IPI Academy

MANAGEMENT forum



Presented by
Management Forum

ICH GCP E6 (R3): What You Need to Know for Implementation and Inspection

This one-day course focuses on what inspectors look for in clinical trials, covering ICH GCP E6(R3), the EU Clinical Trials Regulation, common findings, and practical steps to ensure compliance and inspection readiness.

Format:
Bespoke training

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

Certificate of completion

Course overview

This course focuses on the practical implications of the ICH GCP E6 (R3) guideline and what it means for the conduct and oversight of modern clinical trials. While the core principles of GCP remain unchanged, E6 (R3) introduces clearer expectations around risk-based approaches, oversight, roles and responsibilities, and the use of data and technology.

The course is designed to help clinical research professionals understand what has changed, how organisations are applying the new requirements in practice, and what regulators and inspectors are likely to focus on during GCP inspections.

Through practical examples, the session explores how sponsors, CROs and investigator sites are adapting governance, oversight, and decision-making under E6 (R3), with particular focus on inspection readiness.

By the end of the course, delegates will have a clear understanding of how to implement ICH GCP E6 (R3) in a practical and inspection-ready way.

Benefits of attending

By attending this course, delegates will be able to:

- **Understand** the key updates and principles of ICH GCP E6 (R3)
- **See** how the E6 (R3) requirements are being implemented by companies, using practical examples
- **Understand** how GCP inspections are evolving under E6 (R3) and what inspectors are likely to focus on
- **Apply** risk-based approaches to clinical trial design, oversight, and monitoring
- **Share** experiences and insights with peers facing similar implementation challenges

Who should attend

This course is suitable for anyone involved in the planning, conduct, oversight, or support of clinical trials. It is applicable across Pharma, Biopharma, Clinical Research Organisations (CROs), service providers, and investigator sites. While focused on ICH GCP E6(R3), the principles discussed are relevant to any organisation involved in the conduct and oversight of regulated clinical investigations.

The course is particularly relevant for:

- Clinical Research Associates (CRAs)
- Clinical Trial Managers and Project Managers
- Regulatory Affairs Professionals
- Quality Assurance Personnel
- Investigators and Site Staff
- Data Managers
- Anyone involved in the design, execution, oversight, or governance of clinical trials

Reviews

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Denise Johnson
Clinical Research Manager
SOMNOmedics AG
Jan 29 2026

★★★★☆

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Denise Johnson
Clinical Research Manager
SOMNOmedics AG
Jan 29 2026

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:



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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.

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