Human Research News

July 27, 2020

1) Updated Guidance:
   - Right to Try Policy and Guidance
   - Right to Try Consent Form template
   - Funding Considerations for Federally-Funded and Industry Sponsored Human Research
   - Genetics Research
   - Deception or Incomplete Disclosure
   - Use of Devices in Clinical Research and Treatment
   - Consent Form Checklist for Reliance on External IRBs
   - Steps to Rely on Another IRB – Flow Chart
   - Relying Investigator Responsibility Checklist
   - When UCLA Relies information on ancillary reviews, amendments, and continuing reviews
   - Steps to serve as the Reviewing IRB – Flow Chart
   - Reviewing Investigator Responsibility Checklist
   - NCI CIRB reliance information on ancillary reviews, amendments, and continuing reviews
   - NCI CIRB Boilerplate Consent Language Checklist
   - Family Educational Rights and Privacy Act (FERPA)
   - Brief Overview of webIRB Submission Procedures
   - Tip Sheet: Minimal Risk
   - Closure of Human Subjects Research studies
   - Communication of Results of IRB Review
   - Investigator Responsibilities

2) OHRPP’s “Learn at Lunch”

   Topic: “Access Aggregate Counts (i2b2, LADR, & ACT) and Patient-level Data for Research through UCLA CTSI Informatics Program”
   Presenters: Carina V. Hampp, & Amando Do, Informatics Program, UCLA CTSI

   Date: July 30, 2020
   Time: noon-1pm
   Location: zoom (Register for this meeting)

3) OHRPP’s Office Hours:

   ➢ Bring your questions (“how do I get started applying to the IRB?”, “Can you help me better understand this guidance document?”, etc…) and we’ll do our best to get you answers.
Sessions are *every other Tuesday morning*

Register once to join any session:

Upcoming Office Hours:
  - August 4 at 8:30am

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