

Evaluation of World Health Organisation (WHO) cardiovascular disease (CVD)-risk management package - scenario one (Sri Lanka)

Submission date 06/04/2005	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 07/06/2005	Overall study status Completed	☐ Statistical analysis plan ☐ Results
Last Edited 26/03/2008	Condition category Circulatory System	☐ 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 S Mendis

Contact details
20, Avenue Appia
Geneva-27
Switzerland
CH 1211
mendiss@who.int

Additional identifiers

Protocol serial number
RPC050

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

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Hypertension

Interventions

20 primary health care facilities.

10 facilities to apply the study protocol based on Scenario One of the WHO CVD-Risk Management Package (intervention sites).

Other 10 to continue with conventional treatment or usual care (control sites).

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

07/05/2005

Eligibility

Key inclusion criteria

1. Age 30 - 70 years
2. Currently not under treatment for hypertension
3. Systolic blood pressure between 140 and 179 mmHg measured twice with a 5 to 10 minute interval
4. Informed consent given

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Pregnancy
2. Trauma (injury) as presenting complaint
3. Renal diseases: nephropathy, renal artery stenosis
4. Endocrinological (hormonal) disorders: phaeochromocytoma, Cushing syndrome, Conn syndrome, acromegaly
5. Coarctation of the aorta
6. Use of steroids and/or non-steroidal anti-inflammatories (NSAIs)
7. All acute conditions presenting with bleeding, pain, diarrhoea, vomiting, breathing disorders and circulatory disorders
8. Inability to comply with the follow-up requirements
9. Inability to provide informed consent

Date of first enrolment

07/05/2004

Date of final enrolment

07/05/2005

Locations**Countries of recruitment**

Sri Lanka

Switzerland

Study participating centre

20, Avenue Appia

Geneva-27

Switzerland

CH 1211

Sponsor information

Organisation

The Department of Cardiovascular Diseases (CVD)/World Health Organization (WHO)
(Switzerland)

ROR

<https://ror.org/01f80g185>

Funder(s)**Funder type**

Research organisation

Funder Name

The Department of Cardiovascular Diseases (CVD)/World Health Organization (WHO)
(Switzerland)

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

Not provided at time of registration