

MicroPort Scientific Corporation | 2019 Interim Results

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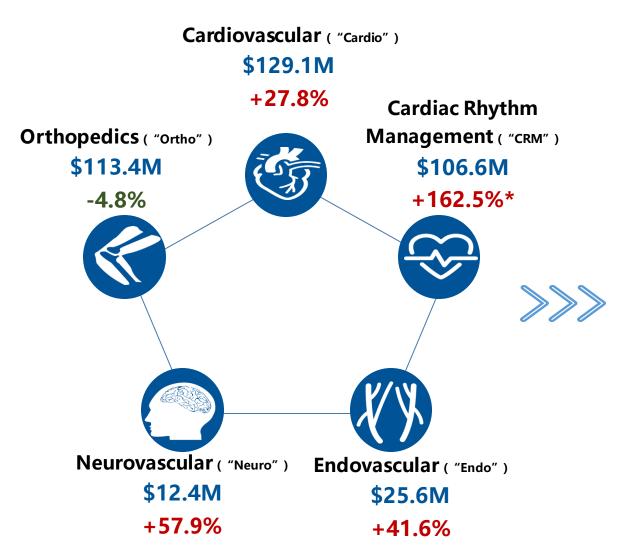
- 1 INTERIM RESULTS HIGHLIGHTS
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- 3 BUSINESS REVIEW
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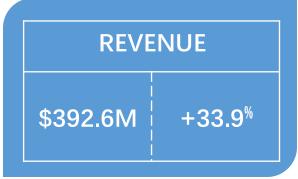
1 INTERIM RESULTS HIGHLIGHTS

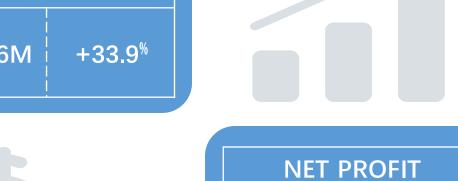
- Financial Highlights
- Business Highlights
- Product and Pipeline Progress

• Revenue Growth Rates and Historic Revenue Numbers Adjusted to Exclude Foreign Exchange Impact

(USD: Million)











INTERIM RESULTS HIGHLIGHTS (CONT'D)

FINANCIAL HIGHLIGHTS

- (USD: Million)
- Revenue: \$392.6m, **33.9%** YOY[↑], mainly attributable to the consolidation of CRM and the robust growth of key segments and core products, slightly offset by short-term fluctuation in non-China Ortho
- Consolidation of CRM for full six months and vigorous growth driven by domestic-made pacemakers led to 162.5% YOY ↑ for CRM segment
- Cardio: 27.8% YOY ↑, domestic Firehawk™ 59.6% YOY ↑, overseas Firehawk™
 52.2% YOY ↑, global Firebird2™ 8.0% YOY ↑
- China Ortho: 53.3% YOY ↑, knee products 45.4% YOY ↑, hip products 78.0%
 YOY ↑
- Endo: 41.6% YOY ↑; Neuro: 57.9% YOY ↑
- Non-China Ortho: 8.9% YOY ↓
- □ Gross Profit: \$281.6m, 28.4% YOY ↑ and GP Margin of 71.7%, increased by 90 bps, mainly due to
- Optimization of product sales structure
- □ Operating cost: \$249.4m, 41.7% YOY ↑, mainly due to
- The acquisition of CRM business
- The increased investments in the on-going and newly established R&D projects
- Increase in sales promotion and post-launching clinical trials
- One-time gain of \$55.8m (net of tax) from the disposal of partial equity interests in EP

NET PROFIT

attributable to equity shareholders

\$65.5M **175.5%** YOY¹

BUSINESS HIGHLIGHTS

- □ Substantial progress in R&D and pipeline development in 2019 1H
- **7** products obtained NMPA* approvals: AspirationTM, SoSuperiorTM, Firehawk NovaTM, MinosTM, ARBORESTM kyphoplastic Balloon Catheter, etc.
- 18 products obtained 25 registration approvals from overseas
- 1 product granted Green Path for NMPA approval: 3D Electronic Laparoscope
- Firehawk™ TARGET AC trial two-year data further proved that it can achieve identical clinical efficacy and safety with the first-in class DES and received reimbursement in France
- **Firesorb**® enrolled>300 patients in FUTURE II and completed 3-year follow-up of FIM, which proved that the Firesorb® based on PLLA is one of the viable, safe and effective solutions to the patients with single de novo coronary artery lesions as compared with the 1st generation biodegradable scaffolds
- VitaFlow® obtained NMPA approval and production license in July 2019 and completed its first implantation on Aug 2019
- VitaFlow® showed low all-cause mortality and no major stroke in 2-year follow-up study
- VitaFlow®II conducted multiple clinical trials in China and EU
- **BonaFire MRI conditional passive lead** finalized the protocol of K-B MRI clinical study
- The laparoscopic surgical instrument control system initiated the type testing process

Endovastec™ successfully launched its initial public offering on the Science and Technology Innovation Board of the Shanghai Stock Exchange on July 22, 2019 Endovaste

INTERIM RESULTS HIGHLIGHTS (CONT'D) Product and Pipeline Progress

CARDIO

- FirehawkTM NOVA
- Firehawk LibertyTM, WALTZ CoCr Coronary Stent System, Firefighter PTCA Balloon Dilatation Catheter, Firefighter NC PTCA Balloon Catheter, Y-connector, Inflation Device
- Two-Year data from Target AC trial of FirehawkTM showed non-inferiority to the internationally recognized stent
- FiresorbTM completed 3-year follow-up of FIM clinical trial and FUTURE II trial enrolled over 300 patients

CRM

- BonaFire MRI conditional passive lead finalized the protocol of K-B MRI clinical study
- NavigoTM left ventricular pacing leads entered clinical validation phase

ORTHO

- Domestically made AspirationTM
 Medial Stability Total Knee
 Replacement System
- Domestically made SoSuperiorTM Medial Stability Total Knee Replacement System,
- Domestically made femoral part
- ARBORESTM kyphoplastic Balloon Catheter
- BIOLOX® Delta Options system, Slo-Con Total Knee instruments, PRIME® Acetabular system, Knee Tensioner Instrument system, Evolution® NitrXTM Medial-Pivot Knee, Evolution® CS Stemmed Femur

NEURO

- Fastrack Microcatheter System
- Vertebral artery stent has been grant NMPA deficiency letter

ENDO

- Minos™ Ultra Low Profile AAA Stent-Graft
- Talos[™] Thoracic Stent-Graft System entered the phase of follow-up study of clinical trials
- Fontus™ Branched Surgical Stent Graft System entered the phase of follow-up study of clinical trials
- Reewarm[™] PTX Drug Coated Balloon passed panel review

Heart Valve

■ VitaFlow® Transcather Arotic Valve and Delivery System

VitaFlow[®] II conducted multiple clinical trials in China and EU

Surgical Robot

- 3D Electronic Laparoscope
- The laparoscopic surgical instrument control system has initiated the type testing process













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2 FINANCIAL REVIEW

- Consolidated Financial Performance
- Operating Expenses

• Revenue Growth Rates and Historic Revenue Numbers Adjusted to Exclude Foreign Exchange Impact

FINANCIAL REVIEW – Consolidated Financial Performance

Revenue

(USD: Million)

	non-CRM Business	CRM Business	Total	Growth
2018 1H	266.1	43.8	309.9	22.0%
2019 1H	286.0	106.6	392.6	33.9%

Operating Expenses

(USD: Million)

	non-CRM Business	CRM Business	Total	Growth
2018 1H	149.5	26.5	176.0	41 7 0/
2019 1H	165.5	83.9	249.4	41.7%

Net Profit attributable to equity shareholders

(USD: Million)

	non-CRM Business	CRM Business	Total	Growth
2018 1H	26.6	(2.8)	23.8	17F F0/
2019 1H	78.4	(12.9)	65.5	175.5%

Gross Profit Margin

(bps)

	GPM	Growth by bps
2018 1H	70.8%	00
2019 1H	71.7%	90

FINANCIAL REVIEW – Operating Expenses

Sales and Marketing Expenses

(USD: Million)

	2018 1H	20191H
Non-CRM Business	76.1	79.2
CRM Business	15.8	47.3
Total	91.9	126.5
S&M% of Sales	29.7%	32.2%

- Sales & Marketing expenses increased by 34.6m, 37.6% YOY ↑
- The acquisition of CRM business
- Increase in sales promotion & postlaunching clinical trials expenses
- Increase in staff cost

Administrative Expenses

(USD: Million)

	2018 1H	20191H
Non-CRM Business	38.7	43.3
CRM Business	3.6	11.7
Total	42.3	55.0
Ad% of Sales	13.6%	14.0%

- Administrative expenses increased by 12.7m, 30.0% YOY ↑
- The acquisition of CRM business
- Increase in staff cost

Research and Development Expenses

(USD: Million)

	2018 1H	20191H
Non-CRM Business	34.7	43.0
CRM Business	7.1	25.0
Total	41.8	68.0
R&D% of Sales	13.5%	17.3%

- Research & Development expenses increased by 26.2m, 62.6% YOY ↑
- The acquisition of CRM business
- Increased investments in the ongoing and newly established R&D projects

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3 BUSINESS REVIEW

- Cardiovascular
- Orthopedics
- CRM
- Endovascular
- Neurovascular
- Heart Valve & Surgical Robot
- Revenue Growth Rates and Historic Revenue Numbers Adjusted to Exclude Foreign Exchange Impact

BUSINESS REVIEW – Cardiovascular

2019 1H Revenue

(USD: Million)

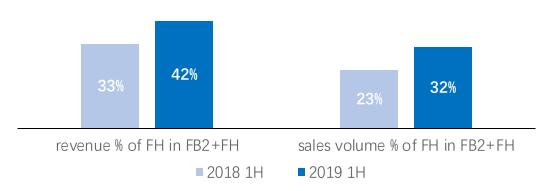
Global: 27.8% †; Domestic: 25.7% †; Overseas: 66.0% †



Highlights on Sales

- □ Global revenue: \$129.1m, 27.8% YOY ↑
- □ Domestic revenue: \$120.4m, strong growth of 25.7% YOY ↑, driven by:
- Firehawk[™]: 59.6% YOY ↑, Firebird2[™]: 6.6% YOY ↑; Balloon Products: 56.6% YOY ↑
- Domestic Hospital Coverage:
- Firehawk[™] hospital coverage 44.0% YOY ↑; Firebird2[™] hospital coverage 16.2% YOY ↑; Balloon Products hospital coverage 13.2% YOY ↑
- □ Overseas revenue: \$8.6m, 66.0% YOY ↑
- Obtained 14 registration approvals from 6 countries/regions during 2019 1H
- FirehawkTM received approvals in 3 additional countries
- FirehawkTM was included in French medical insurance reimbursement lists and completed first implantation in July in France
- Sales covered 29 countries/regions in total

Percentage of FirehawkTM in DES Sales



Highlights on Products

- □ Stent Products: currently, 4 stents in sales portfolio and 1 stents under R&D
- Firehawk LibertyTM obtained CE certification in March
- Firehawk NovaTM received NMPA approval in May
- **FirehawkTM** 's results of Target AC trial at 24 months further proved it can achieve identical clinical efficacy and safety as the internationally recognized stent
- **Firesorb**® completed 3-year follow-up of FIM clinical trial proved that it is one of the viable safe and effective solutions for the single de novo coronary artery lesions patients as compared with the first generation biodegradable scaffolds
- Balloon Products: currently, 3 balloon catheters being sold and 1 balloon catheter was newly approved
- FirefighterTM NC PTCA Balloon Catheter obtained CE certification in March

BUSINESS REVIEW – Orthopedics Non-China

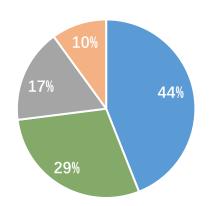
2019 1H Revenue

(USD: Million)



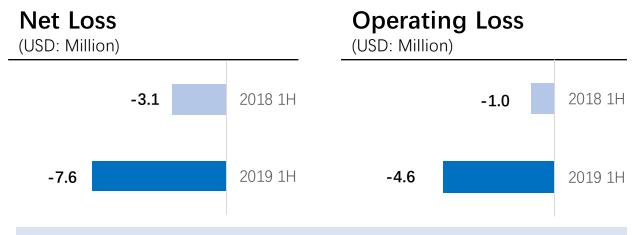
2019 1H Revenue by Geographic Areas

(USD: Million)



	Revenue	Growth
US	45	-11.5%
EMEA	29	-6.5%
Japan	18	3.9%
Others	10	-21.2%





- Revenue: \$101.9m, 8.9% YOY ↓
- US: 11.5% YOY \downarrow , mainly due to the loss of a large distributor in 2018 Q3
- Japan showed continuing solid growth of 3.9% YOY ↑, driven by a focus on sales execution and customer development
- EMEA: 6.5% YOY \downarrow , mainly due to the decrease of direct sales from subsidiaries
- □ Net Loss: -\$7.6m, expanded by \$4.5m YOY

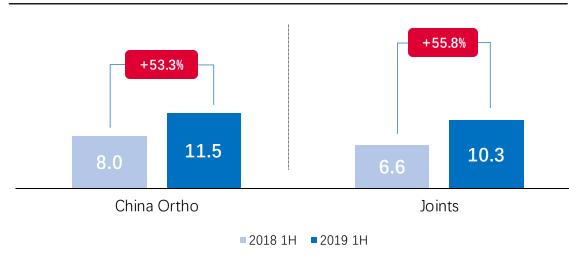
Progress in Restoration and future product development

- Progress in restoring sales :
- Added 4 new distributors representing a significant potential annual incremental surgical cases in US
- Progress in Registration:
- US: Evolution® NitrXTM Medial-Pivot Knee & Evolution® CS Stemmed Femur (both approved in Canada as well) & Knee Tensioner Instrument system
- Japan: PRIME® Acetabular system & Slo-Con Total Knee instruments
- EU: BIOLOX® Delta Options system
- Successful new product campaigns
- Launch of the Evolution® CS Stemmed Femur in March 2019, which will support the expansion of Evolution® Medial-Pivot Knee system

BUSINESS REVIEW – Orthopedics China

Revenue and Growth

(USD: Million)



Progress in China

New Hospital Penetration

SuperPath™	36
Hip Products	81
Advance®	49
Evolution ®	22



China Business Highlights

- □ Revenue: \$11.5m, 53.3% YOY ↑
- Revenue of joints: \$10.3m, 55.8% YOY ↑, driven by:
- Revenue of imported Knee products increased by 45% YOY 1
- Revenue of imported Hip products increased by 78% YOY ↑
- SuperPathTM covered over 100 hospitals with surgical cases increased by 9% YOY ↑
- Revenue of Spine and Trauma: \$1.1m, 18.4% YOY ↑, driven by:
- Launch of new products
- Orderly progress in market admittance in various provinces
- Surgical Instrument
- Developed 90 units of instruments and produced 20 sets of domestically designed ICE Evolution knee instruments
- Global Supply Center ("GSC")
- Processed over 60,000 product delivery in 33 countries/regions

Development on product portfolio

- Domestically made AspirationTM and SoSuperiorTM Medial Stability Total Knee Replacement System gained NMPA approvals and completed their first procedures respectively
- Domestically made Total Hip Replacement System is expected to gain NMPA approval by the end of 2019
- Improved product portfolio in joints will elevate the competitiveness of the China Ortho business

BUSINESS REVIEW – CRM

Product Launch for non-China Business



Non-China Business Highlights

- □ Revenue: \$103.1m, 153.1% YOY[↑], mainly due to
- Consolidation of CRM non-China business for full six months
- New product launch:
- ENOTM, TEOTM and OTOTM, world's smallest 1.5T & 3T MRI Compatible Pacemakers with a volume of 8cc, contributed more than 50% MicroPort CRM pacemakers implants in EU
- Lightweight, modular tablet-based programmer SmartTouchTM with Bluetooth[®] capability
- Business in Japan: sales team is ready to continue serving our customers and introduce our product
- □ Projects under R&D to enhance competitiveness
- InvictaTM defibrillation lead, NavigoTM left ventricular pacing leads and new pacemaker platform with Bluetooth[®] communication capability

2019 1H China Business Revenue

(USD: Million)



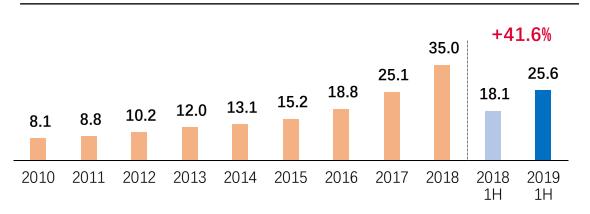
China Business Highlights

- Revenue: \$3.5m, 269.0% YOY[↑]. Significant growth driven by:
- Consolidation of CRM China business for full six months
- Domestically made pacemaker RegaTM family contributed 47% of China business revenue
- Continuous growth from imported pacemakers of 27% vs 6 months of 2018
- Hospital coverage and distribution channel
- Pacemakers covered more than 250 hospitals, 88% YOY 1
- Newly developed over 20 distributors
- Implantation update
- Over 900 implants of RegaTM family pacemakers achieved in 2019 1H
- Over 350 BEFLEXTM pacing lead implants
- Projects under R&D led to "Innovated in China"
- The protocol of K-B MRI clinical study for BonaFire MRI conditional passive lead has been finalized

BUSINESS REVIEW – Endovascular

2019 1H Revenue

(USD: Million)



Extensive Product Pipeline

	2019 A&E	2020E	2021E	Green Path
Minos™ Ultra Low Profile AAA Stent-Graft	✓			Mar 2017
Reewarm™ PTX Drug Coated Balloon	✓			Dec 2015
Fontus™ Branched Surgical Stent Graft System		✓		Aug 2018
Talos™ Thoracic Stent-Graft System			✓	Sep 2017

Endovastec[™] went public on the Science and Technology Innovation Board of Shanghai Stock Exchange on July 22, 2019





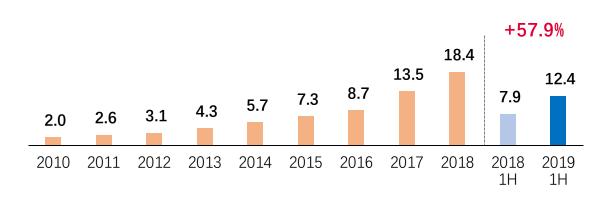
Highlights

- Revenue: \$25.6m, 41.6% YOY[↑], driven by:
- Fast-growing Chinese market with CGAR over 15%
- Castor[™], the world's first thoracic branch stent-graft system, maintained rapid growth, completed over 1000 implants cumulatively
- Continuous focuses and effective promotion in tier 2&3 cities
- Penetrated additional 36 hospitals
- New catalyst for sustainable growth
- Minos™ Ultra Low Profile AAA Stent-Graft received NMPA approval in March
- Reewarm™ PTX Drug Coated Balloon expected to obtain NMPA approval by end of 2019

BUSINESS REVIEW – Neurovascular

2019 1H Revenue

(USD: Million)

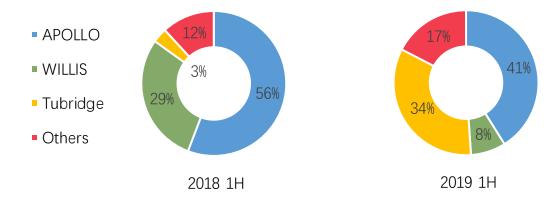


Extensive Product Pipeline

	2019	2020E	Green Path
Vertebral artery stent		✓	Mar 2018
Coils		✓	
Clot Retrieval Device		✓	
Fastrack Microcatheter system	√		

Revenue by Products

(USD: Million)



Highlights

- Revenue: \$12.4m, 57.9% YOY ↑, mainly due to:
- Significant contribution from Tubridge™, representing 33.7% of the revenue
- Revenue decrease of WILLS[™] partly due to launch of competing product and substitution effect of Tubridge[™]
- Hospital Coverage
- APOLLO™: newly penetrated 102 hospitals
- Tubridge™: newly penetrated 65 hospitals
- Registration and clinical progress
- Fastrack Microcatheter system gained NMPA approval in August 2019
- Clinical trials for coils has completed and showed promising results

BUSINESS REVIEW – Heart Valve & Surgical Robot

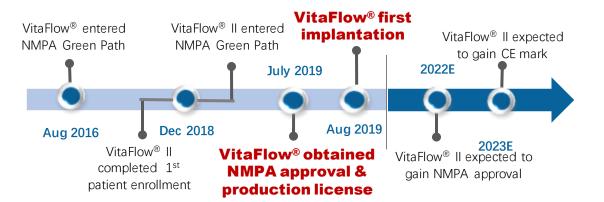
Heart Valve

VitaFlow[®]

- VitaFlow® is designed to provide solution for aortic valve stenosis and has demonstrated safety and effectiveness in treating severe calcified aortic stenosis
- Low all-cause mortality and no major stroke in 2-year follow-up study
- VitaFlow® received NMPA approval & production license in July 2019 and completed its first implantation on Aug 28, with a total of 4 implantations completed on the same day

VitaFlow® II

- □ VitaFlow® II is equipped with retrievable delivery system
- "Retrievable" feature will provide solution to the challenging positioning issue, thereby improving precision and success rate
- While achieving the retrievable feature, VitaFlow® II maintains its remarkable deployment stability and ability in preventing PVL
- Registration and clinical progress
- VitaFlow® II has entered NMPA Green Path in December 2018
- Expected to gain NMPA approval by 2022 and CE mark by 2023
- Clinical trials conducted in China and EU



Surgical Robot

Solid progress in R&D

- 3D Electrical Laparoscope
- The Company's self-developed 3D Electrical Laparoscope was granted NMPA Green Path, becoming the Company's 16th product entering the NMPA Green Path
- The 3D Electrical Laparoscope provides high-resolution instant images which help doctors to conduct minimally invasive surgeries
- Since there is no domestic 3D laparoscopic product currently in China, the grant of Green Path will assist in popularizing cutting -edge technology and boost the development of connected industries
- Laparoscopic Surgical Robotic Control System
- The laparoscopic surgical instrument control system has initiated the type testing process





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4 OUTLOOK

- Continuous Product Pipeline Fueling Long-term Growth

NMPA

OUTLOOK -**Continuous Product Pipeline Fueling Long-term Growth**

Cardio	Firefighter™ NC NMPA & CE	Firebird2™ Nova NMPA & CE	Firesorb™ NMPA	Fantasy Drug-Eluting Stent NMPA & CE	
Ortho	Domestically made hip system NMPA	Domestically made spine system NMPA	Domestically made trauma system NMPA		
CRM	Navigo™ left ventricular pacing leads CE	Kora100™ pacemaker family NMPA	Pacemaker platform with Bluetooth CE	Invicta TM defibrillation lead CE	TPG (external temporary pacemaker) NMPA
Endo	Reewarm™ PTX Drug Coated Balloon NMPA	Fontus ™ Branched Surgical Stent Graft System NMPA	Talos™ Thoracic Stent-Graft System NMPA		
Neuro	Vertebral Artery Stent NMPA	Coils NMPA	Clot Retrieval Device NMPA	-	•
Heart Valve	VitaFlow TM II NMPA &CE				
Surgical Robot	Robot for joint operation NMPA	Robot for endoscope operation NMPA	3D Electronic Laparoscope system II NMPA		

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5 APPENDIX: FINANCIAL STATEMENTS

- Consolidated Income Statement
- Consolidated Balance Sheet

APPENDIX: FINANCIAL STATEMENTS - Consolidated Income Statement

Unit: USD'000	1H 2019	1H 2018	Var.
Revenue	392,607	309,867	27%
Cost of sales	(110,969)	(90,438)	23%
Gross profit	281,638	219,429	28%
Other net income	8,613	4,274	102%
Research and development costs	(67,968)	(41,791)	63%
Distribution cost	(126,465)	(91,902)	38%
Administrative expenses	(54,974)	(42,291)	30%
Other operating costs	(5,860)	(7,942)	-26%
Profit from operations	34,984	39,777	-12%
Finance cost	(9,560)	(8,708)	10%
Gain on disposal of subsidiaries	63,105	-	n.a
Gain on deemed disposal of a joint venture	-	4,133	-100%
Share of losses of associates	(1,318)	(848)	55%
Share of losses of a joint venture	0	(202)	-100%
Profit before taxation	87,211	34,152	155%
Income tax	(26,362)	(9,949)	165%
Profit for the period	60,849	24,203	151%
Attributable to: Equity shareholders of the Company	65,476	23,769	175%

APPENDIX: FINANCIAL STATEMENTS - Consolidated Balance Sheet

Unit: USD'000	30 June. 2019	31 Dec. 2018	Var.
Non-current assets			
Investment properties	7,470	5,451	37%
Other property, plant and equipment	349,463	336,263	4%
Right-of-use assets	46,274	_	n.a
Land use right	14,708	15,087	-3%
Intangible assets	118,100	117,489	1%
Prepayments for non-current assets	10,122	6,222	63%
Goodwill	162,105	162,673	0%
Interest in associates	50,391	12,291	310%
Interest in a joint venture	5,100	5,100	0%
Other financial assets	18,314	11,910	54%
Deferred tax assets	14,506	15,291	-5%
Other non-current assets	41,374	31,979	29%
Total non-current assets	837,927	719,756	16%
Current assets			
Inventories	182,956	175,957	4%
Trade and other receivables	250,892	245,143	2%
Pledged deposits and time deposits	2,032	3,537	-43%
Cash and cash equivalents	95,374	130,054	-27%
Derivative financial assets			
Total current assets	531,254	554,691	-4%
Current liabilities			
Trade and other payables	221,943	236,813	-6%
Contract liabilities	8,754	10,060	-13%
Lease liabilities	9,824	_	n.a
Interest-bearing borrowings	92,751	100,901	-8%
Covertible bonds	86,707	86,834	0%
Income tax payable	17,908	5,782	210%
Total current liabilities	437,887	440,390	-1%
Net current assets	93,367	114,301	-18%

APPENDIX: FINANCIAL STATEMENTS - Consolidated Balance Sheet (cont' d)

Unit: USD'000	30 June. 2019	31 Dec. 2018	Var.
Non-current liabilities			
Interest-bearing borrowings	145,680	137,829	6%
Lease liabilities	37,618	-	n.a
Deferred income	18,465	23,905	-23%
Convertible bonds	4,533	3,571	27%
Contract liabilities	28,069	27,766	1%
Other payables	88,050	84,819	4%
Net defined benefit obligation	8,778	8,806	0%
Deferred tax liabilities	7,513	7,775	-3%
Financial liabilities carried at fair value	13,000	10,640	22%
Total non-current liabilities	351,706	305,111	15%
CAPITAL AND RESERVES			
Share Capital	16	16	0%
Reserves	498,330	442,780	13%
Total equity attributable to equity shareholders of the Comp	498,346	442,796	13%
Non-controlling interests	81,242	86,150_	-6%
TOTAL EQUITY	579,588	528,946	10%

