

Assessment of the lipid lowering agents on plasma homocysteine concentration and renal function

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 17/04/2015	Condition category Circulatory System	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0211136321

Study information

Scientific Title

Assessment of the lipid lowering agents on plasma homocysteine concentration and renal function

Study objectives

Are the benefits of fibrate therapy offset by an increase in plasma homocysteine?

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

Health condition(s) or problem(s) studied

Cardiovascular: Hyperlipidemia

Interventions

3 groups:

1. Group 1 = Fibrate 1
2. Group 2 = Fibrate 2
3. Group 3 = Statin (Control)

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Lipid lowering agents

Primary outcome measure

Measurements of renal function, plasma homocysteine and lipids over an 8-month period

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/11/2003

Completion date

30/09/2005

Eligibility

Key inclusion criteria

Adult patients attending the lipid clinic with raised cholesterol and triglycerides

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

60

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/11/2003

Date of final enrolment

30/09/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Royal Surrey County Hospital Trust
Guildford

United Kingdom
GU2 5XX

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

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

Funder(s)

Funder type

Government

Funder Name

Royal Surrey County Hospital 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