Suggested Training for Clinical Research Coordinators

This educational resource is designed for both new and experienced Clinical Research Coordinators, and other clinical research staff at UCLA who are seeking training resources, including written material, classes and computer based learning modules.

Clinical Research Coordinators (CRCs) are responsible for the organization, coordination, and overall integrity of a research project with humans. Principal and co-investigators provide the overall direction in a clinical study, but CRCs have significant roles in clinical study activities, including **Clinical Trial Study Start-Up.**

Human Research Protection Training

Human Research Protection Training Note: These modules can take several days to complete.

Good Clinical Practice (GCP) Basics

Good Clinical Practice (GCP) Information: The CITI Good Clinical Practice (GCP) Basic Course is an overview of GCP. Some units on campus require this course, but it is optional for the OHRPP. Be sure to check with your departmental GCP training requirements.

CareConnect

CareConnect training for Clinical Coordinators These modules take about 7 hours according to the APEX training staff.

OnCore Training?

- For system security and compliance, training is required for access to OnCore. Registration for online and in-person trainings is available through the CareConnect Training Management System
- To request an account and view training requirements for the different levels of access, view the research recommended training section available from the UCLA CareConnect website
- For training questions, contact crishelp@mednet.ucla.edu or call 310-267-2273.
- Please visit ResearchGo for more information

Data Management Tools

Data Management Tools available online include RedCap, and the Cohort Selection Tool. REDCap is a secure, web-based application for building and managing online surveys and databases.

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