



Course Overview

This GDPR course provides a practical overview of data protection, focusing on clinical research compliance. It covers key principles, lawful processing, special category data, and roles and responsibilities, plus links to the Clinical Trials Regulation and Good Clinical Practice. You'll also learn about records of processing, consent, data subject rights, security, retention, and breach reporting, ending with a final exam to consolidate learning.

Learning Objectives

- Understand what is meant by Data Protection and the privacy principles
- Gain an understanding of GDPR
- Understand the implications of GDPR for citizens including data subject rights
- Understand the implications of GDPR for Clinical Research
- Discover what you need to do to ensure compliance with the GDPR

Module One: Data Protection and GDPR

- GDPR: Legislation and Background
- CTR & CTIS – GDPR compliance points
- GDPR vs GCP
- Article 6
- Article 9
- Key GDPR Concepts and Principles
- Article 5
- Roles and Responsibilities

Module Two: GDPR for Clinical Trials

- ROPA
- GDPR Around the World
- Data Protection Impact Assessments (DPIAs)
- Rules for Consent
- Data Subject Rights
- Data Security and Anonymisation
- Data retention and deletion
- Breach notification and incident

Duration: Approx. 3 hours

As with all of our courses, you are able to take this in your own time, on any device and are able to pause and restart from the place you left off. Time taken will vary depending on your experience and how many of the many optional links and activities you choose to view.