Collaborative Institutional Training Initiative (CITI)
UCF Training Courses

Humans Subjects Protection

Investigators (PI and key research personnel who have contact or communication with subjects' or access to private identifiable information for research purposes and/or are involved in the consent process) are required to complete human research protections training through CITI before conducting human research.

For more information see http://www.research.ucf.edu/Compliance/IRB/Investigators/training.html

Responsible Conduct of Research (RCR)

(NSF) Postdoctoral researchers, graduate, and undergraduate students on NSF awards performing research or training activities on NSF awards must take a RCR course when added to payroll.

(NIH) Trainees, fellows, participants, and scholars receiving support through any NIH training, career development (individual or institutional), research education grant, and dissertation research grant must receive RCR training. Online training in CITI may be a component of the required training. (UCF College of Graduate Studies) Doctoral program students are required to take RCR training.

For more information see http://www.rcr.ucf.edu/resources.html

Conflicts of Interest (COI)

Investigators (PI, Co-PI, project director, and any other person who is responsible for the design, conduct, or reporting of research) are required to complete the COI Basic Course prior to engaging in sponsored research and the COI Refresher Course every four (4) years thereafter.

For more information see http://www.coi.ucf.edu/training.html

Animal Welfare Learning Group

All personnel listed on an animal protocol and/or involved in animal research are required to take Animal Welfare training.

For more information see http://www.research.ucf.edu/Research/OfficeOfAnimalWelfare.html

Export Controls

Export Controls training is required for researchers engaging in restricted research programs. Researchers (PI and project personnel) subject to export controls will be contacted by UCF Export Control office when training is required.

For more information see http://www.research.ucf.edu/ExportControl/training.html

Good Clinical Practice (GCP)

Good Clinical Practice (GCP) includes four basic courses tailored to the different types of clinical research. These courses are acceptable GCP training for NIH requirements.

IRB Administration

IRB Administration is a role-specific optional course designed for IRB administrators, directors, coordinators, and other staff that support the IRB and Human Research Protection Program (HRPP).