

Evaluating Transcranial Magnetic Stimulation as a Biomarker for Mild Traumatic Brain Injury: An mTBI-Predict Consortium Study

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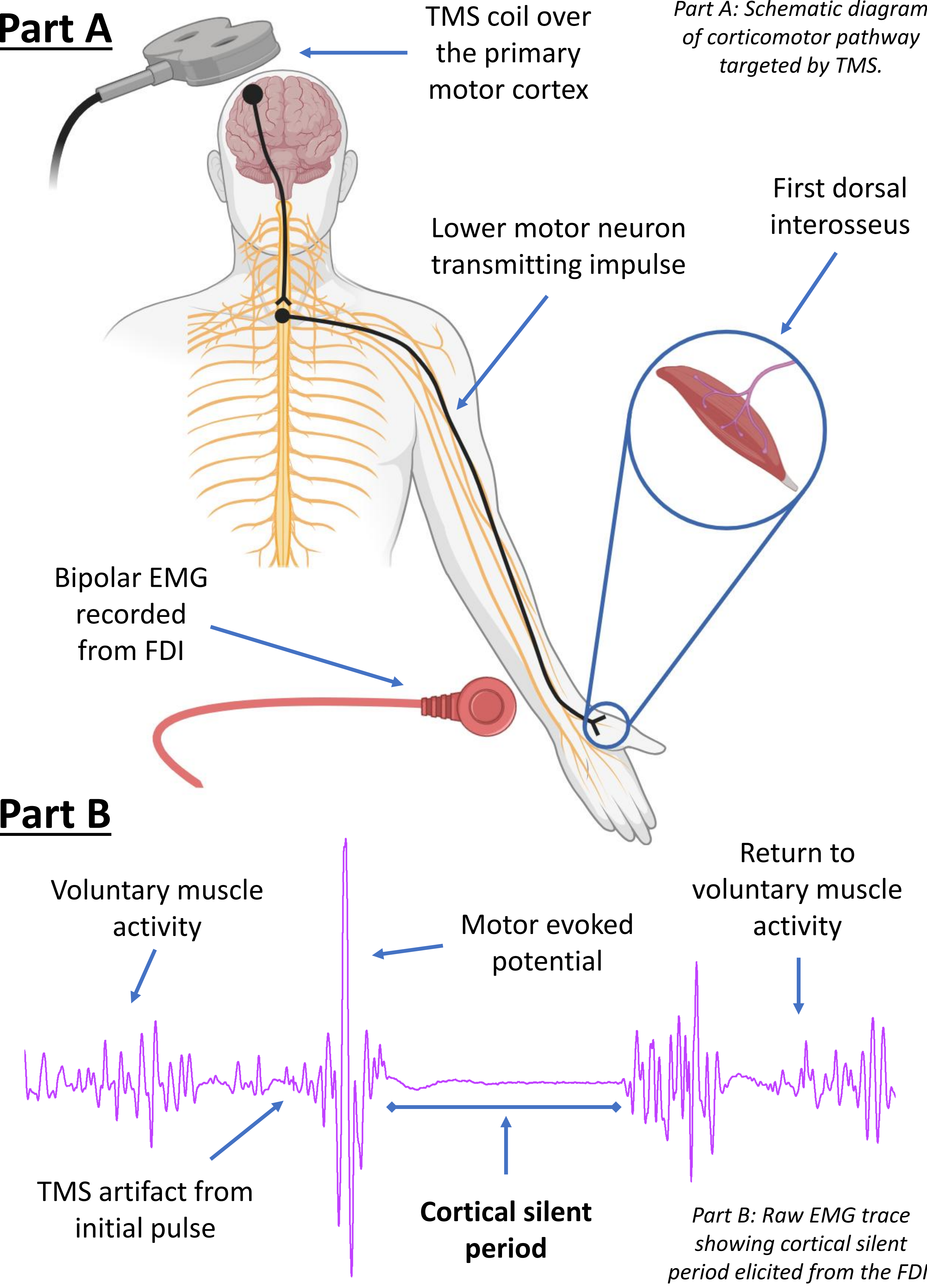
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Background

- Mild traumatic brain injury (mTBI) is estimated to cost \$17 billion USD annually in the USA, with over 2 million cases each year.
- Within the UK, 1.2 million cases are seen annually, with a third of patients unable to work within a year of their injury.
- New biomarkers are emerging in an attempt to assess recovery following mTBI.
- The cortical silent period (CSP) has been suggested as one such biomarker.
- CSP is a temporary suppression of voluntary muscle activity seen in the electromyography (EMG) of a contracted muscle following a single pulse of transcranial magnetic stimulation (TMS).

Study aim: To assess the variability and utility of the cortical silent period as a characterising biomarker for mild traumatic brain injury.

Methods (1)



Methods (2)

- 11 mTBI patients and 13 healthy control volunteers were recruited.
- Each participant completed four visits across 12 days.
- Single-pulse TMS was delivered to the motor cortex.
- Bipolar EMG was recorded from the first dorsal interosseus (FDI).
- During a handgrip squeeze at 30% maximal voluntary contraction, CSP was elicited from the FDI.
- Two trained operators manually extracted CSP duration.
- Within-subject and between-subject variability was assessed to determine biomarker suitability.

Results

Table 1. Participant characteristics	mTBI Patients (n = 11)	Healthy control volunteers (n = 13)
Age (years)	31 ± 11	34 ± 13
Sex (n, %), Male	6 (55)	8 (62)
Body mass index (kg/m ²)	27.87 ± 4.99	27.01 ± 5.57
Time since injury (days)	29 ± 4	N/A
Total visits analysed (n)	42	49

Table 2. CSP variability	mTBI Patients (n = 11)	Healthy control volunteers (n = 13)
Mean duration (ms)	138.9 ± 34.4	133.3 ± 32.9
Within-subject variability (CV _I , %)	16.04	17.62
Between-subject variability (CV _G , %)	18.86	17.26

- Demographic characteristics were similar across mTBI and healthy groups.
- Primary outcome variables of within-subject and between-subject variability did not show differences.
- Variability was also similar between and within participants.

Discussion

- Due to similar variability between mTBI and healthy individuals, CSP alone may not be a strong biomarker for characterising mTBI.
- CSP duration was slightly higher in mTBI, but further work is needed to investigate significance.
- Ongoing work will assess the usefulness of CSP as a prognostic biomarker for mTBI alongside other potential biomarkers such as vestibular, sleep, and biofluid derived variables.

Funding and conflicts

UK mTBI-Predict consortia contract from the Ministry of Defence (MOD) and Department of War (DoW). NIHR BRC Birmingham. Approved by MOD Research Ethics Committee (REC-2217MODREC23). The views and conclusions contained herein are those of the authors and should not be interpreted as necessarily representing the official policies or endorsements, either expressed or implied, of the U.S. Government. The material has been reviewed by the Defense Health Agency. There is no objection to its presentation and/or publication. The opinions or assertions contained herein are the private views of the authors and are not to be construed as official or as reflecting the views of the DoW.