Synthesis of Functional Silver Nanoparticles and Microparticles with Modifiers and Evaluation of Their Antimicrobial, Anticancer, and Antioxidant Activity

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Abstract: An accumulating body of evidence reports the synthesis and biomedical applications of silver nanoparticles. However, the studies regarding the use of maleic acid and citric acid in the synthesis of nano-sized silver particles (AgNPs) and micro-sized silver particles (AgMPs) as well as their antibacterial, antifungal, and anticancer activities have not been reported. In the current study, we synthesized AgNPs and AgMPs using maleic acid and citric acid as capping agents and have characterized them by UV-Vis, energy-dispersive X-Ray spectroscopy (EDS), X-Ray diffraction (XRD), and scanning electron microscope (SEM) analysis. The capped silver particles were examined for their antimicrobial activity and cytotoxicity against bacteria, fungi, and brine shrimp. Additionally, the anticancer activity of these particles was tested against human breast and liver cancer cell lines. The free radical scavenging activity of capped silver particles was evaluated by 2,2-diphenyl-1-picrylhydrazyl (DPPH) assay. SEM analysis revealed a round plate-like morphology of maleic acid capped particles with an average size of $39 \pm 4$ nm, whereas citric acid capped particles display flower-shaped morphology with rough surfaces and an average size of $250 \pm 5$ nm. The uncapped AgMPs were hexagonal with $500 \pm 4$ nm size. EDS and XRD analysis confirmed the presence of Ag and face-centered cubic crystalline nature, respectively. Functionally, capped silver particles exhibited antibacterial activity against Gram-positive (Staphylococcus aureus, Bacillus subtilis, and Micrococcus luteus) and Gram-negative bacteria (Salmonella typhimurium, Enterobacter aerogenes, and Agrobacterium tumefaciens). The bactericidal activity was more active against Gram-negative bacteria with minimum inhibitory concentration (MIC) as low as 5 ppm as compared to 25 ppm for Gram-positive. Similarly, the silver particles demonstrated antifungal activity by inhibiting the growth of five fungal strains (Mucor species, Aspergillus niger, Aspergillus flavus, Aspergillus fumigatus, and Fusarium solani) up to 50% at the concentration of 500 ppm. Additionally, these particles showed substantial toxicity against brine shrimp and also significantly inhibited the
proliferation of breast cancer (MCF7) and liver cancer (HePG2) cell lines (IC₅₀ 8.9–18.56 µM). Uncapped AgMPs were less effective, inhibiting only the proliferation of MCF7 cells with IC₅₀ 46.54 µM. Besides cytotoxicity, these particles acted as potential antioxidants, showing free radical scavenging up to 74.4% in a concentration-dependent manner. Taken together, our results showed that the modifiers affect the shape and size of silver particles and may, in part, contribute to the antimicrobial and antioxidant activity of silver particles. However, the contribution of maleic acid and citric acid in enhancing the antimicrobial, anticancer, and antioxidant potential independent of silver nano and microparticles needs to be studied further. In vivo experiments may determine the therapeutic effectiveness of silver particles capped with these modifiers.

**Keywords:** silver nanoparticles; silver microparticles; functional nanomaterials; anticancer; antimicrobial; antioxidant; cytotoxicity; therapeutic efficacy

1. Introduction

The need for achieving materials of preferred morphology and architecture at a nanometric scale holds great importance largely because methods used in manufacturing influence the therapeutic properties of nanoparticles strongly [1,2]. In the last few years, rigorous research has been dedicated to the fabrication of metal nanostructures due to their distinctive characteristics different from their source materials [3]. Amongst the metal nanomaterials, silver nanoparticles (AgNPs) have emerged as a playful delivery platform in the field of nanotechnology. These important materials have attained considerable interest lately because of their good conductivity [4], chemical stability [5], catalytic [6], and antimicrobial activity [7,8]. It has been shown that AgNPs exhibit strain-specific antimicrobial activity against different pathogenic bacterial strains [9], as well as against toxigenic species of fungi [10,11]. Similarly, due to their distinctive features, these nanoparticles have prognostic applications against various diseases, such as those caused by bacteria, fungi, and viruses, including tuberculosis, acquired immunodeficiency syndrome, and retinal neovascularization [12]. Lately, AgNPs are being proposed as treatment options for cancers due to the anticancer activities of these nanoparticles [13,14]. It has also been reported that their accumulation in rat tissues and organs, which may pose a risk to certain cell populations in remote sites [15], could be translocated to the blood circulation and distributed to organs [16]. However, it has also been shown that the silver content of AgNPs in rat tissues gradually decreases during the 4-month recovery period, indicating tissue clearance of the accumulated silver [17].

The resistance developed by pathogens against antibiotics and other available drugs, so-called antibiotic resistance or drug resistance, currently poses a serious health concern. The combination of AgNPs with antibiotics may have profound antibacterial effects against drug-resistant strains [18]. It has been proposed that by merging modern technologies, such as nanotechnology with the material science, i.e., utilizing the metals’ inherent antimicrobial activity, could overcome such challenges. Metal and metal oxide nanoparticles exemplify a group of materials, which are being explored with respect to their antimicrobial effects. It has been shown that the particle size was the critical factor, which exhibits the antimicrobial efficacy of the metal nanoparticles [19].

Production of novel silver nanoparticles with improved activity for medical uses is a matter of endless concern [20]. Various methods have been established to synthesize silver and silver oxide nanoparticles. For instance, the wet chemical approach and chemical reduction in silver ions in aqueous solutions with or without stabilizing agents represent the methods tailored with definite morphology and controlled biochemical properties as well as antimicrobial activities [21,22].

The stabilizing agents or modifiers play an important role in preventing the clustering of particles in the suspension. Non-ionic polymers lead to steric stabilization of the particles by forming a coating on their surface, whereas ionic polymers add further electrostatic stability to the colloids [23]. Some polymers, such as polyvinylpyrrolidone (PVP), or organic salts, such as vinyl sulfonate, exhibit a tremendous capacity to regulate the growth of inorganic crystallites, including silver.
Therefore, the molecules which potentially control the particle dimensions and, consequently, their self-organization, are considered to be very important [24]. Previously, citrate capped AgNPs have been shown to exhibit inhibitory effects against planktonic and sessile bacteria and cytotoxic effect against osteoblasts [25].

Recent reports demonstrate that a photodynamic therapeutic agent conjugated with fatty acids, such as oleic acid (OA), is effective against metastatic cancer cells [26]. Of particular note, the AgNPs can be incorporated into polymers, which act as deposits of silver (Ag) ions, and show their activity upon exposure or when released into the medium [27,28]. AgNPs-containing polymer composites are currently being produced for their therapeutic use and are found to be active with antimicrobial and cytotoxic effects [22]. A limited number of studies have reported the synthesis of AgNPs with acid copolymers alone [29,30] or with fruit extracts [31]. However, studies regarding the synthesis of AgNPs capped with maleic acid and citric acid are scarce. The antibacterial, antifungal, and anticancer activities of maleic acid and citric acid capped AgNPs have not been reported.

In the current study, silver nanoparticles (AgNPs) and silver microparticles (AgMPs) were prepared and characterized using two different modifiers, such as maleic acid and citric acid, and were named citric acid-capped or uncapped AgMPs and maleic acid-capped AgNPs. The antimicrobial activity and cytotoxicity effects of AgNPs and AgMPs were evaluated against bacteria, fungi, and brine shrimp and on human breast and liver cancer cell lines. Additionally, the free radical scavenging activity of these particles was also performed. Our results show that silver particles exhibit antibacterial activity against three different species of Gram-positive and Gram-negative bacteria and also inhibited the growth of at least five different fungal strains. Their cytotoxicity was also significant against brine shrimp, breast cancer (MCF7), and liver cancer (HePG2) cell lines. Besides cytotoxicity, these nanoparticles acted as antioxidants in a dose-dependent manner. Overall, maleic acid-capped silver nanoparticles were found more effective compared to citric acid-capped and uncapped particles. These modified silver particles could be effective candidates to be explored further in the drug delivery experiments.

2. Materials and Methods

2.1. Synthesis of Silver Nanoparticles and Microparticles

All the chemicals of analytical grade, including silver nitrate (AgNO3), iron (II) sulfate heptahydrate (FeSO4·7H2O), maleic acid (C4H4O4), and citric acid (C6H8O7), were purchased from Sigma–Aldrich (Sarawak, Malaysia). To synthesize nano- and micro-sized silver particles (i.e., AgNPs and AgMPs), 0.1 M silver nitrate solution was prepared in 100 mL volume, and then iron sulfate heptahydrate was added to the silver nitrate solution to give a final concentration of 0.05 M of iron sulfate heptahydrate, as described previously [32]. Then, the maleic acid and citric acid were added to the solution in separate beakers to attain the final concentration of (0.02 M) under vigorous shaking using the magnetic stirrer at room temperature for 30 min. During this process, the change in color was observed. After centrifugation at 6000 rpm for 30 min and washing with distilled water three times, the nanoparticles were dried in the hot air oven at 60 °C, and the resultant powder was saved for further analysis.

2.2. Characterization of Silver Nanoparticles and Microparticles

The optical absorption features of AgNPs and AgMPs were recorded in the range of 300 to 600 nm wavelength by UV-visible spectrophotometer (UV 1602 BMS spectrophotometer, Spain). The size and morphology of the samples were investigated using scanning electron microscopy (SEM) (TESCAN SEM, VEGA 3, Warrendale, PA, USA), operating at the voltage of 20 kV (maximum) with the counting frequency of 2368 cps (maximum). Chemical composition was confirmed by energy-dispersive X-ray (EDS) (Oxford Instruments, Concord, MA, USA) coupled with the SEM as plugin hardware. Magnified micrographs were taken up to the resolution of 10 μm in the scale bar. The samples for X-ray diffraction (XRD) analysis were performed by taking a small amount of solution and drying it on a quartz plate (XRD D8 Advance, Bruker GmbH, Germany). Furthermore, the
crystallite parameters of prepared AgNPs and AgMPs were calculated using following Debye Scherrer’s relation,

\[ D = \frac{0.9 \lambda}{\beta \cos \theta} \]  \hspace{1cm} (1)

where D represents crystalline domain size perpendicular to the reflecting planes, \( \lambda \) is the wavelength of incident X-ray (1.5406 Å), \( \beta \) is the angular full width at half maximum in radians, and \( \theta \) is the angle of diffraction also called Bragg’s angle [33].

2.3. Antibacterial Assay

The bactericidal property of the synthesized AgNPs and AgMPs was investigated by the disc diffusion method following the reported methodology [34]. There were seven bacterial strains used in the study, including three Gram-positive, such as Staphylococcus aureus (ATCC 6538), Bacillus subtilis (ATCC 6633), and Micrococcus luteus (ATCC 10240), and four Gram-negative, i.e., Escherichia coli (ATCC 15224), Salmonella sethual (ATCC 19196), Enterobacter aerogenes (ATCC 13048), and Agrobacterium tumefaciens (AT-10). Briefly, bacterial suspension (100 µL) cultured in Luria Broth (10^5 to 10^7 CFU/mL of bacterial cultures) was spread on a nutrient agar plate. Then, the discs supplemented with synthesized particle suspension were placed on the labeled plates to determine the antibacterial properties at different concentrations (100, 50, 25, 5, 2.5 ppm). Further incubation was carried out at 37 °C in the incubator (WIG-105, Focused Photonics Inc., Hangzhou, China) for 24 to 48 h. The zones of inhibition were calculated after incubation and minimum inhibitory concentration (MIC) values were determined [35,36].

2.4. Antifungal Assay

The antifungal assay was performed by the agar tube dilution method, as reported earlier, against five fungal strains, including Mucor species, Aspergillus niger, Aspergillus flavus, Aspergillus fumigatus, and Fusarium solani. Briefly, 4 mL of Sabouraud dextrose agar medium was poured into test tubes with screw caps and subjected to the autoclave. After cooling to 50 °C, the media was loaded with 100 µL of particles’ stock suspension to make 500 ppm final concentration. Fungal strains were refreshed on sterile Sabouraud Dextrose Agar, and the resultant spores were inoculated into sterile normal saline, and the turbidity was adjusted to a 0.5 McFarland standard equivalent to yield 1 × 10^6 CFU/mL. The suspension was further diluted in a 1:10 ratio in Sabouraud Dextrose Broth to give a turbidity of 5 × 10^5 CFU/mL [37]. One hundred microliters of this suspension was added to the tubes, which were subjected to a slanted position and allowed to solidify at room temperature. Inoculation of fungal cultures was done in each tube from a 7-day-old fungal culture. Distilled water and terbinafine were added to the media to be used as negative and positive control, respectively. Incubation of the tubes was carried out at 28 °C in the incubator (WIG-105, Photonics Inc., Hangzhou, China) for 7 days and results were recorded by measuring linear growth (cm), and growth inhibition was calculated with reference to the negative control with the given formula;

\[ \text{Growth inhibition} = \frac{100 \times \text{Fungal growth in the negative control} - \text{fungal growth in the tube containing sample}}{\text{fungal growth in the negative control}} \]  \hspace{1cm} (2)

2.5. Brine Shrimp Lethality Assay for Cytotoxicity Screening

The preliminary cytotoxicity of synthesized AgNPs and AgMPs was evaluated by the brine shrimp lethality assay [38]. Artificial seawater was prepared by dissolving 34 g sea salt in one liter distilled water with continuous stirring. This was further utilized for hatching Brine shrimp (Artemia salina) eggs between 21 and 28 °C. After getting the hatched eggs, 5 mL of seawater was added to the labeled vials along with 100, 50, 25, 5 µM final concentrations of each test sample. After 2 days, hatched shrimp were transferred to these vials (15 shrimp per vial). The number of surviving shrimps was counted after 24 h [38]. Percentage viability was calculated by the following formula.
peak indicates the formation of polydispersed AgNPs spectrum analysis, which showed a strong indicating the synthesis of with the modifiers, the visually confirmed.

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Results

2.6. MTT Assay on Human Cancer Cell Lines

MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide) assay was performed to test the cytotoxic effect of synthesized AgNPs and AgMPs on breast cancer (MCF7) and liver cancer (HePG2) cell lines (ATCC: CCL-2™ and HB-8065™, respectively) according to the reported procedure [39]. Briefly, cells were cultured in DMEM supplemented with 10% FBS and 1% antibiotics (streptomycin and penicillin). All the components were purchased from Gibco (Invitrogen, Carlsbad, CA, USA). Cells at a density of ~100,000 were seeded in 96-well flat-bottom plates (Corning, USA) and incubated for 24 h at 37 °C in the incubator (Shel Lab, Cornelius, OR, USA) with 5% CO₂ supply. After 24 h, the test material (in triplicate) was added at different concentrations (50, 40, 30, 20, 10 µM) to the wells containing the MCF7 and HePG2 cell lines. Untreated cells and the blank samples (cells with DMSO used as solvent) were used as controls. The plate was incubated again at 37 °C with 5% CO₂ in the incubator for 24 hrs. Cell viability was then determined as follows: 10 µL MTT reagent (5mg/mL) was added to each treated sample, blank, and untreated cells and incubated at 37 °C for 3 h. After that, 100 µL of the solubilization solution (to dissolve formazan crystals) was added to the wells, and incubation was done in the dark at room temperature for 2-4 h. Finally, the absorbance of the samples was measured at 570 nm in the plate reader FLUO star Omega (BMG Labtech, Ortenberg, Germany).

2.7. 2,2-diphenyl-1-picrylhydrazyl (DPPH) Free Radical Scavenging Assay

The free radicals scavenging activity of AgNPs and AgMPs was evaluated using the 96-well plate, following the method reported earlier [40]. Each nanoparticle sample with a final concentration of 80, 40, 20 µM was mixed with 0.1 mM DPPH (190 µL) in a 96-well microtiter plate. Deionized water and ascorbic acid were used as negative and positive controls, respectively. The reaction was carried out in triplicate, and mixtures were incubated in the dark at 37 °C for 30 min, and then the absorbance of samples was measured at 517 nm by a microplate reader (BioTek, Elx 800, Shoreline, WA, USA). IC₅₀ was calculated using table curve software (version 4), and the free radical scavenging potential of the sample was calculated with the help of the following formula.

\[
\text{Percentage scavenging} = \frac{\text{Absorb. control} - \text{test Absorb.}}{\text{Absorb. control}} \times 100
\]

2.8. Statistical Analysis

All experiments and measurements were carried out in triplicate, and the values are presented as the mean ± standard error of the mean. The collected data were statistically analyzed using two-way ANOVA with the GraphPad Prism (V.5) software. The p-value of less than 0.05 was considered statistically significant.

3. Results and Discussion

3.1. Characterization of Silver Nanoparticles and Microparticles

3.1.1. Confirmation of Synthesis

First, the synthesis of silver nanoparticles (AgNPs) and silver microparticles (AgMPs) was visually confirmed. When silver nitrate and the reductant salts were mixed in particular ratios along with the modifiers, the color of the reaction turned dark grey within 10 min with continuous stirring, indicating the synthesis of silver particles. Then the synthesis was further confirmed by UV-Vis spectrum analysis, which showed a strong, broad peak at 400 nm, whereas the broadening of the peak indicates the formation of polydisperse AgNPs and AgMPs (Figure 1A). UV–Vis spectroscopy

\[
\text{Percentage viability} = \frac{\text{Control} - \text{test}}{\text{control}} \times 100
\]
is extensively used for structural characterization of silver particles, which are known to exhibit a UV–visible absorption maximum in the range of 400–500 nm because of surface plasmon resonance excitation [34]. The peak of the citric acid-capped and uncapped AgMPs was broader than that of the maleic acid-capped AgNPs. This difference in peak could be because of their larger size range [41]. The spectrum did not contain any additional peak, thus ruling out the presence of contaminants.

Generally, for spherical shaped nanoparticles, the single peak in the UV-Vis spectra is observed [42]. The appearance of two peaks in the case of citric acid capped and uncapped AgMPs could be attributed to the anisotropic morphologies of the particles. Nanoparticles contain optical properties that are sensitive to shape size agglomeration state, concentration, and surface chemistry of nanoparticles, which makes UV-Vis spectroscopy a valuable tool for the identification and characterization of nanoparticles.

3.1.2. Confirmation of Elemental Composition

Additionally, the elemental composition of the silver particles was examined by an energy-dispersive X-ray spectrocope (EDS) equipped with an SEM. The EDS profile showed a strong silver signal indicating the silver (Ag) as a major component of synthesized particles. The interpretation of the EDS spectrum was that synthesized samples are pure silver (with distinct Ag peaks) without contaminated elements. The elemental composition of particles predicted by EDS is shown in Figure 1B and the Supplementary Figures 1–3. Peaks matching other than silver are due to the elements of the carbon-coated copper grid in the SEM measurement, and peaks corresponding to Fe maybe because of the residues of iron sulfate heptahydrate left in the suspension.

![Figure 1](image)

**Figure 1.** UV-Vis spectroscopy and energy-dispersive X-ray spectroscopy (EDS) analysis of synthesized silver nanoparticles and microparticles. (A) The spectra show the absorption peaks of maleic acid capped AgNPs (red), citric acid capped AgMPs (green), and uncapped AgMPs (blue) in the range of 350–450 nm; (B) The spectrum shows silver (Ag) as the major component of nanoparticles and microparticles. AgNPs: silver nanoparticles, AgMPs: silver microparticles.

3.1.3. Characterization for Structure and Size

The structural features regarding phase composition, the crystallographic orientation of AgNPs and AgMPs were further characterized by X-ray diffraction (XRD), which has long been used to outline and identify bulk materials and nanoparticles [43]. The identified peaks [38.23 (1 1 1), 44.41 (2 0 0), 64.38 (2 2 0), 77.5 (3 1 1)], revealed the crystalline nature of synthesized particles (Figure 2). The XRD outcome was found in accordance with the standard ICSD No. 98-018-0878 [40,44]. Furthermore, through Debye Scherrer’s equation and broadening of the Bragg’s peaks, it was revealed that the approximate crystallite size of maleic acid capped AgNPs are 6 nm as compared to citric acid capped AgMPs of 15 nm and uncapped AgMPs of 19 nm. These findings confirm that the synthesized AgNPs and AgMPs are made up of multiple smaller crystallite units.
Figure 2. X-ray diffraction (XRD) analysis of synthesized silver nanoparticles and microparticles. The spectrum shows face-centered cubic (FCC) crystalline metallic silver nanoparticles (AgNPs) and microparticles (AgMPs). The intensity on the vertical axis is measured in counts per second (CPS), and the diffraction angle (2 theta) measured is taken along the horizontal axis. The value of wavelength in angstrom (a.u.) is also indicated.

3.1.4. Morphology by Scanning Electron Microscopy

The morphology of the synthesized AgNPs and AgMPs was examined by SEM analysis. A typical SEM micrograph of the AgNPs and AgMPs is shown in Figure 3. The uncapped AgMPs were found to be hexagonal (Figure 3A) with an average size of 500 nm ± 4. It was observed that maleic acid and citric acid controlled the shape and size of the synthesized silver particles slightly differently. When maleic acid was used as a modifier, the nanoparticles exhibited round plate-like morphology with sharp multiple facets with an average size of 39 nm ± 4 (Figure 3B). However, when citric acid was used as the modifier, the particles demonstrated a flower-like shape with rough surfaces with an average size of particles 250 nm ± 4 (Figure 3C). The ratio of the silver ions to stabilizing agents controls the size of particles produced. Hence, different factors, including the concentration of the reducing agents, metal salts, as well as time, may have played an important role in the formation of AgNPs [45]. Apart from that, stabilizing agents and modifiers also play a significant role in controlling the shape of the particles by preventing their agglomeration [23]. Modifiers in the current study include maleic acid and citric acid, which influenced the shape and size of the synthesized silver particles.

Maleic acid, being an unsaturated dibasic acid, contains one double bond, typically showing reactions of both carboxylic acids and olefins. Furthermore, there are three carboxyls and one hydroxyl in citric acid, showing typical reactions of carboxylic acids. Thus, they get absorbed on the surface of silver nanoparticles through double bonds and carboxylic acid groups. There are reports which state that silver possesses the lowest surface energy in the 111-plane, and when appropriate additives get adsorbed there, such as citrate and maleic acid [2,20], it might further decline the surface energy and could make synthesized silver plates more stable with the 111-plane as a basal plane. Both citrate and maleic acid favored the growth orientation of crystals of silver seeds. Hence, numerous plate-like silver petals accumulated and shaped secondary particles in agreement with previous reports [20,46]. Moreover, these face-centered cubic crystalline AgNPs with a (111) basal plane have previously been proven to be strong antimicrobial, cytotoxic, and antioxidant agents [36,47,48], and similar findings were observed in the current study.
Figure 3. Scanning electron microscopy (SEM) micrographs of silver nanoparticles and microparticles: (A) uncapped silver microparticles (AgMPs); (B) maleic acid capped silver nanoparticles (AgNPs); and (C) citric acid capped silver microparticles (AgMPs). The images are presenting the shape and size of the synthesized nanoparticles. Scale bar = 5 µm.

3.2. Antibacterial Activity

Currently, nanoparticles are regarded as a practicable substitute for antibiotics and appear to have great potential for resolving the issues related to the development of multidrug resistance against bacteria [49]. The AgNPs are currently considered a potential candidate for the fabrication of a novel course of antimicrobials providing a novel approach to fight against a wide range of pathogens [41,50–52]. In the current study, synthesized AgNPs and AgMPs showed significant antibacterial activity (Figure 4). All the synthesized particles were bactericidal against studied bacterial strains, including both Gram-positive and Gram-negative.

Additionally, minimum inhibitory concentration (MIC) values were determined. Results showed that AgNPs and AgMPs appeared more effective against Gram-negative bacteria (MIC as low as 5 ppm) as compared to Gram-positive (MIC 25 ppm) (Table 1). This can be attributed to less peptidoglycan in the case of the Gram-negative bacterial cell wall [36].

Figure 4. Antibacterial activity of synthesized silver nanoparticles and microparticles: Antibacterial activity of synthesized silver nanoparticles (AgNPs) and silver microparticles (AgMPs) is shown against (A) M. leutus; (B) S. setubal; (C) E. aerogenes; (D) A. tumefaciens. M.A: maleic acid capped AgNPs, C.A: citric acid capped AgMPs, C: Control or uncapped AgMPs. +ve control is streptomycin, while –ve control is distilled water.
Furthermore, our data show that the maleic acid capped AgNPs’ smaller size and round platelike, yet sharp facets were more bactericidal than the citric acid capped and uncapped AgMPs with relatively larger size and rough surfaces. Previous reports have demonstrated the size- and shape-dependent activity of the AgNPs, where smaller nanoparticles appear to have more ability to infiltrate into bacteria, and they have a greater tendency to release more silver ions than larger particles to kill more pathogens [53–55]. It has been shown that the bactericidal activity of AgNPs is dependent on size and the dose. Previously, the citrate capped small-sized (10 nm) AgNPs at the concentration of 20 ppm showed bactericidal effect against S. aureus (14%), E.coli (53%), and B. subtilis (84%) [56]. When the size of the particles was further reduced to 5 nm, the bactericidal activity of AgNPs was increased. Likewise, nanoparticles made up of smaller crystallite units exhibit more antimicrobial potential, probably due to greater surface energy with higher reactivity because of condensed crystal size. The comparative antimicrobial performance obtained in our study are well aligned with this hypothesis and also well reported in previous investigations [33]. However, in the current study, the size of the citrate capped AgMPs was comparatively greater. Therefore, they showed higher MIC values of 100 ppm for S. aureus and 50 ppm for B. subtilis. This size-dependent activity is related to the enhanced surface area to volume ratio [56].

However, the antimicrobial activity could also be attributed to organic acids themselves used in this study, i.e., maleic acid and citric acid. It has been previously reported that maleic acid and citric acid exhibit antimicrobial effects [57–60]. The antimicrobial potential of organic acid is mainly related to the reduction in microbial cell pH by ionization of undissociated acid molecules and the disruption of the transport of the substrate by changing the permeability of the cell membrane [57]. It has also been reported previously that such bactericidal activity is because of the physical interaction of AgNPs with the bacterial cell surfaces, particularly in the case of Gram-negative bacteria, where the accumulated nanoparticles were reported on bacterial surfaces [61].

However, studies have also reported that AgNPs can cause damage to the cellular membrane resulting in altered structure and thus rendering bacteria more penetrable [62,63]. This influence is highly predisposed by the nanoparticles’ morphology and concentration [62,64]. Interaction of AgNPs with Escherichia coli [63] enables the mechanism of action, proving that accumulated nanoparticles on cellular membrane disrupt the bilayer integrity by creating a gap. This makes the membrane more permeable, and as a result, bacterial death occurs [65]. Since enhanced permeability causes more AgNPs to enter the bacterial cell, it may cause more damage to the intracellular structures. Consequently, protein synthesis is terminated as ribosomes might have been denatured, and translation and transcription can be obstructed by targeting the bacterial cell genetic material [66–68].

Furthermore, it is also reported that smaller nanoparticles seem to have a superior ability to penetrate bacteria where they interact with the membrane and results in cell death. Moreover, positively charged silver ions interact with the negatively charged nucleic acid and proteins, causing structural changes and damage to the cell wall, cell membrane, and nucleic acid, as well [66,67]. This indicates that more caution will be needed in biomedical applications of silver nanoparticles.

### 3.3. Antifungal Activity

The synthesized AgNPs and AgMPs were also found effective against five fungal strains (Figure 5). In contrast, uncapped AgMPs did not show inhibitory activity against any fungal strain. At a concentration of 500 ppm, the highest activity was observed against Aspergillus flavus 50.56% and 40.11% (growth inhibition) by maleic acid capped and citric acid capped particles, respectively. Whereas the least activity was observed against Aspergillus niger at the same concentration (500 ppm), showing 25.72% and 18.2% inhibition with maleic acid capped and citric acid capped AgNPs and AgMPs, respectively. Compared to the inhibition of bacterial species, the fungal species showed comparatively less growth inhibition. This could be because of the rigidity of the fungal cell wall, which is composed of chitin than peptidoglycan containing cell wall of bacteria [36].

Similar to their antibacterial activity, the maleic acid capped silver nanoparticles were also found to be more fungicidal, mainly because of their smaller size range and round shape with multiple
sharp facets providing more surface area to interact with pathogens. This enables them to act more effectively as compared to citric acid capped silver microparticles, which were comparatively larger in size and rough flower-shaped. This is supported by previous studies [34].

Several reports support the findings that small spherical nanoparticles have better antimicrobial potential than other morphologies as these are provided with a greater surface area to volume ratio, thus proven more effective [36,42]. Large surface to volume ratio and high-atomic-density (111) aspects possibly boosted the pathogen killing proficiency [42]. It is also understood that more silver ions are released by smaller sized particles than larger ones to kill more pathogens [55]. Other reports also support these findings confirming that antimicrobial activity is reliant on the size and shape of silver particles, mainly because diverse morphologies offer different areas to intermingle with microbes and, as a consequence, end up with different competence [55,69]. The effect of capping agents in enhancing the antifungal activity cannot be ruled out as organic acids, including citric acid, have been reported to have antifungal activity [70].

Moreover, the interaction of AgNPs with microorganisms (bacteria, fungi, and viruses) releases silver ions (Ag⁺) that may damage the microorganism by different means; for example, they target the microbial cell wall (negatively charged) to disable cellular enzymes and interrupt membrane penetrability. Subsequently, cell disintegrates, and cell death takes place [71]. Another plausible justification could be that the silver ions also interact with enzymes and proteins’ thiol groups, which play a vital part in its antimicrobial act [67]. As nanoparticles’ resistance against pathogens has not been reported yet, thus they may have a significant benefit over traditional antimicrobial agents [72].
Table 1. Antibacterial activity of maleic acid capped nano-sized silver particles (AgNPs) and citric acid capped or uncapped micro-sized silver particles (AgMPs) against different bacterial strains.

| AgNPs Conc. (ppm) | Gram-positive Strains | Gram-negative Strains | Zone of Inhibition (cm) ± S.E |
|------------------|-----------------------|-----------------------|-------------------------------|
|                  | M. luteus | S. aureus | B. subtilis | A. tumefaciens | S. Setubal | E. aerogenes |
| 100              | M.A. | 1.25 ± 0.88 | 0.4 ± 1.21 | 0.63 ± 0.71 | 1.31 ± 0.74 | 0.7 ± 1.31 | 0.68 ± 1.33 | 0.66 ± 1.33 | 2.4 ± 0.5 |
|                  | C.A. | 0.9 ± 0.5 | 0.75 ± 0.23 | 0.89 ± 0.60 | 1.33 ± 1.1 | 1.3 ± 0.5 | 1.3 ± 1.3 | 0.34 ± 1.3 | 0.9 |
| 50               | M.A. | 0.98 ± 0.56 | 0.68 ± 1.15 | 0.64 ± 1.45 | 0.52 ± 1.55 | 0.85 ± 1.16 | 0.66 ± 1.24 | 1.4 ± 2.24 | - |
|                  | C.A. | 1.1 ± 1.21 | 0.66 ± 1.15 | 0.98 ± 1.55 | 0.89 ± 1.11 | - | 1.3 ± 1.12 | - | - |
| 25               | M.A. | 0.74 ± 0.42 | 0.46 ± 0.4 | 0.4 ± 0.78 | 0.43 ± 0.44 | 0.54 ± 0.76 | 0.45 ± 0.76 | 0.72 ± 1.76 | - |
|                  | C.A. | 1.7 ± 0.82 | 1.3 ± 0.78 | 1.2 ± 0.78 | 0.98 ± 0.78 | 1.2 ± 0.78 | - | 0.98 ± 0.98 | - |
| 5                | - | - | - | - | - | - | - | - | - |
| 2.5              | - | - | - | - | - | - | - | - | - |
| AgNO₃           | - | - | - | - | - | - | - | - | - |

Negative Control (Distilled H₂O) - - - - - - - - - -
Positive Control (Streptomycin) 2.1 2.42 2.4 3.4 4.72 4.2

Abbreviations: AgNPs—silver nanoparticles, AgMPs—silver microparticles, conc—concentration, S.E—standard error, M.A—maleic acid, C.A—citric acid, C—uncapped control, M. luteus—Micrococcus luteus, S. aureus—Staphylococcus aureus, B. subtilis—Bacillus subtilis, A. tumefaciens—Agrobacterium tumefaciens, S. Setubal—Salmonella Setubal, E. aerogenes—Enterobacter aerogenes.
Figure 5. Antifungal activity of capped and uncapped silver nanoparticles and microparticles. Percentage antifungal activity of maleic acid capped silver nanoparticles (AgNPs, blue) and citric acid capped silver microparticles (AgMPs, red) is shown against different fungal species, including Mucor species, Aspergillus niger, Aspergillus flavus, Aspergillus fumigatus, and Fusarium solani. Terbinafine (purple) was used as positive control, and distilled water was used as negative control. Data are presented as the mean of triplicate readings ± standard deviation of the mean. M.A: maleic acid capped AgNPs, C.A: citric acid capped AgMPs, C: control, i.e., uncapped AgMPs.

3.4. Cytotoxicity by Brine Shrimp Lethality and MTT Assay In Vitro

Brine shrimp (Artemia) have extensive use in testing marine ecotoxicity and are a widely used model organism for use in toxicological assays as an acceptable alternative to the toxicity testing of mammals in the laboratory. In the current study, the cytotoxic impact of synthesized silver particles was examined on brine shrimp nauplii at four different concentrations, including 100, 50, 25, 5 µM. The viability of the nauplii was significantly decreased to 22% in the case of maleic acid capped, 44.3% in case of citric acid capped, and 75.3% in case of uncapped silver particles at the concentration of 100 µM (Figure 6A). The cell viability was decreased in a concentration-dependent manner, indicating that a higher concentration of silver particles is more toxic to the shrimp.

Statistically, the effect of synthesized silver particles was found to be significant ($p < 0.001$, Table 2) with IC$_{50}$ 19.14 and 30.41 µM for maleic acid and citric acid capped silver microparticles, respectively. Whereas uncapped silver microparticles were found to be less toxic with a greater IC$_{50}$ value of 45.1 µM. Previously, silver oxide nanoparticles synthesized using the leaf extract of Callistemon lanceolatus also showed dosage and time-dependent cytotoxicity against brine shrimp nauplii with an IC$_{50}$ value of 85.32 ppm and LC$_{90}$ value 221.8 ppm [73].

Table 2. Analysis of variance for factors affecting the viability of brine shrimp.

| Source of Variation | Df | Sum-of-squares | Mean Square | F-Value | p-Value | Significant |
|---------------------|----|----------------|-------------|---------|---------|-------------|
| Interaction         | 9  | 3091           | 343.5       | 117.4   | <0.0001 | Yes         |
| Types of particles  | 3  | 14280          | 4761        | 1628    | <0.0001 | Yes         |
| Concentration       | 3  | 6649           | 2216        | 757.7   | <0.0001 | Yes         |
| Residual            | 32 | 93.60          | 2.925       |         |         |             |

As a review for cell viability testing, an MTT assay was performed. It was observed that the viability of cultured cells reduced with an increase in the concentration of particles. At the concentration of 10 µM maleic acid capped and citric acid capped particles, the viability of breast cancer (MCF-7) cells was 69% and 76.4%, respectively (Figure 6B). However, at the concentration of 50 µM, the viability of MCF-7 cells diminished to about 25.8% and 41.3% in the presence of maleic
acid and citric acid capped AgNPs and AgMPs, respectively, indicating a concentration-dependent effect. In contrast, the uncapped AgMPs at the same highest concentration, i.e., 50 µM, was found to be less toxic, which decreased viability of MCF7 cells to only 65.2%, unlike maleic acid and citric acid capped AgNPs and AgMPs, respectively, where the viability was 25.8% and 41.3%.

Figure 6. Cytotoxicity and antioxidant activity of silver nanoparticles and microparticles: (A) Percentage (%) cytotoxicity of silver particles against brine shrimp; (B) Antiproliferative activity against MCF7 cell line; (C) Antiproliferative activity against HePG2 cell line; (D) Free radical scavenging or antioxidant activity determined by 2,2-diphenyl-1-picrylhydrazyl (DPPH) assay.

Similarly, the proliferation of liver cancer HePG2 cells was also downregulated in a concentration-dependent manner (Figure 6C). However, the uncapped silver particles were not found effective against this cancer cell line. Shape and size played an important role in defining the cytotoxicity of the studied particles. The impact was found highly significant ($p < 0.001$, Table 3; Table 4) with an IC$_{50}$ value of 8.9 µM for maleic acid capped and 14.6 µM for citric acid capped and 46.54 µM for uncapped silver particles, respectively, for MCF7 cells. The IC$_{50}$ values for HePG2 cells were found to be 9.4 to 18.5 µM in a similar respect. The concentration of the silver particles was found to be a significant factor for inhibiting the proliferation of both MCF7 and HePG2 cells. The higher the concentration of particles, the fewer viable the cells were found, and vice versa [74]. However, the capping agent might have played an important role in enhancing the antiproliferative potential of silver nano- and microparticles as maleic acid and citric acid have been reported to have anticancer properties [75–77].

Table 3. Analysis of variance for factors affecting the viability of MCF7 Cells.

| Source of Variation | Df | Sum-of-squares | Mean Square | F-Value | p-Value | Significant |
|---------------------|----|----------------|-------------|---------|---------|-------------|
| Interaction         | 12 | 2227           | 185.6       | 97.86   | <0.0001 | Yes         |
| Types of particles  | 3  | 22690          | 7562        | 3988    | <0.0001 | Yes         |
| Concentration       | 4  | 5420           | 1355        | 714.6   | <0.0001 | Yes         |
| Residual            | 40 | 75.84          | 1.896       |         |         |             |
According to ISO standards, a reduction in cell viability by more than 30% is considered to indicate cytotoxicity [78]. It has been observed that silver nanoparticles induce toxicity because of the generation of reactive oxygen species (ROS) and oxidative stress, and it has also been reported that toxicity is dependent on the size of the nanoparticles [79–81]. Another report described that the development of the immune response by silver nanoparticles is size-dependent. In that study, treatment of various sized silver nanoparticles (4, 20, and 70 nm) was given to human macrophages (U-937), and it was observed that nanoparticles which were the smallest in size showed the highest pro-inflammatory activity (by releasing cytokines and inducing oxidative stress) [81,82].

Apoptosis induced by silver nanoparticles of different tumor cells has already been reported in vitro and ex vivo, which possibly occurred because of the translocation of the silver within the nucleus [83–86]. Franco-Molina et al. reported that MCF-7 breast cancer cells treated with silver colloids exhibited considerably decreased dehydrogenase activity, causing NADH/NAD+ reduction [87]. Consequently, reduced mitochondrial membrane potential leads to cell death. Another report elaborated that phagocytized silver nanoparticles totally block the cell cycle in the S-phase and excite inflammatory signaling through ROS generation, which finally induces the secretion of tumor necrosis factor-alpha (TNF-α) [48]. Silver nanoparticles synthesized by green chemistry have been shown to exhibit anticancer activity against MCF7 and HePG2 cell lines, whereas silver oxide nanoparticles were found cytotoxic to HePG2 and Chang liver cells [48]. In the current study, the inhibition of liver cancer (HePG2) and breast cancer cells (MCF7) was caused by maleic acid- and citric acid-capped silver particles.

Moreover, it has been reported that the efficiency of silver particles can be enhanced by coating their surfaces, which not only modify their surface charge but also influence the solubility. Various reports suggest that the toxicity and fate of nanoparticles are determined by their surface coatings [88,89]. Previously different coatings have been applied to silver nanoparticles, including trisodium citrate (CT-AgNPs) and polyvinylpyrrolidone (PVP-AgNPs), which improve the physicochemical properties of silver nanoparticles, such as biocompatibility and stability, as compared to uncoated particles [90]. The modifications in the surface charge of nanoparticles through coating agents may influence their toxicity in the recipient cells. For instance, nanoparticles which have a positive charge, such as Ag+, are considered more effective drug delivery tools for anticancer drugs as compared to negatively charged ones due to their longer retention time in the bloodstream [43].

Table 4. Analysis of variance for factors affecting the viability of HePG2 Cells.

| Source of Variation | Df | Sum-of-squares | Mean Square | F-Value | p-Value | Significant |
|---------------------|----|----------------|-------------|---------|---------|-------------|
| Interaction         | 12 | 1676           | 139.6       | 100.5   | <0.0001 | Yes         |
| Types of particles  | 3  | 10340          | 3447        | 2852    | <0.0001 | Yes         |
| Concentration       | 4  | 1571           | 392.7       | 374.5   | <0.0001 | yes         |
| Residual            | 40 | 205.3          | 5.133       |         |         |             |

3.5. Antioxidant Activity

The silver particles fabricated with two different capping agents showed DPPH free radical scavenging activity (Figure 6D). The antioxidant activity of the synthesized silver nanoparticles was found to be 74.4% (maleic acid-coated nanoparticles), 55.2% (citric acid-coated silver particles) at 80 µM. It was found to be significant with \( p < 0.001 \) (Table 5), having a decreasing trend in a concentration-dependent manner. The low IC50 value of 32.03 µM was observed for smaller particles, i.e., maleic acid capped silver nanoparticles (AgNPs) as compared to larger particles, i.e., citric acid capped ones (AgMPs), which showed an IC50 value of 49.62 µM. However, the uncapped silver particles did not show scavenging activity.

DPPH is a stable free radical and accepts an electron or hydrogen radical to become a stable diamagnetic molecule. The antioxidants’ impact on DPPH is mainly because of their hydrogen donating activity [91]. It has been generally utilized to evaluate the radical scavenging ability of synthesized nanoparticles [40]. Kumar et al. reported good antioxidant activity of green synthesized silver nanoparticles (>78%) at 0.1mM [40].
Another report described the antioxidant effect of silver nanoparticles against DPPH, showing 75.16 ± 0.04% inhibition [47]. The antioxidant activity of silver particles in our study might be because of the capping agents (maleic acid and citric acid) adsorbed at the nanoparticles’ surface, which have been reported to possess antioxidant activities due to their ability to chelate metals, and are, therefore, classified as “preventive” or “synergistic” [92]. At higher concentrations, the nanoparticles scavenged the free radicals more than at lower concentrations, possibly because more capping agents were adsorbed at the surface of nanoparticles in the case of a higher concentration, as proposed previously [74].

Taken together, our results showed that the modifiers act as stabilizers, and they may also be involved in enhancing the antimicrobial and anticancer potential of nanoparticles, and may, in part, contribute to the antioxidant activity of nanoparticles to scavenge or remove the free radicals (Figure 7). However, further experimentation may confirm their respective significance in the field of nanomedicine. The use of these modifiers could be a step forward to stabilize the silver nano- and microparticles and may enhance the therapeutic efficacy of silver particles capped with maleic acid and citric acid.

4. Conclusions

An effective and reproducible chemical reduction method for the synthesis of well-defined nano- and micro-sized silver particles has been presented, which implies that the addition of modifiers plays an important role in controlling the shape and size of the synthesized nanoparticles.
The current study showed that modifiers act as stabilizers of nano- and microparticles and affect their size and shape. These modifiers may, in part, contribute to enhancing the antimicrobial and anticancer potential of silver particles and could contribute to the antioxidant activity of nanoparticles to scavenge or remove the free radicals. However, maleic acid and citric acid need to be evaluated separately for their efficacy against the studied bacterial and fungal strains and their cytotoxicity against brine shrimp and studied cancer cell lines. This study may offer an opportunity to explore the therapeutic applications of nano- and micro-sized silver particles further. However, keeping in view the accumulation of silver particles in organs or tissues, more cautions will be needed in biomedical applications of silver particles, particularly, their repeated or long-term uses.

**Supplementary Materials:** The following are available online at www.mdpi.com/2079-4983/11/4/76/s1. Figure S1: Scanning electron microscope (SEM) image showing target (A) of energy dispersive X-Ray spectroscopy (EDS) spectra of uncapped silver microparticles (AgMPs) with (B). Figure S2: Scanning electron microscope (SEM) image showing target (A) of energy dispersive X-Ray spectroscopy (EDS) spectra of maleic acid capped silver nanoparticles (AgNPs) (B). Figure S3: Scanning electron microscope (SEM) image showing target (A) of energy dispersive X-Ray spectroscopy (EDS) spectra of citric acid capped silver microparticles (AgMPs) (B).

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**References**

1. Cölfen, H.; Mann, S. Higher-order organization by mesoscale self-assembly and transformation of hybrid nanostructures. *Angew. Chem. Int. Ed.* 2003, 42, 2350–2365.
2. Cai, X.; Zhai, A. Preparation of microsized silver crystals with different morphologies by a wet-chemical method. *Rare Met.* 2010, 29, 407–412.
3. Xia, Y.; Yang, P.; Sun, Y.; Wu, Y.; Mayers, B.; Gates, B.; Yan, H. One-dimensional nanostructures: Synthesis, characterization, and applications. *Adv. Mater.* 2003, 15, 353–389.
4. Jiang, H.; Moon, K.-S.; Li, Y.; Wong, C. Surface functionalized silver nanoparticles for ultrahigh conductive polymer composites. *Chem. Mater.* 2006, 18, 2969–2973.
5. Mafuné, F.; Kohno, J.-Y.; Takeda, Y.; Kondow, T.; Sawabe, H. Structure and stability of silver nanoparticles in aqueous solution produced by laser ablation. *J. Phys. Chem. B* 2000, 104, 8333–8337.
6. Jain, P.K.; Huang, X.; El-Sayed, I.H.; El-Sayed, M.A. Noble metals on the nanoscale: Optical and photothermal properties and some applications in imaging, sensing, biology, and medicine. *Acc. Chem. Res.* 2008, 41, 1578–1586.
7. Panáček, A.; Kvitek, L.; Pruczek, R.; Kolář, M.; Vedečová, R.; Pizírová, N.; Zbořil, R. Silver colloid nanoparticles: Synthesis, characterization, and their antibacterial activity. *J. Phys. Chem. B* 2006, 110, 16248–16253.
8. Duran, N.; Duran, M.; de Jesus, M.B.; Seabra, A.B.; Favaro, W.J.; Nakazato, G. Silver nanoparticles: A new view on mechanistic aspects on antimicrobial activity. *Nanomedicine* 2016, 12, 789–799, doi:10.1016/j.nano.2015.11.016.
9. Ruparelia, J.P.; Chatterjee, A.K.; Duttagupta, S.P.; Mukherji, S. Strain specificity in antimicrobial activity of silver and copper nanoparticles. *Acta Biomater.* 2008, 4, 707–716, doi:10.1016/j.actbio.2007.11.006.
10. Bocate, K.P.; Reis, G.F.; De Souza, P.C.; Junior, A.G.O.; Durán, N.; Nakazato, G.; Furlaneto, M.C.; De Almeida, R.S.; Panagio, L.A. Antifungal activity of silver nanoparticles and simvastatin against toxigenic species of Aspergillus. *Int. J. Food Microbiol.* 2019, 291, 79–86, doi:10.1016/j.ijfoodmicro.2018.11.012.
11. Mussin, J.E.; Roldan, M.V.; Rojas, F.; Sosa, M.L.A.; Pellegrí, N.; Giusiano, G. Antifungal activity of silver nanoparticles in combination with ketoconazole against Malassezia furfur. *AMB Express* 2019, 9, 131, doi:10.1186/s13568-019-0857-7.

12. Hosseinpour-Mashkani, S.M.; Ramezani, M. Silver and silver oxide nanoparticles: Synthesis and characterization by thermal decomposition. *Mater. Lett.* 2014, 130, 259–262.

13. De Matteis, V.; Cascione, M.; Toma, C.C.; Leporatti, S. Silver Nanoparticles: Synthetic Routes, In Vitro Toxicity and Theranostic Applications for Cancer Disease. *Nanomaterials (Basel)* 2018, 8, 319, doi:10.3390/nano8050319.

14. Chugh, H.; Sood, D.; Chandra, I.; Tomar, V.; Dhawan, G.; Chandra, R. Role of gold and silver nanoparticles in cancer nano-medicine. *Artif. Cells Nanomed. Biotechnol.* 2018, 46, 1210–1220, doi:10.1080/21694101.2018.1449118.

15. Wiemann, M.; Vennemann, A.; Blaske, F.; Sperling, M.; Karst, U. Silver Nanoparticles in the Lung: Toxic Effects and Focal Accumulation of Silver in Remote Organs. *Nanomaterials (Basel)* 2017, 7, 441, doi:10.3390/nano7120441.

16. Tang, J.; Xiong, L.; Wang, S.; Wang, J.; Liu, L.; Li, J.; Yuan, F.; Xi, T. Distribution, translocation and accumulation of silver nanoparticles in rats. *J. Nanosci. Nanotechnol.* 2009, 9, 4924–4932, doi:10.1166/jnn.2009.1269.

17. Lee, J.J.; Kim, Y.S.; Song, K.S.; Ryu, H.R.; Sung, J.H.; Park, J.D.; Park, H.M.; Song, N.W.; Shin, B.S.; Marshak, D.; et al. Biopersistence of silver nanoparticles in tissues from Sprague-Dawley rats. *Part. Fibre Toxicol.* 2013, 10, 36, doi:10.1186/1743-8977-10-36.

18. Shahverdi, A.R.; Fakhimi, A.; Shahverdi, H.R.; Minaian, S. Synthesis and effect of silver nanoparticles on the antibacterial activity of different antibiotics against Staphylococcus aureus and Escherichia coli. *Nanomedicine* 2007, 3, 168–171, doi:10.1016/j.nano.2007.02.001.

19. Dizaj, S.M.; Lotfpour, F.; Barzegar-Jalali, M.; Zarrintan, M.H.; Adibkia, K. Antimicrobial activity of the metals and metal oxide nanoparticles. *Mater. Sci. Eng. C* 2014, 44, 278–284.

20. Wiley, B.; Sun, Y.; Mayers, B.; Xia, Y. Shape-controlled synthesis of metal nanostructures: The case of silver. *Chem. A Eur. J.* 2005, 11, 454–463.

21. Sun, Y.; Yin, Y.; Mayers, B.T.; Herricks, T.; Xia, Y. Uniform silver nanowires synthesis by reducing AgNO3 with ethylene glycol in the presence of seeds and poly (vinyl pyrrolidone). *Chem. Mater.* 2002, 14, 4736–4745.

22. Quintero-Quiróz, C.; Botero, L.E.; Zárate-Triviño, D.; Acevedo-Yepes, N.; Escobar, J.S.; Pérez, V.Z.; Riano, I.J.C. Synthesis and characterization of a silver nanoparticle-containing polymer composite with antimicrobial abilities for application in prosthetic and orthotic devices. *Biomater. Res.* 2020, 24, 13, doi:10.1186/s40824-020-00191-6.

23. Eiden-Assmann, S.; Widoniak, J.; Maret, G. Synthesis and characterization of porous and nonporous monodisperse colloidal TiO2 particles. *Chem. Mater.* 2004, 16, 6–11.

24. Sondi, I; Goia, D.V.; Matijević, E. Preparation of highly concentrated stable dispersions of uniform silver nanoparticles. *J. Colloid Interface Sci.* 2003, 260, 75–81.

25. Flores, C.Y.; Miñán, A.G.; Grillo, C.A.; Salvarezza, R.C.; Vericat, C.; Schilardi, P.L. Citrate-capped silver nanoparticles showing good bactericidal effect against both planktonic and sessile bacteria and a low cytotoxicity to osteoblastic cells. *ACS Appl. Mater. Interfaces* 2013, 5, 3149–3159.

26. Lee, S.; Na, K. Oleic acid conjugated polymeric photosensitizer for metastatic cancer targeting in photodynamic therapy. *Biomater. Res.* 2020, 24, 1–8.

27. Helaly, F.; El-Sawy, S.; Hashem, A.; Khattab, A.; Mourad, R. Synthesis and characterization of nanosilver-silicone hydrogel composites for inhibition of bacteria growth. *Contact Lens Anterior Eye* 2017, 40, 59–66.

28. Kim, J.H.; Park, H.; Seo, S.W. In situ synthesis of silver nanoparticles on the surface of PDMS with high antibacterial activity and bio-safety toward an implantable medical device. *Nano Converg.* 2017, 4, 33.

29. Samoilova, N.; Kurskaya, E.; Krayukhina, M.; Askadsky, A.; Yamskov, I. Copolymers of maleic acid and their amphiphilic derivatives as stabilizers for silver nanoparticles. *J. Phys. Chem. B* 2009, 113, 3395–3403, doi:10.1021/jp806683m.

30. Jiang, X.C.; Chen, C.Y.; Chen, W.M.; Yu, A.B. Role of citric acid in the formation of silver nanoplates through a synergistic reduction approach. *Langmuir* 2010, 26, 4400–4408, doi:10.1021/la903470f.

31. Umadevi, M.; Bindhu, M.R.; Sathe, V. A Novel Synthesis of Malic Acid Capped Silver Nanoparticles using Solanum lycopersicum Fruit Extract. *J. Mater. Sci. Technol.* 2013, 29, 317–322, doi:10.1016/j.jmst.2013.02.002.

32. Bing, A.; Cai, X.-H.; Wu, F.-S.; Wu, Y.-P. Preparation of micro-sized and uniform spherical Ag powders by novel wet-chemical method. *Trans. Nonferrous Met. Soc. China* 2010, 20, 1530–1534.

33. Maqbool, Q.; Nazar, M.; Naz, S.; Hussain, T.; Jabeen, N.; Kausar, R.; Anwaar, S.; Abbas, F.; Jan, T. Antimicrobial potential of green synthesized CeO2 nanoparticles from Olea europaea leaf extract. *Int. J. Nanomed.* 2016, 11, 5015.

34. Van Dong, P.; Ha, C.H.; Kasbohm, J. Chemical synthesis and antibacterial activity of novel-shaped silver nanoparticles. *Int. Nano Lett.* 2012, 2, 9.
35. Shabbir, M.; Ahmad, I.; Ismail, H.; Ahmed, S.; McKee, V.; Akhter, Z.; Poczai, P. Pharmacological, electrochemical and drug–DNA interaction aspects of tridentate Schiff bases and their triphenylphosphine nickel (II) complexes. *Polhedron* 2017, 133, 270–278.

36. Kumar, C.M.K.; Yugandhar, P.; Savithramma, N. Biological synthesis of silver nanoparticles from Adansonia digitata L. fruit pulp extract, characterization, and its antimicrobial properties. *J. Intercult. Ethnopharmacol.* 2016, 5, 79.

37. Ohkhena, F.U.; Wintola, O.A.; Afolayan, A.J. Evaluation of the antibacterial and antifungal properties of Phragmanthera capitata (Sprengel) Balle (Loranthaceae), a mistletoe growing on rubber tree, using the dilution techniques. *Sci. World J.* 2017, 1–8.

38. Ahmad, M.S.; Hussain, M.; Hanif, M.; Ali, S.; Qayyum, M.; Mirza, B. Di-and Triorgonat (IV) Esters of 3, 4-Methylenedioxyphenylpropenoic Acid: Synthesis, Spectroscopic Characterization and Biological Screening for Antimicrobial, Cytotoxic and Antitumor Activities. *Chem. Biol. Drug Des.* 2008, 71, 568–576.

39. Dilshad, E.; Zafar, S.; Ismail, H.; Waheed, M.T.; Cusido, R.M.; Palazon, J.; Poczai, P. Effect of rol genes on polyphenols biosynthesis in Artemisia annua and their effect on antioxidant and cytotoxic potential of the plant. *Appl. Biochem. Biotechnol.* 2016, 179, 1456–1468.

40. Kumar, B.; Smita, K.; Cumbal, L.; Debut, A. Green synthesis of silver nanoparticles using Andean blackberry fruit extract. *Saud J. Biol. Sci.* 2017, 24, 45–50.

41. Manikandan, V.; Velmurugan, P.; Park, J.-H.; Chang, W.-S.; Park, Y.-J.; Jayanthi, P.; Cho, M.; Oh, B.-T. Green synthesis of silver oxide nanoparticles and its antibacterial activity against dental pathogens. *3 Biotech.* 2017, 7, 72.

42. Raza, M.A.; Kanwal, Z.; Rauf, A.; Sabri, A.N.; Riaz, S.; Naseem, S. Size-and shape-dependent antibacterial studies of silver nanoparticles synthesized by wet chemical routes. *Nanomaterials* 2016, 6, 74.

43. Zhang, X.-F.; Liu, Z.-G.; Shen, W.; Gurunathan, S. Silver nanoparticles: Synthesis, characterization, properties, applications, and therapeutic approaches. *Int. J. Mol. Sci.* 2016, 17, 1534.

44. Martínez-Castañón, G.; Niño-Martínez, N.; Loyola-Rodríguez, J.; Patiño-Marin, N.; Martínez-Mendoza, J.; Ruiz, F. Synthesis of silver particles with different sizes and morphologies. *Mater. Lett.* 2009, 63, 1266–1268.

45. Khatoon, N.; Ahmad, R.; Sardar, M. Robust and fluorescent silver nanoparticles using Artemisia annua: Biosynthesis, characterization and antibacterial activity. *Biochem. Eng. J.* 2015, 102, 91–97.

46. Sun, Y.; Gates, B.; Mayers, B.; Xia, Y. Crystalline silver nanowires by soft solution processing. *Nano Lett.* 2002, 2, 165–168.

47. Nagaich, U.; Gulati, N.; Chauhan, S. Antioxidant and antibacterial potential of silver nanoparticles: Biogenic synthesis utilizing apple extract. *J. Pharm.* 2016, 2016, 1–8.

48. Karunagaran, V.; Rajendran, K.; Sen, S. Optimization of biosynthesis of silver oxide nanoparticles and its anticancer activity. *Int. J. Nanosci.* 2017, 16, 1750018.

49. Rai, M.; Deshmukh, S.; Ingle, A.; Gade, A. Silver nanoparticles: The powerful nanoweapon against multidrug-resistant bacteria. *J. Appl. Microbiol.* 2012, 112, 841–852.

50. Dos Santos, C.A.; Rai, M.; Ingle, A.P.; Gupta, I.; Galdiero, S.; Galdiero, M.; Gade, A.; Rai, M. Silver nanoparticles: Therapeutic uses, toxicity, and safety issues. *J. Pharm. Sci.* 2014, 103, 1931–1944.

51. Rai, M.; Deshmukh, S.D.; Ingle, A.P.; Gupta, I.R.; Galdiero, M.; Galdiero, S. Metal nanoparticles: The protective nanoshield against virus infection. *Crit. Rev. Microbiol.* 2016, 42, 46–56.

52. Lara, H.H.; Garza-Treviño, E.N.; Ixten-Turrent, L.; Singh, D.K. Silver nanoparticles are broad-spectrum bactericidal and virucidal compounds. *J. Nanobiotechnol.* 2011, 9, 30.

53. Tamayo, L.; Zapata, P.; Vejar, N.; Azocar, M.; Gulppi, M.; Zhou, X.; Thompson, G.; Rabagliati, F.; Páez, M.A. Release of silver and copper nanoparticles from polyethylene nanocomposites and their penetration into Listeria monocytogenes. *Mater. Sci. Eng. C* 2014, 40, 24–31.

54. Wu, D.; Fan, W.; Kishen, A.; Gutmann, J.L.; Fan, B. Evaluation of the antibacterial efficacy of silver nanoparticles against Enterococcus faecalis biofilm. *J. Endod.* 2014, 40, 285–290.

55. Pal, S.; Tak, Y.K.; Song, J.M. Does the antibacterial activity of silver nanoparticles depend on the shape of the nanoparticle? A study of the gram-negative bacterium Escherichia coli. *Appl. Environ. Microbiol.* 2007, 73, 1712–1720.

56. Agnihotri, S.; Mukherji, S.; Mukherji, S. Size-controlled silver nanoparticles synthesized over the range 5–100 nm using the same protocol and their antibacterial efficacy. *RSC Adv.* 2014, 4, 3974–3983.

57. Organization, W.H. *Surface Decontamination of Fruits and Vegetables Eaten Raw: A Review*; World Health Organization: Geneva, Switzerland, 1998.
58. Ballal, N.V.; Yegneswaran, P.P.; Mala, K.; Bhat, K.S. In vitro antimicrobial activity of maleic acid and ethylenediaminetetaacetic acid on endodontic pathogens. *Oral Surg. Oral Med. Oral Pathol. Oral Radiol. Endodontology* 2011, 112, 696–700.

59. Esvaranandam, S.; Hettiarachchy, N.; Johnson, M. Antimicrobial activity of citric, lactic, malic, or tartaric acids and nisin-incorporated soy protein film against Listeria monocytogenes, Escherichia coli O157: H7, and Salmonella gaminella. *J. Food Sci.* 2004, 69, FMS79–FMS84.

60. In, Y.W.; Kim, J.J.; Kim, H.J.; Oh, S.W. Antimicrobial activities of acetic acid, citric acid and lactic acid against S. nigella species. *J. Food Saf.* 2013, 33, 79–85.

61. Franci, G.; Falanga, A.; Galdiero, M.; Palomba, L.; Rai, M.; Morelli, G.; Galdiero, M. Silver nanoparticles as potential antibacterial agents. *Molecules* 2015, 20, 8856–8874.

62. Periasamy, S.; Joo, H.-S.; Duong, A.C.; Bach, T.-H.L.; Tan, V.Y.; Chatterjee, S.S.; Ching, G.Y.C.; Otto, M. How Staphylococcus aureus biofilms develop their characteristic structure. *Proc. Natl. Acad. Sci. USA* 2012, 109, 1281–1286.

63. Lazar, V. Quorum sensing in biofilms–how to destroy the bacterial citadels or their cohesion/power? *Anaerobe* 2011, 17, 280–285.

64. Rolim, J.P.; Melo, M.A.S.; Guedes, S.F.; Albuquerque-Filho, F.B.; De Souza, J.R.; Nogueira, N.A.P.; Zanin, I.C.; Rodrigues, L.K.A. The antimicrobial activity of photodynamic therapy against Streptococcus mutans using different photosensitizers. *J. Photochem. Photobiol. B Biol.* 2012, 106, 40–46.

65. Rai, M.; Kon, K.; Ingle, A.; Duran, N.; Galdiero, S.; Galdiero, M. Broad-spectrum bioactivities of silver nanoparticles: The emerging trends and future prospects. *Appl. Microbiol. Biotechnol.* 2014, 98, 1951–1961.

66. Morones, J.R.; Elechiguerra, J.L.; Camacho, A.; Holt, K.; Kouri, J.B.; Ramirez, J.T.; Yacaman, M.J. The bactericidal effect of silver nanoparticles. *Nanotechnology* 2005, 16, 2346.

67. Jung, W.K.; Koo, H.C.; Kim, K.W.; Shin, S.; Kim, S.H.; Park, Y.H. Antibacterial activity and mechanism of action of the silver ion in Staphylococcus aureus and Escherichia coli. *Appl. Environ. Microbiol.* 2008, 74, 2171–2178.

68. Bury, N.R.; Wood, C.M. Mechanism of branchial apical silver uptake by rainbow trout is via the proton-coupled Na+ channel. *Am. J. Physiol. Regul. Integr. Comp. Physiol.* 1999, 277, R1385–R1391.

69. Martinez-Castanon, G.; Nino-Martinez, N.; Martinez-Gutierrez, F.; Martinez-Mendoza, J.; Ruiz, F. Synthesis and antibacterial activity of silver nanoparticles with different sizes. *J. Nanopart. Res.* 2008, 10, 1343–1348.

70. Hassan, R.; El-Kadi, S.; Sand, M. Effect of some organic acids on some fungal growth and their toxins production. *Int. J. Adv. Biol.* 2015, 2, 1–11.

71. Feng, Q.L.; Wu, J.; Chen, G.; Cui, F.; Kim, T.; Kim, J. A mechanistic study of the antibacterial effect of silver ions on Escherichia coli and Staphylococcus aureus. *J. Biomed. Mater. Res. B* 2005, 72, 662–668.

72. Ahmad, N.; Sharma, S.; Singh, V.; Shamsi, S.; Fatma, A.; Mehta, B. Biosynthesis of silver nanoparticles from Desmodium triflorum: A novel approach towards weed utilization. *Biotecnol. Res. Int.* 2011, 2011, 1–8.

73. Ravichandran, S.; Paluri, V.; Kumar, G.; Loganathan, K.; Kokati Venkata, B.R. A novel approach for the biosynthesis of silver oxide nanoparticles using aqueous leaf extract of Callistemon lanceolatus (Myrtaceae) and their therapeutical potential. *J. Exp. Nanosci.* 2016, 11, 445–458.

74. Gomaa, E.Z. Antimicrobial, antioxidant and antitumor activities of silver nanoparticles synthesized by Allium cepa extract: A green approach. *J. Genet. Eng. Biotechnol.* 2017, 15, 49–57.

75. Chiriac, A.; Brzezinski, P. Topical malic acid in combination with citric acid: An option to treat recalcitrant warts. *Dermatol. Ther.* 2015, 28, 336–338.

76. Ying, T.-H.; Chen, C.-W.; Hsiao, Y.-P.; Hung, S.-J.; Chung, J.-G.; Yang, J.-H. Citric acid induces cell-cycle arrest and apoptosis of human immortalized keratinocyte cell line (HaCaT) via caspase-and mitochondria-dependent signaling pathways. *Anticancer Res.* 2013, 33, 4411–4420.

77. Ren, J.-G.; Seth, P.; Ye, H.; Jian-Guo, R.; Hanai, J.-I.; Husain, Z.; Sukhatme, V.P. Citrate suppresses tumor growth in multiple models through inhibition of glycolysis, the tricarboxylic acid cycle and the IGF-1R pathway. *Sci. Rep.* 2017, 7, 1–13.

78. Kurt, A.; Erkose-Genc, G.; Uzun, M.; Emrence, Z.; Ustek, D.; Isik-Ozol, G. The antifungal activity and cytotoxicity of silver containing denture base material. *Niger. J. Clin. Pract.* 2017, 20, 290–295.

79. Carlson, C.; Hussain, S.M.; Schrand, A.M.; Braydich-Stolle, L.K.; Hess, K.L.; Jones, R.L.; Schlager, J.J. Unique cellular interaction of silver nanoparticles: Size-dependent generation of reactive oxygen species. *J. Phys. Chem. B* 2008, 112, 13608–13619.

80. Kim, S.; Choi, J.E.; Choi, J.; Chung, K.-H.; Park, K.; Yi, J.; Ryu, D.-Y. Oxidative stress-dependent toxicity of silver nanoparticles in human hepatoma cells. *Toxicol. Vitr.* 2009, 23, 1076–1084.
81. Park, M.V.D.Z.; Neigh, A.M.; Vermeulen, J.P.; De La Fonteyne-Blankestijn, L.J.; Verharen, H.W.; Briedé, J.J.; Van Loveren, H.; De Jong, W.H. The effect of particle size on the cytotoxicity, inflammation, developmental toxicity and genotoxicity of silver nanoparticles. *Biomaterials* **2011**, *32*, 9810–9817.

82. Park, J.; Lim, D.-H.; Lim, H.-J.; Kwon, T.; Choi, J.-S.; Jeong, S.; Choi, I.-H.; Cheon, J. Size dependent macrophage responses and toxicological effects of Ag nanoparticles. *Chem. Commun.* **2011**, *47*, 4382–4384.

83. Banerjee, K.; Das, S.; Choudhury, P.; Ghosh, S.; Baral, R.; Choudhuri, S.K. A novel approach of synthesizing and evaluating the anticancer potential of silver oxide nanoparticles in vitro. *Chemotherapy* **2011**, *57*, 279–289.

84. Park, E.-J.; Yi, J.; Kim, Y.; Choi, K.; Park, K. Silver nanoparticles induce cytotoxicity by a Trojan-horse type mechanism. *Toxicol. Vit.*** **2010**, *24*, 872–878.

85. Yen, H.J.; Hsu, S.H.; Tsai, C.L. Cytotoxicity and immunological response of gold and silver nanoparticles of different sizes. *Small*** **2009**, *5*, 1553–1561.

86. Netchareonsirisuk, P.; Puthong, S.; Dubas, P.; Palaga, T.; Komolpis, K. Effect of capping agents on the cytotoxicity of silver nanoparticles in human normal and cancer skin cell lines. *J. Nanopart. Res.* **2016**, *18*, 322.

87. A Franco-Molina, M.; Mendoza-Gamboa, E.; A Sierra-Rivera, C.; A Gómez-Flores, R.; Zapata-Navarrete, P.; Castillo-Tello, P.; Alcocer-Gonzalez, J.M.; Miranda-Hernandez, D.F.; Tamez-Guerra, R.S.; Rodriguez-Padilla, C. Antitumor activity of colloidal silver on MCF-7 human breast cancer cells. *J. Exp. Clin. Cancer Res.* **2010**, *29*, 1.

88. Sharma, V.K.; Siskova, K.M.; Zboril, R.; Gardea-Torresdey, J.L. Organic-coated silver nanoparticles in biological and environmental conditions: Fate, stability and toxicity. *Adv. Colloid Interface Sci.* **2014**, *204*, 15–34.

89. Nguyen, K.C.; Seligy, V.L.; Massarsky, A.; Moon, T.W.; Rippstein, P.; Tan, J.; Tayabali, A.F. Comparison of toxicity of uncoated and coated silver nanoparticles. In *Proceedings of Journal of Physics: Conference Series*; IOP Publishing: Redcliffe, UK; 2013; p. 012025.

90. Jurasić, D.D.; Ćurlin, M.; Capjak, I.; Crnković, T.; Lovrić, M.; Babić, M.; Horák, D.; Vrček, I.V.; Gajović, S. Surface coating affects behavior of metallic nanoparticles in a biological environment. *Beilstein J. Nanotechnol.* **2016**, *7*, 246–262.

91. Johnson, A.; Obot, I.; Ukpong, U. Green synthesis of silver nanoparticles using Artemisia annua and Sida acuta leaves extract and their antimicrobial, antioxidant and corrosion inhibition potentials. *J. Mater. Environ. Sci.* **2014**, *5*, 899–906.

92. Kazemi, M.; Hadavi, E.; Hekmati, J. Effect of salicylic acid, malic acid, citric acid and sucrose on antioxidant activity, membrane stability and ACC-oxidase activity in relation to vase life of carnation cut flowers. *J. Agric. Technol.* **2012**, *8*, 2053–2063.

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