Certificate Program for
Clinical Research Associate (CRA)

PHYSIS
GLOBAL ACADEMY

About This Course

A Clinical Research Associate (also known as Clinical Trial Monitor) is primarily responsible to monitor, administrate and supervise the overall progress of a trial. Effective clinical trial monitoring is essential to ensure smooth conduct and progress of clinical trial activities. This ultimately helps to generate quality research data for analysis and trial result.

CPCRA is an online & self-paced certificate program which will provide you a comprehensive training on scientific, financial, practical, ethical and technical concepts of clinical trial monitoring. This 10 weeks training course covers an in-depth training for the role of clinical research associate (CRA) in 15 modules covering 150+ topics and subtopics. Each module is well explained in detail with the help of illustrations, mock training, templates, examples, and flowcharts.

Course Highights

- 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 the 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 and develop the required skill set to effectively monitor and administer clinical trial sites
- Identify and define all the procedure to successfully plan, execute, monitor and control the activities of a trial
- Independently plan and execute global monitoring visits
- Learn the process of safety reporting requirements
- Identify, manage and report the trial deviations
- Effectively manage pre, during and after trial job functions in a time-bound manner

Course Curriculum

- Module 1: Introduction to Clinical Research
- Module 2: Evolution of Ethics, Regulations & Guidelines
- Module 3: Essential Clinical Trial Documents
- Module 4: Investigator Site Selection & Assessment
- Module 5: Development of Monitoring Plan
- Module 6: Site Initiation, Documents Review & Delegation of Duties
- Module 7: Clinical Trial Monitoring
- Module 8: Inventory Planning & Tracking
- Module 9: Source Document Verification (SDV)
- Module 10: CRF Review, Collection & Coordination of Data Mgmt.
- Module 11: SAE Review & Regulatory Compliance
- Module 12: IP Accountability & Management
- Module 13: Escalation, Prevention & Management of Violation/ Deviations
- Module 14: Tracking of Enrolments, Payments & Ongoing Correspondence
- Module 15: Site Closure

Who Should Enroll?

- Clinical Research Coordinators, Research Assistants, and Nurses looking to advance their career as Clinical Research Associate.
- Newly appointed Clinical Research Associate (CRA)
- Experienced CRAs interested in certification

Testimonials

4.4

Average Rating

★★★★½

I loved the mock hands-on-training part of clinical trial monitoring course. It was first of its kind experience.

Michael Dar, Study Coordinator

The course modules were very easy to understand and the information is very well presented.

Dave Shaw, Research Coordinator

Great and very informative for all the researchers looking advance their career in clinical research.

Javier Guetta, Trial Investigator

Covers enough content for beginners & the exam structure was very good.

Sunita Chawla, Research Assistant

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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