



Study of structural and optical properties of cupric oxide nanoparticles

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Abstract In this study, cupric oxide (CuO) nanoparticles were synthesized via sonochemical method. The samples were characterized by X-ray diffraction, Fourier transform infrared spectroscopy, scanning electron microscope, and transmission electron microscopy. The spherical CuO nanoparticles were dispersed in sodium hexametaphosphate under sonication (25 kHz) to analyze the particle size distribution and UV absorption spectra. Using these absorption spectra, we further examined the CuO nanoparticle to explore the possibility of using them as a material for applications such as solar cell and textile production.

Keywords Cupric oxide · Sonochemical method

Introduction

Nanocrystalline semiconductor particles have drawn considerable interest in recent years due to their interactive properties such as large surface-to-volume ratio and distinctive electronic and optical properties as compared to bulk materials (Huang et al. 2010; Son et al. 2009; Xu

et al. 2007; Lim et al. 2012; Qi et al. 2007; Kidowaki et al. 2012). Cupric oxide (CuO) is a transition metal oxide. It has a monoclinic structure and a narrow bandgap of 1.2 eV (indirect) with p-type semiconductor material. It is also an attractive candidate for light-harvesting applications due to its band gap energy of 1.4 eV (Son et al. 2009; Xu et al. 2007; Lim et al. 2012). CuO compounds are technologically well-known materials that have their applications in areas (Qi et al. 2007) such as solar energy materials, electronic materials, gas sensor, magnetic media, optical devices, batteries and catalyst (Kidowaki et al. 2012; Lang et al. 2006; Teng et al. 2008; Stewart et al. 2004; Wang et al. 2007; Morales et al. 2005), and constructing junction devices such as p-n diode (Muhibullah et al. 2003), as well as for photoconductive, photothermal, and photoelectrochemical applications (Chiang et al. 2012).

Attempts are made to prepare CuO nanoparticles using different methods such as spray pyrolysis (Chiang et al. 2012), electrochemical techniques (Chen et al. 2009), hydrothermal treatments (McAuleya et al. 2008), sonochemical method (Gandhi et al. 2010), and wet chemical methods (Mahapatra et al. 2008) with different morphologies. Among these methods, sonochemical preparation method is used to break the chemical bond of the solution compound. In general, formation, development, and implosive collapse of microcavities are the three steps involved in the production of nanoparticles through sonochemical method (Gandhi et al. 2010). Most inorganic materials such as Al₂O₃, ZnO, MnO₂, SnO₂, Y₂O₃, and clay particles may be more dispersed in sodium hexametaphosphate. Sodium polyphosphate (Calgon) is also a suitable dispersing agent (Greenwood 2003). These are the solvents used to readily dissolve in water and they are normally added at approximately 1 wt %.

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The optical behavior of CuO nanomaterials is mainly assessed through UV–vis and photoluminescence techniques to explore the electronic transitions in semiconductors including band-edge or near-band-edge transitions. Only a few works based on UV–vis analysis are available on the absorption and band gap studies of CuO nanoparticles (Cho 2013; Lin et al. 2004; Kaur et al. 2006; Tauc et al. 1966). However, optical properties of CuO nanoparticles are not revealed extensively.

In this article, using a simple and low-cost approach, CuO nanoparticles were synthesized using sonochemical method and their optical properties were investigated using different characterization techniques.

Experimental

Materials

Analytic grade (AR) chemicals namely copper (II) nitrate [Cu(NO₃)₂], sodium hydroxide (NaOH), and sodium hexametaphosphate [Na(PO₃)₆] obtained from Merck, and deionized water, obtained in our laboratory, were used without any further purification.

Preparation of copper oxide nanoparticles

In a typical procedure, 50 ml NaOH solution was dropwise added into 100 ml aqueous Cu(NO₃)₂·2H₂O solution under sonication process. Meantime, the mixed solution was kept under sonication at a constant interval of 60 s and frequency of 25 kHz for 60 min and pH was maintained at 12. The Cu(OH)₂ solution thus formed was blackish brown. The precipitate was centrifuged at room temperature at 10,500 rpm and then washed with deionized water several times. Finally, the obtained solution was washed with ethanol to remove ionic impurities and then annealed at 673 K under vacuum for 4 h. Finally, black CuO nanoparticles were obtained.

Characterization

The phase identification was carried out using powder X-ray diffraction (XRD; X'Pert PRO PW-1830; Philips, Germany). The functional group present in the sample was recorded using Fourier transform infrared spectroscopy (FTIR; Spectrum 100; PerkinElmer). The microscopical analysis of the samples was carried out by a scanning electron microscope (SEM; JSM-200; JEOL, USA) equipped with energy-dispersive spectroscopy (EDS), and the powder sample was mixed with ethanol to analyze the sample through transmission electron microscopy (TEM;

JEM-2100; JEOL) along with selected area electron diffraction (SAED) pattern. The CuO nanoparticles were dispersed in (NaPO₃)₆ solution at a ratio of 1:100 and then kept under ultrasonic irradiation (25 kHz) to obtain a homogeneous mixture solution, which was used to analyze particle size distribution by dynamic light scattering technique using a particle size analyzer (PSA; Nanophox, Germany) and optical properties were determined by recording the absorption spectra using a UV spectrometer (Lamda 20; PerkinElmer).

The optical band gap (E_g) was obtained from the absorption coefficient (α), which was calculated from the transmittance spectra of CuO nanoparticles. The value of α is given as follows (Khalil et al. 2011; Sumangala et al.

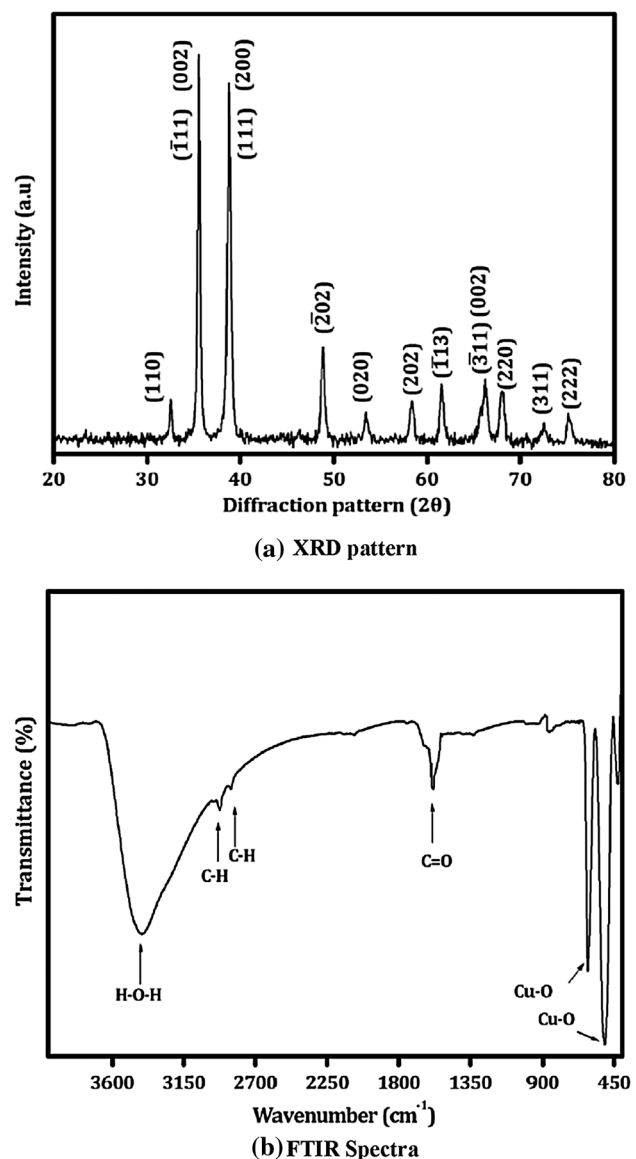


Fig. 1 Characterisation of CuO nanoparticles a XRD Pattern and b FTIR Spectra

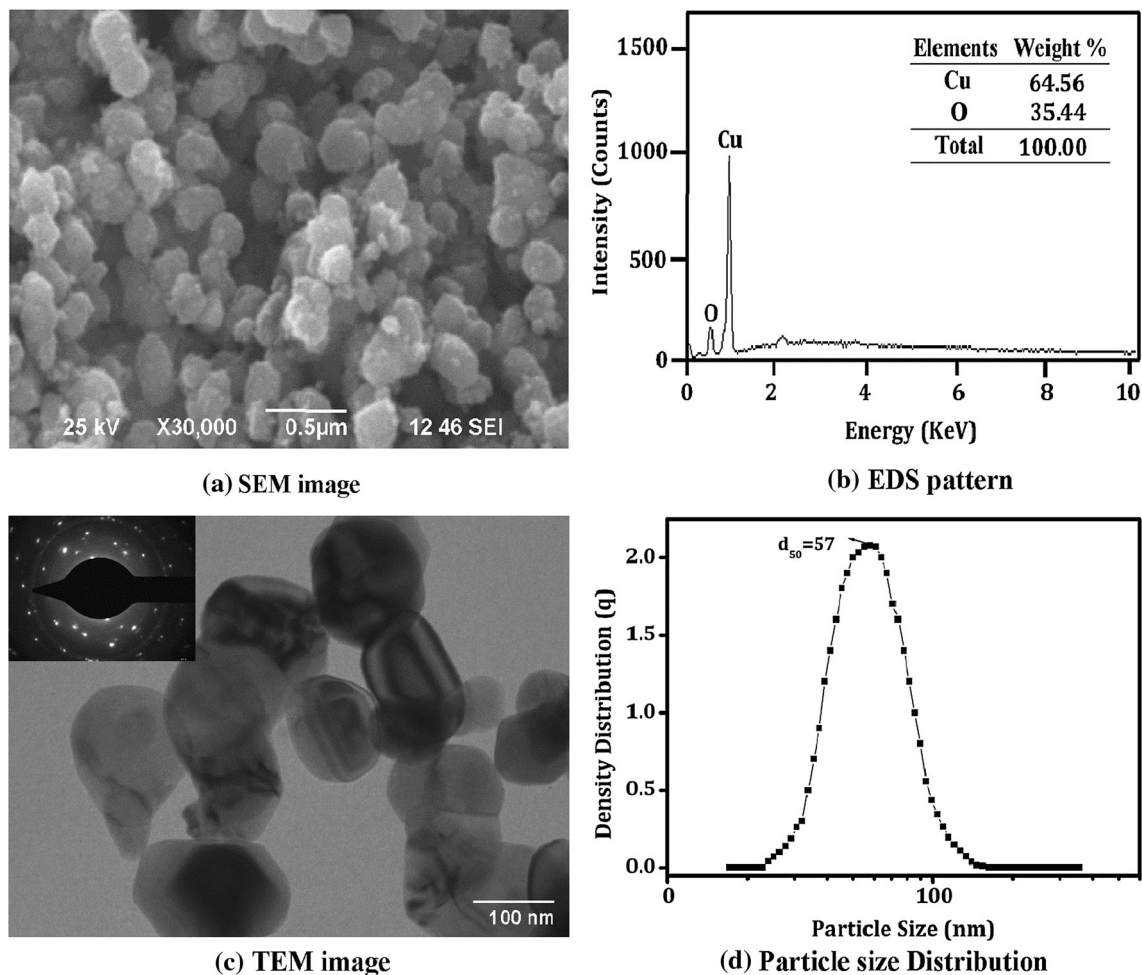


Fig. 2 Characterisation of CuO nanoparticles **a** SEM image, **b** EDS pattern, **c** TEM image and **d** particle size distribution

2005; Mishra and Mishra 2009; Hamad et al. 2014; Growth 2011):

$$\alpha = \frac{1}{d} \ln(1/T)$$

where d is the film thickness, T the transmittance, and α the absorption edge. CuO is a wide band gap semiconductor material with direct band gap. The optical band gap (E_g) is given by

$$(\alpha h\nu)^2 = A(h\nu - E_g)$$

where α is the absorption edge and $h\nu$ the photon energy. The refractive index (n) and extinction coefficient (k) of CuO nanoparticles were calculated from the following expressions (Sumangala et al. 2005):

$$n = \left(\frac{1+R}{1-R} \right) + \sqrt{\left(\frac{4R}{(1-R)^2} - k^2 \right)}$$

$$k = \frac{\alpha \lambda}{4\pi}$$

where R is reflectance, λ the wavelength, and k the extinction coefficient.

The Urbach tail of the particles can be determined from the following relation (Growth 2011):

$$\alpha = \alpha_0 \exp\left(\frac{h\nu}{E_U}\right)$$

where α is absorption edge, α_0 a constant, $h\nu$ the photon energy, and E_U the Urbach energy.

Results and discussion

Analysis of structural and functional properties

The crystal phase of CuO nanoparticles is consistent with JCPDS card no 05-0661 of all diffraction peaks, as shown in Fig. 1a. The sharp observed planes (110), (−111), (111), (202), (020), (220), (−113), and (022) show the monoclinic phase of CuO nanoparticles. Further, it is evident that there

are no peaks corresponding to Cu_2O or $\text{Cu}(\text{OH})_2$ phases. The absence of impurities is also reflected from the XRD pattern. The average crystallite size of the CuO nanoparticles obtained from full-width at half maximum of the diffraction peak is 21 nm.

Figure 1b shows the functional property of FTIR spectra. The obtained spectra show an absorbed band of approximately 640 cm^{-1} , which shows the characteristic band of monoclinic phase of pure CuO similar to that obtained in an earlier report (Gandhi et al. 2010). The bands obtained at 1585 and 1644 cm^{-1} show the carbonyl $\text{C}=\text{O}$ stretching bonds. Similarly, the $\text{C}-\text{H}$ stretching bonds occur in the region of $3300\text{--}2800\text{ cm}^{-1}$ (Xu et al. 2007). Moreover, the broad band centered at 3413 and 3392 cm^{-1} is attributed to the stretching and bending vibrations of absorbed water and surface hydroxyls (Gandhi et al. 2010).

Figure 2 shows the SEM and TEM images of the sample. Figure 2a shows the SEM image of CuO nanoparticles, which consists entirely of flake-like morphology. The fine nanoparticles aggregate due to their high surface energy. EDS analysis of the flake-like CuO architecture suggests that the sample contains only the identified elements of Cu

and O with the atomic and weight percentages as shown in the inset of Fig. 2b. It also confirms that the grown nanoparticles are composed of CuO without any impurities. TEM image suggests that CuO nanoparticles show relatively uniform diameter in the range of $30\text{--}60\text{ nm}$ (Fig. 2c). SAED pattern shows that the CuO nanoparticles comprise small nanocrystalline structures with different orientation of the single crystal diffraction pattern. Therefore, the CuO nanoparticles are crystalline in nature, which closely matches with the XRD pattern. The distribution of CuO nanoparticles is in the range between 35 and 125 nm . However, the average distribution (d_{50}) of the nanoparticles is 57 nm (Fig. 2d).

Analysis of optical properties

The absorption edge is found so that the type of transition and bandgap value can be determined (Essic and Mather 1993). The absorption spectra were used to study the energy band and the type of electronic transitions. Absorption spectra of CuO nanoparticles are shown in Fig. 3a, which show a strong fundamental absorption edge

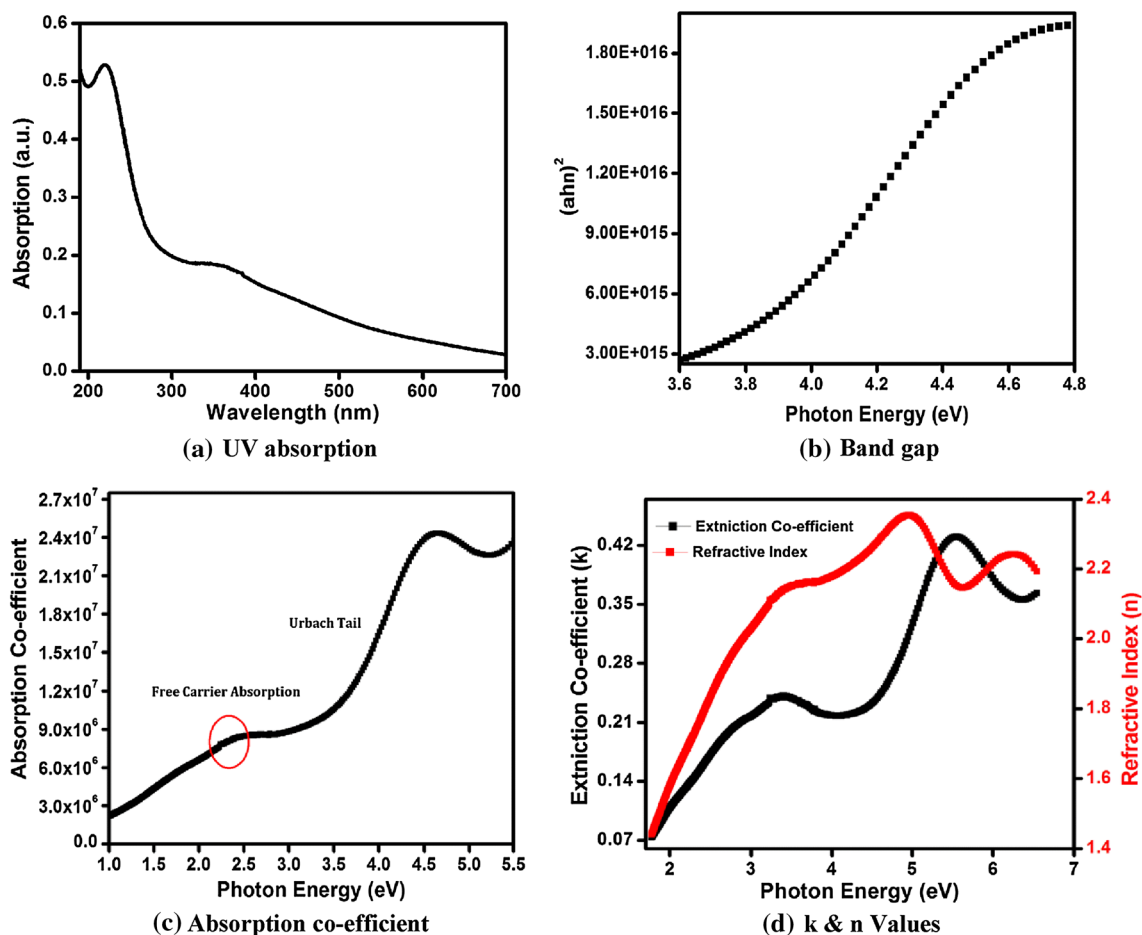


Fig. 3 Optical properties of CuO nanoparticles **a** UV absorption, **b** band gap, **c** absorption coefficient and **d** k & n values

approximately 219 nm due to direct transition of electrons. Optical absorption shows that the direct band gap compared to indirect band gap permits us to determine the crystallinity of a material. The functional relationship between $\alpha h\nu$ and photon energy for CuO nanoparticles is presented in Fig. 3b. The E_g value can be obtained by extrapolating the linear portion to the photon energy axis. If the direct band gap is higher than the indirect band gap, the materials will be crystalline (Radhakrishnan and Beena 2014). The calculated direct band gap value was 3.85 eV, which was higher than the bulk band gap value (3.5 eV). Here, only direct transition-related absorption was observed, and there was no indirect transition absorption peak (Fig. 3b). The observed increasing band gap could be ascribed to the presence of intragap states and quantum confinement effect.

The variation of absorption coefficient of CuO nanoparticles as a function of photon energy is shown in Fig. 3c. The figure clearly shows that the absorption coefficient tends to decrease exponentially as the wavelength increases. This behavior is typical for many semiconductors and could be due to a variety of reasons, such as internal electric fields within the crystal, deformation of lattice due to strain caused by imperfection, and inelastic scattering of charge carriers by phonons (Moss et al. 1973; Honsi et al. 2006; Almqvist 1996). The highest absorption coefficient is observed in the UV region and is in the order of 10^6 – 10^7 from UV to visible region. Elongation of band gap in the form of band tail and absorption due to free carriers are observed from the absorption coefficient.

The k and n values against photon energy plot of CuO nanoparticles are shown in Fig. 3d. From the image, the refractive index of CuO nanoparticles is found to increase with an increase in photon energy, whereas it is found to have lesser value than that for the bulk CuO ($n = 2.6$). For instance at band gap region, the refractive index values range from 2.1 to 2.37. In general, refractive index of the semiconductor is a measure of its transparency to incident spectral radiation. The assessment of the refractive index of the optical material is notably important for applications in integrated optic devices (Xue et al. 2008). The extinction coefficient of CuO nanoparticles increases with increase in the photon energy. The observed extinction coefficient values are very low (0.42) in the absorption region, and it specifies the smoothness of the surface and homogeneity of the particles.

Optical conductivity of CuO nanoparticles with respect to the photon energy is shown in Fig. 4a, which describes the free charges (Madhup et al. 2010; Millis et al. 2005; Lupi et al. 2005). Optical conductivity increases with respect to the photon energy in the UV region and it in turn increases the free carriers. The optical conductivity

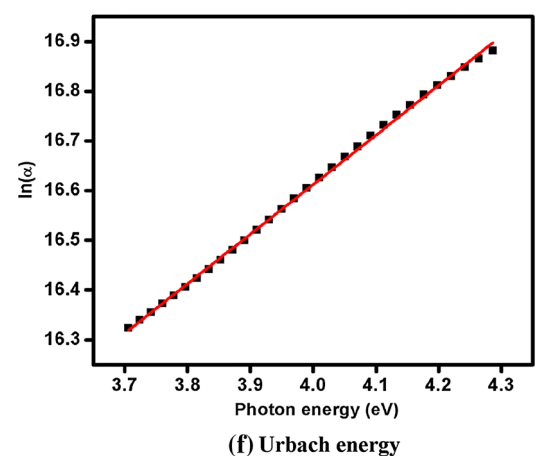
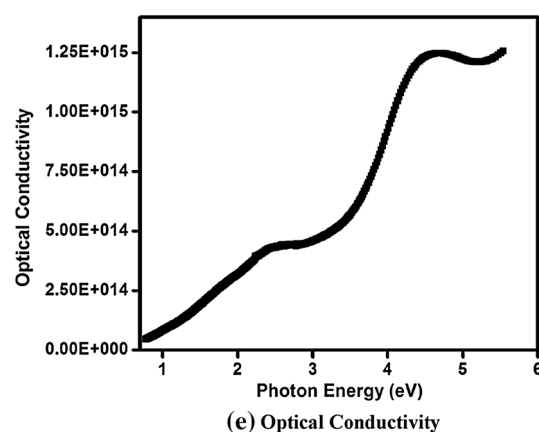


Fig. 4 Optical properties of CuO nanoparticles: *e* optical conductivity and *f* Urbach energy

dramatically increases in the higher energy region (UV absorption), which indicates the availability of free carriers of photon energy and hence, the maximum of optical conductivity is observed. However, an enormous decrease in the optical conductivity in the visible region is observed due to trapping of free carriers.

The plot of $\ln(\alpha)$ versus photon energy is linear, as shown in Fig. 4b. Urbach energy is obtained from the inverse of the slope (see the figure) and the calculated Urbach energy value of CuO nanoparticles is found to be 1.01 eV, which is higher than that of the other metal oxides. Basically, Urbach energy depends on the static and induced disorder, temperature, strong ionic bonds, and average photon energies (Gandhi et al. 2010). Urbach band tail calculation can be performed to confirm the change in band gap energy and the interband formation in CuO nanoparticles. The optical band structure and optical transitions are affected by the width of the localized states available in the band gap, which is called the Urbach tail. Therefore, the CuO nanoparticles can be used in semiconductor devices such as photoamplifier, photovoltaic cell, and photo detectors.

Conclusion

The CuO nanoparticles were successfully prepared via sonochemical method. The crystallite size of CuO was found to be 21 nm. Distribution of spherical CuO nanoparticles was observed from micrographical analysis. The average particle size was clearly visualized and measured 57 nm, respectively, through TEM and PSA studies. Optical properties of CuO nanoparticles were analyzed/clarified through UV absorption. The direct band gap of CuO nanoparticles was found to be large. Moreover, the absorption coefficient, optical conductivity, refractive index along with extinction coefficient and Urbach energy were calculated through UV absorption. The CuO nanoparticles could be used as an extensive semiconductor, optical devices, solar cell applications, and so on.

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References

- Almqvist N (1996) Fractal analysis of scanning probe microscopy images. *Surf Sci* 355:221–228
- Chen LB, Lu N, Xu CM, Yu HC, Wang TH (2009) Electrochemical performance of polycrystalline CuO nanowires as anode material for Li ion batteries. *Electrochim Acta* 54:4198–4201
- Chiang CY, Aroh K, Ehrman SH (2012) Copper oxide nanoparticle made by flame spray pyrolysis for photoelectrochemical water splitting—Part I. CuO nanoparticle preparation. *Int J Hydro Ener* 37:4871–4879
- Cho S (2013) Optical and electrical properties of CuO thin films deposited at several growth temperatures by reactive RF magnetron sputtering. *Met Mater Int* 19:1327–1333
- Essic J, Mather R (1993) Characterization of a bulk semiconductors band gap via near-absorption edge optical transmission experiment. *Am J Phys* 61:646–649
- Gandhi S, Subramani RHH, Ramakrishnan T, Sivabalan A, Dhanalakshmi V, Nair MRG, Anbarasan R (2010) Ultrasound assisted one pot synthesis of nano-sized CuO and its nanocomposite with poly(vinyl alcohol). *J Mater Sci* 45:1688–1694
- Greenwood R (2003) Review of the measurement of zeta potentials in concentrated aqueous suspensions using electroacoustics. *Adv Colloid Interface Sci* 106:55–81
- Growth A (2011) structure and optical characterization of Al-doped ZnO nanoparticle thin films. *Crys Res Tech* 46:1086–1092
- Hamad TK, Yusop RM, Al-Taay WA, Abdullah B, Yousif E (2014) Laser Induced Modification of the Optical Properties of Nano-ZnO Doped PVC Films. *Int J Poly Sci* 2014:787595–787603
- Honsi HM, Fayek SA, El-Sayed SM, Roushdy M, Soliman MA (2006) Optical properties and DC electrical conductivity of Ge₂₈–xSe₇₂S_{bx} thin films. *Vacuum* 81:54–59
- Huang CY, Chatterjee A, Liu SB, Wu SY, Cheng CL (2010) Photoluminescence studies on a single CuO nanowire. *Appl Surf Sci* 256:3688–3692
- Kaur M, Muthe KP, Deshpande SK (2006) Growth and branching of CuO nanowires by thermal oxidation of copper. *J Cryst Growth* 289:670–675
- Khalil SG, Ameen MA, Jubier NJ (2011) Effect of laser irradiation on the optical properties of SnO₂ films deposited by post oxidation of metal films. *Baghdad Sci J* 8:1–8
- Kidowaki H, Oku T, Akiyama T, Suzuki A, Jeyadevan B, Cuya J (2012) Fabrication and characterization of CuO-based solar cells. *J Mater Sci Res* 1:138–143
- Lang XY, Zheng WT, Jiang Q (2006) Size and interface effects on ferromagnetic and antiferromagnetic transition temperatures. *Phys Rev B* 73:224444–224450
- Lim YF, Choi JJ, Hanrath T (2012) Facile Synthesis of Colloidal CuO Nanocrystals for Light-Harvesting Applications. *J Nanomater* 2012:393160–393166
- Lin HH, Wang CY, Shih HC, Chen JM, Hsieh CT (2004) Characterizing well-ordered CuO nanofibrils synthesized through gas-solid reactions. *J Appl Phys* 95:5889
- Lupi S, Ortolani M, Baldassarre L, Calvani P (2005) Optical conductivity and charge ordering in Na_xCoO₂. *Phys Rev B* 72:024550–024554
- Madhup DK, Subedi DP, Chimouriy SP (2010) Optical characterization and thickness estimation of Al³⁺ ion doped ZnO nanofilms from transmittance spectra. *J Optoelectron Adv Mat* 12:1035–1044
- Mahapatra O, Bhagat M, Gopalakrishnan C, Arunachalam KD (2008) Ultrafine dispersed CuO nanoparticles and their antibacterial activity. *J Exp Nanosci* 3:185–193
- McAuleya CB, Dub Y, Wildgoosea GG, Comptona RG (2008) The use of copper(II) oxide nanorod bundles for the non-enzymatic voltammetric sensing of carbohydrates and hydrogen peroxide. *Sen Acta B* 135:230–235
- Millis AJ, Zimmers A, Lobo RPSM, Bontemps N, Homes CC (2005) Mott physics and the optical conductivity of electron-doped cuprates. *Phys Rev B* 72:224517–224521
- Mishra RL, Mishra SK (2009) Optical and gas sensing characteristics of tin oxide nano-crystalline thin film. *J Ovonic Res* 5:77–85
- Morales J, Sanchez L, Martin F, Ramos-Barrado J, Sanchez M (2005) Use of low-temperature nanostructured CuO thin films deposited by spray-pyrolysis in lithium cells. *Thin Solid Films* 474:133–139
- Moss TS, Burrell GJ, Ellis B (1973) *Semiconductor opto-electronics*. Butterworth & Co Ltd., London
- Muhibullah M, Hakim MO, Choudhury MGM (2003) Studies on Seebeck effect in spray deposited CuO thin film on glass substrate. *Thin Solid Films* 423:103–110
- Qi JQ, Tian HY, Li LT, Lai H, Chan W (2007) Fabrication of CuO nanoparticle interlinked microsphere cages by solution method. *Nanoscale Res Lett* 2:107–111
- Radhakrishnan AA, Beena BB (2014) Structural and optical absorption analysis of CuO nanoparticles. *Ind J Adv Chem Sci* 2:158–161
- Son DI, You CH, Kim TW (2009) Structural, optical, and electronic properties of colloidal CuO nanoparticles formed by using a colloid-thermal synthesis process. *Appl Surf Sci* 255:8794–8797
- Stewart SJ, Multigner M, Marco JF, Berry FJ, Hernando A, Gonzalez JM (2004) Thermal dependence of the magnetization of antiferromagnetic copper (II) oxide nanoparticles. *Solid State Commun* 130:247–251
- Sumangala D, Amma D, Vaidyan VK, Manoj PK (2005) Structural, electrical and optical studies on chemically deposited tin oxide films from inorganic precursors. *Mater Chem Phys* 93:194–201
- Tauc J, Grigorovici R, Vancu A (1966) Optical properties and electronic structure of amorphous germanium. *Phys Status Solidi B* 15:627–634

- Teng F, Yao W, Zheng Y, Yutao M, Teng Y, Xu T, Liang S, Zhu Y (2008) Synthesis of flower-like CuO nanostructures as a sensitive sensor for catalysis. *Sens Actu B* 134:761–768
- Wang SQ, Zhang JY, Chen CH (2007) Dandelion-like hollow microspheres of CuO as anode material for lithium-ion batteries. *Scr Mater* 57:337–340
- Xu Y, Chen D, Jiao X, Xue K (2007) CuO microflowers composed of nanosheets: synthesis, characterization and formation mechanism. *Mater Res Bull* 42:1723–1731
- Xue SW, Zu XT, Zhou WT (2008) Effects of post-thermal annealing on the optical constants of ZnO thin film. *J. Alloys Compd* 448:21–26