

## RESEARCH ARTICLE

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# Mental imagery-induced attention modulates pain perception and cortical excitability

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## Abstract

**Background:** Mental imagery is a powerful method of altering brain activity and behavioral outcomes, such as performance of cognition and motor skills. Further, attention and distraction can modulate pain-related neuronal networks and the perception of pain. This exploratory study examined the effects of mental imagery-induced attention on pressure pain threshold and cortical plasticity using transcranial magnetic stimulation (TMS). This blinded, randomized, and parallel-design trial comprised 30 healthy right-handed male subjects. Exploratory statistical analyses were performed using ANOVA and t-tests for pain and TMS assessments. Pearson's correlation was used to analyze the association between changes in pain threshold and cortical excitability.

**Results:** In the analysis of pain outcomes, there was no significant interaction effect on pain between group versus time. In an exploratory analysis, we only observed a significant effect of group for the targeted left hand (ANOVA with pain threshold as the dependent variable and time and group as independent variables). Although there was only a within-group effect of mental imagery on pain, further analyses showed a significant positive correlation of changes in pain threshold and cortical excitability (motor-evoked potentials via TMS).

**Conclusions:** Mental imagery has a minor effect on pain modulation in healthy subjects. Its effects appear to differ compared with chronic pain, leading to a small decrease in pain threshold. Assessments of cortical excitability confirmed that these effects are related to the modulation of pain-related cortical circuits. These exploratory findings suggest that neuronal plasticity is influenced by pain and that the mental imagery effects on pain depend on the state of central sensitization.

**Keywords:** Pain, Attention, Mental imagery, Pain catastrophizing, Cortical excitability

## Background

Mental imagery is the process of envisioning specific physical or cognitive activities or perceptual experiences with the intention of altering the facilitation of neuronal networks [1]. Mental imagery is a powerful tool in improving the performance of motor skills [2-4], cognitive performance, and memory [5] and is widely used in psychological/psychiatric treatments for such disorders as schizophrenia, social phobia, and post-traumatic stress disorder [1].

Mental imagery modulates pain, and certain chronic pain syndromes are altered significantly by mental imagery, such as phantom limb pain [6]. The motor cortex is one neural circuit that can be altered with mental imagery to affect pain sensation (Figure 1a and d). There is increasing evidence of the relationship between the motor cortex and pain modulation (Figure 1a) [7].

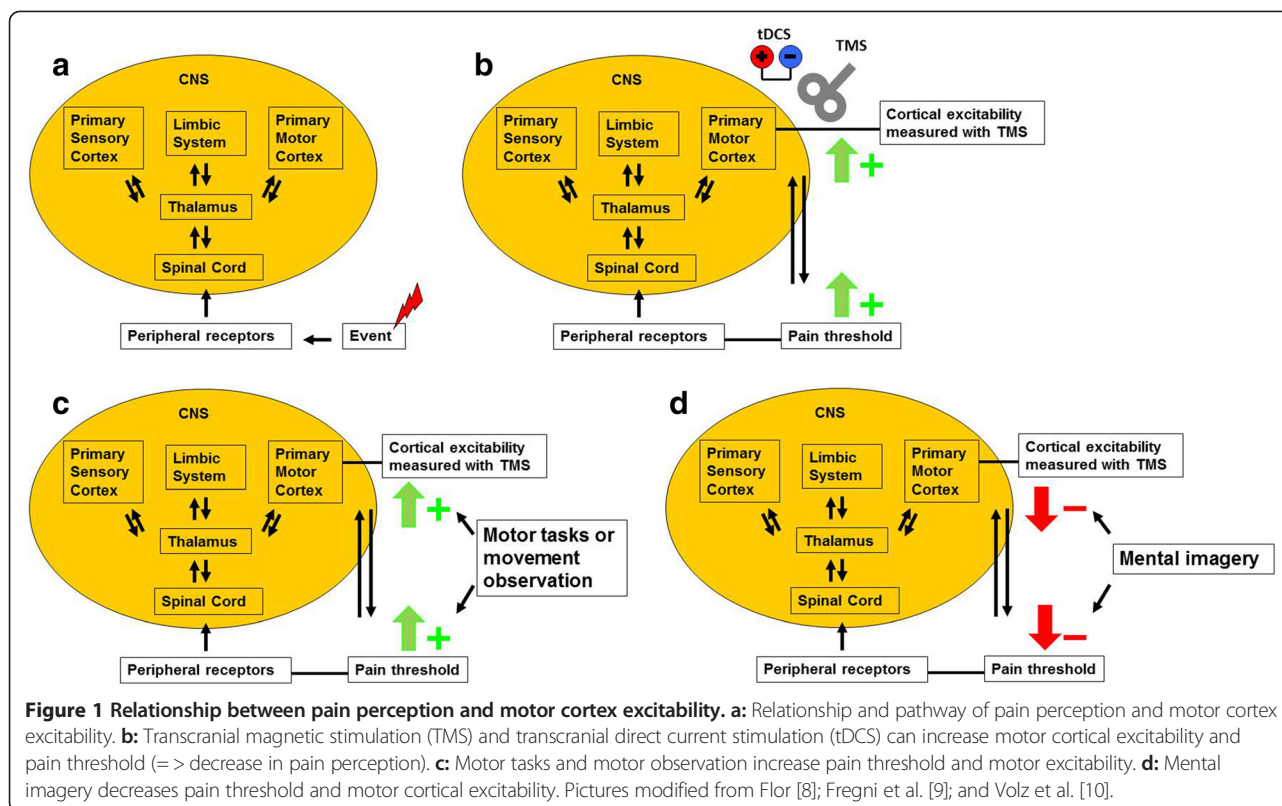
Based on data from invasive and noninvasive brain stimulation techniques, such as transcranial magnetic stimulation (TMS) (Figure 1b), there is a bidirectional relationship between pain and motor cortex excitability. Pain perception modifies TMS-indexed cortical excitability in various areas of the brain, including the motor cortex [10-14] (Figure 1a), and modification of motor cortex excitability with repetitive TMS mitigates pain [12,15,16] (Figure 1b). Thus, we hypothesized that mental imagery that is focused toward a specific hand will significantly modify TMS-indexed cortical

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excitability and pain perception as indexed by quantitative sensory testing (Figure 1d).

Several studies have demonstrated enhanced cortical excitability during mental imagery tasks by measuring motor-evoked potentials (MEPs) using TMS [17-19]. Thus, mental imagery is a proven cognitive tool that changes neuronal plasticity, allowing potential changes in cortical excitability to be correlated with modulations in pain.

In this study, we tested whether mental imagery-induced attention toward a painful stimulus of the hand alters the perception of pain, as measured by pressure pain threshold. We also measured changes in cortical excitability using transcranial magnetic stimulation (TMS) to examine the neurophysiological mechanisms of pain attention. Based on the number of outcomes in this study, we tested this hypothesis in an exploratory manner.

## Results

None of the participating subjects experienced any adverse effects. Because our experimental setting demanded good compliance and attention, we monitored *sleepiness* and *engagement* during the experiment using a questionnaire. One subject was excluded from the experiment, because he fell asleep several times, and experimental adherence was not secured. There were no significant differences between groups in *sleepiness* or *engagement* (*sleepiness*: mental imagery group:  $3.71 \pm 2.46$ , control:  $3.2 \pm 2.78$ ,

$p = 0.6$ ; *engagement*: mental imagery group:  $6.0 \pm 2.61$ , control:  $7.6 \pm 1.76$ ,  $p = 0.06$ ; unpaired *t*-test).

Moreover, we studied various age groups (18–40 years and 41–62 years) and found no significant differences (*t*-test:  $p > 0.05$ , two-tailed unpaired *t*-test) in pain or TMS outcome. Furthermore we assessed whether there were baseline differences as well as treatment related differences in VAS-anxiety and motor function as indexed by Purdue Pegboard test. All these analyses did not show significant results, confirming that these variables could not explain our results (See Table 1 for statistical details).

## Pain threshold

We analyzed the primary outcome of pain threshold as follows:

- i. We initially tested all pain results together. By ANOVA with multiple factors for time, group, and hand, there were no significant interactions ( $p > 0.05$ ) (Table 2). However, group had a main effect ( $F_{(1,27)} = 7.40$ ,  $p = 0.0079$ ), confirming our initial hypothesis that mental imagery-induced attention has a significant effect on the perception of pain, regardless of hand and time (Figure 2).
- ii. We then analyzed both hands separately, because only the left hand was targeted in our mental imagination experiment—the right hand served as

**Table 1 Statistical analyses of results of VAS-anxiety and motor function as indexed by Purdue pegboard test**

VAS for anxiety		
	Baseline/pre intervention	Post intervention
Mental imagery group	1.32 ± 1.20	0.96 ± 1.25
Control group	0.8 ± 0.94	0.43 ± 0.62
Two-tailed unpaired <i>t</i> -test comparing values between groups	P = 0.20	P = 0.13
Motor function as indexed by the Purdue pegboard test		
Left hand		
	Baseline/pre intervention	Post intervention
Mental imagery group	11.9 ± 2.5	12.24 ± 2.26
Control group	11.98 ± 2.45	12.02 ± 2.51
Two-tailed unpaired <i>t</i> -test comparing values between groups	P = 0.98	P = 0.79
Right hand		
	Baseline/pre intervention	Post intervention
Mental imagery group	12.74 ± 2.49	13.33 ± 2.48
Control group	12.89 ± 2.44	13.27 ± 2.50
Two-tailed unpaired <i>t</i> -test comparing values between groups	P = 0.87	P = 0.95

Visual analog scale = VAS. Expressed as: mean ± standard deviation.

an intraindividual control condition. We noted a significant result for the left hand for group (ANOVA,  $F_{(1,27)} = 6.35$ ,  $p = 0.018$ ), indicating that mental imagery versus controls had disparate effects on pain thresholds. We repeated the same analysis for the right hand (which was not targeted in the experiment) and found no significant results (ANOVA,  $F_{(1,27)} = 1.56$ ,  $p = 0.22$ ), confirming that the effects of pain threshold changes in the target hand were induced by the intervention (Table 2).

- iii. Pain threshold of the left hand changed in the mental imagery group by -0.63 kg ( $n = 30$ ; pre: 13.12 kg ± 2.06 kg; post: 12.48 kg ± 2.90 kg) versus +0.24 kg in the control group ( $n = 30$ ; pre: 14.12 kg ± 4.54 kg; post: 14.36 kg ± 4.18 kg). However, by unpaired *t*-test, this difference was not significant (*t*-tests: left hand:  $p = 0.17$ ; right hand:  $p = 0.59$ ). Note that the baseline thresholds did not differ in either hand in any group (*t*-tests: left hand:  $p = 0.70$ ; right hand:  $p = 0.86$ ).

#### Transcranial magnetic stimulation

Because we targeted the left hand in the experiments, TMS was assessed only over the right nondominant hemisphere. All TMS analysis results are listed in Table 2.

We first tested whether cortical excitability changed significantly with of group and time as factors by ANOVA (hand was not a factor, because only one hemisphere was examined). By ANOVA, there were significant results for both groups (MEP amplitude:  $F_{(1,26)} = 7.93$ ,  $p = 0.0091$ ).

Notably, baseline values did not differ between groups with regard to MEP amplitude [ $p = 0.597$  (pre intervention values: mental imagery group: 1.493 mV ± 0.809 mV; controls: 1.700 mV ± 1.117 mV; post intervention: mental imagery group: 1.414 mV ± 0.813 mV; controls: 1.663 mV ± 1.009 mV)] and MEP integral [ $p = 0.816$  (pre intervention: mental imagery group: 0.0214 mV\*s ± 0.0141 mV\*s; controls: 0.0224 mV\*s ± 0.0169 mV\*s; post intervention: mental imagery group: 0.0193 mV\*s ± 0.0127 mV\*s; controls: 0.0218 mV\*s ± 0.0155 mV\*s)], indicating that the effects were not due to a baseline difference between groups. Overall, MEP decreased over time (mental imagery group: from 1.493 mV to 1.414 mV). Expressed as percentages; MEP in the mental imagery group decreased by 5.33% versus 2.2% in the control group. Individual changes in MEP are shown in Figure 3.

We then determined whether the changes in pain threshold level were associated with modulations in cortical excitability by correlation analysis. Shifts in pain threshold and MEP amplitude correlated significantly (Pearson's correlation:  $r = 0.46$ ;  $p = 0.015$ ), suggesting that a decline in pain threshold (ie, greater sensitivity to pain perception) decreases cortical excitability (Figure 4).

The mean and standard deviation of MEP amplitude and integral before and after the intervention of both groups are listed in Table 3. MEP amplitudes in mV pre- and post-intervention of both groups are shown as graphs in Figure 3.

We analyzed whether the effects were due to the experimental group. By ANOVA with of group and time as factors for all cortical silent periods (CSP) with intensities of 110% (mental imagery group: pre intervention: 0.069 s ±

**Table 2 Values of statistical analyses using ANOVA**

Pain outcome both hands			TMS outcome: MEP		
Factors	Degree of freedom	p-Value	Factors	Degree of freedom	p-Value
Time (pre vs. post) Group (mental imagery vs. control) Hand (right vs. left)	$F_{(1,28)} = 7.40$	<b>0.0079</b>	Time (pre vs. post) Group (mental imagery vs. control)	$F_{(1,26)} = 7.93$	<b>0.0091</b>
Interaction of time and hand	$F_{(1,27)} = 1.43$	0.2359	Interaction of time and group	$F_{(1,26)} = 7.99$	0.3753
Interaction of time and group	$F_{(1,27)} = 0.47$	0.4947			
Time (pre vs. post) Hand (right vs. left)	$F_{(1,27)} = 2.87$	<b>0.0939</b>			
			<b>TMS outcome: CSP 110%</b>		
			Time (pre vs. post) Group (mental imagery vs. control)	$F_{(1,27)} = 17.40$	<b>0.0003</b>
			Interaction of time and group	$F_{(1,27)} = 0.2$	0.6611
<b>Pain outcome left hand</b>			<b>TMS outcome: CSP 120%</b>		
Group(mental imagery vs. control) Time (pre vs. post)	$F_{(1,27)} = 6.34$	<b>0.0178</b>	Time (pre vs. post) Group (mental imagery vs. control)	$F_{(1,27)} = 63.57$	<b>0.0001</b>
Interaction of time and group	$F_{(1,27)} = 0.84$	0.3679	Interaction of time and group	$F_{(1,27)} = 0.3$	0.5883
<b>Pain outcome right hand</b>			<b>TMS outcome: CSP 130%</b>		
Group(mental imagery vs. control) Time (pre vs. post)	$F_{(1,27)} = 2.89$	0.1001	Time (pre vs. post) Group (mental imagery vs. control)	$F_{(1,27)} = 58.35$	<b>0.00001</b>
Interaction of time and group	$F_{(1,27)} = 0.01$	0.9248	Interaction of time and group	$F_{(1,27)} = 0.74$	0.3966
			<b>TMS outcome: ICF</b>		
			Time (pre vs. post) Group (mental imagery vs. control)	$F_{(1,27)} = 9.86$	<b>0.004</b>
			Interaction of time and group	$F_{(1,27)} = 0.62$	0.4363
			<b>TMS outcome: SICI</b>		
			Time (pre vs. post) Group (mental imagery vs. control)	$F_{(1,27)} = 1.76$	0.1948
			Interaction of time and group	$F_{(1,27)} = 0.29$	0.5919

Cortical silent periods = CSP; Motor-evoked potentials = MEP; Short intracortical inhibition = SICI; Intracortical facilitation = ICF.

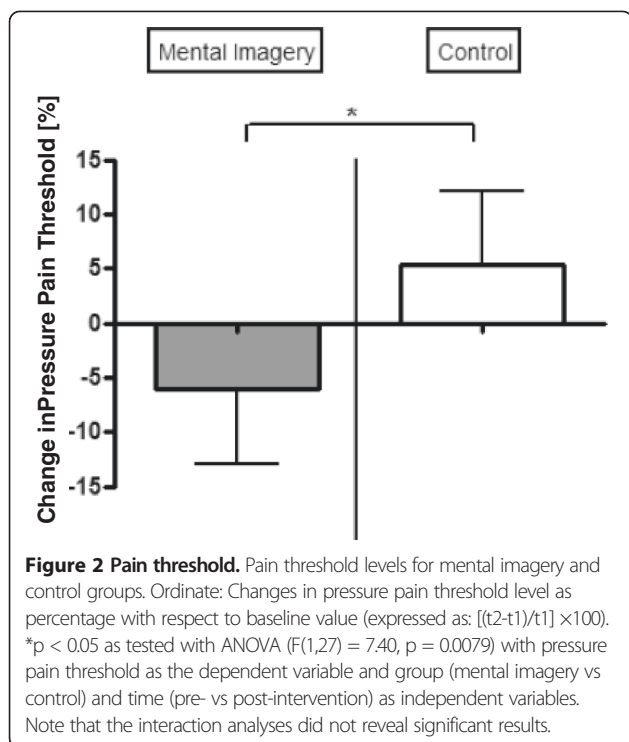
0.032 s; post intervention: 0.07 s  $\pm$  0.035 s; control group: pre intervention: 0.065 s  $\pm$  0.014 s; post intervention: 0.068 s  $\pm$  0.019 s), 120% (mental imagery group: pre intervention: 0.085 s  $\pm$  0.039 s; post intervention: 0.087 s  $\pm$  0.041 s; control group: pre intervention: 0.082 s  $\pm$  0.022 s; post intervention: 0.087 s  $\pm$  0.024 s), and 130% (mental imagery group: pre intervention: 0.095 s  $\pm$  0.043 s; post intervention: 0.097 s  $\pm$  0.044 s; control group: pre intervention: 0.099 s  $\pm$  0.028 s; post intervention: 0.106 s  $\pm$  0.029 s), there were significant results for group, confirming our findings from the pain analysis (ANOVA for CSP 110%:  $F_{(1,27)} = 17.40$ ,  $p = 0.0003$ ; ANOVA for CSP 120%:  $F_{(1,29)} = 63.57$ ,  $p = 0.0001$ ; ANOVA for CSP 130%  $F_{(1,27)} = 58.35$ ,  $p = 0.00001$ ). This result indicates that the groups differed significantly in changes in TMS measures. By ANOVA of ICF, we noted significant results for the experimental group ( $F_{(1,27)} = 9.86$ ,  $p = 0.0040$ ), demonstrating that the cortical excitability in the mental imagery group changed disparately than in the control group.

SICI was unchanged in both groups using t-tests ( $p > 0.05$  for all analyses) as well as interaction analyses (ANOVA).

We also performed linear regression analyses to test for confounders, but none revealed any significant results ( $p > 0.05$ ). Thus, potential confounders, such as age, race, education level, state of engagement and sleepiness, anxiety level, and motor function ability (ie, Purdue pegboard test), did not influence the results. No subject had a score that was higher than 6 (out of 63, mean score: 0.6  $\pm$  1.6), reflecting the absence of depressive symptoms (per [20]: a score between 11–17/63 indicates mild depressive symptoms, and a score over 18/63 is defined as clinically relevant).

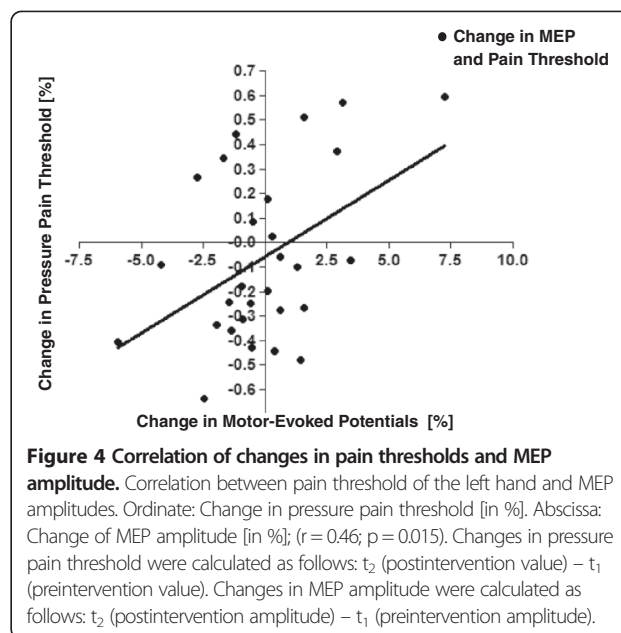
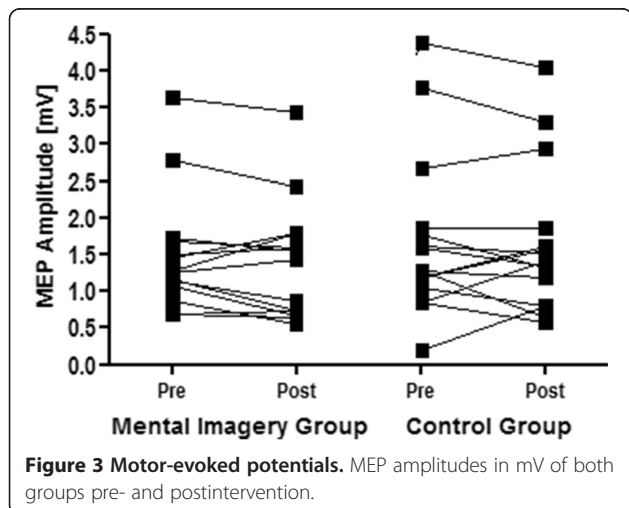
## Discussion

In this exploratory study, we did not observe any significant effect of mental imagery on pain in our primary analysis. However, we noted a small effect in the exploratory



within-group analyses. The targeted left hand experienced a decrease in pressure pain threshold, indicating a rise in pain perception. In contrast, pressure pain threshold rose in the controls, although, this within-group effect was modest. Further analyses of TMS assessments revealed a significant positive correlation of changes in pain and alterations in cortical excitability, suggesting that a decline in pain threshold decreases cortical excitability.

Our results can be interpreted as unexpected, because the literature claims that MEPs increase significantly during voluntary or imagined movement of the finger [21]. Also, studies on phantom limb pain suggest that



the use of mental imagery is an effective method of reducing pain [22-25]. Nevertheless, our findings demonstrate the opposite.

One explanation is that we included healthy subjects with an experimental pain model—not patients who were suffering from actual pain. Because chronic pain patients have a deficient pain matrix and altered pain-related neural networks, such methods as mental imagery might induce differential effects in patients versus healthy subjects [8,16,25,26]. In previous studies, we found that motor tasks, sensory stimuli, and movement observation can change pressure pain threshold levels and cortical excitability [10,27,28] (Figure 1c), in which active tasks for one hand can ameliorate the perception of pain in the targeted hand. In contrast, we also found that the untargeted hand experienced a decrease in pain threshold, indicating greater perception of pain [10,27].

In the current exploratory study, we generated data that suggest that the opposite effects are occurring, because attention and expecting pain indicated enhanced perception of pain. Thus, our findings implicated a close, reciprocal relationship of these two emotional and alertness states. Consistent with our modest effects on pain, these effects have been confirmed by several studies [29-34].

In a separate study, we demonstrated that changes in pain correlate significantly with modulations in TMS assessments, as in our present report [10,35]. In that study, we tested movement observation by showing a video, which was intended to be a distraction to a painful stimulus [10] (Figure 1c). Pain threshold levels increased, reflecting a decline in the perception of pain [10]. This finding is consistent with our current results, which yielded the opposite effects in both outcomes. Subjects paid attention to the



**Table 3 Results of motor-evoked potentials**

Mental imagery group					Control group				
Time	Pre		Post		Time	Pre		Post	
Subject	Amplitude	Integral	Amplitude	Integral	Subject	Amplitude	Integral	Amplitude	Integral
Number	(mV)	(mV*s)	(mV)	(mV*s)	Number	(mV)	(mV*s)	(mV)	(mV*s)
1	0.68	0.01	0.73	0.01	3	2.68	0.04	2.94	0.04
2	1.25	0.01	1.43	0.01	6	1.85	0.03	1.87	0.03
4	MD	MD	MD	MD	7	0.85	0.01	1.42	0.01
5	1.28	0.01	1.79	0.03	9	1.28	0.01	0.64	0.01
8	0.86	0.01	0.55	0.01	14	1.19	0.01	1.56	0.02
10	1.46	0.02	1.80	0.03	15	1.58	0.02	1.34	0.02
11	1.51	0.02	1.60	0.02	16	1.62	0.02	1.53	0.02
12	1.67	0.03	1.60	0.02	17	1.18	0.02	1.62	0.03
13	1.16	0.02	0.73	0.00	19	4.38	0.06	4.04	0.05
18	1.07	0.02	0.66	0.01	21	0.84	0.01	0.57	0.01
20	2.78	0.04	2.42	0.03	22	1.76	0.02	1.31	0.01
24	1.72	0.02	1.54	0.02	23	3.77	0.06	3.29	0.05
25	0.70	0.01	0.64	0.01	27	0.20	0.00	0.80	0.01
26	3.63	0.06	3.43	0.05	28	1.28	0.01	1.18	0.01
29	1.14	0.02	0.86	0.01	30	1.06	0.01	0.81	0.01
<b>Mean</b>	1.49	0.02	1.41	0.02	<b>Mean</b>	1.70	0.02	1.66	0.02
<b>SD</b>	0.81	0.01	0.81	0.01	<b>SD</b>	1.12	0.02	1.01	0.02

Baseline values did not differ between groups: MEP amplitudes:  $t$ -test:  $p = 0.597$  (mental imagery group: mean:  $1.493 \text{ mV} \pm 0.809 \text{ mV}$ ; controls: mean:  $1.700 \text{ mV} \pm 1.117 \text{ mV}$ ); MEP integral:  $t$ -test:  $p = 0.816$  (mental imagery group: mean:  $0.0214 \text{ mV*s} \pm 0.0141 \text{ mV*s}$ ; controls: mean:  $0.0193 \text{ mV*s} \pm 0.0127 \text{ mV*s}$ ). Mean and standard deviation of MEP amplitudes and integral before and after the intervention in both groups (MD = missing data).

hand and focused on the painful stimuli, which effected a moderate decrease in pressure pain levels, thus indicating pain sensitization. Further, previous evidence has shown that cortical excitability and pain significantly correlate in chronic pain patients [16,36-38], supporting our exploratory findings.

The modest within-group effects seen in this article suggest that complex processes, such as distraction and concentration, and influences of attentional processes have an impact on pain perception [32,39,40]. For instance, paying attention to pain, focusing on a painful body part, and rumination of painful stimuli are components of pain catastrophizing [41]. In our experiment, we mimicked attention to a specific body part and placed the focus on painful stimuli on the same area. Thus, we might be able to mimic one part of the complex mechanism of pain catastrophizing and attention-modified pain perception in healthy subjects.

Moreover, cortical excitability is significantly associated with pain coping strategies, such as pain catastrophizing, supporting our findings that cortical excitability results correlate significantly with pain outcome [35]. A subsequent study with a larger sample size should be performed to determine the predictors of pain catastrophizing and test interventions to prevent attention, focusing, and rumination on pain. The mechanism of pain

augmentation must be determined to develop novel targeted therapies for chronic pain conditions.

There are some limitations in our study. First, we assessed TMS on one hemisphere, because only the left hand was the target in our experimental setting—cortical excitability changes were not measured in the other hemisphere. However, because there was no significant effect of pain threshold in the non-targeted right hand, we assumed that there was no such effect on the non-targeted left hemisphere. Further, studies with larger sample sizes that include chronic pain patients are necessary to confirm the results and to show the differences in pain processing between healthy subjects and chronic pain patients.

Also, the age difference between subjects could be a limitation. Although there no significant differences between younger and older subjects, we cannot exclude that age has an influence on our results. In addition, with regard to limitations due to the statistical results: we did not find any significant interactions—only a modest within-group effect on pain outcome was noted.

## Conclusions

Mental imagery-induced attention and focusing on a painful stimulus of a specific body part might enhance the perception of pain. Our findings highlight the effects

and influence of attentional processes on pain perception, which might be components of the mechanisms of pain catastrophizing and chronification, because both phenomena include recurring attention on pain and a painful body part. Further, cortical plasticity changes in the same direction as those in pain perception. These exploratory findings suggest that neuronal plasticity is governed by pain and that pain-related neural networks are altered by the attention state.

**Methods**

**Experimental design**

This study was a blinded, randomized, controlled, parallel-design trial. Thirty healthy right-handed male subjects were enrolled. The participants were randomized into 1 of 2 groups (15 volunteers in both study arms; in total, 30 participants). Both groups underwent the same procedures, including determination of pressure pain threshold and measurements of cortical excitability with transcranial magnetic stimulation before and after the intervention (Figure 5). The intervention was mental imagery of hand movements or a control task (see below).

Other assessment scales were administered to control for sleepiness and engagement during the experiment, in addition to the visual analog scale (VAS) for anxiety and the Purdue pegboard test. The VAS for anxiety [42] is a 10-point rating system (with 0 indicating no anxiety and 10 indicating the worst possible anxiety). The same system was used for the sleepiness and engagement questionnaire (with 0 indicating no sleepiness or engagement and 10 indicating the greatest sleepiness or engagement). The Purdue pegboard test assessed motor function throughout the experiment [43]. The questionnaire for sleepiness and engagement was given after the intervention period, pain threshold measures, and TMS assessment.

This study was approved by the institutional review board of Spaulding Rehabilitation Hospital (Harvard Medical School, Boston, USA) and was conducted per the ethical principles of the World Medical Association/Declaration of Helsinki. The ClinicalTrials.gov identifier is NCT01404039.

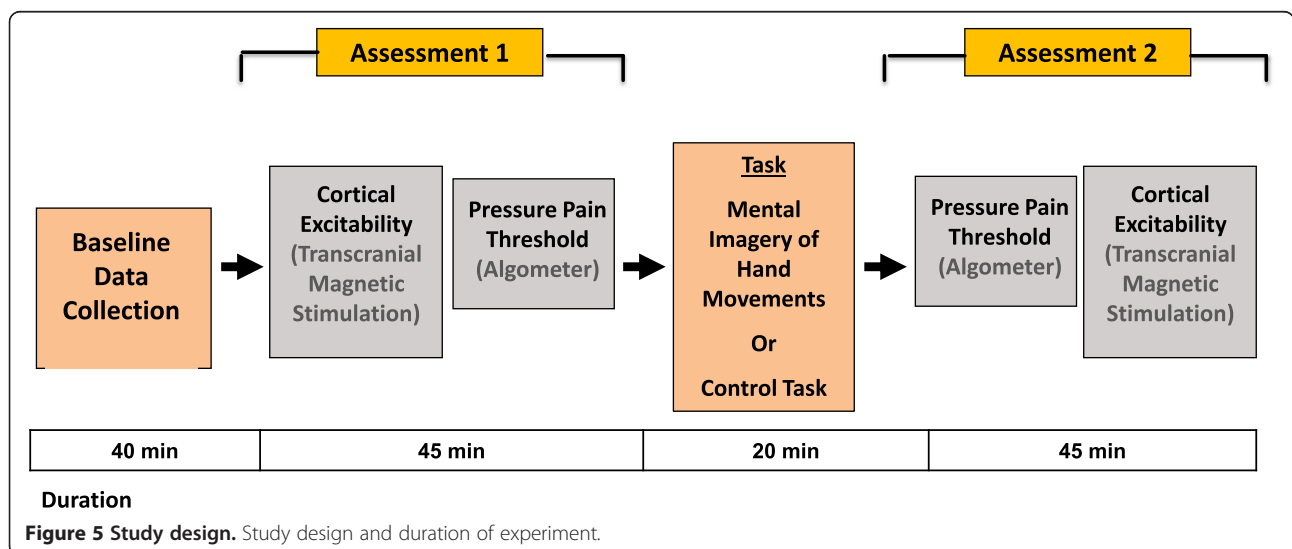
**Intervention: mental imagery group**

Subjects (mean age: 40 years  $\pm$ 12.59 years, range: 19–62 years) were asked to practice mental imagery of motor practice, consisting of sequential and repetitive finger movements of the left hand, for 10 minutes [1,3]. The subjects were seated in a chair and asked to keep their arm and hand muscles fully relaxed, which was first controlled visually by 1 of the experimenters and also controlled by surface electromyographic activity (EMG) recordings.

Participants were instructed to pay attention to the left hand. Further, they were asked to focus on the painful stimulus during the pressure pain threshold measurements, which were performed immediately before and after the period of mental imagery. Subjects were instructed to imagine repetitive movements of the left index finger to the left thumb for 5 minutes (thumb to second finger). Subjects were then asked to consecutively imagine sequential movements of the remaining fingers to the left thumb (thumb to third, fourth, and fifth fingers) for 5 minutes. They were told to concentrate and focus on the left hand. In all subjects, surface (EMG) was recorded simultaneously from the flexor digitorum superficialis (FDS) and opponens pollicis (OP) bilaterally.

**Control group**

The control for mental imagery consisted of performing a simple mental calculation, such as adding or subtracting a 1-digit number from a starting number (eg, 1 + 1 = 2; 2 + 1 = 3; 3 + 1 = 4, etc.). In all subjects (mean age: 36.8 years



$\pm 14.37$  years, range: 20–60 years), EMG activity was recorded simultaneously from the FDS and OP bilaterally. EMG activity was recorded simultaneously from the flexor digitorum superficialis (FDS) and opponens pollicis (OP) bilaterally.

### Subjects

Thirty healthy right-handed male subjects (mean age: 38.1 years  $\pm 13.24$  years, range: 18–62 years) were recruited through postings in public places and the internet. Participants who fulfilled the following criteria were eligible to participate: (1) male; (2) aged between 18 and 65 years; (3) right-handed, indexed per the Edinburgh Handedness Inventory [44]; (4) no neurological or psychiatric disorders, as assessed by Beck Depression Inventory [20] (mean score:  $0.6 \pm 1.6$ ); (5) no use of central nervous system medications; (6) no contraindications to TMS [45]; (7) no rheumatologic disease; and (8) no history of alcohol or substance abuse within the last 6 months.

All 30 subjects provided written informed consent. To create a homogeneous study population, we enrolled only right-handed male participants, because female hormones influence cortical excitability and the dominance of the hemisphere [46,47].

### Pain assessment: pressure pain threshold

Pressure pain thresholds were determined with a Commander algometer (JTech Medical Industries, Salt Lake City, USA). The algometer has a 1-cm<sup>2</sup> rubber probe, which was pressed against the hand (the thenar area of each hand). The applied velocity was 1 kg/cm<sup>2</sup>s. Subjects reported when the pressure stimulus became painful [48]. Because testing pressure pain threshold is operator-dependent, only 1 experienced researcher measured pain to avoid interrater variability and to ensure the same velocity of the increase in pressure. The investigator was blinded to the intervention and unable to view the pressure intensities.

Three repetitions were measured, the thresholds for which were averaged. The area in which pressure was applied minimally differed for each repetition to avoid habituation. One measure took approximately 1 minute per test, totaling roughly 6 minutes. Pressure pain thresholds were determined for both hands. Pain assessments were conducted immediately before and after the period of mental imagery to avoid disrupting the mental imagery process and the subject's concentration.

### Cortical excitability: transcranial magnetic stimulation (TMS)

TMS was assessed using a Bistim2 stimulator and a figure-eight coil (Magstim Company LTDA, UK). Ag/AgCl electrodes (ADInstruments, Colorado Springs, CO, USA) were placed over the first dorsal interosseus muscle (FDI), and a ground electrode was placed over the subject's forearm.

EMG recordings were processed using Powerlab 4/30 (ADInstruments, Colorado Springs, CO, USA) with a band pass filter of 20–2000 kHz. Offline analyses were performed on a private computer using LabChart (ADInstruments, Colorado Springs, CO, USA). First, head measures were taken to identify the approximate spot of the motor cortex (using the vertex as the reference). Then, the TMS coil was held tangentially over the motor cortex at an angle of 45° with respect to the sagittal line of the head. The hotspot was determined by carefully eliciting the most stable and highest MEP amplitudes over the FDI. The best location was marked with a pen on a swim cap, which was worn by each of the subjects.

We defined the following TMS parameters for the assessments. Cortical silent periods (CSPs) are a measure of intracortical inhibition, changes in which are related to GABA activity [49]. In chronic pain, the CSP declines [50]. Our hypothesis was that mental imagery would increase the CSP. Further, motor-evoked potentials (MEPs) are a direct measure of corticospinal excitability [49] that rise in association with alleviation of pain [51]. We hypothesized that mental imagery would increase MEPs. Short intracortical inhibition (SICI) is believed to be controlled by presynaptic GABA<sub>B</sub> [49]. We hypothesized that SICI would be enhanced with mental imagery. Intracortical facilitation (ICF) is linked to NMDA receptor activation [49], and we hypothesized that ICF decreases during mental imagery, because it is reduced with pain treatment.

TMS was evaluated on the right hemisphere and the contralateral, left FDI, which reflected the nondominant hemisphere in all subjects. Resting motor threshold (MT) was determined by eliciting 3 of 5 MEPs with a minimal peak-to-peak amplitude of 100  $\mu$ V [27,52,53]. MEPs were excited with 130% of the individual MT. CSPs were measured at intensities of 110%, 120%, and 130% of the individual MT. Subjects were instructed to perform isometric voluntary contraction during CSP recordings with 15% of maximum contraction force, controlled by a mechanical pinch gauge (Baseline® Evaluation Instruments, Chattanooga, TN, USA) [27,53,54].

TMS measurements included SICI with an interstimulus interval (ISI) of 3 ms and ICF with an ISI of 10 ms [55]. For paired-pulse measurements, the first stimulus was set to 70% of the individual MT, and the second stimulus was set to the individual MEP intensity. Fifteen recordings of each TMS assessment protocol were randomly elicited. Offline analyses included measures of peak-to-peak amplitude, the area-under-the-curve of all MEPs, and the relative duration of CSPs (time from last MEP until normal muscle activity was re-achieved).

### Further assessments

The Beck Depression Inventory (BDI) is a 21-item test that is presented in multiple-choice format that



measures the presence and degree of depression in adults [20].

The Purdue pegboard test measures finger dexterity and monitors motor skills by assessing changes over time through the speed of performance [56,57]. The Purdue pegboard test also assesses motor function [43]. The subject is seated comfortably at a normal-height table. A pegboard is placed in front of the person with a row of cups at far end. The cup on the pegboard contains 25 pins that subjects must place in the correct order (starting with the top hole) as fast as possible. Only one pin at a time can be picked up. If a pin is dropped during the test, the subjects should continue picking up another pin. The entire procedure takes 30 seconds. Each participant repeated the task for 3 times, and median was calculated [58].

### Statistical analyses

Data are presented as mean  $\pm$  standard.

Analyses were performed using STATA (version 11.0, College Station, Texas, US) and GraphPad Prism (version 4.00 for Windows, GraphPad Software, La Jolla, CA, USA).

Mixed ANOVA models with hand, time, and group as factors were used to analyze changes in pain outcome for the effects of hand (left hand, which was the target of mental imagery, and right hand, which was not the target), group (mental imagery versus control group), and time (before and after intervention). TMS data were analyzed with a mixed ANOVA model using measures of cortical excitability (MEP, CSP, SIC1, ICF) to test for time (before and after intervention) and group (mental imagery group versus control group) as factors.

Pearson's correlations were conducted to examine the relationship between changes in motor cortical excitability via TMS measurements and pain outcome, as assessed by pressure pain threshold.

To identify potential confounding variables and to detect any association with dependent (pressure pain threshold) and independent variables (such as age, race, education level, state of engagement and sleepiness, anxiety level, motor function ability—ie, Purdue pegboard test), we performed multiple regression analyses. In addition, two-tailed unpaired t-tests were used to control for differences in baseline characteristics and assessment scores between groups.

Significance was considered at a two-sided level of  $p < 0.05$ . We did not correct for the significance threshold in the multiple comparisons, given the exploratory nature of this study and the number of outcomes. For the TMS measurements, based on the number of tests, it is likely that at least 1 of the significant results is due to chance.

### Competing interests

The authors declare that they have no competing interests. This study was also supported by a NIH grant 1R21DK81773-1A1.

### Authors' contributions

VMS made substantial contributions to conception and design, acquisition of data, analysis and interpretation of data and wrote the manuscript as well as gave the final approval of the version to be published. SCV made substantial contributions to conception and design, acquisition and interpretation of data and was revising the manuscript as well as gave the final approval of the version to be published. SA made substantial contributions to conception and design, acquisition of data and was revising the manuscript as well as gave the final approval of the version to be published. FF made substantial contributions to conception and design, interpretation of data and was revising the manuscript critically for important intellectual content as well as gave the final approval of the version to be published.

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### References

- Pearson DG, Deepröse C, Wallace-Hadrill SM, Burnett Heyes S, Holmes EA. Assessing mental imagery in clinical psychology: a review of imagery measures and a guiding framework. *Clin Psychol Rev.* 2012;33:1–23.
- Schuster C, Butler J, Andrews B, Kischka U, Ettlin T. Comparison of embedded and added motor imagery training in patients after stroke: study protocol of a randomised controlled pilot trial using a mixed methods approach. *Trials.* 2009;10:97.
- Malouin FRC, Doyon J, Desrosiers J, Belleville S. Training mobility tasks after stroke with combined mental and physical practice: a feasibility study. *Neurorehabil Neural Repair.* 2004;18:66–75.
- Hoyek N, Di Rienzo F, Collet C, Hoyek F, Guillot A. The therapeutic role of motor imagery on the functional rehabilitation of a stage II shoulder impingement syndrome. *Disabil Rehabil.* 2014;36(13):1113–9. doi:10.3109/09638288.2013.833309. Epub 2014 Feb 28.
- Hussey EP, Smolinsky JG, Piryatinsky I, Budson AE, Ally BA. Using mental imagery to improve memory in patients with Alzheimer disease: trouble generating or remembering the mind's eye? *Alzheimer Dis Assoc Disord.* 2011;26:124–34.
- MacIver K, Lloyd DM, Kelly S, Roberts N, Nurmikko T. Phantom limb pain, cortical reorganization and the therapeutic effect of mental imagery. *Brain.* 2008;131:2181–91.
- Castillo Saavedra L, Mendonca M, Fregni F. Role of the primary motor cortex in the maintenance and treatment of pain in fibromyalgia. *Med Hypotheses.* 2014;83(3):332–6. doi:10.1016/j.mehy.2014.06.007. Epub 2014 Jun 17.
- Flor H. Cortical reorganisation and chronic pain: implications for rehabilitation. *J Rehabil Med.* 2003;41 (Suppl):66–72.
- Fregni F1, Freedman S, Pascual-Leone A. Recent advances in the treatment of chronic pain with non-invasive brain stimulation techniques. *Lancet Neurol.* 2007;6(2):188–91.
- Volz MS, Suarez-Contreras V, Portilla AL, Illigens B, Bèrmphol F, Fregni F. Movement observation-induced modulation of pain perception and motor cortex excitability. *Clin Neurophysiol.* doi:10.1016/j.clinph.2014.09.022. [Epub ahead of print].
- Flor H, Diers M, Andoh J. The neural basis of phantom limb pain. *Trends Cogn Sci.* 2013;17(7):307–8. doi:10.1016/j.tics.2013.04.007. Epub 2013 Apr 19.
- Lefaucheur JP. Transcranial magnetic stimulation in the management of pain. *Suppl Clin Neurophysiol.* 2004;57:737–48.
- Poreisz C, Csifcsak G, Antal A, Levold M, Hillers F, Paulus W. Theta burst stimulation of the motor cortex reduces laser-evoked pain perception. *Neuroreport.* 2008;19:193–6.
- Ihle K, Rodriguez-Raecke R, Luedtke K, May A. tDCS modulates cortical nociceptive processing but has little to no impact on pain perception. *Pain.* 2014;155:2080–7.
- Zaghi S, Thiele B, Pimentel D, Pimentel T, Fregni F. Assessment and treatment of pain with non-invasive cortical stimulation. *Restor Neurol Neurosci.* 2011;29:439–51.

16. Lefaucheur JP, Drouot X, Menard-Lefaucheur I, Keravel Y, Nguyen JP. Motor cortex rTMS restores defective intracortical inhibition in chronic neuropathic pain. *Neurol.* 2006;67:1568–74.
17. Fourkas ADA, Urgesi C, Aglioti SM. Corticospinal facilitation during first and third person imagery. *Exp Brain Res.* 2006;168:143–51.
18. Kasai TKS, Kawanishi M, Yahagi S. Evidence for facilitation of motor evoked potentials (MEPs) induced by motor imagery. *Brain Res.* 1997;744:147–50.
19. Izumi SFT, Ikai T, Andrews J, Daum M, Chino N. Facilitatory effect of thinking about movement on motor-evoked potentials to transcranial magnetic stimulation of the brain. *Am J Phys Med Rehabil.* 1995;74:207–13.
20. Beck AT, Steer RA. Internal consistencies of the original and revised beck depression inventory. *J Clin Psychol.* 1984;40:1365–7.
21. Liang N, Funase K, Takahashi M, Matsukawa K, Kasai T. Unilateral imagined movement increases interhemispheric inhibition from the contralateral to ipsilateral motor cortex. *Exp Brain Res.* 2014;232(6):1823–32. doi:10.1007/s00221-014-3874-4. Epub 2014 Feb 22.
22. Raffin E, Mattout J, Reilly KT, Giroux P. Disentangling motor execution from motor imagery with the phantom limb. *Brain.* 2012;135:582–95.
23. Raffin E, Giroux P, Reilly KT. The moving phantom: motor execution or motor imagery? *Cortex.* 2011;48:746–57.
24. Beaumont G, Mercier C, Michon PE, Malouin F, Jackson PL. Decreasing phantom limb pain through observation of action and imagery: a case series. *Pain Med.* 2011;12:289–99.
25. Flor H. Maladaptive plasticity, memory for pain and phantom limb pain: review and suggestions for new therapies. *Expert Rev Neurother.* 2008;8:809–18.
26. Zhuo M. Long-term potentiation in the anterior cingulate cortex and chronic pain. *Philos Trans R Soc Lond B Biol Sci.* 2013;369:20130146.
27. Volz MS, Suarez-Contreras V, Mendonca ME, Pinheiro FS, Merabet LB, Fregni F. Effects of sensory behavioral tasks on pain threshold and cortical excitability. *PLoS One.* 2013;8:e52968.
28. Volz MS, Mendonca M, Pinheiro FS, Cui H, Santana M, Fregni F. Dissociation of motor task-induced cortical excitability and pain perception changes in healthy volunteers. *PLoS One.* 2012;7:e34273.
29. Blackwell SE, Rius-Ottenheim N, Schulte-van Maaren YW, Carlier IV, Middelkoop VD, Zitman FG, et al. Optimism and mental imagery: a possible cognitive marker to promote well-being? *Psychiatry Res.* 2012;206:56–61.
30. Malloy KM, Milling LS. The effectiveness of virtual reality distraction for pain reduction: a systematic review. *Clin Psychol Rev.* 2010;30:1011–8.
31. Ossipov MH, Dussor GO, Porreca F. Central modulation of pain. *J Clin Invest.* 2010;120:3779–87.
32. Legrain V, Mancini F, Sambo CF, Torta DM, Ronga I, Valentini E. Cognitive aspects of nociception and pain: bridging neurophysiology with cognitive psychology. *Neurophysiol Clin.* 2012;42:325–36.
33. Dietrich C, Walter-Walsh K, Preissler S, Hofmann GO, Witte OW, Miltner WH, et al. Sensory feedback prosthesis reduces phantom limb pain: proof of a principle. *Neurosci Lett.* 2011;507:97–100.
34. Johnson MI, Bjordal JM. Transcutaneous electrical nerve stimulation for the management of painful conditions: focus on neuropathic pain. *Expert Rev Neurother.* 2011;11:735–53.
35. Volz MS, Medeiros LF, Tarragó Mda G, Vidor LP, Dall'Agnol L, Deitos A, et al. The relationship between cortical excitability and pain catastrophizing in myofascial pain. *J Pain.* 2013;14(10):1140–7. doi:10.1016/j.jpain.2013.04.013. Epub 2013 Jun 27.
36. Mhalla A, de Andrade DC, Baudic S, Perrot S, Bouhassira D. Alteration of cortical excitability in patients with fibromyalgia. *Pain.* 2010;149:495–500.
37. Schwenkreis P, Janssen F, Rommel O, Pleger B, Volker B, Hosbach I, et al. Bilateral motor cortex disinhibition in complex regional pain syndrome (CRPS) type I of the hand. *Neurol.* 2003;61:515–9.
38. Schwenkreis P, Maier C, Tegenthoff M. Motor cortex disinhibition in complex regional pain syndrome (CRPS)-a unilateral or bilateral phenomenon? *Pain.* 2005;115:219–20. author reply 220–211.
39. Sullivan MJ, Lynch ME, Clark AJ. Dimensions of catastrophic thinking associated with pain experience and disability in patients with neuropathic pain conditions. *Pain.* 2005;113:310–5.
40. Schoth DE, Yu K, Lioffi C. The role of threat expectancy in attentional bias and thermal pain perception in healthy individuals. *J Health Psychol.* 2014;19(5):653–63. doi:10.1177/1359105313476976. Epub 2013 Mar 19.
41. Leung L. Pain catastrophizing: an updated review. *Indian J Psychol Med.* 2013;34:204–17.
42. Facco E, Stellini E, Bacci C, Manani G, Pavan C, Cavallin F, et al. Validation of visual analogue scale for anxiety (VAS-A) in preanesthesia evaluation. *Minerva Anestesiol.* 2013;79(12):1389–95. Epub 2013 Jul 9.
43. Tiffin J, Asher EJ. The Purdue pegboard; norms and studies of reliability and validity. *J Appl Psychol.* 1948;32:234–47.
44. Oldfield RC. The assessment and analysis of handedness: the Edinburgh inventory. *Neuropsychologia.* 1971;9:97–113.
45. Rossi S, Hallett M, Rossini PM, Pascual-Leone A. Safety, ethical considerations, and application guidelines for the use of transcranial magnetic stimulation in clinical practice and research. *Clin Neurophysiol.* 2009;120:2008–39.
46. Hattemer K, Knake S, Reis J, Rochon J, Oertel WH, Rosenow F, et al. Excitability of the motor cortex during ovulatory and anovulatory cycles: a transcranial magnetic stimulation study. *Clin Endocrinol (Oxf).* 2007;66:387–93.
47. Smith MJ, Keel JC, Greenberg BD, Adams LF, Schmidt PJ, Rubinow DA, et al. Menstrual cycle effects on cortical excitability. *Neurol.* 1999;53:2069–72.
48. Chesterton LS, Barlas P, Foster NE, Lundeberg T, Wright CC, Baxter GD. Sensory stimulation (TENS): effects of parameter manipulation on mechanical pain thresholds in healthy human subjects. *Pain.* 2002;99:253–62.
49. Paulus W, Classen J, Cohen LG, Large CH, Di Lazzaro V, Nitsche M, et al. State of the art: pharmacologic effects on cortical excitability measures tested by transcranial magnetic stimulation. *Brain Stimul.* 2008;1:151–63.
50. Vidor LP, Torres IL, Medeiros LF, Dussan-Sarria JA, Dall'agnol L, Deitos A, et al. Association of anxiety with intracortical inhibition and descending pain modulation in chronic myofascial pain syndrome. *BMC Neurosci.* 2014;15:42.
51. Nardone R, Höller Y, Brigo F, Orioli A, Tezzon F, Schwenker K, Christova M, Golaszewski S, Trinka E. Descending motor pathways and cortical physiology after spinal cord injury assessed by transcranial magnetic stimulation: a systematic review. *Brain Res.* 2014. doi:10.1016/j.brainres.2014.09.036. [Epub ahead of print].
52. Filipovic SR, Ljubisavljevic M, Svetel M, Milanovic S, Kacar A, Kostic VS. Impairment of cortical inhibition in writer's cramp as revealed by changes in electromyographic silent period after transcranial magnetic stimulation. *Neurosci Lett.* 1997;222:167–70.
53. Kojima S, Onishi H, Sugawara K, Kirimoto H, Suzuki M, Tamaki H. Modulation of the cortical silent period elicited by single- and paired-pulse transcranial magnetic stimulation. *BMC Neurosci.* 2013;14:43.
54. Richter MA, de Jesus DR, Hoppenbrouwers S, Daigle M, Deluce J, Ravindran LN, et al. Evidence for cortical inhibitory and excitatory dysfunction in obsessive compulsive disorder. *Neuropsychopharmacology.* 2011;37:1144–51.
55. Heide G, Witte OW, Ziemann U. Physiology of modulation of motor cortex excitability by low-frequency suprathreshold repetitive transcranial magnetic stimulation. *Exp Brain Res.* 2006;171:26–34.
56. Desrosiers J, Hebert R, Bravo G, Dutil E. The Purdue pegboard test: normative data for people aged 60 and over. *Disabil Rehabil.* 1995;17:217–24.
57. Mack JL. Validity of the Purdue pegboard as a screening test for brain damage in a psychiatric population. *Percept Mot Skills.* 1969;28:832–4.
58. Nielsen H, Knudsen L, Daugbjerg O. Normative data for eight neuropsychological tests based on a Danish sample. *Scand J Psychol.* 1989;30(1):37–45.

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