VALESTA ACADEMY

Share Knowledge. Develop Talent.

Clinical Research Associate Workshop

Practical information: **Number of participants: Between 3 and 6**
Duration: 2 days
Price per person @ Valesta office: €1300

Course Overview

As a Clinical Research Associate (CRA), it is important during all site visits you are self-confident, knowledgeable and a trustworthy partner for all stakeholders throughout all stages of a study. A CRA is involved in all stages of a clinical trial, including identifying and setting up an investigational site, monitoring of the trial and then closing it down. The CRA will also act as the main line of communication between Sponsor and Investigator and is the primary contact for the site.

In this course, you will gain insight into basic monitoring tasks, how to communicate and present a study to site staff and act with confidence in your CRA position.

The Clinical Research Associate Workshop has been created for Junior CRAs (those with little or no experience as a CRA) or those that wish to pursue a CRA function in their career. 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 CRA throughout a clinical trial, such as:

- Investigator selection, interaction and evaluation
- Site evaluation and set up
- Presenting trial protocols
- Monitoring the trial throughout its duration
- Close out sites on completion of the trial
- Preparation of monitoring reports

Course Content

This two-day Clinical Research Associates Workshop is both theoretically and practically orientated to ensure that the valuable learning achieved can be put into practice effectively.

Topics that are being covered during this training are:

- Authorisation by Competent Authorities/Ethics Committee submission approval
- Clinical trial setup and conduct
- Preparation and performance of the monitoring visit
- Source Data Verification and Drug Accountability
- Review of a monitoring visit report
- Close out visit

During the workshop, participants will learn and experience the responsibilities and tasks of a CRA by performing the many simulations and exercises. Participants will share knowledge and gain hands-on experience through group activities with active participation and exchange of information being the primary focus of this method.

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!