Structural and optical characteristics of 1-(4-Methylsulfonyl Phenyl)-3-(4-n, n Dimethyl (amino Phenyl)-2-Prope n- 1-One (MSPPP) Chalcone doped ZnO nanoparticles

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Abstract: The purpose of this paper is to investigate the structural and optical characteristics of 1-(4-Methylsulfonyl Phenyl)-3-(4-n, n Dimethyl (amino Phenyl)-2-Prope n- 1-One (MSPPP) Chalcone doped in ZnO nanoparticles. Part of the aim is to study the characterization of chalcone doped ZnO nanoparticles by several techniques such as X-ray diffraction, Scanning electron microscope, FTIR spectroscopy, and diffuse reflection spectra. All doped samples showed a hexagonal wurtzite structure. This study has shown that the crystallite size of pure ZnO varied from 33.50 to 27.45 nm and when increasing the chalcone percentage by 0.5 and 1.5%, has increased the crystallite sizes in the range of 33.40–33.80 nm and 33.80–36.20 nm, respectively. The value of the energy gap (Eg) for ZnO nanoparticles was 3.14 eV. For 0.5 and 1.5% chalcone doped ZnO, the energy gap decreased by an order of magnitude 0.16 eV.

Keywords: ZnO, Spectroscopic studies, optical, Structural

1. Introduction

Zinc oxide (ZnO) nanoparticles plays a vital role and are extensively utilized in different applications because of their properties [1,2]. ZnO nanoparticle own a broad bandgap of 3.37 eV, a large exciton binding energy of 60 m eV and is an n-type semiconductor material [3,6]. Owing all these properties, it has found widely utilize in different technological application, like laser, photocatalyst, gas sensors and solar cell [7,8]. Additionally, compared to other metal oxides it is economical, easy to fabricate and environmentally [9]. By doping with various materials, the properties of ZnO can be tuned to acquire particular properties [10]. Doping can lead fundamental changes in ZnO’s physical behavior [6]. Most studies noted that variations happen by incorporation of transition metal ions into ZnO lattice [11,12]. Chalcone derivatives have a functionality of a carbonyl group in conjugation with a carbon-carbon double bond, which is known as α, β-unsaturated keto group. The desirable properties of the chalcones are: adding/removing donor can modify high conjugation level; and the photophysical characteristics of these chalcones or acceptor groups the skeleton [13, 14]. The doping of ZnO with other materials (heterostructures) has attracted significant interests from their predominant properties to combine two kinds of materials with lower photoexcitation threshold than individual components and decrease the recombination rate of electron–hole pairs by their physically separated band gap [15,16]. The essential purpose of the present work is to report on the photophysical properties of ZnO nanoparticles doped with the chalcone (MSPPP) in order to obtain novel properties through alloying and doping materials. This investigation is an attempt to highlight the behavior of chalcones doped ZnO nanoparticles as a novel material for optoelectronics applications.

2 Experimental Section

2.1 Chalcone synthesis

Chalcone material of 1-(4-methylsulfonyl phenyl)-3-(4-N, N dimethyl (amino phenyl)-2-prope n-1-one (MSPPP) was performed using a simple reaction of 4- dimethylamino benzaldehyde (0.01 mol) and acetophenones derivatives (0.01 mol) in a basic solution of sodium hydroxide (1.0 gm) and ethanol (50 ml). The product was recrystallized from ethanol and water, filtered, washed with water, and left to dry [17].
The molecular structure of the chalcone is given in Fig. 1. The product confirmed by H1 NMR and UV-Vis spectroscopy. 400 MHz, from 0 to 11 ppm) 
\[ \delta \] 2.86 (6H, s), 3.07 (3H, s), 6.47 (1H, d, J = 15.6 Hz), 6.84 (2H, ddd, J = 8.2, 1.2, 0.4 Hz), 7.47-7.58 (3H, 7.51 (d, J = 15.6 Hz), 7.55 (ddd, J = 8.2, 1.5, 0.5 Hz)), 7.81 (2H, ddd, J = 7.9, 1.9, 0.5 Hz), 8.01 (2H, ddd, J = 7.9, 1.5, 0.5 Hz). \[ \lambda_{\text{max}} \] (methanol 419 nm).

2.2 ZnO nanoparticle synthesis and doping mechanism

A certain amount of zinc acetate dihydrate was dissolved in 75 mL of methanol, after stirring for 50 min. Further, an amount of double distilled water solution (50 mL) of tartaric acid was added dropwise to the previous mixture. The gel formed rapidly [10 minutes] at room temperature under vigorous magnetic stirring and dried at 100°C for 12 h. Subsequently, the dry gel was first grinded and annealed at 550°C for 1 h. A 0.5 and 1.5 % of MSPPP chalcone were added to the prepared ZnO nanoparticle and milled for 5 h and annealed at 110°C for 1 h.

3 Result and Discussion

3.1 characterization of pure ZnO, 0.5 and 1.5 % chalcone doped ZnO

3.1.1 XRD analysis

Figure 2 displayed the structure of pure and chalcone doped ZnO nanoparticles using X-ray diffraction patterns. The peak positions of pure ZnO were found at 2 \( \theta \) of 31.55°, 34.32°, 36.19°, 47.33°, 56.43°, 62.77° and 67.91°. These peaks correspond to the (100), (002), (101), (102), (110), (103) and (112) planes of hexagonal wurtzite structure of ZnO, respectively [18].

| Sample                      | Orientation | \( \theta \) | Crystal size (nm) | Lattice parameters | c/a     |
|-----------------------------|-------------|--------------|-------------------|--------------------|---------|
|                             |             |              |                   | \( a_{100} \)      | \( c_{100} \) |
| ZnO                         | (100)       | 31.62        | 25.73             | 3.2646             | 1.600   |
|                             | (002)       | 34.27        | 23.48             |                    |         |
|                             | (101)       | 36.12        | 27.46             |                    |         |
| 0.5 % MSPPP chalcone - ZnO  | (100)       | 31.66        | 33.40             | 3.2605             | 1.602   |
|                             | (002)       | 34.31        | 33.55             |                    |         |
|                             | (101)       | 36.15        | 33.80             |                    |         |
| 1.5 % MSPPP chalcone - ZnO  | (100)       | 31.71        | 36.16             | 3.2566             | 1.602   |
|                             | (002)       | 34.36        | 33.83             |                    |         |
|                             | (101)       | 36.19        | 35.66             |                    |         |
The doped samples showed peaks similar to ZnO nanoparticle patterns and no new peak has been detected in the XRD patterns. The average crystallite size ($D$) was calculated using Scherrer’s [19].

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

where $\lambda$ is the wavelength of Cu Kα radiation, $\beta$ is the full width half maximum (FWHM) of the diffraction peak and $\theta$ is the Bragg peak angle. Table 1 displayed the average crystallite size ($D$). The crystallite size of pure ZnO varied from 23.48 to 27.46 nm. Increasing the chalcone percentage by 0.5 and 1.5% has increased the crystallite sizes in the range of 33.40–33.80 nm and 33.83–36.16 nm, respectively.

The lattice parameters $a$ and $c$ can be calculated by using Eqs. (2) and (3) [20]:

$$a = \frac{\lambda}{\sqrt{3 \sin \theta}}$$  \hspace{1cm} (2)

$$c = \frac{\lambda}{\sin \theta}$$  \hspace{1cm} (3)

### 3.1.2 SEM of materials

Undoped, 0.5 and 1.5% MSPPP chalcone doped ZnO nanoparticles SEM images are shown in Fig. 3 (a – c). The high magnification SEM images show that the entangled ZnO nanoparticles appear in the form of a clear hexagonal structure. The morphology of the chalcone doped ZnO samples changed and the surface area decreased with increasing the particle size. This decrease in the surface area can be attributed to the molecular structural disorder and lattice strains attributed to chalcone dopant. However, the dopant samples showed more surface porosity compared with ZnO.

### 3.1.3 FTIR of ZnO and MSPPP chalcone doped – ZnO

FTIR was used to measure the vibration and bending modes of the structure of ZnO, 0.5 and 1.5% doped-ZnO at room temperature. The spectra are shown in Fig. 4. The broad absorption band at 3439.39, 1077.74 cm$^{-1}$, can be attributed to the normal polymeric O–H stretching vibration of H$_2$O, in ZnO lattices [21]. Other sharp peaks observed at 1621.45 and 1615.60 cm$^{-1}$ can be attributed to H–O–H bending vibration, which in turn can be assigned to the small amount of H$_2$O in the ZnO nanocrystal, also, the absorption band observed between 2300 and 2400 cm$^{-1}$ are due to the existence of CO$_2$ molecule in the air. The vibration band at 446.31 cm$^{-1}$ assigned to the stretching mode of pure ZnO [21] while the band shifted to 445.51 and 450.18 cm$^{-1}$ for the dopant 0.5 and 1.5%, respectively. This shift might be attributed to the formation of a new bond between the chalcone and ZnO (Table 2).
3.1.4 Optical Properties

The optical absorption spectra are shown in Fig. 5 with wavelength range 300 nm – 1100 nm. When the ZnO was doped with the chalcone by a ratio of 0.5%, the absorption edge (395 nm) red shifted to 415 nm. When the dopant was increased to 1.5%, the absorption edge of the spectrum remained unchanged. The band gaps of the fabricated nanoparticles were calculated using Tauc Eq. [22].

\[
\alpha h\nu = B (h\nu - E_g)^n
\]

where \(\alpha\) is absorption coefficient, \(E_g\) represent optical band gap, \(B\) is band tailing parameter, \(h\nu\) is the photon energy, and \(n = 0.5\) for direct band gap. The energy band gap is determined by extrapolating the straight portion of the curve between \((\alpha h\nu)^2\) and \(h\nu\) when \(\alpha\) equal to zero. The value of the energy gap \((E_g)\) for ZnO nanoparticles was 3.14 eV. This value is in a good agreement with some previous studies [4, 23]. For 0.5 and 1.5% chalcone doped ZnO, the energy gap decreased by an order of magnitude 0.16 eV as seen in Fig. 6. However, the introduction of chalcone on ZnO increased the optical density of the absorption spectrum in the visible region. The good absorption in the UV and visible regions for the chalcone doped ZnO can open a new window for solar cell applications.
Table 2 FTIR for all specimens

| Assignment                        | Wavenumber (cm⁻¹) |
|-----------------------------------|-------------------|
|                                   | Pure ZnO | 0.5% MSPPP chalcone - ZnO | 1.5% MSPPP chalcone - ZnO |
| O – H stretching modes            | 3443.34   | 3435.01                     | 3435.25                     |
| CO₂ stretching modes              | 2343.19   | 2351.45                     | 2357.68                     |
| Zn – O stretching vibration       | 445.98    | 447.55                      | 453.21                      |
| H – O – H bending vibration       | 1077.67   | 1031.26                     | 1023.12                     |

Figure 5 UV – Vis diffuse reflection spectra of ZnO and MSPPP chalcone doped ZnO

Figure 6 (ahv)² against photon energy (eV) in ZnO and chalcone doped ZnO
4 Conclusion

A chalcone derivative, of 1-(4-Methylsulfonyl Phenyl)-3-(4-n, n Dimethyl (amino Phenyl)-2-Propen-1-One (MSPPP) was prepared and added to pure ZnO nanoparticles with 0.5 and 1.5 wt% doping percentages. It was found that the value of the energy gap (Eg) for ZnO nanoparticles was 3.14 eV and for 0.5 and 1.5% chalcone doped ZnO, the energy gap decreased by an order of magnitude 0.16 eV. Also, it was found that the increasing dopant percentage distorted the hexagonal shape of ZnO nanoparticles. Further, the lattice parameters and microstrain were decreased while the crystal size was increased. Chalcone loading on ZnO has decreased the energy bandgap.

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Authors’ contribution

This work was carried out in collaboration among all authors. Authors Mohana Attia, Abdelrahman A. Elbadawi conceived of the presented idea. Author MFA developed the theory and performed the computations and verified the analytical methods. All authors discussed the results and contributed to the final manuscript.

Competing Interests

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Yes

Conflict of interest

The author declares that there is no conflict of interest anywhere.

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