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A randomised crossover comparison of the Cobra Perilaryngeal Airway (CobraPLA®) and Classic Laryngeal Mask Airway™ during anaesthesia in ventilated patients

Submission date
30/09/2005

Recruitment status
Stopped

Registration date
30/09/2005

Overall study status
Stopped

Last Edited
06/03/2009

Condition category
Surgery

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[X] Results
[ ] Individual participant data
[ ] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
N0212140555

Study information

Scientific Title

Study objectives
Does the CobraPLA® perform as well as the Classic Laryngeal Mask Airway™ during anaesthesia?

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: Anaesthesia

Interventions
CobraPLA® and Classic Laryngeal Mask Airway™

Intervention Type
Procedure/Surgery

Phase
Not Applicable

Primary outcome(s)
Not provided at time of registration

Key secondary outcome(s))
Not provided at time of registration

Completion date
13/08/2004

Reason abandoned (if study stopped)
Terminated due to two cases of pulmonary aspiration during this and a related study which lead to the decision that the risk to the participants of continuing the study outweighed the potential benefits.

Eligibility

Key inclusion criteria

1. Patients undergoing anaesthesia with ventilation of the lungs
2. Patients have been included if they were undergoing elective anaesthesia for surgery

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
01/05/2004

Date of final enrolment
13/08/2004

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Anaesthesia
Bath
United Kingdom
BA1 3NG

Sponsor information

Organisation
Department of Health

Funder(s)

Funder type
Government

Funder Name
Royal United Hospital Bath NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/08/2005		Yes	No