Phase III, double-blind, randomised, placebocontrolled trial to evaluate efficacy against radiological pneumonia and invasive pneumococcal disease in Gambian infants

Submission date	Recruitment status No longer recruiting	Prospectively registered	
10/11/2004		☐ Protocol	
Registration date 05/04/2005	Overall study status Completed	Statistical analysis plan	
		[X] Results	
Last Edited 09/11/2007	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 RPC096

Study information

Scientific Title

Acronym

PVT

Study objectives

To assess the efficacy of a nine-valent pneumococcal conjugate vaccine in children.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Phase III, double-blind, randomised, placebo-controlled trial

Primary study design

Interventional

Study type(s)

Health condition(s) or problem(s) studied

Pneumococcus/vaccines

Interventions

Treatment group:

Nine-valent pneumococcal/polysaccharide protein conjugate vaccine which contains 2 µg of type 1, 4, 5, 9V, 14, 19F, 23F polysaccharides, 4 µg of type 6B polysaccharide and 2 µg of type 18 oligosaccharide linked to the diphtheria toxoid protein CRM197, reconstituted with DPT-Hib (Tetramune™) from the same manufacturer

Placebo group:

Lyophilised placebo cake reconstituted with Tetramune™

Intervention Type

Other

Phase

Phase III

Primary outcome(s)

First episode of radiological pneumonia

Key secondary outcome(s))

- 1. Clinical or severe clinical pneumonia
- 2. Invasive pneumococcal disease
- 3. All-cause admissions

Completion date

01/01/2004

Eligibility

Key inclusion criteria

- 1. Infants aged at least 6 weeks and less than one year
- 2. Resident in study area
- 3. Written informed consent obtained from mother

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Lower age limit

6 weeks

Upper age limit

1 years

Sex

Αll

Key exclusion criteria

- 1. Aged less than 6 weeks or more than 1 year
- 2. Not resident in study area
- 3. Planning to move out of study area within 4 months
- 4. Previous receipt of Diphtheria, Pertussis, Tetanus (DPT)-Haemophilus influenzae type b (Hib) vaccine
- 5. Uncertain prior vaccination record
- 6. Serious chronic illness
- 7. Inclusion in previous vaccine trial
- 8. Failure of family to give consent

Date of first enrolment

01/08/2000

Date of final enrolment

01/01/2004

Locations

Countries of recruitment

Gambia

Switzerland

Study participating centre 20, Avenue Appia Geneva-27 Switzerland CH 1211

Sponsor information

Organisation

National Institute of Allergy and Infectious Diseases - Division of Microbiology and Infectious Diseases (USA)

ROR

https://ror.org/043z4tv69

Funder(s)

Funder type

Research organisation

Funder Name

National Institutes of Health (NIH) (USA)

Alternative Name(s)

US National Institutes of Health, Institutos Nacionales de la Salud, NIH, USNIH

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United States of America

Funder Name

World Health Organization (WHO)/Department of Immunisation, Vaccines and Biologicals (IVB) (Switzerland)

Funder Name

Bill and Melinda Gates Foundation (USA)

Alternative Name(s)

Bill & Melinda Gates Foundation, Gates Foundation, Gates Learning Foundation, William H. 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

United States Agency for International Development (USAID) (USA)

Alternative Name(s)

U.S. Agency for International Development, Agency for International Development, USAID

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United States of America

Funder Name

Medical Research Council (UK)

Alternative Name(s)

Medical Research Council (United Kingdom), UK Medical Research Council, MRC

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results	01/03/2005		Yes	No