



Facile synthesis of new composite, Ag-Nps-loaded core/shell CdO/Co₃O₄ NPs, characterization and excellent performance in antibacterial activity

A. M. Nassar^{1,2} · Z. A. Alrowaili³ · Ahmed A. M. Ahmed^{4,5} · B. A. Cheba^{6,7} · Sultan Akhtar⁸

Received: 11 August 2020 / Accepted: 23 October 2020
© King Abdulaziz City for Science and Technology 2020

Abstract

Here, we report for the first time the synthesis of new composite Ag-NPs-loaded core/shell mixed oxides NPs, CdO/Co₃O₄. First, nano-precursors (1) and (2) consist of mixed (cadmium oxalate/cobalt oxalate) and (silver oxalate/cadmium oxalate/cobalt oxalate), respectively, have been synthesized and characterized. Then, thermal treatment of 1 and 2 at 400 °C resulted in the formation of composites CdO/Co₃O₄ (3) and Ag-NPs@CdO/Co₃O₄ (4), respectively. The materials have been characterized by IR, SEM, TEM, TGA and XRD. The morphologies of the materials were described from SEM images. TEM images revealing the formation of core-shell structure with average particles size 5–20 nm. The crystal sizes were calculated from XRD patterns and resulted in 14.21, 14.65, 15.06 and 15.79 nm for 1, 2, 3 and 4, respectively. All the materials undergo tests as antibacterial agents against Gram-positive bacteria (*Bacillus subtilis* and *Enterococcus spp.*) and Gram-negative bacteria (*E. coli* and *Pseudomonas aeruginosa*). Precursor 2 and composite 4 impart higher inhibition against all the bacterial strains compared to precursor 1 and composite 3.

Keywords Mixed oxides · Cadmium · Cobalt · Ag-NPs · Composite · Antibacterial

✉ A. M. Nassar
amnassar@ju.edu.sa

¹ Department of Chemistry, College of Science, Jouf University, Sakaka, Saudi Arabia

² Department of Chemistry, Faculty of Science, Al-Azhar University, Nasr-City, Cairo, Egypt

³ Department of Physics, College of Science, Jouf University, Sakaka, Saudi Arabia

⁴ Department of Basic Science, Common First Year Deanship, Jouf University, Sakaka, Saudi Arabia

⁵ Department of Chemistry, Faculty of Science, Cairo University, Giza, Egypt

⁶ Department of Biology, College of Science, Jouf University, Sakaka, Saudi Arabia

⁷ Department of Biotechnology, College of Nature and Life Sciences, University of Science and Technology of Oran Mohamed-Boudiaf (USTOMB), BP 1505 Al Mnaouar, 31000 Oran, Algeria

⁸ Department of Biophysics, Institute for Research and Medical Consultations (IRMC), Imam Abdulrahman Bin Faisal University, P. O. Box. 1982, Dammam 31441, Saudi Arabia

Introduction

Metal oxides nanomaterials are well-known components for different applications (Bayal and Jeevanandam 2012). The main attention of these materials resulted from their heightened electronic, optical, magnetic, chemical and physical properties than their bulk counter parts (Yuan et al. 2014). Mixed metal oxides are subjected in different areas such as superconductors (Williams et al. 1988), photocatalysts (Koohestani 2019), sensors (Zakrzewska 2001), ground-water remediation (Basu and Ghosh 2013), removal of air toxicity (Li et al. 2011) and pharmaceuticals (Abureesh et al. 2018).

As a significant n-type semiconductor metal-oxide, cadmium oxide (CdO) which has a direct band gap 1.3–2.5 eV (Balamurugan et al. 2016). CdO is included in several applications such as electrical, antibacterial, medical and optical fields (Manjula et al. 2015). Recently, a composite of glucose/cadmium oxide nanoparticles was fabricated and showed potential therapeutic value as an effective anti-glycation agent (Zahera et al. 2020). Gadolinium-doped cadmium oxide thin films were prepared and showed high-performance optoelectronic properties with high-transmittance in visible light (Sakthivel

et al. 2020). A composite of cadmium oxide@graphene was synthesized via a green route based on onion extracts. The nanocomposite imparted antibacterial activity higher than that of pure CdO NPs (Sadhukhan et al. 2019). A. Salem studied the fabrication of silver nanoparticles-doped cadmium oxide composite with face-centered cubic structure and its optical properties. The study demonstrated that the composite present a promising candidate for optoelectronic devices (Salem 2014).

Spinel cobalt oxide, Co_3O_4 is known as a magnetic p-type semiconductor material with unique shape (Salam et al. 2020). It is environment-friendly and has excellent chemical and physical stability (Vennela et al. 2019). It is widely studied as promising materials for several applications including the medical nanotechnology and biotechnology, photocatalytic degradation of organic pollutants, electrochemistry, lithium-ion batteries, gas sensors, and energy storage devices (Assim et al. 2017).

Loading of silver nanoparticles, Ag-NPs on the materials surface or doping in their crystalline matrix significantly impacts on the development of the materials physical and chemical properties (Nassar et al. 2018). Among the activities of Ag-NPs, their antimicrobial activity and high sterilization performance are widely well discussed (Ren et al. 2019; Gong et al. 2007; Kooti et al. 2013). The action mechanism of the Ag-NPs antibacterial activity explained by their high permeability through the bacteria cell wall membrane due to Ag-NPs has a small size as well as their large specific surface area (Liu et al. 2013).

According to the best of our knowledge, recently, only one work has reported the fabrication and characterization of CdO/ Co_3O_4 composite for the first time via a simple solvothermal method (Wang et al. 2020). In this work and based on the biological potential of CdO, Co_3O_4 , and Ag-NPs, our focus is to fabricate, characterize and study the antibacterial activity of mixed oxides CdO/ Co_3O_4 and its loaded silver nanoparticles novel composite Ag-NPs@CdO/ Co_3O_4 .

Experimental

Materials and reagents

The materials that were used in this work without involving any purification are cadmium nitrate tetrahydrate, cobalt nitrate hexahydrate, silver nitrate and oxalic acid dihydrate from Sigma-Aldrich (USA). Milli-Q water was used in all experiments.

Synthesis of mixed cadmium oxalate/cobalt oxalate (precursor 1)

To aqueous mixed solutions of cadmium nitrate tetrahydrate (3.085 g, 0.01 mol), and cobalt nitrate hexahydrate (2.910 g,

0.01 mol), an aqueous solution of oxalic acid dehydrate as precipitant (2.52 g, 0.02 mol) was added with continues stirring. A pink precipitate was formed after 1 min from addition. After stirring for 2 h at 80 °C, it was filtered, washed with hot water and acetone several times before being dried using a dry furnace at 110 °C. Precursor 1 was stored in a desiccator until used.

Synthesis of mixed silver oxalate/cadmium oxalate/cobalt oxalate (precursor 2)

To aqueous mixed solutions of silver nitrate (3.4 g, 0.02 mol), (3.085 g, 0.01 mol), and cobalt nitrate (2.910 g, 0.01 mol), an aqueous solution of the oxalic acid as precipitant (3.78 g, 0.03 mol) was added under continues stirring. A pale pink precipitate was formed immediately and then stirred for 2 h at 80 °C. Then, the solid formed was filtered, washed with hot water and acetone several times, and dried using a dry furnace at 110 °C. Precursor 2 was stored in a desiccator until used.

Fabrication of CdO/ Co_3O_4 (3) and Ag-NPs@CdO/ Co_3O_4 (4)

Composites 3 and 4 were fabricated via the thermal degradation of precursors 1 and 2, respectively. 1 g from precursor in porcelain mortar was stated in muffle furnace at 400 °C with heating rate 10 °C/min, then kept 2 h at this degree to remove the crystalline water and oxalate residues. A black powder of composite 3 (0.45 g, 97.83% yield) and deep gray powder of composite 4 (0.50 g, 94.33% yield) were formed and kept in a desiccator until used.

Instrumentation

SEM images were detected by Thermo Scientific Quattro S. FT-IR spectra were measured by IRTracer-100 SHIMADZU spectrophotometer. X-ray patterns were recorded using XRD-7000 SHIMADZU, with a $\text{CuK}\alpha 1$ radiation source. TGA measurements were performed using TGA-51SHIMADZU with a heating rate of 10 °C/min. TEM images were recorded using (FEI, Morgagni, 268 at 80 keV) analysis with dispersion in ethanol and deposited onto the TEM grids having carbon support.

Antibacterial experiments

The antibacterial activity of the materials 1–4 were determined using agar-well diffusion method (Perez 1990). In summary, the bacterial isolates have been cultivated using a nutrient broth medium that contained 5 g/L of yeast extract, 10 g/L bactotryptone and 10 g/L NaCl for 18 h at 37 °C using sterile cotton swabs in a sterile condition; each

bacterial strain was uniformly swabbed onto separate agar plates, and the agar media were punched using 8-mm-diameter wells and filled with 500 μ l of nanoparticles aqueous solutions. The plates were, then, refrigerated for 2 h before being incubated at 37 $^{\circ}$ C for 24 h., where the antibacterial activity of the samples could be accessed by measuring the diameter of growth prevention halo (zone of inhibition). The tested Bacteria were *Bacillus subtilis* (A), *Enterococcus sp.* (B), *E.coli* (C), and *Pseudomonas aerogenosa* (D).

Results and discussion

In this work, a composite of core-shell mixed oxides CdO/Co₃O₄ and its loaded silver nanoparticles new composite are fabricated by thermal degradation of the corresponding mixed oxalate precursors, schemes 1, 2. Several physicochemical techniques have been used to characterize the materials. The antibacterial activity of precursors and composites has been studied.

Materials characterization

TGA analysis

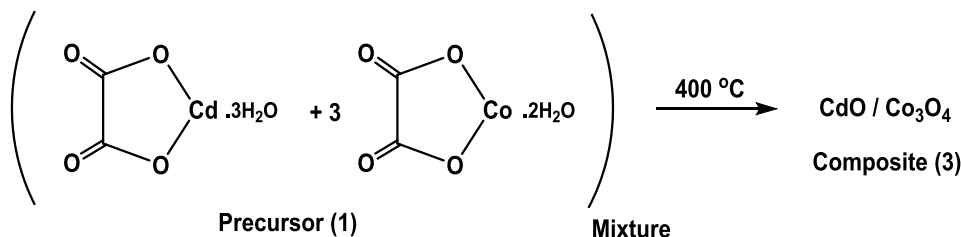
The thermal gravimetric analysis of 1 and 2 were investigated, Fig. 1 to assess the suitable temperature required for conversion of precursors 1 and 2 to composites 3 and 4 and to estimate the composition of the formed residues after thermal degradation. The samples were heated from 25 to 600 $^{\circ}$ C with heating rate 10 $^{\circ}$ C/min. The precursors are stable up to \approx 180 $^{\circ}$ C indicating the absence of adsorbed water molecules. Curve (TG) of precursor 1, (CdC₂O₄.3H₂O + 3CoC₂O₄.2H₂O), Fig. 1a represents two steps of decomposition in the range 180–390 $^{\circ}$ C with mass

loss 54.10% which is close to the theoretical value. The residual part 45.90% corresponds to the successful formation of CdO/Co₃O₄. Figure 1b shows the thermograms of precursor 2, (CdC₂O₄.3H₂O + 3CoC₂O₄.2H₂O + Ag₂C₂O₄). Two degradation steps in the range 180–315 $^{\circ}$ C with mass loss 43.50% leaving residual part 46.50% which agreed well with the theoretical calculations for the formation of 2Ag⁰/CdO/Co₃O₄.

FT-IR spectroscopy

Figure 2a–d shows the FT-IR spectra of materials 1–4, respectively. The precursors FT-IR spectra match the findings of previous studies on the oxalate salts (Nayaka et al. 2016). Figure 2a shows band \sim 3350 cm^{-1} which was assigned to ν O–H of crystalline water molecules (Elseman et al. 2017). The oxalate molecules exhibit transmittance at 1625, 1310, 830 cm^{-1} which are characteristics of bending and stretching vibrations of O–C–O oxalate moiety (Sun and Qiu 2012). The two bands in the far infrared region at 492 and 435 cm^{-1} may be assignable to ν Co–O and ν Cd–O, respectively. Figure 2b exhibits the same bands observed in Fig. 2a as well as the new one at 520 cm^{-1} which could be a result of the ν O–Ag (Nassar et al. 2018). Figure 2c, d shows the FT-IR spectra of composites 3 and 4, respectively. Two apparent and sharp bands at 540–550 cm^{-1} and 640–650 cm^{-1} are characteristic of ν_1 and ν_2 for spinel Co₃O₄, respectively. (Makhlouf et al. 2013). The first band was due to cobaltic oxide vibration in the octahedral hole, and the second band was assigned to cobaltous oxide vibration in the tetrahedral hole in the spinel Co₃O₄ lattice (Fouad et al. 2011). The spectra exhibit a strong band at 750 cm^{-1} . This band is assignable to Cd–O stretching vibration (Małecka and Łącz 2008). No absorptions around 3400 and 1680 cm^{-1} which assignable to H₂O stretching and

Scheme 1 Fabrication of composite 3 from precursor 1



Scheme 2 Fabrication of composite 4 from precursor 2

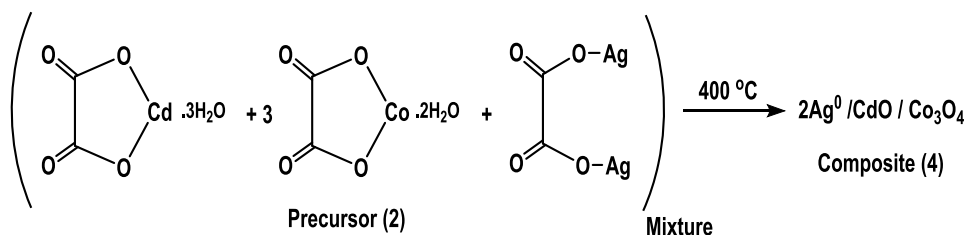


Fig. 1 TGA of precursors 1 (a) and 2 (b)

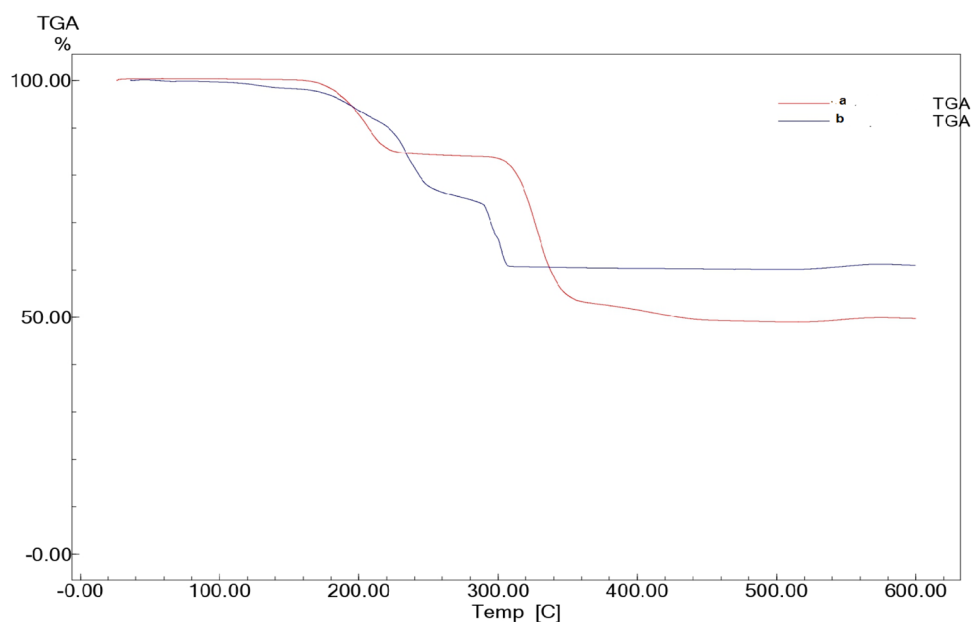
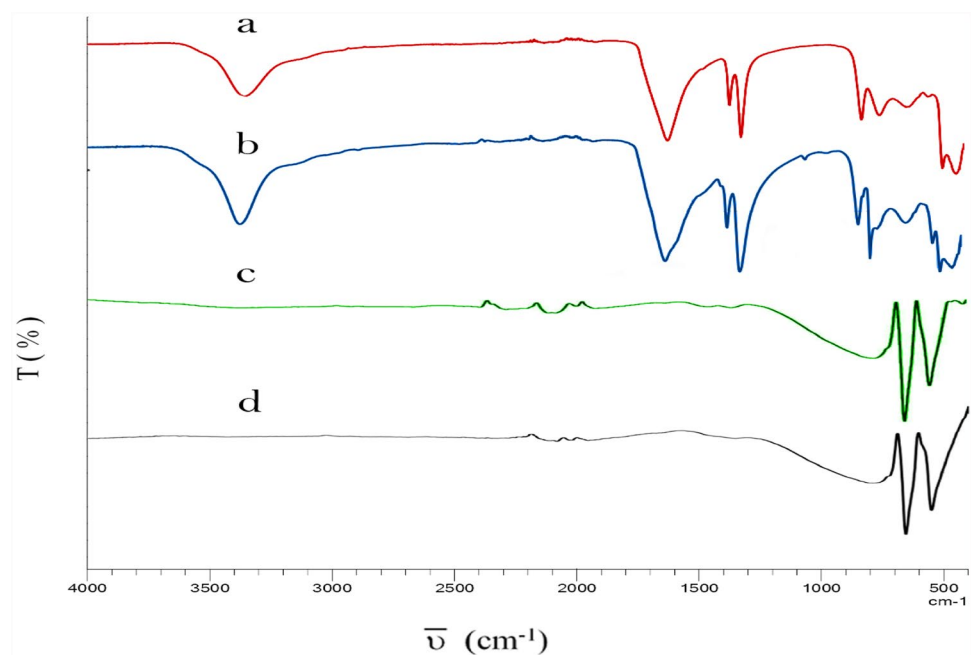


Fig. 2 FT-IR spectra of precursor 1 (a), precursor 2 (b), composite 3 (c) and composite 4 (d)



bending vibrations, respectively, indicating the absence of chemically or physically bonded water molecules in composites structures (Zou et al. 2008).

XRD analysis

Figure 3a–d shows the XRD diffractogram of precursors and their thermal decomposition products. Figure 3a exhibits the characteristic peaks of cobalt oxalate (Du

et al. 2010) and cadmium oxalate (Raj et al. 2008). The peaks at $2\theta = 18.75^\circ$, 22.65° , 30.15° and 44.50° are due to (202), (004), (400) and (224) crystallographic planes which can be indexed as $\beta\text{-CoC}_2\text{O}_4 \cdot 2\text{H}_2\text{O}$. The peaks observed at $2\theta = 18.95^\circ$, 24.48° , 30.20° , 33.60° , 39.70° , 44.56° and 47.75° are referring to planes (111), (012), (200), (210), (213), (030) and (302) indicating the formation of hydrated cadmium oxalate crystals. The main XRD patterns of silver oxalate appeared at $2\theta = 17.10^\circ$,

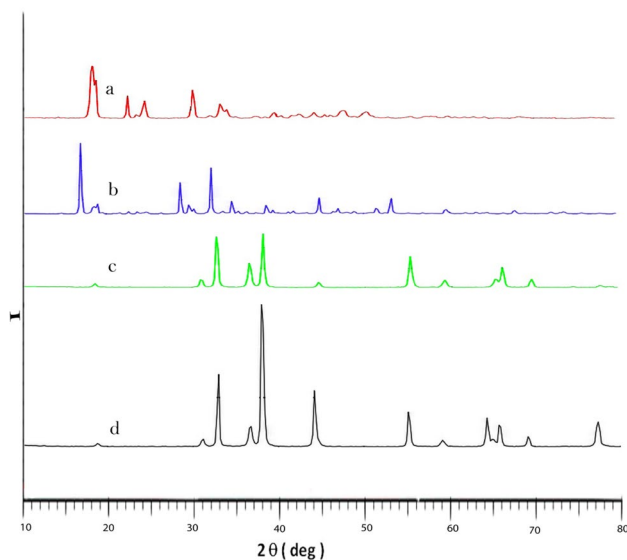


Fig. 3 XRD patterns of precursor 1 (a), precursor 2 (b), composite 3 (c) and composite 4 (d)

19.05°, 28.78°, 29.91°, 32.18°, 36.45°, 38.79°, 39.45°, 47.10°, 51.59° and 67.55°. The X-ray data are in good agreement with the JCPDS No.022-1335 for silver oxalate (Yang et al. 2018). In Fig. 3c, the detected peaks are related to CdO and spinel Co₃O₄ phases. The peaks located at $2\theta = 18.90^\circ, 31.15^\circ, 36.50^\circ, 44.90^\circ, 55.55^\circ, 59.30^\circ$ and 65.33° are assigned to diffraction patterns of crystallographic planes (111), (220), (311), (400), (422), (511), and (440), respectively. All peaks are well matched for Co₃O₄ phase (JCPDS No:76-1802) (Li et al. 2008). The strong diffraction peaks at 2θ values of $33.05^\circ, 38.45^\circ, 65.90^\circ$ and 69.24° correspond to the (111), (200), (311) and (222) planes, respectively. These peaks were indexed to a cubic CdO pattern (JCPDS card no.65-2908) (Usharani et al. 2015). Same peaks of CdO and Co₃O₄ are observed in Fig. 3d as well as the characteristic peaks of silver nanoparticles corresponding to (111), (200), (220), and (311) planes were observed at 2θ values of $38.14, 44.35, 65.00,$ and $77.5,$ respectively (Nassar et al. 2019). No diffraction peaks resulted from the CoO or the metallic cobalt were detected confirming the obtained product from thermal decomposition is pure Co₃O₄ phase. The average crystalline size of each material was calculated using the Scherrer Eq. (1) (Abo Zeid et al. 2020,2019):

$$D = \frac{K\lambda}{\beta \cos \theta} \quad (1)$$

where D is the average particle size, λ is the alpha radiation wavelength of Copper K, β is the whole width of the diffraction peak, $k = 0.9$ is the shape-sensitive coefficient and θ is the diffraction angle. The average crystalline size was found to be 14.21, 14.65, 15.06 and 15.79 nm for materials 1, 2, 3 and 4, respectively. According to G. Gomes et al., silver loading results in larger and more particle sizes (Gomes et al. 2018).

Morphological studies

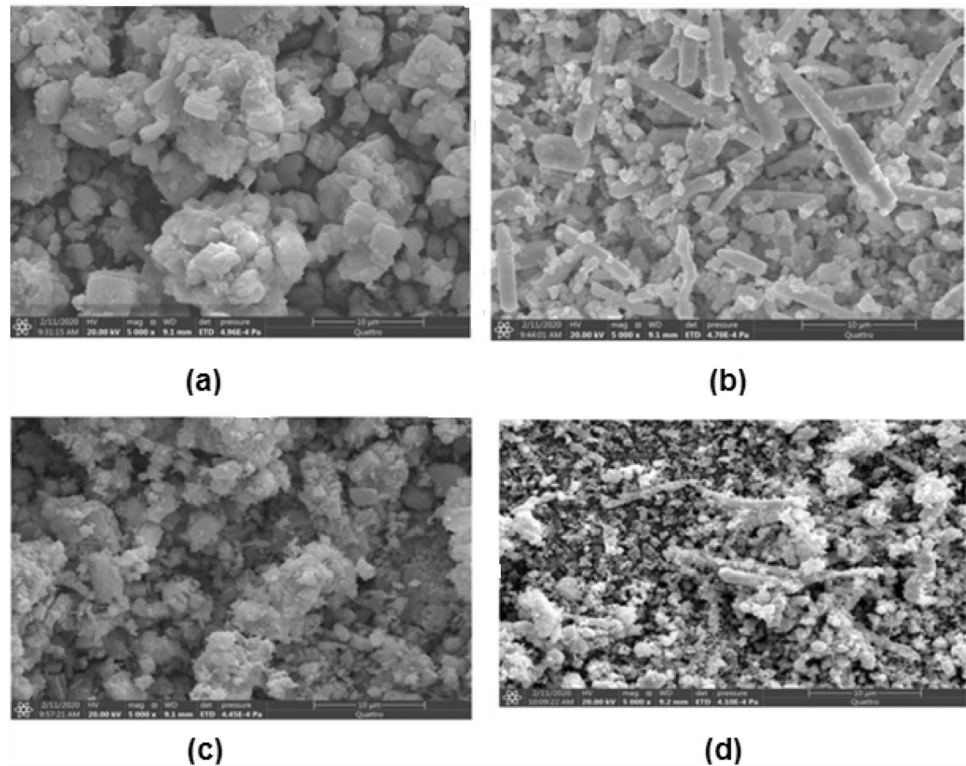
SEM and TEM images of materials were captured and collected in Fig. 4a–d and Fig. 5a–h, respectively. The morphologies of the precursors 1 and 2 show the elongated rod particle shape of cobalt and cadmium oxalates, columnar morphology of silver oxalate and nanosphere of silver ions. The composites 3 and 4 show nanocrystalline structure of CdO/Co₃O₄. In precursor 2 and composite 4 images, the silver nanospheres show physically and uniformly surface dispersion. TEM images of the materials show the average particles size 5–20 nm which was nearly in consonance with the crystals size obtained from XRD. Composite 3 shows core–shell structure as CdO@Co₃O₄. Composite 4 shows the mixed oxides core–shell structure is enveloped with Ag-Nps as Ag@CdO/Co₃O₄. The photos of precursor 2 and composite 4 reveal that the silver particles are predominantly spherical and circular and not physically in contact with each other.

Antibacterial activity

The antibacterial results are provided in Table.1 and illustrated in Figs. 6, 7. The results showed generally that all tested materials exhibited high antibacterial activity following the order $2 > 4 > 1 > 3$ in comparison to both Gram-positive and -negative bacteria. The precursor 2 and composite 4 displayed the higher bacterial growth reduction expressed as larger growth inhibition zone diameter (mm); this stranger antibacterial potential may be due to the presence of silver in their structure.

The most affected bacterial species by the tested nanocomposites were determined to be in the following descending order *Enterococcus sp.* > *E. coli* > *Bacillus subtilis* > and *Pseudomonas aeruginosa*. Similar results obtained showed that CdO NPs and Co₃O₄ NPs exhibited high antibacterial activity (Sadhukhan et al. 2019; Elango et al. 2018). Co₃O₄ and CdO NPs antibacterial activity might be due to generating the reactive oxygen species (ROS) from the surface of NPs that reacted and destroyed the bacterial cell structure. The higher antibacterial potential of silver containing materials might be due to their structural morphology and high

Fig. 4 SEM images of precursor 1 (a), precursor 2 (b), composite 3 (c) and composite 4 (d)



surface area, which apparently well interacted with bacteria and provided a better superficial release of the reactive oxygen species (Morones et al. 2005; Rai et al. 2012).

To sum up, the analysis proved that the precursor and composite supplemented with silver were significantly efficient regarding inactivating some human bacterial pathogens.

The minimum inhibitory concentration (MIC) values of 2 and 4 at different concentration 0.08, 0.16, 0.32 and

0.64 $\mu\text{g}/\text{mL}$, were recorded by standard serial dilution method against *Bacillus subtilis* and *E. coli* using disk diffusion technique (Alsohaimi et al. 2020). The results are summarized in Table 2 and illustrated in Fig. 8. The precursor 2 exhibited the higher antibacterial potential compared to composite 4 expressed as larger inhibition zone diameter (mm), whereas the MIC was found to be 0.08 and 0.32 $\mu\text{g}/\text{mL}$ for 2 and 4, respectively.

Fig. 5 TEM images (a, b) precursor 1, c, d precursor 2, e, f composite 3 and g, h composite 4

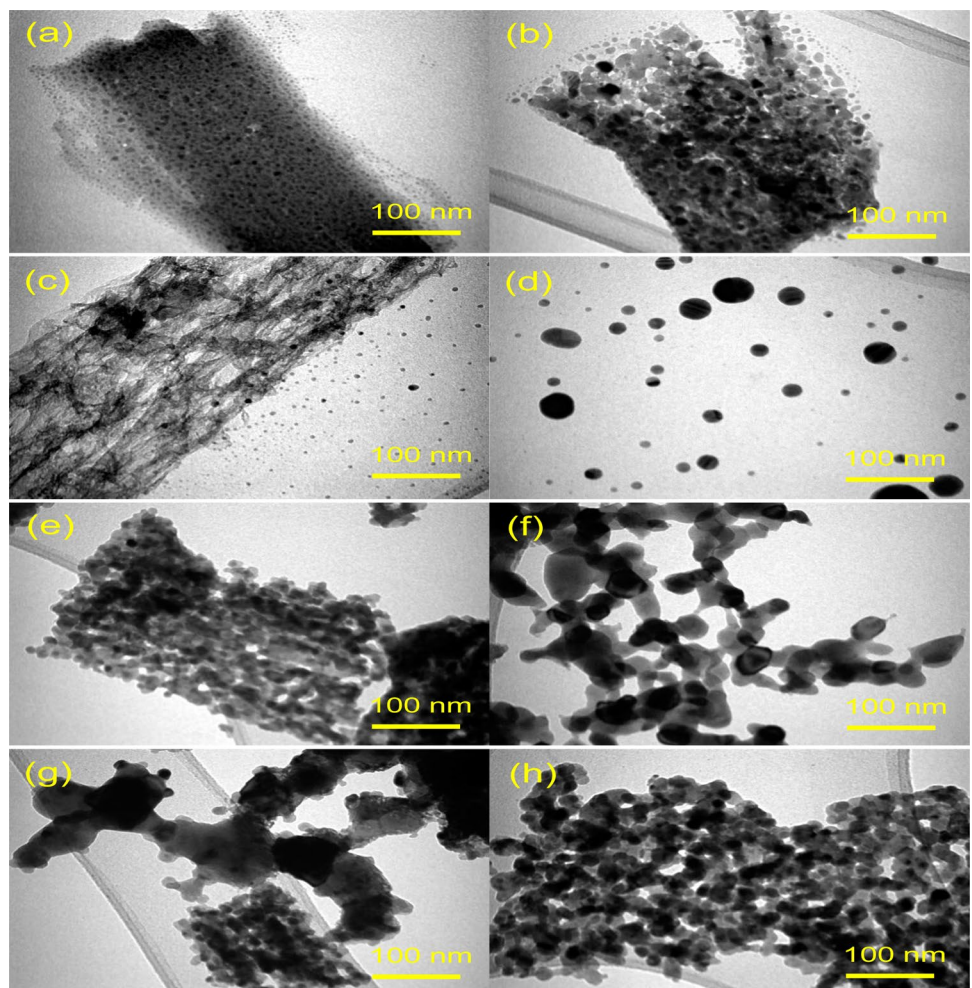


Table 1 Antibacterial activity of materials 1, 2, 3, 4 tested against *Bacillus subtilis* (A), *Enterococcus sp.* (B), *E. coli* (C), and *Pseudomonas aerogenosa* (D)

N	Tested bacteria	Inhibition zone diameter (mm) average ± SD			
		1	2	3	4
A	<i>Bacillus subtilis</i>	31 ± 0.52	32.5 ± 1	25 ± 0.5	28 ± 1.6
B	<i>Enterococcus sp.</i>	27 ± 1.78	40.5 ± 2.3	27 ± 1.75	31.5 ± 1
C	<i>E.coli</i>	26.4 ± 1	32 ± 1	20.5 ± 0.5	42 ± 2.73
D	<i>Pseudomonas aerogenosa</i>	23.45 ± 0.5	33.5 ± 1	20.5 ± 1	29.5 ± 2

Fig. 6 Chart of antibacterial activity of materials 1, 2, 3, 4 tested against *Bacillus subtilis*, *Enterococcus sp.*, *E.coli* and *Pseudomonas aerogenosa*

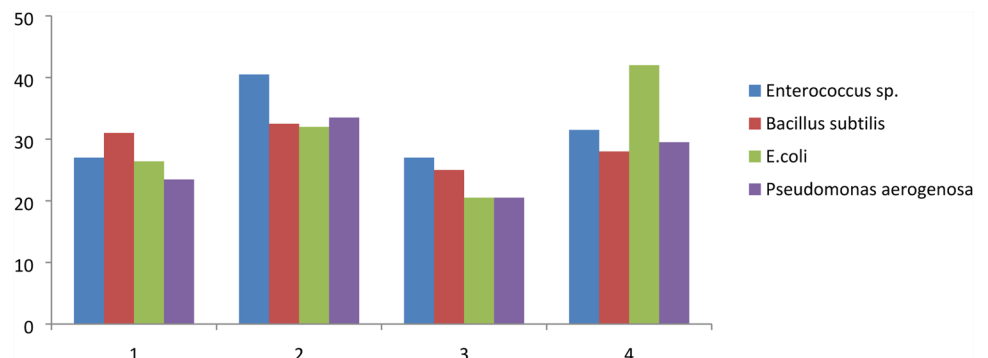


Fig. 7 Images of the results of the antibacterial test for materials 1, 2, 3, 4 against *Bacillus subtilis* (a), *Enterococcus sp.* (b), *E. coli* (c), and *Pseudomonas aerogenosa* (d) after 24 h incubation

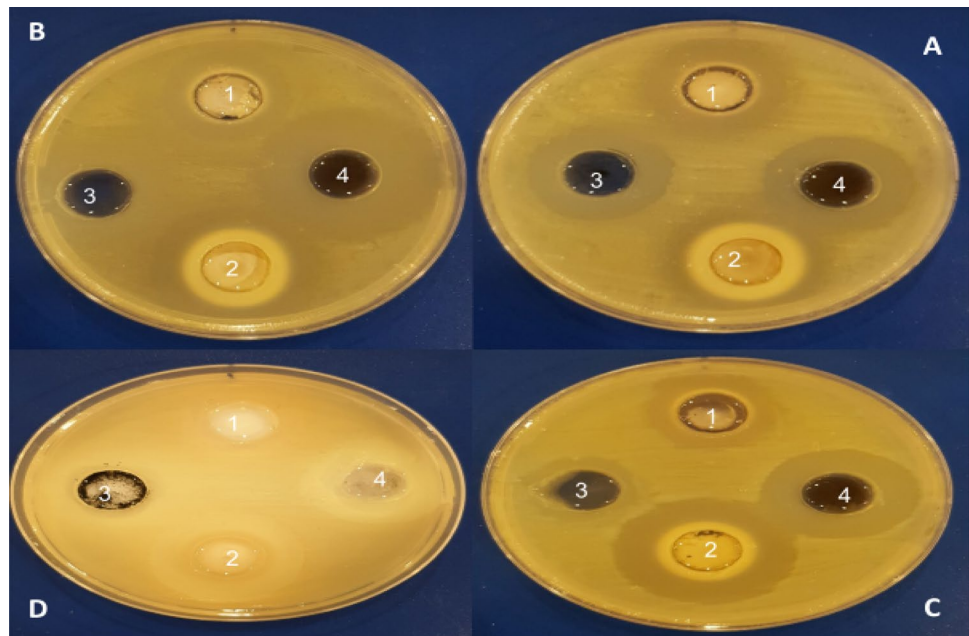
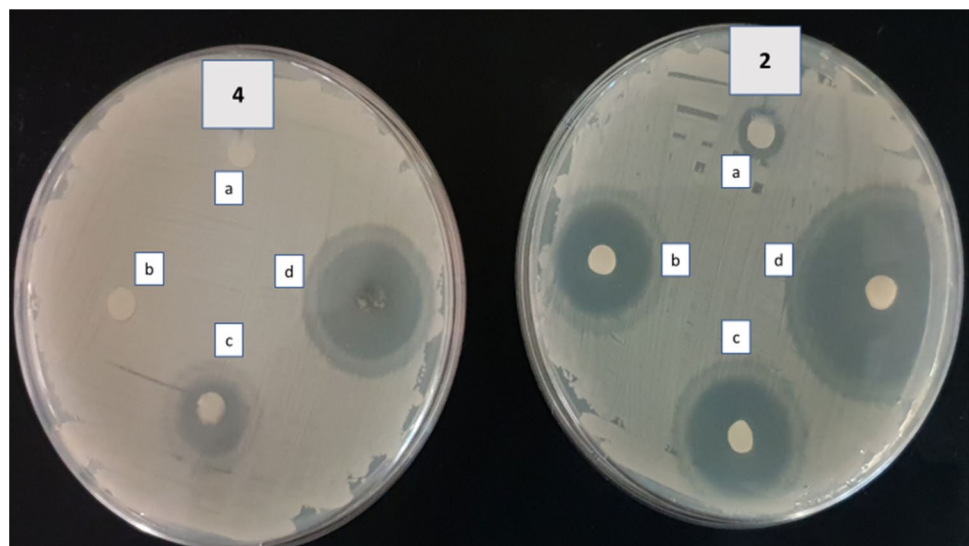


Table 2 MIC values in ($\mu\text{g}/\text{mL}$) of materials 2 and 4 against *Bacillus subtilis* and *E. coli*

Material number	Inhibition zone diameter (mm)							
	2				4			
Material concentration ($\mu\text{g}/\text{mL}$)	0.08	0.16	0.32	0.64	0.08	0.16	0.32	0.64
<i>Bacillus subtilis</i>	10	23	27	41	0	0	19	27
<i>E. coli</i>	10	12	16	40	0	0	10	30

Fig. 8 Results of the antibacterial activity of 2 and 4 as determined by minimum inhibitory concentration (MIC) tested against *Bacillus subtilis* after 24 h incubation. a, b, c, d are abbreviations for concentrations of 0.08, 0.16, 0.32, and 0.64 ($\mu\text{g}/\text{mL}$), respectively



Conclusion

In the present work, a simple route for fabrication of core/shell $\text{CdO}/\text{Co}_3\text{O}_4$ mixed oxides and new composite

$\text{Ag-NPs}@ \text{CdO}/\text{Co}_3\text{O}_4$ by thermal degradation of oxalate precursors was discussed. The nanoparticles of precursors and composites were characterized by physicochemical techniques. The biological activity of the materials was analyzed in terms of the antibacterial activity against two

Gram-positive and two Gram-negative bacterial strains. High inhibition of bacterial growth was resulted from all tested materials particularly from materials contain silver in their structures which are exhibited higher antibacterial activity.

Acknowledgment The authors are thankful for Jouf University for supporting this work by chemicals, laboratories, and analysis.

Compliance with ethical standards

Competing interests No competing interest.

References

- Abo Zeid EF, Ibrahim IA, Ali AM, Mohamed WA (2019) The effect of CdO content on the crystal structure, surface morphology, optical properties and photocatalytic efficiency of p-NiO/n-CdO nanocomposite. *Results in Physics* 12:562–570
- Abo Zeid EF, Nassar AM, Hussein MA, Alam MM, Asiri AM, Hegazy HH, Rahman MM (2020) Mixed oxides CuO-NiO fabricated for selective detection of 2-aminophenol by electrochemical approach. *J Mater Res Technol* 9(2):1457–1467
- Abureesh MA, Oladipo AA, Mizwari ZM, Berksel E (2018) Engineered mixed oxide-based polymeric composites for enhanced antimicrobial activity and sustained release of antiretroviral drug. *Int J Biol Macromol* 116:417–425
- Alsohaimi IH, Nassar AM, Seaf Elnasr TA, Cheba BA (2020) A novel composite silver nanoparticles loaded calcium oxide stemming from egg shell recycling: A potent photocatalytic and antibacterial activities. *J. Clen Prod* 248:119274
- Assim K, Schulze S, Pügner M, Uhlemann M, Gemming T, Giebeler L, Lang H (2017) Co (II) ethylene glycol carboxylates for Co₃O₄ nanoparticle and nanocomposite formation. *J Mater Sci* 52(11):6697–6711
- Balamurugan S, Balu AR, Usharani K, Suganya M, Anitha S, Prabha D, Ilangovan S (2016) Synthesis of CdO nanopowders by a simple soft chemical method and evaluation of their antimicrobial activities. *Pacific Sci Rev A: Nat Sci Eng* 18(3):228–232
- Basu T, Ghosh UC (2013) Nano-structured iron (III)–cerium (IV) mixed oxide: synthesis, characterization and arsenic sorption kinetics in the presence of co-existing ions aiming to apply for high arsenic groundwater treatment. *Appl Surf Sci* 283:471–481
- Bayal N, Jeevanandam P (2012) Synthesis of CuO@ NiO core-shell nanoparticles by homogeneous precipitation method. *J Alloy Compd* 537:232–241
- Du H, Wang J, Wang B, Cang D (2010) Preparation of cobalt oxalate powders with the presence of a pulsed electromagnetic field. *Powder Technol* 199(2):149–153
- Elango M, Deepa M, Subramanian R, Saraswathy G (2018) Synthesis, structural characterization and antimicrobial activities of polyindole stabilized Ag-Co₃O₄ nanocomposite by reflux condensation method. *Mater Chem Phys* 216:305–315
- Elseman AM, Shalan AE, Rashad MM, Hassan AM, Ibrahim NM, Nassar AM (2017) Easily attainable new approach to mass yield ferrocenyl Schiff base and different metal complexes of ferrocenyl Schiff base through convenient ultrasonication-solvothermal method. *J Phys Org Chem* 30(6):e3639
- Fouad OA, Makhlof SA, Ali GA, El-Sayed AY (2011) Cobalt/silica nanocomposite via thermal calcination-reduction of gel precursors. *Mater Chem Phys* 128(1–2):70–76
- Gomes GA, da Costa GL, da Silva Figueiredo ABH (2018) Synthesis of ferrite nanoparticles Cu_{1-x}Ag_xFe₂O₄ and evaluation of potential antibacterial activity. *J Mater Res Technol* 7(3):381–386
- Gong P, Li H, He X, Wang K, Hu J, Tan W, Yang X (2007) Preparation and antibacterial activity of Fe₃O₄@ Ag nanoparticles. *Nanotechnology* 18(28):285604
- Koohestani H (2019) Characterization of TiO₂/WO₃ composite produced with recycled WO₃ nanoparticles from WNiFe alloy. *Mater Chem Phys* 229:251–256
- Kooti M, Saiahi S, Motamedi H (2013) Fabrication of silver-coated cobalt ferrite nanocomposite and the study of its antibacterial activity. *J Magn Magn Mater* 333:138–143
- Li Y, Tan B, Wu Y (2008) Mesoporous Co₃O₄ nanowire arrays for lithium ion batteries with high capacity and rate capability. *Nano Lett* 8(1):265–270
- Li X, Zhu Z, Zhao Q, Wang L (2011) Photocatalytic degradation of gaseous toluene over ZnAl₂O₄ prepared by different methods: a comparative study. *J Hazard Mater* 186(2–3):2089–2096
- Liu J, Wang Z, Luo Z, Bashir S (2013) Effective bactericidal performance of silver-decorated titania nano-composites. *Dalton Trans* 42(6):2158–2166
- Makhlof SA, Bakr ZH, Aly KI, Moustafa MS (2013) Structural, electrical and optical properties of Co₃O₄ nanoparticles. *Superlattices Microstruct* 64:107–117
- Małecka B, Łącz A (2008) Thermal decomposition of cadmium formate in inert and oxidative atmosphere. *Thermochim Acta* 479(1–2):12–16
- Manjula N, Pugalenthil M, Nagarethinam VS, Usharani K, Balu AR (2015) Effect of doping concentration on the structural, morphological, optical and electrical properties of Mn-doped CdO thin films. *Mater Sci-Poland* 33(4):774–781
- Morones JR, Elechiguerra JL, Camacho A, Holt K, Kouri JB, Ramírez JT, Yacaman MJ (2005) The bactericidal effect of silver nanoparticles. *Nanotechnology* 16(10):2346
- Nassar AM, Zeid EA, Elseman AM, Alotaibi NF (2018) A novel heterometallic compound for design and study of electrical properties of silver nanoparticles-decorated lead compounds. *New J Chem* 42(2):1387–1395
- Nassar AM, Elseman AM, Alsohaimi IH, Alotaibi NF, Khan A (2019) Diaqua oxalato strontium (II) complex as a precursor for facile fabrication of Ag-NPs@ SrCO₃, characterization, optical properties, morphological studies and adsorption efficiency. *J Coord Chem* 72(5–7):771–785
- Nayaka GP, Pai KV, Santhosh G, Manjanna J (2016) Recovery of cobalt as cobalt oxalate from spent lithium ion batteries by using glycine as leaching agent. *J Environ Chem Eng* 4(2):2378–2383
- Perez C (1990) Antibiotic assay by agar-well diffusion method. *Acta Biol Med Exp* 15:113–115
- Rai MK, Deshmukh SD, Ingle AP, Gade AK (2012) Silver nanoparticles: the powerful nanoweapon against multidrug-resistant bacteria. *J Appl Microbiol* 112(5):841–852
- Raj AME, Jayanthi DD, Jothy VB (2008) Optimized growth and characterization of cadmium oxalate single crystals in silica gel. *Solid State Sci* 10(5):557–562
- Ren YY, Yang H, Wang T, Wang C (2019) Bio-synthesis of silver nanoparticles with antibacterial activity. *Mater Chem Phys* 235:121746
- Sadhukhan S, Ghosh TK, Roy I, Rana D, Bhattacharyya A, Saha R, Chattopadhyay D (2019) Green synthesis of cadmium oxide decorated reduced graphene oxide nanocomposites and its electrical and antibacterial properties. *Mater Sci Eng, C* 99:696–709
- Sakthivel P, Asaithambi S, Karuppaiah M, Yuvakkumar R, Hayakawa Y, Ravi G (2020) Improved optoelectronic properties of Gd doped

- cadmium oxide thin films through optimized film thickness for alternative TCO applications. *J Alloy Compd* 820:153188
- Salam MA, Abu Khadra MR, Mohamed AS (2020) Effective oxidation of methyl parathion pesticide in water over recycled glass based-MCM-41 decorated by green Co₃O₄ nanoparticles. *Environ Pollut* 259:113874
- Salem A (2014) Silver-doped cadmium oxide nanoparticles: synthesis, structural and optical properties. *The Euro Physical J Plus* 129(12):1–12
- Sun L, Qiu K (2012) Organic oxalate as leachant and precipitant for the recovery of valuable metals from spent lithium-ion batteries. *Waste Manage* 32(8):1575–1582
- Usharani K, Balu AR, Nagarethinam VS, Suganya M (2015) Characteristic analysis on the physical properties of nanostructured Mg-doped CdO thin films—doping concentration effect. *Progress Nat Sci: Mater Int* 25(3):251–257
- Vennela AB, Mangalaraj D, Muthukumarasamy N, Agilan S, Hemalatha KV (2019) Structural and optical properties of Co₃O₄ nanoparticles prepared by sol-gel technique for photocatalytic application. *Int J Electrochem Sci* 14:3535–3552
- Wang X, Yang Y, Zhang F, Tang J, Guo Z (2020) Facile synthesis of Co₃O₄/CdO nanospheres as high rate performance supercapacitors. *Mater Lett* 261:127141
- Williams JM, Beno MA, Carlson KD, Geiser U, Kao HI, Kini AM, Thorn RJ (1988) High transition temperature inorganic oxide superconductors: synthesis, crystal structure, electrical properties, and electronic structure. *Acc Chem Res* 21(1):1–7
- Yang W, Wang C, Arrighi V (2018) Silver oxalate ink with low sintering temperature and good electrical property. *J Electron Mater* 47(5):2824–2835
- Yuan C, Wu HB, Xie Y, Lou XW (2014) Mixed transition-metal oxides: design, synthesis, and energy-related applications. *Angew Chem Int Ed* 53(6):1488–1504
- Zahera M, Khan SA, Khan IA, Sharma RK, Sinha N, Al-Shwaiman HA, Khan MS (2020) Cadmium oxide nanoparticles: an attractive candidate for novel therapeutic approaches. *Colloids Surf, A* 585:124017
- Zakrzewska K (2001) Mixed oxides as gas sensors. *Thin Solid Films* 391(2):229–238
- Zou D, Xu C, Luo H, Wang L, Ying T (2008) Synthesis of Co₃O₄ nanoparticles via an ionic liquid-assisted methodology at room temperature. *Mater Lett* 62(12–13):1976–1978

Publisher's Note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.