Synthesis and Characterization of ZnO Nanostructures by Polymeric Precursor Route

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Zinc oxide nano- and submicrostructures have been synthesized controllably by polymeric precursor method (Pechini). In this approach, zinc acetate $\text{Zn}($CH$_3$COO$\text{)}_2$·2H$_2$O, citric acid and ethylene glycol were used as the source of Zn$^{2+}$, the chelating agent and the connecting agent, respectively. The microstructure of the ZnO nano- and submicrostructures was characterized by X-ray diffractometry and scanning electron microscopy and the optical property was investigated by the room temperature photoluminescence spectra. The effect of ethylene glycol to citric acid mole ratio on the morphology and structure of the products was discussed.

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1. Introduction

Scientists have paid a growing attention to the fabrication of ZnO because of wide range of practical applications in optics, sensors, catalysts, solar cells, data storage, drug delivery and so on [1]. Vapor phase methods involve special equipment complex process control or high temperatures, they are unfavorable for an industrialized process. The chemical methods appear to be of particular interest since they offer the potential of facile scale up, and have flexible processing chemistry. Among chemical methods, sol–gel is an attractive one for preparing nanoparticles because of its easy control, low cost and low fabricating temperature. Pechini’s method is one of the sol–gel methods. In this method, some alpha-hydroxycarboxylic, such as citric acid, forms polybasic acid chelates with metal cations and then the chelates polymerize with a polyhydroxy alcohol and forms polyesters. After calcining the gel, nanometer size powders are achieved [2]. To the best of our knowledge, this is the first report on the synthesis of ZnO nano- and submicrostructures with varying ethylene glycol concentration via Pechini’s method.

2. Experimental

In a typical experiment, 1 g $\text{Zn(AC)}_2$·2H$_2$O and 1 g citric acid monohydrate (CA) were dissolved in deionized water, and appropriated amounts of ethylene glycol (EG) were then added to form a sol at 50°C for 1 h. A white solution was obtained and further heated at 80°C for 1 h to remove excess water. During continued heating at 150°C for 1 h, the solution became more and more viscous and finally became a xerogel. To complete drying, xerogel was placed at 250°C for 1 h. The result powder is a precursor. In the furnace, we heat-treated the precursor at 600°C in air, in a ceramic boat, and then cooled it to reach room temperature.

3. Results and discussion

Figure 1 shows the X-ray diffraction (XRD) patterns of the as-synthesized powders prepared at different mole ratios of EG to CA. As can be seen in Fig. 1a, the relatively low signal/noise ratio and some unknown diffraction peaks indicate that the as-synthesized powder prepared through the use of EG:CA mole ratio 2:1 consisted of insufficiently developed crystalline ZnO and impurities, because the inadequate fuel could not react completely with the zinc nitrate to release enough heat to form the well-developed phase of ZnO. Figure 1b shows the XRD pattern of the product at EG:CA mole ratio 4:1. A search-match data base analysis confirmed that this as-synthesized powder has the phase composition of wurtzite ZnO with $P\overline{6}3m$ structure. The rather stronger (002) peak compared with a standard spectrum in Fig. 1b suggests that the structure of ZnO has a preferential orientation along (002). This unusual diffraction peak of (002) indicates a preferential orientation along the c axis and a rod-like shape. The rod-like shape was confirmed by transmission electron microscopy (TEM) photographs to be discussed later. When the EG:CA mole ratio increases to 7:1, obviously, the crystallinity of the products will improve, as shown in Fig. 1c. Almost all products displayed the XRD peaks corresponding to zincite (JCPDS No. 36-1451).

The effects of the mole ratio EG to CA in the starting solution on the morphology and shape of ZnO powders are shown in Fig. 2. This figure shows the powder scan-
Fig. 1. XRD patterns of ZnO powders obtained with different EG:CA mole ratio (a) 2:1, (b) 4:1 and (c) 7:1.

Fig. 2. SEM images of ZnO powders obtained with different EG:CA mole ratio (a) 2:1, (b) 4:1 and (c) 7:1 (scale bar is 1 µm).

Fig. 3. PL spectra of ZnO: (a) nanoparticles, (b) rod-like and (c) hexagonal submicrorods.
to the overall reaction, the reaction will favor the side opposing the addition of the species. Likewise, the subtraction of a species would cause the reaction to fill the “gap” and favor the side where the species was reduced. This observation is supported by the collision theory. As the concentration of ethylene glycol is increased, the frequency of successful collisions of that reactant would increase also, allowing for an increase in forward reaction, and generation of the product. Even if a desired product is not thermodynamically favored, the end product can be obtained if it is continuously removed from the solution [6].

4. Conclusions

The ZnO nano- and submicrostructures such as rod-like, spherical and hexagonal submicrorods have been fabricated by a simple Pechini approach. Rod-like and spherical ZnO samples show strong near-band edge (NBE) emission generated by the free exciton recombination and very weak deep level emission (DPE) caused by the impurities and structural defects such as oxygen vacancies and interstitials of zinc. A very strong UV emission at ≈ 384 nm and weak green emission at ≈ 510 nm are observed in the PL spectra of one kind of ZnO nanocrystals, i.e. hexagonal ZnO submicrorods, which might indicate the presence of a greater fraction of oxygen vacancies in this sample. It is anticipated that method presented in this work, offers a solution to controlling the morphology of metal oxide crystals.

References