

Haloperidol as an antiemetic following abdominal hysterectomy - a comparison with Zofran®

Submission date 12/09/2003	Recruitment status Stopped	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Stopped	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 17/09/2012	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

Protocol serial number
N0558111116

Study information

Scientific Title

Study objectives

Following withdrawal of Droperidol can Haloperidol, which has been used in palliative care, give good antiemetic effect following acute surgery?

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)

Treatment

Health condition(s) or problem(s) studied

Post operative antiemetics

Interventions

Prospective randomised trial.

17/09/2012: Please note that this trial was stopped as the trial objectives were no longer viable

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Haloperidol, ondansetron (Zofran®)

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

30/09/2003

Reason abandoned (if study stopped)

Objectives no longer viable

Eligibility

Key inclusion criteria

All American Society of Anesthesiologists (ASA) I - III females undergoing abdominal hysterectomy who give their consent.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/10/2001

Date of final enrolment

30/09/2003

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Milton Keynes General NHS Trust

Milton Keynes

United Kingdom

MK6 5LD

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Milton Keynes General NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration