

January 29, 2015

Subject: Revised IRB Forms – Jan 29, 2015

In an effort to keep the research community informed of new and revised Office of Human Research (OHR) policies and forms, a summary of recent changes is being provided. Please remember to always access the most current policies and forms on the [Division of Human Subjects Protection page](#) of the OHR website.

Thank you

The Office of Human Research

## Forms

Number	Title	Version
NA	<b>Overview of IRB Submissions for New Studies</b>	<b>01/29/15</b>
Changes: For all new studies a completed OHR-1 must be included with your submission. A link to a description of the IRB submission process has been added. The address of the IRB has been added.		

Number	Title	Version
OHR-3	<b>Request for Waiver of Subject Authorization to Collect Protected Health Information</b>	<b>01/29/15</b>
Changes: You must select from a list of possible reasons that the waiver is being requested.		

Number	Title	Version
OHR-4	<b>Record/Chart Review/Computer Database Research Study</b>	<b>01/29/15</b>
Changes: Clarification: For all NEW studies a completed OHR-1 must be included with your submission.		

Number	Title	Version
OHR-8	<b>Informed Consent Document for Human Subjects Research</b>	<b>01/29/15</b>
Changes: You are instructed to modify the section about possible side effects according to your study.		

Number	Title	Version
OHR-15	<b>Request for Human Biological Specimens for Research</b>	<b>01/29/15</b>
Changes: Clarification: For all NEW studies a completed OHR-1 must be included with your submission.		

Number	Title	Version
OHR-16	<b>Tissue and Genetic Research</b>	<b>01/29/15</b>
Changes: Clarification: For all NEW studies a completed OHR-1 must be included with your submission.		

If you have any questions or comments about these changes, please contact [patrick.herbison@jefferson.edu](mailto:patrick.herbison@jefferson.edu) .