Calgary Centre for Clinical Research

New CITI Canada "Clinical Research Coordinator" Training Course Available

We are pleased to announce N2’s release of a new CITI training course - the Clinical Research Coordinator course with Canadian content.

What is the Clinical Research Coordinator Course about? This course provides clinical research staff with the foundational skills needed to successfully and confidently operationalize clinical research studies. The course includes thirteen modules that provide clinical research staff with the skills on how to review a protocol, conduct informed consent and manage finances and resources. Below is a complete list of the course modules:

- Planning Research
- Funding, Financial Management, and Budgeting
- Working with the Research Ethics Board (REB)
- Protocol Review and Approvals
- Principal Investigator (PI) Responsibilities
- Clinical Research Coordinator (CRC) Responsibilities
- Sponsor Responsibilities
- Informed Consent
- Site Management, Quality Assurance, and Public Information
- Overview of the Clinical Trial Agreement (CTA)
- Coordinating U.S. Regulated Research Studies – What to Consider?

More information about each module can be found in the attached summary.

Does the Course require any Prerequisites? While there are no prerequisites, we recommend to learners to review the CITI course on Biomedical Research Ethics which provides an excellent background on research ethics involving human participants, prior to completing the Clinical Research Coordinator course.
Who should take this course: The course is recommended for onboarding new Clinical Research Coordinators and other clinical research staff.

It would also be helpful for new investigators as well as the experienced investigators who are interested in learning the practical components of a clinical trial.

You can access the course by logging into your CITI account. Please contact linda.longpre@ucalgary.ca with any questions you have.

Please click here to learn more about the core modules