

Diploma in Clinical Research

- 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 the **Swiss Association of Pharmaceutical Professionals (SwAPP) & Swiss Society of Pharmaceutical Medicine**.
- Job Assistance** : Placement support would be provided to the successful candidates. CVs of successful candidates would be forwarded to CROs Hospitals and Pharmaceutical companies and, Interviews would be organized.
- 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
- Program Details** :
This is a foundation educational program, designed for all the candidates aspiring to work in the field of clinical research or allied professions like central labs. The program provides complete overview and practical environment in the field of clinical research. The program would allow candidates to upgrade their knowledge about ICH GCP Guidelines, regulatory issues and other major aspects of clinical trials management. Select important topics from the following groups would be covered in the program:

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India : Delhi | Ahmedabad | Bangalore | Bhopal | Chennai | Coimbatore | Cochine-Trivandram | Hyderabad | Jammu | Kolkata | Lucknow | Mumbai | Pune | Vijayawada
International : US | UK

- Clinical Research Introduction
- Roles & Responsibilities of Key Stakeholders
- Preparations & Planning for Clinical Trials
- Essential Documentation in Clinical Research & Regulatory Submissions
- Clinical Trials Project Planning & Management
- Study Start Up Process
- Clinical Monitoring Essentials
- Compliance, Auditing & Quality Control in Clinical Research

Course Objective :

- ❖ To provide a comprehensive introduction to the clinical research process.
- ❖ Learn the skills, knowledge and competencies of a candidate for the Clinical Research Associate / Clinical Research Coordinator position.
- ❖ Become more familiar with roles/jobs as part of the study team.