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# A single-centre randomised study to assess the need for performing a line change when a peritoneal dialysis patient presents with peritonitis

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| **Submission date**<br>12/09/2003 | **Recruitment status**<br>No longer recruiting | [_] Prospectively registered<br>[_] Protocol<br>[_] Statistical analysis plan<br>[_] Results<br>[_] Individual participant data<br>[_] Record updated in last year |
| **Registration date**<br>12/09/2003 | **Overall study status**<br>Completed | |
| **Last Edited**<br>06/11/2014 | **Condition category**<br>Infections and Infestations | |

**Plain English summary of protocol**
Not provided at time of registration

## Contact information

**Type(s)**
Scientific

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## Additional identifiers

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

### Secondary identifying numbers
N0176079528

# Study information

### Scientific Title

### Study objectives
Is the risk of a recurrent infection reduced by performing a line change on a peritoneal dialysis (PD) patient when patients present with peritonitis?

### 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)
Not specified

### Study type(s)
Not Specified

### Participant information sheet

### Health condition(s) or problem(s) studied
Infections and Infestations: Peritonitis

### Interventions
Antibiotics and no line change compared to antibiotics and a line change

### Intervention Type
Other

### Phase
Not Applicable

### Primary outcome measure

To evaluate whether recurrence of peritonitis is more common in patients who present with peritonitis and receive antibiotics and no line change compared to patients who receive antibiotics and a line change.

### Secondary outcome measures
Not provided at time of registration

### Overall study start date
01/06/2000

### Completion date
01/06/2003

## Eligibility

### Key inclusion criteria
Not provided at time of registration

### Participant type(s)
Patient

### Age group
Not Specified

### Sex
Not Specified

### Target number of participants
40-50 patients and 40-50 control patients, total 100

### Key exclusion criteria
Not provided at time of registration

### Date of first enrolment
01/06/2000

### Date of final enrolment
01/06/2003

## Locations

### Countries of recruitment
England

United Kingdom

### Study participating centre

**Renal Unit**
Oxford
United Kingdom
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# Sponsor information

**Organisation**
Department of Health (UK)

**Sponsor details**
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

**Sponsor type**
Government

**Website**
http://www.doh.gov.uk

# Funder(s)

**Funder type**
Government

**Funder Name**
Oxford Radcliffe Hospitals NHS Trust (UK)

# 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