

Facile Synthesis and Optical Properties of Rod-Like Copper Oxide Nanoparticles

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Abstract: Copper oxide (CuO) nanoparticles (NPs) were synthesized by the sol-gel method, followed by calcination at 600°C for 2 h. The XRD pattern indicated that the synthesized CuO NPs had a monoclinic structure, with an average crystallite size of 53 nm. The FT-IR spectra showed that surfactant molecules were adsorbed on the surface of the CuO nanoparticles, along with the presence of Cu-O bonding. The TEM analysis revealed rod-like CuO NPs with diameters of about 50 nm and lengths ranging from 150 to 200 nm. The XPS analysis confirmed that copper and oxygen were synthesized as the main components with Cu²⁺ and O²⁻ oxidation states. The optical band gap of CuO was calculated to be 2.0 eV. The maximum PL emission was recorded at 430 nm for the 365 nm excitation wavelength, and the change in PL intensity and peak shift was calculated as a function of excitation wavelength. Reflectance spectra show increasing reflectance with wavelength, while Mie calculations indicate decreasing spectral efficiencies, along with a reduction in the real refractive index and an increase in the imaginary part. A slight size-dependent enhancement in optical response is also observed.

Keywords: CuO, Nanoparticles, Sol-gel, Photoluminescence, Mie analysis.

1. INTRODUCTION

Nanoparticles and nanostructured materials are currently gaining popularity due to their high surface-area-to-volume ratio [1, 2]. Metal oxide nanoparticles (NPs) have remarkable optical, electrical, magnetic, catalytic, and biological capabilities due to their distinct physical and chemical properties that differ greatly from those of the bulk metal [1, 3]. CuO NPs are the most abundant transition metal oxide binary compound, as well as the most chemically and physically stable. It has a monoclinic structure and a narrow bandgap of 1.3–1.7 eV with *p*-type semiconductor material [4]. The multivalent states of CuO NPs also have interesting physicochemical properties. CuO NPs were of great interest due to their potential applications in a wide variety of areas, including electronic and optoelectronic devices, such as lithium-ion batteries, photocatalytic activities, gas sensors, biomedical applications, photovoltaics, and supercapacitor materials [5–10]. CuO NPs, with their extensive redox reactions and coloring effects, can alter optical characteristics and up-conversion [11]. CuO has also been employed as a photocatalyst and host material for various luminous emissions [11]. Recently, researchers explored the optical properties of CuO nanostructures [12–14]. However, a report on the luminescence property is limited. The properties of CuO NPs

highly depend on their synthesis methods [15]. The synthesis methods of CuO NPs can control particle size, morphology, and crystallinity. CuO NPs were prepared using a variety of processes, including sol-gel [16], solution-combustion method [6], ball milling [7], precipitation [14], hydrothermal [17], chemical bath deposition [18], solid-state [19], and green synthesis [20]. Of these methods, the sol-gel method is straightforward, suitable for mass production, achieves relatively low temperatures and ambient conditions, and is an economical solution-based growth technology. This method, which regulates precursor concentrations, reaction time, and temperature, is especially suitable for high-quality crystal formation, a greater surface area, and a wide range of shapes. Despite the extensive literature on the synthesis of CuO nanostructures via solution-based methods, the systematic investigations linking sol-gel synthesis conditions to the photoluminescence (PL) behavior of CuO nanoparticles are not fully understood. This study presents the optimized and controlled sol-gel synthesis of CuO NPs to study their structural, morphological, and optical properties.

2. EXPERIMENTAL PROCEDURES

All the reagents used in this chemical synthesis were supplied without any further purification. For the synthesis of CuO NPs, 0.97 g of cupric

nitrate trihydrate ($\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$) was dissolved in 25 mL of DI water. This solution was kept under vigorous stirring for 3 h at room temperature. During this stirring, 0.456 g of citric acid dissolved in 10 mL of DI water was added dropwise. The resulting solution was kept in the oven at 125°C for 3 h. The dried gel was then calcined at 600°C for 2 h in the muffle furnace and collected for further characterization.

The crystal structure of the samples was determined by powder X-ray diffraction (XRD, Buckler D8 Advance) using $\text{Cu K}\alpha$ ($\lambda = 1.5418 \text{ \AA}$) radiation. The transmittance property of CuO was studied using Perkin Elmer Spectrum 100 Fourier-transformed infrared spectroscopy (FTIR). Morphology and lattice diffraction analyses were carried out using a transmission electron microscope (TEM; JEOL, JEM-2010F) operating at a 200 kV accelerating voltage.

The oxidation states of the synthesized CuO NPs were determined using X-ray photoelectron spectroscopy (XPS) with a PHI 5400 XPS spectrometer using a monochromatic $\text{Al K}\alpha$ X-ray source (1486.6 eV, 15 kV, 300 W) and a hemispherical sector analyzer. The reflectance UV-vis spectrum was obtained using a Shimadzu UV-3600 spectrometer with an integrating sphere. PL spectra of the synthesized CuO NPs were recorded at room temperature on a fluorescence spectrophotometer (F-7000) using a 150 W xenon lamp as an excitation source and a grating to select a suitable wavelength for excitation.

3. RESULTS AND DISCUSSION

3.1. Structural Analysis

The XRD spectrum of the synthesized CuO NPs is illustrated in Fig. 1. The diffraction peaks observed at 2θ are attributed to the (110), (002), (111), (-112), (-202), (020), (202), (-113), (-311), (320), (311), and (222) planes associated with the monoclinic structure of CuO NPs. This result is consistent with prior reports [5, 21, 22]. The observed XRD pattern also matches the standard JCPDS Card no. 45-0937 of the pure CuO phase [23, 24]. The Miller-indexed (002) and (111) reflections were the strongest, suggesting that the NPs produced by the sol-gel method have preferred crystal planes. The sharp and highly intense diffraction peaks in the XRD profile indicate that the CuO NPs are highly crystalline. In addition, no impurities were noticed. Thus, the XRD pattern

strongly supports the formation of CuO nanoparticles.

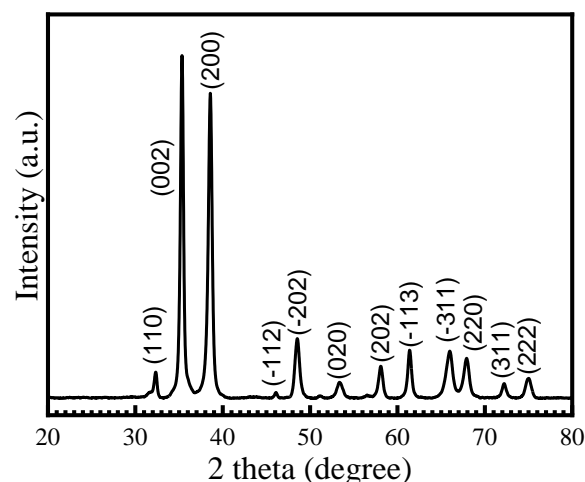


Fig. 1. The XRD spectrum of CuO NPs after calcined at 600°C for 2 h

The average crystallite size (D) was calculated from the XRD pattern using Scherrer's equation [25]:

$$D = \frac{0.9\lambda}{\beta \cos\theta} \quad (1)$$

The dislocation density (δ), which gives the number of defects in a crystal, and the micro-strain (ϵ) were determined using the following relations [14]:

$$\delta = \frac{1}{D^2} \quad (2)$$

$$\epsilon = \frac{\beta}{4 \tan\theta} \quad (3)$$

The lattice constants ($a \neq b \neq c$) and unit cell volume (V) are also calculated from the strong diffraction peaks of the (002), (111), and (-202) planes in the case of the monoclinic crystal [26] using

$$\frac{1}{(d_{hkl})^2} = \frac{1}{\sin^2\gamma} \left(\frac{h^2}{a^2} + \frac{k^2 \sin^2\gamma}{b^2} + \frac{l^2}{c^2} - \frac{2hkl \cos\gamma}{ac} \right) \quad (4)$$

$$V = abc \sin\gamma \quad (5)$$

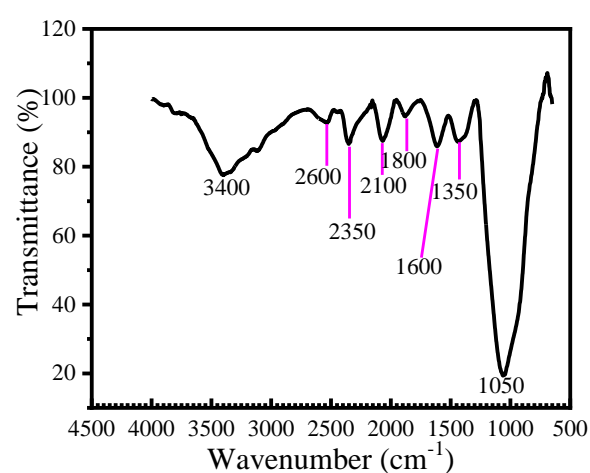
where λ is the wavelength of the X-ray used, which is $\text{CuK}\alpha$ radiation ($\lambda = 1.5406 \text{ \AA}$), β is the full width at half maximum (in radians), and θ is the Bragg's angle, where a , b , and c are the lattice constants, d_{hkl} is the interplanar distance, γ is the interfacial angle ($\gamma = 99.41^\circ$), and h , k , and l are the Miller indices. The calculated average crystallite size is found to be 53 nm, as shown in Table 1. Table 1 also displays the results for peak position and estimated structural parameters. The obtained dislocation density is $3.6 \times 10^{14} \text{ m}^{-2}$, and the micro-strain is 8.3×10^{-3} . The calculated lattice parameters were $a = 4.6336 \text{ \AA}$, $b = 3.4296 \text{ \AA}$, and $c = 5.1251 \text{ \AA}$, and the unit cell volume was 80.33 \AA^3 which is in good agreement with previous reports [6, 14, 17].

Table 1. XRD peak position and calculated structural parameters of CuO NPs

Miller indices (hkl)	2θ	FWHM	d-spacing (Å)	Lattice constants (Å)			Unit cell Volume (Å ³)	Average crystallite size (nm)	Average micro-strain	Dislocation density (m ⁻²)	Urbach energy (eV)
				a	b	c					
(002)	35.3	0.1574	2.5275	4.6336	3.4296	5.1251	80.33	53	8.3×10 ³	3.6×10 ¹⁴	0.84
(111)	38.57	0.1574	2.3168								
(-202)	48.5	0.2558	1.8718								

3.2. FT-IR Spectroscopy

FT-IR spectroscopy is an important instrument for identifying the presence of various functional groups in the synthesized powder. Figure 2 shows the FTIR spectrum of CuO NPs. The band at 3400 cm⁻¹ corresponds to the O–H hydroxyl group stretching vibration [27]. This O–H vibration arises from hydrogen bonding between CuO and adsorbed water molecules, as well as interactions with their surfaces, indicating that the surface of the NPs absorbs water from the environment. Peaks in the range of 2100–2600 cm⁻¹ may be associated with C≡C stretching vibrations or the adsorption of CO₂ on the surface of CuO NPs [28]. The vibration band at 1800 cm⁻¹ is attributed to the C=O stretching mode [27]. The band at 1600 cm⁻¹ can be attributed to the Cu–O stretching vibration, confirming the formation of CuO NPs [29]. The weak peak at 1350 cm⁻¹ usually appears in the spectrum and denotes the C–H stretching mode. The peak at 1050 cm⁻¹ is due to O–H bending and C–O and O–Cu pair stretching [30].

**Fig. 2.** FTIR spectrum of CuO NPs

3.3. Microstructure

The particle size and morphology of the sample

are analyzed by transmission electron microscopy (TEM). Figure 3(a) displays a picture of the calcined powder. The black color of the CuO product is clearly visible. Figures 3(b) and 3(c) show the TEM images of the prepared CuO NPs at different magnifications, allowing for a more detailed investigation of their structural features. These images showed how NPs get agglomerated and have irregular forms [31, 32]. Two distinct particle shapes: spherical and rod-shaped, were shown by TEM images. The rod-shaped particles had a diameter of roughly 50 nm and a length of 150–200 nm. The majority of nanoparticles detected in the micrographs are roughly spherical, with a small number of elongated particles. The particle size of a spherical morphology was in the range of 50–60 nm, which is in good agreement with the calculated crystallite size in the Scherrer formula. Figure 3(d) depicts the selected area electron diffraction (SAED) pattern of CuO NPs. The diffraction rings are indexed as (002), (111), and (-202) planes, confirming the formation of the polycrystalline monoclinic crystal of the synthesized CuO NPs [14, 33]. The TEM study also supports the XRD result.

3.4. XPS Analysis

The chemical oxidation state of the CuO NPs was investigated by X-ray photoelectron spectroscopy (XPS), and the results are presented in Fig. 4. The peaks of the XPS survey spectrum in Fig. 4(a) indicated the presence of Cu, O, and C (used as references) elements in the synthesized CuO NPs [5]. Other contaminants did not show up as clearly on the survey scan. As seen in Fig. 4(b), the XPS spectrum for the Cu 2p core level displays two peaks at 934 eV and 953.9 eV, which correspond to Cu 2p_{3/2} and Cu 2p_{1/2}, respectively. The distance between Cu 2p_{3/2} and Cu 2p_{1/2} was measured to be 19.9 eV, indicating the presence of Cu²⁺ ions in the synthesized NPs [34].

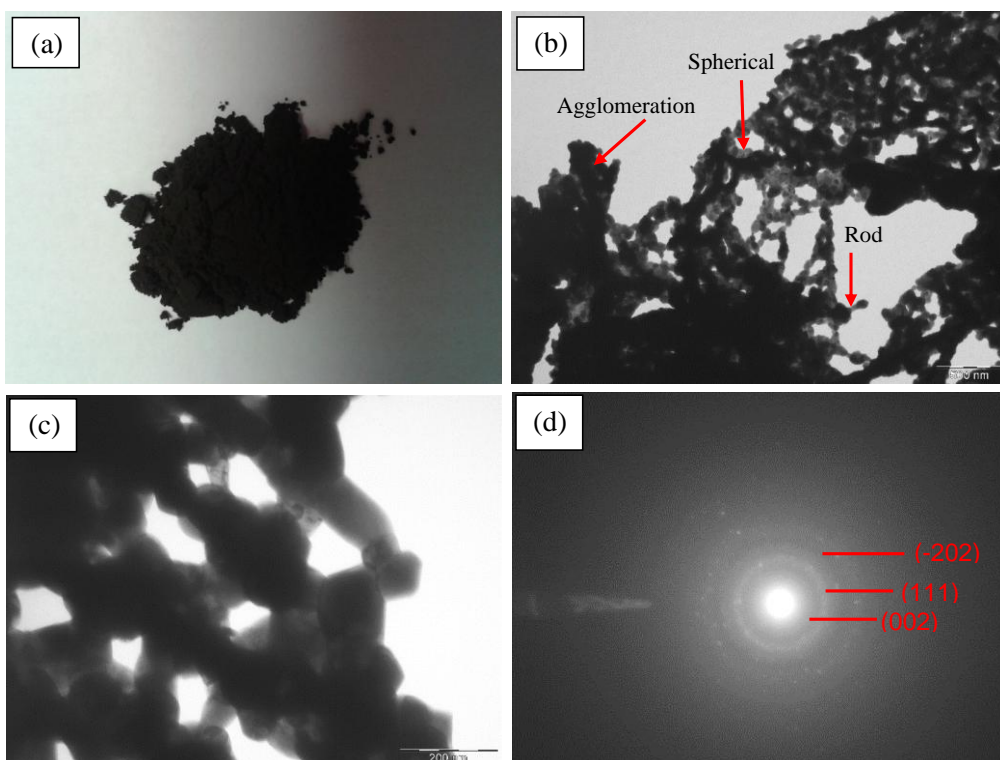


Fig. 3. a) Photograph of calcined powder, b,c) TEM images of the synthesized NPs at different magnifications, d) SAED pattern CuO NPs

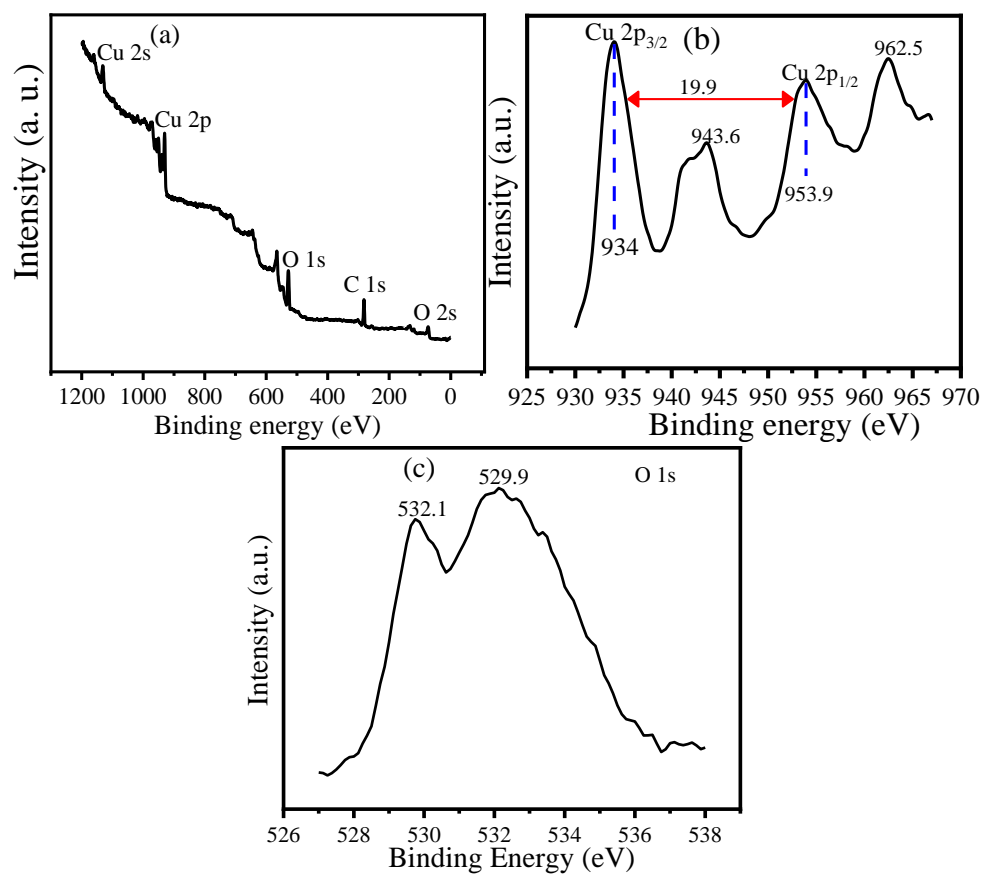


Fig. 4. a) XPS survey spectrum, b) Cu 2p, c) O1s core-levels of CuO NPs

The presence of the Cu^{2+} chemical state was confirmed by the detection of two additional satellite peaks at 943.6 and 962.5 eV in addition to the Cu 2p peaks. Figure 4(c) shows the O1s spectrum of CuO NPs. The O1s spectrum is now deconvoluted into distinct components, where the lower binding energy peak at ~ 529.9 eV corresponds to lattice oxygen (O^{2-}) in CuO, while the higher binding energy component at ~ 532.1 eV is assigned to surface hydroxyl groups and/or adsorbed water molecules [35, 36]. This result is in agreement with the previous reports [37, 38].

3.5. Optical Properties

Figure 5(a) depicts the diffuse reflectance spectrum of CuO NPs recorded at room temperature in the range of 200–800 nm. The sample is extremely reflective in the visual region, as indicated in Figure 5(a) [12]. A significant absorption edge is found at around 354 nm, which is consistent with previous reports for CuO NPs [39, 40]. The reflectance data were converted to the absorption coefficient $F(R)$ values using the Kubelka-Munk equation [41]:

$$\alpha = F(R) = \frac{(1-R)^2}{2R} \quad (6)$$

where $F(R)$ is the Kubelka–Munk function, α is the absorption coefficient, and R is the reflectance (%). The band gap energy (E_g) of the CuO NPs

was evaluated using Tauc's plot in accordance with the equation [42]:

$$\alpha h\nu = A(h\nu - E_g)^n \quad (7)$$

where $h\nu$ represents the photon energy, A is a proportionality constant, E_g is the bandgap energy, and n is the exponential constant index, which depends on the nature of the transition ($n=2$ for an indirect allowed transition). Figure 5(b) depicts the Tauc plot of $(\alpha h\nu)^{1/2}$ vs. $h\nu$, which is based on the indirect transition. The energy band gap is calculated by extrapolating a line tangent to the $h\nu$ axis. The estimated optical energy band gap was approximately 2.0 eV, indicating its potential in photocatalysis. This value and the reported values agree quite well [14, 43]. However, the obtained E_g value for synthesized CuO NPs is substantially higher than that of bulk CuO (1.85 eV) [44, 45] but lower than previously reported values [24, 46, 47]. The quantum confinement effect brought on by the nanosized crystals found in TEM examination is responsible for the high E_g value [47].

One useful method for examining the atomic structure of a crystal is optical conductivity. The optical response of a material is mainly studied in terms of the optical conductivity (σ), which is given by the relation [48]:

$$\sigma = \frac{anc}{4\pi} \quad (8)$$

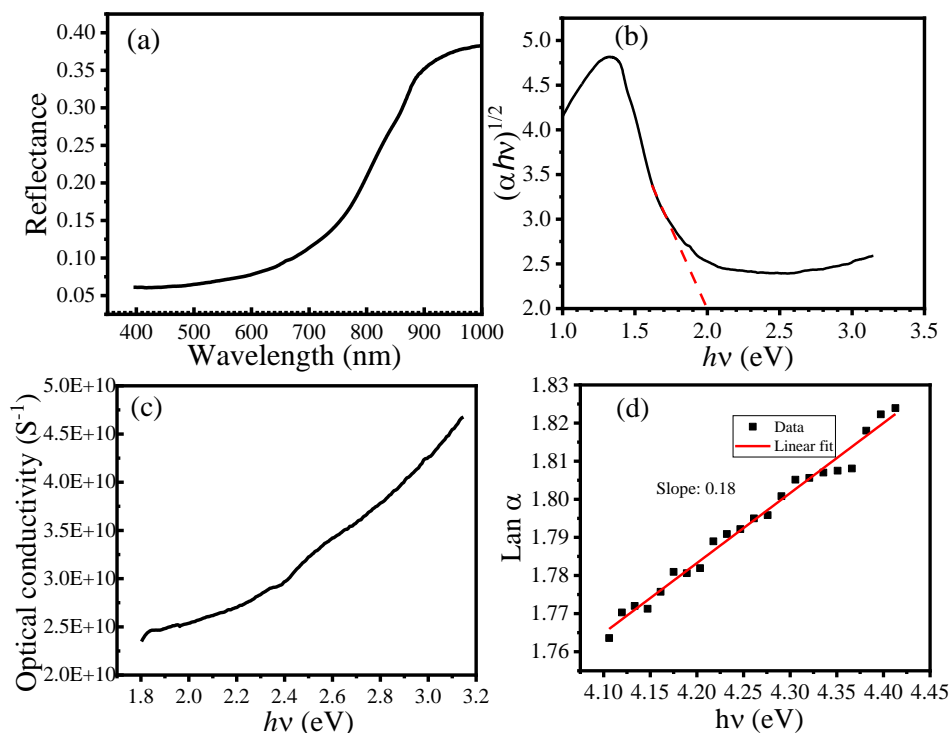


Fig. 5. a) Diffuse reflectance spectrum, b) Tauc plot, c) optical conductivity, d) Urbach energy of the synthesized CuO NPs

where c is the velocity of light, α is the absorption coefficient, and n is the refractive index. The optical conductivity of CuO NPs as a function of photon energy is displayed in Fig. 5(c). Optical conductivity generally increases with photon energy because higher-energy photons can excite electrons across larger energy gaps and into higher conduction states, thereby enhancing the material's ability to absorb and transport optical energy. This trend is often consistent with the decrease in reflectance at higher photon energies, since more photons are absorbed rather than reflected, leading to stronger electronic transitions [49]. The Urbach energy can also be calculated using [50]:

$$\alpha = \alpha_0 e^{\frac{hv}{E_U}} \quad (9)$$

where α is the absorption coefficient, α_0 is a constant, hv is the incident photon energy, and E_U is the Urbach energy, which indicates the width of the band tails of the localized states in the gap region and is used to characterize the degree of disorder in crystalline solids. Fig. 5(d) shows $\ln\alpha$ as a function of photon energy hv . Urbach energy is derived from the inverse of the slope of $\ln\alpha$ vs hv . The slope was calculated to be 0.18. The computed Urbach energy value of CuO NPs was 5.56 eV [48, 51], which is higher than that of other metal oxides. The Urbach energy, which characterizes the width of the exponential absorption tail near the band edge, provides insight into the degree of disorder and defect states in the material. A larger Urbach energy indicates more localized states that facilitate sub-bandgap absorption, while at higher photon energies, the optical conductivity dominates as band-to-band transitions become more pronounced. Thus, the interplay between optical conductivity, reflectance, and Urbach energy highlights how intrinsic electronic structure and disorder collectively govern the material's optical response.

Figure 6 shows the PL emission spectra of CuO NPs recorded at room temperature with excitation wavelengths of 325, 340, and 365 nm. Every spectrum has an emission peak that is easily seen. Violet emission peaks are visible in the spectra at 428, 428.6, and 430.4 nm at excitation wavelengths of 325, 340, and 365 nm, respectively. The plot indicates that as the excitation wavelength increases, the emission's intensity rises linearly. The excitation source at 365 nm produced the greatest emission peak. Emission intensity rises with increasing excitation wavelength because the unrelaxed

photoexcited species separate from the lower-energy excited species more clearly [52]. The increase in emission intensity at higher excitation wavelengths was consistent with the findings of I. Abdullahi et al. [11]. A high recombination rate is indicated by the strong band. Conversely, a decrease in peak heights suggests that band levels are easily accessible for electron migration and that electron-hole couples are well separated [33]. The intensity shift was attributed to radiative recombination caused by surface defect passivation [53]. The recombination process involves moving the electron back to the valence band while simultaneously emitting energy. The luminescence bands in the violet region (428–430 nm) may be attributed to the band edge emission or defect levels in CuO caused by oxygen vacancy and Cu interstitial [12, 54]. Additionally, the violet emission peak is a result of the charge transfer transition from Cu and O ions in the CuO lattices [55]. The emission peaks are also slightly red-shifted, which is directly linked to the average grain size. Because 365 nm light has less energy than 320 nm and 340 nm light, the spectra show a significant red shift when compared to those excited at lower wavelengths. In general, visible emissions are strongly dependent on the synthesis conditions, nanoparticle size, and shape. The PL value achieved here is consistent with previous reports [54, 56].

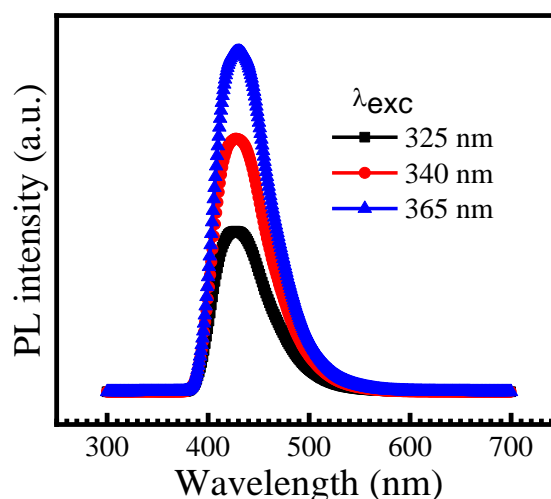


Fig. 6. PL emission spectra of CuO NPs excited by different wavelength using a 150 W xenon lamp as an excitation source

3.6. Mie Analysis

The optical properties of spherical nanoparticles can also be studied using the Mie theory [58]. In

Mie scattering theory, optical scattering efficiency (Q_{sca}) and extinction efficiency (Q_{ext}) [58] can be calculated using

$$Q_{sca} = \frac{2}{x^2} \sum_{n=1}^{\infty} (2n+1) (|a_n|^2 + |b_n|^2)$$

$$Q_{ext} = \frac{2}{x^2} \sum_{n=1}^{\infty} (2n+1) \text{Re}(a_n + b_n) \quad (10)$$

where $x = \frac{2\pi r}{\lambda}$ is the size parameter, r is the radius of the nanoparticle, and a_n and b_n are the Mie scattering coefficients, which are calculated numerically. For a simpler analysis, the directional scattering behavior can be described by the

average cosine of the scattering angle, known as the asymmetry factor (g) [59], given by

$$g = \langle \cos\theta \rangle = \frac{\int_0^\pi 2\pi S_{11}(\theta) \cos\theta \sin\theta d\theta}{\int_0^\pi 2\pi S_{11}(\theta) \sin\theta d\theta} \quad (11)$$

where $S_{11}(\theta)$ is the scattering matrix and θ is the scattering angle. The efficiency factors Q_{sca} , Q_{ext} , Q_{abs} , the real and imaginary (n, k) index of refractions, and the asymmetry (g) were determined using the Mie scattering theory with the help of Mie Plot v.4.3 software. The changes in optical characteristics for particle sizes of 50, 60, and 70 nm are shown in Fig. 7.

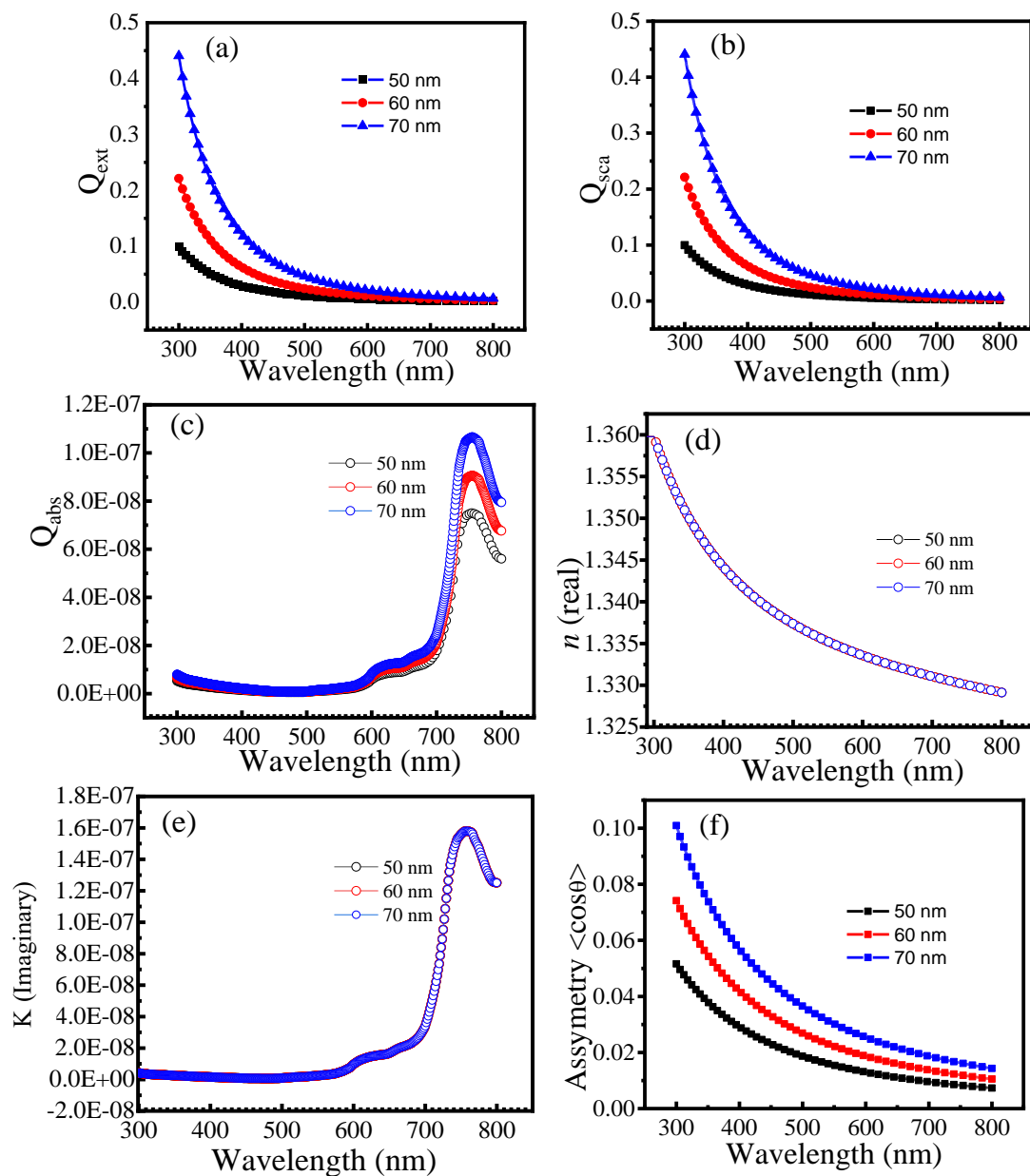


Fig. 7. Mie calculations of the a) extinction efficiency, b) scattering efficiency, c) absorption efficiency, d) real refractive index, e) imaginary refractive index and f) asymmetry of CuO NPs for different particle sizes (50–70 nm)

The optical characteristics of CuO nanoparticles exhibit a clear dependence on both wavelength and particle size. With increasing wavelength, the scattering efficiency and extinction efficiency display a decreasing trend with wavelength, reflecting the reduced contribution of scattering and attenuation in this spectral region (Figs. 7(a) and 7(b)). As wavelength increases (lower photon energy), fewer electronic transitions are excited, so the polarization response weakens, and the real index decreases (Fig. 7(d)). When considering particle size within the 50–70 nm range, both extinction efficiency and scattering efficiency increase with increasing nanoparticle diameter.

This size-dependent enhancement arises from the larger photon–matter interaction and stronger polarization effects in bigger particles, which increases the magnitude of these optical parameters. In contrast, the absorption efficiency (Fig. 7(c)), reflectance, and the imaginary part of the refractive index (Fig. 7(e)), all show a pronounced enhancement with wavelength, indicating stronger photon–matter interactions and more effective electronic transitions. Figure 7(f) displays the asymmetry parameter (g) of CuO NPs at 50–70 nm as a function of wavelength. The forward scattering effect was enhanced with a positive value of g , and the g value also rose as the NP size grew [60]. Furthermore, for CuO NPs, large values of g indicate light scattering in a narrow wavelength range [61].

4. CONCLUSIONS

CuO NPs were successfully prepared using the sol-gel method, which involves calcination at 600°C for 2 h. The XRD pattern confirmed the monoclinic CuO phase and a crystallite size of about 53 nm. The FTIR spectrum verified the chemical bonding of CuO NPs. The TEM investigation confirmed the spherical shape and the rod-like morphology. The spherical particle size was approximately 50–60 nm, while the rod-shaped particles ranged in length from 150 to 200 nm with an average diameter of about 50 nm. The distinct diffraction rings observed in the SAED pattern confirm the polycrystalline structure of the CuO NPs. The binding energies obtained from XPS analysis are associated with the Cu²⁺ and O²⁻ oxidation states. CuO has a direct optical bandgap of 2.0 eV, as shown by the Tauc plot, which is blue-shifted from the bulk value of

1.8 eV. The PL spectra as a function of excitation wavelength revealed that raising the excitation wavelength from 320 nm to 365 nm resulted in enhanced violet emission intensity centered at 428–430 nm and a red shift in peak emission wavelength. The Mie theory–based analysis of the optical properties of CuO NPs in the 50–70 nm size range was employed as a function of wavelength. The calculated results show that spectral efficiencies decrease with increasing wavelength, while the real part of the refractive index decreases and the imaginary part increases, indicating reduced absorption at longer wavelengths. A size-dependent enhancement in optical response is also observed. This may be due to the larger light-matter interaction at a larger size. The combined experimental and theoretical results revealed that CuO NPs exhibit tunable optical behavior.

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