



Presented by
Management Forum

Navigating ICH GCP E6 (R3): What You Need to Know

12 September 2025
+ 15 January 2026

This brand new course on the recently published and long-awaited ICH GCP E6 (R3) is designed to help participants stay compliant and implement the new guideline effectively.



Format:
Live online



CPD:
6 hours for your records



Certificate of
completion

Course overview

The ICH GCP E6 (R3) guideline represents a significant evolution in the conduct of clinical trials, emphasising flexibility, risk-based approaches, and the integration of innovative technologies. This one-day training course is designed to provide clinical research professionals with a comprehensive understanding of the updated guidelines, ensuring compliance and enhancing the quality of clinical trials.

Through a combination of expert-led lectures and group discussions, you will engage in practical exercises to reinforce key concepts and application to real-world scenarios. This training is ideal for all clinical research professionals committed to maintaining GCP compliance in their work. By the end of the course, you will be well-prepared to navigate the regulatory requirements and effective implementation of the new GCP requirements.

Benefits of attending

- **Understand** the key updates and principles of ICH GCP E6 (R3)
- **Recognise** the roles and responsibilities of stakeholders, including sponsors, investigators, and ethics committees
- **Implement** risk-based approaches in clinical trials
- **Prepare** for regulatory inspections and ensure ongoing compliance
- **Stay** compliant and ahead of GCP regulatory changes
- **Gain** practical strategies to implement ICH GCP (R3)
- **Network** and share insights with peers

Who should attend?

This session will be beneficial to anybody who is involved in clinical research. It is applicable to pharma, clinical research organisations (CROs), service providers and investigator sites.

This course is ideal 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 planning, execution, or oversight of clinical trials

Programme

Introduction to ICH GCP E6 (R3)

Core principles of ICH GCP E6 (R3)

Stakeholder responsibilities

- Roles of Institutional Review Boards (IRBs)/Independent Ethics Committees (IECs)
- Investigator obligations
- Sponsor responsibilities
- Service providers

Risk-based approaches

Data governance and integrity

Data and records

Investigator brochures

Protocols

Essential records

Preparing for regulatory inspections

Implementation considerations

Presenter




Laura Brown

Dr Laura Brown is an independent QA and training consultant and was director of the MSc in Clinical Research, School of Pharmacy, University of Cardiff. Laura has many years experience of managing GCP inspections in the pharmaceutical industry and has worked for several leading companies including GSK, Hoechst Marion Roussel, Good Clinical Research Practices and Phoenix International. She has worked as a clinical research manager, audit director and head of a training department. She is an international expert on GCP and clinical trial requirements and was chair of the Institute of Clinical Research GCP Forum for six years. Laura writes regularly on clinical research regulatory requirements and has written a chapter in International Pharmaceutical Product Registration and several articles on the EU Clinical Trial Regulation, and ICH GCP R2 and R3.


Course dates


12 September 2025	Live online 09:30-17:00 UK (London) (UTC+01) <i>Course code 15666</i>	GBP 649 749 EUR 909 1,049 USD 1,043 1,199 Until 08 Aug
15 January 2026	Live online 09:30-17:00 UK (London) (UTC+00) <i>Course code 15667</i>	GBP 649 749 EUR 909 1,049 USD 1,043 1,199 Until 11 Dec

How to book

 **Online:**
ipi.academy/3261

Alternatively contact us to book, or if you have any queries:

 **Email:**
info@ipiacademy.com

 **Phone:**
[+44 \(0\)20 7749 4749](tel:+442077494749)

Discounts

- Booking more than one delegate on any one date qualifies for a **30% discount** on the second and subsequent places.
- Most events qualify for an **early booking discount** prior to 6 weeks before the course date. Be sure to check on our website, where the latest discounts will be shown.

Further information

Fee
The fee includes all meals and refreshments for the duration of the course (for venue-based courses) and a complete set of course materials (provided electronically). If you have any particular requirements, please advise customer services when booking.

Please note
IPI Academy (and our training partners) reserve the right to change the content and timing of the programme, the speakers, the date and venue due to reasons beyond their control. In the unlikely event that the course is cancelled, we will refund the registration fee and disclaim any further liability.

Terms and conditions
The rest of our terms, the event cancellation policy and the terms and conditions are on our website, please visit ipi.academy/content/terms-and-conditions

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IPI
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