Advantage Clinical

Good Clinical Practice (GCP) E6 R2

Clinical Research Sites

Good Clinical Practice training serves as the pre-requisite for any individual participating in clinical research at any level. Advantage Clinical is proud to offer industry recognized GCP training in our comprehensive custom learning management system.

Advantage Clinical's Good Clinical Practice series goes beyond the guidance and provides students with real world applications and skills to apply GCP in their roles as clinical research professionals.

Within this GCP course we include role-based course branching, where students can focus on how GCP is applied to their particular role as a clinical research professional at your organization. This course includes pathways for:

- Principal and Sub-Investigators
- Clinical Research Coordinators
- Clinical Support Staff

This ICH E6 GCP Investigator Site Training meets the Minimum Criteria for ICH GCP Investigator Site Personnel Training identified by TransCelerate BioPharma as necessary to enable mutual recognition of GCP training among trial sponsors. TransCelerate Biopharma is an industry body representing all of the major pharmaceutical and medical device trial sponsors.

Course Components

Intro to GCP	Components of GCP	Sponsor Responsibilities
History and Evolution	Principles of GCP	Monitoring and Safety
ICH Parties	IRB and Ethics Boards	Protocols and Amendments
Key Objectives of GCP	Investigator Responsibilities	Essential Documentation

For more information on the Good Clinical Practice Training Program and more course offerings please visit www.Advantage-Clinical.com or contact us at:

Email: info@advantage-clinical.com
Telephone: 1-800-674-8802

This course is part of our Organizational Subscriptions program.