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Webinar: How to Ensure Study Compliance and Integrity through Training and Delegation of Authority

Date
Tuesday, July 11, 2017 -9:00am to10:00am
Location
Association of Clinical Research Professionals (ACRP)

This webinar is offered for a fee by the Association of Clinical Research Professionals (ACRP). **Learn more and register here.**[1]

Description: According to the FDA and OHRP, the Principal Investigator (PI) is responsible for all study conduct. However, in the day of complex study designs it is impossible for the PI to directly conduct all aspects of a research study. Therefore, qualified and reliable staff must be employed to ensure the study is carried out according to the protocol.

Attendees of this webinar will understand the regulatory requirements of investigator oversight, as well as gain tips for ensuring investigators are properly engaged with delegated duties; study staff entrusted with study conduct are properly trained; and delegation of authority documentation is maintained.

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Links:
[1] https://www.acrpnet.org/courses/webinar-ensure-study-compliance-integrity-training-delegation-authority/