IRBMED SEMINAR SERIES
Multi-Site Research - The Accepting and Ceding of IRB Oversight: What Investigators and Study Teams Need to Know

Ford Auditorium
April 10, 2016
9:00am – 11:30pm

AGENDA

Welcome & Introduction  Judith Birk, JD  9:00 am

Institutional Decisions for Ceding and Accepting IRB Oversight  Lois Brako, Ph.D. and Judy Birk, JD  9:10 am

Ceding IRB Oversight  Angela Faber, BS, CIP  9:40 am

Break and snacks  ------------------  10:05 am

Research Pharmacy: An Ancillary Review  Amy Skyles, PharmD  10:20 am

Accepting IRB Oversight  Judith Birk, JD and Robin Sedman, MSN, M.Ed.  10:40 am

Mock IRB  11:00 am

This installment in our series of mock IRB meetings provides insight into IRB review of a serious adverse event at external site when IRBMED is serving as the IRB of Record for the site. This will be followed by a Q&A session with IRBMED Board Members comprising the mock IRB:

Ann Dillon, BS, CIP – Regulatory Team  Alan Sugar, MD, Chair
Amy Filbrun, MD, Vice Chair
Amy Skyles, PharmD
Corey Zolondek, PhD
Duke Morrow, MDiv, DMin
Judith Avery, MA

End  ------------------  11:30 am