

Laparoscopic assisted reversal of ileostomy versus traditional open reversal

Submission date 06/06/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 11/11/2010	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 30/10/2013	Condition category Surgery	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
NA

Study information

Scientific Title
Laparoscopic assisted reversal of ileostomy versus traditional open reversal: a pilot study

Acronym
LARI

Study objectives

To show that laparoscopic assisted reversal of ileostomy versus traditional open reduces hospital stay and morbidity.

Discharge criteria: having tolerated 2 large meals without nausea/vomiting, passed a bowel motion and adequate pain control on oral analgesia

Please note that as of 14/09/2012, the following changes were made to the record:

1. The scientific title of this trial was changed from "Laparoscopic assisted reversal of ileostomy versus traditional open reversal: a randomised control trial"
2. The target number of participants was changed from 46 to 70

Ethics approval required

Old ethics approval format

Ethics approval(s)

The Adelaide and Meath incorporating the National Children's Hospital (AMNCH) Ethics Committee, Tallaght Hospital approved on the 23rd of February 2010

Study design

Single centre two arm randomised active controlled parallel group trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Ileostomy

Interventions

Patients undergoing surgery for reversal of ileostomy will be randomised to receive either:

1. Traditional: Ileostomy mobilised from skin. Ileostomy excised followed by side to side anastomosis using GIA. Fascia and peritoneum closed and wound left to heal by secondary intention.
2. Laparoscopic assisted: Ileostomy mobilised from skin. Ileostomy excised followed by side to side anastomosis using GIA. One 5 mm port inserted contra-laterally with pneumoperitoneum. Reversal site checked for any omentum or bowel stuck to sutures. The whole abdominal cavity will be checked for any evidence of metastatic disease. Extra 5mm ports inserted and any adhesions present will be divided as well as any further intervention.

Intervention Type

Procedure/Surgery

Phase

Not Applicable

Primary outcome(s)

Length of stay

Key secondary outcome(s))

Morbidity, Diagnostic tool

Completion date

30/06/2011

Eligibility

Key inclusion criteria

1. Any person with a loop ileostomy who has been passed fit for surgery
2. Age 18-85

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Upper age limit

85 years

Sex

All

Key exclusion criteria

1. Inability to give informed consent (eg. Dementia)
2. American Society of Anaesthesiologists (ASA) greater than 3

Date of first enrolment

01/10/2009

Date of final enrolment

30/06/2011

Locations

Countries of recruitment

Ireland

Study participating centre

Dept of Surgery

Dublin

Sponsor information

Organisation
Adelaide and Meath Hospital, Incorporating the National Children's Hospital (AMNCH) (Ireland)

ROR
<https://ror.org/01fvmtt37>

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Adelaide and Meath Hospital, Incorporating the National Children's Hospital (AMNCH) (Ireland) - departmental and personal funds

Results and Publications

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	01/09/2013		Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes