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Think to Move: a Neuromagnetic Brain-Computer Interface (BCI) System for Chronic Stroke

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Abstract

Background and Purpose—Stroke is a leading cause of long-term motor disability among adults. Present rehabilitative interventions are largely unsuccessful in improving the most severe cases of motor impairment, particularly in relation to hand function. Here we tested the hypothesis that patients experiencing hand plegia as a result of a single, unilateral subcortical, cortical or mixed stroke occurring at least 1 year previously, could be trained to operate a mechanical hand orthosis through a brain-computer interface (BCI).

Methods—Eight patients with chronic hand plegia resulting from stroke (residual finger extension function rated on the Medical Research Council scale = 0/5) were recruited from the Stroke Neurorehabilitation Clinic, Human Cortical Physiology Section of the National Institute for Neurological Disorders and Stroke (NINDS) (n = 5) and the Clinic of Neurology of the University of Tübingen (n = 3). Diagnostic MRIs revealed single, unilateral subcortical, cortical or mixed lesions in all patients. A magnetoencephalography-based BCI system was used for this study. Patients participated in between 13 to 22 training sessions geared to volitionally modulate μ rhythm amplitude originating in sensorimotor areas of the cortex, which in turn raised or lowered a screen cursor in the direction of a target displayed on the screen through the BCI interface. Performance feedback was provided visually in real-time. Successful trials (in which the cursor made contact with the target) resulted in opening/closing of an orthosis attached to the paralyzed hand.

Results—Training resulted in successful BCI control in 6 of 8 patients. This control was associated with increased range and specificity of μ rhythm modulation as recorded from sensors overlying central ipsilesional (4 patients) or contralesional (2 patients) regions of the array. Clinical scales used to rate hand function showed no significant improvement after training.

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Disclosures

None.

Conclusions—These results suggest that volitional control of neuromagnetic activity features recorded over central scalp regions can be achieved with BCI training after stroke, and used to control grasping actions through a mechanical hand orthosis.

Keywords

brain-computer interface; MEG; motor; plasticity; stroke

One year after stroke a third of all affected patients have poor or nonexistent residual hand motor function despite intensive treatment and rehabilitation.¹ Significant functional recovery after this initial year is rare^{2,3} despite novel interventional approaches recently applied in the chronic stage, like bilateral arm training or constraint-induced movement therapy.^{4,5} These treatments are based on the ability of the patient to perform actions with the affected arm or hand, and therefore require a moderate degree of residual motor function. There are, however, many patients who do not have such residual function and therefore cannot use the plegic hand at all for training purposes. At present, there is no treatment available for this condition.

Over the past 15 years, an increasing number of brain-computer interface (BCI) systems have been developed.⁶ All of these systems record, decode, and ultimately translate some measurable neurophysiological signal into an effector action or behavior. Existing BCI systems have used invasive microelectrode arrays to record single-unit spiking activity and local field potentials,^{7,8} and subdural electrode arrays to record electrocorticography.⁹ Noninvasive signal recording approaches have used electroencephalography (EEG),¹⁰ magnetoencephalography (MEG),¹¹ blood-oxygen-level dependent functional MRI,¹² and near infrared spectroscopy.¹³ End-user applications developed for human BCI systems have included 1-D, 2-D, and 3-D (point-and-click) screen cursor control, as well as spelling software for communication. In addition, using microelectrode implants in various cortical areas, monkeys have been trained to control robotic arms for reaching and grasping during feeding.¹⁴ A few attempts have been made to apply these technologies to patient groups, with these attempts primarily focusing on patients with amyotrophic lateral sclerosis or tetraplegia.^{6,10}

Here, we describe a BCI system that uses MEG activity evoked by the patient's intent to move a completely paralyzed hand to control grasping motions of a mechanical orthosis attached to the affected hand.

Methods

Patients

Eight patients with chronic hand plegia resulting from stroke were recruited from the Stroke Neurorehabilitation Clinic, Human Cortical Physiology Section of the National Institute for Neurological Disorders and Stroke (NINDS) (n = 5; mean age=58.2 ± 7.0 years; mean hand plegia duration=25.2 ± 11.6 months) and the Clinic of Neurology of the University of Tübingen (n = 3; mean age=41.7 ± 26.6 years; mean hand plegia duration=16.7 ± 6.4 months). One patient recruited from the University of Tübingen, patient MD, experienced a

pediatric stroke. Patients were included in the study if they had a history of a single stroke, with residual finger extension weakness rated as 0/5 on the Medical Research Council (MRC) scale (Table). Therefore, all patients included in this study were completely unable to induce any voluntary movements in extensors of the plegic hand. Spasticity of shoulder, elbow and finger flexors and extensors was rated as 3 or less on the Modified Ashworth Scale¹⁵ in eligible patients to ensure that their arm could maintain a comfortable posture while seated in the MEG chair, and that their fingers could be passively manipulated by the hand orthosis (Figure 1). Medical and neurological screening history and examination excluded patients with major cognitive deficits (Folstein Mini Mental Status Test¹⁶ lower than 23), major depressive disorder, or other uncontrolled illness. Anatomic MRI of the brain was used to exclude patients with cerebellar or brain stem lesions, but otherwise, lesions of the suprapontine corticospinal tract of varying sizes and extent were included as long as they resulted in a plegic hand. Patients provided written informed consent and the study was approved by the Institutional Review Board of the NINDS and the Ethical Committee of the Faculty of Medicine of the University of Tübingen.

BCI Training

MEG Recordings—Neuromagnetic activity recorded from a 275-channel (6 patients) or 153-channel (2 patients) MEG array (VSM Medtech) was used to control a BCI as previously described,¹¹ at both the NIH and the University of Tübingen. Both MEG apparatuses were housed in a magnetically shielded room and used synthetic 3rd gradient balancing to reduce interference from environmental noise. Recordings from all MEG channels were antialiased with a 200 Hz cut-off, low-pass filter, and digitally sampled at 600 Hz.

During recording, patients were seated alone in the shielded MEG room with the lights slightly dimmed, and their head centrally positioned within the sensor array. A closed-circuit video system was used to constantly monitor the patients, while instructions were given during rest periods via an intercom system. Patients were instructed to refrain from extraneous movement while engaged in experimental tasks (especially with the healthy arm) to reduce artifacts. Adherence to these instructions was monitored during recording via the video system, as well as online electromyography recordings obtained from the brachioradialis muscle of both arms.

MEG was chosen to drive this initial proof of principle study of BCI in chronic stroke because of its noninvasiveness and exquisite spatial and temporal resolution. Two additional features of MEG made it desirable relative to EEG. The magnetic fields generated by brain activity are minimally distorted by brain lesions, making MEG particularly appropriate for studies in stroke.^{17,18} Furthermore, the collection of MEG data does not require the attachment of scalp electrodes or related cleaning procedures used to reduce electrode impedance. This latter point in particular allowed patients to start the task rapidly after arriving at the laboratory, avoiding the fatigue inherent to long periods of experimental preparation. Together, these features made MEG an ideal source of on-line recording and localization of dynamic cortical rhythm changes.

μ Rhythm-Based BCI—Amplitude modulation of the μ rhythm was used to control this BCI system,¹¹ which was based on the BCI2000 software platform¹⁹ (www.bci2000.org). The μ rhythm is typically found over the sensorimotor cortex with a base frequency of 9 to 12 Hz. Its arc-shaped waveform includes a strong first harmonic in the β band at 20 to 24 Hz. The terms synchronization and desynchronization are commonly used to describe increases and decreases in μ rhythm amplitude relative to some baseline, respectively. μ rhythm desynchronization has been observed during the planning, execution, or even imagination of limb movements.^{20–23} In particular, the substantial and relatively somatotopic μ rhythm amplitude modulation observed during engagement in motor imagery tasks made it appealing for use in stroke patients, who could perform imagined movements or even attempt to move their plegic hand in the absence of any motor function.

During BCI training, μ rhythm amplitude estimates were derived from 3 to 4 MEG-sensors from the array (Table). The cluster(s) of MEG sensors chosen as BCI controllers were identified after an initial session (described in detail below) during which subjects imagined grasping movements of the plegic hand. The sensors chosen for BCI control were the ones that showed the highest correlated modulation of μ rhythm amplitude between conditions (see supplemental data for detailed description, available online at <http://stroke.ahajournals.org>).

Hand Orthosis—During all BCI training sessions, a mechanical orthosis was attached to the plegic hand. Fingers 2 to 5 (index, middle, ring, and little fingers) were individually inserted into ring-like fasteners that grasped each digit at the first phalanx, and fixated by a screw-adjustable shoe to prevent slippage. Each fastener was connected to a plastic Bowden cable that allowed for hand grasping or hand opening motions. In order to minimize magnetic artifacts in the MEG environment, these cables were extended and retracted by opening and closing computer-controlled pneumatic valves. The orthosis had 2 possible movement motions: flexion or extension of fingers 2 to 5 in a hand grasping or hand opening fashion, respectively (Figure 1). All 4 fingers were synchronously moved in the same direction. The orthosis actions were synchronized with the BCI training task described below through parallel port communication with a custom control circuit.

Real-Time Feedback and BCI Training Task—During each session, patients performed between 150 to 250 trials of a goal-oriented, visual feedback task (supplemental Figure I, available online at <http://stroke.ahajournals.org>). The task was designed to help them achieve volitional control of μ rhythm amplitude, and thus control of the orthosis.

Each trial was initiated by the presentation of a target on either the upper or lower half of the right side of a visual display (Figure 1). The target was a visual representation of an acceptable range of μ rhythm amplitudes for the desired orthosis action. A square screen-cursor would then begin moving at a fixed rate from left-to-right across the display, with the cursor feedback updated every 300 ms. The vertical height of the cursor was a transformation of the recorded μ rhythm amplitude. The goal for the patient was to volitionally modulate the μ rhythm amplitude in such a way so that the cursor contacted the target once it reached the right edge of the screen. The BCI software maintained a history of the mean μ rhythm amplitude estimate from each trial and assigned this to a distribution

representing observations for each target (or orthosis action) condition. The classification threshold, defined as the midpoint between the means of these 2 distributions, was adaptive to account for changes in the shapes of these distributions over the course of training.

At the conclusion of each trial in which the patient was successful at producing the appropriate modulation of μ rhythm amplitude (meaning the cursor hit the target), a simultaneous change in target color (red to yellow) and orthosis action occurred providing reinforcement (Figure 1). If the cursor failed to hit the target, no reinforcement was provided (meaning no orthosis manipulation of hand posture occurred).

μ Rhythm Amplitude/Orthosis Action Coupling—The coupling of μ rhythm synchronization/desynchronization to the resulting orthosis action was determined by patient preference. Patients generally achieved μ rhythm synchronization via passive relaxation imagery, and desynchronization via motor imagery of some hand action. In 5 of the 8 patients, opening of the hand by the orthosis was associated with μ rhythm synchronization, whereas grasping motions were associated with desynchronization. The other 3 patients chose the opposite coupling. This alternative choice was most likely related to the greater degree of spasticity present in muscles of the affected hand and arm of these patients, as their hands normally displayed a more grip-like posture in their passive state. Thus, all patients chose to relate the more passive form of imagery with their passive hand posture state.

Experimental Design

Initial Session—In the initial session, which lasted for approximately 1 hour, patients were familiarized with the MEG environment as well as with the hand orthosis. They sat upright fixating a screen located 50 cm in front of their eyes. During that period of time, they were instructed to perform the following tasks in a randomized order: (1) repeated grasping motions of the intact hand at 0.5 Hz rate guided by a visual metronome stimulus on the screen, (2) motor imagery of the same hand motions without actually moving the intact hand, (3) motor imagery of comparable movements of the plegic hand, and (4) fixation of the metronome stimulus in a “resting” state. The instructions were displayed on the screen in front of the patients with trials separated by 2 second intervals. Twenty-four trials were recorded for motor movements and 48 trials were recorded for imagined movements during this initial session. As stated above, these data were used to determine parameters for the subsequent BCI-training sessions.

BCI Training Sessions—Over the course of approximately 3 to 8 weeks, patients participated in 13 to 22 training sessions (separated by at least 24 hours; Table). The training frequency was highly determined by each patient’s tolerance to fatigue, or additional time commitments, and ranged between 3 to 5 times per week (Table). During these sessions, they performed the training task described above. Each training session lasted 1 to 2 hours and was implemented on an outpatient basis at both locations.

Data Analysis

Behavioral Data—The success rate (the proportion of trials in which patients were successful at contacting the target with the cursor, or alternatively, producing the requested μ rhythm amplitude modulation) was computed for trial presentations during a single training session, and used as a performance measure. A trial was considered a “success” when the cursor arrived at the requested target over the time of a trial (see supplemental Figure I). A significant hit-rate increase from “chance” levels of 50% indicated that volitional control of μ rhythm modulation at the desired MEG sensor locations was achieved (Figures 2 and 3A).

MEG Data—Off-line analysis of all training sessions included computation of spectral power differences (Figure 3B) and statistical maps of μ rhythm amplitude correlations with target condition/orthosis action,²⁴ for each MEG sensor and frequency band. Topographical maps plotted for a single frequency band use spatial information about the location of areas displaying more prominent and consistent synchronization/desynchronization patterns between task conditions (Figure 3C).

Statistical Analysis—The Wilcoxon signed-rank test, the nonparametric homologue to the paired Student *t* test, was used to compare changes in group performance during training.²⁵ To assess changes in individual patient performance during training, the change-point test was used.²⁵ The change-point test assumes the null hypothesis that no time trend exists in the series of performance data. Based on this assumption, each session performance should rank on average near the median, and the cumulative sum of ranks should increase approximately linearly with session number. The maximal deviation from this expected linear increase in rank is considered as a potential “change-point” and is used to divide the time series into 2 components. These components are then compared using a Kolmogorov-Smirnov test to determine statistical significance.

Results

All 8 patients at the 2 sites, NIH and University of Tübingen, successfully completed the study. On average, the group performance rate improved with training to $72.48 \pm 18.36\%$ (median \pm interquartile range) during the final session (Figure 2), as compared to an initial median performance rate of $52.84 \pm 20.59\%$ (paired Wilcoxon signed rank test; $P < 0.05$).

Despite this general and encouraging improvement, there was substantial variability in the ability of different patients to improve their hit-rate (and consequently their μ rhythm desynchronization control, Figure 3). A majority of patients showed exponential performance increases during training with variable growth rates and delays of onset. After 15 training sessions, patient BA did not show any increase in performance, which remained near chance levels of 50%. In contrast, patient WF showed initial high rates of performance above 80% that then declined before becoming stable at approximately 70%. The reason for this decrease in performance is not known.

The majority of patients displayed μ rhythm desynchronization in the 3 Hz-wide control frequency band (9 Hz central frequency for PA, 12 or 12.5 Hz for all other patients) for the “grasping” orthosis action. Furthermore, these differences were greatest in areas surrounding

the control sensor locations (with the exceptions of BW who showed minimal modulation and MD, whose stroke occurred at a very early age, who showed diffuse modulation across the majority of the array). Collectively, the R^2 statistical maps reveal that modulation of the trained μ -rhythm feature was more strongly related to the task in segregated regions of the array that surrounded the control sensor locations (Figure 3C). Four of the 8 participants were able to achieve voluntary μ -rhythm control over central ipsilesional regions of the array, whereas 3 participants achieved control using sensors from central contralesional areas of the array. Post-training MRC scores of finger extension strength remained 0/5 for all patients, indicating that the training had no effect on gross hand motor function.

Discussion

These data demonstrate that most patients with chronic stroke and complete hand paralysis, in this small sample, can learn to modulate μ rhythm amplitude to achieve binary control of an orthosis that passively manipulates the grasping posture of the plegic hand. Furthermore, this control can be achieved using MEG signals recorded over the ipsilesional hemisphere. Patients achieved success rates that varied between 65% and 90% by the end of the 13 to 22 sessions training period, and 6 of the 8 patients showed a significant performance improvement over this period. Of the 2 patients (WF and BW) who did not show improvements, patient WF displayed a high success rate at the outset of training (86%) and his failure to improve may be attributed to a ceiling effect. This surprising degree of voluntary control of cortical rhythms, despite predominantly extensive subcortical lesions that in the case of 4 of the 8 patients expanded into cortical tissue, suggests that this strategy could be effective in patients with various lesion types.

It should be kept in mind that for this study, we only included patients that were completely unable to move the paretic hand because they represent the patient group with very few rehabilitation options available. The BCI approach was used here to induce hand grasping and opening in patients unable to elicit voluntary movements because of the potential importance of these motions for activities of daily living.²⁶ More studies are clearly needed to evaluate the extent to which our conclusions apply to patients with different lesion locations, extension, etiology, or even chronicity. We also do not know if the behavioral gains demonstrated in this study consolidate over time, or fade in the absence of constant reinforcement, as has been reported in some motor learning paradigms in healthy subjects.²⁷ Although these represent important areas of future research, our results clearly indicate that voluntary control of μ rhythm amplitude recorded over central cortical regions (either ipsilesional or contralesional) can be used to control biphasic grasping motions of a plegic hand through a hand orthosis.

Although present MEG technology is not practical for long-term or portable brain control of an orthosis, our results suggest that similar control may be achieved with EEG. Recording μ rhythm from 3 to 4 MEG sensor sites was successful in driving the BCI orthosis capable of a bimodal grasping-opening of a completely paralyzed hand. It is then theoretically possible that properly referenced EEG electrodes placed on these crucial locations could be similarly successful in driving the BCI. If so, there is the potential that relatively inexpensive and

portable EEG-orthosis systems could be developed in the future to operate in home or chronic care settings.

Two forms of BCI systems have been described in the past in humans: invasive⁸ and noninvasive.²⁸ Noninvasive approaches, comparable to ours, have been used to allow communication in locked-in or severely paralyzed amyotrophic lateral sclerosis patients²⁹ and after tetraplegia,³⁰ but to our knowledge not after stroke. Our findings now provide conclusive data demonstrating the potential usefulness of BCI-like approaches in patients with severe motor disability resulting from stroke.

Although the full analysis of the electrophysiological data recorded over the training period in our patients is clearly beyond the boundaries of this report, it is likely that this form of imagery training led to cortical reorganization in our patients, consistent with previous findings.³¹

In summary, these results demonstrate that patients with chronic stroke and complete hand paralysis can learn to control μ rhythm synchronization and desynchronization through motor imagery of the paralyzed hand. Harnessing the cortical activity generated by such imagery through a noninvasive BCI device, can then be used to elicit hand grasping/opening motions of an orthosis attached to the paralyzed hand.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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References

1. Lai SM, Studenski S, Duncan PW, Perera S. Persisting consequences of stroke measured by the stroke impact scale. *Stroke*. 2002; 33:1840–1844. [PubMed: 12105363]
2. Duncan PW, Goldstein LB, Horner RD, Landsman PB, Samsa GP, Matchar DB. Similar motor recovery of upper and lower extremities after stroke. *Stroke*. 1994; 25:1181–1188. [PubMed: 8202977]
3. Duncan PW, Lai SM. Stroke recovery. *Topics in Stroke Rehabilitation*. 1997; 4:51–58.
4. Wolf SL, Winstein CJ, Miller JP, Taub E, Uswatte G, Morris D, Giuliani C, Light KE, Nichols-Larsen D. Effect of constraint-induced movement therapy on upper extremity function 3 to 9 months after stroke: The EXCITE randomized clinical trial. *JAMA*. 2006; 296:2095–2104. [PubMed: 17077374]
5. Whitall J, McCombe Waller S, Silver KH, Macko RF. Repetitive bilateral arm training with rhythmic auditory cueing improves motor function in chronic hemiparetic stroke. *Stroke*. 2000; 31:2390–2395. [PubMed: 11022069]

6. Birbaumer N, Cohen LG. Brain-computer interfaces: communication and restoration of movement in paralysis. *J Physiol.* 2007; 579:621–636. [PubMed: 17234696]
7. Nicoletis MA. Actions from thoughts. *Nature.* 2001; 409:403–407. [PubMed: 11201755]
8. Hochberg LR, Serruya MD, Friehs GM, Mukand JA, Saleh M, Caplan AH, Branner A, Chen D, Penn RD, Donoghue JP. Neuronal ensemble control of prosthetic devices by a human with tetraplegia. *Nature.* 2006; 442:164–171. [PubMed: 16838014]
9. Leuthardt EC, Miller KJ, Schalk G, Rao RP, Ojemann JG. Electrocorticography-based brain computer interface—the Seattle experience. *IEEE Trans Neural Syst Rehabil Eng.* 2006; 14:194–198. [PubMed: 16792292]
10. Wolpaw JR, Birbaumer N, McFarland DJ, Pfurtscheller G, Vaughan TM. Brain-computer interfaces for communication and control. *Clin Neurophysiol.* 2002; 113:767–791. [PubMed: 12048038]
11. Mellinger J, Schalk G, Braun C, Preissl H, Rosenstiel W, Birbaumer N, Kubler A. An MEG-based brain-computer interface (BCI). *Neuroimage.* 2007; 36:581–593. [PubMed: 17475511]
12. Weiskopf N, Veit R, Erb M, Mathiak K, Grodd W, Goebel R, Birbaumer N, et al. Physiological self-regulation of regional brain activity using real-time functional magnetic resonance imaging (fMRI): methodology and exemplary data. *Neuroimage.* 2003; 19:577–586. [PubMed: 12880789]
13. Sitaram R, Zhang H, Guan C, Thulasidas M, Hoshi Y, Ishikawa A, Shimizu K, Birbaumer N. Temporal classification of multichannel near-infrared spectroscopy signals of motor imagery for developing a brain-computer interface. *NeuroImage.* 2007; 34:1416–1427. [PubMed: 17196832]
14. Taylor DM, Tillery SI, Schwartz AB. Direct cortical control of 3D neuroprosthetic devices. *Science.* 2002; 296:1829–1832. [PubMed: 12052948]
15. Bohannon RW, Smith MB. Interrater reliability of a modified Ashworth scale of muscle spasticity. *Phys Ther.* 1987; 67:206–207. [PubMed: 3809245]
16. Folstein MF, Folstein SE, McHugh PR. “Mini-mental state”: a practical method for grading the cognitive state of patients for the clinician. *J Psychiatr Res.* 1975; 12:189–198. [PubMed: 1202204]
17. Tecchio F, Zappasodi F, Tombini M, Caulo M, Vernieri F, Rossini PM. Interhemispheric asymmetry of primary hand representation and recovery after stroke: a MEG study. *Neuroimage.* 2007; 36:1057–1064. [PubMed: 17543542]
18. Huang JC, Nicholson C, Okada YC. Distortion of magnetic evoked fields and surface potentials by conductivity differences at boundaries in brain tissue. *Biophys J.* 1990; 57:1155–1166. [PubMed: 2393701]
19. Schalk G, McFarland DJ, Hinterberger T, Birbaumer N, Wolpaw JR. BCI2000: a general-purpose brain-computer interface (BCI) system. *IEEE Trans Biomed Eng.* 2004; 51:1034–1043. [PubMed: 15188875]
20. Pfurtscheller G, Lopes da Silva FH. Event-related EEG/MEG synchronization and desynchronization: basic principles. *Clin Neurophysiol.* 1999; 110:1842–1857. [PubMed: 10576479]
21. McFarland DJ, Miner LA, Vaughan TM, Wolpaw JR. Mu and beta rhythm topographies during motor imagery and actual movements. *Brain Topogr.* 2000; 12:177–186. [PubMed: 10791681]
22. Kubler A, Nijboer F, Mellinger J, Vaughan TM, Pawelzik H, Schalk G, McFarland DJ, Birbaumer N, Wolpaw JR. Patients with ALS can use sensorimotor rhythms to operate a brain-computer interface. *Neurology.* 2005; 64:1775–1777. [PubMed: 15911809]
23. Pfurtscheller G, Neuper C, Brunner C, da Silva FL. Beta rebound after different types of motor imagery in man. *Neurosci Lett.* 2005; 378:156–159. [PubMed: 15781150]
24. Wolpaw JR, McFarland DJ. Multichannel EEG-based brain-computer communication. *Electroencephalogr Clin Neurophysiol.* 1994; 90:444–449. [PubMed: 7515787]
25. Siegel, S., Castellan, NJ. *Nonparametric Statistics for the Behavioral Sciences.* New York: McGraw-Hill; 1988.
26. Al Snih S, Markides KS, Ottenbacher KJ, Raji MA. Hand grip strength and incident ADL disability in elderly Mexican Americans over a seven-year period. *Aging Clin Exp Res.* 2004; 16:481–486. [PubMed: 15739601]

27. Caithness G, Osu R, Bays P, Chase H, Klassen J, Kawato M, Wolpert DM, Flanagan JR. Failure to consolidate the consolidation theory of learning for sensorimotor adaptation tasks. *J Neurosci*. 2004; 24:8662–8671. [PubMed: 15470131]
28. Wolpaw JR. Brain-computer interfaces as new brain output pathways. *J Physiol*. 2007; 579:613–619. [PubMed: 17255164]
29. Birbaumer N, Ghanayim N, Hinterberger T, Iversen I, Kotchoubey B, Kubler A, Perelmouter J, Taub E, Flor H. A spelling device for the paralyzed. *Nature*. 1999; 398:297–298. [PubMed: 10192330]
30. Pfurtscheller G, Guger C, Muller G, Krausz G, Neuper C. Brain oscillations control hand orthosis in a tetraplegic. *Neurosci Lett*. 2000; 292:211–214. [PubMed: 11018314]
31. Sharma N, Pomeroy VM, Baron JC. Motor imagery: a backdoor to the motor system after stroke? *Stroke*. 2006; 37:1941–1952. [PubMed: 16741183]

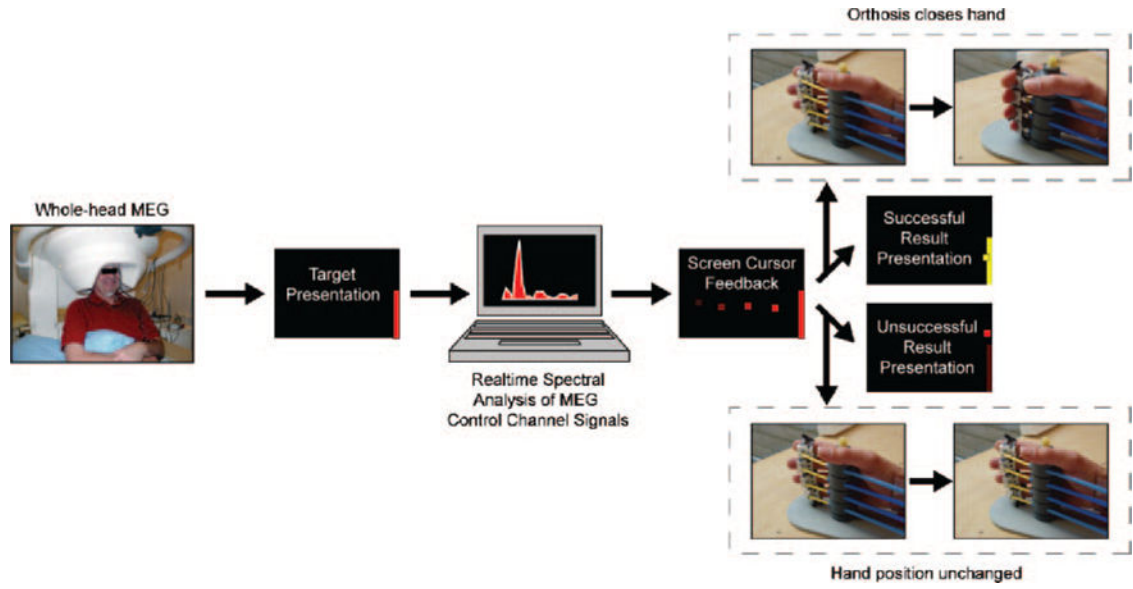


Figure 1. Trial description for BCI training. Whole-head MEG data (153 or 275-channels) was continuously recorded throughout each training block. At the initiation of each trial, 1 of 2 targets (top-right or bottom-right edge of screen) appeared on a projection screen positioned in front of the subject. Subsequently, a screen cursor would appear at the left edge of the screen, and begin moving toward the right edge at a fixed rate. A computer performed spectral analysis on epochs of data collected from a preselected subset of the sensor array (3 to 4 control sensors). The change in power estimated within a specific spectral band was transformed into the vertical position of the screen cursor feedback projected onto the screen. At the conclusion of the trial, if the subject was successful in deflecting the cursor upwards (net increase in spectral power over the trial period) or downwards (net decrease in spectral power over the trial period) to contact the target, 2 simultaneous reinforcement events occurred. The cursor and target on the visual feedback display changed colors from red to yellow. At the same time, the orthosis initiated a change in hand posture (opening or closing of hand). If the cursor did not successfully contact the target, no orthosis action was initiated.

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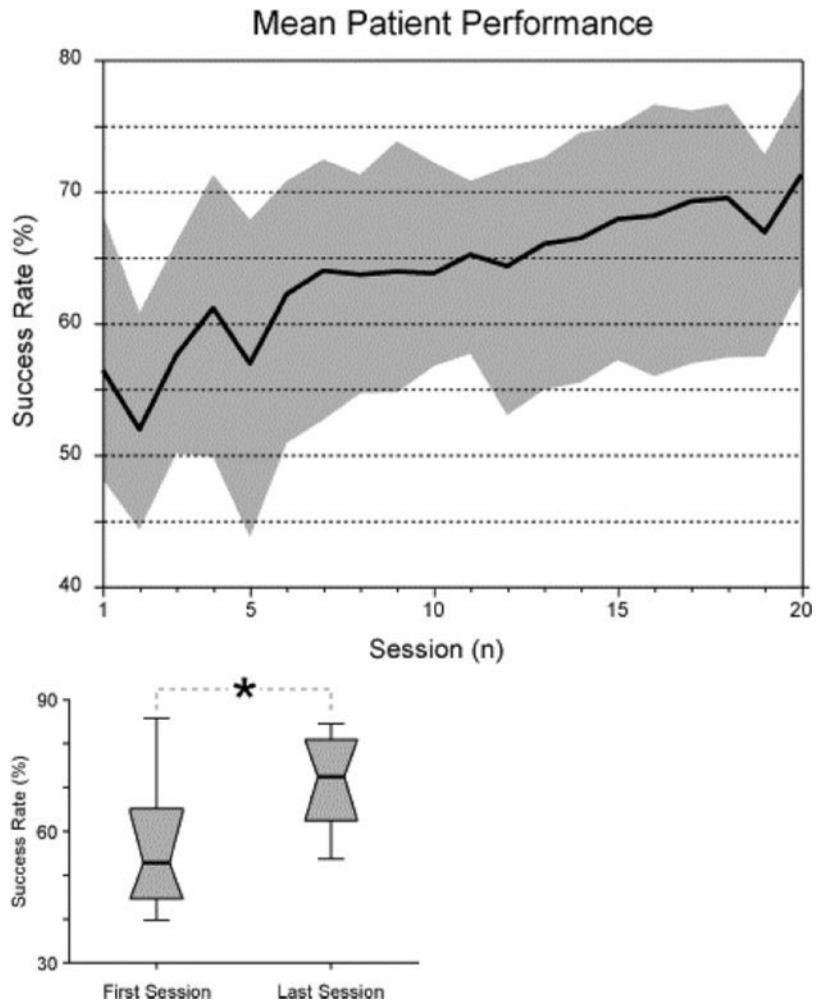


Figure 2. Average group success rate as a function of training session. The average success rate for the last training session is $72.48 \pm 18.36\%$ (median \pm interquartile range). As the total number of training sessions completed by patients was unique, the time-series for each individual was resampled and normalized to 20 sessions (the mode of the session duration across the patient group) using linear interpolation, before being averaged. The gray shaded area represents the 95% CI of the median estimate, which was computed using a bootstrap technique repeated 10 000 times. The boxplot (pre and post-training median and interquartile range) inset shows a significant group increase in success rate between the first and last training sessions.

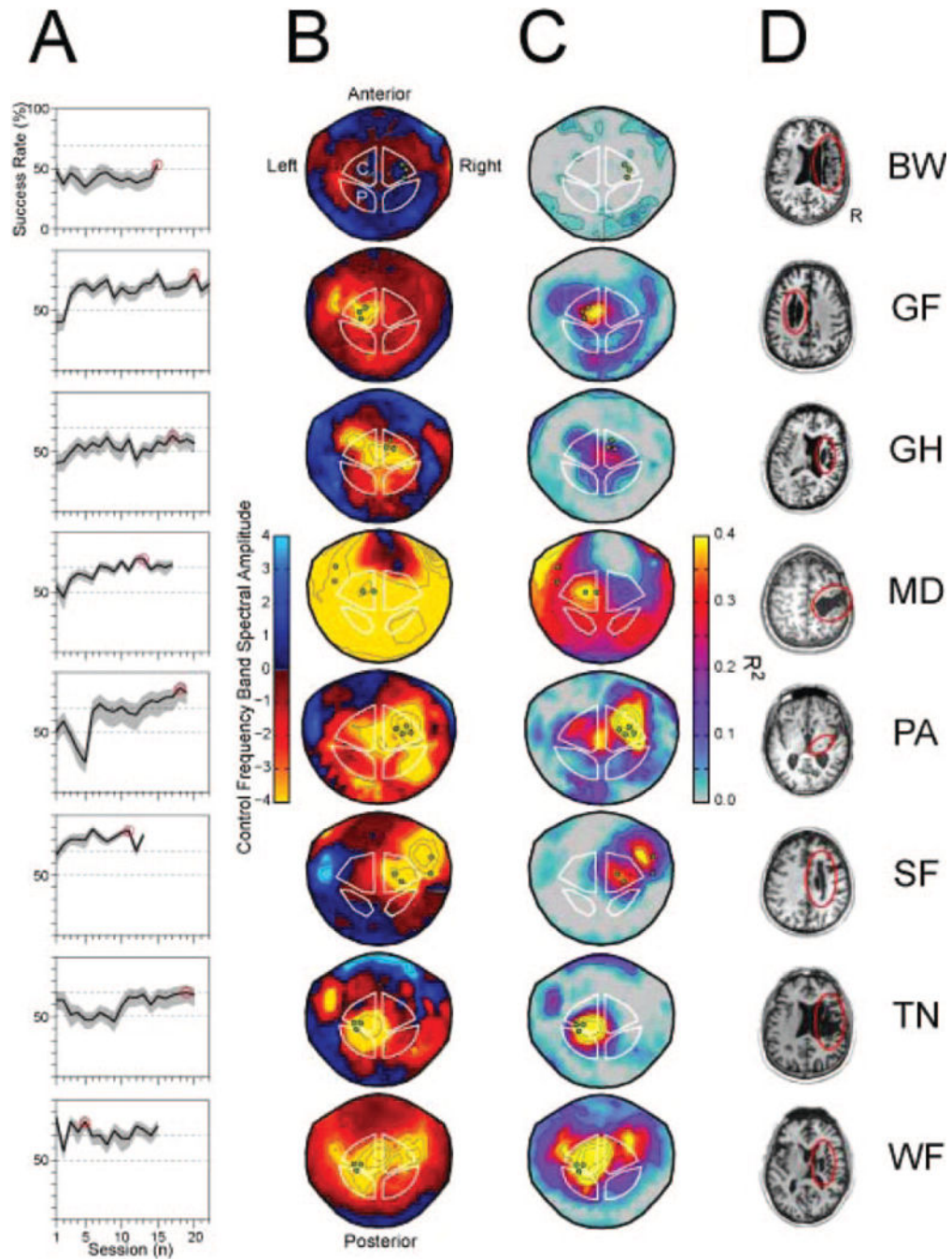


Figure 3. Individual subject performance, task-related brain activity, and lesion representations. Each row displays individual data for study participants. (From Left to Right): Column A shows the session performance for each patient. The gray shaded represents the 95% CI of the mean, which was computed using a bootstrap technique repeated 10 000 times. Columns B and C display task-related MEG brain activity from the sessions indicated by the red circle in Column A. With the exception of WF, whose performance peaked within the first 5 sessions of training, these represent the session with the highest performance that occurred

within the final 4 sessions of training. Column B displays a flat map of the spectral amplitude difference across the MEG array between both target conditions. The sensor locations used to generate feedback and control the orthosis action are highlighted by the green-filled circles. The locations of central and parietal sensors within the right and left hemispheres of the arrays are outlined in white (labeled in the top row of Figure 3B as “C” and “P”, respectively). Column C displays a statistical map (R^2) of the correlation of μ rhythm amplitude across the MEG array with target location/orthosis action. Column D displays single axial images from T1-weighted, high resolution MRI scans obtained for each subject (neurological convention). The red circles highlight the location of each patient’s lesion. All but patient GF had right hemisphere lesions.

Patient and Training Characteristics

Table

Patient	Age, y	Sex	Lesion Type	Duration of Hand Plegia, mo	Total Training Sessions, n	Training Frequency, Mode, Sessions/wk	Training Sensor Location, Mode	BCI Control Frequency, Mode, Hz
BW	63	F	Right subcortical (BG, centrum semiovale, wallerian degeneration of PT)	30	15	5	Ipsilesional central	12
GF	53	M	Left subcortical and cortical (corona radiata, put, insula, wallerian degeneration of PT)	12	22	3	Ipsilesional central	12
GH	55	M	Right subcortical (BG)	41	20	4	Ipsilesional central	12
MD	14	M	Right subcortical and cortical (put and adjacent white matter, pre- and postcentral gyri and adjacent regions, partially retained PT in the IC)	12	17	5	Contralesional central/temporal	12.5
PA	67	M	Right subcortical (thalamus and BG)	14	19	5	Ipsilesional central	9
SF	44	F	Right subcortical (put and adjacent white matter)	24	13	5	Ipsilesional central/temporal	12.5
TN	68	M	Right subcortical and cortical (encephalomalacia involving posterior frontal lobe, insula, and BG)	37	20	5	Contralesional central/parietal	12
WF	52	M	Right subcortical (BG and adjacent white matter)	16	15	3	Contralesional central/parietal	12

BG indicates basal ganglia; PT, pyramidal tract; Put, putamen; IC, internal capsule.