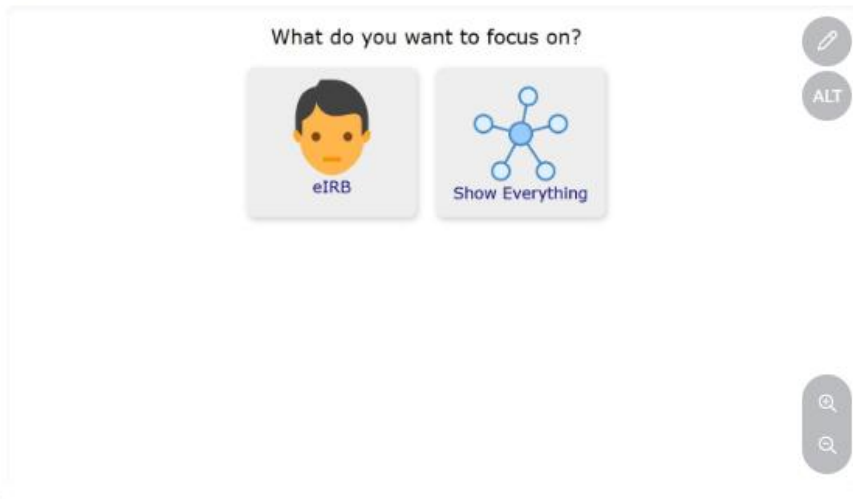


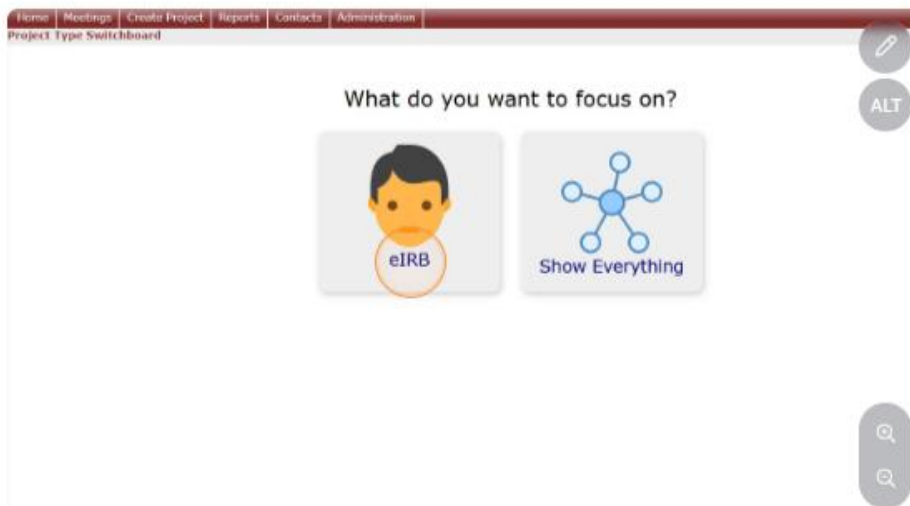
How To Submit An IRB Continuing Review Application

Learn the process for submitting an IRB continuing review through the eIRB portal.

- 1 After successfully logging into eIRB you should see this landing page.



- 2 Click "eIRB"



3

Click "Projects"

The screenshot shows the eIRB system interface. At the top, there is a navigation bar with tabs: Home, Meetings, Create Project, Reports, Contacts, and Administration. Below this, the 'My eIRB' section displays several metrics: 0 Reviews, Me (user profile), 15 Projects (highlighted with an orange circle), 9 xForms, and 24 Events. There are also buttons for 'Search Projects', 'Export to Excel', and 'Start xForm'. Below the metrics, there are two tabs: '14 PI' and '2 Faculty Advisor'. The main area displays a grid of project cards, each with a title, status, expiration date, and a brief description. The projects listed are:

- 99999-UTRGV**: New From PI, Exp 05/27/2027, Allen, Eric, IRB General Testing
- IRB-2023-TEST-UTRGV**: New From PI, Exp 06/13/2027, Allen, Eric, Demonstration Project
- IRB-2026-002-UTH**: Open, Exp Exempt, Allen, Eric, IRB Testing Initial Exempt - Funding
- IRB-2026-003-UTRGV**: Open, Exp Exempt, Allen, Eric, IRB Testing Initial Exempt - No Funding
- IRB-2026-004-UTRGV**: Open, Exp 03/30/2027, Allen, Eric, IRB Testing Initial Expedited CR - Funding
- IRB-2026-005-UTRGV**: Open, Exp 04/08/2027, Allen, Eric, IRB Testing Initial Expedited CR - No Funding
- IRB-2026-006-UTRGV**: Open, Exp Expedited Expiration, Allen, Eric, IRB Testing Initial Expedited No CR - With Funding
- IRB-2026-007-UTRGV**: Open, Exp Expedited Expiration, Allen, Eric, IRB Testing Initial Expedited No CR - No Funding
- IRB-2026-008-UTRGV**: Open, Exp 04/12/2027, Allen, Eric, IRB Testing Initial Full Board CR Funding

4

Note only click "projects" once

This screenshot is identical to the one above, showing the eIRB system interface. The 'Projects' tab is highlighted with an orange circle. The dashboard metrics and project list are the same as in the previous screenshot.

- 5 Click on the project you wish to submit a continuing review for "IRB-2023-TEST-UTRGV".

Home Meetings Create Project Reports Contacts Administration eIRB

My eIRB

0 Reviews 15 Projects 9 xForms 24 Events

Search Projects Export to Excel Start xForm

14 PI 2 Faculty Advisor

99999-UTRGV New From PI Exp 05/27/2027 Allen, Eric IRB General Testing	IRB-2023-TEST-UTRGV New From PI Exp 06/13/2027 Allen, Eric Demonstration Project	IRB-2026-002-UTH Open Exp Exempt Allen, Eric IRB Testing Initial Exempt - Funding
IRB-2026-003-UTRGV Open Exp Exempt Allen, Eric IRB Testing Initial Exempt - No Funding	IRB-2026-004-UTRGV Open Exp 03/30/2027 Allen, Eric IRB Testing Initial Expedited CR - Funding	IRB-2026-005-UTRGV Open Exp 04/08/2027 Allen, Eric IRB Testing Initial Expedited CR - No Funding
IRB-2026-006-UTRGV Open Exp Expedited Expiration Allen, Eric IRB Testing Initial Expedited No CR - With Funding	IRB-2026-007-UTRGV Open Exp Expedited Expiration Allen, Eric IRB Testing Initial Expedited No CR - No Funding	IRB-2026-008-UTRGV Open Exp 04/12/2027 Allen, Eric IRB Testing Initial Full Board CR Funding

- 6 Click "xForms"

ACCORDS

Project
Update
Add Contact
Add Project-Site
Project-Site
Update
Add Attachment
Add Contact
Add Event
Add Note
Add Related Project-Site
Add Animal
Expirations
Generate Doc
Send Email
Start xForm
xForms (3)
Misc
Contact History
Doc Templates
Notifications
Run Project Report
Run Project-Site Report
Project Audit
Project Sub Screen
Project-Site Audit
Project-Site Sub Screen
Deleted Events
Done
Recent Items
IRB-2023-TEST-

Project: **IRB-2023-TEST**
Committee: IRB
Category:
Department: College of Fine Arts
Agent Types: Focus Groups
Title: Demonstration Project
Additional Risk: No
Related Information:
Expedited Categories: Category 6. Collection of data from voice, video, digital or image recordings made for research purposes.
Funding Status: Yes
International Research Status: No
IRB Clinical Research Status: No
IRB Project Review Type: Expedited
Name of External IRB:
Name of Site Relying on UTRGV:
Regulatory Oversight: HHS
Thesis/Dissertation Status: None of the Above
Project-Site
Site(s): **UTRGV - University of Texas Rio Grande Valley**
Status: New From PI
Approval: June 14, 2025 for 12 months
Initial Approval: June 14, 2025
Additional

Sponsor(s): DOE (Primary)
Sponsor Id:
Grants:
CRO:
Year: 2026
Exempt Categories:
External IRB FWA Number:
International Research Locations:
IRB approved number of participants:
IRB Clinical Trial Status:
Multisite Status:
Name of Other Funding Source:
Other Committee Oversight Status:
Study Populations:
VA Status:

7 Click "IRB Continuing Review"

The University of Texas
Rio Grande Valley

Home Meetings Create Project Reports Contacts Administration Find Project (Ctrl+F)

Forms on Project-Site IRB-2023-TEST-UTRGV

Help Filter: 

Actions
Set as move target
Show Deleted
Done

Recent Items
IRB-2023-TEST-UTRGV
99999-UTRGV
Allen, Eric
Sanchez, Jillian
IRB
IRB-2026-011-UTRGV
IRB-2026-010-UTRGV

My Docs & xForms
0 Attachments
11 xForms

Associated xForms

Action	Form	Identifier	Stage/Status	Started	Submitted
 	IRB Continuing Review	Demonstration Project	Data Entry	23 hours ago	
	IRB Reportable Event Form	Demonstration Project	PI Review	Tuesday at 6:15 PM ET	Res
	IRB Amendment Form	Demonstration Project	PI Review	05/29/2026 at 6:24 PM ET	Res

8 Verify information on this page then click "next" this button.

Project Number Add Note View Audit
IRB-2023-TEST

Title Add Note View Audit
Demonstration Project

PI Add Note View Audit
Allen, Eric
Email: eric.allen@utrgv.edu Phone:

Faculty Advisor Add Note View Audit
Note: if blank, there is not faculty advisor for this project
Name Role

Display Other Committee Oversight Status Add Note View Audit

[Next](#) [Save for Later](#) [More](#)

Copyright ©2000-2026 Tech Software. All Rights Reserved.
2026.5.8472.0/Release/1030d2c | GCWAWST | 2026-06-05 19:23:23Z | 0.388s
Powered By  IRBManager

- 9 Select the appropriate option for your research project, in this example I chose "Enrollment and data collection continues".

The University of Texas
Rio Grande Valley

Collaborators

Continuing Review Details

Page 2 of 2

IRB Continuing Review -- Continuing Review Details

Status of the Project (Required) Add Note View Audit

Study related interventions include blood draws, study treatment, questionnaires, or etc.

☒ Enrollment and data collection continues

☐ Enrollment is complete but participants are still receiving study related interventions

☐ De-identified data analysis is ongoing

☐ Paused

☐ Completed

Please attach the informed consent form(s) (i.e. parental permission or child assent) currently being used for this project. Add Note View Audit

Note: If your study does not have any informed consent form(s) skip this question.

Add Attachment

Does this project involve an FDA regulated drug or device? (Required) Add Note View Audit

☐ Yes

☒ No

Does the research dataset contain identifiers? (Required) Add Note View Audit

☐ Yes

☒ No

Additional Continuing Review Details Add Note View Audit

Have any of the approved members of the research team been removed from the study? (Required)

☐ Yes

☐ No

Have there been any changes in funding? (Required)

- 10 Due to the previous response you will need to attach a copy of the consent documents, click "add attachment".

The University of Texas
Rio Grande Valley

Collaborators

Continuing Review Details

Page 2 of 2

IRB Continuing Review -- Continuing Review Details

Status of the Project (Required) Add Note View Audit

Study related interventions include blood draws, study treatment, questionnaires, or etc.

☒ Enrollment and data collection continues

☐ Enrollment is complete but participants are still receiving study related interventions

☐ De-identified data analysis is ongoing

☐ Paused

☐ Completed

Please attach the informed consent form(s) (i.e. parental permission or child assent) currently being used for this project. Add Note View Audit

Note: If your study does not have any informed consent form(s) skip this question.

Add Attachment

Does this project involve an FDA regulated drug or device? (Required) Add Note View Audit

☐ Yes

☒ No

Does the research dataset contain identifiers? (Required) Add Note View Audit

☐ Yes

☒ No

Additional Continuing Review Details Add Note View Audit

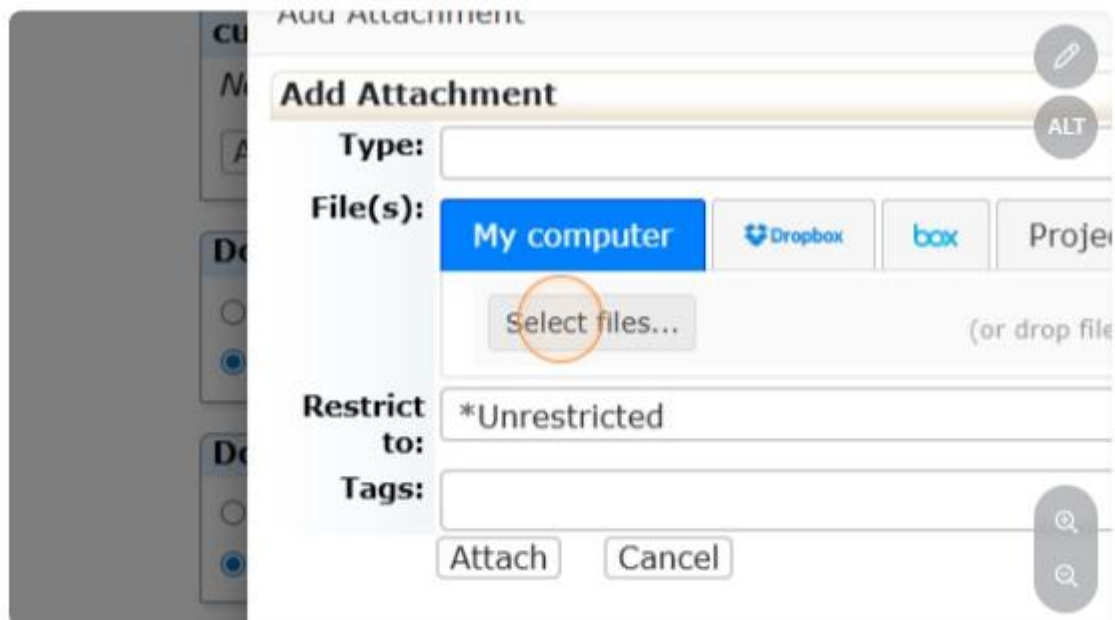
Have any of the approved members of the research team been removed from the study? (Required)

☐ Yes

☐ No

Have there been any changes in funding? (Required)

- 11 Select the appropriate file from your computer by clicking "Select files..."



Add Attachment

Type:

File(s): **My computer**   Project

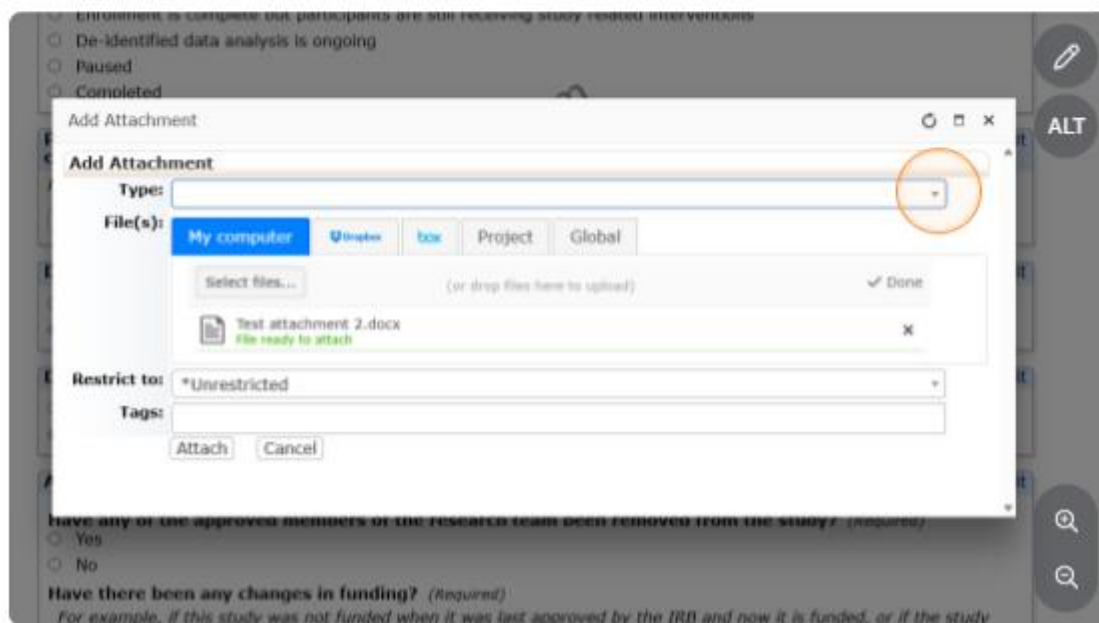
Select files... (or drop files here to upload)

Restrict to: *Unrestricted

Tags:

Attach **Cancel**

- 12 At the end of the "Type" field click the down arrow and select the appropriate title for your attachment.



Add Attachment

Type: 

File(s): **My computer**   Project Global

Select files... (or drop files here to upload) **Done**

 Test attachment 2.docx
File ready to attach **X**

Restrict to: *Unrestricted

Tags:

Attach **Cancel**

13 Click "Attach"

Add Attachment

Type: **Consent Form**

File(s): **My computer** Project Global

Select files... (or drop files here to upload)

Test attachment 2.docx
File ready to attach

Restrict to: *Unrestricted

Tags:

Attach Cancel

14 Select the appropriate answer, in this example I selected "No".

Draft Form

☒ Enrollment and data collection continues
☐ Enrollment is complete but participants are still receiving study related interventions
☐ De-identified data analysis is ongoing
☐ Paused
☐ Completed

Please attach the informed consent form(s) (i.e., parental permission or child assent) currently being used for this project. Add Note View Audit
Note: If your study does not have any informed consent form(s) skip this question.
Add Attachment
 Test attachment 2.docx Consent Form

Does this project involve an FDA regulated drug or device? (Required) Add Note View Audit
☐ Yes
☒ No

Does the research dataset contain identifiers? (Required) Add Note View Audit
☐ Yes
☒ No

Additional Continuing Review Details Add Note View Audit
Have any of the approved members of the research team been removed from the study? (Required)
☐ Yes
☐ No
Have there been any changes in funding? (Required)
For example, if this study was not funded when it was last approved by the IRB and now it is funded, or if the study was funded when it was last approved by the IRB and the award period has lapsed. For the purpose of this question, No Cost-Extensions are not considered a change in funding.
☐ Yes
☐ No

17 Select the appropriate answer, in this example I selected "No".

☐ Yes
☒ No

Does the research dataset contain identifiers? (Required) [Add Note](#) [View Audit](#)

☒ Yes
☐ No

Additional Continuing Review Details [Add Note](#) [View Audit](#)

Have any of the approved members of the research team been removed from the study? (Required)

☐ Yes
☒ No

Have there been any changes in funding? (Required)
For example, if this study was not funded when it was last approved by the IRB and now it is funded, or if the study was funded when it was last approved by the IRB and the award period has lapsed. For the purpose of this question, No Cost-Extensions are not considered a change in funding.

☐ Yes
☒ No

Have study activities been conducted since this study's last approval date? (Required)

☐ Yes
☐ No

Does/did your study involve enrollment of subjects at any point? (Required)
This question refers to whether your project planned enrollment since the beginning of the study.

☐ Yes
☐ No

Have any subjects been withdrawn from study at any time since the initial approval? (Required)

☐ Yes
☐ No

Were there any subject deaths during the course of the study? (Required)

☐ Yes
☐ No

18 Select the appropriate answer, in this example I selected "Yes".

The University of Texas
Rio Grande Valley

Collaborators

Continuing Review Details

Page 2 of 2

IRB Continuing Review -- Continuing Review Details

Have any of the approved members of the research team been removed from the study? (Required)

☐ Yes
☒ No

Have there been any changes in funding? (Required)
For example, if this study was not funded when it was last approved by the IRB and now it is funded, or if the study was funded when it was last approved by the IRB and the award period has lapsed. For the purpose of this question, No Cost-Extensions are not considered a change in funding.

☐ Yes
☒ No

Have study activities been conducted since this study's last approval date? (Required)

☒ Yes
☐ No

Does/did your study involve enrollment of subjects at any point? (Required)
This question refers to whether your project planned enrollment since the beginning of the study.

☐ Yes
☐ No

Have any subjects been withdrawn from study at any time since the initial approval? (Required)

☐ Yes
☐ No

Were there any subject deaths during the course of the study? (Required)

☐ Yes
☐ No

Since the last IRB approval, have you had any problems recruiting subjects? (Required)

☐ Yes
☐ No

Have there been any reportable events since the last continuing review? (Required)
A reportable event includes any meeting the following definition: Any untoward or unfavorable medical

19



+

20

ALT

21 Select the appropriate answer, in this example I selected "No".

Provide a narrative summary of interim findings from the data collected since the last IRB approval. (Required)

This question refers to the findings obtained from the data you collected as part of the research procedures of this study. Findings must be documented in study records and supporting data must be available for review as part of the post approval monitoring.

So far the preliminary data,....

Does/did your study involve enrollment of subjects at any point? (Required)

This question refers to whether your project planned enrollment since the beginning of the study.

☒ Yes

☐ No

Is this a multi-site study in which UTRGV is the IRB of Record and the other sites' IRBs are relying on UTRGV IRB? (Required)

A multi-site research study is when there are many participating sites, each institution is enrolling its own subjects and carrying out the protocol at its own site. For non-exempt human subjects research that includes multiple sites, cooperative agreements will be needed (i.e., Single IRB or Reliance Agreement).

☐ Yes

☒ No

Have any subjects been withdrawn from study at any time since the initial approval? (Required)

☐ Yes

☐ No

Were there any subject deaths during the course of the study? (Required)

☐ Yes

☐ No

Since the last IRB approval, have you had any problems recruiting subjects? (Required)

☐ Yes

☐ No

Have there been any reportable events since the last continuing review? (Required)

22 Provide the appropriate answer.

Does/did your study involve enrollment of subjects at any point? (Required)

This question refers to whether your project planned enrollment since the beginning of the study.

☒ Yes

☐ No

Is this a multi-site study in which UTRGV is the IRB of Record and the other sites' IRBs are relying on UTRGV IRB? (Required)

A multi-site research study is when there are many participating sites, each institution is enrolling its own subjects and carrying out the protocol at its own site. For non-exempt human subjects research that includes multiple sites, cooperative agreements will be needed (i.e., Single IRB or Reliance Agreement).

☐ Yes

☒ No

How many subjects have enrolled and participated since this study was first initiated? (Required)

How many subjects were enrolled since the last approval date? (Required)

Have any subjects been withdrawn from study at any time since the initial approval? (Required)

☐ Yes

☐ No

Were there any subject deaths during the course of the study? (Required)

☐ Yes

☐ No

Since the last IRB approval, have you had any problems recruiting subjects? (Required)

☐ Yes

☐ No

Have there been any reportable events since the last continuing review? (Required)

A reportable event includes any event meeting the following definition: Any untoward or unfavorable medical occurrence in a human subject, including any abnormal sign (for example, abnormal physical exam or laboratory finding), symptom, or disease, temporally associated with the subject's participation in research, whether or not considered related to the subject's participation in the research. Reportable events encompass both observed and

23 Provide the appropriate answer.

Does/did your study involve enrollment of subjects at any point? (Required)
This question refers to whether your project planned enrollment since the beginning of the study.

☒ Yes
☐ No

Is this a multi-site study in which UTRGV is the IRB of Record and the other sites' IRBs are relying on UTRGV IRB? (Required)
A multi-site research study is when there are many participating sites, each institution is enrolling its own subjects and carrying out the protocol at its own site. For non-exempt human subjects research that includes multiple sites, cooperative agreements will be needed (i.e., Single IRB or Reliance Agreement).

☐ Yes
☒ No

How many subjects have enrolled and participated since this study was first initiated? (Required)
13

How many subjects were enrolled since the last approval date? (Required)
1

Have any subjects been withdrawn from study at any time since the initial approval? (Required)
☐ Yes
☒ No

Were there any subject deaths during the course of the study? (Required)
☐ Yes
☒ No

Since the last IRB approval, have you had any problems recruiting subjects? (Required)
☐ Yes
☒ No

Have there been any reportable events since the last continuing review? (Required)
A reportable event includes any event meeting the following definition: Any untoward or unfavorable medical occurrence in a human subject, including any abnormal sign (for example, abnormal physical exam or laboratory finding), symptom, or disease, temporally associated with the subject's participation in research, whether or not considered related to the subject's participation in the research. Reportable events encompass both physical and psychological harms. They occur most commonly in the context of biomedical research, although on occasion, they can occur in the context of behavioral and social research.

☐ Yes
☒ No

+

24 Select the appropriate answer, in this example I selected "Yes".

Does/did your study involve enrollment of subjects at any point? (Required)
This question refers to whether your project planned enrollment since the beginning of the study.

☒ Yes
☐ No

Is this a multi-site study in which UTRGV is the IRB of Record and the other sites' IRBs are relying on UTRGV IRB? (Required)
A multi-site research study is when there are many participating sites, each institution is enrolling its own subjects and carrying out the protocol at its own site. For non-exempt human subjects research that includes multiple sites, cooperative agreements will be needed (i.e., Single IRB or Reliance Agreement).

☐ Yes
☒ No

How many subjects have enrolled and participated since this study was first initiated? (Required)
13

How many subjects were enrolled since the last approval date? (Required)
1

Have any subjects been withdrawn from study at any time since the initial approval? (Required)
☒ Yes
☐ No

Were there any subject deaths during the course of the study? (Required)
☐ Yes
☒ No

Since the last IRB approval, have you had any problems recruiting subjects? (Required)
☐ Yes
☒ No

Have there been any reportable events since the last continuing review? (Required)
A reportable event includes any event meeting the following definition: Any untoward or unfavorable medical occurrence in a human subject, including any abnormal sign (for example, abnormal physical exam or laboratory finding), symptom, or disease, temporally associated with the subject's participation in research, whether or not considered related to the subject's participation in the research. Reportable events encompass both physical and psychological harms. They occur most commonly in the context of biomedical research, although on occasion, they can occur in the context of behavioral and social research.

☐ Yes
☒ No

25 Provide the appropriate response.

Does/did your study involve enrollment of subjects at any point? (Required)

This question refers to whether your project planned enrollment since the beginning of the study.

- ☒ Yes
☐ No

Is this a multi-site study in which UTRGV is the IRB of Record and the other sites' IRBs are relying on UTRGV IRB? (Required)

A multi-site research study is when there are many participating sites, each institution is enrolling its own subjects and carrying out the protocol at its own site. For non-exempt human subjects research that includes multiple sites, cooperative agreements will be needed (i.e., Single IRB or Reliance Agreement).

- ☐ Yes
☒ No

How many subjects have enrolled and participated since this study was first initiated? (Required)

13

How many subjects were enrolled since the last approval date? (Required)

3

Have any subjects been withdrawn from study at any time since the initial approval? (Required)

- ☒ Yes
☐ No

Subjects Withdrawn Since Study Start (Required)

1

Subjects Withdrawn Since Last Approval (Required)

0

Please enter the number of subjects that voluntarily withdrew from the study. (Required)

0

Please summarize the reasons why subjects withdrew or were withdrawn by the investigator from the study. (Required)

Please do not include any direct or indirect identifying information about the subjects.

+

26 Provide the appropriate response.

Does/did your study involve enrollment of subjects at any point? (Required)

This question refers to whether your project planned enrollment since the beginning of the study.

- ☒ Yes
☐ No

Is this a multi-site study in which UTRGV is the IRB of Record and the other sites' IRBs are relying on UTRGV IRB? (Required)

A multi-site research study is when there are many participating sites, each institution is enrolling its own subjects and carrying out the protocol at its own site. For non-exempt human subjects research that includes multiple sites, cooperative agreements will be needed (i.e., Single IRB or Reliance Agreement).

- ☐ Yes
☒ No

How many subjects have enrolled and participated since this study was first initiated? (Required)

13

How many subjects were enrolled since the last approval date? (Required)

3

Have any subjects been withdrawn from study at any time since the initial approval? (Required)

- ☒ Yes
☐ No

Subjects Withdrawn Since Study Start (Required)

1

Subjects Withdrawn Since Last Approval (Required)

0

Please enter the number of subjects that voluntarily withdrew from the study. (Required)

0

Please summarize the reasons why subjects withdrew or were withdrawn by the investigator from the study. (Required)

Please do not include any direct or indirect identifying information about the subjects.

27 Provide the appropriate response.

Does/did your study involve enrollment of subjects at any point? (Required)

This question refers to whether your project planned enrollment since the beginning of the study.

☒ Yes

☐ No

Is this a multi-site study in which UTRGV is the IRB of Record and the other sites' IRBs are relying on UTRGV IRB? (Required)

A multi-site research study is when there are many participating sites, each institution is enrolling its own subjects and carrying out the protocol at its own site. For non-exempt human subjects research that includes multiple sites, cooperative agreements will be needed (i.e., Single IRB or Reliance Agreement).

☐ Yes

☒ No

How many subjects have enrolled and participated since this study was first initiated? (Required)

13

How many subjects were enrolled since the last approval date? (Required)

3

Have any subjects been withdrawn from study at any time since the initial approval? (Required)

☒ Yes

☐ No

Subjects Withdrawn Since Study Start (Required)

1

Subjects Withdrawn Since Last Approval (Required)

0

Please enter the number of subjects that voluntarily withdrew from the study. (Required)

0

Please summarize the reasons why subjects withdrew or were withdrawn by the investigator from the study. (Required)

Please do not include any direct or indirect identifying information about the subjects.

28 Click in the textbox and provide the appropriate response.

How many subjects have enrolled and participated since this study was first initiated? (Required)

13

How many subjects were enrolled since the last approval date? (Required)

3

Have any subjects been withdrawn from study at any time since the initial approval? (Required)

☒ Yes

☐ No

Subjects Withdrawn Since Study Start (Required)

1

Subjects Withdrawn Since Last Approval (Required)

0

Please enter the number of subjects that voluntarily withdrew from the study. (Required)

0

Please summarize the reasons why subjects withdrew or were withdrawn by the investigator from the study. (Required)

Please do not include any direct or indirect identifying information about the subjects.

Were there any subject deaths during the course of the study? (Required)

☐ Yes

☐ No

Since the last IRB approval, have you had any problems recruiting subjects? (Required)

☐ Yes

☐ No

Have there been any reportable events since the last continuing review? (Required)

A reportable event includes any event meeting the following definition: Any untoward or unfavorable medical occurrence in a human subject, including any abnormal sign (for example, abnormal physical exam or laboratory

29 Select the appropriate answer, in this example I selected "No".

4

Subjects Withdrawn Since Last Approval (Required)

0

Please enter the number of subjects that voluntarily withdrew from the study. (Required)

0

Please summarize the reasons why subjects withdrew or were withdrawn by the investigator from the study. (Required)

Please do not include any direct or indirect identifying information about the subjects.

The participant relocated.....

Were there any subject deaths during the course of the study? (Required)

☐ Yes
 ☒ No

Since the last IRB approval, have you had any problems recruiting subjects? (Required)

☐ Yes
 ☒ No

Have there been any reportable events since the last continuing review? (Required)

A reportable event includes any event meeting the following definition: Any untoward or unfavorable medical occurrence in a human subject, including any abnormal sign (for example, abnormal physical exam or laboratory finding), symptom, or disease, temporally associated with the subject's participation in research, whether or not considered related to the subject's participation in the research. Reportable events encompass both physical and psychological harms. They occur most commonly in the context of biomedical research, although on occasion, they can occur in the context of social and behavioral research. Reportable events can also include incidents such as a breach of confidentiality due to the loss of laptop including encrypted data etc.

☐ Yes
 ☒ No

Have there been any protocol deviations, no matter how minor, to any part of this study including the IRB approved forms? (Required)

Note: A protocol deviation is unplanned or unapproved changes to the IRB approved research procedures.

☐ Yes
 ☒ No

30 Select the appropriate answer, in this example I selected "No".

Please enter the number of subjects that voluntarily withdrew from the study. (Required)

0

Please summarize the reasons why subjects withdrew or were withdrawn by the investigator from the study. (Required)

Please do not include any direct or indirect identifying information about the subjects.

The participant relocated.....

Were there any subject deaths during the course of the study? (Required)

☐ Yes
 ☒ No

Since the last IRB approval, have you had any problems recruiting subjects? (Required)

☐ Yes
 ☒ No

Have there been any reportable events since the last continuing review? (Required)

A reportable event includes any event meeting the following definition: Any untoward or unfavorable medical occurrence in a human subject, including any abnormal sign (for example, abnormal physical exam or laboratory finding), symptom, or disease, temporally associated with the subject's participation in research, whether or not considered related to the subject's participation in the research. Reportable events encompass both physical and psychological harms. They occur most commonly in the context of biomedical research, although on occasion, they can occur in the context of social and behavioral research. Reportable events can also include incidents such as a breach of confidentiality due to the loss of laptop including encrypted data etc.

☐ Yes
 ☒ No

Have there been any protocol deviations, no matter how minor, to any part of this study including the IRB approved forms? (Required)

Note: A protocol deviation is unplanned or unapproved changes to the IRB approved research procedures.

☐ Yes
 ☒ No

Provide a narrative summary of benefits experienced by participants in the next year. (Required)

31 Select the appropriate answer, in this example I selected "No".

Were there any subject deaths during the course of the study? (Required)

☐ Yes

☒ No

Since the last IRB approval, have you had any problems recruiting subjects? (Required)

☐ Yes

☒ No

Have there been any reportable events since the last continuing review? (Required)

A reportable event includes any event meeting the following definition: Any untoward or unfavorable medical occurrence in a human subject, including any adverse event (for example, abnormal physical exam or laboratory finding), symptom, or disease, temporally associated with the subject's participation in research, whether or not considered related to the subject's participation in the research. Reportable events encompass both physical and psychological harms. They occur most commonly in the context of biomedical research, although on occasion, they can occur in the context of social and behavioral research. Reportable events can also include incidents such as a breach of confidentiality due to the loss of laptop including encrypted data etc.

☐ Yes

☒ No

Have there been any protocol deviations, no matter how minor, to any part of this study including the IRB approved forms? (Required)

Note: A protocol deviation is unplanned or unapproved changes to the IRB approved research procedures.

☐ Yes

☒ No

Provide a narrative summary of benefits experienced by participants in the past year. (Required)

Note: If participants did not experience any benefits, please state that here.

Does/did your study involve analysis of records, data or biological specimens at any point? (Required)

This question refers to the use of information about subjects from records, data or biospecimens that are either existing or have been prospectively collected as part of your study (i.e., medical charts, educational records, etc.).

☐ Yes

☒ No

32 Select the appropriate answer, in this example I selected "No".

☒ No

Since the last IRB approval, have you had any problems recruiting subjects? (Required)

☐ Yes

☒ No

Have there been any reportable events since the last continuing review? (Required)

A reportable event includes any event meeting the following definition: Any untoward or unfavorable medical occurrence in a human subject, including any adverse event (for example, abnormal physical exam or laboratory finding), symptom, or disease, temporally associated with the subject's participation in research, whether or not considered related to the subject's participation in the research. Reportable events encompass both physical and psychological harms. They occur most commonly in the context of biomedical research, although on occasion, they can occur in the context of social and behavioral research. Reportable events can also include incidents such as a breach of confidentiality due to the loss of laptop including encrypted data etc.

☐ Yes

☒ No

Have there been any protocol deviations, no matter how minor, to any part of this study including the IRB approved forms? (Required)

Note: A protocol deviation is unplanned or unapproved changes to the IRB approved research procedures.

☐ Yes

☒ No

Provide a narrative summary of benefits experienced by participants in the past year. (Required)

Note: If participants did not experience any benefits, please state that here.

Does/did your study involve analysis of records, data or biological specimens at any point? (Required)

This question refers to the use of information about subjects from records, data or biospecimens that are either existing or have been prospectively collected as part of your study (i.e., medical charts, educational records, etc.).

☐ Yes

☒ No

33 Click in the textbox and provide the appropriate response.

finding), symptom, or disease, temporally associated with the subject's participation in research, whether or not considered related to the subject's participation in the research. Reportable events encompass both physical and psychological harms. They occur most commonly in the context of biomedical research, although on occasion, they can occur in the context of social and behavioral research. Reportable events can also include incidents such as a breach of confidentiality due to the loss of laptop including encrypted data etc.

☐ Yes
☒ No

Have there been any protocol deviations, no matter how minor, to any part of this study including the IRB approved forms? (Required)
Note: A protocol deviation is unplanned or unapproved changes to the IRB approved research procedures.

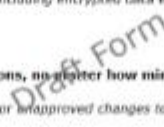
☐ Yes
☒ No

Provide a narrative summary of benefits experienced by participants in the past year. (Required)
Note: If participants did not experience any benefits, please state that here.

Does/did your study involve analysis of records, data or biological specimens at any point? (Required)
This question refers to the use of information about subjects from records, data or biospecimens that are either existing or have been prospectively collected as part of your study (i.e., medical charts, educational records, etc.).

☐ Yes
☒ No

Please summarize any annual reports, multi-center reports or similar documents. This includes any Data Safety Monitoring reports, if applicable. (Required)
Note: If not applicable please type N/A.








+

34 Select the appropriate answer, in this example I selected "No".

psychological harms. They occur most commonly in the context of biomedical research, although on occasion, they can occur in the context of social and behavioral research. Reportable events can also include incidents such as a breach of confidentiality due to the loss of laptop including encrypted data etc.

☐ Yes
☒ No

Have there been any protocol deviations, no matter how minor, to any part of this study including the IRB approved forms? (Required)
Note: A protocol deviation is unplanned or unapproved changes to the IRB approved research procedures.

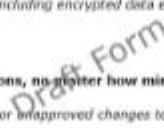
☐ Yes
☒ No

Provide a narrative summary of benefits experienced by participants in the past year. (Required)
Note: If participants did not experience any benefits, please state that here.

Does/did your study involve analysis of records, data or biological specimens at any point? (Required)
This question refers to the use of information about subjects from records, data or biospecimens that are either existing or have been prospectively collected as part of your study (i.e., medical charts, educational records, etc.).

☐ Yes
☒ No

Please summarize any annual reports, multi-center reports or similar documents. This includes any Data Safety Monitoring reports, if applicable. (Required)
Note: If not applicable please type N/A.








- 35 Click in the textbox and provide the appropriate response.

psychological harms. They occur most commonly in the context of biomedical research, although on occasion, they can occur in the context of social and behavioral research. Reportable events can also include incidents such as a breach of confidentiality due to the loss of laptop including encrypted data etc.

☐ Yes
☒ No

Have there been any protocol deviations, no matter how minor, to any part of this study including the IRB approved forms? (Required)
Note: A protocol deviation is unplanned or unapproved changes to the IRB approved research procedures.

☐ Yes
☒ No

Provide a narrative summary of benefits experienced by participants in the past year. (Required)
Note: If participants did not experience any benefits, please state that here.

(Benefits.....)

Does/did your study involve analysis of records, data or biological specimens at any point? (Required)
This question refers to the use of information about subjects from records, data or biospecimens that are either existing or have been prospectively collected as part of your study (i.e., medical charts, educational records, etc.).

☐ Yes
☒ No

Please summarize any annual reports, multi-center reports or similar documents. This includes any Data Safety Monitoring reports, if applicable. (Required)
Note: If not applicable please type N/A.

+

- 36 Select the appropriate answer, in this example I selected "No".

Does/did your study involve analysis of records, data or biological specimens at any point? (Required)
This question refers to the use of information about subjects from records, data or biospecimens that are either existing or have been prospectively collected as part of your study (i.e., medical charts, educational records, etc.).

☐ Yes
☒ No

Please summarize any annual reports, multi-center reports or similar documents. This includes any Data Safety Monitoring reports, if applicable. (Required)
Note: If not applicable please type N/A.

N/A

Have you received any complaints about the study since the last IRB approval? (Required)

☐ Yes
☒ No

Since the last IRB continuing review, has there been any other relevant information and/or recent literature regarding this study, especially information about risks associated with the research procedures? (Required)

☐ Yes
☐ No

Provide an assessment of whether the relationship of risks to potential benefits has changed based on the study results obtained since the IRB last approval. (Required)
This question refers to benefits of the study, such as knowledge gained.

You will need to click "Next" and then "Submit" to send this form to the next stage in the review process. Add Note

- 37 Select the appropriate answer, in this example I selected "No".

☐ Yes
☒ No

Please summarize any annual reports, multi-center reports or similar documents. This includes any Data Safety Monitoring reports, if applicable. (Required)
Note: If not applicable please type N/A.

N/A

Have you received any complaints about the study since the last IRB approval? (Required)
☐ Yes
☒ No

Since the last IRB continuing review, has there been any other relevant information and/or recent literature regarding this study, especially information about risks associated with the research procedures? (Required)
☐ Yes
☒ No

Provide an assessment of whether the relationship of risks to potential benefits has changed based on the study results obtained since the IRB last approval. (Required)
This question refers to benefits of the study, such as knowledge gained.

You will need to click "Next" and then "Submit" to send this form to the next stage in the review process. Add Note

Previous Next Save for Later More ▾

- 38 Click in the textbox and provide the appropriate response.

☐ Yes
☒ No

Please summarize any annual reports, multi-center reports or similar documents. This includes any Data Safety Monitoring reports, if applicable. (Required)
Note: If not applicable please type N/A.

N/A

Have you received any complaints about the study since the last IRB approval? (Required)
☐ Yes
☒ No

Since the last IRB continuing review, has there been any other relevant information and/or recent literature regarding this study, especially information about risks associated with the research procedures? (Required)
☐ Yes
☒ No

Provide an assessment of whether the relationship of risks to potential benefits has changed based on the study results obtained since the IRB last approval. (Required)
This question refers to benefits of the study, such as knowledge gained.

You will need to click "Next" and then "Submit" to send this form to the next stage in the review process. Add Note

Previous Next Save for Later More ▾

- 39 Review the red text, then click "Next".

NOTE: At this appropriate position type N/A.

N/A

Have you received any complaints about the study since the last IRB approval? (Required)

☐ Yes

☒ No

Since the last IRB continuing review, has there been any other relevant information and/or recent literature regarding this study, especially information about risks associated with the research procedures? (Required)

☐ Yes

☒ No

Provide an assessment of whether the relationship of risks to potential benefits has changed based on the study results obtained since the IRB last approval. (Required)

This question refers to benefits of the study, such as knowledge gained.

The research team learned.....

You will need to click "Next" and then "Submit" to send this form to the next stage in the review process. Add Note

Previous **Next** Save for Later More ▾

Copyright © 2000-2020 Tech Software, All Rights Reserved
2000-2020 Tech Software, All Rights Reserved | 2000-2020 Tech Software, All Rights Reserved | 2000-2020 Tech Software, All Rights Reserved
Powered by Tech Software

- 40 Click "Submit" to send the application to the next stage. (e.g. Faculty advisor review, PI review, or IRB review)

The University of New
Rio Grande Valley
Form Complete

Form Completed

You've completed the form. You can now either save the form for later revision, or submit it.

Go Back Save for Later Print **Submit**