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Synthesis of novel mesoporous silica nanoparticles functionalized with succinic dihydrazone Schiff-base metal complexes and a study of their biological activities

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For the first time, functionalized mesoporous silica nanoparticles (MSNs) with dinuclear Schiff-base complexes were synthesized as attractive organic–inorganic hybrids and their capability investigated for loading antibiotic drugs and immobilization of enzymes. 2D-hexagonal MCM-41 nanoparticles were synthesized by a sol–gel method and amino propyl trimethoxy silane (APTMS) anchored on the surface of the MSNs as a linker. MSN-APS nanoparticles were coordinated with dihydrazone Schiff-base complexes of copper(II) and nickel(II) by the nitrogen atom of APS for making MSN-APS-Cu₂L and MSN-APS-Ni₂L hybrids, respectively (L is the Schiff-base ligand). These novel mesoporous silica nanoparticles were characterized by various techniques, such as FT-IR, LA-XRD, FE-SEM, TEM, EDX, BET and TGA. The results show that the synthesized hybrids have high potential for loading gentamicin and immobilizing enzymes such as DNase, coagulase and α -amylase.

Introduction

Mesoporous materials with a pore size of 2–50 nm have attracted the attention of various scientists since their discovery in 1992.^{1–3} About a decade later, other mesoporous silica nanoparticles were also synthesized and studied.⁴ The remarkable characteristics of these new materials are: controllable pore and particle size, stable suspension for long periods, high surface area, mechanical and chemical stability and low toxicity.^{4–7} Mesoporous silica nanoparticles (MSNs) are divided into five categories: 2D-hexagonal MCM-41 (MCM-41s), swollen pore MCM-41 (SMCM-41s), hollow mesoporous nanoparticles (HNPs), rod-like MCM-41 (RMCM-41s) and radial mesoporous silica nanoparticles (RNPs).^{8–12} Among them, MCM-41 with a particle size of 50–200 nm has many interesting features, such as biocompatibility, stability, good interactions with many drugs for use as antibacterial agents, ease of functionalization and high surface area.^{13–16} In recent years, organic groups or inorganic compounds (in the form of metal complexes) were grafted on the surface of MSNs. Metal complexes supported on

MSNs are utilized in various fields, such as catalysis, drug delivery, fluorescence, pollutant removal, photodynamic therapy and tooth bleaching, as well as enzyme immobilization.^{9,17–28} Metal complexes of Ni²⁺, Co²⁺ and Cu²⁺ have been grafted on carbon nanotubes, quantum dots and various nanoparticles, such as Au, Fe₂O₃ and SiO₂.^{29–32} Grafting of metal complexes on nanoparticles has resulted in an increase in the stability and lifetimes of the enzymes.³³ Among the ligands and complexes, bis-acyl-hydrazones and their metal complexes have shown excellent properties for catalytic, anti-cancer, antibacterial and antifungal activities.^{34–51} Therefore, with regard to the importance of MSNs, bis-acyl-hydrazones and their complexes, in this research work we report the synthesis of new organic–inorganic hybrids and their characterization, and an investigation of the capability of these compounds to show biological activities. 3-(Aminopropyl) triethoxysilane (APTMS) was functionalized on MSN and grafted onto bis-acyl-hydrazone Schiff-base complexes of Ni²⁺ and Cu²⁺. These novel hybrids, including binuclear metal complexes grafted on the surface of MSN through linkers, have exhibited attractive properties, such as high surface area, biocompatibility, thermal and mechanical stability, and a high number of active sites (metallic centers) for stabilizing enzymes and loading therapeutic agents. The ability of the new hybrids to load gentamicin, and their antibacterial activities as well as the immobilization of α -amylase, DNase and coagulase enzymes were investigated.

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Highlight

Experimental

Synthesis of bis(2-hydroxybenzaldehyde) succinic dihydrazone(H_4L)

1 mmol (0.146 g) of succinic dihydrazide was dissolved in 25 mL of absolute ethanol in a flask and heated and stirred for 30 min. Then 0.3 mL (2 mmol) of salicylaldehyde was slowly added to the solution. The mixture was refluxed for 3 h. The obtained white precipitate was filtered, washed with absolute ethanol and dried in an oven at 80 °C for 5 h. This compound was characterized by FT-IR.

Synthesis of bis-acyl-hydrazone complexes Cu_2L and Ni_2L

A mixture of 0.28 mmol (0.1 g) of H_4L and 25 mL of absolute ethanol was added to a flask and heated and stirred for 30 min. Then 0.56 mmol of acetate salts of Cu^{2+} and Ni^{2+} was dissolved in ethanol. The solution of salts was added to the ligand solution to obtain Cu_2L and Ni_2L . The mixtures were refluxed for 3 h. The obtained green (Cu_2L) and brown (Ni_2L) precipitates were filtered, washed with absolute ethanol and dried at room temperature. These compounds were characterized with FT-IR.

Synthesis of MSNs and MSN-APS

1 g of CTAB was dissolved in 480 mL of deionized water and stirred vigorously. Then 3.5 mL of 2M NaOH was added to this solution and it was heated at 80 °C for 4 h. 5 mL of TEOS was added dropwise into the solution and stirred for 2 h at 80 °C to form a white gel. This white gel was collected, washed several times with deionized water and dried at 80 °C in an oven overnight. The obtained MNS product was dispersed in acidic ethanol and refluxed for 24 h. To prepare MSN-APS, 0.1 g of pre-synthesized MSN was dispersed in 80 mL of ethanol, and 1 mL of 3-aminopropyl triethoxysilane (APTMS) was slowly added to this suspension and stirred. This mixture was refluxed for 3 h. Then it was filtered, thoroughly washed with ethanol and dried in an oven at 80 °C for 12 h. The MSNs were characterized by FT-IR, FE-SEM, EDX, LA-XRD, BET.

Synthesis MSN-APS- Cu_2L and MSN-APS- Ni_2L

0.1 g of MSN-APS was dispersed in 20 mL of absolute ethanol and stirred for 30 min. A solution containing 0.1 mmol of Schiff-base complexes Cu_2L (0.044 g) or Ni_2L (0.045 g) was sonicated in 10 mL of absolute ethanol for 10 min and gradually added to the MSN-APS solution. The mixture was stirred at room temperature for 12 h. The nanoparticles were filtered and washed several times with ethanol until the solution under filtration became colorless. The collected products were dried in an oven at 80 °C for 24 h. Different stages of the synthesis of hybrid mesoporous silica nanoparticles are shown in Fig. 1. MSN-APS- Cu_2L and MSN-APS- Ni_2L nanoparticles were characterized by FT-IR, FE-SEM, TEM, EDX, LA-XRD, BET and TGA techniques.

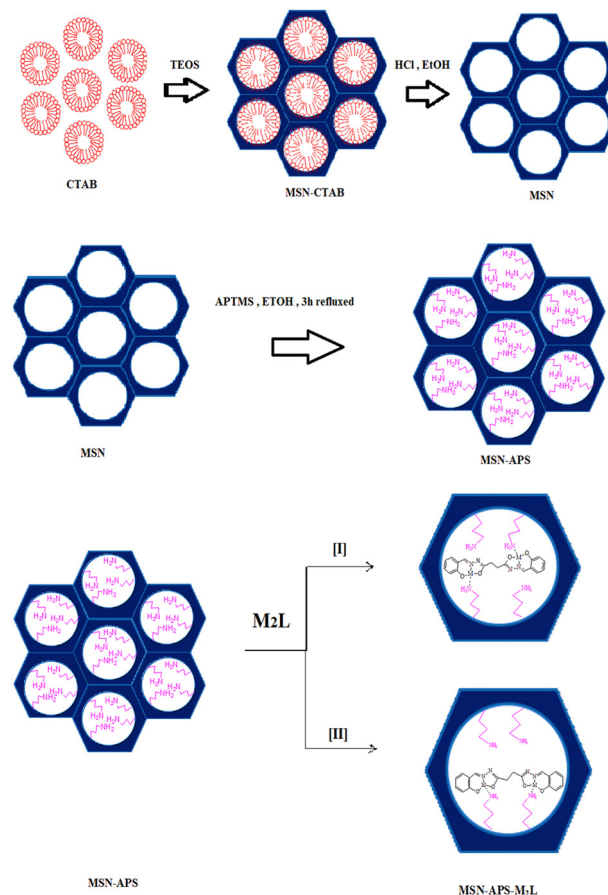


Fig. 1 Schematic of synthesized mesoporous silica nanoparticles.

Biological experiments

Several microbial tests and enzyme immobilization were accomplished in this research work. In the first part the intrinsic antibacterial activity of MSN-APS- Cu_2L and MSN-APS- Ni_2L was registered against *Escherichia coli* (ATCC 25922) and *Staphylococcus aureus* (ATCC 6538) bacteria in Mueller–Hinton broth culture medium. 100 μ L of bacterial suspension that had grown until 0.5 McFarland turbidity was added to 20 mg of each compound in 5 mL of Mueller–Hinton broth and incubated at 37 °C (Merck, Germany). Simultaneously, a culture was prepared from each bacterial species without nanoparticles and a culture was prepared with nanoparticles without bacterial inoculation. The optical density of the different samples was measured with an ELISA microplate reader at 665 nm over 24 h. In the second part the loading of MSN, MSN-APS, MSN-APS- Cu_2L and MSN-APS- Ni_2L nanoparticles was examined with gentamicin antibiotic and their inhibitory activities were investigated against four bacterial species. For this test, 10 mg of gentamicin was dissolved in 1 mL of MQ water and added to 20 mg of nanoparticles. For proper loading, samples were kept in an incubator for 24 h at 37 °C. After three washings of the nanoparticles with MQ-water, the antibacterial activity of the nanocomposites and Gen@nanocomposites was investigated against Gram-positive bacteria, i.e., *Staphylococcus aureus*.



(ATCC 6538) and *Bacillus subtilis* (ATCC 6633), and two Gram-negative species, i.e., *Escherichia coli* (ATCC 25922) and *Pseudomonas aeruginosa* (ATCC 9027) according to the standard Kirby-Bauer disc diffusion method recommended by CLSI guidelines. In the third part, the capacity of enzyme immobilization of these new organic-inorganic hybrids was studied. Three enzymes were used for this examination: α -amylase (Sigma Company), coagulase and DNase (from *S. aureus*). The concentration of α -amylase was 2 mg in 1 mL of PBS and those of coagulase and DNase were 1 mL with respect to 20 mg of each composite. In case of α -amylase, a solution of pure enzyme (2 mg mL⁻¹ PBS) was prepared and 20 mg of each nanocomposite was added to 1 mL of this solution and incubated at 37 °C for 24 h. These nanoparticles were then washed three times with PBS and dispersed in 500 μ L sterile distilled water. Finally, the enzyme loadings on these compounds were evaluated through culturing on starch agar, a plasma coagulation test and clear halo zone formation on DNase agar.

Results and discussion

FT-IR spectroscopy

The FT-IR spectra of H₄L, Cu₂L, Ni₂L, MSN-APS-Cu₂L and MSN-APS-Ni₂L are shown in Fig. 2. Information about the FT-IR spectra of MSN and MSN-APS was reported in our earlier article.^{52,53} In the FT-IR spectra of H₄L appeared bands of -OH (3428 cm⁻¹), -NH (3201 cm⁻¹), aromatic -CH (3054 cm⁻¹) and aliphatic -CH (2857–2928 cm⁻¹). Also the bands of C=O and C=N were observed at 1662 cm⁻¹ and 1622 cm⁻¹.^{54,55} In the IR spectra of binuclear Schiff-base complexes Cu₂L and Ni₂L, carbonyl bands had vanished and the imine groups had decreased to 1618 cm⁻¹ and 1606 cm⁻¹, respectively, because of coordination of the imine of the ligand

to transition metals ions. In the Cu₂L spectra, the bands at 491 cm⁻¹ and 597 cm⁻¹ were related to the vibrations of Cu-N and Cu-O. The vibrations for Ni₂L were observed at 525 cm⁻¹ and (Ni-N) 616 cm⁻¹ (Ni-O). The FT-IR spectra of organic-inorganic hybrids of MSN-APS-Cu₂L and MSN-APS-Ni₂L showed that the MSN-APS and binuclear Schiff-base complexes were grafted together, because the bands of the silanol groups (Si-O-Si) at 1085, 802, 465 cm⁻¹ and the imine of bis-acyl-hydrazone complexes were seen for MSN-APS-Cu₂L and MSN-APS-Ni₂L.

FE-SEM and TEM

As shown in the FE-SEM images, MSN-APS-Cu₂L and MSN-APS-Ni₂L had monotonous spherical morphology and their mean diameter was estimated as approximately less than 100 nm by ImageJ software (Fig. 3(a)–(c)). Also, the spherical morphology of the initial MSN was preserved for these organic-inorganic nanohybrids, which demonstrated that the Schiff-base complexes had settled on the surface of the nanoparticles or on the surface of channel-like pores. Mesoporous channels were clearly observed in the TEM images of MSN-Cu₂L, with a hexagonal ordered arrangement. Obvious channel-like pores parallel to each other can be seen (Fig. 4).

EDX

The elemental content of MSN, MSN-APS-Cu₂L and MSN-APS-Ni₂L was investigated by using energy dispersed X-ray (EDX) spectroscopy. The EDX spectral patterns of the nanoparticles are shown in Fig. 5: MSN (Si: 43.64% and O: 56.36%), MSN-APS-Cu₂L (Si: 14.61%, O: 48.71%, N: 7.62%, C: 28.50% and Cu: 1.26%) and MSN-APS-Ni₂L (Si: 16.91%, O: 41.41%, N: 6.13%, C: 35.14% and Ni: 0.40%). The EDX results indicate the presence of the expected elements in the structure of the nanoparticles and prove their formation.

XRD

The low-angle powder X-ray diffraction in the 0.8° < 2 θ < 10° range for MSN-APS-Cu₂L and MSN-APS-Ni₂L can be seen in Fig. 6. In our previous work, the patterns for MSNs had four reflection peaks with different intensities that indicate Bragg peaks by indexing planes (100), (110), (200) and (210) at 2 θ = 2.27°, 3.88°, 4.50°, and 5.94°, respectively.⁵² This XRD pattern exhibited a hexagonal arrangement of pores and the 2D-MCM-41 structure of the nanoparticles.⁵⁶ The XRD patterns of MSN-APS-Cu₂L and MSN-APS-Ni₂L compared with MSN show two peaks at 2 θ = 2.01° and 2.11° (high intensity) and at 4.2° and 4.08° (low intensity), while the other peaks had vanished because the mesoporous channels were filled with binuclear Schiff-base complexes as well as a decreased order of mesopores (Fig. 6(a) and (b)). The position of index peak (100) was used to calculate the interplanar spacings d_{100} for the materials. The d_{100} values were used to calculate the distances between pore centers a_0 ($a_0 = \frac{2d_{100}}{\sqrt{3}}$). The calculated values of a_0 for MSN, MSN-APS-Cu₂L and MSN-APS-Ni₂L are 4.48 nm, 5 nm and 4.82 nm, respectively. The values of FWHM were

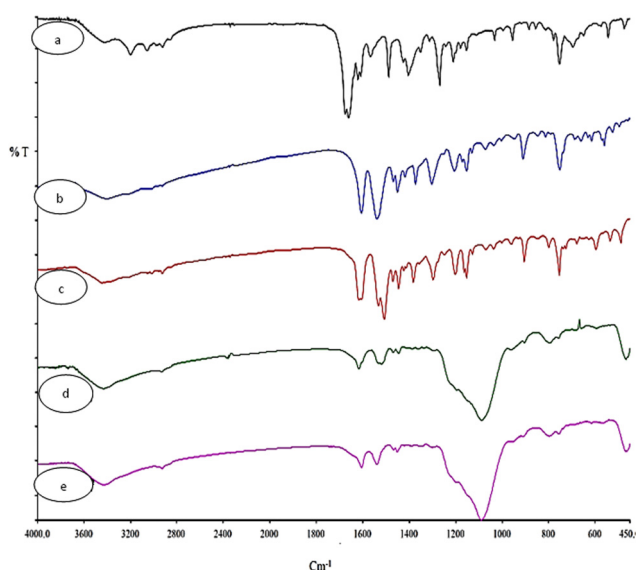


Fig. 2 FT-IR spectra of synthesized nanohybrids (a) L, (b) Ni₂L, (c) Cu₂L, (d) MSN-APS-Cu₂L and (e) MSN-APS-Ni₂L.



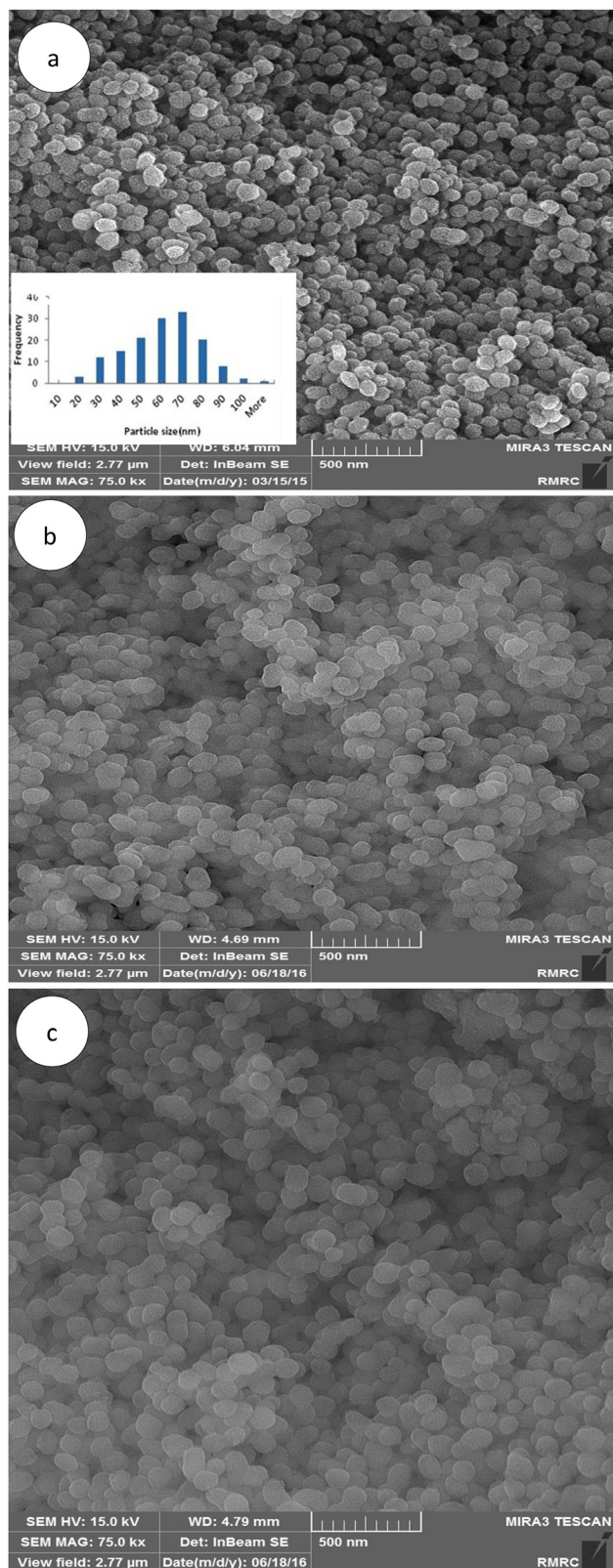


Fig. 3 FE-SEM images of the MSNs and functionalized MSNs: (a) MSN, (b) MSN-APS-Cu₂L and (c) MSN-APS-Ni₂L.

extracted from the curves and used to calculate the particle size of the crystallite (D). According to the Debye-Scherrer equation,

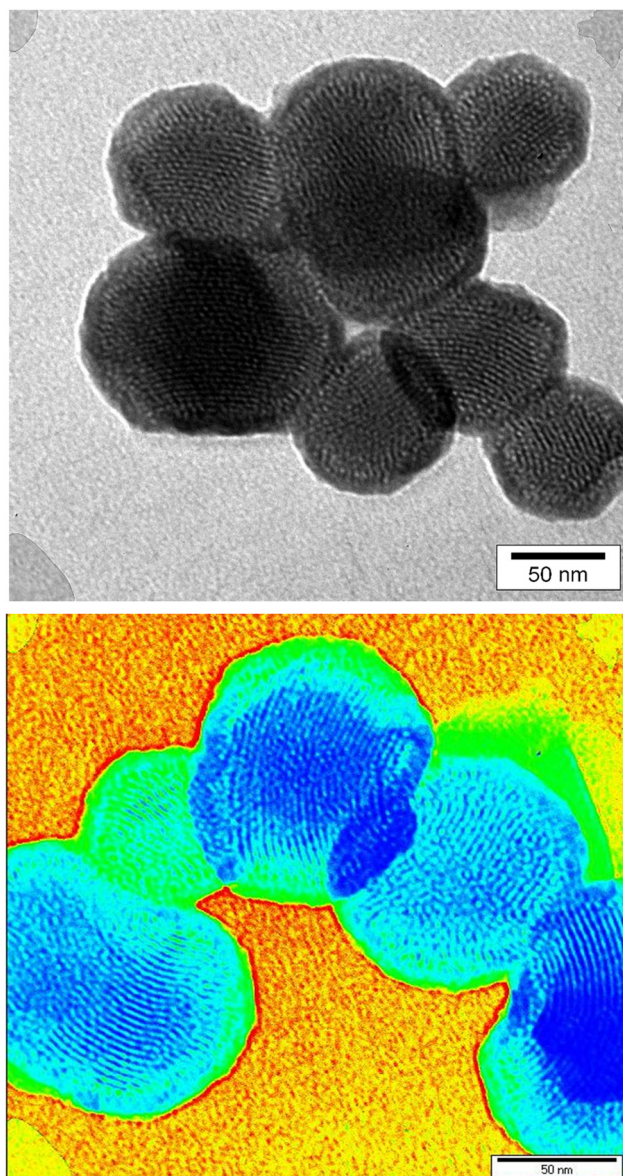


Fig. 4 TEM images of MSN-APS-Cu₂L.

the D value for MSN-APS-Cu₂L is 23.89 nm and that for MSN-APS-Ni₂L is 24.31 nm.

TGA

According to the thermogravimetric curves of MSN-APS-Cu₂L and MSN-APS-Ni₂L two main weight loss steps were observed from 25 °C to 800 °C at a heating rate of 10 °C min⁻¹ (Fig. 7). The first step of decreasing weight was related to loss of physical water or organic solvents on the surface. The second step (22% loss weight) was attributed to the decomposition of the organic linker and the organic parts of binuclear Schiff-base complexes on MSNs from 250 to 450 °C. The amount of binuclear complexes grafted on MSN was estimated as 0.53 mmol g⁻¹. The residual mass up to 450 °C is related to the inorganic parts, including metals and silica components.



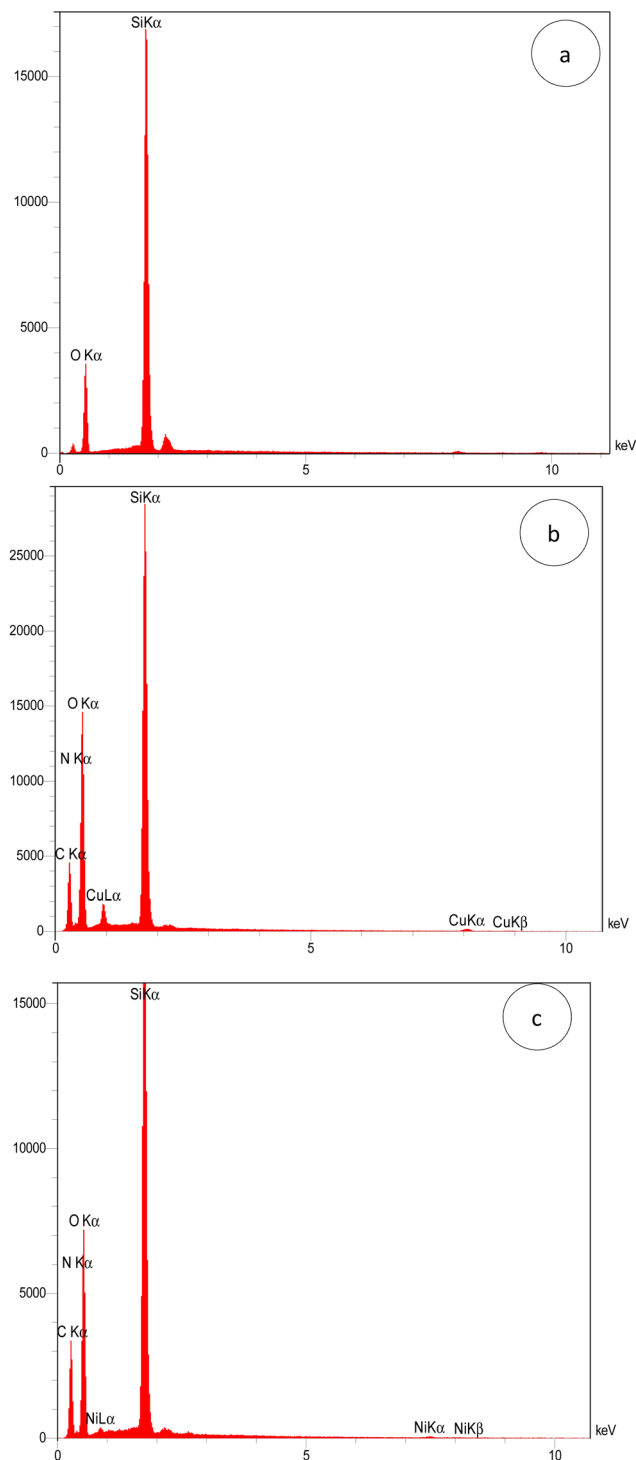


Fig. 5 EDX patterns of (a) MSN, (b) MSN-APS-Cu₂L and (c) MSN-APS-Ni₂L.

BET

The isotherms of N₂ absorption-desorption for MSN, MSN-APS-Cu₂L and MSN-APS-Ni₂L are displayed in Fig. 8. The patterns of these nanoparticles exhibit type IV isotherms, as expected for mesoporous silica with cylindrical and very uniform pores. The sharp rise at a relative pressure of 0.2–0.4 showed that the pores of the nanoparticles are very narrow. The specific surface area

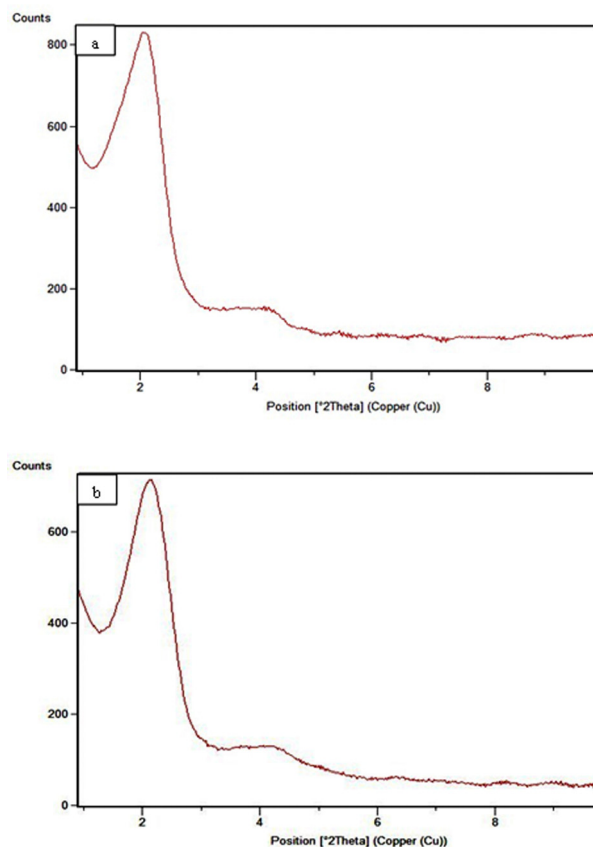


Fig. 6 Low-angle XRD patterns: (a) MSN-APS-Cu₂L and (b) MSN-APS-Ni₂L.

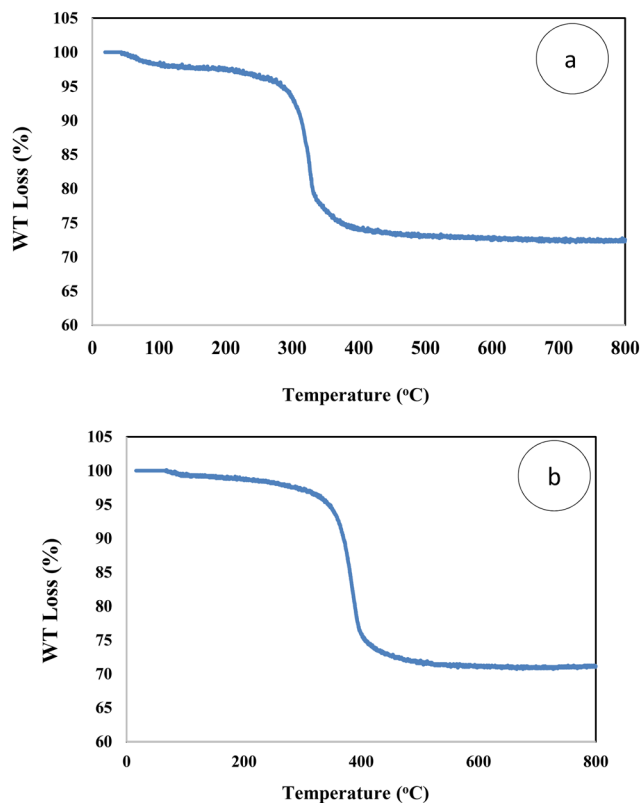


Fig. 7 TGA curves of (a) MSN-APS-Cu₂L and (b) MSN-APS-Ni₂L.



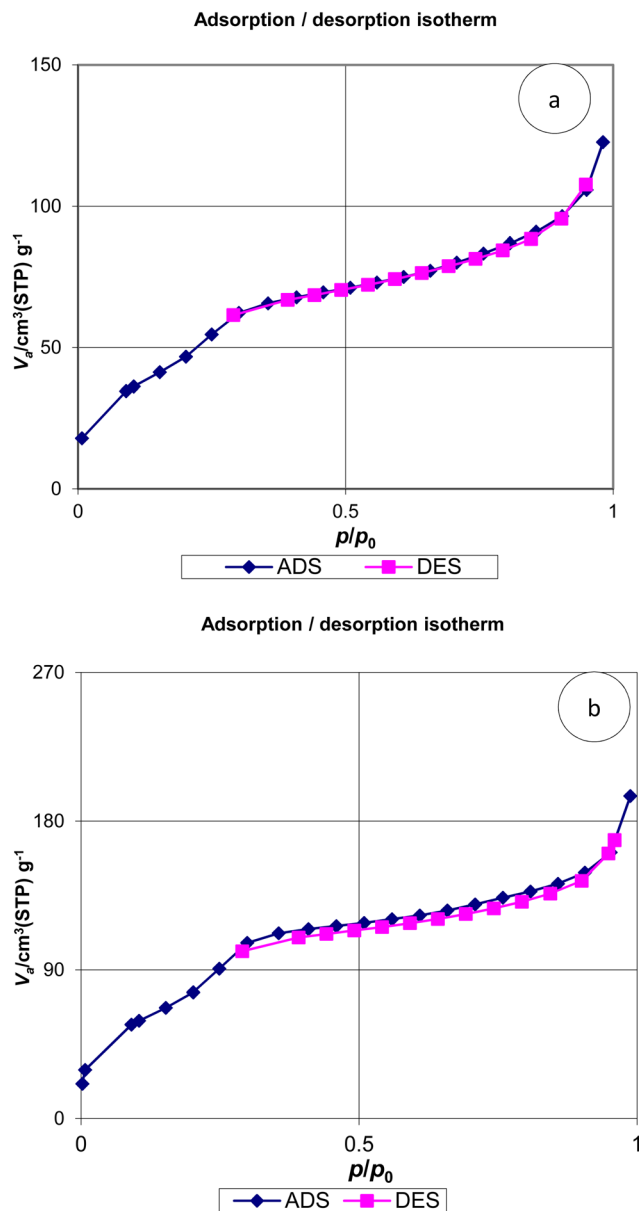


Fig. 8 N_2 adsorption-desorption isotherms of (a) MSN-APS- Cu_2L and (b) MSN-APS- Ni_2L .

(s), pore volume (v_p) and pore radius (r_p) of the nanoparticles were calculated by the BET and BJH methods. The data obtained from these methods for MSN-APS- Cu_2L were $s = 179.34 \text{ m}^2 \text{ g}^{-1}$, $v_p = 0.174 \text{ cm}^3 \text{ g}^{-1}$, and $r_p = 1.21 \text{ nm}$ and for MSN-APS- Ni_2L were $s = 305.6 \text{ m}^2 \text{ g}^{-1}$, $v_p = 0.278 \text{ cm}^3 \text{ g}^{-1}$, and $r_p = 1.21 \text{ nm}$. The values of v_p and r_p decreased compared to those of MSN ($s = 789 \text{ m}^2 \text{ g}^{-1}$, $v_p = 0.782 \text{ cm}^3 \text{ g}^{-1}$, $r_p = 1.21 \text{ nm}$) because of the anchoring of binuclear Schiff-base complexes on the surface of the pores of MSNs.

Biological activity

In this work, the new organic-inorganic hybrids were studied for their antibacterial activity, gentamicin loading potential and enzyme immobilization. Investigation of the growth of treated

Table 1 Antibacterial effect of synthesized nanohybrids

Compounds	Inhibition zone (mm)			
	<i>S. aureus</i>	<i>B. subtilis</i>	<i>E. coli</i>	<i>P. aeruginosa</i>
MSN	10 ^a	0	0	10 ^a
MSN-APS	10 ^a	0	0	10 ^a
MSN-APS- Cu_2L	10 ^b	13 ^b	0	13 ^a
MSN-APS- Ni_2L	10 ^a	10 ^a	0	10 ^a
MSN@Gen	10 ^b	9 ^b	10 ^b	8 ^b
MSN-APS@Gen	13 ^b , 15 ^a	15 ^b	16 ^b	10 ^b
MSN-APS- Cu_2L @Gen	20 ^b	19 ^b	20 ^b	15 ^b
MSN-APS- Ni_2L @Gen	15 ^b	17 ^b	20 ^b	18 ^b

^a Inhibitory effect. ^b Bactericidal effect.

bacteria showed that MSN-APS- Cu_2L had bactericidal effects against *S. aureus* and *B. subtilis*. The effect of gentamicin-loaded mesoporous nanoparticles was discovered by a disc diffusion method. The results of this assay for nanoparticles alone and nanoparticles@Gen are presented in Table 1. MSN, MSN-APS- Cu_2L and MSN-APS- Ni_2L loaded gentamicin had more antibacterial activity than MSN or MSN-APS because, due to their bis-hydrazone groups and active metal ion centers, they have good interaction with the $-NH_2$ and $-OH$ groups of gentamicin.⁵⁶ The results of the inhibition zone of composites and nanoparticles@Gen against *S. aureus*, *B. subtilis*, *E. coli* and *P. aeruginosa* are presented in Table 1 and Fig. 9 shows the effect of antibacterial activity of nanohybrids@Gen against *B. subtilis* and its comparison with MSN@Gen and MSN-APS@Gen. The results of enzyme immobilization for these compounds are presented in Tables 2 and 3. MSN-APS- Cu_2L and MSN-APS- Ni_2L showed glorious potential for immobilization of coagulase, DNase and α -amylase. Fig. 10, for example, shows a large clear zone around MSN-APS- Ni_2L @Am on starch agar, which means efficient immobilization of amylase and its diffusion in the culture medium. Also, coagulase-loaded MSN-APS- Cu_2L and MSN-APS- Ni_2L nanoparticles caused plasma clot formation over 1–2 h following addition to plasma. The good loading and stabilization of these enzymes on MSN-APS- Cu_2L and MSN-APS- Ni_2L were due to the interaction of the $-NH_2$ and $-COO^-$

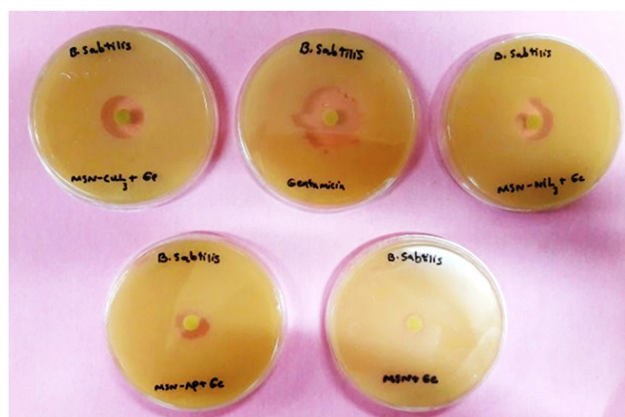


Fig. 9 Comparison of the antibacterial effects of MSN-APS- Ni_2L @Gen (top right), MSN-APS- Cu_2L (top left), gentamicin (top middle), MSN@Gen (bottom right) and MSN-APS@Gen (bottom left).



Table 2 The effect of loading of coagulase enzyme

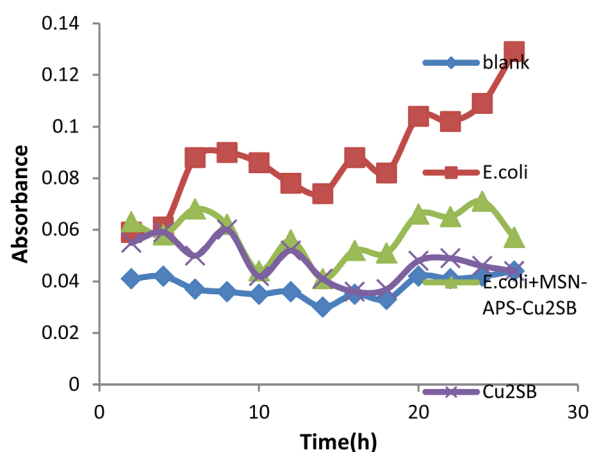
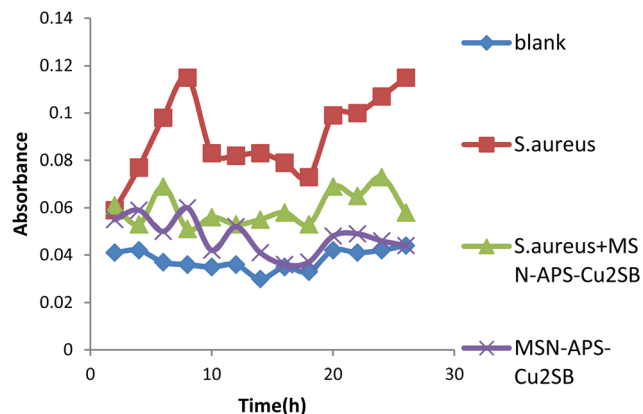
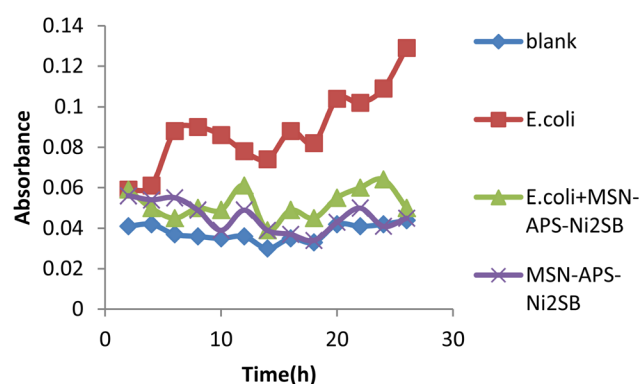
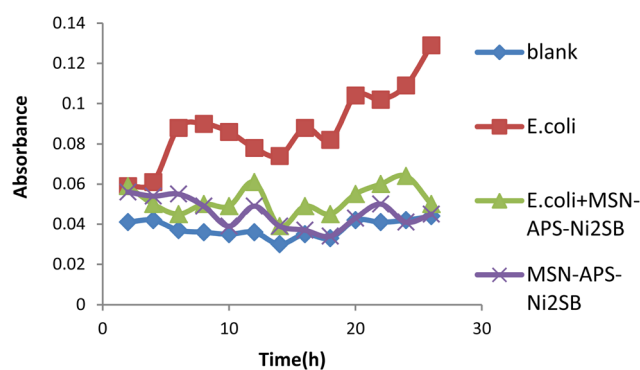
hour	MSN@Coag	MSN-APS@Coag	MSN-APS-Cu ₂ L@Coag	MSN-APS-Ni ₂ L@Coag
1–2	–	–	+	+
18–24	+	+	+	+

Table 3 The effect of loading DNase enzyme

	MSN@Am	MSN-APS@Am	MSN-APS-Cu ₂ L@Am	MSN-APS-Ni ₂ L@Am
Effect zone	0	0	47	43

Fig. 10 Effect zone of MSN-APS-Ni₂L@Am.

groups of the enzymes with the empty coordination sites of metal complexes on MSN.^{56–62} These hybrid nanoparticles also had greater effects as the bishydrazone Schiff-base ligands and binuclear complexes efficiently increased enzyme immobilization on MSN. Fig. 11–14 show that new MSNs functionalized by metal dinuclear compounds had a killing effect against *E. coli* and *S. aureus* bacteria in Mueller–Hinton broth. All these

Fig. 11 Antibacterial activity of MSN-APS-Cu₂L against *E. coli* in Mueller–Hinton broth medium.Fig. 12 Antibacterial activity of MSN-APS-Cu₂L against *S. aureus* in Mueller–Hinton broth medium.Fig. 13 Antibacterial activity of MSN-APS-Ni₂L against *E. coli* in Mueller–Hinton broth medium.Fig. 14 Antibacterial activity of MSN-APS-Ni₂L against *S. aureus* in Mueller–Hinton broth medium.

microbial tests determined the importance and good potential applications of these new hybrids of MSNs.

Conclusions

New mesoporous silica nanoparticles with universal formulae MSN-APS-Cu₂L and MSN-APS-Ni₂L were synthesized by a



grafting method with success. These mesoporous silica nanoparticles are made from three important parts: MCM-41 nanoparticles, APTMS (linker) and Cu₂L and Ni₂L (binuclear Schiff-base complexes). These novel functionalized MSNs with dihydrazone complexes of Cu²⁺ and Ni²⁺ had interesting properties, such as keeping the particle sizes of the initial support, having a high surface area, suitable pore size, and metallic active sites for interaction with gentamicin antibiotic and DNase, coagulase and α -amylase enzymes. The antibacterial activity of MSN-APS-Cu₂L and MSN-APS-Ni₂L showed a bactericidal effect against *E. coli* and *S. aureus*. These compounds carried gentamicin antibiotic well and compared to MSN and MSN-APS showed better operation of loaded gentamicin. The results of enzyme immobilization showed that these two novel nanoparticles also stabilized DNase, coagulase and α -amylase.

Conflicts of interest

There are no conflicts to declare.

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