



Clinical Research Coordinator and GCP

Get Trained by Experts.

ABOUT THIS CLASS

This 2-day class covers the responsibilities and actions of a clinical research coordinator (study coordinator). It includes conceptual lectures on roles and responsibilities of the site and sponsor, a summary of the current regulatory environment, overview of Good Clinical Practices (GCP), study initiation activities, monitoring activities auditing activities, and SOP requirements. The class is a mixture of lecture and break-out team exercises designed to challenge the participants in the various roles and responsibilities of a clinical research coordinator.

FUNCTIONALITY COVERED ...

- Regulatory Requirement
- GCP Overview
- Informed Consent Process
- Study Implementation
- Monitoring Activities
- Auditing Activities
- Standard Operating Procedures (SOP)

WHO SHOULD ATTEND?

This class is designed for people who demonstrate an interest in conducting clinical research for a site or investigator or who are currently in this role and wish to broaden their understanding and knowledge.

OUR INSTRUCTORS

Jacqueline Busheikin, RN, CCRP -
President of JANA Research Consulting,
Inc.

Helen R. Darwin, CCRP - President of
Darwin Site Management Services, Ltd.

PUBLIC TRAINING COST

- \$500/student.
- Please contact us for on-site training costs.

TO REGISTER:

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