

# Effect of dialystate sodium concentration on intracellular and extracellular fluid volumes in regular chronic haemodialysis (HD) patients

<b>Submission date</b>	<b>Recruitment status</b>	[X] Prospectively registered
28/09/2007	No longer recruiting	<input type="checkbox"/> Protocol
<b>Registration date</b>	<b>Overall study status</b>	<input type="checkbox"/> Statistical analysis plan
28/09/2007	Completed	[X] Results
<b>Last Edited</b>	<b>Condition category</b>	<input type="checkbox"/> Individual participant data
07/09/2012	Urological and Genital Diseases	

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

Dr Andrew Davenport

### Contact details

Department of Nephrology / Transplantation

Royal Free Hampstead NHS Trust

Pond Street

Hampstead

London

United Kingdom

NW3 2QG

+44 020 7794 0500 x 5689 or PA 5457

[andrew.davenport@royalfree.nhs.uk](mailto:andrew.davenport@royalfree.nhs.uk)

## Additional identifiers

### Protocol serial number

N0256183912

## Study information

**Scientific Title****Study objectives**

Does dialysate sodium concentration affect sodium balance during haemodialysis, by altering the amount of sodium available for diffusion?

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Added June 2008: approved by University College Hospital Ethics Committee

**Study design**

Prospective cross over design

**Primary study design**

Interventional

**Study type(s)**

Treatment

**Health condition(s) or problem(s) studied**

Urological and Genital Diseases: Haemodialysis

**Interventions**

Each patient will dialyse against three different dialysate sodium concentrations. The sequence will be randomised.

Until June 2008, trial dates were 01/04/2006 to 01/10/2007.

**Intervention Type**

Drug

**Phase**

Not Specified

**Drug/device/biological/vaccine name(s)**

dialystate sodium

**Primary outcome(s)**

Intracellular and extracellular fluid volume changes, effluent dialysate sodium, interdialytic weight gain, predialysis blood pressure, intradialytic blood pressure, intradialytic blood volume changes.

**Key secondary outcome(s))**

Not provided at time of registration

**Completion date**

31/08/2009

# Eligibility

## Key inclusion criteria

Added June 2008:

1. 18 years or older
2. Stable on dialysis
3. 3 x week HD patients
4. Able to provide informed consent

## Participant type(s)

Patient

## Healthy volunteers allowed

No

## Age group

Not Specified

## Lower age limit

18 years

## Sex

Not Specified

## Key exclusion criteria

Added June 2008:

1. Unable to provide informed consent
2. Unstable on dialysis
3. Dialysis frequency < or > 3 x per week

## Date of first enrolment

01/03/2008

## Date of final enrolment

31/08/2009

# Locations

## Countries of recruitment

United Kingdom

England

## Study participating centre

Department of Nephrology / Transplantation

London

United Kingdom

NW3 2QG

# Sponsor information

## Organisation

Record Provided by the NHSTCT Register - 2007 Update - Department of Health

## Funder(s)

### Funder type

Government

### Funder Name

The Royal Free Hampstead NHS Trust (UK)

### Funder Name

Special Trustees

# Results and Publications

## Individual participant data (IPD) sharing plan

### IPD sharing plan summary

Not provided at time of registration

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	results	01/10/2010		Yes	No
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes