

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 Accredited ISO 13485:2016 Lead Auditor Certification**.

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 !

For more information: K. Durgankitha | +91 9281079255 | trainings@mqci.in



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