School of Dental Medicine Research News

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Clinical Research Training Classes

New to clinical research? Have 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 aspe of clinical research. Sign up for all or a few. Topics include:

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

Training and presentations on additional topics are available as well through UH and CWRU. Please visit https://research.case.edu/researchapps/education/onlinecalendar.cfm for 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 port).

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- Delegation is now allowed for FRPPRs. When the NIH instituted the policy of 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