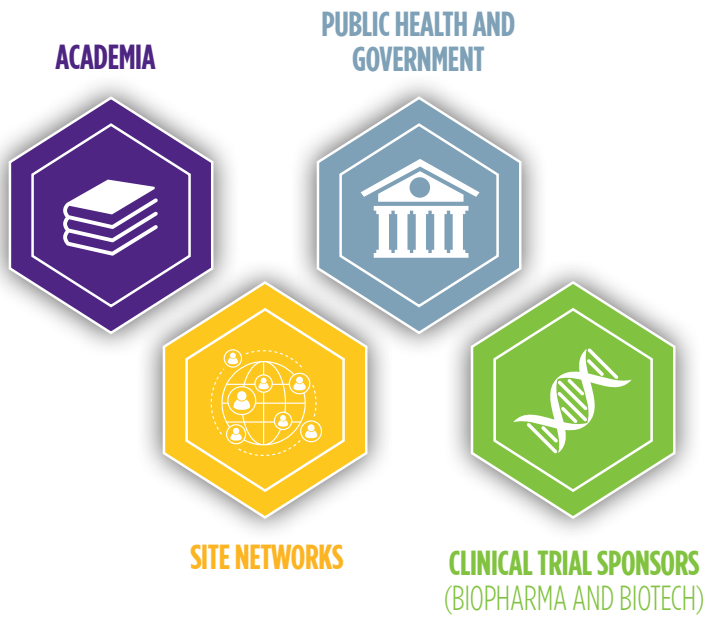
OVERVIEW

SiteCoach: Advanced Clinical Research Training Program for Emerging Sites

Structure:

SiteCoach is a instructor-led virtual training program that was built around the Joint Task Force Competency framework from Clinical Research Professionals. It is aligned to the current Good Clinical Practices guidelines and illustrated by real life scenarios and interactive activities. It 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. Graduates receive the certificate after the course is completed.

Who is SiteCoach for?



How will SiteCoach prepare my site?

Global Outreach

Translated into Spanish, Chinese and Japanese, with capabilities to deliver in Portuguese. SiteCoach is also accredited for CME in South Africa.

PPD's SiteCoach Training Program is designed to support Investigational Sites on their path to excellence by coaching each staff member to become a successful contributor to the conduction of clinical trials.

- Interactive activities, critical thinking scenarios and case study reviews
- Access to a robust toolbox that contains a wide variety of support documents, from regulations and guidelines to site document templates and trial management tools

WHAT OUR CUSTOMERS ARE SAYING

Agree SiteCoach is a worthwhile investment

Are very satisfied with SiteCoach training

Agree SiteCoach training increased their confidence in conducting clinical trials

Agree SiteCoach training encouraged their repeat participation in clinical trials

COURSE STRUCTURE

Four Modules Tailored for Your Organization

Introduction to Clinical Research

Provides an overview of the history and ethics of clinical research, the key steps in the drug development process together with the associated key players. Further, it will get trainees acquainted with the clinical research entities and will guide participants through the trial design, treatment assignment and study monitoring.

Ready, Steady, Recruit!

Focused on the site activation phase and will include information about essential documents, study protocol and its elements, qualification of the site and the monitoring tasks associated with the site initiation visit.

Ongoing Trial Activities

Provides insight about the responsibilities of the Principal Investigator, including communication with the CRA and Clinical Research Coordinator, investigational product cycle at the site, subject safety, audits, etc.

Effective Trial Management

Focuses on communication and strategies for effective study and site management, communication and teamwork. It supports site staff in becoming successful leaders and professionals in conducting clinical trials.

VIRTUAL OR FACE-TO-FACE WORKSHOPS

For more information please contact us at +1 877 643 8773, +1 919 456 5600
or at ppdinfo@ppd.com. www.ppd.com

PPD®