










Certificate Program in Pharmacovigilance

About This Course

With the globalization of the pharmaceutical industry, the need to maintain universal standards as well as the demand for qualified and trained professionals to provide valuable inputs on Indian, European Union and American Regulatory, Research and Safety requirements in Pharmacovigilance has propelled the need for a trained and certified manpower in this industry.

This self-paced certificate program will provide you a comprehensive training on scientific, practical, ethical and technical concepts on drug safety and management. CPPV is a 6 months course and covers all the pharmacovigilance concepts and its applications in 11 modules covering 250+ topics. Each module is well explained in detail with the help of illustrations, examples, and flowcharts.

Course Highlights

-  Mode of Learning: Online/ Distance
-  Course Timings: Self-paced
-  Course Duration: 6 Months
-  Total Efforts: 60 Hours Approximately
-  Assessment: Online Single Exam at the End of the Course
-  Award: Certificate of Completion and a Grade Card
-  Course Access: Lifetime

Please visit our website physislearningacademy.com for information regarding course fee and current batch

Learning Objectives

At the end of the course, you should be able to learn about the:

- ⑤ Concepts and processes involved in pharmacovigilance
- ⑤ Global guidelines governing pharmacovigilance
- ⑤ Global reporting forms and their procedure to report
- ⑤ Hands-on-training through case studies
- ⑤ Knowledge & skills required to start career in pharmacovigilance industry

Course Curriculum

- ▶ Module 1: Pharmacovigilance Terminology
- ▶ Module 2: Introduction to Pharmacovigilance
- ▶ Module 3: Global Perspective of PV and ADR Reporting
- ▶ Module 4: Guidelines & Standards Governing PV
- ▶ Module 5: Global Adverse Events Reporting System & Reporting Forms
- ▶ Module 6: Individual Case Safety Reports (ICSRs)
- ▶ Module 7: Public Safety Update Reports (PSURs)
- ▶ Module 8: Signal Detection
- ▶ Module 9: Medical Dictionary for Drug Regulatory Activities (MedDRA)
- ▶ Module 10: Expedited Reporting and Requirements
- ▶ Module 11: Pharmacovigilance Inspections
- ▶ Assessment and Evaluation through Exam

Who Should Enroll?

- ③ Health-science students/ working professionals (Pharmacy, Life-Science, Medicine, AYUSH, Dental, Nursing, Physiotherapy, Ph.D. etc.) who are looking to gain knowledge & start their career in the pharmacovigilance industry.
- ③ Fresher, newly appointed and experienced pharmacovigilance personnel.

Testimonials

4.6

Average Rating



Great course on Pharmacovigilance and easy methodology.

Bina Mathur

All course modules are very easy to understand & the team was really supportive.

Deepika Rawat

It helped me in understanding the basic understanding along with some advance concepts of Pharmacovigilance with the help of case studies.

Manoj Mishra

Excellent course and covers all the concepts of Pharmacovigilance.

Ashish Gupta

How This Works?



Upon enrollment, the course participants will get the login details or course material via post before the start of batch via email.



Upon receipt of course content, the participant can read or log in to take the course modules anytime and from any device.



After reading all the course modules, the course participant is required to attempt and submit an exam.



Upon evaluation of exam, the certificate of completion will be issued to all the successful participants of the batch.

Contact Us

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