

To ascertain the usefulness of the Aintree Catheter as a means of intubation and to compare its usage via a laryngeal mask (LMA) and an intubating laryngeal mask (ILMA)

Submission date 30/09/2004	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 15/04/2015	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

Contact name

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Contact details

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United Kingdom
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0190137105

Study information

Scientific Title

To ascertain the usefulness of the Aintree Catheter as a means of intubation and to compare its usage via a laryngeal mask (LMA) and an intubating laryngeal mask (ILMA)

Study objectives

To compare ease of intubation using an Aintree catheter through a laryngeal mask (LMA) and an intubating laryngeal mask (ILMA).

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: Anaesthesia

Interventions

Aintree Catheter vs laryngeal mask (LMA) and intubating laryngeal mask (ILMA)

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Time taken to achieve intubation

Secondary outcome measures

Not provided at time of registration

Overall study start date

30/10/2004

Completion date

30/06/2006

Eligibility

Key inclusion criteria

Patients undergoing General Anaesthesia

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

n = 60, power = 80%

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

30/10/2004

Date of final enrolment

30/06/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

The Queen Victoria Hospital NHS Trust

East Grinstead

United Kingdom

RH19 3DZ

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

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

Funder(s)

Funder type

Government

Funder Name

Queen Victoria Hospital NHS Trust (UK)

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