Bacterial Leakage of Mineral Trioxide Aggregate, Calcium-Enriched Mixture and Biodentine as Furcation Perforation Repair Materials in Primary Molars

Nahid Ramazani a*, Parisa Sadeghi b

a Children and Adolescent Health Research Center, Oral and Dental Disease Research Center, Department of Pediatric Dentistry, School of Dentistry, Zahedan University of Medical Sciences, Zahedan, Iran; b Dentist, Zahedan University of Medical Sciences, Zahedan, Iran

ARTICLE INFO

Article Type: Original Article

Introduction: Adequate seal of iatrogenically perforated area within the root canal system can improve the long term treatment prognosis. This in vitro study evaluated the sealing ability of mineral trioxide aggregate (MTA), calcium-enriched mixture (CEM) cement and Biodentine in repair of furcation perforation in primary molars. Methods and Materials: A total of 61 freshly extracted primary mandibular second molars were randomly divided into three groups (n=17) and 10 teeth were put in negative (without perforation, n=5) and positive (perforated without repair, n=5) control groups. Turbidity was used as the criteria of bacterial leakage, when detected in the model of dual-chamber leakage. Data were analyzed using the Chi-Square and Kaplan-Meier survival analysis in SPSS software. The level of significance was set at 0.05. Results: All positive samples showed turbidity, whereas none of the negative samples allowed bacterial leakage. There was no significant difference between the number of turbidity samples in repaired teeth with all test materials (P=0.13). No significant difference was also detected in the mean survival time (P>0.05). Conclusion: CEM cement and Biodentine showed promising results as perforation repair materials and can be recommended as suitable alternatives of MTA for repair of furcation perforation of primary molars.

Keywords: Biodentine; Biomaterial; Calcium-Enriched Mixture; Furcation Perforation; Mineral Trioxide Aggregate; Perforation Repair; Sealability

Introduction

Dental caries is the most prevalent infectious disease during childhood [1]; most of which necessitate pulp treatment. The main objective of pulp therapy in primary teeth is to maintain the integrity of oral structures, guide permanent teeth to erupt properly and finally ensure general well-being of the child [2]. Because of the complexity of treatments and behavior challenges in children [3], procedural accidents such as perforation and creating an artificial opening in the furcation area are not uncommon. Perforation has been reported to occur in 2-12% of cases [4]. This procedural accident influences the prognosis of endodontic treatment [5, 6].

Repair of the perforation with an optimum sealing biomaterial has clinical significance in preventing the consequences and eventual tooth loss [6, 7]. In recent years, the sealing ability of the perforation repair material has been the subject of many investigations. Although it cannot be considered as the only criterion, providing adequate seal at the interface of material-dentin wall is one of the characteristics of ideal repair material [7]. Different biomaterials show different degree of sealing ability and biocements like mineral trioxide aggregate (MTA) and calcium-enriched mixture (CEM) cement have been successfully used for this purpose [5, 6, 8].

MTA has resulted in successful outcomes in furcation repair. Despite many good properties [9], MTA has long setting time
and difficult handling [5, 10-12]. These are important considerations for clinical application in pediatric patients. CEM is cement also provides good results when used for perforation repair [5, 10]. This cement has short setting time and offers good sealing ability and handling [5, 8, 10, 13, 14]. Biodentine is another popular biomaterial in endodontics with promising results. It is a new dentine substitute containing tricalcium silicate with good handling and mechanical properties. It has also good sealing ability and short setting time [4, 11, 12, 15-19]. In two separate studies, Hagghoo et al. [5, 8], found no significant difference between MTA and CEM cement as perforation repair materials in primary molars. There are also sparse papers that have evaluated the applications of Biodentine as a posterior restorative material [15], the capping agent in vital pulp therapies [18, 20, 21] and root end filling [17].

Given the serious implications of furcation perforation as well as sparse data comparing the sealing ability of MTA, CEM cement and Biodentine for repairing the perforated primary teeth, the purpose of this in vitro study was to compare the bacterial leakage of MTA, CEM cement and Biodentine in repairing the simulated furcation perforations in primary molars using dual-chamber bacterial leakage model.

Materials and Methods

Ethics Committee of Zahedan University of Medical Sciences approved the study protocol (Grant No: 7121). In this in vitro study, 61 freshly extracted primary mandibular second molars were used. The sample size was calculated based on previous similar studies [8, 22, 23] using Minitab statistical software. The inclusion criteria were as follows: normal furcation (with completely distinct roots), minimal caries (at least 4 mm caries included) and difficult handling [5, 10-12]. These are important considerations for clinical application in pediatric patients. CEM is cement also provides good results when used for perforation repair [5, 10]. This cement has short setting time and offers good sealing ability and handling [5, 8, 10, 13, 14]. Biodentine is another popular biomaterial in endodontics with promising results. It is a new dentine substitute containing tricalcium silicate with good handling and mechanical properties. It has also good sealing ability and short setting time [4, 11, 12, 15-19]. In two separate studies, Hagghoo et al. [5, 8], found no significant difference between MTA and CEM cement as perforation repair materials in primary molars. There are also sparse papers that have evaluated the applications of Biodentine as a posterior restorative material [15], the capping agent in vital pulp therapies [18, 20, 21] and root end filling [17].

In each tooth, the access cavity was prepared using a #5 diamond bur (D&Z Co., Wies Baden, Germany) mounted in a high speed water cooled handpiece. Cavity preparations with 2 mm depths were also made at root ends. Then, orifices and the prepared apical-end cavities of roots were filled with light-cured glass ionomer (Fuji II LC, GC Corporation, Tokyo, Japan). Except for the negative control group, the floor of pulp chamber was perforated using a #010 round bur (D&Z Co., Wies Baden, Germany) installed on a high speed handpiece with constant water spray. The size of perforation was the same as the bur size (1 mm in diameter) in all samples. The bur was replaced with a new one after making every six perforations.

The samples were randomly assigned into five groups (three experimental and two control groups). In groups I, II and III (n=17), perforations were sealed with either ProRoot MTA (Dentsply, Tulsa Dental, Tulsa, OK, USA), CEM cement (BioniqueDent, Tehran, Iran) or Biodentine (Septodont, Saint-Maur-des-Fosses, France), respectively. In the positive control group (n=5), no repairing material was used. In the negative control group (n=5), the perforation area was covered with two coats of nail varnish. After irrigation of samples with 10 mL normal saline, repairing materials were mixed according to the manufacturers’ instructions and placed by a carrier gun on the perforation site. Biomaterials were packed with moist cotton pellets while the samples were positioned in wet soft sponges. Condensing upon the sponge simulated the clinical condition in the oral cavity. At the end, all samples were placed in an incubator at 37°C and 100% humidity for 24 h to allow the biomaterials being fully set. Subsequently, teeth were coated by two layers of nail varnish except for the perforation site and nearly 1 mm around it.

In this experiment, a dual-chamber anaerobic bacterial leakage apparatus was used. The upper chamber was assembled by 3 mL plastic Eppendorf cylinder (Sigma-Aldrich Co., Hamburg, Germany) after cutting off 5 mm from its end. The samples were mounted in the cylinder so that the external surface of perforation area was left outside and accessible. The gaps between the sample and the inner side of cylinder were completely sealed with sticky wax. The apparatus was sterilized with ethylene oxide for 8 h. The upper chamber was inserted in 10 mL glass vial (Pouyan Teb Co., Tehran, Iran) as the lower chamber which previously was filled with 5 mL of sterile Phenol Red Broth (PRB, Merck, Darmstadt, Germany). The upper-lower chamber interface was tightly sealed with parafilm (Supa Co., Tehran, Iran). It was checked that the perforation area was immersed in PRB. The whole assembling was incubated at 37°C in 100% humidity for 3 days. An amount of 9x10⁸ CFU/mL of Enterococcus faecalis (E. faecalis) (PTCC1778) compatible with 0.5 McFarland standard was used to inoculate 2 mm of PRB. This bacterial suspension was added to the upper chamber every two days. The vials glasses were daily observed for turbidity (red to yellow color conversion) as the indicator of bacterial growth throughout 90 days of experiment.

In this study, all procedures were done by the same experienced practitioner. Data were recorded for each experimental or control samples and finally analyzed using the Chi-Square and Kaplan-Meier survival analysis tests with SPSS software (SPSS version 18.0, SPSS, Chicago, IL, USA). The level of significance was set at 0.05.
Results

During the entire observational period, control samples behaved as were expected. All positive samples exhibited red to yellow color changes. Whiles, no color conversion was recorded with the negative control group.

The majority of MTA samples remained without turbidity throughout the monitoring period. On the 34th day of apparatus assembling, color conversion of one sample in MTA group was detected. There was another sample with color change recorded by day 48 in this group.

Turbidity did not occur until day 27th in CEM samples. During the experimental period, three additional samples from CEM group exhibited bacterial contamination on day 40. Two of the remaining samples showed turbidity on day 49th. The first two samples of Biodentine group showed leakage observed by day 23. The other turbidity samples were added by day 25 (one sample), day 48 (two samples) and day 62 (two samples).

Throughout the experiment, the turbidity results remained unchanged from day 62 to the end. Totally, as presented in Table 1, MTA showed the less number of turbidity followed by CEM cement and Biodentine, respectively. However, the Chi-Square test failed to detect a statistical significant difference among three experimental groups (P=0.13). The means of survival time in different groups are shown in Table 2. According to the Kaplan-Meier survival analysis, there was no significant difference in mean survival time of study groups (P>0.05).

Discussion

The focus point of the current study (comparison of MTA, CEM cement and Biodentine based on the sealing ability as perforation repair materials) have not been explored in earlier studies. Totally, 2, 6 and 7 samples of turbidity were recorded with MTA, CEM cement and Biodentine, respectively. Although, based on statistical tests, throughout the 90-day experiment, perforation repair material had no significant effect on the microbial leakage.

Different techniques have been proposed to assess the sealing ability of various perforation repair materials [9, 24, 25]. Microleakage dye penetration model is one of the traditional methods with advantages such as easy manipulation and inexpensiveness [16, 26]. However, chemical characteristics, pH and low molecular size of dye and it’s dissolution by repairing material may affect the depth of dye penetration and cause leakage to be over- or under-estimated [8, 13, 26]. Moreover, it has to be noted that no more than one plan of dye penetration can be detected [26]. These make the above mentioned method not clinically relevant and it is logical to adopt a new standard, valid and reliable model. So in the current study, microleakage analysis method using a dual-chamber microleakage apparatus utilizing E. faecalis, was applied. This method is an improvement over microleakage dye penetration model. By using this assembling, the clinical bacterial contamination can be simulated [13, 27]. E. faecalis was chosen because it is the commonly detected microorganism in post endodontic failure [27]. This gram positive, facultative, anaerobic organism has the ability to persist with inadequate source of nutrition and can invade the dentinal tubules [5, 9, 28, 29].

Several studies have reported the ability of MTA to prevent leakage in a variety of applications [8, 14, 20, 24, 30]. Moreover, they reported its superiority compared to other dental materials. For this reason, we included MTA as a standard perforation repair material for better comparison. This material is routinely used to repair perforation defects mainly due to its moist compatibility [31-33]. In the study by Sahebi et al. [6], regarding the sealing ability of different materials, significantly more microleakage of MTA was reported compared to CEM cement. However, the present research found no detectable difference between MTA and CEM cement in this regard. The differences in methodology can be the reason that makes direct comparison difficult. Perhaps the techniques used to evaluate microleakage and the type of tooth material (primary versus permanent teeth) may be related to the different findings.

In another in vitro study CEM cement exhibited no significant difference from MTA as root-end filling and sealing materials [25]. Although in that study a method other than dual-chamber apparatus was used. However, in one recent study by Zarenejad et al. [14], CEM cement and MTA showed similar behaviors when applied as intra-orifice barrier during nonvital bleaching.

| Table 1. | Turbidity in different experimental groups and the time of detection and number of samples added |
|---------|--------------------------------------------------------------------------------------------------|
| Group   | Turbidity | Day (N)   |
|---------|-----------|-----------|
| MTA     | 2         | 34 (1)    |
|         |           | 48 (1)    |
| CEM cement | 6     | 27 (1)    |
|         |           | 40 (3)    |
|         |           | 49 (2)    |
| Biodentine | 7     | 23 (2)    |
|         |           | 25 (1)    |
|         |           | 48 (2)    |
|         |           | 62 (2)    |

| Table 2. | Mean (SD) of survival time in different experimental groups [Confidence Interval (CI) =95%] |
|---------|--------------------------------------------------------------------------------------------------|
| Group   | Mean (SD)                                        |
|---------|--------------------------------------------------|
| MTA     | 84.235 (3.873)                                   |
| CEM cement | 72.647 (5.797)                                   |
| Biodentine | 70.059 (6.319)                                   |
| Total   | 75.647 (3.253)                                   |
The findings of the current study confirm those of two previous investigations by Haghgoo et al. [5, 8] who evaluated the sealing ability of CEM cement and MTA by using bacterial leakage and dye penetration models in primary teeth. They concluded that the two tested biomaterials demonstrate similar capacities as furcation perforation materials [5, 8].

The results of the study by Shahi et al. [26], using protein leakage model in permanent teeth, confirms the findings of the present study regarding the good sealing ability of MTA as perforation repair material. In addition, in another study by Samiee et al. [34], MTA and CEM allowed similar resistance to leakage at the material-dentinal wall interface of repaired perforations. The results obtained from that study is consistent with the result yielded in the present investigation. CEM cement when compared to MTA as driven from different studies [13, 35, 36], and the current one revealed no detectable difference or even statistically superior performance. Considering the result obtained together with the benefits such as short setting time, low toxicity and low price [5, 27, 29, 37] CEM cement can be proposed as potential substitute of MTA.

Different applications have been proposed for Biodentine [15, 17, 18, 20, 21, 38]. Some studies investigated the performance of this new dentine substitute in restoration of posterior teeth [15, 16]. Koubi et al. [16], used glucose diffusion microleakage method and found that the material performed as well as resin modified glass ionomer. Additionally, less marginal discoloration and good handling has been attributed to Biodentine [15]. In one study, it was showed that when Biodentine was used as root-end filling material, significantly better marginal adaptation was observed compared to MTA [17]. However, according to Soundappan et al. [12], Biodentine could not compete with MTA as root end filling material. Considering the biocompatible entity of Biodentine and its ability to induce odontoblast differentiation the bacterial leakage resistance of this calcium-silicate cement after repair of perforation must be assessed. Given the good properties of Biodentine [11, 20] together with our findings addition of this cement to the list of primary tooth perforation repair materials is crucial.

MTA, CEM cement and Biodentine participate in hydroxyl apatite formation at the material-dentine interface [14, 39]. After adding liquid to powder, these three formulations form small sized non structured hydrate gels, which may flow to better accessing gaps and spread and fit into the dentinal tubules by wetting of dentin surface which in turn prevents bacterial leakage. Moreover, they exhibited a slight post-setting expansion [5, 12, 15]. The amount of bacterial leakage is proportional to the size of perforation [16]. In order to achieve a valuable comparison, all perforation defects were made similarly and with the same size.

Despite the promising results regarding the sealing ability of CEM cement and Biodentine, it should be kept in mind that in vitro studies, due to many inherent drawbacks cannot simulate oral condition completely. On the other hand, because of no expression of full clinical characteristic of the repairing material under in vitro conditions, the long term prognosis of perforation sealed teeth are unknown. So future clinical studies on accidentally perforated primary molars are recommended to evaluate the long term prognosis.

**Conclusion**

Within the limitation of the present study, CEM cement and Biodentine had no notable difference compared to MTA in terms of in vitro bacterial leakage.

**Acknowledgment**

The authors wish to thank the Deputy of Research and Technology of Zahedan University of Medical Sciences for their support.

Conflict of Interest: 'None declared'.

**References**

1. Ramazani N. Child dental neglect: a short review. Int J High Risk Behav Addict. 2014;3(4):e21861.
2. Guideline on pulp therapy for primary and young permanent teeth. Pediatr Dent. 2008;30(7 Suppl):170-4.
3. Ramazani N, Mohammadi A, Amirabadi F, Ramazani M, Ehsani F. In vitro investigation of the cleaning efficacy, shaping ability, preparation time and file deformation of continuous rotary, reciprocating rotary and manual instrumentations in primary molars. J Dent Res Dent Clin Dent Prospects. 2016;10(1):49-56.
4. Kakani AK, Veeramachaneni C, Majeti C, Tummala M, Khiyani L. A Review on Perforation Repair Materials. J Clin Diagn Res. 2015;9(9):Ze09-13.
5. Haghgoo R, Niyakan M, Nazari Moghaddam K, Asgary S, Mostafaloo N. An In vitro Comparison of Furcal Perforation Repaired with Pro-root MTA and New Endodontic Cement in Primary Molar Teeth- A Microleakage Study. J Dent (Shiraz). 2014;15(1):28-32.
6. Sahebi S, Moazami F, Sadat Shojae N, Layeghehghad M. Comparison of MTA and CEM Cement Microleakage in Repairing Furcal Perforation, an In Vitro Study. J Dent (Shiraz). 2013;14(1):31-6.
7. Aggarwal V, Singla M, Miglani S, Kohli S. Comparative evaluation of push-out bond strength of ProRoot MTA, Biocement, and MTA Plus in furcation perforation repair. J Conserv Dent. 2013;16(5):462-5.
8. Haghgoo R, Arfa S, Asgary S. Microleakage of CEM Cement and ProRoot MTA as Furcal Perforation Repair Materials in Primary Teeth. Iran Endod J. 2013;8(4):187-90.
9. Samiei M, Aghazade M, Farhadi F, Shahveghar N, Torab A, Vahid Pakdel SM. Sealing Efficacy of Single-cone Obturation Technique with MTA and CEM Cement: An in Vitro Bacterial Leakage Study. J Dent Res Dent Clin Dent Prospects. 2014;8(2):77-83.

10. Adl A, Sobhnamayan F, Kazemi O. Comparison of push-out bond strength of mineral trioxide aggregate and calcium enriched mixture cement as root end filling materials. Dent Res J (Isfahan). 2014;11(5):564-7.

11. Jang YE, Lee BN, Koh JT, Park YJ, Joo NE, Chang HS, Hwang IN, Oh WM, Hwang YC. Cytotoxicity and physical properties of tricalcium silicate-based endodontic materials. Restor Dent Endod. 2014;39(2):89-94.

12. Soundappan S, Sundaramurthy JL, Raghu S, Natanasabapathy V. Biodentine versus Mineral Trioxide Aggregate versus Intermediate Restorative Material for Retrograde Root End Filling: An Invitro Study. J Dent (Tehran). 2014;11(2):143-9.

13. Tabrizizade M, Asadi Y, Sooratgar A, Moradi S, Sooratgar H, Afshar S, Sayedi M. Sealing ability of mineral trioxide aggregate and calcium-enriched mixture cement as apical barriers with different obturation techniques. Iran Endod J. 2014;9(4):261-5.

14. Zarenejad M, Asgary S, Ramazani N, Haghshenas MR, Rafiee M, Ramazani M. Coronal microleakage of three different dental biomaterials as intra-orifice barrier during nonvital bleaching. Dent Res J (Isfahan). 2015;12(6):581-8.

15. Koubi G, Colon P, Franquic JM, Hartmann A, Richard G, Faure MO, Lambert G. Clinical evaluation of the performance and safety of a new dentine substitute, Biodentine, in the restoration of posterior teeth - a prospective study. Clin Oral Investig. 2013;17(1):243-9.

16. Koubi S, Elmerini H, Koubi G, Tassery H, Camps J. Quantitative evaluation by glucose diffusion of microleakage in aged calcium silicate-based open-sandwich restorations. Int J Dent. 2012;2012:105863.

17. P VR, Vemisetty H, K D, Reddy SJ, D R, Krishna MJ, Malathi G. Comparative Evaluation of Marginal Adaptation of Biodentine(TM) and Other Commonly Used Root End Filling Materials- An Invitro Study. J Clin Diagn Res. 2014;8(3):243-5.

18. Villat C, Grosogeat B, Seux D, Farge P. Conservative approach of a symptomatic carious immature permanent tooth using a tricalcium silicate cement (Biodentine): a case report. Restor Dent Endod. 2013;38(4):258-62.

19. Daryaeian M, Nazarimoghadam K, Ramazani N. Comparison of pH changes in external simulated root defects after insertion of Biodentine and Modified Portland Cement in intracanal space. J Mash Dent Sch. 2015;40(1):37-46.

20. Niranjani K, Prasad MG, Vasa AA, Divya G, Thakur MS, Saujanya K. Clinical Evaluation of Success of Primary Teeth Pulpotomy Using Mineral Trioxide Aggregate((R)), Laser and Biodentine(TM)– an In Vivo Study. J Clin Diagn Res. 2015;9(4):Zc35-7.

21. Kusum B, Rakesh K, Richa K. Clinical and radiographical evaluation of mineral trioxide aggregate, biodentine and propolis as pulpotomy medicaments in primary teeth. Restor Dent Endod. 2015;40(4):276-85.

22. Ferris DM, Baumgartner JC. Perforation repair comparing two types of mineral trioxide aggregate. J Endod. 2004;30(6):422-4.

23. Singh P, Paul J, Al-Khuraiti AA, Vellappally S, Halawany HS, Hashim M, Abraham NB, Jacob V, Thavarajah R. Sealing ability of mineral trioxide aggregate, calcium phosphate cement, and glass ionomer cement in the repair of perforation perforations. Acta Medica (Hradec Kralove). 2013;56(3):97-103.

24. Tselnik M, Baumgartner JC, Marshall JG. Bacterial leakage with mineral trioxide aggregate or a resin-modified glass ionomer used as a coronal barrier. J Endod. 2004;30(11):782-4.

25. Asgary S, Eghbal MJ, Parirrokh M, Torabzadeh H. Sealing ability of three commercial mineral trioxide aggregates and an experimental root-end filling material. Iran Endod J. 2006;1(3):101-5.

26. Shahi S, Rahimi S, Hasan M, Shiezadeh V, Abdolrahimi M. Sealing ability of mineral trioxide aggregate and Portland cement for furcal perforation repair: a protein leakage study. J Oral Sci. 2009;51(4):601-6.

27. Moradi S, Dusfani R, Ghazvini K, Lomee M. Sealing Ability of Orthoapge MTA and CEM Cement in Apically Resected Roots Using Bacterial Leakage Method. Iran Endod J. 2013;8(3):109-13.

28. Razmi H, Aminsoobhani M, Bolhari B, Shamshirgar F, Shahsavas S, Shamshiri AR. Calcium Enriched Mixture and Mineral Trioxide Aggregate Activities against Enterococcus Faecalis in Presence of Dentin. Iran Endod J. 2013;8(4):191-6.

29. Bidar M, Naderinasab M, Talati A, Ghazvini K, Asgari S, Hadizadeh B, Gharechahi M, Mashadi NA. The effects of different concentrations of chlorhexidine gluconate on the antimicrobial properties of mineral trioxide aggregate and calcium enrich mixture. Dent Res J (Isfahan). 2012;9(4):466-71.

30. Chang SW. Chemical characteristics of mineral trioxide aggregate and its hydration reaction. Restor Dent Endod. 2012;37(4):188-93.

31. Kerner S, Bronneck F. Conservative treatment of a large facial midroot perforation. Case Rep Dent. 2015;2015:326302.

32. Haghgoo R, Abbasi F. Treatment of Furcal Perforation of Primary Molars with ProRoot MTA versus Root MTA: A Laboratory Study. Iran Endod J. 2013;8(2):52-4.

33. Nagpal R, Manuja N, Pandit IK, Rallan M. Surgical management of iatrogenic perforation in maxillary central incisor using mineral trioxide aggregate. BMJ Case Rep. 2013;2013.

34. Samiei M, Eghbal MJ, Parirrokh M, Abbas FM, Asgary S. Repair of furcal perforation using a new endodontic cement. Clin Oral Invest. 2010;14(6):653-8.

35. Milani AS, Shakouie S, Borna Z, Sighar Deljavan A, Asghari Jafarabadi M, Pournaghi Azar F. Evaluating the Effect of Resection on the Sealing Ability of MTA and CEM Cement. Iran Endod J. 2012;7(3):134-8.

36. Hasheminia M, Loriaei Niai S, Asgary S. Sealing Ability of MTA and CEM Cement as Root-End Fillings of Human Teeth in Dry, Saliva or Blood-Contaminated Conditions. Iran Endod J. 2010;5(4):151-6.

37. Mozamati F, Salehi S, Jamshidi D, Alavi A. The long-term effect of calcium hydroxide, calcium-enriched mixture cement and mineral trioxide aggregate on dentin strength. Iran Endod J. 2014;9(3):185-9.

38. Nayak G, Hasan MF. Biodentine-a novel dentinal substitute for single visit apexification. Restor Dent Endod. 2014;39(2):120-5.

39. Arman M, Khalilak Z, Rajabi M, Esnaashari E, Saati K. In Vitro Spectrophotometry of Tooth Discoloration Induced by Tooth-Colored Mineral Trioxide Aggregate and Calcium-Enriched Mixture Cement. Iran Endod J. 2015;10(4):226-30.

Please cite this paper as: Ramazani N, Sadeghi P. Bacterial Leakage of Mineral Trioxide Aggregate, Calcium-Enriched Mixture and Biodentine as Furcation Perforation Repair Materials in Primary Molars. Iran Endod J. 2016;11(3):214-8. doi: 10.7508/iej.2016.03.013.