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

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

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

#### Study design

Randomised controlled trial

#### Primary study design

Interventional

#### 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: phaechromocytoma, 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