





WEBINAR Applying the FDA Framework in Conducting IRB Review

Zoom Link

Monday, May 15th 1:00 pm - 2:15 pm

Ada Sue Selwitz, MA

Executive Integrity/Compliance Advisor Office of the Vice President for Research & Associate Director, Regulatory Support & Research Ethics,

Center for Clinical and Translational Sciences

Belinda M. Smith, MS, RD, CCRC

Office of Research Integrity, Research Education Specialist

Objectives:

- Examine when an activity falls under FDA regulations
- Outline a step-by-step framework for applying FDA regulations in clinical investigations