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Randomised controlled trial of complete bowel preparation vs distal bowel cleansing in the management of patients referred under fast track guidelines for colorectal cancer

Submission date 30/09/2005	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol
Registration date 30/09/2005	**Overall study status** Completed	[_] Statistical analysis plan [_] Results
Last Edited 02/02/2018	**Condition category** Cancer	[_] Individual participant data [_] 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0249132150

Study information

Scientific Title
Randomised controlled trial of complete bowel preparation vs distal bowel cleansing in the management of patients referred under fast track guidelines for colorectal cancer

Study objectives
Whis is preferable in the management of patients undergoing flexible sigmoidoscopy - complete bowel preparation or distal bowel cleansing?

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)
Treatment

Participant information sheet
Not available in web format, please use the contact details to request a patient information sheet

Health condition(s) or problem(s) studied
Cancer: Colorectal

Interventions
1. Complete bowel preparation
2. Distal bowel cleansing

Intervention Type
Procedure/Surgery

Primary outcome measure

1. Patient satisfaction
2. 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
200 patients undergoing flexible sigmoidoscopy following fast track referral

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
200

Key exclusion criteria
Inability to self-administer enema, 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

Taunton & Somerset Hospital
Taunton
United Kingdom
TA1 5DA

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
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SW1A 2NL
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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 (UK) 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