

# Comparison of Non-invasive Finger Blood Pressure and Accutorr oscillometer Blood Pressure

Published: 25-11-2008

Last updated: 06-05-2024

The study should determine whether non-invasive continuous blood pressure values are comparable to intermittent oscillometric blood pressure determinations.

<b>Ethical review</b>	Approved WMO
<b>Status</b>	Pending
<b>Health condition type</b>	Central nervous system vascular disorders
<b>Study type</b>	Observational non invasive

## Summary

### ID

NL-OMON32670

### Source

ToetsingOnline

### Brief title

Non-invasive Finger Blood Pressure and Accutorr

### Condition

- Central nervous system vascular disorders

### Synonym

faint, syncope

### Research involving

Human

### Sponsors and support

**Primary sponsor:** Academisch Medisch Centrum

**Source(s) of monetary or material Support:** Ministerie van OC&W

## Intervention

**Keyword:** Blood pressure, evaluation, non-invasive

## Outcome measures

### Primary outcome

Oscillometrically determined blood pressure.

### Secondary outcome

Not applicable.

## Study description

### Background summary

Continuous blood pressure monitoring is important in syncope evaluation.

### Study objective

The study should determine whether non-invasive continuous blood pressure values are comparable to intermittent oscillometric blood pressure determinations.

### Study design

In consecutive patients of the Syncope-Unit, non-invasive continuous blood pressure values (Nexfin, BMEYE, Amsterdam) will be compared to oscillometric blood pressure values.

### Study burden and risks

None.

## Contacts

### Public

Academisch Medisch Centrum

Postbus 22660  
1100 DD Amsterdam  
Nederland  
**Scientific**  
Academisch Medisch Centrum

Postbus 22660  
1100 DD Amsterdam  
Nederland

## Trial sites

### Listed location countries

Netherlands

## Eligibility criteria

### Age

Adults (18-64 years)

Elderly (65 years and older)

### Inclusion criteria

Patients visiting the Syncope Unit older than 18 years of age.

### Exclusion criteria

Age under 18 years old.

## Study design

### Design

**Study type:** Observational non invasive

Masking: Open (masking not used)

Control: Uncontrolled

Primary purpose: Diagnostic

## Recruitment

NL  
Recruitment status: Pending  
Start date (anticipated): 01-11-2008  
Enrollment: 35  
Type: Anticipated

## Ethics review

Approved WMO  
Application type: First submission  
Review commission: METC Amsterdam UMC

## Study registrations

### Followed up by the following (possibly more current) registration

No registrations found.

### Other (possibly less up-to-date) registrations in this register

No registrations found.

### In other registers

Register	ID
CCMO	NL24969.018.08