UCLA-CDU CFAR
Clinical Science Core
Fee Schedule

INSTITUTIONAL REVIEW BOARD (IRB) INITIAL SUBMISSION **$1273.66**

Initial application for IRB approval of a new study. Includes review of study design, informed consent procedures and forms, plan for privacy/ confidentiality and data security, assistance with a recruitment plan and materials, and other items.

IRB AMENDMENT **$372.99**

Modifications to IRB approved research studies. Includes changes to the protocol, informed consent, and/or recruitment materials.

IRB CONTINUING REVIEW **$191.04**

An IRB progress report of ongoing human subjects research (also known as “renewal”). Required once a year for ongoing research studies.

IRB POST APPROVAL REPORTING (PAR) **$191.04**

Reports informing the IRB of items pertaining to research conduct or participant safety, such as adverse events, violations, deviations/incidents, and updated study safety information.

IRB STUDY CLOSURE **$191.04**

A request submitted to the IRB when all research activity and data analysis is completed.

INSTITUTIONAL BIOSAFETY COMMITTEE (IBC) **$682.32**

The IBC is an ancillary regulatory committee that oversees research involving infectious agents, recombinant/ synthetic nucleic acids, transgenic animals, etc. The purpose of IBC is to ensure researchers have safety and reporting plans in place to prevent spills/accidents, as well as if spills/accidents happen. (Note: If your study involves blood draws or processing lab samples from HIV positive participants, you will need IBC review.)

RSVP QUERY AND UNLIMITED EMAIL REFERRALS (NO CALLS) **$109.18**

RSVP is a volunteer registry of HIV positive and negative individuals from the greater Los Angeles area interested in participating in research. Use of the registry requires specific IRB approval.

Updated: 08/31/23

For questions, please contact Stephanie Buchbinder at sbuchbinder@mednet.ucla.edu