ISRCTN57453322 https://doi.org/10.1186/ISRCTN57453322

Long-term prognosis following cerebrovascular events of different etiologies

Submission date 14/04/2006	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 13/06/2006	**Overall study status** Completed	
Last Edited 13/06/2006	**Condition category** Circulatory System	

Plain English summary of protocol
Not provided at time of registration

Study website
http://www.uni-essen.de/neurologie/stroke/

Contact information

Type(s)
Scientific

Contact name
Dr Christian Weimar

Contact details
Hufelandstr. 55
Essen
Germany
45122
+49 (0)201 723 2495
stroke.med@uni-essen.de

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
DI 327/9-1

Study information

Scientific Title

Study objectives
To determine the rate of recurrence in patients with cerebrovascular events of defined etiology

Ethics approval required
Old ethics approval format

Ethics approval(s)
Ethics Committee of the University of Essen reviewed the protocol of this trial and confirmed that ethical approval was not required.

Study design
Prospective observational cohort study

Primary study design
Interventional

Secondary study design
Cohort study

Study setting(s)
Not specified

Study type(s)
Diagnostic

Participant information sheet

Health condition(s) or problem(s) studied
Cerebral stroke and TIA

Interventions
Patients are followed-up via telephone interview to determine the long-term rate of recurrence in patients with predefined stroke etiologies

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Recurrent stroke

Secondary outcome measures

Mortality of any cause

Overall study start date
01/09/2002

Completion date
31/12/2009

Eligibility

Key inclusion criteria
Patients with acute cerebrovascular events of defined etiology including:
1. Cardiac right-to-left shunt
2. Transient ischemic attack (TIA) with symptoms lasting <1 hour
3. Intracerebral bleeding
4. Intracranial stenosis
5. Dissection of brain supplying arteries
6. Coagulation disorder
7. Previous history of migraine
8. Basilar artery occlusion
9. Vasculitis
10. Arteriovenous malformation
11. Moyamoya syndrome

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
3000

Key exclusion criteria
Inability or refusal to provide consent for study participation

Date of first enrolment
01/09/2002

Date of final enrolment
31/12/2009

Locations

Countries of recruitment
Germany

Study participating centre
Hufelandstr. 55
Essen
Germany
45122

Sponsor information

Organisation
German Research Foundation (Deutsche Forschungsgemeinsschaft) (DFG)

Sponsor details
Kennedeyallee 40
Bonn
Germany
53175
theodora.hogenkamp@dfg.de

Sponsor type
Research organisation

Website
http://www.dfg.de

ROR
https://ror.org/018mejw64

Funder(s)

Funder type
Research organisation

Funder Name
German Research Foundation (Deutsche Forschungsgemeinsschaft) (DFG) DI 327/9-1

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration