Pressure and cold pain threshold in healthy subjects undergoing interferential current at different amplitude modulated frequencies*

Avaliação do limiar de dor à pressão e ao frio em indivíduos saudáveis submetidos à corrente interferencial em diferentes frequências de amplitude modulada

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ABSTRACT

BACKGROUND AND OBJECTIVES: The interferential current is an electrical stimulation method widely used with analgesic purposes; however, there are controversies regarding their usage parameters. The aim of this study was to compare different amplitude modulated frequencies of the interferential current on the pressure and cold pain threshold in healthy volunteers.

METHODS: This is a crossover, randomized and blinded evaluators and evaluated in quantitative character test. Participants were 20 volunteers divided into four groups, who for four weeks were subject to different modulated frequencies. Thresholds of pressure and cold pain in four different times were evaluated - at the beginning, right after electric stimulation, 20 and 60 minutes after. As a form of electrical stimulation, interferential current was used at modulated amplitude frequencies of 1Hz, 10Hz and 100Hz with base frequency of 4,000Hz, being that in a particular week volunteers received placebo (0Hz).

RESULTS: There has been no significant difference in comparisons within frequencies or between them and the placebo group. **CONCLUSION:** The use of amplitude modulated frequencies of interferential current of 1Hz, 10Hz and 100Hz did not change cold and pressure pain thresholds, similar to what has been found for the placebo group

Keywords: Analgesia, Pain measurement, Transcutaneous electric nerve stimulation.

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RESUMO

JUSTIFICATIVA E OBJETIVOS: A corrente interferencial é um método de eletroestimulação bastante utilizado com finalidades analgésicas, porém, existem controvérsias com relação aos seus parâmetros de uso. O objetivo do estudo foi comparar diferentes frequências de amplitude modulada, da corrente interferencial, sobre o limiar de dor à pressão e ao frio, em voluntários saudáveis.

MÉTODOS: Trata-se de um ensaio cruzado, aleatório e com avaliadores e avaliados encobertos, de caráter quantitativo. Participaram deste estudo 20 voluntários subdivididos em quatro grupos, que durante quatro semanas passaram por diferentes frequências de amplitude modulada. Foram avaliados os limiares da dor à pressão e ao frio em quatro momentos distintos – no início, logo após eletroestimulação, 20 e 60 minutos após. Como forma de eletroestimulação foi utilizada a corrente interferencial nas frequências de amplitude modulada de 1Hz, 10Hz e 100Hz, com frequência de base de 4.000Hz, sendo que em uma determinada semana os voluntários recebiam placebo (0Hz).

RESULTADOS: Não houve diferença significativa nas comparações dentro das frequências, nem entre elas e o grupo placebo. **CONCLUSÃO**: O uso da corrente interferencial nas frequências de amplitude modulada de 1Hz, 10Hz e 100Hz não produziu alterações nos limiares de dor ao frio e à pressão, semelhante ao encontrado para o grupo placebo.

Descritores: Analgesia, Estimulação elétrica nervosa transcutânea, Mensuração da dor.

INTRODUCTION

The International Society for the Study of Pain (IASP) defines pain as "a sensory, emotional and unpleasant experience, related to tissue injury or described in such terms". It is a manifestation involving physical, psychic and cultural mechanisms, should be considered the fifth vital sign and as such should be measured and recorded¹. So, methods are used to quantify pain intensity, among them pressure algometry, which is a power transducer requiring perceptive response of the evaluated patient, being very useful in the evaluation of clinical trials².

Other experimental pain models are used to represent pain, such as cold-induced pain, which is a simple method involving minimum risk of tissue injury and pain stops after removal of the stimulus. During such test, a painful sensation is generated by temperature receptors which start sending stimuli of a possible tissue injury through peripheral and central pathways, resulting in cold-induced pain sensation³.

Since pain is still the primary reason for looking for medical aid, analgesic techniques are developed and studied, such as the use of interferential current. This is commonly used as electric stimulation, the primary clinical use of which is to relieve pain^{4,5}. This current has effect on decreasing pain intensity by significantly increasing pain threshold and may even block nervous conduction⁶⁻⁸. However, there are controversies about interferential current effectiveness with some studies showing positive results⁹ and others negative results¹⁰, which is often caused by different pain evaluation methods or by differences in stimulation parameters.

So, there is need for studies evaluating current efficacy on different pain threshold types, as well as the diversity of therapeutic parameters commonly used for interferential current, such as, whether low amplitude modulated frequencies (AMF) have any kind of influence on analgesic mechanisms. So, this study aimed at comparing the effect of different interferential current AMFs on pressure and cold pain threshold, determining the effectiveness of the electric current to promote analgesia by increasing pain threshold in healthy individuals.

METHODS

This is a crossover, randomized and quantitative trial with blind evaluating and evaluated individuals. Participated in the study 20 volunteers, being 13 females and 7 males, with mean age of 20.35±1.725 years, weight 63.95±12.03 kg, height 1.695±0.1073 m and body mass index (BMI) of 22.13±2.714 kg/m². Volunteers have signed the Free and Informed Consent Term (FICT), where they agreed in participating in the study without any compensation and were informed that they could quit the study at any time by free will.

Individuals with any contraindication to the use of electric stimulation, in addition to contraindications to cold and heat were excluded.

Volunteers were randomly divided in four groups of 5 individuals each: group 1 (G1), group 2 (G2), group 3 (G3) and group 4 (G4). Individuals have always performed the same activities within the group, in four consecutive weeks and in crossover manner, that is, each group has gone through all different types of frequencies in different weeks. This way, all volunteers were submitted to all types of electric stimulation and placebo.

Evaluation protocol

Different transcutaneous AMFs were evaluated: 0Hz, 1Hz, 10Hz and 100Hz, with bipolar technique and base frequency of 4000 Hz, on pressure and cold pain thresholds. For such, volunteers have participated in an initial evaluation (EV1),

before being undergoing electric stimulation.

Previously to initial evaluation, volunteers have immersed their left arm until the elbow joint interline, in a container with water at 38° C (Inconterm® thermometer) for 5 minutes, aiming at homogenizing sample temperature. Soon after arm drying, pressure pain threshold was evaluated (Algometer – Kratos® with capacity to produce up to 50 Kgf, with sharp metal tip). It was explained that they should report the exact moment when they felt pain. Sites to apply the algometer were: 1cm distal to pisiform bone in hypothenar region and 1 cm distal to scaphoid bone in thenar region. Pressure was gradually increased and removed as soon as the volunteer reported pain.

Cold-induced pain was evaluated three minutes after pressure pain threshold evaluation. Volunteers immersed the left hand in a container with water and ice (5° C) until wrist joint interline. Individuals were asked to report pain sensation as soon as they felt it, time was measured as from the moment volunteers immersed their hand in cold water and as soon as they said the word "pain", this time was recorded as cold pain threshold.

Such evaluations were repeated immediately after using interferential current (EV2), 20 (EV3) and 60 (EV4) minutes after. In all evaluation moments, the evaluator did not know with AMF frequency the volunteer would receive or had received.

Electric stimulation protocol

Rubber-silicone 8 cm² electrodes were used, coupled to patients with water-soluble gel and fixed with adhesive tape; equipment used was Neurovector (Ibramed®). Since this was a crossover study, volunteers were initially distributed in one of 4 groups. In the first week, they went through a certain frequency, in the following week they were submitted to a different frequency, and so on, until the end of the 4 proposed weeks:

- 0Hz placebo moment when no current was emitted to volunteer;
- 1Hz equipment produced 1Hz AMF;
- 10Hz 10Hz AMF;
- 100Hz -100Hz AMF.

Electric stimulation was performed for 20 minutes, with electrodes located on elbow joint, being one positioned on the ulnar nerve sulcus (ulnar nerve superficiality) and the other on the cubital face (median nerve superficiality), with intensity considered high, however not painful. Asepsis was performed with cotton soaked in 70% alcohol on application site, previous to electrodes placement.

Statistical analysis

Data normality was analyzed with Kolmogorov-Smirnov test and comparisons were analyzed by ANOVA for repeated measures with significance level of 5%.

The study was approved by the Ethics and Research Committee, State University of Western Paraná 143/2013.

Table 1. Values for algometer pressure in grams (g), for different groups

	0Hz	1Hz	10Hz	100Hz
AV1	0.776±0.412	0.835±0.425	0.777±0.458	0.813±0.455
AV2	0.764±0.368	0.814±0.349	0.695±0.353	0.836±0.459
AV3	0.767±0.357	0.780±0.380	0.760±0.448	0.747±0.401
AV4	0.738±0.382	0.747±0.351	0.755±0.397	0.779±0.423

(0Hz: placebo, 1Hz, 10Hz and 100Hz), in different evaluation moments (AV1: pre-stimulation; AV2: immediately after stimulation; AV3: 20 minutes after; AV4: 60 minutes after); with pressure on hypothenar eminence.

Table 2. Values for algometer pressure in grams (g), for different groups

	0 Hz	1 Hz	10 Hz	100 Hz
AV1	0.599±0.293	0.638±0.300	0.614±0.283	0.782±0.630
AV2	0.654±0.319	0.658±0.310	0.561±0.269	0.681±0.338
AV3	0.718±0.363	0.713±0.355	0.641±0.263	0.649±0.323
AV4	0.626±0.362	0.654±0.352	0.684±0.307	0.670±0.319

(0Hz: placebo, 1Hz, 10Hz and 100Hz), in different evaluation moments (AV1: pre-stimulation; AV2: immediately after stimulation; AV3: 20 minutes after; AV4: 60 minutes after); with pressure on thenar eminence.

Table 3. Values for cold pain threshold in seconds (s), for different groups

	0 Hz	1 Hz	10 Hz	100 Hz
AV1	31.45±20.87	43.95±37.49	28.90±31.33	28.35±21.42
AV2	46.10±72.65	35.40±22.76	28.50±30.93	24.00±18.57
AV3	34.75±25.99	40.15±23.67	33.25±32.68	33.75±51.35
AV4	35.15±23.38	39.40±26.61	34.55±50.83	43.10±63.42

(0Hz: placebo, 1Hz, 10Hz and 100Hz), in different evaluation moments (AV1: pre-stimulation; AV2: immediately after stimulation; AV3: 20 minutes after; AV4: 60 minutes after).

RESULTS

There has been no significant difference in pressure pain threshold among different evaluations and also among different frequencies, both for thenar region pressure (p=0.2714) (Table 1) and for hypothenar region (p=0.7645) (Table 2). Similarly, there has been no significant difference for cold pain threshold (p=0.5687) (Table 3).

DISCUSSION

This study looked at comparing different interferential current AMFs on pressure and cold pain threshold, with strong, however pleasant stimulation. There were no significant differences in pressure and cold pain thresholds, regardless of the evaluation moment. This confirms previous study where authors have reported no significant changes in A delta and C fibers threshold when interferential current was applied to healthy individuals¹¹. Similarly, a different study inducing cold pain in healthy individuals has not observed significant effect on pain threshold of individuals submitted to interferential current¹². It is believed that this lack of positive effects (analgesia) is due exactly to the fact of not stimulating with painful current, that is, stimulation was not painful and this way there has been no inhibitory effect by descending pathways.

Notwithstanding studies showing poor results of interferen-

tial current to control experimental pain, there are hints that it works as complementary therapy for pain relief, being more effective than placebo. However, since studies substantiating this are widely different, such hints should be taken with care⁴. A study using interferential current with painful and non-painful stimulations in patients with chronic pain due to knee osteoarthritis, has observed that interferential current was effective for analgesia, but high intensity (painful) stimulation has produced more hypoalgesic effects as compared to sensory stimulation⁶.

A study evaluating the effect of interferential current on pressure and cold pain models, has observed hypoalgesia by electric stimulation, differently from what has been observed in our study, and authors mention that pain thresholds evaluation is enough for the observation of the effects of the current, with no need for higher intensities, which generate more discomfort and risks for injuries? This way, in our study, although with different results from those already mentioned, we decided for evaluating just pain thresholds.

In healthy individuals, stimulation below pain threshold in cases of electric stimulation may be the differential between the production or not of hypoalgesic effects by using conventional TENS² and also its burst form¹³. A study evaluating Bernard diadynamics in healthy individuals with non-painful intensities, also points to the lack of hypoalgesic effects of electric stimulation¹⁴. That is, with non-painful intensities,

accepted as major stimuli for ascending analgesia, via gate, non-noxious stimulation was unable to produce pain threshold changes. However, authors show evidences that 80 Hz TENS increases pressure algometry threshold in healthy participants when a strong, however painless sensation is produced in the experimental pain site¹⁵.

In our study, we decided for two experimental ways to evaluate pain: pressure and cold. These ways have some advantages, such as better control of stimuli and conditions, in addition to helping the recording, which is important; however emotional aspects of individuals with clinical pain are not addressed. So, we emphasized that the study had as limitation the fact that participants had no pain and also that the intensity used has not reached pain threshold. So it is suggested that further studies are carried out with individuals with chronic pain and/or using higher current intensities.

CONCLUSION

Interferential current at 1Hz, 10Hz and 100Hz AMFs has not produced cold and pressure pain threshold changes, similarly to what was found for the placebo group.

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