



Malawi-Liverpool-Wellcome Trust
Clinical Research Programme
P.O Box 30096, Chichiri, Blantyre 3,
Malawi.
Tel. +265 1 876444 Fax +265 1 875774

Title: Creating demand for Fishermen's schistosomiasis and HIV services (FISH): piloting and delivery of a 3-arm cluster randomized control trial (cRCT) in Malawi

Principal Investigator: Dr. Augustine Choko

Participant ID

[The following text must be read to the participant, who must have their own copy to take home]

Introduction

Hello. My name is and I am working with Malawi Liverpool Wellcome Trust (MLW) on behalf of Dr. Augustine Choko and colleagues. We thank you for accepting to answer a few questions about yourself. In this component of the study, we are interested in understanding a few things about fishermen who are part of a cluster randomized trial about increasing demand for HIV and schistosomiasis services.

Request for your Voluntary Participation

I would like to ask you to voluntarily participate in this interview. You have been identified because you are a fisherman here in Mangochi. Please note that you were not selected to participate in this interview because you are HIV positive or negative, or that because you have schistosomiasis. We consider that your participation in this project would help us understand the best ways to serve fishing communities with HIV and schistosomiasis services. This is very important because the Ministry of Health may consider implementing successful strategies studied here nationally.

Procedure

Please note that we are carrying out these procedures on everyone who accepts to participate in this interview. All participants are asked a few questions about themselves and/or their social and sexual networks.

Your participation is entirely voluntary. If you decide to take part, you may withdraw from the interview any time. You also have a right not to answer any particular question or questions that will be asked. Declining to participate in this interview will not affect any health services that you or any person related to you may be currently receiving or may require in future.

Confidentiality

All personal information collected in this study will be kept strictly confidential. I will not share the information you provide with anyone who is not part of this research. But it may be shared with fellow researchers and may also be published through meetings or journals in a manner that does not reveal your identity. Before sharing in this manner,

C02a: Participant Information and Consent Form for Baseline Interview - Eng v1.0; 18 Feb 2020

the information from you will be combined with that from other research participants. Information which could identify you or anyone related to you will never be released. This also means that names of study participants, including your own will not be included when sharing the data. Data collection equipment and the data collected will be kept with identifiers, locked, and only accessible to people that have authorised access.

Risks

You may be uncomfortable with some of the questions that I will ask. You are perfectly entitled to refuse to discuss issues that you do not want to.

Benefits

There are no direct benefits to you in your taking part in this interview. However, what we learn from this study would help the Ministry of Health to make important decisions regarding how best to serve fishing communities with HIV and schistosomiasis services. It is not yet known if services including the offer of HIV self-test kits would increase demand for HIV and schistosomiasis services. Therefore, your participation would benefit many others in the future.

Compensation

You will not receive payment for participating in the study.

Contact details

This research has been approved by the College of Medicine Research Ethics Committee (COMREC) and the Liverpool School of Tropical Medicine Research Ethics Committee. If you have any questions regarding your rights as a research participant, or concerns on how you have been treated in the study, please feel free to contact **Dr. Augustine Choko** [+265 (0) 999 577 452] or [augutc@gmail.com]. If you have any questions regarding your rights as a research participant, or concerns on how you have been treated in the study, please feel free to contact COMREC Secretariat, College of Medicine, Private Bag 360, Chichiri, Blantyre 3 or call on 01871911 ext 334.

Consent Declaration

If you agree to voluntarily participate in the study, please sign or write your initial or your thumb print below to show that you understand the information above and that your consent is given voluntarily.

		Participant Initials / thumb print
1	I have received and read or had read to me the information sheet provided by the Researcher that explains in detail the reasons for the study.	-----
2	I have understood the purpose of the research.	-----
3	I have asked all the questions that I have about the purpose of the research and feel that I have enough information about it.	-----
4	I understand the reasons for this study.	-----
5	I am willing to take part in the study.	-----
6	I understand what I will be required to do if I participate in the study.	-----
7	I know that I have the right to leave the study at any time or to refuse to answer any questions.	-----
8	If I do not agree to take part in this study, I understand that I will not be penalized for doing so by the researcher nor by any medical service providers in the future.	-----
9	I voluntarily agree to take part in this study	-----

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 Participant name Date Signature or thumb print

If the participant gave verbal consent, please enter the name of person who witnessed the consent here, and their signature:

----- / ----- / -----
 Name of Witness (BLOCK CAPITALS) Date Signature or thumb print

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 Name of person obtaining consent Date Signature