



## UNIVERSITY CERTIFICATION

Delaware Valley University issues a Certificate of Completion that demonstrates your qualification and CRMC course completion. Additionally, this course qualifies health professionals with 33 ANCC credits obtained.

### Skills and Knowledge to Manage your job

Upon completion, you will have the basic training to participate in the drug development process, identify ethical issues in clinical research, and understand sound clinical practices as they relate to studies designed to support medications awaiting world-wide marketing; in general, manage a Phase III trial.

## TARGET AUDIENCE

Persons already in a science or healthcare field or seeking entry. Health care professionals such as: LPNs, RNs, BSNs, Physicians, Dentists, Pharmacists, Therapists, Scientists, or Laboratory technician and Medical Technicians and all other Science professionals, who want to work in the pharmaceutical industry. It is an excellent refresher for those already working in the clinical research field who need updated GCP training in the US and International standards.

### Course Objectives

Upon completion of this course, participants will be able to:

- Describe the drug development process and roles/opportunities in Clinical Research.
- Understand the latest Good Clinical Practices.
- Identify ethical issues and their impact on the development of new products.
- Monitor clinical sites and study trials
- Coordinate all data, records, and activities involved in a clinical study.
- Learn details of Clinical Data Management.
- Institute and obtain proper Informed Consent.
- Employ efficient subject recruitment methods.
- Develop a suitable clinical trial protocol and suitable study budget.
- Initiate and examine ongoing study activities
- Identify adverse effects and proper reporting format.
- Learn to about audit & inspection techniques.
- Develop critical understanding of Regulatory Approval expectations

### Frequently Asked Questions

This course is offered in collaboration with Applied Quality Solutions. Applied Quality Solutions is an independent education company that designs, produces and presents continuing education courses to the pharmaceutical and biotech industries across the country. The course is nationally accredited through the ANCC. For specific course content questions, please contact Juliet at [appliedqualitysolutions@yahoo.com](mailto:appliedqualitysolutions@yahoo.com).



## CLINICAL RESEARCH MONITORING AND COORDINATION FAQ

### **How is the course set up and organized?**

Your multimedia lectures and live slides are viewable on any PC or smartphone. There are ten chapters and two exams. Those chapters start with a video introduction and then followed by a slide show with instructor narration voice over. Each chapter has a home work assignment.

### **How much does the course cost?**

The cost of the entire program is \$1,875.

### **Am I eligible for a refund if I don't complete the course?**

Once you receive course materials, refunds are not available.

### **How much are the textbooks? Where can I purchase them?**

The price of the course includes all required course materials. Textbooks will be sent via USPS priority mail upon registering for the course.

### **How long does it take? How long do I have to finish the course?**

It should take the average learner about 35 hours. You can spread those 35 hours over a period of time as needed. The maximum time the University allows is four months.

### **What payment options are available?**

We accept checks, credit cards, and purchase orders.

### **Do you offer Financial Aid?**

This university does not offer financial aid for continuing education courses.

### **Can I get a student loan?**

Yes, as long as the lender gives you the check and they make the check out to either you or the university. Because you are not a matriculated student, student loans that have to be processed through the Office of Financial Aid are not available.

### **Which credit cards do you take?**

American Express, Discover, Mastercard and Visa.

## CONTINUING EDUCATION CREDITS

### **Will I receive Nursing Continuing Education contact hours (CE) for this program?**

Yes, this program is approved for 33 Nursing Continuing Education contact hours by the ANCC.

### **How many college credits is this course?**

This is a non-credit continuing education course.

**Is this an accredited course?**

Yes. This course is designed for professionals wishing to enter the clinical research field or to update their resumes. It can be used as a prep course prior to taking one of the various industry group certification exams. Most of these certifications require that individuals must meet additional experience requirements before qualifying to sit for the exam. The FDA does recommend continuing education, but does not require specific type of certification to work in research.

**How long does it take to get my certificate?**

After you have successfully completed the course, it takes about two weeks for the University to send you the certificates

**What does this course cover to advance my skills in clinical research?**

This course provides a thorough overview of the roles and responsibilities of the clinical research monitor and coordinator. It has been created to provide you with the key skills, job criteria and industry expectations of both job roles. Upon completion, you will have the basic training to participate in the drug development; identify ethical issues in clinical research; and understand good clinical practices as they relate to studies designed to support medication awaiting worldwide marketing. This course will also provide you with a certificate that shows potential employers and the FDA that you have current training from MSU College of Nursing, a nationally recognized and accredited university.

**What does a clinical research coordinator do?**

Clinical Research Coordinators work directly with physicians during clinical trials. These CRCs ensure proper treatment and care of patients who participate in the tens of thousands of ongoing worldwide research trials.

**What does a clinical research associate or monitor do?**

Clinical Research Associates (CRAs) or Monitors are critical pharmaceutical team representatives who oversee ethical treatment of patients while assuring quality conduct of the trials: exciting travel and professional networking highlight the breadth of this career opportunity.

**I am not sure I need to take the CRCM course. Who should take this course?**

Life Science majors, nurses and physicians are the major audience. Other science degreed individuals should consider enrollment since the pharmaceutical industry employs such persons that have a diverse understanding of scientific principles needed for the wide-range of therapeutics and medical devices currently being developed. Other professionals that may consider this course include: Chemists, Engineers, Pharmaceutical Sales Representatives, Medical Technologists, Laboratory Technicians, Clinical Research Associates, Physicians, Dentists, Physician Assistants, Medical Monitors, Pharmacists, Pharmacologists, Physical Therapists, Respiratory Therapists, Psychologists, Biologists, Medical Writers and Data Managers.

**Are there any prerequisites for this course?**

A Bachelors degree is typically required to be hired as a CRA. If you have some type of life sciences background, you will be able to better understand the medical science side of the course, but it is not strictly necessary. You do not need medical training to be a CRA. If you do not have a Bachelors Degree, contact [appliedqualitysolutions@yahoo.com](mailto:appliedqualitysolutions@yahoo.com) discuss your eligibility.