ABC OF QUALITY CONTROL IN CLINICAL RESEARCH

VENUE

Ground Floor Training Room
31 Princess of Wales Terrace
Parktown, Johannesburg, 2193

Courses are also offered in Durban on a regular basis.

On-site training for groups of 20 or more will be considered , subject to viability.

COURSE FEE

R 3,200 incl VAT (Non -WHC)
R 2,300 excl VAT (WHC Divisions)

COURSE DURATION

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

BOOKING

Please contact us at:
+27 11 274 9256/9327/9200
training@academicadvance.co.za
http://www.academicadvance.co.za

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 1 ½ 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.

TARGET AUDIANCE

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.

ACCREDITATION AND REGISTRATION

The course is HPCSA accredited- 8 CPD points and 2 Ethics points

COURSE CONTENT

- Site Organisation
- Study preparation
- Study Conduct
- Study Close Out R Archiving
- Audits & Inspections

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