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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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
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

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
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 measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/03/2002

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

Age group
Not Specified

Sex
Female

Target number of participants
40

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

England

United Kingdom

Study participating centre
Department of Anaesthesia
Manchester
United Kingdom
M23 9LT

Sponsor information

Organisation
Department of Health (UK)

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

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
South Manchester University 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