Video Article

MRI-guided dmPFC-rTMS as a Treatment for Treatment-resistant Major Depressive Disorder

Katharine Dunlop¹, Pauline Gaprielian⁶, Daniel Blumberger^{5,7}, Zafiris J. Daskalakis^{5,7}, Sidney H. Kennedy^{2,3,5}, Peter Giacobbe^{2,3,5}, Jonathan Downar^{2,3,4,5}

¹Institute of Medical Sciences, University of Toronto

²MRI-Guided rTMS Clinic, University Health Network

³Department of Psychiatry, University Health Network

⁴Toronto Western Research Institute, University Health Network

⁵Department of Psychiatry, University of Toronto

⁶Faculty of Arts and Science, University of Toronto

⁷Temerty Centre for Therapeutic Brain Intervention, Centre for Addiction and Mental Health

Correspondence to: Jonathan Downar at jonathan.Downar@uhn.ca

URL: <http://www.jove.com/video/53129>

DOI: [doi:10.3791/53129](https://doi.org/10.3791/53129)

Keywords: Medicine, Issue 102, Neuroscience, Magnetic Resonance Imaging (MRI) Guided, Repetitive Transcranial Magnetic Stimulation (rTMS), Dorsomedial Prefrontal Cortex (dmPFC), Major Depressive Disorder (MDD)

Date Published: 8/11/2015

Citation: Dunlop, K., Gaprielian, P., Blumberger, D., Daskalakis, Z.J., Kennedy, S.H., Giacobbe, P., Downar, J. MRI-guided dmPFC-rTMS as a Treatment for Treatment-resistant Major Depressive Disorder. *J. Vis. Exp.* (102), e53129, doi:10.3791/53129 (2015).

Abstract

Here we outline the protocol for magnetic resonance imaging (MRI) guided repetitive transcranial magnetic stimulation (rTMS) to the dorsal medial prefrontal cortex (dmPFC) in patients with major depressive disorder (MDD). Technicians used a neuronavigation system to process patient MRIs to generate a 3-dimensional head model. The head model was subsequently used to identify patient-specific stimulatory targets. The dmPFC was stimulated daily for 20 sessions. Stimulation intensity was titrated to address scalp pain associated with rTMS. Weekly assessments were conducted on the patients using the Hamilton Rating Scale for Depression (HamD₁₇) and Beck Depression Index II (BDI-II). Treatment-resistant MDD patients achieved significant improvements on both HAM-D and BDI-II. Of note, angled, double-cone coil rTMS at 120% resting motor threshold allows for optimal stimulation of deeper midline prefrontal regions, which results in a possible therapeutic application for MDD. One major limitation of the rTMS field is the heterogeneity of treatment parameters across studies, including duty cycle, number of pulses per session and intensity. Further work should be done to clarify the effect of stimulation parameters on outcome. Future dmPFC-rTMS work should include sham-controlled studies to confirm its clinical efficacy in MDD.

Video Link

The video component of this article can be found at <http://www.jove.com/video/53129/>

Introduction

Repetitive transcranial magnetic stimulation (rTMS) is a form of indirect focal cortical stimulation. rTMS employs brief, focal electromagnetic field pulses that penetrate the skull to stimulate target brain regions. rTMS is thought to engage the mechanisms of synaptic long-term potentiation and long-term depression, thereby increasing or decreasing the cortical excitability of the region stimulated¹. Generally, the rTMS pulse frequency determines its effects: higher frequency stimulation tends to be excitatory, while lower frequency is inhibitory. Non-invasive stimulatory procedures are also widely used as a causal probe to induce temporary 'cortical lesions', and establish neural-behavior relationships or functional regions by temporarily disabling the function of a desired cortical region²⁻⁴.

Therapeutic rTMS involves multiple stimulation sessions, usually applied once daily over several weeks, to treat a variety of disorders, including major depressive disorder (MDD)⁵, eating disorders⁶, and obsessive-compulsive disorder⁷. rTMS for MDD is a potential option for medically refractory patients, and allows the clinician to noninvasively target and alter the excitability of a cortical region directly involved with depressive etiology or pathophysiology. The conventional cortical target for MDD-rTMS is the dorsolateral prefrontal cortex (DLPFC)⁸. However, convergent evidence from neuroimaging, lesion, and stimulation studies identifies the dorsomedial prefrontal cortex (dmPFC) as a potentially important therapeutic target for MDD⁹ and a variety of other psychiatric disorders characterized by deficits in self-regulation of thoughts, behaviors, and emotional states¹⁰. The dmPFC is a region of consistent activation in emotional regulation¹¹, behavioral regulation^{12,13}. The dmPFC is also associated with neurochemical¹⁴, structural¹⁵, and functional¹⁶ abnormalities in MDD.

Described here is the procedure for 20 sessions (4 weeks) of magnetic resonance imaging (MRI) guided rTMS to the dmPFC bilaterally, as a treatment for major depressive disorder. In addition to a conventional 10 Hz protocol applied over 30 min, an intermittent theta burst stimulation protocol (TBS) is discussed, which applies 50 Hz triplet bursts at 5 Hz over a 6 min session¹⁷. Both protocols are thought to be excitatory, with

the TBS protocol having the potential to achieve comparable effects using a much shorter session¹⁸. In both protocols, anatomical MRIs as well as clinical assessments are acquired prior to rTMS. Neuronavigation uses the anatomical scans to account for anatomical variability of dmPFC and optimize the location of rTMS. A relatively new 120°-angled fluid-cooled rTMS coil was also used in order to stimulate deeper midline cortical structures. Finally, rTMS intensity titration was used over the first week of rTMS sessions to ensure that patients could habituate to the higher pain levels associated with dmPFC stimulation as compared to conventional DLPFC stimulation.

Protocol

This study was approved by the Research Ethics Board at the University Health Network.

1. Subject Selection

1. Conduct an initial assessment on a prospective patient. The inclusion criteria included the presence of a current depressive episode that is resistant to at least 1 adequate trial of medication, and a Diagnostic and Statistical Manual of Mental Disorders, Fifth Edition, (DSM-5) diagnosis of MDD as established by the assessing psychiatrist. Confirm the diagnosis with a standardized Mini Mental State Examination (MINI).
2. Ensure that patients are on a stable medication or are washed out of their medication routine for at least 4 weeks prior to their first rTMS treatment session. Do not alter this medication regimen throughout rTMS treatment to help disambiguate the cause of any observed clinical improvement or deterioration.
3. Exclude patients who may have a potential contraindication to rTMS or MRI, including seizure history, cardiac arrhythmia, implanted or foreign devices/metal particulates, unstable medical conditions, or pregnancy. Patients with comorbid post-traumatic stress disorder, obsessive-compulsive disorder, other anxiety disorders, attention deficit hyperactivity disorder, bulimia nervosa or binge eating disorder, or moderate Cluster B personality features are also suitable for this treatment and need not be excluded. Patients with bipolar disorder rather than MDD may also be suitable for this treatment. Patients with psychotic disorders, active substance use, a primary diagnosis of borderline or antisocial personality disorder, or persistent depressive disorder (dysthymia) may be less suitable for treatment and may require exclusion.

2. Acquiring Magnetic Resonance Images

1. Acquire patients' MRIs at any time prior to treatment. Here, use a 3 Tesla scanner with an 8-channel phased-array head coil (refer to Table of Materials), or any scanner capable of created a 3D representation of a patient's brain.
2. Adhering to local site protocol, acquire a T1-weighted fast spoiled gradient-echo anatomical scan. Use the following parameters: TE = 12 msec, TI = 300 msec, flip angle = 20°, 116 sagittal slices, thickness = 1.5 mm, no gap, 256 x 256 matrix, FOV 240 mm. This scan will be used for real-time rTMS neuronavigation during motor thresholding and treatment sessions.

3. Preprocessing Anatomical Scans for Real-time Neuronavigation

1. Prepare for MRI guidance using a neuronavigation system.
Note: The following steps employ the Visor 2.0 neuronavigation system (refer to Table of Materials), but other navigation systems such as the Brainsight TMS Navigation, StealthStation, Aimnav, and NBS System 4 use similar procedures.
2. Segment anatomical MRIs into its scalp and brain components. Register the two segments into standard stereotactic space, such as Talairach and Tournoux space¹⁹.
3. Place target markers by selecting the following points on the MRI: Nasion; Left and right ear, targeting the tragus; Anterior commissure; Posterior commissure; Interhemispheric point (point between the two hemispheres); the anterior most point of the brain; the posterior most point of the brain; the superior most point of the brain; and the left and right most point of the brain.
4. Reconstruct the surfaces of the patient's scalp and brain in standard space to create a three-dimensional surface-based head model – this image will be used to identify stereotactic scalp coordinates overlying the dmPFC (Talairach and Tournoux coordinate X0, Y+60, Z+60) for optimal coil vertex placement during treatment.
Note: This method uses population coordinates to identify the stimulation target. Other methods to identify a stimulation target, outlined in the Discussion, include single-subject anatomy or fMRI activation maps.
5. Register brain and scalp coordinates from stereotactic space to patient space for individualized coil placement.

4. Motor Threshold Assessment

1. Seat patient in the treatment chair, adjusting the camera for an unobstructed view of the patient.
2. Place the headband with the marker clip attached to it around the patient's head. The marker clip should sit above bridge of nose.
3. Preprocess the anatomical scan for the patient as described above in Step 3.
4. Load the preprocessed anatomical scans to the neuronavigation program and turn on the camera.
5. Using a neuronavigation pen, highlight each scalp target point on the patient. The movements made with the neuronavigation pen will be projected on the television screen in the form of red lines.
6. Assess patients' motor thresholds, the minimum intensity needed to globally excite the motor pathway, prior to rTMS treatment. For this step, begin by having the patient's lower limbs extended and supported from below, using a stool or a chair equipped with an extensible leg rest.
7. For motor threshold determination, under neuronavigation, target the medial primary motor cortex. Place the coil vertex over the sagittal fissure, 0.5-1.0 cm anterior to the central sulcus. Use an angled or double-cone coil for deeper pulse penetration into medial areas. Use stimulator equipped with a fluid-cooled coil, whose windings are angled at 120° to allow deeper penetration of the pulses (refer to Table of Materials).

8. Perform motor thresholding separately for the left and right hemispheres. Orient the coil laterally to direct rTMS-evoked current flow to the desired hemisphere²⁰. For example, to stimulate the left hemisphere, orient the coil with the handle pointing rightwards and the direction of current flow toward the left hemisphere. Observe the contralateral (right) lower limb for movements during this procedure.
9. Determine threshold and elicited motor movement visually by the hallucus longus muscle of the big toe.
Note: Unlike conventional motor threshold testing that targets the hand muscle, stimulating the medial wall of the motor cortex will target the toe muscle. Motor evoked potentials (MEPs) may also be used as a more accurate determination of motor threshold, however it is a much lengthier approach.
 1. Begin by stimulating at 55% of maximum machine intensity, then adjust upwards or downwards in increments of ~5% depending on whether a response is observed. Reduce the increment size steadily to ~1% as the motor threshold is approached, as previously described²¹. Stimulate no more frequently than 0.2 Hz (once per 5 sec) to avoid inhibitory or excitatory effects over time.
 2. Once a motor threshold is established, move the vertex 1-2 cm anteriorly and posteriorly, in exploratory increments of 2-3 mm, to determine whether any alternative site offers a lower motor threshold. Use the lowest threshold achieved along this arc for each side.

5. rTMS Treatment & Adaptive Titration

1. Perform a course of neuronavigated dmPFC-rTMS, using a total of 20-30 daily sessions over 4-6 weeks. For treatments, use the 120° angled, fluid-cooled coil and the parameters listed below for dmPFC stimulation in each treatment session (refer to **Table of Materials**).
2. Seat the patient in the treatment chair, adjusting the camera for an unobstructed view of the patient.
3. Place a headband with a marker clip attached to it around the patient's head (placed laterally so as not to block the rTMS coil placement over the medial target site) as described above. Using a camera, the neuronavigation system, will detect the marker clip and will allow for preprocessing and neuronavigation.
4. Load the preprocessed anatomical scans to the neuronavigation program and turn on the camera.
5. Using a neuronavigation pen, highlight each scalp target point on the patient. The movements made with the neuronavigation pen will be projected on the television screen in the form of red lines.
6. Place the coil over the dmPFC target under MRI guidance using the neuronavigation system. For verification purposes, this point should lie close to 25% of the distance from nasion toinion. Laterally. Orient the coil laterally, with the handle pointing away from the hemisphere to be stimulated. Stimulate the left hemisphere, then re-orient the coil by 180° to stimulate the right hemisphere, maintaining the vertex in the same location over the dmPFC scalp site.
7. Ensure that the scalp site for dmPFC remains in close contact with the coil itself throughout treatment. Ensure that the patient and operator wear earplugs or other hearing protection during treatment.
8. For 10 Hz stimulation, use a duty cycle of 5 sec on, 10 seconds off for a total of 60 trains (3,000 pulses) per hemisphere per session. Perform this protocol of the left then right hemisphere by orienting the coil laterally, as previously described²⁰.
Note: The described protocol for 10 Hz rTMS is outside international safety guidelines (Rossi *et al.*, 2009). There is evidence for its safety^{18,22}.
9. For TBS stimulation, use a duty cycle of 2 sec on, 8 sec off for a total of 600 pulses per hemisphere per session. Perform this protocol of the left then right hemisphere by orienting the coil laterally, as previously described²⁰.
10. Adaptively titrate the rTMS stimulus intensity upwards from an initial value of 20% maximum stimulator intensity, to allow the patient to habituate to the pain and scalp discomfort associated with rTMS during the initial sessions²³. Increment the stimulation intensity by 2-5% on each train of stimulation, as tolerated.
 1. To assess tolerability, have the patient rate pain on a verbal analogue scale (VAS) from 0 to 10 (0 = no pain, 10 = limit of tolerability without emotional distress) after each train of stimulation is delivered.
11. Begin with a higher stimulation intensity on each session, using a level associated with moderate tolerability (VAS 5-6) from the previous session, until the patient is starting at the target intensity of 120% of resting motor threshold on each hemisphere. Maintain a verbal analogue scale of less than 9 throughout treatments during this titration process. Titration is typically completed in 2-5 days.
12. Monitor the patient for other adverse effects during treatment.
Note: The most common treatment-interrupting adverse effect is a syncopal episode, arising during the first or second session of treatment in ~1% of patients. The patient may recount feeling dizzy, faint, or disoriented, and may transiently (~10 sec) lose consciousness. Regular, repeated convulsive movements or post-episode confusion lasting more than a few seconds should be absent, however. In the event of a syncopal episode, lower the headrest on the chair if possible and encourage the patient to remain still until recovered. The session may proceed if the patient is recovered and willing to go on after a few min.
13. Monitor the patient for a generalized tonic-clonic seizure during treatment.
Note: These events are rare, and we have not observed a seizure in ~8,000 sessions of dmPFC-rTMS across >200 individual patients to date. Regular, rhythmic, vigorous convulsive movements lasting 10-40 sec, initially around 3 Hz and becoming progressively less rapid, accompanied by unresponsiveness, are suggestive of seizure rather than syncope. However, the two may be difficult to distinguish for an untrained observer.
 1. Use video monitoring during all treatments so that the episode can be reviewed by a neurologist at subsequent assessment, if necessary. In the event of such an episode, apply standard seizure first aid steps, including clearing the area of objects with the potential to cause injury, placing the patient on the ground if possible or lowering the treatment chair to the horizontal position if not, laying the patient on the left side if possible, ensuring a clear airway, and ensuring that someone remains with the patient until the seizure terminates and the person regains full alertness.
 2. Call emergency services if the seizure does not self-terminate after ~60 sec.

6. Clinical Data Collection

1. Collect standardized self-reported questionnaires at baseline, weekly throughout treatment and at follow-up (e.g., 2, 4, 6, 12, and 26 weeks post-treatment). Collect the following self-report data: Beck Depression Inventory (BDI-II)²⁴, and Beck Anxiety Inventory²⁵ on a daily basis throughout treatment.
2. Collect depression severity scores via the clinician-rated 17-item Hamilton Rating Scale for Depression score²⁶ (HamD₁₇) at baseline, weekly during treatment, and at 2, 4, 6, 12 and 26 weeks post-treatment in follow-up.

Representative Results

In previous work, HamD₁₇ was used as a measure of treatment response for 10 Hz dmPFC-rTMS. **Table 1** displays the pre- and post-treatment HamD₁₇ scores in a previously published case series²⁷. Among all subjects, pre-treatment HamD₁₇ score was 21.66.9 that significantly decreased by 4.331% to 12.58.2 post-rTMS ($t_{22} = 6.54$, $p < 0.0001$)²⁷. Using a remission criterion of HamD₁₇ ≤ 7 , 8 of 23 subjects remitted following treatment. **Table 2** displays the pre- and post-treatment BDI-II scores in the same case series²⁷. Pre-treatment BDI-II was 32.59.9 and significantly decreased by 34.231.7% to 22.012.8 post-rTMS ($t_{22} = 5.11$, $p < 0.001$). HamD₁₇ and BDI-II percent improvement was correlated to determine whether the same subjects responded on both measures ($r = 0.72$, $p = 0.0001$).

Adaptive titration was reported in a larger subset of 47 MDD patients undergoing 10 Hz dmPFC-rTMS²³. In a case series that included this subset of patients, subjects achieved the target stimulus intensity in 0.91.8 sessions and were able to complete an entire rTMS session at the intended intensity at 4.53.7 sessions²³. Adaptive titration was not correlated to treatment improvement.

A comparison of TBS to 10 Hz dmPFC stimulation was recently performed in a recent 185-subject chart review¹⁸. Outcomes did not differ significantly between groups. On the HamD₁₇, 10 Hz patients had a 50.6% response and 38.5% remission rate, while TBS patients achieved a 48.5% response and 27.9% remission rate. On the BDI-II, 10 Hz patients had a 40.6% response and 29.2% remission rate, while TBS patients achieved a 43.0% response and 31.0% remission rate¹⁸.

Subject #	Pre-Treatment HAMD	Post-Treatment HAMD	% Improvement
11	21	1	95.24
6	18	2	88.89
4	28	4	85.71
2	12	2	83.33
9	22	4	81.82
25	19	4	78.95
12	20	5	75.00
10	20	5	75.00
14	14	4	71.43
16	26	10	61.54
7	19	8	57.89
24	17	9	47.06
3	19	11	42.11
8	21	14	33.33
5	36	24	33.33
17	23	16	30.43
15	37	27	27.03
23	12	9	25.00
19	28	21	25.00
13	29	22	24.14
1	12	10	16.67
21	13	12	7.69
18	23	22	4.35
22	21	22	-4.76
20	22	24	-9.09
Mean	21.28	11.68	46.28

Standard Dev.	6.68	8.24	31.81
---------------	------	------	-------

Table 1: Individual subject HamD₁₇ improvement, using baseline and post-treatment HamD₁₇ scores.

Subject #	Pre-rTMS BDI	Post-rTMS BDI	% Improvement
11	26	3	88.46
6	21	4	80.95
4	45	9	80.00
2	17	4	76.47
16	36	13	63.89
5	35	17	51.43
3	30	15	50.00
12	26	14	46.15
14	22	12	45.45
1	33	19	42.42
10	34	20	41.18
23	32	19	40.63
9	22	15	31.82
15	57	40	29.82
19	38	28	26.32
7	25	22	12.00
18	45	41	8.89
20	45	43	4.44
17	25	24	4.00
13	44	44	0.00
22	36	37	-2.78
21	30	32	-6.67
8	24	31	-29.17
Mean	32.52	22.00	34.16
Standard Dev.	9.86	12.83	31.70
	TTEST	3.99713E-05	5.114221135

Table 2: Individual subject BDI-II improvement, using baseline and post-treatment BDI-II scores.

Discussion

Here, MRI-guided dmPFC-rTMS was applied for treatment-resistant MDD. In general, rTMS at this site was well tolerated, with mild scalp discomfort and pain at the site of stimulation that was adequately managed using adaptive titration. In open-label trials and a chart review, both 10 Hz and theta burst stimulation resulted in significant improvements in depressive severity as measured by the HamD₁₇ and BDI-II.

There are two critical steps worth noting in the rTMS treatment procedure for optimal dmPFC stimulation. First, an angled, double-cone coil allows for optimal stimulation of deeper structures within the medial aspect of the prefrontal cortex²⁸. Second, a treatment stimulation intensity of 120% resting motor threshold at this medial site is well-tolerated and without serious adverse events, despite the relatively high intensity of the applied stimulation in absolute terms when compared to the lower absolute intensities required for conventional DLPFC-rTMS. This same intensity also appears to be safe and tolerable for TBS protocols with dmPFC-rTMS, notwithstanding the significantly lower values of 80% active motor threshold more commonly used with TBS¹⁸. As previously mentioned, significant pain and discomfort is associated with anterior medial prefrontal stimulation at higher intensities²⁹. Adaptive titration was quickly and successfully used to aide in rTMS-related discomfort adaptation. In sum, the use of an angled rTMS coil and relatively high stimulation intensity (with adaptive titration) may allow for deeper penetration of stimulation to the medial prefrontal and underlying cingulate cortices²⁸, without incurring higher risks of seizure of intolerable scalp pain.

Neuronavigation is often used for precise individualized anatomical landmarking for coil vertex placement. However, one problem with MRI-guided neuronavigation is that it potentially omits the functional relationships of the desired stimulation target to other brain regions in favor of anatomical specificity across subjects. Indeed, there is significant functional connectivity variability found in association cortices, including regions of prefrontal cortex, which may impede treatment efficacy³⁰. For example, a recent study used resting-state functional connectivity to

show that left DLPFC-rTMS treatment efficacy in MDD was dependent on left DLPFC connectivity to the subgenual cingulate cortex³¹. Patients that improved with left DLPFC-rTMS tended to have anticorrelated functional connectivity between the DLPFC and the subgenual cingulate cortex at baseline. Therefore, resting-state functional connectivity could be harnessed to further optimize target placement and identify potential biomarkers once the functional characteristics of response are identified³².

One major limitation of rTMS as a treatment is that it is unclear how certain stimulation parameters influence its treatment efficacy. There is substantial variability in the parameters of conventional left DLPFC stimulation for MDD across studies, and there is also increasing evidence of substantial inter-individual variability in how some rTMS parameters affect cortical excitation and inhibition or treatment efficacy^{33,34}. For example, the effects of 10 Hz stimulation on motor evoked potential (MEP) was recently shown to vary considerably across subjects, with some showing *decreases* rather than increases in MEP strength after stimulation³⁵. Other rTMS treatment parameters that potentially require further optimization (or individualization) to maximize treatment efficacy include the number of pulses per session, the number of sessions per day, stimulation intensity and the duty cycle (how many seconds stimulation is on and off per cycle).

There are also general limitations to rTMS as a treatment. These include the logistical requirements for patients to make multiple visits to hospital for treatment, limited access to treatment for patients in remote areas, the high cost of treatment (>\$250 per session) with conventional parameters, and the low volumes of patients who can be treated per device using conventional parameters (1-2 per hr at most). Parameter optimization may help to address some of these problems in future. Other forms of non-invasive stimulation, such as transcranial direct current stimulation (tDCS), may also come to serve as a less expensive alternative to rTMS, suitable for use at home rather than in the clinic³⁶.

Despite its technical limitations, dmPFC-rTMS is clinically promising for treatment-resistant MDD. rTMS, and dmPFC-rTMS in particular, may also prove to be a promising option in other medication-resistant psychiatric illnesses including eating disorders¹⁰, obsessive-compulsive disorder³⁷, and post-traumatic stress disorder³⁸. Identifying good treatment candidates for these disorders may require additional tools other than traditional symptom-based diagnostic classification schemas – in particular, neuroimaging. Acquiring patient neuroimaging data before and after treatment allows for the identification of potential biological pre-treatment predictors and mechanisms of treatment response. Dorsomedial and subgenual cingulate resting-state functional connectivity have been identified as potential predictors to treatment response²⁷. Additionally, graph theoretical measures such as betweenness centrality have been shown to differentiate dmPFC-rTMS responders and non-responders at baseline based on subscales for hedonic responses²³. Neuroimaging also points to anterior mid-cingulate cortex and dorsomedial thalamic resting state functional connectivity change that correlates to treatment response²⁷. In sum, functional neuroimaging may become a useful clinical tool as potential predictors and mechanisms of treatment response are identified.

Since current dmPFC-rTMS studies have used an open-label design, future directions should include the creation of a sham-controlled trial to assess its therapeutic efficacy in MDD versus sham and conventional stimulation. However, creating a convincing sham-control arm is technically challenging, particularly for simulating somatosensory or nociceptive sensations, as well as convincingly blinding the rTMS technician³⁹. In a recent meta-analysis, over half of patients were able to correctly guess their treatment arm³⁹. In another meta-analysis, placebo effects were large, but comparable to escitalopram trials⁴⁰. Future studies involving a rTMS sham arm should consider a design that addresses all sensory aspects of rTMS for both the patient and the technician. Nonetheless, augmenting magnetic stimulation techniques through TBS⁴¹, priming stimulation⁴² or adjunctive cognitive behavioral therapy⁴³ or pharmacotherapy⁴⁴ may also help to optimize the therapeutic effects of rTMS. TBS in particular has the potential to achieve significant improvements in treatment duration and thus in patient volumes, access times, and treatment cost, while achieving equivalent outcomes to much longer conventional protocols^{18,45}.

In summary, rTMS of the dmPFC is a promising novel approach to therapeutic brain stimulation for treatment-resistant MDD. By incorporating the use of a MRI-guided neuronavigation system, a fluid-cooled, 120° angled stimulation coil, a high stimulation intensity and an adaptive titration schedule, dmPFC-rTMS can be safely and accurately delivered to deep targets in the medial prefrontal cortex. As these regions are central to the pathophysiology of many neuropsychiatric disorders, this approach may have promising applications not only for MDD, but also for a variety of other psychiatric conditions that are resistant to standard treatments.

Disclosures

Authors Ms. Dunlop and Ms. Gapirellian have no disclosures to report.

Acknowledgements

The authors wish to thank Aisha Dar, Vanathy Niranjani, and Dr. Umar Dar for technical assistance with rTMS delivery and data collection. The authors also wish to acknowledge the generous support of the Toronto General and Western Hospital Foundation, the Buchan Family Foundation, and the Ontario Brain Institute in funding this work.

References

1. Fitzgerald, P. B., Fountain, S., Daskalakis, Z. J. A comprehensive review of the effects of rTMS on motor cortical excitability and inhibition. *Clinical Neurophysiology*. **117**, 2584-2596 (2006).
2. Pascual-Leone, A., Gates, J. R., Dhuna, A. Induction of speech arrest and counting errors with rapid-rate transcranial magnetic stimulation. *Neurology*. **41**, 697-702 (1991).
3. Young, L., Camprodou, J. A., Hauser, M., Pascual-Leone, A., Saxe, R. Disruption of the right temporoparietal junction with transcranial magnetic stimulation reduces the role of beliefs in moral judgments. *Proceedings of the National Academy of Sciences of the United States of America*. **107**, 6753-6758 (2010).
4. Hilgetag, C. C., Théoret, H., Pascual-Leone, A. Enhanced visual spatial attention ipsilateral to rTMS-induced "virtual lesions" of human parietal cortex. *Nature neuroscience*. **4**, 953-957 (2001).

5. Berman, R. M., *et al.* A randomized clinical trial of repetitive transcranial magnetic stimulation in the treatment of major depression. *Biological psychiatry*. **47**, 332-337 (2000).
6. Van den Eynde, F., *et al.* Repetitive transcranial magnetic stimulation reduces cue-induced food craving in bulimic disorders. *Biological psychiatry*. **67**, (8), 793-795 (2010).
7. Berlim, M. T., Neufeld, N. H., Vanden Eynde, F. Repetitive transcranial magnetic stimulation (rTMS) for obsessive-compulsive disorder (OCD): an exploratory meta-analysis of randomized and sham-controlled trials. *Journal of psychiatric research*. **47**, (8), 999-1006 (2013).
8. Fitzgerald, P. B., *et al.* A randomized trial of unilateral and bilateral prefrontal cortex transcranial magnetic stimulation in treatment-resistant major depression. *Psychological Medicine*. **41**, 1187-1196 (2011).
9. Downar, J., Daskalakis, Z. J. New targets for rTMS in depression: A review of convergent evidence. *Brain Stimulation*. **6**, 231-240 (2013).
10. Downar, J., Sankar, A., Giacobbe, P., Woodside, B., Colton, P. Unanticipated Rapid Remission of Refractory Bulimia Nervosa, during High-Dose Repetitive Transcranial Magnetic Stimulation of the Dorsomedial Prefrontal Cortex: A Case Report. *Frontiers in psychiatry*. **3**, (30), 1-5 (2012).
11. Gallinat, J., Brass, M. Keep Calm and Carry On": Structural Correlates of expressive suppression of emotions. *PLoS ONE*. **6**, e1-e4 (2011).
12. Langner, R., Cieslik, E. C., Rottschy, C., Eickhoff, S. B. Interindividual differences in cognitive flexibility: influence of gray matter volume, functional connectivity and trait impulsivity. *Brain structure, & function*. (2014).
13. Jung, Y. -C., *et al.* Synchrony of anterior cingulate cortex and insular-striatal activation predicts ambiguity aversion in individuals with low impulsivity. *Cerebral cortex*. **24**, (5), 1397-1408 (2014).
14. Auer, D. P., Pütz, B., Kraft, E., Lipinski, B., Schill, J., Holsboer, F. Reduced glutamate in the anterior cingulate cortex in depression: An in vivo proton magnetic resonance spectroscopy study. *Biological Psychiatry*. **47**, 305-313 (2000).
15. Bora, E., Fornito, A., Pantelis, C., Yucel, M. Gray matter volume in major depressive disorder: a meta-analysis of voxel-based morphometry studies. *Psychiatry research*. **211**, (1), 37-46 (2013).
16. Sheline, Y. I., Price, J. L., Yan, Z., Mintun, M. A. Resting-state functional MRI in depression unmasks increased connectivity between networks via the dorsal nexus. *Proceedings of the National Academy of Sciences of the United States of America*. **107**, 11020-11025 (2010).
17. Huang, Y. -Z., Edwards, M. J., Rounis, E., Bhatia, K. P., Rothwell, J. C. Theta burst stimulation of the human motor cortex. *Neuron*. **45**, 201-206 (2005).
18. Bakker, N., *et al.* rTMS of the dorsomedial prefrontal cortex for major depression: safety, tolerability, effectiveness, and outcome predictors for 10 Hz versus intermittent theta-burst stimulation. *Brain Stimulation*. **In Press**, 1-22 (2014).
19. Talairach, J., Tournoux, P. Co-planar stereotaxic atlas of the human brain: 3-dimensional proportional system: an approach to cerebral imaging. *Neuropsychologia*. **39**, 145 <http://books.google.com/books?id=ssEbmvcJT8C> (1988).
20. Terao, Y., *et al.* A single motor unit recording technique for studying the differential activation of corticospinal volleys by transcranial magnetic stimulation. *Brain Research Protocols*. **7**, 61-67 (2001).
21. Schutter, D. J. L. G., van Honk, J. A standardized motor threshold estimation procedure for transcranial magnetic stimulation research. *The journal of ECT*. **22**, 176-178 (2006).
22. Downar, J., Geraci, J., *et al.* Anhedonia and Reward-Circuit Connectivity Distinguish Nonresponders from Responders to Dorsomedial Prefrontal Repetitive Transcranial Magnetic Stimulation in Major Depression. *Biological psychiatry*. 1-26 (2013).
23. Downar, J., Geraci, J., *et al.* Anhedonia and Reward-Circuit Connectivity Distinguish Nonresponders from Responders to Dorsomedial Prefrontal Repetitive Transcranial Magnetic Stimulation in Major Depression. *Biological Psychiatry*. **76**, (3), 176-185 (2014).
24. Beck, A. T., Steer, R. A., Brown, G. K. *Manual for the Beck depression inventory-II*. 1-82 Psychological Corporation San Antonio, TX (1996).
25. Beck, A. T., Epstein, N., Brown, G., Steer, R. A. An inventory for measuring clinical anxiety: psychometric properties. *Journal of consulting and clinical psychology*. **56**, 893-897 (1988).
26. Hamilton, M. C. Hamilton Depression Rating Scale (HAM-D). *REDLOC*. **23**, 56-62 (1960).
27. Salomons, T. V., *et al.* Resting-State Cortico-Thalamic-Striatal Connectivity Predicts Response to Dorsomedial Prefrontal rTMS in Major Depressive Disorder. *Neuropsychopharmacology official publication of the American College of Neuropsychopharmacology*. **39**, 488-498 (2014).
28. Hayward, G., *et al.* Exploring the physiological effects of double-cone coil TMS over the medial frontal cortex on the anterior cingulate cortex: an H2(15)O PET study. *The European journal of neuroscience*. **25**, 2224-2233 (2007).
29. Vanneste, S., Ost, J., Langguth, B., De Ridder, D. TMS by double-cone coil prefrontal stimulation for medication resistant chronic depression: a case report. *Neurocase*. **20**, (1), 61-68 (2014).
30. Mueller, S., *et al.* Individual Variability in Functional Connectivity Architecture of the Human Brain. *Neuron*. **77**, 586-595 (2013).
31. Fox, M. D., Buckner, R. L., White, M. P., Greicius, M. D., Pascual-Leone, A. Efficacy of transcranial magnetic stimulation targets for depression is related to intrinsic functional connectivity with the subgenual cingulate. *Biological Psychiatry*. **72**, 595-603 (2012).
32. Fox, M. D., Liu, H., Pascual-Leone, A. Identification of reproducible individualized targets for treatment of depression with TMS based on intrinsic connectivity. *NeuroImage*. **66**, 151-160 (2013).
33. Kedzior, K., Azorina, V., Reitz, S. More female patients and fewer stimuli per session are associated with the short-term antidepressant properties of repetitive transcranial magnetic stimulation (rTMS): a meta-analysis of 54 sham-controlled studies published between 1997-2013. *Neuropsychiatric disease and treatment*. **10**, 727-756 (2014).
34. Lee, J. C., Blumberger, D. M., Fitzgerald, P. B., Daskalakis, Z. J., Levinson, A. J. The Role of Transcranial Magnetic Stimulation in Treatment-Resistant Depression: A Review. *Current Pharmaceutical Design*. **18**, 5846-5852 (2012).
35. Maeda, F., Keenan, J. P., Tormos, J. M., Topka, H., Pascual-Leone, A. Interindividual variability of the modulatory effects of repetitive transcranial magnetic stimulation on cortical excitability. *Experimental Brain Research*. **133**, 425-430 (2000).
36. Brunoni, A. R., Ferrucci, R., Fregni, F., Boggio, P. S., Priori, A. Transcranial direct current stimulation for the treatment of major depressive disorder: a summary of preclinical, clinical and translational findings. *Progress in neuro-psychopharmacology, & biological psychiatry*. **39**, 9-16 (2012).
37. Mantovani, A., Simpson, H. B., Fallon, B. A., Rossi, S., Lisanby, S. H. Randomized sham-controlled trial of repetitive transcranial magnetic stimulation in treatment-resistant obsessive-compulsive disorder. *The international journal of neuropsychopharmacology / official scientific journal of the Collegium Internationale Neuropsychopharmacologicum (CINP)*. **13**, 217-227 (2010).
38. Watts, B. V., Landon, B., Graft, A., Young-Xu, Y. A sham controlled study of repetitive transcranial magnetic stimulation for posttraumatic stress disorder. *Brain Stimulation*. **5**, 38-43 (2012).

39. Berlim, M. T., Broadbent, H. J., Van den Eynde, F. Blinding integrity in randomized sham-controlled trials of repetitive transcranial magnetic stimulation for major depression: a systematic review and meta-analysis. *The international journal of neuropsychopharmacology / official scientific journal of the Collegium Internationale Neuropsychopharmacologicum (CINP)*. **16**, 1173-1181 (2013).
40. Brunoni, A. R., Lopes, M., Kaptchuk, T. J., Fregni, F. Placebo response of non-pharmacological and pharmacological trials in major depression: a systematic review and meta-analysis. *PLoS One*. **4**, e4824 (2009).
41. Chistyakov, A. V., Rubicsek, O., Kaplan, B., Zaaroor, M., Klein, E. Safety tolerability and preliminary evidence for antidepressant efficacy of theta-burst transcranial magnetic stimulation in patients with major depression. *The international journal of neuropsychopharmacology / official scientific journal of the Collegium Internationale Neuropsychopharmacologicum (CINP)*. **13**, 387-393 (2010).
42. Iyer, M. B., Schleper, N., Wassermann, E. M. Priming stimulation enhances the depressant effect of low-frequency repetitive transcranial magnetic stimulation). *The Journal of neuroscience the official journal of the Society for Neuroscience*. **23**, 10867-10872 (2003).
43. Vedeniapin, A., Cheng, L., George, M. S. Feasibility of simultaneous cognitive behavioral therapy and left prefrontal RTMS for treatment resistant depression. *Brain Stimulation*. **3**, 207-210 (2010).
44. Rumi, D. O., et al. Transcranial magnetic stimulation accelerates the antidepressant effect of amitriptyline in severe depression: A double-blind placebo-controlled study. *Biological Psychiatry*. **57**, 162-166 (2005).
45. Platz, T., Rothwell, J. C. Brain stimulation and brain repair--rTMS: from animal experiment to clinical trials--what do we know. *Restorative neurology and neuroscience*. **28**, 387-398 (2010).