



R 1,500 incl VAT (Non-WHC) R 975 excl VAT (WHC)





Classroom based (available online from March 2021)



4 Hours

Medical research involving human subjects is strictly controlled by the South African Health Products Regulatory Authority (SAHPRA); DOH and Ethics Committees. It is mandated that the research team be well versed with the International (ICH) and SA Good Clinical Practice guidelines. Each member of the team must be in possession of a valid, accredited GCP certificate.

In addition to the clinical personnel, a research team may include a number of support staff who recruit participants, take informed consents, counsel participants, transport participants, carry out laboratory procedures, capture data, etc. It is imperative that this category of staff understands the concept of research on human subjects, the guidelines that govern the conduct of this research, the roles of the study team and the importance of their role in the greater team.

The course has 4 modules including a "Test your knowledge" at the end of each module:

- Introduction
- Module 1: The development of Good Clinical Practice (GCP)
- Module 2: GCP in South Africa



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

Module 4: Topical Issues in Clinical Research

There are 50 multiple choice questions.

Module 3: Audits and Inspections

- 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.



This course is a revision of Good Clinical Practice principles and processes for all non-clinical support staff who have previously attended GCP training. The course is interactive and discussion around current issues is encouraged.



All non-clinical support staff, such as counsellors, fieldworkers, recruiters, data capturers, study administrators, drivers, etc, who have previously.



Parktown, Johannesburg

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



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