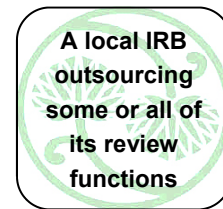


New England IRB: Central IRB Providing Study and Site Transition

New England IRB is dedicated to the protection of human subjects in clinical trials. Over the past 21 years, we have worked with sponsors, CROs, hospitals and academic institutions across the U.S. in transitioning their clinical trials for protocol and site review.

This has included studies being transitioned from:



We have developed a streamlined study and site transition process. Combined with the industry's fastest review turnaround, this has been instrumental in assisting sponsors, CROs and PIs to continue research while keeping transitional delays to a minimum.

- 🕒 1-week protocol review; 24-hr site review
- 🕒 Overnight notification
- 🕒 Electronic submission process
- 🕒 Approval letters posted online
- 🕒 Single point of contact

Contact Information

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We understand that high quality and fast turnaround are priorities in selecting a central IRB. We also recognize the importance of providing these with a sensitivity to each organization's communication needs.

New England IRB has full-AAHRPP accreditation status and is in good standing with FDA and OHRP.

Contact us to learn how we can provide support in transitioning protocols and sites to New England IRB for review.

