



DATE: Effective Date is September 1, 2023

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Subject: IRB Fee Schedule for FY2024\*

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<b>Billing Code</b>	<b>Activity</b>	<b>Amount</b>
IR	<b>Full Board Initial Review of Protocol:</b> Includes protocol, investigators' brochure, Informed consent document(s), recruitment materials, review of Investigator Credentials	<b>\$ 3,000</b>
PR	<b>Periodic Review (Full Board Continuing Review)</b>	<b>\$ 1,500</b>
FBA	<b>FULL BOARD Revisions/Amendments/Changes to Research:</b> Protocol amendments/revisions; Consent Form research-related modifications (note: simultaneous change to consent form based on protocol revision to protocol is considered a "single change").	<b>\$ 750</b>
FR	<b>Final Report/Close-out</b>	<b>\$ 500</b>
EXP	<b>EXPEDITED REVIEW (Initial Review)</b>	<b>\$ 1,000</b>
EXE	<b>EXEMPT Category Determination/Review (Initial Review)</b>	<b>\$ 1,000</b>
NHR	<b>Formal NHR or QI/QA determination</b>	<b>\$ 350</b>
EXA	<b>EXPEDITED/EXEMPT Revisions/Amendments/Changes to Research:</b> All other changes to protocol or consent document not requiring review by convened IRB but requiring IRB Chair/Designee review	<b>\$500</b>
HUD	<b>Humanitarian Use Device Review</b> Continuing Review of HUD/HDE	<b>\$ 2,000</b> <b>\$1,000</b>
ADA	<b>Administrative Letters, Administrative Changes to Protocol:</b> Not requiring consent form modifications	<b>\$ 250</b>
CIRB	<b>Commercial IRB on-site fee (UNTHSC investigators only)</b> <b>UNTHSC delegation to another non-federal IRB</b> <u>One time charge</u> for managing local institutional human subject protection oversight and IRB-related regulatory issues across all project years	<b>\$ 5,000</b>
RE	<b>Reportable Events Review</b>	<b>\$ 250</b>



\*The above schedule applies to for-profit, industry-sponsored projects (e.g., sponsored clinical trials), as well as institutions/organizations not affiliated with the North Texas Regional IRB (with the exception of the Commercial IRB on-site fee which applies to UNTHSC investigators only).