

Cite this: *RSC Adv.*, 2019, 9, 28894Received 19th June 2019
Accepted 4th September 2019

DOI: 10.1039/c9ra04606b

rsc.li/rsc-advances

Facile one-pot synthesis of Mg-doped g-C₃N₄ for photocatalytic reduction of CO₂†

Xinyue Dong,^a Suicai Zhang,^b Hualin Wu,^b Zhuo Kang ^{*bc} and Li Wang^{*ad}

Graphitic carbon nitride (g-C₃N₄) has attracted wide attention due to its potential in solving energy and environmental issues. However, rapid charge recombination and a narrow visible light absorption region limit its performance. In our study, Mg-doped g-C₃N₄ was synthesized through a facile one-pot strategy for CO₂ reduction. After Mg doping, the light utilization efficiency and photo-induced electron–hole pair separation efficiency of the catalysts were improved, which could be due to the narrower band gap and introduced midgap states. The highest amounts of CO and CH₄ were obtained on Mg-CN-4% under ultraviolet light illumination, which were about 5.1 and 3.8 times that of pristine g-C₃N₄, respectively; the yield of CO and CH₄ reached 12.97 and 7.62 μmol g⁻¹ under visible light irradiation. Our work may provide new insight for designing advanced photocatalysts in energy conversion applications.

Introduction

Semiconductor photocatalysis technology is regarded as a green and promising strategy that uses solar energy to solve the energy crisis and environmental pollution problems faced by human beings.^{1–3} Fujishima and Honda successfully used a TiO₂ photocatalyst for water splitting in 1972,⁴ which marked the beginning of a new era in the field of photocatalysis. At present, photocatalysts such as titanium dioxide (TiO₂),⁵ zinc oxide (ZnO),⁶ graphene carbonitride (g-C₃N₄),⁷ bismuth oxide (Bi₂O₃),⁸ cadmium sulfide (CdS),⁹ and layered metal hydroxides,¹⁰ have been used in the field of photocatalysis.¹¹

g-C₃N₄ is a potential 2D layered structure polymer organic semiconductor photocatalyst with a band gap width of 2.7 eV, which has attracted widespread attention of researchers.^{12–14} The advantages of low-cost, abundant and stability make it widely applied in water decomposition,¹⁵ CO₂ reduction,¹⁶ NO removal,¹⁷ and organic pollutant degradation.¹⁸ However, the low specific surface area, high photo-generated electron–hole pairs recombination, and low visible-light harvesting capacity limit its development.^{19,20}

Strategies for g-C₃N₄ modification, including metal doping (K,²¹ Cu,²² Pd,²³ Mn²⁴), non-metal doping (B,²⁵ N,²⁶ S,²⁷ P²⁸), semiconductor recombination (V₂O₅,²⁹ MnO₂,³⁰ CeO₂,³¹ TiO₂ (ref. 32)), and so on, have been developed to suppress the recombination of photo-generated electron–hole pairs, expand the visible light response range, and improve the photocatalytic performance.^{33,34} Among these strategies, element doping was a hot research. For example, Kumar *et al.* synthesized Se-doped g-C₃N₄ by facile one-pot two-step strategy and the catalysts demonstrated high photocatalytic performance in CO₂ reduction.³⁵ Wang *et al.* synthesized K-doped g-C₃N₄ by thermal polymerization, and the photocatalytic hydrogen evolution rate of K-doped g-C₃N₄ was improved.²¹ Tang *et al.* synthesized Mg/g-C₃N₄ by hydrothermal deposition method and the results show that Mg element plays an important role in increasing electron–holes pairs separation and extending the range of light response.³⁶ However, doping of Mg into carbon nitride by one-pot method has not been explored so far.

In this manuscript, Mg-doped g-C₃N₄ photocatalysts with various doping amount have been successfully prepared by one-pot method with magnesium chloride as Mg source. The catalysts were characterized by a series of methods such as, XRD, TEM, XPS, UV-vis, and PL. The catalytic performance towards photocatalytic CO₂ reduction was evaluated in the gas phase reactor.

Results and discussion

The morphologies of samples were observed by SEM. As shown in Fig. 1(a), the SEM image of g-C₃N₄ displays typical two-dimensional layered structure with wrinkles and irregular stacks.³⁷ After doping Mg (Fig. 1(b)), the morphology of g-C₃N₄ changes to some extent, which may be attributed to

^aSchool of Energy and Environmental Engineering, University of Science and Technology Beijing, Beijing 100083, China. E-mail: wangli@ces.ustb.edu.cn

^bState Key Laboratory for Advanced Metals and Materials, School of Materials Science and Engineering, University of Science and Technology Beijing, Beijing 100083, P. R. China

^cBeijing Municipal Key Laboratory of New Energy Materials and Technologies, University of Science and Technology Beijing, Beijing 100083, P. R. China

^dKey Laboratory of Resource-Oriented Treatment of Industrial Pollutants, Beijing 100083, China

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c9ra04606b



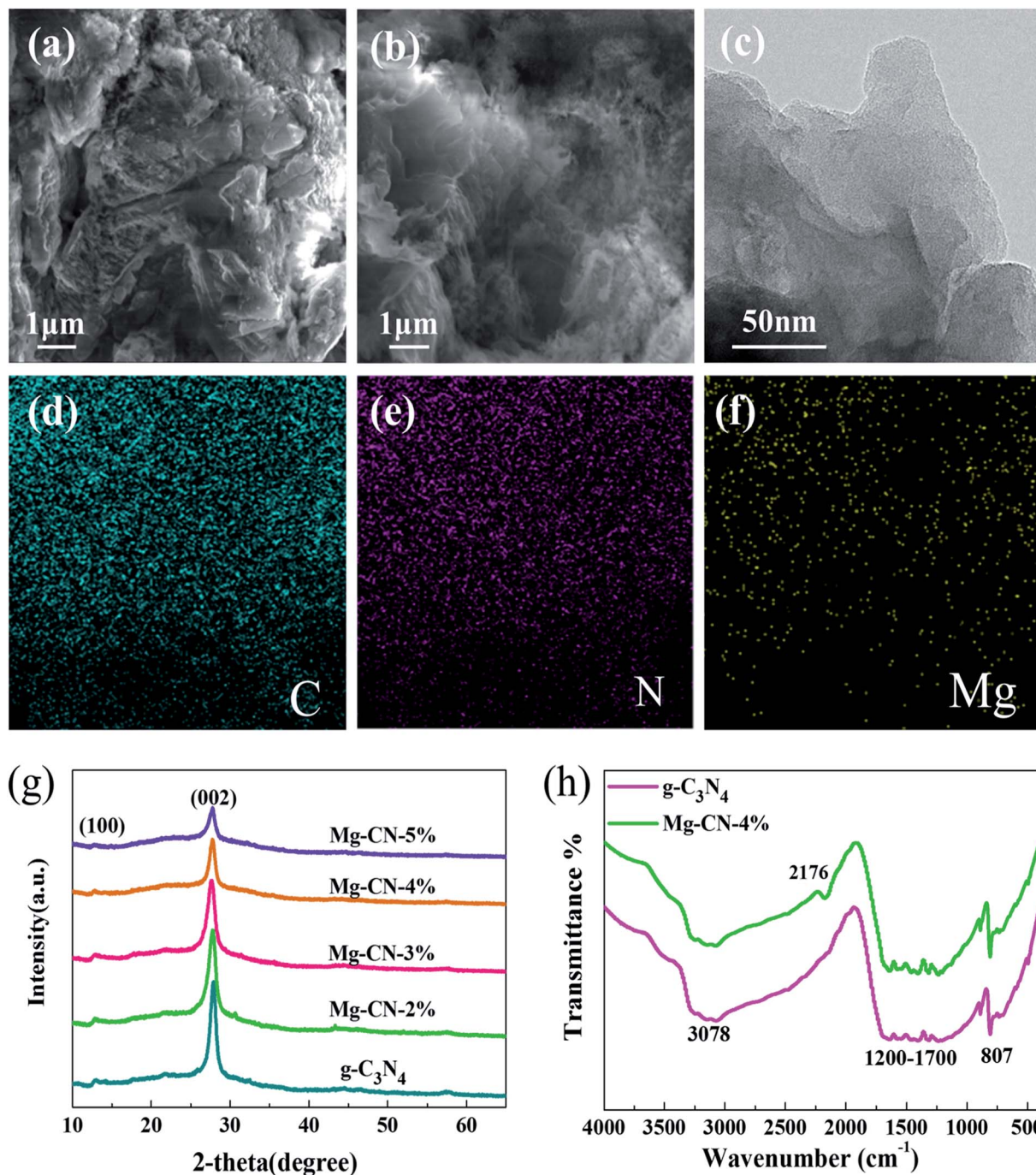


Fig. 1 SEM images of (a) g-C₃N₄, (b) Mg-CN-4%; (c) TEM image of Mg-CN-4%; (d) elemental mapping images of C, (e) N, (f) Mg of Mg-CN-4%; (g) XRD spectra of samples; (h) FTIR spectra of g-C₃N₄ and Mg-CN-4%.

delamination effect.³⁸ Compared with g-C₃N₄, the layers of Mg-CN-4% become thinner and curlier. The Mg-CN-4% having optimum photocatalytic activity was selected for TEM characterization. The TEM image of Mg-CN-4% shows a typical layered structure in Fig. 1(c).³⁹ The EDX spectrum (Fig. S1†) and elemental mapping of the Mg-CN-4% indicates that the sample consisted of C, N and Mg elements.

The crystalline structures of samples were analyzed by XRD. Fig. 1(g) shows XRD patterns of g-C₃N₄ and Mg-doped

g-C₃N₄. For g-C₃N₄ sample, there are two typical diffraction peaks at 12.86° and 27.79° corresponding to (100) and (002) facet of g-C₃N₄. The peak at 12.86° is assigned to in-plane repeating tri-s-triazine units, and the sharp peak at 27.79° is ascribed to interlayer stacking of conjugated aromatics.⁴⁰ Mg-doped g-C₃N₄ has similar diffraction peaks with g-C₃N₄, indicating that Mg doping does not change the graphitic-like structure of g-C₃N₄. No diffraction peaks of the Mg species were observed in the Mg-CN-4%, manifesting that



Mg was successfully doped into $g\text{-C}_3\text{N}_4$. After doping Mg, the peak intensity of the catalyst decreases, indicating that Mg doping can reduce the crystallinity of $g\text{-C}_3\text{N}_4$.⁴¹

The doping state and bonding characteristics of $g\text{-C}_3\text{N}_4$ and Mg-CN-4% was confirmed by FTIR spectra. In Fig. 1(h), the representative peak appeared at 807 cm^{-1} belonged to the characteristic breathing vibration mode for tri-*s*-triazine rings of $g\text{-C}_3\text{N}_4$,⁴² and the peaks at $1200\text{--}1700\text{ cm}^{-1}$ are ascribed to the typical CN heterocycles stretching modes.⁴³ The peaks at $1240\text{--}1400\text{ cm}^{-1}$ and $1480\text{--}1650\text{ cm}^{-1}$ are assigned to the stretching modes of C–N and C=N groups, respectively.²¹ Another broad peak at around 3078 cm^{-1} is caused by N–H bonds, which is derived from uncondensed amine groups.⁴⁴ Compared with $g\text{-C}_3\text{N}_4$, a distinct peak appeared at 2176 cm^{-1} in Mg-CN-4%, which can be attributed to C≡N group. Due to Mg-induced surface defects, some sp^2 C–N bonds were converted into C≡N bonds in Mg-CN-4%.^{45,46}

To further examine the surface chemical composition and elemental valence of samples, XPS measurements was performed. Fig. 2(a) illustrates XPS survey spectra of $g\text{-C}_3\text{N}_4$ and Mg-CN-4%, elements of C, N, O were detected in both catalysts. Mg elements can be observed in Mg-CN-4%, which

is consistent with the results of XRD. The N1s peaks of $g\text{-C}_3\text{N}_4$ can be deconvoluted into four characteristic peaks with binding energy at 398.68 eV, 399.35 eV, 400.87 eV, and 404.61 eV, respectively (Fig. 2(b)). The most intense peak was at 398.68 eV, corresponding to sp^2 hybridized N in C–N=C groups. The peaks at 399.35 eV and 400.87 eV are ascribed to tertiary N atoms bonded of N–(C)₃ groups and amino functional groups⁴⁷ and the peak at 404.61 eV is attributed to typical π -excitations in the heterocycles. The C1s spectra of $g\text{-C}_3\text{N}_4$ are shown in Fig. 2(c), from which two peaks can be observed at 284.94 eV and 288.25 eV, corresponding to C–C bonds and N–C–N coordination in the graphitic structure, respectively.⁴⁸ The Mg2p binding energy is 50.81 eV, which usually belongs to Mg^{2+} .^{49,50} In the O1s XPS spectra (Fig. S2†), no peaks appeared at $\sim 528\text{--}531\text{ eV}$, which correspond to O in metal oxides.²³ Therefore, magnesium is doped into the CN skeleton in an ionic state. Compared with $g\text{-C}_3\text{N}_4$, the C1s and N1s peaks of Mg-CN-4% show a slight shift, indicating that the surface chemical structure of Mg-doped $g\text{-C}_3\text{N}_4$ changes.⁵¹

The optical absorption properties of $g\text{-C}_3\text{N}_4$ and Mg-CN-4% were studied by UV-vis diffuse reflectance spectra. As

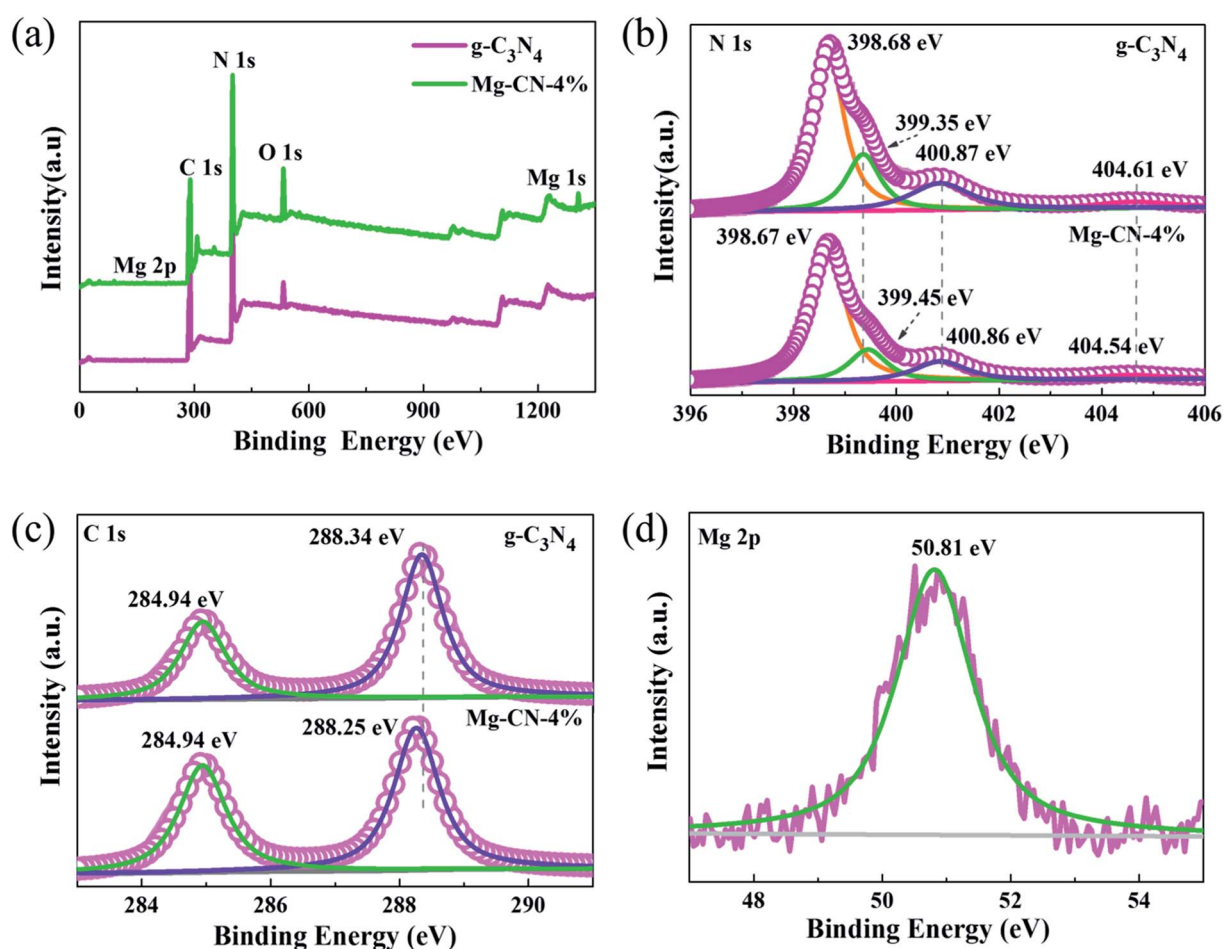


Fig. 2 (a) Wide survey scans, (b) N1s high-resolutions, (c) C1s high-resolutions of $g\text{-C}_3\text{N}_4$ and Mg-CN-4%, (d) Mg2p high-resolutions of Mg-CN-4%.



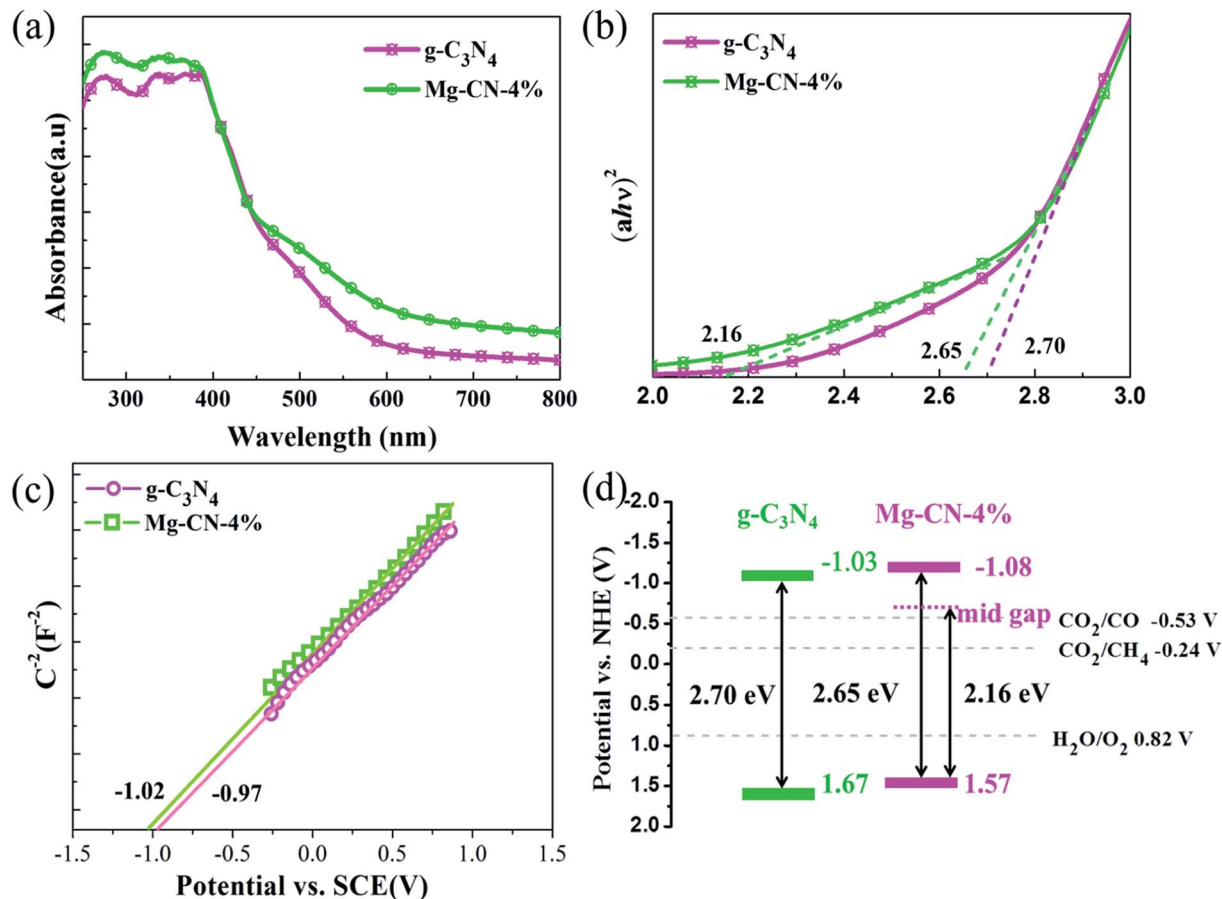


Fig. 3 (a) UV-vis diffuse reflectance spectra, (b) the plots of $(ah\nu)^2$ versus $h\nu$ for the band gap energies, (c) Mott-Schottky plots, (d) band gap structure of $g\text{-C}_3\text{N}_4$ and Mg-CN-4%.

seen in Fig. 3(a), the intensity of light absorption of Mg-CN-4% was higher than $g\text{-C}_3\text{N}_4$ from ultraviolet light to visible light region. The optical band gap energies of as-prepared photocatalysts have been calculated by using the following formula:

$$ah\nu = A(h\nu - E_g)^n$$

where a , h , ν , A and E_g are the absorption coefficient, Planck's constant, light frequency, proportionality constant and bandgap, respectively.⁵² The optical band gap energies for samples were obtained from a plot of $(ah\nu)^2$ vs. $h\nu$ ($n = 1/2$ for a direct transition, $n = 2$ for an indirect transition). As shown in Fig. 4(b), the band gap of $g\text{-C}_3\text{N}_4$ was estimated to be 2.70 eV, which is equal to the band gap in the previously reported paper.⁵³ It is clearly revealed that Mg doping reduced the band gap of Mg-CN-4% to 2.65 eV. The UV-vis spectra of Mg-CN-4% presented a tail absorption which indicates the formation of the midgap states.³⁵ It is reported that such midgap states is due to the existence of disorders/defects within the band gap.⁵⁴ The Mott-Schottky curves of $g\text{-C}_3\text{N}_4$ and Mg-CN-4% are shown in Fig. 3(c). The flat band potentials of $g\text{-C}_3\text{N}_4$ and Mg-CN-4% were confirmed as -0.97 V and -1.02 V vs. SCE, which correspond to -0.73 V and -0.78 V vs. NHE, respectively. According to the rule

of thumb, the conduction band (CB) is more negative by about -0.3 eV than the flat band potential.⁴² The CB of $g\text{-C}_3\text{N}_4$ and Mg-CN-4% could be estimated to be -1.03 V and -1.08 V. According to their band gap, the valence band (VB) position of $g\text{-C}_3\text{N}_4$ and Mg-CN-4% can be calculated as 1.67 V and 1.57 V, respectively. Combined UV-vis spectrum with the Mott-Schottky curves, Fig. 3(d) shows the VB and CB positions of Mg-CN-4%. The midgap states energy from VB to midgap states for Mg-CN-4% was determined to be 2.16 eV. Therefore, the exact position of midgap states for Mg-CN-4% was calculated as -0.59 V, which is higher than CO_2 reduction level (CO_2/CO , -0.53 V vs. NHE; CO_2/CH_4 , -0.24 V vs. NHE). Due to the presence of midgap states, Mg-CN-4% could harvest long-wavelength visible light to stimulate electrons in the VB to midgap states.^{55,56} Moreover, the excited electrons in CB could be temporarily trapped by midgap states to suppress electron-hole pairs recombination.^{57,58}

PL analysis is commonly performed to study the migration, transfer and recombination of electron-hole pairs produced by the photocatalyst. The PL spectra of $g\text{-C}_3\text{N}_4$ and Mg-CN-4% with excitation wavelength of 325 nm are shown in Fig. 4(a). It can be seen that the spectra of Mg-CN-4% is similar to $g\text{-C}_3\text{N}_4$ with a strong peak around 465 nm. When



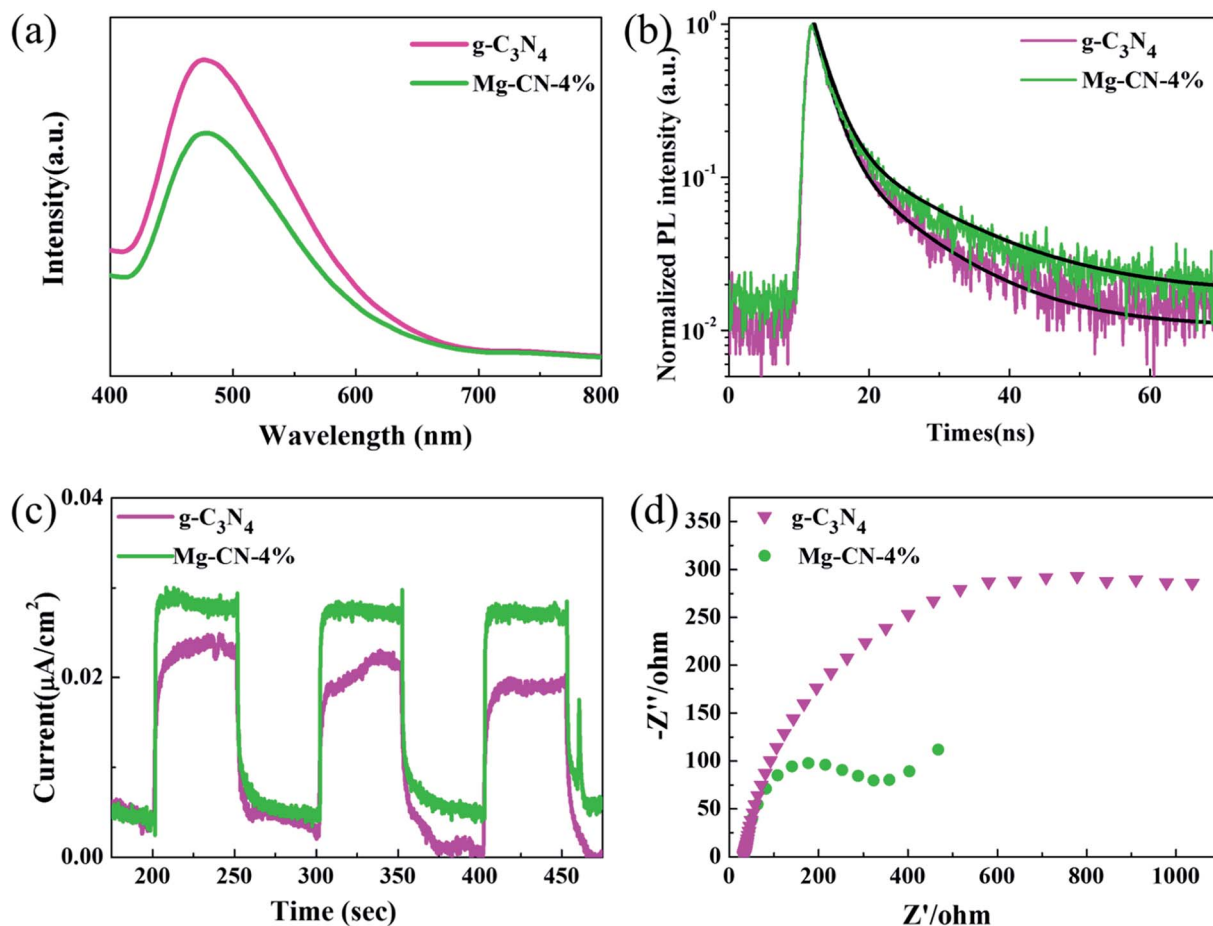


Fig. 4 (a) Photoluminescence spectra, (b) time-resolved PL spectra, (c) transient photocurrent–time curves, (d) electrochemical impedance spectra of $g\text{-C}_3\text{N}_4$ and Mg-CN-4%.

Mg was added, the PL intensity clearly decreases, indicating that Mg-doped $g\text{-C}_3\text{N}_4$ can suppress the recombination of photo-generated electron–hole pairs. The midgap states might act as the center for entrapment of electrons, and enhances photocatalytic activity of catalyst.³⁵ As displayed in Fig. 4(b), a longer recombination lifetime of Mg doped $g\text{-C}_3\text{N}_4$ was examined by time-resolved photoluminescence, indicating that the recombination of carriers was suppressed, which is consistent with the PL results.

Fig. 4(c) shows the transient photocurrent–time ($I-t$) curves for two samples with interval 50 s light on/off cycle of visible light irradiation. Notably, photocurrent intensity of Mg-CN-4% reached $0.028 \mu\text{A cm}^{-2}$, which is higher than that of $g\text{-C}_3\text{N}_4$. The photocurrent intensity sharply increased from the light-off to light-on state, which is attributed to fast photo-generated electrons transport on the sample surface.⁵⁹ Hence, a higher photocurrent response indicates a lower electron–hole recombination. Fig. 4(d) shows electrochemical impedance spectroscopy of $g\text{-C}_3\text{N}_4$ and Mg-CN-4%. It can be clearly observed that the arc radius of EIS Nyquist plot become smaller after doping Mg, which reflects more effective separation of electron–hole pairs.⁶⁰

Evaluation of photocatalytic activity

The ultraviolet photocatalytic activity of $g\text{-C}_3\text{N}_4$ and $g\text{-C}_3\text{N}_4$ doped with different magnesium amount are shown in Fig. 5(a) and (b). Under the combined action of ultraviolet irradiation, water vapor and carbon dioxide, the products of CO and CH_4 were detected by gas chromatography after 8 h photocatalytic reaction, and the accumulation of CO was much higher than that of CH_4 . The optimum doping amount of the samples is Mg-CN-4%, and the yield of CO and CH_4 can reach up to 57.90 and $9.12 \mu\text{mol g}^{-1}$, which is 5.1 and 3.8 times of that produced by $g\text{-C}_3\text{N}_4$. When Mg-CN-2%, Mg-CN-3%, Mg-CN-5% were used as catalysts, the accumulation of CO can reach to 25.35, 47.23, $26.54 \mu\text{mol g}^{-1}$, and the yield of CH_4 is 4.66, 8.25, $5.49 \mu\text{mol g}^{-1}$. The visible light photocatalytic activity of $g\text{-C}_3\text{N}_4$ and Mg-CN-4% is shown in Fig. 5(c). After 8 h of visible light irradiation, the yield of CO were 6.35, $12.97 \mu\text{mol g}^{-1}$, and yield of CH_4 were 2.01 and $7.62 \mu\text{mol g}^{-1}$, respectively.

According to the analysis above, the light utilization efficiency of the $g\text{-C}_3\text{N}_4$ has been improved after Mg doping, which could due to the narrowed down band gap and introduced midgap states. The proper amount of Mg doping can suppress the recombination of photo-induced electron–hole pairs, thus improving photocatalytic activity.



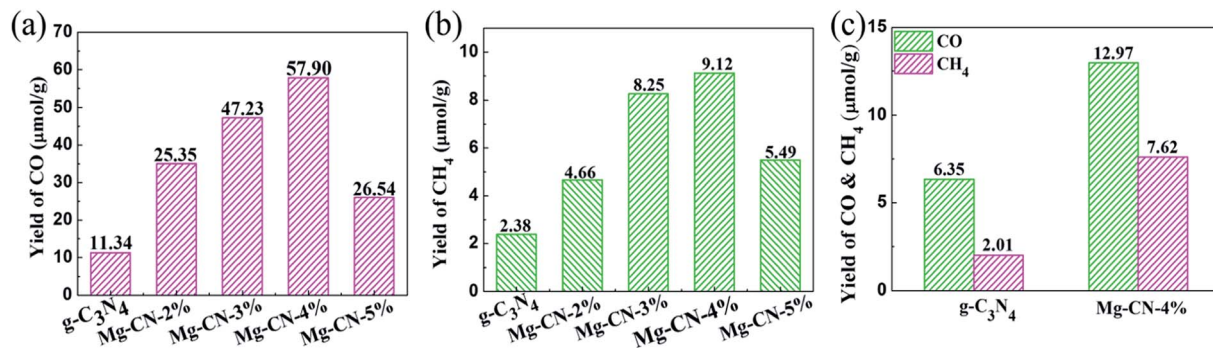


Fig. 5 UV-light photocatalytic activity of g-C₃N₄ and Mg-doped g-C₃N₄ samples (a) the yield of CO, (b) the yield of CH₄; (c) visible-light activity of g-C₃N₄ and Mg-CN-4%.

Conclusion

In summary, a facile one-pot strategy is used for the fabrication of Mg-doped g-C₃N₄ which exhibit considerable UV-light and visible-light photocatalytic CO₂ reduction activity. The optimum amount of the dopant was discovered to be 4 wt%, and its yield of CO and CH₄ were about 5.1 and 3.8 times that of pristine g-C₃N₄ under UV-light, and reached 12.97 and 7.62 μmol g⁻¹ under visible light irradiation. Such kind of enhancement could be attributed to the increased light utilization efficiency as well as the promoted photo-induced charge carrier separation due to the narrowed down band gap and introduced midgap states after effective Mg doping. This work is expected to open up a new insight to design highly active metal-free photocatalysts for solar-light-driven CO₂ reduction.

Experimental

Preparation of bulk g-C₃N₄

A certain amount of melamine was weighed into a crucible, and calcined in a muffle furnace at 550 °C for 4 h at a heating rate of 2.5 °C min⁻¹. After cooling to room temperature, the yellow-colored product was grounded into powder.

Synthesis of Mg-doped g-C₃N₄ catalysts

6.3 g of melamine and a certain amount of MgCl₂·6H₂O were dissolved in 80 mL of an alcohol-water mixed solution (2 : 1 by volume), and ultrasonically dispersed for 1 h, followed by 3 h stirring. The mixture was then dried at 80 °C to remove the solvent. Finally, the solid mixture was placed in a crucible and calcined at 550 °C for 4 h in a muffle furnace. After cooling to room temperature, it was grounded into powder. The synthesized product is labeled as Mg-CN-*X*, where *X* is the molar ratio of Mg to melamine (*X* = 2%, 3%, 4%, 5%).

Characterization

X-Ray diffraction (XRD) patterns were carried out by an Ultima IV X-ray diffractometer. The scanning electron microscopy (SEM) analyses were conducted using a Quanta 3D FEG apparatus. Transmission electron microscopy (TEM) image was

obtained by a JEM-2100 microscope. Fourier transform infrared (FTIR) spectrum was performed on a Nicolet iS50. X-ray photoelectron spectroscopy (XPS) was acquired in Thermo scientific ESCALAB 250Xi X-ray photoelectron spectrometer. The UV-vis diffuse reflectance spectroscopy (DRS) of the samples was measured using an Agilent Cary 5000 spectrometer. The photoluminescence (PL) spectrum of the samples were measured with an F-4600 fluorometer at excitation wavelength of 325 nm. Time-resolved photoluminescence spectra of samples were recorded on a FLS920 Full-featured fluorescence spectrometer at excitation wavelength of 330 nm.

Photoelectrochemical measurements

The photocurrent was measured by using a CHI1030B electrochemical workstation in a standard three-electrode system. A 300 W xenon lamp was used to provide visible light. For the prepared working electrode, 10 mg of the obtained sample was dispersed in ultrapure water, then 1 mL of Nafion solution (0.5 wt%) was added. 10 μL of the suspension was dropped on fluorine-doped tin oxide (FTO) glass, and dried at room temperature. Mott-Schottky plots were measured on the above-mentioned three-electrode system with a frequency of 1000 Hz. The electrochemical impedance spectroscopy (EIS) of samples was performed by using a CHI760D with potassium ferricyanide solution as electrolyte solution.

Photocatalytic experiment

The photocatalytic CO₂ reduction of the samples was carried out in a self-made rectangular reactor. Firstly, 20 mg of the catalyst was plated on the bottom of the reactor, then high purity CO₂ (99.999%) was passed through a water vapor reactor at a flow rate of 15 ml min⁻¹. In order to exhaust air, the mixed gas (CO₂, H₂O) was passed through the photocatalytic reactor for 20 minutes. Finally, close the air outlet, then reduce the flow rate and stop ventilation after 10 minutes. The 300 W xenon lamp was used to provide the UV light and visible light. After irradiation for 8 h, the mixed gas in the reactor was analyzed by a gas chromatograph.



Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the National Natural Science Foundation of China (No. 51702014) and the Fundamental Research Funds for the Central Universities (FRF-AS-17-002).

References

- 1 C. Hu, W. Hung, M. Wang and P. Lu, *Carbon*, 2018, **127**, 374.
- 2 Z. Sun, H. Wang, Z. Wu and L. Wang, *Catal. Today*, 2018, **300**, 160.
- 3 G. Liao, J. Fang, Q. Li, S. Li, Z. Xu and B. Fang, *Nanoscale*, 2019, **11**, 7062.
- 4 A. Fujishima and K. Honda, *Nature*, 1972, **238**, 37.
- 5 A. Khalilzadeh and A. Shariati, *Sol. Energy*, 2018, **164**, 251.
- 6 W. Li, G. Wang, Y. Feng and Z. Li, *Appl. Surf. Sci.*, 2018, **428**, 154.
- 7 J. Hu, H. Li, C. Huang, M. Liu and X. Qiu, *Appl. Catal., B*, 2013, **598**, 142.
- 8 J. Wen, J. Xie, X. Chen and X. Li, *Appl. Surf. Sci.*, 2017, **391**, 72.
- 9 M. Zhou, J. Chen, C. Hou, Y. Liu, S. Xu, C. Yao and Z. Li, *Appl. Surf. Sci.*, 2019, **470**, 908.
- 10 M. Zhang, Z. Luo, M. Zhou, G. Zhang, K. A. Alamry, L. A. Taib, A. M. Asiri and X. Wang, *Appl. Catal., B*, 2017, **210**, 454.
- 11 B. Fang, A. Bonakdarpour, K. Reilly, Y. Xing, F. Taghipour and D. P. Wilkinson, *ACS Appl. Mater. Interfaces*, 2014, **6**, 15488.
- 12 G. Liao, Y. Gong, L. Zhang, H. Gao, G. Yang and B. Fang, *Energy Environ. Sci.*, 2019, DOI: 10.1039/C9EE00717B.
- 13 D. Huang, Z. Li, G. Zeng, C. Zhou, W. Xue, X. Gong, X. Yan, S. Chen, W. Wang and M. Cheng, *Appl. Catal., B*, 2019, **240**, 153.
- 14 S. Hu, L. Ma, J. You, F. Li, Z. Fan, G. Lu, D. Liu and J. Gui, *Appl. Surf. Sci.*, 2014, **311**, 164.
- 15 Y. Zeng, H. Li, J. Luo, J. Yuan, L. Wang, C. Liu, Y. Xia, M. Liu, S. Luo, T. Cai, S. Liu and J. C. Crittenden, *Appl. Catal., B*, 2019, **249**, 275.
- 16 J. Tang, R. Guo, W. Zhou, C. Huang and W. Pan, *Appl. Catal., B*, 2018, **237**, 802.
- 17 M. Zhou, G. Dong, F. Yu and Y. Huang, *Appl. Catal., B*, 2019, **256**, 1037.
- 18 G. Li, B. Wang, J. Zhang, R. Wang and H. Liu, *Appl. Surf. Sci.*, 2019, **478**, 1056.
- 19 Z. Xiong, H. Wang, N. Xu, H. Li, B. Fang, Y. Zhao, J. Zhang and C. Zheng, *Int. J. Hydrogen Energy*, 2015, **40**, 10049.
- 20 J. Wu, N. Li, X. Zhang, H. Fang, Y. Zheng and X. Tao, *Appl. Catal., B*, 2018, **226**, 61.
- 21 Y. Wang, S. Zhao, Y. Zhang, J. Fang, Y. Zhou, S. Yuan, C. Zhang and W. Chen, *Appl. Surf. Sci.*, 2018, **440**, 258.
- 22 S. Le, T. Jiang, Q. Zhao, X. Liu, Y. Li, B. Fang and M. Gong, *RSC Adv.*, 2016, **6**, 38811.
- 23 N. Wang, J. Wang, J. Hu, X. Lu, J. Sun, F. Shi, Z. Liu, Z. Lei and R. Jiang, *ACS Appl. Energy Mater.*, 2018, **1**, 2866.
- 24 J. Wang, C. Cui, Q. Kong, C. Ren, Z. Li, L. Qu, Y. Zhang and K. Jiang, *Chem. Eng. J.*, 2019, **359**, 723.
- 25 S. C. Yan, Z. S. Li and Z. G. Zou, *Langmuir*, 2010, **26**, 3894.
- 26 Y. Zhou, L. Zhang, W. Huang, Q. Kong, X. Fan, M. Wang and J. Shi, *Carbon*, 2016, **99**, 111.
- 27 G. Liu, P. Niu, C. Sun, S. C. Smith, Z. Chen, G. Lu and H. Cheng, *J. Am. Chem. Soc.*, 2010, **132**, 11642.
- 28 M. Bellardita, E. I. García-López, G. Marci, I. Krivtsov and J. R. García, *Appl. Catal., B*, 2018, **220**, 222.
- 29 Y. Hong, Y. Jiang, C. Li, W. Fan, X. Yan, M. Yan and W. Shi, *Appl. Catal., B*, 2016, **180**, 663.
- 30 M. Wang, M. Shen, L. Zhang, J. Tian, X. Jin, Y. Zhou and J. Shi, *Carbon*, 2017, **120**, 23.
- 31 M. Li, L. Zhang, M. Wu, Y. Du, X. Fan, M. Wang, L. Zhang, Q. Kong and J. Shi, *Nano Energy*, 2016, **19**, 145.
- 32 L. Zhou, L. Wang, J. Lei, Y. Liu and J. Zhang, *Catal. Commun.*, 2017, **89**, 125.
- 33 B. Fang, Y. Xing, A. Bonakdarpour, S. Zhang and D. P. Wilkinson, *ACS Sustainable Chem. Eng.*, 2015, **3**, 2381.
- 34 L. Ye, D. Wu, K. H. Chu, B. Wang, H. Xie, H. Y. Yip and P. K. Wong, *Chem. Eng. J.*, 2016, **304**, 376.
- 35 A. Kumar, R. K. Yadav, N. Park and J. Baeg, *ACS Appl. Nano Mater.*, 2018, **1**, 47.
- 36 J. Tang, W. Zhou, R. Guo, C. Huang and W. Pan, *Catal. Commun.*, 2018, **107**, 92.
- 37 S. Liu, H. Zhu, W. Yao, K. Chen and D. Chen, *Appl. Surf. Sci.*, 2018, **430**, 309.
- 38 Q. Yan, G. Huang, D. Li, M. Zhang, A. Pan and W. Huang, *Mater. Sci. Technol.*, 2018, **34**, 2515.
- 39 X. Wu, J. Cheng, X. Li, Y. Li and K. Lv, *Appl. Surf. Sci.*, 2019, **465**, 1037.
- 40 F. Guo, W. Shi, C. Zhu, H. Li and Z. Kang, *Appl. Catal., B*, 2018, **226**, 412.
- 41 W. Fang, J. Liu, L. Yu, Z. Jiang and W. Shangguan, *Appl. Catal., B*, 2017, **209**, 631.
- 42 Q. Li, Z. Sun, H. Wang and Z. Wu, *J. CO₂ Util.*, 2018, **28**, 126.
- 43 S. Hua, L. Ma, J. You, F. Li, Z. Fan, G. Lu, D. Liu and J. Gui, *Appl. Surf. Sci.*, 2014, **311**, 164.
- 44 Z. Sun, J. M. Theresa Agatha Fischer, Q. Li, J. Hu, Q. Tang, H. Wang, Z. Wu, M. Hankel, D. J. Searles and L. Wang, *Appl. Catal., B*, 2017, **216**, 146.
- 45 W. Oh, V. W. C. Chang, Z. Hu, R. Goei and T. Lim, *Chem. Eng. J.*, 2017, **323**, 260.
- 46 B. Yue, Q. Li, H. Iwai, T. Kako and J. Ye, *Sci. Technol. Adv. Mater.*, 2011, **12**, 034401.
- 47 E. Liu, J. Chen, Y. Ma, J. Feng, J. Jia, J. Fan and X. Hu, *J. Colloid Interface Sci.*, 2018, **524**, 313.
- 48 F. Raziq, Y. Qu, M. Humayun, A. Zada, H. Yu and L. Jing, *Appl. Catal., B*, 2017, **201**, 486.
- 49 M. A. Behnajady, B. Alizade and N. Modirshahla, *Photochem. Photobiol.*, 2011, **87**, 1308.
- 50 J. Liu, H. Yang, W. Tan, X. Zhou and Y. Lin, *Electrochim. Acta*, 2014, **129**, 459.
- 51 H. Huang, K. Xiao, N. Tian, F. Dong, T. Zhang, X. Du and Y. Zhang, *J. Mater. Chem. A*, 2017, **5**, 17452.



- 52 S. Pany and K. M. Parida, *Phys. Chem. Chem. Phys.*, 2015, **17**, 8070.
- 53 X. Wang, S. Blechert and M. Antonietti, *ACS Catal.*, 2012, **2**, 1596.
- 54 H. Yaghoubi, Z. Li, Y. Chen, H. T. Ngo, V. R. Bhethanabotla, B. Joseph, S. Ma, R. Schlaf and A. Takshi, *ACS Catal.*, 2015, **5**, 327.
- 55 L. Jiang, X. Yuan, G. Zeng, J. Liang, Z. Wu, H. Yu, D. Mo, H. Wang, Z. Xiao and C. Zhou, *J. Colloid Interface Sci.*, 2019, **536**, 17.
- 56 J. Ran, T. Y. Ma, G. Gao, X. Du and S. Z. Qiao, *Energy Environ. Sci.*, 2015, **8**, 3708.
- 57 R. Ding, S. Cao, H. Chen, F. Jiang and X. Wang, *Colloids Surf., A*, 2019, **563**, 263.
- 58 W. Tu, Y. Xu, J. Wang, B. Zhang, T. Zhou, S. Yin, S. Wu, C. Li, Y. Huang, Y. Zhou, Z. Zou, J. Robertson, M. Kraft and R. Xu, *ACS Sustainable Chem. Eng.*, 2017, **5**, 7260.
- 59 J. Zhang, S. Liu, J. Yu and M. Jaroniec, *J. Mater. Chem.*, 2011, **21**, 14655.
- 60 L. Yang, J. Huang, L. Shi, L. Cao, H. Liu, Y. Liu, Y. Li, H. Song, Y. Jie and J. Ye, *Appl. Catal., B*, 2018, **221**, 670.

