

POST GRADUATE CERTIFICATE PROGRAM IN DRUG SAFETY & MEDICAL REVIEW FOR PHYSICIANS

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Pharmacovigilance has grown significantly in recent years and its importance in the healthcare system has been recognized worldwide. There are considerable issues which need to be addressed to ensure the safety of medicines and medical devices.

Medical professionals play an important role in the global pharmacovigilance system. They require substantial understanding and expertise in the field of drug and device safety which will successfully contribute to this area through early recognition, management, and reporting of the drug and device safety issues. It is important that medical professionals should be formally trained in the pharmacovigilance. They should possess a combination of training and skills in this area.

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 and off shoring for every pharmaceutical and medical device company. Large numbers of global drug and device 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 entry level positions for medical doctors, viz. Drug Safety Physician, Medical Reviewer, and Safety Scientist, and this leads to higher positions over the years at fast pace.

For medical professionals pharmacovigilance career offers

- Career in Medical Science
- Substantial career growth
- Attractive salaries
- 5 days a week job
- International training and job opportunities
- Fixed holidays and annual vacations

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

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

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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.

Cliniminds Program covers the entire spectrum of Drug Safety Physician / Medical Reviewer Role. Following are the key topics

1. Introduction to Pharmacovigilance
2. Global Pharmacovigilance Regulations
3. Reportability Criteria & Case reports
4. Safety reporting and Processing ICSR Medical Evaluation of ICSR
5. Aggregate Report Writing
6. Signal Detection & Risk Management

Eligibility	MBBS, MD, MS, DNB
Mode	Online E learning
Duration	03 Months
Methodology	Online, eLearning Webinars
Placement Guidance	CV Preparation, Mock Interviews and sending CVs to Recruiters
Examination	Online MCQs
Certificate	Accredited from Accreditation Council for Clinical Research Education, USA
Payment	Fee payable using credit / debit card via paypal

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