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BiOBr nanoparticle-modified $\text{Ti}_3\text{C}_2\text{T}_x$ MXenes for photocatalytic degradation of organic arsenic in wastewater

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Arsenic (As) contamination in water remains a serious concern due to its high toxicity and harmful effects on human health and the environment. Herein, we successfully synthesized a novel photocatalyst (BiOBr/Ti₃C₂) by *in situ* inserting BiOBr nanoparticles into Ti₃C₂T_x MXenes for the photocatalytic degradation of organic arsenic roxarsone (3-nitro-4-hydroxyphenylarsonic acid) in wastewater. BiOBr/Ti₃C₂ exhibited a unique morphology characterized by uniform BiOBr nanoparticles within more dispersed Ti₃C₂T_x layers. Heterojunctions were formed between Ti₃C₂T_x and BiOBr, which were conducive to photogenerated charge separation and electron transfer in Ti₃C₂T_x layers. An optimal BiOBr/Ti₃C₂ photocatalyst achieved a removal efficiency of 1.27 mg g_{cat}⁻¹ h⁻¹ for 2 mg L⁻¹ of roxarsone wastewater within 0.5 h and a higher removal rate (about 1.5 times at 3 h) than pure Ti₃C₂T_x. In addition, BiOBr/Ti₃C₂ exhibited an apparent quantum yield (AQY) of 29.5% and good reusability for 4 cycles. The enhanced photocatalytic performance was mainly attributed to the intercalation of BiOBr nanoparticles within Ti₃C₂T_x layers, which increased reaction space and improved the separation and transport of photocarriers. Holes (h⁺) and ·OH in the valence band (VB) of BiOBr/Ti₃C₂ were involved in the main route of roxarsone oxidative mineralization. The present BiOBr/Ti₃C₂ system provides fundamentals for the sustainable photocatalytic treatment of wastewater containing organic arsenic.

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Introduction

Roxarsone is an organic arsenic feed additive that has been widely used in the poultry and pig industries for decades to promote animal growth and treat coccidiosis.¹ A portion of the roxarsone added to the feed cannot be absorbed by poultry and pigs, and eventually remains in the faeces. Since roxarsone is highly water-soluble, it is easily transferred from animal feces to the surrounding aqueous environment, such as surface water or groundwater.^{2–6} Organic arsenic has low toxicity, but in the natural environment, it can be decomposed into highly toxic inorganic arsenic through biological or photooxidation reactions.^{7–9} Furthermore, inorganic arsenic can be divided into As(III) and As(V).¹⁰ As(III) and As(V) can be rapidly absorbed from the gastrointestinal tract, causing inevitable damage to the human body, further increasing their potential risks to human health and ecology.^{3,11–16} Aromatic amine feed additives, including organic arsenic, have been banned worldwide because

of the concerns about arsenic contamination in the environment.^{2,17} However, after decades of use, organic arsenic has been accumulated in large quantities in the environment. The degradation, transformation, migration and impact of organic arsenic on organisms have attracted the attention of many countries around the world. At present, the main reported organic arsenic removal technologies are categorized as biological methods,^{18–21} adsorption,^{22–26} and advanced oxidation processes such as UV-based processes,^{27–30} sulfite (S(IV)) oxidation processes,³¹ and Fenton/Fenton-like processes.^{32–34} However, these techniques have some limitations. The biological degradation of organic arsenic is slower and may produce more toxic alkyl arsenic compounds.¹⁸ Adsorption often requires long contact times and high costs.^{22,23} Advanced oxidation processes are introduced into the treatment of arsenic-containing water. As an advanced oxidation process, photocatalytic oxidation is widely used in the oxidative removal of organic or inorganic pollutants because it can use clean and pollution-free light energy,³⁵ produce a large number of strong oxidizing active substances and have no secondary pollution. However, highly toxic inorganic arsenic compounds will cause serious pollution to the environment, in case the inorganic arsenic compounds produced after oxidation are not treated reasonably in time. Therefore, the current photocatalysis technology has been widely considered and put into practice. The problems of current photocatalytic systems for the

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degradation of organic arsenic include the low utilization of visible light and the fast charge recombination speed, which limit their applications. Therefore, various photocatalysts, such as TiO_2 ,³⁶ ZnO ,³⁷ CuO ³⁸ and goethite³⁹ were developed to enhance the photodegradation of organic arsenic. Up to now, the MXene family has not been reported in the photocatalytic degradation of organic arsenic.

MXenes are a very famous family of two-dimensional (2D) materials,⁴⁰ and have been widely used in photocatalysis, batteries and supercapacitors.^{41–43} $\text{Ti}_3\text{C}_2\text{T}_x$, as a member of MXenes, has excellent electron transportability, a large specific surface area and adjustable active sites, which may promote the separation and transport of photoexcited electron–hole pairs.⁴⁴ The construction of composites based on $\text{Ti}_3\text{C}_2\text{T}_x$ is very simple because of the flexibility and unique layered structure of $\text{Ti}_3\text{C}_2\text{T}_x$. In $\text{Ti}_3\text{C}_2\text{T}_x$ composites, $\text{Ti}_3\text{C}_2\text{T}_x$ not only increases the absorption of light but also causes rapid carrier migration and inhibits its recombination.⁴⁵ Therefore, $\text{Ti}_3\text{C}_2\text{T}_x$ is often used as a noble metal⁴¹ co-catalyst to improve the photocatalytic activity of complexes.^{40,46–48} It has also been noted that $\text{Ti}_3\text{C}_2\text{T}_x$ with –F and –OH end groups, originating from exfoliation with HF, possess a narrow bandgap and also have potential photocatalytic properties.⁴⁹ However, the photocatalytic performance of MXene $\text{Ti}_3\text{C}_2\text{T}_x$ was not satisfactory. Its application is limited by rapid charge recombination. Various methods have been developed to overcome this problem. MXenes have been coupled to various semiconductors (such as TiO_2 and $\text{g-C}_3\text{N}_4$) to form heterojunctions, thereby facilitating photogenerated electron–hole separation and improving light absorption in photocatalyzed reactions. Considering that the properties of 2D accordion-like MXene $\text{Ti}_3\text{C}_2\text{T}_x$ can be improved by changing the elemental composition and adjusting the surface functional groups, it is an effective approach to introduce heteroatoms to modify the surface and improve the photocatalytic property.

Currently, many efficient photocatalyst materials have emerged.^{50,51} Among them, BiOBr is a typical bismuth oxyhalide material (BiOX , X = Cl, Br, I) that has been widely used in photocatalysis because of its suitable energy band structure and chemical stability.⁵² However, the photocatalytic performance of BiOBr is limited by the low separation efficiency of the charge carriers. Therefore, various strategies have been adopted to improve its photocatalytic performance,⁵³ such as bismuth-rich strategy, element doping,^{50,51,54,55} defect engineering and heterostructure construction.⁵⁶ Coupling with other semiconductors is one of the most common ways to accelerate the separation efficiency of photogenerated electron–hole pairs. It was found that there are almost no reports on BiOBr nanoparticle-modified MXene $\text{Ti}_3\text{C}_2\text{T}_x$ materials, even their use in the photocatalytic degradation of organic arsenic.

In this work, we successfully synthesized a novel composite of BiOBr nanoparticle-modified MXene $\text{Ti}_3\text{C}_2\text{T}_x$, which was used in the photocatalytic removal of roxarsone from wastewater. A heterojunction was attempted to establish between MXene $\text{Ti}_3\text{C}_2\text{T}_x$ and BiOBr nanoparticles to improve the degradation efficiency of roxarsone. The ultra-fine and highly dispersed BiOBr nanoparticles effectively expanded the specific surface area, enhanced light absorption, and promoted carrier separation on

the $\text{Ti}_3\text{C}_2\text{T}_x$ matrix. Compared with pure $\text{Ti}_3\text{C}_2\text{T}_x$, the degradation efficiency of roxarsone in the BiOBr/ Ti_3C_2 composite was significantly enhanced. A mechanistic study revealed that the holes (h^+) and $\cdot\text{OH}$ in the VB of BiOBr/ Ti_3C_2 primarily contributed to the oxidative mineralization of roxarsone.

Experimental

Chemicals

MXene $\text{Ti}_3\text{C}_2\text{T}_x$ was purchased from Jilin Yiyi Technology Co., Ltd. KBr and $\text{C}_2\text{H}_5\text{OH}$ were purchased from Tianjin Tianli Chemical Reagent Co., Ltd. $\text{Bi}(\text{NO}_3)_3 \cdot 5\text{H}_2\text{O}$ and 4-hydroxy-2,2,6-tetramethylpiperidinyloxy (TEMPO) were purchased from Macklin Chemical Co., Ltd. Dimethyl sulfoxide (DMSO) was purchased from Tianjin Fuyu Fine Chemical Co., Ltd. Phloroglucinol ($\text{C}_6\text{H}_6\text{O}_3$) was purchased from Tianjin Fuchen Co., Ltd. Ethylenediaminetetraacetic acid (EDTA) was purchased from Tianjin Beilian Fine Chemicals Development Co., Ltd. Isopropyl alcohol (IPA) was purchased from Aladdin Co., Ltd.

Preparation of BiOBr

0.669 g of KBr was dissolved in 75 mL of deionized water and heated to 90 °C. 2.429 g of $\text{Bi}(\text{NO}_3)_3 \cdot 5\text{H}_2\text{O}$ was dissolved in 100 mL of $\text{C}_2\text{H}_5\text{OH}$ by ultrasonication to form a suspension. KBr solution was added to the suspension. After 30 min of the reaction, a pale yellow precipitate was obtained. The yellow precipitate was collected by centrifugation, washing, and drying at 60 °C for 12 h to obtain BiOBr.

Preparation of BiOBr nanoparticles

1.07 g of the as-prepared BiOBr and 0.27 g of $\text{C}_6\text{H}_6\text{O}_3$ were dissolved in 40 mL of deionized water, stirred at 60 °C for 2 h, and then subjected to hydrothermal treatment at 200 °C for 12 h. The obtained product was washed 3 times with water and ethanol and dried at 60 °C for 6 h to obtain BiOBr nanoparticles.

Preparation of BiOBr/ Ti_3C_2

0.05 g of MXene $\text{Ti}_3\text{C}_2\text{T}_x$ and the required mass of BiOBr nanoparticles were dispersed into 40 mL of DMSO, stirred at room temperature for 12 h, and then treated by ultrasonication for 1 h. After centrifugation, washing with water and ethanol for 3 times, and drying at 60 °C for 8 h, a series of products labeled as BiOBr- x / Ti_3C_2 (x wt% = 5 wt%, 10 wt%, 20 wt% loading of BiOBr nanoparticles) was finally obtained.

Characterizations

X-ray diffraction (XRD) patterns were recorded using a Rigaku D/Max 2400 diffractometer with $\text{Cu K}\alpha$ radiation. Fourier transform infrared (FT-IR) spectroscopy was performed on a Nicolet Thermo 360 spectrometer using the KBr pellet method. Scanning electron microscopy (SEM) was performed using a JSM-7500 F field emission SEM (JEOL, Japan). Transmission electron microscopy (TEM) images were obtained using a FEI Tecnai G2 F20 field emission transmission electron microscope. X-ray electron spectroscopy (XPS, THERMO) was

used to analyze the composition and electronic states of the elements, and the scanning spectra of Ti 2p, C 1s, Bi 4f, Br 3d and O 1s were obtained. Photocurrent response and electrochemical impedance spectroscopy (EIS) were performed on an electrochemical workstation (CHI650D, China) with a 300 W xenon lamp and a 420 nm cut-off filter as the visible light source. UV-vis diffuse reflectance spectra were obtained on a JASCO V-770 spectrometer with BaSO₄ as the internal standard. Photoluminescence (PL) spectroscopy and time-resolved photoluminescence (TRPL) were performed using a fluorescence spectrometer (Edinburgh Instrument FLS980) with an excitation wavelength of 373 nm.

Photocatalytic activity tests

50 mg of the photocatalyst sample was dispersed in 50 mL of roxarsone aqueous solution containing 2 mg L⁻¹ of the initial concentration (unless otherwise noted). The system temperature was controlled at 0 °C using a water bath and was operated under continuous stirring. Before photoreaction, the reactant was controlled in the dark to reach the adsorption–desorption equilibrium. A 300 W xenon lamp with a cut-off filter of 420 nm was used as the visible light source. The suspension was then sampled and centrifuged at the required time interval. The absorbance of roxarsone was measured using the UV-vis spectrophotometer. Eqn (1) was used to calculate the removal efficiency of roxarsone. The liquid residual inorganic products of roxarsone after photocatalytic degradation were analyzed by high-performance liquid chromatography-inductively coupled plasma mass spectrometry LC-ICP-MS (LC, Prin-Cen, Elspe-2; ICP-MS, NeXion1000 G, PerkinElmer, USA). By changing the loading of BiOBr on Ti₃C₂T_x, the reaction temperature, pH value and initial concentration of roxarsone were determined according to the same procedure. The recycle test was also performed following the same procedure in 1 h using the regenerated photocatalyst in every cycle. By adding 2 mmol of the scavengers, a trapping experiment was carried out to study the radical contribution, during which EDTA, DMSO, IPA and TEMPOL were used as scavengers of h⁺, e⁻, ·OH and ·O₂⁻, respectively. The apparent quantum yields (AQY) of the photocatalytic degradation of roxarsone were measured using a monochromatic lamp with bandpass filters at 320, 450, 550, 600 and 750 nm. The irradiation intensity was measured as 10.95 W m⁻² using a light power detector (SM206-SOLAR). The irradiation area was controlled at 13.5 cm². The AQY of the photocatalytic degradation of roxarsone for 1 h was calculated using eqn (2).

$$D = (C_0 - C_t)/C_0 = (A_0 - A_t)/A_0 \quad (1)$$

where D is the removal efficiency, C_0 is the initial concentration of roxarsone, C_t is the concentration of roxarsone at t min, A_0 is the absorbance of roxarsone at the initial concentration, A_t is the absorbance of roxarsone at t min.

$$\text{AQY (\%)} = (N_e/N_p) \times 100\% \quad (2)$$

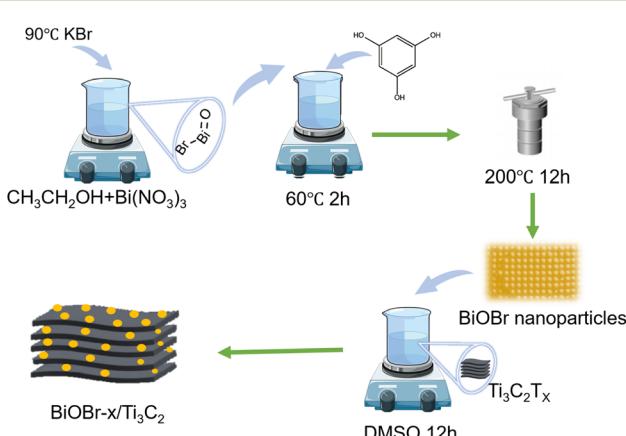
where N_e is the total number of electrons transferred during the reaction and N_p is the number of incident photons.

Results and discussions

Morphology and structure of materials

Scheme 1 illustrates the synthetic strategy of BiOBr/Ti₃C₂. The morphological changes and elemental distribution of BiOBr nanoparticles, Ti₃C₂T_x, and BiOBr-x/Ti₃C₂ were observed and compared by SEM and TEM images, as shown in Fig. 1a–l. The preparation of BiOBr/Ti₃C₂ was divided into two steps. After heat treatment with distilled water and phloroglucinol,⁵⁷ BiOBr became nanoparticles, which favored adhering to the surface of Ti₃C₂T_x. Subsequently, the BiOBr nanoparticles and Ti₃C₂T_x matrix were put into DMSO to make the MXene interlayer open and disperse more evenly. Finally, BiOBr nanoparticles were more easily loaded within the interlayers, and they further formed heterojunctions with MXene Ti₃C₂T_x. In the SEM images, pristine BiOBr agglomerates (Fig. 1a) and pristine MXene Ti₃C₂T_x do not show obvious delamination (Fig. 1b). For BiOBr-10/Ti₃C₂, DMSO showed a polar effect and increased the distance of the MXene Ti₃C₂T_x layers, so that BiOBr nanoparticles could be loaded and highly dispersed in Ti₃C₂T_x interlayers (Fig. 1c). BiOBr-10/Ti₃C₂ retained the accordion shape and exhibited a rough surface (Fig. 1c), indicating the strong combination of BiOBr nanoparticles with Ti₃C₂T_x and the formation of a heterojunction. The TEM images (Fig. 1d and e) show that Ti₃C₂T_x is well-laminated and loaded with ultrafine and uniform BiOBr nanoparticles. In the HRTEM image (Fig. 1f), BiOBr nanoparticles were embedded between the Ti₃C₂T_x lattices, indicating the existence of strong interactions and interfaces. The concave and convex surface of Ti₃C₂T_x provides a good platform for the formation of BiOBr/Ti₃C₂ heterojunctions. The elemental mapping images further confirmed the successful binding of BiOBr nanoparticles with Ti₃C₂T_x (Fig. 1g–i). Ti and C are widely distributed throughout the entire region of the composite as the main elements of the matrix, whereas Bi, Br, and O are evenly distributed and have relatively low density. The co-existence and overlapping distribution of the elements indicate a great opportunity to form a heterojunction on the surface of Ti₃C₂T_x.

XRD and FT-IR spectra were used to investigate the structure of the BiOBr-x/Ti₃C₂ composites with different BiOBr loads, as



Scheme 1 Schematic of the preparation of BiOBr/Ti₃C₂.



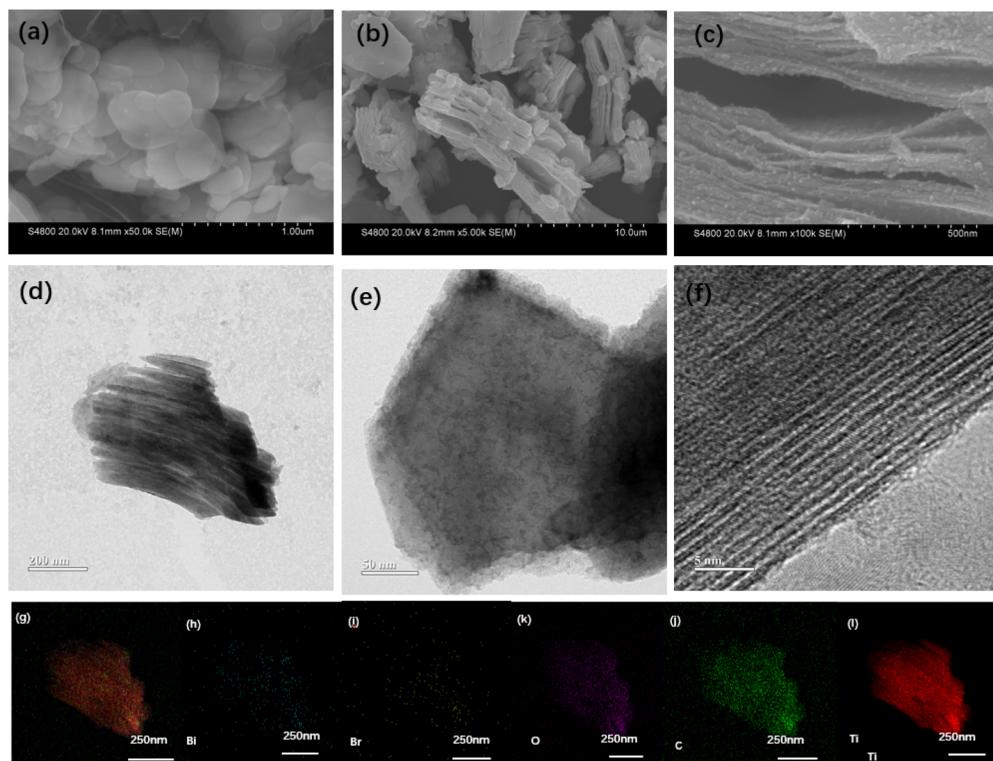


Fig. 1 SEM images of (a) BiOBr, (b) $\text{Ti}_3\text{C}_2\text{T}_x$ and (c) BiOBr-10/ Ti_3C_2 ; (d) TEM and magnified TEM (e) images of BiOBr-10/ Ti_3C_2 ; (f) HRTEM image of BiOBr-10/ Ti_3C_2 ; elemental mapping images of (g) overall distribution, (h) Bi, (i) Br, (j) C, (k) O and (l) Ti in BiOBr-10/ Ti_3C_2 .

shown in Fig. 2a and b, respectively. In Fig. 2a, pure BiOBr shows five peaks at 2 theta of 13.24° , 27.76° , 34.72° , 50.12° , and 60.08° , corresponding to the five crystal planes of (001), (200),

(040), (040), and (200), respectively.⁵² Pure $\text{Ti}_3\text{C}_2\text{T}_x$ shows four peaks at 2 theta of 10.06° , 19.72° , 25.2° , and 62.2° , corresponding to the four crystal planes of (001), (110), (111), and (220), respectively.⁵⁸ For the BiOBr- x / Ti_3C_2 composites, the crystal planes of (001), (200), (040), (040) and (200) become more obvious with the increase of BiOBr loading. The results indicate that BiOBr nanoparticles were successfully loaded onto $\text{Ti}_3\text{C}_2\text{T}_x$, but their size was larger and uneven at higher loading. $\text{Ti}_3\text{C}_2\text{T}_x$ was treated with DMSO to make its layers more dispersed and uniform and to increase the layer spacing. Therefore, the peaks intensity of the $\text{Ti}_3\text{C}_2\text{T}_x$ phase peaks increased with BiOBr loading in the composite, indicating that the $\text{Ti}_3\text{C}_2\text{T}_x$ matrix maintained the original accordion-like layer structure and provided larger layer spacing for BiOBr loading. However, the peak intensity of $\text{Ti}_3\text{C}_2\text{T}_x$ for BiOBr-20/ Ti_3C_2 did not increase because the $\text{Ti}_3\text{C}_2\text{T}_x$ layers were partially filled by overloaded BiOBr. The obvious delamination of $\text{Ti}_3\text{C}_2\text{T}_x$ and the increase in the contact area with BiOBr nanoparticles increase the tendency to form heterojunctions, which is conducive to the transport of photogenerated charges.

FT-IR spectroscopy was used to compare the variation of the main groups in the BiOBr/ Ti_3C_2 composites, as shown in Fig. 2b. The FT-IR spectrum of pure $\text{Ti}_3\text{C}_2\text{T}_x$ has a Ti-C corresponding vibration band at 587 cm^{-1} .⁵⁹ The wide peaks at 3450 cm^{-1} belong to the O-H tensile vibrations, which may be caused by surface water molecules or terminal hydroxyl groups.⁵⁹ The FT-IR spectrum of pure BiOBr shows a peak at 517 cm^{-1} , which is attributed to the symmetric stretching

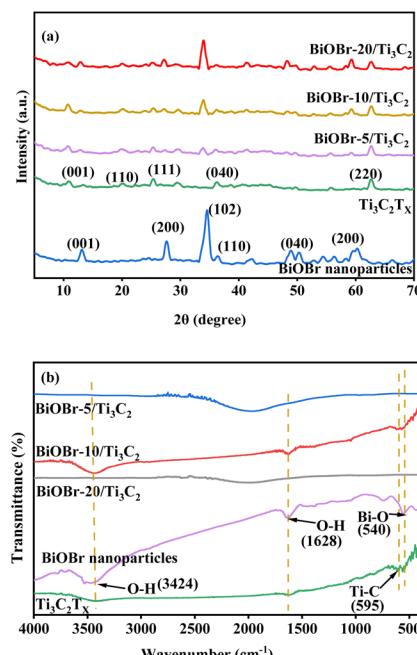


Fig. 2 (a) XRD patterns and (b) FT-IR spectra of BiOBr, $\text{Ti}_3\text{C}_2\text{T}_x$ and BiOBr- x / Ti_3C_2 .



vibration of the Bi–O bond.⁶⁰ In addition, the peaks observed at 1630 and 3430 cm^{-1} in the BiOBr- x /Ti₃C₂ composites correspond to O–H vibrations of adsorbed water molecules. The BiOBr- x /Ti₃C₂ composites also show Bi–O and Ti–C vibration peaks, indicating the simultaneous existence of Ti₃C₂T_x and BiOBr phases.

XPS was employed to investigate the surface elemental states of the photocatalysts, which provided evidence for the successful combination of BiOBr and Ti₃C₂T_x. XPS survey spectra and high-resolution XPS spectra of BiOBr, Ti₃C₂T_x and BiOBr- x /Ti₃C₂ samples are presented in Fig. 3a–f. In Fig. 3a, Ti, C, Bi, O, and Br are detected in the XPS spectra of BiOBr- x /Ti₃C₂ composites supporting 10 wt% and 20 wt% of BiOBr. In the Bi 4f spectrum shown in Fig. 3b, two distinct peaks with binding energies of 159.5 and 164.7 eV are assigned to Bi 4f_{5/2} and Bi 4f_{7/2}, respectively.⁶¹

In Fig. 3c, each O 1s spectrum can be deconvoluted into three peaks. The peak at 530.4 eV corresponds to lattice oxygen, while the peaks at 531.8 eV and 533.3 eV are attributed to oxygen vacancies (OVs) and chemisorbed oxygen (hydroxyl groups), respectively.⁶² In Fig. 3d, the peaks at 68.5 and 69.4 eV are associated with Br 3d_{5/2} and Br 3d_{3/2}, respectively.⁶¹ In Fig. 3e, the Ti 2p spectrum of each BiOBr- x /Ti₃C₂ sample can be deconvoluted into five peaks located at 454.7, 456.1, 459.2, 459.8 and 464.8 eV, corresponding to Ti–C, Ti³⁺ (Ti 2p_{3/2}), Ti–O (2p_{3/2}), Ti⁴⁺ (Ti 2p_{3/2}) and Ti–O (2p_{1/2}), respectively.^{58,59} In contrast, the Ti 2p spectrum of pure Ti₃C₂T_x exhibits only four peaks. This difference is primarily attributed to the peak at 459.2 eV, which

indicates the formation of Ti–O bonds between the O in BiOBr and the Ti in MXene Ti₃C₂T_x. In Fig. 3f, the C 1s spectrum is deconvoluted into four peaks at 281.6, 284.9, and 286.7 eV, corresponding to C–Ti bonds, sp² carbon C=C, and C–O bonds, respectively.⁵⁸ Compared to pure BiOBr, the Bi 4f, O 1s, and Br 3d peaks of the BiOBr- x /Ti₃C₂ samples exhibited a significant shift toward lower binding energies (Fig. 3b–d), while the Ti 2p peaks shifted toward higher binding energies (Fig. 3e). The observations indicate that electrons are gained during the oxidation process of BiOBr, leading to observed shifts in the binding energies of Bi, O, and Br to lower fields. Simultaneously, Ti and C in the composite material lose electrons, resulting in a shift to higher fields.

Photoelectric properties of materials

The photocurrent and EIS reflect the ability of photo-excited carriers to separate and transport under illumination. Fig. 4a shows a comparison of the photocurrent response of the BiOBr- x /Ti₃C₂ composites with different BiOBr loadings. Compared with the original Ti₃C₂T_x, the transient photocurrent response of a BiOBr/Ti₃C₂ sample was significantly increased, indicating that the interfacial charge transfer ability of Ti₃C₂T_x was promoted due to the formation of heterojunctions between the MXene Ti₃C₂T_x and BiOBr nanoparticles. Among the samples, BiOBr-10/Ti₃C₂ exhibited the highest photocurrent density owing to its highly dispersed BiOBr nanoparticles and laminated Ti₃C₂T_x layers (Fig. 1c). As a result, the sample can achieve more efficient photogenerated charge separation and transfer, which helps improve its photocatalytic performance. In addition, the photocurrent of each sample shows stability in intensity with time extension, indicating that the composites can provide a stable number of electrons and holes during the irradiation process and further exhibit stable photocatalytic activity. Fig. 4b shows the comparison of the EIS of BiOBr- x /Ti₃C₂ composites with different BiOBr loadings. As shown in the figure, the order of the capacitance arc semicircle for the samples is that Ti₃C₂T_x > BiOBr > BiOBr-5/Ti₃C₂ > BiOBr-20/

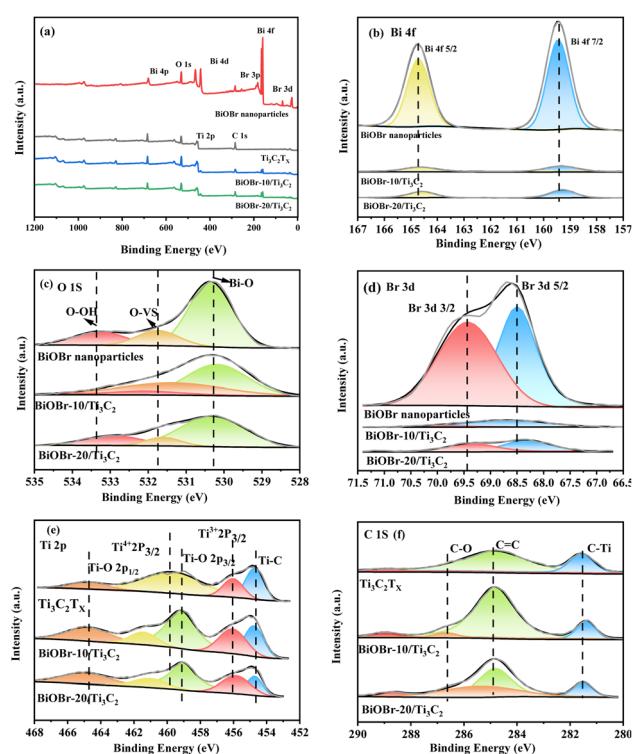


Fig. 3 (a) XPS survey spectra; high-resolution XPS spectra of (b) Bi 4f, (c) O 1s, (d) Br 3d, (e) Ti 2p and (f) C 1s for BiOBr, Ti₃C₂T_x and BiOBr- x /Ti₃C₂.

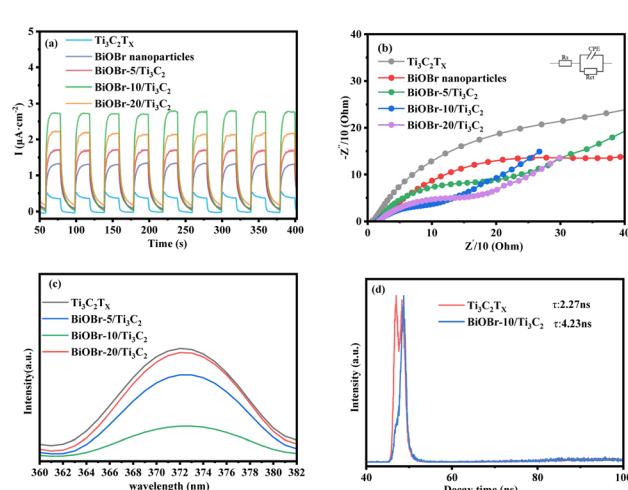


Fig. 4 (a) Transient photocurrent density, (b) Nyquist plots, (c) PL spectra and (d) TRPL spectra of Ti₃C₂T_x, BiOBr and BiOBr- x /Ti₃C₂.



$\text{Ti}_3\text{C}_2 > \text{BiOBr-10/Ti}_3\text{C}_2$. The smaller charge transfer impedance between $\text{Ti}_3\text{C}_2\text{T}_x$ layers and the BiOBr nanoparticles is conducive to the separation of $\text{e}^+ - \text{h}^+$ pairs in the photocatalytic process. The capacitance arc semicircle of BiOBr-10/Ti₃C₂ is the smallest, indicating that a more efficient interface between BiOBr nanoparticles and $\text{Ti}_3\text{C}_2\text{T}_x$ is constructed, which provides more space for the separation and transfer of $\text{e}^+ - \text{h}^+$ pairs and makes it have the best photocatalytic activity.

The state and the lifetime of active electron-hole pairs were further analyzed by PL and TRPL spectra, respectively. Fig. 4c shows the PL spectra of the photocatalyst samples. The pristine $\text{Ti}_3\text{C}_2\text{T}_x$ shows the strongest emission peak at 373 nm. The emission peak intensity of the BiOBr-x/Ti₃C₂ sample was lower than that of pristine $\text{Ti}_3\text{C}_2\text{T}_x$. In particular, BiOBr-10/Ti₃C₂ shows the weakest emission peak, indicating that appropriately loaded BiOBr nanoparticles reduced the speed of $\text{e}^+ - \text{h}^+$ pairs recombination in $\text{Ti}_3\text{C}_2\text{T}_x$. Fig. 4d shows the TRPL spectra of $\text{Ti}_3\text{C}_2\text{T}_x$ and BiOBr-10/Ti₃C₂. The average fluorescence lifetimes of $\text{Ti}_3\text{C}_2\text{T}_x$ and BiOBr-10/Ti₃C₂ were 2.27 and 4.23 ns, respectively, which were calculated by eqn (3). The longer the fluorescence lifetime, the more conducive it is to carrier separation, which is conducive to photogenerated carriers participating in the photocatalytic reaction. Based on the photocurrent, EIS, PL and TRPL spectra, it can be concluded that the interfacial electron transfer was improved after the formation of the BiOBr/Ti₃C₂ heterojunction.

$$\tau_a = (A_1\tau_1^2 + A_2\tau_2^2)/(A_1\tau_1 + A_2\tau_2) \quad (3)$$

where τ and A are the emission lifetimes of each component and amplitude, respectively.

Photocatalytic performance of roxarsone degradation

The photocatalytic activity of the samples was evaluated by the photocatalytic degradation of roxarsone under visible light. Fig. 5a shows a comparison of the photocatalytic degradation of roxarsone by different photocatalysts. Pure $\text{Ti}_3\text{C}_2\text{T}_x$ exhibits basal photocatalytic activity for roxarsone degradation, achieving 38.21% removal for 3 h, and $1.06 \text{ mg g}_{\text{cat}}^{-1} \text{ h}^{-1}$ of degradation efficiency at 0.5 h, which is almost twice as high as that of pure BiOBr. Compared with pure $\text{Ti}_3\text{C}_2\text{T}_x$, the BiOBr-x/Ti₃C₂ sample exhibits a significant enhancement in the photocatalytic activity for roxarsone degradation. The activity order of the BiOBr-x/Ti₃C₂ samples was BiOBr-10/Ti₃C₂ > BiOBr-5/Ti₃C₂ > BiOBr-20/Ti₃C₂. BiOBr-10/Ti₃C₂ as the optimal sample achieved 57.92% removal for 3 h, and $1.27 \text{ mg g}_{\text{cat}}^{-1} \text{ h}^{-1}$ of degradation efficiency at 0.5 h. The results indicate that modification of an appropriate amount (10 wt%) of BiOBr for $\text{Ti}_3\text{C}_2\text{T}_x$ can promote the separation of photogenerated carriers and the transfer of electrons to effectively participate in the photocatalytic reaction. However, BiOBr-20/Ti₃C₂ exhibited lower photocatalytic degradation ability for roxarsone than pure $\text{Ti}_3\text{C}_2\text{T}_x$. As discussed above, excess BiOBr in the composite results in the coverage of the active sites within the $\text{Ti}_3\text{C}_2\text{T}_x$ layers, which lowers the photogenerated carrier density and restrains the photoelectron transfer and the separation of $\text{e}^+ - \text{h}^+$ pairs. Fig. 5b shows the effect of medium pH on the photocatalytic degradation of

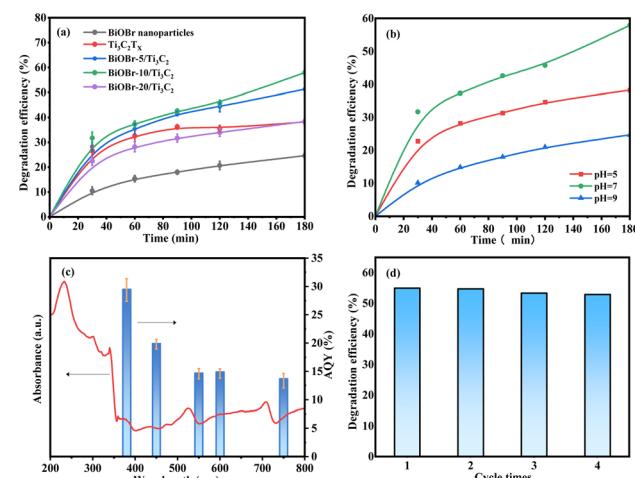


Fig. 5 (a) Photocatalytic degradation of roxarsone over different photocatalysts (conditions: 300 W Xe lamp, 0.05 g catalyst, 50 mL 2 mg L^{-1} of roxarsone solution, pH = 7.0); (b) effect of the medium's pH on photocatalytic degradation of roxarsone over BiOBr-10/Ti₃C₂ (conditions: 300 W Xe lamp, 0.05 g catalyst, 50 mL 2 mg L^{-1} of roxarsone solution); (c) UV-vis absorption spectrum of BiOBr-10/Ti₃C₂ (left axis) and wavelength dependent AQY (%) of photocatalytic degradation of roxarsone (right axis); and (d) recycling ability of BiOBr-10/Ti₃C₂ (conditions: 300 W Xe lamp, 0.05 g catalyst, 50 mL 2 mg L^{-1} of roxarsone solution, pH = 7.0, 3 h).

roxarsone. The most efficient degradation of roxarsone can be achieved in a neutral environment. Acidic condition inhibits the immobilization of As^{5+} products, both reducing the degradation efficiency and increasing the risk of secondary pollution. Under alkaline conditions, As^{5+} tends to form soluble arsenate with OH^- , and the active sites of photocatalysts are poisoned, enhancing arsenic migration. Fig. 5c shows the AQY values for the photocatalytic degradation of roxarsone by BiOBr-10/Ti₃C₂ at different wavelengths. The AQY values at 380, 450, 550, 600, and 750 nm were calculated to be 29.5%, 19.9%, 14.7%, 14.9%, and 13.7%, respectively. The trend of the AQY value agreed well with the diffuse reflectance spectrum. Fig. 5d shows the recyclability of BiOBr-10/Ti₃C₂ for the photocatalytic degradation of roxarsone. The spent photocatalyst in each cycle was sufficiently washed with water and ethanol. After treatment, the photocatalyst exhibited good recyclability and could be reused for 4 cycles.

Photocatalytic mechanism

UV-vis diffuse reflectance spectra were used to investigate the band structure of the BiOBr/Ti₃C₂ heterojunction, as shown in Fig. 6a. Pure BiOBr shows absorption bands at 231 and 358 nm, and the absorption range does not exceed 450 nm, which limits the use of the visible part of solar energy. Pure $\text{Ti}_3\text{C}_2\text{T}_x$ exhibits stronger absorption in the ultraviolet region (200–400 nm) and reduced absorption in the visible region (500–800 nm). With the increase of BiOBr content, the absorption strength of the BiOBr-x/Ti₃C₂ composite is increased in the visible region, indicating that BiOBr combined with $\text{Ti}_3\text{C}_2\text{T}_x$ effectively improves the light absorption range and ability of MXene. Based on Fig. 6a, the



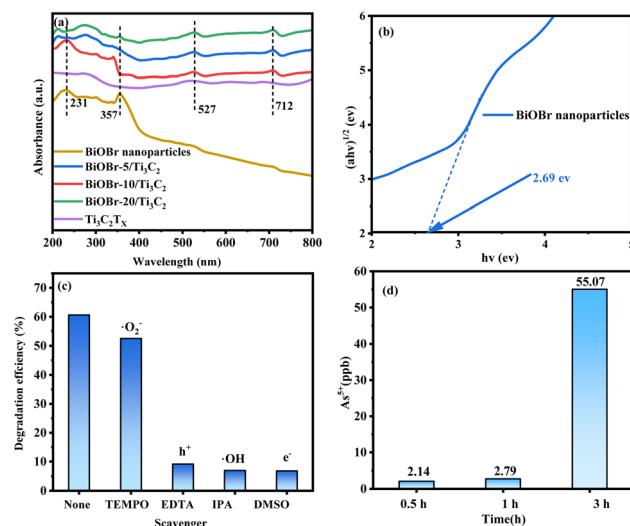
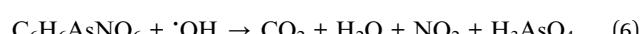
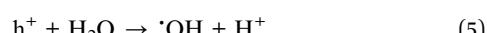
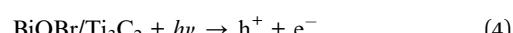


Fig. 6 (a) UV-vis absorption spectra of BiOBr, Ti₃C₂T_x and BiOBr-x/Ti₃C₂; (b) Kubelka-Munk plot of BiOBr; (c) photocatalytic degradation of roxarsone over BiOBr-10/Ti₃C₂ (conditions: 300 W Xe lamp, 0.05 g catalyst, 50 mL 2 mg L⁻¹ of roxarsone solution, pH = 7, 1 h); (d) final product detection for roxarsone degradation at different times (conditions: 300 W Xe lamp, 0.05 g catalyst, 50 mL 2 mg L⁻¹ of roxarsone solution, pH = 7.0).

bandgaps (E_g) of the samples were determined by the transformation of the Kubelka-Munk function to the UV-vis diffuse reflectance.^{58,63} Fig. 6b shows the $(ahv)^{1/2}$ vs. (hv) plot for BiOBr nanoparticles. The bandgap of BiOBr was estimated to be 2.69 eV. The values of $E(\text{H}_2\text{O}/\cdot\text{OH})$ and $E(\text{O}_2/\cdot\text{O}_2^-)$ were 2.8 and -0.33 eV,⁶⁴ respectively. In addition, the E_{VB} position of BiOBr was approximately 3.02 eV, whereas the E_{CB} position was not more negative than the $E(\text{O}_2/\cdot\text{O}_2^-)$ value.⁶⁵ Therefore, $\cdot\text{OH}$ was significantly generated and contributed to the redox reaction instead of $\cdot\text{O}_2^-$.

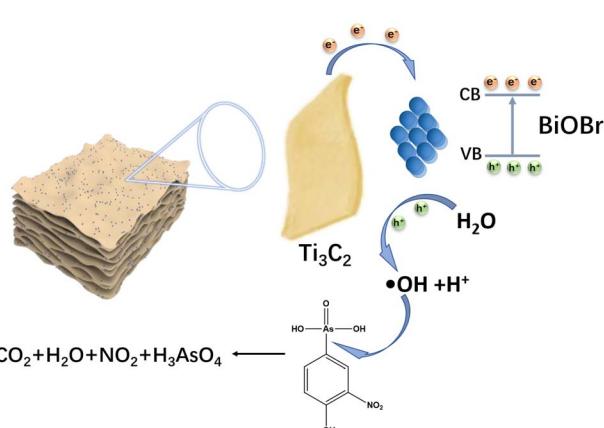
To elucidate the mechanism of the photocatalytic degradation of roxarsone, active species capture experiments were performed with EDTA, DMSO, IPA and TEMPOL as scavengers

of h^+ , e^- , $\cdot\text{OH}$ and $\cdot\text{O}_2^-$, respectively, as shown in Fig. 6c. The $\cdot\text{OH}$ and h^+ scavengers (DMSO, EDTA) had obvious inhibitory effects on the degradation of roxarsone, indicating that $\cdot\text{OH}$ is an active substance essential for the degradation of roxarsone, while h^+ plays an important role in $\cdot\text{OH}$ generation. The inhibition by $\cdot\text{O}_2^-$ scavenger (TEMPOL) was very weak, indicating that $\cdot\text{O}_2^-$ is not the main active substance for the photocatalytic degradation of roxarsone. Although the scavenger (DMSO) has the most remarkable inhibitory effect, the electrons only contribute to the separation of $\text{e}^-\text{-h}^+$ pairs to produce more holes for roxarsone degradation, instead of direct participation in the reaction. The residual inorganic arsenic forms in the liquid phase were confirmed by LC-ICP-MS to analyze the reaction details and assess the safety after water treatment. As shown in Fig. 6d, As⁵⁺, as the final product of roxarsone, can be detected after each reaction period, and its concentration increases with the reaction time. From the residual roxarsone concentration (2–50 ppb, Fig. 6d), it can be assessed that the treated water by the present system was at a safe level, according to the specified level (0.01 mg L⁻¹) by the World Health Organization (WHO). Based on these findings, we propose the mechanism for the photocatalytic degradation of roxarsone over BiOBr/Ti₃C₂, as illustrated by eqn (4)–(6) and Scheme 2. Upon absorption of light energy by BiOBr/Ti₃C₂, electrons are excited from the VB to the CB, which generates abundant holes in the VB of the photocatalyst (eqn (4)). The holes oxidize water molecules to generate abundant hydroxyl radicals ($\cdot\text{OH}$) (eqn (5)).⁶⁶ Finally, $\cdot\text{OH}$ participates in the mineralization and oxidation of roxarsone to form inorganic products (eqn (6)).



Conclusions

In summary, BiOBr nanoparticles were successfully loaded onto MXene Ti₃C₂T_x to form a heterostructure for the photocatalytic degradation of roxarsone. The proposed construction strategy effectively addressed the issue of low photocatalytic efficiency of Ti₃C₂T_x in environmental protection areas. The uniform dispersion of BiOBr nanoparticles on the surface of Ti₃C₂T_x layers promoted the formation of an interface between the two phases, significantly enhancing the electron transport capability, electron donor density, and the separation efficiency of electron–hole pairs. The optimized BiOBr/Ti₃C₂ material exhibited excellent photocatalytic performance for roxarsone degradation, along with an AQY of 29.5% and good reusability for at least 4 cycles. A series of in-depth mechanism studies proposed that h^+ and $\cdot\text{OH}$ in the VB of the BiOBr/Ti₃C₂ system are involved in the main pathway of the oxidative mineralization of roxarsone. This work is expected to elucidate more feasible and scalable photocatalytic systems for organoarsenic-



Scheme 2 Proposed mechanism for the photocatalytic degradation of roxarsone over BiOBr/Ti₃C₂.



contaminated wastewater treatment with high efficiency and sustainability.

Data availability

Data will be made available upon request.

Author contributions

Yixin Guo: methodology, data curation, software, and writing – original draft. Ya-Nan Wang: formal analysis and methodology. Jinsong Peng: project administration. Haiyan Song: project administration, methodology, data curation, writing – original draft, and writing – review & editing. Chunxia Chen: project administration and methodology.

Conflicts of interest

There are no conflicts to declare.

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