The antimicrobial activity of biogenic silver nanoparticles synthesized from extracts of Red and Green European pear cultivars

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\section*{ABSTRACT}

Green nanotechnology stands amongst the leading giants of innovation for the twenty first century technological advances. More interesting, is the use of natural products as reducing agents. These could be recyclable materials from fruits and vegetables to produce nanoparticles (NPs) with novel properties. In the current study, silver NPs (AgNPs) were synthesized using the water extracts from the peel and flesh of two \textit{Pyrus communis} L. cultivars, namely, the Forelle (Red) Pears (RPE) and Packham Triumph (Green) Pears (GPE). The AgNPs were characterized by UV-Vis spectrophotometry, Dynamic Light Scattering (DLS), High Resolution Transmission Electron Microscopy (HRTEM) and Fourier Transform Infra-Red Spectroscopy (FTIR). The antibacterial activities of the AgNPs were evaluated using agar well diffusion and microdilution assays. The cytotoxicity of the AgNPs was investigated on a rat macrophage (RAW 264.7) cells using MTT assay. Both the RPE and GPE were capable of synthesizing the AgNPs at high temperatures (70 and 100 °C). The AgNPs exhibited antibacterial activity against the test strains, and also had low toxicity towards the RAW 264.7 cells. Thus, the synthesized AgNPs have a potentially viable use in bio-applications for treatment of bacterial infections.

\section*{Introduction}

Green nanotechnology has brought a lot of exciting opportunities that could be beneficial to waste recycling in the agricultural sector \cite{1,2}. A large volume of fruit and vegetable wastes are discarded daily. Since these waste products contain useful phytochemicals, they could instead be recycled to harness the bio-active compounds and incorporate them into new products \cite{2,3}. The use of fruit, vegetable or plant extracts in the biological synthesis of nanoparticles (NPs) is an attractive approach as plant materials are readily available \cite{4}. Plant extracts reported in the synthesis of NPs contain antioxidants such as phenolic acids and flavonoids \cite{4,5}, with valuable functional groups such as hydroxyl, carboxyl, carboxyl and phenol groups. These functional groups are involved in the reduction of various metals (silver, gold and palladium) into their respective metallic NPs \cite{6}.

Metallic NPs, such as silver nanoparticles (AgNPs) are of particular interest. These NPs exhibit a number of properties that are useful in food sciences \cite{1,2}, biosensors \cite{2,7} nanobiotechnology, nanomedicine, textile, etc. The NPs at a size range of 1–100 nm \cite{8,9} have been shown to have unique and novel properties \cite{8}. Due to their unique physical and chemical properties, AgNPs are widely used in nanomedicine as antimicrobial \cite{3,10}, as antiseptic agents in the clothing industry, as well as in the development of skincare products \cite{11}.

The current study explores the synthesis of AgNPs using the European pear or \textit{Pyrus communis} L., This pear fruit is common in temperate zones worldwide, and has over 2000 variants \cite{12,13}. The Packham’s Triumph (Green) and Forelle (Red) are among pear variants available in South Africa \cite{14}. The red-skinned appearance of Forelle pears is due to the high amounts of anthocyanins \cite{15}. Anthocyanins are flavonoids that are responsible for the various colours in flowers, vegetables and fruits. They have been shown to have a number of properties, which include antibacterial, antioxidant, and anticancer bioactivities \cite{16}. Anthocyanin-rich fruits, such as blackberry (\textit{Rubus fruticosus} L.), strawberry (\textit{Fragaria vesca} L.) and raspberry (\textit{Rubus idaeus} L.), have been used to synthesize AgNPs with an antimicrobial activity using green nanotechnology \cite{17}.

The pear industry is the third-largest in the South African fruit industry after citrus and apples \cite{18}. The \textit{Pyrus} spp. is extremely nutritious and a rich source of phytochemicals associated with health benefits (nutraceuticals). These phytochemicals are excellent candidates for the synthesis of bioactive AgNPs \cite{19,20}. While the synthesis of both gold NPs...
[21] and AgNPs [19,20,22] have been demonstrated using Asian pear variants, such as the Huangguan pear (*Pyrus bretschneideri* Rehder); the European pears have not been used previously to synthesize AgNPs. Both the appearance and phytochemical composition of the Asian and European pears differ significantly. Asian pear variants contain more glucose compared to European pears, which have higher fructose to glucose ratio and a higher concentration of sorbitol [23]. It is, therefore, possible that the characteristics of the AgNPs produced from the green and red European pear variants will be different. In this study, we demonstrate the synthesis of AgNPs from aqueous extracts of Packham’s Triumph and Forelle pears and show their antibacterial activity against several bacterial strains.

Materials and methods

**Preparation of the pear water extract**

The Green (Packham Triumph) and the Red (Forelle) pears were a kind gift from Kromco (Pty) Ltd (Grabouw, South Africa). Five pears each from green and red pears were washed with distilled water (dH2O) and dried with a paper towel. The peel and flesh were separated and individually blended in 400 mL boiled dH2O. The crushed material was centrifuged at 9000 rpm for 30 min using the Sorval Lynx 6000 centrifuge (Thermo Fisher Scientific, USA). The supernatant was vacuum filtered using Whatman 0.45 µm filters (Merck, Germiston, South Africa). The filtrates were frozen at −80 °C and freeze-dried using a freeze dryer (Virtis, Gardiner, NY, USA). The extracts were dissolved in dH2O to obtain Green and Red Pear Peel (GPP and RPP, respectively) and Pear Flesh (GPF and RPF, respectively) extracts.

**Phytochemical analysis of the pear-extracts**

The Total Phenolic Content (TPC) analysis was evaluated in the RPE and GPE following the Folin-Ciocalteau assay using a previously described protocol [24]. The TPC was expressed as µg/mL equivalents of gallic acid.

The reducing sugars in the RPE and GPE were analysed following the phenol-sulfuric acid reaction method using 0–25 mg/mL D-glucose (Sigma) as a standard, following a previously described protocol [25].

**Synthesis of AgNPs**

Synthesis of AgNPs was performed by reducing AgNO3 (Sigma, St. Louis, MO, USA) with the GPE and RPE. The AgNPs were synthesised by mixing the pear extracts and AgNO3 solution in a 1:3 (v/v) ratio (i.e. 0.5 mL extract: 1.5 mL AgNO3). Synthesis was performed using different concentrations of the extracts (1.563–50 mg/mL), AgNO3 solution (0.5, 1, 2 and 3 mM), temperatures (25, 70 and 100 °C), and pH ranges (4, 5, 6, 7, 8, 9 and 10). All the reactions were incubated for 16 h at their set conditions, with shaking. After the incubation period, all the samples were inspected for a colour change and imaged using a digital camera. The reactions were performed in 2 mL tubes in an Eppendorf Thermomixer Comfort (Hamburg, Germany) while shaking at 400 rpm.

**Characterization of the AgNPs**

The AgNPs were centrifuged using an Eppendorf centrifuge 5417 R (Hamburg, Germany) at 14,000 rpm for 10 min to remove unreacted extract and AgNO3 before analysis. The AgNPs were characterized by UV-Vis using a POLARstar Omega Plate reader (BMG Labtech, Offenburg, Germany) at a wavelength ranging from 300 to 700 nm, Dynamic Light Scattering (DLS) using a Malvern NanoZS90 Zetasizer (Malvern Panalytical Ltd., Enigma Business Park, UK), FTIR spectra read on an FTIR spectrophotometer (Perkin Elmer, Waltham, MA, USA) in a 500–4000 cm−1 wavenumber range, and HRTEM using Tecnai F20 Field Transmission Electron Microscope (FEI Company, Oregon, USA), as previously described [26].

**Bioactivity of the AgNPs**

**Antibacterial activity of the AgNPs**

*Staphylococcus aureus*, MRSA, *Pseudomonas aeruginosa*, and *Escherichia coli* were used to assess the antibacterial activity of the pear extracts and AgNPs. The bacterial strains were purchased from American Tissue Culture Collection (ATCC; Manassas, VA, USA). The bacterial cultures were adjusted to 0.5 McFarland standard at OD600nm between 0.08 and 0.1. The cultures were then diluted to 1:150 with fresh Müller Hinton Broth (MHB, Sigma), and used for agar well diffusion and microdilution assays, as previously described [5,26].

**Agar well diffusion assay**

Microorganisms were streaked on the Müller Hinton Agar (MHA, Sigma) plates. Then 50 µL each of the negative control (MHB), positive control (1:1 penicillin-streptomycin (Pen-Strep) cocktail at 10 U/mL) (Gibco, Bleiswijk, The Netherlands), pear extracts and the AgNPs were added in triplicates into their respective wells.

**Determination of minimum inhibitory concentration (MIC)**

The microorganisms were exposed to the pear extracts and AgNPs at concentrations of 0 to 500 µg/mL, for 24 h. The MIC was recorded for each treatment, and the OD600nm was measured for each treatment using the POLARstar Omega Plate reader.

**Evaluation of the cytotoxicity of the AgNPs**

Cytotoxicity of the pear extracts and AgNPs was evaluated in RAW 264.7 cells (ATCC) using the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT, Sigma) assay, as previously described [27]. The cells were grown in Dulbecco’s Modified Eagles Medium (Gibco, Roche, Germany) supplemented with 10% foetal bovine serum (Gibco) and 1% Pen-Strep. The cells were treated with 0–500 µg/mL pear
extracts and AgNPs. Doxorubicin (Sigma) at a concentration of 50 µg/mL was used as a positive control.

Statistical analysis
The data are expressed as mean ± standard error of the mean of three independent experiments, with each condition performed in triplicate. Statistical analysis was performed by two-way ANOVA using Graphpad™ Prism v6 software, and t-test using Microsoft Excel (2010). The differences between the samples were considered statistically significant when the p < .05.

Results and discussion
Green synthesis of AgNPs
Green nanotechnology provides safer, inexpensive, and eco-friendly methods for the synthesis of NPs using natural products as reducing, capping and stabilizing agents [5, 26]. These green chemistry principles-based methods replace the toxic reducing agents that are used in chemical synthesis methods. The natural products derived from either microorganisms or plants can be biomolecules and/or secondary metabolites that have the ability to reduce metal precursors into metallic NPs [28, 29]. In recent years, green nanotechnology has shifted interest to the use of food wastes, including, fruits and vegetables for the synthesis of NPs. These waste products were shown to produce bio-active NPs [3, 30–32] that are biocompatible and have selective toxicity [31].

Pears have high quantities of nutraceuticals, such as phenolic acid, flavonoids, and triterpenes. The phenolic acids include gallic acid, ferulic acid, ursolic acid, etc. [13, 33]. Similar phenolic acids have been implicated in the green synthesis of NPs [31, 34, 35]. The nutraceuticals certainly vary among the pear variants [13, 33]. As shown in Table 1, the RPE (both the peel and flesh) had a higher reducing sugar content and lower TPC compared to the GPE. The phenolic compounds are the major phytochemicals in plant extracts and prospective reducing agents in the green synthesis of metallic NPs [36–38], while the sugars can be used as reducing and/or stabilizing agents [39]. Their presence in the pear extracts substantiates their role in the reduction of AgNO3 and stabilization of the AgNPs.

Analysis of TPC and reducing sugars in the pear-extracts.

| Pear extracts | TPC (µg/mL) | Reducing sugars (µg/mL) |
|---------------|------------|------------------------|
| RPP           | 16         | 75                     |
| RPF           | 38         | 45                     |
| GPP           | 127        | 51                     |
| GPF           | 65         | 36                     |

Table 1.

Effect of extract concentration and temperature on AgNP synthesis
To determine the optimal extract concentration and temperature for the synthesis of AgNPs, various concentrations of RPE and GPE were incubated with 1 mM AgNO3. Changes in colour from clear to yellow and then brown indicated successful synthesis of the AgNPs, and were further confirmed by UV-Vis analysis (Figure 1). As shown in Figure 1(A), there was no colour change in the samples that were incubated at room temperature (∼25 °C) for all concentrations. This was further confirmed by UV-Vis spectra of these samples, indicating the absence of the characteristic peak around 400–500 nm, thus further indicating that there was no formation of RP-AgNPs at 25 °C. These variable effects of temperature have been reported previously and attributed to the different types of phytochemicals in a particular plant extract. While the *Acinetobacter calcoaceticus* extracts were unable to synthesise AgNPs at 20 °C [40], the extracts from *Salvia africana-lutea*, *Sutherlandia frutescens* [26] and *Terminalia mantaly* [5] produced AgNPs at 25 °C.

RPE-synthesized AgNPs were visible from 6.25 to 50 mg/mL at 70 °C (Figure 1(B)), and in all tested concentrations for reactions at 100 °C (Figure 1(C)). The corresponding UV-Vis spectra showed surface plasmon resonance (SPR) peaks around 400–450 nm. SPR is defined as a phenomenon where the electrons of the metal involved in the NP synthesis (Ag in this case) become excited by the photons at a certain angle of incidence [41]. AgNPs display a characteristic SPR peak around 400–450 nm [5, 26, 42]; which is also used to profile NP size, shape, stability, and estimate NP concentration [43].

Similar effects of temperature and extract concentrations were observed for GP-AgNPs, as shown in Figure 2. At 25 °C, there was no colour change or UV-Vis spectra in all the GPE-AgNO3 reactions. Colour changes to light yellow, orange and brown were observed at 70 °C and 100 °C. This was demonstrated by the absence or presence of the SPR peaks in the UV-Vis spectra around 400–450 nm. In essence, the colour change and presence of SPR suggested the formation of AgNPs [5, 26, 44].

The intensity of the colour, as well as the absorbance, are directly proportional to the production rate and the concentration of the NPs [26, 43, 45]. These two parameters were used in the selection of the optimum pear extract concentration and temperature. The 100 °C was the optimal temperature for the synthesis of the RP-AgNPs and GP-AgNPs. At this temperature, all the RPE and GPE concentrations produced AgNPs with more intense colours than at 70 °C. Moreover, the AgNPs at 70 °C produced broader UV-Vis peaks compared to 100 °C, signifying the production of polydispersed AgNPs at 70 °C. Based on these parameters, the optimum concentrations of the pear extracts were 3.125 mg/mL for the peel extract and 6.25 mg/mL for the flesh extract.

Effect of pH on AgNP synthesis
A yellow to brown colors were observed at all pH conditions, confirming the synthesis of the AgNPs. As shown in Figure 3,
the peel extracts produced narrow UV-Vis spectra (A and C), while the flesh produced broad spectra (B and D). This further indicates that the peel extracts at high pH synthesize AgNPs that have a narrow size distribution and uniform shapes, whereas, the low pH conditions synthesized larger AgNPs that are likely to be polydispersed. The pH of the samples is often adjusted to control NP sizes and reaction rate. AgNPs synthesized at low pH aggregate over time and have a wide range of shapes and sizes. On the other hand, AgNPs synthesized at high pH are smaller in size and have uniform shapes [43,46]. Using Lawsonia inermis leaf extracts, AgNPs could only be synthesized at pH 9 and not at pH 4 [46]. Parachlorella kessleri synthesized smaller AgNPs with a core size range of 10–25 nm at pH 10 and larger AgNPs (10–60 nm core size) at pH 4 [43]. Similarly, ascorbic acid and citrate-reduced AgNPs were smaller with uniform shapes at high pH [47]. Based on the results for all pear extracts, pH 10 appeared to be the optimum pH for AgNP synthesis. The UV-Vis spectra for the AgNPs at their native pH (pH 4 for the RPE and GPP extracts, pH 5 for the GPF extracts) were broad and had a low SPR which directly corresponds to non-uniform AgNP shapes and low concentration, respectively [40].

**Effect of AgNO₃ concentration on AgNP synthesis**

The concentration of the metal precursor also plays a crucial role in the synthesis of metallic NPs [40]. Some studies reported a positive correlation between the precursor concentration and the amount of NPs formed. By increasing the concentration of AgNO₃ the rate of Gum tragacanth AgNPs formation was increased [45]. Contrary, more AgNPs were synthesised from Acinetobacter calcoaceticus extracts at low (0.5 and 0.7 mM) than at higher (5 mM) AgNO₃ concentrations. Moreover, lower concentrations of AgNO₃ were associated with smaller and uniform AgNPs compared to higher concentrations [40]. A positive correlation between the concentration of AgNO₃ and AgNPs formation was observed in Figure 4. Although the SPR in the GPF-AgNPs was not as profound when compared to the SPR of the other AgNPs (Figure 4(D)), it was apparent that AgNPs were present and was
supported by the color change. The optimum AgNO₃ concentration for the synthesis of the AgNPs was determined as 1 mM.

**Synthesis and characterization of optimized AgNPs**

The AgNPs were then synthesized using the optimum conditions, that is, 3.125–6.25 mg/mL extract concentration, pH 10, 1 mM AgNO₃, and 100 °C. Figure 5(A) shows the UV-Vis spectra of the RP-AgNPs and GP-AgNPs with their SPR peaks at 408–432 nm. The SPR of the AgNPs is indicated in Table 2. There was a red-shift in the SPR of the RP-AgNPs (432 nm for RPP-AgNPs and 424 nm for RPF-AgNPs) and a blue-shift for the GP-AgNPs (408 nm for GPP-AgNPs, 416 nm for GPF-AgNPs). The absorbance of GPP-AgNPs and RPP-AgNPs at SPR was higher than that of GPF-AgNPs and RPF-AgNPs, which indicated that the concentration of the AgNPs synthesized from the peel extracts was higher than AgNPs from the flesh extracts [42].

The majority of the AgNPs were spherical in shape and had varying core size distributions, as indicated by HRTEM images (Figure 5(B,C)). The RPE produced AgNPs with larger core sizes than the GPE. The core size distribution of the RPE-AgNPs was in the size range of 10–70 nm, with the average core size of 43 nm and 46 nm for RPP-AgNPs and RPF-AgNPs, respectively (Table 2). The core size distributions of the GPE-AgNPs were at 5–40 nm range, with an average core size of 20 nm and 25 nm for GPP-AgNPs and GPF-AgNPs, respectively. The hydrodynamic size distribution of the AgNPs determined by DLS showed larger sizes. (Figure 5(D)). The Z (d.nm) average of the peel AgNPs was smaller than that of the flesh AgNPs (Table 2). These sizes were different from the core size obtained from the HRTEM images (Figure 5(B)).

Figure 2. Effect of temperature and concentration of GPE on the synthesis of GP-AgNPs. The GP-AgNPs were synthesized at 25 °C (A), 70 °C (B), and 100 °C (C).
is due to the fact that the hydrodynamic diameter of the AgNPs was augmented by the phytochemicals on the surface of the AgNPs [5,26], which made their average hydrodynamic size much higher than that of the metal core size [5,26,48]. The size difference in the core and hydrodynamic diameters could be attributed to the content of phytochemical components in the pear extracts. Phenolic acids and flavonoids occur in different combinations and concentrations and can influence the size of the AgNPs. This is supported by Krishnaraj et al. who synthesized AgNPs in the size range of

Figure 3. Effect of pH on the synthesis of AgNPs. UV-Vis spectra of RPP-AgNPs (A), RPF-AgNPs (B), GPP-AgNPs (C) and GPF-AgNPs (D) at 100 °C.

Figure 4. Effect of AgNO₃ concentrations on the synthesis of AgNPs. The AgNPs were synthesized at optimal pear extract concentrations, temperature and different concentrations of AgNO₃. The UV-Vis spectra represent (A) RPP-AgNPs, (B) RPF-AgNPs, (C) GPP-AgNPs and (D) GPF-AgNPs at 100 °C.
20–30 nm using Acalyphia indica plant extracts [49], whereas Banala et al. synthesized AgNPs in the size range of 5–50 nm using Carica papaya leaf extracts [50].

The Pdi of the AgNPs was below 0.6 (Table 2). This also supports the TEM and DLS data that showed that their size distribution range was not too broad. All the AgNPs had negatively charged ζ-potential values (Table 2), thus indicating strong repulsive forces between the AgNPs which will prevent aggregation of AgNPs in solution [48].

**Table 2.** The physico-chemical properties of the optimized GP-NPs and RP-NPs.

| AgNPs      | [Extract] mg/mL | SPR (nm) | Absorbance at SPR | Core size distribution (nm) | Z average (d nm) | Pdi    | ζ-potential (mV) |
|------------|-----------------|----------|-------------------|-----------------------------|-----------------|--------|-----------------|
| RPP-AgNPs  | 3.125           | 432      | 0.852             | 10–70                       | 117.2           | 0.244  | −9.5            |
| RPF-AgNPs  | 6.25            | 424      | 0.425             | 30–65                       | 190             | 0.460  | −1.1            |
| GPP-AgNPs  | 3.125           | 408      | 0.529             | 10–40                       | 157.1           | 0.383  | −7.7            |
| GPF-AgNPs  | 6.25            | 416      | 0.324             | 5–30                        | 168.1           | 0.545  | −3.4            |

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**FTIR analysis of optimized AgNPs**

The FTIR analysis of the pear extracts and AgNPs in Figure 6 shows that the extracts contain the same functional groups.
as those in their respective AgNPs. This indicated that the functional groups present in the pear extracts played a role in the reduction of AgNO₃ and stabilization of the AgNPs [51]. Two prominent absorption peaks were observed at ~1650 and 3300 cm⁻¹. The peaks at ~1650 cm⁻¹ represent the involvement of an amide-bond (NH–C=O) which might probably come from proteins [52]. The peaks at ~3300 cm⁻¹ suggest the presence of hydroxyl or amine groups, which could indicate the interaction between the silver ions (Ag⁺) and hydroxyl or amine groups of the RPE and GPE. These

Figure 6. Comparison of the FTIR spectra of the pear extracts and AgNPs. FTIR of RPP-AgNPs versus RPP (A), RPF-AgNPs versus RPF (B), GPP-AgNPs versus GPP (C), GPF-AgNPs versus GPF (D). Each peak in the AgNPs indicates the functional group of the phytochemical involved in the NP synthesis.
functional groups are among the ones identified in the AgNPs synthesized from the Huangguan pear juice extracts [20], which could be equivalent to the AgNPs synthesized using RPF and GPF extracts. This further implies that proteins, in addition to the reducing sugars and phenolic compounds, found in pear extracts were involved in the synthesis of the AgNPs.

Antibacterial effects of AgNPs

The antibacterial effects of the AgNPs were investigated using agar well diffusion against two Gram-positive (S. aureus, MRSA) and two Gram-negative (E. coli and P. aeruginosa) bacterial strains. The presence of ZOIs was used as a sign of bacterial growth inhibition. The RPE and GPE had no activity on the four strains (data not shown), while they were susceptible to the effects of Pen-Strep, which was used as the positive control. Pen-Strep is a cocktail of two commercial antibiotics, penicillin and streptomycin with well-known antimicrobial activity [53]. AgNPs synthesized from the pear flesh showed higher antibacterial activity than the AgNPs synthesized from the pear peels. As shown in Table 3, the AgNPs synthesized from the Forelle pear extracts (RP-AgNPs) had enhanced antibacterial activity when compared to the AgNPs synthesized from the Packham’s Triumph pear extracts (GP-AgNPs). Huang et al. also demonstrated that AgNPs synthesized from the juice of the Asian Huangguan pear inhibit the growth of P. aeruginosa, E. coli and S. aureus [20]. Huang et al. further demonstrated that the ZOIs were size-dependent and larger than AgNPs produced by citrate reduction [20].

The MICs, lowest concentrations of the samples that inhibited visual growth of the bacteria are recorded in Table 4. The pear extracts showed no effect on bacterial growth. The GPF-AgNPs were the most active among the AgNPs. This was

Table 3. ZOIs for bacteria treated with the AgNPs.

| Bacterial strains | Pen-Strep | GPP-AgNPs | GP-AgNPs | RPP-AgNPs | RPF-AgNPs |
|-------------------|-----------|-----------|----------|-----------|-----------|
| E. coli           | 8         | 1.8       | 3.0      | 2.5       | 4.1       |
| S. aureus         | 6         | 1.5       | 3.5      | 2.9       | 4.0       |
| P. aeruginosa     | 9         | 2.2       | 3.1      | 3.0       | 4.3       |
| MRSA              | 5         | 1.3       | 2.8      | 2.3       | 3.7       |

Table 4. Evaluation of the MICs for the pear extracts and AgNPs.

| Bacterial strains | GPP | GPP-AgNPs | GP  | GPF-AgNPs | RPP | RPP-AgNPs | RPF | RPF-AgNPs |
|-------------------|-----|-----------|-----|-----------|-----|-----------|-----|-----------|
| E. coli           | >500| >500      | >500| >500      | >500| >500      | >500| >500      |
| S. aureus         | >500| >500      | >500| >500      | >500| >500      | >500| >500      |
| P. aeruginosa     | >500| >500      | >500| >500      | >500| >500      | >500| >500      |
| MRSA              | >500| >500      | >500| >500      | >500| >500      | >500| >500      |

Figure 7. Growth inhibitory effects of the AgNPs against the selected bacterial strains. (A) RPP-AgNPs (B) RPF-AgNPs, (C) GPP-AgNPs and (D) GPF-AgNPs. *Statistically significant at p < .05, **p < .001 and p < .0001.
based on the findings that these AgNPs inhibited visible growth for all the bacteria at 500 μg/mL. Of the four strains, *P. aeruginosa* was the only strain that was susceptible to all the AgNPs (Table 4). The antibacterial effects of the test samples were further confirmed by measuring the optical density of the treated bacteria at 600 nm (Figure 7). MRSA was most resistant to the effects of the AgNPs, especially the RPF-AgNPs and GPP-AgNPs which did not affect their growth rates, even at higher concentrations. *S. aureus*, *E. coli*, and to a lesser extent *P. aeruginosa*, showed susceptibility to the AgNPs in a dose-dependent manner. Growth of *P. aeruginosa* was completely inhibited by all the AgNPs at 500 μg/mL. This represents the minimum bactericidal concentration (MBC, lowest concentration that inhibits bacterial growth) of the AgNPs on *P. aeruginosa*. The antibacterial effects were statistically significant from 31.25 μg/mL for *E. coli* and *S. aureus*, and from 250 μg/mL for the other bacterial strains. Overall, the antibacterial effects of the RP and GP AgNPs followed similar trends in all the bacterial strains, except for MRSA. The RPP-AgNPs and the GPF-AgNPs are the only AgNPs that were effective against MRSA. The observed differential antibacterial effects could be due to the differences in the phytochemical composition in the RPE and GPE. However, while the main difference in the two pear variants is that RPE has a higher concentration of anthocyanins than GPE [15], it is unlikely that anthocyanins are the only phytochemicals playing a role in the antibacterial activity of these AgNPs.

### Cytotoxicity of AgNPs

The biocompatibility of the AgNPs was evaluated in RAW 264.7 cells using the MTT assay. The assay is used to quantifying the percentage of metabolically active cells by measuring mitochondrial enzymatic activities. Only viable cells are able to process the yellow MTT dye to produce an insoluble formazan product, which can be solubilized by the addition of DMSO to produce a purple colour that is proportional to the number of live cells [54]. The anthocyanin-rich RP extracts and AgNPs were less toxic to the cells when compared to GP extracts and AgNPs (Figure 8). The RPP and GPF extracts showed negligible toxicity towards the cells (Figure 8(A,D)). In contrast, their respective AgNPs seemed to offer protection to the cells. Significant toxicity was only observed in GPP extracts and AgNPs at concentration ≥250 μg/mL.

### Conclusion

This study demonstrated the successful green synthesis of AgNPs using the aqueous peel and flesh extracts of the Packham’s Triumph and Forelle pear variants. The method of
synthesis used was rapid, cost-effective, and eco-friendly. The GPF and RPP AgNPs exhibited significant antimicrobial activities against the multi-drug-resistant strains, that is, MRSA and P. aeruginosa. The synthesized AgNPs showed differential effects on the proliferation of RAW 264.7 cells, where the RP-AgNPs. Thus, the synthesized AgNPs have a potential for use in bio-applications, such as skincare therapeutics, cosmetics, and the food industry to prevent spoilage. However, the safety of these AgNPs warrants further investigations for their use.

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Disclosure statement

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Data availability statement

The data for the study is presented in the paper as figures and tables.

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