Fabrication, characterization, and visible-light photocatalytic performance of ternary plasmonic composites

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HIGHLIGHTS

- W-Ti-SBA15 composites with low TiO\textsubscript{2}/SiO\textsubscript{2} mass ratios and their Ag-contained analogs were fabricated.
- These composites showed enhanced photocatalytic performance in comparison to those with high TiO\textsubscript{2}/SiO\textsubscript{2} mass ratios.
- Photocatalysis mechanism of these Ag-W-Ti-SBA15 plasmonic composites was proposed.

ABSTRACT

A group of mesoporous WO\textsubscript{3}-TiO\textsubscript{2}-contained SBA15 composites (W-Ti-SBA15) and their Ag-contained plasmonic analogs were fabricated and then systematically characterized by a collection of analytical techniques. Original mesoporous structures with highly ordered channels were well-maintained and WO\textsubscript{3}-TiO\textsubscript{2} could form spherical clusters that were gradually reduced in size accompanying with the decrease of TiO\textsubscript{2}/SiO\textsubscript{2} mass ratios. Deposited Ag species were confirmed as zero-valence Ag by X-ray photoelectron spectroscopy analysis and exerted almost no effect on the morphology, microstructure, and textural but optical property, by which the visible-light harvesting ability of composites was remarkably strengthened owing to the localized surface plasmonic resonance (LSPR) induced by the metallic Ag. Under visible-light irradiation, W-Ti-SBA15 composites showed satisfactory photocatalytic performance that could be further enhanced by using their Ag-contained ternary plasmonic analogs. The enhancement of photocatalytic efficiency was mainly attributed to the favorable mesoporous morphology, the suitable band alignment, and the LSPR effect. Eventually, a possible photocatalysis mechanism was primarily proposed on the base of reactive radical species trapping experiments.

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1. Introduction

It is well recognized that anatase (TiO\textsubscript{2}) is an excellent heterogeneous photocatalyst to mediate environmental contaminants
removal, evolution of gaseous hydrogen, and selective organic conversion in ultraviolet region, owning to the favorable physicochemical properties [1–3]. For practical applications, the TiO2 should be modified in terms of microstructure and morphology to pursue the extension of light absorption range, the enhancement of photocatalytic performance, and the ease of recovery after utilization [4]. Among several modification strategies that currently intensively researched, the construction of a composite containing TiO2 and another semiconductor component with an appropriate band structure is an effective manner to strengthen the visible-light harvesting and photocatalytic capability as a result of the inhibited recombination of photoinduced charge carriers [5].

Tungsten oxide (WO3) with a relatively narrow band gap (−2.4 eV) has been extensively applied in various fields such as electrochromic devices, dye-sensitized solar cells, gas sensors, and photocatalytic processes [6,7]. It is usually adopted as a suitable candidate to fabricate composites together with TiO2, attributing to the well-matched band gap structure [8–12]. The generated charge carries are able to migrate across heterojunction interfaces towards different components and the recombination of charge carries is thus effectively retarded, hereby improving photocatalytic efficiency [13–15]. In addition, the integration of WO3 into composites induces the enhancement of surface acidity, by which some species with unpaired electrons like O2−, H2O, and OH− are prone to access active sites and convert to oxidative O2− and*OH radicals, thus boosting photocatalytic performance [16–18]. If the produced composite incorporates into an appropriate matrix like SBA15, much more active sites can be exposed originating from the decreased particle size and a large amount of pollutant molecules tend to gather around active sites as a result of the strengthened adsorption ability from support, further benefiting photocatalytic processes [19–22].

The deposition of noble metal nanoparticles on surface of semiconductors is another efficient manner to promote photocatalytic efficiency through the formation of schottky junction at the metal-semiconductor interfaces and the LSPR effect by metallic noble elements [23–25]. The former is able to induce the directional migration of electrons and hereby causes the separation of charge carriers [26,27]. The later can enhance the visible-light adsorption and boost the generation of charge carriers. Among mostly used noble metals such as Ag, Pt, Pb, Au, Ag is quite promising due to the low toxicity, antibacterial activity, easy preparation and controllable morphology [28].

In a previous study, the synthesis and photocatalytic assessment of mesoporous W-Ti-SBA15 composites were investigated, and these heterojunction-structured composites showed enhanced photocatalytic efficiency in comparison to single-component samples [29]. However, the photocatalytic performance was not as good as we had expected, mainly attributing to the serious aggregation of TiO2 particles since the excess amount of titanium precursor was added during synthesis. In order to achieve the further advancement of photocatalytic performance, in this study we tried to reduce the mass ratio of TiO2/SiO2 for the sake of the maximal exposure of active sites to contaminant molecules. In addition, silver nanoparticles were introduced on the surface of above composites to construct a ternary Ag-W-Ti-SBA15 plasmonic system. As far as we know, there is no relevant investigation regarding the construction and photocatalytic evaluation of ternary Ag-W-Ti-SBA15 plasmonic composites.

In the current study, a series of W-Ti-SBA15 composites and their Ag-decorated counterparts were fabricated and were fully characterized. The original mesoporous structure was well conserved even after decorating metallic Ag species. W-Ti-SBA15 composites showed satisfactory photocatalytic performance toward the degradation of dye rhodamine B (Rhb) under visible-light irradiation. As expected, Ag-contained Ag-W-Ti-SBA15 plasmonic composites exhibited relatively high photocatalytic efficiency comparing with their W-Ti-SBA15 analogs. The enhancement of catalytic outcome was discussed and a possible photocatalysis mechanism was proposed basing upon active species trapping experiments.

2. Materials and methods

2.1. Materials

Pluronic P123 (PEO20PPO70PEO20) and sodium tungstate (Na2WO4·2H2O) were purchased from Aldrich Chemical Co. and ACROS Organics, respectively. Hydrochloric acid (HCl, 36–38%, AR), silver nitrate (AgNO3, AR), titanium sulfate (Ti(SO4)2·9H2O, CP), tetraethyl orthosilicate (TEOS, Si(OCH3)4, AR), nitroblue tetrazolium (NBT, AR), and other chemicals involved were obtained from Sinopharm Chemical Reagent Co., Ltd. (Shanghai, China) and used directly as received without further purification. Rhodamine B (Rhb, AR) was purchased from Shanghai SSS Reagent Co., Ltd. (Shanghai, China). Deionized water was used throughout the experiment.

2.2. Synthesis of W-Ti-SBA15 composites

The synthetic procedure was similar to the previous study except that the amount of titanium and tungsten precursors was greatly reduced [29]. A series of W-Ti-SBA15 mesoporous composites were prepared and nominated as 5%WTSx, where 5% and x were defined as theoretical mass ratios of formed WO3/TiO2 and TiO2/SiO2, respectively. Sample WO3/TiO2 with a mass ratio of 5% was fabricated according to a same procedure as above except the absence of silicon source and denoted as 5%WT. In addition, N-doped TiO2 (N-TiO2) was synthesized in the light of a previous report for comparison [30].

2.3. Synthesis of Ag-W-Ti-SBA15 plasmonic composites

Since the as-synthesized sample 5%WT50.1 showed the best photocatalytic behavior and was specifically selected to prepare Ag-W-Ti-SBA15 plasmonic composites. In a typical procedure, the sample 5%WT50.1 (1 g) in deionized water (40 mL) was ultrasonicated for 30 min to make a suspension that was subsequently added with a desired amount of AgNO3. After a vigorous stir of 900 r/min (JB-2A constant temperature agitator, Shanghai INESA Scientific Instrument Co., Ltd, China) at room temperature for 5 h in dark, methanol (20 mL) was added into the reaction system that was then exposed to a 400 W halogen lamp (Institute for Electric Light Source, Beijing) equipped with a NaNO2 solution (2 M) to remove ultraviolet-light (<400 nm) for 20 min. The resultant precipitate was collected by centrifugation, washed with ethanol and deionized water for at least five times to completely remove silver cations, and dried at 60 °C for 12 h to provide final products. For simplicity, these composites were denoted as xAgWTS, and x herein was referred to a theoretical mass ratio of Ag versus WTS.

2.4. Characterizations

Powder X-ray diffraction (XRD) patterns were recorded on a Bruker D8 Advance X-ray diffractometer (Bruker AXS, Germany) using a Cu Kα radiation source (λ = 1.5406 Å). The X-ray photoelectron spectroscopy (XPS) measurements were conducted on a Thermo Scientific ESCALAB 250XI system. Binding energies were calibrated with the containment carbon (C1s = 284.6 eV). The morphology and microstructure of obtained samples were recorded on transmission electron microscopy (TEM, JEOL, JEM-2011) and the chemical composition of samples was determined by X-ray energy dispersion spectroscopy (EDS). The specific surface areas were mea-
sured by nitrogen adsorption-desorption isotherms at 77 K using a Micromeritics 3Flex apparatus. Before measurement, the samples were degassed at 423 K under vacuum over 6 h. UV–vis diffuse reflectance spectra (UV–vis DRS) were performed on a Shimadzu UV–2600 spectrophotometer using BaSO₄ as a reference.

2.5. Photocatalytic activity assessment

Photocatalytic evaluation experiments were conducted through the degradation of dye RhB under visible-light irradiation using a 300 W xenon lamp (CEL-HFX300, AuLigh, Beijing) with a 420–780 nm cut-off filter. The photocatalyst (40 mg) was dispersed into an RhB aqueous solution (15 mg L⁻¹, 80 ml). Prior to the light irradiation, the suspension was vigorously stirred (900 r/min, JB–2A constant temperature agitator, Shanghai INESA Scientific Instrument Co., Ltd., China) for 1 h to reach an adsorption saturation of organic molecules onto the surface of catalyst particles. After degradation reactions occurred, 4 mL aliquot was sampled, diluted, and centrifuged at 10000 rpm for 5 min (TGL–20 B centrifugal machine, Shanghai Anting Scientific Instrument Factory, China) to remove catalyst particles at every 30 min. The residue concentration of RhB was evaluated through a UV–vis spectrophotometer (Purkinje General T6) at 544 nm.

Several active radical species possibly generated during photocatalytic reactions were detected in order to propose a mechanism. Specifically, 1.0 mM isopropanol alcohol (IPA) or disodium ethylenediaminetetraacetate (EDTA–2Na) was introduced to check the presence of hydroxyl radicals (•OH) or produced holes (h⁺) [31,32]. In addition, 25 μM nitroblue tetrazolium (NBT) was added into reaction system in place of RhB to analyze superoxide radicals (•O₂⁻) that might probably produce [33].

Each reported value as above was the average of at least three parallel tests and the corresponding standard deviation was calculated within ±3%.

3. Results and discussion

Low-angle and wide-angle XRD patterns of obtained 5%WTSx composites with variable TiO₂/SiO₂ mass ratios are represented in Fig. 1A and B, respectively. All samples in Fig. 1A exhibit three well-resolved peaks in low-angle X-ray diagrams, assigning to characteristic reflections of (100), (110), and (200) crystal planes that originate from mesoporous silicon-based materials with p6mm hexagonal symmetry [34]. It is obvious that the involvement of external species has almost no effect on the SBA15 original structure. In Fig. 1B, a series of peaks at approximately 25.6°, 38.2°, 48.4°, 54.4°, 55.4°, 63.1° assigned to the extra-framework tetragonal anatase TiO₂ phase [JCPDS No.21-1272] [10,16] are clearly observed in all samples, even the one with quite low TiO₂/SiO₂ mass ratio of 0.025, possibly attributing to the much higher condensation rate and thus the easier generation of titania clusters in aqueous environment than silica [35]. The intensity of peaks is gradually reduced with the abatement of TiO₂/SiO₂ mass ratios, suggesting the stepwise decrease of the amount and crystallinity of anatase. A broad band at around 23° is attributed to amorphous or microcrystalline phase of silica [36]. However, there are no diffraction reflections corresponding to WO₃ phase and solid solution of WₓTi₁₋ₓO₂ [37], revealing the uniform distribution of WO₃ onto TiO₂ matrix and the inexistence of other impurities [9]. The integration of metallic Ag species into WTS samples exerts almost no effect on the original mesoporous structures since three feature diffraction signals are prominent in Fig. 1C. In addition, quite similar diffraction patterns of Ag-contained ternary composites to those WTS series are observed in Fig. 1D and the absence of metallic Ag signals is mainly due to the uniform dispersion or the overlap of (111) crystal plane in metallic Ag phase at 38.1° by the (004) crystal plane in anatase TiO₂ at 38.2° [38].

The morphology and microstructures of as-prepared composites were observed by TEM images and are depicted in Fig. 2. All WTS samples with low TiO₂/SiO₂ mass ratios possess extremely ordered and uniform mesoporous microstructures. As seen in Fig. 2A–D, both the quantity and size of the black spherical clusters anatase TiO₂ are gradually decreased along with the decrease of Ti species concentration, which is in good accordance with the XRD analyses. The EDS spectrum of the sample 5%WTS0.1 in Fig. A1 shows four elemental species W, Ti, Si, and O and mass ratios of WO₃/TiO₂ and TiO₂/SiO₂ are estimated as 0.042 and 0.094 that are quite close to theoretical values. After the decoration of metallic Ag, the original mesoporous structure with regular parallel channels is well maintained in sample 5%AgWTS, as shown in Fig. 2E. Two distinct lattice fringes of 0.350 nm and 0.242 nm are observed in HRTEM image and correspond to crystallographic planes of (101) and (111) in anatase TiO₂ phase and metallic Ag in Fig. 2F. However, no obvious lattice fringes assigning to WO₃ are found in such circumstance.

The N₂ adsorption-desorption isotherms were checked to investigate the textural property of as-synthesized composites, as represented in Fig. 3. It is observed that all composites display the identical type IV isotherms with a typical H1 hysteresis loop in a relative pressure P/P₀ range from 0.5 to 0.9, suggesting of regular mesoporous microstructures with hexagonal cylindrical pores at capillary condensation step [39], which is in good agreement with the analytic results from XRD patterns and TEM images. Specific surface area (S₅₄⁰) and pore volume values of these mesoporous composites are collected in Table 1. As expected for WTS series, with the reduction of TiO₂/SiO₂ mass ratio, specific surface area and pore volume values are gradually increased, revealing that Ti and W alien species mainly exist as large clusters on the external surface of SBA-15 support and the higher TiO₂/SiO₂ mass ratio induces the less ordered mesoporous materials with the lower specific surface area [40]. As to composites AgWTS with the increase of Ag precursor addition, specific surface area values are slightly decreased while the average pore diameters keeps almost in variant, indicating that the presence of small amount of Ag exerts an inconspicuous effect on the textural property.

The optical property of obtained composites was analyzed by UV–vis diffuse reflectance spectra and shown in Fig. 4. It is evident that the photoresponse of WTS series is merely in UV–light region and the adsorption edge is close to 400 nm, corresponding to the O²⁻–Ti⁴⁺ charge transition of extra-framework anatase TiO₂. The intensity of the band at about 330 nm increases along with the raise of anatase content, owning to the low structural integrity caused by the comparably high concentration of alien species [41]. A weak band at about 210–230 nm in all samples except 5%WTS0.2 is relevant to the isolated tetrahedral Ti⁴⁺ cations in framework [41], suggesting the relatively high TiO₂/SiO₂ mass ratio tends to promote the formation of extra-framework anatase TiO₂. In addition, with the abatement of TiO₂/SiO₂ mass ratios, the adsorption edge of WTS series is gradually blue-shifted in Fig. 4A, mainly relating to the size shrinkage of anatase TiO₂ clusters in SBA-15 matrix. As soon as metallic Ag was deposited, the photoresponse range of samples is able to extend to the visible-light region, ascribing to the

<table>
<thead>
<tr>
<th>Composites</th>
<th>S₅₄⁰ (m² g⁻¹)</th>
<th>Pore volume (cm³ g⁻¹)</th>
<th>Band gap (eV)</th>
</tr>
</thead>
<tbody>
<tr>
<td>5%WTS0.2</td>
<td>590.9</td>
<td>0.666</td>
<td>2.95</td>
</tr>
<tr>
<td>5%WTS1</td>
<td>719.7</td>
<td>0.694</td>
<td>2.97</td>
</tr>
<tr>
<td>5%WTS0.5</td>
<td>899.4</td>
<td>0.852</td>
<td>2.98</td>
</tr>
<tr>
<td>5%WTS0.025</td>
<td>838.1</td>
<td>0.711</td>
<td>3.12</td>
</tr>
<tr>
<td>5%AgWTS</td>
<td>743.3</td>
<td>0.726</td>
<td>3.24</td>
</tr>
<tr>
<td>10%AgWTS</td>
<td>712.5</td>
<td>0.714</td>
<td>3.26</td>
</tr>
<tr>
<td>15%AgWTS</td>
<td>695.4</td>
<td>0.663</td>
<td>3.26</td>
</tr>
<tr>
<td>20%AgWTS</td>
<td>682.7</td>
<td>0.739</td>
<td>3.28</td>
</tr>
</tbody>
</table>
LSPR effect of plasmonic metallic Ag species [42]. Band gap energies ($E_g$) of these semiconductors were estimated via an empirical equation $\alpha hv = A(hv - E_g)^n$ and $\alpha$, $v$, $E_g$ and $A$ refers to the absorption coefficient, light frequency, band gap energy, and a constant, respectively. It is generally realized that anatase TiO$_2$ undergoes an indirect transition and correspondingly $n$ is set as 4 [31]. As listed in Table 1, $E_g$ values increase with the decrease of TiO$_2$/SiO$_2$ mass ratio in WTS series and the increase of Ag deposition in AgWTS series.

To further confirm the presence of Ag species and check the surface valence states, sample 5%AgWTS was selected for XPS analysis with C1s at 284.6 eV as a reference. Clearly, the survey spectrum provides five elements Ti, O, Si, Ag, and W in Fig. 5A, confirming the indeed presence of ternary components as anticipated. There are two peaks with binding energies located at 373.63 eV and 367.63 eV in high resolution spectrum of the Ag 3d spectrum in Fig. 5B, assigning to Ag 3d3/2 and Ag 3d5/2, respectively. The existence of both Ag peaks with the description of 6.0 eV verifies the formation of zero-valence Ag species instead of Ag ions in composites [43]. Fig. 5C displays two prominent signals at around 458.5 eV and 464.2 eV that are indexed to the Ti 2p$_{3/2}$-Ti 2p$_{1/2}$ spin-orbit components of Ti$^{4+}$ species in anatase TiO$_2$ lattice [8,15]. W species can be found in Fig. 5D and the band was deconvoluted to two peaks centered at 35.8 eV and 37.9 eV in binding energy owning to the spin-orbit splitting of W (4f) components [44]. In addition, the former peak at 35.8 eV is generally considered as an integration signal composed of the Ti$^{4+}$ species (Ti 3p) in TiO$_2$ phase and W(4f) ingredients [15].

Photocatalytic performance of obtained composites was estimated by the decomposition of dye RhB under visible-light illumination and repeated at least twice. As depicted in Fig. 6A, the direct photolysis of RhB can be ignored even after 180 min, revealing the sufficient stability of RhB upon such condition. The standard sample N-TiO$_2$ is able to supply a quite weak photocatalytic removal efficiency of around 23%. Without the mesoporous silicate matrix, the sample 5%WTS can catalytically destroy around 39% of RhB molecules that is much lower than those from WTS series with variable TiO$_2$/SiO$_2$ mass ratios. Particularly, under the identical condition the best candidate 5% WTS0.1 has an optimal photocatalytic degradation efficiency of 63% that is significantly better than that of 5%WTS0.7, a sample with a large TiO$_2$/SiO$_2$ mass ratio of 0.7. The enhancement of photocatalytic behavior of composites with low TiO$_2$/SiO$_2$ mass ratios is mainly attributed to the enlarged specific surface areas of composites and good exposure of active sites induced by the decreased size of semiconductor clusters, which facilitates the adsorption of organic molecules near to SBA-15 matrix and further easy access to active sites to undergo degradation reactions. As a result, the sample 5% WTS0.1 was typically selected to construct Ag-contained ternary plasmonic AgWTS composites. Under the same condition, all Ag-contained composites exhibit relatively high photocatalytic performance comparing
to samples without the Ag decoration, as seen in Fig. 6B. The composite with an Ag mass ratio of 5% shows the best photocatalytic outcome among several Ag-deposited systems, indicating that 5% is a suitable mass ratio and excess amount of Ag deposition might role as recombination center of charge carriers, hereby inhibiting the photocatalytic process [45].

In order to discriminate the active radicals during degradation and further propose a photocatalysis mechanism, active radicals trapping experiments were carried out over the sample 5%AgWTS by means of the introduction of suitable scavenger such as IPA and EDTA-2Na for hydroxyl radicals *OH and holes, respectively. Remarkably, the addition of EDTA-2Na has no effect on the degrada-
tion efficiency in Fig. 7A. However, the photocatalytic performance is greatly retarded as soon as IPA was added, indicating radicals •OH instead of holes exert dominant role in catalytic decomposition. In addition, the remarkable decrease of NBT concentration is indicative of the generation of superoxide radicals \( \cdot \text{O}_2^- \). On the basis of above results, radicals •\text{O}_2^- and •OH are predominate species that greatly promote the photocatalytic efficiency. The standard deviation bar of each value was added in Figs. A2–A4.

According to active radicals entrapping experiments, a possible mechanism was eventually proposed as shown in Fig. 8. Under visible-light irradiation (\( \lambda > 420 \text{ nm} \)), the component WO\(_3\) tends to excite and promote electrons to transfer to the conduction band,
leaving holes on the valence band. TiO$_2$ is unable to excite because of the wide band gap. The photogenerated electrons on WO$_3$ conduction band are insufficient to reduce dissolved oxygen to produce radicals *O$_2^{-}$ instead, they can migrate and transfer to metallic Ag sites via the Schottky barrier at interface between metal and semiconductor phase. In addition, the holes collected on WO$_3$ valence band can thermodynamically transfer to valence band of TiO$_2$ and further oxidatively convert OH$^-$ or water to radicals •OH since the potential of valence band maximum $E_{VBM}$ ($+2.95$ eV) is more positive than $E$(•OH/OH$^-$) ($+1.99$ eV) and $E$(•OH/H$_2$O) ($+2.68$ eV) [46,47]. In addition, metallic Ag tends to excite under visible light irradiation as a result of LSPR effect and the produced hot electrons can inject to the conduction band of TiO$_2$ and thus react with dissolved oxygen to form radicals *O$_2^{-}$ since the potential of conduction band minimum $E_{CBM}$ ($-0.29$ eV) is much negative than $E$(•O$_2^{-}$/O$_2$) ($-0.046$ eV) [47]. The formation of radicals *O$_2^{-}$ that was unavailable in WTS composites [32] can be successfully achieved in metallic Ag-contained WTS analogs, mainly attributing to the presence of LSPR generated hot electrons that are of high reductive capability. In addition, the retained electrons on the WO$_3$ valence band can disperse onto the SBA-15 matrix by means of electron delocalization effect [20], thus promoting the photocatalytic performance through the efficient inhibition of recombination process.
4. Conclusion

In this study, a series of mesoporous W-Ti-SBA15 composites and their Ag-deposited analogs were constructed and fully characterized by various techniques. It was found that original mesoporous structures were well remained and semiconductor clusters in SBA-15 matrix were gradually decreased in size along with the debatement of TiO2/SiO2 mass ratio. The deposition of Ag had almost no effect on the morphology, microstructure, and textural properties of composites. However, the visible-light absorption capability of composites was significantly enhanced due to the LSPR effect induced by the metallic Ag deposited. As for the degradation of dye RhB under visible light irradiation, as-synthesized WTS composites showed satisfactory photocatalytic efficiency that could be further enhanced by using Ag-contained ternary plasmonic composites. The enhancement of photocatalytic performance was discussed and a possible photocatalysis mechanism was finally proposed basing upon reactive radical species trapping experiments.

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Appendix A.
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