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Randomised controlled trial of delayed colonoscopy following complete bowel preparation

Submission date	Recruitment status	[ ] Prospectively registered
30/09/2005	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
30/09/2005	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
19/10/2016	Surgery	[ ] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Mr Chun Kheng Khoo

Contact details
29 Showell Park
Taunton
United Kingdom
TA2 6BY
+44 (0)7734 204888
filhos@hotmail.com

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0249132151

Study information

Scientific Title
Randomised controlled trial of delayed colonoscopy following complete bowel preparation

Study objectives
The aim of this study is to evaluate the effect of increasing the delay between bowel preparation and colonoscopy.

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)
Hospital

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Colonoscopy

Interventions
1. Colonoscopy 12 hours after bowel preparation
2. Colonoscopy 24 hours after bowel preparation

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
Adverse effects, discomfort, bowel preparation, efficacy

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/11/2003

Completion date
01/06/2006

Eligibility

Key inclusion criteria
100 patients undergoing colonoscopy

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
100

Key exclusion criteria
Diabetes mellitus (unless solely diet controlled)

Date of first enrolment
01/11/2003

Date of final enrolment
01/06/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
29 Showell Park
Taunton
United Kingdom
TA2 6BY

Sponsor information

Organisation
Department of Health

Sponsor details
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
Taunton and Somerset Research and Development Consortium

Funder Name
NHS R&D Support Funding

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