Course Structure

Certificate Course in Clinical Trials and Contract Research

Course-Paper-I: Introduction to Clinical Trials

Course-Paper-II: ICH, Ethics, IRB and Regulations for Clinical Trials

Course-Paper-III: Contract Research in Clinical Trials

Course-Paper-IV: Biostatistics in Clinical Trials

Certificate Course in Medical and Scientific Documentation Writing

Course-Paper-I: Basics of Good Medical Writing

Course-Paper-II: Scientific Writing: Concepts and Applications

Course-Paper-III: Protocol Writing for Clinical Trials

Course-Paper-IV: Case Report and Publication Writing

Certificate Course in Clinical Trials Data Management

Course-Paper-I: Introduction to Clinical Trials Data Management

Course-Paper-II: Biostatistics, SAS and Data Management

Course-Paper-III: Computerized Systems for E- Case Report Form and Data Acquisition and Validation

Course-Paper-IV: ICH, Ethics, IRB and Regulations for Clinical Trials

Comprehensive Program in Clinical Trials, Regulations and Data Management

Course-Paper-I: Introduction to Clinical Trials

Course-Paper-II: ICH, Ethics, IRB and Regulations for Clinical Trials

Course-Paper-III: Contract Research in Clinical Trials

Course-Paper-IV: Biostatistics in Clinical Trials

Course-Paper-V: Introduction to Clinical Trials Data Management

Course-Paper-VI: Computerized Systems for E- Case Report Form and Data Acquisition and Validation

Course-Paper-VII: Bioavailability and Bioequivalence Studies

Course-Paper-VIII: Case Study Report

Professional Designation in Clinical Trials and Ethics

Course-Paper-I: Introduction to Clinical Trials

Course-Paper-II: ICH, Ethics, IRB and Regulations for Clinical Trials