

## Review Article

# Reparative Dentinogenesis Induced by Mineral Trioxide Aggregate: A Review from the Biological and Physicochemical Points of View

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This paper aims to review the biological and physicochemical properties of mineral trioxide aggregate (MTA) with respect to its ability to induce reparative dentinogenesis, which involves complex cellular and molecular events leading to hard-tissue repair by newly differentiated odontoblast-like cells. Compared with that of calcium hydroxide-based materials, MTA is more efficient at inducing reparative dentinogenesis *in vivo*. The available literature suggests that the action of MTA is attributable to the natural wound healing process of exposed pulps, although MTA can stimulate hard-tissue-forming cells to induce matrix formation and mineralization *in vitro*. Physicochemical analyses have revealed that MTA not only acts as a “calcium hydroxide-releasing” material, but also interacts with phosphate-containing fluids to form apatite precipitates. MTA also shows better sealing ability and structural stability, but less potent antimicrobial activity compared with that of calcium hydroxide. The clinical outcome of direct pulp capping and pulpotomy with MTA appears quite favorable, although the number of controlled prospective studies is still limited. Attempts are being conducted to improve the properties of MTA by the addition of setting accelerators and the development of new calcium silicate-based materials.

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## 1. Introduction

The aim of direct pulp capping is to maintain the vitality and function of the dental pulp following its exposure to the external environment. Although calcium hydroxide-based materials have been extensively used for this procedure because of their potential to induce hard-tissue repair and subsequent dentin bridge formation [1], mineral trioxide aggregate (MTA) has recently received much attention as a good substitute for calcium hydroxide-based materials and has demonstrated promising clinical outcomes (reviewed in [2]).

MTA is a bioactive material that was developed in the early 1990s, originally as a retrograde filling material, and first appeared in the dental scientific literature in 1993 [3]. Since then, the indications for MTA have expanded significantly. Currently, MTA is used to seal off exposed pulps

and various communications between the root canal system and surrounding tissues for a variety of indications such as root-end filling, perforation repair, and apexification [4]. MTA is a modified preparation of Portland cement [5–8], which is the basic ingredient of concrete and mortar that may never have been used as a dental material before the development of MTA. Currently, two different preparations of MTA are available: the original preparation is grey-colored (GMTA); whereas, a white preparation (WMTA) was recently introduced to address esthetic concerns.

A large number of studies have disclosed that MTA shows favorable biocompatibility and has physical properties suitable for dental application such as good sealing ability. However, the basic question of why such materials induce the hard-tissue repair of exposed pulps has not yet fully been answered. Thus, the purpose of this paper is to summarize the biological process of pulp tissue repair and then review

the available literature regarding the ability of MTA to induce reparative dentinogenesis from both the biological and physicochemical points of view.

## 2. Pulp Wound Healing and Tissue Repair

Dental pulp possesses a natural tissue repair potential, which leads to the formation of reparative dentin. It has been well documented that dental pulp possesses the ability to form a hard-tissue barrier (dentin bridge) after direct pulp capping or pulpotomy. During reparative dentinogenesis, the original odontoblasts at the exposure site are destroyed and replaced by newly differentiated odontoblast-like cells [1, 9–11]. Pulpal wound healing involves stem/progenitor cells migration to the injured site and their subsequent proliferation and differentiation into odontoblast-like cells. Reparative dentinogenesis is often initiated by the formation of a fibrodentin matrix, which is atubular and/or irregular and is associated with cuboidal cells. The formation of a tubular dentin-like matrix by elongated and polarized odontoblast-like cells takes place later (Figure 1).

Despite extensive studies, the molecular signaling involved in cell differentiation during reparative dentinogenesis has still not been fully characterized. During tooth development, odontoblast differentiation is controlled by specific basement membrane-mediated epithelial-mesenchymal interactions [12–14]. Fibronectin, an extracellular matrix glycoprotein found in association with the dental basement membrane, appears to play a crucial role in the terminal differentiation of odontoblasts [15, 16]. On the other hand, during reparative dentinogenesis when the basement membrane or dental epithelium is absent, the adhesion of progenitor cells to an appropriate surface (scaffold) may be a critical requirement for the differentiation of hard-tissue-forming cells [17]. When calcium hydroxide is applied to the exposed pulp tissue, a layer of dystrophic calcification associated with cellular degeneration may be the surface to which the pulp cells migrate and attach and where they subsequently differentiate into odontoblast-like cells. This process is considered to be mediated by the fibronectin deposited on this layer, which is structurally comparable to the basement membrane [18]. The formation of the tubular dentin-like matrix is preceded by the deposition of fibrodentin, which contains fibronectin [18]. The fibrodentin may be comparable to the mantle dentin observed in developing teeth and has been suggested to play an important role in the terminal differentiation of odontoblasts [13]. Recently, calcium ions released from calcium hydroxide have been shown to stimulate fibronectin gene expression in dental pulp cells [19].

During reparative dentinogenesis after pulp capping, bone sialoprotein (BSP) and osteopontin (OPN) have been detected at the exposure site and in the fibrous matrix (fibrodentin), but not in the tubular dentin-like matrix [20, 21] (Figure 1), while odontoblast-like cells were shown to express dentin sialoprotein [22, 23]. BSP and OPN are suggested to control the mineralization process [24].

These noncollagenous proteins may be associated with the initially formed calcified layer underneath the superficial necrotic zone. In addition, OPN is implicated in diverse biological events, including wound healing [25]. Interestingly, OPN gene expression in human dental pulp cells is enhanced by fibronectin [19]. The colocalization of fibronectin and OPN at the pulp exposure site suggests their role in the migration of progenitors and their differentiation into odontoblast-like cells during reparative dentinogenesis.

Transforming growth factor- $\beta$  (TGF- $\beta$ ) and other members of this family of growth factors have been implicated in tooth development and dental tissue repair [26]. TGF- $\beta$  isoforms and their receptors have been identified in odontoblasts of healthy and carious teeth [27, 28], and the TGF- $\beta$  type I receptor was identified in an animal model of pulp capping [20]. TGF- $\beta$ 1 has been demonstrated to bind to immobilized fibronectin [29]. Extracellular matrices like proteoglycans have been suggested to bind to many growth factors and modulate their activities [30]. During reparative processes after pulp capping, the fibronectin-rich matrix serves as a reservoir of growth factors as well as a substrate for cell migration and attachment. Such growth factors including TGF- $\beta$  and other inductive molecules expressed in the pulp tissue might be involved in odontoblast-like cell differentiation.

The derivation of the stem/progenitor cells during reparative dentinogenesis is still unclear. Pulp tissue contains a population of cells with stem-cell-like properties. Dental pulp stem cells have been found in human permanent teeth [31, 32] and exfoliated human deciduous teeth [33]. These stem cell populations have been suggested to reside in the microvasculature [34]. Hard tissue formation occurs in the pulp cavity after tooth replantation and transplantation, where dentin- and bone-like tissues are seen [35, 36]. Transplantation of a green fluorescent protein (GFP-) transgenic rat tooth into a wild-type rat socket demonstrated that bone-like tissue was formed by both host and donor cells. On the other hand, all cells lining the dentin-like matrix expressed GFP, suggesting that this matrix was formed by surviving donor odontoblasts and/or pulp cells capable of differentiating into odontoblast-like cells [37]. These results suggest that the dental pulp contains two types of progenitor cells capable of differentiating into either odontoblast- or osteoblast-like cells. This has been confirmed by allogenic tooth transplantation into the sublingual region using lacZ-transgenic ROSA26 mice [38]. Both types of progenitor cells might also be involved in reparative dentinogenesis. It is likely that the superficial layer and the fibrodentin of the newly formed dentin bridge matrix have osteogenic characteristics. The involvement of other cells including inflammatory cells from the bloodstream in pulpal wound healing is not ruled out.

Current studies are focusing on the derivation and phenotypes of the cells involved in nonspecific reparative dentinogenesis and the molecular mechanisms that regulate their cytodifferentiation. Such approaches will ultimately lead to regenerative therapy and tissue engineering of the dentin-pulp complex.

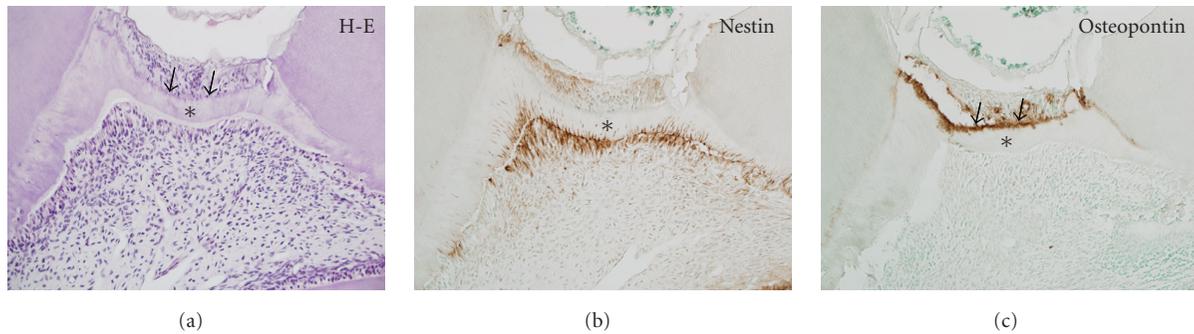


FIGURE 1: Dentin bridge formation in rat molar at 14 days after direct pulp capping with MTA: H-E staining (a), immunohistochemistry of nestin (b), and osteopontin (c). (a) A thin layer of fibrous matrix (arrows) is followed by a dentin-like matrix (\*) with tubular structures pulpally lined with odontoblast-like cells. (b) The odontoblast-like cells intensely express nestin, an intermediate filament expressed in differentiated odontoblasts. Their processes also show immunoreactivity for nestin in the tubular matrix (\*). (c) Osteopontin immunoreactivity is detected in the superficial fibrous matrix (arrows), but not in tubular dentin-like matrix (\*).

### 3. Cellular Reactions to MTA In Vitro

A number of in vitro studies have been conducted to evaluate the biocompatibility of MTA by measuring various parameters such as proliferation and viability using different types of cells in direct and/or indirect contact with MTA. Overall, the results suggest that the cytotoxicity of set MTA is less than that of traditional materials (reviewed by Camilleri and Pitt Ford [39] and by Roberts et al. [40]). For example, studies using cultured osteoblastic cells have demonstrated that MTA is less toxic than amalgam [41–43], Super EBA [42, 43], and intermediate restorative material (IRM) [41], since cells in contact with MTA showed higher viability and/or proliferative activity. However, MTA in its freshly mixed state shows a higher cytotoxicity [44, 45], which could be due to its high pH [46–49].

In vitro experiments have also demonstrated that MTA has the capacity to stimulate cell differentiation/activation, which may contribute to hard tissue matrix formation and/or mineralization. Incubation of gingival and periodontal ligament fibroblasts with MTA causes the induction of osteogenic phenotypes such as alkaline phosphatase, osteonectin, osteopontin, and osteonidgen [43]. MTA also stimulates the production of bone morphogenetic protein (BMP-) 2 and TGF- $\beta$ 1 from human gingival fibroblasts [50] and causes the upregulation of type I collagen and osteocalcin mRNA expression in an osteoblast-like cell line (MC3T3-E1) [51]. In a more recent study in which a cementoblast cell line was used, WMTA extracts at lower concentrations induced biomineralization of these cells and caused the upregulation of the mRNA expression of type I collagen and bone sialoprotein [52].

Recent studies using pulp cells have also suggested the capacity of MTA to stimulate matrix formation and mineralization during dentinogenesis. Cultured rat pulp cells stimulated with MTA through transwell inserts showed an increased mineralization and upregulation of BMP-2 mRNA and protein [53]. Thus, BMP-2 may be involved in MTA-induced mineralization. The MTA-induced mineralization together with the alkaline phosphatase activity and dentin

sialophosphoprotein (DSPP-) and BSP-expressions of cultured pulp cells was further enhanced when enamel matrix derivative (EMD) was added [54]. Thus, a combination of MTA and EMD may promote the differentiation of pulp cells more rapidly than MTA alone. In addition, rat clonal pulp cells stimulated with MTA upregulate cyclooxygenase-2 and inducible nitric oxide synthase mRNA expression via the nuclear factor kappa B signaling system [55]. Thus, prostaglandins and nitric oxide could be involved in various MTA-induced pulp tissue reactions including inflammation and hard tissue formation.

Overall, in vitro studies indicate that MTA is a biocompatible material and possesses the capacity to stimulate hard tissue-forming cells to induce matrix formation and mineralization. As will be discussed below, a significant part of the bioactivity of MTA in vitro may arise from its reactions with the surrounding environment, for example, the culture media and/or serum [56], which form biocompatible byproducts, including carbonated apatite [57, 58].

### 4. In Vivo Reparative Dentinogenesis by MTA

Subcutaneous and intraosseous implantation studies have consistently shown that MTA elicits less severe tissue reactions compared with those of traditional materials such as amalgam and Super EBA (reviewed by Camilleri and Pitt Ford [39] and Roberts et al. [40]). Moreover, subcutaneous implantation of MTA causes dystrophic calcification in the connective tissue adjacent to this material [59, 60].

The ability of MTA to induce reparative dentinogenesis or dentin bridge formation has been consistently demonstrated in animal studies in which direct pulp capping or pulpotomy was performed in mechanically exposed pulps [21, 61–70]. These studies have also shown that MTA causes limited pulp tissue necrosis shortly after its application. Thus, MTA seems less causative compared with calcium hydroxide, which is known to cause the formation of a necrotic layer along the material-pulp interface [1, 9–11]. Compared with calcium hydroxide, MTA induces reparative

dentin formation at a greater rate and a superior structural integrity [61, 62].

Also, the majority of studies in which MTA capping was carried out in mechanically pulp-exposed healthy human teeth showed that MTA provides higher frequencies of dentin bridge formation, a better quality (thickness, completeness, and/or integrity) dentin bridge, and milder pulp inflammation compared with calcium hydroxide-based materials [23, 71–78]. In one representative study involving 33 healthy third molars [75], direct pulp capping was performed with MTA or a hard-setting calcium hydroxide cement (Dycal) in 20 and 13 teeth, respectively, and histological, ultrastructural, and quantitative analyses were carried out after 1 week to 3 months. The MTA-capped pulps were mostly free from inflammation, and hard tissue bridges of steadily increasing length and thickness were formed. Dycal-capped pulps, however, showed the formation of less consistent barriers, which were frequently accompanied by tunnel defects, and pulp inflammation often persisted, even after 3 months. Moreover, a recent study has reported that MTA pulpotomy in 12 human permanent molars with irreversible pulpitis resulted in complete dentin bridge formation in all cases after 2 months [79].

On the other hand, the cellular and molecular events involved in MTA-induced reparative dentinogenesis have been addressed in a limited number of *in vivo* studies. In one study, early pulpal cell response after capping with GMTA was examined in mechanically exposed dog pulps [64]. GMTA initially induced the formation of a zone of crystalline structure and an arrangement of pulp cells with the morphological features of increased biosynthetic activity, for example, nuclear and cytoplasmic polarization and developed cytoplasmic organization. Then, the deposition of fibroblasts, followed by reparative dentin formation, which was characterized by the presence of polarized odontoblast-like cells and a tubular dentin-like matrix, was seen. Thus, the stereotypic pulp defense mechanism by which fibroblasts triggers the expression of the odontoblastic potential of pulp cells [10, 13] may be involved in MTA-induced reparative dentinogenesis. In another study, the reparative process of mechanically exposed rat molar pulps capped with WMTA was investigated by immunohistochemistry [21]. The reparative process involved initial deposition of osteopontin in the superficial layer of the pulpal matrix followed by increased cell proliferation and the appearance of nestin-immunoreactive newly differentiated odontoblast-like cells. Thus, the reparative dentinogenesis that occurs following MTA capping is primarily governed by the natural healing process of exposed pulps, which involves the proliferation and migration of progenitors followed by their differentiation into odontoblast-like cells. Osteopontin could play a triggering role in the initiation of this process. The expression of dentin sialoprotein, a noncollagenous protein expressed exclusively by odontoblasts, has been detected in newly differentiated odontoblast-like cells after direct pulp capping of human teeth with MTA [23]. Stronger dentin sialoprotein expression was observed in MTA-capped teeth than in Dycal-capped teeth, suggesting a superior dentinogenic effect of MTA.

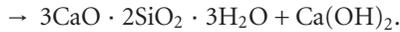
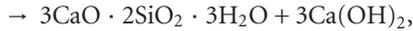
Based on the histologic investigations mentioned above, MTA appears superior to calcium hydroxide-based materials with regard to its capacity to stimulate reparative dentinogenesis, as far as mechanically exposed healthy pulps are concerned. MTA and calcium hydroxide share a mechanism of hard tissue formation that can be regarded as the natural healing process of exposed pulps. Thus, MTA should be regarded as “the current gold standard material”, for *in vivo* pulp capping experiments aimed at investigating the cellular and molecular mechanisms involved in nonspecific reparative dentinogenesis. Based on the *in vitro* capacity of MTA to stimulate hard tissue-forming cells, however, the possibility that MTA has specific actions for stimulating dentinogenesis cannot be ruled out.

## 5. MTA as a “Calcium Hydroxide-Releasing Material”

The major component of MTA is essentially a refined preparation of Portland cement, which is a mixture of dicalcium silicate ( $2\text{CaO}\cdot\text{SiO}_2$ ), tricalcium silicate ( $3\text{CaO}\cdot\text{SiO}_2$ ), tricalcium aluminate ( $3\text{CaO}\cdot\text{Al}_2\text{O}_3$ ), gypsum, and tetracalcium aluminoferrite ( $4\text{CaO}\cdot\text{Al}_2\text{O}_3\cdot\text{Fe}_2\text{O}_3$ ) [5–8]. Gypsum is added for setting retardation. Trace amounts of  $\text{SiO}_2$ ,  $\text{CaO}$ ,  $\text{MgO}$ ,  $\text{K}_2\text{SO}_4$ , and  $\text{Na}_2\text{SO}_4$  are also present [5–7]. The greatest difference in composition between MTA and Portland cement is that MTA contains bismuth oxide as a radiopacifier [5–8]. Moreover, the particles of MTA are more uniform and smaller than those of Portland cement [7]. MTA also contains fewer toxic heavy metals and has a longer working time [80, 81]. Thus, MTA has undergone modification and purification from Portland cement to make it more suitable for clinical use. The major compositional difference between the grey and white versions of MTA is that the levels of chromophores (mainly ferric compounds such as tetracalcium aluminoferrite) are greatly reduced in the white version.

The compressive strength of MTA gradually increases after initial setting and is comparable to those of the Super EBA and IRM cements after 21 days [82]. Thus, MTA possesses sufficient physical strength for endodontic use and is stronger than calcium hydroxide-based materials. Increasing the water-to-powder ratio causes increases in the porosity and solubility of MTA [46]. Too large a condensation pressure causes decreases in surface hardness and compressive strength [83].

The setting reaction of MTA is basically similar to that of Portland cement involving the hydration of anhydrous mineral oxide compounds via the dissolution of the compounds followed by the crystallization of hydrates [49, 84, 85]. During the hydration process of calcium silicate components, a portlandite phase, which is mainly composed of calcium hydroxide crystals, is produced together with less basic calcium silicate hydrate ( $3\text{CaO}\cdot 2\text{SiO}_2\cdot 3\text{H}_2\text{O}$ ) as follows [49, 84, 85]:



MTA is hydrophilic and requires moisture to set, which is a favorable property when there is potential for moisture contamination in the clinical setting; moisture from the surrounding tissue may assist the setting [82, 86]. Blood contamination has little impact on the degree of leakage [87]. One less than ideal property of MTA is that it is a slow-setting material like Portland cement. MTA requires approximately three hours for initial setting [7, 82], and the reaction continues slowly for weeks [82, 88, 89] and probably months.

Hydrated MTA is alkaline, and its pH rises from 10.5 to 12.5 three hours after mixing [6, 82]. This pH rise is due to the progression of calcium hydroxide formation during the hydration process. When set MTA is immersed in water, it shows solubility of less than 3% of weight loss in 24 hours [46, 81, 82, 90–92], which is lower than that of zinc oxide-eugenol cement [82]. Moreover, Ca ions are continuously released, and the medium maintains a high pH [46–49]. Such dissolution of calcium hydroxide may be a key mechanism behind the biological properties of MTA.

The dissolution of calcium hydroxide may negatively influence the physical properties of MTA. SEM analysis of water-immersed MTA revealed an increased porosity, which may have been caused by the dissolution of calcium hydroxide and other hydration products [46]. Water immersion of MTA results in the formation of a subsurface layer of low Ca concentration (Ca-leached layer) [93]. A porosity increase and the formation of a Ca-leached layer have also been reported for Portland cement [94, 95] and thus these properties are derived from the parent material. Nevertheless, MTA has advantages over traditional calcium hydroxide preparations in terms of its structural stability and sealing ability, since calcium hydroxide preparations show a high degree of dissolution and lack sealability [96–98]. In addition, Si and Al show an increased concentration within the Ca-leached layer [93], probably resulting from the formation and/or accumulation of insoluble components such as calcium silicate hydrate and ettringite. Thus, the dissolution process may not be one way, but rather involves a “self-reparative” mechanism that compensates for Ca dissolution.

## 6. Interaction of MTA with the Surrounding Environment: A Basis for Its Bioactivity

MTA and Portland cements are virtually devoid of phosphorus [5–7]. However, when these cements are immersed in phosphate-containing solutions such as phosphate-buffered saline (PBS), they interact with the medium and produce apatite crystals on their surfaces [57, 58, 99–102] (Figure 2). The portlandite phase is not detectable on the

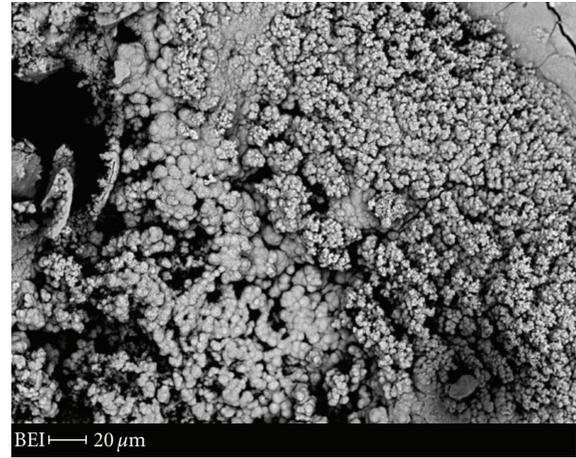


FIGURE 2: SEM photograph of the surface of white MTA immersed in PBS for 10 days, showing precipitates of various morphologies. Wavelength-dispersive X-ray spectroscopy analysis revealed that the precipitates contained Ca and P as their main elemental components.

upper surface of PBS-immersed Portland cement, but is found only in the interior [102]. This indicates that the calcium ions supplied by portlandite dissolution interact with the phosphate ions in the medium, allowing for the formation of apatite crystals. Such properties of apatite formation are considered to be important for explaining the biocompatibility and/or bioactivity of MTA, since the surface precipitation of biocompatible material(s) may be a basis for the bioactivity of several inorganic biomaterials [103, 104]. The precipitates are also formed at the MTA-dentin interface [57, 58, 99, 105], and thus may play a role in the achievement of a good marginal seal, as will be described below.

Sarkar et al. [99] first reported the formation of white precipitates with a globular ultrastructure on GMTA following immersion in PBS solution. Using X-ray diffraction (XRD) analysis, the authors identified the crystals as hydroxyapatite, although their calcium-to-phosphorus ratios were different from that reported for hydroxyapatite. This finding was confirmed by Bozeman et al. [100], who also used XRD and SEM and analyzed both WMTA and GMTA that had been subjected to PBS immersion. They concluded that the crystal precipitates on both MTA materials were chemically and structurally similar to hydroxyapatite. In addition, the authors also found that GMTA produces twice as many crystals as WMTA, suggesting that the two MTA materials do not possess the same level of bioactivity [100]. On the other hand, Tay et al. [57] analyzed the crystal precipitates on PBS-immersed white Portland cement with SEM, XRD, TEM, and Fourier transformation-infrared spectroscopy. They identified the precipitates as calcium-deficient poorly crystalline carbonated apatite, which had been transformed from an initially formed amorphous calcium phosphate. Most recently, Reyes-Carmona et al. [58] examined the precipitates on various PBS-immersed MTA preparations

and Portland cements by means of SEM and XRD and found the presence of amorphous calcium phosphate crystals of different morphologies and Ca/P ratios, which may act as precursors during the formation of carbonated apatite. The formation of crystals mainly composed of Ca and P has also been demonstrated on the pulpal surface of MTA that had been capped on exposed dog pulps *in vivo* [64].

Taken together, studies have consistently reported that MTA and Portland cement are able to interact with phosphate-containing fluids to form apatite deposits on their surfaces. This appears to be a common characteristic of calcium silicate-containing biomaterials [106, 107]. Since carbonated apatite represents the biological apatite phases found in bone, cementum, and dentin, this apatite layer may play a triggering role in the dentinogenic activity of MTA by supporting new tissue formation and its integration into dentin-like tissue.

## 7. Sealing Ability

A large number of studies have demonstrated that MTA has a better sealing ability than that of traditional materials such as amalgam, glass ionomer cement, and zinc oxide-eugenol cement by means of dye leakage, bacterial leakage, and fluid infiltration tests (reviewed by Roberts et al. [40]). However, the question of why MTA exhibits such a good seal has not yet fully been resolved.

MTA shows slight expansion upon setting [108–110], which may contribute, at least in part, to its good sealing ability. The marginal adaptation of MTA is in general better than that of traditional materials [111–114], although one study [114] reported that marginal adaptation did not correlate with leakage. WMTA exposed to a water-soluble dye before achieving full set showed poorer adaptation and more leakage compared with those of IRM and Super EBA cements [115].

GMTA-dentin bond failure usually occurs cohesively within the MTA material [116]. This indicates the presence of certain bonding mechanisms that may contribute to its sealing ability. One explanation for this mechanism is the above-mentioned ability of MTA to spontaneously produce apatite precipitates in the presence of phosphate-containing fluids [57, 58, 99, 100, 102], since materials with an apatite layer are known to form a chemical bond with calcified tissues such as bone [103, 107, 117]. Additionally, the formation of apatite-like materials that fill the MTA-dentin interfacial space has been demonstrated [58, 99]. The apatite was deposited within collagen fibrils, and the interfacial layer composed of apatite was accompanied with tag-like structures that extended into the dentinal tubules [58]. Taken together, the MTA-dentin interfacial layer formation that results from the capacity of MTA to induce spontaneous apatite formation may contribute to minimizing leakage not only by filling the gap along the interface but also via interactions with dentin such as intrafibrillar apatite deposition to promote mineral nucleation on dentin.

## 8. Antimicrobial Activity

Antimicrobial capacity due to high pH is considered one of the advantages of the calcium hydroxide-based materials used for direct pulp capping, since this procedure is often carried out on pulps that have already been bacterially contaminated and/or carry a potential risk of bacterial leakage along the restoration margins [96, 97, 118]. Various MTA preparations show antibacterial [119–125] and antifungal [124, 126–128] activities against different microbial strains. Similar to calcium hydroxide-based materials, the antimicrobial action of MTA is most likely associated with elevated pH resulting from ionization that releases hydroxyl ions. However, this activity of MTA is limited against some facultative bacteria and has no effect on strict anaerobic bacteria [119], and it is also weaker than the actions of zinc oxide-eugenol cements [119, 124]. Sealapex, a calcium hydroxide-based sealer, shows similar to better anti-microbial activity compared with those of various MTA preparations and/or Portland cements [121, 124, 125]. Taken together, the antimicrobial activity of MTA may not be as strong as those of traditional calcium hydroxide-based cements and sealers, although it does contribute to the reduction of bacterial contamination in pulpal wounds. The weaker antimicrobial activity of MTA may be compensated for by its good sealing ability.

## 9. Clinical Performance of MTA as a Pulp Capping Medicament and Pulpotomy Dressing

Two studies have investigated the clinical performance of MTA applied to the pulp capping of cariously exposed permanent teeth and reported success rates of 93% [129] and 98% [130]. In one study [130], 53 teeth with carious exposures diagnosed as reversible pulpitis were capped with MTA. Forty-nine out of 53 teeth were recalled in a mean period of 3.94 years, and 98% of the cases presented a normal radiographic appearance, no symptoms, and a normal response to cold testing. Regarding primary teeth, one study evaluated the clinical outcome of direct pulp capping with either WMTA or Dycal in 25 symmetrical pairs of carious primary molars [131]. None of the teeth exhibited clinical or radiographical failure during the follow-up period of up to 24 months.

Four studies regarding MTA pulpotomies in cariously exposed permanent teeth have reported high success rates ranging from 93%–100% [132–135]. In one prospective study, the success rates of partial pulpotomies using either GMTA or calcium hydroxide were compared with a mean follow-up period of 34.8 months [135]. Fifty-one teeth in 34 patients were available for recall, and there was no statistically significant difference in the success rate between GMTA and calcium hydroxide (93% and 91%, resp.). In addition, a case report in which partial pulpotomies were conducted on 2 cases of dens evaginatus and histologic examination was conducted after 6 months; complete dentin bridge formation without pulp inflammation was demonstrated [136].

MTA is also used as a pulpotomy dressing for primary teeth and is considered an appropriate alternative to formocresol, since studies comparing MTA and formocresol consistently showed that MTA gave similar to better results both clinically and radiographically [137–143].

Overall, the clinical outcome of direct pulp capping and pulpotomy with MTA seems quite favorable, although the number of controlled prospective studies is still limited.

## 10. Modification of MTA: A Research Trend

The handling properties of MTA are recognized to be less than ideal, since the working time is limited to a few minutes—even though this slow-setting material requires approximately three hours for initial setting [7, 82]—and the cement mixture is somewhat grainy and sandy. Thus, attempts were made to improve these drawbacks by using additives to accelerate setting. Calcium chloride (2% to 15%) has been widely studied as a setting accelerator: it reduces the setting time [144–147], increases the sealing ability [148], and maintains a high pH [144, 147]. Moreover, the addition of calcium chloride did not affect the formation of dentin bridges following pulpotomies in dog teeth [70], and thus may not deteriorate the biologic properties of MTA. However, calcium chloride reduces the compressive strength of set MTA [144]. One study recommended an admix of 1% methylcellulose and 2% calcium chloride because it improved the handling properties of MTA without reducing its compressive strength [145].  $\text{Na}_2\text{HPO}_4$  also reduces the setting time [149] while maintaining biocompatibility in vitro [150]. A recent report has documented that the addition of  $\text{Na}_2\text{HPO}_4$  to WMTA creates a more biocompatible material, as demonstrated by subcutaneous implantation [60]. However, MTA with the addition of resinous components to allow light curing did not stimulate mineralization when implanted into rat connective tissues [151].

Attempts have also been made to improve the working (and physical) properties of MTA by developing new calcium silicate-based materials [108, 152–157]. Among these, NEC (new endodontic cement) is a novel endodontic material consisting of different calcium compounds (i.e., calcium oxide, calcium phosphate, calcium carbonate, calcium silicate, calcium sulfate, calcium hydroxide, and calcium chloride) [155–157]. NEC is reported to show a shorter setting time [155], better handling properties, and a similar sealing ability [157] compared with those of MTA. When applied to exposed dog dental pulps, both NEC and MTA show favorable responses characterized by the formation of dentin bridges, while Dycal showed inferior responses accompanied by incomplete dentin bridge formation and pulp inflammation [156].

## 11. Conclusions

The available literature suggests that MTA is more efficient at inducing reparative dentinogenesis in vivo compared with calcium hydroxide-based materials. However, MTA and

calcium hydroxide share several biological properties that contribute to the induction of reparative dentinogenesis, mostly due to the fact that set MTA acts as a “calcium hydroxide-releasing material.” Thus, the dentinogenic mechanism of MTA may be attributable to the natural wound healing process of exposed pulps, which is considered to be the mechanism involved in calcium hydroxide-induced reparative dentinogenesis. Nevertheless, in vitro studies suggest the presence of dentinogenic mechanisms specific to MTA, since MTA can stimulate hard tissue-forming cells to induce matrix formation and mineralization. MTA has several beneficial physical properties over calcium hydroxide, including a good sealing ability, a lower degree of dissolution, and a higher structural stability. MTA also has the ability to interact with phosphate-containing fluids to spontaneously form apatite precipitates, which not only explains its biocompatibility and bioactivity but may also contribute to its sealing ability. Thus, the capacity of MTA to induce hard tissue repair of exposed pulps may depend heavily on its ability to create a local environment in which the inherent wound healing capacity of the pulp is not deteriorated.

The clinical outcome of direct pulp capping with MTA seems quite favorable, although the number of controlled prospective studies is still limited. Attempts are being conducted to improve the working properties of MTA via the addition of setting accelerators and the development of new calcium silicate-based materials.

## References

- [1] U. Schröder, “Effects of calcium hydroxide-containing pulp-capping agents on pulp cell migration, proliferation, and differentiation,” *Journal of Dental Research*, vol. 64, pp. 541–548, 1985.
- [2] D. E. Witherspoon, “Vital pulp therapy with new materials: new directions and treatment perspectives—permanent teeth,” *Journal of Endodontics*, vol. 34, no. 7, supplement, pp. S25–S28, 2008.
- [3] S.-J. Lee, M. Monsef, and M. Torabinejad, “Sealing ability of a mineral trioxide aggregate for repair of lateral root perforations,” *Journal of Endodontics*, vol. 19, no. 11, pp. 541–544, 1993.
- [4] M. Torabinejad and N. Chivian, “Clinical applications of mineral trioxide aggregate,” *Journal of Endodontics*, vol. 25, no. 3, pp. 197–205, 1999.
- [5] N. K. Sarkar, R. Caicedo, P. Ritwik, R. Moiseyeva, and I. Kawashima, “Physicochemical basis of the biologic properties of mineral trioxide aggregate,” *Journal of Endodontics*, vol. 31, no. 2, pp. 97–100, 2005.
- [6] J. Camilleri, F. E. Montesin, K. Brady, R. Sweeney, R. V. Curtis, and T. R. P. Ford, “The constitution of mineral trioxide aggregate,” *Dental Materials*, vol. 21, no. 4, pp. 297–303, 2005.
- [7] T. Dammaschke, H. U. V. Gerth, H. Züchner, and E. Schäfer, “Chemical and physical surface and bulk material characterization of white ProRoot MTA and two Portland cements,” *Dental Materials*, vol. 21, no. 8, pp. 731–738, 2005.
- [8] S. Asgary, M. Parirokh, M. J. Eghbal, and F. Brink, “Chemical differences between white and gray mineral trioxide aggregate,” *Journal of Endodontics*, vol. 31, no. 2, pp. 101–103, 2005.

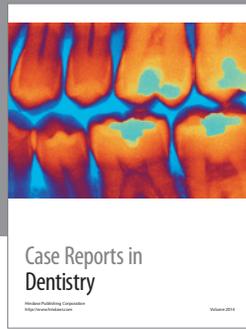
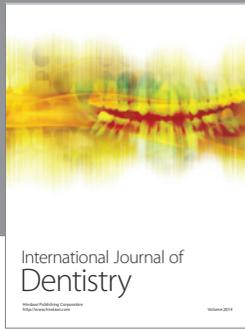
- [9] T. Yamamura, "Differentiation of pulpal cells and inductive influences of various matrices with reference to pulpal wound healing," *Journal of Dental Research*, vol. 64, pp. 530–540, 1985.
- [10] D. Tziafas, "Mechanisms controlling secondary initiation of dentinogenesis: a review," *International Endodontic Journal*, vol. 27, no. 2, pp. 61–74, 1994.
- [11] M. Goldberg and A. J. Smith, "Cells and extracellular matrices of dentin and pulp: a biological basis for repair and tissue engineering," *Critical Reviews in Oral Biology & Medicine*, vol. 15, no. 1, pp. 13–27, 2004.
- [12] I. Thesleff and K. Hurmerinta, "Tissue interactions in tooth development," *Differentiation*, vol. 18, no. 2, pp. 75–88, 1981.
- [13] J. V. Ruch, "Odontoblast differentiation and the formation of the odontoblast layer," *Journal of Dental Research*, vol. 64, no. 9, pp. 489–498, 1985.
- [14] J. V. Ruch, "Odontoblasts: developmental aspects," in *Dynamic Aspects of Dental Pulp*, R. Inoki, T. Kudo, and L. Olgart, Eds., pp. 29–50, Chapman and Hall, London, UK, 1990.
- [15] H. Lesot, M. Osman, and J. V. Ruch, "Immunofluorescent localization of collagens, fibronectin, and laminin during terminal differentiation of odontoblasts," *Developmental Biology*, vol. 82, no. 2, pp. 371–381, 1981.
- [16] H. Lesot, M.-D. Kubler, J. L. Fausser, and J.-V. Ruch, "A 165 kDa membrane antigen mediating fibronectin-vinculin interaction is involved in murine odontoblast differentiation," *Differentiation*, vol. 44, no. 1, pp. 25–35, 1990.
- [17] A. Veis, "The role of dental pulp—thoughts on the session on pulp repair processes," *Journal of Dental Research*, vol. 64, pp. 552–554, 1985.
- [18] K. Yoshida, N. Yoshida, H. Nakamura, M. Iwaku, and H. Ozawa, "Immunolocalization of fibronectin during reparative dentinogenesis in human teeth after pulp capping with calcium hydroxide," *Journal of Dental Research*, vol. 75, no. 8, pp. 1590–1597, 1996.
- [19] M. Mizuno and Y. Banzai, "Calcium ion release from calcium hydroxide stimulated fibronectin gene expression in dental pulp cells and the differentiation of dental pulp cells to mineralized tissue forming cells by fibronectin," *International Endodontic Journal*, vol. 41, no. 11, pp. 933–938, 2008.
- [20] Y.-C. Hwang, I.-N. Hwang, W.-M. Oh, J.-C. Park, D.-S. Lee, and H.-H. Son, "Influence of TGF- $\beta$ 1 on the expression of BSP, DSP, TGF- $\beta$ 1 receptor I and Smad proteins during reparative dentinogenesis," *Journal of Molecular Histology*, vol. 39, no. 2, pp. 153–160, 2008.
- [21] M. Kuratate, K. Yoshida, Y. Shigetani, N. Yoshida, H. Ohshima, and T. Okiji, "Immunohistochemical analysis of nestin, osteopontin, and proliferating cells in the reparative process of exposed dental pulp capped with mineral trioxide aggregate," *Journal of Endodontics*, vol. 34, no. 8, pp. 970–974, 2008.
- [22] R. N. D'Souza, T. Bachman, K. R. Baumgardner, W. T. Butler, and M. Litz, "Characterization of cellular responses involved in reparative dentinogenesis in rat molars," *Journal of Dental Research*, vol. 74, no. 2, pp. 702–709, 1995.
- [23] K.-S. Min, H.-J. Park, S.-K. Lee, et al., "Effect of mineral trioxide aggregate on dentin bridge formation and expression of dentin sialoprotein and heme oxygenase-1 in human dental pulp," *Journal of Endodontics*, vol. 34, no. 6, pp. 666–670, 2008.
- [24] W. T. Butler, "Dentin matrix proteins," *European Journal of Oral Sciences*, vol. 106, no. 1, supplement, pp. 204–210, 1998.
- [25] J. Sodek, B. Ganss, and M. D. McKee, "Osteopontin," *Critical Reviews in Oral Biology & Medicine*, vol. 11, no. 3, pp. 279–303, 2000.
- [26] A. J. Smith, J. B. Matthews, and R. C. Hall, "Transforming growth factor-beta1 (TGF- $\beta$ 1) in dentine matrix. Ligand activation and receptor expression," *European Journal of Oral Sciences*, vol. 106, supplement 1, pp. 179–184, 1998.
- [27] A. J. Sloan, H. Perry, J. B. Matthews, and A. J. Smith, "Transforming growth factor- $\beta$  isoform expression in mature human healthy and carious molar teeth," *Histochemical Journal*, vol. 32, no. 4, pp. 247–252, 2000.
- [28] A. J. Sloan, M. L. Couble, F. Bleicher, H. Magloire, A. J. Smith, and J. C. Farges, "Expression of TGF- $\beta$  receptors I and II in the human dental pulp by in situ hybridization," *Advances in Dental Research*, vol. 15, pp. 63–67, 2001.
- [29] D. L. Mooradian, R. C. Lucas, J. A. Weatherbee, and L. T. Furcht, "Transforming growth factor- $\beta$ 1 binds to immobilized fibronectin," *Journal of Cellular Biochemistry*, vol. 41, no. 4, pp. 189–200, 1989.
- [30] E. Ruoslahti and Y. Yamaguchi, "Proteoglycans as modulators of growth factor activities," *Cell*, vol. 64, no. 5, pp. 867–869, 1991.
- [31] S. Gronthos, M. Mankani, J. Brahim, P. G. Robey, and S. Shi, "Postnatal human dental pulp stem cells (DPSCs) in vitro and in vivo," *Proceedings of the National Academy of Sciences of the United States of America*, vol. 97, no. 25, pp. 13625–13630, 2000.
- [32] S. Gronthos, J. Brahim, W. Li, et al., "Stem cell properties of human dental pulp stem cells," *Journal of Dental Research*, vol. 81, no. 8, pp. 531–535, 2002.
- [33] M. Miura, S. Gronthos, M. Zhao, B. Lu, L. W. Fisher, P. G. Robey, and S. Shi, "SHED: stem cells from human exfoliated deciduous teeth," *Proceedings of the National Academy of Sciences of the United States of America*, vol. 100, no. 10, pp. 5807–5812, 2003.
- [34] S. Shi and S. Gronthos, "Perivascular niche of postnatal mesenchymal stem cells in human bone marrow and dental pulp," *Journal of Bone and Mineral Research*, vol. 18, no. 4, pp. 696–704, 2003.
- [35] H. Ohshima, K. Nakakura-Ohshima, H. Yamamoto, and T. Maeda, "Alteration in the expression of heat shock protein (Hsp) 25-immunoreactivity in the dental pulp of rat molars following tooth replantation," *Archives of Histology and Cytology*, vol. 64, no. 4, pp. 425–437, 2001.
- [36] A. Hosoya, K. Yoshida, N. Yoshida, K. Hoshi, M. Iwaku, and H. Ozawa, "An immunohistochemical study on hard tissue formation in a subcutaneously transplanted rat molar," *Histochemistry and Cell Biology*, vol. 119, no. 1, pp. 27–35, 2003.
- [37] C. Zhao, A. Hosoya, H. Kurita, et al., "Immunohistochemical study of hard tissue formation in the rat pulp cavity after tooth replantation," *Archives of Oral Biology*, vol. 52, no. 10, pp. 945–953, 2007.
- [38] Y. Takamori, H. Suzuki, K. Nakakura-Ohshima, et al., "Capacity of dental pulp differentiation in mouse molars as demonstrated by allogenic tooth transplantation," *Journal of Histochemistry & Cytochemistry*, vol. 56, no. 12, pp. 1075–1086, 2008.
- [39] J. Camilleri and T. R. Pitt Ford, "Mineral trioxide aggregate: a review of the constituents and biological properties of the material," *International Endodontic Journal*, vol. 39, no. 10, pp. 747–754, 2006.
- [40] H. W. Roberts, J. M. Toth, D. W. Berzins, and D. G. Charlton, "Mineral trioxide aggregate material use in endodontic

- treatment: a review of the literature," *Dental Materials*, vol. 24, no. 2, pp. 149–164, 2008.
- [41] Q. Zhu, R. Haglund, K. E. Safavi, and L. S. W. Spångberg, "Adhesion of human osteoblasts on root-end filling materials," *Journal of Endodontics*, vol. 26, no. 7, pp. 404–406, 2000.
- [42] G. A. Pelliccioni, G. Ciapetti, E. Cenni, et al., "Evaluation of osteoblast-like cell response to Proroot MTA (mineral trioxide aggregate) cement," *Journal of Materials Science: Materials in Medicine*, vol. 15, no. 2, pp. 167–173, 2004.
- [43] S. Bonson, B. G. Jeansonne, and T. E. Lallier, "Root-end filling materials alter fibroblast differentiation," *Journal of Dental Research*, vol. 83, no. 5, pp. 408–413, 2004.
- [44] R. Haglund, J. He, K. E. Safavi, L. S. W. Spångberg, and Q. Zhu, "Effects of root-end filling materials on fibroblasts and macrophages in vitro," *Oral Surgery, Oral Medicine, Oral Pathology, Oral Radiology, and Endodontics*, vol. 95, no. 6, pp. 739–745, 2003.
- [45] H. A. Balto, "Attachment and morphological behavior of human periodontal ligament fibroblasts to mineral trioxide aggregate: a scanning electron microscope study," *Journal of Endodontics*, vol. 30, no. 1, pp. 25–29, 2004.
- [46] M. Fridland and R. Rosado, "Mineral trioxide aggregate (MTA) solubility and porosity with different water-to-powder ratios," *Journal of Endodontics*, vol. 29, no. 12, pp. 814–817, 2003.
- [47] M. A. Duarte, A. C. C. Demarchi, J. C. Yamashita, M. C. Kuga, and C. Fraga Sde, "pH and calcium ion release of 2 root-end filling materials," *Oral Surgery, Oral Medicine, Oral Pathology, Oral Radiology, and Endodontics*, vol. 95, no. 3, pp. 345–347, 2003.
- [48] M. Fridland and R. Rosado, "MTA solubility: a long term study," *Journal of Endodontics*, vol. 31, no. 5, pp. 376–379, 2005.
- [49] J. Camilleri, "Characterization of hydration products of mineral trioxide aggregate," *International Endodontic Journal*, vol. 41, no. 5, pp. 408–417, 2008.
- [50] G. Guven, Z. C. Cehreli, A. Ural, M. A. Serdar, and F. Basak, "Effect of mineral trioxide aggregate cements on transforming growth factor  $\beta$ 1 and bone morphogenetic protein production by human fibroblasts in vitro," *Journal of Endodontics*, vol. 33, no. 4, pp. 447–450, 2007.
- [51] N. Tani-Ishii, N. Hamada, K. Watanabe, Y. Tujimoto, T. Teranaka, and T. Umemoto, "Expression of bone extracellular matrix proteins on osteoblast cells in the presence of mineral trioxide," *Journal of Endodontics*, vol. 33, no. 7, pp. 836–839, 2007.
- [52] S. S. Hakki, S. B. Bozkurt, E. E. Hakki, and S. Belli, "Effects of mineral trioxide aggregate on cell survival, gene expression associated with mineralized tissues, and biomineralization of cementoblasts," *Journal of Endodontics*, vol. 35, no. 4, pp. 513–519, 2009.
- [53] Y. Yasuda, M. Ogawa, T. Arakawa, T. Kadowaki, and T. Saito, "The effect of mineral trioxide aggregate on the mineralization ability of rat dental pulp cells: an in vitro study," *Journal of Endodontics*, vol. 34, no. 9, pp. 1057–1060, 2008.
- [54] K.-S. Min, S.-H. Yang, and E.-C. Kim, "The combined effect of mineral trioxide aggregate and enamel matrix derivative on odontoblastic differentiation in human dental pulp Cells," *Journal of Endodontics*, vol. 35, no. 6, pp. 847–851, 2009.
- [55] H. Minamikawa, Y. Deyama, K. Nakamura, et al., "Effect of mineral trioxide aggregate on rat clonal dental pulp cells: expression of cyclooxygenase-2 mRNA and inflammation-related protein via nuclear factor  $\kappa$  B signaling system," *Journal of Endodontics*, vol. 35, no. 6, pp. 843–846, 2009.
- [56] M. C. Tingey, P. Bush, and M. S. Levine, "Analysis of mineral trioxide aggregate surface when set in the presence of fetal bovine serum," *Journal of Endodontics*, vol. 34, no. 1, pp. 45–49, 2008.
- [57] F. R. Tay, D. H. Pashley, F. A. Rueggeberg, R. J. Loushine, and R. N. Weller, "Calcium phosphate phase transformation produced by the interaction of the Portland cement component of white mineral trioxide aggregate with a phosphate-containing fluid," *Journal of Endodontics*, vol. 33, no. 11, pp. 1347–1351, 2007.
- [58] J. F. Reyes-Carmona, M. S. Felipe, and W. T. Felipe, "Biomineralization ability and interaction of mineral trioxide aggregate and white Portland cement with dentin in a phosphate-containing fluid," *Journal of Endodontics*, vol. 35, no. 5, pp. 731–736, 2009.
- [59] R. Holland, V. De Souza, M. J. Nery, J. A. Otoboni Filho, P. F. E. Bernabé, and E. Dezan Jr., "Reaction of rat connective tissue to implanted dentin tubes filled with mineral trioxide aggregate or calcium hydroxide," *Journal of Endodontics*, vol. 25, no. 3, pp. 161–166, 1999.
- [60] M. Lotfi, S. Vosoughhosseini, M. A. Saghiri, M. Mesgari-abbasi, and B. Ranjkesh, "Effect of white mineral trioxide aggregate mixed with disodium hydrogen phosphate on inflammatory cells," *Journal of Endodontics*, vol. 35, no. 5, pp. 703–705, 2009.
- [61] T. R. Pitt Ford, M. Torabinejad, H. R. Abedi, L. K. Bakland, and S. P. Kariyawasam, "Using mineral trioxide aggregate: as a pulp-capping material," *Journal of the American Dental Association*, vol. 127, no. 10, pp. 1491–1494, 1996.
- [62] I. M. Faraco Jr. and R. Holland, "Response of the pulp of dogs to capping with mineral trioxide aggregate or a calcium hydroxide cement," *Dental Traumatology*, vol. 17, no. 4, pp. 163–166, 2001.
- [63] R. Holland, V. de Souza, S. S. Murata, et al., "Healing process of dog dental pulp after pulpotomy and pulp covering with mineral trioxide aggregate or Portland cement," *Brazilian Dental Journal*, vol. 12, no. 2, pp. 109–113, 2001.
- [64] D. Tziafas, O. Pantelidou, A. Alvanou, G. Belibasakis, and S. Papadimitriou, "The dentinogenic effect of mineral trioxide aggregate (MTA) in short-term capping experiments," *International Endodontic Journal*, vol. 35, no. 3, pp. 245–254, 2002.
- [65] W. E. Andelin, S. Shabahang, K. Wright, and M. Torabinejad, "Identification of hard tissue after experimental pulp capping using dentin sialoprotein (DSP) as a marker," *Journal of Endodontics*, vol. 29, no. 10, pp. 646–650, 2003.
- [66] M. S. Dominguez, D. E. Witherspoon, J. L. Gutmann, and L. A. Opperman, "Histological and scanning electron microscopy assessment of various vital pulp-therapy materials," *Journal of Endodontics*, vol. 29, no. 5, pp. 324–333, 2003.
- [67] R. Menezes, C. M. Bramante, A. Letra, V. G. G. Carvalho, and R. B. Garcia, "Histologic evaluation of pulpotomies in dog using two types of mineral trioxide aggregate and regular and white Portland cements as wound dressings," *Oral Surgery, Oral Medicine, Oral Pathology, Oral Radiology and Endodontology*, vol. 98, no. 3, pp. 376–379, 2004.
- [68] M. Parirokh, S. Asgary, M. J. Eghbal, and F. Brink, "A comparative study of white and grey mineral trioxide aggregate as pulp capping agents in dog's teeth," *Dental Traumatology*, vol. 21, no. 3, pp. 150–154, 2005.
- [69] S. Simon, P. Cooper, A. Smith, B. Picard, C. Naulin Ifi, and A. Berdal, "Evaluation of a new laboratory model for pulp

- healing: preliminary study," *International Endodontic Journal*, vol. 41, no. 9, pp. 781–790, 2008.
- [70] E. A. Bortoluzzi, N. J. Broon, C. M. Bramante, et al., "Mineral trioxide aggregate with or without calcium chloride in psulpotomy," *Journal of Endodontics*, vol. 34, no. 2, pp. 172–175, 2008.
- [71] M. Aeinehchi, B. Eslami, M. Ghanbariha, and A. S. Saffar, "Mineral trioxide aggregate (MTA) and calcium hydroxide as pulp-capping agents in human teeth: a preliminary report," *International Endodontic Journal*, vol. 36, no. 3, pp. 225–231, 2003.
- [72] V. Chacko and S. Kurikose, "Human pulpal response to mineral trioxide aggregate (MTA): a histologic study," *Journal of Clinical Pediatric Dentistry*, vol. 30, no. 3, pp. 203–209, 2006.
- [73] C. E. Iwamoto, E. Adachi, C. H. Pameijer, D. Barnes, E. E. Romberg, and S. Jefferies, "Clinical and histological evaluation of white ProRoot MTA in direct pulp capping," *American Journal of Dentistry*, vol. 19, no. 2, pp. 85–90, 2006.
- [74] M. d. L. R. Accorinte, R. Holland, A. Reis, et al., "Evaluation of mineral trioxide aggregate and calcium hydroxide cement as pulp-capping agents in human teeth," *Journal of Endodontics*, vol. 34, no. 1, pp. 1–6, 2008.
- [75] P. N. R. Nair, H. F. Duncan, T. R. Pitt Ford, and H. U. Luder, "Histological, ultrastructural and quantitative investigations on the response of healthy human pulps to experimental capping with mineral trioxide aggregate: a randomized controlled trial," *International Endodontic Journal*, vol. 41, no. 2, pp. 128–150, 2008.
- [76] M. L. R. Accorinte, A. D. Loguercio, A. Reis, et al., "Response of human dental pulp capped with MTA and calcium hydroxide powder," *Operative Dentistry*, vol. 33, no. 5, pp. 488–495, 2008.
- [77] L. Sawicki, C. H. Pameijer, K. Emerich, and B. Adamowicz-Klepalska, "Histological evaluation of mineral trioxide aggregate and calcium hydroxide in direct pulp capping of human immature permanent teeth," *American Journal of Dentistry*, vol. 21, no. 4, pp. 262–266, 2008.
- [78] M. L. R. Accorinte, A. D. Loguercio, A. Reis, et al., "Evaluation of two mineral trioxide aggregate compounds as pulp-capping agents in human teeth," *International Endodontic Journal*, vol. 42, no. 2, pp. 122–128, 2009.
- [79] M. J. Eghbal, S. Asgary, R. A. Baglue, M. Parirokh, and J. Ghoddsi, "MTA pulpotomy of human permanent molars with irreversible pulpitis," *Australian Endodontic Journal*, vol. 35, no. 1, pp. 4–8, 2009.
- [80] D. Abdullah, T. R. Pitt Ford, S. Papaioannou, J. Nicholson, and F. McDonald, "An evaluation of accelerated Portland cement as a restorative material," *Biomaterials*, vol. 23, no. 19, pp. 4001–4010, 2002.
- [81] I. Islam, H. Kheng Chng, and A. U. J. Yap, "Comparison of the physical and mechanical properties of MTA and portland cement," *Journal of Endodontics*, vol. 32, no. 3, pp. 193–197, 2006.
- [82] M. Torabinejad, C. U. Hong, F. McDonald, and T. R. Pitt Ford, "Physical and chemical properties of a new root-end filling material," *Journal of Endodontics*, vol. 21, no. 7, pp. 349–353, 1995.
- [83] M. H. Nekoofar, G. Adusei, M. S. Sheykhrezae, S. J. Hayes, S. T. Bryant, and P. M. H. Dummer, "The effect of condensation pressure on selected physical properties of mineral trioxide aggregate," *International Endodontic Journal*, vol. 40, no. 6, pp. 453–461, 2007.
- [84] J. Camilleri, "Hydration mechanisms of mineral trioxide aggregate," *International Endodontic Journal*, vol. 40, no. 6, pp. 462–470, 2007.
- [85] Y.-L. Lee, B.-S. Lee, F.-H. Lin, A. Y. Lin, W.-H. Lan, and C.-P. Lin, "Effects of physiological environments on the hydration behavior of mineral trioxide aggregate," *Biomaterials*, vol. 25, no. 5, pp. 787–793, 2004.
- [86] L. Gancedo-Caravia and E. Garcia-Barbero, "Influence of humidity and setting time on the push-out strength of mineral trioxide aggregate obturations," *Journal of Endodontics*, vol. 32, no. 9, pp. 894–896, 2006.
- [87] A. M. Montellano, S. A. Schwartz, and T. J. Beeson, "Contamination of tooth-colored mineral trioxide aggregate used as a root-end filling material: a bacterial leakage study," *Journal of Endodontics*, vol. 32, no. 5, pp. 452–455.
- [88] R. A. VanderWeele, S. A. Schwartz, and T. J. Beeson, "Effect of blood contamination on retention characteristics of MTA when mixed with different liquids," *Journal of Endodontics*, vol. 32, no. 5, pp. 421–424, 2006.
- [89] S. R. Sluyk, P. C. Moon, and G. R. Hartwell, "Evaluation of setting properties and retention characteristics of mineral trioxide aggregate when used as a furcation perforation repair material," *Journal of Endodontics*, vol. 24, no. 11, pp. 768–771, 1998.
- [90] G. Danesh, T. Dammaschke, H. U. V. Gerth, T. Zandbiglari, and E. Schäfer, "A comparative study of selected properties of ProRoot mineral trioxide aggregate and two Portland cements," *International Endodontic Journal*, vol. 39, no. 3, pp. 213–219, 2006.
- [91] C. Poggio, M. Lombardini, C. Alessandro, and R. Simonetta, "Solubility of root-end-filling materials: a comparative study," *Journal of Endodontics*, vol. 33, no. 9, pp. 1094–1097, 2007.
- [92] A. Bodanezi, N. Carvalho, D. Silva, et al., "Immediate and delayed solubility of mineral trioxide aggregate and Portland cement," *Journal of Applied Oral Science*, vol. 16, no. 2, pp. 127–131, 2008.
- [93] M. Kuratate, Y. Shigetani, L. Han, and T. Okiji, "Compositional change of mineral trioxide aggregate immersed in water: alteration of elemental distribution in the surface layer," *Japanese Journal of Conservative Dentistry*, vol. 52, no. 4, pp. 348–354, 2009.
- [94] C. Carde, R. François, and J.-M. Torrenti, "Leaching of both calcium hydroxide and C-S-H from cement paste: modeling the mechanical behavior," *Cement and Concrete Research*, vol. 26, no. 8, pp. 1257–1268, 1996.
- [95] K. Haga, M. Shibata, M. Hironaga, S. Tanaka, and S. Nagasaki, "Change in pore structure and composition of hardened cement paste during the process of dissolution," *Cement and Concrete Research*, vol. 35, no. 5, pp. 943–950, 2005.
- [96] P. Hørsted-Bindslev and H. Løvschall, "Treatment outcome of vital pulp treatment," *Endodontic Topics*, vol. 2, pp. 24–34, 2002.
- [97] G. Bergenholtz, "Evidence for bacterial causation of adverse pulpal responses in resin-based dental restorations," *Critical Reviews in Oral Biology & Medicine*, vol. 11, no. 4, pp. 467–480, 2000.
- [98] A. H. B. Schuur, R. J. M. Gruythuysen, and P. R. Weselink, "Pulp capping with adhesive resin-based composite vs. calcium hydroxide: a review," *Endodontics and Dental Traumatology*, vol. 16, no. 6, pp. 240–250, 2000.

- [99] N. K. Sarkar, R. Caicedo, P. Ritwik, R. Moiseyeva, and I. Kawashima, "Physicochemical basis of the biologic properties of mineral trioxide aggregate," *Journal of Endodontics*, vol. 31, no. 2, pp. 97–100, 2005.
- [100] T. B. Bozeman, R. R. Lemon, and P. D. Eleazer, "Elemental analysis of crystal precipitate from gray and white MTA," *Journal of Endodontics*, vol. 32, no. 5, pp. 425–428, 2006.
- [101] F. R. Tay and D. H. Pashley, "Guided tissue remineralisation of partially demineralised human dentine," *Biomaterials*, vol. 29, no. 8, pp. 1127–1137, 2008.
- [102] P. Taddei, A. Tinti, M. G. Gandolfi, P. L. Rossi, and C. Prati, "Vibrational study on the bioactivity of Portland cement-based materials for endodontic use," *Journal of Molecular Structure*, vol. 924–926, pp. 548–554, 2009.
- [103] T. Kokubo, "Bioactive glass ceramics: properties and applications," *Biomaterials*, vol. 12, no. 2, pp. 155–163, 1991.
- [104] T. Kasuga, "Bioactive calcium pyrophosphate glasses and glass-ceramics," *Acta Biomaterialia*, vol. 1, no. 1, pp. 55–64, 2005.
- [105] M.-K. Wu, E. G. Kontakiotis, and P. R. Wesselink, "Long-term seal provided by some root-end filling materials," *Journal of Endodontics*, vol. 24, no. 8, pp. 557–560, 1998.
- [106] X. Liu, C. Ding, and P. K. Chu, "Mechanism of apatite formation on wollastonite coatings in simulated body fluids," *Biomaterials*, vol. 25, no. 10, pp. 1755–1761, 2004.
- [107] K. Ohura, T. Nakamura, T. Yamamuro, et al., "Bone-bonding ability of P<sub>2</sub>O<sub>5</sub>-free CaO.SiO<sub>2</sub> glasses," *Journal of Biomedical Materials Research*, vol. 25, no. 3, pp. 357–365, 1991.
- [108] H. K. Chng, I. Islam, A. U. Yap, Y. W. Tong, and E. T. Koh, "Properties of a new root-end filling material," *Journal of Endodontics*, vol. 31, no. 9, pp. 665–668, 2005.
- [109] I. Islam, H. K. Chng, and A. U. J. Yap, "Comparison of the physical and mechanical properties of MTA and portland cement," *Journal of Endodontics*, vol. 32, no. 3, pp. 193–197, 2006.
- [110] B. Storm, F. C. Eichmiller, P. A. Tordik, and G. G. Goodell, "Setting expansion of gray and white mineral trioxide aggregate and Portland Cement," *Journal of Endodontics*, vol. 34, no. 1, pp. 80–82, 2008.
- [111] M. Torabinejad, P. W. Smith, J. D. Kettering, and T. R. Pitt Ford, "Comparative investigation of marginal adaptation of mineral trioxide aggregate and other commonly used root-end filling materials," *Journal of Endodontics*, vol. 21, no. 6, pp. 295–299, 1995.
- [112] E. Gondim Jr., A. A. Zaia, B. P. F. A. Gomes, C. C. R. Ferraz, F. B. Teixeira, and F. J. Souza-Filho, "Investigation of the marginal adaptation of root-end filling materials in root-end cavities prepared with ultrasonic tips," *International Endodontic Journal*, vol. 36, no. 7, pp. 491–499, 2003.
- [113] G. Shipper, E. S. Grossman, A. J. Botha, and P. E. Cleaton-Jones, "Marginal adaptation of mineral trioxide aggregate (MTA) compared with amalgam as a root-end filling material: a low-vacuum (LV) versus high-vacuum (HV) SEM study," *International Endodontic Journal*, vol. 37, no. 5, pp. 325–336, 2004.
- [114] C. B. Xavier, R. Weismann, M. G. de Oliveira, F. F. Demarco, and D. H. Pozza, "Root-end filling materials: apical microleakage and marginal adaptation," *Journal of Endodontics*, vol. 31, no. 7, pp. 539–542, 2005.
- [115] S. I. Tobón-Arroyave, M. M. Restrepo-Pérez, J. A. Arismendi-Echavarría, Z. Velásquez-Restrepo, M. L. Marín-Botero, and E. C. García-Dorado, "Ex vivo microscopic assessment of factors affecting the quality of apical seal created by root-end fillings," *International Endodontic Journal*, vol. 40, no. 8, pp. 590–602, 2007.
- [116] P. Yan, B. Peng, B. Fan, M. Fan, and Z. Bian, "The effects of sodium hypochlorite (5.25%), chlorhexidine (2%), and glyde file prep on the bond strength of MTA-dentin," *Journal of Endodontics*, vol. 32, no. 1, pp. 58–60, 2006.
- [117] S. Shinzato, M. Kobayashi, W. F. Mousa, et al., "Bioactive bone cement: effect of surface curing properties on bone-bonding strength," *Journal of Biomedical Materials Research*, vol. 53, no. 1, pp. 51–61, 2000.
- [118] R. L. Leung, W. J. Loesche, and G. T. Charbeneau, "Effect of Dycal on bacteria in deep carious lesions," *The Journal of the American Dental Association*, vol. 100, no. 2, pp. 193–197, 1980.
- [119] M. Torabinejad, C. U. Hong, T. R. P. Ford, and J. D. Kettering, "Antibacterial effects of some root end filling materials," *Journal of Endodontics*, vol. 21, no. 8, pp. 403–406, 1995.
- [120] T. J. Stowe, C. M. Sedgley, B. Stowe, and J. C. Fenno, "The effects of chlorhexidine gluconate (0.12%) on the antimicrobial properties of tooth-colored ProRoot mineral trioxide aggregate," *Journal of Endodontics*, vol. 30, no. 6, pp. 429–431, 2004.
- [121] C. R. Sipert, R. P. Hussne, C. K. Nishiyama, and S. A. Torres, "In vitro antimicrobial activity of Fill Canal, Sealapex, Mineral Trioxide Aggregate, Portland cement and EndoRez," *International Endodontic Journal*, vol. 38, no. 8, pp. 539–543, 2005.
- [122] A. U. Eldeniz, H. H. Hadimli, H. Ataoglu, and D. Ørstavik, "Antibacterial effect of selected root-end filling materials," *Journal of Endodontics*, vol. 32, no. 4, pp. 345–349, 2006.
- [123] K. Al-Hezaimi, T. A. Al-Shalan, J. Naghshbandi, S. Oglesby, J. H. S. Simon, and I. Rotstein, "Antibacterial effect of two mineral trioxide aggregate (MTA) preparations against *Enterococcus faecalis* and *Streptococcus sanguis* in vitro," *Journal of Endodontics*, vol. 32, no. 11, pp. 1053–1056, 2006.
- [124] M. Tanomaru-Filho, J. M. Tanomaru, D. B. Barros, E. Watanabe, and I. Y. Ito, "In vitro antimicrobial activity of endodontic sealers, MTA-based cements and Portland cement," *Journal of Oral Science*, vol. 49, no. 1, pp. 41–45, 2007.
- [125] S. Asgary and F. A. Kamrani, "Antibacterial effects of five different root canal sealing materials," *Journal of Oral Science*, vol. 50, no. 4, pp. 469–474, 2008.
- [126] S. Al-Nazhan and A. Al-Judai, "Evaluation of antifungal activity of mineral trioxide aggregate," *Journal of Endodontics*, vol. 29, no. 12, pp. 826–827, 2003.
- [127] K. Al-Hezaimi, K. Al-Hamdan, J. Naghshbandi, S. Oglesby, J. H. S. Simon, and I. Rotstein, "Effect of white-colored mineral trioxide aggregate in different concentrations on *Candida albicans* in vitro," *Journal of Endodontics*, vol. 31, no. 9, pp. 684–686, 2005.
- [128] K. Al-Hezaimi, J. Naghshbandi, S. Oglesby, J. H. S. Simon, and I. Rotstein, "Comparison of antifungal activity of white-colored and gray-colored mineral trioxide aggregate (MTA) at similar concentrations against *Candida albicans*," *Journal of Endodontics*, vol. 32, no. 4, pp. 365–367, 2006.
- [129] N. Farsi, N. Alamoudi, K. Balto, and A. Al Mushayt, "Clinical assessment of mineral trioxide aggregate (MTA) as direct pulp capping in young permanent teeth," *Journal of Clinical Pediatric Dentistry*, vol. 31, no. 2, pp. 72–76, 2006.
- [130] G. Bogen, J. S. Kim, and L. K. Bakland, "Direct pulp capping with mineral trioxide aggregate: an observational study," *Journal of the American Dental Association*, vol. 139, no. 3, pp. 305–315, 2008.

- [131] D. Tuna and A. Ölmez, "Clinical long-term evaluation of MTA as a direct pulp capping material in primary teeth," *International Endodontic Journal*, vol. 41, no. 4, pp. 273–278, 2008.
- [132] K. M. Barrieshi-Nusair and M. A. Qudeimat, "A prospective clinical study of mineral trioxide aggregate for partial pulpotomy in cariously exposed permanent teeth," *Journal of Endodontics*, vol. 32, no. 8, pp. 731–735, 2006.
- [133] O. A. S. El Meligy and D. R. Avery, "Comparison of mineral trioxide aggregate and calcium hydroxide as pulpotomy agents in young permanent teeth (apexogenesis)," *Pediatric Dentistry*, vol. 28, no. 5, pp. 399–404, 2006.
- [134] D. E. Witherspoon, J. C. Small, and G. Z. Harris, "Mineral trioxide aggregate pulpotomies: a case series outcomes assessment," *Journal of the American Dental Association*, vol. 137, no. 5, pp. 610–618, 2006.
- [135] M. A. Qudeimat, K. M. Barrieshi-Nusair, and A. I. Owais, "Calcium hydroxide vs mineral trioxide aggregates for partial pulpotomy of permanent molars with deep caries," *European Archives of Paediatric Dentistry*, vol. 8, no. 2, pp. 99–104, 2007.
- [136] E. T. Koh, T. R. Pitt Ford, S. P. Kariyawasam, N. N. Chen, and M. Torabinejad, "Prophylactic treatment of dens evaginatus using mineral trioxide aggregate," *Journal of Endodontics*, vol. 27, no. 8, pp. 540–542, 2001.
- [137] H. A. Agamy, N. S. Bakry, M. M. F. Mounir, and D. R. Avery, "Comparison of mineral trioxide aggregate and formocresol as pulp-capping agents in pulpotomized primary teeth," *Pediatric Dentistry*, vol. 26, no. 4, pp. 302–309, 2004.
- [138] S. E. Jabbarifar, D. D. Khademi, and D. D. Ghasemi, "Success rates of formocresol pulpotomy vs mineral trioxide aggregate in human primary molar tooth," *Journal of Research in Medical Sciences*, vol. 9, no. 6, pp. 304–307, 2004.
- [139] N. Farsi, N. Alamoudi, K. Balto, and A. Mushayt, "Success of mineral trioxide aggregate in pulpotomized primary molars," *Journal of Clinical Pediatric Dentistry*, vol. 29, no. 4, pp. 307–311, 2005.
- [140] G. Holan, E. Eidelman, and A. B. Fuks, "Long-term evaluation of pulpotomy in primary molars using mineral trioxide aggregate or formocresol," *Pediatric Dentistry*, vol. 27, no. 2, pp. 129–136, 2005.
- [141] L. Peng, L. Ye, H. Tan, and X. Zhou, "Evaluation of the formocresol versus mineral trioxide aggregate primary molar pulpotomy: a meta-analysis," *Oral Surgery, Oral Medicine, Oral Pathology, Oral Radiology and Endodontology*, vol. 102, no. 6, pp. e40–e44, 2006.
- [142] M. Aeinehchi, S. Dadvand, S. Fayazi, and S. Bayat-Movahed, "Randomized controlled trial of mineral trioxide aggregate and formocresol for pulpotomy in primary molar teeth," *International Endodontic Journal*, vol. 40, no. 4, pp. 261–267, 2007.
- [143] A. B. Fuks, "Vital pulp therapy with new materials for primary teeth: new directions and treatment perspectives," *Journal of Endodontics*, vol. 34, no. 7, supplement, pp. S18–S24, 2008.
- [144] P. Kogan, J. He, G. N. Glickman, and I. Watanabe, "The effects of various additives on setting properties of MTA," *Journal of Endodontics*, vol. 32, no. 6, pp. 569–572, 2006.
- [145] B. S. Ber, J. F. Hatton, and G. P. Stewart, "Chemical modification of ProRoot MTA to improve handling characteristics and decrease setting time," *Journal of Endodontics*, vol. 33, no. 10, pp. 1231–1234, 2007.
- [146] K. B. Wiltbank, S. A. Schwartz, and W. G. Schindler, "Effect of selected accelerants on the physical properties of mineral trioxide aggregate and Portland cement," *Journal of Endodontics*, vol. 33, no. 10, pp. 1235–1238, 2007.
- [147] E. A. Bortoluzzi, N. J. Broon, C. M. Bramante, W. T. Felipe, M. Tanomaru Filho, and R. M. Esberard, "The influence of calcium chloride on the setting time, solubility, disintegration, and pH of mineral trioxide aggregate and white Portland cement with a radiopacifier," *Journal of Endodontics*, vol. 35, no. 4, pp. 550–554, 2009.
- [148] E. A. Bortoluzzi, N. J. Broon, C. M. Bramante, R. B. Garcia, I. G. de Moraes, and N. Bernardineli, "Sealing ability of MTA and radiopaque Portland cement with or without calcium chloride for root-end filling," *Journal of Endodontics*, vol. 32, no. 9, pp. 897–900, 2006.
- [149] T.-H. Huang, M.-Y. Shie, C.-T. Kao, and S.-J. Ding, "The effect of setting accelerator on properties of mineral trioxide aggregate," *Journal of Endodontics*, vol. 34, no. 5, pp. 590–593, 2008.
- [150] S. J. Ding, C. T. Kao, M. Y. Shie, C. Hung Jr., and T. H. Huang, "The physical and cytological properties of white MTA mixed with Na<sub>2</sub>HPO<sub>4</sub> as an accelerant," *Journal of Endodontics*, vol. 34, no. 6, pp. 748–751, 2008.
- [151] J.E. Gomes-Filho, M. D. de Faria, P. F. E. Bernabé, et al., "Mineral trioxide aggregate but not light-cure mineral trioxide aggregate stimulated mineralization," *Journal of Endodontics*, vol. 34, no. 1, pp. 62–65, 2008.
- [152] J. Camilleri, "Modification of mineral trioxide aggregate. Physical and mechanical properties," *International Endodontic Journal*, vol. 41, no. 10, pp. 843–849, 2008.
- [153] M. G. Gandolfi, S. Pagani, F. Perut, et al., "Innovative silicate-based cements for endodontics: a study of osteoblast-like cell response," *Journal of Biomedical Materials Research A*, vol. 87, no. 2, pp. 477–486, 2008.
- [154] M. G. Gandolfi, F. Perut, G. Ciapetti, R. Mongiorgi, and C. Prati, "New Portland cement-based materials for endodontics mixed with articaine solution: a study of cellular response," *Journal of Endodontics*, vol. 34, no. 1, pp. 39–44, 2008.
- [155] S. Asgary, S. Shahabi, T. Jafarzadeh, S. Amini, and S. Kheirieh, "The properties of a new endodontic material," *Journal of Endodontics*, vol. 34, no. 8, pp. 990–993, 2008.
- [156] S. Asgary, M. J. Eghbal, M. Parirokh, F. Ghanavati, and H. Rahimi, "A comparative study of histologic response to different pulp capping materials and a novel endodontic cement," *Oral Surgery, Oral Medicine, Oral Pathology, Oral Radiology and Endodontology*, vol. 106, no. 4, pp. 609–614, 2008.
- [157] S. Asgary, M. J. Eghbal, and M. Parirokh, "Sealing ability of a novel endodontic cement as a root-end filling material," *Journal of Biomedical Materials Research A*, vol. 87, no. 3, pp. 706–709, 2008.



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