

OHRP Research Community Forum

Agenda

This will be a two day event full with keynote presentations, interactive workshops, and exhibits, plus lots of opportunities for professional networking.

> Click here for session descriptions

Day 1: Applied / Worksh	p Day - Se	ptember 6, 2017
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Time	Group A	Group B	
7:00-7:30	Breakfast and Registration		
7:30–9:00	Group A: Brief Welcome "When Does my Protocol Need IRB Review: Impact of OHRP Regulations on IRB Review"		
9:00-9:15	Break (15 min)	Group B: Networking Exhibitor review	
9:15–11:15	Group A: "Thinking through Case Studies: Bridging the Gap between Study Design & IRB Review"		
11:15–11:30	Break (15 min)		
11:30-12:30	Plenary Presentation: Michele Russell-Einhorn "Consent in 2017: A New Landscape"		
12:30-1:15	Lunch		
1:15–2:15	"I Thought You'd Never AskA Conversation with IRB Chairs" Local Panel Terry Malone, Bob Frenck, Mike Linke, and Sara Harnish		
2:15-2:30	Group A:	Break (15 min)	
2:30-3:45	Evaluation and Transition to Networking Exhibitor review	Group B: Brief Welcome "When Does my Protocol Need IRB Review: Impact of OHRP Regulations on IRB Review"	
3:45-4:00		Break (15 min)	

Time	Group A	Group B
4:00-6:00		"Thinking through Case Studies: Bridging the Gap between Study Design & IRB Review"
6:00-6:15		Evaluation & Day 1 Closing Remarks

Day 2: Theoretical / Conference Day – September 7, 2017				
Time	Track 1 (ethics / regulatory)	Track 2 (community / CBPR)	Track 3 (novel / technology / mobile)	
8:00-9:00	Registration and Breakfast			
9:00-9:15	Welcome and Opening Remarks			
9:15–10:15	Keynote Address: "Jazz, Blues, and Improvisation - the Changing Rhythm of Ethical Clinical Research" - Christine Grady			
10:15–10:30	Break			
10:30-11:15	Plenary 1: Dr. Jerry Menikoff – "Overview of the 2018 Changes to the Common Rule"			
11:15–12:15	Plenary 2: Joe Konstan – "Improvised Exemptions: A Soloist's Perspective"			
12:15–1:00	Lunch			
1:00-2:00	"Investigator Responsibilities: Striking a Chord for Compliant Innovation" – Stuart Goldstein and Jonathan Thompson	"Learning Health Networks - Harmonic Collaborations of Patients, Clinicians, Researchers and Health Systems" – Peter Margolis	"Research Consent Using Telemedicine: Acute Stroke as a Case Study" – Opeolu Adeoye	
2:00-2:15	Break			
2:15–3:15	"Ethical Considerations of Working with Drug-Using Patients in Clinical Research" – Bill Stoops	"Strategies for Rapid and Robust Ethics Review of Disaster Research" – Angela Morley	"Conducting Human Subjects Research in a Basic Research Department" – Jason Heikenfeld	
3:15–3:30	Break			
3:30-4:30	"Why does it have to be So Difficult? Let's Just Measure 'Understanding'" - Rick Ittenbach and Jeremy Corsmo	"Human subject protection: perspectives from a community-based participatory researcher" – Nancy Schoenberg	"From Improvisational Riffs to Symphonic Harmonies: TeleICU Research - Searching for Melodies in Chaotic Cacophonous Critical Care" – Ralph Panos	
4:30-5:00	Ask the Feds			

5:00-5:15

Evaluation & Closing Remarks