

Neuropsychological rehabilitation: Modular cognitive retraining versus compensatory skills training

Submission date 01/08/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 27/10/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 14/08/2009	Condition category Injury, Occupational Diseases, Poisoning	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Peter Frommelt

Contact details
Medical Director
Asklepios Klinik Schaufling
Haustein 2
Schaufling
Germany
94571
+49 (0)9904 775571
p.frommelt@asklepios.com

Additional identifiers

Protocol serial number
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

Study type(s)

Treatment

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)
additional for both therapies
"living with neuropsychological impairment" and attention training

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Not provided at time of registration.

Key secondary outcome(s)

Not provided at time of registration.

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

Healthy volunteers allowed

No

Age group

Adult

Sex

All

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)

Funder(s)

Funder type

Research organisation

Funder Name

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

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration