ISRCTN74962920 https://doi.org/10.1186/ISRCTN74962920

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

Submission date 12/09/2003	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 12/09/2003	**Overall study status** Completed	
Last Edited 06/11/2014	**Condition category** 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
OX3 7LJ

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