Evaluation of Vascular care in Alzheimer's disease.

No registrations found.

Ethical review	Positive opinion
Status	Recruitment stopped
Health condition type	-
Study type	Interventional

Summary

ID

NL-OMON22112

Source
NTR

Brief title
EVA

Health condition

Alzheimer's disease

Sponsors and support

Primary sponsor: ZONMw Doelmatigheidsonderzoek
Source(s) of monetary or material Support: ZONMw Doelmatigheidsonderzoek

Intervention

Outcome measures

Primary outcome

Change in impairments after two years of follow up in activities of daily living as measured by the Interview for Deterioration in Daily life in dementia (IDDD).

Secondary outcome

Change in cognitive functioning, behavioral abnormalities, blood pessure, body weight. New lesions on MRI after two years.

Study description

Background summary

Frequently patients with a clinical diagnosis of Alzheimer's disease also harbour cerebrovascular lesions on neuroimaging. It is not clear to what extent these lesions contribute to (1) functional impairments, and (2) the prognosis of these patients. In this open multicentre clinical trial the hypothesis is tested that intensive vascular care is efficacious in these patients, resulting in slower deterioration after a follow up of two yeras in patients receiving such care, in comparison to patients receiving standard care.

Study objective

Intensive vascular care, aimed at secundary prevention of cerebrovascular disease, is efficacious in patients with Alzheimer's disease that have vascular lesions on MRI.

Study design

N/A

Intervention

Multicomponent intervention consisting of a combination of measures aimed at prevention of cerebrovascular disease (platelet aggregation inhibitors, statin, strict regulation of blood pressure, diabetes, life style interventions concerning smoking, body weight, excercise). Patients in the control group will recieve 'regular care' i.e. less frequent visits, without specific attention to vascular risk factors.

Contacts

Public
Academic Medical Center (AMC), Department of Neurology (H2-217),
P.O. Box 22660
W.A. Gool, van
Meibergdreef 9
Amsterdam 1100 DD
The Netherlands
+31 (0)20 5664044

Scientific
Academic Medical Center (AMC), Department of Neurology (H2-217),
P.O. Box 22660
W.A. Gool, van
Meibergdreef 9
Amsterdam 1100 DD
The Netherlands
+31 (0)20 5664044

Eligibility criteria

Inclusion criteria

Patients with early Alzheimer's disease according to clinical criteria, that have on MRI either cerebral infarcts or significant white matter abnormalities.

Exclusion criteria

1. Severe dementia;

2. Limited life expectancy.

Study design

Design

Study type:	Interventional
Intervention model:	Parallel
Allocation:	Randomized controlled trial
Masking:	Open (masking not used)
Control:	Active

Recruitment

NL

Recruitment status:	Recruitment stopped
Start date (anticipated):	01-06-2002

Enrollment:	130
Type:	Actual

Ethics review

Positive opinion Date:	30-03-2006
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	NL590
NTR-old	NTR646
Other	: N/A
ISRCTN	ISRCTN74902933

Study results

Summary results
N/A