

Cost-effectiveness of low salt diet in lowering blood pressure in Hong Kong Chinese

Submission date 14/10/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 14/10/2002	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 01/07/2009	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

Contact name
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Contact details
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
811028

Study information

Scientific Title

Study objectives

Not provided at time of registration

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)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

High blood pressure

Interventions

All patients will undergo a 4-week placebo run-in and drug washout phase. After baseline measurements, each patient will be randomised to conventional treatment or low sodium diet for 6 months.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/10/1998

Completion date

01/09/2000

Eligibility

Key inclusion criteria

1. Age between 18 and 75 inclusive
2. Patient has seated diastolic blood pressure between 90 and 110 mmHg inclusive; or seated systolic blood pressure of greater than 140 mmHg
3. Patient must be willing to give full informed consent

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/10/1998

Date of final enrolment

01/09/2000

Locations

Countries of recruitment

Hong Kong

Study participating centre

Department of Medicine

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Hong Kong

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Sponsor information

Organisation

Hong Kong Health Services Research Fund (Hong Kong)

Sponsor details

Health Welfare and Food Bureau
Government Secretariat, HKSAR
20th floor Murray Building
Garden Road

-

Hong Kong

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+852 (0)2973 8288
hsrf@hwfb.gov.hk

Sponsor type

Government

Website

http://www.fhb.gov.hk/grants/english/funds/funds_hhsrf/funds_hhsrf_abt/funds_hhsrf_abt.html

ROR

<https://ror.org/03qh32912>

Funder(s)

Funder type

Government

Funder Name

Hong Kong Health Services Research Fund (Hong Kong)

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