Effect of laryngeal mask airway (LMA) in small airway function in the postoperative period compared with the endotracheal tube (ETT)

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
30/04/2018	Surgery	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Prof Iain T Campbell

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

Effect of laryngeal mask airway (LMA) in small airway function in the postoperative period compared with the endotracheal tube (ETT)

Study objectives

To compare the effect of laryngeal mask airway versus end tracheal tube on small airway function in the post operative period.

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

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Surgery: Small airway function

Interventions

Patients will be randomised to either LMA or ET for anaesthesia.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Changes in airway resistance in the lung.

Secondary outcome measures

If there are marked differences it will add a new dimension to the choice of airway device used in patients with constrictive/obstructive airways disease.

Overall study start date

05/02/2007

Completion date

05/02/2009

Eligibility

Key inclusion criteria

- 1. ASA Grades 1 or 2 (ie healthy patients of patients with well controlled chronic conditions) for elective surgery expected to last >30mins, on peripheral or body surfaces. We expect this would mean mainly orthopaedic or plastic surgery patients but could include vascular (varicose veins)
- 2. Smokers and non-smokers
- 3. BMI < 30
- 4. Patients consenting to the research
- 5. No history of gastric reflux

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

80 participants; 2 groups of 40. 40 patients recruited as of August 2008.

Key exclusion criteria

- 1. ASA 3 or higher
- 2. BMI >30
- 3. Patients suffering with gastro-oesophageal reflux
- 4. Patients unable to consent to the research or those who refuse to participate

Date of first enrolment

05/02/2007

Date of final enrolment

05/02/2009

Locations

Countries of recruitment

England

United Kingdom

Study participating centre University Hospital of South Manchester NHS Foundation Trust Manchester United Kingdom M23 9LT

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 dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

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

Funder(s)

Funder type

Government

Funder Name

University Hospital of South Manchester NHS Foundation Trust (UK), NHS R&D Support Funding

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