



Post Graduate Diploma in Clinical Trials Management

This is an extensive educational program, designed for the Pharma professionals/Life Sciences/Medical Practitioners aspiring to work in the field of clinical research or allied professions like central labs, CROs, Sponsor Company & Pharmacovigilance. 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 Pharma professionals/Life Sciences/Medical Practitioners in developing career in the field of clinical research.

Program Details : The program would cover

	Clinical Research Introduction & Terminology Healthcare Management Issues Introduction to Pharmaceutical Industry & Global Challenges Global Clinical Research Environment & Opportunities Regulatory Affairs Principles of Pharmacology & Drug Discovery & Development 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 Clinical Data Management, Biostatistics, Analysis & Reporting
• • •	Clinical Data Management, Biostatistics, Analysis & Reporting Pharmacovigilance Bioavailability and Bioequivalence Studies Management of Cancer Clinical Trials Organizational Behavior & Human Resource Management
: Onli	Financial Management ne/Distance Learning Ionths
: MD Grae Lab	, MS, MBBS, BDS, BHMS, BAMS, BUMS, BPT, B.Pharms, Graduate/Post duate Degree in Life Sciences, Mathematics, Pharmacology, Pharmacy, Medical oratory, Nursing, Biochemistry, Microbiology, Biotechnology and all professionals king with Pharmaceutical companies, CROs and Hospitals.
: Prin	ted Training Modules; Online e-learning System

Examination : Online MCOs

Mode

Duration

Eligibility

Methodology

C-101 First Floor, Sector-2, Noida - 201301 (U.P), India

 Mobile: +91-9311172560, +91-9810068241 Email: info@cliniminds.com
 Website: www.cliniminds.com

 India: Delhi - NCR | Ahmadabad | Bangalore| Bhopal | Chandigarh | Chennai |Coimbatore | Hyderabad | Kolkata | Mumbai | Trivandrum International: US | UK

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





Assuring and Advancing Quality in Clinical Research Education since 1965

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.
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	 To provide a comprehensive introduction to the clinical research process, conduct & management of clinical trials. Learn the skills, knowledge and competencies of a candidate for the Clinical Research Associate and other key positions. Become more familiar with roles/jobs as part of the study team. Extensive Knowledge & application in different aspects of Clinical Research.

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