IPI Academy

MANAGEMENT forum



Presented by
Management Forum

A Regulatory Update on the Medical Device and In-Vitro Diagnostic Regulations in the EU and UK

This course will provide an excellent opportunity to hear the latest developments from regulatory experts, and enable participants to discuss the implications of the regulations on working practices and the potential impact on future regulatory strategies.

Format:
Bespoke training

CPD:
6 hours for your records (depending on your requirements)

Certificate of completion

Course overview

Stay informed on the significant changes since the adoption of the Medical Device Regulation (MDR) and the In-Vitro Diagnostic Regulation (IVDR).

The landscape of medical device and in-vitro diagnostic regulations in the EU and UK has undergone significant transformation in recent years. With the adoption of the Medical Device Regulation (MDR) and the In-Vitro Diagnostic Regulation (IVDR),, the regulatory framework has become more rigorous and complex. Staying abreast of these changes is crucial for professionals involved in regulatory affairs, quality assurance, and related fields to ensure compliance and maintain market access.

This course is essential for those seeking to stay fully updated on the latest regulatory developments.

Key topics to be addressed:

- Comprehensive insights into the rephrased implementations of MDR and IVDR
- Understand the new guidelines
- Learn about the role and output of the Medical Device Coordination Group (MDCG)
- Recent medical device regulatory developments in the UK, providing a comparative perspective on the evolving regulatory landscape
- Explore potential future developments and the evolving role of the European Medicines Agency (EMA) in the context of medical device regulations

This course offers a unique opportunity to hear from regulatory experts, discuss the implications of current and forthcoming regulations on working practices, and understand the potential impact on future regulatory strategies.

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Please note: This programme has been designed to reflect the most current understanding of the regulatory landscape for medical devices and IVDs. The content will be delivered using the latest available information at the time of presentation, adapting to any regulatory developments that emerge beforehand. Attendees can expect a clear overview of current requirements, insight into forthcoming regulatory changes, and informed discussion of future developments, including evolving controls on software, AI-supported devices, and healthcare applications.

Benefits of attending

- **Stay updated** on MDR and IVDR implementations, new guidelines, and recent UK regulatory developments
- **Explore** the changes in the UK regulation as its system becomes independent and moves away from the EU rules
- **Navigate** the increasingly complex world of device and diagnostic regulations
- **Gain** insights into future changes to the regulations and how "digital medicine" may be regulated in the future
- **Learn** from leading regulatory experts and gain practical advice

Who should attend

This training course has been designed for all those working or interested in the medical device and diagnostic market in the EU and the UK, including medical device and in-vitro diagnostic professionals from the following areas:

- Regulatory affairs
- Clinical studies
- Vigilance
- Post-market surveillance
- Quality systems
- Technical support and business development

Reviews

★★★★★

Was very useful. I feel I accomplished further knowledge about the implementation of MDR and IVDR and also what is on the horizon in terms of the UK regulations from the course. The content of update on IVDR was great as it was directly related to my work. I also enjoyed learning more about the history of the regulations/legislative process and gained some useful background knowledge on combination products.

Liz Gillman
Clinical Compliance Project Lead
LifeScan Scotland
Apr 30 2025

★★★★★

Very good course.

Richard Jones
Senior Medical Device Auditor
Eurofins E&E CML Limited
Oct 22 2024

★★★★★

Very well run, met my objectives, and speakers were interesting, approachable and very knowledgeable.

Lucie Green
QA Manager
Vyaire Medical Products
Sep 24 2019

★★★★★

Excellent training performed by very knowledgeable speakers.

Marie-Pierre Hontas
Senior Director Scientific & Clinical Affairs
Vexim SA – Stryker IVS
Mar 6 2019

Run this programme in-house for your whole team

Coming to IPI Academy for your in-house training provides an all-inclusive service which gives you access to a wide variety of content, learning platforms and delivery mechanisms as well as your own personal training adviser who will work with you from the initial enquiry through to feedback and follow-up after the programme.

With over 600 trainers, all practitioners and experts across a huge range of fields, we can provide the training you need, where you need it, when you need it, and at a price which suits your budget. Our approach to tailored learning and development consists of designing and delivering the appropriate solution for each client.

For your FREE consultation and to find out more about how we can work with you to solve your training needs, please contact our training advisers:



ALEKSANDRA BEER
Tel: +44 (0)20 7749 4749
Email:
inhouse@ipiacademy.com



YESIM NURKO
Tel: +44 (0)20 7749 4749
Email:
inhouse@ipiacademy.com



Harry ALTAMONT
Tel: +44 (0)20 7749 4749
Email:
inhouse@ipiacademy.com

IPI Academy

IPI Academy is a training initiative of Falconbury and Management Forum; leading providers of industry training for over 30 years, based in the UK.

10-12 Rivington Street
London EC2A 3DU

ipi.academy
Tel: +44 (0)20 7749 4749
Email: info@ipiacademy.com