Attention Device Trial Investigators! UAB has a new procedures document which enables device trial PIs to directly sign the 1572-like investigator agreement (IA) provided by the sponsor and incorporate the signed document into the regulatory binder for the study. If the sponsor’s IA refers to the Declaration of Helsinki, please contact Adam McClintock in the Office of the IRB.

Please review the process document carefully to understand your responsibilities.

Do you plan to engage in Data Coordinating Center (DCC) activities? If so, this document is intended for you! UAB has launched a new Assessment Group through the Clinical Trials Administrative Committee (CTAC) to review these DCC opportunities to ensure their purpose aligns with the mission of the University relative to scholarship and advancement of knowledge.

Be sure to review the Charter thoroughly and reach out to Drs. Kimberly or Nichols (co-chairs) with any questions.

Did You Know? The CITP on the Go podcast now includes five brief video ‘how-to’ guides on Clinicaltrials.gov.

Check them out here (scroll down the page).

Thank you for reading,
UAB Clinical Trials Administration Committee