Programme as of April 28, 2025



An event by 🔅 MedTech Europe

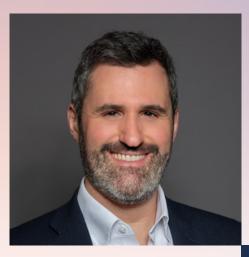
13 – 15 May > Lisbon

#MTF2025 | themedtechforum.eu



In cooperation with

WELCOME INTRODUCTION



Dear participants, Dear speakers, Dear sponsors,

Following the great success of The MedTech Forum 2024 in Vienna, which surpassed all previous records, I am delighted to welcome you to the next chapter of our journey - The MedTech Forum 2025, which will be held in Lisbon from 13 to 15 May.

The 2025 programme will feature leading voices in the medical technology community, including industry experts and key stakeholders. Together, we will tackle the ever- evolving opportunities and challenges for our industry, from innovation and digital transformation to regulatory developments and sustainability.

I encourage you to take full advantage of the many networking opportunities throughout the event. Whether it's sharing ideas, forging new partnerships, or reconnecting with colleagues, these moments are where some of the most valuable insights and collaborations are born.

Best regards,

Oliver BISAZZA Chief Executive Officer MedTech Europe

PROGRAMME AT A GLANCE

TUESDAY 13 MAY 2025

18:30-22:00 Welcome Cocktail Reception

PÁTIO DA GALÉ

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for Medtech Innovation Hospital Walls Commercial—Insights from BCCS 4th Milkman Study Sponsored by BCG Thealth Data Space In the US and European Ma AVANIA EXHIBITION AREA 1150-12:20 NETWORKNO BREAK From Evaluation to Action: The Future of MDR and IVDR 220-13:10 SPONSORED SESSION MedTech Roberts beter and safer care 12:20-13:10 SPONSORED SESSION Navigating global business challenge and ModTech Roberts beter and safer care 12:20-13:10 SPONSORED SESSION Navigating global business challenge and ModTech Roberts beter and safer care 12:20-13:10 SPONSORED SESSION Navigating global business challenge and ModTech Roberts beter and safer care 12:20-13:10 SPONSORED SESSION Navigating global business challenge and ModTech Roberts beter and safer care Safe ALS matter Care: ransforming European Informating European Informating European Informating European Informating Diagnostics 14:10-115:00 PARALLEL SESSION UK - EU: ready for a reset? 14:10-115:00 PARALLEL SESSION UK - EU: ready for a reset? 15:10-18:00 PARALLEL SESSION UK - EU: ready for a reset? 15:10-18:00 PARALLEL SESSION UK - EU: ready for a reset? 15:10-18:00 PARALLEL SESSION UK - EU: ready for a reset? 15:10-18:00 PARALLEL SESSION UK - EU: ready for a reset? 15:10-18:00 PARALLEL SESSION UK - EU: ready for a reset? 15:10-18:00 PARALLEL SESSION Navigating US: Prove the Roberts of the Parallenge commistration opooting or theads 15:10-18:00 <t< td=""><td>1:00-11:50 PARALLEL SESSION</td><td>11:00-11:50 PARALLEL SESSION</td><td>11:00-11:50 SPONSORED SESSION</td><td>11:00-11:50 PARALLEL SESSION</td><td>11:00-11:50 ASK THE EXPER</td></t<>	1:00-11:50 PARALLEL SESSION	11:00-11:50 PARALLEL SESSION	11:00-11:50 SPONSORED SESSION	11:00-11:50 PARALLEL SESSION	11:00-11:50 ASK THE EXPER
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Safe AI, Smarter Care::::::::::::::::::::::::::::::::::::	EXHIBITION AREA	13:10-14:10 LUNCH BREAK			
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	EXHIBITION AREA	10:20-10:50 NETWORKING BREAK			

10:50-11:40 PARALLEL SESSION

Global Market Focus: Brazil

Advancing the Assessment of

Digital Health to Foster Innovation

13:40-14:30 PARALLEL SESSION

A New Cardiovascular Health Plan for Europe, a Game Changer for Patients

14:40-15:30 PARALLEL SESSION

Navigating Interoperability in the EHDS: Roles and

Responsibilities for Manufacturers

PARALLEL SESSION

11:50-12:40

PARALLEL SESSION

SPONSORED SESSION

PARALLEL SESSION

Navigating Litigation: Securing

MedTech's Future in Europe

Closing the Loop: Turning Customer Insights Into Product Excellence

Sponsored by SMARTEEVA

Cutting Edge: The Future of Robotic Surgery

14:40-15:30 PARALLEL SESSION

Engaging with Impact: Stories of MedTech and Patient Collaboration

11:50-12:40

10:50-11:40 SPONSORED SESSION

Aligning PLM & QMS Processes for

MedTech Excellence

Sponsored by VEEVA MedTech

11:50-12:40 PARALLEL SESSION

Shaping the Future of Wound Care

13:40-14:30 PARALLEL SESSION

Fast-track health technology assessment for in vitro diagnostics

APIFARMA

 14:40-15:30
 PARALLEL SESSION

 Measure Your Readiness with the Self-Assessment Tool
 14:40-15:30
 ASK THE EXPERT

 Market Intelligence
 Market Intelligence
 Market Intelligence

10:50-11:40

11:50-12:40

13:40-14:30

ASK THE EXPERT

ASK THE EXPERT

ASK THE EXPERT

Everything You Wanted To Know

About IHI (But Were Afraid To Ask)

How AI & PLM Drive Smarter

Compliance in MedTech

PTC/DELOITTE

Unlocking Health Data: The EU Data Act's Impact on Digital Health

REEDSMITH

15:40-16:00	PLENARY
Conclusions	

10:50-11:40 PARALLEL SESSION

Re-imagining Cyber Secure Health

Implementing the EU Green Deal in Healthcare

EXHIBITION AREA

Innovating for Planetary and Human Health: Sustainability

Trends at Global Level

14:40-15:30 PARALLEL SESSION

The MDR/IVDR 5-year recertification: Urgent question,

strategic choices

11:50-12:40

PARALLEL SESSION

TUESDAY 13 MAY

18:30-22:00

WELCOME COCKTAIL RECEPTION

WELCON opiformo We are ver



We are very pleased to welcome you to Lisbon and to Pátio da Galé, the chosen venue for the Medtech Forum Welcome Reception.

Located in one of Europe's largest squares, Pátio da Galé was once part of the Royal Palace, blending rich history with modern elegance.

With its grand arcades, open-air courtyards, and views of the Tagus river, Pátio da Galé is truly a unique location for cultural exhibitions or corporate events.

DIRECTION: GALE PATIO, PRAÇA DO COMÉRCIO, 11 00 LISBON, PORTUGAL

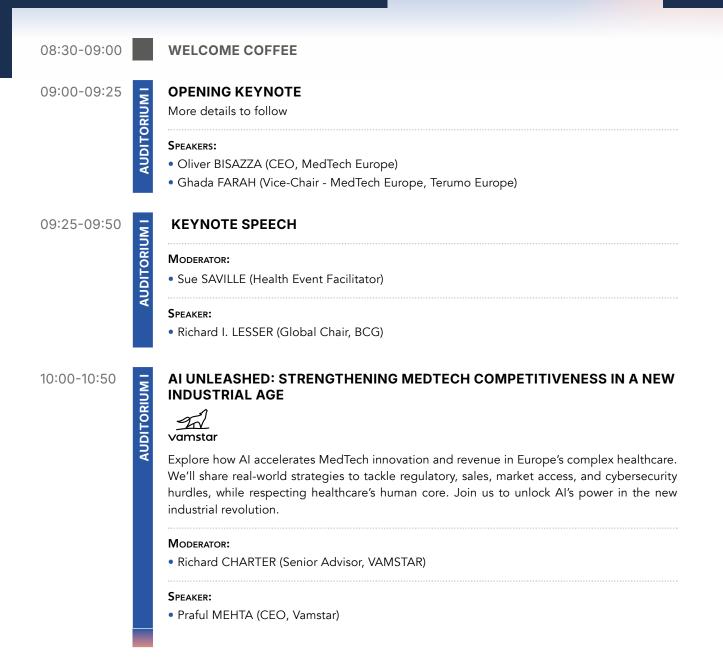
By Car /Taxi / Uber (no more than 7€)

By public transportation:

To get from the Lisbon Congress Center to Pátio da Galé using public transport, follow these steps:

- **Tram:** Walk to the «Centro de Congressos» stop and take tram 15E towards «Praça da Figueira».
- **Stop:** Get off at «Praça do Comércio».
- **Walk:** From Praça do Comércio, Pátio da Galé is just a 2-minute walk away.





AUDITORIUM II

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ROOM

ROOM

10:00-10:50

MEDTECH POLICY TRENDS: NAVIGATING A CHANGING GEOPOLITICAL LANDSCAPE

An insightful session exploring how the evolving geopolitical landscape – the 2024-2029 EU mandate, as well as looking West (the US) and East (China and India) – is shaping the future of the medtech sector. Dive into the opportunities and risks emerging in this dynamic environment and discover how our industry can adapt and thrive.

MODERATOR:

• Christian CLARUS (Director Global Government Affairs, B. Braun Group)

SPEAKERS:

- Benish ASLAM (Lead, Government Affairs and Policy, Asia Pacific Medical Technology Association (APACMed))
- Trevor GUNN (Vice-President International Relations, Medtronic)
- Mathew SHEARMAN (APCO)

FUTURE OF BUSINESS INTERACTION: COMMERCIALIZING DIGITAL SOLUTIONS IN MEDTECH **Deloitte.**

The MedTech industry is shifting towards digital solutions. Healthcare providers seek intelligent, scalable tools and services that leverage innovative tech to improve patient outcomes. Join industry voices to explore opportunities and tensions in this new commercial landscape.

SPEAKERS:

- Ivo ANTÃO (CTO, Luz Saúde)
- Pedro GARCIA DA SILVA (CIO, Champalimaud Foundation)
- Stefan HOLZER (Senior Director, Tis Digital Health Solutions, Terumo EMEA)
- Georg SELLE (Deloitte)

RESILIENT PROCUREMENT, STRONGER HEALTHCARE: THE PROCURE RESULTS

This session will present the final results of the ambitious EU ProCure project aiming to make current public procurement practices on medtech more resilient and efficient all over the EU, and to ensure that public health systems are ready for whatever crises the future brings.

MODERATOR:

• Hans BAX (Senior Advisor Value & Innovation-based Access, MedTech Europe)

- Raquel ARES (Manager, Science and Innovation Link Office (SILO))
- Bérénice CLEUET (European Projects Manager, Reseau des Acheteurs Hospitaliers (RESAH))
- Camille SERRES (European Projects Manager, Reseau des Acheteurs Hospitaliers (RESAH))

10:00-10:50

ASK THE EXPERT: MEDTECH MEETS SOCIAL MEDIA: EU COMPLIANCE UPDATE

Hogan Lovells

AUDITORIUM III

AUDITORIUM

Recent EU enforcement updates highlight increased regulatory scrutiny over how medical device companies interact with healthcare professionals (HCPs) and promote their products on social media – This session will provide some tools to mitigate risks and avoid enforcement actions at the EU Member State level.

SPEAKERS:

- Fabien ROY (Partner Global Regulatory, Hogan Lovells)
- Arne THIERMANN (Hogan Lovells)

11:00-11:50

BOOSTING EUROPE'S ATTRACTIVENESS FOR MEDTECH INNOVATION

Empowering patients, inspiring innovation! Europe is attractive for its innovative research ecosystem, accessible healthcare system and growing efforts towards value-based healthcare. However, action is needed to retain its historic place as global 'epicenter' of medtech innovation. Let's discuss!

SPEAKERS:

- Ben DESMET (Partner Deloitte Life Sciences Practice, Deloitte)
- Clara SATTLER (Head of EMEA, Philips)

TRANSFORMING HEALTHCARE BEYOND HOSPITAL WALLS

Outside-of-hospital and hybrid care models have proved their ability to improve patient outcomes, increase access, and reduce the burden on hospitals and healthcare systems. The session will address the healthcare providers' point of view and practical industry initiatives to scale up these models.

- Nadine JAMOUS (Director, EMEA Market Access, Health Economics & Reimbursement, Zimmer Biomet)
- Frederic NOEL (Vice President WE Enterprise Accounts & EurAsia Integrated Health Solutions, Medtronic)

ROOM

ROOM

11:00-11:50

THE AI EDGE IN MEDTECH COMMERCIAL—INSIGHTS FROM BCG'S 4TH MILKMAN STUDY

Join us as we unveil BCG's latest 'Milkman' Commercial Benchmarking Study, offering fresh insights into the evolution of MedTech commercial models. With Qiagen's Thorsten Harzer, we will explore real-world AI in action and share practical steps toward AI-enabled commercial model transformation.

MODERATOR:

• Goetz GERECKE (Managing Director & Senior Partner, BCG)

SPEAKERS:

- Thorsten HARZER (Head of Digital Business Solutions, Qiagen)
- Basir MUSTAGHNI (Managing Director & Partner, BCG)
- Can SCHNIGULA (Partner, BCG)

PAVING THE WAY FOR THE EUROPEAN HEALTH DATA SPACE: FROM POLICY TO PRACTICE

As the EHDS transitions from policymaking to practical implementation, this session brings together key stakeholders to discuss strategies, challenges, and best practices for its implementation. Panelists will highlight essential factors that accelerate adoption and help realize the full potential of the EHDS.

MODERATOR:

• Petra WILSON (Chair, Digital Health and Innovation Centre in Scotland)

SPEAKERS:

- Enrique MARTINS (Associate Professor in Health Management and Leadership FCS-UBI and ISCTE-IUL (University of Lisbon)
- Amélie SCHÄFER (European Projects Manager, Health Data Hub)
- Bruno SICRE (Director Product Management, Digital Provider Solutions, Resmed)
- Daniela SPIESSBERGER (EU Policy Advisor, Gematik)

ASK THE EXPERT: STATUS OF DIGITAL HEALTH REIMBURSEMENT IN THE US AND EUROPEAN MARKETS



AUDITORIUM

Join market access experts Meike Bomhof and Stephen Hull to explore digital health reimbursement in the USA, UK, France, Germany, and the Netherlands. Get updates on policies, coding, and payer evidence needs for AI, remote monitoring, and robotic tech across hospitals, data centers, and homes.

SPEAKER:

• Stephen HULL (Senior Vice President, Avania Market Access)

AUDITORIUM

11:50-12:20

NETWORKING BREAK

12:20-13:10

FROM EVALUATION TO ACTION: THE FUTURE OF MDR AND IVDR

As we approach the 8th birthday of the MDR and IVDR and take stock of their targeted evaluation, this timely discussion will examine the consensus between actors and stakeholders for short-term solutions, including implementing acts, and the long-term vision for sustainable regulatory improvements.

MODERATOR:

• Bassil AKRA (Chief Executive Officer, AKRA TEAM GmbH)

SPEAKERS:

- Kelvin OKUNDAYE (Accredited assistant to Dr. Peter Liese Member of the European Parliament, Coordinator for the EPP Group in the Committee on the Environment, Public Health and Food Safety, Group of the European People's Party (Christian Democrats))
- Flora GIORGIO (Head of Unit, European Commission)
- Manoja RANAWAKE (Vice President Regulatory Affairs EMEA, WWIPD & IP OUS, BD)
- Rui SANTOS IVO (President, INFARMED)

CANCER PATIENT JOURNEY HOW MEDTECH FOSTERS BETTER AND SAFER CARE

Medical technologies directly affect patient life and are essential for health care professionals in working side by side for cancer prevention, early detection, diagnosis and efficient and safe patient care. In this panel, stakeholders discuss and suggest on how optimizing medtech solutions enables better patient care, accelerates precision oncology and ensures patient safety.

MODERATOR:

Maciej GAJEWSKI (Exact Sciences)

SPEAKERS:

- Antonella CARDONE (CEO, Cancer Patients Europe)
- Dennis A. OSTWALD (CEO, WifOR Institute)
- Luca QUAGLIATA (Vice-President, Medical Affairs Thermo Fisher Scientific)
- Paola THELLUNG DE COURTELARY (Policy and Scientific Manager, European Society of Pathology)

AUDITORIUM II

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ROOM

SOOM

12:20-13:10

FROM HYPE TO REALITY: UNLOCKING AI'S POTENTIAL IN MEDICAL DEVICES

zühlke

Al offers immense potential for medical devices, from early disease detection to robotic surgery. Yet, clinical deployment remains challenging. This talk explores how best practices, risk-based approaches, and regulatory alignment can unlock Al's benefits while ensuring safety and reliability.

SPEAKER:

Gabriel KRUMMENACHER (Director Data Science, Zuehlke)

NAVIGATING GLOBAL BUSINESS CHALLENGES IN MEDTECH OPERATIONS TO PROVIDE CONSISTENT PATIENT CARE



In today's landscape, healthcare organizations face unique challenges, such as evolving trade policies, supply chain disruptions, technology advancements and global market expansion. Industry leaders will discuss strategies and insights for designing MedTech operations that address these challenges and ensure seamless global patient care.

MODERATOR:

• Jesus RUEDA RODRIGUEZ (Director General - Strategies, Special Projects & International Affairs - MedTech Europe)

SPEAKERS:

- Christian CLARUS (Director Global Government Affairs B. Braun Group)
- Daniele FAZIO (VP & GM, Medical Devices and Drug Delivery, Flex)
- Wilfred VAN ZUILEN (Zimmer Biomet)

ASK THE EXPERT: AI-DRIVEN TRANSLATION IN MEDTECH: THE POWER OF QUALITY ESTIMATION AND HUMAN EXPERTISE

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AUDITORIUM III

Discover how Al-driven translation technology is transforming the Medical Devices industry by ensuring compliance and efficiency. This session explores Al-powered machine translation, automatic quality estimation (AQE), and expert post-editing. Learn how glossaries, quality frameworks, and Al optimize workflows with real-world case studies.

SPEAKER:

• María ILLESCAS (Director of Innovation and AI Solutions, SEPROTEC)

13:10-14:10

NETWORKING LUNCH

14:10-15:00

SAFE AI, SMARTER CARE: TRANSFORMING EUROPEAN HEALTHCARE SYSTEMS

The panel seeks to bring together stakeholders from across the healthcare ecosystem to discuss how the safe uptake of AI can revolutionise healthcare delivery in Europe.

The panel will gather experts from hospitals, healthcare professionals, industry and academia to provide for a multi-perspective discussion.

SPEAKERS:

- Mariana MADUREIRA (Coordinator Medical Device Sector, INFARMED)
- Eric SUTHERLAND (OECD)
- Véronique TORDOFF (Image-Guided Therapy Leader Europe, Philips)

EMPOWERING EUROPEAN INNOVATORS WITH EARLY FEASIBILITY STUDIES

HEU-EFS is an IHI project which aims at developing an EU harmonised framework to improve the uptake of Early Feasibility Studies (EFS) for medical devices in the European Union. EFS are a crucial part of the evidence-generation cycle for medical devices. Conducting these studies in a coordinated manner across Europe will bring significant benefits to patients, innovators, and the entire healthcare system.

MODERATOR:

• Niklas BLOMBERG (Executive Director, Innovative Health Initiative (IHI))

SPEAKERS:

- Alexandra POULSSON (Senior Advisor, Division for Health Services Norwegian Institute of Public Health)
- Laura SAMPIETRO-COLOM (Innovation Deputy Director Hospital Clinic Barcelona)
- Rosanna TARRICONE (Associate Dean at SDA Bocconi Associate Professor at the Department of Social and Political Sciences)
- Fanny VAN DER LOO (Senior Director Government Affairs EMEA, Canada, Latin America Edwards Lifesciences)
- Yasemin ZEISL (Project Coordinator European Patients' Forum (EPF))

UK - EU: READY FOR A RESET?

A reset in the relationship between the UK and the EU is underway, with agreement that leaders will meet in an EU-UK Summit. What can realistically be expected from the reset and what it could imply for market access for medical technologies? This session will explore the current market access environment in the UK and what opportunities the reset of the bilateral relations could bring for the medtech industry.

SPEAKER:

ROOM A

Peter ELLINGWORTH (Chief Executive - ABHI)

AUDITORIUM I

ROOM

AUDITORIUM III

AUDITORIUM I

14:10-15:00

FROM DATA TO DIAGNOSIS: DIGITAL TOOLS TRANSFORMING DIAGNOSTICS

Join us to explore the transformative potential of digital solutions in the IVD sector for clinical evidence generation. These technologies enable efficient data collection, analysis, and validation, improving diagnostic accuracy and supporting personalized medicine. Let's discuss how MedTech Europe members can shape these trends.

SPEAKER:

Chaohui GUO (Head of Clinical Validation, Roche)

ASK THE EXPERT: ARE YOU WINNING OR LOSING MARKET SHARE? THE VALUE OF PRECISE MARKET DATA

Ask the expert session on MTE's flagship publication and golden standard data trackers. The place to ask direct questions comming from all type of data users and stakeholders. Explore the value of precise, actionable market data in shaping business strategies. Learn how MedTech Europe's data services provide companies with the tool to uncover trends, benchmark performance, and identify growth opportunities.

SPEAKERS:

- Teodora ANGELOVA (Senior Manager Market Data, MedTech Europe)
- Georgiy BOGDANOV (Manager Market Data, MedTech Europe)

15:10-16:00

CLIMATE MEETS ECONOMICS: BRIDGING GREEN GOALS AND COMPETITIVE EDGE IN HEALTHCARE

Are decarbonisation and competitiveness friends or foe? Do the Draghi Report and EU Clean Industrial Deal deliver for MedTech? How to leverage the power of medtech for healthcare system transformation and competitiveness?

MODERATOR:

• Sue SAVILLE (Health Event Facilitator)

- Martin FUHRER (Senior Vice President Head of Diagnostics Central & Eastern Europe, Central Asia; member of MedTech Europe Board, Siemens Healthcare Diagnostics GmbH)
- Véronique TORDOFF (Image-Guided Therapy Leader Europe, Philips)

ROOM A

15:10-16:00

HTAR IN MEDICAL DEVICES - BOOSTING OR STIFLING INNOVATION IN EUROPE

Boosting or stifling innovation in Europe? This session will provide timely insights on the status of implementation of the Health Technology Assessment Regulation (HTAR) four months after its date of application with a focus on its impact on the availability of innovation in Europe.

MODERATOR:

• Andrea RAPPAGLIOSI (Senior Vice President Public Affairs EMECLA, Edwards Lifesciences)

SPEAKERS:

- Burçak AYDIN (Senior Manager HAG INSIGHT Consortium)
- Marco MARCHETTI (Vice Chair HTA Coordination Group / Direttore UOC HTA Agenas)
- Leslie PIBOULEAU (Directorate-General for Health and Food Safety, European Commission)
- Piotr SZYMAŃSKI (Chairman of the Regulatory Affairs Committee at European Society of Cardiology, Consultant Cardiologist and Head of the Clinical Cardiology Department - CSK MSWiA)

SHAPING THE FUTURE OF CLINICAL INVESTIGATIONS IN EUROPE

In light of the pilot coordinated assessment for clinical investigations of medical devices (MDR Art.78) which has been launched in early February by the Member States and the European Commission, this session will take stock of the process so far, clarify the possibilities offered to sponsors by this pilot and answer any remaining questions. The session will also look to the future at next steps.

MODERATOR:

Carine COCHEREAU (Integra)

- Benedicte NUYTTENS (Head of Clinical Investigation Entity, Federal Agency for Medicines and Health Products)
- Nebojsa SERAFIMOVIC (Senior Assessor for Medical Devices, Austrian Federal Office for Safety in Health Care)

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ROOM

15:10-16:00

EU AND MDSAP - A DREAM OR A VIABLE OPPORTUNITY?

The Medical Device Single Audit Program (MDSAP) is arguably the landmark example of international regulatory cooperation. Learn why it is so timely for the EU to join it as a full member, accelerating European patients' access to medical technologies and fostering EU's competitiveness and innovation.

MODERATOR:

• Olga VAN GROL-LAWLOR (Senior Global Regulatory Intelligence & Advocacy Manager, Boston Scientific)

SPEAKERS:

- Oliver BOHLE (Regulatory Affairs Manager, Spectaris)
- Tracey DUFFY (First Assistant Secretary, Australian Government, Department of Health and Aged Care)
- Flora GIORGIO (Head of Unit, European Commission)
- Graeme TUNBRIDGE (Senior Vice President Global Regulatory and Quality, BSI)

ASK THE EXPERT: NAVIGATING U.S. ANTI-BRIBERY COMPLIANCE UNDER THE TRUMP ADMINISTRATION

ReedSmith Driving progress through partnership

This session focuses on how multinational companies can manage risk related to the U.S. Anti-Kickback Statute, False Claims Act, and Foreign Corrupt Practices Act in light of the changing priorities under the Trump Administration, and how understanding U.S. enforcement activity can help companies prepare for how similar trends may play out in Europe.

SPEAKERS:

- Ali ISHAQ (Partner, ReedSmith)
- Rosanne KAY (Partner, ReedSmith)
- Jeff LAYNE (Partner, ReedSmith)

16:00-16:30

NETWORKING BREAK

AUDITORIUM I

AUDITORIUM I

ROOM A

16:30-17:20

NAVIGATING SUSTAINABLE CORPORATE GOVERNANCE: CHALLENGES AND OPPORTUNITIES

Join us for a session on implementing the CSRD and CSDDD in MedTech. Explore challenges, opportunities, and potential changes amid economic and global competition. Gain insights on staying ahead in sustainability while remaining competitive. Engage with experts on the future of sustainable corporate governance.

MODERATOR:

• Céline FAEH (Director Government Affairs EMEA - Zimmer Biomet)

SPEAKERS:

- Alexandre BARROS (Hydrumedical)
- Celia DOUSSINEAU (Siemens Healthineers)
- Greta KOCH (Parliamentary Assistant to MEP Axel Boss, European Parliament)

WHAT IF WE DESIGNED HEALTHCARE FOR WOMEN?

Challenging the systems, science, and silence that have failed women: thought leaders explore how medtech can advance diagnostics, personalized care, reproductive health, and gender equity—driving solutions through collaboration.

MODERATOR:

Peter ELLINGWORTH (Chief Executive, ABHI)

SPEAKERS:

- Tanja BRYCKER (Vice President, Strategic Development, Breast & Skeletal Health and Gynae Surgical Solutions, Hologic)
- Bejal PANDYA (Congenital Cardiologist & Clinical Director of Specialised Cardiology, Barts Heart Centre)
- Nina WILSON (CEO & Founder, One Women Health)

EUDAMED: THE CENTRAL APPROACH FOR GATHERING DATA ON MEDICAL DEVICES

The European Medical Device Database EUDAMED will become the mandatory IT system to use for the MDR and IVDR in only 6 months' time after this session. What changes and new considerations will this central approach bring to manufacturers and other users in data submission and data access?

- Flora GIORGIO (Head of Unit, European Commission)
- Silvia OSTUNI (Legal and Policy Officer, DG SANTE, Medical Device Unit)

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ROOM

16:30-17:20

TIME TO ACT: THE NEED FOR ACCELERATED VALUE PATHWAYS FOR BREAKTHROUGH INNOVATIONS

Edwards

This panel explores Europe's next steps to accelerate patient access to breakthrough innovations through early evidence, streamlined regulations, adaptive HTA, and innovative procurement using Real-World Evidence.

MODERATOR:

• Andrea RAPPAGLIOSI (Senior Vice President Public Affairs EMECLA - Edwards Lifesciences)

SPEAKERS:

- Marco MARCHETTI (Vice Chair HTA Coordination Group / Direttorre UOC HTA, Agenas)
- Laura SAMPIETRO-COLOM (Innovation Deputy Director Hospital Clinic Barcelona)
- Piotr SZYMANSKI (Chairman of the Regulatory Affairs Committee at European Society of Cardiology - Consultant Cardiologist and Head of the Clinical Cardiology Department - CSK MSWiA)
- Rosanna TARRICONE (Associate Dean at SDA Bocconi Associate Professor at the Department of Social and Political Sciences)

ASK THE EXPERT: THE US REGULATORY LANDSCAPE – MAKING SENSE OF THE CURRENT CHAOS

Hogan Lovells

AUDITORIUM

Between staff reductions, executive orders, and tariffs, it has been a tumultuous time at the FDA. In this session, hear the key initiatives impacting the US FDA are and how these could affect the regulation of medical devices in the US. Discuss best practices to continue doing business in the US.

SPEAKER:

• Michael HEYL (Partner Global Regulatory, Hogan Lovells)

AUDITORIUM I

17:30-18:10

PIONEERING FUSION RESEARCH – INSIGHTS FROM THE CHAMPALIMAUD FOUNDATION

In her keynote address Leonor Beleza will introduce the pioneering work of the Champalimaud Foundation and its distinctive concept of fusion research—an integrated model that unites advanced biomedical research with translational, interdisciplinary clinical care.

Her remarks will highlight the Foundation's commitment to innovation, scientific excellence, and sustainability in healthcare, offering a source of inspiration for those shaping the future of care. Drawing from her extensive career, Leonor Beleza will share key reflections on leadership, research-driven impact, and the evolving role of global collaboration in healthcare advancement.

The keynote will be followed by a fireside chat with Oliver Bisazza, providing further perspectives on the Foundation's mission and the broader implications of its work for the medical technology and research communities.

SPEAKERS:

- Leonor BELEZA (President, Fundacao Champalimaud)
- Oliver BISAZZA (CEO, MedTech Europe)

18:10-19:40

NETWORKING RECEPTION



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AUDITORIUM I

AUDITORIUM I

08:00-08:30

WELCOME COFFEE

08:30-09:20

CEO #NOFILTER

A high-level discussion with executives from four leading companies on trends and key issues, including priorities in serving patients and customers, how to speed up care pathways, navigating the European regulatory landscape, encouraging investment in innovation and how to attract top talent.

MODERATOR:

• Sue SAVILLE (Health Event Facilitator)

SPEAKERS:

- Annette BRÜLS (Corporate Vice-President, EMEACLA, Edwards Lifesciences)
- Roland GOETTE (Executive Vice President EMEA, BD)
- Urmi Prasad RICHARDSON (President EMEA, Thermo Fisher Scientific)
- Gavin WOOD (Company Group Chairman, Johnson & Johnson MedTech EMEA)

09:30-10:20

WHY EUROPE MATTERS: EXPLORING MEDTECH INVESTMENT POTENTIAL

How can Europe remain an attractive destination for MedTech investments? Hear from top investors as they explore the sector's potential and what it needs to thrive in a competitive global market.

MODERATOR:

• Stephen LEVIN (Editor in chief, MedTech Strategist)

- Xavier BERTRAND (SVP & President, EMEA, Boston Scientific BSCI)
- Craig COOPER (Partner, Serpentine ventures)
- Janke DITTMER (General Partner, Gilde Healthcare)
- Giovanni MONTI (Founder, Option 5 Health Ltd)

AUDITORIUM II

ROOM A

ROOM

09:30-10:20

DRIVING SECURE HEALTH DATA ACCESS VIA THE IDERHA IHI PROJECT

The IDERHA (http://www.iderha.org/) consortium will create a federated data space for secure health data access and develop policy recommendations for RWE acceptance in regulatory decision making. The panel will explore the importance and opportunities of health data access to enable innovative healthcare solutions.

MODERATOR:

• Hugh LAVERTY (Head of Scientific Operations, Innovative Health Initiative (IHI))

SPEAKERS:

- Flora GIORGIO (Head of Unit, European Commission)
- Philip GRIBBON (Fraunhofer ITMP)
- Christian MUEHLENDYCK (Scientific Partnerships Lead, Johnson&Johnson MedTech EMEA, Germany)
- Gözde SUSUZLU (Senior Programme Manager, The European Patient Forum)

PROTECTING PRIVACY IN THE AGE OF AI-POWERED HEALTHCARE

Delve into the critical role of privacy in the rapidly evolving landscape of Al-driven healthcare. This session will explore the intersection of cutting-edge digital health innovations and robust privacy frameworks, including the EU Al Act and GDPR. Learn how to implement ethical Al practices and privacy-preserving technologies to build trust and drive compliant innovation.

MODERATOR:

Peter BLENKINSOP (Partner Faegre Drinker)

SPEAKERS:

- Benjamin MEANY (Senior Manager Digital, Software and AI Regulation, MedTech Europe)
- Chantal VETS (Sr. Legal Program Director, Digital Regulations Medtronic)

SECURE AND SEAMLESS: THE NEXT GENERATION OF TECHNICAL DOCUMENTATION

This panel will delve into modernising Technical Documentation through a harmonised, itemlevel approach, enabling secure, system-agnostic data exchange with standardised formats and nomenclature. The discussion will highlight the importance of supporting SMEs by removing financial barriers, while preserving flexibility for manufacturers, paving the way for sustainable and forward-thinking regulatory practices.

MODERATOR:

James SHEARN (Director Regulatory & Quality Compliance, STERIS)

- Tanja KNAUER (Vice President, Regulatory Affairs & Standards and Digital Operations and Strategy, Siemens Healthineers)
- Sandy WRIGHT (Head of Devices, Scarlet)

AUDITORIUM III

AUDITORIUM

09:30-10:20

ASK THE EXPERT: INNOVATIVE PAYMENT SCHEMES – LESSONS LEARNED FROM PHARMA AND MEDICAL (DIGITAL) DEVICES

MedTech Europe's Reimbursement and Funding Working Group has carefully developed twenty 'Key Success factors' and relevant metrics to operationalise and allow the qualitative assessment of the suitability and effectiveness of IPS across Europe. During the session, the speakers will present the ey findings.

SPEAKERS:

- Hatim ABDULHUSSEIN (Chief Executive Officer Health Innovation Kent Surrey Sussex, NHS)
- Oriana CIANI (Associate Professor of Practice, SDA Bocconi School of Management)
- Jorge Juan FERNÁNDEZ GARCÍA (Chief Innovation Officer, Hospital Clínic de Barcelona)

10:20-10:50

10:50-11:40

NETWORKING BREAK

RE-IMAGINING CYBER SECURE HEALTH

Multistakeholder panel discussion: representatives from hospitals, industry and regulators will debate cybersecurity in healthcare, aiming to share best practices and insights on how to reimagine the health ecosystem to make sure it is cybersecure.

MODERATOR:

Miroslav PALAT (President, CzechMed)

SPEAKERS:

- Manan HATHI (Senior Manager, Digital Health Regulatory Policy and Intelligence, Stryker)
- Saila RINNE (Head of Unit, eHealth, Well-Being and Ageing, Directorate-General Communications Networks, European Commission)
- Lino SANTOS (National Coordinator, Portuguese National Cybersecurity Centre (CNCS))

NAVIGATING LITIGATION: SECURING MEDTECH'S FUTURE IN EUROPE

Explore Europe's evolving litigation landscape, from class actions under the Representative Actions Directive to the impact of Third Party Litigation Funding. Learn proactive strategies to navigate these challenges and ensure resilience in a rapidly changing legal environment.

SPEAKERS:

- Carolyn BLAKE (European Policy Consultant, U.S. Chamber of Commerce, Institute for Legal Reform)
- Brian KRUID (Senior Litigation Counsel, International Stryker)
- Patrick REILLY (Partner Faegre Drinker Biddle & Reath LLP)
- Sofia VAZ SAMPAIO (Morais Leitão)

AUDITORIUM

21 | PROGRAMME

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ROOM

10:50-11:40

GLOBAL MARKET FOCUS: BRAZIL

Brazil is projected a solid growth of medical technology market over the next five years. Its evolving healthcare market, particularly for innovative technologies presents a wealth of opportunities for medtech manufacturers. In this session, experts will explore the trends and policies driving the growth of the medtech market, discuss the challenges and opportunities and what it takes to succeed in Brazil.

MODERATOR:

• Carlos GOUVEA (Executive President, Câmara Brasileira de Diagnóstico (CBDL))

SPEAKERS:

- Kristin CIRIELLO POTHIER (Life Sciences Sector Leader, Global Deal Advisory and Strategy Leader, Healthcare and Life Sciences, KPMG LLP)
- Nataly TREJOS (Associate Director, LATAM Regulatory Policy Johnson&Johnson MedTech)

ALIGNING PLM & QMS PROCESSES FOR MEDTECH EXCELLENCE

Veeva MedTech

Unlocking product lifecycle excellence starts with aligning PLM and QMS. Learn how MedTech companies streamline operations, enhance product integrity, and accelerate time-to-market with a process-first approach balancing innovation, safety, and efficiency.

SPEAKERS:

- Mandy BLOCHER (Head of PLM Digitalization & Regulatory Intelligence, B.Braun (Division Aesculap))
- Arielle FAKHRAEE (Director, MedTech Quality Strategy, Veeva Systems)

ASK THE EXPERT: EVERYTHING YOU WANTED TO KNOW ABOUT IHI (BUT WERE AFRAID TO ASK)

IKOP, IKAA, Financial Contribution, 3A? If it does not ring a bell, join this session! IHI projects offer many benefits such as funding and networking, to name a few. People participating in the first projects will highlight why they have joined this public-private partnership on health innovation.

SPEAKERS:

- Hugh LAVERTY (Head of Scientific Operations, Innovative Health Initiative (IHI))
- Peter ZANDBERGEN (Director Public-Private Partnerships, Innovation Partnerships Europe Innovation & Strategy, Philips)



ROOM

AUDITORIUM I

AUDITORIUM

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ROOM

11:50-12:40

IMPLEMENTING THE EU GREEN DEAL IN HEALTHCARE

Tackling Challenges, Seizing Opportunities - How is the medical technology sector navigating the Green Deal maze? How to connect sustainability dots for better health and patient outcome? What's coming next?

MODERATOR:

• Sue SAVILLE (Health Event Facilitator)

SPEAKERS:

- Bill DOHERTY, Executive Vice President, COOK MEDICAL
- Bert HARTOG (Member of the Advisory Board, DiCE)
- Maureen MAZUREK (Chief EHS & Sustainability Officer, BD)

CLOSING THE LOOP: TURNING CUSTOMER INSIGHTS INTO PRODUCT EXCELLENCE

smarleeva

Join us to explore the transformative potential of digital solutions and AI for enhanced product performance, customer experience, safety, and post-market surveillance. Discover impactful projects and discuss how the MedTech Industry can shape these trends.

SPEAKERS:

- Michael SCHAETZLE (Head of Data Insights and Process Excellence Post-Market Quality, Roche Diagnostics)
- Raffel ZUBER (Data Scientist, Roche Diagnostics)

ADVANCING THE ASSESSMENT OF DIGITAL HEALTH TO FOSTER INNOVATION

The focus of this session will be to shine a spotlight on EU-level initiatives currently supporting the development of inclusive and fit-for-purpose European assessment frameworks for digital health technologies. Understanding the interplay between different initiatives and their potential to be effectively applied in practice is expected to advance the assessment of digital health to foster timely access to innovation for European patients.

MODERATOR:

• Katarzyna MARKIEWICZ-BARREAUX (AI Strategic Intelligence Lead, Philips)

- Rubén CASADO ARROYO (Healthcare Professional, Université Libre de Bruxelles-Erasme Hospital)
- Marco MARCHETTI (Vice Chair HTA Coordination Group / Direttore UOC HTA, Agenas)
- Juan Carlos REJON PARRILLA (Researcher, Andalusian HTA Agency (AETSA))
- Emmanouil TSIASIOTIS (Academia representative, Graduate school of health economics and management (ALTEMS), Università Cattolica del Sacro Cuore)

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ROOM

AUDITORIUM

AUDITORIUM I

11:50-12:40

SHAPING THE FUTURE OF WOUND CARE

The session will examine the burden of wounds on the healthcare systems and policy trends impacting wound management. How better wound care policy can address inequalities in healthcare and drive sustainable healthcare.

MODERATOR:

Judith HARGREAVES (Molnlycke)

SPEAKER:

• Andrea KEADY (Solventum)

ASK THE EXPERT: HOW AI & PLM DRIVE SMARTER COMPLIANCE IN MEDTECH



MedTech companies face rising pressure from fast-evolving global rules. Join Deloitte & PTC to see how AI and PLM turn compliance into an asset - enabling smarter decisions, continuous monitoring, and full lifecycle coverage. Move from caution to confidence.

SPEAKER:

• René ZOELFL (Senior Director, Global Advisor of MedTech PTC)

12:40-13:40

13:40-14:30

NETWORKING LUNCH

INNOVATING FOR PLANETARY AND HUMAN HEALTH: SUSTAINABILITY TRENDS AT GLOBAL LEVEL

Environmental challenges do not halt at national or regional level while supply chains in the medical technology sector are globally intertwined. As many governments are looking to advance sustainability in the healthcare sector, this session aims at providing a snapshot of common trends.

MODERATOR:

• Sue SAVILLE (Health Event Facilitator)

- Benish ASLAM (Regional Lead, Government Affairs and Policy and ESG, Asia Pacific Medical Technology Association (APACMed))
- Addie MACGREGOR (Sustainability Manager, ABHI)
- Elena VILLALOBOS PRATS

AUDITORIUM II

ROOM A

13:40-14:30

CUTTING EDGE: THE FUTURE OF ROBOTIC SURGERY

The value of Robotic-assisted surgery (RAS) will be the centre of the discussions among experts and stakeholder organisations. Speakers will evaluate the unmet healthcare needs of European citizens and discuss strategic policy initiatives to address them.

MODERATOR:

• Laura HAMPUNEN (Stryker)

SPEAKERS:

- Dirk GHADAMGAHI (Head of Medical Affairs Johnson&Johnson MedTech EMEA)
- Kris MAES (Director of the Urology Service and Coordinator of the Centre for Minimally Invasive and Robotic Surgery, Hospital da Luz, Lisbon)
- Franca MELFI (Prof. of Thoracic Surgery, University of Calabria)
- James PORTER (Chief Medical Officer Robotic Surgery and Digital Technology, Medtronic)

A NEW CARDIOVASCULAR HEALTH PLAN FOR EUROPE, A GAME CHANGER FOR PATIENTS

The session will be entitled "A new Cardiovascular Health Plan for Europe, a game changer for patients", and will bring together key representatives from the cardiovascular community, patient organizations, industry, and policymakers. The discussion will explore how to accelerate early detection, innovation, and equitable access to treatment, and how these elements can be integrated into the upcoming EU Cardiovascular Health Plan.

MODERATOR:

Alexander OLBRECHTS (Director Digital Health, MedTech Europe)

- Birgit BEGER (CEO, European Heart Network)
- Marta BRAGAGNOLO (EU Projects Manager, Global Heart Hub)
- Frederic CLEMENT (Patient Care Solutions: International Chief Marketing & Strategy Officer, GE Healthcare)
- Aoife DELMAS (Chief Development Officer European Society of Cardiology)

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ROOM

13:40-14:30

FAST-TRACK HEALTH TECHNOLOGY ASSESSMENT FOR IN VITRO DIAGNOSTICS

apifarma

The evolution of in vitro diagnostics (IVDs) underscores the need for an optimized health technology assessment (HTA) framework. A BMJ Innovations study used design thinking to engage stakeholders, proposing a fast-track HTA system to enhance IVD evaluation, adoption, and healthcare efficiency.

MODERATOR:

• Fernando ALMEIDA (President, INRJ)

SPEAKERS:

- Carlos CATALÃO (Medical and Health Policy Director, Roche)
- Luis MIGUEL SANTIAGO (MD, PhD Professor and Researcher, Faculdade de Medicina da Universidade de Coimbra - Centro de Estudo e Investigação em Saúde da Universidade de Coimbra (CEISUC))
- Tamara MILAGRE (EVITA)
- Helena MONTEIRO (Director of the Medical and Digital Health Technologies Unit, INFARMED)
- Guilherme VITORINO (Vice-Director of NOVA IMS, UNIVERSIDADE NOVA)

ASK THE EXPERT: UNLOCKING HEALTH DATA: THE EU DATA ACT'S IMPACT ON DIGITAL HEALTH

ReedSmith

AUDITORIUM III

Driving progress through partnership

The EU Data Act aims to foster fair and secure data sharing across the EU. It will affect any connected medical device or health wearable that generates or collects data from its users and where those services are supported by the cloud, could implicate cloud portability requirements under the Act. In this presentation, we will explore the main features and implications of the Data Act for the digital health sector, as well as the key issues and challenges that need to be addressed, such as deadlines, obligations towards patients/users, compliance requirements.

- Friederike WILDE-DETMERING (Counsel, ReedSmith)
- Cynthia O'DONOGHUE (Partner, Reed Smith LLP)

AUDITORIUM I

AUDITORIUM

14:40-15:30

THE MDR/IVDR 5-YEAR RECERTIFICATION: URGENT QUESTION, STRATEGIC CHOICES

As MDR and IVDR certificates near expiry, a system-wide bottleneck looms. Can the current framework withstand the pressure, or are we heading toward gaps in patient access to life saving devices? This high-level panel explores the strategic and urgent questions behind the 5-year recertification cycle, at the heart of the EU's ongoing regulatory evaluation.

MODERATOR:

• Amanda MAXWELL (European Regulatory Affairs Editor, Citeline)

SPEAKERS:

- Bassil AKRA (Chief Executive Officer AKRA TEAM GmbH)
- Frank MATZEK (Vice President, Biotronik)
- Alexey SHIRYAEV (President Team-NB)

ENGAGING WITH IMPACT: STORIES OF MEDTECH AND PATIENT COLLABORATION

This plenary session highlights powerful examples of how medtech companies are engaging with patients and patient associations to drive innovation and improve outcomes. Hear firsthand from company representatives and patient advocates as they share their stories, challenges, and the power of working together to shape a patient-centered future.

MODERATOR:

• Sandra SA (Patient Experience and Partnership Lead, Roche)

- Kalina BOZHKOVA (Director, European Alliance for Patient Access)
- Hugh LAVERTY (Head of Scientific Operations, Innovative Health Initiative (IHI))
- Roi SHTERNIN (Founder & Chief Patient Officer, Chronically)

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MOOS

14:40-15:30

NAVIGATING INTEROPERABILITY IN THE EHDS: ROLES AND RESPONSIBILITIES FOR MANUFACTURERS

This session explores new interoperability requirements introduced by the EHDS, examining their impact on digital health solutions and medical technologies, and highlighting the importance of collaboration with standards organizations such as IHE to ensure seamless interoperability in this new health data ecosystem.

MODERATOR:

• Shweta BHARDWAJ (Director, Global R&D and Digital Policy, Johnson and Johnson, Johnson & Johnson)

SPEAKERS:

- Alexander BERLER (Strategic Business Development Director at IHE Catalyst AISBL, HL7 Hellas)
- Sabine DÖRHÖFER (Business Owner Standardization Roche Diagnostics International)
- Andreas NEOCLEOUS (Project Manager, National eHealth Authority Cyprus)
- Janos VINCZE (Vendor Co-Chair, IHE Europe)

MEASURE YOUR READINESS WITH THE SELF-ASSESSMENT TOOL

This session will introduce a self-assessment tool for medical technology suppliers to assess their internal readiness to engage with Value-Based Healthcare systems, co-developed by Prof. Laing (Swansea University) and a MedTech Europe ad hoc working group.

MODERATOR:

Klaus ANDERSEN (Managing Partner, Aquilo Consulting)

- Stephanie FRIDD (Director Value Based Care, Philips)
- Dominique GILSOUL (Global Market Access Leader, Solventum)
- Hamish LAING (Director, Swansea University Value-Based Health and Care Academy)

AUDITORIUM III

AUDITORIUM I

14:40-15:30

ASK THE EXPERT: REMOVING THE GUESSWORK FROM IVD MARKET INTELLIGENCE

Ask the expert on the best in-class report and data collection for the IVD industry. Participants can discover the transformative value of precise, actionable market data in driving business success. Gain insights into how MedTech Europe's data services empower companies to uncover emerging trends, benchmark performance against peers, and identify untapped growth opportunities—all while shaping effective, forward-thinking strategies. Don't miss this opportunity to learn from the leaders in market data excellence!

SPEAKER:

• Teodora ANGELOVA (Senior Manager Market Data, MedTech Europe)

15:40-16:00

CONCLUSIONS

SPEAKERS:

• Oliver BISAZZA (CEO, MedTech Europe)

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