

A comparison of the effects of Lisinopril and Bendrofluazide on cardiac baroreceptor sensitivity and large artery function following recent ischaemic cerebral stroke

Submission date 30/09/2004	Recruitment status No longer recruiting	☐ Prospectively registered
Registration date 30/09/2004	Overall study status Completed	☐ Protocol
Last Edited 02/05/2018	Condition category Circulatory System	☐ 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

N0123138457

Study information

Scientific Title

A comparison of the effects of Lisinopril and Bendrofluazide on cardiac baroreceptor sensitivity and large artery function following recent ischaemic cerebral stroke

Study objectives

The aim is to investigate the effects on the cardiovascular system of two, commonly-used, blood pressure-lowering treatments following a recent stroke. The objective is to establish the effects of the thiazide diuretic, Bendrofluazide, and the angiotensin Lisinopril

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

Stroke

Interventions

Clinical Study - Human Physiology -Inpatient Admission| - Multicentre Study: National|NHS Patients|Study of Non-Therapeutic Procedure

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Bendrofluazide, lisinopril

Primary outcome(s)

Cardiac baroreceptor sensitivity

Key secondary outcome(s)

Not provided at time of registration

Completion date

05/05/2004

Eligibility

Key inclusion criteria

1. The onset of first ischaemic stroke between 2 and 8 weeks prior to randomisation.
2. A mean brachial blood pressure (BP) level measured on 2 different occasions a week apart between 140/90 mmHg and 180/110 mmHg.
3. The presence of sinus rhythm.
4. Age >18 years

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 Years

Sex

Not Specified

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

27/05/2003

Date of final enrolment

05/05/2004

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

University Hospitals of Leicester

Leicester

United Kingdom

LE1 4PW

Sponsor information

Organisation

Department of Health

Funder(s)**Funder type**

Hospital/treatment centre

Funder Name

University Hospitals of Leicester NHS Trust (UK)

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration