

Study Coordinator Leadership: From Theory to Action

Thursday, January 30, 2020 8:00am-4:00pm UW Medicine South Lake Union Seattle, WA

Conference Schedule

8:00-8:30am	Breakfast & Check-In	C-Building Lobby
8:30-8:45am	 Welcome & Introduction to the Day Amy Good, Implementation Manager, UW Research Coordination Center/Clinical Trials Office Alana Clark, Regulatory Manager, UW Research Coordination Center/Clinical Trials Office 	Orin Smith Auditorium
8:45-9:25am	 Keynote Session – Walk the Path: Tips & Tricks Aimee Schantz, Research Coordinator, UW Department of Pediatrics 	Orin Smith Auditorium
9:25-9:35am	Break	
9:35-10:35am	BREAKOUT SESSION 1	
Option 1A	Ensure a Successful Trial: Things to Think About When Establishing Timeline and Feasibility Assessments for Your PI and Research Team • Lara Schiff, Research Implementation Manager, UW	C-123AB
	Establishing Timeline and Feasibility Assessments for Your PI and Research Team	C-123AB E-130AB
Option 1B	Establishing Timeline and Feasibility Assessments for Your PI and Research Team • Lara Schiff, Research Implementation Manager, UW A Single IRB: The Promise and the Reality • Adrienne Meyer, Assistant Director of Reliances, UW	

10:45-11:45am	BREAKOUT SESSION 2	
Option 2A	 Inclusiveness from Theory to Application Kelly Shipman, Clinical Research Associate, Treuman Katz Center for Pediatric Bioethics, Seattle Children's 	E-130AB
Option 2B	 Placebos, Randomization, and Financial Incentives – oh my! Balancing Ethical Considerations in Clinical Research Kathryn Porter, Research Scientist, Seattle Children's Research Institute Stephanie Kraft, Associate Professor, Seattle Children's Research Institute 	Orin Smith Auditorium
Option 2C	 HIPAA & Scenarios for Critical Thinking Shelia Ganti, Research Coordinator, Seattle Children's 	F-106
11:45am-12:50pm	Networking Lunch	E-130AB
12:50-1:50pm	BREAKOUT SESSION 3	
Option 3A	 Quality Improvement Culture: How to Continue Evaluation Through the Plan Do Check Adjust Approach Cody Blankenship, Regulatory Project Manager, Oregon Health Sciences University 	Orin Smith Auditorium
Option 3B	 Regulatory Year in Review Karen Moe, Director, Human Subjects Division, Asst. Vice Provost for Research, University of Washington 	E-130AB
Option 3C	 All These Dang Rules: Why Do We Have to Follow Them? Eli Reis, Communications & Training, Clinical Trials Office, University of Washington 	F-106
1:50-2:00pm	Break	
2:00-2:50pm	 How to Talk to Your PI About Good Clinical Practices Tara Bumgarner, Clinical Research Coordinator, Seattle Children's Cody Hammer, Clinical Research Manager, UW Hematology Division 	Orin Smith Auditorium
3:00-4:00pm	Post-Conference Networking Hour and Raffle Drawing	C-Building Lounge







