Certificate Program in Medical Writing

PHYSIS
LEARNING ACADEMY

About This Course

As per reports, the Indian drug development outsourced market size at little over Rs 1,500-1800 crores. Medical writing contributes to about three-four percent of this with an estimated growth rate of 15-20 percent. India has a huge amount of experienced and qualified researchers who are equally proficient in the English language. This, in addition, makes India as one of the preferred destinations for medical writing jobs.

This self-paced certificate program will provide you a comprehensive training on procedures and strategies to design different clinical trial documents in compliance with applicable regulatory and GCP guidelines. CPMW is a 6 months course and covers all the medical writing process in 10 modules covering 200+ topics. Each module is well explained in detail with the help of illustrations, examples, and flowcharts.

Course Highights

- 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:

- Process and strategies to design clinical trial documents
- Knowledge and skills required to start career as medical writer
- Applicable guidelines & policies to develop essential trial documents
- Strategies to improve the quality of documents

Course Curriculum

- Module 1: Introduction to Clinical Research
- Module 2: Introduction to Medical Writing
- Module 3: Essential Clinical Trial Documents
- Module 4: Developing Standard Operating Procedures (SOPs)
- Module 5: Developing Clinical Trial Protocol
- Module 6: Developing Informed Consent Document
- Module 7: Developing Case Report Forms (CRFs)
- Module 8: Developing Data Management Plan
- Module 9: Developing of Clinical Study Report (CSR)
- Module 10: Clinical Research Glossary
- Assessment and Evaluation through Exam

Who Should Enroll?

- 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 as full-time or freelance medical writer.
- Fresher, newly appointed and experienced clinical research personnel.

Testimonials

4.5

Average Rating

★★★★½

The inclusion of all the topics on clinical trial medical writing is great.

Abhilasha Kumari

The detailed explanation of processes and strategies discussed are good learning.

Kanchan Seth

All the modules are well presented in order and well explained.

Ravi Gopal

Great course. I would recommend this course, who are willing to take in-depth knowledge on “How to write and design clinical trial documents”?

Ajay Mishra

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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