

Myocardial stunning during paediatric dialysis and the effects of cooling the dialystate during haemodialysis

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| Submission date 29/04/2010 | Recruitment status No longer recruiting | <input type="checkbox"/> Prospectively registered |
| Registration date 29/04/2010 | Overall study status Completed | <input type="checkbox"/> Protocol |
| Last Edited 12/04/2017 | 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
5119

Study information

Scientific Title
Myocardial stunning during paediatric dialysis and the effects of cooling the dialystate during haemodialysis: a randomised interventional treatment trial

Study objectives

Original hypotheses:

In comparison to standard 37°C dialysate, dialysate temperature 0.5°C below core body temperature reduces:

1. HD-induced regional LV dysfunction (myocardial stunning)
2. The incidence of IDH and therefore improves the potential for fluid removal during HD (UF)
3. The side effects of IDH and therefore improves the dialysis experience for the child
4. Circulating surrogate markers of myocardial injury

Aim of this pilot:

Primary aim was to determine whether using dialysate 0.5°C cooler than body temperature is associated with a decrease in acute left ventricular (LV) dysfunction as measured by 2D speckle tracking within limits of patient tolerability of feeling cold.

Ethics approval required

Old ethics approval format

Ethics approval(s)

National Research Ethics Service, Central London REC 2, 31/07/2007, ref: 07/Q0508/64

Study design

Randomised interventional treatment trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Topic: Renal and Urogenital; Subtopic: Renal and Urogenital (all Subtopics); Disease: Renal

Interventions

The patient will be randomised to cooled dialysate or dialysate temperature of 37°C.

Follow-up length: 24 months

Study entry: single randomisation only

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Regional LV dysfunction; patients will undergo echocardiogram assessments at 0, 15 and 240 minutes post-dialysis

Key secondary outcome(s)

To determine the differences between cooled and standard dialysate in the:

1. Incidence of hypotensive episodes (IDH) defined as a blood pressure (BP) below the 5th percentile adjusted for age and sex

2. Incidence of self-reported intradialytic and postdialytic symptoms
3. Quality of the dialysis procedure based on a specific questionnaire designed to assess their treatment experience and symptoms related to feeling cold during dialysis
4. UF achieved, calculated as the ratio of actual UF volume and desired UF volume

Completion date

01/07/2011

Eligibility

Key inclusion criteria

1. Children on conventional 4 hours three times/week dialysis
2. Stable on dialysis for 2 months
3. Male and female, lower age limit of 1 month

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Lower age limit

1 months

Sex

All

Key exclusion criteria

Difficult echocardiogram assessments

Date of first enrolment

01/12/2007

Date of final enrolment

01/07/2011

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

30 Guilford Street
London
United Kingdom
WC1N 1EH

Sponsor information

Organisation

Great Ormond Street Hospital for Children (UK)

ROR

<https://ror.org/03zydm450>

Funder(s)

Funder type

Charity

Funder Name

Kids Kidney Research (UK)

Alternative Name(s)

KKR

Funding Body Type

Private sector organisation

Funding Body Subtype

Other non-profit organizations

Location

United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration