

Propofol and remifentanil requirements after acute ethanol intake in clinical settings

Submission date 30/06/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 08/07/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 14/09/2009	Condition category Surgery	<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

Study information

Scientific Title

Acronym
PRAECS

Study objectives

Null hypothesis: Patients receiving intravenous ethanol would not require smaller doses of propofol and remifentanil for induction and maintenance of anaesthesia.

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)

Not Specified

Health condition(s) or problem(s) studied

Interaction of acute ethanol intravenous intake with propofol (general anesthetic) and remifentanil (opioid)

Interventions

Comparison of two groups (with different ethanol doses) versus control (no ethanol), regarding propofol and remifentanil requirements

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Propofol, remifentanil and acute ethanol

Primary outcome(s)

Reduction of propofol and remifentanil requirements

Key secondary outcome(s)

Degree of reduction, side-effects

Completion date

07/06/2004

Eligibility

Key inclusion criteria

Female surgical patients on general anesthesia

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

Alcoholic patients, patients in abstinence

Date of first enrolment

02/02/2004

Date of final enrolment

07/06/2004

Locations**Countries of recruitment**

Greece

Study participating centre

Korytsas 14

Thessaloniki

Greece

55133

Sponsor information**Organisation**

Theagenio Anticancer Hospital of Thessaloniki (Greece)

ROR

<https://ror.org/004hfxk38>

Funder(s)**Funder type**

Hospital/treatment centre

Funder Name

Theagenio Anticancer Hospital of Thessaloniki (Greece)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration