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| **Official Use Only**Protocol number |

 

 **OFFICE OF RESEARCH, INNOVATION AND DEVELOPMENT**

 **Ethics Committee for the Humanities (ECH)**

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| **PROTOCOL CONSENT FORM**  |

**HOW TO USE THIS CONSENT FORM:**

This document is the standard ECH-approved template to assist you with designing a written informed consent form. You may adapt the template as you see fit, but remember that **the document must address the participant directly**, have some information about each heading, and include the UG logo at the top. **Please write in SIMPLE, NON-TECHNICAL language**.

The text written in **[RED]** is for **guidance only** and should be removed before finalising the document. **Answers provided under each heading should be in phrases and not bulleted**. Also, **this information box should be deleted before the document is finalised.**

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| **Section A- BACKGROUND INFORMATION** |

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| Title of Study:  | **The title must be consistent on the proposal and all other documents** |
| Principal Investigator: | **For students and researchers, state the lead person’s name**  |
| Certified Protocol Number |  |

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| **Section B– CONSENT TO PARTICIPATE IN RESEARCH** |

**General Information about Research**

* State clearly the purpose of the study in simple words (avoid the use of jargon and technical language. Remember the consent form is for the participant – please do not copy and paste from your research proposal).
* Indicate the expected duration that will be required of participants in the study.
* Give a description of the procedures/methods to be followed and the identification of any experimental procedures and what the participant(s) is supposed to do.

**Benefits/Risks of the study**

* Indicate specifically the benefits and risks associated with the study. Include all physical, social, and psychological risks and benefits anticipated. (*State the direct benefits to participants that could be expected from their participation in the study. If the participants will not benefit directly, clearly state this fact. Also, state (realistically) the potential benefits, if any, to society expected from this study.)*
* Indicate any hazards to participants and what steps will be taken to minimize the risks (e.g., referral for counselling or therapy, etc.)

**Confidentiality**

* Describe the extent to which confidentiality of records identifying the participants will be maintained.
* Indicate all groups that may have direct access to the research records at any particular time; thus, by signing or thumbprinting a written consent form, the participant or their representative is authorizing such access.

**Compensation**

* State clearly if there are any compensation packages either in cash or kind available for participants who participate in the study.
* The exact amount or gift to be given must be clearly spelt out.
* The conditions for receiving the package and when it will be made should also be indicated (a token of appreciation should be given at the end of the study).

**Withdrawal from Study**

* State that participation is voluntary and participants may withdraw at any time without penalty.
* More specifically, state that the participant will not be adversely affected if he/she declines to participate or later withdraws from the study.
* Assure that the participant or the participant's legal representative will be

informed promptly if information becomes available that may be relevant to the

participant's willingness to continue participating in the study or withdraw from the study.

* Any circumstances and/or reasons under which participation may be terminated should be stated clearly.

**Contact for Additional Information**

* Provide contact details of persons assigned to respond to questions about the research and respond to cases of research-related injury.
* Names, Institutional affiliation, addresses, email, and telephone numbers (including mobile numbers) should be made accessible to all participants.
* If you have any questions about your rights as a research participant in this study you may contact the Administrator of the Ethics Committee for Humanities, ISSER, University of Ghana at ech@ug.edu.gh  or 00233- 303-294-0531.

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| **Section C- PARTICIPANT AGREEMENT** |

**"I have read or have had someone read all of the above, asked questions, received answers regarding participation in this study, and am willing to give consent for me, my child/ward to participate in this study. I will not have waived any of my rights by signing this consent form. Upon signing this consent form, I will receive a copy for my personal records."**

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Name of Participant

\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Signature or mark of Participant Date

**If participant can neither read nor understand the form themselves, a witness must sign here:**

I was present while the benefits, risks, and procedures were read to the volunteer. All questions were answered and the volunteer has agreed to take part in the research.

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Name of witness

\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Signature of witness / Mark Date

I certify that the nature and purpose, the potential benefits, and possible risks associated with participating in this research have been explained to the above individual.

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Name of Person who Obtained Consent

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Signature of Person Who Obtained Consent Date