Title: Reliability, sensitivity and predictive value of fMRI during multiple object tracking as a marker of cognitive training in stroke survivors.

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Abstract

Computerized cognitive training (CCT) combined with transcranial direct current stimulation (tDCS) has showed some promise in alleviating cognitive impairments in patients with brain disorders, but the robustness and possible mechanisms are unclear. In this prospective double-blind randomized clinical trial, we investigated the feasibility and effectiveness of combining CCT and tDCS, and tested the predictive value of and training-related changes in fMRI-based brain activation during attentive performance (multiple object tracking) obtained before, during and after the three-weeks intervention in chronic stroke patients (> 6 months since hospital admission). Patients were randomized to one of two groups, receiving CCT and either (1) tDCS targeting left dorsolateral prefrontal cortex (1 mA), or (2) sham tDCS, with 40s active stimulation (1 mA) before fade out of the current. 77 patients were enrolled in the study, 54 completed the cognitive training, and 48 completed all training and MRI sessions. We found significant improvement in performance across all trained tasks, but no additional gain of tDCS. fMRI-based brain activation showed high reliability, and higher cognitive performance was associated with increased tracking-related activation in the dorsal attention network (DAN) and default mode network (DMN) as well as anterior cingulate after compared to before the intervention. We found no significant associations between cognitive gain and brain activation measured before training or in the difference in activation after intervention. Combined, these results show significant training effects on trained cognitive tasks in stroke survivors, with no clear evidence of additional gain of concurrent tDCS.

53 **Introduction**

54 While stroke mortality has decreased during the last decades in western countries, it remains a
55 leading cause of disabilities (Feigin et al., 2017), and many stroke survivors experience
56 persistent cognitive sequelae. Indeed, prevalence of neurocognitive disorders in post stroke
57 survivors have been reported to as high as 54% (Barbay et al., 2018), and may increase risk of
58 dementia (Pendlebury, 2012). Reduced working memory capacity is among the most common
59 impairments (Teasell, 1992; Zinn et al., 2007), with high prevalence at the chronic stage
60 (Mahon et al., 2017; Schaapsmeerders et al., 2013), and potentially large impact on daily
61 functioning (Synhaeve et al., 2014). Stroke lesions influence brain activation as measured
62 using brain imaging (Altamura et al., 2009), and working memory deficits have been linked to
63 aberrant brain activation during task engagement (Ziemus et al., 2007).

64 Computerized cognitive training (CCT) has shown some promise in augmenting
65 cognitive rehabilitation. A recent meta-analysis revealed small but significant improvements
66 on cognitive functioning in response to CCT in healthy participants (Au et al., 2015). Similar
67 findings have been reported in patients with Parkinson's disease (Leung et al., 2015), mild
68 cognitive impairment (Maffei et al., 2017), brain injury (Weicker et al., 2016), and stroke (De
69 Luca et al., 2018b; Peers et al., 2018), as well as schizophrenia (Prikken et al., 2018) and
70 major depression (Motter et al., 2016). However, the underlying mechanisms remain elusive.

71 Furthermore, non-invasive electrical stimulation of the brain has been suggested to
72 augment training benefits (Kang et al., 2015; Marquez et al., 2015). Indeed, evidence suggests
73 amplified motor recovery post-stroke, when physical therapy is paired with transcranial direct
74 current stimulation (tDCS) (Kang et al., 2015), and the beneficial effects have been suggested
75 to generalize to a cognitive rehabilitation context (Andrews et al., 2011; Fregni et al., 2005;
76 Lawrence et al., 2018). Despite some positive findings, the overall effectiveness of tDCS has
77 been questioned (Medina and Cason, 2017). Also, the potential mechanisms, if any, are

unclear, yet enhancing neuronal excitability by lowering the membrane-potential has been suggested as one of the candidate means (Giordano et al., 2017).

Functional MRI has become an increasingly used non-invasive brain imaging method linking behavior to brain activation patterns. Studies investigating alterations in brain activation in response to CCT have reported mixed results. In healthy controls, training gains have been associated with decreased activity in medial and dorsal frontal areas, including anterior cingulate (Heinzel et al., 2016; Miró-Padilla et al., 2018; Motes et al., 2018), with increased efficiency reflected in lowered activity as a suggested cognitive mechanism (Constantinidis and Klingberg, 2016). For patient cohorts the current evidence is sparser, but increased activity in frontal and parietal areas in response to cognitive training has been reported in patients with schizophrenia (Haut et al., 2010; Ramsay and MacDonald, 2015; Subramaniam et al., 2014). In stroke patients, CCT has been associated with increased functional connectivity (FC) between hippocampus, frontal and parietal areas, which was associated with improved cognitive performance (Lin et al., 2014; Yang et al., 2014).

Despite some promising findings, the effects of CCT (Buitenweg et al., 2018; De Luca et al., 2018a; Melby-Lervåg and Hulme, 2013; Melby-Lervåg et al., 2016) and tDCS (Dedoncker et al., 2016; Hill et al., 2016) are still unclear. Importantly, more knowledge about the origins of individual differences in training gains is key to evaluate and improve selection to fully realize the clinical potential of these interventions. Investigations of the neural underpinnings of cognitive improvement in response to CCT and tDCS may help clarify the benefits and limitations for future clinical application. It is therefore key to better understand mechanisms besides behavioral transfer-effects before evaluating the full effect of these programs.

To this end, we investigated differences in training-related gains of a commonly used working memory computerized training program in combination with tDCS, and tested the predictive value and sensitivity of fMRI-based brain activation as a marker for training-

104 related gain in task performance. Specifically, we estimated activation patterns during
105 multiple object tracking (MOT), which broadly recruits brain networks involved in sustained
106 attention and working memory (Alnæs et al., 2014; Dørum et al., 2016; Pylyshyn and Storm,
107 1988) at baseline, before initiating training (on average 4 weeks after baseline measure), and
108 after a three-week intervention. To assess specific associations with tDCS and following a
109 double-blind design, in addition to CCT each participant was pseudo-randomized into two
110 groups: 50% (n=27) receiving active tDCS directed towards the left dorsolateral prefrontal
111 cortex and 50% (n=27) receiving sham stimulation.

112 We tested the following hypotheses: "(1) Variability in cognitive training gain is
113 reflected in differential slopes of increased performance across participants, (2) level of
114 improvement is affected by tDCS stimulation protocol, (3) fMRI-based brain activation
115 patterns prior to training are predictive of training gain, and (4) individual differences in task
116 improvement are reflected in differential patterns of brain activation after the three-weeks
117 intervention, in particular involving frontal and parietal brain regions.

118

119 **Methods**

120 *Sample and exclusion criteria*

121 Figure 1 depicts the study recruitment pipeline. Participants were recruited from two major
122 hospitals located in Oslo, Norway (Stroke unit at Oslo University hospital, and Geriatric
123 department at Diakonhjemmet Hospital, Oslo, Norway). Hospital staff identified suitable
124 participants. Inclusion criterion was identifiable stroke with ischemic or hemorrhagic
125 etiology. Patients with transient ischemic attacks (TIA) were not included. Exclusion criteria
126 included MRI contraindications, other neurological diseases diagnosed prior to the stroke,
127 severe psychiatric diagnosis including bipolar disorder, schizophrenia or drug abuse.
128 Approximately 900 invitations were sent out, and 250 responded; among those, 77 patients
129 were eligible for participation and were initially included in the study (Ulrichsen et al., 2019).

130 19 participants withdrew from the study prior to (n=14) or during (n=5) the cognitive training
131 due to the labor-intensity of the intervention. Four participants were excluded during the
132 course of the intervention due to newly occurring medical issues violating the inclusion
133 criteria. Two participants were included without radiological findings when admitted to the
134 stroke unit, but displayed neurological symptoms. Initial MRI diffusion weighted imaging
135 revealed possible acute infarctions and the patients were treated with thrombolysis.
136 Subsequent MRI T2-weighted imaging revealed no certain radiological findings, but
137 symptoms persisted upon discharge and they were diagnosed by a neurologist with ICD-10
138 code I63.9 Cerebral infarction, and therefore included in the final sample. In total, 54
139 participants completed the cognitive training program. For timepoint 1, one participant had
140 corrupted behavioral data and two participants were not able to complete the task during the
141 fMRI assessment, and were excluded from all MRI analysis. For MRI timepoint 2 and 3, one
142 participant had poor alignment to the MNI-template, and two participants were not able to
143 complete the task, yielding a total of 51 participants baseline and 48 with full MRI at all
144 timepoints. In this final sample, 26 participants received active stimulation, and 22 received
145 sham stimulation.
146 The study was approved by the Regional Committee for Medical and Health Research
147 Ethics South-East Norway (2014/694 and 2015/1282), and all participants provided written
148 informed consent.
149

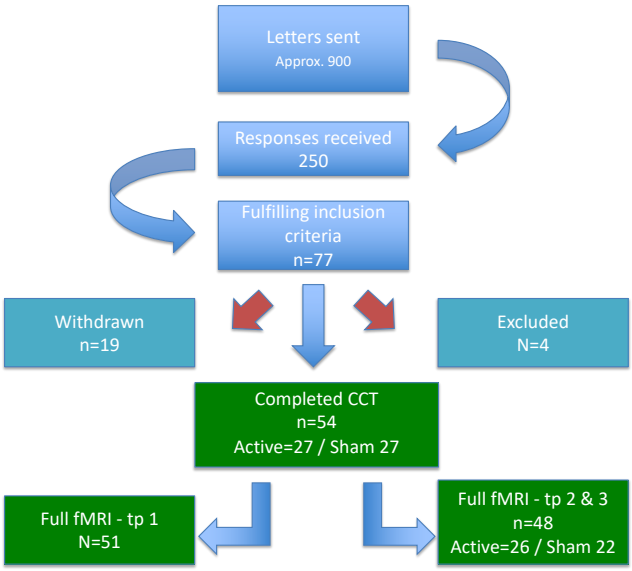

Figure 1. Flowchart displaying inclusion pipeline

Patient characteristics and lesion demarcation

Sample characteristics for the participants completing the intervention (n=54) are described in Table 1, including National Institutes of Health Stroke Scale (NIHSS) (Goldstein et al., 1989), Trial of Org 10172 in Acute Stroke Treatment (TOAST) (Adams et al., 1993), Mini-Mental State Exam (MMSE) (Strobel and Engedal, 2008) and IQ derived from Wechsler Abbreviated Scale of Intelligence (WASI) (Ørbeck and Sundet, 2007). Briefly, the sample mean age was 69 years (sd=7.3), comprised 74% males, and average IQ of 110 (sd=16).

For each participant, lesion demarcation was performed by a trained assistant, utilizing the semi-automated toolbox clusterize, implemented for SPM8 (Clas et al., 2012; de Haan et al., 2015), and guided by radiological descriptions. Figure 2 displays a probabilistic lesion map across participants. Supplementary Figure 1 provides an overview of individual lesions.

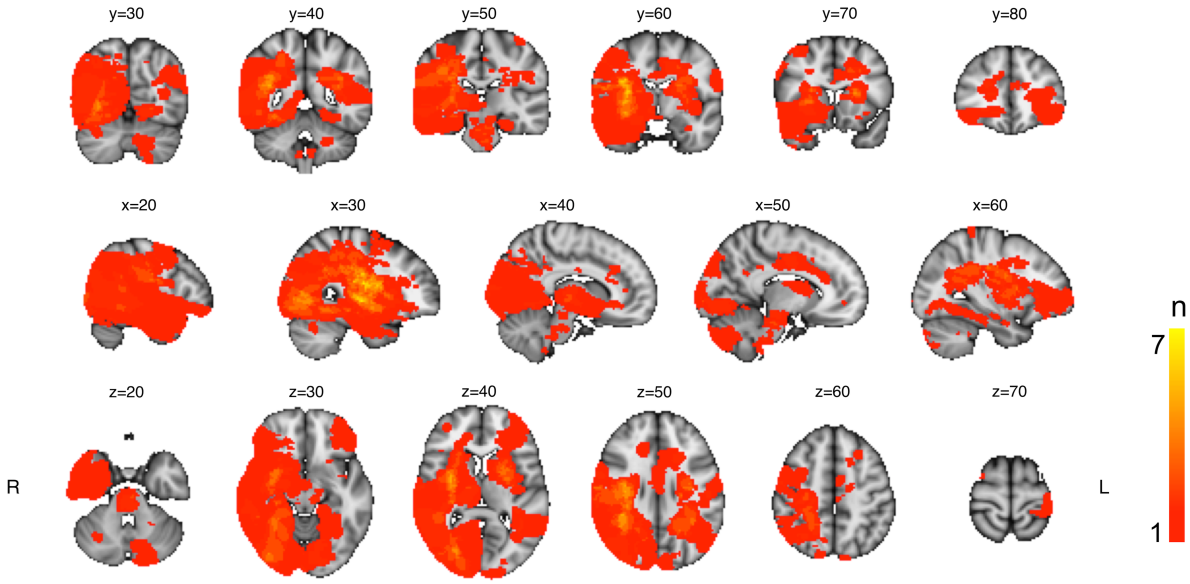

Figure 2. Heatmap displaying lesion overlap across included participants. The color scale reflect the number of overlapping lesions in each voxel (ranging from 0 to 7).

	Mean	sd	Min	Max
Participant descriptives (current)				
Age at inclusion	69.13	7.374	47	81
Males (%)	74.07	-		
Time between stroke and inclusion (months)	25.74	9.17	6	45
Self-reported education in years	14.39	3.71	9	30
MMSE at inclusion	27.98	1.855	22	30
IQ	110.45	16.906	66	136
Days between inclusion and training	33.78	12.986	19	74
Days between MRI assessment pre/post training	31.72	6.257	19	49
Clinical variables during hospitalization				
NIHSS at discharge	1.33	1.53	0	7
TOAST classification	Large vessel disease n=20 Small vessel disease n=18 Cardioembolic disease n=6 Other n=10			

Table 1. Sample characteristics

Study protocol

The protocol have previously been described in detail (Richard et al., 2020). Briefly, after inclusion the participants completed MRI assessment at three timepoints: timepoint 1, at

timepoint 2 on average 3-4 weeks after timepoint 1 without additional contact, and at timepoint 3, after the participant completed the training protocol.

Cognitive training

The patients performed a computerized working memory training program (Cogmed QM, Cogmed Systems AB, Stockholm, Sweden), comprising 12 auditory-verbal and visual-spatial working memory tasks. The patients performed a computerized working memory training program (Cogmed QM, Cogmed Systems AB, Stockholm, Sweden), comprising 12 auditory-verbal and visual-spatial working memory tasks. The implementation of the protocol has previously been reported in detail (Richard et al., 2020). Briefly, the training comprised five weekly training sessions over 3-4 weeks, with 17 training sessions for each participant in total. Each participant completed two weekly tDCS stimulation sessions, with a total of 6 tDCS sessions with a minimum of 48 hours between each stimulation. Level of Cogmed task difficulty was automatically adjusted according to each participant's performance level. The initial two first sessions for each task was discarded from statistical analysis to ensure each participant was at a desired level of difficulty. The same task setup was used for all participants, and tasks completed three times or less over the course of training were excluded. Eight Cogmed subtests were included in further analysis, including Grid, Cube, Digits, Hidden objects, Sort, Rotation, Twist, and 3D-cube.

tDCS

Participants were randomly assigned to either active or sham condition, using codes provided by the manufacturer and implemented by an in-house Matlab script, pseudo-randomizing participants while maintaining balance in groups of 20. tDCS stimulation was applied at six occasions for each participant, with a minimum of 48 hours between stimulations, aiming at an average of two stimulations per week. Stimulation current was 1000 μA, stimulation time

199 was 20 minutes for the active group with a ramp-up time of 120 seconds and fade-out time of
200 30s. The current density was limited to 1mA to limit the risk of adverse events due to the
201 electrical stimulation. The sham stimulation consisted of ramp-up followed by 40 seconds of
202 active stimulation and then fade-out, following factory settings. Stimulation was delivered via
203 a battery-driven direct current stimulator (neuroConn DC stimulator plus, Germany), through
204 5 x 7 cm rubber pads, yielding a 28.57 μA/cm2 current density at skin surface. Impedance
205 threshold at start-up of stimulation was < 20 kΩ, and was regulated by the device with an
206 absolute threshold at 40 kΩ for automatically terminating stimulation. The pads were covered
207 with high-conductive gel (Abralyt HiCl, Falk Minow Services Herrsching, Germany) to
208 enhance conductance.
209 Electrode location was determined by following the 10-20 system, where the anodal
210 electrode was located at F3 corresponding to left DLPFC and cathodal at O2 corresponding to
211 right occipital/cerebellum, and fixated with rubber bands.
212
213 *fMRI task*
214 Participants performed a blocked version of the MOT (Pylyshyn and Storm, 1988), as
215 previously described in detail (Alnæs et al., 2015b). Briefly, the participants were presented
216 with ten identical circles on a grey background. All objects were initially blue, before zero
217 (passive viewing), one (load 1) or two (load 2) of the objects turned red for 2.5 seconds,
218 designating them as targets. After targets had returned to blue, the objects started moving
219 randomly around the screen for 12 seconds. After the 12 second tracking period, the objects
220 became stationary, one of the objects turned green, and the participants were instructed to
221 respond “yes” or “no” to whether or not the green probe had been designated as a target in
222 that trial. Participants were instructed to keep fixation on a fixation point at the center of the
223 screen during both attentive tracking and passive viewing. The task consisted of 24 semi-
224 randomized trials in total, ensuring the same condition was not repeated consecutively.

225 *MRI acquisition*

226 MRI data was acquired with a 3T GE 750 Discovery MRI scanner (32-channel head coil)
227 located at Oslo University Hospital. T1-weighted images was acquired using a 3D IR-
228 prepared FSPGR (BRAVO) with the following parameters: scan time: 4:43 minutes and
229 included 188 sagittal slices, repetition time (TR): 8.16 ms, echo time: 3.18 ms, flip angle: 12º,
230 voxel size: 1 × 1 × 1 mm, field of view: 256 x 256 mm,

231 *Functional MRI sequences* was acquired with a BOLD-sensitive gradient echo planar
232 sequence, with TR of 2.25 s, TE 30 ms, field of view 256 x 256 mm, slice thickness 3 mm
233 with 0.5 mm gap acquired ascending interleaved, 90º flip angle, with R/L orientation. The
234 initial first five volumes were removed before analysis.

235

236 *fMRI data processing*

237 fMRI data was processed using FMRI Expert Analysis Tool (FEAT) Version 6.00, from
238 FMRIB's Software Library (FSL; Jenkinson et al., 2012; Smith et al., 2004), and included the
239 following steps: correction for motion using MCFLIRT (Jenkinson et al., 2002), linear trend
240 removal and high-pass filtering (0.01 Hz), removal of non-brain tissue using BET (Smith,
241 2002), spatial smoothing with a Gaussian kernel of full width at half maximum (FWHM) of 6
242 mm (SUSAN ; Smith and Brady, 1997), and linear registration using FLIRT (Jenkinson and
243 Smith, 2001) to Montreal Neurological Institute (MNI) 152 standard space using the T1-
244 weighted scan as an intermediate. In the same manner, normalization parameters were
245 estimated for each participants T2-weighted image, and applied to the demarked lesion.

246

247 *Statistical analysis of behavioral data*

248 *Cogmed main effects of time:* To investigate the trajectory of Cogmed performance across
249 participants and time points, linear mixed effects (lme) models were estimated for each test,
250 using the lme function from the nlme-package (Pinheiro et al., 2017) in R (Team, 2016). Task

251 performance was entered as dependent variable, with training session as independent variable,
252 and age, sex, tDCS-group, educational level in years (self-reported), and interaction between
253 training session and tDCS, as fixed factors, and participant as random factor. To assess
254 normality in the data according to criteria for lme, we plotted distribution of random effects
255 intercepts, see supplementary Figure 7. Visual inspection did not indicate severe violations of
256 the criteria.

257 *Cogmed individual trajectories:* To quantify individual trajectories in Cogmed
258 performance across the training-period, we estimated for each test and each participant a
259 linear model with performance as dependent variable and session number as independent
260 variable, yielding a beta-estimate (slope) for each test for each participant reflecting changes
261 in performance over time. As a measure of overall performance for each participant, we used
262 the average task performance across sessions for each test for each participant. To investigate
263 task homogeneity in beta-estimates and average performance we performed multivariate
264 outlier detection with the aq.plot function in the mvoutliers package in R (P. Filzmoser and
265 Gschwandtner, 2018), with an alpha of 0.001 and 90% of the sample to estimate the minimum
266 covariance determinant. Two subtests ("hidden objects" and "Digits") were found to have a
267 high number of outliers compared to the other subtests, and were excluded from further
268 analysis (See Supplementary Figure 2 for details).

269 To derive a common score for average Cogmed performance and gain, we performed
270 a principal component analysis (PCA) on beta-estimates and average performance,
271 respectively. All scores were zero-centered and standardized prior to running the PCA. We
272 used the first factor from the beta PCA as a *Cogmed change score*, and the first factor from
273 the average performance PCA as a *Cogmed average performance score (*see Supplementary
274 Figure 3 for scree-plot derived from the PCA performed on both with and without the
275 excluded tasks*)*.

To explore potential association between initial cognitive capacity and Cogmed change score, we correlated performance on the cognitive performance at baseline assessed with the Cabpad battery (https://www.cognisoft.info) at TP1, described in detail previously (Richard et al., 2019) with the Cogmed change score. To test for associations between initial cognitive performance and response to tDCS we estimated linear models using the same baseline cognitive performance from Cabpad as the dependent variable, and Cogmed change score and tDCS group as independent variables. Models were estimated using the lm function in the stats package in R (Team, 2013).

To assess the association between Cogmed scores and lesion severity, we estimated linear models for both Cogmed change and average performance scores, using lesion volume, number of lesions and their interaction as explanatory variables. Models were estimated using the lm function in the stats package in R (Team, 2013).

fMRI behavioral analysis: To investigate performance during the MOT task across timepoints, linear mixed effects (lme) models were estimated for both load 1 and load 2, using the lme function from the nlme-package (Pinheiro et al., 2017) in R. Task performance was entered as dependent variable, with training session as independent variable, and participant as random factor. To assess for pairwise differences across the timepoints, post hoc analysis was performed using the ghlt function from the multcomp package (Hothorn et al., 2008) in R, and corrected for multiple comparisons using bonferroni correction.

fMRI data analysis

First-level GLMs for each participant and scanning session were estimated using FEAT (Woolrich et al., 2001), including regressors for passive viewing, load 1, load 2, responses, as well as the standard and extended motion parameters estimated during preprocessing. The following contrasts were estimated: passive viewing, load 1, load 2, tracking ([load 1 + load 2] – passive viewing) and load (load 2 – load 1).

302 To estimate main effects of experimental conditions across individuals, contrasts of
303 parameter estimates (COPEs) from each participant's first scan session (baseline) were
304 submitted to randomise (Winkler et al., 2014) performing a one sample t-test for each
305 condition separately. Multiple comparisons were corrected for using permutation testing and
306 threshold free cluster enhancement (TFCE (Smith and Nichols, 2009)).
307 To estimate associations with task performance with brain activation during task
308 engagement, individual COPE maps for load 1 and load 2 were stacked separately, and
309 submitted to randomise, including sex, age and task performance as covariates.
310 To test for differences in brain activation between scan sessions, we first performed
311 fixed effects analysis for each participant separately, including all COPEs from first level
312 GLM, specifying the contrasts session 3 minus session 2, session 3 minus session 1, and
313 session 2 minus session 1. Next, all individual COPE maps were submitted to randomise to
314 test for group-level main effects while correcting for multiple comparisons using permutation
315 testing and threshold free cluster enhancement (TFCE).
316 To quantify within-subject reliability in brain activation across time we estimated
317 intraclass correlation coefficient (ICC) for each voxel between assessment two and three
318 utilizing the variance from one-way random ANOVA, for the contrasts passive viewing, load
319 1 and load 2, as implemented in the ICCest function from the ICC R package (Wolak et al.,
320 2012).
321 To test for associations between Cogmed change and average performance scores and
322 brain activation, individual level COPE maps for timepoint 1 and the difference between
323 timepoint 3 and 2 including the conditions passive viewing, load1, load 2, tracking as well as
324 load were stacked separately and submitted to randomise. The model included sex, age, and
325 either Cogmed change score or Cogmed average performance score (obtained from the PCA).
326 In the same manner to explore differences associated with tDCS stimulation protocol, the
327 same COPE´s from the difference between timepoint 3 and 2 including all conditions were

submitted to randomise, with the model including sex, age and tDCS-condition. All models were corrected for multiple comparisons across space using 5000 permutations and thresholded using TFCE.

Results

Cogmed performance

Figure 3 shows Cogmed performance over time for each group and Table 2 displays corresponding summary statistics. Lme revealed a significant group-level improvement on all Cogmed tests included in the analysis. In addition, we found a significant association with age on mean performance across all tests, indicating lower average performance with higher age. We found no significant associations between Cogmed performance and sex, tDCS group (sham/experimental), or years of education. We found a significant interaction between time and tDCS on the twist task, suggesting augmented training gain in the participants in the active tDCS stimulation group. However, after removal of one outlier defined by visual inspection, the result did not reach significance (see Supplementary Table 1 and Supplementary Figure 4 for corresponding statistics and visualization).

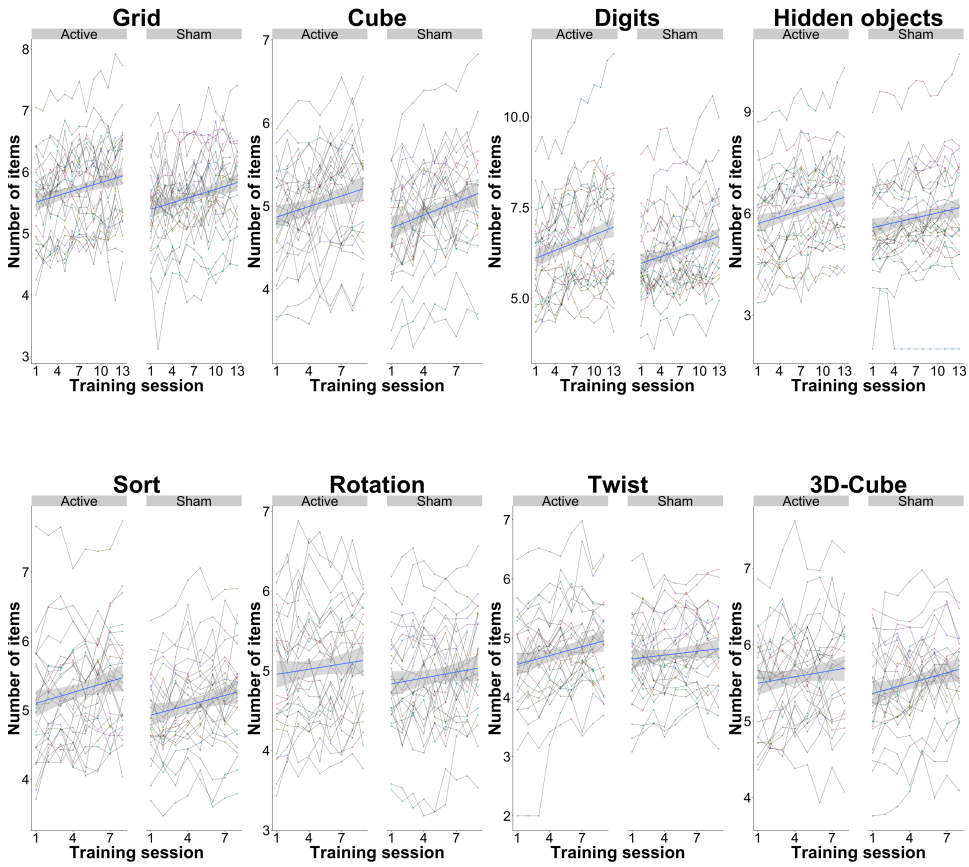

Figure 3. Individual level task performance during the course of the intervention period for each group (tDCS sham/active). The fit lines are based on a linear fit within each group.

	Grid	Sort	Digits	Cube	Hidden objects	Twist	3D-Cube	Rotation
Session	**8.66** ***(<0.001)***	**6.40** ***(<0.001)***	**12.79** ***(<0.001)***	**7.44** ***(<0.001)***	**11.03** ***(<0.001)***	**6.45** ***(<0.001)***	**3.16** ***(0.002)***	**2.21** ***(0.028)***
Age	**-3.05** ***(0.004)***	**-2.11** ***(0.041)***	**-6.07** ***(<0.001)***	**-2.54** ***(0.015)***	**-4.86** ***(<0.001)***	**-3.62** ***(0.001)***	**-3.90** ***(<0.001)***	**-3.41** ***(0.001)***
Sex	-0.89 (0.377)	-0.33 (0.741)	-2.29 (0.027)	-1.22 (0.23)	-2.25 (0.029)	-1.46 (0.15)	-1.42 (0.162)	-1.34 (0.187)
tDCS	-0.65 (0.521)	-1.03 (0.309)	-0.06 (0.951)	-0.53 (0.599)	0.12 (0.908)	1.00 (0.324)	-0.82 (0.414)	-0.81 (0.422)
Education	0.06 (0.951)	0.96 (0.34)	0.90 (0.374)	0.14 (0.886)	1.57 (0.124)	0.01 (0.991)	-0.15 (0.884)	0.33 (0.744)
Session x tDCS	0.2 (0.842)	-0.39 (0.7)	-1.53 (0.128)	0.46 (0.644)	-1.79 (0.074)	**-2.83** **(0.005)**	1.72 (0.087)	1.23 (0.22)

Table 2. Summary statistics (t/(p))from linear mixed effects models testing for associations between task performance and session, age, sex, tDCS, education and the interaction between session and tDCS. t-vaules along with raw p-vaules are displayed, significant findings are highlighted (FDR-corrected, alpha = 0.05).

We found no significant correlation between Cogmed average performance and Cogmed change score (r=.17, t=1.29, p=.2). We found a significant association between lesion volume and Cogmed change score (t -2.5, p=.017), indicating a negative correlation between lesion volume and training response, which did not retain significance after removal of one outlier (t=-.58, p=.563). See Supplementary Table 2 for corresponding statistics.

Linear models revealed no significant associations between cognitive capacity as measured using Cabpad at baseline and gain in Cogmed performance. Furthermore, there were no significant interactions between Cabpad score at baseline and tDCS on Cogmed change score. See supplementary figure 6 and supplementary table 4 for corresponding statistics.

Behavioral fMRI results

LME analysis revealed main effect of time on task performance during both load 1 and load 2. Post hoc analysis revealed significant differences between timepoint 3 and 1, as well as between timepoint 2 and 1, but not between timepoint 3 and 2. See Table 3 and Figure 4 for details.

	Main effect		**Tp2 - Tp1**		**Tp3 – Tp1**		**Tp3 – Tp 2**	
Model	**F**	**p**	**t**	**p**	**t**	**p**	**t**	**p**
Load 1	**9.22**	**2.20e-04**	**3.52**	**1.28e-03**	**3.89**	**3.03e-04**	0.36	1.00
Load 2	**8.24**	**5.04e-04**	2.31	0.06	**4.05**	**1.56e-04**	1.73	0.25

Table 3. Summary statistics from linear mixed effects models testing for associations between task performance and session for the MOT task. Bonferroni-corrected p-values are displayed, significant results are highlighted.

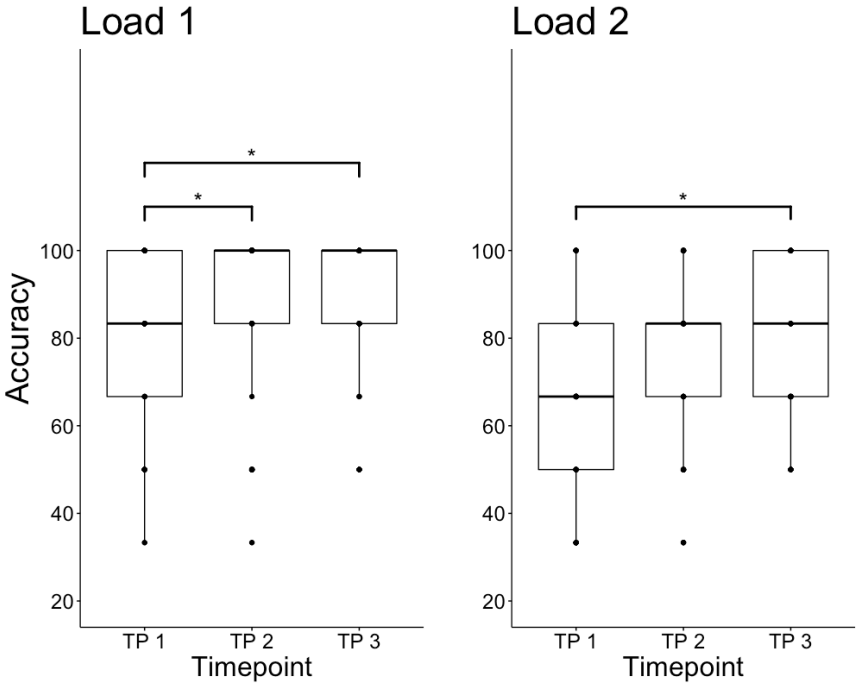

Figure 4. Task performance during the MOT task at the three different timepoints. Significant differences in group means is marked *, Table 3 displays corresponding statistics.

Association between task performance and fMRI activation.

Linear models revealed no significant association between in-scanner task performance and fMRI activation during neither load 1 or load 2 during the first assessment (TP 1). In-scanner behavioral performance was therefore not included in further analysis.

fMRI main effects of task

Figure 5 shows main effects of task condition at baseline, and Table 4 summarizes anatomical locations. Passive viewing was associated with increased activity in the dorsal attention network (DAN) and the inferior and superior occipital cortex, and decreased activation in pre and post-central gyrus, supplementary motor area (SMA), anterior cingulate, as well as default mode network (DMN), including posterior cingulate gyrus (PCC), medial prefrontal gyrus (mPFC), and superior and medial temporal gyrus (S/MTG).

Load 1 was associated with increased activity in DAN, extending into the inferior and superior occipital cortex, cerebellum, and precentral gyrus, in the cingulo-opercular (CO) network, including thalamus, frontal operculum and insula, as well as in the ventral attention (VAN) network including temporoparietal junction and supramarginal gyrus, middle frontal gyrus, and frontal operculum. Decreased activity was seen in DMN nodes including PCC, mPFC and S/MTG. Load 2 showed a similar pattern to Load 1, with additional increased activation in right anterior middle frontal gyrus. Tracking over passive viewing revealed same activation pattern as load 1 and load 2, with additional bilateral activation of anterior middle frontal gyrus. Load-related (load 2 –load1) increases in activation were seen in superior and inferior parietal lobe, as well as right anterior middle frontal gyrus, with no significant load-dependent decreases in activation.

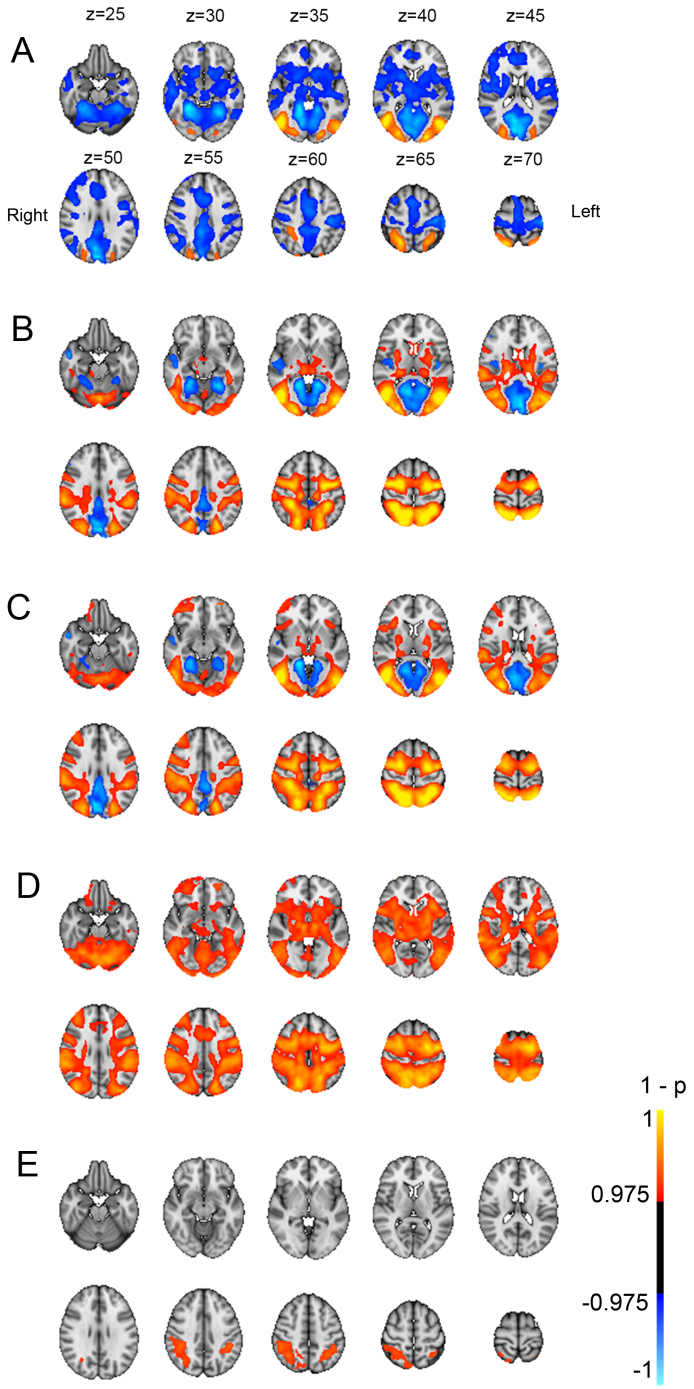

Figure 5. Main effects from task-fMRI for (A) passive viewing > baseline, (B) load 1 > baseline, (C) load 2 > baseline, (D) tracking > passive viewing, and (E) increased cognitive load (load 2 > load 1). Panel A display corresponding coordinates in MNI, and left / right orientation. P-values (1-p) have been multiplied with the sign of the effect in relation to the main contrast.

Condition	Direction	size	X	Y	Z	Area
PV	pos	4029	33	19	30	Lateral occipital cortex
	pos	3861	55	18	32	Lateral occipital cortex
	pos	1784	29	41	61	Superior parietal lobe
	neg	64337	33	35	21	Cerebellum
Load1	pos	57023	59	29	8	Cerebellum
	neg	12615	33	39	23	Cerebellum
	neg	1494	19	62	24	Middle temporal gyrus
	neg	256	65	53	42	Central opercular cortex
	neg	153	20	78	48	Middle frontal gyrus
	neg	19	22	71	50	Inferior frontal gyrus
Load2	pos	66242	59	29	8	Cerebellum
	pos	411	65	71	37	Frontal operculum
	pos	8	65	80	55	Middle frontal gyrus
	neg	10269	31	40	26	Occipital fusiform cortex
	neg	694	18	63	26	Superior temporal gyrus
Tracking	pos	101656	59	29	8	Cerebellum
Load	pos	4590	24	42	55	Inferior parietal lobe
	pos	1709	67	42	56	Inferior parietal lobe
	pos	245	24	79	54	Middle frontal gyrus

Table 4. *Clusters associated with different MOT condition at baseline (all p<.05, corrected)*

fMRI – reliability

Figure 6 shows voxel-wise ICC for COPE values for passive viewing, load 1 and load 2, between timepoint 2 and 3. The results revealed ICC>0.4 across major portions of the brain, including occipital, temporal, parietal lobe for all conditions, as well as and a large part of the frontal lobes for load 1.

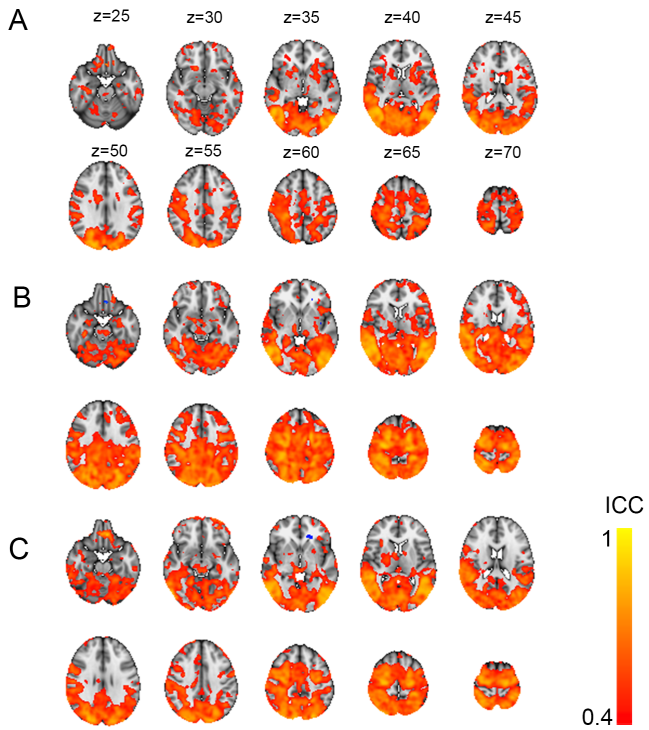

Figure 6. Voxel-wise ICC for condition passive viewing (A), load 1 (B), and load 2 (C). Thresholded at ICC > 0.4.

Pairwise comparison of fMRI estimates across timepoints

Comparison of fMRI signal between the timepoints revealed no significant differences between timepoint 3 and 1, timepoint 2 and 1 as well timepoint 3 and 2, for neither passive viewing, load 1, load 2, tracking nor load, indicating fluctuating patterns, but not in a time-dependent manner.

Cogmed –fMRI associations

We found no significant associations between the Cogmed change score and brain activation at timepoint 1, or between Cogmed change score and the difference in brain activation between timepoint 2 and 3.

Furthermore, results revealed no significant associations between the Cogmed average score and brain activation at timepoint 1. However, between timepoint 2 and 3 we found a significant association between Cogmed average performance score and the difference in activation during attentive tracking, indicating larger activation increases in the anterior and posterior parts of the cingulate cortex and left precuneus, superior parietal lobe, lateral occipital cortex, as well as cerebellum with greater average Cogmed performance. See Table 5 and Figure 7 for corresponding statistics. Using a Cogmed average performance score based on all 8 subtests (from the PCA, now including two sub-tests previously excluded), revealed slightly weaker but highly similar effects in the (Supplementary Figure 5 and Table 3), suggesting that excluding the two subtests did not bias the results.

	Direction	Size	x	y	z	Location
Frontal	pos	120	30.1	64.4	61.4	Middle frontal gyrus
	pos	46	59.5	63.9	60.5	MFG L
Limbic	pos	1269	41.7	74.2	53.3	ACC
	pos	261	43.8	52.3	53.8	PCC
Parietal	pos	83	23	36.8	61.9	SPL
	pos	207	55	30.2	48.7	Precuneus cortex
Occipital	pos	117	21.5	25.9	39.3	Lateral occipital cortex
	pos	100	61.6	22.7	44.4	Lateral occipital cortex
	pos	33	35.8	15.3	36.3	Occipital pole
Cerebellar	pos	54	60	26	13	Cerebellum
	pos	16	41.1	37.2	23.5	Cerebellum

Table 5. Cogmed average score clusters associated with difference in activation between timepoint 2 and 3. (all $p<.05$, corrected).

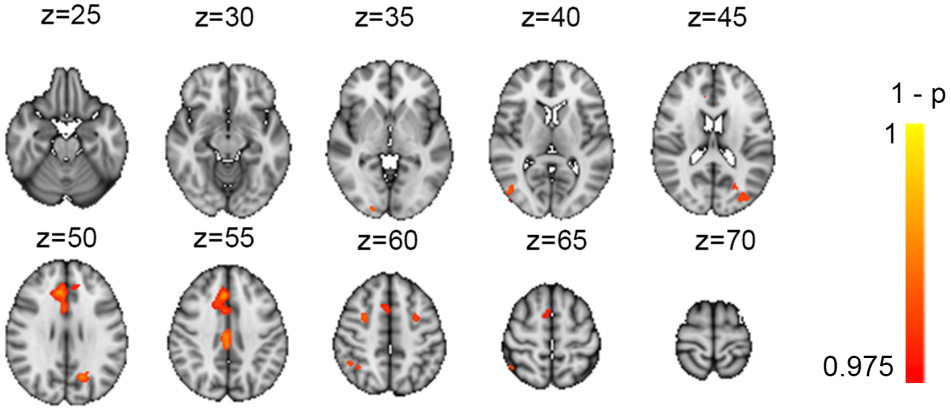

Figure 7. (A) Association between Cogmed average performance score, and difference in task-related activation (L1 & L2 > PV) between timepoint 3 and 2.

tDCS –fMRI associations

Results revealed no difference in fMRI-activation between the active and sham tDCS stimulation groups.

Discussion

In this study, we tested the feasibility and effectiveness of a commercially available working memory training package paired with tDCS in a heterogeneous group of chronic stroke survivors. Further, we assessed the predictive value of fMRI-based brain activation pre-training on training related gains, as well as changes in task activations pre and post training.

The results revealed significant increase in performance on the trained tasks, with no additional benefits from tDCS on training gain. Our analysis revealed robust activation in response to the MOT task in line with previous findings. It is important to note that the current study investigated possible markers for individual training outcome, and a comprehensive evaluation of the potential behavioral and clinical transfer effects is beyond

482 the scope of the current study. Importantly, investigation of reliability across timepoints
483 supported the feasibility of utilizing task-fMRI for delineating neural correlates of behavioral
484 training in a clinical group. Following the promising reliability in brain activation across the
485 different fMRI assessments, we found no significant association between cognitive
486 performance gain during the cognitive training and brain activation prior to training, or with
487 brain activation change between pre- and post training. However, higher average performance
488 across the training period was associated with higher load-dependent increase in activation in
489 the precuneus, and furthermore associated with increased activation in the DAN and DMN as
490 well as the ACC during active task performance after compared to before intervention.
491 Tailoring behavioral interventions for alleviating cognitive difficulties following
492 stroke calls for investigation of both treatment response, as well as delineating factors
493 predicting favorable outcome. Cognitive abilities prior to training as well as motivation (Guye
494 et al., 2017; Jaeggi et al., 2014) are currently among our best predictors for beneficial effects.
495 Our results revealed significant improvement on all included tasks from the training program,
496 in line with previous reports (Spencer-Smith and Klingberg, 2015; Westerberg et al., 2007).
497 Our results support that repeated practice, where level of difficulty is adjusted during training,
498 improves performance on the trained tasks. It is important to note that the current study
499 investigates possible biomarkers for individual training outcome, and its beyond our aim to
500 evaluate potential behavioral transfer effects. However, the generalizability of training-effects
501 following CCT has been debated (Au et al., 2015; Melby-Lervåg and Hulme, 2013; Morrison
502 and Chein, 2011; Shipstead et al., 2012; von Bastian and Oberauer, 2014). Indeed, recent
503 large-scale reviews suggest a lack of reliable transfer beyond tasks that share properties with
504 the trained tasks (Melby-Lervåg et al., 2016; Sala and Gobet, 2018). The lack of transfer
505 effects is suggested to be caused by the failure of targeting common mechanisms underlying
506 fluid intelligence such as working memory and attention, and rather targeting task specific
507 mechanisms through repeated practice (Melby-Lervåg et al., 2016). Despite our promising

behavioral results on trained tasks, the lack of transfer-effects reported in the literature calls for caution before implementation in a clinical setting.

Our results revealed no significant interaction between cognitive improvement and tDCS stimulation with the exception of one of the trained tasks, which did not reach significance after removal of an outlier. Although anodal tDCS has shown beneficial effects on cognitive training compared with sham stimulation in healthy adults (Martin et al., 2013) and have been associated with steeper learning curves (Ruf et al., 2017), a recent meta-analysis provided no support of a beneficial effect of tDCS on cognitive training compared to sham (Medina and Cason, 2017). The current literature investigating the added effect of tDCS when combined with working memory training in stroke patients is scarce. Single session anodal tDCS towards the left DLPFC improved recognition accuracy during an N-back task (Jo et al., 2009), and longitudinal concurrent anodal tDCS and CCT increased accuracy on a non-trained continuous performance test (CPT) compared to sham (Park et al., 2013). However, divergence in study protocols and results hampers a direct comparison (daSilva Filho et al., 2017). Beyond robust improvement on all trained tasks during the course of the intervention, our results provide no evidence of additional beneficial effects of tDCS stimulation.

Our fMRI results for task activation during the first assessment revealed increased activation of the DAN, along with deactivation of DMN for passive viewing, load 1 and load 2. Further, contrasting tracking to passive viewing, and load 2 to load 1 revealed that attentive tracking increased DAN activation in a load-dependent manner, in line with previous reports based on healthy samples (Alnæs et al., 2014; Dørum et al., 2016).

Importantly, longitudinal studies utilizing fMRI for investigation of within-subject changes depends on the reliability of brain measurements across time in order to pick up the relevant signal. Previous investigations suggest favorable reliability in task based compared to resting-state fMRI (Specht, 2020), indicating higher reliability when the task elicits strong

534 patterns of activation (Johnstone et al., 2005) increasing contrast to noise ratio (Bennett and
535 Miller, 2010). Our results are in line with these reports, as results revealed fair (>0.4) to good
536 (>0.75) reliability across task-associated brain regions following guidelines for ICC
537 interpretation (Cicchetti, 1994), with poor reliability in non-task associated areas.
538
539 Our results revealed no significant interaction between activation during passive
540 viewing, load 1, load 2, tracking nor load, and time. Although we did not find any predictive
541 value of fMRI-activation patterns on Cogmed performance improvement, our results revealed
542 an association between average performance during the Cogmed training period and increased
543 tracking-related activity in key nodes of the DAN and DMN as well as ACC. DMN is
544 commonly suppressed during task engagement, with increasing suppression with increased
545 task demands (Anticevic et al., 2012). Furthermore, cognitive abilities are negatively
546 correlated with degree of de-activation (Lipp et al., 2012), most likely due to need for higher
547 effort to resolve the task. In line with these findings our results indicate less suppression of
548 PCC and precuneus during task engagement, with greater average performance.
549 Working memory capacity, reflecting both executive control as well as cognitive
550 flexibility, has been linked to general cognitive abilities (Conway et al., 2003). Furthermore,
551 higher cognitive abilities have been associated with increased activity in frontal and parietal
552 cortices (Basten et al., 2015), and cognitive flexibility have been linked to brain activation in
553 ventrolateral prefrontal cortex, as well as anterior and posterior cingulate (Dajani and Uddin,
554 2015). As our results indicate that greater cognitive capacity is associated with an increase in
555 activation during task engagement over time, it may reflect compensation and learning to
556 more efficiently resolve the task.
557 The current findings should be interpreted in light of several limitations. Based on the
558 clinical ratings (NIHSS) at hospital discharge, the patients were sampled from the less severe
559 part of the stroke severity spectrum. Further, patient drop-out during the intervention was

560 relatively high, partly due to the labor-intensity of the training. Hence, our sample is biased
561 towards higher functioning stroke patients, rendering the generalizability of our results to
562 more severe cases unclear. The lack of a control group for the CCT does not allow us to
563 disentangle effects of the cognitive training from the effect of time, anticipation, or other
564 confounders. As our main aim was to identify potential fMRI-based markers for training
565 outcome, we did not assess possible cognitive and clinical transfer effects, which will be
566 relevant for evaluating clinical relevance, and further studies are needed to establish the
567 premises for optimizing the clinical potential for working memory interventions in individual
568 stroke patients. It has furthermore been reported that potential beneficial effects of tDCS may
569 emerge at a delayed stage after intervention (Goodwill et al., 2016; Li et al., 2019). As our last
570 assessment was performed shortly after the final stimulation, any delayed effects may have
571 been lost. Future follow-up assessments of the same patients may provide relevant
572 information about potential long-term effects of CCT and tDCS. Potential effects also may be
573 dependent of stimulation frequency (Boggio et al., 2007; Yun et al., 2015). Increasing
574 frequency beyond current protocol may have altered our results. Lesion size and location
575 may influence current flow and cortical excitability (Marquez et al., 2013), and future studies
576 may test for associations between lesion characteristics and response to tDCS. While the
577 degree of cognitive impairment at baseline is likely to influence the response to cognitive
578 training and rehabilitation in general, we found no significant association between baseline
579 cognitive performance and training gain., see response above. However, the lack of pre-
580 stroke neuropsychological evaluation complicates the assessment of stroke-related cognitive
581 impairment. The group level task activation patterns in the current study largely mirrored
582 those reported in previous studies in healthy controls (Alnæs et al., 2015a; Dørum et al.,
583 2016). However, the current study design did not allow us to make an explicit comparison
584 (D'Esposito et al., 2003). Here we only considered brain activation in response to task
585 demands. Future studies should also pursue a promising line of research utilizing imaging

indices of structural and functional connectivity to investigate behavioral correlates of stroke, which may supplement and increase our understanding of stroke and stroke recovery.

In conclusion, we have investigated response to a computerized cognitive training program in 54 stroke survivors in combination with tDCS, as well as the predictive value of neural activation on training outcome, and neural alterations following training. Our results revealed increased performance across all trained tasks, with no additional benefit of tDCS. Brain activation prior to the training was not predictive for training outcome, nor was training gains reflected in altered brain activation. The generalizability of the reported beneficial effects of CCT remains uncertain, and future studies may be able to assess the transfer value of for relevant cognitive and clinical variables in stroke patients.

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