The N-methyl-D-aspartate (NMDA)-antagonist memantine affects training induced motor cortex plasticity: a study using transcranial magnetic stimulation

Submission date	Recruitment status No longer recruiting	Prospectively registered		
06/05/2005		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
09/05/2005	Completed	[X] Results		
Last Edited	Condition category	[] Individual participant data		
15/02/2008	Other			

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

Study objectives

Training of a repetitive synchronised movement of two limb muscles leads to short-term plastic changes in the primary motor cortex, which can be assessed by transcranial magnetic stimulation (TMS) mapping.

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)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Training induced motor cortex plasticity

Interventions

Memantine versus placebo either given as a single dose or daily over 8 days.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Memantine

Primary outcome measure

The differential effects of different treatment regimens.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/06/1998

Completion date

31/01/2005

Eligibility

Key inclusion criteria

Healthy subjects (adults, either sex)

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

20

Key exclusion criteria

- 1. Left-handedness
- 2. Neurological disorders
- 3. Intake of central acting drugs

Date of first enrolment

01/06/1998

Date of final enrolment

31/01/2005

Locations

Countries of recruitment

Germany

Study participating centre Department of Neurology

Bochum

Sponsor information

Organisation

BG-Kliniken Bergmannsheil (Germany) - Department of Neurology

Sponsor details

Buerkle-de-la-Camp-Platz 1 Bochum Germany 44789

Sponsor type

Hospital/treatment centre

ROR

https://ror.org/04j9bvy88

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

BG-Kliniken Bergmannsheil (Germany) - Department of Neurology; main funders

Funder Name

Merz Pharmaceuticals GmbH (Germany) - funded the measurement of the memantine serum levels

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 summaryNot provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results	12/05/2005		Yes	No