



ISO 13485 Certification Training: Master Medical Devices Quality Management System

28 Apr - 02 May 2026
Madrid



ISO 13485 Certification Training: Master Medical Devices Quality Management System

Ref.: 103600269_44692 **Date:** 28 Apr - 02 May 2026 **Location:** Madrid **Fees:** 5700 **Euro**

Course Overview

ISO 13485 certification is a globally recognized standard for quality management systems in medical devices. This course offers a comprehensive understanding of ISO 13485 compliance, regulatory requirements, and best practices.

Participants will learn how to implement ISO 13485 to ensure their organizations meet regulatory demands and enhance product safety. The training covers key topics including differences between ISO 13485 and FDA 21 CFR 820, risk management, documentation, and audit processes.

Target Audience

- Quality Managers responsible
- Regulatory Affairs Specialists
- ISO 13485 Consultants
- Medical Device Engineers
- Internal & Lead Auditors
- Manufacturing & Production Managers

Targeted Organizational Departments

- Quality Assurance & Control
- Regulatory Compliance
- Manufacturing & Production
- Research & Development
- Supply Chain & Procurement

Targeted Industries

- Medical Device Manufacturing
- Pharmaceuticals & Biotechnology
- Healthcare & Hospitals
- Regulatory & Certification Bodies
- Testing & Calibration Laboratories

Course Offerings

By the end of this course, participants will be able to:

- Understand ISO 13485 standard requirements and its role in medical device quality assurance
- Implement ISO 13485 best practices to enhance product safety
- Conduct internal audits using the ISO 13485 audit checklist
- Differentiate between ISO 13485 vs FDA 21 CFR 820 compliance requirements
- Develop and manage ISO 13485 documentation requirements
- Gain insights into ISO 13485 accreditation and certification steps
- Become an ISO 13485 lead auditor or ISO 13485 lead implementer

Training Methodology

- Case Studies: Real-world examples of ISO 13485 compliance failures & successes
- Workshops & Exercises: Hands-on ISO 13485 certification training activities
- Group Discussions: Peer-to-peer knowledge exchange on ISO 13485 QMS training
- Mock Audits: Practice applying ISO 13485 audit training techniques

Course Toolbox

- ISO 13485 implementation guide & templates
- ISO 13485 compliance checklist for internal audits
- ISO 13485 regulatory requirements reference guide
- ISO 13485 certification exam prep materials

Course Agenda

Day 1: Introduction to ISO 13485 & Regulatory Framework

- **Topic 1:** Overview of ISO 13485 certification and its importance in the medical devices industry
- **Topic 2:** Understanding the medical devices quality management system QMS
- **Topic 3:** Key regulatory requirements and compliance standards for ISO 13485
- **Topic 4:** ISO 13485 vs FDA 21 CFR 820: Key differences and alignment strategies
- **Topic 5:** ISO 13485 documentation requirements: Essential records and reports
- **Topic 6:** ISO 13485 certification process: Step-by-step guide to implementation
- **Reflection & Review:** Key takeaways, Q&A, and discussion on regulatory challenges



Day 2: Implementation of ISO 13485 Quality Management System

- **Topic 1:** ISO 13485 standard requirements: Understanding the structure and clauses
- **Topic 2:** Developing an ISO 13485 implementation guide for organizations
- **Topic 3:** Risk management and preventive measures in ISO 13485 compliance
- **Topic 4:** Internal audits and ISO 13485 compliance checklist for effective monitoring
- **Topic 5:** Managing ISO 13485 accreditation and certification requirements
- **Topic 6:** Integrating ISO 13485 with other quality standards for seamless compliance
- **Reflection & Review:** Group activity and discussion on implementation challenges

Day 3: ISO 13485 Lead Auditor & Internal Audit Processes

- **Topic 1:** Understanding the role of an ISO 13485 lead auditor and internal auditor
- **Topic 2:** Preparing an ISO 13485 audit checklist for internal and external audits
- **Topic 3:** Conducting ISO 13485 audit training: Key skills and techniques
- **Topic 4:** Handling non-conformities and corrective action processes
- **Topic 5:** Reporting and documentation of audit findings in ISO 13485 compliance
- **Topic 6:** ISO 13485 certification exam preparation and mock assessment
- **Reflection & Review:** Interactive session on audit best practices and case studies

Day 4: Advanced ISO 13485 Implementation & Process Optimization

- **Topic 1:** Best practices for ISO 13485 QMS training and workforce development
- **Topic 2:** Supplier management and control in ISO 13485 compliance
- **Topic 3:** Enhancing medical device quality assurance through continuous improvement
- **Topic 4:** ISO 13485 lead implementer strategies for effective QMS deployment
- **Topic 5:** Documentation and record-keeping requirements for regulatory audits
- **Topic 6:** Addressing ISO 13485 certification cost and budget considerations
- **Reflection & Review:** Discussion on real-world implementation success stories

Day 5: Certification Exam & Practical Application

- **Topic 1:** Final preparation for the ISO 13485 certification exam
- **Topic 2:** Conducting a full-scale internal audit simulation
- **Topic 3:** Common mistakes and challenges in ISO 13485 implementation
- **Topic 4:** ISO 13485 consulting services and career opportunities
- **Topic 5:** Developing a roadmap for ISO 13485 accreditation and long-term compliance
- **Topic 6:** Continuous improvement and maintaining compliance post-certification
- **Reflection & Review:** Course wrap-up, certification guidance, and Q&A

FAQ

- **What specific qualifications or prerequisites are needed for participants before enrolling in the course?**



No prior certification is required, but experience in medical device quality assurance or ISO 13485 compliance is recommended.

- **How long is each day's session, and what is the total course duration?**

Each session lasts 4-5 hours, with breaks and interactive discussions. The total course spans five days, approximately 20-25 hours.

- **What is the cost of obtaining ISO 13485 certification, and what factors affect it?**

The ISO 13485 certification cost varies based on factors such as training provider fees, company size, and accreditation body charges. Contact us for detailed pricing.

How This Course is Different from Other ISO 13485 Training Courses

This ISO 13485 certification training goes beyond standard courses by offering:

- Real-world case studies on ISO 13485 compliance
- Hands-on exercises and mock audits for skill development
- Training for roles like ISO 13485 lead auditor and internal auditor
- Flexible online learning options

With expert trainers and a structured curriculum, this course prepares professionals for ISO 13485 accreditation and medical device industry certification.



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**Environment &
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**Governance, Risk and
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WHO WE ARE

Agile Leaders is a renowned training center with a team of experienced experts in vocational training and development. With 20 years of industry experience, we are committed to helping executives and managers replace traditional practices with more effective and agile approaches.

OUR VISION

We aspire to be the top choice training provider for organizations seeking to embrace agile business practices. As we progress towards our vision, our focus becomes increasingly customer-centric and agile.

OUR MISSION

We are dedicated to developing value-adding, customer-centric agile training courses that deliver a clear return on investment. Guided by our core agile values, we ensure our training is actionable and impactful.

WHAT DO WE OFFER

At Agile Leaders, we offer agile, bite-sized training courses that provide a real-life return on investment. Our courses focus on enhancing knowledge, improving skills, and changing attitudes. We achieve this through engaging and interactive training techniques, including Q&As, live discussions, games, and puzzles.



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