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# Construction of CeCu<sub>x</sub>-BTC/CN S-type heterojunctions and photocatalytic CO<sub>2</sub> reduction to CO and CH<sub>4</sub>

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S-type heterojunction photocatalysts (CeCu<sub>x</sub>-BTC/CN) of cerium-copper bimetallic organic framework (CeCu<sub>x</sub>-BTC) and graphitic carbon nitride (g-C<sub>3</sub>N<sub>4</sub>) were constructed by a simple solvothermal method using cerium nitrate, copper nitrate, and urea as the raw materials, and 1,3,5-benzene-tricarboxylic acid as the ligand for the photocatalytic CO<sub>2</sub> reduction to CO and CH<sub>4</sub>. The results show that the built-in electric field constructed by Fermi energy level flattening transfers the electrons in an S-type manner, which not only preserves the strong reducing properties of the electrons in the material but also provides the maximum redox capacity and enables the composite samples to obtain higher visible-light trapping capacity and improve the separation efficiency of the carriers while refining the crystal particles. With the addition of only 1 mL of H<sub>2</sub>O as the proton supply source, CeCu<sub>0.05</sub>-BTC/CN exhibits the optimal photocatalytic performance. The CO and CH<sub>4</sub> yields were 64.44 and 0.5575 μmol g<sup>-1</sup>, which were 7.56 and 2.42 times higher than those of g-C<sub>3</sub>N<sub>4</sub>, respectively, and the catalytic performances were basically stable after cycling tests.

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## 1 Introduction

Since the industrial revolution, the greenhouse effect has become more pronounced as carbon dioxide emissions have continued to increase, and the frequency of extreme weather events has led to increasingly severe global environmental problems.<sup>1,2</sup> The emission of fossil fuels (coal, oil, natural gas, *etc.*) is the main source of the increase of carbon dioxide in the atmosphere.<sup>3</sup> Therefore, it is imperative to reduce the consumption of fossil fuels and the content of carbon dioxide in the atmosphere. Carbon dioxide reduction using solar-driven semiconductor photocatalysts is an efficient and promising technology for achieving long-term solar-fuel conversion.<sup>4,5</sup> Graphite carbon nitride (g-C<sub>3</sub>N<sub>4</sub>) is a 2D layered semiconductor,<sup>6</sup> which is often regarded as a very promising semiconductor material due to its suitable band gap energy, good chemical-thermal stability, and low cost.<sup>7-9</sup> However, the small specific surface area, the presence of rapid recombination of photogenerated electrons and holes, and the limited absorption and photoresponsivity to visible light limit the practical use of this material for photocatalytic carbon dioxide reduction

applications. At present, several aspects of catalyst modification are common in the market, such as chemical doping,<sup>10</sup> the construction of heterojunction,<sup>11</sup> noble metal loading,<sup>12</sup> and the regulation of its morphology.<sup>13</sup> These include metal oxides, sulfides, carbon-nitrides, metal-organic skeletons (MOFs), *etc.*<sup>14,15</sup>

During research in the semiconductor field, MOFs have attracted the attention of a wide range of researchers due to their diverse and controllable crystal structure, ultra-high specific surface area,<sup>16,17</sup> and tunable pores.<sup>18</sup> Cerium (Ce) is an excellent catalytic additive as a rare earth element with unfilled electron 4f orbitals, capable of reversible changes between Ce<sup>4+</sup> and Ce<sup>3+</sup>, and surface oxygen vacancy defects that can be spontaneously generated and eliminated. Ye *et al.*<sup>19</sup> obtained oxygen vacancy-rich and N-doped MOF-derived electrocatalysts by pyrolysis of melamine-containing CuCe-BTC with high selectivity and good electrocatalytic activity compared to a reversible hydrogen electrode with a NH<sub>3</sub> yield of 44.5 μg h<sup>-1</sup> mg<sub>cat.</sub><sup>-1</sup> at 0.5 V. Zhang *et al.*<sup>20</sup> prepared CuO/CeO<sub>2</sub> catalysts synthesized from MOFs using *in situ* synthesis, mechanical mixing, and impregnation methods and found that the CuCeO-ETH catalysts had higher than 98% selectivity for CO oxidation and a wide temperature activity window. Zhang *et al.*<sup>21</sup> achieved efficient CO<sub>2</sub>-CH<sub>4</sub> electroconversion with a methane local current density of 138.6 mA cm<sup>-2</sup> by pyrolysis of a metal-organic framework (MOF) precursor for the preparation of cerium carbide-copper oxide complexes (Cu/CeO<sub>2</sub>@C). Therefore, the heterostructured photocatalytic system constructed

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from the two may better improve the photocatalyst activity and CO<sub>2</sub> reduction efficiency.

Currently, although some Cu-BTC, Ce-BTC and related materials of CeCu-BTC from different preparation methods have been synthesized, little research has been reported on bimetallic organic frameworks (CeCu-BTC) and graphite carbon nitride (g-C<sub>3</sub>N<sub>4</sub>) composite heterostructures for photocatalytic carbon dioxide reduction. In the present study, composite heterostructures of bimetallic organic frameworks (CeCu<sub>x</sub>-BTC) and graphite carbon nitride (g-C<sub>3</sub>N<sub>4</sub>) were synthesized by a simple solvothermal method using cerium nitrate and urea as raw materials and 1,3,5-benzotricarboxylic acid as a ligand. The results show that Ce-BTC microrods surrounded by Cu-doped microspheres can form beneficial MOFs materials to synergistically improve the photocatalytic CO<sub>2</sub> reduction activity, and this finding also paves the way for further improvement of the photocatalytic CO<sub>2</sub> reduction performance and a small amount of low-cost composite modification of g-C<sub>3</sub>N<sub>4</sub>.

## 2 Experimental section

### 2.1 Materials

Cerium nitrate hexahydrate (Ce(NO<sub>3</sub>)<sub>3</sub>·6H<sub>2</sub>O), copper nitrate trihydrate (Cu(NO<sub>3</sub>)<sub>2</sub>·3H<sub>2</sub>O), *N,N*-dimethylformamide (C<sub>3</sub>H<sub>7</sub>NO), 1,3,5-benzene-tricarboxylic acid (C<sub>9</sub>H<sub>6</sub>O<sub>6</sub>), all analytically pure AR, Aladdin Reagents Ltd urea (CH<sub>4</sub>N<sub>2</sub>O), anhydrous ethanol (C<sub>2</sub>H<sub>6</sub>O), both analytically pure AR, Sino-pharm Group Chemical Reagent Co. Ltd acetone (C<sub>3</sub>H<sub>6</sub>O), analytically pure AR, Tianjin Jindong Tianzheng Fine Chemical Reagent Factory.

### 2.2 Preparation of g-C<sub>3</sub>N<sub>4</sub>, Ce-BTC and Cu-BTC

g-C<sub>3</sub>N<sub>4</sub> and Ce-BTC were prepared by simple thermal polymerization and solvothermal methods, respectively. The temperature was raised to 550 °C at a rate of 5 °C min<sup>-1</sup> and held for 4 hours. After grinding, a light yellow g-C<sub>3</sub>N<sub>4</sub> powder was obtained.

3 mmol (0.63 g) of 1,3,5-benzene tricarboxylic acid (H<sub>3</sub>BTC) and 10 mmol (2.17 g) of Ce(NO<sub>3</sub>)<sub>3</sub>·6H<sub>2</sub>O were dissolved in 50 mL (1 : 1) of a mixture of dimethylformamide (DMF) and ethanol, and the mixture was ultrasonicated for 30 min and then sealed in an autoclave at 100 mL, held at 130 °C for 24 h, naturally cooled to room temperature, washed sufficiently, and then dried under vacuum at 80 °C for 12 h. The Ce-BTC powder material was ground. Cu-BTC was prepared by replacing Ce(NO<sub>3</sub>)<sub>3</sub>·6H<sub>2</sub>O with 0.13 g of Cu(NO<sub>3</sub>)<sub>2</sub>·3H<sub>2</sub>O as described above.

### 2.3 Preparation of CeCu<sub>x</sub>-BTC/CN composite photocatalysts

CeCu<sub>x</sub>-BTC/CN composite photocatalysts were prepared by a simple solvent-thermal method. The CeCu<sub>x</sub>-BTC/CN composite photocatalysts were prepared similarly to Ce-BTC, except that Cu(NO<sub>3</sub>)<sub>2</sub>·3H<sub>2</sub>O and g-C<sub>3</sub>N<sub>4</sub> were added to the reaction solution. As an example, CeCu<sub>0.05</sub>-BTC/CN was weighed, and 0.63 g of 1,3,5-benzotricarboxylic acid (H<sub>3</sub>BTC), 2.17 g of Ce(NO<sub>3</sub>)<sub>3</sub>·6H<sub>2</sub>O, and 0.13 g of Cu(NO<sub>3</sub>)<sub>2</sub>·3H<sub>2</sub>O were dissolved in a 50 mL (1 : 1) mixture of dimethylformamide

(DMF) and ethanol. And 500 mg of g-C<sub>3</sub>N<sub>4</sub> was taken in a certain amount of anhydrous ethanol. After ultrasonic dispersion for 30 min, the two were mixed and strongly stirred and sealed in a 100 mL polytetrafluoroethylene stainless steel autoclave at 130 °C for 24 h, naturally cooled to room temperature and centrifuged, washed several times with anhydrous ethanol and acetone, and then vacuum-dried for 12 h at 80 °C and milled to obtain the CeCu<sub>0.05</sub>-BTC/CN powder material. Wherein, the obtained CeCu<sub>x</sub>-BTC/g-C<sub>3</sub>N<sub>4</sub> catalysts were named as CeCu<sub>0.05</sub>-BTC/CN, CeCu<sub>0.1</sub>-BTC/CN, CeCu<sub>0.15</sub>-BTC/CN, CeCu<sub>0.2</sub>-BTC/CN, respectively. CeCu-BTC was prepared with *x* = 0.05 as an example, and in all the synthesised CeCu<sub>x</sub>-BTC/CN, *x* denotes a Cu content of 5%.

### 2.4 Characterization

The crystal structure was analyzed using an X-ray diffraction analyzer with a scanning range of 5 to 70°, Bruker diffractometer type D8-ADVANCE, Germany. GAIA3(XMU/XMH) Ga-ion double beam scanning electron microscope (FIB-SEM) and high-resolution field emission scanning electron microscope Hitachi SU8010 to observe the surface micro-morphology of Czechoslovakia and Japan. Transmission electron microscopy (TEM) images were taken with a FEI-Talos F200S instrument with an accelerating voltage of 200 kV, energy spectrum: X-Twin, origin Netherlands. FT-IR spectrometer to analyze the molecular structure and chemical components of the samples, NICOLET iS50FT-IR type. X-ray electron spectroscopy to analyze the elemental composition and chemical state of the samples, ESCALAB 205Xi type, all of the above are from Thermo Fisher Scientific, USA. UV-Vis Diffuse Reflectance Spectroscopy, test wavelength range 200–800 nm, Model U-3900, Hitachi High-Technologies Corporation, Japan. PL Fluorescence Spectroscopy, testing wavelength of 285 nm, Model F-4600, Hitachi High-Technologies, Japan. Beijing Perfectlight High Performance Xenon Light Source simulating daylight (PLS-SXE300E/UV), power 300 W, Beijing Perfectlight Technology Co. Fully automated system gas chromatography for photocatalytic carbon dioxide reduction testing, Model GC79720, Beijing CEC Jinyuan Technology Co. Electrochemical workstation for electrochemical performance testing of samples, Model 1260A+1287A, Sully Strong, UK. EPR electron paramagnetic resonance spectrometer using the Bruker model E500, Germany.

### 2.5 Photocatalytic CO<sub>2</sub> reduction

The photocatalytic CO<sub>2</sub> reduction experiments were carried out in a sealed 250 mL quartz glass reactor with a 28# standard ball mill and flange O-ring seal, which was connected to an automatic on-line photocatalytic analysis system using a PLS-SXE300 xenon lamp with a wavelength of 320–780 nm as the light source. Specifically, 50 mg of photocatalyst was put into the reaction bottle to build a platform, and 1 mL of deionized water was used as the reaction solution. Before illumination, vacuum the reaction system 2–3 times until the pressure gauge shows −91.7 kPa, and then charge pure CO<sub>2</sub> gas with high-purity argon as the carrier gas. Use the circulating water



chiller to maintain the temperature in the bottle and carry out the photocatalytic reaction under stable temperature conditions. The trace gas in the reaction bottle was extracted by automatic sampling every 60 min, and the yield of CO and CH<sub>4</sub> was obtained by a GC7920 gas chromatograph (equipped with FID, TCD, ECD, and FPD detectors).

## 2.6 Electrochemical testing

The electrochemical tests were carried out using a three-electrode system. Glymercury electrode was used as a reference electrode, a platinum electrode as a counter electrode, a catalyst membrane electrode as a working electrode, and a 0.2 mol L<sup>-1</sup> Na<sub>2</sub>SO<sub>4</sub> aqueous solution as an electrolyte. Catalyst membrane electrode: 10 mg of photocatalyst was weighed and mixed with 0.2 mL of ethanol, 0.7375 mL of water, and 0.0625 mL of 5% Nafion solution and sonicated for at least 30 min. Next, using a micropipette, 10 μL of the combined solution was applied to a 4 cm<sup>2</sup> conductive glass surface, creating a center area with a volume of 1 cm<sup>3</sup>, which was then left to dry naturally. The transient photocurrent test potential in the photocurrent test was 0.1 V, and the test time was 540 s with a light–dark cycle of 60 s. Electrochemical impedance frequency range 100 kHz to 0.01 Hz.

## 2.7 Electron paramagnetic resonance (EPR)

5,5-Dimethyl-1-pyrroline-N-oxide (DMPO) was used as a spin trapping agent and methanol as a hole-sacrificing agent at room temperature and atmospheric pressure. Mix the solution to be tested with 100 mm of capturing agent in a ratio of 1 : 1 to obtain 1 mL of the mixed solution. After full mixing, suck 30 μL with a capillary glass tube and put it into the EPR-specific test tube. Record the spectral signal in dark and light conditions for 15 minutes.

# 3 Results and discussion

## 3.1 Characterization of physical phases and micromorphology

Fig. 1(a–c) shows the XRD spectra of the samples, the bright diffraction peaks of the samples indicate that all the samples are well crystallized. There are two distinct diffraction peaks at 13.05° and 27.34° for g-C<sub>3</sub>N<sub>4</sub>, corresponding to the (100) and (002) crystal planes of the graphite-like phase, respectively, which are attributed to the in-plane stacking and interlayer stacking of the typical triazine ring in g-C<sub>3</sub>N<sub>4</sub> (ref. 22) (JCPDS 87-1526). The stronger diffraction peak crystal spacing is typically 0.326 nm. The diffraction peaks of Ce-BTC were 8.5°, 10.6°, and 18.1°, respectively, and the XRD data were consistent with the previous report<sup>23</sup> (CCDC ID: 773084), in which the peaks of CeCu-BTC and Ce-BTC remained consistent. It is noteworthy that a part of CeO<sub>2</sub> was generated in the samples under hydrothermal conditions, with diffraction peaks of 28.55°, 33.09°, and 47.49°, corresponding to the (111), (200), and (220) crystal planes of CeO<sub>2</sub>, respectively. The Cu-BTC diffraction peaks were 6.7°, 9.5°, 11.6°, 13.5°, and 19.3°, respectively. Some of the synthesised samples contain very small amounts of

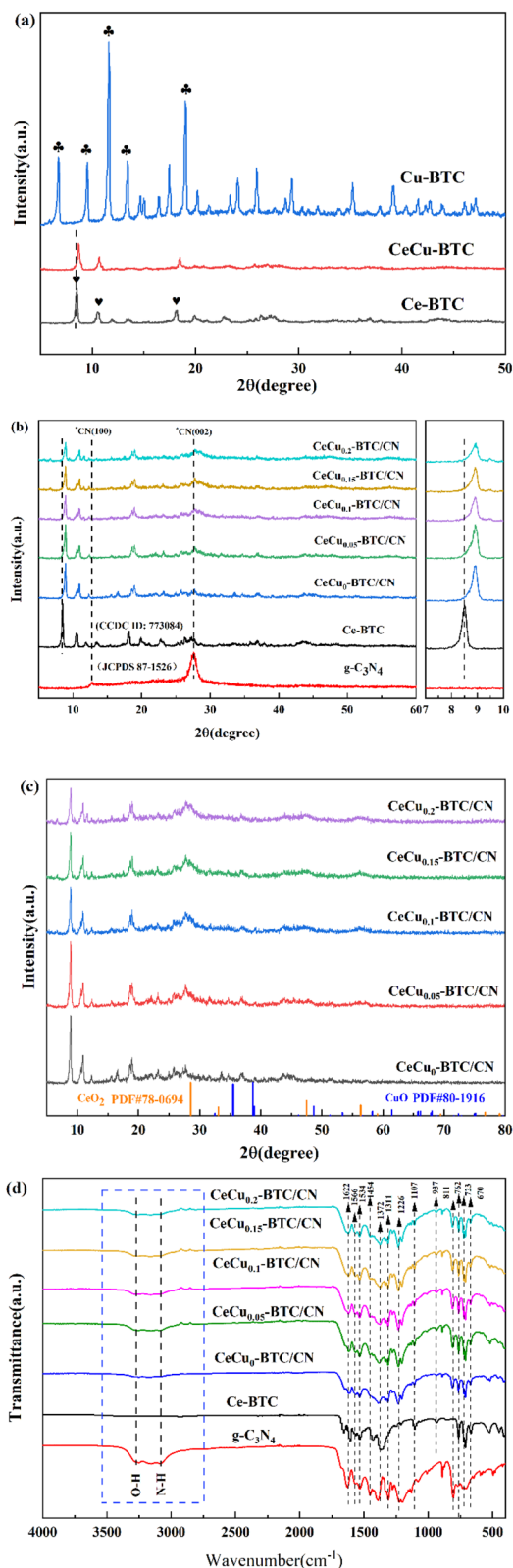


Fig. 1 XRD patterns of Ce-BTC, Cu-BTC and CeCu-BTC (a), g-C<sub>3</sub>N<sub>4</sub>, Ce-BTC and CeCu<sub>x</sub>-BTC/CN (b), XRD spectra of CeO<sub>2</sub> and CuO (c), FT-IR spectra of g-C<sub>3</sub>N<sub>4</sub>, Ce-BTC and CeCu<sub>x</sub>-BTC/CN (d).



copper oxide impurities, and the diffraction peaks at  $35.5^\circ$  and  $38.9^\circ$  correspond to the (111) and (200) crystal planes of CuO, respectively.<sup>19,20</sup> Due to the difference between the ion radii of  $\text{Ce}^{3+}$  (1.02 Å) and  $\text{Cu}^{2+}$  (0.73 Å), the doping of Cu causes the XRD pattern to move in a higher  $2\theta$  direction, indicating that the nodes of the  $\text{Ce}^{3+}$  skeleton are partially and isomorphically replaced by  $\text{Cu}^{2+}$  ions to form CeCu-BTC. With the increasing amount of Cu doping in the heterojunction, the corresponding (101) diffraction peak intensity gradually decreases, the full width of the half-peak increases, and the grain size gradually decreases, indicating that the crystal growth process of Ce-BTC is hindered by the high content of Cu,<sup>24</sup> and the refined grain size also helps to increase the specific surface area of the composite photocatalyst, which in turn improves its photocatalytic performance.

Fig. 1(d) shows the FT-IR spectra of the samples, and all  $\text{CeCu}_x\text{-BTC/CN}$  samples show almost the same absorption bands, and the broad absorption bands at  $3500\text{--}2820\text{ cm}^{-1}$  can be attributed to the bonded or physically adsorbed hydroxyl ( $\text{-OH}$ ) stretching vibrations of the bonded or physically adsorbed water molecules and the N-H bond stretching vibrations.<sup>25</sup> In the characteristic frequency region, the absorption bands at  $1625\text{--}1534\text{ cm}^{-1}$  and  $1454\text{--}1372\text{ cm}^{-1}$  belong to the asymmetric stretching vibration (vas) and symmetric stretching vibration (vs) of the carboxyl group ( $\text{O-C=O}$ ) in homobenzoic acid,<sup>26,27</sup> respectively. In the fingerprint region, the absorption bands at  $1300\text{--}1000\text{ cm}^{-1}$  belong to C-O stretching vibration, the absorption band at  $1107\text{ cm}^{-1}$  corresponds to C-H in-plane bending vibration, and the absorption peak at  $810\text{ cm}^{-1}$  corresponds to the out-of-plane bending vibration of the heptane-triazole ring in  $g\text{-C}_3\text{N}_4$ . It is also found that the absorption intensity of the peak at this location gradually increases with the increase of Cu content in the heterojunction. The absorption bands at  $767$  and  $721\text{ cm}^{-1}$  correspond to C-C ring out-of-plane bending vibrations, which confirm the aromatic ring structure of the BTC ligand.<sup>28</sup> The absorption band between  $700$  and  $450\text{ cm}^{-1}$  can be attributed to the Ce-O stretching vibration of Ce-BTC samples.

By comparison, it can be seen that the organic ligand BTC has formed a bridge structure with rare earth ions and Cu ions. For the offset phenomenon of some absorption peaks, it also shows that the compound of BTC and  $g\text{-C}_3\text{N}_4$  has a significant impact on the chemical environment of the  $g\text{-C}_3\text{N}_4$  molecule, which fully shows that the two have successfully constructed a heterojunction, which is consistent with the XRD analysis results.

Fig. 2 is the SEM image of the sample. Fig. 2(a) shows that  $g\text{-C}_3\text{N}_4$  presents a clearly visible irregular two-dimensional folded layered structure, and the adjacent thin sections may show a good pore structure.<sup>29</sup> Ce-BTC in Fig. 2(b) shows a typical rod- or bar-like structure with different sizes and lengths, mostly around  $15\text{--}20\text{ }\mu\text{m}$ , presenting a smooth and textured surface, and the crystal edges can be clearly seen. Fig. 2(c) shows the tight and ordered stacking of Ce-BTC and  $g\text{-C}_3\text{N}_4$  cladding after the sonicated  $g\text{-C}_3\text{N}_4$  structure has been dispersed. Some surface aggregates of copper oxide with microspherical structure and micron flower-like  $g\text{-C}_3\text{N}_4$  are

shown in Fig. 2(d). In addition, when the Cu content was further increased, such as in  $\text{CeCu}_{0.1}\text{-BTC/CN}$  samples, more Cu-BTC particles appeared (Fig. 2(h)), but the high content of Cu caused the crystal growth of Ce-BTC to be hindered, which might lead to a corresponding decrease in the particle size of Cu-BTC (Fig. 2e-g).<sup>30</sup> To sum up, Cu-BTC and  $g\text{-C}_3\text{N}_4$  are evenly dispersed in the structure of Ce-BTC to form a relatively complex structure. It is inferred that heterostructures may be formed and play a certain role in promoting the separation of charges in photoexcitation.<sup>31</sup>

Fig. 3 shows the TEM image of the sample, and Fig. 3(a) shows that some coalesced spherical Cu-BTC and rod-shaped Ce-BTC are uniformly distributed on the smooth carbon nitride surface and are in close contact on the surface of  $g\text{-C}_3\text{N}_4$ . Further, it is clearly proved that Cu in Ce-BTC is successfully doped and coexists with  $g\text{-C}_3\text{N}_4$ , forming a more complex structure, which proves that the sample is successfully compounded. As reported in previous studies,  $g\text{-C}_3\text{N}_4$ , with its functional amino groups and large specific surface area, is also a good carrier for loading nanocatalysts, which is more favorable for the construction of heterojunction<sup>32,33</sup> structure. Thus, in summary, it was confirmed that the  $\text{CeCu}_x\text{-BTC/CN}$  composite heterojunction was successfully constructed, and the composition of this heterojunction may be conducive to the rapid transfer of photoexcited electrons at the interface,<sup>34</sup> which may improve charge migration and photocatalytic efficiency. Fig. 3(b and c) clearly show lattice spacings of  $0.32$ ,  $0.31$ ,  $0.19$ , and  $0.21\text{ nm}$  corresponding to the (002) crystallographic plane of  $g\text{-C}_3\text{N}_4$ , the (111) and (220) crystallographic planes of  $\text{CeO}_2$ , and the (200) crystallographic plane of CuO grains, with well-matched lattice planes and good agreement with the XRD data. Fig. 3(d) shows its dark field image. By collecting scattered (diffracted) electronic signals for imaging, it can be seen that the larger and thicker the sample, the stronger the scattering and the brighter the sample area in the dark field. Fig. 3(e-h) clearly shows the elemental distribution of the samples, with brighter points in the elemental mapping region indicating higher concentrations of the matched elements. It is worth noting that the distribution of C, N, O, Ce, and Cu elements is relatively uniform. The total distribution of each element is shown in Fig. 3(i), which indicates the good coexistence of Ce-BTC, Cu-BTC, and  $g\text{-C}_3\text{N}_4$ , and it also indicates that the structure of the prepared samples is well dispersed and thermally stable, and it also proves once again that the samples are successful in constructing heterojunctions.

Fig. 4 shows the surface element and chemical valence composition of  $g\text{-C}_3\text{N}_4$ , CeCu-BTC, and  $\text{CeCu}_{0.05}\text{-BTC/CN}$ . Pollution carbon ( $284.8\text{ eV}$ ) was selected as the internal standard for all elements in the figure, and the spectral correction has been completed. Fig. 4(a) shows the full spectrum of the sample. It is found that there are mainly energy spectrum peaks related to C 1s, N 1s, and O 1s elements in  $g\text{-C}_3\text{N}_4$ . CeCu-BTC and  $\text{CeCu}_{0.05}\text{-BTC/CN}$  mainly have energy spectrum peaks related to C 1s, N 1s, O 1s, Ce 3d, and Cu 2p elements without other impurities, which corresponds to XRD data. The element content in  $\text{CeCu}_{0.05}\text{-BTC/CN}$  also indicates that  $g\text{-C}_3\text{N}_4$  and  $\text{CeCu}_x\text{-BTC}$  are successfully combined.



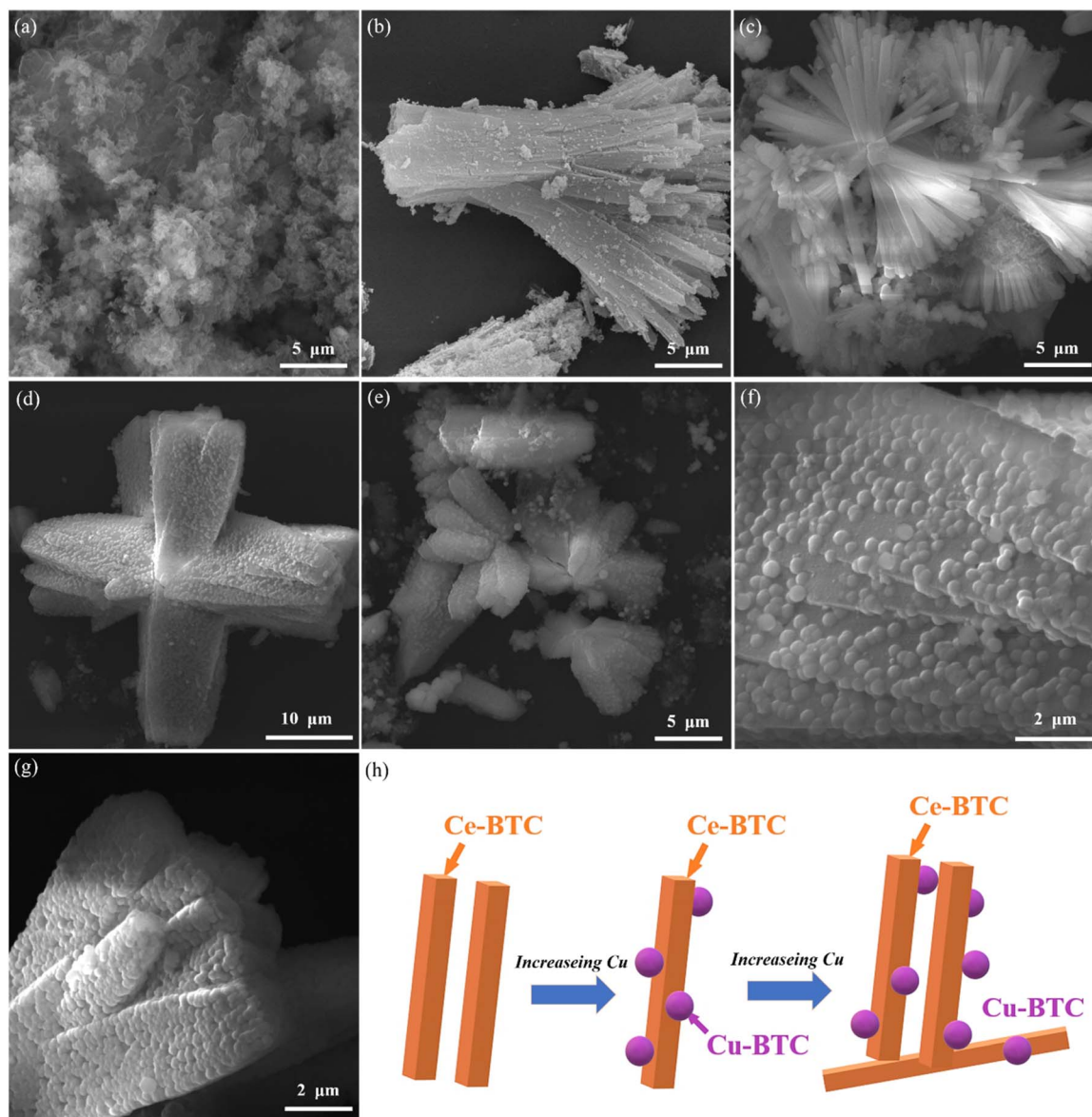


Fig. 2 SEM images of  $g\text{-C}_3\text{N}_4$  (a), Ce-BTC (b),  $\text{CeCu}_0\text{-BTC/CN}$  (c),  $\text{CeCu}_{0.05}\text{-BTC/CN}$  (d),  $\text{CeCu}_{0.1}\text{-BTC/CN}$  (e),  $\text{CeCu}_{0.05}\text{-BTC/CN}$  sample enlarged view (f),  $\text{CeCu}_{0.1}\text{-BTC/CN}$  sample enlarged view (g), schematic diagram of sample crystal growth (h).

As shown in Fig. 4(b), the C 1s spectrum is deconvoluted into three characteristic peaks, of which the characteristic peak of 284.8 eV corresponds to the amorphous carbon (C-C) of the sample, and the characteristic peaks at 285.7 eV and 288.12 eV correspond to C-O-C and C-C=O<sup>35,36</sup> respectively. The characteristic peak at 287.84 eV corresponds to the characteristic peak of SP<sup>2</sup> hybridized carbon in  $g\text{-C}_3\text{N}_4$  (N=C=N),<sup>19</sup> The peak is shifted by 0.28 eV to reach 288.12 eV at high binding energy in the composite samples, suggesting that there is a change in charge transport or chemical environment at the surface of the triazine ring, which may be related to the introduction of  $\text{CeCu}_x\text{-BTC}$ , and also indicates that the density of the electron cloud around the Ce and Cu atoms is increased. Fig. 4(c) shows that the peak of the N 1s energy level appears at the C-N binding energy of 398.35 eV, which corresponds to the pyridine nitrogen

in the tricyclic (C=N-C) shown in Fig. 4(d) at binding energies of 529.6 eV, 531.63 eV, and 533.1 eV, respectively, can be classified into three component peaks: lattice oxygen (Olat), surface chemisorbed oxygen (OAds), and surface reactive oxygen (OSur), which correspond positively to C=O and C-O.<sup>19,36</sup> Among them, surface active oxygen (OSur) is mainly classified as metal oxide-bonded (Cu-O or Ce-O), which is derived from oxygen defects with lower coordination properties or weaker surface oxygen bonding and possesses better oxidative activity against toluene at low temperatures,<sup>37,38</sup> so it is speculated that the fabricated samples may possess better catalytic activity. As an example, Ce 3d of  $\text{CeCu}_{0.05}\text{-BTC/CN}$  in Fig. 4(e). The Ce 3d spectrum is deconvoluted into two spin-orbit peaks v and u and assigned to Ce 3d<sub>3/2</sub> vs. Ce 3d<sub>5/2</sub>. Notate v<sup>0</sup> (882.5 eV), v<sup>1</sup> (885.7 eV), v<sup>2</sup> (888.4 eV), v<sup>3</sup> (898.1 eV), u<sup>0</sup> (900.6 eV), u<sup>1</sup> (903.9 eV), u<sup>2</sup> (906.7 eV), u<sup>3</sup>



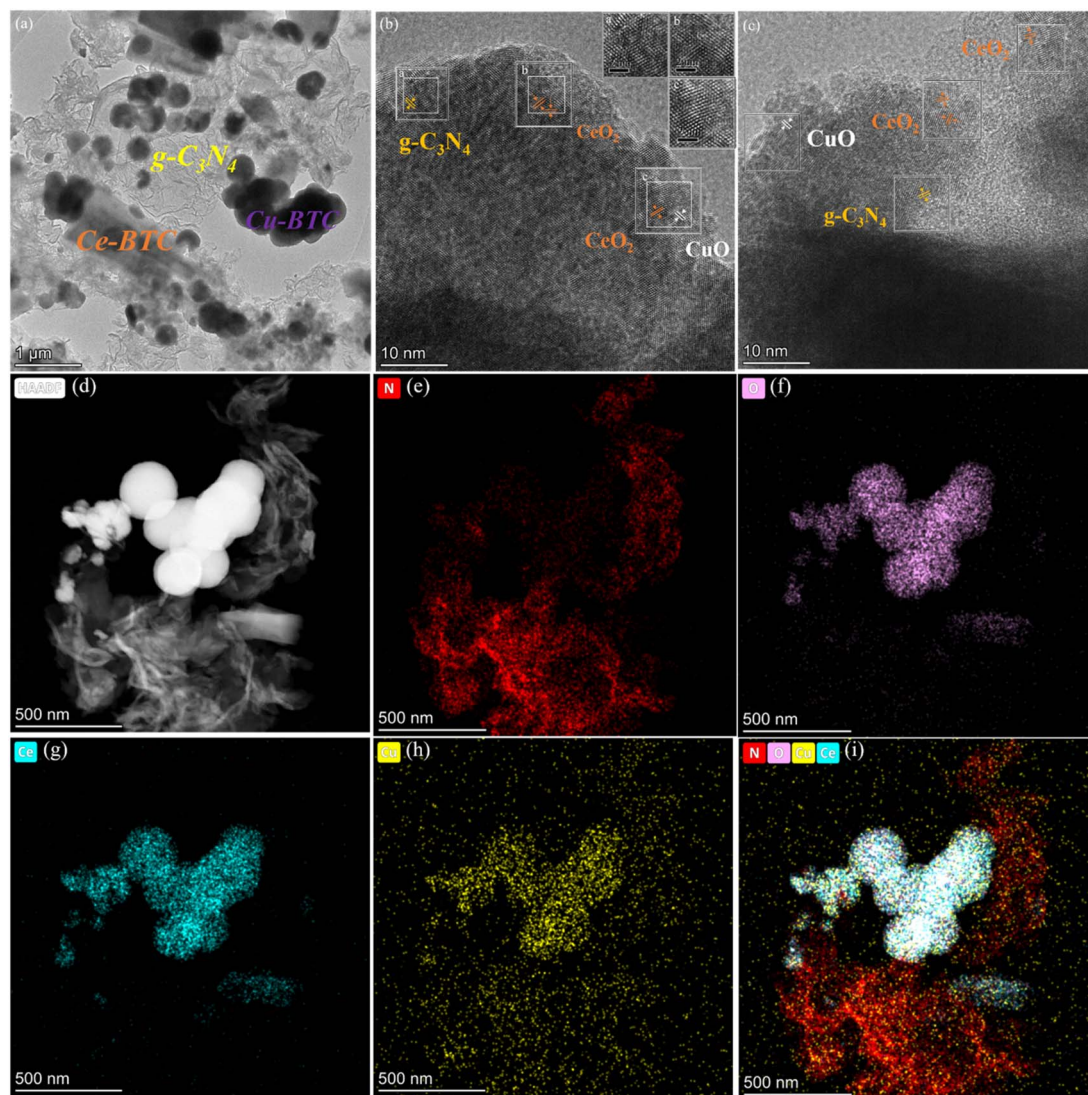


Fig. 3 TEM (HR-TEM) (a–c) of  $\text{CeCu}_{0.05}\text{-BTC/CN}$ , dark field image of  $\text{CeCu}_{0.05}\text{-BTC/CN}$  (d), EDS mapping image: N (e), O (f), Ce (g), Cu (h), general diagram of element distribution (i).

(916.7 eV). The four characteristic peaks  $u^0$ ,  $u'$ ,  $u''$ , and  $u'''$  are attributed to Ce  $3d_{3/2}$  spectral peaks, and  $v^0$ ,  $v'$ ,  $v''$ , and  $v'''$  are attributed to Ce  $3d_{5/2}$  spectral peaks.  $u^0$ ,  $u''$ , and  $u'''$  with  $v^0$ ,  $v'$ , and  $v'''$  can be attributed to  $\text{Ce}^{4+}$ , and the remaining two characteristic peaks correspond to  $\text{Ce}^{3+}$ .<sup>20,30,38</sup> Therefore,  $\text{Ce}^{3+}$  and  $\text{Ce}^{4+}$  coexisted on the surface of this sample, and it was hypothesized that the abundant photocatalytic properties could also be attributed to the renewable conversion between  $\text{Ce}^{3+}$  and  $\text{Ce}^{4+}$ . In Fig. 4(f), the Cu 2p spectrum is deconvoluted into two spin orbital peaks, Cu  $2p_{3/2}$  and Cu  $2p_{1/2}$ , corresponding to binding energies of 933.1, 934.3 eV, and 952.5, 956.3 eV (ref. 19 and 20) respectively. The Cu element in  $\text{CeCu}_x\text{-BTC}$  mainly exists in the form of  $\text{Cu}^+$  and  $\text{Cu}^{2+}$ , while the peaks of Ce 3d and Cu 2p are shifted in the direction of low binding energy, which indicates that there are interactions and electron transfer between  $\text{CeCu-BTC}$  and  $\text{g-C}_3\text{N}_4$ . It indicates that the photo-generated electrons on the  $\text{g-C}_3\text{N}_4$  surface are transferred to

$\text{CeCu-BTC}$ , and the electron–hole pair complex is effectively suppressed. This is in agreement with the XRD and TEM data.

Fig. 5(a and b) shows the UV-Vis absorption spectra of the samples in the range of 250–800 nm, which were calculated using the Tauc plot algorithm and the energy band eqn (1).

$$\alpha h\nu^{1/n} = A(h\nu - E_g) \quad (1)$$

where  $\alpha$  is the absorbance index,  $n$  is determined by the type of optical jump within the semiconductor,  $\nu$  is the frequency,  $h$  is Planck's constant ( $4.1356676969 \times 10^{-15}$ ), and  $E_g$  is the semiconductor bandgap width. Apparently,  $\text{CeCu}_{0.05}\text{-BTC/CN}$  shows a significant enhancement of visible light absorption compared to  $\text{g-C}_3\text{N}_4$ , while the presence of heterojunction results in red-shifted absorption bands, suggesting that the sample is capable of attracting more visible-light photons in the UV and visible ranges to obtain the active  $e^-/h^+$ , which subsequently



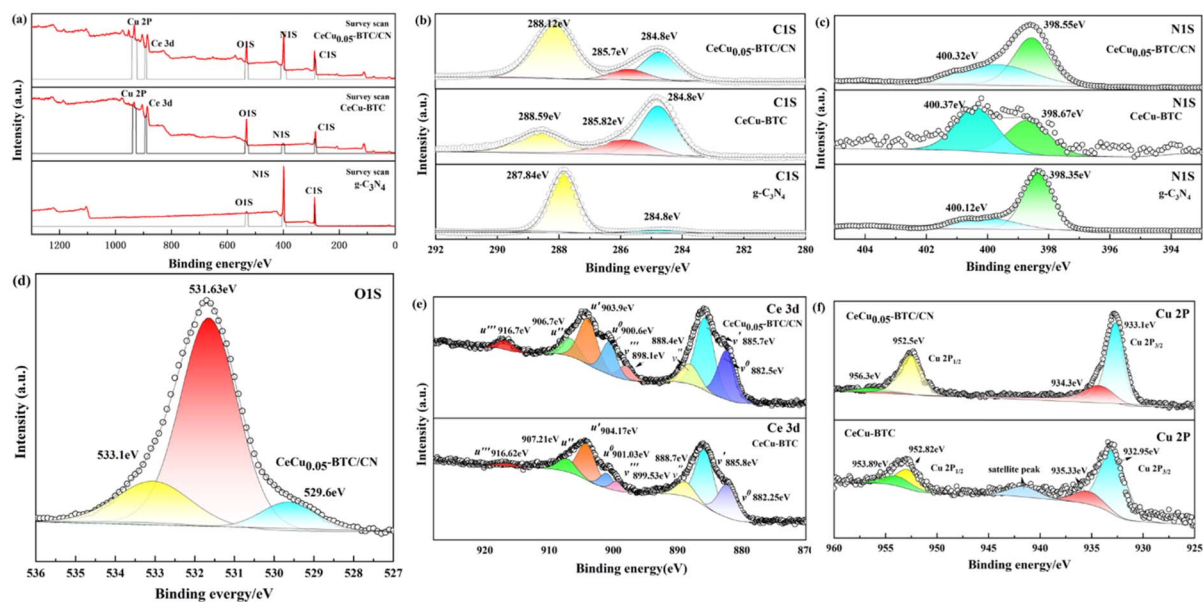


Fig. 4 XPS spectra of the  $g\text{-C}_3\text{N}_4$ , CeCu-BTC and CeCu<sub>0.05</sub>-BTC/CN: total elemental spectra (a), C 1s (b), N 1s (c) O 1s (d), Ce 3d (e), Cu 2p (f).

improves the photocatalytic performance. As shown in Fig. 5(a and b), the visible absorption band edges of  $g\text{-C}_3\text{N}_4$ , CeCu-BTC and, CeCu<sub>0.05</sub>-BTC/CN are about 455 nm, 534 nm, and 509 nm, which is in agreement with earlier reports.<sup>39</sup> The optical band gap energy ( $E_g$ ) valuations of 2.72, 2.32, and 2.44 eV for  $g\text{-C}_3\text{N}_4$ , CeCu-BTC, and CeCu<sub>0.05</sub>-BTC/CN, respectively, were deduced from eqn (1). The results show that the introduction of Cu narrows the optical band gap value of CeCu<sub>0.05</sub>-BTC/CN composites, and the lower  $E_g$  values indicate that the samples

may produce more photogenerated carriers and possess higher visible light trapping ability.<sup>34</sup>

Fig. 5(c) shows the separation of photogenerated carriers generated at the contact interface of different samples at the 315 nm excitation wavelength. The stronger the PL peak, the higher the  $e^-/h^+$  recombination rate, and the lower the corresponding separation efficiency, which is more conducive to the photocatalytic reaction. This is also one of the reasons why the original  $g\text{-C}_3\text{N}_4$  has low capture ability for visible light and is

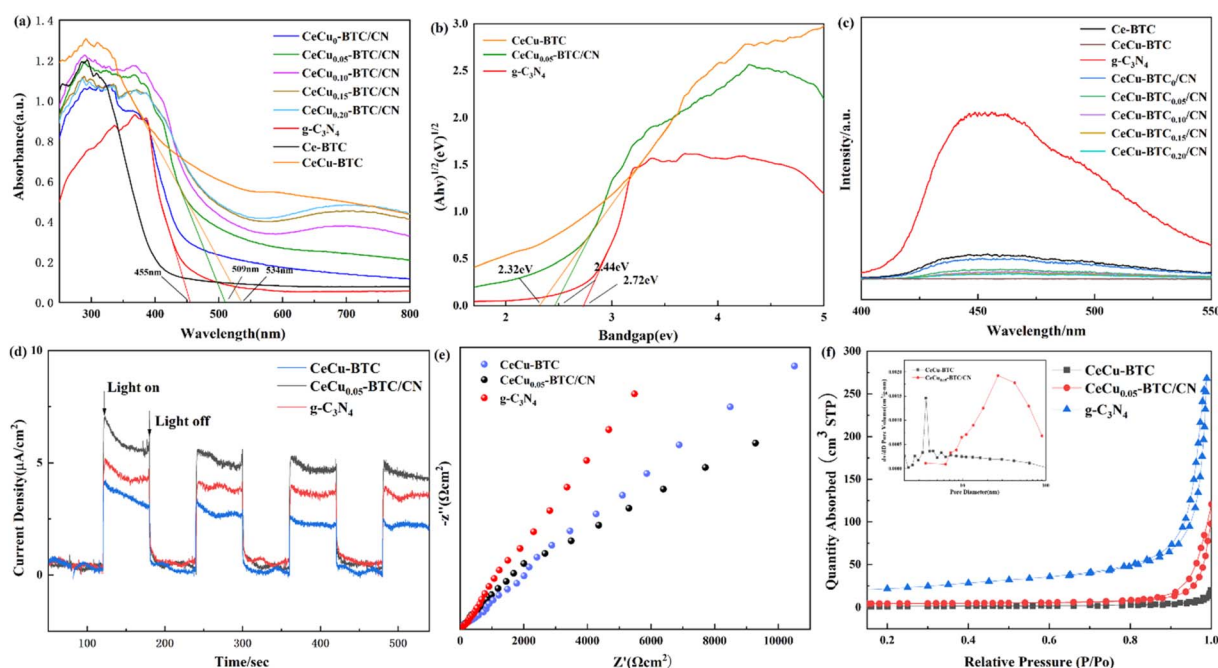


Fig. 5 UV-Vis absorption spectrum (a) and Tauc curve (b), fluorescence spectrum (c), transient photocurrent response spectrum (d), Nyquist plots (e) of the sample, N<sub>2</sub> adsorption and desorption isothermal curves and pore size distribution curves (f).



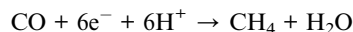
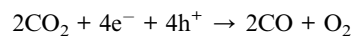
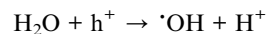
easy to compound. CeCu-BTC has no fluorescence in this band, which also shows that the introduction of the metal organic framework helps to reduce the carrier recombination. At the same time, it is speculated that the phenomenon of improving the photoelectron separation efficiency and reducing the recombination rate should also be related to the synergistic effect between *g*-C<sub>3</sub>N<sub>4</sub> and Cu doping in CeCu-BTC and Ce.<sup>30</sup> In summary, the construction of this heterojunction not only enhances the photoelectron transport capacity but also effectively suppresses the recombination rate and, at the same time, has a contributing effect on the improvement of photocatalytic performance.<sup>40,41</sup>

Fig. 5(d) shows the transient current response of CeCu-BTC, CeCu<sub>0.05</sub>-BTC/CN, and *g*-C<sub>3</sub>N<sub>4</sub>. The intensity of the photocurrent reflects the separation efficiency of photogenerated electrons and holes.<sup>42</sup> The transient photocurrent intensity of CeCu<sub>0.05</sub>-BTC/CN was significantly higher than that of CeCu-BTC, and *g*-C<sub>3</sub>N<sub>4</sub>, showing a periodic and stable reciprocating law, indicating that the carrier separation efficiency in the synthesized heterostructure is higher and the photocatalytic reaction is more efficient. The radius of the arc area in the EIS Nyquist plot shown in Fig. 5(e) indicates the resistance to the transfer of charge on the surface of the semiconductor electrode.<sup>43</sup> With the decrease in semicircular radius, the generation efficiency of electron hole pairs increases. It is observed that the arc radius of CeCu<sub>0.05</sub>-BTC/CN Nyquist is smaller than that of other samples, which indicates that the former has a smaller transmission impedance and lower resistance to carrier separation, which is more conducive to charge transfer. Fig. 5(f) shows the N<sub>2</sub> adsorption-desorption experiments of the samples at 77 K. The *g*-C<sub>3</sub>N<sub>4</sub>, CeCu-BTC, and CeCu<sub>0.05</sub>-BTC/CN were found to be H<sub>3</sub>-type hysteresis loops with type IV adsorption isotherms. When the sample is at  $P/P_0 > 0.8$ , the hysteresis loop may be due to the adsorption of N<sub>2</sub> in the slit pores of MOFs, and the H<sub>3</sub> hysteresis loop is mainly found in the aggregates of layered structure, so it is speculated that it may be the intermolecular graphite-like structure of *g*-C<sub>3</sub>N<sub>4</sub>. This type of hysteresis loop also shows that most of the materials have a mesoporous structure and the pore distribution is uneven. The specific surface area, pore volume, and average pore size of different samples can be obtained in Table 1. It is found that the specific surface area and average pore size of CeCu<sub>0.05</sub>-BTC/CN are significantly increased compared with CeCu-BTC. It is inferred that the doping of Cu and the recombination of *g*-C<sub>3</sub>N<sub>4</sub> may lead to more defects and affect the growth of crystals, which is also consistent with the results of electron microscopy. At the same time, we infer that a more porous structure and relatively large specific surface area also provide more active sites and the possibility of reagent

adsorption for the reaction, thereby enhancing the photocatalytic ability of the sample.

### 3.2 Photocatalytic carbon dioxide reduction

Fig. 6 shows the photocatalytic reduction test of carbon dioxide under simulated sunlight. *g*-C<sub>3</sub>N<sub>4</sub> and CeCu<sub>*x*</sub>-BTC/CN products are both CO and CH<sub>4</sub>, where CO is designated as the main product, accompanied by a small amount of methane production. The photocatalytic carbon dioxide reduction experiments were carried out under gas-solid phase conditions with 1 mL of pure water as the proton-providing source and no sacrificial agent, and the reaction mechanism of the different intermediates produced by participating in the carbon dioxide reduction can be represented by the following reaction equations.<sup>44</sup>



For the original *g*-C<sub>3</sub>N<sub>4</sub>, the CO yield is only 8.522 μmol g<sup>-1</sup> and the CH<sub>4</sub> yield is 0.232 μmol g<sup>-1</sup> under this condition, which may be due to the higher charge-carrier recombination rate and rapid recombination. As can be seen in Fig. 6(a), the CO and CH<sub>4</sub> reduction capacities of CeCu<sub>*x*</sub>-BTC/CN, which successfully constitutes the heterostructure, are both higher than those of *g*-C<sub>3</sub>N<sub>4</sub>. Among them, CeCu<sub>0.05</sub>-BTC/CN CO yield was 64.44 μmol g<sup>-1</sup>, which was 7.56 times of *g*-C<sub>3</sub>N<sub>4</sub>, and the CO yields of other samples were 15.07, 36.39, 34.43, and 31.07 μmol g<sup>-1</sup> (*x* = 0, 0.1, 0.15, 0.2), which were 1.77, 4.27, 4.04, and 3.65 times of *g*-C<sub>3</sub>N<sub>4</sub>, respectively. The yield of the heterostructure CeCu<sub>0.05</sub>-BTC/CN shown in Fig. 6(b) was 0.5575 μmol g<sup>-1</sup>, which was 2.42 times higher than that of *g*-C<sub>3</sub>N<sub>4</sub>, and the CH<sub>4</sub> yields of the other samples were 1.149, 0.557, 0.552, and 0.542 μmol g<sup>-1</sup> (*x* = 0, 0.1, 0.15, 0.2), which were 5 times higher than that of *g*-C<sub>3</sub>N<sub>4</sub>, respectively, 2.42, 2.4, and 2.36 times that of *g*-C<sub>3</sub>N<sub>4</sub>, respectively. It also shows that the content of Cu doping is the key to realizing the high photocatalytic activity of the composite heterostructure material. The suitable Cu content makes it well dispersed on Ce-BTC and *g*-C<sub>3</sub>N<sub>4</sub> surfaces, which facilitates carrier transfer and separation.<sup>45,46</sup> In addition, when the content of Cu was higher than 5.0 wt%, a further increase in the proportion of Cu led to a decrease in the photocatalytic CO<sub>2</sub> reduction performance.

Fig. 6(c) shows that in the absence of a photocatalyst, light source, or N<sub>2</sub> atmosphere instead of CO<sub>2</sub>, the reaction did not produce CO and CH<sub>4</sub>, indicating that CO<sub>2</sub> was the only carbon source involved in the reduction reaction, with no other factors influencing it. Fig. 6(d) shows the cycling test of sample CeCu<sub>0.05</sub>-BTC/CN, in which the CO and CH<sub>4</sub> yields were found to remain good after centrifugal recovery and activation several times. In addition, the recovered XRD spectra (Fig. 6(e)) and FT-IR spectra (Fig. 6(f)) also proved that the main crystal structure and chemical functional groups had no obvious change after the stability test, indicating that it had good photocatalytic stability.

Table 1 Comparison of S<sub>BET</sub>, pore volume and average pore size

Sample	S <sub>BET</sub> (m <sup>2</sup> g <sup>-1</sup> )	Pore volume (cm <sup>3</sup> g <sup>-1</sup> )	Average pore size (nm)
<i>g</i> -C <sub>3</sub> N <sub>4</sub>	73.87	0.41	22.40
CeCu-BTC	4.4905	0.03	26.04
CeCu <sub>0.05</sub> -BTC/CN	16.510	0.13	35.53





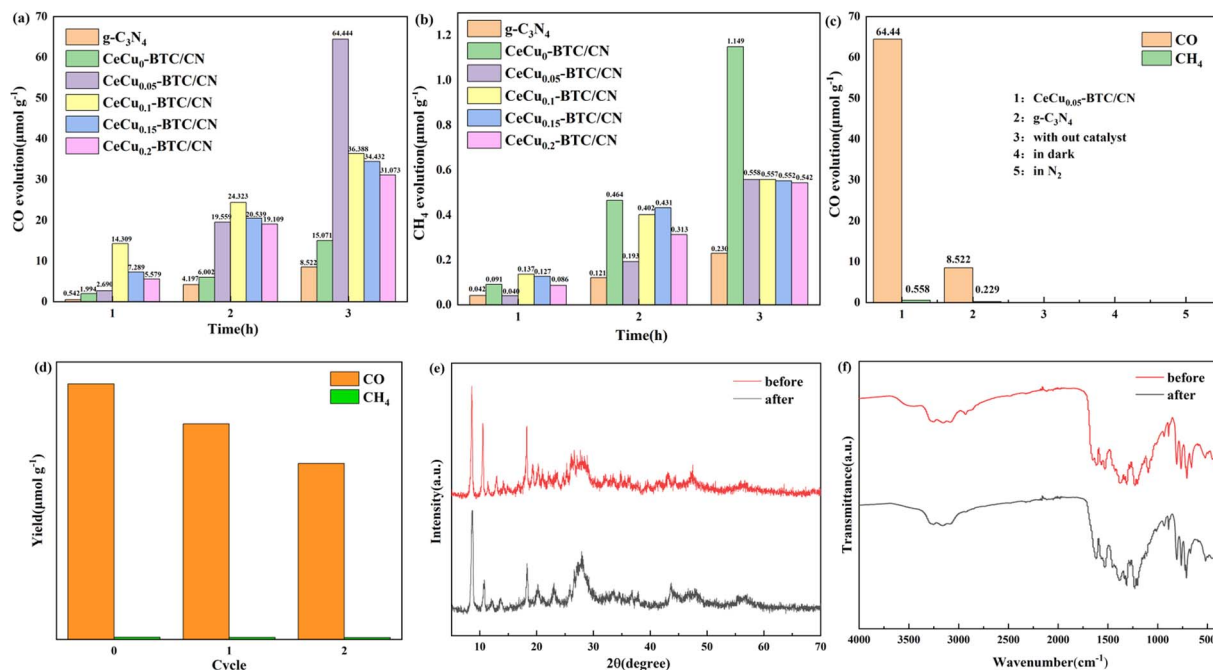


Fig. 6 Photocatalytic CO<sub>2</sub> reduction CO yield plot (a), CH<sub>4</sub> yield plot (b), control experiment (c), CeCu<sub>0.05</sub>-BTC/CN cycle test (d), XRD and FT-IR changes of the samples before and after illumination (e and f).

### 3.3 Research on photocatalytic mechanism

To further explain the sample mechanism, electron paramagnetic resonance (EPR) was used to analyze the direction of electron transfer in the system. In Fig. 7(a and b), DMPO-<sup>•</sup>OH was not seen in any of the dark reaction conditions, and CeCu<sub>0.05</sub>-BTC and CeCu<sub>0.05</sub>-BTC/CN were observed in the light reaction, showing a standard fourfold peak and four peaks equidistant with peak heights of 1 : 2 : 2 : 1. The larger DMPO-<sup>•</sup>OH may be attributed to its VB potential to be corrected, that is, the VB potential of CeCu-BTC and CeCu<sub>0.05</sub>-BTC/CN should be greater than 1.99 eV (ref. 47) (OH<sup>-</sup>/-OH 1.99 eV). On the contrary, DMPO-<sup>•</sup>OH was not found in g-C<sub>3</sub>N<sub>4</sub>, presumably due to the weak oxidation potential of the photogenerated holes, which also suggests that the photogenerated holes remain in the VB of CeCu-BTC and are not transferred to the VB of g-C<sub>3</sub>N<sub>4</sub>. According to the linear extrapolation results, the  $E_{\text{VB-XPS}}$  of g-C<sub>3</sub>N<sub>4</sub> and CeCu-BTC in Fig. 7(c and d) are 1.65 and 2.03 eV, respectively.

In order to obtain the actual VB potential of the samples more accurately and efficiently, the samples were estimated according to eqn (2) and (3).  $\phi$  is the work function of the XPS instrument, with a value of 4.2 eV, and the NHE at pH = 7 is 4.44 eV. As a result, the values of  $E_{\text{VB-NHE}}$  for g-C<sub>3</sub>N<sub>4</sub> and CeCu-BTC are obtained as 1.41 and 1.79 eV. Eqn (3) yields ECB's of -1.31 and -0.53 eV for g-C<sub>3</sub>N<sub>4</sub> and CeCu-BTC, respectively.

$$E_{\text{VB-NHE}} = \phi + E_{\text{VB-XPS}} - 4.44 \quad (2)$$

$$E_{\text{CB}} = E_{\text{VB}} - E_{\text{g}} \quad (3)$$

Finally, to verify the accuracy of the above potential calculations, we also utilised the semiconductor electronegativity

theory (shown in eqn (4)) to calculate the VB potential.  $X$  is the geometric mean of the absolute electronegativity of the atoms within the semiconductor, and  $E_{\text{c}}$  is a constant with respect to

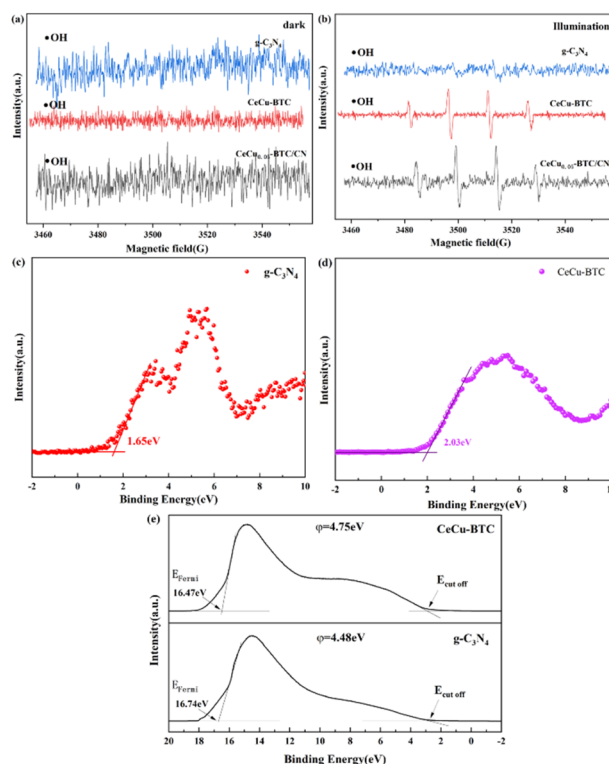


Fig. 7 EPR spectra of DMPO-<sup>•</sup>OH spin capture (a and b), VB-XPS (c and d), UV electron spectroscopy UPS (e).



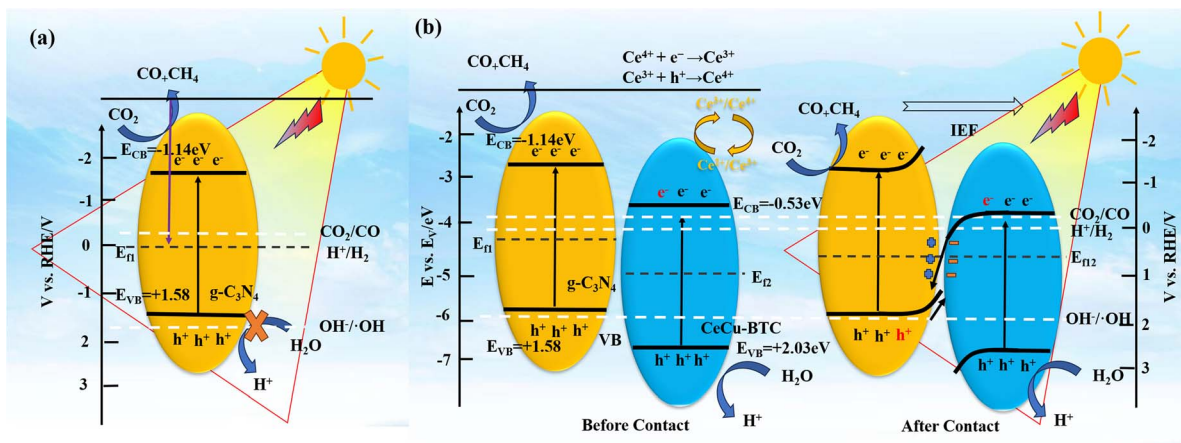


Fig. 8 Photocatalytic mechanism diagram of g-C<sub>3</sub>N<sub>4</sub> (a), the work functions of CN and CeCu-BTC before/after contact and the formation of internal electric field (b).

the standard H electrode, generally taken to be 4.5 eV, with  $X$  for g-C<sub>3</sub>N<sub>4</sub> being 4.73 eV,<sup>48</sup> as an example. The  $E_{VB}$  of g-C<sub>3</sub>N<sub>4</sub> is 1.58 eV and the  $E_{CB}$  is -1.14. Meanwhile, the results obtained by the two calculation methods were found to be approximate, and the semiconductor electronegativity theory was further used for the explanation of the photocatalytic mechanism.

$$E_{VB} = X - E_e + 0.5E_g \quad (4)$$

$$\phi = 21.22 - \Delta E \quad (5)$$

In addition, Fig. 7(e) shows the UPS spectrum of the sample. The work function ( $\phi$ ) of the sample is obtained through formula (5), and  $\Delta E$  is the distance from the upper starting point to the cut-off edge of the secondary electron after the bias voltage is applied. The  $\phi$  of g-C<sub>3</sub>N<sub>4</sub> and CeCu-BTC are 16.74 and 16.47 eV, respectively. The results indicate that CeCu-BTC has a larger  $\phi$  value and a relatively small Fermi energy level, while the opposite is true for g-C<sub>3</sub>N<sub>4</sub>. Therefore, when g-C<sub>3</sub>N<sub>4</sub> and CeCu-BTC are in close contact to form a CeCu-BTC/CN heterojunction, the work function difference between the two samples causes the free electrons of CN to spontaneously transfer to CeCu-BTC. With the Fermi level of g-C<sub>3</sub>N<sub>4</sub> gradually decreasing, the Fermi level of CeCu-BTC gradually increases until the interface  $E_{f12}$  reaches equilibrium. Similarly, a space charge region, *i.e.* an built-in electric field, will be formed at the interface between the two. The existence of the built-in electric field will bend the Fermi level and pull up the energy band of g-C<sub>3</sub>N<sub>4</sub> and lower the energy band of CeCu-BTC. Therefore, the photogenerated electrons of g-C<sub>3</sub>N<sub>4</sub> have to overcome the potential energy barrier to fall back into CeCu-BTC driven by the built-in electric field and interface chemical bonds (Fig. 8(a and b)). Finally, by analyzing the photocatalytic CO<sub>2</sub> reduction mechanism diagram of g-C<sub>3</sub>N<sub>4</sub> and the electron paramagnetic resonance (EPR) capture results of hydroxyl radicals (<sup>•</sup>OH) shown in Fig. 7(a), and combined with the position where the energy bands of the photocatalysts are located, it is preferred that the photogenerated electrons on the CB of CeCu-BTC will recombine with the photogenerated holes of the VB of g-C<sub>3</sub>N<sub>4</sub>,

which also verifies the successful construction of the S-type heterojunction.

The above conclusions indicate that CeCu-BTC and CN change from a steady state to an excited state upon irradiation with visible light larger than the bandgap of both, producing photogenerated electrons and holes. Due to the interaction of IEF and Coulomb at the interface, the photogenerated electrons on the CB of CeCu-BTC will combine with the photogenerated holes on the VB of g-C<sub>3</sub>N<sub>4</sub>, and then the photogenerated electrons accumulated on the CB of g-C<sub>3</sub>N<sub>4</sub> will react with the CO<sub>2</sub> adsorbed on the surface of the catalyst to form CO and CH<sub>4</sub>. Meanwhile, the CB potential of CeCu-BTC and g-C<sub>3</sub>N<sub>4</sub> is more negative than that of Cr(VI)/Cr(III) (+0.51 eV vs. NHE), which indicates that the reduction of adsorbed Cr(VI) to Cr(III) is thermodynamically beneficial to the CB of the photocatalyst. The coexistence of Ce<sup>3+</sup> and Ce<sup>4+</sup> can also aid in the transfer of electrons from the catalyst surface, thus prolonging the reaction time between electrons and carbon dioxide.

## 4 Conclusions

In summary, The S-type heterojunction photocatalysts (CeCu<sub>x</sub>-BTC/CN) with cerium-copper bimetallic organic framework (CeCu<sub>x</sub>-BTC) and graphite-phase carbon nitride (g-C<sub>3</sub>N<sub>4</sub>) were constructed by a simple solvothermal method for photocatalytic CO<sub>2</sub> reduction to CO and CH<sub>4</sub>. The results show that the built-in electric field constructed by Fermi energy level flattening transfers the electrons in an S-type manner, which not only preserves the strong reducing properties of the electrons in the material but also provides the maximum redox capacity and enables the composite samples to obtain higher visible-light trapping capacity and improve the separation efficiency of the carriers while refining the crystal particles. CeCu<sub>0.05</sub>-BTC/CN possessed the optimal photocatalytic performance with the addition of only 1 mL of H<sub>2</sub>O as the proton provision source. The CO and CH<sub>4</sub> yields were 64.44 μmol g<sup>-1</sup> and 0.5575 μmol g<sup>-1</sup>, which were 7.56 and 2.42 times higher than that of g-C<sub>3</sub>N<sub>4</sub>, respectively, and the catalytic performance was basically stable after the cyclic test,



and this heterojunction also paved the way for a small amount of low-cost composite modification of g-C<sub>3</sub>N<sub>4</sub>.

## Data availability

Crystallographic data for g-C<sub>3</sub>N<sub>4</sub>, Ce-BTC, have been deposited at the JCPDS/CCDC under JCPDS 87-1526, CCDC ID: 773084 and can be obtained from <http://dx.doi.org/10.1002/smll.202003597>, <https://doi.org/10.1016/j.matchemphys.2018.04.031>, <http://dx.doi.org/10.1016/j.apsusc.2017.06.235>. The authors confirm that the data supporting the findings of this study are available within the article and/or as its supplementary materials. The data that support the findings of this study are available from the first or corresponding author (Hui Zhang or Rui fen Wang) upon reasonable request.

## Author contributions

H. Z. and Y. H. S. completed the sample synthesis and testing. R. F. W., S. L. A., and R. H. G. conceived the idea and accompanied the test and guidance at all stages; H. Z. and Y. H. S. wrote the manuscript; Y. W. M. was responsible for the final review; and all the authors contributed to the manuscript.

## Conflicts of interest

The authors declare no competing interests.

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