Protecting Patients, Not Trying Patience

Your IRB, Simplified

Univo IRB is restoring the IRB industry by returning focus to the founding principle of protecting patients in research. We're led by IRB review process and clinical operations veterans, bringing decades of experience and a modern approach to trials designed for today.

We understand the challenges placed on sites, and we know that the best patient protections come from strong partnerships founded on excellence in customer service. Whether you're managing Phase I-IV or noninterventional trials, when you work with us, you benefit from resources and tools that are specifically designed to ensure your research is safe and compliant.

Challenges? ***Overcome.*** Patients? ***Protected.*** ***That's our purpose, and that's our promise to you.***

Uniquely Univo

What sets us apart from other IRBs?

- **Patient education and advocacy:** Better informed patients, expanding clinical research knowledge and opportunities to more individuals in need
- **Forward-thinking approach:** Easier, faster IRB submission process leveraging modern technology and real-time dashboards
- **Service-first process:** Direct access through live chat and direct calls to your dedicated account personnel managing your studies
- **Seasoned IRB experts:** Decades of regulatory and clinical operations experience
- **Fair pricing:** No upcharges and free advisory reviews

ONEVERSE®

IRB Submissions Made Easy

OneVerse is an industry-first IRB platform that streamlines your document access and submission. It's built for efficiency and transparency, simplifying the IRB review process with easy-to-use features.

Want to get your IRB approval faster? Crush your time lines using a tool that's purpose-built for today's trials.

Simplify IRB Review With Univo IRB

You now have IRB options. Why should you partner with Univo IRB?

For agile solutions and a forward-thinking approach that makes the IRB submission process straightforward and transparent.

Because our goal is to protect your patients, not try your patience.

NEED FASTER APPROVALS?

Our seasoned experts and streamlined IRB review process delivers review decisions in as few as **4-5 BUSINESS DAYS** for new studies reviewed by the full board, **1-2 BUSINESS DAYS** for new sites added to approved research, and **2-3 BUSINESS DAYS** for recruitment and minor changes to research. ***Meet your milestones on time.***

Discover how we can help get your study started faster.

EXPEDITE YOUR REVIEW

ABOUT UNIVO IRB

Univo IRB is a next generation institutional review board (IRB) specializing in the ethical review of Phase I-IV pharmaceutical, device, biologic, behavioral, and psycho-social research. We offer agile study approaches, expert solutions powered by industry-leading study technology, and a service-first approach. Univo IRB holds accreditation through the Association for the Accreditation of Human Research Protection Programs (AAHRPP). With support from senior advisors and 60+ years of industry experience, Univo IRB guides your study to approval while respecting the rights and welfare of patients every step of the way.