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

Submission date
12/09/2003

Recruitment status
No longer recruiting

Registration date
12/09/2003

Overall study status
Completed

Last Edited
06/01/2009

Condition category
Surgery

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[X] Results
[_] Individual participant data

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
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

Primary study design
Interventional

Study design
Randomised double blind study

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