ISRCTN96045625 https://doi.org/10.1186/ISRCTN96045625

Angio-seal versus vaso-seal for haemostasis

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	[X] Results
Last Edited	**Condition category**	[ ] Individual participant data
24/08/2012	Haematological Disorders	

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
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

26/02/2003

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