Clinical Project Management Training

Training Online 4U

About This Course

This interactive and engaging course was developed by industry experts with a wealth of clinical trial experience. This course will explain in everything you need to know about becoming an effective project manager within this field, from study start up right through to closeout. It also includes aspects of GCP and Good Documentation Practice/Essential documents.

Who is the course suited to?

It is aimed at those working within clinical research as a project manager, or those wishing to progress in to a project management role in this field. The content of the course includes, but is not limited to the topics outlined below:

Module 1: Introduction
- What is Project Management?
- The Role of a Project Manager
- Clinical Trial Phases
- Ethics Committees
- Regulatory and Advisory Bodies

Module 2: Start-up
- Before a Trial
- Project Planning
- Site Selection
- Essential Documents
- Overseeing Monitors
- Effective Teams
- Budget Management
- Vendor Management

Module 3: During study
- During a Trial
- Site Initiation Visits
- Supporting Monitors
- Patient Safety & Safety Reporting
- Communication
- Issue Management
- Audits & Inspections

Module 4: Closeout
- Trial Closeout
- Closeout Visit
- Communication
- Documentation
- Archiving
- Key Attributes, Tips & Tricks

Price: £349 inc. VAT (discount available for 6 or more users).

Duration: Approx. 8hours

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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