Silver Nanoparticles as Chlorhexidine and Metronidazole Drug Delivery Platforms: Their Potential Use in Treating Periodontitis

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Purpose: Periodontal disease (PD), defined as oral inflammation caused by dental plaque, is an emerging problem. PD may lead to tooth loss, and treatment options are limited. In this study, we designed, synthesized, and characterized silver nanoparticles (AgNPs) conjugated with chlorhexidine (AgNPs-CHL) or metronidazole (AgNPs-PEG-MET) to determine whether they can be used to treat PDs.

Materials and Methods: AgNPs were synthesized and characterized by transmission electron microscopy, UV-vis spectrometry, thermogravimetric analyses, and dynamic light scattering. We determined the safety and the antimicrobial and anti-inflammatory properties of synthesized AgNPs in an in vitro model of periodontitis. Antimicrobial properties were determined by measuring the minimum inhibitory concentration (MIC) and minimum biofilm eradication concentration (MBEC) on reference strains of bacteria and fungi. Human gingival fibroblast (HGF-1), murine macrophage (RAW264.7) and human foetal osteoblast (hFOB1.19) cells were used in the study. Lipopolysaccharide (LPS) was used to induce inflammation. Cytokine levels were measured using an enzyme-linked immunosorbent assay; metalloproteinase expression was measured using Western blotting.

Results: The synthesized AgNPs were spherical and narrow-dispersed with an average diameter of $13.4 \text{ nm} \pm 3.0 \text{ nm}$ in the case of AgNPs-CHL and $3.72 \text{ nm} \pm 0.72 \text{ nm}$ in the case of AgNPs-PEG-MET. Both types of AgNPs were active against bacteria and fungi. AgNPs-CHL proved to be a more potent antimicrobial agent, although they were more cytotoxic than AgNPs-PEG-MET; however, both demonstrated beneficial properties in nontoxic concentrations. AgNPs-CHL and AgNPs-PEG-MET decreased the production of proinflammatory cytokines IL-1 β , IL-6, IL-8 and TNF α . Both agents also decreased the levels of metalloproteinases MMP3 and MMP8, which may indicate that they will inhibit tissue degradation.

Conclusion: AgNPs-CHL and AgNPs-PEG-MET may be possible therapeutic options for PD, as they have antibacterial and anti-inflammatory properties. However, to fully understand the potential of AgNPs, our in vitro findings must be evaluated in an in vivo model.

Keywords: drug delivery platforms, gingival fibroblast, macrophages, osteoblasts, periodontal disease, periodontitis

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Introduction

Periodontal disease (PD) are an emerging clinical and epidemiological problem, the incidence of which is rising.¹ PD is divided into two categories: gingivitis and periodontitis. Gingivitis is defined as inflammation of the oral cavity caused by dental plaque produced by oral microflora. Periodontitis is a more severe disease which destroys the

Graphical Abstract

structure of the teeth, such as ligaments, bones, and soft tissues, and can cause tooth loss.² Oral microbiota seems to play a crucial role in PD pathophysiology. The bacteria associated with PD are predominantly Gram-negative anaerobic bacteria (ie *Porphyromonas* spp., *Treponema* spp., *Tannerella* spp. and *Aggregatibacter* spp.).³ However, the initiation and further progression of PD is strictly associated with biofilm formation and imbalances between microbiota and immune response. Furthermore, evidence on the role of these bacteria in PD development is still growing. For example, *Streptococcus mutans* are frequently isolated from patients with severe PD.⁴ A high prevalence of *Staphylococcus epidermidis* in periodontal pockets and subgingival sites in PD patients has also been noted, while the presence of *Staphylococcus aureus* is considered critical for the exacerbation of the disease.^{5,6} On the other hand, biofilm-associated *Acinetobacter* spp. are associated with suppuration and the advance of PD into a chronic state.^{7,8} Fungi such as *Candida albicans* are also indicated as critical for PD persistence and progression, although its role has not yet been established.⁹ Contemporary dentistry has yet to find an effective method of resolving inflammation caused by oral infections, possibly because the mechanism of the

disease is not yet fully understood. As the role of the formation of bacterial plaque in PD is well described, the mechanical removal of plaque and antiseptics are often used to manage these diseases. First, exfoliative and subgingival stone remedies are used. In more advanced cases, surgical procedures using materials that regenerate the tissues surrounding the tooth are necessary. Finally, special restorations are conducted to stabilize the teeth.¹⁰

In most cases, pharmacotherapy with bacteriostatic and bactericidal agents (parahydroxybenzoate, chlorhexidine, hyaluronic acid), antifungal agents (nystatin, fluconazole) and antibiotics (metronidazole clindamycin, amoxicillin) are also necessary.^{11–15} Bacteria eradication is often unsuccessful due to biofilm formation with a complex structure, which is poorly penetrated by drugs. Bacteria in biofilm are 100–1000 more resistant to treatment than planktonic ones.¹⁶ The Federal Drug Administration (FDA) has approved only a few new antimicrobial agents in recent decades.¹⁷ Treating PD is a daunting challenge, and novel therapies are needed.

More uses for nanotechnology are being discovered every year, and it is being applied in many scientific fields, including medicine. Due to miniaturization, nanoparticles

(NPs), defined as particles sized 1–100 nm, have different properties than their macro equivalent, such as a large surface-to-volume ratio, the ability to easily penetrate biological and structural barriers, etc.¹³ AgNPs have beneficial properties which may be used in dentistry and are often studied in that regard,^{12,18} primarily due to their antimicrobial efficacy, even against multidrug-resistant strains.¹⁹ Importantly, AgNPs are also effective against pathogens in biofilm.²⁰ Craciunescu et al recently reported that AgNPs biosynthesized in collagen gel demonstrate antimicrobial activity against pathogens related to PD in non-toxic concentrations.²¹ In addition to their antimicrobial activity, AgNPs may have anti-inflammatory properties, which are especially beneficial in treating PD.^{19,22–24} Reports also suggest that AgNPs have antitumor properties against oral cancer cells in in vitro models,¹⁹ and they have been successfully used as drug delivery platforms.^{25,26}

Novel therapies using AgNPs and their conjugates with drugs can provide an excellent solution to PD treatment challenges. This work reports on the results of the synthesis of conjugates of AgNPs with two drugs: chlorhexidine and metronidazole. The chlorhexidine molecules in the obtained conjugates are directly connected to the silver surface (AgNPs-CHL), whereas the metronidazole molecules are attached to the silver surface via a polyethylene glycol (PEG) linker (AgNPs-PEG-MET). We examined the cytotoxicity, antimicrobial and anti-inflammatory properties of these conjugates in PD in an in vitro model.

Chlorhexidine is an antibacterial agent commonly used in dentistry due to 'its ability to disrupt bacterial cell walls in both gram-positive and gram-negative bacterial strains.²⁷ Metronidazole, on the other hand, is effective against protozoa and anaerobic bacteria. When internalized by a pathogen, metronidazole molecules create free radicals that can destroy the DNA of microorganisms.²⁸

Recently published reports show that combining AgNPs with chlorhexidine and metronidazole produces excellent bacteriostatic and bactericidal effects.²⁹ A synergic effect was observed when AgNPs were combined with chlorhexidine to treat an infected wound.³⁰ However, to date, there has been no study of conjugates built of chlorhexidine/metronidazole attached to the surface of AgNPs. Rather than simply combining the drugs with AgNPs, conjugating them allows medicine to be delivered to the proper cells together at the same rate and enables bioactive components to be released gradually.

In this work, we synthesized and examined antimicrobial and anti-inflammatory properties of AgNPs conjugated with chlorhexidine or metronidazole to demonstrate that they can be used to treat PD. We evaluated their biological properties in periodontitis in an in vitro model.

Materials and Methods

Materials

All compounds used in our studies had a purity of > 95%, as determined by HPLC analyses. Chlorhexidine ($\geq 99.5\%$), silver nitrate (ACS reagent, $\geq 99\%$), sodium borohydride ($\geq 98\%$), metronidazole (pharmaceutical secondary standard; $\geq 98\%$), EDC (N-(3-Dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride), HS-PEG-COOH poly (ethylene glycol) 2-mercaptoethyl ether acetic acid (average $M_n=1000$ Da) and all solvents were purchased from Sigma-Aldrich (Germany). Milli-Q ultrapure water (resistivity: $18.2 \text{ M}\Omega \text{ cm}^{-1}$, Millipore-Merk) was used in all experiments. Dialysis cellulose membranes (Nadir, Bionovo) with a molecular weight cut-off (MWCO) of 10,000 Da were used to purify the synthesized NPs.

Six strains of microorganisms were used in the study. *Acinetobacter lwoffii* PCM 2235 was acquired from the Polish Collection of Microorganisms (Polish Academy of Sciences, Wroclaw), while *Candida albicans* ATCC 10231, *Staphylococcus epidermidis* ATCC 14990, *S. epidermidis* ATCC 2532, *Stenotrophomonas maltophilia* ATCC 12716 and *Streptococcus mutans* ATCC 29176 were obtained from the American Type Culture Collection (ATCC). The ATCC provided all the cell lines used in this study. Human gingival fibroblast (HGF-1), human foetal osteoblast (hFOB1.19) and murine macrophage (RAW264.7) cells were used in the study.

Synthesis of Silver Nanoparticles Coated with Chlorhexidine

Eighty milligrams (79 μmol) of chlorhexidine were placed in a round-bottom flask and 40 mL of ultra-pure water was added. The flask was immersed in a cooling ice-water bath on a magnetic stirrer and intensively stirred for 30 minutes. Afterwards, 78 μL of an aqueous solution of AgNO_3 (1M) was added and the solution was stirred for 15 minutes. A solution of NaBH_4 in water (16 mg in 10 mL), previously cooled in a refrigerator (4°C), was subsequently added to the flask one drop at a time while the mixture was vigorously stirred. As the reducing agent was added, the originally colourless mixture gradually became yellow, then light brown and finally dark brown. Once all the

NaBH₄ had been added, the mixture was stirred at a slower rate for the next 2 hours. The prepared NPs were purified of inorganic compounds (derived from the oxidation of NaBH₄) and non-attached chlorhexidine by dialysis (using ultrapure water, 1 h, MWCO 10 kDa). The yield of the reaction calculated based on the ICP MS analysis was 68%.

Synthesis of Conjugates of Silver Nanoparticles with Metronidazole

The synthesis was performed in two steps: 1) coupling metronidazole with HS-PEG-COOH via ester bonds; and 2) synthesising AgNPs using the compounds prepared in the first step as ligands and NaBH₄ as a reducing agent. The carboxyl groups in PEG molecules were activated in the esterification process by EDC molecules to improve performance.³¹

Eighty milligrams (80 µmol) of HS-PEG-COOH, 27 mg (160 µmol) of metronidazole and 34 mg (176 µmol) of EDC were placed in a round-bottom flask and dissolved in 10 mL of ultra-pure water. The flask was placed on a magnetic stirrer and stirred for 48 hours. Next, the flask was immersed in a cooling ice-water bath on a magnetic stirrer, 40 µL of an aqueous solution of AgNO₃ (1M) was added, and the mixture was stirred for 15 minutes. A solution of NaBH₄ in water (9 mg in 6 mL), previously cooled in a refrigerator, was subsequently added to the flask one drop at a time while the mixture was vigorously stirred. As the reducing agent was added, the originally colourless mixture gradually became yellow, then light brown and finally a transparent dark brown. Once all the NaBH₄ had been added, the mixture was stirred at a slower rate for the next 24 hours. Finally, the NPs were purified of inorganic compounds (derived from the oxidation of NaBH₄) and non-attached ligands by dialysis (using ultrapure water, 24 h, MWCO 10 kDa). The final solution was dark brown, and the yield of the reaction, calculated based on the ICP MS analysis, was 27%.

Characterisation of the Nanoparticles

UV-Vis Spectrophotometry

UV-Vis spectra were recorded using a Cary 50 Conc UV/Vis spectrophotometer (Varian). The spectra were recorded in aqueous solutions.³²

Dynamic Light Scattering

Dynamic Light Scattering (DLS) measurements were performed with a Zetasizer Nano series apparatus (Malvern) equipped with a He-Ne laser (4 mW) at 632.8 nm and

a thermostatted cell holder. The hydrodynamic diameters of the particles present in the aqueous solutions were measured at 25°C. The solutions were equilibrated for 10 minutes before measurement.³³

Thermogravimetric Analyses (TGA)

Thermogravimetric (TG) measurements were performed using a TA Instruments DSC Q20 apparatus (TA instruments) with thermobalance (precision ± 0.4%; minimal mass 0.02 mg) in an N₂ atmosphere with a scanning rate of 10 K min⁻¹.³²

Laser Ablation Inductively Coupled Plasma Mass Spectrometry

The concentrations of silver in the aqueous solutions of the synthesized AgNPs were determined using a high-resolution laser ablation inductively coupled plasma mass spectrometer (LA-ICP-MS; Atom ES-HR-ICP-MS-Nu Instruments). The spectrometer is connected to the Teledyne CETAC Aridus3 Desolvating Nebulizer System, a specialized liquid sample introduction accessory for ICP-MS.³³

Antimicrobial Activity

Microorganisms

All strains were stored at -80°C in ROTI Store cryo vials. Before the tests the strains were transferred to fresh Mueller-Hinton broth (Biocorp) for bacteria or RPMI-1640 (Sigma-Aldrich) for fungi and incubated for 24 hours at 37°C while being shaken at 120 rpm. The cultures were then seeded on Mueller-Hinton agar (BioMaxima) or Sabouraud dextrose agar (BioMaxima) plates and incubated as described. These agar cultures were used for further microbiological assays. Cell densities for all assays were adjusted spectrophotometrically (Multiskan GO Microplate Spectrophotometer, Thermo Fisher Scientific) at 600 and 530 nm for bacteria and fungi, respectively.

Activity Against Planktonic Cultures

Minimum inhibitory concentrations (MICs) were determined using the broth microdilution method on 96-well polystyrene plates, according to Clinical and Laboratory Standards Institute recommendations.^{34,35} To do this, initial inoculums of bacteria (5 × 10⁵ CFU/mL) in Mueller-Hinton Broth and fungi (2 × 10³ CFU/mL) in RPMI-1640 with 2% D-glucose were exposed to the ranging concentration of tested compounds (0.03125–32 µg/mL) and incubated at 37°C for 18 and 24 hours, respectively. The MIC was taken as the lowest concentration at which

noticeable growth of microorganisms was inhibited. All experiments were conducted in triplicate.³⁶

Activity Against Biofilm

Minimum biofilm eradication concentrations (MBECs) were assessed on 96-well polystyrene flat-bottom plates according to the protocol described above.^{37,38} For this purpose, after 24 hours of growth, the cultures of microorganisms were diluted to obtain a final density of 5.0×10^5 CFU/mL and 2.0×10^5 CFU/mL of bacteria and fungi, respectively. Microorganisms were diluted in Mueller-Hinton broth or RPMI-1640 with 2% D-glucose (bacteria and fungi, respectively). Briefly, 100 μ L of cell suspension was added to the test plates and incubated for 24 hours at 37°C. The wells were then rinsed three times with PBS (phosphate-buffered saline) (pH 7.4) to remove nonadherent cells. Subsequently, 100 μ L of the test compounds in a concentration range of 0.03125–32 μ g/mL were added to each well. After 24 hours of incubation at 37°C, 20 μ L of a cell viability reagent was added (resazurin, 4 g/L; Sigma Aldrich). The MBEC was read after 1 hour of incubation with shaking at 120 rpm. The determined values were recorded as the lowest concentration at which the reduction of resazurin (from blue to pink) was $\leq 10 \pm 0.5\%$ compared to the positive (100%) and negative (0%) controls. The reduction was monitored by measuring absorbance at 570 nm (reduced) and 600 nm (oxidized) using a microplate spectrophotometer (Multiskan GO Microplate Spectrophotometer, Thermo Fisher Scientific). All experiments were conducted in triplicate.³⁶

Transmission Electron Microscopy

To characterise the NPs, transmission electron microscopy (TEM) analyses were performed using a JEM 1400 JEOL Co. microscope (JEOL) at 120 kV acceleration voltage. The aqueous solution of the sample (with a concentration 1 mg/mL) was cast onto a carbon-coated copper microgrid (200 mesh) and air-dried for 24 hours.³²

A different TEM protocol was used to investigate NP internalization and ultrastructural changes in hFOB1.19 cells, *S. mutans* and *S. maltophilia*. The hFOB1.19 cells were plated in a 6-well plate. After 24 hours, the cells were incubated with AgNPs-CHL (1 μ g/mL) or AgNPs-PEG-MET (1 μ g/mL). *S. maltophilia* and *S. mutans* were grown overnight (37°C, shaken). The bacterial cells were centrifuged at 3500 rpm for 7 minutes and resuspended in a fresh Mueller-Hinton medium to obtain a dense cell suspension of 5×10^5 CFU/mL. The cells were then

exposed to NPs at MIC concentrations and incubated for 18 hours at 37°C. After treatment, bacterial cells were centrifuged at 3500 rpm for 10 minutes and washed twice with PBS. The hFOB1.19 cells, *S. maltophilia* and *S. mutans* were fixed in 2.5% electron microscopy grade glutaraldehyde (Polysciences) in 0.1 M PBS (pH 7.4), post-fixed in 1% osmium tetroxide (Polysciences), dehydrated in a graded series of ethanol (30%–100%) and embedded in Epon (Sigma). Ultrathin (65 nm) sections were cut using a Leica UC7 Ultramicrotome, stained with UranylLess (Delta Microscopies) and Reynold's lead citrate (Delta Microscopies), and examined on a Tecnai G2 Spirit BioTWIN TEM at 120 kV.³⁷

Cell Culture

HGF-1 (ATCC CRL-2014) and RAW264.7 (ATCC TIB-71) cells were cultured in Dulbecco's Modified Eagle's Medium. The hFOB1.19 cells (ATCC CRL-11372) were cultured in a 1:1 mixture of Ham's F12 Medium Dulbecco's Modified Eagle's Medium and 2.5 mM L-glutamine. Media for all cell lines were supplemented with 10% foetal bovine serum and 1% antibiotics (penicillin/streptomycin). We followed the provider's recommendations for all cell cultures. The cells were kept at 37°C in a humidified atmosphere of 5% CO₂. The cells were cultured under sterile conditions and maintained in a 75 cm² tissue culture flask. The medium was replaced every 48 hours. When confluent, the HGF-1 and hFOB1.19 cells were detached using trypsin-EDTA solution and cultured in a new flask. The RAW264.7 cells were detached by scraping them.³⁹

MTT Assay

The cells were plated in a 96-well plate, incubated overnight and treated with AgNPs-CHL or AgNPs-PEG-MET for 24 hours. The NPs solutions were prepared *ex tempore* in FBS-free media. Sonication was performed to ensure that the NPs were distributed equally. Control cells were treated with media but without FBS. AgNPs-CHL were tested in concentration ranges of 0.2–3 μ g/mL and AgNPs-PEG-MET were tested in concentration ranges of 0.2–10 μ g/mL. After 24 hours the media was supplemented with water-soluble tetrazolium salt to a final concentration 0.5 mg/mL and incubated for 2 hours. After incubation, the media was removed and crystals were dissolved in DMSO. Absorbance was measured with a Synergy H1 microplate reader (Biotek), using 540 nm and 630 nm as references. The variability of the cells is presented as

percentage of control. Absorbances were corrected with blank NPs. The MTT assay protocol used is described above.³⁹

Measurement of Cytokine Levels

The IL-1 β , IL-6, IL-8, and TNF α levels were measured in cell culture medium or cell lysate using commercially available enzyme-linked immunosorbent assays (DuoSet ELISA, R&D Systems). Cells were plated in a 6-well plate, incubated overnight and then treated with AgNPs-CHL or AgNPs-PEG-MET for 24 hours. The NPs solutions were prepared *ex tempore* in FBS-free media. Sonication was performed to ensure equal distribution of NPs. The control cells were treated with FBS-free media. Two hours later, the cells were stimulated with 1 μ g/mL of LPS (Sigma-Aldrich). AgNPs-CHL was used in concentrations of 0.5, 1.5 and 3 μ g/mL, while AgNPs-PEG-MET was used in concentrations of 1, 5 and 10 μ g/mL. After 24 hours the media were collected, aliquoted and stored at -80°C . The cells were lysed in a lysis buffer with protease inhibitors. The cell lysate was aliquoted and stored at -80°C . All tests were performed according to the manufacturers' protocols. Samples, standard curves, and negative controls were measured using a Synergy H1 microplate reader (Biotek) at 450 nm with a wavelength correction of 540 nm. The stimulation protocol for RAW264.7 was based on Moro et al⁴⁰ and the use of DuoSet ELISA was previously established by Kim and Lee.⁶

Detection of Reactive Oxygen Species

The cells were plated in a 6-well plate, incubated overnight and treated with AgNPs-CHL or AgNPs-PEG-MET for 24 hours. NPs solutions were prepared *ex tempore* in FBS-free media. Sonication was performed to ensure equal distribution of NPs. The control cells were treated with FBS-free media. Two hours later, the cells were stimulated with 1 μ g/mL of LPS (Sigma-Aldrich). AgNPs-CHL were used in concentrations of 0.5, 1.5 and 3 μ g/mL, while AgNPs-PEG-MET were used in concentrations of 1, 5 and 10 μ g/mL. After 24 hours, the media were supplemented with 10 μ M 2,7-dichlorofluorescein diacetate (DCF-DA, Sigma Aldrich). After 30 minutes the cells were detached and samples were prepared. The fluorescence of the oxidized DCF was measured using flow cytometry (excitation wavelength: 480 nm; emission wavelength: 525 nm). Data were expressed as percentage

of untreated cells using a BD FACS Calibur cytometer with dedicated software (Cell Quest Pro).³⁶

Western Blotting

The levels of metalloproteinases MMP3, MMP8 and MMP9 were measured using Western blotting. Cells were plated in 100 mm Petri dishes, incubated overnight and treated with AgNPs-CHL or AgNPs-PEG-MET for 24 hours. NP solutions were prepared *ex tempore* in FBS-free media. Sonication was performed to ensure equal distribution of NPs. The control cells were treated with FBS-free media. Two hours after NPs treatment, the cells were stimulated with 10 μ g/mL of LPS (Sigma-Aldrich). AgNPs-CHL was used in concentrations 0.5, 1.5 and 3 μ g/mL, while AgNPs-PEG-MET was used in concentrations 1, 5 and 10 μ g/mL. After 24 hours the medium was removed, the cells were washed, detached and lysed and the protein levels were measured using the Bradford method. Once the samples were prepared, they were subjected to electrophoresis; the proteins were then transferred to nitrocellulose membranes (Protran, Schleicher and Schuell BioScience) and detected using antibodies (Abcam). β -Actin (Abcam) was used as the loading control. Immunoactive proteins were detected using an enhanced chemiluminescence Western blotting detection kit (Amersham Biosciences). Protein levels were quantified using densitometry software (ImageLab, Bio-Rad). The method was previously established and described by our team,³⁹ and our stimulation protocol was based on those of Feng et al, Sun et al, De Colli et al, Li et al and Yu et al.^{41–45}

Statistical Analysis

Each experiment was replicated independently three times. Data are presented as mean $\pm$ standard deviation. Statistical analyses were performed using Prism 9 software (GraphPad), one-way analysis of variance (ANOVA) and either Tukey's or Dunnett's post hoc test. The IC₅₀ was calculated by analysing a nonlinear regression log (inhibitor) vs normalized response.³⁹

Results

In this study, we synthesized and characterized novel conjugates of AgNPs with chlorhexidine (AgNPs-CHL) and metronidazole (AgNPs-PEG-MET). We evaluated them as potential therapeutic options for periodontitis by determining their antimicrobial and anti-inflammatory potential. We

also assessed the safety of the potential application of the synthesized AgNPs conjugates as a novel nanotherapy.

AgNPs-CHL was tested as an aqueous solution with the starting concentration of silver determined from an ICP-MS analysis at 114 ppm containing 0.3 mg/mL of the conjugate. The UV-vis spectra of the synthesized AgNPs-CHL were recorded after the addition of NaBH₄ and after dialysis (Figure 1).

AgNPs-PEG-MET was employed as an aqueous solution with the starting concentration of silver determined from an ICP-MS analysis at 71 ppm containing 0.2 mg/mL of the conjugate, 15% of which consisted of metronidazole (determined from TGA).

Thin Layer Chromatography (TLC)⁴⁶ was used to verify that the nanoparticle solution was free of impurities.

Characterisation of the Nanoparticles

UV-Vis Spectroscopy and TEM

The UV-Vis spectra recorded for the aqueous solutions of AgNPs-CHL and AgNPs-PEG-MET are presented in Figure 1.

In the spectrum of AgNPs-CHL, which was recorded after adding a reducing agent to the reaction system, the intensive band of surface plasmon resonance (SPR) at 409 nm was symmetrical and narrow. However, after excess chlorhexidine had been removed (via dialysis), the observed band widened, indicating that some of the NPs increased in size during this process. This phenomenon occurred because chlorhexidine is only weakly connected with the surface, so the combination of NPs could run

during dialysis. Thus, this process cannot be carried out for a long time (aggregation was observed after 5 h).

The UV-VIS spectrum of AgNPs-PEG-MET (Figure 1) indicates that their metallic cores are spherical, small and equal in size. It was reported that for spherical AgNPs, only one band of SPR is observed, and if it is narrow, the polydispersity is low; furthermore, if the maximum SPR band is close to 400 nm, the diameter of the NPs is in the range of a few nanometres.⁴⁷

In the spectrum of AgNPs-PEG-MET recorded before excess metronidazole was removed via dialysis, a characteristic band for this compound was visible at 316 nm. This band disappeared after dialysis, as the amount of metronidazole was too small to be visible.

The SPR band in the UV-VIS spectrum recorded for AgNPs-PEG-MET solution was located at a shorter wavelength than that for AgNPs-CHL, so the first-mentioned NPs should be smaller.

The TEM results were consistent with those obtained from UV-VIS spectroscopy. Micrographs of the synthesized NPs and the histograms drawn based on them are presented in Figure 2. For the AgNPs-CHL sample, the average diameter of the NPs was $d = 13 \pm 3$ nm. It is most likely that the tiny fraction of NPs with a larger diameter ($16 \text{ nm} < d < 25 \text{ nm}$) visible in Figure 2A were responsible for the slight broadening of the SPR band (Figure 1). Thus, the widening of the SPR band results from the size dispersity of NPs visible in the microscopic images.

The average diameter of the NPs in the AgNPs-PEG-MET sample was significantly lower ($d = 3.7 \pm 0.7$ nm). The differences between the size of NPs follow their other connection of ligand with the silver surface. Chlorhexidine connects directly to the surface of the silver via the relatively weak interaction between nitrogen atoms and metal atoms. In contrast, metronidazole is connected through a PEG linker attached to the nanoparticle via the significantly stronger thiolate Ag-S bonds. These semi-covalent bonds offer good protection for silver and gold NPs against aggregation.⁴⁸ Therefore, during the synthesis of AgNPs-PEG-MET, the ligands containing metronidazole bind to the metal surface fast and stably, preventing further nanoparticle growth. It is worth mentioning that in the case of thiolate-stabilized AgNPs, bioactivity may be limited due to the metallic core of the densely covered nanoparticle. However, our results and those of other studies indicate that such AgNPs can exhibit high antimicrobial activity.^{32,49–51}

Figure 1 UV-Vis spectra of AgNPs-CHL and AgNPs-PEG-MET recorded in water solutions.

Figure 2 TEM micrographs of the synthesized NPs: (A) AgNPs-CHL; (B) AgNPs-PEG-MET; (C) AgNPs-CHL; (D) AgNPs-PEG-MET. The histograms were prepared based on these images.

DLS Analyses

DLS measurements allowed us to analyse the size and dispersity of the synthesized NPs, including the stabilization layer attached to the metal surface. The size dispersity curves (calculated by the number of NPs) and the average hydrodynamic diameters determined are shown in Figure 3. For AgNPs-PEG-MET, the average size of the NPs was 6.39 nm and the dispersity was narrow; for the AgNPs-CHL sample, the main peak was narrow, with an average of 27.93 nm, but a small fraction of larger NPs was visible. UV-VIS spectra and TEM analyses also

revealed the presence of a small fraction of larger NPs in this material (see above). It is worth adding that DLS shows that the stabilizing layer was very thick for both types of synthesized NPs, which offers better protection against aggregation. Considering the average diameter obtained using the DLS measurements (which determined the whole nanoparticle size) and that using the TEM measurements (in which only the metal core was visible), we calculated the thickness of the stabilizing layer for both synthesized NPs: AgNPs-PEG-MET was 2.67 ± 0.55 nm, while AgNPs-CHL was 14.53 ± 1.53 nm.

Figure 3 The number averaged hydrodynamic diameter distribution of the synthesized NPs obtained from DLS measurements.

TGA Analyses

The content of the medicines attached to the surface of the NPs was determined using TGA, and the results of these analyses were used to evaluate the stability of the synthesized nanomaterials. The TGA curves for the synthesized AgNPs-CHL and the drug used for their preparation are presented in Figure 4.

Thermal decomposition of chlorhexidine (CHL) was completed below 700°C and no residual mass was observed. The process comprises three main steps, with

Figure 4 TGA curves recorded during the heating AgNPs-CHL and CHL not attached on the silver surface under nitrogen atmosphere (solid lines) and their derivatives with temperature (dotted lines with the same colours).

maximums of 211°C, 470°C and 656°C. The final mass obtained after heating the AgNPs-CHL to 800°C corresponded to the silver content in this material (36% of the initial mass). Based on this value, the concentration of chlorhexidine in a solution can be easily calculated if its silver content is known (for example, from ICP MS), as the AgNPs-CHL consists of only these two components. The thermal stability of fabricated AgNPs-CHL is higher than that of pure chlorhexidine, and the main steps of its decomposition must be carried out at higher temperatures (Figure 4).

The TGA curves recorded for metronidazole, the AgNPs coated with this drug (AgNPs-PEG-MET) and the PEG derivatives used to link the silver and the drug are shown in Figure 5.

The composition of AgNPs-PEG-MET was determined using the TGA measurements. As shown in Figure 5, metronidazole decomposes completely below 300°C, while PEG begins decomposing above 300°C. The percentage of mass, 85%, corresponds to the part of the sample after decomposition of metronidazole, meaning that its content is 15% w/w. The residue above 800°C corresponds to the silver content in the sample (36%). In the sample, the PEG connecting the medicine to silver comprised 49%.

The thermal decomposition of the metronidazole attached to AgNPs occurs at a lower temperature than the non-attached (maximum decomposition occurs at 180°C in the case of AgNPs-PEG-MET and at 254°C for

Figure 5 TGA curves recorded during the heating of AgNPs-PEG-MET. MET is metronidazole not attached to the silver surface, and PEG is a derivative used to link the silver and the drug. The measurements were taken under a nitrogen atmosphere (solid lines); derivatives of the curves with temperature are represented as dotted lines with the same colours.

pure metronidazole). Decomposition probably requires less energy when these molecules are combined with PEG molecules via ester bonds. Differences are also visible in the decomposition of connected and free PEG: that of the PEG attached to the NPs requires several stages and ends at 800°C (Figure 5).

Antimicrobial Activity

Antimicrobial activity was evaluated using 5 reference strains of bacteria: 3 gram-positive strains (*S. maltophilia*, *S. mutans*, and 2 strains of *S. epidermidis*); 1 gram-negative strain (*A. lwoffii*); and 1 reference strain of fungi (*C. albicans*). The strains of microorganisms were chosen based on human oral microflora composition.

Both AgNPs-CHL and AgNPs-PEG-MET exhibited high antimicrobial activity against all the strains included in this study. The highest activity was noticed for AgNPs-CHL and *S. epidermidis*, for which growth inhibition (Table 1) and biofilm eradication (Table 2) occurred below the concentration used in the assays (≤ 0.03125 $\mu\text{g/mL}$). Generally, AgNPs-CHL was characterized by the highest activity. For other Gram-positive bacteria, MIC values were also below the concentration range; however, the MBEC values were 4 times (0.125 $\mu\text{g/mL}$). In the case of *A. lwoffii* and *C. albicans*, growth was inhibited at 0.125 and 0.25 $\mu\text{g/mL}$, respectively, and MBEC values were 8 times higher. The AgNPs-PEG-

MET inhibited the growth of all tested strains; however, contrary to AgNPs-CHL, it was more active against the Gram-negative strain and fungi. It also failed to eradicate the biofilm of *S. epidermidis* strains at the range of concentrations used in the experiments (> 32 $\mu\text{g/mL}$). The MBEC values for *S. maltophilia* and *S. mutans* were also relatively high, at 16 and 32 $\mu\text{g/mL}$, respectively. We observed a synergistic effect of drugs and NPs. AgNPs-CHL and AgNPs-PEG-MET demonstrated better antimicrobial activity than drugs alone in all cases. The antimicrobial activity of AgNPs-PEG-MET was particularly strong, as metronidazole inhibited the growth of only *A. lwoffii*, while AgNPs-PEG-MET inhibited the growth of all tested bacteria other than *S. epidermidis*. Our TEM measurements did not reveal ultrastructural changes caused by NPs in bacterial cells, but we observed that AgNPs-CHL can be found in *S. mutans* and *S. maltophilia*. Interestingly, AgNPs-PEG-MET were found to be localized around the bacterial cell wall (Figure 6).

Cytocompatibility

To ensure that the agents were safe to use, we determined the impact of NPs on the viability of human gingival fibroblasts, human foetal osteoblasts and murine macrophage cell lines using MTT assay, which measures the NAD(P)H-dependent oxidoreductase activity in mitochondria.

Table 1 Minimum Inhibitory Concentrations ($\mu\text{g/mL}$) Against Tested Strains

	<i>A. lwoffii</i> PCM 2235	<i>C. albicans</i> ATCC 10231	<i>S. epidermidis</i> ATCC 14990	<i>S. epidermidis</i> ATCC 2532	<i>S. maltophilia</i> ATCC 12716	<i>S. mutans</i> ATCC 29176
AgNPs-CHL	0.125	0.25	≤ 0.03125	≤ 0.03125	≤ 0.03125	≤ 0.03125
AgNP-PEG-MET	0.5	0.25	2	16	8	8
Chlorhexidine	0.125	2	≤ 0.03125	0.25	0.125	0.125
Metronidazole	0.125	> 32	> 32	> 32	> 32	> 32

Table 2 Minimum Biofilm Eradication Concentrations ($\mu\text{g/mL}$) Against Tested Strains

	<i>A. lwoffii</i> PCM 2235	<i>C. albicans</i> ATCC 10231	<i>S. epidermidis</i> ATCC 14990	<i>S. epidermidis</i> ATCC 2532	<i>S. maltophilia</i> ATCC 12716	<i>S. mutans</i> ATCC 29176
AgNPs-CHL	1	4	≤ 0.03125	0.125	0.125	0.125
AgNP-PEG-MET	1	4	> 32	> 32	16	32
Chlorhexidine	16	8	0.5	1	0.125	1
Metronidazole	1	> 32	> 32	> 32	> 32	> 32

Figure 6 Ultrastructure of *S. maltophilia* and *S. mutans* after treatment with AgNPs for 24 hours: (A) *S. maltophilia* control; (B) *S. maltophilia* treated with AgNPs-CHL; (C) *S. maltophilia* treated with AgNPs-PEG-MET; (D) *S. mutans* control; (E) *S. mutans* treated with AgNPs-CHL; (F) *S. mutans* treated with AgNPs-PEG-MET. The scale bar is present on the left side of each picture. Arrows indicate NPs.

AgNPs-CHL, at a concentration range of 0–0.5 $\mu\text{g/mL}$, did not impact the viability of the hFOB1.19, HGF1 and RAW264.7 cells. AgNPs-PEG-MET at a concentration range of 0–10 $\mu\text{g/mL}$ did not decrease the viability of HGF-1, hFOB1.19 or RAW264.7. The cytotoxicity of the tested NPs was type- and concentration-dependent. AgNPs-PEG-MET were more biocompatible than AgNPs-

CHL. The IC_{50} values for both types of NPs are presented in Table 3, and the results are represented graphically in Supplementary Figure 1.

To further investigate the safety of AgNPs-CHL and AgNPs-PEG-MET, we calculated the selectivity index (SI). The higher a compound's SI value, the safer it is. Mean selectivity was calculated as mean $\text{IC}_{50}/\text{MIC}$ and

Table 3 Half Maximal Inhibitory Concentration (IC₅₀) for AgNPs-CHL and AgNPs-PEG-MET

NPs Type	hFOB1.19	HGF-1	RAW264.7
AgNPs-CHL	2.332 µg/mL	2.246 µg/mL	8.425 µg/mL
AgNPs-PEG-MET	>10 µg/mL	>10 µg/mL	>10 µg/mL

mean IC₅₀ for HGF-1 cells was calculated to 2.246 (µg/mL) for AgNPs-CHL. Due to the lack of cytotoxicity for AgNPs-PEG-MET, we used IC₅₀ higher than 10 µg/mL in the calculations. We observed high SI values for tested NPs, even above 71 for AgNPs-CHL and above 40 for AgNPs-PEG-MET. Our results were similar when we used the mean of IC₅₀ for hFOB1.19 cells in our calculations. The SI values we obtained were high: above 109 for AgNPs-CHL and above 40 for AgNPs-PEG-MET. The exact SI values are presented in Table 4. TEM analysis also revealed that both types of NPs were present in hFOB1.19 cells (Figure 7). We observed that AgNPs were found in different cellular compartments. However, neither AgNPs-CHL nor AgNPs-PEG-MET caused major changes in the cells' ultrastructure. AgNPs-CHL was localized in the cytosol (Figure 7C and D), while AgNPs-PEG-MET was found in the autophagic vacuoles (Figure 7E and F). Our NPs did not affect the ultrastructure of the mitochondria or the nucleus.

Anti-Inflammatory Properties

To assess the anti-inflammatory properties of NP drugs we examined their impact on cytokine production in LPS-stimulated murine macrophages. After stimulation with LPS, we observed increased levels of IL-1β, IL-6 and TNFα secreted by RAW264.7 cells to the medium. LPS stimulation did not increase the expression of IL-8. AgNPs-CHL in concentrations of 1.5 and 3 µg/mL significantly decreased the levels of IL-1β, IL-6 and TNFα. Interestingly, 3 µg/mL of AgNPs-CHL decreased the TNFα concentration below the levels produced by unstimulated control. AgNPs-PEG-MET at a concentration of

10 µg/mL significantly decreased the levels of IL-1β and TNFα produced by stimulated RAW264.7 cells (Figure 8).

AgNPs-CHL and AgNPs-PEG-MET also decreased the intracellular levels of cytokines in stimulated RAW264.7 cells. AgNPs-CHL in concentrations of 0.5–3 µg/mL decreased levels of IL-1β and IL-8 in RAW264.7 lysate. A similar effect was observed for IL-6, but in concentrations of 1.5–3 µg/mL AgNPs-CHL. AgNPs-CHL did not impact the intracellular production of TNFα. AgNPs-PEG-MET decreased the production of IL-6 only (Figure 9).

We also assessed the impact of NP drugs on the intracellular levels of selected cytokines in LPS-stimulated HGF-1 cell lysates. However, neither AgNPs-CHL nor AgNPs-MET-PEG impacted intracellular IL-1β, IL-6, IL-8 or TNFα levels (Supplementary Figure 2).

To further examine the anti-inflammatory properties of synthesized NPs, we determined their influence on ROS production using LPS-stimulated murine macrophages (RAW264.7). As intracellular production of ROS increases when macrophages are stimulated to produce an inflammatory response,⁵² we hypothesized that decreased production of ROS in LPS-stimulated macrophages may be one of the mechanisms of AgNPs-CHL's and AgNPs-PEG-MET's anti-inflammatory properties. After stimulation with LPS, we observed a statistically significant increase in ROS production in the RAW264.7 cells. Treatment with only the highest concentration of AgNPs-CHL (3 µg/mL) increased ROS production in LPS-stimulated RAW264.7 cells. AgNPs-PEG-MET did not impact ROS production in LPS stimulated RAW264.7 cells (Figure 10). This indicates that our NPs did not exert anti-inflammatory properties on ROS production.

Impact on Metalloproteinase Expression

The destruction of tissues is important in the pathophysiology of periodontitis. Metalloproteinases are important proteins involved in this process, so we assessed the effect of AgNPs-CHL and AgNPs-PEG-MET on the expression of these enzymes. LPS stimulation increased the intracellular

Table 4 Mean SI – Selectivity index (Mean IC₅₀/MIC) for AgNPs-CHL and AgNPs-PEG-MET. The Mean IC₅₀ Value for HGF-1 Cells (Table 3) and MIC Values (Table 1) Were Used in These Calculations. Values Were Rounded to the Nearest Decimal

NPs Type	SI <i>A. lwoffii</i> PCM 2235	SI <i>C. albicans</i> ATCC 10231	SI <i>S. epidermidis</i> ATCC 14990	SI <i>S. epidermidis</i> ATCC 2532	SI <i>S. maltophilia</i> ATCC 12716	SI <i>S. mutans</i> ATCC 29176
AgNPs-CHL	17,97	8.98	>71,87	>71,87	>71,87	>71,87
AgNPs-PEG-MET	>20	>40	>5	>0.625	>1,25	>1,25

Figure 7 Ultrastructure of the hFOB1.19 cells after 24 hours of incubation with AgNPs: (A and B) control cells; (C and D) cells treated with 1 µg/mL AgNPs-CHL; (E and F) cells treated with 1 µg/mL AgNPs-PEG-MET. Arrows indicate NPs. *Autophagic vacuoles; V–vacuoles; N – nucleus; M – mitochondria. The scale bar is present on the left side of each picture.

production of MMP3, MMP8 and MMP9 in hFOB1.19 cells. AgNPs-CHL at a concentration of 3 µg/mL and AgNPs-PEG-MET at a concentration of 10 µg/mL decreased the production of MMP3 and MMP8 (Figure 11).

LPS stimulation did not significantly affect metalloproteinase production in the HGF-1 cells. However, treatment with AgNPs-PEG-MET at a concentration of 10 µg/mL decreased MMP3 levels in the stimulated HGF-1 cells ([Supplementary Figure 3](#)).

Figure 8 Effect of AgNPs-CHL and AgNPs-PEG-MET on cytokine secretion by RAW264.7 cells to medium. Cells were treated with AgNPs-drugs for 24 hours and LPS (1 μg/mL) was used to induce inflammation. Secretions: (A) IL-1β; (B) IL-6; (C) TNFα; (D) IL-8. Data are presented as mean ± SD. ###*p* < 0.001 (vs untreated control), ****p* < 0.001 (vs control treated with LPS).

Discussion

The aim of this study was to assess the properties of AgNPs as drug delivery platforms to treat PD. Two types of NPs were examined: AgNPs-CHL and AgNPs-PEG-MET. We determined that AgNPs can be effective drug delivery platforms for chlorhexidine and metronidazole. Importantly, they have a synergistic antimicrobial effect and they are non-toxic for mammalian cells at low concentrations. We also found that our NPs have anti-inflammatory properties. The conjugates of AgNPs with chlorhexidine and metronidazole were synthesized and

tested in an in vitro model for the first time. We used three cell lines which are commonly used to study PD in our experiments: HGF-1 and hFOB1.19, which were used as models of periodontal cells, and RAW264.7, which are commonly used to assess the anti-inflammatory properties of different compounds.^{53–55}

Nanoparticles as Drug Delivery Platforms

Various types of NPs have been described as drug delivery platforms. The implementation of NPs should ensure positive properties, such as reduced toxicity and cost of

Figure 9 Effect of AgNPs-CHL and AgNPs-PEG-MET on intracellular production of cytokines by RAW264.7 cells. Cells were treated with AgNPs-drugs for 24 hours and LPS (1 μ g/mL) was used to induce inflammation: (A) production of IL-1 β ; (B) production of IL-6; (C) production of TNF α ; (D) production of IL-8. Data are presented as mean $\pm$ SD. ## P < 0.01 (vs untreated control), * P < 0.05, ** P < 0.01, *** P < 0.001 (vs control treated with LPS).

treatment, while increasing drug effectiveness.⁵⁶ Most existing NPs drug delivery platforms are based on gold and AgNPs.^{57–59} We decided to use AgNPs, as published data suggests that they have stronger antimicrobial effects than gold NPs.^{60,61} NPs are often functionalized using glutathione (GSH) or polyethylene glycol (PEG) to increase their biocompatibility. PEG improves water solubility and prevents protein coating of NPs, which increases their half-life in the bloodstream.^{62,63} We therefore

decided to use PEG to link metronidazole molecules to the surfaces of the NPs.

Antimicrobial Properties

AgNPs are well known for their antimicrobial and antifungal properties.^{23,24,64} The mechanisms of this activity include release of Ag ions, DNA damage, bacterial cell wall lysis, reactive oxygen species generation and enzyme inhibition.^{65–67} AgNPs can also affect the peptidoglycan

Figure 10 Effect of AgNPs-CHL and AgNPs-PEG-MET on intracellular production of reactive oxygen species in RAW264.7 cells. Cells were treated with AgNPs-drugs for 24 hours and LPS (1 µg/mL) was used to induce inflammation. Data are presented as mean $\pm$ SD. * $P < 0.05$, ** $P < 0.01$ (vs unstimulated control).

structures of the cell wall by disrupting the β -bonds between N-acetylglucosamine and N-acetylmuramic acids.⁶⁸ We observed synergistic antimicrobial effects caused by the drugs and NPs, indicating that our NPs have strong antimicrobial properties and were more effective than drugs alone in all cases. AgNPs-CHL and AgNPs-MET-PEG were found to be effective against bacteria and fungi, both in planktonic form and as biofilm. MBEC values were higher than MIC values. This is a typical observation, as pathogens in a biofilm are more resistant to treatment due to the complex nature of biofilm.¹⁶

Different factors may influence the antibacterial properties of AgNPs. Pal et al determined that shape is an important modulator of this activity and proved that *Escherichia coli* is more susceptible to triangular AgNPs compared to spherical and rod-shaped AgNPs.⁶⁹ Raza et al proved that AgNPs' antimicrobial activity against *Pseudomonas aeruginosa* and *E. coli* is size- and shape-dependent. Interestingly, spherical AgNPs with smaller sizes were more potent antimicrobials than triangular ones.⁷⁰ AgNPs' antimicrobial activity is also functionalization dependent. Our results indicated that pegylated NPs were less effective. There are a few possible explanations for this. First, we did not examine the effect of AgNPs-PEG-MET on anaerobic bacteria strains, but metronidazole is typically more effective on these bacteria than it is on aerobic strains. Second, as mentioned earlier, functionalization by thiolate combined ligands might reduce AgNPs' antimicrobial activity.⁷¹

Data on AgNPs as antimicrobial drug delivery platforms are limited. To the best of our knowledge, this is the first

study to combine AgNPs with chlorhexidine or metronidazole. Deng et al determined that AgNPs and different antibiotics (ampicillin, penicillin, enoxacin, kanamycin, neomycin and tetracycline) had a synergistic antimicrobial effect on *Salmonella typhimurium*, but they found no chemical bond between the AgNPs and antibiotic molecules, indicating that it was simply a co-exposition. Interestingly, a mixture of AgNPs and antibiotics was found to be more effective than AgNPs or antibiotics alone.⁷² Esmaeillou et al synthesized spherical AgNPs stabilized with thioglycolic acid and conjugated with vancomycin (VAN), expecting the NPs to be effective against multi-antibiotic resistant (MDR) strains.⁷³ Combining NPs with an antibiotic significantly decreased the MIC values of the tested strains. The AgNPs-VAN were found to be more potent antimicrobial agents than AgNPs or VAN alone against *Staphylococcus aureus*, *E. coli* (VRE) or *S. epidermidis* (MRSE).⁷³ Hur et al achieved similar results, although they found that in some cases conventional antibiotics were more effective than AgNPs conjugated with VAN.⁷⁴ AgNPs were conjugated with ceftriaxone and ciprofloxacin in both studies, the results of which indicated that conjugates of AgNPs and drugs were more effective than the drugs or AgNPs alone.^{75,76} Interestingly, our previous studies showed that functionalizing some AgNPs with capping agents did not severely impact their antimicrobial activity.⁷¹

Anti-Inflammatory Properties

Both AgNPs-CHL and AgNPs-PEG-MET have anti-inflammatory properties, although those of the former are stronger. Few studies have reported on the anti-inflammatory properties of chlorhexidine. Hour-

Figure 11 Effect of AgNPs-CHL and AgNPs-PEG-MET on metalloproteinase levels in hFOB1.19 cells. Cells were treated with AgNPs-drugs for 24 hours and LPS (10 μg/mL) was used to induce inflammation: **(A)** representative Western blots; **(B)** quantification of MMP3 levels; **(C)** quantification of MMP8 levels; **(D)** quantification of MMP9 levels. Data are presented as mean ± SD. #*P*<0.05 (vs unstimulated control), **P*<0.05 (vs control treated with LPS).

Haddad et al reported that chlorhexidine can decrease the levels of TNFα and INF γ in BALB/c mice with experimentally induced PD and decrease leucocytosis.⁷⁷ We did not find any reports where metronidazole alone demonstrated anti-inflammatory properties in an *in vitro* or *in vivo* model, and some reports indicated that metronidazole does not have anti-inflammatory properties.⁷⁸

Several studies have assessed the anti-inflammatory properties of AgNPs. Two studies confirmed that AgNPs might have stronger anti-inflammatory properties than drugs (specifically, sodium diclofenac).^{79,80} Singh et al reported that 66 nm green-synthesized AgNPs inhibited the inflammatory response in LPS-stimulated RAW264.7.⁸¹ They observed decreased levels of nitric oxide (NO) production, cyclooxygenase (COX-2) mRNA, and the expression of the inflammation-related proteins p-IkB, NF-κB p-38, p-JNK and p-ERK.⁸¹ David et al also showed that 40 nm green-synthesized AgNPs could decrease the production of IL-1β in ultraviolet-light-stimulated human keratinocytes (HaCaT).⁸² AgNPs may decrease ROS, nitric oxide and TNFα production in LPS-stimulated N9 and N27 neuronal cells by the induction of hydrogen-sulfide-synthesizing enzymes.⁸³ This effect was more prominent for AgNPs than AgNO₃, which indicates that nano-size is an important modulator of biological properties.⁸³ Ten-nanometre AgNPs were also reported to decrease levels of TNFα and IL-12 and to decrease the expression of COX-2 in normal human dermal fibroblasts and normal human keratinocytes.⁸⁴ However, Huang et al reported that 3–5 nm AgNPs may trigger inflammation in murine microglia (BV-2) and mouse neuroblastoma cells (N2a): they observed increased levels of IL-1β and translations of proinflammatory genes.⁸⁵ These observations about the anti-inflammatory properties of AgNPs were also confirmed in *in vivo* studies.^{86,87} Similarly, glutathione-stabilized AgNPs may increase the production of IL-6, IL-8 and TNFα in HGF-1 cells.⁵⁵ TNFα expression may also be increased when RAW264.7 cells are treated with 68.9 nm AgNPs.⁸⁸ Interestingly, a recent study examined the role of highly biocompatible biomimetic scaffold embedding AgNPs in the treatment of oral lesions.⁸⁹ Synthesized material demonstrated both antimicrobial and anti-inflammatory properties, inhibiting the growth of *F. nucleatum* and *P. gingivalis* and suppressing production of IL-6, IL-8 and TNFα in stimulated THP1 cells (human monocytic leukaemia).⁸⁹

We also observed that AgNPs-CHL and AgNPs-PEG-MET decreased levels of MMP3 and MMP8, which suggests that AgNPs inhibit tissue degeneration and may slow disease progression. Both drugs used in this study can impact MMP activity. Chlorhexidine inhibits MMP2, MMP8 and MMP9,⁹⁰ while metronidazole decreases the activity of proMMP-3.⁹¹ Frankova et al reported that 10 nm AgNPs did not affect MMP1 and MMP3 expression in normal human dermal fibroblasts and normal human keratinocytes.⁸⁴ AgNPs 68.9 nm in size may increase

mRNA levels of MMP3, MMP11 and MMP19 in RAW264.7 cells.⁸⁸ AgNPs embedded in a specific polysaccharide (EPS) can decrease the enzymatic activity of MMP2 MMP9 from SKBR3 (human breast adenocarcinoma) cells.⁹² Two- to four-nanometre AgNPs can also reduce the enhancement of MMP1 and MMP2 from platelet-rich plasma. This effect was capping-agent-dependent and glutathione stabilized, and pegylated AgNPs were found to be more potent than AgNPs capped with liponic acid.⁹³

Cytocompatibility

The synthesized conjugates exhibited relatively low cytotoxicity at relatively high concentrations. Importantly, AgNPs-CHL and AgNPs-PEG-MET have beneficial properties in nontoxic concentrations. Several reports have indicated that AgNPs can decrease the viability of human cells. AgNPs are cytotoxic to cell types such as gingival fibroblasts, osteoblasts, dental pulp, muscles and others found in the oral cavity and gastrointestinal tract.^{36,94–96} The main cytotoxicity mechanisms of AgNPs are mitochondrial disruption and ROS generation, which lead to DNA and protein damage.⁹⁷ AgNPs can also affect membrane potential and have a lytic effect.⁹⁷ Dysfunction of important cellular processes may cause homeostasis imbalance and programmed cell death.

Several factors influence AgNPs cytotoxicity. Smaller AgNPs are typically more cytotoxic than larger ones.⁹⁸ Their shape is also important. Soleimani et al reported that AgNPs with sharper geometry were more cytotoxic, while spherical ones had the least effect on cell viability.⁹⁹ Functionalization may also impact the safety profile of AgNPs, which is essential in our case. We noticed that AgNPs-PEG-MET had a weaker effect on cell viability than AgNPs-CHL. Pegylated AgNPs are usually safer than non-thiolate-functionalized AgNPs, which was proven by our team in several cellular models (gingival fibroblasts, human umbilical vein endothelial cells and human platelets).^{71,93} This observation is consistent with the data reported in the literature. Soenen et al proved that pegylated gold NPs (AuNPs) have a lesser effect on cell morphology, ROS generation and cell death induction in HUVEC cells than non-pegylated ones.¹⁰⁰ Yu et al reported a similar observation, but for iron oxide NPs.¹⁰¹

Several studies have suggested that AgNPs can be internalized by eukaryotic cells. Diffusion through cellular membrane, endocytosis or phagocytosis are thought to be the main internalization mechanisms of AgNPs.⁹⁷

Sengstock et al proved that human mesenchymal stem cells can uptake 80-nm AgNPs, which were found inside these cells but not in the nucleus.¹⁰² A similar observation was described for 5-nm AgNPs, but for human mesenchymal stem cells and osteoblasts.¹⁰³ However, 30-nm AgNPs can enter HaCaT cells and penetrate the nucleus.¹⁰⁴ We observed that AgNPs-CHL and AgNPs-PEG-MET were internalized by the cells we tested, but the NPs did not penetrate the nucleus and did not cause major ultrastructure changes.

Interestingly, pegylated NPs were found in the autophagic vacuoles, whereas AgNPs-CHL were localized in the cytosol. Guo et al recorded a similar observation, although they used a higher concentration than we did.¹⁰⁵ This phenomenon may be partially explained by the differences in cytotoxicity of the two types of NPs. Enclosing AgNPs-PEG-MET in autophagic vacuoles may limit their contact with other organelles, thus having a protective effect, and the internalization of the NPs may coat them with proteins, which will influence their surface properties.¹⁰⁶

Safety of Potential Applications

We are aware that the cytotoxicity of AgNPs-CHL may be concerning, but it must be remembered that no clinically used substances are free of side effects. AgNPs may thus be beneficial drug delivery platforms. Due to the synergistic effects of drugs and AgNPs, lower concentrations of pharmaceuticals may be used, which will potentially decrease their side effects. We also observed high selectivity index values. There is no consensus regarding which SI values should be the cut-off point: some studies suggest that compounds with an SI > 3 are worth further investigation, while others suggest that an SI > 10 is preferable.^{107,108} Both types of AgNPs used in this study have at least two SI values > 10.

The antibacterial agents used in this study, chlorhexidine and metronidazole, are cytotoxic. Babich et al examined the impact of chlorhexidine on epithelial gingival cell line (S-G), human gingival fibroblast (HGF-1) and found that chlorhexidine decreased the viability of the cell lines tested in a concentration- and time-dependent manner by disrupting cell membranes and calcium homeostasis disruption.¹⁰⁹ Hidalgo et al reported that chlorhexidine could decrease ATP production and proliferation rate in human dermal fibroblasts.¹¹⁰ Even as concentrations of chlorhexidine as low as 0.0001% depleted ATP production in the cells.¹¹⁰ Chlorhexidine can also cause apoptosis and

necrosis and damage DNA and ROS generation, which was proven in a RAW264.7 macrophage model.¹¹¹

Ferreira et al reported time- and concentration-dependent metronidazole cytotoxicity against human gingival fibroblasts (FMM1). Concentrations of metronidazole as low as 5 mg/L decreased the viability of FMM1 cells. However, visible changes in cell morphology were observed following treatment with 150 mg/L.¹¹²

Conclusion

We designed and synthesized novel conjugates of AgNPs with chlorhexidine or metronidazole. AgNPs-CHL and AgNPs-PEG-MET exhibit potent antimicrobial, antibiofilm and anti-inflammatory effects. The safety of their potential clinical usage was evaluated in in vitro models, which showed that NPs may have beneficial properties in non-toxic concentrations. Our results prove that AgNPs are effective drug delivery platforms for chlorhexidine and metronidazole. AgNPs-CHL and AgNPs-PEG-MET may be possible therapeutic options for PD. Our findings need to be evaluated in in vivo studies.

Abbreviations

AgNPs-CHL, silver nanoparticles conjugated with chlorhexidine; AgNPs-PEG-MET, silver nanoparticles stabilized with poly(ethylene glycol) and conjugated with metronidazole; AgNPs, silver nanoparticles; DLS, dynamic light scattering; IC₅₀, half-maximal inhibitory concentration; IL-1 β , interleukin1 β ; IL-6, interleukin 6; IL-8, interleukin 8; LPS, lipopolysaccharide; MBEC, minimum biofilm eradication concentration; MIC, minimum inhibitory concentration; MMP, metalloproteinase; NPs, nanoparticles; PD, periodontal disease; PEG, poly(ethylene glycol); ROS, reactive oxygen species; SI, selectivity index; TEM, transmission electron microscopy; TGA, thermogravimetric analysis; TNF α , tumour necrosis factor.

Ethical Statement

According to local and European Union law, this study protocol did not require ethical approval from any institutional board. We received written consent for Somersault1824's team (I. Lahortiga, and L. Cox) to use the images they designed under a CC BY open access license in all printed and digital formats.

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Disclosure

The authors report no conflicts of interest in this work.

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