cliniminds™
Institute of Health Sciences Training & Management
An ISO 9001:2008 Certified Academy

Think i

ACCRE
Assuring and Advancing Quality in Clinical Research Education since 1965

# Advanced Post Graduate Diploma in Clinical Trials Management

## Our Industry Training Partners – Thinki and CRQA

### Hands on Software Training

### Program Accredited by Accreditation Council for Clinical Research Education, USA

**100% Placement Record | Industry Accredited Program | Specialization | Scholarship | Internship with Leading Companies | Technical as well as Management Training | Expert Faculty | Hi- Tech Classroom**

## Specialization in Clinical Research, Pharmacovigilance, Data Management, SAS, Medical and Scientific Content Writing, Drug Regulatory Affairs

Clinical Research, Pharmacovigilance, Data Management, Regulatory and Medical Writing sector is rapidly growing industry globally with growth rate of over 20% per annum. Over 50,000 professionals would be required in the next 2 – 3 years in India alone. Global industry size is over US$50 billion, and in India industry is expected to touch the turnover of US$2 - 3 billion in the next 3 years. Other key global markets like China, South East Asia, Europe and Americas continue to grow in double digit. MNC companies are outsourcing businesses like Pharmacovigilance, data management, medical writing and SAS from India. Thousands of new jobs are being created in Clinical Data Management; Pharmacovigilance; Medical Writing and other areas of life sciences.

This growth is opening up massive opportunity for the medical, pharmacy and life sciences graduates and post graduates. Sector requires highly specialized and skilled professional workforce, with applied clinical research, pharmacovigilance, regulatory, medical writing, quality assurance and data management skills. Cliniminds has been at the forefront of providing clinical research training and consulting solutions to the life sciences industry for the last several years.

Cliniminds flagship program, Post Graduate Diploma in Clinical Trials Management program is in existence for the last 7 years and over 25 batches have passed and have been placed in the pharmaceutical companies, CROs and other clinical research organizations. Our students have been placed with the leading companies. We have already trained over 2,000 professionals and successfully placed them in the industry.

**Program Structure**

- 5 months Intensive Full time Classroom Training; 6 months Internship
- Add on Specialization in Pharmacovigilance, Clinical Data Management, Clinical Operations, Drug Regulatory affairs, Medical Writing

**The program would cover:**

- Clinical Research
- Pharmacovigilance
- Clinical Data Management
- Statistical Analysis System (SAS)
- Drug Regulatory Affairs
- Medical and Scientific Content Writing
- **English Communication & Personality Development**

C-101 First Floor, Sector-2, Noida - 201301 (U.P), India
**Mobile:** +91-9910068241, +91-9810068241, +91-9311172560 **Email:** info@cliniminds.com **Website:** www.cliniminds.com
**India: Delhi - NCR | Gurgaon | Bangalore| Bhopal | Chennai | Kolkata | Mumbai | Trivandrum | Varanasi | Visakhapatnam International: Russia | Saudi Arabia | US |UK**

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**In following area Students would be eligible to work after completion of the Program:-**

- Clinical Research Operations
- Pharmacovigilance
- Clinical Data Management
- Medical & Scientific Writing
- Regulatory Affairs

**Advantages of Cliniminds Program**

- Industry Accredited / Certified
- Application of Clinical Research & Pharmacovigilance in real business like environment.
- Completely Job Oriented – Hands-on Training
- Internship Certificate will be provided by leading CRO/SMO/Hospital
- Accredited by the by Accreditation Council for Clinical Research Education, USA (ACCRE) & Certified by Pharmaceutical Society of India
- 100% placement – Excellent Placement Record
- Training by the team of industry experts – both full time and visiting senior faculty
- Small batch – 15 Seats

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| **Faculty** | **:** | Training would be imparted by the full time cliniminds faculty & visiting experts from the industry. |
| **Mode** | **:** | Classroom/Online/Webinar |
| **Duration** | **:** | 11 Months |
| **Eligibility** | **:** | MD, MS, MBBS, BDS, BHMS, BAMS, BUMS, BPT, B.Pharm, 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** | **:** | 5 Months classroom and 6 months internship, Hands on training, online access to study materials, Printed study materials and Workshops. |
| **Examination** | **:** | Classroom/online exams & Project work |
| **Certificate** | **:** | Certificate would be awarded upon successful completion of the program. Program is certified by the Pharmaceutical Society of India. |
| **Accreditation** | **:** | Program is accredited by **Accreditation Council for Clinical Research Education**, **USA.** |
| **Job Assistance** | **:** | Extensive Placement support would be provided to the successful candidates. 100% Past Placement Track Record. |

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**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 the company bank account. We also accept Credit/ Debit Cards.

**International Payments :** Through Debit/Credit cards using Paypal or wire payment through banks.

**Course Objectives:**

- Extensive applied / practical knowledge imparted to equip you to work at any global clinical research organization or pharmaceutical company.
- Learn the skills, knowledge and competencies of a candidate for the Clinical Research Associate and Pharmacovigilance jobs.
- Practical aspects of important Pharmacovigilance activities as per the global standards like medical evaluation, casualty assessment, expectedness assessment, case narratives, MedDRA, case processing preparation of safety report etc.
- Learning detailed functions of clinical data management, regulatory affairs and medical and scientific writing.
- Balance of academic and job orientation.