IRBMED SEMINAR SERIES

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

MEDICAL SCHOOL
UNIVERSITY OF MICHIGAN

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**

Seminar Series