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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 [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 28/09/2007	**Overall study status** Completed	
Last Edited 17/10/2016	**Condition category** Surgery	

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
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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
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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