Original Article

Synthesis and Characterization of Silver Nanoparticles for Antibacterial Application against Bacillus Subtilis and Pseudomonas Aeruginosa

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Abstract: In this study, the stable silver nanoparticles (AgNPs) were synthesized by reducing silver nitrate (AgNO3) using trisodium citrate (TSC). The product was characterized by Ultraviolet-Visible spectroscopy, Fourier-transform infrared spectroscopy (FTIR), transmission electron microscopy (TEM) and X-ray diffraction analysis (XRD). UV-Vis spectrum showed a peak around 420 nm. TEM analysis revealed the homogeneity in the size of AgNPs (35-45 nm), well-dispersed quasi-spherical in water. The prepared AgNPs exhibited high antibacterial activity against Bacillus subtilis and Pseudomonas aeruginosa bacteria. The average zones of inhibition were 20 mm and 17 mm for Pseudomonas aeruginosa and Bacillus subtilis bacteria, respectively. The inhibition zone of AgNPs was also compared to the reference antibiotics drugs such as ampicillin and natamycin. This research exhibits an efficient and eco-friendly synthesis of silver nanoparticles with potent antimicrobial and antibacterial performance.

Keywords: Silver nanoparticles, chemical synthesis, antibacterial activity, antimicrobial agent, cell inhibition.
1. Introduction

The development of new effective antimicrobial reagents based on metal nanomaterials has replaced the conventional antibiotics due to emergence of resistant microbes. Silver nanoparticles (AgNPs) have long been known to have strong antibacterial effects as well as antimicrobial activities \([1, 2]\) and could be considered as a potential alternative for existing drugs. AgNPs have also wide applications in the biomedical and environmental fields such as biological labeling, drug delivery, cancer therapy, molecular imaging, diagnostics, textile, catalysis, photography, electronics and food industry \([3-6]\). In one of our previous works \([7]\), the antibacterial activity against Escherichia coli and Staphylococcus aureus of silver nanoparticles was reported, where the colloidal silver solution was successfully synthesized in a wet chemical process using silver nitrate (AgNO\(_3\)), sodium borohydride (NaBH\(_4\)) and starch as starting agents. It has been proven that AgNPs exhibited an excellent antibacterial activity against Gram-negative bacteria Escherichia coli (E. coli) and Gram-positive Staphylococcus aureus. It is also reported that the average inhibition zone of AgNPs was 7.7 mm for Escherichia coli and 7 mm for Staphylococcus aureus bacteria. In this study, AgNPs were synthesized from silver nitrate (AgNO\(_3\)) and trisodium citrate dihydrate (TSC) and the average inhibition zones of AgNPs were found to be 17 mm for Bacillus subtilis and 20 mm for Pseudomonas aeruginosa bacteria. Several proposals have been developed to explain the inhibitory effects of AgNPs on bacteria. It is believed that reaction of heavy metal with proteins leads to the inactivation of the proteins in bacteria \([8]\) and interaction between silver atoms with the sulfur-containing proteins plays an essential role in cellular damage \([9]\). It is revealed that the silver atoms can destroy microorganisms completely through oxidative catalysis mechanism \([10]\). Experimental results may provide useful information for understanding the antibacterial effects and inhibitive process of silver atoms although the mechanism of antimicrobial effects of silver atoms on bacteria is complicated. It is noticeable that AgNPs are non-toxic to animal cells but highly toxic to bacteria, so they are very promising antibacterial systems.

2. Materials and Methods

Silver nitrate (AgNO\(_3\), 99.8%), trisodium citrate dihydrate (TSC, Na\(_3\)C\(_6\)H\(_5\)O\(_7\).2H\(_2\)O, 99%) supplied by Sigma-Aldrich were of analytical grade and used as received. Deionized water prepared with a Milli-Q water purification system was used to prepare the solutions and to clean all glassware and experimental tools. The synthesis process of colloidal silver was carried out as follows. Firstly, AgNO\(_3\) was dissolved in the deionized water to obtain 30 ml of 1 mM AgNO\(_3\) solution. Then, it was heated until boiling. After that, 5 ml of 30 mM TSC solution was added drop-wise to the AgNO\(_3\) solution under vigorous stirring. The molar ratio of TSC/AgNO\(_3\) was 5:1, the volume ratio of TSC/AgNO\(_3\) was 1:6 and injection time was 6 min. The reaction was maintained at the boiling status for 14 min in the dark to avoid direct exposure to the natural light. After the reaction was completed, the stock solution was cooled down to the room temperature. The chemical reaction was the reduction of silver ions by TSC:

\[
4\text{AgNO}_3 + \text{Na}_3\text{C}_6\text{H}_5\text{O}_7 + 2\text{H}_2\text{O} \rightarrow 4\text{Ag} \downarrow + \text{H}_2\text{C}_6\text{H}_5\text{O}_7 + 3\text{NaNO}_3 + \text{H}_2\text{O} \uparrow
\]

The change of the color of the reaction solution from transparent to pale yellow was an indicator for the reduction of the Ag\(^+\) ions into metallic silver. UV-Vis spectrophotometer (HITACHI-U2900, Japan) was used to characterize the structure of silver nanoparticles (wavelength of 250-600 nm). Transmission electron microscopy (TEM, JEM-1010 JEOL) with an electron kinetic energy of 80 keV was used to investigate the morphology and size of the products. The size-distribution profile of AgNPs was studied by using dynamic light scattering (DLS) measurements conducted with a particle size analyzer (PSA, Delta Nano C, Beckman). The Fourier-Transform Infrared (FTIR) spectra recorded on the Spectrum
FTIR Affinity-1S (SHIMADZU) in the range of 500-4000 cm\(^{-1}\) at a scan speed of 16 cm/s was used to detect the functional groups on the AgNPs. X-ray diffraction pattern was performed using an X-ray diffractometer (Siemens D5005) with the wavelength of Cu-K\(_\alpha\) radiation of \(\lambda = 1.5417\ \text{Å}\) to analyze the phase structure and exact material identification of samples. Bacillus subtilis and Pseudomonas aeruginosa bacteria were selected as the test micro-organisms to evaluate the antibacterial activity of AgNPs. All cultures were procured from Thai Nguyen University of Sciences (Vietnam) and were maintained at 4°C. A sterile agar plate was perforated to form a well with the diameter of 6 mm. The antibacterial performance of AgNPs was checked by agar well diffusion assay. 15 ml of nutrient medium was poured onto the plate then allowed to solidify for 5 min. After that, 100 µl of inoculum suspension containing bacteria was spread uniformly on the plate and allowed to dry for 5 min. Finally, 70 µl of AgNPs colloidal solution was loaded into the well on the plate. After incubating at 37°C for 24 h, the diameter of inhibition zone was measured and recorded as a mean value of the triplicate experiment.

3. Results and Discussion

The growth mechanism of AgNPs obtained by the oxidation-reduction reaction between sodium citrate and silver nitrate was described by the diagram in Figure 1a. The metallic silver (Ag\(^0\)) was first formed from the Ag\(^+\) reduction by citrate. The Ag atoms aggregated to form silver cluster and the initial seeds began to grow via the Ostwald ripening, in which the larger particles grew at the expense of the smaller ones. The UV-Vis absorption spectrum of the synthesized AgNPs is shown in Figure 1b. It can be seen that the plasmon absorption band is located at 420 nm with the value of the absorbance intensity of about 2.83 (arbitrary unit) corresponding to the typical characteristic surface plasmon resonance (SPR) of the spherical AgNPs. In addition, the spectrum has only one peak, indicating that the particles are mainly spherical with small size and regular distribution. This result is in good agreement with previous findings, that the number of surface plasmon resonance (SPR) peaks increases as the symmetry of particles decreases [11]. The full width at half maximum (FWHM) is of 90 nm. The SPR peak is narrow showing the reductive process of Ag\(^+\) ions was completed and obtained AgNPs were uniform.

TEM image of the AgNPs and their size distribution is shown in Figure 2a and Figure 2b, respectively. It is clearly revealed that AgNPs have a uniform spherical shape of about 35-45 nm, the average size is 40 nm in diameter. Figure 3 shows the color of the colloidal AgNPs solution at different reaction times. The color changed from translucent to darker when the reaction time increased.

Figure 1. (a) Growth mechanism, (b) UV-Vis absorption spectrum of the AgNPs.
Figure 2. (a) TEM image, (b) The size distribution of AgNPs.

Figure 3. The change of the AgNPs solution color according to reaction time.

Figure 4. (a) The UV-Vis absorbance spectra of AgNPs prepared for different reaction time, (b) Dependence of absorption peak intensity on the reaction time.

The UV-Vis absorption spectra of the samples synthesized for different reacting times are shown in Figure 4a. The intensity of the surface plasmon band increases quasi linearly with respect to the reaction time in the first 14 min and remains unchanged when the reaction time is prolonged to 42 min (Figure 4b). All SPR peaks typically located between 420-430 nm, characteristic of silver nanoparticles [12]. This observation shows that the nucleation and growth reaction were completed after 14 min.

Figure 5a displays the XRD pattern of the synthesized AgNPs. Four characteristic peaks were observed at 37.68°, 44.56°, 65.07° and 76.62° corresponding to the diffraction of (111), (200), (220) and (311) crystalline planes, respectively, which can be readily indexed to a face-centered cubic (FCC) lattice structure of silver (JCPDS No.04-0783) [13]. Figure 5b depicts the FTIR spectrum of AgNPs. The strong absorption peaks at 1590, 1398 and 1372 cm\(^{-1}\) wavenumbers represent the presence of NO\(_2\) group coming from the AgNO\(_3\) solution. The peaks at 3413 and 2926 cm\(^{-1}\) associate with the OH functional group. The peak at 1058 cm\(^{-1}\) was due to stretching of C-OH binding [2].

It is known that if AgNPs are unstable and aggregated, the plasmon peak intensity at 420 nm will decrease. So the UV-Vis spectroscopy can be used for monitoring the stability of AgNPs and the result is shown in Figure 6. It can be seen that the plasmon peak intensity slightly decreased after more than 40 days, which indicated the stability of the sample over time. Therefore, the sample could be kept and used for antibacterial activity for a long time.
It is known that the antibacterial effect of AgNPs depends on several parameters such as the size, shape and the surface charge of the particles. AgNPs can be attached to the surface of the bacterial cell membrane by interacting with the sulfur-containing proteins, disturbing permeability and respiration function of the cell, resulting in cell death [14]. AgNPs can also penetrate the cell wall into the bacteria, causing the deterioration of the membrane and cell destruction [15]. AgNPs may interact with the thiol group compound in the respiratory enzyme of bacterial cell, thus inhibiting the respiration process in the bacteria [14]. AgNPs can condense the DNA and prevent the replication of the DNA as well as the reproduction of the cells [16]. The silver ions released from AgNPs may penetrate inside the cell and interact with phosphorus in protein and DNA of bacteria, resulting in the loss of the cell viability, and ultimately damaging the cell.
mm). Similarly, the inhibition zone of Bacillus subtilis was found to be wider than that of Staphylococcus aureus (12 mm) [7]. The inhibition zone of AgNPs was also compared against the reference antibiotics drugs containing ampicillin. The results show that the antibacterial activity for Bacillus subtilis of the 70 µl of colloidal AgNPs solution was approximately equal to 90% that of 0.875 mg ampicillin antibiotics, and the antibacterial activity for Pseudomonas aeruginosa of AgNPs and ampicillin was almost the same.

4. Conclusions

We have used a method of reduction Ag⁺ to Ag⁰ by TSC for synthesizing uniform spherical AgNPs with average diameter of 40 nm in size and stable for long term (over 40 h). The obtained results show that the synthesized AgNPs exhibited an excellent antimicrobial activity against the microorganism disease, such as Bacillus subtilis and Pseudomonas aeruginosa, known as multidrug resistant and hazardous bacteria. The average zones of inhibition for Bacillus subtilis and Pseudomonas aeruginosa were of 17 mm and 20 mm, respectively. Such AgNPs constitute a potential candidate for antibacterial applications.

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