Assessing the impact of an intervention to implement the referral care component of Integrated Management of Childhood Illnesses (IMCI) in district hospitals

Submission date	Recruitment status No longer recruiting	Prospectively registered		
20/11/2008		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
21/11/2008	Completed	[X] Results		
Last Edited	Condition category	[] Individual participant data		
15/01/2015	Other			

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Mike English

Contact details

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Additional identifiers

Protocol serial number SSC911

Study information

Scientific Title

A randomised, parallel group trial of a multifaceted intervention to improve care of children in Kenyan district hospitals

Study objectives

That the quality of paediatric inpatient care at district hospitals in Kenya can be improved by an intervention over 18 months comprising evidence based guidelines, training, supervision, feedback and facilitation of local problem solving.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved by the KEMRI National Ethical Review Committee on 21st September 2005

Study design

Randomised, parallel, controlled intervention study

Primary study design

Interventional

Study type(s)

Quality of life

Health condition(s) or problem(s) studied

Quality of paediatric inpatient care

Interventions

Intervention arms:

Baseline survey followed by training, provision of guidelines and job aides (guideline booklets, standard admission record forms), external supervision and feedback after surveys and local facilitation provided over 18 months. Cross-sectional surveys conducted at 6, 12 and 18 months post-baseline.

Control arm:

Baseline survey followed by limited training, provision of guidelines and job aides (guideline booklets, standard admission record forms), written feedback after surveys over 18 months. Cross-sectional surveys conducted at 6, 12 and 18 months post-baseline.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Proportion of children with admission assessment, classification and treatment that is adherent to guidelines provided for pneumonia, diarrhoea/dehydration and malaria. Measurements will be available at 6, 12 and 18 months post-baseline with the primary endpoint being 18 months.

Key secondary outcome(s))

Proportion of key clinical signs documented for pneumonia, diarrhoea/dehydration and malaria admissions at 6, 12 and 18 months post-baseline with the primary endpoint being 18 months and will use a facility inventory to describe the proportion of items on a pre-specified checklist that are available at the time of the survey also at 6, 12 and 18 months post-baseline with the primary endpoint being 18 months.

Completion date

31/05/2009

Eligibility

Key inclusion criteria

- 1. Hospitals purposefully selected to represent diversity of Kenyan context and fulfilling minimum size requirements of 1,000 paediatric admissions per annum.
- 2. Random selection of all childhood admissions:
- 2.1. Aged 0 59 months, either sex
- 2.2. In participating hospitals with a diagnosis covered by inpatient IMCI guidelines (malaria, pneumonia, meningitis, malnutrition, diarrhoea/dehydration, neonatal sepsis, prematurity/low birthweight)

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Lower age limit

0 months

Upper age limit

59 months

Sex

All

Key exclusion criteria

Children aged 0 - 59 months in participating hospitals with non-IMCI diagnoses

Date of first enrolment

01/07/2006

Date of final enrolment

31/05/2009

Locations

Countries of recruitment

Study participating centre KEMRI/Wellcome Trust Programme Nairobi Kenya 00100

Sponsor information

Organisation

The Wellcome Trust (UK)

ROR

https://ror.org/029chgv08

Funder(s)

Funder type

Charity

Funder Name

The Wellcome Trust (UK) (grant ref: 076827) - Senior Research Fellowship awarded to Dr. Mike English

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	25/11/2011		Yes	No
Results article	results	18/07/2014		Yes	No

Other publications	discussion paper on study design	01/06/2008	Yes	No
Participant information sheet	Participant information sheet	11/11/2025 11/11 /2025	No	Yes