IRB Submission Forms

All Human Subjects Research studies must be submitted to the USA IRB through IRBNet. To provide the USA IRB with essential project details, every IRBNet package must include a completed supplemental form generated from the links below.

These links route directly to interactive web-based forms. Detailed guidance and templates are included in each link. Complete the form online, download your final summary document, and upload it into your IRBNet submission package for IRB review.

The pdf created must be attached to your project in IRBNet so the IRB can review and provide approval / feedback.

IRB Submission Forms	
Form Name & Access Link	**Purpose & Overview**
New Study Application	All new projects must have a new study application attached to the project in IRBNet.
Annual Renewal / Check-In Form	All active studies require renewal with the IRB on a yearly basis, at minimum. Frequency and additional information can be found on the project's initial approval letter.
Amendments	Use this form to report any changes to an active study, including changes to the protocol, submission of new materials, personnel changes, recruitment or consent changes, etc. Changes cannot be implemented until formal approval has been granted by the IRB.
Reportable Events	Use this form to inform the IRB about any reportable events. This will vary by study type, and includes Adverse Events, Protocol Deviations, Protocol Exceptions, and Participant Complaints.
Emergency Use Post-Use Report	Required following the emergency use of an investigational drug or device to report safety and administration details to the IRB.
Study Closure	Every study must be closed with the IRB once all participant interactions, data collection, and identifiable analyses are complete.

Additional Resources:

- **New to USA IRB Submissions?** See our Submission and Review Guidance.
- **Need help navigating IRBNet?** Explore the IRBNet Tutorials on the ORCI website.
- **Have you completed all required training?** Reference training requirements on the ORCI website.
- **Building a consent form from scratch?** Use our consent templates to provide a basic structure and ensure you are using all federally and institutionally required language.
- **Receive a consent form from a sponsor / collaborator?** Reference our Consent Local Context Language to ensure all institutionally required language is inserted.

Still have questions? Contact us! We're here to help.