



Research Ethics and Integrity

Application for Level 2 Approval

This application form should be completed for research that involves a laboratory experiment, field/online survey, any form of contact with healthy adult human participants, or sensitive information (such as, but not restricted to medical, genetic, financial, personnel, criminal or administrative records and test results including academic achievements), but does **not** entail i) potential risk of harm or discomfort to the participants, researchers, or any other individuals; or ii) a lack of informed consent from any individuals involved.

Research should not commence until written approval has been received from the Ethics Committee. This should be borne in mind when setting a start date for the project. Should the research change such that the responses to the questions in this form are no longer applicable, you must seek further ethical approval.

Applications should be made on this form, and submitted electronically to the Convener of the Ethics Committee. Applications will be assessed by the Convener in the first instance, and may then be passed to members of the Ethics Committee at the School of Economics. For a project that uses the BLUE subject pool and/or involves an experiment conducted at the BLUE lab, the application will also be reviewed by the Director of BLUE.

Any necessary supporting documentation should also be attached to this form. (Tick those that apply):

consent form

information and instructions to participants

questionnaire

other - specify:

1. Details of Research and Investigator(s)

1.1. Title:

1.2. Type of project: Staff research project Student research project

1.3. Principal Investigator(s) (students should also include the name of their supervisor):

- Name:
 Affiliation:
- Name:
 Affiliation:
- Name:
 Affiliation:

1.4. Does your research involve collaboration with other academic/non-academic partners or employees?

Yes No

If Yes, what steps will be taken to ensure that all individuals adhere to UoE research ethics and integrity standards?

1.5. Proposed start date and duration:

1.6. Funding:

Is this project externally funded?

Yes No

If Yes, please list the names of the funding bodies.

If your answer above is Yes, does the project comply with the ethics guidelines of the funding bodies?

Yes No

If No, contact the Ethics Convener before submitting your application

1.7. Confirmations:

I confirm I have read the CAHSS guidance on Research Ethics and Integrity

I confirm I have completed the online GDPR and Information Security training

2. Brief description of the proposed project (maximum 200 words)

3. Participants - please fill in if the project involves human participants

3.1. How many participants will be involved in the study?

3.2. Are there any potential risks of harm (physical, psychological, social, legal or economic) or discomfort to participants or researchers associated with the proposed project? These include, but are not restricted to, intentionally upsetting questions, disturbing images, noise, or physical contact.

Yes No

If Yes, please fill in the Level 3 application form.

3.3. Does the data collection involve deception or instructions/questions that intentionally mislead the participants?

Yes No

If Yes, please fill in the Level 3 application form.

3.4. Are the participants from the BLUE subject pool?

Yes No

If No, please answer the following two questions.

3.5. Do the participants include individuals below the age of 18, vulnerable adults, or adults with a limited capacity to give informed consent?

Yes No

If Yes, please fill in the Level 3 application form.

3.6. Will you clearly explain to the participants a) the purpose of the data collection, expected duration, and procedures; b) the right to withdraw from the research at any point and consequences of withdrawal; c) potential risks, discomfort, or other adverse effects; and d) whom to contact for any questions?

Yes No

If No, please fill in the Level 3 application form.

4. Data Management - please fill in if the project involves data that are not publicly available

4.1. Will the stored data be fully anonymised, such that the data do not contain any information that could lead to an individual being identified?

Yes No

4.2. If the answer to the previous question is No, will you a) explain potential identification of the participants' identity to the individuals involved; and b) obtain explicit consent?

Yes No

If No, please fill in the Level 3 application form.

4.3. If the project involves data collection from human participants, please indicate what *personal information* you will collect. E.g. gender, income, political views, race, family background.

4.4. Potential Privacy Risks: Please assess the following risks of participant identification.

Identifiable due to...	Likelihood	Severity of Harm
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Data linkage:

Low participant numbers

Geographical location

Transfer of data

Access of data

Identify measures to reduce or eliminate risks identified as possible/significant or probable/severe.

4.5. It is a requirement of the General Data Protection Regulation to ensure individuals are aware of how information about them will be managed. If the project involves personal data collection from human participants, will you inform the participants that the collected information will only be used for research purposes?

Yes No

If No, please fill in the Level 3 application form.

4.6. How, where, and for how long will the data collected be stored? Who will have access? What data security arrangements will be put in place?

Please provide a comprehensive data management plan, from collection (in any format) to destruction or permanent storage of the data. If you intend to share or make any version of the data publicly available, please refer to it here.

4.7. Will information containing personal, identifiable data be transferred to, shared with, supported by, or otherwise available to third parties outside the European Economic Area?

Yes No

If yes, what safeguard will apply (e.g. US/EU Privacy Shield)?

5. Prevent Duty: does your research concern groups which may be construed as terrorist or extremist?

Yes No

If Yes, please fill in the Level 3 application form.

6. Potential Risks to Researchers

6.1. Researcher safety: does your research imply any risk to your personal safety?

Yes No

If Yes, please explain how you would minimize the risks, as well as details of the insurance you will be covered by, and associated risk assessments that have been conducted.

6.2. To the best of your knowledge, could any institutional or personal conflicts of interest arise from this research?

Yes No

If Yes, please elaborate on the nature of this, and the steps they are taking to ensure the integrity of the research.

7. BLUE Lab: please answer the following questions if you use the BLUE subject pool and/or conduct an experiment at the BLUE Lab.

7.1. Please confirm that you have read and will comply with the BLUE Lab Ethical Guidelines, and the BLUE Lab Usage Guidelines.

Yes No

7.2. Please outline the proposed timeline of the project, such as dates for a pilot(s), the actual experiment/survey, etc.

7.3. Please briefly describe the expected duration and compensation (e.g. average per participant, minimum and maximum). If the participants are incentivized, briefly explain also how the payment is determined.

7.4. (Optional) Please describe the experimental/survey protocol.

8. Any Other Issues Please use the space below for any issues you would like to bring to the attention of the Ethics Committee; or your response to comments and recommendations from the Committee.

ETHICS DECLARATION

I confirm that the proposed research will comply with the Ethics Guidelines of the College of Arts, Humanities and Social Sciences and the School of Economics.

Signed

Date:

Signed (supervisor)

Date:

(student project only)

Please enclose any relevant documentation (consent forms, survey questions, experimental instructions, etc).

APPROVAL

Ethics Committee Member:

Signed

Date:

Ethics Committee Member:

Signed

Date:

APPROVAL BY CONVENER'S ACTION

Ethics Committee Convener:

Signed

Date:

APPROVAL (for use of BLUE lab or BLUE subject pool)

Director of the BLUE lab:

Signed

Date: