



QTECH-SOL PROFESSIONAL DEVELOPMENT CENTER LLC

Somerset New Jersey USA

Private Vocational School

Training Division of Qtech Solutions Inc.



ADVANCED Clinical Research Associate Training

Our Mission

Our mission is to provide the best-in-class job oriented career development Elearning training courses and programs in clinical research, SAS data management and business analysis for students and professionals requiring a skills refresh – or the development of new skills and experience for job entry, advancement, and placement.

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 new skill set?

If your answer is yes to any of the questions above, starting advanced CRA training program might be a right decision. How quickly you will be able to move up grades surely depends on your work experience, but also motivation, and possessing advanced CRA skills.

OVERVIEW OF THE CURRICULUM

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



LIST OF TOPICS

*** The real time scenario case is 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

What is the Difference Between Basic CRA and Advanced CRA ?	
CRA	ACRA
Basic and core topics and exercises	Advanced topics and exercises
Certain educational background is strongly recommended to enter the industry	Prior training or relevant experience is required
Designed for non experienced individuals to enter into field	Designed for experienced professionals to add new skills set
In depth knowledge of roles and responsibilities of CRA + theoretical aspect with projects which help to understand industry requirements	Exposure to diverse CRA exercises, real time documentation + advanced learning supported by practical case scenario tasks and follow up process

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.

Main Roles and Responsibilities of Mid-Level Clinical Research Associate -

- Performing site selection, initiation, monitoring and close-out visits, plus maintaining appropriate documentation
- Supporting the development of a subject recruitment plan
- Establishing regular lines of communication plus administering protocol and related study training to assigned sites
- Evaluating the quality and integrity of site practices – escalating quality issues as appropriate
- Managing progress by tracking regulatory submissions, recruitment, case report form (CRF) completion, and data query resolution

TYPICAL CAREER PATH OF Clinical Research Associate



Do You Know...?

Hiring Clinical Research professionals:

- 99% of the employers choose candidates with clinical research degree or experience
- 75% of the employers choose candidates with pharmacology or toxicology degree
- 80% of the employers choose candidates with nursing degree
- 40% of the employers choose candidates with Biological Science degree
- 80% of the employers choose candidates with public health degree
- 65% of the employers choose candidates with Biotechnology degree
- 90% of the employers choose candidates with Medicine degree

Do you want to know more? Call us!



Contact Information

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