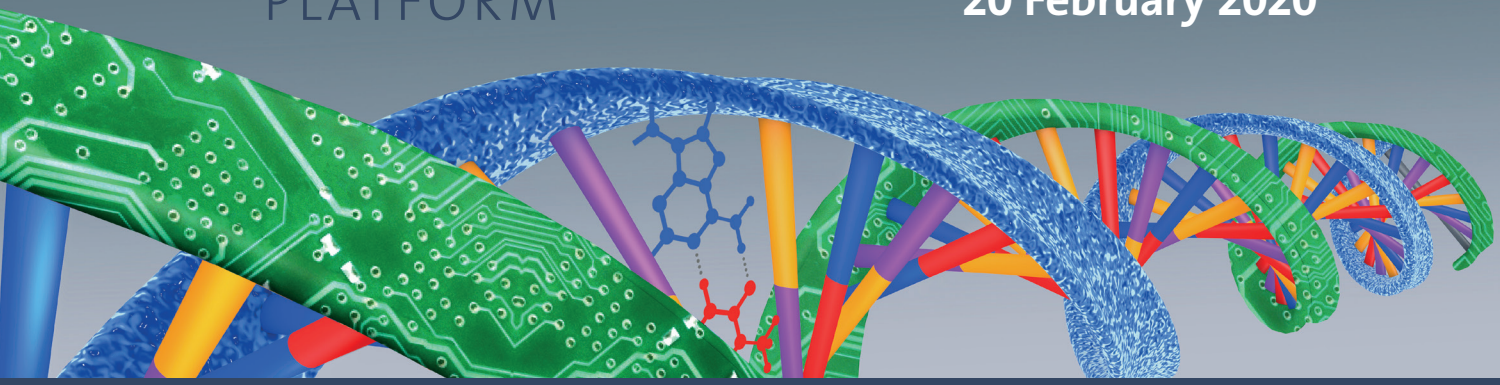
MEDTECH & PHARMA PLATFORM

1-Day Training Course
20 February 2020



Overview of Medical Devices Regulation and Impacts for Medtech and Pharmaceutical Industry

Special Focus: MDR implementation for medical devices and combination products

20 February 2020
9:00 am - 5:00 pm
Basel, Switzerland

Information and registration:
www.medtech-pharma.com

Objectives

This training aims at providing a practical, hands-on and 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 2020 this training shall also provide an update on the most recent developments and implementation timelines to support you in being ready for May this year.

Workshops and practical examples will complement the training.

Course leaders

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

Registration fee

Standard rate: 500 CHF*
Member rate: 400 CHF*
*Fees subject to VAT.

Agenda

Introduction to the MDR

- Highlight of the key changes for the MDR 2017/745
- Latest news from the European Commission
- Timelines, transitional provisions and Notified Body (NB) designation update
- MDR implementation project and gap analysis

Special focus session: MDR Implementation from an industry perspective (pharma and medical device)

- Combination products
- Medical devices

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

Benefit from recognized leaders in an interactive, accessible and timely format
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