



ISO 37301 Compliance Management Lead Auditor Training

11 - 15 May 2027
Dubai



ISO 37301 Compliance Management Lead Auditor Training

Ref.: 103600629_82208 **Date:** 11 - 15 May 2027 **Location:** Dubai **Fees:** 6500 **Euro**

Course Overview:

The ISO 37301 Lead Auditor Training for Compliance Management Systems is an advanced professional programme designed to develop the knowledge and practical auditing competence required to assess a Compliance Management System CMS against ISO 37301 requirements. This ISO 37301 Lead Auditor Course combines Compliance Management System Training with structured ISO 37301 Audit Training, enabling participants to understand compliance principles, evaluate CMS requirements, prepare audits, collect and assess audit evidence, report findings, and manage audit programmes.

The course addresses the complete audit lifecycle, from understanding organizational context, compliance obligations, risk assessment, governance, leadership, controls, monitoring, and continual improvement to preparing and conducting first-, second-, and third-party audits. ISO 37301 emphasizes a systematic, risk-based approach to managing legal, regulatory, contractual, and voluntary obligations, supported by governance, monitoring, and continuous improvement.

Participants in this ISO 37301 Auditor Training programme develop practical capabilities in audit planning, Stage 1 and Stage 2 audit activities, interviewing, sampling, evidence evaluation, nonconformity reporting, closing meetings, corrective-action follow-up, and audit programme management. The structure also reflects the seven competency domains identified for ISO 37301 Lead Auditor professionals, covering CMS principles, requirements, audit concepts, audit preparation, audit execution, audit closure, and audit programme management.

Target Audience:

- Compliance Managers and Compliance Officers
- Internal Auditors and Lead Auditors
- Risk and Governance Professionals
- Legal and Regulatory Compliance Specialists
- Management System Auditors
- Ethics and Integrity Professionals
- GRC Professionals
- Quality and Integrated Management System Professionals
- Consultants involved in Compliance Management System Audit activities
- Professionals preparing for ISO 37301 Lead Auditor responsibilities
- Managers responsible for Corporate Compliance Training and oversight
- Professionals involved in Compliance Monitoring Training and assurance



Targeted Organizational Departments:

- Compliance and Ethics
- Internal Audit and Assurance
- Governance, Risk, and Compliance
- Legal and Regulatory Affairs
- Enterprise Risk Management
- Quality and Management Systems
- Corporate Governance
- Internal Controls
- Policy and Standards
- Business Integrity and Investigations

Targeted Industries:

- Banking, Financial Services, and Insurance
- Oil, Gas, Energy, and Utilities
- Healthcare and Pharmaceuticals
- Telecommunications and Technology
- Government and Public Sector
- Construction and Infrastructure
- Manufacturing and Industrial Operations
- Transportation and Logistics
- Professional and Business Services

Course Offerings:

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

- Interpret the fundamental principles and requirements of ISO 37301.
- Understand the structure and operation of an effective Compliance Management System.
- Evaluate organizational context, compliance obligations, risks, governance, and controls.
- Apply professional auditing principles to a Compliance Management System Audit.
- Differentiate between first-party, second-party, and third-party audits.
- Establish audit objectives, scope, criteria, and feasibility.
- Prepare audit plans, working documents, and evidence-gathering approaches.
- Conduct Stage 1 and Stage 2 ISO 37301 audit activities.
- Apply interviewing, observation, sampling, document review, and evidence evaluation techniques.
- Identify conformity, nonconformity, observations, and areas requiring improvement.
- Prepare professional audit findings, conclusions, and audit reports.
- Conduct opening meetings, closing meetings, and audit follow-up activities.

Training Methodology:

The ISO 37301 Lead Auditor Training uses an applied, scenario-based methodology that combines technical interpretation with practical audit activities. Instructor-led sessions introduce the principles of ISO 37301, the structure of a Compliance Management System, compliance risks, governance arrangements, controls, monitoring, and continual improvement. Participants then apply these concepts through guided audit scenarios based on realistic organizational situations.

Case studies are used to examine compliance obligations, risk registers, control environments, leadership responsibilities, reporting mechanisms, and compliance performance. Group exercises enable participants to define audit objectives, scope, and criteria; prepare audit plans; review documented information; develop interview questions; and assess objective evidence. Role-based simulations allow participants to practise opening meetings, auditor interviews, audit-team discussions, conflict management, presentation of findings, and closing meetings.

The methodology also includes practical exercises in distinguishing major and minor nonconformities, drafting audit findings, evaluating corrective actions, and preparing audit reports. The PECB Lead Auditor framework places substantial emphasis on conducting audits, preparing audits, applying audit principles, and evaluating evidence, which is reflected throughout the learning activities.

Throughout the ISO 37301 Training Course, participants receive structured feedback to strengthen professional judgment, evidence-based decision-making, and confidence in conducting ISO 37301 Compliance Audit activities.

Course Toolbox:

Participants will receive practical guidance, examples, and insights related to:

- ISO 37301 clause interpretation reference structure
- Compliance obligations register examples
- Compliance risk assessment examples
- Compliance control matrix examples
- CMS audit planning examples
- Audit objectives, scope, and criteria guide
- Stage 1 audit review framework
- Stage 2 audit activity examples
- Audit interview question examples
- Audit evidence classification guide

Note: Tools, software, templates, or proprietary systems are **not provided** as part of the course.

Course Agenda:

Day 1: ISO 37301 and Compliance Management System Fundamentals

- **Topic 1:** ISO 37301 Lead Auditor role, certification context, and audit responsibilities
- **Topic 2:** Principles, scope, terminology, and structure of ISO 37301
- **Topic 3:** Compliance obligations, organizational context, and interested parties
- **Topic 4:** Compliance risk assessment and the risk-based approach to CMS
- **Topic 5:** Leadership, governance, compliance culture, ethics, and accountability
- **Topic 6:** ISO 37301 requirements for planning, support, operations, evaluation, and improvement
- **Reflection & Review:** Connect ISO 37301 requirements with the responsibilities of an auditor when evaluating the design, implementation, and effectiveness of a CMS.

Day 2: Audit Principles and ISO 37301 Audit Preparation

- **Topic 1:** Fundamental audit concepts, principles, ethics, and professional conduct
- **Topic 2:** First-party, second-party, and third-party Compliance Management System Audits
- **Topic 3:** Risk-based auditing, materiality, reasonable assurance, and professional judgment
- **Topic 4:** Defining audit objectives, scope, criteria, feasibility, and audit engagement
- **Topic 5:** Building the audit plan, audit timetable, working documents, and test plans
- **Topic 6:** Audit-team roles, technical experts, responsibilities, and communication protocols
- **Reflection & Review:** Review the decisions required to transform an audit mandate into a structured and risk-focused ISO 37301 audit plan.



Day 3: Conducting the ISO 37301 Audit

- **Topic 1:** Stage 1 audit activities and review of CMS documented information
- **Topic 2:** Opening meetings, audit communication, guides, observers, and stakeholder interaction
- **Topic 3:** Stage 2 audit activities and evaluation of CMS implementation
- **Topic 4:** Audit interviews, observation, document review, analysis, and technical verification
- **Topic 5:** Audit sampling, evidence collection, corroboration, and objective evaluation
- **Topic 6:** Managing audit challenges, conflicting evidence, communication issues, and emerging risks
- **Reflection & Review:** Evaluate how auditors move from documented information to reliable, sufficient, and objective audit evidence.

Day 4: Audit Findings, Reporting, and Closing the Audit

- **Topic 1:** Evaluating evidence and developing defensible audit findings
- **Topic 2:** Conformity, major and minor nonconformity, anomalies, and observations
- **Topic 3:** Drafting clear, evidence-based nonconformity statements
- **Topic 4:** Preparing audit conclusions and conducting quality review of audit documentation
- **Topic 5:** Closing meetings, presenting findings, and communicating audit conclusions
- **Topic 6:** Audit reporting, certification recommendations, corrective actions, and follow-up
- **Reflection & Review:** Review how audit evidence is converted into accurate findings, conclusions, and professional audit reports.

Day 5: Managing the ISO 37301 Audit Programme and Lead Auditor Practice

- **Topic 1:** Establishing and managing an ISO 37301 audit programme
- **Topic 2:** Applying the PDCA cycle to audit programme management
- **Topic 3:** Managing audit resources, records, confidentiality, and documented information
- **Topic 4:** Evaluating auditor competence, performance, professional behaviour, and audit-team effectiveness
- **Topic 5:** Combined audits, surveillance audits, recertification, and certification lifecycle considerations
- **Topic 6:** Integrated Lead Auditor case study covering planning, conducting, closing, and following up an ISO 37301 audit
- **Reflection & Review:** Consolidate the complete ISO 37301 Lead Auditor process and identify the key decisions required to manage effective, credible, and risk-based CMS audits.

FAQ:



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

There are no universal mandatory prerequisites for attending the training course itself; however, participants will benefit from prior knowledge of compliance management, governance, risk management, internal audit, legal or regulatory compliance, or management-system standards. Familiarity with ISO 37301 requirements and general auditing principles is strongly recommended for participants seeking to progress toward professional Lead Auditor responsibilities.

Professional certification requirements are separate from course attendance. For example, the PECB Lead Auditor credential requires relevant professional and audit experience in addition to passing the applicable examination.

How long is each day's session, and is there a total number of hours required for the entire course?

Each day's session is generally structured to last around 4-5 hours, with breaks and interactive activities included. The total course duration spans five days, approximately 20-25 hours of instruction.

What is the difference between understanding ISO 37301 and being able to audit a Compliance Management System?

Understanding ISO 37301 focuses primarily on interpreting the requirements of the standard and how a Compliance Management System should be established and operated. An ISO 37301 Lead Auditor must go further by applying audit principles, defining audit scope and criteria, evaluating risks, collecting objective evidence, conducting interviews, applying sampling techniques, identifying nonconformities, preparing audit conclusions, reporting findings, and managing audit follow-up activities.

This distinction is reflected in the Lead Auditor competency framework, which separates knowledge of CMS principles and requirements from the practical competencies required to prepare, conduct, close, and manage ISO 37301 audits.



How This Course is Different from Other ISO 37301 Lead Auditor Courses:

The ISO 37301 Lead Auditor Training for Compliance Management Systems is designed around the real work of an auditor rather than limiting the learning experience to clause-by-clause interpretation. While many ISO 37301 Certification Training programmes focus heavily on theoretical requirements, this course integrates Compliance Management System Training with practical Compliance Audit Training across the complete audit lifecycle.

Participants work with the relationship between compliance obligations, compliance risks, governance arrangements, policies, controls, monitoring mechanisms, investigations, internal audit, management review, and continual improvement. This reflects how an actual CMS operates as an integrated organizational system. ISO 37301-based compliance management depends not only on documented requirements but also on leadership, risk assessment, controls, monitoring, culture, and continual improvement.

The course also places strong emphasis on audit preparation, professional judgment, Stage 1 and Stage 2 activities, evidence collection, sampling, interviewing, nonconformity reporting, audit conclusions, corrective-action evaluation, and audit programme management. These areas align directly with the seven competency domains identified for ISO 37301 Lead Auditor professionals.



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