



SYNTHESIS AND CHARACTERIZATION OF PVA CAPPED CdS NANOPARTICLES

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ABSTRACT

CdS is an important II-VI semiconductor and has been highlighted as an efficient emitter because of its high quantum yield and direct band gap in the visible range. In the present work, PVA capped CdS nanoparticles have been synthesized using Simple Chemical route. This paper presents the study of the structural and optical properties of PVA capped cadmium sulphide nanoparticles. XRD pattern showed the hexagonal phase of CdS nanoparticles. The band gap of the nano crystalline material is determined from the UV spectrograph. The calculated grain size is found to be 3.62 nm.

Keywords: Nanocrystals, PVA, SEM, XRD, UV-Vis spectroscopy, FTIR.

1. INTRODUCTION

The semiconductor nanoparticles exhibit structural, optical, electronic, Luminescence and photo-conducting properties [1] very different from their bulk properties. It is also attractive because of their possible application in solar cell, photo-detector, Laser, LED, high density magnetic information storage and many others in semi conducting industries [2]. Quantum dots of II-VI semiconductors have d particular attention, because they are easy to synthesize in the size range required for quantum confinement. So II-VI semiconductors with dimension in the nanometer range have generated considerable interest for researcher and scientific community. CdS nanoparticles exhibit size dependent properties it has a band gap energy E_g of 2.42 eV at room temperature and pressure. Since CdS has wide band gap, it is used as window material for heterojunction solar cells to avoid the recombination of photo-generated carriers which improves the solar cell efficiency [3].



Semiconductor nanoparticles are themselves highly unstable, and in the absence of capping agent, they agglomerate very rapidly. For this reason, bonding of capping agents to nanoparticles is necessary to provide chemical passivation and also to improve the surface state which has substantial influence on the optical and the electronic properties of nanoparticles. Polymer capped nanoparticles composites have been increasingly studied [4].

2. SYNTHESIS: MATERIALS AND METHODS

Cadmium Sulphide nanoparticles were grown by Chemical Bath method using PVA as a capping agent. All the chemical reagents used were of analytical grade. For the undoped CdS nanoparticles, the solution of CdSO₄ (0.5 M) was prepared by dissolving CdSO₄ in DI water. The matrix solution was prepared by adding the CdSO₄ solution to an aqueous solution (2%) of PVA with constant stirring at room temperature to form a well dispersed PVA capped Cd²⁺ ion solution which is a transparent liquid indicating the complete dissolution of CdSO₄. The pH of the solution was maintained at around 8 by slowly adding Ammonium Buffer Solution drop by drop to the above matrix solution to form the metallic complex. The thiourea as S²⁻ ion source was then added to the above metallic complex solution to form colloidal solution of CdS nanoparticles.

3. CHARACTERIZATION

The prepared sample was taken for XRD study and UV studies. The structure of the sample is determined by X-ray diffraction measurements have been performed by using Xpert PANalytical instrument operating at 40 kV and current of 30 mA with CuK α radiation. FTIR measurements are recorded using FTIR-8400s SHIMADZH (IR solution). UV-Visible spectrum of the nanoparticles is recorded using PERKIN ELMER UV-Vis spectrophotometer.

4. RESULTS AND DISCUSSION

SEM ANALYSIS:

Scanning electron microscope (SEM) study: SEM is a type of electron microscope that images the sample surface by scanning it with a high-energy beam of electrons in a raster scan pattern. Here, it is used to characterize the morphology of the nanoparticles.

Fig.1 shows the SEM micrographs of the CdS nanoparticles. The grain size was measured from SEM photograph by keeping the photograph under traveling microscope having high accuracy. It was observed that the grain size have nano-metric dimensions ranging from 10.26 nm to 22.12 nm.

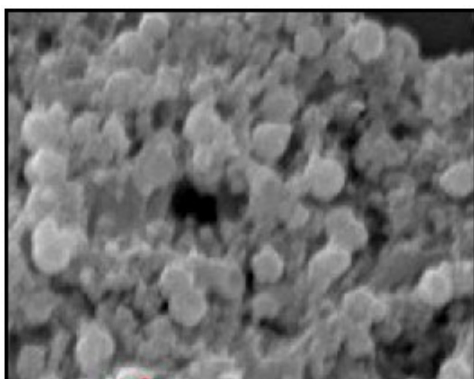


Figure 1: SEM image of PVA capped CdS sample.

UV- Visible STUDY:

Considering the band gap energy of bulk CdS as 2.42 eV, large blue shift is observed in both CdS and Cu doped CdS samples as shown in figure.. This indicates a clear quantum confinement in PVA capped CdS nanostructures. C.M. Janet *et al* reported that when the size of CdS nanocrystal becomes smaller than the exciton radius a remarkable quantum size effect leads to a size dependent increase in the band gap and a blue shift in the absorption onset [5].

Large blue shift is observed in CdS sample with respect to bulk CdS. This indicates a clear quantum confinement in PVA capped CdS nanostructures. C.M. Janet et al also reported that, the size of CdS nanocrystal becomes smaller than the exciton radius a remarkable quantum size effect leads to a size dependent increase in the band gap and a blue shift in the absorption onset [6].

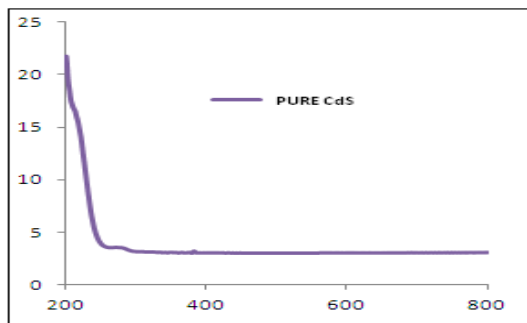


Figure 2. Absorption spectra of PVA capped CdS nanoparticles
XRD STUDY:

The structure of the prepared sample was investigated by X-Ray Diffractometer technique. The XRD pattern of the sample is shown in fig. The particle size was calculated by using Debye- Scherrer formula.

$$D = k\lambda / \beta \cos\theta$$

Where, k is a dimensionless constant, λ is the wavelength of X-Ray used, β = Full width at Half maxima (FWHM) of the diffraction peak and θ is the diffraction angle for the (h, k, l) plane Bragg's angle. XRD shows hexagonal CdS nanoparticles with size in a few nanometers range. Significant peaks were obtained at 2θ angles 26° , 29° , 32° , 36° , 44° , 51.5° , 61.5° corresponds to the reflections at (002), (101), (200), (102), (110), (112), (202) planes.

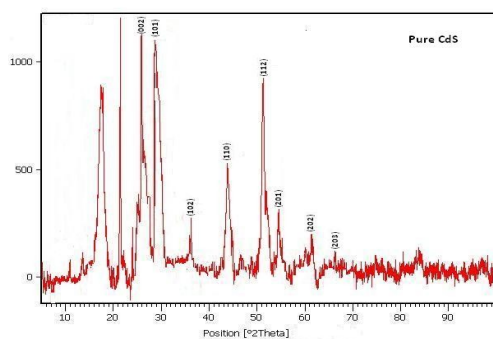


Figure 3: XRD spectra for PVA capped CdS sample.

FTIR STUDY:

In FTIR spectrum, the peak at 3431 cm^{-1} is assigned to O-H stretching of absorbed water on the surface of the sample and the peak at 1438 cm^{-1} is attributed to bending vibrations of Poly Vinyl Alcohol used in the process. The C-O stretching vibration of absorbed PVA molecule gives its intense peak at 1025 cm^{-1} . In addition to surface coverage of CdS by PVA, presence of trace amount of template ligand namely PVA is also evident, its ring C-H vibration occurs at about 3074 cm^{-1} which is a very weak peak. The FT-IR spectra for PVA capped CdS sample is shown in figure 4.

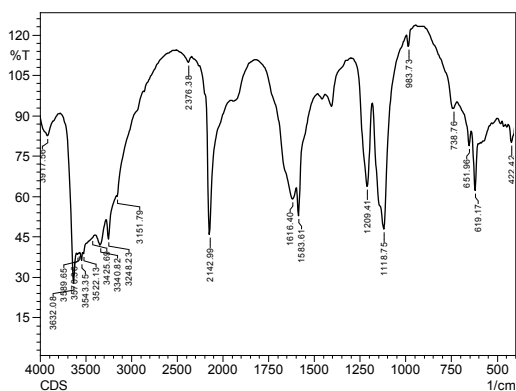


Figure 4: FTIR spectra for PVA capped CdS sample.



5. CONCLUSIONS

CdS nanoparticles have been synthesized in the PVA matrix through simple chemical route. X-ray diffraction confirms the hexagonal structure. UV-visible absorption showed a blue shift indicating quantum confinement of the particle. The obtained results can be useful for the started point of photonic device fabrication.

6. REFERENCES

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