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### **INFORMED CONSENT FORM: RANDOMISED CONTROLLED TRIAL**

*This document outlines the research study and expectations for potential participants. It should be written in layman terms and typed on MUST-IRC letterhead. The wording should be directed to the potential participant NOT to IRC. If a technical term must be used, define it the first time it is used. Also, any abbreviation should be spelled out the first time it is used.*

**NB: All the sections of this document must be completed without any editing or deletions**

*Please use a typing font that is easily distinguishable from the questions of the form*

### **Study Title: It should be the same as on all other documents related to the study AFFECTING THE EPIDEMIOLOGY OF HIV IN UGANDA THROUGH OLDER ADOLESCENTS**

#### **Principal Investigator(s):**

**Emmanuel Kyagaba, Principal Investigator, Mbarara University of Science and Technology and Internet Solutions for Kids, Uganda**  
**Michele Ybarra, Ph.D., Principal Investigator and President, Center for Innovative Public Health Research (CiPHR)**

### **INTRODUCTION**

HIV continues to be a significant public health challenge in Uganda.

#### **What you should know about this study:**

- You are being asked to join a research study.
- This consent form explains the research study and your part in the study
- Please listen to it carefully and take as much time as you need to decide
- You are a volunteer. You can choose not to take part. And if you join, you may quit at any time. There will be no penalty if you decide to quit the study.

**Purpose of the research project:** *Include a statement that the study involves research, estimated number of participants, an explanation of the purpose(s) of the research procedure and the expected duration of the subject's participation.*

#### **Background/ Purpose**

We are developing a sexual health and HIV prevention program for young adults in Uganda. The program will be sent to young people through text messages. This research study is

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MUST-IRC Stamp:	APPROVAL DATE: APPROVED CONSENT IRB VERSION NUMBER: PI NAME: IRB NO: