Long-term prognosis following cerebrovascular events of different etiologies

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

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

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Additional identifiers

Protocol serial number 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

Study type(s)

Diagnostic

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(s)

Recurrent stroke

Key secondary outcome(s))

Mortality of any cause

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

Healthy volunteers allowed

No

Age group

Adult

Sex

All

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)

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

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?
Study website	Study website	11/11/2025	11/11/2025	No	Yes