Antibiotic prophylaxis in varicose vein surgery

Submission date [] Prospectively registered Recruitment status 30/09/2004 No longer recruiting [] Protocol [] Statistical analysis plan Registration date Overall study status 30/09/2004 Completed [X] Results [] Individual participant data Last Edited Condition category 05/01/2010 Surgery

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

Protocol serial number

N0084125086

Study information

Scientific Title

A double blinded randomised controlled trial of antibiotic prophylaxis in varicose vein surgery

Study objectives

To compare antibiotic prophylaxis with no antibiotic, with primary endpoint being the rate of wound infection.

Please note that as of 13/02/2009 this record was extensively updated. All amendments can be found in the relevant field under the above update date.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Added 13/02/2009: Hull and East Riding Local Research Ethics Committee gave approval on the 23rd September 2002 (ref: LREC/08/02/135)

Study design

Double blinded randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Surgery: Varicose vein

Interventions

- 1. Antibiotics (Augmentin 1.2 g)
- 2. No antibiotics

The treatment was a one-off treatment (varicose vein surgery), which lasted a range of 44 - 65 minutes. Patients were seen once at 14 days post-operation.

Intervention Type

Procedure/Surgery

Phase

Not Applicable

Primary outcome(s)

Added 13/02/2009:

The degree of wound complications, determined by an adapted version of the ASEPSIS wound scoring system.

Key secondary outcome(s))

Added 13/02/2009:

- 1. Visit to the General Practitioner for a wound-related problem
- 2. The requirement of antibiotics in the post-operative period for a perceived wound infection

Completion date

01/06/2006

Eligibility

Key inclusion criteria

Added 13/02/2009:

- 1. Aged 18 years or older, either sex
- 2. Undergoing groin surgery for varicose veins. All patients with varicosities of the greater saphenous vein (GSV) listed for saphenofemoral ligation, stripping of the GSV and phlebectomies were eligible to enter the trial.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

All

Key exclusion criteria

Added 13/02/2009:

- 1. Patients whose surgery did not include a groin incision
- 2. Patients below the age of 18 years
- 3. Pregnancy or lactation
- 4. Penicillin allergy
- 5. Receiving antibiotics for other indications

Date of first enrolment

09/04/2003

Date of final enrolment

01/06/2006

Locations

Countries of recruitment

United Kingdom

England

Study participating centre Dept of Vascular Surgery

Hull United Kingdom HU3 2JZ

Sponsor information

Organisation

Department of Health

Funder(s)

Funder type

Government

Funder Name

The North and South Bank Research and Development Consortium (UK)

Results and Publications

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 revi	ewed? Patient-facing?
Results article	results	01/01/2010	Yes	No
Participant information sheet	Participant information sheet	11/11/2025 11/11	/2025 No	Yes