

## **Status Report - General Protocol**

Thank you for your response! The IRB staff will start working on the continuation review. Please contact us at irb@usu.edu with any questions.

Below is a summary of your responses

Download PDF

Protocol Title

Status Report (Closure)--Excellent Example!

Protocol Number

99999

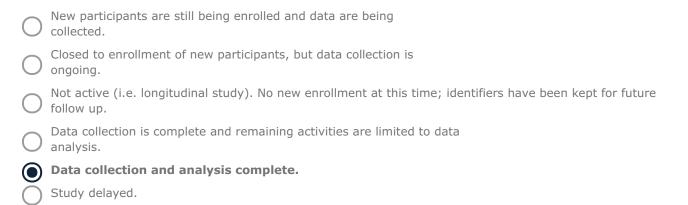
Name of the Principal Investigator (PI)

Jane Researcher

Study cancelled

If you are not the PI on this study, please enter your name and email address below.

Please indicate the status of this project:





Signed consent documents must be maintained for a minimum of 3 years after the completion of the study. Has other identifying information (i.e. names, addresses, dates of birth, link lists, A#s, etc.) been destroyed?

Yes, identifiers have been destroyed.

No, we still have identifiers.

We never collected identifiers.

Approximate destruction date:

5.1.22

If there is any additional information you would like the IRB office to know about identifiers in your study, please explain below.

As per the consent form, we are keeping audio recordings of people who allowed us to for future research. A new protocol will be submitted in Kuali when we are ready to use those recordings in another study.

Did you collect video, audio, or photographs for use in the research?

	Yes
$\bigcirc$	No

Video, audio, and photographs can be de-identified by blurring faces and distorting voices. Have these records been de-identified or destroyed?



If there is any additional information you would like the IRB to know about these sorts of records in your study, please explain below.

See above.

Did you collect bio-specimens?

Yes

Are you	request	ing to	close	this	study?
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## [Learn More about Closing Protocols]

$oldsymbol{O}$	I would like to close this study.
$\bigcirc$	I would like to keep this study open.

Have additional risks or benefits to participants been identified since the last review?



Have any grievances, complaints, or adverse events been identified since the last review?

$\bigcirc$	Yes
$\bigcirc$	No

Since the last review, have any participants withdrawn from the research?

$\bigcirc$	Yes
	No

Since the last review, have you discovered additional information or risks that could influence an individual's willingness to participate?

$\bigcirc$	Yes
	No

How many publications have resulted from this research?

1

Have there been any modifications or revisions in the methods, procedures, or participant population that have not already been approved by the IRB?

$\bigcirc$	Yes
	No

Have research assistants been added or removed without approval through an amendment?



Do all members of the research team have current CITI certifications? Check here.

NOTE: The renewal of your study will not be approved if active members of your research team have expired CITI training.



Please enter the CITI Training expiration date for all members of your research team, including yourself.

Format your response as [Name: Expiration Date]

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Researcher 1: 12/31/24; Researcher 2: 6/22/25; Researcher 3: 7/7/23; Researcher 4: 7/22/25; Researcher 5: 5/15/24
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Total Number of Participants Accrued (Please specify numbers for each population, as applicable)

35

If you have accrued a substantial number of participants (15%) over or under the number that is approved in your protocol, please explain why in the field below.

Forced to stop due to COVID-19. Re-evaluated study purposes and determined that we would cease data collection for this project. Accrual was lower than anticipated (35 actual; 80 anticipated). A reportable event was submitted on 7/5/22 to document the deviation.

Is there an active protocol for this project at another institution or agency?



Please write a brief summary of key research findings. Include a list of publications if applicable.

Individuals with dysarthria did not modify their speech production to align more closely with a healthy model. Speech production modifications are predicted by communicative participation.