Good Documentation Practice and Data Integrity

Training Online 4U

About This Course

This professionally developed Good Documentation Practice (GDocP) course explores data integrity and document requirements in detail and will show how GDocP can be implemented into a number of roles. Good Documentation Practice is mandatory to ensure that documentation (and ultimately products) meet industry standards and other legal responsibilities in the pharmaceutical sector.

Who is the course suited to?

This is a professionally developed, interactive and engaging online course essential for anyone working in Clinical Research. It is suitable for all levels of experience.

Module 1

- Introduction to GDocP
- Documents
- Data
- Document Requirements

Module 2

- Data Integrity
- Documentation Control
- Electronic Records

Module 3

- Corrections
- Common Errors
- Security and Data Protection

Price: £99 inc. VAT (discount available for 6 or more users).

Duration: Approx. 2 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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Finished Product - A medicinal product which has undergoneall stages of production, including packaging in its final container.

Manufacture- All operations of purchase of materials andproducts, production, Quality Control, release,

storage, distribution of medicinal products and the relatedcontrols.

Production – All operations involved in the preparation of amedicinal product, from receipt of materials,

through processing and packaging, to its completion as afinished product.

Validation- Action of proving, in accordance with the principles ofGood Manufacturing Practice, that any procedure, process, equipment, material, activity or system leads to expected results.