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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

Registration date
30/09/2004

Overall study status
Completed

Last Edited
02/05/2018

Condition category
Circulatory System

[ ] 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

Contact name
Dr David Eveson

Contact details
University Hospitals of Leicester
c/o Research and Development Office
Leicester General Hospital NHS Trust
Leicester
United Kingdom
LE1 4PW
+44 (0)116 258 4109
david.eveson@uhl-tr.nhs.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
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

Study design
Randomised controlled trial

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
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 measure
Cardiac baroreceptor sensitivity

Secondary outcome measures
Not provided at time of registration

Overall study start date
27/05/2003

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

Age group
Adult

Lower age limit
18 Years

Sex
Not Specified

Target number of participants
Not provided at time of registration

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
England

United Kingdom

Study participating centre
University Hospitals of Leicester
Leicester
United Kingdom
LE1 4PW

Sponsor information

Organisation
Department of Health

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

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
University Hospitals of Leicester 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