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Synthesis of homo- and hetero-metallic cobalt and zinc nano oxide particles by a calcination process using coordination compounds: their characterization, DFT calculations and capacitance behavioural study†

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Nano cobalt and porous zinc–cobalt oxide particles were synthesized using the concept of coordination compounds of the type $[M(II)L'L']$ (where $M(II) = Co(II) \& Zn(II)$, $L = 4$ -hydroxy benzaldehyde and $L' =$ piperazine) and were thoroughly characterized. Because the precursors are coordination compounds possessing specific geometry in the crystal lattice, uniform and appropriately sized homo- and heterometallic nanocrystals of Co_3O_4 and $ZnO \cdot Co_3O_4$ were obtained after a thermal process. The homo and hetero composite particles were characterized by transmission electron microscopy (TEM), scanning electron microscopy (SEM), energy dispersive X-ray analysis (EDX), X-ray diffraction (XRD), FT IR spectroscopy and electrochemistry. The paramagnetic chemical shift of the methyl protons in DMSO due to the nanoparticles was studied by NMR spectroscopy, which indicated that the cobalt particles were ferromagnetic. The structural design modification and surface area of Co_3O_4 was improved by adding the ZnO component. DFT calculations were done to validate the nano structure. Supercapacitance ability of the nanoparticles was studied by cyclic voltammetry, and electrochemical calculations were performed to determine the microelectronic characteristics of the material. The specific capacitance was estimated at 207.3 and 51.1 $F\ g^{-1}$ for the $ZnO \cdot Co_3O_4$ and Co_3O_4 electrodes, respectively. Clearly, $ZnO \cdot Co_3O_4$ exhibited a much higher specific capacitance than the Co_3O_4 nanocrystal, which was attributed to better conductivity and higher surface area. The capacitance activity showed multifold enhancement due to the porous nature of Zn oxide in the heterometallic nano $ZnO \cdot Co_3O_4$ composite.

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1. Introduction

Transition metal oxides have gained considerable interest in recent years owing to their interesting magnetic, optical field emission and biomedical applications. Transition metal ions exist in variable oxidation states, which make them versatile precursor materials for use in nanoscale chemistry. Among the prominent mixed valence oxides, spinel type cobalt oxides (Co_3O_4) have attracted considerable attention in the areas of environmental science, catalysis and medicine. For example, many Co_3O_4 NPs have been utilized for the degradation of

pollutants and in electro catalysis for oxygen hydrogen generation, and very recently, cobalt nano oxides were used as probes in medical diagnostic devices.^{1–4} A glucose sensor of cobalt oxide nano rods was prepared by Kuo-Chuan Ho *et al.* for the non-enzymatic detection of glucose.⁵ The interesting applications of nano transition oxides depend on their size and different structural morphologies including nanotubes, nanorods, nano cubes and meso porous structures.

Many synthetic techniques and routes have been utilized to prepare these nanomaterials, including sol–gel methods,⁶ solvothermal synthesis,⁷ thermal decomposition of cobalt precursors,⁸ sonochemical methods,⁹ co-precipitation¹⁰ and microwave-assisted methods.¹¹ Most of these methods are not feasible for large-scale production owing to the expensive and toxic chemicals required and the use of complex instruments. Researchers are looking for more facile synthetic routes to obtain new nanomaterials of mixed valence oxides by choosing appropriate precursors, which have potential advantages including high yield

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of pure products, the absence of solvents, low energy consumption and functional efficiency. Herein, we have undertaken the task of preparing homo and hetero-metallic Co_3O_4 , $\text{ZnO} \cdot \text{Co}_3\text{O}_4$ mixed-valence oxides possessing different structural morphologies and electrochemical behaviour.^{12,13} In this work, we report a new modified calcination process that uses the coordination chemistry concept of employing piperazine and aldehyde with metal salts to obtain a uniform single crop of nano Co_3O_4 and porous $\text{ZnO} \cdot \text{Co}_3\text{O}_4$ crystals. The heterobimetallic oxides show multifold enhanced activities (catalytic and capacitance properties) compared to monometallic nano oxide.¹⁴ The obtained heterobimetallic nanomaterials have mixed oxidation states, which helps build up the inner electric field at the junction interface and create more pores in the porous material.^{15,16}

2. Experimental

2.1. Materials

Chemicals, including CoCl_2 , $\text{Zn}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ (4-hydroxy benzaldehyde), and piperazine were purchased from Sigma Aldrich, USA. Power X-ray diffraction (XRD) of the products was measured using a Philips X'Pert PRO MPD diffractometer at a scanning rate of 4° min^{-1} , with 2θ ranging from 10° to 70° , using $\text{Cu K}\alpha$ radiation ($\lambda = 1.5406 \text{ \AA}$). The morphologies of the samples were studied by scanning electron microscopy (SEM) (JEOL SM5600LV) at 20 kV. The powders were ultrasonicated in ethanol, and a drop of the suspension was dried on a carbon-coated microgrid. Transmission electron microscopy (TEM) observations were performed with a JEM 100CX-II microscope operated at 100 kV. NMR spectra were recorded in $\text{DMSO-}d_6$ on a Jeol 400 MHz NMR spectrometer. Thermal studies were performed using a TGA/SDTA 851e (Mettler Toledo) thermogravimetric analyser in ambient atmosphere from 20°C to 700°C at a heating rate of $10^\circ \text{C min}^{-1}$. Electrochemical measurements were performed using a potentiostat (AutolabPGSTAT101) in a standard three-electrode setup, with a working electrode of CoZnO_2 and Co_3O_4 nanocrystals ($50 \mu\text{g}$ dispersed in water and isopropanol solution) loaded on a carbon paper substrate (SIGRACET®, grade GDL-24BC, geometric area $1 \times 1 \text{ cm}^2$), as well as a Pt mesh and a saturated calomel electrode (SCE) as the counter and reference electrodes, respectively.

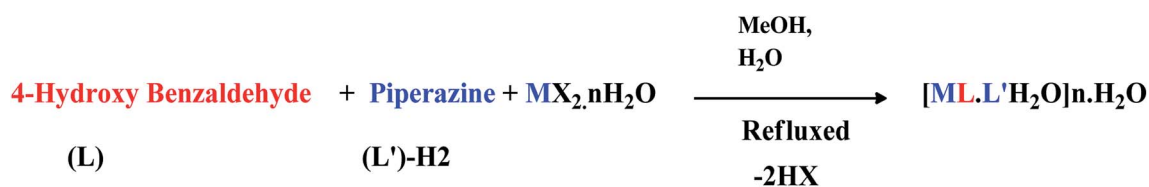
2.2. Synthesis of precursors and nanoparticles

The mono and heterobimetallic nanoparticles were prepared by a modified coordination chemistry procedure. A methanolic solution of the metal salt $\text{Co}(\text{II})/\text{Zn}(\text{II})$, aldehyde and piperazine

at a 1 : 1 : 1 molar ratio was refluxed for 2 h in a 100 mL round bottom flask (Scheme 1). The blue cobalt complex and white powder of the zinc compound were obtained, and adducts were washed with methanol and hexane and were dried under vacuum. The prepared coordination compounds were characterized by FTIR and mass spectrometry on the basis of the preliminary characterization (Scheme 1). $[\text{Co}(\text{II})\text{benzaldehyde} \cdot \text{piperazine} \cdot \text{H}_2\text{O}]0.5\text{H}_2\text{O}$; mp 300°C dm/z 292.15(293.20) $[\text{M}-\text{L}-\text{L}'/1.5\text{H}_2\text{O} + \text{H}]$. $[\text{Zn}(\text{II})\text{benzaldehyde} \cdot \text{piperazine} \cdot \text{H}_2\text{O}]$; mp 235°C dm/z 289.65(293.19) $[\text{Zn}-\text{L}-\text{L}' \cdot \text{H}_2\text{O}-3\text{H}^+]$. The FTIR bands at 498, 877, 998, 1224, 1342, 1413, 1450, 1584, 2825, 3196 and 3556 cm^{-1} are due to the $\text{C}=\text{O}$, $\text{C}-\text{C}$, $\text{C}-\text{H}$, $\text{C}-\text{N}$ and H_2O vibration and bending modes. The broad bands in the $3556-3196 \text{ cm}^{-1}$ range in the spectra of the precursors have been attributed to the stretching vibrations of H_2O , OH and NH. The band at 2825 cm^{-1} was due to the $\text{C}-\text{H}$ stretching mode, and the bands at 1342 and 1584 cm^{-1} were assigned to the bonding of $\text{M}(\text{II})$ with $\text{N}-\text{H}$ and $\text{C}-\text{O}$. Another band due to $\text{C}-\text{O}$ stretching was observed at 1224 cm^{-1} in the spectra of the precursors. The low-frequency absorptions at 498 and 877 cm^{-1} were attributed to the $\text{M}-\text{O}$ stretching and $\text{M}-\text{O}$ bending vibrations in the spectra of the complexes.¹⁷ The $[\text{Co}(\text{II})\text{benzaldehyde} \cdot \text{piperazine} \cdot \text{H}_2\text{O}]0.5\text{H}_2\text{O}$ was homogenized by ultrasonication. The powder was thoroughly washed with anhydrous ethanol to remove impurities. The compound was dried in an air oven at 60°C for 12 h and then calcinated at $500-600^\circ \text{C}$ for 6 h in an electric furnace. Co_3O_4 nanoparticles were prepared with 0.75 g of the complex $[\text{Co}(\text{II})\text{benzaldehyde} \cdot \text{piperazine} \cdot \text{H}_2\text{O}]0.5\text{H}_2\text{O}$ in a porcelain crucible and placed in the furnace. The compound was heated to 500°C at a rate of $10^\circ \text{C min}^{-1}$. The cobalt nanoparticles were characterized by various spectroscopic methods. The heterometallic nanocomposite was formed by mixing of $[\text{Co}(\text{II})\text{benzaldehyde} \cdot \text{piperazine} \cdot \text{H}_2\text{O}]0.5\text{H}_2\text{O}$ and $[\text{Zn}(\text{II})\text{benzaldehyde} \cdot \text{piperazine} \cdot \text{H}_2\text{O}]$, coordination compounds at a 1 : 5 ratio. The high concentration of the zinc coordination compound in methanol was used because Zn forms a porous material; hence, pockets of Co_3O_4 will be formed, which will increase the surface area in the $\text{ZnO} \cdot \text{Co}_3\text{O}_4$ composite material. The mixture was sonicated for approximately 25 minutes in methanol, filtered, washed with hexane and dried in an oven at 60°C . The nanoconjugate was prepared by calcination, as reported for homometallic Co_3O_4 .

2.3. Computational method

We carried out a series of theoretical calculations of $\text{ZnO} \cdot \text{Co}_3\text{O}_4$ molecular aggregates in order to find out how closely they could



Where $\text{M}=\text{Co}(\text{II})/\text{Zn}(\text{II})$, $\text{X}=\text{Cl}/\text{NO}_3$, and $n=0/0.5$

Scheme 1 Synthetic route of homo and heterometallic nano particles.



approach each other and which atoms and their interactions would be involved in the adduct formation. Herein we adopt a three step molecular modeling protocol, (1) generation of Co_3O_4 and ZnO nano particles molecular coordinates from the X-ray crystal structures in two different sizes *e.g.* $[\text{Co}_3\text{O}_4]_4$ (9.00 nm), $[\text{Co}_3\text{O}_4]_{10}$ (12 nm), $[\text{ZnO}]_4$ (5 nm), and $[\text{ZnO}]_{20}$ (10 nm). (2) Geometry optimization of $[\text{Co}_3\text{O}_4]_n$, $n = 4, 10$ with $[\text{ZnO}]_m$, $m = 4, 20$, by employing molecular mechanics force field which includes van der Waals (Lennard-Jones potential), hydrogen bonding, desolvation and electrostatic terms and treats the intramolecular bonds and bond angles of both the molecules as rigid. (3) DFT calculation of three geometrically optimized $[\text{Co}_3\text{O}_4]_n \cdot [\text{ZnO}]_m$ adducts (where $n = 4, 10$ and $m = 4, 20$), (a) $[\text{Co}_3\text{O}_4]_4 [\text{ZnO}]_4$, (b) $[\text{Co}_3\text{O}_4]_{10} [\text{ZnO}]_4$, and (c) $[\text{Co}_3\text{O}_4]_{10} [\text{ZnO}]_{20}$. All the molecular mechanic energy-minimization of corresponding molecular adducts were done using Autodock 4.2 software.¹⁸ All reported DFT computations were performed using ORCA computational package¹⁹ for previously optimized structures. The single point energy calculation carried out by unrestricted B3LYP functional²⁰ using Aldrich's def2-TZVP basis set for all the atoms²¹ to calculate the HOMO and LUMO

energies. To speed up the calculations we have used the resolution of identity (RI) approximation with the decontracted auxiliary def2-TZV/J Coulomb fitting basis sets and the chain-of-spheres (RIJCOSX) approximation to exact exchange as executed in ORCA. DFT calculation utilizes the atom-pairwise dispersion correction with the Becke–Johnson damping scheme (D3BJ).²²

3. Results and discussion

The precursors of homo- and heterometallic nano oxides were prepared by using the coordination chemistry concept with a low-cost starting material. The characterization and morphology of the nano Co_3O_4 and $\text{ZnO} \cdot \text{Co}_3\text{O}_4$ were studied by TEM and SEM images. The TEM images showed that Co_3O_4 and $\text{ZnO} \cdot \text{Co}_3\text{O}_4$ particles possessed a rectangular unit cell with several vertices of Co_3O_4 (size 19.14–56.20 nm) and porous ZnO crystalline (22.79 nm pore size) pockets filled with Co_3O_4 . The particle size of the rectangular pyramidal units was 14.14–16.13 nm. On the basis of the TEM images, it can be concluded that particle overlap occurred and that different long nanoconjugates were obtained by interfacial reactions and agglomeration. The SEM results

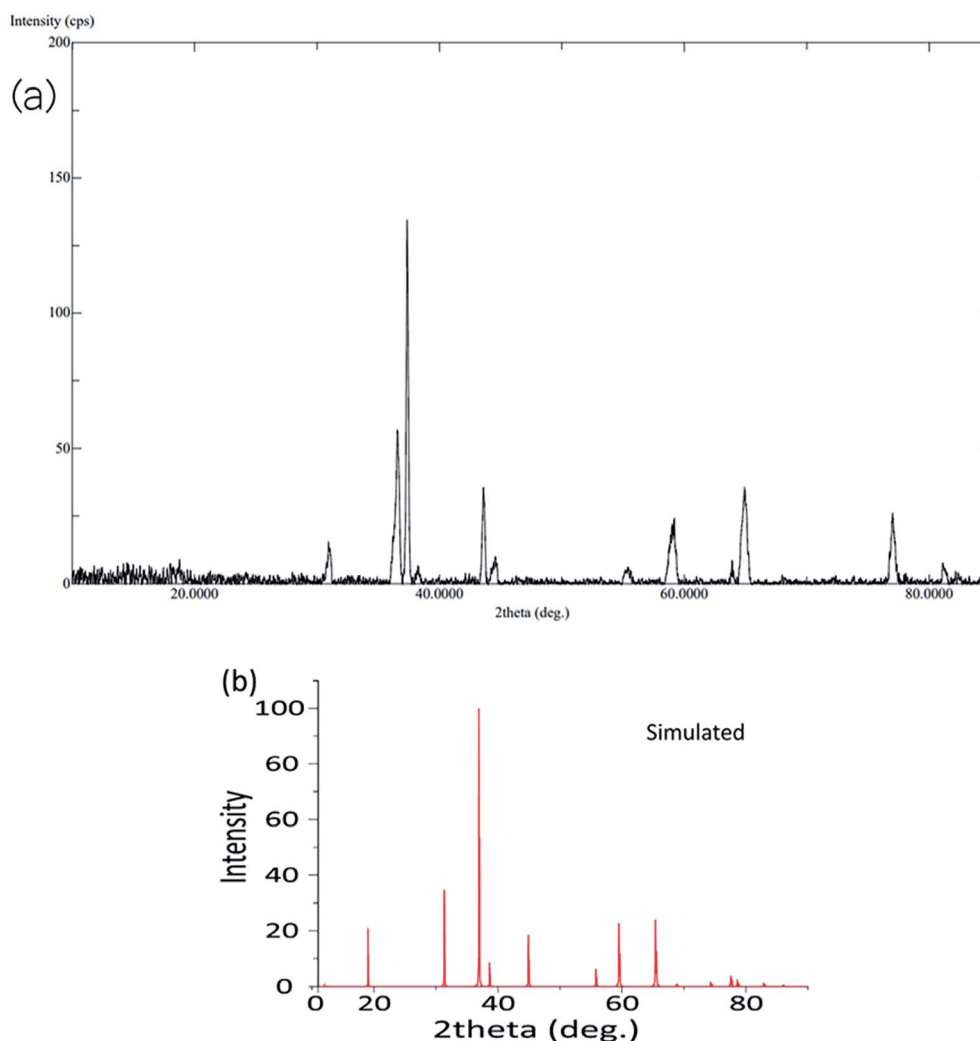


Fig. 1 Experimental and simulated XRD pattern of Co_3O_4 nanoparticles.



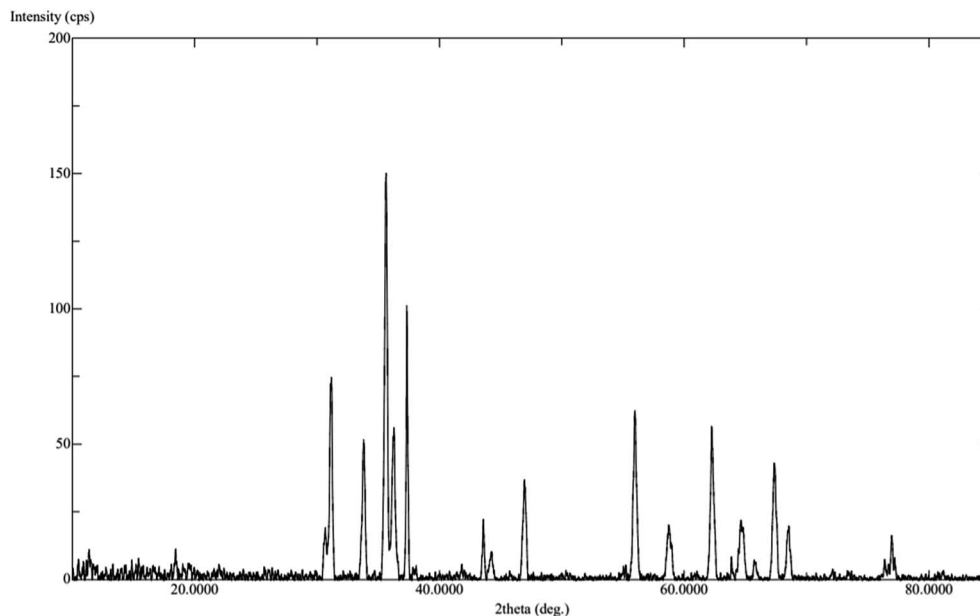


Fig. 2 XRD pattern of ZnO·Co₃O₄ nanoparticles.

corroborated well with the TEM results for the Co₃O₄ and ZnO·Co₃O₄ nanoparticles. This procedure is important because we can obtain identical nano oxide particles with a defined geometry by repeating the procedure.

3.1. Infrared spectroscopy

FTIR spectra were recorded for cobalt and zinc nanoparticles to confirm the bonding of oxides to metal ions in the nanocrystal

materials and the cationic position in the structure. The FTIR spectra (Fig. S1 and S2[†]) showed two characteristic stretching vibrations bands of the M–O bonds in Co₃O₄. A sharp band appeared at 583 cm⁻¹ due to the vibration of Co(III) ions in the octahedral void of oxide ions. The second band appeared at 665 cm⁻¹, confirming the presence of Co²⁺ in the tetrahedral hole, which resulted in the formation of pure nanocrystals.²³ A similar FTIR was obtained for ZnO·Co₃O₄, with additional weak bands at a lower frequency of 580 cm⁻¹, which indicated the

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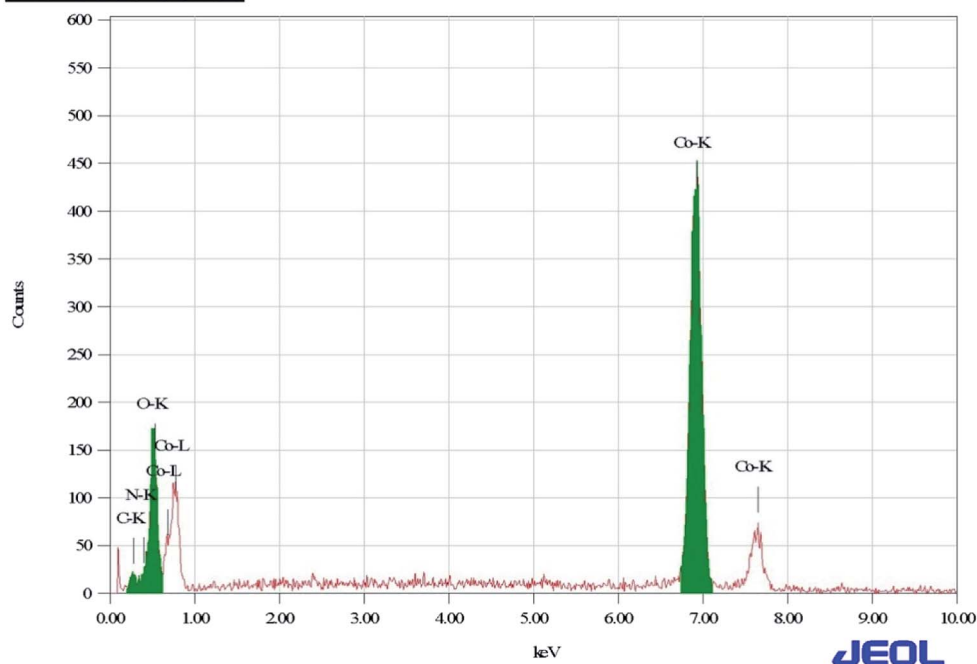
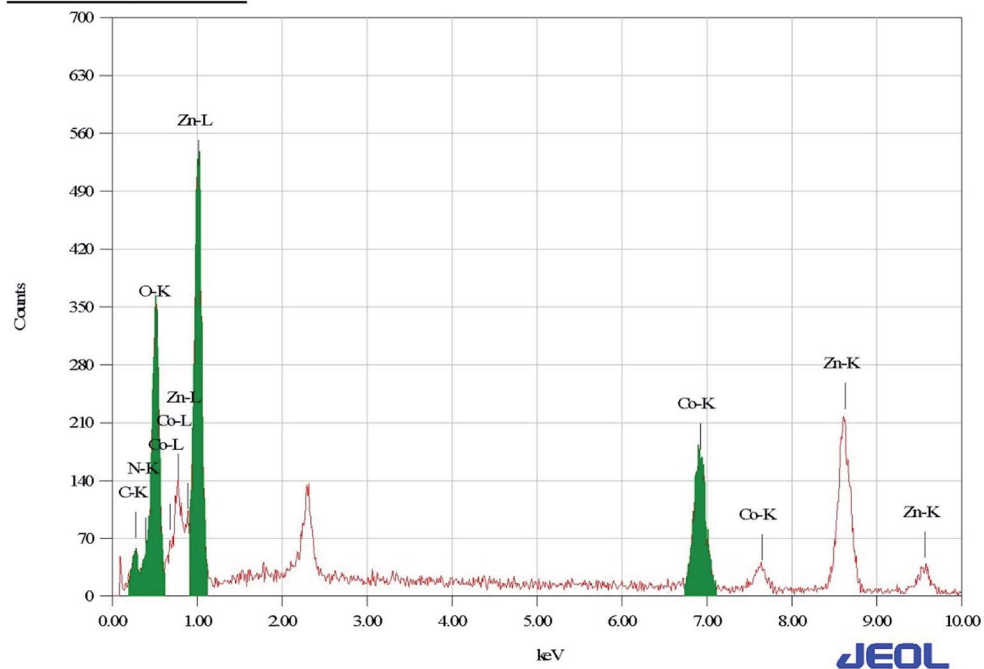
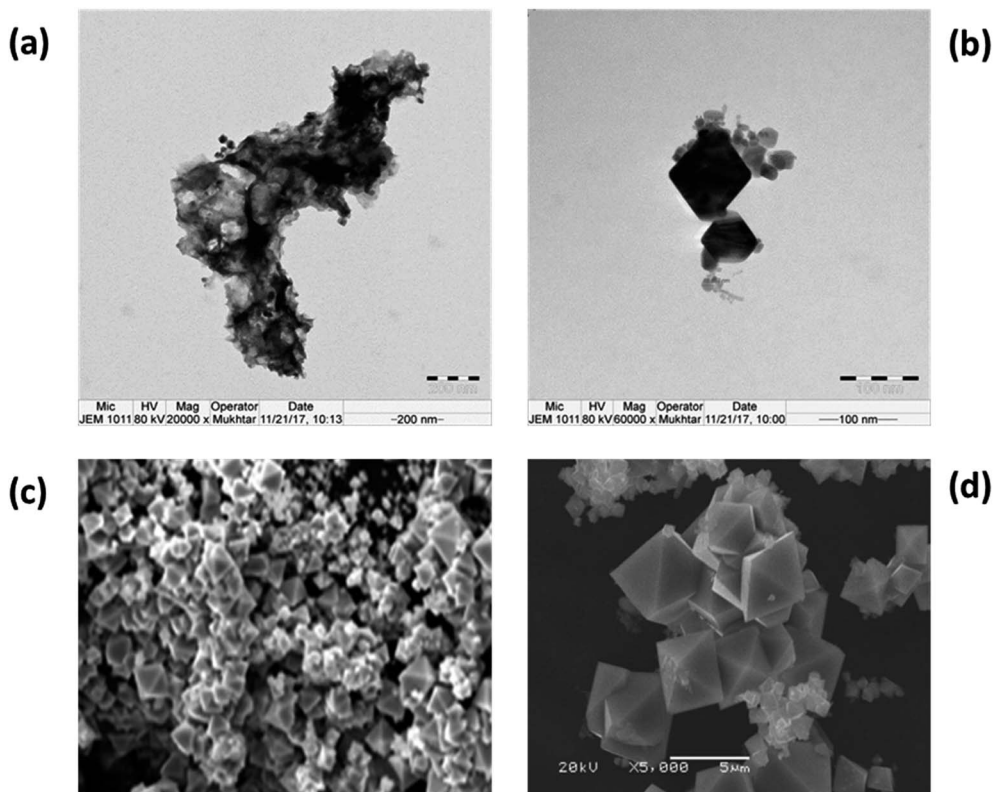


Fig. 3 EDX of Co₃O₄.



JED-2200 SeriesFig. 4 EDX of ZnO·Co₃O₄.Fig. 5 Typical TEM images: (a) Co₃O₄ nanoparticles and (b) the shape and size of the particles. Typical SEM images of a Co₃O₄ (c) bulk particle and (d) the geometry of Co₃O₄ particles.

presence of Zn oxide with Co_3O_4 .²⁴ The cobalt was present as Co^{2+} , and two cobalt ions were in the Co^{3+} oxidation state. Mixed oxidation states (divalent and trivalent ions) provide crystal field stabilization at the octahedral (Co^{3+}) and tetrahedral (Co^{2+}) sites of Co_3O_4 . No other band was observed, thus confirming the purity of the oxide nanoparticles.

3.2. Thermo gravimetric analysis (TGA)

The TGA analysis of the $[\text{M}(\text{II})\text{benzaldehyde}\cdot\text{piperazine}\cdot\text{H}_2\text{O}]n\text{H}_2\text{O}$, with $\text{M} = \text{Co}(\text{II})/\text{Zn}(\text{II})$, indicated a double-step weight loss between 573–763 K. The weight loss indicated the decomposition of $[\text{M}(\text{II})\text{benzaldehyde}\cdot\text{piperazine}\cdot\text{H}_2\text{O}]$ into the metal oxides $\text{Co}_3\text{O}_4/\text{ZnO}$. The weight loss was found to be approximately 68%, which is equivalent to the loss of water, aldehyde and piperazine

compounds to form cobalt oxide/zinc oxide. The observed weight loss of cobalt oxide was close to the theoretical value. Upon the calcination of the obtained $[\text{Co}(\text{II})\text{benzaldehyde}\cdot\text{piperazine}\cdot\text{H}_2\text{O}]n\text{H}_2\text{O}$ and the mixture of $[\text{Co}(\text{II})\text{benzaldehyde}\cdot\text{piperazine}\cdot\text{H}_2\text{O}]n\text{H}_2\text{O}$, $[\text{Zn}(\text{II})\text{benzaldehyde}\cdot\text{piperazine}\cdot\text{H}_2\text{O}]$ at 773 K in air, both metal complexes were converted into Co_3O_4 and $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ nanoparticles. The TGA (Fig. S3†) curves of the Co_3O_4 oxide nanoparticles show no weight loss, indicating the purity and stability of the particles. The XRD pattern of the sample also clearly indicated that nano Co_3O_4 particles are pure oxides.

3.3. XRD measurement

The calcination process was performed directly at 500–600 °C to convert the mono metallic compound and mixture of two

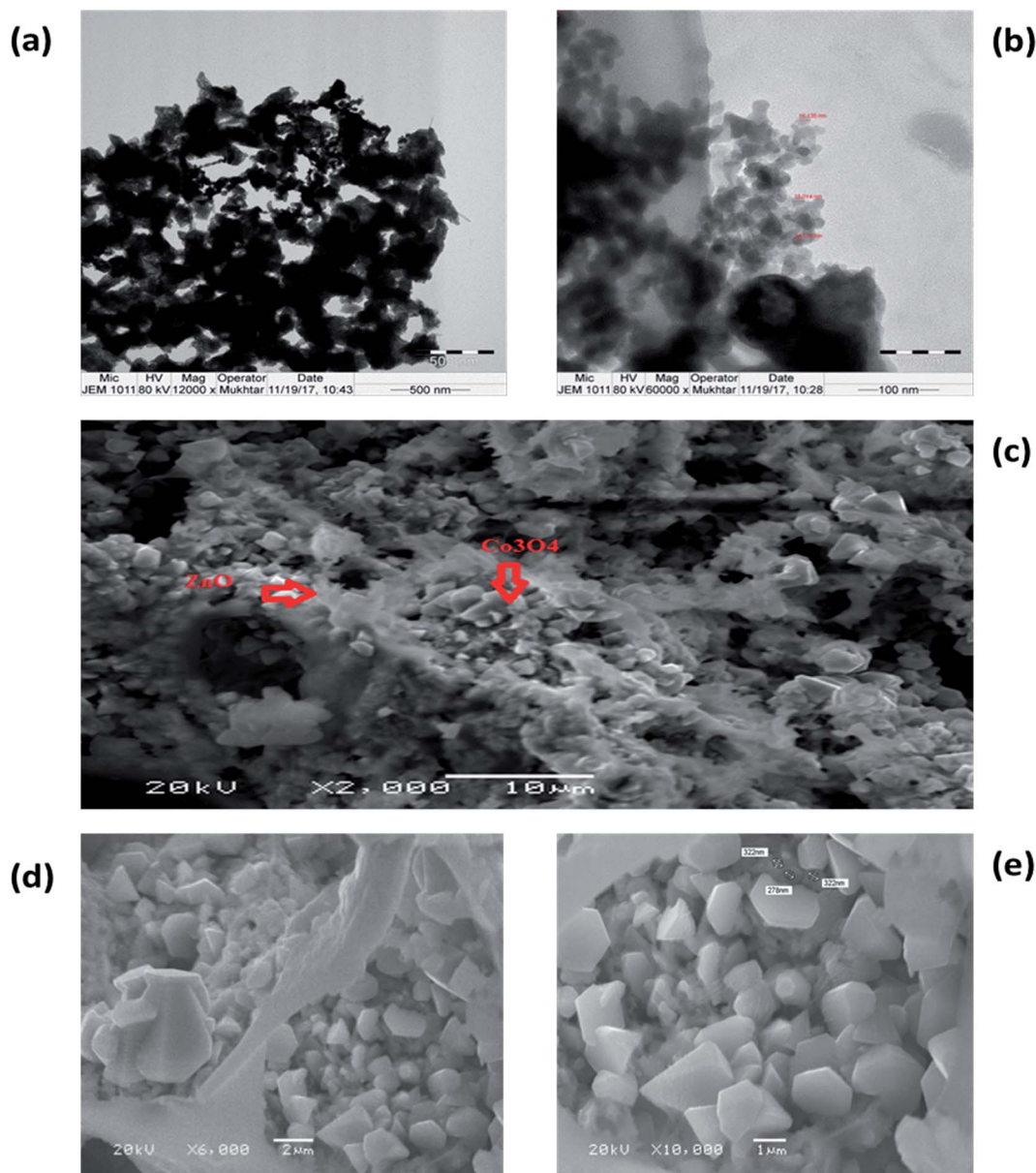


Fig. 6 Typical TEM images, (a) and (b) of the structure and size of $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ particles. SEM images of $\text{ZnO}\cdot\text{Co}_3\text{O}_4$, showing (c) the porous structure of ZnO with, (d) and (e), pockets of Co_3O_4 .



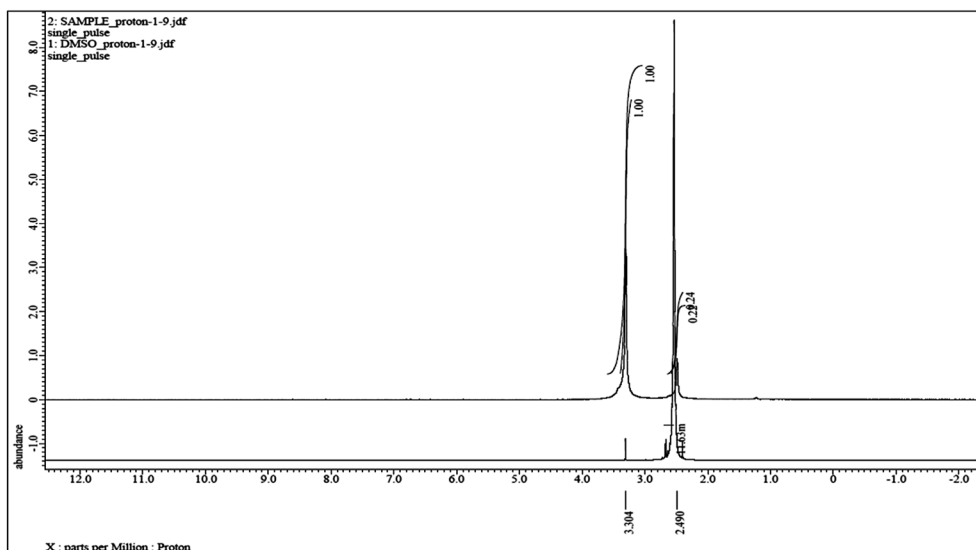


Fig. 7 ^1H NMR chemical shift due to Co_3O_4 .

complexes to get homo and heterometallic oxides. The nano Co_3O_4 and $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ products maintained the crystalline phases of the oxides over time (Fig. S4†). The X-ray diffraction confirmed that both the nano Co_3O_4 and $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ composite were pure. The peaks of the Co_3O_4 and $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ were observed at $2\theta = 36.56^\circ, 37.56^\circ, 38.24^\circ, 43.58^\circ, 64.92^\circ,$ and 65.04° due to the 111, 220, 311, 400, 440 and 511 planes for Co_3O_4 . For $\text{ZnO}\cdot\text{Co}_3\text{O}_4$, peaks were observed at $2\theta = 30.50^\circ, 31.20^\circ, 31.35^\circ, 33.78^\circ, 35.56^\circ, 36.30^\circ, 36.56^\circ, 37.34^\circ, 37.56^\circ, 43.58^\circ, 46.96^\circ, 55.96^\circ, 62.24^\circ,$ and 64.92° , due to ZnO 100, 002, 101, 102, 110 and 103. Hexagonal ZnO with a lattice along the Co_3O_4 diffraction lines 111, 220, 313, 400, 422, 440, and 511 corresponded to a rectangular pyramidal crystal supported by JCPDS data (Fig. 1). Our results are in good agreement with previously reported results.²⁴ No peaks signifying other combinations of cobalt were obtained, which confirms the purity and crystalline structure of the Co_3O_4 nanoparticle and $\text{ZnO}\cdot\text{Co}_3\text{O}_4$. Additionally, the simulated XRD pattern of Co_3O_4 structure is in good agreement with the experimentally observed XRD pattern also indicated the purity of Co_3O_4 nanoparticles (Fig. 2). The size ($\sim 15\text{--}31$ nm) of the crystalline phase Co_3O_4 nanocrystal and the Zn porous crystal containing Co_3O_4 was calculated from the XRD data using the Scherrer equation (a).²⁵ The similar range of the size of the particle was measured by TEM, which supports the stable phase of the crystalline material at room temperature after calcination.

3.4. EDX analysis

The Co_3O_4 and mixed oxide $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ nanoparticles prepared by calcination of the $[\text{Co}/\text{Zn}(\text{II})\text{benzaldehyde}\cdot\text{piperazine}\cdot\text{H}_2\text{O}]$ complex at 500°C were characterized by EDX, SEM and TEM. The analysis data supported the formation of one type of aggregated nanoparticles with an average size of $15\text{--}31$ nm. The rectangular pyramidal morphology and size of the particles were further calculated by the XRD spectrum to validate the

TEM results. Polycrystalline Co_3O_4 and porous $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ particles were analysed by energy-dispersive X-ray (EDX) spectra of Co_3O_4 (Fig. 3) and mixed $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ (Fig. 4). In the spectra, Co and O peaks were detected in both Co_3O_4 and $\text{ZnO}\cdot\text{Co}_3\text{O}_4$, and Zn peaks were observed in the $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ spectra. This reveals that cobalt oxide aligned with ZnO to yield the composite. The EDX spectra of the oxides and element maps obtained showed the composition and mass percentages of $\sim 21\%$ oxygen and 76% metals for the elements O, Co, and Zn, respectively. Carbon signals originating from grid background were observed at a negligible percentage.

3.5. Morphologies of CoCo_3 and $\text{ZnO}\cdot\text{Co}_3\text{O}_4$

TEM and SEM images of nanomaterials prepared at 500°C were obtained. The shape of the particles depends on the chemical environment precipitation and aggregation method. Homo and hetero metal oxides aggregated to minimize the interfacial energy are cubic nanoparticles with an average particle size of $15\text{--}31$ nm (Fig. 5a–d). The morphology of Co_3O_4 , with an average diameter of 31.0 nm, in a regular crystalline phase indicated that CoCo_3 contains nanocrystals with a single shape. It was observed that the hexagonal crystalline porous ZnO cavity was uniformly filled with Co_3O_4 particles. The magnified SEM images show a uniform morphology (Fig. 6a–c). Porous hetero nanocrystals are important for electrochemical studies. To confirm the effect that the nature of the porous ZnO has on Co_3O_4 deposition, comparative cyclic voltammetry experiments were performed.

Transmission electron microscopy (TEM) images of the homo and hetero nano structure of Co_3O_4 and $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ (Fig. 5 and 6a, b) show low-magnification images of Co_3O_4 and mixed $\text{ZnO}\cdot\text{Co}_3\text{O}_4$. The magnified images of Co_3O_4 exhibited a rectangular pyramidal type, as shown in Fig. 9a, with directional edges. The clear shape of the crystals was studied by SEM. The $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ material was scanned by TEM, and it was



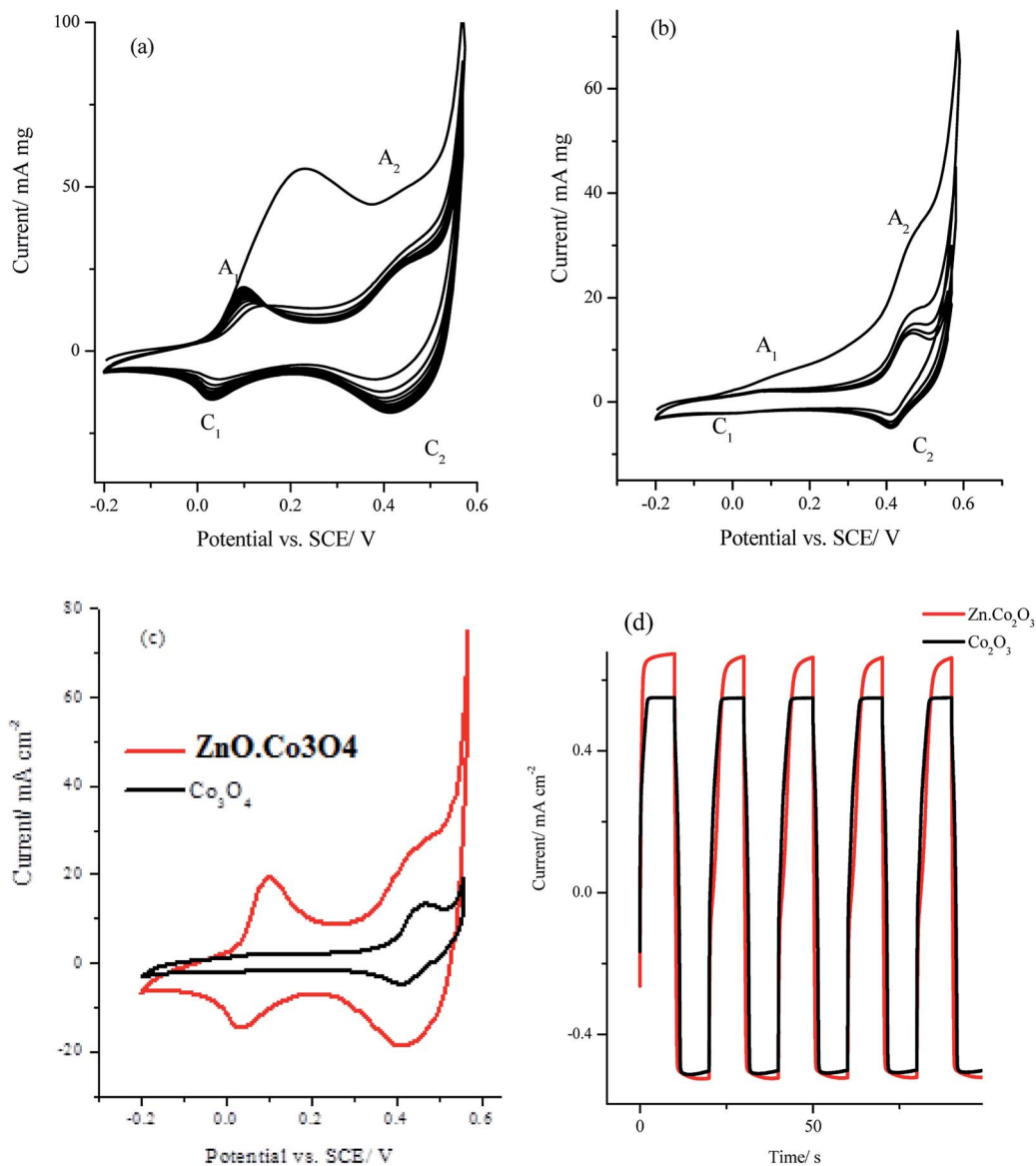


Fig. 8 Multi-cyclic voltammetry at 50 mV s^{-1} in 1.0 M KOH for (a) for $\text{ZnO} \cdot \text{Co}_3\text{O}_4$, (b) Co_3O_4 nanocrystals and (c) comparison for the stable cycle, (d) galvanostatic charge–discharge curves of $\text{ZnO} \cdot \text{Co}_3\text{O}_4$ and Co_3O_4 electrodes at charge current of 4.0 A g^{-1} .

observed that the hetero nanostructure is a porous conjugate of mixed oxide. Fig. 10a and b also reveal nanopolycrystalline Co_3O_4 deposited in the single crystalline ZnO porous cavities. The TEM observation was further validated by SEM analysis (Fig. 6c and d).²⁶

3.6. Paramagnetic character evaluation of nanoparticle Co_3O_4

NMR was employed to study the paramagnetic character of metal in nanophases in solution. The ^1H NMR chemical shift due to Co_3O_4 was monitored by Evan's method. Evan's NMR studies^{27,28} showed a paramagnetic chemical shift of ~ 0.8 ppm in the DMSO dimethyl proton signal at 2.49 ppm. The sharp change in the chemical shift at 3.3 ppm in the NMR spectrum clearly indicated that the particle exhibits paramagnetism

(Fig. 7). The small shift in the NMR signals indicates that Co_3O_4 nanoparticles have weak ferromagnetic behaviour.

3.7. Electrochemistry

The cyclic voltammetry (CV) measurements for Co_3O_4 and $\text{ZnO} \cdot \text{Co}_3\text{O}_4$ nanocrystals in 1.0 M KOH at a scan rate of 50 mV s^{-1} are depicted in Fig. 8. The cyclic voltammograms of the nanocrystals of Co_3O_4 and $\text{ZnO} \cdot \text{Co}_3\text{O}_4$ (Fig. 8a and b) exhibited two redox peaks of (A_1/C_1) and (A_2/C_2) that could be attributed to the redox couples $\text{Co(II)}/\text{Co(III)}$ and $\text{Co(III)}/\text{Co(IV)}$ and are located at the mid-peak potential of 65 and 430 mV vs. SCE, respectively.²⁹ However, for the hetero-bimetallic nano oxide $\text{ZnO} \cdot \text{Co}_3\text{O}_4$, the redox peaks were well resolved and had a significantly higher current than did the Co_3O_4 electrode as shown in Fig. 8c. Moreover, the oxidation current during the



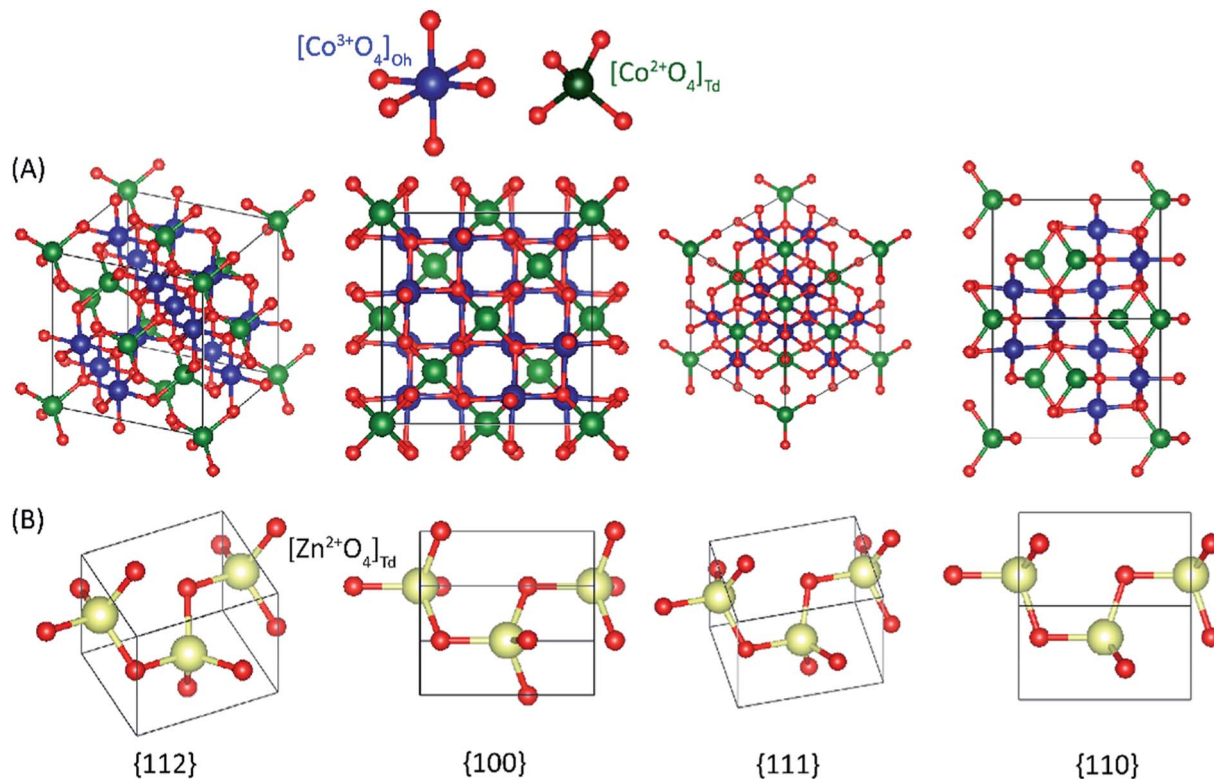


Fig. 9 The illustration of atom configurations of the {100}, {110}, {111}, and {112} crystal planes of the Co_3O_4 spinel and ZnO structure.

first anodic scan was much higher before it stabilized in the successive scans for both oxides, apparently due to a complete conversion of the $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ and Co_3O_4 to higher oxidized species, which was not completely reversible in the alkaline solution.^{30–32} The specific capacitance of the $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ and Co_3O_4 electrodes can be calculated from the CV using eqn (1):

$$C_s = Q/2m\Delta V \quad (1)$$

where C_s is the specific capacitance (F g^{-1}), Q is the volumetric charge under the CV in coulomb, m is the mass of the oxide materials in grams and ΔV is the potential window of the cyclic voltammetry. Using the cyclic voltammetry in Fig. 8c, the specific capacitance values were estimated to be 207.3 and 51.1 F g^{-1} for $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ and Co_3O_4 electrodes, respectively. Moreover, Fig. 8d illustrates the galvanostatic charge–discharge curves of $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ and Co_3O_4 electrodes at charge current of 4.0 A g^{-1} and in 1.0 M KOH solution. The specific capacitance obtained from each discharge curve is equal 210.5 and 54.6 F g^{-1} for $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ and Co_3O_4 electrodes respectively which is in good agreement with the values obtained from the cyclic voltammetry. These observations indicate that heterometallic $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ exhibited a much higher specific capacitance than did a monometallic Co_3O_4 nanocrystal, which could be attributed presumably to the mesoporous structure and higher surface area that allow fast ion diffusion and better conductivity as a result of ZnO incorporation into the Co_3O_4 structure.³³

3.8. Computational modelling

Ambient chemical transformations between nanoparticles of Co_3O_4 and ZnO leading to hybrid molecular adducts that preserve structure and topology are poorly explored area in material science. Atomically precise nanoparticles of Co and Zn metals, often called nanoclusters, which constitute an exploding discipline in nanomaterials because of their well-defined structures and drastic changes in their properties, in comparison to their bulk form, arising due to electronic confinement. Hence, theoretical calculations of $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ molecular aggregates have been performed in order to find out how closely they could approach each other and which atoms and their interactions would be involved in the adduct formation.

A complete search over the relevant rotational and translational degrees of freedom of Co_3O_4 nanocluster with respect to ZnO nanocluster in the $\text{Co}_3\text{O}_4\text{ZnO}$ adduct with DFT is unfeasible due to the computational cost. Therefore, we used a combined approach utilizing the highly efficient force-field based method to identify a global minimum energy geometry of the $\text{Co}_3\text{O}_4\text{ZnO}$ adducts, e.g. model 1: $[\text{Co}_3\text{O}_4]_4 [\text{ZnO}]_4$, model 2: $[\text{Co}_3\text{O}_4]_{10} [\text{ZnO}]_4$, and model 3: $[\text{Co}_3\text{O}_4]_{10} [\text{ZnO}]_{20}$ and then performed single point energy calculations using DFT method to calculate the electronic properties. We have used the reported crystal structure coordinates of Co_3O_4 and ZnO, without any structural relaxation, as the initial coordinates for geometry optimizations. The atom configurations of the {100}, {110}, {111}, and {112} crystal planes of the Co_3O_4 spinel and ZnO adducts are depicted in Fig. 9. Similarly, the crystal planes (100,



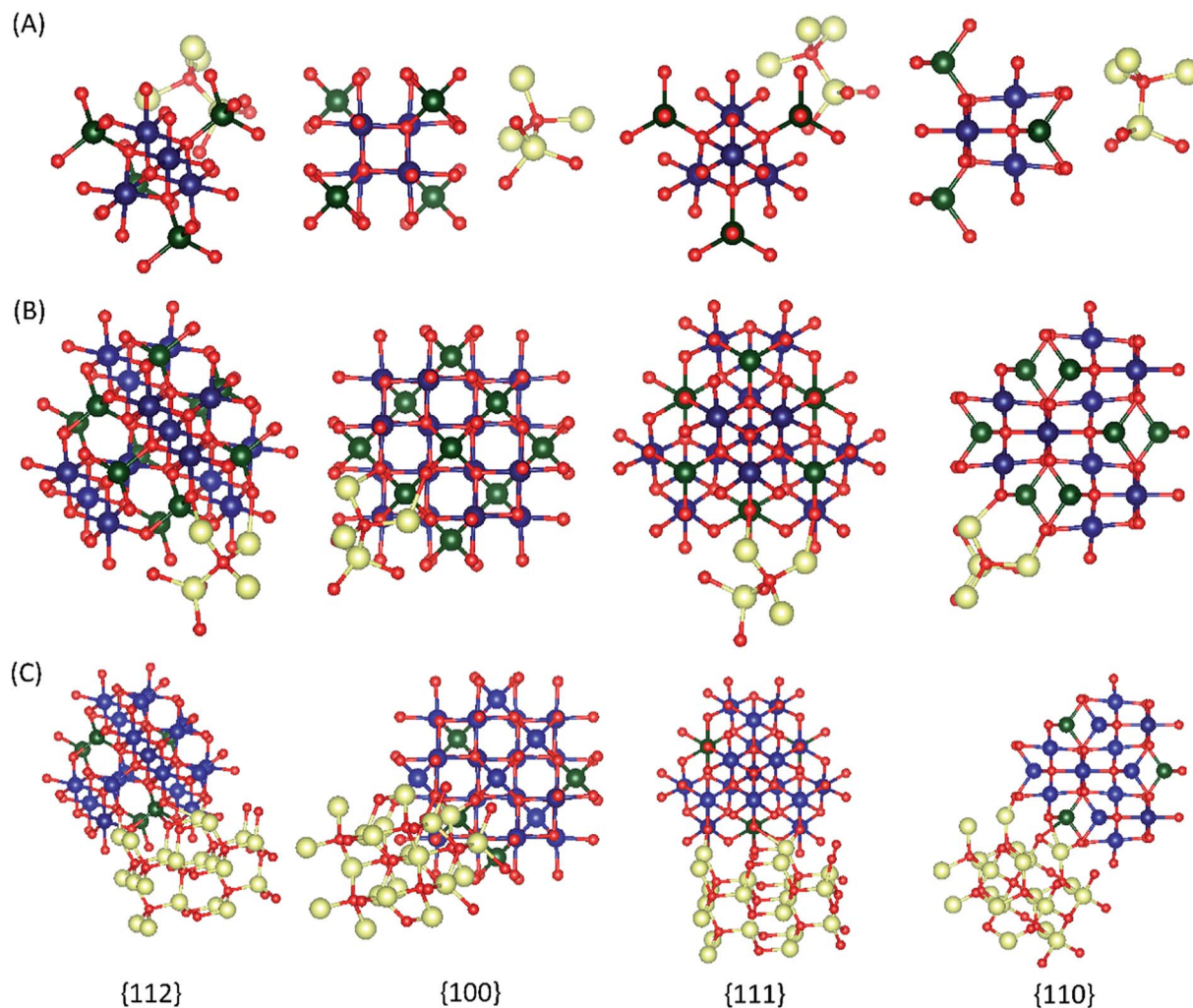


Fig. 10 The illustration of atom configurations of the {100}, {110}, {111}, and {112} crystal planes of the global minimum energy geometry of $\text{Co}_3\text{O}_4\text{ZnO}$ adducts, (A) model 1, (B) model 2, and (C) model 3.

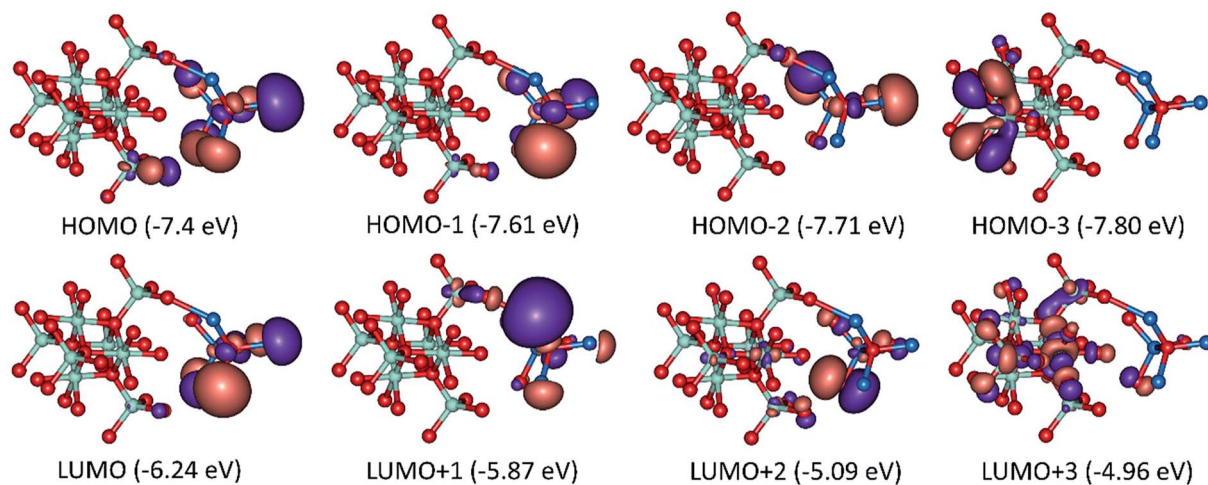


Fig. 11 Frontier molecular orbitals and their energies of model 1: $[\text{Co}_3\text{O}_4]_4 [\text{ZnO}]_4$ adduct.



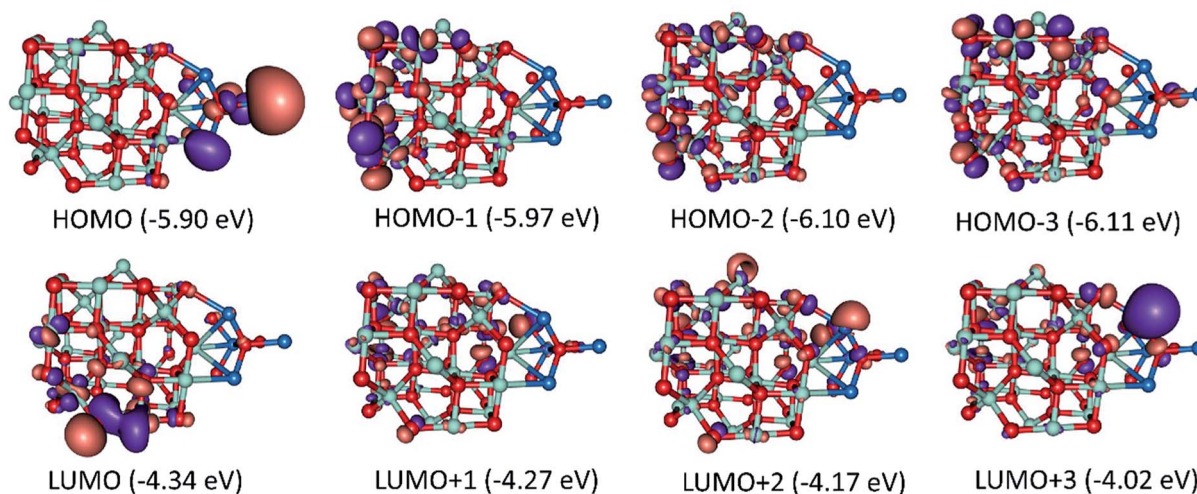


Fig. 12 Frontier molecular orbitals and their energies of model 2: $[\text{Co}_3\text{O}_4]_{10} [\text{ZnO}]_4$ adduct.

110, 111 and 112) of the global minimum energy geometries of the three adducts of Co_3O_4 and ZnO e.g. model 1: $[\text{Co}_3\text{O}_4]_4 [\text{ZnO}]_4$, model 2: $[\text{Co}_3\text{O}_4]_{10} [\text{ZnO}]_4$, and model 3: $[\text{Co}_3\text{O}_4]_{10}$

$[\text{ZnO}]_{20}$ are illustrated in Fig. 10 and S5.† From the force-field global minimum geometries (FFGMG) of adducts, we identified that the significant changes are observed in the

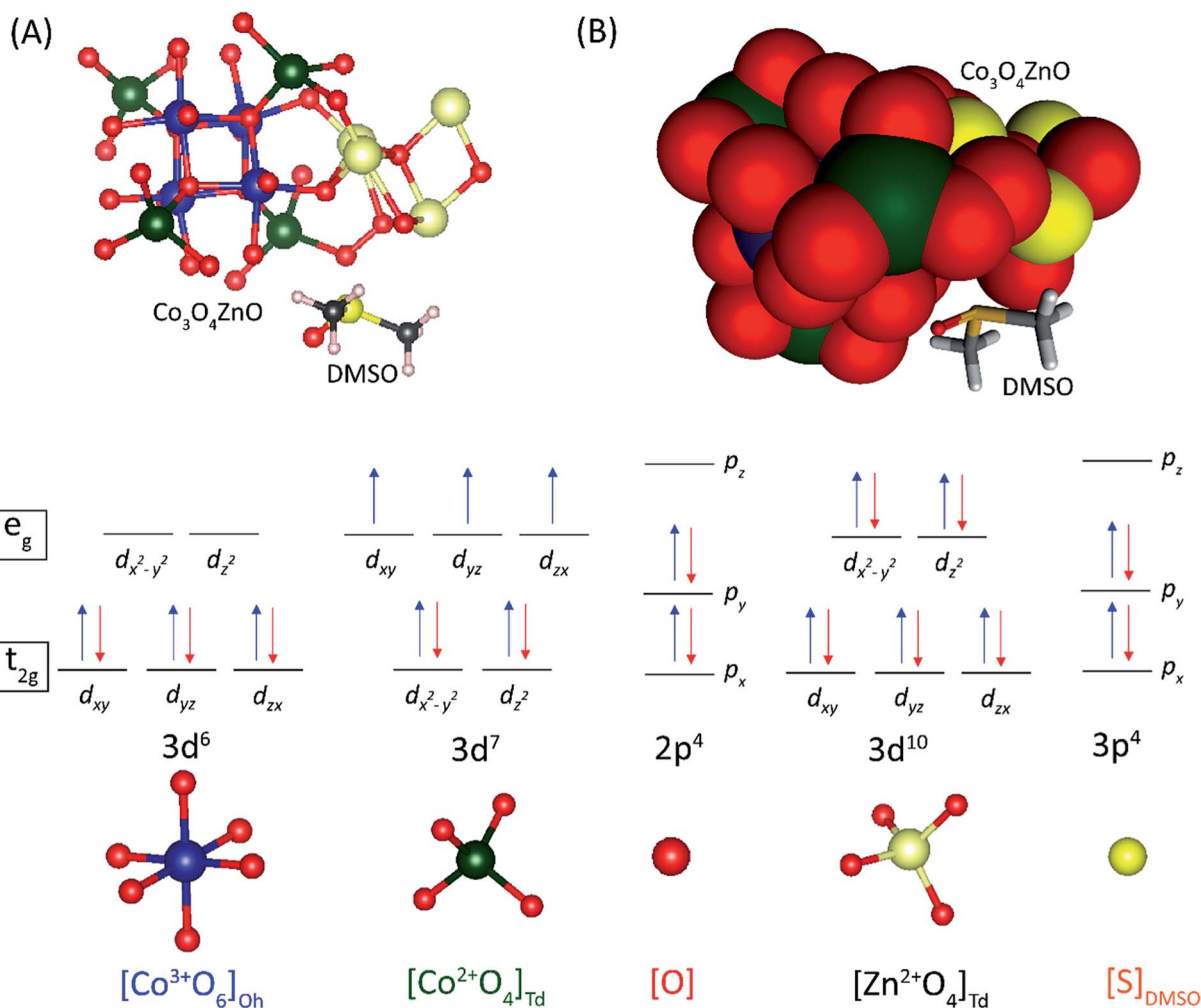


Fig. 13 The schematic representation of crystal field diagram of the global minimum energy geometry of $[\text{Co}_3\text{O}_4]_4 [\text{ZnO}]_4$ DMSO adduct.



orientations and distances between the Co_3O_4 and ZnO adducts. In model 1, no bond formation observed and the nearest distance between the Co_3O_4 and ZnO adducts is found to be 2.527 Å ($\text{Zn}\cdots\text{O}$). Whereas in model 2, Co_3O_4 and ZnO nanoclusters are approached so near to each other that they are linked together through the two Zn–O bonds (2.065 and 2.149 Å) between a bridging oxygen atom of $[\text{Co}_3\text{O}_4]_{10}$ and a zinc atom of $[\text{ZnO}]_4$ with Co–O–Zn oxo-linkage of 142.24° and 120° angles. Other two more close contacts with 2.24 Å also found between Zn and O atoms. Interestingly, in model 3, two different bonds, Zn–O: 1.935, 2.552 and 1.960 Å, between the zinc atoms of $[\text{ZnO}]_{20}$ and oxygen atoms of $[\text{Co}_3\text{O}_4]_{10}$; Co–O: 2.197 Å, between the cobalt atom of $[\text{Co}_3\text{O}_4]_{10}$ and oxygen atom of $[\text{ZnO}]_{20}$ are observed. The values of oxo-linkages angles Zn–O–Co: 144°, 149° and 111° and 94°: Co–O–Zn are observed. Additionally, a short O–O (1.463 Å) contact is also observed between the oxygen atoms of both adducts. Thus, from the computed structures of the adducts clearly suggested that as size of the interacted adducts increases more tightly they bound to each other *via* coordinate and covalent bonding and formed a single phase heterometallic nanomaterial. Further we have also demonstrated the corresponding frontier molecular orbitals of the model 1 and model 2, to explore the HOMO–LUMO gap and electronic properties (Fig. 11 and 12). The HOMO–LUMO gap is found to be 1.16 eV in model 1 while 1.56 eV in model 2. In the model 1, HOMO, HOMO–1, HOMO–2, LUMO, LUMO+1 and LUMO+2 are localized on the ZnO adduct while HOMO–3 and LUMO+3 on Co_3O_4 adduct (Fig. S5† and 11). Whereas in model 2, HOMO and LUMO are localized on the ZnO and Co_3O_4 adducts, respectively (Fig. 12).

We have also performed the geometry optimization of model 1 with the DMSO molecule, to find out the paramagnetic nature of the $\text{Co}_3\text{O}_4\text{ZnO}$ adduct which cause the shifting of $^1\text{H-NMR}$ peaks of DMSO. The Co_3O_4 structure comprises a cubic close-packed array of O^- where 1/8 of the tetrahedral interstices are occupied by high-spin Co^{2+} , whereas half of the octahedral interstices are occupied by low-spin Co^{3+} . Each Co^{2+} ($e_g^4 t_{2g}^3$) is surrounded by four nearest neighbours of opposite spin, giving rise to an antiferromagnetic network. In contrast, the Co^{3+} exhibited a closed-shell configuration (t_{2g}^6) and nil magnetic moment, as depicted in Fig. 13. The global minimum energy geometry of model 1 with DMSO is illustrated in Fig. 13. The global minimum energy geometry indicated that in the presence of DMSO molecule the ZnO and Co_3O_4 adducts joined through the various Zn–O–Co linkage. Interestingly, Zn atom of ZnO adduct joined through the oxygen atoms of both types of Co^{2+} (T_d) and Co^{3+} (O_h) ions of Co_3O_4 adduct (Fig. 13). Such kind of linkage may perturbed the antiferromagnetic coupling between the Co^{2+} (T_d) ions or ligand field stabilization, which cause the paramagnetic nature of the $\text{Co}_3\text{O}_4\text{ZnO}$ adduct and this paramagnetism cause the shifting of $^1\text{H-NMR}$ peaks of DMSO as observed experimentally.

4. Conclusion

The present method for preparing nanoparticle precursors for homo- and heterometallic cores is more superior to other

methods because we have used Werner's coordination theory reaction and reaction environment. One type of the nanocrystal was obtained, in comparison to other methods, which produce a mixture of nanoparticles with different shapes and sizes. This method is simple and suitable for nanocrystal design and tailoring. Co_3O_4 nanoparticles can be produced at a low temperature in the absence of solvent, surfactant and expensive or complicated equipment. The pure nanostructure Co_3O_4 and $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ particles with a size of 18–20 nm was successfully synthesized by the thermal decomposition of $[\text{M}(\text{II})\text{L}_2\text{L}']$ (where $\text{M} = \text{Co}(\text{II}), \text{Zn}(\text{II})$ $\text{L} = 4$ -hydroxy benzaldehyde and $\text{L}' =$ piperazine) complexes as new precursors. Nanoparticles were formed by redox reactions among the piperazine, benzaldehyde and counter ions. The rectangular/cubic rhomboid Co_3O_4 nanoparticles were obtained with agglomeration. DFT calculations support the formation and structure of the $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ and Co_3O_4 . The model has been generated to understand the nano structures. Weak ferromagnetic behaviour was observed in the NMR chemical shift of methyl proton signals. The specific capacitance of the $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ and Co_3O_4 electrodes was calculated from the CV; $\text{ZnO}\cdot\text{Co}_3\text{O}_4$ exhibited high capacitance which attributed to the better conductivity and surface area. DFT calculations with DMSO molecule further validated the experimental results displaying the paramagnetic nature of the $\text{Co}(\text{II})$ ion in Co_3O_4 adduct which caused the shifting of $^1\text{H-NMR}$ peaks of DMSO.

Conflicts of interest

There is no conflicts to declare.

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References

- 1 L. Hu, Q. Peng and Y. Li, *J. Am. Chem. Soc.*, 2008, **130**, 16136–16137.
- 2 G. Wang, H. Wang, W. Li and J. Bai, *RSC Adv.*, 2011, **1**, 1585–1592.
- 3 T. A. Taton, *Trends Biotechnol.*, 2002, **20**, 277–279.
- 4 R. Atchudan, T. N. J. I. Edison, D. Chakradhar, N. Karthik, S. Perumal and Y. R. Lee, *Ceram. Int.*, 2018, **44**, 2869–2883.
- 5 C.-W. Kung, C.-Y. Lin, Y.-H. Lai, R. Vittal and K.-C. Ho, *Biosens. Bioelectron.*, 2011, **27**, 125–131.
- 6 M. El Baydi, G. Poillerat, J.-L. Rehspringer, J. L. Gautier, J.-F. Koenig and P. Chartier, *J. Solid State Chem.*, 1994, **109**, 281–288.
- 7 J. Ma, S. Zhang, W. Liu and Y. Zhao, *J. Alloys Compd.*, 2010, **490**, 647–651.
- 8 A. Ruplecker, F. Kleitz, E.-L. Salabas and F. Schüth, *Chem. Mater.*, 2007, **19**, 485–496.
- 9 A. Askarinejad and A. Morsali, *Ultrason. Sonochem.*, 2009, **16**, 124–131.



- 10 T.-L. Lai, Y.-L. Lai, C.-C. Lee, Y.-Y. Shu and C.-B. Wang, *Catal. Today*, 2008, **131**, 105–110.
- 11 C. Sun, X. Su, F. Xiao, C. Niu and J. Wang, *Sens. Actuators, B*, 2011, **157**, 681–685.
- 12 P. Poizot, S. Laruelle, S. Grugeon, L. Dupont and J. M. Tarascon, *Nature*, 2000, **407**, 496.
- 13 C. Klingshirn, *ChemPhysChem*, 2007, **8**, 782–803.
- 14 A. Sulciute, J. Baltrusaitis and E. Valatka, *J. Appl. Electrochem.*, 2015, **45**, 405–417.
- 15 Z. Zhang, C. Shao, X. Li, C. Wang, M. Zhang and Y. Liu, *ACS Appl. Mater. Interfaces*, 2010, **2**, 2915–2923.
- 16 C. W. Na, H.-S. Woo, I.-D. Kim and J.-H. Lee, *Chem. Commun.*, 2011, **47**, 5148–5150.
- 17 M. Salavati-Niasari, A. Khansari and F. Davar, *Inorg. Chim. Acta*, 2009, **362**, 4937–4942.
- 18 G. M. Morris, R. Huey, W. Lindstrom, M. F. Sanner, R. K. Belew, D. S. Goodsell and A. J. Olson, *J. Comput. Chem.*, 2009, **30**, 2785–2791.
- 19 (a) F. Neese, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2012, **2**, 73–78; (b) F. Neese, *Orca, An ab Initio, Density Functional and Semiempirical Program Package version*, 2009.
- 20 C. Lee, W. Yang and R. G. Parr, *Phys. Rev. B*, 1988, **37**, 785–789.
- 21 (a) F. Weigend and R. Ahlrichs, *Phys. Chem. Chem. Phys.*, 2005, **7**, 3297–3305; (b) A. Schaefer, C. Huber and R. Ahlrichs, *J. Chem. Phys.*, 1994, **100**, 5829–5835; (c) A. Schaefer, H. Horn and R. Ahlrichs, *J. Chem. Phys.*, 1992, **97**, 2571–2577.
- 22 (a) S. Grimme, J. Antony, S. Ehrlich and H. Krieg, *J. Chem. Phys.*, 2010, **132**, 154104; (b) C. Steffen, K. Thomas, U. Huniar, A. Hellweg, O. Rubner and A. Schroer, *J. Comput. Chem.*, 2010, **31**, 2967–2970.
- 23 B. Pejova, A. Isahi, M. Najdoski and I. Grozdanov, *Mater. Res. Bull.*, 2001, **36**, 161–170.
- 24 R. K. Sharma and R. Ghose, *J. Alloys Compd.*, 2016, **686**, 64–73.
- 25 B. D. Cullity and S. R. Stock, *Elements of X-ray Diffraction*, Prentice-Hall, New York, 3rd edn, 2001.
- 26 Y. Tak and K. Yong, *J. Phys. Chem. C*, 2008, **112**, 74–79.
- 27 D. F. Evans, *J. Chem. Soc.*, 1959, 2003–2005.
- 28 S. K. Sur, *J. Magn. Reson.*, 1989, **82**, 169–173.
- 29 N. Spataru, C. Terashima, K. Tokuhiko, I. Sutanto, D. A. Tryk, S.-M. Park and A. Fujishima, *J. Electrochem. Soc.*, 2003, **150**, E337–E341.
- 30 W.-J. Zhou, M.-W. Xu, D.-D. Zhao, C.-L. Xu and H.-L. Li, *Microporous Mesoporous Mater.*, 2009, **117**, 55–60.
- 31 J. Xu, L. Gao, J. Cao, W. Wang and Z. Chen, *Electrochim. Acta*, 2010, **56**, 732–736.
- 32 V. Srinivasan and J. W. Weidner, *J. Power Sources*, 2002, **108**, 15–20.
- 33 Y. Shang, T. Xie, C. Ma, L. Su, Y. Gai, J. Liu and L. Gong, *Electrochim. Acta*, 2018, **286**, 103–113.

