TMS SMART – Scalp Mapping of Annoyance Ratings and Twitches caused by Transcranial Magnetic Stimulation

Lotte Meteyard[1]

Nicholas Holmes[2]

[1] School of Psychology and Clinical Language Sciences, University of Reading, Whiteknights, RG6 6AL. email: l.meteyard@reading.ac.uk. Tel: +44(0)118 378 8142

ORCID ID 0000-0003-1184-1908

[2] School of psychology, University Park, University of Nottingham, Nottingham, NG7 2RD. Email:

npholmes@neurobiography.info

Key Words

TMS, experimental control, peripheral effects, TMS side-effects, control conditions.

Acknowledgements

Abstract

The magnetic pulse generated during Transcranial magnetic stimulation [TMS] also stimulates cutaneous nerves and muscle fibres, with the most commonly reported side effect being muscle twitches and sometimes painful sensations. These sensations affect behaviour during experimental tasks, presenting a potential confound for 'online' single-pulse TMS studies. Our objective was to systematically map the degree of disturbance (ratings of annoyance, pain, and muscle twitches) caused by TMS at 43 locations across the scalp. Ten participants provided ratings whilst completing a choice reaction time task, and ten participants provided ratings whilst completing a 'flanker' reaction time task. TMS over frontal and inferior regions resulted in the highest ratings of annoyance, pain, and muscle twitches caused by TMS. In separate analyses we predicted the difference in reaction times (RT) under TMS by scalp location and subjective ratings. Frontal and inferior scalp locations showed the greatest cost to RTs under TMS (i.e., slowing), with midline sites showing no or minimal slowing. Increases in subjective ratings of disturbance predicted longer RTs under TMS. Critically, ratings were a better predictor of the cost of TMS than scalp location or scalp-to-cortex distance, and the more difficult 'flanker' task showed a greater effect of subjective disturbance. The peripheral sensations and discomfort caused by TMS pulses significantly and systematically influence RTs during single-pulse, online TMS experiments. We provide the data as an online resource (www.tms-smart.info) so that researchers can select control sites that account for the level of general interference in task performance caused by online single-pulse TMS.

1.0 Introduction

A transcranial magnetic stimulation (TMS) machine produces a rapidly-changing magnetic field which, when positioned on the scalp, can be used to disrupt function temporarily in a specific brain area (Sparing & Mottaghy, 2008; Wagner, Valero-Cabre, & Pascual-Leone, 2007; Walsh & Cowey, 2000). This provides a non-invasive means to draw inferences about the contribution of specific brain regions to a given behavioural task. (Bolognini & Ro, 2010; Ruff et al., 2009).

TMS causes peripheral auditory and cutaneous sensations (Bestmann et al., 2005; Borckardt et al., 2006; Nikouline, Ruohonen, & Ilmoniemi, 1999; Starck et al., 1996; Wasserman, 1998). Tension headaches and scalp discomfort are frequently reported side-effects of undergoing TMS (Anderson et al., 2006; Benninger et al., 2011; Maizey et al., 2013; O'Reardon, and et al., 2007; VonLoh et al., 2013).

These peripheral sensations differ by scalp location. TMS is more disturbing over prefrontal cortex than over parietal cortex (Abler et al., 2005). Crying has been reported as an adverse effect for participants in speech studies that stimulated left prefrontal areas (Wasserman, 1998). Greater discomfort over frontal regions of the scalp is likely due to a greater density of muscles and nerves (Maizey et al., 2013; Wassermann, 1998; Machii et al., 2006; Loo et al., 2008). Discomfort is related to the intensity and frequency of stimulation but even single-pulse TMS can cause notable discomfort, especially with participants naïve to TMS (Maizey et al., 2013). Notably, ratings of scalp discomfort caused by TMS positively correlate with the number of errors made on a delayed match to sample task (Abler et al., 2005).

Appropriate control conditions are needed to ensure that results are due to the changes in brain activity caused by TMS, not as a consequence of peripheral effects (Arana et al., 2008; Duecker & Sack, 2013 & 2015; Sandrini et al., 2011). There are several options for control conditions in TMS: the use of additional behavioural tasks or conditions, a no TMS condition (e.g., Tamè & Holmes, 2016), the use of sham TMS and application of TMS to a control location on the scalp.

Duecker & Sack (2015) provided a review of sham TMS, for which a number of conditions and devices have been evaluated and developed (e.g., Arana et al., 2008; Rossi et al., 2007; Sommer et al., 2006). Sham TMS is designed to mimic the sound and sometimes the peripheral sensations of TMS, without changing cortical activity (Borckardt et al., 2008; Duecker & Sack, 2015), and it is

possible to create a sham condition under which participants cannot reliably judge whether they received sham or real rTMS (Duecker & Sack, 2015; Herwig et al., 2010). However, sham TMS has to deliver high intensity cutaneous sensations (70% of participant's maximum pain tolerance) to be matched to real TMS (Arana et al, 2008).

Control locations for TMS should attempt to control for peripheral sensations, yet to our knowledge, this control is not often implemented. Ideally, a selected control site should give the same level of comfort or discomfort as the experimental site. The vertex (Cz, at the top of the head) is often used, under the assumption that the brain tissue is deeper under the scalp, attenuating cortical stimulation (e.g., Kalbe et al., 2010; Sandrini et al., 2011; Silvanto et al., 2008; Whitney et al., 2010; Vetter et al., 2015; but see Fox et al., 1997; Jung et al., 2016; Loo et al., 2000). It is not clear that this location controls well for cutaneous sensations and discomfort, particularly when the experimental site is frontal and known to be more painful (Wasserman, 1998).

The best-practice in TMS studies may be to have more than one control condition, with multiple stimulation sites or stimulation time points (Duecker & Sack, 2105; Miall et al,. 2008; Tamè & Holmes, 2016) and a control task (Sandrini et al., 2011). However, the selection of control conditions remains sub-optimal.

The aim of the study reported here is to map the subjective discomfort of TMS across the scalp. Rather than comparing a small number of sites, we aimed for coverage of the entire scalp, in order to create a detailed map of the frequency of muscle twitches, and participant ratings of annoyance and pain (extending Abler et al, 2005; Arana et al., 2008; Loo et al., 2000). We used two different tasks to test whether task difficulty would interact with the application of. That is, does greater discomfort under TMS affect a difficult task more than an easier task? Together with an accompanying website (http://www.tms-smart.info), this data provides a resource that researchers can use to control for peripheral sensations caused by TMS.

2.0 Method

2.1 Participants

Twenty individuals took part. Ten completed a choice-RT (CRT) ask and ten completed a 'flanker' task. All participants provided written informed consent and underwent screening in line with the TMS Rules of Operation at the Centre for Integrative Neuroscience and Nuerodynamics, University of Reading. The study received a favourable opinion for conduct from the University of Reading Research Ethics Committee (UREC Project 13/03).

For the choice RT task, mean age was 31.6 years (SD=5.5), 6 were female and all were right handed by self-report. Mean resting motor threshold, examined by visual inspection of muscle twitches in the hand, was 51.9% (SD=7.7%) of maximum stimulator output. For the flanker task, mean age was 25.2 years (SD=3.5 years), 6 were female and 8 were right handed by self-report. Their mean resting motor threshold was 48.5% (SD=6.4%) of maximum stimulator output (Rossini et al., 1994). All participants completed the ratings task. No participants or other data were excluded from the final dataset prior to analysis.

2.2 Scalp locations

We aimed for coverage of the entire scalp. To begin, we transferred locations from a 32 channel EEG cap onto a Lycra skull cap (see Herwig et al., 2003 and Homan et al., 1987 for the 10:20 system being used position the TMS coil). The 32 locations were marked using BrainSight Frameless Stereotaxy (Rogue Resolutions, Cardiff, Wales), and registered to a T1 structural scan (author NPH). This registration made clear that the occipital, temporal, and cerebellar areas were not fully covered, so we added 5 sites per hemisphere and one midline site. This gave a total of 24 scalp locations in each hemisphere (5 in the midline, and 19 covering each hemisphere; Figure 1). For occipital sites, we used 6 cm distance measures as a rough guide from point to point marked on the cap using the 10:20 system. For the midline we took a line from Cz to Oz (our location: Inion). We also took lines from Pz to O2/O3 (our location: Lateral Inion 1 & 2), P3 to P7/P8 (our location: Lateral Occipital 1 & 2). For other sites, we used the registration data available in BrainSight to add locations approximately corresponding to the following: Lateral-Occipital (3 & 4), Temporal-Parietal Junction (TPJ 1 & 2) and the Anterior Temporal Lobes (ATL 1 & 2; see Figure 1).

Figure 1: Locations of cortical sites stimulated with TMS and axes of stimulation.

Legend/Key: Scalp locations on the Lycra cap were assigned numbers to support randomisation and blinding during the experiment. For each participant Cz was aligned to their vertex and this alignment was checked periodically during testing.
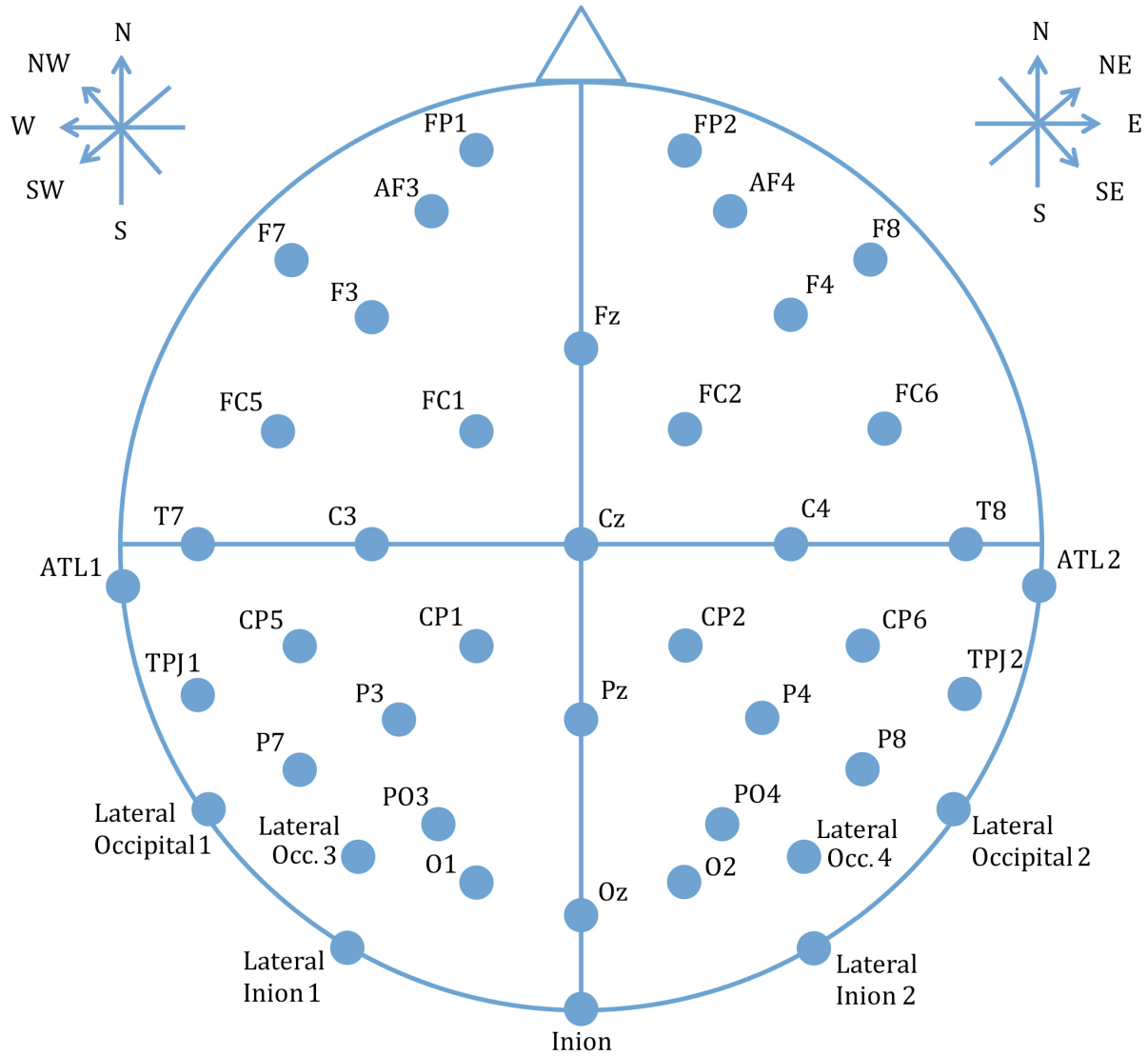
2.3 Coil orientations

To measure the influence of coil orientation we used a cardinal reference system with North taken as the direction faced by the participant. This was preferable to any allocentric (e.g., superior, inferior, anterior, posterior) or neuraxis-centred (e.g., ventral, dorsal, rostral, caudal) reference system, since coil orientation did not change as the coil moved along the scalp. 'South' was always with the handle pointing along the surface of the scalp following 'lines of longitude' towards the inion; 'East' was always with the handle pointing along the surface of the scalp following similar lines of longitude (not latitude) towards the right ear. Each location was stimulated biphasically using four coil orientations that mapped 45 degree rotations around a circle. Orientations were North-South (along the midline), North East-South West (45° going clockwise), East-West (along the ear-to-ear axis, 90°), and North West-South East (135°, see Figure 1). The TMS coil handle was aligned to the above axes, usually oriented towards the back of the participant's head, and away from the face. It was rotated to lie in the opposite direction along the axis if this position was not possible (e.g., if the shoulder was in the way for inferior and lateral sites).

2.4 TMS intensity

We kept the intensity constant at 50% of the maximum stimulator output. This is the approximate mean intensity for eliciting MEPs and visible muscle contractions with this equipment in our laboratory. Researchers using TMS often change the intensity of stimulation on a participant-by-participant basis, for example, by setting the intensity to a fixed proportion of the resting (RMT) or active motor threshold (AMT). This approach is appropriate when stimulating over motor cortex in order to affect or elicit motor-evoked potentials in a contralateral muscle. A similar approach may be useful for eliciting phosphenes by stimulating over occipital cortex (Siniatchkin et al., 2011). However, much less is known about the relationship between TMS intensity and assumed cortical activation for all other brain areas (e.g. Peurala et al., 2008). We chose 50% intensity to provide a realistic level of stimulation (at approximate RMT) within the typical range for a TMS study, and therefore reflecting a typical level of discomfort. This level of stimulation is likely to have stimulated the cortex of many, if

not all of the sites used in our experiment. However, because of differences in scalp to brain distance between participants and sites (Table 1), the amount of cortical stimulation likely differed across sites. We had two options for setting TMS intensity: 1) adjusted for each participant and site, or 2) kept constant across participants and sites.

We could have attempted to set the TMS intensity to obtain approximately the same level of cortical stimulation across all participants and sites. Using MEPs as an output variable, systematically mapping over the motor cortex to find the site and coil orientation that induces MEPs at the lowest intensity, then found the RMT and AMT for each participant, then decreased the intensity so that no effect of TMS was visible on the electromyograph (EMG). Using this ineffectual TMS intensity, we could then have applied a scalp to brain distance adjustment (Stokes et al., 2007) to adjust the TMS intensity for all remaining TMS sites. This method would have the advantage of being less likely to have stimulated the cortex, but it would have likely resulted in TMS intensities of 20-30% of the maximum stimulator output, much lower than typically used in experiments. Most importantly, this method of TMS intensity adjustment per participant and site would have confounded the purpose of the experiment: to measure the relative annoyance of TMS between scalp sites at levels of stimulation typically used in experiments. In addition, different TMS intensities at different scalp locations would render the ratings of annoyance, pain and twitches meaningless for further analysis. In order to systematically map the difference in subjective discomfort across scalp sites, TMS intensity needed to be constant whilst we differed scalp location.

Thus, we maintained TMS intensity constant, at 50% of maximum stimulator output. Ours is a study of the scalp, not of the brain. We accept, of course, that stimulation of the underlying cortex was very likely for many, if not all scalp locations and participants. Nevertheless, it is parsimonious to assume that our results relate best to regions of the scalp, targeted with constant-intensity TMS, rather than to regions of the brain, not targeted and with variable-intensity TMS. We also completed post-hoc analyses to control for scalp-to-brain distance (see below).

2.5 Tasks and stimuli

All stimuli were presented using Matlab and Psychtoolbox 3 (Brainard, 1997) in black text on a white background, font size 20, screen resolution 1920 x 1080 pixels. Stimuli were presented using a CRT monitor with 75 Hz refresh rate.

2.5.1 Choice reaction time task

This task was a basic spatial-compatibility, choice RT task designed to be simpler than the flanker task (below). Stimuli were the left and right chevrons (< and >). A fixation cross was presented centrally. Three chevrons appeared on the left or right of the fixation cross, displaced 100 pixels horizontally from the central fixation cross. These could be left or right facing, as above. Therefore, there were two congruent conditions (<<< + , + >>>) and two incongruent conditions (>>> +, + <<<). Participants were instructed to press C with one finger if the chevron appeared on the left, and N with another finger if it appeared on the right. These are spatially-compatible responses as the C key is on the left and the N key on the right of the keyboard. Speed and accuracy were both encouraged. When the data were inspected, three participants had responded to arrow direction rather than the side of the screen on which the arrow appeared. As this still constituted a choice RT task with spatial compatibility (the congruence conditions were still valid), as the behavioural task was not the primary purpose of the present report, and as their data were comparable to other participants' (mean RT, proportion correct, and size of congruency effect, see Appendix A), these three participants' data were included in the final analysis. Removing these data points and re-analysing did not change the conclusions drawn.

2.5.2 Flanker task (Eriksen & Eriksen, 1974)

Stimuli were the left and right chevrons (< and >). Three chevrons were presented centrally, and faced left (<<<) or right (>>>). Flanker chevrons facing left or right were presented either side, displaced 100 pixels horizontally from the centre of the screen. Therefore, there were two congruent conditions (<<< <<< <<<, >>> >>> >>>) and two incongruent conditions (<<< >>> <<<, >>> <<< >>>).

Participants were instructed to press the C key with one finger if the central chevrons pointed left, and N with another finger if they pointed right. Speed and accuracy were both encouraged.

2.5.3 Ratings Task

Participants were asked to rate the annoyance, pain, and peripheral muscle twitches in the head, face and neck caused by TMS. Each was rated on a scale of 0 to 10, with 0 being the minimum and 10 the maximum. For annoyance ("How annoying were the TMS pulses?"), anchor points (printed on a sheet of paper for reference) were "not at all annoying" (0) to "highly distracted and unable to complete the task" (10). For pain ("If painful, how intense was the pain from the TMS pulses?"), anchors were "no pain at all" (0) to "the worst pain they could tolerate during the experiment" (10). We specified pain intensity, so that participants would focus on the sensory aspects of the pain (intensity, quality) rather than their emotional response (i.e., pain unpleasantness) that would already be partly captured by the rating of annoyance (Melzack & Casey, 1968). For twitches ("If there were any twitches, how strong were the muscle twitches from the TMS pulse?"), anchors were "no twitches" (0) to "a very strong cramp" (10). In addition to the subjective ratings provided by the participants, a second experimenter recorded whether any visible twitches were observed in the face, head or neck that were coincident with the TMS pulse (0 to 5 twitches observed in each block). To facilitate this, a mirror was angled in front of the participant so the researcher could sit or stand behind them and observe. The participant was not able to observe themself in the mirror.

2.6 Design and procedure

The Lycra cap with scalp locations was fitted to the participant with Cz aligned to their vertex, and periodically checked during the experiment. Resting motor threshold was first established for each participant (Rossini et al., 1994). We identified the approximate location of M1 by measuring 5cm lateral and 1cm anterior from vertex for the hemisphere to be tested (right or left). Participants rested their ipsilateral hand on the desk, clearly visible to the experimenters. Test TMS pulses were delivered around this location until a visible twitch was seen in the thumb, index, or middle fingers (typically originating from the 1st or 2nd dorsal interosseous, adductor pollicis, or opponens pollicis muscles). Stimulator intensity was varied until a visible twitch was seen on 5 out of 10 consecutive

pulses delivered at approximately 0.2Hz. This stimulator intensity was recorded as the resting motor threshold (RMT).

For the behavioural experiment, stimulation strength was set at 50% of the maximum stimulator output (biphasic). As confirmed by our data, this level was selected to be approximately at RMT. Single-pulse TMS studies typically stimulate at 110-140% RMT. However, since RMT assessed visually (as we did) is likely to be 11% higher than RMT assessed with EMG (Westin et al., 2014), this intensity of stimulation was likely above threshold for stimulating cortex. One hemisphere per participant was stimulated at all scalp locations during the experiment. The order of testing Scalp Location was fully randomised for each participant. Coil orientation was nested within location and then randomised (i.e., a randomly selected permutation of all possible orders of the four coil orientations was selected once location had been selected). Half of the participants had their left, and half their right hemisphere stimulated. This between-subjects manipulation of hemisphere was a pragmatic decision that allowed data collection to be completed in one two-hour session (partly to meet the requirements of our local ethics committee). To account for any potential confound from this manipulation, data analyses included Hemisphere and the interaction between Hemisphere and Scalp Location as predictors of interest.

Participants provided button press responses using the hand ipsilateral to the TMS, to minimise any chance of cortical stimulation from the TMS influencing hand motor control and response times. Single pulse TMS was delivered on each trial using a figure-of-eight coil.

Each task had identical timings and trial and block structure. Participants completed 60 practice trials with no TMS (coil disabled, away from head; 3 blocks of 20 trials, with each block split into 4 mini-blocks of 5 trials to reflect the actual experiment). They were then introduced to the rating task and response scales. To explain the rating procedure, participants were asked to remember the TMS pulses delivered during the RMT procedure and consider the sensations (annoyance, pain, and twitches) that they may have experienced. Thus, these pulses acted as reference pulses for participants' ratings of annoyance, pain, and twitches. During the experiment, participants completed 5 experimental trials for each combination of coil orientation and scalp location. The experiment was structured so that a location was randomly selected, and then each of 4 randomly-ordered coil orientations was completed for that location (4 blocks of 5 trials for each location; 20 trials per

location). In each block of 5 trials, a fixation cross appeared for a randomly selected duration between 500 and 1000ms, followed by the imperative stimulus. A single TMS pulse was delivered at the same time as the imperative stimulus appeared on screen. Stimuli appeared on screen until the participant responded or until 2 seconds, whichever was sooner. There was an inter-trial-interval of 2 seconds. In each block of 5 trials, participants saw one of each trial type (congruent left, incongruent left, congruent right, incongruent right) and a randomly selected 5th trial. After completing the 5 trials of the task, they were asked to give ratings for the annoyance, pain, and twitches caused by the preceding 5 TMS pulses. Participants were asked to provide ratings after 5 trials to allow them to complete the behavioural task whilst experiencing more than one pulse at that location and with that coil orientation. The total number of trials per participant was 480 (5 trials x 4 orientations x 24 locations). In between each block of 5 trials, participants were able to take a break for as long as they wished.

2.7 Analysis

Ratings data were taken from all 20 participants and averaged across participants to give values for each coil orientation at each scalp location. Mean ratings (annoyance, pain, twitches) and measurement of observed visible twitches were strongly positively correlated with each other (all correlation coefficients>=0.85). This precludes them being used as separate fixed effect predictors due to collinearity. The self-reported intensity of twitches correlated the most strongly with all other variables (all correlation coefficients>=0.9), and was therefore used in further analyses as a fixed effect predictor.

RTs were taken from correct trials only. We used RTs from correct practice trials for baseline RT (the first 20 trials were discarded as true practice trials - we assumed that RT would be stable after these initial trials). The mean practice RT was taken for each trial type (congruent left, incongruent left, congruent right, incongruent right) and this was subtracted from the RT for that trial type during the TMS experiment, on a trial-by-trial basis. This gave us the *difference* in RT for a specific trial type, when the participant was undergoing TMS. Therefore, all analyses of RTs looked at how the predictor variables changed the impact of TMS on RTs. Inspection of initial linear mixed-effects models showed that residuals were not normally distributed, caused by a rightward skew typical in reaction time data (Baayen & Milin, 2010; Ratcliff, 1993). Following Baayen & Milin (2010), data trimming and

transformations, followed by model inspection, were explored to best normalize residuals for the models. The best results were achieved with a single cut-off value, excluding all reaction time differences longer than 400ms (the minimum reaction time difference was -389ms, so an upper cut-off was selected to mirror the minimum value). This removed 2.9% of the data.

To allow analysis of Scalp Location as a within subjects variable, we coded homologous scalp locations across left and right hemispheres together to look at the effect of scalp location on RT. For example, FP1 (left) and FP2 (right) were coded together, the two ATL sites were coded together, and so on.

There are three analyses. (1) A descriptive analysis of ratings across scalp locations and orientations. (2) Predicting RT by scalp location and orientation and (3) Predicting RT differences from subjective ratings. The second and third analyses were completed using linear mixed-effect models implemented in R (R Core Team, 2014); using the packages lme4 (Bates, Maechler, Bolker, & Walker, 2014), multcomp (Hothorn, Bretz, & Westfall, 2008), lmerTest (Kuznetsova, Brockhoff, & Bojesen, 2014), ggplot2 (Wickham, 2009) and corrplot (Wei, 2013). For these models, the reference level for scalp location was set to Cz/vertex, illustrating the effect of TMS across scalp locations relative to a commonly used control.

3.0 Results

3.1 Task performance

We compared error rates between Flanker and CRT tasks and found no significant difference (median errors Flanker=5 (range 0-44), median errors CRT=5.5 (range 0-66); Kruskal-Wallis=0.0230, df=1, p-value=0.879). Within task (Wilcoxon signed ranks test), there was no significant difference between congruent and incongruent conditions for the Flanker task (median errors congruent=1.5 (range 0-6), median errors incongruent=2 (range 0-41); W=35, p=0.264) and no significant difference for the CRT task (median errors congruent=3 (range 0-29), median errors incongruent=2.5 (range 0-37); W=49.5, p>0.900).

3.2 Scalp mapping of annoyance ratings and twitches

For participant ratings of subjective annoyance, pain, and twitches, and for the experimenter assessments of visible twitches, descriptive and inferential statistics were derived for each location, orientation, and hemisphere. A full account of these rich data is not possible here. Instead, we report the two dependent variables that we suggest are the most useful: The median ratings of muscle twitches and the mean effect of TMS on RT. Ratings of annoyance, pain, and twitches were strongly co-linear; twitches were most representative (see below). The effect of TMS on RT is likely the most useful measure for behavioural studies. The raw data are available at www.tms-smart.info for viewing and download, along with interactive maps of all available dependent variables and statistics, and a facility to find suggested control sites for each TMS location (10:20 system) and for individual MNI coordinates.

Median ratings of twitches (Figure 2, upper panels) varied from 0 (at superior sites on or adjacent to the midline) to 7 out of 10 (inferior frontal and anterior temporal sites). Ratings were symmetrical across the midline, and included four peaks – over the left and right anterior temporal/inferior frontal cortices, and over the left and right lateral occipital cortices.

Mean effects of TMS on RT (Figure 2, lower panels) showed a broadly comparable distribution across the scalp, with the largest impairments in RT (up to 181ms) seen at inferior frontal, anterior temporal, and lateral occipital sites. The smallest changes in RT occurred along and adjacent to the superior midline, while TMS-induced improvements in RT (up to 61ms) were seen at superior parietal and occipital sites in the left hemisphere. The following section assesses the relationships between participant ratings and the effects of TMS on behaviour.

Figure 2: Interpolated maps of the effects of TMS location on the median participant rating of muscle twitches (upper panels) and the mean reaction time (RT, lower panels), averaged across the four coil orientations.
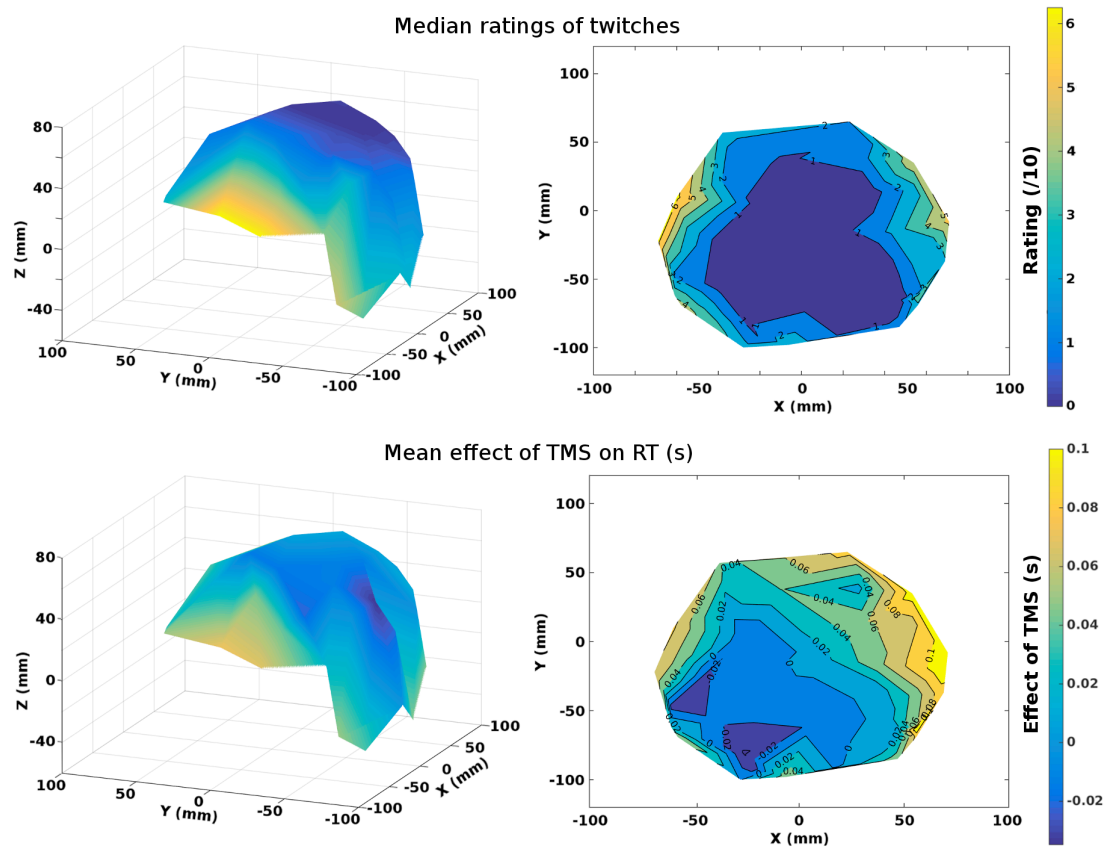
Legend/Key: The maps in the left column show a 3D interpolation and rendering, viewed from behind the participant's left ear, of the raw data mapped to the approximate Montreal Neurological Institute (MNI) coordinates (X, Y, Z, mm), as calculated in FSL, using a 12 degree-of-freedom affine transformation in FLIRT. The maps in the right column show a 2D plan view (from above) of the same data, and include the numerical values in the contour maps to show the scale. Dark blue shows low ratings (upper panels) or RT decrease (lower panels); while bright yellow shows higher ratings (upper panels) or RT increases (lower panels).

3.3 Predicting reaction times by scalp location and coil orientation

Random intercepts were fit for each participant to account for participant-by-participant variation in RTs (variance=1055ms). Following Barr et al (2013) we fit random effects to control for the within subjects variation in the effects of Scalp Location (variance=1360ms), Trial (trial within each scalp location, a number between 1-20; variance=425ms), Coil Orientation (variance=15ms), and Congruence (variance=581ms). To check that Scalp Location could be coded as homologous locations across Hemispheres, we tested the interaction between Scalp Location and Hemisphere. The interaction did not significantly improve model fit when compared against a model with Scalp Location as a main effect only (X^2 goodness of fit=21.884, df=24, p=0.59). However, three scalp locations showed significant differences between hemispheres. CP5&6 (estimate=57.23ms, Standard Error=24.88ms, df=73, t=2.301, p=0.024), P7&8 (estimate=50.27ms, SE=25.00ms, df=75, t=2.011, p=0.048) and FC1&2 (estimate=55.54ms, SE=24.88, df=73, p=0.029). In all cases, RT costs under TMS were greater for the right hemisphere. Therefore, the interaction of Scalp Location and Hemisphere was retained in the final model as a control. To control for the effect of preceding trials (i.e. if the previous trial had been particularly painful, or an error trial) we included the reaction time from the previous trial. This was taken from within each block of 5 trials, so trial 1 had a value of 0 for the previous trial RT, trial 2 had the RT from trial 1, and so on, up to trial 5. This significantly improved model fit (X^2=158.87, df=1, p<0.001; estimate=0.09, SE=0.01, df=861.7, t=12.57, p<0.001). The main effect of Task improved model fit (X^2=4.443, df=1, p=0.035), the Flanker task showed a marginally greater impact of TMS on RTs than the CRT task (estimate=34.97ms, SE=16.99, df=17, t=2.06, p=0.055). The main effect of Congruence was not significant (p=0.366) nor did it improve model fit (X^2=0.8785, df=1, p=0.35). There was no significant interaction between Task and Congruence (p>0.1) and the interaction term did not improve model fit (X^2=1.92, df=2, p=0.38). The effect of Coil Orientation was not significant and did not improve model fit (X^2=4.86, df=3, p=0.18); there were no significant differences between any TMS coil orientations (Tukey Contrasts all p>0.6). Coil Orientation and Congruence were not included as fixed effects predictors in the final model. With random effects noted as re(), the final model that best explained the data was:

RT Difference under TMS ~ Scalp Location + Scalp Location x Hemisphere + Task + Previous RT + re(Participant) + re(Location x Participant) + re(Axes x Participant) + re(Congruence x Participant) + re(Trial x Participant)

The fixed effects alone had an R^2 of 0.14, and the full model had an R^2 of 0.43 (effect size of 0.80; Champely, 2016; Cohen et al., 2003; Selya et al., 2012; Lefcheck, 2015).

The effect of TMS on RTs varied across Scalp Locations, Figure 3 plots the mean RT difference data for each location, and Table 1 presents the estimated mean RT effect for each location (least squares means and standard errors, estimated difference against zero) and the approximate MNI location. Multiple comparisons (Tukey contrasts) comparing each Scalp Location to each other Scalp Location are provided in Appendix B. Here we present the estimated differences between each location and Cz/vertex, as this is a commonly used control location in TMS studies. Researchers wishing to re-reference the data to other sites and/or orientations can do so with the freely-available raw dataset at www.tms-smart.info. A number of sites did not differ significantly from Cz/vertex, these were C3&4, CP1&2, CP5&6, AF, PZ, P3&4, P7&8, FZ, F3&4, FC1&2, OZ, 01&2, TPJ, Lateral Occipital 3&4, Lateral Inion, and Inion. Six sites showed a greater cost of TMS than Cz/vertex. These were FC5&6 (est. mean=42.62ms, SE=18.58ms, t(383)=2.293, p=0.02); Lateral Occipital 1&2 (est. mean=53.43ms, SE=18.66ms, t(388)=2.864, p=0.004); ATL (est. mean=7.01ms, SE=18.56ms, t(381)=3.781, p=0.0002); T7&8 (est. mean=39.55ms, SE=18.62ms, t(386)=2.124, p=0.034), FP (est. mean=43.47ms, SE=18.70ms, t(392)=2.325, p=0.02), and F7&8 (est. mean=59.69ms, SE=18.64ms, t(387)=3.203, p=0.001). One site had faster RTs under TMS, significantly faster when compared against Cz/vertex. This was PO3&4 (est. mean=-36.69ms, SE=18.56ms, t(380)=-1.977, p=0.048). Figure 4 presents the RT difference in milliseconds for each scalp location (TMS RT – No TMS RT).

Table 1 presents least squares means for the effect of TMS on RTs, comparing each site against zero. The sites showing the biggest costs were frontal and lateral sites, some of which showed very high RT costs for TMS. The sites with the largest slowing of RTs under TMS were the ATL (~75ms cost), F7&8 (~70ms cost), Lateral-Occipital 1&2 (~65ms cost) and T7&8 (~55ms cost). The sites with the smallest cost were sites in the midline, some of which showed negligible changes. The standard error across sites was around 10ms, and the following sites showed a less than 10ms slowing under TMS: Cz (~2ms), F3&4 (~4.5ms), FC1&2 (~6ms), C3&4 (~3ms), CP5&6 (~5ms). The complete least squares means broken down by hemisphere and location are provided in Appendix B.

Figure 3: Effect of TMS by Scalp Location. Legend/Key: A positive difference means slower RTs under TMS, a negative difference means faster RTs under TMS. Error bars are 95% confidence intervals.
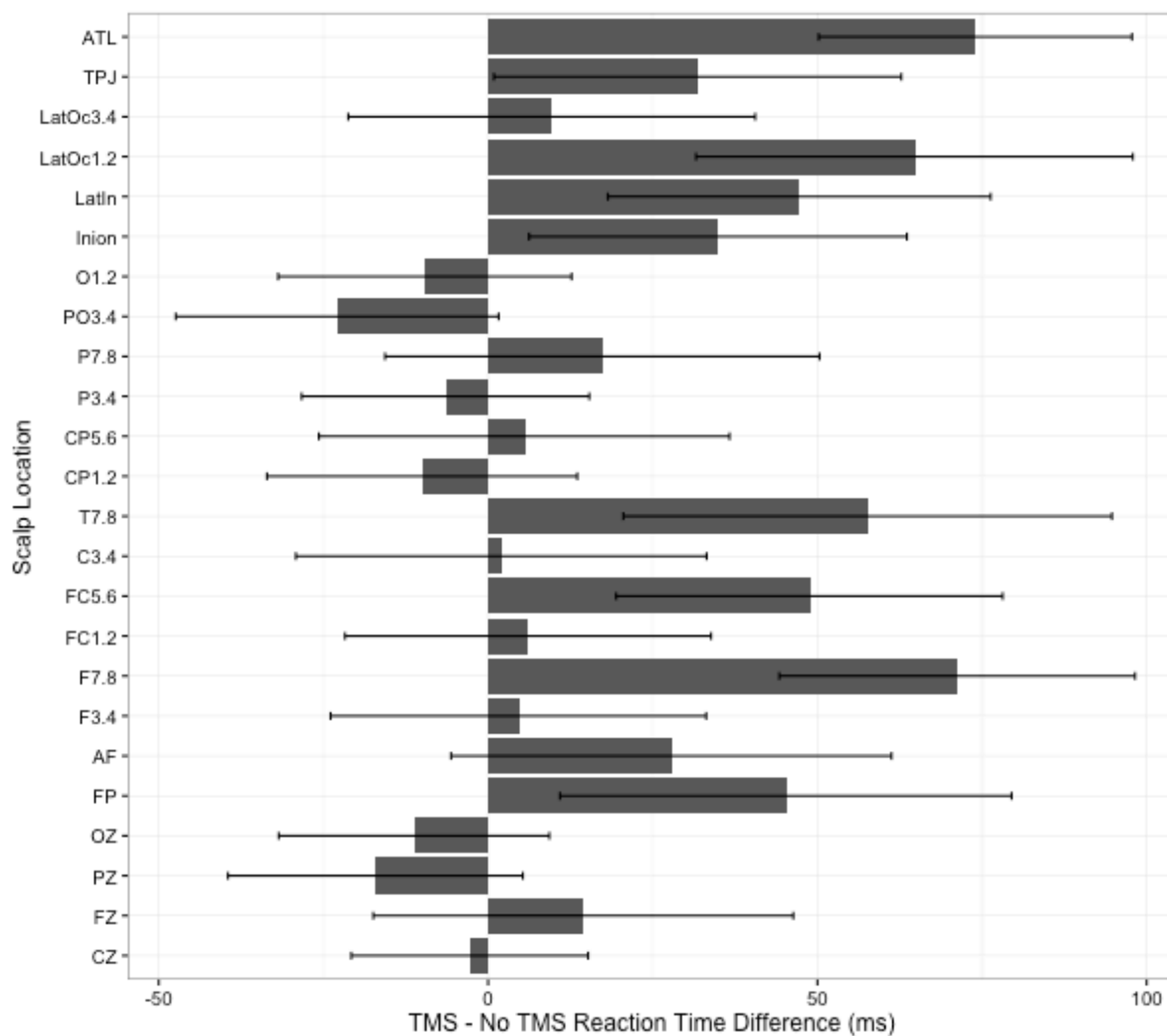
Table 1: Least squares estimated means[1] for the effect of TMS on reaction times (ms) by scalp location with approximate MNI co-ordinates (mean of left and right hemispheres, with absolute X coordinates) and scalp-cortex distance (mm)[2]

Location	Estimated mean (ms)	t-value [1]	Approximate MNI	Mean+/-SD scalp-cortex distance (mm)
CZ	-1.96	-0.14	(6,-13,99)	17.7 (16.7,18.6)
FZ	13.96	1.03	(9,44,73)	15.0 (14.0,16.0)
PZ	-16.83	-1.24	(4,-68,95)	21.2 (19.7,22.6)
OZ	11.08	-0.82	(17,-102,42)	13.6 (12.5,14.6)
FP	45.83	3.37*	(30,77,25)	13.1 (12.1,14.2)
AF	27.51	2.03	(37,64,41)	14.4 (13.3,15.4)
F3&4	4.60	0.34	(41,32,66)	14.2 (13.3,14.9)
F7&8	71.27	5.26*	(71,27,21)	13.4 (12.3,14.4)
FC1&2	6.28	0.46	(28,16,86)	15.1 (14.1,16.0)
FC5&6	48.48	3.58*	(67,5,54)	13.8 (12.9,14.7)
C3&4	2.92	0.21	(47,-18,86)	14.4 (13.3,15.4)
T7&8	56.49	4.15*	(78,-26,39)	11.6 (10.6,12.5)
CP1&2	-10.46	-0.77	(28,-48,96)	17.1 (16.0,18.1)
CP5&6	5.44	0.40	(65,-47,68)	16.3 (15.2,17.5)
P3&4	-6.25	-0.46	(39,-70,83)	18.9 (17.7,20.0)
P7&8	16.89	1.25	(60,-77,45)	13.6 (12.8,14.3)
PO3&4	-22.88	-1.69*	(37,-87,66)	16.0 (14.9,17.1)
O1&2	-9.57	-0.71	(33,-98,43)	13.4 (12.4,14.4)
Inion	35.38	2.61	(20,-115,-41)	21.4 (19.3,23.8)
Lateral-inion	47.74	3.52*	(51,-98,-46)	17.4 (15.9,19.1)
Lateral-Occipital 1&2	65.33	4.80*	(58,-66,-42)	15.3 (14.1,16.7)
Lateral Occipital 3&4	9.93	0.73	(59,-101,-11)	11.8 (10.9,12.6)
~TPJ	31.99	2.35	(47,-55,17)	11.7 (11.1,12.4)
~ATL	74.56	5.50*	(63,-14,-2)	11.4 (10.7,12.1)

Table 1 Key: * p<0.002 Bonferroni corrected p-value

Table 1 Key continued:

[1] Least squares means testing whether RT difference (TMS-No TMS) for each scalp location is significantly different to zero. Positive numbers indicate slower reaction times under TMS, negative numbers indicate faster reaction times under TMS. Least squares means have been calculated for a model that includes Scalp Location X Hemisphere, Task and Trial as predictors.

[2] Measured in a separate group of 20 participants for whom we had an MRI scan available. The brain and TMS locations used in the main experiment was registered to each of 19 other brain scans, and mean scalp-cortex distance was estimated using FLSView.

3.4 Predicting reaction times by ratings of annoyance, pain, twitches, and by visible twitches

To test whether Twitch Ratings are a better explanation for the variation in RTs across Scalp Locations, we added Twitch Ratings as a fixed effect predictor to the existing model (detailed above). This significantly improved model fit (X^2(1)= 28.81, p<0.001) and showed a significant main effect of Twitch Ratings (est. mean=14.99ms, SE=2.77, df=5411, t=5.406, p<0.0001). For every increment in rated Twitch intensity, there was an RT cost of around 15ms under TMS. We tested the interaction between Twitch Ratings and Task, to explore whether subjective discomfort differentially affected the two tasks. The interaction was significant (est. mean=7.75ms, SE=2.26ms, df=557, t=3.43, p=0.0007), with a greater cost in RTs for the Flanker task than the CRT task. The interaction term significantly improved model fit (X^2(1)=12.454, p<0.0005). To illustrate the interaction, Figure 4 plots the effect of Twitch Ratings on reaction times (ms), by task.

Figure 4: Reaction time difference (TMS – No TMS) by subjective rating of Twitch intensity for CRT and Flanker Tasks.

Legend/Key: A positive value on the y axis means slower RTs under TMS. Error bars are 95% confidence intervals.
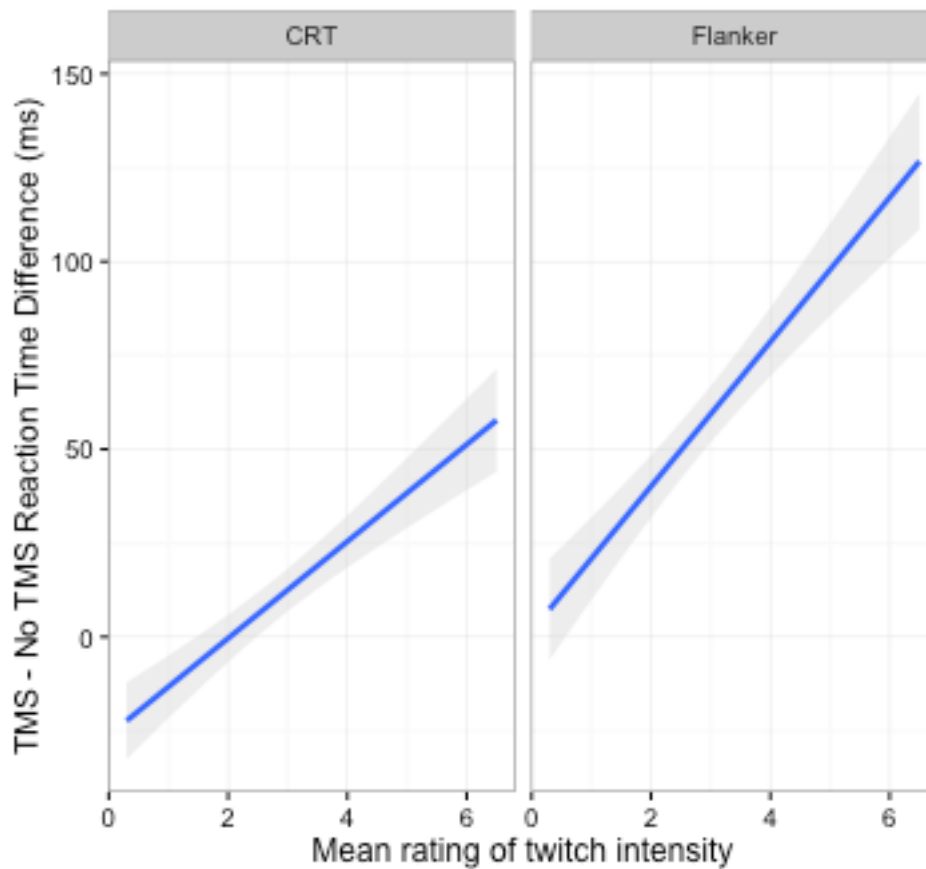
Once Twitch Rating was included in the model, significant variation in RTs as predicted by Scalp Location was substantially reduced. In this model, only one Scalp Location showed a significant difference to Cz. This was P03&4 (est. mean=-39.77, SE=18.49, df=381, t=-2.150, p<0.05), which retained significantly speeded RTs under TMS relative to Cz. All Scalp Locations which had shown slowing under TMS relative to CZ in the previous model were no longer significant. For this model, the fixed effects had an R^2 of 0.14 (effect size 0.15) and the full model (i.e., both fixed and random effects) had an R^2 of 0.43 (effect size of 0.81).

Appendix D provides details of a parallel analysis that uses scalp to brain distance instead of Scalp Location, where we found identical results. Thus, when subjective discomfort is taken into account (i.e., the inclusion of Twitch Ratings as a predictor), neither Scalp Location nor scalp to brain distance predict the cost of TMS on RTs.

To check whether individual variation in resting motor threshold (RMT) influenced the results, we added each participant's RMT as a predictor to this model, both as a main effect (i.e. does a higher or lower RMT change how much TMS influences overall RTs in the task) and with an interaction between RMT and Scalp Location (i.e. does the influence of TMS on RTs change for particular scalp locations when the participant has a higher or lower RMT?). It could be argued that individuals with lower RMTs would be more likely to experience cortical stimulation from TMS (fixed at 50% of maximum stimulator output). We might see that a lower RMT would lead to greater changes in RTs under TMS (a main effect) or particular cortical locations would show greater RT changes under TMS for individuals with lower RMT (an interaction). We found no such effects, that is, there was no significant main effect of RMT and no interaction between RMT and Scalp Location.

4.0 Discussion

When mapped across the scalp, subjective ratings and differences in RTs showed minimal changes along the midline but high interference from TMS at inferior frontal, anterior temporal, and lateral occipital sites. This confirms previous reports that TMS is more uncomfortable – and often painful – over frontal scalp locations (Abler et al., 2005; Loo et al., 2008; Machii et al., 2006; Maizey et al., 2013; Wassermann, 1998). This experiment is the first to provide data that systematically maps these effects across the scalp.

We found significant differences in how TMS affected reaction times when the pulse was applied to different scalp locations. Analyses compared each scalp location against Cz/vertex, a commonly used control site in TMS studies (e.g., Jung et al., 2016), and six locations produced longer RTs under TMS as compared to Cz/vertex. These results are in line with the argument that TMS over Cz seemingly does not affect behaviour (e.g., Jung et al., 2016; Silvanto et al., 2008), as Cz showed no substantial change in RTs when on-line TMS was applied. This makes Cz/Vertex a suitable control for other similarly benign scalp locations (e.g., FZ, FC1&2, C3&4, etc., see Table 1 in Results). However Cz/Vertex will only control for limited dimensions of TMS (e.g., the audible click, presence of the

experimenter) when compared against scalp locations that are more uncomfortable under TMS (e.g., Lateral-Occipital, ATL, T7&8).

We found that subjective ratings of annoyance, pain, and twitches were highly correlated; this is perhaps unsurprising given that participants provided these ratings consecutively. It does however support previous findings that the discomfort caused by TMS is primarily due to the effect of the TMS pulse on the skin, nerves, and muscle of the scalp (Abler et al., 2005; Arana et al., 2008; Loo et al., 2000). Once ratings of muscle twitches was added as a predictor variable, the effect of Scalp Location disappeared for all areas where TMS caused longer RTs. This means that the increase in RTs seen for many Scalp Locations under TMS is due to the fact that these locations are more uncomfortable under TMS. This finding did not change when we included scalp-to-cortex distance rather than scalp location. Abler and colleagues (2005) found a positive correlation between the number of errors on a task and subjective ratings of discomfort caused by TMS. Our data provides further support for the relationship between discomfort under TMS and changes in behavioural performance.

One question that we have been asked is "How do you know it wasn't brain stimulation that caused the differences in behavioural response times?" It is likely that cortical stimulation did occur at most TMS sites. However, the degree of cortical stimulation across different locations was, at best, poorly controlled due to varying scalp to brain distances (Stokes et al, 2005; Sack et al, 2009). When compared against other means of localising cortical sites for stimulation (i.e. fMRI guided, MRI guided and Talairach) the 10:20 anatomical system shows the smallest effect size for effects of TMS (versus sham) on RTs (Sack et al., 2009). We found no effect of scalp to brain distance and no effect of resting motor threshold on RT once subjective discomfort was accounted for. The most parsimonious explanation for the cause of the clear and consistent effects of TMS on RTs is peripheral stimulation and discomfort.

The one scalp location that showed speeded RTs under TMS (PO3&4) remained significant in the analysis that included subjective ratings. Previous studies have shown that TMS over parietal sites affects visual attention (Chambers & Heinen, 2010; Taylor & Thut, 2012), so this result could be explained by cortical stimulation broadly enhancing visual attention or orienting during the task.

Interestingly, for the parietal sites with faster RTs under TMS (PO3&4) median ratings of annoyance, pain, and twitches were close to zero. Therefore, at these sites the TMS pulse at stimulus onset may have acted as a general alerting signal that reduced RTs (Drager et al., 2004; Duecker et al., 2013; Duecker & Sack, 2013; Marzi et al., 1998; Nikouline et al., 1999; Sawaki et al., 1999). Our data is not able to distinguish between these two explanations.

The data also showed that greater discomfort (i.e., subjective ratings) produces a greater cost on task performance, and this cost differed by task. For every unit increase in annoyance there was a 14ms increase in RTs for the CRT task, and a 21ms increase in RTs for the Eriksen flanker task. Thus, more difficult tasks are affected more by the TMS-induced discomfort. Models explained ~45% of the variance in RTs. This is comparable, for example, to the size of the effect of scalp to brain distance on resting motor threshold (Stokes et al., 2007). Because of this relationship control tasks need to be carefully matched to experimental tasks in TMS experiments in order for confounds to be managed (Rossi et al., 2009). We should stress that our findings apply only to online single pulse TMS studies (i.e., the conditions that we tested), where the TMS pulse is delivered during task performance, close to stimulus presentation and the preparation and execution of a behavioural response. Whilst online and offline rTMS, paired-pulse, quadripulse, and other TMS protocols are all likely to induce similar scalp sensations, their influence on task performance needs to be determined.

We did not measure the participant's ratings of auditory or visual interference of TMS at each site (although participants' wore ear-plugs to mitigate the auditory interference). For sites close to the ears, the TMS-related sounds would have been louder and more lateralized than sites further away. For sites close to the eyes, participants' peripheral vision would have been partly occluded by the TMS coil. These variables may have explained additional variance in the data, and should be studied further.

The data we have presented systematically maps the peripheral effects of single-pulse TMS across the scalp. We cannot generalise these results to other TMS stimulation parameters, but note that studies which use bursts of TMS will tend to result in higher levels of subjective discomfort, which can lead to even more "side-effects" and annoyance. As noted above, best practice in TMS studies is to have more than one control condition for a reliable, specific effect of TMS (Sandrini et al., 2011). Our data provides a tool to select control sites for on-line TMS experiments (www.tms-smart.info) when

researchers want to control for annoyance, pain and twitches. In an ideal case, the control site will be a different brain region/scalp location, selected to match for subjective ratings (annoyance, pain, or twitches), or the effect on RTs. For example, if the experimental TMS site is over the right TPJ with the coil handle oriented approximately 45 degrees behind the inter-aural axis (SE-NW, median rating of twitches=3), suitable control sites using the same coil orientation, include the right lateral inion (7.91cm away, median twitches=3), the right lateral occipital 2 (6.81cm away, median twitches=3), or FC6 (6.80cm away, median twitches=2.5). The tools on the website allow you to customise the search for control locations using all our available data, the 10:20 system, and/or MNI co-ordinates.

Once target and control sites have been selected, we would then recommend a systematic pilot exploration of the effects of TMS intensity and coil orientation on participant-reported levels of annoyance, twitches, and pain, and on RT. We have provided an example of a systematic exploration that could be used to select appropriate TMS intensities for different sites (Appendix C). Interestingly, for scalp locations shown to be the most annoying from our initial data, the effect of TMS intensity on ratings was linear (see Appendix C). Our data provides a tool to select control sites for on-line TMS experiments (www.tms-smart.info) when researchers want to control for annoyance, pain and twitches. The tools on the website allow you to customise the search for control locations using all our available data, the 10:20 system, and/or MNI co-ordinates. We hope this data and the online resource will be useful for future TMS studies to properly control for the peripheral effects of TMS, particularly when those peripheral effects are extremely annoying.

References

Abler, B., Walter, H., Wunderlich, A., Grothe, J., Schönfeldt-Lecuona, C., Spitzer, M., & Herwig, U. (2005). Side effects of transcranial magnetic stimulation biased task performance in a cognitive neuroscience study. *Brain Topography*, *17*(4), 193-196. doi: 10.1007/s10548-005-6028-y

Anderson, B., Mishory, A., Nahas, Z., Borckardt, J. J., Yamanaka, K., Rastogi, K., & George, M. S. (2006). Tolerability and safety of high daily doses of repetitive transcranial magnetic stimulation in healthy young men. *Journal of ECT*, *22*(1), 49-53.

Arana, A. B., Borckardt, J. J., Ricci, R., Anderson, B., Li, X., Linder, K. J., Long, J. M., Sackeim, H. A., & George, M. S. (2008). Focal electrical stimulation as a sham control for repetitive transcranial magnetic stimulation: Does it truly mimic the cutaneous sensation and pain of active prefrontal repetitive transcranial magnetic stimulation? *Brain Stimulation*, *1*(1), 44-51. doi: 10.1016/j.brs.2007.08.006

Baayen, R. H., & Milin, P. (2010). Analyzing reaction times. *International Journal of Psychological Research*, *3*(2), 12-28.

Bates, D., Maechler, M., Bolker, B., & Walker, S. (2014) lme4: Linear mixed-effects models using Eigen and S4. R package version 1.1-6. http://CRAN.R-project.org/package=lme4

Barr, D. J., Levy, R., Scheepers, C., & Tily, H. J. (2013). Random effects structure for confirmatory hypothesis testing: Keep it maximal. *Journal of memory and language*, *68*(3), 255-278.

Bestmann, S., Baudewig, J., Siebner, H. R., Rothwell, J. C., & Frahm, J. (2005). BOLD MRI responses to repetitive TMS over human dorsal premotor cortex. *NeuroImage*, *28*(1), 22-29. doi: 10.1016/j.neuroimage.2005.05.027

Benninger, D. H., Berman, B. D., Houdayer, E., Pal, N., Luckenbaugh, D.A., Schneider, L., Miranda, S., & Hallett, M. (2011). Intermittent theta-burst transcranial magnetic stimulation for treatment of Parkinson disease. *Neurology*, 76(7), 601-609. doi: 10.1212/WNL.0b013e31820ce6bb

Bolognini, N., & Ro, T. (2010). Transcranial magnetic stimulation: disrupting neural activity to alter and assess brain function. *Journal of Neuroscience*, *30*(29), 9647-9650. doi: 10.1523/JNEUROSCI.1990-10.2010

Borckardt, J. J., Nahas, Z. H., Teal, J., Lisanby, S. H., McDonald, W. M., Avery, D., Durkalski, V., Pavlicova, M., Long, J. M., Sackeim, H. A., & George, M.S. (2013). The painfulness of active, but not sham, transcranial magnetic stimulation decreases rapidly over time: results from the double-blind phase of the opt-tms trial. *Brain Stimulation*, 6(6), 925-928. doi: 10.1016/j.brs.2013.04.009

Brainard, D. H. (1997). The psychophysics toolbox. *Spatial Vision*, 10(4), 433-436. doi: 10.1163/156856897X00357

Cardoso, E. F., Fregni, F., Maia, F. M., Boggio, P. S., Myczkowski, M. L., Coracini, K., Vieira, A. L., Melo, L. M., Sato, J. R., Marcolin, M. A., Rigonatti, S. P., Cruz, A. C. Jr., Barbosa, E. R., & Amaro Jr., E. (2008). rTMS treatment for depression in Parkinson's disease increases BOLD

responses in the left prefrontal cortex. *International Journal of Neuropsychopharmacology*, 11(2), 173-183. doi: *10.1017/S1461145707007961*

Champely, S. (2016). pwr: Basic Functions for Power Analysis. R package version 1.2-0. http://CRAN.R-project.org/package=pwr

Chambers, C. D., & Heinen, K. (2010). TMS and the functional neuroanatomy of attention. *cortex*, *46*(1), 114-117.

Cohen, J., Cohen, P., West, S. G., & Aiken, L. S. (2003). *Applied multiple regression/correlation analysis for the behavioral sciences*. 3rd Edition, Routledge.

Drager, B., Breitenstein, C., Helmkei U., Kamping, S., & Knecht, S. (2004). Specific and nonspecific effects of transcranial magnetic stimulation on picture-word verification. *European Journal of Neuroscience*, 20(6), 1681-1687. doi: 10.1111/j.1460-9568.2004.03623.x

Duecker, F., de Graaf, T. A., Jacobs, C., & Sack, A. T. (2013). Time-and task-dependent non-neural effects of real and sham TMS. *PloS ONE*, *8*(9), e73813. doi: 10.1371/journal.pone.0073813

Duecker, F., & Sack, A. T. (2013). Pre-stimulus sham TMS facilitates target detection. *PLoS ONE*, *8*(3), e57765. doi: 10.1371/journal.pone.0057765

Duecker, F., & Sack, A. T. (2015). Rethinking the role of sham TMS. Frontiers in psychology, 6, 210.

Eriksen, B. A., & Eriksen, C. W. (1974). Effects of noise letters upon the identification of a target in a nonsearch task. *Perception & Psychophysics*, 16(1), 143-149. doi: 10.3758/BF03203267

Fox, P., Ingham, R., George, M. S., Mayberg, H., Ingham, J., Roby, J., Martin, C., & Jerabek, P. (1997). Imaging human intra-cerebral connectivity by PET during TMS. *NeuroReport*, 8(12), 2787-2791.

de Graaf, T. A., Koivisto, M., Jacobs, C., & Sack, A. T. (2014). The chronometry of visual perception: review of occipital TMS masking studies. *Neuroscience & Biobehavioral Reviews*, *45*, 295-304. doi: 10.1016/j.neubiorev.2014.06.017

Elke, K, Schlegel, M., Sack, A. T., Nowak, D. A., Dafotakis, M., Bangard, C., Brand, M., Shamay-Tsoory, S., Onur, O. A., & Kessler, J.. (2010). Dissociating cognitive from affective theory of mind: a TMS study. *Cortex,* 46(6), 769-780. doi: 10.1016/j.cortex.2009.07.010

Herwig, U., Cardenas-Morales, L., Connemann, B. J., Kammer, T., & Schönfeldt-Lecuona, C. (2010). Sham or real—Post hoc estimation of stimulation condition in a randomized transcranial

magnetic stimulation trial. *Neuroscience Letters*, *471*(1), 30-33. doi: 10.1016/j.neulet.2010.01.003

Herwig, U., Satrapi, P., & Schönfeldt-Lecuona, C. (2003). Using the international 10-20 EEG system for positioning of transcranial magnetic stimulation. *Brain Topography*, *16*(2), 95-99.

Homan, R. W., Herman, J., & Purdy, P. (1987). Cerebral location of international 10–20 system electrode placement. *Electroencephalography and Clinical Neurophysiology*, *66*(4), 376-382.

Hothorn, T., Bretz, F., & Westfall, P. (2008). Simultaneous inference in general parametric models. *Biometrical Journal*, 50(3), 346-363. doi: 10.1002/bimj.200810425

Janicak, P. G., O'Reardon J. P., Sampson, S. M., Husain, M. M., Lisanby, S. H., Rado, J. R., Heart, K. L., & Demitrack, M. A. (2008). Transcranial magnetic stimulation in the treatment of major depressive disorder: A comprehensive summary of safety experience from acute exposure, extended exposure, and during reintroduction treatment. *Journal of Clinical Psychiatry*, 69(2), 222-232.

Jung, J., Bungert, A., Bowtell, R., & Jackson, S. R. (2016). Vertex stimulation as a control site for transcranial magnetic stimulation: A concurrent TMS/fMRI study. *Brain Stimulation*, *9*(1), 58-64. doi: 10.1016/j.brs.2015.09.008

Kuznetsova, A., Brockhoff, P. B., & Bojesen, R. H. (2014) lmerTest: Tests for random and fixed effects for linear mixed effect models (lmer objects of lme4 package). R package version 2.0-6. http://CRAN.R-project.org/package=lmerTest

Lefcheck, J. S. (2015) piecewiseSEM: Piecewise structural equation modeling in R for ecology, evolution, and systematics. Methods in Ecology and Evolution. 7(5): 573-579. DOI: 10.1111/2041-210X.12512

Lisanby, S. H., Gutman, D., Luber, B., Schroeder, C., & Sackeim, H. A. (2001). Sham TMS: intracerebral measurement of the induced electrical field and the induction of motor-evoked potentials. *Biological Psychiatry*, *49*(5), 460-463. doi: 10.1016/S0006-3223(00)01110-0

Loo, C. K., Taylor, J. L., Gandevia, S. C., McDarmont, B. N., Mitchell, P. B., & Sachdev, P. S. (2000). Transcranial magnetic stimulation (TMS) in controlled treatment studies: Are some "sham" forms active? *Biological Psychiatry*, 47(4), 325–331. doi: 10.1016/S0006-3223(99)00285-1

Machii, K., Cohen, D., Ramos-Estebanez, C., & Pascual-Leone, A. (2006). Safety of rTMS to non-motor cortical areas in healthy participants and patients. *Clinical Neurophysiology*, 117(2), 455-471. doi: 10.1016/j.clinph.2005.10.014

Maizey, L., Allen, C. P., Dervinis, M., Verbruggen, F., Varnava, A., Kozlov, M., Adams, R. C., Stokes, M., Klemen, J., Bungert, A., & Hounsell, C. A. (2013). Comparative incidence rates of mild adverse effects to transcranial magnetic stimulation. *Clinical Neurophysiology*, *124*(3), 536-544. doi: 10.1016/j.clinph.2012.07.024

Marzi, C. A., Miniussi, C., Maravita, A., Bertolasi, L., Rothwell, J. C., Sanes, J. N., & Zanette, G. (1998). Transcranial magnetic stimulation selectively impairs interhemispheric transfer of visuo-motor information in humans. *Experimental Brain Research*, *118*(3), 435-438. doi: 10.1007/s002210050299

Melzack, R., & Casey, K. L. (1968). Chapter: 20. Sensory, motivational, and central control determinants of pain: A new conceptual model. In Kenshalo, D., R. (Ed). *The Skin Senses,* pp. 423-439. Charles C Thomas, Springfield.

Miall, R. C., Christensen, L. O. D., Cain, O., & Stanley, J. (2008). Disruption of state estimation in the human lateral cerebellum. *PLoS Biology*, 5(11), e316. doi: 10.1371/journal.pbio.0050316

Nikouline, V., Ruohonen, J., & Ilmoniemi, R. J. (1999). The role of the coil click in TMS assessed with simultaneous EEG. *Clinical Neurophysiology*, 110(8), 1325-1328. doi: 10.1016/S1388-2457(99)00070-XO'Reardon, J. P., Solvason, H. B., Janicak, P. G., Sampson, S., Isenberg, K. E., Nahas, Z., McDonald, W. M., Avery, D. H., Fitzgerald, P. B., Loo, C. K., Demitrack, M. A., George, M. S., & Sackeim, H. A. (2007). Efficacy and safety of transcranial magnetic stimulation in the acute treatment of major depression: A multsite randomized controlled trial. *Biological Psychiatry*, 62(11), 1208-1216. doi: 10.1016/j.biopsych.2007.01.018

Peurala, S. H., Müller-Dahlhaus, J. F. M., Arai, N., & Ziemann, U. (2008). Interference of short-interval intracortical inhibition (SICI) and short-interval intracortical facilitation (SICF). *Clinical Neurophysiology*, 119(10):2291-2297. doi. 10.1016/j.clinph.2008.05.031

R Core Team (2014). R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria, 2012.

Ratcliff, R. (1993). Methods for dealing with reaction time outliers. *Psychological Bulletin*, *114*(3), 510. doi: 10.1037/0033-2909.114.3.510

Rossi, S., Hallett, M., Rossini, P. M., Pascual-Leone, A., & Safety of TMS Consensus Group. (2009). Safety, ethical considerations, and application guidelines for the use of transcranial magnetic stimulation in clinical practice and research. *Clinical Neurophysiology*, *120*(12), 2008-2039. doi: 10.1016/j.clinph.2009.08.016

Rossi, S., Pasqualetti, P., Zito, G., Vecchio, F., Cappa, S. F., Miniussi, C., Babiloni, C., & Rossini, P. M. (2006). Prefrontal and parietal cortex in human epsidodic memory: an interference study by repetitive transcranial magnetic stimulation. *European Journal of Neuroscience*, 23(3), 793-800. doi: 10.1111/j.1460-9568.2006.04600.x

Rossi, S., Ferro, M., Cincotta, M., Ulivelli, M., Bartalini, S., Miniussi, C., Giovannelli, F., & Passero, S. (2007). A real electro-magnetic placebo (REMP) device for sham transcranial magnetic stimulation (TMS). *Clinical Neurophysiology*, *118*(3), 709-716. doi: 10.1016/j.clinph.2006.11.005

Rossini, P. M., Barker, A. T., Berardelli, A., Caramia, M. D., Caruso, G., Cracco, R. Q., Dimitrijevic, M. R., Hallett, M., Katayama, Y., Lucking, C. H., Maertens de Nordhout, A. L., Marsden, C. D., Murray, N. M. F., Rothwell, J. C., Swash, M., & Tomberg, C. (1994). Non-invasive electrical and magnetic stimulation of the brain, spinal cord and roots: basic principles and procedures for routine clinical application. Report of an IFCN committee. *Electroencephalography and Clinical Neurophysiology*, *91*(2), 79-92.

Ruff, C. C., Driver, J., & Bestmann, S. (2009). Combining TMS and fMRI: from 'virtual lesions' to functional-network accounts of cognition. *Cortex*, *45*(9), 1043-1049. doi: 10.1016/j.cortex.2008.10.012

Sack, A. T., Kadosh, R. C., Schuhmann, T., Moerel, M., Walsh, V., & Goebel, R. (2009). Optimizing functional accuracy of TMS in cognitive studies: a comparison of methods. *Journal of Cognitive Neuroscience*, *21*(2), 207-221.

Sandrini, M., Umiltà, C., & Rusconi, E. (2011). The use of transcranial magnetic stimulation in cognitive neuroscience: a new synthesis of methodological issues. *Neuroscience & Biobehavioral Reviews*, *35*(3), 516-536. doi: 10.1016/j.neubiorev.2010.06.005

Satow, T., Mima, T., Hara, H., Oga, T., Ikeda, A., Hashimoto, N. & Shibasaki, H. (2002). Nausea as a complication of low-frequency repetitive transcranial magnetic stimulation of the posterior fossa. *Clinical Neurophysiology*, 113, 1441-1443. doi: 10.1016/S1388-2457(02)00187-6

Sawaki, L., Okita, T., Fujiwara, M., & Mizuno, K. (1999). Specific and non-specific effects of transcranial magnetic stimulation on simple and go/no-go reaction time. *Experimental Brain Research*, *127*(4), 402-408. doi: 10.1007/s002210050808

Selya, A. S., Rose, J. S., Dierker, L. C., Hedeker, D., & Mermelstein, R. J. (2012). A practical guide to calculating Cohen's f2, a measure of local effect size, from PROC MIXED. Frontiers in psychology, 3, 111.

Silvanto, J., Cattaneo, Z., Battelli, L., & Pascual-Leone, A. (2008). Baseline cortical excitability determines whether TMS disrupts or facilitates behavior. *Journal of Neurophysiology*, *99*(5), 2725-2730. doi: 10.1152/jn.01392.2007

Siniatchkin, M., Schlicke, C., Stephani, U. (2011). Transcranial magnetic stimulation reveals high test–retest reliability for phosphenes but not for suppression of visual perception. *Clinical Neurophysiology*, 122(12):2475-2481. doi: 10.1016/j.clinph.2011.05.003

Sommer, J., Jansen, A., Dräger, B., Steinsträter, O., Breitenstein, C., Deppe, M., & Knecht, S. (2006). Transcranial magnetic stimulation—a sandwich coil design for a better sham. *Clinical Neurophysiology*, *117*(2), 440-446. doi: 10.1016/j.clinph.2005.09.025

Sparing, R., & Mottaghy, F. M. (2008). Noninvasive brain stimulation with transcranial magnetic or direct current stimulation (TMS/tDCS)—from insights into human memory to therapy of its dysfunction. *Methods*, *44*(4), 329-337. doi: 10.1016/j.ymeth.2007.02.001

Stokes, M. G., Chambers, C. D., Gould, I. C., English, T., McNaught, E., McDonald, O., & Mattingley, J. B. (2007) Distance-adjusted motor threshold for transcranial magnetic stimulation. *Clinical Neurophysiology*, 118(7):1617-1625. doi: 10.1016/j.clinph.2007.04.004

Taylor, P. C. J., & Thut, G. (2012). Brain activity underlying visual perception and attention as inferred from TMS–EEG: A review. *Brain stimulation*, *5*(2), 124-129.

Tamè, L., Holmes, N. P. (2016). Involvement of human primary somatosensory cortex in vibrotactile detection depends on task demand. *NeuroImage*.

Terao, Y., Ugawa, Y., Suzuki, M., Sakai, K., Hanajima, R., Gemba-Shimizu, K. & Kanazawa, I. (1997). Shortening of simple reaction time by peripheral electrical and submotor-threshold magnetic cortical stimulation. *Experimental Brain Research*, 115, 541-545. doi: 10.1007/PL00005724

Vetter, P., Grosbras, M. H., & Muckli, L. (2015). TMS over V5 disrupts motion prediction. *Cerebral Cortex*, *25*(4), 1052-1059. doi: 10.1093/cercor/bht297

VonLoh, M., Chen, R., & Kluger, B. (2013). Safety of transcranial magnetic stimulation in Parkinson's disease: A review of the literature. *Parkinsonism and Related Disorders*, 19, 573-585. doi: 10.1016/j.parkreldis.2013.01.007

Walsh, V., & Cowey, A. (2000). Transcranial magnetic stimulation and cognitive neuroscience. *Nature Reviews Neuroscience*, *1*(1), 73-80. doi: 10.1038/35036239

Wagner, T., Valero-Cabre, A., & Pascual-Leone, A. (2007). Noninvasive human brain stimulation. *Annual Review of Biomedical Engineering*, *9*, 527-565. doi: 10.1146/annurev.bioeng.9.061206.133100

Wassermann, E.M. (1998). Risk and safety of repetitive transcranial magnetic stimulation: report and suggested guidelines from the International Workshop on the Safety of Repetitive Trancranial Magnetic Stimulation, June 5-7,1996. *Electroencephalography and Clinical Neurophysiology*, 108(1), 1-16.

Westin GG, Bassi BD, Lisanby SH, Luber B (2014) Determination of motor threshold using visual observation overestimates transcranial magnetic stimulation dosage: safety implications. Clinical Neurophysiology, 125(1):142-147. doi: 10.1016/j.clinph.2013.06.187

Wickham, R. (2009). ggplot2: elegant graphics for data analysis. Springer New York, 2009.

Wei, T (2013) corrplot: Visualisation of a correlation matrix. R package version 0.73, https://github.com/taiyun/corrplot

Appendix A

Table A1: Task by participant summary of mean RT, proportion correct and congruency effects.

Task	Participant	Mean correct RT (ms)	Proportion correct	Congruency effect (Incongruent – Congruent, ms)
Flanker	1	437	0.99	14.6
	2	428	0.91	49.9
	3	563	0.99	44.7
	4	644	0.99	42.1
	5	578	0.99	-6.2
	6	467	0.97	37.0
	7	558	1.00	13.5
	8	592	0.96	44.5
	9	622	1.00	40.2
	10	604	0.97	49.4
Mean (SD)		**549 (77.6)**	**0.98 (0.03)**	**33.0 (18.9)**
CRT	11	412	0.96	12.1
	12	330	1.00	3.8
	13	295	0.86	3.7
	14	417	0.98	18.4
	15	385	0.98	5.5
	16*	629	0.99	20.8
	17	286	0.98	-3.3
	18	494	0.99	26.4
	19*	431	0.99	23.9
	20*	485	0.99	50.5
Mean (SD)		**416 (103)**	**0.97 (0.04)**	**16.2 (15.6)**

* Participants who responded to arrow direction rather than side of presentation, data were recoded to reflect their responses.

Appendix B

Multiple comparisons for all scalp locations.

Adjusted P values using Tukey HSD to control for family wise error rate.
For each comparison, the null hypothesis is that the difference between the Test Location and Reference is 0.
A positive estimate means that the Test Location shows a greater cost of TMS (i.e. longer RTs under TMS) than the Reference. A negative estimate means that the Test Location shows less cost (i.e. shorter RTs under TMS) than the Reference.

ADJUSTED P VALUES						
Test Location	**Reference for comparison**	Estimate	Std Error	z-value	Pr(>|z|)	Sig. codes: 0 = *** 0.001 = ** 0.01 = * 0.05 = .
FC5.6	CZ	44.8541	20.1587	2.225	0.8585	
Inion	CZ	13.2431	20.2043	0.655	1	
LatIn	CZ	35.6462	20.1804	1.766	0.9873	
LatOc	CZ	55.2387	20.2466	2.728	0.4928	
V5	CZ	-2.9647	20.1519	-0.147	1	
TPJ	CZ	13.9263	20.1827	0.69	1	
ATL	CZ	71.6026	20.1584	3.552	0.0647	.
C3.4	CZ	-20.3417	20.2203	-1.006	1	
T7.8	CZ	40.288	20.2311	1.991	0.949	
CP1.2	CZ	-16.2257	20.1627	-0.805	1	
CP5.6	CZ	-21.3591	20.168	-1.059	1	
P3.4	CZ	-16.4929	20.1526	-0.818	1	
P7.8	CZ	-7.7011	20.1813	-0.382	1	
PO3.4	CZ	-41.1535	20.1698	-2.04	0.9358	
O1.2	CZ	-23.8884	20.1737	-1.184	1	
FP	CZ	42.5712	20.2215	2.105	0.9129	
FZ	CZ	-8.6627	20.2111	-0.429	1	
AF	CZ	22.5426	20.185	1.117	1	
F3.4	CZ	6.1978	20.1776	0.307	1	
F7.8	CZ	62.4987	20.2189	3.091	0.2381	
FC1.2	CZ	-20.0655	20.1582	-0.995	1	
PZ	CZ	-20.8473	20.1747	-1.033	1	
OZ	CZ	-23.5049	20.157	-1.166	1	
Inion	FC5.6	-31.6111	20.1877	-1.566	0.9974	
LatIn	FC5.6	-9.2079	20.1642	-0.457	1	
LatOc	FC5.6	10.3845	20.2299	0.513	1	
V5	FC5.6	-47.8188	20.1359	-2.375	0.7682	
TPJ	FC5.6	-30.9278	20.1668	-1.534	0.9981	

ATL	FC5.6	26.7485	20.142	1.328	0.9998	
C3.4	FC5.6	-65.1958	20.2027	-3.227	0.1715	
T7.8	FC5.6	-4.5661	20.2147	-0.226	1	
CP1.2	FC5.6	-61.0799	20.1458	-3.032	0.275	
CP5.6	FC5.6	-66.2132	20.1514	-3.286	0.1456	
P3.4	FC5.6	-61.3471	20.1358	-3.047	0.2653	
P7.8	FC5.6	-52.5552	20.1649	-2.606	0.5932	
PO3.4	FC5.6	-86.0076	20.1526	-4.268	<0.01	**
O1.2	FC5.6	-68.7425	20.1568	-3.41	0.1023	
FP	FC5.6	-2.2829	20.2058	-0.113	1	
FZ	FC5.6	-53.5168	20.1937	-2.65	0.5569	
AF	FC5.6	-22.3115	20.1684	-1.106	1	
F3.4	FC5.6	-38.6563	20.1616	-1.917	0.9665	
F7.8	FC5.6	17.6446	20.2022	0.873	1	
FC1.2	FC5.6	-64.9197	20.1417	-3.223	0.1738	
PZ	FC5.6	-65.7014	20.1578	-3.259	0.1559	
OZ	FC5.6	-68.359	20.1404	-3.394	0.1078	
LatIn	Inion	22.4031	20.2097	1.109	1	
LatOc	Inion	41.9956	20.2772	2.071	0.9252	
V5	Inion	-16.2077	20.1818	-0.803	1	
TPJ	Inion	0.6832	20.2122	0.034	1	
ATL	Inion	58.3595	20.1877	2.891	0.3692	
C3.4	Inion	-33.5848	20.2481	-1.659	0.9942	
T7.8	Inion	27.045	20.2601	1.335	0.9998	
CP1.2	Inion	-29.4688	20.1915	-1.459	0.9991	
CP5.6	Inion	-34.6022	20.1969	-1.713	0.9913	
P3.4	Inion	-29.736	20.1821	-1.473	0.999	
P7.8	Inion	-20.9442	20.2109	-1.036	1	
PO3.4	Inion	-54.3966	20.1986	-2.693	0.5215	
O1.2	Inion	-37.1315	20.203	-1.838	0.9792	
FP	Inion	29.3282	20.2506	1.448	0.9992	
FZ	Inion	-21.9058	20.2401	-1.082	1	
AF	Inion	9.2995	20.2145	0.46	1	
F3.4	Inion	-7.0453	20.2069	-0.349	1	
F7.8	Inion	49.2557	20.2481	2.433	0.7278	
FC1.2	Inion	-33.3086	20.1868	-1.65	0.9947	
PZ	Inion	-34.0904	20.2034	-1.687	0.9928	
OZ	Inion	-36.748	20.1859	-1.82	0.9814	
LatOc	LatIn	19.5924	20.2527	0.967	1	
V5	LatIn	-38.6109	20.1581	-1.915	0.9662	
TPJ	LatIn	-21.7199	20.1885	-1.076	1	
ATL	LatIn	35.9564	20.1644	1.783	0.9854	
C3.4	LatIn	-55.9879	20.2259	-2.768	0.4606	
T7.8	LatIn	4.6418	20.2373	0.229	1	
CP1.2	LatIn	-51.872	20.1684	-2.572	0.6191	
CP5.6	LatIn	-57.0053	20.1741	-2.826	0.4153	

P3.4	LatIn	-52.1391	20.1586	-2.586	0.6073	
P7.8	LatIn	-43.3473	20.1874	-2.147	0.8946	
PO3.4	LatIn	-76.7997	20.1753	-3.807	0.0278	*
O1.2	LatIn	-59.5346	20.1796	-2.95	0.3267	
FP	LatIn	6.925	20.2276	0.342	1	
FZ	LatIn	-44.3089	20.2161	-2.192	0.8757	
AF	LatIn	-13.1036	20.191	-0.649	1	
F3.4	LatIn	-29.4484	20.1831	-1.459	0.9991	
F7.8	LatIn	26.8525	20.2245	1.328	0.9998	
FC1.2	LatIn	-55.7118	20.1639	-2.763	0.4669	
PZ	LatIn	-56.4935	20.1803	-2.799	0.4352	
OZ	LatIn	-59.1511	20.1628	-2.934	0.3396	
V5	LatOc	-58.2033	20.2249	-2.878	0.3782	
TPJ	LatOc	-41.3123	20.2564	-2.039	0.9359	
ATL	LatOc	16.364	20.2312	0.809	1	
C3.4	LatOc	-75.5803	20.2919	-3.725	0.0374	*
T7.8	LatOc	-14.9506	20.3028	-0.736	1	
CP1.2	LatOc	-71.4644	20.2348	-3.532	0.0701	.
CP5.6	LatOc	-76.5978	20.2412	-3.784	0.0294	*
P3.4	LatOc	-71.7316	20.2247	-3.547	0.0665	.
P7.8	LatOc	-62.9398	20.2539	-3.108	0.2289	
PO3.4	LatOc	-96.3922	20.2407	-4.762	<0.01	***
O1.2	LatOc	-79.127	20.2462	-3.908	0.0183	*
FP	LatOc	-12.6674	20.2938	-0.624	1	
FZ	LatOc	-63.9013	20.2822	-3.151	0.2078	
AF	LatOc	-32.696	20.2566	-1.614	0.9961	
F3.4	LatOc	-49.0409	20.2497	-2.422	0.7342	
F7.8	LatOc	7.2601	20.2914	0.358	1	
FC1.2	LatOc	-75.3042	20.2307	-3.722	0.0379	*
PZ	LatOc	-76.0859	20.2473	-3.758	0.0308	*
OZ	LatOc	-78.7436	20.2292	-3.893	0.0201	*
TPJ	V5	16.891	20.1607	0.838	1	
ATL	V5	74.5673	20.1365	3.703	0.039	*
C3.4	V5	-17.377	20.1978	-0.86	1	
T7.8	V5	43.2527	20.2093	2.14	0.8988	
CP1.2	V5	-13.2611	20.1399	-0.658	1	
CP5.6	V5	-18.3944	20.1456	-0.913	1	
P3.4	V5	-13.5283	20.1304	-0.672	1	
P7.8	V5	-4.7365	20.1591	-0.235	1	
PO3.4	V5	-38.1888	20.1472	-1.895	0.9705	
O1.2	V5	-20.9237	20.1511	-1.038	1	
FP	V5	45.5359	20.1998	2.254	0.8424	
FZ	V5	-5.698	20.1886	-0.282	1	
AF	V5	25.5073	20.163	1.265	0.9999	
F3.4	V5	9.1624	20.1557	0.455	1	
F7.8	V5	65.4634	20.1965	3.241	0.163	

FC1.2	V5	-17.1009	20.136	-0.849	1	
PZ	V5	-17.8826	20.1523	-0.887	1	
OZ	V5	-20.5402	20.1346	-1.02	1	
ATL	TPJ	57.6763	20.1661	2.86	0.3892	
C3.4	TPJ	-34.268	20.2277	-1.694	0.9924	
T7.8	TPJ	26.3617	20.2391	1.303	0.9999	
CP1.2	TPJ	-30.152	20.1706	-1.495	0.9987	
CP5.6	TPJ	-35.2854	20.1759	-1.749	0.9888	
P3.4	TPJ	-30.4192	20.1609	-1.509	0.9985	
P7.8	TPJ	-21.6274	20.1896	-1.071	1	
PO3.4	TPJ	-55.0798	20.1779	-2.73	0.4959	
O1.2	TPJ	-37.8147	20.1818	-1.874	0.974	
FP	TPJ	28.6449	20.23	1.416	0.9994	
FZ	TPJ	-22.589	20.2197	-1.117	1	
AF	TPJ	8.6163	20.1935	0.427	1	
F3.4	TPJ	-7.7285	20.1857	-0.383	1	
F7.8	TPJ	48.5724	20.2263	2.401	0.7487	
FC1.2	TPJ	-33.9918	20.1663	-1.686	0.993	
PZ	TPJ	-34.7736	20.1827	-1.723	0.9906	
OZ	TPJ	-37.4312	20.1648	-1.856	0.9761	
C3.4	ATL	-91.9443	20.2032	-4.551	<0.01	**
T7.8	ATL	-31.3146	20.2145	-1.549	0.9978	
CP1.2	ATL	-87.8284	20.1463	-4.36	<0.01	**
CP5.6	ATL	-92.9617	20.1517	-4.613	<0.01	**
P3.4	ATL	-88.0955	20.1362	-4.375	<0.01	**
P7.8	ATL	-79.3037	20.1649	-3.933	0.0171	*
PO3.4	ATL	-112.7561	20.1535	-5.595	<0.01	***
O1.2	ATL	-95.491	20.1573	-4.737	<0.01	***
FP	ATL	-29.0314	20.2059	-1.437	0.9993	
FZ	ATL	-80.2653	20.1946	-3.975	0.0149	*
AF	ATL	-49.06	20.1687	-2.432	0.7266	
F3.4	ATL	-65.4048	20.1609	-3.244	0.1629	
F7.8	ATL	-9.1039	20.2022	-0.451	1	
FC1.2	ATL	-91.6681	20.1417	-4.551	<0.01	**
PZ	ATL	-92.4499	20.1585	-4.586	<0.01	**
OZ	ATL	-95.1075	20.1406	-4.722	<0.01	***
T7.8	C3.4	60.6297	20.2743	2.99	0.2982	
CP1.2	C3.4	4.1159	20.2072	0.204	1	
CP5.6	C3.4	-1.0174	20.2129	-0.05	1	
P3.4	C3.4	3.8488	20.1975	0.191	1	
P7.8	C3.4	12.6406	20.2265	0.625	1	
PO3.4	C3.4	-20.8118	20.2143	-1.03	1	
O1.2	C3.4	-3.5467	20.2186	-0.175	1	
FP	C3.4	62.9129	20.2673	3.104	0.232	
FZ	C3.4	11.679	20.2558	0.577	1	
AF	C3.4	42.8843	20.229	2.12	0.9071	

F3.4	C3.4	26.5395	20.2227	1.312	0.9998	
F7.8	C3.4	82.8404	20.2629	4.088	<0.01	**
FC1.2	C3.4	0.2761	20.2032	0.014	1	
PZ	C3.4	-0.5056	20.2194	-0.025	1	
OZ	C3.4	-3.1632	20.2015	-0.157	1	
CP1.2	T7.8	-56.5138	20.2194	-2.795	0.4394	
CP5.6	T7.8	-61.6471	20.2253	-3.048	0.2648	
P3.4	T7.8	-56.781	20.2092	-2.81	0.4306	
P7.8	T7.8	-47.9891	20.2389	-2.371	0.7698	
PO3.4	T7.8	-81.4415	20.226	-4.027	0.0119	*
O1.2	T7.8	-64.1764	20.2301	-3.172	0.1964	
FP	T7.8	2.2832	20.2781	0.113	1	
FZ	T7.8	-48.9507	20.2675	-2.415	0.739	
AF	T7.8	-17.7454	20.2418	-0.877	1	
F3.4	T7.8	-34.0902	20.2342	-1.685	0.993	
F7.8	T7.8	22.2107	20.2754	1.095	1	
FC1.2	T7.8	-60.3536	20.2147	-2.986	0.3051	
PZ	T7.8	-61.1353	20.2311	-3.022	0.2805	
OZ	T7.8	-63.7929	20.2134	-3.156	0.2034	
CP5.6	CP1.2	-5.1334	20.1553	-0.255	1	
P3.4	CP1.2	-0.2672	20.1398	-0.013	1	
P7.8	CP1.2	8.5246	20.1691	0.423	1	
PO3.4	CP1.2	-24.9278	20.1568	-1.237	0.9999	
O1.2	CP1.2	-7.6626	20.161	-0.38	1	
FP	CP1.2	58.797	20.21	2.909	0.355	
FZ	CP1.2	7.5631	20.1984	0.374	1	
AF	CP1.2	38.7683	20.1724	1.922	0.9648	
F3.4	CP1.2	22.4235	20.1655	1.112	1	
F7.8	CP1.2	78.7245	20.2057	3.896	0.0205	*
FC1.2	CP1.2	-3.8398	20.1457	-0.191	1	
PZ	CP1.2	-4.6216	20.1624	-0.229	1	
OZ	CP1.2	-7.2792	20.1441	-0.361	1	
P3.4	CP5.6	4.8662	20.1458	0.242	1	
P7.8	CP5.6	13.658	20.1747	0.677	1	
PO3.4	CP5.6	-19.7944	20.1626	-0.982	1	
O1.2	CP5.6	-2.5293	20.1668	-0.125	1	
FP	CP5.6	63.9303	20.2151	3.162	0.201	
FZ	CP5.6	12.6964	20.2038	0.628	1	
AF	CP5.6	43.9017	20.1784	2.176	0.8822	
F3.4	CP5.6	27.5569	20.1712	1.366	0.9997	
F7.8	CP5.6	83.8578	20.2112	4.149	<0.01	**
FC1.2	CP5.6	1.2936	20.151	0.064	1	
PZ	CP5.6	0.5118	20.1677	0.025	1	
OZ	CP5.6	-2.1458	20.1499	-0.106	1	
P7.8	P3.4	8.7918	20.1591	0.436	1	
PO3.4	P3.4	-24.6606	20.1473	-1.224	0.9999	

O1.2	P3.4	-7.3955	20.1512	-0.367	1	
FP	P3.4	59.0642	20.1998	2.924	0.3449	
FZ	P3.4	7.8302	20.1884	0.388	1	
AF	P3.4	39.0355	20.1626	1.936	0.9619	
F3.4	P3.4	22.6907	20.1554	1.126	1	
F7.8	P3.4	78.9917	20.1964	3.911	0.0186	*
FC1.2	P3.4	-3.5726	20.1357	-0.177	1	
PZ	P3.4	-4.3544	20.1522	-0.216	1	
OZ	P3.4	-7.012	20.1344	-0.348	1	
PO3.4	P7.8	-33.4524	20.1763	-1.658	0.9944	
O1.2	P7.8	-16.1873	20.1806	-0.802	1	
FP	P7.8	50.2724	20.2292	2.485	0.6876	
FZ	P7.8	-0.9616	20.2177	-0.048	1	
AF	P7.8	30.2437	20.1917	1.498	0.9986	
F3.4	P7.8	13.8989	20.1839	0.689	1	
F7.8	P7.8	70.1999	20.2257	3.471	0.0847	.
FC1.2	P7.8	-12.3644	20.1645	-0.613	1	
PZ	P7.8	-13.1462	20.1816	-0.651	1	
OZ	P7.8	-15.8038	20.1635	-0.784	1	
O1.2	PO3.4	17.2651	20.1684	0.856	1	
FP	PO3.4	83.7247	20.2163	4.141	<0.01	**
FZ	PO3.4	32.4908	20.2057	1.608	0.9963	
AF	PO3.4	63.6961	20.1797	3.156	0.2053	
F3.4	PO3.4	47.3513	20.1725	2.347	0.7862	
F7.8	PO3.4	103.6523	20.214	5.128	<0.01	***
FC1.2	PO3.4	21.088	20.1531	1.046	1	
PZ	PO3.4	20.3062	20.1693	1.007	1	
OZ	PO3.4	17.6486	20.1512	0.876	1	
FP	O1.2	66.4596	20.2207	3.287	0.1418	
FZ	O1.2	15.2257	20.2096	0.753	1	
AF	O1.2	46.431	20.1842	2.3	0.815	
F3.4	O1.2	30.0862	20.1766	1.491	0.9988	
F7.8	O1.2	86.3871	20.2159	4.273	<0.01	**
FC1.2	O1.2	3.8229	20.1569	0.19	1	
PZ	O1.2	3.0411	20.1732	0.151	1	
OZ	O1.2	0.3835	20.1555	0.019	1	
FZ	FP	-51.2339	20.2579	-2.529	0.6551	
AF	FP	-20.0286	20.2321	-0.99	1	
F3.4	FP	-36.3735	20.2251	-1.798	0.9841	
F7.8	FP	19.9275	20.2658	0.983	1	
FC1.2	FP	-62.6368	20.205	-3.1	0.2316	
PZ	FP	-63.4185	20.2214	-3.136	0.2141	
OZ	FP	-66.0762	20.2036	-3.271	0.1525	
AF	FZ	31.2053	20.2208	1.543	0.9979	
F3.4	FZ	14.8605	20.2142	0.735	1	
F7.8	FZ	71.1614	20.2548	3.513	0.0722	.

FC1.2	FZ	-11.4028	20.1941	-0.565	1	
PZ	FZ	-12.1846	20.2105	-0.603	1	
OZ	FZ	-14.8422	20.1929	-0.735	1	
F3.4	AF	-16.3448	20.188	-0.81	1	
F7.8	AF	39.9561	20.229	1.975	0.9536	
FC1.2	AF	-42.6081	20.1682	-2.113	0.9108	
PZ	AF	-43.3899	20.1849	-2.15	0.8942	
OZ	AF	-46.0475	20.1669	-2.283	0.8259	
F7.8	F3.4	56.301	20.2213	2.784	0.449	
FC1.2	F3.4	-26.2633	20.1608	-1.303	0.9998	
PZ	F3.4	-27.0451	20.1777	-1.34	0.9998	
OZ	F3.4	-29.7027	20.1599	-1.473	0.999	
FC1.2	F7.8	-82.5643	20.202	-4.087	<0.01	**
PZ	F7.8	-83.346	20.2185	-4.122	<0.01	**
OZ	F7.8	-86.0037	20.2008	-4.257	<0.01	**
PZ	FC1.2	-0.7818	20.1579	-0.039	1	
OZ	FC1.2	-3.4394	20.1399	-0.171	1	
OZ	PZ	-2.6576	20.1566	-0.132	1	

Appendix B

Least squares means for all scalp locations and scalp locations x hemisphere.

Uncorrected p values and Bonferroni correction are reported.
For each comparison, the null hypothesis is that the value for each Scalp Location is 0.
A positive estimate means that the Scalp Location shows a greater cost of TMS (i.e. longer RTs under TMS). A negative estimate means that the Scalp Location shows an advantage under TMS (i.e. shorter RTs under TMS).

Scalp Location	Estimate	St. Err	DF	t-value	Lower CI	Upper CI	Uncorrected p value	Sig. with Bonferroni corrected p < 0.002
CZ	-1.955	13.538	73.3	-0.14	-28.934	25.02	0.886	
FC5.6	48.482	13.557	73.8	3.58	21.467	75.5	6.00E-04	*
Inion	35.381	13.553	73.7	2.61	8.374	62.39	0.011	
LatIn	47.743	13.55	73.6	3.52	20.741	74.74	7.00E-04	*
LatOc	65.332	13.604	74.8	4.8	38.229	92.43	2E-15	*
V5	9.928	13.547	73.5	0.73	-17.067	36.92	0.466	
TPJ	31.986	13.585	74.4	2.35	4.918	59.05	0.021	
ATL	74.564	13.552	73.7	5.5	47.559	101.57	2E-15	*
C3.4	2.917	13.581	74.3	0.21	-24.143	29.98	0.831	
T7.8	56.49	13.623	75.2	4.15	29.353	83.63	1.00E-04	*
CP1.2	-10.46	13.536	73.3	-0.77	-37.435	16.52	0.442	
CP5.6	5.44	13.548	73.6	0.4	-21.557	32.44	0.689	
P3.4	-6.252	13.533	73.2	-0.46	-33.222	20.72	0.645	
P7.8	16.892	13.552	73.6	1.25	-10.112	43.9	0.216	
PO3.4	-22.888	13.532	73.2	-1.69	-49.857	4.08	0.095	
O1.2	-9.569	13.53	73.2	-0.71	-36.533	17.39	0.482	
FP	45.827	13.603	74.8	3.37	18.727	72.93	0.001	*

	Estimate	St. Err	DF	t-value	Lower CI	Upper CI	Uncorrected p value	
FZ	13.96	13.578	74.2	1.03	-13.094	41.01	0.307	
AF	27.506	13.547	73.5	2.03	0.511	54.5	0.046	
F3.4	4.604	13.553	73.7	0.34	-22.403	31.61	0.735	
F7.8	71.272	13.562	73.9	5.26	44.248	98.3	2E-15	*
FC1.2	6.28	13.542	73.4	0.46	-20.707	33.27	0.644	
PZ	-16.83	13.533	73.2	-1.24	-43.799	10.14	0.218	
OZ	-11.081	13.531	73.2	-0.82	-38.047	15.89	0.415	

Scalp Location x Hemisphere	Hemisphere	Estimate	St. Err	DF	t-value	Lower CI	Upper CI	Uncorrected p value	Sig. with Bonferroni corrected p < 0.001
CZ	Left	-4.112	19.149	73.4	-0.21	-42.273	34.05	0.831	
FC5.6	Left	40.742	19.132	73.1	2.13	2.614	78.87	0.037	
Inion	Left	9.131	19.18	73.9	0.48	-29.087	47.35	0.635	
LatIn	Left	31.535	19.156	73.5	1.65	-6.638	69.71	0.104	
LatOc	Left	51.127	19.226	74.6	2.66	12.824	89.43	0.01	
V5	Left	-7.076	19.126	73	-0.37	-45.194	31.04	0.713	
TPJ	Left	9.815	19.158	73.5	0.51	-28.362	47.99	0.61	
ATL	Left	67.491	19.132	73.1	3.53	29.362	105.62	7.00E-04	*
C3.4	Left	-24.453	19.197	74.1	-1.27	-62.702	13.8	0.207	
T7.8	Left	36.176	19.209	74.3	1.88	-2.096	74.45	0.064	
CP1.2	Left	-20.337	19.136	73.2	-1.06	-58.474	17.8	0.291	
CP5.6	Left	-25.471	19.142	73.3	-1.33	-63.618	12.68	0.187	
P3.4	Left	-20.605	19.126	73	-1.08	-58.722	17.51	0.285	
P7.8	Left	-11.813	19.156	73.5	-0.62	-49.987	26.36	0.539	
PO3.4	Left	-45.265	19.144	73.3	-2.36	-83.416	-7.11	0.021	

O1.2	Left	-28	19.148	73.4	-1.46	-66.159	10.16	0.148	
FP	Left	38.46	19.199	74.2	2	0.206	76.71	0.049	
FZ	Left	-12.774	19.187	74	-0.67	-51.006	25.46	0.508	
AF	Left	18.431	19.16	73.6	0.96	-19.75	56.61	0.339	
F3.4	Left	2.086	19.152	73.5	0.11	-36.081	40.25	0.914	
F7.8	Left	58.387	19.196	74.1	3.04	20.14	96.63	0.003	
FC1.2	Left	-24.177	19.132	73.1	-1.26	-62.306	13.95	0.21	
PZ	Left	-24.959	19.149	73.4	-1.3	-63.12	13.2	0.197	
OZ	Left	-27.617	19.13	73.1	-1.44	-65.742	10.51	0.153	
CZ	Right	0.202	19.142	73.3	0.01	-37.944	38.35	0.992	
FC5.6	Right	56.221	19.214	74.4	2.93	17.94	94.5	0.004	
Inion	Right	61.63	19.154	73.5	3.22	23.461	99.8	0.002	
LatIn	Right	63.951	19.17	73.7	3.34	25.752	102.15	0.001	*
LatOc	Right	79.537	19.253	75	4.13	41.182	117.89	1.00E-04	*
V5	Right	26.933	19.191	74	1.4	-11.305	65.17	0.165	
TPJ	Right	54.157	19.268	75.2	2.81	15.775	92.54	0.006	
ATL	Right	81.638	19.2	74.2	4.25	43.383	119.89	1.00E-04	*
C3.4	Right	30.287	19.217	74.4	1.58	-8	68.57	0.119	
T7.8	Right	76.804	19.322	76	3.97	38.321	115.29	2.00E-04	*
CP1.2	Right	-0.582	19.149	73.4	-0.03	-38.742	37.58	0.976	
CP5.6	Right	36.351	19.177	73.8	1.9	-1.86	74.56	0.062	
P3.4	Right	8.1	19.151	73.4	0.42	-30.065	46.26	0.674	
P7.8	Right	45.597	19.174	73.8	2.38	7.391	83.8	0.02	
PO3.4	Right	-0.511	19.132	73.1	-0.03	-38.64	37.62	0.979	
O1.2	Right	8.862	19.121	73	0.46	-29.246	46.97	0.644	
FP	Right	53.194	19.275	75.3	2.76	14.798	91.59	0.007	
FZ	Right	40.694	19.218	74.4	2.12	2.406	78.98	0.038	
AF	Right	36.582	19.156	73.5	1.91	-1.592	74.76	0.06	

F3.4	Right	7.121	19.18	73.9	0.37	-31.098	45.34	0.712	
F7.8	Right	84.157	19.164	73.6	4.39	45.969	122.35	2E-15	*
FC1.2	Right	36.737	19.172	73.8	1.92	-1.466	74.94	0.059	
PZ	Right	-8.701	19.127	73.1	-0.45	-46.821	29.42	0.65	
OZ	Right	5.455	19.142	73.3	0.28	-32.691	43.6	0.776	

Appendix C / Supplementary materials 3: Effect of varying TMS intensity on ratings of annoyance, pain, and muscle twitches

Following reviewers' comments on an earlier version of this manuscript, we studied, in one volunteer, the effect of varying TMS intensity on subjective ratings of annoyance, pain, and twitches, and on visible twitches. This approach could be used to further refine the choice of control site for an experiment.

Methods

From the original dataset, we chose 6 scalp locations that spanned the full range of median twitch ratings (Table C1).

Table C1: Sites chosen for control study of TMS intensity

		Median twitch rating (n=10 per hemisphere)	
Rank	Site	Left	Right
1	ATL1/2	6.4	5.5
5	TPJ1/2	3.6	2.8
10	AF3/4	2.4	1.3
14	C3/4	1.1	1.5
19	Cz	.1	
23	CP1/2	0	0

One participant who had taken part in the main experiment was recruited. The participant received 5 single pulses of biphasic TMS at ~0.2Hz at each location (in both left and right hemispheres for all sites except Cz), with the handle of the coil pointing South. After each set of 5 TMS pulses, the participant rated the subjective annoyance, pain, and twitches just as in the main experiment. An experimenter observed the participant, and noted down how many of the trials were accompanied by observable muscle twitches in the face, scalp, neck, or body of the participant.

After each set of 5 trials, the TMS coil was repositioned and the intensity of the TMS was changed according to a fully-randomised sequence. Five TMS intensities were used: 30, 40, 50, 60, and 70% of the maximum stimulator output. The mean ratings as a function of TMS intensity are shown in

Figures C1 (mean across all 11 sites for 4 different ratings) and C2 (mean across left and right hemispheres for each site, excluding Cz).

Results & Discussion

For each variable averaged across TMS sites, mean ratings increased linearly as a function of TMS intensity – r^2s ranged between 0.90 (twitches) to 0.99 (visible twitches), all two-tailed ps<.05. The effect of TMS intensity was also approximately linear for ATL1/2 (r(3)=.912, p=.031), AF1/2 (r(3)=.982, p=.0029), TPJ1/2 (r(3)=.979, p=.0036, CP1/2 and Cz (both r(3)=.884, p=.047), but not for C3/4 (r(3)=.707, p=.18.

Data like this could be used, for example, to set TMS intensity for the control site to produce a comparable level of annoyance or muscle twitches to that of the target site (e.g., if control site cannot be varied freely). Further, these data support the assumption of a linear effect of TMS intensity on subjective annoyance, pain, and twitches, although for sites associated with low annoyance (midline and/or superior scalp locations), there may be an additional threshold of TMS intensity under which TMS is not at all annoying.
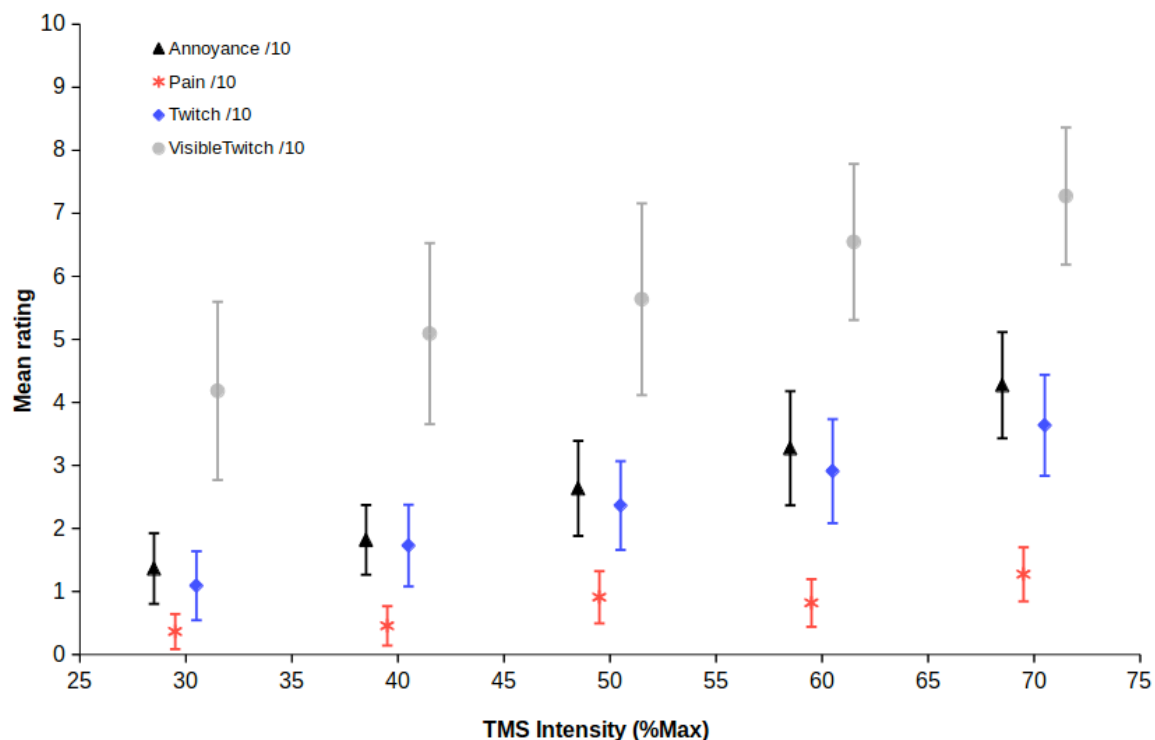
Figures

Figure C1: Mean across TMS sites as a function of TMS intensity (%Max) for subjective ratings of annoyance (black triangles), pain (red asterisks), and muscle twitches (blue diamonds), as well as observed twitches (grey circles). Visible twitches have been multiplied by two to show on the same scale.
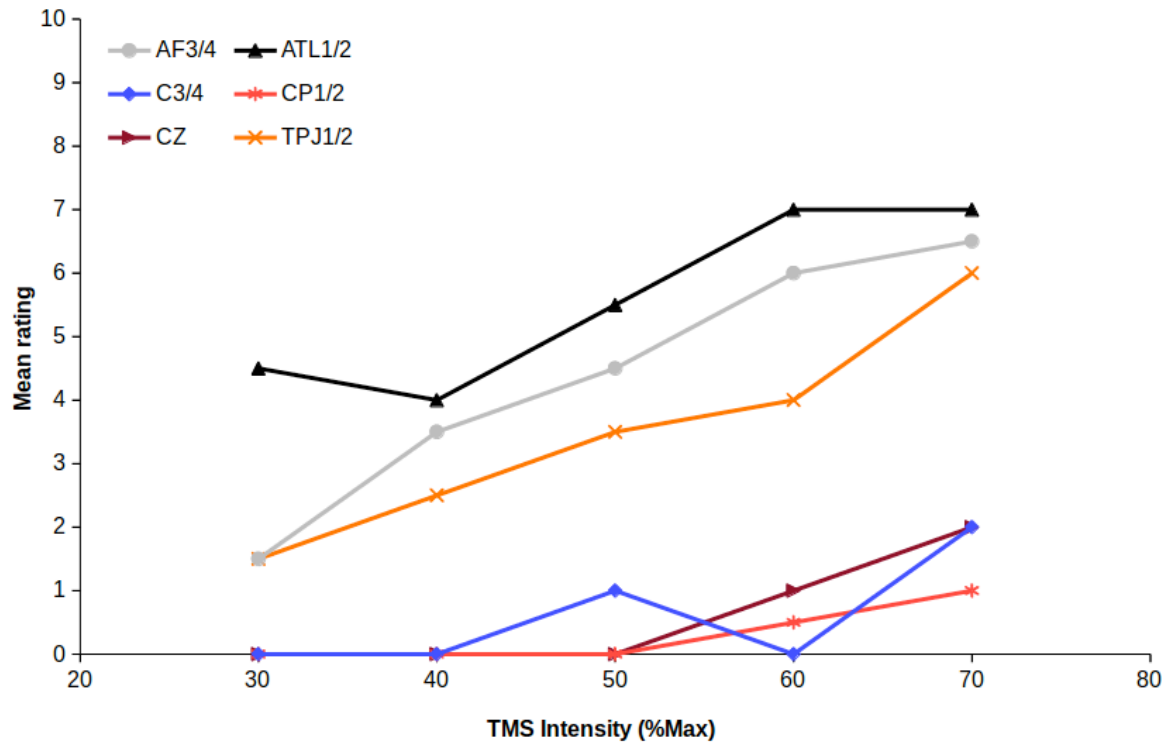
Figure C2: Mean across left and right hemispheres for each TMS site as a function of TMS intensity (%Max) for subjective ratings of annoyance. Black triangles: ATL1/2; Grey circles: AF3/4; Orange crosses: TPJ1/2; Blue diamonds: C3/4; Red asterisks: CP1/2; Maroon trianges: Cz.

Appendix D / Supplementary materials 4: Scalp-to-brain distance for the 43 sites stimulated

Methods

Twenty participants from an MRI dataset including the original TMS-SMART participant (NPH) were selected (10 females with a mean±SE age of 25.6±2.2 years; 10 males with a mean±SE age of 25.6±2.6 years).

Each participant's anatomical MRI scan (MPRAGE, 1x1x1mm) was registered to NPH's brain using 12 degree of freedom affine transformation in FLIRT (66). The resulting transform matrix was inverted using InvertXFM, then this inverted transform was applied to the 43 original locations used for TMS-SMART using ApplyXFM. This resulted, for each participant, in the approximate locations of scalp and brain sites targeted in the original experiment.

For each of 20 participants, the closest voxel on the scalp of the participant's MRI to the transformed target location was estimated (x, y, and z coordinates in scanner anatomical space). From this scalp voxel, the closest voxel of grey matter (cortical or cerebellar) was estimated. The distance between scalp and grey matter was calculated, and plot in the images below.

Results 1: Scalp to brain distances and correlations with Twitches and RTs

The mean±SE distance from scalp to brain across all 43 locations was 14.76±0.39 cm. The means and bootstrapped 95% confidence intervals are given in Table 1, main text.
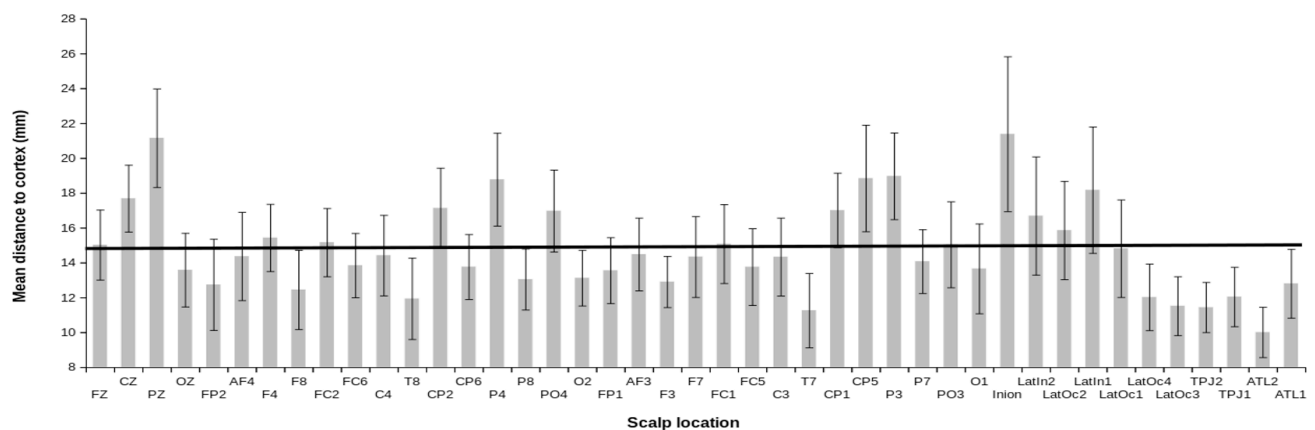

Figure D1. Mean scalp-to-brain distance for all 43 target locations. N=20. Solid horizontal line shows group mean. Error bars show 99.9% confidence intervals (i.e., p<.05, bonferroni corrected for 43 locations).

The mean across coil orientations of the median ratings of twitches (rated on a scale of 0-10) from the main experiment, and the mean effect of TMS on reaction times (in seconds) were correlated across the locations targeted. These data were taken from a different groups of participants (i.e. those who participated in the original study).

These two variables were significantly correlated with scalp-to-brain distance. For twitches, there was a significant negative correlation ($r(41)=-.525$, $p=.0003$); as mean distance to cortex increases, ratings for twitches decrease (Figure D2). For the RT difference caused by TMS, there was a significant negative correlation ($r(41)=-.400$, $p=.008$); as mean distance to cortex increases, the RT difference under TMS decreases.
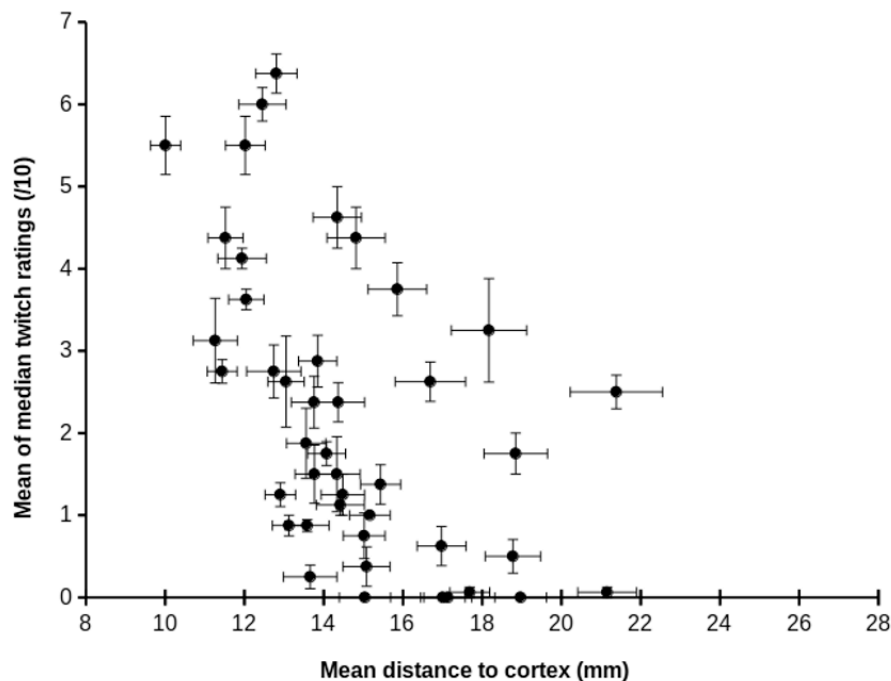

Figure D2. Mean±SE scalp-to-brain distances (mm) for 43 target locations (N=20) and mean±SE of median twitch ratings across the four coil orientations in the main experiment. Error bars show standard error. The relationship between the two variables was significant, r(41)=-.525, p=.0003.
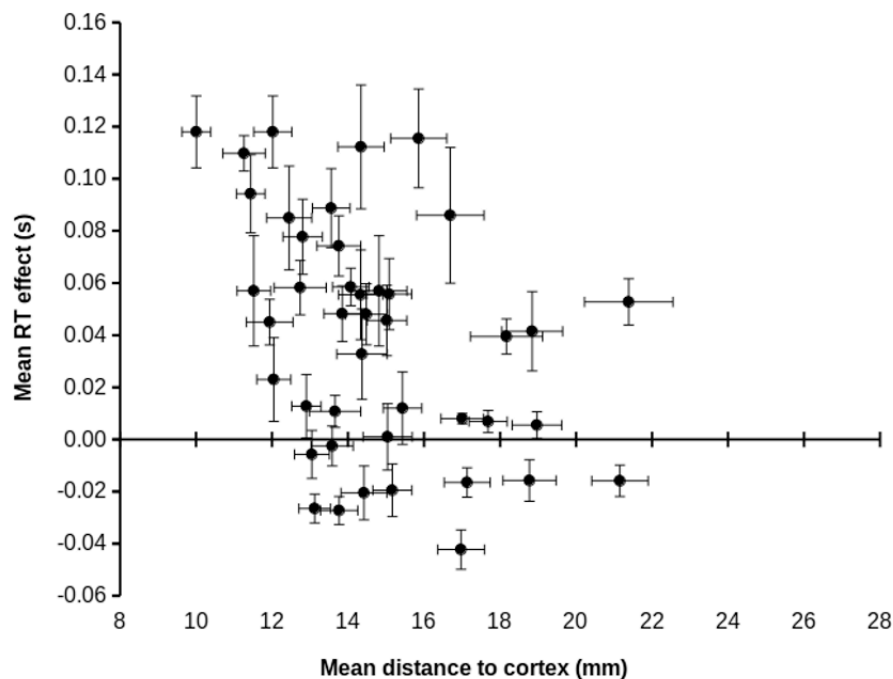
Figure D3. Mean±SE scalp-to-brain distances (mm) for 43 target locations (N=20) and mean±SE RT effects across the four coil orientations in the main experiment. Error bars show standard error. The relationship between the two variables was significant, r(41)=-.400, p=.008.

Results 2: Scalp to brain distance as a model covariate

To check whether scalp to brain distance provides a better explanation for RT differences under TMS than subjective ratings of Twitches, we repeated our analyses (detailed in 3.3 and 3.4 main text). We replaced Scalp Location with Scalp to brain distance. It was not possible to enter Scalp Location and Scalp to brain distance together, as the two variables are perfectly collinear (i.e. there is one scalp to brain distance measure per location, so the two variables are perfectly predictable from each-other). For these models, Scalp to brain distance and Twitches were scaled to reduce collinearity.

The models replicated our previous finding. When Scalp to brain distance was entered alone, it was a significant predictor of RT differences under TMS (see Table D1; estimate = -10.70, 95% CI = -18.52 - -2.88). As Scalp to brain distance decreases, the RT effects of TMS increase. Just as in our original models, Scalp Location accounted for significant variation in RT differences under TMS. The effect of Task was marginal, with the Flanker task having a trend for longer RTs (estimate = 34.87, 95% CI = -10.54 – 80.27).

Table D1: Model including Scalp-to-brain distance

Predictor	Estimate	SE	df	t	p
Fixed Effects					
Intercept	-0.26	12.66	18.00	-0.02	0.98
Scalp to brain Distance	-10.70	3.09	426.00	-3.47	0.0006*
Scalp to brain x Hemisphere interaction	0.79	4.37	429.00	0.18	0.856
Previous Trial RT	0.09	0.01	8623.00	12.81	<0.0001*
Task (Flanker > CRT)	34.87	17.91	18.00	1.95	0.067
Random Effects (intercepts)		*Variance*	*SD*		
Subject		1193.68	34.55		
Scalp Location x Subject		1890.29	43.48		
Trial x Subject		426.16	20.64		
Coil Orientation x Subject		14.28	3.78		
Congruence x Subject		580.20	24.09		

Model formula: RT Difference ~ re(Subject) + re(Scalp Location x Subject) + re(Coil Orienation x Subject) + re(Congruence x Subject) + re(Trial x Subject) + scale(Scalp to brain distance) + scale(Scalp to brain distance) x Hemisphere + previous trial RT + Task

We then added Twitch ratings to this model as a main effect and its interaction with Task (Table D2). In this model, Scalp to brain distance was no longer a significant predictor (estimate = 1.33, 95% CI = -6.10 – 8.76). Twitch ratings significantly predicted the RT difference under TMS (estimate = 20.10, 95% CI = 13.14 – 26.95), for every unit increase in Twitch ratings the RT cost of TMS increased by ~20ms. The interaction of Twitch ratings and Task was also significant (estimate = 12.25, 95% CI = 2.87 – 21.63). A unit increase in Twitch ratings had a greater cost on the Flanker task, increasing RTs by ~13ms more than for the CRT task (estimate = 12.25, 95% CI = 2.86 – 21.63).

Table D2: Model including Scalp-to-brain distance and Twitch ratings

Predictor	Estimate	SE	df	t	p
Fixed Effects					
Intercept	-0.35	13.10	18.00	-0.03	0.98
Scalp to brain Distance	1.33	2.80	436.00	0.48	0.64
Scalp to brain x Hemisphere interaction	-0.35	3.77	422.00	-0.09	0.93
Previous Trial RT	0.09	0.01	8643	12.52	<0.0001*
Twitch Ratings	20.10	2.60	632.00	7.71	<0.0001*
Twitch Ratings x Task interaction (Flanker > CRT)	12.25	3.54	605.00	3.46	0.0006*
Random Effects (intercepts)		*Variance*	*SD*		
Subject		1327.26	36.43		
Scalp Location x Subject		1309.03	36.18		
Trial x Subject		434.05	20.83		
Coil Orientation x Subject		15.71	3.96		
Congruence x Subject		581.26	24.11		

Model formula: RT Difference ~ re(Subject) + re(Scalp Location x Subject) + re(Coil Orienation x Subject) + re(Congruence x Subject) + re(Trial x Subject) + scale(Scalp to brain distance) + scale(Scalp to brain distance) x Hemisphere + previous trial RT + scale(Twitch ratings) + Task x scale(Twitch ratings)

Conclusion

We found that Scalp to brain distance was significantly correlated with both subjective ratings of twitches and RT differences under TMS. This is likely a product of physiology – areas of the brain closest to the scalp (frontal and inferior sites) also have a greater density of muscles and nerve fibres; this leads to greater discomfort. However, in models where both are taken into account, subjective discomfort is the stronger predictor of RT costs under TMS. This supports our original analysis (see 3.3 and 3.4 in the main text).

Additional Reference

66 Jenkinson, M., Bannister, P., Brady, J. M. and Smith, S. M. (2002). Improved Optimisation for the Robust and Accurate Linear Registration and Motion Correction of Brain Images. *NeuroImage*, 17(2), 825-841.