

Activity, Lifestyle And Nutrition and Therapy study

Submission date 04/08/2005	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 04/08/2005	Overall study status Completed	☐ Statistical analysis plan ☒ Results
Last Edited 04/02/2013	Condition category Circulatory System	☐ Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Helma IJzelenberg

Contact details
De Boelelaan 1085
Amsterdam
Netherlands
1081 HV
+31 (0)20 598 9951
helma.ijzelenberg@falw.vu.nl

Additional identifiers

Protocol serial number
NTR42

Study information

Scientific Title

Acronym
ALANT

Study objectives

To evaluate cost-effectiveness of a lifestyle intervention in patients with cardiovascular disease.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics approval received from the local ethics committee

Primary study design

Interventional

Study design

Randomised, active controlled, parallel group trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Cardiovascular disease

Interventions

Patients of the VU University Medical Centre will be randomised into two groups. One group will receive a six-month lifestyle intervention which includes a physical activity program and when appropriate smoking cessation and weight loss. The other group will continue to receive usual care.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Cost-effectiveness of a lifestyle intervention in patients with cardiovascular disease

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/03/2008

Eligibility**Key inclusion criteria**

1. The patient has manifest cardiovascular suffering in combination with at least one lifestyle related risk factor (physical inactivity, smokes, obesity)
2. The patient must be able to participate in a training program
3. The patients' insurance company will cover the costs of the intervention program

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. A recent cardiovascular incident (within less than 3 months)
2. Unstable (co)-morbidity
3. A planned operation or undergoing investigations which can directly lead to an operation
4. Younger than 18 years
5. The patient has already attended the intervention program
6. Insufficient knowledge of the Dutch language

Date of first enrolment

01/04/2005

Date of final enrolment

01/03/2008

Locations**Countries of recruitment**

Netherlands

Study participating centre

De Boelelaan 1085

Amsterdam

Netherlands

1081 HV

Sponsor information**Organisation**

Vrije University Medical Centre (VUMC) (The Netherlands)

ROR

<https://ror.org/00q6h8f30>

Funder(s)

Funder type

Research organisation

Funder Name

The Netherlands Organisation for Health Research and Development (ZonMw) (The Netherlands)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	10/09/2012		Yes	No