

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

Submission date 28/09/2007	Recruitment status No longer recruiting	☐ Prospectively registered
		☐ Protocol
Registration date 28/09/2007	Overall study status Completed	☐ Statistical analysis plan
		☐ Results
Last Edited 17/10/2016	Condition category Surgery	☐ Individual participant data
		☐ 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

Protocol serial number

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

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Treatment

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(s)

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

Key secondary outcome(s)

No secondary outcome measures

Completion date

01/01/2004

Eligibility

Key inclusion criteria

Subjects with history of 3rd degree haemorrhoids.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

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**

United Kingdom

England

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

Funder(s)

Funder type

Government

Funder Name

Leeds Teaching Hospitals NHS Trust

Results and Publications

Individual participant data (IPD) sharing plan**IPD sharing plan summary**

Not provided at time of registration