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photocatalytic activity for the degradation of

Synthesis of $Cu_x Co_{3-x}O_4$ nanoparticles by

structural and optical properties and

methylene blue

a sonochemical method and characterization of

In the present investigation, $Cu_xCo_{3-x}O_4$ (x = 0, 0.02, 0.04, 0.06, 0.1) nanoparticles have been synthesized by the sonochemical method and subsequent calcination at high temperature. The synthesized nanoparticles were further characterized using X-ray diffraction, UV-vis spectroscopy, FT-IR spectroscopy, Raman spectroscopy, transmission electron microscopy and energy-dispersive X-ray spectroscopy. The analysis results showed that nanoparticle size, band gap and photocatalytic activity of $Cu_xCo_{3-x}O_4$ compounds change by altering the Cu doping content. According to the XRD study, no impurity peaks associated with copper or copper oxide phases were observed in the diffractograms of doped samples. The highest degradation of methylene blue dye under visible light was observed in the presence of $Cu_{0.02}Co_{2.98}O_4$ nanoparticles and found to be 87.51% after 330 minutes. This value increased with increasing the amount of photocatalyst and with decreasing dye concentration.

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1 Introduction

Cobalt oxide (Co_3O_4) nanoparticles are one of the interesting metal oxides because they have unique physical and chemical properties. They are widely used in gas sensors, pH sensors, catalysts, anode materials in Li-ion rechargeable batteries, supercapacitors, electrochemical sensors, photocatalysts, solar absorbers and electrochromic materials.¹⁻⁹ Co₃O₄ is a p-type antiferromagnetic semiconductor and mixed valence compound ($CoO \cdot Co_2O_3$). It adopts the normal spinel structure. So far, several synthesis methods have been employed to produce Co₃O₄ nanoparticles including hydrothermal/solvothermal,^{10,11} precipitation,¹² sonochemical,¹³ sol-gel,¹⁴ solution combustion,15 successive ionic layer adsorption and reaction (SILAR),¹⁶ chemical spray pyrolysis¹⁷ and thermal decomposition.¹⁸ According to reports, the incorporation of metal dopants is an effective method for improving electrical conductivity and catalytic activity of oxide nanoparticles. Rehman et al.19 prepared Ce and Zn doped CuO nanoparticles by co-

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precipitation method and a remarkable increment in degradation efficiency for methylene blue (81.64%) under visible light was observed in the presence of these nanocatalysts compared to undoped samples (38.77%). Degradation efficiency was found to be 78% for methylene blue dye under UV-A light using 3% Cu-doped NiO nanoparticles obtained by green synthesis combined with a sol-gel method.20 Several metals are used to modify the Co₃O₄ nanoparticles including Fe, Mn, Cu, Ni, Mo,^{20,21} Ag, and Pd.^{22,23} Jincy and Meena synthesized copperdoped cobalt oxide nanoparticles by a hydrothermal method and they observed that these nanoparticles exhibited extreme sensitivity towards NH₃ gas at room temperature even at a lower concentration of 5 ppm in comparison to undoped Co₃O₄.²⁴ In another work, Cu-doped Co₃O₄ nanosheets obtained by a solvothermal method show enhanced electrochemical performance due to the reduced particle size and larger surface area.25 Dong and co-workers fabricated mesoporous Ag-doped Co₃O₄ nanowire arrays supported on an FTO substrate by an electrodeposition-hydrothermal process for water oxidation in 0.5 M H₂SO₄ and it has been determined that the addition of silver as a dopant significantly improves both the conductivity and stability of the electrocatalyst in an acidic environment.26

As far as we know, there have been no previous reports in the literature regarding the synthesis of Cu-doped Co_3O_4 nanoparticles using the sonochemical method. In this study, we carried out the synthesis of undoped Co_3O_4 nanoparticles and

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various concentrations of Cu-doped Co_3O_4 nanoparticles using the sonochemical method, followed by thermal treatment at high temperature. For synthesis, we chose the sonochemical method due to its simplicity and low cost. We selected copper as a dopant for our study because the ionic radius of Cu^{2+} is close to the ionic radius of Co^{2+} and therefore Cu^{2+} ions can easily enter into the Co_3O_4 matrix. The influence of dopant concentration on the structural, optical properties and photocatalytic activity of Co_3O_4 nanoparticles was examined in detail.

2 Experimental

2.1 Materials and instrumentation

All chemicals [cobalt nitrate $(Co(NO_3)_2 \cdot 7H_2O)$, copper(II) chloride(CuCl₂ · 2H₂O), sodium hydroxide (NaOH), polyvinyl alcohol (PVA), hydrogen peroxide (H₂O₂) and methylene blue (MB) (C₁₆H₁₈ClN₃S) were of analytical grade and were used without any further purification. Bath type ultrasonicator (VCX500) was used for the synthesis of Co₃O₄ nanoparticles.

The structural analysis of the synthesized oxide powders was analyzed using Rigaku Mini Flex 600 X-ray diffractometer ($\lambda =$ 1.54060 Å) using Ni-filtered Cu K α radiation with a step width of 0.1° and a scan speed/duration time of 5° per minute over a range of 2 θ from 10° to 90°. Optical properties of the samples were assessed using Specord 250 Plus UV-vis Spectrophotometer. Fourier transform infrared (FTIR) measurement has been carried out within 400–4000 cm⁻¹ region on a Varian 3600 FTIR spectrometer. Transmission electron microscopy (TEM) measurements were conducted on the JEOL JEM-1400 transmission electron microscope. EDX study was carried out on X-Max Energy dispersive spectrometer. Raman spectroscopy measurements were recorded on an EnSpectr R532 spectrometer in the spectral range of 100–800 cm⁻¹.

2.2 Preparation of undoped and Cu-doped Co₃O₄ nanoparticles

Undoped Co_3O_4 nanoparticles were synthesized by the sonochemical method and subsequent calcination process with some modifications reported in our previous research.²⁷ In this experiment, sodium hydroxide was used instead of sodium borohydride as a precipitating agent. For the synthesis, 100 ml of 0.1 M $\text{Co}(\text{NO}_3)_2 \cdot 7\text{H}_2\text{O}$ aqueous solution was prepared and stirred with PVA solution as a stabilizer agent (2.0 wt%) for 15 min to obtain a homogenous solution. Then, as-prepared 100 ml of 0.2 M NaOH solution was added drop by drop into the above solution under the sonication process using an ultrasonic probe. This solution was irradiated with 20 kHs ultrasound wave for an hour. The pink precipitate was obtained at the end of the reaction. Finally, the precipitate was centrifuged at 6000 rpm, washed three times with distilled water and ethanol in the end, and dried at room temperature. Thereafter, the product was annealed at 500 °C for 4 h in a Muffle furnace and the color of it turned black powder. For preparing the different concentrations of Cu-doped Co3O4 nanoparticles $(Cu_x Co_{3-x}O_4, x = 0.02, 0.04, 0.06, 0.1)$, a stoichiometric amount of CuCl₂·2H₂O was dissolved in 100 ml distilled water and added to cobalt nitrate solution. The rest of the experiments is the same as described above. The schematic representation of the synthesis process of $Cu_x Co_{3-x}O_4$ nanoparticles is given in Fig. 1.

2.3 Photocatalytic activity measurement

The photocatalytic activity of undoped Co₃O₄ and Cu-doped Co_3O_4 nanoparticles was investigated on methylene blue (MB) dye under visible light irradiation. For experiments, initially, 10 mg catalyst was added to the 10 ppm dye solution. Before irradiation, the stirring was performed in the dark for 45 min to get an adsorption-desorption equilibrium between the dye and catalyst. Then 0.4 ml of 450 mM hydrogen peroxide (H2O2) solution as an oxidant was added to the solution and the solution was exposed to visible light irradiation on a magnetic stirrer. 116 W halogen lamp was used as a source of visible light. The experiments were performed at room temperature. At predetermined time intervals, 3 ml of dye solution was taken out from the beaker and centrifuged at 6000 rpm for 5 min to completely precipitate particles. Then UV-vis adsorption spectra were measured in the range of 400-800 nm. The experiments were also performed by varying the initial dye concentration (5 ppm and 20 ppm) and amount of catalyst (20 mg and 30 mg). Degradation efficiency was calculated by Beer-Lambert law using the following equation:22,28



Fig. 1 Schematic representation of the experimental work.

Degradation efficiency
$$(R) = \frac{C_0 - C}{C_0} \times 100 = \frac{A_0 - A}{A_0} \times 100\%$$
(1)

where C_0 and A_0 -initial concentration and absorbance of dye solution, C and A-concentration and absorbance of dye solution after the time interval, respectively.

3 Results and discussion

3.1 Structural analysis

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To elucidate the formation mechanism of Cu_rCo_{3-r}O₄ nanoparticles, structural analysis was conducted before thermal annealing. Fig. 2a shows the XRD patterns of undoped and different concentrations (2%, 4%, 6% and 10%) of Cu-doped samples before thermal annealing. The low-intensity peaks occurred at $2\theta = 19.23^{\circ}$, 38.27° corresponding to (001) and (101) planes in the diffractogram of the undoped sample. These results match the JCPDS-30-0443 standard data, confirming the formation of a brucite-like phase of β -Co(OH)₂ with a hexagonal crystal structure. It should be noted, there were no diffraction peaks attributed to α -Co(OH)₂, Cu(OH)₂, or impurities, signifying a high-purity final product. The size and interplanar spacing of the β -Co(OH)₂ nanoparticles were determined as 5.17 nm and 4.61183 Å, respectively, based on the most intense peak corresponding to the (001) plane. The lattice parameters were calculated as follows: a = b = 3.17 Å, c = 4.61 Å, and $\alpha =$ $\beta = 90^{\circ}$, $\gamma = 120^{\circ}$. The Debye–Scherrer formula was employed to calculate the crystallite size:29,30

$$D = 0.89\lambda/\beta\cos\theta \tag{2}$$

where θ is Bragg's angle, λ and β are the X-ray wavelength and the full width at half maximum intensity, respectively.

Interplanar spacing (d) is evaluated by Bragg's Law equation:³⁰

$$d_{h,k,l} = \frac{n\lambda}{2\sin\theta} \tag{3}$$

where *n* is an integer, θ -is a diffraction angle and λ is the wavelength of X-rays.

The lattice constants of β -Co(OH)₂ nanoparticles with a hexagonal crystal structure are calculated by the following equation:³¹

$$\frac{1}{d_{h,k,l}^2} = \frac{4}{3} \left(\frac{h^2 + hk + k^2}{a^2} \right) + \frac{l^2}{c^2}$$
(4)

The values of lattice parameters "a" and "c" were calculated from (101) and (001) reflections, respectively, and the obtained values are consistent with the standard data.

One diffraction peak appeared at $2\theta = 38.36^{\circ}$ corresponding to the (101) plane in the XRD pattern of 2% Cu doped sample before thermal annealing. In the XRD pattern of 4% Cu doped sample, the small intensity peaks appearing at $2\theta = 19.36^{\circ}$, 38.52° and 51.68° values are ascribed to the (001), (101) and (102) crystal planes, respectively. One diffraction peak at $2\theta =$ 38.64° corresponding to (101) plane appeared in the diffractogram of the 6% Cu doped sample, two diffraction peaks at $2\theta =$ 19.65° and 38.06° values corresponding to (001) and (101) planes were observed in the diffractogram of the 10% Cu doped sample. Before thermal annealing, the XRD patterns of Cudoped samples show peaks associated with β -Co(OH)₂, with no observable peaks related to Cu(OH)2. The results reveal that as the concentration of copper increases in doped β -Co(OH)₂, the diffraction peak corresponding to the (001) plane shifts to a higher angle. The size and interplanar spacing of 4% Cu doped β -Co(OH)₂ nanoparticles were estimated as 5.26 nm and



Fig. 2 XRD patterns of undoped and 2%, 4%, 6%, 10% Cu doped (a) β -Co(OH)₂ and (b) Co₃O₄ nanoparticles.

4.57881 Å, respectively. However, the peak corresponding to the (001) plane was not detected in 2% and 6% Cu-doped $Co(OH)_2$ nanoparticles, and the intensity of this and other peaks in 10% Cu-doped $Co(OH)_2$ nanoparticles was minimal, preventing an accurate size determination.

Fig. 2b illustrates the XRD patterns of undoped and various concentrations (2%, 4%, 6% and 10%) of Cu-doped samples after thermal annealing at 500 °C for 4 h. (111), (220), (311), (222) and (440) reflection planes were observed at $2\theta = 18.91^{\circ}$, 31.25° , 36.84° , 38.56° , 65.19° diffraction angles in the XRD pattern of the undoped sample. This result aligns well with JCPDS-00-009-0418 standard data, confirming the formation of Co₃O₄ with a cubic crystal structure. The space group is identified as Fd-3m (227), and the lattice parameters are a = b = c = 8.0849 Å; $\alpha = \beta = \gamma = 90^{\circ}$. The lattice constant (*a*) was calculated using the interplanar spacing value with the following equation,³² yielding a value close to the standard (*a* = 8.084 Å):

$$d = \frac{a}{\sqrt{h^2 + k^2 + l^2}}$$
(5)

The lattice constant and nanoparticle size were determined using the most intense peak in the diffractogram, which corresponds to the (311) plane. The crystallite size of undoped, 2%, 4%, 6% and 10% Cu-doped Co_3O_4 nanoparticles is 22.55 nm, 29.17 nm, 21.87 nm, 34.19 nm and 31.26 nm, respectively. It can be observed that the size of Co_3O_4 nanoparticles undergoes irregular changes with increasing Cu concentration.

No additional peaks related to the copper or copper oxide were detected in the diffractograms of Cu-doped Co₃O₄ nanoparticles. This observation suggests that Cu atoms effectively replaced Co atoms, leading to a well-dispersed distribution of Cu within the Co₃O₄ lattice.^{33,34} The similarity in ionic radii between Co^{2+} (0.72 Å) and Cu^{2+} (0.71 Å) in tetrahedral coordination facilitates the easy incorporation of Cu²⁺ ions into the Co_3O_4 matrix, substituting the Co^{2+} ions. As a result of Cu substitution, the lattice constant of the Co₃O₄ nanoparticles decreased compared to the undoped samples. Furthermore, the peaks in the diffractograms of Cu-doped Co₃O₄ nanoparticles exhibited a shift towards higher angles compared to the undoped sample. To further investigate the impact of Cu doping on the Co_3O_4 nanoparticles, lattice strain (ε), dislocation density (δ) , and unit cell volume were calculated from the XRD data using the equations provided, and the corresponding values are presented in Table 1.

$$\epsilon = \frac{\beta}{4\tan\theta} \tag{6}$$

$$\delta = \frac{n}{D^2} \tag{7}$$

$$V = a^3 \tag{8}$$

As the nanoparticle size increases, the lattice strain value decreases, and conversely, as the nanoparticle size decreases, the lattice strain value increases. Below are the possible chemical reactions that could occur:

$$Co(NO_3)_2 \cdot 7H_2O \rightarrow Co^{2+} + 2NO_3^- + 7H_2O$$

$$NaOH \rightarrow Na^+ + OH^-$$

$$Co^{2+} + 2OH^- + xH_2O \rightarrow Co(OH)_3 \cdot xH_2O$$

$$\operatorname{Co}(\operatorname{OH})_2 \cdot x \operatorname{H}_2 \operatorname{O} \xrightarrow{t} \operatorname{Co}_3 \operatorname{O}_4 + x \operatorname{H}_2 \operatorname{O}$$

$$\operatorname{CuCl}_2 \cdot 2\operatorname{H}_2\operatorname{O} \to \operatorname{Cu}^{2+} + 2\operatorname{Cl}^- + 2\operatorname{H}_2\operatorname{O}$$
$$\operatorname{Co}^{2+} + \operatorname{Cu}^{2+} + 2\operatorname{OH}^- + x\operatorname{H}_2\operatorname{O} \to \operatorname{Cu}_x\operatorname{Co}_{1-x}(\operatorname{OH})_2 \cdot x\operatorname{H}_2\operatorname{O}$$

$$\operatorname{Cu}_{x}\operatorname{Co}_{1-x}(\operatorname{OH})_{2} \cdot x\operatorname{H}_{2}\operatorname{O} \xrightarrow{t} \operatorname{Cu}_{x}\operatorname{Co}_{3-x}\operatorname{O}_{4} + x\operatorname{H}_{2}\operatorname{O}_{5}$$

3.2 UV-vis spectroscopy

The optical absorption spectra of undoped and different concentrations of Cu-doped Co_3O_4 nanoparticles were recorded at room temperature in the wavelength range of 250 nm and 900 nm. To perform UV-vis spectroscopy measurements, polyvinyl alcohol (PVA) films were prepared based on all synthesized samples by mechanical stirring. Fig. 3a shows the optical absorption spectra of all samples thermal annealed at 500 °C. Three absorption peaks in the ultraviolet, visible and near-infrared regions were observed for all samples. These peaks were found at 272.2 nm, 469.2 nm, and 773.8 nm in the spectrum of undoped Co_3O_4 nanoparticles. The peak at 272.2 nm is attributed to the PVA. As for the Cu-doped Co_3O_4 nanoparticles,

Table 1 The values of crystal structure parameters of $Cu_xCo_{3-x}O_4$ (x = 0, 0.02, 0.04, 0.06, 0.1) nanoparticles

Sample	2θ (°)	Crystallite size (nm)	Interplanar distance (Å)	Lattice strain	Dislocation density (10 ⁻³)	Lattice constant $(a = b = c)$ (Å)	Unit cell volume (Å) ³
Undoped Co ₃ O ₄	36.84	22.55	2.43768	0.0051	1.967	8.0849	528.47
$Cu_{0.02}Co_{2.98}O_4$	36.94	29.17	2.43131	0.0039	1.175	8.0637	524.33
Cu _{0.04} Co _{2.96} O ₄	36.86	21.87	2.43653	0.0052	2.091	8.0810	527.71
Cu _{0.06} Co _{2.94} O ₄	36.94	34.19	2.43131	0.0033	0.855	8.0637	524.33
Cu _{0.1} Co _{2.9} O ₄	36.95	31.26	2.43093	0.0037	1.023	8.0625	524.09



Fig. 3 (a) The absorption spectrums and (b) the band gap energy plots for $Cu_xCo_{3-x}O_4$ nanoparticles; (c) FT-IR spectra of undoped (2) and 10% Cu doped Co_3O_4 nanoparticles (1).

the absorption peaks were detected at the following wavelengths: 449.4 nm and 762 nm for 2% Cu-doped Co_3O_4 , 485.6 nm and 777.8 nm for 4% Cu-doped Co_3O_4 , 491.2 nm and 778.2 nm for 6% Cu-doped Co_3O_4 , and 534 nm and 826.4 nm for 10% Cu-doped Co_3O_4 nanoparticles. The results indicate that a blue shift, or a shift towards a lower value of wavelength, was observed in the absorption curve of 2% Cu-doped Co_3O_4 nanoparticles compared to undoped Co_3O_4 nanoparticles. Conversely, the absorption curves of 4%, 6%, and 10% Cudoped Co_3O_4 nanoparticles showed a red shift, or a shift towards a higher value of wavelength, compared to undoped Co_3O_4 nanoparticles. The optical band gap values (E_g) of samples have been determined using the following Tauc's equation:

$$\alpha = [B(h\nu - E_g)^n]/h\nu \tag{9}$$

where α is the absorption coefficient, $h\nu$ is the photon energy, *B* is a constant, and the value of *n* depends on the nature of transition: n = 2 for indirect allowed transition and n = 1/2 for direct allowed transition, n = 3/2 for direct forbidden transition, n = 3 for indirect forbidden transition. The $(\alpha h\nu)^2$ versus $h\nu$ curves for samples annealed at 500 °C are shown in Fig. 3b. It

reveals two band gaps associated with the electronic band structure of cobalt oxide. The band gap energy of samples was estimated by extrapolating the linear part of the curves to zero absorption ($\alpha = 0$).³⁵ The direct E_g values are given in Table 2. The presence of two band gaps is ascribed to the electronic band structure of cobalt oxide. The first band gap (E_{g1}) is attributed to the O²⁻ Co³⁺ charge transfer (with the Co³⁺ level located below the conduction band), whereas the higher band gap (E_{g2}) is assigned to the O²⁻ Co²⁺ charge transfer process.^{35,36}

The E_g value of pure Co₃O₄ nanoparticles increased from 1.58 eV and 3.33 eV to 1.70 eV and 3.43 eV when doped with 2% Cu. However, with higher Cu content, the band gap decreased.

Table 2 The band gap energy values and degradation efficiency in 330 min for $Cu_xCo_{3-x}O_4$ nanoparticles

Sample	The band gap (eV) (for samples annealed at 500 °C)	Degradation efficiency (%)
Undoped Co ₃ O ₄	1.58 and 3.33	79.20
$Cu_{0.02}Co_{2.98}O_4$	1.70 and 3.43	87.51
Cu _{0.04} Co _{2.96} O ₄	1.45 and 3.26	74.66
Cu _{0.06} Co _{2.94} O ₄	1.44 and 3.21	85
Cu _{0.1} Co _{2.9} O ₄	1.26 and 3.08	65.60

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The increase in the band gap (E_g) of 2% Cu-doped Co₃O₄ nanoparticles can be attributed to the Moss–Burstein effect. The energy band broadening occurs when the Fermi level lifts into the conduction band according to Moss–Burstein effect. The decrease in band gap at higher Cu dopant concentrations is associated with crystal defects. The incorporation of more Cu²⁺ ions into the Co₃O₄ lattice leads to an increase in defects.³⁷ When Cu²⁺ ions enter the crystal lattice of cobalt oxide in large quantities, the obtained mixed oxide shows combined physicochemical characteristics of both copper oxide and cobalt oxide.

3.3 FT-IR spectroscopy

IR spectra of undoped and 10% Cu-doped Co_3O_4 nanoparticles are given in Fig. 3c and d, respectively. Two sharp absorption peaks appeared at 661 cm⁻¹ (v_1) and 568 cm⁻¹ (v_2) in the spectrum of undoped Co_3O_4 nanoparticles. The v_1 band is attributed to the tetrahedral coordinated Co^{2+} ions, while the v_2 band is assigned to the octahedral coordinated Co^{3+} ions.³⁸ These characteristic peaks were observed at 662 cm⁻¹ and 572 cm⁻¹ in the spectrum of 10% Cu-doped Co_3O_4 manoparticles. So, this result confirms the formation of Co_3O_4 with spinel structure. The absence of a peak related to Cu–O bond in the spectrum of Cu-doped Co_3O_4 nanoparticles can be explained by the overlapping of Cu–O peak with Co–O peak due to their closely similar vibration wavelengths.^{25,39}

3.4 TEM and EDX studies

TEM measurements were conducted to determine the size and morphology of obtained samples. The TEM images of pure Co_3O_4 nanoparticles are presented in Fig. 4A. For TEM measurements, the sample was dispersed in ethanol by ultrasonic vibration and was dropped onto the surface of carboncoated copper grids. It can be seen that nanoparticles show spherical-like, hexagonal-like and irregularly structured morphology and the average diameter of Co_3O_4 nanoparticles is in the range of 5.57–79.15 nm. Fig. 4B displays TEM images of 10% Cu-doped Co_3O_4 nanoparticles. These nanoparticles exhibit various morphologies, including spherical-like, hexagonal-like, rhombohedral-like, and irregular structures. The size of nanoparticles is in the range of 15.78–78.31 nm.

To determine the elemental composition, stoichiometry and purity of samples EDX analysis was carried out. EDX analysis results of undoped Co_3O_4 nanoparticles (Fig. 5A) indicate the presence of cobalt and oxygen elements, with each element constituting 50% of the composition. The EDX spectrum depicted in Fig. 5B confirms the presence of cobalt, copper and oxygen elements in the synthesized 2% Cu-doped Co_3O_4 nanoparticles and atomic percentages of these elements are 48.43%, 0.31% and 50.32%, respectively. Based on this result, the percentage ratio of Cu/Co atoms is found to be 0.006, which this value is close to the theoretical ratio of Cu/Co atoms (0.007) in the $Cu_{0.02}Co_{2.98}O_4$ sample. For 4% Cu-doped Co_3O_4



Fig. 4 TEM images of (A) undoped Co_3O_4 and (B) 10% Cu-doped Co_3O_4 nanoparticles.



Fig. 5 Element mapping and EDX spectrum of (A) undoped Co₃O₄ and (B) 2% Cu-doped Co₃O₄ nanoparticles.

nanoparticles (Fig. 6A), atomic percentages of cobalt, copper and oxygen elements are 49.47%, 0.53% and 50%, respectively. The percentage ratio of Cu/Co atoms is equal to 0.01 and this value coincides with the theoretical ratio of Cu/Co atoms (0.01) in the Cu_{0.04}Co_{2.96}O₄ compound. For 6% Cu-doped Co₃O₄ nanoparticles (Fig. 6B), atomic percentages of cobalt, copper and oxygen elements are 48.98%, 1.02% and 50%, respectively. According to the result, the percentage ratio of Cu/Co atoms is equal to 0.02 and this value coincides with the theoretical ratio of Cu/Co atoms (0.02) in the Cu_{0.06}Co_{2.94}O₄ compound.

EDX analysis results of 10% Cu-doped Co_3O_4 nanoparticles (Fig. 6C) reveal the presence of cobalt, copper, oxygen, sodium, chlorine elements and the atomic percentages of these

elements are 43.97%, 2.90%, 48.57%, 3.39% and 1.17%, respectively. The occurrence of trace amounts of sodium and chlorine elements can be attributed to the initial reagents utilized in the synthesis process. Based on the results, the percentage ratio of Cu/Co atoms is equal to 0.06, which is close to the theoretical ratio of Cu/Co atoms (0.03) in the Cu_{0.1}Co_{2.9}O₄ sample. When examining the element mapping of the Cudoped Co₃O₄ nanoparticles, it becomes evident that the distribution of Co, Cu, and O elements coincides.

3.5 Raman spectroscopy

The Raman spectra of pure, 2% Cu-doped Co_3O_4 and 10% Cudoped Co_3O_4 nanoparticles are presented in Fig. 7. Three

Fig. 6 Element mapping and EDX spectrum of (A) 4% Cu-doped, (B) 6% Cu-doped and (C) 10% Cu-doped Co₃O₄ nanoparticles.

Raman peaks at 184 cm⁻¹, 464 cm⁻¹ and 666 cm⁻¹ are observed for pure Co₃O₄ nanoparticles. These peaks are correspond to the F_{2g} , E_g and A_{1g} modes, respectively, which they are in good agreement with the reported literatures.40-42 The strong band at 666 cm⁻¹ (A_{1g} mode) is assigned to the characteristics of the octahedral sites, while F_{2g} and E_g modes are attributed to the combined vibrations of the tetrahedral site and octahedral oxygen motions.⁴² As can be seen from the figure, the intensity of Raman peaks decreased after doping of Co3O4 with Cu. The Raman peaks at 514 cm⁻¹ and 674 cm⁻¹ are observed for 2% Cu-doped Co₃O₄ nanoparticles and the Raman peaks at 512 cm⁻¹ and 674 cm⁻¹ are observed for 10% Cu-doped Co₃O₄

Fig. 7 Raman spectra of (a) undoped, (b) 2% Cu-doped Co₃O₄ and (c) 10% Cu-doped Co₃O₄ nanoparticles.

nanoparticles. A slight shift of the peak observed at A_{1g} mode to higher wavenumbers occurred after doping.

3.6 Photocatalytic activity

Fig. 8 illustrates the photocatalytic degradation of methylene blue in the presence of $Cu_x Co_{3-x}O_4$ (x = 0, 0.02, 0.04, 0.06, 0.1) nanoparticles. Initially, we examined the dye's degradation efficiency under visible light over time in the presence of H_2O_2 without a catalyst, for comparative purposes (Fig. 8b). The MB dve exhibits its highest absorption peak at a wavelength of 663.4 nm. The figure clearly indicates a time-dependent reduction in the intensity of the absorption curve. Based on the experimental findings, the degradation efficiency for MB dye in the presence of undoped Co₃O₄ nanoparticles and H₂O₂ (Fig. 8a) was determined using the eqn (1) and was 79.20% after 330 min. Conversely, in the absence of a catalyst, the dye's degradation was minimal, with an efficiency of only 2.72% after 330 minutes. According to reports, the decomposition of MB dye without H₂O₂ proceeds with difficulty when Co₃O₄ is used as a photocatalyst. The degradation efficiency improves significantly by introducing an oxidant and carefully choosing its concentration. While a higher concentration of the oxidizing agent enhances the degradation rate by increasing hydroxyl radicals (OH'), there are instances where this process may lead to the counterproductive scavenging of valuable OH' radicals, resulting in the formation of weaker hyperoxyl radicals (HO₂).43 The pH of the dye solution plays a critical role in the degradation. Researchers found that using Co₃O₄ nanoparticles as a catalyst, an alkaline medium lead to the decomposition of H_2O_2 into water and oxygen, while in an acidic medium, the number of OH' radicals decreases with the formation of positively charged H⁺ ions, resulting in decreased degradation

Fig. 8 Time-dependent photocatalytic degradation of MB dye (a) in the presence of undoped Co_3O_4 nanoparticles and H_2O_2 ; (b) in the presence of H_2O_2 without catalyst under visible light irradiation; in the presence of (c) 2%, (d) 4%, (e) 6%, (f) 10% Cu-doped Co_3O_4 nanoparticles and H_2O_2 under visible light irradiation.

efficiency.⁴³ Thus, for our experiments, a neutral pH environment was chosen. The mechanism of the photocatalytic degradation process is as follows: when the light energy is equal to or exceeds the band gap energy of the catalyst, electrons are excited from the valence zone to the conduction zone. This generates positively charged holes in the valence zone and negatively charged electrons in the conduction zone. The positively charged holes produce OH' radicals through water oxidation or by being captured by OH⁻. Electrons in the conduction zone react with oxygen adsorbed on the catalyst's surface and form superoxide anion radicals ($'O_2^-$). Consequently, these radicals decompose the dye solution.^{28,43} The

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generation of hydroxyl radicals occurs according to the Photo-Fenton reaction mechanism. Based on this complex reaction mechanism, Co^{2^+} ions react with $\mathrm{H_2O_2}$ to form OH' radicals, and Co^{2^+} ions are regenerated as a consequence of the reduction of Co^{3^+} ions together with the ligand present in the solution. The detailed reaction mechanism is as follows:⁴³

$$Co_{3}O_{4} + hv \rightarrow h_{vb}^{+} + e_{cb}^{-}$$

$$H_{2}O + h_{vb}^{+} \rightarrow OH^{+} + H^{+}$$

$$O_{2} + e_{cb}^{-} \rightarrow O_{2}^{-}$$

$$O_{2}^{-} + H_{2}O \rightarrow H_{2}O_{2} + e_{cb}^{-} \rightarrow OH^{-}$$

$$P_{(ads)}^{2+} + H_{2}O_{2} \rightarrow Co^{3+}_{(ads)} + OH^{-} + OH^{-}$$

$$Co^{3+} - L + hv \rightarrow Co^{2+} + L^{-}$$

Methylene blue dye + $h_{vb}^{+} \rightarrow$ oxidation product

Methylene blue dye + OH' \rightarrow degradation product

The degradation efficiency of MB dye in the presence of Cu_{0.02}Co_{2.98}O₄ nanoparticles improved. In the presence of $Cu_{0.02}Co_{2.98}O_4$, $Cu_{0.04}Co_{2.96}O_4$, $Cu_{0.06}Co_{2.94}O_4$, $Cu_{0.1}Co_{2.9}O_4$ nanoparticles, the degradation efficiency of MB dye increased from 46.77% to 87.51%, 38.76% to 74.66%, 44.95% to 85% and 31.47% to 65.60% for 90-330 min irradiation time, respectively. As can be seen from the results, the highest degradation was observed in the presence of Cu_{0.02}Co_{2.98}O₄ nanoparticles and the weak degradation was observed in the presence of nanoparticles. When $Cu_{0.04}Co_{2.96}O_4$ $Cu_{0.1}Co_{2.9}O_4$ and Cu_{0.1}Co_{2.9}O₄ nanoparticles are used, the degradation efficiency decreases in comparison to undoped Co₃O₄ nanoparticles. Cu_{0.02}Co_{2.98}O₄ nanoparticles have a wide band gap and as a result of their use as a catalyst, an increase in the degradation efficiency occurs with decreasing electron-hole pair recombination and the generation of free electrons and holes for the formation of radicals. However, when the concentration of copper is high, the photocatalytic activity diminishes. This can be ascribed to excessive oxygen vacancies and the high amount of the dopant, which acts as recombination centers for photoinduced electrons and holes.34 The experiments were also performed for the degradation of MB in the presence of

Fig. 9 Time-dependent photocatalytic degradation of MB dye in the presence of $Cu_{0.02}Co_{2.98}O_4$ nanoparticles: (a) amount of photocatalyst is 20 mg, (b) amount of photocatalyst is 30 mg, (c) the concentration of the dye is 5 ppm and (d) the concentration of the dye is 20 ppm.

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 $Cu_{0.02}Co_{2.98}O_4$ nanoparticles by varying the initial dye concentration (5 ppm and 20 ppm) and amount of photocatalyst (20 mg and 30 mg). When using 20 mg and 30 mg $Cu_{0.02}Co_{2.98}O_4$ nanoparticles, the degradation efficiency of MB dye was found to be 90.94% and 93.19% in 330 min, respectively. For these experiments, the concentration of the dye was taken as 10 ppm. When using 5 ppm and 20 ppm dye concentrations, the value of the degradation efficiency of MB dye was 90.48% and 86.44% in 330 min, respectively. For these experiments, the amount of photocatalyst was taken as 10 mg (Fig. 9).

4 Conclusion

In the present study, $Cu_x Co_{3-x}O_4$ (x = 0, 0.02, 0.04, 0.06, 0.1) nanoparticles were synthesized by sonochemical method and subsequent calcination at high temperature. Based on the XRD analysis, the size of Co₃O₄ nanoparticles exhibits irregular changes as the concentration of Cu increases. The band gap value of undoped Co3O4 nanoparticles initially rises from 1.58 eV and 3.33 eV to 1.70 eV and 3.43 eV with a 2% Cu doping content, but subsequently decreases with higher Cu content. The increase in the band gap of $Cu_{0.02}Co_{2.98}O_4$ nanoparticles can be attributed to the Moss-Burstein effect and the decrease of band gap at higher dopant concentrations can be explained with crystal defect. According to TEM measurements, Cu_{0.1}Co_{2.9}O₄ nanoparticles exhibit spherical-like, hexagonallike, rhombohedral-like and irregularly structured morphology and the size of nanoparticles is in the range of 15.78-78.31 nm. EDX results reveal that the percentage ratio of Cu/Co atoms is near to the theoretical ratio of Cu/Co atoms in the $Cu_xCo_{3-x}O_4$ compound and the distribution of Co, Cu and O elements coincide. FT-IR and Raman results also confirm the formation of Co₃O₄. The highest degradation efficiency for MB dye (87.51%) was observed in the presence of $Cu_{0.02}Co_{2.98}O_4$ nanoparticles and this value increased with increasing the amount of photocatalyst and with decreasing of dye concentration to 93.19% and 90.48%, respectively.

Conflicts of interest

There are no conflicts to declare.

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