ONLINE
05TH-06TH DECEMBER 2024

15:00CET YouTube

2ND ANNUAL MEDICAL DEVICES REGULATIONS & SAFETY 2024 CONFERENCE

VNA INTERNATIONAL
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Industry Events 10times

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Welcome to the 2nd Annual Medical Devices Regulations & Safety 2024 Conference!

We are thrilled to bring together industry leaders, regulatory experts, and stakeholders for two days of insightful discussions, collaborative learning, and networking opportunities. Throughout the conference, attendees will gain valuable insights into the latest regulatory updates, emerging safety standards, and innovative strategies for compliance.

- 8+ DIFFERENT TOPICS
- Q&A SESSIONS
- PANEL DISCUSSION
- POST-CONFERENCE MATERIALS
- CERTIFICATE OF ATTENDANCE

From risk management methodologies to quality assurance best practices, our diverse range of sessions will equip you with the knowledge and tools needed to navigate the complex landscape of medical device regulations. Join us as we explore the future of medical device safety and regulation together!

Key poins:

- Updates on medical device regulations and safety standards.
- Discussions on improving safety practices.
- Strategies for navigating regulatory complexities.
- Effective risk mitigation techniques.
- Ensuring product integrity through quality assurance.
- Exploring the impact of emerging technologies on regulations.
- Networking opportunities and collaboration for industry professionals and regulators.

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The 2nd Annual Medical Devices Regulations & Safety 2024 Conference is tailored for professionals across the medical device industry seeking to enhance their understanding of regulatory compliance and quality assurance practices:

- Medical device manufacturers and suppliers: Engaged in the production and distribution of medical devices.
- Regulatory affairs professionals: Responsible for ensuring compliance with regulations and standards.
- Quality assurance and quality control personnel: Tasked with maintaining product quality standards.
- Healthcare professionals: Interested in understanding regulatory requirements impacting patient care.
- Researchers and innovators: Developing new medical technologies and curious about regulatory implications.
- Government regulators: Overseeing compliance with medical device regulations.
- Legal and compliance professionals: Guiding organizations through regulatory landscapes.
- Consultants and advisors: Providing expertise in navigating regulatory challenges in the medical device industry.

This comprehensive gathering ensures insightful discussions and valuable networking opportunities across various domains.

Expected Speakers:

- **Medtronic:** A leading medical device company known for its innovative technologies in areas such as cardiac and vascular health, diabetes management, and surgical solutions.
- **Johnson & Johnson:** A multinational corporation with a diverse range of medical devices, pharmaceuticals, and consumer healthcare products.
- **Abbott Laboratories:** Known for its wide range of medical devices, diagnostics, nutrition, and pharmaceutical products, Abbott is a global healthcare company committed to improving people's lives.
- **Boston Scientific:** Specializing in medical devices for interventional medical specialties, Boston Scientific focuses on innovations for cardiology, urology, endoscopy, and neuromodulation.
- **Siemens Healthineers:** A leading medical technology company offering products and services in diagnostic imaging, laboratory diagnostics, and advanced therapies.

Benefits of attendance:

- Meet new connections in groups or one-on-one;
- Further develop and strengthen your network;
- Expand your knowledge
- Stay up to date
- Grow your personal brand
- Enjoy, create memories and have fun

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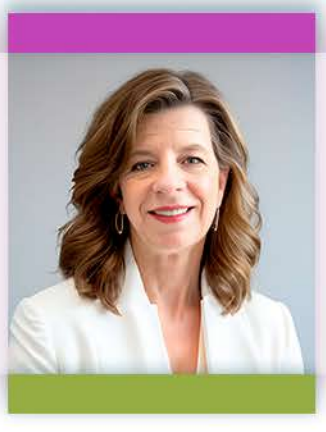
Speakers

Richard Houlihan

CEO

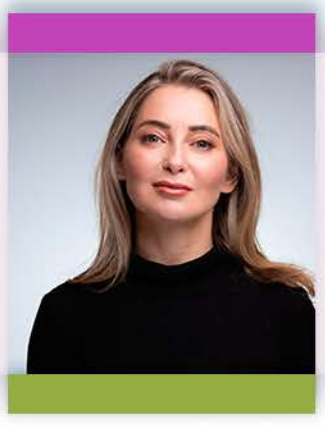
EirMed



Ann Vu

JD, RAC
Sr. VP, Quality, Regulatory
and Clinical
(NASDAQ: ZIMV)

ZimVie



Jacqueline van Druten

(MICR.CIM.RD)
Clinical & Regulatory
Affairs Director

CLINR+

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Day 1

15:00-15:05 Opening Remarks

15:05-15:30 **"Recent Updates in Medical Device Regulations: Global Perspectives"**

15:30-15:35 Q&A Session (5 mins)

15:35-16:00 **Richard Houlihan**
"EUDAMED is nearly here, what now?"

16:00-16:05 Q&A Session (5 mins)

16:05-16:15 Coffee Break (10 mins)

16:15-16:40 **Ann Vu**
"Navigating Regulatory Complexities: Strategies for Compliance Success"

16:40-16:45 Q&A Session (5 mins)

16:45-17:10 **"Effective Risk Management in Medical Device Development and Deployment"**

17:10-17:15 Q&A Session (5 mins)

17:15-17:45 **Panel Discussion:**
"Collaborative Approaches to Addressing Regulatory Challenges in the Medical Device Industry: Perspectives from Manufacturers, Regulators, and Healthcare Providers"*

Day 2

15:00-15:05 Opening Remarks

15:05-15:30 **"Ensuring Product Integrity: Quality Assurance in the Medical Device Industry"**

15:30-15:35 Q&A Session (5 mins)

15:35-16:00 **Jacqueline van Druten**
Ace your submission - Focus on EU MDR Clinical Evaluation Conformity routes. What you need to know for your legacy and new products.

16:00-16:05 Q&A Session (5 mins)

16:05-16:15 Coffee Break (10 mins)

16:15-16:40 **"Patient Safety and Regulatory Compliance: Key Considerations for Healthcare Professionals"**

16:40-16:45 Q&A Session (5 mins)

16:45-17:10 **"Legal and Ethical Implications in Medical Device Regulation and Safety"**

17:10-17:15 Q&A Session (5 mins)

17:15-17:20 Closing Remarks

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Speakers

Richard Houlihan

Richard Houlihan is an international speaker and guest university lecturer on EUDAMED. His time running the European Commission IT teams developing EUDAMED has given him unrivalled insights into how EUDAMED affects the MedTech industry. This EUDAMED experience and his 28 years in IT give him a real advantage when helping companies prepare for EUDAMED, FDA GUDID, and other regulatory submissions.

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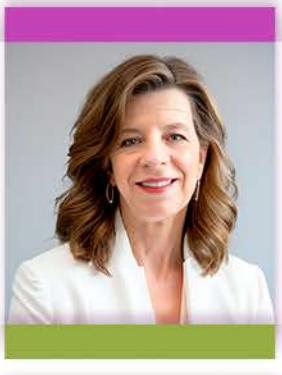
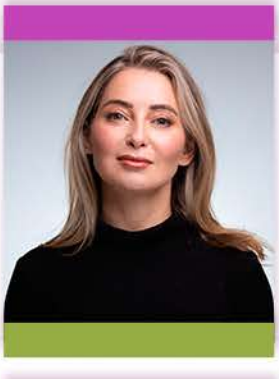


Ann Vu

Ann leads quality and regulatory compliance across the ZimVie global organization. She brings more than 25 years of transformative quality, operations and regulatory experience in the medical device, pharmaceutical, nutritional, and biologics sectors. Ann joined ZimVie from Zimmer Biomet, where she led Quality and Regulatory Affairs for Robotics and several other business units. She has also held progressive leadership roles with global companies, including Ethicon, Ortho Clinical Diagnostics, Bausch + Lomb, and Steris.

ZimVie



Jacqueline van Druten

Jacqueline van Druten is the EMEA & NA Clinical & Regulatory Affairs Director at CLIN-r+, she leads a team of consultants and advisors who provide data-driven solutions for medical device regularory submissions. This includes but is not limited to compiling Dossiers/Technical Documents, Bioconpatability Evaluations, clinical development strategies (CDP) and post-market surveillance automation solustions for MedTech manufacturers. With over two decades of experience in the healthcare industry, she has a strong background in clinical research, medical innovation (pharma, MedTech, biotech and nutraceuticals) and marketing, as well as a particular interest in MedTech validations, MedTech sustainability

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