ISRCTN13770180 https://doi.org/10.1186/ISRCTN13770180

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

Submission date 30/09/2005	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 30/09/2005	**Overall study status** Completed	
Last Edited 12/02/2010	**Condition category** 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

N0249142477

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 summary

Not 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