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GREEN SYNTHESIS OF SILVER NANOPARTICLES USING *ARGEMONE MEXICANA* LEAF EXTRACT AND EVALUATION OF THEIR ANTIMICROBIAL ACTIVITIES

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There is an increasing commercial demand for nanoparticles due to their wide applicability in various areas such as electronics, catalysis, chemistry, energy, and medicine. Metallic nanoparticles are traditionally synthesized by wet chemical techniques, where the chemicals used are quite often toxic and flammable. In this work, we describe a cost effective and environment friendly technique for green synthesis of silver nanoparticles from 5mM $AgNO_3$ solution through the extract of *Argemone maxicana* leaf extract as reducing agent as well as capping agent. Nanoparticles were characterized using UV–Vis absorption spectroscopy, FTIR, XRD and SEM. X-ray diffraction and SEM analysis showed the average particle size of 30 nm as well as revealed their structure. Further these biologically synthesized nanoparticles were found to be highly toxic against different bacterial spp. The most important outcome of this work will be the development of value-added products from *Argemone maxicana* (a potential weed of India) for biomedical and nanotechnology based industries. This is for the first time that *Argemone maxicana* weed leaf extract was used for the synthesis of nanoparticles.

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1. Introduction

The field of nanotechnology is one of the most active areas of research in modern material sciences. Nanoparticles exhibit completely new or improved properties based on specific characteristics such as size, distribution and morphology. Nanotechnology is a field that is burgeoning day by day, making an impact in all spheres of human life. New applications of nanoparticles and nanomaterials are emerging rapidly [1, 2, 3]. Nanocrystalline silver particles have found tremendous applications in the field of high sensitivity biomolecular detection and diagnostics4, antimicrobials and therapeutics [5,6], Catalysis [7] and micro-electronics8. However, there is still need for economic, commercially viable as well environmentally clean synthesis route to synthesize silver nanoparticles.

A number of approaches are available for the synthesis of silver nanoparticles for example, reduction in solutions9, chemical and photochemical reactions in reverse micelles10, thermal decomposition of silver compounds [11], radiation assisted [12], electrochemical [13], sonochemical14, microwave assisted process [15] and recently via green chemistry route [16,17,18].

Biological methods of synthesis have paved way for the "greener synthesis" of nanoparticles and these have proven to be better methods due to slower kinetics, they offer better manipulation and control over crystal growth and their stabilization. This has motivated an upsurge in research on the synthesis routes that allow better control of shape and size for various nanotechnological applications. The use of environmentally benign materials like plant extract19, bacteria20, fungi21 and enzymes [22] for the synthesis of silver nanoparticles offer numerous

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benefits of eco-friendliness and compatibility for pharmaceutical and other biomedical applications as they do not use toxic chemicals for the synthesis protocol. Chemical synthesis methods lead to presence of some toxic chemical absorbed on the surface that may have adverse effect in the medical applications. Green synthesis provides advancement over chemical and physical method as it is cost effective, environment friendly, easily scaled up for large scale synthesis and in this method there is no need to use high pressure, energy, temperature and toxic chemicals.

Silver has long been recognized as having inhibitory effect on microbes present in medical and industrial process [23, 24]. The most important application of silver and silver nanoparticles is in medical industry such as topical ointments to prevent infection against burn and open wounds [25].

Here in, we report for the first time synthesis of silver nanoparticles, reducing the silver ions present in the solution of silver nitrate by the aqueous extract of Argemone leaf. Further these biologically synthesized nanoparticles were found highly toxic against different pathogenic bacteria and fungi.

2. Materials and method

2.1 Plant material and preparation of the extract

Green Argemone leaves were used to make the aqueous extract. Argemone leaves weighing 5g were thoroughly washed in distilled water, cut into fine pieces and were boiled into 100 ml sterile distilled water and filtered through Whatman No.1 filter paper (pore size 25 µm).

2.2 Synthesis of silver nanoparticles

5mM aqueous solution of Silver nitrate ($AgNO_3$) was prepared and used for the synthesis of silver nanoparticles. 10 ml of Argemone leaf extract was added into 90 ml of aqueous solution of 5 mM Silver nitrate for reduction into Ag^+ ions and kept at room temperature for 4 hours.

2.3 UV-Vis Spectra analysis

The reduction of pure Ag^+ ions was monitored by measuring the UV-Vis spectrum of the reaction medium at 3 hours after diluting a small aliquot of the sample into distilled water. UV-Vis spectral analysis was done by using UV-Vis spectrophotometer.

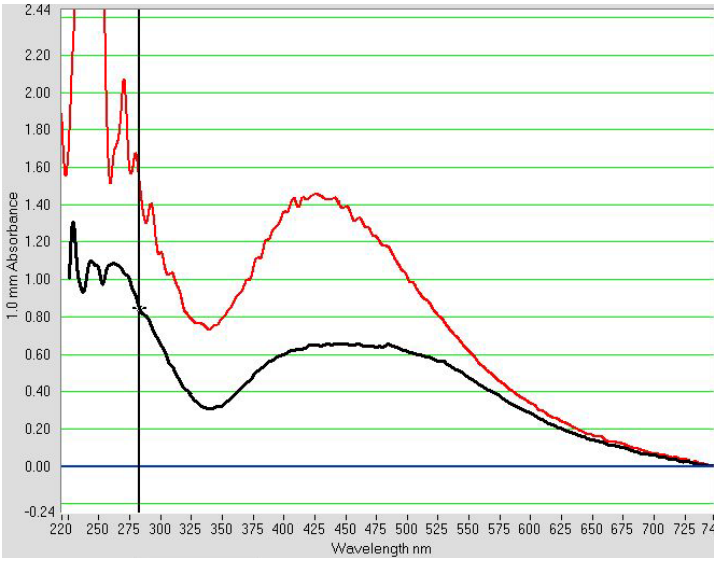


Fig. 1. UV-Vis absorption spectra of silver nanoparticles synthesized by Argemone leaf extract after 2 hrs (Black) and 4 hrs (Red).

2.4 FTIR analysis of dried biomass after bioreduction

To remove any free biomass residue or compound that is not the capping ligand of the nanoparticles, the residual solution of 30 ml after reaction was centrifuged at 10000 rpm for 10 min and the resulting suspension was redispersed in 2 ml sterile distilled water. The centrifuging and redispersing process was repeated three times. Thereafter, the purified suspension was freeze dried to obtain dried powder. Finally, the dried nanoparticles were analyzed by FTIR Nicolet Avatar 660 (Nicolet, USA).

2.5. XRD measurement

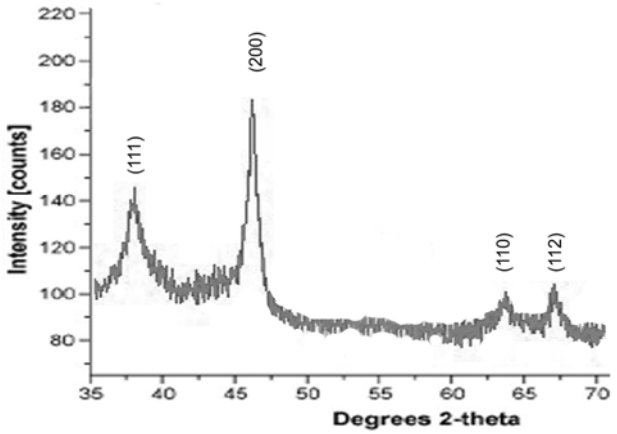
The silver nanoparticle solution thus obtained was purified by repeated centrifugation at 5000 rpm for 20 min followed by redispersion of the pellet of silver nanoparticles into 10 ml of deionized water. After freeze drying of the purified silver particles, the structure and composition were analyzed by XRD and SEM. The dried mixture of silver nanoparticles was collected for the determination of the formation of Ag nanoparticles by an X'Pert Pro x-ray diffractometer (PAN analytical BV, The Netherlands) operated at a voltage of 40 kV and a current of 30 mA with Cu Kα radiation in a θ- 2 θ configuration. The crystallite domain size was calculated from the width of the XRD peaks, assuming that they are free from non-uniform strains, using the Scherrer formula.

$$D = 0.94\ \lambda / \beta \cos \theta$$

where D is the average crystallite domain size perpendicular to the reflecting planes, λ is the X-ray wavelength, β is the full width at half maximum (FWHM), and θ is the diffraction angle. To eliminate additional instrumental broadening the FWHM was corrected, using the FWHM from a large grained Si sample.

$$\beta \text{ corrected} = (FWHM^2_{sample} - FWHM^2_{si})^{1/2}$$

This modified formula is valid only when the crystallite size is smaller than 100 nm [26].



S. No.	2Θ value	Plane	Element	Phase
1	38.114	111	Ag	Cubic
2	64.526	110	Ag	Hexagonal
3	67.529	112	Ag	Hexagonal

Fig. 2: XRD pattern of silver nanoparticles

2.6. SEM analysis of silver nanoparticles

Scanning Electron Microscopic (SEM) analysis was done using Hitachi S-4500 SEM machine. Thin films of the sample were prepared on a carbon coated copper grid by just dropping a very small amount of the sample on the grid, extra solution was removed using a blotting paper and then the film on the SEM grid were allowed to dry by putting it under a mercury lamp for 5 min.

2.7. Antibacterial assays

The antibacterial assays were done on *Escherichia coli* and *Pseudomonas aeruginosa* by standard disc diffusion method. Briefly Luria Bertani (LB) agar medium was used to cultivate bacteria. Fresh overnight cultures of inoculum (100 µl) of each culture were spread on to LB agar plates. Sterile paper discs of 3mm diameter (containing 50mg/litre silver nanoparticles) along with four standard antibiotic containing discs were placed in each plate.

2.8. Antifungal Assays

The antifungal assays were done on *Aspergillus flavus* by food poisoning method. Potato dextrose agar (PDA) medium was used in the study. The medium of each Petri dish contains 30 ppm silver nanoparticles were inoculated each alone at the centre with 5mm Inoculum disc of each pathogenic fungus and incubated at 25 °C for 7 days. The medium with Inoculum disc of each fungus but without silver nanoparticles served as control.

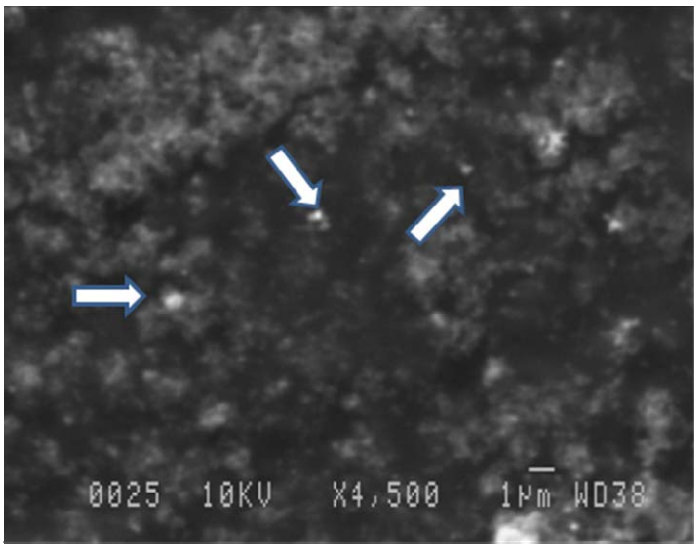


Fig. 3. SEM Micrograph of Silver Nanoparticles

3. Results and discussion

It is well known that silver nanoparticles exhibit yellowish brown color in aqueous solution due to excitation of surface plasmon vibrations in silver nanoparticles [27]. As the Argemone leaf extract was mixed in the aqueous solution of the silver ion complex, it started to change the color from watery to yellowish brown due to reduction of silver ion; which indicated formation of silver nanoparticles. It is generally recognized that UV–Vis spectroscopy could be used to examine size- and shape-controlled nanoparticles in aqueous suspensions [28]. Figure 1

shows the UV-Vis spectra recorded from the reaction medium after 4 hours. Absorption spectra of silver nanoparticles formed in the reaction media has absorbance peak at 440 nm, broadening of peak indicated that the particles are polydispersed.

The biosynthesised silver nanostructure by employing Argemone leaf extract was further demonstrated and confirmed by the characteristic peaks observed in the XRD image (Figure 2) and the structural view under the scanning electron microscope (Figure 3). The XRD pattern showed three intense peaks in the whole spectrum of 2θ value ranging from 10 to 80. Average size of the particles synthesized was 20 nm with size range 10 to 50nm with cubic and hexagonal shape. The typical XRD pattern (Fig. 2) reveled that the sample contains a mixed phase (cubic and hexagonal) structures of silver nanoparticles. The average estimated particle size of this sample was 20 nm derived from the FWHM of peak corresponding to 111 plane (figure 2). The SEM image showing the high density silver nanoparticles synthesized by the Argemone leaf extract further confirmed the development of silver nanostructures.

Further the nanoparticles syntheses by green route are found highly toxic against pathogenic bacteria and fungi at a concentration of 30 ppm (Table 1). Antibacterial effects of Ag nanoparticles obeyed a dual action mechanism of antibacterial activity, *i.e.,* the bactericidal effect of Ag^+ and membrane-disrupting effect of the polymer subunits.

Table1. Effect of silver nanoparticles on human pathogens

Name of the pathogen	**Disease**	**Zone of Inhibition (mm)**	
Bacteria		Food poisoning method (50 ppm Ag NPs)	
		Treated	Control
Escherichia coli	Cholecystitis, Bactremia, Cholangitis, Diarrhea	15 ± 0.4	45 ± 0.2
Pseudomonas syringae	Urinary tract infection, Ventilor associated Pneumonia	10 ± 0.5	40 ± 0.3
Fungi		Food poisoning method (50 ppm Ag NPs)	
		Treated	Control
Aspergillus flavus	Mycotoxicosis	10 ± 0.2	30 ± 0.1

Reduction of silver ions present in the aqueous solution of silver complex during the reaction with the ingredients present in the Argemone leaf extract observed by the UV-Vis spectroscopy revealed the presence of silver nanoparticles may be correlated with the UV-Vis spectra. UV-Vis spectroscopy is well known to investigate shape and size controlled of nanoparticles. The XRD and SEM analysis showed the particle size between 25-50 nm as well the cubic structure of the nanoparticles. FTIR analysis confirmed that the Bioreduction of Ag+ ions to silver nanoparticles are due to the reduction by capping material of plant extract. The Silver

nanoparticles synthesized via green route are highly toxic to multidrug resistant bacteria hence has a great potential in biomedical applications. The present study showed a simple, rapid and economical route to synthesized Silver nanoparticles.

4. Conclusions

In conclusion, the bio-reduction of aqueous Ag+ ions by the Argemone leaf extract has been demonstrated. The reduction of the metal ions through leaf extracts leading to the formation of silver nanoparticles of fairly well-defined dimensions. But the capabilities of Argemone leaf extract as a capping and reducing agent is not tested and not well defined. In the present study we found that Argemone can be also good source for synthesis of silver nanoparticles. This green chemistry approach toward the synthesis of silver nanoparticles has many advantages such as, ease with which the process can be scaled up, economic viability, etc. Applications of such eco-friendly nanoparticles in bactericidal, wound healing and other medical and electronic applications, makes this method potentially exciting for the large-scale synthesis of other inorganic materials (nanomaterials). Toxicity studies of silver nanoparticles on human pathogen opens a door for a new range of antibacterial agents. The reduction of silver ions and stabilization of the silver NPs was thought to occur through the participation of Leaf proteins and metabolites. Most importantly, the reaction was simple and convenient to handle, and it is believed that it has advantages over other biological syntheses.

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