

Recovery profile and side effects of remifentanyl-based anaesthesia with desflurane or propofol for prolonged surgery

Submission date 12/09/2003	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 12/09/2003	Overall study status Completed	☐ Statistical analysis plan ☐ Results
Last Edited 09/12/2019	Condition category Surgery	☐ 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
N0190101946

Study information

Scientific Title

Recovery profile and side effects of remifentanil-based anaesthesia with desflurane or propofol for prolonged surgery

Study objectives

To compare the recovery profiles, post-operative analgesic requirements and side effects of remifentanil-based anaesthesia with desflurane or with propofol in cases of prolonged surgery.

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

Remifentanil-based anaesthesia with desflurane or with propofol.

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

15/10/2003

Eligibility

Key inclusion criteria

Patients having maxillofacial or plastic surgical procedures with an operating time greater than 4 hours

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

15/10/2001

Date of final enrolment

15/10/2003

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

The Queen Victoria Hospital NHS Trust

East Grinstead

United Kingdom

TN22 4BY

Sponsor information**Organisation**

Department of Health (UK)

Funder(s)**Funder type**

Government

Funder Name

Queen Victoria Hospital NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

The data sharing plans for the current study are unknown and will be made available at a later date

IPD sharing plan summary

Data sharing statement to be made available at a later date