Mixed effectiveness of rTMS and retraining in the
treatment of focal hand dystonia

Teresa J. Kimberley, University of Minnesota
Rebekah L. S. Schmidt, University of Minnesota
Mo Chen, University of Minnesota
Dennis D. Dykstra, University of Minnesota
Cathrin Buetefisch, Emory University

Journal Title: Frontiers in Human Neuroscience
Volume: Volume 9
Publisher: Frontiers | 2015-07-09, Pages 385-385
Type of Work: Article | Final Publisher PDF
Publisher DOI: 10.3389/fnhum.2015.00385
Permanent URL: https://pid.emory.edu/ark:/25593/r5nbf

Final published version: http://dx.doi.org/10.3389/fnhum.2015.00385

Copyright information:
© 2015 Kimberley, Schmidt, Chen, Dykstra and Buetefisch.
This is an Open Access article distributed under the terms of the Creative
Commons Attribution 4.0 International License (http://creativecommons.org/licenses/by/4.0/), which permits making multiple
copies, distribution of derivative works, distribution, public display, and publicly
performance, provided the original work is properly cited. This license requires
credit be given to copyright holder and/or author, copyright and license notices
be kept intact.

Accessed November 20, 2018 8:04 AM EST
Mixed effectiveness of rTMS and retraining in the treatment of focal hand dystonia

Teresa J. Kimberley1, Rebekah L. S. Schmidt1, Mo Chen1, Dennis D. Dykstra2 and Cathrin M. Buetefisch3

1 Program in Physical Therapy, Brain Plasticity Laboratory, Department of Physical Medicine and Rehabilitation, University of Minnesota, Minneapolis, MN, USA, 2 Department of Physical Medicine and Rehabilitation, University of Minnesota, Minneapolis, MN, USA, 3 Department of Neurology and Rehabilitation Medicine, Emory University, Atlanta, GA, USA

Though the pathophysiology of dystonia remains uncertain, two primary factors implicated in the development of dystonic symptoms are excessive cortical excitability and impaired sensorimotor processing. The aim of this study was to determine the functional efficacy of an intervention combining repetitive transcranial magnetic stimulation (rTMS) and sensorimotor retraining. A randomized, single-subject, multiple baseline design with crossover was used to examine participants with focal hand dystonia (FHD) (n = 9). Intervention: 5 days rTMS + sensorimotor retraining (SMR) vs. Five days rTMS + control therapy (CTL) (which included stretching and massage). The rTMS was applied to the premotor cortex at 1 Hz at 80% resting motor threshold for 1200 pulses. For sensorimotor retraining, a subset of the Learning-based Sensorimotor Training program was followed. Each session in both groups consisted of rTMS followed immediately by 30 min of the therapy intervention (SMR or CTL). Contrary to our hypothesis, group analyses revealed no additional benefit from the SMR training vs. CTL. When analyzed across group however, there was significant improvement from the first baseline assessment in several measures, including tests of sensory ability and self-rated changes. The patient rated improvements were accompanied by a moderate effect size suggesting clinical meaningfulness. These results provide encouragement for further investigation of rTMS in FHD with a need to optimize a secondary intervention and determine likely responders vs. non-responders.

Keywords: transcranial magnetic stimulation, writer’s cramp, rehabilitation, sensory, human, neuromodulation

Introduction

Focal dystonia is a movement disorder that can affect any body part and severely impair a person’s ability to function in their daily life. In focal hand dystonia (FHD) there is an involuntary activation of agonist and antagonist muscles in the hand and forearm. In task specific hand dystonia, the dystonia manifests when a person attempts to execute specific tasks such as writing, typing, or playing a musical instrument. Most often these tasks are related to a repetitive action, often done in the context of an occupation, thus, impairing the ability to work and reducing quality of life. The individual with FHD otherwise has a normal neurological exam and normal function of the hand with other tasks. Although the pathophysiology of FHD remains unclear, and may differ among the different types of dystonias, there is considerable evidence for abnormal
inhibition in primary motor cortex (M1), premotor cortex (PMC), spinal cord, and brainstem (Siebner et al., 1999; Butefisch et al., 2005; Hallett, 2006, 2011; Quartarone et al., 2014). For example, it has been shown that participants with task specific dystonia have disturbances in task dependent inhibition in M1 (Butefisch et al., 2005) and intracortical inhibition (Kimberley et al., 2009). While dystonia is mainly a motor problem, mild abnormalities of sensation have been reported in patients with dystonia, even in body parts not affected by the dystonia. There is also evidence to suggest that the problem may be related to faulty sensorimotor integration (for review, Abbruzzese and Berardelli, 2003; Quartarone et al., 2014). Another abnormality in FHD is that of plasticity. It has been demonstrated that decreases in M1 inhibition facilitate the induction of plasticity (Hess et al., 1996; Di Lazzaro et al., 2006), thus, the abnormally increased plasticity in dystonia could be related to the abnormally decreased motor cortical inhibition that is found in patients with dystonia (Hallett, 2011).

The evidence for abnormal inhibition in patients with dystonia is derived from experiments using transcranial magnetic stimulation (TMS) to M1 as a testing tool to measure the level of intracortical inhibition in the GABAnergic system (Ridding et al., 1995; Butefisch et al., 2005). Most related to this proposal, the cortical silent period (CSP) is attributed to both spinal and cortical inhibition and is a test of GABA\(_B\) activity (Chen et al., 1999; Kimberley et al., 2009). CSP is the duration of EMG quiescence following a TMS pulse to M1 that is superimposed upon a low level of voluntary muscle contract in the muscle of interest (Kimberley et al., 2009).

While TMS provides a means to measure these abnormalities in M1 excitability, when applied repetitively (repetitive TMS, rTMS) it can also be used as a tool to modulate excitability of the stimulated neuronal tissue. When applied at low frequencies, it results in increased inhibition in the stimulated area of the brain that persists for more than 20 min post intervention (Chen et al., 2003). This rTMS protocol has been applied in persons with FHD in a single session and a temporary reduction in dystonic posturing and increased inhibition was recorded with a maximal effect when stimulation was applied to the PMC (Murase et al., 2005). These findings have been extended by applying low-frequency (1 Hz) rTMS to the PMC in persons with FHD for 5 days, resulting in improved handwriting velocity and changes in brain excitability (Borich et al., 2009). Additional work examined whether rTMS applied during performance of an active, but non-dystonic, hand “writing” task would enhance the beneficial effects (Kimberley et al., 2013). The multiple sessions of rTMS strengthened intracortical inhibition causing a prolongation of CSP after 3 days of intervention and pen force was reduced at day 1 and 5; notably, 68% of patients self-reported as “responders” after 5 days of intervention, suggesting potential beneficial effect of the inhibitory neuromodulation (Kimberley et al., 2013). Together, evidence suggests that rTMS induced PMC inhibition is an encouraging approach for neuromodulation therapy in people with FHD. Neuromodulation alone may not be adequate to produce clinically meaningful changes, however.

Beyond botulinum toxin, which has a variable and temporary benefit, there are few opportunities for treatment for people with FHD. Rehabilitation interventions based upon the hypothesis of aberrant learning and maladaptive plasticity have had some modest success [for review (Altenmuller and Jabusch, 2010)]. Sensorimotor training is one of these interventions that has been shown to modify brain activation and produce some functional improvements (Byl and McKenzie, 2000; Zeuner et al., 2002, 2003; Byl et al., 2003, 2009). The objective of the training is to redefine spatial and temporal processing capacities through guided activities that emphasize different aspects of sensory feedback (e.g., somatosensation, proprioception, kinesthesia) in order to improve sensorimotor integration and restore task-specific skills. A significant limitation of this training is the duration of treatment, however, which ranges from 8 to 12 weeks and requires several hours of therapy/day. Additionally, although improvements have been shown to occur in some cases, participants do not achieve premorbid levels of function and it is not universally beneficial.

The purpose of this study was to investigate whether the pairing PMC rTMS with sensorimotor retraining is more effective in improving dystonia and abnormal inhibition when compared to PMC rTMS with a control intervention. Combining neuromodulation treatment by means of rTMS or transcranial direct current stimulation with motor training is an approach that has been used in the rehabilitation of motor stroke (Hao et al., 2013) and builds on the evidence that plastic changes are either enhanced or blocked depending on the rTMS protocol (Butefisch et al., 2004). Considering the evidence for abnormally enhanced plasticity in people with FHD, executing a sensorimotor training in a more normal neurophysiological state (i.e., improved M1 inhibition) may enhance the effect of the training. Given the previous work that demonstrated increased inhibition in people with FHD following rTMS to PMC (Borich et al., 2009), as well as positive clinical changes from a prolonged intervention protocol, we hypothesized that rTMS applied to the PMC followed by sensorimotor training (SMR) would result in reduced dystonia symptoms and increased M1 inhibition, compared to rTMS alone.

Methods

Experimental Design

A randomized, crossover study design was used. Participants were randomly assigned to one of two initial treatment phases: rTMS + sensorimotor retraining (SMR) or rTMS with no specialized retraining. In order to control for the unspecific effects of an intervention such as emotional support, attention etc., subjects in the rTMS without specialized retraining condition were exposed to a control therapy (CTL, rTMS + CTL) with non-specific stretching and massage. Each phase schedule consisted of 2 days of baseline testing, five intervention sessions, 2 days of post-test testing and one follow-up session (1-week after post-test) (Table 1). After the completion of the first phase of intervention, follow-up test, and a 1-month washout period, participants crossed over to the other intervention. In other words, the people that were randomly assigned to SMR first
crosses over to receive CTL (SMR-CTL), and the people that received the CTL first, crosses to receive SMR (CTL-SMR).

The baseline and post-test testing sessions consisted of a series of behavioral and motor cortex corticospinal excitability outcome measures. Behavioral outcome measures included digitized and physician rated handwriting, self-reported symptom severity, clinical hand dystonia assessment, and sensory testing. The TMS measure used to investigate intracortical inhibition was the CSP as this has been found to be the most sensitive to change (Daskalakis et al., 2006). Digitized handwriting and TMS measures were collected at each testing session (two baseline and two post-test) while all other measures were collected once during baseline testing and once during post-test testing. The crossover design was selected to maximize participant recruitment in this rare disorder and allow all participants to receive both therapies. The two baseline- and post-test-testing sessions were chosen to determine a more stable response to the TMS and handwriting measures by averaging over 2 days.

Participants
Nine participants (3 females; mean age: 46 ± 10.6 years) (Table 2) with FHD were recruited and eight completed the entire study. One subject dropped out after the first phase of intervention due to moving out of state (Figure 1). Exclusion criteria included any neurologic conditions other than dystonia, botulinum injection within the past 4 months, medications with effects on the central nervous system and contraindications to rTMS (Rossini et al., 2007). Participants gave written, informed consent prior to participation according to the Declaration of Helsinki (World Medical Association, 2013) and study approval was granted by the University of Minnesota Clinical and Translational Science Institute and Institutional Review Board.

Behavioral Measures
Participant Rated Measures
Symptom severity was assessed using the global rating of change (GROC) (Kamper et al., 2009) and Arm Dystonia Disability scale (ADDS) (Fahn, 1989) and SF-36 (Ware et al., 2000). For the GROC, participants were asked to identify between one and three functions most impacted by FHD. At post-test 1 and follow-up, they were then asked to select a rating of perceived change that represented the level of function compared to baseline. Perceived change consisted of a ±7 point Likert scale (+7 = a very great deal better, 0 = no change, −7 = a very great deal worse) (Siebner et al., 1999; Borich et al., 2009). The ADDS was adapted from previous work (Fahn, 1989) where participants completed a survey of task difficulty for activities such as writing, handling utensils, and buttoning on a scale of 1–4 (1 = no difficulty, 4 = not able or marked difficulty). Participants completed the full SF-36 assessment with subsections of interest: “physical functioning” and “role of limitations due to physical health” and “emotional well-being.” ADDS and SF-36 were collected at baseline 1, post-test 1, and follow-up.

Clinical Hand Dystonia Assessment
Video recordings were made as participants wrote on a pad of paper with pen. Participants were asked to draw a series of 10 loops across the pad of paper followed by “The dog is barking”

<table>
<thead>
<tr>
<th>TABLE 1</th>
<th>Study schedule example.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Day 1</td>
<td>Day 2</td>
</tr>
<tr>
<td>Week 1</td>
<td>Baseline 1</td>
</tr>
<tr>
<td>Week 2</td>
<td>Intervention</td>
</tr>
<tr>
<td>Week 3</td>
<td>Follow-up</td>
</tr>
</tbody>
</table>

Participants completed each phase of treatment according to a schedule that included two baseline sessions, five intervention sessions, two post-test sessions and a follow-up session 1-week later.

<table>
<thead>
<tr>
<th>TABLE 2</th>
<th>Participant demographics.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Participant</td>
<td>Group</td>
</tr>
<tr>
<td>1</td>
<td>CTL-SMR</td>
</tr>
<tr>
<td>2</td>
<td>SMR-CTL</td>
</tr>
<tr>
<td>3</td>
<td>SMR-CTL</td>
</tr>
<tr>
<td>4</td>
<td>CTL-SMR</td>
</tr>
<tr>
<td>5</td>
<td>SMR-CTL</td>
</tr>
<tr>
<td>6</td>
<td>SMR-CTL</td>
</tr>
<tr>
<td>7</td>
<td>CTL-SMR</td>
</tr>
<tr>
<td>8</td>
<td>CTL-SMR</td>
</tr>
<tr>
<td>9</td>
<td>SMR-CTL</td>
</tr>
</tbody>
</table>

M, male; F, female; R, right; L, left; N, no; Y, yes; y, years; Sx, symptoms; FHD, focal hand dystonia; MCP, Metacarpophalangeal; DIP, Distal Interphalangeal; CTL, control; SMR, sensorimotor retraining.
and their signature, each repeated four times. A physician blinded to participant allocation rated recordings. Scoring criteria were adapted from a standardized writer’s cramp rating scale (WCRS) (Wissel et al., 1996), rating pathological flexion or extension at their wrist, fingers and elbow, presence of tremor, dystonic posture, writing speed and latency of dystonic symptoms. Final scores are expressed as two separate ratings listed as a movement score and speed score.

Handwriting
Digitized handwriting was assessed at each baseline, post-test and follow up visit. Handwriting samples were collected using a computerized tablet (WACOM Co., Ltd., Japan) with MovAlyzeR® (Neuroscript LLC, Tempe, AZ) hardware and software. Participants used a custom modified digitized pen (Kiko Software, Netherlands) to write in a self-selected pace and style on the tablet with real-time visual feedback. Writing tasks included “My country tis of thee” at a self-selected pace, repeated eight times. Data were sampled at 215 Hz (resolution: 5080 lpi, accuracy: ±0.01 pressure range: 0–800 g). Writing samples were segmented by points of minimal velocity into single strokes for analysis. Variables of interest for each stroke were automatically calculated within the software and included: mean peak vertical velocity (Zeuner et al., 2007), axial pen pressure, and normalized jerk which is a measure of dysfluency (Caligiuri et al., 2006, 2009). Data for each measure was exported for statistical analysis. One participant that did not display symptoms affecting handwriting and did not participate in the handwriting analysis.

Sensory Testing
Sensory discrimination was assessed at baseline, post-test and follow-up. Examinations included two point discrimination and the Byl–Cheney–Boczai Sensory Discriminator (BCB) (Byl et al., 2002). Two-point discrimination threshold was completed using a Disk-Criminator™. Participants were asked to reply “one” or “two” after each presentation. Static and dynamic stimuli were presented to the index and ring fingers bilaterally, meaning it was presented as a static stimuli or it was slowly swept across the skin (dynamic) (Dellon and Kallman, 1983). The BCB test was completed using a custom-made set of cubes with metal beads arranged in unique shapes such as a circle, triangle, and arrow. Each cube was fitted in a wood platform and the participant’s finger passed across the cube by the examiner. Participants were then given a visual list of shapes and asked to select the corresponding shape they felt. The index and ring finger were tested bilaterally. The BCB has been designed to assess somatosensory function and sensory discrimination in a clinical setting (Byl et al., 2002).

TMS
Corticospinal excitability was assessed with the CSP. The CSP was selected as the TMS outcome measure of interest because it is known to be altered in people with FHD (Kimberley et al., 2009) and has been shown to be the most reliable measure of excitability (Daskalakis et al., 2006). For TMS testing and intervention, participants were comfortably seated in a semi-reclined chair. The hand region has previously been described along a mid-hypotenuse line from marks 5 cm lateral and 5 cm anterior to the vertex with a 45° posterolateral orientated TMS coil (Borich et al., 2009). This measurement was used to guide coil placement over the hand motor region. TMS evoked electromyography (EMG) data were collected from the first dorsal interosseous (FDI) muscle on the affected hand. Silver chloride disk electrodes were placed on the target muscle in a belly-tendon montage with a bandpass filter 20–20,000 Hz (Cadwell Laboratory, Washington). To find the optimal position of the TMS coil (hotspot) to activate the FDI muscle, a 70-mm figure-of-eight TMS coil connected to a Magstim BiStim2 magnetic stimulator (Magstim Co., Whitland, UK) was used. Single-pulse TMS pulses were delivered while the participant was at rest until a coil position was identified that elicited the largest motor evoked potentials (MEP) with the least intensity of the maximum stimulator output. The intensity was then decreased until the minimum intensity to elicit an MEP amplitude greater than 50 µV (peak-to-peak) in at least three of five trials in the resting target muscle was determined and defined as the resting motor threshold (RMT) (Rossini et al., 1999).

CSP testing was completed during a voluntary isometric contraction of the target muscle whereby the MEP is followed by a short duration of EMG quiescence. The maximal voluntary contraction for finger abduction was recorded using a custom

FIGURE 1 | Consort diagram of screened and enrolled participants.
Participants were randomly assigned to either the rTMS + SMR or rTMS + CTL intervention first and then crossed over to receive the other intervention after 1-month washout. (SMR, sensorimotor retraining intervention; CTL, control intervention; rTMS, repetitive transcranial magnetic stimulation).
strain gauge placed around the index finger. Real-time visual feedback was given on a laptop screen to project the force produced by the participant and 20% of the maximum of three trials was calculated and displayed on a target line. For the CSP, participants were asked to contract until the target line was met, then a single TMS pulse was delivered to the hotspot at 120% RMT. Ten trials were collected with a short rest period to prevent fatigue.

CSP duration was calculated in milliseconds (ms). CSP EMG data were first rectified, and then a 10-ms moving average calculation was applied to the data. The onset of the CSP was set as the time point of the delivery of the TMS stimulus. The average of the pre-stimulus moving average data (−25 to 0 ms) was used as a threshold to determine the off-set of the CSP, defined as the point that the moving average value returned to the pre-stimulus level. The average of the 10 trials CSP was calculated.

All interventions sessions included rTMS, which consisted of 1200 pulses (20 min) delivered by a Magstim Rapid² magnetic stimulator (Magstim Co., Whitland, UK) at 1 Hz with the intensity of 80% of RMT to PMC which replicated previous studies assess the effect of rTMS without rehabilitation intervention (Borich et al., 2009; Kimberley et al., 2013, 2015). The PMC was defined as 1 cm medial and 2 cm rostral to the hotspot of the first dorsal interosseous muscle (Fink et al., 1997).

Training
Sensorimotor retraining was based on aspects of the Learning-based Sensorimotor Training program (Byl et al., 2003). The program was administered for 30 min immediately following the rTMS during the training phase. Participants were guided though supervised practice of sensory discrimination training. For example, with vision obstructed, the participant was required to discriminate grades of sandpaper as well as perform various stereognosis and graphesthesia tasks. Sensory decimation training focused primarily on the digits associated with the individual’s clinical pattern (Table 2) but was also performed with all digits, palm and dorsum and on the unimpaired hand. Tasks were primarily dynamic, meaning, the participant actively moved their digit across a surface and then used another digit to move across different choices to select the matching texture. Other tasks required the participant to select and pull out small safety pins from bowls of rice. Tasks were made progressively more difficult throughout the treatment.

The CTL intervention was also delivered immediately following the rTMS and consisted of 30 min of active and passive generalized stretching to wrist, shoulder and finger muscles and massage to the wrist, hand and shoulder musculature. All participants were instructed to avoid tasks most affected by dystonic symptoms as much as possible for the duration of the treatment. No home exercise instruction was given.

Data Processing
For measures that were collected on two pre-test sessions or two post-test sessions (handwriting and TMS), averages were calculated from the two baselines or the two post-tests, respectively. This approach was intended to represent the true baseline and post-test values. To make the data comparable between groups for all data points, the normalized change scores were calculated using a linear transformation as: Normalized change score = (X_{post} - X_{baseline})/X_{baseline} where, X_{post} is the average value of the two post-tests; X_{baseline} is the mean value of the two baselines. The only exception to the change score analysis was the WCRS, which was assessed with raw scores, due to the potential of the baseline to have a 0 score.

Statistical Analysis

Group Analysis
Pre-planned comparisons for all dependent measures (Table 3 for list) included two-factor repeated measures analysis of variance (RM ANOVA) to determine a group (SMR and CTL) × time [change from baseline (average of both baseline days) to post-test (average of post-test days 1 and 2) and follow-up] interaction of change from baseline for each treatment phase (p < 0.05). If a lack of group effect was found, data would be collapsed between groups to assess change from baseline to both follow up 1 and follow up 2 to determine longitudinal effect. Normality was assessed with Shapiro–Wilk W-test (p < 0.05). Initial comparison was with RM ANOVA (p < 0.05) and then, given lack of normality when data were collapsed, Friedman analysis of variance by ranks (p < 0.05) was applied to determine longitudinal effect over time, including both phases of the experiment. In this case, change from initial baseline to two assessment points (1. follow-up of phase one, 2. follow-up of phase two) was calculated and used for the Friedman test. Additionally, effect sizes are expressed through Kendall’s coefficient of concordance which is a non-parametric statistic associated with the Friedman test that makes no assumptions regarding the probability distribution (Kendall and Smith, 1939). As indicated, post hoc Wilcoxon signed rank test corrected for multiple comparisons (two repeated measures, p < 0.025) was used to accompany Friedman analysis.

The use of effect size in clinical intervention studies has been increasingly advocated for to improve interpretation of results, given the large variability and small sample sizes that often accompany interventional based studies (Ottenbacher and Maas, 1999; Musselman, 2007; Gomes-Osman and Field-Fote, 2015).

Single Subject Analysis
Given the small sample size and the variable response to rTMS (López-Alonso et al., 2014), a priori defined single subject analyses were also conducted. Single subject responses were evaluated using a 2 SD band method (Backman et al., 1997; Deng et al., 2013). GROC scores were assessed using minimal clinically important differences (MCID) with changes of ≥5 considered clinically meaningful (Kamper et al., 2009). Physican’s ratings of WCRS >1 point was considered meaningful. Two-point discrimination changes were considered meaningful if ≥2 testing levels on the Disk-Criminator™.
TABLE 3 | Results repeated measures ANOVA.

<table>
<thead>
<tr>
<th>Outcome</th>
<th>$F (n, df)$</th>
<th>Group x time</th>
<th>Sphericity</th>
</tr>
</thead>
<tbody>
<tr>
<td>CSP</td>
<td>1.132 (8, 2)</td>
<td>0.336</td>
<td>Assumed</td>
</tr>
<tr>
<td>2pt Non-dom ring</td>
<td>2.743 (8, 2)</td>
<td>0.081</td>
<td>Assumed</td>
</tr>
<tr>
<td>2pt Non-dom index</td>
<td>0.248 (8, 2)</td>
<td>0.782</td>
<td>Assumed</td>
</tr>
<tr>
<td>2pt Dom index</td>
<td>0.241 (8, 2)</td>
<td>0.787</td>
<td>Assumed</td>
</tr>
<tr>
<td>2pt Dom ring</td>
<td>1.028 (8, 2)</td>
<td>0.370</td>
<td>Assumed</td>
</tr>
<tr>
<td>ADDS</td>
<td>0.272 (8, 2)</td>
<td>0.763</td>
<td>Assumed</td>
</tr>
<tr>
<td>2pt Static dom index</td>
<td>0.651 (8, 1.4)</td>
<td>0.476</td>
<td>Rejected, GG $p = 0.014$</td>
</tr>
<tr>
<td>2pt Static dom ring</td>
<td>0.548 (8, 1.4)</td>
<td>0.524</td>
<td>Rejected, GG $p = 0.018$</td>
</tr>
<tr>
<td>2pt Static non-dom index</td>
<td>0.008 (8, 1.3)</td>
<td>0.964</td>
<td>Rejected, GG $p = 0.004$</td>
</tr>
<tr>
<td>2pt Static non-dom ring</td>
<td>0.524 (8, 2)</td>
<td>0.598</td>
<td>Assumed</td>
</tr>
<tr>
<td>BCB right ring</td>
<td>1.278 (8, 2)</td>
<td>0.294</td>
<td>Assumed</td>
</tr>
<tr>
<td>Emotional well-being</td>
<td>0.338 (8, 1.4)</td>
<td>0.635</td>
<td>Rejected, GG $p = 0.012$</td>
</tr>
<tr>
<td>Physical functioning</td>
<td>0.115 (8, 1.2)</td>
<td>0.785</td>
<td>Rejected, GG $p = 0.0001$</td>
</tr>
<tr>
<td>Role of limitations due</td>
<td>1.988 (8, 1.1)</td>
<td>0.177</td>
<td>Rejected, GG $p = 0.0001$</td>
</tr>
<tr>
<td>to physical health</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Handwriting jerk</td>
<td>0.373 (8, 2)</td>
<td>0.693</td>
<td>Assumed</td>
</tr>
<tr>
<td>Handwriting pressure</td>
<td>0.553 (8, 1.4)</td>
<td>0.523</td>
<td>Rejected, GG $p = 0.031$</td>
</tr>
<tr>
<td>Handwriting velocity</td>
<td>0.161 (8, 1.0)</td>
<td>0.705</td>
<td>Rejected, GG $p = 0.0001$</td>
</tr>
<tr>
<td>WCRS movement score</td>
<td>0.337 (8, 2)</td>
<td>0.717</td>
<td>Assumed</td>
</tr>
<tr>
<td>WCRS writing speed score</td>
<td>2.302 (8, 1.5)</td>
<td>0.135</td>
<td>Rejected, GG $p = 0.045$</td>
</tr>
</tbody>
</table>

CSP: cortical silent period; Dom, dominant; Non-dom, non-dominant; 2pt, two-point discrimination; ADDS, Arm dystonia disability scale; BCB, Byl–Cheney–Bocca Sensory Discriminator; WCRS, writer’s cramp rating scale; GG, Greenhouse–Geisser correction for sphericity. No significant interactions detected.

Results

Group Results

Effect of the Two Different Interventions on Behavior and Corticospinal Excitability

Results of the two factor RM ANOVA revealed no significant intervention (CTL vs. SMR) × time (Pre-test, Post-test, Follow up) interactions in any measure, indicating there was no superior benefit or neurophysiologic change to the sensorimotor retraining intervention with rTMS compared to control with rTMS (Table 3).

In contrast to our expectations, mean GROC scores did not return to zero following the washout period, which would indicate a return to the previous level of function (Figure 2). This suggests a carryover effect from the participants’ first intervention into the beginning of the second intervention. There was no difference in carryover effect between treatments.

One participant (#7) in the CTL-SMR group reported a worsening of symptoms at the first post-test (after CTL) that returned to normal at follow-up (GROC change of -4). This participant reported significant worsening of symptoms (GROC change of -4) in the subsequent SMR phase baseline and a further worsening at both post-test and follow-up (GROC change of -6). However, it appeared that the participant did not understand the rating instructions with the GROC measure, and there were conflicting responses when compared to the other self-reported measures. For example, the subject did not report worsening in the ADDS assessment which asked the same question regarding function as the identified GROC variable. For qualitative assessment this participant (#7) was removed from the visual inspection of the GROC in Figure 2, due to concerns mentioned, but is included for all other analyses.

Overall, the results of ADDS (Figure 3) suggest a perceived improvement by participants during the first treatment, regardless of group (all participants).

Overall Effect of Interventions on Corticospinal Excitability and Behavior

Given lack of overall group effect, data were collapsed across groups to determine the effects of the treatment across the entire experiment within each participant. For this analysis, normalized change scores were calculated from Phase 1 (average of the two baseline assessments for handwriting and TMS measures) to Follow up 1 and Follow up 2. Data were found to be non-normal ($p < 0.05$ Shapiro–Wilk W-test). Friedman test was conducted to evaluate differences in medians of the three test points. The results are listed in Table 4. Of note, there were significant findings in two self-reported measures; emotional well-being $X^2(2, N = 8) = 11.21, p = 0.004$, Kendall’s coefficient of concordance $= 0.70$; and ADDS: $X^2(2, N = 8) = 6.1, p = 0.048$, Kendall’s coefficient of concordance $= 0.38$, as well as four different sensory tests (see Table 4). Size of effect were considered: 0.1 = small, 0.3 = medium, and >0.5 indicate a large effect (Field, 2009) (Table 4), thus the effects were large.
and medium, respectively. Post hoc Wilcoxon signed rank test corrected for multiple comparisons (two repeated measures, \( p < 0.025 \)) revealed improvements in 2pt discrimination of the dominant ring finger (\( N = 8, Z = 2.39, p < 0.017 \)) between baseline and follow up 1 and SF36 emotional well-being between baseline and follow up 2 (\( N = 8, Z = 2.37, p < 0.018 \)), and BCB ring finger (\( N = 8, Z = 2.38, p < 0.017 \), with a trend in ADDS (\( p = 0.028 \)).

### Single Subject Analysis

Given the small number of patients in this study and known variable response to rTMS, a single subject analysis was also done. The benefit of single subject analysis in small \( n \) clinical studies, is that it allows for detailed analysis of within subject variability and response that is masked by group level statistics (Kimberley and Di Fabio, 2010). All subjects displayed changes in at least one measure but changes did not consistently reflect improvement in all measures with rTMS + SMR training, as hypothesized (Table 5). Two subjects in the SMR group experienced a clinically meaningful improvement in function. One subject had evidence of handwriting improvement despite only reporting a minimal change in symptoms by the GROC and no change in handwriting pressure (Figure 4).

### Discussion

The results suggest rTMS paired with sensorimotor training is not superior to rTMS with control. The findings further suggest a potential longitudinal benefit associated with two bouts of

<table>
<thead>
<tr>
<th>Measure</th>
<th>Friedman's ( X^2 ) (2, ( N = 8 ))</th>
<th>Baseline to follow-up 1</th>
<th>Baseline to follow-up 2</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Chi-square ( p )-values</td>
<td>Kendall's W</td>
<td>Mean (95% CI) ( p )-values</td>
</tr>
<tr>
<td>2pt Dynamic dom index</td>
<td>7.182 0.028* 0.449</td>
<td>-0.31 (-0.62, 0.00) 0.050</td>
<td>-0.32 (-0.66, 0.01) 0.074</td>
</tr>
<tr>
<td>2pt Dynamic dom ring</td>
<td>2 0.368 0.125</td>
<td>-0.46 (-0.33, 0.24) 0.713</td>
<td>-0.17 (-0.47, 0.13) 0.168</td>
</tr>
<tr>
<td>2pt Dynamic non-dom index</td>
<td>1.727 0.422 0.108</td>
<td>-0.18 (-0.60, 0.25) 0.309</td>
<td>-0.13 (-0.61, 0.35) 0.462</td>
</tr>
<tr>
<td>2pt Dynamic non-dom ring</td>
<td>3.769 0.152 0.236</td>
<td>-0.26 (-0.50, -0.02) 0.062</td>
<td>-0.36 (-0.76, 0.04) 0.062</td>
</tr>
<tr>
<td>2pt static dom index</td>
<td>6.522 0.038* 0.408</td>
<td>-0.27 (-0.46, -0.08) 0.027</td>
<td>-0.22 (-0.57, 0.12) 0.176</td>
</tr>
<tr>
<td>2pt static Dom Ring</td>
<td>7 0.030* 0.438</td>
<td>-0.29 (-0.48, -0.10) 0.017*</td>
<td>-0.38 (-0.70, -0.05) 0.043</td>
</tr>
<tr>
<td>2pt Static non-dom index</td>
<td>2.273 0.321 0.142</td>
<td>-0.11 (-0.30, 0.08) 0.276</td>
<td>-0.10 (-0.78, 0.58) 0.395</td>
</tr>
<tr>
<td>2pt Static non-dom ring</td>
<td>2.24 0.326 0.140</td>
<td>-0.05 (-0.32, 0.23) 0.932</td>
<td>-0.24 (-0.63, 0.15) 0.172</td>
</tr>
<tr>
<td>BCB left index</td>
<td>0.929 0.629 0.058</td>
<td>0.10 (-0.48, 0.27) 0.345</td>
<td>0.04 (-0.46, 0.54) 0.833</td>
</tr>
<tr>
<td>BCB left ring</td>
<td>2.385 0.304 0.149</td>
<td>0.02 (-0.34, 0.38) 0.865</td>
<td>0.53 (-0.37, 1.43) 0.367</td>
</tr>
<tr>
<td>BCB right index</td>
<td>1.5 0.472 0.094</td>
<td>0.49 (0.02, 0.96) 0.035</td>
<td>0.48 (-0.30, 1.25) 0.176</td>
</tr>
<tr>
<td>BCB right ring</td>
<td>6 0.050* 0.375</td>
<td>0.17 (-0.16, 0.50) 0.439</td>
<td>0.54 (0.17, 0.90) 0.017*</td>
</tr>
<tr>
<td>Sentence jerk</td>
<td>4.571 0.102 0.327</td>
<td>-0.14 (-0.43, 0.16) 0.263</td>
<td>-0.28 (-0.62, 0.07) 0.091</td>
</tr>
<tr>
<td>Sentence pressure</td>
<td>0.857 0.651 0.061</td>
<td>-0.15 (-0.46, 0.16) 0.575</td>
<td>-0.19 (-0.84, 0.27) 0.310</td>
</tr>
<tr>
<td>Sentence velocity</td>
<td>3.429 0.18 0.245</td>
<td>6.68 (-5.34, 18.71) 0.208</td>
<td>-0.14 (-5.90, 5.62) 0.735</td>
</tr>
<tr>
<td>CSP</td>
<td>4.75 0.093 0.297</td>
<td>-0.06 (-0.25, 0.12) 0.441</td>
<td>-0.19 (-0.39, 0.01) 0.036</td>
</tr>
<tr>
<td>Emotional well-being</td>
<td>11.214 0.004* 0.701</td>
<td>0.06 (-0.06, 0.19) 0.161</td>
<td>0.17 (0.06, 0.28) 0.018*</td>
</tr>
<tr>
<td>Physical functioning</td>
<td>4.625 0.099 0.289</td>
<td>0.02 (-0.10, 0.15) 0.893</td>
<td>0.10 (-0.10, 0.29) 0.109</td>
</tr>
<tr>
<td>Role of limitations due to physical health</td>
<td>3.8 0.15 0.239</td>
<td>0.41 (-0.35, 1.16) 0.102</td>
<td>0.33 (-0.57, 1.24) 0.655</td>
</tr>
<tr>
<td>ADDS</td>
<td>6.091 0.048* 0.381</td>
<td>0.13 (0.00, 0.25) 0.028</td>
<td>0.18 (-0.11, 0.48) 0.116</td>
</tr>
<tr>
<td>WCRS movement score</td>
<td>5.636 0.06 0.352</td>
<td>3.33 (-0.61, 7.27) 0.157</td>
<td>1.25 (-0.88, 3.38) 0.010</td>
</tr>
<tr>
<td>WCRS writing speed</td>
<td>5.2 0.074 0.325</td>
<td>0.22 (-0.12, 0.56) 0.046</td>
<td>0.25 (-0.14, 0.64) 0.18</td>
</tr>
</tbody>
</table>

CSP: cortical silent period; Dom, dominant; Non-dom, non-dominant; 2pt, two-point discrimination; ADDS, arm dystonia disability scale; BCB, Byl-Cheney–Boczai sensory discriminator; WCRS, writer’s cramp rating scale. No significant interactions detected. *\( p < 0.05 \), and **\( p < 0.025 \).
rTMS separated by 1 month, as indicated by subjective rating improvements.

The inhibitory rTMS applied here was intended to promote a favorable inhibited cortical state with rehabilitation applied during the improved neural environment following the neuromodulation. SMR was selected as a likely beneficial treatment because evidence of clinical improvement has been observed with this intervention (Byl et al., 2009). Yet, this therapy is very time intensive and may be challenging for patients to undertake, thus adjunct therapies that may enhance the effects of SMR would be advantageous. The SMR treatment performed here was brief in comparison to previous work, however which could have contributed to the lack of superior beneficial effect observed. Interestingly, despite the short duration of training, there was a significant improvement in dynamic sensory discrimination as measured by the BCB and two point static and dynamic discrimination testing on some fingers. This suggests that even a small amount of training may be able to improve this ability and it may be associated with perceived symptom improvement.

The control intervention was selected to deliver equal dose of intervention time and match investigator attention provided to the participant, but was not intended to be a specific therapy for FHD, as there is no evidence that general stretching and massage should improve FHD. Active digit and hand movement and strengthening were avoided as a control intervention, as it has been shown that there was no difference in improvement of FHD with either a task specific training or a more generic motor intervention (Zeuner et al., 2008). The evidence here is unable to determine if the control therapy gave additional benefit beyond what would be delivered with rTMS alone, as there was no group that received only rTMS.

It is interesting to note that there was no effect of increased inhibition after rTMS as measured by an increased CSP in the group analysis. Our single subject analysis suggests variable responses between subjects. It is important to note, however, that our measurements were taken at least 24 h after conclusion of the rTMS. It has been previously reported, that the increased inhibitory effect of rTMS may only be transiently observable and not persist beyond immediately post intervention (Kimberley et al., 2015). It is unknown if a more comprehensive TMS assessment may have revealed other changes.

Another consideration is that the training practice began immediately following the neuromodulation which may have influenced the potency of training effects, as there is evidence to support that a preceding bout of neuromodulation may alter the effect of motor training (Jung and Ziemann, 2009). Further, it is unknown what the ideal time frame is for rTMS effects to optimally manifest or whether treatment tasks should precede or follow the neuromodulation for the most effective capitalization of neurolasticity. It is noteworthy to consider that the vast majority of literature characterizing the effects of non-invasive brain stimulation are based on a healthy response, yet it has been shown that people with dystonia have an abnormal plastic response to both behavioral and neuromodulatory perturbations.
(Weise et al., 2011; Meunier et al., 2012; Belvisi et al., 2013). This
deficient homeostatic plasticity may then augment the cortical
response secondary to rTMS combined with behavior training,
unlike in a healthy individual. Future work should recognize
that the ideal protocol for learning in a healthy individual may
not clearly translate to an intervention in dystonia. Inclusion
of a healthy control group may be advantageous to compare
responses to neuromodulation interventions and further our
understanding of how rTMS in healthy adults may support or
differ in regards to persons with dystonia.

Future Work and Limitations

The paramount limitation to this study was a small number
of participants. The small sample may have underpowered
the study and as a result, real change between groups went
undetected. The high variance seen in our single subject analysis
is not unusual as a current conundrum in neuromodulation
is the variable response between subjects to many different
types of neuromodulation (Lopez-Alonso et al., 2014). Notably,
individual participants did display significant changes in TMS
outcomes as determined by the single subject analysis, yet when
averaged as a group, the effects were nullified. Of note, however,
is that the physiologic TMS measures did not agree with the clinical
measures. This has finding has been supported in other work as
well (Pomeroy et al., 2007; Kimberley et al., 2013; Sadnicka et al.,
2014).

Additionally, self-rated improvements did not agree with
other measures. This may indicate that the outcome measures
used are not sensitive enough to capture the therapeutic effects;
however, a placebo effect must also be considered. The likelihood
of placebo effect is weakened though when one considers that
the beneficial effect had a moderate effect size. A disagreement
between objective measurement and self-rated improvement has
been reported in musician’s dystonia where patients tended
to rate their improvement higher than an objective keyboard
assessment task (Van Vugt et al., 2014). Van Vugt et al. (2014)
speculated that this was due to patient self-selection of tasks
during their day that less severely triggered dystonic symptoms,
leading to an overestimation of ability. A similar phenomenon
may be at play in these results.

Clinical studies in dystonia are often characterized by
small samples and a heterogeneity of responses require careful
consideration in the study design (Galpern et al., 2011). It
is interesting that self-reported change continued to improve
despite ongoing intervention. This finding suggests that the
choice of a cross-over design in FHD or rTMS trials may not
be ideal. This design was selected to maximize recruitment in
a rare disease, reduce between subject variability, and give all
participants the chance to experience the active intervention. Due
to variability in the disorder on a day-to-day basis and variability
in baseline TMS responses, a two-group design would better
control for a potential carryover effect. Accordingly, factors that
predict a participants’ response should be further evaluated to
investigate the role of non-invasive brain stimulation in those
with focal dystonia and multicenter trials should be considered
to achieve higher recruitment.

This design did not assess for potential benefit from the SMR
alone with sham rTMS, nor a sham-sham effect (sham rTMS +
control intervention). Given the current state of the literature that
reports highly variable response to SMR training, and only after
extensive practice, we felt it was unlikely that 30 min of training
alone would be sufficient to demonstrate benefit, however this
could be additionally investigated. Also, the effect of rTMS to
PMC vs. sham rTMS has been studied with the same parameters
as used in this trial, thus, if large physiologic or behavioral
changes were observed a historical comparison could have been
made (Murase et al., 2005; Borich et al., 2009; Kimberley et al.,
2013, 2015). Finally, there is sufficient literature that suggests that
in some people with FHD, inhibitory rTMS to the PMC may have
beneficial effects, but other potential targets to stimulation should
be explored.

Conclusion

The results suggest rTMS paired with sensorimotor training is
not superior to rTMS paired with control therapy, but there are
potentially beneficial therapeutic effects of rTMS combined with
non-specific rehabilitation in the treatment of FHD according
to self-perceived measures. The majority of participants (6 of
9, 67%) reported improvement in self-rated function; however,
two subjects experienced no change and one subjects reported
worsening of symptoms. The variability of responses suggests
rTMS interventions may not be effective in all participants with
symptoms of dystonia and the optimal intervention and timing
of synergist intervention remains to be determined.

Acknowledgments

We thank Byron Eddy, MD for assistance. Funding provided
by the National Institutes of Health, Office of Rare Diseases
Research in the National Center for Advancing Translational
Sciences under the award number NS065701. This study has
been registered at ClinicalTrials.org under trial identification
NCT01738581. Research reported in this publication was also
supported by the National Center for Advancing Translational
Sciences of the National Institutes of Health Award Number
UL1TR000114. The content is solely the responsibility of the
authors and does not necessarily represent the official views of
the National Institutes of Health.

References

Abbruzzese, G., and Berardelli, A. (2003). Sensorimotor integration in
movement disorders. Mov. Disord. 18, 231–240. doi: 10.1002/mds.10327


Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2015 Kimberley, Schmidt, Chen, Dykstra and Buetefisch. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) or licensor are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.