

ACRP 2011 Webinar Listing



	DATE	SPEAKER	TITLE
	2011 WEBINARS		
01	Friday January 14	Barbara Davis, David Montgomery, Lynn Van Dermark	2011 Global Conference Speakers Orientation: Best Practices (FREE)
02	Wednesday January 19	Barbara Davis	If They Don't Get It, You Don't Got It: Informed Consent
03	Wednesday January 26	Brian Martens	Recruiting for Difficult Studies on Small Budgets: Lessons Learned from Bipolar Studies
04	Wednesday February 16	Eunice Franklin-Becker and Michael Hill	Is Your Site Ready for Research?
05	Wednesday February 23	Veronique Brisson	Facing the Challenge: Informed Consent in Vulnerable Populations
06	Wednesday March 2	Esther Daemen	From CRA Training to CRA Performance in a Global Context
07	Wednesday March 9	Esther Daemen	Maximizing Training Value with Cost-Effective Job Aids and Checklists
08	Wednesday March 23	Veronique Brisson	Staying Ahead of the Curve: Avoid Being Caught Off-Guard by GCP Audits
09	Wednesday April 6	Rosemary Truman	Using Best in Class Clinical Trial Development Practices to Achieve Breakthrough Commercialization
10	Wednesday April 27	Carol Bogner	Seven Mistakes Commonly Made During a Monitoring Visit
11	Wednesday May 11	Carmen Gonzalez	Latino Patient Recruitment: A Tailored Approach for Maximizing Enrollment
12	Saturday May 14	Nancy Thomas	Item Writers Workshop: Item Writing Training (FREE)
13	Friday May 20	Shay Brill and Bridget-Anne Kirwan	How to Submit a 2012 Global Conference Proposal (FREE)
14	Wednesday June 1	Christine Drabick	Avoiding Common Mistakes in Clinical Research: An FDA Perspective
15	Friday June 3	Jennifer Witebsky	Certification: Navigating the Application Process and the Exam (FREE)
16	Wednesday June 8	Stephanie Christopher	Learning to Communicate Effectively in Multidisciplinary Teams
17	Wednesday June 15	Sandra Chase	Global Recruitment & Retention: Indispensable Strategies for Reducing Clinical Trial Timelines
18	Wednesday July 13	Marjorie Shulman	FDA Premarket Notification (510(k)) Update
19	Wednesday August 10	Tim Davis	The ePRO Choice: Understanding the Impact of Recent FDA Guidance on Your ePRO Tool Selection
20	Wednesday August 17	Joseph Kim and Samuel Whitaker	Site Selection, Patient Recruitment, and Patient Stipend Management: Industry Data and Insights
21	Wednesday August 31	Stephanie Dubois	Placebo Response in Psychiatric Clinical Research
22	Wednesday September 7	Kelly Larrabee	Overcoming Sponsor/CRO Partnership Challenges

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	DATE	SPEAKER	TITLE
23	Tuesday September 20	Mukesh Kumar	September 2011 FDA Safety Reporting Requirements for Clinical Trials: Tips for Compliance
24	Wednesday October 5	Rodd Schlerf	Standard Digital Signatures and How They Can Enhance Clinical Operations
25	Wednesday October 19	Brenda Reese & Arthur Czech	Reduce or Eliminate Changes of Scope: The Clinical Trial Budget Secret
26	Wednesday November 2	Dr. Michael Drues	Pharmacogenomics: The Future of Clinical Trials, New Product Development, and the Practice of Medicine
27	Wednesday November 9	Stuart Horowitz & Jeffrey Cooper	Research Misconduct: Lessons Learned and Practical Approaches to Problems
28	Wednesday November 16	Dr. Nagina Parmar	Ethical Requirements & Development Guidelines for Conducting Pediatric Trials in Developing Countries
29	Wednesday November 30	Dr. Michael Drues	Combination Products: An Overview of the Advantages & Challenges for Clinical Trial Professionals
30	Wednesday December 7	Jeffrey Zigler and Christopher Gingras	Challenges in Clinical Trials: Coverage, Reimbursement, and Compliance Considerations
31	Tuesday December 13	Mukesh Kumar	September 2011 FDA Safety Reporting Requirements for Clinical Trials: Tips for Compliance
32	Wednesday December 14	Robert Romanchuk	Vulnerable Subjects: What the Regulations Don't Say