

### **2020 Annual Results Presentation**

March 31 2021

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### Content





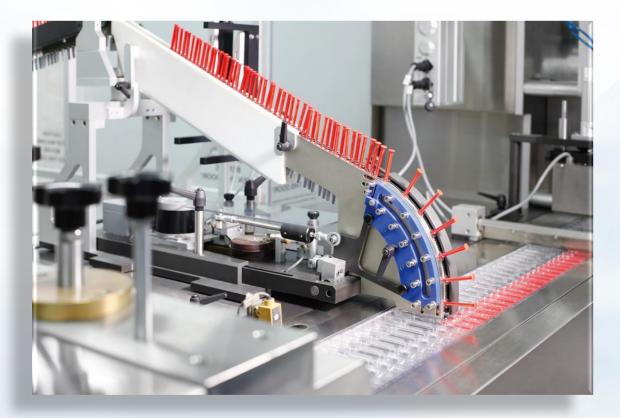
- 2020 Highlights
- R&D & Pipeline
- Financial Performance

• Q & A



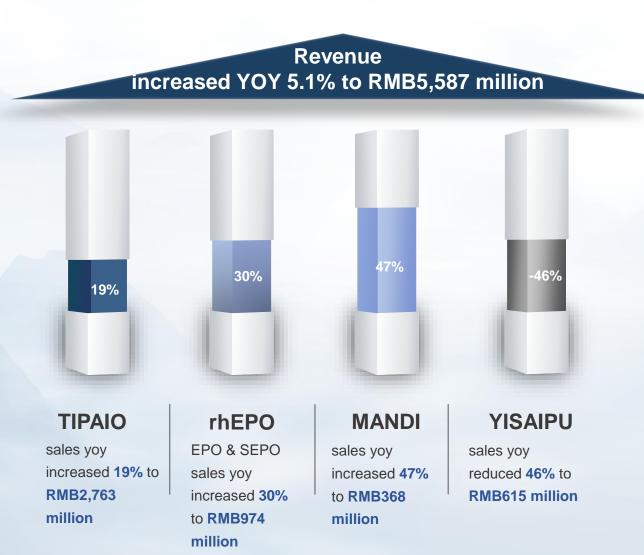
# **2020 Highlights**

Chairman, Director and CEO Dr. Jing LOU

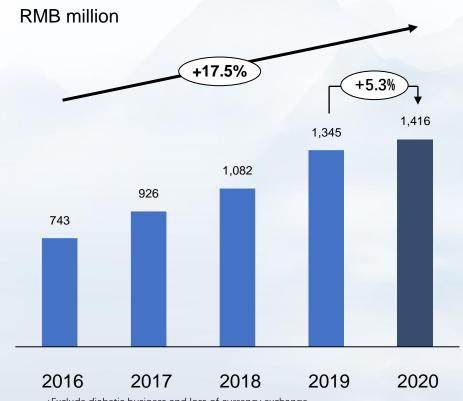


### **Performance Summary**





#### Adjusted Normalized Net Profit Attributable to Owners of the Parent



\*Exclude diabetic business and loss of currency exchange

### **2020 Highlights**



R&D



#### Manufacture

- 302H (HER2) NDA approval and product launched to the market
- 601A (VEGF) AMD and DME completed the patient enrollment in phase I trial, RVO and mpCNV initiated the preparation of Phase II clinical trial
- 602 (EGFR) entered into Phase II clinical trial
- 609A (PD1) completed the patient enrollment in Phase I in U.S. and proceeding patient enrollment in Phase I in China
- 608 (IL-17A) completed patient enrollment in Phase I and preparing for Phase II clinical trial
- 610 (IL-5) IND approved and completed patient enrollment in Phase la
- 611 (IL-4) IND approved by FDA and NMPA, patient enrollment in Phase Ia in U.S.
- SSS06 (rESP) completed patient enrollment in Phase II clinical study
- ~6 small molecules entered into BE

complete



- Construction of the 3<sup>rd</sup> workshop of Shenyang Base was accomplished and entered into equipment testing process
- Phase I of CDMO Project in Shenyang Desheng is progressing smoothly and about to commence the Phase II Construction of the R&D and CDMO Base in Shenzhen
- Songshan Lake entered into the final stage

- Sales Network: 3,263 sales and marketing employees, 797 distributors, and 2,263 third-party promoters
- NRDL: TPIAO sales increased 19%, national negotiation within expectation; Cipterbin was approved, launched, and successfully be included into NRDL in within the same year
- rhEPO sales rebounded to RMB974 million
- Mandi sales yoy rises of 47% to CNY368 million as the online revenue soared ~100%



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Co-developed 5 kinds of multi-specific antibodies and about to import the 1st type of antibody

Selected VSIG-4 and PSGL-1 targeted antibodies partnered MCM in immunooncology and commenced pre-clinical study

Remitch (TRK-820) almost completed the patient enrollment of Phase III bridging trial

Pegsiticase entered into Phase III clinical trial in U.S. and received US\$4 million milestone fee

Invested in GenSight, a world leading gene therapy specialized in ophthalmology company, and its core products have been approved and filed NDA in Europe

Collaboration

GenSight

**TORAY** 

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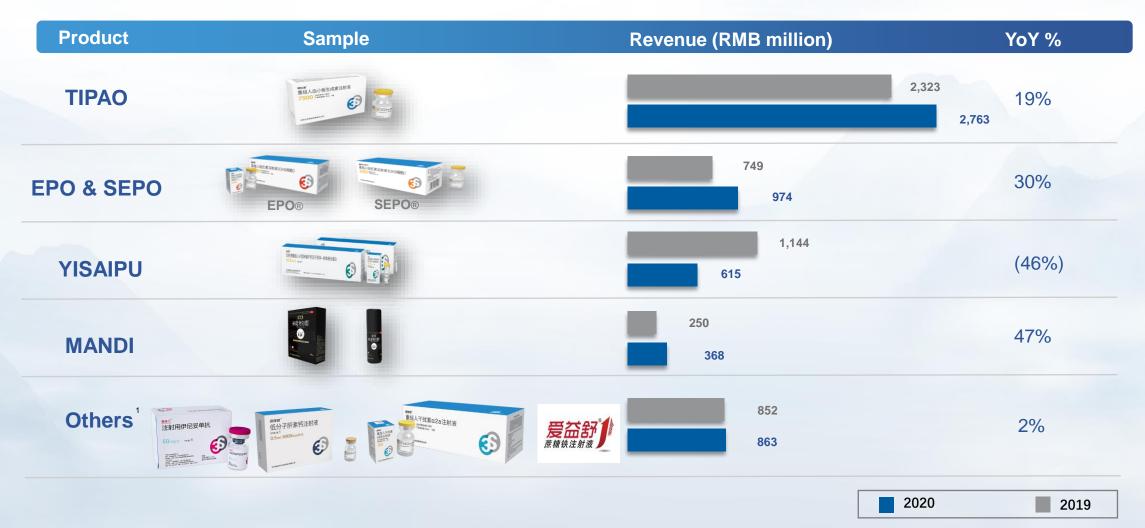
Biosciences

MSCI ESG Rating from BBB to A and surpassed 78% global biotech companies

Sales



### **Core Products Maintained Significant Growth**



1. Except for the above four core products, there are other products including but not limited to Sparin (injectable low-molecular-weight heparin calcium), Aiyishu (Iron sucrose), and Humulin (recombinant insulin) which had exceeded over RMB100 million sales in 2020.

### **TPIAO**

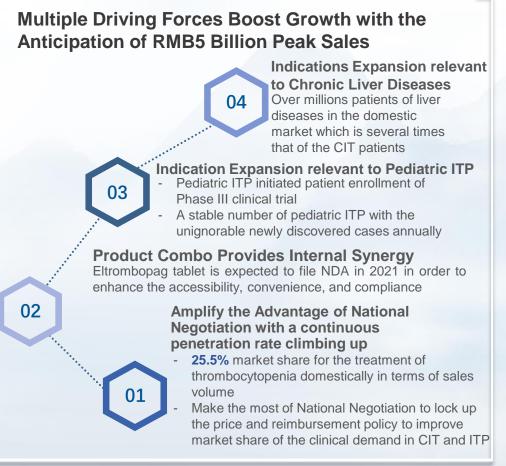
### Performance

- Sound sales growth to RMB2,763 million yoy increased 19% to a new level
- First-to-market and the only commercialized rhTPO product in the world which is recommended by more than 4 Chinese Experts Consensus<sup>1-4</sup> as the firstchoice recommendation for boosting platelet in China
- Sales value with a dominated leading position as the market share of **72.7%**<sup>5</sup>
- National Hospital Ranking Top 11 vs Top 15 in 2019



- . Recommended by Chinese Expert Consensus on Prevention and Treatment of CIT in Malignant Lymphoma
- 2. Recommended by The Expert Consensus for Diagnosis and Treatment of Thrombocytopenia in Adult Critical Illness in China

### Outlook



3. The first-choice recommendation for boosting platelet production as Recommended by *The Expert Consensus for Diagnosis and Treatment of Thrombocytopenia in China* 

4. Recommended drugs for emergency treatment of children with ITP in the *Guidelines for the Diagnosis* and *Treatment of Primary Immune Thrombocytopenia in Children* 

5. Source: IQVIA 2020 and the market definition includes rhTPO, IL-11, and oral treatments

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### rhEPO (EPO & SEPO)

#### Performance

- Sales rebounded yoy 30% to RMB974 million
- Acceleration in the applications of EPIAO in IV and V CKD patients and in CIA oncology indication in Mainland China due to the benefit of NRDL coverage
- A product of National Essential Medicines List (NEML) with a steady incremental penetration rate in the grassroots market domestically
- Dual-brand strategy stabilized the pricing system
- Market share consistently ranked **Top 1**



. Source: IQVIA 2020 and the market definition includes rhEPO and oral treatments.

#### Outlook

Brand Image & Production Capacity Reinforce the Competence While Combination of Innovation and Imitation Extend Products Lifespan

**Expansion of Production Capacity** Newly built facilities in Shenyang and Shenzhen 04 supports the future demands generated from national and provincial VBP Advance NRDL and NEML benefits Continue to solve the NRDL coverage of Long-acting the reduction of allogeneic blood products Iteration transfusion in surgery patients 03 Strengthen the implementation of NEML Deployed two types of long-acting products to lengthen product the life cycle 01 02 **Product Synergy** Intend to file NDA of oral treatment Roxadustat in 2023 - Self-developed SSS17HIF is in Phase I clinical trial

### **YISAIPU**

### Performance

- Dual impact of COVID-19 pandemic and fierce competition, sales slumped 46% to RMB615 million
- New patients increased after Initiated 50% price cut while the DOT was extended within the existing patients. A significantly growth showed in sales volume in Q4 2020
- Currently covered 3,700 hospitals and medical institutes, including Tier 1 hospitals ~1,700
- Market Share in TNFα remained Top 1
- Accumulated treated over few hundred thousands patients



- 1. Source: IQVIA 2020
- 2. Source: Wind, National Bureau of Statistics
- 3. Source: 2018 Rheumatoid Arthritis Treatment Guidelines

### Outlook

Promote County Level Strategy, Accelerate Penetration to Expand Incremental Market

 The number of domestic patients of RA in the grassroots hospitals boosted from 17% in 2010 to 49%<sup>2</sup> in 2019



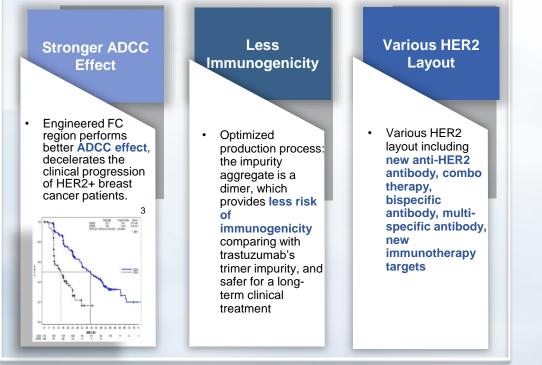
 Domestic DMARDS utilization rate is extremely lower than well-developed countries<sup>3</sup>



### **CIPTERBIN**

#### Performance

- Approved by CDE and was included in NRDL in the same year
- Self-developed, China first innovated under the new label Inetetamab was included in Clinical Guidance<sup>1</sup> and Chinese Experts Consensus<sup>2</sup> after launching to the market



#### Outlook

Seize the opportunity of NRDL, Position in the Core Track, and Estimated Peak Sales reach RMB1.5 billion

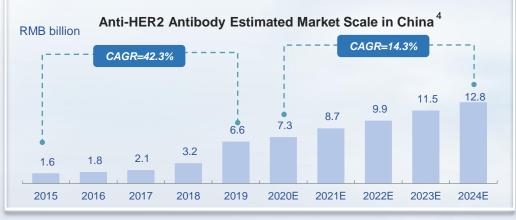
Successfully be included in NRDL with a competitive price to end users

Completed 140+ core hospital entrance



Develop multiple varieties & comb therapy

• Develop innovative anti-HER2 drug resistance treatments, accumulate real-world data from precision treatments, and layout future indications roadmap



- 1. Source: The Principle of Innovative Anti-tumor Drug Clinical Guidance (2020) issued by China National Health Commission
- 2. Source: China Advancing Period of Breast Cancer Experts Consensus Guidance 2020 (CABC3)
- 3. The chart indicates the test result of HER2 positive breast cancer patients after taking the treatment of Cipterbin
- 4. Source: Frost & Sullivan

### MANDI

#### Performance

- Sales surged yoy 47% to RMB368 million
- Market share took up **75%**, occupied the absolute dominate position
- Online sales yoy soared ~100%, Ranked Top 1 in the same category



### Outlook

**Aesthetic** 

Medicine

Consumer

Take a Lead in the Hair Loss Frontier Segment and Aesthetic Medicine Enhances Peak Sale Expectation

#### Market Space

DRGs-free, consumer product hashtag, non-NRDL category with broader market space with at least 200 million potential users domestically

#### Brand Marketing

Fully utilize the first-mover advantage of brand awareness to bond the platform for deeper cooperation

#### **Channel Conversion**

Diversify online platforms and accelerate online customer conversion rate

#### **Dosage alternatives**

- Foam type about to launch in order to enhance convenience

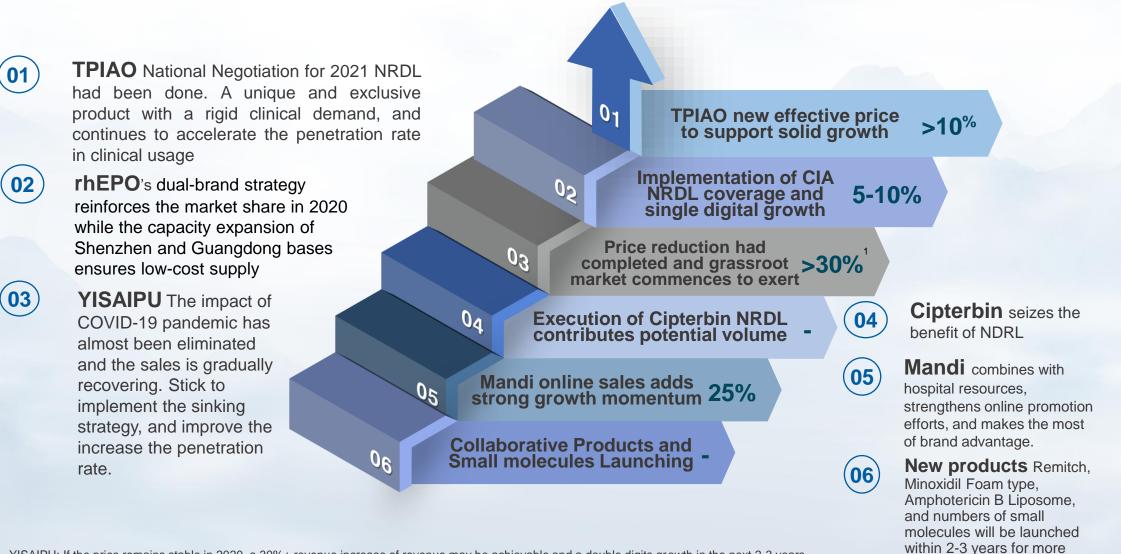
- Expand product line and enrich alternatives

#### Accompany Service

Optimize user experience, extend DOT, and improve compliance and customer relationship

1 Source: CPA 2020

### **Multiple Catalysts Expedite Sustainable Growth in the Recent 2-3 Years**



incremental sales

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## **R&D** and **Pipeline**

Sunshine Guojian President Dr. Zhenping ZHU



### **R&D Progress**

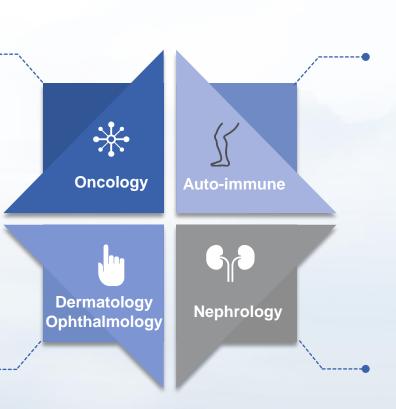
- 302H (HER2) NDA approval and product launched to the market
- 304R (CD20) in the progress of Phase III trial site inspection
- 602 (EGFR) entered into Phase II clinical trial
- 609A (PD1) completed the patient enrollment in Phase I in U.S. and proceeding patient enrollment in Phase I in China
- 612 (HER-2) Filed IND

verseau

705 (Bispecific) Filed Pre-IND

Selected VSIG-4 and PSGL-1 targets partnered MCM in immuno-oncology for the China domestic authorization and commenced pre-clinical study

- MN709 (Minoxidil Foam) completed patient enrollment of Phase III clinical trial
- 601A (VEGF) AMD and DME completed the patient enrollment in phase I trial and initiated RVO and pmCNV preparation of Phase II clinical trial



- 608 (IL-17A) completed patient enrollment in Phase I
- 610 (IL-5) IND approved and completed patient enrollment in Phase Ia clinical trial
- 611 (IL-4) IND approved by FDA and NMPA, and proceeding patient enrollment in Phase la clinical trial in U.S.
- 613 (IL-1β) Filed Pre-IND
- Equity Investment:

Sensorion

GenSight

Core product GS010 have been approved and filed NDA in Europe

Core product SENS401 is proceeding Phase II clinical trial of SSNHL1 and preclinical studies multiple preventive indications

- SSS17 (HIF-117) initiated patient enrollment of Phase I clinical trial
- SSS06 (rESP) completed patient enrollment in Phase II clinical trial
- RD001 (rESP) proceeding patient enrollment in Phase II clinical trial
- **TPIAO (rhTPO)** completed surgery patients with hepatic dysfunction at the risk of thrombocytopenia Phase I trial and recruiting patients for pediatric ITP Phase III clinical trial Remitch (TRK-820) almost

milestone fee



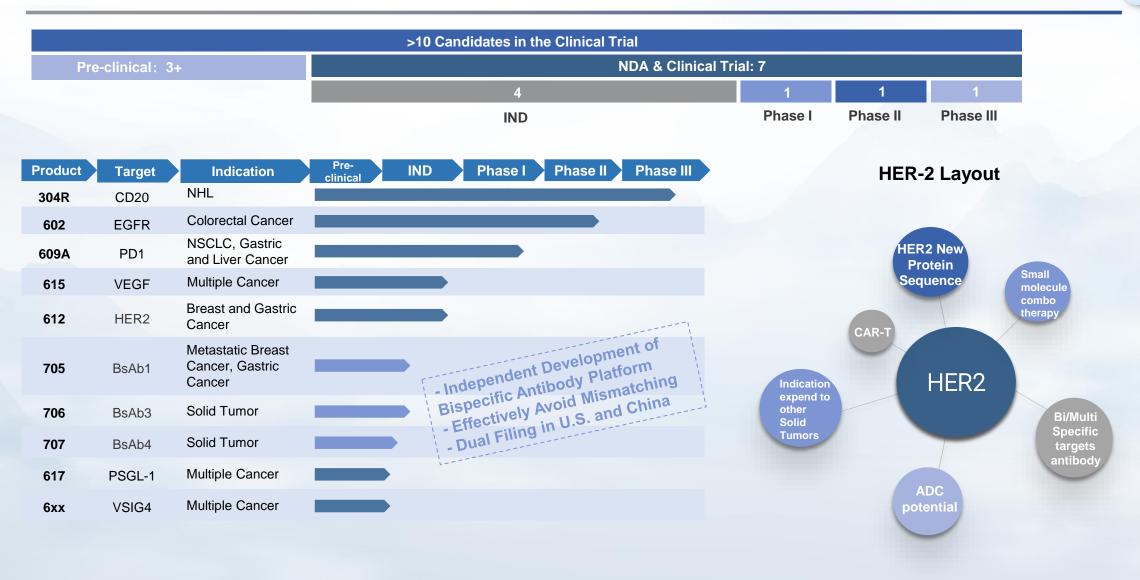
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completed the patient enrollment of Phase III bridging trial Pegisiticase entered into Phase III clinical trial in U.S. **Biosciences** and received US\$4 million

### **Robust and Innovative Product Pipeline**

	Pre-clinical	IND	Phase I		Phase II	F	Phase III & BE	NDA
	6	4	7		5		10	2
Therapeutic	Product Candidate	Pre-clinical	Clinical Trial Application	Phase I	Phase II	Phase III	New Drug Registration	Product Approv
	SSS06 Long-acting EPO							
	RD001 Long-acting EPO							
Nenhaeleeu	SSS17 HIF inhibitor							
Nephrology	SSS12				BE			
	SSS13				BE			
	TRK-820/Remitch							
	304R anti-CD20 antibody							
	602 anti-EGFR antibody							
	609A anti-PD1 antibody			Uni	ted States			
	612 anti-HER2 antibody							
	615(SB8) anti-VEGF antibody							
Oncology	705 BsAb1							
	706 BsAb2							
	707 BsAb3							
	617 anti-PSGL-1 antibody							
	6xx anti-VSIG-4 antibody							
	SSS24				BE			
	301S TNFR-Fc Fusion Protein							
	TPO-105 Pediatric ITP							
	TPO-106 CLD							
	SSS07 anti-TNFα antibody							
	SSS11 Pegsiticase							
	608 anti-IL17 antibody							
Auto-Immune	610 antio-IL5 antibody							
and others	611 anti-IL4R antibody			United Sta	ates			
	613 anti-IL1β antibody							
	601A anti-VEGF antibody							
	AP506				BE			
	SSS20				BE			
	SSS32				BE			
Dermatology,	TK805							
Ophthalmology	MN709							

### **Biologics Pipeline in Oncology**



### **Biologics Pipeline in Nephrology, Auto-immune, and Others**

herapeutic	Product	Target	Indication	Pre- Clinical	IND	Phase I	Phase II	Phase III	NDA	FOCU	IS ON GLOBAL BLOCKBUST	
	301S	$TNF_{\alpha}$	RA, PS, AS									
Auto-	SSS07	TNFa	RA, AS, Ulcerative colitis							608	<ul> <li>New amino acid sequence targetir</li> <li>17A</li> <li>Global sales of Secukinumab with</li> </ul>	
immune	610	IL5 $\alpha$	Asthma, COPD								same target is approximately US\$	
	613	IL1b	Multiple auto- immune diseases								billion in 2020	
	SB16	RANKL	Bone diseases <sup>2</sup>								- New amino acid sequence targetir	
	608	IL17A	PS, AS, Psoriatic arthritis							611	<ul> <li>IL-4Rα</li> <li>Dual filing in both U.S. and China</li> <li>Clabal calca of Duplizumab with the</li> </ul>	
Dermat ology	611	IL4Rα	Atopic dermatitis, Asthma								<ul> <li>Global sales of Duplizumab with the same target exceeded US\$4 billion 2020</li> </ul>	
	ТРО	-	Pediatric ITP								2020	
Nephro	ТРО	-	Surgery patients with hepatic dysfunction at the risk of thrombocytopenia	_	_	_				610	<ul> <li>New variable region sequence antibo targeting IL-5</li> <li>The 1<sup>st</sup> domestic application and no similar launched product in China</li> </ul>	
logy	SSS06	EpoR	Anemia associated with CKD							010	<ul> <li>Global sales of Mepolizumab with the same target exceeded US\$1.2 billio</li> </ul>	
	RD01	EpoR	Anemia associated with CKD				•				2020	
	SSS11	Uric Acid	Hyperuricemia, Refractory gout								- Global multi-center clinical trials	
Ophthal mology	601A	VEGF	Ophthalmologic diseases (AMD,DME,				•			601A		
			RVO, pmCNV)									

1. Source: Global sales of each drug are consolidated from public data

2. Postmenopausal female with osteoporosis at high fracture risk

18<sup>2.</sup>

### **Small Molecule Combo**



#### Sevelamer Carbonate (SSS13)

- · Oral treatment of Hyperphosphatemia
- NDA filed in 2020

#### Cinacalcet Hydrochloride (SSS12)

- Oral treatment of Hyperthyroidism
- NDA: 2021

#### Roxadustat (SSS38)

- Oral treatment of rhEPO which included into the Top Notch Guidance as recommendation, creates synergy with **EPO**
- NDA: 2023



#### Apremilast (AP506)

- Oral treatment of PS and Psoriatic Arthritis
- Synergy: YISAIPU
- NDA: 2021

#### Tofacitinib (SSS32)

- Oral treatment of RA
- Synergy: YISAIPU
- NDA: 2021

#### Eltrombopag (SSS20)

- Oral treatment of ITP creates
   synergy with **TPIAO**
- NDA: 2021



Trifluridine and Tipiracil Hydrochloride Tablets (SSS24)

- Oral treatment of Colorectal Cancer
- Synergy: SB8, 602
- NDA: 2022

#### Dermatology



#### Minoxidil Foam (MN709)

- Foam Type
- Synergy: MANDI
- NDA: 2021

#### Apremilast (AP506)

- Indication: PS
- Synergy: YISAIPU

### **Integrated International Strategic Collaborations**



Cooperated with Sensorion, the world's leading new therapy company for inner ear diseases, and obtained the pre-emption right to buy its products in Greater China

Sensorion

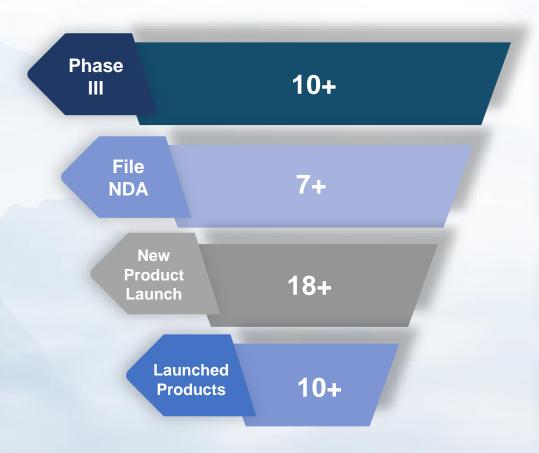
1.

Great China Region includes China mainland, Hong Kong, Macau, and Taiwan

### **R&D Outlook**



### Estimation in 2025



	601A (RVO/pmCNV)	611	612	613 7	05,706,707	617	VSIG-4	SSS17
Estimated Peak Sales (RMB bn)	16 <sup>1</sup>	30	10	30	30	10-	20 -	15 <sup>2</sup>
	01A DME) 6	<b>02</b> (	609A China)	610	SB16	RD001	SSS06	Other possible candidates
1	6 <sup>1</sup>	5	15 <sup>3</sup>	15	-	20 <sup>2</sup>	20 <sup>2</sup>	
TPIAC	0 <sup>4</sup> Remitc	h 301S	304R	615	5 608	609A (U.S.)	601A (AMD)	>10 small molecule biosimilar
-	-	-	10-13	6	15	15 <sup>3</sup>	16 <sup>1</sup>	
TPIAO	EPO	Mandi	Yisaipu	Cipte	rbin Xeno	opax A	mpholipad	Existed launched products

- 1. Total peak sales for all indications covered by 601A
- 2. Total peak sales for rhEPO products
- 3. Total peak sales for all indications covered by 609A
- 4. Indications of Pediatric ITP and relevant liver diseases for TPIAO

### **Integrated R&D Centers and Platforms**



### **Comprehensive Manufacturing Capabilities Adhere to Int'l Quality Standards**

- All 10 production lines for different dosage forms are certified by GMP in 2010
- QA personnel represent 20%+ of all manufacturing employees at the site
- General manager has 10+ years' experience of pharmaceutical R&D, manufacturing and quality control
- Suzhou High-Tech Park Future CMO Facility
- Serves world-renowned companies such as Mylan and Sanofi
- QA personnel represent nearly 40% of all manufacturing employees at the site
- EU GMP certified production lines in Italy
- All existing and new production lines were granted GMP certification in 2013 and in 2016
- QA personnel represent 20%+ of all manufacturing employees at the site
- QA director has 10 years' experience of pharmaceutical R&D, manufacturing and quality control
- Shenzhen Song Shan Lake Facility(EPO, R&D, CDMO)



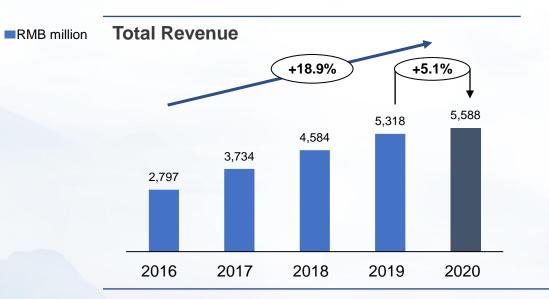
## Financial Performance

Chief Financial Officer Mr. Fei WANG

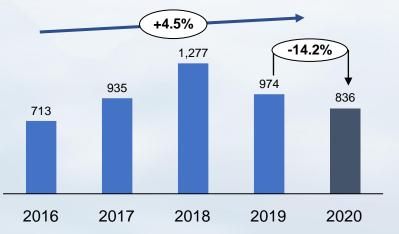


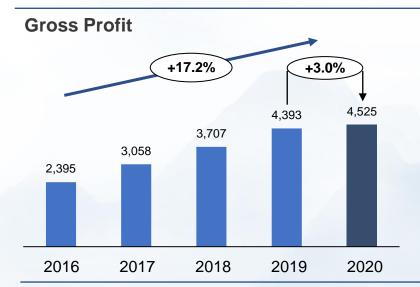


# Main Revenue Continuously Maintains Steady Growth while Gross Profit retains 80%+

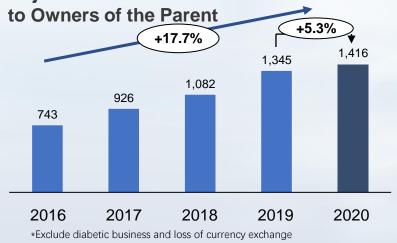


#### Net Profit Attributable to Owners of the Parent

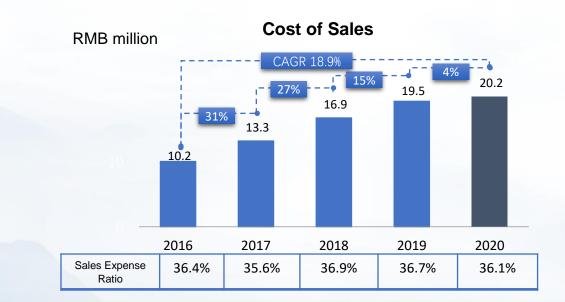


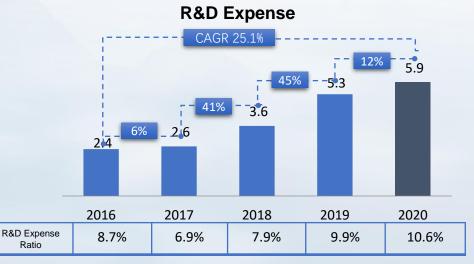


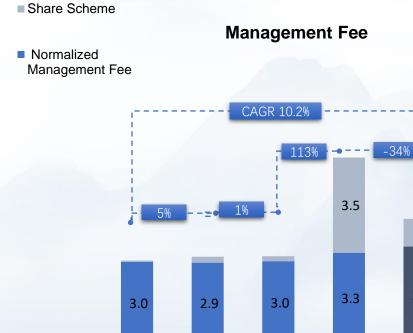
**Adjusted Normalized Net Profit Attributable** 



### Continuously Invest into R&D while Sales and Management Fees are Gradually Reduced







2017

8.4%

10.8%

2018

2019

6.9%

2016

Management

Cost Ratio

1.0

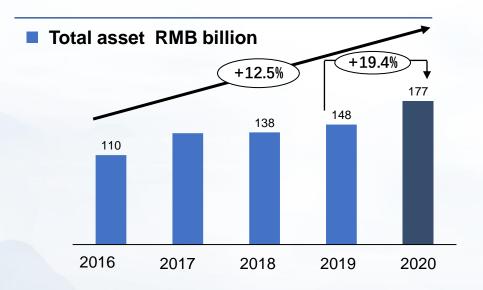
3.5

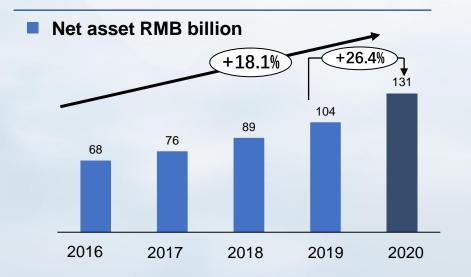
2020

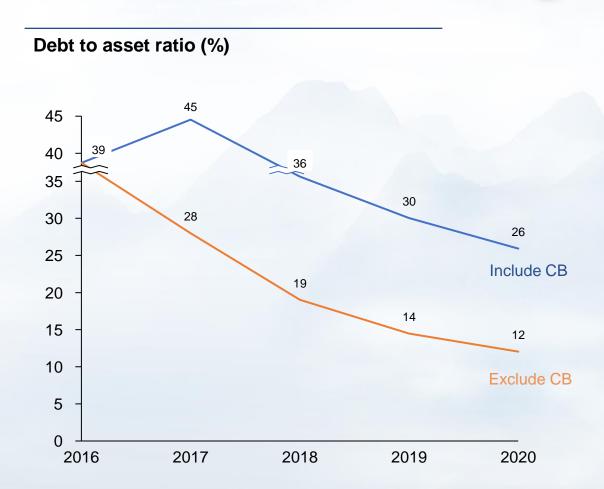
8.1%

12.7%

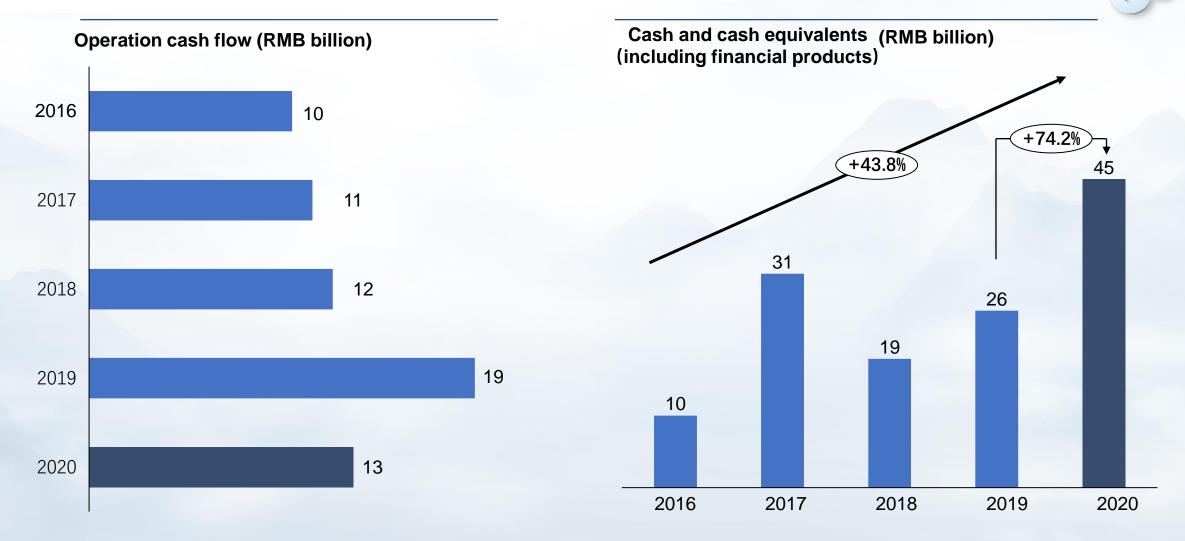
# Significant growth of both net assets and total assets in 2020 while the debt to asset ratio continuously decreases







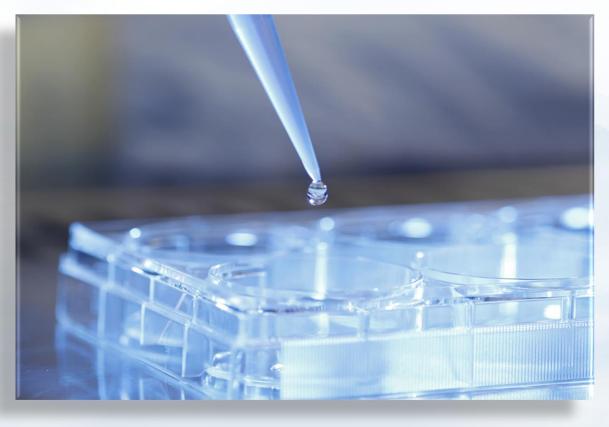
### **Excellent Cash Flow and Abundant Cash Reserves**





# Q & A

### Management Team





## THANKS

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