

Does furosemide improve renal function in patients with renal impairment?

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 02/11/2016	Condition category Urological and Genital Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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
N0060110641

Study information

Scientific Title
Does furosemide improve renal function in patients with renal impairment?

Study objectives

Does furosemide improve renal function in patients with renal impairment?

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

Urological and Genital Diseases: Renal function

Interventions

Randomised, controlled trial, parallel design.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Furosemide

Primary outcome(s)

Change in creatinine clearance after 24 hours of receiving, or not, furosemide infusion.

Key secondary outcome(s)

Other outcome measures include rate of requirement of renal replacement therapy, peak creatinine, requirement for mineral replacement.

Completion date

31/07/2007

Eligibility**Key inclusion criteria**

100 patients, 2 per week.

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

11/02/2002

Date of final enrolment

31/07/2007

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Chelsea & Westminster Hospital

London

United Kingdom

SW10 9NH

Sponsor information**Organisation**

Department of Health

Funder(s)**Funder type**

Government

Funder Name

Chelsea and Westminster Healthcare NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration