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Analysis of the duration of combination therapy that is necessary for hepatitis C (HCV) genotype 1 eradication

Submission date	Recruitment status	[_] Prospectively registered
30/09/2004	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
30/09/2004	Completed	[_] Results
Last Edited	**Condition category**	[_] Individual participant data
18/07/2016	Infections and Infestations	[_] Record updated in last year

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
N0263132921

Study information

Scientific Title
Analysis of the duration of combination therapy that is necessary for hepatitis C (HCV) genotype 1 eradication

Study objectives
What is the duration of combination therapy required in order to produce a sustained viral clearance in hepatitis C infected patients

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 contact details to request a participant information sheet

Health condition(s) or problem(s) studied
Infections and Infestations: Hepatitis C

Interventions
Randomised Controlled Trial
1. Chemotherapy 1
2. Chemotherapy 2

Intervention Type
Drug

Phase

Not Applicable

Primary outcome measure
Viral clearance

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/10/2003

Completion date
01/10/2005

Eligibility

Key inclusion criteria
50 patients from Hepatology

Participant type(s)
Patient

Age group
Not Specified

Sex
Both

Target number of participants
50

Key exclusion criteria
Does not match inclusion criteria

Date of first enrolment
01/10/2003

Date of final enrolment
01/10/2005

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

University College London
London
United Kingdom
WC1E 6HX

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
Hospital/treatment centre

Funder Name
University College London Hospitals 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