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Propofol and remifentanil requirements after acute ethanol intake in clinical settings

Submission date	Recruitment status	[ ] Prospectively registered
30/06/2005	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
08/07/2005	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
14/09/2009	Surgery	[ ] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Fotis Kanakoudis

Contact details
Korytsas 14
Thessaloniki
Greece
55133
+30 2310437531
fkanak@otenet.gr

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

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

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
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 measure
Reduction of propofol and remifentanil requirements

Secondary outcome measures
Degree of reduction, side-effects

Overall study start date
02/02/2004

Completion date
07/06/2004

Eligibility

Key inclusion criteria
Female surgical patients on general anesthesia

Participant type(s)
Patient

Age group
Adult

Sex
Female

Target number of participants
60

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)

Sponsor details
Al. Symeonidis 2
Thessaloniki
Greece
54007
+30 2310898213
amitragas@yahoo.com

Sponsor type
Hospital/treatment centre

Website
http://www.theagenio.gr

ROR
https://ror.org/004hfxk38

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Theagenio Anticancer Hospital of Thessaloniki (Greece)

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