ABC OF QUALITY CONTROL IN CLINICAL RESEARCH

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

- The course is HPCSA accredited — 10 CPD points

Booking

Please contact us at.

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http://www.academicadvance.co.za

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.

Course Duration

1 ½ days

(Start and end times may vary slightly, confirmation will be provided on registration)

Course Fee

R 3,000.00 incl VAT (Non-WHC)

R 2,150.00 ex VAT (WHC Divisions/Syndicates)

Venue

BEESA Conference Centre,

Unit 3, Sherborne Square,

5 Sherborne Rd, Parktown, 2193

Courses are also offered in Durban on a regular basis and in Cape Town subject to a minimum number of attendees.

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