OSP Roundtable

Thursday, December 14, 2017
9:30 - 10:30 AM
Plant Science Building, Room 404

New NIH Human Subjects Requirements

Presenters

Guilaine Senecal, Assistant Director, Office of Research Integrity and Assurance
Jamie Sprague, Sr. Grant & Contract Officer/Federal Team Lead, Office of Sponsored Programs

Earlier this fall, the National Institutes of Health (NIH) issued new compliance requirements for NIH-funded studies with human participants. These changes to the definition of clinical trials, use of a single institutional review board (sIRB), and certificates of confidentiality policies will affect the submission of proposals, management of awards, and completion of human participant review and compliance at Cornell.

Join us in two weeks to learn about these areas of human subjects research compliance and how Cornell researchers and research administrators will need to respond to these new requirements. **Special attention will be paid to the new PHS Human Subjects and Clinical Trials Information form required of all proposals submitted to NIH, the Centers for Disease Control and Prevention (CDC), the Food and Drug Administration (FDA), and the Agency for Healthcare Research and Quality on or after January 25, 2018.**

Feel free to send me your questions about NIH's new human subjects requirements, including the new PHS Human Subjects and Clinical Trials Information form prior to the session, so I can provide them to our presenters in advance of the Roundtable.

We look forward to seeing you in two weeks on Thursday, December 14th!

Thank you,
Christine