

Evaluation of the practice of long saphenous vein stripping in preventing varicose vein recurrence following varicose vein surgery

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 type="checkbox"/> Results
Last Edited 29/04/2016	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

N0263107648

Study information

Scientific Title

Evaluation of the practice of long saphenous vein stripping in preventing varicose vein recurrence following varicose vein surgery

Study objectives

Is stripping of the long saphenous vein necessary in order to prevent recurrence?

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

Surgery: Varicose vein recurrence

Interventions

Stripping vs usual practice

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Varicose vein recurrence

Secondary outcome measures

Not provided at time of registration

Overall study start date

22/01/2002

Completion date

01/06/2005

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

150 patients from Vascular Surgery.

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

22/01/2002

Date of final enrolment

01/06/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

The Middlesex Hospital
London
United Kingdom
W1N 8AA

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

University College London 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