

HCCA's Research Compliance Conference April 21-24, 2010 Dallas, TX

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Pre-Conference		
		RP1: (2 CDs - \$20) Research Compliance 101 – Kevin Eskew, Fred Herman
		RP2: (2 CDs - \$20) FDA's New Enforcement Agenda: What It Means to Clinical Investigators and IRBs – Rachel Nosowsky, Rick Robinson
		RP3: (2 CDs - \$20) Faculty Training for Research Compliance Professionals – Angelique Dorsey
		RP4: (2 CDs - \$20) Raising the Bar on Adequate Human Subject Protection – Cynthia Gates, Jeffrey A. Cooper, MD
General Sessions		
		RGS1: Opening Remarks / 2010 Update from OHRP and ORI – Jo An Rochez, Laura Odwazny
		RGS2: International Research – Dr. Melody Lin
		RGS3: Opening Remarks / Research Compliance: A Year in Review – Lisa Murtha, Kendra Dimond
		RGS4: An FDA Report: Physician Initiated Device Studies at Academic Medical Centers – Anne T. Hawthorn
Concurrent Session I		
		R102: Case Studies in Clinical Research Fraud Enforcement – Jesse Witten, Gary W. Eiland
		R103: Developing a Research Compliance Program in the Context of Corrective Actions to an OHRP Investigation – Michael Roach, Ron Sagritalo
Concurrent Session II		
		R201: Managing Regulatory Compliance for Investigator-Initiated Research – Leah R. Kendall, Thomas Bechert
		R202: Ensuring a Sound, Compliant Animal Care and Use Program in a Changing (and Challenging) Landscape – Kathy Wadsworth
		R203: Data Security in Research: Is the IRB Responsible? – Russell Opland, Marian Hughlett

MP3	CD	Session Name
Concurrent Session III		
		R301: Conflicts of Interest in Research: Ethical, Regulatory, and Practical Considerations – Suzanne M. Rivera, PhD., Ann N. James, PhD.
		R302: Developing an Effective Anti-Bribery and Corruption Compliance Program in an Environment of Heightened Enforcement – Jay Perlman, Joel Rush
		R303: The Learnings of a Developing Clinical Trials Office Within an Established Teaching Hospital – Eve Sakran, Ljudmila Hadzikadunic
Concurrent Session IV		
		R401: Compliance Challenges in Establishing and Using Clinical Databases – Melissa (Lisa) Thompson, Betsy Hall
		R402: Unanticipated Problems in Human Subject Research: Adverse Events and Beyond – Keren Dunn, Andra M. Popa
		R403: Research Misconduct: Detection and Risk Mitigation Solutions – Susan S. Night, Sheryl Tatar Dacso
Concurrent Session V		
		R501: Managing Export Control Compliance in Biomedical Research – Don Rischer
		R502: Off-Label Use vs. Clinical Trial Use of Devices: FDA Regulatory Issues – Neil O'Flaherty
		R503: The Ten Steps to an Effective Research Compliance Program: A Practicum for Research Compliance Professionals – Luanna Putney, Juliann Tenney
Saturday, Post-Conference Sessions		
		RW1: (2 CDs - \$20) Research Risks: What's the Assessment? – Margaret Hambleton, John E. Steiner Jr.
		RW2: (2 CDs - \$20) The Closer: Resolve the Risky Business of Billing Compliance – Kathleen Hurtado, Kelly Willenberg

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