 UNIVERSITY OF Baguio	Reference No.: QF-QAO-016	Revision No: 00	Effectivity Date: March 1, 2020
	POLICIES, PROCEDURES AND GUIDELINES		


Title	
REC PROTOCOL NO	

UBRDC-RCR-022 Form 2. RESEARCH PROPOSAL ETHICS REVIEW FORM FOR RESEARCHES WITH NO HUMAN PARTICIPANTS

To the ethics reviewer: Evaluate the presence of the important ethical considerations in the application by checking the appropriate column.

COMPONENTS	YES	NO	REMARKS
1. A justification of the need to conduct the study given the current state of knowledge is discussed.			
2. An assurance that humans will not be adversely affected by the study is mentioned.			
3. How the researcher will obtain the materials to be used in the experiment/study is discussed.			
4. The benefits that humans will derive from the research during and after the study are explicitly stated.			
5. A discussion on the precautions to minimize the negative impact of the study on humans and the environment is provided.			
6. An information dissemination mechanism is sufficiently described by the proponent/s.			
7. The proposed methods for testing the stated hypothesis (if applicable) are appropriate.			
8. In general, there is a detailed satisfactory description of the procedures, invasive or not.			
9. In studies involving animals and plants, there is a provision made for extended care of the species.			
10. The research design is appropriate for the study.			
11. The study is NOT a duplication of a previous study.			
12. The selection of subjects/specimens/materials ensures objective and non-discriminatory procedures.			
13. There is a provision made for the people in the community where the specimens/subjects/materials were obtained to be informed of the results of the study.			
14. There is evidence that the species of the animals/plants/materials used in the study will not endanger health of the investigator.			
15. The participants will NOT be spending or using their resources for the research.			
16. There is evidence of the contribution of the study to improvement of life, systems and processes and most especially to the groups involved.			



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In general, the study is

Low risk

Medium risk

High risk

COMMENTS/SUGGESTIONS:

Action Taken

The study is approved to commence

The study needs minor revisions prior to commencement

The study needs major revisions prior to commencement

Evaluated by:

 (Signature over printed name)
 Member, ERC

 Date

