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

Submission date 12/09/2003	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 12/09/2003	**Overall study status** Completed	
Last Edited 18/10/2016	**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

Protocol serial number
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

Study type(s)
Treatment

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(s)
1. Access site complications
2. Time to mobilisation
3. Patient satisfaction

Key secondary outcome(s))
Not provided at time of registration

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

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Not Specified

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
United Kingdom

England

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

Sponsor information

Organisation
Department of Health (UK)

Funder(s)

Funder type
Government

Funder Name
Barts and The London NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration