Clinical Research

ABC of **Quality** Control for Clinical **Trial** Sites

Introduction

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.

Course Overview

This practical 1½ day course designed and presented by Professor Lesley Burgess 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.

Course Content

- Site Organisation
- Study Preparation
- Study Conduct
- Study Close Out & Archiving
- Audits & Inspections

Accreditation and Registration

- Wits Health Consortium is a SACRA registered GCP training provider
- The course is HPCSA accredited — 10 CPD points

Target Audience

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.

Registration Fee

R2 700 per person (including VAT)

Course Duration

Day 1: 08h30-15h30
Day 2: 08h30-12h30
(Start and end times may vary slightly, confirmation will be provided on registration)

Venue

These courses are usually held at the Wits Health Consortium (Pty) Ltd Parktown offices. Special requests will be considered, subject to viability.

Registration

Please complete the course registration form, e-mail to **training@academicadvance.co.za**

Contact Details

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Further Information

For further information, please go to our website:

http://www.academicadvance.co.za