QTECH-SOL PROFESSIONAL DEVELOPMENT CENTER LLC

Somerset New Jersey USA

Private Vocational School

Clinical Research Associate (CRA)

CLIENT SETTING PROGRAM

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

APPROVED

Clinical Research Associate 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

Website: www.qtechelearncenter.com **Email:** qpdc@qtechelearncenter.com

About Clinical Research

Clinical research is a rapidly expanding field, creating exciting job opportunities. Clinical Research Associates work in a broad range of research settings, including: academic health centers, government agencies and departments, contract research organizations, pharmaceutical, biotechnology and medical device firms.

INT - CLIENT SETTING PROGRAM

Program was designed by industry experts for students and young professionals. This Training program provides in depth knowledge of roles and responsibilities of Clinical Research Associate performed at various Pharmaceutical and Research Organizations. Interested Student must enroll into the program for participation. Recent graduate students who have completed their graduation (past 12 months) or Currently pursuing their graduation are eligible for this Client Setting program. Proof of eligibility must be submitted when requested.

Lessons

Preclinical Research
Introduction to Clinical Trials
SOP Development
In House CRA Responsibilities
Protocol Design and Development
Subject Recruitment Process and Informed Consent
Selection of Investigation Site
Site Monitoring
Study Close Out

DURATION : 30 Days

Delivery

This Training program is ONLINE. Each student will be provided access to Learning Material for 30 Days. The Student must spend min 3 hours/day (Mon thru Fri). During this period they will attend scheduled Online classes and will submit the exercise questions by lesson for evaluation using our Online Learning Management System. All Questions and Queries during the Training program will be addressed via email. This is a Long Distance Training program. Project Assignments will be provided for delivery. QPDC Terms and Conditions Apply for participation into this program and QPDC reserves the right for admission and determines the eligibility for participation.

Certification and Post Training Assistance

At the end of this Training program student will take a Scenario Based Objective Online Exam to get course completion certificate. Exam fee applies. The Student must secure minimum 75% score in final exam. The Student can purchase reading material book for their future reference. After the course completion, students will be assisted in their resume preparation, narrative writing and mock interviews for applying jobs.

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Typical Duties of Clinical Research Associate

- ✓ Writes, edits and assists in writing protocols
- ✓ Coordinates protocol review for clinical studies
- ✓ Writes and reviews patient informed consents
- ✓ Designs and develops case report forms (CRFs)
- ✓ Helps, writes, assembles, and distributes investigator brochures, SOP documents and study/CRF instruction manuals
- ✓ Prepares and submits documents required to meet regulatory, GCP, and SOP requirements
- ✓ Determines order, ships, and tracks investigational drug supplies
- ✓ Evaluates and selects investigators (sites), plans and conducts investigator meetings
- ✓ Develops study budgets and grant payment schedules
- ✓ Sends study packages such as protocols, brochures, and contracts/agreement letters to field monitors for site initiation
- ✓ Assures all adverse events (AEs) are reported according to regulation and company policy
- ✓ Prepares final study reports and assist in the response or reporting to any FDA inspections

Who Can Qualify ?

Education Qualification:

- ✓ 99% of the employers choose candidates with **Clinical research** degree
- ✓ 99% of the employers choose candidates with **Bachelors Degree in Science**.
- ✓ 75% of the employers choose candidates with **pharmacology or toxicology** degree
- ✓ 80% of the employers choose candidates with **nursing** degree
- ✓ 80% of the employers choose candidates with **public health** degree
- ✓ 75% 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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