## **Navicent Health Institutional Review Board**



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## 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		In-Person	Clinical Research	One-time Course
(includes a Research Billing Compliance Module)		Class	Coordinators (RN and Non-RN)	*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