



## ACRP Webinar Titles 2009

- The Massachusetts Gift Ban Law
- 2009 Global Conference Speaker Orientation
- Health Canada – Regulatory Review
- Managing Up How Study Coordinators Get the Best from Their Principal Investigators
- Financial Management for Clinical Trials: What Every Study Coordinator Should Know – PART I
- Financial Management for Clinical Trials: What Every Study Coordinator Should Know – PART II
- Strategies for Increasing Ethnic Diversity in Clinical Drug Trials
- How to Write a Top-Notch Proposal for the ACRP Global Conference
- Health Canada – Clinical Trial Inspections
- Advanced Analysis of Electronic Healthcare Data to Address Trial Design, Site Selection and Recruitment
- Investigator Responsibility: Essential for Drug Safety and Study Integrity
- The Declaration of Helsinki
- The Informed Consent Process: Achieving Its Purpose or Achieving Our Obligations?
- Optimal Handling of Laboratory Samples - improve Your data Quality: Guidance for Clinical Research
- Too Important To Ignore: How to Market, Promote, and Brand Your Site
- People, Process, and Technology: The Three-Legged Stool of Clinical Research
- The CRO-Site-Sponsor Triangle
- Forecasting Study Success at the Site
- No More Hand-Me-Downs: Research Designed for Children
- Adaptive Clinical Trials: What Do They Offer?
- The Changing Landscape of FDA Expectations: Investigator Supervisory Responsibilities
- ACRP Online Maintenance Made Easy
- Overcoming Recruitment Barriers for Long-Term Studies
- Improving Clinical Data Quality Through Cooperation Between Study Sites and Data Management Groups
- First in Man Clinical Trials of Investigational Medicinal Products (IMPs)
- Monitoring for Quality: Risk-Based Clinical Monitoring