










Certificate Program in GCP & Clinical Research

About This Course

Clinical Research industry offers a number of options for health science students & working professionals to excel their career as Clinical Research Coordinator, Trial Coordinator, Research Assistance, Research Nurse, Research Scientist etc. The average salary for a fresher level position of Clinical Research Coordinator is estimated to be worth approximately \$50000 to \$60000.

This online & self-paced certificate program CPGCPCR will provide you a comprehensive training on scientific, practical, ethical and technical concepts of the clinical trial along with the application of GCP guidelines. This 10 weeks course covers all the clinical research concepts and its applications in 16 modules covering 250+ topics. Each module is well explained in detail with the help of illustrations, templates, examples and flowcharts.

Course Highlights

-  Mode of Learning: Online and from any Device
-  Course Timings: Self-paced
-  Course Duration: 10 Weeks
-  Total Efforts: 60 Hours Approximately
-  Assessment: Online Single Exam at the End of Course
-  Award: Certificate of Completion and a Grade Card
-  Course Access: Lifetime

Please visit our website physisglobalacademy.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:

- ① Concepts and processes involved in clinical research
- ① GCP guidelines and its application in clinical trials
- ① Roles & responsibilities of clinical trial personnel and stakeholders
- ① Maintenance and management of clinical research documents
- ① Process of safety reporting and quality assurance

Course Curriculum

- ▶ Module 1: Introduction to Clinical Research
- ▶ Module 2: Evolution of Ethics, Regulations & ICH-GCP
- ▶ Module 3: GCP and Code of Federal Regulations (CFR) Guidelines
- ▶ Module 4: Roles and Responsibilities of CR Personnel
- ▶ Module 5: Clinical Study Process
- ▶ Module 6: Institutional Review Board (IRB)
- ▶ Module 7: Informed Consent Process
- ▶ Module 8: Study Subject Recruitment & Retention
- ▶ Module 9: Safety Reporting & Adverse Event Management
- ▶ Module 10: Monitoring and Quality Assurance
- ▶ Module 11: Research Protocol
- ▶ Module 12: Essential Research Documents
- ▶ Module 13: Maintenance of Source Documents
- ▶ Module 14: IND/ NDA/ ANDA/ AADA
- ▶ Module 15: Site Closure
- ▶ Module 16: Research Misconduct

Who Should Enroll?

- ① Medical Researchers & health science professionals who are looking to gain knowledge & start their career in clinical research.
- ① Fresher, newly appointed and experienced clinical research personnel (CRC, CRA, Project Manager, Research Nurse, Trial Assistant, Investigator, QA Personnel etc.)

Testimonials

4.6

Average Rating



The course covers in-depth knowledge on subject matter. It was fun learning this course.

Amy Moore, Trial Assistant

I really found this course very interesting. This really increased my wealth of knowledge in the clinical research domain.

McKay Michelle, Researcher

It was well put together. Great job! Keep it up.

Robert Williams, CRA

Contains quality information for clinical researchers.

Xiao Yang, PhD Researcher

How This Works?



Upon enrollment, the course participant will get the course login details before the start of batch via email.



Upon receipt of course login details, the participant can 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 online 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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