

Locating Stamped Consent Forms

On your dashboard, click "View All" to view your studies.

The dashboard shows various study statuses: In-Draft, Awaiting Authorization, Pre-Review, and Under Review. The 'My Studies' table lists several studies, and the 'View All' button is highlighted with an orange arrow and text overlay.

Select the study in order to find the stamped consent form.

Study ID	Study Name	Status	Reviewer	Approval Date	Expiration Date
IRB-FY21-63	study #4	Unsubmitted	Diane Director-IRB	07-28-2021	N/A
IRB-FY21-57	test study #3	Unsubmitted	Diane Director-IRB	07-12-2021	N/A
IRB-FY21-55	test study	Unsubmitted	Diane Director-IRB	06-28-2021	N/A
IRB-FY21-54	new study	Unsubmitted	Diane Director-IRB	06-28-2021	N/A
IRB-FY21-47	How to Avoid Common Pitfalls on the Road to IRB Approval	Under Review	Diane Director-IRB	05-13-2021	N/A
IRB-FY21-40	Bad Research	Under Review	Mary Beth Goodnight	04-12-2021	N/A
IRB-FY21-35	MRI Review Test	Approved	Diane Director-IRB	01-25-2021	01-24-2022
IRB-FY21-32	CL Test	Unsubmitted	Diane Director-IRB	01-22-2021	N/A
IRB-FY21-22	Another Internal Use	Under Review	Diane Director-IRB	12-04-2020	N/A
IRB-FY21-21	For internal use only?	Approved	Diane Director-IRB	12-04-2020	N/A
IRB-FY21-15	test2	Under Review	Mary Beth Goodnight	12-01-2020	N/A

At the bottom, click on "Attachments" to find your most recently stamped consent form.

The study details page shows metadata such as Approval Date, Expiration Date, and Organization. At the bottom, the 'Attachments' tab is highlighted with an orange arrow and text overlay.

Double click on the form to download it.

Role: Researcher | Diane Director-IRB

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Approval Date: 01-25-2021	Expiration Date: 01-24-2022	Organization: Users loaded with unmatched Organization affiliation. Current Policy Post-2018 Rule	Active Submissions: Modification Incident Incident	Population Flags:	Additional Flags:
Admin Check-In Date: N/A	Closed Date: N/A	Sponsors: N/A			

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Filename	Uploaded By	Date Uploaded	+ Upload Attachments
Consent - Signature (4) (1) (2).pdf		07-19-2021 2:31 PM	

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Double click on the form to download it

Stamps are located at the bottom right of the page.

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Purpose: The purpose of this research study is to increase our understanding of xxx by xxx.

Description of Study: Provide brief overview using lay language (4-6th grade reading level). Provide detail about the research procedures and specify what participants will be requested to do. State clearly any procedures that are experimental and safe use has not been documented. Include the type of procedures, if they will be audio or video taped, and where they will take place. Format such as headings, bullet points, indents, bolded type, etc. may be used to enhance clarity and facilitate explanation of information.

Number of Participants: 99.

Length of Participation: The overall time commitment will be no longer than xxx. However if you decide to stop participating in the study, we encourage you to tell the researchers.

Inclusion / Exclusion Criteria: list the inclusion and exclusion criteria for participants to qualify for this study.

Possible Risks: Instructions: Describe what is at risk. (Is there a possibility the participant will become tired, frustrated, bored, above and beyond what could be expected to occur in daily living). Describe how the investigator will respond if the risk occurs--(participant becomes tired or uncomfortable).

If appropriate: Participants will be informed about new research that provides additional information about risks or that may influence their decision to continue participation in this research.

Possible Benefits to the Participant: Payments to participants/monetary incentives should not be listed in this section.

Alternatives to Participation: Individuals may choose not to participate

Adult Consent

UT Dallas - Train
 IRB-PV21-35
 Approved on 1-25-2021
 Expires on 1-24-2022

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