

THE ASSESSMENT OF CORONAL TOOTH DISCOLORATION WITH USE OF MICROMEGA MTA OR MTA+ AS THE PULP-CAPPING MATERIAL#

PULPA KAPAKLAMA MATERYALİ OLARAK MİKROMEGA MTA VE CERKAMED KULLANIMIYLA MEYDANA GELEN KORONAL DİŞ RENKLENMESİNİN **DEĞERLENDİRİLMESİ**[#]

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ABSTRACT

Aim: Mineral trioxide aggregate and calcium silicate cements have common usage in endodontics and restorative dentistry. However, MTA has some disadvantages such as long setting time, discoloration and cost. The treatment can not finish in a single visit because of long setting time. The aim of this study is to determine the discoloration of different materials on the coronal dentine.

Material and Methods: In this study, fifty bovine teeth were prepared and filled with Proroot MTA, MM-MTA, Biodentine, MTA+ (Cerkamed) placed and then sealed with a translusent composite. Unfilled samples were determined as a control groub. The specimens were kept at 37 °C in a dark environment. The color assessment was performed a with a spectrophotometer at different intervals (1th day, 1 month, 3 months, 9 months). The statistical analysis was performed by using One way Anova and Post-hoc Tukey tests.

Results: In the analysis of the tooth discoloration, the materials (Pro Root MTA, MM-MTA and MTA+ Cerkamed) were showed discoloration by the time. Biodentine showed tooth color stability because of zirconium oxide as a radiopacifier. The discoloration dearee is very high between 1th month and 3th month.

Conclusions: Different discoloration degrees of materials which are used in vital pulp treatments have been determined in this study. This criteria should also be taken into consideration if a dental vital pulp treatment isplanned in a tooth which may be anesthetically anxious.

Keywords: MTA, discoloration, vital pulp treatment

ÖΖ

Amac: Mineral trioksit agregat (MTA) ve kalsiyum silikat bazlı simanlar endodonti ve restoratif tedavisinde sıklıkla kullanılmaktadır. Ancak MTA'nın sertleşme süresinin uzun olması, dişlerde renklenmeye sebep olması ve maliyetinin fazla olması gibi dezavantajları bulunmaktadır. Bu çalışmanın amacı farklı materyallerin koronal materyallerin koronal dentinde yaptığı renklenmevi değerlendirmektir.

Gereç ve Yöntem: Bu çalışmada elli adet sığır dişinden örnekler hazırlanıp 5 gruba ayrıldı. Gruplara Proroot MTA, Biodentin, MM-MTA, MTA+ (Cerkamed) yerleştirildi ve daha sonra translusent kompozit ile restore edildi. Kontrol grubundaki örneklere herhangi bir işlem uygulanmadı. Örnekler karanlık ortamda 37°C'de bekletildi. Renk değişimleri bir spektrofotometre ile farklı zaman aralıklarında (1. gün, 1 ay, 3 ay, 9 ay) ölçüldü. İstatistiksel analiz Tek yönlü Anova ve Post-hoc Tukey testleri kullanılarak yapıldı.

Bulgular: Çalışmada elde edilen verilere göre; Pro Root MTA, MM-MTA ve MTA+ (Cerkamed) kullanılan gruplarda zamanla renklenme gözlendi. Biodentinin iceriğinde bulunan zirkonyum oksit nedeniyle renklenme meydana gelmedi. Renklenme en fazla 1. av ve 3. avda görüldü.

Sonuç: Vital pulpa tedavilerinde kullanılan materyallerin diş renklenmesi üzerine etkisinin değerlendirildiği bu çalışmada; materyallerin farklı renklendirme dereceleri belirlenmiştir. Estetik açısından kaygı oluşabilecek bir dişte vital pulpa tedavisi planlanıyorsa bu kriter de göz önüne alınmalıdır.

Anahtar Kelimeler: MTA, renklenme, vital pulpa

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INTRODUCTION

Mineral trioxide aggregate (MTA) is very popular for vital pulp treatment, apexification of immature teeth and perforation repair1-3. MTA is a biologically safe material, but has some major disadvantages, namely its long setting time and the discoloration that it causes to both teeth and restorative materials^{4,5}. A new calcium silicate-based material called Biodentine (Septodont, Saint-Maurdes Fossés, France) has been developed to overcome these disadvantages^{6,7}. Biodentine contains calcium carbonate, tricalcium silicate liquid containing calcium chloride and zirconium oxide radiopacifier8. The developers added chloride to Biodentine to shorten its setting time to about 12 minutes⁹. The zirconium oxide guarantees the color stability of Biodentine¹⁰. The rapid setting time means that treatment can be completed within a short period.

Micromega MTA (MM-MTA) is another newly developed calcium silicate cement, composed of dicalcium silicate, calcium sulfate dehydrate, tricalcium aluminate, tricalcium silicate, magnesium oxide, bismuth oxide and calcium carbonate (CaCO3)¹¹. The CaCO₃ provides MM-MTA with a short setting time of about 20 minutes. MM-MTA can be used for practical applications because it is encapsulated. These beneficial characteristics make MM-MTA a good choice as a pulp-capping material^{12, 13}.

MTA+ is a new pulp-capping material composed of calcium oxide, silicon oxide, sodium oxide, iron oxide, aluminum oxide, potassium oxide, bismuth oxide, magnesium oxide, zirconium oxide and calcium phosphate. According to the manufacturer, MTA+ can be used in direct pulp-capping, root canal wall perforation, intracanal resorption, posterior root apex filling, pulp amputation, and the treatment of teeth with interrupted root development. The manufacturer claims that MTA+ is the only MTA product made of nanoparticles, and that the MTA+ particles are three times smaller than those of other MTAs. There are studies in the literature on the discoloration of Biodentine and ProRoot MTA. However, there is only one study concerning the discoloration of MM-MTA, and there is no information about MTA+ discoloration.

The aim of this *in vitro* study was to assess of the discoloration of MTA with nanoparticles (MTA+) when it is applied as a pulp-capping material.

MATERIALS AND METHODS Sample preparation

The materials used in this in vitro study and their chemical compounds are shown in Table 1. We obtained 50 bovine central incisors and examined them under a microscope to check for fractures or structural defects (×3.5 magnification). The crowns and roots were separated; then, the crowns were cleaned and polished with a polishing paste. We prepared 10 x 10 mm disks from the facial surfaces of the crowns. We used a digital caliper to ensure that the thickness of each disk was standardized to 2.00-2.03 mm. Cavities were induced (diameter=5 mm, depth=1.5 mm) in the centers of the dentinal surfaces (Figure 1). The specimens were then immersed in 1% sodium hypochlorite for 30 minutes, washed with water, and placed in 20% EDTA (pH 7.7) for 2 minutes¹⁴. The specimens were washed and dried with cotton pellet then were randomly assigned to five subgroups (n = 10).

 Table 1. Materials used in the study

| Material/manufacturer | Chemical compounds |
|--|---|
| Proroot MTA (Dentsply, Tulsa, OK, ABD) | Tricalcium silicate, dicalcium silicate, tricalcium aluminate, tetracalcium aluminoferrite. |
| MTA+ (Cerkamed Medical Company, Stalowa, Poland) | Potassium oxides, aluminium oxides, ferric oxides, potassium sulphate, sodium sulphate |
| Biodentine (Septodont, Saint- Maurdes-Fosses, France) | Powder: tricalcium silicate, dicalcium silicate, calcium carbonate and oxide, iron oxide, zirconium oxide Liquid: calcium chloride, hydrosoluble polymer |
| MM-MTA (Micro Mega, Besançon, France | Tricalcium silicate, dicalcium silicate, tricalcium oxide, tricalcium aluminate, tetracalcium aluminoferrite, calcium sulphate, silicate oxide, bismuth oxide, calcium carbonate |



Figure 1. Dentin disk preparation

Group 1 was the control group, in which no MTA was applied. Groups 2–5 cavities were filled as follows: Group 2, ProRoot MTA; Group 3, MTA+; Group 4, Biodentine; and Group 5, MM-MTA. We then placed moist pellets on the samples and incubated them at 37°C for 24 hours.

After the 24 hour incubation period, dentinal surfaces were etched 37% orthophosphoric acid for 30

seconds, washed with water and dried. We applied adhesive (single-bond universal 3M ESPE, St. Paul MN, USA) and cured the samples using an LED light curing device (VALO Cordless LED curing light; Ultradent Products Inc., South Jordan, UT, USA). We then sealed the entire dentinal surface with a translucent composite (Amaris Translucent; Voco GmbH, Cuxhaven, Germany) and stored the specimens in distilled water at 37°C in a dark environment.

After 24 hours, we dried the teeth with a cotton pellet and conducted color measurements of all of the specimens using a spectrophotometer (Easyshade, Vita Zahnfabrik, Bad Sackingen, Germany). We calibrated the instrument prior to taking the measurements for each group, which were repeated after 30, 90 and 180 days. The colors were evaluated with respect to the CIE formula (L*a* b*) (14); the differences (ΔE_{ab*}) between the four color measurements were calculated as follows: $\Delta E_{ab*} = [(\Delta L*)^2 + (\Delta a*)^2 + (\Delta b*)^2]^{\frac{1}{2}}$.

Statistical analysis

We performed the data analysis SPSS software (ver. 17.0; IBM Corp., Armonk, NY, USA). We conducted a one-way analysis of variance (ANOVA) to evaluate whether the mean differences between ΔE for each group were statistically significant. When the p-values in the one-way ANOVA were statistically significant, we used a post-hoc Tukey HSD test to establish the groups responsible for the differences. A p value of less than 0.05 was noted to be statistically significant.

RESULTS

Table 2 indicates ΔE measurements regarding for groups and follow-up times. Figure 2 shows a image of a specimen from each group at the different time point.

| | 1" day | 1"month | 3 rd month | 9 th month |
|------------|--------|---------|-----------------------|-----------------------|
| Control | (A) | | | |
| Biodentine | | | | N |
| ProRoot | 1 | | | |
| MM-MTA | | | | |
| MTA+ | | | | |

Figure 2. A spectrophotometric image of a tooth from each group at the different time poin

There were no statistically differences between the ΔE levels of the control and Biodentine groups at each time point (p = 0.075/ p=0.361).

There were statistically significant differences among ProRoot, MTA+ and MM-MTA groups in ΔE levels at each time point (p <0.001). The samples in the ProRoot group showed more discoloration after 3 months than after 1 month (p <0.001). In the MTA+ and MM-MTA groups, there was more discoloration after 3 and 9 months, respectively, than after 1 month (p <0.001).

Table 2. ΔE measurements regarding for groups and follow-up times

| | 1 st day | 1 st month | 3 rd month | 9 th m onth | p-value 🕈 |
|------------|--------------------------|---------------------------|--------------------------------------|-------------------------------|-----------|
| Control | 2.18±0.84 ^{A,a} | 3.34±2.66 ^{A,a} | 6.16±4.20 ^{A,a} | 8.41±6.46 ^{A,B,a} | 0.075 |
| ProRoot | 3.84±2.40 ^{A,a} | 11.08±3.71 ^{B,b} | $15.76 \pm 2.01^{\text{B},\text{c}}$ | 15.24±3.01 ^{A,C,b,c} | <0.001 |
| MTA+ | 4.04±3.97 ^{A,a} | 17.16±3.24 ^{C,b} | 23.02±3.72 ^{C_f} | 22.76±4.35 ^{Dg} | <0.001 |
| Biodentine | 2.33±1.42 ^{A,a} | 2.92±1.50 ^{A,a} | 5.12±3.64 ^{A,a} | 5.58±4.29 ^{B,a} | 0.361 |
| MM-MTA | 2.40±2.02 ^{A,a} | 7.99±2.75 ^{B,b} | 15.72±3.34 ^B [#] | 17.96±5.22 ^{C,D,c} | <0.001 |
| p-value ‡ | 0.239 | <0.001 | <0.001 | <0.001 | |

DISCUSSION

MTA is the best pulp-capping material currently available. Histologically, MTA forms a better dentin bridge than Ca(OH)¹⁵. However, tooth discoloration, which is undesirable in esthetic terms, is inevitable when conventional MTA is used as the pulp-capping material. In this in vitro study, we investigated the discoloration effects of four pulp-capping materials. There are many reports of tooth discoloration following MTA treatment in the literature, but there are no studies concerning discoloration due to MM-MTA and MTA+.

Both bovine and human incisors have been used in MTA discoloration studies^{9, 16}. In this study, we decided not to use human tissue for ethical reasons. Hence, we used bovine incisors taken from animals that had been slaughtered for consumption. The texture of bovine teeth tissue is like to human teeth, and it is easier to prepare standard specimens from bovine teeth.

ISO 9241-302 recommends calculating color differences using the three-dimensional CIE L*a*b*and CIE 2000 color measurement formulas; we used the former formula to evaluate discoloration in this study. Although CIE2000 is newer, the L*a*b* formula is still used in many dentistry color studies.

The results of our study showed that MM-MTA, MTA+ and ProRoot MTA all cause significant discoloration, which increases over time. In these cases, the discoloration may be caused by bismuth oxide, which all three of these MTAs contain (serving as a radiopacifier). The oxidation of heavy metal oxides, such as iron oxide and, particularly, bismuth oxide may cause discoloration. There are several factors that contribute to discoloration in the case of bismuth oxide: first, oxidized bismuth oxide may react with carbon dioxide in the air to produce bismuth carbonate, resulting in discoloration. It has also been reported that bismuth oxide undergoes thermal dissociation at high temperatures, yielding metallic bismuth and oxygen¹⁷, and that bismuth oxide is converted to a black precipitate when it interacts with dentin collagen¹⁸. We used a high-intensity light curing device to polymerize the bonding agent in this study. This may have contributed to the increase in temperature, and hence the increase in discoloration.

Only one study reported that Biodentine may show discoloration after 1 year. The other studies reported no discoloration of Biodentine, which accords with the present study. The radiopacifier in Biodentine is zirconium oxide, which does not cause discoloration. To prevent discoloration, it may be preferable to use zirconium oxide instead of bismuth oxide in future silicate cements.

There was more discoloration in the MTA+ group than in the Pro Root and MM-MTA groups in this study, especially during the first month, in which dramatic discoloration occurred. As mentioned previously, the manufacturer claims that MTA+ is composed of nanoparticles that are three times smaller than those in other MTAs. The nanoparticles in MTA+ may penetrate deeper into the dentin, and then react to cause more discoloration.

MM-MTA is a newly developed, fast-setting, encapsulated form of MTA that is easy to use. There have been some studies on MM-MTA¹¹⁻¹³, but to the best of our knowledge only one report has detailed MM-MTA discoloration. In contrast to this study, Dettwiler *et al.*¹⁵ reported no significant color change in MM-MTA or other calcium silicate cements containing bismuth oxide. However, their results contradict many other studies. In most previous studies on MTA, the samples were set by being placed on moist pellets and then incubated at 37° C under 100% humidity for 24 hours, whereas Dettwiler *et al.*¹⁵ sealed the cavities with a dual-curing composite resin self-adhesive immediately after applying the MTA, i.e., without waiting for it to set. This immediate sealing of the MTA may have reduced oxidation of the bismuth oxide, thus preventing discoloration.

CONCLUSION

In this *in vitro* study, we found that, in terms of discoloration, it is safer to use zirconium oxide rather than bismuth oxide as a radiopacifier for MTAs. In particular, we observed more discoloration when using a nanoparticle-based MTA containing bismuth oxide.

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