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SYNERGISTIC EFFECT OF BIO-CONJUGATE SILVER NANOPARTICLES WITH ANTIBIOTIC: ANTIBACTERIAL AGENT AGAINST MULTI-DRUG RESISTANCE BACTERIA

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ABSTRACT: Green synthesis of silver nanoparticles from naturally occurring biomaterials provides an alternative, eco-friendly and cost-effective means of obtaining nanoparticles. It is an important thrust area for all scientists and has gained importance because it prevents the environment from pollution. Multi-drug resistance is a growing problem in the treatment of infectious diseases and the widespread use of broad-spectrum antibiotics has produced antibiotic resistance for many human bacterial pathogens. Advancement in nanotechnology has opened a new field in nanomedicine, allowing the synthesis of nanoparticles and evaluation of their synergistic effect with pre-existing antibiotics that can be assembled into a complex structure. Novel studies and technologies are developed to understanding the mechanisms of disease for the design of new drugs, but unfortunately, infectious diseases continue to be a major health problem worldwide. Since ancient times, silver was known for its anti-bacterial effects and for centuries it has been used for prevention and control of disparate infections. Now-a-days vancomycin resistance in gram-positive Cocci is a growing global health problem. In the early 1950s, vancomycin was used as the primary antibacterial agent for the treatment of infections caused by methicillin-resistant Staphylococcus aureus (MRSA). Progress in the development of new drugs to use vancomycin has been improved by using nanomedicines. Silver nanoparticles are attracting much interest because of their potent antibacterial activity. Many studies have also shown an important activity of silver nanoparticles against bacterial infections. This paper explains the new ideas to overcome current challenges and solutions in the treatment of infectious diseases, particularly the use of nanosilver particles as antimicrobials. The main objective of this work is to prepare silver nanoparticles from lichen, then prepare nanoconjugates with vancomycin antibiotics and explore their antibacterial properties. In the present study, we report a simple method of synthesis of bioconjugates of silver nanoparticles from ethanolic extract of lichen and their characterization by Ultraviolet-Visible (UV-vis), Fourier Transform Infrared (FTIR) spectroscopy, PL, XRD and TEM) analyses. Nanoparticles thus obtained were tested for antimicrobial activity against grampositive bacteria and gram-negative bacteria.

Keywords: Biosynthesis, Lichen, Silver nanoparticles, Bioconjugates, Antimicrobial activity, XRD, TEM, MDRB

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INTRODUCTION: Metal nanoparticles (NPs) have attracted much attention during recent years owing to their excellent properties which are different from bulk material.



These particles gained importance during recent years owing to their broad-spectrum application in a number of processes such as agriculture, cosmetics, healthcare, drug or gene delivery, medical devices, biosensor and catalysis ^{1, 2, 3, 4, 5, 6, ^{7, 8, 9} beside their antimicrobial properties ^{10, 11}. Many metal NPs are essential nutrients to the living system while some are toxic ¹². Their efficiency depends on their shape and size. Among the coinage metals silver has highest thermal and electrical conductivity.}

They may have a multidimensional structure such as nano-tubes and nano-wires. The green method of NP synthesis using plant extracts, bacteria, actinomycetes, fungi and enzymes are therefore, frequently used because of their environmentfriendly nature and biocompatibility ^{13, 14, 15, 16}. The lichen, belonging to the Usneaceae family, grows like moss on trees in a temperate climate. They are slowest growing plants living in symbiosis with algae, fungi and perennial trees. Different genera of lichens are used in curing dyspepsia, amenorrhea and vomiting. Lichens produce secondary metabolites that are used as crude drugs. Silver nanoparticles have become a focus of interest because they play a significant role in biological systems, living organisms, and medicine ^{17, 18}.

In the present medical industry, silver NPs are very useful in preventing infection in burns and open wounds ¹⁹. Silver nanoparticles have also been reported to possess antifungal ²⁰ antiviral ²¹, and anti-platelet activity ²². They are also beneficial as antimicrobials and therapeutic agents ^{23, 24} and the development of Ag NPs as an antibacterial agent is underway. Silver nanoparticles (AgNPs) have found plenty of applications as antibacterial agents because of their biological activities and safety. Their synthesis is generally carried out through the reduction of Ag³⁺ using inorganic agents or through biogenic approaches.

Numerous functional bioconjugate NPs such as the single-strand DNA²⁵, anti-bodies, ciprofloxacin²⁶ phospholipids, and various proteins capped NPs, have emerged in the past decade for the development of surface modification techniques on chemical structure, allowing useful application in the fields of drug design, drug delivery, biosensors, and so on ²⁷. In our study vancomycin was conjugated to the biosynthesized silver nanoparticles, starch was used as a bridge linker and then its antimicrobial activities against vancomvcin-resistant MDR pathogens were determined.

EXPERIMENTAL:

Materials and Methods: AgNO₃ (Merck, India Ltd.), ethanol (AR grade) and double distilled water were used. An aqueous solution of starch was used as a bridge linker. Lichen was procured from the medicinal garden of ITM University, Gwalior,

India. UV-Vis spectral measurements were done with Perkin Elmer Spectrophotometer between 200 and 800 nm. FTIR spectra were recorded with Perkin-Elmer Spectrometer, FTIR spectrum one, in 4000-400 cm⁻¹ region as KBr disc. TEM Images of Ag NPs were obtained using JEOL, JEM 2100 transmission electron microscope at 190 KV.

Samples were prepared using a drop of colloidal solution of Ag NPs on a carbon-coated copper grid and to allow the above sample to completely dry in a vacuum desiccator. The sediment particles obtained were used for scanning FTIR spectra. TEM images were obtained with a TEM electron microscope. Doubly distilled and deionized water was used as the solvent for preparing the stock solutions of all reagents. Buffer and all other chemicals were prepared according to standard laboratory procedures and the manufacturer's guidelines.

Preparation of Plant Extract: Lichens plants were collected from the medicinal garden of ITM University, Gwalior, Madhya Pradesh. Lichens were then washed thoroughly with distilled water and 20% alcohol then air-dried, homogenized. The sample is then filtered and dried by using a rotary evaporator. The dried sample was collected and stored at 4 °C for further use.

Phytochemical Screening of the Plant Extract: The extract was then subjected to qualitative phytochemical screening, which was done to understand the phytochemical constituents of the plant extract chosen for the study. For this, some specific tests were performed to evaluate the presence of particular phytochemicals. The tests performed for checking the availability of flavonoids, alkaloids, glycosides, steroids, phenols, saponins, terpenoid, cardiac glycosides and tannins²⁸.

Synthesis of Silver Nanoparticles: Aqueous solution of silver nitrate 0.01M and thiosemicarbazide 0.01 M were prepared in separate beakers by adding 10 ml distal water and vigorous magnetic stirring for about 10 min. An ethanolic solution of lichen was added in silver nitrate solution and stirred for about 1 h. The solution of thiosemicarbazide is now added slowly dropwise and further stirred for another 30 min at

60 °C. The solution was then centrifuged at 4000 rpm for 20 min. The obtained NPs were washed several times using ethanol and dried at room temperature for 36 h in atmospheric condition was collected and allowed to centrifuge at 12000 rpm. Pellets were collected and then washed thrice by means of ethanol and then kept for drying on the dry bath. The sample is then collected and allowed for characterization.

Preparation of Bioconjugate of Silver Nanoparticles: Conjugates were prepared to disperse Ag-NPs in 25 ml phosphate buffer (50mM, pH 6.5) in the presence of starch as a binding agent. After 30 min, 20 mg of vancomycin was added to the mixture and stirred at room temperature for 24 h under vigorous mixing. Excess vancomycin and other chemicals were eliminated by removing the supernatant after centrifugation at 12000 rpm for 20 min, and resuspended in deionized water and stored at 4 °C.

Characterization of Silver Nanoparticles: UV-Vis spectral measurements were done with a Perkin Elmer Spectrophotometer between 200 and 800 nm. FTIR spectra were recorded with Perkin-Elmer Spectrometer, in 4000-400 cm⁻¹ region as KBr disc. TEM Images of NPs were obtained using JEOL, JEM 2100 transmission electron microscope at 190 KV. Samples were prepared using a drop of colloidal solution of Ag NPs on a carbon-coated copper grid and to allow the above sample to completely dry in a vacuum desiccator. The sediment particles obtained were used for scanning FTIR spectra.

Bacterial Strains and Antimicrobial Testing Conditions: Bacterial strains and their relevant attributes are mentioned in **Table 1**. Strains were received from PGI Chandigarh, India and maintained in sterile broth medium.

TABLE 1: BACTERIAL STRAINS USED IN OURSTUDY

S. no.	Bacterial strains	Strain no.
1	Escherichia coli	MTCC-1563
2	Stayphylococcus aureus	MTCC- 3160
3	Enterococcus faecalis	MTCC-439

To determine the minimum inhibitory concentration (MIC) of silver nanoparticles and vancomycin conjugate AgNPs strains were grown in Muller Hinton Broth with aeration at 37 °C for

24h. The effects of nanoparticles, pure vancomycin, and vancomycin capped AgNps on individual bacterial isolates were determined ²⁹. The according to the following method minimum inhibitory concentration (MIC) determined by the microdilution method with serially diluted samples. The samples were diluted to get a series of concentrations from 0.19 mg/ml to 100 mg/ml in sterile nutrient broth.

The bacterial inoculums were prepared by adjusting the turbidity to 0.5Mc Farland $(1.5 \times 10^8 \text{ CFU/ml})$ standard. The microorganism suspension of 50 microlitres was added to the broth dilutions. These were incubated for 18 h at 37°C. Then 10 microlitre of each suspension was added to the solution to obtain a final concentration. The lowest preventing concentration of antibiotic the appearance of turbidity is considered to be UV spectro-photometer at 660 nm. Similarly, this protocol was performed for plant extract, AgNPs, pure vancomycin and bioconjugate of AgNPs. The results were then compared to 30 .

RESULTS AND DISCUSSION:

Qualitative Analysis of Phytochemicals Present in Plant Extract: The plant extract chosen for the study showed the presence of the following phytochemicals, discussed in Table 2.

TABLE 2: QUALITATIVE ANALYSIS OF PHYTO-
CHEMICALS PRESENT IN PLANT EXTRACT

S. no.	Phytochemicals	Ethanolic extract of lichen		
1	Steroids	Present		
2	Alkaloids	Present		
3	Flavonoids	Absent		
4	Terpenoids	Absent		
5	Tannins	Present		
6	Saponins	Present		
7	Phenol	Present		
8	Quinone	Absent		
9	Coumarin	Present		
10	Protein	Present		
11	Sugar	Present		
12	Gum	Present		

Visible Analysis: Silver nitrate solution has turned dark brown in color, which indicated the formation of AgNPs **Fig. 1**. This color change is due to the property of quantum confinement which is a size-dependent property of nanoparticles that affects the optical property of the nanoparticles. The color change in silver nitrate solution treated with Plant leaf extract.



FIG. 1: SYNTHESIZED SILVER NANOPARTICLES FROM PLANT EXTRACT

UV Spectra: It is known that when nanoparticles are formed, the colour of the solution containing both the NPs and plant extract turns dark brown depending on the presence of organic molecules in the extract. In our study, AgNO₃ was mixed with ethanolic extract of lichens incubated at room temperature; its color turned dark brown after 24h. The UV-Vis spectrum of this colloidal solution was run from 200 to 800 nm at room temperature which displayed peaks at 350 and 400 nm. The highest peak at 385 nm has been attributed to the excitation of surface plasmon resonance (SPR) of AgNPs.



FIG. 2: ULTRAVIOLET-VISIBLE ABSORPTION SPECTRUM OF SYNTHESIZED SILVER NANO-PARTICLES USING ALCOHOLIC EXTRACT OF LICHEN

Photo-oxidation of chemical constituents present in the extract may also have occurred ²⁹. Profile of the UV-Vis spectrum depends on the concentration of substrate, silver ions). Colour of these NPs remained unchanged even after several weeks ³⁰. It has been observed that when the colloidal solution containing Ag NP is slowly heated up to 60 °C the color intensity increases with increasing temperature and NPs are quickly formed. It demonstrates the effect of temperature on the

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biosynthesis of NPs. Absorption peaks in the UV-Vis spectrum are related to the shape of NPs. According to the criterion of Zhang and Nogues^{31,} the peaks at 385 nm correspond to cubical Ag NPs. Since we have observed a major peak at 385 nm AgNPs are supposed to be mainly spherical though the presence of a small amount of other types of NPs cannot be ignored.

FT-IR Spectral Analysis: FT-IR spectrum was run to identify the involvement of biomolecules present in lichen for the reduction of AgNO₃ to AgNPs. It is known to contain phenol, amines, aldehydes, and ketones besides many other compounds in traces. Since, all these compounds are excellent reducing agents, they undergo changes in stretching frequencies of their functional groups as a consequence of the reduction of AgNO₃ to AgNPs. IR spectrum is very complicated because of the overlap of frequencies in the same region.



FIG. 3: FT-IR SPECTRA OF SILVER NANOPARTICLES

However, we have attempted to identify the shifts in stretching vibrations after the formation of Ag NPs. Primary amines exhibit two N-H stretching frequencies in 3500-3300 cm⁻¹ region which has been found to appear at 3410 and 3455 cm⁻¹ in the NPs containing lichen extract. The band is 1600-1500 cm⁻¹ is due to CO stretching but amide bands also appear in the same region of the spectrum.

We have observed amide I and amide II bands at 1658 and 1540 cm⁻¹. A band at 1560 cm⁻¹ has been assigned to (C=O) stretching frequency. The COO⁻ group generally appears above 1600 cm⁻¹ but overlaps with amide II band ³². These spectral results indicate the involvement of organic molecules in the reduction of AgNO₃ leading to the formation of Ag NPs.

Fluorescence Analysis: The emission spectra of biosynthesized silver nanoparticles have been measured in the ethanolic solution at room temperature in a very dilute solution $(10^{-3} \text{ to } 10^{-5} \text{ mol})$. The fluorescence spectra of AgNPs are shown in **Fig. 4**. The AgNPs showed an emission band at 395 nm, 450 nm upon photoexcitation at 384 and a weak band at 695 nm.



FIG. 4: FLUORESCENCE SPECTRA OF SILVER NANOPARTICLES

XRD Analysis: The synthesized AgNPs were further characterized using X-ray diffractometry Fig. 5. The X-ray diffraction pattern showed four intense peaks (38.2° , 44.3° , 64.5° , 77.5°) in the whole spectrum of 2θ values ranging from 20° to 80°, which correspond to the (111), (200), (220), and (311) crystallographic planes of face-centered cubic silver, respectively. The crystallite domain size was calculated using the Debye-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 wavelength of Cu_{ka}, β is the full width at half maximum, and θ is the Bragg diffraction angle. The average crystallite size of the synthesized AgNPs was 0.54 nm, derived from the full width at half maximum of the peak corresponding to the (111) plane, which is in a good agreement with the particle size determined by TEM analysis³¹.



FIG. 5: XRD SPECTRA OF SILVER NANO-PARTICLES

 TABLE 3: DETERMINATION OF CRYSTALLINE SIZE OF BIOSYNTHESIZED AgNPs BY USING DEBYE

 SCHERRER'S EQUATION

S. no.	2-theta (deg.)	d (angle)	FWHM(deg.)	Particle size
1.	38.2	2.36	0.652	0.300nm
2.	44.3	2.054	1.40	0.21 nm
3.	64.5	1.44	0.84	1.2 nm
4.	77.5	1.234	0.91	0.46nm

Size and Morphology of Biosynthesized Silver Nano Particles & Its Bioconjugate: The shape and size morphology of biosynthesized Ag- nanoparticles and its bioconjugate were characterized using Transmission Electron Microscopic (TEM) study as shown in **Fig. 6A-B** which demonstrated the formation of slightly spherical AgNPs and irregularly shaped its bioconjugate. The size of the AgNPs & its bioconjugate is 50 nm 200 nm.

In-vitro Antibacterial Screening: Bacteriological examination was performed in sterile nutrient for both liquid systems. The conjugates exhibited antibacterial activity against MDR bacterial strains. The MIS of plant extract, pure vancomycin and vancomycin capped silver nanoparticles for these

bacteria was calculated as the lowest concentration at which bacterial growth was inhibited ³².

The MIC of pure vancomycin and AgNPs and vancomycin capped AgNPs for these bacteria was calculated as the lowest concentration at which bacterial growth was inhibited. The MIC of pure vancomycin for gram-negative strains was $\geq 3.2 \mu g/ml$. When pure AgNPs were employed, the MIC was 0.186 $\mu g/ml$, 0.197 $\mu g/ml$ & 0.051 $\mu g/ml$. Furthermore, when NPs are used as a capping agent in combination with vancomycin, MIC decreased considerably. In this condition, the MIC were 0.009 $\mu g/ml$, 0.053 $\mu g/ml$ & 0.039 0 $\mu g/ml$. The MIC corresponded to the MBC in all bacterial species. The results are shown in **Table 4**.



FIG. 6: TEM MICROGRAPH OF (A) BIOSYNTHESIZED Ag NANOPARTICLES (B) ANTIBIOTIC CAPPED AgNPs

TABLE 4: EFFECTS OF PLANT EXTRACT, BIOSYNTHESISED AgNPs, PURE VANCOMYCIN AND VANCOMYCIN CAPPED SILVER NANOPARTICLES

S.	Bacterial	Plant	Biosynthesised	Pure	Vancomycin
no.	strains	extract	AgNPs	vancomycin	capped AgNPs
1	Escherichia coli (MTCC -1563)	0.25	0.186	> 3.2	0.009
2	Staphylococcus aureus (MTCC-31620)	0.20	0.197	3.2	0.053
3	Enterococcus faecalis (MTCC- 439)	0.197	0.051	3.2	0.006



FIG. 7: IN-VITRO ANTIBACTERIAL ACTIVITY (A) PLANT EXTRACT (B) BIOSYNTHESIS SILVER NANOPARTICLES (C) VANCOMYCIN CAPPED SILVER NANOPARTICLES

The literature revealed that AgNPs with larger surface area provides better contact with microorganisms ³³. Thus, these particles may penetrate the bacterial cell membrane or attach to the bacterial surface and inhibit their replication ^{34, 35}. In our experiment, AgNPs have been found to be most effective against *Enterococcus faecalis*. It has been reported that antibacterial efficiency is increased by lowering the particle size ³⁶. Usually, NPs attach to the cell wall of bacteria and damage membrane and respiration system leading to cell death ³⁷. The toxicity of smaller NPs was greater than those of larger ones because the smaller ones can easily adhere to bacterial cell wall ³⁸.

Mode of Action of Silver Ions: Silver ions penetrate into the cytoplasm; denature the ribosome leading to the suppression of enzymes and proteins

which eventually arrest their metabolic function resulting in apoptosis of bacteria. Bactericidal activity is due to silver ions released from Ag NPs as a consequence of their interaction with microbes ¹¹. However, four possible mechanisms of antibacterial activity of AgNPs have been proposed (a) interference during cell wall synthesis (b) suppression during protein biosynthesis (c) disruption of the transcription process and (d) disruption of primary metabolic pathways ¹⁷.

Each mechanism involves structural changes, biochemical changes and charges on both the silver ions and biomolecules in the microbial cells. AgNPs also inhibit the proliferation of cancer cell lines by different modes of action ³⁷. They mediate and amplify the death signal by triggering the activation of Caspase-3 molecule. The DNA splits

into fragments by Caspase-3. Ag NPs may interfere with the proper functioning of cellular proteins and induce subsequent changes in cellular chemistry. Sometimes Ag NPs alter the function of mitochondria by inhibiting the catalytic activity of lactate dehydrogenase. AgNPs may also cause proliferation of cancer cells by generating ROS which ultimately leads to DNA damage ⁴⁰.

CONCLUSION: The present study aims to employ the use of plants as a source of synthesizing silver nanoparticles with less hazards towards the environment, easily scaled up, stable and economically viable. The silver nanoparticles were synthesized by using lichen extract. The biosynthesized nanoparticles are then allowed to characterize by means of visible observation, UV Spectroscopy, XRD, & TEM analysis.

The present study provides new insight into the synthesis of stable silver metal nanoparticles with less hazards along with exploring its antibacterial activity in the development of a potential tool in biomedics. In our study, we obtained the UV-Vis absorption of the AgNPs around 385 nm that shows the plasmaon resonance peak. The shape and size of AgNPs and the context in which the assay is performed are important factors in its capacity to annihilate superbugs.

This work explained that vancomycin can be conjugated with AgNPs to enhance antibacterial activity against gram-positive bacteria, but not gram-negative bacteria which can be justified because it is associated with the outer membranes of the gram-negative bacteria. *In- vitro* bactericidal examinations showed that the vancomycin capped with AgNPs exhibited substantial activity against MDRB. According to these results, we can say that such a nano-conjugate system may provide a very important method for the development of a new generation of effective antibacterial agents.

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