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# Construction of a crossed-layer-structure MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> heterojunction with enhanced photocatalytic performance†

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A novel crossed-layer-structure MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> (graphitic carbon nitride) was synthesized by a facile method, and was characterized by a collection of analytical techniques: X-ray diffraction patterns, FT-IR spectra, SEM, TEM, and XPS. The crossed-layer-structure morphologies of MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> could be readily tailored by adjusting the molar ratio of MoS<sub>2</sub> to dicyandiamide; with an increase in the molar ratio from 0.08% to 4%, a morphology transformation was observed. It can be used as a photocatalyst in degrading methyl orange under simulated solar light irradiation. These MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> samples demonstrated 2 to 3 times higher performance than pure g-C<sub>3</sub>N<sub>4</sub>. We also found that the highest phenol degradation activity was with MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-4 under visible light. The high performance was attributed to the unique morphology that provided low recombination losses of photogenerated charges.

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## 1. Introduction

Due to the global energy crisis and environmental pollution,<sup>1–5</sup> various technologies have been used to cope with the crisis. Photocatalysis is considered to be one of the most promising technologies to produce renewable fuels and remove pollutants *via* solar energy. Photocatalytic degradation of pollutants in water by using semiconductor catalysts has received extensive and increasing attention in the past few decades, and various kinds of catalyst suitable for use in natural light have been developed.<sup>6,7</sup>

The key to achieving solar pollution degradation is to develop stable, efficient, nontoxic and inexpensive photocatalysts that are capable of working in the visible light, which occupies *ca.* 50% of the incoming solar irradiation on earth. In the last few years, a graphite-like organic polymeric carbon nitride (g-C<sub>3</sub>N<sub>4</sub>), a metal-free photoactive semiconductor consisting of abundant and nontoxic elements, has ignited increasing interest owing to its photocatalytic activity for degradation of pollutants in water under visible-light irradiation, and its cheap, stable, environmental friendly, metal-free features. In fact, g-C<sub>3</sub>N<sub>4</sub>, one of the oldest synthetic polymers, can be traced back to 1834, and reported for the first time by Liebig and named “melon”.<sup>8</sup> Unlike graphite, g-C<sub>3</sub>N<sub>4</sub> is a semiconductor material with an optical band gap of 2.7 eV.<sup>9</sup>

Recently, g-C<sub>3</sub>N<sub>4</sub> derivatives with various heteroatom dopants (*e.g.* C, B, F, S, P, O, I),<sup>10–17</sup> have been fabricated by co-thermal condensation of suitable additives with the precursor of g-C<sub>3</sub>N<sub>4</sub>. In comparison, post-functionalization of g-C<sub>3</sub>N<sub>4</sub> has 40 extra advantages of introducing much more functional groups or dopants into g-C<sub>3</sub>N<sub>4</sub> after the lattice is formed at elevated temperatures. However, the high chemical stability and bad solubility in common solvents of g-C<sub>3</sub>N<sub>4</sub> constitute a vast barrier to post-modify g-C<sub>3</sub>N<sub>4</sub> material.<sup>18</sup> Great efforts have been made to enhance the photocatalytic activity of g-C<sub>3</sub>N<sub>4</sub>, such as doping,<sup>19–23</sup> copolymerization,<sup>18,24</sup> texturization,<sup>25,26</sup> supermolecular assembly,<sup>27,28</sup> surface heterojunction design.<sup>29</sup> Highly condensed g-C<sub>3</sub>N<sub>4</sub> with fewer defects favors enhanced the photocatalytic performance by increasing the crystallinity and charge carrier mobility.<sup>30</sup> Therefore, it is urgent for us to obtain highly efficient g-C<sub>3</sub>N<sub>4</sub> with enhanced crystallinity and charge carrier separation efficiency in a novel way.<sup>31</sup> To date, various nanoarchitectural g-C<sub>3</sub>N<sub>4</sub>, ranging from one-dimensional nanorods and nanowires to two-dimensional nanosheets and three-dimensional mesoporous structures, have been designed by many methods and strategies, such as hard/soft template synthesis,<sup>32–34</sup> solvothermal/molten-salt technology,<sup>35,36</sup> ods,<sup>37,38</sup> and supramolecular chemistry.<sup>27,39</sup>

Transition-metal dichalcogenides (TMDs), especially the ones with atomic thickness, have been regarded as a new class of nanomaterials for fundamental studies and promising applications because of their special properties. TMDs can be metals, semiconductors, or insulators,<sup>40–42</sup> exhibiting diverse properties. MoS<sub>2</sub> is one of the most promising semiconductors owing to its inherent and thickness-dependent band gap as well as its abundance in nature as molybdenite.<sup>43</sup> Moreover, bulk MoS<sub>2</sub> crystals can be exfoliated to single- or few-layer

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nanosheets, which exhibit unusual physical and electronic properties. Specifically, bulk MoS<sub>2</sub> is a semiconductor with narrow band gap of about 1.3 eV, while the isolated MoS<sub>2</sub> monolayer possesses a large band gap of 1.8–1.9 eV.<sup>43</sup>

It is worth noting that the MoS<sub>2</sub> nanosheets,<sup>44</sup> as an electron acceptor, can efficiently increase the visible-light absorbance, charge separation, specific surface area and active sites for reaction, which result in an enhanced photocatalysis ability. MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> heterojunction photocatalyst has been synthesized, using basically layered self-assembly technology and the impregnation method.<sup>45–53</sup> In this paper, a facile method, impregnation–calcination, was applied to synthesize crossed-layer-structure MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>. The activities of as-prepared photocatalysts were evaluated by photodegradation of methyl orange (MO) under simulated solar light illumination. The as-fabricated MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> heterojunction photocatalyst exhibited enhanced photoreactivity compared with the pure g-C<sub>3</sub>N<sub>4</sub>.

## 2. Experimental section

### 2.1 Reagents

Dicyandiamide (C<sub>2</sub>H<sub>4</sub>N<sub>4</sub>, CP), isopropanol (IPA, AR), sodium molybdate (Na<sub>2</sub>MoO<sub>4</sub>, CP), thiourea (CN<sub>2</sub>H<sub>4</sub>S, CP), disodium ethylenediaminetetraacetate dehydrate (EDTA-2Na, AR), 1,4-benzoquinone (BQ, AR), methyl orange (MO, AR), and other chemicals involved were purchased from Sinopharm Chemical Reagent Co., Ltd. (China) and used directly for experimental without further purifications. All aqueous solutions throughout were prepared with the deionized water.

### 2.2 Catalysts preparation

**2.2.1 Preparation of bulk g-C<sub>3</sub>N<sub>4</sub>.** Graphitic carbon nitride was synthesized using dicyandiamide. In a typical run, the precursor was put in a lidded high quality alumina crucible, then placed inside a tube furnace and heated at 5 °C minute<sup>-1</sup>, and finally held at 550 °C for 4 hours in Ar flow. The resultant powders were then washed with water, HCl, and once again with water to remove all unreacted and potentially detrimental surface species. The products were denoted as g-C<sub>3</sub>N<sub>4</sub>.

**2.2.2 Preparation of MoS<sub>2</sub> sheets.** Bulk MoS<sub>2</sub> (BMS) was synthesized by the hydrothermal method. 242 mg sodium molybdate and 342.5 mg thiourea were dissolved in 60.0 mL distilled water and further ultrasonicated for 1 h, and then transferred into a Teflon-lined stainless steel autoclave. The autoclave was heated in the oven at 200 °C for 24 hours and cooled to room temperature naturally. The as-obtained product was washed with water and ethanol, and precipitated by centrifuge with a speed of 10 000 rpm. Then dried in an oven at 60 °C for 24 h. 1 g bulk MoS<sub>2</sub> powder was put into a 100 mL glass bottle and ultrasonicated for 48 h, then put the suspension 4 mg mL<sup>-1</sup>.

**2.2.3 Preparation of crossed-layer-structure MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> heterojunction photocatalyst.** In a typical synthesis of MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> hybrid, 10 g of dicyandiamide and 0.5 (1, 2.5, 5, 12.5, 25) mL MoS<sub>2</sub> suspension were immersed into 50 mL aqueous solution. These were then mixed with stirring and

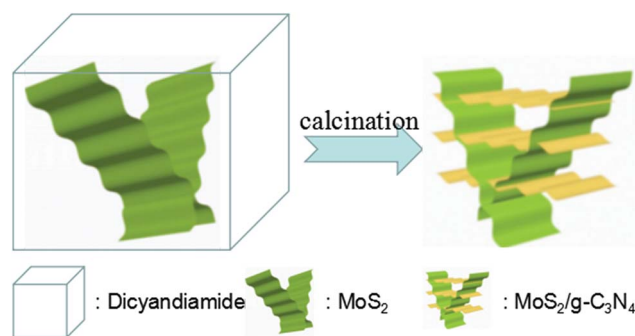
ultrasonication for 60 min as-prepared precursor was obtained after evaporation and drying in an oven at 60 °C for 24 h. Then calcination of these as-prepared precursor stabilized at 550 °C for 4 h with a ramp rate of 5 °C min<sup>-1</sup> in Ar flow from room temperature, as shown in Scheme 1. The resultant powders were then washed with water, HCl, and once again with water to remove all unreacted and potentially detrimental surface species. The obtained samples were designated as MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-2, MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-4, MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-10, MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-20, MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-50 and MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-100 respectively.

### 2.3 Catalysts characterization

X-ray diffraction (XRD) of the samples was carried out on a Bruker AXS D8 X-ray diffractometer with a Cu-Kα X-ray source operating at 40 kV and 100 mA. The morphologies of the samples were observed using a scanning electron microscope (SEM, JEOL JSM-6490LV) and a transmission electron microscope (TEM, FEI Tecnai G2). X-ray photoelectron spectroscopic (XPS) measurements were made on an Escalab 250Xi system. UV-vis diffuse reflection spectra of the as-prepared products were carried out using an Evolution 220 UV-vis spectrophotometer (Thermo Fisher) from 200 to 800 nm. Fourier transform infrared (FT-IR) spectra of the samples were measured by a Nicolet 5700 Fourier transform infrared spectrometer. The photoluminescence spectra (PL) of the samples were obtained using a fluorescence spectrometer (Hitachi F-4500) at 293 K. The photocurrent curves and electrochemical impedance spectroscopy (EIS) plots were collected by a CHI660 electrochemical analyzer.

### 2.4 Photocatalytic experiments

The activity of the photocatalysts was evaluated *via* the photocatalytic degradation of MO under simulated solar light irradiation. In a typical experiment, 0.05 g photocatalyst was suspended in 100 mL 10 mg L<sup>-1</sup> MO solution. The suspension was first dispersed by sonication for 30 min and stirred for 30 min in the dark to obtain adsorption–desorption equilibrium between the MO and the photocatalyst. The suspension was then stirred and irradiated under a 500 W Xe lamp. During the irradiation process, about 3 mL of suspension was taken from the reaction cell every 30 min and centrifuged to remove the photocatalyst. The absorbance of the MO solution in degradation was detected on TU-1900 UV-vis spectrophotometer.



Scheme 1 Schematic illustration for the preparation of crossed-layer-structure MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> heterojunction photocatalyst.



## 2.5 Electrochemical measurements

Photoelectrochemical measurements were performed in a three-electrode, single compartment quartz cell on an electrochemical station (CHI 660). Samples on ITO glass with an active area of *ca.* 1.0 cm<sup>2</sup> (1 mg of photocatalyst) were prepared as the working electrode. The platinum sheet and saturated calomel electrode (SCE) were served as the counter electrode and reference electrode, respectively. Besides, a bias voltage of 0.5 V was used for driving the photo-generated electrons transfer from the working electrode to the platinum electrode. The light source (300 W xenon lamp) with ultraviolet filter ( $\lambda > 400$  nm) was mounted 10.0 cm away from the photoelectrochemical cell. A 0.50 M Na<sub>2</sub>SO<sub>4</sub> aqueous solution worked as the electrolyte.

## 3. Results and discussion

The XRD patterns of the as-prepared pure g-C<sub>3</sub>N<sub>4</sub>, MoS<sub>2</sub>, and MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> nanocomposites were shown in Fig. 1. These two peaks at 13.057° and 27.76° matched well with the (100) and (002) crystal planes of graphitic carbon nitride. This result

demonstrate that the synthesized samples were graphitic carbon nitride. The small angle peak at 13.057°, which corresponded to a *d*-spacing of 0.677 nm, was due to the in-plane structural packing motif. The strong peak at 27.76° was a characteristic interlayer stacking peak and corresponded to an interlayer distance of *d* = 0.321 nm, as shown in Fig. 1a. Existing two diffraction peaks at 12.91° and 14.17° corresponded to g-C<sub>3</sub>N<sub>4</sub> (100) peak and MoS<sub>2</sub> (002) peak in MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-50 and MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-100, however no distinct diffraction peaks corresponding to other MoS<sub>2</sub> co-catalysts could be observed. This might be due to the small amount of MoS<sub>2</sub> contents, having small amount of MoS<sub>2</sub> contents in the surface of g-C<sub>3</sub>N<sub>4</sub> and highly dispersion in the interior of polymeric g-C<sub>3</sub>N<sub>4</sub> photocatalysts. The presence of MoS<sub>2</sub> in the MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> composite samples could be confirmed by TEM, SEM and XPS analyses, as discussed later. From the inset, the (002) peak of MoS<sub>2</sub> co-catalysts shifted to left comparing with pure g-C<sub>3</sub>N<sub>4</sub>.<sup>51</sup> Moreover, the diffraction peaks at 14.0°, 33.4° and 58.7° were in good agreement with the (002), (100) and (110) planes of the hexagonal phase of MoS<sub>2</sub> in Fig. 1b.

The structural information of the as-developed g-C<sub>3</sub>N<sub>4</sub> and MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> samples were confirmed by the FT-IR analysis (Fig. 2). FT-IR spectrum showed the presence of polar oxygen-containing functionalities at 1062 cm<sup>-1</sup> (C–O–C stretching) and 1205 cm<sup>-1</sup> (phenolic C–OH stretching). Further observation, the characteristic peaks of MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> were found to be almost identical to the pure g-C<sub>3</sub>N<sub>4</sub>, inferring that the impregnation of MoS<sub>2</sub> did not destroy the in-plane tri-s-triazine units. The MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> sample revealed almost similar characteristic features to the pure g-C<sub>3</sub>N<sub>4</sub>, verifying that the structural integrity of g-C<sub>3</sub>N<sub>4</sub> remained intact after the incorporation with MoS<sub>2</sub>. Several strong absorption bands in the range of 1200–1650 cm<sup>-1</sup> was originated from the skeletal stretching of C–N heterocycles with peaks positioned at 1632, 1573, 1423, 1329 and 1245 cm<sup>-1</sup>, comprising both trigonal N–(C)<sub>3</sub> (full condensation) and bridging C–NH–C units (partial condensation). This exemplified the successful development of the extended C–N–C

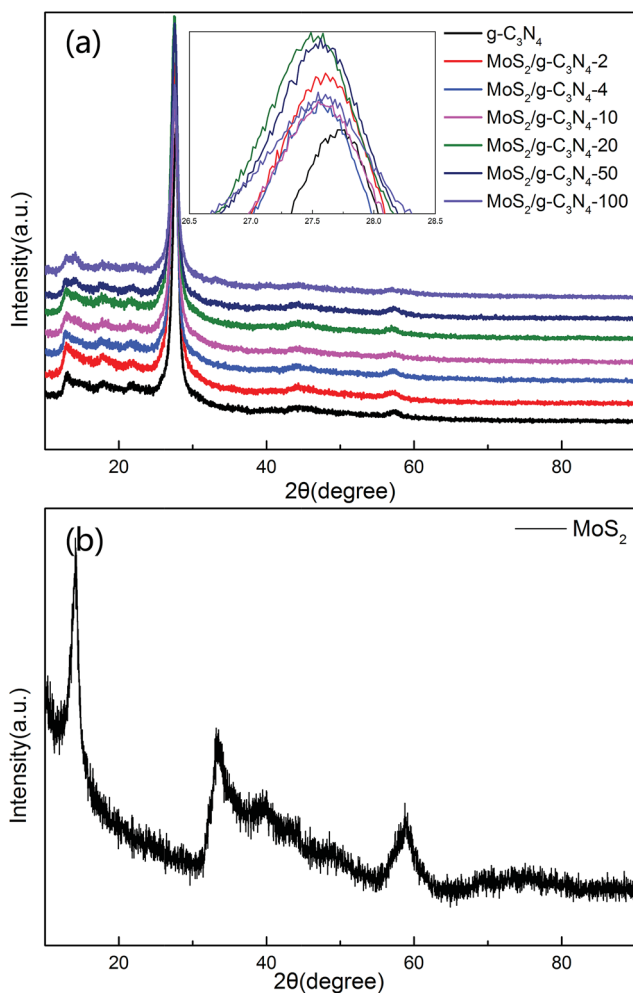


Fig. 1 XRD patterns of different photocatalyst: (a) pure g-C<sub>3</sub>N<sub>4</sub> and a series of MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> (b) MoS<sub>2</sub>.

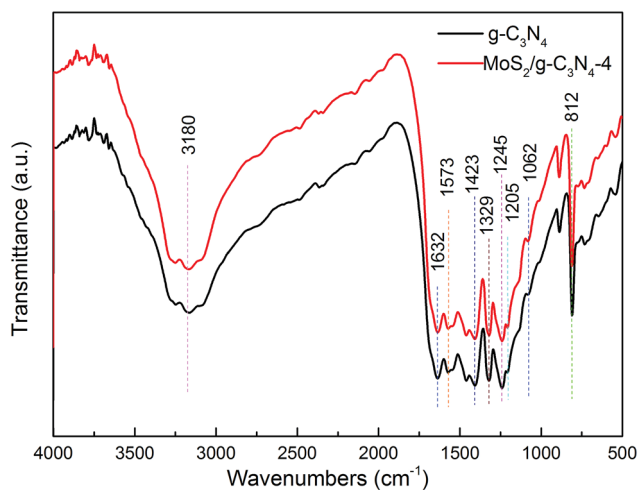


Fig. 2 FT-IR spectra of pure g-C<sub>3</sub>N<sub>4</sub> and MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-4 heterojunction photocatalyst.



network. The broad peak ranging from 3000 to 3700  $\text{cm}^{-1}$  was assigned to the N-H and O-H stretches due to the free amino groups and adsorbed hydroxyl species, respectively, whereas the sharp band at 812  $\text{cm}^{-1}$  was derived from the breathing vibration of tri-s-triazine units.<sup>54</sup> The virtually identical X-ray diffraction patterns (Fig. 1) and FT-IR spectra (Fig. 2) of  $g\text{-C}_3\text{N}_4$  and  $\text{MoS}_2/g\text{-C}_3\text{N}_4$  reveal that loading with  $\text{MoS}_2$  does not change the bulk structure of  $g\text{-C}_3\text{N}_4$ .

XPS of  $\text{MoS}_2/g\text{-C}_3\text{N}_4$  samples were measured on an Escalab 250Xi system and shown in Fig. 3. However, the C/N atomic ratio increased from 0.77 for  $g\text{-C}_3\text{N}_4$  to 0.84 for  $\text{MoS}_2/g\text{-C}_3\text{N}_4$ -4 (Fig. 3a). There were mainly two carbon species presented in the samples (Fig. 3b): one (284.6 eV) was  $\text{sp}^2$  C-C bonds, and the other one (288.0 eV) was  $\text{sp}^2$ -hybridized carbon in N-containing aromatic ring (N-C=N). The latter was considered as the major carbon species in  $\text{C}_3\text{N}_4$  layer. The high resolution N1s spectra could be also deconvoluted into three different peaks at binding energies of 398.7, 400.2, 401.3 eV, respectively (Fig. 3c). These peaks could be ascribed to the  $\text{sp}^2$ -hybridized nitrogen involved in triazine rings (C-N=C), the tertiary nitrogen N-(C)<sub>3</sub> groups, and the free amino groups (C-N-H), respectively. A weak O1s peak at 532.1 eV in the  $\text{MoS}_2/g\text{-C}_3\text{N}_4$ -4 was attributed to the adsorbed  $\text{H}_2\text{O}$  or  $\text{CO}_2$ , which was a common phenomenon found in literatures (Fig. 3d).<sup>55</sup> For Mo3d spectra, two peaks, accredited to the doublet  $\text{Mo}3\text{d}_{5/2}$  and  $\text{Mo}3\text{d}_{3/2}$ , were located at 229.4 and 232.5 eV (Fig. 3e). The other peak, corresponding to the  $\text{S}2\text{p}_{3/2}$  orbital of divalent sulfide ions ( $\text{S}^{2-}$ ), was observed at 162.3 eV (Fig. 3f). Those results indicate the existence of  $\text{Mo}^{4+}$

and  $\text{S}^{2-}$ , with an atomic composition ratio for Mo and S of 1 : 2. Of course, existing two peaks in the Fig. 5C corresponding to  $\text{S}2\text{p}_{3/2}$  and  $\text{S}2\text{p}_{1/2}$  orbital of tetravalent sulphide ions ( $\text{S}^{4-}$ ), were observed at 167.5 and 168.6 eV.<sup>56</sup> This is contributed to high-temperature calcination, which make divalent sulfide ions ( $\text{S}^{2-}$ ) be reduced to tetravalent sulfide ions ( $\text{S}^{4-}$ ), and produce a lot sulfide vacancy, which lead to great improvement of catalyst activity.

Transmission electron microscopy (TEM) image were showed in Fig. 4. Compared with pure  $g\text{-C}_3\text{N}_4$  (Fig. 4c), the as-prepared  $\text{MoS}_2/g\text{-C}_3\text{N}_4$  samples with laminar morphology exhibited nearly transparent feature with black lines and black cycles, indicating its crossed-layer-structure (Fig. 4a and b). Transparent layer structure of  $\text{MoS}_2$  was observed (Fig. 4d), indicating that the black lines and black cycles in  $\text{MoS}_2/g\text{-C}_3\text{N}_4$  samples (Fig. 4a and b) were  $\text{MoS}_2$ . The morphology of crossed-layer-structure  $\text{MoS}_2/g\text{-C}_3\text{N}_4$  was also investigated *via* scanning electron microscopy (SEM). As shown in Fig. 5a and b, crossed-layer-structure  $\text{MoS}_2/g\text{-C}_3\text{N}_4$  samples still maintained loose and irregular cross-like 3D morphology, although a great portion of  $g\text{-C}_3\text{N}_4$  nanosheets was stacked. Its morphology was quite different from that of the pure  $g\text{-C}_3\text{N}_4$  with a typical layer structure stacked layer by layer (Fig. 5c) and  $\text{MoS}_2$  with sheet structure (Fig. 5d). To further verify the crossed-layer-structure, EDX elements mappings were carried out and the results were shown in Fig. 6. Elements mapping of  $\text{MoS}_2/g\text{-C}_3\text{N}_4$ -4 reveals that the sample mainly contains five elements (C, N, O, S and Mo), which is coincident with the results from XPS (Fig. 3). Of course, the results also confirmed further the existence of the crossed-layer-structure.

The BET surface areas and porous structures of the photocatalysts were investigated by nitrogen adsorption-desorption. Fig. 7a and b showed the nitrogen adsorption-desorption isotherms and the corresponding pore-size distribution curves of pure  $g\text{-C}_3\text{N}_4$  and  $\text{MoS}_2/g\text{-C}_3\text{N}_4$  heterojunction catalysts ( $\text{MoS}_2/g\text{-C}_3\text{N}_4$ -2,  $\text{MoS}_2/g\text{-C}_3\text{N}_4$ -4 and  $\text{MoS}_2/g\text{-C}_3\text{N}_4$ -50). It was obvious that the nitrogen adsorption-desorption isotherms for

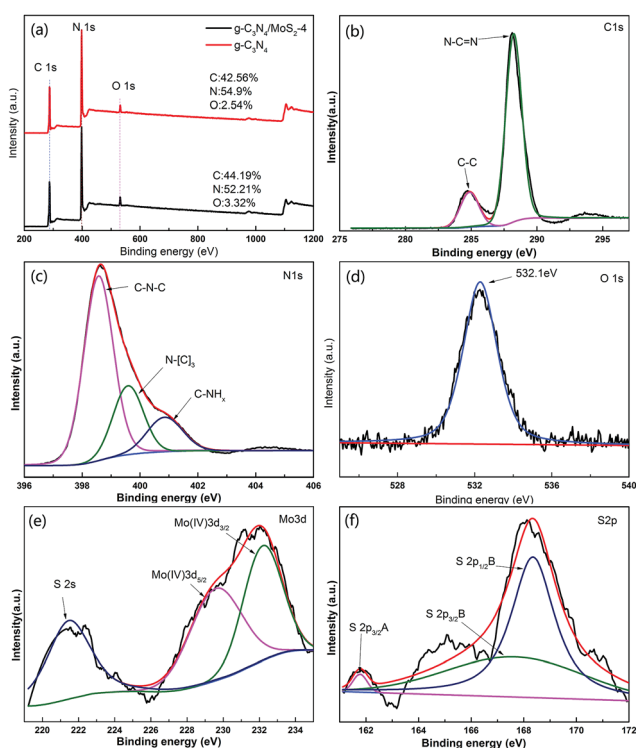


Fig. 3 XPS spectra of as-fabricated photocatalysts: (a) survey of  $g\text{-C}_3\text{N}_4$  and  $\text{MoS}_2/g\text{-C}_3\text{N}_4$ -4 and (b) C1s; (c) N1s; (d) O1s; (e) S2p; (f) Mo3d of  $\text{MoS}_2/g\text{-C}_3\text{N}_4$ -4.

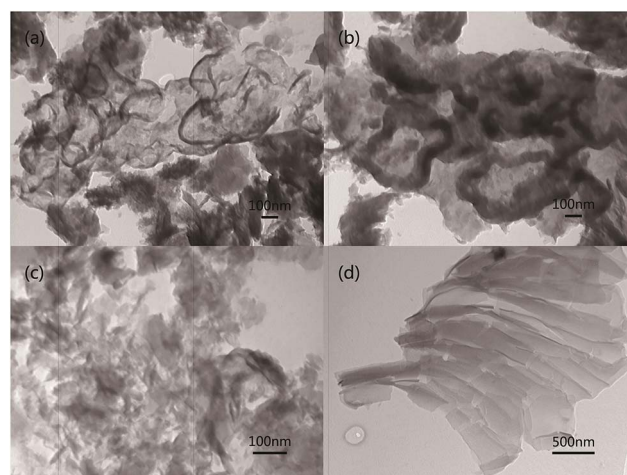


Fig. 4 Typical TEM images of (a)  $\text{MoS}_2/g\text{-C}_3\text{N}_4$ -4, (b)  $\text{MoS}_2/g\text{-C}_3\text{N}_4$ -100, (c)  $g\text{-C}_3\text{N}_4$ , (d)  $\text{MoS}_2$  samples.



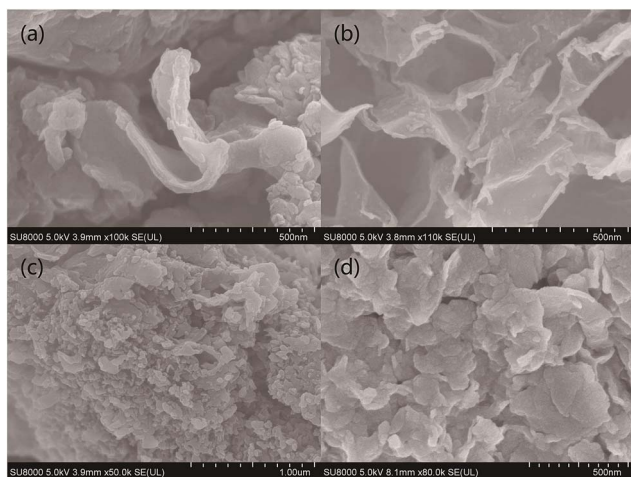


Fig. 5 Typical SEM images of (a)  $\text{MoS}_2/\text{g-C}_3\text{N}_4\text{-4}$ , (b)  $\text{MoS}_2/\text{g-C}_3\text{N}_4\text{-100}$ , (c) pure  $\text{g-C}_3\text{N}_4$  and (d)  $\text{MoS}_2$  samples.

pure  $\text{g-C}_3\text{N}_4$  and  $\text{MoS}_2/\text{g-C}_3\text{N}_4$  heterojunction catalysts belonged to type IV, indicating the presence of macropores. The increase of the surface areas was accompanied by the increase of the pore volumes, resulting from the increased size of the generated pores as suggested by the slight shift of the hysteresis onset in the corresponding nitrogen sorption isotherms to higher  $P/P_0$  as well as by the results of pore size analysis (Fig. 7a and b). Pure  $\text{g-C}_3\text{N}_4$  catalyst had a Brunauer–Emmett–Teller (BET) surface area of  $34.38 \text{ m}^2 \text{ g}^{-1}$  and a pore volume of  $0.077 \text{ cm}^3 \text{ g}^{-1}$ . While the BET surface areas and pore volumes of  $\text{MoS}_2/\text{g-C}_3\text{N}_4$  samples

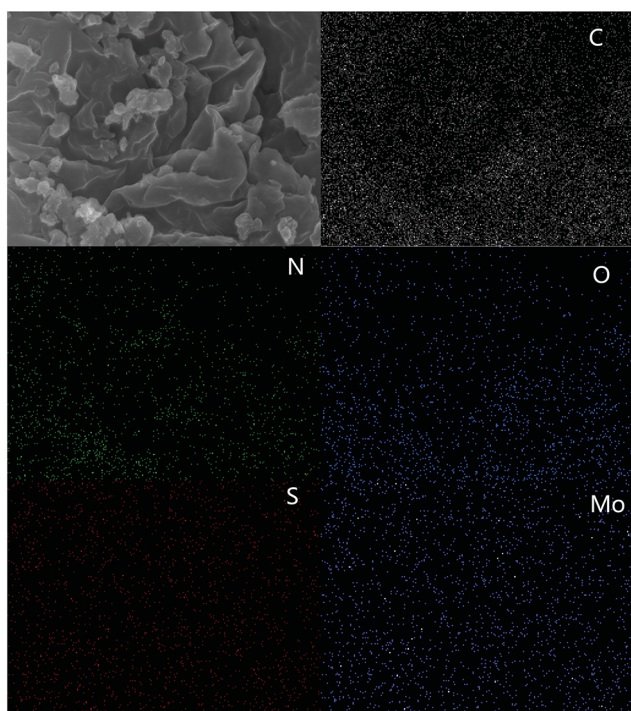


Fig. 6 SEM images and corresponding mapping images of  $\text{MoS}_2/\text{g-C}_3\text{N}_4\text{-4}$ .

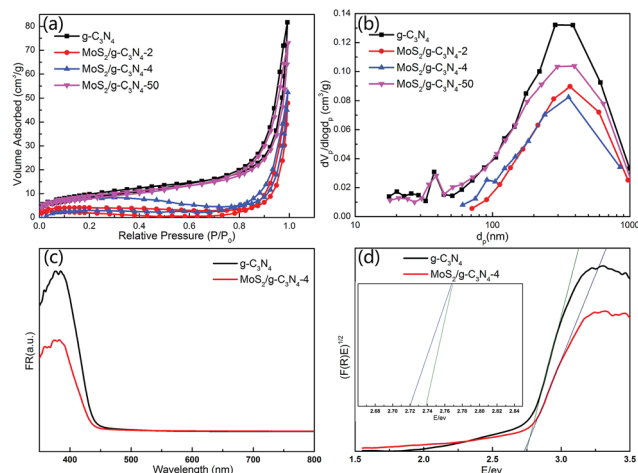


Fig. 7 (a)  $\text{N}_2$  adsorption isotherms and (b) pore size distribution of the  $\text{g-C}_3\text{N}_4$  and  $\text{MoS}_2/\text{g-C}_3\text{N}_4$ . (c) UV-vis spectra of  $\text{g-C}_3\text{N}_4$  and  $\text{MoS}_2/\text{g-C}_3\text{N}_4\text{-4}$  composites. (d) The plot of  $(F(R)E)^{1/2}$  versus energy ( $E$ ) for the band gap energy of  $\text{g-C}_3\text{N}_4$  and  $\text{MoS}_2/\text{g-C}_3\text{N}_4\text{-4}$ .

were less than pure  $\text{g-C}_3\text{N}_4$ . For  $\text{MoS}_2/\text{g-C}_3\text{N}_4$  samples, with the mass of  $\text{MoS}_2$  increasing, the BET surface areas and pore volume were increasing (Table 1).

The UV-Vis diffuse reflectance spectra (DRS) and the band gap values calculated by plots of  $(F(R)E)^{1/2}$  versus photo energy of  $\text{g-C}_3\text{N}_4$  and  $\text{MoS}_2/\text{g-C}_3\text{N}_4$  were shown in Fig. 7c and d. The band gap energy of  $\text{g-C}_3\text{N}_4$  in the onset is about 2.75 eV, corresponding to an absorption edge of 450 nm. After hybridization with  $\text{MoS}_2$ , the band gap of  $\text{MoS}_2/\text{g-C}_3\text{N}_4\text{-4}$  was about 2.72 eV, with an absorption edge of 455 nm. The composite photocatalyst can improve the utilization of the visible light.

The charge carrier separation and recombination rates of the photoexcited carriers were investigated using room temperature photoluminescence spectra with an excitation wavelength of 431 nm. As shown in Fig. 8,  $\text{g-C}_3\text{N}_4$  exhibited a broad emission peak that was centered at 450 nm due to the band–band PL phenomenon with the energy of light, which was approximately equal to the bandgap energy of  $\text{g-C}_3\text{N}_4$ . The emission peak of  $\text{MoS}_2/\text{g-C}_3\text{N}_4\text{-4}$  shifted to *ca.* 455 nm, which was consistent with DRS results. The PL intensity for  $\text{MoS}_2/\text{g-C}_3\text{N}_4\text{-4}$  decreased significantly compared to that of  $\text{g-C}_3\text{N}_4$ . In general, a decrease in the PL intensity indicates a suppressed electron–hole pair recombination, which makes  $\text{MoS}_2/\text{g-C}_3\text{N}_4\text{-4}$  to generate more photoelectrons and holes to participate in the photocatalytic reaction. This may be because the heterojunction formed at the interface and interior between  $\text{g-C}_3\text{N}_4$  and  $\text{MoS}_2$  with a high rate

Table 1 BET specific surface area and pore volume of the  $\text{g-C}_3\text{N}_4$  and  $\text{MoS}_2/\text{g-C}_3\text{N}_4$  samples

Samples	$\text{g-C}_3\text{N}_4$	$\text{MoS}_2/\text{g-C}_3\text{N}_4\text{-2}$	$\text{MoS}_2/\text{g-C}_3\text{N}_4\text{-4}$	$\text{MoS}_2/\text{g-C}_3\text{N}_4\text{-50}$
BET ( $\text{m}^2 \text{ g}^{-1}$ )	34.3807	11.3649	24.527	31.109
Pore volume ( $\text{cm}^3 \text{ g}^{-1}$ )	0.0778	0.0368	0.0465	0.0717



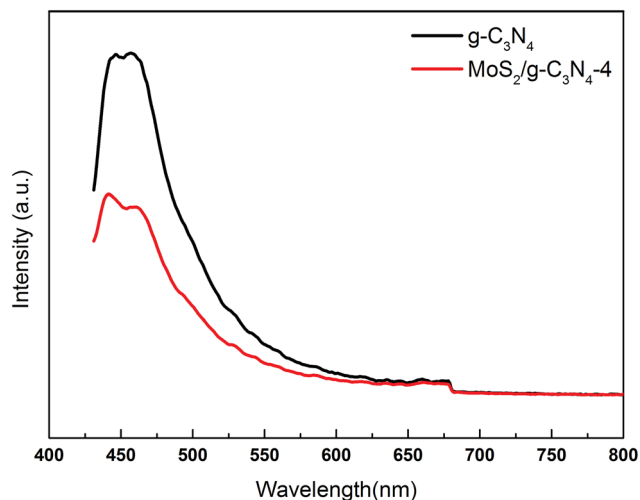


Fig. 8 PL spectra of pure  $g\text{-C}_3\text{N}_4$  and  $\text{MoS}_2/g\text{-C}_3\text{N}_4\text{-4}$  composites under 431 nm excitation at room temperature.

of carrier mobility can restrain the recombination of photo-generated charge effectively and accelerate charge transport.

To gain deeper insights into the charge transport behavior in the  $\text{MoS}_2/g\text{-C}_3\text{N}_4$  system, we conducted electrochemical impedance spectroscopy (EIS) measurements (Fig. 9). In the Nyquist diagram, the radius of each arc is associated with the charge-transfer process at the corresponding electrode/electrolyte interface with a smaller radius corresponding with a lower charge-transfer resistance.<sup>49,56</sup> The  $\text{MoS}_2/g\text{-C}_3\text{N}_4\text{-4}$  exhibited the smallest charge transfer resistance among all  $g\text{-C}_3\text{N}_4$ -based electrodes under visible light irradiation, suggesting that effective shuttling of charges between the electrode and the electrolyte, and faster interfacial charge transfer occurred at the composite interface and interior owing to the formation of the crossed-layer-structure.

Semiconductors exhibit unique photoelectric properties owing to their essential band-gap structure, which provides

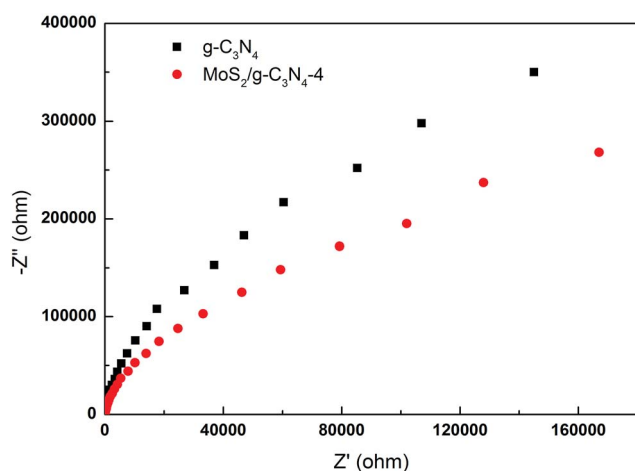


Fig. 9 Electrochemical impedance spectroscopy of the  $g\text{-C}_3\text{N}_4$  and  $\text{MoS}_2/g\text{-C}_3\text{N}_4\text{-4}$  samples.

a simple and economical light-to-electric conversion approach for various energy-related applications. Separation of electrons and holes plays a significant role in the photocatalytic reaction, and the photocurrent derives from electrons stimulated from the valence band to the conduction band under visible-light irradiation. Therefore, a higher photocurrent response indicates better separation efficiency of electrons and holes, which boosts the photocatalytic activity of the photocatalysts.<sup>57</sup> Fig. 10 displayed the transient photocurrent response of as-fabricated product printed on indium tin oxide (ITO) glass plates in a 0.5 M  $\text{Na}_2\text{SO}_4$  aqueous solution under 300 W Xe lamp light irradiation ( $\lambda > 400$  nm). The transient photocurrents of the samples were measured during repeated ON/OFF illumination cycles at 0.5 V. All of the samples exhibit prompt and reproducible photocurrent responses on each illumination. When the irradiation was interrupted, the photocurrent rapidly dropped to almost zero (steady-state value), and the photocurrent reverted to the original value once light was switched back on again. The photocurrent density of the  $\text{MoS}_2/g\text{-C}_3\text{N}_4\text{-4}$  is about 1.93 times higher than that of the pure  $g\text{-C}_3\text{N}_4$ . In addition, the transient photocurrent is widely regarded as the most efficient evidence to characterize charge separation.<sup>58</sup> The unique crossed-layer-structure can further shorten the photo-electron transport distance, thus facilitating fast electron transfer across the crossed-layer-structure. These results suggest that the enhanced light harvesting, efficient separation of photogenerated electron-hole pairs and the crossed-layer-structure of the  $\text{MoS}_2/g\text{-C}_3\text{N}_4$  have a significant role in improving the overall photoelectrochemical performance.

The photocatalytic activities of the as-prepared  $g\text{-C}_3\text{N}_4$ , and  $\text{MoS}_2/g\text{-C}_3\text{N}_4$  were evaluated by the degradation of MO in aqueous solution under visible light irradiation. We studied the influence of the  $\text{MoS}_2$  amount in  $\text{MoS}_2/g\text{-C}_3\text{N}_4$  on the photocatalytic activity under the same conditions. As shown in Fig. 11, the removal rate of MO using  $\text{MoS}_2/g\text{-C}_3\text{N}_4$  as the photocatalyst was higher than that of pure  $g\text{-C}_3\text{N}_4$ , and it improved gradually with the increase of the  $\text{MoS}_2$  amount. When the

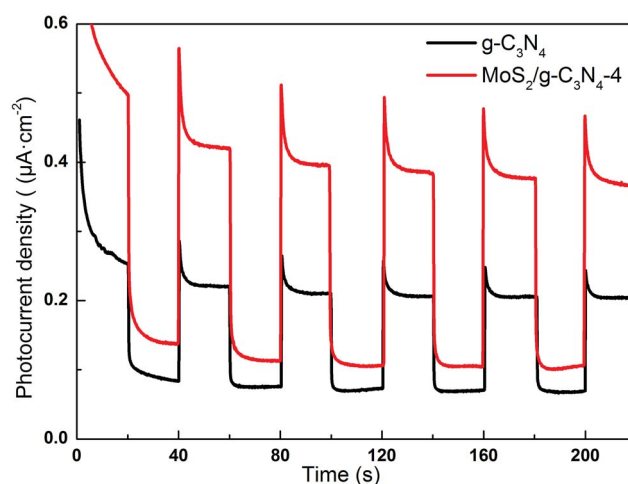


Fig. 10 Transient photocurrent response for the  $g\text{-C}_3\text{N}_4$  and  $\text{MoS}_2/g\text{-C}_3\text{N}_4\text{-4}$  sample.



percent of MoS<sub>2</sub> reached 0.16% (MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-4), it exhibited the highest photocatalytic activity. However, photocatalytic activity of MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-20 (0.8%) was significantly lower than that of MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-4 (0.16%). The reason might be that the too many active sites of g-C<sub>3</sub>N<sub>4</sub> were covered by MoS<sub>2</sub>, leading to weaken the absorption of visible light by g-C<sub>3</sub>N<sub>4</sub>, and causing the decrease of photocatalytic activity. Yet according to the results of BET analysis, surface areas and pore volumes were not main reason for enhancing photocatalytic degradation activity. We inferred that the enhanced activities were due to the synergistic effect between g-C<sub>3</sub>N<sub>4</sub> and MoS<sub>2</sub>. The optimum photocatalytic degradation efficiency of MO was 68% for MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-4 (0.16%), which is much higher than that of g-C<sub>3</sub>N<sub>4</sub>. In other words, it demonstrated 3.4 times higher performance than g-C<sub>3</sub>N<sub>4</sub>. Of course, we also tested the visible light activity for degradation of colorless pollutant (phenol) in Fig. S1.† Compared with degradation MO under stimulated solar light in Fig. 11, we observed that the tendency was consistent and MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-4 showed optimal activity among the MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> heterojunction catalysts. The result of photocatalytic degradation of phenol by thermal treatment g-C<sub>3</sub>N<sub>4</sub> was added in the Fig. S2,† there were no obvious change in degradation phenol under visible light, so in the photocatalytic degradation phenol process, the thermal treatment could not promote the photocatalytic activity of g-C<sub>3</sub>N<sub>4</sub>. In summary, the high performance was attributed to the enhanced light harvesting efficient separation of photogenerated electron-hole pairs, and the crossed-layer-structure of the MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>.

In order to confirm the main active species for the photocatalytic degradation of MO over MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>, the scavengers BQ, disodium EDTA and IPA were added to quench the possible active species <sup>•</sup>O<sub>2</sub><sup>-</sup>, hole (h<sup>+</sup>) and <sup>•</sup>OH, respectively.<sup>41-44</sup> As shown in Fig. 12, when disodium EDTA was added, the removal rate of MO had no obvious decrease, implying that h<sup>+</sup> did not dominate the photocatalytic reaction with MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>. However, the dramatic decline of MO removal was achieved in the presence of BQ and IPA, suggesting that <sup>•</sup>O<sub>2</sub><sup>-</sup> and <sup>•</sup>OH were

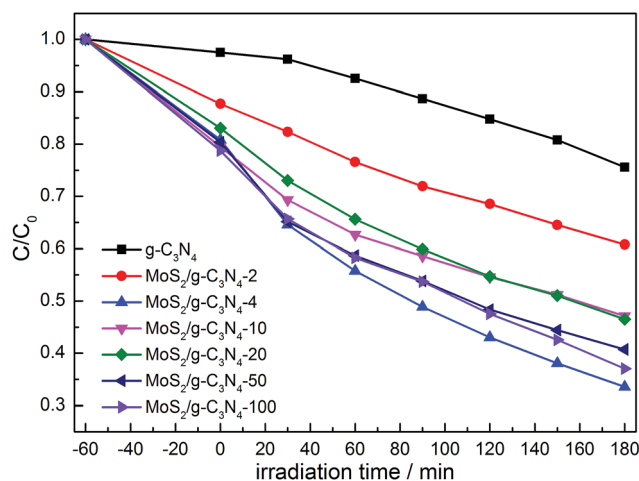


Fig. 11 Photocatalytic degradation of MO by g-C<sub>3</sub>N<sub>4</sub> and various MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> samples under simulated solar light.

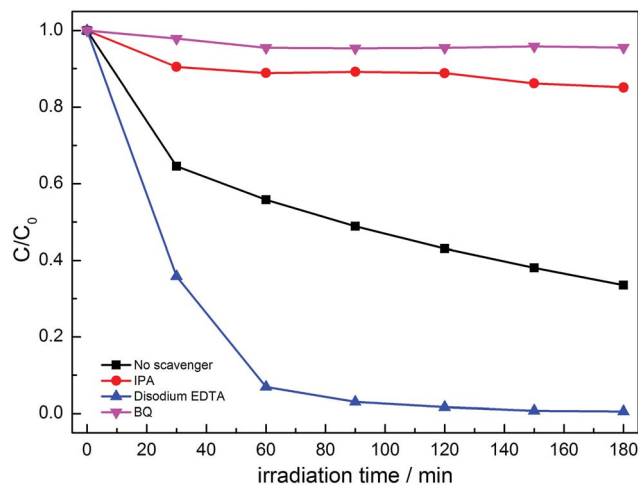


Fig. 12 Photocatalytic performances for the degradation of MO over MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub>-4 composite with different sacrificial agents under simulated solar light irradiation.

the main active species. According to the above results, we inferred that the enhanced activities stemmed from the synergistic effect of <sup>•</sup>O<sub>2</sub><sup>-</sup> and <sup>•</sup>OH active species between g-C<sub>3</sub>N<sub>4</sub> and MoS<sub>2</sub>.

The photocatalytic mechanisms are illustrated in Fig. 13. The photoexcited electrons would transfer from g-C<sub>3</sub>N<sub>4</sub> to MoS<sub>2</sub> due to the lower CB positions,<sup>53,58</sup> and the separated electrons on the surface of MoS<sub>2</sub> would combine with adsorbed O<sub>2</sub> to produce <sup>•</sup>O<sub>2</sub><sup>-</sup> radicals (the presence of <sup>•</sup>O<sub>2</sub><sup>-</sup> was confirmed by previous report<sup>59</sup>). Subsequently, it will combine with H<sup>+</sup> to produce H<sub>2</sub>O<sub>2</sub> which finally decomposes into hydroxyl radicals.<sup>60,61</sup> At the same time, the accumulated holes on the surface of g-C<sub>3</sub>N<sub>4</sub> would oxidize H<sub>2</sub>O to <sup>•</sup>OH, then <sup>•</sup>OH would oxidize dye molecules to H<sub>2</sub>O/CO<sub>2</sub>. In Fig. 11, disodium EDTA, as a sacrificial agent of hole, was added to the reaction system, with the holes decreasing, meanwhile, the balance moves to the direction which can produce holes and electrons. The mentioned factor would produce a large amount of electrons and improve the separation efficiency of photogenerated

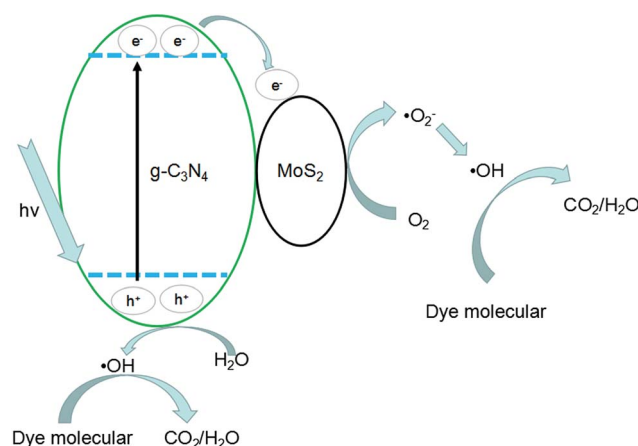


Fig. 13 Photocatalytic mechanism of MoS<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> heterostructures.



electron-hole pairs. In the reaction of dye degradation, the addition of disodium EDTA made its reaction rate increasing greatly. The observed MO degradation was more than 90% in an hour. When IPA and BQ were added into the reaction system, the  $\cdot\text{OH}$  and  $\cdot\text{O}_2^-$  were captured, which led to active component of dye degradation sharply reducing and we could observe the degradation ability had fallen dramatically. Compared with pure  $\text{g-C}_3\text{N}_4$ , the  $\text{MoS}_2/\text{g-C}_3\text{N}_4$ , the  $\text{MoS}_2/\text{g-C}_3\text{N}_4$  heterostructure had extended the lifetime of separated electron-hole pairs, resulting in the improvement of photocatalytic degradation efficiency.

## 4. Conclusions

We have demonstrated a novel method to prepare crossed-layer-structure  $\text{MoS}_2/\text{g-C}_3\text{N}_4$  heterojunction photocatalyst by impregnation-calcination, which have been evidenced by TEM, SEM, XRD, XPS, IR and UV-vis DRS analysis. Under simulated solar light irradiation for 3 h, the optimum photocatalytic degradation efficiency of MO was 68% for  $\text{MoS}_2/\text{g-C}_3\text{N}_4$ -4 (0.16%), which is much higher than that of  $\text{g-C}_3\text{N}_4$ . In other words, it demonstrated 3.4 times higher performance than  $\text{g-C}_3\text{N}_4$ . Of course,  $\text{MoS}_2/\text{g-C}_3\text{N}_4$ -4 also showed optimal activity in phenol degradation under visible light. The high performance was attributed to the enhanced light harvesting efficient separation of photogenerated electron-hole pairs and the crossed-layer-structure of the  $\text{MoS}_2/\text{g-C}_3\text{N}_4$ , which play a significant role in improving the overall photocatalytic performance.

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## Notes and references

- R. Dagherir, P. Drogui and D. Robert, *Ind. Eng. Chem. Res.*, 2013, **52**, 3581–3599.
- F. E. Osterloh, *Chem. Soc. Rev.*, 2013, **42**, 2294–2320.
- Y. Qu and X. Duan, *Chem. Soc. Rev.*, 2013, **42**, 2568–2580.
- S. Cao, J. Low, J. Yu and M. Jaroniec, *Adv. Mater.*, 2015, **27**, 2150–2176.
- P. Wang, S. Sun, X. Zhang, X. Ge and W. Lü, *RSC Adv.*, 2016, **6**, 33589–33598.
- B. Choudhury and P. K. Giri, *RSC Adv.*, 2016, **6**, 24976–24984.
- S. Le, T. Jiang, Q. Zhao, X. Liu, Y. Li, B. Fang and M. Gong, *RSC Adv.*, 2016, **6**, 38811–38819.
- J. Liebig, *Ann. Pharm.*, 1834, **10**, 1–47.
- J. Wen, X. Li, H. Li, S. Ma, K. He, Y. Xu, Y. Fang, W. Liu and Q. Gao, *Appl. Surf. Sci.*, 2015, **358**, 204–212.
- H. Zhang, L. Zhao, F. Geng, L.-H. Guo, B. Wan and Y. Yang, *Appl. Catal., B*, 2016, **180**, 656–662.
- G. Liu, P. Niu, C. Sun, S. C. Smith, Z. Chen, G. Q. Lu and H. M. Cheng, *J. Am. Chem. Soc.*, 2010, **132**, 11642–11648.
- S. Guo, Z. Deng, M. Li, B. Jiang, C. Tian, Q. Pan and H. Fu, *Angew. Chem.*, 2016, **55**, 1830–1834.
- S. C. Yan, Z. S. Li and Z. G. Zou, *Langmuir*, 2010, **26**, 3894–3901.
- G. Dong, K. Zhao and L. Zhang, *Chem. Commun.*, 2012, **48**, 6178.
- J. Li, B. Shen, Z. Hong, B. Lin, B. Gao and Y. Chen, *Chem. Commun.*, 2012, **48**, 12017.
- Y. Wang, Y. Di, M. Antonietti, H. Li, X. Chen and X. Wang, *Chem. Mater.*, 2010, **22**, 5119–5121.
- G. Zhang, M. Zhang, X. Ye, X. Qiu, S. Lin and X. Wang, *Adv. Mater.*, 2014, **26**, 805–809.
- X. Du, G. Zou, Z. Wang and X. Wang, *Nanoscale*, 2015, **7**, 8701–8706.
- Z.-F. Huang, J. Song, L. Pan, Z. Wang, X. Zhang, J.-J. Zou, W. Mi, X. Zhang and L. Wang, *Nano Energy*, 2015, **12**, 646–656.
- X. Ding, W. Ho, J. Shang and L. Zhang, *Appl. Catal., B*, 2016, **182**, 316–325.
- Z. Ni, F. Dong, H. Huang and Y. Zhang, *Catal. Sci. Technol.*, 2016, **6**, 6448–6458.
- F. Dong, Z. Zhao, Y. Sun, Y. Zhang, S. Yan and Z. Wu, *Environ. Sci. Technol.*, 2015, **49**, 12432–12440.
- T. Xiong, W. Cen, Y. Zhang and F. Dong, *ACS Catal.*, 2016, **6**, 2462–2472.
- D. Zheng, C. Pang, Y. Liu and X. Wang, *Chem. Commun.*, 2015, **51**, 9706–9709.
- K. Ai, Y. Liu, C. Ruan, L. Lu and G. M. Lu, *Adv. Mater.*, 2013, **25**, 998–1003.
- J. Zhang, M. Zhang, C. Yang and X. Wang, *Adv. Mater.*, 2014, **26**, 4121–4126.
- M. Shalom, S. Inal, C. Fettkenhauer, D. Neher and M. Antonietti, *J. Am. Chem. Soc.*, 2013, **135**, 7118–7121.
- Y. Zheng, L. Lin, B. Wang and X. Wang, *Angew. Chem., Int. Ed.*, 2015, **54**, 12868–12884.
- H. Wang, L. Zhang, Z. Chen, J. Hu, S. Li, Z. Wang, J. Liu and X. Wang, *Chem. Soc. Rev.*, 2014, **43**, 5234–5244.
- J. Zhu, P. Xiao, H. Li and S. A. Carabineiro, *ACS Appl. Mater. Interfaces*, 2014, **6**, 16449–16465.
- W.-J. Ong, L.-L. Tan, Y. H. Ng, S.-T. Yong and S.-P. Chai, *Chem. Rev.*, 2016, **116**, 7159–7329.
- H. Yan, *Chem. Commun.*, 2012, **48**, 3430–3432.
- Y. Wang, X. Wang, M. Antonietti and Y. Zhang, *ChemSusChem*, 2010, **3**, 435–439.
- J. Sun, J. Zhang, M. Zhang, M. Antonietti, X. Fu and X. Wang, *Nat. Commun.*, 2012, **1139**, DOI: 10.1038/ncomms2152.
- Y. Cui, Z. Ding, X. Fu and X. Wang, *Angew. Chem.*, 2012, **51**, 11814–11818.
- M. K. Bhunia, K. Yamauchi and K. Takanabe, *Angew. Chem., Int. Ed.*, 2014, **53**, 11001–11005.
- P. Niu, L. Zhang, G. Liu and H.-M. Cheng, *Adv. Funct. Mater.*, 2012, **22**, 4763–4770.
- Z. Lin and X. Wang, *Angew. Chem.*, 2013, **52**, 1735–1738.
- Y. S. Jun, J. Park, S. U. Lee, A. Thomas, W. H. Hong and G. D. Stucky, *Angew. Chem.*, 2013, **52**, 11083–11087.
- H. Zhang, *ACS Nano*, 2015, **9**, 9451–9469.
- M. Chhowalla, H. S. Shin, G. Eda, L. J. Li, K. P. Loh and H. Zhang, *Nat. Chem.*, 2013, **5**, 263–275.
- G. R. Bhimanapati, Z. Lin, V. Meunier, Y. Jung, J. Cha, S. Das, D. Xiao, Y. Son, M. S. Strano, V. R. Cooper, L. Liang,



- S. G. Louie, E. Ringe, W. Zhou, S. S. Kim, R. R. Naik, B. G. Sumpter, H. Terrones, F. Xia, Y. Wang, J. Zhu, D. Akinwande, N. Alem, J. A. Schuller, R. E. Schaak, M. Terrones and J. A. Robinson, *ACS Nano*, 2015, **9**, 11509–11539.
- 43 X. Zhang, Z. Lai, C. Tan and H. Zhang, *Angew. Chem.*, 2016, **55**, 8816–8838.
- 44 S. S. Chou, B. Kaehr, J. Kim, B. M. Foley, M. De, P. E. Hopkins, J. Huang, C. J. Brinker and V. P. Dravid, *Angew. Chem., Int. Ed.*, 2013, **52**, 4160–4164.
- 45 Q. Li, N. Zhang, Y. Yang, G. Wang and D. H. L. Ng, *Langmuir*, 2014, **30**, 8965–8972.
- 46 M. Li, L. Zhang, X. Fan, M. Wu, Y. Du, M. Wang, Q. Kong, L. Zhang and J. Shi, *Appl. Catal., B*, 2016, **190**, 36–43.
- 47 D. Zheng, G. Zhang, Y. Hou and X. Wang, *Appl. Catal., A*, 2016, **521**, 2–8.
- 48 X. Lu, Y. Jin, X. Zhang, G. Xu, D. Wang, J. Lv, Z. Zheng and Y. Wu, *Dalton Trans.*, 2016, **45**, 15406–15414.
- 49 Y. Hou, A. B. Laursen, J. Zhang, G. Zhang, Y. Zhu, X. Wang, S. Dahl and I. Chorkendorff, *Angew. Chem.*, 2013, **52**, 3621–3625.
- 50 H. Yu, P. Xiao, P. Wang and J. Yu, *Appl. Catal., B*, 2016, **193**, 217–225.
- 51 J. Li, E. Liu, Y. Ma, X. Hu, J. Wan, L. Sun and J. Fan, *Appl. Surf. Sci.*, 2016, **364**, 694–702.
- 52 W.-C. Peng and X.-Y. Li, *Catal. Commun.*, 2014, **49**, 63–67.
- 53 L. Ge, C. Han, X. Xiao and L. Guo, *Int. J. Hydrogen Energy*, 2013, **38**, 6960–6969.
- 54 W.-J. Ong, L.-L. Tan, S.-P. Chai, S.-T. Yong and A. R. Mohamed, *Nano Energy*, 2015, **13**, 757–770.
- 55 D. J. Martin, K. Qiu, S. A. Shevlin, A. D. Handoko, X. Chen, Z. Guo and J. Tang, *Angew. Chem., Int. Ed.*, 2014, **53**, 9240–9245.
- 56 L. Ma, H. Fan, J. Wang, Y. Zhao, H. Tian and G. Dong, *Appl. Catal., B*, 2016, **190**, 93–102.
- 57 J. Yan, Z. Chen, H. Ji, Z. Liu, X. Wang, Y. Xu, X. She, L. Huang, L. Xu, H. Xu and H. Li, *Chemistry*, 2016, **22**, 4764–4773.
- 58 Y. Hou, Z. Wen, S. Cui, X. Guo and J. Chen, *Adv. Mater.*, 2013, **25**, 6291–6297.
- 59 X. Bai, L. Wang, R. Zong and Y. Zhu, *J. Phys. Chem. C*, 2013, **117**, 9952–9961.
- 60 Q. Li, C. Zhang, J. Ma, G. Wang and D. H. L. Ng, *ChemCatChem*, 2014, **6**, 1392–1400.
- 61 Q. Li, J. Bian, L. Zhang, R. Zhang, G. Wang and D. H. L. Ng, *ChemPlusChem*, 2014, **79**, 454–461.

