

A guide to completing Low and Negligible
risk iRECs modifications for researchers in
the School of Psychology

iRECs MODIFICATION Guide

For researchers submitting
MODIFICATIONS to a project
approved by the HREAP-C
before AUG 2023

HREAP-C Behavioural Sciences

Where do I find out more about iRECs?

Everything provided by UNSW about iRECs should be available here: <https://research.unsw.edu.au/irecs-new-ethics-online-updates>

How do I modify a project that was approved by HREAP-C before AUG 2023

- 1) Email DVCR RECS Human Ethics humanethics@unsw.edu.au to request your application is unlocked for modification.
 - a. Please use BOTH the App number (e.g., 1292) and the number in the project title (e.g., 3232) to minimise confusion.

Request to unlock/modify Human Ethics App 1292: 3232-Complex Decision Making

 Kristy Martire
To: DVCR RECS Human Ethics

[↩ Reply](#) [↩ Reply All](#) [→ Forward](#)

Mon 18/09/2023

Dear RECS,

Could you please unlock Human Ethics Legacy App 1292: 3232-Complex Decision Making for modification.

Thank you,
Kristy


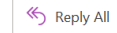
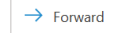

Kristy Martire
Professor

School of Psychology
Room 1007, Level 10, Mathews Building
UNSW SYDNEY 2052

2) Follow the link in your email from RECS to edit the unlocked app

iRECS Application unlocked for modification (3232)

 donotreply@infonetica.net
To: Kristy Martire
Cc: Kristy Martire

 Reply  Reply All  Forward 
Mon 18/09/2023 12:52 PM

Dear Prof Kristy Martire,

Reference Number: 3232
Project Title: 3232 - Complex Decision Making

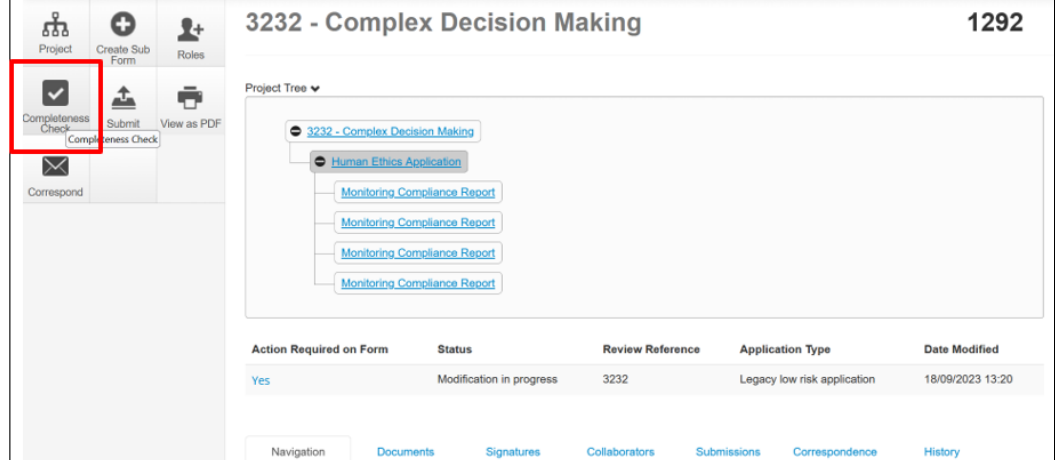
Your Legacy low risk application application has been unlocked and you can now submit your modification in iRECS.

You can access your application modification via <https://irecs.unsw.edu.au/Project/Index/1737>.

Kind regards,

Research Ethics & Compliance Support (RECS)
UNSW Sydney
NSW 2052 AUSTRALIA
E: humanethics@unsw.edu.au

3) Choose 'Completeness check' in the top left hand panel



3232 - Complex Decision Making 1292

Project Tree

- 3232 - Complex Decision Making
 - Human Ethics Application
 - Monitoring Compliance Report
 - Monitoring Compliance Report
 - Monitoring Compliance Report
 - Monitoring Compliance Report

Action Required on Form	Status	Review Reference	Application Type	Date Modified
Yes	Modification in progress	3232	Legacy low risk application	18/09/2023 13:20

Navigation Documents Signatures Collaborators Submissions Correspondence History

4) Choose: 'Legacy Modification Summary'

Completeness Check ✕

Incomplete: Please complete the following questions

- Is there Research Grant Funding associated with this Project?
- Chief Investigator Chief Investigator Appointment Type
- Chief Investigator Responsibilities
- Legacy Clinical Trials
- Legacy Modification Summary
- Participants & Area of Research: Indicate the type of research that will be conducted
- Participants & Area of Research: Indicate the target population for this research
- Access to Existing Collections: Does the research involve access to existing collections of data or biospecimens?
- Legacy Low Risk Modification Declaration

Close

5) Provide the following information in the 'Legacy Modification Summary'

EXISTING APPROVAL:

[Provide a brief description of the approved project to contextualise your modification request.

E.g., We were originally approved to recruit participants from SONA-1. They completed a [specific] task/answered a series of questions etc. The study took 30 mins and participants received 0.5 SONA-1 credits in recompense for their participation.]

REQUESTED CHANGES:

[provide a description of the changes you would like to make to the project.

E.g., We would like to recruit participants from SONA-P; Add a new investigator to the project; Add a new questionnaire; Change the study duration/recompense etc.]

IF A NEW METHOD OF RECRUITMENT IS REQUESTED: See 1-3 here

https://unsw.sharepoint.com/:w:/s/HREAP-C2/EaGwUYTcTH5Lhj4y4_glnr0BTWzCwAXcAo6SYLwAEySfqA

IF A NEW QUESTIONNAIRE OR STIMULUS IS REQUESTED:

[Attach all new questionnaires/ examples of stimuli etc]

IF A NEW DURATION OR RATE OF RECOMPENSE IS REQUESTED:

[Update the Participant Information Statement and Consent Form to include the updated duration/recompense information (tracked or highlighted changes are helpful) and attach.

ETHICAL ISSUES ARISING:
 [All requests: provide details about any possible ethical implications from the proposed changes or explicitly state that “no ethical issues are anticipated”]

6) Upload/attach all relevant supporting documentation here file:

Clean Legacy Project Document Upload

Type	Document Name	File Name	Version Date	Version	Size	View	Delete
PROTOCOL	UPLOAD New_Revised_PISCFS_Stumili_RecruitmentDocs_etc_HERE	UPLOAD New_Revised_PISCFS_Stumili_RecruitmentDocs_etc_HERE.pdf	18/09/2023	1	59.2 KB	Download	Delete

[Upload Document](#)

7) Accept the terms or transfer the modification to the CI so they can accept the terms. (instructions are provided in iRECS for this)

Declaration 0

If you have created this application on behalf of a Chief Investigator (CI), you must transfer this project to the CI to complete the final (Declaration and Submission) steps. Instructions for transferring a project can be found in the help section of iRECS.

As the submitting investigator of the research, I confirm that:

- The information provided in the submission is accurate, correct, and complete.
- I will ensure that the investigators and study personnel conduct the research following the National Statement on Ethical Conduct in Human Research (updated 2018) requirements.
- I will ensure that the investigators and study personnel will not commence recruitment, data collection or access data from existing collections (if applicable) without written confirmation of human ethics approval.
- Qualified research personnel will conduct all research procedures for both training and experience.
- I will ensure that all research personnel follow the approved protocol, procedures, terms, and conditions specified by the HREC/HREAP when conducting this human research once approved.
- I will ensure that approval will be sought for all modifications made to the research before implementing them in the conduct of the research.
- I will ensure that any conditions of approval will be met, and any requisite approvals, permits or regulatory processes relevant to the research will be obtained before recruitment, and data collection commences.
- I have read and understood the applicable UNSW Workplace Health and Safety policies. Therefore, we will undertake all appropriate training in Workplace Health and Safety as dictated by UNSW policies.
- The human research proposal has been provided to the head of the school for approval or their information before submission.

Accept

8) Work your way through the remaining items in the 'completeness check' section. Click the link in the completeness check. Update the info that is requested and then press save. Then go back and forth between the check and the save until it's all complete.

a) Please see the HREAP-C guidance here

<https://acrobat.adobe.com/link/track?uri=urn:aaid:scds:US:b6f8eb7a-d900-3dcf-92c6-eb634e2a1ca9> in relation to Participants & Area of Research.

Completeness Check



Incomplete: Please complete the following questions

- [Is there Research Grant Funding associated with this Project?](#)
- [Chief Investigator Chief Investigator Appointment Type](#)
- [Chief Investigator Responsibilities](#)
- [Legacy Clinical Trials](#)
- [Participants & Area of Research: Indicate the type of research that will be conducted](#)
- [Participants & Area of Research: Indicate the target population for this research](#)
- [Access to Existing Collections: Does the research involve access to existing collections of data or biospecimens?](#)

Close

9) Go back to 'Completeness check' and select 'Legacy Low Risk Modification Declaration'

Completeness Check



Incomplete: Please complete the following questions

- [Is there Research Grant Funding associated with this Project?](#)
- [Chief Investigator Chief Investigator Appointment Type](#)
- [Chief Investigator Responsibilities](#)
- [Legacy Clinical Trials](#)
- [Participants & Area of Research: Indicate the type of research that will be conducted](#)
- [Participants & Area of Research: Indicate the target population for this research](#)
- [Access to Existing Collections: Does the research involve access to existing collections of data or biospecimens?](#)
- [Legacy Low Risk Modification Declaration](#)

Close

10) Choose to 'Submit' from the completeness check window or the top left hand panel.

iRECS (Applicant) Work Area Contacts Help Prof Kristy Martire

Human Ethics Application

Project Title: 3232 - Complex Decision Making

Version: 3.0

Compare with Previous Submission

Submission

Note: Below is some helpful guidance completing this form. Please note that the session will time out after 30 minutes of inactivity. It is advised that you regularly save to ensure no content is lost.

- Save** To save your form progress select the **'Save'** button. Note: iRECS sessions will expire after 30 minutes of inactivity.
- Completeness Check** To perform a completeness check (checks mandatory questions are answered), select the **'Completeness Check'** button.
- Submit** To submit your completed form, select the **'Submit'** button.

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