
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

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| 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 |
| Objectives | <p>Aim 1: Determine the effect of a 1-week course of intensive 1 Hz stimulation on anxiety potentiated startle</p> <p>Aim 2: Determine the effect of a 1-week course of intensive 1 Hz stimulation on attention control.</p> |
| Number of Subjects | 10 |
| Main Inclusion and Exclusion Criteria | Anxiety patients, between 18 – 50 years old, free of psychological and neurological conditions, free of contra-indications for TMS and MRI |

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

**Statistical
Methodology**

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.

**Data and Safety
Monitoring Plan**

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.