

MedTech Summit

11-15 June 2018
Hotel NH Brussels Bloom
Brussels

EU MDR & IVDR: THE RACE IS ON...

Including Clinical Strategies,
PMS & Vigilance, Legal,
Combination Products,
Sterilisation, and
Emerging Markets Requirements



DAY ONE MONDAY 11 JUNE	DAY TWO TUESDAY 12 JUNE	DAY THREE WENESDAY 13 JUNE	DAY FOUR THURSDAY 14 JUNE	DAY FIVE FRIDAY 15 JUNE
EU Medical Device Regulation	EU Medical Device Regulation	Medical Device Regulatory Affairs in Emerging Markets Asia	Medical Device Regulatory Affairs in Emerging Markets RUSSIA & CEE	Medical Device Regulatory Affairs in Emerging Markets LATAM, Middle East & Africa
	Clinical Evaluations & Investigations	Clinical Evaluations & Investigations	Drug Device Combination Products	Drug Device Combination Products
	Post Market Surveillance & Vigilance	Post Market Surveillance & Vigilance	IVD Regulatory & Strategic Forum	IVD Regulatory & Strategic Forum & CDx
EU Medical Device Law	EU Medical Device Law		Sterilisation and Reprocessing of Medical Devices	Sterilisation and Reprocessing of Medical Devices

MAIN CONFERENCE DAY ONE • MONDAY 11 JUNE 2018

07:50	<i>Registration</i>
08:50	Chairperson's opening remarks
09:00	Scientific and technical challenges of the novel EU medical devices legislative framework Arnd Hoeveler, Head of Unit of JRC (Joint Research Centre), European Commission, Italy
09:20	SME perspective: Where are we with EU MDR implementation? Susana de Azevedo Wäsch, Head of Global Regulatory Affairs, Geistlich Pharma AG, Switzerland
09:40	Competent Authority perspective: Practical advice and expectations for industry in preparing for EU MDR implementation Graeme Tunbridge, Group Manager – Devices Regulatory Affairs, MHRA, UK
10:00	Perspective on Notified Bodies: Preparing for EU MDR implementation and expectations for industry Gert Bos, Executive Director & Partner, Qserve Group, The Netherlands
10:20	IMPLEMENTATION PANEL DISCUSSION: Sharing experiences with interpreting and implementing the EU MDR in practice Gert Bos, Qserve Group Elizabeth Gfoeller, MED-EL Hans-Heiner Junker, TÜV SÜD Product Service Juliette Cook, Cochlear AG Alexander Natz, EUCOPE Rainer Voelksen, Occlutech Lincoln Tsang, Arnold & Porter
11:00	<i>Morning coffee break</i>

Speaking, panellist, moderator and webinar opportunities

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MAIN CONFERENCE DAY ONE • MONDAY 11 JUNE 2018 *(continued)*

	EU Medical Device Regulation	EU Medical Device Law
11:40	Notified Body perspective: Preparing for re-designation and latest status of the application process Julien Sénac, Certification Project Manager, LNE/G-MED, USA	Practical implications of the new regulations and case law on commercial relationships Jackie Mulryne, Counsel, Arnold & Porter, UK Ewan Townsend, Counsel, Arnold & Porter, UK
12:15	PANEL DISCUSSION: Sharing latest experiences and concerns on the Notified Body re-designation process Guy Buijzen, DEKRA Bassil Akra, TÜV SÜD Product Service GmbH Julien Sénac, LNE/G-MED	Understanding new liability rules under the MDR Sean Fahey, Partner, Pepper Hamilton LLP, USA
12:50	Networking Lunch	
14:00	Outlining significant changes in the responsibilities and obligations of economic operators under the EU MDR Juliette Cook, Regional Regulatory Director EMEA, Cochlear AG, Switzerland	Person Responsible for Regulatory Compliance Bjorn Delbeecke, Regulatory Affairs Specialist EMEA, Agfa HealthCare, Belgium
14:35	Latest updates on the MDSAP program including Canada's 1 January 2019 deadline and impacts on industry Maham Ansari, Director of Regulatory Affairs, Synaptive Medical, Canada	Presentation to be delivered by White & Case LLP Axel Schulz, Partner, White & Case LLP, Belgium
15:10	Afternoon tea break	
15:50	Understanding the role of CEN and CENELEC (and their European standards) in the context of the new EU MDR and their priorities for ensuring a smooth transition Catherine Vigneron, Customer Service Specialist, Standardization and Digital Solutions, CEN-CENELEC Management Centre, Belgium	Exploring key legal and contractual considerations for economic operators under the EU MDR Erik Vollebregt, Partner, Axon Lawyers, The Netherlands
16:25	End of conference day one	Notified Body liability and legal status under the EU MDR Heike Wachenhausen, Partner, Wachenhausen Rechtsanwälte Partnerschaft mbB, Germany
		17:00 End of conference day one

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MAIN CONFERENCE DAY TWO • TUESDAY 12 JUNE 2018

09:00	Chairperson's Opening Remarks			
	EU Medical Device Regulation	EU Medical Device Law	Clinical Evaluations & Investigations	Post Market Surveillance and Vigilance
09:10	EUDAMED: Latest timelines and updates Céline Bourguignon, Director, Government Affairs EMEA Medical Device & IVDs Department, <i>Johnson & Johnson, Belgium</i>	Exploring recent EU case law related to the EU MDR Emmanuel Garnier, Managing Associate, <i>Simmons & Simmons, France</i>	CASE STUDY: Exploring the key impacts of the EU MDR and EUDAMED developments on clinical evaluations and investigations Carine Cochereau, QRA and Clinical Director EMEA, <i>Cardinal Health, Belgium</i>	Exploring the changes in the new MDR relating to post market surveillance (PMS) and vigilance Hans-Heiner Junker, Senior International Affairs Manager, <i>TÜV SÜD Product Service, Germany</i>
09:45	CASE STUDY: Practically getting into compliance with UDI Unique Device Identification (UDI): Latest status, requirements and practically preparing in Europe Bianca M. Gravenhorst Greve, Senior RA Manager, <i>Coloplast A/S, Denmark</i> Inge Ørnhoj, Senior Master Data Manager, <i>Coloplast A/S, Denmark</i>	Exploring the legal implications of the General Data Protection Regulation (GDPR) and impact on manufacturers Wolfgang Werner, VP Regulatory and Quality, <i>Biovotion AG, Switzerland</i>	Competent Authority perspective: Expectations from industry on new requirements for clinical evaluations and investigations under the MDR Tom Melvin, Health Products Regulatory Authority, <i>Ireland</i>	CASE STUDY: Developing an ISO guidance document for PMS Arjan van Drongelen, Scientific Officer, <i>National Institute for Public Health and the Environment (RIVM), The Netherlands</i>
10:20	Morning coffee break			
11:00	Clarifying the relationship between ISO 13485:2016 – Quality Management Systems (QMS) and the EU MDR Robert van Bortel, Principal Consultant, <i>Medical Device Project B.V., The Netherlands</i>	PANEL DISCUSSION: Sharing experiences with preparing for requirements of the GDPR Wolfgang Werner, <i>Biovotion AG</i> Olivier Mignolet, <i>Simmons & Simmons</i>	Competent Authority perspective: Expectations for clinical investigations and GCP under the MDR Tim De Schutter, Inspector Medical Devices, <i>FAMHP, Belgium</i>	EXTENDED PANEL DISCUSSION: How are different companies practically implementing new PMS and vigilance requirements under the new MDR? Fayez Abou Hamad, <i>Terumo Europe N.V.</i> Nathalie Gerard, <i>Terumo Europe N.V.</i> Andrea Sparti, <i>Cendres+Métaux SA</i>
11:35	Outlining significant changes for manufacturers falling under the self-certification process (Class I and custom made devices) Katrien Martens, Inspector, <i>FAMHP, Belgium</i>	PANEL DISCUSSION: The global courtroom: Strategies for managing cross-border products liability litigation Lori G. Cohen, Shareholder; Chair, Pharmaceutical, Medical Device & Health Care Litigation Group; Chair, Trial Practice Group, <i>Greenberg Traurig LLP, USA</i> Maria H. Bragaglia, Partner, <i>Demarest Advogados, Brazil</i> Ashley L. Paterson, Associate, <i>Bennett Jones LLP, Canada</i> Aimee H. Wagstaff, Partner, <i>Andrus Wagstaff, USA</i>	Presentation to be delivered by Indegene Representative speaker from <i>Indegene</i>	
12:10	Networking Lunch			

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MAIN CONFERENCE DAY TWO • TUESDAY 12 JUNE 2018 *(continued)*

	EU Medical Device Regulation	EU Medical Device Law	Clinical Evaluations & Investigations	Post Market Surveillance and Vigilance
13:30	Exploring the regulatory qualification, classification and future of medical device software Koen Cobbaert, AGFA Healthcare Mette Kraag Luxhøj, Radiometer	Understanding the classification of borderline products and associated legal issues John P. Lavelle, Jr., Partner, Morgan, Lewis & Bockius LLP, USA	Notified Body perspective: Practical advice for industry on meeting new requirements for clinical evaluations and investigations under the MDR Bassil Akra, Vice President – Global Focus Teams (Cardiovascular, Orthopaedic and Clinical), TÜV SÜD Product Service GmbH, Germany	Exploring the impact the obligations of economic operators will have on manufacturers' implementation of the PMS process Andrea Sparti, Regulatory Affairs Manager, Cendres+Métaux SA, Switzerland
14:05		Reviewing the current status of advertising and claims for medical devices Bill Kurani, MSRA, MSEE, Head of RA/QA, Genomics, Agilent Technologies, USA	Clinical Evaluations: Moving from MDD to MDR Vincent Legay, Senior Project Manager, NAMSA, France	CASE STUDY: Compiling post market surveillance (PMS) plans and meeting requirements of the new MDR Andrea Castañeda, Associate Manager - International Quality Compliance, Edwards Lifesciences, Spain
14:40	Afternoon tea break			
15:20	OPEN FLOOR Q&A INTERACTIVE SESSION Gert Bos, Qserve Group Hans-Heiner Junker, TÜV SÜD Product Service Elizabeth Gfoeller, MED-EL		Post Market Clinical Follow-Up (PMCF): Exploring Competent Authority and Notified Body expectations Bassil Akra, Vice President – Global Focus Teams (Cardiovascular, Orthopaedic and Clinical), TÜV SÜD Product Service GmbH, Germany	
15:55			CASE STUDY: Practically implementing Post Market Clinical Follow-Up (PMCF) activities under the new MDR Jean-François Lataste, Sr. Clinical Research Manager, Medtronic, Switzerland	
16:30	End of conference day two			

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MAIN CONFERENCE DAY THREE • WEDNESDAY 13 JUNE 2018

09:00	Chairperson's Opening Remarks		
	Clinical Evaluations & Investigations	Post Market Surveillance and Vigilance	Regulatory Affairs in Emerging Markets
09:10	Notified Body perspective: MEDDEV 2.7.1 Revision 4 expectations and observations from industry on clinical evaluations Ito Udofia , Head of Notified Body, UL International (UK) Ltd., UK	CASE STUDY: Ensuring best practice for Field Safety Corrective Action (FSCA) reporting Maite Llácer , Director Quality Compliance INT'L, Edwards Lifesciences, Spain	Clarification and exploration of the expectations of the CFDA for medical device registration Jack Wong , Director, Asia Regulatory Professional Association, Singapore
09:45	CASE STUDY: Practically applying the new MEDDEV 2.7.1 rev 4 guidance for clinical evaluations Avanti Kulkarni , Global Regulatory Affairs – Principal Clinical Evaluation Manager, Abbott Vascular, USA Vinuta Rau , Principal Clinical Research Scientist, Clinical Affairs, Abbott Vascular, USA	CASE STUDY: Successfully implementing a robust complaint handling and surveillance system Anne Mørkeberg Larsen , Senior Product Surveillance Advisor, Novo Nordisk A/S, Denmark Lavinia Marinescu Pedersen , Senior Product Defect and Recall Advisor, Novo Nordisk A/S, Denmark	Understanding China's CFDA clinical evaluation requirements Ed Woo , Asia Pacific Regulatory Affairs Director, Varian Medical, Hong Kong
10:20	INTERACTIVE PANEL DISCUSSION: Sharing experiences with writing a CER following MEDDEV 2.7.1 rev 4 Ilona Anderson , Director of Clinical Research, MED-EL GmbH, Austria Representative speaker from Cactus Communications	How to integrate Complaint Trending with Risk Management Ruben Roijers , Post Market Surveillance, Philips, The Netherlands	CASE STUDY: Navigating China's regulatory processes for successful product registration Wen Peng , Director, Regulatory Affairs, Asia Pacific, Edwards Lifesciences, China
10:55	Morning coffee break		
11:30	CASE STUDY: Practically preparing for stricter clinical evidence requirements for legacy products under the new MDR Leo Hovestadt , International Regulatory Affairs and Quality Assurance Director, Elekta, The Netherlands	Outlining new timelines for post market reporting and practically meeting requirements Tabitha Reed , Senior Manager, Regulatory Post Market Surveillance, Intuitive Surgical, Inc., USA	INTERACTIVE ROUNDTABLE DISCUSSION: Sharing experiences with successfully registering products in China Jack Wong , Director, Asia Regulatory Professional Association, Singapore
12:05	State of the art best practices and literature review using distiller Laurie Mitchell , President, Criterion Edge Inc., USA Peter O'Blenis , President, Evidence Partners, Canada	CASE STUDY: Best strategies for leveraging post market surveillance (PMS) data to become better informed about your products Patrick Caines , Senior Director, Quality & Compliance, Baxter, USA	Software as a Medical Device (SaMD) and Cybersecurity in China Tobias Schreiegg , Quality & Regulatory Affairs Manager, Siemens Healthcare, Germany
12:40	Networking Lunch		
12:50	Networking Lunch		

MAIN CONFERENCE DAY THREE • WEDNESDAY 13 JUNE 2018 *(continued)*

	Clinical Evaluations & Investigations	Post Market Surveillance and Vigilance	Regulatory Affairs in Emerging Markets
14:00	CASE STUDY: Practically applying risk based monitoring for medical device clinical investigations Nurcan Coskun, Global Risk Based Monitoring & Technology Solutions Program Manager, Medtronic , <i>Switzerland</i>	INTERACTIVE SESSION: How can I be compliant with the MDR Post Market activities and reports? Philippe Auclair, Abbott Laure-Anne Thieren, Johnson & Johnson Medical Devices	The implementation progress of the ASEAN Medical Device Directive (AMDD) Jack Wong, Director, Asia Regulatory Professional Association, <i>Singapore</i>
14:35	CASE STUDY: Implementing a successful clinical investigation / post market clinical follow up (PMCF) strategy Dorota Johansson, Clinical and Research Director, Bactiguard , <i>Sweden</i>		Practical guidance on ASEAN labelling requirements Dr. Jennifer Neff, Director Regulatory & Medical Affairs, bess AG , <i>Germany</i>
15:10	Afternoon tea break		
15.50	CASE STUDY:Successfully outsourcing clinical investigations Frank Van Leeuwen, Focus Clinical Consultancy, <i>Belgium</i>	Generating the necessary evidence through Life Cycle Communication strategy Patrice Becker, Global Director Scientific Communications, Medical Affairs, Medtronic , <i>France</i>	INTERACTIVE ROUNDTABLE SESSION: Sharing experiences with medical device registration in ASEAN countries Dr. Jennifer Neff, Director Regulatory & Medical Affairs, bess AG , <i>Germany</i>
16.25	Reviewing requirements of the new MDR for demonstrating equivalence for clinical evaluations Regina Moeckel, Medical Devices, TÜV Rheinland LGA Products GmbH, <i>Germany</i>	End of the Day Q&A: A chance for any remaining questions	Exploring an overview of the regulatory environment in Hong Kong Ed Woo, Asia Pacific Regulatory Affairs Director, Varian Medical , <i>Hong Kong</i>
17:00	Outlining requirements for clinical evidence for class III devices in the MDR Eddie Van Eeckhoven, Medical Device Regulatory and Clinical Advisor, Eeckhoven bvba , <i>Belgium</i>	End of conference day three	The impact of the EU Medical Device Regulation (MDR) on emerging markets Elena Mateo-Buesa, Regulatory Affairs Manager – Global Strategy, Ethicon (part of J&J family of companies) , <i>Germany</i>
17:35	End of conference day three		

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MAIN CONFERENCE DAY FOUR • THURSDAY 14 JUNE 2018

09:00	Chairperson's Opening Remarks			
	Sterilisation and Reprocessing of Medical Devices	Drug Device Combination Products	IVD Regulatory & Strategic Forum	Regulatory Affairs in Emerging Markets
09:10	Understanding the impact of the EU Medical Device Regulation (MDR) on sterilisation processes Jeff Vest, Principal Specialist – MHS (Technical Reviewer), TÜV SÜD Product Service, UK	Notified Body Perspective: Practically implementing new requirements of the EU MDR for drug device combination products (DDCPs) Daniel Shoukier, Lead Auditor, SQS, Switzerland	Practically implementing the IVDR: Competent Authority perspective Stephen Lee, Senior Regulatory Policy Advisor – IVDR, MHRA, UK	Shedding light on Russian requirements for manufacturing site inspections Maria Bogorad, Regulatory, Quality & Continuous Improvement, Johnson & Johnson, Russia
09:45	Latest updates on national implementation of reprocessing single-use devices and implications of Common Specifications for industry Dan Vukelich, President, Association of Medical Device Reprocessors (AMDR), USA	Industry Perspective: Practically implementing new requirements of the EU MDR for drug device combination products Hilde Viroux, Global Head EU MDR Compliance, Alcon, USA	Practically implementing the IVDR: Notified Body perspective Nick Baker, IVD Technical Manager, LRQA, UK	Understanding the Russian Federation government procurement system and the implications for industry Yana Pruss-Galynsky, CEO, Zephyr Biomedical, Israel
10:20	Exploring new and existing requirements for reprocessing of devices for sterile use Hagen Thielecke, Director R&D, Vanguard AG, Germany	EU MDR Rule 21: Evaluating the impact on substance-based devices Jörg Plessl, Director EU Affiliates, Regulatory Affairs, Norgine, Germany	INTERACTIVE PANEL DISCUSSION: Working with Notified Bodies and Competent Authorities to meet IVDR requirements Stephen Lee, Senior Regulatory Policy Advisor – IVDR, MHRA, UK Nick Baker, IVD Technical Manager, LRQA, UK Sue Spencer, Head of Global Medical Device Services, UL, UK Catherine Holzmann, Certification Division Manager In Vitro Diagnostics, LNE/G-MED, France Simon Richards, VP, Regulatory Affairs, EMER, Alere, UK	Safety monitoring: Best practises for successfully conducting post-market surveillance in Russia
10:55	Morning coffee break			
11:25	Exploring new EU MDR requirements for classification of class I reusable medical devices Jan Havel, Team Leader Nonactive Medical Devices Generic Group, TÜV SÜD Product Service GmbH, Germany	Understanding the requirements for biocompatibility as outlined in ISO 10993 for drug device combination products Clemens Günther, Director Nonclinical Safety, Bayer, Germany	Practically implementing the IVDR: Industry perspective Simon Richards, VP, Regulatory Affairs, EMER, Alere, UK	EXTENDED INTERACTIVE SESSION: Practical guidance on successfully registering medical devices in Russia A case study from industry followed by a collaborative discussion Chris Dark, Head of Regulatory Affairs, Arkray Factory Ltd, United Kingdom Marina Semenova, QA Manager, Arkray ooo, Russia
12:00	The MDR requires evidence regarding the qualification and competency of personnel in sterilization – how is the Medical Device industry going to achieve this? Arthur Dumba, Medical Device Sterilization Consultant, International Irradiation Association (iia), Switzerland	Drug-device compatibility: Where is the guideline? Lise Vanderkelen, Department Head Pharma Services, Toxikon Europe, Belgium	Understanding the new classification system for IVDs under the IVDR Mark Thomson, Senior Regulatory Expert, Medical Device and Equipment Team, Life Science Regulatory	
12:35	Networking Lunch			



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MAIN CONFERENCE DAY FOUR • THURSDAY 14 JUNE 2018 *(continued)*

	Sterilisation and Reprocessing of Medical Devices	Drug Device Combination Products	IVD Regulatory & Strategic Forum	Regulatory Affairs in Emerging Markets
14:00	CASE STUDY: Best practice for successfully carrying out Ethylene Oxide (EO) sterilisation Jan Douglas , Engineering Manager, Cook Medical, Denmark	CHMP Meetings: How to prep for, manage and excel at these EU regulatory meetings Bert Regeer , MD, Senior Scientific Lead, 3D Communications, USA	New requirements of clinical evidence under the IVDR Sue Spencer , Head of Global Medical Device Services, UL, UK	Updates on the progress of the Eurasian Economic Union (EAEU) regulations and the impact on industry Alexey Stepanov
14:35	Exploring Ethylene Oxide (EO) residual testing per ISO 10993-7 Pamela Barrows , Director, Controlled Environments and Sterilization Control, Getinge Group, USA	Updates on Canadian regulation of drug device combination products Douglas Watson , Senior Policy Analyst, Health Canada, Canada Subject to Final Confirmation	Implications of the IVDR clinical evidence requirements on legacy products Maurizio Suppo , Vice President Regulatory Affairs, QARAD, Italy	Understanding labelling requirements for medical devices in Russia Bulat Magdiev , Regulatory Associate, RB, Russia
15:10	Afternoon tea break			
15:40	CASE STUDY: Exploring sterilization validation approaches for Gamma and ETO sterilization Bjoern Wiese , Director, Sterilization Technology and Analytical Testing EMEA, Zimmer Biomet, Switzerland	Updates from the FDA on new regulations regarding drug device combination products Barr Wiener , Associate Director Combination Products Office, FDA, USA	INTERACTIVE ROUNDTABLE SESSION: Sharing experience with increased clinical evidence requirements under the IVDR Chris Weatherall , Senior Director Quality Assurance & Regulatory Affairs, Meridian Bioscience Inc., UK	Practically exploring updates to medical device regulations in Kazakhstan Yana Pruss-Galynsky , CEO, Zephyr Biomedical, Israel
16:15	Exploring latest updates to the Radiation Sterilization standard ISO 11137 and implications for industry Arne Miller , Professor, Risø High Dose Reference Laboratory, Denmark	Practically approaching the US FDA GMP guidelines for combination products Stephan Affolter , Head of Quality System & Regulatory Affairs, Ypsomed, Switzerland	Overview and updates to EUDAMED and implications to IVDs Céline Bourguignon , Director, Global regulatory policy, Government Affairs & Policy, Johnson & Johnson, Chair of the MedTech Eudamed Taskforce, MedTech Europe, Belgium	The adoption of an EU Medical Device Directive (MDD)-based regulation system in Ukraine Alexander Schapovalov , Foreign Affairs Manager, TÜV SÜD, Germany
16:50	End of conference day four	Comparing EU MDR and US FDA drug device combination product regulations to identify effective regulatory strategies Winifred Wu , President and Principal Advisor, Strategic Regulatory Partners, USA	Unique Device Identifiers (UDI) requirements for IVDs Melissa Finocchio , Project Portfolio Leader, Global Quality, bioMérieux, France	The Product Tracking System (UTS): Turkey's new registration database and other upcoming changes Ilker Yilmaz , Regulatory Affairs & Market Access Manager, Meril, Turkey
	17.25 End of conference day four			

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MAIN CONFERENCE DAY FIVE • FRIDAY 15 JUNE 2018

09:00	Chairperson's Opening Remarks			
	Sterilisation and Reprocessing of Medical Devices	Drug Device Combination Products	IVD Regulatory & Strategic Forum	Regulatory Affairs in Emerging Markets
09:10	Moving from the MDD to the MDR – what will be the impact on sterile packaging, including latest status updates to ISO 11607 – Packaging for terminally sterilized medical devices Thierry Wagner , Regulatory Affairs Director - Europe, Middle-East & Africa, DuPont Medical and Pharmaceutical Protection , <i>Luxembourg</i>	Practically overcoming regulatory and practical challenges with the design, manufacture, and distribution of pre-filled syringes Ann Jans , Quality Assessor, FAMHP , <i>Belgium</i>	Clarification of the role of the Person Responsible for Regulatory Compliance (PRRC) under the IVDR from a practical standpoint Petra Kaars-Wiele , Senior Director International Regulatory Affairs & Global Labeling, Abbott , <i>Germany</i>	ANVISA perspective: Understanding the latest regulatory framework in Brazil and the implications for industry
09:45	CASE STUDY: Handling sterile packaging waste Karina Engels , Product Manager, Sterilisation & InteguSeal EMEA, Halyard Health , <i>Belgium</i>	Practically approaching the co-development of drug and device components of drug device combination products Sanjay Jain , Drug Delivery Systems Principal Engineer, Janssen , <i>USA</i>	The effects the IVDR will have on economic operators Chris Dark , Head of Regulatory Affairs, Arkray Factory Ltd , <i>UK</i>	CASE STUDY: How to pass the ANVISA GMP audit Gloria Schneider-Ferrer , Manager, Global Strategy - Brazil, DePuy Synthes , <i>Switzerland</i>
10:20	Global sterility assurance challenges and opportunities Byron Lambert , Sr Assoc Fellow, Sterilization, Abbott Vascular , <i>USA</i>	Considerations for how increased clinical data requirements as proposed in the EU MDR impacts combination products Florian Tolkmitt , Clinical Evaluation & Regulatory Affairs Expert, <i>Germany</i>	New requirements of post-market surveillance under the IVDR Gary Carpenter , Director of Clinical and Regulatory Affairs, BioSure , <i>UK</i>	MDSAP pilot update: The impact of the MDSAP on the Brazilian regulatory environment Georg Bauer , Department Manager Foreign Affairs, TÜV SÜD , <i>Germany</i>
10:55	Morning coffee break			
11:25	Assessing global expectations for EO-sterilisation Julio Morote Garcia , RA & Sterilization Specialist, Bentley InnoMed GmbH , <i>Germany</i>	An overview of recent changes and updates in human factors engineering guidance and the implications for drug device combination products Molly Story , Senior Director, Global Usability Engineering and Risk Management, Sanofi , <i>USA</i>	Incident reporting for IVDs under the IVDR Caroline Alexander , Head of Quality and Regulatory Affairs, QuantuMDx Group Ltd. , <i>UK</i>	Review of updates to INMETRO certification requirements: How to ensure compliance Luciano Oliveira Ferreira , Head of Quality & Regulatory Affairs, QA/RA Works , <i>Brazil</i>
12:00	Understanding the liquid sterilisation of medical devices with derivatives of animal origin Wilhelm Erdbrügger , Sen. Manager Process Development, BIOTRONIK , <i>Switzerland</i>	INTERACTIVE SESSION: A hands-on exploration of human factors testing for combination products Yvonne Limpens , Managing Human Factors Specialist, UL , <i>Netherlands</i> Frauke Schuurkamp , Managing Human Factors Specialist, UL , <i>Netherlands</i>	Comparing US and EU regulatory requirements for IVDs	Updates to COFEPRIS regulations in Mexico and guidance for implementation Ana Luisa Fritschy , Manager Regulatory Affairs, Aptar Stelmi , <i>France</i>
12:35	Networking Lunch			

MAIN CONFERENCE DAY FIVE • FRIDAY 15 JUNE 2018 *(continued)*

	Sterilisation and Reprocessing of Medical Devices	Drug Device Combination Products	IVD Regulatory & Strategic Forum	Regulatory Affairs in Emerging Markets
14:00	Microbiological aspects <i>Ciming Li, Product Safety Expert, Switzerland</i>	Understanding the challenges associated with packaging and labelling for drug device combination products <i>Judit Jávör, Regulatory Affairs Expert & Compliance Director, PharmaPatent, Hungary</i>	The impacts of the IVDR for companion diagnostics manufacturers <i>Alex Laan, Principal Certification Manager, Dekra Certification, Netherlands</i>	New requirements for post-market surveillance and vigilance in Latin America <i>Andrea Sisneros, Director, Riodove Services Ltd, UK</i>
14:35	Exploring sterilisation techniques for 3D printed products <i>James McCoy, Staff Quality Scientist, Global Sterilization Research and Innovation, BD, USA</i>	Identifying latest regulatory requirements and practical approaches to designing generic or biosimilar/device combination products <i>Alice Maden, Associate Director, Regulatory Affairs, BD, France</i>	How the IVDR impacts both pharmaceutical and diagnostics manufacturers <i>Claudia Dollins, Head of Global Regulatory Affairs, Biomarkers & Diagnostics, Merck, Germany</i>	PANEL DISCUSSION: Exploring the latest regulatory requirements across Latin America <i>Ana Luisa Fritschy, Manager, Regulatory Affairs, Aptar Stelmi, France</i> <i>Patricia Pascale, Regulatory Affairs Director, LATAM, ConvaTec, Brazil</i>
15:10	Afternoon tea break			
15:40	CASE STUDY: Best practice sterilisation processes for drug device combination products <i>Jeff Vest, Principal Specialist – MHS (Technical Reviewer), TÜV SÜD Product Service, UK</i>		INTERACTIVE ROUNDTABLE SESSION: How is the companion diagnostics industry preparing for changes with the IVDR? <i>Tricia Carrigan, Head, Companion Diagnostics, Regulatory Affairs Oncology, Bayer, Germany</i>	A practical overview of Saudi Arabia's pre-market process <i>Naif Alhussinan, Acting Section Head for Licensing Importers and Distributors, SFDA, Saudi Arabia</i>
16:15	INTERACTIVE Q&A SESSION: Sterilisation of drug device combination products <i>Tim Carlson, Sterilization Program Manager, Global Sterilization Assurance, BD, USA</i>		Discussion of the implications of Brexit on the IVD industry <i>Fabien Roy, Counsel, Hogan Lovells International LLP, Belgium</i>	PANEL DISCUSSION: Navigating requirements and strategies for registering medical devices in the Middle East and North Africa (MENA) <i>Naif Alhussinan, Acting Section Head for Licensing Importers and Distributors, SFDA, Saudi Arabia</i> <i>Reem El-Sayed, Regulatory Affairs Director Turkey, Middle East & Saudi Arabia, GE Healthcare, UAE</i> <i>Mamoona Firdous, Regulatory Affairs Manager, Julphar, UAE</i>
16:50	End of conference day five			

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