Comparative Evaluation of Antibacterial Activity of Silver Nanoparticles Biosynthesized Using Fruit Juices

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Abstract

Synthesis of nanosized particles with antimicrobial property is of significance in therapeutic applications of nanotechnology. The present study reports an environmental friendly and rapid method for biosynthesis of silver nanoparticles and their antibacterial activity. This green synthetic method used various fruit juices like sweet lime, lime and orange as a reducing and capping agent for silver nitrate. Different thermal reduction methods-microwave oven and hot plate were used for synthesizing silver nanoparticles. The silver ions were reduced into silver nanoparticles within few minutes of reaction. Silver nanoparticles so prepared were characterized using UV–visible spectrophotometer and scanning electron microscope. An effort was also been made to predict the size of the silver nanoparticles using UV-visible spectra by Mie Scattering protocol. Scanning electron micrograph (SEM) revealed useful information about the morphology of silver nanoparticles. The synthesized silver nanoparticles showed antibacterial property against pathogenic bacteria Escherichia coli, Salmonella typhimurium, Klebsiella pneumoniae and Staphylococcus aureus. Agar well diffusion method was adopted to assay the nanoparticles for bactericidal activity against test organisms on nutrient agar plates. Silver nanoparticles biosynthesized from orange fruit juice were more effective as antibacterial agent. The maximum zones of inhibition of 8.0 mm, 6.0 mm, 8.0 mm and 5.0 mm were observed against Escherichia coli, Salmonella typhi, Klebsiella pneumoniae and Staphylococcus aureus respectively. The study shows that gram negative bacteria are more susceptible to antibacterial action of silver nanoparticles. Such studies are crucial in the demonstration of therapeutic importance of silver nanoparticles.

Keywords: Fruit juice; Biosynthesis; Silver nanoparticles; Characterization; Antibacterial effect

Introduction

The development of reliable green processes for the synthesis of silver nanoparticles is an important aspect of current nanotechnology research. Noble metal nanoparticles have been the subject of focus due to their unique optical, electronic, mechanical, magnetic and chemical properties. Preparation of silver nanoparticles has attracted particularly considerable attention due to their diverse properties and uses like magnetic and optical polarizability, catalysis [1], electrical conductivity [2] antimicrobial activities [3-5], DNA sequencing [6] and surface-enhanced Raman scattering [7]. Nanomaterials such as Ag, Au, Pt, and Pd have been synthesized by different methods, including hard template [8], using bacteria [9], fungi [10] and plants [11]. The strong toxicity of silver nanoparticles have been recently shown to be promising antimicrobial material. Sondi et al. studied the antimicrobial activity of silver nanoparticles against Escherichia coli as a model of Gram-negative bacteria [12]. The health benefits provided by fruits and vegetables can be attributed to many biologically active phytochemicals present. These biological activities include functions like detoxification, immuno-protection, antiviral, anticancer and antioxidant properties [13]. Employing nano-biotechnological protocols have observed an increase in the area of synthesis of nanomaterials employing microbes, plants/plant parts and conglomerate of nutrients/biochemicals like honey. A quest for an environmentally sustainable synthesis process has led to a few biomimetic approaches. Sometimes the synthesis of nanoparticles using plants or parts of plants can prove advantageous over other biological processes by eliminating the elaborate processes of culturing of microorganisms. Among metal nanoparticles, silver nanoparticles (AgNPs) have been known to have inhibitory and antimicrobial activity. Considering the antioxidant properties of fruit juices, we have made an attempt to use fruit juice as conglomerate of metabolites as reducing and stabilizing agent for the biosynthesis of metal nanoparticles.
The contents were taken in a Teflon container and placed in a microwave oven for three minutes at medium heat setting. The color change from colorless to brown was observed. Similar microwave irradiation was carried out in a glass container. The color change was observed from colorless to brown for 1 mM and colorless to dark brown for 10 mM. Similarly for hot plate method, silver nitrate base (1 mM and 10 mM) and lemon juice were mixed in a conical flask at the ratio of 4:1 and 2:1. The conical flask was kept on a hotplate and heated at about 80°C for 10-15 minutes. Color change was observed from colorless to yellow for 1 mM and colorless to dark brown for 10 mM. Same process was applied for all the three juices (orange, sweet lime and lime).

**Characterization by UV-Vis spectroscopy**

Nanoparticles are made directly as powders and must be stored in a liquid medium for suspension. If the nanoparticles are aggregated, they will build a solid bridge causing them to lose their nanoparticle properties. All the samples are prepared and stored in double distilled water. The silver nanoparticles were characterized by UV-Vis Spectroscopy, one of the most widely used techniques for structural characterization of silver nanoparticles [14]. Aliquots of solutions of different concentrations (25% and 50%) were ultrasonicated to disperse particles and were measured using UV-Vis spectrophotometer (Double beam, Make: Labomed, USA). An aliquot of the sample was also used for scanning electron microscopic studies, 25 µl of sample was sputter coated on copper stub and the images of nanoparticles were studied using SEM.

**Characterization by SEM**

The synthesized silver nanoparticles were characterized using scanning electron microscope. Usually scanning electron microscope is used to characterize the internal properties like exact size, shape, dimensions of nanoparticles. A thin layer of AgNPs solution was placed on sample holder (silica piece). The care was taken that bubbles should not be present. Next the sample holder was kept in the dessicator for half an hour. Further the sample holder containing dry sample was exposed to gold sputtering. This is done to maintain the electron channeling. When sample containing silica piece is mounted into the SEM equipment, the electron beam was made to fall on sample for characterization. If the beam is of high intensity then flow of electron channeling is not be present. Next the sample holder containing dry sample was kept in the dessicator for 2 hours. Further the sample holder containing dry sample was used for characterization using STM. Different images were obtained which confirmed the presence of AgNPs.

**Characterization by scanning tunneling microscope**

The synthesized silver nanoparticles were characterized using scanning tunneling microscope (Nano Surf Easy Scan 2). Usually scanning tunneling microscope (STM) is used to characterize the superficial properties of nanoparticles. A thin layer of AgNPs solution was placed on sample holder. The care was taken that bubbles should not be present. Next the sample holder was kept in the dessicator for two hours. Further the sample holder containing dry sample was used for characterization using STM. Different images were obtained which confirmed the presence of AgNPs.

**Evaluation of antibacterial activity**

Silver nanoparticles biosynthesized from lemon juice, orange juice and sweet lime juice were tested individually against test organisms for antibacterial activity by agar well diffusion method [15]. For this study both Gram positive (Staphylococcus aureus ATCC 6538P) and Gram negative (Escherichia coli ATCC 8739, Salmonella typhimurium ATCC 23564 and Klebsiella pneumonia ATCC 10031) organisms were used. This was performed by determining ZOI (zone of inhibition), which is rapid and inexpensive to determine the susceptibility of a particular test organism to antimicrobial agent. This was executed by measuring the zone of inhibition using a vernier caliper. Pure cultures were subcultured into nutrient broth and incubated at 37°C for 24-48 hours. Three wells of 5 mm diameter were made on pre-incubated nutrient agar plates using gel puncher. Each test organism (10⁶ cfu/mL) was spread uniformly onto the individual plates using spread plate technique. Using sterile micropipette tips, 0.1 mL (100 µL) of the sample of silver nanoparticle solution was pipetted into each of the wells in all the plates. After incubation, the diameters of zone of inhibition were measured in triplicate.

**Results and Discussion**

**Biosynthesis of silver nanoparticles**

It is well known that silver nanoparticles exhibit a yellowish-brown color in aqueous solution due to excitation of surface plasmon vibrations of silver nanoparticles [16]. Reduction of silver ions to silver nanoparticles could be followed by a color change and UV-Vis spectra. The technique outlined above has proven to be very useful for the analysis of nanoparticles [17]. Therefore, the progress in conversion reaction of silver ions to silver nanoparticles was followed by a color change and spectroscopic techniques. The photograph of sample solutions containing silver nitrate and silver nitrate in the presence of optimal amounts of fruit juices after completion of the reaction shows appearance of a yellowish-brown color which confirms the existence of silver nanoparticles (photo not shown). Several approaches have been employed to obtain better synthesis of silver nanoparticles. Recently, synthesis of silver nanoparticles using plant extracts getting more attraction [18,19]. Citrus fruits like orange and lemon are rich source of antioxidants. The compounds present in these fruit juices possess a broad spectrum of chemical and biological activities including radical scavenging properties [20]. Flavonoids present have been reported to exhibit antioxidant, anti-carcinogenic, antihypertensive and antimicrobial properties. The orange juice contains oxido-reductively labile ascorbic acid and citric acid. These two compounds along with polyphenols might have played a pivotal role in nano-conversion. In a recent study, the function of citrate ions in the synthesis of silver nanoparticles through a synergistic reduction approach in ambient conditions was explored and it was found that the citrate ions can play multiple roles in the synthesis process including a reducing agent, a stabilizer, and a complex agent and they show some unique features under the reported conditions [21]. The silver nanoparticles were characterized by UV-Vis spectroscopy. The absorption spectra (Figures 1-3) of silver nanoparticle solution prepared with the proposed method showed a surface plasmon absorption band with a maximum of 420-430 nm, indicating the presence of spherical Ag nanoparticles. These structural details were confirmed by SEM images. Scanning electron microscopy provided further insight into the morphology and size details of the silver nanoparticles. Comparison of experimental results showed that the diameter of prepared nanoparticles in the solution was about 45-50 nm. Figures 4-6 show scanning tunneling microscopic images of silver nanoparticles biosynthesized from lime juice, sweet lime juice and orange juice respectively. Similarly Figures 7-9 shows the scanning electron micrograph of silver nanoparticles obtained from orange juice, lime juice and sweet lime juice respectively. Chemical synthesis methods may lead to the presence of some toxic chemical species adsorbed on the surface that may have adverse effects in its
Bactericidal activity of silver nanoparticles

After 24 hours of incubation, the inhibitory effect of AgNPs from orange juice was significant as compared to the other two. Zone of inhibition (ZoI) was used as a measure for comparing bactericidal activity of these AgNPs. AgNPs from orange showed about 8 mm zone against the test organisms: *E. coli* and *K. pneumoniae*. Similarly, the AgNPs from orange showed 5 mm and 6 mm ZoI against test organisms.
organisms: S. aureus and S. typhimurium respectively. (Table 1 and Figure 10-12). Table 1 gives the ZoI of the formed silver nanoparticles against pathogens E. coli, K. pneumoniae, S. aureus and S. typhimurium.

Figure 6: Scanning Tunneling Microscopic image of silver nanoparticles biosynthesized from sweet-lime juice.

Figure 7: Scanning electron micrograph of silver nanoparticles biosynthesized from lime juice.

Figure 8: Scanning electron micrograph of silver nanoparticles biosynthesized from orange juice.

Figure 9: Scanning electron micrograph of silver nanoparticles biosynthesized from sweet-lime juice.

Figure 10: Comparison of antibacterial activity of silver nanoparticles biosynthesized from lime juice against A) Escherichia coli, B) Klebsiella pneumoniae, C) Staphylococcus aureus and D) Salmonella typhimurium.

Figure 11: Comparison of antibacterial activity of silver nanoparticles biosynthesized from orange juice against A) Escherichia coli, B) Klebsiella pneumoniae, C) Staphylococcus aureus and D) Salmonella typhimurium.
However, such aggregation will lead to loss of properties associated of nanoparticles [26]. In our study also, we observed such aggregation authors that, smaller the particle sizes more will be the cytotoxic effect cytotoxic nature of nanoparticles. It was reported earlier by many agglomeration, particle size also plays important role in defining the is an important parameter for toxicological studies. Apart from nanoparticles application is their tendency to aggregate because of small size and large surface area. The degree of agglomeration is an important parameter for toxicological studies. Apart from agglomeration, particle size also plays important role in defining the cytotoxic nature of nanoparticles. It was reported earlier by many authors that, smaller the particle sizes more will be the cytotoxic effect of nanoparticles [26]. In our study also, we observed such aggregation of nanoparticles if stored in the aqueous form for long time period. However, such aggregation will lead to loss of properties associated with nano-scale nature of nanoparticles. The present study has not involved any cytotoxic assessment of the nanoparticles.

Conclusion

The present approach of biosynthesis of silver nanoparticles used fruit juices as reducing and capping agents. The method is cost effective and environmental friendly. The formed silver nanoparticles were well characterized by UV-Vis spectra, scanning tunneling micrographs and scanning electron micrographs. The antimicrobial activity of these nanoparticles was well demonstrated against both gram positive and gram negative bacteria. The study also confirms that gram positive bacteria are relatively resistant to the bactericidal action than gram negative bacteria. Silver nanoparticles biosynthesized using orange juice were more effective in their antibacterial activity than those from other fruit juices. The approach of use of fruit juices for biosynthesis of nanoparticles is novel and can be adapted to biosynthetic approaches of other metal nanoparticles.

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Table 1: Zone of inhibition (mm) of silver nanoparticles biosynthesized from fruit juices against test microorganisms.

| Microorganism | Lime juice | Orange juice | Sweet-lime juice |
|---------------|------------|--------------|-----------------|
| *Escherichia coli* | 7 | 8 | 6 |
| *Klebsiella pneumoniae* | 6 | 8 | 5 |
| *Staphylococcus aureus* | 5 | 5 | 4 |
| *Salmonella typhimurium* | 6 | 6 | 4 |

Figure 12: Comparison of antibacterial activity of silver nanoparticles biosynthesized from sweet-lime juice against A) *Escherichia coli*, B) *Klebsiella pneumoniae*, C) *Staphylococcus aureus* and D) *Salmonella typhimurium*.
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