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The safety and comfort of early mobilisation following day case coronary angiography

Submission date 29/09/2006	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 29/09/2006	**Overall study status** Completed	
Last Edited 23/04/2015	**Condition category** Surgery	

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
N0236164472

Study information

Scientific Title
The safety and comfort of early mobilisation following day case coronary angiography

Study objectives
It is hypothesised that complications will not be increased by early mobilisation and that patient comfort will be improved.

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
Surgery: Coronary angiography

Interventions
Randomisation of patients to either control or experimental group. Control group will mobilise at 4 hours (current practice) experimental group will mobilise at 1.5 hours. Other than ambulation time, each group will be treated identically.

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
1. Haematoma size
2. Complication rate: pseudoaneurysm, presence of bleeding.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/11/2004

Completion date
30/09/2005

Eligibility

Key inclusion criteria
All patients able to attend for day case angiography.

Participant type(s)
Patient

Age group
Adult

Sex
Not Specified

Target number of participants
Not provided at time of registration

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/11/2004

Date of final enrolment
30/09/2005

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
St George's Hospital
London
United Kingdom
SW17 0QT

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2006 Update - Department of Health

Sponsor details
The Department of Health, Richmond House, 79 Whitehall
London
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dhmail@doh.gsi.org.uk

Sponsor type
Government

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

Funder(s)

Funder type
Government

Funder Name
St George's Healthcare NHS Trust (UK), NHS R&D Support Funding

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