



NUREC Research Protocol Review Checklist

Components of the Research Protocol	Required Information	Yes <input type="checkbox"/> No <input type="checkbox"/> Partially done <input type="checkbox"/>	Reviewer Comments
1. Title of the Study	Clear, concise, and reflective of the study objectives.		
2. Principal Investigator and Research Team	Names, qualifications, affiliations, roles, and responsibilities.		
3. Abstract / Summary	Brief overview of the background, objectives, methods, and significance of the study.		
4. Background and Rationale	Scientific justification, existing literature, justification of the conceptual/theoretical framework chosen, problem statement, and the need for the study.		
5. Objectives / Research Questions	Clearly stated primary and secondary objectives or hypotheses and research questions.		
6. Study Design and Methodology	Justification for the study design and the type of methodology (qualitative/quantitative/mixed), sampling, tools, data collection and analysis.		
7. Study Population and Sample Size	Description of participants, inclusion/exclusion criteria, and justification for sample size (calculation of the sample size for quantitative research and justification for sample size for qualitative research is needed).		
8. Recruitment and Consent Procedures	How participants will be identified, approached, and informed consent obtained.		
9. Informed Consent Documents	Participant information sheets and informed consent forms in appropriate languages.		
10. Confidentiality and Data Protection	Measures to protect privacy, data storage, coding, and access control.		

11. Risk-Benefit Assessment	Potential harms, discomforts, and benefits to participants and society.		
12. Compensation and Incentives	Any payments, reimbursements, or other participant benefits. Note that, no participants shall be paid during any research undertaking in Zambia because it jeopardizes the research outcomes and there is no framework approved by NHRA. Exception is given only to data collectors hired, consultants and local guides. However, once NHRA develops the framework for compensation, considerations will be made and appropriate revisions to this SoP shall be made.		
13. Ethical Considerations	Protection of vulnerable populations, risk mitigation, and adherence to ethical guidelines.		
14. Community Engagement	Plans to involve local communities or stakeholders in culturally sensitive ways.		
15. Monitoring and Safety	Procedures for monitoring the study, handling adverse events, and reporting.		
16. Dissemination of Results	Plans for sharing findings with stakeholders, participants, and the scientific community.		
17. Conflict of Interest Disclosure	Declaration of any financial or other conflicts by the researcher or team.		
18. Funding Source and Budget Summary	Identification of funding bodies and how the funds will be used.		
19. Institutional Approvals and Permissions	1. Proof of University Introduction Letter Proof 2. Proof of approvals from relevant organizations, ministries, institutions, or local authorities.		
20. Timeline of the Study	Realistic timeline showing key stages from start to completion.		
21. Annexes / Supporting Documents	Tools, questionnaires, letters of support, CVs, translations, etc.		