



Design and development of anisotropic inorganic/polystyrene nanocomposites by surface modification of zinc oxide nanoparticles



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ABSTRACT

Anisotropic yolk/shell or Janus inorganic/polystyrene nanocomposites were prepared by combining miniemulsion polymerization and sol–gel reaction. The morphologies of the anisotropic composites were found to be greatly influenced by surface modification of zinc oxide (ZnO) nanoparticle seeds. Two different types of the oleic acid modified ZnO nanoparticles (OA-ZnO) were prepared by post-treatment of commercial ZnO powder and homemade OA-ZnO nanoparticles. The morphologies and properties of the nanocomposites were investigated by transmission electron microscope (TEM), thermogravimetric analysis (TGA), Fourier transform infrared spectroscopy (FT-IR), dynamic light scattering (DLS), and energy dispersive X-ray spectroscopy (EDX). It was found that both post-treated OA-ZnO and in-situ prepared OA-ZnO nanoparticles resulted in the yolk–shell and Janus structure nanocomposites, but with varied size and morphology. These nanocomposites showed stable and strong fluorescence by introducing quantum dots as the co-seeds. The fluorescent anisotropic nanocomposites were decorated separately with surface carboxyl and hydroxyl groups. These composites with unique anisotropic properties will have high potential in biomedical applications, particularly in bio-detection.

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1. Introduction

The concept of “Janus particles”, which named after the two-faced Roman God Janus, was first raised by P. G. de Gennes in his Nobel Prize address in 1991 [1]. “Janus” particles are asymmetric and anisotropic in structure, shape, and surface characteristics with each surface having distinctively different physical and chemical properties [2]. Due to the unique geometrical asymmetry, Janus nanoparticles can be used as ideal drug delivery carriers, catalysis, surfactants, and biosensor [3–5]. Several strategies have been developed to synthesize asymmetrical nanoparticles. These include the layer-by-layer self-assembly [6,7], evaporated metal particles [8–11], Pickering emulsion method [12], biphasic electrified jetting [13,14], photo-polymerization in a microfluidic channel [15,16], and polymer self-assembly [17,18]. Janus particles can assume different geometries: spherical, dumbbell, half raspberry-like, cylindrical, disk, and a variety of other shapes [2]. Besides Janus composites, yolk–shell like nanocomposites was also a newly emerging system for biomedical applications [19,20]. The functional core and hollow shell

of the yolk–shell particles can be tailored to the requirements of the nanoreactors [21] and biomedicine [22]. The common and typical synthesis of the yolk/shell particles includes the following, namely, selective etching or dissolution, bottom-up or soft-template, ship-in-bottle, Ostwald ripening or galvanic replacement process, and Kirkendall diffusion methods [22]. The Janus or yolk–shell nanoparticles are usually composed of inorganic–inorganic dimmer or inorganic–polymer composites [23–26]. Among these, the silica/polystyrene composites is one of the most frequently studied systems [27,28] and the silica/magnetite/polymer composites have been developed into the ternary system [29–31]. The straightforwardness and yield of the reaction have been important factors in the development of the synthetic methods. We have previously reported a one-step facile method based on oleic acid modified iron oxide nanoparticles via combined miniemulsion polymerization and sol–gel reaction for preparing the ternary magnetic composites with Janus or yolk–shell morphologies [32,33]. Such magnetic nanomaterials have demonstrated some advantages in bio-detection of rare target by magnetic collection utilizing the active surface functionality. However, the role of the oleic acid modified inorganic nanoparticles was not well understood in the formation of the ternary composites. Few literatures have been reported about the bilateral system, particularly on the influence of the surface chemistry of Titania hydrosols on the stability and morphology of the hybrid nanoparticles [34].

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In this study, we investigated the effect of surface modification of the oleic acid modified zinc oxide (OA-ZnO) nanoparticles on the composite size and morphologies using a one-pot synthesis process. Two different particle systems were investigated: the OA-ZnO nanoparticles were developed in an in-situ synthesis with simultaneous surface modification, and the other by post-modification of commercialized zinc oxide powder. The oleic acid modified zinc oxide was used as the inorganic seeds. Polystyrene/ZnO@SiO₂ nanocomposites were prepared via a combined process of miniemulsion polymerization and sol-gel reaction. For bio-detection applications, carboxyl group was introduced to the surface of the polymer component for conjugation of antibody. The ternary system without magnetic component was further developed into a fluorescent probe by conjugating with the hydrophobic quantum dots for immune detection of biomarkers via a sandwich method.

2. Materials and experiments

2.1. Materials

All reagents used were analytical grade and available commercially. Zinc oxide (ZnO) was purchased from Emperor Nano Material Co. (Nanjing, China). Oleic acid (OA, 90%) was purchased from Shanghai Chemical Reagents Co. (Shanghai, China). Styrene (St), ammonium hydroxide (NH₄OH, 28 wt %), and Sodium hydroxide (NaOH) were purchased from Sinopharm Chemical Reagent Co. (Shanghai, China). Tetraethoxysilane (TEOS), sodium dodecyl sulfate (SDS) and 4, 4'-Azobis (4-cyanovaleic acid) (ACVA) were purchased from Sigma-Aldrich (USA). Hexadecane (HD) was purchased from Acros Organics (USA). Hydrophobic quantum dots (CdSSe/ZnS core shell nanocrystals with particle size of 9 nm and molecular weight of 7.5×10^5) were purchased from Nanjing technology Co. (Hangzhou, China).

The aforementioned chemicals were used as received. Styrene was washed first with 5 wt % sodium hydroxide solution and stored at -20°C before using. High-purity nitrogen gas, provided by a local supplier, was used for sparging solutions without further purification. Deionized (DI) water was used throughout the entire experiment.

2.2. Synthesis of oleic acid-modified zinc oxide nanoparticles

OA-ZnO-a nanoparticles were fabricated by surface modification of the commercial ZnO powder. With the index of particles' size by Dynamic Laser Scattering (DLS) measurement, the orthogonal experiments were carried out to optimize liquid–solid ratio, mixing time, and the reaction temperature and time for surface modification of the commercial ZnO powder. The optimum modification condition was determined as follows. Briefly, oleic acid (1.94 mL) was dissolved in ethanol (50 mL) with mechanical stirring. After adding ZnO nanoparticles (0.25 g), the dispersion was heated to 60°C which under 200 W sonication (KQ-400KDV, Kunshan Ultrasonic Instruments Co, Ltd.) for 3 h. Finally, the dispersion was cooled down to room temperature.

Another type of OA-ZnO nanoparticles (OA-ZnO-b) was synthesized via a modified process according to Heyon's work [35]. Both types of OA-ZnO nanoparticles were collected by a centrifugal machine (3000 rpm/min, 10 min), and the precipitate was dispersed in 50 mL ethanol and DI water with ultrasonication for 10 min. The washing procedure was repeated for five times and the precipitate was dried by freezing-drying for further use. The extra oleate and oleic acid was removed from both OA-ZnO-a and OA-ZnO-b nanoparticles by washing.

2.3. Synthesis of polystyrene/ZnO@SiO₂ nanocomposites

Anisotropic polystyrene/ZnO@SiO₂ nanocomposites were prepared via a combined process of miniemulsion polymerization and sol-gel reaction reported by our group [32]. The water phase was prepared by dissolving 0.092 g SDS in 39 g of DI water. The oil phase was prepared

by the following steps: HD (0.4 g) was first dissolved in St monomer (7 g) by 3 min sonication, then OA-ZnO nanoparticles powder (0.025 g) was then dissolved in the monomer phase. TEOS (2 g) was subsequently resolved into the mixture of the hydrophobic components. The water phase and oil phase were mixed by dropping the oil dispersion dropwise to the water phase under 200 W sonication (KQ-400KDV, Kun Shan Ultrasonic Instruments Co., Ltd., China), followed by 250 rpm mechanical stirring for 30 min. Miniemulsion droplets system was developed by sonicating the emulsions at 500 W in an ice bath with a duty cycle of 50% for 20 min by a Scientz-IID sonifier (JY92-2D, Ningbo Xinzhi Bio-technology Co., Ltd., China). Upon neutralizing 0.12 g of ACVA by 1 mL of NaOH solution (0.856 M) and added into the flask, the system was deoxygenized by nitrogen bubbling for 0.5 h with 250 rpm mechanical stirring. With a condenser under nitrogen protection, the flask was moved into 70°C water bath to initiate St monomer polymerization. After reaction for 1.5 h, 50 μL of NH₄OH was added into the reaction. After heating for another 5 h, the reaction system was stirred at room temperature overnight. The final polystyrene/ZnO@SiO₂ nanocomposites were obtained by centrifugal separation.

2.4. Synthesis of QDs@ polystyrene/ZnO@SiO₂ nanocomposites

The fluorescent nanocomposites were developed by doping a controlled amount of hydrophobic QDs (5 mg) with the OA-ZnO nanoparticles into the oil phase. The steps were kept the same as synthesis of polystyrene/ZnO@SiO₂ nanocomposites.

2.5. Characterization

Transmission electron microscopy (TEM). The washed composite particles were dispersed in DI water and dried onto carbon-coated copper grids before examination. TEM images and energy dispersive X-ray (EDX) analysis were obtained with a Philips Tecnai 20 electron microscope.

Thermogravimetric analysis (TGA). A thermogravimetric analyzer (Netzsch TG209, Germany) was used to analyze the surface modification of the OA-ZnO. The measurement was performed under a nitrogen atmosphere from ambient temperature to 900°C with a heating rate of $10^\circ\text{C}/\text{min}$.

Fourier transform infrared spectroscopy (FT-IR). FT-IR spectra of unmodified ZnO and the OA-ZnO nanoparticles were obtained by using the infrared spectrometer (Bruker Tensor 27, USA).

Dynamic light scattering (DLS). Hydrodynamic size of the nanocomposites and purified OA-ZnO nanoparticles were measured by the DLS instrument (Zetasizer Nano-ZS, Malvern, U.K.).

Fluorescence spectra. The fluorescence of QDs and composite nanoparticles were measured using a spectrophotometer (LS-55, Perkin-Elmer, USA) at room temperature. Fluorescent composites were dropped on glass slide.

Fluorescent imaging system. After dropping the fluorescent sample onto a glass slide, the fluorescence imaging was performed by using an inverted fluorescence microscope (Nikon ECLIPSE Ti, Japan).

3. Results and discussion

Fig. 1 shows the TEM images and DLS data of OA-ZnO-a and OA-ZnO-b. As shown in Fig. 1A, OA-ZnO-a is of irregular shape with an average size of 25 nm. In Fig. 1B, OA-ZnO-b mainly consists of mono-dispersed spherical particles of a size about 15 nm. Note that in the former (OA-ZnO-a), the oleic acid was functionalized by post-treatment of the commercial ZnO powder and in the latter (OA-ZnO-b), the nanocrystals were surface-functionalized during the in-situ synthesis. With oleic acid coating, DLS data of the two kinds of OA-ZnO nanoparticles show good dispersion in the organic solvent. Upon surface modification, the hydrophobicity of the commercial ZnO particle was dramatically improved. The hydrodynamic average diameter of OA-ZnO-a (Fig. 1C) is

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