



ASK THE COMPLIANCE OFFICE

Presents:

*Let's Talk About the Consent Process
with*

*Kris West, J.D.,
Director, Office of Research Compliance*

When: July 29, 2009

Where: Emory Hospital Auditorium, 2nd Floor

Time: 11:30am – 1:00pm



One in four FDA Warning Letters involving clinical trials cite deficiencies with the consent process.

DON'T BE NEXT!!

Learn how to properly obtain and document informed consent for clinical trials.

Register using Peoplesoft (requires Emory ID and password). There is no charge for registering; insert 00000 the cost field. The course number 265003.

Bring your lunch. Dessert and water will be supplied.