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# Growth of narrow-bandgap Cl-doped carbon nitride nanofibers on carbon nitride nanosheets for high-efficiency photocatalytic H<sub>2</sub>O<sub>2</sub> generation†

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Heterojunction construction has been proved to be an effective way to enhance photocatalysis performance. In this work, Cl-doped carbon nitride nanofibers (Cl–CNF) with broadband light harvesting capacity were *in situ* grown on carbon nitride nanosheets (CNS) by a facile hydrothermal method to construct a type II heterojunction. Benefiting from the joint effect of the improved charge carriers separation efficiency and a broadened visible light absorption range, the optimal heterostructure of Cl–CNF/CNS exhibits a H<sub>2</sub>O<sub>2</sub> evolution rate of 247.5 μmol g<sup>−1</sup> h<sup>−1</sup> under visible light irradiation, which is 3.4 and 3.1 times as much as those of Cl–CNF (72.2 μmol g<sup>−1</sup> h<sup>−1</sup>) and CNS (80.2 μmol g<sup>−1</sup> h<sup>−1</sup>), respectively. Particularly, the heterojunction nanostructure displays an apparent quantum efficiency of 23.67% at 420 nm. Photoluminescence spectra and photocurrent measurements both verified the enhanced charge carriers separation ability. Our work provides a green and environmentally friendly strategy for H<sub>2</sub>O<sub>2</sub> production by elaborate nanostructure design.

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## 1. Introduction

H<sub>2</sub>O<sub>2</sub>, as an important chemical reagent, has been widely used in medical disinfection, industrial production, daily sterilization and decontamination.<sup>1,2</sup> In addition, due to its high energy density (2.1 MJ kg<sup>−1</sup> for 60% aqueous H<sub>2</sub>O<sub>2</sub>), harmless products emission and convenient storage and transportation conditions, H<sub>2</sub>O<sub>2</sub> is regarded as a promising energy carrier.<sup>3,4</sup> H<sub>2</sub>O<sub>2</sub> has also been added as a favorable electron acceptor to boost charge carrier separation to enhance the photocatalytic performance of photocatalysts.<sup>5–7</sup> At present, H<sub>2</sub>O<sub>2</sub> is predominantly prepared by industrial oxidation of anthraquinone, which currently accounts for more than 95% of the total production of H<sub>2</sub>O<sub>2</sub>. However, the anthraquinone oxidation method is environmentally hazardous, and consumes extensive energy. Therefore, it is urgent to develop an environmentally friendly and low energy consumption method for H<sub>2</sub>O<sub>2</sub> synthesis.<sup>8</sup> Photocatalysis, which is driven by solar energy, has been considered to be one of the effective ways to solve the energy and environmental crisis.<sup>9</sup> Among various kinds of photocatalyst, g-C<sub>3</sub>N<sub>4</sub>, because of its unique features, such as

a suitable bandgap for light harvesting, facile preparation process, favorable heat and chemical stability and metal-free characteristics,<sup>10–12</sup> has attracted extensive interest in photocatalytic production of H<sub>2</sub> (ref. 13–16) and H<sub>2</sub>O<sub>2</sub>,<sup>17–19</sup> nitrogen fixation,<sup>20–22</sup> organic pollutants decomposition,<sup>23–25</sup> CO<sub>2</sub> reduction<sup>26–28</sup> and organic reactions.<sup>29,30</sup> However, several shortcomings, such as a narrow light absorption range, serious recombination of photogenerated charge carriers and low surface charge transfer efficiency still exist in g-C<sub>3</sub>N<sub>4</sub>, which limit its photocatalytic activity.<sup>31–34</sup>

Based on the above problems, a lot of methods have been employed to improve the photocatalytic performance, including defect construction, heteroatom doping, morphology optimization, *etc.*<sup>35–40</sup> For example, the photocatalytic H<sub>2</sub>O<sub>2</sub> generation activity was significantly enhanced by constructing carbon vacancies<sup>18</sup> or nitrogen vacancies<sup>19</sup> on the surface of g-C<sub>3</sub>N<sub>4</sub> due to the improved light harvesting capacity. Heteroatom doping, for example, Fe and non-metal S, P doping, has been proved to be an effective approach to boost the photocatalytic hydrogen generation activity and pollutants degradation.<sup>5–7,33,41</sup> Similarly, the halogen (F, Cl and Br)-doped photocatalysts exhibited some advantages of improving optical absorption, promoting charge migration and tuning band positions for photocatalytic applications.<sup>42</sup> However, the photoexcited charge carriers recombination problem is always difficult to be alleviated in single-component photocatalyst. Accordingly, heterojunction construction is developed and has been proved to be an effective strategy to inhibit the recombination of charge carriers during photocatalysis process.

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Among different types of heterojunction, the type II heterojunction can take both advantages of the two semiconductors to broaden the light absorption range. Moreover, the staggered band structure in type II heterojunction is supposed to be favorable for charge carriers separation, thus is ideal for enhancing photocatalytic performance. Polymeric graphitic carbon nitride has been reported to combine with an all-inorganic cesium lead halide perovskite ( $\text{CsPbBrCl}_2$ ) to prepare a type II heterojunction.<sup>43</sup> Due to the delay of the photogenerated electron–hole pairs recombination, the synthesized hybrid presented remarkably improved performance in toxic cationic and anionic dyes degradation as compared to individual building units. Hao *et al.* reported on the preparation of a ZnS/g- $\text{C}_3\text{N}_4$  heterostructure with abundant zinc vacancy defects modified on the surface of ZnS.<sup>44</sup> The heterostructure possessed expanded light absorption range according to its absorption spectra. Moreover, the staggered band structure and large number of zinc vacancies could boost the separation of photogenerated charge carriers. Due to the joint effect of broadened light absorption capacity and boosted charge carriers separation ability, the photocatalytic hydrogen production activity of ZnS/g- $\text{C}_3\text{N}_4$  is more than 30 times than that of pure g- $\text{C}_3\text{N}_4$ . The calculation results by Wei *et al.* also show that the formation of g- $\text{C}_3\text{N}_4/\text{r-TiO}_2$  heterojunction can expand its absorption range toward visible light.<sup>45</sup> In addition, the staggered alignments of band structure and built-in electric field could efficiently promote the separation of photoinduced electron–hole pairs, therefore leading to an improved  $\text{H}_2\text{S}$  splitting performance. Particularly, the 1D/2D heterostructure could realize the unique dimensionality-dependent advantages of rapid carrier transfer along the 1D nanostructures and superior electron mobility of 2D materials and simultaneously alleviate the drawbacks of the single units, such as the low surface area of 1D nanofibers and the tendency of 2D nanosheets toward restacking.<sup>46</sup> Combining the advantages of doping and heterojunction, the construction of Cl-doped carbon nitride nanofibers on carbon nitride nanosheets is expected to show superior photocatalytic activity.

Herein, we fabricated a type II heterojunction nanostructure by growing narrow-bandgap 1D Cl-CNF on 2D CNS *via* a simple hydrothermal synthesis procedure. A variety of characterization methods were applied to verify the formation of the heterostructure. The light absorption ability of the heterostructure was improved because of the introduction of Cl-CNF, which exhibits broadband light harvesting capacity. In addition, the photoluminescence spectra and photocurrent measurements jointly demonstrated that the charge carriers generation and separation ability was significantly enhanced in the heterojunction. Therefore, a  $\text{H}_2\text{O}_2$  generation rate of  $247.5 \mu\text{mol g}^{-1} \text{h}^{-1}$  was obtained in the optimal sample of Cl-CNF/CNS under visible light irradiation ( $\lambda > 420 \text{ nm}$ ), which is 3.4 and 3.1 times as much as those of single-component Cl-CNF ( $72.2 \mu\text{mol g}^{-1} \text{h}^{-1}$ ) and CNS ( $80.2 \mu\text{mol g}^{-1} \text{h}^{-1}$ ), respectively. Especially, the optimal type II heterojunction nanostructure achieves an apparent quantum efficiency of 23.67% at 420 nm benefiting by the synergistic effect the Cl-CNF and CNS.

## 2. Experimental

### 2.1. Preparation of the photocatalysts

The CNS powder was prepared by polymerization of dicyandiamide in air, accompanied by using ammonium chloride as a gas template.<sup>31</sup> Typically, dicyandiamide (2 g) and ammonium chloride (10 g) were first ground in a mortar to mix homogeneously and then the mixture was transferred to a covered crucible. The thermal condensation was accomplished by heating the crucible in a muffle furnace with a heating rate of  $1 \text{ }^\circ\text{C min}^{-1}$  and a maintaining temperature of  $550 \text{ }^\circ\text{C}$  for 3 h. To prepare a uniformly-distributed heterojunction nanostructure, the CNS sample should be acidified in advance. The acidification of CNS was proceeded by adding the CNS powder (0.5 g) into a hydrochloric acid aqueous solution ( $30 \text{ mL}$ ,  $3 \text{ mol L}^{-1}$ ). The mixture was stirred for 12 h to make sure the acidification was finished completely. Then the dispersion was centrifuged and washed with water for several times until the pH of the supernatant is close to neutral. The acidified powder was finally collected after drying the precipitate in an oven. For comparison, melamine-derived carbon nitride and dicyandiamide-derived carbon nitride were prepared under the same preparation condition as that of CNS by polymerization of sole dicyandiamide and melamine, respectively.

The narrow-bandgap Cl-CNF semiconductor with broadband light harvesting capacity was prepared by a hydrothermal method according to a previous report.<sup>35</sup> Typically, melamine (0.252 g) and cyanuric chloride (0.738 g) were added into acetonitrile solution (30 mL) in sequence, and then the mixture was stirred for 24 h at room temperature and hydrothermally reacted for another 24 h at  $180 \text{ }^\circ\text{C}$ . The precipitate was collected and respectively washed with water and ethanol for several times to remove the impurities on the catalyst surface. Finally, the precipitate was dried in an oven at  $60 \text{ }^\circ\text{C}$ .

The Cl-CNF/CNS nanohybrids were prepared under the same hydrothermal condition with that of Cl-CNF, except for the introduction of acidified CNS. After the acetonitrile solution containing melamine and cyanuric chloride was stirred for 12 h, acidified CNS was added and then the mixture was stirred for another 12 h to ensure that the Cl-CNF precursors were uniformly adsorbed on the acidified CNS nanosheets surface. The Cl-CNF/CNS samples were then obtained after the suspension mixture hydrothermally reacted at  $180 \text{ }^\circ\text{C}$  for 24 h. The grafting density of Cl-CNF on the CNS surface can be adjusted by changing the addition amount of acidified CNS. In the abbreviation of Cl-CNF/CNS- $x$ ,  $x$  is the addition mass of acidified CNS. Unless otherwise specially mentioned, the Cl-CNF/CNS sample represents Cl-CNF/CNS-0.4. As control samples, Cl-CNF/melamine-derived carbon nitride and Cl-CNF/dicyandiamide-derived carbon nitride with carbon nitride addition amount of 0.4 g were prepared similarly with that of Cl-CNF/CNS for  $\text{H}_2\text{O}_2$  generation.

### 2.2. Characterizations

The surface charges of the samples and precursors were measured by their zeta potentials on Zetasizer Nano (Malvern



Instruments) in their reaction solutions. The morphologies of the samples were investigated on scanning electron microscope (SEM, Quanta 250 FEG FEI) equipped with an energy-dispersive X-ray (EDX) detector and transmission electron microscope (TEM) (Tecnai G2 20 TEM) with an accelerating voltage of 200 kV. The light harvesting capacities of the samples were characterized by their absorption spectra on a Hitachi ultraviolet/visible/near-infrared (UV-vis-NIR) spectrophotometer equipped with an integrating sphere with BaSO<sub>4</sub> as a base. The crystalline characteristics of the samples were investigated by collecting their X-ray diffraction (XRD) patterns on a Bruker D8 Advance diffractometer with tube voltage and current of 40 kV and 40 mA, respectively. The Fourier transform infrared (FTIR) spectra of the samples were collected on a Thermo Scientific Nicolet iS5 infrared spectrophotometer using potassium bromide pellet technique. X-ray photoelectron spectroscopy (XPS) measurements were recorded on an Escalab 250 Xi (Thermo Fisher Scientific) system with Al K $\alpha$  ( $h\nu = 1486.6$  eV) as the X-ray source. The photoluminescence (PL) measurements (Hitachi F-4600 spectrophotometer) were carried out to investigate the radiative recombination of the photogenerated charge carriers. The transient-state photoluminescence (TRPL) spectra were recorded on Edinburgh Instruments FLS980 spectrophotometer with excitation and emission wavelength of 350 and 455 nm, respectively. N<sub>2</sub> adsorption-desorption isotherms, which were used to calculate the Brunauer-Emmett-Teller (BET) specific surface areas, were collected on a Micromeritics ASAP 2020 analyzer.

### 2.3. Photocatalytic performance measurements

The photocatalytic performances of the photocatalysts were evaluated through the photogeneration of H<sub>2</sub>O<sub>2</sub> in a photo-reactor by employing a 300 W Xe lamp (CEL-HXF300, CEALIGHT, Beijing, China) equipped with a cut-off filter of 420 nm as the light source. Typically, the photocatalyst (50 mg) was first dispersed into an ethanol aqueous solution (50 mL, 10 vol% ethanol in deionized water). The mixture was subjected to ultrasonic treatment for 10 min to obtain a uniformly mixed suspension, and then bubbled with O<sub>2</sub> for 20 min to create an oxygen-rich environment under stirring before light was turned on. During photocatalytic reaction process, every 1.5 mL of the reaction solution was taken out after 0, 0.5, 1.0, 1.5 and 2.0 h of light irradiation, respectively. The concentration of H<sub>2</sub>O<sub>2</sub> generated in the reaction solution was determined by a Ce(SO<sub>4</sub>)<sub>2</sub> titration method<sup>1</sup> based on that yellow Ce<sup>4+</sup> solution can be reduced by H<sub>2</sub>O<sub>2</sub> to colorless Ce<sup>3+</sup> solution:  $2\text{Ce}^{4+} + \text{H}_2\text{O}_2 \rightarrow 2\text{Ce}^{3+} + 2\text{H}^+ + \text{O}_2$ . The generation of H<sub>2</sub>O<sub>2</sub> can therefore be quantitatively calculated by the decrease amount of Ce<sup>4+</sup>. Moreover, the evolution amount of H<sub>2</sub>O<sub>2</sub> under monochromatic light was measured with a bandpass filter under the same reaction condition. The power intensity of monochromatic light was measured with an optical power meter (CEL-NP2000-2, CEALIGHT, Beijing, China). The apparent quantum yield (AQY) was estimated by the H<sub>2</sub>O<sub>2</sub> amount generated under monochromatic light and corresponding power density according to the equation:  $\text{AQY} (\%) = (\text{number of reacted}$

electrons/number of incident photons)  $\times 100 = (2 \times \text{number of produced H}_2\text{O}_2 \text{ molecules})/(\text{number of incident photons}) \times 100$ .

### 2.4. Photoelectrochemical measurements

The photoelectrochemical measurements were carried out on a CHI 760E electrochemical workstation (Shanghai Chenghua) by using a standard three-electrode system, in which the sample electrode, the Ag/AgCl electrode and the Pt plate electrode were served as the working electrode, the reference electrode and the counter electrode, respectively. Notably, the working electrode was prepared by deposition of photocatalyst suspension (10 mg mL<sup>-1</sup> in 5.0 vol% Nafion ethanol solution) on a cleaned fluorine-doped tin oxide (FTO) glass substrate with a surface area of  $1 \times 1$  cm<sup>2</sup>. The aqueous Na<sub>2</sub>SO<sub>4</sub> solution (0.5 M, pH = 6.6) was used as electrolyte.<sup>47</sup> The normal hydrogen electrode (NHE at pH = 0) can be converted from the reference electrode potential (vs. Ag/AgCl) according to  $E_{\text{NHE}} = E_{\text{Ag/AgCl}} + 0.059 \times \text{pH} + E_{\text{Ag/AgCl}}^0$ , where  $E_{\text{Ag/AgCl}}^0$  is 0.198 V at 25 °C.<sup>48</sup> The Mott-Schottky curves for flat potential confirmation were measured by impedance-potential method. The photocurrents were measured at a potential of 0.2 V (vs. Ag/AgCl) under on-off visible light irradiation. The electrochemical impedance spectroscopy (EIS) tests were performed in the frequency range of 0.01 Hz to 100 kHz.

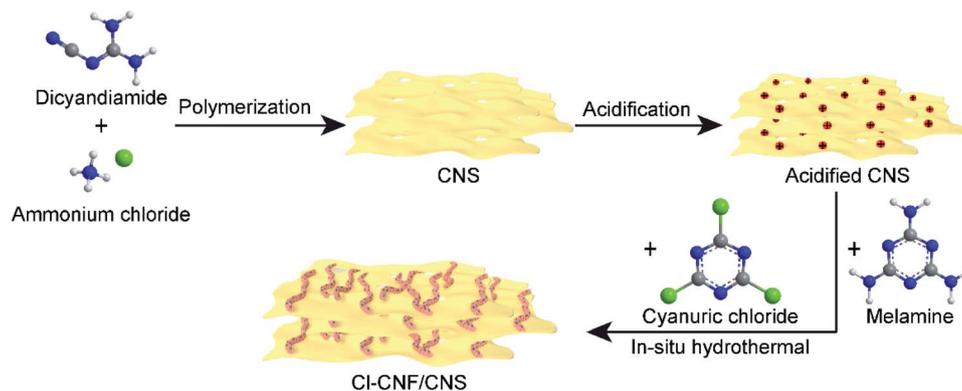
## 3. Results and discussion

### 3.1. Morphology, structure and property

The Cl-CNF/CNS heterostructure was synthesized by an *in situ* hydrothermal method in the presence of acidified CNS (Scheme 1). Due to the high negative potential of cyanuric chloride, the CNF precursors cannot be well adsorbed onto the negative-charged CNS surface (Fig. S1†). We therefore acidified CNS by HCl to change its surface potential (sample acidified CNS in Fig. S1†) in advance to ensure the adsorption of the Cl-CNF precursors. Pure Cl-CNF and CNS were also prepared for control experiments. As shown in Fig. 1a, the Cl-CNF sample synthesized by hydrothermal method exhibits a fibrous nanostructure with a diameter of about tens of nanometers and a length of several hundred nanometers. The EDX spectrum of Cl-CNF verified the successful doping of Cl element in the framework (Fig. S2†). The doping of Cl has been reported to be beneficial to adjust the energy band structure and inhibit the recombination of photogenerated charge carriers,<sup>40,49</sup> thus endow Cl-CNF with broadband spectral response. The CNS sample, obtained by calcining a mixture of dicyandiamide and ammonium chloride, exhibits wrinkled nanosheet characteristic (Fig. 1b). The sheet-like CNS surface can enlarge its specific surface area and provide sufficient space for Cl-CNF anchoring.<sup>25</sup> No Cl element was observed in the EDX spectrum of CNS (Fig. S2†), indicating ammonium chloride worked as a gas template other than a dopant during the calcination process.

In the Cl-CNF/CNS nanohybrid, Cl-CNF was tightly grown on the surface of CNS (Fig. 1c), indicating the successful





Scheme 1 Schematic illustration for the synthesis of the heterostructure.

formation of the heterostructure. The tightly contact interface formed between Cl-CNF and CNS is beneficial for the charge carrier migration and transfer in photocatalytic reaction. Moreover, the Cl content of Cl-CNF/CNS is remarkably decreased compared with that of pure Cl-CNF because of the existence of CNS substrate (Fig. S2 and Table S1<sup>†</sup>). The SEM elemental mapping images in Fig. 1d demonstrate that the C, N and Cl elements are uniformly distributed on the surface of the heterostructure, suggesting Cl-CNF is homogeneously bound

on the surface of CNS. In addition, the ratio of Cl-CNF to CNS can be adjusted by changing the amount of CNS added. However, Cl-CNF cannot be well loaded on CNS surface when CNS adding amount is too low or too high. When the CNS amount is too low, scattered Cl-CNF can be observed beyond the CNS surface (Fig. S3a<sup>†</sup>). On the other hand, when the CNS amount is too high, the morphology of Cl-CNF will be damaged during the hydrothermal process (Fig. S3b and c<sup>†</sup>).

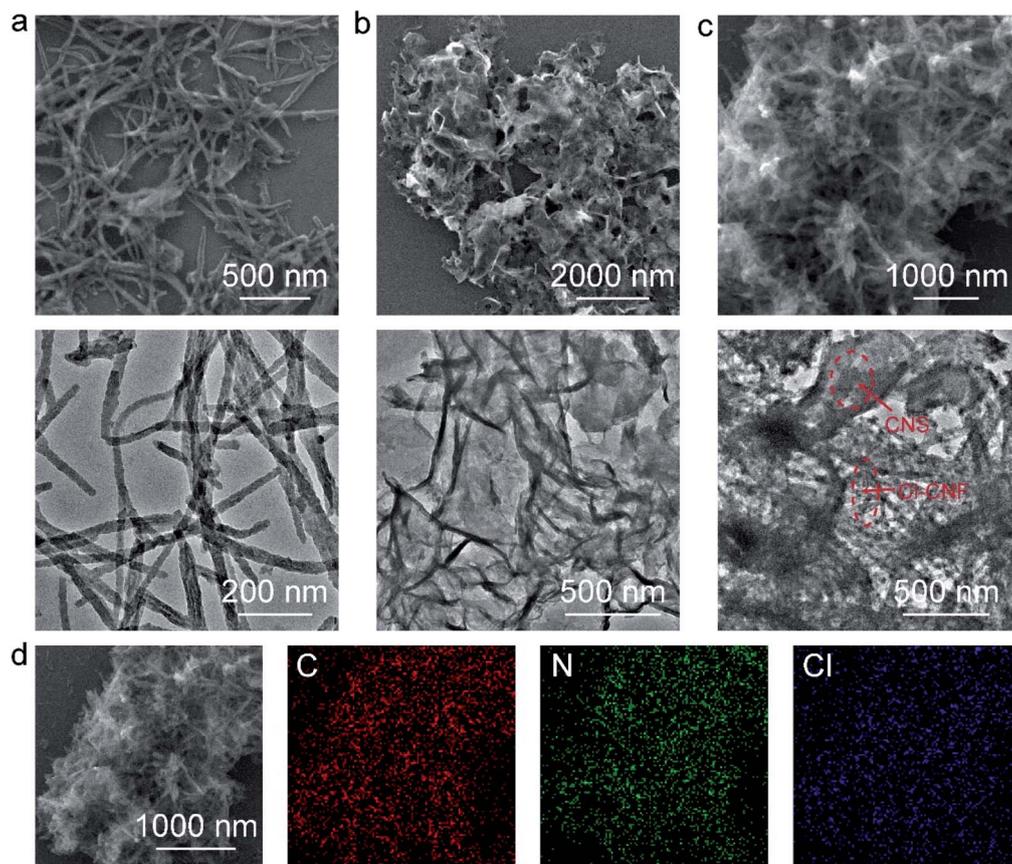


Fig. 1 Morphology of the photocatalysts. (a) SEM and TEM images of Cl-CNF. (b) SEM and TEM images of CNS. (c) SEM and TEM images of Cl-CNF/CNS. (d) SEM elemental mapping.



The crystal structures of the samples were investigated by their XRD patterns (Fig. 2a). CNS shows two characteristic diffraction peaks of  $13.0^\circ$  and  $27.6^\circ$  corresponding to  $g\text{-C}_3\text{N}_4$ , which can be indexed to the in-planar packing (1 0 0) and interlayer stacking (0 0 2) plane, respectively.<sup>50</sup> However, the peak of (1 0 0) plane in Cl-CNF vanishes, suggesting that the repeatability of the tri-*s*-triazine unit in the layer is significantly destroyed, which is probably caused by the Cl doping. In addition, the intensity of (0 0 2) peak in Cl-CNF decreases and the peak width at half height broadens, suggesting that the crystallinity of Cl-CNF is weakened intensively.<sup>14</sup> Moreover, compared with pure CNS, the (0 0 2) peak of Cl-CNF/CNS shifts to  $27.1^\circ$ , indicating an increase in the distance between interlayers. The intensities of the (1 0 0) and (0 0 2) peaks in Cl-CNF/CNS are between those in Cl-CNF and CNS, demonstrating that Cl-CNF and CNS are well integrated. Fig. 2b presents the FTIR spectra of the samples. The peaks range from 3000 to  $3500\text{ cm}^{-1}$  can be assigned to the stretching vibration of the N-H bond caused by the incomplete condensation of the precursor or the O-H bond of the adsorbed water molecules.<sup>51</sup> The peaks range from 1100 to  $1700\text{ cm}^{-1}$  can be indexed to the N=C-N heterocyclic stretches of tri-*s*-triazine unit. The peak at  $814\text{ cm}^{-1}$  represents the out-of-plane bending mode of tri-*s*-triazine unit,<sup>52,53</sup> which can only be observed in the periodic structure of CNS and the Cl-CNF/CNS nanohybrid, further verifying the disordered structure of Cl-CNF. However, no obvious N-Cl or C-Cl bonds appeared in the nanohybrid, probably because these bands overlap with the C-N stretching band.

The light harvesting abilities of the samples were characterized by their UV-vis absorption spectra (Fig. 2c). Due to its unique structure and Cl doping, Cl-CNF exhibits strong light absorption in the visible light region, and its absorption tail can even extend to  $\sim 628\text{ nm}$ , endowing it a superior semiconductor to enhance the light absorption ability of the heterostructure. The orange color of the Cl-CNF powder also reflects its strong light absorption ability and narrow bandgap characteristic (Fig. 2c, inset). On the contrary, CNS exhibits moderate visible light absorption capacity with a bandgap of 2.80 eV (Fig. 2c and S4<sup>†</sup>). After hybridizing, the light absorption ability of CNS was significantly enhanced in Cl-CNF/CNS, which is favorable for photocatalytic applications.

The valence states and charge transfer behaviors of the samples were analyzed by their XPS spectra. The trend of the Cl

amount in the survey spectra (Fig. 3a) is consistent with the EDX.

The results in Table S1,<sup>†</sup> indicating the successful hybridization of Cl-CNF and CNS. Fig. 3b shows the high-resolution C 1s spectra of the sample, the peaks of Cl-CNF in 288.3, 285.9 and 284.5 eV are respectively indexed to  $\text{sp}^2\text{ C}$  (N=C=N) in the tri-*s*-triazine unit, C-NH<sub>x</sub> groups on the edge of the framework and carbon contamination (C-C, C=C species) introduced during the condensation process. In CNS, the main peak of  $\text{sp}^2\text{ C}$  located at 287.9 eV, lowering than the binding energy value in the Cl-CNF/CNS nanohybrid. That is, the binding energy corresponding to  $\text{sp}^2\text{ C}$  in CNS was increased after it interacted with Cl-CNF. The results indicate that the electron density of CNS is reduced in the Cl-CNF/CNS nanohybrid, in other words, electrons are being transferred from CNS to Cl-CNF at the Cl-CNF/CNS interface. In the high-resolution N 1s spectra (Fig. 3c), the peaks of Cl-CNF centered at 398.7, 400.0, 400.9 and 405.2 eV are respectively assigned to  $\text{sp}^2\text{ N}$  (C=N=C), tertiary carbon-nitrogen bonds [N-(C)<sub>3</sub>], NH<sub>x</sub> groups and charge effect ( $\pi \rightarrow \pi^*$ ). Similar with the result of the high-resolution C 1s spectra of the samples, the  $\text{sp}^2\text{ N}$  peak of CNS also shows a lower binding energy relative to Cl-CNF/CNS, further demonstrating the electron is transferred from CNS to Cl-CNF. In addition, the obvious peaks of Cl element in Cl-CNF/CNS suggest Cl was successfully doped into the conjugate heptazine units (Fig. 3d).<sup>11,54</sup> The Cl 2p spectrum could be deconvoluted into two possible Cl sites of substitutional and interstitial sites (Cl<sub>sub</sub> and Cl<sub>int</sub>). The peaks centered at 202.3 and 200.6 eV can be indexed to the 2p<sub>1/2</sub> and 2p<sub>3/2</sub> peaks of Cl<sub>sub</sub> dopants stemmed from the C-Cl bonds in the framework. The other two peaks at 198.7 and 197.0 eV can be attributed to the Cl<sub>int</sub> dopants derived from the Cl<sup>-</sup> ions.<sup>55</sup> The photocatalytic activity is expected to be improved by Cl-doping due to its effect of broadening optical absorption and promoting charge migration.<sup>42</sup> Benefiting from the role of ammonium chloride as a gas template, the surface area of CNS ( $56.76\text{ m}^2\text{ g}^{-1}$ ) is higher than that of Cl-CNF ( $19.97\text{ m}^2\text{ g}^{-1}$ ) according to their nitrogen adsorption-desorption isothermal curves (Fig. S5<sup>†</sup>). Particularly, the Cl-CNF/CNS nanohybrid also possesses a large specific surface area ( $54.53\text{ m}^2\text{ g}^{-1}$ ), which can provide abundant active sites for photocatalytic reactions. Combining with the improved light absorption capacity, Cl-CNF/CNS is expected to show superior performance for H<sub>2</sub>O<sub>2</sub> generation.

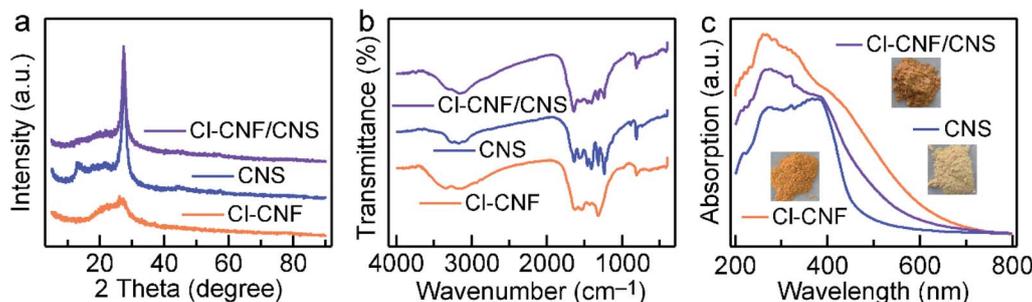


Fig. 2 Structures and properties of the samples. (a) XRD patterns. (b) FTIR spectra. (c) Absorption spectra of the samples. Inset shows the photos of the sample powders.



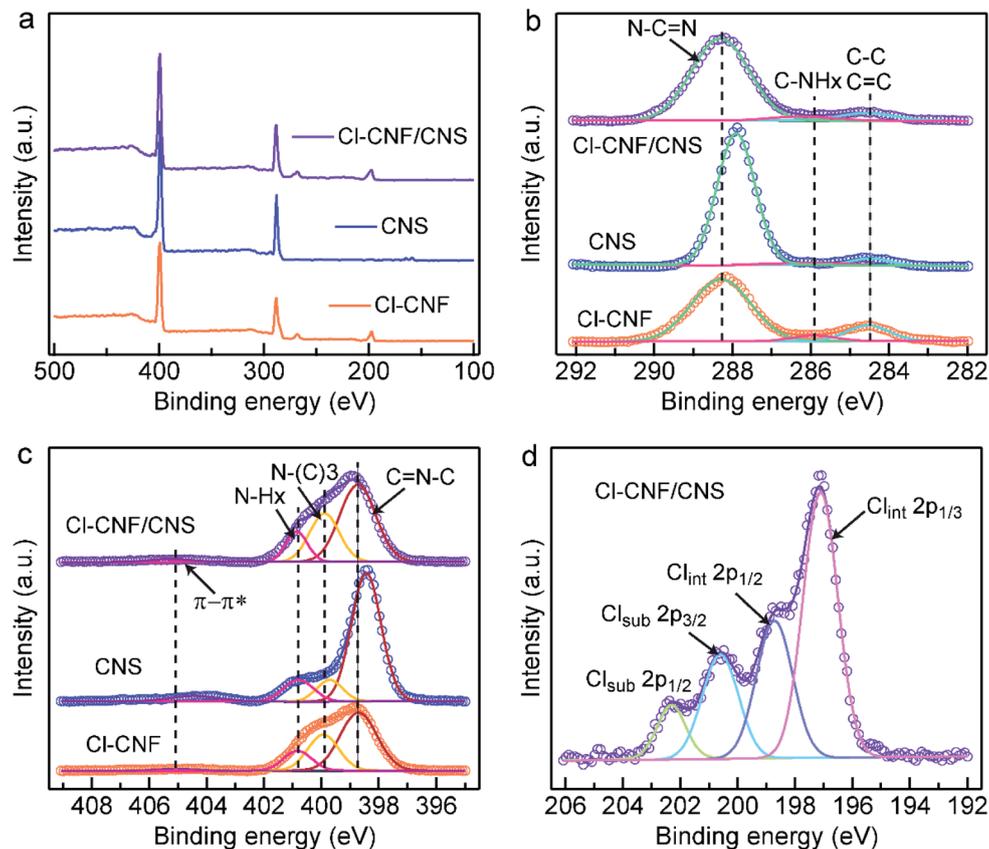


Fig. 3 XPS spectra of the samples. (a) XPS survey spectra the three samples. (b) High-resolution C 1s spectra of CI-CNF, CNS and CI-CNF/CNS. (c) High-resolution N 1s spectra of CI-CNF, CNS and CI-CNF/CNS. (d) High-resolution Cl 2p spectrum of CI-CNF/CNS.

### 3.2. Photocatalytic H<sub>2</sub>O<sub>2</sub> evolution

The photocatalytic activity of the samples was evaluated by H<sub>2</sub>O<sub>2</sub> generation with ethanol as an electron donor. In our work, H<sub>2</sub>O<sub>2</sub> was assumed to be produced through the two-electron reduction reaction of oxygen molecules. The H<sub>2</sub>O<sub>2</sub> concentration of all the samples increased with prolonging the reaction time (Fig. 4a). Through screening experiment, the CI-CNF/CNS-0.4 (always abbreviated as CI-CNF/CNS) nano-hybrid is found to show the highest H<sub>2</sub>O<sub>2</sub> production rate of 247.5  $\mu\text{mol g}^{-1} \text{h}^{-1}$  under visible light irradiation ( $\lambda > 420 \text{ nm}$ ), which is about 3.4 and 3.1 times as much as those of CI-CNF and CNS (Fig. 4a and b). The detectable activity of pure CI-CNF and CNS toward H<sub>2</sub>O<sub>2</sub> generation demonstrates that the conduction bands of both components are more negative than the redox potential of O<sub>2</sub>/H<sub>2</sub>O<sub>2</sub> (0.68 V vs. NHE at pH = 0). Moreover, the photocatalytic activities of the four nano-hybrids with different CNS contents toward H<sub>2</sub>O<sub>2</sub> generation are higher than those of pure CI-CNF and CNS. The enhanced activity of the nano-hybrids is believed to be caused by the synergistic effect of the improved photo-generated charge carriers separation efficiency and the enhanced light harvesting ability in the heterostructure.

The photocatalytic activity of the samples was evaluated by H<sub>2</sub>O<sub>2</sub> generation with ethanol as an electron donor. In our work, H<sub>2</sub>O<sub>2</sub> was assumed to be produced through the two-electron reduction reaction of oxygen molecules. The H<sub>2</sub>O<sub>2</sub>

concentration of all the samples increased with prolonging the reaction time (Fig. 4a). Through screening experiment, the CI-CNF/CNS-0.4 (always abbreviated as CI-CNF/CNS) nano-hybrid is found to show the highest H<sub>2</sub>O<sub>2</sub> production rate of 247.5  $\mu\text{mol g}^{-1} \text{h}^{-1}$  under visible light irradiation ( $\lambda > 420 \text{ nm}$ ), which is about 3.4 and 3.1 times as much as those of CI-CNF and CNS (Fig. 4a and b). The detectable activity of pure CI-CNF and CNS toward H<sub>2</sub>O<sub>2</sub> generation demonstrates that the conduction bands of both components are more negative than the redox potential of O<sub>2</sub>/H<sub>2</sub>O<sub>2</sub> (0.68 V vs. NHE at pH = 0). Moreover, the photocatalytic activities of the four nano-hybrids with different CNS contents toward H<sub>2</sub>O<sub>2</sub> generation are higher than those of pure CI-CNF and CNS. The enhanced activity of the nano-hybrids is believed to be caused by the synergistic effect of the improved photo-generated charge carriers separation efficiency and the enhanced light harvesting ability in the heterostructure.

To investigate the effect of CI-CNF and CNS in CI-CNF/CNS for H<sub>2</sub>O<sub>2</sub> generation activity improvement, we have also prepared other control samples for comparison. Among the carbon nitrides synthesized by different precursors, the H<sub>2</sub>O<sub>2</sub> generation rates of melamine-polymerized carbon nitride and dicyandiamide-polymerized carbon nitride are both lower than that of CNS (Fig. S6<sup>†</sup>), which is probably due to the suitable energy band structure and large specific surface area of CNS, verifying the crucial role of ammonium chloride. Expectedly,



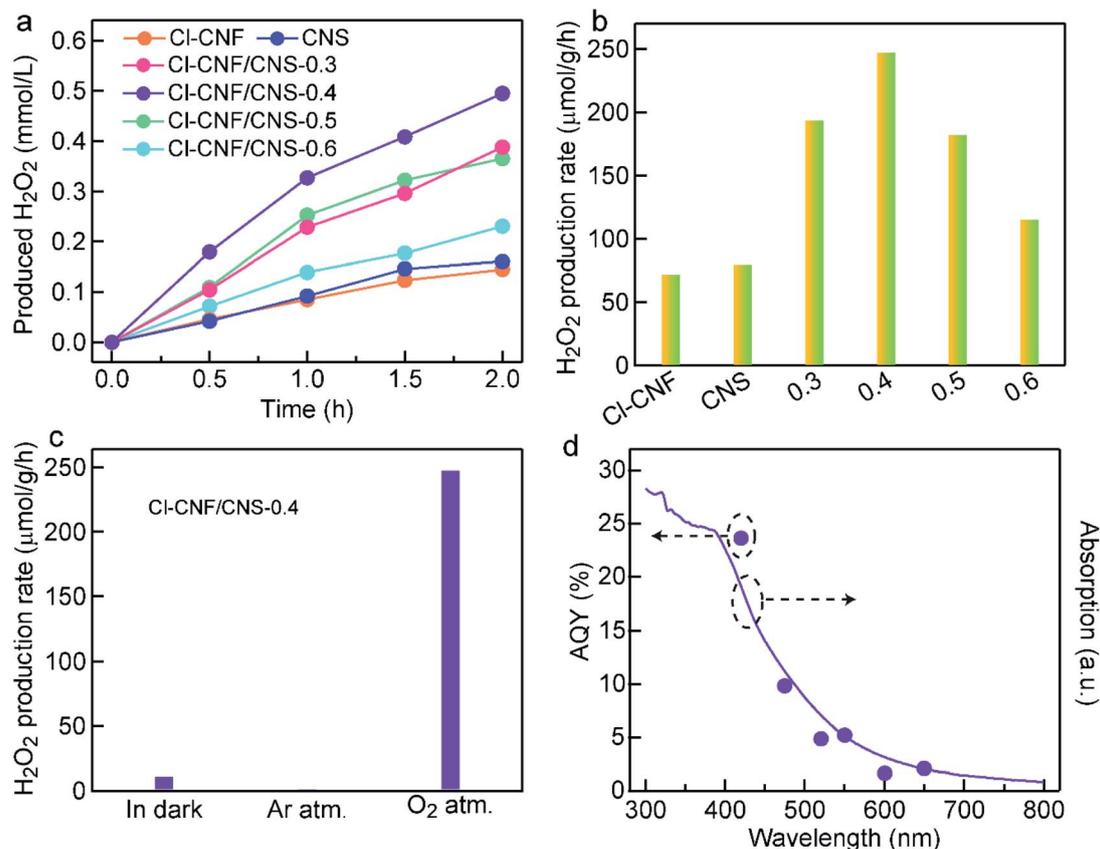


Fig. 4 Photocatalytic H<sub>2</sub>O<sub>2</sub> generation under visible light. (a) Time courses of the samples for H<sub>2</sub>O<sub>2</sub> generation. (b) Photocatalytic H<sub>2</sub>O<sub>2</sub> generation rates of the various samples. The samples of 0.3, 0.4, 0.5 and 0.6 represent Cl-CNF/CNS-0.3, Cl-CNF/CNS-0.4, Cl-CNF/CNS-0.5 and Cl-CNF/CNS-0.6. (c) Photocatalytic H<sub>2</sub>O<sub>2</sub> generation rates of Cl-CNF/CNS under different conditions. (d) AQYs and absorption spectrum of Cl-CNF/CNS.

the H<sub>2</sub>O<sub>2</sub> generation rate of Cl-CNF/melamine-derived carbon nitride (nanohybrid of Cl-CNF and melamine-polymerized carbon nitride) and Cl-CNF/dicyandiamide-derived carbon nitride (nanohybrid of Cl-CNF and dicyandiamide-polymerized carbon nitride) are both enhanced compared with their pure components, further implying Cl-CNF is a favorable semiconductor to construct heterojunctions for H<sub>2</sub>O<sub>2</sub> generation. Moreover, the photocatalytic activity of Cl-CNF/CNS toward H<sub>2</sub>O<sub>2</sub> generation is the highest among the nanohybrids with the same carbon nitride addition content, indicating the abundant active sites and strong light absorption ability are both vital and indispensable for photocatalysis performance enhancement.

In order to verify our assumption that H<sub>2</sub>O<sub>2</sub> was generated through two-electron reduction reaction of oxygen in our work, the H<sub>2</sub>O<sub>2</sub> generation reactions were carried out under different conditions (Fig. 4c). A trace amount of H<sub>2</sub>O<sub>2</sub> was detected in darkness, demonstrating that light is the energy source to drive the photocatalytic reaction. Negligible amount of H<sub>2</sub>O<sub>2</sub> was detected under Ar atmosphere, indicating the presence of O<sub>2</sub> is a prerequisite for H<sub>2</sub>O<sub>2</sub> production. Control experiments under different atmospheres verify that H<sub>2</sub>O<sub>2</sub> was produced *via* light-driven two-electron reduction process of oxygen in our work. H<sub>2</sub>O<sub>2</sub> decomposition is an important factor to restrict the activity improvement for H<sub>2</sub>O<sub>2</sub> generation, thus the

decomposition behavior of H<sub>2</sub>O<sub>2</sub> over Cl-CNF/CNS was investigated under visible light irradiation. As shown in Fig. S7,† Cl-CNF/CNS can lead to a small amount decomposition of H<sub>2</sub>O<sub>2</sub> under light irradiation due to the instability of H<sub>2</sub>O<sub>2</sub>. The result also explained why the growth rate of H<sub>2</sub>O<sub>2</sub> concentration in the reaction solution gradually decreased with prolonging the reaction time in Fig. 4a. Under monochromatic light irradiation, the AQY of Cl-CNF/CNS was measured and compared with its absorption spectrum (Fig. 4d). The trend of the AQY values agrees well with the absorption spectrum, indicating that the two-electron oxygen reduction reaction in the formation of H<sub>2</sub>O<sub>2</sub> is light-driven. Especially, the AQY value reaches 23.67% at 420 nm, which is higher than many reported values (Table S2†), manifesting the Cl-CNF/CNS heterostructure is a superior catalyst for photocatalytic H<sub>2</sub>O<sub>2</sub> generation.

The cycle test was performed to investigate the stability of Cl-CNF/CNS. As shown in Fig. 5a, the Cl-CNF/CNS heterostructure exhibits favorable stability for photocatalytic H<sub>2</sub>O<sub>2</sub> evolution. The small reduction of H<sub>2</sub>O<sub>2</sub> generation rate in the second and third run is probably caused by the catalyst loss during centrifugation. According to the XRD patterns and FTIR spectra comparisons between the fresh and used Cl-CNF/CNS samples in Fig. 5b and c, no obvious change was occurred in the crystal and chemical structure of Cl-CNF/CNS after cyclic



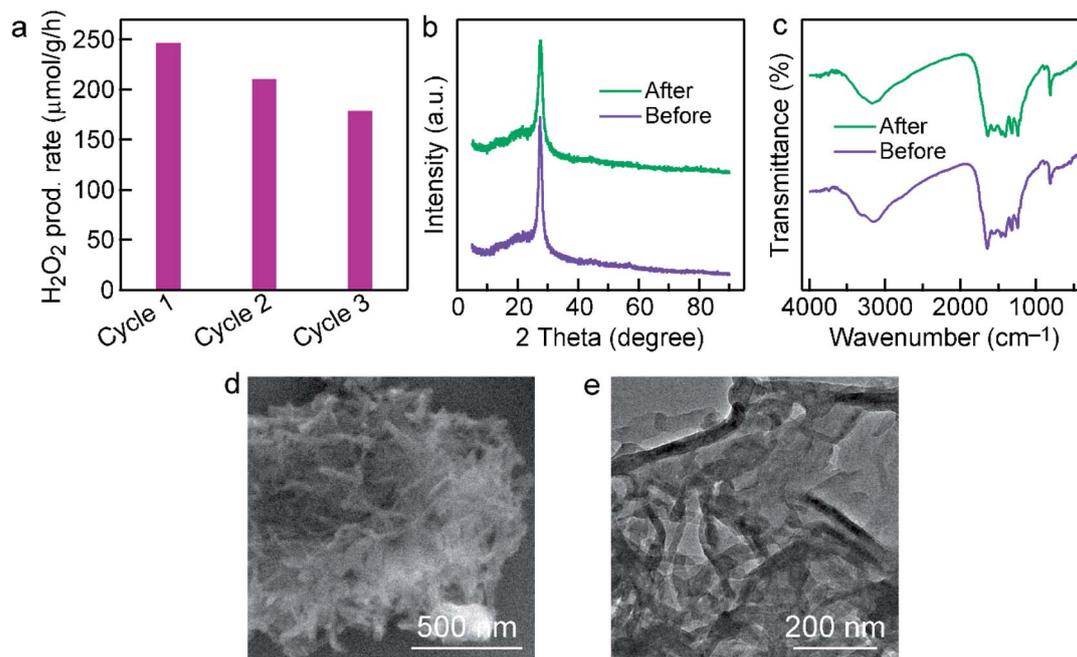


Fig. 5 Stability test of Cl-CNF/CNS. (a) Cyclic test of Cl-CNF/CNS for H<sub>2</sub>O<sub>2</sub> generation. (b) XRD patterns and (c) FTIR spectra of Cl-CNF/CNS before and after stability test. (d) SEM and (e) TEM images of Cl-CNF/CNS after stability test.

test. Additionally, there is no obvious change in the morphology of Cl-CNF/CNS after stability test (Fig. 5b and c), further confirming the stability of the photocatalyst.

### 3.3. Mechanism analysis

To uncover the mechanism for photocatalytic activity improvement in the heterostructure, photocurrent measurements of the three samples (Cl-CNF, CNS and Cl-CNF/CNS) were first carried out under discontinuous visible light ( $\lambda > 420$  nm) to investigate the generation and separation of photoexcited charge carriers (Fig. 6a). The results show that the current density of Cl-CNF/CNS is higher than those of Cl-CNF and CNS, indicating that the generation and separation efficiency of photogenerated charge carriers was significantly improved compared with pure Cl-CNF and CNS electrodes, the enhanced charge carriers separation efficiency is beneficial for photocatalytic H<sub>2</sub>O<sub>2</sub> generation. The EIS curves of the samples show that the Cl-CNF/CNS heterojunction processes the smallest charge transfer resistance whether the measurements were performed under light irradiation (Fig. 6b) or in dark (Fig. S8<sup>†</sup>), indicating the fastest charge transfer and migration rate in Cl-CNF/CNS.

The separation and transfer properties of the photo-generated charge carriers were further investigated by the PL spectra (Fig. 6c). The PL intensity of CNS is the strongest, suggesting its serious radiative recombination of photogenerated electron-hole pairs.<sup>56</sup> The PL peak position of Cl-CNF redshifts to  $\sim 525$  nm, corresponding to its bandgap excitation of 2.36 eV. The PL intensity of Cl-CNF/CNS decreases in comparison with that of CNS, demonstrating the radiative recombination of the photogenerated charge carriers was effectively inhibited. As shown in Fig. 6d, the TRPL spectra were also collected to

calculate the lifetime of the charge carriers by a fitted bi-exponential equation:  $I(t) = A_1 \exp(-t/\tau_1) + A_2 \exp(-t/\tau_2)$ , where  $\tau_1$ ,  $\tau_2$  are the emission lifetime values, and  $A_1$ ,  $A_2$  are the corresponding amplitudes. The calculated lifetime values of the samples are shown in Table 1. The lifetime of Cl-CNF/CNS is shortened compared with that of CNS, indicating the improved charge transfer kinetic for photocatalytic reactions.

The Mott-Schottky curves of Cl-CNF and CNS were measured to calculate their band structures and further depict the charge transfer direction (Fig. S9<sup>†</sup>). Their positive slopes indicate that they are both n-type semiconductors.<sup>53</sup> The flat potentials of Cl-CNF and CNS, which are obtained according to the intersection of the X-axis and the slopes at different frequencies, are directly deemed as their conduction band. The conduction band potentials of Cl-CNF and CNS are calculated to be  $-1.21$  and  $-1.71$  V (*vs.* Ag/AgCl), respectively, which are respectively  $-0.60$  and  $-1.10$  V *vs.* NHE at pH = 0. The pure Cl-CNF and CNS catalysts can therefore reduce oxygen molecules to H<sub>2</sub>O<sub>2</sub>. Combining with the bandgap values in Fig. S4<sup>†</sup>, the band structures of Cl-CNF and CNS were obtained and presented in Fig. 5e. We therefore proposed the charge transfer mechanism for photocatalytic H<sub>2</sub>O<sub>2</sub> production over Cl-CNF/CNS. Upon light excitation, the photogenerated electrons transferred from the conduction band of CNS to that of Cl-CNF due to the more negative conduction band potential of CNS, which is consistent with the XPS results. The electrons on the conduction band of Cl-CNF then react with oxygen molecules through two-electron reduction reaction to generate H<sub>2</sub>O<sub>2</sub>. On the contrary, photogenerated holes left on the valence band of Cl-CNF transferred to that of CNS for ethanol oxidation (Fig. 6e). Thus, the photogenerated charge carriers can be



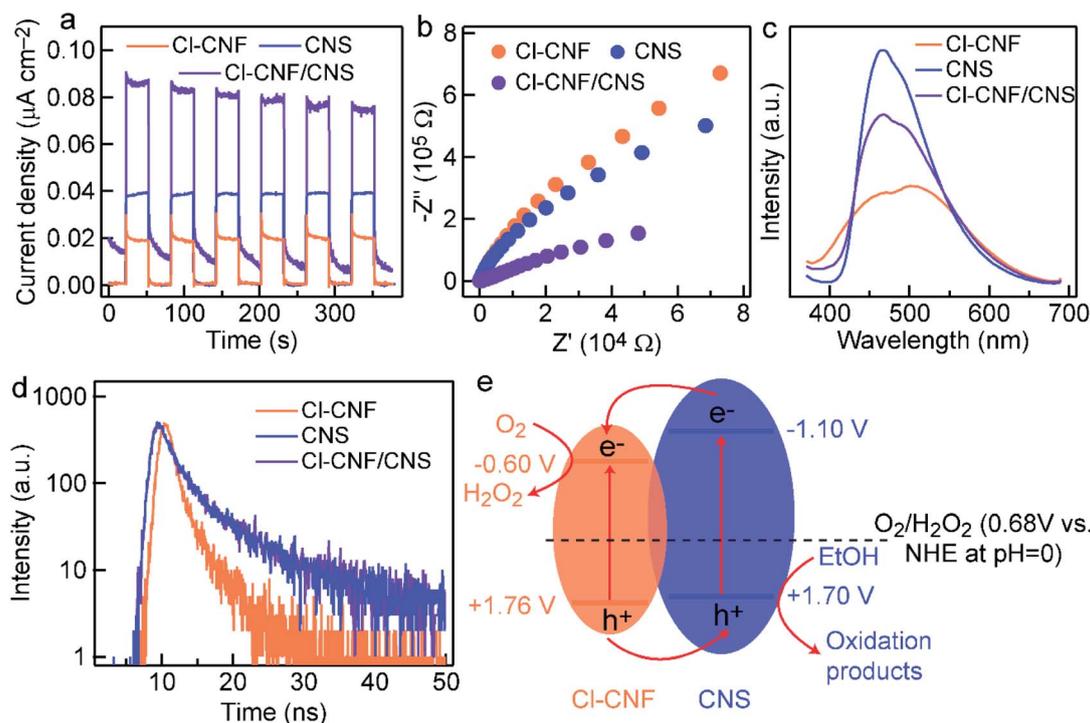


Fig. 6 Proposed charge transfer mechanism in the Cl-CNF/CNS heterostructure. (a) Photocurrent densities of the sample electrodes under chopped visible light illumination. (b) EIS curves of the electrodes under light irradiation. (c) PL and (d) TRPL spectra of the samples. (e) The possible charge transfer behaviors at the Cl-CNF/CNS interface.

Table 1 The charge carriers lifetime of the samples

	$\tau_1$ (ns)	$\tau_2$ (ns)	$A_1$ (%)	$A_2$ (%)
Cl-CNF	1.1354	4.0053	74.21	25.79
CNS	2.2483	10.8263	58.09	41.91
Cl-CNF/CNS	2.0915	10.1601	53.15	46.85

effectively separated to participate the redox reactions and improve the photocatalytic activity.

## 4. Conclusions

In summary, a 1D/2D heterostructure was synthesized by grafting narrow-bandgap Cl-CNF on CNS substrate through an *in situ* hydrothermal method. The light harvesting ability of CNS was significantly enhanced due to the introduction of Cl-CNF with broadband spectral response. Moreover, the heterojunction shows a favorable photogenerated charge carriers separation efficiency, which is beneficial to improve its photocatalytic performance. The optimal Cl-CNF/CNS sample exhibits a H<sub>2</sub>O<sub>2</sub> generation rate of 247.5 μmol g<sup>-1</sup> h<sup>-1</sup> under visible light irradiation, which is 3.4 and 3.1 times as much as those of Cl-CNF and CNS, respectively. Especially, Cl-CNF/CNS reaches a high AQY value of 23.67% at 420 nm. Our work provides a feasible method for constructing heterojunctions with different dimensions to improve the photogenerated charge carriers separation ability and the photocatalytic performance.

## Conflicts of interest

There are no conflicts of interest to declare.

## Acknowledgements

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## References

- 1 L. Chen, L. Wang, Y. Wan, Y. Zhang, Z. Qi, X. Wu and H. Xu, *Adv. Mater.*, 2020, **32**, 1904433.
- 2 S. Wu, H. Yu, S. Chen and X. Quan, *ACS Catal.*, 2020, **10**, 14380–14389.
- 3 H. Zhang, L. Jia, P. Wu, R. Xu, J. He and W. Jiang, *Appl. Surf. Sci.*, 2020, **527**, 146584.
- 4 P. Zhang, Y. Tong, Y. Liu, J. J. M. Vequizo, H. Sun, C. Yang, A. Yamakata, F. Fan, W. Lin, X. Wang and W. Choi, *Angew. Chem., Int. Ed.*, 2020, **59**, 16209–16217.
- 5 P. Raizada, A. A. P. Khan and P. Singha, *Sep. Purif. Technol.*, 2020, **247**, 116957.
- 6 A. Sudhaik, P. Raizada, S. Thakur, R. V. Saini, A. K. Saini, P. Singh, V. K. Thakur, V.-H. Nguyen, A. A. P. Khan and A. M. Asirif, *J. Taiwan Inst. Chem. Eng.*, 2020, **113**, 142–154.



- 7 A. Sudhaik, P. Raizada, P. Singh, A. Hosseini-Bandegharaei, V. K. Thakur and V.-H. Nguyen, *J. Environ. Chem. Eng.*, 2020, **8**, 104483.
- 8 C. Feng, L. Tang, Y. Deng, J. Wang, J. Luo, Y. Liu, X. Ouyang, H. Yang, J. Yu and J. Wang, *Adv. Funct. Mater.*, 2020, **30**, 2001922.
- 9 A. Ullah, J. Khan, M. Sohail, A. Hayat, T. K. Zhao, B. Ullah, M. Khan, I. Uddin, S. Ullah, R. Ullah, A. U. Rehman and W. U. Khan, *J. Photochem. Photobiol., A*, 2020, **401**, 112764.
- 10 J. Fu, J. Yu, C. Jiang and B. Cheng, *Adv. Energy Mater.*, 2018, **8**, 1701503.
- 11 L. Shi, L. Yang, W. Zhou, Y. Liu, L. Yin, X. Hai, H. Song and J. Ye, *Small*, 2018, **14**, 1703142.
- 12 B. Wu, L. Zhang, B. Jiang, Q. Li, C. Tian, Y. Xie, W. Li and H. Fu, *Angew. Chem., Int. Ed.*, 2021, **60**, 4815–4822.
- 13 Q. Xu, B. Zhu, C. Jiang, B. Cheng and J. Yu, *Sol. RRL*, 2018, **2**, 1800006.
- 14 Y. Chen, X. Liu, L. Hou, X. Guo, R. Fu and J. Sun, *Chem. Eng. J.*, 2020, **383**, 123132.
- 15 H. Yu, R. Shi, Y. Zhao, T. Bian, Y. Zhao, C. Zhou, G. I. N. Waterhouse, L. Z. Wu, C. H. Tung and T. Zhang, *Adv. Mater.*, 2017, **29**, 1605148.
- 16 H. Yu, H. Ma, X. Wu, X. Wang, J. Fan and J. Yu, *Sol. RRL*, 2021, **5**, 2000372.
- 17 J. Jia, W. Huang, C. Feng, Z. Zhang, K. Zuojiào, J. Liu, C. Jiang and Y. Wang, *J. Alloys Compd.*, 2019, **790**, 616–625.
- 18 S. Li, G. Dong, R. Hailili, L. Yang, Y. Li, F. Wang, Y. Zeng and C. Wang, *Appl. Catal., B*, 2016, **190**, 26–35.
- 19 X. Li, J. Zhang, F. Zhou, H. Zhang, J. Bai, Y. Wang and H. Wang, *Chinese. J. Catal.*, 2018, **39**, 1090–1098.
- 20 X. Li, X. Sun, L. Zhang, S. Sun and W. Wang, *J. Mater. Chem. A*, 2018, **6**, 3005–3011.
- 21 W. Wang, H. Zhang, S. Zhang, Y. Liu, G. Wang, C. Sun and H. Zhao, *Angew. Chem., Int. Ed.*, 2019, **58**, 16644–16650.
- 22 Y. Liu, X. Huang, Z. Yu, L. Yao, S. Guo and W. Zhao, *ChemistrySelect*, 2020, **5**, 7720–7727.
- 23 Q. Wang, Y. Shi, Z. Du, J. He, J. Zhong, L. Zhao, H. She, G. Liu and B. Su, *Eur. J. Inorg. Chem.*, 2015, **2015**, 4108–4115.
- 24 Y. Wu, H. Liao and M. Li, *Micro. Nano. Lett.*, 2018, **13**, 752–757.
- 25 J. Niu, K. Wang, Z. Ma, F. Yang and Y. Zhang, *ChemistrySelect*, 2020, **5**, 12353–12364.
- 26 J. Fu, B. Zhu, C. Jiang, B. Cheng, W. You and J. Yu, *Small*, 2017, **13**, 1603938.
- 27 L. Cheng, H. Yin, C. Cai, J. Fan and Q. Xiang, *Small*, 2020, **16**, 2002411.
- 28 Y. Xie, Y. Zhuo, S. Liu, Y. Lin, D. Zuo, X. Wu, C. Li and P. Wong, *Sol. RRL*, 2020, **4**, 1900440.
- 29 I. Camuossi, B. Mannucci, A. Speltini, A. Profumo, C. Milanese, L. Malavasi and P. Quadrelli, *ACS Sustainable Chem. Eng.*, 2019, **7**, 8176–8182.
- 30 N. D. Shcherban, P. Mäki-Arvela, A. Aho, S. A. Sergiienko, P. S. Yaremov, K. Eränenband and D. Y. Murzin, *Catal. Sci. Technol.*, 2018, **8**, 2928–2937.
- 31 Y. Guo, B. Chang, T. Wen, S. Zhang, M. Zeng, N. Hu, Y. Su, Z. Yang and B. Yang, *J. Colloid Interface Sci.*, 2020, **567**, 213–223.
- 32 H. Li, G. Huang, H. Xu, Z. Yang, X. Xu, J. Li, A. Qu and Y. Chen, *J. Solid State Chem.*, 2020, **292**, 121705.
- 33 N. T. T. Truc, T. D. Pham, M. V. Nguyen, D. V. Thuan, D. Q. Trung, P. Thao, H. T. Trang, V. N. Nguyen, D. T. Tran, D. N. Minh, N. T. Hanh and H. M. Ngoc, *J. Alloys Compd.*, 2020, **842**, 155860.
- 34 X. Wang, D. Han, Y. Ding, J. Liu, H. Cai, L. Jia, X. Cheng, J. Wang and X. Fan, *J. Alloys Compd.*, 2020, **845**, 156293.
- 35 Y. Cui, Z. Ding, X. Fu and X. Wang, *Angew. Chem., Int. Ed.*, 2012, **51**, 11814–11818.
- 36 Q. Gu, Y. Liao, L. Yin, J. Long, X. Wang and C. Xue, *Appl. Catal., B*, 2015, **165**, 503–510.
- 37 D. Liu, D. Chen, N. Li, Q. Xu, H. Li, J. He and J. Lu, *Angew. Chem., Int. Ed.*, 2020, **59**, 4519–4524.
- 38 Q. Zhu, B. Qiu, M. Du, J. Ji, M. Nasir, M. Xing and J. Zhang, *ACS Sustainable Chem. Eng.*, 2020, **8**, 7497–7502.
- 39 P. H. Linh, P. D. Chung, N. V. Khien, L. T. M. Oanh, V. T. Thu, T. N. Bach, L. T. Hang, N. M. Hung and V. D. Lam, *Diam. Relat. Mater.*, 2021, **111**, 108214.
- 40 D. Wang, X. Huang, Y. Huang, X. Yu, Y. Lei, X. Dong and Z. Su, *Colloids Surf., A*, 2021, **611**, 125780.
- 41 A. Sudhaik, P. Raizada, S. Thakur, A. K. Saini, P. Singh, A. Hosseini-Bandegharaei, J.-H. Lim, D. Y. Jeong and V.-H. Nguyen, *Appl. Nanosci.*, 2020, **10**, 4115–4137.
- 42 Z. Cheng, K. Y. Zheng, G. Y. Lin, S. Q. Fang, L. Y. Li, J. H. Bi, J. N. Shen and L. Wu, *Nanoscale Adv.*, 2019, **1**, 2674–2680.
- 43 T. Paul, D. Das, B. K. Das, S. Sarkar, S. Maiti and K. K. Chattopadhyay, *J. Hazard. Mater.*, 2019, **380**, 120855.
- 44 X. Hao, J. Zhou, Z. Cui, Y. Wang, Y. Wang and Z. Zou, *Appl. Catal., B*, 2018, **229**, 41–51.
- 45 S. Wei, F. Wang, P. Yan, M. Dan, W. Cen, S. Yu and Y. Zhou, *J. Catal.*, 2019, **377**, 122–132.
- 46 Z. L. He, J. Zhang, X. Li, S. N. Guan, M. C. Dai and S. G. Wang, *Small*, 2020, **16**, 2005051.
- 47 S. Hong, S. Lee, J. Jang and J. Lee, *Energ Environ. Sci.*, 2011, **4**, 1781–1787.
- 48 W. Li, P. Da, Y. Zhang, Y. Wang, X. Lin, X. Gong and G. Zheng, *ACS Nano*, 2014, **8**, 11770–11777.
- 49 F. Guo, M. Li, H. Ren, X. Huang, K. Shu, W. Shi and C. Lu, *Sep. Purif. Technol.*, 2019, **228**, 115770.
- 50 Q. Gao, J. Xu, Z. Wang and Y. Zhu, *Appl. Catal., B*, 2020, **271**, 118933.
- 51 C. Zhao, C. Ding, C. Han, X. Yang and J. Xu, *Sol. RRL*, 2021, **5**, 2000486.
- 52 G. Tzvetkov, M. Tzvetkov and T. Spassov, *Superlattices Microstruct.*, 2018, **119**, 122–133.
- 53 W. Wang, H. Zhang, S. Zhang, Y. Liu, G. Wang, C. Sun and H. Zhao, *Angew. Chem., Int. Ed.*, 2019, **58**, 16644–16650.
- 54 X. Fang, S. Cai, M. Wang, Z. Chen, C. Lu and H. Yang, *Angew. Chem., Int. Ed.*, 2021, **60**, 7046–7050.
- 55 M. Y. Cao, K. Wang, I. Tudela and X. F. Fan, *Appl. Surf. Sci.*, 2021, **536**, 147784.
- 56 S. Hu, X. Sun, Y. Zhao, W. Li, H. Wang and G. Wu, *J. Taiwan Inst. Chem. Eng.*, 2020, **107**, 129–138.

