Randomised controlled trial on the use of multimodal strategies to accelerate recovery of patients undergoing colorectal resection

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	[X] Results		
Last Edited	Condition category	[] Individual participant data		
12/02/2010	Surgery			

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Mr Chun Kheng Khoo

Contact details

c/o The Doctors' Residences
Taunton & Somerset Hospital
Musgrove Park
Taunton
United Kingdom
TA1 5DA
+44 (0)7734 204888
filhos@hotmail.com

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

Study objectives

Does the use of multimodal strategies accelerate recovery of patients undergoing bowel resection?

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

Health condition(s) or problem(s) studied

Surgery: Colorectal resection

Interventions

- 1. Multimodal strategies for accelerated recovery (eg relates to fluids, analgesia, early mobilisation, etc).
- 2. Control: 'usual care'

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

- 1. EORTC QLQ-C30
- 2. Pain
- 3. Length of stay

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/04/2003

Completion date

31/10/2004

Eligibility

Key inclusion criteria

90 patients undergoing colorectal resection

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

90

Key exclusion criteria

- 1. Age under 16
- 2. Inability to walk 100m
- 3. Contraindication to epidural anaesthesia

Date of first enrolment

01/04/2003

Date of final enrolment

31/10/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre c/o The Doctors' Residences Taunton United Kingdom TA1 5DA

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 (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 summaryNot provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/06/2007		Yes	No