PRESBYTERIAN

The PHS HRPO charges an IRB review fee to human research projects and requests for exempt determination. All fees are assessed according to the established fee schedule.

PHS IRB Fee Structure

New Study Review - Industry Sponsored/Federally Funded/Granted Trials

$2000	Initial Review of the protocol, investigator brochure, consent form(s), study materials, etc.
$2200	Cede Review Fee - request to cede review to an external IRB (includes initial review of the protocol, investigator brochure, consent form(s), study materials, etc.). There is an additional IRB Authorization Agreement processing fee (see below)
$750	Continuing Review
$400	Protocol Amendment – major changes
$300	Informed Consent revision – major changes
$100	Revisions to subject related material, investigational brochure, package insert, recruitment materials, advertising, dear inv. letters, change in Principal Investigator (PI), etc.

Exempt Determination of a funded study

$500	Flat fee for the duration of the study

Unaffiliated Investigator Initiated, Unfunded Studies

$500	Initial Review of protocol, investigator brochure, consent form(s), study materials, etc. - one flat fee for the duration of the study, show proof of (no) funding

Other Services

$350	Additional charge for late continuing review submission
$1,000	Processing of an IRB Authorization Agreement (when utilizing an external IRB). This fee is in addition to the cede review fee.
No charge	Serious Adverse Event Reporting
No charge	DSMB or misc. reports from the sponsor
No charge	Emergency Use or Compassionate Use request
No charge	Study Closure
No charge	Student/Pharmacy Resident lead projects.