UCDAVIS HEALTH | Clinical and Translational Science Center

UCDAVIS LIBRARY

Regulatory/IRB Overview

Clinical Trial Office Regulatory Dept: April Cuellar, CCRC
Laura Pulido-Ramirez, BS, CCRC

IRB Education Program Lead: Nicole Walters, BS, CIP, PMP

DATES:
June 12, 2020
July 10, 2020
August 14, 2020
September 11, 2020

TIME: 12 to 2 p.m.

PLACE: https://ucdavis.zoom.us/j/596870816

Complimentary assistance with IRB and regulatory questions.

GET HELP ON THE FOLLOWING:

- Does my project require IRB review?
- How do I obtain IRB approval?
- How do I submit a protocol deviation to the IRB?
- How do I obtain ancillary approvals, e.g., IT evaluation, Radiation Use Committee, etc.?
- How do I obtain IRB approval for recruitment methods, such as social media, email blasts, and telephone scripts?

No appointment necessary!

Sending your question(s) ahead of time is strongly encouraged and can be emailed to hs-ctoregulatory@ucdavis.edu.

2020 Clinical Researcher Office Hours

Email general questions about office hours to the CTSC Clinical Research Education Program:
hs-CTOeducation@ucdavis.edu

Email questions about Regulatory/IRB office hours and support to:
hs-ctoregulatory@ucdavis.edu

For updated listings of CTSC seminars and events, please visit our calendar:
health.ucdavis.edu/ctsc/eventcalendar

The UC Davis CTSC is supported by award TR001860 from the NIH National Center for Advancing Translational Sciences.