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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

Contact name
Dr Shirley Hulme

Contact details
Diabetes Centre
Whiston Hospital
Prescot
United Kingdom
L35 5DR
+44 (0)151 426 1600

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
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

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Treatment

Participant information sheet

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 measure
1. Patient adherence to treatment
2. Blood pressure
3. Proteinuria
4. Adverse events
5. Treatment satisfaction

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/11/2001

Completion date
01/04/2004

Eligibility

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

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
24

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
England

United Kingdom

Study participating centre
Diabetes Centre
Prescot
United Kingdom
L35 5DR

Sponsor information

Organisation
Department of Health (UK)

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

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

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration