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

Submission date 12/09/2003	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 12/09/2003	**Overall study status** Completed	
Last Edited 09/12/2019	**Condition category** Surgery	

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