

A pilot study of Combivir® therapy for patients with primary biliary cirrhosis

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 15/11/2011	Condition category Digestive System	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0265092544

Study information

Scientific Title

Study objectives

This study will further test the hypothesis that there is an infectious aetiology involved in the development of primary biliary cirrhosis (PBC) by undertaking a randomised, controlled, phase II pilot study of Combivir® therapy in patients with PBC.

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

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Treatment

Participant information sheet**Health condition(s) or problem(s) studied**

Primary biliary cirrhosis (PBC)

Interventions

This investigation is designed as a randomised, controlled, phase II pilot study of Combivir® in approximately 60 patients with PBC. It is expected that the majority of PBC patients will already be taking ursodeoxycholic acid and patients enrolled in the study will have been on this treatment for at least 6 months.

Patients will be randomised to continue with ursodeoxycholic acid alone or in combination with Combivir®. The clinical, virological, histological and immune effects of the study drug will be examined. The clinical end point of the study will be 1 year of therapy or evidence for developing end stage liver disease. All PBC patients except for those with decompensated liver disease will be enrolled in the study after obtaining an informed written consent.

The PBC patients will already be on ursodeoxycholic acid at an adjusted dose of 13 - 15 mg/kg of body weight/day in 2 - 3 divided doses. Patients treated with Combivir® will receive one tablet twice a day: Lamivudine 150 mg and Zidovudine 300 mg twice a day. Those patients not on ursodeoxycholic acid at the start of the study will be treated with ursodeoxycholic acid at the dose indicated for a period of 6 months prior to randomisation to ursodeoxycholic acid alone or in combination with Combivir® twice a day.

At enrolment, each patient with PBC will be assessed for the inclusion criteria. Prior to therapy, patients will have a thorough history taken to assess symptoms. An objective graded clinical parameter scale will include the development, presence or worsening of pruritus, fatigue, sicca syndrome or right upper quadrant pain. At the same time patients will be examined for the presence or development of overt clinical signs such as jaundice, splenomegaly or hepatomegaly. At this point, the baseline blood tests will include: full blood count (FBC), reticulocyte count, prothrombin time (PT), erythrocyte sedimentation rate (ESR), blood urea nitrogen (BUN), creatinine, sodium, potassium, calcium, phosphate, albumin, total protein, bilirubin, aspartate aminotransferase (AST), alanine aminotransferase (ALT), alkaline phosphatase, cholesterol, creatine kinase (CK), amylase, immunoglobulins, antinuclear antibodies (ANA), and quantitated AMA. All of these are laboratory assessments that are clinically useful in following patients with PBC.

Unless problems develop in the interim, patients will be seen at months 1, 3, 6, 9 and 12 after initiation of therapy. At the initial clinic visit, blood will be drawn for BUN, creatinine, electrolytes, amylase, bilirubin, AST, ALT, alkaline phosphatase, albumin, PT, serum lactate and FBC. Subsequently, samples will be drawn for hepatic biochemistry, as well as virological and immunological studies. Patients will undergo a physical exam at each visit and also be questioned about changes in symptoms. Liver biopsies will be performed as clinically indicated.

Response to therapy will be based on changes in symptomatology, development of overt clinical signs, immunological parameters, improvement of liver function and biochemistry. Immunological studies include quantitative AMA levels. Reverse transcription polymerase chain reaction (RT-PCR) and Western blot virological studies will be performed on serum samples before and after therapy in the Hepatitis Research Laboratory, Alton Ochsner Medical Foundation, USA.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Ursodeoxycholic acid, Combivir® (lamivudine and zidovudine)

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/10/2001

Completion date

01/05/2005

Eligibility

Key inclusion criteria

Patients will be recruited from the Liver Out-Patients Department at the Queen Elizabeth Hospital:

1. Patients greater than 20 years old of either sex
2. Elevated alkaline phosphatase or alanine aminotransferase (ALT) within 3 months prior to the start of therapy
3. Positive serum anti-mitochondrial antibodies (AMA) (titre greater than 1:20)
4. Liver biopsy histology compatible with PBC

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

60

Key exclusion criteria

1. Patients treated with immunosuppressive or anti-inflammatory agents
2. Advance liver disease: Child's class B or C
3. Patients with secondary hepatological diagnosis
4. Alcohol abuse (greater than 50 g of alcohol per day)
5. Other significant co-morbidity (e.g. cardiac or renal failure)
6. Pregnancy or breast feeding
7. Sexually active female of child bearing age not using effective contraception

Date of first enrolment

01/10/2001

Date of final enrolment

01/05/2005

Locations**Countries of recruitment**

England

United Kingdom

Study participating centre

Liver Medicine

Birmingham

United Kingdom

B15 2TH

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Industry

Funder Name

University Hospital Birmingham NHS Trust (UK)

Funder Name

NHS R&D Support Funding

Funder Name

GlaxoSmithKline (GSK) (UK)

Alternative Name(s)

GlaxoSmithKline plc., GSK plc., GSK

Funding Body Type

Government organisation

Funding Body Subtype

For-profit companies (industry)

Location

United Kingdom

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?
Results article	results	01/12/2004		Yes	No