

Randomised trial of Polytetrafluoroethylene (PTFE) patch saphenoplasty in patients having surgery for recurrent long saphenous varicose veins

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 checked="" type="checkbox"/> Results
Last Edited 17/03/2008	Condition category Surgery	<input type="checkbox"/> Individual participant data

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

N0106092280

Study information

Scientific Title

Study objectives

To compare the use of a PTFE patch with normal repair for surgery for recurrent varicose veins.

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

Health condition(s) or problem(s) studied

Surgery: Long saphenous varicose veins

Interventions

1. Polytetrafluoroethylene patch
2. Flush ligation

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

To reduce the re-recurrence rate after reoperative surgery for recurrent long saphenous varicose veins.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/03/2003

Completion date

28/02/2007

Eligibility

Key inclusion criteria

At least 50 patients having surgery for recurrent long saphenous varicose veins.

Participant type(s)

Patient

Age group

Senior

Sex

Not Specified

Target number of participants

50

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/03/2003

Date of final enrolment

28/02/2007

Locations

Countries of recruitment

England

United Kingdom

Study participating centre**Consultant Surgeon**

Gloucester

United Kingdom

GL1 3NN

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

Gloucestershire R&D Consortium (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	01/09/2007		Yes	No

