

Perioperative anxiety: a comparison between diazepam and propranolol

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 11/07/2016	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0280123688

Study information

Scientific Title

Perioperative anxiety: a comparison between diazepam and propranolol

Study objectives

Both drugs are anxiolytic, but propranolol should not cause cognitive/psychomotor dysfunction postoperatively, possibly allowing earlier patient discharge

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Prospective randomised double-blind controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Mental and Behavioural Disorders: Perioperative anxiety

Interventions

Patients undergoing surgery will be randomised to receive

1. Diazepam
2. Propranolol

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

1. Preoperative anxiety - Hospital Anxiety and Depression Score (HADS) and Visual Analogue Scale (VAS)

2. Pulse/blood pressure - postoperatively
3. VAS - postoperative nausea, vomiting, pain, sedation and satisfaction
4. Tests of cognitive/psychomotor dysfunction - postoperative

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/08/2003

Completion date

31/07/2004

Eligibility

Key inclusion criteria

80 inpatients

Participant type(s)

Patient

Age group

Not Specified

Sex

Both

Target number of participants

80

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/08/2003

Date of final enrolment

31/07/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Anaesthetics
Wirral
United Kingdom
CH49 5PE

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Wirral Hospitals NHS Trust (UK)

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration