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SCHONFELD, Lisa; Jahanshahi, Ali; LEMMENS, Evi; Bauwens, Matthias; Heschem, Sarah-Anna; Schipper, Sandra; Lagiere, Melanie; HENDRIX, Sven & Temel, Yasin (2017) Motor cortex stimulation does not lead to functional recovery after experimental cortical injury in rats. In: RESTORATIVE NEUROLOGY AND NEUROSCIENCE, 35(3), p. 295-305.

DOI: 10.3233/RNN-160703

Handle: <http://hdl.handle.net/1942/24310>

Motor cortex stimulation does not lead to functional recovery after experimental cortical injury in rats

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Abstract.

Background: Motor impairments are among the major complications that develop after cortical damage caused by either stroke or traumatic brain injury. Motor cortex stimulation (MCS) can improve motor functions in animal models of stroke by inducing neuroplasticity.

Objective: In the current study, the therapeutic effect of chronic MCS was assessed in a rat model of severe cortical damage.

Methods: A controlled cortical impact (CCI) was applied to the forelimb area of the motor cortex followed by implantation of a flat electrode covering the lesion area. Forelimb function was assessed using the Montoya staircase test and the cylinder test before and after a period of chronic MCS. Furthermore, the effect of MCS on tissue metabolism and lesion size was measured using [¹⁸F]-fluorodesoxyglucose (FDG) μ PET scanning.

Results: CCI caused a considerable lesion at the level of the motor cortex and dorsal striatum together with a long-lasting behavioral phenotype of forelimb impairment. However, MCS applied to the CCI lesion did not lead to any improvement in limb functioning when compared to non-stimulated control rats. Also, MCS neither changed lesion size nor distribution of FDG.

Conclusion: The use of MCS as a standalone treatment did not improve motor impairments in a rat model of severe cortical damage given our specific treatment modalities.

Keywords: Motor cortex stimulation, motor impairment, rehabilitation, behavioral tests, PET

1. Introduction

Cortical damage due to traumatic brain injury (TBI) or stroke often leads to persistent functional impairments if the motor cortex is part of the traumatized or infarcted brain region. The resulting motor impairments are disabling and form a major socio-economic burden (Parker, Wade, & Langton Hewer, 1986; Walker & Pickett, 2007). Thus far, the only

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clinically proven therapy for patients with motor deficits is physical rehabilitation therapy; still, many patients do not achieve full recovery. To enhance the efficacy of physical rehabilitation therapy, motor cortex stimulation (MCS) has been proposed as a potential therapeutic approach (Brown, Lutsep, Weinand, & Cramer, 2006; R. Levy et al., 2008). Recently, the results of a multicenter study were reported, in which stroke patients suffering from hemiplegia received six weeks of MCS via implanted epidural electrodes concurrent with physical rehabilitation therapy (R. M. Levy et al., 2015). The authors reported a promising recovery course of the patients that was still present six months after cessation of the therapy (R. M. Levy et al., 2015). MCS has also been applied in rodent and non-human primate models of ischemic infarcts and resulted in improved limb function, again when being combined with physical rehabilitation therapy (Adkins et al., 2006; Baba et al., 2009; Plautz et al., 2003). However, limited data are available investigating the effect of MCS on its own without an additional intervention.

One of the mechanisms explaining the therapeutic effect of MCS on motor recovery is neuroplasticity. In a previous study, we found that MCS applied to naïve rats increased cell proliferation in the subventricular zone (SVZ) compared to non-stimulated controls (Jahanshahi, Schönfeld et al., 2013a). Furthermore, a higher number of neural stem and progenitor cells (NSPCs) and mature neurons were detected in the motor cortex underneath the electrode (Jahanshahi, Schönfeld et al., 2013a). This finding could be explained by a process called electrotaxis, where cells migration is induced by an electrical field (Jahanshahi, Schönfeld, Lemmens, Hendrix, & Temel, 2013). In other studies using different forms of electrical stimulation, this increase of NSPCs at the side of stimulation was further corroborated, strengthening the hypothesis of electrotaxis *in vivo* (Morimoto et al., 2011; Rueger et al., 2012).

Animal models can be used to mimic clinical symptoms in a standardized way. With a controlled cortical impact (CCI), a cortical lesion can be created in rats that results in long-lasting functional deficits (L. M. Schönfeld et al., 2016). Similar to humans, rats possess a topographic organization of the motor cortex, where distinct cortical areas control the function of specific body parts (Nishibe, Barbay, Guggenmos, & Nudo, 2010; Starkey et al., 2012) and a CCI lesion in the forelimb area of the motor cortex can cause deficits in motor functions specific to the contralateral forelimb (L.M. Schönfeld et al., 2016).

In the present study, we tested whether MCS as a standalone treatment is able to achieve functional recovery in a rat model of severe CCI in the forelimb area of the motor cortex. To document functional recovery, we measured the effect of MCS on forelimb function and metabolic brain activity.

2. Materials & methods

2.1. Subjects

All animal experiments were conducted according to the directive 2010/63/EU on the protection of animals used for scientific purposes and had been approved by the local ethical committee for animal experiments at Maastricht University. Forty male Sprague-Dawley rats (Charles River, France), ten weeks old and weighing approximately 400 g at the time of surgery, were housed in pairs under a reversed 12 h light/dark cycle. Housing and testing facilities were kept at a constant temperature of 22° C and a humidity of 40–60%. Animals received standard laboratory chow (Sniff, Germany) and acidified water (pH 2.3–2.7) *ad libitum*, if not specified otherwise. Each behavioral assessment took place during the dark phase of the reversed night-day cycle (between 7 am and 7 pm), which is the active period of the rats.

2.2. CCI induction and electrode implantation

Induction of CCI was performed as previously described in detail (L.M. Schönfeld et al., 2016). Shortly, a craniotomy was made above the forelimb area of the motor cortex (coordinates AP 0–3.5 mm anterior to bregma, ML 0.5–4 mm lateral to bregma) contralateral to the dominant paw, as determined by baseline performance in the Montoya staircase test. All rats received a CCI using an electromagnetically driven impactor device (Leica Impact One, Leica Biosystems, USA) with an impactor tip of 3 mm diameter, an impact depth of 5 mm and a velocity of 3 m/s (Fig. 1c). Polyurethane-isolated flat electrodes (3.4 × 3 mm; Medtronic, USA) with six exposed monopolar platinum/iridium contact points, were positioned on top of the CCI lesion (Fig. 1a-b) and a reference wire was anchored to the contralateral skull (coordinates AP 1.75 mm anterior to bregma and ML 2.25 mm lateral to bregma). Both the electrode and the reference wire were fixed with dental cement (Paladur; Heraeus, Germany) exposing two

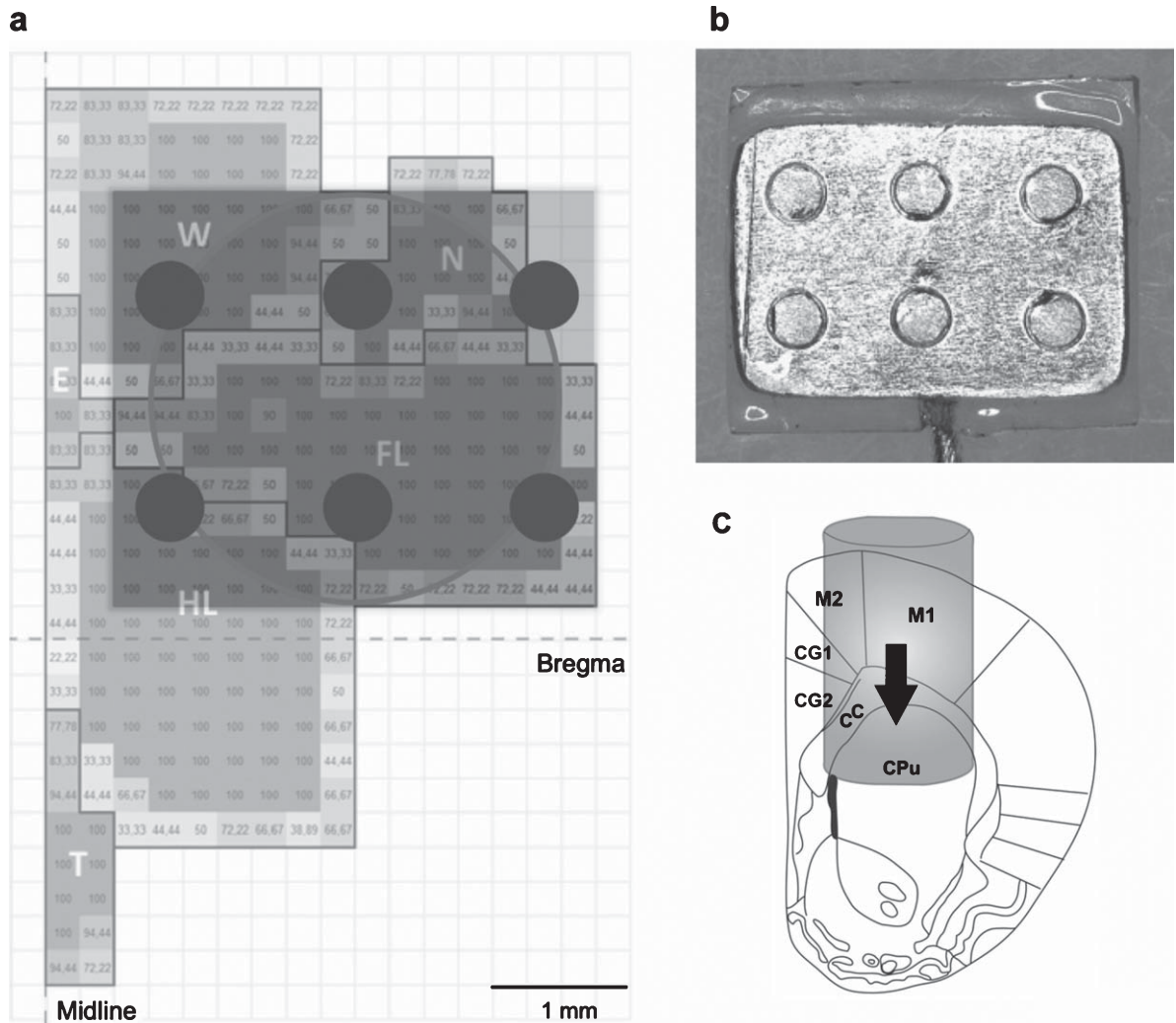


Fig. 1. Location of the CCI and electrode placement. A schematic representation of the somatotopic organization of the rat motor cortex is shown, modified from Fonoff et al. (2016; a). The CCI targeted large parts of the forelimb area (FL; CCI area represented by a red circle), followed by electrode placement on top of the lesion (blue square). Dark circles represent the monopolar electrode contacts that delivered the current. On the right, a scaled-down picture of the electrode lead is shown (real size 3.4×3 mm; b). CCI on the motor cortex damaged large parts of the cortex and the dorsal striatum (c; image shown at 1.7 mm anterior to bregma). CCI: controlled cortical impact; W: whiskers; N: neck muscles; FL: forelimb; HL: hindlimb; T: tail; E: eyes; M1: primary motor cortex; M2: secondary motor cortex; CG1: cingulate cortex 1; CG2: cingulate cortex 2; CC: corpus callosum; CPu: caudate putamen (striatum).

134 electrode contact pins (Multi-Contact, Switzerland)
 135 on the back of the animal's head to allow connection
 136 to an external electrical stimulator. Animals from the
 137 control group received non-functional dummy elec-
 138 trodes of the same size and material. After surgery,
 139 rats were left to recover for two weeks.

140 2.3. Motor cortex stimulation

141 MCS was applied daily for 2 h during a period
 142 of 31 consecutive days. Stimulation parameters were

chosen based on previous studies (Baba et al., 2009;
 Jahanshahi, Schönfeld et al., 2013b; Teskey, Flynn,
 Goertzen, Monfils, & Young, 2003) and consisted of
 a frequency of 30 Hz, 1 ms pulse width and biphasic
 constant current set at 50% of the current that
 evoked a motor threshold. MCS was delivered by an
 external digital stimulator (DS8000, World Precision
 Instruments, Germany) while rats stayed individually
 in stimulation chambers (width 28 cm, depth 50 cm,
 height 47.8 cm) and were allowed to move around
 freely under conditions similar to their home cage.

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154 Control animals underwent the same procedure with-
155 out any current being delivered.

156 2.4. Behavioral testing

157 Reaching and grasping abilities of both forelimbs
158 were assessed with the Montoya staircase test. In
159 short, rats have to retrieve sucrose pellets (Test Diet,
160 USA) lying on each step of a staircase located on the
161 left and on the right hand side of a platform inside a
162 narrow translucent box (Montoya, Campbell-Hope,
163 Pemberton, & Dunnett, 1991). All steps, except the
164 highest one, were baited with three sucrose pellets.
165 Rats were habituated to the staircase boxes and then
166 trained daily until they retrieved a minimum of 55%
167 of the pellets from at least one of the staircases
168 (Windle et al., 2006). Testing sessions lasted 15 min-
169 utes and took place on two consecutive days, twice
170 daily with a minimal inter-trial interval of three hours
171 resulting in a total of four testing sessions per time
172 point of behavioral assessment. During the training
173 and testing period, rats were food deprived to 85–90%
174 of their free-feeding weight to increase their motiva-
175 tion for pellet reaching. Data are presented as the total
176 number of eaten pellets for the impaired and healthy
177 forelimb separately, as well as using a *difference score*
178 defined as the score of the healthy forelimb subtracted
179 from the score of the impaired forelimb ($\text{pellets}_{\text{imp}} -$
180 $\text{pellets}_{\text{heal}}$). Behavioral testing in the Montoya stair-
181 case test was performed at three time points: Before
182 CCI, two weeks after CCI and after four weeks of
183 (sham) MCS.

184 Paw use during vertical exploration was measured
185 by the cylinder test as described previously (L.M.
186 Schönfeld et al., 2016). Rats were transferred to
187 Perspex cylinders on an illuminated platform and
188 recorded from above during 10 minutes (GoPro Hero
189 4, GoPro, USA), while they explored the cylinder
190 by rearing and leaning against the wall. Based on
191 the video footage, the first twenty wall contacts were
192 scored and used for analysis. Wall contacts were made
193 using both paws individually ('impaired', 'healthy')
194 or using both paws at the same time ('both'). Data
195 are presented as the percentage of the wall contacts
196 with either the impaired or the healthy forelimb rela-
197 tive to the total twenty wall contacts ($\text{contacts}_{\text{imp}}/20$
198 $* 100$ and $\text{contacts}_{\text{heal}}/20 * 100$, respectively). In
199 addition, the *difference score* of the percentages is
200 shown to visualize asymmetry between both fore-
201 limbs ($\% \text{contacts}_{\text{imp}} - \% \text{contacts}_{\text{heal}}$). Behavioral
202 testing using the cylinder test was performed at four
203 time points: Before CCI, two weeks after CCI, two

weeks after initiation of MCS or sham stimulation
and after four weeks of MCS or sham stimulation.

206 2.5. Functional imaging

207 Distribution of ^{18}F -fluorodesoxyglucose as an
208 indirect indicator of glucose-related metabolic activ-
209 ity in the central nervous system was visualized
210 *in vivo* using a μPET scanner (μPET Focus,
211 Siemens, the Netherlands). Rats were anesthetized
212 with Isoflurane and received 10–20 mBq ^{18}F -
213 fluorodesoxyglucose (FDG; GE Healthcare, the
214 Netherlands) intravenously, immediately followed by
215 scanning the entire brain for 30 minutes. Thereafter,
216 a static image was reconstructed using OSEM2D.
217 Each animal underwent μPET twice; on the first day
218 rats from the MCS group were scanned with stimula-
219 tion off, whereas on the second day stimulation was
220 switched on 10 minutes before as well as throughout
221 the entire duration of scanning in order to visualize
222 potential acute effects of MCS by using autoradio-
223 graphy. Control animals were scanned twice under
224 the same conditions, thus without any stimulation
225 being delivered. The lesion volume was calculated
226 by delineating the virtual CCI area, identified as the
227 cortical area without any visual presence of FDG
228 (Fig. 3a). The CCI lesion was delineated through-
229 out its full length in static images of sequential brain
230 slices using pmod image analysis software (PMOD
231 2.9, pmod Technologies, Switzerland).

232 Autoradiography was performed after the second
233 μPET scan to visualize FDG distribution at a high
234 spatial resolution. After transcardial perfusion with
235 4% paraformaldehyde, brains were frozen and cut
236 into 50 μm thick sections. Autoradiography phos-
237 phor plates (GE Healthcare, the Netherlands) were
238 exposed to the frozen brain sections during approx-
239 imately 2 hours and read for each animal with a
240 Typhoon FLA7000IP scanner (GE Healthcare, the
241 Netherlands). For each animal, the entire lesioned
242 and healthy hemispheres of three sections at a com-
243 parable bregma level were delineated separately and
244 the intensity of the FDG signal for the individ-
245 ual hemispheres was measured using ImageQuant
246 TL software (GE Healthcare, the Netherlands). Sig-
247 nal intensity (arbitrary units, a.u.) depended on the
248 amount of radioactive counts present in the delin-
249 eated area and was corrected for the injected amount
250 of MBq, animal weight, the time from injection
251 until exposure to the autoradiography plate and the
252 duration of exposure on the autoradiography plate.
253 To correct for inter-individual fluctuations, intensity

values were expressed by dividing the intensity values measured within the lesioned hemisphere by the intensity values measured within the healthy hemisphere for each individual animal. In addition, intensity values for each hemisphere are shown separately.

2.6. Statistical analysis and artwork

Data are presented as mean \pm standard error of the mean and were analyzed with repeated-measures ANOVA (SPSS 20, IBM, US) with time (baseline, post CCI and post stimulation) as within-subjects factor and group (MCS and control) as between-subjects factor. Imaging data were analyzed with one-way ANOVA and p -values below 0.05 were considered significant. Values below or above 1.5 interquartile ranges were identified as outliers by SPSS and excluded. In addition, animals that lost their electrodes during the course of the experiment and therefore could not undergo stimulation during 31 days were excluded from the analysis.

Figures were created using GraphPad Prism 5 (GraphPad Software, US), Adobe Illustrator CS6 (Adobe, US) and Microsoft PowerPoint 2011 (Microsoft, US).

3. Results

3.1. Chronic motor cortex stimulation failed to recover grasping skills and paw asymmetry during vertical exploration behavior

The Montoya staircase test was used to assess the recovery of reaching and grasping skills that were impaired by severe CCI. Repeated measures ANOVA revealed that the number of eaten pellets with the healthy paw increased after CCI compared to the number of eaten pellets before CCI for both groups [$F(2, 44) = 14.27, p < 0.001$; Fig. 2a] without any significant difference between rats that received MCS and rats that received sham stimulation. The number of eaten pellets with the impaired paw decreased after CCI [$F(2, 46) = 85.18, p < 0.001$; Fig. 2b] and a significant difference was detected between the groups at all time points [$F(1, 23) = 6.18, p < 0.05$]. This difference between the groups, however, was constant at all timepoints (2.41, 2.45 and 2.87 pellets, representing the mean number of pellets eaten before CCI, after CCI and after four weeks of MCS or sham stimulation), which indicates a lack of functional

improvement caused by MCS. The *difference score* was calculated by subtracting the number of pellets eaten with the healthy paw from the number of pellets eaten with the impaired paw. Using the *difference score*, a decline over time was shown implying worse pellet retrieval with the impaired paw in both groups [$F(2, 46) = 71.44, p < 0.001$; Fig. 2c]. The analysis of the *difference score* also revealed that MCS treatment did not affect the number of pellets eaten after CCI. These results indicate that MCS did not have any effect on grasping and reaching behavior after CCI.

The cylinder test was performed to measure vertical exploration behavior with the individual forelimbs. After CCI, all animals showed an increased reliance on the healthy paw to lean against the cylinder wall [$F(3, 69) = 19.30, p < 0.001$; Fig. 2d] at the expense of using their impaired paw [$F(3, 66) = 6.77, p < 0.001$; Fig. 2e]. However, MCS treatment did not restore usage of the impaired paw. Analysis of the *difference score* showed a stronger asymmetry in paw use after CCI compared to paw use before CCI [$F(1, 8, 66) = 19.73, p < 0.001$; Fig. 2f]. Yet, treatment with MCS could not resolve this asymmetry in wall contacts between both forelimbs. Taken together, these results indicate that treatment with MCS could not restore forelimb use for vertical exploration behavior.

3.2. Lesion volume and glucose metabolism in the lesioned hemisphere did not change despite motor cortex stimulation

Delineation of the lesion area in the reconstructed μ PET images was performed to estimate the amount of histological damage that was present after chronic application of MCS (Fig. 3a). The lesion volume of animals that received chronic MCS was not significantly different from the lesion volume of non-stimulated controls [$F(1,8) = 2.98, p > 0.05$; Fig. 3b], which indicated a lack in overt tissue recovery.

Imaging the distribution of FDG in brain slices was performed by means of autoradiography to measure functional recovery that may have occurred at the tissue level (Fig. 3c1-2). The ratio of intensity values between the healthy and the lesioned hemisphere did not differ between rats that received MCS and control rats that received sham stimulation [$F(1,7) = 0.12, p > 0.05$; Fig. 3d]. When comparing the intensity values of the delineated lesioned and healthy hemisphere separately, no difference was detected between the animal groups, either [lesioned hemisphere: $F(1,7) = 0.64, p > 0.05$; healthy hemi-

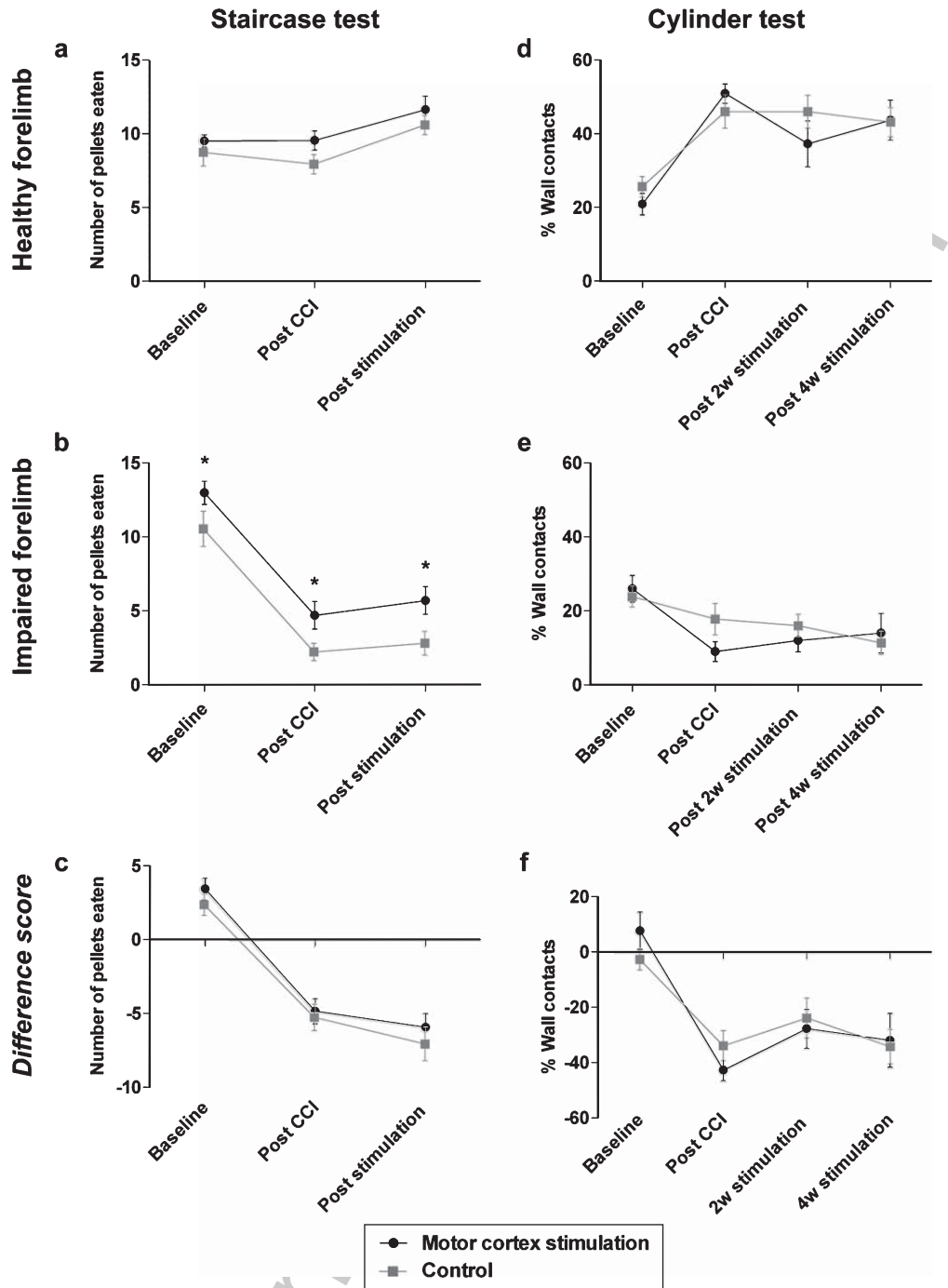


Fig. 2. Chronic motor cortex stimulation after a unilateral CCI neither recovered reaching and grasping skills nor paw use during vertical exploration. CCI to the forelimb area of the motor cortex did not affect pellet retrieval with the healthy paw (a) whereas it resulted in significantly less pellet retrieval with the impaired paw (b), as measured in the Montoya staircase test. This behavioral impairment was unaltered by MCS. Concerning the impaired paw, an overall significant difference between groups at all time points was detected, which was unrelated to the application of MCS. The *difference score* also reflected the tendency of less pellets eaten with the impaired paw (c). Use of the cylinder test showed that after a unilateral CCI rats increased the use of their ipsilateral paw to lean against the cylinder walls (d), while neglecting the paw contralateral to the CCI lesion (e). The *difference score* also show a decreased use of the impaired forelimb to lean against the cylinder wall (f). This effect of CCI on motor impairment was not restored by MCS. CCI=controlled cortical impact; MCS= motor cortex stimulation; * $p < 0.05$ comparing MCS with control.

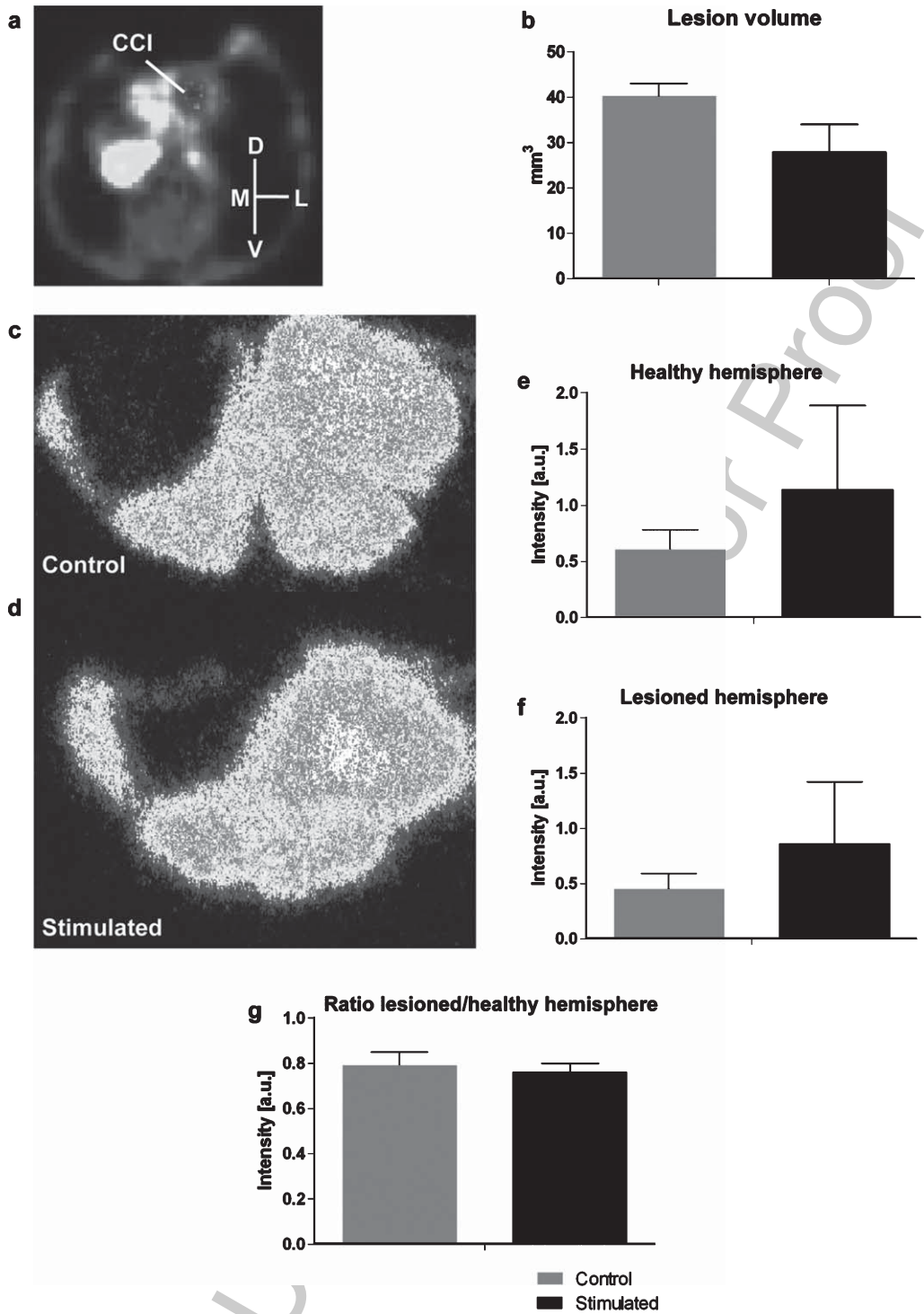


Fig. 3. Neither lesion volume nor glucose distribution was affected by chronic motor cortex stimulation. A representative μ PET image is shown in a horizontal plane, used to delineate the lesion area (white line, a). Chronic MCS of the lesion area did not significantly change the size of the CCI lesion (b). Representative autoradiography images are shown of a control rat (c) and a rat that received motor cortex stimulation (d). The radioactive signal intensity (arbitrary units, a.u.) for the healthy (e) and lesioned hemisphere (f) separately did not differ between control rats and rats that were stimulated. Also no difference between groups was present in the ratio of the signal intensity between the lesioned hemisphere and the healthy hemisphere. CCI: controlled cortical impact; M: medial; L: lateral; D: dorsal; V: ventral.

sphere: $F(1,7)=0.61, p>0.05$; Fig. 3e-f], showing again that MCS treatment did not influence FDG distribution.

4. Discussion

Motor impairments are among the most debilitating consequences of stroke or TBI and have a strong impact on a patient's day-to-day activities. Electrical stimulation of the motor cortex has been shown to cause functional improvements in animal models of ischemic stroke and the ability of MCS to improve motor recovery in humans is currently under investigation (R. M. Levy et al., 2015; Shin, Dixon, Okonkwo, & Richardson, 2014). In rodent models of stroke, functional recovery was measurable as improved limb placement in response to a sensory cue, grasping or balance (Cheng et al., 2012; Moon et al., 2009). An increased formation of new blood vessels and dendritic sprouting has been found in addition to more NeuN-positive cells in the ischemic cortex (Cheng et al., 2012; Kang et al., 2013).

Although stroke and TBI have a different cause, both result in strikingly similar effects at the cellular level, such as excitotoxicity, oxidative stress and inflammatory responses (Lopez, Dempsey, & Vemuganti, 2015); therefore treatments effective in animal models of stroke might also be applicable to animal models of TBI. However, compared to stroke, endogenous plasticity processes are more limited after CCI lesions and more diverse behavioral rehabilitation training is necessary to induce morphological and functional rehabilitation (Combs, Jones, Kozlowski, & Adkins, 2016; Jones et al., 2012). In the specific case of MCS, stimulation parameters that induced robust recovery in stroke models were less effective in a TBI model with comparable motor impairments (Jefferson et al., 2015).

In the current study the potential of chronic MCS to achieve functional recovery after a cortical lesion was assessed. To induce the cortical lesion, rats received a CCI in the forelimb area of the motor cortex contralateral to the preferred limb. MCS was applied to freely moving rats during a period of 31 consecutive days and different aspects of forelimb function were assessed before and after the stimulation period. We measured fine motor skills with the Montoya staircase test and the cylinder test and both tests are sensitive ways to detect asymmetrical paw use after a cortical lesion (MacLellan, Langdon, Botsford, Butt, &

Corbett, 2013; L.M. Schönfeld et al., 2016; Windle et al., 2006). Furthermore, comparable behavioral tests were used in studies where a regenerative effect of MCS was detected (Morimoto et al., 2011; Zhou et al., 2010). In addition, we visualized the distribution of a radioactive glucose analog, FDG, to detect potential changes in brain metabolism after chronic MCS.

In line with previous research, we found that severe CCI created long-lasting motor impairments specific to the contralateral forelimb (L.M. Schönfeld et al., 2016), which could be detected for up to eight weeks after the insult. Motor impairments observed in patients with a cortical lesion usually have a chronic course; therefore modeling long-lasting motor impairments in animals is essential and can be achieved by severe CCI.

In the current study, we wanted to test the therapeutic potential of MCS as a standalone treatment administered in a home cage-environment. This procedure was different compared to previous research, where MCS was always administered together with physical rehabilitation training. In the current study, MCS as an independent treatment did not cause any improvement of motor impairments. After CCI, reaching and grasping skills with the impaired limb were equally affected in rats that received MCS compared to sham-stimulated control rats. Also, after CCI rats predominantly used their healthy forelimb during vertical exploration whereas usage of the other, impaired, forelimb was not restored after chronic MCS.

In line with these findings, no changes in either lesion size or FDG distribution were detected after the application of MCS. Changes in FDG distribution can be used as an indicator of neural activity (Gobel, Oltmanns, & Chung, 2013); therefore increased FDG distribution after MCS would have been an indirect measure of tissue restoration, whereas the absence of any MCS-induced change suggests a lack of treatment effect on energy metabolism.

A few studies on TBI in rats have reported a therapeutic effect of MCS (Jefferson et al., 2015; Yoon, Cho, Kim, Lee, & Lee, 2015; Yoon et al., 2012). In those studies, MCS was co-administered with behavioral rehabilitation training during a period ranging from two (Yoon et al., 2015; Yoon et al., 2012) to nine weeks (Jefferson et al., 2015). In all studies an improvement of forelimb function at the end of the stimulation period was reported (Jefferson et al., 2015; Yoon et al., 2015; Yoon et al.,

2012), together with an increase in size of the cortical area responsible for wrist movement, as assessed by intracortical microstimulation mapping (Jefferson et al., 2015). However, the TBI lesions created in these studies were considerably smaller compared to our lesions, which may have increased the likelihood of regeneration. Also, in these studies the implanted electrodes did not only cover the damaged cortex, but also stimulated spared cortical regions, which could facilitate re-mapping of lost functions onto the surrounding cortex. Lastly, MCS was always administered together with behavioral training, which is representative of the clinical situation, but does not provide any information about the therapeutic effect of MCS on its own. It is of note: three factors might explain why MCS did not induce functional regeneration in our study. First, the present CCI lesion was very severe and damaged the corpus callosum and parts of the striatum in addition to the entire forelimb area of the motor cortex. We chose for inducing such a severe CCI since previously we have been able to measure long-lasting behavioral impairments after this specific form of CCI (L.M. Schönfeld et al., 2016). In studies using milder CCI lesions in the motor cortex, spontaneous recovery of motor functions was measured, starting already at 2 weeks after the lesion (Goffus, Anderson, & Hoane, 2010; Nishibe et al., 2010; Shijo, Ghavim, Harris, Hovda, & Sutton, 2015). Taken together, behavioral improvement might be more likely after a milder lesion since the area to regenerate is smaller and surrounding cortical regions are spared which might allow functional re-mapping of the lost area.

Second, in numerous studies, in which MCS caused functional improvements, it was not chosen to stimulate the damaged brain area directly; instead stimulation electrodes were implanted on top of the lesion penumbra (Adkins et al., 2006; Boychuk, Adkins, & Kleim, 2011; Jefferson et al., 2015; Moon et al., 2009; O'Bryant et al., 2014; Plautz et al., 2003; Zhou et al., 2010). Electrical stimulation of the spared surrounding cortex may induce plasticity processes in contrast to the stimulation of a damaged brain region that during the course of several weeks develops into a large morphological cavity (L.M. Schönfeld et al., 2016). In a number of studies the therapeutic effect of MCS was not explained by tissue restoration at the lesion side, but by remapping of the lost functions onto the spared cortex around the lesion (Boychuk et al., 2011; Jefferson et al., 2015; Teskey et al., 2003). In the current study, the lesion area was stimulated directly to clarify whether the influence of MCS on

NSPCs found in an earlier experiment (Jahanshahi, Schonfeld, et al., 2013b) could rescue the damaged cortical tissue. In a previous study we showed an increased amount of NSPCs at the stimulated cortex of naïve rats, which presumably had migrated from the SVZ (Jahanshahi, Schonfeld et al., 2013b). Under *in vitro* conditions, electrical fields can induce migration of NSPCs towards the current source, a process known as electrotaxis (Babona-Pilipos, Droujinine, Popovic, & Morshead, 2011; Babona-Pilipos, Pritchard-Oh, Popovic, & Morshead, 2015; Feng et al., 2012; Liu et al., 2015; McCaig, Rajnicek, Song, & Zhao, 2005), and *in vitro* electrotaxis is a widely proven phenomenon occurring in different cells types, including NSPCs (Babona-Pilipos et al., 2015; Feng et al., 2012; Liu et al., 2015). Also, stimulation of the striatal penumbra after ischemic stroke in rats has been shown to increase the number of proliferating cells in the vicinity of the electrode and this finding co-occurred with a decreased lesion size and behavioral improvement (Morimoto et al., 2011). However, contrary findings have been reported as well. In a different study, a non-invasive form of cortical stimulation was applied in naïve rats and did not lead to directional migration of labeled neural stem cells in response to an electrical field (Keuters et al., 2015). The authors concluded that accumulation of neural stem cells at the stimulated cortical area is rather due to local cell proliferation and not to cell migration from neurogenic regions (Keuters et al., 2015). These contrary findings indicate, that *in vivo* electrotaxis first needs to be reliably demonstrated, before its role in stimulation-induced motor recovery can be investigated.

Third, in previous studies MCS has been delivered while the animals underwent rehabilitative therapy in the form of repetitive reaching with the impaired forelimb (Boychuk et al., 2011; Jefferson et al., 2015; Teskey et al., 2003). Probably an additional behavioral stimulus, that is stronger and more specific to the impaired forelimb than mere locomotion in a home cage, is necessary to cause improvement through MCS. Pairing of MCS with rehabilitation therapy may result in a synergistic effect and there are only a few reports showing a therapeutic effect of MCS without any additional intervention (Adkins-Muir & Jones, 2003; Adkins et al., 2006; Zhou et al., 2010).

In conclusion, the use of MCS as a standalone treatment did not improve motor impairments in a rat model of severe cortical damage given our specific treatment modalities.

Acknowledgments

This work was supported in part by a grant from Hasselt University (BOF11603BOFMO) to SH and a grant from the Hersenstichting Nederland and Medtronic Europe, (A1196117) to YT and AJ.

Disclosure statement

No competing financial interests exist.

References

- Adkins-Muir, D.L., & Jones, T.A. (2003). Cortical electrical stimulation combined with rehabilitative training: Enhanced functional recovery and dendritic plasticity following focal cortical ischemia in rats. *Neurological Research*, *25*(8), 780-788. doi: 10.1179/016164103771953853
- Adkins, D.L., Campos, P., Quach, D., Borromeo, M., Schallert, K., & Jones, T.A. (2006). Epidural cortical stimulation enhances motor function after sensorimotor cortical infarcts in rats. *Experimental Neurology*, *200*(2), 356-370. doi: 10.1016/j.expneurol.2006.02.131
- Baba, T., Kameda, M., Yasuhara, T., Morimoto, T., Kondo, A., Shingo, T., ... & Date, I. (2009). Electrical stimulation of the cerebral cortex exerts antiapoptotic, angiogenic, and anti-inflammatory effects in ischemic stroke rats through phosphoinositide 3-kinase/Akt signaling pathway. *Stroke*, *40*(11), e598-605. doi: 10.1161/STROKEAHA.109.563627
- Babona-Pilipos, R., Droujinine, I.A., Popovic, M.R., & Morshead, C.M. (2011). Adult subependymal neural precursors, but not differentiated cells, undergo rapid cathodal migration in the presence of direct current electric fields. *PLoS One*, *6*(8), e23808. doi: 10.1371/journal.pone.0023808
- Babona-Pilipos, R., Pritchard-Oh, A., Popovic, M.R., & Morshead, C.M. (2015). Biphasic monopolar electrical stimulation induces rapid and directed galvanotaxis in adult subependymal neural precursors. *Stem Cell Research and Therapeutics*, *6*, 67. doi: 10.1186/s13287-015-0049-6
- Boychuk, J.A., Adkins, D.L., & Kleim, J.A. (2011). Distributed versus focal cortical stimulation to enhance motor function and motor map plasticity in a rodent model of ischemia. *Neurorehabilitation and Neural Repair*, *25*(1), 88-97. doi: 10.1177/1545968310385126
- Brown, J.A., Lutsep, H.L., Weinand, M., & Cramer, S.C. (2006). Motor cortex stimulation for the enhancement of recovery from stroke: A prospective, multicenter safety study. *Neurosurgery*, *58*(3), 464-473. doi: 10.1227/01.NEU.0000197100.63931.04
- Cheng, X., Li, T., Zhou, H., Zhang, Q., Tan, J., Gao, W., ... & Duan, Y.Y. (2012). Cortical electrical stimulation with varied low frequencies promotes functional recovery and brain remodeling in a rat model of ischemia. *Brain Research Bulletin*, *89*(3-4), 124-132. doi: 10.1016/j.brainresbull.2012.07.009
- Feng, J.F., Liu, J., Zhang, X.Z., Zhang, L., Jiang, J.Y., Nolta, J., & Zhao, M. (2012). Guided migration of neural stem cells derived from human embryonic stem cells by an electric field. *Stem Cells*, *30*(2), 349-355. doi: 10.1002/stem.779

- Gobel, B., Oltmanns, K.M., & Chung, M. (2013). Linking neuronal brain activity to the glucose metabolism. *Theoretical Biology and Medical Modeling*, *10*, 50. doi: 10.1186/1742-4682-10-50
- Goffus, A.M., Anderson, G.D., & Hoane, M. (2010). Sustained delivery of nicotinamide limits cortical injury and improves functional recovery following traumatic brain injury. *Oxidative Medicine and Cellular Longevity*, *3*(2), 145-152. doi: 10.4161/oxim.3.2.11315
- Harris, N.G., Mironova, Y.A., Hovda, D.A., & Sutton, R.L. (2010). Chondroitinase ABC enhances pericontact axonal sprouting but does not confer robust improvements in behavioral recovery. *Journal of Neurotrauma*, *27*(11), 1971-1982. doi: 10.1089/neu.2010.1470
- Jahanshahi, A., Schönfeld, L., Janssen, M.L., Heschem, S., Kocabicak, E., Steinbusch, H.W., ... & Temel, Y. (2013). Electrical stimulation of the motor cortex enhances progenitor cell migration in the adult rat brain. *Experimental Brain Research*, *231*(2), 165-177. doi: 10.1007/s00221-013-3680-4
- Jahanshahi, A., Schönfeld, L.M., Lemmens, E., Hendrix, S., & Temel, Y. (2013). In vitro and in vivo neuronal electrotaxis: A potential mechanism for restoration? *Molecular Neurobiology*. doi: 10.1007/s12035-013-8575-7
- Jefferson, S.C., Clayton, E.R., Donlan, N.A., Kozłowski, D.A., Jones, T.A., & Adkins, D.L. (2015). Cortical stimulation concurrent with skilled motor training improves forelimb function and enhances motor cortical reorganization following controlled cortical impact. *Neurorehabilitation and Neural Repair*. doi: 10.1177/1545968315600274
- Kang, C., Yang, C.Y., Kim, J.H., Moon, S.K., Lee, S., Park, S.A., ... & Zhang, L.Q. (2013). The effect of continuous epidural electrical stimulation on neuronal proliferation in cerebral ischemic rats. *Annals of Rehabilitation Medicine*, *37*(3), 301-310. doi: 10.5535/arm.2013.37.3.301
- Keuters, M.H., Aswendt, M., Tennstaedt, A., Wiedermann, D., Pikhovych, A., Rothues, S., ... & Rueger, M.A. (2015). Transcranial direct current stimulation promotes the mobility of engrafted NSCs in the rat brain. *NMR in Biomedicine*, *28*(2), 231-239. doi: 10.1002/nbm.3244
- Levy, R., Ruland, S., Weinand, M., Lowry, D., Dafer, R., & Bakay, R. (2008). Cortical stimulation for the rehabilitation of patients with hemiparetic stroke: A multicenter feasibility study of safety and efficacy. *Journal of Neurosurgery*, *108*(4), 707-714. doi: 10.3171/JNS/2008/108/4/0707
- Levy, R.M., Harvey, R.L., Kissela, B.M., Winstein, C.J., Lutsep, H.L., Parrish, T.B., ... & Venkatesan, L. (2015). Epidural electrical stimulation for stroke rehabilitation: Results of the prospective, multicenter, randomized, single-blinded everest trial. *Neurorehabilitation and Neural Repair*. doi: 10.1177/1545968315575613
- Liu, J., Zhu, B., Zhang, G., Wang, J., Tian, W., Ju, G., ... & Song, B. (2015). Electric signals regulate directional migration of ventral midbrain derived dopaminergic neural progenitor cells via Wnt/GSK3beta signaling. *Experimental Neurology*, *263*, 113-121. doi: 10.1016/j.expneurol.2014.09.014
- Lopez, M.S., Dempsey, R.J., & Vemuganti, R. (2015). Resveratrol neuroprotection in stroke and traumatic CNS injury. *Neurochemistry International*, *89*, 75-82. doi: 10.1016/j.neuint.2015.08.009

- 667 MacLellan, C.L., Langdon, K.D., Botsford, A., Butt, S., &
668 Corbett, D. (2013). A model of persistent learned nonuse
669 following focal ischemia in rats. *Neurorehabilitation
670 and Neural Repair*, 27(9), 900-907. doi: 10.1177/
671 1545968313496323
- 672 McCaig, C.D., Rajnicek, A.M., Song, B., & Zhao, M. (2005).
673 Controlling cell behavior electrically: Current views and
674 future potential. *Physiological Reviews*, 85(3), 943-978. doi:
675 10.1152/physrev.00020.2004
- 676 Montoya, C.P., Campbell-Hope, L.J., Pemberton, K.D., & Dunnett,
677 S.B. (1991). The "staircase test": A measure of independent
678 forelimb reaching and grasping abilities in rats. *Journal of
679 Neuroscience Methods*, 36(2-3), 219-228.
- 680 Moon, S.K., Shin, Y.I., Kim, H.I., Kim, H., Lee, J.O., &
681 Lee, M.C. (2009). Effect of prolonged cortical stimula-
682 tion differs with size of infarct after sensorimotor cortical
683 lesions in rats. *Neuroscience Letters*, 460(2), 152-155. doi:
684 10.1016/j.neulet.2009.05.029
- 685 Morimoto, T., Yasuhara, T., Kameda, M., Baba, T., Kuramoto, S.,
686 Kondo, A., ... & Date, I. (2011). Striatal stimulation nurtures
687 endogenous neurogenesis and angiogenesis in chronic-phase
688 ischemic stroke rats. *Cell Transplantation*, 20(7), 1049-1064.
689 doi: 10.3727/096368910X544915
- 690 Nishibe, M., Barbay, S., Guggenmos, D., & Nudo, R.J. (2010).
691 Reorganization of motor cortex after controlled cortical
692 impact in rats and implications for functional recovery. *Journal of Neurotrauma*, 27(12), 2221-2232. doi:
693 10.1089/neu.2010.1456
- 694 O'Bryant, A.J., Adkins, D.L., Sitko, A.A., Combs, H.L.,
695 Nordquist, S.K., & Jones, T.A. (2014). Enduring post-
696 stroke motor functional improvements by a well-timed
697 combination of motor rehabilitative training and cortical stimu-
698 lation in rats. *Neurorehabilitation and Neural Repair*. doi:
699 10.1177/1545968314562112
- 700 Parker, V.M., Wade, D.T., & Langton Hewer, R. (1986). Loss
701 of arm function after stroke: Measurement, frequency, and
702 recovery. *International Rehabilitation Medicine*, 8(2), 69-73.
- 703 Plautz, E.J., Barbay, S., Frost, S.B., Friel, K.M., Dancause, N.,
704 Zoubina, E.V., ... & Nudo, R.J. (2003). Post-infarct cortical
705 plasticity and behavioral recovery using concurrent cortical
706 stimulation and rehabilitative training: A feasibility study in
707 primates. *Neurological Research*, 25(8), 801-810.
- 708 Rueger, M.A., Keuters, M.H., Walberer, M., Braun, R., Klein,
709 R., Sparing, R., ... & Schroeter, M. (2012). Multi-session
710 transcranial direct current stimulation (tDCS) elicits inflam-
711 matory and regenerative processes in the rat brain. *PLoS One*,
712 7(8), e43776. doi: 10.1371/journal.pone.0043776
- Schönfeld, L.M., Jahanshahi, A., Lemmens, E., Schipper, S., Doo-
714 ley, D., Joosten, E., ... & Hendrix, S. (2016). Long-term motor
715 deficits after controlled cortical impact in rats can be detected
716 by fine motor skill tests but not by automated gait analysis.
717 *Journal of Neurotrauma*. doi: 10.1089/neu.2016.4440
718
- 719 Shijo, K., Ghavim, S., Harris, N.G., Hovda, D.A., & Sutton, R.L.
720 (2015). Glucose administration after traumatic brain injury
721 exerts some benefits and no adverse effects on behavioral and
722 histological outcomes. *Brain Research*, 1614, 94-104. doi:
723 10.1016/j.brainres.2015.04.022
- 724 Shin, S.S., Dixon, C.E., Okonkwo, D.O., & Richardson,
725 R.M. (2014). Neurostimulation for traumatic brain
726 injury. *Journal of Neurosurgery*, 121(5), 1219-1231. doi:
727 10.3171/2014.7.JNS131826
- 728 Starkey, M.L., Bleul, C., Zorner, B., Lindau, N.T., Mueggler,
729 T., Rudin, M., & Schwab, M.E. (2012). Back seat driving:
730 Hindlimb corticospinal neurons assume forelimb control fol-
731 lowing ischaemic stroke. *Brain*, 135(Pt 11), 3265-3281. doi:
732 10.1093/brain/aws270
- 733 Teskey, G.C., Flynn, C., Goertzen, C.D., Monfils, M.H., & Young,
734 N.A. (2003). Cortical stimulation improves skilled forelimb
735 use following a focal ischemic infarct in the rat. *Neurological
736 Research*, 25(8), 794-800.
- 737 Walker, W.C., & Pickett, T.C. (2007). Motor impairment after
738 severe traumatic brain injury: A longitudinal multicenter
739 study. *Journal of Rehabilitation Research and Development*,
740 44(7), 975-982.
- 741 Windle, V., Szymanska, A., Granter-Button, S., White, C., Buist,
742 R., Peeling, J., & Corbett, D. (2006). An analysis of four
743 different methods of producing focal cerebral ischemia with
744 endothelin-1 in the rat. *Experimental Neurology*, 201(2), 324-
745 334. doi: 10.1016/j.expneurol.2006.04.012
- 746 Yoon, Y.S., Cho, K.H., Kim, E.S., Lee, M.S., & Lee, K.J. (2015).
747 Effect of Epidural Electrical Stimulation and Repetitive Tran-
748 scranial Magnetic Stimulation in Rats With Diffuse Traumatic
749 Brain Injury. *Annals of Rehabilitation Medicine*, 39(3), 416-
750 424. doi: 10.5535/arm.2015.39.3.416
- 751 Yoon, Y.S., Yu, K.P., Kim, H., Kim, H.I., Kwak, S.H., & Kim,
752 B.O. (2012). The effect of electric cortical stimulation after
753 focal traumatic brain injury in rats. *Annals of Rehabilitation
754 Medicine*, 36(5), 596-608. doi: 10.5535/arm.2012.36.5.596
- 755 Zhou, Q., Zhang, Q., Zhao, X., Duan, Y.Y., Lu, Y., Li, C., &
756 Li, T. (2010). Cortical electrical stimulation alone enhances
757 functional recovery and dendritic structures after focal cere-
758 bral ischemia in rats. *Brain Research*, 1311, 148-157. doi:
759 10.1016/j.brainres.2009.11.022