

Left prefrontal activity and the voluntary control of social emotional behaviour

Published: 12-01-2010

Last updated: 04-05-2024

The aim of this study is to investigate whether the vIPFC is crucially involved in the voluntary control of social emotional behaviour. Presumably, the congruency effect will be larger (i.e. higher reaction times in the incongruent condition) on the...

Ethical review	Approved WMO
Status	Recruitment stopped
Health condition type	Other condition
Study type	Interventional

Summary

ID

NL-OMON35035

Source

ToetsingOnline

Brief title

The PFC and the control of social motivational behaviour

Condition

- Other condition

Synonym

not applicable

Health condition

niet van toepassing

Research involving

Human

Sponsors and support

Primary sponsor: Radboud Universiteit Nijmegen

Source(s) of monetary or material Support: VIDI-subsidie van Dr. K. Roelofs

Intervention

Keyword: Approach-avoidance, Left prefrontal cortex, Social emotional behaviour, Theta-burst stimulation

Outcome measures

Primary outcome

Reaction times and error rates on the approach-avoidance task will be used to test whether the congruency effect in the cTBS condition applied over the vlPFC is larger than in the other TMS conditions.

Secondary outcome

Aarterial spin labelling during task performance is used to investigate the effects of the different TMS conditions on cerebral blood flow. Moreover, hormonal levels of cortisol and testosterone are measured in order to control for differences in performance on the approach-avoidance-task between subjects.

Study description

Background summary

The left ventrolateral prefrontal cortex (vlPFC) is found to be involved in the control of social emotional behaviour. This social emotional behaviour could be quantified by means of the approach-avoidance (AA) task, in which happy or angry facial expressions should be either approached or avoided by pulling or pushing a joystick, respectively. It is found that people are more likely to approach positive and to avoid negative facial expressions (affect-congruent condition). In the affect-incongruent condition (i.e. avoid positive and approach negative faces), on the other hand, reaction times are found to be longer because the automatic emotional tendencies have to be suppressed (the so-called congruency effect). The aim of this project is to investigate whether

activity of the vIPFC is crucial for the control of social emotional behaviour. In order to infer a causal relationship, Transcranial Magnetic Stimulation (TMS) will be applied over the vIPFC. We hypothesize that the congruency effect will increase after inhibition of the left vIPFC.

Study objective

The aim of this study is to investigate whether the vIPFC is crucially involved in the voluntary control of social emotional behaviour. Presumably, the congruency effect will be larger (i.e. higher reaction times in the incongruent condition) on the approach-avoidance task after inhibition of the left vIPFC by TMS. This finding would indicate a crucial role for the vIPFC in overruling automatic emotional tendencies.

In previous studies, endogenous testosterone and cortisol were found to modulate social emotional behaviour. Therefore, hormonal measures will be taken into account. We expect to find a positive correlation between cortisol and the congruency effect and a negative correlation between testosterone and the congruency effect.

Furthermore, to investigate (and control for) the influence of TMS on cerebral blood flow, Arterial Spin Labeling (ASL) will be performed. By means of this technique, it is possible to magnetically label the blood supply to the region of interest (PFC), which gives an indication of the cerebral blood flow in the specific region. We expect to measure less cerebral blood flow in the left vIPFC after inhibition with TMS.

Study design

The study will consist of four sessions, each separated by approximately a week. During the first session, the motor threshold (MT) will be determined (to determine the stimulation intensity in the later TMS-sessions) and a structural MRI-scan will be made (in order to localize the TMS coil on the left vIPFC in the other TMS-sessions). Furthermore, during this first session, a short cTBS protocol (20 sec.) will be applied on the vIPFC at a low intensity (80% of aMT). By this means, subjects will be allowed to indicate whether they would like to proceed with the experiment. In the other three sessions, an off-line TMS-protocol will be used in which Theta Burst Stimulation (TBS) or repetitive TMS (5 Hz.) will be administered. In order to control for the effects of TMS on the cerebral blood flow, subjects will perform the AA-task during the measurement of ASL. Moreover, because endogenous testosterone and cortisol were found to influence approach-avoidance behaviour, these hormones will also be taken into account.

Intervention

All participants will receive two different theta-burst stimulations of 600 pulses and one repetitive TMS (5Hz.) of 200 pulses at 80% of active motor

threshold on three different sessions:

1. Continuous TBS (inhibition) over the vIPFC (experimental intervention).
2. Repetitive TMS (5Hz.) over the vIPFC (to control for stimulation).
3. Continuous TBS over the vertex (to control for site).

All TMS protocols will take 40 sec.

Study burden and risks

TMS is not painful at the level of intensity used in this project (i.e. 80% of active motor threshold). From the previous literature we know that, in rare cases, subjects could report a (light) headache, which could be treated easily with paracetamol. On the basis of incidental epileptic seizures triggered by TMS in early 90*s, safety-guidelines were established to set up the maximum duration of TMS stimulation (Wassermann, 1998). Therefore, our protocols will follow these safety TMS guidelines. Furthermore, all subjects will be pre-screened for relevant medical history, drug abuse, head trauma, neurological or psychiatric illness, pregnancy, heart disease, cardiac pacemakers, medication pumps, tricyclic antidepressants, neuroleptics, family history of neurological illness, psychiatric illness or epilepsy. Because the risk associated with participation can be considered negligible and the burden can be considered minimal, we do not expect adverse events during the project.

Contacts

Public

Radboud Universiteit Nijmegen

P.O. Box 9101
6500 HB Nijmegen
NL

Scientific

Radboud Universiteit Nijmegen

P.O. Box 9101
6500 HB Nijmegen
NL

Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adults (18-64 years)

Elderly (65 years and older)

Inclusion criteria

Healthy, right-handed males aged 18-35 years. All subjects will have normal or corrected-to-normal vision.

Exclusion criteria

Contra-indications for TMS and fMRI: drug abuse, head trauma, neurological or psychiatric illness, pregnancy, heart disease, claustrophobia, cardiac pacemakers, metal objects in the body, medication pumps, tricyclic antidepressants, neuroleptics and a family history of neurological illness, psychiatric illness or epilepsy.

Study design

Design

Study type: Interventional

Masking: Open (masking not used)

Control: Uncontrolled

Primary purpose: Other

Recruitment

NL

Recruitment status: Recruitment stopped

Start date (anticipated): 29-03-2010

Enrollment: 40

Type:

Actual

Ethics review

Approved WMO

Date:

23-03-2010

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

NL29916.091.09