Solvothermal growth of single-crystal CdS nanowires

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Abstract. Cadmium sulfide (CdS) nanowires (NWs) were prepared by the solvothermal method using ethylenediamine as a solvent. Two sets of CdS NWs were synthesized at 160 and 200 °C for various reaction durations (3.5, 7 and 24 h). Scanning/tunneling electron microscopy was used to examine the surface morphology of the grown NWs. Their dimensions are found to depend on the reaction temperature and duration. The CdS NWs grown at 200 °C for all durations are longer than those prepared at 160 °C, with diameters ranging from 15 to 40 nm. A three-armed structure is exhibited by all the samples. The grown CdS NWs display a hexagonal wurtzite phase and grows along the (001) direction. The optical absorption of the grown NWs shows a sharp absorption edge with a blueshift, which indicates an expansion of the optical band gap. All prepared samples show two emission peaks in their photoluminescence spectra. The emission peak location depends on the reaction temperature and duration. The CdS NWs prepared at 160 °C show a sharp band–band emission compared with those prepared at 200 °C. Raman analysis indicates that the optical properties of the grown NWs are enhanced with increased temperature and reaction duration.

Keywords. Cadmium sulfide; nanowire; solvothermal method.

1. Introduction

Considerable interest is focused on the synthesis of nanocrystalline materials because of their unique physical and chemical properties that distinguish them from bulk-phase materials. The particle size and shape of materials significantly affect their properties. Thus, control of the size and morphology of nanocrystalline materials can lead to the discovery of new physical and chemical properties (Wang *et al* 2011). One-dimensional (1D) semiconductors such as nanowires (NWs), nanorods and nanoribbons have several unique advantages, including high crystallinity, self-assembly, high surface-to-volume ratio, quantum confinement effects and slow electron–hole recombination (Choi *et al* 2010; Mahdi *et al* 2012a; Rai *et al* 2012). The ability to grow 1D, single-crystal and nanostructured semiconductors has led to the fabrication of functional nanoscale devices (Li *et al* 2005, 2010; Yingkai *et al* 2006; Xi *et al* 2010; Mahdi *et al* 2012a). Among group II–VI semiconductor materials, cadmium sulfide (CdS; direct band gap = 2.42 eV), is considered to be an excellent material for various optoelectronic applications within the visible range of the electromagnetic spectrum (Yan *et al* 2009; Mahdi *et al* 2012b). 1D CdS has been prepared using

various methods, such as D.C. electrochemical techniques (Mondal *et al* 2007a), thermal evaporation (Kar and Chaudhuri 2006), chemical vapour deposition (Ma *et al* 2007) and hydro/solvothermal methods (Qingqing *et al* 2005; Xu *et al* 2005; Nirmala Jothi *et al* 2011). Solvothermal synthesis is widely used to prepare semiconductor NWs, particularly CdS. This synthesis can enable the control wire morphology and dimensions via reaction variables such as solvent type, temperature and duration (Datta *et al* 2009). The solvo/hydrothermal method is also simple, easy to scale up, low-energy consuming and cost-effective (Yan *et al* 2009; Chen and Wu 2011). In the solvothermal method, preparation parameters such as reaction temperature, duration and solvent type have crucial functions in determining the nature of the product as well as its physical properties. The solvothermal method can produce high-purity nanocrystalline materials with a high degree of crystallinity compared with other solution-based methods (Dalvand *et al* 2011). In the present work, CdS NWs were grown by the solvothermal method. The effects of the reaction temperature and duration on NW morphology, crystalline structure and optical properties were investigated.

2. Materials and methods

Cadmium nitrate [Cd(NO₃)₂·4H₂O] and sulfur (S) powder were used as Cd²⁺ and S^{2−} ion sources, respectively. Approximately 0.22 g of [Cd(NO₃)₂·4H₂O] and 0.066 g of S

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powder were dissolved in 50 ml of ethylenediamine (en) and stirred for 10 min. The solution was poured into a 100 ml teflon-lined stainless steel autoclave filled with en to 60% of its total volume capacity. The autoclave was sealed and subsequently loaded into an oven. Two sets of samples were prepared at different reaction temperatures and durations. The first set was prepared at 160 °C for 3.5, 7 and 24 h, whereas the other set was grown at 200 °C for the same durations. After removing the autoclave from the oven and cooling naturally, the resulting yellow precipitates were filtered, washed several times with absolute ethanol as well as distilled water and then air dried at 60 °C for 2 h. A

field-emission scanning electron microscopy (FESEM) system (FEI Nova NanoSEM 450) was used to determine the morphology of the products. The microstructures were investigated by a transmission electron microscopy (TEM) system (Libra 120-Carl Zeiss). The crystalline structure of the prepared CdS nanostructures was investigated by X-ray diffraction (XRD) using a PANalytical X'Pert PRO MRD PW3040 instrument with CuK α radiation. Optical properties such as absorption, photoluminescence (PL) and Raman spectra were measured using a Shimadzu UV-Vis spectrophotometer (UV-1800) and Horiba Jobin Yvon H 800 UV equipment.

Figure 1. FESEM images of CdS nanowires (A, B and C) prepared at 160 °C, (D, E and F) at 200 °C for 3.5, 7.0 and 24 h.

3. Results and discussion

3.1 Surface morphology

Figure 1 shows the FESEM images of the CdS NWs grown at 160 and 200 °C for various reaction durations. Figure 1(A–C) shows the CdS NWs prepared at 160 °C for 3.5, 7 and 24 h, respectively; figure 1(D–F) displays the CdS NWs prepared at 200 °C for the same durations. The results show that the dimensions of the grown CdS NWs depend on the reaction temperature and duration. The grown NWs exhibit uniform shapes and similar diameters, as clearly shown by the TEM images (figure 2). The TEM image of the sample prepared at 160 °C for 3.5 h contains a number of nanoparticles in addition to the NW structure. In the solvothermal method, two

mechanisms are involved in the formation of the 1D structure. Cd^{2+} ions interact with the lone pair of electrons of the en nitrogen atoms to form a Cd(en)_2^{2+} complex. S^{2-} ions are then slowly released into the solution, consequently, increasing the temperature and resulting in the combination of the Cd^{2+} and S^{2-} ions to form a two-dimensional (2D), CdS-0.5en complex structure, as shown in the following reaction:

This complex has an organic–inorganic lamellar structure with inorganic CdS layers separated by organic en spacers (Kar *et al* 2008). The 2D complex structure becomes unstable with increased temperature until around 120 °C; thus, the

Figure 2. TEM images of CdS nanowires (A, B and C) prepared at 160 °C, (D, E and F) at 200 °C for 3.5, 7.0 and 24 h.

complex fragments into a 1D needle-like structure to achieve stability. The needle-like structure then grows to form NWs due to the high surface energy. Mahdi *et al* (2012c) grown CdS nanostructures by solvothermal and they noted that the CdS prepared at 100 and 120 °C grew into nanosheets, and those prepared at 160 and 200 °C developed a nanowire structure. On the other hand, a number of uncoupled Cd²⁺ ions directly react with S²⁻ ions to form CdS nanoparticles. These nanoparticles also serve as nuclei for the growth of CdS NWs in solution. A three-armed nanorod structure is also shown in the TEM images (figure 3). This three-armed structure frequently appears in all the samples prepared at 160 and 200 °C. The 1D CdS structure grows into nanorods, as clearly shown by the hexagonal shape in figure 3. The three-armed structure is due to the presence of CdS nanoparticles in the solution, which acts as nuclei for the growth of CdS nanorods. Chen *et al* (2002) grew CdS nanorods by solvothermal synthesis using CdS nanoparticles as seeds to nucleate CdS nanorods, and they obtained tetrapod structures. The NW diameter distribution analysis shows that the diameters of CdS NWs grown at 160 °C for 3.5 and 24 h range from 10 to 30 nm, whereas those of CdS nanorods prepared for 7 h range from 15 to 40 nm (figure 4). This difference in diameter may be due to the higher concentration of the three-armed structure in the sample prepared for 7 h than the other samples. Interestingly, the NWs shorten with increased reaction duration to 24 h and some CdS nanosheets appear. The longer duration may have caused the wire to break into shorter pieces. The appearance of a number of CdS nanosheets may have been due to the growth of sheets that do not break into the needle structure. However, the reason for the absence of nanosheet structures in the sample prepared for 7 h is unclear. Figure 2(D–F) show that the CdS NWs grown at 200 °C for 3.5, 7 and 24 h, respectively, are longer than those prepared at 160 °C. The wire diameters of these NWs range from 15 to 35 nm, as indicated by the wire diameter distribution results in figure 4.

Figure 3. TEM image of three-arm CdS nanorod prepared at 160 °C for 7.0 h.

Jang *et al* (2007) prepared 1D CdS by the solvothermal method at different reaction temperatures and durations. They noted that with increased duration, CdS NWs grown at 160 °C increase in length but not in diameter. Moreover, Datta *et al* (2009) studied the reaction temperature (150–250 °C) and duration (5–48 h) effect on the morphology, crystallinity and the optical properties of CdS NWs prepared via solvothermal on Cd foil substrate. They noted that the wires were grown vertically on the substrate and the diameter average was 200–400 nm. The elemental ratios (Cd/S) of all grown CdS NWs range from 1.10 to 1.12, as determined by energy dispersive X-ray measurements.

3.2 Crystalline structure

The crystalline structure of the solvothermally prepared CdS NWs was determined by XRD analysis, and the results are shown in figure 5(A and B). Compared with a standard XRD database (PDF-4, 00-001-0780), the grown CdS NWs exhibit hexagonal wurtzite structures belonging to the C_{6v}^4 CdS space group. The preferred orientation of the grown NWs can be predicted from the following formula (Phuruangrat *et al* 2010):

$$P_{(hkl)} = \frac{I_{(hkl)} \sum I'_{(hkl)}}{I'_{(hkl)} \sum I_{(hkl)}}, \quad (2)$$

where $I_{(hkl)}$ and $I'_{(hkl)}$ are the integrated intensities of the experimental and random (simulated) XRD diffractions, respectively. For a random orientation, P equals one. For a particular (hkl) plane with a preferred orientation, P is greater than one. The calculated P_{hkl} values of the grown CdS NWs prepared at 160 and 200 °C are listed in table 1. The preferred orientation of the NWs prepared at 160 °C for 3.5 h is along the $\langle 001 \rangle$ direction. This preferred orientation changes to the $\langle 100 \rangle$ direction with increased reaction duration to 7 and 24 h. The XRD patterns of the CdS NWs prepared at 200 °C indicate that the samples are more crystalline than those prepared at 160 °C (figure 5(B)). The preferred orientation of the CdS NWs grown at 200 °C also depends on the reaction duration. The preferred orientation of the CdS NWs prepared for 3.5 h is along the $\langle 110 \rangle$ direction. Increasing the reaction duration to 7 and 24 h changes this orientation to the $\langle 100 \rangle$ direction. Dalvand and Mohammadi (2011) studied the effect of preparation parameters on the crystalline structure, morphology and optical properties of solvothermally grown CdS NWs using en as a solvent as well as $\text{Cd}(\text{Ac})_2 \cdot 2\text{H}_2\text{O}$ and S powder as ion sources. They concluded that the preferred orientation of the NWs prepared at 160 °C for 12 h was along $\langle 110 \rangle$. This orientation changes to the $\langle 100 \rangle$ direction with increased reaction temperature to 200 °C. They also noted that the preferred orientation changes from $\langle 110 \rangle$ to $\langle 100 \rangle$ with increased reaction duration from 12 to 24 h. Datta *et al* (2009) noted that the CdS nanowires grown along $\langle 001 \rangle$ direction and the wires became more crystalline when the temperature of the preparation increased from 150 to 250 °C. Surface atoms have higher energies than interior atoms and

Figure 4. Diameter distribution of CdS nanowires (A, B and C) prepared at 160 °C, (D, E and F) at 200 °C for 3.5, 7.0 and 24 h.

thus significantly affect the growth direction. For the hexagonal structure, the atomic density is higher on the (001) surface than on the (100) and (110) surfaces. Therefore, atoms on the (002) face have significantly lower excess energy than atoms on other faces (Phuruangrat *et al* 2009). The preferred orientation in all prepared CdS NWs is along the $\langle 100 \rangle$ or $\langle 110 \rangle$ direction, which makes a 90° angle with $\langle 001 \rangle$. Hence, the growth direction of the grown CdS NWs is along $\langle 001 \rangle$. The (002) peak intensity is higher than those of the (100) and (101) planes of the CdS NWs grown at 160 °C for 3.5 h, which may be due to the lower aspect ratio (Zhou *et al* 2011). Growth along the c -axis of the NW wurtzite structure benefits optoelectronic and electro-optic applications due to the polarized light emission along the c -axis of the wire (Thupakula *et al* 2012). Growth along the $\langle 001 \rangle$ direction

indicates that the grown NWs are bipolar because of the non-centrosymmetric charge distribution in the wurtzite crystal along the c -axis (Thupakula *et al* 2012). The lattice constants a and c for the hexagonal structure are calculated using the following equation (Nien *et al* 2008):

$$\frac{1}{d_{hkl}^2} = \frac{4}{3} \left(\frac{h^2 + hk + k^2}{a^2} \right) + \frac{l^2}{c^2}, \quad (3)$$

where d_{hkl} is the interplanar spacing of the atomic planes; and h , k and l are the Miller indices.

The calculated lattice parameter values of the prepared samples are almost identical to standard data (table 1). The particle size, P_s can be calculated from the XRD peak

Figure 5. XRD patterns of CdS nanowires prepared (A) at 160 °C and (B) 200 °C for various times.

broadening using the Debye–Scherer equation (Thupakula *et al* 2012):

$$P_S = \frac{K \times \lambda}{\beta \cos \theta}, \quad (4)$$

where K is a shape factor, λ the X-ray radiation wavelength (~ 0.15 nm), β the full-width at half-maximum (FWHM) of the XRD diffraction peak and θ the half of the diffraction angle. The CdS NW growth direction is along $\langle 001 \rangle$. Thus, the FWHM of the (002) plane is used to calculate P_S . The diameters (R_w) of the grown CdS NWs are estimated using $P_S = 2R_w$. The estimated R_w values are listed in table 1. The calculated R_w found close to the dominated diameter of the grown CdS NWs that estimated by TEM images. Thus, XRD can be used as a method to estimate the diameter of the grown wires.

3.3 Optical properties

3.3a Optical absorption: Optical studies were performed by measuring the absorption of the prepared CdS NWs

within the 300–900 nm wavelength range. Figure 6 shows the UV–Vis absorption of the CdS NWs prepared at 160 and 200 °C for different durations. The absorption edge of CdS NWs grown at 160 °C for 3.5 and 7 h is observed at around 480 nm; this peak shifted to ~ 485 nm for the NWs prepared for 24 h (figure 6A). Thus, the optical band gap for the NWs prepared for 3.5 and 7.0 h is 2.58 eV, and that for the NWs prepared for 24 h is 2.55 eV. The absorption band peaks for the NWs grown at 200 °C for all durations peak at ~ 485 nm (figure 6B). The obtained optical band gap values are higher than the bulk value (2.42 eV and ~ 515 nm) (Mahdi *et al* 2012b). The sharp absorption peak of the CdS NWs can be attributed to their size uniformity and high crystallinity (Jang *et al* 2007; Yan *et al* 2009). Using the solvothermal method, Yan *et al* (2009) prepared CdS NWs with diameters ranging from 23 to 37 nm and an optical energy gap of 2.56 eV.

3.3b Photoluminescence spectra analysis: The room-temperature PL emission spectra of the CdS NWs prepared at 160 and 200 °C are shown in figure 7. The CdS NWs prepared at 160 °C for 3.5 and 7 h show a sharp PL emission band at around 496 nm (2.5 eV), which correspond to a typical band–band transition. The observed PL band at 496 nm is blue shifted from bulk CdS (500–700 nm). However, the emission bands comes either from band edge or recombination of surface states energy level (Sadhu *et al* 2008). The PL emission of the NWs prepared for 24 h redshifts to 519 nm (2.39 eV), as shown in figure 7(A). A broad yellow emission is observed at 668 nm (1.85 eV) for the NWs prepared for 3.5 and 7 h; this emission peaks at ~ 690 nm (1.8 eV) for the NWs grown for 24 h. Excess Cd in CdS can lead to the formation of an acceptor level within the CdS optical band gap. Thus, the defect band is attributed to the transition of Cd-interstitial donors to the valence band (Tsai *et al* 1996). Ahmad-Bitar (2000) attributed the appearance of the defect emission band to recombination via surface localized states, the transition from I_{Cd} to the valance band, and the transition from the interstitial Cd–Cd vacancy complex that acts as a donor to an acceptor level. The shift in the defect band position may also be related to the effect of the concentration of the defect levels. Interestingly, the CdS NWs prepared at 200 °C for all durations show a sharp PL emission band at 512 nm and another broad emission band at 690 nm. The PL intensities of the CdS NWs prepared for 3.5 h at 160 and 200 °C are the highest among all the samples. Table 2 summarizes the optical properties of the CdS NWs prepared at 160 and 200 °C. However, Datta and his workers (2009) noted that the solvothermally grown CdS NWs showed a distinct blue shift of the PL peak position which centered at ~ 530 nm from low temperature (180 °C) to the higher temperature (250 °C) prepared samples.

3.3c Raman spectra analysis: Figure 8(A and B) shows the room-temperature Raman spectra of all prepared CdS NWs excited by 515 nm Ar^+ laser light. The two observed

Table 1. Crystalline parameters of prepared CdS NWs prepared at 160 °C and 200 °C for 3.5, 7.0 and 24 h.

Preparation temperature (°C)	Time (h)	Orientation face	P_{hkl}	a (Å)	c (Å)	R_w (nm)
160	3.5	(002)	2.51	4.14	6.71	20
=	7.0	(100)	1.55	4.15	6.73	21
=	24	(100)	1.42	4.11	6.71	18
200	3.5	(110)	2.42	4.13	6.72	17
=	7.0	(100)	1.29	4.13	6.72	20
=	24	(100)	1.41	4.13	6.71	23

(A)

(B)

Figure 6. Optical absorption of CdS nanowires prepared (A) at 160 °C and (B) at 200 °C for various times.

peaks in the Raman spectra correspond to the first-order (1LO) and second-order (2LO) longitudinal optical phonon modes of CdS, which are polarized in the x - z face and strongly coupled to the exciton along the c -axis (Arguello *et al* 1969; Wang *et al* 2002; Pan *et al* 2005; Mondal *et al* 2007b). For all the samples prepared at 160 °C, the 1LO peak is located at 298 cm^{-1} . The 2LO peak of the samples prepared for 3.5 and 7 h appear at 600 cm^{-1} and redshifts to 590 cm^{-1} for the sample prepared at 24 h (figure 8(A)). The

(A)

(B)

Figure 7. Photoluminescence spectra of CdS nanowires prepared (A) at 160 °C and (B) at 200 °C for various times.

Raman spectra show that the 1LO and 2LO peaks of all the samples prepared at 200 °C are located at 299 and 600 cm^{-1} , respectively. For bulk CdS, the fundamental 1LO and overtone 2LO modes are located at 305 and 605 cm^{-1} , respectively (Mahdi *et al* 2012b). Thus, the redshift in the positions of the 1LO and 2LO modes in all prepared CdS NWs samples occurs because of the phonon confinement effect (Owens and Poole 2008). The exciton-phonon coupling strength in semiconductors can be determined from

Table 2. Optical results of CdS NWs prepared at 160 °C and 200 °C for 3.5, 7.0 and 24 h.

Preparation temperature (°C)	Preparation time (h)	Absorption peak (nm)	PL peak (nm)	Defect peak (nm)	$I_{\text{Main}}/I_{\text{Defect}}$	$I_{2\text{LO}}/I_{1\text{LO}}$
160	3.5	480	496	668	14.37	0.69
=	7.0	480	496	668	1.77	0.71
=	24	485	519	690	0.38	1.03
200	3.5	485	510	690	2.60	0.70
=	7.0	485	512	690	1.10	1.32
=	24	485	512	690	1.00	1.20

(A)

(B)

Figure 8. Raman spectra of CdS nanowires prepared (A) at 160 °C and (B) at 200 °C for various times.

the intensity ratio of the phonon overtone to the fundamental ($I_{2\text{LO}}/I_{1\text{LO}}$) (Mahdi *et al* 2012d). The CdS NWs prepared at 160 °C for 24 h show higher ($I_{2\text{LO}}/I_{1\text{LO}}$) intensity ratios than the samples prepared for 3.5 and 7 h. On the other hand, the NWs prepared at 200 °C for 7 h have the highest ($I_{2\text{LO}}/I_{1\text{LO}}$) intensity ratio (table 2). NWs with high ($I_{2\text{LO}}/I_{1\text{LO}}$) intensity ratios exhibit strong exciton–LO phonons because of phonon confinement in the transverse directions and the transfer of elementary excitation (carriers, excitons and phonons) to the longitudinal direction (Cao *et al* 2007). These single-crystal

CdS NWs possess high optical properties and are promising building blocks for photonic devices (Yan *et al* 2009).

4. Conclusions

CdS NWs/nanorods were prepared by the solvothermal method at 160 and 200 °C for 3.5, 7 and 24 h and using en as a solvent. The dimensions of the grown CdS NWs are found to depend on the reaction temperature and duration. Analysis of the diameter distribution of the CdS NWs grown at 160 and 200 °C shows that the reaction duration does not significantly affect the wire diameter. A three-armed CdS nanorod structure frequently appears in all the samples prepared at 160 and 200 °C. Appearing three-armed CdS nanorod structure can be attributed to the CdS nanoparticles in the solution, which acts as nuclei for the growth of this structure. The CdS NWs grown at 200 °C for 3.5, 7 and 24 h are longer than those prepared at 160 °C. The crystalline structure of all grown CdS NWs is wurtzite, and the NW/nanorod growth direction is along the $\langle 001 \rangle$ plane. The NW diameters estimated from the XRD results are almost identical to those calculated from the TEM images, thus the XRD method can be used efficiently to calculate the diameter of the NWs. Sharp absorption peaks of the CdS NWs with blueshifts in the absorption edge are observed in all the prepared samples. The optical band gap of the NWs prepared at 160 °C decreases from 2.58 to 2.55 eV with increased reaction duration from 3.5 to 24 h. However, the band gaps of the samples grown at 200 °C are unaffected. The PL emission spectra of the CdS NWs prepared at 160 and 200 °C for 3.5 h show a sharp green emission. The ratios of the band–band emission to the defect intensity for the samples prepared at 200 °C are lower than those for the samples prepared at 160 °C. Raman results show that the optical properties of the nanostructures are enhanced with increased reaction duration for all prepared CdS NWs.

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