

Investigating Cytochrome c Oxidase Activity and Hemodynamic Responses During TMS Using Broadband NIRS

Violeta Casero¹, Saurabh Sonkusare¹, Yifan Cui¹, Nicholas Gregory^{2,3}, Nathaniel Tung¹, Gemma Bale^{2,3}, Qiaoting Huang¹, Valerie Voon¹

Department of Psychiatry, University of Cambridge¹; Department of Physics, University of Cambridge²; Department of Engineering, University of Cambridge³

Study Overview

Introduction: Broadband Near-Infrared Spectroscopy (bNIRS) is a non-invasive, non-ionizing technique that enables bedside, in vivo monitoring of the hemodynamic and metabolic activity of the cerebral cortex [1].

Objective: This study aims to evaluate the efficacy of bNIRS in measuring functional responses to transcranial magnetic stimulation (TMS).

Methods: A miniature broadband NIRS system was employed across three TMS sessions targeting the right frontal pole. Thirty-one healthy participants (N = 31) underwent two intervention arms—continuous theta burst stimulation (cTBS) and intermittent theta burst stimulation (iTBS)—against a sham comparator.



Figure 1: Schematic of bNIRS-TMS pairing during session procedure. bNIRS monitoring was conducted over a 20-minute period, with TMS intervention administered at the 10-minute mark.



Figure 2: Flow chart illustrating the bNIRS data analysis pipeline, from raw data acquisition to statistical analysis.

Results and Discussions

cTBS Effects:

- Increased ΔHBO₂, ΔoxCCO (P<0.001)
- Decreased ΔHHB (P<0.001)
- Suggests neural activation and effective modulation [2].
- Excitatory-like effects may result from altered neurovascular coupling or compensatory excitatory responses within the frontal pole [3].

iTBS Effects:

- Decreased ΔHBO₂, ΔHHB (P<0.001)
- Increased ΔoxCCO (P<0.001)
- Suggests heterogeneity in neurovascular-metabolic responses across individuals [4].



Figure 3: Mean changes (lines) and standard deviations (shaded areas) in oxyhemoglobin (Δ HBO₂), deoxyhemoglobin (Δ HHB), and cytochrome c oxidase (Δ oxCCO) following cTBS (blue), iTBS (red), and sham (grey) conditions. Data was smoothed using a Savitzky-Golay filter for visualisation purposes only.

Conclusion

This study demonstrates that bNIRS effectively captures neural modulation following TMS. Findings highlight the utility of bNIRS in monitoring neurovascular and metabolic changes during brain stimulation and underscore the importance of accounting for regional differences in neurovascular coupling and individual variability in future neuromodulation protocols.

References:

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