

Certificate Program for Clinical Trial Investigators & Site Personnel

This is a comprehensive educational program, designed for the Clinical Investigators & Site Personnel aspiring to work in the field of clinical research or allied professions like Investigator Site & Ethics Committee. The program provides complete overview and practical environment in the field of clinical research. The program would candidates to upgrade their knowledge about ICH GCP Guidelines, regulatory issues and other major aspects of clinical trials management. Program would help Clinical Investigators & Site Personnel in developing career in the field of clinical research.

Program Details : The program would cover

- Introduction to Clinical Research & Phases of Clinical Trials
- Introduction to ICH GCP / Schedule Y
- Ethical Considerations in Clinical Research
- Investigator Roles & Responsibilities
- "Informed Consent Process & Documentation"
- Safety Reporting
- Study Drug Accountability
- Site Selection Process and Pre-Study Visits
- Routine Site Monitoring Visits
- Site Audits
- Site Close Out

Mode : Online/Distance Learning

Duration : 6 Months

Eligibility : MD, MS, MBBS, BDS, BHMS, BAMS, BUMS, BPT, B.Pharms, Graduate/Post Graduate Degree in Life Sciences, Mathematics, Pharmacology, Pharmacy, Medical Laboratory, Nursing, Biochemistry, Microbiology, Biotechnology and all professionals working with Pharmaceutical companies, CROs and Hospitals.

Methodology : Printed Training Modules; Online e-learning System

Examination : Online MCQs

Certificate : Certificate would be awarded upon successful completion of the program. Program is Certified & Accredited by the **Pharmaceutical Society of India**.

Accreditation : Accreditation would be awarded upon successful completion of the program. Program is Certified & Accredited by **Accreditation Council for Clinical Research Education, US**.

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Disclaimer - This course brochure is for the purpose of creating an awareness about the program and career options. The exact information - This course structure would be given to you at the time of orientation, and may vary from this brochure.

- Job Assistance** : Placement support would be provided to the successful candidates. CVs of successful candidates would be forwarded to relevant organization.
- Fee payment** : Fee Payable by Cash, Cheque/Bank draft in the name of “**TENET HEALTH EDUTECH PVT LTD.**” payable at Delhi. Fee can also be deposited in company bank account. We also accept Credit/Debit Cards.
- International Payments** : Through Debit/Credit cards using Paypal or wire payment through banks
- Course Objectives** : • This program consists of modules on topics related to Good Clinical Practice (GCP), applicable to clinical research. This course is intended to provide the necessary training for investigators and other senior site personnel, to better understand the regulations regarding clinical research in human subjects, and a better understanding of roles and responsibilities in planning and conducting clinical trials. The program also covers the regulatory aspects as well