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Randomised comparison of radial versus femoral access for coronary angioplasty in the ReoPro era

Submission date	Recruitment status	[ ] Prospectively registered
12/09/2003	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
12/09/2003	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
18/10/2016	Surgery	[ ] 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
N0205116657

Study information

Scientific Title
Randomised comparison of radial versus femoral access for coronary angioplasty in the ReoPro era

Acronym
DA7RAD study

Study objectives
When using ReoPro during percutaneous coronary intervention (PCI):
1. Is the vascular complication rate less than when radial access rather than femoral access?
2. Is mobilisation early safe, is patient satisfaction 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)
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
Percutaneous coronary angioplasty

Interventions
Randomised prospective clinical trial: radial versus femoral access for coronary angioplasty.

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
1. Access site complications
2. Time to mobilisation
3. Patient satisfaction

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/11/2002

Completion date
01/06/2004

Eligibility

Key inclusion criteria
Patients undergoing percutaneous coronary angioplasty at Barts and The London NHS Trust

Participant type(s)
Patient

Age group
Not Specified

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
01/11/2002

Date of final enrolment
01/06/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
St Bartholomew's Hospital
London
United Kingdom
EC1A 7BE

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
Government

Funder Name
Barts and The London 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