

New to the CITI Program? Read the getting started guide or watch the getting started video.

Need Help? <u>Support Center</u> Status Page

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LOG IN

LOG IN THROUGH MY ORGANIZATION

REGISTER

Organizations listed here use "Single Sign On" (SSO) for CITI Program access.

SSO requires a username and password issued by the organization.

If your organization is not listed here, it does not use Single Sign On. Click on the "Log In" tab (if you already have a CITI Program account) or the "Register" tab (if you are new to CITI Program and creating an account for the first time).

To find your organization, enter its name in the box below, then pick from the list of choices provided.

University of Louisiana

University of Louisiana at Lafayette

See our full list of SSO-enabled organizations

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University of Louisiana at Lafayette

Continue To SSO Login / Instructions

See our full list of SSO-enabled organizations



Secure Login

This secure site uses UL Lafayette credentials to allow students, faculty and staff to access various systems used on campus, such as Ulink, Moodle, and Webmail.

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I forgot my password | Activate my account

SIGN IN

Learner Tools for University of Louisiana at Lafayette

Add a Course

- Remove a Course
- View Previously Completed Coursework
- Update Institution Profile
- View Instructions Page
- Remove Affiliation

 SUPPORT
 LEGAL

 888.529.5929
 Accessibility

9:00 a.m. – 7:00 p.m. ET <u>Copyright</u>

Monday - Friday

Privacy and Cookie Policy

Contact Us Statement of Security Practices

Terms of Service



Steps: 1 2 3 4 5 6 7
Select Curriculum
* indicates a required field. You will be provided a series of enrollment questions. Your responses will determine the curriculum for the courses you are going to take. Please read the questions carefully. Please read the responses carefully to make the best choice.
Question 1
Responsible Conduct of Research Please make your selection below to receive one of the courses in the Responsible Conduct of Research. Choose one answer Biomedical Responsible Conduct of Research Course Social and Behavioral Responsible Conduct of Research Course Physical Science Responsible Conduct of Research Course Humanities Responsible Conduct of Research Course Responsible Conduct of Research for Engineers Responsible Conduct of Research for Administrators Biology Summer REU Students Not at this time.
Question 2
Would you like to take the Conflicts of Interest course? Choose one answer Yes No
Question 3
CITI US Export Control Please make your selection below to receive the CITI US Export Control Regulations course. Choose one answer O CITI Export Controls Not at this time.

CITI - Learner Registration - University of Louisiana at Lafayette

Question 4
Biosafety/Biosecurity Please make your selection below to receive the courses in the Biosafety/Biosecurity Course.
Choose one answer
O Introduction to Biosafety
O Introduction to Biosafety for Students
O Basic Biosafety Training
O Biosafety Retraining
O Animal Biosafety
Shipping and Transport of Regulated Biological Materials
OSHA Bloodborne Pathogens
O Select Agents, Biosecurity and Bioterrorism
Emergency and Incident Response to Biohazard Spills and Releases
O Human Gene Transfer Trials
O NIH Recombinant DNA (rDNA) Guidelines
OSHA Personal Protective Equipment Training
O Institutional Biosafety Committee Member
O Biosafety Complete Training
O Dual Use Research of Concern (DURC)
Not at this time.
Question 5
If you want to take Health Information Privacy and Security (HIPS) ontionally, please make your selection below

Question 5
If you want to take Health Information Privacy and Security (HIPS) optionally, please make your selection below.
Health Information Privacy and Security (HIPS) Course - Information for Clinicians
\square Health Information Privacy and Security (HIPS) Course - Information for Investigators
☐ Health Information Privacy and Security (HIPS) Course - Information for Students or Instructors
☐ Health Information Privacy and Security (HIPS) Course - Information for Fundraisers
☐ Health Information Privacy and Security (HIPS) Course - Information for Marketers
✓ Not at this time.

Question 6
Laboratory Animal Research
Do you conduct studies that use Lab animals? 1. If YES, then you must complete the Basic course and the appropriate species specific modules.
2. If you are an IACUC Member you should complete the "Essentials for IACUC Members".
3. Choose the appropriate species specific electives according to your research interests.
Choose all that apply
\square "Working with the IACUC Course" is required if you plan to use lab animals in your work.
\square If you are an IACUC Member you are required to complete the "Essentials for IACUC Members" course now.
☐ Institutional Official: Animal Care and Use
☐ IACUC Community Member
☐ Antibody Production
☐ If you are planning to do aseptic surgery on animals, you may want to complete the "Aseptic Surgery" course now. Your Institution may require this.
☐ If you plan to conduct studies that have the potential to cause "more than momentary pain and distress" in Mice or Rats you should complete the module on "Minimizing Pain and Distress".
Choose the appropriate species specific electives depending on your work or interests.
☐ I work with Mice. Family: Muridae Cricetidae
☐ I work with Rats. Genus: Rattus
☐ I work with Frogs, Toads or other Amphibians
☐ I work with Hamsters. Family: Muridae
☐ I work with Gerbils
☐ I work with Guinea Pigs
☐ Working With Ferrets in Research Settings
☐ I work with Rabbits, Family: Leporidae
☐ I work with Fish
☐ I work with Cats
☐ I work with Dogs
☐ I work with Swine
☐ I work with Non-Human Primates (NHP)
☐ Working With Animals In Biomedical Research - Refresher Course
Question 7
Would you like to take the IRB Chair course?
Choose one answer
O Yes
Not at this time.
Who the trial time.

Question 8

Human Subjects Research

Please choose one learner group below based on your role and the type of human subjects activities you will conduct. You will be enrolled in the Basic Course for that group.

Revised Common Rule

Please make your selection below if you wish to be enrolled in the Revised Common Rule course.

Revised Common Rule

Not at this time.

	Question 9
	Would you like to take the Institutional/Signatory Officials for Human Subjects Research course? Choose one answer Yes Not at this time
	Question 10
	Revised Common Rule Please make your selection below if you wish to be enrolled in the Revised Common Rule course. Choose one answer O Revised Common Rule Not at this time.
	Question 11
	Question 1
	Good Clinical Practice (GCP) Please make the appropriate selection if you are required to complete the Good Clinical Practice (GCP) course. Choose one answer
(GCP for Clinical Trials with Investigational Drugs and Medical Devices (U.S. FDA Focus) GCP for Clinical Investigations of Devices GCP for Clinical Trials with Investigational Drugs and Biologics (ICH Focus) GCP – Social and Behavioral Research Best Practices for Clinical Research Not at this time.

Webinars
Please choose the Webinars you would like to review:
Choose all that apply
☐ Ethics & Policy Issues in CRISPR Gene Editing New Content
☐ The Challenge of Medicare Advantage Plans and Local Coverage Determinations
☐ GDPR & Human Subject Research in the U.S.
☐ FERPA: A Quick Review of the Law for Researchers and IRBs
☐ Preparing for Single IRB (sIRB) under the Common Rule
\square Transitioning Research to the Revised Common Rule: The What, How, and Why
Revised Common Rule: Overview of Revisions
Revised Common Rule: Revisions to Informed Consent
Revised Common Rule: Revisions to Definitions
☐ Building a ClinicalTrials.gov Compliance Program – Tips for Investigators and Institutions
☐ Running a Virtual IRB Meeting
☐ Medical Marijuana: A Budding Field of Research
\square Understanding Consent Requirements and "Key Information" Under the Revised Rule
☐ Artificial Intelligence (AI) and Human Subject Protections
☐ Social Media and Research Recruiting
☐ International Students in Focus at U.S. Higher Education Institutions (HEIs)
☐ Importance of Peer Review and Data Validation in Research
COVID-19: Supporting Ethical Care and Responding to Workforce Concerns in a Public Health Emergency
☐ "Nuts & Bolts" of Running a Virtual IRB Meeting
COVID-19 and Human Research Protection Programs
☐ FERPA and Online Learning in the Time of COVID-19
☐ Informed Consent: A Focus on the Process
☐ Working with Your IRB
☐ Title IX and the New Regulations
Accreditation 101 for New and Adjunct Faculty
Export Compliance: An Overview for Staff, Students, and Faculty
☐ IRB Administrator Professional Development and Self-Advocacy
\square Research with Native American Communities: Important Considerations When Applying Federal Regulation
Health Disparities: Promoting Equity and Diversity in Clinical Research
☐ Informed Consent and Research with Wearable Tech
\square The Playbook: Successfully Developing and Deploying Digital Clinical Measures
☐ Managing Conflict with Your Dissertation Chair
\square Getting Started in Grant Writing: An Introduction for Graduate Students, Postdocs, and New Faculty
Remote Informed Consent: The Same, but Different, but Still the Same
Understanding Decentralized Clinical Trials (DCTs) and Virtual Study Visits
\square Principles and Practices for Managing Undue Foreign Influence in an Academic Environment
☐ Blockchain and Higher Education
☐ GDPR: Top Noncompliance Risks and Mitigation Strategies
Race in Clinical Research: Ethics and IRB Decision Making
☐ Decentralized Clinical Trials (DCTs) and Your Workforce
☐ Intellectual Property and Working With Your Technology Transfer Office
Cost Allowability on Federally Sponsored Awards
Privacy and Ethical Considerations for Connected and Automated Vehicles (CAVs)
\square Leveraging IT Insight in IRB Review: Why Technology-Based Expertise is Critical to Human Subjects
☐ IRB Review of Observational Research
Research with Audio-Visual Mobile Data Collection Tools: Ethics and Regulations
U.S. Department of Defense (DoD) Regulations & Requirements for Human Subject Research
A Beginner's Guide to Being a Sponsor-Investigator
☐ Partnering with Technology Companies
☐ Bring Your Own Device (BYOD) Studies
☐ Data Management and Security for Student Researchers: An Overview
☐ Facial Recognition Considerations for Researchers
\square IRB Protocol Noncompliance: When Research Goes Rogue, What Next?
☐ Managing Your Grant as Systems: A Guide for Grant Management Success

Subject Research
A Beginner's Guide to Being a Sponsor-Investigator
Partnering with Technology Companies
Bring Your Own Device (BYOD) Studies
Data Management and Security for Student Researchers: An Overview
Facial Recognition Considerations for Researchers
IRB Protocol Noncompliance: When Research Goes Rogue, What Next?
Managing Your Grant as Systems: A Guide for Grant Management Success
Not at this time.





Welcome, Heather

Add Institutional Affiliation Register as Independent Learner





Institutional Courses

Institutional Courses are available to learners who have an affiliation with one or more subscribing institutions. If an institution with which you are affiliated is not listed, you may want to add an affiliation. If you are no longer associated with a listed institution, you may want to remove an affiliation.

University of Louisiana at Lafayette	View Courses
Would you like to affiliate with another Institution?	Add Affiliation
Would you like to remove an existing affiliation?	Remove Affiliation

Independent Learner

Register as an independent learner to purchase course content. Before you purchase a course, please make sure you do not already have access to that course through an Institutional affiliation. Please note that refunds are not available for courses purchased as an Independent Learner.

Register as an Independent Learner



University of Louisiana at Lafayette

Active Courses

Learner Tools



Assurance Statement

Basic Human Subjects Research - Basic

CITI Program's <u>Terms of Service</u> and <u>Privacy and Cookie Policy</u> include the following provisions for learners. Please read them carefully.

Account Security: I will keep my username and password secure, and I will not share them or allow anyone else to access my account. I will contact CITI Program Support if I believe my account has been compromised.

Work Integrity: I will complete all required quizzes and any other assessments using only my own work. I will not engage in any activities that would dishonestly improve my results, or improve or hurt the results of other learners.

Quiz Sharing: I will not share CITI Program quiz questions or answers on any website, via email, photocopying, or by any other means.

Recordkeeping: I understand that CITI Program keeps account activity logs, including computer IP address, time spent in each content area, number of quiz attempts, and quiz scores. Indications of inappropriate use will be investigated, and may be reported to organizations with which I am affiliated.

✓ I AGREE to the above, the <u>Terms of Service</u>, and the <u>Privacy and Cookie Policy</u>, in order to access CITI Program materials.



Dominique Rosado ID 6192128

Basic Human Subjects Research

University of Louisiana at Lafayette

- · Complete all 11 required modules
- · Complete 1 of 2 elective modules
- Achieve an average score of at least 80% on all quizzes associated with this course's module requirements



modules complete





Required Modules

Complete all 11 required modules.

Modules	Completed	Score	
Belmont Report and Its Principles (ID 1127)	24-Jan-2020	100%	Review
History and Ethical Principles - SBE (ID 490)	24-Jan-2020	80%	Review
Defining Research with Human Subjects - SBE (ID 491)	Incomplete	-	Start
The Federal Regulations - SBE (ID 502)	Incomplete	-	Start
Assessing Risk - SBE (ID 503)	Incomplete	- [Start
Informed Consent - SBE (ID 504)	Incomplete	- [Start
Privacy and Confidentiality - SBE (ID 505)	Incomplete	- [Start
Populations in Research Requiring Additional Considerations and/or Protections (ID 16680)	Incomplete	- [Start
nternet-Based Research - SBE (ID 510)	Incomplete	- [Start
Social and Behavioral Research (SBR) for Biomedical Researchers (ID 4)	Incomplete	-	Start
Conflicts of Interest in Human Subjects Research (ID 17464)	Incomplete	- [Start