
02. Pre-Clinical Research, Phases of Clinical Trials.

03. International Conference on Harmonization and good Clinical Practices guidelines (ICH-GCP guidelines)

04. Ethics in Clinical Research

05. Review and Approval Processes of Investigational New Drug/New Drug Application

06. Study Design & Essential Documents

07. Clinical Site Selection,

08. CRA responsibilities & activities

09. Safety Monitoring and QA audit

10. Pharmacovigilance

11. Protocol writing

12. Basics of Epidemiology

13. Basics of Biostatistics