Synthesis of Mn-doped CdTe nanoparticles and their application as fluorescence sensors

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Abstract:
Mn-doped CdTe nanoparticles (NPs) were synthesized via a novel, facile method at low temperature. The doped NPs were directly synthesized in aqueous solution by mixing CdCl₂·2.5H₂O, fresh NaHTe solution, thioglycolic acid (TGA) and MnCl₂·4H₂O under suitable conditions. Mn-doped CdTe NPs evaluated as fluorescence sensors for bovine serum albumin (BSA) in aqueous solution. Experiment results showed that the fluorescence emission of Mn-doped CdTe NPs was enhanced significantly by BSA, while other substances exhibited no significant effect on NPs. Based on the distinct optical properties of Mn-doped CdTe NPs with BSA, Mn-doped CdTe NPs can be developed as potential identified fluorescence sensors for determination of bovine serum albumin (BSA) in aqueous solution.

Key words: Mn-doped CdTe QDs, Fluorescence, thioglycolic acid, Sensors, bovine serum albumin (BSA)

Introduction
Semiconductor nanoparticles (or quantum dots, QDs), with their unique properties such as good optical characteristics, large surface-to-volume ratio, high electron-transfer efficiency, biocompatibility, dimensional similarities with biological macromolecules and high surface reaction activity, have experienced huge development and a great widening of the fields of applications [1-6]. As a result of tremendous progress made in the understanding of many properties of semiconductors nanoparticles, there have been considerable interests from researchers to go beyond the pure nanoparticles and investigate materials that are intentionally doped with impurities [7–14]. Doped metal nanoparticles (NPs) without doubt have been among the most exhilarating research topics in recent years. Due to quantum confinement of charge carriers in tiny spaces, NPs show some unique and fascinating properties. By ity provides a means of coupling the optical and magnetic properties of these materials via sp-d exchange interaction [15]. Ma et al. [16] have reported the Mn-doped ZnSe QDs for the determination of L-tyrosine in enzyme catalysis system. He et al. [17] have reported the Mn-doped ZnS QDs for the detection of enoxacin in biological fluids. Much effort has been made to realize Mn doping in ZnS, CdS, ZnSe and CdSe NPs in order to produce new materials.

However, Mn-doped CdTe nanoparticles are not fully developed, despite their high potential over a range of applications. Until now, as far as we know, there is no researcher reported on the synthesis of Mn-doped CdTe NPs and their application as fluorescence sensors for determination bovine serum albumin (BSA). A new method for the determination of BSA has been developed based on enhancement of the fluorescence of Mn-doped CdTe quantum dots by BSA in aqueous solution.

Experimental

Synthesis of CdTe NPs
NaHTe solution was prepared using a modified literature procedure by treatment of Te powder with excess NaBH₄ at a molar ratio of 1:20 [18]. Sodium borohydride was added to a small flask with 5mL doubly distilled water for dissolving and connected to the nitrogen to drive off the oxygen dissolving in water. Then, adding Te powder into the oxygen-free water. During the course of the reaction, a small outlet connected to the flask was kept open to discharge the pressure from the resulting H₂. The NaHTe solution was obtained when the color of the solution changed to transparent (Te powder is completely dissolved.)
Synthesis of TGA-capped CdTe nanoparticles

The synthesis of the CdTe NPs was at the molar ratio 1:0.5:2.4 of Cd^{2+}/Te^{2–}/TGA in our experiments. In a three-necked round-bottomed flask, 2 mmol of CdCl_{2}·2.5H_{2}O was dissolved with 200 mL double distilled water, and added 4.8 mmol of TGA under stirring. Then adjusted the pH of the solution to 6 by the addition of 1.0 mol/L NaOH solution dropwise. The resulted solution was drive off dissolved O_{2} in the solution by N_{2} bubbling for 30 min. Under vigorous stirring, oxygen-free NaHTe was injected into the above solution. The reaction between Cd^{2+} and NaHTe took place immediately upon the injection of NaHTe solution. The colorless solution changed to golden yellow.

Synthesis of Mn-doped CdTe nanoparticles

Mn-doped CdTe nanoparticles (NPs) were directly synthesized in aqueous solution by mixing TGA-capped CdTe nanoparticles, 0.01mM MnCl_{2}·4H_{2}O under suitable conditions. After boiling the solution for 8 min, the color of solution changed to orange. Keep the solution at 60°C for 40 min to obtain the Mn-doped CdTe NPs with long fluorescence lifetime.

Results and discussion

The comparison of optical properties

![Fluorescence emission spectra](image)

Fig. 1: Fluorescence emission spectra of (a) TGA–CdTe nanoparticles and (b) Mn-doped TGA–CdTe nanoparticles with excitation wavelength at 351 nm.

The optical properties of Mn-doped TGA–CdTe NPs and undoped NPs were characterized by using fluorescence spectroscopy (Fig. 1). The emission peaks both center at 555 nm, and the spectra are characterized by good symmetry and relatively narrow spectral width. The most striking evidence for the doping is the enhancement in the emission intensity of the Mn-doped CdTe in relation to the pure CdTe nanoparticles, which will be helpful for further research.

Characterization by XRD

The XRD pattern (Fig. 2) was obtained from Mn-doped TGA–CdTe powders precipitated from an aqueous solution of Mn-doped CdTe NPs with excess isopropanol. The XRD spectrum which scans over the 2 theta (θ) range of 10–80° shows diffractive peaks at 25.43° and 42.89°, corresponding to the crystal planes 101 and 103, confirming that it has a hexagonal wurtzite crystalline structure. According to the Debye–Scherrer equation, D=0.89\lambda/\beta \cos \theta, the average size of Mn-doped TGA–CdTe NPs is 23.746 nm.

Characterization by TEM

In order to investigate the morphology and size of the prepared nanoparticles, TEM measurement was carried out (Fig. 3). As shown in Fig. 3, Mn-doped CdTe nanoparticles were nearly monodisperse spherical in shape with an average particle size about 50 nm, which was a little bigger than the size estimated by the Scherrer formula calculations based on XRD pattern.

![TEM images](image)

Fig. 3: TEM images of Mn-doped CdSe.
Influence of pH

The influence of pH on the fluorescence intensity of Mn-doped CdTe system in the absence and presence of BSA was shown in Fig. 4. The low fluorescence intensity in acidic medium is the result of dissociation of the nanoparticles due to protonation of the surface dinding thiolates [19]. With the increase of pH, the deprotonation of the thiol group in the thioglycolic acid molecule occurs. This deprotonation could strengthen the covalent bond between Cd and thioglycolic acid molecule, which bring about the fluorescence intensity enhance with pH increasing. Besides, alkali medium is suitable for both pure Mn-doped CdTe and the presence of BSA. However, the fluorescence intensity begins to decline with the further increase of pH value. Therefore, the exploration of the most suitable pH value is ongoing.

Feasibility of the Mn-doped CdTe QDs for the detection of BSA.

To explore the potential of the Mn-doped CdTe QDs for the detection of BSA, we investigated the effect of the Mn-doped CdTe QDs in the presence of different concentrations of BSA. Spectroscopic measurements were performed in aqueous solution. Fig. 5 shows the enhancement response of the Mn-doped CdTe QDs to BSA in aqueous solution. The addition of BSA enhanced the fluorescence intensity of Mn-doped CdTe QDs.

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References


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