WHAT YOU NEED TO KNOW ABOUT CLINICALTRIALS.GOV AND THE NEW COMMON RULE

Who?
Any Investigator who:
- is also the Sponsor (Investigator-initiated)
  AND
- has a study that meets the requirements to register a study on ClinicalTrials.gov

When?
Any clinical trial approved by the IRB on or after January 21, 2019

What?
Upload an IRB-approved consent form (ICF) to ClinicalTrials.gov anytime after recruitment is closed, but no later than 60 days after the last subject visit

New Requirement: Posting Clinical Trial Consent Form

What else?
- Only one version of the ICF needs to be posted per study
- Redaction of proprietary or sensitive information is allowed

For questions or additional information, please click here https://www.utmb.edu/research/home/office-of-clinical-research/clinicaltrials-gov-resources or call the Office of Clinical Research at 409-772-1985