Effectiveness of benzocaine in reducing deep cavity restoration and post-extraction stress in dental patients

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ABSTRACT

benzocaine 20% للتحقق من فعالية التخدير الموضعي

الأهداف: لتخفيف الألم والإجهاد في المرضى بعد ترميم حفرة سنية عميقة (LA). وقلل الأسنان تحت التخدير الموضعي.

الطريقة: أجريت هذه التجربة السريرية الاستطلاعية من أكتوبر 2014 حتى أبريل 2015 بكليّة طب الأسنان، جامعة طيبة، المدينة المنورة، المملكة العربية السعودية. تم شمل 45 مريضاً في مجموعة saline و46 في المجموعة benzocaine 20%.

الطريقة: تم تقييم إجهاد الأسنان قبل العملية وبعدها باستخدام مقياس التوتر البصري (VAS). علاوة على ذلك، تم تسجيل حالات الانزعاج بعد كل علاج من قبل المرضى باستخدام مقياس VAS.

النتائج: كانت هناك فروق مهمة ذات دلالة إحصائية بين متوسط درجات التوتر للمريض في المجموعة saline وعمق السنوي السفلي، والخشي السفلي، والذي تتأثر به جماعته التي تحت التخدير الموضعي (benzocaine 20%) في موقع العملية.

الخاتمة: أثبتت هذه الدراسة أنه يمكن تحقيق التخدير الموضعي 20% في موقع العملية تحت التخدير الموضعي، وقلل الأسنان تحت التخدير الموضعي.

أعمال: يتم استخدام التخدير الموضعي 20% في علاج الأسنان تحت التخدير الموضعي في موقع العملية، وقلل الأسنان تحت التخدير الموضعي.

المتى: تم القيام بالتجارب السريرية الاستطلاعية في المملكة العربية السعودية من أكتوبر 2014 حتى أبريل 2015.

استنتاج: هذه الدراسة أظهرت أن التخدير الموضعي 20% يمكن أن يقلل من التوتر والإجهاد في المرضى بعد ترميم حفرة سنية عميقة (LA).

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Anxiety is one of the most common causes of dental fear.1 There have been 3 stable and reliable factors, which contribute to dental fear. The first factor is related to patterns of dental avoidance and anticipatory anxiety, the second, related to fear, which is associated to specific dental stimuli and procedures, and the third is concerned with physiologic arousal during dental treatment.2 Fear develops through the interaction of 3 phenomena, those which are instinctive, dependent on maturation, and
developed through learning from individual and social experience. It is unusual for modern local anesthetics to fail completely unless they have been injected incorrectly, but it is common for patients to retain some sensation of pressure around the tooth. Dental stress and anxiety may lead them to interpret this as pain, and cooperation is lost. Application of topical anesthetic for post-operative pain relief has not usually been considered as a part of routine clinical dental practice. However, infiltration local anesthesia (LA) has been used in other surgical disciplines with good effect. The field of anesthesia has witnessed many developments throughout the last century, and a number of different techniques and agents have been developed. Anesthesia has become an essential procedure in every surgical operation and treatment. In practice, topical anesthesia including cocaine, amethocaine, lignocaine, and prilocaine is applied to skin, eye, ear, nose, and mouth. Their application was reported to be useful and effective for reducing pain sensation. When used to produce topical anesthesia, they usually have a rapid onset of action (5-10 minutes [mins]), and a moderate duration of action (30-60 mins). A rapid onset of action of 5 mins or less has also been reported for 10% and 20% benzocaine gels for the relief of spontaneous toothache pain. Various preparations of lignocaine are available as aqueous solutions (4%), or in water-miscible bases, such as gels, ointments, creams, and sprays (2-10%). Intact skin works as a barrier preventing the diffusion of local anesthetics so high concentrations of anesthetic agents (for example, 20% benzocaine or 4% lignocaine) are required. The use of LA in dentistry is standard practice. No studies were found in the literature, which looked at the possibility of using benzocaine soaked rolls/swabs to reduce the post operative stress in adult patients following simple dental extractions, or deep cavity restoration. This study concentrated on the effectiveness of topical anesthetic 20% benzocaine for relieving pain of local anesthetic injection, and post-operative stress in patients following extraction of teeth, or deep restoration under LA.

Methods. This prospective study was conducted from October 2014 to April 2015 after obtaining the approval from the ethical committee. Patients who attended Taibah University, College of Dentistry (TUCoD), Al Madinah Al Munawarah, Kingdom of Saudi Arabia scheduled for extraction of teeth or deep cavity restoration under LA were considered for inclusion in the study. Using convenience-sampling pattern, 100 patients were selected to one of the 2 groups. Patients who fulfilled the following criteria were eligible for inclusion into the study: 1) male, aged 16-70 years of age. 2) scheduled for extraction of between 1-2 teeth or one deep cavity restoration. 3) American Society of Anesthesiologists I or II patients. 4) where the patient was able to understand and cooperate with the requirements of the protocol, and were able and willing to provide an appropriate written informed consent. Patients who were excluded from the study were hypersensitive or allergic to topical anesthesia, needed more than 3 teeth extraction, disliked the taste of the anesthetic, and refused to continue, or had vomiting reflex. Prior to the study, a researcher allocated the sequence of patient identity numbers to either the test or control group. Slips of paper with the test group or the control group were placed in opaque envelopes, and sealed by a secretary who was not associated with the study. These envelopes had been numbered sequentially on their outside with the patient identity number and was attached to the patient’s dental hospital treatment record.

On the dental chair, once the patient signed the consent a dental assistant placed the swab over the site of injection in the mouth for 20 second. If the slip in the envelope said that the patient was in the study group, small swabs were painted with topical anesthetic paste 20% benzocaine (Sky-Caine Gel, Skydent Inc., NY, USA). If the slip in the envelope said control group, the patient had the swab impregnated with cold normal saline mixed with one lozenge honey and lemon flavored Strepsils (Reckitt Benckiser Healthcare International Ltd., Nottingham, England). The slip was placed back in the envelope, which was placed back into the patients’ records. This ensured that the patient and the dentist carrying out the assessment were blind. Once the swabs were removed, the patient received 1.8 ml mepivacaine 2% with epinephrine 1:100,000. For the upper teeth, patients received buccal and palatal infiltrations. For lower teeth, they had either buccal and lingual infiltrations, or inferior alveolar nerve block additional to buccal injections. This was the usual clinical practice. Standard extraction techniques using elevators and dental forceps were employed. For deep caries, cavities prepared with high-speed contra angle hand piece.

The researcher was just an observer. Each patient was assessed for levels of stress, and these assessments

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were recorded using the visual analogue scale (VAS). The scores given were 0 if no sign of stress, or 100 for very severe stress. Evaluation of stress was made pre-operatively and immediately post-extraction. The discomfort of the injections were recorded by the patients after each treatment on standard 100 mm VAS, tagged at the endpoints with “no pain” (0 mm) and “unbearable pain” (100 mm). The researcher who made all the stress and pain observation was completely independent of the whole process. Statistical analysis was performed using Statistical Package for Social Sciences, version 21 (IBM Corp., Chicago, IL, USA). T-test was carried out to know significance level and \( p \)-value was adjusted to \( p<0.05 \). Sample size calculation was made for this study based on a study by Gazal and Mackie.\(^\text{10}\)

**Results.** Out of the 100 patients, 6 were excluded because they fall into the exclusion criteria. Ninety-four patients were randomly allocated to one of the 2 groups. However, at the end of the study there were missing data for 3 patients (one in the normal saline group who fainted after local anesthetic injection, and 2 in the benzocaine group because their teeth extraction was performed surgically), giving 45 in the benzocaine group (study) and 46 in the normal saline group (control). Thus, the final sample size included 91 patients aged between 16-66 years (mean 37.70 years, standard deviation 14.50). The teeth had deep caries for restoration, or it was extracted due to gross caries, which could not be restored.

There were no statistically significant differences between the mean stress scores for the treatment (20% benzocaine) and control (normal saline) groups at the baseline assessment pre-operatively \((p=0.97)\) (Table 1). However, there were statistically significant differences between the mean stress scores for the benzocaine and normal saline groups post-operatively \((p=0.002)\) (Table 1). For both benzocaine and normal saline groups, changes in stress scores from the pre-operative score to the post-operative score were made using the paired sample t-test. There were significant decreases in stress scores between the pre-operative \((p=0.003)\), and post-operative scores \((p=0.008)\) (Table 2).

The discomfort of the injections was recorded by the patients after each treatment on standard 100 mm VAS,
tagged at the endpoints with “no pain” (0 mm) and “unbearable pain” (100 mm). There were significant differences between the mean pain scores for patients in the post buccal injection \( (p=0.001) \), post palatal injection \( (p=0.01) \), and the post inferior alveolar nerve block groups \( (p=0.02) \). These figures are illustrated in Table 3, Figures 1 A & B, and Figure 2 A & B.

Patients in the benzocaine group were more comfortable during administration of local anesthetic than the normal saline group. However, there were no significant differences between the mean pain scores for the patients in benzocaine and normal saline groups post lingual injection \( (p=0.075) \) (Table 3). Clinically, lingual injections were more painful for patients in the normal saline group than in the benzocaine group. This finding did not achieve a statistical significance because the number of patients who received lingual injections was small. Thus, a large sample size of patients in the lingual injection group might have significant differences in pain injection scores.

**Discussion.** Injection of LA is considered one of the main reasons for dental fear and dental visiting avoidance. Efforts have been performed to reduce pain perception of needle injection.\(^{11,12}\) Dental treatment fear and anxiety gradually builds up during childhood as results of a bad experience caused by dentist misbehavior, painful tooth preparation, or needle injection.\(^{13}\) Local anesthesia is necessary for a painless dental treatment. However, patients sometimes feel the needle puncturing the mucosa, or the pressure of depositing the local anesthetic solutions is painful. Different techniques have been employed in dentistry to minimize the pain on injection. Use of topical anesthesia, low-pressure

![Figure 1 - Pain scores compared between patients in benzocaine and normal saline groups regarding post buccal and palatal injections (A & B).](image)

![Figure 2 - Pain scores compared between patients in benzocaine and normal saline groups regarding post lingual injection inferior alveolar nerve block.](image)
injection, narrow, sharp needles, and a slow injection rate were not enough to achieve a completely painless injection under all circumstances. A large number of dentists believe that fearful patients are difficult and more stressful to treat, and they are most likely to attend appointments irregularly.

The result of this study showed that there was a statistically significant difference in mean pain and stress scores between the patients in the topical benzocaine and those in the normal saline groups (p<0.05). The application of 20% benzocaine as a topical anesthetic for intraoral injections did reduce the pain of needle insertion, post-operative stress, and uneasiness connected with dental extractions under LA. The results of the current study are consistent with earlier findings of previous studies. All topical anesthetics are similarly reducing pain associated with needle puncture. When EMLA-60 was applied on oral mucosa, it seemed to have longer duration of action. Hodosh et al conducted a double-blind randomized control trial on 100 patients to investigate the effectiveness of a combination of different types of topical local anesthetics for reducing pain during maintenance visits. A 35% potassium nitrate; 20% benzocaine; 10% tetracaine/aqueous hydroxyethyl cellulose gel was applied to the teeth and gingiva prior to hygienist-administered maintenance treatments. The outcome of this study revealed that the application effectively anesthetized the teeth and gingiva, so the dental work was performed with comfort. A study demonstrated that the combination of EMLA cream with propofol is beneficial and providing safe and effective pain management for lumbar punctures.

On the contrary, a study was conducted to evaluate the patient’s perceived pain response to the injection and anesthetic deposition for the greater palatine nerve block. Their findings revealed that the use of pressure and topical analgesia (20% benzocaine) did not reduce posterior palatal pain. However, in the current study, the patients in the benzocaine group reported less pain scores following palatal injection compared with those that are in the control group. From the clinical point of view, the palatal infiltration technique used in this study is more comfortable than the greater palatine nerve block because it minimizes the risk of direct contact of the needle with greater palatine nerve.

In this study, it was noticed by the dental surgeon who carried out all teeth extractions, that there was a group of patients who recorded a high level of dental stress at the baseline assessment; they scored less after completing the tooth extraction, or deep cavity preparation procedure. One possible explanation was given by these patients is that their needle phobia was eased off by having painless dental injections. Similarly, psychological processes, such as social appearance, anxiety, and blood injection phobia may affect the phobia of dental procedures. Targeting these points related psychological constructs may improve the management of dental anxiety among adult patients.

Despite using topical benzocaine there was a group of patient who scored mild stress scores pre-operatively, they reported high level of stress post-operatively. These patients stated that seeing blood on the dental gauze and hearing the noise of surgical instruments during extraction procedure scared them and they became stressful post-operatively. However, dental injection pain can sometimes be a result of too rapid administration of local anesthetic, or due to the difference in pH value between the local anesthetic solution and the soft tissues in the mouth. Rapid injection can tear the tissue and result in immediate pain followed by soreness. The duration of discomfort varies a lot between different injections sites and techniques. Administration of oral ibuprofen pre-operatively in patients with irreversible pulpitis increased the success rates of inferior alveolar nerve block anesthesia and eased needle pain.

In conclusion, this investigation has demonstrated that the post-operative stress associated with dental extractions and deep cavity restoration under LA can be reduced by the application of topical anesthetic (20% benzocaine) at the surgical site for intraoral injections. Topical anesthetic can be of value to help patients with needle phobia and reduce their level of fear. It is strongly recommended for all dental clinics and institutions to administer topical anesthetic before giving local anesthetic injections. In the study sample only males were included as only male patients are treated in TUCoD, so females should be included in future studies for more valid and stronger results.

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### Ethical Consent

All manuscripts reporting the results of experimental investigations involving human subjects should include a statement confirming that informed consent was obtained from each subject or subject’s guardian, after receiving approval of the experimental protocol by a local human ethics committee, or institutional review board. When reporting experiments on animals, authors should indicate whether the institutional and national guide for the care and use of laboratory animals was followed.