

# Chilenje Infant Growth, Nutrition and Infection Study

<b>Submission date</b> 28/06/2005	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 05/08/2005	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
<b>Last Edited</b> 13/01/2015	<b>Condition category</b> Nutritional, Metabolic, Endocrine	<input type="checkbox"/> 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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WC1E 7HT

## Additional identifiers

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

**Secondary identifying numbers**  
N/A

## Study information

**Scientific Title**

Chilenje Infant Growth, Nutrition and Infection Study

**Acronym**

CIGNIS

**Study objectives**

Feeding for 12 months with a fortified complementary food will decrease the proportion of stunted children by 30%.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Added 02/09/10:

1. Approved by the University of Zambia ethics committee
2. Approved by the London School of Hygiene and Tropical Medicine ethics committee

**Study design**

Randomised controlled trial

**Primary study design**

Interventional

**Secondary study design**

Randomised controlled trial

**Study setting(s)**

Other

**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**

Infant growth faltering

**Interventions**

One of two locally developed complementary foods fed for 12 months

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome measure**

Prevalence of stunting

**Secondary outcome measures**

1. Body composition
2. Indices of iron (Fe), zinc (Zn), copper (Cu), selenium (Se) and vitamin A status
3. Morbidity
4. Gastrointestinal permeability
5. Response to oral polio vaccine
6. Behavioural development
7. Development of chronic childhood viral infections

**Overall study start date**

20/07/2005

**Completion date**

30/06/2008

**Eligibility**

**Key inclusion criteria**

Infants aged 6 months and resident in Chilenje, Zambia whose mothers agree:

1. To prepare and feed their infants for 12 months the food supplied to them
2. To attend the specified clinic or home visits
3. To let their infants undergo specified urine collection and blood sampling
4. To permit the infants to be tested for human immunodeficiency virus (HIV)

**Participant type(s)**

Patient

**Age group**

Child

**Lower age limit**

6 Months

**Sex**

Both

**Target number of participants**

800

**Key exclusion criteria**

Non-fulfilment of inclusion criteria or evidence of chronic disease (active tuberculosis [TB], symptomatic HIV)

**Date of first enrolment**

20/07/2005

**Date of final enrolment**

30/06/2008

# Locations

## Countries of recruitment

England

United Kingdom

Zambia

## Study participating centre

**Nutrition and Public Health Interventions Research Unit**

London

United Kingdom

WC1E 7HT

# Sponsor information

## Organisation

Bill and Melinda Gates Foundation (USA)

## Sponsor details

P.O. Box 23350

Seattle WA

United States of America

98102

## Sponsor type

Charity

## ROR

<https://ror.org/0456r8d26>

# Funder(s)

## Funder type

Charity

## Funder Name

Bill and Melinda Gates Foundation (USA) - (ref: 37253)

## Alternative Name(s)

Bill & Melinda Gates Foundation, Gates Foundation, BMGF, B&MGF, GF

**Funding Body Type**

Government organisation

**Funding Body Subtype**

Trusts, charities, foundations (both public and private)

**Location**

United States of America

**Funder Name**

DSM (South Africa)

## 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?
<a href="#">Results article</a>	results	01/05/2011		Yes	No
<a href="#">Results article</a>	results	01/02/2012		Yes	No
<a href="#">Results article</a>	results	01/03/2012		Yes	No
<a href="#">Results article</a>	results	01/11/2014		Yes	No