IRB

Oversight Without Obstacles

Institutional Review Board (IRB) Services

Regardless of your project's scope, therapeutic niche or number of investigators, Advarra is your partner for the conduct of efficient, responsible research.

IRB Solutions to Empower Better Research Programs

Experienced Project Managers

Advarra's coordinators help you streamline tasks and reach a full understanding of compliance recommendations.

Thorough Review

You can trust Advarra to provide high-integrity reviews and feedback regardless of project size, scope or complexity.

Deep Therapeutic Expertise

Our IRB has expertise across all major therapeutic areas and sets the standard in both new and expanding fields.

Real-Time Reporting

With web-based technologies that are available anytime, anywhere, you can log in to view projects and plan ahead.

Streamlined Communication

Advarra combines a dedicated project management team with powerful online tools to simplify collaboration.

Rapid Turnaround

Advarra provides prompt, accurate and thorough reviews and works with you to meet demanding timelines.

Simplified Communication

Advarra is committed to getting your trial started on time and helping you reach critical study milestones quickly.

Review Item	Submission to Decision
New protocol and initial informed consent for multisite studies (full board review)	4 business days
New protocol and initial informed consent (minimal risk review)	1-2 business day
New site for a multisite study	1 business day

Therapeutic Specialization

In addition to offering expert guidance in all research phases and across all major therapeutic areas, Advarra also provides highly specialized review services for areas such as oncology and neurology. Our IRB members and operational staff include distinguished scientists and industry leaders with a deep understanding of and appreciation for the regulatory and scientific framework in which we work.

Technology-Enabled Solutions

The Advarra Center for IRB Intelligence (CIRBI) Platform sets the gold standard in review quality, submission turnaround time and document accessibility. Enjoy unmatched transparency, accessibility and quality oversight. With real-time notifications and status postings, you'll always know exactly where your study is in the review process.

- Shorten your site activation timeline with Advarra's IRB-Ready® approach and the Advarra CIRBI Platform
- Improve your communication and access projects anytime, anywhere in real time
- Make informed decisions with powerful project status metrics

Services and Capabilities

- Daily IRB meetings
- Ad-hoc review when needed
- Specialized services for Phase I and minimal risk research
- eConsent enabled and proficient in all major platforms
- Draft review and advisory opinion capabilities
- Dedicated support for institutions and therapeutic research consortia
- Coordinated IRB and IBC review for recombinant DNA research
- HRP/IRB consulting
- Central point of contact for every project
- Native US and Canadian review capabilities

Ready to make your research altogether better?

Contact BusinessDevelopment@advarra.com to get started.

ADVARRA

advarra.com