Pediatric Research Alliance

Children's | EMORY UNIVERSITY | Georgia Tech | MOREHOUSE SCHOOL OF MEDICINE

Understanding the New NIH Clinical Trial Requirements for Grants and Contracts

Friday, January 12th
12:00pm – 1:00pm
Egleston Classrooms 5-7

Presented by Emory's Office for Clinical Research:

Holly Sommers, BA
Director, Office of Sponsored Programs

Rebecca Rousselle, BA, CIP
Director, Institutional Review Board

Jennifer Prozonic, MPH, CCR, CHRC
Supervisor, Clinical Trial Compliance

Learning Objectives:

1) Determine if your study meets the NIH definition of a clinical trial & the implications of that determination.
2) Determine what NIH Funding Opportunities allow clinical trials & understand the new required paperwork for submission.
3) Determine who needs Good Clinical Practice training and how often.
4) Determine what studies will be required to use a single IRB and what is entailed.
5) Determine what studies are automatically covered under a Certificate of Confidentiality.
6) Determine what studies require ClinicalTrials.gov registration & reporting, & the penalties for noncompliance.

To RSVP: Visit www.pedsresearch.org calendar and select 1/12/18
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