

Does applying limb compression before long saphenous vein stripping result in a better outcome than applying it after stripping?

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 14/06/2017	Condition category Surgery	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0143157279

Study information

Scientific Title

Does applying limb compression before long saphenous vein stripping result in a better outcome than applying it after stripping?

Study objectives

To find out whether applying limb compression before the long saphenous vein is stripped results in a better postoperative outcome than applying it at the end of the procedure.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Double-blinded block randomisation pilot study

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: Varicose vein

Interventions

Limb compression before the long saphenous vein is stripped compared to after.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Post-operative outcome.

Secondary outcome measures

Not provided at time of registration

Overall study start date

17/05/2004

Completion date

01/04/2006

Eligibility

Key inclusion criteria

25 patients requiring varicose vein surgery

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

25

Key exclusion criteria

Patients requiring saphenopopliteal ligation as well

Date of first enrolment

17/05/2004

Date of final enrolment

01/04/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Watford General Hospital

Watford

United Kingdom

WD18 0HB

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

West Hertfordshire 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