VALESTA ACADEMY

Share Knowledge. Develop Talent.

Clinical Trial Nurse Workshop

Practical information: **Number of participants: Between 4 and 10**
Duration: 1 day (09:00 – 17:00)
Price per person @ Valesta office: €750

Course Overview

Clinical trials are an important part of advancing patient care, and the coordinators of clinical trials require specialised training. As a Clinical Trial Nurse (CTN), you are the link between the site and subject and between the site and sponsor, so you have a crucial role in the setup and conduct of a study to ensure the high quality of the data.

If you are starting as a CTN and need extra guidance to fully understand your role in clinical research, this course will provide you with knowledge on the CTN tasks so you feel confident to deliver high quality work and data for both the subject and sponsor.

At this workshop we will not only explain the tasks involved, but you will also experience the responsibilities of a CTN during hands-on exercises throughout the day. To enroll in this course, participants are required to have a Good Clinical Practice certification. This course can be combined with the “Introduction to Good Clinical Practice” course provided by Valesta Academy.

By attending this course you will gain:

An understanding of the main responsibilities and tasks of a Clinical Trial Nurse and receive hands-on experience on several tasks. You will have a better understanding in:

- The purpose and the content of the Informed Consent Form
- How to explain an Informed Consent to a patient
- How to complete a Case Report Form according to the source documents
- The purpose and quality standards of the Essential Documents
- Safety reporting in a clinical trial

Course Content

During this interactive and practically oriented training, participants will be engaging in several possible Clinical Trial Nurse tasks and responsibilities.

Topics covered during the training are:

- Drug/device development cycle
- Competent Authority and Ethics Committee submission and approval
- Good Clinical Practice (GCP) implications for the Principal Investigator and his team
- Start-up and conduct of a clinical trial
- Informed consent: content, process and documentation requirements
- Essential documents: purpose and quality
- Case Report Form (CRF) completion
- Safety Reporting process and requirements

This intensive one day course will comprehensively guide you through the tasks and responsibilities of a CTN. The course will also provide theory and interactive exercises and cases, making this training the ideal foundation for a career as a CTN.

For more information or to reserve a seat on this training course please contact us today!

We are flexible! Should you require training to be carried out in-house, we would be happy to discuss your needs and come up with a suitable solution for you.
Call our office on +32 15 28 15 05!