

Unraveling OCD: Using Brain Stimulation to Investigate Control

Submission date 15/04/2024	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 03/05/2024	Overall study status Completed	<input checked="" type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 27/01/2025	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

Obsessive-compulsive disorder (OCD) is a serious mental health condition where individuals experience persistent, unwanted thoughts (obsessions) and feel compelled to perform repetitive behaviors (compulsions). These obsessions and compulsions can greatly disrupt daily life. While therapies like counseling and certain medications are successful for about half of patients, others may need alternative options like different medications or deep brain stimulation, which involves surgery and carries risks. Research is ongoing to explore new treatments, such as repetitive transcranial magnetic stimulation (rTMS), which has shown promise in treating OCD. In this study, we will investigate the effectiveness of a specific type of rTMS on reducing OCD symptoms and its impact on brain activity. To accurately assess its effectiveness, we will include a placebo stimulation in the study, where some participants will receive a simulated treatment that we do not expect to have an effect.

Who can participate?

Patients with OCD, 18 to 70 years old with no contraindications for rTMS

What does the study involve?

The participant was assigned by a computer program to one of two groups:

- a) receiving active 20Hz ACDC TMS, set at 100% of their motor threshold, consisting of 40 rounds of stimulation with 50 pulses each, lasting 18 minutes, totaling 2000 pulses per session, for 1 session;
 - b) receiving sham ACDC TMS, where a sham coil was applied to the mPFC for 1 session.
- Immediately after the stimulation, the participant underwent EEG testing while performing the Stroop task, and any side effects were assessed.

What are the possible benefits and risks of participating?

If you grant Consent, you will be contributing to the collection of data that will help decide whether TMS is an effective method in the treatment of OCD. At the same time, the conditions for which you are being treated may be improved. There is no honorarium for participating in the research.

The risks of rTMS are generally low, with the main risk being the induction of an epileptic seizure. Even this risk has so far been very rare, usually in people who are predisposed to

seizures. Therefore, people who have or have had epilepsy (even a single seizure) or suspected epilepsy, or who belong to a risk group for epileptic seizures, i.e. people after a serious head injury with unconsciousness, after a stroke, with increased intracranial pressure and, according to individual assessment, with other neurological diseases affecting the brain (brain tumours, multiple sclerosis, Parkinson's disease, etc.), cannot participate in rTMS (and this project). Due to the magnetic field induction, TMS cannot be performed on persons with metallic material in the head area (vascular clips, stimulator, cochlear implants, foreign bodies, etc.) except in the mouth. If any of the conditions and situations described above apply to you, please inform the researcher who will contact you.

Where is the study run from?

National Institute of Mental Health (Czechia)

When is the study starting and how long is it expected to run for?

November 2022 to October 2023

Who is funding the study?

Charles University (Czechia)

Who is the main contact?

Dr Olga Laskov, olga.laskov@nudz.cz

Contact information

Type(s)

Public, Principal investigator

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Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

140223

Study information

Scientific Title

The anterior cingulate double cone transcranial magnetic stimulation (ACDC TMS) as an instrument for the investigation of inhibitory control in patients with obsessive compulsive disorder (OCD)

Acronym

ACDC OCD

Study objectives

A single active ACDC TMS application in a patient population with OCD will lead to:

1. Enhancement of the inhibitory control performance as measured by the Stroop test - decrease in the interference,
2. Lower reaction times to incongruent stimuli (as previous studies showed),
3. Lower error count,
4. Decrease of the ERN, P3, and N2 amplitude in response to errors,
5. Increase of P3 and N2 latencies in response to errors,
6. Reduction of the theta amplitude in resting EEG compared to sham dTMS and
7. At the same time, we expect that the behavioral and electrophysiological changes induced by the application of active dTMS will be positively correlated with Y-BOCS rates and negatively correlated with disease duration.

Ethics approval required

Ethics approval required

Ethics approval(s)

approved 04/11/2022, Ethics committee of the Third Faculty of Medicine, Charles University (Ruska 87, Prague, 10000, Czech Republic; +420 267102111; marek.vacha@lf3.cuni.cz), ref: GAUK140223

Study design

Randomized placebo-controlled double-blind study

Primary study design

Interventional

Study type(s)

Other, Treatment

Health condition(s) or problem(s) studied

Inhibitory control in patients with obsessive compulsive disorder

Interventions

The participant was randomly (by computer program) allocated to one of the two groups: a) active 20Hz ACDC TMS, 100% of motor threshold, 40 trains, 50 pulses per train, 18 minutes duration; 2000 pulses per session; 1 session; b) sham ACDC TMS with a sham coil applied either to mPFC; 1 session. Right after the stimulation: EEG with Stroop task and side effect assessment took place.

Intervention Type

Device

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

ACDC TMS was administered using a MagPro R30 stimulator with a cool D-B80 A/P coil

Primary outcome(s)

Inhibitory control measured via the interference effect in the Stroop task, measured as the reaction time to incongruent minus the reaction time to congruent stimuli right after the active and sham ACDC TMS stimulation

Key secondary outcome(s)

Measured right after the active and sham ACDC TMS stimulation:

1. Inhibitory control measured via the reaction times change and error count in the Stroop task (as less sensitive markers)
2. Electrophysiological effect of the active and sham ACDC TMS stimulation measured using EEG

Completion date

31/10/2023

Eligibility

Key inclusion criteria

1. Male and female inpatients or outpatients from the daycare center entering six-week CBT at age 18-70 years
2. Meet DSM-V criteria for OCD as determined by Structured Clinical Interview for DSM-5 (SCID-5)
3. The mental ability to understand and sign Informed Consent Form

4. Being on a stable and adequate dose of antidepressants and/or antipsychotics (monotherapy or combination) for at least one week prior to enrolment and if clinically appropriate to continue on an unchanged dose of medication during the trial

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Upper age limit

70 years

Sex

All

Total final enrolment

30

Key exclusion criteria

1. Psychiatric comorbidity on axis I and II according to DSM V in six months before enrollment to the study
2. Personality disorder that makes participation in the trial difficult
3. History of substance dependence in the last year except nicotine
4. Contraindications of rTMS (history of epilepsy or any neurologic condition likely to increase risk of seizure, mass brain lesions, cerebrovascular accident, metal in the head, a history of major head trauma with unconsciousness longer than 5 minutes)
5. Pregnancy or breast-feeding
6. Patients with severe somatic disorders (cardiovascular disease, neoplasms, endocrinology disorders etc.)
7. Patients treated with electroconvulsive therapy less than 3 months before enrollment or suffering from neurologic disorder (e.g., epilepsy, head trauma with loss of consciousness) and patients using any treatment which can strongly affect EEG
8. Substantial suicidal risk as judged by the treating psychiatrist
9. Sensory and motor impairment precluding the participation in computer tests

Date of first enrolment

01/12/2022

Date of final enrolment

31/10/2023

Locations

Countries of recruitment

Czech Republic

Study participating centre

National Institute of Mental Health

Topolova 748

Klecany

Czech Republic

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Sponsor information

Organisation

Charles University

ROR

<https://ror.org/024d6js02>

Funder(s)

Funder type

University/education

Funder Name

The Charles University Grant Agency (GA UK)

Results and Publications

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study will be available upon request from Dr. Olga Laskov olga.laskov@nudz.cz

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Participant information sheet	in Czech		17/04/2024	No	Yes
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes

[Statistical Analysis Plan](#)

[Study website](#)

Study website

17/04/2024	No	No
11/11/2025	11/11/2025	No
		Yes