

2023 FY Results

March 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 staff
- ~ 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

2023 FY Highlights

5 new drug approvals:

Mingfule(AIS)

- R&D ·
 - Covid-19 mRNA vaccine (EUA)
 - Narlumosbart for injection
 - Irinotecan liposome injection
 - COVID-19 bivalent mRNA vaccine (EUA)

7 applications for marketing approval :

China

- Amphotericin B liposome
- Prugliptin tablets (DPP-4)
- Omalizumab for injection
- Meloxicam nanocrystal injection
- Enlonstobart for injection

North America

- Amphotericin B liposome for injection
 - Irinotecan liposome injection

Business

Bu

46 IND approvals :

China

17 for the first indication24 for additional indications

• Re **North America** 5 for the first indication

- Revenue increased by 1.7% to RMB31.45B
- Underlying profit attributable to shareholders* (see page 6) increased by 2.8% to RMB6.28B

BD

- Nectin-4ADC: licensed-out the rights in the US, EU, UK, Canada, Australia, Iceland, Liechtenstein, Norway and Switzerland to Corbus, with US\$7.5M upfront payment, potentially US\$685M milestone payment and royalty.
- Obtained the exclusive promotion rights of Glumetinib (c-MET inhibitor) from Haihe Biopharma. The product has been approved for marketing in March 2023
- Signed a strategic partnership agreement with Pfizer to launch a local brand of the COVID-19 oral therapeutic treatment Nirmatrelvir/Ritonavir in China







Part 01

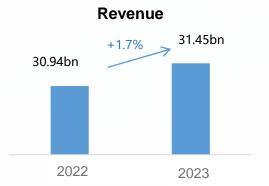
Financial Highlights



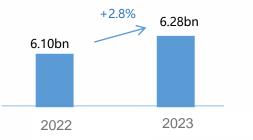
Financial Highlights

Unit: RMB MM

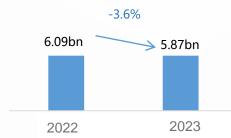
	2023	2022	Change
Revenue	31,450	30,937	+1.7%
Gross profit	22,177	22,256	-0.4%
Gross profit margin	70.5%	71.9%	-1.4%
R&D expenses	4,831	3,987	+21.2%
Underlying profit attributable to shareholders*	6,275	6,106	+2.8%
Profit attributable to shareholders	5,873	6,091	-3.6%
Basic earnings per share (RMB cents)			
 Based on underlying profit attributable to shareholders 	52.86	51.23	+3.2%
Based on profit attributable to shareholders	49.47	51.11	-3.2%



Underlying profit attributable to shareholders



Profit attributable to shareholders



*Note:

Underlying profit attributable to shareholders, a non-HKFRS measure, represents profit excluding fair-value losses on financial assets recognised in profit or loss at fair value, share-based employee remuneration expenses and gains on deemed disposal of partial interest in an associate and a joint venture.



Revenue by product category Unit: RMB MM

	2023	2022	Change
Finished drugs	25,637	24,520	+4.6%
Bulk vitamin C	1,929	2,529	-23.7%
Bulk antibiotics	1,712	1,503	+13.9%
Functional Food and Others	2,172	2,385	-8.9%

Finished drug revenue Unit: RMB MM

2022 Change 2023 Nervous system 9,089 8,108 +12.1% Oncology 6,139 7,340 -16.4% **Anti-infective** 3,540 4,236 +19.7% Cardiovascular 2,440 2,889 -15.5% **Respiratory system** 1,560 696 +124.0% Digestion & metabolism 889 755 +17.8% Others 1,249 1,006 +24.1% License fee income 186 -81.3% 35



Unit: RMB MM

	2023	2022	Change	2023 OPM	2022 OPM	Change
Finished drug	6,700	6,068	+10.4%	26.1%	24.7%	+1.4%
Bulk vitamin C	5	443	-98.9%	0.3%	17.5%	-17.3%
Bulk antibiotics	154	114	+35.2%	9%	7.6%	+1.4%
Functional Food and Others	561	647	-13.3%	25.9%	27.1%	-1.2%

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 Drug 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 3.5% 4.9%	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 6.1% Cardi- 共计: Nervous 35.5%	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% (不含授权费 收入)	Cardio- vascular	 Major products include Xuanning (levamlodipine maleate tablets and dispersible tablets), Mingfule (recombinant human TNK tissue-type plasminogen activator for inj`ection), Encun (clopidogrel bisulfate tablets), Daxinning (dronedarone hydrochloride tablets), Abicang (Aspirin enteric-coated tablets), Yishuning (nifedipine controlled-release tablets), and Meiluolin (Ticagrelor tablets)
Anti-infective 16.5% 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)
24%	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 national talks in 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 23.7%, mainly due to the weakening price of vitamin C products

Bulk antibiotics

- Major products: 7-ACA (intermediate), cefazolin sodium, penicillin potassium, penicillin sodium, azithromycin, and ertapenem sodium
- Sales of antibiotic products increased by 13.9%, driven by the growth in sales volume

Functional food and others

- During the period, there was a certain decline in the price of caffeine products, while the sales volume maintained a stable growth
- 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 FY 2023: RMB 4.83 B

Technology Platform

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

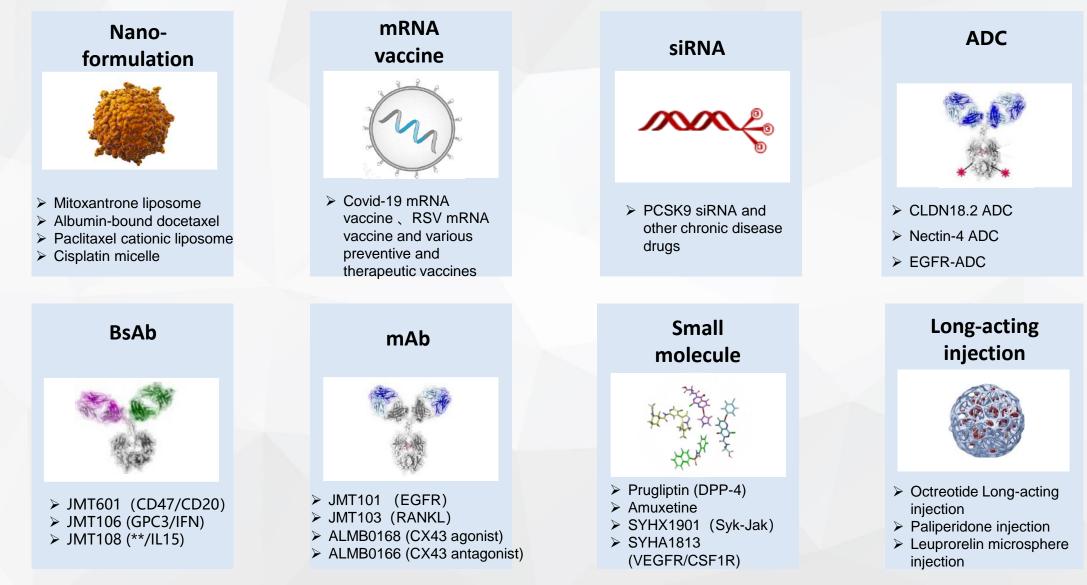
Project under Development & IPs

- Approx.300 projects under development (approx. 130 innovative drug projects)
- 1806 IPs applications
- 906 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

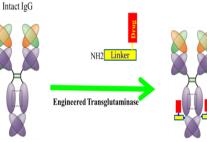
Various linker design for stability, PK and other purpose benefical for ADCs

- 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

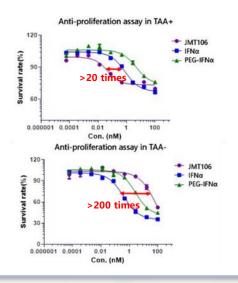
infiltration and MHCI presentation

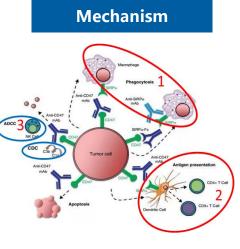
Lower production cost : high efficiency, low toxicity, and simple process



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)		(15	5)	(1	6)	(7)
\geq	Phl		Phil	(POC)	Phll / III pi	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	JMT601 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 Phil)	Paclitaxel cationic liposome	JMT103 bone metastasis	Albumin-bound Docetaxel	Irinotecan liposome (US)
SYHX2009 NTRK/ROS1	SYH2038 SOS1	SYH2043 CDK2/4/6	Albumin-bound Sirolimus	Octreotide long- acting injection	Albumin-bound Paclitaxel II	Mitoxantrone hydrochloride liposome (NPC)	Amphotericin B Liposome (US)
SYH2045 PRMT5	SYH2051 ATM	SYH2053 PCSK9	Alprostadil liposome		Clevidipine injectable emulsion	Daunorubicin cytarabine liposome	(03)
SYHA1908		Cisplatin micelle					

Biological Agents Chemical Drugs New formulations

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				H 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				
	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				

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				1 1 1 1 1	
Digestion & metabolism	TG103	GLP-1	Obesity, Diabetes					
	Omalizumab biosimilar	IgE	Allergic asthma, CIU			1 1 1		BL
	Ulsinumab	IL-12/IL-23	Moderate to severe plaque psoriasis					
	Batoclimab	FcRn	MG					
Immune	CM310	IL-4Rα	Asthma, COPD					
ininune	CM326	TSLP	Asthma, COPD					
	NBL-012*	IL-23p19	Psoriasis, HS, IBD					
	Secukinumab	IL-17A	psoriasis					

Pipeline - Chemical Drugs

Over 40 small molecule new drugs under development: 1 filed NDA, 23 under clinical trial stage (2 under Phase III / 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						
metabolis m	SYH2053	PCSK9	FH, Mixed hyperlipidemia					- - - - - - - - - - - - - - - - - - -	
	SYHA1805	FXR Agonist	NASH(MASH)					1 1 1 1 1	
	NBP soft capsule*		VaD, Ischemic stroke(US)						
Nervous system	SYHA1402	ARI	Diabetic neuropathy					, 1 1 1 1	
	Amuxetine hydrochloride enteric tablets	5-HT, SNDRI	Anti-depressant					1 1 1 1	
Immune	SYHX1901	Syk-Jak	RA, SLE, COVID-19						
	Simmitinib	FGFR、KDR	Gastric cancer, cholangiocarcinoma, SQCC					- 1 1 1 1	
	SYHA1813	VEGFR/CSF1R	Relapsed or advanced solid tumour						
Oncology	SYHX2005	FGFR4	Advanced solid tumor					- 	
0,	SYH2043	CDK2/4/6	Breast cancer					1 1 1 1	
	SYH2045	PRMT5	Advanced malignant tumors					1 1 1 1	
	SYH2051	ATM	Solid tumor						

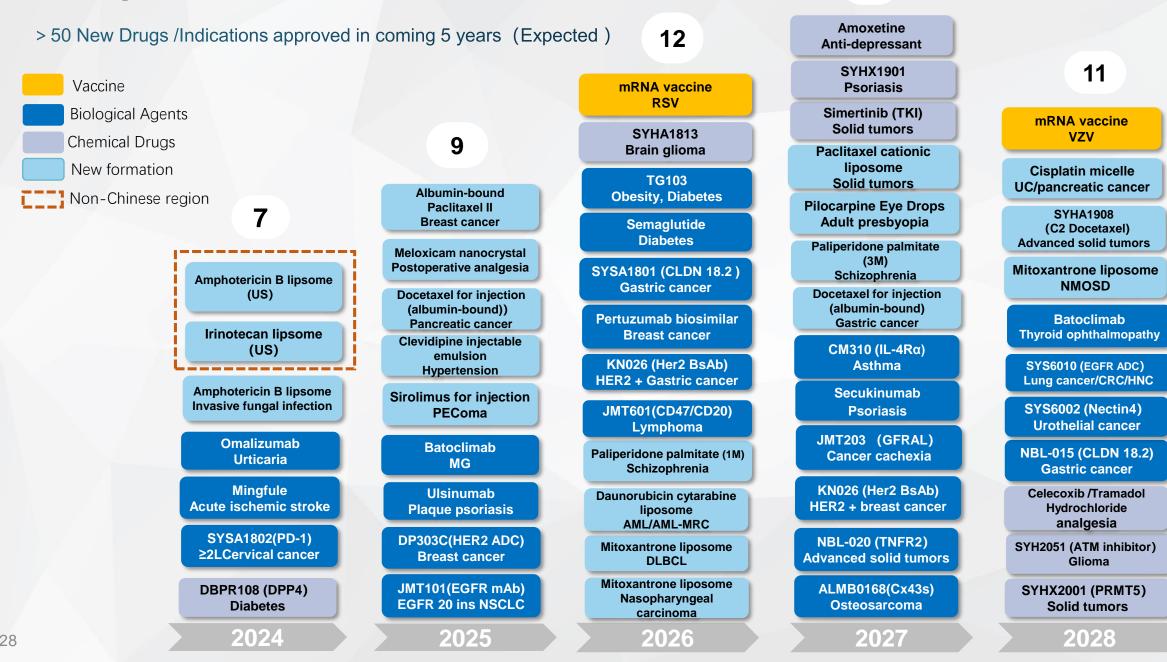
Pipeline - New formulations

Over 30 new preparations under development: 2 applied marketing approval, 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					★ Marketed((P	TCL)
	Irinotecan liposome injection*	pancreatic cancer, Breast cancer					★ Marketed	
	Daunorubicin cytarabine liposome	Leukemia					(pancreatic c	ancer)
	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					NDA a	ccepted
Analgesia	Meloxicam nanocrystal injection	Moderate-to-severe pain						
Anti-infective	Amphotericin B liposome *	Invasive fungal infection						۲
Cardio- cerebrovascular	Clevidipine injectable emulsion	Hypertension					NDA ad	ccepted
	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
1	Dapagliflozin tablets	Digestion & Metabolism	2024
2	Rabeprazole sodium enteric-coated tablets (10mg)	Digestion & Metabolism	2024
3	Olaparib tablets	Oncology	2024
4	Palbociclib tablets (125mg)	Oncology	202
5	Lenalidomide capsules (5mg, 10mg)	Oncology	2024
6	Peramivir injection (150mg/15ml)	Anti-infective	202
7	Aprepitant injection	Others	202
8	Dexrazoxane for injection	Others	202
9	Roxadustat capsules	Others	202
10	Peramivir Injection (300mg/60ml)	Anti-infective	202
11	Regorafenib tablets	Oncology	202
12	Ilaprazole enteric-coated tablets	Digestion & Metabolism	202
13	Terezolamide phosphate tablets	Anti-infective	202
14	Oseltamivir phosphate for oral suspension	Anti-infective	202
15	Mesalazine enteric-coated tablets	Autoimmunity	202
16	Vonorazone fumarate tablets	Digestion & Metabolism	202
17	Adenosine cobalamin capsules	Others	202

IND Approvals Obtained as now

IND approval for the 1st indication (22)	
SYH2045 (Solid tumors)	Meloxicam nanocrystal injection (moderate-to-severe pain for adults)
Clevidipine injectable emulsion (Hypertension)	Octreotide long-acting injection (acromegaly)
NBL-020 (Advanced solid tumors)	SYS6010 (advanced solid tumors)
SYH2051 (Solid tumors)	JMT203 (tumor cachexia)
Semaglutide injection (Type 2 diabetes)	NBL- 028 (Advanced tumors)
SYS6006.32 (Bivalent COVID-19 mRNA vaccine)	Secukinumab injection (Psoriasis)
SYS6011 (Solid tumors)	SYH2038 (Advanced solid tumor)
Dextromethorphan hydrobromide quinidine sulfate tablets (Pseudobulbar paralysis)	SYH2053 injection (Primary hypercholesterolemia or mixed dyslipidemia in adults)
Dexmedetomidine hydrochloride nasal spray (Sedation before invasive examination)	JMT106 injection (Solid tumor) (U.S.)
CPO301 (Advanced solid tumors) (U.S. & Canada)	NBL- 028 injection (U.S.), NBL- 020 injection (U.S.)

IND Approvals Obtained as now

IND approval for additional indications (24)

KN026 injection (in combination with docetaxel albumin for treatment of first-line HER2 positive recurrent and metastatic breast cancer)	Docetaxel for injection (albumin-binding), five indications were approved
JMT101 injection (in combination with docetaxel albumin for treatment of second-line squamous cell non-small cell lung cancer)	Enransubezumab injection (SG-001) (Combined chemotherapy for first-line cervical cancer)
Paclitaxel cationic liposome for injection (Advanced solid tumor - arterial infusion chemotherapy)	Simetinib (Combined with SG001for solid tumor, combined with irinotecan liposomes for second-line and above esophageal cancer)
JMT101 injection (Combined with SG001 and irinotecan for colorectal cancer)	ALMB-0166 (Acute ischemic stroke)
Bartolizumab (Proliferative lupus nephritis)	CM326 (COPD) 、CM310 (COPD)
JMT601 injection (Combined with different regimes for CD20- positive diffuse large B-cell lymphoma)	SYHX1901 tablets (Vitiligo, alopecia areata)
Octreotide Long-acting Injection (Gastrointestinal and Pancreatic Neuroendocrine Tumors)	SYSA1801 injection (Combined with SG001 and CAPOX for the first-line treatment of CLDN18.2-positive, HER2-negative, unresectable locally progressive or metastatic gastric or gastroesophageal junction adenocarcinoma)
Sirolimus for injection (albumin-binding) in combination with endocrine therapy for second-line and above HR-positive HER2- negative advanced breast cancer	Cisplatin micelles (Combined with paclitaxel for advanced solid tumors)



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 2023

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

 Emphasize areas like idiopathic pulmonary fibrosis (IPF), COPD/asthma, and cough, exploring innovative targets, drug-device combinations, and drug delivery systems.
 Focus on high-end antibiotics effective

- Focus on high-end antibiotics effective against clinically resistant bacteria.
- Address conditions such as atopic dermatitis, systemic lupus erythematosus, and inflammatory bowel disease (IBD).

Respiratory, Autoimmune & Anti-Infective Field 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.



· Expand into major population-

based psychiatric disorders

schizophrenia, focusing on

novel targeted drugs with

the layout and collaboration of

improved efficacy, safety, and

Emphasize fast-acting nasal

like depression and

compliance.

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.

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

Nuclear Medicine Field

• Focus on unmet clinical 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



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

35

- 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