Can Repellents Prevent Malaria in Africa?

Submission date 29/01/2010	Recruitment status No longer recruiting
Registration date 18/02/2010	Overall study status Completed
Last Edited 21/08/2014	Condition category Infections and Infestations

[] Prospectively registered

[] Protocol

- [] Statistical analysis plan
- [X] Results
- [] Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s) Scientific

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers NIMR/HQ/R8a/VolIX/780

Study information

Scientific Title

Low cost repellents for use in rural Africa: a short-term efficacy, effectiveness and perceived benefit survey in Kilombero, Tanzania

Acronym

CRPMA

Study objectives

As Tanzania progresses towards the goals of the Abuja declaration and insecticide-treated bed nets (ITN) coverage becomes almost universal, there is likely to be a selection pressure on malaria mosquitoes to feed outdoors and earlier in the evening when hosts are available. This coupled with changes in lifestyle such as increased access to electricity so people stay awake later means that the relative exposure of the population to infectious mosquito bites is likely to switch to earlier in the evening. A topical insect repellent containing deet can dramatically reduce malaria in South America and Southern Asia where vectors feed early in the evening. The project aims to measure the impact of such a repellent on clinical episodes of malaria in rural Africa.

Ethics approval required

Old ethics approval format

Ethics approval(s)

1. Ifakara Health Institute Institutional Review Board, 10/11/2008, ref: IHRDC/IRB/No. A46

2. National Institute of Medical Research, Tanzania, 06/03/2009, ref: NIMR/HQ/R8a/VolIX/780

Study design

Cluster controlled randomised trial

Primary study design Interventional

Secondary study design Randomised controlled trial

Study setting(s) Other

Study type(s) Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied Malaria

Interventions

Long lasting insecticide treated nets (Olyset) + 15% deet repellent Long lasting insecticide treated nets (Olyset) + placebo lotion Total duration of intervention: 44 weeks Total duration of follow-up: 1 month after the trial ends

Intervention Type

Drug

Phase Not Applicable

Drug/device/biological/vaccine name(s)

Deet-containing insect repellent

Primary outcome measure

1. Malaria incidence

2. Clinical episodes of malaria

Data is continually collected on a daily basis through passive case detection at a local clinic throughout the trial.

Secondary outcome measures

Malaria prevalence.

Data is continually collected on a daily basis through passive case detection at a local clinic throughout the trial.

Overall study start date

30/08/2009

Completion date 30/07/2010

Eligibility

Key inclusion criteria Household head over 18 years, either sex

Participant type(s) Patient

Age group Adult

Lower age limit 18 Years

Sex Both

Target number of participants

4819

Key exclusion criteria Under six months of age

Date of first enrolment 30/08/2009

Date of final enrolment 30/07/2010

Locations

Countries of recruitment England

Tanzania

United Kingdom

Study participating centre Disease Control and Vector Biology Unit (DCVBU) London United Kingdom WC1E 7HT

Sponsor information

Organisation Ifakara Health Institute (Tanzania)

Sponsor details Box 53 Ifakara Tanzania 53

Sponsor type Research organisation

Website http://www.ihi.or.tz/

ROR https://ror.org/04js17g72

Funder(s)

Funder type Research organisation

Funder Name Population Services International (PSI) (Tanzania) - Innovations Grant

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<u>Results article</u>	results	16/08/2014		Yes	No