MVP Roadshow Presentation

July 2020





Who we are

- Pharmaceutical and devices company based in Victoria, Australia
- ASX Listed since 2003 (ASX: MVP)
- Market Capitalisation A\$440m
- 2 Key operating segments Pharmaceutical and Medical Devices
- Also have a significant API development project ongoing in partnership with the CSIRO
- Growing geographically
- Investing in product development



Penthrox®

A world class opportunity

Penthrox®

- Market Leader for trauma pain
- Opiate sparing, fast-acting inhalational analgesic
- 85% of patients reach clinical analgesia within 6-10 breaths
- Is a solution to a significant unmet clinical need
- Demonstrated safety and efficacy profile for 30+ years
- Manufactured in Australia









Coffey (2014)- STOP!: A Randomised, Double-blind, Placebo-controlled Study Of The Efficacy & Safety Of Methoxyflurane For The Treatment Of Acute Pain



Future for Penthrox® Australia Azerbaija Croatia Canada Greece Malta Germany Norway Australia New Zealand Hungary Croatia Liechtens Australia New Zealand New Zealand New Zealand Monaco Singapore Greece New Zealand Monaco Australia Singapore Singapore Saudi Arabi Ireland Malta Saudi Arai Australia New Zealand Moldova Moldova South Africa South Africa Norway New Zealand Australia Moldova Azerbaijan Hungary Australia Azerbaijan South Africa UAE Liechtenstein New Zealand Moldova Azerbaijan Georgia UAE New Zealand Georgia Azerbaijan Georgia Ukraine Moldova Ukraine Saudi Arabia Georgia Guatemala Guatemala Azerbaijan Ukraine Kazakhstan Vatican City South Africa Ukraine Portugal Guatemala Azerbaijan Kuwait Kazakhstan Georgia Oman South Korea South Korea Luxemburg Bahrain Czech Republic Lebanon Guatemala Zimbabwe Botswana Macedonia MVP continues to negotiate with interested Zimbabwe Botswana

MVP continues to negotiate with interested parties from around the world in terms of registering and selling Penthrox®, whilst concurrently pursuing other important international regulatory submissions and preparations in countries including USA, China, Russia, Iran, Iraq, Thailand and South Korea.



Penthrox® in USA

FDA Update

With FDA for review:

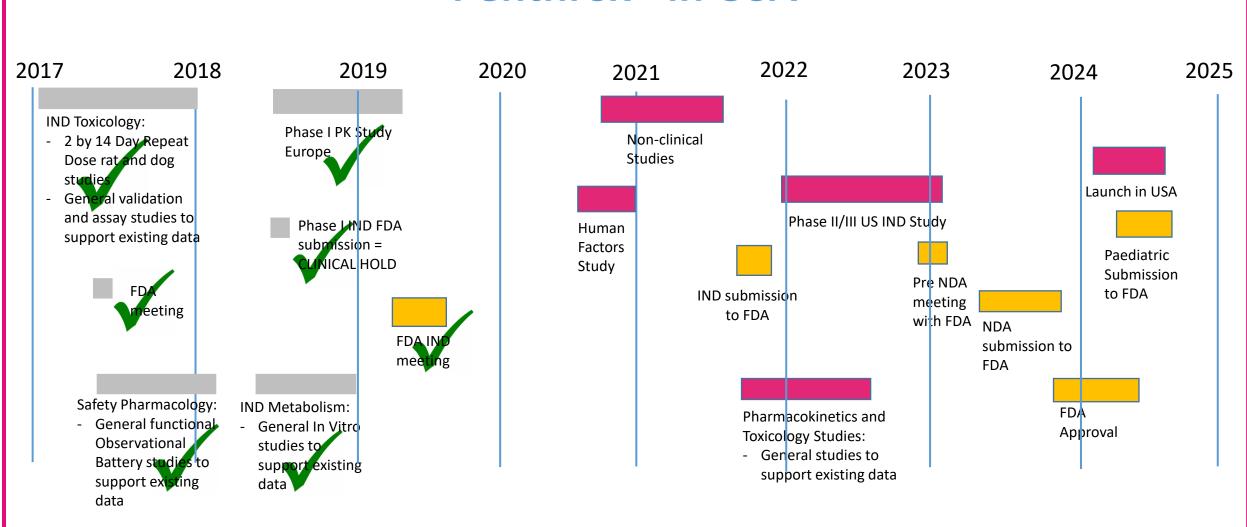
- Pre-Clinical protocol which mimics human dosing regimen
- CRO ready to commence trial once FDA response received
- Responses to device questions

Human Factors study protocol – ready to commence

Completion of PASS in Europe – awaiting final Clinical Study Report in Jun/Jul 2020 IND submission targeted for 2021.



Penthrox® in USA





Penthrox® in China

MVP concluded a significant transaction with Daiichi Sankyo for the Chinese, Thai and Vietnamese markets in 2018.

MVP received an upfront payment of circa \$21m for the commercialisation rights

The China IND was approved in November 2019

- Phase I PK
- Phase III Bridging Trauma
- Phase III Bridging Acute Pain

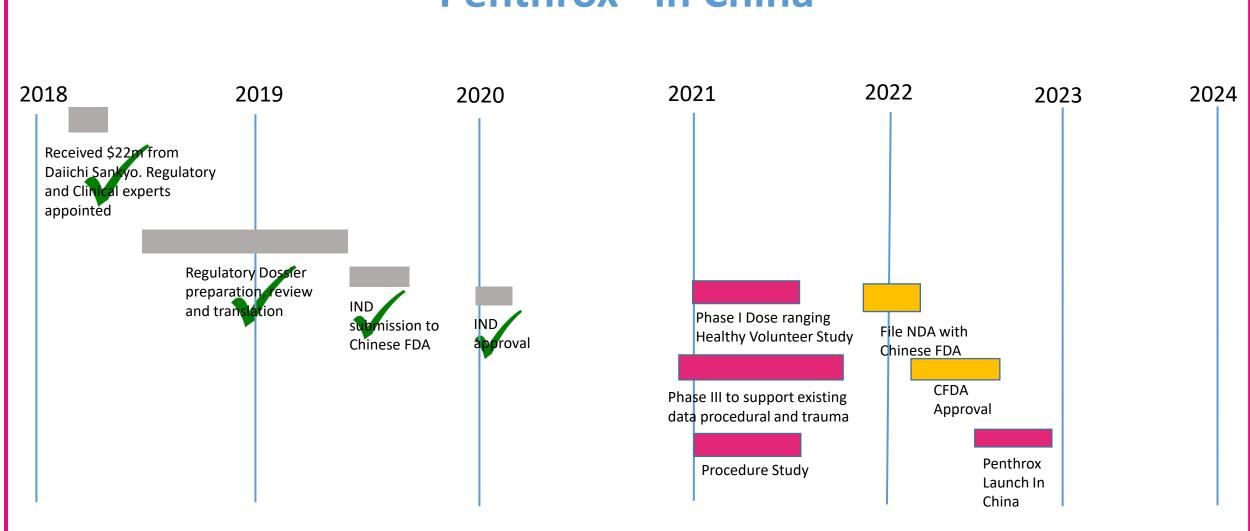
MVP site selections complete and protocol training provided to partner

Recruitment on hold due to COVID-19

NDA approval expected 2022



Penthrox® in China

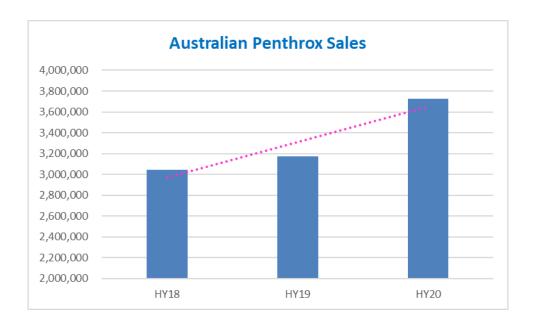




Financial Performance – H1 FY20

Penthrox® Sales in Australia

- In market" sales in Australia grew 18%
- New Mundipharma distribution agreement delivering strong sales growth in Australia particularly in GP/hospitals
- Sales for Australian 'Dr's Bags' up 54%

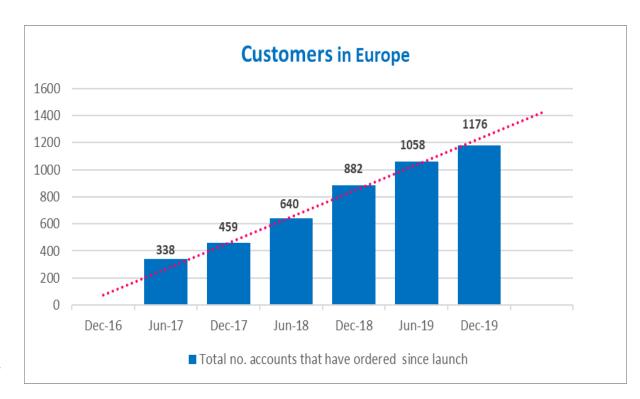




Financial Performance – H1 FY20

Penthrox® Sales in Europe

- UK in-market sales grew 42%
- 131 hospitals using in the UK
- Additional ambulance services targeted
- In-market European sales grew 35%
- 386 customers in France
- Launched in Italy, Nordics, Switzerland
- A number of major countries in Europe still to launch including Germany and Spain





Continuous Flow

Our own API - Methoxyflurane

Benefits of methoxyflurane manufacture under Continuous Flow vs Batch Process

- 1. Better overall process control
 - 40% better reaction conversion
 - 90% better overall yield conversion
 - Significantly reduced impurity profile
- 2. Fast scale-up capability
 - Increase output up x10+ on same footprint
- 3. Safer Environment
 - Reduced manual handling
 - Controlled energetic process (exotherm)
 - Waste Streams reduced



Continuous Flow

Future API technologies

MDI is developing its core flow technology into several generic APIs, currently manufactured under standard batch processing.

Examples of this are:

LIDOCAINE (USP): Estimated USD \$3.5 billion global sales

DICLOFENAC: Estimated USD \$6.0 billion global sales

SALBUTAMOL: Estimated USD \$6.0 billion global sales

• ISO/DES/SEVOFLURANE: Estimated USA \$3.0 billion global sales market. (Significant

improvements in handling highly toxic & corrosive Fluoride

intermediates under safe flow conditions)

• **LEFLUNOMIDE**: Early stage development under flow



Continuous Flow

Our Continuous Flow technology delivers:

- Increased yields through better process conversion
- Increased purity through better process control
- Better control over entire process test in real time
- Lower cost of production
- Lower CAPEX
- Less waste
- Less carbon footprint "greener technology"
- Smaller footprint
- Quicker to scale-up
- Safer



Global Respiratory – H1 FY20

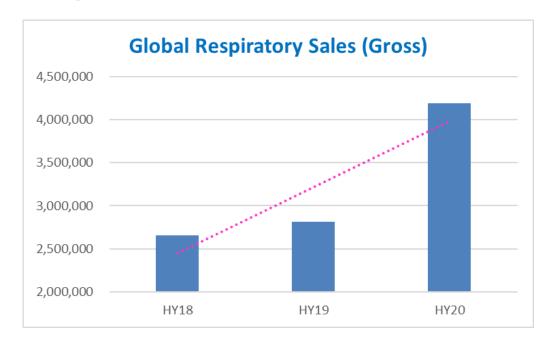
Global Respiratory Sales

Overall

- Global sales grew 49%
- Improved European sales
- Canada sales grew 207% strong cardboard spacer sales

Australia

- Sales growth in Australia of 44%
- Breath-A-Tech sales grew 31%





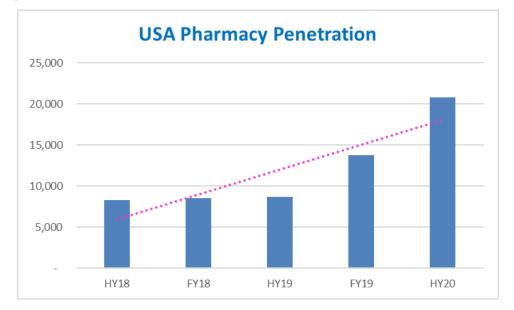


Respiratory - USA

Respiratory Sales in USA

- Sales to USA grew 49% now ranged in over 20,000 pharmacies
- Walgreens launched into 4,750 stores in Feb
- Private label launch into over 4,500 Walmart stores expected in Aug 2020















Outlook

MVP's ambition is to globalise Penthrox, and in doing so, make it the mainstream analgesic of choice around the world. This process has begun. Over the next 12 months+ we expect to:-

- begin selling Penthrox into numerous new markets New EU launches, Thailand, Mexico, etc;
- Continued expansion/growth in the UK/Ireland territory;
- progress work on gathering the clinical data needed to submit a "New Drug Application" to the Food & Drug Administration in the USA, including opening of the IND;
- Conclude the human trial associated with our China registration program; and
- Work toward extending the 'indications for use' for Penthrox.

Whilst at the same time:

- Continue to grow our respiratory business via new product launces and greater pharmacy reach in the USA; and
- Commercialisation of first Continuous Flow technology for Lidocaine.



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