Making IRB easier: Flexibilities for Informed Consent

Linda Halstead, MA - Director of Research Protection Programs

Tuesday, January 10, 2017
12pm-1pm

- What are the different kinds of consent waivers and when can they be used?
- What is the difference between a waiver of consent and waiver of consent documentation?
- When is a waiver of HIPAA Authorization possible?
- Does it make a difference if the study is reviewed by Full Board or Expedited Review?
- Is consent necessary for Exempt studies?
- What is involved in getting a waiver approved by the IRB?

For the Research Affairs noontime Seminar Program
Research Affairs Main Conference Room
24887 Taylor Street, Suite 201P
***For lunch and credit you**
must register in OWL

Click link to OWL Register

Linda Halstead

Linda G. Halstead, M.A., is Director of Loma Linda University's Research Protection Programs, and serves as IRB Administrator. Having founded LLU's pre-award research service for the University in 1975, she has been responsible for implementing institutional policies and federal regulations for human subject protections at Loma Linda since the early days of the U.S. Office for Protection from Research Risks (now known as the Office for Human Research Protections -- OHRP). Ms. Halstead served as the President of the international Society of Research Administrators in 1983-84.

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