HURON IRB

THE BUSINESS OF RESEARCH IS ONLY GETTING MORE COMPLEX AND CHALLENGING

The fiercely competitive research environment challenges institutions to maintain a world-class research enterprise that moves society forward, promotes innovation and advances knowledge — all without adding additional administrative burden to researchers.

Facilitate high quality, compliant reviews with an easy-to-install, comprehensive and extensible electronic IRB solution from the leader in IRB automation. Huron's IRB solution embeds compliant IRB operating practices from its popular HRPP Toolkit into proven Huron software. The solution combines Huron's policy expertise and human research protection best practices with electronic workflow design patterns based on 10+ years of successful software implementations. Currently, more than 80 leading research institutions use Huron IRB to improve their compliance efforts.

Deploy your solution with speed and flexibility. You don't have to choose between rapid deployment and configuration flexibility with Huron IRB. Providing a full-featured comprehensive solution out of the box, the system can be rapidly configured to be up and running quickly and efficiently. Huron IRB is part of the Huron Research Suite, a comprehensive suite of software solutions to facilitate communication, relieve administrative burden and free up time for what matters most — your research mission.

ORGANIZATIONAL BENEFITS	SOFTWARE FEATURES
Meet AAHRPP accreditation standards	Huron HRPP toolkit SOPs, checklists, worksheets and AAHRPP reporting
Keep up with regulatory changes	Incorporation of policy updates as they occur
Manage the entire study lifecycle end-to-end	Support for initial submissions, modifications, reportable new information and continuing reviews
Assign the appropriate IRB of record for each study	Support for multiple study models, including single-site studies, collaborative studies and multi-site studies
Manage relationships with other institutions	Tracking for authorization agreements, communication plans, accreditation status and points of contact profiles for institutions with whom your institution collaborates on research studies
Maintain regulatory compliance with the NIH Final Policy and the HHS Final Common Rule on the use of a single IRB of record for multi-site research	Full support for the single IRB (sIRB) of record and participating sites (pSite) in collaborative and multi-site studies
Significantly reduce the administrative burden and staff time required to maintain multi-site studies	Built-in integration with the Huron IRB Exchange, a cloud-based subscription service that facilitates the exchange of data between a sIRB and pSites in multi- site and collaborative research
Eliminate paper processes	Comprehensive form sets with validations and document attachments
Speed collaborative application preparation by study teams and reduce delays in routing and review	Electronic workflow routing and review, including parallel ancillary reviews
Accelerate processing of meeting discussions and decisions into compliant records and correspondence	Integrated meeting management facility

ORGANIZATIONAL BENEFITS	SOFTWARE FEATURES
Keep everyone notified of important milestones and the need to attend to their part of the process	Automated reminders
Simplify required communications	Correspondence generation
Ensure the study team uses only the latest, approved consent form and your own watermarks	Consent form stamping and watermarking
Manage the audit trail for all document versions	Electronic document management
Speed system adoption	Integrated on-line help and comprehensive user documentation
Reduce errors and compliance risk, and facilitate rapid access to remote data	Built-in system integration with Huron's COI, Grants and CTMS solutions
Simplify eIRB setup and ongoing operation for IT-overburdened organizations	Hosted solution also available

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