

# ISO 13485:2016 Lead Auditor Training program on Medical Device Quality Management System

Become a Global Leader in  
Medical Device Quality!

Unlock your potential in the fast-growing **Medical Device Industry** with the **Exemplar Global International Auditors and Training Provider Certification Body**.

**ELIGIBILITY:** Medical Device Manufacturers, Quality professionals and engineers, Regulatory Professionals, Internal and External Auditors and consultants.

**MODE:** Online | **DURATION:** 5 Days

## KEY CONTENTS:

- Overview of ISO 13485:2016 – Structure, Clauses & Requirements
- Audit Planning, Execution, Reporting, and Follow-up
- Non-Conformance Identification and Corrective Action Management
- Roles & Responsibilities of Lead Auditors and Audit Teams

**Advance Your Expertise. Lead With Quality. Become Globally Certified !**

