

JOIN



FAST TRACK

ADVANCED POST GRADUATE PROGRAM IN CLINICAL RESEARCH & PHARMACOVIGILANCE

**FAST TRACK INTENSIVE PROGRAM – 6 WEEKS AND 2 WEEKS PROJECT WORK
100% PLACEMENT PLAN: INTERNSHIPS : INDUSTRY CERTIFICATION**

We understand your need for the educational program which is of shorter duration yet effective and job oriented, and provides with you with the excellent job opportunity at the same time. In today's competitive world the time is only most important resource.

The program has been designed in a way that you would learn Clinical research and Pharmacovigilance in a shorter span, equivalent to what M.Sc. or a Post Graduate Diploma student would learn in 1 – 2 years.

Strengths of our Program:

- Shorter Duration
- Intensive Program – Full Day – 6 Weeks and 2 Weeks Project Work
- Mentoring by Medical Doctors, Clinical Research & Pharmacovigilance Experts
- Skill Development, Placement & Career Development Focus
- Job Ready
- Favourable Cost Benefit
- Limited Seats

Clinical research is rapidly growing industry globally with growth rate of over 40% per annum. Sector requires highly specialized and skilled professional workforce, with applied Clinical research & Pharmacovigilance. Over 50,000 professionals would be required in the next 2 – 3 years. Global industry size is over US\$40 billion, and in India industry is expected to touch the turnover of US\$2 billion in the next two years. Other key global markets like China, South East Asia, Europe and Americas continue to grow in double digit.

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.

Regulatory bodies such as the USFDA and EMEA are intensifying safety regulations, thereby boosting the adoption rates of Pharmacovigilance systems by pharmaceutical companies. Several countries, including India are in the process of implementing stringent regulations for adoption of Pharmacovigilance. There is significant potential for outsourcing/off shoring for mid-sized companies as well. Large number of global drug companies have starting off shoring their Pharmacovigilance activities to the markets like India. In a typical Pharmacovigilance department or specialized Pharmacovigilance company, there are several positions. Pharmacovigilance offers excellent growth prospects. Some of the positions are Drug Safety Associate; Drug Safe Scientist; Aggregate Report Scientist; Team Leaders.

- ✎ **Cliniminds** has been at the forefront of providing clinical research training and consulting solutions to the life sciences industry for the last several years. We have already trained over 2,000 professionals and successfully placed them in the industry.
- ✎ **Cliniminds** flagship program, Post Graduate Diploma in Clinical Research & Pharmacovigilance, is in existence for the last 6 years and over 18 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.

Program Structure:

Six weeks comprehensive practical knowledge to the Clinical research processes, Drug development process, Regulatory affairs, Essential documentation, Roles and Responsibilities, Ethics, Monitoring, Conduct and Management of trials, Pharmacovigilance, Data management and various other related issues in intensive Full time Classroom Training.

Certificate :

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

Accreditation :

Program is Certified & Accredited by the Swiss Association of Pharmaceutical Professionals (SwAPP) & Swiss Society of Pharmaceutical Medicines.

Internship option with various pharmaceutical companies, CROs, and hospital research sites.

Entry Level Career Options Upon Completion of the Program:

- ☞ Clinical Research Operations
- ☞ Clinical Research Co-coordinator
- ☞ Clinical Trial Assistant.
- ☞ In-house CRA
- ☞ Pharmacovigilance Officer
- ☞ Drug Safety Associate
- ☞ Drug Safety Officer
- ☞ Data Management Professional

Program Details :

- Clinical Research Introduction
- 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
- Overview of Clinical Data Management & Biostatistics
- Clinical Data Management Software Training
- General Overview of Pharmacovigilance
- Medical Dictionary for Drug Regulatory Activities MedDRA
- Regulatory Aspects in Pharmacovigilance
- Diagnosis And Management of Adverse Drug Reactions
- Medical Evaluation of Adverse Events
- Quality System in Pharmacovigilance
- Periodic Safety Update Reports (PSURs) For Marketed Drugs (ICH E2C)
- Pharmacovigilance Database And Signal Detection Tools Risk Assessment, Evaluation And Management
- EudraVigilance
- Suspected Unexpected Serious Adverse Reaction (SUSAR) Reporting
- Expedited Reporting Requirements
- Pharmacovigilance Software Overview

Mode : Classroom/Online/Webinar

Duration : 6 Weeks and 2 Weeks Project Work

Eligibility:

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

Methodology :

Full time class Room + Hands on training, Online access to study materials, Printed study materials and Workshops.

Examination : Classroom/online exams & Project work

Job Assistance :

Extensive Placement support would be provided to the successful candidates.

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 Payment :

Through Debit/Credit cards using Paypal or wire payment through banks.

For more information and admission, please contact admissions@cliniminds.com

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