










Certificate Program in  
**Clinical Trial Project Management**

# About This Course

Nowadays, efficient and cost-effective projects are the lifeblood of every clinical research organization. Independent of type and phase, meeting project timelines effectively requires special knowledge and skills to manage each aspect of a project. In clinical trials, a project manager is primarily responsible to plan, design, monitor and execute all the project activities with help of project team.

CPCTPM is an online & self-paced certificate program which will provide you a comprehensive training on scientific, financial, practical, ethical & technical concepts of clinical trial project management. This 10 weeks training course covers all the applications and concepts of project management in 13 modules. 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 the Course
-  Award: Certificate of Completion and a Grade Card
-  Course Access: Lifetime

Please visit our website [physisglobalacademy.com](https://physisglobalacademy.com) for information regarding course fee and current batch

# Learning Objectives

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

- ① Develop the required skills to effectively plan and manage a clinical trial
- ② Identify and define all the procedure to successfully plan, execute, monitor and control the activities of a project
- ③ Draft budget strategies, tracking and implementation
- ④ Learn the process of safety reporting requirements
- ⑤ Create an effective communication channel for project team members and stakeholders
- ⑥ Effectively manage the quality of clinical trial activities.

# Course Curriculum

- ▶ Module 1: Introduction to Clinical Research
- ▶ Module 2: Introduction to Trial Project Management
- ▶ Module 3: Regulatory Requirements & Submission
- ▶ Module 4: Project Planning & Forecasting
- ▶ Module 5: Clinical Trial Design
- ▶ Module 6: Trial Budgeting & Financial Planning
- ▶ Module 7: Vendor Selection & Contracting
- ▶ Module 8: Trial Communication & Coordination
- ▶ Module 9: Project Set-up & Documentation
- ▶ Module 10: Patient Recruitment & Retention Strategy
- ▶ Module 11: Project Tracking, Monitoring & Management
- ▶ Module 12: Regulatory Inspections & Audit
- ▶ Module 13: Study Close-out

# Who Should Enroll?

- ④ Clinical researchers looking to develop skills in project management of trials
- ④ Sr. CRAs and aspiring project managers
- ④ Researchers who are recently appointed as managers
- ④ Experienced project managers looking to gain skills & knowledge to provide fundamental skills to execute their job functions efficiently

# Testimonials

## 4.4

Average Rating



All the clinical researchers should take this course to learn the project management of a study. This course will definitely help in making research activities more efficient and compliant.

Elizabeth Alderson, Investigator

The course was very accessible and easy to take at your own time. The person was very good in response to few of my questions.

Iliana Benzakein, Clinical Research Associate

It meets my expectations and covers enough information on all the activities.

Satya Jayprakash, CRC

# 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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