WCG Avoca's Training Programs

WCG Avoca offers on-demand web based training programs on a variety of topics to support clinical research professionals in their roles to provide high quality clinical trials. This training solution is easy to deploy, and saves time and money.

Each program is broken down into courses with interactive lessons to ensure attendees are engaged and comprehending key information. WCG Avoca's solution is a flexible and convenient way to train teams.

OVERVIEW OF TRAINING PROGRAMS

TRAINING TOPIC	COURSES	TARGET COMPANIES	FUNCTIONAL TEAM
SPONSOR TRAINING	26	Sponsors/CROs	All staff: Clinical Operations, study teams, quality, risk management, procurement
INTRO TO RESEARCH	17	Biotech/Sites	Clinical Trial Team
GCP	5	Sponsors/CROs/Sites	Study Staff
GLOBAL RESEARCH	8	Sponsors/CROs/Sites	Regulatory Affairs/Compliance, Roles, Sponsors, CROs, Sites
SITE TRAINING	25	Research Sites	All staff participating in clinical trials
REGULATORY	16	Sponsors/CROs/Sites	Regulatory and Compliance professionals
INSPECTION READINESS	7	Sponsors/CROs/Sites	All staff
MEDICAL DEVICE	3	Sponsors/CROs	Med Dev
RESEARCH QUALITY MANAGEMENT	1	Sponsors CROs/Sites	Quality Professionals/Operational Leaders
RESEARCH ETHICS	4	Sponsors/CROs/Sites	New or routine IRB/EC Professional Training
FSP/VENDOR MANAGEMENT OVERSIGHT	1	Sponsors/CROs	Vendor Management Professionals

Our training programs utilize the InvestigatorSpace training platform. InvestigatorSpace is designed by a team of software engineers specifically for use in the pharmaceutical and clinical trial fields. The system is used around the globe by industry professionals at clinical sites, sponsor, and research organizations. All training activity is logged and monitored, and training certifications are available for download by the end-users. Comprehensive Training Reports are also available for training monitoring and audit history.

Please reach out for more information on our course offerings and pricing. WCG Avoca Quality Consortium (AQC) Members get a discounted rate for all Training Programs.

PROGRAM DESCRIPTIONS

SPONSOR TRAINING PROGRAM

For Sponsors/CROs

This program focuses on foundational sponsor training offerings, providing an inclusive curriculum that contributes to a culture of quality and compliance in clinical trial conduct.

INTRODUCTION TO RESEARCH TRAINING PROGRAM

For Biotech companies embarking on their first clinical trial

This program teaches those who are new to the clinical research space the basics of clinical trial execution as it relates to quality, compliance, and patient safety.

GOOD CLINICAL PRACTICE TRAINING PROGRAM

For Sponsors, CROs, Sites

This program is for organizational routine GCP training and GCP training for study staff.

GLOBAL RESEARCH TRAINING PROGRAM

For Regulatory Affairs/Compliance Roles, Sponsors, CROs, Sites

This program is for those involved in global clinical research at Sponsors, CROs, and Sites.

WCG Avoca enables clinical service providers to accelerate their employees' learning and stay up to date on the latest industry regulations and standards.

SITE TRAINING PROGRAM

For Research Sites

This program focuses on foundational site training offerings, providing an inclusive curriculum that contributes to a culture of quality and compliance in clinical trial conduct.

REGULATORY TRAINING PROGRAM

For Sponsors/CROs/Sites, Regulatory and Compliance Professionals

This program is for regulatory and compliance professionals or anyone desiring to have a solid research regulatory training program.

INSPECTION READINESS TRAINING PROGRAM

For Sponsors/CROs/Sites

This program is for any team desiring to stay inspection ready.

MEDICAL DEVICE TRAINING PROGRAM

For Medical Device Professionals at Sponsors/CROs/Sites

This program is for those who are new to or need brushing up on medical device regulations. Upon completing this program, attendees will be able to recognize the requirements for conducting medical device research and identify recommended procedures to follow during a regulatory authority inspection of medical device trials including the ability to recognize the consequences of noncompliance.

RESEARCH QUALITY MANAGEMENT TRAINING PROGRAM

For Sponsors/CROs/Sites, Quality Professionals, Operational Leaders

This program is for research quality professionals or those planning to implement or maintain a quality management system. It provides key foundational elements for focus on GCP quality management systems and processes.

RESEARCH ETHICS

For New or Routine IRB/EC Professional Training, Sponsors/CROs/Site Professionals

This program is for those seeking foundational research ethics training.

FSP/VENDOR MANAGEMENT

For Sponsors/CROs/Sites, Vendor Management Professionals

This program is for those individuals who are involved in any level of vendor selection, qualification, management, and oversight. It highlights key ICH regulations and best practices in the vendor management lifecycle.

Contact Dawn.Auerbach@theavocagroup.com to discuss your training needs.