

Prospective randomised trial of laparoscopic versus closed insertion of Tenckhoff catheters for peritoneal dialysis access

Submission date 12/09/2003	Recruitment status Stopped	☐ Prospectively registered ☐ Protocol
Registration date 12/09/2003	Overall study status Stopped	☐ Statistical analysis plan ☐ Results
Last Edited 22/11/2011	Condition category Surgery	☐ Individual participant data ☐ Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Mr S Sudhindran

Contact details
Box No 210
Transplant Unit
Addenbrooke's NHS Trust
Cambridge
United Kingdom
CB2 2QQ
+44 (0)7623 851784
sudhindran@bigfoot.com

Additional identifiers

Protocol serial number
N0544093499

Study information

Scientific Title

Study objectives

Laparoscopic versus closed insertion of peritoneal dialysis catheters

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

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Peritoneal dialysis

Interventions

Patients who are admitted to Addenbrooke's Hospital for insertion of peritoneal dialysis catheters will be randomised to undergo the procedure by one of the two methods:

1. Percutaneous closed insertion under local anaesthesia in the ward
2. Laparoscopic insertion under general anaesthesia in the main operating theatre

The failure rates and complications of the two procedures will be compared. Both these techniques are normally done in this hospital. The only additional requirement from the patient is consent for randomisation.

Trial stopped due to poor recruitment.

Intervention Type

Procedure/Surgery

Phase

Not Applicable

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

11/09/2003

Reason abandoned (if study stopped)

Participant recruitment issue

Eligibility

Key inclusion criteria

100 patients (PROJ)

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

11/09/2000

Date of final enrolment

11/09/2003

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Box No 210

Cambridge

United Kingdom

CB2 2QQ

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

Cambridge Consortium - Addenbrooke's (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration