

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
Registration date 12/09/2003	Overall study status Completed	☐ Statistical analysis plan
		☐ Results
Last Edited 18/10/2016	Condition category Surgery	☐ Individual participant data
		☐ 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

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