

## Review Article

# Silver Nanoparticles in Dental Biomaterials

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Silver has been used in medicine for centuries because of its antimicrobial properties. More recently, silver nanoparticles have been synthesized and incorporated into several biomaterials, since their small size provides great antimicrobial effect, at low filler level. Hence, these nanoparticles have been applied in dentistry, in order to prevent or reduce biofilm formation over dental materials surfaces. This review aims to discuss the current progress in this field, highlighting aspects regarding silver nanoparticles incorporation, such as antimicrobial potential, mechanical properties, cytotoxicity, and long-term effectiveness. We also emphasize the need for more studies to determine the optimal concentration of silver nanoparticle and its release over time.

## 1. Introduction

Silver (Ag) ions or salts are known to have a wide antimicrobial effect [1–4] and they have been used for years [5], in different fields in medicine, including wound dressings [6], catheters [7], and prostheses [8]. Besides being a potent antimicrobial, Ag has many advantages, such as low toxicity and good biocompatibility with human cells [9], long-term antibacterial activity, due to sustained ion release [10], and low bacterial resistance [11].

With the advent of nanotechnology, silver nanoparticles (AgNPs) have been synthesized, and they have shown potent antimicrobial properties [12, 13]. AgNPs have demonstrated unique interactions with bacteria and fungi species [12, 14]; thereafter, they are widely used in medical arena, such as in wound sutures [15], endotracheal tubes [16], surgical instruments [17], and bone prostheses [18].

AgNPs have also been applied in several areas of dentistry, as endodontics [19, 20], dental prostheses [21], implantology [22, 23], and restorative dentistry [24–26]. AgNPs incorporation aims to avoid or at least to decrease the microbial colonization over dental materials, increasing oral health levels and improving life quality.

Because of their small size, AgNPs possess chemical, physical, and biological properties distinctive from those presented by traditional bulk materials [27]. Their smaller particles and large surface area provide potent antibacterial effects at a low filler level, diminishing Ag particle concentration necessary for its efficacy [25, 28–30] and avoiding negative influence on mechanical properties [10, 31, 32].

Other advantage provided by the small size is the possibility of AgNPs to penetrate through cell membranes more readily, resulting in higher antimicrobial activity [33], which is especially important since microorganisms in biofilms are more resistant to antimicrobial agents than planktonic pathogens [34].

The antimicrobial mechanism of AgNPs has been extensively investigated but it remains unclear [35]. It seems that silver ions interact with the peptidoglycan cell wall [34] causing structural changes, increased membrane permeability and, finally, cell death [3]. Further, AgNPs could interact with the exposed sulfhydryl groups in bacterial proteins, avoiding DNA replication [36].

Another important aspect to be studied is the toxicity and biocompatibility of AgNPs. Considering their unique physical and chemical properties, it is likely that these

nanoparticles also possess unique toxicity mechanisms [37]. Because of that, a better understanding of AgNPs safety is needed, in order to increase their clinical use [38].

Nanotechnology provides a wide range of possibilities to develop new antimicrobial materials [3]. However, there are disadvantages, for example, color change, an important property of dental materials [39]. In this review, we discuss AgNPs incorporation into dental materials, such as composite resin and adhesive systems, acrylic resin, root canal fillings, and implants, highlighting aspects regarding microorganism growth inhibition, cytotoxicity, and physical properties of these modified materials.

## 2. AgNPs Characterization

One important step for the development of AgNPs-containing materials is their characterization. Many studies have analyzed the Ag dispersion [26, 28, 32, 40], through Transmission Electron Microscopy (TEM). This technique allows visualizing how AgNPs spread into the tested material, as well as to verify the particle size.

According to Cheng et al. [28], NAg particles of ~3 nm were clearly visible and well dispersed throughout the polymer matrix. These results were confirmed in a subsequent study [29], in which authors reported NAg sizes ranging from 2 to 5 nm. This very small size allows NAg penetration on dentinal tubules [40], which can represent the possibility of inactivating residual bacteria on dentine. Besides that, it has also been shown that AgNPs were well dispersed in the material with minimal appearance of nanoparticle aggregates [26, 32].

Another important feature to be analyzed is the minimum inhibitory concentration (MIC) of AgNPs. MIC is defined as the lowest concentration of antimicrobial agent at which 90% growth is observed in the medium [46]. Hernández-Sierra et al. [47] used the liquid dilution method to find the MIC of 25 nm-AgNPs against *S. mutans* strains, and the results showed an average MIC of  $4.86 \pm 2.71 \mu\text{g/mL}$ , suggesting a higher antimicrobial effect of AgNPs.

In a similar study, Espinosa-Cristóbal et al. [48] tested different AgNPs sizes (8.4 nm, 16.1 nm, and 98 nm) and they reported higher MICs than the abovementioned study:  $101.98 \pm 72.03 \mu\text{g/mL}$ ,  $145.64 \pm 104.88 \mu\text{g/mL}$ , and  $320.63 \pm 172.83 \mu\text{g/mL}$ , respectively. This probably occurred because of the methodology, which included sucrose for *S. mutans* growing. Authors also verified that MICs were directly proportional to the particle size; it means, as bigger the size as higher the MIC.

## 3. Forms of Incorporation

AgNPs used in dental materials are incorporated through distinct ways, depending on the type of material. For composite resin and adhesive systems, the most common technique is adding a monomer, usually 2-(tert-butylamino)ethyl methacrylate, in order to improve Ag salt solubility in the resin solution [28, 32, 40]. For dental implants, the process is totally different: Titanium samples are soaked in  $\text{AgNO}_3$

solutions, rinsed with deionized water, dried, and irradiated with UV light from a high-pressure Hg lamp. This process allows producing samples with different Ag concentrations, depending on the  $\text{AgNO}_3$  solution concentration [49].

Another difference is related to the form of AgNP obtainment. In some studies the particles are commercially available, so they are obtained directly from the producer [50–52]. In others, AgNPs are prepared by reduction of  $\text{AgNO}_3$ , with  $\text{NaBO}_4$  [22], polyvinylpyrrolidone [21], sodium citrate [44], and gallic acid [48], among others.

## 4. Composite Resin and Adhesive System

Dental caries is still the most common and widespread oral disease, having as the main etiologic agent the acidic attack from cariogenic bacteria, such as *Streptococcus mutans* and *Lactobacillus* spp. [51]. Currently, the most widely dental material used to treat caries lesions is composite resin, especially because of its esthetics and load-bearing properties [53–55]. Hence, many studies have been performed, in order to improve quality and durability of polymeric restorative materials [56–58].

In spite of the notable advances obtained, composite restorations accumulate more biofilm than other restorative materials [59–62]. This is especially important in cases of failures on restoration margin [63]. Actually, although it is wanted, the perfect sealing between the restorative material and the cavity wall often does not occur [32]. It has been shown that there is microleakage on restoration margins, and these gaps can be colonized by oral bacteria, resulting in secondary caries [64], which makes necessary the restoration replacement.

In order to prevent or to diminish biofilm accumulation over composite and in the restorations margins, antimicrobial restorative materials have been developed, especially through the incorporation of AgNPs to composite resins [24–26] and adhesive systems [29, 32, 40–42, 50, 52, 65, 66]. These materials are multiphase substances composed of an organic polymer matrix, filler particles, coupling agent (silane), and the initiator-accelerator of polymerization [67], and AgNPs incorporation is based on the modification in the filler components [61].

A research developed by Cheng et al. [26] reported the effect of AgNPs incorporation, at different concentrations, to a composite resin, in order to investigate its mechanical properties and biofilm formation. In this study, composites were synthesized with AgNPs at 0.028, 0.042, 0.088, and 0.175%. Mechanical properties of composites with AgNPs at 0.028% and 0.042% were similar to those with no AgNPs. Besides that, counts of colony forming units (CFU) for total streptococci and *S. mutans*, using AgNPs at 0.042%, were 75% smaller than the control group without AgNPs. These data suggest that AgNPs incorporation to composite resins enables good mechanical properties and notable antimicrobial potential, even at low concentration.

In order to evaluate the influence of AgNPs incorporation on bond strength to dental substrate, Melo et al. [41] added AgNPs, at 0.1% by mass, to an adhesive system. The results

have shown that AgNPs did not compromise the bond strength ( $P > 0.1$ ), at the same time that it decreased metabolic activity on biofilm, compared to control group without AgNPs. In this study it was also observed reduction of CFU for total microorganisms, total streptococci, and mutans streptococci ( $P < 0.05$ ).

Li et al. [32] performed a study incorporating of AgNPs, at 0.05% by mass, to an adhesive system, aiming to assess bacterial inhibition provided by this antimicrobial, in both short and long distance. It has been reported that AgNPs reduced CFU number and acid lactic production on biofilm over and away to the adhesive surface, evidencing that AgNPs-containing adhesives enable long-distance antibacterial potential.

Another important aspect to be assessed is the biocompatibility of AgNPs-containing restorative materials. Accordingly, Zhang et al. [42] have studied the effects of AgNPs incorporation, at 0.05% by mass, to a primer and an adhesive, regarding human gingival fibroblast viability. It has been shown that AgNPs addition did not affect the cytotoxicity of primer and adhesive tested, evidencing the clinical applicability of this antimicrobial.

Based on abovementioned studies, it is possible to say that the antibacterial effects of AgNPs-containing restorative materials might decrease the development of recurrent caries, to increase the longevity of tooth restorations, and to be effective in decreasing the formation of bacterial biofilms on teeth and restorations, without compromising mechanical properties and cytotoxicity of composite resins and adhesive systems.

## 5. Acrylic Resin

Dentures, mostly constituted by poly(methyl methacrylate) (PMMA) acrylic resin [68], have their inner surface considerably rough [69], and this roughness, allied to other factors (e.g., poor hygiene, xerostomy, and HIV infection), contributes to the emergence of denture stomatitis [70, 71]. This pathology, characterized by red focal area, mostly localized in palatal mucosa, is present in 50–70% of complete denture wearers [72, 73], and it is frequently associated with *Candida* species colonization. These fungi colonize denture surfaces forming a biofilm [74], which acts as a key-factor to denture stomatitis development [75].

The treatment of denture stomatitis is based on topical or systemic antifungal drugs, for example, fluconazole and nystatin [76–78]. However, this infection is often persistent, since antifungal resistance has been reported in *Candida* biofilms [75]. Moreover, it has been observed that *Candida* species present in biofilms are less susceptible to antifungal drugs than planktonic cells [79–81]. Another problem related to denture stomatitis is that many geriatric prosthetic wearers present difficulties on keeping the denture clean, due to their reduced motor dexterity, memory loss, and cognitive impairment [21].

Considering the aforementioned factors, denture stomatitis represents a challenge for dentistry, and methods for its prevention, should be encouraged. Accordingly, AgNPs

have been satisfactorily incorporated into polymers used as tissue conditioners and as denture base [82–84]. The action mechanisms of AgNPs-incorporated polymers is still unclear, since some authors attribute the antimicrobial effectiveness to the silver ions release [85, 86] and others to the direct contact between the material and the microorganisms [87].

Acosta-Torres et al. [43] developed a PMMA containing 1  $\mu\text{g/mL}$  of AgNPs and they compared this new compound to unmodified PMMA. It has been observed that PMMA-AgNPs specimens showed significantly less *Candida albicans* adherence compared to PMMA ( $P < 0.05$ ), demonstrating the antifungal potential of AgNPs incorporated to acrylic resin. Besides that, they evaluated the activity of mouse fibroblasts and human lymphocytes, and it has been shown that PMMA-AgNP compound does not present cytotoxicity or genotoxicity. These results suggest that the novel acrylic resin incorporated with AgNPs could be developed as a denture base.

In a study performed by Monteiro et al. [44] AgNPs were incorporated in a commercial acrylic resin, in different concentrations (0.05%, 0.5%, and 5% of AgNPs, by mass). The authors evaluated the mechanical properties of the modified resin, as well of the unmodified one (0% of AgNPs). Thereunto, the flexural strength test was performed, and it was observed that all the groups presented very similar flexural resistance values, suggesting that AgNPs incorporation does not affect the mechanical properties of acrylic resin.

When dentures are ill-fitted is recommended recovering his base with tissue conditioners, which are easily degradable with time and occasionally susceptible to microbial colonization [88]. Thus, AgNPs incorporation could also be profitable in this material and not only in dentures base.

Accordingly, Nam [21] has incorporated AgNPs into a commercial tissue conditioner, in the following concentrations: 0.1%, 0.5%, 1.0%, 2.0%, and 3.0%. Their inhibitory effect was evaluated against *Staphylococcus aureus*, *Streptococcus mutans*, and *Candida albicans* after 24 h and 72 h. The authors have reported that the modified tissue conditioner presented antimicrobial properties even at lower concentrations, that is, 0.1% (for *S. mutans* and *S. aureus*) and 0.5% (for *C. albicans*).

## 6. Endodontic Materials

Several studies have demonstrated that bacteria are the main etiologic agent of pulpal infection and periradicular lesion formation [89–91]. The microbiota of infected root canals is polymicrobial and is dominated by Gram-negative anaerobes [92, 93]. It has been demonstrated that the presence of residual bacteria in root canal is connected with significantly higher rates of treatment failure [94].

Since elimination of bacteria in root canals is the key to treatment success [95], endodontic materials should ideally provide some antimicrobial activity [96, 97], in order to improve the prognosis of endodontically treated teeth [98]. Various materials have been used as root canal fillings, among which gutta-percha is one of the most used [95]. This material has been proved to present slight antibacterial property, provided by the zinc oxide in its components; however, this

does not provide to gutta-percha an effective bactericidal potential [98].

Accordingly, Iranian researchers [45] have introduced nanosilver-gutta-percha, as an attempt to improve the antibacterial effect of gutta-percha. The new material, which is standard gutta-percha coated with AgNPs, has demonstrated significant effect against *Enterococcus faecalis*, *Staphylococcus aureus*, *Candida albicans*, and *Escherichia coli*.

Besides that, Shantiaee et al. [99] have tested the biocompatibility of this new material, by comparing the cytotoxicity of nanosilver-coated gutta-percha and normal gutta-percha on mouse fibroblasts. In this study, after 24 hours, nanosilver-coated gutta-percha presented cytotoxicity similar to normal gutta-percha and, after one week, it reached the lowest level of cytotoxicity among the tested materials.

Other important step in the endodontic treatment is the chemomechanical debridement of pulpal tissue and pathogenic bacteria. In this stage, irrigant solutions should be used, for dissolving tissue and disinfecting the root canal system [100]. For this purpose, sodium hypochlorite (NaOCl) has been used for more than 70 years, and it remains as one of the most common solutions [101]. However, if NaOCl passes beyond the apex, it is extremely toxic to the periapical tissues [102].

In this context, Lotfi et al. [20] performed a study comparing the antibacterial effect of NaOCl and AgNP solution against *Enterococcus faecalis*, which is a bacterium often isolated from failed endodontic treatment cases [103]. Authors have observed that there were no significant differences among 5.25% NaOCl and 0.005% AgNPs, suggesting that this solution, in a remarkably lower concentration, possesses the same bactericidal effect as 5.25% NaOCl; hence, it could be used as a new intracanal irrigant.

Another important endodontic material is the mineral trioxide aggregate (MTA), used in many indications such as perforations sealing, external/internal root resorption repair, and apexification [104, 105]. In spite of being a material of wide application, the antimicrobial properties of MTA are controversial, and they seem to be limited [106, 107].

Aiming to improve its antimicrobial potential, Samiei et al. [19] modified MTA by adding AgNPs, at 1% weight. Its effect against oral bacteria and fungi species was assessed. Results have showed that AgNPs-containing MTA possesses higher antimicrobial effect against *Enterococcus faecalis*, *Candida albicans*, and *Pseudomonas aeruginosa*, compared to unmodified MTA.

Although AgNP is a promising antimicrobial, there are only a few studies employing it in endodontic materials. And considering that endodontic treatment success is highly connected to the bacteria elimination, researches involving AgNPs incorporation to root canal filling materials and intracanal irrigators should be encouraged.

## 7. Titanium Implants

Titanium (Ti) implants, widely used in dentistry, usually present infection around their surface, which remains one of the most important complications in Implantology [49, 108].

Several measures have been proposed to avoid bacterial contamination, such as implant disinfection and aseptic surgical protocols; nevertheless, bacterial invasion often occurs after surgery [109].

In order to prevent biofilm formation over implants surface, antibacterial coatings have been developed; however, most of them present poor long-term antibacterial action and also the possibility of generating resistant strains after prolonged use [110–112]. In this context, AgNPs incorporation to implant surface has been suggested [109, 113], since it would be possible to produce coatings with long-term antibacterial properties by controlling Ag release [23].

In study performed by Zhao et al. [23], AgNPs were incorporated into titania nanotubes (TiO<sub>2</sub>-NTs) on Ti implants, in a process involving silver nitrate immersion and ultraviolet radiation. The antibacterial effect against *Staphylococcus aureus* was assessed, and results have shown inhibition of planktonic bacteria during the first several days. Moreover, AgNPs-coating Ti implants have presented ability to prevent bacteria adhesion for up to 30 days, which are considered sufficient time to prevent post-infection in early stages.

In a similar study, Flores et al. [22] have evaluated the antibacterial activity of AgNPs against *Pseudomonas aeruginosa*. It has been reported that the number of total cells found on AgNP-modified implants represents only 20% of those attached to unmodified surfaces. This data suggests that the incorporation of AgNPs on Ti implants is an efficient method to protect implant surface against pathogen colonization.

As important as the antibacterial potential is the biocompatibility of these modified implants. Aiming to evaluate this property, Lu et al. [114] have tested Ti implants incorporated with different concentrations of AgNPs (0.5, 1, 1.5, 2 M). For all the tested concentrations, osteoblasts started to adhere on the coatings after 1 day of culture and spread well until 7 days of culture. However, after this, the inhibitory effect of 1 M Ag on cell proliferation became significant, suggesting that AgNP coatings with low amounts of silver were more favorable for osteoblasts growth.

## 8. Future Perspectives

As shown in the previous paragraphs, AgNPs-containing dental materials present good antimicrobial properties (Table 1). However, much is still to be discovered. One of the most important experiments to be performed is to apply the bench results on *in vivo* studies [29, 52], since laboratory conditions do not exactly reproduce oral conditions. Other aspect to be investigated is the long term effectiveness of AgNPs applied on dental materials [32, 51], whereas a long lasting antimicrobial potential of them is desirable.

## 9. Conclusions

In this review, the antimicrobial effect of AgNPs incorporation into dental materials was investigated, such as composite resin, endodontic materials, acrylic resin, and implants. Several studies have shown that silver, in its nanoparticulated form, possesses an inhibitory effect against many bacteria

TABLE I: Antimicrobial activity of AgNPs-containing dental materials.

Material studied	AgNPs concentration	Antimicrobial effectiveness	Reference
Composite resin	0.028 wt%, 0.042 wt%, 0.088 wt%, 0.175 wt%	Good inhibitory activity against <i>S. mutans</i> , at 0.042 wt%	[25]
Adhesive system	0.1 wt%	Reduction of CFU for total microorganisms, total streptococci, and <i>S. mutans</i>	[41]
Adhesive system	0.05 wt%	Reduction of CFU and acid lactic production for <i>S. mutans</i>	[32]
Primer and adhesive	0.05 wt%	Good inhibitory activity against total microorganisms, total streptococci, and <i>S. mutans</i>	[42]
Acrylic resin	1 $\mu$ g/mL	Reduction on <i>C. albicans</i> adherence	[43]
Acrylic resin	0.05 vol%, 0.5 vol% and 5 vol%	Good efficacy against <i>C. albicans</i> , at 5 vol%	[44]
Tissue conditioner	0.1%, 0.5%, 1.0%, 2.0% and 3.0%.	Antimicrobial properties against <i>S. mutans</i> and <i>S. aureus</i> at 0.1% and against <i>C. albicans</i> at 0.5%	[21]
Intracanal irrigant	0.005%	Bactericidal effect against <i>E. faecalis</i>	[20]
Gutta-percha	Not mentioned	Significant effect against <i>E. faecalis</i> , <i>S. aureus</i> , <i>C. albicans</i> and <i>E. coli</i> .	[45]
MTA	1 wt%	High antimicrobial effect against <i>E. faecalis</i> , <i>C. albicans</i> , and <i>P. aeruginosa</i>	[19]
Titanium implants	0.5 M, 1.0 M, 1.5 M, 2.0 M	Prevention of <i>S. aureus</i> adhesion for up to 30 days	[23]
Titanium implants	$3.16 \times 10^{-2}$ mg Ag/mL	Reduction of <i>P. aeruginosa</i> adhesion	[22]

and fungi, including *S. mutans*, *C. albicans*, *P. aeruginosa*, *E. faecalis*, and *S. aureus*, among others, which could decrease the occurrence of secondary caries, fungal infection, fails on endodontic treatment, and dental implant losses. Although AgNP is a promising antimicrobial to be used in dentistry, its application on some areas, as endodontics and implantology, remains scarce; thereafter, we mostly encourage studies on these fields.

AgNP has also been proved to be biocompatible with mammalian cells, suggesting that its application on dental materials does not represent a threat to human health. However, more studies are necessary to determine the optimal concentration of this silver compound, in order to guarantee the antimicrobial effect without increasing its cytotoxicity. Moreover, further studies are needed to investigate the Ag ion release and long-term properties of the new AgNP-containing dental materials. We also encourage researchers to study and elucidate the best ways of silver incorporation as well as the possible negative influence of its addition in dental materials, especially regarding color changes and mechanical properties.

## Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

## References

- [1] A. Panáček, L. Kvítek, R. Prucek et al., "Silver colloid nanoparticles: synthesis, characterization, and their antibacterial activity," *The Journal of Physical Chemistry B*, vol. 110, no. 33, pp. 16248–16253, 2006.
- [2] J. S. Kim, E. Kuk, K. N. Yu et al., "Antimicrobial effects of silver nanoparticles," *Nanomedicine: Nanotechnology, Biology, and Medicine*, vol. 3, no. 1, pp. 95–101, 2007.
- [3] S. Pal, Y. K. Tak, and J. M. Song, "Does the antibacterial activity of silver nanoparticles depend on the shape of the nanoparticle? A study of the gram-negative bacterium *Escherichia coli*," *Applied and Environmental Microbiology*, vol. 73, no. 6, pp. 1712–1720, 2007.
- [4] K.-J. Kim, W. S. Sung, S.-K. Moon, J.-S. Choi, J. G. Kim, and D. G. Lee, "Antifungal effect of silver nanoparticles on dermatophytes," *Journal of Microbiology and Biotechnology*, vol. 18, no. 8, pp. 1482–1484, 2008.
- [5] I. Chopra, "The increasing use of silver-based products as antimicrobial agents: a useful development or a cause for concern?" *Journal of Antimicrobial Chemotherapy*, vol. 59, no. 4, pp. 587–590, 2007.
- [6] V. Alt, T. Bechert, P. Steinrücke et al., "An in vitro assessment of the antibacterial properties and cytotoxicity of nanoparticulate silver bone cement," *Biomaterials*, vol. 25, no. 18, pp. 4383–4391, 2004.
- [7] U. Samuel and J. P. Guggenbichler, "Prevention of catheter-related infections: the potential of a new nano-silver impregnated catheter," *International Journal of Antimicrobial Agents*, vol. 23, no. 1, pp. S75–S78, 2004.
- [8] G. Gosheger, J. Hardes, H. Ahrens et al., "Silver-coated megaendoprostheses in a rabbit model—an analysis of the infection rate and toxicological side effects," *Biomaterials*, vol. 25, no. 24, pp. 5547–5556, 2004.
- [9] T. V. Slenters, I. Hauser-Gerspach, A. U. Daniels, and K. M. Fromm, "Silver coordination compounds as light-stable, nanostructured and anti-bacterial coatings for dental implant and restorative materials," *Journal of Materials Chemistry*, vol. 18, no. 44, pp. 5359–5362, 2008.

- [10] C. Damm, H. Münstedt, and A. Rösch, "Long-term antimicrobial polyamide 6/silver-nanocomposites," *Journal of Materials Science*, vol. 42, no. 15, pp. 6067–6073, 2007.
- [11] S. L. Percival, P. G. Bowler, and D. Russell, "Bacterial resistance to silver in wound care," *Journal of Hospital Infection*, vol. 60, no. 1, pp. 1–7, 2005.
- [12] J. R. Morones, J. L. Elechiguerra, A. Camacho et al., "The bactericidal effect of silver nanoparticles," *Nanotechnology*, vol. 16, no. 10, pp. 2346–2353, 2005.
- [13] M. Rai, A. Yadav, and A. Gade, "Silver nanoparticles as a new generation of antimicrobials," *Biotechnology Advances*, vol. 27, no. 1, pp. 76–83, 2009.
- [14] C.-N. Lok, C.-M. Ho, R. Chen et al., "Proteomic analysis of the mode of antibacterial action of silver nanoparticles," *Journal of Proteome Research*, vol. 5, no. 4, pp. 916–924, 2006.
- [15] C. Damm, H. Münstedt, and A. Rösch, "The antimicrobial efficacy of polyamide 6/silver-nano- and microcomposites," *Materials Chemistry and Physics*, vol. 108, no. 1, pp. 61–66, 2008.
- [16] D. J. Stickler, "Biomaterials to prevent nosocomial infections: is silver the gold standard?" *Current Opinion in Infectious Diseases*, vol. 13, no. 4, pp. 389–393, 2000.
- [17] M. S. Cohen, J. M. Stern, A. J. Vanni et al., "In vitro analysis of a nanocrystalline silver-coated surgical mesh," *Surgical Infections*, vol. 8, no. 3, pp. 397–403, 2007.
- [18] Z. Zhang, M. Yang, M. Huang, Y. Hu, and J. Xie, "Study on germicidal efficacy and toxicity of compound disinfectant gel of nanometer silver and chlorhexidine acetate," *Chinese Journal of Health Laboratory Technology*, vol. 17, pp. 1403–1406, 2007.
- [19] M. Samiei, M. Aghazadeh, M. Lotfi, S. Shakoei, Z. Aghazadeh, and S. M. V. Pakdel, "Antimicrobial efficacy of mineral trioxide aggregate with and without silver nanoparticles," *Iranian Endodontic Journal*, vol. 8, no. 4, pp. 166–170, 2013.
- [20] M. Lotfi, S. Vosoughhosseini, B. Ranjkesh, S. Khani, M. Saghiri, and V. Zand, "Antimicrobial efficacy of nanosilver, sodium hypochlorite and chlorhexidine gluconate against *Enterococcus faecalis*," *African Journal of Biotechnology*, vol. 10, no. 35, pp. 6799–6803, 2011.
- [21] K.-Y. Nam, "In vitro antimicrobial effect of the tissue conditioner containing silver nanoparticles," *Journal of Advanced Prosthodontics*, vol. 3, no. 1, pp. 20–24, 2011.
- [22] C. Y. Flores, C. Diaz, A. Rubert et al., "Spontaneous adsorption of silver nanoparticles on Ti/TiO<sub>2</sub> surfaces. Antibacterial effect on *Pseudomonas aeruginosa*," *Journal of Colloid and Interface Science*, vol. 350, no. 2, pp. 402–408, 2010.
- [23] L. Zhao, H. Wang, K. Huo et al., "Antibacterial nano-structured titania coating incorporated with silver nanoparticles," *Biomaterials*, vol. 32, no. 24, pp. 5706–5716, 2011.
- [24] J. Durner, M. Stojanovic, E. Urcan, R. Hickel, and F.-X. Reichl, "Influence of silver nano-particles on monomer elution from light-cured composites," *Dental Materials*, vol. 27, no. 7, pp. 631–636, 2011.
- [25] L. Cheng, M. D. Weir, H. H. K. Xu et al., "Antibacterial amorphous calcium phosphate nanocomposites with a quaternary ammonium dimethacrylate and silver nanoparticles," *Dental Materials*, vol. 28, no. 5, pp. 561–572, 2012.
- [26] L. Cheng, M. D. Weir, H. H. K. Xu et al., "Effect of amorphous calcium phosphate and silver nanocomposites on dental plaque microcosm biofilms," *Journal of Biomedical Materials Research Part B: Applied Biomaterials*, vol. 100, no. 5, pp. 1378–1386, 2012.
- [27] S. Eckhardt, P. S. Brunetto, J. Gagnon, M. Priebe, B. Giese, and K. M. Fromm, "Nanobio silver: its interactions with peptides and bacteria, and its uses in medicine," *Chemical Reviews*, vol. 113, no. 7, pp. 4708–4754, 2013.
- [28] Y. J. Cheng, D. N. Zeiger, J. A. Howarter et al., "In situ formation of silver nanoparticles in photocrosslinking polymers," *Journal of Biomedical Materials Research—Part B Applied Biomaterials*, vol. 97, no. 1, pp. 124–131, 2011.
- [29] L. Cheng, K. Zhang, M. A. S. Melo, M. D. Weir, X. Zhou, and H. H. K. Xu, "Anti-biofilm dentin primer with quaternary ammonium and silver nanoparticles," *Journal of Dental Research*, vol. 91, no. 6, pp. 598–604, 2012.
- [30] C. Fan, L. Chu, H. R. Rawls, B. K. Norling, H. L. Cardenas, and K. Whang, "Development of an antimicrobial resin—a pilot study," *Dental Materials*, vol. 27, no. 4, pp. 322–328, 2011.
- [31] T.-O. Peulen and K. J. Wilkinson, "Diffusion of nanoparticles in a biofilm," *Environmental Science and Technology*, vol. 45, no. 8, pp. 3367–3373, 2011.
- [32] F. Li, M. D. Weir, J. Chen, and H. H. K. Xu, "Comparison of quaternary ammonium-containing with nano-silver-containing adhesive in antibacterial properties and cytotoxicity," *Dental Materials*, vol. 29, no. 4, pp. 450–461, 2013.
- [33] H.-J. Park, S. Park, J. Roh et al., "Biofilm-inactivating activity of silver nanoparticles: a comparison with silver ions," *Journal of Industrial and Engineering Chemistry*, vol. 19, no. 2, pp. 614–619, 2013.
- [34] K. Chaloupka, Y. Malam, and A. M. Seifalian, "Nanosilver as a new generation of nanoparticle in biomedical applications," *Trends in Biotechnology*, vol. 28, no. 11, pp. 580–588, 2010.
- [35] E. T. Hwang, J. H. Lee, Y. J. Chae et al., "Analysis of the toxic mode of action of silver nanoparticles using stress-specific bioluminescent bacteria," *Small*, vol. 4, no. 6, pp. 746–750, 2008.
- [36] D. Seth, S. R. Choudhury, S. Pradhan et al., "Nature-inspired novel drug design paradigm using nanosilver: efficacy on multi-drug-resistant clinical isolates of tuberculosis," *Current Microbiology*, vol. 62, no. 3, pp. 715–726, 2011.
- [37] X. Chen and H. J. Schluesener, "Nanosilver: a nanoparticle in medical application," *Toxicology Letters*, vol. 176, no. 1, pp. 1–12, 2008.
- [38] A. G. R. Targino, M. A. P. Flores, V. E. dos Santos Junior et al., "An innovative approach to treating dental decay in children. A new anti-caries agent," *Journal of Materials Science: Materials in Medicine*, vol. 25, no. 8, pp. 2041–2047, 2014.
- [39] G. Chladek, A. Mertas, I. Barszczewska-Rybarek et al., "Antifungal activity of denture soft lining material modified by silver nanoparticles—a pilot study," *International Journal of Molecular Sciences*, vol. 12, no. 7, pp. 4735–4744, 2011.
- [40] M. A. S. Melo, L. Cheng, M. D. Weir, R.-C. Hsia, L. K. A. Rodrigues, and H. H. K. Xu, "Novel dental adhesive containing antibacterial agents and calcium phosphate nanoparticles," *Journal of Biomedical Materials Research, Part B: Applied Biomaterials*, vol. 101, no. 4, pp. 620–629, 2013.
- [41] M. A. S. Melo, L. Cheng, K. Zhang, M. D. Weir, L. K. A. Rodrigues, and H. H. K. Xu, "Novel dental adhesives containing nanoparticles of silver and amorphous calcium phosphate," *Dental Materials*, vol. 29, no. 2, pp. 199–210, 2013.
- [42] K. Zhang, F. Li, S. Imazato et al., "Dual antibacterial agents of nano-silver and 12-methacryloyloxydodecylpyridinium bromide in dental adhesive to inhibit caries," *Journal of Biomedical Materials Research Part B: Applied Biomaterials*, vol. 101, no. 6, pp. 929–938, 2013.
- [43] L. S. Acosta-Torres, I. Mendieta, R. E. Nuñez-Anita, M. Cajero-Juárez, and V. M. Castaño, "Cytocompatible antifungal acrylic

- resin containing silver nanoparticles for dentures,” *International Journal of Nanomedicine*, vol. 7, pp. 4777–4786, 2012.
- [44] D. R. Monteiro, L. F. Gorup, A. S. Takamiya, E. R. de Camargo, A. C. R. Filho, and D. B. Barbosa, “Silver distribution and release from an antimicrobial denture base resin containing silver colloidal nanoparticles,” *Journal of Prosthodontics*, vol. 21, no. 1, pp. 7–15, 2012.
- [45] O. Dianat and M. Ataie, “Gutta-percha coated with nanosilver particles,” Invention registered number: 56019, 2008.
- [46] T. Liu, X. Song, Z. Guo, Y. Dong, N. Guo, and X. Chang, “Prolonged antibacterial effect of silver nanocomposites with different structures,” *Colloids and Surfaces B: Biointerfaces*, vol. 116, pp. 793–796, 2014.
- [47] J. F. Hernández-Sierra, F. Ruiz, D. C. Cruz Pena et al., “The antimicrobial sensitivity of *Streptococcus mutans* to nanoparticles of silver, zinc oxide, and gold,” *Nanomedicine: Nanotechnology, Biology, and Medicine*, vol. 4, no. 3, pp. 237–240, 2008.
- [48] L. F. Espinosa-Cristóbal, G. A. Martínez-Castañón, R. E. Martínez-Martínez, J. P. Loyola-Rodríguez, J. F. Reyes-Macías, and F. Ruiz, “Antibacterial effect of silver nanoparticles against *Streptococcus mutans*,” *Materials Letters*, vol. 63, no. 29, pp. 2603–2606, 2009.
- [49] L. Zhao, P. K. Chu, Y. Zhang, and Z. Wu, “Antibacterial coatings on titanium implants,” *Journal of Biomedical Materials Research Part B: Applied Biomaterials*, vol. 91, no. 1, pp. 470–480, 2009.
- [50] K. Zhang, M. A. S. Melo, L. Cheng, M. D. Weir, Y. Bai, and H. H. K. Xu, “Effect of quaternary ammonium and silver nanoparticle-containing adhesives on dentin bond strength and dental plaque microcosm biofilms,” *Dental Materials*, vol. 28, no. 8, pp. 842–852, 2012.
- [51] M. A. S. Melo, S. F. F. Guedes, H. H. K. Xu, and L. K. A. Rodrigues, “Nanotechnology-based restorative materials for dental caries management,” *Trends in Biotechnology*, vol. 31, no. 8, pp. 459–467, 2013.
- [52] L. Cheng, K. Zhang, M. D. Weir, H. Liu, X. Zhou, and H. H. K. Xu, “Effects of antibacterial primers with quaternary ammonium and nano-silver on *Streptococcus mutans* impregnated in human dentin blocks,” *Dental Materials*, vol. 29, no. 4, pp. 462–472, 2013.
- [53] T. J. Fruits, J. A. Knapp, and S. S. Khajotia, “Microleakage in the proximal walls of direct and indirect posterior resin slot restorations,” *Operative Dentistry*, vol. 31, no. 6, pp. 719–727, 2006.
- [54] F. H. Coelho-de-Souza, G. B. Camacho, F. F. Demarco, and J. M. Powers, “Fracture resistance and gap formation of MOD restorations: influence of restorative technique, bevel preparation and water storage,” *Operative Dentistry*, vol. 33, no. 1, pp. 37–43, 2008.
- [55] S. P. Samuel, S. Li, I. Mukherjee et al., “Mechanical properties of experimental dental composites containing a combination of mesoporous and nonporous spherical silica as fillers,” *Dental Materials*, vol. 25, no. 3, pp. 296–301, 2009.
- [56] S. C. Bayne, J. Y. Thompson, E. J. Swift Jr., P. Stamatiades, and M. Wilkerson, “A characterization of first-generation flowable composites,” *The Journal of the American Dental Association*, vol. 129, no. 5, pp. 567–577, 1998.
- [57] D. C. Watts, A. S. Marouf, and A. M. Al-Hindi, “Photopolymerization shrinkage-stress kinetics in resin-composites: methods development,” *Dental Materials*, vol. 19, no. 1, pp. 1–11, 2003.
- [58] J. L. Drummond, “Degradation, fatigue, and failure of resin dental composite materials,” *Journal of Dental Research*, vol. 87, no. 8, pp. 710–719, 2008.
- [59] M. M. Zalkind, O. Keisar, P. Ever-Hadani, R. Grinberg, and M. N. Sela, “Accumulation of *Streptococcus mutans* on light-cured composites and amalgam: an in vitro study,” *Journal of Esthetic Dentistry*, vol. 10, no. 4, pp. 187–190, 1998.
- [60] L. Papagiannoulis, A. Kakaboura, and G. Eliades, “In vivo vs in vitro anticariogenic behavior of glass-ionomer and resin composite restorative materials,” *Dental Materials*, vol. 18, no. 8, pp. 561–569, 2002.
- [61] S. Imazato, “Antibacterial properties of resin composites and dentin bonding systems,” *Dental Materials*, vol. 19, no. 6, pp. 449–457, 2003.
- [62] N. Beyth, A. J. Domb, and E. I. Weiss, “An in vitro quantitative antibacterial analysis of amalgam and composite resins,” *Journal of Dentistry*, vol. 35, no. 3, pp. 201–206, 2007.
- [63] J. M. Antonucci, D. N. Zeiger, K. Tang, S. Lin-Gibson, B. O. Fowler, and N. J. Lin, “Synthesis and characterization of dimethacrylates containing quaternary ammonium functionalities for dental applications,” *Dental Materials*, vol. 28, no. 2, pp. 219–228, 2012.
- [64] E. A. Kidd, F. Toffenetti, and I. A. Mjör, “Secondary caries,” *International Dental Journal*, vol. 42, no. 3, pp. 127–138, 1992.
- [65] K. Zhang, L. Cheng, S. Imazato et al., “Effects of dual antibacterial agents MDPB and nano-silver in primer on microcosm biofilm, cytotoxicity and dentine bond properties,” *Journal of Dentistry*, vol. 41, no. 5, pp. 464–474, 2013.
- [66] F. Li, M. D. Weir, A. F. Fouad, and H. H. K. Xu, “Effect of salivary pellicle on antibacterial activity of novel antibacterial dental adhesives using a dental plaque microcosm biofilm model,” *Dental Materials*, vol. 30, no. 2, pp. 182–191, 2014.
- [67] J. L. Ferracane, “Resin composite—state of the art,” *Dental Materials*, vol. 27, no. 1, pp. 29–38, 2011.
- [68] A. M. Diaz-Arnold, M. A. Vargas, K. L. Shaull, J. E. Laffoon, and F. Qian, “Flexural and fatigue strengths of denture base resin,” *Journal of Prosthetic Dentistry*, vol. 100, no. 1, pp. 47–51, 2008.
- [69] K. Bulad, R. L. Taylor, J. Verran, and J. Fraser McCord, “Colonization and penetration of denture soft lining materials by *Candida albicans*,” *Dental Materials*, vol. 20, no. 2, pp. 167–175, 2004.
- [70] N. Boscato, A. Radavelli, D. Faccio, and A. D. Loguercio, “Biofilm formation of *Candida albicans* on the surface of a soft denture-lining material,” *Gerodontology*, vol. 26, no. 3, pp. 210–213, 2009.
- [71] H. F. Oliveira Paranhos, C. H. Silva-Lovato, R. F. de Souza et al., “Effect of three methods for cleaning dentures on biofilms formed in vitro on acrylic resin,” *Journal of Prosthodontics*, vol. 18, no. 5, pp. 427–431, 2009.
- [72] E. Budtz-Jørgensen, P. Mojon, J. M. Banon-Clément, and P. Bachni, “Oral candidosis in long-term hospital care: comparison of edentulous and dentate subjects,” *Oral Diseases*, vol. 2, no. 4, pp. 285–290, 1996.
- [73] Z. N. Al-Dwairi, “Prevalence and risk factors associated with denture-related stomatitis in healthy subjects attending a dental teaching hospital in North Jordan,” *Journal of the Irish Dental Association*, vol. 54, no. 2, pp. 80–83, 2008.
- [74] L. P. Samaranyake and R. G. Nair, “Oral *Candida* infections—a review,” *Indian Journal of Dental Research*, vol. 6, no. 3, pp. 69–82, 1995.

- [75] J. Chandra, P. K. Mukherjee, S. D. Leidich et al., "Antifungal resistance of Candidal biofilms formed on denture acrylic in vitro," *Journal of Dental Research*, vol. 80, no. 3, pp. 903–908, 2001.
- [76] L. F. Perezous, C. M. Flaitz, M. E. Goldschmidt, and R. L. Engelmeier, "Colonization of *Candida* species in denture wearers with emphasis on HIV infection: a literature review," *Journal of Prosthetic Dentistry*, vol. 93, no. 3, pp. 288–293, 2005.
- [77] C. R. Sims, L. Ostrosky-Zeichner, and J. H. Rex, "Invasive candidiasis in immunocompromised hospitalized patients," *Archives of Medical Research*, vol. 36, no. 6, pp. 660–671, 2005.
- [78] R. Rowan, M. McCann, and K. Kavanagh, "Analysis of the response of *Candida albicans* cells to Silver(I)," *Medical Mycology*, vol. 48, no. 3, pp. 498–505, 2010.
- [79] L. J. Douglas, "Candida biofilms and their role in infection," *Trends in Microbiology*, vol. 11, no. 1, pp. 30–36, 2003.
- [80] C. J. Seneviratne, L. Jin, and L. P. Samaranayake, "Biofilm lifestyle of *Candida*: a mini review," *Oral Diseases*, vol. 14, no. 7, pp. 582–590, 2008.
- [81] D. R. Monteiro, L. F. Gorup, A. S. Takamiya, A. C. Ruvollo-Filho, E. R. de Camargo, and D. B. Barbosa, "The growing importance of materials that prevent microbial adhesion: antimicrobial effect of medical devices containing silver," *International Journal of Antimicrobial Agents*, vol. 34, no. 2, pp. 103–110, 2009.
- [82] G. Dhir, D. W. Berzins, V. B. Dhuru, A. R. Periathamby, and A. Dentino, "Physical properties of denture base resins potentially resistant to *candida* adhesion," *Journal of Prosthodontics*, vol. 16, no. 6, pp. 465–472, 2007.
- [83] L. A. Casemiro, C. H. G. Martins, F. D. C. P. Pires-de-Souza, and H. Panzeri, "Antimicrobial and mechanical properties of acrylic resins with incorporated silver-zinc zeolite—part I," *Gerodontology*, vol. 25, no. 3, pp. 187–194, 2008.
- [84] Y. Abe, M. Ishii, M. Takeuchi, M. Ueshige, S. Tanaka, and Y. Akagawa, "Effect of saliva on an antimicrobial tissue conditioner containing silver-zeolite," *Journal of Oral Rehabilitation*, vol. 31, no. 6, pp. 568–573, 2004.
- [85] M. Z. Kassaei, A. Akhavan, N. Sheikh, and A. Sodagar, "Antibacterial effects of a new dental acrylic resin containing silver nanoparticles," *Journal of Applied Polymer Science*, vol. 110, no. 3, pp. 1699–1703, 2008.
- [86] H. Kong and J. Jang, "Antibacterial properties of novel poly(methyl methacrylate) nanofiber containing silver nanoparticles," *Langmuir*, vol. 24, no. 5, pp. 2051–2056, 2008.
- [87] S.-J. Ahn, S.-J. Lee, J.-K. Kook, and B.-S. Lim, "Experimental antimicrobial orthodontic adhesives using nanofillers and silver nanoparticles," *Dental Materials*, vol. 25, no. 2, pp. 206–213, 2009.
- [88] N. Okita, D. Ørstavik, J. Ørstavik, and K. Østby, "In vivo and in vitro studies on soft denture materials: microbial adhesion and tests for antibacterial activity," *Dental Materials*, vol. 7, no. 3, pp. 155–160, 1991.
- [89] A. Byström and G. Sundqvist, "Bacteriologic evaluation of the efficacy of mechanical root canal instrumentation in endodontic therapy," *Scandinavian Journal of Dental Research*, vol. 89, no. 4, pp. 321–328, 1981.
- [90] L. Fabricius, G. Dahlén, A. E. Ohman, and A. J. Möller, "Predominant indigenous oral bacteria isolated from infected root canals after varied times of closure," *Scandinavian Journal of Dental Research*, vol. 90, no. 2, pp. 134–144, 1982.
- [91] G. Sundqvist, D. Figdor, S. Persson, and U. Sjögren, "Microbiologic analysis of teeth with failed endodontic treatment and the outcome of conservative re-treatment," *Oral Surgery, Oral Medicine, Oral Pathology, Oral Radiology, and Endodontics*, vol. 85, no. 1, pp. 86–93, 1998.
- [92] J. C. Baumgartner and W. A. Falkler, "Bacteria in the apical 5 mm of infected root canals," *Journal of Endodontics*, vol. 17, no. 8, pp. 380–383, 1991.
- [93] J. C. Baumgartner, S.-U. Khemaleelakul, and T. Xia, "Identification of spirochetes (treponemes) in endodontic infections," *Journal of Endodontics*, vol. 29, no. 12, pp. 794–797, 2003.
- [94] U. Sjögren, D. Figdor, S. Persson, and G. Sundqvist, "Influence of infection at the time of root filling on the outcome of endodontic treatment of teeth with apical periodontitis," *International Endodontic Journal*, vol. 30, no. 5, pp. 297–306, 1997.
- [95] Y. Shantiaee, F. Maziar, O. Dianat, and F. Mahjour, "Comparing microleakage in root canals obturated with nanosilver coated gutta-percha to standard gutta-percha by two different methods," *Iranian Endodontic Journal*, vol. 6, no. 4, pp. 140–145, 2011.
- [96] 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.
- [97] 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.
- [98] J. Kreth, D. Kim, M. Nguyen et al., "The antimicrobial effect of silver ion impregnation into endodontic sealer against *Streptococcus mutans*," *The Open Dentistry Journal*, vol. 2, pp. 18–23, 2008.
- [99] Y. Shantiaee, O. Dianat, H. Mohammad Khani, and A. Akbarzadeh Baghban, "Cytotoxicity comparison of nanosilver coated gutta-percha with Guttaflow and normal gutta-percha on L929 fibroblast with MTT assay," *Beheshti University Dental Journal*, vol. 29, pp. 62–68, 2011.
- [100] D. Kandaswamy and N. Venkateshbabu, "Root canal irrigants," *Journal of Conservative Dentistry*, vol. 13, no. 4, pp. 256–264, 2010.
- [101] R. M. Clarkson and A. J. Moule, "Sodium hypochlorite and its use as an endodontic irrigant," *Australian Dental Journal*, vol. 43, no. 4, pp. 250–256, 1998.
- [102] J. W. Harrison, T. A. Svec, and J. C. Baumgartner, "Analysis of clinical toxicity of endodontic irrigants," *Journal of Endodontics*, vol. 4, no. 1, pp. 6–11, 1978.
- [103] I. N. Rôças, J. F. Siqueira Jr., and K. R. N. Santos, "Association of *Enterococcus faecalis* with different forms of periradicular diseases," *Journal of Endodontics*, vol. 30, no. 5, pp. 315–320, 2004.
- [104] F. Abbasipour, V. Akheshteh, A. Rastqar, H. Khalilkhani, S. Asgari, and M. Janahmadi, "Comparing the effects of mineral trioxide aggregate and calcium enriched mixture on neuronal cells using an electrophysiological approach," *Iranian Endodontic Journal*, vol. 7, no. 2, pp. 79–87, 2012.
- [105] S. Sahebi, M. Nabavizade, V. Dolatkhah, and D. Jamshidi, "Short term effect of calcium hydroxide, mineral trioxide aggregate and calcium-enriched mixture cement on the strength of bovine root dentin," *Iranian Endodontic Journal*, vol. 7, no. 2, pp. 68–73, 2012.
- [106] 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.

- [107] E. P. Hernandez, T. M. Botero, M. G. Mantellini, N. J. McDonald, and J. E. Nör, "Effect of ProRoot MTA mixed with chlorhexidine on apoptosis and cell cycle of fibroblasts and macrophages *in vitro*," *International Endodontic Journal*, vol. 38, no. 2, pp. 137–143, 2005.
- [108] R. O. Darouiche, "Treatment of infections associated with surgical implants," *The New England Journal of Medicine*, vol. 350, no. 14, pp. 1422–1429, 2004.
- [109] J. Hardes, H. Ahrens, C. Gebert et al., "Lack of toxicological side-effects in silver-coated megaprotheses in humans," *Biomaterials*, vol. 28, no. 18, pp. 2869–2875, 2007.
- [110] J. G. E. Hendriks, J. R. Van Horn, H. C. Van Der Mei, and H. J. Busscher, "Backgrounds of antibiotic-loaded bone cement and prosthesis-related infection," *Biomaterials*, vol. 25, no. 3, pp. 545–556, 2004.
- [111] D. Campoccia, L. Montanaro, and C. R. Arciola, "The significance of infection related to orthopedic devices and issues of antibiotic resistance," *Biomaterials*, vol. 27, no. 11, pp. 2331–2339, 2006.
- [112] C. R. Arciola, L. Baldassarri, D. Campoccia et al., "Strong biofilm production, antibiotic multi-resistance and high *gelE* expression in epidemic clones of *Enterococcus faecalis* from orthopaedic implant infections," *Biomaterials*, vol. 29, no. 5, pp. 580–586, 2008.
- [113] W. Chen, Y. Liu, H. S. Courtney et al., "In vitro anti-bacterial and biological properties of magnetron co-sputtered silver-containing hydroxyapatite coating," *Biomaterials*, vol. 27, no. 32, pp. 5512–5517, 2006.
- [114] X. Lu, B. Zhang, Y. Wang et al., "Nano-Ag-loaded hydroxyapatite coatings on titanium surfaces by electrochemical deposition," *Journal of the Royal Society Interface*, vol. 8, no. 57, pp. 529–539, 2011.



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