

QTECH-SOL PROFESSIONAL DEVELOPMENT CENTER

Clinical Science Training Institute



ADVANCED Clinical Research Online Program (CROP)

Are you currently working as junior CRA or on other entry level CRA position?

Are you ready for career advancement to the higher level of CRA role?

Did you complete basic CRA training, but you wish to add a new skill set?

Self-Paced Online Training

Our Mission

Our mission is to provide the best-in-class job oriented certification and skill based courses towards Clinical research, Drug Safety, Pharmacovigilance, Clinical Data Management, Clinical SAS Data Analytics and Healthcare business. We offer Entry-Mid and Senior programs for students and professionals, looking for skills refresh or career advancement

CAREER FOCUSED PROGRAM

(Learning for Job)

Website: www.qtech-solutions.com Email: qpdc@qtech-solutions.com

OVERVIEW OF THE CURRICULUM

Designed by industry experts for professionals looking to add new skill set.

LIST OF TOPICS

* Real time scenario cases are assigned to each topic:

- 1. Clinical Trial Budget
- 2. Investigator Selection
- 3. Pre-Study Visit
- 4. Protocol
- 5. Informed Consent Preparation
- 6. Investigational New Drug (IND) Application
- 7. Institutional Review Board (IRB) Regulatory Correspondence
- 8. Case Report Form (CRF)
- 9. Site Monitoring
- 10. Co-Monitoring Visits
- 11. Study Initiation Visit
- 12. Clinical Trial Management Systems (CTMS)
 Tracking Recording
- 13. Trial Master File
- 14. Database Lock
- 15. Audit

Education Preferred by Employers Hiring for CRA positions:

Pharmacy - 40%
Pharmacology or Toxicology - 60%
Nursing - 50 %
Public health - 40%
Biotechnology - 30%
Medicine - 30%
Clinical Research - 99%

LIST OF TOPICS: THEORETICAL ASPECT OF THE FIELD

- 1. Risk Based Monitoring (RBM)
- 2. FDA Audit process for Clinical Research
- 3. Clinical Trial Project Management (CTPM) and Time-lines
- 4. Development of Monitoring Plan
- 5. Protocol Deviation/Violation Management
- 6. Trial Master File and QC Management
- 7. Clinical Data Reconciliation and Archiving
- 8. Management and Reconciliation of Investigational Product
- 9. Advanced Clinical Research Management -1
- 10. Advanced Clinical Research Management -2
- 11. Advanced Clinical Research Management -3
- 12. Advanced Clinical Research Management -4
- 13. Planning and Conducting Global Clinical Trials
- 14. Management of a Successful Clinical Research Site A
- 15. Management of a Successful Clinical Research Site B

The typical career pathway of CRA begins from entry level position working as Clinical Research Assistant.

The promotion to mid level positions comes with progress and experience while the opportunities remain open for Clinical Research Associate and Documentation Specialist.

TYPICAL CAREER PATH OF Clinical Research Associate

CRA (Monitor, In House); Clinical Research Documentation Specialist; Project Coordinator; CRA I, II; Senior CRA; Principal CRA



Entry Level Mid Level



Clinical Research Trial Monitor; Clinical Project Manager; Clinical Research Manager; Associate Director of Clinical Research; Director of Clinical Research

Management Level

What is the Difference Between Basic CRA and Advanced CRA?

C R A

Basic and core topics and exercises

Certain educational background is strongly recommended to enter the industry

Designed for non experienced individuals to enter into field

In depth knowledge of roles and responsibilities of CRA + theoretical aspect with projects which help to understand industry requirements Advanced

Advanced topics and exercises

Prior training or relevant experience is required

Designed for experienced professionals to add new skills set

Exposure to diverse CRA exercises, real time documentation + advanced learning supported by practical case scenario tasks and follow up process



Contact Information

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