

Dynamics of language-related neuroplasticity in stroke: a longitudinal study

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The present study examines stroke patients* language abilities as well as the recruitment of additional brain areas in language use over time to compensate for stroke-induced language deficits.

Ethical review	Approved WMO
Status	Recruiting
Health condition type	Other condition
Study type	Observational non invasive

Summary

ID

NL-OMON49365

Source

ToetsingOnline

Brief title

LangPlast

Condition

- Other condition
- Structural brain disorders

Synonym

CVA, stroke

Health condition

aphasia and language disorders through brain damage

Research involving

Human

Sponsors and support

Primary sponsor: Radboud Universiteit Nijmegen

Source(s) of monetary or material Support: Ministerie van OC&W

Intervention

Keyword: aphasia, Plasticity, stroke

Outcome measures

Primary outcome

Main study parameter is the task-related brain activity measured with electroencephalography and how it differs over time, together with patients* functional communication and memory performance for the included behavioural tasks.

Secondary outcome

To assess the effort that patients often experience in language use after stroke lesions, measured by tracking the heart rate and skin conduction due to sweat responses (GSR).

Study description

Background summary

Suffering from a stroke in the left hemisphere very often impairs patients* language functions. Recent research suggests that, during recovery, additional brain areas might be recruited to overcome deficits induced by the lesion. However, the underlying processes of this neural recruitment and reorganization of language functions in the brain are still largely unknown. The present study will investigate the language-related recruitment and reorganization in the brain at different time points after stroke. We will employ electroencephalography combined with various language and memory tasks to gain more insight into brain areas that are recruited for language functioning after stroke lesions, and its effect on behavioural language performance.

Study objective

The present study examines stroke patients* language abilities as well as the recruitment of additional brain areas in language use over time to compensate for stroke-induced language deficits.

Study design

Longitudinal study at two time points to assess patients* memory and communication abilities and language-related brain activity at approximately two and seven months after stroke onset.

Study burden and risks

The current procedures of electroencephalography and magnetic resonance imaging and experimental measures implicate minimal burden and insignificant risk to participants. Patients do not directly benefit from their participation in terms of therapeutic gain, but they do obtain extensive information about their language and communicative abilities and change over time, which they appreciate. Participants can withdraw from the study at any time point. The findings of this study will yield further insights into the reorganization of the language network after stroke and possibly aid to create more individualized patient therapy and care.

Contacts

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Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adults (18-64 years)

Elderly (65 years and older)

Inclusion criteria

- Between 18 - 78 years of age
- Ischemic or haemorrhagic cerebral infarction
- Postonset of approx. 2 months (subacute stage) for session 1, and 7 months (chronic stage) for session 2
- Native speaker of Dutch
- Pre-morbidly right handed
- Willingness and ability to give written informed consent and willingness and ability to understand the nature and content, to participate and to comply with the study requirements.

Exclusion criteria

- History of previous stroke
- Pre-existent cognitive problems
- Severe aphasia (unable to understand instructions or provide consent). This will be predetermined by the referring neurologist or speech language pathologist, according to their usual practice
- Use of psychotropic medication or recreational drugs
- Pregnancy
- Serious head trauma
- Neurological or psychiatric disorders (other than stroke or epilepsy as a consequence of the stroke)
- Large or ferromagnetic metal parts in the head (except for a dental wire)
- Implanted cardiac pacemaker or neurostimulator
- Claustrophobia

Study design

Design

Study type:	Observational non invasive
Intervention model:	Other
Allocation:	Non-randomized controlled trial
Masking:	Open (masking not used)
Control:	Active
Primary purpose:	Other

Recruitment

NL	
Recruitment status:	Recruiting
Start date (anticipated):	08-02-2021
Enrollment:	75
Type:	Actual

Ethics review

Approved WMO	
Date:	18-05-2020
Application type:	First submission
Review commission:	CMO regio Arnhem-Nijmegen (Nijmegen)

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
CCMO	NL72600.091.20