

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.

The course has 5 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



This course is a revision of Good Clinical Practice principles and processes for all clinical trial personnel who have previously attended GCP training, incorporating current ICH and SA GCP Guidelines.



All clinical research personnel, who have previously completed a recognised GCP course: Investigators, Study Coordinators, Managers etc.

- Module 3: Audits and Inspections
- Module 4: Genetic Research Trends
- Module 5: Litigation in Clinical Trials



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





+27 11 274 9327/9256/9368