

Conflicts of Interest in Clinical Research

New financial relationship disclosure requirements for the pharmaceutical and medical device industries are driving a renewed interest in identifying and managing conflicts of interest in clinical research. This session will discuss new regulatory requirements, ways for research sites to identify both investigator and institutional conflicts, and practical methods for managing those conflicts. At the conclusion of this session, participants will understand their own obligations to adhere to regulatory requirements to help manage conflict of issues in their clinical research.

Speakers: Beth Schermer, JD and Kristen Rosati, JD
Coppersmith Schermer & Brockelman PLC
Tuesday March 2, 2010 1:00–4:00 pm

For questions regarding registration, please call AzHHA's Education Department at 602-445-4300 or email edservices@azhha.org.

Early registration deadline is Tuesday, February 23, 2010. Registration made after this date will incur a \$25 late fee.

Cancellation Policy: A \$50 nonrefundable processing fee will be charged for each cancellation. Cancellations must be made in writing five business days prior to the scheduled event. No refunds will be made after this date. If you register and find later that you can not attend, substitutions are permitted and encouraged.

For additional information about AzTransNet, visit: www.azabrc.gov/aztransnet



PHOENIX CHILDREN'S
Hospital

This activity has been planned and implemented in accordance with the Essential Areas and policies of the Accreditation Council for Continuing Medical Education (ACCME) through joint sponsorship of Phoenix Children's Hospital and AzTransNet. Phoenix Children's Hospital is accredited by the Accreditation Council for Continuing Medical Education (ACCME) to sponsor continuing medical education for physicians.

Phoenix Children's Hospital designates this educational activity for a maximum of 3 AMA PRA Category 1 Credits™. Physicians should only claim credit commensurate with the extent of their participation in the activity.

Provider approved by the California Board of Registered Nursing, Provider Number CEP12830 for 3 Contact Hours.



Who should attend?

Clinical research directors, clinical researchers, clinical research coordinators, IRB chairs, IRB coordinators, compliance officers, privacy and security officers, and attorneys will be interested in these sessions.

Continuing Education?

Accredited CMEs, CLEs and CEUs available for up to 3 hours each session.

How much?

Each session is \$125 per attendee for the half-day session, including materials. Attendees that register for all five sessions will get a discounted registration fee of \$500.

Registration?

Online registration is available at <http://events.signup4.com/compliance>

Location?

Phoenix Children's Hospital
1919 E. Thomas Road,
Phoenix, AZ 85016
Melvin L. Cohen Conference Center

Questions?

Contact Julie Robbins
E-mail: robbinsj@battelle.org
Phone: (602) 595-3770
Fax: (602) 595-4760



Compliance Challenges for Clinical Research Sites

An Educational Workshop Series

This workshop series is designed and coordinated by the Arizona Translational Resource Network (AzTransNet) [a project of the Arizona Biomedical Research Commission], Phoenix Children's Hospital and The Arizona Hospital and Healthcare Association

■ Clinical Trials Reimbursement and Payment

Speakers: Julie Nelson, JD and Kristen Rosati, JD, Coppersmith Schermer & Brockelman PLC (CSB) and Ashley Lopez, MS, Director of Research, Phoenix Children's Hospital
■■■ February 9, 2010 ■

■ Clinical Trials Contracting

Speakers: Mayan Tahan, JD and Kristen Rosati, JD, CSB and Jeremy Stoloff, JD, MS, Associate General Counsel, Banner Health
■■■ February 23, 2010 ■

■ Conflicts of Interest in Clinical Research

Speakers: Beth Schermer, JD and Kristen Rosati, JD, CSB
■■■ March 2, 2010 ■

■ Clinical Research Privacy and Security

Speaker: Kristen Rosati, JD, CSB
■■■ April 6, 2010 ■

■ Biospecimens in Clinical Research

Speakers: Kristen Rosati, JD, CSB and Joan Shapiro, PhD, Vice President of Clinical Research, St. Joseph's Hospital and Medical Center
■■■ April 27, 2010 ■

Clinical research sites across Arizona continue to face challenges in complying with the plethora of federal laws that govern clinical research and in handling the risks involved in conducting clinical trials. The workshop series is designed to provide in-depth training and an opportunity for discussion with other clinical research sites on some of the most sticky compliance challenges for clinical research sites. Each half-day seminar will come with written materials, including template forms, and will be taught by attorneys from Coppersmith Schermer & Brockelman PLC and clinical research specialists from Arizona hospitals. The series will be held at Phoenix Children's Hospital in the Cohen Conference Center.



Who should attend?

Clinical research directors, clinical researchers, clinical research coordinators, IRB chairs, IRB coordinators, compliance officers, privacy and security officers, chief financial officers, billing personnel, and attorneys will be interested in these sessions.

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