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Neuropsychological rehabilitation: Modular cognitive retraining versus compensatory skills training

Submission date	Recruitment status	[ ] Prospectively registered
01/08/2003	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
27/10/2003	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
14/08/2009	Injury, Occupational Diseases, Poisoning	[ ] 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

B8

Study information

Scientific Title

Study objectives
A randomised controlled trial on neuropsychological rehabilitation

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Other

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Severe head trauma, stroke

Interventions
Compensatory skills training versus retraining basic cognitive impairments (either memory skills training or problemsolving)
addititional for both therapies
"living with neuropsychological impairment" and attention training

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Not provided at time of registration.

Secondary outcome measures
Not provided at time of registration.

Overall study start date
01/04/2002

Completion date
30/04/2005

Eligibility

Key inclusion criteria
1. Neuropsychological impairments in at least two of the cognitive domains (cf. International Classification of Functioning, Disability and Health [ICF], World Health Organisation [WHO]): attention, memory, or problem solving
2. Age 16 to 55 years
3. Participation in the therapy programme for 5 weeks
4. Barthel Index at least summary score 40

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
Not provided at time of registration.

Key exclusion criteria
1. Prior brain injury to the actual trauma
2. Psychiatric disorder before trauma medically documented
3. Substance abuse
The participants are all inpatients at the neuropsychological rehabilitation centres. They are no longer in emergency units but in therapeutic care for the reintegration in job, family and daily life.

Date of first enrolment
01/04/2002

Date of final enrolment
30/04/2005

Locations

Countries of recruitment
Germany

Study participating centre
Medical Director
Schaufling
Germany
94571

Sponsor information

Organisation
Bavarian Network for Rehabilitation Research (Rehabilitationswissenschaftlicher Forschungsverbund Bayern) (Germany)

Sponsor details
RFB-Geschäftsstelle
Wilmar Igl
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Sponsor type
Research organisation

Funder(s)

Funder type
Research organisation

Funder Name
Bavarian Network for Rehabilitation Research (Rehabilitationswissenschaftlicher Forschungsverbund Bayern) (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