



Office of the Human Research Protection Program

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HUMAN RESEARCH NEWS: UPDATE ON EDUCATION AND TRAINING

Date: 2010-05-13

Noontime Luncheon Series: OHRPP now offers two types of hour-long training sessions for UCLA research staff and faculty. Both are held once a month during lunch hours. See below for descriptions, locations, target audiences, and schedule through 2010. You are welcome to bring your lunch.

Please RSVP: We recommend but do not require that you RSVP if you are planning to attend so that there will be enough handouts for everyone. To RSVP, please either call 310-206-7081 or e-mail OHRPPEducation@research.ucla.edu and indicate your name, your department, and your contact number.

	NOONTIME EDUCATION SERIES	LEARN AT LUNCH
Who should attend?	Anyone interested in the topic	Clinical Coordinators, Research Associates, and anyone interested in the conduct of clinical research
When is it?	3rd Wednesday of the month	1st Thursday of the month
What time is it?	Noon to one	Noon to one
Where is it?	11000 Kinross, Suite 210	Bel Air Room, 17-323 CHS
Who presents?	OHRPP Staff	OHRPP Staff and invited speakers

2010 Schedule for Noontime Education Series at Kinross:

Date	Title
May 19	Investigational Drugs?What to Consider and Include in the IRB Application
June 16	Children in Research?What to Consider when Preparing and IRB Application
July 21	Tips for Speeding Up Approvals of Biomedical Applications
August 18	Conducting a Risk Assessment?Determining Level of IRB Review
September 15	Tips for Converting Your Paper Application to webIRB
October 20	Tips for Speeding Up Approvals of Social Behavioral Applications
November 17	Post IRB Approval Requirements?Amendments, Continuing Review, Reporting Unanticipated Problems and Updated Safety Information

2010 Schedule for Learn at Lunch at CHS:

Date	Title
June 3	Post-Approval Reporting Requirements (Unanticipated Problems, Adverse Events and Updated Safety Information)
August 5	HIPAA for Clinical Researchers
September 2	UCLA Investigational Drug Service
October 7	Data Safety Monitoring Plans
November 4	Overview of FDA Drug Approval Process

Survey Monkey for Educational Needs for Conducting Human Research: Please click on this link for the [Survey Monkey](#) if you are interested in providing feedback about your education and training needs. This survey should take about five minutes to complete and will help the OHRPP office make the appropriate training available to the human research community.