

Clinical Research Curriculum Certificate (CRCC)

Application Checklist

(The CRCC program is currently only for NIH staff, trainees, and contractors.)

Please decide which type of certificate to apply for and have all documentation before submitting your application package to crcc@mail.nih.gov. All requirements must have been completed within 2 years of your application date, unless stated otherwise.

For the CRCC CERTIFICATE:

[Introduction to the Principles and Practice of Clinical Research \(IPPCR\) Course](#)
certificate of completion

[Ethical and Regulatory Aspects of Clinical Research Course](#)
certificate of completion

[NIH CITI Training](#) – Good Clinical Practice Course (US FDA Focus)
certificate of completion showing the following 5 modules:

- 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

Attendance at one virtual IRB meeting as a guest observer. View the [OHSRP event calendar](#) for IRB meetings and contact crcc@mail.nih.gov to schedule your attendance.

For the CRCC CERTIFICATE WITH COMMENDATION:

All requirements for the CRCC Certificate **and** one of the options below:

[Principles of Clinical Pharmacology Course](#) certificate of completion **or**

Successful completion of an [FAES](#) course in statistics or epidemiology in the last 5 years, verified by an unofficial transcript showing no withdrawal or drop of the course.