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BiOCl/SnS$_2$ hollow spheres for the photocatalytic degradation of waste water

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Abstract

The doping effect of a new dopant to BiOCl as photocatalyst is studied. SnS$_2$ as a narrow-band-gap semiconductor is used to tune the electronic and band structure of wide-band-gap BiOCl. BiOCl/SnS$_2$ hollow spheres are synthesized by a facile and economic one-pot hydrothermal method. The shell of the hollow spheres is composed by BiOCl and SnS$_2$, but the outermost layer is BiOCl. BiOCl/SnS$_2$ hollow spheres exhibit much higher photocatalytic activity than pure SnS$_2$ and BiOCl in the degradation of Rhodamine-B under visible light and sun-light irradiation. The improvement is explained by the band structure tuning effect by SnS$_2$ and the morphological benefits of the hollow sphere structure.

Keywords:
Halogen bismuth oxide; Tin disulfide; Hollow spheres; Photocatalysis

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1. Introduction

Photocatalysis is a hot topic in solar energy conversion and photo-oxidation of organic pollutants.[1-6] A contradiction exists for photocatalysts: wide-band-gap semiconductors usually have high photocatalytic performance but poor response to visible light, which leads to low utilization efficiencies of full solar energy.[7-9] Band-structure engineering is expected to tune the electronic and band structures of wide-band-gap semiconductors to extend the photo response from the ultra-violet (UV) region to the visible light region.[2, 10-13] Bismuth oxychloride (BiOCl), a layered wide-band-gap ternary semiconductor, has recently attracted considerable attention in photocatalysis.[14-16] BiOCl is an ideal parent semiconductor to be doped by other semiconductors.[17, 18] Doped BiOCl has advantages such as high reduction potential of the holes, low electron-hole recombination rates and the easy formation of oxygen vacancies.[17, 19] Doping is an effective way for band-structure tuning, for example anion such as Iodine doping is reported to broaden the light response range.[10, 19, 20] SnS$_2$ is a kind of n-type semiconductor with band gap of 2.18-2.44eV, the electronic property and band potentials matches well with BiOCl, in this work SnS$_2$ is chosen as the dopant to BiOCl to achieve high visible-light-responsive ability.[21-23] Both the valence band and conduction band potentials of SnS$_2$ are more negative than those of BiOCl, this thermodynamically allows the photogenerated electron transfer from the conduction band of SnS$_2$ to the conduction band of BiOCl under visible light irradiation, which can enhance the separation of photogenerated electrons and holes in SnS$_2$ and increase sensitivity of BiOCl.[24-26] Thus, the SnS$_2$-incorporated BiOCl composites are expected to perform higher visible-light-driven photocatalytic activity than individual SnS$_2$ and BiOCl.

The burgeoning nanotechnology also sheds light to the possibility of tuning the band structure by constructing superstructures which can make use of the advantage of the wide-band-gap semiconductors and reduce the shortcomings of doping materials.[22-26] BiOCl compounds usually exist in tetragonal matlockite structure, which consists of [Bi$_2$O$_2$]$^{2+}$ layers sandwiched between two sheets of Cl$^-$ ions.[15, 16, 18, 19] Therefore, it is difficult to control the morphologies of BiOCl crystals. In most reports BiOCl are layered sheets in micrometer scale.[27, 28] The property of nano-material is greatly influenced by morphology, and shape control becomes vital in the preparation of nano-sized functional materials. Here for the first time we constructed a totally new BiOCl-based nanostructure: hollow spheres with BiOCl at the outmost surface. The
hierarchical architecture can allow multiple reflections of light, which enhances 
light-harvesting efficiency and thus increase the quantity of photogenerated electrons 
and holes available to participate in the photocatalytic reaction. [29, 30]

In this work, a general and facile one-pot hydrothermal synthesis of a novel 
BiOCl/SnS$_2$ hollow spheres is proposed. The resulted composites show enhanced light 
harvesting ability and high performance in the degradation of organic pollutants under 
visible light and sun-light irradiation.

2. Experimental

All the reagents were purchased from Aladdin reagent (China) Co., Ltd and were used 
as received without further purification. All water was deionized water prepared in 
lab.

2.1 Synthesis

Synthesis of hexagonal SnS$_2$ nanoflakes

In a typical preparation process, 1 mmol SnCl$_4$·5H$_2$O, 4.0 mmol Thiourea and 0.2 g 
Polyvinyl pyrrolidone (PVP) were dissolved in 40mL deionized water. The mixture 
was vigorously stirred for 30 minutes to ensure homogeneous dispersion. The solution 
was transferred to a 50mL Teflon-lined stainless steel autoclave and maintained at 
120℃ for 12h. After cooling to room temperature the product was collected by 
centrifugation, followed by washing thoroughly with deionized water and absolute 
ethanol. The final product was dried at 60℃ in vacuum for 8h.

Synthesis of BiOCl nanoflakes

The procedures are the same with the synthesis of SnS$_2$ nanoflakes except the reactant 
changed into 1 mmol Bi(NO$_3$)$_3$·5H$_2$O and 1 mmol KCl dissolved in 40 ml deionized 
water.

Synthesis of SnS$_2$-incorporated BiOCl hollow microspheres

The same procedures are applied, the reactant changed into the mixture of 1 mmol 
SnCl$_4$·5H$_2$O, 4.0 mmol Thiourea, 1 mmol Bi(NO$_3$)$_3$·5H$_2$O, 1 mmol KCl, 1 mmol 
citric acid and 0.2 g PVP.

To study the role of PVP and citric acid, parallel experiments are performed without 
citric acid.

2.2 Physical characterizations

The morphologies of the samples were characterized by field emission scanning 
electron microscope (FESEM) (JEOL JSM-T300, operated at 10kV) and field 
emission transmission electron microscope (FETEM) (JEOL SM-6330F, operated 200
kV). Energy dispersive X-ray spectroscopy (EDX) (JEOL JSM-6330F equipped with an X-ray energy dispersive spectrometer) was used to analyze the element composition. The crystal structure of the samples were determined on an X-ray diffractometer (XRD) (MSALXD-2 with Cu-Kα radiation, D/Max-III A, Rigaku Co., Japan, CuK1, λ=1.54056 Å) and selected area electron diffraction (SAED). X-ray photoelectron spectroscopy (XPS) spectra were acquired using an ESCA Lab 250 (Thermo VG) with 200 W Al Kα radiation in twin anode and the distance between X-ray gun and sample is about 1 cm. The analysis chamber pressure is about $2\times10^{-7}$ Pa and the pass energy is 20 eV for high resolution scans.

2.3. Photocatalytic performance characterizations

Photocatalytic activities of the samples were evaluated by photo-bleaching of Rhodamine B (RhB) under UV-vis (385-740 nm) and visible light (λ>420 nm) irradiations. Experiments were carried out in a 250 mL cylindrical-shaped-glass reactor at room temperature in the air and at neutral pH conditions. A suspension of 10 mg photocatalyst and 100 ml aqueous solution of RhB (10 mg L$^{-1}$) was magnetically stirred for at least 2 hours before irradiation to establish an adsorption-desorption equilibrium. The optical system used for the photocatalytic reaction consisted of a Xenon lamp (XQ350W) and a 420 nm cutoff filter, which was placed under the reaction cell to completely remove all incoming light with wavelengths shorter than 420 nm to ensure irradiation with visible light only. At specific time intervals, 4 ml suspension were sampled and centrifugated at 5000 rpm for 10 min to remove the photocatalyst powder. The concentration of remnant RhB in the solution after irradiation was analyzed by UV-vis spectrophotometer by recording the variations of the absorbance band maximum at λ=554 nm (RhB).

2.4 Photoelectrochemical Measurements

In the fabrication of the photo anode, 20 mg as-prepared photocatalyst powder and 2 ml ethanol solution were mixed homogeneously. The mixture was sprayed on an FTO (fluorine doped tin oxide) glass and allowed to dry under ambient conditions. Photocurrent was measured by an electrochemical workstation(SP-150 from Bio-Logic Science Instruments) in a standard three-electrode system with the as-prepared samples as the working electrodes with an active area of ca. 6 cm$^2$, a Pt wire as the counter electrode, and a saturated calomel electrode (SCE) as reference electrode. A 350 W Xe lamp equipped with an ultraviolet cutoff filter (λ>420 nm) was utilized as the visible-light source. A 0.1 M Na$_2$SO$_4$ aqueous solution was used as the
3. Results and discussion

3.1 Characterization of the separately prepared samples

FESEM images (Fig. S1a and b) clearly show that SnS$_2$ are hexagon nanoflakes with a uniform size of 200–300 nm in length and ~20 nm in thickness. PVP plays an important role in the shape control. As capping agent PVP has long alkyl chains with a molecular weight of 55,000 and the carbonyl groups in the backbone structure determine the shape of SnS$_2$ by the effect of the Van der Waals force and hydrogen bond. Trace amount of PVP can be selectively adsorbed on certain facets, promoting overgrowth in the specific energetically favorable direction. The microstructure was further characterized by HRTEM. As shown in Fig. S1c and d, the same morphology of hexagon nanoflake was observed which is in good accordance with the FESEM images. The atomic arrangement was observed by HRTEM of a selected part of a single nanoflake shown in Fig. S1d. Clear lattice planes with high crystallinity were observed. The resolved lattice fringe is 0.316 nm and corresponds to the d-spacing of (100) lattice planes of hexagonal phase SnS$_2$. The SAED inset in Fig.S1d shows hexagonal array of spots, which revealed single crystal structure of the nanoflake. From SAED it is concluded that the [001] orientation is the preponderant growth direction of the SnS$_2$ nanoflake, which is parallel to the (100) facet.

In this work BiOCl is prepared by a simple hydrothermal reaction of the mixture of Bi(NO$_3$)$_3$·5H$_2$O and KCl. Since no morphology control is applied, the morphology of as-prepared BiOCl nanoflakes was asymmetric polygon nanoflakes with a diameter of about 0.4–4µm as shown in the FESEM and FETEM images of Fig. S2. The clear lattice stripe of SAED (inset in Fig.S2c) exhibits the single crystalline nature of the nanoflakes. The diffraction spots of the typical tetragonal phase could be indexed to (110) and (200) facets. The interplanar distance of the lattice is 0.274 nm, which is consistent with the (110) planes of the tetragonal system of BiOCl. The HRTEM in Fig.S2d also confirms that the growth direction of BiOCl flake is along the [001] orientation, which is parallel to the (110) plane.

3.2 Physical characterization of the BiOCl/SnS$_2$ hollow spheres

From above results it is found that when SnS$_2$ or BiOCl are prepared separately both are flakes. The difference is SnS$_2$ are hexagon nanoflakes with smaller size while BiOCl flakes are bigger and have irregular shape. The idea of this work is to extend the light absorption edge of BiOCl to the visible light region by forming BiOCl/SnS$_2$
composites. It is found citric acid plays a crucial role in the morphology development of the BiOCl/SnS$_2$ composites. As shown in Fig. 1a and 1b, when no citric acid added the morphology of as-prepared BiOCl/SnS$_2$ composites is a flower like architecture which is composed by layers of flakes. The flakes have irregular shapes with a thickness of about 50 nm and a width of about 0.5-1.0µm and the shapes are different from either SnS$_2$ or BiOCl. In the presence of citric acid, the FESEM image (Fig. 1c) demonstrates that the as-obtained product is consisted of monodispersed hollow spheres with a good uniformity and narrow size distribution. The spheres have an average diameter of 500-600 nm and have a hole on the surface, showing hollow structure inside. A magnified image of the hollow spheres (Fig. 1d) exhibits detailed surface information. The shell was composed by nanosheets with a thickness of about 15nm as building unit, the nanosheets form a porous and loose packed surface. At the same time some broken spheres can be found, also proving the hollow structure.

XRD is used to confirm the composition of the BiOCl/SnS$_2$ composites. Fig.2 shows the XRD patterns of as-synthesized pure SnS$_2$, pure BiOCl and SnS$_2$-incorporated BiOCl composites. From Fig.2a and b hexagonal SnS$_2$ phase and tetragonal BiOCl phase are identified from standard card JCPDS 89-2028 and JCPDS 82-0485, respectively. Both flower-like and hollow spheres of BiOCl/SnS$_2$ composites are composed by tetragonal BiOCl phase and hexagonal SnS$_2$ phase, which is proved by the coexistence of all characteristic peaks of SnS$_2$ and BiOCl phases (Fig. 2c and d). Comparing Fig.2c and d, it is found the full width at half maximum (FHWM) of the XRD peaks in Fig.2d is much wider than that of Fig.2c. According to calculation with Scherrer equation$^{[31]}$ the crystal size of hollow spheres of BiOCl/SnS$_2$ composites is smaller than the flower-like structure. It is also interesting to note that the ratio between BiOCl (001) peak and SnS$_2$ (001) peak is different for BiOCl/SnS$_2$ hollow spheres and flower-like structure, showing difference in crystalline growth. Energy dispersion spectrum (EDS, Fig.S3) also proves the coexistence of Sn, S, Bi, O and Cl elements and the atomic ratio of the elements is in good accordance with the stoichiometric ratio of BiOCl and SnS$_2$.

Compared with the flower-like structure, the hollow spheres of BiOCl/SnS$_2$ composites have the advantages of smaller particle size revealed by SEM in Fig.1 and smaller crystal size revealed by XRD in Fig.2. The BiOCl/SnS$_2$ is expected to have better performance in application of photocatalysis. The questions of how BiOCl and SnS$_2$ form the hollow spheres, which is on the surface and the details of the structure
of the spheres must be answered before studying the photocatalytic property. Fig.3a and b are TEM images of the BiOCl/SnS₂ spheres. The size and morphology of the sphere are in accordance with the SEM images in Fig.1. Fig.3b is an enlarged picture of the spheres, from which it is observed that the inner part of the sphere is grayer than the shell, proving the hollow structure. On the spheres in the middle of Fig.3a (circled by red ring) some bright parts in the gray area are found, which corresponds to the holes found on the surface of the sphere in Fig. 1c and d. The thickness of the shell of the hollow spheres is found to be about 80 nm from Fig.3b. Fig.3b also shows the spheres are composed by layered nanosheet units connected to each other. The crystal property of the shell of the spheres is studied by HRTEM as shown in Fig.3c. Two d values of lattice spacing are indexed to be 0.215 and 0.343 nm, corresponding to the spacing of the (102) crystal plane of SnS₂ and (101) crystal plane of BiOCl, respectively. This indicates the presence of the mixed phase of SnS₂ and BiOCl in the hierarchical hollow spheres, which is in consistent with XRD results, the (102) peak of SnS₂ and (101) peak BiOCl are presented in the XRD of Fig.2. The SAED pattern (Fig. 3d) indexed out the diffraction patterns of (110) and (101) plane for BiOCl. The (110) plane is parallel to the growth direction of BiOCl and the (101) plane is also observed in the HRTEM in Fig.3c. For SnS₂ only (102) and (011) facet are indexed, (100) facet information is missing. The diffraction rings from the (110) Bragg reflections of tetragonal BiOCl overlaps with the diffraction spots from the (011) Bragg reflections of the hexagonal SnS₂ due to the coincidence of lattice spacing of the two crystals.

In TEM characterization, long time bombardment of the sample surface by high energy electron beam will melt the selected area and make it grow bigger.\cite{32, 33} Based on this observation we randomly selected a flake on the surface of the hollow sphere as labeled by the red ring in Fig.4. After 60s of high energy electron beam bombardment the flake grow bigger (inside the red ring in Fig.4). HRTEM image in Fig.4 shows uniform crystal property of the onsite-grown crystal, the d distance of 0.343 nm corresponds to the (101) facet of BiOCl. This result proves the surface of the hollow sphere is BiOCl. From TEM results a conclusion could be reached: the shell of the hollow sphere is composed by both BiOCl and SnS₂, but the outer most is BiOCl.

To gain detailed information about the dispersion of SnS₂ and BiOCl throughout the whole sphere, Energy dispersive X-ray spectroscopy (EDS) mapping were conducted.
EDS mapping of a typical hollow sphere shows the distribution of the Bi, O, Cl, Sn and S elements (Fig. 5). It is noted that Bi, O, Cl, Sn and S elements are uniformly distributed throughout the sphere. EDS mapping can only show the distribution of the elements, the chemical state and the structure of the sphere still need further characterizations.

The X-ray photoelectron spectroscopy (XPS) is a powerful tool to get information of the chemical state and valence band states of the elements on the surface of the materials. In this work, XPS results can also provide useful information on the structure of the material. XPS spectrum in Fig.6a disclosed the presence of Bi, O, Cl, Sn and S components, as well as C. The result is in accordance with the XRD and EDS results. Typical high-resolution XPS spectra for Bi 4f and Sn 3d are shown in Fig. 6b and 6e. It is clear that the binding energies of Bi 4f$_{7/2}$ (159.6 eV) and Bi 4f$_{5/2}$ (164.9 eV) in SnS$_2$/BiOCl composites are higher than those of pure BiOCl (159.1 eV and 164.4 eV), which is characteristic of Bi$^{3+}$.[18, 25, 27] Nevertheless, the binding energies of Sn 3d$_{5/2}$ and Sn 3d$_{3/2}$ peaks of SnS$_2$/BiOCl are located at 486.2 eV and 494.9 eV which lower than those of pure SnS$_2$ (486.8 eV and 495.3 eV). This is explained by the fact that the Fermi levels of SnS$_2$ are lower than that of BiOCl, so that the electrons on Bi atom can transfer to the Sn atom in SnS$_2$ dispersed on the surface of BiOCl, which causes the change in the outer electron cloud density of Bi and Sn ions and makes the Bi 4f$_{7/2}$ and Bi 4f$_{5/2}$ binding energies increase and Sn 3d$_{5/2}$ and Sn 3d$_{3/2}$ binding energies decrease. The interaction of the Fermi level electron and electron cloud may change the property of individual BiOCl and SnS$_2$, leading to new properties. Fig. 6c shows the high-resolution XPS spectrum of O 1s, which can be fitted into two peaks. The main peak at 530.7 eV belongs to the Bi-O bonds in (BiO)$_2^{2+}$ an BiOCl, and the peak at 532.1 eV is assigned to the hydroxyl groups on the surface.[27] In the high-resolution spectra of the Cl 2p (Fig. 6d), two peaks at 197.9 and 199.4 eV are attributed to Cl 2p$_{3/2}$ and Cl 2p$_{1/2}$ respectively, they can be ascribed to Cl$^-$ in BiOCl and SnS$_2$-incorporated BiOCl composites.[27] Two peaks at 161.5 eV and 162.7 eV shown in Fig. 6f are attributed to S 2p$_{3/2}$ and S 2p$_{1/2}$, which are characteristic of $S^{2-}$ in SnS$_2$.[25] In conclusion, according to the XPS results, the coexistence of SnS$_2$ and BiOCl in the composites is confirmed and the Fermi level electron and electron cloud interact with each other.

3.3 Discussion on the growth mechanism and structure details of the hollow spheres

From TEM of Fig.3b the SnS$_2$-incorporated BiOCl composite is composed by hollow
spheres. EDS mapping results of Fig.5 proves the distribution of the elements is uniform throughout the spheres. But how SnS$_2$ and BiOCl form the spheres is not clear. High-angle angular dark field scanning transmission electron microscopy (HAADF-STEM) and bright field scanning transmission electron microscopy (BF-STEM) are used to further verify the structure of the spheres. The BF-STEM images (Fig. S4a and b) are similar to TEM images in Fig.3. However in the HAADF-STEM images (Fig. S4c and d) the outer wall of the hollow spheres has a strong brightness contrast compared with the low brightness contrast of the inner part of the sphere. In the HAADF mode the high brightness contrast shows the presence of heavier elements such as Bi and the low brightness contrast shows the presence of light elements such as Sn. This is a strong proof of the structure of the hollow spheres: BiOCl is in the outer layer and SnS$_2$ in the inner layer, both of them form the shell of the hollow spheres. This structure is also proved by the high energy electron beam bombardment of the sample as discussed in Fig.4. XPS results also support this conclusion. From Fig.6e and f it is noted that on the surface of the SnS$_2$-incorporated BiOCl hollow spheres the peaks of Sn3d and S2p are hard to find with very small peak height. This is a direct proof that the outer layer of the hollow spheres is mainly composed by BiOCl layer.

Based on above analysis, the growth mechanism of SnS$_2$-incorporated BiOCl hollow spheres is illustrated in Scheme 1. In the beginning of the crystal growth, the nucleation of BiOCl and SnS$_2$ crystals are modulated by the absorbance and complex of PVP on the surface. PVP leads to the formation of nanoplates and the nanoplates form the flower-like structure by self-assembly which is driven by van der Waals forces and hydrogen bond among the organic molecules on the surface of nanomaterials.$^{[27, 28]}$ Citric acid as a biological ligand for metal ions can form strong complexes with Bi$^{3+}$, Al$^{3+}$, Ca$^{2+}$, Fe$^{3+}$, Zn$^{2+}$, Mg$^{2+}$, etc.$^{[28]}$ Citric acid can be hydrolyzed into [C$_6$O$_7$H$_5$]$^{3-}$ and H$^+$ which will affect the nucleation, growth and self-assembly of crystals. The [C$_6$O$_7$H$_5$]$^{3-}$ ions can chelate with Bi$^{3+}$ to form [C$_6$O$_7$H$_5$]$^{3-}$Bi$^{3+}$. $^{[14, 27]}$ Hydrogen bond can be formed between the carboxyl of citric acid and the hydroxyl ions in [C$_6$O$_7$H$_3$]$^{3-}$Bi$^{3+}$, and between the hydroxyl ions in [C$_6$O$_7$H$_3$]$^{3-}$Bi$^{3+}$. The self-assembly growth of SnS$_2$-BiOCl nanoplates is controlled by the interaction between [C$_6$O$_7$H$_3$]$^{3-}$Bi$^{3+}$ and hydrogen bond. In the presence of citric acid nanoplates of BiOCl and SnS$_2$ are formed first. Citric acid is then absorbed on the surface of the nanoplates, with the other end forming [C$_6$O$_7$H$_5$]$^{3-}$Bi$^{3+}$ with Bi$^{3+}$ in
the hollow structure is the result of the interaction between the
$[\text{C}_6\text{O}_7\text{H}_5\text{Bi}_3]$ complex ion and hydrogen bond. The surface $[\text{C}_6\text{O}_7\text{H}_5\text{Bi}_3]$ complex
ion forms BiOCl which forms the outer layer of the shell of the hollow sphere.

3.4 Characterization of the photocatalytic activity of the BiOCl/SnS$_2$ hollow spheres

The photocatalytic activity of the BiOCl/SnS$_2$ hollow spheres was evaluated by the
degradation of RhB, a typical organic contaminant. Fig. 7 presents the variation of
RhB concentration ($C/C_0$) as the function of irradiation time for experiments carried
out under irradiation at wavelengths in the range of 385-740 nm and visible light
irradiations at $\lambda>420$ nm, respectively. For comparison, the photocatalytic activity of
Degussa’s P25 and the degradation of RhB without photocatalysts were tested under
the same conditions. The blank test without photocatalysts shows that RhB is very
stable and does not decompose even after long-time illumination with UV-vis or
visible light irradiations. A rapid decrease in the concentration of RhB with time is
realized in the presence of the as-prepared SnS$_2$-incorporated BiOCl photocatalysts.

For comparison, the photocatalytic property of pure SnS$_2$ and BiOCl were also tested.
All the samples exhibited much higher photocatalytic activity under UV-vis and
visible light irradiation compared to the standard (Degussa’s P25). Among them the
BiOCl/SnS$_2$ hollow spheres showed the highest RhB removal rate with total
decomposition within only 50 min under UV-vis light irradiation and within only 70
min under visible light irradiation, which is a triple enhancement in comparison to the
standard (Degussa’s P25). After 90 minutes of visible light irradiation the degradation
efficiency of RhB on pure SnS$_2$ and BiOCl are less than 21% and 43%, respectively,
which is significantly less than that of the BiOCl/SnS$_2$ hollow spheres. The
remarkable improvement of the photocatalytic performance of the BiOCl/SnS$_2$ hollow
spheres is explained by three reasons. The first point is the tuning effect of SnS$_2$. Both
the valence band and conduction band potentials of SnS$_2$ are more negative than those
of BiOCl. This thermodynamically allows the photogenerated electron transfer from
the conduction band of SnS$_2$ to the conduction band of BiOCl, which can enhance the
separation of photogenerated electrons and holes in SnS$_2$ and increase the
sensitization of BiOCl. Secondly the hierarchical assemble of nanoflakes greatly
increases the surface area.$^{[27, 28]}$ Thirdly, the hierarchical architecture can allow
multiple reflections of light, which enhances light-harvesting efficiency and thus
increase the quantity of photogenerated electrons and holes available to participate in
the photocatalytic reaction.$^{[27, 28]}$ It should also be pointed out that the BiOCl/SnS$_2$
hollow spheres show much higher activity than the flower-like structure. This is
due to at least two factors. The bigger size of the flower-like structure leads to
smaller surface area and reduces the efficiency. The second is the benefit of the
hollow structure: both the inner and outside wall of the sphere can be utilized in the
photocatalysis, which greatly improves the efficiency. This explanation is supported
by the Brunauer-Emmett-Teller (BET) specific surface areas of the SnS$_2$-incorporated
BiOCl samples with different morphologies by using nitrogen adsorption-desorption
isotherms. The BET specific surface area of SnS$_2$-incorporated BiOCl hollow spheres
was calculated to be 32.5639 m$^2$g$^{-1}$, which was much larger than the flower-like
structure without adding citric acid (23.4383 m$^2$g$^{-1}$) (Fig. S5 and Table 1 in the
supporting materials). The pores of the SnS$_2$-incorporated BiOCl hollow spheres are
mesopores, which provide much better access of the solution to the reaction sites than
the micropores of the flower-like structure. Besides the higher activity, the
SnS$_2$-incorporated BiOCl hollow spheres also show strong durability during the
photocatalytic degradation of RhB. After ten times of cycle test the efficiency only
drops by 5%, which reaches the best durability reported in literature.$^{[19,28]}$

To further investigate the photocatalytic mechanism, the transient photocurrent
responses of the samples were recorded via several irradiation on-off cycles. The
representative plots are shown in Fig. 9. When the light was turned on the
photocurrent reached a high point and followed by a fast decline, forming a spike,
then decreased gradually until a constant value is reached. The photocurrent was
instantaneously close to zero as long as the light was switched off. The initial anodic
photocurrent spike is caused by the separation of electron-hole pairs at the
BiOCl/SnS$_2$ composite.$^{[31,32]}$ Holes move to the BiOCl surface, where they are trapped
or captured by reduced species in the electrolyte. At the same time the electrons are
transported to the conduction band of SnS$_2$. The decay of the photocurrent is caused
by the recombination of photogenerated electron-hole pairs. Instead of capturing
electrons from the electrolyte, a portion of the holes at the surface of BiOCl
recombine with electrons from the conduction band of SnS$_2$ or just accumulate at the
surface.$^{[34,35]}$ The electrons in the conduction band of SnS$_2$ take part in the reduction
of the photogenerated oxidized species in the electrolyte, which results in the decay of
the photocurrent. When the separation and recombination of the electron-hole pairs
reach an equilibration, a constant current is reached. When the light is turned off, the
holes accumulated at the BiOCl surface and the electrons in SnS$_2$ conduction band are
recombined simultaneously, leading to a fast drop of the photocurrent to zero. It can
be seen from Fig. 9 that BiOCl/SnS$_2$ exhibits a prompt generation of photocurrent with
good reproducibility. The much higher anodic photocurrent of the BiOCl/SnS$_2$
composite compared with BiOCl or SnS$_2$ alone is a direct proof of a much higher
efficiency of the separation of electron-hole pairs, which is the origin of the synergetic
effect resulting in the high photocatalytic performance of the BiOCl/SnS$_2$ composite.

3.5 Discussion on the origin of the improved photocatalytic property of the
BiOCl/SnS$_2$ composite

UV-vis diffuse reflectance spectroscopy of unmodified SnS$_2$, BiOCl and BiOCl/SnS$_2$
were used to study the band gap of the samples. As shown in Fig. S6, BiOCl absorbs
the ultraviolet light with an absorbance edge of about 360 nm, whereas SnS$_2$ and
BiOCl/SnS$_2$ exhibit an obvious red shift and show good absorptions in the visible
light range with an absorbance edge around 550 and 450 nm, respectively, suggesting
that the incorporation of SnS$_2$ can effectively extend the light absorption edge of
BiOCl from ultraviolet into visible region. For the crystalline semiconductor, the band
gap ($E_g$) can be calculated according to the following equation:

$$\alpha h v = A (h v - E_g)^{1/2}$$

where $\alpha$, $h$, $v$, $E_g$, $A$ are the absorption coefficient, Plank constant, light frequency,
band gap energy, and a constant, respectively. Hence, $E_g$ can be estimated by
extrapolating the linear portion of $(\alpha h v)^2$ vs. $(h v)$ curves (Fig. S6B). The $E_g$ value of
BiOCl, BiOCl/SnS$_2$ and SnS$_2$ are calculated to be about 3.20, 2.38, 2.17 eV,
respectively. The fact that the $E_g$ value of BiOCl/SnS$_2$ is between those of pure SnS$_2$
and BiOCl is a direct proof of the formation of the BiOCl/SnS$_2$ p-n junction.

The photoelectrochemical method is used to determine the conduction band (CB) of
SnS$_2$ and valence band (VB) of BiOCl semiconductors. The photocurrent spectra of
SnS$_2$ and BiOCl are shown in Fig. S7 a and b. In the photocurrent spectra of SnS$_2$, the
flat band potential is at -0.98 V versus SCE. For n-type semiconductor, the flat band
potential is closely related to the bottom of the CB. The VB of SnS$_2$ is calculated to be
1.19 V according to the corresponding $E_g$ value (2.17 eV). The photocurrent onset
potential (conduction band edge) of BiOCl is at +0.88 V versus SCE. As the band gap
energy of BiOCl is 3.20 eV, the CB edge is calculated to be -2.32 V. The energy band
structure diagram of SnS$_2$ and BiOCl is schematically represented in Fig. S8. BiOCl is a p-type semiconductor with Fermi level close to the VB, whereas SnS$_2$ is a n-type semiconductor with Fermi level close to CB. When the Fermi levels of p-BiOCl and n-SnS$_2$ reach equilibration, an internal electric field directed from n-SnS$_2$ to p-BiOCl is built, preventing the charge migrating from n-SnS$_2$ into p-BiOCl. The energy band positions of BiOCl in BiOCl/SnS$_2$ heterojunction shift upward along the Fermi level ($E_{f,n}$) and those of SnS$_2$ shift downward along its Fermi level ($E_{f,p}$). As a result, the band positions of BiOCl and SnS$_2$ in the heterojunction have a type-II band structure in which the CB and VB of SnS$_2$ lie below those of BiOCl. The formation of heterojunction between BiOCl and SnS$_2$ results in a pseudo band (PB in Fig.S8) between VB and CB of BiOCl, which can accept photogenerated electrons from VB. Thus the band gap of BiOCl is narrowed, leading to response to visible light. The electrons in the PB can continue to move to the CB of BiOCl when absorbing more visible light. Due to the effect of internal electric field, the photoelectrons on the CB bottom of BiOCl will transfer to that of SnS$_2$ and be trapped by oxygen (O$_2$) to form reactive super oxide radicals (O$_2^-$) or OH$^-$, and the holes remained in the VB of BiOCl will react with the reactants adsorbed on the surface of BiOCl/SnS$_2$. Thus, the photogenerated electron-hole pairs will be effectively separated because of the formation of p-n junction formed between BiOCl and SnS$_2$, resulting in a reduced electron-hole recombination rate.

4. Conclusions

In summary, a novel BiOCl/SnS$_2$ hollow spheres is synthesized for the first time by a facile one-pot hydrothermal process. A triple enhancement in the visible light photocatalytic performance of the hollow spheres is observed, which is explained by the hierarchical structure and the doping effect of SnS$_2$ to BiOCl. The easy synthesis and high efficiency makes the novel composite a very promising candidate for
application in photocatalysis. The idea of this work can also be used to tune electronic
and band structure of other wide-band-gap semiconductors for application in both
solar energy conversion and photocatalysis in environmental protection.

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Figures and caption:

Fig. 1 FESEM images of as prepared SnS$_2$-incorporated BiOCl flower-like structure without citric acid (a, b) and hollow spheres with citric acid (c, d).

Fig. 2 XRD patterns of (a) pure SnS$_2$, (b) pure BiOCl, (c) SnS$_2$-incorporated BiOCl flower-like structure without citric acid and (d) SnS$_2$-incorporated BiOCl hollow spheres with citric acid.

Fig. 3 FETEM images (a, b and c) and SEAD pattern (d) of as prepared SnS$_2$-incorporated BiOCl HMSs. microspheres.

Fig. 4 Onsite growth of crystal by electron beam bombardment, (a): selected flake before bombardment, (b and c): after bombardment, (d): HRTEM of the bombarded area.

Fig. 5 EDS mapping of SnS$_2$-incorporated BiOCl hollow sphere, the elements presented are (a), Bi (b), O (c), Cl (d), Sn (e), S (f).

Fig. 6 XPS spectra of BiOCl, SnS$_2$ and SnS$_2$-incorporated BiOCl hollow spheres, survey spectrum (a), Bi 4f (b), O 1s (c), Cl 2p (d), Sn 3d (e), S 2p (f).

Scheme 1. Proposed formation mechanism of SnS$_2$-incorporated BiOCl hierarchical nanostructures.

Fig. 7 Comparison of visible-light photocatalytic activities, (a) SnS$_2$-incorporated BiOCl hollow spheres, (b) SnS$_2$-incorporated BiOCl hollow spheres of flower-like structure, (c) pure SnS$_2$, (d) pure BiOCl, (e) P$_{25}$, (f) adsorption in dark, (g) degradation of 10 mg L$^{-1}$ RhB without photocatalyst under UV-vis (385-740 nm) (A) and visible light ($\lambda$>420 nm) irradiations (B).

Fig. 8 Cycling degradation curves of SnS$_2$-incorporated BiOCl hollow spheres on the degradation of 10 mg L$^{-1}$ RhB under visible light ($\lambda$>420 nm) irradiations.

Fig.9 Comparison of transient photocurrent response of pure SnS$_2$, pure BiOCl and BiOCl/SnS$_2$ in 0.1 M Na$_2$SO$_4$ aqueous solution under visible-light irradiation vs SCE.
Fig. 1 FESEM images of as prepared SnS$_2$-incorporated BiOCl flower-like structure without citric acid (a, b) and hollow spheres with citric acid (c, d).
Fig. 2 XRD patterns of (a) pure SnS$_2$, (b) pure BiOCl, (c) SnS$_2$-incorporated BiOCl flower-like structure without citric acid and (d) SnS$_2$-incorporated BiOCl hollow spheres with citric acid.
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Fig. 9 Comparison of transient photocurrent response of pure SnS₂, pure BiOCl and BiOCl/SnS₂ in 0.1 M Na₂SO₄ aqueous solution under visible-light irradiation vs SCE.
A kind of novel SnS$_2$-incorporated BiOCl hollow spheres are synthesized. The band structure tuning effect and the morphological benefits of the hollow sphere structure with great improved performance in photocatalysis.