DIODE LASER IRRADIATION COMBINED WITH PROPOLIS APPLICATION AS A TREATMENT FOR DENTINE HYPERSENSITIVITY

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ABSTRACT

INTRODUCTION: Dentine hypersensitivity (DH) is an extensive clinical condition, which is still of need to find a long-term treatment resulting in pain reduction.

OBJECTIVES: The aim of this study was to evaluate and compare efficacy of sodium fluoride varnish application, diode laser (810 nm) irradiation, ethanolic extract of propolis application, and combined diode laser (810 nm) irradiation with ethanolic extract of propolis application for DH reduction in periodontal maintenance patients.

MATERIAL AND METHODS: In total, 104 periodontal maintenance patients (416 teeth) completed this study. Dentine hypersensitivity was assessed using visual analogue scale (VAS). The patients were randomly divided into four groups according to the treatment: 1 group – diode laser (810 nm) irradiation, 2 – application of ethanolic extract of propolis, 3 – application of ethanolic extract of propolis, followed by diode laser (810 nm) irradiation, and 4th group – sodium fluoride varnish application. The treated teeth were evaluated after treatment session and at one week, one month, three months, and six months post-operatively. The evaluations were analyzed using IBM SPSS Statistics, version 25.0 (IBM Corp., Armonk, NY, USA). Mean and standard deviation were calculated for each group. Wilcoxon’s rank sum test and Friedman’s test were used, and values of p < 0.05 were considered as statistically significant.

RESULTS: All four groups showed a significant reduction in the VAS scores immediately after treatment. The most long-term results were obtained in group with combined treatment.

CONCLUSIONS: The present study demonstrates that better result has been noted in group with combined application of ethanolic extract of propolis with diode laser (810 nm) irradiation.

KEY WORDS: propolis, LLLT, lasers, dentine sensitivity.

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INTRODUCTION

Dentine hypersensitivity (DH) is a common clinical finding, which has been defined as “a short, sharp pain arising from exposed dentine in response to stimuli typically thermal, evaporative, tactile, osmotic, or chemical, and which cannot be ascribed to any other form of dental pathology” [1, 2]. Depending on the study, DH is a pervasive condition with a prevalence ranging from 3% to 57% of the general population. It is estimated to affect 15% of adult population, mostly within 30 and 40 years of age. It is known that women are affected more than men [3, 4].
Etiology of DH includes multiple factors. Various aspects, such as recession of gingiva, enamel loss, and tooth wear as well as patients’ habits could be associated with DH. Moreover, frequent etiological factors are related to periodontal disease and consequences of its treatment. Chronic periodontitis is characterized by formation of periodontal pockets and alveolar bone resorption. As known in physiological conditions, enamel normally covers the dentine in the crown and cementum covers it in the root, which makes the tooth insensitive to direct stimulation. Due to chronic periodontitis and its treatment, such as scaling and root planning, the dentine might be uncovered. Consequently, DH occurs with an uncovered exposure of diameters of dentinal tubules [5,6].

The mechanism of DH-related pain is described by several theories. Most widely acknowledged hypothesis is the so-called ‘hydrodynamic theory’. In the beginning of the twentieth century, Alfred Gysi, without any scientific evidence, determined that there is an outward fluid flow inside dentinal tubules. Gysi assumed that the application of some appropriate stimuli on the dentine surface could extend the changes of fluid movement and activate the pulpal nerves. In 1963, Martin Brännström has published a lot of studies supporting this theory. The studies were performed on both human and animal models. This concept is based on an evidence that the fluid flow in dentinal tubules is stimulus-induced. Thermal, tactile, or chemical stimuli change the movement of fluid and induce deformation of nerve endings in pulp, causing pain [5-7].

There are various methods of DH treatment. Several of them are based on a decrease of fluid movement due to dentinal tubules sealing or nerve activity blockage. According to publications from recent years, no completely efficient treatment for DH has been found yet [3,4,8]. Propolis, known as the bee-glue, is a natural resinous substance collected from plant buds or exudates by bees. Most often, propolis contains a mixture of resin, essential oils, wax, amino acids, minerals, ethanol, complex of A, E, B vitamins, zinc, pollen, and bioflavonoids also known as the highly active biochemical substance. Biological activity of propolis is mainly associated with well-known plant compounds, such as bioflavonoids. Propolis is widely used not only in medicine, but also in contemporary dentistry because of its anti-inflammatory, anti-oxidant, anti-bacterial, anti-viral, and anti-fungal properties. According to literature review, propolis has been used for DH treatment and showed a significant reduction of pain [9-11].

Many papers have been published on wide usage of DH laser-based treatment. Variety of types of lasers are recommended for effective DH treatment, of which is a low-level laser therapy (LLLT). In recent studies, the LLLT was suggested as an effective treatment. Low-level laser irradiation increases the metabolic activity of pulp cells, resulting in the production of tertiary dentine [12-14].

In recent years, the most common desensitizer agent was sodium fluoride. A range of desensitizers based on sodium fluoride are recommended for an efficient reduction of pain. The mechanism of action is explained by occluding dentine tubules with calcium fluoride crystals. Moreover, obliteration of dentinal tubules by calcium fluoride crystals reduce the fluid movement. Due to a hydrodynamic theory, this may help to decrease the level of pain [1,4].

**OBJECTIVES**

The aim of this study was to compare efficacy of sodium fluoride varnish application, diode laser (810 nm) irradiation, ethanolic extract of propolis application, and combined diode laser (810 nm) irradiation with ethanolic extract of propolis application for DH reduction in periodontal maintenance patients at long-term follow-up.

**MATERIAL AND METHODS**

One hundred and twenty patients with periodontal diseases affected by DH were included into the present study. Patients’ recruitment was performed within the Department of Therapeutic Dentistry, Faculty of Dentistry, Kharkiv National Medical University. Inclusion criterion was DH of teeth in patients with chronic periodontal disease. Exclusion criteria were carries lesions or restorations, any professional DH therapy during the last 6 months, usage of desensitizing tooth-paste within the last 3 months, and pregnancy.

Firstly, patients were provided with oral and written information about the intention and design of the study. Secondly, patients were included in our study only after signing an informed consent form. The ethical and bioethical committee of the Kharkiv National Medical University (minutes No. 6 of 4 October 2017) approved the study protocol and related consent forms.

The vitality of all experimental teeth was examined at the beginning and the end of the study by an electric pulp tester (Averon, EOT 1.1, Russia) to exclude pulp pathology. The degree of sensitivity was determined by an examiner using visual analogue scale (VAS). Each tooth response was measured by thermal evaporative stimulus as air blast. A cold air blast was given with an air syringe for 1-2 seconds at approximately 1 cm at right angle to the buccal site from the surface of tooth. Adjacent teeth were isolated with cotton rolls to prevent false positive results. Air stimulus time was controlled by chronometer, and the distance was measured by a periodontal pocket probe (Surgicon Pvt. Ltd., Pakistan). All patients were requested to record their level of dentine hypersensitivity using VAS scale ranging from 0 to 10, where 0 was “no pain” and 10 “unbearable pain”. A separate sheet of paper with pre-printed VAS scale was provided to patients for their scores. For every assessment of DH level, the patients filled in a new line of VAS to prevent any biased evaluation, which could be influenced by previ-
The patients were provided with instructions, such as monthly, three months, and six months post-operatively. (for immediate assessment of results), at one week, one month, three months, and six months post-operatively. The included teeth were evaluated after the procedure ending of our study. For the evaluation of results, only one procedure per patient in all groups at the beginning of our study. For the evaluation of results, only one tooth with DH was randomly chosen in every quadrant (four teeth for one patient). Values were collected before the treatment (baseline values), and the included teeth were evaluated after the procedure (for immediate assessment of results), at one week, one month, three months, and six months post-operatively. The patients were provided with instructions, such as not to rinse, eat, or drink for 30 minutes after the treatment, and to avoid using any other desensitizing agent during the study. All data were collected using Microsoft Excel sheet. Statistical analysis was performed by IBM SPSS Statistics for Windows, version 25.0 (IBM Corp., Armonk, NY, USA). Firstly, mean and standard deviation were calculated for each group. Since the data were not distributed normally, non-parametric tests were selected. Wilcoxon's rank sum test was used to evaluate the differences within groups at each time point. Intragroup time-dependent data were analyzed by Friedman’s test. Values of p < 0.05 were considered as statistically significant.

RESULTS

Of 120 patients included in our study, 104 completed the 6-month study period. There were 34 males and 70 females, with a mean age of 40.91 ± 9.03 years. Mean age and standard deviation in all groups are presented in Table 1. Due to reduced number of patients who completed the full study, the number of patients in each group became 26. The total number of teeth included in our study was 416. Table 2 presents teeth distribution.

Complications such as adverse pulp effects or allergic reactions were not observed throughout the study. Responses of the patients to thermal evaporative stimulus by visual analogue scale (VAS) throughout the study and the effects of treatments in the four groups at different time points are shown in Figure 1.

| Table 1. Mean and standard deviation of patients’ age |
|-----------------------------------------------------|
| **Group** | **Age** |
| 1 | 40.38 ± 9.37 |
| 2 | 40.81 ± 9.24 |
| 3 | 41.04 ± 9.04 |
| 4 | 41.42 ± 8.98 |

| Table 2. Distribution of teeth included in study |
|-------------------------------------------------|
| **Teeth** | **Number of teeth** |
| Maxillary central incisors | 25 |
| Maxillary lateral incisors | 24 |
| Maxillary canine | 51 |
| Maxillary premolars | 81 |
| Maxillary molars | 27 |
| Mandibular central incisors | 26 |
| Mandibular lateral incisors | 26 |
| Mandibular canine | 55 |
| Mandibular premolars | 76 |
| Mandibular molars | 25 |
Mean VAS scores and standard deviations at different times of the study in the four groups are demonstrated in Table 3. Based on intragroup comparisons, the treatment of DH was significantly effective in all four groups (Table 3). Intragroup comparisons showed that differences between baseline scores and immediate, 1 week, 1, 3, and 6 months after the treatment were statistically significant ($p < 0.05$). In all four groups, immediate responses for thermal evaporative stimulus showed a significant reduction of level of pain. Responses in group 3 revealed significant differences among the scores of baselines, immediately after the treatment, and at 1 week, 1, 3, and 6 months post-operatively (Table 3).

Paired comparisons showed significant differences between the treatment groups, with $p < 0.05$. No significant differences between the treatment groups were found at baseline ($p > 0.05$). Additionally, no significant differences between group 1 and 4, and group 3 and 4 for thermal evaporative stimulus immediately after the treatment were observed.

Comparison between groups showed that the lowest VAS scores for thermal evaporative stimulus were obtained in group 3 during the study (Figure 1).

### DISCUSSION

DH is a common condition for patients with periodontal disease, because of exposed root surfaces and gingival recession. However, with multiple variants of treatments, there is still no gold standard for reducing DH [4, 8]. The present clinical study compares the efficacy of sodium fluoride varnish, ethanolic extract of propolis application, diode laser (810 nm) irradiation, and the combination of two methods, such as application of ethanolic extract of propolis and diode laser (810 nm) irradiation in decreasing the symptoms of DH.

Patient’s pain and discomfort caused by dentine hypersensitivity is individual and highly subjective. In most studies, scales are used for assessment of DH. The most frequent are visual analogue scale (VAS) and Schiff sensitivity scale [4]. According to literature review, scientists evaluated DH with the use of thermal evaporative stimuli, such as cold air blast provided with an air syringe. Findings of these studies demonstrated that evaporative stimuli dehydrated the dentine surface, which increased the movement of fluid in dentine tubules [6].

In our study, sodium fluoride varnish was used, because sodium fluoride is the most frequent agent of desensitizers and we sought to compare its efficiency with usefulness of our design of DH treatment. The results of our study are in accordance with other researches and confirm that application of sodium fluoride desensitizer shows immediate result in the reduction of DH-related pain, however, with a short-term effect [4, 12]. The 6 months VAS scores did not return to the baseline values but were significantly higher when compared to a combined treatment.

Here, ethanolic extract of propolis was chosen as a natural desensitizing agent and because of the occurrence in many studies, in which showed a high efficacy in DH treatment [15-19]. In our study, the immediate VAS scores were significantly lower than the baseline values. However, during the period of study, the meaning of VAS scores increased. Studies on propolis comparing other desensitizers provided significant results in reducing DH [2, 20]. In the present study, a significant difference between the baseline and 6-month VAS scores was also obtained.

In recent times, laser irradiation was proven efficient for DH treatment [21-24]. In our study, diode laser was chosen as an LLIT appliance, and it was successfully implemented. Many studies reported the same results. However, most of them described a higher decrease in the 6-month VAS scores, which was not obtained.
in our study. It could be explained that all these studies were performed using different diode lasers, with a wide range of wavelengths of 635-830 nm and dosages of 2-10 J/cm². The usage of all these parameters is acceptable and does not affect the morphology of enamel or dentine surface [12-14, 21-24].

According to literature review, in this study, we wanted to combine all the impacts of application of propolis ethanolic extract with diode (810 nm) laser irradiation for DH treatment. We considered that this design of DH treatment can improve and prolong the outcomes in decreasing the level of pain.

The mechanism of action of both methods are not completely clear. For example, scientists observed that the dentine binds to propolis flavonoids and forms crystals with an ability to adhere to the dentine surface. As a result, the dentine tubules are occluded. On the other hand, propolis consists of various natural resinous substances. It is confirmed that propolis could be resemble dental adhesive substances as varnishes or composite resins. The dentine tubules are occluded because mechanical interlocking could prevent fluid flow. Moreover, propolis may deeply diffuse inside dentinal tubules and be difficult to remove, which could sufficiently prolong the result of treatment [15-17]. However, the mechanism of action of propolis requires more studies for clarity. Other studies showed that the production of transforming growth factor (TGF)-β1 is stimulated by propolis. Moreover, TGF-β1 is important for odontoblast-like cell differentiation. According to studies, in a case of direct pulp-capping, propolis paste partial dentine bridge formation was observed after four weeks. This is explained by reparative dentinogenesis stimulated by propolis flavonoids, which can produce tertiary dentine and affect dentinal fluid movement [18, 19].

LLLT or photobiomodulation (PBM) is known for using light energy to elicit biological responses from the cell and to normalize its function [25, 26]. A small fraction of laser energy at 810 nm wavelength reaches the pulp and transmits through the dental hard tissues. Therefore, the physiological cellular functions may normally be stimulated by LLLT. In addition, laser energy may promote the production of sclerotic dentine and the internal occlusion of dentine tubules. However, the results of in-vitro studies revealed that even such a small dosage could melt the dentine, producing a double-layer structures of tubules. Therefore, narrowing of the dentine tubules decreases the internal fluid movement, which could explain intermediate analgesic effect caused by depressing nerve transmission that stops the C-fiber afferents depolarization [13, 14].

The present study was designed based on the abovementioned facts. Even though our research showed good results in reducing DH, experimental studies are still necessary for understanding the exact mechanism of suggested type of DH treatment.

CONCLUSIONS

In our study, the efficacy of four types of DH treatment was assessed. All groups of patients showed statistically significant results. The highest results were obtained after a combined DH treatment. Therefore, combined application of ethanolic extract of propolis with diode laser (810 nm) irradiation can be recommended for DH treatment because of immediate and prolong reduction of the level of pain without any adverse reactions.

CONFLICT OF INTEREST

The authors declare no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

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