Home and Community Management of Malaria and Pneumonia

Submission date	Recruitment status No longer recruiting	Prospectively registered		
14/06/2010		☐ Protocol		
Registration date 04/03/2011	Overall study status Completed	Statistical analysis plan		
		[X] Results		
Last Edited 29/12/2020	Condition category Infections and Infestations	Individual participant data		

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number HS 72

Study information

Scientific Title

An integrated Management of Malaria and Pneumonia in children under five at home and community level

Acronym

HCMMP

Study objectives

Prompt use and compliance with efficacious treatment for both malaria and pneumonia provided at home and community level reduces under five morbidity and mortality significantly and is more cost effective than giving antimalarials alone.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Uganda National Council for Science and Technology approved on the 17th August 2005 (ref: HS 72)

Study design

Cluster randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Malaria and pneumonia

Interventions

The intervention arm has combined home and community malaria and pneumonia treatment where pre-packaged drugs (anti-malarial and antibiotics) are distributed through trained community medicine distributors. Children who present with fever are given antimalarials (Coartem®) and when they in addition have rapid breathing, they are given antibiotics (amoxycillin).

For the control arm, antimalarials (Coartem®) are distributed by trained community medicine distributors to the children who present with fever. Those who present with rapid breathing are referred to health facilities. The difference between the intervention and control arms is that in the intervention, there is community distribution of antibiotics in addition to the antimalarials while in the control, there is only distribution of antimalarials alone.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Mortality in children under five

Key secondary outcome(s))

Determined at the end of the two year period:

1. Mortality preceded by an acute febrile illness

- 2. Severe anaemia
- 3. Cost-effectiveness of the interventions
- 4. Equity analysis

Completion date

01/12/2011

Eligibility

Key inclusion criteria

Children aged 4 to 59 months

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Lower age limit

4 months

Upper age limit

59 months

Sex

All

Key exclusion criteria

Children below 4 months of age or those above 59 months old

Date of first enrolment

02/12/2009

Date of final enrolment

01/12/2011

Locations

Countries of recruitment

Uganda

Study participating centre

Makerere University School of Public Health

Kampala Uganda 4147072

Sponsor information

Organisation

Uganda National Council for Science and Technology (UNCST) (Uganda)

ROR

https://ror.org/05jv1pq83

Funder(s)

Funder type

Research organisation

Funder Name

United Nations Children's Fund (UNICEF)/United Nations Development Programme (UNDP) /World Bank/World Health Organization (WHO) - Special Programme for Research and Training in Tropical Diseases (TDR) (ref: A20141)

Results and Publications

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	22/09/2013	29/12/2020	Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes