

# The effects of doxycycline treatment on inflammation and endothelial function in advanced atherosclerosis

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-OMON23404

### Source

Nationaal Trial Register

### Brief title

N/A

### Health condition

atherosclerosis, peripheral arterial disease, inflammation, Doxycycline

## Sponsors and support

**Primary sponsor:** none

**Source(s) of monetary or material Support:** none

## Intervention

## Outcome measures

### Primary outcome

Endothelial function

### Secondary outcome

## Study description

### Background summary

Patients with mild to moderate peripheral arterial disease were treated with doxycycline 100mg daily or placebo.

Reduction in vascular inflammation was assessed by measuring vascular endothelial function and peripheral plasma markers of inflammation.

### Study objective

Doxycycline will ameliorate the vascular condition and inflammatory status in patients with mild to moderate atherosclerotic disease

### Study design

t=0, follow up after 30, 60, 90 and 120 days

### Intervention

Patients were (if necessary) pre-treated with or switched to simvastatin 40 mg daily at least four weeks prior to trial start.

Patients were treated with 100 mg doxycycline for four weeks

## Contacts

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## Eligibility criteria

### Inclusion criteria

1. Patients suffering from mild to moderate stable peripheral arterial disease who do not require carotis or femoral endarterectomy (Fontaine IIa)

### Exclusion criteria

1. Known hypersensitivity for tetracycline derivatives
2. Patients treated with antibiotics in last 3 months for other diseases then lower urinary tract infections
3. Patients treated with immunosuppressive agents (including steroids)
4. Patients treated with fibrates
5. Patients with Diabetes Mellitus regulated with drug therapy
6. Signs of kidney or liver failure

## Study design

### Design

Study type:	Interventional
Intervention model:	Crossover
Allocation:	Randomized controlled trial
Masking:	Double blinded (masking used)
Control:	Placebo

### Recruitment

NL

Recruitment status:	Recruitment stopped
Start date (anticipated):	01-01-2006
Enrollment:	15
Type:	Actual

## Ethics review

Positive opinion	
Date:	23-07-2008
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	NL1331
NTR-old	NTR1389
Other	METC LUMC : P05.189
ISRCTN	ISRCTN wordt niet meer aangevraagd

## Study results

### Summary results

In progress