School of Dental Medicine Research News

July 2018
Distributed via email 7/6/2018

Contents:

Χ

ne Training oportunities poortunities

Af) as at the nate of the transfer of the same of the second state of the second secon

ed when transferring protected health information (PHI), including o another.

o known as a Confideat Disclosure Agreement CDA) is used e party to another for review only (no further use or dissemination), ating the potential for a future relationship between the parties.

combined into a single agreement that defines the rights, he providing and receiving parties regarding issues such as use, otects you as agreement and Case from various claims, but it also eing used without your consent or compensation, if warranted

Clinical Research Training Classes

New to clinical research ave questions about what makes a good (and compliant!) research project?

Beginning in August, UH will be presenting a series of sessions titled "The Basics" that discuss different aspect of clinical research. Sign up for all or a few. Topics include:

- x Module 1-FDA Regulations and Good Clinical Practice
- x Module 2-Required Regulatory Documentation and Essential Documents
- x Module 3-Study Feasibility through Study Activation
- x Module 4-Coverage Analysis, Research Billing Compliance, and Subject Calendar
- x Module 5-Prescreening, Eligibility, and Enrollment Process
- x Module 6-Informed Consent and REonsent
- x Module 7-Good Documentation Practices and Clinical Data Management
- x Module 8-Adverse Events and Protocol Deviations
- x Module 9-Avoiding NonCompliane Findings

Training and presentations on additional topics are available as well through UH and CWRU. Please visit https://research.case.edu/researchapps/education/onliatendar.cfmfor further details and registration.

NIH Reporting Updates

Over the past month, the NIH has made some changes that will impact reporting in the RPPR (Research Performance Progress Report) and the FRPPR (Final Research Performance Progress Re

Χ

X Delegation is now allowed for FRPPRs. When the NIH institute to the requiring FRPPRs instead of a final progress report upload, the ability to delegate to allow others to work on the FRPPR was not included, meaning only the PI/PD had access to the report. That has now changed in the NIH