

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

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 18/10/2016	Condition category Surgery	<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

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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