REVIEW ARTICLE

A Novel Intervention Technology for Cerebral Palsy: Brain Stimulation

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Abstract

A common pediatric disorder with posture and motor dysfunction in neurological diseases is known as cerebral palsy (CP). Recently, a series of effective techniques have been developed for treatment of CP. These promising methods need high-tech equipment for brain stimulation and mainly classified into invasive and no-invasive approaches. This study aimed to introduce these techniques for treatment of patients who suffer from CP. The potential and performance of currently available brain stimulation techniques have been mentioned in detail. Moreover, the clinical application, safety, efficacy and challenges of these methods have been discussed. Here we review the recent advances in the CP treatment with an emphasis on brain stimulation techniques

Keywords: Cerebral palsy; Brain stimulation; Pediatric disorder

Introduction

A common pediatric disorder with posture and motor dysfunction in neurological diseases is known as cerebral palsy (CP) (1-3). Neurological disorders commonly appeared in early stage of human life statically reported about 3 to 4 cases in 1000 newborn (4, 5). These patients mainly suffer from other problems like orthopedically disorders followed by neurological dysfunctions, unfortunately, that affects their normal life (1). Around 50% of these patients show cognitive deficits. Furthermore, one-third of children suffer from seizure attacks (6, 7). Nowadays, advanced technologies in the field of brain imaging and stimulation have been introduced to medicine for diagnosis and treatment. In addition, cellular therapy methods associated with novel advanced techniques can be useful for treating these disabilities (8-15).

The aim of this review is to propose effectiveness, profits, and detriments of brain stimulation techniques in CP. This paper runs over the various research efforts within this paradigm reported to date and attempts a selection of the appropriate investigation in this field that are shown in Table 1. Brain stimulation can play a role in remedying CP. Brain tissue can be directly stimulated with electricity. In these non-invasive techniques electrodes directly put in the target site in the scalp and then magnetic field created in the head using magnetic circles (9). We aimed to introduce all advanced techniques clinically used for brain stimulation in neurological disorders such as CP.

Invasive procedure for brain stimulation

Deep brain stimulation (DBS)

In this technique, precise electrical impulses directly conducted into the defect area using quad polar electrodes implanted into the brain. Electrical stimulation by this method modulates cellular functions via cell-cell signaling and communication and signaling molecules and chemical mediators. Currently available DBS devices in the market composed of pulse generator set and two electrode leads. Pulse generator set usually fixes in subclavicular region adjusted externally and two electrode leads fix in the brain (16-18). This devise act similar to cardiac pacemaker with stimulating and inhibiting activity (16). This adjustment was managed with patient symptoms. The DBS device can be used for improving diatonic and tremor symptoms in Parkinson disorders. Additionally, this approach has been recommended for obsessive-compulsive disease. The adverse sign of neurodegenerative disorders diminishes following the application of this procedure. On the other

hand, DBS is significantly better than the ablative surgery via safety and efficacy (17).

Chronic pain was decreased using DBS for thalamic stimulation (19, 20). Moreover, group of scientists worked on decrease of tremor by thalamic stimulation. DBS had effective impact on tremor and pain (21, 22). In addition, application of this technique on subthalamic nucleus (STN) significantly decreased bradykinesia, tremor and rigidity (23, 24). The satisfied results have been described for the use of DBS in hippocampus and STN for treatment of epilepsy (25). In recent years, high percent of children with dystonia had effective response to this therapeutic method (15, 26). In 2008, a study described the effect of DBS on patient with secondary dystonia. The symptom of disease was improved following using the DBS method. Therefore, this method has been recommended for treatment of various forms of dystonia (27, 28). Although the issue of DBS application for child has received critical attention, but it is in quiet in early stage.

The last decades have shown a growing trend toward using DBS technique for dystonia complaints in pediatric population (6). The healing potential of DBS was investigated in secondary dystonia associated with CP. Globus pallidus stimulation by DBS method promoted the movement disabilities and dyskinesia in comparison with patients treated with traditional methods such as drug therapy (16). In addition, bilateral pallial DBS could be an alternative method for patients who suffer from dystonia-chorea and CP (29).

The optimum therapeutic spot in these group of sick people is the posterior lateroventral region of Globus pallidus internus (GPi). Diffusion of

the stimulation to adjacent structures (mainly Globus pallidus externus), may bring out the little recovery (30). DBS can offer significant changes in multiple domains of general health, dysfunctions and disabilities. Thus, the sequential assessments to evaluate the clinical utilities following DBS via rating scales particularly in children with CP are obligatory (26, 31, 32).

One of the main obstacles faced by many experiments is superficial or deep hematoma after DBS treatment. Additionally, local infection and erosion have existed as a health problem caused by DBS intervention. Another major issue that can impair medical impediment is the digging the electrodes from the skull to the trunk to insert the device in the chest (17).

Non-invasive procedure for brain stimulation

Direct brain Stimulation

The brain can be provoked with direct method via transcranial fascinated much attention. The past twenty years showed increasingly rapid advances in the field of rehabilitative interventions (33, 34). The direct brain stimulation (TDCS) has more effective than the other approaches for brain stimulation. This method is completely non-invasive and consists of two electrodes, one of them provokes cortical part and the other provokes inhibitory function, to stimuli brain in contact with scalp (35).

Recently an advanced model of TDCS system has been developed and showed some advantages comparing to other methods including safety, cost-effectiveness and portable, permitting neurologist to apply brain stimulation associated with exercise therapy in rehabilitation centers (36-39). The TDCS system targets the regional synaptic in the brain to control

the cortical excitability. Not only local cortical stimulation induced by TDCS is not stable, but also it is prompted through weak electrical stimuli to the scalp basically through employment of electrodes (35, 37, 40). A considerable amount of literature have been published on applying the TDCS technique for treatment of neurological disorders (40). Furthermore, this technique has been identified as safe and efficient for pediatric patients. Recently erythematous rash was identified as a side effect in pediatric TDCS treatment (41).

In an original investigation of the efficacy of TDCS treatment on children (between 5 to 12 yr old), the treatment regime was 2 mA for 30 min in 10 sessions. The result of this study represented that children mainly suffered from tingling and itching, mood disorders and also irritability with 28.6%, 42.9% and 36%, respectively (42). The intensity of stimulation was examined in TDCS system. The optimization of stimuli signals via TDCS according to the age of recipient have been recommended especially for children (43). However, research was conducted on child with perinatal ischemic stroke and hemiparesis disorder. Other parameters such as dose play an essential role in child-specific TDCS protocols. In this experiment, magnetic resonance imaging (MRI) was used to evaluate the current flow. They checked some important factors including electrode size and location, dose strength and time period. Computational modeling is necessary to calculate effective dose in order to enhance healing in pediatric stroke through the TDCS guidance (44). Much of available literature on the potential capacity of TDCS technique to increase motor learning in adults and also high clinical studies have reported positive data on motor recovery and reestablishing the equilibrium of the motivation between two hemispheres (45,

46). Case studies on patients containing damaged neurons in hemisphere showed that the TDCS technique can successfully stimulate damaged site in the brain without any side effect on normal tissue. Moreover, many studies have been carried out to restore motor learning disorders thought contralesional stimulation and neuroplasticity pathways (47-49). The mutual impact of anodal TDCS and virtual reality for accelerating walk were evaluated in patient with spastic deformities. In addition, a pilot clinical trial study was deliberate to assess the efficacy of this method in a rehabilitation center. The treated group with both anodal stimulation and virtual reality had satisfied results and improvement. The anodal stimulation together with virtual reality can significantly recover movement in CP patient with spastic di-paretic (26). Combination therapy for example using TDCS with 1 mA strength ended the conquering primary motor cortex and physical training via treadmill for twenty min for 10 d accelerate motor function in patient with spastic diparetic cerebral palsy. The positive impact of the TDCS technique combined with physical training examined on twenty four patients with spastic diparetic. The temporal function of treated children has been increased (9). The TDCS method associated with physical training augmented some circuits our brain including mediolateral and anterior-posterior.

This research also established the functional effects of the anodal TDCS technique over primary motor cortex through movement exercise (28). Additionally, the impact of the cathodal TDCS was examined to improve the voluntary activities in patients suffer from the dystonia. Hand function with controlled manner was increased followed by cathode stimulation of healthy hemisphere (50, 51). Moreover, this technique cannot be clinically

performed for improving dystonia problem. They carried out a double-blind experiment to assess the impact of the TDCS method on dystonia treatment. The TDCS regime was programmed using 2 mA intensity on cortex for about ten min. The prescribed regime resulted in the symptoms lessening in some patients (51, 52). Gillick et al showed some advantages of the TDCS methods for movement rehabilitation in young and old patient in a control trial study (8, 53). Taken together, many studies and guidelines have been published to validate the efficacy, safety and practicability of the TDCS technique in human. The currently existing data approve the admissibility and the applicability of the TDCS method in CP patient. The published experiments have suggested some signs such as short-lived tingling or slight itching (33, 36, 54). The prescription of this method in CP patient is recommended for many rehabilitation centers.

Magnetic Stimulation of brain

In recent years, a novel approach has been introduced in the field of neuroscience known as transcranial magnetic stimulation (55). Abundant studies have been carried out using TMS for investigating neuroplasticity after brain injuries. Magnetic stimulation of brain is a very simple and safe technique. Electromagnetic field is created following TMS induction in the brain tissue (56). In this system, a figure-eight conductive coil was used to conduct electrical current to the neural cells. This magnetic stimulation induces local electrical variations in cortical neural cells based on Faraday's Law. Cortical excitability and eternal modifications in neural behavior can be regulated via monotonous stimulation with TMS (57). However, cortical excitability was assisted or limited via modulation of the intensity of the

repetitive TMS (rTMS) pulses (58, 59).

The main principle of the rTMS is decreasing of the brain function (60) and augments cortical excitability through interhemispheric limitation (61). In early times, Magnetic stimulation of brain was used to facilitate the prediction and improvement after stroke attacks (62, 63) and also neuropsychiatric diseases (64). However, many studies have documented the unknown complications after using the rTMS technique for patients with severe neurological, physiological and psychiatric disorders (64, 65). The application of this system in children is recommended for management of hyperactivity disorder (ADHD) and autism (ASD)(66, 67).

The repetitive TMS has been suggested in high frequency (>5-10 Hz) for treatment of stroke (68-70). Up to now, there is no evidence about any adverse symptoms for rTMS usage (71-73). The use of this advanced technique in grownups (74-77) and patient with stroke are benign and acceptable (78). Both methodological and safety concern are considered as the main obstacle for using rTMS in children. Although it is more effective for movement improvement results in stroke attacks (72, 73, 79, 80), more common disorders in pediatric, treated with rTMS technique, and are pediatric stroke and spasticity (72). The therapeutic effects were tested of the low-frequency rTMS on patient (n=10) suffered from chronic subcortical Arterial Ischemic

Stroke (AIS) with motor neuron dysfunction and dyskinesia. In this project patient were divided into two groups, half of them received lowfrequency rTMS and the others were considered as a control group. The patients in experiment group were treated with 1200 stimuli for about 7 days. The adverse side effects were not reported in this study and functional improvements, as well as appropriate clinical outcomes, were reported after 7 d follow up post-treatment (78). Cortical stimulation was suggested for children spasticity using the rTMS technique. A study on seventeen patient showed spastic quadriplegia symptoms using repetitive TMS with low and high-frequency intensity (81). rTMS- excited cortical neurons act normally and spasticity improvement can be seen after optimized stimulation(82, 83). Additionally, in 2012 a phase II clinical trial for evaluation of the rTMS technique plus occupational therapy on patients suffered from chronic hemiparesis was conducted (84).

Seizure known as an adverse side effect which was no reported following the use of the rTMS technique for pediatric but headache and local scalp pain were seen in some patient. This method causes transient tinnitus and threshold and further evaluation should be done (85-87).

The application of this technique has been inhibited for patients having implants in their bodies.

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Table 1: A brief summary of the recent papers on brain stimulation

Author (date)	Sample	Method	Results
Warren et al (2009)	Child with sec- ondary dystonia	DBS	Improvement of motor function
Vidailhet et al (2009)	Child with dystonia-chore- athesosis CP	Bilateral pallidal-DBS	Improvement of motor
Berweck et al (2009)	Child with dystonia-chore- athesosis CP	Bilateral pallidal-DBS	Improvement of motor function
Young et al (2013)	11 patients with dystonia aged 7-19 yrs	Cathodal TDCS, over motor cortex at C_3 or C_4 , 1 ma for 18 min with a 20 min pause interval	Reduction of involuntary over flow activity in a sub- set of children
Auvichayapat et al (2014)	46 CP patients Aged 8-18 yrs	Anodal TDCS, over left primary motor cortex at M ₁ , 1 ma for 20 min on five consecutive days	Sign. Reduction in CP- related spasticity (but not in PROM)
Duarte et al (2014)	24 CP children Aged 5-12 yrs	Anotal TDCS + treadmill training, over primary motor cortex at M _{1,} 1 ma for 20 min on five weekly sessions for 2 weeks	Sign. Improvements In static balance and functional performance
Grecco et al (2015)	20 children with diparetic CP Aged 5-10 yrs	Anotal TDCS + vitual reality, over primary motor cortex at M _{1,} 1 ma for 20 min on five weekly sessions for 2 weeks	Sign. Improvement in gait velocity, cadence, gross motor function and inde- pendent mobility Sign. Increase in MEP
Gillick et al (2015)	20 patients with hemiparesis Aged 8-21yrs	Ipsilesional anodal and contralesional cathodal TDCS + CIMT, over primary motor cortex at M _{1,} 0.7 ma for ten 2-hours sessions. TDCS was performed in first 20 min of period.	Sign. Improvement in motor functions
Bhanpuri Et al (2015)	9 patients with dystonia Aged 10-21yrs	Anodal and cathodal TDCS, over motor cortex at C_3 or C_4 , 2 ma, 9 minutes per day for 5 days	No sign. Changes in dystonia
Bernadette et al (2015)	19 patients with congenital hemiparesis aged 8-17 yrs	5 treatment alternate weekday over 2 weeks rtms for either combined with modified constraint induced-movement therapy(mcimt) or sham rtms combined with mcimt	Minor but not major adverse events such as headaches, cast irritation were found in both experimental and sham groups.
Valle et al (2007)	17 children with spastic CP aged 6-12 yrs quadri- paresia	High frequency (5 Hz) rtms, low frequency (1 Hz) rtms , sham rtms over CL ABP motor area, for 5 days	Improvement in elbow movement by high frequency rtms

In conclusion, the present study provides additional evidence with respect to brain stimulation techniques in patient with neurodegenerative disorders. Neuroscientists reported satisfied results about using advanced technologies including DBS, TDCS, and rTMS for treatment of cerebral stroke, spasticity and patients with abnormal motor function. On the other hand, the safety, efficacy and also no adverse side effects such as seizure following the use of these advanced methods have been discussed in the literature. Further research regarding the role of brain stimulation would be of great help in treatment of neurological disorders.

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Author's contribution

Naser Amini(designing the study), Agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved, Final approval of the version to be published

Arash Sarveazad (data acquisition), Fateme Khodaei(analysis), Sara Ramezani (interpretation of data): Substantial contributions to the conception or design of the work; or the acquisition, analysis, or interpretation of data for the work

Hosein Safakheil, Peiman Brouki Milan: Drafting the work or revising it critically for important intellectual content

Lena Mohebi, Final approval of the version to be published:

All authors agreed to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Conflict of interest

The authors declare that there is no conflict of interests.

References

- 1. Aisen ML, Kerkovich D, Mast J, et al. Cerebral palsy: clinical care and neurological rehabilitation. Lancet Neurol 2011: 10: 844-852.
- 2. Pervin R, Ahmed S, Hyder RT, Yasmeen BN, Rahman M, Islam F. Cerebral palsy-an update. Northern Int Med College J 2013: 5: 293-296.
- 3. Reid LB, Rose SE, Boyd RN. Rehabilitation and neuroplasticity in children with unilateral cerebral palsy. Nature Rev Neurol 2015: 11: 390.
- 4. Stanley FJ, Blair E, Alberman E. Cerebral palsies: epidemiology and causal pathways: Cambridge University Press, 2000.
- 5. Yeargin-Allsopp M, Braun KVN, Doernberg NS, Benedict RE, Kirby RS, Durkin MS. Prevalence of cerebral palsy in 8-year-old children in three areas of the United States in 2002: a multisite collaboration. Pediatrics 2008: 121: 547-554.
- 6. Kirby RS, Wingate MS, Braun KVN, et al. Prevalence and functioning of children with cerebral palsy in four areas of the United States in 2006: a report from the Autism and Developmental Disabilities Monitoring Network. Res Develop Disabil 2011: 32: 462-469.
- 7. Sigurdardottir S, Vik T. Speech, expressive language, and verbal cognition of preschool children with cerebral palsy in Iceland. Develop Med Child Neurol 2011: 53: 74-80.
- 8. Gillick B, Menk J, Mueller B, et al. Synergistic effect of combined transcranial direct current stimulation/constraint-induced movement therapy in children and young adults with hemiparesis:

- study protocol. BMC Pediatr 2015: 15: 178.
- 9. Grecco LAC, Duarte NdAC, Mendonça ME, et al. Transcranial direct current stimulation during treadmill training in children with cerebral palsy: a randomized controlled double-blind clinical trial. Res Develop Disabil 2014: 35: 2840-2848.
- 10. Taub E, Uswatte G, Elbert T. New treatments in neurorehabiliation founded on basic research. Nature Rev Neurosci 2002: 3: 228.
- 11. Wolf SL, Winstein CJ, Miller JP, et al. Retention of upper limb function in stroke survivors who have received constraint-induced movement therapy: the EXCITE randomised trial. Lancet Neurol 2008: 7: 33-40.
- 12. Sarveazad A, Babahajian A, Bakhtiari M, et al. The combined application of human adipose derived stem cells and Chondroitinase ABC in treatment of a spinal cord injury model. Neuropeptides 2017: 61: 39-47.
- 13. Amini N, Vousooghi N, Hadjighassem M, et al. Efficacy of human adipose tissue-derived stem cells on neonatal bilirubin encephalopathy in rats. Neurotoxicity Res 2016: 29: 514-524.
- 14. Yousefzadeh-Chabok S, Ramezani S, Reihanian Z, Safaei M, Alijani B, Amini N. The role of early posttraumatic neuropsychological outcomes in the appearance of latter psychiatric disorders in adults with brain trauma. Asian J Neurosurg 2015: 10: 173.
- 15. Amini N, Vousooghi N, Soleimani M, et al. A new rat model of neonatal bilirubin encephalopathy (kernicterus). J Pharmacol Toxicol Methods 2017: 84: 44-50.
- Marks WA, Honeycutt J, Acosta F, Reed M.
 Deep brain stimulation for pediatric movement disorders. Seminars in Pediat Neurol 2009: 90-98.
- 17. Kupsch A, Benecke R, Müller J, et al. Palli-

- dal deep-brain stimulation in primary generalized or segmental dystonia. N Eng J Med 2006: 355: 1978-1990.
- 18. Schwalb JM, Riina HA, Skolnick B, Jaggi JL, Simuni T, Baltuch GH. Revision of deep brain stimulator for tremor. J Neurosurg 2001: 94: 1010-1012.
- 19. Hosobuchi Y, Adams JE, Rutkin B. Chronic thalamic stimulation for the control of facial anesthesia dolorosa. Arch Neurol 1973: 29: 158-161.
- 20. Mazars G, Merienne L, Cioloca C. Treatment of certain types of pain with implantable thalamic stimulators. Neuro-Chirurgie 1974: 20: 117.
- 21. Benabid AL, Pollak P, Hoffmann D, et al. Long-term suppression of tremor by chronic stimulation of the ventral intermediate thalamic nucleus. The Lancet 1991: 337: 403-406.
- 22. Burchiel KJ. Thalamotomy for movement disorders. Neurosurgery Clinics of North America 1995: 6: 55-71.
- 23. Hamani C, Richter E, Schwalb JM, Lozano AM. Bilateral subthalamic nucleus stimulation for Parkinson's disease: a systematic review of the clinical literature. Neurosurgery 2005: 56: 1313-1324.
- 24. Yu H, Neimat JS. The treatment of movement disorders by deep brain stimulation. Neurotherapeutics 2008: 5: 26-36.
- 25. Halpern CH, Samadani U, Litt B, Jaggi JL, Baltuch GH. Deep brain stimulation for epilepsy. Neurotherapeutics 2008: 5: 59-67.
- 26. Gimeno H, Tustin K, Selway R, Lin J-P. Beyond the Burke–Fahn–Marsden Dystonia Rating Scale: deep brain stimulation in childhood secondary dystonia. Euro J Paediatr Neurol 2012: 16: 501-508.
- 27. Katsakiori P, Kefalopoulou Z, Markaki E, et al. Deep brain stimulation for secondary dystonia:

- results in 8 patients. Acta Neurochirurgica 2009: 151: 473.
- 28. Ostrem JL, Starr PA. Treatment of dystonia with deep brain stimulation. Neurotherapeutics 2008: 5: 320-330.
- Berweck S. BP-DBS for dystonia-choreoathetosis cerebral palsy. Lancet Neurol 2009: 8: 692-693.
- 30. Vidailhet M, Yelnik J, Lagrange C, et al. Bilateral pallidal deep brain stimulation for the treatment of patients with dystonia-choreoathetosis cerebral palsy: a prospective pilot study. Lancet Neurol 2009: 8: 709-717.
- 31. Krauss JK, Loher TJ, Weigel R, Capelle HH, Weber S, Burgunder J-M. Chronic stimulation of the globus pallidus internus for treatment of non-dYT1 generalized dystonia and choreoathetosis: 2-year follow up. J Neurosurg 2003: 98: 785-792.
- 32. Preto T, Dalvi A, Kang U, Penn R. A prospective blinded evaluation of deep brain stimulation for the treatment of secondary dystonia nad primary torticollis syndromes. J Neurosurg 2008: 109: 405-409.
- 33. Brunoni AR, Nitsche MA, Bolognini N, et al. Clinical research with transcranial direct current stimulation (tDCS): challenges and future directions. Brain Stimulation 2012: 5: 175-195.
- 34. Jacobson L, Koslowsky M, Lavidor M. tDCS polarity effects in motor and cognitive domains: a meta-analytical review. Experimental Brain Res 2012: 216: 1-10.
- 35. Zaghi S, Acar M, Hultgren B, Boggio PS, Fregni F. Noninvasive brain stimulation with low-intensity electrical currents: putative mechanisms of action for direct and alternating current stimulation. The Neuroscientist 2010: 16: 285-307.
- 36. Brunoni AR, Amadera J, Berbel B, Volz MS, Rizzerio BG, Fregni F. A systematic review on

- reporting and assessment of adverse effects associated with transcranial direct current stimulation. Int J Neuropsychopharmacol 2011: 14: 1133-1145.
- 37. Nitsche MA, Paulus W. Excitability changes induced in the human motor cortex by weak transcranial direct current stimulation. J Physiol 2000: 527: 633-639.
- 38. Nitsche MA, Cohen LG, Wassermann EM, et al. Transcranial direct current stimulation: state of the art 2008. Brain Stimulation 2008: 1: 206-223.
- 39. Poreisz C, Boros K, Antal A, Paulus W. Safety aspects of transcranial direct current stimulation concerning healthy subjects and patients. Brain Res Bulletin 2007: 72: 208-214.
- 40. Moura RCF, Santos C, Collange Grecco L, et al. Effects of a single session of transcranial direct current stimulation on upper limb movements in children with cerebral palsy: A randomized, sham-controlled study. Develop Neurorehabil 2017: 20: 368-375.
- 41. Auvichayapat N, Amatachaya A, Auvichayapat P. Reduction of spasticity in cerebral palsy by anodal transcranial direct current stimulation. J Med Assoc Thai 2014: 97: 954-962.
- 42. Erwin B, Jr MM, Baker KK. Mechanisms of deep brain stimulation and future technical developments. Neurol Res 2000: 22: 259-266.
- 43. Moliadze V, Schmanke T, Andreas S, Lyzhko E, Freitag CM, Siniatchkin M. Stimulation intensities of transcranial direct current stimulation have to be adjusted in children and adolescents. Clin Neurophysiol 2015: 126: 1392-1399.
- 44. Gillick BT, Kirton A, Carmel JB, Minhas P, Bikson M. Pediatric stroke and transcranial direct current stimulation: methods for rational individualized dose optimization. Frontiers Human Neurosci 2014: 8: 739.

- 45. Fregni F, Boggio PS, Mansur CG, Wagner T, Ferreira MJ, Lima MC et al. Transcranial direct current stimulation of the unaffected hemisphere in stroke patients. Neuroreport 2005;16(14):1551-5.
- 46. Reis J, Robertson EM, Krakauer JW, et al. Consensus: can transcranial direct current stimulation and transcranial magnetic stimulation enhance motor learning and memory formation? Brain Stimulation 2008: 1: 363-369.
- 47. Marquez J, van Vliet P, McElduff P, Lagopoulos J, Parsons M. Transcranial direct current stimulation (tDCS): does it have merit in stroke rehabilitation? A systematic review. Int J Stroke 2015: 10: 306-316.
- 48. Fritsch B, Reis J, Martinowich K, et al. Direct current stimulation promotes BDNF-dependent synaptic plasticity: potential implications for motor learning. Neuron 2010: 66: 198-204.
- 49. Wittenberg GF. Neural plasticity and treatment across the lifespan for motor deficits in cerebral palsy. Develop Med Child Neurol 2009: 51: 130-133.
- 50. Young SJ, Bertucco M, Sanger TD. Cathodal transcranial direct current stimulation in children with dystonia: a sham-controlled study. J Child Neurol 2014: 29: 232-239.
- 51. Young SJ, Bertucco M, Sheehan-Stross R, Sanger TD. Cathodal transcranial direct current stimulation in children with dystonia: a pilot open-label trial. J Child Neurol 2013: 28: 1238-1244.
- 52. Bhanpuri NH, Bertucco M, Young SJ, Lee AA, Sanger TD. Multiday transcranial direct current stimulation causes clinically insignificant changes in childhood dystonia: a pilot study. J Child Neurol 2015: 30: 1604-1615.
- 53. Gillick B, Menk J, Mueller B, et al. Synergis-

- tic effect of combined transcranial direct current stimulation/constraint-induced movement therapy in children and young adults with hemiparesis: study protocol. BMC Pediatr: 15: 178.
- 54. Poreisz C, Boros K, Antal A, Paulus W. Safety aspects of transcranial direct current stimulation concerning healthy subjects and patients. Headache 2008: 48: 484.
- 55. Barker AT, Jalinous R, Freeston IL. Non-invasive magnetic stimulation of human motor cortex. The Lancet 1985: 325: 1106-1107.
- 56. Hallett M. Transcranial magnetic stimulation and the human brain. Nature 2000: 406: 147.
- 57. Kirton A, Gunraj C, Chen R. Cortical excitability and interhemispheric inhibition after subcortical pediatric stroke: plastic organization and effects of rTMS. Clin Neurophysiol 2010: 121: 1922-1929.
- 58. Berardelli A, Inghilleri M, Rothwell J, et al. Facilitation of muscle evoked responses after repetitive cortical stimulation in man. Exp Brain Res 1998: 122: 79-84.
- 59. Chen R, Classen J, Gerloff C, et al. Depression of motor cortex excitability by low frequency transcranial magnetic stimulation. Neurology 1997: 48: 1398-1403.
- 60. Kim Y-H, You SH, Ko M-H, et al. Repetitive transcranial magnetic stimulation—induced corticomotor excitability and associated motor skill acquisition in chronic stroke. Stroke 2006: 37: 1471-1476.
- 61. Pal PK, Hanajima R, Gunraj CA, et al. Effect of low-frequency repetitive transcranial magnetic stimulation on interhemispheric inhibition. J Neurophysiol 2005: 94: 1668-1675.
- 62. Mori F, Koch G, Foti C, Bernardi G, Centonze D. The use of repetitive transcranial magnetic stimulation (rTMS) for the treatment of spastici-

- ty. Progress Brain Res 2009: 175: 429-439.
- 63. George MS, Nahas Z, Kozel F, Goldman J, Molloy M, Oliver N. Improvement of depression following transcranial magnetic stimulation. Current Psychiatry Reports 1999: 1: 114-124.
- 64. Koh T, Eyre J. Maturation of corticospinal tracts assessed by electromagnetic stimulation of the motor cortex. Arch Dis Childhood 1988: 63: 1347-1352.
- 65. Concerto C, Lanza G, Cantone M, et al. Repetitive transcranial magnetic stimulation in patients with drug-resistant major depression: a six-month clinical follow-up study. Int J Psychiatr Clin Practice 2015: 19: 252-258.
- 66. Croarkin PE, Wall CA, Lee J. Applications of transcranial magnetic stimulation (TMS) in child and adolescent psychiatry. Taylor & Francis, 2011
- 67. Mineyko A, Kirton A. The black box of perinatal ischemic stroke pathogenesis. J Child Neurol 2011: 26: 1154-1162.
- 68. Mercuri E, Barnett A, Rutherford M, et al. Neonatal cerebral infarction and neuromotor outcome at school age. Pediatrics 2004: 113: 95-100.
- Lisanby SH, Gutman D, Luber B, Schroeder C, Sackeim HA. Sham TMS: intracerebral measurement of the induced electrical field and the induction of motor-evoked potentials. Biol Psychiatr 2001: 49: 460-463.
- 70. Quintana H. Transcranial magnetic stimulation in persons younger than the age of 18. J ECT 2005: 21: 88-95.
- 71. Rossini P, Desiato M, Caramia M. Age-related changes of motor evoked potentials in healthy humans: non-invasive evaluation of central and peripheral motor tracts excitability and conductivity. Brain Res 1992: 593: 14-19.
- 72. Lin K-L, Pascual-Leone A. Transcranial mag-

- netic stimulation and its applications in children. Chang Gung Med J 2002: 25: 424-436.
- 73. Takeuchi N, Chuma T, Matsuo Y, Watanabe I, Ikoma K. Repetitive transcranial magnetic stimulation of contralesional primary motor cortex improves hand function after stroke. Stroke 2005: 36: 2681-2686.
- 74. Naeser MA, Martin PI, Nicholas M, et al. Improved naming after TMS treatments in a chronic, global aphasia patient—case report. Neurocase 2005: 11: 182-193.
- 75. Kauffmann CD, Cheema MA, Miller BE. Slow right perfrontal transcranial magnetic stimulation as a reatment for medication-resistant depression: A double-blind, placebo-controlled study. Depression and Anxiety 2004: 19: 59-62.
- 76. Klein E, Kolsky Y, Puyerovsky M, Koren D, Chistyakov A, Feinsod M. Right prefrontal slow repetitive transcranial magnetic stimulation in schizophrenia: a double-blind sham-controlled pilot study. Biol Psychiatr 1999: 46: 1451-1454.
- 77. Pascual-Leone A, Rubio B, Pallardó F, Catalá MD. Rapid-rate transcranial magnetic stimulation of left dorsolateral prefrontal cortex in drug-resistant depression. The Lancet 1996: 348: 233-237.
- 78. Kirton A, Chen R, Friefeld S, Gunraj C, Pontigon A-M. Contralesional repetitive transcranial magnetic stimulation for chronic hemiparesis in subcortical paediatric stroke: a randomised trial. Lancet Neurol 2008: 7: 507-513.
- 79. Mansur C, Fregni F, Boggio P, et al. A sham stimulation-controlled trial of rTMS of the unaffected hemisphere in stroke patients. Neurology 2005: 64: 1802-1804.
- 80. Hsu W-Y, Cheng C-H, Liao K-K, Lee I-H, Lin Y-Y. Effects of repetitive transcranial magnetic stimulation on motor functions in patients with stroke: a meta-analysis. Stroke 2012: 43: 1849-

1857.

- 81. Valle AC, Dionisio K, Pitskel NB, et al. Low and high frequency repetitive transcranial magnetic stimulation for the treatment of spasticity. Develop Med Child Neurol 2007: 49: 534-538.
- 82. Valero-Cabré A, Oliveri M, Gangitano M, Pascual-Leone A. Modulation of spinal cord excitability by subthreshold repetitive transcranial magnetic stimulation of the primary motor cortex in humans. Neuroreport 2001: 12: 3845-3848.
- 83. Quartarone A, Bagnato S, Rizzo V, et al. Distinct changes in cortical and spinal excitability following high-frequency repetitive TMS to the human motor cortex. Exp Brain Res 2005: 161: 114-124.
- 84. Yasuhara A. Changes in EEG after transcrani-

- al magnetic stimulation in children with cerebral palsy. Functional Neuroscience: Evoked Potentials and Magnetic Fields 1999: 49: 233-238.
- 85. Gilbert DL, Garvey MA, Bansal AS, Lipps T, Zhang J, Wassermann EM. Should transcranial magnetic stimulation research in children be considered minimal risk? Clin Neurophysiol 2004: 115: 1730-1739.
- 86. Miguel Angel C-C, Ignacio M-M, Cordero G CG, et al. Transcranial magnetic stimulation and acoustic trauma or hearing loss in children. Neurol Res 2001: 23: 343-346.
- 87. Garvey MA, Kaczynski KJ, Becker DA, Bartko JJ. Subjective reactions of children to single-pulse transcranial magnetic stimulation. J Child Neurol 2001: 16: 891-894.