



PROTOCOL CONSENT FORM
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Section A- BACKGROUND INFORMATION
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Title of Study:	
Principal Investigator:	
Certified Protocol Number	

Section B- CONSENT TO PARTICIPATE IN RESEARCH
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Please seek the consent of the research participants by informing them (research participants) about your research using the guide below. Develop your form as would be used on the field.

**General Information about Research**

- State clearly the purpose of the study in easily-understood words (avoid the use of jargons and technical language).
- 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 procedures which are experimental and what the participant(s) is supposed to do.

**Benefits of the study**

Indicate specifically the benefits associated with the study. Include all physical, social and psychological benefits anticipated.

**Risk of the study**

Indicate specifically the risks associated with the study. Include all physical, social and psychological risks anticipated.

## **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 they signing or thumb printing 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

## **Withdrawal from Study**

- State that participation is voluntary and participants may withdraw at any time without any penalty.
- More specifically, state that the participant will not be adversely affected if he/she declines to participate or later stops participating.
- Provide assurance that the participant or the participant's legal representative will be informed in a timely manner if information becomes available that may be relevant to the participant's willingness to continue participation or withdraw.
- Any circumstances and/or reasons under which the participant's participation may be terminated should be stated clearly.

## **Contact for Additional Information**

- This statement should indicate whom to contact for answers to any questions about the research and whom to contact in case of research-related injury.
- Names, addresses and telephone numbers (including mobile numbers) should be made accessible to all participants.
- If you have any issues on your rights as a participant you can contact the address below:

**Administrator, Ethics Committee for Basic and Applied Sciences**  
**College of Basic and Applied Sciences**  
**University of Ghana**  
**P. O. Box LG 68**  
**Legon – Accra**  
**IP No.: 3014**  
**Email: [ethicscbas@ug.edu.gh](mailto:ethicscbas@ug.edu.gh)**

Section C- VOLUNTEER AGREEMENT

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

\_\_\_\_\_  
Name of Volunteer

\_\_\_\_\_  
Signature or mark of volunteer

\_\_\_\_\_  
Date

**If volunteers cannot read 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.

\_\_\_\_\_  
Name of witness

\_\_\_\_\_  
Signature of witness

\_\_\_\_\_  
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.

\_\_\_\_\_  
Name of Person who obtained Consent

\_\_\_\_\_  
Signature of Person who obtained Consent

\_\_\_\_\_  
Date