WORLD HEALTH ORGANIZATION
REGIONAL OFFICE FOR AFRICA (AFRO)

For administrative use

CONFIDENTIAL
(exception: see 1.1-1.5 below)

JOINT AFRO, EDCTP & TDR SMALL GRANTS PROGRAMME

PROPOSAL

PART I. ADMINISTRATIVE INFORMATION (*Please print or type)*

Selected information from this box (1.1-1.5) may be released to the general public if this proposal is selected for funding		
1. Name of Principal Investigator and Institution affiliation	Title:	Sex: (M/F)
Last Name:	First name:	
Highest Degree obtained and year:		
Full postal address of Principal Investigator to be used for professional correspondence:		
Telephone:	Email (1):	
Mobile:	Email (2) or Website:	
2. Title of Project: *(120 characters maximum)*		
2.1a Country of research site:	**2.1b Leveraged amount: Yes / No** **If yes, amount in USD:**	
2.2a Proposed starting date:	**2.2b Estimated duration:**	
2.3 List of investigator(s), gender and institution(s) / department(s) collaborating on this project with the principal investigator(s) *(if any)*		
2.4 Name(s) of investigator(s) employed by this grant		

2.5 Summary of project: *(Do not exceed 300 words)*

3.1 Background and rationale (maximum 1 page):

3.2 Study Objectives (General and specific objectives - these must be specific, measurable, achievable within the budget and time context, resourced and time bound):

3.3 Materials and methods: (**maximum 3 pages)** to include:

i. Study area/setting
ii. Study subjects, design
iii. Sample size
iv. Sampling technique
v. Data collection instruments/methods
vi. Data management
vii. Analysis plans
viii. Potential risks to the project and mitigation actions

4. Expected outputs and dissemination of results

4.1 What are the expected outputs (publications) from this project?

4.2 How will data that are generated in this study be made openly available to use and analyse by third parties free of charge?

4.3 What other arrangements will there be to disseminate the findings?

4.4 How will this study impact on practice and policy?

4.5 Do you agree to share your results on the WHO website? Yes ☐ No ☐

4.6 Do you agree that all publications should be open access? Yes ☐ No ☐

4.7 Do you agree to send related peer review publication to WHO? Yes ☐ No ☐

4.8 Do you agree to acknowledge the financial support from WHO and EDCTP in all of your reports and publications? Yes ☐ No ☐

5. Ethical considerations:

The principal investigators are required to get clearance from a national, institutional or any other official Ethical Review Committee (*Required if the proposal involves research on human subjects, including collection of information or biological material)*

5.1 Is ethical clearance required? Yes ☐ No ☐

If yes:

is institutional ethical clearance document attached? Yes ☐ No ☐

is there a national ethical review body in your country? Yes ☐ No ☐

is national ethical clearance document attached? Yes ☐ No ☐

If you do not think that ethical clearance is needed for your study, please explain why not?

5.2 Summarise the potential risks and benefits of your study to individuals, communities or the country concerned.

5.3 How will informed consent be obtained?

5.4 How will confidentiality of the data gathered be ensured?

6. National government approval

Is national government approval required? Yes ☐ No ☐

If "yes", is the approval document attached? Yes ☐ No ☐

7. Budget

Have you attached your budget which should clearly link your expected results to the amount requested?

Yes ☐ No ☐