



Clinical Research Curriculum Certificate (CRCC) Application Checklist

The CRCC program is currently only for those employed at the NIH as staff, trainees, and contractors.

TO APPLY:

1. Review the requirements below and choose your certificate type.
2. Complete the requirements and retain documentation. All requirements must have been completed within 2 years of your application date, unless stated otherwise.
3. Email crcc@mail.nih.gov to request a link to upload your CRCC application documentation.



CRCC Certificate

- ☐ **Introduction to the Principles and Practice of Clinical Research (IPPCR) Course**
Submit a certificate of completion
- ☐ **Ethical and Regulatory Aspects of Clinical Research: Live Course or Asynchronous Course**
Submit a certificate of completion
- ☐ **NIH CITI Training - Good Clinical Practice Course - US FDA Focus**
Submit a completion report showing the following 5 modules at minimum:
 - CITI Good Clinical Practice Course for Clinical Trials Involving Drugs and Devices
 - Conducting Investigator-Initiated Studies According to FDA Regulations and GCP
 - Investigator Obligations in FDA-Regulated Research
 - Audits and Inspections of Clinical Trials
 - Detecting and Evaluating Adverse Events
- ☐ **Attend one virtual IRB meeting as a guest observer.**
Contact crcc@mail.nih.gov to schedule your attendance. Attendance will be confirmed directly with OHSRP (no need to submit documentation).



CRCC Certificate with Commendation

- ☐ **Submit all required documents of the CRCC Certificate as outlined above.**
- ☐ **Principles of Clinical Pharmacology Course** or **FAES course in statistics or epidemiology**
Submit a certificate of completion *Submit an unofficial transcript showing course within the last 5 years and no withdrawal or drop of the course.*