

LigaSure technique of haemorrhoidectomy versus Ferguson Clinic haemorrhoidectomy: a randomised prospective study

Submission date 28/09/2007	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 28/09/2007	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 17/10/2016	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

N0436165617

Study information

Scientific Title

LigaSure technique of haemorrhoidectomy versus Ferguson Clinic haemorrhoidectomy: a randomised prospective study

Study objectives

The study is designed to compare Ferguson clinic closed haemorrhoidectomy with LigaSure haemorrhoidectomy. The primary end points of the study is the assessment of postoperative pain, morbidity and time to full recovery.

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

Health condition(s) or problem(s) studied

Surgery: Haemorrhoidectomy

Interventions

Ferguson clinic closed haemorrhoidectomy vs LigaSure haemorrhoidectomy.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

1. Visual analogue of postoperative pain on a daily basis for one week, followed by pain scores at two and four weeks as well as overall satisfaction
2. Pre and postoperative anorectal physiology and endoanal ultrasonography (12 weeks post operative)
3. Postoperative complications: a) infection; b) wound dehiscence; c) haemorrhage

Secondary outcome measures

No secondary outcome measures

Overall study start date

01/01/2003

Completion date

01/01/2004

Eligibility

Key inclusion criteria

Subjects with history of 3rd degree haemorrhoids.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/01/2003

Date of final enrolment

01/01/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
LIMIT - Leeds Institute for Minimally Invasive The
Leeds
United Kingdom
LS1 3EX

Sponsor information

Organisation

Record Provided by the NHSTCT Register - 2007 Update - Department of Health

Sponsor details

The Department of Health, Richmond House, 79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
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Sponsor type

Not defined

Website

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

Funder(s)

Funder type

Government

Funder Name

Leeds Teaching Hospitals 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