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## Biogenic synthesis of a silver nanoparticle– SnZrMoP nanocomposite and its application for the disinfection and detoxification of water

Rupinder Kaur,<sup>a</sup> Sandeep Kaushal<sup>b</sup> and Prit Pal Singh<sup>\*b</sup>

A silver nanoparticle (AgNP)–tin zirconium(iv) molybdophosphate (SnZrMoP) nano-composite was synthesized by a superficial and green synthetic approach using mulberry leaf extract, with the plant extract playing the role of a reducing agent as well as a capping agent. The nanocomposite was characterized by Fourier-transform infrared spectroscopy (FTIR), scanning electron microscopy (SEM), transmission electron microscopy (TEM), energy dispersive X-ray spectroscopy (EDX), X-ray diffraction (XRD), and thermogravimetric analysis (TGA). The chemical stability of the nanocomposite and the effect of calcination on the ion-exchange capacity of the nanocomposite were also investigated. The nanocomposite was observed to be highly selective for Ba<sup>2+</sup> and Sr<sup>2+</sup> ions on the basis of distribution coefficient studies. The applications of the nanocomposite as an antimicrobial agent and for the quantitative and selective separation of toxic Ba<sup>2+</sup> and Sr<sup>2+</sup> ions from industrial effluents were explored.

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### Introduction

The natural habitats of living organisms have been depleted with the illimitable flourishing of industrialization and unrestrained pollution. The expulsion of industrial waste into the environment has added many major pollutants to ecosystems, such as pesticides, antibiotics, and heavy metals. The presence of even traces of heavy metals in natural water bodies can cause major health hazards to humans and animals.<sup>1</sup> The metals of particular concern in industrial effluents are cadmium, chromium, lead, mercury, silver, arsenic, aluminium, copper, cobalt, manganese, and barium. Being a divalent-alkaline earth metal, barium forms salts with carbonates, nitrates, chlorides, and sulphides and thus has been widely employed in different fields, such as brick-making, ceramic glaze, cement, glass, paint, and plastic industries.<sup>2–5</sup> Barium can be rapidly absorbed from the gastrointestinal tract and deposited in the muscles, lungs, and bones.<sup>6</sup> A low dose of barium can act as a muscle stimulant, but higher doses of barium can have an adverse impact on the nervous system and can lead to cardiac irregularities, weakness, anxiety, shortness of breath, and paralysis. Industrial emissions are the main source of barium in the atmosphere.

Nanomaterials of different sizes, shapes, and compositions have found comprehensive applications. Metallic nanoparticles

among these materials are of great interest because of their inimitable optical, electrical, and electrochemical properties.<sup>7,8</sup> Among the various metal nanoparticles, silver nanoparticles (AgNPs) are of immense interest because of their versatile applications as antimicrobial agents, biosensor materials, composite fibers, cryogenic superconductor materials, cosmetic products, and electronic components.

The synthesis of AgNPs can be carried out by a number of techniques, like laser ablation, gamma irradiation, electron irradiation, chemical reduction, photochemical methods, microwave processing, and biological synthetic methods.<sup>9</sup> However, the majority of routes for the synthesis of nanoparticles encompass the use of hazardous chemicals or high energy requirements. To overcome these demerits, biogenic nanoparticles synthesized from plant extracts without making use of any poisonous chemicals would be considered attractive for nanotechnology.<sup>10</sup> The phytochemicals present in plants are water soluble and can cause the immediate reduction of metal ions.<sup>11</sup>

A number of medicinal plants, such as *Acorus calamus*, *Alternanthera dentate*, *Ocimum sanctum*, *Azadirachta indica*, *Brassica rapa*, *Coccinia indica*, *Vitex negundo*, *Solanum xanthocarpum*, *Croton bonplandianum*, *Andrographis paniculata*, and *Cleome viscosa*, have been reported in literature for the synthesis of biogenic Ag nanoparticles.<sup>12–20</sup> In this paper, we have reported a low-cost facile approach for the synthesis of silver nanoparticles from the mulberry leaf extract. Nanosized inorganic particles and cluster-based nanocomposites represent materials with the most appropriate properties for various applications.<sup>21,22</sup> Inorganic cation exchangers have already

<sup>a</sup> Guru Nanak Dev Polytechnic College, Ludhiana, Punjab, India<sup>b</sup> Department of Chemistry, Sri Guru Granth Sahib World University, Fatehgarh Sahib, Punjab, India. E-mail: dhillonps2003@gmail.com, kaushalsandeep33@gmail.com; Fax: +91-1763-234236; Tel: +91-84270-00415, +91-7009795652

been employed as a support to disperse and stabilize numerous nanoparticles.<sup>23,24</sup>

In the present work, AgNPs were synthesized from mulberry leaf extract in an environment friendly process. The AgNP–SnZrMoP composite was synthesized by the incorporation of biogenic silver nanoparticles (GAgNPs) in an SnZrMoP ion exchanger. The nanocomposite was found to be highly selective for Ba<sup>2+</sup> and Sr<sup>2+</sup> ions. The as-synthesized nanocomposite was employed for the efficient removal of Ba<sup>2+</sup> ions from industrial effluents, and as an antimicrobial agent to inhibit the growth of *E. coli* and *Bacillus*. To the best of our knowledge, there is no report in literature regarding the use of a hybrid nanocomposite of biogenic silver nanoparticles and SnZrMoP ion exchanger for the disinfection and detoxification of water.

## Experimental

### Reagents and instruments

Zirconium oxychloride, stannic chloride, sodium molybdate, sodium dihydrogen phosphate, ethylene diamine tetra acetic acid, silver nitrate, and all other chemicals and reagents were purchased from Loba Chemie, India and used as received. A single electrode pH meter (Toshniwal, India) was used for the pH studies. Fourier transform infrared (FTIR) spectra were recorded on a PerkinElmer RX1 FTIR spectrometer. X-ray diffraction (XRD) patterns were obtained using a PAN analytical system DY 3190 X-ray diffractometer. Scanning electron microscopy (SEM) images were obtained from an S-4800 field emission SEM system (JEOL scanning electron microscope) equipped to perform elemental chemical analysis by energy dispersive X-ray spectroscopy (EDX). Transmission electron microscopy (TEM) studies were performed using the MIC JEM 2100. Thermal analysis was performed by heating the nanocomposite material up to 900 °C at a constant increment of 10 °C per minute in a nitrogen atmosphere, using Hitachi SGA7400 thermogravimetric analyzer. UV-vis spectra were recorded in a quartz cuvette using a Shimadzu-UV 2600 spectrophotometer.

### Preparation of the reagents

Solutions of zirconium oxychloride and stannic chloride (0.1 M) were prepared in 1 M HCl solution. Sodium dihydrogen phosphate (0.1 M), sodium molybdate (0.1 M), silver nitrate

(0.1, 0.2, 0.3, and 0.4% w/v) and EDTA (0.01 M) solutions were prepared in double-distilled water (DDW).

### Synthesis of an inorganic gel of tin zirconium(IV) molybdophosphate (SnZrMoP)

Tin zirconium(IV) molybdophosphate samples were prepared at room temperature (25 ± 2 °C) by mixing 0.1 M solutions of each of sodium molybdate, sodium dihydrogen phosphate, stannic chloride, and zirconium oxychloride in different volume ratios. Ethylene diamine tetra acetic acid (0.01 M EDTA) was used as a capping agent.<sup>25</sup> Each solution of the salts was added into the EDTA solution kept under a constant stirring with a magnetic stirrer. The pH of the mixture was adjusted to 0–1 by adding aqueous ammonia or hydrochloric acid with constant stirring, and a white colored gel was obtained as the final product.

### Synthesis of green silver nanoparticles (GAgNPs)

The leaf extract was prepared from air dried and finely crushed mulberry leaves. The extract was then filtered using Whatman no. 1 filter paper, to remove the particulate matter. Different concentrations of AgNO<sub>3</sub> solution (0.1, 0.2, 0.3, 0.4, and 0.5% w/v) were prepared for optimization. For the reduction of Ag<sup>+</sup> ions to Ag, mulberry leaf extract was added to silver nitrate (AgNO<sub>3</sub>) solution with continuous stirring for 1 h at room temperature. Due to the surface plasmon resonance phenomenon, the color of the solution changed from colorless to pale yellow, and finally reddish brown, indicating the complete reduction of AgNO<sub>3</sub>.

### Synthesis of the GAgNPs–SnZr(IV) MoP nanocomposite

Four different samples of GAgNPs–SnZrMoP nanocomposite were prepared by varying the molar concentration of silver nitrate solution while keeping the volume ratio of different components of the inorganic counterpart constant (Table 1). The resultant slurry was kept for 24 h and then washed repeatedly with DDW. The material was filtered off and washed thoroughly with DDW to remove the excess acid. The filtered gel was dried at a temperature of 40 °C in a hot air oven. The dried product was immersed in DDW to obtain small granules. The resultant product was saturated with hydrogen ions by keeping the granules in 0.1 M HCl solution for 24 h with periodic stirring, and by changing the solution to get better results. After 24 h, the granules were filtered out and washed a number of times to remove the excess acid, and then dried at 50 ± 2 °C till a constant weight was obtained.

**Table 1** Optimization of conditions for the synthesis of the GAgNPs–SnZrMoP nanocomposite

Sample Code	Synthesis conditions						Wt of mulberry leaves (g)	%age (w/v) of AgNO <sub>3</sub> (100 mL)	pH	Properties		
	ZC (0.1 M) (mL)	SC (0.1 M) (mL)	SM (0.1 M) (mL)	EDTA 0.01 M (mL)	SDP (0.1 M) (mL)	Color of granules				Yield (g)	IEC (meq g <sup>-1</sup> )	
S-1	100	100	100	100	300	—	—	1	White	5.20	1.95	
S-2	100	100	100	100	300	3.00	0.10	1	Shiny grey	5.55	2.16	
S-3	100	100	100	100	300	3.00	0.20	1	Shiny grey	5.87	2.25	
S-4	100	100	100	100	300	3.00	0.30	1	Shiny grey	6.12	2.48	
S-5	100	100	100	100	300	3.00	0.40	1	Shiny grey	6.44	2.36	

ZC – zirconium oxychloride; SC – stannic chloride; SDP – sodium dihydrogen phosphate; SM – sodium molybdate; M – mol L<sup>-1</sup>.



### Ion-exchange capacity (IEC)

The standard column method was used to determine the ion-exchange capacity of the nanocomposite. The nanocomposite material (0.5 g) in H<sup>+</sup> form was placed in a glass column containing glass wool support at the bottom. The hydrogen ions eluted with 0.5 M NaNO<sub>3</sub> solution were estimated by titration against a standard sodium hydroxide (0.1 M) solution. The hydrogen ions released were then estimated by the formula:

$$\text{IEC} = \frac{N \times V}{W} \text{ meq g}^{-1}$$

where  $N$  and  $V$  are the normality and volume in mL of NaOH, respectively, and  $W$  is the weight in grams of the nanocomposite.<sup>26</sup>

### Effect of calcination on the ion-exchange capacity

The nanocomposite cation exchange material in H<sup>+</sup> form was heated at various temperatures for 1 h in the muffle furnace from 100 °C to 900 °C, in increments of 10 °C. The Na<sup>+</sup> ion-exchange capacity was determined by a batch process, after cooling the material to room temperature.<sup>26</sup>

### pH-Titrations studies and functionality of the exchanger

The Topp and Pepper method<sup>27</sup> was used for the pH-titration studies of the GAgNPs–SnZrMoP nanocomposite. An equimolar mixture of alkali and alkaline earth metal chlorides and their corresponding hydroxide solutions in different volume ratios (total volume being 50 mL) was added to each of the several 250 mL conical flasks containing 0.5 g of the nanocomposite in H<sup>+</sup> form. The pH of the solution was recorded every 24 h till the attainment of equilibrium.

### Chemical stability

The nanocomposite cation exchanger (0.2 g) in H<sup>+</sup> form was treated separately with 50 mL of different concentrations of strong acid (HCl), strong base (NaOH), and organic solvents, like a 10% solution of each of acetone and ethyl alcohol for 24 h, with occasional shaking. After removal of the excess reagent, the nanocomposite was dried in a hot air oven at 40 ± 1 °C and its ion exchange capacity was determined by the batch method.

### Distribution studies

The distribution coefficients ( $K_d$ ) for various metal ions were determined in different solvent systems. The nanocomposite cation exchange material in H<sup>+</sup> form (0.2 g) was placed separately in contact with 20 mL solutions of each of the different metal ions for 24 h, with occasional shaking to attain equilibrium. A standard solution of EDTA (0.01 M) was used for estimation of the amount of metal ions in solution, before and after equilibrium. The distribution coefficient ( $K_d$ ) represents the ratio of the amount of metal ions in the exchanger phase and the solution phase, and is given by:

$$K_d = \frac{I - F}{F} \times \frac{V}{M} \text{ mL g}^{-1}$$

where  $I$  is the initial concentration of metal ions in the solution phase,  $F$  is the final concentration of metal ions in the solution

phase,  $V$  is the volume of solution (mL), and  $M$  is the amount of ion exchanger taken (g).

### Binary separations

**Quantitative separation of metal ions in binary synthetic mixtures.** The cation exchanger (1 g) in H<sup>+</sup> form was taken in a glass column having glass wool support at the bottom. A mixture of two metal ions (0.01 M each) was passed through the column (flow rate: 0.25 mL min<sup>-1</sup>) till the solution level was just above the surface of the exchanger. The unexchanged metal ions were removed by rinsing the column with DDW. The adsorbed metal ions on the exchanger were eluted individually using appropriate eluting reagents. The fractions (10 mL) of the eluent were collected and the metal ions were estimated by titration against a standard solution of EDTA (0.01 M).

**Selective separation of Ba(II) ions from a synthetic mixture.** The selective separation of three different amounts of Ba<sup>2+</sup> and Sr<sup>2+</sup> ions (separately) from the synthetic mixtures containing a fixed amount of Mg<sup>2+</sup>, Ca<sup>2+</sup>, Co<sup>2+</sup>, Fe<sup>3+</sup>, and Ni<sup>2+</sup> was achieved on the column of the GAgNPs–SnZrMoP nanocomposite.

**Quantitative separation of Ba(II) ions from wastewater samples.** Water soluble barium salts are used in various industrial processes and laboratories. In the chemistry lab, barium chloride solution is used as a sulfate ion indicator in the presence of hydrochloric acid, while barium carbonate is chiefly used in brick-making and ceramic industries, and barium nitrate is used to manufacture high-quality glass with a low co-efficient of thermal expansion. Environmental remediation by the GAgNPs–SnZrMoP nanocomposite was checked on wastewater samples collected from the paint industry, brick-making industry, glass industry, and chemistry laboratory. The effluent samples were filtered to remove any suspended particles and neutralized. The color-producing substances were removed by adsorption on charcoal. The waste sample (100 mL) was passed over the column of GAgNPs–SnZrMoP and the effluent collected was recycled three times at a flow rate of 0.5 mL min<sup>-1</sup>. The metal ions were eluted with 0.5 M HNO<sub>3</sub> and were determined titrimetrically with 0.01 M EDTA solution.

### Antimicrobial studies

The antibacterial potential of GAgNPs, SnZrMoP, and the GAgNPs–SnZrMoP nanocomposite was investigated separately. At first, the cultures for *E. coli* and *Bacillus* were grown uninventively, in an incubator preserved at 37 °C with a shaking velocity of 200 rpm. The medium-term developed culture was shifted to flasks containing 20 mL Luria Broth. For both *E. coli* and *Bacillus*, a control experiment was set up without the addition of any chemical. Then, 500 µL of dimethyl sulfoxide (DMSO) was added in both *E. coli* and *Bacillus* cultures in another control experiment. In the test experiment, 50 mg of GAgNPs, SnZrMoP, and GAgNPs–SnZrMoP was added in different flasks, followed by 1 h of growth in a shaker. Thereafter, a 1 mL culture was withdrawn from every flask after a time span of 2 h. The optical density (OD 600 (nm)) was documented separately for individual samples and a graph was plotted of the OD 600 (nm) versus time (h) to measure the rate of growth inhibition.



## Results and discussion

A novel nanocomposite cation exchange material, GAgNPs–tin-zirconium(IV) molybdophosphate, was synthesized by embedding biogenic silver nanoparticles (GAgNPs) into a novel four-component inorganic cation exchanger tin-zirconium(IV) molybdophosphate. In the present investigation, green silver nanoparticles were synthesized from mulberry leaf extract without using any toxic chemicals, to reduce the environmental impact caused by various conventional techniques. The synthesis conditions were optimized by varying the %age (w/v) concentration of silver nitrate solution and by keeping the volume ratios of the inorganic counterparts constant (Table 1). The ion-exchange capacity increased with the increase in doping of silver nanoparticles up to a certain extent. The sample C-4 was selected for further studies due to its highest ion-exchange capacity. The higher ion-exchange capacity of the GAgNPs–SnZrMoP nanocomposite might be due to the interaction between the GAgNPs and SnZrMoP. The synthesis was carried out in the presence of EDTA, which acted as a capping agent and a polymer, and may also be responsible for the enhanced ion-exchange capacity.

### Characterization of the nanocomposite

The GAgNPs–SnZrMoP nanocomposite was characterized by UV-vis spectrophotometry (UV-vis), Fourier transform infrared (FTIR) spectroscopy, X-ray diffraction (XRD), scanning electron microscopy (SEM), transmission electron microscopy (TEM), energy dispersive X-ray (EDX), and thermogravimetric analysis (TGA-DTA) studies.

UV-vis spectroscopy is the simplest and most important technique to confirm the formation of nanoparticles. The absorbance spectrum of the sample in the range of 400–800 nm, using a Shimadzu-UV 2600 spectrometer confirmed the reduction of silver ions to silver nanoparticles (Fig. 1). The absorption maximum of the bioreduced GAgNPs at different time intervals was observed as a sharp band at 460 nm in the UV-vis spectra.

The FTIR spectrum of SnZrMoP (Fig. 2a) showed a broad band around 3431.77  $\text{cm}^{-1}$ , which could be attributed to symmetric

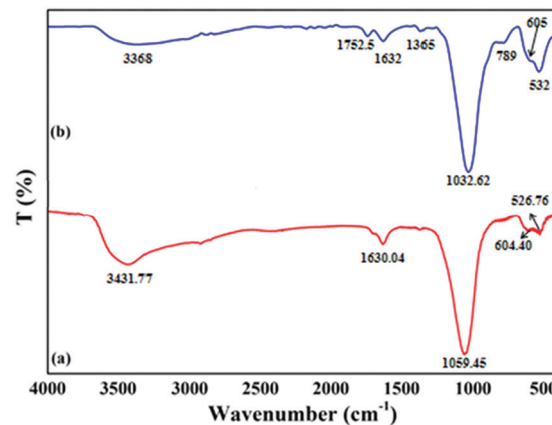


Fig. 2 FTIR spectra of (a) SnZrMoP and (b) GAgNPs–SnZrMoP nanocomposite.

and asymmetric –OH stretching. The absorption band at 1630.04  $\text{cm}^{-1}$  was due to the H–O–H bending of interstitial water in the composite material. A band in the region of 1059.45  $\text{cm}^{-1}$  corresponded to ionic phosphate stretching. The band at  $\sim 526.75 \text{ cm}^{-1}$  was attributed to the Zr–O bond. A sharp peak at 601.40  $\text{cm}^{-1}$  was ascribed to the presence of a molybdate group.<sup>28</sup> The FTIR spectrum of the GAgNPs–SnZrMoP (Fig. 2b) showed a broad band in the region of 3368  $\text{cm}^{-1}$ , which could be attributed to symmetric and asymmetric –OH stretching, and this peak may be due to the presence of alcohols and phenols in the plant extract. The peak at 1752  $\text{cm}^{-1}$  indicated the stretching vibrations of C=O group. The band at 1632  $\text{cm}^{-1}$  was attributed to N–H bending vibrations of primary amines. The band at 1365  $\text{cm}^{-1}$  corresponded to the C–N stretching of aromatic ring compounds. The broad band at 1051  $\text{cm}^{-1}$  may be due to the presence of  $\text{PO}_4^{3-}$ ,  $\text{HPO}_4^-$ , and  $\text{H}_2\text{PO}_4^{4-}$  groups. The band at  $\sim 532 \text{ cm}^{-1}$  may be due to the presence of Zr–O bond and Sn–O bond. The sharp peak at 605  $\text{cm}^{-1}$  was attributed to the molybdate group. The appearance of three new bands at 1752, 1365, and 789  $\text{cm}^{-1}$  may be due to the incorporation of green silver nanoparticles in the hybrid material. The variation in the wave numbers of different peaks may also be due to the embedding of silver nanoparticles into the tetravalent metal acid salt.

The X-ray diffraction pattern of the SnZrMoP ion exchanger showed characteristic peaks that match the JCPDS card number 00-024-1491, indicating the semicrystalline nature of the ion exchanger (Fig. 3a). The XRD spectrum of GAgNPs (Fig. 3b) showed four characteristic diffraction peaks at  $2\theta$  of 28.21 (111), 42.73 (200), 64.54 (220), and 76.63 (311). This pattern matches with JCPDS card number 89-3722, and corresponds to the fcc lattice of silver.<sup>29</sup> The XRD pattern of GAgNPs depicted its crystalline nature. The variations in the intensities of different peaks may be due to the presence of organic compounds in the plant extracts. The XRD pattern of the nanocomposite material GAgNPs–SnZrMoP (Fig. 3c) revealed its polycrystalline nature. The appearance of characteristic diffraction peaks at  $2\theta$  of 28.01 (111), 46.44 (200), and 66.54 (220) clearly demonstrated the incorporation of GAgNPs into the inorganic ion-exchanger.

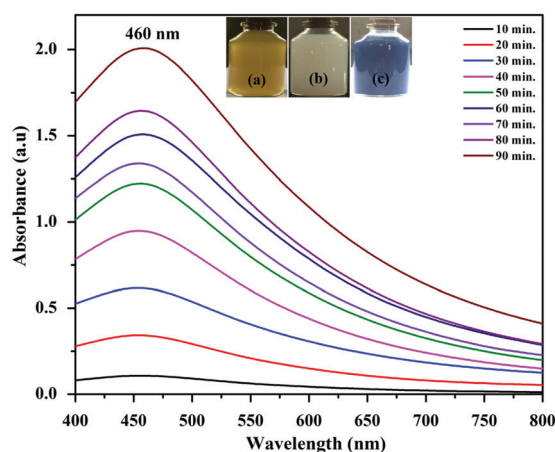


Fig. 1 UV-vis spectra of GAgNPs at different reaction times. (Inset: Images depicting the color of the synthesized (a) GAgNPs (b) SnZrMoP (c) GAgNPs–SnZrMoP.)





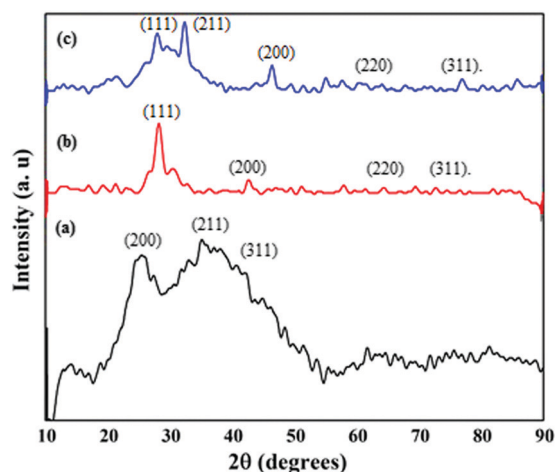


Fig. 3 X-ray diffraction patterns of (a) SnZrMoP; (b) GAgNPs; (c) GAgNPs-SnZrMoP nanocomposite.

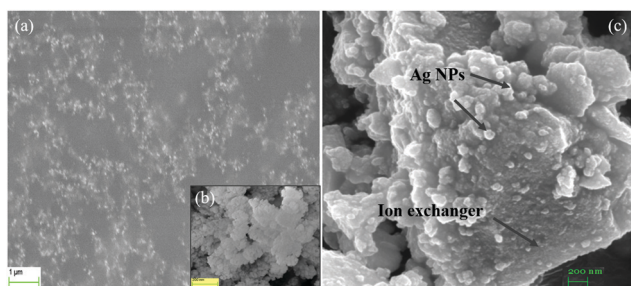


Fig. 4 FESEM images of (a) GAgNPs; (b) SnZrMoP; (c) GAgNPs-SnZrMoP nanocomposite.

The FESEM image (Fig. 4a) of the GAgNPs revealed that the particles were spherical in shape and agglomerated. The FESEM image of SnZrMoP (Fig. 4b) revealed that the inorganic ion-exchanger contained globular particles and there were areas of aggregations and agglomerations. The surface morphology of the nanocomposite material revealed that the material contained aggregates of silver nanoparticles, which were uniformly embedded in the matrix of the inorganic ion exchanger SnZrMoP (Fig. 4c).

The typical HRTEM images (Fig. 5a and b) revealed that the biogenic silver nanoparticles were spherical in shape. These nanoparticles were lying apart due to the reduction and capping

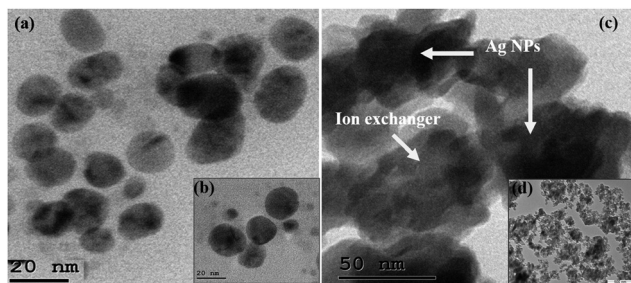


Fig. 5 HRTEM images of (a and b) GAgNPs; (c) GAgNPs-SnZrMoP nanocomposite; and (d) SnZrMoP ion-exchanger.

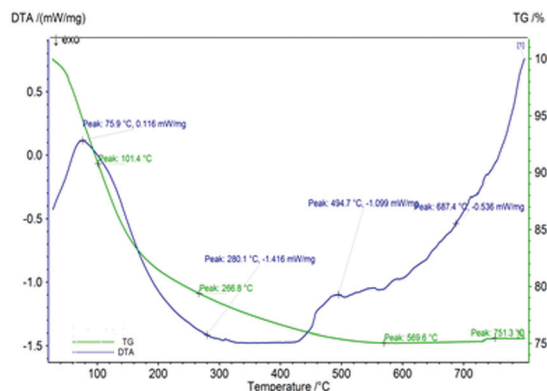


Fig. 6 TGA-DTA curves of the GAgNPs-SnZrMoP nanocomposite ion exchanger.

of silver nanoparticles with the phytochemicals present in the mulberry leaf extract. The average particle size of the GAgNPs was  $\approx 15$  nm, which was in good agreement with the particle size calculated from the XRD analysis. It could be observed from the images in Fig. 5d that the inorganic material was composed of a network of nearly spherical particles. It is clear from the HRTEM images of the synthesized AgNPs-SnZrMoP composite (Fig. 5c) that a large number of well-dispersed AgNPs were attached to the surface of the ion exchanger.

TGA/DTA studies of the GAgNPs-SnZrMoP nanocomposite hybrid ion exchanger are shown in Fig. 6. The nanocomposite ion exchanger was heated from room temperature to 800 °C, with an increment of 10 °C min<sup>-1</sup>. The weight loss up to 101 °C was  $\approx 8.99\%$ , which was due to loss of water molecules from the sample. This change was confirmed from the endothermic hump in the same region of the DTA curve. The weight loss in the 101–266 °C temperature range was  $\approx 11\%$ , which was due to the loss of the water of crystallization. The weight loss in the temperature range 266–569 °C was  $\approx 4.75\%$ , which was due to the conversion of inorganic phosphate into pyrophosphate.<sup>30</sup> There was no significant weight loss in the temperature range 594.7–800 °C which established high thermal stability of the nanocomposite material.

### Physiochemical properties

The ion-exchange capacity of alkali metals and alkaline earth metals follows the order:  $K^+ > Na^+ > Li^+$  and  $Ba^{2+} > Sr^{2+} > Ca^{2+} > Mg^{2+}$ , respectively (Table 2). The decrease in ion-exchange capacity of both the alkali and alkaline earth metals is in accordance

Table 2 IEC of the GAgNPs-SnZrMoP nanocomposite for alkali and alkaline earth metals

S. no.	Exchanging ion	Ionic radii (Å)	Hydrated ionic radii (Å)	IEC (meq g <sup>-1</sup> )
1	Li <sup>+</sup>	0.68	3.40	2.42
2	Na <sup>+</sup>	0.97	2.76	2.48
3	K <sup>+</sup>	1.33	2.32	2.55
4	Mg <sup>2+</sup>	0.78	10.80	2.62
5	Ca <sup>2+</sup>	1.06	9.60	2.73
6	Sr <sup>2+</sup>	1.27	6.30	2.80
7	Ba <sup>2+</sup>	1.43	5.90	2.89



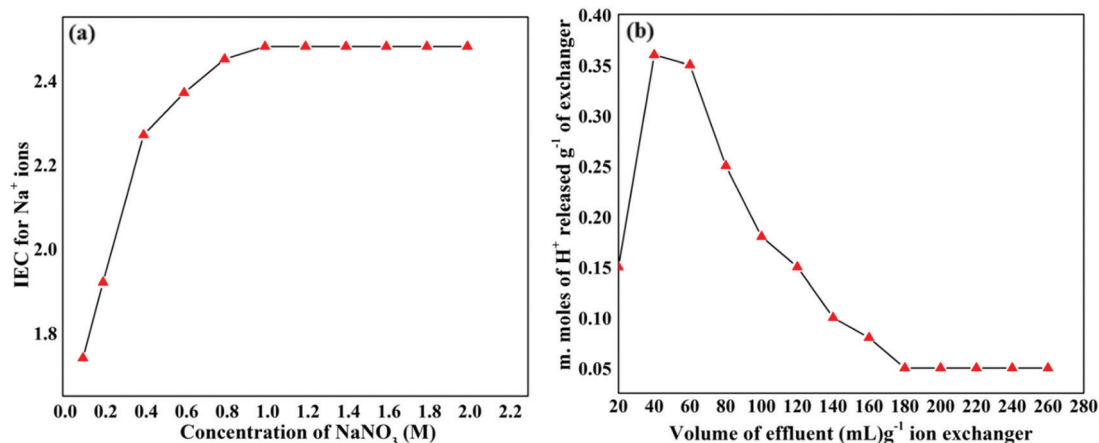


Fig. 7 (a) Effect of the eluent concentration; (b) elution behavior of the GAgNPs–SnZrMoP nanocomposite ion exchanger.

with decrease in their hydrated ionic radii. Coulomb's law is obeyed by ionic crystals, according to which a small cation will be attracted with a greater force and held more strongly than a bigger ion of the same charge. Also, ions having smaller hydrated ionic radii can easily penetrate into the pores of the exchanger and interact with the active sites for the exchange of H<sup>+</sup> ions, resulting in a higher ion-exchange capacity.<sup>24</sup>

The effect of eluent concentration on the ion-exchange capacity of the GAgNPs–SnZrMoP nanocomposite is shown in Fig. 7(a). It was observed that the degree of elution also depended on the concentration of the eluent. The optimum concentration of NaNO<sub>3</sub> as an eluent was observed to be 0.8 M for 1.0 g of the cation exchanger. The column elution experiment reflected the column efficiency and predicts the volume of eluent required for the complete elution of H<sup>+</sup> ions. It was observed from the elution behavior that the exchange rate is faster in the beginning and a maximum of H<sup>+</sup> ions were eluted in the first 150 mL of NaNO<sub>3</sub> solution (1 M). It was observed that about 250 mL of NaNO<sub>3</sub> solution (1 M) was sufficient to elute all the exchangeable H<sup>+</sup> ions from 500 mg of the sample in ~4 h (Fig. 7b).

The ion-exchange capacity of the nanocomposite was also affected by calcination (Table 3). The nanocomposite was heated in the temperature range of 100 °C to 800 °C for 1 h.

It was observed that there was a significant decrease in the ion-exchange capacity of the nanocomposite cation exchanger at higher temperature, which might be due to the condensation of phosphate groups to pyrophosphate at elevated temperature.

However, the ion exchanger was found to be quite stable to heat as the composite retained about 86.28% of its mass and 85.50% of its ion-exchange capacity up to a temperature of 300 °C.

It was observed that the nanocomposite had a reasonably good stability in different solvents. The chemical stability may be due to the binding of GAgNPs with the inorganic matrix. This can prevent the dissolution of the nanohybrid material or the leaching of any of its constituent elements in the solution. The GAgNPs–SnZrMoP was resistant to dissolution even in 1 M HNO<sub>3</sub> solution. The material showed negligible dissolution in organic solvents, while very feeble dissolution was observed in alkaline medium, which overall shows that the material exhibited good chemical stability (Table 4).

Distribution studies were performed for 15 different metal ions in diverse solvent systems to explore the potential of the nanocomposite for the separation of metal ions. It was observed that the *K<sub>d</sub>* values varied with the nature of the solvent. It was also observed (Table 5) that Ba<sup>2+</sup> was strongly adsorbed, and Sr<sup>2+</sup>, Pb<sup>2+</sup>, and Hg<sup>2+</sup> were significantly adsorbed, while other metal ions were poorly adsorbed on the surface of the nanocomposite.

Some analytically important binary separations were performed on the basis of the separation factors, such as: Cu(II)–Ba(II); Ca(II)–Ba(II); Mg(II)–Ba(II); Fe(III)–Ba(II); Ni(II)–Ba(II); Hg(II)–Ba(II). The order of elution and the eluents used for the binary separations are given in Table 6.

During the separation of Ba<sup>2+</sup> ions from the synthetic mixtures of different metal ions, nearly more than 90% of

Table 3 Effect of temperature on the ion-exchange capacity of the GAgNPs–SnZrMoP nanocomposite

S. no.	Heating temp. (°C)	Color of granules	Weight retained (%)	IEC for Na <sup>+</sup> ions (meq g <sup>-1</sup> )	Retained ion-exchange capacity (%)
1	50	Shiny grey	99.60	2.40	100
2	100	Blackish brown	94.12	2.30	97.56
3	200	Blackish brown	90.20	2.00	93.35
4	300	Black grey	86.28	1.50	85.50
5	400	Dark grey	78.43	1.45	82.40
6	500	Silvery grey	64.71	1.20	70.00
7	600	Silvery grey	64.71	1.05	52.81
8	700	Silvery grey	66.67	0.85	42.81
9	800	Silvery grey	66.67	0.85	36.32



Table 4 Chemical stability of the GAgNPs–SnZrMoP nanocomposite ion exchanger

S. no.	Solvent	Mass loaded (mg)	Mass retained (mg)	Mass retained (%)	Color	IEC for Na <sup>+</sup> ions (meq g <sup>-1</sup> )
1	1 M HNO <sub>3</sub>	200	200	98.80	Shiny grey	2.45
2	1 M HCl	200	198	99.20	Shiny grey	2.46
3	1 M H <sub>2</sub> SO <sub>4</sub>	200	198	98.80	Shiny grey	2.45
4	1 M NaOH	200	127	68.55	Shiny grey	1.70
5	5% Acetone	200	200	100	Shiny grey	2.48
6	5% Ethanol	200	196	99.20	Shiny grey	2.46

Table 5 *K<sub>d</sub>* values of some metal ions on the GAgNPs–SnZrMoP nano-hybrid ion-exchanger column in different solvent systems

Metal ions	DMW	0.1 M CH <sub>3</sub> COOH	0.1 M HNO <sub>3</sub>	0.5 M HNO <sub>3</sub>
Mg <sup>2+</sup>	29.50	22.71	19.61	13.57
Ca <sup>2+</sup>	64.94	47.08	39.65	29.22
Sr <sup>2+</sup>	890	627.18	548.77	268.69
Ba <sup>2+</sup>	1640	1399	1179	770
Cr <sup>3+</sup>	210	166.32	152	98.64
Fe <sup>3+</sup>	64.67	51.22	45.94	32.05
Co <sup>2+</sup>	5.71	4.80	4.33	1.97
Ni <sup>2+</sup>	14.00	10.64	10.36	5.58
Cu <sup>2+</sup>	29.13	22.48	21.86	11.00
Zn <sup>2+</sup>	66.40	54.86	41.71	19.88
Cd <sup>2+</sup>	54.84	41.13	22.32	13.33
Hg <sup>2+</sup>	400	349.32	326.68	198.80
Pb <sup>2+</sup>	550	469	396.55	198.83
Al <sup>3+</sup>	9.73	8.21	6.99	1.65

the metal ions were recovered successfully. The percentage recovery of Ba<sup>2+</sup> ions remained almost constant (>88%) upon increasing the loading of Ba<sup>2+</sup> ions (Table 7). These results confirm the high efficiency of the GAgNPs–SnZrMoP nanocomposite column.

Set 1: Ba<sup>2+</sup> (1.37 mg), Co<sup>2+</sup> (0.59 mg), Ni<sup>2+</sup> (0.58 mg), Ca<sup>2+</sup> (0.20 mg), Mg<sup>2+</sup> (0.24 mg), Fe<sup>3+</sup> (0.56 mg).

Set 2: Ba<sup>2+</sup> (2.74 mg) and the amounts of rest of metals were kept same as in Set 1.

Set 3: Ba<sup>2+</sup> (4.11 mg) and the amounts of rest of metals were kept same as in Set 1.

Similarly, the selective separation of Sr<sup>2+</sup> ions was achieved on the column of GAgNPs–SnZrMoP, from the following binary synthetic mixtures of metal ions: Ca(II)–Sr(II); Mg(II)–Sr(II); Co(II)–Sr(II); Cu(II)–Sr(II); Ni(II)–Sr(II), and the recovery of Sr(II) was more than 89% (Table 8).

Set 1: Sr<sup>2+</sup> (0.59 mg), Zn<sup>2+</sup> (0.65 mg), Ni<sup>2+</sup> (0.58 mg), Mg<sup>2+</sup> (0.24 mg), Cu<sup>2+</sup> (0.63 mg).

Set 2: Sr<sup>2+</sup> (1.18 mg), and the amounts of rest of metals were kept same as in Set 1.

Set 3: Sr<sup>2+</sup> (1.77 mg), and the amounts of rest of metals were kept same as in Set 1.

## Applications

**Removal of Ba(II) ions from wastewater samples.** From the above observations, it is evident that the synthesized material is

Table 6 Quantitative separation of metal ions from binary mixtures using the GAgNPs–SnZrMoP column

Material	Separations achieved	Separation factor	Eluent used	Metal ions (mg)			Efficiency (%)	Error (%)	Volume of eluent required (mL)
				Loaded	Eluted <sup>a</sup>	Efficiency (%)			
GAgNPs–SnZrMoP	Cu(II)	56.30	0.1 M HNO <sub>3</sub>	2.35	2.15	91.49	–8.51	60	
	Ba(II)		0.5 M HNO <sub>3</sub>	22.52	20.40	90.59	–9.41	50	
	Ca(II)	25.25	0.1 M HNO <sub>3</sub>	2.16	1.78	82.40	–17.6	60	
	Ba(II)		0.5 M HNO <sub>3</sub>	22.52	20.64	91.65	–8.35	40	
	Mg(II)	55.59	0.1 M HNO <sub>3</sub>	1.31	1.23	93.89	–6.11	40	
	Ba(II)		0.5 M HNO <sub>3</sub>	22.52	20.46	90.85	–9.15	60	
	Fe(III)	25.36	0.1 M HNO <sub>3</sub>	6.03	5.50	91.21	–8.79	30	
	Ba(II)		0.5 M HNO <sub>3</sub>	22.52	20.74	92.04	–7.96	60	
	Ni(II)	117.14	0.1 M HNO <sub>3</sub>	1.64	1.46	89.02	–10.98	40	
	Ba(II)		0.5 M HNO <sub>3</sub>	22.52	20.60	91.47	–8.53	60	
	Hg(III)	26.35	0.1 M HNO <sub>3</sub>	16.04	14.10	87.91	–12.09	60	
	Ba(II)		0.5 M HNO <sub>3</sub>	22.52	20.25	89.92	–10.08	60	

<sup>a</sup> An average of three replicate determinations.

Table 7 Selective separation of Ba<sup>2+</sup> from a synthetic mixture of Ba<sup>2+</sup>, Co<sup>2+</sup>, Ni<sup>2+</sup>, Ca<sup>2+</sup>, Mg<sup>2+</sup>, and Fe<sup>3+</sup> ions on the column of GAgNPs–SnZrMoP

Metal ions	Set no.	Amount loaded (mg)	Amount recovered <sup>a</sup> (mg)	Recovery (%)	Error (%)	Volume of eluent used (mL)	Eluent used
Ba <sup>2+</sup>	1	1.37	1.21	88.32	–11.68	60	0.5 M HNO <sub>3</sub>
	2	2.74	2.45	89.42	–10.58	60	0.5 M HNO <sub>3</sub>
	3	4.11	3.684	89.54	–10.46	60	0.5 M HNO <sub>3</sub>

<sup>a</sup> Average of three replicate determinations.



**Table 8** Selective separation of Sr<sup>2+</sup> ions from a synthetic mixture of Ca<sup>2+</sup>, Mg<sup>2+</sup>, Co<sup>2+</sup>, Cu<sup>2+</sup>, and Ni<sup>2+</sup> ions on the column of GAgNPs–SnZrMoP

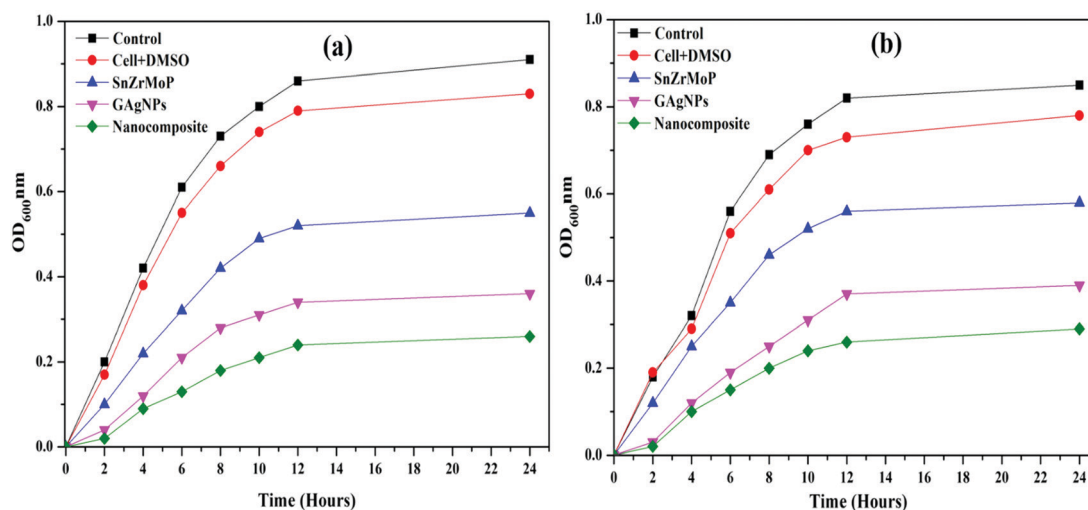
Metal ions	Set no.	Amount loaded (mg)	Amount recovered <sup>a</sup> (mg)	Recovery (%)	Error (%)	Volume of eluent used (mL)	Eluent used
Sr <sup>2+</sup>	1	0.59	0.53	89.83	−10.17	60	0.5 M HNO <sub>3</sub>
	2	1.18	1.07	90.67	−9.33	60	0.5 M HNO <sub>3</sub>
	3	1.77	1.62	91.52	−8.48	60	0.5 M HNO <sub>3</sub>

<sup>a</sup> Average of three replicate determinations.

**Table 9** Separation of Ba<sup>2+</sup> ions from wastewater samples collected from glass, paint, and brick-making industries, and a chemistry lab on a column of the GAgNPs–SnZrMoP nanocomposite

Source of wastewater sample	Metal ions	Eluent used	Metal ions obtained		
			Amount loaded (mg)	Amount <sup>a</sup> eluted (mg)	Recovery (%)
Glass industry	Ba <sup>2+</sup>	0.5 M HNO <sub>3</sub>	1.62	1.45	89.50
Paint industry		0.5 M HNO <sub>3</sub>	1.55	1.38	89.03
Brick-making industry		0.5 M HNO <sub>3</sub>	1.73	1.56	90.17
Chemistry laboratory		0.5 M HNO <sub>3</sub>	0.85	0.75	88.24

<sup>a</sup> Average of three replicate determinations.

**Fig. 8** Growth curves of (a) *E. coli*; (b) *Bacillus* bacteria.

an efficient candidate for the effective removal of Ba<sup>2+</sup> ions from wastewater samples collected from different sources. The nanocomposite was used for the removal of Ba(II) ions from wastewater samples collected from different industrial units. The Ba<sup>2+</sup> ions were quantitatively separated and recovered from wastewater containing 1.45, 1.38, 1.56, and 0.75 mg of Ba(II) ions in effluents obtained from glass, paint, and brick-making industries, and from a chemistry lab, respectively (Table 9).

### Antimicrobial study

It was revealed from the antibacterial studies that all three materials inhibited the growth of both G<sup>+</sup> and G<sup>−</sup> bacterial cultures to an astonishing level in the order: GAgNPs–SnZrMoP > GAgNPs > SnZrMoP. However, both the bacteria showed regular growth under the control experiment conditions (Fig. 8). The inhibition in the growth of both G<sup>+</sup> and G<sup>−</sup> bacterial cultures might be due to strong interactions of the nanocomposite with the

cell membrane due to which the cellular and molecular functions of the cell are interrupted by interfering with its permeability. This may finally result in a crumbling of the layer structure that ultimately causes the death of the cells.<sup>31</sup>

## Conclusion

In the present study, a Ba(II) selective nanocomposite cation exchanger GAgNPs–tin zirconium(IV) molybdophosphate was synthesized by a sol–gel method, which had a good ion-exchange capacity (2.48 meq g<sup>−1</sup>) as compared to its inorganic counterpart tin-zirconium(IV) molybdophosphate (1.95 meq g<sup>−1</sup>). From the HR-TEM, interactions between the green silver nanoparticles and polybasic acid salt (tin-zirconium molybdophosphate) are very clear and are responsible for its high thermal and chemical stability. It is clear that the particle sizes were in the range of 10–20 nm. The nanohybrid cation exchanger, GAgNPs–SnZrMoP





was successfully used for inhibition of the growth of both bacterial cultures, *i.e.*, *E. coli* (Gram (–) ve) and *Bacillus* (Gram (+) ve). The nanohybrid cation exchanger, GAgNPs–SnZrMoP had better antibacterial properties as compared to the inorganic ion exchanger (SnZrMoP) and metal nanoparticles (GAgNPs). On the basis of distribution coefficient studies, some analytically important binary separations were also performed. The recovery of metal ions was very sharp, and more than 88% of metals was recovered successfully. On the basis of the above results, it can be concluded that the nanocomposite ion exchanger GAgNPs–SnZrMoP can be used as an efficient candidate for environmental remediation.

## Conflicts of interest

There are no conflicts to declare.

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