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

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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

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

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

Remifentanil-based anaesthesia with desflurane or with propofol.

Intervention Type

Procedure/Surgery

Phase

Not Applicable

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

15/10/2001

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

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

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

England

United Kingdom

Study participating centre The Queen Victoria Hospital NHS Trust

East Grinstead United Kingdom TN22 4BY

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL

Sponsor type

Government

Website

http://www.doh.gov.uk

Funder(s)

Funder type

Government

Funder Name

Queen Victoria Hospital NHS Trust (UK)

Results and Publications

Publication and dissemination plan

A post-operative comparison between Desflurane and Propofol used as part of a Remifentanil based anaesthetic. Nene S, Patel C, Fenlon S. Poster presentation at Eurosiva meeting, Madrid June 2006

Intention to publish date

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