Water-dispersible TiO\textsubscript{2} nanoparticles via a biphasic solvothermal reaction method

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Abstract

A biphasic solvothermal reaction method has been used for the synthesis of TiO\textsubscript{2} nanoparticles (NPs). In this method, hydrolysis and nucleation occur at the interface of organic phase (titanium (IV) n-propoxide and stearic acid dissolved in toluene) and water phase (tert-butylamine dissolved in water) resulting in the nucleation of the stearic acid-capped TiO\textsubscript{2} NPs. These NPs are hydrophilic due to hydrophobic stearic acid ligands and could be dispersed in toluene, but not in water. These stearic acid-capped TiO\textsubscript{2} NPs were surface-modified with 2,3-dimercaptosuccinic acid (DMSA) in order to make them water soluble. The resultant TiO\textsubscript{2} NPs were easily redispersed in water without any noticeable aggregation. The Rietveld profile fitting of X-ray diffraction (XRD) pattern of the TiO\textsubscript{2} NPs revealed highly crystalline anatase structure. The average crystallite size of TiO\textsubscript{2} NPs was calculated to be 6.89 nm, which agrees with TEM results. These results have important implications for the use of TiO\textsubscript{2} in biomedical, environmental, and industrial applications.

Keywords: TiO\textsubscript{2}; X-ray diffraction; UV-vis absorption; Fluorescence spectra

Background

TiO\textsubscript{2} nanoparticles (NPs) have been widely investigated in the recent past due to their applications in a wide range of fields including solar cells [1], water photolysis for hydrogen production [2], sensors [3], and antireflective and photochromic devices [4]. TiO\textsubscript{2} has three well-known crystallographic phases in nature: anatase, rutile, and brookite. Among these, anatase has been proved to have excellent chemical and physical properties for environmental remediation [5] and many other uses [6-8]. Numerous methods for the synthesis of TiO\textsubscript{2} NPs have been developed, such as hydrolytic sol-gel process [9], nonhydrolytic sol-gel process [10], hydrothermal methods [11], solvothermal methods [12], and so on. The synthesis of TiO\textsubscript{2} nanoparticles generally involves hydrolysis and condensation of titanium precursors. The titanium precursors are extremely water sensitive; therefore, in conventional aqueous/alcohol-phase/sol-gel method in conventional solution-phase synthetic routes, small amount of water is used to inhibit the hydrolysis. However, prepared TiO\textsubscript{2} NPs suffer from poor crystallinity and inferior material properties as compared to those prepared through high-temperature, nonhydrolytic methods. Furthermore, these methods of synthesis suffer from problems of aggregation, size, and low monodispersity and post-treatment procedures (for converting amorphous phase to crystalline TiO\textsubscript{2} phase) which greatly affect the desired properties of the nanoparticles and restrict their large-scale production and applicability. The properties of TiO\textsubscript{2} are highly dependent on surface area, crystalline phase, and single crystallinity. The high-quality TiO\textsubscript{2} NPs prepared through nonhydrolytic methods are insoluble in aqueous medium, which make their utilization toward biological/biomedical applications impossible. At present, the synthesis methods for production of water-dispersible TiO\textsubscript{2} NPs with a tunable size is challenging to the researchers.

In this letter, we present the preparation of water-soluble and biocompatible highly crystalline TiO\textsubscript{2} NPs through biphasic solvothermal interface reaction method.

Methods

The following chemicals were used as purchased: titanium (IV) n-propoxide, tert-butylamine, 2,3-dimercaptosuccinic acid (DMSA) and stearic acid (SA) (Sigma-Aldrich, Steinheim, Germany) and toluene (Penta, Chrudim, Czech Republic). All the chemicals were of analytical grade.
purity. Deionized water (Millipore) was used to prepare aqueous solutions (≥18 MΩ). In biphasic solvothermal reaction method, the reaction occurs at the interface of water phase and organic phase at elevated temperature. In the synthesis procedure, the organic phase consists of 90 μL of titanium (IV) n-propoxide and 0.5 g of SA dissolved in 10 mL of toluene. The water phase contains 100 μL of tert-butylamine dissolved in 10 mL of deionized (DI) water. First, water phase was added to a Teflon-lined steel autoclave. Then, the organic phase was added slowly into the Teflon-lined steel autoclave without any stirring. The autoclave was sealed and heated to 170°C for 6 h. The reaction mixture was then cooled to room temperature, and methanol was added to precipitate the TiO2 NPs. TiO2 NP precipitates were recovered by centrifugation and washed several times with methanol to remove the excess of surfactant. This resulted in hydrophobic SA-coated TiO2 NPs, which are dispersible in toluene. The water dispersibility of TiO2 NPs was achieved by treating the SA-coated TiO2 NPs in a solution of ethanol and toluene containing 2,3-DMSA for 24 h with vigorous stirring. This resulted in DMSA-coated TiO2 NPs which were recovered via centrifugation. Then, the final NPs were easily dispersed in water.

The crystal structure and morphology of as-synthesized nanoparticles were investigated with X-ray diffraction (XRD) using monochromatic Cu Kα radiation (λ = 1.5418 Å) and transmission electron microscope (TEM). The crystalline nature of the NPs was then examined by TEM measurements. The optical properties were investigated by UV-visible (UV-vis) absorption and fluorescence spectra at room temperature.

Results and discussion

During heating, hydrolysis and nucleation of the titanium (IV) n-propoxide occur at the interface of organic phase and water phase resulting in simultaneous nucleation of TiO2 NPs. The XRD pattern of TiO2 NP sample prepared at 170°C was analyzed with Rietveld profile fitting method using FullProf program [13] within anatase 141/amd space group. The Rietveld profile fitting of XRD pattern of prepared TiO2 NPs illustrated in Figure 1 shows the good quality of the fit. No impurity phases were found in the XRD patterns of TiO2 NP samples. The diffraction peaks were indexed with powder diffraction standard data (ICDD 21-1272). The crystallite size of TiO2 NPs is estimated from broadening of anatase (101) peak using the Debye-Scherrer formula [14]. The calculated crystallite size for TiO2 nanoparticles prepared at 170°C is 6.89 nm. The nanoparticles were also prepared at lower temperatures (140°C, 150°C, and 160°C) and higher temperatures (180°C and 190°C). NPs prepared at lower temperatures have smaller crystallite size but the product yield is low, while NPs prepared at higher temperatures have higher yield but the crystallite size is bigger. The optimum temperature is 170°C for the preparation of TiO2 NPs with narrow size distribution and nearly 100% yield.

The morphology of SA-coated and DMSA-coated TiO2 NPs was examined by TEM measurements. As shown in Figure 2a,b, the resulting TiO2 NPs (SA-coated and DMSA-coated) appear as spherical particles with good monodispersity. The size distribution of the nanoparticles is in Additional file 1: Figure S2, calculated by measuring hundred particles, shows that the TiO2 NPs have an average size of 6 nm, which is in good accordance with the size of TiO2 NPs observed through XRD measurement. The inset of Figure 2a,b presents the SAED pattern of TiO2 NPs, confirming that anatase crystal structure can be indexed with (101), (103), (200), (105), (213), (116), (107), and (008) crystallographic planes.

UV-vis absorption spectra of TiO2 nanoparticles dispersed in toluene and DI water are measured and shown in Figure 3. The absorption coefficient (α) was determined from the optical spectrum using the formula

\[ \alpha = \frac{2.3026A}{t} \]

where A and t are the measured absorbance and thickness of the sample, respectively. The optical bandgap energy (Eg) was evaluated from the absorption spectrum, and the optical absorption coefficient (α) near the absorption edge is given by following equation:

\[ ahv = B(hv-E_g)^n \]

where h, ν, B, and E_g are Plank’s constant, frequency of incident photons, constant, and optical bandgap energy,
respectively. $E_g$ was estimated by plotting $h\nu$ versus $(\alpha h\nu)^{1/2}$ and extrapolating linear portion near the onset of absorption edge to the energy axis as shown in the inset of Figure 3. The determined value of $E_g$ is 3.06 and 3.1 eV for TiO$_2$ nanoparticles dispersed in toluene and DI water, respectively.

Fluorescence spectra of SA-coated TiO$_2$ NPs in toluene and DMSA-coated TiO$_2$ NPs in DI water with an excitation wavelength of 325 nm were recorded at room temperature and are shown in Figure 3a,b. The broad emission spectra which are observed from 400 to 500 nm arise from indirect bandgap and surface recombination processes [15]. After multipeak Gaussian fitting of fluorescence spectra in Figure 3a,b, we found that Gaussian curves fit original curves perfectly. The peak positions of Gaussian bands in Figure 4a are located at about 384, 407, 440, 480, and 525 nm, respectively. The peak positions of Gaussian bands in Figure 4b are located at about 394, 418, 445, 485, and 540 nm,
respectively. All these peaks are red shifted due to the light-induced relaxation of polar molecules [16]. The prepared TiO2 NPs with high surface-to-volume ratio favor the existence of large quantities of oxygen vacancies. The observed fluorescence bands may be the result of emission from radiative recombination of self-trapped excitons localized within TiO6 octahedra and oxygen vacancies [17]. Oxygen vacancies have been considered as the most common defects and usually act as radiative centers in the luminescence processes [18]. The emission peak at about 384/394 nm is attributed to the emission of near bandgap transition of anatase. This is consistent with the Eg calculated by UV measurement technique (i.e., approximately 3.1 eV). The emission bands at 407 and 418 nm were ascribed to electron transition mediated by defect levels in the bandgap [19]. In addition, the signals observed in wavelength range from 440 to 540 nm arise from the excitonic PL, which mainly results from surface oxygen vacancies and defects. The peaks at 440 and 445 nm are attributed to band edge free excitons, and the other peaks at 480 and 485 nm correspond to bound excitons [20].

Conclusions

A facile route for the synthesis of TiO2 NPs through biphasic solvothermal interface reaction method has been reported. The XRD pattern of TiO2 NPs revealed the anatase structure. The average XRD crystallite size was calculated as 6.89 nm using the Scherrer formula. The optical studies showed that the bandgap is 3.1 eV. The results show that synthesized nanoparticles are monodispersed with long-term stability. This synthesis method is simple with very high production yield and does not require any post-treatment procedure, and products can be collected from organic phase which effectively avoids TiO2 grain aggregation.

Additional file

Additional file 1: Synthesis, size distribution, XRD patterns, and FTIR spectra of TiO2 nanoparticles. Figure S1: Schematic of TiO2 nanoparticles synthesis via a biphasic solvothermal interface reaction method. Figure S2: The size distribution of the nanoparticles. Figure S3: The XRD patterns of the TiO2 nanoparticles prepared at different temperatures. Figure S4: FTIR spectra of the SA-capped TiO2 nanoparticles.

Competing interests

The authors declare that they have no competing interests.

Authors’ contributions

RM carried out the synthesis and characterization. JD improved the manuscript and participated in the studies. JH directed and coordinated the present study as principal investigator. All authors read and approved the final manuscript.

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