

HRPP Investigator Guidance Series

This page serves as a hub for all of the Investigator Guidance Series documents. Each document is an abbreviated version of one of our [HRPP Policies and Procedures](#) intended for investigators, coordinators, and other study team members. This page is a good starting point for any study team member with a question about a policy on a specific topic. A link to the full policy/procedure is included in each document.

Document	Updated
Advertisements (3.5)	12/13/2023
Authority of the IRB	12/13/2023
Change Requests	12/13/2023
cIRB	12/13/2023
Closure of Research	12/13/2023
Compensation	12/09/2024
Confidentiality	12/13/2023
Continuing Review	02/05/2024
Contraception Requirements	12/13/2023
Data and Safety Monitoring	12/13/2023
Data Registries	12/15/2023
Emergency Research - Waiving Consent	12/13/2023
Emergency Use of a Test Article	01/25/2024
Employees as Subjects	12/13/2023
Ethical Access	12/09/2024
Exempt Research	01/25/2024
Expanded Access to Investigational Drugs and Devices	12/14/2023
Expedited Review	12/13/2023
Financial COIs	01/25/2024

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<u>Full Board Review</u>	12/13/2023
<u>Humanitarian Use Device (HUD)</u>	12/13/2023
<u>Incidental Findings</u>	12/13/2023
<u>Increased monitoring, interim Continuing Review, and verification</u>	12/15/2023
<u>Informed Consent</u>	12/13/2024
<u>International Research</u>	12/13/2023
<u>Investigational and Marketed Devices</u>	01/26/2024
<u>Investigational and Marketed Drugs</u>	12/13/2023
<u>IRB Approval Criteria</u>	12/15/2023
<u>Obtaining Informed Consent for Non-English Speaking Persons</u>	12/13/2023
<u>Obtaining Informed Consent for Persons with Additional Needs</u>	12/13/2023
<u>PI Qualifications and Responsibilities (job description)</u>	12/13/2023
<u>Placebos</u>	12/13/2023
<u>Post-Approval Monitoring of Research</u>	12/09/2024
<u>Pregnancy Testing</u>	12/13/2023
<u>Privacy</u>	01/26/2024
<u>Recruitment</u>	12/13/2023
<u>Reimbursement</u>	12/13/2023
<u>Research Involving Children</u>	12/13/2023
<u>Research Involving Decisionally Impaired Persons</u>	12/13/2023
<u>Research Involving Neonates</u>	12/13/2023
<u>Research Involving Pregnant Women and Fetuses</u>	12/13/2023
<u>Research Involving Prisoners</u>	01/25/2024
<u>Research Personnel Qualifications & Responsibilities (study team job descriptions)</u>	12/13/2023
<u>Short Form Consent</u>	12/13/2023
<u>sIRB</u>	12/13/2023

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<u>Students as Subjects</u>	12/13/2023
<u>Study Hold</u>	12/13/2023
<u>Suspension</u>	12/13/2023
<u>Termination</u>	12/13/2023
<u>Using PHI in Research</u>	12/13/2023
<u>Vulnerable Populations - Additional Protections</u>	12/13/2023
<u>Waiving Consent Process</u>	12/13/2023
<u>Waiving Signed Consent</u>	12/13/2023
<u>Wash Out</u>	12/13/2023
<u>What requires IRB review and approval?</u>	01/10/2024

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