



OHRPP Updates

September 10, 2020

OHRPP Updates

**Investigator
Responsibilities**



Learn at Lunch



**OHRPP Office
Hours**



**Human Research
News**



Investigator Responsibilities - overview

Qualifications to serve as a PI are set by UCLA Policy

- [UCLA Policy 900](#)

PI Duties and Responsibilities are set by UCLA Policy

- [UCLA Policy 991](#)

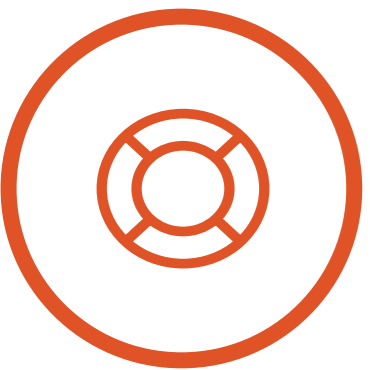
OHRPP guidance and procedure

- [PI Responsibilities](#)



Investigator Responsibilities - highlights

- As PI, have **sufficient training/qualifications** to safely oversee this specific type of research project:
 - **Clinical medical expertise to oversee biomedical clinical trial**
 - **Cultural expertise to oversee research in specific communities (such as in foreign countries, with indigenous communities, etc...)**

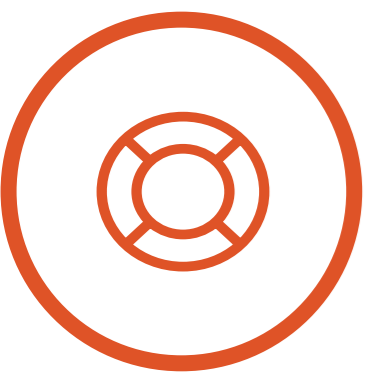


Investigator Responsibilities - highlights

- **Ensure that all *staff on the research* team are:**
 - ☑ **Qualified** to do the tasks they are assigned
 - ☑ Have **completed all required training** (as directed by IRB, other UCLA entities, and as required by the funding agency/Sponsor, if applicable)
 - ☑ Are appropriately **supervised**
 - To successfully conduct the tasks they are qualified to do
 - To **not** conduct tasks for which they are not qualified

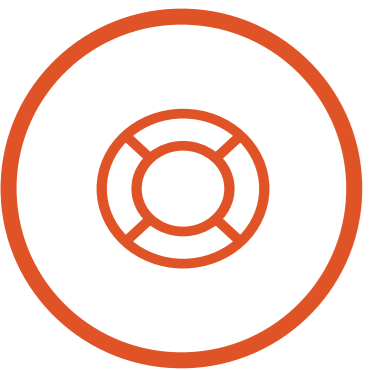
Investigator Responsibilities - highlights

- **Design the study to**
 - maximize benefits and minimize risks
 - ensure equitable subject selection
 - Obtain legally effective consent or waiver from the IRB
 - Obtain HIPAA authorization or waiver from the IRB (acting as the privacy board for research), if applicable



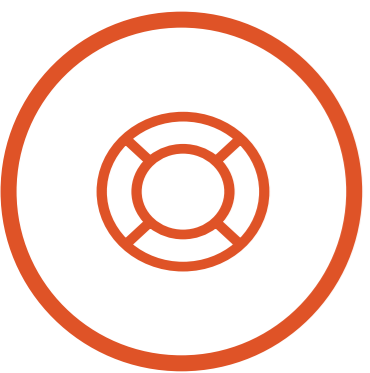
Investigator Responsibilities - highlights

- **Begin research only after *IRB and all other institutional/regulatory requirements* are met. Other approvals/reviews might include:**
 - Research Advisory Panel of the CA Attorney General’s Office (for schedule 1 or 2 drug research)
 - CIRC
 - ISPRC
 - MRSC
 - Etc...



Investigator Responsibilities - highlights

- **Conduct the research as described in the IRB-approved application**
- **Submit Amendment applications and receive approval before implementing changes to the protocol**
 - *Except for deviations from the approved protocol to prevent immediate hazards to participants*
- **Submit PAR applications as required per the PAR Guidance**
- **Submit continuing review applications or complete annual PI assurances to continue the research**



Investigator Responsibilities - highlights

- When in doubt about IRB requirements, *ask OHRPP directly*

Learn at Lunch - September

“Investigator Responsibilities for single site (UCLA) and central/lead site (UCLA + other sites) under a reliance agreement: best practices and how to avoid pitfalls”

Presenter: Moore Rhys, OHRPP

Date/time: September 24, noon-1pm

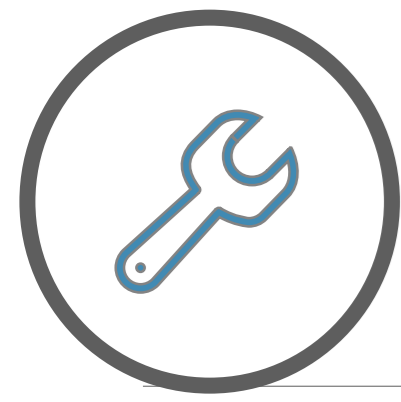
[Register in advance](#)



OHRPP's "Office Hours"

- OHRPP Quality Improvement Unit staff are hosting *half-hour open Q/A sessions every other week* to answer your questions
- *Upcoming sessions*
 - Tuesday, September 15, 2020 8:30am
 - Tuesday, September 29, 2020 8:30am

Register once and you can join any session.



Reminder - Subscribe to Human Research News

To be in the know when OHRPP releases updated guidance and offers training opportunities, please subscribe to Human Research News

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Any Questions?

Contact Information

OHRPP Website

<https://ohrpp.research.ucla.edu>

Moore Rhys, OHRPP Asst. Director

Phone: x46339

Email: moore.rhys@research.ucla.edu