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

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

Study design

Randomised controlled trial

Primary study design

Interventional

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