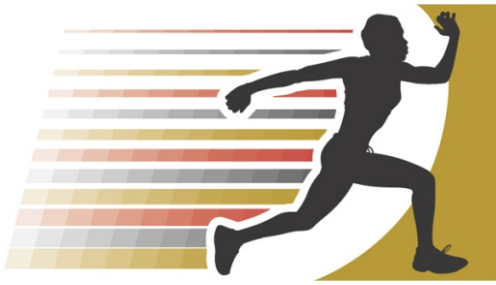
cliniminds™
Academy for Clinical Research and Healthcare Training & Management

An ISO 9001:2008 Certified Academy

MEDRIO

FAST TRACK



JOIN

Now put your career on Fastrack with Cliniminds 3 months Clinical Research & Pharmacovigilance Program

Why to spend time and money on 1 or 2 years programs, when your objective is to get the right placement?

20th Batch of Clinical Research &Pharmacovigilance Program

100% Placement of Last Batch

ADVANCED POST GRADUATE DIPLOMA IN CLINICAL RESEARCH & PHARMACOVIGILANCE

FAST TRACK INTENSIVE PROGRAM –3 MONTHS (INCLUDING PROJECT WORK) : 100% PLACEMENT PLAN: INTERNSHIPS: INDUSTRY CERTIFICATION: ACCOMMODATION INCLUSIVE

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 & 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
- Case study based practical training approach
- Emphasis on Personality Development & Communication Skills
- Intensive Program – Full Day – 3 Months (Including Project Work)
- Mentoring by Medical Doctors, Clinical Research, Pharmacovigilance, Data Management Experts, IT Professionals
- Skill Development, Placement & Career Development Focus
- 100% Placement Support Upon Successful Completion of Program
- 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.

GET AHEAD OF THE REST IN THIS RAPIDLY GROWING CLINICAL RESEARCH INDUSTRY.....

Program Structure:

- Total duration of the program is 3 months.
- Classroom training : 3 Months (Including Project Work).

Last Date of Registration: - 20th April 2012

Class Commencement Date : Sunday, 29th April 2012

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.

Advantages of Cliniminds Program :

- Industry Accredited / Certified - Completely Job Oriented
- Accredited by the Swiss Pharmaceutical Professionals Association (SwAPP) & Pharmaceutical Society of India
- 100% placement – Excellent Placement Record
- Dual Certification
- Program Superior than M.Sc. and MBA in Clinical
- Training by the team of industry experts – both full time and visiting senior faculty
- Small batch

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

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 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.
- Balance of academic and job orientation.

Mode : Classroom & Webinar

Duration : 3 Months (Including 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.