PUBLIC RESPONSIBILITY IN MEDICINE AND RESEARCH

Putting Together the HRPP Puzzle

FACULTY

Name, Designation
Title
Institution

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Research compliance is an institution-wide undertaking. From investigators and research coordinators to institutional review board members and staff to institutional officials—all members of an institution's human research protections program (HRPP) are key to ensuring high quality, ethical research, and clinical trials. This intensive, half-day version of the program will provide participants with the tools they need to be effective members of their HRPP by highlighting best practices and common oversight challenges. The course is designed for all members of an institution's HRPP but places special emphasis on addressing the role and function of researchers and research staff in relation to the institutional review board.

After attending this course, you will be able to:

- Define various aspects of an HRPP and how those aspects interact with each other
- Understand how the HRPP fits into a comprehensive compliance program
- Identify benefits of being involved in a larger institutional program
- Identify common compliance challenges and collaboratively discuss solutions

Agenda

7:30-8:30 AM	*Registration*
8:30-8:35 AM	**Welcome and Introduction**
8:35-9:15 AM	**Understanding Your Institution's HRPP and How it Ensures High-Quality, Ethical Research** **Putting it Together: Defining the Pieces that Make Up the HRPP and Why They Are Important**
9:15-9:45 AM	**Interactions and Challenges: How Does Your Office/Position Interact With Each Part of the HRPP, and What are the Obstacles and Potential Solutions?**
9:45-10:00 AM	*Break*
10:00-11:30 AM	**Interactions and Challenges (*Continued*)**
11:30–12:00 PM	**Wrap Up Discussion**
12:00 PM	*Adjournment*

Please note, this agenda is subject to change.