Clinical Research Associate (CRA) Training

Training Online 4U

About This Course

This interactive and engaging course was developed by industry experts with a wealth of clinical trial experience. It goes through the whole clinical trial process and also includes aspects of Good Clinical Practice and Good Documentation Practice/Essential Documents.

Who is the course suited to?

It is aimed at those working within clinical research in roles such as Clinical Trial Associate/Admin who wish to progress to, or have recently become, a CRA/monitor and want to receive full training/certification.

Module 1: Introduction

- The role of a CRA
- Clinical Trial Phases
- Ethics Committees
- Regulatory and Advisory Bodies
- Good Clinical Practice
- Good Documentation Practice

Module 2: Start-up

- Before a Trial
- Monitoring Visits
- Site Selection
- Essential Documents
- Document Control
- Regulatory/Ethics submissions
- Informed Consent

Module 3: During Study

- During a Trial
- Site Initiation Visits
- Monitoring Approaches
- Patient Safety
- Safety Reporting
- Documentation/ Audits
- Communication

Module 4: Closeout

- Trial Closeout
- Closeout Visit
- Communication (End of Trial)
- Documentation
- Archiving
- Attributes, Tips & Tricks
- Career Progression

Duration: Approx. 8 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.

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