Course Description:

This combination Classroom Course/Study Group for ACRP Certification provides guidance, for candidates eligible to take the exam, on how to prepare for an ACRP Certification (e.g., CCRA®, CCRC®, ACRP-CP®, CPI®).

N.B.: This course will not cover the medical device or project management subspecialty designations.

The certification programs are now aligned with the Clinical Trial Competency Framework developed by the Joint Task Force for Clinical Trial Competency. Knowledge areas reflect current practice as a monitor, study coordinator and investigator. The exam covers the 6 relevant guidelines: Declaration of Helsinki, ICH E2A, ICH E6, E8, E9 and E11.

The course will help you familiarize yourself with the format of the exam, tackle example questions, and conduct a personal gap analysis to ensure you are fully primed to earn your ACRP Certification. The Detailed Content Outlines (DCOs) will be used to customize your personal needs. Please bring a copy of the DCO with you.

This course includes presentation/hand-outs, interactive classroom exercises, discussions and question-and-answer sessions. No contact hours are offered.

Course Learning Objectives:

1. Be able to list and use the benefits of certification to drive their exam preparation efforts
2. Be familiar with the question-style and layout of ACRP’s Certification exams, including how to approach their multiple-choice format
3. Be able to identify and describe at least three (3) strategies for preparing for ACRP’s Certification exams
4. Be able to use and reference the six (6) relevant guidelines used to create the ACRP’s Certification exams:
   a. The Declaration of Helsinki
   b. ICH E2A—Safety Definitions
   c. ICH E6(R2)—Good Clinical Practice
   d. ICH E8—General Considerations for Clinical Trials
   e. ICH E9—Statistical Principles for Clinical Trials
   f. ICH E11—Clinical Trials in the Pediatric Population
5. Be able to assess their individual level of readiness by practicing exam questions based on the Detailed Content Outline (DCO) under training conditions (NOT exam conditions)
Agenda:

8.30AM  
Registration Sign-in and Light Breakfast

9:00–9.15  
Welcome and Introduction

9:15–10:00  
Module 1: Format of Exam– What to Expect

10:00–11:00  
Module 2: Overview of the 6 Relevant Guidelines

11:00–11:15  
Refreshment Break

11:15–12:00PM  
Module 3: Personal Analysis of DCO

12:00–12:45  
Lunch provided

12:45–2:30  
Module 4: Kick-Start of Study Group/Sample Questions

2:30–2:45  
Refreshment Break

2:45–4:15  
Module 4: Kick-Start of Study Group/Sample Questions

4:15–4:30  
Wrap Up Questions and Answers, Adjourn

Target Audience:

Eligible candidates for the CCRA®, CCRC®, ACRP-CP®, and CPI® ACRP Certification Exam.


For more information about application and exam fees visit: [www.acrnet.org/certification](http://www.acrnet.org/certification)

Questions About Certification Benefits, Requirements, and Preparation: [www.acrnet.org/certification](http://www.acrnet.org/certification)

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Event Registration Information: Membership not required for registration. Bring your email confirmation to the event. No refunds or transfers. No contact hours offered.

Event Manager For Course Registration or Cancellations: Lisette Gonzalez, CCRA, New York Metropolitan Chapter of ACRP, lgonzalez@phase2phaseconsulting.com or call 917-881-4295.

Questions About This Classroom Course: Janet Ellen Holwell, CCRC, CCRA, FACRP, TIACR, Communications Chair, New York Metropolitan Chapter of ACRP, jholwell@aol.com or call 718-263-4160.

Sponsorship Provided by

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