Upcoming Education Opportunities

There are upcoming educational opportunities related to the myIRB submission process. Please join me on the 2 December at 9 a.m. to be educated on the myIRB revision submission process. The training will be in room HSC C1-7 of the Communicore Building.

One week later, I will be presenting on the Continuing Review Submission process on the 9th of December at 10 a.m. in room HSC C1-7 of the Communicore Building.

There is no need to RSVP. Each session should last less than ninety minutes.

On the 11 December from noon—1:30, Dr. Wajeeh Bajwa will be presenting the December Brown Bag on “Oversight of Clinical Investigations.” The objectives are:

For participants to better understand compliance requirements at UF.

For participants to better understand the challenges in fulfilling compliance requirements.

For participants to learn about resources/tools available to achieve compliance.

This is our monthly education session. CME and CEU credits are available for attendees.

Please RSVP for this event to tiffany.danielle@ufl.edu.

Office Closures

IRB-01 and IRB-02 Main Offices will be closed on Thursday November 28 and Friday November 29 in accordance with the University of Florida holiday calendar.

You may still submit to IRB-01 electronically via myIRB. IRB-02 is not currently accepting myIRB submissions.

Both offices will reopen on Monday December 2, 2013.

What is noncompliance in human subjects research?

Failure to follow federal regulations; state and local laws; institutional policies governing human subject research; or the requirements or determinations of the Institutional Review Board. This can include failure to follow the VHA handbook for VA regulated research.

Examples of noncompliance:

- Failure to obtain IRB approval prior to conducting human subjects research.
- Continuation of research activities after a study has expired.
- Failure to obtain informed consent of research subjects.
- Failure to follow the Protocol as approved by the IRB.