



Management Presentation

2020 Interim Results

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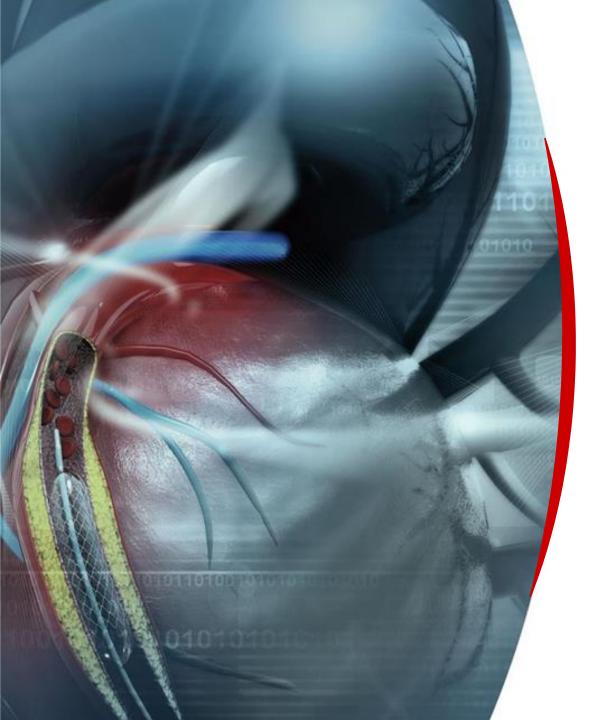
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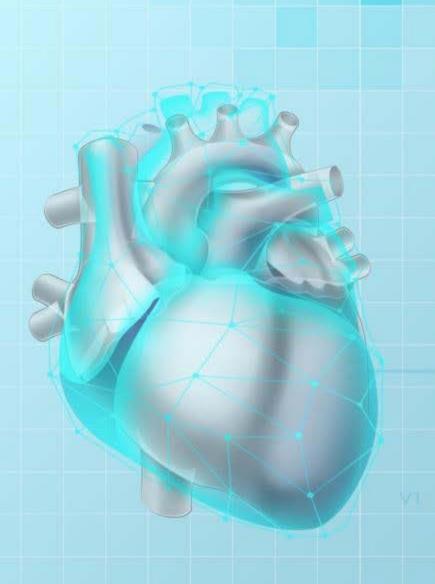






01

Summary





Leading Innovative Player Delivering Interventional Procedural Medical Device Solutions

Our Focus



Transcatheter Valve Therapeutic
Medical Devices



Neurointerventional Procedural Medical Devices

2020 Interim Highlights



01

TaurusOne®: Getting Ready for NMPA Registration

02

TaurusElite®: Highly Acclaimed by KOLs during Clinical Trial

03

Decent Revenue Growth of Neuro Business amid COVID Headwind

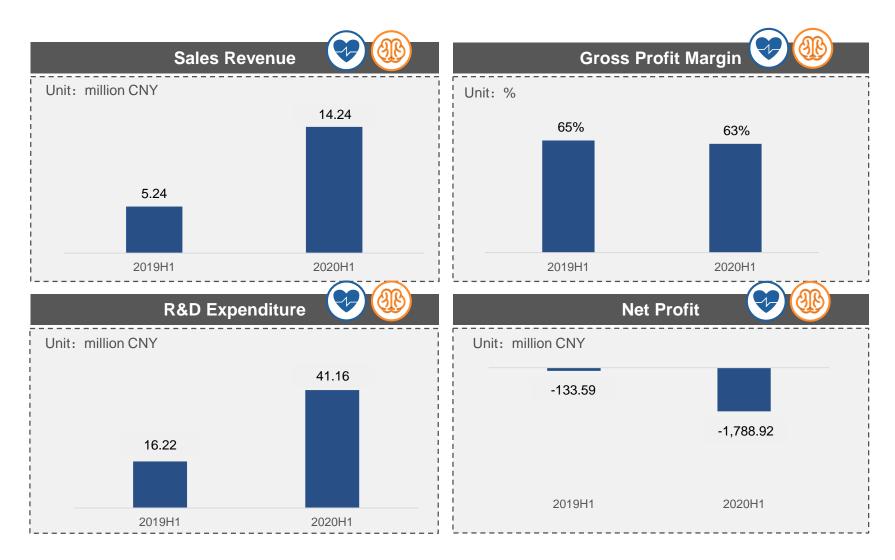
04

Launching First Ischemia Stroke Neurointerventional Product

05

Systematic Preparation for TAVR Commercialization

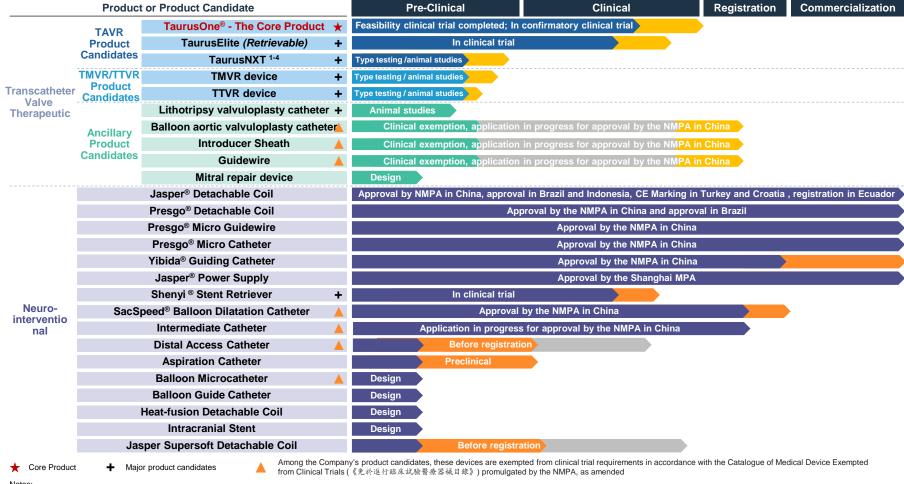
Interim Financial Overview



Note: Consolidated numbers. The operating results of the Neuro-interventional business were only consolidated as of 29 March 2019

Comprehensive Portfolio of Products and Product Candidates in Various Stages of Development

Seven registered products and 19 products under development (as of 2020.08.28), a variety of advanced features are tailored to the needs of Chinese doctors and patients

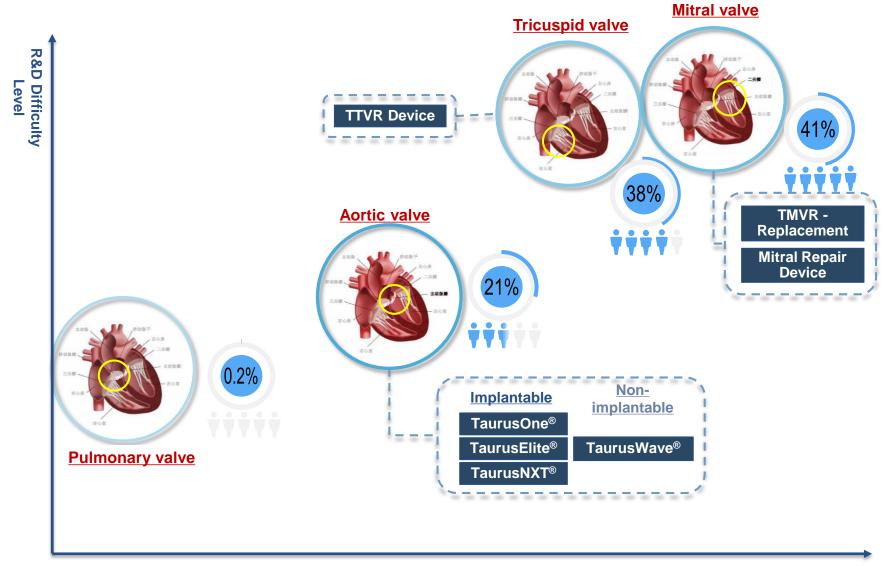


Notes:

- 1. Retrievable, steerable and glutaraldehyde-free anti-calcification
- 2. The "retrievable" function allows physicians to retrieve the valve during a TAVR procedure if the initial release position of the valve is not ideal
- 3. The "steerable" function allows physicians to steer the position and orientation of the valve during a TAVR procedure
- 4. The "glutaraldehyde-free anti-calcification" technology can effectively resist valve calcification, and significantly improve the durability of the valve



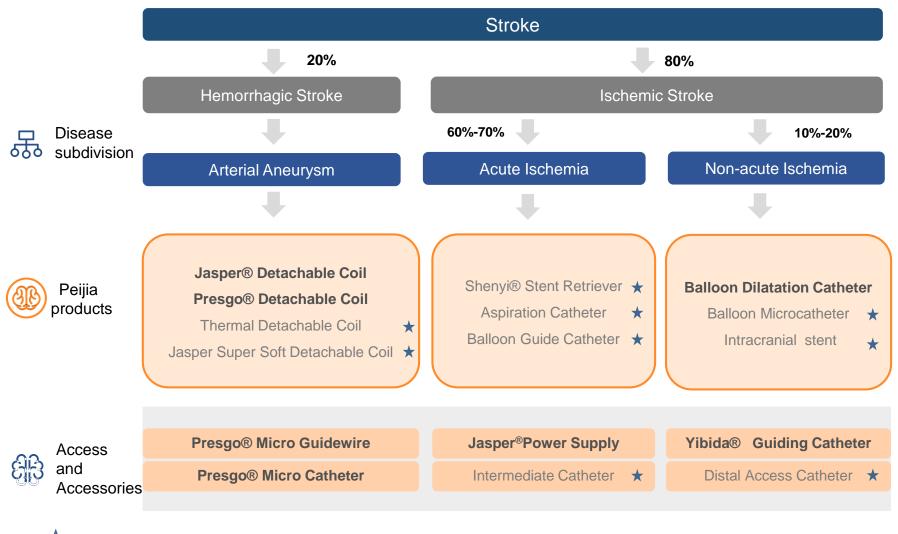
Strategic Product and Pipeline Coverage in Heart Valve Diseases



Note: Valve diseases above mainly include stenosis and regurgitation, data from Frost & Sullivan

Number of Patients

Strategic Product and Pipeline Coverage in Stroke



Note: The star label products are in the design/clinical/registration stage and have not been licensed for registration. Please refer to the product schedule for details.



Recent Progress of Major Products



TaurusOne® : Getting ready to submit application to NMPA

- Concluded the follow-up on all patients in April 2020
- Have compiled statistical reports
- Obtaining stampings / sign-offs from all major and sub trial centers
- NMPA application will be ready once all the above sign-offs are received



TaurusElite®: Received positive feedbacks from KOLs during clinical trial

- · Conducting clinical trial in 10 centers
- Partial data were presented at China Valve Hangzhou conference by Han Yaling's team
- Initial performance received positive feedback from KOLs
- TaurusElite® entered publicity period for "innovative medical device" green path







- Design Features:
 - ① Distal clot capturing basket
 - 2 Longer net
 - 3 Bigger mesh, no kink at the turn
 - 4 Entirely visible
- 11 trial centers
- Positive feedback from KOLs

- Design Features:
 - Rapid exchange structure
 - 2 Longer delivery system
 - 3 Laser welding distal end
 - 4 Full size specification
 - ⑤ Small profile
- Obtained NMPA registration certificate in August 2020



Shenyi ® Stent Retriever: currently in clinical trial



SacSpeed® Balloon Dilatation
Catheter: Received NMPA approval



Commercialization



- Sales recruitment: Plan to hire 6-7 regional managers and 20-30 front-line sales by 2023
- Professional marketing and medical team:
 The Marketing Department and the Sales
 Department cooperate to carry out training program and marketing education. Building word of mouth among KOLs for TaurusOne® and TaurusElite® ahead of time
- Plan to build a flat sales structure to cover the top center efficiently
- Set up Department of Medicine and CoreLab



- 2020 marks the first year in which Peijia launches their first product for ischemia
- SacSpeed® balloon dilatation catheter was approved by NMPA in August 2020
- Shenyi ® Stent Retriever as well as the aspiration catheter are in the pipeline



Training Plan



- Develop standard curriculum for marketing and medical teams
- Establish and improve the training and certification for clinical support specialist (CS) and the mid and long-term development plan
- Establish the doctor education program "Yijia College for Valve"(《医 嘉学院携辦行》)

Expansion of business in hemorrhagic stroke



- Peijia is the only domestic player who can offer all the necessary accessories to complete an aneurysm embolization operation
- Revenue from commercialized products have entered an inflection point
- The new product was gradually admitted to hospital
- Despite the negative impact of COVID-19, sales in the first half of 2020 still achieved significant growth compared with the same period of the previous year







02

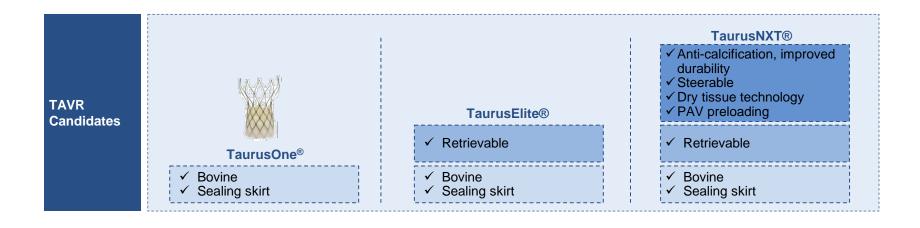
Business Overview





Heart Valve Business

TAVR: Product Pipeline



	1st Gen TAVR: TaurusOne [®] Preparing for Application	2nd Gen TAVR: TaurusElite® In Clinical Trial	3rd Gen TAVR: TaurusNXT® Preclinical Preparation
2020H1 Schedule	 All patients were followed up for 30 days, 6 months and 12 months by April 2020 Complete the clinical data collation & analysis, signing in the main research Units and sub-centers Expected to submit for registration Soon 	 Started clinical trial in December, 2019 Partial clinical data were released at China Valve Hangzhou in July, 2020 Received positive feedback from KOLs In August 2020, TaurusElite® entered publicity period for "innovative medical device" green path 	In third party testing / animal studies Plan to start clinical trials in 2021H1

Notes

^{1.} The only one in the TAVR product market in China that has conducted both feasibility and confirmatory clinical trials for the first generation TAVR product with clinical trial protocols approved by the NMPA (IND)



TaurusOne® Progress

Feb 2017

- · TaurusOne® is recognized by NMPA as "Innovative Medical Device"
- NMPA approved single-center feasibility clinical trials and multicenter confirmatory clinical trials

Sept 2017

Initiate multicenter confirmatory clinical trials to confirm safety and efficacy

Apr 2019

· The final patient in the confirmatory clinical trial was enrolled













NMPA preclinical review

Single center feasibility clinical trial

Multicenter confirmatory clinical trial

Registration

NMPA Approval

Aug 2017

- · Complete a single center feasibility clinical trial to assess safety
- · All-cause mortality was zero and cardiac function significantly improved

Apr 2020

· For confirmatory clinical trial, we completed 20 days, 6 months and 12 months of follow-up for all patients

Expected



1st Gen TaurusOne® clinical data provide evidence of safety & efficacy



 The only TAVR player that conducted NMPA pre-approved feasibility and confirmatory clinical trials

(Sept 2014 《Principles for Clinical Trial Review for Transcatheter Aortic Valve Implantation (Draft) //



More stringent in enrolling patients in our clinical trials

	Male N=58	Female N=62	Total N=120*
Age	76.95	78.16	77.58
STS Score	9.81	10.06	9.94
Agatston Score	560.9	324.0	438.5

6 hospitals

(Beijing Fuwai Hospital is the main research institution)

97.6%

Operative successful rate 6.67%*

All-cause mortality within 12 months (Primary safety endpoint)

Note: Exclude 5 roll-in patients



TaurusOne® Academic conference publication and journal submission

Released at academic conferences



In PCR e-Course at 24/06/2020 Hot line/ Late-Breaking Trials, TaurusOne® TAVR system for the treatment of patients with severe calcific aortic valve stenosis was first published by Prof Wu in a multicenter, prospective, single-group, target-controlled clinical trial for 12 months

Contribute to top academic journals



Plan to submit to EuroIntervention



Taurus Elite® Progress Overview

Dec 2019

• TaurusElite® begins clinical trials with the first patient enrolled

July 2020

· TaurusElite® Clinical trial of progressed smoothly, some data is released at HZ Valve Conference

Aug 2020

• TaurusElite® is recognized by NMPA as "Innovative Medical Device"

2020 Q3

· Complete the TaurusElite® clinical trial and patients enrollment























Confirmatory Clinical Trial

Registration

NMPA

TaurusElite®: Preliminary data from clinical trials

Data presented at China Valve Hangzhou 2020 on July 24th*

Primary Endpoint

The incidence of complex events within 30 days

Baseline

	N=46
Age,Year	75.72±3.72
STS, %	9.36±4.22
HU850, mm ³	588.38±403.10

30Day Outcome

	TaurusElite®™ Experiment(N=31)	
	Postoperation	30 Days
All-Cause Mortality	3.23%(1)	3.23%(1)
Disabling stroke	0.00%(0)	0.00%(0)
MI	0.00%(0)	0.00%(0)
New PPI	9.68%(3)	9.68%(3)

Approval

Expected

TaurusElite® of Academic conference & Live surgery broadcast



Some data were published at 2020 HZ Valve Society



Live surgery in Second Affiliated Hospital of ZJU



CIT2020, Live surgery in General Hospital of Northern Theater



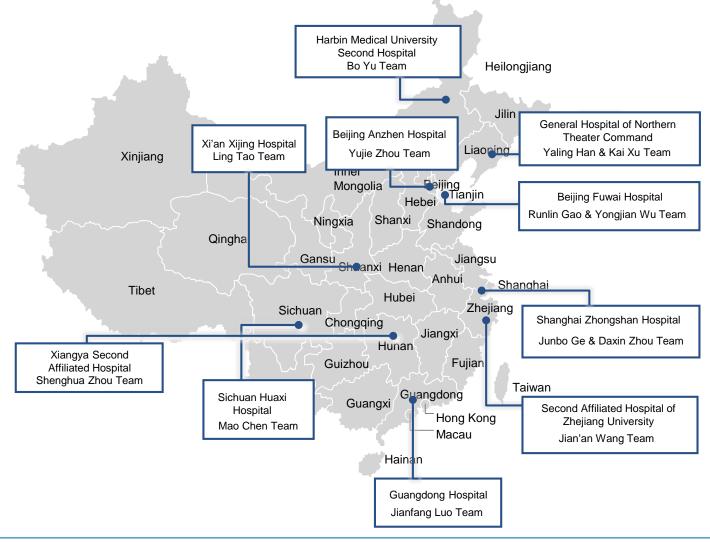
Live surgery in Guangdong Provincial People's Hospital

Notes: *The enrollment is still ongoing. The data listed above is only for reference. It is not the final statistic data



Taurus Elite® Clinical Trial: Received Positive Feedbacks from KOLs

TaurusElite® is conducting clinical trial in 10 centers



Other Heart Valve Products Development Progress and Pipelines

Major Product Candidates



Minimally invasive procedure for mitral regurgitation



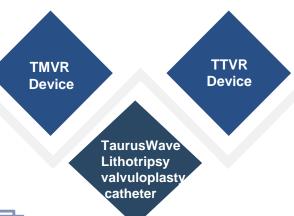
Separate the anchorage and sealing skirt designs from the valve function to address mitral valve's characteristic of complex structure



One of the few companies with TMVR product candidates under development in China¹



In type testing and animal studies



+

Soften calcification on leaflets to alleviate stenosis



In animal studies, expected to enter into the clinical trial stage in the fourth quarter of 2020 or first quarter of 2021



Potentially non-implantation solution for aortic stenosis



Minimally invasive procedure for tricuspid regurgitation



Separate the anchorage and sealing skirt designs from the valve function to address tricuspid valve's characteristic of complex structure



One of the few companies with TTVR product candidates under development in China²



In type testing and animal studies

Other Transcatheter Valve Therapeutic Products

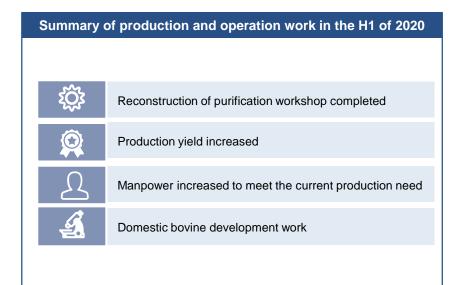
Product category	Features and applications	Development Status	
Balloon aortic valvuloplasty catheter+	Used for the dilatation of heart valves or vena cava stenosis		
Introducer sheath+	Used to intrude into the artery percutaneously in an interventional surgery and establish a passageway for introducing a catheter into the blood vessel	Registration	
Guidewire+	Used to establish a passageway from the puncture position to the lesion or to the distal end through the lesion to assist other devices in positioning		
Mitral repair device	Used to repair the mitral valve when treating mitral regurgitation(Transatrial septum)	Design	

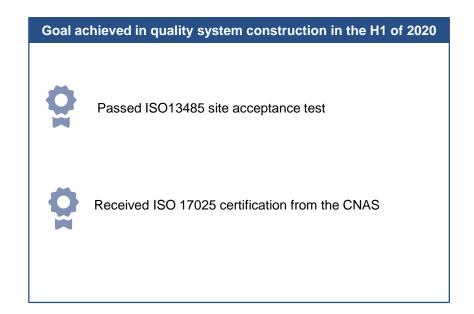
Notes 1&2: Source from Frost & Sullivan



^{+:} Exempted from clinical trial requirements

Heart Valve Business: Production Plan & Progress





Expansion in Production Capacity



Current capacity (TAVR): 3,000 sets /year

The standard labor-hour required for fixation is reduced from 13hrs to 10hrs

Heart Valve Business: Marketing and Academic Events





- In PCR2020, Professor Yongjian Wu announced the clinical trial data of TaurusOne® globally on June 25
- . Company hold the Peijia Special Session of "Demonstration, TAVR Way" in NCF2020 on June 10
- TaurusOne® clinical trials announced in China on July 4
- Summit forum of "Heart road with valve " was held in China Valve (Hangzhou) 2020 on July 27





TaurusElite®™ is recyclable, excellent and reliable



live broadcasts, all using the TaurusElite®™ recyclable TAVR system.

4

Platforms. Different audiences, different ways.Live broadcast of the whole network for the public, and live broadcast of professional meetings for doctors



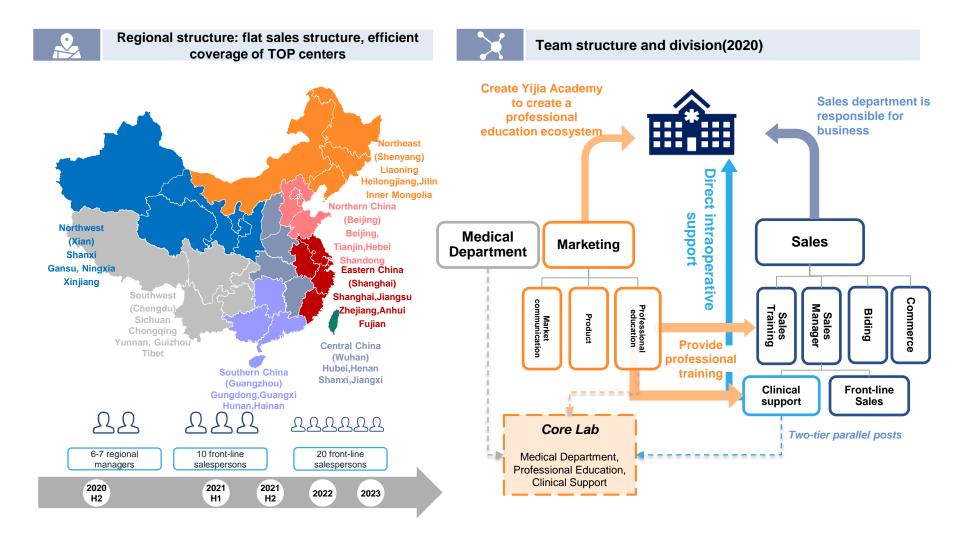
centers (Fuwai, Huaxi, SAHZU, General Hospital of Northern Theater Command(GHNTC), Second Xiangya Hospital of Central South University, Anzhen), share their experience to let more doctors and experts understand Peijia Valve



- The team of chairman of SAHZU-Wang Jianan held operations live for the public on 2 June
- The team of chairman of SAHZU-Wang Jianan held operations live on 18 June as one of series of activities of China Structural Heart Disease Academic Activity Week.
- During CIT2020, the team of Professor Xu Kai from the General Hospital of the Northern Theater Command held operation live on 3 July
- At the China Valve (Hangzhou) 2020, six centers (Fuwai, Huaxi, SAHZU, GHNTC, Second Xiangya Hospital of Central South University, , Anzhen) held nationwide combined operation live in 27-30 July



Preparation for TAVR Commercialization: Marketing Team Structure



Preparation for TAVR Commercialization: Establish a Professional Marketing and Medical Team

Clinical Specialist (CS) Training System and Long-term Development Plan

	Level	Level 0/1	Level 2	Level 3
1	Fraining period	1Week~1 month	3~6 months	6~18 months
	mber of surgical assisting cases	0~10 cases/>10 cases	>30 cases	>80 cases
C	Clinical role and positioning	Taurus operation follower	TAVR clinical technical specia	alist TAVR clinical expert
C	Anatomical cognition	Understand	Skilled	Proficient
linical	Image recognition	Understand	Analysis	Proficient
techni	Product information	Product performance parameters	Competitive product differentiation comparison	R&D design ideas
cal spe	Operation Knowledge	Master the procedure of Taurus surgery	Different valve implantation procedures	Patient and valve selection recommendations
cialist t	Skill & application	Standard process skills application	Techniques analysis of complex lesion	Technical support for special circumstances
Clinical technical specialist training module	Strategic Analysis	Understand the main points of preoperative discussion	Master the use of intraoperative strategies	Provide intraoperative coping strategies
modu	Perioperative management	Anesthesia/Medication/Imaging Team	Familiar with TAVR team building	Provide new center TAVR program
е	Evidence-based frontier	Understand the evidence-based progress of TAVR	Follow up on TAVR cutting-edge hotspots	Participate in the formulation of prospective clinical programs



Preparation for TAVR Commercialization: To Create a Professional Education Ecosystem

Customized projects for the needs of hierarchical customers

Leading hospitals:

8 centers completed 200+ TAVR cases

Tier1 Leading hospitals TAVR industry KOL Mature TAVR teacher

spokesman

Tier 2 Key hospitals Carry out TAVR independently > 50 cases/year

Operator reeducation

Tier 3 TOP hospitals TAVR <50 cases/year Need teaching

Operator Training

Establish the doctor education system (《医嘉学院携辦行》)



- Online courses
- Ultrasound + CT + key steps of surgery + case intensive reading



- Combining leading center teachers and TAVR training system, integrating Peijia characteristics, focusing on CT evaluation and case intensive reading and review
- Planning to cooperate with Cardiovascular Alliance to academically support standardized training of Valve Center



Suzhou headquarters is under planning







Neurointerventional Business

Neurointerventional Business: Market Launch of the First Product Targeting Ischemic Stroke

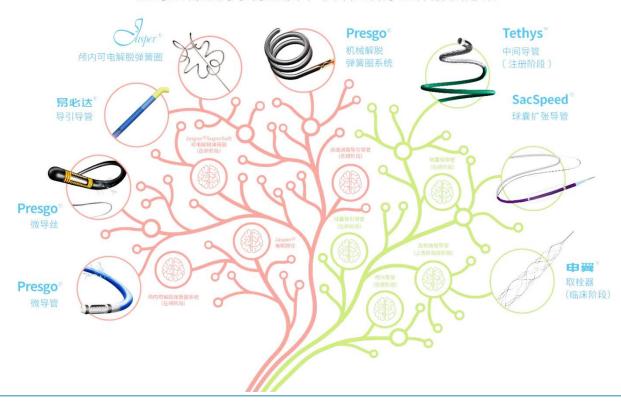


- ✓ Achieva is committed to provide a comprehensive solution for stroke treatment
- √ We have constructed product lines of hemorrhagic products, ischemic products and access products



迭代,蝶变。新声

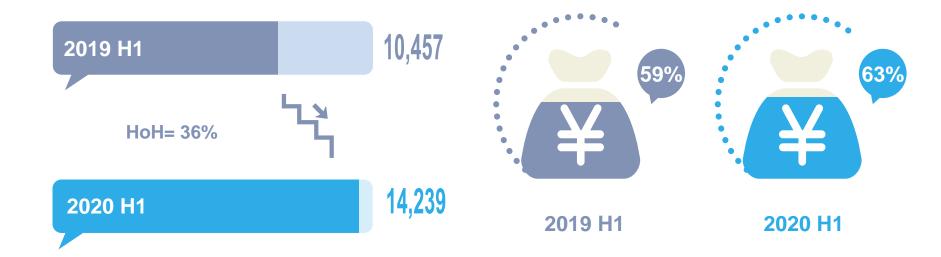
加奇生物致力于打造脑卒中介入治疗整体解决方案



Neurointerventional Business: 2020 Interim Highlights

Revenue (in thousand CNY)

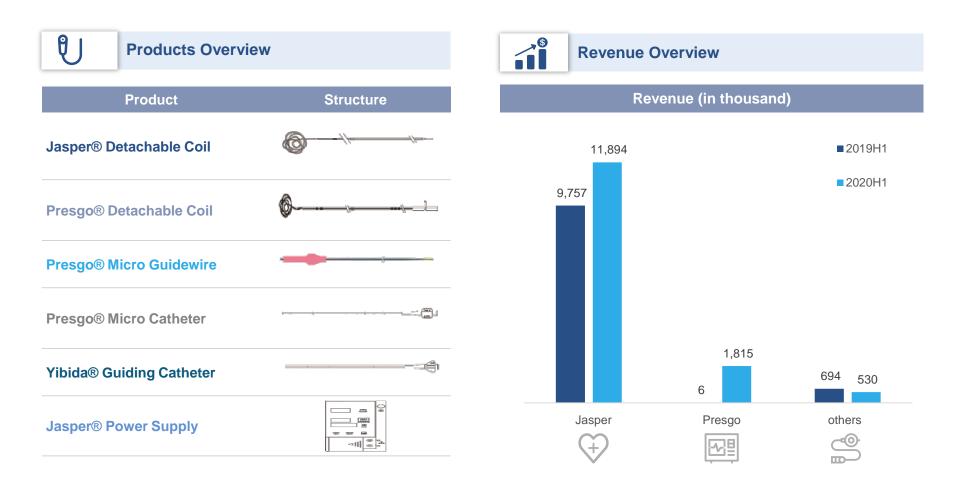
Gross Margin



Note: The data on this page is the parent financial statement of Achieva Medical from January to June 2020 and January to June 2019, rather than the consolidated statement. Achieva Medical 's operating results have been consolidated since March 29, 2019.



Neurointerventional Business: Commercialized Products



Note: The data on this page is the parent financial statement of Achieva Medical from January to June 2020 and January to June 2019, rather than the consolidated statement. Achieva Medical 's operating results have been consolidated since March 29, 2019.



Shenyi® Stent Retriever+ Progress Overview

September 2018

Shenyi® Stent Retriever completed animal experiments

July 2019

Started clinical trial enrollment of patients

August 2020

In clinical trial enrollment, 168 patients have been enrolled as of August 26

December 2020

Complete clinical trials and all patients enter into groups

2021 Q2

Submit registration applications to NMPA

Prospective, multi-center, randomized and control clinical trial

Submit registration to NMPA

Approved by NMPA

The "2019 Update on 2018 Guidelines on the Early Management of Acute Ischemic Stroke. AHA/ASA" proposes that for large vessel occlusion and ASPECTS≥6 within 6 hours of onset, it is recommended to perform mechanical thrombus removal based on the results of CT and CTA (or MRI and MRA) without other imaging evaluations.

expected

Clinical Design and Research Center of Shenyi® Stent Retriever

- Prospective, multi-center, randomized and control trial; test groupShenyi® Stent Retriever, control group Solitaire FR, 1:1, 236 cases in total
- Primary endpoint: success rate of reperfusion after occlusion of vascular (mTICI ≥ 2b, independent evaluation by core laboratory)

11 participating centers

(listed on the right)

Centers Changhai Hospital Jianmin Liu 01 The First Affiliated Hosptal of 02 Qi Fang Soochow University The First Affiliated Hospital of Jinan Li'an Huang University 04 Zhongshan People's Hospital Zhian Han Henan Provincial People's Hospital Tianxiao Li 05 The Affiliated Hospital of Xuzhou 06 Xinchun Ye **Medical University** 07 Maoming People's Hospital Geng Liao 80 Nanjing First Hospital Hongchao Shi The Second Affiliated Hospital of 09 Guodong Xiao Soochow University Chenzhou No.1 People's Hospital 10 Xiaoxi Yao Guangdong Provincial People's 11 Chengbo Dai Hospital

Operators' Assessment

"Shenyi® Stent Retriever has a unique and novel design, which guarantees the success of the operation. For example, the "net pocket" design of the front section of the stent reduces embolism "escape" during the long pull-back process; the stent also has a longer working length, a larger mesh design, and good adhesion and chimeric thrombosis ability to increase the success rate of the operation. Overall the stent is well visualized and has good visibility, thereby improving the controllability and safety."

----Neurology Dept., Guangdong Provincial People's Hospital

"Shenyi® Stent Retriever has a net pocket in the front end, which can partially prevent thrombus from escaping. The Stent Retriever can visualize the opening of the lesion in the whole process and provide information for treatment by assisting in judging the presence of vascular stenosis in situ and the degree of chimeric thrombus, etc. The Stent Retriever can be delivered through a 0.017 inch micro catheter, which is a big advantage."

— Cerebrovascular Intervention Dept., Zhongshan People's Hospital



SacSpeed® Balloon Dilatation Catheter

August 2020

Obtained the NMPA Registration Certificate

Registered examination

Submit registration to NMPA

Approved by NMPA

<u>Defining a new era of intracranial artery stenosis</u> <u>rapid exchange and precise expansion</u>





RCT research shows no worse than imported products



Better passability than similar products



More complete specifications and sizes

Target Market



Prefecture-level city stroke center



Neurology is the main department with interventional department and neurosurgery department supplemented



Mainly treat ischemic stroke

Other Neurointerventional Procedural Products

1. NMPA Registration Stage

Intermediate catheter* Used to introduce interventional procedural devices and diagnostic devices into peripheral and neuro vessel system

2. Other pre-application products candidates



Aspiration catheter



Thermal detachable coil

- To be applied to rebuild blood flow
- Used for the treatment of cerebral aneurysm and dural arteriovenous fistula

Intracranial stent

Used in cooperation with balloon dilatation catheter to alleviate ischemia

Jasper Supersoft detachable coil

Used for the treatment of cerebral aneurysm and dural arteriovenous fistula



 Used to introduce interventional procedural devices and diagnostic devices into peripheral and neuro vessel system



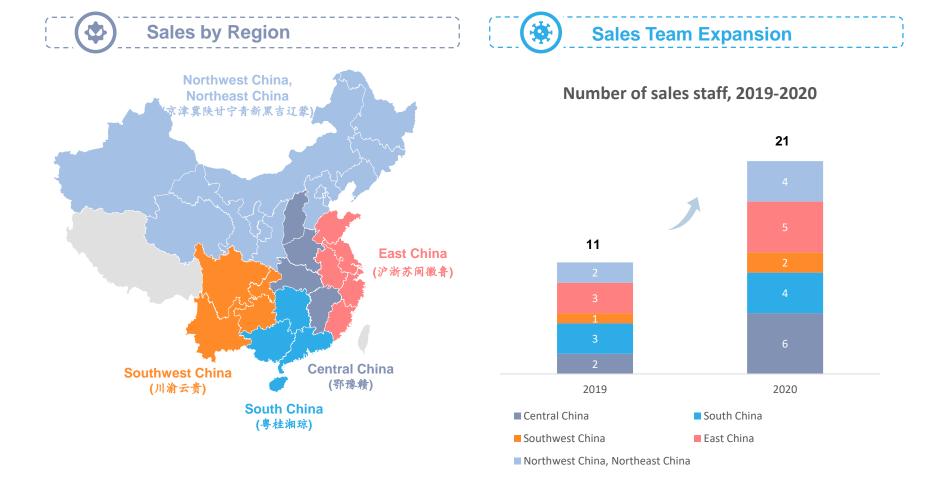
 Used for dilating stenosis to help with intracranial blood supply, and used to provide the channel for other diagnostic or therapeutic medical devices



 To assist the introduction of catheter to peripheral or neuro blood vessels. The balloon may temporarily block the vessel for angiography

Note: + means being exempted from clinical trial requirements

Neurointerventional Business: Sales Revenue Overview



Note: The number of sales staff is as of June 30, 2020



Neurointerventional Business: Production



Shanghai Production Facility



Competitive salary





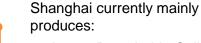


Suzhou Production Facility



Attract more potential employees

Shanghai Production Facility





- Jasper Detachable Coil
- Presgo Detachable Coil
- Presgo Micro Guidewire
- Presgo Micro Catheter
- Jasper Power Supply

Suzhou Production Facility



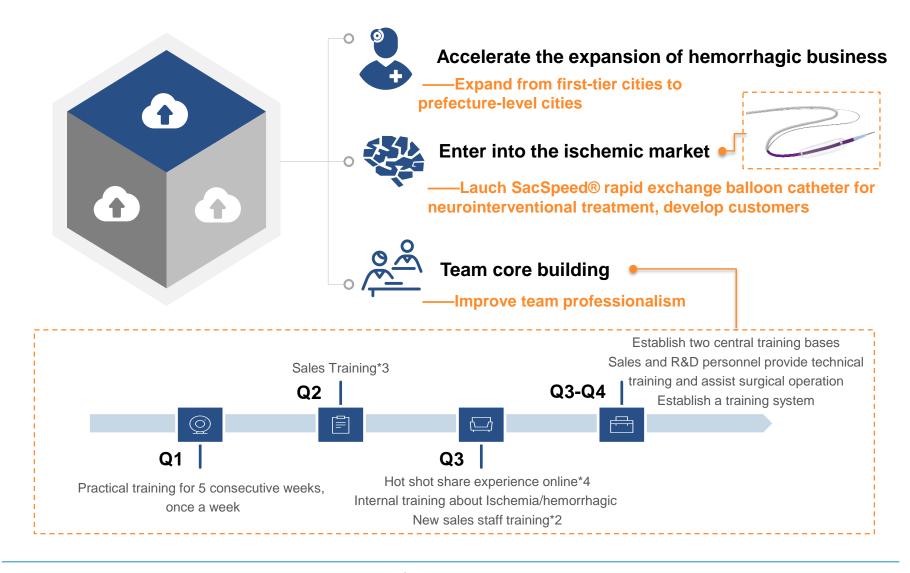
Aug 2020, Achieva Suzhou facility obtained Jasper ® commissioned production license

- So far, we have obtained the production license for the Yibida Guiding Catheter and commissioned production license for Jasper® in Suzhou.
- It is planned to relocate most of the production to the Suzhou facility within this year. The Suzhou facility will become our main manufacturing facility gradually.

The pilot program of the Jiangsu medical device registrant system: eligible medical device registration applicants can apply for a medical device registration certificate separately, and then commission the production to a qualified and capable production enterprise



Neurointerventional Business: 2020H2 Marketing Strategies & Focuses





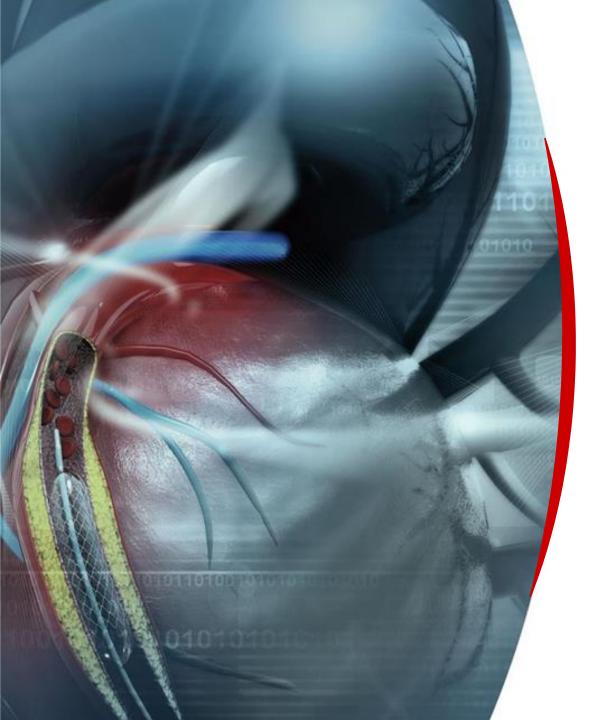


Impact of COVID-19

Impact of COVID-19

 	TaurusOne [®]	The stamping process of the clinical branch center report was delayed and took about one month longer than expected. Still expect to submit the application in Q3, 2020.
 	TaurusElite®	The enrollment speed of clinical patients has exceeded expectations, and the enrollment process may be completed earlier than originally expected.s
R&D	TaurusNXT®	The new travel restrictions have caused animal experiments to be transferred back to China. The plan to start human clinical trials in the first half of next year remains unchanged
***	TMVR	The new travel restrictions have caused animal experiments to be transferred back to China for a slight delay. It is expected that the human clinical program will not change next year
 	TaurusWave	COVID-19 has delayed the development of human clinical trials, and it is being evaluated whether to transfer back to China or use remote teaching
 	Shenyi® Stent Retrieve	r The enrollment speed of clinical patients has been slowed down due to COVID-19
Commercialization	Heart Valve Business	Recruitment of marketing department, sales department, and medical technology team continues without being impacted
¥	Neurointerventional Business	Bidding was stopped for a period of time, which had a negative impact on the speed of admission of new products; The coil business still has significant growth, which has a significant advantage over competing products. It is expected that the second-generation coil will continue to increase hospital coverage in the second half of the year, and it will also be able to exert strength
Production		Peijia was the first batch of Suzhou companies to resume work, and has resumed work completely by now; Increased the safety stock of raw materials during COVID-19; Currently everything is in order with external suppliers



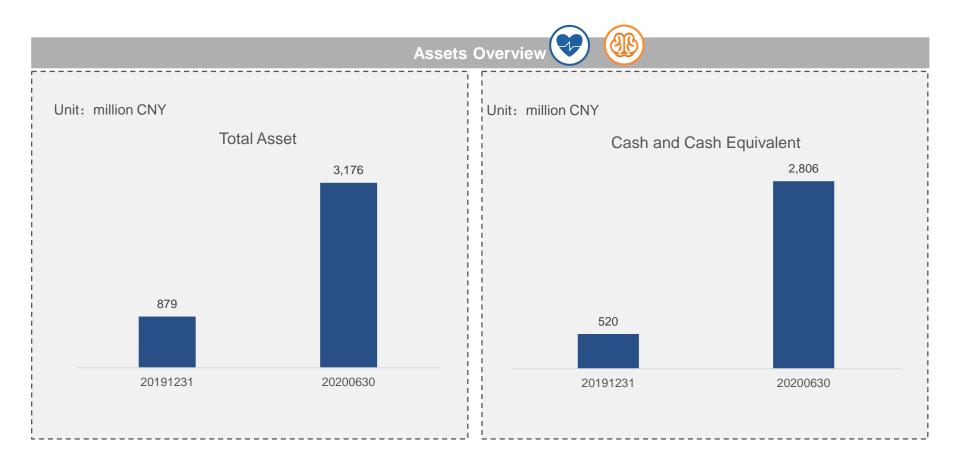




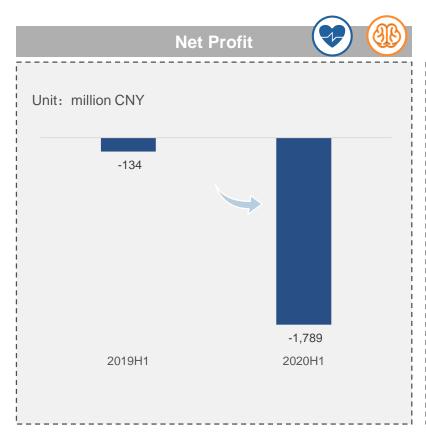
03

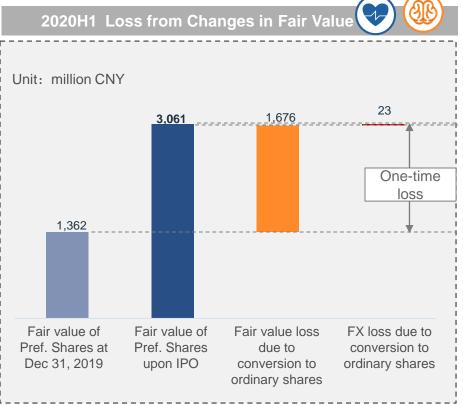
Financial Overview

Financial Overview – the Group

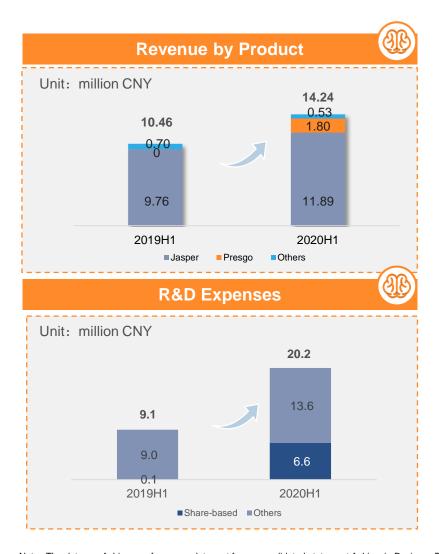


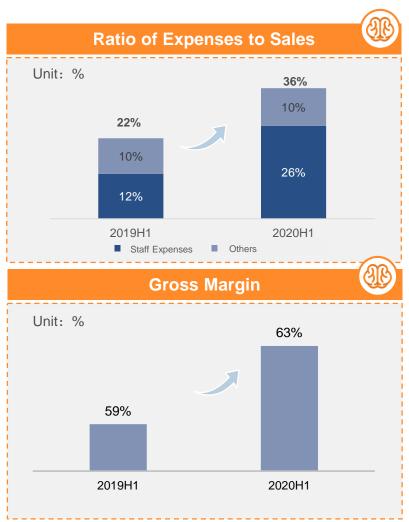
Net Profit & Loss from Changes in Fair Value – the Group





Financial Overview – Neurointerventional Business (Achieva)

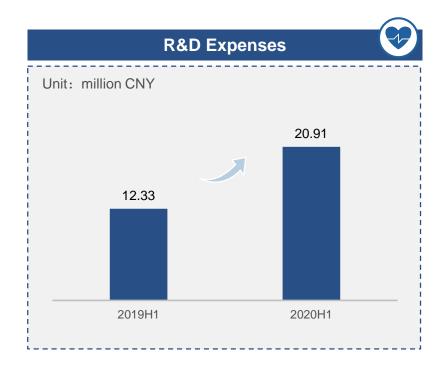


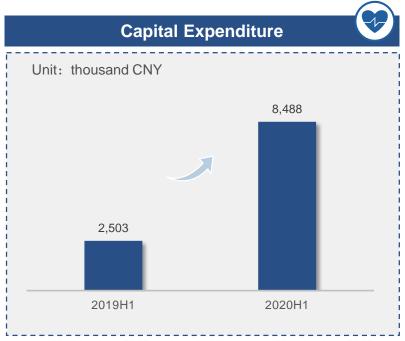


Notes: The data are Achieva performance data, not from consolidated statement. Achieva's Business Performance has been merged since March 2019.



Financial Overview – Heart Valve Business (Peijia)









04

Appendix

Focusing on the High-growth Transcatheter Valve Therapeutic and Neurointerventional Procedural **Medical Device Markets in China**



The Transcatheter Valve Therapeutic Medical Device Market in China

Market Size of TAVR Products in China, 2017-2025E

(US\$MM⁽¹⁾, at Ex-Factory Price Level)



- At early stage of development
- Significant growth potential
- No single dominating player(2)
- Continue to be led by a few domestic players⁽²⁾



Kev Success Factor

To develop advanced products with features tailored to the needs of Chinese patients and physicians



The Neurointerventional Procedural Medical Device Market in China

Market Size of Embolization Coil in China, 2014-2025E

(US\$MM(1), at Ex-Factory Price Level)

Source: Literature review and Frost & Sullivan analysis



(US\$MM(1))

2014-2025E

Market Size of MT Device in China,



- Dominated by several global medical device giants
- Domestic players are expected to gradually increase their market shares
- Technology advancements
- Improvements in products
- More favorable policies



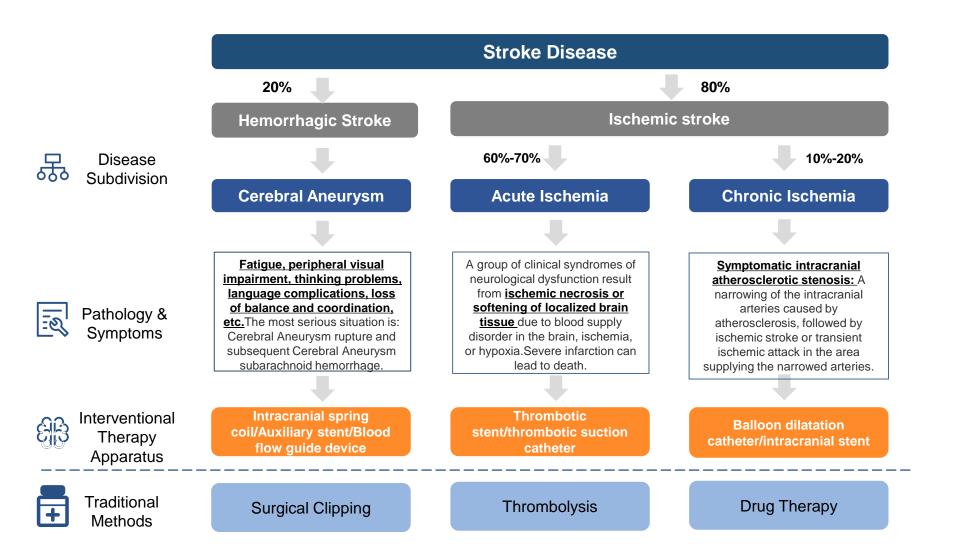
Kev Success Factor

To develop a comprehensive product portfolio tailored to the needs of Chinese patients and physicians

Notes:

- 1. The translations between Renminbi and U.S. dollars were made at the rate of CNY6.8845 to US\$1.00 as of January 15, 2020
- 2. According to Frost & Sullivan

Neurointerventional Business: Stroke Disease Overview







05

Financial Statements

Consolidated Statement of Comprehensive Loss

January - June 2020

CNY'000, (Unaudited)	6 months ended 30 June	
	2020	2019
Revenue	14,239	5,239
Cost of sales	(5,285)	(1,831)
Gross profit	8,954	3,408
Selling and distribution expenses	(5,162)	(1,301)
Administrative expenses	(65,325)	(28,938)
Research and development expenses	(41,164)	(16,221)
Other income	7,157	1,413
Other gains/(losses) - net	(2,788)	(1,591)
Operating loss	(98,328)	(43,230)
Finance income	7,908	50
Finance costs	(22,980)	(4,373)
Finance costs – net	(15,072)	(4,323)
Fair value change in financial		
instruments issued to investors	(1,675,526)	(86,037)
Loss before income tax	(1,788,926)	(133,590)
income tax expense	_	-
Loss for the year/period	(1,788,926)	(133,590)
Other comprehensive income:		
Items that will not be reclassified to profit or loss:		
- Fair value change relating to preferred shares due to own credit risk	-	(11,019)
Other comprehensive income for the year/period, net of tax		(11,019)
Total comprehensive loss for the year/period	(1,788,926)	(144,609)
Total comprehensive loss attributable to:		
- Owners of the Company	(1,788,926)	(144,609)
- Non-controlling interests	-	-
Total comprehensive loss for the year/period	(1,788,926)	(144,609)



Consolidated Balance Sheets

As of 30 June 2020

CNY'000 —	As at	As at	
	30 June 2020	31 December 2019	
Non-current assets			
Right-of-use assets	5,790	6,394	
Property, plant and equipment	75,602	70,241	
Investment properties	21,857	22,460	
Intangible assets	217,243	219,308	
Prepayments and other receivables	2,809	3,455	
Total non-current assets	323,301	321,858	
Current assets			
Inventories	15,361	11,163	
Financial assets at fair value through profit or loss	210,000	15,000	
Prepayments and other receivables	31,562	26,836	
Cash and cash equivalents	2,595,607	504,627	
Total current assets	2,852,530	557,626	
Total assets	3,175,831	879,484	
Current liabilities			
Lease liabilities	1,262	1,233	
Trade and other payables	40,220	47,641	
Contract liabilities	620	1,313	
Total current liabilities	42,102	50,187	
Non-current liabilities			
Financial instruments issued to investors	-	1,362,309	
Lease liabilities	490	1,129	
Deferred tax liabilities	20,320	20,320	
Deferred income	3,519	3,591	
Trade and other payables	154	154	
Total non-current liabilities	24,483	1,387,503	
Equity			
Equity attribute to owners of the Company			
Share capital and share premium	5,522,442	79,563	
Other reserves	37,860	35,298	
Accumulated losses	(2,451,056)	(673,067)	
Total equity	3,109,246	(558,206)	



Consolidated Statement of Cash Flows

January - June 2020

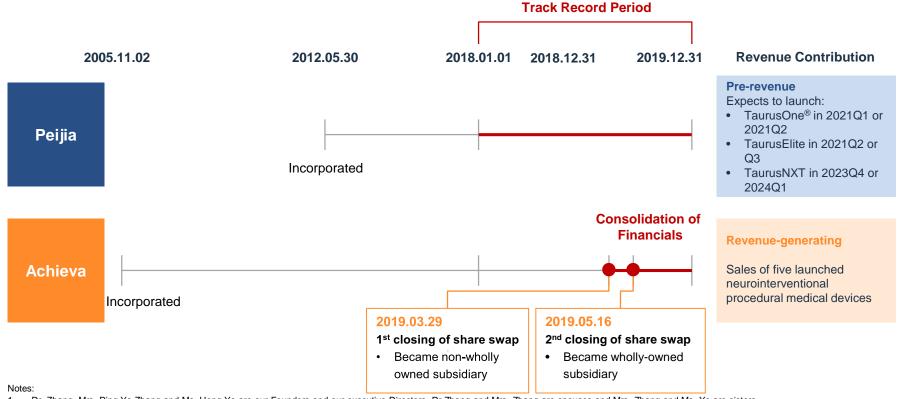
ONNESS	6 months ended 30 June	
CNY'000 —	2020	2019
Cash flows from operating activities		
Cash used in operations	(114,381)	(29,705)
Interest received	8,309	50
Interest paid	(51)	(59)
Net cash flow from operating activities	(106,123)	(29,714)
Cash flows from investing activities		
Payments for property, plant and equipment	(10,430)	(3,517)
Payments for intangible assets	(170)	-
Payments for financial assets at fair value through profit or loss	(532,000)	(29,000)
Proceeds from disposals of financial assets at fair value through profit or loss	337,000	-
Interest income received from financial assets at fair value through profit or loss	1,792	221
Cash acquired from acquisition of subsidiaries	-	59,622
Proceeds from disposal of property, plant and equipment	312	231
Net cash (outflow)/inflow from investing activities	(203,496)	27,557
Cash flows from financial activities		
Capital contribution from shareholders	4	21,567
Payments for shares bought back	-	(19,217)
Proceeds from issuance of ordinary shares	2,361,292	-
Proceeds from option exercise	24,520	-
Payments for listing expenses	(5,143)	-
Interest paid to borrowings from related parties	(691)	(390)
Repayment of borrowings from related parties	(2,301)	-
Interest paid for a bank loan	-	-
Principal elements of lease payments	(610)	(849)
Net cash inflow from financing activities	2,377,071	1,111
Net increase in cash and cash equivalents	2,067,452	(1,046)
Cash and cash equivalents at beginning of the year/period	504,627	94,762
Exchange gains on cash and cash equivalents	23,528	146
Cash and cash equivalents at end of the year/period	2,595,607	93,862



Financial Contribution

Achieva's results of operations has been consolidated since March 29, 2019

- Our Founders⁽¹⁾ have been the single largest group of shareholders of Achieva Medical with over 30% equity interests since its incorporation
- The operations of our Company and Achieva have been managed and overseen by our Founders since their respective inception
- A majority of the shareholders of our Company and of Achieva held equity interests in both our Company and Achieva prior to the Share Swap



- 1. Dr. Zhang, Mrs. Ping Ye Zhang and Ms. Hong Ye are our Founders and our executive Directors. Dr. Zhang and Mrs. Zhang are spouses and Mrs. Zhang and Ms. Ye are sisters
- 2. Due to COVID-19 situation, for conservative's sake, Company expected launch dates to move one quarter later for TaurusOne® (from 2020Q4 or 2021Q1 to 2021Q1 or 2021Q2) and TaurusElite (from 2021Q1 or 2021Q2 to 2021Q2 or 2021Q3)

