



ASX:MVP

FY23 Full Year Results

31 August 2023



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- difficulties or delays in manufacturing;
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Non-IFRS Financial Information

This presentation uses non-IFRS information including underlying revenue, underlying EBIT and underlying adjustments. These measures are key performance measures used by MDI, the investment community, and peers with similar business portfolios. MDI uses these measures for its internal management reporting as it better reflects what MDI considers to be its underlying performance. Underlying revenue and EBIT are used to measure segment performance and have been extracted from the segment information disclosed in the Consolidated Full Year Financial Report.



FY23 Full Year

Key messages

Robust growth for Penthrox and Respiratory in all markets

group revenue up 47%

Gross margin improved

\$5.4 million delivered from pricing and volume growth in FY23

Primary investment phase complete

driving delivery of growth strategy

Strong momentum toward positive operating cashflow in FY25

continued growth in key segments and \$6 million from pricing and efficiency expected in FY24

US market entry advancing

commercial opportunity for Penthrox of US\$300-\$400 million identified through market assessment; partner search commenced



FY23 Full Year Financial highlights

Revenue

\$32.3m¹

+47%

Pain Management Revenue

\$20.4m¹

+54%

Respiratory Revenue

\$11.7m

+43%

Underlying EBIT

\$18.3m (loss)

(pcp \$14.7m loss)

Underlying Adjustments

\$10.3m² (gain)

(pcp \$1.2m loss before tax)

NPAT

\$5.6m (loss)

(pcp \$12.4m loss)

Operational highlights

Commercial milestones

- Expansion of Pentrox into Australian hospital emergency departments
- Strong growth for Pentrox in key partner markets
- Spend in France scaled back in light of market conditions
- Market share gains in the Respiratory segment

Market registrations and product development

- Clinical trial in China discontinued
- UK paediatric trial closed; submission expected Q3 FY24
- Funding award of \$1.5 million to support development of next generation inhaler
- US partner search underway; commercial market assessment complete



FINANCIAL PERFORMANCE

Earnings summary

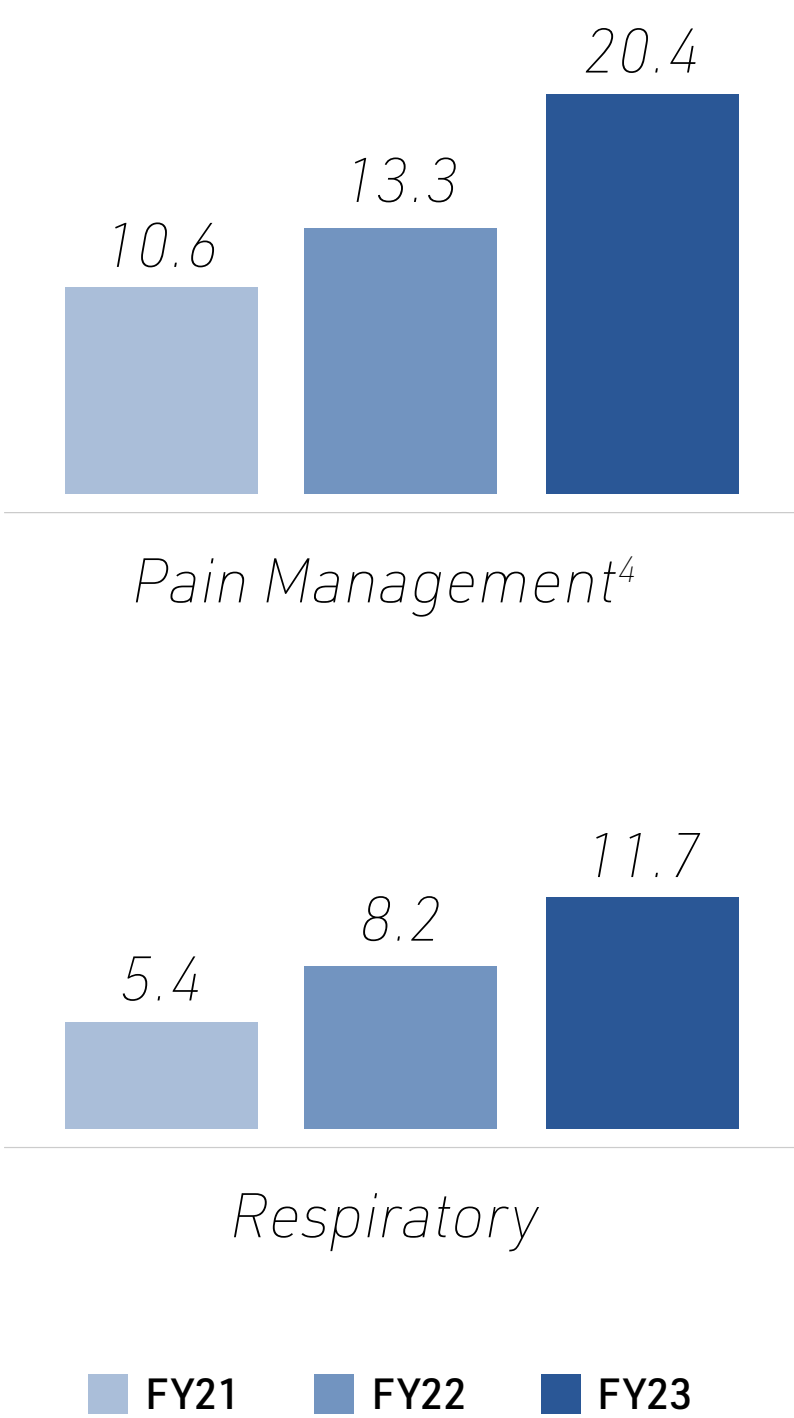
FY23 full year results

\$million	FY22	FY23	Change \$m
Revenue ¹	21.9	32.3	10.4
Underlying EBIT	(14.7)	(18.3)	(3.6)
Underlying Adjustments (before tax) ²	(1.2)	10.3	11.5
Reported EBIT	(15.9)	(8.0)	7.9
NPAT	(12.4)	(5.6)	6.8

Commentary

- Group revenue up 47%
- Higher volumes in key markets
- Higher prices
- Underlying EBIT lower, with higher costs from capability build
- Capability build complete, efficiencies to be delivered in future periods
- Non-cash Underlying Adjustments of \$10.3 million (gain before tax), mostly arising from cessation of clinical trial preparations in China, drive improvement in Reported EBIT and NPAT

Segment Revenue³
(\$million)



1. Excludes Contract termination revenue of \$18.9 million (refer to the Full Year Consolidated Financial Report).
 2. Underlying adjustments are detailed on page 22 in the Appendix.
 3. Excludes other segment revenues relating to discontinued businesses (FY23: \$0.2 million; FY22: \$0.4 million; FY21: \$0.4 million).

Pain Management segment revenue

Revenue up 54%, higher volumes and pricing

\$million	FY22	FY23	Change %
Europe	4.0	5.5	39%
Australia	7.4	9.6	30%
Rest of World	1.9	4.6	141%
Product revenue	13.3	19.7	49%
Milestone and other revenue ¹	-	0.7	-
Pain Management	13.3	20.4	54%

Commentary

Europe

- In-market volumes up 39% with growth in all countries
- UK and Ireland in-market volumes up 34% with encouraging momentum
- France up 33% against a challenging backdrop
- 1H23 deferred deliveries recovered in 2H23

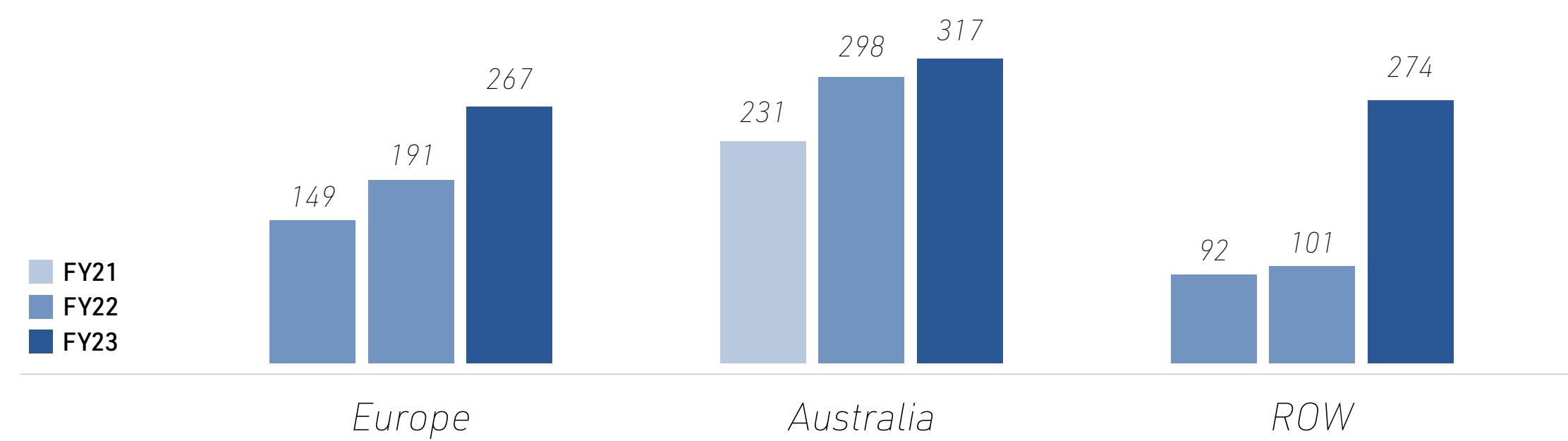
Australia

- Volumes up 6% with solid demand from ambulance
- Growing penetration in emergency departments
- Growth in procedural segments
- Prices increased from November 2022

Rest of World (ROW)

- Significant uplift from inventory stocking for relaunch of Pentrox in Canada
- Growth in Middle East, South Africa and Asia

Penthrox units² 000s



1. Excludes Contract termination revenue of \$18.9 million (refer to the Full Year Consolidated Financial Report).
2. European volumes reflect “in-market” sales units, which may differ from units sold to distribution partners in the period (and recognised in revenue). The Company believes this measure improves the transparency of underlying demand.

Respiratory segment revenue

Share gains drive revenues 43% higher

\$million	FY22	FY23	Change %
Europe	1.7	2.3	40%
Australia	3.2	3.8	19%
USA	2.9	4.6	59%
Rest of World	0.4	1.0	150%
Respiratory	8.2	11.7	43%

Commentary

Europe and Rest of World

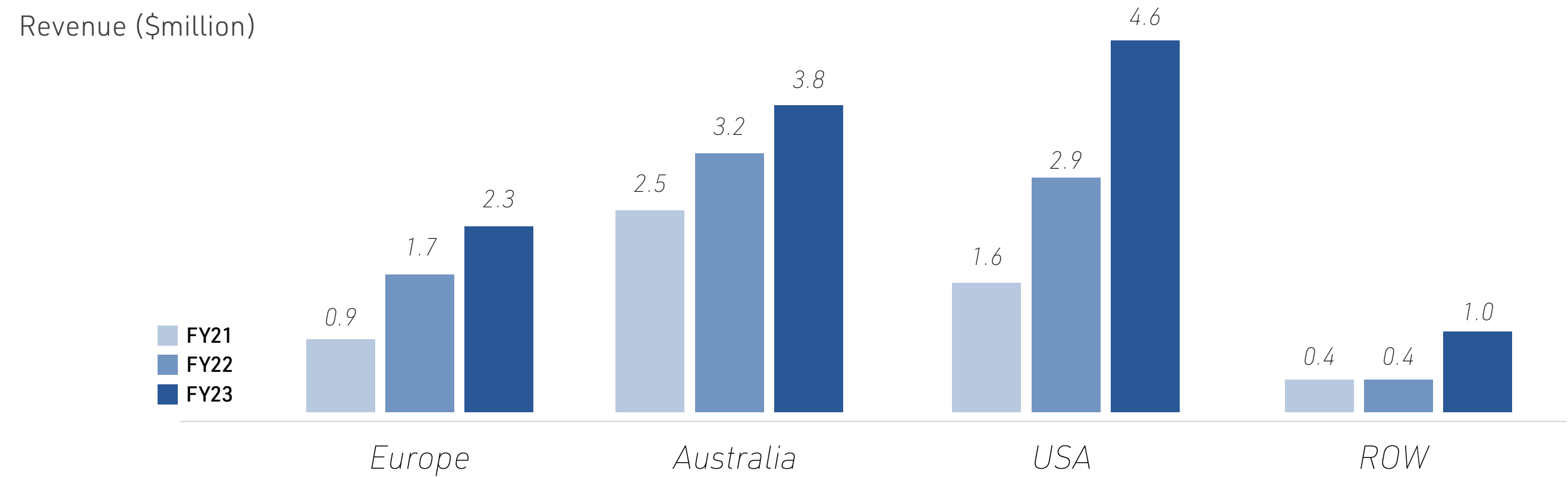
- Strong underlying demand
- Increased engagement with partners
- Prices increased

Australia

- Strong underlying demand
- Increased engagement with direct banner groups
- Penetration into hospital channel
- Prices increased

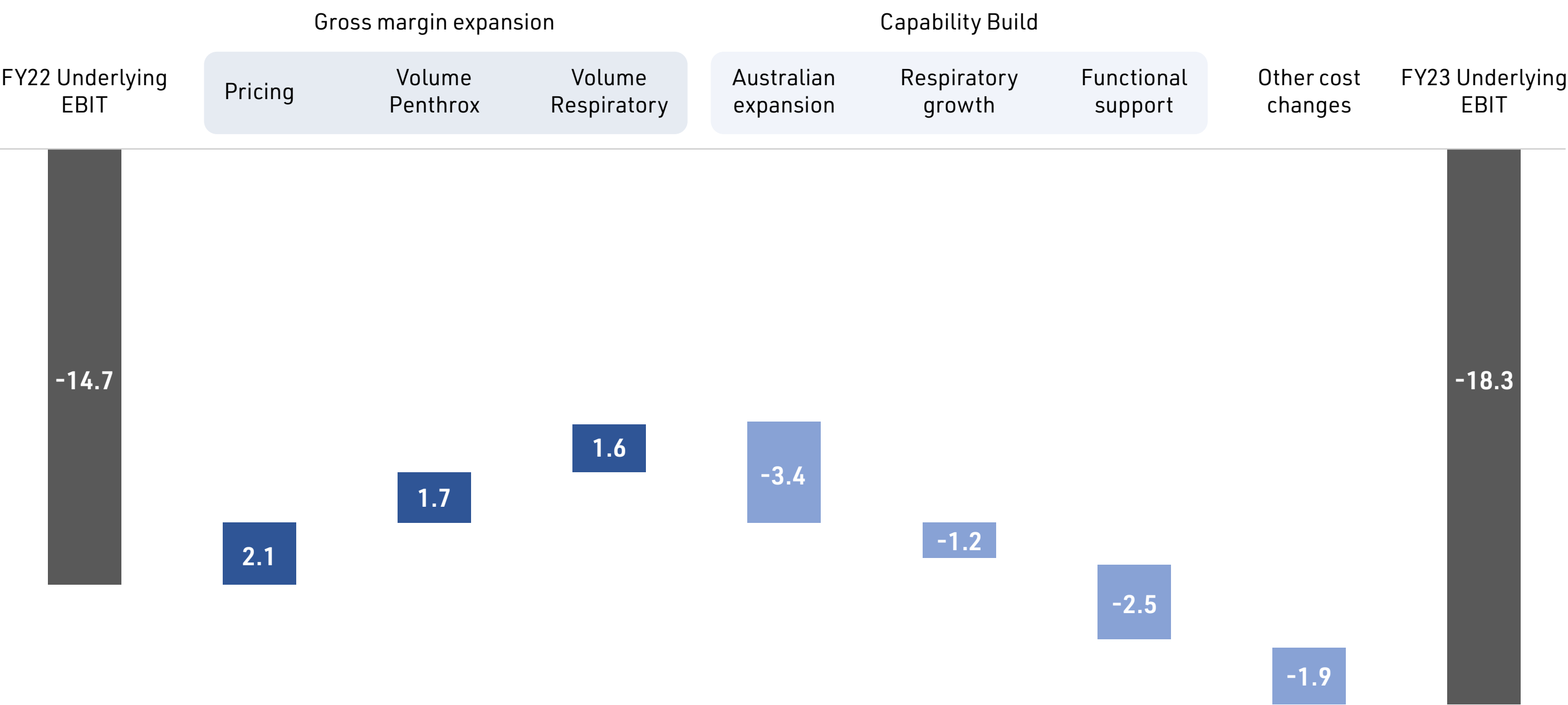
USA

- Market share growth driven by investment in third party infield resources and strong partner engagement
- Continued expansion into pharmacy banner/wholesaler groups



Volume and pricing drive margin expansion

Underlying EBIT bridge (\$millions)



Commentary

Pricing

- Stronger pricing in Pain Management and Respiratory segments, particularly for Pentrox in Australia

Volumes

- Growth in Pentrox volumes in all markets
- Share gains and strong underlying demand in Respiratory

Capability build

- Investment in Australian Pentrox field team
- Increased commercial resources in Respiratory
- Leadership and functional capability

Other costs

- Inflationary impacts
- Contract termination costs relating to scale down in France

Balance sheet and cashflow

Primary investment phase complete, improved cashflows expected

\$'000	FY22	FY23	Change \$m
Underlying EBITDA	(11.7)	(15.1)	(3.4)
Non-cash items	1.0	0.7	(0.3)
Change in working capital	(2.4)	(1.8)	0.6
Change in other assets and liabilities	0.1	(0.8)	(0.9)
Income tax received	2.3	-	(2.3)
Interest paid	(0.1)	(0.1)	0.0
Operating cash flow	(10.8)	(17.1)	(6.3)
Working capital / sales %	17%	17%	-
Capital expenditure	(5.2)	(7.7)	(2.5)
Cash	20.4	24.7	4.3

Commentary

Change in working capital

- Investment to support sales growth in both Pain Management and Respiratory segments
- Working capital to sales % in line with prior year

Capital expenditure

- Plant and equipment (\$1.8m), mostly relating to manufacturing projects
- Intangible assets (\$5.9m), mostly relating to development of the next generation inhaler ("Selfie"), the UK paediatric trial, and registration activities for US and China (China now discontinued)
- Capital expenditure in FY24 (including spend on trials and market registration activity) is expected to be approximately \$5 million

Cash

- Closing cash balance of \$24.7 million



FY24 PRIORITIES

FY24 priorities

Deliver improved earnings and cashflow and advance US market entry

1. Improve margins through pricing and operational efficiency
2. Increase penetration of Pentrox in Australian hospital emergency departments (EDs)
3. Review go-to-market model for Europe
4. Progress partner search and finalise pathway for US market entry
5. Drive continued growth in Respiratory



FY24 priorities

Improve margins through pricing and efficiency

Incremental benefits of \$6 million expected in FY24

Pricing benefits

- Higher pricing in Australia
- Opportunity to reset commercial arrangements in other markets

Cost efficiencies

- European restructuring
- Supply chain efficiency
- Scale efficiencies

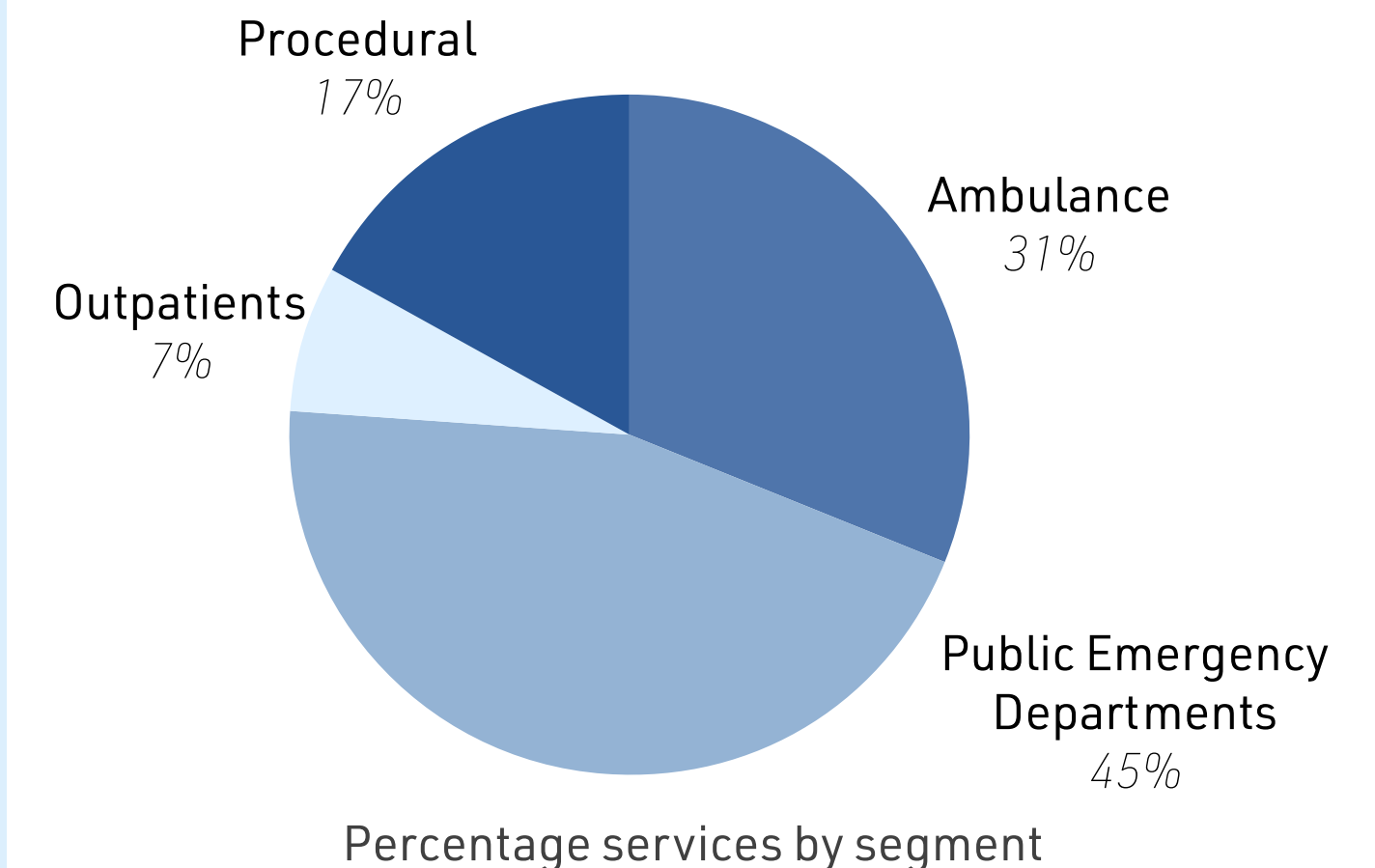


Increase penetration of Penthrox in Australian hospital emergency departments

National Account Management Team delivering encouraging progress

- Volume in Australia up 6% in FY23
- Progressing inclusion of Penthrox on public hospital formularies
- Expanded use of Penthrox in private hospitals
- Growth in procedural segment being realised

Attractive market opportunity¹



Opportunity to broaden market penetration

- ~75% of Penthrox volumes currently from Ambulance segment
- Significant opportunity to grow in emergency departments and procedural

FY24 priorities

Review go-to-market model for Europe

Realise potential of Penthrox in key European markets through a more cost-effective approach

- In-market field team in France scaled back, service to existing customers remains a priority
- Leverage attractive position established in France (annual demand ~65,000 units, >300 customers)
- Progress update to be provided at the AGM in October 2023



FY24 priorities

Progress US partner search and finalise pathway for market entry

Commencement of clinical pathway expected in FY25



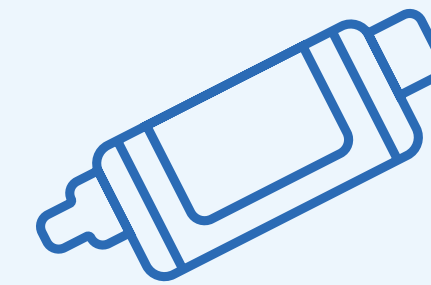
Funding strategy

US adviser appointed to support search for partner(s)



Clinical pathway

Development of clinical pathway advancing



Commercialisation

Development of next generation device ("Selfie") on track, expected to be used in clinical trial and commercial launch



Commercial market assessment

Annual in-market revenue potential for Pentrox of US\$300-US\$400 million¹

Large addressable market

- 20 million adult patients² in acute pain and trauma, 75% in emergency departments (ED) and emergency medical services (EMS) segments
- Further 10 million adult patients² in outpatient procedures, reached with label expansion
- Paediatric opportunity with label expansion

Attractive pricing potential

- Estimated in-market pricing between US\$35-US\$75 per unit
- Urgent care centres most price sensitive, ED least price sensitive



FY24 OUTLOOK

FY24 outlook

The Company expects underlying EBIT in FY24 to improve on the prior year, driven by:

- Higher Pentrox volumes in Australian hospital emergency departments;
- Share growth in the Respiratory segment; and
- Incremental margin improvements of \$6 million from pricing and efficiency.

Capital expenditure in FY24 (including spend on trials and market registration activity) is expected to be approximately \$5 million.





Appendix

Reconciliation between underlying EBITDA and net loss after tax

\$million	FY22	FY23
Underlying EBITDA	(11.7)	(15.1)
Depreciation and amortisation expense	(3.0)	(3.2)
Underlying EBIT	(14.7)	(18.3)
Contract termination revenue - Pain Management segment ¹	-	18.9
Impairment losses - Capitalised Registration Costs ²	-	(6.7)
Commercial Market Assessment Costs ³	-	(1.9)
Impairment losses - Veterinary segment ⁴	(0.6)	-
Finalisation of costs for the CSIRO Continuous Flow technology program ⁵	(0.6)	-
Total underlying adjustments	(1.2)	10.3
Reported EBIT	(15.9)	(8.0)
Net interest	-	0.5
Net loss before tax	(15.9)	(7.5)
Income tax benefit	3.5	1.9
Net loss after tax	(12.4)	(5.6)

Notes

FY23

1. Contract termination revenue arising from the termination of agreements for the distribution of Pentrox in China (\$18.5 million), and other countries where revenue opportunities are not being pursued (\$0.4 million).
2. Impairment of capitalised registration costs following the cessation of market activities in China of \$5.7 million, and an additional \$0.9 million in other countries where revenue opportunities are not being pursued. There was also a \$0.1 million impairment in relation to patents and trademarks.
3. Costs to complete a comprehensive assessment of the commercial potential for Pentrox in the US (\$1.9 million) which are not of a capital nature.

FY22

4. Impairment losses recognised following the Group's decision to discontinue the Veterinary business (\$0.6 million).
5. Finalisation costs for the CSIRO Continuous Flow technology program (\$0.6 million).

Company overview

- Medical Developments International Limited (ASX:MVP) is an Australian based pain management and respiratory company
- The Company's lead product, Pentrox (The Green Whistle®), an inhaled, needle-free, non opioid analgesic, is manufactured in Australia and sold globally
- The Company also has a portfolio of respiratory products for sufferers of asthma and COPD¹
- The Company's strategic focus is to accelerate penetration of Pentrox through direct in-market capability in Australia and Western Europe, and to grow its Respiratory segment through market share gains
- Longer term growth for Pentrox will be underpinned by entry into the USA market



Penthrox

Over 8 million used worldwide

The superior efficacy, safety and administration benefits of Penthrox deliver improved patient outcomes and lower overall costs

- Inhaled needle-free analgesic¹
- Non-opioid¹
- Portable, self administered device¹
- Effective pain relief within 6–10 breaths¹⁻⁴ and rapid offset
- Established safety profile with over 8 million uses
- Well tolerated, with the majority of adverse events mild and transient^{1,2}
- Approved for use in children in Australia¹
- Efficiency benefits of Penthrox in hospital emergency departments illustrated in British study⁵

The iconic *Green Whistle*



Helping patients manage asthma and COPD¹

Providing pharmacies, medical clinics and hospitals with a range of respiratory devices which are designed to assist patients to manage asthma and COPD¹

- Space chambers
- Portable nebulisers
- Silicon Face Masks





ASX:MVP

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