

Effectiveness perifocal deltamethrin spray versus insecticide treated curtains for Aedes control

Submission date 23/07/2012	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 07/09/2012	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 18/01/2019	Condition category Infections and Infestations	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Dengue is a common viral infection spread by mosquitoes. It is widespread in tropical and sub-tropical regions, mainly affecting urban areas but extending into more populated rural areas. The mosquitoes breed in water storage containers and other deposits containing relatively clean water. Attempts at eradicating the mosquitoes failed during the 1960s in Latin America and today more than 50% of houses are infested with mosquitoes in many endemic areas. This research is studying new methods to control dengue mosquitoes. We are studying the acceptance, effectiveness and cost-effectiveness of insecticide-treated curtains and of spraying the insecticide deltamethrin in and around the house.

Who can participate?

Houseblocks are selected at random in the city of Santiago de Cuba among those with the highest infestation rates in 2009-2010, and all families living in these areas are invited to participate

What does the study involve?

Groups of houses are randomly allocated to one of three groups. Group 1 receive the routine mosquito control activities of the Ministry of Health programme. Group 2 receive insecticide-treated curtains (maximum of three per house) on top of the routine mosquito control activities of the Ministry of Health programme. Group 3 receive every four months a spraying of deltamethrin in and around their houses on top of the routine mosquito control activities of the Ministry of Health programme.

What are the possible benefits and risks of participating?

Participants receive the mosquito control methods for free. As they have shown already good results in other studies, the mosquitoes in the house will decrease if the tool is correctly used. The curtains that will be distributed didn't show any side effects in other studies, except for a brief feeling of itching and sneezing in 7% of the people, lasting for at most 1 day. The spraying

is a procedure that has already been used in Santiago de Cuba by the Ministry of Health. In this study it's another insecticide, widely used in Africa and Latin-America, but up to now not yet used in this area in this form.

Where is the study run from?

Institute of Tropical Medicine (Belgium)

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

May 2011 to December 2013

Who is funding the study?

1. Directorate-General for Development Cooperation (Belgium)

2. Institute of Tropical Medicine (Belgium)

3. Ministry of Health (Cuba)

Who is the main contact?

Prof. Patrick Van der Stuyft

Contact information

Type(s)

Scientific

Contact name

Prof Patrick Van der Stuyft

Contact details

Institute of Tropical Medicine

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

B300201111923

Study information

Scientific Title

Effectiveness and cost-effectiveness of perifocal residual deltamethrin spraying and of long lasting deltamethrin treated curtains distribution for Aedes control

Study objectives

1. Evaluate the uptake and acceptability of tools in both intervention arms (ITC and PFS)
2. Evaluate the effectiveness of both tools, on top of the routine, in the control of Aedes aegypti (in comparison to the effect of the routine programme alone)
3. Compare the cost-effectiveness of both tools with the routine Aedes control strategy

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethical Committee, University Hospital Antwerp, 14/11/2011 ref: 11/34/227

Study design

Interventional cluster-randomized controlled trial single-center study

Primary study design

Interventional

Secondary study design

Cluster randomised trial

Study setting(s)

Home

Study type(s)

Prevention

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

Effect of intervention on Aedes aegypti infestation level (vector of dengue fever)

Interventions

Control group: routine aedes control programme (entomological surveillance, source reduction, larviciding and selective adulticiding, health education)

Insecticide treated curtains: made from long-lasting, insecticide-treated (pyrethroid deltamethrin is applied during manufacture) polyester netting that requires no re-impregnation (PermaNet®; Vestergaard-Frandsen company). PermaNet materials are special UV protected and retain their insecticidal properties and efficacy for about 2 years (information from producer). The material has been approved by WHOPES for use as bednets.

Deltamethrin perifocal spraying (PFS): K-Othrine 25 WG, supplied by Bayer Environmental Sciences co. (25% deltamethrin formulation) will be sprayed every 4 months (3 times/year). It is a granular formulation that need to be solved into water (20 gram in 8 Liter of water, sufficient to

treat 200 m² and attaining 25 mg a.i./m²). It has a long lasting residual activity. Where deposits remain undisturbed, residual activity depends upon the nature of the surface. Sustained residual activity beyond 12 weeks post-application is observed on non-porous surfaces. Deltamethrin is photostable and the particulate suspension enhances availability to insects.

The insecticide will be sprayed on the outside of the ground-level water tanks and the walls behind them; and on the adult Aedes resting sites (for example under beds, in and under closets) in the intradomestic area.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

1. Effectiveness: Breteau Index (number of containers positive for immature aedes stages per 100 inspected houses)
2. House index (number of houses positive for at least one container with aedes immature stages per 100 houses inspected)
3. Pupae per person index (number of pupae per inhabitant), (adult mosquito infestation), confirmed Dengue cases (if any)

Secondary outcome measures

1. Uptake and use of ITC
2. Acceptance of PFS and ITC (and identification of underlying lay dimensions of acceptability)
3. Change in intra-and extradomiciliary risks for Aedes infestation
4. Cost- effectiveness of ITC and of PFS
5. Residual insecticidal activity of deltamethrin applied in ITC and in PFS

Overall study start date

01/05/2011

Completion date

31/12/2013

Eligibility**Key inclusion criteria**

All households of houseblocks, chosen at random in urban Santiago de Cuba

Participant type(s)

All

Age group

All

Sex

Both

Target number of participants

63 clusters of 250 households each

Key exclusion criteria

1. Houseblocks without community approval
2. Houses without household approval

Date of first enrolment

01/05/2011

Date of final enrolment

31/12/2013

Locations**Countries of recruitment**

Belgium

Cuba

Study participating centre

Institute of Tropical Medicine

Antwerp

Belgium

2000

Sponsor information**Organisation**

Institute of Tropical Medicine (Belgium)

Sponsor details

General Epidemiology and Disease Control

Public Health Department

Nationalestraat 155

Antwerp

Belgium

2000

Sponsor type

University/education

Website

<http://www.itg.be/itg/>

ROR

<https://ror.org/03xq4x896>

Funder(s)

Funder type

Government

Funder Name

Directorate-General for Development Cooperation (Belgium) ref: 95900

Funder Name

Institute of Tropical Medicine (Belgium)

Funder Name

Ministerio de Salud Pública [MINSAP] (Cuba)

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?
Results article	results	01/05/2016	18/01/2019	Yes	No
Results article	results	08/11/2017	18/01/2019	Yes	No
Results article	results	02/01/2018	18/01/2019	Yes	No