

# Human Subjects Research Regional Conference October 20, 2008

**Hotel Murano  
1320 Broadway Plaza - Tacoma, WA**

Time	Topic and Speaker(s)	
7:00 am – 8:00 am	Institutional Officials Private Meeting	
7:30 am – 8:15 am	Breakfast and Registration	
8:15 am – 8:20 am	<p style="text-align: center;"><b>Welcome</b></p> Royce Morrison, MD; President, Board of Directors, Northwest Association for Biomedical Research Director, Clinical Strategy, Charles River Clinical Services Northwest	
8:20 am – 9:45 am	<p style="text-align: center;"><b>Improving the Effectiveness of Clinical Research</b></p> H. Cliff Lane, MD; Chair, Barriers to Clinical Research Steering Committee Clinical Director, National Institute of Allergy and Infectious Disease, National Institutes of Health	
9:45 am – 10:00 am	Break	
10:00 am – 11:30 pm	<p style="text-align: center;"><b>Track 1: Basic IRB Fundamentals</b></p>	<p style="text-align: center;"><b>Track 2: Advanced IRB Fundamentals</b></p>
	<p style="text-align: center;"><b>Consent Process Risk Benefit Analysis</b></p> Charlotte Shupert, PhD, CIP; Associate Director, Research Integrity Office, Oregon Health & Science University	<p style="text-align: center;"><b>Post Approval Monitoring</b></p> Lynette Schenkel, Assistant Vice President, Office for Responsible Conduct of Research, University of Oregon
11:30 pm – 12:45 pm	Lunch	
	<p style="text-align: center;"><b>Assessing Research Risk</b></p> Annette Rid, MD; Post-doctoral Fellow, Department of Bioethics National Institutes of Health	
12:45 pm – 2:00 pm Breakout session 1	<p style="text-align: center;"><b>Working with Prisoners</b></p> Lynette Schenkel, Assistant Vice President, Office for Responsible Conduct of Research, University of Oregon And Deanne Unruh, PhD; Research Associate, Department of Special Education and Clinical Sciences, College of Education University of Oregon	<p style="text-align: center;"><b>Cognitive Impairment Roundtable</b></p> David Forster, JD, MA, CIP; Vice President, Compliance, Western Institutional Review Board

2:00 pm – 2:15 pm	Break	
2:15 pm – 3:30 pm Breakout session 2	<p><b>Is it research? Is it human subjects research? Is it exempt or expeditable or do we need to take it to the board?</b></p> <p>Charlotte Shupert, PhD, CIP; Associate Director, Research Integrity Office, Oregon Health &amp; Science University And Estela Hamblen, MHA; Manager, Institutional Review Office, Swedish Medical Center</p>	<p><b>Tissue and Data Repositories Roundtable</b></p> <p>Kathryn Schuff, MD; Associate Professor, Endocrinology, Director, Regulatory Support Services, Oregon Clinical and Translational Research Institute, Oregon Health &amp; Science University And Malia “Stephanie” Fullerton, PhD; Assistant Professor, Department of Medical History and Ethics, University of Washington School of Medicine And Kara Manning Drolet, PhD; IRB Co-Chair, Research Integrity Office, Oregon Health &amp; Science University</p>
3:30 pm – 3:45 pm	Break	
3:45 pm – 4:45 pm	<p><b>From Washington to Washington: A Coast-to-Coast Update from OHRP</b></p> <p>Elyse Summers, JD Acting Director, Division of Education and Development Office for Human Research Protections</p>	
4:45 pm – 5:00 pm	<p><b>Conclusion</b></p> <p>Royce Morrison, MD; President, Board of Directors, Northwest Association for Biomedical Research Director, Clinical Strategy, Charles River Clinical Services Northwest</p>	
5:00 pm	Adjourn	
Evening	<p>Association of Clinical Pharmacology Units Annual Meeting Welcome Reception</p> <p>All conference attendees are invited</p>	