Preparation of an ultrathin 2D/2D rGO/g-\(\text{C}_3\text{N}_4\) nanocomposite with enhanced visible-light-driven photocatalytic performance†

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A simple solvent method was developed to construct an ultrathin 2D/2D rGO/g-\(\text{C}_3\text{N}_4\) nanocomposite using a suspension of thermally exfoliated g-C\(_3\)N\(_4\) nanosheets and graphene oxide (GO) followed by a NaHSO\(_4\) reducing process. Different from the g-C\(_3\)N\(_4\) bulk, the results revealed that the g-C\(_3\)N\(_4\) nanosheets and rGO components in the as-prepared ultrathin nanocomposite have a strong interfacial interaction and abundant coupling interfaces. Moreover, improved visible light absorption properties and fast charge carrier separation efficiency were observed in the ultrathin 2D/2D rGO/g-\(\text{C}_3\text{N}_4\) nanocomposite, as compared to the pure g-\(\text{C}_3\text{N}_4\) nanosheets, which ensures its enhanced photocatalytic activity for methyl orange (MO) degradation and \(\text{CO}_2\) photoreduction. It was confirmed that the thermally exfoliated g-C\(_3\)N\(_4\) nanosheet is a good 2D material for the construction of 2D/2D heterostructures.

1. Introduction

Nowadays, energy and environmental issues have become a realistic and urgent problem all over the world.\(^1\)–\(^3\) Photocatalysis is regarded as one of the most promising technologies to settle both environmental and energy problems simultaneously through photochemical reactions.\(^4\)–\(^6\) However, low light utilization efficiency, fast recombination of charge carriers and the instability of numerous photocatalysts still limit the practical application of photocatalysis.

As a visible-light-driven photocatalyst, g-C\(_3\)N\(_4\) has attracted immense research interest in the past decade due to its tunable \(\text{configuration and good stability,}^{7\)–\(^9\) which makes it an excellent candidate for solar energy conversion and pollution abatement. However, pure g-C\(_3\)N\(_4\) suffers from shortcomings such as too small surface area, inefficient use of visible light and fast recombination of charge carriers.\(^10\)\(^,\)\(^11\) To overcome these problems, many methods have been developed to improve the photocatalytic performance of pure g-C\(_3\)N\(_4\) including metal doping,\(^12\)–\(^14\) non-metal doping,\(^15\)–\(^18\) morphology tailoring\(^19\)\(^,\)\(^20\) and semiconductors coupling\(^21\)\(^–\)\(^23\) and so on. Notably, as g-C\(_3\)N\(_4\) is a lamellar material, it provides a good 2D surface platform to obtain the highly efficient composite through constructing the unique 2D/2D heterostructures between g-C\(_3\)N\(_4\) and the other 2D species. Compared with 0D/2D and 1D/2D heterostructures, the 2D/2D heterostructures have a rather larger surface area and more abundant coupling heterointerfaces, which consequently enhances its photocatalytic performance.\(^24\)–\(^26\) So far, several 2D/2D heterostructures based on g-C\(_3\)N\(_4\) bulks have been developed, such as SnNb\(_2\)O\(_6\)/g-C\(_3\)N\(_4\),\(^26\) BiOBr/g-C\(_3\)N\(_4\) (ref. 27) and g-C\(_3\)N\(_4\)/Ca\(_2\)Nb\(_2\)TaO\(_{10}\).\(^28\) All 2D/2D heterostructures demonstrated the high photocatalytic activity for pollutants elimination and \(\text{H}_2\) production.\(^29\)–\(^28\) More recently, a hybrid catalyst of 2D/2D rGO/g-C\(_3\)N\(_4\) was prepared by directly heating a mixture of melamine and graphene oxide (GO) in air. It was observed that the photocatalytic activity of this 2D/2D composite for rhodamine B degradation was 2.6 times higher than that of pure g-C\(_3\)N\(_4\).\(^30\) Afterwards, the 2D/2D rGO/g-C\(_3\)N\(_4\) composite was also fabricated by incorporating rGO and protonated g-C\(_3\)N\(_4\) bulks through ultrasonic dispersion and electrostatic self-assembly strategy.\(^30\)–\(^31\) Since graphene and g-C\(_3\)N\(_4\) possess the similar carbon network and \(\text{sp}^2\) conjugated \(\pi\) structure, they have the excellent compatibility to form composite, which may possess an outstanding photocatalytic performance due to its superior interfacial interaction, fast electrons transfer and high charges separation efficiency.\(^32\) However, in most of cases, g-C\(_3\)N\(_4\) bulks were used as the matrix. The distribution and the effective coupling interfaces of as-prepared 2D/2D composites may be subject to the large size and tiny surface area of g-C\(_3\)N\(_4\) bulks. Neither heating melamine nor ultrasonic exfoliation of g-C\(_3\)N\(_4\) bulks could easily obtain the ultrathin 2D g-C\(_3\)N\(_4\) nanosheets with small size and large surface area. As a result, it is highly required to design and construct the ultrathin 2D/2D rGO/g-
C₃N₄ heterostructures based on 2D g-C₃N₄ nanosheets, which may also provide a new insight on the synthesis of the other composite materials for various practical applications.

Herein, a thermal-exfoliation method was used to fabricate ultrathin 2D g-C₃N₄ nanosheets at first.³⁵ Then, the acid pre-treatment with HCl can easily alter the g-C₃N₄ surface to possess positive polarity.³⁶ Upon reduction of GO using NaHSO₃ as a reducing agent, the resulting rGO/g-C₃N₄ heterostructures could be successfully obtained by the electrostatic attraction of oppositely charged materials. Finally, the as-obtained rGO/g-C₃N₄ was treated by vacuum cooling drying method to keep the stability of ultrathin 2D g-C₃N₄ structure. Compared with g-C₃N₄ bulks and the ultrathin g-C₃N₄ nanosheets, the photocatalytic activity of as-obtained 2D/2D rGO/g-C₃N₄ heterostructures for CO₂ photoreduction and methyl orange (MO) degradation has been significantly improved. The photocatalytic mechanism of this kind of 2D/2D heterostructures was also discussed.

2. Experimental

2.1 Synthesis of g-C₃N₄ bulks and g-C₃N₄ nanosheets

The thermal polymerization method was used for the preparation of g-C₃N₄ bulks. Typically, the precursor was composed by 3.5 g of urea and 1.5 g of dicyanodiamine which was put in a ceramic crucible covered with its lid. Then the precursor was sintered at 550 °C for 4 h in the furnace with the heating ramp of 2 °C min⁻¹. Until the furnace was cooled to room temperature, the yellow g-C₃N₄ bulks were collected. Next, the as-prepared g-C₃N₄ bulks were further sintered at 520 °C for 2 h with the heating ramp of 5 °C min⁻¹ in order to prepare ultrathin g-C₃N₄ nanosheets. Different from the preparation process of g-C₃N₄ bulks, the crucible in the thermal-exfoliation process was not covered with its lid. The white g-C₃N₄ nanosheets were collected when the furnace was cooled to room temperature.

2.2 Synthesis of 2D/2D rGO/g-C₃N₄ nanocomposite

Firstly, 1 g of as-prepared g-C₃N₄ nanosheets was added to 200 ml of HCl solution with concentration 0.5 mol l⁻¹ and was sonicated for 0.5 h to obtain slurry, which was continuously stirred for another 3.5 h to achieve completely proton-functionalized surface. Then the slurry was filtered and washed with deionized water and alcohol for several times.

Secondly, the GO dispersion (purchased from XFNANO, China) was diluted with water to obtain 100 ml of solution and sonicated for 0.5 h. Then 200 mg of as-obtained proton-functionalized g-C₃N₄ nanosheets was added to the solution and sonicated for another 0.5 h. After that, the solution was vigorously stirred for 1 h to achieve homogeneous suspension. The weight fractions of rGO in the 2D/2D rGO/g-C₃N₄ nanocomposite were set as 1, 3, 5, and 7 wt% respectively.

Thirdly, a certain amount of NaHSO₃ as reduce agent was added to the suspension, which was then transferred to an oil bath, heated to 95 °C and kept the temperature for 1.5 h to reduce GO. Finally, the suspension was filtered and vacuum dried at -60 °C for 6 h to obtain the 2D/2D rGO/g-C₃N₄ nanocomposite. According to the weight fractions of rGO in the nanocomposite, the as-prepared samples were denoted as 1rGO/g-C₃N₄, 3rGO/g-C₃N₄, 5rGO/g-C₃N₄ and 7rGO/g-C₃N₄ respectively. The diagram of detailed preparation process was shown in Fig. 1.

2.3 Characterization

X-Ray diffraction (XRD) patterns of the as-prepared nanocomposites were performed on an X-ray diffractometer (Bruker Advanced D8) using Ni-filtered CuKα radiation over the diffraction angle (2θ) ranging from 5° to 45°. The scan rate was set as 10° min⁻¹. The morphology and particle size of nanocomposites were characterized via scanning electron microscopy (SEM, Quanta 450) and transmission electron microscopy (TEM, FEI Tecnai G20). A Fourier transform infrared spectroscopy (FTIR) was acquired from Thermo-Nicolet iS10 with a standard KBr pellet method. The optical absorbance spectra of the nanocomposites were obtained using an ultraviolet-visible (UV-Vis) spectrophotometer (PerkinElmer, Lambda 750S). The photoluminescence (PL) spectra were carried out on an F-7000 spectrophotometer using a Xe lamp as excitation source. The excitation wavelength was 370 nm. The surface areas of the samples were measured using nitrogen adsorption–desorption by an automatic analyzer (ASAP 2020).

2.4 Evaluation of photocatalytic activity

The photocatalytic activity of as-prepared nanocomposites was evaluated by CO₂ photoreduction and MO photodegradation.

The photoreduction of CO₂ was conducted in a cylindrical steel reactor, which was vertically irradiated by a 300 W xenon lamp with a 400 nm cutoff filter. The schematic diagram of the photoreactor was demonstrated in Fig. S1.† The light spectrum of 400 nm UV light cutoff filter was provided in Fig. S2.† The distance from sample to 300 W Xenon lamp was 18 cm. In each experiment, 50 mg of nanocomposite powder was placed in the bottom of the cylindrical steel reactor with 120 ml of volume and 9 cm² of floor space. Besides, a minimal unsealed glass bottle with 1 ml deionized water was also put in the bottom of the cylindrical steel reactor simultaneously. Then, CO₂ gas with high purity (99.999%) was pushed into the reactor until the pressure of reactor reached 0.5 Mpa. In order to eliminate the air component, the reactor was washed 3 times by high purity CO₂ gas purging. Afterward, the light was turned on to initiate the reaction. About 0.5 ml of gas was taken out from the reactor.
for the analysis of CH₄ concentration using a gas chromatograph (GC-7920 with FID detector) after 3 h. Photodegradation of MO solution was also conducted with the same light source as CO₂ photoreduction. For each experiment, 50 ml of MO aqueous solution with a concentration of 20 ppm was mixed with 50 mg of nanocomposite. Before irradiation, the solution was put in a sealed glass beaker and ultrasonicated for 15 min, and then stirred for another 30 min in the dark to ensure adsorption–desorption equilibrium. 5 ml of the reaction suspension was taken out at 10 min intervals, centrifuged and examined at 464 nm using a UV-vis spectrophotometer (UV-2102PC).

3. Results and discussion

Fig. 2 shows the XRD patterns of as-prepared ultrathin g-C₃N₄ nanosheets and the series of 2D/2D rGO/g-C₃N₄ nanocomposites. Two apparent diffraction peaks are observed in all nanocomposites. As shown, the weak diffraction peak at 2θ = 13.11° with d value of 0.671 nm is related to the in-plane ordering of tri-s-triazine units, while the strong diffraction peak at 2θ = 27.31° is ascribed to the interlayer stacking of aromatic segments with d value of 0.324 nm. The two diffraction peaks can be indexed to (100) and (002) diffraction planes of g-C₃N₄ phase. From the results of XRD patterns, it can be found that the lattice structures of g-C₃N₄ are well-maintained after thermal exfoliation, acid pre-treatment and conjugation with GO process. In addition, no characteristic diffraction peak of GO at 2θ = 9.51° is observed in the series of rGO/g-C₃N₄ nanocomposites which may be attributed to the reduction of GO to rGO. However, the diffraction of rGO at 2θ = 24.51° is also not found, which may be due to the low content and fairly low diffraction intensity of rGO.

SEM and TEM measurements were conducted to characterize the surface morphologies and structures of g-C₃N₄ nanosheets and 3rGO/g-C₃N₄ nanocomposite in Fig. 3. Upon thermal exfoliation of g-C₃N₄ bulks, the thermal exfoliated g-C₃N₄ nanosheets present a thin layered structure without obvious domains of g-C₃N₄ bulks as shown in Fig. 3a and b. Meanwhile, the thermal exfoliated ultrathin g-C₃N₄ nanosheets comprised of rare layers are observed in the TEM images shown in Fig. 3c and d. The ultrathin g-C₃N₄ nanosheets display a porous structure (indicated by the white dotted circles) in Fig. 3c, which may be originated from the thermal polymerization of urea and dicyanodiamine. By employing these thermal-exfoliated ultrathin g-C₃N₄ nanosheets, we would get an ultrathin 2D/2D structure of g-C₃N₄ with rGO. Fig. 3e and f show the morphology of 3rGO/g-C₃N₄ nanocomposite. It can be observed that a 2D/2D structure has been successfully constructed. The porous material indicated by the white dotted circles in Fig. 3e may be g-C₃N₄, which is totally wrapped by rGO nanosheets with smooth surface. It indicates that there is an abundant interfacial contact area between g-C₃N₄ and rGO, which will in turn enhance the interfacial interaction and improve the charge carriers’ separation efficiency, thus increasing the photocatalytic reaction active sites for enhanced catalytic performance.

The N₂ adsorption–desorption isotherms of different samples were demonstrated in Fig. S3.† The surface areas of ultrathin g-C₃N₄ nanosheets, 3rGO/g-C₃N₄ and g-C₃N₄ bulks were calculated and determined as 158.56, 92.12 and 51.34 m² g⁻¹, respectively. It can be observed that the surface areas of ultrathin g-C₃N₄ and 3rGO/g-C₃N₄ are much higher than that of g-C₃N₄ bulks. Owing to the large surface area, it has been investigated that the photocatalytic activity of ultrathin g-C₃N₄ nanosheets was much higher than g-C₃N₄ bulks. Therefore, the photocatalytic activity of rGO/g-C₃N₄ nanocomposites based on g-C₃N₄ nanosheets may be further enhanced for this reason.

Fig. 4 shows the FTIR spectra of as-prepared ultrathin g-C₃N₄ nanosheets and the series of rGO/g-C₃N₄ nanocomposites. The broad absorption band around 3200 cm⁻¹ is originated from the N–H stretching vibration of uncondensed aminogroups and O–H stretches of adsorbed hydroxyl species. A series of peaks in the range from 1050 to 1800 cm⁻¹ are attributed to the typical stretching modes of C–N and C–NH heterocycles which
comprise both trigonal \( \text{N}-(\text{C})_3 \) (full condensation) and bridging \( \text{C}–\text{NH}–\text{C} \) units.\(^{38,39}\) The peak centered at 808 cm\(^{-1}\) is ascribed to the characteristic breathing mode of triazine units in g-C\(_3\)N\(_4\).\(^{31}\) Notably, the position of this peak in pure g-C\(_3\)N\(_4\) nanosheets exhibits a slight shift from 808.74 cm\(^{-1}\) to 811.59 cm\(^{-1}\) relative to 7rGO/g-C\(_3\)N\(_4\), indicating that a strong interaction is formed after the g-C\(_3\)N\(_4\) nanosheets are assembled with rGO by the electrostatic adsorption.\(^{40}\)

UV-Vis diffuse reflectance spectra of ultrathin g-C\(_3\)N\(_4\) nanosheets and the series of rGO/g-C\(_3\)N\(_4\) nanocomposites are demonstrated in Fig. 5. As shown in Fig. 5a, all rGO/g-C\(_3\)N\(_4\) nanocomposites exhibit the stronger visible light absorption compared with the ultrathin g-C\(_3\)N\(_4\) nanosheets, verifying that the introduction of rGO evidently improves the optical property of g-C\(_3\)N\(_4\) nanosheets. Noticeably, with the increase of rGO weight fraction in rGO/g-C\(_3\)N\(_4\) nanocomposites, a red shift to a longer wavelength is observed in the absorption edge of nanocomposites, which may result from the band gap narrowing of the rGO/g-C\(_3\)N\(_4\) nanocomposites. The plots of the transformed Kubelka–Munk value as a function of light energy are shown in Fig. 5b. It is clearly that the band gap of nanocomposite is narrowed in contrast to the ultrathin g-C\(_3\)N\(_4\) nanosheets. As the stronger visible light absorption of rGO/g-C\(_3\)N\(_4\) nanocomposites may improve the photo excitation efficiency of g-C\(_3\)N\(_4\), it is anticipated that the overall photocatalytic activity of nanocomposites will be enhanced. However, the high rGO weight fraction in nanocomposites may bring the negative effects because most of the incident light may be absorbed by rGO, while the available light for g-C\(_3\)N\(_4\) component obviously decreases. Therefore, for improving the light utilization, the appropriate rGO weight fraction in nanocomposites should be critically controlled.

PL was performed in order to investigate the transferring, trapping and separation efficiency of photo-generated charge carriers in ultrathin g-C\(_3\)N\(_4\) nanosheets and the series of rGO/g-C\(_3\)N\(_4\) nanocomposites in the photocatalytic reaction.\(^{38}\) As can be seen from Fig. 6, the PL intensity is found to following the sequence: ultrathin g-C\(_3\)N\(_4\) nanosheets > 1rGO/g-C\(_3\)N\(_4\) > 3rGO/g-C\(_3\)N\(_4\) > 5rGO/g-C\(_3\)N\(_4\) > 7rGO/g-C\(_3\)N\(_4\). Compared with the ultrathin g-C\(_3\)N\(_4\) nanosheets, the intensity of emission peak in the PL spectra of rGO/g-C\(_3\)N\(_4\) nanocomposites are obviously weakened, implying that the separation efficiency of photo-generated electron–hole pairs is much higher than that of pure g-C\(_3\)N\(_4\) nanosheets. Because rGO can act as an effective electron collector, the recombination of photo-generated electrons and holes in nanocomposites is evidently decreased with the increase of rGO weight fraction. As a result, the extent of the fluorescence quenching is noticeably increased with the increase of weight rGO fraction. Generally, an appropriate rGO weight fraction would enhance the photo-generated charge carriers’ separation efficiency of nanocomposites and improve its light utilization. However, an excessive rGO may weaken the visible light utilization of g-C\(_3\)N\(_4\) component owing to its light screening effect. Furthermore, the surface active sites on g-C\(_3\)N\(_4\) may be occupied by rGO nanosheets, which in turn results in a reduced photocatalytic performance of nanocomposite.\(^{41}\)
To evaluate the photocatalytic properties of ultrathin g-C$_3$N$_4$ nanosheets and the series of rGO/g-C$_3$N$_4$ nanocomposites under visible light illumination, MO removal in aqueous system with concentration of 20 ppm is firstly evaluated and depicted in Fig. 7a. The absorbance spectra of MO solution over ultrathin g-C$_3$N$_4$ nanosheets and 3rGO/g-C$_3$N$_4$ nanocomposite have been given in Fig. S4.† MO degradation efficiency over 1rGO/g-C$_3$N$_4$, 3rGO/g-C$_3$N$_4$, 5rGO/g-C$_3$N$_4$, 7rGO/g-C$_3$N$_4$, and ultrathin g-C$_3$N$_4$ is 86.32%, 97.5%, 87.93%, 80.42% and 77.45% respectively after 60 min of visible light irradiation. All rGO/g-C$_3$N$_4$ nanocomposites present the enhanced photocatalytic activity for MO degradation compared with ultrathin g-C$_3$N$_4$. In particular, the degradation efficiency of MO initially increases, then decreases with the increase of rGO weight fraction in rGO/g-C$_3$N$_4$ nanocomposites. The highest photocatalytic activity is achieved when 3rGO/g-C$_3$N$_4$ nanocomposite is used as photocatalyst. The corresponding kinetic constants ($k$) were calculated and shown in Fig. 7b. Thereinto, the MO degradation kinetic curves accord with pseudo first order by linear transforms $\ln \left( \frac{C_0}{C_t} \right) = kt$, where $C_0$ is the initial concentration of MO, $C_t$ is the concentration of MO at time $t$, and $k$ is the kinetic constant. The reaction rate constant ($k$) of 3rGO/g-C$_3$N$_4$ is 0.05191 min$^{-1}$/$\text{C}_0$, which is about 2.19 times higher than that of ultrathin g-C$_3$N$_4$ nanosheets.

Photocatalytic performance of ultrathin g-C$_3$N$_4$ nanosheets and the series of rGO/g-C$_3$N$_4$ nanocomposites in gaseous system under visible light illumination are secondly evaluated and shown in Fig. 8. As it can be found in Fig. 8a, the rGO hybridization has a substantial effect on the photocatalytic activity of rGO/g-C$_3$N$_4$ for the photoreduction of CO$_2$. The ultrathin g-C$_3$N$_4$ nanosheets exhibit a minimal total CH$_4$ evolution of 7.49 $\mu$mol/10 g catalyst over 3 h. Meanwhile, 3rGO/g-C$_3$N$_4$ demonstrates a total CH$_4$ evolution of 17.36 $\mu$mol/10 g catalysts over 3 h, which is about 2.32 times higher than that of ultrathin g-C$_3$N$_4$ nanosheets. To investigate the stability of nanocomposite, cyclic performance toward the photoreduction of CO$_2$ to CH$_4$ through consecutive five test cycles was carried out over the 3rGO/g-C$_3$N$_4$ nanocomposite. As shown in Fig. 8b, the 3rGO/g-C$_3$N$_4$ nanocomposite maintains its high photocatalytic activity for the CO$_2$ photoreduction after five cycles under the same conditions, demonstrating that the developed photocatalyst is stable with the prolonged reaction duration.

Many 2D/2D heterostructures based on g-C$_3$N$_4$ have been developed and demonstrated the enhanced photocatalytic activity for photocatalytic applications compared with pure g-C$_3$N$_4$, such as SnNb$_2$O$_6$/g-C$_3$N$_4$, BiOBr/g-C$_3$N$_4$, g-C$_3$N$_4$/Ca$_2$Nb$_2$TaO$_{10}$ (ref. 28) and rGO/g-C$_3$N$_4$. However, all these works are based on g-C$_3$N$_4$ bulks. In this work, the g-C$_3$N$_4$ bulks were thermal exfoliated to ultrathin g-C$_3$N$_4$ nanosheets. As the g-C$_3$N$_4$ nanosheets present the larger surface area than that of g-C$_3$N$_4$ bulks, it has been verified that the photocatalytic activity of g-C$_3$N$_4$ nanosheets was much higher than that of g-C$_3$N$_4$ bulks. Furthermore, we demonstrated the enhanced photocatalytic activity for MO degradation and CO$_2$ photoreduction for ultrathin 2D/2D rGO/g-C$_3$N$_4$ nanocomposites as compared to the g-C$_3$N$_4$ nanosheets with the same photocatalytic reaction parameters and the model pollutant molecules in this experiment. Therefore, the photocatalytic activity of 2D/2D heterostructures based on g-C$_3$N$_4$ nanosheets may present the further...
enhanced photocatalytic performance in contrast to g-C$_3$N$_4$ bulks.

Based on the above discussion, the proposed mechanism for MO degradation and CO$_2$ reduction under visible light irradiation over the rGO/g-C$_3$N$_4$ nanocomposite is shown in Fig. 9. When the rGO/g-C$_3$N$_4$ nanocomposite is irradiated with the visible light source, the g-C$_3$N$_4$ component is excited and the photo-generated electrons would transfer from its valence band (VB) to the conduction band (CB). Because of the good conductivity and remarkable electron storage capacity of rGO, the photo-generated electrons then migrate from g-C$_3$N$_4$ to rGO nanosheet in nanocomposite. The probability of electron–hole recombination is greatly reduced due to this transferring process, thus leading to the enhancement of photoinduced charges separation efficiency. On the other hand, the photo-generated holes remained in the VB of g-C$_3$N$_4$ may react with the adsorbed H$_2$O molecules or OH$^-$ ions to generate ·OH reactive radicals. Subsequently, the holes and the generated ·OH reactive radicals will directly degrade the MO molecules.\textsuperscript{43,44}

As for photoreduction of CO$_2$, it is widely known that two processes usually undergo in this process. Firstly, the photo-generated holes oxidize water to obtain hydrogen ions (H$^+$) via the half-reaction process (2H$_2$O $\rightarrow$ 4H$^+$ + O$_2$ + 4e$^-$). Secondly, the photogenerated electrons reduce CO$_2$ to CH$_4$ via acquiring 8-electrons process (CO$_2$ + 8H$^+$ + 8e$^-$ $\rightarrow$ CH$_4$ + 2H$_2$O).\textsuperscript{45,46} As the edges of VB and CB of g-C$_3$N$_4$ are determined to be +1.4 eV and –1.3 eV respectively, the above two processes can be completed in the rGO/g-C$_3$N$_4$ nanocomposites.\textsuperscript{19} It is worth mentioning that the photoreduction of CO$_2$ to CH$_4$ would be promoted by the enriched electron density on the rGO nanosheets.

In summary, the as-prepared ultrathin 2D/2D rGO/g-C$_3$N$_4$ nanocomposites have a large surface area and the abundant contact interfaces, which in turn promote the light absorption and increase the photogenerated charges separation efficiency. As a result, the high photocatalytic performance for MO degradation and CO$_2$ reduction over rGO/g-C$_3$N$_4$ nanocomposite is achieved by modifying ultrathin g-C$_3$N$_4$ with rGO nanosheets to form a unique ultrathin 2D/2D heterostructures. However, an optimum rGO weight fraction in rGO/g-C$_3$N$_4$ nanocomposites is should be critically controlled as the low rGO content in the rGO/g-C$_3$N$_4$ nanocomposite could not provide adequate electron storage sites to accelerate the separation of charges, while an excess of rGO may significantly weaken the visible light utilization of g-C$_3$N$_4$ component and occupy the surface active sites on g-C$_3$N$_4$, leading to a reduced photocatalytic performance.

4. Conclusions

In conclusion, we have successfully prepared an ultrathin 2D/2D rGO/g-C$_3$N$_4$ nanocomposite with thermal exfoliated g-C$_3$N$_4$ nanosheets and graphene oxide (GO) solution followed by a NaHSO$_3$ reducing process. To keep the structure stability of the ultrathin 2D/2D rGO/g-C$_3$N$_4$ nanocomposite, vacuum cooling drying method was used in this work. The ultrathin 2D/2D nanocomposite exhibited the enhanced visible light absorption, the strong interfacial interaction and the abundant coupling interfaces as supported by SEM, TEM, DRS and FTIR studies. Compared with the pure ultrathin g-C$_3$N$_4$ nanosheets, the as-prepared 2D/2D nanocomposite shows the significantly enhanced photocatalytic activity for MO degradation and CO$_2$ photoreduction. The ultrathin 2D/2D rGO/g-C$_3$N$_4$ nanocomposite with 3 wt% of rGO demonstrated the highest photocatalytic activity, which was 2.19 times and 2.32 times higher than that of pure ultrathin g-C$_3$N$_4$ nanosheets respectively. This enhanced visible light photocatalytic activity may be ascribed to the strong visible light absorption and interfacial interaction between g-C$_3$N$_4$ nanosheets and rGO component, which in turn facilitates the separation efficiency of charge carriers.

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Notes and references

1 M. E. Boot-Handford, J. C. Abanades, E. J. Anthony, M. J. Blunt, S. Brandani, N. Mac Dowell, J. R. Fernández, M. C. Ferrari, R. Gross, J. P. Hallett, R. S. Haszeldine, P. Heptonstall, A. Lyngfelt, Z. Makuch, E. Mangano, R. T. J. Porter, M. Pourkashanian, G. T. Rochelle, N. Shah, J. G. Yaoa and P. S. Fennell, Energy Environ. Sci., 2014, 7, 130–189.

2 N. M. P. Bocken, S. W. Short, P. Rana and S. Evans, J. Cleaner Prod., 2014, 65, 42–56.

3 Z. Liu, D. Guan, W. Wei, S. Davis, P. Ciais, J. Bai, S. Peng, Q. Zhang, K. Hubacek, G. Marland, R. J. Andres, D. Crawford-Brown, J. Lin, H. Zhao, C. Hong, T. A. Boden, K. Feng, G. P. Peters, F. Xi, J. Liu, Y. Li, Y. Zhao, N. Zeng and K. He, Nature, 2015, 524, 335–338.
C. Yu, W. Zhou, L. Hong, L. Yuan and D. D. Dionysiou, *Chem. Eng. J.*, 2016, 287, 117–129.

C. Acar, I. Dincer and C. Zamfirescu, *Int. J. Energy Res.*, 2014, 38, 1903–1920.

D. Sudha and P. Sivakumar, *Chem. Eng. Process.*, 2015, 97, 112–133.

S. Ye, R. Wang, M. Z. Wu and Y. P. Yuan, *Appl. Surf. Sci.*, 2015, 358, 15–27.

J. Zhu, P. Xiao, H. Li and S. A. Carabineiro, *ACS Appl. Mater. Interfaces*, 2014, 6, 16449–16465.

S. Cao, J. Low, J. Yu and M. Jaroniec, *Adv. Mater.*, 2015, 27, 2150–2176.

J. Liu, H. Wang and M. Antonietti, *Chem. Soc. Rev.*, 2016, 45, 2308.

G. Mamba and A. K. Mishra, *Appl. Catal., B*, 2016, 198, 347–377.

W. J. Ong, L. L. Tan, S. P. Chai and S. T. Yong, *Dalton Trans.*, 2014, 44, 1249–1257.

S. Hu, R. Jin, G. Lu, D. Liu and J. Gui, *RSC Adv.*, 2014, 4, 24863–24869.

G. Jiang, X. Li, M. Lan, T. Shen, X. Lv, F. Dong and S. Zhang, *Appl. Catal., B*, 2017, 205, 532–540.

T. Xiong, W. Cen, Y. Zhang and F. Dong, *ACS Catal.*, 2016, 6, 2462–2472.

Z. F. Huang, J. Song, L. Pan, Z. Wang, X. Zhang, J. J. Zou, W. Mi, X. G. Zhang and L. Wang, *Nano Energy*, 2015, 12, 646–656.

K. Wang, Q. Li, B. Liu, B. Cheng, W. Ho and J. Yu, *Appl. Catal., B*, 2015, 176–177, 44–52.

Y. P. Zhu, T. Z. Ren and Z. Y. Yuan, *ACS Appl. Mater. Interfaces*, 2015, 7, 16850–16856.

W. Zhang, Z. Zhao, F. Dong and Y. Zhang, *Chin. J. Catal.*, 2017, 38, 372–378.

Z. Huang, F. Li, B. Chen and G. Yuan, *RSC Adv.*, 2015, 5, 14027–14033.

L. Y. Chen and W. D. Zhang, *Appl. Surf. Sci.*, 2014, 301, 428–435.

H. Shi, G. Chen, C. Zhang and Z. Zou, *ACS Catal.*, 2016, 4, 3637–3643.

W. Jiang, W. Luo, J. Wang, M. Zhang and Y. Zhu, *J. Photochem. Photobiol., C*, 2016, 28, 87–115.

H. Huang, K. Xiao, N. Tian, X. Du and Y. Zhang, *Colloids Surf., A*, 2016, 511, 64–72.

F. Dong, T. Xiong, Y. Sun, Y. Zhang and Y. Zhou, *Chem. Commun.*, 2015, 51, 8249–8252.

Z. Zhang, D. Jiang, D. Li, M. He and M. Chen, *Appl. Catal., B*, 2016, 183, 113–123.

Y. Sun, W. Zhang, T. Xiong, Z. Zhao, F. Dong, R. Wang and W. K. Ho, *J. Colloid Interface Sci.*, 2014, 418, 317–323.

S. Thaweesak, M. Lyu, P. Peerakiatkhajohn, T. Butburee, B. Luo, H. Chen and L. Wang, *Appl. Catal., B*, 2017, 202, 184–190.

B. Yuan, J. Wei, T. Hu, H. Yao, Z. Jiang, Z. Fang and Z. Chu, *Chin. J. Catal.*, 2015, 36, 1009–1016.

W. J. Ong, L. L. Tan, S. P. Chai, S. T. Yong and A. R. Mohamed, *Nano Energy*, 2015, 13, 757–770.

S. Hu, W. Zhang, J. Bai, G. Lu, L. Zhang and G. Wu, *RSC Adv.*, 2016, 6, 25695–25702.

C. Jian, X. Xu, L. Tao, K. Pandiselvi and J. Wang, *Sci. Rep.*, 2016, 6, 37318.

Y. Li, M. Wang, S. J. Bao, S. Lu, M. Xu, D. Long and S. Pu, *Ceram. Int.*, 2016, 42, 18521–18528.

F. Dong, T. Xiong, Y. Sun, Y. Zhang and Y. Zhou, *Chem. Commun.*, 2015, 51, 8249–8252.

Z. Zhang, D. Jiang, D. Li, M. He and M. Chen, *Appl. Catal., B*, 2016, 183, 113–123.

Y. Sun, W. Zhang, T. Xiong, Z. Zhao, F. Dong, R. Wang and W. K. Ho, *J. Colloid Interface Sci.*, 2014, 418, 317–323.

S. Thaweesak, M. Lyu, P. Peerakiatkhajohn, T. Butburee, B. Luo, H. Chen and L. Wang, *Appl. Catal., B*, 2017, 202, 184–190.

B. Yuan, J. Wei, T. Hu, H. Yao, Z. Jiang, Z. Fang and Z. Chu, *Chin. J. Catal.*, 2015, 36, 1009–1016.

W. J. Ong, L. L. Tan, S. P. Chai, S. T. Yong and A. R. Mohamed, *Nano Energy*, 2015, 13, 757–770.

S. Hu, W. Zhang, J. Bai, G. Lu, L. Zhang and G. Wu, *RSC Adv.*, 2016, 6, 25695–25702.

C. Jian, X. Xu, L. Tao, K. Pandiselvi and J. Wang, *Sci. Rep.*, 2016, 6, 37318.

Y. Li, M. Wang, S. J. Bao, S. Lu, M. Xu, D. Long and S. Pu, *Ceram. Int.*, 2016, 42, 18521–18528.

Y. Li, R. Jin, F. Xu, Y. Yang, Y. Man, X. Liu, Y. Xing and S. Song, *J. Hazard. Mater.*, 2016, 313, 219–228.

M. Fu, Q. Jiao and Y. Zhao, *Mater. Charact.*, 2013, 86, 303–315.

K. Zhang, L. L. Zhang, X. S. Zhao and J. Wu, *Chem. Mater.*, 2010, 22, 1392–1401.

Q. Cai, J. Shen, Y. Feng, Q. Shen and H. Yang, *J. Alloys Compd.*, 2015, 628, 372–378.

R. C. Pawar, V. Khare and C. S. Lee, *Dalton Trans.*, 2014, 43, 12514–12527.

M. Xu, L. Han and S. Dong, *ACS Appl. Mater. Interfaces*, 2013, 5, 12533–12540.

H. Gu, T. Zhou and G. Shi, *Talanta*, 2015, 132, 871–876.

S. Hu, W. Zhang, J. Bai, G. Lu, L. Zhang and G. Wu, *RSC Adv.*, 2016, 6, 25695–25702.

A. Siahvashi, D. Chesterfield and A. A. Adesina, *Chem. Eng. Sci.*, 2013, 93, 313–325.

S. C. Yan, Z. S. Li and Z. G. Zou, *Langmuir*, 2010, 26, 3894–3901.

C. Han, G. Lei, C. Chen, Y. Li, X. Xiao, Y. Zhang and L. Guo, *Appl. Catal., B*, 2014, 147, 546–553.

E. Karamian and S. Sharifnia, *J. CO2 Util.*, 2016, 16, 194–203.

Y. Bai, T. Chen, P. Wang, L. Wang, L. Ye, X. Shi and W. Bai, *Sol. Energy Mater. Sol. Cells*, 2016, 157, 406–414.