

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

<b>Submission date</b> 12/09/2003	<b>Recruitment status</b> Stopped	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 12/09/2003	<b>Overall study status</b> Stopped	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 22/11/2011	<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

**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