing capabilities for building blocks, API and drug products









1 Payload platform

Multiple payload types for different cancer treatment mechanism of actions

- microtubule inhibitor
- Topoisomerase inhibitors
- Protein degrading agent
- DNA alkylating agent
- DNA repair inhibitors
- Immunoagonists
- Immunosuppressants
- hormone

3 Conjugation platform

Including mainstream conjugation technology and proprietary site-specific conjugation technology with IP rights.

- Conventional lysine and cysteine conjugation platforms
- SmartQTag enzyme-mediated conjugation technology
- Glycan-engineered site-specific conjugation technology

2 Linker platform

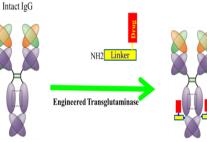
Various linker design for stability, PK and other purpose benefical for ADCs

- Hydroflex platform ensures hydrophilicity, stability and PK
- TMEC the TME payload released platform overcomes the problem of non-internalize targets
- Novel ESIM self-immolation platform enhances drug release efficiency.
- The HighDAR platform makes it possible to use low potent payloads in ADC

4 New molecular entity

Multiple ADC drug molecules provide more clinical treatment options, and overcome the drug resistance of current existing therapies.

| ISAC Immune-stimulating antibody conjugate | DAC Degrader-antibody conjugate | dpAC Dual-paylood anfibody conjugate | CA-ADC Conditional-activated-ADC | Bs-ADC Bispecific-ADC | PDC Peptide-drug conjugate |
|--|---------------------------------------|--|---|--|---|
| Immune-stimulator | Protein degrader | Synthetic lethal | pH-activated mAb | Dual-epitope | Peptide |
| | | • | | | 0-• |
| | Immune-stimulating antibody conjugate | Immune-stimulating antibody conjugate Degrader-antibody conjugate | Immune-stimulating antibody conjugate Degrader-antibody conjugate Dual-payload antibody conjugate | Immune-timulating antibody Degrader-antibody conjugate Dual-pol/oad antibody Conjugate Conditional-activated-ADC | Immune-stimulating antibody Designate antibody conjugate Designate antibody conjugate Constituand-activated -ADC Magnetic-ADC Immune-stimulator Protein degrader Synthetic lethal pH-activated mAb Dual-epitope |



Bispecific Antibody Platform

Antibody-interferon fusion protein platform

Structural advantages

Synergistic binding effect when targeting the same cell
 Smaller molecular weight (smaller than that of

The activity of interferon-containing impurities is far lower than that of conventional bispecific antibodies.
No serious safety risk of interferon-containing impurities

Stronger target selectivity

The optimizing of interferon mutation further improves

More stable product

Significantly reduces the interferon breakage during the

production and in vivo, resulting in high yield and low risk

Limited binding activity to receptors on TAA – cells,

demonstrating a wider therapeutic window

conventional antibodies) Safer impurities

from the production process

More efficient in CMC development

target selectivity

- High expandability : quickly forms new molecules
 - Better safety profile : much safer than interferon
 - More effective in tumor killing: much stronger than interferon
- remodels the tumor immune micro environment: enhances immune cell

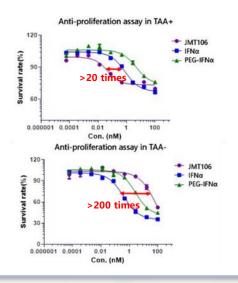
infiltration and MHCI presentation

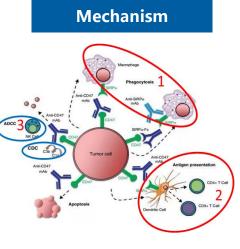
Lower production cost : high efficiency, low toxicity, and simple process



Left-arm: high-affinity, tumortargeting semi-antibody Right-arm: Low-affinity SIRPα-Fc fusion protein

- Does not bind to TAA-/CD47+cells, including erythrocytes, platelets, etc.
- TAA-dependency enhances CD47 competitive binding
- High expandability: various types of tumor-targeting antibodies could be used as the left-arm
- ✓ Wider safety window
- Lower molecular weight, better suits solid tumors
- Simple production process
- Possession of intellectual property right





hole



Candidates under Clinical Trial Stage

| | (26) | | (1 | 5) | (1 | 8) | (7) |
|------------------------|-------------------------|----------------------|----------------------------|--------------------------------------|---------------------------------------|---|---------------------------------------|
| > | Phl | | Phll | (POC) | Phll / III piv | votal trial | NDA |
| NBL-012 IL23-P19 | NBL-015 CLDN18.2 mAb | NBL-020 TNFR2 | CLDN18.2 ADC | CM326 TSLP | JMT101 EGFR mAb | CM310 IL4R | SYSA1802 PD-1 |
| NBL-028 CLDN6-CD137 | SYS6002 Nectin-4 ADC | SYS6010 EGFR ADC | ALMB0166 Cx43i mAb | ALMB0168 Cx43s mAb | KN026 Her2 BsAb | TG103 Fc-GLP1 | Omalizumab biosimilar |
| SYS6011 | JMT203 GFRAL | Secukinumab | ЈМТ601 CD20/CD47 | SYHX1901 JAK/SYK | Pertuzumab | Ulsinumab | DBPR108 DDP4 |
| SYHA1801 BRD4 | SYHA1803 Pan-FGFR | SYHA1805 FXRs | Simmitinib TKI | SYHA1813 VEGFR/CSF1R | DP303C HER2 ADC | NBP Capsule (VaD) | Amphotericin B Liposome |
| SYHA1807 LSD1 | SYHA1811 BTK | SYHA1815 FGFR/RET | Amuxetine 5-HT/NE | SYHA1402 ARi | Batoclimab | Semaglutide | Meloxicam nanocrystal injection |
| SYHX1903 CDK9 | SYHX2001 PRMT5 | SYHX2005 FGFR4 | NBP Capsule (US PhII) | Paclitaxel cationic liposome | JMT103 bone metastasis | Pregabalin extended-release tablets | Irinotecan liposome (US) |
| SYHX2009 NTRK/ROS1 | SYH2038 SOS1 | SYH2043 CDK2/4/6 | Albumin-bound Sirolimus | Octreotide long- acting injection | Albumin-bound Paclitaxel II | Pilocarpine hydrochloride eye drops | Amphotericin B Liposome (US) |
| SYH2045 PRMT5 | SYH2051 ATM | SYH2053 PCSK9 | Alprostadil liposome | | Clevidipine injectable emulsion | Daunorubicin cytarabine liposome | |
| SYHA1908 | | Cisplatin micelle | | | Albumin-bound Docetaxel | Mitoxantrone hydrochloride liposome (NPC) | Biological Age |
| | | | | | | | Chemical Dru |
| | | | | | | | New formula |

Pipeline - Biological Agents

Over 40 new biologic drugs under development: 2 filed BLA, 23 under clinical trial stage(9 under pivotal trial stage) and over 10 under pre-clinical stage

| ТА | Major Candidates | Target | Indication (s) | Pre- clinical | Phase I | Phase II | Ph II / III NDA pivotal trial |
|----------|-----------------------|------------------|-------------------------------------|------------------|---------|----------|----------------------------------|
| | JMT103 | RANKL | GCTB, osteoporosis, bone metastasis | | | | Marketed(GCTB) |
| | SYSA1802 (SG001) | PD-1 | Tumors | | | | BLA |
| | JMT101 | EGFR | Multiple solid tumors | | | | |
| Oncology | DP303c | HER2 ADC | Breast cancer | | | | |
| | Pertuzumab biosimilar | HER2 | Breast cancer | | | | |
| | KN026 | HER2 BsAb | Gastric cancer, breast cancer | | | | |
| | SYSA1801* | Claudin 18.2 ADC | Gastric cancer, pancreatic cancer | | 1 | | |
| | ALMB0168 | CX43 agonist | Bone cancer, cancer bone metastasis | | | | |
| | JMT601* | CD47/CD20 | NHL& multiple hematologic tumors | | | | |
| | SYS6002* | Nectin-4 ADC | Tumors | | | | |
| | NBL-028 | CLDN6-CD137 | Advanced tumors | | | | |
| | NBL-020* | TNFR2 | Advanced solid tumor | | | | |
| | SYS6010* | EGFR ADC | Tumors | | | | |
| | SYS6011 | Undisclosed | Solid tumors | | | | |
| | NBL-015* | Claudin 18.2 mAb | Advanced solid tumors | | | | |
| | JMT203 | GFRAL | Cancer cachexia | | | | |

*The product was developed in both China and the U.S.

Pipeline - Biological Agents

| ТА | Major Candidates | Target | Indication (s) | Pre-clinical | Phase I | Phase II | Ph II / III pivotal trial | NDA |
|------------------------|-----------------------|-----------------|-------------------------------------|--------------|---------|-----------------------|------------------------------|-----------------------|
| Nervous system | ALMB0166 | CX43 antagonist | Spinal cord injury, AIS | | | | | |
| Digestion & metabolism | TG103 | GLP-1 | Obesity, Diabetes | | | 1 1 1 | | |
| | Omalizumab biosimilar | IgE | Allergic asthma, CIU | | | | | BLA |
| | Ulsinumab | IL-12/IL-23 | Moderate to severe plaque psoriasis | | | | | |
| | Batoclimab | FcRn | MG | | | | | |
| Immune | CM310 | IL-4Rα | Asthma, COPD | | | | | |
| mmune | CM326 | TSLP | Asthma, COPD | | | | | |
| | NBL-012* | IL-23p19 | Psoriasis, HS, IBD | | | 1 1 1 1 1 | | 1 1 1 1 1 |
| | Secukinumab | IL-17A | Psoriasis | | | | | 1 1 1 1 |

Pipeline - Chemical Drugs

Over 40 small molecule new drugs under development: 1 filed NDA, 23 under clinical trial stage (4 under pivotal trial stage) and about 20 under pre-clinical stage

| ТА | Major Candidates | Target | Indication (s) | Pre-clinical | Phase I | Phase II | Ph II / III pivotal trial | NDA |
|----------------|---|--------------------|---|--------------|---------|----------------|------------------------------|-------|
| | DBPR108 | DPP-4 | Diabetes | | | | | ★ NDA |
| Digestion & | Semaglutide | GLP-1 | Type2 diabetes | | | | | |
| metabolism | SYH2053 | PCSK9 | FH, Mixed hyperlipidemia | | | | | |
| | SYHA1805 | FXR Agonist | NASH(MASH) | | | | | |
| Ophthalmology | Pilocarpine hydrochloride eye drops | AChR | Presbyopia | | | | | |
| | NBP soft capsule* | | VaD, Ischemic stroke(US) | | | | | |
| Nervous system | Pregabalin extended- release tablets | γ-GABA analogue | Neuropathic pain associated with diabetic peripheral neuropathy | | | | ▶ | |
| | SYHA1402 | ARI | Diabetic neuropathy | | | | | |
| | Amuxetine hydrochloride enteric tablets | 5-HT, SNDRI | Anti-depressant | | | | | |
| Immune | SYHX1901 | Syk-Jak | RA, SLE, COVID-19 | | | | | |
| | Simmitinib | FGFR、KDR | Gastric cancer, cholangiocarcinoma, SQCC | | | | | |
| | SYHA1813 | VEGFR/CSF1R | Relapsed or advanced solid tumour | | | | | |
| Oncology | SYHX2005 | FGFR4 | Advanced solid tumor | | | | | |
| | SYH2043 | CDK2/4/6 | Breast cancer | | | | | |
| | SYH2045 | PRMT5 | Advanced malignant tumors | | | | | |
| 26 | SYH2051 | ATM | Solid tumor | | | | | |

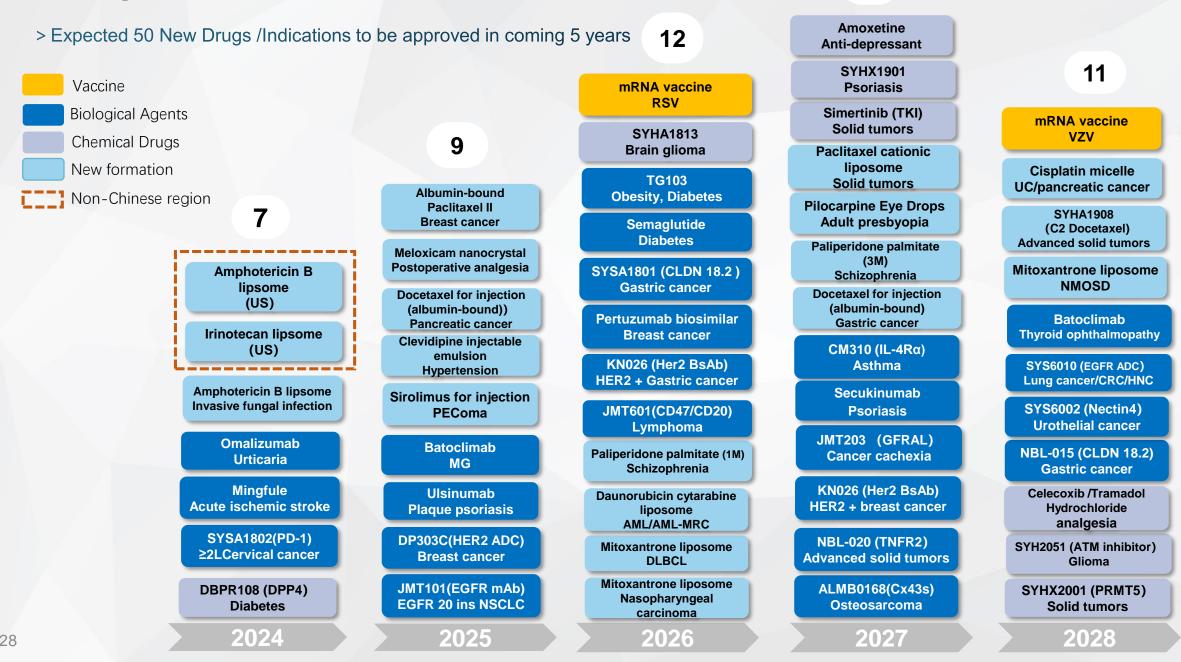
Pipeline - New formulations

Over 30 new preparations under development: 2 flied NDA, 11 under clinical trial (5 under pivotal trial stage), and over 20 under pre-clinical stage

| ТА | Major Candidates | Indication(s) | Pre-clinical | Phase I | Phase II | Phase III / Pivotal Clinical Trial | NDA |
|-----------------|---|---|--------------|----------------|----------|--|---------------------|
| Oncology | Mitoxantrone hydrochloride liposome injection * | Multiple hematologic neoplasms & solid tumors | | | | 7 | Marketed((PTCL) |
| | Irinotecan liposome injection* | pancreatic cancer, Breast cancer | | | | | Marketed |
| | Daunorubicin cytarabine liposome | Leukemia | | | | | (pancreatic cancer) |
| | Albumin-bound Paclitaxel II | Multiple solid tumors | | | | | |
| | Docetaxel for injection (albumin-bound) * | Gastric cancer, pancreatic cancer | | | | | |
| | Paclitaxel cationic liposome | Advanced solid tumors | | | | | |
| | Sirolimus for injection (albumin-bound) | Multiple hematologic cancers & solid tumors | | | | | |
| | SYHA1908 for injection (class 1 new drug+nano drug) | Solid tumors | | | | | |
| | Cisplatin micelle | Multiple solid tumors | | | | | |
| Analgesia | Meloxicam nanocrystal injection | Moderate-to-severe pain | | | | | 📩 🕇 NDA |
| Anti-infective | Amphotericin B liposome * | Invasive fungal infection | | | | | NDA 🛧 |
| Cardio- | Clevidipine injectable emulsion | Hypertension | | | | | |
| cerebrovascular | Alprostadil liposome | Vasodilation | | | | | |
| Endocrinology | Octreotide long-acting injection | Acromegaly | | | | | |

Pipeline Products Launch Plan

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Common Generics Launch Plan

17 applications have been filed for marketing approvals, expecting to receive approval in 2024-2025; Over 20 candidates under pharmaceutical research, expecting to receive approval in 2026-2027

| No. | Product | Therapeutic Area | Expected to be launched | Launched or not |
|-----|--|---------------------------|-------------------------|-----------------|
| 1 | Dapagliflozin tablets | Digestion & Metabolism | 2024 | \checkmark |
| 2 | Rabeprazole sodium enteric-coated tablets (10mg) | Digestion & Metabolism | 2024 | \checkmark |
| 3 | Olaparib tablets | Oncology | 2024 | |
| 4 | Palbociclib tablets (125mg) | Oncology | 2024 | |
| 5 | Lenalidomide capsules (5mg, 10mg) | Oncology | 2024 | |
| 6 | Peramivir injection (150mg/15ml) | Anti-infective | 2024 | \checkmark |
| 7 | Aprepitant injection | Others | 2024 | |
| 8 | Dexrazoxane for injection | Others | 2024 | |
| 9 | Roxadustat capsules | Others | 2024 | |
| 10 | Peramivir Injection (300mg/60ml) | Anti-infective | 2025 | |
| 11 | Regorafenib tablets | Oncology | 2025 | |
| 12 | Ilaprazole enteric-coated tablets | Digestion & Metabolism | 2025 | |
| 13 | Terezolamide phosphate tablets | Anti-infective | 2025 | |
| 14 | Oseltamivir phosphate for oral suspension | Anti-infective | 2025 | |
| 15 | Mesalazine enteric-coated tablets | Autoimmunity | 2025 | |
| 16 | Vonorazone fumarate tablets | Digestion & Metabolism | 2025 | |
| 17 | Adenosine cobalamin capsules | Others | 2025 | |

IND Approvals Obtained since the beginning of 2024

| IND approval for the 1st indication (7+2) | |
|--|---|
| JMT202 (mAb) | Lower triglyceride (TG) levels in patients with hypertriglyceridaemia |
| SYS6023 (ADC) | Advanced solid tumors |
| SYH2039 tablets (MAT2A) | Advanced malignant tumors |
| Dexmedetomidine hydrochloride nasal spray | Sedation before invasive procedures |
| Pilocarpine hydrochloride eye drops | Presbyopia |
| Pregabalin extended-release tablets | Neuropathic pain associated with diabetic peripheral neuropathy |
| Semaglutide injection | Weight management |
| JMT106 injection (GPC3/IFN) -US | Advanced solid tumors |
| SYH2039 tablets (MAT2A enzyme inhibitor) -US | Advanced malignant tumors |

IND Approvals Obtained since the beginning of 2024

| IND approval for additional indications (9) | |
|---|---|
| SYSA1801 injection | In combination with CAPOX and SG001 or with irinotecan hydrochloride liposome injection for first-line and second-line treatment of Claudin18.2-positive gastric cancer |
| JMT101 injection | In combination with docetaxel (albumin-bound) for treatment of second-line and above EGFR lung squamous cell carcinoma |
| Simmitnib hydrochloride tablets | In combination with irinotecan liposome for the treatment of second- line advanced esophageal cancer |
| Sirolimus for injection (albumin-bound) | In combination with endocrine therapy for the treatment of second- line and above HR-positive HER2-negative advanced breast cancer after failure of standard therapy |
| Sirolimus for injection (albumin-bound) | In combination with gumetinib for the treatment of locally advanced/recurrent or distant metastasis MET-overexpression non- small cell lung cancer |
| SYH2043 tablets | In combination with fulvestrant for the treatment of advanced breast cancer |
| Cisplatin micelle injection | In combination with paclitaxel for the treatment of advanced solid tumors |
| Octreotide long-acting injection | Gastroenteropancreatic neuroendocrine tumors |
| Irinotecan liposome injection | In combination with oxaliplatin and tegafur for adjuvant treatment of pancreatic cancer |



BD Strategic Layout and Path of Advancement

Focusing on strategic domains, deepening BD strategies, and establishing an international BD ecosystem

BD Product Positioning: Aligning closely with clinical needs, emphasizing clinical benefits, grasping international cutting-edge technology and product trends, strengthening areas of group advantage, focusing on key clinical stage products in the mid to late phases.

BD Technology Platforms: Actively exploring collaboration and development of early-stage products with AI pharmaceuticals, nucleic acid drug antigen screening platforms, gene therapy technologies, novel vaccine development platforms, and intratumoral injection technology platform.

Internationalization of BD: Pursuing a dual strategy of both licensing in and out, expanding international projects with leading multinational pharmaceutical companies and Belt and Road initiatives, reinforcing strategic relationships with fund institutions having overseas resources, and advancing the connection and collaboration of global projects.

BD Ecosystem Construction : BD Ecosystem Construction: Leveraging the advantages of group clinical development, product registration, and commercialization resources, adopting a Pharma+Biotech win-win model, engaging in extensive and in-depth collaboration with Biotech companies or research institutions that possess innovative advantages, including practical and feasible merger and acquisition models, to continue supporting the group's external innovation.

BD Work Completion Status for 2023

License-out:

- Granted Corbus Pharmaceuticals the development and commercialization rights for SYS6002 (Nectin-4ADC, Phase 1) in the United States, European Union countries, the United Kingdom, Canada, Australia, Iceland, Liechtenstein, Norway, and Switzerland.
- License-in:
 - Obtained Pfizer's exclusive authorization to locally market the oral antiviral COVID-19 treatment medication, Namatavir Tablets/Litoconavir Tablets, in China



BD Key Therapeutic Area Strategy for 2024

Reinforce Leading Position in Established Areas

Comprehensive Management of Stroke Disease, with a Focus on the Strategic Positioning and Collaboration of Innovative Drug Projects in Vascular Recanalization. Neuroprotection. and Anti-Inflammation that Synergize with the Company's Existing Resources

Attention to Late-stage Clinical or Newly Approved Drugs in the Alzheimer's Disease (AD) Field, as well as Emerging Novel Targeted Therapeutics.

Neurology Field

Strengthen differentiation in hematologic malignancies, lung cancer, and breast

- cancer, focusing on targeted therapies,
- new immunotherapies, and combination
- treatments.
- Explore innovative drugs in areas such as digestive tract tumors, gynecological tumors, and urological tumors.

Oncology Field

· Focus on challenging areas like refractory hypertension, hyperlipidemia, and heart failure.Pay attention to long-acting, oral diabetes/weight reduction innovative products.

 Address thyroid diseases and innovative treatments related to gout.

Cardiovascular & Endocrinology Field

drug-device combinations, and drug delivery systems.

> Address conditions such as atopic dermatitis, systemic lupus erythematosus,

Respiratory, Autoimmune & Anti-Infective Field



- Emphasize areas like idiopathic pulmonary fibrosis (IPF), COPD/asthma. and cough, exploring innovative targets,
- Focus on high-end antibiotics effective against clinically resistant bacteria.
- and inflammatory bowel disease (IBD).

Explore Novel Therapeutic Areas and Technology Platforms

 Address primary and secondary kidney diseases like IgA nephropathy and diabetic nephropathy. Focus on complications of kidney diseases like renal anemia, hyperphosphatemic kidney disease, hypertension,

and kidney-related itching.

Nephrology Field

· Expand into major populationbased psychiatric disorders like depression and schizophrenia, focusing on the layout and collaboration of novel targeted drugs with improved efficacy, safety, and compliance. Emphasize fast-acting nasal

spray formulations.

Psychiatry Field

 Concentrate on wellestablished companies with mature late-stage ophthalmology pipelines. Focus on products for treating retinal diseases like AMD using new targets, long-acting formulations, nanomedicines, and gene therapies, with a special focus on geographic atrophy indications.

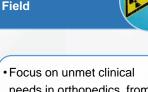
Nuclear Medicine Ophthalmology Field

 Chronic Pain: Focus on innovative drug projects that provide better pain relief, higher safety, and nonaddictive properties.

 Acute Pain: Concentrate on innovative projects that extend postoperative pain relief duration while maintaining higher safety.

Pain Management Field

• Focus on therapeutic nuclear medicine, breaking through new targets, new indications, and new isotopes while avoiding homogenization. Focus on larger market opportunities in solid tumor such as prostate cancer, lung cancer, breast cancer, gastrointestinal tumors under new nuclides and new target combination



needs in orthopedics, from completely innovative products to products that improve patient accessibility, and tap opportunities in orthopedics

 Focus on spine surgeryrelated drugs, osteoporosis iterations

Orthopedics Field



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Aim to Become an ESG Leader in Pharmaceutical Industry

- Awarded "China's Top Healthiest Workplace" by Mercer, "AAA Enterprise with Harmonious Labour Relations in Hebei Province" and "National Advanced Enterprise in Employment"
- Achieved "Five Zeros and One Low"*
- The major shareholder of the Group granted 220M conditional shares to over 300 employees in 2022
- Improving board diversity continuously

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- Structural reduction of carbon emissions the ratio of innovative drugs /formulations increasing and the ratio of APIs decreasing
- Invested RMB763m in smart manufacturing, equipment upgrade and modification
- Plan to invest more than RMB100m per year to support the upgrade of environment protection center
- Ouyi, NBP, CSPC Innovation, Factory in Taizhou (our subsidiaries) are recognized as "Green Factories" by the MIIT

People orientated, Win-win Future

> Environmentallyfriendly, Carbon Emissions Reduction

- Adhere to the procurement principle of "fair, impartial, green and transparent"
- Online bidding and procurement; supplier integrity commitment; Blacklist Management System for Dishonesty
 - Employee assistance: In 2023, a total of 81 employees and their families were helped
 - CSPC Education Assistant Fundhelped 2024 college students in 2023
 - Cancer and critical illness patients assisting program- assisted 42 patients in 2023
 - Medical care program for poor children- helped 42 children with hemopathy in 2023
 - Patient assisting program: the number of aid recipients reached 59300 in 2023.

Continuous MSCI ESG rating upgrade

Sep-19 Aug-20 Apr-21 Dec-21 Nov-22 Oct-23

Environmental Protection Plan 2025

- ✓ Reduce greenhouse gas emissions per unit of revenue by 50.66%
- Reduce the emission of non-hazardous waste (general solid waste) per unit of revenue by 70.7%
- Reduce the discharge of hazardous waste per unit of revenue by 26%
- *The emission reduction target is based on the emission in 2017

- Reduce the comprehensive energy consumption by 56.57%
- Reduce the water consumption per unit of revenue by 35.52%

*Five Zeros and One Low- zero cases of death, serious injuries, multiple injuries, occupational disease and poisoning incident as well as low incident rate of minor injuries

WeChat of CSPC IR Team:



Thanks!

SUPPORT

GUIDANCE