

Standard versus rapid initiation of anti-hypertensive therapy

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 16/09/2014	Condition category Circulatory System	<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
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