

# Randomised controlled trial of delayed colonoscopy following complete bowel preparation

<b>Submission date</b> 30/09/2005	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 30/09/2005	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
<b>Last Edited</b> 19/10/2016	<b>Condition category</b> Surgery	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> 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**  
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  
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+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

