Optimising portal haemodynamics and liver parenchymal perfusion during orthotopic liver transplantation (OTL) using an implantable pump

Submission date	Recruitment status	Prospectively registered
30/09/2005	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2005	Completed	Results
Last Edited	Condition category	Individual participant data
15/02/2018	Surgery	[] Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Prof Brian R Davidson

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0256107980

Study information

Scientific Title

Optimising portal haemodynamics and liver parenchymal perfusion during orthotopic liver transplantation (OTL) using an implantable pump

Study objectives

Whether hepatic tissue oxygenation and perfusion of hepatic microcirculation can be improved with a short term mechanical augmentation of portal flow.

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)

Hospital

Study type(s)

Treatment

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

Liver transplantation

Interventions

Randomised controlled trial

Intervention Type

Procedure/Surgery

Phase

Not Applicable

Primary outcome measure

Service outcomes development

Secondary outcome measures

Not provided at time of registration

Overall study start date

15/01/2003

Completion date

30/09/2003

Eligibility

Key inclusion criteria

12 patients (6 patients with portal pump and 6 controls)

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

12

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

15/01/2003

Date of final enrolment

30/09/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Royal Free Hampstead NHS Trust

London

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type

Government

Funder Name

The Royal Free Hampstead NHS Trust (UK)

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