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This webcast program is intended to enhance the knowledge and skills of a clinical researcher in the conduct of clinical trials in order to evaluate a drug’s, biologic’s and/or medical device’s safety and effectiveness in treating, preventing, or diagnosing a specific disease or condition.

This activity was originally presented as a LIVE activity in Seattle, WA on April 30 – May 3, 2011 at the ACRP 2011 Global Conference. If you received credit for a session as a result of attendance in Seattle, you are not eligible to receive credit for that same session within the online version, but you are welcome to use it as a resource. The duration of each presentation is 60-120 minutes. Initial Release Date is June 6, 2011 and Expiration Date is June 6, 2013.

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Learning Objectives:

- Develop strategies and tools for managing the clinical research process such as investigator meetings, site selection, patient recruitment, project management, budgeting at the site, and preparation for regulatory inspections
• Define ethical and regulatory considerations involved in the conduct of research involving human subjects

• Identify implications for global clinical research, such as recruitment efforts within special populations, consenting vulnerable populations and regional regulatory and legal issues
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