

Investigating the Anticonvulsant Effects of Repetitive Transcranial Magnetic Stimulation on Perforant Path Kindling Model in Rats

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Article information	Abstract
<p>Article history: Received: 23 Dec 2012 Accepted: 10 Feb 2013 Available online: 19 May 2013 ZJRMS 2015 Feb; 17(2): 37-41</p> <p>Keywords: Cranial magnetic stimulation Epilepsy Path kindling Dentate gyrus</p> <p>*Corresponding author at: Department of Medical Physics, Faculty of Medical Sciences, Tarbiat Modares University, Tehran, Iran. E-mail: pourmir@modares.ac.ir</p>	<p>Background: Almost 20% of epileptics are drug resistant. Studies have shown that low frequency repetitive transcranial magnetic stimulation (rTMS) is with therapeutic effects on epilepsy-affected laboratory models. Anticonvulsant effects of rTMS depend on several parameters among which radiation frequency is the most important one. In this study, the therapeutic impacts of 1 and 2 Hz rTMS on convulsing parameters in epileptic model of electrical kindling stimulation of the perforant path were investigated.</p> <p>Materials and Methods: In this experimental study 21 rats were randomly divided into three groups, namely '1 Hz treatment group' and '2 Hz treatment group' and 'kindling group'. The kindling group only received kindling stimulations for seven days. One Hz and 2 Hz frequency treatment groups received maximally 5 min rTMS after termination of kindling stimulation per day for a week. Stimulation and stability electrodes had been placed, in turn, on perforant path and dentate gyrus. For quantifying the duration of the subsequent discharge waves, two-way ANOVA test and Bonferroni post-test were employed. In addition, for quantifying the convulsive behaviors, Kruskal-Wallis and the Mann-Whitney <i>U</i> tests were used.</p> <p>Results: The results showed that 1 Hz and 2 Hz frequency rTMS have considerable inhibitory impact on the development of convulsive phases. Anticonvulsive effect was observed from the first day after rTMS was undertaken. In addition, the animals did not show fourth and fifth convulsive stages, and a significant reduction was evident in their recorded peak discharge waves compared with kindle group.</p> <p>Conclusion: Low frequency rTMS possesses significant anticonvulsive effects which depend upon sTMS stimulation frequency.</p> <p>Copyright © 2015 Zahedan University of Medical Sciences. All rights reserved.</p>

Introduction

Epilepsy is a neurological disease from which about 1.5% of global human populations are suffering. This disease is categorized based on various electrographic recurrent attacks. The representation of these attacks lies in the brain and progressively spread from seizure origin to different areas, and even whole brain, electrophysiologically. The varieties of this disease share a single feature: epilepsy increases the cortical excitability of the brain [1]. Temporal lobe is the most prevalent epilepsy [2]. Today, anticonvulsive medications are largely used to tackle epilepsy; however, about 20% of epileptics are drug-resistant [3]. In such cases, surgery is used. Nevertheless, surgery is an invasive method that in several cases is accompanied with irreversible side effects, leading to neuronal damages [4].

The previous studies have shown that low frequency rTMS can have therapeutic effects on epilepsy-affected laboratory models. For example, Akamatsu et al. showed that rTMS of 1000 pulses at 0.5 Hz led to a prolonged latency for seizure development and decreased frequent seizures after intraperitoneal injection of pentylenetetrazol in Wistar rats [5]. Roteaberg et al. investigated antiepileptic effect of rTMS on kainic acid epilepsy model

at 0.75, 0.5, and 0.25 frequencies. They found out that seizure duration decreased at 0.5 and 0.75 Hz frequencies, but 0.25 Hz frequency had no impact on it [6].

These observations demonstrate that anticonvulsive effect of rTMS depends on its frequency. Kindling is a standard model for investigation into the impacts of different therapeutic factors on epilepsy and its resulted seizures [1]. Applying physical parameters such as using 50 Hz magnetic field has shown a weak inhibitory influence on seizure process of kindling [7]. Even though, no report of the rTMS effects, with different frequencies and intensities, on neural excitability parameters, in seizure state, has been published.

Inhibitory and stimulatory effects of rTMS depend on field intensity, duration of the applied pulse train, time interval between each pulse train and applied frequency in particular [7-9]. Due to the huge advantage of this technique (rTMS is a painless and cheap procedure without side effects), obtaining its optimal parameters, especially the applying frequency, can bring beneficial results in considering this method as a supplement or an alternative for conventional epilepsy treatment technique that are with several side effects. In this study, the

therapeutic impacts of 1 and 2 Hz rTMS on convulsing parameters in epileptic model of electrical kindling stimulation of the perforant path were investigated.

Materials and Methods

In this clinical trial, 21 male Wistar rats weighing 270-300 g (from animal house of Tarbiat Modarres University of Medical Sciences) were used. The rats were kept under 12:12 h light-dark-cycle at a constant temperature. The lights were turned on at 7 in the morning. The rats were kept in separate cages with straw-covered floors. They had free access to food and water. After preparation of sterilized surgical instruments, the animal underwent anesthesia using pentobarbital sodium (50 mg/kg, intraperitoneal). The rat, with shaved head, was put into stereotaxy. After the animal's head was fixed and washed with Betadine, a midline incision was made with a surgical blade and the surface of the skull was cleaned with alcohol to identify the bregma point. The position of perforant path on the right hemisphere from the bregma was determined for placing stimulatory electrodes (in millimeter: AP=-6.9, L=4.1, and V=2.4 to 2.7 from the surface of the skull bone). In addition, the position of dentate gyrus from the bregma was determined for placing stability electrodes (in millimeter: AP=-3, L=1.8, and V=3.2 to 3.5 from the surface of the skull bone) [10]. After fastening anchor bolts, uni-polar electrodes, and stimulating and stability electrodes were set in certain places. Using stimulator, an electric excitation, with 50 μ A to 1 mA intensity, was applied on perforant path through stimulating electrode. If the electrodes had been set at appropriate places, following application of stimulation with a single-pulse, the potential of field excitatory post-synaptic potentials (fEPSP) would have been recorded. Otherwise, stimulating and stability electrodes were relocated until an fWSP with maximum amplitude (between 3 to 10 mV) was recorded. In addition, by applying paired-pulse stimulations with 70 ms intrapulse interval and observing paired pulse facilitation phenomenon, as the salient feature of dentate gyrus, the electrode location was confirmed physiologically as well. Then, electrodes and bolts were fixed over the animal's skull using dental cement. For healing the wounds and training the animal, it was given 10 days to rest. Whole research project was in consistent with ethical considerations of keeping and using laboratory animals under the observation of Ethics Committee of Tarbiat Modarres University of Medical Sciences, Tehran, Iran.

Animal Stimulation:

A) Stimulating animals for rapid kindling: Animals were stimulated using rapid kindling technique. First, the subsequent discharge wave threshold for each animal was determined. In this regard, the perforant path was stimulated with a 20 μ A flow. In case the subsequent discharge waves were recorded (at least for 10 seconds), this intensity was taken to be threshold current. Otherwise, in a 5 min interval-cycle, the flow rate was gradually increased for 10 μ A every time to obtain

stimulation threshold. In his procedure, the animals were stimulated for 5 seconds with a single-phase square wave at frequency of 50 Hz, subsequent discharge wave threshold intensity, and 1 ms pulse duration [11]. These stimulations were undertaken twelve times per day in a 5-min interval and continued until the fifth stage of convulsion was observed. Convulsive attack stages were divided into five categories based in Racine pattern (reference number). For generating this stimulating wave, a stimulator (Nihon Kohden, Japan) and a constant-current stimulus isolation unit (SS-202J; Nihon Kohden, Japan) were employed.

B) rTMS stimulations: for investigating the effect of rTMS stimulation frequency, the animals underwent anesthesia with a little amount of CO₂ for 5 min after kindling stimulation, every day. Then, they were fixed in the holder and exposed to magnetic radiation in previously determined spatial coordinates. This spatial coordinates were determined in a way that the maximum field intensity acted on dentate gyrus. All animals were exposed to the magnetic radiation with an intensity equating 80% of motor threshold of each animal. Motor threshold of each animal was behaviorally determined. For keeping the animal immobilized during rTMS stimulation, a holding chamber made of pressured glass (plexiglas) was used. After recovery from surgery, the animals were kept in holding chamber for 15 min per day to lower their stress and get them accustomed to the experimental environment. After 4 to 5 days in holding chamber, animal's head gradually became fixed. After fixing the animal's head, electric coil was located on the motor cortex of the dorsal muscles in a way that the highest field intensity was acting on this region. Low intensities of stimulation pulses applied to the animal, and then contractile responses of dorsal muscles were studied. Field intensity was increased gradually to reach the first contractile response of the animal's dorsal muscle. This output intensity of the device was taken to be 100% of the animal's motor threshold (at rest).

Experimental groups: the animals were divided into two groups, namely 'kindle, and 'kindle+rTMS'. Kindle group received kindling stimulations twelve times per day under 5-min interval. Kindle+rTMS group followed the same protocol, except that they received therapeutic stimulations five min after the last kindling stimulation. A scheme of research plan is presented in figure 1. The parameter investigated in this study included the highest subsequent discharge wave duration per day (maximum ADD; mADD) and highest convulsive phase (maximum Stage of Seizure: mSS).

There were 6 animals in each experimental group. Due to the very low variance of kindling technique, this number of animals was quite adequate. For comparing mADD changes in different groups, two-way ANOVA test and Bonferroni subsequent test were used. The comparison between behavioral phases of seizure was performed using Kruskal-Wallis and Mann-Whitney *U* tests. Significant distance of 0.05 has been set for all statistical tests ($p < 0.05$), unless the contrary has been stated. Prism 5 was employed for statistical analyses.

Results

There was no significant difference between mean subsequent discharge wave duration after first stimulation and stimulation threshold intensity in animals in different groups. That is, the intensity of neural system excitability was equal in different groups. Applying kindling stimulation to kindle group caused animals to reach fifth stage of convulsion after 5.4 ± 0.5 days, in average. The intensity of anticonvulsive effects of different rTMS frequencies in various studies was compared with the data from this group of animals over 7 days.

In order to investigate the impact of rTMS frequency on the intensity of anticonvulsive effects, 1 Hz and 2 Hz rTMS were applied. A sample of subsequent discharge waves of kindle and kindle+rTMS groups are presented in

figure 2. The comparison of mADD was performed using two-way ANOVA test in 7 days between kindle and kindle+rTMS groups, receiving rTMS at different frequencies. The results indicated significant difference in that regard between the mentioned groups ($p < 0.001$). Bonferroni subsequent test showed a significant difference between kindle+rTMS group at 1 Hz and 2 Hz frequencies and kindle group. However, there was no significant difference between 1 and 2 Hz group (Fig. 3).

Comparison between behavioral phases of convulsion revealed a significant difference between groups receiving 1 Hz and 2 Hz rTMS and kindle group, using Kruskal-Wallis test. Mann-Whitney *U* test showed a significant difference between kindle+rTMS and kindle groups from the 2 day. However, there was no significant difference between the two rTMS groups, $p < 0.001$ (Fig. 4).

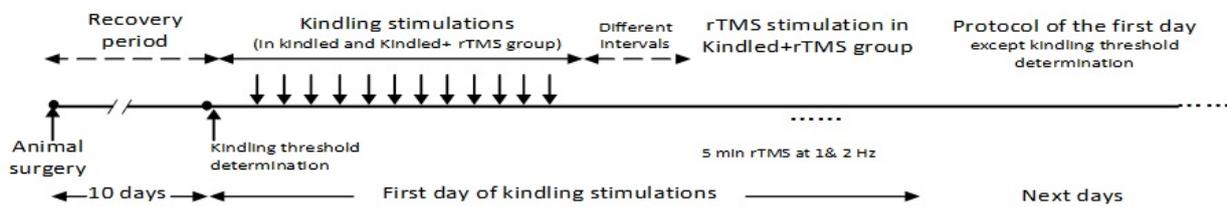


Figure 1. Timing diagram showing the process of kindling stimulation and kindle + rTMS

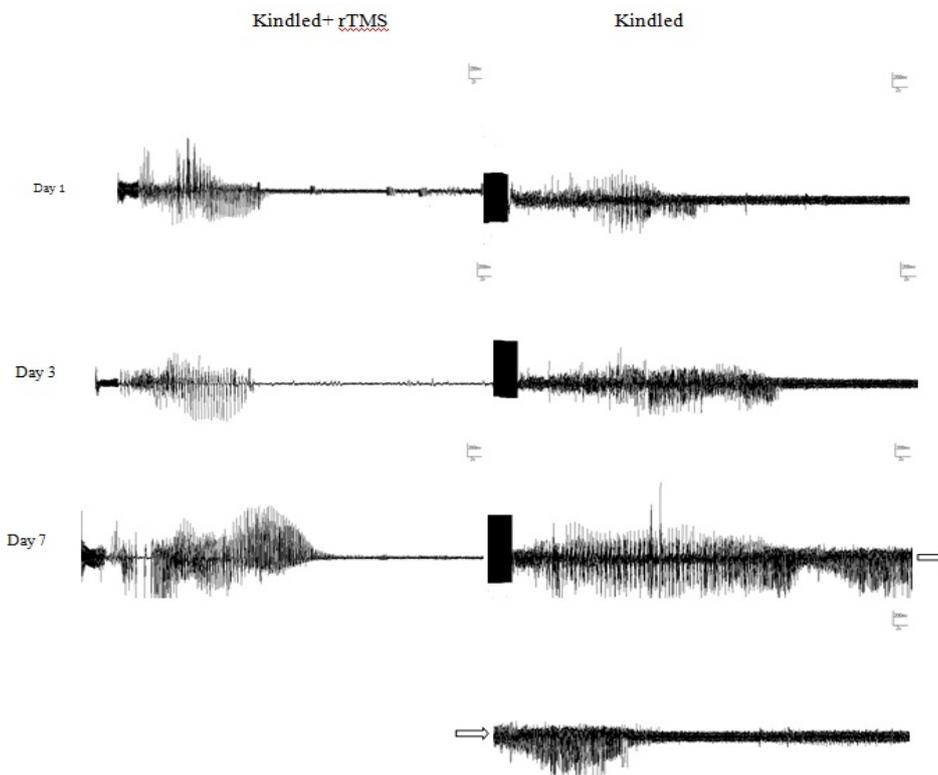


Figure 2. Sample of discharge waves after Kindling group and Kindling + rTMS group. (Kindling stimulation was applied for 2 seconds at the start of registration and scale of all these figures were for 2 seconds for the horizontal axis and 200 mV for the vertical axis that are not clear due to lack of space)

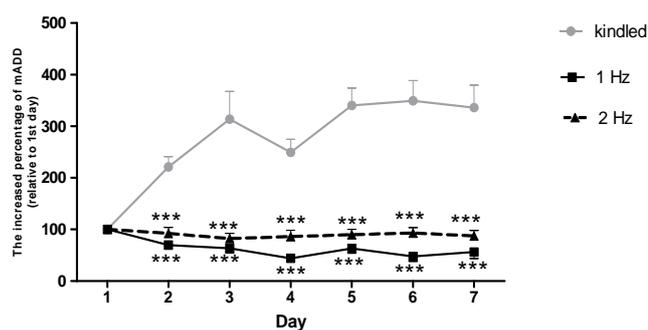


Figure 3. Effect of rTMS with different frequencies during the perforant path kindling. Data are shown as mean±standard error of the mean. * $p < 0.01$ and *** $p < 0.001$ is compared with Kindle group (N = 6).

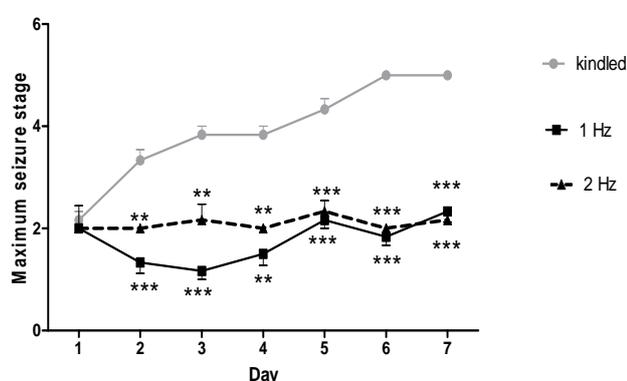


Figure 4. Effect of rTMS with different frequencies during the perforant path kindling on the incidence of behavioral seizures. Data are shown as mean±standard error of the mean. * $p < 0.01$ and *** $p < 0.001$ is compared with Kindling group (N = 6).

Discussion

The findings of this study showed that applying rTMS at 1 and 2 Hz frequencies has remarkable impact on inhabitation of convulsive stages. In that, this anticonvulsive effect was observable from the first day after applying rTMS. In addition, those animals that had received rTMS did not show fourth and fifth convulsive stages, and the highest discharge wave observed in them was with a considerable reduction compared with kindle group. This study was in consistent with previous ones. For example, Akamatsu et al. demonstrated that applying 1000 pulse of rTMS at 0.5 Hz frequency caused mitigation of epileptic seizures in successive epilepsy model, after injection of pentylenetetrazol to peritoneum of epileptic rats [5]. Rotenberg et al., revealed that rTMS at 0.5 and 0.75 Hz has anticonvulsive effects on rats with kainic acid-induced epilepsy [6]. Ke et al. and Huang et al. studied the frequency-dependent effect of rTMS in pilocarpine epileptic model. They applied rTMS to the rats at different frequencies for two weeks. Then, pilocarpine was injected and it was observed that 1, 0.8, 0.5, and 0.3 Hz frequencies postponed the onset of seizure [12, 13]. It has been demonstrated that low frequency stimulations generate antiepileptic effect through

decreasing NMDA receptor and subsequent reduction of neuron excitability [14, 15].

In a preliminary study by Anschel et al., it was demonstrated that 1 Hz rTMS has anticonvulsive effect [16]. In the mentioned experiment, the injection of cerebrospinal fluid of depressed patients, exposed to 1 Hz rTMS, to the ventricle of the rats decreased kindling rate through increasing seizure threshold. rTMS was stimulated for eight daily sessions of 26.6 min. The intensity of stimulation was equal to 90% of individual's motor threshold. The same protocol was employed for 10 Hz rTMS. A total of twenty pulse trains, each for 8 seconds with 22 seconds interval and total radiation duration of 22 min were applied. The results showed that in contrast to 1 Hz frequency, 10 Hz frequency did not reduce seizure threshold. These findings conform to the results from the present study in that 1 Hz rTMS inhibits and suppresses neuronal activity. However, duration of stimulation and also the type of protocol (injection of cerebrospinal fluid of radiated sample to the studied sample) were different from the present study. Regarding the above, it could be argued that rTMS radiation causes secretion and increased concentration of endogenous anticonvulsive substances in cerebrospinal fluid. These cases possess anticonvulsive effects and increase in their concentration can last for several minutes after rTMS radiation [16]. It has been suggested that the time-varying magnetic field at some frequencies and intensities cause release of melatonin, an anti-epileptic compound, in the brain tissue [17, 18].

There are other theories regarding the mechanism of this technique. Modulating the voltage-dependent sodium ion channels [19, 20] and the release of neurotransmitters at interneuron in cortical and cerebrospinal regions [21, 22] are among the mechanisms of the drugs, with anticonvulsive effects on rTMS variables suggested for the patients with different types of epilepsy and normal samples. However, due to the fact that all generated physical and physiological effects of this technique on brain tissue and intranetworking impacts of different points of brain on each other are not quite determined, then the mechanism produced by this technique in the brain is not exactly obvious.

In order to establish this promising technique as an alternative or supplementary option for curing treatment-resistant epilepsy, on the one hand, the exact simulation of physical parameters of generated electric field in the brain tissue and performing experiments at molecular level can be helpful in clarifying these mechanisms. On the other hand, further studies to investigate the effects of other frequencies, to obtain optimal frequencies with therapeutic impacts, and to study the effects of other parameters of this technique such as intensity of magnetic stimulation radiation, spatial shape of radiation coils, duration of magnetic stimulation, and spatial distance and direction of radiation coil can help determine the optimal radiation conditions for curing treatment-resistant epilepsies. Due to the advantages of this technique,

obtaining optimal parameters for treating epileptics can be a step towards its clinically popularization.

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Authors' Contributions

All authors had equal role in design, work, statistical analysis and manuscript writing.

Conflict of Interest

The authors declare no conflict of interest.

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