ISRCTN16286916 https://doi.org/10.1186/ISRCTN16286916

Standard versus rapid initiation of anti-hypertensive therapy

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 16/09/2014	**Condition category** Circulatory System	

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
N0237101856

Study information

Scientific Title

Study objectives

The aim of the study is to evaluate the impact on blood pressure control, patient satisfaction and well being and adherence to medication of conventional gradual titration of anti-hypertensive treatment versus rapid introduction and dose escalation of four anti-hypertensive agents in subjects with diabetic nephropathy.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised controlled trial, open design

Primary study design
Interventional

Study type(s)
Treatment

Health condition(s) or problem(s) studied
Cardiovascular: Hypertension

Interventions
Standard versus rapid initiation of anti-hypertensive therapy

Intervention Type
Other

Phase
Not Applicable

Primary outcome(s)
1. Patient adherence to treatment
2. Blood pressure
3. Proteinuria
4. Adverse events
5. Treatment satisfaction

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

Completion date
01/04/2004

Eligibility

Key inclusion criteria
24 subjects recruited from new patients entering the nephropathy clinic

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
01/11/2001

Date of final enrolment
01/04/2004

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Diabetes Centre
Prescot
United Kingdom
L35 5DR

Sponsor information

Organisation
Department of Health (UK)

Funder(s)

Funder type
Government

Funder Name
St Helens and Knowsley Hospitals NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration