QTECH-SOL PROFESSIONAL DEVELOPMENT CENTER LLC

Somerset New Jersey USA

Private Vocational School

Clinical Research Associate Workshops

Our Mission

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

APPROVED

Our Clinical Science Professional training program is approved by State of New Jersey, Department of Education and Department of Labor and Workforce Development and listed on Eligible Training Provider List

STATE OF NEW JERSEY
DEPARTMENT OF EDUCATION

STATE OF NEW JERSEY
DEPARTMENT OF LABOR
AND WORKFORCE DEVELOPMENT

- **Qtech-Sol Professional Dev Center LLC (QPDC)** is a Private Vocational School (PVS) with its corporate headquarters in New Jersey and regional offices in India.
- We provide professional job-oriented Elearning training courses and Workshop programs for students and industry professionals globally. Our clinical research training courses are approved by **the State of New Jersey Department of Education and Department of Labor and Workforce Development** and listed on New Jersey Eligible Training Provider List.
- We are committed and completely focused on fulfilling the specialized skills and knowledge needs of professionals in the following fields:

  - ✓ **Clinical Research**
  - ✓ **Drug Safety and Pharmacovigilance**
  - ✓ **Clinical SAS Programming**
  - ✓ **Financial Banking**
  - ✓ **Health Care Insurance**

OUR PROGRAMS

Rather than providing generic training programs like many other marketplace training services companies, we decided to take our combined clinical, banking and learning development expertise to the highest level and design the best in the industry training programs to ramp up quickly the skills, knowledge, and practical experience of our career-oriented global customers.

OUR TEAM

Our Training Development and Delivery Staff consists of an outstanding team of industry professionals and consultants with years of clinical science and business experience who will work with students and industry professionals for the training courses and programs needed to help achieve their job and career goals.

QUALITY

Our professional clinical science, business, and training staff including consultants and advisors bring their scientific, regulatory, and business acumen to produce industry-focused career training.

Clinical Research Associate Workshops

The following are the workshops offered at Qtech-Sol Professional Development Center. Each of the workshop provided is a 3 day program. Access to Online workshop material is provided to enrolled participant for workshop delivery. Enrolled Participants will attend WebEx Meeting to understand the workshop class and will submit their learning's as MCQ, Discussion Based Questions – DBQ and Exercises for delivery. The Exercises are evaluated for Feedbacks and Program Certification. The Program is designed by industry experts in domain, for students and young professionals. The training provides in-depth knowledge pertaining to the role and responsibilities performed by "**Clinical Research Associate**" for conducting Clinical Trials.

Workshop Listing Offered	
1. Role of Clinical Research Associate	17. In House CRA Responsibilities
2. Drug Discovery and Research Process	18. Clinical Protocol Design and Development
3. Pre Clinical Research	19. Clinical Trial Budget
4. Introduction to Clinical Trials	20. Selection of Investigator
5. Phase I Clinical Trials.	21. Roles and Responsibilities Of Investigator
6. Phase II Clinical Trials.	22. Selection Of Investigation Site
7. Phase III Clinical Trials.	23. Study Initiation
8. Phase IV Clinical Trials.	24. Selection And Vendor Management
9. FDA Regulations	25. CRF Design And Data Capture
10. Good Clinical Practice and ICH Guidelines	26. Investigator Meetings And Timelines
11. Institutional Review Board (IRB)	27. Introduction To Adverse Event Reporting And Classification
12. SOP Development	28. Trial Master File (TMF)
13. A 6-Month Process for Planning Multinational Clinical Trials	29. In-House Monitoring & Reporting
14. Communication With Cross Functional Team	30. Informed Consent Preparation and Review
15. Overview of Protocol	31. Preparing For Internal Audit
16. Medical Dictionary – MedDRA Coding	32. Clinical Trial Management System (CTMS)

Do you want to know more? Connect with us!

Contact Information

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