Huron's IRB Interim Staffing Solution

A Trusted and Proven Solution for Growing and Sustaining Operations of Human Research Protection Programs

Huron's IRB Services Interim Staffing Solution provides professional leadership and submission processing support for Human Research Protection Programs (HRPPs) and Institutional Review Boards (IRBs). Our previous experience and regulatory expertise give us the opportunity to quickly place trained staff and provide institutions with the capacity to address local resource shortages and/or eliminate backlogs. Our team of seasoned experts understands and appreciates the challenges and unique situations HRPPs face, enabling us to be an effective partner, advisor, and resource for your institution.

Interim Staffing Services	
Leadership	**Submission Processing**
Supply interim leadership to keep organization running until a permanent leader is identified	Pre-review incoming items to ensure review readiness
Mentor and evaluate IRB staff, reviewers, and chairs to maximize team performance	Execute Not Human Subjects Research and Exempt Determinations
Hire and re-train IRB staff	Review minimal risk and full board biomedical and social behavioral research studies
Assist with AAHRPP accreditation preparation	Provide expertise in the Revised Common Rule Subparts B, C, and D, and additional regulatory criteria across all federal agencies
Provide regulatory consultation to the research community	Train and deliver continuing education for IRB members
Facilitate regulatory audits and inspections	Draft and authorize meeting minutes and decision letters
Develop and refine policies and procedures	Support study teams with navigating the IRB submission and review process

What sets Huron Apart?

Huron's IRB Services Team has been providing interim staffing support to research institutions of all sizes across the United States for nearly two decades. Our team's experience in developing a customizable Toolkit of policies and procedures that has been utilized across hundreds of institutions highlights our mastered understanding of the federal regulations governing human subjects research. This, combined with our team's exposure and adaptability to numerous IRB software and Clinical Trials Management Systems (CTMS), equips us with the capability to fill any role quickly so that you organization can begin to see immediate results.

The single most important factor that our clients will attest to is the competency and level of service provided by the entire project team. As your partner, the Huron team will work to eliminate backlogs and sustain fluid review operations through its commitment to tangible turnaround time milestones, while continually providing a customer-serviced-based approach to your research community. We will share accountability for your success and dedication to achieving the results aspired to by your institution.

HURON

huronconsultinggroup.com