



IACUC THREE I's	
IBC	
IRB BIOSECURITY RA	
RI COMPLIANCE REGULATORY	

THREE I's: and RESEARCH INTEGRITY™: COMPLIANCE, ETHICS and BIOSECURITY ... Building Resilience

Day 3 **WEDNESDAY, APRIL 29, 2026**

DRAFT CONFERENCE AGENDA

	WELCOME TO DAY 3		
10:00 AM - 10:45 AM	BREAKOUT SESSIONS		
	<p style="text-align: center;">IACUC</p> <p style="text-align: center;">COMPLIANCE USE STANDARD PROCEDURE (CUSP)</p> <p style="text-align: center;">CUSP - THE FEDERAL DEMONSTRATION PARTNERSHIP</p> <p style="text-align: center;">PANEL</p>	<p style="text-align: center;">IRB</p> <p style="text-align: center;">PREVENTING AND IDENTIFYING FRAUDULENT RESPONDENTS IN ONLINE SURVEY RESEARCH</p> <p style="text-align: center;">JULIE SIMPSON, PhD DIRECTOR RESEARCH INTEGRITY SERVICES UNIVERSITY of NEW HAMPSHIRE</p> <p>Researchers have used the internet to conduct research for decades. More recently, however, online research has become a target for fraudulent respondents, particularly when researchers offer monetary incentives for participation in a study. This session will explore the current environment for online data collection from human participants, outline strategies for researchers and Institutional Review Boards (IRBs) to help minimize and identify fraudulent responses and protect data integrity and examine the challenges of balancing data integrity with participant welfare.</p>	<p style="text-align: center;">COMPLIANCE</p> <p style="text-align: center;">RESEARCH SECURITY: EVOLVING RISK AND REGULATION</p> <p style="text-align: center;">MINAL MICHAEL CARON, COUNSEL ROPES & GRAY LLP</p> <p>This presentation offers a comprehensive overview of the rapidly evolving U.S. research security landscape, focusing on recent regulatory changes, enforcement actions, and compliance challenges affecting federally funded research institutions. Attendees will learn about the status of key "mandates" such as NSPM-33, the CHIPS and Science Act, and agency-specific requirements for disclosure, cybersecurity, and research security training. The session will provide practical recommendations to help research compliance and administration professionals strengthen institutional research security programs and navigate heightened federal scrutiny.</p>

10:45 AM – 11:45 AM	GENERAL SESSION		
	<p>ALL I'S RESEARCH INTEGRITY & RESEARCH ADMINISTRATION INVITED!</p> <p>SPOT THE ISSUES ...ETHICS CHALLENGE</p> <p>KATHRYN A HOLTHAUS, MS, MA DIRECTOR OF RESEARCH SUBJECTS PROTECTION AND LABORATORY SAFETY COMPLIANCE RESEARCH OPERATIONS BRIGHAM & WOMEN'S HOSPITAL</p> <p>TED MYATT, ScD ASSOCIATE VICE PROVOST OF RESEARCH INTEGRITY TUFTS UNIVERSITY</p> <p>ROSS HICKEY, JD CIP CPIA ASSISTANT PROVOST FOR RESEARCH INTEGRITY UNIVERSITY OF SOUTHERN MAINE DIRECTOR OF THE MAINE REGULATORY ETHICS AND TRAINING CENTER (MERTEC) AT USM</p> <p>FBI</p>		
11:45 AM – 11:55 AM	BREAK		
11:55 AM – 12:40 PM			
	<p>IACUC</p>	<p>IBC</p> <p>SUMMARY OF THE NIH BIOSAFETY MODERNIZATION INITIATIVE</p> <p>JAMES W. KLENNER, MSC, MPH, MPA, RBP, CBSP ASSOCIATE DIRECTOR, OFFICE OF RESEARCH INTEGRITY BIOSAFETY OFFICER BALL STATE UNIVERSITY</p> <p>This presentation will discuss the development of the NIH initiative to strengthen biosafety policies, practices, and oversight with an emphasis on the evolving risks related to future advancements in science and technology. The goal is to develop a more effective, transparent, and modern biosafety policy.</p>	<p>IRB</p> <p>NAVIGATING THE ETHICAL REVIEW OF PARTICIPATORY AND COMMUNITY-ENGAGED RESEARCH</p> <p>JOSH MANGIN RESEARCH COMPLIANCE TRAINING MANAGER, UNIVERSITY OF NEW ENGLAND</p> <p>Participatory Research (PR) and Community-Engaged Research (CErR) models fundamentally shift the relationship between researcher and participant, moving toward equitable partnership and shared decision-making. This shift creates unique ethical considerations that challenge traditional Institutional Review Board (IRB) review processes, particularly concerning power dynamics, informed consent, and data ownership. This session is designed to provide IRB members, administrators, and Human Research Protection Program (HRPP) staff with</p>

			<p>practical guidance and frameworks for the robust and responsible review of PR/CEnR protocols. This session will explore key elements such as defining PR and CEnR, discussing consent processes, and navigating complexities in data sharing and publication. The goal is to equip reviewers with the tools necessary to facilitate ethical research that respects community autonomy while adhering to regulatory standards.</p>
12:45 PM – 1:15 PM	LUNCH		
1:15 PM – 2:00 PM			
	TBA		