UWIScholar

Research Ethics Application

*** Title of the Research Project**

*** Research Start Date**

dd/mm/yyyy

*** Research End Date**

dd/mm/yyyy

* 1. Abstract

Summarise the Aims, Methodology, Location, Data Storage/Access, Time Frame, and Confidentiality Statement or Declaration for the study

300 words left

* 2. Background and Rationale for the study

Please include the literature review

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* 3. Aims and Objectives

300 words left

4. Research Objectives/ Research Questions/ Hypotheses (If Applicable)

300 words left

5. Methodological Design

*** 5.1 Overall Study Design (Theoretical Framework)**

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*** 5.2 Location and Time Frame of the study**

5.3 Justification for Participants or Subjects

*** 5.3.1.1 Inclusion Criteria**

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*** 5.3.1.2 Exclusion Criteria**

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*** 5.3.2 Special/Vulnerable Populations**

*** 5.3.3 Research Related Justification for Sample Size**

*** 5.3.4 Recruitment of Subjects**

*** 5.4 Research Procedures/Protocols - Data Collection/Research Intervention Procedures**

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5.5 Data Collection Instruments

Please upload the data collection instruments that will be used. These can include:

- Measurements
- Questionnaires
- Focus group questions/topics
- Interview questions/topics

Enter a description for the instrument and upload the file

Description	File (doc; docx; pdf)
	Choose File No file chosen

Add Row

*** 5.6 Methods of Data Analysis**

6. Confidentiality

*** 6.1 Methods for storing and securing study/biological data**

*** 6.2 Methods for protecting participants' confidentiality**

7. Risk/Benefit

*** 7.1 Indicate what is the level of risk associated with this research**

No more than minimal risk

*** 7.2 Please describe risk, discomfort (physical/psychological), inconvenience, side effects, and financial costs to participants (include measures to mitigate these risks/discomforts)**

*** 7.3 Indicate direct benefits to participants**

7.4 Impacts of the study on human groups that are not participants in the study (positive/negative, where applicable)

7.5 Impacts of the study on the environment (positive/negative, where applicable)

* 8. Compensation, rewards or other incentives for participants

9. Informed Consent

*** 9.1 Describe process for informed consent. Indicate if waiver of written consent is requested with justification or waiver of consent all together with justification.**

10. Funding

10.1 State sources of Funding. Indicate any potential for conflict of interests between researcher and funder.

10.2.1 What is the budget for the study (enter details below)?

10.2.2 Or upload a file containing the budget details

Choose File No file chosen

11. Expected outcomes and impact of the study

*** 11.1 How the results will be disseminated?**

11.2 How the results will be acted upon for both the participants and the community?

11.3 Limitations

* 12. References

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13. Document Upload

13.1 Research Proposal (required for clinical trials)

Choose File No file chosen

13.2 SPIRIT/WHO guidelines checklist for clinical trials

Choose File No file chosen

13.3 Recruitment Materials

Choose File No file chosen

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13.4 Letters to institutions for permission to access research sites and approvals as required

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13.5 Letters and Consent Forms

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13.6 Other Documents

Description	File
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