



Welcome to the Imricor Annual Meeting

Wednesday, 4 May 2022 (AEST) Tuesday, 3 May 2022 (CDT)



### Please submit questions at any time throughout the meeting

To ask a question select the 'Q&A' icon Q&A Q&A **Broadcast** Vote **Documents** Your questions(s) Select the topic your question relates to. Type your question into the chat box at the bottom of the screen and press 'Send' You may enter a question using the field below. 3 Adoption of Remuneration Report ▼ Your question will be sent immediately for Enter your question here Send review 24 character(s)



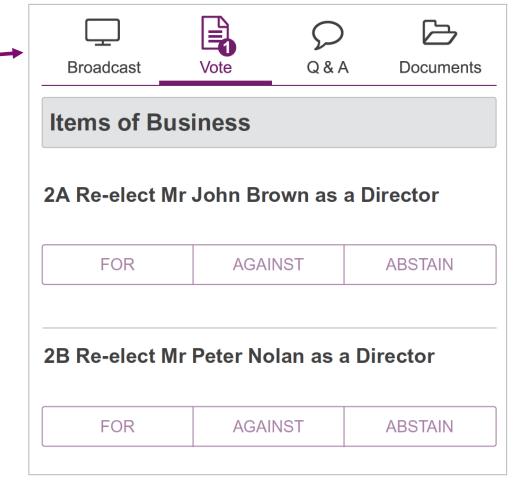
### Voting online at the virtual meeting

When the Chair declares the poll open, select the 'Vote' icon and the voting options will appear on your screen



To vote, select your voting direction. A tick will appear to confirm receipt of your vote

To change your vote, select 'Click here to change your vote' and press a different option to override.





### Imricor's Board of Directors



Steve Wedan President and CEO, and Chair



Mark Tibbles
Deputy Chair and Lead
Independent Director



Anita Messal Non-executive Director



Peter McGregor
Non-executive Director

# Chair's Address





### Delivering on our Strategic Plan



More sites to do procedures



**Grow Installed Base** 



**Expand indications** 



More procedures per site



**Expand geographies** 



Higher ASP and margin improvement



**New product development** 



### FY21 Highlights

- Contracted five new sites across Europe, bringing the total number of sites to fourteen
- Performed procedures as restrictions allowed through COVID waves
- Current contracted site status:

Current Site Status	Number of Sites
Operational	6
Installation complete (preparing to commence or re-commence procedures)	2
Preparing for installation (e.g. ordering equipment, scheduling training, etc.)	7

Two additional sites are under construction, but are not yet under contract with Imricor



### FY21 Highlights continued

- Appointed Regional Health Care Group (RHCG) in Australia to help facilitate TGA and Medsafe approvals
  - Medsafe approval received for all Imricor products in New Zealand
  - Received TGA approval on Imricor's Advantage-MR System, Vision-MR catheter under review
- Entered into an exclusive distribution agreement with RHCG for consumables, and non-exclusive distribution of capital equipment in Australia and New Zealand
- Filed an application for an Investigational Device Exemption (IDE) from the US Food and Drug Administration (FDA)
- Successfully raised A\$16.5 million via an oversubscribed institutional placement at \$1.00 per CDI
- Successfully raised A\$1 million in October via an oversubscribed security purchase plan (SPP) offered to eligible CHESS Depositary Interest (CDI) holders in Australia and New Zealand
- Entered into two Sales Distribution Agreements with:
  - NordicNeuroLab AS (NordicNeuroLab), a leading maker of MRI compatible in-room monitors
  - MiRTLE Medical, LLC (MiRTLE), maker of an MRI-compatible 12 lead ECG system and
- Completed a strategic investment in MiRTLE



### Site expansion plans underpinned by strong pipeline

Helios Hospital Berlin-Buch, Germany



Semmelweis University Heart and Vascular Centre, Hungary



**German Heart Centre Berlin, Germany** 



Charité Medical University Virchow-Klinikum Campus, Germany



**Henry Dunant Hospital Centre, Greece** 



Policlinico Casilino, Italy (new in 2022)



Dresden Heart Centre, Germany

Helios Leipzig Heart Centre, Germany

Lübeck University Hospital, Germany

Rhön Clinic Bad Neustadt Campus, Germany

> Münster University Hospital, Germany

South Paris Cardiovascular Institute, France Maastricht University Medical Centre, The Netherlands

Amsterdam University Medical Center, The Netherlands

> Haga Hospital, The Netherlands

The Lausanne University Hospital, Switzerland

The Heart and Diabetes Centre NRW, Germany



### Partnering to drive growth in clinical sites

Imricor has entered into a number of agreements that promote future iCMR site adoption





## Developing product lines to support expanded indications & margin improvement



#### Steerable Sheath and Transseptal Needle

- Preclinical trials underway
- Undergoing final device testing
- Preparing for approval submission as part of VT trial in 2022
- Preparing for CE mark submission in parallel with VT trial

The steerable sheath and transseptal needle are intended to be used together in procedures where access to the left side of the heart is required and the physician opts to access the left side by crossing the intra-atrial septum



#### **Diagnostic Catheter**

- Utilising advancements from the next generation ablation catheter to create a more consistent product line for physicians and reduce production costs to drive a higher gross margin
- Currently in 3<sup>rd</sup> round of technical documentation review with Notified Body
- CE mark expected in 2022

A diagnostic catheter can sense electrical signals flowing through the heart and provide cardiac stimulation but is not used for ablation. A diagnostic and ablation catheter is required to perform atrial flutter procedures - currently sold by Imricor in a two-catheter set comprising two ablation catheters.







# A strong and growing market in cardiac ablation

A large global addressable market with high growth potential supported by favourable growth drivers

#### **Drivers of Global Catheter Ablation Market**



Increased incidence of cardiac disease



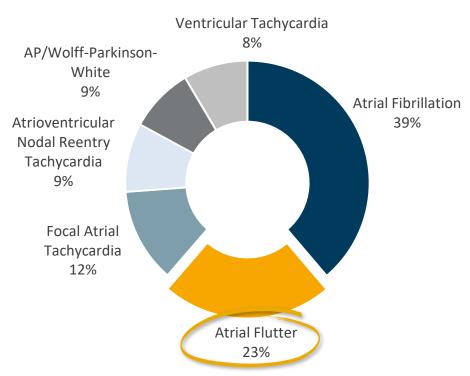
• Shift towards minimally invasive procedures



 Cost effectiveness of catheter ablation as treatment option

#### **Breakdown of Ablation Procedures**

Ablation Procedure Types EU 2019





### Focus Initiatives for 2022

#### **Site Growth**

- Increased engagement with Electrophysiologists to drive demand for iCMR labs
- Increased engagement with **Philips and Siemens** sales forces to help deliver the pipeline of iCMR labs

#### Utilisation

- Initiate procedures at each contracted site
- Increase procedure volume at each contracted site

#### **Market Expansion**

- Progress approval of VT indications through preclinical and clinical trials
- Progress TGA approval for Imricor's Vision-MR Ablation Catheter in Australia
- Progress Investigational Device Exemption (IDE) from the FDA in the US to begin clinical trial

#### **Products**

- Submit second-generation ablation catheter for CE mark approval
- Submit steerable sheath and transseptal needle for CE mark approval
- Receive CE mark approval for diagnostic catheter
- Progress development of MRI compatible biopsy system



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