Abstract
Green procedure for synthesizing silver nanoparticles (AgNPs) is currently considered due to its economy and toxic-free effects. Several existing works on synthesizing AgNPs using leaves extract still involve the use of physical or mechanical treatment such as heating or stirring, which consume a lot of energy. To extend and explore the green extraction philosophy, we report here the synthesis and antibacterial evaluations of a purely green procedure to synthesize AgNPs using *Carica papaya, Manihot esculenta, and Morinda citrifolia* leaves extract without the aforementioned additional treatment. The produced AgNPs were characterized using the ultraviolet–visible spectroscopy (UV-vis), field emission scanning electron microscopy (FESEM), energy-dispersive X-ray spectroscopy (EDX), Fourier transform infrared spectroscopy (FTIR), and antibacterial investigations. For antibacterial tests, two bacteria namely *Escherichia Coli* and *Bacillus Cereus* were selected. The presently employed method has successfully produced spherical AgNPs having sizes ranging from 9 to 69 nm, with plasmonic characteristics ranging from 356 to 485 nm, and energy-dispersive X-ray peak at...
approximately 3 keV. In addition, the smallest particles can be produced when *Manihot esculenta* leaves extract was applied. Moreover, this study also confirmed that both the leaves and synthesized AgNPs exhibit the antibacterial capability, depending on their concentration and the bacteria type.

**Keywords:** Carica papaya, *Manihot esculenta*, Morinda citrifolia, green synthesis, silver nanoparticles

**Introduction**

Silver nanoparticles (AgNPs) have been the attraction for various active research subjects in recent years due to their appealing electrical [1], optical [2, 3], physical [4], and thermal behaviors [5]. Their applications are widely found in the biological products [6], photovoltaics [7], catalysts [8], and chemical sensors [9]. Of interest also are their utilization in fabrics [10], antimicrobial coatings [6, 11], biomedical devices [12, 13], and wound dressing owing to their bacterial-repelling property [14, 15]. It is well known that their properties depend strongly on their sizes, species, shapes, and procedures of the synthesis. Studies on the toxicity properties of AgNPs on the bacterial growth have been well-documented in the literature. Colony forming unit (CFU) and inhibition zone are common approaches used to investigate their antibacterial capability. It is established that the nanoparticles have a large surface area compared to micro or macroparticles. Thus, nanoparticles have a higher tendency of interaction with the bacterial cells compared to the bigger particles. AgNPs with the size of 5 nm demonstrated the greatest antibacterial capability against *Escherichia coli* (*E. coli*) MTCC 443 and *Staphylococcus aureus* NCIM 5201 compared to the larger particles such as 7 nm and 10 nm sizes at similar bacterial concentrations [16]. Another study also found that smaller AgNPs with the sizes ranging from 15 to 50 nm exhibited more antibacterial activity than larger particles ranging from 25 to 70 nm, 30 to 80 nm, and 30 to 200 nm against *Pseudomonas aeruginosa* and *E. coli*. In addition, the shape of AgNPs also affected their antibacterial capability [17]. The minimum inhibitory concentration of AgNPs in other shapes such as nanocubes, nanospheres, and nanowires were found to be 37.5, 75, and 100 µg mL⁻¹ when a bacterial concentration of 10⁴ CFU mL⁻¹ was employed. The high antibacterial activity of these anisotropic-shaped AgNPs was due to the basal plane with high-atom-density facets acting as the maximum reactivity sites [17].

Since AgNPs are proven to be toxic, their existence and applications in the environment have become a hot topic for research [18-23]. The presence of AgNPs in the environment is commonly attributed to their release from commercial products. For instance, about 5 to 95% of the total amount of AgNPs in the consumer
products were expelled to sewage treatment plants. In addition, they were released from six types of socks containing AgNPs with the concentrations ranging from 1.5 to 650 µg in 500 mL of the wash water [24]. Shirts, medical masks and cloths, toothpaste, shampoo, detergent, towels, teddy bear toys, and two humidifiers were also found to release AgNPs to the wash water with the concentration of about 45 µg product$^{-1}$ [25]. Furthermore, about 85 g d$^{-1}$ of AgNPs were discharged from the laundry into the sewer system and the municipal wastewater treatment plant [26]. It is of importance to note that AgNPs in the environment can be transformed into different forms, such as ionic Ag, Ag$_2$O, and Ag$_2$S depending on the environmental conditions via physical and chemical events [27].

There exist many works on synthesizing and characterizing numerous properties of AgNPs in the past several years [28-34]. Since understanding how their various properties change corresponding to different extracting techniques is vital to achieve their most desirable performance, several production approaches were continually proposed and revised. Although many extracting methods are possible, the current trend focuses on how this can be carried out in a cheaper and greener manner. Currently, synthesis of AgNPs using plant extracts is highly considered because of their simplicity, abundant availability, and potential to eliminate a complicated preparation such as the use of microorganism as presented by Mishra et al. [34]. The use of leaves, seeds, roots, and fruits in the green synthesis of AgNPs have been continually proposed and remain an active subject for research investigation. Among these, the exploration of leaves extract as the reducing and stabilizing agent to synthesize nanoparticles of AgNPs is a preferable approach due to their wide obtainability in the environment. Table 1 lists the recent pattern of AgNPs synthesis using numerous leaf extracting approaches [4, 31, 32, 35-41]. Several works, for instance Salem et al. [40], Vijay Kumar et al. [41], Muthukrishnan et al. [38], and He et al. [33], on the production of AgNPs still employed physical or mechanical treatments such as heating or stirring, which consume a lot of energy.

Therefore, the aim of the present study is to evaluate the suitability of the use of Carica papaya (C. papaya), Manihot esculenta (M. esculenta), and Morinda citrifolia (M. citrifolia) leaves extract without the aforementioned additional treatment to synthesize AgNPs. It is essential to provide a purely green procedure for future exploration to minimize the use of energy, its massive consumption of which is evidenced in other alternative approaches. C. papaya is a climacteric tropical plant originated from the Southern Mexico. It is also generally well-populated in the tropical climate such as Brazil, Malaysia, and Indonesia. In addition, M. esculenta, generally known as Cassava, is well recognized as a carbohydrate source. In the tropical region, it is the third largest source of carbohydrates after rice and maize [42]. Furthermore, M. citrifolia is a medical plant
that is widely grown in Malaysia and Indonesia. In these countries, it is well known as Mengkudu widely
explored for numerous medical applications. Leaves of these plants are commonly utilized as a vegetable source
as well [43, 44]. For comprehensive details, the chemical compositions of these three types of leaf are offered in
Table 2 [42, 45-48]. Many existing works have confirmed that the biomolecules contained in leaves such as
proteins, enzymes, polysaccharides, amino acids, and vitamins can act as bioreductant from reducing metal ions
to the formation of AgNPs in the solution [49, 50]. The use of all currently proposed leaves is due to their
possession of these favorable properties as presented in Table 2. Also, since previous studies have proven that
these leaves have antibacterial capability, their exploration either as extract or capping on AgNPs are interesting
for further research pursuit. Such study can be highly beneficial for future medical applications.

Materials and Methods

Materials

The basic compound under study, silver nitrate (AgNO₃, QReC, Auckland, New Zealand), was used as the silver
salt. C. papaya, M. esculenta, and M. citrifolia leaves were collected from the surrounding area of Universiti
Teknologi Malaysia, Johor Bahru, Malaysia. 0.45 µm nylon membrane (Whatman® Nylon membrane, Sigma-
Aldrich, St. Louis, MO, USA) was used as the filter. For synthesizing process, solutions were prepared using the
ultrapure water (resistivity 18.2 MΩ cm) (Arium Ultrapure Water System, Sartorius Malaysia Sdn Bhd, Kuala
Lumpur, Malaysia). Bacteria, E. coli and Bacillus cereus (B. cereus), were obtained from the Faculty of
Biosciences and Medical Engineering, Universiti Teknologi Malaysia. In addition, agar powder (OXOID
CM0003, Oxoid Ltd, Cheshire, England) was used. Broth powder (Merck VM447243, Merck KGaA,
Darmstadt, Germany) was also employed in this study.

Preparation of leaves extract

To remove impurities, fresh leaves were first washed using the tap water and followed by the ultrapure water
three times each. Then, 10 g of leaf was mixed with 300 mL ultrapure water in a 500 mL Erlenmeyer flask. The
mixture was heated to 250 °C for 30 min, before cooled at the room temperature. To obtain a pure leaves extract,
the mixture was then filtered through a nylon membrane filter of 0.45 µm. The solution passing the membrane
was then stored in a fridge at a temperature of 7 °C for future use. The same extraction procedure was then
applied for all proposed leaves.
Synthesis of AgNPs

Firstly, AgNO$_3$ solution (100 mL in volume) was prepared using AgNO$_3$ and the ultrapure water with a concentration of 0.15 M in a 500 mL Erlenmeyer flask. To initiate the synthesis process, 100 mL leaves extract was added slowly into the AgNO$_3$ solution. The mixtures were then left overnight to perform the reduction of metal ions to the formation of AgNPs in the solution. For control, a 100 mL AgNO$_3$ solution with the same concentration without leaves extract was also prepared. For AgNPs harvesting, the mixtures were then centrifuged at 8000×g for 45 min and the pellet obtained in this process was collected. To purify AgNPs production, the pellet was then cleansed with the ultrapure water before centrifuged at 8000×g for 45 min. The purification process was repeated for three times. Next, the pellet was air dried. AgNPs obtained using this procedure were then stored for the characterization. The same procedure was carried out for synthesizing AgNPs using other leaves extract.

Plasmonic investigation

Plasmonic property of AgNPs was characterized using the UV-Vis spectrometer (Perkin–Elmer, No. 101N4110104) installed with the Lambda 25 software. It was operated at a resolution of 1 nm, a scan speed of 960 nm min$^{-1}$, and the electromagnetic wavelength in the range of 300 to 700 nm. 2 mL sample mixtures were injected into the UV-vis tube. In this inspection, the ultrapure water was used as a blank.

FTIR characterization

For this observation, AgNPs were mixed with potassium bromide (1:100) to produce the specimen in a pellet form. The pellet was pressed hydraulically (Specac model with a serial number of N29850) at 10 tons in pressure. The pressed pellet was then taken and placed into the FTIR holder. Biomolecules bonding of the synthesized AgNPs was identified using the FTIR spectrometer (PerkinElmer Frontier-GPOB model 96046) installed with the PerkinElmer Spectrum software. The spectrometer used OptKBr (7800 to 400 cm$^{-1}$) as beam splitter and MIR TGS (15000 to 370 cm$^{-1}$) as detector. This characterization was conducted using a spectrum wavelength in the range of 650 to 4000 cm$^{-1}$ at a resolution of 4 cm$^{-1}$ and accumulations of 10 scans at room temperature.
FESEM and SEM-EDX

AgNPs morphologies were also characterized by means of the field emission scanning electron microscopy (FESEM ZEISS Supra 35VP). This apparatus was supplied by Carl Zeiss Sdn Bhd. It was operated at an accelerating of 5 kV with a magnification of 50000×. AgNPs elements were then confirmed by SEM-EDX (HITACHI S-3400N) equipped with the Bruker Quantax software. It was operated at a voltage of 15 kV.

Preparation of agar and broth nutrients

To produce agar nutrient, 14 g of the nutrient agar powder was mixed with 500 mL of ultrapure water. In addition, 4 g of the broth powder was prepared with the same water mixture to produce the broth nutrient solution. In the preparation, all solutions were sterilized using the autoclave ALP (model CL-40M no.805415) at a temperature of 121 °C for 2 h.

Colony forming test

Colony forming test was prepared using the previous study by Mueller and Hinton [51] as basis. AgNO₃ and leaves extract were freshly prepared and mixed for this test. A 10-mL AgNO₃ solution (0.15 M) was prepared using the ultrapure water in a 50 mL plastic tube. A variation in the leaf quantity was made such that ratios of 5:2 and 5:3 for AgNO₃ and leaves extract were obtained. AgNO₃ solution with the same concentration was also prepared as a control case. Bacterial cultures of about $10^5$ × colony were taken and about 0.1 mL of inoculum was mixed with the solution and incubated at 37 °C for 24 h. For comparison purpose, the pure leaves extract was also studied. Then, the colony of the survived bacteria was counted. This test procedure was carried out for all considered leaves.

Inhibition zone test

In this investigation, antibacterial activity was determined by using the paper disk assay method as basis [52]. In this test, AgNO₃ and leaves extract were freshly prepared and mixed analogously to the method employed in the colony forming test. Bacterial cultures of about $5 \times 10^1$ colony were taken and about 0.1 mL of inoculum was spread on each agar plate. Filter paper (2 mm in diameter) was steeped in AgNPs solution with the similar variation as the colony forming test for 1 min and then put onto the agar plate. As comparison, the filter paper was also steeped in AgNO₃ solution. Next, the plate was incubated at 37 °C for 24 h. An inhibition zone can be described as the clear area surrounding the filter paper containing AgNP solution deposited on the plate. The
diameter that defines inhibition zone was then measured using a ruler. This test procedure was similarly performed for all other test samples.

Results and Discussions

Plasmonic property

Surface plasmon resonance is well known as the collective oscillation of the electrons in the conduction band when the particles absorb the electromagnetic wave [53]. Theoretically, AgNP aggregation and dispersion phenomena can be identified with the UV-vis absorption spectra. UV-vis spectra of AgNPs synthesized using different leaves extract are depicted in Fig. 1. For comparison purpose, UV–vis spectra for AgNO₃ solution were also included. Figs. 2a-2c show the solution color of AgNPs synthesized using *C. papaya*, *M. esculenta*, and *M. citrifolia*, respectively. It is clear that the reduction of silver ion to the formation of AgNPs in the solution occurred as indicated by the change in solution color from yellow to brown or reddish yellow to deep red (see Fig. 2).

From UV-vis spectra, the peak absorbance can be found at 485, 356, and 471 nm for AgNPs synthesized using *C. papaya*, *M. esculenta*, and *M. citrifolia*, respectively (see Fig. 3). These findings are in agreement with those found in the previous works [4, 54]. This observation has confirmed that AgNPs reduced and stabilized using different leaves extract exhibit different characteristics in their plasmonic property. Theoretically, AgNPs have spectra in the visible region ranging from 380 to 480 nm due to the excitation of localized surface Plasmon resonance [55, 56]. It is well known that their surface plasmon oscillation is extremely affected by their size, shape, and surrounding media as well as treatment employed [3, 57]. AgNPs in spherical shape can be correlated with a single peak in the UV-vis spectrum [58]. On the other hand, AgNPs in the irregular shapes have two or more peaks depending on their symmetry. The spectra characteristics exhibited in Fig. 1 suggest that AgNPs synthesized using *C. papaya*, *M. esculenta*, and *M. citrifolia* were spherical in nature.

For a comprehensive overview, AgNPs synthesized using *C. papaya* shows the highest maximum peak and followed by those using *M. esculenta* and *M. citrifolia* (see Fig. 3). The different maximum peaks of their UV-vis characteristics can be associated with the size of AgNPs. For instance, increasing the size of spherical nanoparticles from 8 to 99 nm increased their maximum peak spectra from 517 to 575 nm [59]. It was measured from the current study that the absorbance at \( A_{\text{max}} \) of AgNPs synthesized using *C. papaya*, *M. esculenta*, and *M. citrifolia* are 1.56, 1.24, and 0.82 a.u, respectively. The different absorbance of the UV-vis spectra can be
specifically related to the agglomeration of AgNPs. The increase in the absorbance spectra indicates a higher production of AgNPs [60].

FTIR characteristics

FTIR was inspected to analyze biomolecule compounds, which can act as the capping agent for stabilizing AgNPs production. This method has previously been established and proved to be reliable [4, 61]. For this purpose, the FTIR spectra of AgNPs synthesized using different leaves extract are depicted in Fig. 4. Several prominent peaks are observed around the wave numbers ranges of 1042 to 1084 cm$^{-1}$ (region I), 1384 to 1394 cm$^{-1}$ (region II), 1590 to 1619 cm$^{-1}$ (region III), and 3301 to 3444 cm$^{-1}$ (region IV).

IR bands around 1042 to 1084 cm$^{-1}$ are characterized as the phosphorus compounds. It was obvious in Table 2 that all leaves used in this study contained phosphorus mineral. FTIR spectra around 1384 to 1394 cm$^{-1}$ are associated with the nitro compounds, which are reported also in AgNPs synthesized using *Cleistanthus collinus* leaves extract [62]. In addition, IR bands around 1590 to 1619 cm$^{-1}$ are associated with the C=C stretching modes of vibration [63]. Finally, spectra peaks within 3301 to 3444 cm$^{-1}$ are the -NH stretching modes, which were also reported by Jeyaraj et al. [61], which used *Sesbania grandiflora* leaves extract.

In region I of the wave number, AgNPs synthesized using *M. esculenta* are more intense in terms of transmittance compared to the others. The similar characteristics were also observed in region II. In region III, the transmittance of AgNPs synthesized using *C. papaya* is significantly intense compared with those nanoparticles synthesized using *M. esculenta*, and *M. citrifolia*. In the region IV, AgNPs synthesized using *M. esculenta* did not have a prominent peak. It is apparent that there were slightly different peak intensities for all synthesized AgNPs. Such difference and obvious characteristics can be correlated with the concentration of biomolecule on the surface of AgNPs [64].

Morphology and size

Morphology and size of AgNPs synthesized by chemical, physical, and biological approaches are affected by the reducing agent, stabilizer, and surrounding medium. In biological synthesis particularly that using leaves extract, their properties are extremely affected by the chemical composition of the leaves. In the present work, the shape of AgNPs synthesized using *C. papaya*, *M. esculenta*, *M. citrifolia* leaves extract is found to be spherical (see Figs. 5a-5c). It is confirmed that the single peak in the UV-vis spectra (see Fig. 1) for all synthesis solution is related to the spherical AgNPs.
Specifically, this work found that AgNP sizes synthesized using *C. papaya* were concentrated within 13 to 69 nm with an average of 40.8 nm (see Fig. 6a). In addition, AgNPs sizes synthesized using *M. esculenta* were in the range of 13 to 38 nm with an average of 23.0 nm (see Fig. 6b). For those synthesized using *M. citrifolia*, the range was 9 to 54 nm with an average of 26.5 nm (see Fig. 6c). By this procedure, the smallest AgNPs size was produced when *M. esculenta* leaves extraction was applied, followed by those extracted using *M. citrifolia* and *C. papaya*. These findings match with their previously discussed plasmonic properties.

In general, results from this study enhance the understanding of the effectiveness of the use of local leaves for synthesizing AgNPs in a purely green fashion at room temperature. It is noteworthy to see that AgNP sizes obtained using the presently employed procedure are comparable with those employing physical or mechanical treatment (see Table 1). Moreover, for certain studies such as proposed by Muthukrishnan et al. [38] and Balan et al. [35], results from this study are preferable in terms of their size. Also, the present green procedure has successfully produced AgNPs smaller than those obtained by Dipankar and Murugan [36], Prakash et al. [39], as well as the recent study by Kharat and Mendhulkar [37] and Ashraf et al. [4], who also synthesized AgNPs using leaves extract without additional treatment.

**Energy dispersive characteristic**

Energy dispersive X-ray spectroscopy, which is sometimes abbreviated as EDS, EDX, or XEDS is a common procedure for analyzing the basic elements on the surface of sample. Their characteristics generally depend on the source of X-ray excitation and the sample. The SEM-EDX spectra of all synthesized AgNPs can be observed in Figs. 7a-7c. The spectra characteristics have confirmed the presence of AgNPs. The sharp signal peak of the spectrum exhibits that the reduction from AgNO$_3$ to AgNPs using *C. papaya*, *M. esculenta*, and *M. citrifolia* were successfully carried out.

Specifically, the percentages of AgNPs synthesized using *C. papaya*, *M. esculenta*, and *M. citrifolia* observed in the EDX spectra are 94.69, 92.44, and 99.22%, respectively. It is ratified that AgNPs are dominant in the samples compared to other elements. The spherical AgNPs synthesized using *C. papaya* exhibit the EDX peak at approximately 3 keV. In addition, similar characteristics were also observed for those synthesized using *M. esculenta* and *M. citrifolia*. Such peaks are typical absorptions of metallic AgNPs due to the surface plasmon resonance [65]. These findings are in line with the results from previous works, which also synthesized AgNPs using leaves extract [4, 66]. The highest peaks in Figs. 3a-3c at approximately 3 keV confirm that metal AgNPs are dominant element compared to others. In addition, other smaller elemental peaks shown in Figs. 6a-6c are
possibly due to the contribution from enzymes or proteins present within *C. papaya*, *M. esculenta*, and *M. citrifolia* leaves. In general, this study has proven that the synthesis using different leaves extract such as *C. papaya*, *M. esculenta*, and *M. citrifolia* can produce AgNPs with different properties in terms of plasmonic, molecule bonding, morphology, and energy dispersive. It is well-known that the synthesis of AgNPs can be divided into three categories namely physical, chemical, and biological. In the physical approach, synthesis AgNPs using laser ablation, small ceramic heater, and thermal decomposition methods has been established. For the chemical approach, this requires three main ingredients namely a silver salt, a reducing agent, and a stabilizing or capping agent. In the biological approach, the reducing and stabilizing agents are replaced using biomolecules obtained from the living organism such as plants that are explored in this work. Proteins, enzymes, and vitamins are the biomolecules contained in the employed leaves extract (see Table 2) that can act as bioreductant from reducing metal ions to the formation of AgNPs in the solution. Moreover, the proposed purely green procedure is environmentally friendly compared to chemical or physical approach since the latter uses the chemical as reducing and stabilizing agent that may dispel toxin to the environment, in addition to their employment of huge amount of energy in the production.

**Antibacterial investigation**

*Antibacterial activity by all leaves*

Various studies had reported that *C. papaya*, *M. esculenta*, and *M. citrifolia* leaves extract can be used as antibacterial agents [67, 68]. Their studies confirmed that the antibacterial activity is strongly affected by the medium, extraction process, and type of bacteria. As summarized in Table 3, it can be seen that *C. papaya* inhibited bacteria colony growth of *E. coli* better than *M. esculenta* and *M. citrifolia*. These are supported by the inhibition zone observation also shown in Table 3. *C. papaya* leaves extract is able to inhibit *E. coli* growth to about 1 cm in zone diameter. Recently, *C. papaya* leaves extract was reported to act as an effective antibacterial component with the zones of inhibition in the range of 14 to 16 mm against *Staphylococcus aureus*, *E. coli*, *B. cereus*, and *Pasteurellamultocida* [67].

In inhibiting the *B. Cereus* growth, this study notices that *M. esculenta* is the most effective. It inhibits the growth of *B. Cereus* to about 1.2 cm in diameter. *M. esculenta* leaves extract can be used as an effective antibacterial against gram-positive bacteria such as *Corynebacterium diphtheriae*, *Listeria monocytogenes*, *Pseudomonas aeruginosa* and gram-negative bacteria such as *Vibrio cholera*, *Shigella flexneri*, and *Salmonella typhi* [69]. Their extract is also effective as an antibacterial against *Pseudomonas aeruginosa*, *E. coli*,
Enterobacter cloacae, Klebsiella pneumoniae, Providencia stuartii, and Enterobacter aerogenes [68]. Phenolic compounds such as acubin, l-asperuloside, alizarin, and scopoletin are the chemical composition that act towards their antimicrobial activity. On the other hand, it is worthwhile to see that M. citrifolia did not show any antibacterial activity, a phenomenon evidenced in both colony forming and inhibition zone studies.

Antibacterial activity by synthesized AgNPs

AgNPs have been widely proven as an antibacterial and an antimicrobial agent [6, 15]. Table 4 lists the outcomes from the colony forming unit test of synthesized AgNPs against E. coli and B. cereus. It is apparent that AgNPs have completely (100%) inhibited the bacteria colonies of E. coli and B. cereus from $10^{15}$ initial colony to 0 colony as observed in the agar plate. In comparison, about $60 \times 10^1$ bacterial colonies were still existed when the AgNO$_3$ solution was employed against B. cereus for control. In addition, the AgNO$_3$ solution also completely inhibited the colony of E. coli. To the best of the authors knowledge, this is the first evidence that shows that AgNPs synthesized using local plants such as C. papaya, M. esculenta, and M. citrifolia can act as an antibacterial agent against E. coli and B. cereus. Further proof is shown by the inhibition zone test in Table 5. It is noticed that AgNPs synthesized using C. papaya have inhibition zones varying from 1.8 to 2.6 cm. In addition, AgNPs synthesized using M. esculenta and M. citrifolia have inhibition zones varying from 1.7 to 2.0 cm and 1.6 to 2.2 cm, respectively. As a control, the AgNO$_3$ solution has inhibition zones about 0.7 and 1.0 cm against E. coli and B. cereus, respectively. Moreover, AgNPs synthesized using C. papaya show the largest average inhibition zone compared to others.

Although the mechanism of the bactericidal effect of AgNPs is still not well understood, several works provided some initial evidence that AgNPs can be used as an effective antibacterial agent. In terms of toxicity effect, the bacterial cell death was caused by the interaction between AgNPs and constituents of the bacteria membrane as a result of its structural change and the eventual damage [18]. AgNPs can be categorized as a hydrophobic material. Although they are hydrophobic, their bioaccumulation mechanism takes the role of their antibacterial properties [70]. AgNPs tend to accumulate at the bacterial membrane and further form aggregates once they are in contact with the organism. Consequently, the diminishment of the bacterial membrane integrity and its damage lead to bacteria cellular death [70]. There is also an implication that AgNPs can penetrate into the bacteria membrane [71]. Moreover, AgNPs can inhibit bacteria respiratory enzyme, which then facilitates the generation of reactive oxygen and consequently damages the cell [72]. Alternatively, the toxicity of AgNPs
is surface charge-dependent affected by the capping agent and synthesis procedure. The more negative charged AgNPs is less toxic compared with those nanoparticles having a more positive charge [64].

Conclusion

The aim of this investigation is to evaluate the effectiveness of the green procedure to synthesize AgNPs using C. papaya, M. esculenta, and M. citrifolia leaves extract without additional treatment such as physical or mechanical. This study has identified that the plasmonic property of AgNPs differs according to their size and reducing agent as well as stabilizing agent with AgNPs synthesized using C. papaya having the highest maximum peak and absorbance compared to the others. In addition, FTIR characteristics have confirmed the biomolecule bonding on AgNPs. FESEM investigations revealed that spherical AgNPs ranging from 9 to 69 nm were produced using this simple procedure. Specifically, the smallest nanoparticles can be produced when M. esculenta leaves were employed. Also, both leave extracts and synthesized AgNPs showed great potential as an antibacterial component against E. coli and B. cereus. In detail, C. papaya leaves extract was the most effective antibacterial agent against E. coli compared to the others. For B. cereus, M. esculenta leaves extract was the most effective to inhibit the bacteria. AgNPs synthesized using all considered leaves extract were found to completely inhibit all bacteria colonies. In addition, AgNPs synthesized using C. papaya showed the largest inhibition zone compared with those nanoparticles synthesized using M. esculenta, and M. citrifolia.

This work extends the knowledge for the exploration of natural resources particularly C. papaya, M. esculenta, and M. citrifolia as reducing and stabilizing agents for a green synthesis of AgNPs. These findings have significant implications for future medical applications particularly for providing potential antibacterial agent from natural resources and synthesized AgNPs. A further study could assess the long-term stability of AgNPs using this procedure and investigate their antibacterial performance on other bacteria.

Acknowledgements

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Conflict of Interest:

The authors declare that they have no conflict of interest.
References


Figure Captions

Fig. 1. UV-vis spectra of AgNPs synthesized by different leaves extract

Fig. 2. The solution color of AgNPs synthesized using (a) C. papaya, (b) M. esculenta, and (c) M. citrifolia

Fig. 3. (a) absorbance and (b) wavelength of all solutions. Note that I, II, and III refer to the AgNO$_3$+ $M.$
citrifolia, AgNO$_3$+ $M.$ esculenta, and AgNO$_3$+ C. papaya, respectively.

Fig. 4. FTIR spectra of AgNPs synthesized using different leaves extract

Fig. 5. FESEM of AgNPs synthesized using (a) C. papaya, (b) M. esculenta, and (c) M. citrifolia

Fig. 6. Size distribution histograms of all synthesized AgNPs

Fig. 7. EDX spectra of AgNPs synthesized using (a) C. papaya, (b) M. esculenta, and (c) M. citrifolia
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Fig. 7. EDX spectra of AgNPs synthesized using (a) *C. papaya*, (b) *M. esculenta*, and (c) *M. citrifolia*
Table captions

Table 1. Shape and size of AgNPs synthesized using various leaves extract

Table 2. Chemical compositions of *C. papaya*, *M. esculenta*, and *M. citrifolia* [42, 45-48]

Table 3. Effects of different leaves extract on the colonies and inhibition zones of *E. Coli* and *B. Cereus*

Table 4. Effects of AgNPs presence on the colonies of *E. Coli* and *B. Cereus*

Table 5. Effects of AgNPs presence on the inhibition zones of *E. Coli* and *B. Cereus*
Table 1. Shape and size of AgNPs synthesized using various leaves extract

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<th>Additional treatment</th>
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<td><em>Mimusops elengi</em></td>
<td>No</td>
<td>55 to 83</td>
<td>Prakash et al. [39]</td>
</tr>
<tr>
<td><em>Aloe vera</em></td>
<td>No</td>
<td>5 to 85</td>
<td>Ashraf et al. [4]</td>
</tr>
<tr>
<td><em>Elephantopus scaber</em></td>
<td>No</td>
<td>78</td>
<td>Kharat and Mendhulkar [37]</td>
</tr>
<tr>
<td><em>Carica papaya</em></td>
<td>No</td>
<td>13 to 69</td>
<td>Present work</td>
</tr>
<tr>
<td><em>Manihot esculenta</em></td>
<td>No</td>
<td>13 to 38</td>
<td>Present work</td>
</tr>
<tr>
<td><em>Morinda citrifolia</em></td>
<td>No</td>
<td>9 to 54</td>
<td>Present work</td>
</tr>
</tbody>
</table>
Table 2. Chemical compositions of *C. papaya*, *M. esculenta*, and *M. citrifolia* [42, 45-48]

<table>
<thead>
<tr>
<th>Leaves extract</th>
<th>Proximate</th>
<th>Mineral</th>
<th>Vitamin</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>C. papaya</em></td>
<td>Lipid</td>
<td>Calcium</td>
<td>Vitamin A</td>
</tr>
<tr>
<td></td>
<td>Protein</td>
<td>Magnesium</td>
<td>Vitamin B12</td>
</tr>
<tr>
<td></td>
<td>Crude fibre</td>
<td>Phosphorus</td>
<td>Vitamin C</td>
</tr>
<tr>
<td></td>
<td>Moisture</td>
<td>Iron</td>
<td>Vitamin E</td>
</tr>
<tr>
<td></td>
<td>Ash</td>
<td></td>
<td>Niacin</td>
</tr>
<tr>
<td></td>
<td>Carbohydrate</td>
<td></td>
<td>Thiamine</td>
</tr>
<tr>
<td></td>
<td>Fat</td>
<td></td>
<td>Riboflavin (Vitamin B2)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Beta-carotene</td>
</tr>
<tr>
<td><em>M. esculenta</em></td>
<td>Caloric content</td>
<td>Calcium</td>
<td>Ascorbic acid</td>
</tr>
<tr>
<td></td>
<td>Water</td>
<td>Copper</td>
<td>Niacin</td>
</tr>
<tr>
<td></td>
<td>Protein</td>
<td>Iron</td>
<td>Riboflavin (Vitamin B2)</td>
</tr>
<tr>
<td></td>
<td>Lipid</td>
<td>Magnesium</td>
<td>Thiamin</td>
</tr>
<tr>
<td></td>
<td>Carbohydrates</td>
<td>Manganese</td>
<td>Vitamin A</td>
</tr>
<tr>
<td></td>
<td>Fiber</td>
<td>Phosphorous</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Ash</td>
<td>Potassium</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Sodium</td>
<td></td>
</tr>
<tr>
<td><em>M. citrifolia</em></td>
<td>Water</td>
<td>Calcium</td>
<td>Ascorbic acid</td>
</tr>
<tr>
<td></td>
<td>Protein</td>
<td>Phosphorous</td>
<td>Niacin</td>
</tr>
<tr>
<td></td>
<td>Fat</td>
<td>Iron</td>
<td>Riboflavin (Vitamin B2)</td>
</tr>
<tr>
<td></td>
<td>Carbohydrate</td>
<td></td>
<td>Thiamin</td>
</tr>
<tr>
<td></td>
<td>Fiber</td>
<td></td>
<td>Beta-carotene</td>
</tr>
<tr>
<td></td>
<td>Ash</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
Table 3. Effects of different leaves extract on the colonies and inhibition zones of *E. Coli* and *B. Cereus*

<table>
<thead>
<tr>
<th>Leaves extract</th>
<th>Colony</th>
<th>Inhibition zone (cm)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td><em>E. Coli</em></td>
<td><em>B. Cereus</em></td>
</tr>
<tr>
<td><em>Carica papaya</em></td>
<td>$2 \times 10^{11}$</td>
<td>$29 \times 10^{9}$</td>
</tr>
<tr>
<td><em>Manihot esculenta</em></td>
<td>$66 \times 10^{11}$</td>
<td>$4 \times 10^{9}$</td>
</tr>
<tr>
<td><em>Morinda citrifolia</em></td>
<td>$8 \times 10^{11}$</td>
<td>$4 \times 10^{15}$</td>
</tr>
</tbody>
</table>
Table 4. Effects of AgNPs presence on the colonies of *E. Coli* and *B. Cereus*

<table>
<thead>
<tr>
<th>Type of Leaf</th>
<th><em>E. Coli</em></th>
<th></th>
<th></th>
<th><em>B. Cereus</em></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>AgNO$_3$ + and</td>
<td>leaves extract (5:2)</td>
<td></td>
<td>AgNO$_3$ + and</td>
<td>leaves extract (5:3)</td>
<td></td>
</tr>
<tr>
<td><em>C. papaya</em></td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td><em>M. esculenta</em></td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td><em>M. citrifolia</em></td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>
**Table 5. Effects of AgNPs presence on the inhibition zones of *E. Coli* and *B. Cereus***

<table>
<thead>
<tr>
<th>Type of Leaf</th>
<th><em>E. Coli</em></th>
<th></th>
<th><em>B. Cereus</em></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>AgNO$_3$ + and leaves extract (5:2)</td>
<td>AgNO$_3$ + and leaves extract (5:3)</td>
<td>AgNO$_3$ + and leaves extract (5:2)</td>
<td>AgNO$_3$ + and leaves extract (5:3)</td>
</tr>
<tr>
<td><em>C. papaya</em></td>
<td>1.8 cm</td>
<td>2.6 cm</td>
<td>1.7 cm</td>
<td>1.8 cm</td>
</tr>
<tr>
<td><em>M. esculenta</em></td>
<td>1.7 cm</td>
<td>2.0 cm</td>
<td>1.7 cm</td>
<td>2.0 cm</td>
</tr>
<tr>
<td><em>M. citrifolia</em></td>
<td>1.6 cm</td>
<td>2.2 cm</td>
<td>1.3 cm</td>
<td>2.4 cm</td>
</tr>
</tbody>
</table>