



**IRB Fee Schedule
Single Site Studies***

SERVICE	Fee
Initial Review - expedited or convened review, as required by the regulations, may include review of protocol, one informed consent document, assent form (if applicable), drug/device brochure (if applicable), recruitment materials, subject materials, and IRB application for the principal investigator. <i>There is a vetting process for first time investigators (no additional charge).</i>	\$2,480
Additional Informed Consent Form(s) – each (Beyond the first informed consent document submitted with Initial Review e.g., blood draw consent form, sub-study consent form)	\$385 each
Review of Application for Exempt Determination or Not Human Subject Determination <i>Note: Projects submitted for exempt determination but do not qualify may need expedited or convened review. Only the applicable review fee will apply.</i>	\$1,105
Continuing Review – interval as determined by the committee	\$2,480
Review of Annual Status Report (Only for studies <u>not</u> requiring continuing review under 2018 requirements)	\$550
Review of Modifications to previously approved research** Examples include but are not limited to the following: Amendments Package insert Change/add location Safety letters ICF Revisions PDR Protocol exceptions SAEs IDB/IB updates Safety letters Safety Reports	\$605
Change of Principal Investigator Includes but is not limited to vetting procedure, ensuring researcher is qualified by training and experience, reviewing conflicts of interest, confirming HSP training, completing a license check as applicable, and updating informed consent form.	\$1,105
Addition / Change of Key Personnel Includes but is not limited to reviewing any conflicts of interest, confirming HSP training, and completing a license check as applicable.	\$420

CONFIDENTIAL



Review of Study Materials (after initial submission) ** Examples include but are not limited to the following: Advertisements Subject diary DMC/DSMB reports Recruitment materials Letters to subjects	\$385
Study Closeout - review of final closeout report	\$605
Reportable Events (e.g., adverse events, reportable external adverse events (IND), deviations) <i>Note: IND Reports - BRANY IRB review is required only for reports that meet the definition of an unanticipated problem.</i>	\$110 per report
Foreign language translations coordinated by BRANY (upon request) Includes securing estimates from vendor, coordinating sponsor/CRO approval, ensuring certificate of accuracy is received, administrative IRB review, preparation of final IRB documentation, and circulating final acknowledgement letter. <i>(Note: the cost of translation is billed on a per word basis)</i>	Cost of Translation with handling fee plus \$605
Late submission Fee – when receipt of Continuing Review is after the submission deadline, which is less than 14 calendar days from the IRB approval expiration date, a late fee will apply in addition to the Continuing Review Fee. (e.g., Continuing Review Fee + \$220= \$2,700)	Add \$220 to Continuing Review Fee
Withdrawal of Submission – fee applies if study has been withdrawn after being placed on an IRB meeting agenda or provided to an IRB reviewer	\$330
Rush Fee Full Board Initial Review accelerated outside scope of normal time frame. Does not guarantee IRB approval. (Initial Review Fees + \$825 = \$3,305)	Add \$825 to Initial Review fee

**This fee schedule will apply to studies having up to 4 study sites. Each site is processed as a single IRB submission and the related IRB fees will apply. For studies with five sites or more please contact BRANY for a multisite IRB fee schedule.*

**** Multiple documents submitted together that require independent review may receive more than one charge.**

Note: These fees are subject to change without advance notice.

CONFIDENTIAL