

# HRPP Investigator Guidance Series

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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.

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<a href="#">Advertisements</a>	12/13/2023
<a href="#">Authority of the IRB</a>	12/13/2023
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<u>Full Board Review</u>	12/13/2023
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<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
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<u>Research Involving Children</u>	12/13/2023
<u>Research Involving Decisionally Impaired Persons</u>	12/13/2023
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<u>Research Involving Pregnant Women and Fetuses</u>	12/13/2023
<u>Research Involving Prisoners</u>	01/25/2024
<u>Research Personnel Qualifications &amp; Responsibilities (study team job descriptions)</u>	12/13/2023
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