COVARTIM®

STEP-UP Program

Register by
January 20, 2021

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MDR QMS Update Workshop

Get your QMS compliant to MDR

Objectives

Understand the MDR impact on your ISO 13485:2016 QMS

Upgrade your QMS to be compliant with the MDR

Duration: 2,5 days

Location: to be defined

Prerequisite: Your are a manufacturer of a medical device

Your QMS must be already certified ISO 13485:2016

Participants: QA/RA Manager, QA/RA Associate, Management Representative

Registration fee: 5.000 € VAT excl. / Company

DAY 1 (week #5, 2021)	DAY 2 (Week #8, 2021)	DAY 3 (TBD)
Review of the MDR impact on QMS Checklist for compliance Case studies Templates Breakout rooms: ↳ Work on your own QMS ↳ Get support from COVARTIM's experts	Experience sharing Breakout rooms: ↳ Finalize your own QMS ↳ Get support from COVARTIM's experts	On site (or remote) mock audit: check-up of your QMS compliance to the MDR requirements

IMPORTANT NOTES:

Two representatives per company can participate. This workshop includes **2 collective sessions** (Day 1, Day 2) and **1 individual session** (Day 3). During Day 1 and Day 2, COVARTIM's experts will provide individual coaching throughout the program. Participants must get access to their own QMS. **Templates** will be provided. Participants will receive a **certificate** after the full completion of the workshop.

Number of seats limited. Payment must be made upon registration.