



# QTECH-SOL PROFESSIONAL DEVELOPMENT CENTER LLC

Somerset New Jersey USA  
Private Vocational School



## Professional Development and 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**



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





- **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.



# QPDC Workshop

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 and evaluation. 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 (OR) Drug Safety Associate at Job .

## WORKSHOP LEARNINGS

### CLINICAL SCIENCE WORKSHOPS

#### **Clinical Research Associate Training and Workshop Learning's**

The candidate will be working on various workshops to perform the following tasks. Case 1: Introduction To Clinical Trial , Case 2: FDA EMEA Regulations , Case 3: Institutional Review Board (IRB) , Case 4: Protocol Design and Development, Case 5: Clinical Trial Budget , Case 6: Case Report Form (CRF) Design , Case 7: Investigator Meeting , Case 8: Site Management and Initiation , Case 9: Informed Consent Preparation , Case 10: Trial Master File , Case 11: Adverse Event Monitoring and Reporting, Case 12: Audit

#### **Drug Safety Associate Training and Workshop Learning's**

The candidate will be working on various workshops to perform the following tasks. CASE 1. Introduction to Adverse Events , CASE 2. Role of DSA, CASE 3. Characteristics of Case , CASE 4. Medical Record Extraction, CASE 5. Basics of Coding , CASE 6. Triage, CASE 7. Case Narratives , CASE 8. SAE Reconciliation, CASE 9. Drug Safety Database and Software AND Other workshop learning on Medical Record Extraction , Adverse Events Case Processing, CIOMS Line Listing , Case processing and FDA Reporting for Medical Devices, Revision of SOP Quality Control Procedure, SAE Reconciliation, PSUR - Periodic Safety Update Reporting, Triage, Data Entry, Signal Detection, Labeling Edit check, Quality Control Procedure, Resolution of queries of pending cases , SUSAR – Suspected Unexpected Serious Adverse Reaction

#### **Clinical Data Management Training and Workshop Learning's**

The candidate will be working on various workshops to perform the following tasks. CASE 1. Introduction to Clinical Trials, CASE 2. Protocol Design and Development, CASE 3. Data Management Plan, CASE 4. Data Cleaning and Data Validation., CASE 5. Query Management, CASE 6. Coding of Adverse Events, CASE 7. SAE Reconciliation., CASE 8. Elements of CRF, CASE 9. e-CRF designing, Data tracking from CRF.

## ANALYTICS & REPORTING WORKSHOP

### Clinical SAS Oriented - Data Management Workshop Learning's

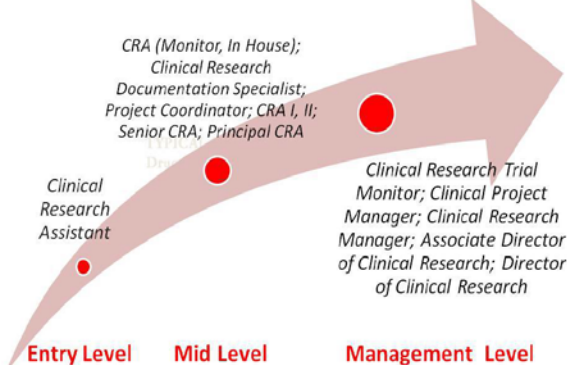
The candidate will be working on various workshops to perform the following tasks.(1) SAS Efficiency Programming (2) Data based Validation (3) Protocol based Validation, Pharmacokinetics and Pharmacodynamics (5) Preparing Analysis Datasets CDISC (6) Oncology project Phase-I (7) Ophthalmology Project Phase-II (8) Cardiology Project Phase-III (9) Central Nervous System (CNS) Project (Phase-IV)(10) Aggregate Reporting Process (Pharmacovigilance) (11) Open CDISC Validation.

## HEALTHCARE BUSINESS ANALYST WORKSHOP

### Healthcare Business Analyst Workshop Learning's

The candidate will be working on various workshops to perform the following tasks. (1) Overview of US Healthcare system (2) Healthcare Claims (3) Healthcare Claims Forms (4) Public Health Coverage (5) HIPAA – Health Insurance portability and Accountability Act (6) International Classification of Diseases (ICD) (7) Healthcare Codes (8) Overview of Medicaid Management Information System (MMIS) (9) NCPDP – National Council for Prescription Drug Product (10) Healthcare Data warehouse.

### TYPICAL CAREER PATH OF Clinical Research Associate



### TYPICAL CAREER PATH OF Drug Safety Associate



Do you want to know more? Connect with us!

### Contact Information

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