Inquisitive Thoughts...Do Ethics or Protections Ever Need Deliberation or Specification? IT DEPENDS!

A One-day Educational Workshop Co-Sponsored by The Office for Human Research Protections (OHRP), St. John Medical Center, and the Oklahoma Medical Research Foundation (OMRF)

October 31, 2019 8AM - 4:45PM

St. John Medical Center, Mary K Chapman Health Plaza, Robinson Lafortune Room, Tulsa, OK with simulcast at the Oklahoma Medical Research Foundation's Wileman Center, Oklahoma City, OK

TIME	DURATION	SESSION
8:00 – 8:30	30 min	Registration and Breakfast
8:30 – 8:40	10 min	Welcome and Introductions (Organizers & OHRP)
8:40 – 9:55	75 min	Overview of the Regulations and How to Apply Them Misti Ault Anderson, OHRP
9:55 – 10:50	55 minutes	Evidence Based Practice (EBP) and Research Under the Common Rule** Stacie Merritt, St. John Medical Center Jaime O. Hernandez, OHRP
10:50-11:00	10 min	Break
11:00 – 12:00	60 minutes	The Subject Perspective: A Panel Discussion** Stacie Merritt, St. John Medical Center
12:00 – 13:00	60 min	Networking lunch
13:00 – 14:00	60 min	Secondary Research with Data, Biospecimens, and Private Information Jaime O. Hernandez, OHRP
14:00 – 14:10	10 min	Break
14:10 – 15:10	60 min	Critical Review of a Clinical Research Protocol ** Yvonne Lau, OHRP Misti Ault Anderson, OHRP
15:10 – 16:15	65 min	Informed Consent for Clinical Trials: Putting it into Practice** Yvonne Lau, OHRP
16:15 – 16:30	15 min	Closing Remarks & Final Questions

Food and beverages provided courtesy of Complion, St. John Medical Center, and the Oklahoma Medical Research Foundation. OHRP funding does not support this portion of the event.

^{**}Workshop sessions that are marked with a double asterisk (**) on the agenda meet the criteria in the Certified IRB Professional (CIP) recertification guidelines at www.primr.org/Subpage.aspx?id=1579. These sessions are eligible as accredited continuing education units for CIP and SOCRA accreditation. A maximum of 4.0 hours of continuing education credits can be claimed.