
Open-label pilot to test a novel 1 Hz intensive repetitive transcranial magnetic stimulation paradigm in patients with anxiety.

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Previous Versions

Study Summary

Title	Open-label pilot to test a novel 1 Hz intensive repetitive transcranial magnetic stimulation paradigm in patients with anxiety.
Short Title	Open-label 1 Hz rTMS pilot

IRB Number

Phase	Pilot Study
Methodology	Open-label
Study Duration	1 years

Study Center(s)	Single-center
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Objectives	Aim 1: Determine the effect of a 1-week course of intensive 1 Hz stimulation on anxiety potentiated startle Aim 2: Determine the effect of a 1-week course of intensive 1 Hz stimulation on attention control.
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Number of Subjects	10
Main Inclusion and Exclusion Criteria	Anxiety patients, between 18 – 50 years old, free of psychological and neurological conditions, free of contraindications for TMS and MRI

Statistical Methodology

Aim 1: Outcome measures. Anxiety potentiated startle (APS) Statistical Analysis. The main analysis will be a paired-sample t-test (pre vs. post stimulation).

Data and Safety Monitoring Plan

Aim 2: Outcome measures. Accuracy and reaction time during the visual short term memory task (VSTM). Statistical Analysis. The main analysis will be a 2 [stimulation: pre-stimulation vs. post-stimulation] by 2 [load: low vs. high] repeated-measures ANOVA.

The Principal Investigator is responsible for detecting, documenting, and reporting unanticipated problems (UPs), adverse events (AEs), including serious adverse events (SAEs), and deviations in accordance with IRB requirements and federal regulations.