March 21, 2018

ClinicalTrials.gov studies reassigned effective April 1, 2018

Effective April 1, 2018, the registration and results submission for ClinicalTrials.gov studies has been reassigned to the following departments:

Revised IRB Forms

The IRB forms below have been revised. Since forms and guidance may be updated at any time to reflect new regulations and IRB accreditation, be sure to download the latest versions as needed from the IRB Toolkit for Investigators to avoid delays...

REMINDER of COI for IRB Process

Conflict of Interest disclosure and training must be completed for all research personnel responsible for the design, conduct or reporting of a study prior to the study being reviewed, per federal requirements.

Are original signatures always required on forms submitted through Research Affairs?

Signatures of Principal Investigators and their Department Chairs are frequently required on submissions to Research Affairs (IRB, IACUC, grants, etc.).