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Communicative diagnostics and communicative strategies of people with aphasia and their dependents

Submission date	Recruitment status	[ ] Prospectively registered
14/06/2007	Stopped	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
16/07/2007	Stopped	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
02/09/2009	Signs and Symptoms	[ ] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Study website
http://www.refonet.de

Contact information

Type(s)
Scientific

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56588

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
05002

Study information

Scientific Title

Acronym
KAPUA

Study objectives
1. The communication training of the dependents improves the communicative capacity of the people with aphasia
2. The improvement of the communicative capacity also occurs for people with aphasia, who show no improvement in the symptoms
3. The improvement of the communicative capacity is more intense for people with aphasia who also show an improvement in the symptoms than for affected persons who show no change in the symptoms

Ethics approval required
Old ethics approval format

Ethics approval(s)
Approval received from the local ethics committee (Landesärztekammer Rheinland-Pfalz, Mainz) on the 30th October 2006.

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Aphasia

Interventions
Please note that as of 02/09/09 the status of this record was changed to 'stopped' due to recruitment problems. The trial officially stopped on 15/04/2008.

1. Communication training for people with aphasia and their dependents; training twice for two days within two months and homework

2. Tests (Aachen Aphasia Test [AAT], Aphasia Check List [ACL], Communicative Effectiveness Index [CETI], KAPUA [communication test for people with aphasia and their dependents]) to compare at the beginning, two months later and after eight months from the start.

Control:
Control group without training. Only tests to compare at the beginning, two months later and eight months from the start.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
To prove the improvement of communicative capacity of people with aphasia in special consideration of the communicative part of the dependents.

Secondary outcome measures
To establish an approved and evaluated training program for dependents of people with aphasia.

Overall study start date
01/03/2007

Completion date
01/05/2009

Reason abandoned (if study stopped)
Participant recruitment issue

Eligibility

Key inclusion criteria
1. German native speaker
2. Aphasia since at least six months
3. Live together with the dependant in the same household

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
120 people with aphasia with one dependant each

Key exclusion criteria

1. Severe apraxia of speech
2. Severe dysarthria
3. Severe dementia

Date of first enrolment
01/03/2007

Date of final enrolment
01/05/2009

Locations

Countries of recruitment
Germany

Study participating centre
Buchenstrasse 6
Waldbreitbach
Germany
56588

Sponsor information

Organisation
Refonet (Germany)

Sponsor details
Postfach 10 07 63
Bad Neuenahr-Ahrweiler
Germany
53445

Sponsor type
Industry

Website
http://www.refonet.de

ROR
https://ror.org/04yeh2x21

Funder(s)

Funder type
Industry

Funder Name
Refonet (Germany)

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