Graded motor imagery for pathologic pain
A randomized controlled trial
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Abstract—Background: Phantom limb and complex regional pain syndrome type 1 (CRPS1) are characterized by changes in cortical processing and organization, perceptual disturbances, and poor response to conventional treatments. Graded motor imagery is effective for a small subset of patients with CRPS1. Objective: To investigate whether graded motor imagery would reduce pain and disability for a more general CRPS1 population and for people with phantom limb pain. Methods: Fifty-one patients with phantom limb pain or CRPS1 were randomly allocated to motor imagery, consisting of 2 weeks each of limb laterality recognition, imagined movements, and mirror movements, or to physical therapy and ongoing medical care. Results: There was a main statistical effect of treatment group, but not diagnostic group, on pain and function. The mean (95% CI) decrease in pain between pre- and post-treatment (100 mm visual analogue scale) was 23.4 mm (16.2 to 30.4 mm) for the motor imagery group and 10.5 mm (19 to 19.2 mm) for the control group. Improvement in function was similar and gains were maintained at 6-month follow-up. Conclusion: Motor imagery reduced pain and disability in these patients with complex regional pain syndrome type I or phantom limb pain, but the mechanism, or mechanisms, of the effect are not clear.

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Complex regional pain syndrome type 1 (CRPS) is considered a pathologic pain syndrome because the pain does not seem to reflect the underlying tissue pathology. However, the pathophysiology of CRPS1 is not well understood: peripheral and central changes have been observed and altered central representation of perceptual, motor, and autonomic systems have been implicated. If such cortical mechanisms underpin the disease, it would seem reasonable to target them in treatment—“train the brain.” One such approach, graded motor imagery, reduced pain and disability in a relatively homogenous sample of patients with CRPS1 after wrist fracture, all of whom had motor dysfunction as part of their condition. Although those clinical trials appear encouraging, about 50% of subjects were excluded, so whether the approach is effective for a wider CRPS1 population is not known. The first aim of the current study was to resolve this issue.

Graded motor imagery might also be effective in those with phantom limb pain, which is also considered a pathologic pain syndrome and is also thought to be dominated by altered cortical mechanisms. The similarities between phantom limb pain and CRPS1, which have been noted elsewhere, suggest that graded motor imagery may be effective for phantom limb pain as well as CRPS1. The second aim of the current study was to determine if this is the case.

Methods. Design. A single blind randomized trial with 6-month follow-up was conducted (figure 1).

Eligible participants were drawn from three patient groups: patients with phantom limb pain after amputation of one limb, phantom limb pain (within deafferented zone, according to results of previous quantitative sensory testing, not verified here) after
brachial plexus avulsion injury of one arm, and patients with complex regional pain syndrome, type 1 (CRPS1). Sixty-nine eligible patients (37 F) were contacted via hospital physiotherapy department, neurology, and pain clinic waiting lists. Patients were excluded if they had been diagnosed with any other neurology, psychopathology, or muscle disorder or dyslexia; had difficulty performing a rapid naming task; were visually impaired; had any other limb pathology or pain; or lived outside the immediate metropolitan area of the host department. Nine patients were excluded according to those criteria. Nine patients with CRPS1 who did not fulfill recognized diagnostic criteria were also excluded, which left 51 patients (32 F). This sample size would detect an appropriate bank of images classified by each subject into one of four categories, according to the level of pain that would be imparted by the control treatment. An independent investigator who was blind to group and assessment occasion. All assessments were undertaken at prerandomization and at 6 weeks (completion of the treatment period). Pain VAS and function NRS were also undertaken at 6 months follow-up.

**Active treatment—motor imagery program.** The first 2 weeks were the limb laterality recognition phase. Forty photographs of a right hand, matched to gender, and in various positions and alignments, were digitally mirrored to create a bank of 80 images. Twenty-four photographs of a right foot, matched to gender, and in various positions and alignments, were digitally mirrored to create a bank of 48 images. Thus, there were four banks of images: upper and lower limbs, male and female. Every image in the appropriate bank of images was classified by each subject into one of four categories, according to the level of pain that would be imparted by the control treatment. An independent investigator who was blind to group and assessment occasion. All assessments were undertaken at prerandomization and at 6 weeks (completion of the treatment period). Pain VAS and function NRS were also undertaken at 6 months follow-up.

**Table 1. Protocol for training load during each phase of the motor imagery program.**
Subjects. All data were collected over 32 months. One female subject in the control group withdrew from the study because she sustained an unrelated injury. There were no other dropouts or withdrawals. Subject characteristics according to group are shown in Table E-1 on the Neurology Web site at www.neurology.org. There was no difference between the groups by gender (p = 0.56 by x² test), age, duration of disease, VAS pain score, or NRS score (p > 0.50 for all, by two-sample t-tests).

Pain and function at post-program. Statistical results are described here, NNTs in Table 2 and results for specific diagnoses are presented in Figure 2. The regressions showed effects for pain VAS [ANCOVA F(3,46) = 6.77, p = 0.001] and for function NRS [ANCOVA F(3,46) = 8.95, p < 0.001]. For pain intensity at post-program, there was a

Table 2 Mean (95% CI) for number needed to treat (NNT) to achieve a preset change in pain or function or both

<table>
<thead>
<tr>
<th>Response at post-program</th>
<th>NNT to get a 50% decrease in pain</th>
<th>NNT to get a 4-point increase in function</th>
<th>NNT for both criteria</th>
</tr>
</thead>
<tbody>
<tr>
<td>Response at 6-mo. follow-up</td>
<td>3 (2-6)</td>
<td>4 (2-11)</td>
<td>4 (2-17)</td>
</tr>
</tbody>
</table>

Pain = average pain intensity over previous 2 days; Function = ability to perform five patient-selected activities.

Because one proposed mechanism of effect of graded motor imagery involves normalization of cortical organization, which in turn is related to duration of symptoms, a secondary analysis investigated the relationship between the duration of symptoms and the response to treatment, via two linear regressions between duration and change in pain VAS at post-treatment and at follow-up. For all analyses, significance was set at α = 0.05.
main effect of treatment group (unstandardized B = −1.298, p = 0.002). That is, the mean (95% CI) decrease in average pain over the last 2 days, as measured on the 100 mm VAS, was 23.4 mm (16.2 to 30.4 mm) for the MIP group and 10.5 mm (1.9 to 19.2 mm) for the control group.

For function NRS, there was a main effect of treatment group (unstandardized B = 1.532, p = 0.001). That is, the mean (95% CI) increase in average score for each patient-specific task, measured using a 0 to 10 NRS, was 2.2 points (1.3–3.0 points) for the MIP group and 0.6 points (0.2–1.0 points) for the control group.

**Pain and function at 6-month follow-up.** The regressions showed effects for pain VAS [ANCOVA F(3,46) = 8.701, p < 0.001] and for function NRS [ANCOVA F(3,46) = 7.327, p = 0.001]. For pain VAS at follow-up, there was a main effect of treatment group (unstandardized B = −2.07, p = 0.001) (figure 2B). That is, the mean (95% CI) decrease in pain VAS between pretreatment and follow-up was 32.1 mm (23.8–40.3 mm) for the MIP group and 11.6 mm (2.4–20.7 mm) for the control group. The effect size at 6-month follow-up, but not at post-program, was consistent with the estimated effect size.

For function NRS, there was a main effect of treatment group (unstandardized B = 2.18, p = 0.001) (figure 2B). That is, the mean (95% CI) increase in task-specific NRS VAS between pretreatment and follow-up was 3.7 points (2.7–4.6 points) for the MIP group and 1.5 points (1–2.2 points) for the control group.

**Outcome and response to treatment at follow-up: Pain VAS ≤ 3, function NRS ≥ 5, or both.** NNT was calculated for pain VAS ≤ 3, function NRS ≥ 5, and for both, and are shown, with 95% CI, in table 2. The duration of symptoms did not relate to change in pain VAS at post-treatment (p = 0.224), nor to change in pain VAS at follow-up (p = 0.071).

**Participation in home program.** Participation with the home program was 75% throughout the treatment period. There was no difference between treatment groups in any phase (p > 0.211 for all).

**Treatment during follow-up period.** Only 11 patients in the treated group, but all patients in the control group, reported that they sought treatment for their pain during the follow-up period (p < 0.001 by χ² test). For the treated group, this constituted between 3 and 11 (median = 6) physiotherapy treatments, including motor imagery, tactile discrimination training, and functional exposure (9 patients), a multidisciplinary pain management program (1 patient), and between 2 and 4 general practitioner visits (3 patients). For the control group, treatments sought were physiotherapy treatment (motor imagery, desensitization, tactile discrimination, functional exposure) for 14 patients (median number of treatments = 12, range 1 to 18), multidisciplinary pain management program for 7 patients, general practitioner visits for 10 patients (median number of visits = 6, range 2 to 1), spinal cord stimulator for 1 patient.

**Discussion.** Graded motor imagery reduced pain and disability in a wider CRPS1 population and in those with phantom limb pain after amputation or brachial plexus avulsion injury. The following results support this position: 1) a significant effect of treatment group on both primary outcome measures (pain and function), such that pain decreased and function increased for the motor imagery group, relative to the control group; 2) NNTs for response to treatment at 6 months of about 3.

Graded motor imagery has been shown to reduce pain and disability in a relatively homogeneous group of patients with chronic CRPS1 and mirror therapy alone has shown efficacy for those with acute CRPS1. The current data corrobore those studies, although the mean magnitude of pain reduction was about 50% less in the current work than it was in the earlier studies using graded motor imagery. A likely contributor to the reduced mean effect is the relative heterogeneity of the current sample. First, phantom limb pain and CRPS1, although both categorized here as pathologic pain disorders, and although changes in cortical organization and some clinical findings are common to both, are fundamentally different: whereas CRPS usually occurs after minor injury and involves no demonstrable nerve injury, phantom limb pain occurs after major trauma and undeniable nerve injury. Further to that, phantom pain after brachial plexus avulsion injury may involve different mechanisms to phantom pain after amputation. That is, the current design may conceal stronger effects in one group than another and that the effect occurred regardless of diagnostic group does not imply that the same mechanisms underpin that effect in each group. This study was underpowered to systematically evaluate different response profiles between the diagnostic groups.

Even the current sample of patients with CRPS1 was more heterogeneous than previous samples: previous studies limited inclusion to CRPS1 initiated by wrist fracture, limited disease duration to more than 6 months, and excluded those without motor signs and symptoms. The final issue is particularly important because diagnosis of CRPS1 is based on presenting signs and symptoms, not on mechanisms, and many contributing mechanisms exist. Perhaps subclasses of CRPS1 exist and some are better suited to motor imagery training than others.

The control group here was intended to reflect current practice, which is dominated by drug and physical therapies, implemented according to a cognitive-behavioral model. That the medical component of the control group treatment constituted ongoing medical care and therefore was not new probably reduced its effect because most treatment is effective to some extent when it is first started. Importantly, however, the commencement of conventional physiotherapy in addition to ongoing medical care had no discernible effect on pain or function. This seems discouraging, although the current study limited the duration and frequency of physical therapy, which suggests that firm conclusions are probably premature.

Although the patients used here were broadly similar to those involved in other studies including therapeutic trials, it is not possible to di-
rectly compare the current results to established data from CRPS1 or phantom limb pain because robust trials of other therapies are lacking. This lack of data probably explains why tricyclic antidepressants, antiepileptics, and opioids remain the mainstay of treatment, particularly with phantom limb pain—these drugs have demonstrated some efficacy in other types of neuropathic pain. That said, the demonstrated effect of such drugs in those groups is not better than the current results. For example, a systematic review of drug therapies for peripheral neuropathic pain reported NNTs of between 2.2 and 5.0 for tricyclics, antiepileptics, and opioids, and between 2.9 and 7.7 for selective serotonin reuptake inhibitors. The present study found NNTs of about 3 for similar outcome parameters.

How progressive motor imagery reduced pain and disability is not known. This study was based on the idea of applying established principles of rehabilitation to training the brain. That limb laterality recognition activates premotor cortices but not primary motor cortex, whereas imagined movements activate both, and that the order of hand laterality recognition, imagined movements, and mirror movements seems to be important in the effect motor imagery, seem consistent with this possibility. Perhaps practicing laterality recognition is to limb movement as practicing knee movement is to walking. This seems sensible in principle, but the neural mechanisms by which it might occur are unknown.

An alternative possible explanation for the effect of motor imagery is that it promotes sustained attention to the affected limb. That probably does not explain the effect in patients with CRPS1 post-wrist fracture, because undertaking the treatment components in a different order does not work, but it may explain the effect observed here. Such an explanation would seem consistent with several findings in the literature. For example, patients with CRPS1 and with phantom limb pain take longer to recognize the laterality of a pictured limb if it corresponds to their own affected limb. Acute experimental pain, and the expectation of acute experimental pain, delays recognition of the laterality of the opposite limb, which implies an information processing bias toward the body part in pain, yet patients with CRPS1 and phantom limb pain show the opposite effect, which implies an opposite information processing bias—away from the body part in pain. If so, progressive motor imagery may simply serve to reverse that tendency. That speculation remains to be verified, but other signs and symptoms in CRPS1 and after amputation or brachial plexus avulsion are consistent with some sort of neglect: symptoms of foreignness and neglect, the perception that their affected limb is bigger than it is, which also occurs after anesthesia.

The question, then, is why would a neglect-like disorder cause pain? Pain is not characteristic of the usual parietal neglect syndromes. The cortical model of pathologic pain is relevant here because it poses that the changes to cortical proprioceptive representation, themselves related to the neglect-like phenomena outlined above, underpin the pain because they evoke an incongruence between motor commands and sensory feedback. It is established that 1) cortical proprioceptive representation is altered in CRPS1 and in phantom limb pain, 2) that the extent of reorganization relates to the duration of symptoms (it is notable that although the current study did not find a relationship between duration of symptoms and response to motor imagery, it was probably underpowered), and 3) that resolution of symptoms is associated with normalization of cortical reorganization. There is a large amount of literature on sensory-motor incongruence, typified by the reafore principle, whereby an exact copy of the command for movement is subtracted from sensory input about the actual movement to yield an error signal. However, only recently has sensory-motor incongruence been empirically investigated with regard to pain, and there are insufficient data to conclude whether motor imagery might remove sensory-motor incongruence. If it does, this mechanism may also mediate the effect of sensory discrimination training in phantom limb pain and of sensory-motor retuning in CRPS1, for which there is preliminary evidence. Alternatively, perhaps the neglect-like phenomena and the pain are not causally related but both result from, for example, altered central representations of somatic input in the thalamus or cortex, in which case motor imagery might primarily serve to normalize these central representations. Thus, although evidence is emerging that treatments such as graded motor imagery and sensory discrimination training can be effective for pathologic pain, further studies are needed to replicate the current data and elucidate the mechanisms involved.

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References