Synthesis and Characterization of Chromium Doped Zinc Sulfide Nanoparticles

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Abstract
Nanoparticles of pure and Zn$_{1-x}$Cr$_x$S with concentrations (x = 0.00, 0.02, 0.08 and 0.10) were prepared successfully by co-precipitation technique. Crystallographic study was intervened by X-ray diffraction spectroscopy. There were no extra peaks observed in the XRD pattern. From XRD peaks it is confirmed that the samples prepared were highly pure and cubic in structure. The particle size calculated from XRD data was around 3 nm. It confirms that the quantum dots were formed. Surface morphology and crystal size were verified by scanning electron microscopic (SEM) and transmission electron microscopic (TEM) studies. The aggregation of the nanoparticles was observed, which is due to the quantum dots. Chemical species present in the prepared samples were identified by FTIR study. Using UV-Vis spectrometer (Ultra Violet Visible), absorption spectra were obtained for x = 0.00, 0.02, 0.08 and 0.1. It is confirmed from UV-Vis that the detected absorption peaks were shifted towards higher wavelengths. The energy band gap values were estimated using UV-Vis spectrometer and found lying in the range of 3.33 eV - 3.02 eV. The energy band gap decreases with increasing Cr concentrations i.e. the red shift was observed.

Keywords
Nanoparticles, Co-Precipitation, Energy Band Gap, Red Shift, Quantum Dots

Subject Areas: Nanometer Materials

1. Introduction
ZnS is one of the most important wide band gap semiconductors from II-VI group elements [1]. Among the most studied host materials for diluted magnetic semiconductor (DMS), it means some magnetic materials with appropriate content doped in semiconducting materials known as (DMS). ZnS is a potential candidate for device

applications because of its wide band gap of 3.7 eV, large exciton binding energy of 40 meV, high index of refraction of 2.27 at 1 μm and ZnS has been identified as an excellent host semiconductor for supporting room-temperature ferromagnetism when doped with a variety of 3d transition metal ions [2]. It is chemically more stable, nontoxic and environmentally safer than other II-VI compound semiconductors. Hence it finds more potential applications in biological detection [3]. Doping with proper element is widely used as an effective method to tune surface states, energy levels, electrical, optical and magnetic properties of semiconducting materials. The optical and magnetic properties of doped semiconductor nanostructures are sensitive to size, morphology, crystal structure, and effects, which may provide new opportunities for investigation and application of ZnS-based nanomaterials.

The zinc chalcogenides ZnS, ZnSe and ZnTe are semiconductors with zinc blende structure at ambient conditions. The stability of high pressure phases of these zinc chalcogenides ZnX (X = S, Se, Te) appears to be more complex. Chalcogenides such as ZnSe, ZnS have been widely used as important candidates due to their better chemical stability [4]-[6] for photoluminescence (PL), electroluminescence (EL) and cathodoluminescence (CL) devices. ZnS is one of the most promising luminescent materials among II-VI groups, because it exhibits wide optical transparency from the visible light of 0.4 μm to the deep infrared region of 12 μm. Doped ZnS semiconductor materials have a wide range of applications in phosphors, light emitting displays, optical sensors and also are used in optoelectronic applications such as reflector, dielectric filter and window materials.

Efficient doping of rare-earth ions into II-VI semiconductor host is not favourable compared to the transition metals [5]. In view of this, transition metal doped ZnS, like ZnS:Mn, ZnS:Cu, ZnS:Co and ZnS:Ni has been widely investigated. However, surprisingly, in spite of both applied and academic interest, Cr doped ZnS has been least studied. There are only a few reports [7] [8] on Cr doped ZnS and all these reports are on ZnS:Cr prepared by physical methods only. Surprisingly none of the earlier authors have used the chemical synthesis, although chemical route is simple and capable of yielding nanoparticles of higher purity.

In the present study we have chosen chemical co-precipitation route for synthesis of Cr doped ZnS nanocrystals because this route is easy to handle and cost-effective and takes less time. It was investigated from XRD; samples were cubic in structure and highly pure. Fourier Transform Infra-Red (FTIR) spectra confirm that Cr goes into ZnO lattice structure. Scanning electron microscopy (SEM) outlines, aggregation of the particles in all samples were observed. It is confirmed from transmission electron microscopic study reports that the crystal size obtained from XRD data and that obtained from TEM study are almost the same, which is around 3 nm. The SAED patterns consist of three concentric sharp rings, which correspond to the (111), (220) and (311) planes of diffraction peaks which belong to cubic structure. In the optical study it is observed that energy band gap decreases from 3.33 eV to 3.02 eV with increasing Cr concentration; red shift in the band gap is detected. Based on density-function calculations Cr is theoretically proposed to be a reliable dopant in ZnS [9]. Such results on Cr doped and undoped ZnS nanoparticles are reported in this paper qualitatively and quantitatively.

2. Materials and Methods

Samples Zn_{1-x}Cr_xS nanoparticles were synthesized by chemical co-precipitation route at room temperature. The starting materials were Zinc nitrate, chromium nitrate and sodium sulfide. Chemicals used without further purification were of analytical grade. The chemical used as solvent was methanol. The appropriate amounts of zinc nitrate, chromium nitrate and sodium sulfide respectively were dissolved separately in 100 ml methanol and stirred at room temperature for 2 h. Chromium nitrate solution was added into zinc nitrate and stirred for 20 min. Sodium sulfide solutions then added into as prepared solutions of zinc nitrate and chromium nitrate drop wise with vigorous stirring to get homogeneous solution. Throughout the sample preparation, pH-10 of the solution was maintained. The light sky-blue coloured precipitate was obtained; the colour of the precipitate becomes darken as Cr concentration increases. The precipitate was continuously stirred for 2 h again to get homogeneous particle size. The precipitate was collected and washed several times by deionised water followed by methanol. The washed precipitate was dried at 50°C for 4 hours in the oven and ground for 15 minutes to get Cr doped ZnS nanoparticles finally.

3. Results, Discussion

3.1. Structural Study

The X-ray diffraction (XRD) patterns for pure and Cr doped ZnS samples were recorded at room temperature as
shown in Figure 1. From the peak broadening diffraction it is confirmed that the crystallites were nanosized. The X-ray diffraction patterns of pure and Cr doped ZnS nanoparticles were recorded by X-ray diffraction spectrometer (make: Bruker) using CuKα radiation (λ = 1.5406 Å) with 2θ ranging 20° - 80°. The observed diffraction peaks corresponded to (111), (220) and (311) planes are in good agreement with the standard diffraction peaks cubical structure. As well as the d-spacing values and relative intensities of the peaks coincide with the JCPDS data (00-001-0792) of cubic structure. The prepared samples were highly pure because no extra peaks observed in XRD pattern. The lattice parameters, volume cell, X-ray density, grain size were determined using XRD data & their values are summarized in Table 1.

The (111) diffraction peaks were shifted to higher 2θ values as Cr concentration increases into ZnS than pure ZnS sample. It may be due to the small ionic radii of Cr³⁺ (0.63 Å) as compared to that of Zn²⁺ (0.74 Å), which have entered into ZnS. A. Rahdar [10] reported that the Cr doped ZnS nanoparticles synthesized by co-precipitation route was wurtzite (hexagonal) structure. But in this present work the XRD pattern is in good agreement

![Figure 1. XRD pattern of pure and Cr doped ZnS nanoparticles.](image_url)

<table>
<thead>
<tr>
<th>Lattice parameter “a” (nm)</th>
<th>Volume (Å³)</th>
<th>X-ray density (gm/cm³)</th>
<th>Grain size (nm)</th>
<th>% of crystallinity</th>
<th>% of amorphous</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.53913</td>
<td>156.151</td>
<td>1.0366</td>
<td>3.853</td>
<td>57.4</td>
<td>42.6</td>
</tr>
<tr>
<td>0.53758</td>
<td>155.360</td>
<td>1.0391</td>
<td>3.520</td>
<td>57.6</td>
<td>42.4</td>
</tr>
<tr>
<td>0.53777</td>
<td>155.466</td>
<td>1.0101</td>
<td>3.653</td>
<td>57.6</td>
<td>42.4</td>
</tr>
<tr>
<td>0.53423</td>
<td>152.475</td>
<td>1.0470</td>
<td>3.886</td>
<td>56.6</td>
<td>43.4</td>
</tr>
</tbody>
</table>
with the diffraction data of cubical structure and the cell parameter “a” is very close to the reported value in JCPDS card data.

Grain size of the nanoparticles was determined by the Debye-Scherrer’s equation,

\[ D = \frac{K \lambda}{\beta \cos \theta} \]  

where \( K \) is shaping factor and having value 0.9.

Figure 2 shows the plots (a) lattice constant “a” verses Cr concentration (b) volume of unit cell verses Cr concentration (c) X-ray density verses Cr concentration and (d) grain size verses Cr concentration. Lattice constant “a” and volume of the unit cell decreases gradually from pure ZnS to Cr concentration 0.10 except 0.08 X-ray density increases with Cr concentration except 0.08 and the average grain size of nanoparticles for Cr concentration 0.02 is less than pure ZnS, and Cr concentrations 0.08 and 0.10 as shown in Figure 2.

Two more parameters are recorded by X-ray diffraction spectrometer are crystallinity % and amorphousness % of the prepared samples. From these recorded values it is clear that the crystallinity % is more than the value amorphousness % of the pure and Cr doped ZnS nanoparticles.

3.2. Morphological Study

Scanning electron microscopy was employed to get scanning electron micrographs of pure and Cr doped ZnS
samples with 5 µm magnification as shown in Figure 3. It was clearly seen from SEM images that as synthesized samples exhibit the aggregation of the particles at room temperature. It may be due to the smaller particle size which is around 3 nm. The transmission electron microscopy (TEM) images were obtained using JEOL-2010 transmission electron microscope at an acceleration voltage of 200 kV with magnification 2 nm and 50 nm for pure and Cr doped ZnS nanoparticles as shown in the Figure 4.

The selected area electron diffraction (SAED) patterns for each sample are shown Figure 4. The selected area electron diffraction (SAED) patterns consist of three concentric sharp rings, which corresponded to the (111), (220) and (311) planes of diffraction belong to the cubic structure. Therefore, it can be further confirmed that the
nanoparticles are in cubical structure. It is highly confirmed that as synthesized samples are pure and no secondary phases confines. It is found that the Cr goes to the ZnS sites. The results from Transmission electron microscopic TEM study are in good agreement with XRD results.

4. Optical Study

4.1. UV-Vis Investigation

The optical study of pure and Cr doped ZnS samples were carried out by ultra violet visible spectrometer. The optical measurements were done with UV spectrometer [Make: JASCO UV-VIS-NIR Spectrophotometer Model: V-670 Serial No. A009761154] in the wavelength range of 200 nm to 1000 nm. The UV spectra of pure and Cr doped ZnS samples are plotted. The absorption spectra for pure and Cr doped ZnS is as shown in Figure 5. Tauc’s plots are as shown in Figure 6. The energy band gap was calculated from the Tauc’s plots. The energy band gap values are in the range 3.33 eV - 3.02 eV. The band gap increase with increasing Cr concentration is attributed to size quantization effect due to the smaller size of the particles [11]. In this present work even though the particles are quantum dots the band gap from Tauc’s plots of pure ZnS and Cr doped ZnS samples were 3.33 eV, 3.20 eV, 3.05 eV and 3.02 eV for Cr concentrations 0.00, 0.02, 0.08 and 0.1 respectively. It means

![Figure 5. Absorption pattern of pure and Cr doped ZnS nanoparticles.](image)

![Figure 6. Tauc’s plots for pure and Cr doped ZnS nanoparticles.](image)
that the band gap decreases with increase in Cr concentration in ZnS lattice. This is due to the s-p and d interactions and the quantum confinement effect. The red shift in the band gap confirms the uniform substitution of Cr ions in the ZnS lattice structure.

4.2. FTIR Spectra Analysis

Identifying the absorbed species into the crystal surface and to confirm the formation of crystalline ZnO nanoparticles Fourier transform infrared spectroscopy study was carried out. FTIR (Make: JASCO FTIR, JAPAN) spectra were recorded at room temperature to obtain information about the chemical bonding and chemical species in a material. FTIR absorption spectra of pure and Cr doped ZnO nanoparticles are as shown in the Figure 7 and the peak values are listed in the Table 2.

Broad bands around 3444 cm\(^{-1}\) and 3480 cm\(^{-1}\) appear typically due to O-H vibrations of EDTA and water, because all FTIR spectra were carried out by mixing samples with KBr, some water vapour may be absorbed as KBr is hygroscopic. The additional weak bands and shoulders are observed at 1626 cm\(^{-1}\) and 1639 cm\(^{-1}\) it may be due to micro structural formation of the samples. The C-H bending of methylene in ME is observed at 1407 cm\(^{-1}\) band. The spectra at 1149 cm\(^{-1}\) and 1141 cm\(^{-1}\) were observed, it due to characterization size frequency inorganic ions. The bands 1011 cm\(^{-1}\) and 1040 cm\(^{-1}\) observed due C-O stretching. The spectra at 657 cm\(^{-1}\) and

![Figure 7. FTIR spectra for pure and Cr doped ZnS nanoparticles.](image)

<table>
<thead>
<tr>
<th>FTIR peaks observed</th>
<th>X = 0.00</th>
<th>X = 0.02</th>
<th>X = 0.08</th>
<th>X = 0.1</th>
</tr>
</thead>
<tbody>
<tr>
<td>OH-vibrations</td>
<td>3444.36</td>
<td>3457.47</td>
<td>3455.30</td>
<td>3480.313</td>
</tr>
<tr>
<td>CO(_2) mode</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>2103.36</td>
</tr>
<tr>
<td>Additional weak bands</td>
<td>1626.22</td>
<td>1626.22</td>
<td>1630.51</td>
<td>1639.68</td>
</tr>
<tr>
<td>C-H bending</td>
<td>-</td>
<td>1366.313</td>
<td>1365.636</td>
<td>1407.78</td>
</tr>
<tr>
<td>Inorganic ions</td>
<td>1141.17</td>
<td>1141.17</td>
<td>-</td>
<td>1149.24</td>
</tr>
<tr>
<td>CO stretching</td>
<td>1011.09</td>
<td>-</td>
<td>1013.11</td>
<td>1040.52</td>
</tr>
<tr>
<td>ZnS band</td>
<td>657.14</td>
<td>657.14</td>
<td>668.70</td>
<td>-</td>
</tr>
</tbody>
</table>
are ZnS bands that correspond to sulfide. One extra band observed in FTIR spectra for Cr concentration 0.10 at 2103 cm\(^{-1}\) assigned due to CO\(_2\) mode, it may be due to the CO\(_2\) absorbed from atmosphere while doing FTIR measurements. FTIR spectra of the pure and 10% Cr doped ZnS nanoparticle samples of present investigation are in good agreement with the reported literature [8] [9] [12] [13].

5. Conclusion

Cr doped ZnS nanocrystals were prepared using co-precipitation route successfully. No extra phases were observed in XRD patterns. The samples were cubic in structure and highly pure; it is confirmed from XRD study. From FTIR study it is intervened that the prepared samples were highly pure. The aggregation of the particles in all samples was observed form scanning electron microscopy. From transmission electron microscopic study it is reported that the crystal size obtained from XRD data and that obtained from TEM study are almost the same, which is around 3 nm. The SAED patterns consist of three concentric sharp rings, which correspond to the (111), (220) and (311) planes of diffraction peaks which belong to cubic structure. In the optical study it is observed that energy band gap decreases from 3.33 eV to 3.02 eV with increasing Cr concentration; red shift in the band gap is detected.

References


