

<b>IRB Fee Schedule Single Site Studies*</b>
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SERVICE	Fee
<b>Initial Review</b> - 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>	\$2,360
<b>Additional Informed Consent Form(s)</b> – each (Beyond the first informed consent document submitted with Initial Review e.g., blood draw consent form, sub-study consent form)	\$365 each
<b>Review of Application for Exempt Determination or Not Human Subject Determination</b>  <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>	\$1,050
<b>Continuing Review</b> – interval as determined by the committee	\$2,360
<b>Review of Annual Status Report</b> (Only for studies <i>not</i> requiring continuing review under 2018 requirements)	\$525
<b>Review of Modifications to previously approved research**</b> (e.g., amendments, ICF revisions, IDB/IB updates, Package Insert, PDR, safety letters, change in site, protocol exceptions, safety reports, safety letters, SAEs)	\$575
<b>Change of Principal Investigator</b> 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.	\$1,050
<b>Addition / Change of Key Personnel</b> Includes but is not limited to reviewing any conflicts of interest, confirming HSP training, and completing a license check as applicable.	\$400
<b>Review of Study Materials (after initial submission) **</b> (e.g., advertisements, recruitment materials, subject diary, letters to subjects, DMC/DSMB reports)	\$365



<b>Study Closeout</b> - review of final closeout report	\$575
<b>Reportable Events</b> (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>	\$105 per report
<b>Foreign language translations coordinated by BRANY</b> (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 \$575
<b>Late submission Fee</b> – 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 + \$210= \$2,570)	Add \$210 to Continuing Review Fee
<b>Withdrawal of Submission</b> – fee applies if study has been withdrawn after being placed on an IRB meeting agenda or provided to an IRB reviewer	\$315
<b>Rush Fee</b> Full Board Initial Review accelerated outside scope of normal time frame. Does not guarantee IRB approval. (Initial Review Fees + \$785 = \$3,145)	Add \$785 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**