Investor Presentation – AngelMed Guardian FDA Approval

Hydrix Limited (ASX: **HYD**) attaches an investor presentation in relation to the recent FDA approval of the AngelMed Guardian device, as announced to the market on 28 June 2021.

A recorded presentation given by Executive Chairman Gavin Coote, which provides a narrative to the Investor Presentation slides, can be accessed at the following link: Video link

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Authorisation: This announcement has been authorised by the Board of Hydrix Limited.

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About Hydrix Limited

Hydrix Limited (ASX: HYD) is a powerful product innovation company. Hydrix purpose is to enhance the health, safety, and wellbeing of one Billion lives. The company leverages its powerful product innovation capability across multiple growth platforms: Hydrix Services design and engineer client products which transform industries; Hydrix Ventures generate equity returns through investing in high potential companies; and Hydrix Medical bring innovative medical technologies to market.

Hydrix Limited

Hydrix brings technologies to life which improve people's health, safety and well-being

AngelMed FDA Approval ASX Announcement on 28 June

MedTech/Cardiac business poised for strong product revenue growth

- Diversified medtech product design services and distribution business – 18-year history, global customers
- "One stop Services shop" bringing medical technology concepts to commercialisation
- Exclusive APAC rights to distribute *world's first/only implantable heart attack alert device*
  - Commenced first in world commercial sales/implants in SNG August 2020 *(special access scheme)*
  - Announced US FDA Approval 28 June
  - Australia TGA and Singapore HSA approvals to come
- Positioned for high growth selling Cardiac devices in APAC beginning 2H FY22

Hydrix Medical – Portfolio overview

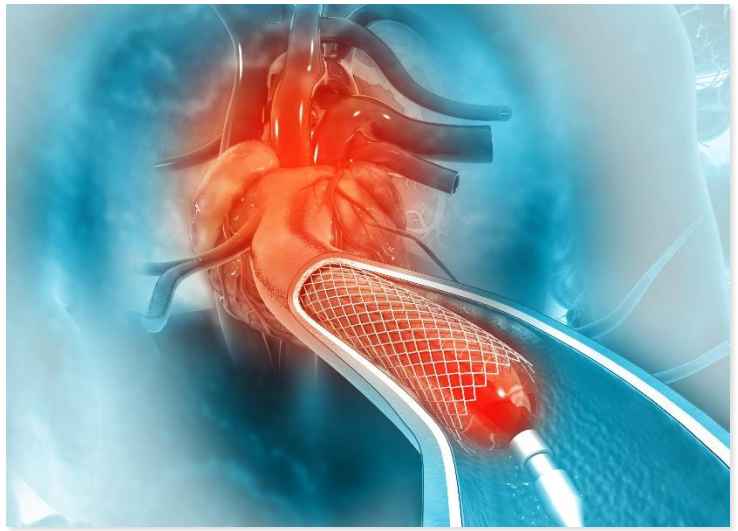
AngelMed Guardian

A 3% capture rate of patient market = $100m rev p.a. (15k units)

Implanted Heart Monitor

- 3rd party distribution agreement (7 years +2 years).
- An implantable continuous heart monitoring device that detects Acute Coronary Syndrome (ACS) events, including silent heart attacks.
- A global 'first of kind' device, approved by the FDA who concluded the AngelMed Guardian "satisfies an unmet need."

Phyzhon



A 15% capture rate of PCI surgery market = $25m rev p.a. (20k units)

Interventional cardiology device

- Exclusive rights to distribute in Australia & New Zealand.
- Single-use device for interventional cardiology surgery to manage coronary artery blockages.
- High potential to disrupt current products and procedures, improve patient and cardiologist outcomes, reduce operating time and lower hospital costs.

Indicative Milestone Timelines / Potential News Flow

1H FY22

- Submit TGA (AUS) and HSA (SNG) regulatory approval for AngelMed Guardian
- First implants of AngelMed Guardian in Australia under Special Access Scheme (SAS)
- Commence First in Human (FIH) Trials for Phyzhon in Melbourne, Australia
- Appoint APAC sub-distributors for AngelMed Guardian
- Potential Hydrix Ventures portfolio updates and revaluation events
- Ongoing evaluation of several high-growth potential “Buy, Build, Invest” initiatives

2H FY22

- AngelMed Guardian TGA (AUS) & HSA (SNG) regulatory approval
- AngelMed Guardian AUS health reimbursement approval
- Commence commercial sales of AngelMed in AUS, NZ & SNG
- CE Mark approval of Phyzhon
- Sales of Phyzhon under authorized prescriber scheme in Melbourne Australia
- TGA regulatory approval for Phyzhon once CE Mark approved
- Commence commercial sales of Phyzhon Guidewire in AUS & NZ

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