Medical Device Regulation (EU MDR) 2017/745

If you already sell medical devices into Europe, you probably already know that CE marking is changing. But do you fully understand what the impact is for your business, products and your supply chain?

If you are not 100% sure or would like to refresh your knowledge, this 2 day course is most definitely for you.

Created by industry leading experts, this course has been designed to help medical device manufacturers understand the additional requirements of the standard, so you know what you need to do for your business and products to meet the new MDR.

Our expert trainers will take you through the MDR, not just from an industry perspective, but also from a Notified Body perspective, so you understand what both sides are looking for.

"Fantastic to have an instructor with subject knowledge that was able to relate to the experience of participants. Presented and delivered at a good pace to allow understanding and ample opportunity for specific questions."

Upon successful completion of the course you will receive a Meddev Solutions course certificate.

Find out more:
info@meddevsolutions.co.uk
www.meddevsolutions.co.uk/training

Course Content

- Introduction to the EU MDR
- Medical Devices covered by the EU MDR
- Placing a device on the market
- Device Classification
- Unique Device Identifiers (UDI)
- Routes to Conformity
- Clinical Data Evaluation
- Technical File documentation
- GSPR & Risk Management
- Post Market Surveillance & Vigilance

Who should attend?

This EU MDR training course has great value for anyone involved in implementing or maintaining a QMS.

- Quality Assurance professionals
- Quality Engineers
- Research and Design Engineers
- Internal Auditors
- Quality Managers
- Manufacturing Engineers
- Regulatory professionals

Available as
In-House • Public • Virtual • eLearning

Bespoke
Our experts can customise this course purely for your business needs.