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Transcranial direct current stimulation efficacy in trigeminal neuralgia

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Abstract

Trigeminal neuralgia is a severe, disabling pain and its deafferentation remains a

challenge for health providers. Transcranial direct current stimulation is a non-invasive

stimulation technique which finds new utility in managing pain. Therefore, the

introduction of alternative, non-invasive, safe, and effective methods should be

considered in treating patients with trigeminal neuralgia unresponsive to conventional

treatment.

TO THE EDITOR

We read with interest a case report by Tao JC et al, who presented one patient with

herpetic neuralgia after herpes zoster ophthalmicus who was treated with patient-

controlled intravenous analgesia (PCIA) with esketamine [1]. The procedure resulted in a

significant pain relief without adverse reactions.

We support authors' statement that primary trigeminal neuralgia can achieve

satisfactory curative effects through medical treatment and interventions like

radiofrequency. However, people who suffer from side effects to oral medication or

more complicated cases (such as neuralgia caused by varicella-zoster virus infection of

the trigeminal nerve) are not satisfactorily managed and require more aggressive

surgical treatments. Deep brain stimulation and motor cortex stimulation are off label,

"last treatment option" techniques which may offer relief to trigeminal neuralgia that is otherwise refractory to pharmacological management and surgery^[2, 3]. Nevertheless, (PCIA) with esketamine could be an alternative, non-invasive, safe, and effective method.

We would like to suggest transcranial direct current stimulation (tDCS) as an additional potential effective method which could offer relief in intractable trigeminal neuralgia. A 51-year-old woman with idiopathic trigeminal neuralgia, was hospitalized towards modifying the therapeutic strategy and ameliorating the adverse events of medical treatment. She presented daily paroxysmal attacks with a mean duration of 1,5-2 h, reoccurring patterns of few minutes (up to 3-4 minutes) that created intense, needle-like pain on the dental, submental, and periocular areas of the right side of the face. Refractory period was not established, and she also reported night awakening with instant feeling of intense pain. She scored 9 in the Visual Analogue Scale (VAS)^[4]. Her medical history was unremarkable. Physical examination, imaging and laboratory investigations were normal. Peros medication (including carbamazepine) proved to be ineffective and was not tolerated.

Relying on current literature, we decided to apply tDCS^[5, 6]. The placement of the electrodes followed the Electroencephalography System 10-20 with the anode (+) occurring at the motor cortex and the cathode (-) at contralateral motor area-cortex and at the contralateral mastoid bone. The Sooma™ tDCS device was used, with the intensity settings for the active tDCS being at 2mA, lasting 30 minutes per session, with 30 s ramp up and down, for a total of 10 days (2 wk). The electrical conduction took place through 2 electrodes 5 X 5 cm (contact surface 25 cm²). No severe adverse events were reported. Anodal tDCS for 10 days ameliorated the intensity of pain (the VAS score reduced from 9 to 2). During the post-stimulation period, the sleep and the quality of life were improved, as they were reflected in Short-Form (36) Health Survey (SF-36), Athens Insomnia Scale (AIS) (the SF-36 and the AIS scores changed from 32 to 60 and from 17 to 5, respectively)^[Z]. The patient displayed sustained efficacy with this management for 3 months.

A number of studies have demonstrated that anodal tDCS modulates the cortical excitability of the motor cortex. However, the effects of tDCS on cortical excitability can spread to distant cortical areas. The main mechanism of action of tDCS is the modulation of the membrane potential of neurons in the stimulated cortical area, mediated by N-methyl-D-aspartate receptors (NMDA-R). Nonetheless, it has been shown that the modulation of nicotinic receptors, BDNF polymorphisms and sex hormonal variations also affect brain plasticity^[9].

The results described may provide additional evidence for the effectiveness of tDCS as a therapeutic instrument in treating patients with trigeminal neuralgia unresponsive to conventional treatment. The three-month efficacy reported is positive, however, studies with longer-term follow-up data to assess the sustainability of the tDCS intervention should be scheduled.

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