

# Training Program on European Union Medical Device Regulations

Become EU MDR-ready and elevate your role as  
a Global Medical Device Professional!

Advance Your Regulatory Skills - Empower Global Compliance!

## Training Modules:

- ◆ General Overview of EUMDR
- ◆ Classification of Medical Devices
- ◆ Economic Operators
- ◆ Quality Management System (QMS) Requirements
- ◆ Technical Documentation
- ◆ Clinical Evaluation & Investigation
- ◆ Conformity Assessment
- ◆ UDI and EUDAMED
- ◆ Labelling and Instructions for Use (IFU)
- ◆ Post-Market Surveillance (PMS)

**Eligibility:** Medical Device  
Manufacturers, Quality and Regulatory  
Professionals, Service Providers in  
Healthcare sector & Researchers  
Academics.

**Mode:** Online  
**Duration:** 2 Days



For registration:  
[www.ibsc-amtz.in](http://www.ibsc-amtz.in)

For Admissions:

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