# Synthesis and characterization of ZnSe: Ag/SiO<sub>2</sub> nanoparticles

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**Abstract.** Argentum-doped zinc selenide/silicon dioxide nanoparticles (expressed as ZnSe: Ag/SiO<sub>2</sub>) were synthesized by Stöber method. The structure, morphology and fluorescence properties of the quantum dots were characterized by X-ray powder diffraction, transmission electron microscopy, infrared spectrum, ultraviolet-visible spectrum and fluorescence spectrum. The results show that the as-prepared ZnSe: Ag/SiO<sub>2</sub> nanoparticles are spherical, most of which are about 30 nm in size, and have good fluorescence properties. Compared with that of ZnSe: Ag nanoparticles, the stability of ZnSe: Ag/SiO<sub>2</sub> nanoparticles is enhanced obviously. The ZnSe: Ag/SiO<sub>2</sub> nanoparticles will have potential applications in biological fluorescence analysis.

# **1 INTRODUCTION**

II-VI group quantum dots (QDs), also known as semiconductor nanocrystals, have attracted much attention due to their unique physical and chemical properties, such as wide and continuous absorption spectrum, narrow and symmetrical emission spectrum and photochemical stability.[1-3] Over the years, II-VI group quantum dots have been widely used in biological analysis.[4-6] Zinc selenide (ZnSe) quantum dots are II-VI semiconductor fluorescence materials with low toxicity, wide band gap (2.69eV) and unique fluorescence properties, which are beneficial to their applications in the field of biological analysis[7]. In 2000, Meijerink et al. [8] reported for the first time that Mn doped ZnSe quantum dots synthesized by organometallic synthesis method had excellent fluorescence properties. The fluorescence properties of quantum dots can be improved by doping transition metal ions in the quantum dots.[9] However, in a complex biological environment, the molecules modified on the surface of quantum dots tend to fall off slowly, resulting in a decrease in the stability and fluorescence intensity of quantum dots, which is not conducive to the application of quantum dots in the complex environment.[10] Therefore, in order to improve the stability of the fluorescent quantum dots, it is a good solution to coat the surface of quantum dots with non-toxic, chemically inert and optically transparent silica shell layer. In addition, there are abundant hydroxyl functional groups on the surface of SiO2 shell, which can be modified by silane coupling agent to make the surface of nanoparticles contain amino groups, so as to better realize the application of nanoparticles in biomolecular detection. The method commonly used to prepare SiO2-coated nanoparticles is Stöber method, which has mild and simple synthesis conditions. For instance, the SiO2-coated zinc based nanoparticles (e.g. ZnSe/ZnS/SiO<sub>2</sub>[11], ZnS:

Mn/SiO<sub>2</sub>[12], Mn/ZnS@SiO<sub>2</sub>-NH<sub>2</sub>[13] and ZnS/SiO<sub>2</sub>[14]) were prepared by the Stober method. Up to now, to the best of our knowledge, the synthesis of SiO<sub>2</sub>-coated Ag-doped ZnS nanoparticles (i.e. ZnSe: Ag/SiO<sub>2</sub>) by Sober method has not been reported. In this paper, ZnSe: Ag/SiO<sub>2</sub> nanoparticles were prepared by Stober method by coating SiO<sub>2</sub> on ZnSe: Ag quantum dots reported in our laboratory [15], and the characterization and fluorescence properties of the ZnSe: Ag/SiO<sub>2</sub> nanoparticles were studied.

# 2 Experimental

## 2.1 Instruments and reagents

The X-ray powder diffraction (XRD) patterns were performed by using an X'pert-MPD Model (Philips, Andover, MA). Infrared (IR) spectra were measured with an Equinox 55 FTIR spectrometer (Bruker, Bremen, Germany). The ultraviolet-visible (UV-Vis) spectra were obtained by a Cary 50 UV-visible spectrometer (Varian, Palo Alto, CA). Fluorescence spectra experiments were carried out by using an F-1000 spectrometer photometer (Hitachi, Chiyoda, Japan). Transmission electron microscope experiment were carried out by using a HT7700 Transmission Electron Microscope (Hitachi Corporation, Japan).

All the chemicals including Sodium borohydride (NaBH<sub>4</sub>), Zinc acetate dihydrate (Zn (Ac)<sub>2</sub>·2H<sub>2</sub>O), Selenium (Se), Silver nitrate dihydrate (AgNO<sub>3</sub>·2H<sub>2</sub>O), L-cysteine (L-Cys), Sodium hydroxide (NaOH), tetraethoxy silane (TEOS), ammonium hydroxide (NH<sub>3</sub>·H<sub>2</sub>O), 3-aminopropyl triethoxy silane (APTES) and Anhydrous ethanol (C<sub>2</sub>H<sub>5</sub>OH) were purchased from Guoyao Chemical Reagent. The water used in this experiment is ultrapure water. The preparation of 0.05 mo/ L NaHSe solution was performed according to the procedure from

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the literature [16].

#### 2.2 Synthesis of ZnSe:Ag/SiO<sub>2</sub> nanoparticles

First, under a N2 atmospere, a certain amount of Lcysteine, 0.1 mol/L Zn  $(OAc)_2$  solution and 0.01 mol/L AgNO3 solution were sequentially added into a three-neck flask under magnetic stirring, and the pH value of the mixed solution was adjusted to 10.0 by adding 2.0 mol/L NaOH solution. When the temperature of oil bath was heated to 100°C, a freshly prepared nitrogen-saturated solution of 0.05 mol/L NaHSe was quickly injected into the mixed solution and the reaction solution was subjected to a reflux for 1 hr under vigorous stirring to obtain Lcysteine-modified ZnSe: Ag QDs colloidal solution. The solution was then cooled to 30 °C.

Then, L-cysteine modified ZnSe: Ag quantum dots solution was added to a three-necked flask, the pH value of the solution was adjusted to 10.0 with 5 wt% ammonia water under magnetic stirring for 30 minutes, and a certain volume of tetraethyl orthosilicate solution (TEOS: ethanol =1:3) and a certain volume of 3-aminopropyl triethoxysilane (APTES) solution were added. The above reaction mixture solution reacted in a water bath at 30°C for 12 hours to obtain ZnSe: Ag/SiO<sub>2</sub> nanoparticle colloidal solution. The as-prepared colloidal solution was concentrated by a rotary evaporator and then purified by ethanol precipitation procedure to obtain purified ZnSe: Ag/SiO<sub>2</sub> nanoparticles powder.

# 3 Results and discussion

## 3.1 XRD analysis

The XRD patterns of the as-prepared L-cysteine-modified ZnSe: Ag quantum dots and ZnSe: Ag/SiO<sub>2</sub> nanoparticles are presented in Fig.1. Fig. 1a shows that the three diffraction peaks at  $2\theta$ =28.20°, 46.54° and 54.91° correspond to the (111), (220) and (311) planes of the standard cubic crystalline ZnSe (JCPDS No.80-0021), respectively, which indicates that the as-prepared ZnSe: Ag quantum dots belong to the cubic zinc blende structure. It can be seen from Fig.1 b that there are three diffraction peaks at  $2\theta$ = 27.45°, 46.47° and 54.43° corresponding to ZnSe: Ag quantum dots, and another wide diffraction peak at  $2\theta$ = 23° corresponding to amorphous silicon dioxide (JCPDS No. 29-0085) [17], which indicates that the SiO<sub>2</sub> is coating on the surface of ZnSe: Ag quantum dots.



Figure 1. XRD patterns of L-cysteine-modified ZnSe: Ag quantum dots (a) and ZnSe: Ag/SiO<sub>2</sub> nanoparticles (b)

#### 3.2 TEM analysis

Fig.2 shows the TEM images of  $ZnSe: Ag/SiO_2$  nanoparticles. It can be seen that Ag:  $ZnSe/SiO_2$  nanoparticles have a spherical structure with average particle size of about 30nm.



Figure 2. TEM image of the ZnSe: Ag/SiO2 nanoparticles

#### 3.3 Infrared spectrum analysis

The Infrared spectra of ZnSe: Ag quantum dots and ZnSe: Ag/SiO<sub>2</sub> nanoparticles are shown in Fig. 3 (a), it is can be found from Fig.3a, IR(KBr): 3416.3cm-1(v<sub>O-H</sub>), 2924.4cm-1(v<sub>N-H</sub>), 1587.7cm-1(v<sub>C=O</sub>), 1409.4cm-1(v<sub>C=O</sub>), 1256.7cm-1(v<sub>C-O</sub>), 1060 cm-1(v<sub>C-N</sub>) and 646 cm-1(v<sub>C-S</sub>). From Fig.3 b, it can be seen that the above vibration peaks have shifted slightly, and three new absorption peaks appeared at 1070cm-1(v<sub>O-Si-O</sub>), 921.3cm-1(v<sub>Si-OH</sub>), 463.9cm-1(v<sub>Si-O</sub>), respectively. The three new peaks are all characteristic peaks of silica, indicating that silica is successfully coated on the ZnSe: Ag quantum dots.



Figure 3. IR spectra of L-cysteine-modified ZnSe:Ag quantum dots (a) and ZnSe:Ag/SiO<sub>2</sub> nanoparticles (b)

#### 3.4 UV-Vis absorption spectrum analysis

Fig.4 shows the UV-Vis absorption spectra of the asprepared ZnSe: Ag quantum dots (a) and ZnSe:  $Ag/SiO_2$ nanoparticles (b). As can be seen from Fig.4, compared with the absorption edge of ZnSe: Ag quantum dots, the absorption edge of ZnSe: Ag/SiO<sub>2</sub> nanoparticles show an obvious red shift, which is attributed to the quantum size effect. [18]



Figure 4. UV-Vis absorption spectra of ZnSe: Ag quantum dots (a) and ZnSe: Ag/SiO<sub>2</sub> nanoparticles(b)

#### 3.5 Fluorescence spectrum analysis

Fig.5 presents the fluorescence emission spectra of ZnSe: Ag quantum dots (a) and ZnSe: Ag/SiO<sub>2</sub> nanoparticles (b). Fig.5a shows that ZnSe: Ag has a strong emission peak at 440nm. Compared with the emission peak of ZnSe: Ag quantum dots, the emission peak of ZnSe: Ag/SiO<sub>2</sub> shows a certain degree of red shift, while the fluorescence intensity decreases slightly. The red shift is caused by the formation of SiO<sub>2</sub> shell. The decrease of fluorescence intensity may be due to the effect of SiO<sub>2</sub> shell on the optical refractive index of the fluorescent quantum dot and the shielding effect on partial emitted light.[10]



Figure 5. Fluorescence spectra of ZnSe quantum dots (a) and ZnSe: Ag/SiO<sub>2</sub> nanoparticles (b)

# **4 CONCLUSION**

In this paper, ZnSe:  $Ag/SiO_2$  composite nanoparticles are successfully prepared by Stöber method. X-ray powder diffraction analysis, transmission electron microscopy, infrared spectrum analysis, ultraviolet-visible absorption spectrum analysis and fluorescence spectrum analysis show that SiO<sub>2</sub> is coated on ZnSe: Ag fluorescent quantum dots, and the ZnSe: Ag/SiO<sub>2</sub> spherical composite nanoparticles with an average particle size of 30 nm have good fluorescence properties. The functional groups on the silicon dioxide shell of the composite nanoparticles will provide the possibility for the combination of the nanoparticle with biomacromolecules, which lays the foundation for their application in biological analysis.

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