



Atrium Health Navicent Pre-requisite CITI training for Conduct of Clinical Research

CITI Training Requirements for anyone involved in the Conduct of Clinical Research at Navicent Health in accordance with Atrium Health Office of Clinical and Translational Research requirements (Effective June 1, 2019)

****Note:** The specific modules required are pre-determined based on your area of research involvement using the following categories:

Course	Mode	Participant	Maintenance	
Clinical Research				
CITI Basic Course	Biomedical Research	CITI module	Investigator, Support Staff**	Refresher, every 3 years
	Shipping Transport of Regulated Biologic Material	CITI module	Support Staff**	Refresher, every 2 years
	Social and Behavioral Research	CITI module	Investigator, Support Staff**	Refresher, every 3 years
	Conflict of Interest	CITI module	Investigator, Support Staff**	Every 3 years
	Good Clinical Practice	CITI module	Investigator, Support Staff**	Refresher, every 2 years
	Data Management, Integrity & Security	CITI module	Support Staff**	Every 3 years
	Clinical Trial Billing Compliance (CTBC)	CITI module	Research Managers and Directors, SPA personnel, and others involved in the research budget/billing/coding processes	Every 3 years
Clinical Research Coordinator 101 (includes a Research Billing Compliance Module)	In-Person Class	Clinical Research Coordinators (RN and Non-RN)	One-time Course *Must be completed before participating in clinical research activity	
Biosafety/Biosecurity (select applicable modules – animal, human, recombinant DNA)	CITI module	Investigators, all staff listed on IBC protocol – unless determined exempt by IBC Chair	Annual refresher	
IRB Members	CITI module	IRB Members	Refresher, every 3 years	