

Approved Documents

How to find stamped consent and assent forms and other approved documents

1. The easiest place to locate currently approved research documents, such as the consent and assent forms, is on the Study Documents tab in the study workspace. Note that the *study workspace* is different from application workspaces; to navigate back to the main study workspace from an application workspace, click on the title of the Initial Application.

ARROW
Application Review for Research Oversight at Wisconsin

Hello, uwirb pi1

My Home IRB IACUC Biosafety SCRO HELP

APPROVED

NO ACTION REQUIRED:
Approved by the IRB.

[VIEW APPLICATION](#)

Print Form | View Differences

[New Continuing Review](#)

[New Change](#)

[New Personnel Change](#)

[New Reportable Event](#)

Sample Protocol Application

Initial Application: [Sample Protocol Application](#)

APPLICATION DETAILS

ID: 2017-0047
PI: uwirb pi1
Board: ED/SBS IRB
Staff Reviewer: CASEY PELLIE
Reviewer Contact: casey.pellie@wisc.edu

MILESTONES

Date Submitted: 9/25/2017
Initial Approval: 11/1/2017
Expiration: 9/24/2018

Pre-Submission → IRB Staff Pre-Review → IRB Committee Review → Review Complete

Modifications Requested (between IRB Staff Pre-Review and IRB Committee Review)

History Follow-On Submissions Correspondence **Study Documents** ...

CHANGES
No data to display.

REPORTABLE EVENTS
No data to display.

CONTINUING REVIEWS

ID	State	Modified	Letter
2017-0047-CR001	Awaiting Correspondence	12/6/2017	View

2. From the Study Documents tab on the study workspace, approved documents, including stamped consent and assent forms, will be available by clicking on their title. Note that the only approved documents that will be stamped are *signed* consent and assent forms (unless the study has been determined as Exempt - in which case none of the consent document will be stamped). Recruitment documents, consent/assent information sheets, oral consent/assent scripts, etc. will not be stamped and will not appear here.

3. This information can also be found on your approval letter.

Title: TEST6
Principal Investigator: Uwirb PII
Point-of-contact:
IRB Staff Reviewer: STEPHANIE WILSON

A designated ED/SBS IRB member conducted an expedited review of the above-referenced initial application. The study was approved by the IRB member for the period of 12 months with the expiration date of 3/9/2018. The study qualified for expedited review pursuant to 45 CFR 46.110 and, if applicable, 21 CFR 56.110 and 38 CFR 16.110 in that the study presents no more than minimal risk and involves:

Category 7: Research on individual or group characteristics or behavior (including, but not limited to, research on perception, cognition, motivation, identity, language, communication, cultural beliefs or practices, and social behavior) or research employing survey, interview, oral history, focus group, program evaluation, or quality assurance methodologies

To access the materials approved by the IRB, including any stamped consent forms, recruitment materials and the approved protocol, if applicable, please log in to your ARROW account and view the documents tab in the submission's workspace.

If you requested a HIPAA waiver of authorization, altered authorization and/or partial