

# MicroPort Scientific Corporation

## 2024 Interim Results

30 August 2024

Rega DR  
7202  
SN C8A4K24  
Made in China

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# Strong Results Underpinned by Strategic Focus and Transformation Efforts

## Transformation Underway

Reshaping our business with full P&L responsibility and to create shareholder value








Strategic Priorities	1H2024 Progress
Category Leadership	<ul style="list-style-type: none"> <li>Revenue\$558.7 mn ▲17.0%<sup>YOY</sup></li> <li>Leading market share in domestic for major segments</li> </ul>
Meaningful Innovation	<ul style="list-style-type: none"> <li>31 NMPA, 11 CE Mark, 4 FDA approvals YTD</li> <li>4 Green Paths YTD, 34 in total, No.1 in industry for 9 consecutive years</li> <li>Firesorb® obtained NMPA approval</li> </ul>
Global Expansion	<ul style="list-style-type: none"> <li>Going-abroad business revenue ▲44.0%<sup>YOY</sup></li> <li>Cardio. ▲56.3%<sup>YOY</sup> Endo. ▲65.0%<sup>YOY</sup> Neuro. ▲87.0%<sup>YOY</sup> Structural Heart▲29.2%<sup>YOY</sup> Surgical Robots▲293.2%<sup>YOY</sup></li> </ul>
Unlock Efficiency	<ul style="list-style-type: none"> <li>Total Operating Exp. ratio significantly dropped from 94% to 64%<sup>YOY</sup></li> <li>R&amp;D Exp. ratio substantially reduced from 39% to 21%<sup>YOY</sup></li> </ul>
Narrowed Strategic Focus	<ul style="list-style-type: none"> <li>Divestment of non-core business &amp; asset</li> <li>Shut down R&amp;D projects at early stages based on evaluation</li> </ul>

Non-HKFRS Net Loss Substantially Reduced by ▼63.1%<sup>YOY</sup>

Note: 1. revenue growth rate adjusted to exclude foreign exchange impact, due to appreciation or depreciation of US\$ against functional currencies.

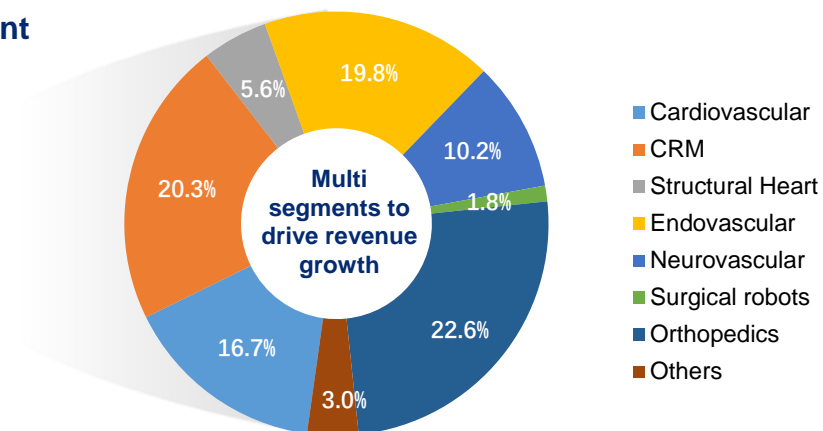
# Positive Momentum Across Business Segments

## Rapid Revenue Growth

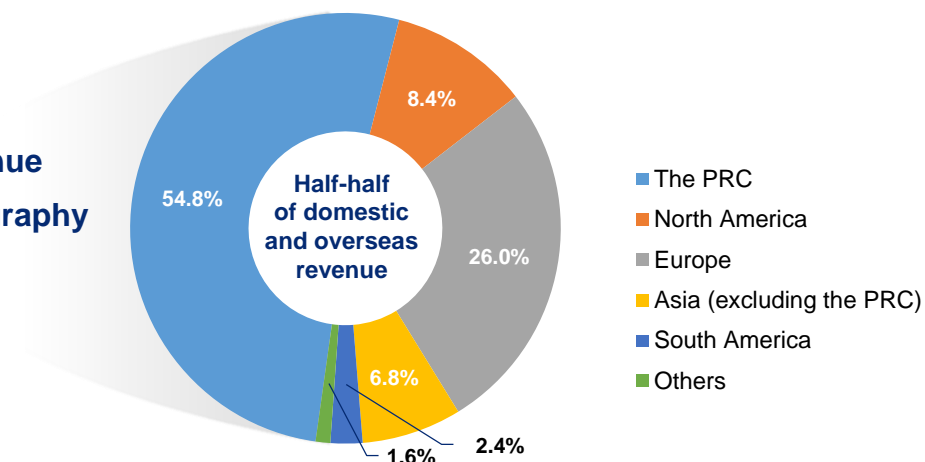
Segment	Revenue	YoY Changes	China Revenue YoY Changes
 Orthopedics	\$126.3mn	▲ 9.0%	▲ 32.5%
 CRM	\$113.4mn	▲ 5.6%	▲ 61.5%
Segment	Revenue	YoY Changes	Going-abroad Revenue YoY Changes
 Cardiovascular	\$93.3mn	▲ 13.4%	▲ 56.3%
 Endovascular	\$110.4mn	▲ 26.3%	▲ 65.0%
 Neurovascular	\$57.1mn	▲ 36.5%	▲ 87.0%
 Structural Heart	\$31.1mn	▲ 26.7%	▲ 29.2%
 Surgical Robots	\$10.0mn	▲ 117.0%	▲ 293.2%

## Diversified Business

### ➤ Revenue by Segment



### ➤ Revenue by Geography

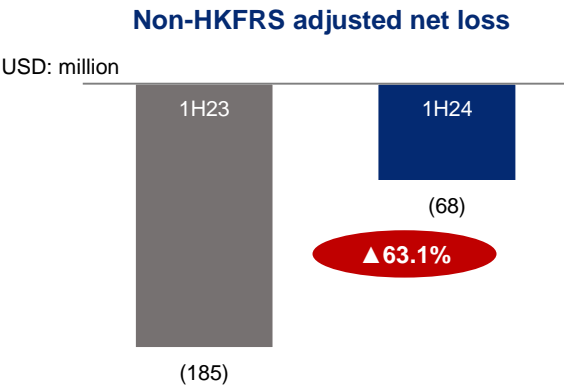
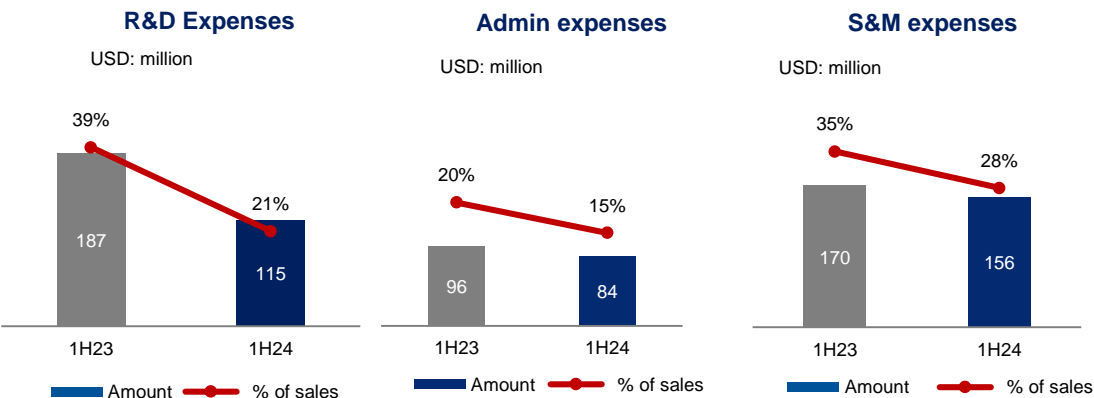
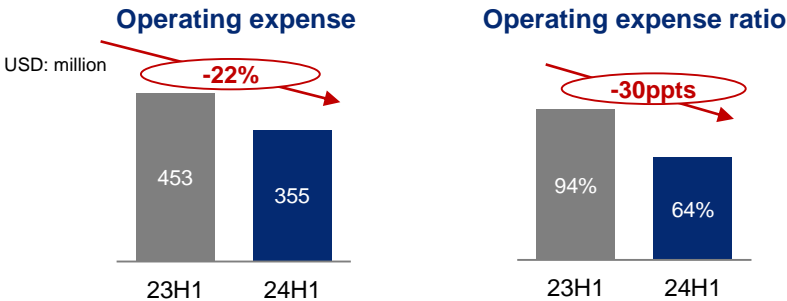


# Substantially Expense Reduced & Cost Effectiveness Increased

Improved Operating Efficiency

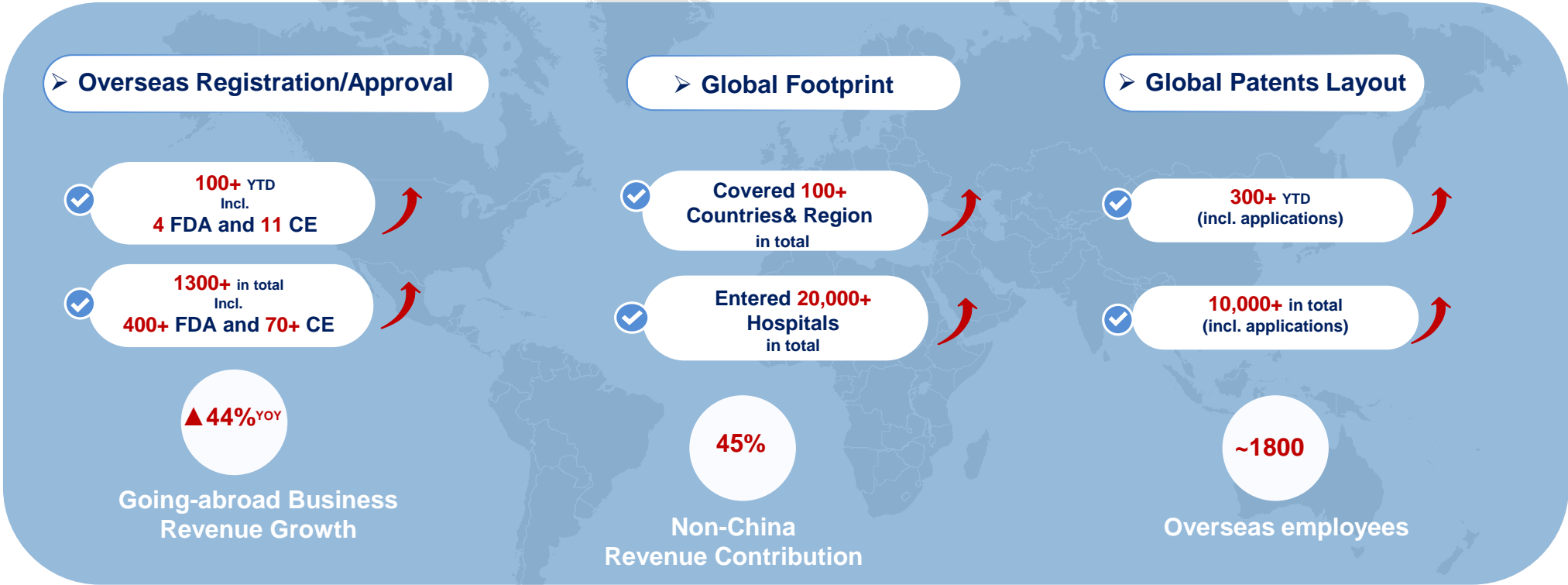
Strict Cost Control Across Key Items

Non-HKFRS Adjusted Net Loss Substantially Narrowed



Note: 1. revenue growth rate adjusted to exclude foreign exchange impact, due to appreciation or depreciation of US\$ against functional currencies.

# Expanding Our International Presence With Broad-based Capabilities



Possession of Comprehensive Product Portfolio

Established Global Innovation Network System

Recognized Overseas Clinical Trial Evidence

Extensive & Profound Distribution Channel

Leveraging Inter-Group Synergy

Tapping into Diversified Markets

Note: 1. revenue growth rate adjusted to exclude foreign exchange impact, due to appreciation or depreciation of US\$ against functional currencies. 2. refers to initial registration/approval; 3. Registration data as of August 30, 2024.

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# Cardiovascular Business: Net Profit Substantially Increased by 279.4%<sup>YOY</sup>

## Most Complete Product Pipeline In Coronary Field

### Key Financials



### Business Highlights



China











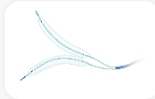





- 1H24 revenue up 4.4%<sup>YOY</sup>, among which the DES products growth are higher than the market level, further enhanced its position of No.1 market share
- Hospital coverage expanding: DES portfolio has accumulatively entered 3,500+ hospitals, achieving stable growth in market share
- The new generation of bioresorbable scaffold Firesorb® obtained NMPA approval
- Atherectomy Wire obtained NMPA approval



Non-China

- 1H24 revenue up 56.3%<sup>YOY</sup>, strong growth achieved in EMEA (up 107.9%<sup>YOY</sup>) Asia-Pacific (excluding China, up 43.4%<sup>YOY</sup>) and Latin America (up 6.1%<sup>YOY</sup>)
- Extensive sales network: DES sales cover over 80+ countries and regions
- Diversified & flexible product portfolio to maintain a leading market position

### Complete & Diversified Product Pipeline

Diagnostic Device	Active		Non-active	
		<b>Argusclarity® Insight-100</b> Optical Coherence Tomography System ◆ NMPA approval		<b>Argusclarity® Insight-100h</b> Optical Coherence Tomography System ◆ NMPA approval
		<b>Soul-Man™</b> Digital Subtracting Angiography System ◆ NMPA approval		<b>Deycypher®</b> Intravascular Ultrasound Imaging System Upcoming approval by NMPA
Therapeutic Device		<b>Integrated Coronary Intravascular Lithotripsy System</b> Upcoming approval by NMPA		<b>Integrated Coronary Transluminal Rotational Atherectomy System</b> Upcoming approval by NMPA
		<b>Firebird2®</b> Rapamycin-Eluting Coronary Cocr Stent System ◆ NMPA approval		<b>Firehawk™</b> Rapamycin Target Eluting Coronary Stent System ◆ NMPA approval
		<b>Firesorb®</b> Bioresorbable Rapamycin Eluting Coronary Scaffold System ◆ NMPA approval		<b>Coronary Sirolimus Eluting Balloon Catheter</b> Upcoming approval by NMPA
		<b>Pioneer™</b> PTCA Balloon Catheter ◆ NMPA approval		<b>Firefighter™</b> PTCA Balloon Catheter ◆ NMPA approval
		<b>AnchorV®</b> Exchange Device ◆ NMPA approval		<b>Coronary Scoring Balloon Catheter</b> Upcoming approval by NMPA
		<b>Interline™</b> Guiding Catheter ◆ NMPA approval		<b>Bilumos®</b> Dual-lumen Catheter ◆ NMPA approval

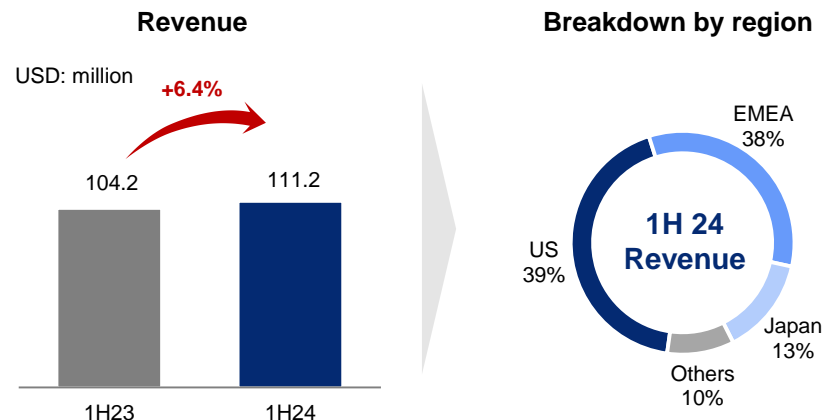
Note: 1. revenue growth rate adjusted to exclude foreign exchange impact, due to appreciation or depreciation of US\$ against functional currencies.

# Orthopedics Business: Positive EBITDA Achieved

## Revenue Increased By 9.0% YoY



### Non-China Business

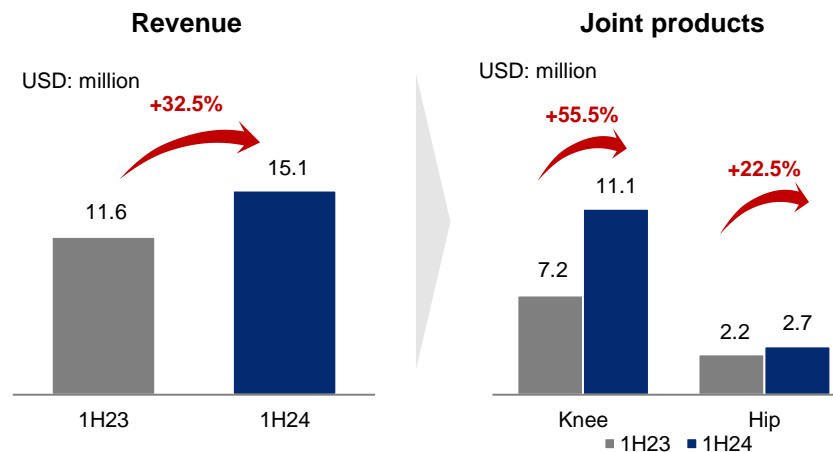


#### Business Highlights

- **1H24 revenue up 6.4%<sup>YoY</sup>**, continuously improving global supply chain with suppliers being proactively diversified, backorders back to normal operating levels
- **Strong growth in the international markets (EMEA up 19.5%<sup>YoY</sup>, Japan up 8.0%<sup>YoY</sup>)** while US declined (-4.5%<sup>YoY</sup>) under short-term pressure for commercial penetration due to delayed impact from historical backorders
- **Sales revenue of Knee portfolio up 12.5%<sup>YoY</sup>**, driven by the growing recognition of the premium Medial Pivot Knee system, as well as the successful execution of SkyWalker™ commercial strategy
- SkyWalker™ & Evolution® application with **~300 case of TKA** surgeries performed in **10+** hospitals/ institutions in US and Europe
- **New Products launched:**
  - (Knee) Kinematic Alignment, CCK Revision, Evolution NitrX
  - (Hip) Xelha AR navigation, Dynasty Dual Mobility, Gladiator HA Coated Modular Stem System



### China Business



#### Business Highlights

- **1H24 revenue up 32.5%<sup>YoY</sup>**, driven by rapid growth in both implantation volume and sales volume of hip and knee joint products
- Further expanded hospital coverage with regional coverage efficiency strategically enhanced
- Strict implementation of cost-control measures, GPM improved **35 pts**
- **New Products launched:**
  - Evolution® CCK Knee System and zirconium-niobium femoral condyle obtained NMPA approval, enhanced ex-VBP product portfolio

■ Products newly approved for marketing

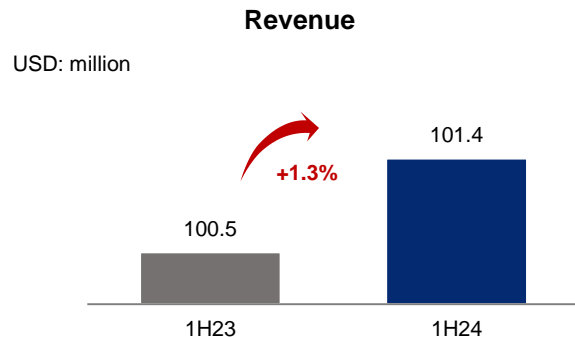
Note: 1. revenue growth rate adjusted to exclude foreign exchange impact, due to appreciation or depreciation of US\$ against functional currencies.

# CRM Business: EBITDA significantly improved

## Revenue Increased By 5.6%<sup>YoY</sup>



### Non-China Business



#### Business Highlights

- **1H24 revenue up 1.3%<sup>YoY</sup>**
  - upstream parts supply problem has been comprehensively solved
- **Expediting market entry:**
  - CE Approval for TALENTIA and ENERGY defibrillators, cardiac resynchronization therapy
  - defibrillators (CRT-Ds), and XFine™ leads
  - TALENTIA ICDs and CRT-Ds approved in Australia
  - EDIS and GALI defibrillator systems approved in Cyprus
- **1st commercial implantation :**

#### High Voltage

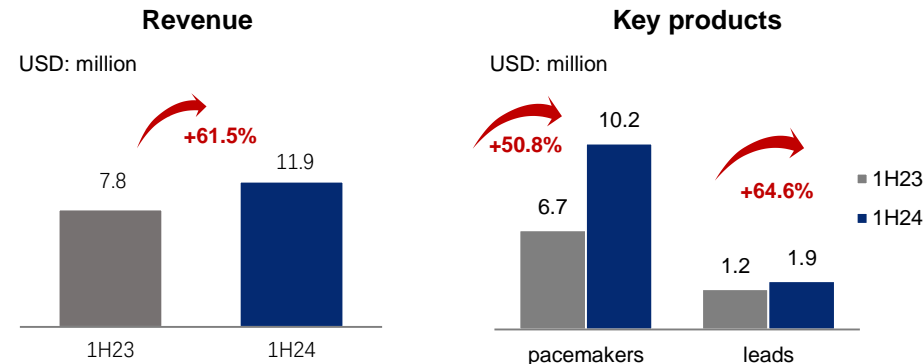
- TALENTIA™ CRT-D in Europe.
- first GALI™ CRT-D SonR® system in Japan

#### Low Voltage

- Alizea™ Bluetooth® pacemaker system in the US



### China Business

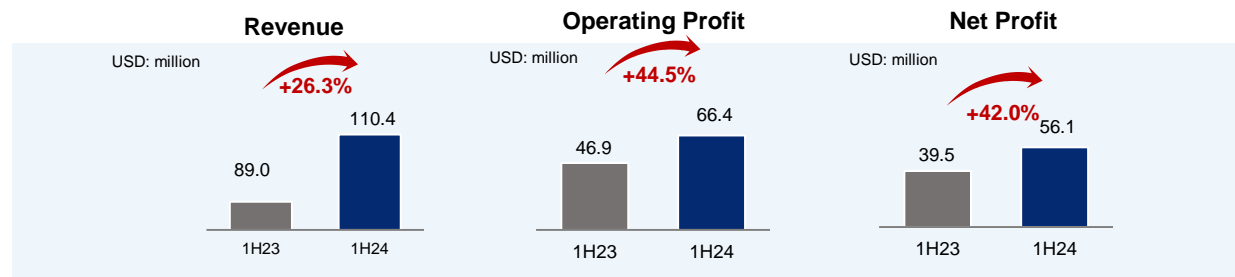


#### Business Highlights

- **1H24 revenue up 61.5%<sup>YoY</sup>**, with revenue of pacemakers **up 50.8%<sup>YoY</sup>**; revenue of leads **up 64.6%**
- **Accelerated market penetration** driven by continuous bids winning in VBP tenders, further consolidate **Top 1** market position among domestic pacemakers
- **GPM improved by 22 ppts** led by proactive dynamic adjustment of product mix and continuous efforts in reduction of costs
- **Further enriched product portfolio strengthens competitive position:** NMPA approval obtained for
  - imported 1.5T/3T full-body MRI compatible ENO™ pacemaker, bridged the intergenerational divide with MNCs, offering premium choices to Chinese patients
  - the domestic 1.5T/3T full-body MRI compatible pacemaker

# Endovascular Business

## Key Financials



## Business Highlights



China

- **1H24 revenue up 23.6%<sup>YOY</sup>**, key products recorded steady growth, accelerated hospital entry of new products
- **Innovative products approved in recent years accelerate market penetration:** Castor<sup>®</sup>, world's first-in-class thoracic branch stent-graft system, covered **1,100+** hospitals, Minos<sup>®</sup> covered **nearly 900** hospitals, Reewarm<sup>®</sup> PTX covered **1000+** hospitals, Fontus<sup>®</sup> covered **200+** hospitals; Talos<sup>®</sup> covered **nearly 300** hospitals since approval
- **Further enriched product pipeline:** L-REBOA<sup>®</sup>, Vewatch<sup>®</sup>, Vepack<sup>®</sup>, Vflower<sup>®</sup>, ReeAmber<sup>®</sup> were approved for marketing by the NMPA



Non-China

- **1H24 revenue up 65.0%<sup>YOY</sup>**, core products accelerate the development and layout of overseas markets
- **Rapid progressing global launch of core products:** Castor<sup>®</sup> entered into **19** countries, Minos<sup>®</sup> entered into **21** countries, Hercules<sup>®</sup> Low Profile entered into **22** countries
- **Swiftly advancing overseas sales channel:** newly developed **3** countries, innovative products covered **34** markets accumulatively, across EU, Latin America and Southeast Asia
- **Achieved significant progress for R&D projects:** **8** new certificates were obtained in overseas markets; Cratos<sup>®</sup> received the EU Customized Certificate

### Aortic Intervention

Product	Pre-clinical	Clinical	Registration
L-REBOA <sup>®</sup> Aortic Occlusion Balloon Catheter	✓ Obtained <b>NMPA</b> approval		
Cratos <sup>®</sup> Branched Aortic Stent-Graft System	• Submitted for <b>NMPA</b> review		
Aegis <sup>®</sup> II Abdominal Aortic Stent-Graft System	• Conducting pre-market clinical trial		
Hector <sup>®</sup> Multi-branched Aortic Stent-Graft System	• Conducting FIM clinical trial		
Aortic Tear Flow-Restriction Stent	• Conducting FIM clinical trial		

### Peripheral Venous Intervention

Vflower <sup>®</sup> Venous Stent System	✓ Obtained <b>NMPA</b> approval		
Vewatch <sup>®</sup> Vena Cava Filter	✓ Obtained <b>NMPA</b> approval		
Vepack <sup>®</sup> Filter Retriever	✓ Obtained <b>NMPA</b> approval		
Fishhawk <sup>®</sup> Mechanical Thrombectomy Catheter	• Completed pre-market clinical trial		

### Peripheral Arterial Intervention

ReeAmber <sup>®</sup> Balloon Dilation Catheter	✓ Obtained <b>NMPA</b> approval		
Ryfle <sup>®</sup> Fibered Embolization Coil	• Submitted for <b>NMPA</b> review		
Below-The Knee Drug Coated Balloon Catheter	• Conducting pre-market clinical trial		

### Oncological Intervention

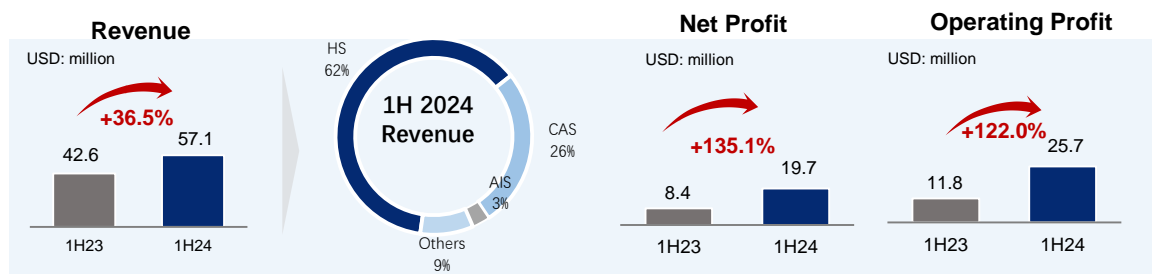
HepaFlow <sup>®</sup> TIPS Stent Graft System	• Completed pre-market clinical trial		
Transjugular Liver Access Set	• Submitted for <b>NMPA</b> review		
Polyvinyl Alcohol (PVA) Embolic Microspheres	• Conducting pre-market clinical trial		

Product admitted to NMPA Green Path

Note: 1. revenue growth rate adjusted to exclude foreign exchange impact, due to appreciation or depreciation of US\$ against functional currencies.

# Neurovascular Business

## Key Financials



## Business Highlights



China

- **1H24 revenue up 33.8%<sup>YOY</sup>**, key products recorded steady growth, accelerated hospital entry of new products
- **Coverage of high-quality hospitals accelerates market penetration:** approximately **300** hospitals have been newly covered, with a total coverage of about **3300** hospitals, accumulated support for approximately **190,000** neuro-interventional surgeries
- **Core product sales boost market share growth:** NUMEN® Coil and APOLLO™ Intracranial Stent benefited from winning the VBP bids, which accelerated the development of new markets; the number of X-track® admitted to hospitals doubled, and clinical use increased by approximately **800%<sup>YOY</sup>**



Non-China

- **1H24 revenue up 87.0%<sup>YOY</sup>**, achieved a breakthrough in international business, especially in Asia Pacific, Latin America and EMEA
- **Worldwide commercialization process accelerated:** a total of 8 products that have been launched into the overseas market, and have been commercialized in **21** overseas countries, covering 9 of the top 10 countries worldwide in terms of the number of neuro-interventional procedures
- **Japan:** NUMEN® Coil has entered more than **240** local hospitals
- **France:** NUMEN® Coil achieved **first** commercial clinical application
- **Brazil:** Tubridge® and Neurohawk® achieved **first** commercial implantation/ usage
- **Argentina:** Tubridge® and X-track® achieved **first** commercial implantation/ usage

### Hemorrhagic Stroke

Product	Pre-clinical	Clinical	Registration
Tubridge Plus® Flow-diverting Stent			✓ NMPA approved in Aug 2024
Numen Silk® 3D Electroically Detachable Coil			✓ NMPA approved • FDA approval expected in 2024
Numen Uni® Electroically Detachable Coil			• NMPA approval expected in 2024
NuFairy™ Absorbable Coil Embolization System			• Conducting clinical trial
Rebridge® Intracranial Visualized Stent			• Conducting clinical trial

### CAS

Safecer™ Embolic Protection Device			✓ NMPA approved in Jun 2024
Pathfinder™ Carotid Artery Dilatation Catheter			✓ NMPA approved in Apr 2024
Intracranial Drug-Coated Balloon Catheter System			• Conducting clinical trial
Intracranial Autodistensible Drug Stent			• Design Validation
Intracranial Bulbar Expansion Drug Stent			• Design Validation

### AIS

Neurohawk® Pass17/21 Stent Thrombectomy Device			✓ NMPA approved in Jul 2024
NeuroGuard® Balloon Protection Guide Catheter			✓ NMPA approved in Jan 2024

### Access Product

17 Microcatheter			• Design Validation
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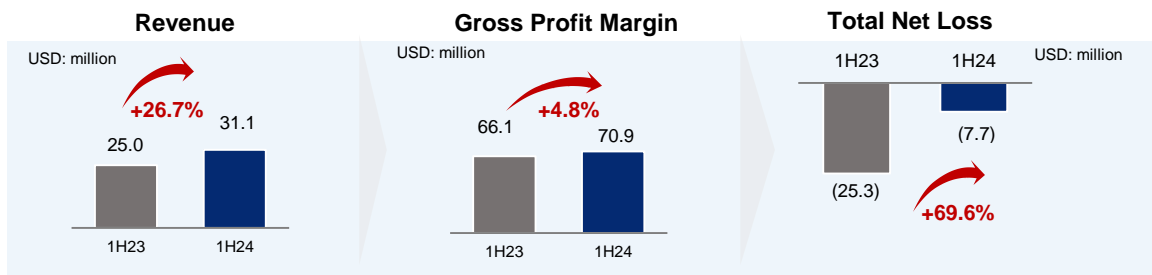
Product admitted to NMPA Green Path

Note: 1. revenue growth rate adjusted to exclude foreign exchange impact, due to appreciation or depreciation of US\$ against functional currencies.



# Structural Heart Business

## Key Financials



## Business Highlights



China

- **1H24 revenue up 26.4%<sup>YOY</sup>**, primarily attributable TAVI products and procedural accessories in the PRC owing to the increased hospital penetration
- **Focusing on enhancement of TAVI accessibility in China:** new access to **50** hospitals brought the Company's coverage to more than **600** hospitals, implantation volume increased by around 10%<sup>YOY</sup>
- **Official business expansion into stroke prevention in patients with nonvalvular atrial fibrillation:** acquired 51% equity of MP CardioAdvent, and the first batch commercial implantations of the self-developed AnchorMan® LAAC System has been completed up to date after receiving approval by the NMPA in January, 2024



Non-China

- **1H24 revenue up 29.2%<sup>YOY</sup>**, core products accelerate the development and layout of overseas markets
- **Rapid progressing global launch of core products:** **VitaFlow Liberty®** obtained **CE mark** and achieved commercial implantation; also received registration approvals in Hong Kong, Saudi Arabia, Belarus and Malaysia; TAVI products have entered nearly **100** hospitals in Argentina, Colombia, Thailand, Russia, Chile and Switzerland
- **Successfully integrated LAA portfolio:** Alwide® Plus reached milestone achievements in emerging markets; CE registration of Alwide® Plus, AnchorMan® LAAC System and AnchorMan® LAA Access System entered key approval process

	Product	Pre-clinical	Clinical	Registration
TAVI	VitaFlow Liberty®	✓ Obtained <b>NMPA</b> approval ✓ Obtained <b>CE</b> approval		
	VitaFlow Liberty® Flex	• NMPA Registration in progress		
	VitaFlow® IV	• Design stage		
	VitaFlow® Balloon Expandable	• Design stage		
Accessories	Alwide® Plus Balloon Catheter	• CE Marking and registration in emerging markets in progress		
	AccuSniper™ Double Layer Balloon Catheter	✓ Obtained <b>NMPA</b> approval		
	Alpass® Catheter Sheath II	• Submitted for <b>NMPA</b> review		
TMV	Replacement product (Self-developed)	• Progressing FIM study		
	AltaValve™ - Replacement product (Partnership with 4C Medical)	• Received breakthrough and IDE approval of FDA		
TTV	Replacement product	• Design Stage		
LAA products	AnchorMan® Left Atrial Appendage Closure System	✓ Obtained <b>NMPA</b> approval • Submitted for <b>CE</b> review		
	AnchorMan® Left Atrial Appendage Access System	✓ Obtained <b>NMPA</b> approval • Submitted for <b>CE</b> review		

# Surgical Robot Business

## Key Financials



## Business Highlights

- 1H24 revenue up **65.3%**<sup>YOY</sup>, recorded robust sales growth and rapid pace of commercialization

- Leading position among domestic brands



Toumai®	R-ONE®
<ul style="list-style-type: none"> <li>7 units of new commercial installation</li> <li>Overall commercial installation base reached <b>20</b> units to date, <b>No.1</b> among domestic brands</li> </ul>	<ul style="list-style-type: none"> <li>2 units of commercialized installation following the NMPA approval in Dec 23</li> </ul>

- 1H24 revenue up **293.2%**<sup>YOY</sup>, overseas markets reached a milestone in commercialization, **Toumai® obtained CE Mark**

- Ambitious Overseas Penetration



Toumai®	SkyWalker™
<ul style="list-style-type: none"> <li>Successfully completed the <b>first 2</b> commercial installations with revenue breakthrough</li> <li>Received over <b>10</b> units of overseas orders up to date</li> <li>Conducted <b>~100</b> clinical operations in overseas hospitals up to date</li> <li>Created <b>over 20</b> records of 1st remote surgery in China and globally</li> </ul>	<ul style="list-style-type: none"> <li>Sales unit <b>doubled</b> YoY on the leverage of synergies with Orthopedics Business</li> <li>Accumulatively received nearly <b>30</b> overseas orders up to date</li> <li>Expanded to cover <b>20+</b> countries across <b>5</b> continents</li> </ul>

	Product	Pre-clinical	Clinical	Registration
Laparoscopic	Toumai® Laparoscopic Surgical Robot	✓ Obtained NMPA approval ✓ Obtained CE marking		
	Toumai® Remote Laparoscopic Surgical Robot		• Conducting Clinical trial/ Clinical Evaluation	
	Toumai® Single-arm Laparoscopic Surgical Robot		• Submitted for NMPA review	
Orthopedic	DFVision® 3D Electronic Laparoscope	✓ Obtained NMPA approval ✓ Obtained CE marking		
	SkyWalker™ Orthopedic Surgical Robot	✓ Obtained NMPA approval ✓ Obtained CE& FDA& ANVISA& TGA & CDSCO approval		
Natural Orifice	Trans-bronchial Surgical Robot		• Conducting Clinical trial/ Clinical Evaluation	
Panvascular	R-ONE® Panvascular Surgical Robot	✓ Obtained NMPA approval		
Percutaneous	iSR'obot® Mona Lisa Robotic Transperineal Prostate Biopsy System	✓ Obtained NMPA approval		

Product admitted to NMPA Green Path

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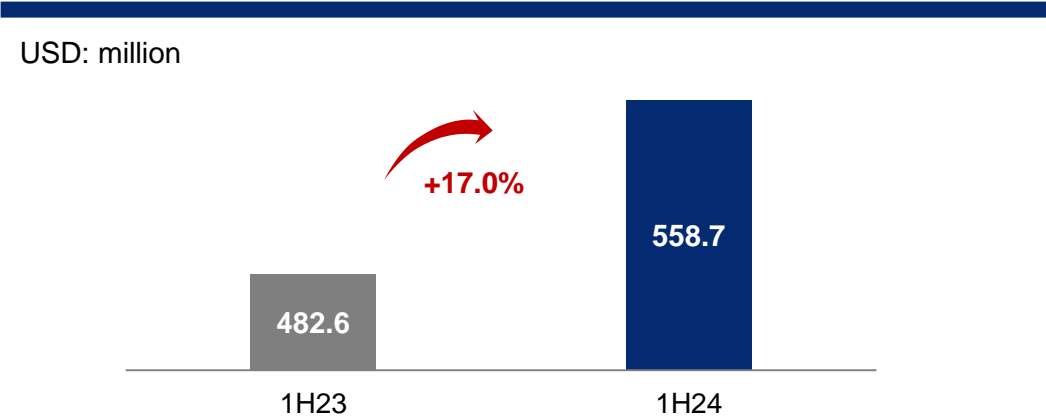
**Financial Review**

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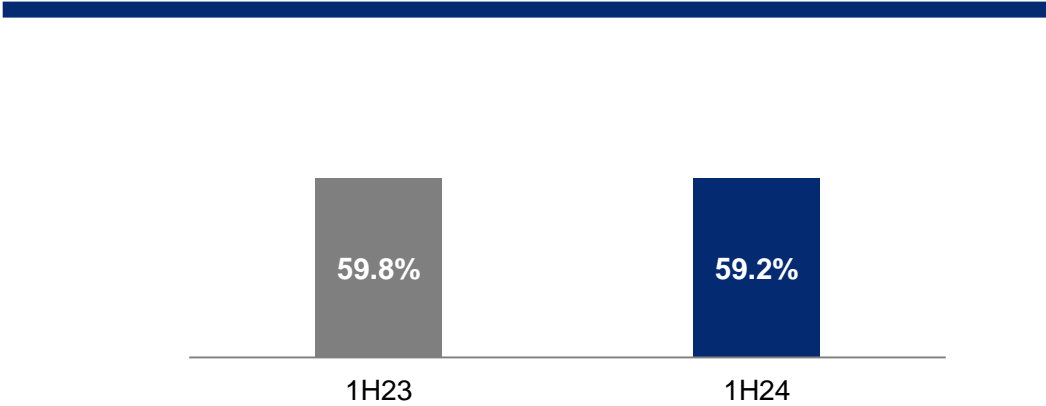
**Appendix – Financial Statements**

# Consolidated Financial Performance

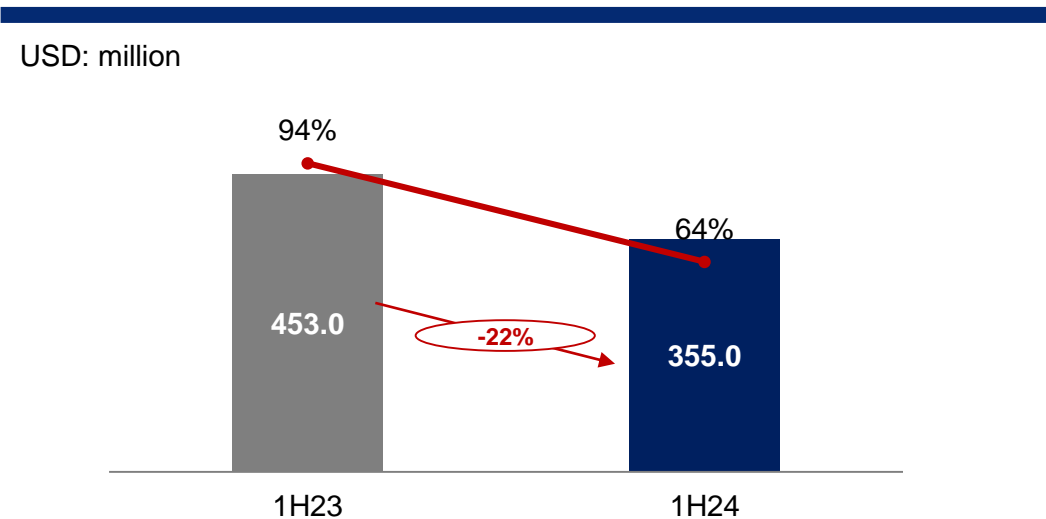
## Revenue



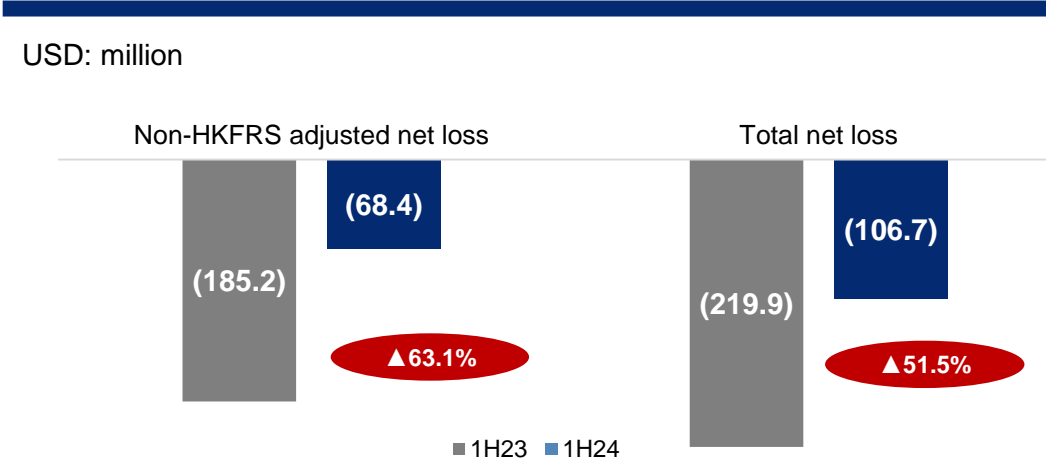
## Gross Profit Margin



## Operating Expenses



## Non-HKFRS Adjusted Net Loss & Total Net Loss



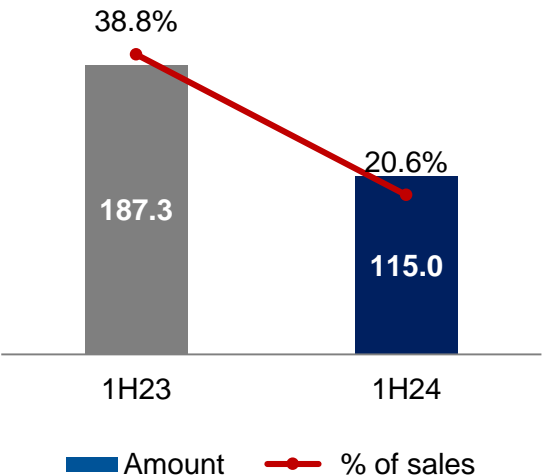
■ Amount      —●— % of sales

Note: revenue growth rate adjusted to exclude foreign exchange impact, due to appreciation or depreciation of US\$ against functional currencies

# Operating Expenses

## R&D Expenses

USD: million

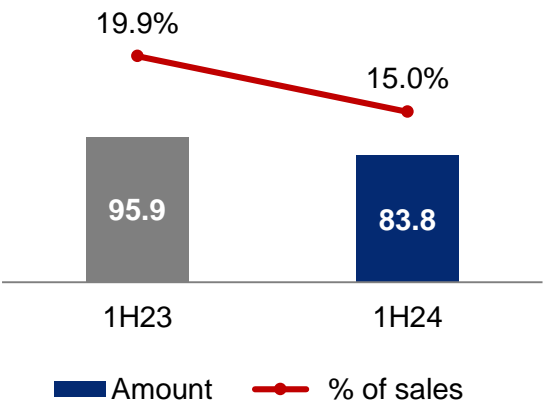


**R&D expenses decreased by 38.6%<sup>YOY</sup>**

- ♦ Due to the proactive cost control and resource focus measures to prioritize and focus on core projects and improve R&D efficiency

## Administrative Expenses

USD: million

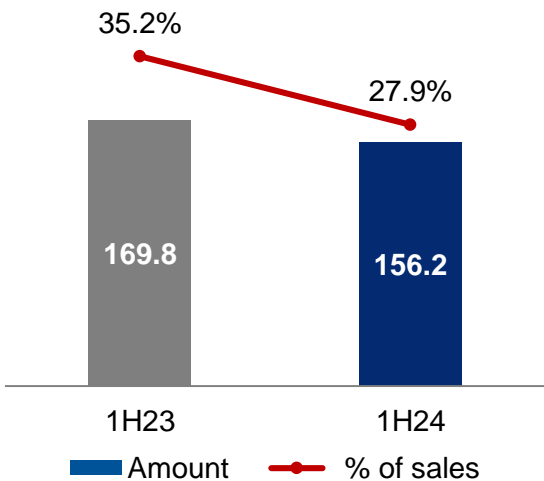


**Administrative expenses decreased by 12.6%<sup>YOY</sup>**

- ♦ Due to the Group's effective cost control and scale of operations

## Sales & Marketing Expenses

USD: million



**Sales & marketing expenses decreased by 8.0%<sup>YOY</sup>**

- ♦ Due to efforts to strengthen the synergy and interconnectivity between overseas and domestic sales platforms.





# Consolidated Income Statement

USD'000	2024 1H	2023 1H	Var.
<b>Revenue</b>	558,702	482,605	15.8%
Cost of sales	(228,122)	(194,189)	17.5%
<b>Gross profit</b>	<b>330,580</b>	<b>288,416</b>	<b>14.6%</b>
Other net income	(68)	17,039	-100.4%
Research and development costs	(115,033)	(187,334)	-38.6%
Distribution cost	(156,150)	(169,800)	-8.0%
Administrative expenses	(83,785)	(95,890)	-12.6%
Other operating costs	(12,348)	(12,374)	-0.2%
<b>Loss from operations</b>	<b>(36,804)</b>	<b>(159,943)</b>	<b>-77.0%</b>
Finance cost	(48,416)	(37,256)	30.0%
Gain on disposal of subsidiaries	6,922	2,845	143.3%
Gain on deemed disposal of interest in equity-accounted investees	-	5,437	-100.0%
Share of profits less losses of equity-accounted investees	(8,146)	(17,258)	-52.8%
<b>Loss before taxation</b>	<b>(86,444)</b>	<b>(206,175)</b>	<b>-58.1%</b>
Income tax	(20,230)	(13,746)	47.2%
<b>Loss for the period</b>	<b>(106,674)</b>	<b>(219,921)</b>	<b>-51.5%</b>
<b>Attributable to: Equity shareholders of the Company</b>	<b>(96,830)</b>	<b>(162,618)</b>	<b>-40.5%</b>

# Consolidated Balance Sheet

USD'000	30 June 2024	31 Dec 2023	Var.
<b>Non-current assets</b>			
Investment properties	6,087	6,256	-3%
Property, plant and equipment	978,657	1,004,573	-3%
Intangible assets	236,699	234,435	1%
Goodwill	147,271	149,393	-1%
Equity-accounted investees	375,085	372,637	1%
Financial assets measured at fair value through profit or loss ("FVPL")	8,479	10,003	-15%
Derivative financial instruments	-	3,574	-100%
Deferred tax assets	30,366	31,382	-3%
Other non-current assets	109,022	109,705	-1%
<b>Total non-current assets</b>	<b>1,891,666</b>	<b>1,921,958</b>	<b>-2%</b>
<b>Current assets</b>			
Inventories	407,912	414,868	-2%
Trade and other receivables	390,504	310,648	26%
Pledged deposits and time deposits	170,948	225,352	-24%
Cash and cash equivalents	740,097	1,019,551	-27%
Financial assets measured at FVPL	196,122	40,028	390%
<b>Total current assets</b>	<b>1,905,583</b>	<b>2,010,447</b>	<b>-5%</b>
<b>Current liabilities</b>			
Trade and other payables	411,693	448,342	-8%
Contract liabilities	18,464	18,770	-2%
Lease liabilities	47,470	46,915	1%
Interest-bearing borrowings	317,891	295,438	8%
Income tax payable	11,648	4,985	134%
Convertible bonds	103,154	549,470	-81%
<b>Total current liabilities</b>	<b>910,320</b>	<b>1,363,920</b>	<b>-33%</b>

## Consolidated Balance Sheet (cont'd)

USD'000	30 June 2024	31 Dec 2023	Var.
<b>Non-current liabilities</b>			
Interest-bearing borrowings	801,523	508,330	58%
Lease liabilities	74,107	85,327	-13%
Deferred income	50,110	42,344	18%
Convertible bonds	341,841	213,267	60%
Contract liabilities	26,733	27,669	-3%
Other payables	275,202	262,865	5%
Deferred tax liabilities	24,343	25,686	-5%
Derivative financial instruments	5,117	-	N/A
<b>Total non-current liabilities</b>	<b>1,598,976</b>	<b>1,165,488</b>	<b>37%</b>
<b>CAPITAL AND RESERVE</b>			
Share capital	18	18	-
Reserves	699,795	757,801	-8%
<b>Total equity attributable to equity shareholders of the Company</b>	<b>699,813</b>	<b>757,819</b>	<b>-8%</b>
Non-controlling interests	588,140	645,178	-9%
<b>Total equity</b>	<b>1,287,953</b>	<b>1,402,997</b>	<b>-8%</b>