ISRCTN78140928 https://doi.org/10.1186/ISRCTN78140928

A randomised study of the optimal bowel preparation for routine capsule endoscopy using Citramag and Senna or Metoclopramide

Submission date
29/09/2006

Recruitment status
No longer recruiting

Registration date
29/09/2006

Overall study status
Completed

Last Edited
13/10/2011

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

Contact name
Dr Christopher Fraser

Contact details
Wolfson Unit for Endoscopy
North West London Hospitals NHS Trust
Watford Road
Harrow
United Kingdom
HA1 3UJ
+44 020 8235 4225
chris.fraser@imperial.ac.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0515171640

Study information

Scientific Title

Study objectives
To determine whether taking bowel preparation (Citramag and Senna) or a medicine to speed up transit through the stomach (Metoclopramide) will improve the quality of images seen, increase the transit through the small bowel, and increase the rate of completion of capsule endoscopy.

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

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
Surgery: Capsule endoscopy

Interventions
Randomised controlled study.

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Citramag and Senna or Metoclopramide

Primary outcome measure
Bowel visualisation, stomach and small intestinal transit time and capsule study completion rates (to the caecum)

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/12/2005

Completion date
01/12/2007

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
150 patients, 50 of which will be in the control group.

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/12/2005

Date of final enrolment
01/12/2007

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Wolfson Unit for Endoscopy
Harrow
United Kingdom
HA1 3UJ

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2006 Update - Department of Health

Sponsor details
The Department of Health, Richmond House, 79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Government

Funder Name
North West London 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/05/2009		Yes	No