A randomised double blind study looking at combination granisetron and cyclizine antiemetic therapy versus single use granisetron or cyclizine in day case gynaecological patients

| Submission date | Recruitment status | Prospectively registered | | |
|-------------------------------|--|--|--|--|
| 12/09/2003 Registration date | No longer recruiting Overall study status | Protocol | | |
| | | Statistical analysis plan | | |
| 12/09/2003 | Completed | [X] Results | | |
| Last Edited | Condition category | [] Individual participant data | | |
| 06/01/2009 | Surgery | | | |

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number N0224112482

Study information

Scientific Title

Study objectives

Is a combination of prophylatic granisetron and cyclizine more effective in the prevention of post-operative nausea and vomiting than use as single agents in day case gynaecological patients?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised double blind study

Primary study design

Interventional

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Surgery: Gynaecological

Interventions

Combination granisetron and cyclizine anti-emetic therapy versus single use granisetron or cyclizine

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome(s)

To see if the combination anti-emetic regime reduces post-operative nausea and vomiting.

Key secondary outcome(s))

Not provided at time of registration

Completion date

01/06/2004

Eligibility

Key inclusion criteria

Patients over 18 years of age undergoing a day case gynaecological procedure.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

Female

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/06/2002

Date of final enrolment

01/06/2004

Locations

Countries of recruitment

United Kingdom

England

Study participating centre Consultant Anaesthetist

Torquay United Kingdom TQ2 7AA

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

South Devon Healthcare NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

| Output type | Details | Date created | Date added | Peer reviewed? | Patient-facing? |
|-----------------|---------|--------------|------------|----------------|-----------------|
| Results article | results | 01/11/2006 | | Yes | No |