ABC of Quality Control

R 3,500 Incl VAT (Non -WHC)
R 2,390 Excl VAT (WHC)
Discounts apply to groups

Classroom based

1½ days

Running a clinical trial is a complex and detailed process. Maintaining accurate records and quality data throughout a clinical trial is a continual, dynamic process. Trial requirements are carefully prescribed in detailed documents such as the protocol, the ICH and SA GCP guidelines, data management plan and the project plan.

The course has 5 modules including a "Test your knowledge" at the end of each module:

- Introduction
- Module 1: Site Organisation
- Module 2: Site Preparation
- Module 3: Site Conduct
- Module 4: Study Close Out and Archiving
- Module 5: Audits and Inspections

This practical 1½ day course designed and presented by Professor Lesley Burges will equip clinical research site personnel with the relevant knowledge and tools to ensure successful study conduct from start to finish, in preparation for monitoring audits and site inspections.

At the end of this course, learners will be expected to complete a final assessment:

- There are 50 multiple choice questions.
- The pass mark is 70%.
- Learners have (2) attempts.
- A certificate will be issued on passing the assessment.
- CPD Points will be issued on completion.

Clinical research personnel such as investigators, study coordinators, and project managers with a valid GCP certificate and preferably some work experience in the clinical research field.

Parktown, Johannesburg

In-house training and travel to other areas will be considered subject to viability.

+27 11 274 9327/9256/9368

http://www.academicadvance.co.za

training@academicadvance.co.za