



BASIC ICH GOOD CLINICAL PRACTICE (GCP)

The ICH Good Clinical Practice (GcP) training is important to develop the necessary skills and competencies to perform Clinical Trials according to the regulations and guidelines. The ICH GCP course will cover an introduction, ethics in clinical trials, PI responsibilities, investigational product management and overview of the practical implementation of GCP in clinical trials to ensure compliance with regulatory and ethics committee requirements.

This course is open to individuals who work in clinical research and need to understand the ICH GcP requirements so that they comply with regulations.

Duration: 1 ½ days.

Date: 24 - 25 July 2023

Venue: Video Conferencing Room, UNAM Library, Main Campus, Windhoek

BASIC ICH GOOD CLINICAL PRACTICE (GCP)

This training is designed to provide practical understanding of the principles and application of GCLP in laboratory and trials settings. The training course will cover the elements in GCP and GCLP required for compliance, including the common GCLP audit findings.

This course is targeted at lab personnel involved in clinical trial sample handling, processing, and sample.

Duration: 1 ½ days.

Date: 25 to 26 July 2023

Venue: Video Conferencing Room, UNAM Library, Main Campus, Windhoek

Please note that course material will be provided. A certificate of attendance will be issued once a minimum of 80% is achieved in the final quiz. CPD points can be claimed. Both courses will be conducted in English.

The training is at no cost to the participants.

Please email details of participants for either course to: rabotse@ltclinicalresearch.co.za / snghoshi@unam.na by no later than 17 July 2023. In your submission, clearly indicate which course you will be attending.



Related enquiries contact: Dr J Sheehama jsheehama@unam.na / Mr. Sibusiso sibu@ltclinicalresearch.co.za