MEDTECH & PHARMA
PLATFORM

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Digital Training Course

Overview of Medical Devices Regulation and Impacts on Combination Products

29 September 2020 | 2:00 pm - 5:00 pm (CET)

Objectives

This training aims at providing a comprehensive overview of the regulatory system and requirements for medical devices in Europe with particular focus on combination products.

With the MDR applying from May 26th 2021 this training shall also provide an update on the most recent developments and implementation timelines to support you in being ready for May next year.

Course leaders

- Karin Schulze, Head of Medical Devices at SFL
- Stephan Affolter, Regulatory and Quality Intelligence Manager at Ypsomed
- Juan Martin Carriquiry, EU MDR Expert, Device Development & Commercialization at Novartis

Registration fee

Standard rate: 300 CHF*
*Fees subject to VAT.

Agenda

- Introduction to the MDR
- Highlight of the key changes in the MDR 2017/745
- Update on the latest changes
- MDR Implementation for Medical Devices – Industry Perspective
- MDR Implementation for Article 117 impacted combination products – Industry Perspective

About Medtech & Pharma Platform (MPP) Association

The Medtech & Pharma Platform (MPP) is a cross-sectoral not-for-profit industry association focusing on combined products. MPP is made up of Medtech, Pharma and Software companies dedicated to enhancing synergies between the sectors and to provide a cross-sectoral forum to exchange knowledge, collaborate in technology and regulatory areas as well as to promote product development and innovation.

The association aims to further strengthen advocacy work for companies to reduce time to market for drugs, devices and combinations thereof, improve access to innovative products and better match patients' needs.

Register now

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