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# Propofol and remifentanil requirements after acute ethanol intake in clinical settings

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

**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

### Primary study design
Interventional

### Study design
Randomised controlled trial

### 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