



Clinical Research Monitoring and GCP

Get Trained by Experts.

ABOUT THIS CLASS

This 2-day class covers the basics of clinical research from the perspectives of a clinical research monitor and the study site. It includes conceptual lectures on roles and responsibilities of the site, sponsor and CRA, overview of Good Clinical Practices (GCP), site initiation activities, monitoring activities and site closeout activities. 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 monitor.

CONTENT COVERED

- GCP Overview
- Clinical Research Monitor Responsibilities
- Study Site Responsibilities
- Study Initiation Activities
- Monitoring Activities
- Site Closeout Activities
- Clinical Research Monitor Training

OUR INSTRUCTORS

Jacqueline Busheikin, RN, CCRP - President of JANA Research Corporation

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

PUBLIC TRAINING COST

- \$1,000/Participant.
- Includes materials and training costs.

TO REGISTER:

- Call (650) 292-5325
- Email training@biopharm.com