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Green Synthesis of Selenium Nanoparticles using *Allium Sativum* Extract and Its Antimicrobial Activity Against Gram Negative Bacteria

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Abstract

Green synthesis of nanoparticles by using the plant extract has an increased benefit because it is eco-friendly, safe strategic and cost effective, even in the industry it produces much less toxic waste. In the present study we investigated the extract of *Allium sativum* which is used for the synthesis of biogenic selenium nanoparticles. The bio reduction of selenium nanoparticles was observed due to colour change of the sodium selenite solution and the studied selenium nanoparticles were characterized by UV-Visible spectrophotometer, Transmission electron microscopy (TEM), Fourier transform spectroscopy (FT-IR) and Energy dispersive X-Ray spectroscopy (EDAX). The selenium nanoparticles in size ranging 7-48nm which found more stable more than two months. The green synthesize selenium nanoparticles significantly inhibited the growth of *Escherichia coli* and *Salmonella typhi*.

Keywords

Allium sativum, Antimicrobial activity, Green synthesis, Selenium nanoparticles

INTRODUCTION

Nanotechnology is mainly depends on the arrangement of materials at the atomic stage to achieve nanoscale matter with unique characteristics [1]. The nanoparticles synthesis is broadly classified in different physical and chemical procedure but this

technique are costly and harmful for environment [2]. Biosynthetic methods are simple for synthesis of eco-friendly, cost effective nanoparticles using microbes, algae and plant materials etc.

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Biosynthesis of metal nanoparticles using plant extract is the most cherished green method of production of nanoparticles and it is exploited to a vast extract because the plants are widely distributed, easily available, safe to handle and with a range of metabolites [3].

Selenium has great potential and exclusive properties in the field of physics, chemistry and biology [4]. The selenium at nano size acts as a potential chemo-preventive agent with reduced toxicity [5]. Selenium a metalloid chalcogens, attracts more attention because of its special physical properties such as the anisotropy of conductivity and catalytic activity [6]. Selenium nanoparticles has been synthesized by different approaches like *Bacillus sp. Mshl* [7], *Klebsiella pneumonia* [8], *Aspergillus terus* [9], *Saccharomyas cerevisiae* [10], *Bougainvillea spectabilis* [11], leaves of lemon [12], rasin extract of grapes [13].

The selenium is most important because of selenium deficiency can lead to heart disease, hypothyroidism and a weakened immune system [14]. Kashin-beck disease is also a result of selenium and iodine deficiency [15]. This disease affects bones and joints of growing children. One symptom is enlargement of cracking of small joints while a more serious symptom is distorted growth of long bones that leads to shorter structure. Recent supplementation of salt with selenium has reduced the occurrence of the disease.

Human beings are often infected by microorganisms such as bacteria, molds, yeast and viruses present in their surroundings. Gram negative *Escherichia coli* and *Salmonella typhi* were widely used to bacterial experiments. These bacteria mostly present on the body surface of mammals and also present in their surroundings but sometimes it occurs infection to them, so therefore *Escherichia coli* and *Salmonella typhi* strain were selected for the antibacterial activity by using selenium nanoparticles synthesized by *Allium sativum* extract in this study by well diffusion method. The selenium nanoparticles showing antimicrobial activity by zone of inhibition and it's increased with increasing concentrations.

MATERIALS AND METHODS

Preparation of Allium sativum extract

Buds of *Allium sativum* 10gm were collected in a clean mortar. Buds were crushed using motor pestle and sufficiently diluted with water to make a thick paste. This paste was filtered through Whatman filter paper. The resulting pest was stored in refrigerator and used for further experiments.

Synthesis of metal nanoparticles

Flask containing 25 ml 5 mM Na2SeO3 solutions was kept on magnetic stirrer. Then drop wise addition of *Allium sativum* extract was made in flask containing Na2SeO3 solution until color of sodium selenite solution changed. From this solution 5 ml was taken which was used as a control. Remaining 20 ml solution was kept in shaker in dark for 72 hrs. After few days the color change of the solution was observed.

UV-Vis spectra analysis

The reduction of metallic selenium ions was observed by measuring the UV-V is spectrum after 10 to 15 min of color change. A small aliquot was drawn from the solution and a wavelength from 250nm to 700nm on UV-Vis spectrophotometer (Optizon Double beam 3220).

TEM analysis

Transmission Electron Microscopic (TEM) analysis was performed with Techni 20 (Philips, Holland). A thin film of the sample was prepared on a carbon coated copper grid by dropping a very small amount of the sample on the grid. The Allium sativum extract containing Se nanoparticles were subjected to centrifugation at 13000 rpm for 10 min. The pellet thus recovered was subjected to washing by its resuspension in de-ionized water followed by centrifugation at 13000 rpm for 10 min, to remove possible organic contamination present in nanoparticles. Finally, pellet was freeze dried using a lyophilizer (Labconco, Kanas, USA).

EDAX analysis

EDAX analysis was carried out on EDAX XL-30 operating at 15-25KeV. Incorporation of selenium nanoparticles in gauze cloth. Nanoparticles suspension was poured on the gauze cloth discs (diameter 1cm) and there discs were dried at $36^{\circ}c$ for 7 days.

Sample preparation for Fourier Transform Spectroscopy (FTIR)

Metal containing *Allium sativum* extract for Fourier Transform Infrared (FT-IR) analysis was prepared by mixing 5 mg metal salt in 10 ml *Allium sativum* extract. This metal containing *Allium sativum* extract was incubated at room temperature for 1 hour. After 1 hour incubation, this metal containing leaf extract was dried in Petri plate. After drying, particles were scraped using blade. So, powder of synthesized nanoparticles was obtained. Then spectral scan analysis was carried out at wave number ranging from 400-4000 cm⁻¹ by using a FT-IR spectrometer (Perkin Elmer, Spectrum GX) with resolution of 0.15

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cm⁻¹ to evaluate functional groups that might be involved in sorption process.

Antibacterial studies

The selenium nanoparticles synthesized from *Allium* sativum extract was tested for their antimicrobial activity by well diffusion method against pathogenic organisms like *Escherichia coli* and *Salmonella typhi*. Each strain was spread uniformly on the individual plates using glass spreader. Well of size 6mm have been made on Muller Hilton agar plates using gel puncture. Using micropipette 25µl, 50µl, 75µl and 100µl of the sample of nanoparticles solution and also Ampicillin as control were poured into wells.

After incubation at 37°c for 24 hrs the different level of zone of inhibition measure.

RESULTS AND DISCUSSION Visual observation

Reduction of metal salts into metal nanoparticles by the bio-molecules is always accompanied by the colour change of reaction medium. In the present study the colourless solution of sodium selenite is changed in light pink colour after drop wise addition of *Allium sativum* extract at zero hour. As the reduction proceed, the colour of reaction medium is gradually changed to dark pink colour after 24 hours.



Fig. 1: Color change of reaction medium, the color of reaction medium is gradually changed to dark pink color after 24 hours.

UV – Visible Spectroscopy

In order to determine the formation of Selenium nanoparticles in the extract of *Allium sativum*, a

spectral scanning procedure was carried out from 250 nm to 700 nm. Colloidal solution exhibited absorption maxima at 400 nm (Fig. 2).

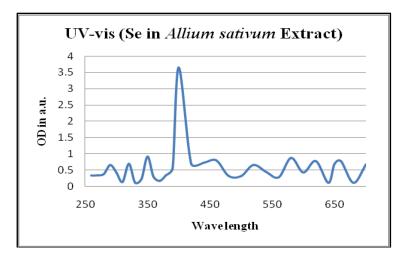


Fig. 2: UV- VIS spectra selenium nanoparticles synthesize by Allium sativum extract and gets peak at 400nm.



Initially the colloidal solution appeared white in color but after incubation of a period of 24 hours, it turned to reddish brown in color. Building of absorbing maximum at 400 nm clearly indicates the gradual formation of particles during the incubation period. **Transmission Electron Microscopy (TEM)** The results obtained from the transmission electron microscopy (TEM) and selected area electron diffraction (SEAD) study gives clear indication about the shape, size and size distribution of the nanoparticles.

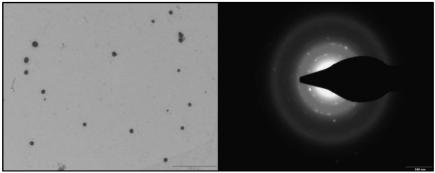


Fig. 3: TEM image (left) and SEAD pattern (right) of Selenium nanoparticles synthesized using extract of *Allium sativum*.

TEM analysis of colloidal solution indicated the formation of selenium nanoparticles. (Figure 3) shows that size of particles, generated using *Allium sativum* extract ranges from 8 – 52 nm. Formation of variable size of particles indicates that particles suggest that *Allium sativum* extract could form polydisperse nanoparticles. Figure 3 shows Selected Area Electron Diffraction (SAED) of selenium nanoparticles results shows that particles are crystalline in nature as diffraction ring appeared which correspond to diffraction angle of (111, 121 and 311).

Fourier Transform Infrared Spectroscopy (FT-IR)

FT-IR analysis was carried out to identify the possible bio molecules and plant extract-metal ions

formation interaction responsible for and stabilization of selenium nanoparticles. The result of FT-IR analysis of Allium sativum extract is presented in figure 4. The figure 4 shows the spectrum of both the sample control (A) and test (B). The fig. 4 (B) shows the spectrum of the sample that contains selenium metal in Allium sativum extract or fig. 4 (A) shows the spectrum of the Allium sativum extract that did not contain metal selenium. Spectra B show the peaks of both control and test, similarly the Fig. 4 (A) is showing transmission peaks of the control sample. Around 600 and 500 may be due to the partial duitriation of amine or carboxyl group.

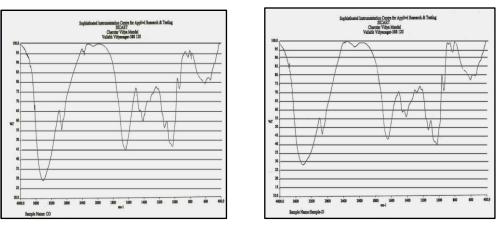


Fig. 4: FTIR spectrum of (a) Allium sativum extract and (b) Selenium nanoparticles synthesized by Allium sativum extract

(a)

(b)



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Two absorption peaks located around 3400 and 4000 can be assigned as the absorption peak of N-H. The peaks located around 3000 and 3200 may be due to the presence of C-H group. The absorption peaks around 2300 and 2000 can be assigned as the peaks of CO2. The absorption peaks around 1500 and 1800 can be assigned as the absorption peaks of C=O / C=N / C=C. The peaks around 1200 and 1100 were attributed to the stretching vibration of carboxyl group (C=O). The peaks around 1100 and 1000 may be due to the presence of C-O group. Two absorption

peaks around 600 and 500 may be due to the partial deuteration of amine or carboxyl group.

Energy Dispersive X-Ray Spectroscopy (EDAX)

EDAX analysis gives qualitative as well as quantitative status of elements that may be involved in formation of nanoparticles. Figure shows the elemental profile of synthesized nanoparticles using *Allium sativum* extract. The analysis revealed the highest proportion of Selenium (55%) in nanoparticles followed by oxygen (15%), sodium (28%) etc.

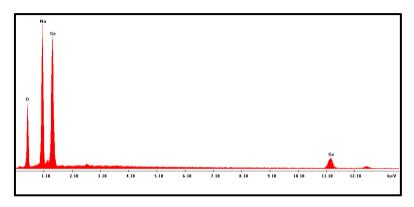
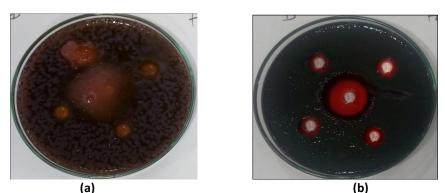


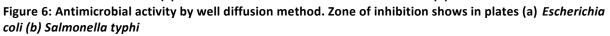
Fig. 5: EDAX spectrum of selenium nanoparticles synthesize using extract of Allium sativum

Antimicrobial activity by well diffusion method

It is well known that selenium ions and nanoparticles are highly toxic to microorganisms. Selenium nanoparticles have been known to have inhibitory and bacterial effect and thus we extend its application as an antibacterial agent. The antimicrobial activity is estimated by the zone of inhibition. Several studies propose that selenium may attach to the surface of the cell membrane disturbing permeability and respiratory function of the cell. It is also possible that selenium nanoparticles not only interact with the surface of membrane but can also penetrate inside the bacteria.

The antimicrobial activity of selenium nanoparticles synthesized by *Allium sativum* extract was investigated against pathogenic organisms such as *Escherichia coli* and *Salmonella typhi* using well diffusion method.







It may be observed that increasing quantity of selenium nanoparticles have comparatively higher antimicrobial activity against *Escherichia coli* and

Salmonella typhi probably due to thinner peptidoglycan layer and presence of porins.

Table 1: Zone of inhibition of Selenium nanoparticles synthesized by Allium sativum extra	ts against:
pathogenic bacteria	

Sample	Escherichia	Salmonella
	coli	typhi
Zone of inhibition (mm)		
25µl	11	13
50µl	16	18
75µl	21	23
100µl	29	27

The susceptibility of pathogenic bacteria to selenium nanoparticles shown in Table 1.

The diameter of inhibition zone (mm) around each well with selenium nanoparticles synthesized by *Allium sativum* were found to have highest antimicrobial activity against *Escherichia coli* (29mm) and *Salmonella typhi* (27mm) at 100µl and the lesser antimicrobial activity of selenium nanoparticles synthesized by *Allium sativum* extract was found at 25µl *Escherichia coli* (11mm) and *Salmonella typhi* (13mm).

CONCLUSIONS

The present study was carried out to synthesis of Selenium nanoparticles using extract of *Allium sativum*. The bio molecules of *Allium sativum* extract acted as stabilizing as well as capping agent leading to the formation of Selenium nanoparticles. UV-Vis Spectra at 400nm with *Allium sativum* extract and observed as hollow and spherical particles in size ranging 7-45nm which is found more stable more than two months. EDAX analysis was carried out to check the presoak of Selenium in nanoparticles. Results of EDAX, confirmed its present. TEM and

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SEAD represented addition evidence of formation of nanoparticles whereas SEAD indicates the particles were crystalline in nature. The zones of inhibition were formed in the antimicrobial screening test indicated, that the selenium nanoparticles synthesized by *Allium sativum* extract in this process has the efficient antimicrobial activity against pathogenic bacteria. The biologically synthesized selenium nanoparticles could be of immense use in medical field for their efficient antimicrobial function.

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