Double-blind randomised controlled study to compare the outcomes of laparoscopic gastric banding and laparoscopic Roux-en-Y gastric bypass in morbidly obese patients attending the Multidisciplinary Morbid Obesity Clinic at King's College Hospital

Submission date	Recruitment status	Prospectively registered
30/09/2005	No longer recruiting	∐ Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2005	Completed	Results
Last Edited	Condition category	Individual participant data
30/04/2018	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

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Additional identifiers

Protocol serial number

Study information

Scientific Title

Double-blind randomised controlled study to compare the outcomes of laparoscopic gastric banding and laparoscopic Roux-en-Y gastric bypass in morbidly obese patients attending the Multidisciplinary Morbid Obesity Clinic at King's College Hospital

Study objectives

1. To compare the early outcome (first six months post-operatively) of the two operations with that of a matched group of morbidly obese patients who are on a low calorie diet only (controls) 2. To determine if there is any clinical and statistical difference in the outcomes of the two surgical procedures in the early, medium and long term (up to 5 years post-operatively)

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Double-blind randomised controlled study

Primary study design

Interventional

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Nutritional, Metabolic, Endocrine: Obesity

Interventions

T compare the outcomes of laparoscopic gastric banding and laparoscopic Roux-en-Y gastric bypass

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s))

Not provided at time of registration

Completion date

Eligibility

Key inclusion criteria

Not provided at time of registration

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

01/09/2003

Date of final enrolment

01/09/2009

Locations

Countries of recruitment

United Kingdom

England

Study participating centre Department of Surgery

London United Kingdom SE5 9RS

Sponsor information

Organisation

Department of Health

Funder(s)

Funder type

Government

Funder Name

Kings College Hospital NHS Trust R&D Consortium (UK)

Funder Name

NHS R&D support funding

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration