

Tuesday 21 May 2024

Pre-conference workshops			
9.00am – 5.00pm	ANDHealth Digital Summit Location: Launch Pad @ Stone & Chalk, Lot Fourteen, Corner North Terrace and Frome Road, Adelaide 5000 Following the success of the inaugural Digital Health Summit in Adelaide last year, ANDHealth are returning with a focus on AI, investment and industry trends! Join leading experts as they ask the question – digital health: hype or here to stay? The full-day event will be packed with engaging thought leaders and feature keynote presentations, panel discussions, and networking opportunities. Topics will include: Al in health: We're asking the wrong questions Health beyond the hype: Is Al here to stay?		
	 Myth-busting: The untapped value of patenting in software Digital health: The investor perspective Cybersecurity and beyond: Building trust in the AI age Bridging the gap between digital promise and clinical practice Regulating at the speed of tech At the conclusion of the Summit, delegates will move to the Adelaide Convention Centre, where the networking reception will be held in the AusMedtech Exhibition Hall from 5:30pm – 7:00pm. 		
8.30am – 12.30pm	ASEAN regulatory requirements - addressing the practical issues The ASEAN region is an important market for Australian medical device companies for a variety of reasons, such as its proximity (example being physically closer to Australia than Europe), large market size (population of 600+ million) and increasing demand for improved standard of care and therefore medical devices. ASEAN medical device regulations are guided by the ASEAN Medical Device Directive, introduced about a decade ago		

In terms of regulatory approvals, ASEAN is different from European CE certification (CE Marking) process, with no version of CE Marking existing in the ASEAN region. Medical device companies need to seek regulatory approvals for each ASEAN country. The effort for medical device Regulatory Affairs team is not to be underestimated.

The objectives of this workshop are to:

- Gain an appreciation of ASEAN medical device regulations and the implementation across each country.
- Get an update on current topics of interests on ASEAN medical device regulations.
- Clarify questions you might have in your preparation for registering medical devices in the region. (Please provide your questions in advance via email).
- Appreciation of the requirements for Good Distribution Practice Medical Devices (GDPMD). GDPMD has been and continues to be, rolled out across the ASEAN countries. GDPMD is a requirement for obtaining Import/wholesale/distribution license.

1.00pm - 5.30pm

AusMedtechInvest 2024

AusMedtechInvest 2024 is Australia's inaugural boutique medtech investment roundtable forum and aims to build meaningful personal connections between innovative medical technology (devices and diagnostic) businesses and investors, and help great ideas attract the capital needed to thrive in a fast-paced, competitive market.



1.30pm - 4.30pm

AusMedtech South Australian Site Tours







Adelaide BioMed City



Tour One: Adelaide BioMed City

Tour time: 1:30 pm – 4:30 pm **Check-in:** 1:00 pm – 1:30 pm

Meeting point: Adelaide Convention Centre - AusMedtech Registration Area

Transport: Walking tour

Tour Two: Lot Fourteen

Tour Time: 1.30 pm – 4.30pm **Check-in:** 1:00 pm – 1:30 pm

Meeting point: Adelaide Convention Centre - AusMedtech Registration Area

Transport: Walking tour and tram

Tour Three: Tonsley Innovation District

Tour Time: 1.30 pm – 4.30pm **Check-in:** 1:00 pm – 1:30 pm

Meeting point: Adelaide Convention Centre - AusMedtech Registration Area **Transport:** Coach to Tonsley and returning to the Adelaide Convention Centre.

Refreshments will be provided.

Tour Four: Mawson Lakes | Australian National Fabrication Facility - South Australia

Tour Time: 1.30 pm – 4.30pm **Check-in:** 1:00 pm – 1:30 pm

Meeting point: Adelaide Convention Centre - AusMedtech Registration Area

Transport: Coach and walking tour. Refreshments will be provided.

5.30pm -7.00pm	Pre-Conference welcome reception
	AusMedtech 2024 Welcome reception
	Sponsored by:
	FB RICE
	The IP Navigators
7.30pm	Clinical trials dinner AusBiotech and its AusBiotech Clinical Trials Advisory Group warmly invite those working in the clinical trials sector to join them for this unsponsored, informal dinner and to connect face-to-face before AusMedtech 2024 conference officially begins.
	Clinical trials contribute to Australia economically and socially and are a critical component in the development process of bringing new therapies, devices and diagnostics to patients. However, the Australian clinical trial ecosystem is complex, involving many parts and stakeholders; this dinner is an opportunity to unite under the same roof and celebrate the sector and its progress.



Wednesday 22 May 2024

07.30am –	Delegate registration
08.30am	
8.30am –	Opening & welcome
8.50am	
	Conference opening and welcome address
	Welcome to country, Major Sumner AM, Ngarrindjeri/Kaurna Elder
	• The Hon Susan Close MP, Deputy Premier for South Australia, Minister for Industry, Innovation and Science, Minister for Climate, Environment and Water, and Minister for
	Defence and Space Industries
	Rosanne Hyland, Chief Operating Officer, AusBiotech
8.50am-	Keynote session
9.15am	
	National Health Medical Research Council (NHMRC) update
	Chair: Gavin Fox-Smith, Chief Executive Officer, Omnigon
	Professor Steve Wesselingh, Chief Executive Officer, National Health Medical Research Council (NHMRC)
	Further details to be confirmed

9.15am –	Plenary session			
10.00am	Australian success stories			
	This panel discussion will celebrate the 2023/24 successful expansion and growth of medtech companies. Followed by a Q&A with the audience.			
	Chair: Robyn Lindner, General Manager, AusBioNSW			
Panellists: • Sam Lanyon, Executive Director, Lumos				
	 Sam Lanyon, Executive Director, Lumos Marjan Mikel, Chief Executive Officer & Managing Director, Respiri Paul Anderson, Chief Executive Officer & Managing Director, Orthocell 			
	Jerneen Williams, Director of Operations, Bellberry Limited			
10.00am – 10.25am	Keynote session			
	Medical device regulation: an update on the latest			
	With sweeping reforms occurring across the globe in medical device regulation, this session focuses on how the Therapeutic Goods Administration (TGA) is supporting the industry through these changes, along with the challenges of regulating new and emerging technologies and what this means for Australian patients.			
	Chair: Karen Parr, Director Policy & Communications, AusBiotech			
	Professor Anthony Lawler, Deputy Secretary, Medical Devices & Product Quality Division, Therapeutic Goods Administration			
	Professor Antifory Lawrer, Deputy Secretary, Medical Devices & Product Quanty Division, Therapeutic Goods Administration			
10.25am –	Morning tea in the exhibition			
10.55am 10.55am-	Plenary session			
11.55am	Tienary session			
	Harnessing the pipeline: Partnering with big medtech			
	For medtech companies commercialising biomedical research across the ecosystem, there is an interdependent relationship between small and large companies. Multinational companies are a critical part of the Australian landscape, with their ability to provide the resources, experience, and infrastructure necessary to support R&D, clinical development,			
	manufacturing, and distribution of devices and diagnostics.			
	This plenary session provides an invaluable opportunity to hear from big medtech on partnering, prospecting, and support work in the Australian environment.			
	Chair: Dell Kingsford Smith, AusBiotech Board; Vice President Medical Affairs, Market Access & Government Affairs, Asia Pacific, Cochlear			
	Panellists:			
	Maurice Ben-Mayor, President & Managing Director, Stryker			
	Mick Trevaskis, Chief Executive Officer, Device Technologies			
	Rebecca Cortiula, Senior Managing Director Australasia, Varian			
	Jane Crowe, Managing Director, ANZ Cardinal Health			
	Pat Williams, Vice President & Country Manager ANZ & Korea, Edwards Lifesciences			
11.55am –	Room move break			
12.00pm				

12.00	Seed on A	Cassian D	Cassian C
12.00pm – 12.55pm	Session A	Session B	Session C
	Europe continues to face a dynamic and challenging regulatory environment. In effect since 26 May 2021, the European Union's (EU) medical device (and In-Vitro Device Regulation (IVDR)) regulations mandate that medical device manufacturers targeting the EU market must adhere to new standards. Key changes introduced by these regulations include: New requirements for translations Stricter clinical evaluation and post-market surveillance Expanded product scope Introduction of unique device identification (UDI) systems Greater transparency and data reporting Reinforced patient safety measures, and Enhanced role of notified bodies This panel of seasoned experts will discuss the issues facing the sector and pitfalls new players often come across. Chair: George Loizou, Director, CMS SciDoc Pty Ltd Panellists: Julie Winson, Quality and Regulatory Director, LBT Innovations Anthony Skeats, Chief Operating Officer, Micro-X Chris Henry, Managing Director, Actis Medical Hwee EE Tan, Principal Consultant, DH RegSys Consulting Pty Ltd	Building workforce skills and capabilities for successful medical device commercialisation In this session facilitated by ARCS Australia, the focus is on workforce skills and capabilities crucial for the effective commercialisation of medical devices. The session will feature representatives from industry, academia, and government perspectives to provide cross sector insights. Facilitated discussions will emphasise cross-sector collaborations, aiming to empower attendees with actionable insights for navigating the intricate landscape of medical device commercialisation. This session aspires to foster collaboration and skill enhancement, ensuring successful market entry for medical innovations. Chair: Dr Tim Boyle, Chief Executive Officer, ARCS Panellists: Dana Bell, Director Partnership SA, MTPConnect Shan-Shan Wang, Founder & Chief Executive Officer, Roam Technologies (RoamTech.ai) Ajay Nair, President APAC, Mullings Group Paul Cohen, Managing Director & Founder, Paul Cohen Consulting In partnership with:	Capital markets: is the 'old normal' actually the 'new normal'? After almost 2 years of steep declines in valuation, many industry players are talking about a "return to normal". But were the heady days of no-low due diligence, FOMO driven investments and "frothy" valuations really normality? Or are we just longing for a moment in time to return? Chair: Bronwyn Le Grice, Managing Director & Chief Executive Officer, ANDHealth Panellists: • Dr Chris Nave, Founding Partner & Managing Director, Brandon Capital • Dr Steve Burnell, Managing Director, TenMile • Elyse Shapiro, Healthcare Analyst, Canaccord Genuity
12.55pm - 1.55pm	Lunch in the exhibition		

Concurren	Concurrent sessions				
1.55pm 2.50pm	Session D	Session E	Early-Stage Innovation Forum		
	Ethics and AI – has the horse already left the stable? Artificial intelligence has been heralded as the largest technological change facing our generation, but concerns about ethics, privacy and consumer protection are still front of mind. As the Australian Government contemplates regulation of AI, we have to consider whether this might be a case of shutting the stable door after the horse has bolted. Chair: Grace Lethlean, Chief Product Officer, ANDHealth Panellists: • Daniela Cecere-Palazzo, Senior Lawyer, Youlegal • Angie Corbo, Informatics Global Catalyst Innovation & Architecture, Roche • Liesl Yearsley, Chief Executive Officer & Founder, Akin • Justine Lacey, Director, Responsible Innovation Future Science Platform, CSIRO In partnership with:	Set yourself up for funding success by learning about the suite of programmes funded by the Federal Government that support medtech innovation. Hear about the purpose and intention of each programme, eligibility criteria, who they are best suited for, the type of projects they are looking to support, and upcoming dates. Chair: Dr Tracey Wilkinson, Director Stakeholder Engagement WA, MTPConnect Panellists: Associate Professor Tracey Laba, Director Frontiers Program, Medical Research Future Fund, Dept of Health & Aged Care Representative, National Reconstruction Fund David Chuter, Executive Director, Industry Growth, Department of Industry, Science and Resources Liz Crompton, SME Connect Program Advisor, CSIRO	The Early-Stage Innovation Forum will feature rapid fire rounds of quick-pitch presentations on early-stage technologies and projects from local research institutes, universities, hospitals, and pre-series A companies in the areas of medical devices and diagnostics, digital health and enabling technologies. Expert panel: • Michelle Gallaher, Chief Executive Officer, Cerulea • Eddie Walker, Partner, FB Rice • Sarah Meibusch, Partner, OneVentures • Steve Burnell, Executive Chair, Tenmile • Mark Lawler, Country Manager ANZ, West Pharmaceuticals Presentation 1: ARIA, the world's first commercially scalable Non-Invasive Bionic Vision System Presented by: Warren Bingham, Global Vice President, ARIA Research Presentation 2: Diagnosis of early cardiovascular disease via a blood qPCR test Presented by: Prof Gemma Figtree, University of Sydney, Kardiomics Presentation 3: IMAGENDO: Building a revolutionary healthcare solution for women health using Al Presented by: Jodie Avery, Senior Research Fellow, The University of Adelaide Presentation 4: JiffyStent Inserter - redefining stent insertion for the relief of kidney stone pain, saving significant pain, time, and money Presented by: Joseph Ischia, Co-Founder & Chief Executive Officer, JiffyStent		

3.45pm- 4.15pm	Afternoon tea in the exhibition			
Concurrent sessions				
4.15pm- 5.10pm	Session H	Session I	Early-Stage Innovation Forum	
	Reimbursement Are Australia's medtech reimbursement pathways appropriately designed and sufficiently rewarding to attract both local and global innovation? Chair: Dell Kingsford Smith, VP Medical Affairs, market access and Government Affairs, Cochlear Panellists: Penny Shakespeare, Deputy Secretary, Department of Health and Aged Care Polo Guilbert – Wright, Senior Director Government Affairs ANZ, Edwards Lifesciences George Papadopoulos, Director & Partner, Lucid Healthcare Consulting Sarah Griffin, Director, MedTechnique Consulting Nicola Leavold, Commercial Director Australasia, BXTA	The growing importance of environmental, social, and governance (ESG) and how does the medtech board react to this? The Boston Consulting Group states "Health care accounts for 5% of total global carbon emissions, and medical devices and technology are responsible for a large portion of that. Much of this comes from the manufacturing operations and supply chains of medtech companies and their suppliers. At the provider level, medtech generates tonnes of unrecycled waste through single-use disposable products and packaging." There is a growing demand from customers, shareholders and employees for Boards to implement ESG initiatives within organisations. Medtech companies must choose to balance current competitiveness and future investments while connecting ESG to economic value. This panel of experienced executives and Board members will tackle the thorny issue of how to do this whilst generating value. Chair: Melissa McBurnie, Partner & Head of Impact, Brandon Capital Panellists: Olivia Pitt, Head of ESG, Ellerston Capital Lis Boyce, Partner, Piper Alderman Elizabeth Dallimore, Executive Chair, Inspiring Holdings Dr David Brookes, Executive Chair, Anatara Lifesciences		
7.00pm– 7.30pm	Pre-dinner drinks			
7.30pm – 10.00pm	AusMedtech conference dinner Rosanne Hyland, Chief Operating Officer, AusBiotech Dinner speakers: Scott Stirling, Chief Executive Officer, Red Dust, and Jonathon Lindsay-Tjapaltjarri Hermawan, Director of Programs and Strategy Lead, Red Dust Facilitator: Rebekah Cassidy, Chief Executive Officer, AusBiotech			

With Australia's indigenous communities overrepresented in low socioeconomic figures, Australia's First Nations people face a greater risk of poor health, higher rates of illness, disability, and death, and live shorter lives than people from higher socioeconomic groups.

Founded in the Northern Territory, not-for-profit Red Dust draws on the strengths of both western health knowledge and traditional cultural knowledges to create a positive influence on young people and improve outcomes for communities. Providing a 'community-as-family' model of health and well-being programmes, Red Dust works alongside community leaders and elders to create a stronger future for indigenous youth and their families.

Join us for a fireside chat with Red Dust, Director of Programs and Strategy Lead Jonathan Hermawan and CEO Scott Stirling, as they discuss their personal journey over the past seven years growing Red Dust and developing a cooperative model that is charting a new course and leading to improved health and wellbeing outcomes.

Sponsored by:





Thursday 23 May 2024

8.30am – 9.00am	Delegate registration		
9.00am – 9.15am	Hon Stephen Dawson MLC, Minister for Emergency Services; Innovation and the Digital Economy; Science; Medical Research; Minister Assisting the Minister for State and Industry Development, Jobs and Trade, Western Australian Government		
	Chair: Rosanne Hyland, Chief Operating Officer, AusBiotech		
9.15am – 10.00am	Plenary session		
	Issues facing healthcare in Australia Get your caffeine fix from the exhibition hall and start day 2 of the conference with a wide-ranging panel discussion with leaders from our representative industry and government organisations about the burning challenges- and the exciting opportunities- facing the Australian healthcare sector that are keeping them awake at night. Chair: Stuart Dignam, Chief Executive Officer, MTPConnect		
	Panellists: Rebekah Cassidy, Chief Executive Officer, AusBiotech Dr Anna Lavelle, Chair, Medicines Australia Ian Burgess, Chief Executive Officer, MTAA Dr Tim Boyle, Chief Executive Officer, ARCS Bronwyn Le Grice, Managing Director & Chief Executive Officer, ANDHealth		

10.00am – 10.45am	Keynote session			
	What women don't want Much is made about the progress of corporate Australia in the diversity stakes, however 2023 saw some disturbing stories about women in corporate Australia sitting on the receiving end of verbal and physical harassment and abuse. The digital harassment of women leaders is now a frequent occurrence for women in politics and advocacy, academia, media, and private industry, with insults, abuse, threats, trolling and doxing now a commonplace for women leaders.			
	A play on the 2000 film What Women Want, this panel will reflect on their personal experiences as they shine a spotlight on the treatment of professional women, and challenge industry to become champions of change in ending discrimination against women in the workplace. Chair: Bronwyn Le Grice, Managing Director & Chief Executive Officer, ANDHealth			
	 Rebecca Wilson, Chair, Alcidion & LBT Innovations & NED Hansen Technologies Dr Elaine Stead, Principal, Main Sequence Ventures Michelle Gallagher, Chief Executive Officer, Cerulea 			
10.45am – 11.15am	Morning tea in the exhibition			
11.15am – 12.10pm	Session J	Session K	Session L	
·	Al in medical imaging – from promise to practice The promise of artificial intelligence (AI) in medical imaging analysis has been discussed for many years and is now a reality. Was it smooth sailing for this seemingly ideal use case? Are there lessons to be shared which will inform the next wave of technologies? In this panel discussion we'll hear from those who have been at the forefront of the development and adoption of AI in medical imaging, they'll share what's worked, what didn't, the challenges that remain and where to next. Chair: Dr Ludovic Labat, Chief Executive Officer, Neo-Bionica Panellists: • Dr Michelle Perugini, Chief Executive Officer, Presagen Dr Mark Phillips, Head of Clinical Research &	Exploring the emerging material science manufacturing and investment opportunities to reduce medical plastic waste Hospital generated plastic waste is a worldwide issue. When first introduced into hospitals, single-use plastics were an attractive option as it allowed for maintenance of a sterile environment and infected plastic material could be easily disposed via landfill waste. For equipment such as syringes this is vital. However, the sheer quantity of single use plastics being used in hospitals is becoming alarming, particularly considering the overuse or unnecessary use of single-use plastics. In Australia, the healthcare industry is responsible for around 8% of Australia's carbon emissions and generates large amounts of non-recyclable plastic waste. Environmental Social Governance (ESG) factors are increasingly becoming a factor in procurement and related commercial decisions.	As an early stage medtech company or founder it is often difficult to know exactly what is required to get a product/asset through the clinical phase. This session will explore the things you need to consider when running a clinical trial and before you start writing the protocol. You have a product, you know what you want it to do, you've even got it to the point it's ready for clinical testing and met all the manufacturing regulatory requirements. But wait, clinical research has its own regulatory standards and requirements. How on earth do I navigate this world? The panel will discuss the considerations for embarking on the clinical development phase of a trial, when you should start to plan, what to look out for, whether to hire or contract expertise and how to engage a site(s) and clinician(s). Chair: Natalie Barber, Director, Clinical Operations, Chrysalis	
	Medical Affairs, Annalise.ai Dr Wenji Pang, Chief Scientific Officer - Imaging & AI, Resonance Health Alison Deslandes, President, Australasian Society of Ultrasound in Medicine	This diverse panel session will discuss ways to address these issues and funding assistance that is available. Chair: Stuart Anderson, Clinical Translation and Commercialisation Medtech Program Manager, MTAA	Panellists: Simon Cook, Executive Director, Operations, Eudaemon Technologies Helen Plummer, Research Manager, Cercare Prof Les Bokey, Institute Director, Ingham Institute	

		 Panellists: Jane Crowe, Managing Director, ANZ Cardinal Health Dr Adam Walczak, Director, HNE Innovation Office, Hunter New England Local Health District Pauline Salib, Antimicrobial Stewardship Pharmacist, Western Health Leonie Walsh, Interim Chair, Solving Plastic Waste CRC Dr Carly Hollier, A/Senior Manager, Sustainability, Delivery Excellence, Partnerships & Projects, HealthShare NSW In partnership with: 	for Applied Medical Research • Simon Belcher, General Manager, Three Peaks Medical
12.10pm – 1.05pm	Session M	Session N	Session O
	Strategic considerations in manufacturing for medtech With a strong push and incentives for sovereign and local manufacturing capacity, how might medtech companies think about whether to buy, build or partner to develop the capabilities, skillsets and tools required to execute on manufacturing at scale. This session will explore the strategic thinking required for manufacturing business decisions, including the impact of technologies such as additive manufacturing, Industry 4.0, robotics, AI and other emerging technologies. Chair: Nick Northcott, Managing Partner, Chrysalis Advisory Panellists: Lisa Henretty, Chief Operating Officer, Enersol Louisa De Vries, Consulting Manager, Bosch Australia Manufacturing Solutions Val Valentine, Director, Edwards Lifesciences Pablo Solis, Chief Executive Officer & Co-Founder, Protego Medical	What angel investors look for in a MedTech startup company Many early-stage companies require external funding. One source of funding can come from Angel Investors, who typically provide a cash injection in exchange for equity in the company. Startups, particularly in the medtech sector, can be a risky proposition for an investor, and there are key elements that an Angel Investor will look for when deciding whether to invest in that company. Such considerations may include the strength of the team behind the startup, the business model, what intellectual property the startup has, and ultimately, what sort of return the Angel Investor can obtain on their investment. This panel session, which comprises angeliInvestors who are active in investing in startups in the medeech industry, will discuss the key elements that they look for in a startup when deciding whether or not to invest, and provide insight into how you, as a startup, can best position yourself so as to make yourself as attractive to	Are medtech and digital product clinical trials - same or different? Following on from the previous panel session that explored how early stage medtech companies navigate the clinical phase, in this session the invited speakers explore more specifically the commonalities and differences in designing, running and managing clinical trials between digital products and devices, as well as digitally enabled devices. Founder will share how they navigated the sometimesunclear requirements for Software as a Medical Device trials in both Australia and the US, and what they learned from the more established medical device trials. We also explore how to work with CROs, and how to assess who would be the right partner. The goal of the session is to get clarity on what common pitfalls are, and highlight relevant factors and strategies to accelerate the clinical phase for both devices and digital products.

		an angel investor as possible. Some of the panel members have also been in the position of forming a startup and themselves seeking investment, and can provide valuable advice from their own experiences. The session will be chaired by Dr Milena Dryza, Senior Associate of Madderns Patent and Trade Mark Attorneys. Milena is a patent attorney with many years of experience in private practice, as well as an in-house patent counsel for leading biotechnology company, CSL. Chair: Dr Milena Dryza Senior Associate, Madderns Patent & Trademark Attorneys. Panellists: • Dr Anabela Correia, Chief Executive Officer, LiVac • Dr Nick Haan, Chief Executive Officer. Seonix Bio • David Saint, Chair, Southern Angels • Dr Amandeep Hansra, Lead Investor and Cofounder of Medical Angels Sponsored by: Madderns Sponsored by:	Chair: Dr Katja Beitat, Head of Health Tech, Cicada Innovations Panellists: • Helen Souris, Chief Executive Officer, CardiHub • Mary-Beth Brinson, Chief Executive Officer, Cyban • Stewart Bartlett, Chief Executive Officer, Ferronova • Eric Davies, Advisor to Roam Technologies (RoamTech.ai)
1.05pm – 2.00pm	Session P	Session Q	Session R
AusMedtech 202	What infrastructure do we need to build a vibrant medtech innovation and commercialisation ecosystem The road to successful commercialisation in medtech is often winding and with many obstacles along the way. Being aware of the pitfalls and shaping your start-up for the best possible outcome is no simple task. It has been said we need the right people, science, infrastructure and money to build a success. Incubator and accelerator programmes are a vital part of the medtech innovation ecosystem but what else is required to help our sector? and provide a rich environment of education, mentorship, industry collaboration, networking and capital raising support. Our panel will look into the roles of education,	How to close a VC round (and retain your sanity) Competition for capital is fierce at the moment, with investors focusing on the stability of their existing portfolio and wary of cash burn in a tight market. So how DO you land a significant investor, and more importantly, what do you need to do to close the deal! Chair: Sarah Meibusch, Partner, OneVentures Panellists: Arthur Shih, Chief Executive Officer, Humanetix Peter Vranes, Chief Executive Officer & Co-Founder, Nutromics	Partnering - the pathway to successor is it? A business partnership is a collaboration between two or more entities that pool resources, technology and/or finances to achieve a generally agreed goal. In medtech this is a widely accepted pathway to develop technology, access markets or ensure access to specific technologies. Whilst there are many instances of this being a success there are also many instances where the expected success did not materialise. This panel of seasoned executives will discuss many of the pitfalls and key decision points where partnerships are useful of inhibitory. Chair: Peter Bradley, Principal, Qatalyst Consulting

	 Richard Horton, Partner, Squire Patton Boggs or Melissa McBurnie, Partner & Head of Impact, Brandon Capital. Richard Horton, Partner, Squire Patton Boggs or Melissa McBurnie, Partner & Head of Impact, Brandon Capital. Peter Rowland, Non-Executive Director, Micro-X or Michael Kavanagh, Chief Executive Officer & President, Nanosonics or President, Nanosonics or President, Nanosonics or President, Chief Executive Officer, LiVac Dr Lilly Bojarski, General Manager, Cicada Health Tech Hub Natalie Rickers, Commercialisation Director, South Australia MTP Connect Kelly Coverdale, Managing Director, Cover Biomedical Professor Sharath Sriram, Director, Discovery to Device Facility, RMIT University; President, Science & Technology Australia
2.05pm – 3.00pm	unch in the exhibition
3.00pm – 3.30pm	Plenary session
	essons from 20 years of collaborations creating meaningful medtech ventures With a global presence and multidisciplinary skills, IDE Group is a well-regarded partner for medtech commercialisation and product development. DE works with its partners to find and assess business opportunities, conduct research, create and implement commercial strategy, gain access to funding, develop new echnology, manufacture quality products and create successful medtech ventures. Since 2003, IDE has grown over 150 medical technology businesses and realised over 500 projects across the medical technology landscape globally. You will hear from IDE Group CEO George Sidis and Eudaemon Technologies Executive Director, Operations, Simon Cook as they discuss the organisation's journey to date and how the supported Australian medtech innovators to generate high-impact ventures and improve health outcomes. Chair: Warren Bingham, Global Vice President, ARIA Research George Sidis, Chief Executive Officer, IDE Group & Simon Cook, Executive Director, Operations, Eudaemon Technologies

3.30pm –	Plenary session
4.15pm	
	Take home lessons from AusMedtech 2024 The two days of AusMedtech are a cornucopia of data, information, knowledge, and wisdom. Many of us will have individual gems that are pertinent to our particular business or problem to take home. However, many of the big picture and relevant issues explored may be missed because of conflicts or competing priorities drawing attendees to other sessions. This panel of hardened Medtech executives have made it a mission to pick the pearls of wisdom from the conference and highlight to the final audience. We will also draw on the collective attendees to reduce these to actionable take home messages from the conference. Chair: Peter Bradley, Principal, Qatalyst Consulting
	Panellists: - Jo Close, Director Adelaide Intermediary Program, MTPConnect - Craig Newton, Chief Executive Officer, Kynetyka - Aisah Sirop, Commercial Manager, The University of Adelaide - Jack Brown, Corporate Development Director, LBT Innovations
4.15pm – 5.00pm	Conference closing reception Exhibition Hall Rosanne Hyland, Chief Operating Officer, AusBiotech