







DEFINITIONS

In this interim report, unless the context otherwise requires, the following expressions shall have the following meanings.

"Asahi Intecc" Asahi Intecc Co., Ltd., a medical devices company incorporated under the laws of

Japan with limited liability on 8 July 1976, and all of its subsidiaries

"Audit Committee" the audit committee of the board

"Award Share" in respect of a Selected Participant, such number of Scheme Shares (or such

number of returned shares) as determined by the Board and awarded to each of the

Selected Participant(s) for the purpose of the Award

"Board" the board of Directors

"Business Day" a day (other than Saturday, Sunday and public holidays) on which the Stock

Exchange is open for trading and on which banks are open for business in Hong

Kong

"BVI" the British Virgin Islands

"CE" French acronym for "Communate Europpene", a certification mark that indicates

conformity with health, safety and environmental protection standards for products

sold within the European Economic Area

"CG Code" the corporate governance code as contained in Appendix 14 to Listing Rules

"Company" or "we" or "us" or

"our"

MicroPort NeuroTech Limited (微創腦科學有限公司), an exempted company

incorporated in the Cayman Islands, the shares of which are listed on the Main

Board of the Stock Exchange (stock code: 2172)

"Director(s)" director(s) of the Company, including all executive, non-executive and independent

non-executive directors

"FDA" the United States Food and Drug Administration

"Frost & Sullivan" Frost & Sullivan (Beijing) Inc., our industry consultant

"Global Offering" the global offering of the shares, details of which are set forth in the Prospectus

"Group" the Company and its subsidiaries

"Hong Kong" the Hong Kong Special Administrative Region of the People's Republic of China

"HKFRSs" Hong Kong Financial Reporting Standards

"KPMG" KPMG, Certified Public Accountants

"Listing" the listing of the shares on the Main Board of the Stock Exchange



"Listing Date"	15 July 2	2022 the date on	which dealings in the	shares on the Main Board first
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"Listing Rules" the Rules Governing the Listing of Securities on The Stock Exchange of Hong Kong

Limited

"Main Board" the stock exchange (excluding the option market) operated by the Stock Exchange

which is independent from and operated in parallel with the Growth Enterprise Market of the Stock Exchange. For the avoidance of doubt, the Main Board

excludes the Growth Enterprise Market of the Stock Exchange

"MicroPort" MicroPort Scientific Corporation (微創醫療科學有限公司), an exempted company

incorporated in the Cayman Islands with limited liability whose shares are listed on the Main Board of the Stock Exchange (stock code: 00853), and one of our

Controlling Shareholders

"Model Code" Model Code for Securities Transactions by Directors of Listed Issuers as contained

in Appendix 10 to the Listing Rules

"NHSA" National Healthcare Security Administration

"Prospectus" the prospectus of the Company dated 29 June 2022

"Prior-year Period" the six months ended 30 June 2022

"MP Scientific" MicroPort Scientific Investment LTD, a company incorporated in the BVI with

limited liability on September 30, 2020 and is a direct wholly owned subsidiary of

MicroPort, and one of our Controlling Shareholders

"NMPA" National Medical Products Administration (國家藥品監督管理局) and its

predecessor, the China Food and Drug Administration (國家食品藥品監督管理總局)

"PRC" the People's Republic of China

"Rapid Medical" Rapid Medical Ltd., a company incorporated in the State of Israel with limited

liability on 12 August 2008, which is primarily engaged in the development, manufacturing and sales of innovative devices for neuro-interventional procedures

and is indirectly owned as to 22.28% equity by the Company

"Reporting Period" for the six months ended 30 June 2023

"RMB" Renminbi, the lawful currency of the PRC

"R&D" Research and development

"Selected Participant" eligible participant selected by the Board to participate in the Scheme

"Share Award Scheme" a share award scheme adopted by the Board on 26 August 2022

Definitions (Continued)

"SFO" the Securities and Futures Ordinance (Chapter 571) of Hong Kong, as amended,

supplemented or otherwise modified from time to time

"share(s)" ordinary share(s) of the Company

"Shareholder(s)" holder(s) of the share(s)

"Stock Exchange" The Stock Exchange of Hong Kong Limited

"subsidiaries" has the meaning ascribed thereto under the Listing Rules

"Trustee" the original trustee and any additional or replacement trustees, being the trustee

or trustees for the time being of the trusts declared in the trust deed entered into between the Company and trustee, which are independent third parties and not

connected with the Company or any of its connected persons

"Vesting Date" the date on which the Trustee may vest the legal and beneficial ownership of

the Awarded Shares (or the relevant portions thereof) in the relevant Selected

Participant

"WE'TRON CAPIAL" WE'TRON CAPITAL LIMITED (中國微創投資管理有限公司), a company incorporated

in Hong Kong with limited liability on October 26, 2005

"%" per cent

CORPORATE INFORMATION



Executive Directors:

Mr. Xie Zhiyong (謝志永)

Mr. Wang Yigun Bruce (王亦群)

Non-Executive Directors:

Mr. Peng Bo (彭博) (Chairperson of the Board)

Mr. Wang Lin (王琳) Ms. Wu Xia (吳夏)

Independent Non-Executive Directors:

Dr. Xu Yi (胥義)

Dr. Zhang Haixiao (張海曉)

Mr. Siu Chi Hung (蕭志雄)

AUDIT COMMITTEE

Mr. Siu Chi Hung (蕭志雄) (Chairperson)

Dr. Xu Yi (胥義)

Dr. Zhang Haixiao (張海曉)

REMUNERATION COMMITTEE

Dr. Xu Yi (胥義) (Chairperson)

Mr. Peng Bo (彭博)

Mr. Siu Chi Hung (蕭志雄)

NOMINATION COMMITTEE

Dr. Zhang Haixiao (張海曉) (Chairperson)

Mr. Xie Zhiyong (謝志永)

Dr. Xu Yi (胥義)

REGISTERED OFFICE

Tricor Services (Cayman Islands) Limited Third Floor, Century Yard, Cricket Square

P.O. Box 902

Grand Cayman, KY1-1103

Cayman Islands

HEAD OFFICE AND PRINCIPAL PLACE OF BUSINESS IN THE PRC

No. 1661 Zhangdong Road Pudong New Area, Shanghai PRC

PRINCIPAL BANKERS

China Construction Bank Shanghai Zhangjiang Branch

220 Keyuan Road Pudong New Area Shanghai

PRC

PRC

Bank of China Shanghai Zhoupu Branch

1st Floor, Wanda Plaza No. 3435 Hunan Road Pudong New Area Shanghai

Shanghai Pudong Development Bank Co., Ltd. Zhangjiang Keji Branch

No. 56 Boyun Road Pudong New Area Shanghai

PRC

PRINCIPAL PLACE OF BUSINESS IN HONG KONG

5/F, Manulife Place 348 Kwun Tong Road Kowloon Hong Kong

AUTHORISED REPRESENTATIVES

Mr. Peng Bo (彭博) Ms. Hui Yin Shan (許燕珊)

COMPANY SECRETARY

Ms. Hui Yin Shan (許燕珊)

AUDITOR

KPMG

Public Interest Entity Auditor registered in accordance with the Accounting and Financial Reporting Council Ordinance 8th Floor, Prince's Building 10 Chater Road Central Hong Kong

LEGAL ADVISER

Clifford Chance

27th Floor, Jardine House One Connaught Place Hong Kong

COMPLIANCE ADVISER

Somerley Capital Limited

20/F, China Building 29 Queen's Road Central Hong Kong

PRINCIPAL SHARE REGISTRAR AND TRANSFER OFFICE IN THE CAYMAN ISLANDS

Tricor Services (Cayman Islands) Limited

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HONG KONG BRANCH SHARE REGISTRAR AND TRANSFER OFFICE

Computershare Hong Kong Investor Services Limited

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WEBSITE

www.medneurotech.com

STOCK CODE

2172

LISTING DATE

15 July 2022

PRESIDENT'S STATEMENT



Executive Director and President Mr. Xie Zhiyong

In the first half of 2023, with the concerted efforts of all the employees, the Group had achieved a breakthrough growth in various operating results. During the Reporting Period, the Group realised revenue of approximately RMB299.2 million, representing a substantial increase of 45.2% over the Prior-year Period; the profit from operations was approximately RMB82.2 million, representing an increase of approximately 505.2% over the Prior-year Period; the net profit reached approximately RMB58.0 million, representing a turnaround from losses to profits as compared to the Prior-year Period.

The Group continues to export innovative and high-quality treatment concepts for cerebral vessel diseases to the industry. During the Reporting Period, our products had established the presence in over 200 new hospitals, and supported approximately 145,000 procedures in more than 2,800 hospitals across the country. As the related products of the Group have won the bids in all the centralized procurement projects carried out in the past, the hospital admission and clinical promotion ushered in a breakthrough. In terms of cultivating low-tier markets, the Group assisted in the establishment of national stroke centers at the grassroots level through the "Eagle & Swallows (神雕飛 燕)" program. In the first half of the year, our products were promoted to enter

more than 100 grassroots hospitals, and had been clinically used in a total of approximately 700 grassroots hospitals, covering more than 200 low-tier cities and counties. Among them, multiple market-share leading products including Tubridge® Flow-diverting Stent performed particularly well in the grassroots market.

In the international market, the Group's overseas revenue amounted to approximately RMB15.0 million in the first half of 2023. As of the end of the Reporting Period, the Group's products have been commercialized in a total of 12 overseas countries, covering half of the countries ranking top 10 worldwide in terms of the number of neuro-interventional procedures. In addition, the Group has successfully implemented the direct sales operation model for the first time in both Ireland and the United Kingdom, which not only better suits the local market demand and marketing habits, but also significantly improves operational efficiency.

As for the research and development, from the beginning of 2023 and up to the date of this report, the Group received the approvals for marketing in China of four new products, including the Tigertriever® Revascularization Device, W-track® Intracranial Aspiration Catheter, Q-track® Microcatheter and Veyronwire™ Neurovascular Guide Wire; three products were under the review process by the NMPA, including the Neurovascular Balloon Guide Catheter, Neurohawk® Stent Thrombectomy Device 2 and Distal Protection Device. Meanwhile, several clinical projects of Tubridge® series flow diverting stents have made significant progress: the PARAT MINI clinical study for the treatment of wide-neck, small and medium-sized aneurysms have completed the enrollment of all patients; the pre-marketing clinical study of Tubridge Plus® Flow-diverting Stent, the new generation product with full visualization has also successfully enrolled all patients.

President's Statement (Continued)

We are committed to establishing a comprehensive quality management system based on the "quality culture, quality system and quality tools (質量文化、質量體系、質量工具)". During the Reporting Period, we have obtained the MDSAP (Medical Device Single Audit Program), a quality system certification accepted in five countries, effectively reducing the auditing cost of products entering overseas markets.

We further accelerated the localization of raw materials and improved the lean production capacity. As of the end of the Reporting Period, the localization rate of raw materials for our products had reached over 90%. At the same time, we had steadily increased the production efficiency, improving the quality and efficiency in an all-round way.

Looking forward to the future, MicroPort NeuroTech™ will continue to execute its development strategy of "global integration vision, sustainable quality development, consolidated position in neuro-intervention, expansion of neuro tech field", firmly adopt the "physician-engineer collaboration (醫工結合)" R&D model, and create the innovative product portfolio with international competitiveness, so as to provide the high-quality and accessible solutions for cerebral vessel diseases to patients around the world.

The Directors, senior management and all employees of the Group will uphold the principles of integrity, diligence and responsibility to pursue a high-quality development. On behalf of all my colleagues in the Company, I hereby would like to extend our heartfelt thanks to all shareholders and partners for the long-term companionship and support.

Executive Director and President

Mr. Xie Zhiyong



For the six months ended 30 June

	2023 RMB′000 (unaudited)	2022 RMB'000 (unaudited)	Change %
Revenue	299,193	205,993	45.2%
Gross profit	232,637	141,547	64.4%
Profit from operations	82,200	13,582	505.2%
Profit/(loss) for the period	57,999	(93,729)	N/A
Earnings/(loss) per share	0.11	(0.20)	N/A

For the six months ended 30 June 2023, the Group's revenue was mainly derived from hemorrhagic stroke products, cerebral atherosclerotic stenosis products, acute ischemic stroke products and access products. During the Reporting Period, the Group recorded the revenue of approximately RMB299.2 million, representing an increase of around 45.2% over that of approximately RMB206.0 million for the Prior-year Period.

The increase was mainly due to: (1) market-share leading products (including Tubridge® Flow-diverting Stent ("Tubridge® Flow-diverting Stent") etc.) continued to increase the market penetration, further consolidating the competitive advantages, and maintaining a positive growth momentum; (2) The continued tendering and admission of several new products launched by the Group in recent years (including NUMEN® Coil Embolization System ("NUMEN® Coil"), Bridge® Rapamycin Target Eluting Vertebral Artery Stent System ("Bridge® Vertebral Artery DES") and U-track® Intracranial Support Catheter System ("U-track® Support Catheter"), etc.) into hospitals has contributed to the exploration of blank markets; newly approved products in 2022 (including Neurohawk® Stent Thrombectomy Device ("Neurohawk® Thrombectomy Device"), Diveer® Intracranial Balloon Dilatation Catheter ("Diveer® Balloon Catheter"), etc.) accelerated the market development, contributing to the Group's revenue growth.

During the Reporting Period, the Group recorded the profit from operations of approximately RMB82.2 million, representing an increase of approximately 505.2% as compared to that of approximately RMB13.6 million for the Prior-year Period, which was mainly attributable to the increase in revenue and gross profit.

During the Reporting Period, the Group recorded the profit for the period of approximately RMB58.0 million, representing the losses turning into profits as compared to the Prior-year Period.

MANAGEMENT DISCUSSION AND ANALYSIS

INDUSTRY OVERVIEW

Stroke is an acute cerebral vessel disease, which is the second major fatal disease in the world and the first major fatal disease in China, with high rates of incidence, disability, mortality and recurrence. According to the research data of the Global Burden of Disease (GDB), China had the largest number of stroke patients in the world, and the proportion of patients younger than 70 years old kept increasing, with a trend toward younger patients. Another research result¹ on the burden of stroke disease in China shows that in 2020, the prevalence of stroke in China was 2.6% among people aged 40 years or older, which was much higher than the global prevalence of stroke, and the number of new strokes in China (approximately 3.4 million) was higher than that in the United States (approximately 0.61 million) and Europe (approximately 1.12 million), representing approximately a quarter of all new stroke cases worldwide each year. The research also shows that there were significant urban-rural differences in the burden of stroke disease in China, with both stroke incidence and mortality rates higher in rural areas than in urban areas.

Thanks to the development of neuroimaging, the neuro-interventional therapy is gradually replacing the traditional surgical craniotomy and conventional drug therapy with its safe, effective and minimal invasive characteristics, and has become an important method of stroke treatment. However, currently the neuro-interventional medical device industry in China is still at an early stage of development, with a relatively low market penetration especially in the vast grassroots areas represented by low tier cities and counties. In recent years, relevant policies have been successively issued to encourage and support the development of the neuro-interventional industry in terms of stroke diagnosis, treatment and prevention, medical device innovation, technical specifications and supervision, etc.

As an important aspect of "Healthy China Construction" (健康中國建設), China has been gradually establishing and improving the prevention, diagnosis and treatment policy for stroke. In 2021, multiple departments including the NMPA jointly formulated the Comprehensive Plan for Strengthening Stroke Prevention and Treatment Work to Reduce Millions of New Disabilities (《加強腦卒中防治工作減少百萬新發殘疾工程綜合方案》), which proposes the overall goal of further improving the prevention and treatment effect of stroke and reducing the incidence rate and disability rate, and clarifies that the phased goals to be achieved by 2022, 2025, and 2030, including the awareness rates of hypertension among residents, the development of intravenous thrombolysis and thrombectomy techniques, etc. Meanwhile, the "Identification and Hierarchical Diagnosis and Treatment Action for the Stroke in China's Thousands of Counties and Ten Thousands of Towns (中國千縣萬鎮卒中識別與分級診療行動)" was accelerated to open up the green channel for stroke treatment, and establish and improve the hierarchical diagnosis and treatment model for stroke as a specific disease. According to the National Health Commission of the PRC, as of the end of August 2023, an aggregate of 1,827 stroke centers were established in the country, including 603 advanced stroke centers (inclusive of construction units), 546 comprehensive prevention stroke centers and 678 prevention stroke centers. With the rapid growth of the number of stroke centers and the continuous improvement of the stroke first-aid maps, the capacity and the coverage of diagnosis and treatment in grassroots areas have been further improved.

Burden of stroke in China in 2020, JAMA Netw Open. 2023;6(3):e231455

In addition, the pharmaceutical and healthcare system reform in the PRC continued to deepen. In March 2023, the NHSA issued the Notice on Pharmaceutical Centralized Procurement and Price Management in 2023 (《關於做好2023 年醫藥集中採購和價格管理工作的通知》), which proposes to conduct a new batch of national organized procurement of high-value medical consumables based on the principle of "one product, one policy", establishing the overall tone for centralized procurement of drugs and medical consumables. In June 2023, Beijing issued the Management Plan on the Linage of DRG Payment and Volume-based Procurement in Medical Institutions (《醫療機構DRG付費和帶 量採購聯動管理方案》), aiming to correlate DRG payment with volume-based procurement (VBP) to bring the winning price of VBP into the consideration of DRG weight. This policy will motivate medical institutions to further optimize cost and resource management and has an important reference significance for the form and direction of future centralized VBPs. In July 2023, six departments including the National Health Commission jointly issued the Key Tasks for Deepening the Reform of the Medical and Health System in the Second Half of 2023 (《深化醫藥衛生體 制改革2023年下半年重點工作任務》), which clarifies the development priorities for deepening the medical reform in the next stage: from the perspective of medical insurance coverage, treatment and surgical projects with clear clinical efficacy and significant technical value will be prioritized to be included into the adjustment scope; from the perspective of payment mode, while adjusting the payment structure through the centralized VBPs of drugs and medical devices as well as the innovative medical insurance negotiations, no less than 70% of the coordinated regions are required to carry out DRG/DIP reform by the end of 2023. Under the reform of diversified and composite medical insurance payment methods, medical devices with the clear clinical value and the strong rigid treatment demand are expected to usher in faster growth, while auxiliary attributes and non-essential varieties are showing a weakening trend, which will bring a long-term and in-depth impact on the structure of clinical medical products.

COMPANY'S BUSINESS

As a pioneer and the largest Chinese company in the neuro-interventional medical device industry in China, the Group is committed to providing innovative and accessible solutions for cerebral vessel diseases to patients and physicians around the world. The Group has a comprehensive portfolio of commercialized products covering three major areas of cerebral vessel diseases, namely hemorrhagic stroke, cerebral atherosclerotic stenosis and acute ischemic stroke. According to Frost & Sullivan, the Group is the only Chinese company among the top five players in China's neuro-interventional medical device market, with the market share of approximately 8% in terms of the sales in 2022, nearly double that of 2020 while ranking the first among all the domestic brands for many years.

Since its establishment, while always adhering to the goal of addressing clinical needs, the Group has been placing key emphasis on R&D and innovation with independent intellectual property rights. After years of experiences, the Group has already mastered a number of core design and manufacturing technology platforms for the R&D and manufacturing of neuro-interventional medical devices. The Group has developed multiple "first" or "only" products, including the first stent system for treating intracranial atherosclerotic diseases in the world, the only intracranial stent graft approved for treating cerebral vessel diseases in the world, the first Chinese-developed flow-diverting stents approved by the NMPA, and the first vertebral artery drug-eluting stent in China that has been admitted to the NMPA's innovative medical device special review and approval procedure (the "Green Path") and approved by the NMPA.

BUSINESS REVIEW

In the first half of 2023, by adhering to the sinking strategy of sales channel, the Group actively responded to changes in the external policy environment and accelerated the global business layout, thus achieving a rapid growth in the operating performance with the profitability greatly improved. During the Reporting Period, the Group achieved the revenue of RMB299.2 million, representing an increase of approximately 45.2% over the Prior-year Period. The profit from operations was RMB82.2 million, representing an increase of approximately 505.2% over the Prior-year Period. The profit for the period was RMB58.0 million, turning losses into profits as compared to the Prior-year Period.

COMMERCIALIZATION CAPABILITIES

The Group has established a team for the promotion of medical solutions with the professional medical background and extensive experiences. The team continues to export innovative neuro-interventional treatment concepts to the market, and provides patients and physicians with an integrated solution to treat cerebral vascular diseases, including the promotion and education regarding the surgical methods and products, recommendations for treatment options, training on surgery and surgical devices, clinical support and postoperative follow-ups, so as to strengthen our leading position as a domestic brand.

As at the end of the Reporting Period, our team for the promotion of medical solutions consisted of nearly a hundred employees with an average industry experience of over 8 years. In addition, the Group has established cooperative relationships with more than 200 distributors and sub-distributors, and our sales channels cover 31 provinces, municipalities and autonomous regions across the country. The Group's products have been clinically used in more than 2,800 hospitals nationwide, covering more than 1,400 tertiary hospitals and all of the top 100 hospitals in China's National Stroke Center, cumulatively supporting approximately 145,000 neuro-interventional procedures.

In the first half of 2023, benefiting from the comprehensive and complete product matrix and the long-term accumulation of practitioner recognition, the Group's products have newly developed more than 200 hospitals, of which more than 100 were county-level hospitals, gradually consolidating the grassroots market. In the meantime, the provincial VBP projects of coils have been implemented gradually in various provinces, and Henan Province has also launched the VBP of neuro-interventional medical consumables among public medical institutions. The related products of the Group have won the bids in all the VBP projects carried out in the past, with hospital admission and clinical promotion ushering in a breakthrough.

In the field of hemorrhagic stroke products, Tubridge® Flow-diverting Stent, the Group's market-share leading product, further accelerated its market penetration with the focus on developing the second-tier and grassroots hospitals. During the Reporting Period, Tubridge® Flow-diverting Stent was newly admitted into more than 150 hospitals, covering more than 940 hospitals in total, which had also led to a rapid increase in the combinatorial usage of Fastrack® Microcatheter System ("Fastrack® Microcatheter"). NUMEN® Coil took the opportunity of winning the VBP bid to accelerate the development of new markets. During the Reporting Period, the product newly entered more than 150 hospitals and had achieved clinical applications in an accumulated number of 730 hospitals. WILLIS® Intracranial Stent Graft ("WILLIS® Stent Graft"), as the world's first and only approved intracranial stent graft, not only has excellent clinical effects in the treatment of complex cranial vascular diseases, but has also won a wide recognition by clinical experts with its continuously expansion of advantages in indications such as vascular rupture in nasopharyngeal carcinoma surgery and cervical dissection aneurysm.

In the field of cerebral atherosclerotic stenosis treatment products, the innovative product Bridge® Vertebral Artery DES has further increased the acceptance of the drug-loaded stent treatment concept by surgeons with its differentiated characteristics such as grooved drug loading design and low long-term restenosis rate. During the Reporting Period, Bridge® Vertebral Artery DES newly entered more than 230 hospitals, covering more than 820 hospitals in total. It had also won the exclusive bid in Henan Province's VBP, with the clinical usage increasing significantly. In addition, APOLLO™ Intracranial Stent System ("APOLLO™ Intracranial Stent") continued to accelerate the pace of hospital admission, and established the presence in more than 140 new hospitals during the Reporting Period, covering more than 2,030 hospitals in total. Diveer® Balloon Catheter has continued to accelerate the market introduction with its advantages of ultra-soft head end, ultra-low outside diameter, and easier passage through highly narrow lesions since its launch in 2022. As of the end of the Reporting Period, Diveer® Balloon Catheter has been listed on the procurement platforms of 29 provinces nationwide and entered more than 100 hospitals in total, further improving the Group's product portfolio in the treatment of atherosclerotic stenosis.

In the field of acute ischemic stroke treatment products and access products, as of the end of the Reporting Period, Neurohawk® Thrombectomy Device and X-track® Distal Catheter, which were newly launched in 2022, both had been listed on the procurement platforms of 27 provinces, and had entered over 250 and 100 hospitals respectively. As a key accessory in aneurysm treatment surgery, driven by the sales volume of related therapeutic products of the Group, U-track® Support Catheter has further leveraged its competitive advantages such as the high clinical adaptability and the complete channel integration, and had realized the year-on-year increase of 150% in terms of clinical usage during the Reporting Period, continuously bringing new momentum to the revenue growth.

As for the grassroots market, the Group actively responded to the national call for establishing primary stroke centers through the Eagle & Swallows (神雕飛燕) program. The Group has been providing the clinical training, follow-up consulting and routine guidance to physicians in hospitals in low-tier cities and counties, thereby helping grassroots hospitals to improve their stroke treatment ability. In the first half of 2023, Eagle & Swallows team established the presence in more than 100 new grassroots hospitals, with a total coverage of approximately 700 hospitals in over 200 lower-tier cities and counties. In addition, the Group continued to promote the high quality medical resources to those local areas through the special fund of "Brain Power" (百腦神通) for cultivating young neuro-interventional physicians, so as to build a platform for technical communication among grassroots clinicians, allowing more local patients with cerebral vessel diseases to benefit from the initiatives. As of the end of the Reporting Period, the Group had launched a total of 13 clinical education bases for the Brain Power program and provided technical trainings to around 160 surgeons.

The Group is committed to improving the global stroke clinical diagnosis and treatment technology and continues to provide professional training to doctors on clinical techniques and standardized diagnosis and treatment processes, gradually building up a customised, systematic and multi-level clinical training system. With the focus on the promotion of our innovative products, namely Tubridge® Stent, NUMEN® Coil and Bridge® Vertebral Artery DES, we have offered a series of innovative clinical therapies through the combination of several product portfolios, including the "AND procedure" (APOLLO™ Intracranial Stent + Neurohawk® Thrombectomy Device + Diveer® Balloon Catheter) for the treatment of large vessel occlusions associated with intracranial atherosclerotic stenosis (ICAS-LVO) and the "NEXT procedure" (Neurohawk® Thrombectomy Device + X-track® Distal Catheter) for the acute thrombectomy surgeries.

INTERNATIONAL BUSINESS

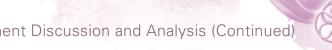
During the Reporting Period, the Group achieved a breakthrough in its international business with the overseas revenue of RMB15.0 million, representing an increase of 27.3% over the Prior-year Period.

As of the end of the Reporting Period, the Group's products have been commercialized in a total of 12 overseas countries, including South Korea, the United States, Brazil, Poland, Spain, Portugal, Chile, Ireland, the United Kingdom, Croatia, Greece and Argentina, covering half of the countries ranking top 10 worldwide in terms of the number of neuro-interventional procedures. In Ireland and the United Kingdom, the Group has successfully implemented a direct sales model, which has significantly improved the operational efficiency while better adapting to local market demand and marketing habits, adding new impetus to the growth of overseas business. In France, the Micro Frame and Micro Fill series of NUMEN® Coils have entered the national medical insurance reimbursement list. In the United States, the Group's NUMEN® Coil has been rapidly promoted through the existing sales channels of our associate company, Rapid Medical. Meanwhile, NUMEN® Coil can be used in conjunction with Rapid Medical's self-owned Comaneci® Embolization Assist Device ("Comaneci® Assist Device") to form a complementary product portfolio in the field of coil embolization procedures — "Numenaneci" (NUMEN® Coil + Comaneci® Embolization Assist Device). In the future, both parties will leverage their complementary strengths in terms of sales channels and product distribution to promote the application of innovative neurovascular disease solutions in the Chinese and global markets.

In terms of product admission and market promotion, multiple innovative products of the Group continued to debut in the international market. In the first half of 2023, NUMEN® Coils were approved successively for commercialization in Australia, Saudi Arabia, Colombia and Argentina, and Tubridge® Flow-diverting Stent and Fastrack® Microcatheter were also approved for marketing in Argentina. In the first half of 2023, the Group carried out a total of 7 overseas surgical training and academic exchange activities, and for the first time at the World Live Neurovascular Conference (WLNC), we promoted and exhibited Tubridge® Flow-diverting Stent and other products on a large scale in the form of independent exhibition stand, attracting many top clinical experts in neuro-intervention from around the world to observe and study.

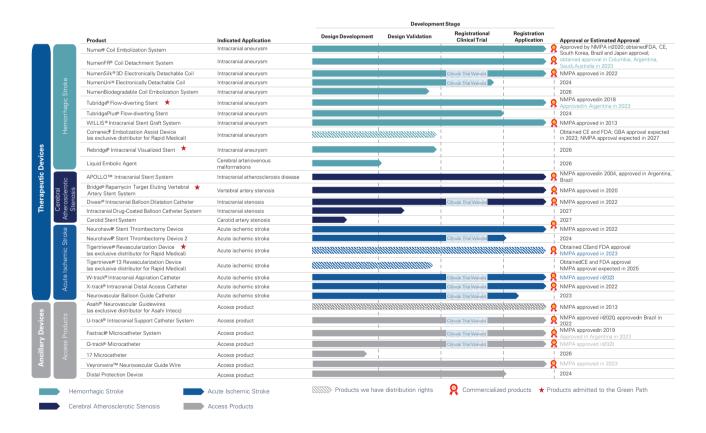
PRODUCT PIPELINE

Since the approval for marketing of the first product in 2004, leveraging its excellent R&D capability and efficient physician-engineer collaboration (醫工結合) model, the Group has built up a diversified portfolio of neuro-interventional products. As of the date of this report, the Group had a total of 17 products that have been approved and commercialized in China, and 13 pipeline products under different development stages. Among them, four products have been approved by the NMPA to be admitted to the Green Path, ranking first among Chinese neuro-interventional medical device companies.



From the beginning of 2023 and up to the date of this report, the Group's R&D projects have achieved fruitful results. Four products including Tigertriever® Revascularization Device ("Tigertriever® Revascularization Device"), W-track® Intracranial Aspiration Catheter ("W-track® Aspiration Catheter"), Q-track® Microcatheter and Veyronwire™ Neurovascular Guide Wire ("Veyronwire™ Guide Wire") have been approved by the NMPA for marketing; the registration applications of Neurovascular Balloon Guide Catheter ("Balloon Guide Catheter"), Neurohawk® Stent Thrombectomy Device 2 and Distal Protection Device have been submitted to the NMPA for approval. In addition, several clinical projects of Tubridge® Flow-diverting Stent have made significant progress: the PARAT MINI clinical study for the treatment of wide-neck, small and medium-sized aneurysms has completed the enrollment of all patients; the pre-marketing clinical study of the new generation of Tubridge Plus® Flow-diverting Stent with full visualization has also successfully completed the enrollment of all patients.

The following chart summarizes our product portfolio and development status as of the date of this report.



Hemorrhagic Stroke Products

Intracranial aneurysm is one of the main causes of hemorrhagic stroke. According to Frost & Sullivan, hemorrhagic stroke products represent the largest segment in terms of sales of neuro-interventional medical devices in China. The Group has a portfolio of eleven products for the treatment of hemorrhagic stroke, of which five products have been approved for commercialisation, including embolization coils, flow-diverting stents and stent grafts, and covering key therapeutic areas of hemorrhagic stroke. According to Frost & Sullivan, the market share of the Group's Tubridge® Flow-diverting Stent was more than 45%, ranking top two in the domestic market share and the first among domestic brands in terms of the number of procedures performed in 2022.

During the Reporting Period, the Group recorded the revenue of hemorrhagic stroke products of RMB206.8 million, representing an increase of 76.0% over the Prior-year Period, which was mainly due to the significant increase in the clinical use of Tubridge® Flow-diverting Stent and the increase in the global sales revenue of NUMEN® Coil.

NUMEN® Coil

NUMEN® Coil is a coil embolization system used to treat intracranial aneurysm. It was approved by the NMPA in September 2020, and was subsequently approved for commercialization in the European Union, South Korea, the United States, Brazil, Japan, Argentina, Australia, Saudi Arabia and Colombia, and has been commercialised in 12 overseas countries or regions, including South Korea, the United States, Brazil, Poland, Spain, Portugal, Chile, Ireland, the United Kingdom, Croatia, Greece and Argentina, all receiving high praise from local clinicians. NUMEN® Coil permits stable framing, smooth filling and finishing, with superb conformability to shapes of aneurysms. Its three models, MicroFrame, MicroFill and MicroFinish, have a total of 177 specifications, providing physicians with a full range of embolization options. In June 2023, the research results of NUMEN® Coil applied to aneurysms less than 5mm were officially published in the journal "BMC Surgery", further demonstrating its safety and effectiveness of application to aneurysms less than 5mm, and the clinical effect has reached the international advanced level.

NUMEN Silk® Coil

NUMEN Silk® Coil is an iterative product developed based on NUMEN® Coil, and was approved by the NMPA in February 2022. As a new generation of ultra-soft electronically detachable coil, NUMEN Silk® Coil features a greater smoothness in the filling stage and finishing stage. The smoothness of the distal-end of its delivery wire improves the microcatheter's stability, to minimize the chance of the kick-back of the microcatheter in the finishing stage, therefore reducing the risk of aneurysm rupture.









Tubridge® Flow-diverting Stent

Tubridge® Flow-diverting Stent was the first neuro-interventional medical device that entered the Green Path, and was also the first Chinese-developed flow-diverting stent approved by the NMPA. Leveraging the principle of haemodynamics, Tubridge® Flow-diverting Stent can alter the blood flow state of the aneurysm to reduce the impact of blood flow on the aneurysm, which allows the endothelial cells to grow along the stent skeleton, gradually repairing the aneurysm neck and curing the aneurysm. Since its launching in 2018, the product has been widely recognised by surgeons in the industry by virtue of its excellent clinical effects. During the Reporting Period, the product was listed in the 2022 Shanghai Biomedical "New and Excellent Medical Devices" Product Catalogue (《2022年度上海市生物醫藥"新優藥械"產品目錄》) and passed the "Shanghai Brand" certification.



During the Reporting Period, the PARAT MINI clinical study of Tubridge® Flow-diverting Stent in the treatment of small and medium-sized and intracranial wide-neck aneurysms completed the enrollment of all cases, in order to expand the indications of this product in the treatment of small and medium-sized aneurysms. Its new-generation product, Tubridge Plus® Flow-diverting Stent ("Tubridge Plus® Flow-diverting Stent"), which aims to improve the smoothness in delivery and stent visibility under angiography, could facilitate the accurate placement of the stent and enhance the safety of procedures. As of the date of this report, the PARAT PLUS study of the pre-marketing clinical trial for this product has completed the enrollment of all cases. The two clinical studies provided a number of evidence-based medical evidences for Tubridge® series flow-diverting stent in the treatment of large and giant aneurysms, medium and small aneurysms, and real-world applications.

WILLIS® Stent Graft

WILLIS® Stent Graft is the first and the only intracranial stent graft approved for treating cerebral vessel diseases in the world. It is also the first neuro-interventional medical device that applies the theory of intracranial parent artery reconstruction in practice, with a focus on the characterised and unique treatment sector, and provides viable solutions for complex neurovascular diseases, including dissecting aneurysms, blood blister-like aneurysms, pseudo-aneurysms as well as carotid-cavernous fistulae.



Comaneci® Assist Device

Comaneci® Assist Device is an adjustable temporary coil embolization assisting stent developed by Rapid Medical. It has received CE Marking in 2014 and FDA approval in 2019, and received FDA Breakthrough Device designation in February 2022 for the treatment of cerebral vasospasm after hemorrhagic stroke. The product is used in the coil embolization of wide-neck or unusually shaped aneurysms to prevent the coil from falling and inadvertently blocking the artery. We are the exclusive distributor in Greater China for Comaneci® Assist Device.

Rebridge® Intracranial Visualized Stent ("Rebridge® Stent")

The Rebridge® Stent is the first Chinese-developed full-visualized coil embolization assisting stent to enter clinical trials. The whole body of the stent is densely braided from radiopaque alloy wires, and thus, when compared with other stents that only have several radiopaque wires, Rebridge® Stent allows physicians to position more precisely for optimal adherent effect after stent expansion.

Intracranial Atherosclerotic Stenosis Products

The Group has a comprehensive product portfolio in the field of treatment of cerebral atherosclerotic stenosis, consisting of five self-developed products, which specifically cover solutions for the three major disease segments including intracranial stenosis, vertebral artery stenosis and carotid artery stenosis. According to Frost & Sullivan, the Group's intracranial stents ranked the first in China, with the market share of over 60% in terms of the volume of procedures performed in 2022.

During the Reporting Period, the Group recorded the revenue of cerebral atherosclerotic stenosis products of approximately RMB55.8 million, representing an increase of approximately 17.1% over the Prior-year Period. The increase was mainly due to the acceleration of marketing of Bridge® Vertebral Artery DES.



APOLLO™ Intracranial Stent

APOLLO™ Intracranial Stent is a balloon-expandable stent system, and was approved by the NMPA in 2004. It is the first stent system in the world to treat intracranial atherosclerotic disease (ICAD), and was approved for marketing in Argentina in 2015. With its excellent safety and efficacy, APOLLO™ Intracranial Stent has maintained the first place in its market share for many years. In recent years, benefiting from the application of stenosis cases in emergency clot retrieval procedure in grassroots hospitals, the market demand for APOLLO™ Intracranial Stent has maintained a stable growth trend.



Bridge® Vertebral Artery DES

Bridge® Vertebral Artery DES is the first approved vertebral artery DES admitted to the Green Path. Bridge® Vertebral Artery DES is designed with single-sided grooved drug-coated stent, and the drug is accurately targeted to release, which can effectively reduce the incidence of in-stent stenosis and avoid the negative impact of drugs on the endothelialization of the stent. The results of pre-marketing clinical trials of the product showed that the success rate of Bridge® Vertebral Artery DES implantation was 98%, and the incidence of in-stent restenosis (≥50%) at 6 months after operation was only 3.7%, which fully proved its clinical safety and effectiveness. During the Reporting Period, the product was listed in the 2022 Shanghai Biomedical "New and Excellent Medical Devices" Product Catalogue (《2022年度上海市生物醫藥"新優藥械"產品目錄》).



Diveer® Balloon Catheter

Diveer® Balloon Catheter is a specialized rapid-exchange intracranial balloon catheter self-developed by the Company, which is used for the interventional treatment of non-acute symptomatic intracranial atherosclerotic stenosis. Its ultra-soft tip reduces the risk of vascular injury, and its low push resistance enables excellent placement and pushability in tortuous vessels and complex lesions. The product was approved by the NMPA in January 2022, further expanding the Group's product line for the treatment of cerebral atherosclerosis stenosis.



Acute Ischemic Stroke Products

In the field of acute ischemic stroke, the Group has four commercialized products and three products under research and development, covering stent thrombectomy devices and aspiration thrombectomy devices. According to Frost & Sullivan, the Company is the only Chinese company with stent thrombectomy devices compatible with different sizes of blood vessels.

During the Reporting Period, the Group recorded the revenue of acute ischemic stroke products of approximately RMB6.9 million, representing an increase of approximately 1,459.5% over the Prior-year Period, mainly due to the revenue growth contributed by the newly-launched Neurohawk® Thrombectomy Device and X-track® Distal Catheter in 2022.

Neurohawk® Thrombectomy Device

Neurohawk® Thrombectomy Device is the Group's self-developed stent retriever with full visualization, which was approved by the NMPA in February 2022. It features a composite mesh design consisting of two meshes with different opening sizes arranged in a staggered spiral pattern, which allows it to better capture large, tough or fragile clots and improves its wall apposition.



X-track® Distal Access Catheter

X-track® Distal Access Catheter is an intermediate catheter product developed by the Group for treating acute ischemic stroke, which was approved by the NMPA in April 2022. The product adopts special polymer material and double-wire braided structure, which can reach the lesion site for multiple times during the operation, and its good anti-fatigue performance can fully address the clinical needs for catheter improvement.





Tigertriever® Revascularization Stent

Tigertriever® Revascularization Stent is the world's first adjustable stent retriever with full visualization, indicated for procedures performed in blood vessels of varying diameters. The product obtained the CE Marking in the European Union in May 2018, obtained the FDA approval in the United States in March 2021 and was approved by the NMPA in August 2023. We were engaged by Rapid Medical as the exclusive distributor in Greater China for Tigertriever® Revascularization Stent, Tigertriever® 13 Stent and all iterations of Tigertriever®. Tigertriever® Stent was admitted into the NMPA's Green Path in May 2020. Tigertriever® 13 Stent is the smallest stent embolectomy device for the treatment of distal vascular occlusion in the world, which was approved by the FDA in July 2022.



W-track® Aspiration Catheter

W-track® Aspiration Catheter is an intracranial aspiration catheter used for clot aspiration. It has a multi-segment transition design to allow its smooth delivery, and its double-wire braided structure with stainless steel enhances the elongation resistance of the catheter while maintaining flexibility. W-track® Aspiration Catheter can reach the target occlusion quickly and smoothly, in particular in tortuous intracranial vessels. The product was approved by the NMPA in August 2023.



Balloon Guide Catheter

Balloon Guide Catheter is a large lumen catheter with a compliant balloon at the distal tip of the catheter, which is designated to facilitate the insertion and guidance of an intravascular catheter while causing temporary distal flow arrest in the artery. We have submitted the registration application for this product to the NMPA.

Access Products

The Group has a product portfolio of seven auxiliary access devices, among which the five commercialized products include Asahi® Neurovascular Guidewires ("**Asahi® Guidewires**"), U-track® Intracranial Support Catheter System ("**U-track® Support Catheter**"), Fastrack® Microcatheter System, Q-track® Microcatheter and Veyronwire™ Guide Wire, and the products under research and development include various models of microcatheter products and distal protection device products.

During the Reporting Period, the Group recorded the revenue of access products of approximately RMB28.6 million, representing a decrease of approximately 27.7% over the Prior-year Period, which was because the production and logistics of self-owned products were affected by the public health event during the Prior-year Periods, leading to the proportion of distributed products in the sales portfolio increased significantly.

Asahi® Guidewires

Asahi® Guidewires are one of the global leading neurovascular guidewires. Asahi® Guidewires feature a unique multistranded coil design at the tip, enhancing torque response, elongation resistance and flexibility. The product was approved by the NMPA in August 2013. The Group has been engaged by Asahi Intecc as the exclusive distributor of Asahi® Guidewires in China since 2016.

Fastrack® Microcatheter

Fastrack® Microcatheter is currently the only microcatheter system with a lumen of 0.029" in China. Its unique large lumen can provide the simplicity of instrument delivery and recovery. The product is designed to reach farther lesions in neurovascular surgery, and support the precise delivery of intracranial interventional devices. The product was approved by the NMPA in July 2019.



U-track® Support Catheter

U-track® Support Catheter can reach remote lesions in neurovascular surgery and support the precise delivery of various neurovascular interventional devices. The product was approved by the NMPA in December 2020 and was approved for marketing in Brazil in September 2022. During the Reporting Period, the first batch of commercial use of this product was completed in Brazil. It was the Company's fourth product entering the Brazilian market and the first access product, which enriched the Company's product portfolio for cerebrovascular diseases in Brazil.



Q-track® Microcatheter

Q-track® Microcatheter was approved by the NMPA in June 2023. The product adopts a non-invasive head end, specially treated transition section design and hydrophilic coating lubrication, which can reach the deep blood vessels of the brain and avoid the stimulation of blood vessels as much as possible. The product has an effective length of 155cm and is compatible with various surgical procedures to meet the needs of different scenarios. In particular, it can effectively remove thrombus when using in conjunction with the Neurohawk® Thrombectomy Device during the treatment of acute ischemic stroke.





RESEARCH AND DEVELOPMENT

The Group has always adhered to the purpose of addressing clinical needs and continued on innovation. After years of accumulation, we have mastered the core design and manufacturing technology platforms for the R&D and manufacturing of neuro-interventional medical devices, including braiding and coiling technology, stent forming and processing technology, balloon technology and catheter technology. We have also established a core R&D team with significant technical expertise in these fields. As of 30 June 2023, the Group had a total of 165 R&D personnel, more than 55% of which have master's degrees or above.

The Group has established a mature project evaluation mechanism to regularly track the development direction of cutting-edge technology in the industry, evaluate market demand and its own technology reserves, so as to provide a foundation for medium-and long-term product development strategy. Through a mature physician-engineer collaboration system, we actively understand the clinical needs of physicians and patients, conduct in-depth exploration of clinical pain points, and regularly evaluate new technologies under development to ensure our products meet clinical needs.

Intellectual Property Rights

The Group insists on R&D and innovation with proprietary intellectual property rights. As of 30 June 2023, the Group had 196 authorized patents, including 44 overseas patents. During the Reporting Period, 18 authorized patents were newly granted, including 1 overseas patent. In addition, the Group has 295 patents under application. With the strategies of branding, marketing and compliance protection, we have completed the layout of domestic and foreign trademarks with 176 registered trademarks and submitted 9 applications for new trademarks during the Reporting Period.

QUALITY MANAGEMENT AND MANUFACTURING

The Group upholds the product quality as its core value. We have established a digital product quality control system covering the entire production process, allowing us to trace the whole life cycle of product design, development, manufacturing and after-sale service. During the Reporting Period, the Group obtained the MDSAP (Medical Device Single Audit Program), a quality system certification accepted in five countries, which effectively reduces the audit cost for products entering overseas markets. As of the end of the Reporting Period, the Group has successively obtained a number of system certifications, covering the relevant regulations and standard requirements of China, the European Union, the United States, Australia, Brazil, Japan, South Korea, Argentina and other countries around the world, forming an international quality management system.

The Group summed up the "Practical Experience on the 'One Core and Three Links (一核三環)' Quality and Health Management Model" in the long-term exploration and practice based on the comprehensive quality management, and was awarded with the honorary title of "Shanghai Quality Benchmark" in July 2023 by virtue of this project. Through the implementation of the "'One Core and Three Links' Quality and Health Management Model", the Group has formed a relatively complete management model in terms of improving product quality, reducing quality costs and improving system operation capabilities, which is conducive to establishing clearer quality evaluation standards, formulating clearer quality management strategies, and promoting our high-quality development.

During the Reporting Period, we further accelerated the localization of raw materials and had completed 80 supply chain improvement and upgrading projects, in order to improve the stability of the supply chain, and significantly optimize production costs. As of the end of the Reporting Period, the localization rate of raw materials for our products had reached over 90%. At the same time, we had established an advanced quality management system, continuously strengthened the construction of the lean system, steadily improved the production yield and the production efficiency, and realized the cost reduction and consumption control.

HUMAN RESOURCES

After a decade of development, the Group has built the largest neuro-interventional industrialization team in China, with a full-cycle operational capabilities in the neuro-interventional medical device industry covering R&D, clinical trials and registration, supply chain management and commercialization. As of 30 June 2023, the Group had a total of 575 employees, more than 50% of which had bachelor's degrees or above.

PROSPECT

Considering the aging population, the increasing number of stroke patients and the improvement of medical infrastructures, the neuro-interventional medical device industry in China is faced with huge development opportunities. In order to seize such opportunities and enhance core competitiveness amidst the market competition, the Group will make full use of its first-mover and scale advantages and implement active business strategies, including but not limited to the following:

1. Continue to enhance innovation capabilities to offer a comprehensive solution for cerebral vessel diseases

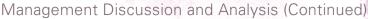
We will continue to expand the depth and breadth of our product portfolio to achieve full product coverage of the cerebrovascular therapeutic area. Through independent development and external cooperation, we will continue with development, innovation and iteration, aligning every step of product improvement with clinical needs to offer stroke patients with a top-quality total solution.

2. Promote the universal and affordable strategy and improve operating efficiency

We will continue to optimize our operating system and quality control system in an all-round way, upgrade our manufacturing technologies, strengthen our training system, and build a global supply chain system to further reduce costs and improve operating efficiency. In addition, we plan to increase our production capacity by expanding our production facilities and teams. Taking advantage of the economies of scale, we will promote universal and affordable neuro-interventional solutions, thereby increasing the level of stroke disease diagnosis and treatment in grassroots medical institutions, and benefiting more patients.

3. Expand the strategic global layout

We will actively expand our global presence and gradually enter the top ten countries and regions in terms of the volume of neuro-interventional procedures. We plan to advance the registration of our innovative products overseas and expand our international team to provide physicians and patients from all over the world with advanced therapeutic products and treatment options. We also plan to establish overseas R&D and production centers to expand our brand visibility globally and attract talents and resources in the neuro-interventional field worldwide. In addition, we will continue to have in-depth cooperation with leading international companies to enlarge our product portfolio and sales network, so as to build an international innovation platform.





FINANCIAL REVIEW

Revenue

During the Reporting Period, the Group's revenue was mainly derived from hemorrhagic stroke products, cerebral atherosclerotic stenosis products, acute ischemic stroke products and access products. The Group's revenue increased by approximately 45.2% from approximately RMB206.0 million for the six months ended 30 June 2022 to approximately RMB299.2 million for the six months ended 30 June 2023.

The increase was mainly because: (1) market-share leading products (including Tubridge® Flow-diverting Stent, etc.) continued to increase the market penetration, further consolidating the competitive advantages, and maintaining a positive growth momentum; (2) the continued tendering and admission of several new products launched by the Group in recent years (including NUMEN® Coil, Bridge® Vertebral Artery DES and U-track® Support Catheter, etc.) into hospitals has contributed to the exploration of blank markets; and (3) multiple newly approved products in 2022 (including Neurohawk® Thrombectomy Device, Diveer® Balloon Catheter, etc.) accelerated the market development, contributing to the Group's revenue growth.

Set out below is the breakdown of revenue by product category:

For the six months ended 30 June

	2023 RMB'000 (unaudited)	2022 RMB'000 (unaudited)	Change %	
Hemorrhagic stroke products	206,837	117,505	76.0%	
Cerebral atherosclerotic stenosis products	55,827	47,677	17.1%	
Acute ischemic stroke products	6,924	444	1,459.5%	
Access products	28,597	39,563	-27.7%	
Other business revenue	1,008	804	25.4%	
Total	299,193	205,993	45.2%	

Cost of Sales

Our cost of sales increased by approximately 3.3% from approximately RMB64.4 million for the six months ended 30 June 2022 to approximately RMB66.6 million for the six months ended 30 June 2023. Such increase was primarily due to an increase in sales volume of various types of products mentioned above.

Gross Profit and Gross Profit Margin

Our gross profit increased by approximately 64.4% from approximately RMB141.5 million for the six months ended 30 June 2022 to approximately RMB232.6 million for the six months ended 30 June 2023. The increase was primarily due to an increase in sales volume of various types of products mentioned above.

The Group's gross profit margin was approximately 77.8% during the Reporting Period. Gross profit margin increased by 9.1 percentage points from 68.7% in the Prior-year Period, primarily due to an increase in the proportion of in-house produced products in the product sales structure, as well as the implementation of multiple supply chain improvement projects and economies of scale to reduce production costs.

Research and Development Costs

Our research and development costs increased by approximately 71.9% from approximately RMB49.2 million for the six months ended 30 June 2022 to approximately RMB84.5 million for the six months ended 30 June 2023, primarily due to the increase in the investment in existing and newly launched R&D projects.

Distribution Costs

Our distribution costs increased by approximately 65.9% from approximately RMB33.7 million for the six months ended 30 June 2022 to approximately RMB55.9 million for the six months ended 30 June 2023, primarily due to the resumption of distribution activities in the Chinese market in the first half of 2023, and an increase in the investment in overseas business distribution as compared with the Prior-year Period.

Administrative Expenses

Our administrative expenses decreased by approximately 11.2% from approximately RMB31.7 million for the six months ended 30 June 2022 to approximately RMB28.2 million for the six months ended 30 June 2023, primarily due to the improvement of efficiency on operating management, as well as the transfer of expenses to other operating departments as the change of premises' usage.

Other Net Income

Our other net income increased by approximately 276.0% from approximately RMB4.8 million for the six months ended 30 June 2022 to approximately RMB18.2 million for the six months ended 30 June 2023, primarily due to: (1) an increase of approximately RMB6.2 million in gains from government grants compared to the Prior-year Period; (2) an increase of approximately RMB5.2 million in interest income compared to the Prior-year Period; and (3) an increase in gains on fair value changes in financial assets of approximately RMB2.1 million.

Other Operating Costs

During the Reporting Period, other operating costs were nil, as compared to approximately RMB18.2 million in the Prior-year Period, primarily due to related listing expenses incurred in the Prior-year Period, and there were no such expenses for the six months ended 30 June 2023.



Our finance costs decreased by approximately 97.8% from approximately RMB89.5 million for the six months ended 30 June 2022 to approximately RMB2.0 million for the six months ended 30 June 2023, primarily due to: the interest of approximately RMB87.0 million on other financial liabilities as a result of preferred shares issued under the series A financing in the Prior-year Period, such interest expense required no payment in cash and no further accrued from the Listing Date of the Group, and there was no such interest expense for the six months ended 30 June 2023.

Share of the Losses of an Associate

During the Reporting Period, the Group's share of the losses of an associate came from Rapid Medical. The Group began to treat Rapid Medical as an associate under equity method from an accounting perspective since May 2021.

Share of the losses of an associate decreased by approximately 7.1% from approximately RMB12.8 million for the six months ended 30 June 2022 to approximately RMB11.9 million for the six months ended 30 June 2023.

Income Tax Expenses

Our income tax expenses increased by approximately 106.1% from approximately RMB5.0 million for the six months ended 30 June 2022 to approximately RMB10.3 million for the six months ended 30 June 2023, primarily due to an increase in profit from operations.

Inventories

Our inventories consist of: (1) raw materials used in production and research and development; (2) work in progress; and (3) finished goods.

Our inventory increased from approximately RMB114.7 million as of 31 December 2022 to approximately RMB173.5 million as of 30 June 2023, primarily due to an increase in reserves of raw materials and finished goods as a result of the continuous expansion of the Group's business scale.

Current Trade and Other Receivables

Our current trade and other receivables primarily consist of: (1) trade receivables; and (2) prepayments and deposits.

Our current trade and other receivables increased from approximately RMB35.3 million as of 31 December 2022 to approximately RMB57.5 million as of 30 June 2023, primarily due to an increase in trade receivables as a result of the growth of the business.

Trade and Other Payables

Our trade and other payables primarily consist of: (1) trade payables due to third-party suppliers and related parties; (2) accrued expenses; (3) accrued payroll; and (4) other payables.

Our trade and other payables increased from approximately RMB188.7 million as of 31 December 2022 to approximately RMB208.6 million as of 30 June 2023, primarily due to: (1) an increase in trade payables due to the increase in procurement of raw materials; and (2) an increase in other payables as a result of the growth of the business.

Lease Liabilities

As of 30 June 2023, the Group recorded lease liabilities of approximately RMB73.1 million, which were primarily in relation to the properties the Group leased for our office premises, manufacturing and R&D facilities. The Group recognizes lease liabilities with respect to all leases, except for short-term leases and leases of low value assets.

Capital Expenditure

The capital expenditure of the Group amounted to approximately RMB14.8 million during the Reporting Period, representing an addition of intangible assets and property, plant and equipment. In particular, the intangible assets of the Group primarily represent the capitalized development costs.

Foreign Exchange Exposure

During the Reporting Period, the Group mainly operated in China and a majority of its transactions were settled in RMB, the functional currency of the Company's primary subsidiaries. As of 30 June 2023, certain portion of the Group's bank balances was denominated in U.S. dollars. The Group currently does not have a foreign currency hedging policy. However, our management monitors foreign exchange exposure and will consider hedging significant foreign currency exposure should the need arise. Except for certain bank balances, trade receivables, trade and other payables, and other amounts denominated in foreign currencies, the Group did not have significant foreign currency exposure from its operations as of 30 June 2023.



Significant Investment

As of 30 June 2023, the Group's significant investment was an investment in an associate Rapid Medical at a cost of approximately US\$27.5 million (equivalent to RMB191.9 million). The issued and fully paid share capital of Rapid Medical is 22.1 million shares, 22.28% of which are held by the Group, and its principal business is the development, manufacture and sale of innovative devices for neuro-interventional procedures. As at 30 June 2023, the Group's interests in associates were all derived from Rapid Medical, amounting to approximately RMB148.8 million, which accounted for approximately 7.9% of the Group's total assets. For the six months ended 30 June 2023, Rapid Medical recorded a loss of approximately US\$7.5 million (equivalent to RMB52.1 million), which was mainly due to the increase in R&D and sales activities expenses of Rapid Medical, and the Group recorded a share of losses of an associate of approximately RMB11.9 million. For details, please refer to the section headed "Acquisition of certain interests in Rapid Medical" in the Prospectus. The Group have been approved to use trademarks of Rapid Medical and became the exclusive agent of Rapid Medical's related products in Greater China, and we have leveraged Rapid Medical's sales network in the United States to facilitate our overseas business. As a strategic investor, the Group will hold our investment in Rapid Medical for the long term.

Contingent Liabilities

As of 30 June 2023, the Group did not have any contingent liabilities.

Capital Management

The Group's objectives in managing capital are to safeguard the Group's ability to continue as a going concern in order to provide returns for the Shareholders and benefits for other stakeholders and to maintain an optimal capital structure to reduce the cost of capital. The Group actively and regularly reviews and manages its capital structure to maintain a balance between the higher Shareholders' returns that might be possible with higher levels of borrowings and the advantages and security afforded by a sound capital position, and makes adjustments to the capital structure in light of changes in economic conditions.

Liquidity and Financial Resources

The Group's cash and cash equivalents were approximately RMB827.5 million as of 30 June 2023, as compared to approximately RMB827.9 million as of 31 December 2022, primarily due to the net cash inflow from operating activities of approximately RMB32.4 million, net cash outflow from investing activities of approximately RMB21.8 million and net cash outflow from financing activities of approximately RMB19.8 million during the Reporting Period. The Group's policy is to regularly monitor its liquidity requirements, to ensure that it maintains sufficient reserve of cash and adequate committed lines of funding from major financial institutions to meet its liquidity requirements in the short and long term.

Borrowings and Gearing Ratio

Total borrowings of the Group, including interest-bearing borrowing as of 30 June 2023 and 31 December 2022 were nil. As of 30 June 2023, the gearing ratio of the Group (calculated as total interest-bearing borrowings and lease liabilities divided by total equity plus other financial liabilities as of the same date) decreased to 4.6%, as compared to 5.7% as of 31 December 2022.

Net Current Assets/Liabilities

As of 30 June 2023, the Group's net current assets were approximately RMB1,141.2 million, as compared to approximately RMB1,040.9 million as of 31 December 2022. Such increase was mainly attributable to the profit from operating activities during the Reporting Period.

Charge on Assets

As of 30 June 2023, there was no charge on assets of the Group.

Material Acquisitions and Disposals of Subsidiaries, Associates and Joint Ventures

During the Reporting Period, the Group did not have any material acquisitions or disposals of subsidiaries, associates and joint ventures.

Future Plans for Material Investments or Capital Assets

As of 30 June 2023, the Group did not have any plans for material investments and capital assets. The Group actively responded to external environment changes and continuously promoted business development. The Group will make further announcements in accordance with the Listing Rules, where applicable, if any investments and acquisition opportunities materialize.

CORPORATE GOVERNANCE AND OTHER INFORMATION

DIRECTORS' AND CHIEF EXECUTIVES' INTERESTS AND SHORT POSITIONS IN SHARES AND UNDERLYING SHARES

As at 30 June 2023, the interests and short positions of Directors and chief executives of the Company in the shares, underlying shares and debentures of the Company or any of its associated corporations (within the meaning of Part XV of the SFO) which were required (a) to be notified to the Company and the Stock Exchange pursuant to Divisions 7 and 8 of Part XV of the SFO (including interests and short positions which they are taken or deemed to have under such provisions of the SFO), or (b) to be and were entered in the register required to be kept by the Company pursuant to section 352 of the SFO, or (c) as otherwise notified to the Company and the Stock Exchange pursuant to the Model Code, are as follows:

(a) Long positions in the shares and underlying shares of our associated corporation:

Name of Director	Name of Associated Corporation	Nature/Capacity of Interest	Percentage of Shareholding
Mr. Peng Bo	MicroPort	Beneficial Owner (1)	0.42%
	MicroPort CardioFlow Medtech Corporation (微創心通醫療科技有限公司) (" MicroPort CardioFlow ")	Beneficial Owner (2)	<0.1%
Mr. Xie Zhiyong	Company	Beneficial Owner (3)	0.02%
	MicroPort	Beneficial Owner (4)	0.05%
Mr. Wang Yiqun Bruce	Company	Beneficial Owner (5)	0.01%
	MicroPort	Beneficial Owner (6)	0.02%

Notes:

- As at 30 June 2023, Mr. Peng Bo was interested in (i) 869,496 shares of MicroPort; and (ii) 6,841,170 underlying shares
 of MicroPort by virtue of the options granted to him under a share option scheme of MicroPort. As Pokang Limited is a
 company wholly owned by Mr. Peng Bo, Mr. Peng Bo is deemed to be interested in such shares by virtue of Part XV of
 the SFO.
- 2. As at 30 June 2023, Mr. Peng Bo was interested in 54,304 shares of MicroPort CardioFlow.
- 3. As at 30 June 2023, Mr. Xie Zhiyong was interested in 125,775 shares of the Company.
- 4. As at 30 June 2023, Mr. Xie Zhiyong was interested in (i) 538,851 shares of MicroPort; and (ii) 316,883 underlying shares of MicroPort by virtue of the options granted to him under a share option scheme of MicroPort.
- 5. As at 30 June 2023, Mr. Wang Yiqun Bruce was interested in 79,063 shares of the Company.
- 6. As at 30 June 2023, Mr. Wang Yiqun Bruce was interested in 405,620 shares of MicroPort.

Save as disclosed above, as at 30 June 2023, none of the Directors or chief executives of the Company had any interests or short positions in the shares, underlying shares and debentures of the Company and its associated corporations (within the meaning of Part XV of the SFO) which are required to be notified to the Company and the Stock Exchange pursuant to Divisions 7 and 8 of Part XV of the SFO (including interests or short positions which they were taken or deemed to have under such provisions of the SFO), or are required, pursuant to Section 352 of the SFO, to be entered in the register referred to therein, or as otherwise notified to the Company and the Stock Exchange pursuant to the Model Code.

SUBSTANTIAL SHAREHOLDERS' INTERESTS AND SHORT POSITIONS IN SHARES AND UNDERLYING SHARES

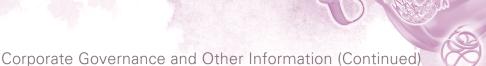
To the best knowledge of Directors, as at 30 June 2023, the following persons (other than Directors or chief executives of the Company), are directly or indirectly, interested in 5% or more of the shares or short positions in the shares and the underlying shares of the Company, which were required, pursuant to Section 336 of the SFO, to be entered in the register referred to therein:

Name of Shareholder	Nature of Interest	Number of Shares (1)	Percentage of Shareholding
MP Scientific Corporation ("MP Scientific") (2)	Beneficial Owner	310,871,340 (L)	53.35%
MicroPort (2)	Interest of controlled corporation	310,871,340 (L)	53.35%
WE'TRON Capital (3)	Beneficial Owner	60,526,500 (L)	10.39%
Maxwell Maxcare Science	Interest of controlled corporation	63,915,000 (L)	10.97%
Foundation Limited ("Maxw	vell		
Maxcare") (3)(4)			

Notes:

- 1. The letter "L" denotes a long position in our shares.
- 2. MP Scientific is directly wholly owned by MicroPort. By virtue of the SFO, MicroPort is deemed to be interested in the shares in which MP Scientific is interested.
- 3. WE'TRON Capital is directly owned as to 100.00% by Maxwell Maxcare. By virtue of the SFO, Maxwell Maxcare is deemed to be interested in the shares held by WE'TRON Capital.
- Maxwell Maxcare is also the sole shareholder of Miracle Medical Limited. Miracle Medical Limited held 3,388,500 shares, representing approximately 0.58%. By virtue of the SFO, Maxwell Maxcare is deemed to be interested in the shares held by Miracle Medical Limited.

Save as disclosed above, as at 30 June 2023, no other interests or short positions in the shares or underlying shares of the Company were recorded in the register which is required to be kept under section 336 of the SFO.



USE OF NET PROCEEDS FROM THE GLOBAL OFFERING

The Company was listed on the Main Board of the Stock Exchange on Listing Date with total net proceeds from the Listing of approximately HK\$278.1 million after deduction of the underwriting commissions, fees and other estimated expenses payable by the Company in connection with the Global Offering. The proceeds from Listing are and will continuously be used in accordance with the plans as disclosed in the section headed "Future Plans and Use of Proceeds" of the Prospectus, namely:

Use of proceeds	Approximate percentage of total amount (%)	Amount of net proceeds allocated upon Listing (HK\$ million)	Utilized amount as at 30 June 2023 (HK\$ million)	Unutilized amount as at 30 June 2023 (HK\$ million)	Expected timeline of full utilization
Research and development of therapeutic and access products for hemorrhagic stroke, cerebral atherosclerotic stenosis and acute ischemic stroke ("AIS")	30%	83.4	83.4	-	Fully utilized
Commercialization of the Company's products for hemorrhagic stroke, cerebral atherosclerotic stenosis and AIS	20%	55.6	55.6	_	Fully utilized
Expansion of the Company's manufacturing facility to increase the scale of the Company's production	15%	41.7	_	41.7	By the year ending 31 December 2023
Expansion of the Company's global presence	20%	55.6	55.6	_	Fully utilized
Advancing the Company's product portfolio through strategic acquisitions, investment, cooperation or a combination of these tactics	10%	27.8	-	27.8	By the year ending 31 December 2023
Working capital and other general corporate purposes	5%	13.9	13.9	_	Fully utilized

Save as disclosed above, since the Listing Date, the Group has not utilized any other portion of the net proceeds and will gradually utilize the remaining net proceeds in accordance with the intended purposes as stated in the Prospectus. The expected timeline is based on the best estimation of future market conditions and business operations made by the Company currently and remains subject to change based on future development of market conditions and actual business needs.

PURCHASE, SALE OR REDEMPTION OF THE COMPANY'S LISTED SECURITIES

During the six months ended 30 June 2023, save for the 517,000 shares purchased by the Trustee of the Share Award Scheme on the Stock Exchange at the total consideration of HK\$9,533,660 (equivalent to RMB8,310,000) pursuant to the terms of the trust deed under the Share Award Scheme, neither the Company nor any of its subsidiaries purchased, sold or redeemed any listed securities of the Company.

SHARE AWARD SCHEME

Reference is made to the annual report of the Company for the year ended 31 December 2022. Unless otherwise provided in the Share Award Scheme, subject to the receipt by the Trustee of within the period stipulated in the vesting notice sent to the relevant Selected Participant by the Board or the Committee, and a confirmation from the Company that all vesting conditions having been fulfilled, the Trustee shall transfer the relevant Award Shares to the Selected Participant(s) or his/her nominee(s) as soon as practicable after the Vesting Date. The Vesting Date shall be on any Business Day at the end of March of any year or any other date as stated in the Offer Letter or may be otherwise determined by the Board. No Award Shares have been granted during the year ended 31 December 2022.

During the six months ended 30 June 2023, 516,717 Award Shares have been granted and been fully vested on the same day.

Categories	Date of grant	Number of Award Share granted as at 1 January 2023	Number of Award Shares granted during the six months ended 30 June 2023
Pirantana			
Directors	0.000/0.0/0.0		405 775
Xie Zhiyong	2023/03/30	_	125,775
Wang Yiqun Bruce	2023/03/30	_	79,063
Other grantees in aggregate			
Employee	2023/03/30	_	311,879



Corporate Governance and Other Information (Continued)

SHARE SCHEME

Save as the Share Award Scheme mentioned above, as of 30 June 2023, no other share scheme was adopted by the Company.

A share scheme ("**Share Scheme**") was approved by the Company's annual general meeting on 28 June 2023. Summary of the Share Scheme was set out in the Appendix II of circular of the Company dated 2 June 2023. During the Reporting Period, there were no options granted to any grantees.

MATERIAL EVENTS AFTER THE REPORTING PERIOD

There were no material events undertaken by the Company or by the Group after the Reporting Period.

COMPLIANCE WITH CORPORATE GOVERNANE CODE

The Company aims to achieve high standards of corporate governance which are crucial to the development and safeguard the interests of the Shareholders. To accomplish this, the Company has adopted the CG Code and the associated Listing Rules after the Listing.

The Board reviewed the Company's corporate governance practices and is satisfied that the Company has complied with all applicable code provisions as set out in the CG Code during the Reporting Period.

The Company will continue to review and enhance its corporate governance practices to ensure compliance with the CG Code.

Corporate Governance and Other Information (Continued)

COMPLIANCE WITH THE MODEL CODE FOR SECURITIES TRANSACTIONS BY DIRECTORS

The Company has adopted the Model Code as its code of conduct regarding securities transactions by the Directors. Upon specific enquiry, all Directors confirmed that they had complied with the requirements as set out in the Model Code as at the date of this interim report.

CHANGES IN THE BOARD AND THE DIRECTORS' INFORMATION

There is no change in information of the Directors which are required to be disclosed pursuant to Rule 13.51B(1) of the Listing Rules since the Company's last published annual report.

CONTINUING DISCLOSURE OBLIGATION PURSUANT TO THE LISTING RULES

Save as disclosed in this interim report, the Company does not have any other disclosure obligations under Rules 13.20, 13.21 and 13.22 of the Listing Rules.

REVIEW BY AUDIT COMMITTEE

The Audit Committee consists of three independent non-executive Directors, namely Mr. Siu Chi Hung (Chairperson), Dr. Xu Yi and Dr. Zhang Haixiao.

The Audit Committee has reviewed together with the management of the Company the accounting principles and policies adopted by the Company, the interim report and the unaudited consolidated financial statements of the Group for the Reporting Period.

REVIEW BY INDEPENDENT AUDITOR

The Group's interim financial report for the Reporting Period is unaudited, but has been reviewed by the Company's independent auditor, KPMG, in accordance with Hong Kong Standard on Review Engagements 2410, "Review of Interim Financial Information performed by the Independent Auditor of the Entity" issued by the Hong Kong Institute of Certified Public Accountants.

INTERIM DIVIDEND

The Board has resolved not to pay any interim dividend for the six months ended 30 June 2023.



EMPLOYEES AND REMUNERATION POLICIES

The Group offers remuneration packages based on individuals' qualifications and experiences and generally match the market rate for salary and bonus to stay competitive in the labor market. The Group also provide extensive training programs to our employees and award incentives to encourage inventions by our R&D team. As required under the PRC regulations, the Group participate in housing fund and various employee social security plan that are organized by applicable local municipal and provincial governments.

On behalf of the Board

MicroPort NeuroTech Limited

Mr. Peng Bo

Chairperson

30 August 2023

INDEPENDENT AUDITOR'S REPORT



Review report to the board of directors of MicroPort NeuroTech Limited

(Incorporated in Cayman Islands with limited liability)

Introduction

We have reviewed the interim financial report set out on pages 39 to 64 which comprises the consolidated statement of financial position of MicroPort NeuroTech Limited (the "Company") as of 30 June 2023 and the related consolidated statement of profit or loss, statement of profit or loss and other comprehensive income and statement of changes in equity and condensed consolidated cash flow statement for the six-month period then ended and explanatory notes. The Rules Governing the Listing of Securities on The Stock Exchange of Hong Kong Limited require the preparation of an interim financial report to be in compliance with the relevant provisions thereof and Hong Kong Accounting Standard 34, *Interim financial reporting*, issued by the Hong Kong Institute of Certified Public Accountants. The directors are responsible for the preparation and presentation of the interim financial report in accordance with Hong Kong Accounting Standard 34.

Our responsibility is to form a conclusion, based on our review, on the interim financial report and to report our conclusion solely to you, as a body, in accordance with our agreed terms of engagement, and for no other purpose. We do not assume responsibility towards or accept liability to any other person for the contents of this report.

Scope of review

We conducted our review in accordance with Hong Kong Standard on Review Engagements 2410, Review of Interim Financial Information Performed by the Independent Auditor of the Entity, issued by the Hong Kong Institute of Certified Public Accountants. A review of the interim financial report consists of making enquiries, primarily of persons responsible for financial and accounting matters, and applying analytical and other review procedures. A review is substantially less in scope than an audit conducted in accordance with Hong Kong Standards on Auditing and consequently does not enable us to obtain assurance that we would become aware of all significant matters that might be identified in an audit. Accordingly we do not express an audit opinion.

Conclusion

Based on our review, nothing has come to our attention that causes us to believe that the interim financial report as at 30 June 2023 is not prepared, in all material respects, in accordance with Hong Kong Accounting Standard 34, *Interim financial reporting*.

KPMG

Certified Public Accountants

8th Floor, Prince's Building 10 Chater Road Central, Hong Kong

30 August 2023



for the six months ended 30 June 2023 — unaudited (Expressed in Renminbi)

Six months ended 30 June

	Note	2023 RMB'000	2022 RMB'000
Revenue	3	299,193	205,993
Cost of sales		(66,556)	(64,446)
Gross profit		232,637	141,547
Other net income	4	18,198	4,840
Research and development costs		(84,531)	(49,183)
Distribution costs		(55,919)	(33,710)
Administrative expenses		(28,185)	(31,749)
Other operating costs	5(b)	_	(18,163)
- m. n			
Profit from operations	F/-)	82,200	13,582
Finance costs Share of losses of an associate	5(a)	(1,963) (11,923)	(89,468) (12,839)
Stidle of losses of all associate		(11,323)	(12,039)
Profit/(loss) before taxation	5	68,314	(88,725)
Income tax	6	(10,315)	(5,004)
Profit/(loss) for the period		57,999	(93,729)
Attributable to:		04.044	(00.050)
Equity shareholders of the Company		64,041	(92,352) (1,377)
Non-controlling interests		(6,042)	(1,377)
Profit/(loss) for the period		57,999	(93,729)
Earnings/(loss) per share	7		
Basic and diluted (in RMB)	/	0.11	(0.20)

The notes on pages 47 to 64 form part of this interim financial report. Details of dividends payable to equity shareholders of the Company are set out in note 14.

CONSOLIDATED STATEMENT OF PROFIT OR LOSS AND OTHER COMPREHENSIVE INCOME

for the six months ended 30 June 2023 — unaudited (Expressed in Renminbi)

Six months ended 30 June

	2023 RMB'000	2022 RMB'000
Profit/(loss) for the period	57,999	(93,729)
Other comprehensive income for the period (after tax and reclassification adjustments):		
Items that will not be reclassified to profit or loss: Exchange differences on translation of financial statements of the Company	46,137	(22,061)
Items that may be reclassified subsequently to profit or loss: Exchange differences on translation of financial statements of foreign subsidiaries	(20,858)	(23,905)
Other comprehensive income for the period	25,279	(45,966)
Total comprehensive income for the period	83,278	(139,695)
Attributable to:		
Equity shareholders of the Company Non-controlling interests	89,320 (6,042)	(138,318) (1,377)
Total comprehensive income for the period	83,278	(139,695)

The notes on pages 47 to 64 form part of this interim financial report.



CONSOLIDATED STATEMENT OF FINANCIAL POSITION

at 30 June 2023 — unaudited (Expressed in Renminbi)

		At 30 June	At 31 December
	Note	2023 RMB'000	2022 RMB'000
	TVOLE	TUVID 000	THVID 000
Non-current assets			
Property, plant and equipment	8	176,236	193,566
Investment property		13,097	13,268
		189,333	206,834
Intangible assets	8	130,486	131,650
Interest in an associate	O	148,820	155,501
Deferred tax assets		12,896	11,642
Other non-current assets	9	27,013	26,688
		508,548	E22 21E
Current assets		500,540	532,315
Financial assets measured at fair value through profit or loss	15	266,444	266,053
Inventories		173,473	114,726
Trade and other receivables	10	57,461	35,256
Time deposits	11	60,079	40,721
Cash and cash equivalents	11	827,466	827,929
		1,384,923	1,284,685
Current liabilities			
Trade and other payables	12	208,567	188,703
Contract liabilities		7,589	11,632
Lease liabilities		24,077	24,725
Derivative financial instruments		_	272
Income tax payables		3,525	18,468
		243,758	243,800

Consolidated Statement of Financial Position (Continued) at 30 June 2023 — unaudited (Expressed in Renminbi)

Note	At 30 June 2023 RMB'000	At 31 December 2022 RMB'000
Net current assets	1,141,165	1,040,885
Total assets less current liabilities	1,649,713	1,573,200
Non-current liabilities Lease liabilities Deferred income Other non-current liabilities	49,031 19,556 8,329	60,519 19,136 7,894
	76,916	87,549
NET ASSETS	1,572,797	1,485,651
CAPITAL AND RESERVES 14 Share capital Reserves	76 1,565,050	76 1,472,727
Total equity attributable to equity shareholders of the Company Non-controlling interests	1,565,126 7,671	1,472,803 12,848
TOTAL EQUITY	1,572,797	1,485,651

Approved and authorised for issue by the board of directors.

Peng BoXie ZhiyongDirectorDirector

Date: 30 August 2023

The notes on pages 47 to 64 form part of this interim financial report.



CONSOLIDATED STATEMENT OF CHANGES IN EQUITY

for the six months ended 30 June 2023 — unaudited (Expressed in Renminbi)

			Attributable to equity shareholders of the Company							
	Note	Share capital RMB'000	Share premium RMB'000	Exchange reserve RMB'000	Capital reserve RMB'000	Statutory general reserve RMB'000	Retained profits RMB'000	Total RMB'000	Non- controlling interests RMB'000	Total (deficit)/ equity RMB'000
Balance at 1 January 2022		60	_	984	(320,594)	24,581	120,029	(174,940)	_	(174,940)
Changes in equity for the six months ended 30 June 2022: Loss for the period		_	_	_	_	_	(92,352)	(92,352)	(1,377)	(93,729)
Other comprehensive income		_	_	(45,966)	_	_	_	(45,966)	_	(45,966)
Total comprehensive income				(45,966)			(92,352)	(138,318)	(1,377)	(139,695)
Capital contributions from non-controlling interests		_	_	_	1,049	_	_	1,049	15,761	16,810
Equity-settled share-based transactions	14	_	_	_	6,274	_	_	6,274	_	6,274
Balance at 30 June 2022 and 1 July 2022		60	_	(44,982)	(313,271)	24,581	27,677	(305,935)	14,384	(291,551)

Consolidated Statement of Changes in Equity (Continued) for the six months ended 30 June 2023 — unaudited

(Expressed in Renminbi)

		Attributable to equity shareholders of the Company						_	
No	Share capital te RMB'000	premium	Exchange reserve RMB'000	Capital reserve RMB'000	Statutory general reserve RMB'000	Retained profits RMB'000	Total RMB'000	Non- controlling interests RMB'000	Total (deficit)/ equity RMB'000
Balance at 30 June 2022 and 1 July 2022	60	_	(44,982)	(313,271)	24,581	27,677	(305,935)	14,384	(291,551)
Changes in equity for the six months ended 31 December 2022: Profit for the period Other comprehensive income	_	_ _	— 34,191	_	_	70,587 —	70,587 34,191	(1,536)	69,051 34,191
Total comprehensive income	_	_	34,191	_		70,587	104,778	(1,536)	103,242
Issuance of ordinary shares Conversion of preferred shares into	2	276,138	_	_	_	_	276,140	_	276,140
ordinary shares 1 Appropriation of statutory general reserve Equity-settled share-based transactions	3 14 — —	1,101,653 — —	- - -	290,286 — 5,867	14,229 —	— (14,229) —	1,391,953 — 5,867	- - -	1,391,953 — 5,867
Balance at 31 December 2022	76	1,377,791	(10,791)	(17,118)	38,810	84,035	1,472,803	12,848	1,485,651



Attributable to equity shareholders of the Company Non-Statutory Share Share Exchange Capital general Retained controlling Total capital premium reserve reserve reserve profits Total interests equity RMB'000 RMB'000 RMB'000 RMB'000 RMB'000 RMB'000 RMB'000 RMB'000 RMB'000 Balance at 1 January 2023 1,377,791 (10,791)(17,118)38,810 84,035 1,472,803 12,848 1,485,651 Changes in equity for the six months ended 30 June 2023: Profit for the period 64,041 64,041 (6,042)57,999 Other comprehensive income 25,279 25,279 25,279 Total comprehensive income 25,279 64,041 89,320 (6,042)83,278 Capital contributions from non-controlling interests 1,665 1,665 865 2,530 Repurchase of shares under share award scheme 14(b) (8,310)(8,310)(8,310)Share granted under share award scheme 14(f) 6.652 6.652 6.652 2,996 Equity-settled share-based transactions 14 2,996 2,996 Balance at 30 June 2023 7,671 1,572,797 76 1,377,791 14,488 (14,115)38,810 148,076 1,565,126

The notes on pages 47 to 64 form part of this interim financial report.

CONDENSED CONSOLIDATED CASH FLOW STATEMENT

for the six months ended 30 June 2023 — unaudited (Expressed in Renminbi)

Six months ended 30 June

	2023 RMB'000	2022 RMB'000
Operating activities		45 700
Cash generated from operations	58,870	45,720
Tax paid	(26,513)	(628)
Net cash generated from operating activities	32,357	45,092
Investing activities		
Payments for the purchase of property, plant and equipment	(8,544)	(19,338)
Payments for intangible assets, including expenditures on capitalised	(0,344)	(10,000)
development costs	(6,305)	(11,160)
Placement of time deposits	(20,000)	(50,000)
Payment for purchase of financial assets at fair value through profit and loss	(57,067)	(50,000)
Proceeds from disposal of property, plant and equipment	(57,067)	_
Interest received	1,399	
Proceeds from redemption of financial assets at fair value through profit and	1,333	_
loss	68,660	_
Net cash used in investing activities	(21,790)	(80,498)
Financing activities		
Capital element of lease rentals paid	(12,136)	(13,208)
Interest element of lease rentals paid	(1,858)	(2,374)
Payment for repurchase of shares under share award scheme	(8,310)	(2,374)
Capital contribution from shareholders	(0,310)	66,669
Capital contributions from non-controlling interests	2,530	16,810
Capital Contributions from non-controlling interests	2,330	10,010
Net cash (used in)/generated from financing activities	(19,774)	67,897
Net (decrease)/increase in cash and cash equivalents	(9,207)	32,491
Cash and cash equivalents at 1 January	827,929	593,287
Effect of foreign exchanges rates changes	8,744	14,224
Cash and cash equivalents at 30 June	827,466	640,002

The notes on pages 47 to 64 form part of this interim financial report.



(Expressed in Renminbi unless otherwise indicated)

1 Basis of preparation

MicroPort NeuroTech Limited (the "Company") was incorporated in the Cayman Islands on 30 September 2020 as an exempted company with limited liability under the Companies Act (As Revised) of the Cayman Islands. The Company and its subsidiaries (together, "the Group") are principally engaged in the research and development, manufacturing and sale of neuro-interventional medical devices. The Company has not carried out any business since the date of its incorporation save for the Group reorganisation below.

During the six months ended 30 June 2023 and 2022, the Group's business was primarily conducted through MicroPort NeuroTech (Shanghai) Co., Ltd. ("**MP NeuroTech Shanghai**") (微創神通醫療科技(上海)有限公司). As part of the Group restructuring (the "**Restructuring**"), the Group obtained control of MP NeuroTech Shanghai in 2022.

This interim financial report has been prepared in accordance with the applicable disclosure provisions of the Rules Governing the Listing of Securities on The Stock Exchange of Hong Kong Limited, including compliance with Hong Kong Accounting Standard ("**HKAS**") 34, *Interim financial reporting*, issued by the Hong Kong Institute of Certified Public Accountants ("**HKICPA**"). It has been reviewed by the audit committee of the Company and was authorised for issue on 30 August 2023.

The interim financial report has been prepared in accordance with the same accounting policies adopted in the 2022 annual financial statements, except for the accounting policy changes that are expected to be reflected in the 2023 annual financial statements. Details of any changes in accounting policies are set out in Note 2.

The preparation of an interim financial report in conformity with HKAS 34 requires management to make judgments, estimates and assumptions that affect the application of policies and reported amounts of assets and liabilities, income and expenses on a year-to-date basis. Actual results may differ from these estimates.

The interim financial report contains condensed consolidated financial statements and selected explanatory notes. The notes include an explanation of events and transactions that are significant to an understanding of the changes in financial position and performance of the Group since the 2022 annual financial statements. The condensed consolidated interim financial statements and notes thereon do not include all of the information required for a full set of financial statements prepared in accordance with Hong Kong Financial Reporting Standards ("HKFRSs").

This interim financial report is unaudited, but has been reviewed by KPMG in accordance with Hong Kong Standard on Review Engagements 2410, *Review of Interim Financial Information Performed by the Independent Auditor of the Entity*, issued by the HKICPA. KPMG's independent review report to the Board of Directors of the Company is included on page 38.

The financial information relating to the financial year ended 31 December 2022 that is included in the interim financial report as comparative information does not constitute the Company's annual consolidated financial statements for that financial year but is derived from those financial statements. The Company's annual consolidated financial statements for the year ended 31 December 2022 are available from the Company's registered office. The auditors have expressed an unqualified opinion on those financial statements in their report dated 29 March 2023.

(Expressed in Renminbi unless otherwise indicated)

2 Changes in accounting policies

The HKICPA has issued the following new and amended HKFRSs and guidance that are first effective for the current accounting period of the Group. Of these, the following developments are relevant to the Group's financial statements:

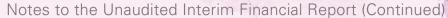
- HKFRS 17, Insurance contracts
- Amendments to HKAS 8, Accounting policies, changes in accounting estimates and errors: Definition of accounting estimates
- Amendments to HKAS 12, Income taxes: Deferred tax related to assets and liabilities arising from a single transaction
- Amendments to HKAS 12, Income taxes: International tax reform Pillar Two model rules

None of these developments have had a material effect on how the Group's results and financial position for the current or prior periods have been prepared or presented in this interim financial report. The Group has not applied any new standard or interpretation that is not yet effective for the current accounting period.

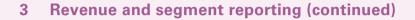
3 Revenue and segment reporting

The Group sells medical devices through appointed distributors.

For the purpose of resources allocation and performance assessment, the Group's management focuses on the operating results of the Group as a whole. As such, the Group's resources are integrated and no discrete operating segment information is available. Accordingly, no operating segment information is presented.



(Expressed in Renminbi unless otherwise indicated)



(a) Disaggregation of revenue

Disaggregation of revenue from contracts with customers by major products or service lines and geographical location of customers is as follows:

	Six months e	nded 30 June
	2023 RMB'000	2022 RMB'000
Revenue from contracts with customers within the scope of HKFRS 15		
Sales of medical devices — point in time	298,185	205,189
Revenue from other sources Gross rentals	1,008	804
	299,193	205,993
Disaggregated by geographical location of customers — the PRC — Outside the PRC	284,158 15,035	194,181 11,812
	299,193	205,993

The geographical analysis above includes property rental income in the PRC for the six months ended 30 June 2023 of RMB1,008,000 (six months ended 30 June 2022: RMB804,000).

4 Other net income

Six	months	ended	30 J	lune

	2023 RMB'000	<i>2022</i> RMB'000
Fair value changes in financial instruments measured at fair value Government grants Interest income on financial assets carried at amortised cost Net loss on disposal of property, plant and equipment Others	2,121 7,731 8,318 – 28	— 1,493 3,124 (31) 254
	18,198	4,840

(Expressed in Renminbi unless otherwise indicated)

5 Profit/(loss) before taxation

Profit/(loss) before taxation is arrived at after charging/(crediting):

(a) Finance costs

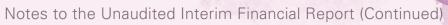
Six months ended 30 June

	2023 RMB'000	2022 RMB'000
Interest on other financial liabilities (Note 13)	_	87,032
Interest on lease liabilities	1,858	2,374
Total interest expenses on financial liabilities not at fair value		
through profit or loss	1,858	89,406
Others	105	62
	1,963	89,468

(b) Other operating costs

Six months ended 30 June

	2023 RMB′000	2022 RMB'000
Listing expenses Donations	Ξ	16,344 1,819
	_	18,163



(Expressed in Renminbi unless otherwise indicated)

5 Profit/(loss) before taxation (continued)

(c) Other items

Six months ended 30 June

	2023 RMB'000	2022 RMB'000
Amortisation of intangible assets	7,697	6,867
Depreciation charge — owned property, plant and equipment and investment	9.000	7 104
property — right-of-use assets	8,909 12,514	7,184 13,938
	21,423	21,122
Less: Capitalised into intangible assets	(817)	(250)
	20,606	20,872
Research and development expenditure	90,409	62,550
Less: Development costs capitalised into intangible assets	(5,878)	(13,367)
	84,531	49,183
(Reversal)/provision of inventories write-down	(176)	231

6 Income tax

Six months ended 30 June

	2023 RMB'000	2022 RMB'000
Current tax — PRC Corporate Income Tax ("CIT") Provision for the period	11,569	6,126
Deferred tax Origination and reversal of temporary differences	(1,254)	(1,122)
	10,315	5,004

(Expressed in Renminbi unless otherwise indicated)

6 Income tax (continued)

Pursuant to the CIT Law of the PRC, all of the Company's PRC subsidiaries are liable to PRC CIT at a rate of 25%, except for MP NeuroTech Shanghai, which is entitled to a preferential income tax rate of 15% as it is certified as a "High and New Technology Enterprise" ("**HNTE**") during the six months ended 30 June 2023 and 2022. According to Guoshuihan 2009 No. 203, if an entity is certified as an HNTE, it is entitled to a preferential income tax rate of 15% during the certified period.

Taxation for overseas subsidiaries is similarly calculated using the estimated annual effective rates of taxation that are expected to be applicable in the relevant countries.

Certain countries which the Group operates in, recently enacted or plan to enact new tax laws to implement the Pillar Two model rules with reference to the framework published by the Organisation of Economic Co-operation and Development ("**OECD**"). The new tax laws will take effect after 1 January 2024. When these laws take effect, the Group expects to be subject to a system of top-up taxes adjustments that results in the total amount of taxes payable on excess profit in each jurisdiction representing at least the minimum rate of 15%. As the new tax laws are not yet effective, the Group does not expect any current tax impact for the year ending 31 December 2023 (2022: nil). The Group has applied the temporary mandatory exception from deferred tax accounting for the top-up tax and would account for the tax as current tax when incurred.

7 Earnings/(Loss) per share

(a) Basic earnings/(loss) per share

The calculation of basic earnings/(loss) per share is based on the profit attributable to equity shareholders of the Company of RMB64,041,000 for the six months ended 30 June 2023 (loss attributable to equity shareholders of the Company of RMB92,352,000 for the six months ended 30 June 2022) and the weighted average of 582,658,100 ordinary shares (six months ended 30 June 2022: 461,397,840 shares).

(b) Diluted earnings/(loss) per share

The calculation of diluted earnings/(loss) per share amounts for the six months ended 30 June 2022 had not included the convertible bonds issued and the preferred shares issued by the Company, as they had an anti-dilutive effect on the basic earnings/(loss) per share amounts.

8 Property, plant and equipment and intangible assets

During the six months ended 30 June 2023, the Group acquired items of property, plant and equipment with a cost of RMB7,474,000 (six months ended 30 June 2022: RMB13,094,000) and capitalised development costs of RMB5,878,000 (six months ended 30 June 2022: RMB13,367,000).



(Expressed in Renminbi unless otherwise indicated)

9 Other non-current assets

	30 June 2023 RMB'000	31 December 2022 RMB'000
Lease deposits Prepayments for property, plant and equipment Others	23,445 2,700 868	23,555 2,723 410
	27,013	26,688

10 Trade and other receivables

As of the end of the reporting period, the ageing analysis of trade debtors (which are included in trade and other receivables), based on the invoice date and net of allowance for doubtful debts, is as follows:

	30 June 2023 RMB'000	31 December 2022 RMB'000
Within 1 month	28,056	5,622
1 to 3 months	1,990	4,155
3 to 12 months	1,203	294
Over 12 months	271	_
	31,520	10,071
Other debtors	4,806	3,283
Deposits and prepayments	21,135	21,902
	57,461	35,256

Trade receivables are generally due within 90 days from the date of billing.

(Expressed in Renminbi unless otherwise indicated)

11 Cash and cash equivalents and time deposits

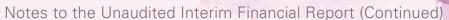
	30 June 2023 RMB'000	31 December 2022 RMB'000
Time deposits Time deposits with original terms over 3 months	60,079	40,721
Cash and cash equivalents Deposits with banks	827,466	827,929

As of the end of the reporting period, cash and cash equivalents and time deposits of the Group situated in the PRC amounted to RMB592,574,000 (31 December 2022: RMB539,663,000). Remittance of funds out of the PRC is subject to the relevant rules and regulations of foreign exchange control.

12 Trade and other payables

As of the end of the reporting period, the ageing analysis of trade payables (which are included in trade and other payables), based on the invoice date, is as follows:

	30 June 2023 RMB'000	31 December 2022 RMB'000
Within 1 month	57,114	35,093
Over 1 month but within 3 months	1,546	2,560
Over 3 months but within 6 months	3,540	368
Over 6 months but within 1 year	2,965	1,306
Over 1 year	972	889
Trade payables	66,137	40,216
Accrued expenses	19,816	22,583
Accrued payroll	30,674	42,333
Other payables	91,940	83,571
	208,567	188,703



(Expressed in Renminbi unless otherwise indicated)

13 Other financial liabilities

In November 2021, the Company and several investors (the "2021 Pre-IPO Investors") entered into a share subscription and purchase agreement, pursuant to which: (i) the 2021 Pre-IPO Investors subscribed for an aggregate of 2,032,495 newly issued series A-2 preferred shares of the Company (the "Series A-2 Preferred Shares") at an aggregate consideration of approximately US\$31.26 million; and (ii) MicroPort Scientific Investment LTD ("MP Scientific", the immediate parent of the Company) transferred 7,720,432 ordinary shares of the Company to the 2021 Pre-IPO Investors at a consideration of approximately US\$118.74 million, whereby the transferred ordinary shares were reclassified and redesignated as Series A-2 Preferred Shares (together the "Pre-IPO Investment").

Upon the completion of the Pre-IPO Investment in November 2021, the convertible bonds issued by the Company were simultaneously exchanged into an aggregate of 11,759,125 series A-1 preferred shares of the Company (the "Series A-1 Preferred Shares") at a price of approximately US\$5.95 per Series A-1 Preferred Shares.

Upon the completion of the Listing in July 2022, 95,625 series A-1 Preferred Shares and 48,764,635 series A-2 Preferred Shares issued by the Group were automatically converted into 107,560,260 ordinary shares of the Company in aggregate, resulting in a transfer of the carrying amount of other financial liabilities of RMB1,408,788,000 to ordinary share capital of RMB14,000, share premium of RMB1,101,653,000, capital reserve of RMB290,286,000 and exchange reserve of RMB16,835,000 (included in other comprehensive income), respectively.

(Expressed in Renminbi unless otherwise indicated)

14 Capital and reserves

(a) Dividends

The directors of the Company did not propose the payment of any dividend during the six months ended 30 June 2023 (six months ended 30 June 2022: nil).

(b) Purchase of own shares

During the six months ended 30 June 2023, the Company purchased its own ordinary shares through the designated trustees under the share award scheme (Note 14(f)) as follows:

Month/year	No. of shares repurchased	Price paid per s Highest HKD	hare Lowest HKD	considerations paid RMB'000
January and April 2023	517,000	20.20	13.72	8,310

Repurchased shares held at the end of reporting period under the share award scheme were classified as treasury shares and presented as a decrease in the capital reserve.

(Expressed in Renminbi unless otherwise indicated)

14 Capital and reserves (continued)

(c) Share options granted by the ultimate controlling party

MicroPort Scientific Corporation ("MPSC"), the ultimate controlling party of the Group, has granted certain share options to the employee of the Group. Each option gives the holder the right to subscribe for one ordinary share of MPSC, while the Group did not have an obligation to settle such transaction.

Apart from the outstanding share options carried forward from 2022, during the six months ended 30 June 2023, MPSC granted 75,496 share options to the employees of the Group (2,470,920 share options were granted during the six months ended 30 June 2022). These share options granted in March 2023 will vest in two instalments, 50% of total share options will vest on 31 March 2025 and the other 50% will vest on last date of each month over an explicit vesting period from April 2025 to March 2027.

During the six months ended 30 June 2023, 106,277 share options were exercised (six months ended 30 June 2022: 63,389) with a weighted average exercise price of HK\$8.90 (equivalent to approximately RMB7.52) (six months ended 30 June 2022: HK\$4.41 (equivalent to approximately RMB3.66)).

(d) Share awards granted by the ultimate controlling party

MPSC has granted certain number of its own ordinary shares to the employee of the Group under the share award scheme approved by the board of MPSC with no vesting conditions attached at nil consideration. MPSC and the Group also entered into a recharge arrangement approximate to the grant-date fair value of this shared-based payment and the recharge is required to be paid after the shares are awarded. The fair value of services received in return for the shares awarded of RMB70,000 and nil for the six months ended 30 June 2023 and 2022, respectively, which is measured by the grant-date share price of MPSC, was recognised as expenses on the grant date with a corresponding increase in trade and other payables due to MPSC.

(e) Employee share purchase plan (the "ESPP")

Since 2015, the Group adopted several ESPPs, pursuant to which, the partnership firms, whose limited partners consisted of employees of the Group, invested in the Group by way of subscribing newly issued equity interests of MP NeuroTech Shanghai. All participants of the ESPPs have purchased equity interests in respective partnership firms at amounts specified in the respective partnership agreements.

All ESPPs contain a service condition. Employees participating in the plan have to transfer out their equity interests if their employments with the Group were terminated within the vesting period, to a person or a party nominated by the general partners of the partnership firms at a price no higher than the amounts specified in the respective partnership agreements. The fair value of the ESPP at the grant date, being the difference between the considerations and the fair value of the equity interests subscribed shall be spread over the vesting period and recognised as staff costs in the profit or loss.

The total expenses recognised in the consolidated statement of profit or loss for the above ESPP are RMB285,000 and RMB162,000 for the six months ended 30 June 2023 and 2022, respectively.

(Expressed in Renminbi unless otherwise indicated)

14 Capital and reserves (continued)

(f) Share award scheme

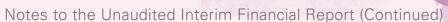
Pursuant to the Share Award Scheme adopted by the Company approved by the Board in 2023, the Company may purchase its own shares and grant such shares to certain employees of the Group at nil consideration. For the six months ended 30 June 2023, the Company granted 516,717 shares (six months ended 30 June 2022: nil) with a fair value of HKD7,544,000 (equivalent to RMB6,652,000, six months ended 30 June 2022: nil) to the Group's executives and certain employees to settle the discretionary bonuses.

15 Financial assets measured at fair value through profit or loss

	30 June 2023 RMB'000	31 December 2022 RMB'000
Wealth management products	266,444	266,053

As at 30 June 2023, the Company held five wealth management products subscribed from five segregated portfolio companies incorporated in the Cayman Islands, with purchase cost amounted to US\$36,450,000 (equivalent to RMB249,368,000) in aggregate at annualised return rate of 1.6% or 1.5%-4.5%. The Company can redeem the wealth management products at any time upon expiration of lock-up period (if any).

The fair values of the wealth management products are within level 3 of the fair value hierarchy as disclosed in Note 16(a).



(Expressed in Renminbi unless otherwise indicated)

16 Fair value measurement of financial instruments

(a) Financial assets and liabilities measured at fair value

Fair value hierarchy

The following table presents the fair value of the Group's financial instruments measured at the end of each reporting period on a recurring basis, categorised into the three-level fair value hierarchy as defined in HKFRS 13, *Fair value measurement*. The level into which a fair value measurement is classified is determined with reference to the observability and significance of the inputs used in the valuation technique as follows:

Level 1 valuations: Fair value measured using only Level 1 inputs i.e. unadjusted quoted prices in

active markets for identical assets or liabilities at the measurement date

— Level 2 valuations: Fair value measured using Level 2 inputs i.e. observable inputs which fail to

meet Level 1, and not using significant unobservable inputs. Unobservable

inputs are inputs for which market data are not available

Level 3 valuations: Fair value measured using significant unobservable inputs

(Expressed in Renminbi unless otherwise indicated)

16 Fair value measurement of financial instruments (continued)

(a) Financial assets and liabilities measured at fair value (continued)

Fair value hierarchy (continued)

The Group has a team that manages the valuations of the financial instruments for financial reporting purpose. The team manages the valuation on a case by case basis. External valuation experts will be involved when necessary.

Fair value at Fair value measurements 30 June as at 30 June 2023 categorised i			
2023 RMB'000	Level 1 RMB′000	Level 2 RMB'000	Level 3 RMB'000
266,444	_	_	266,444
Fair value at 31 December	as at 31 Dece	ember 2022 cate	gorised into
2022 RMB'000	Level 1 RMB'000	Level 2 RMB'000	Level 3 RMB'000
266,053 (272)	_	— (272)	266,053 —
	30 June 2023 RMB'000 266,444 Fair value at 31 December 2022 RMB'000	2023 RMB'000 206,444 Fair value at 31 December 2022 RMB'000 266,053 As at 30 June as at 30 June Level 1 RMB'000 RMB'000 As at 31 December as at 31 December Amber as at 31 December Amber	Section 2023

During the six months periods ended 30 June 2022 and 2023, there were no transfers between Level 1 and Level 2, or transfers into or out of Level 3. The Group's policy is to recognise transfers between levels of fair value hierarchy as at the end of each of the reporting period in which they occur.

(Expressed in Renminbi unless otherwise indicated)



(a) Financial assets and liabilities measured at fair value (continued)

Valuation technique and input used in Level 2 fair value measurement

These foreign currency forward contracts are fair valued by comparing the contracted rate to the exchange rate of relevant currency issued by the bank at the end of reporting period. All movements in fair value are recognised in profit or loss in the period they occur.

Information about Level 3 fair value measurements

	Valuation techniques	Significant unobservable inputs
Wealth management products measured at FVPL	Net asset value	Expected rate of return of 1.6% or 1.5%–4.5% (Note)

Note: The wealth management products measured at FVPL is redeemable at a redemption price equals to the net asset values as stated in the monthly shareholder's statement issued by the administrators, which represents a rate of return of 1.6% or 1.5%-4.5%. The fair value measurement is positively correlated to the expected rate of return. As at 30 June 2023, it is estimated that with all other variables held constant, an increase/decrease of 100 basis points in the expected rate of return would have increase/decrease the Group's profit by RMB1,020,000/RMB1,020,000.

The movement during the six-month period ended 30 June 2023 in the balance of this Level 3 fair value measurement are as follows:

	Wealth management products RMB'000
At 1 January 2023	266,053
Exchange adjustments	9,863
Purchase of wealth management products	57,067
Redemption of wealth management products	(68,660)
Changes in fair value recognised in profit or loss	2,121
At 30 June 2023	266,444

(b) Fair value of financial assets and liabilities carried at other than fair value

The carrying amounts of the Group's financial instruments carried at cost or amortised cost were not materially different from their fair values as at 30 June 2023 and 31 December 2022.

(Expressed in Renminbi unless otherwise indicated)

17 Commitments

Capital commitments in respect of property, plant and equipment and intangible assets outstanding at 30 June 2023 not provided for in the interim financial statements are as follows:

	30 June 2023 RMB'000	31 December 2022 RMB'000
Contracted for Approved but not contracted for	1,371 239,653	4,373 253,801
	241,024	258,174

18 Material related party transactions

(a) Key management personnel remuneration

Six months ended 30 June

	2023 RMB'000	2022 RMB'000
Salaries and other benefits Discretionary bonuses Equity-settled share-based payment expenses	2,710 3,250 —	2,851 2,684 91
	5,960	5,626

(Expressed in Renminbi unless otherwise indicated)

Relationship

18 Material related party transactions (continued)

(b) Related parties

Name of party

Particulars of the Group's other transactions with related parties other than key management personal remuneration during six months ended 30 June 2023 and 2022 are as follows:

MPSC	Ultimate controlling party of the Group	
MicroPort Product Innovation Inc	Subsidiary of MPSC	
MicroPort CRM Japan Co., Ltd.	Subsidiary of MPSC	
MicroPort Scientific Ltd.	Subsidiary of MPSC	
MicroPort Scientific Vascular Brasil Ltda.	Subsidiary of MPSC	
MicroPort Sinica Co., Ltd.* (微創投資控股有限公司)	Subsidiary of MPSC	
Shanghai MicroPort Medical (Group) Co., Ltd.*	Subsidiary of MPSC	
(上海微創醫療器械(集團)有限公司)		
Shanghai Shenyi Medical Technology Co., Ltd.*	Subsidiary of MPSC	
(上海神奕醫療科技有限公司)		
Shanghai ShenTai Medtech Co., Ltd.*	Subsidiary of MPSC	
(上海神泰醫療科技有限公司)		
Shanghai SafeWay Medtech Co., Ltd.*	Subsidiary of MPSC	
(上海安助醫療科技有限公司)	0.1.11. (1.4.000)	
D-pulse Medical (Beijing) Co., Ltd.*	Subsidiary of MPSC	
(龍脈醫療器械(北京)有限公司)	Subsidient of MDCC	
MicroPort Access Medtech (Jiaxing) Co., Ltd.* (微創龍脈醫療科技(嘉興)有限公司)	Subsidiary of MPSC	
「阪 削 能 灰 香 凉 行 X (新 央 / 有 水 ム ロ / Fujian Kerui Pharmaceutical Co., Ltd.*	Subsidiary of MPSC	
(福建科瑞藥業有限公司)	Substituting of tvir SC	
Suzhou Micropet Health Technology Co., Ltd.*	Subsidiary of MPSC	
(蘇州微寵健康科技有限公司)	o abolalary or wire of	
Shanghai Weizhuo Technology Co., Ltd.*	Subsidiary of MPSC	
(上海微琢科技有限公司)	,	
Shanghai MicroPort Endovascular MedTech Co., Ltd.*	Subsidiary of MPSC	
(上海微創心脈醫療科技(集團)股份有限公司		
Suzhou ProSteri Medical Technology Co., Ltd.*	Equity-accounted investee of	
(蘇州諾潔醫療技術有限公司)	MPSC	
Zhejiang Accupath Smart Manufacturing (Group) Co., Ltd.*	Equity-accounted investee of	
(浙江脈通智造科技(集團)有限公司)	MPSC	
AccuTarget MediPharma (Shanghai) Co., Ltd.*	Equity-accounted investee of	
(上海導向醫療系統有限公司)	MPSC	
Shanghai HuaRui Bank Co., Ltd.*	Equity-accounted investee of	
(上海華瑞銀行股份有限公司, "SHRB")	MPSC	
Rapid Medical Ltd.	Equity-accounted investee of the	

^{*} English translation is for identification purpose only.

Group

(Expressed in Renminbi unless otherwise indicated)

18 Material related party transactions (continued)

(c) Leasing arrangement with related parties

In February 2020 and May 2021, MP NeuroTech Shanghai leased out its own properties to a related party and recognised rental income amounted to RMB1,008,000 for the six months ended 30 June 2023 (six months ended 30 June 2022: RMB804,000).

(d) Cash deposits placed in a related party

As at 30 June 2023, the Group has placed cash deposits amounted to RMB90,641,000 in SHRB with interest rate range of 1.8%–3.45% per annum.

(e) Other transactions with related parties

Particulars of the Group's other transactions with related parties during the six months ended 30 June 2022 and 2023 are as follows:

Six months ended 30 June

	2023 RMB'000	2022 RMB'000
Sales of goods to an equity-accounted investee of the Group	3,389	5,978
Sales of materials to a subsidiary of MPSC	19	_
Service fee charged by subsidiaries of MPSC	5,174	3,072
Service fee charged by an equity-accounted investee of MPSC	875	675
Purchase of goods from subsidiaries of MPSC	4,501	3,096
Purchase of goods from an equity-accounted investee of MPSC	5,354	1,554
Purchase of equipment from subsidiaries of MPSC	619	_
Payment on behalf of the Group by subsidiaries of MPSC	7,238	3,139
Payment on behalf of related parties by the Group	148	171