



Home

Contact OHRPP

[Office](#)
[Staff Directory](#)
[Program Feedback](#)
[Kinross Consults](#)

For Researchers

[Clinical Research Toolkit](#)
[Consent Form Templates **New!**](#)
[Emergency Use of a Test Article](#)
[External IRBs](#)
[Getting Started **New!**](#)
[HIPAA](#)
[IRB Feedback Survey](#)
[IRB Review Fees](#)
[Letter to Sponsors](#)
[Meeting Calendars](#)
[ORA Research Portal](#)
[Other Resources](#)
[Participants' Bill of Rights](#)
[Policies & Guidance](#)

What's New

[Human Research News](#)

About OHRPP

[Overview and Purpose](#)
[Organization Chart](#)
[HRPP Program Description](#)

Education & Training

[Certification \(CITI Training\)](#)
[HIPAA Training](#)
[Monthly Training Sessions](#)
[Training Upon Request](#)

Post Approval Reporting

[Overview](#)
[PI Reporting Requirements](#)
[Decision Trees](#)
[Other Post Approval Activities](#)

Quality Improvement

About UCLA IRBs

[IRB Descriptions](#)
[FWA and IRB Registration](#)
[Human Research Policy Board](#)
[IRB Statistics](#)

For and About IRB Members

[Meeting Calendars](#)
[IRB Member Rosters](#)
[Members Checklists](#)
[Member Standards](#)
[References](#)
[Assessing Risk](#)

For Research Participants

[Información Para Participantes](#)
[Participant Survey](#)

HUMAN RESEARCH NEWS: IMPORTANT WEBIRB UPDATES

Date: 2011-01-05

Updated webIRB Rollout

An updated version of webIRB will be released on Monday, January 11, 2011. The updates and revisions are based on feedback from the UCLA research community. Unlike the previous release, very few additional questions will be added. Some questions will be deleted.

The majority of the changes includes added guidance, revised text, and edited branching to accommodate additional types of studies. Other changes include:

- Ability to submit Post Approval Reports along with amendments
- Simplified amendment submissions
- Enhanced e-mail notifications (e.g., ability of study team to send e-mail messages to other study team members)
- Additional information on all Notice templates (i.e., funding, grant number, short title)

webIRB Server Down This Weekend!

In order to deploy the revisions noted above, the webIRB server, including the Sandbox, will be unavailable from 5:00 pm Friday, the 7th to 5:00 pm Sunday, the 9th. We apologize for any inconvenience but this downtime is necessary in order to make the needed upgrades.

webIRB News Rolled into Human Research News

Because all researchers and staff are now required to use webIRB the webIRB News list serve has been canceled. All webIRB updates and notifications will be incorporated into the *Human Research News* list serve.

- *To subscribe* to future announcements, click on the link below. Scroll down to the section titled "Subscribing to Investigators-I" and provide the required information.
- *To unsubscribe* click on the link below and scroll to the bottom of the page.
- <http://lists.ucla.edu/cgi-bin/mailman/listinfo/investigators-I>

Improved IRB Review and Approval Turnaround Times

Since the implementation of revised IRB procedures and webIRB, the turnaround time for IRB review and approval has decreased significantly. See the [ORA Online Research](#) Center portal for an historical overview of the IRB volume and turnaround times. See table below for current average turnaround times.

Important Notes:

1) As these are median times listed below, *allow more time before and after holiday periods.* 2) The clock starts when a *complete* study is received in the OHRPP office. *Incomplete submissions and delayed PI responses increase actual turnaround times.*

Type of Studies and Review	Median Turnaround Time
Full Committee New Studies	45 Days
Full Committee Continuing Reviews	24 Days
Full Committee Amendments	35 Days
Expedited New Studies	15 Days
Expedited Renewals	7 Days
Expedited Amendments	5 Days

[Add comment](#) | [Revise](#)