

## Post Graduate Diploma in Pharmacovigilance

**100% Placement | Three Months Internship | Industry Accredited Program | Specialization | Job Oriented**

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.

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 has 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 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

- 6 Months extensive training on Pharmacovigilance (PV) is provided.
- 3 Months Internship in leading Hospitals, CROs, SMOs, Pharma Companies.
- Online Internship with leading US Company.

### Entry Level Career Options Upon Completion of the Program:

- Phramacovigilance Officer
- Drug Safety Associate
- Drug Safety Officer

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

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India : Delhi | Bangalore| Bhopal | Chandigarh | Chennai |Coimbatore | Hyderabad | Jammu | Kolkata | Mumbai| Pune |Surat | Thrissur | Trivandrum |Vijayawada  
International : US | UK

### 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
- Training by the team of industry experts – both full time and visiting senior faculty
- Small batch – 10 Seats

**Mode** : Classroom/Online/Webinar

**Duration** : 6 months

**Eligibility** : MD, MS, MBBS, BDS, BHMS, BAMS, BUMS, BPT, B.Pharm, Graduate/ Nursing, Biochemistry, and all professionals working with Pharmaceutical companies, CROs and Hospitals.

**Methodology** : 6 Months Weekend class Room + 3 Months Paid Internships, 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 & 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 Medicines.

**Job Assistance** : Extensive 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

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International : US | UK

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

**Program Details:**

- Introduction to Clinical Research & Phases in Clinical Research
- Principles of Good Clinical Practices - ICH GCP
- 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
- Expedited Reporting Requirements
- Periodic Safety Update Reports (PSUR,s) For Marketed Drugs (ICH E2C)
- Pharmacovigilance Database And Signal Detection Tools
- Risk Assessment, Evaluation And Management

**Course Objective :**

- Candidate should able to understand the basic concepts, importance of Pharmacovigilance and Global Pharmacovigilance regulations.
- 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.

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