Angio-seal versus vaso-seal for haemostasis

Submission date Recruitment status Prospectively registered 12/09/2003 No longer recruiting [] Protocol [] Statistical analysis plan Registration date Overall study status 12/09/2003 Completed [X] Results [] Individual participant data Last Edited Condition category 24/08/2012 Haematological Disorders

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0176120413

Study information

Scientific Title

Study objectives

Is using a sealing device superior to femoral compression for haemostasis following angiography or angioplasty?

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 the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Haemostasis

Interventions

Randomised contolled trial. Sealed envelopes will be used with the different randomisations.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

The time to obtain haemostasis and development of any haemotoma during a 1 week period.

Secondary outcome measures

Not provided at time of registration

Overall study start date

Completion date

26/02/2004

Eligibility

Key inclusion criteria

300 patients

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

300

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

26/02/2003

Date of final enrolment

26/02/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Radiology Department

Oxford United Kingdom OX3 9DU

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

Hospital/treatment centre

Funder Name

Oxford Radcliffe Hospitals 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/02/2007		Yes	No