

# Using Mobile Phones to Measure Diarrhoea

<b>Submission date</b> 06/03/2019	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 20/03/2019	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 02/07/2020	<b>Condition category</b> Infections and Infestations	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

### Background and study aims

With diarrhoea being one of the two most significant causes of death in children under five, collection of accurate data on diarrhoea prevalence is of paramount importance. However, collecting this data has been proven difficult. Standard questions used to assess diarrhoea, asking if a child has had diarrhoea two or more times in the past two weeks, has been seen to suffer from recall bias, social desirability, high cost of data collection, and Hawthorne effects. To address this, we will study the impact of shortening recall period to 24 hours, including a multi-level questionnaire, and altering incentive level. We will do this while collecting data using mobile phone text messaging.

### Who can participate?

Households selected will have at least one child under the age of five, live in the study area, and have access to mobile phones.

### What does the study involve?

Participants will be randomised into one of eight cohorts. Each cohort, at random order, will be subjected to every possible combination of incentive (no incentive or TZS1000 incentive), recall period (two week or 24 hour), and questionnaire type (standard question or three level). Over eight two-week rounds of data collection, participants will receive surveys on diarrhoea as per their randomisation, and will be remunerated and incentivised as necessary.

### What are the possible benefits and risks of participating?

Possible harms of the study surround discomfort with questions being asked, however, participants are free to withdraw at any time. Further, there is small risk of a breach of confidentiality. To address this, data is kept separately from identifiers, and all data are encrypted. There are no direct benefits, however, if a participant reports a medically concerning symptom, they will be referred to a local clinic.

### Where is the study run from?

UN-Habitat Regional Office for Africa and the Arab States, Nairobi (Kenya)

### When is the study starting and how long is it expected to run for?

Data collection will begin in April 2019 and end in August 2019.

Who is funding the study?  
University of Warwick, UK

Who is the main contact?  
Ryan Rego (ryan.rego@warwick.ac.uk) and Samuel Watson (s.watson.1@warwick.ac.uk)

## Contact information

**Type(s)**  
Scientific

**Contact name**  
Mr Ryan Rego

**ORCID ID**  
<https://orcid.org/0000-0003-1361-9366>

**Contact details**  
Warwick Medical School  
University of Warwick  
Warwick  
United Kingdom  
CV4 7AL  
+44 07462890990  
ryan.rego@warwick.ac.uk

## Additional identifiers

**Clinical Trials Information System (CTIS)**  
Nil known

**ClinicalTrials.gov (NCT)**  
Nil known

**Protocol serial number**  
REGO2018-2148

## Study information

**Scientific Title**  
Pilot testing of a parent record of child health

**Acronym**  
SMSdiarrhoea

**Study objectives**  
This study aims to evaluate several hypotheses aimed at improving traditional diarrhoea measurement methods, asking if a child has had two or more loose or watery stool in the past two weeks. These are:

- Reducing the recall threshold from two weeks to 24 hours will decrease prevalence
- The use of a three point scale asking further questions on number of schools, treatment, and blood will result in lower prevalence than the standard measure
- That response rates vary over time
- That incentive alters response rate and data quality

### **Ethics approval required**

Old ethics approval format

### **Ethics approval(s)**

Approved 12/01/2018, Warwick Biomedical & Scientific Research Ethics Committee (University House, University of Warwick, Coventry, CV4 7AL; E.C.Dight@warwick.ac.uk; 024 765 73961), ref: REGO-2018-2148

### **Study design**

Individual level crossover factorial randomised controlled trial

### **Primary study design**

Observational

### **Study type(s)**

Screening

### **Health condition(s) or problem(s) studied**

Diarrhoea/Enteric Infection

### **Interventions**

Differing recall periods; questionnaire types; and incentive levels for diarrhoea surveys.

Initially, households will be randomised into one of eight cohorts using block randomisation. At every two week periods, cohorts will switch into new surveying strategies. These strategies are unique combinations of recall period (two week or 24 hour); questionnaire type (three level or one level); and incentive level (no incentive or some incentive). All surveys will take place by mobile phone text messaging, and all cohorts will experience all combinations of treatments at any time - through a random sequence. This study will take place in slums around Mwanza, Tanzania.

### **Intervention Type**

Other

### **Primary outcome(s)**

Diarrhoea Prevalence through the use of a mobile phone survey. This will take place at pre-set time points either daily or every two weeks for a period of 16 weeks.

### **Key secondary outcome(s)**

Trends in diarrhoea prevalence over time

### **Completion date**

01/11/2019

# Eligibility

## Key inclusion criteria

1. Resides in the enumeration area
2. Has a child under five
3. Has access to mobile phone
4. Consents to study

## Participant type(s)

Carer

## Healthy volunteers allowed

No

## Age group

Mixed

## Sex

All

## Total final enrolment

141

## Key exclusion criteria

1. Child under five is exclusively breast fed
2. Does not have their own mobile phone (e.g. shares phone or no access to phone)

## Date of first enrolment

15/03/2019

## Date of final enrolment

01/04/2019

# Locations

## Countries of recruitment

Kenya

Tanzania

## Study participating centre

### UN-Habitat

UN-HABITAT Regional Office for Africa and the Arab States

P.O. Box 30030, GPO

Nairobi

Kenya

00100

# Sponsor information

## Organisation

University of Warwick

## ROR

<https://ror.org/01a77tt86>

# Funder(s)

## Funder type

University/education

## Funder Name

University of Warwick

## Alternative Name(s)

The University of Warwick, Warwick

## Funding Body Type

Private sector organisation

## Funding Body Subtype

Universities (academic only)

## Location

United Kingdom

# Results and Publications

## Individual participant data (IPD) sharing plan

All data generated or analysed during this study will be included in the subsequent results publication.

## IPD sharing plan summary

Other

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	results	30/06/2020	30/06/2020	Yes	No