



Southern Wisconsin Chapter
Local Chapter Meeting for October 11, 2011

GCP training, Part 2: IRB Responsibilities, Sponsor/Monitoring Responsibilities, Elements of Informed Consent and Beyond, Pediatric Consent and Parental Permission, The importance of SOPs for Clinical Research, and Adverse Event and Deviation Reporting

Tuesday, September 13, 2011

Time: 5:00 pm – Light supper and Chapter meeting
5:30 pm – 7:30 pm Presentation

Speakers: **JoAnn Jessen, MSN**
President, Principal Consultant
Pharma Compliance Partners
Suellen Bigaj, BSN
Vice President, Principal Consultant
Pharma Compliance Partners

Location: Aurora Conference Center
Sycamore Room
2920 West Dakota Street
Milwaukee, WI 53215

AUDIENCE: Clinical research professionals

LEARNING OBJECTIVES:

- Review IRB, sponsor and monitoring responsibilities
- Review of the elements of informed consent and beyond, including pediatric consent and parental permission
- The importance of standard operating procedures for a clinical research site
- Review of adverse event and deviation reporting.

COST: \$10 ACRP MEMBERS who want educational credits, payable online

\$25 CHARGE for CEUs if not a NATIONAL ACRP MEMBER

🕒 Dinner included 🕒 Pre-registration is requested by October 4th 🕒 continuing education contact hours have been applied for through ACRP

In order to receive contact hours, you will be responsible to complete the evaluation form. This form will be posted directly to your record on the ACRP website 10 days following the event and will be available to complete for 30 days.

RSVP to Jennifer Fink, Secretary, ACRP of Southern WI, Jennifer.fink@aurora.org