

# Randomised single blind patient controlled trial of trans-illuminated powered suction phlebectomy versus conventional hook phlebectomy

<b>Submission date</b> 12/09/2003	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 12/09/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 13/09/2013	<b>Condition category</b> Circulatory System	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

### 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**

N0192119060

# **Study information**

## **Scientific Title**

## **Study objectives**

To determine whether TRIVEX reduces operative times when compared with hook phlebectomies.

## **Ethics approval required**

Old ethics approval format

## **Ethics approval(s)**

Not provided at time of registration

## **Study design**

Randomised single blind patient 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**

Cardiovascular: Phlebectomy

## **Interventions**

Trans-illuminated powered suction phlebectomy versus conventional hook phlebectomy

## **Intervention Type**

Other

## **Phase**

Not Specified

## **Primary outcome measure**

Operative time, visual analogue scales for pain and bruising and cost of procedure.

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

31/07/2002

**Completion date**

01/04/2004

## 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**

50

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

31/07/2002

**Date of final enrolment**

01/04/2004

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

Department of Vascular Surgery

Nottingham

United Kingdom

NG7 2UH

# 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

Queen's Medical Centre University Hospital 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