Laryngeal mask vs endotracheal tube for bariatric surgery

Submission date	Recruitment status	Prospectively registered
26/08/2008	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
18/09/2008	Completed	Results
Last Edited	Condition category	Individual participant data
18/09/2008	Nutritional, Metabolic, Endocrine	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Ulderico Freo

Contact details

Istituto di Anestesiologia Dipartimento di Farmacologia e Anestesiologia Via C. Battisti, 267 Padova Italy 35121 +39 (0)49 821 3094 ulderico.freo@unipd.it

Additional identifiers

Protocol serial number

1615P

Study information

Scientific Title

Comparison study on ventilation with laryngeal mask and with endotracheal tube for bariatric surgery

Study objectives

Ventilation with laryngeal mask results in better recovery profile than ventilation with endotracheal tube in obese patients undergoing bariatric surgery.

Ethics approval required

Old ethics approval format

Ethics approval(s)

This study was approved by the Ethics Committee, Padua Hospital on 25/07/2008.

Study design

Randomised double-blind (patient, evaluating physician) trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Obesity

Interventions

Ventilation with laryngeal mask vs ventilation with endotracheal tuber

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

1. Time to independent walking after the end of anesthesia

Key secondary outcome(s))

- 1. Ventilatory parameters
- 2. Postoperative morbidity
- 3. Patient and surgeon satisfaction

Completion date

08/01/2011

Eligibility

Key inclusion criteria

- 1. Both males and females, age >18 years
- 2. American Society of Anaesthesiologists (ASA) classification I-III
- 3. Obese patients (body mass index [BMI] >30) undergoing bariatric surgery
- 4. Written informed consent

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

All

Key exclusion criteria

- 1. Psychiatric disorders
- 2. Drug abuse
- 3. Known or possible pregnancy
- 4. Previous airway pathology
- 5. Pulmonary diseases
- 6. History of gastro-oesophageal reflux

Date of first enrolment

08/01/2008

Date of final enrolment

08/01/2011

Locations

Countries of recruitment

Italy

Study participating centre Istituto di Anestesiologia

Padova Italy 35121

Sponsor information

Organisation

Department of Pharmacology and Anaesthesiology, Padua University (Italy)

ROR

https://ror.org/00240q980

Funder(s)

Funder type

University/education

Funder Name

Department of Pharmacology and Anaesthesiology, Padua University (Italy)

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? Participant information sheet 11/11/2025 11/11/2025 No

Participant information sheet

Yes