ISRCTN13275753 https://doi.org/10.1186/ISRCTN13275753

Randomised placebo controlled trial of antihypertensive agents for the prevention of cardiovascular complications of hypertension

Submission date 07/09/2005	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 08/09/2005	**Overall study status** Completed	
Last Edited 19/05/2011	**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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

Study information

Scientific Title

Study objectives
That treatment with diuretic or beta-blocker in hypertensive people aged between 65 and 74 reduces risk of stroke, coronary heart disease and death.

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
Hypertension

Interventions
Amiloride, hydrochlorothiazide compared with placebo; atenolol compared with placebo.

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Amiloride, hydrochlorothiazide and atenolol

Primary outcome measure
Stroke, coronary events, deaths from all causes.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/03/1982

Completion date
01/03/1987

Eligibility

Key inclusion criteria
Men and women aged 65-74 at recruitment in 226 general practices in the Medical Research Council's General Practice Research Framework. Systolic blood pressure 160-209 mmHg and diastolic pressure less than 115 mmHg.

Participant type(s)
Patient

Age group
Senior

Sex
Both

Target number of participants
4396 participants

Key exclusion criteria
Secondary hypertension; already on antihypertensive treatment; accepted indication for antihypertensive treatment; myocardial infarction or stroke within previous three months; impaired renal function; presence of angina, intermittent claudication, diabetes, gout, bronchial asthma, serious intercurrent disease including malignant hypertension.

Date of first enrolment
01/03/1982

Date of final enrolment
01/03/1987

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Non-communicable Disease Epidemiology Unit
London
United Kingdom
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Sponsor information

Organisation
Sponsor not defined - Record provided by the Medical Research Council (UK)

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Sponsor type
Not defined

Funder(s)

Funder type
Research council

Funder Name
Medical Research Council (UK)

Alternative Name(s)
Medical Research Council (United Kingdom), UK Medical Research Council, MRC

Funding Body Type
Government organisation

Funding Body Subtype
National government

Location
United Kingdom

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	13/07/1985		Yes	No