

# Step-down affordable treatment for chronic HEPatitis B infection in Africa

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		<input type="checkbox"/> Protocol
<b>Registration date</b> 31/03/2014	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
<b>Last Edited</b> 10/05/2021	<b>Condition category</b> Infections and Infestations	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

Hepatitis B virus (HBV) is a virus which causes long term liver damage in people all over the world. It is very common in much of sub-Saharan Africa and southeast Asia. Some countries have more than 10% of the entire population with the virus. Zambia is a country with a high HBV prevalence approximately 10%. HBV causes liver cirrhosis and liver cancer after several decades of infection, but now this is preventable. It is known that the best way to treat people who have active HBV infection is to give them powerful drugs that stop the virus replicating. The old drugs that used to be used to treat hepatitis B (such as lamivudine) are very cheap but they are not very powerful and many people develop resistance to them so that they stop working after a period of time. The new drugs that are used to treat hepatitis B are more powerful and most people do not develop resistance. However, the new drugs may have more side effects in the long term and they are much more expensive. In many infections we know that starting treatment with a powerful drug and then reducing the treatment to a weaker drug is very effective. In London a small study of this approach has been tried in patients with hepatitis B with success in most cases. Specifically people who had been treated with the expensive new drug tenofovir and who were responding to treatment were changed to treatment with the weaker, cheaper, probably safer drug lamivudine. The great majority responded very well. The aim of this study is to find out whether this way of treating hepatitis B works in Lusaka, Zambia.

### Who can participate?

80 Zambian adults, both sexes and aged 18 years or more, with HBV infection, already with evidence of some liver damage (using a marker in blood called ALT).

### What does the study involve?

Participants take tenofovir, the best single drug currently available, for 12 months to suppress the virus, then step down to lamivudine treatment. The evidence suggests that after a period of complete viral suppression lamivudine failure may be much less likely. Participants are monitored very closely during the 6 months after stepping down to lamivudine.

### What are the possible benefits and risks of participating?

The main benefit is that participants receive at least 18 months of powerful antiviral therapy. The principal risks are associated with the liver biopsy (bleeding, but only 1 in 3000 have this

problem), and the possibility that the infection may flare up aggressively at the end of the study. Past experience suggests that this is very uncommon. If successful, this approach could lead to a much more affordable treatment approach for Africans with HBV infection, of whom there are many tens of thousands.

Where is the study run from?

University Teaching Hospital, Lusaka (Zambia)

When is the study starting and how long is it expected to run for?

January 2014 to September 2017

Who is funding the study?

Medical Research Council (UK)

Who is the main contact?

1. Prof. Graham Foster, principal investigator
2. Prof. Paul Kelly and Dr Bright Nsokolo, chief investigators in Lusaka

## Contact information

**Type(s)**

Scientific

**Contact name**

Prof Graham Foster

**Contact details**

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## Additional identifiers

**Protocol serial number**

Step HEP

## Study information

**Scientific Title**

Step-down affordable treatment for chronic hepatitis B infection in Africa: feasibility of treatment strategy

**Acronym**

Step HEP

**Study objectives**

That a treatment regime of tenofovir for 48 weeks followed by lamivudine for 24 weeks effectively suppresses hepatitis B virus (HBV) replication in more than 50% of patients.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

University of Zambia Biomedical Research Ethics Committee, 24/05/2013, ref: 005-02-13

**Study design**

Single group evaluation of feasibility of treatment strategy

**Primary study design**

Interventional

**Study type(s)**

Treatment

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

Hepatitis B infection

**Interventions**

Patients will be given Tenofovir 300mg daily (orally) for 52 weeks and then lamivudine 100mg (orally) daily for 26 weeks. Patients will be followed up for 6 months after starting lamivudine.

**Intervention Type**

Drug

**Phase**

Not Applicable

**Drug/device/biological/vaccine name(s)**

Tenofovir, lamivudine

**Primary outcome(s)**

Proportion of patients who successfully step down to lamivudine monotherapy with virological control of replication throughout

**Key secondary outcome(s)**

1. Proportion of patients who, even if there is virological rebound, achieve successful control on re-introduction of tenofovir
2. Accuracy of ALT monitoring in comparison with viral load monitoring
3. Accuracy of HBsAg quantification compared to viral load monitoring

The primary and secondary outcomes will be assessed by virological measurements in blood samples obtained every 3 months.

**Completion date**

01/09/2017

# Eligibility

## Key inclusion criteria

1. HBV viral load >105 copies/ml
2. Alanine aminotransferase (ALT) >1.3 times upper limit of normal (which sets the criterion at 45 i.u./l)
3. Evidence of inflammation on liver biopsy
4. May be either e antigen negative (n=40) or positive (n=40)

## Participant type(s)

Patient

## Healthy volunteers allowed

No

## Age group

Adult

## Sex

All

## Key exclusion criteria

1. Histological or radiological evidence of cirrhosis
2. HIV infection
3. History of alcohol abuse or histological evidence of alcoholic liver disease
4. History of any long-term drug ingestion
5. Histological evidence of metabolic liver disease (haemochromatosis, Wilsons disease, a1-antitrypsin deficiency) or autoimmune liver disease [antibodies to M2 antigen, Perinuclear antineutrophil cytoplasmic antibodies (p-ANCA), nuclear antigens, microsomes or smooth muscle]
6. Histological or radiological evidence of schistosomiasis
7. Histological evidence of hepatitis D virus (HDV) infection
8. Virological evidence of active hepatitis C virus (HCV) or hepatitis E virus (HEV) infection

## Date of first enrolment

06/01/2014

## Date of final enrolment

31/12/2015

# Locations

## Countries of recruitment

Zambia

## Study participating centre

University Teaching Hospital

Lusaka

Zambia

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## Sponsor information

### Organisation

Queen Mary University of London (UK)

### ROR

<https://ror.org/026zzn846>

## Funder(s)

### Funder type

Research council

### Funder Name

Medical Research Council (MR/K007394/1)

### Alternative Name(s)

Medical Research Council (United Kingdom), UK Medical Research Council, Medical Research Committee and Advisory 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?
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<a href="#">Other publications</a>	baseline liver pathology	01/09/2017	10/05/2021	Yes	No
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