

PAPER

[View Article Online](#)
[View Journal](#) | [View Issue](#)
Cite this: *Nanoscale*, 2020, **12**, 10010

The synthesis of interface-modulated ultrathin Ni(II) MOF/g-C₃N₄ heterojunctions as efficient photocatalysts for CO₂ reduction†

 Lina Zhao,^{a,b} Zhenlong Zhao,^a Yuxin Li,^a ^a Xiaoyu Chu,^{a,b} Zhijun Li,^a Yang Qu,^a ^a Linlu Bai ^a and Liqiang Jing ^a

It is highly desirable to improve charge separation and to provide catalytic functions for the efficient photocatalytic CO₂ reduction reaction (CO₂RR) on g-C₃N₄ (CN). Here, dimension-matched ultrathin NiMOF/CN heterojunctions have been successfully constructed by the *in situ* growth of NiMOF nanosheets on hydroxylated and 1,4-aminobenzoic acid (AA) functionalized CN nanosheets, respectively, with ultrasonic assistance. The resultant NiMOF/CN heterojunctions exhibited excellent photocatalytic activities for the CO₂RR to produce CO and CH₄, especially NiMOF/CN-AA, which had photoactivity 18 times higher than that of bare CN. Based on the surface photovoltage responses, wavelength-dependent photocurrent action spectra, electrochemical impedance spectra, and CO₂ electrochemical reduction data, it is clearly confirmed that the exceptional photoactivity mainly resulted from the favorable charge transport properties of ultrathin CN and coupled NiMOF, and from the greatly enhanced charge separation *via* excited high-level electron transfer from CN to NiMOF in the resultant intimately contacted heterojunction caused by the induction effect of AA, and also from the provided catalytic functionality of the central Ni(II) for CO₂ activation. This work provides a feasible synthetic protocol to fabricate MOF-containing dimension-matched heterojunctions with good charge separation for efficient photocatalysis.

Received 31st March 2020,
Accepted 15th April 2020

DOI: 10.1039/d0nr02551h

rsc.li/nanoscale

Introduction

The photocatalytic CO₂ reduction reaction (CO₂RR) into valuable fuels and chemicals is considered as a sustainable and promising pathway to lessen fossil fuel depletion and related environmental pollution.^{1–4} Among various semiconductor photocatalysts, graphitic carbon nitride (CN) possesses a moderate band gap to harvest visible light and a relatively negative conduction band (CB) with sufficient thermodynamic energy^{5,6} and hence becomes a suitable photocatalyst candidate.⁷ Nevertheless, bulk CN always exhibits unfavourable photocatalytic performance due to the low specific surface area, rapid recombination of photogenerated electron–hole pairs, along with weak adsorption and activation of

reactants.^{8,9} Therefore, it is desirable to develop feasible strategies to fabricate highly active CN-based photocatalysts for CO₂RR. Controlling the morphology to obtain ultrathin two-dimensional (2D) CN nanosheets with an enlarged surface area and shortened charge transfer distance might remedy the intrinsic drawbacks of bulk CN to a certain extent. Based on CN nanosheets the construction of heterojunctions with traditional semiconductors like TiO₂ and Fe₂O₃^{10,11} *etc.* or the introduction of noble metals are strategies commonly performed to further improve the photoactivity.^{12–15} However, the dimension mismatching and loosely-contacted interface of the fabricated heterojunctions, still comprehensively exist, which would hinder efficient interfacial charge transfer. Accordingly, it is significant to construct novel dimension-matched g-C₃N₄-based heterojunctions with an intimately contacted interface and catalytic functions.

In recent years, metal–organic frameworks (MOFs), constructed with coordinating inorganic metal ions and organic linkers, have attracted great attention in photocatalysis as promising semiconductor materials. MOFs have characteristic merits such as uniformly scattered active sites as catalytic centres, enriched pore systems and large specific surface areas, which are all adjustable by modulating the multiscale

^aKey Laboratory of Functional Inorganic Materials Chemistry (Ministry of Education), School of Chemistry and Materials Science, International Joint Research Center and Lab for Catalytic Technology, Heilongjiang University, Harbin 150080, P. R. China. E-mail: liyuxin@hlju.edu.cn, lbai2@e.ntu.edu.sg, jinglq@hlju.edu.cn

^bDepartment of Food & Environmental Engineering, East University of Heilongjiang, Harbin 150066, P. R. China

†Electronic supplementary information (ESI) available. See DOI: 10.1039/d0nr02551h

structures.^{16–19} Therefore, it is feasible to incorporate suitable MOFs with CN to construct heterojunctions and meanwhile introducing uniform catalytic sites.^{20–22} To date, some MOF/CN heterojunctions such as UiO-66/g-C₃N₄,²³ ZIF-8/g-C₃N₄,²⁴ BIF-20@g-C₃N₄²⁵ and NH₂-UiO-66/NH_x-CN,²⁶ have already been reported to be effective for photocatalytic CO₂RR. However, the MOFs involved are mostly three-dimensional (3D), hence rendering long-range distances of photogenerated carriers and especially poor dimension-matching with CN nanosheets.²⁷ Moreover, the post-combination of MOF with CN would lead to less contacted interfaces.^{28,29} Therefore, it is necessary to adopt *in situ* synthetic methods to construct 2D/2D MOF/g-C₃N₄ heterojunctions by controllably growing 2D MOFs on CN nanosheets.

Among numerous MOFs investigated, Ni₂(OH)₂(BDC) (BDC = benzene-1,4-dicarboxylate, CCDC no. 985792), briefly named NiMOF, is an ideal coupler choice for CN because it has a laminar morphology, Ni clusters as potential catalytic sites, and relatively good electroconductivity allowing for fast electron transfer.^{30–34} Then the main challenge is how to realize the *in situ* growth of ultrathin 2D NiMOF on CN nanosheets. Naturally, it is reasonable to think of pre-modifying the BDC ligands covalently connected with CN to induce the topological structure of NiMOF to form on the CN surface. However, the conjugated structure makes CN lack relevant binding sites for BDC.²⁶ Thus, it is necessary to functionalize CN in advance by introducing oxygen-containing groups, like hydroxyl and carboxylic groups to avoid destroying the conjugated structure.^{2,35} More importantly, during the process of *in situ* growth of NiMOF, thickening should be prevented to obtain a 2D morphology. A lot of the synthetic routes towards 2D MOFs have been studied, like surfactant-induction or physically assisted methods. By contrast, physical methodologies like ultrasonic treatment are more facile and feasible.

Based on the above, in this work we have fabricated ultrathin dimension-matched NiMOF/CN heterojunctions with intimate interfaces by the *in situ* growth of NiMOF on pre-modified CN by OH and 4-aminobenzoic acid (AA) groups, respectively, with ultrasonic assistance.^{36–39} The as-prepared NiMOF/CN heterojunctions exhibited excellent photocatalytic activities for CO₂RR to produce CO and CH₄ as the main products, among which NiMOF/CN-AA notably reaches 18 times the photoactivity of that for bare CN. With the assistance of the surface photovoltage responses, wavelength-dependent photocurrent action spectra, electrochemical impedance spectra, and CO₂ electrochemical reduction, the mechanism is uncovered in detail showing that the favourable charge transport of ultrathin CN and coupled NiMOF resulted from the greatly-enhanced charge separation *via* the excited high-level electron transfer from CN to NiMOF in the resultant intimately-contacted heterojunction due to the induction effect of AA and then to the provided catalytic functions of Ni clusters for CO₂ activation. This work provides a feasible synthetic protocol to fabricate 2D/2D MOF involved heterojunctions with good charge separation for efficient photocatalysis.

Experimental

All the chemicals and reagents were of analytic grade and used without further purification. Deionized water was used throughout the reaction.

Synthesis of materials

g-C₃N₄ nanosheets (CN). CN was prepared by the urea method. Urea was heated with a heating rate of 0.5 °C min^{−1} in a semi-covered ceramic crucible. After the temperature increased to 550 °C, the crucible was heated for 3 hours at constant temperature, and then slowly cooled to room temperature naturally. A light-yellow powder was obtained and then ground into a fine powder.

OH-functionalized CN (CN-OH). CN was hydroxylated by means of acid treatment. 2 g of the as-prepared CN powder was vigorously stirred for 1 hour in 150 mL 0.5 M nitric acid. Then, the suspension was treated under ultrasonication for 15 min for 3 cycles. Subsequently, the process of washing with water and ethanol was repeated for three cycles, and then vacuum drying was carried out at 60 °C.

AA-functionalized CN (CN-AA). The preparation of AA-functionalized CN was synthesized by two steps.³⁷ The first step is the synthesis of the diazonium salt by the following procedure. 7 mmol (280 mg) NaOH and 7 mmol (960 mg) AA were added into 80 mL water. 7.6 mmol (526 mg) NaNO₂ was then added slowly in an ice bath. Subsequently, the solution was added to 6 mL HCl solution (20%, 6.4 M, 19.2 mmol) and stirred for 1 hour. A solution of phenyl carboxylic diazonium salt with a pale-yellow colour was obtained. Then, the solution was filtered and washed three times with deionized water and ethanol. The obtained powder was dried at 60 °C for 12 hours under vacuum. The second step was to add the prepared CN into the deionized water and then to ultrasound for 1 hour. After mixing the as-prepared diazonium salt and the CN suspension solution, the mixture was stirred for 4 hours in an ice bath and then washed with distilled water, ethanol, DMF and acetone in sequence and repeated for three cycles. Finally, the filtered powder was dried at 60 °C for 12 hours under vacuum.

NiMOF/CN-OH and NiMOF/CN-AA. The *x*NiMOF/CN-OH and *x*NiMOF/CN-AA nanocomposites (*x*% is the weight ratio of NiMOF) were prepared under ultrasonic assistance. 0.5 g CN-OH or CN-AA was dispersed completely into a solvent of 30 mL DMF, 2 mL ethanol and 2 mL deionized water under ultrasound. Then, a certain amount of benzene-1,4-dicarboxylic acid (BDC) was dissolved into the mixed solution under continuous ultrasonication. In this process, the BDC ligand could be dispersed onto the surface of CN-OH or CN-AA. Next, equimolar NiCl₂·6H₂O was added, followed by stirring for 10 minutes. Subsequently, triethylamine (TEA) was quickly injected into the solution. The obtained solution was stirred for 5 minutes to obtain a uniform colloidal suspension which was continuously ultrasonicated for 8 h in airtight conditions.

x NiMOF/CN-OH and x NiMOF/CN-AA were produced by centrifuging, washing by ethanol and water, and freeze-drying under vacuum with the assistance of a large amount of water.

Characterization of materials

The X-ray powder diffraction (XRD) patterns of the samples were conducted by using a Bruker D8 advance diffractometer with $\text{CuK}\alpha$ radiation. The ultraviolet-visible diffuse reflectance spectra (UV-Vis DRS) were tested on a Model Shimadzu UV2550 spectrophotometer with BaSO_4 as the reference. The Fourier-transform infrared (FT-IR) spectra were recorded with a Bruker Equinox 55 spectrometer with KBr as the diluent. The thickness of the samples was analysed by atomic force microscopy (AFM) on a multimode nanoscope VIII instrument (Bruker) with mica as the base. Transmission electron microscopy (TEM) images and data documented the morphology of the samples by using a FEI Tecnai G2 S-Twin instrument with a 200 kV acceleration voltage. The surface photovoltage spectroscopy (SPS) measurements for the samples were implemented on a home-built apparatus, equipped with a lock-in amplifier (SR830, USA) synchronized with a light chopper (SR540, USA). The electron paramagnetic resonance (EPR) measurements were conducted by a Bruker EMX plus model spectrometer operating at the X-band frequency. The X-ray photoelectron spectroscopy (XPS) was carried out with Kratos-AXIS ULTRA DLD, in which Al (Mono) was the X-ray source. CO_2 TPD (temperature-programmed desorption) was tested by the Chemisorption Analyzer (Tp 5080 Chemisorb) with a TCD detector.

Hydroxyl radical measurements

The detection of hydroxyl radicals ($\cdot\text{OH}$) was carried out by a coumarin fluorescence method. A 50 mL coumarin solution with a concentration of 1×10^{-3} M was firstly prepared. 0.05 g catalyst was added into the coumarin solution and stirred for 30 minutes, which ensured the realization of the adsorption-desorption equilibrium. Upon irradiation for 1 h, the suspension was centrifuged and then transferred into a Pyrex glass cell. The fluorescence data was recorded by a PerkinElmer LS55 spectrofluorometer. A 420 nm light filter was utilized to cut off UV light.

Photoelectrochemical and electrochemical measurements

A traditional three-electrode system was utilized to measure the photoelectrochemical (PEC) and electrochemical (EC) reduction, except that the working electrode of the prepared sample, a platinum plate (99.9%) and a saturated KCl Ag/AgCl electrode were used as the counter electrode and reference electrode, respectively. During the experiments, 0.5 M Na_2SO_4 solution was used as the electrolyte, and high purity nitrogen gas was bubbled through. PEC experiments were performed in a quartz cell using a 300 W xenon lamp with a cut-off filter ($\lambda > 420$ nm) as the illumination source. The PEC and EC performance of a series of catalysts was tested on an IVIUM V13806 electrochemical workstation.

Photocatalytic activity evaluation for CO_2 conversion

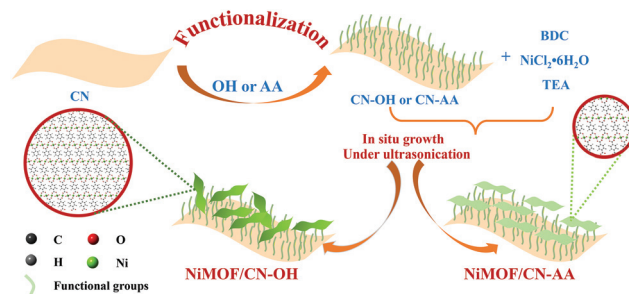
The photocatalytic device contains a cylindrical steel reactor (100 mL volume and 3.5 cm^2 area), a light source (300 W xenon arc lamp), and a gas system (high pure CO_2). Then, the reactor was added by a suspension solution of 0.1 g catalyst and 5 mL water. CO_2 gas was passed through the water and then into the reaction setup. After reaching ambient pressure, the photocatalyst was equilibrated in the $\text{CO}_2/\text{H}_2\text{O}$ system for 1 hour, followed by irradiation for 4 h. During irradiation, 250 μL of the gas produced was taken from the reaction cell at given time intervals for CO and CH_4 concentration analysis using a gas chromatograph (GC-7920 with TCD, Au Light, Beijing) and for O_2 concentration analysis using a gas chromatograph (GC-7900 with TCD, Perfect Light, Beijing).

Results and discussion

Structural characterization

As depicted in Scheme 1, the preparation of NiMOF/CN heterojunction photocatalysts contained two steps. The functionalized CN nanosheets, CN-OH and CN-AA, were firstly obtained under nitric acid and AA treatment, respectively. Next, the NiMOF nanosheets were *in situ* grown on the surface of CN-OH or CN-AA after adding the BDC ligand and $\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$ in sequence under ultrasonication. It should be pointed out that the functionalization treatments with OH or AA to CN rendered different structures and morphologies of the NiMOF grown on CN *in situ*.

As shown in Fig. 1a, the XRD patterns showed that both 4NiMOF/CN-OH and 4NiMOF/CN-AA showed the characteristic peaks of CN located at 13.1° (100) and 27.3° (002), indicating that the structure of CN was preserved during functionalization treatment and *in situ* growth of NiMOF (Fig. 1a).⁴⁰ It was also observed in 4NiMOF/CN-OH and 4NiMOF/CN-AA that tiny peaks at 8.9° , 14.1° , 15.8° , and 17.8° respectively reflected the (200), (001), (201) and (-201) planes of NiMOF (Fig. S1†).⁴¹ The weak XRD signals of NiMOF resulted from the low weight ratio. The coexistence of the characteristic peaks respectively assigned to CN and NiMOF clearly confirmed the successful combination of the two components. Noteworthily, the XRD peak at 8.9° of 4NiMOF/CN-AA was obviously weaker than that



Scheme 1 A schematic illustration of the formation of NiMOF/CN-OH and NiMOF/CN-AA heterojunctions.



Fig. 1 XRD patterns (a) and N₂ sorption isotherm curves (b) from CN, 4NiMOF/CN-OH and 4NiMOF/CN-AA. AFM images of 4NiMOF/CN-OH (c) and 4NiMOF/CN-AA (d).

of 4NiMOF/CN-OH, which implied the thinner morphology and better dispersibility of the NiMOF nanosheets in 4NiMOF/CN-AA.^{42,43}

Given the porous structure of NiMOF, the N₂ physisorption properties were investigated, as shown in Fig. 1b. According to the classification by IUPAC, the N₂ isotherm of 4NiMOF/CN-OH and 4NiMOF/CN-AA were identified as type-II curves with H4-type hysteresis loops, simultaneously possessing the micropores in the NiMOF frameworks and the mesopores originating from the slit-like pores formed by stacking of nanosheets.⁴⁴ The as-measured Brunauer-Emmett-Teller (BET) surface areas of 4NiMOF/CN-OH and 4NiMOF/CN-AA were respectively 40.16 and 52.58 m² g⁻¹, much higher than the 35.14 m² g⁻¹ of CN. These results on the N₂ physisorption properties, along with the above XRD patterns, indicated that the microporous NiMOF is successfully introduced to construct nanocomposites with functionalized CN. In these nanocomposites, CN possessed UV-vis absorption with edges at 460 nm (Fig. S2†), while the OH and AA functionalization and NiMOF coupling did not change the light absorption range. This was further verified by the UV-vis DRS spectrum of the single NiMOF nanosheets as fabricated by the controlled experiment where an indiscernible absorption peak appeared in the visible region ($\lambda > 420$ nm). The band gap energies (E_g) of CN and NiMOF were accordingly calculated to be 2.7 eV and 3.6 eV, respectively. Thus, under visible light CN would act as the light absorber and photogenerated charge producer, and then transfer to the NiMOF nanosheets which could play a significant role in accepting electrons to inhibit charge recombination and then execute catalytic behaviour. To better achieve this process, the NiMOF nanosheets should display good dispersibility and a thinner morphology, which is beneficial for charge transfer among interfaces and NiMOF inner structures.

AFM images depicted the 2D layer-like morphologies of the CN and functionalized CN with lateral sizes from 0.5 to 3 μ m and with corresponding heights of uniformly 4.6 nm (Fig. S3†). After growing NiMOF, the height of 4NiMOF/CN-AA reached about 8.6 nm, with a thickness 4.0 nm thicker than that of CN-AA and a thickness 2.0 nm thinner than that of 4NiMOF/CN-OH. This result illustrated that 2D NiMOF nanosheets grew on the surfaces of functionalized CN nanosheets. In other words, 4NiMOF/CN-OH and 4NiMOF/CN-AA were classified to the dimension-matched heterojunctions. Moreover, given that the cell height of NiMOF is ~ 1.0 nm,³⁰ the NiMOF in the 4NiMOF/CN-OH and 4NiMOF/CN-AA was estimated as six and four layers, respectively. Integrating the layer numbers and weight ratios, it could be inferred that AA functionalization benefited the uniform dispersion of NiMOF on the surface of CN, which was consistent with the above-mentioned result on the XRD patterns. Predictably, the thinner NiMOF nanosheet would be favourable for charge separation and transfer.²⁷

Electron microscope images revealed finer morphological features within the micro-nano scales. In the TEM images, all the prepared samples featured 2D sink-like structures, in which the edges were slightly curled (Fig. 2 and S4†). For 4NiMOF/CN-OH and 4NiMOF/CN-AA, it was hard to obviously observe the existence of bulky NiMOF, which indicated good dispersibility, without obvious aggregation, of NiMOF on the

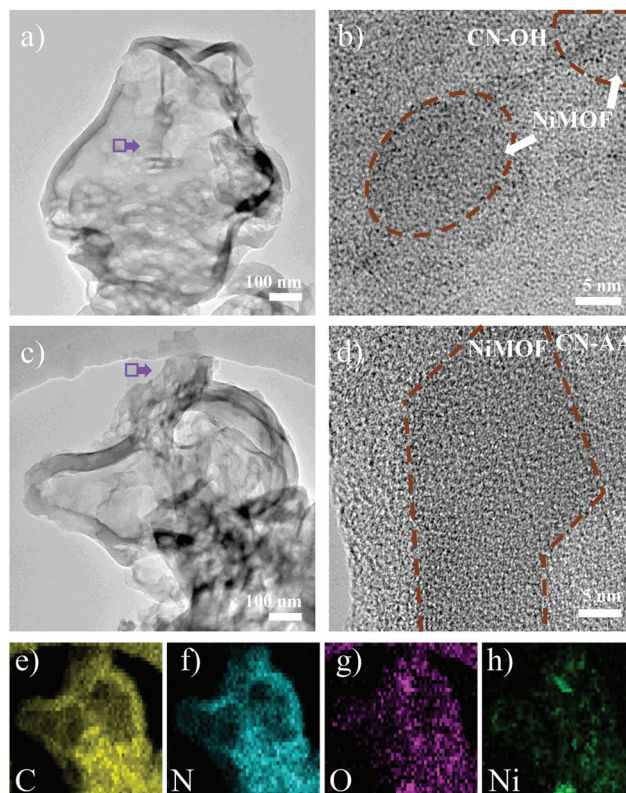


Fig. 2 TEM (a) and HRTEM (b) images of 4NiMOF/CN-OH. TEM (c), HRTEM (d) and elemental mapping images (e–h) of 4NiMOF/CN-AA.

surface of CN at the sub-micrometre scale. Noteworthy, NiMOF was clearly observed in the HRTEM images of the 4NiMOF/CN-OH and 4NiMOF/CN-AA heterojunctions. Fig. 2b presented the nanoscale morphology of 4NiMOF/CN-OH, in which NiMOF nanosheets tightly adhered to the surface of CN-OH. Comparatively, 4NiMOF/CN-AA presented better dispersibility with a larger size and more uniform grayscale, both of which implied a higher dispersion of NiMOF on the surface of CN-AA (Fig. 2d). Predictably, the better dispersibility for 4NiMOF/CN-AA would be helpful for charge transfer from the interface to the whole surface and provided more accessible active sites for the dimension-matched heterojunction photocatalysts.^{27,45} In addition, elemental mapping analysis further proved that the 4NiMOF/CN-AA sample consisted of C, N, O and Ni elements (Fig. 2e–h). The dispersed Ni element confirmed that the NiMOF was uniformly dispersed on the CN surface. All the above information on morphology indicated the successful preparation of the dimension-matched heterojunctions by *in situ* growth of well-dispersed NiMOF onto the functionalized CN.

The above information on morphology proved the successful preparation of the heterojunctions *via in situ* growth of well-dispersed NiMOF nanosheets onto the functionalized CN. Besides, interfacial connection is another significant consideration to influence the efficiency of charge separation and transfer in the dimension-matched heterojunctions. Therefore, the FT-IR and XPS spectra of the prepared samples were tested to finely analyse the interfacial connection between CN and NiMOF. Fig. 3a lists the FT-IR spectra of CN, CN-OH, CN-AA, 4NiMOF/CN-OH and 4NiMOF/CN-AA. All these samples exhibited similar profiles because of the low proportion of NiMOF and high coincidence of the characteristic peaks between CN and NiMOF. Still, valuable detailed information could be

obtained through meticulous analysis. It could be observed in the FT-IR spectrum of CN-OH that the peak at 3040 cm^{-1} obviously strengthened and red-shifted after the hydroxylation, which demonstrated the successful introduction of the surface hydroxylic group on the CN.⁴⁶ For CN-AA, two observations certified the success of AA functionalization. The first one is the red-shift of the peak at 1628 cm^{-1} , which represents introduction of the carboxyl groups.^{35,37} Secondly, two peaks at 2874 and 2890 cm^{-1} were clearly observed, which were attributed to the stretching vibration of the C–H bond from the 1,4-substituted benzene of AA.⁴⁷ In addition, the FT-IR spectra further revealed the connection approach between NiMOF and CN in both 4NiMOF/CN-OH and 4NiMOF/CN-AA. During the process of NiMOF growth, the BDC ligand and Ni^{2+} ion were introduced in sequential order, so it could be inferred that the BDC ligand is preferably connected with the functionalized CN. If not, the broad peak of the surface hydroxylic groups in the range of $3000\text{--}3200\text{ cm}^{-1}$ would obviously shift to 3600 cm^{-1} and be sharper under the coordination between Ni and CN-OH.⁴⁸ However, a slight blue shift occurred and there was no sharpness observed due to the H-bond between the carboxylic group of the BDC ligand and the functional group on CN. This was also verified by the XPS spectra. The N 1s peaks of CN were all shifted to a lower binding energy, indicating that CN related to the NiMOF.² The Ni 2p spectra of 4NiMOF/CN-OH and 4NiMOF/CN-AA possessed indiscernible alternation compared to that of NiMOF fabricated as reference (Fig. 3c). This demonstrated that there were few Ni ions coordinating with CN-OH or CN-AA. According to the above, it was concluded that the functionalized CN and NiMOF were chemically connected by the CN-OH(AA)–BDC–Ni mode.

Based on the above results, dimension-matched NiMOF/CN-based heterojunctions with well-dispersed NiMOF and intimate CN-OH(AA)–BDC–Ni interfaces were synthesized by *in situ* growth of NiMOF onto the functionalized CN. The O atom introduced by functionalization provided more binding sites for the BDC ligand, which is beneficial to disperse the 2D NiMOF nanosheets. Since the AA groups with carboxylic groups have more O atoms than the OH groups, more BDC ligands could be bound on the surface of CN-AA, which renders the formation of NiMOF with better dispersibility and thinner thickness.

Photogenerated charge separation and transfer

Photophysical testing with steady-state surface photovoltage spectroscopy (SS-SPS) is an eminent strategy to reflect the properties of photogenerated charge carriers and was applied to examine the as-synthesized samples. Generally, a stronger SS-SPS response reflects a more efficient charge separation. Among the NiMOF/CN-OH heterojunctions with different amounts of NiMOF, 4NiMOF/CN-OH demonstrated the strongest SS-SPS signal and thereby the most efficient charge separation (Fig. S5†). With a lower proportion of NiMOF, the quantity of charge transfer will be limited, while with a higher proportion, the quality will be negatively influenced by aggregation of NiMOF. With the same loading amount of NiMOF,

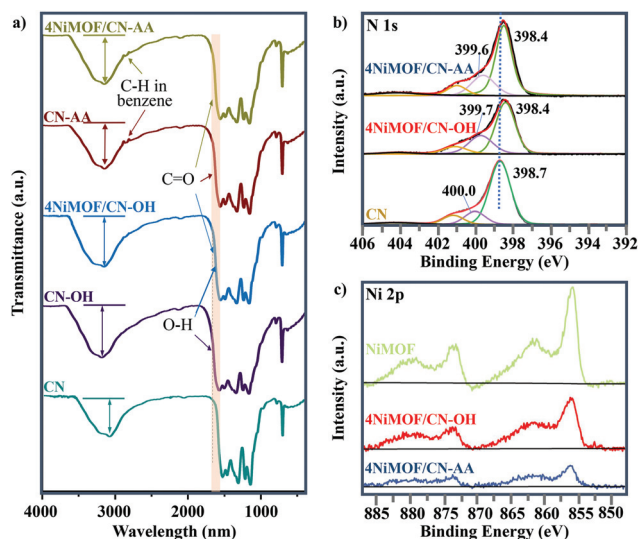


Fig. 3 (a) FT-IR spectra of CN, CN-OH, CN-AA, 4NiMOF/CN-OH and 4NiMOF/CN-AA. (b) N 1s XPS spectra of CN, 4NiMOF/CN-OH and 4NiMOF/CN-AA. (c) Ni 2p XPS spectra of NiMOF, 4NiMOF/CN-OH and 4NiMOF/CN-AA.

4NiMOF/CN-AA showed a further strengthened SS-SPS response in comparison to that for 4NiMOF/CN-OH, indicating an improved charge separation (Fig. 4a). This mainly resulted from more dispersed and thinner NiMOF nanosheets formed by the induction of AA groups as well as the intimate interface, both of which could significantly facilitate the photogenerated charge transfer and separation.

Photochemical analysis toward the amount of produced hydroxyl radicals ($\cdot\text{OH}$) could further verify the separation of photogenerated charge carriers in the photocatalytic process.⁴⁹ A stronger fluorescent signal indicated an increased amount of $\cdot\text{OH}$, and further implied better charge separation. The coumarin fluorescence strategy is a well-accepted method to measure the amount of $\cdot\text{OH}$, where the coumarin reacts with $\cdot\text{OH}$ to produce 7-hydroxyl-coumarin and emits an obvious luminescence at 430–550 nm. As shown in the Fig. 4a inset and S6,[†] the order of the fluorescent intensities of the NiMOF/CN-OH samples was in good agreement with the above photophysical results. In addition, the observed fluorescent intensity order was as follows: CN < 4NiMOF/CN-OH < 4NiMOF/CN-AA, which was also in good accordance with that of the SS-SPS results. Integrating the results of the photophysical and photochemical results, it could be concluded that the photogenerated charge transfer and separation was significantly improved after *in situ* growth of 4% NiMOF onto functionalized CN, especially for CN-AA. Additionally, the transient photocurrent-time (*I*-*t*) curves supported the above results and further presented the stability of the as-prepared photocatalysts (Fig. S7[†]).

Evaluation of photocatalytic activities

The photocatalytic activities for CO₂RR were conducted under visible-light irradiation with a 420 nm cut-off filter. The reduction products, CH₄ and CO, and the oxidation product of O₂ were detected. As shown in Fig. 4b and S8,[†] the photoactivities on yielding CO, CH₄ and O₂ had low activities for CN, NiMOF and functionalized CN, but achieved a significant enhancement when the NiMOF was grown on CN-OH and CN-AA. Among the serial heterojunctions with a proper amount of NiMOF, 4NiMOF/CN-OH displayed the highest photocatalytic performance due to the improvement on charge

separation and transfer (Fig. S8[†]). Impressively, 4NiMOF/CN-AA exhibited the best photoactivity with 11-fold and nearly 2-fold enhancements, respectively, compared with CN and 4NiMOF/CN-OH. For all the samples, the order of the photoactivities was consistent with those of the photophysical and photochemical results, indicating that the enhanced photoactivities would be tightly associated with better NiMOF dispersibility and improved charge separation.

Compared with similar work reported (Table S1[†]),^{23–26} the photocatalytic activities of 4NiMOF/CN-AA in our study exhibited an excellent performance for the aqueous CO₂RR among the MOF/g-C₃N₄-based heterojunction photocatalysts. For 4NiMOF/CN-AA, the amounts of evolved CH₄ and CO linearly increased with increasing the irradiation time, and there was nearly no activity loss after five-cycle reactions (Fig. S9[†]). These results reflected the stable nature of the as-fabricated heterojunction photocatalysts.

Discussion of the mechanism

As described above, the photocatalytic activities of CN can be greatly improved by *in situ* growth of NiMOF with the induction of OH or AA to obtain a series of NiMOF/CN heterojunction photocatalysts. This has been proved as the promoted charge transfer and separation caused by the matched 2D/2D structures and efficient interfacial connection between functionalized CN and NiMOF. This is further supported by electrochemical impedance spectra of three typical samples (Fig. 5a). 4NiMOF/CN-OH and 4NiMOF/CN-AA exhibited an obviously decreased capacitive arc radius compared with CN, implying that the charge recombination was inhibited by transferring the photogenerated electrons from CN to NiMOF. Moreover, 4NiMOF/CN-AA showed the smallest arc radius, which demon-



Fig. 4 SS-SPS responses and FS spectra (inset) related to the amount of produced $\cdot\text{OH}$ (a) and the photocatalytic activities for CO₂ conversion under visible-light irradiation for 4 hours (b) of CN, 4NiMOF/CN-OH and 4NiMOF/CN-AA.



Fig. 5 Electrochemical impedance spectra (a), normalized photocurrent action spectra under different monochromatic light irradiation (b), EPR spectra with UV-vis light irradiation at 98 K (c) and electrochemical reduction curves in the CO₂-bubbled system (d) of CN, 4NiMOF/CN-OH and 4NiMOF/CN-AA.

strated that the well-dispersed NiMOF on CN-AA had a better charge transfer capacity.

The charge separation might be attributed to the photoelectron transfer from CN to the as-grown NiMOF acting as an electron-accepting platform in the NiMOF/CN heterojunctions *via* the well-modulated intimate interfaces. To verify the possibility for CO₂ photoreduction, Mott-Schottky spectra of CN and NiMOF were carried out at the frequency of 500, 1000 and 1500 Hz (Fig. S10†). The resulting positive slopes of the $C^2 - E$ values (*vs.* the applied potentials) were consistent with that of the typical n-type semiconductors, and the intersection at -1.46 eV for CN and -1.53 eV for NiMOF *vs.* Ag/AgCl (*i.e.* -1.26 eV for CN and -1.33 eV for NiMOF *vs.* NHE) was determined as the flat band position. Given that generally for the n-type semiconductor the bottom of the conduction band (CB) for CN or lowest unoccupied molecular orbital (LUMO) for NiMOF is approximately equal to the flat-band potential, the CB of CN and the LUMO of NiMOF were determined as -1.26 eV and -1.33 eV *vs.* NHE, respectively. With the bandgap energy provided by the Tauc plot (Fig. S2†), the valence band (VB) of CN was calculated as $+1.49$ eV, and the highest occupied molecular orbital (HOMO) of NiMOF was $+2.27$ eV. Given the more negative potential of CN and NiMOF in the heterojunctions than the reduction potential of CO₂ reduction, it is theoretically feasible for the photocatalytic CO₂ reduction to CO and CH₄. Because the energy position of the LUMO of NiMOF was slightly more negative than the CB of CN, it was inferred that the electrons could transfer by means of high-level electron transfer.

In order to verify this hypothesis, monochromatic photocurrent action spectra were thus examined. As shown in Fig. 5b, the photocurrent density of 4NiMOF/CN-OH and 4NiMOF/CN-AA gradually increased as the excitation wavelength decreased from 460 to 400 nm with steps of 10 nm. Noticeably, it began to sharply increase under 440 nm excitation, which was the threshold wavelength to facilitate the transfer of visible-light excited appropriate-energy electrons from CN to NiMOF. The difference between the absorption edge at 460 nm and the threshold wavelength at 440 nm verified that the photogenerated electrons should be excited to a higher energy level of the CB of CN so as to realize the photoelectron transfer. Subsequently, a low-temperature EPR technique was employed to explore the process of photoelectron transfer from CN to NiMOF (Fig. 5c). The weak EPR signal of CN at $g = 2.001$ was assigned to the unpaired electrons on the conjugated CN aromatic rings.²³ After growing NiMOF, this signal becomes stronger, especially for 4NiMOF/CN-AA. This indicated that the introduction of NiMOF efficiently inhibited the charge recombination and generated more free electrons. This corresponded well with the results of the fluorescent spectra. It was remarkable that the unpaired electron in Ni(II) was directly observed for 4NiMOF/CN-OH and 4NiMOF/CN-AA. The signals at $g = 2.285$ in the EPR spectra were attributed to the unpaired electron in the $3d_{x^2-y^2}$ orbital of Ni.⁴³ However, Ni(II) complexes were extremely thermally unstable and dispro-

portionately formed Ni(0) and Ni(II) species. The formation of Ni(II) species was due to the partially reduced central Ni(II) in NiMOF by the photoelectrons transferred from CN. In addition, 4NiMOF/CN-AA possessed a much stronger signal at $g = 2.285$ in comparison to 4NiMOF/CN-OH, which confirmed the better capacity of charge transfer and separation for 4NiMOF/CN-AA.

Electrochemical reduction measurements were also performed to explore the mechanism of CO₂ conversion (Fig. 5d and S11†). In both of the N₂ and CO₂ bubbled systems, the potential values had the order 4NiMOF/CN-AA < 4NiMOF/CN-OH < CN. This result confirmed that both 4NiMOF/CN-OH and 4NiMOF/CN-AA were beneficial to improve the photocatalytic activity for the CO₂ reduction. It also demonstrated that the NiMOF possessed catalytic potential. By comparison, the samples in the CO₂ bubbled system had lower potential values than those in the N₂ bubbled system, which illustrated that the reduction behaviour of CO₂ was also much preferred compared with H₂O reduction. This indicated that the coupled NiMOF could provide good catalytic functions for CO₂ reduction, which was further supported by the promoted adsorption of CO₂ based on the CO₂ TPD spectra (Fig. S12†). Thus, it was deduced that the specific central Ni(II) species in the NiMOF/CN heterojunctions were the catalytic sites for activating CO₂ and hence facilitated the reduction by the photoelectrons.

Based on all the above results, a schematic process has been proposed about high-energy-level electron transfer over dimension-matched intimately contacted 4NiMOF/CN-OH and 4NiMOF/CN-AA heterojunction photocatalysts (Fig. 6a). Under visible-light irradiation, the photogenerated electrons on CN were excited to the energy level higher than the CB, and then transferred to the lowest unoccupied molecular orbital (LUMO) of NiMOF *via* the CN-OH/AA-BDC-Ni interface. Subsequently, the Ni(II) species could accept the high-energy-level photoelectrons from the BDC ligand and then catalyse the CO₂RR. In this case, the spatially separated photogenerated holes on CN and electrons on NiMOF would possess sufficient thermodynamic energy to induce redox reactions, respectively, leading to the promoted charge separation and hence to the improved photocatalytic activities. Moreover, the photocatalytic activities for CO₂ conversion could be further improved under the irradiation of UV-visible light. As shown in Fig. 6b, the photocatalytic activity on the total products of CO and CH₄ was $54.5 \mu\text{mol g}^{-1}$ for 4NiMOF/CN-AA, an 18-fold improvement compared with that of CN. Compared with visible light irradiation, the activities under UV-vis irradiation possessed a 6-fold improvement for both 4NiMOF/CN-OH and 4NiMOF/CN-AA. This large improvement further confirmed the as-proposed mechanism on high-electron-level electron transfer in that the electrons were excited to higher energy levels under UV irradiation and were more beneficial to electron transfer.

To clearly certify the influence on the photoactivities by dimension matching, a control experiment was carried out in which a reference sample was prepared, namely 4B-NiMOF/

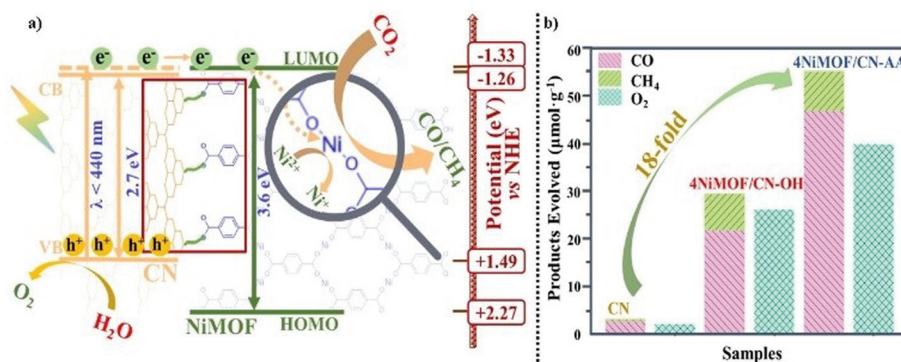


Fig. 6 (a) A schematic diagram of the photogenerated charge transfer process and the induced photochemical reaction in the resultant NiMOF/functionalized CN nanocomposite. (b) Photocatalytic activities for CO₂ conversion under UV-vis irradiation for 4 hours of CN, 4NiMOF/CN-OH and 4NiMOF/CN-AA.

CN-AA, by *in situ* growing 4% bulky NiMOF onto CN-AA under hydrothermal conditions. Besides, 4CoMOF/CN-AA and 4ZnMOF/CN-AA were also prepared by substituting Ni²⁺ ions with Co²⁺ or Zn²⁺ ions during the process of *in situ* growth. These reference samples sharply decreased photoactivity for CO₂ conversion compared with 4NiMOF/CN-AA (Fig. S13†). The SS-SPS spectra certified the weak charge separation of 4B-NiMOF/CN-AA due to the thicker morphology and bad dispersibility (Fig. S5†). The electrochemical reduction spectra revealed the decreased catalytic capacities for 4CoMOF/CN-AA and 4ZnMOF/CN-AA due to their increased potential values as seen in Fig. S14.† These results further confirmed that the excellent photocatalytic properties on CO₂RR were due to the dimension-matched structures, well-modulated interfaces and introduced Ni active centres.

Conclusions

In summary, dimension-matched NiMOF/CN-OH and NiMOF/CN-AA heterojunctions with CN-OH/AA-BDC-Ni interfacial connection were successfully prepared by means of *in situ* growth with ultrasonic assistance. These nanocomposites exhibited excellent photocatalytic activities for the CO₂RR to produce CO and CH₄. It was clear that the exceptional photocatalytic activities mainly resulted from the favourable charge transport within the ultrathin CN and coupled NiMOF nanosheets and from the greatly-enhanced charge separation *via* excited high-level electron transfer from CN to NiMOF in the resultant intimately contacted heterojunction caused by surface oxygen-containing functionalization, and also from the provided catalytic functionality of the Ni clusters for CO₂ activation. In particular, all the structural factors, such as the *in situ* growth approach, ultrasonic assistance, good dispersibility, well-modulated interfaces and provided active sites, are quite important for an efficient photocatalytic CO₂RR. This work offers unique insight into CN-based heterojunction systems, which will be essential for the development of CO₂ photocatalysts.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors are grateful for support from the National Natural Science Foundation of China (U1805255), and Scientific Research Project of East University of Heilongjiang (HDFHX180106).

Notes and references

- 1 Z. Sun, N. Talreja, H. Tao, J. Texter, M. Muhler, J. Strunk and J. Chen, *Angew. Chem., Int. Ed.*, 2018, **57**, 7610.
- 2 J. Bian, J. Feng, Z. Zhang, Z. Li, Y. Zhang, Y. Liu, S. Ali, Y. Qu, L. Bai, J. Xie, D. Tang, X. Li, F. Bai, J. Tang and L. Jing, *Angew. Chem., Int. Ed.*, 2019, **58**, 10873.
- 3 X. Li, Y. Sun, J. Xu, Y. Shao, J. Wu, X. Xu, Y. Pan, H. Ju, J. Zhu and Y. Xie, *Nat. Energy*, 2019, **4**, 690.
- 4 Z. Guo, G. Chen, C. Cometto, B. Ma, H. Zhao, T. Groizard, L. Chen, H. Fan, W. Man, S. Yiu, K. Lau, T. Lar and M. Rober, *Nat. Catal.*, 2019, **2**, 830.
- 5 J. Fu, J. Yu, C. Jiang and B. Cheng, *Adv. Energy Mater.*, 2018, **8**, 1701503.
- 6 C. Tan, X. Cao, X. Wu, Q. He, J. Yang, X. Zhang, J. Chen, W. Zhao, S. Han, G. Nam, M. Sindoro and H. Zhang, *Chem. Rev.*, 2017, **117**, 6225.
- 7 W. Zhang, A. Mohamed and W. Ong, *Angew. Chem., Int. Ed.*, 2020, DOI: 10.1002/ange.201914925.
- 8 Y. Xiao, G. Tian, W. Li, Y. Xie, B. Jiang, C. Tian, D. Zhao and H. Fu, *J. Am. Chem. Soc.*, 2019, **141**, 2508.
- 9 S. Sun and S. Liang, *Nanoscale*, 2017, **9**, 10544.
- 10 X. Zhang, X. Zhang, J. Li, J. Sun, J. Bian, J. Wang, Y. Qu, R. Yan, C. Qin and L. Jing, *Appl. Catal., B*, 2018, **237**, 50.
- 11 J. Wang, C. Qin, H. Wang, M. Chu, A. Zada, X. Zhang, J. Li, F. Raziq, Y. Qu and L. Jing, *Appl. Catal., B*, 2018, **221**, 459.

- 12 W. Ong, L. Tan, Y. Ng, S. Yong and S. Chai, *Chem. Rev.*, 2016, **116**, 7159.
- 13 H. Zhang, X. Gu, P. Liu, J. Song, J. Cheng and H. Su, *J. Mater. Chem. A*, 2017, **5**, 2288.
- 14 L. Kong, Y. Dong, P. Jiang, G. Wang, H. Zhang and N. Zhao, *J. Mater. Chem. A*, 2016, **4**, 9998.
- 15 M. Chu, K. Hu, J. Wang, Y. Liu, S. Ali, C. Qin and L. Jing, *Appl. Catal., B*, 2019, **243**, 57.
- 16 Q. Wang and D. Astruc, *Chem. Rev.*, 2020, **120**, 1438.
- 17 W. Jiang, N. Zhang, X. Zheng, C. Gao, L. Wang, Q. Zhang, Y. Zhu and Y. Xiong, *J. Am. Chem. Soc.*, 2019, **141**, 10924.
- 18 S. Zhong, Y. Xi, Q. Chen, J. Chen and S. Bai, *Nanoscale*, 2020, **12**, 5764.
- 19 J. Duan, Y. Li, Y. Pan, N. Behera and W. Jin, *Coord. Chem. Rev.*, 2019, **395**, 25.
- 20 J. Wang, C. Wang and W. Lin, *ACS Catal.*, 2012, **2**, 2630.
- 21 Q. Yang, Q. Xu and H. Jiang, *Chem. Soc. Rev.*, 2017, **46**, 4774.
- 22 H. Wang, X. Yuan, Y. Wu, G. Zeng, X. Chen, L. Leng and H. Li, *Appl. Catal., B*, 2015, **174–175**, 445.
- 23 L. Shi, T. Wang, H. Zhang, K. Chang and J. Ye, *Adv. Funct. Mater.*, 2015, **25**, 5360.
- 24 S. Liu, F. Chen, S. Li, X. Peng and Y. Xiong, *Appl. Catal., B*, 2017, **211**, 1.
- 25 G. Xu, H. Zhang, J. Wei, H. Zhang, X. Wu, Y. Li, C. Li, J. Zhang and J. Ye, *ACS Nano*, 2018, **12**, 5333.
- 26 Y. Wang, L. Guo, Y. Zeng, H. Guo, S. Wan, M. Ou, S. Zhang and Q. Zhong, *ACS Appl. Mater. Interfaces*, 2019, **11**, 30673.
- 27 M. Zhao, Y. Huang, Y. Peng, Z. Huang, Q. Ma and H. Zhang, *Chem. Soc. Rev.*, 2018, **47**, 6267.
- 28 R. Li, W. Zhang and K. Zhou, *Adv. Mater.*, 2018, **30**, 1705512.
- 29 A. Dhakshinamoorthy, A. Asiri and H. Garcia, *Adv. Mater.*, 2019, **31**, 1900617.
- 30 S. Zhao, Y. Wang, J. Dong, C. He, H. Yin, P. An, K. Zhao, X. Zhang, C. Gao, L. Zhang, J. Lv, J. Wang, J. Zhang, A. Khattak, N. Khan, Z. Wei, J. Zhang, S. Liu, H. Zhao and Z. Tang, *Nat. Energy*, 2016, **1**, 16184.
- 31 J. Duan, S. Chen and C. Zhao, *Nat. Commun.*, 2017, **8**, 1.
- 32 Y. Jiao, J. Pei, C. Yan, D. Chen, Y. Hu and G. Chen, *J. Mater. Chem. A*, 2016, **4**, 13344.
- 33 B. Han, X. Ou, Z. Deng, Y. Song, C. Tian, H. Deng, Y. Xu and Z. Lin, *Angew. Chem., Int. Ed.*, 2018, **57**, 16811.
- 34 A. Cao, L. Zhang, Y. Wang, H. Zhao, H. Deng, X. Liu, Z. Lin, X. Su and F. Yue, *ACS Sustainable Chem. Eng.*, 2019, **7**, 2492.
- 35 G. Zhou, M. Wu, Q. Xing, F. Li, H. Liu, X. Luo, J. Zou, J. Luo and A. Zhang, *Appl. Catal., B*, 2018, **220**, 607.
- 36 H. Wang, C. Qian, J. Liu, Y. Zeng, D. Wang, W. Zhou, L. Gu, H. Wu, G. Liu and Y. Zhao, *J. Am. Chem. Soc.*, 2020, **142**, 4862.
- 37 M. Jahan, Q. Bao, J. Yang and K. Loh, *J. Am. Chem. Soc.*, 2010, **132**, 14487.
- 38 J. Xu, S. He, H. Zhang, J. Huang, H. Lin, X. Wang and J. Long, *J. Mater. Chem. A*, 2015, **3**, 24261.
- 39 B. Li, L. Sun, J. Bian, N. Sun, J. Sun, L. Chen, Z. Li and L. Jing, *Appl. Catal., B*, 2020, **270**, 118849.
- 40 S. Yan, Z. Li and Z. Zou, *Langmuir*, 2009, **25**, 10397.
- 41 A. Mesbah, P. Rabu, R. Sibille, S. Lebegue, T. Mazet, B. Malaman and M. Francois, *Inorg. Chem.*, 2014, **53**, 872.
- 42 Z. Liang, C. Qu, D. Xia, R. Zou and Q. Xu, *Angew. Chem., Int. Ed.*, 2018, **57**, 9604.
- 43 H. Yang, S. Hung, S. Liu, K. Yuan, S. Miao, L. Zhang, X. Huang, H. Wang, W. Cai, R. Chen, J. Gao, X. Yang, W. Chen, Y. Huang, H. Chen, C. Li, T. Zhang and B. Liu, *Nat. Energy*, 2018, **3**, 140.
- 44 J. Duan, S. Chen and C. Zhao, *Nat. Commun.*, 2017, **8**, 15341.
- 45 H. Zhang, *Adv. Mater.*, 2015, **27**, 7372.
- 46 H. Zhao, S. Wang, F. He, J. Zhang, L. Chen, P. Dong, Z. Tai, Y. Wang, H. Gao and C. Zhao, *Carbon*, 2019, **150**, 340.
- 47 A. Kumar, J. Kudva, S. Kumar, U. Vishwanatha, V. Kumar and D. Naral, *J. Mol. Struct.*, 2018, **1167**, 142.
- 48 S. Halder, J. Mondal, J. Ortega-Castro, A. Frontera and P. Roy, *Dalton Trans.*, 2017, **46**, 1943.
- 49 Y. Nakabayashi and Y. Nosaka, *J. Phys. Chem. C*, 2013, **117**, 23832.