

# Non-invasive Assessment of Coronary Artery Disease in Patients with Chest Pain.

No registrations found.

<b>Ethical review</b>	Positive opinion
<b>Status</b>	Recruitment stopped
<b>Health condition type</b>	-
<b>Study type</b>	Interventional

## Summary

### ID

NL-OMON24749

### Source

NTR

### Brief title

N/A

### Health condition

Patients with suspected coronary artery disease

## Sponsors and support

**Source(s) of monetary or material Support:** Netherlands Heart Foundation

## Intervention

## Outcome measures

### Primary outcome

MSCT may improve (as compared to MPI) the diagnosis of patients presenting with chest pain complaints and an intermediate likelihood of CAD.

Particularly in these patients, a non-invasive test with a high specificity (to exclude CAD) is

needed to allow optimal management of patients. Currently, MPI is used for this purpose, but the specificity of MPI is suboptimal (70%).

### **Secondary outcome**

N/A

## **Study description**

### **Background summary**

N/A

### **Study objective**

In patients presenting with chest pain complaints and an intermediate risk of coronary artery disease, Multi-Slice Computed Tomography (MSCT) will have a higher specificity as compared to myocardial perfusion imaging (MPI). Accordingly, MSCT may serve as an accurate first-line evaluation tool.

### **Study design**

N/A

### **Intervention**

MSCT coronary angiography in addition to myocardial perfusion imaging (MPI).

## **Contacts**

### **Public**

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## Eligibility criteria

### Inclusion criteria

Adult patients (having obtained legal majority age) with chest pain complaints and an intermediate pre-test likelihood of CAD (based on the Diamond and Forrester method) with the need for additional imaging studies to evaluate the presence/absence of CAD.

### Exclusion criteria

Fertile women, patients with severe renal failure, patients presenting with a known allergy to iodine contrast media, patients included in another clinical trial, patients under guardianship, and patients whose degree of cooperation is incompatible with carrying out the study.

## Study design

### Design

Study type:	Interventional
Intervention model:	Other
Allocation:	Non controlled trial
Masking:	Open (masking not used)
Control:	N/A , unknown

### Recruitment

NL	
Recruitment status:	Recruitment stopped
Start date (anticipated):	01-10-2004

Enrollment: 100  
Type: Actual

## Ethics review

Positive opinion  
Date: 15-09-2005  
Application type: First submission

## 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
NTR-new	NL446
NTR-old	NTR486
Other	: N/A
ISRCTN	ISRCTN65675235

## Study results

### Summary results

N/A