

## Kelmac Group® Delivers Specialized ISO 13485 Internal Auditor Training for Medical Device Quality Professionals



**Limerick, Ireland Aug 19, 2026 ([IssueWire.com](https://www.issuewire.com))** - Kelmac Group®, a global provider of ISO training, consulting and auditing services, has announced the availability of its specialized [ISO 13485 internal auditor training](#) for professionals working in medical device quality, compliance, regulatory, and quality management functions.

The training is designed to help participants develop the knowledge and practical skills required to conduct effective internal audits within an ISO 13485 quality management system. The programme focuses on understanding applicable requirements, preparing and conducting audits, gathering objective evidence, identifying findings, and supporting corrective action.

**“Effective internal auditing is an important part of maintaining and continually improving a medical device quality management system. Our training is designed to help professionals move beyond understanding requirements and develop the practical skills needed to apply auditing techniques confidently in their organizations,”** said a representative of Kelmac Group®.

**Building Practical ISO 13485 Auditing Skills**

Medical device organizations rely on effective quality management systems to support consistent processes and meet applicable regulatory and customer requirements. Internal auditors play an important role by independently evaluating processes and identifying opportunities for improvement.

Kelmac Group® ISO 13485 internal auditor training addresses key areas of the internal audit process, including:

- Understanding ISO 13485 requirements
- Internal audit planning and preparation
- Developing audit checklists
- Conducting effective audit interviews
- Gathering and evaluating objective evidence
- Identifying and documenting nonconformities
- Preparing audit reports
- Evaluating corrective actions
- Supporting continual improvement

The training is intended for professionals who have responsibilities related to quality management systems, regulatory compliance, quality assurance, internal auditing, and medical device operations.

### **Supporting Medical Device Quality and Compliance**

As organizations continue to strengthen their quality and compliance processes, developing competent internal auditors can help create a more effective approach to monitoring and improving quality management systems.

The ISO 13485 internal auditor training from Kelmac Group® provides professionals with an opportunity to strengthen their auditing capabilities and apply structured audit techniques within medical device quality environments.

The programme can be particularly relevant to quality managers, quality assurance professionals, regulatory and compliance personnel, internal auditors, consultants, and other professionals involved in ISO 13485 implementation and maintenance.

### **About Kelmac Group®**

[Kelmac Group®](#) is a global provider of conformity assessment, regulatory compliance, risk management, ISO consulting, auditing, and professional training services.

Kelmac Group® provides training and professional services across a wide range of international standards and management systems, helping organizations and professionals strengthen their knowledge, competence, compliance, and quality performance.

Through its specialized training programmes, Kelmac Group® continues to support professionals across industries, including organizations operating within the medical device sector.

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