ISRCTN19844007 https://doi.org/10.1186/ISRCTN19844007

Use of erythropoietin for anaemia management, clearance of low and middle molecular weight uraemic toxins, quality of life and cost effectiveness of mid-dilution on-line haemodiafiltration compared to conventional low flux haemodialysis

Submission date
29/09/2006

Registration date
29/09/2006

Last Edited
17/05/2017

Recruitment status
No longer recruiting

Overall study status
Completed

Condition category
Haematological Disorders

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[ ] Results
[ ] Individual participant data
[ ] 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
N0205168926

Study information

Scientific Title
Use of erythropoietin for anaemia management, clearance of low and middle molecular weight uraemic toxins, quality of life and cost effectiveness of mid-dilution on-line haemodiafiltration compared to conventional low flux haemodialysis

Study objectives
To quantify the effect of on-line mid-dilution haemodiafiltration on the use of erythropoietin for anaemia management in patients with end-stage renal disease compared to the technique of conventional haemodialysis.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Primary study design
Interventional

Study design
Randomised controlled trial

Study type(s)
Treatment

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

Interventions
Not provided at time of registration

Intervention Type
Drug

Phase
Not Applicable

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

Primary outcome(s)
Control of haemoglobin and use of erythropoietin

Key secondary outcome(s)
Not provided at time of registration

Completion date
31/07/2007

Eligibility

Key inclusion criteria
Not provided at time of registration

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
27/07/2005

Date of final enrolment
31/07/2007

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Royal London Hospital
London
United Kingdom
E1 1BB

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2006 Update - Department of Health (UK)

Funder(s)

Funder type
Government

Funder Name
Barts and The London NHS Trust (UK)

Funder Name
NHS R&D Support Funding (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration