

Research Compliance Program Manager, Office of Research Administration

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# **Instructions for Lifespan's Research Training Courses**

All research personnel are required to complete this training prior to conducting research. Please ask the PI or your direct supervisor to select which courses are required from the table below.

### Go to: http://www.citiprogram.org

### Create an account:

- 1. Click Register
- 2. Under <u>Select Your Organization Affiliation</u>, enter "Lifespan Corporation" and complete the registration steps. *NOTE: You do not have to use a Lifespan email address to affiliate with our training courses.*
- 3. Refer to the table below to select your curriculum

### Already have an account? You will need to add Lifespan to your account:

- 1. Log in to your existing CITI account
- 2. Go to Affiliate with Another Institution, enter "Lifespan Corporation" and complete the registration steps.
- 3. Refer to the table below to select your curriculum

#### Select courses:

- 1. Under Lifespan Corporation Courses, click on Add a course
- 2. Refer to the table below to select your curriculum

#### Complete the required courses:

- 1. To begin the course, click on the course title from the MAIN MENU
- 2. Click on Complete the Integrity Assurance Statement before beginning the course

# Provide documentation:

- 1. Select View-Print-Share Completion Record from the MAIN MENU
- 2. Send a copy of your completion certificates to your supervisor

Select Curriculum - Lifespan Corporation (ID: 628)				
	Check if required:	Course:	Required for:	Frequency:
Q1		Basic Human Subjects Protection (HSP) - includes HIPAA	Anyone who will work on a human subjects research project	Due once prior to conducting any research activity
		HSP for IRB members	Lifespan IRB members, RPO coordinators, RPO administrators	Due once prior to conducting any research activity
		Recombinant DNA and Biosafety	Anyone who will work with recombinant or synthetic nucleic acid materials	Due once prior to conducting any research activity
Q2		HIPAA Annual Renewal	Anyone who will work on a human subjects research project	Due annually; Note: HIPAA is included in the HSP course; therefore, you can skip this course during the years when HSP is completed.
Q3		Refresher/Re-certification in Human Subject Protection (HSP) - includes HIPAA	Anyone working on a human subjects research project	Due every 3 years from when you completed the first HSP course
Q4		Good Clinical Practice (GCP)	NIH-funded clinical trial personnel who will be engaged in the conduct, oversight, or management of clinical trials	Due once prior to conducting any research activity
		Good Cillical Fractice (GCF)	Anyone who will be involved with recruitment, consent, data collection from research subjects, and data management is encouraged to complete this course	
Q5		Refresher in Good Clinical Practice (GCP)	NIH-funded clinical trial personnel engaged in the conduct, oversight, or management of clinical trials	Due every 3 years from when you completed the first GCP course
			Anyone involved with recruitment, consent, data collection from research subjects, and data management is encouraged to complete this course	
Q6		Working with the IACUC	Anyone who will work on an animal research project	Due once prior to conducting any research activity
		Essentials for IACUC Members	Lifespan IACUC members, IACUC coordinators, IACUC administrators	Due once prior to conducting any research activity
		Annual Rodent	Anyone who will work on an animal research project with rodents	Due prior to conducting any research activity and again annually
		Annual Large Animal	Anyone who will work on an animal research project with large animals	Due prior to conducting any research activity and again annually
Q7	Animal-specific courses:  Mice Guinea Pigs Rats Rabbits Swine Zebrafish Cats Post-Care of Mice and Rats	Animal-specific courses:	Optional	Optional and renewal is not required
		☐ Mice ☐ Guinea Pigs		
		Rats Rabbits		
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Q8		Responsible Conduct of Research	Optional	Optional and renewal is not required
Q9		Conflicts of Interest	PI's and any other person who is responsible for the design, conduct or reporting of research (e.g. Co-Investigators and other senior/key personnel)	Due immediately and again every 4 years
Q10		Empty course		
Q11		Clinical Trial Billing Compliance (CTBC)	Optional	Optional and renewal is not required