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A randomised controlled trial investigating the role of subfascial endoscopic perforator vein surgery (SEPS) in the prevention of recurrence in recurrent long saphenous varicose veins

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 30/09/2016	**Condition category** Circulatory System	[_] 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0211091333

Study information

Scientific Title
A randomised controlled trial investigating the role of subfascial endoscopic perforator vein surgery (SEPS) in the prevention of recurrence in recurrent long saphenous varicose veins

Study objectives
To evaluate the advantages of SEPS to the standard open procedure - recurrent long saphenous 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
Not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied
Cardiovascular: Recurrent long saphenous varicose veins

Interventions
Randomised controlled trial randomised into two groups:
Group 1: subfascial endoscopic perforator vein surgery (SEPS)
Group 2: standard open procedure

Intervention Type
Procedure/Surgery

Primary outcome measure

Improvement in function measured by refill time, incidence of recurrent reflux, clinical recurrence, and cosmetic results.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/09/2000

Completion date
31/03/2004

Eligibility

Key inclusion criteria
Patients with recurrent varicose veins, undergoing surgery, with Duplex proven incompetent perforating veins

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
Not provided at time of registration

Key exclusion criteria
Does not meet inclusion criteria

Date of first enrolment
01/09/2000

Date of final enrolment
31/03/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

The Whiteley Clinic
Guildford
United Kingdom
GU2 7RF

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
Royal Surrey County Hospital NHS Trust

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