# A randomised trial of hepatitis B booster vaccination at the age of 13 to 15 years following infant vaccination to assess efficacy against hepatitis B infection and acute hepatitis

Submission date	Recruitment status  No longer recruiting	[X] Prospectively registered		
21/09/2000		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
21/09/2000	Completed	[X] Results		
Last Edited	Condition category	[] Individual participant data		
02/07/2009	Infections and Infestations			

## Plain English summary of protocol

Not provided at time of registration

## Contact information

## Type(s)

Scientific

#### Contact name

Dr Andrew J Hall

#### Contact details

London School of Hygiene and Tropical Medicine Keppel Street London United Kingdom WC1E 7HT

# Additional identifiers

Protocol serial number G0000531

# Study information

Scientific Title

#### **Acronym**

**GHABS** 

#### **Study objectives**

Added as of 23/04/2007:

A booster dose of Hepatitis B Virus (HBV) vaccine 15 years after infant vaccination will increase the magnitude and duration of immune responses (measured two and 52 weeks after the boost).

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Not provided at time of registration

#### Study design

Randomised controlled trial

## Primary study design

Interventional

#### Study type(s)

Treatment

#### Health condition(s) or problem(s) studied

Hepatitis B Virus

#### **Interventions**

Added as of 23/04/2007: Booster dose HBV vaccine

#### Intervention Type

Other

#### Phase

**Not Specified** 

#### Primary outcome(s)

Added as of 23/04/2007:

Anti-HBs titers after one year.

#### Key secondary outcome(s))

Added as of 23/04/2007:

Proportion without anamnestic response following boosting.

#### Completion date

31/12/2005

# **Eligibility**

#### Key inclusion criteria

Added as of 23/04/2007:

- 1. ID and vaccination data matched with Gambia Hepatitis Intervention Study (GHIS) database
- 2. Born between 1/7/1988 and 31/12/1989
- 3. Living in selected Health Centre (HC) catchment area's
- 4. Informed consent participant and parent/guardian

#### Participant type(s)

**Patient** 

## Healthy volunteers allowed

No

#### Age group

Child

#### Sex

All

### Key exclusion criteria

Added as of 23/04/2007:

- 1. No complete informed consent
- 2. No match found
- 3. Not fully vaccinated in infancy
- 4. Not in age range

#### Date of first enrolment

01/01/2004

#### Date of final enrolment

31/12/2005

## Locations

#### Countries of recruitment

United Kingdom

England

Gambia

Study participating centre
London School of Hygiene and Tropical Medicine
London
United Kingdom
WC1E 7HT

Sponsor information

#### Organisation

Medical Research Council (MRC) (UK)

# Funder(s)

## Funder type

Research council

#### **Funder Name**

Medical Research Council (MRC) (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

Not provided at time of registration

## **Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	15/08/2007		Yes	No