Effect of Pulpotomy Procedures With Mineral Trioxide Aggregate and Dexamethasone on Post-endodontic Pain in Patients with Irreversible Pulpitis: A Randomized Clinical Trial

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ABSTRACT

Objective: Endodontic post-treatment pain continues to be one of the main problems encountered by dental professionals. Therefore, pain control during and after endodontic treatment is one of the most important issues in endodontics. The purpose of this clinical trial was to compare postoperative pain relief achieved with dexamethasone (DEX) and mineral trioxide aggregate (MTA) used as pulp coverage after pulpotomy in human molars with irreversible pulpitis.

Methods: This prospective double-blind study was conducted on 54 patients complaining of dental pain due to irreversible pulpitis. The standard pulpotomy procedure was performed by the same dentist in all patients. At the time of the cotton pellet placement, patients were randomly divided into three groups: those in whom a sterile dry cotton (DC) pellet was used, patients treated with a cotton pellet soaked in MTA, and those who were treated with a cotton pellet soaked in DEX. After completion of the treatment, patients received rescue medication every 6 hours for the first day. Postoperative pain was assessed at 6-hour intervals for 24 hours, and then every day until day 7 using a visual analog scale.

Results: In general, patients treated with MTA suffered the lowest levels of pain at all time intervals. Post-pulpotomy pain was significantly reduced at 18 and 24 hours and from days 2 to 7 post-treatment in the MTA group. DEX lowered the pain level more than the DC pellet. However, the differences observed in the mean pain scores of the DEX and DC pellet groups at all-time intervals were not statistically significant.

Conclusion: Pulpotomy procedures can reduce pain related irreversible pulpitis. Pulpotomy with MTA-soaked cotton pellet significantly reduces pain intensity in patients with irreversible pulpitis.

Keywords: Dexamethasone, irreversible pulpitis, mineral trioxide aggregate, pulpotomy, postoperative endodontic pain, visual analog scale

INTRODUCTION

Pain of odontogenic origin has long been a source of fear and anxiety among the general population (1). It has also been a common source of frustration among dentists who have difficulties to alleviate their patient’s pain (2). Endodontic post-treatment pain continues to be a significant problem facing the dental profession (3). Therefore, pain control during and after endodontic treatment is one of the most important issues in endodontics. Consequently, inhibiting the post-treatment pain that affects both the patients and dentists is an important factor to consider when performing dental treatment.

Endodontic pain is often an emergency situation that necessitates immediate treatment. It has been indicated that emergency pulpotomy is very effective in relieving acute dental pain caused by acute pulpitis. Irreversible pulpitis is the most common reason for performing endodontic treatment, characterized by prolonged sensitivity to cold or heat (4).
Immediate postoperative pain is most likely to occur in patients with irreversible pulpitis following treatment. It has a high occurrence of 25%–40% (5). This pain is influenced by several factors, such as patient anxiety, pulpal status, existence of pre-treatment pain, and manipulation of periapical tissues (6).

Many medications have been prescribed to relieve pain in irreversible pulpitis. Opioid and non-opioid analgesics, benzodiazepines, non-steroidal anti-inflammatory drugs, corticosteroids, or antibiotics, and even usage of mineral trioxide aggregate (MTA), have been prescribed in dentistry (7-10). Among the several treatment options, an emergency pulpotomy is the most reliable way to obtain pain relief in vital pulp cases (4).

Mineral trioxide aggregate has attracted attention in the field of endodontics as a potential medicament for pulpotomy procedures, pulp capping, apexification, repair of root perforation, and repair of resorptive defects (11, 12). It also has favourable sealing ability, biocompatibility, and the ability to induce dentine bridge formation, along with cementum and periodontal ligament regeneration (13).

The pain following endodontic therapy is often linked to the inflammatory process (14). Glucocorticoids inhibit the production by multiple cells or factors that are important in producing the inflammatory response (15). The post-pulpotomy pain-relieving effect of dexamethasone (DEX) as sedative dressing has not been evaluated previously. In addition, most of the studies involving MTA have only investigated its effect on dentine bridge formation rather than its analgesic effects. Therefore, the purpose of this clinical trial was to compare the postoperative pain relief of DEX and MTA after pulpotomy in human permanent molars with irreversible pulpitis. Furthermore, the relationship between pre- and postoperative pain levels was also investigated. The null hypothesis was that there is no significant difference among the three pulp capping agents in terms of post-operative pain.

**MATERIALS AND METHODS**

**Ethical statement**

The research proposal was reviewed and confirmed by the ethics committee of Kerman University of Medical Sciences, Kerman, Iran. Patients involved in the study were formally informed, and each patient signed a consent form prior to enrollment. Consent form provided patient comprehensive information about the purposes and background of the study, clinical procedure, risks and discomfort, potential benefits, costs and payment, confidentiality, use and disclosure of health information, and subjects’ rights and freedom to quit from the study at any time.

This prospective double-blind study was conducted on 54 patients who experienced dental pain due to irreversible pulpitis, which was determined based on clinical examination. Therapeutic interventions were performed by two different clinicians.

All patients with a single painful and restorable molar indicating symptomatic irreversible pulpitis (i.e., a history of spontaneous pain for a few seconds to several hours, pain exacerbating with hot and cold fluids, radiating pain, or reproducible pain with cold testing) were included in the study.

Exclusion criteria were the following: patients who used analgesics for more than 1 day, patients taking antibiotics, patients with prior history of root canal therapy, tenderness to percussion, pulp necrosis as diagnosed by thermal and electrical tests, evidence of periapical radiolucency, active systemic disease, moderate or severe marginal periodontitis, internal/external root resorption in periapical radiograph, patients whose tooth mobility was greater than Grade 1 and diffuse pulp chamber calcification. In this study, those teeth with no bleeding as an indicator of pulp vitality after access cavity preparation were also excluded.

Pulpal diagnosis was determined by cold test and electric pulp test. Periapical diagnosis was based on percussion and palpation examination along with radiographic evaluation. A comprehensive clinical examination was also performed to rule out intra/extraoral swelling, the presence of a sinus tract, or other major pathology.

The standard pulpotomy procedure was performed by the first dental practitioner for all the patients. Inferior alveolar nerve block (for mandibular teeth-2% Lidocaine 1:100,000 epinephrine carpalues (DarouPaksh, Tehran, Iran) and buccal infiltration (for maxillary teeth) were administered, the teeth were isolated with a rubber dam, caries were removed, and then the access cavities were prepared. After the removal of the coronal pulp tissue, the pulp chamber was irrigated with sterile saline solution and dried with sterile cotton pellets. Pulp covering and temporary restoration were performed by a second dental practitioner. At each treatment step and follow-up visit, clinical assessment of the treated tooth, which was only identified by a specified code, was carried out by two experienced dental practitioners who were blinded to the treatment groups.

The patients were randomly divided into three groups. Group 1: sterile dry cotton (DC) pellet; Group 2: a cotton pellet soaked in white MTA (Proroot, USA). Group 3: 8 mg/2 ml DEX (Osveh, Iran). In the MTA group, the cotton pellet was prepared by soaking the pellet in diluted white MTA (3:1 liquid to powder ratio). After placing the cotton pellet in the pulp chamber, the access cavity was properly sealed with Coltosol (Ariadent, Iran).

Patients were given postoperative forms that included instructions on rescue medication guidelines (400 mg Ibuprofen every 6 hours for the first 24 hours and then as needed. In case the patient was unable to take Ibuprofen, 500 mg acetaminophen), and recording of the pain level score. Pain was recorded using a visual analog scale (VAS) for pain from 0 (no pain) to 10 (worst pain) at 6, 12, 18, and 24 hours and 2 to 7 days after treatment.

An emergency telephone number was provided to patients to be able to report any possible problems. Treatment was rendered for patients who required any additional treatment, such as a pulpectomy; however, they were excluded from the study.
Patients were contacted by phone daily and appointments for RCT were scheduled at the end of 7 days. Patients were asked to record their pain severity in the provided charts. After the 7th day, the patients handed in the completed charts, and an appointment was scheduled to complete the treatment procedure.

**Statistical analysis**

Data analysis was done using the SPSS 20.0 software package. All data were expressed as ±1 standard deviation (SD). After testing for normality of pairwise differences with the Shapiro–Wilk normality test, the mean values of data were calculated, and statistical significance was determined with the analysis of variance to compare the effect of both sedative dressings on the severity of postoperative pain between the different groups. A P<0.05 was considered to be statistically significant.

**RESULTS**

A total of 54 patients were enrolled for this clinical trial. Forty-five patients (27 women and 18 men) completed the 7-day follow-up of the study. Nine patients left or were excluded from this study (Diagram 1). The demographic data of three groups of patients with irreversible pulpitis are presented in Table 1.

All patients reported their pain levels at 6-hour intervals for a period of 24 hours and then every 24 hours until day 7 after treatment.

To compare the three therapeutic methods, the average VAS score was calculated for each patient at different time points. Pre-treatment VAS was as follows: 25% ≤5.5, 50% ≤7, 75% ≤8, and 25% ≥8 (Table 2). No significant differences in pre- and postoperative pain scores were seen among the three groups with respect to age, gender, and tooth location (P>0.05) (Tables 1 and 2). However, 24 hours after treatment and on days 6 and 7, female patients complained more of severe pain compared to male patients.

Regardless of the pulpotomy procedure, the mean VAS for pain showed that pulpotomy was efficacious in reducing post-treatment pain in all patients with irreversible pulpitis (P<0.001) (Table 2). It was observed that the differences between the VAS scores of the three treatment methods were significant (P=0.014).

In the group treated with MTA, the patients had no post-treatment severe pain at any time interval except at 6 (20% cases) and 12 (5% cases) hours postoperatively (Fig. 1). Severe pain was reported at all-time intervals by dry cotton- and DEX-treated patients (except 3 and 4 days post-treatment) (Figs. 2 and 3). The percentage of subjects reporting no or mild pain after the treatment period was 20%–100% for MTA-treated, 5%–95% for DEX-treated, and 16%–80% for DC-treated groups (Figs. 1–3).

### Table 1. Demographic data of three groups

| Group/No   | Gender | Age  |
|------------|--------|------|
|            | Male   | Female | Mean±SD | Range |
| DC/15 cases | 4 (26.7%) | 11 (73.3%) | 30.47±10.13 | 18-44 |
| MTA/15 cases| 9 (60.0%) | 6 (40.0%) | 28.87±11.03 | 19-52 |
| DEX/15 cases| 5 (33.3%) | 10 (66.7%) | 26.07±5.65 | 21-36 |
| Total      | 18 (40.0%) | 27 (60.0%) | 28.47±9.21 | 19-52 |

### Table 2. Pre- and postoperative VAS for pain of all patients regardless of treatment

| Time point          | n | Mean±SD | Minimum | Maximum | 25th | 50th | 75th |
|---------------------|---|---------|---------|---------|------|------|------|
| Pre-operative VAS   | 45 | 6.51±1.39 | 3.00    | 9.00    | 5.50 | 7.00 | 8.00 |
| VAS 6 hour          | 45 | 4.02±2.80 | 0.00    | 9.00    | 2.00 | 4.00 | 6.00 |
| VAS 12 hour         | 45 | 3.02±2.58 | 0.00    | 7.00    | 0.00 | 3.00 | 5.00 |
| VAS 18 hour         | 45 | 2.31±2.38 | 0.00    | 7.00    | 0.00 | 2.00 | 4.00 |
| VAS 24 hour         | 45 | 1.93±2.24 | 0.00    | 8.00    | 0.00 | 2.00 | 3.00 |
| VAS 2 day           | 45 | 1.48±2.19 | 0.00    | 9.00    | 0.00 | 0.00 | 2.50 |
| VAS 3 day           | 45 | 1.48±1.98 | 0.00    | 9.00    | 0.00 | 0.00 | 3.00 |
| VAS 4 day           | 45 | 1.60±2.31 | 0.00    | 9.00    | 0.00 | 0.00 | 3.00 |
| VAS day             | 45 | 1.62±2.53 | 0.00    | 9.00    | 0.00 | 0.00 | 3.00 |
| VAS 6 day           | 45 | 1.46±2.37 | 0.00    | 9.00    | 0.00 | 0.00 | 2.00 |
| VAS 7 day           | 45 | 1.46±2.44 | 0.00    | 9.00    | 0.00 | 0.00 | 2.00 |
significant at 6 and 12 hours post-treatment (P>0.05) (Table 3). However, a statistically significant difference at 18 and 24 hours and 2–7 days post-treatment was found in the groups (Fig. 4). The DEX treatment was also associated with lower VAS levels compared with the DC-treated group (Table 3). However, no significant differences were observed in the mean pain scores of the two groups.

**DISCUSSION**

The primarily aim of this study was to determine and compare the post-pulpotomy pain-relieving the effect of administration of DEX and MTA as sedative dressings in patients with irreversible pulpitis. Furthermore, this study was designed to investigate the relationship between pre- and post-treatment pain levels in these patients. The control of postoperative endodontic pain continues to pose a significant challenge for dentists. Among various dental procedures, endodontic treatment is associated with more frequent and severe postoperative pain (16, 17).

New strategies of pain management with optimal analgesic efficacy and minimum incidence of side effects are always needed to prevent and control the occurrence of postop-
Objective: The postoperative pain may also develop when the integrity of the periapical tissues is compromised. Irritation of periapical tissues caused by local trauma, caustic irrigating solutions, or contamination during endodontic treatment may cause an acute inflammatory reaction, potentially leading to pain and/or swelling. Many chemical mediators (prostaglandins, leukotrienes, bradykinin, etc.) have been associated with the inflammatory process. The concentration of bradykinin, a potent pain mediator, was 13 times greater in teeth with irreversible pulpitis compared to the normal pulps (28). These chemical mediators cause pain at the site of inflammation (29). Pain can be induced indirectly by an increase in edema and tissue pressure, or directly through lowering the excitability threshold of the A-delta and C-fibers (18).

DEX is a steroidal anti-inflammatory drug that inhibits phospholipase A2 and consequently reduces the amount of chemical mediators, decreasing the polymorphonuclear leukocyte chemotaxis (30). However, it has been shown that the administration of DEX leads to a partial inhibition of development of inflammatory periapical lesion. In comparison with the DC-treated group, DEX was not able to significantly reduce pain intensity at any time interval. It has been suggested in previous oral pain studies that the absorption and distribution of the medication occur before the initiation of surgical tissue damage and consequent production of inflammatory mediators to provide analgesia to patients before endodontic treatment is started (18).

CONCLUSION

Pulpotomy procedures can reduce pain related irreversible pulpitis. Pulpotomy with MTA-soaked cotton pellet significantly reduces pain intensity in patients with irreversible pulpitis undergo pulpotomy.

Disclosures

Conflict of interest: The authors deny any conflicts of interest related to this study.

Ethics Committee Approval: The research proposal was reviewed and confirmed by the ethics committee of Kerman University of Medical Sciences, Kerman, Iran.

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