

Randomised, double-blind, placebo-controlled trial to identify any morphine-sparing effects of preoperative propacetamol in patients undergoing laparoscopic sterilisation by clip occlusion of fallopian tubes

Submission date 12/09/2003	Recruitment status Stopped	☐ Prospectively registered ☐ Protocol
Registration date 12/09/2003	Overall study status Stopped	☐ Statistical analysis plan ☐ Results
Last Edited 19/10/2011	Condition category Signs and Symptoms	☐ Individual participant data ☐ 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

Protocol serial number

N0226107795

Study information

Scientific Title

Study objectives

To demonstrate that preoperative administration of paracetamol (as intravenous propacetamol) in female patients undergoing laparoscopic sterilisation (by tubal clip occlusion) reduces postoperative morphine requirements.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised double-blind placebo-controlled trial

Primary study design

Interventional

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Signs and Symptoms: Pain

Interventions

1. Propacetamol
2. Placebo

July 2008: trial stopped, drug no longer available and replaced by an alternative agent.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

propacetamol

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s))

Not provided at time of registration

Completion date

30/09/2003

Eligibility

Key inclusion criteria

20 patients undergoing female laparoscopic sterilisation as a day-case and 20 controls.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Female

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/03/2002

Date of final enrolment

30/09/2003

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Department of Anaesthesia

Manchester

United Kingdom

M23 9LT

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

South Manchester University Hospitals NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration