


 Cite this: *RSC Adv.*, 2020, **10**, 39080

# NH<sub>2</sub>-MIL-88B (Fe<sub>α</sub>In<sub>1-α</sub>) mixed-MOFs designed for enhancing photocatalytic Cr(VI) reduction and tetracycline elimination†

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Aiming at solving the issue of wastewater purification, this work synthesized NH<sub>2</sub>-MIL-88B (Fe<sub>α</sub>In<sub>1-α</sub>) photocatalysts by a simple one-pot method, which was employed for photocatalytic reduction of Cr(VI) and oxidation of TC-HCl. Compared with traditional NH<sub>2</sub>-MIL-88B (Fe) photocatalysts, NH<sub>2</sub>-MIL-88B (Fe<sub>0.6</sub>In<sub>0.4</sub>) displayed excellent photocatalytic performance; the photocatalytic redox rate for Cr(VI) and TC-HCl reached 86.83% and 72.05%, respectively. The good photocatalytic performance might be attributed to the metal-to-metal charge transition (MMCT) between Fe–O clusters and In–O clusters formed by doping In(III) into NH<sub>2</sub>-MIL-88B (Fe), which provides effective active sites for the photocatalytic reduction and oxidation routes. Besides, the synergistic effect of the ligand-to-metal charge transition (LMCT) and MMCT expands the separation and transfer of photogenerated carriers and inhibits the recombination of electron–hole pairs, thus effectively improving the photocatalytic performance. Therefore, this work could provide a new method for the construction of mixed metal MOFs for the photocatalytic degradation of pollutants.

 Received 31st August 2020  
 Accepted 18th October 2020

DOI: 10.1039/d0ra07487j

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## Introduction

Heavy metals and residual antibiotics in wastewater have become a serious threat to living creatures and the environment.<sup>1–4</sup> Hexavalent chromium (Cr(VI)), as a heavy metal ion pollutant in wastewater with high toxicity and mobility, has seriously threatened human health and the natural environment.<sup>5,6</sup> If a large amount of Cr(VI) existed in the drinking water system, it would cause damage to the human stomach, kidneys, liver, and retinas and also serious environmental concern.<sup>7,8</sup> Therefore, it is urgent to remove Cr(VI) from wastewater. There are many existing solutions such as chemical precipitation,<sup>9</sup> membrane separation,<sup>10</sup> ion exchange,<sup>11</sup> activated carbon adsorption<sup>12</sup> and photocatalytic reduction.<sup>13–15</sup> Among them, the reduction of Cr(VI) to Cr(III) by semiconductor photocatalysis is a very effective strategy due to the low toxicity of Cr(III).<sup>16–18</sup> Furthermore, tetracycline hydrochloride (TC-HCl), as one of the most common antibiotics, is widely used in various industries such as medicine, agriculture, and animal husbandry. TC-HCl is abundant in the environment due to its stable chemical structure and non-biodegradation, which has already imperiled the ecosystem and human health.<sup>19–21</sup> There are many current

ways to remove TC-HCl, including adsorption,<sup>22</sup> advanced oxidation processes (AOPs),<sup>21</sup> biological degradation,<sup>23</sup> electrolysis,<sup>24</sup> membrane filtration,<sup>25</sup> ion exchange<sup>26</sup> and photocatalytic degradation.<sup>27,28</sup> Photocatalytic degradation is widely used in recent years because of mild reaction conditions, high efficiency, low cost and greenly.<sup>20,29–31</sup>

Metal–organic frameworks (MOFs), as a class of crystalline inorganic–organic hybrid materials, are composed of the interconnection between metal centers and multidentate organic ligands.<sup>32,33</sup> MOFs have been valued by researchers in the field of photocatalysis<sup>34,35</sup> due to the inherent large specific surface area, uniform and adjustable pore structure, and abundant active sites.<sup>36</sup> Under the irradiation of incident light, the photogenerated electrons are excited from the valence band (VB) and transferred to the conduction band (CB) through the ligand-to-metal charge transition (LMCT) process. The separated photogenerated electrons–holes pairs can perform the reduction and oxidation reaction with the reactants adsorbed on the catalyst surface.<sup>37</sup> Traditional MOFs existed some shortcomings, such as small photoresponse range and high electrons–holes recombination rate.<sup>38</sup> It is undoubtedly a good way to change the photoresponse range and photocatalytic performance of MOFs by adjusting the central metal or organic ligands.<sup>39</sup> The strategy of combining two or more transition metals into the same framework to form mixed-metal MOFs has aroused the interest of researchers.<sup>40</sup>

Mixed-metal organic framework (MM-MOFs), containing two or more metal centers, are periodically arranged with a single ligand in the whole framework.<sup>41,42</sup> There is a metal-to-metal

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0ra07487j



charge transition (MMCT) process between multiple metals in MM-MOFs, which facilitates the transfer of photogenerated electrons and inhibits the recombination of electrons–holes pairs. Compared with single-metal MOFs, the synergistic effects between LMCT and MMCT of MM-MOFs can effectively improve the photocatalytic activity in various applications.<sup>43</sup> Also, MM-MOFs can effectively control the electronic structure and bandgap structure while increasing the metal active sites.<sup>44–46</sup> For instance, Vu *et al.* synthesized Fe-Cr-MIL-101 for dye degradation by the hydrothermal method. Mixed-metal MOFs showed better adsorption capacity and higher photocatalytic activity and stability in comparison with Cr-MIL-101.<sup>47</sup> Maite P. *et al.* studied the synthesis of multimetal MOFs doped with Zn(II), Ru(III) and Pd(II) into Cu-MOF and applied to electrochemical reduction of CO<sub>2</sub>.<sup>48</sup> In our previous report, we successfully synthesized NH<sub>2</sub>-MIL-68 (In<sub>α</sub>Fe<sub>1-α</sub>) that displayed good photocatalytic activity.<sup>49</sup> According to reports, NH<sub>2</sub>-MIL-88B (Fe) has a flexible pore structure and a large open structure channel due to its 3D porous structure containing hexagonal channels and bipyramidal cages.<sup>50</sup> The wide and flexible pore structure facilitated other metal ions to smoothly enter the 3D structure to replace the original central metal ions, and can provide adjustment space for structural changes that may be caused by the introduction of other central metal ions, thereby improving related performance.

Based on the above analysis, a novel MM-MOFs material NH<sub>2</sub>-MIL-88B (Fe<sub>α</sub>In<sub>1-α</sub>) was constructed and synthesized *via* the one-pot method directly, which used NH<sub>2</sub>-MIL-88B (Fe) as the structural framework. Under visible light irradiation, the activity of photocatalytic treatment of pollutants in water was tested. The results showed that the efficiency of Cr(VI) reduction and TC-HCl oxidation of NH<sub>2</sub>-MIL-88B (Fe<sub>α</sub>In<sub>1-α</sub>) was 1.7 and 1.6 times that of NH<sub>2</sub>-MIL-88B (Fe). Moreover, a possible photocatalytic mechanism was proposed through characterization and performance and active species testing. This work provides an effective way for the construction of MM-MOFs and the photocatalytic degradation of various pollutants in wastewater.

## Experimental sections

The chemicals and characterization details could be found in ESI.†

### Synthesis of NH<sub>2</sub>-MIL-88B (Fe)

The NH<sub>2</sub>-MIL-88B (Fe) sample was prepared *via* a one-pot method. 2-Aminoterephthalic acid (NH<sub>2</sub>-BDC) (0.54 mmol) and Fe(NO<sub>3</sub>)<sub>3</sub>·9H<sub>2</sub>O (0.48 mmol) were dissolved in DMF (12 mL), added CH<sub>3</sub>CN (12 mL) after mixed well. Then, and the mixture was transferred into a flask, oil bath at 120 °C for 45 min. The suspension was cooled to room temperature naturally and washed with methanol. The sample was dried at 70 °C for 5 h, then NH<sub>2</sub>-MIL-88B (Fe) was gained.

### Synthesis of In-doped NH<sub>2</sub>-MIL-88B (Fe)

In-doped NH<sub>2</sub>-MIL-88B (Fe) was synthesized through the above similar process, except that pure Fe(NO<sub>3</sub>)<sub>3</sub>·9H<sub>2</sub>O was replaced with different ratios Fe(NO<sub>3</sub>)<sub>3</sub>·9H<sub>2</sub>O and In(NO<sub>3</sub>)<sub>3</sub>·xH<sub>2</sub>O. The

products were named as NH<sub>2</sub>-MIL-88B (Fe<sub>α</sub>In<sub>1-α</sub>), and the synthesis diagram was shown in Scheme 1.

### Evaluation of photocatalytic performance

The photocatalytic performance of as-obtained samples was evaluated by removing Cr(VI) and TC-HCl. A certain amount of synthesized NH<sub>2</sub>-MIL-88B (Fe<sub>α</sub>In<sub>1-α</sub>) was added into the Cr(VI) solution and stirring sufficiently. Simultaneously, the pH value of the Cr(VI) solution was adjusted to 2 with HCl solution. Then proceed to photocatalytic degradation. During the experiment, pay attention to the concentration of Cr(VI) in the system. The suspension was stirred for some time under dark conditions. When the concentration of Cr(VI) no longer changed, switch to Xe lamp irradiation. The suspensions taken out regularly are centrifuged and filtered with a 0.22 μm syringe filter. The determination of Cr(VI) was realized by the DPC method with a UV-visible spectrophotometer. The photocatalytic oxidation of TC-HCl is similar to the above process. Where the pH value of the TC-HCl solution was adjusted to 9 with NaOH solution and the determination was realized by the UV-visible spectrophotometer method. The removal adsorption reduction/oxidation efficiency of pollutant is calculated using Lambert Beer's law and the following formula:

$$\text{Efficiency (\%)} = C/C_0 \times 100\% \quad (1)$$

here  $C$  and  $C_0$  are the concentrations of pollutants in the water at  $t$  and  $t_0$ .

## Results and discussions

### Structure and morphology analysis

As revealed in Fig. 1a, the crystalline structure of NH<sub>2</sub>-MIL-88B (Fe) and In-doped NH<sub>2</sub>-MIL-88B (Fe<sub>α</sub>In<sub>1-α</sub>) are confirmed by the XRD pattern. The obvious characteristic peaks of NH<sub>2</sub>-MIL-88B (Fe) located at 9.21°, 10.25°, 13.07°, 16.65°, 18.53°, 20.69°, consistent with the literature reported,<sup>45</sup> verified the successful preparation of NH<sub>2</sub>-MIL-88B (Fe). The doped amounts of In on NH<sub>2</sub>-MIL-88B (Fe) determined by the ICP-AES analysis as shown in Table S1.† Concomitantly, the main characteristic peaks of NH<sub>2</sub>-MIL-88B (Fe<sub>α</sub>In<sub>1-α</sub>) appeared in the same position as NH<sub>2</sub>-MIL-88B, which preliminarily proved that NH<sub>2</sub>-MIL-88B (Fe<sub>α</sub>In<sub>1-α</sub>) was successfully prepared. The as-prepared MM-MOFs mainly exist combination and crystal lattice in the form



Scheme 1 Synthesis diagram of the synthetic processes of NH<sub>2</sub>-MIL-88B (Fe<sub>α</sub>In<sub>1-α</sub>).



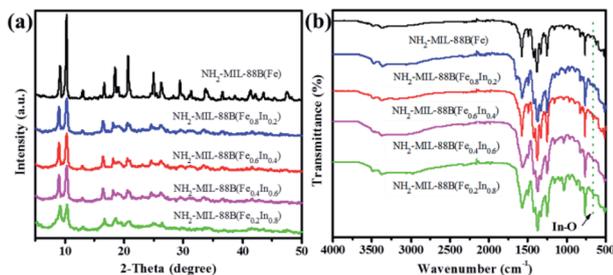


Fig. 1 (a) XRD patterns and (b) FT-IR of  $\text{NH}_2\text{-MIL-88B (Fe)}$  and  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  samples.

of  $\text{NH}_2\text{-MIL-88B (Fe)}$ . As the molar ratio of  $\text{In(III)}$  ions increases, the characteristic peak at  $9.21^\circ$  shifts to  $9.02^\circ$ . The deviation of the characteristic peak might be due to that the ion size of  $\text{In(III)}$  (94.0 pm) in the crystal structure was significantly larger than the ion size of  $\text{Fe(III)}$  (78.5 pm), thereby the size of the trimer composed of  $\text{InO}_6$  octahedron was larger than that of  $\text{FeO}_6$  octahedron in the crystal structure of  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$ . Therefore, the peak corresponding to the characteristic peak in the spectrum could shift to a lower angle due to the increase of the unit cell parameters.<sup>51</sup>

To further determine the chemical structure and investigate the doped of  $\text{In(III)}$  of the prepared samples, the FT-IR spectrum is employed to analyze. As presented in Fig. 1b, the FT-IR spectrum of  $\text{NH}_2\text{-MIL-88B (Fe)}$  is consistent with the previous literature.<sup>52</sup> By comparison, after indium doping, a new characteristic absorption peak generation, which could be attributed to the  $\text{In-O}$  bond appears at  $663\text{ cm}^{-1}$ , indicating that  $\text{In(III)}$  is successfully introduced and the  $\text{In-O}$  bond is formed. Except for the appearance of new characteristic absorption peaks, the characteristic absorption peaks corresponding to other chemical bonds have not been changed, indicating that the introduction of  $\text{In(III)}$  does not affect the chemical structure of  $\text{NH}_2\text{-MIL-88B (Fe)}$ ,  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  was successfully prepared.

Fig. 2a–f displays representative SEM images and length comparison of  $\text{NH}_2\text{-MIL-88B (Fe)}$  and  $\text{In-doped NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$ . As shown in Fig. 2a,  $\text{NH}_2\text{-MIL-88B (Fe)}$  performed a standard spindle with uniform morphology, in which the length is 300–500 nm. Additionally, the spindle is continuously elongated in the longitudinal direction with the increase of  $\text{In(III)}$  ion content as shown in Fig. 2b–f. However, the overall morphology of  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  is unchanged, forming a six-sided double pyramid with a length in the range of 500–1000 nm. The cause of the morphological change may be ascribed to  $\text{In(III)}$  ions replacing  $\text{Fe(III)}$  into the  $\text{NH}_2\text{-MIL-88B (Fe)}$  lattice, which increases the lattice spacing in the crystal.<sup>51</sup> The BET specific surface area test results showed that the BET specific surface areas of  $\text{NH}_2\text{-MIL-88B (Fe}_{0.6}\text{In}_{0.4})$  is  $204\text{ m}^2\text{ g}^{-1}$  less than that of  $\text{NH}_2\text{-MIL-88B (Fe)}$ , as Table S2† in details. The pore structure of  $\text{NH}_2\text{-MIL-88B (Fe)}$  was narrowed after the introduction and the specific surface area was relatively reduced because of the large ion size of  $\text{In(III)}$ .

The morphology of the  $\text{NH}_2\text{-MIL-88B (Fe}_{0.6}\text{In}_{0.4})$  in the TEM image is a hexagonal prism with sharp ends, as shown in

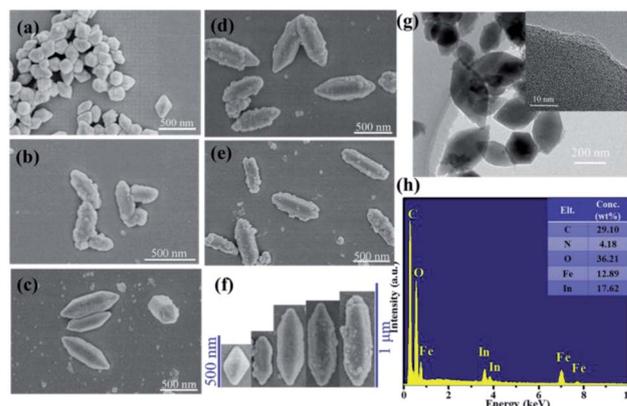


Fig. 2 SEM images of (a)  $\text{NH}_2\text{-MIL-88B (Fe)}$ , (b–e)  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  samples ( $\alpha = 0.8, 0.6, 0.4, 0.2$ , respectively), (f) the compared SEM images of  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$ , (g) TEM image of  $\text{NH}_2\text{-MIL-88B (Fe}_{0.6}\text{In}_{0.4})$  and (h) EDX image of  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$ .

Fig. 2g, which is the same as the result obtained in the SEM image. HRTEM displayed that  $\text{NH}_2\text{-MIL-88B (Fe}_{0.6}\text{In}_{0.4})$  has an ordered pore structure, where is difficult to find nanoparticles or metal oxide clusters outside the catalyst. Thence,  $\text{In(III)}$  has been incorporated into the framework of  $\text{NH}_2\text{-MIL-88B (Fe)}$  instead of being supported on the surface. As shown in Fig. 2h,  $\text{In}$  and  $\text{Fe}$  elements were approved in the EDX images, which even demonstrated the conformation of MM-MOFs  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$ .

### XPS analysis

The XPS spectra of  $\text{NH}_2\text{-MIL-88B (Fe}_{0.6}\text{In}_{0.4})$  composite is shown in Fig. 3a. There are five characteristic peaks in the survey spectrum, which corresponding to the  $\text{C 1s}$ ,  $\text{N 1s}$ ,  $\text{In 3d}$ ,  $\text{O 1s}$ , and  $\text{Fe 2p}$  electron orbitals, respectively. These elements confirm the results of the EDX analysis as displayed in Table S3.† Fig. 3b is the characteristic peak of  $\text{C}$ , all of them are derived from organic ligands. The characteristic peaks at 284.4, 286.2, and 288.2 eV could be attributed to  $\text{C=C}$ ,  $\text{C-N}$  and  $\text{C=O}$  respectively, which correspond to various carbon-containing covalent bonds in  $\text{NH}_2\text{-BDC}$ .<sup>53</sup> In Fig. 3c, the XPS spectrum of  $\text{N 1s}$  can be divided to 396.8 and 397.9 eV, which represents the  $\text{-N}^{\text{=+}}$  and  $\text{-NH}^{\text{+}}$  respectively. It belonged to the amino group extending or protruding into the cavity and the positively charged. There are four characteristic peaks in the XPS spectrum of  $\text{O 1s}$  shown in Fig. 3d. The peaks at 531.7 and 530.8 eV can be attributed to  $\text{-OH}$  and  $\text{C=O}$  of the organic ligand. Also, 529.9 and 529.1 eV correspond to the characteristic peaks of  $\text{In-O}$  and  $\text{Fe-O}$  respectively. The appearance of  $\text{In-O}$  clusters shows that  $\text{In(III)}$  has been doped into the catalyst and connected to the organic ligands, thus successfully constructing  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$ , which is consistent with the results of TEM. Fig. 3e shows the characteristic peaks of  $\text{Fe 2p}$ , where the satellite peak of  $\text{Fe}$  is at 715.2 eV. Among the peaks of  $\text{Fe 2p}_{1/2}$  and  $\text{Fe 2p}_{3/2}$ , 724.0 eV and 710.0 eV represent  $\text{Fe}^{3+}$ , and 722.2 eV and 709.0 eV represent  $\text{Fe}^{2+}$ .<sup>54–56</sup> The appearance of  $\text{Fe}^{2+}$  is due



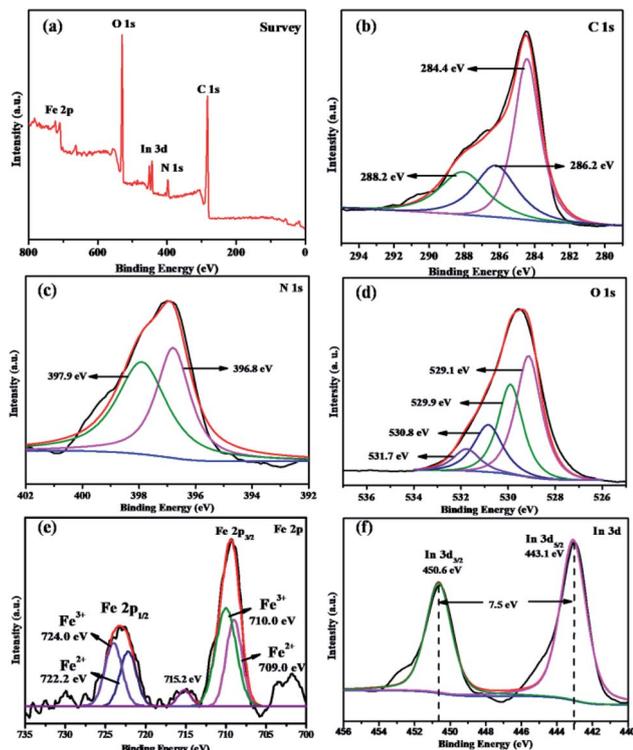


Fig. 3 XPS spectra of  $\text{NH}_2\text{-MIL-88B (Fe}_{0.6}\text{In}_{0.4})$  composite (a). Survey; (b) C 1s; (c) N 1s; (d) O 1s; (e) Fe 2p; (f) In 3d, respectively.

to the combination of photogenerated electrons and  $\text{Fe}^{3+}$ , reducing  $\text{Fe}^{3+}$  to  $\text{Fe}^{2+}$ . The peak at 450.6 and 443.1 eV can be verified to  $\text{In } 3d_{3/2}$  and  $\text{In } 3d_{5/2}$  as presented in Fig. 3f, respectively. This once again proves that In is successfully doped in  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$ . Contacted with the characterization results of XRD, IR, EDS, and XPS, it can be further confirmed that the In-doped  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  bimetallic material has been successfully prepared.

### Optical properties

The UV-vis DRS spectra of the samples were used to demonstrate the optical absorption properties in Fig. 4a. As reduced, the absorption intensity of  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  gradually decreases in the visible light region.  $\text{NH}_2\text{-MIL-88B (Fe}_{0.8}\text{In}_{0.2})$

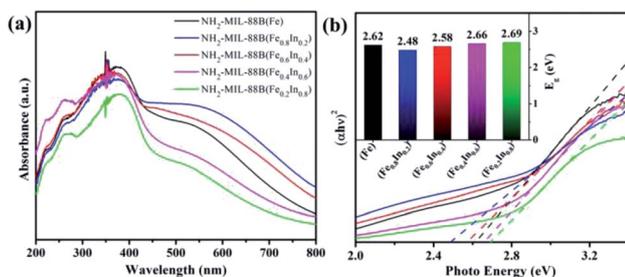


Fig. 4 (a) UV-vis DRS spectrum and (b) plots of  $(\alpha h\nu)^2$  versus photon energy ( $h\nu$ ) of the  $\text{NH}_2\text{-MIL-88B (Fe)}$  and the  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  samples.

has the strongest absorption intensity of visible light, and  $\text{NH}_2\text{-MIL-88B (Fe}_{0.6}\text{In}_{0.4})$  has a slight enhance in visible light absorption intensity. Comparing with  $\text{NH}_2\text{-MIL-88B (Fe)}$ , both have significantly enhanced visible light absorption. Thereby, this indicates that a certain proportion of In(III) doping, which is beneficial to improve the visible light absorption capacity of  $\text{NH}_2\text{-MIL-88B (Fe)}$ .

The corresponding bandgap energy ( $E_g$ ) values could be calculated by following the eqn (1)

$$(\alpha h\nu)^2 = A(h\nu - E_g) \quad (2)$$

here,  $\alpha$ ,  $h$ ,  $\nu$ ,  $A$ ,  $E_g$  corresponds to the absorption coefficient, the Planck's constant, light frequency, constant, the bandgap energy, respectively. As displayed in Fig. 4b, the  $E_g$  values of  $\text{NH}_2\text{-MIL-88B (Fe)}$ ,  $\text{NH}_2\text{-MIL-88B (Fe}_{0.8}\text{In}_{0.2})$ ,  $\text{NH}_2\text{-MIL-88B (Fe}_{0.6}\text{In}_{0.4})$ ,  $\text{NH}_2\text{-MIL-88B (Fe}_{0.4}\text{In}_{0.6})$  and  $\text{NH}_2\text{-MIL-88B (Fe}_{0.2}\text{In}_{0.8})$  were estimated to be 2.62, 2.48, 2.58, 2.66 and 2.69 eV, respectively. The size of the bandgap energy is related to the difficulty of electronic excitation. Under visible light irradiation, the lower bandgap energy can allow more electrons to excite the transition to CB, promote the effective separation of electrons and holes, thereby improving the photocatalytic activity. Among them,  $\text{NH}_2\text{-MIL-88B (Fe}_{0.8}\text{In}_{0.2})$  and  $\text{NH}_2\text{-MIL-88B (Fe}_{0.6}\text{In}_{0.4})$  displayed the lower  $E_g$  and the stronger visible light response-ability, which may perform better photocatalytic performance.

As shown in Fig. S1,† the fluorescence intensity of  $\text{NH}_2\text{-MIL-88B (Fe}_{0.6}\text{In}_{0.4})$  is significantly lower than that of  $\text{NH}_2\text{-MIL-88B (Fe)}$ , which indicates that the addition of In(III) inhibits the recombination of photogenerated  $e^- - h^+$ , thereby allowing more photogenerated carriers to participate photocatalytic reaction.<sup>57,58</sup>

### Photocatalytic activities

For verifying the photocatalytic performance of  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  obtained from the characterization results, we conducted experiments on photocatalytic Cr(VI) reduction under different catalytic conditions. As displayed in Fig. 5a, the Cr(VI) concentration was unaltered significantly with visible light irradiation and no photocatalyst, at the same time, the small range change of Cr(VI) concentration was due to adsorption with photocatalyst and no light, which indicated that visible light irradiation played an important role in the photocatalytic reduction of Cr(VI). The photocatalytic reduction efficiency of  $\text{NH}_2\text{-MIL-88B (Fe}_{0.6}\text{In}_{0.4})$  for reducing Cr(VI) is 86.83%, which is significantly higher 1.7 times than that of  $\text{NH}_2\text{-MIL-88B (Fe)}$  of 50.96%, thereby, which demonstrated that the doping of In(III) effectively improves the photocatalytic reduction performance. As shown in Fig. 5b, the different In(III) doped amount of  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  was carried out to discuss the capability of photocatalytic Cr(VI) reduction. With the doping of In, the photocatalytic reduction effect of the catalyst gradually increases. Among them,  $\text{NH}_2\text{-MIL-88B (Fe}_{0.6}\text{In}_{0.4})$  has the highest photocatalytic reduction rate. By contrast,  $\text{NH}_2\text{-MIL-88B (Fe}_{0.8}\text{In}_{0.2})$  is the lowest one. With the addition of more In(III) content, the catalytic effect gradually decreases. This may be due to more In(III) doping, resulting in lower visible light absorption. And higher bandgap energy is not conducive to the



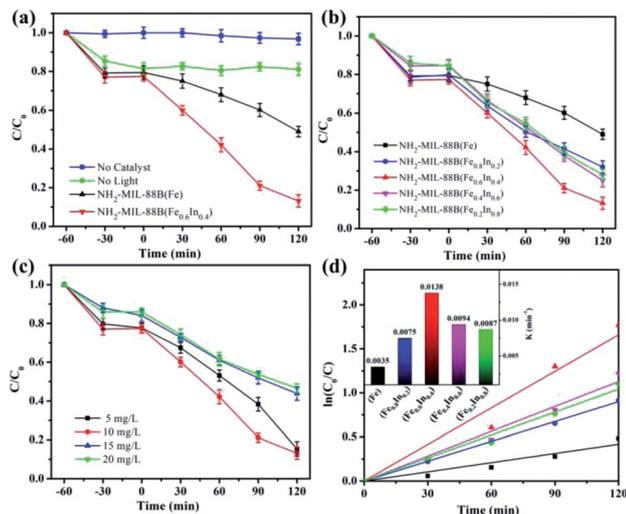


Fig. 5 Photocatalytic reduction of Cr(vi) (a) blank group comparison, (b)  $\text{NH}_2\text{-MIL-88B}(\text{Fe}_\alpha\text{In}_{1-\alpha})$ , (c) Cr(vi) on different concentrations reduction performance, and (d) the fitting of the pseudo-first-order linear line for the photocatalytic reaction. (Reaction conditions:  $0.5 \text{ g L}^{-1}$  photocatalyst,  $3 \text{ mL}$  ethanol, RT,  $\text{pH} = 2$ ).

excitation of electrons on VB, thereby reducing the photocatalytic activity of photocatalytic reduction of hexavalent chromium. Meanwhile, Fig. 5c shows that the photocatalytic Cr(vi) on different concentrations reduction performance of  $\text{NH}_2\text{-MIL-88B}(\text{Fe}_{0.6}\text{In}_{0.4})$  was investigated. With increasing the amount of catalyst, the photocatalytic performance gradually increases, reaching the maximum at  $10 \text{ mg L}^{-1}$ . However, an excess of the catalyst leads to lower photocatalytic reduction efficiency. This may be due to excessive catalyst hindering the light irradiation of the Cr(vi) solution. Fig. 5d displays that the photocatalytic reduction efficiency has a good linear relationship with the reaction time, which conforms to the first-order reaction kinetics. The pseudo-first-order rate constant  $k$  is obtained by the following formula:

$$\ln(C_0/C) = kt \quad (3)$$

here,  $C_0$ ,  $C$ ,  $k$ , and  $t$  corresponds to the initial concentrations of Cr(vi), the concentrations of Cr(vi), the pseudo-first-order rate constant, and time, respectively. The value of  $k$  and  $R^2$  of  $\text{NH}_2\text{-MIL-88B}(\text{Fe})$  and  $\text{NH}_2\text{-MIL-88B}(\text{Fe}_\alpha\text{In}_{1-\alpha})$  was displayed in Tables S4 and S5.†

To investigate the oxidation performance of the prepared photocatalyst, the photocatalytic degradation of TC-HCl by  $\text{NH}_2\text{-MIL-88B}(\text{Fe}_\alpha\text{In}_{1-\alpha})$  under different catalytic conditions was performed. Similar to the photocatalytic reduction effect of Cr(vi), the test result of Fig. 6a shows that the light and photocatalyst are crucial in the reaction system. As shown in Fig. 6b, the photocatalytic degradation efficiency of  $\text{NH}_2\text{-MIL-88B}(\text{Fe}_{0.6}\text{In}_{0.4})$  on TC-HCl under visible light irradiation conditions is 72.05%, which is significantly higher 1.6 times than that of  $\text{NH}_2\text{-MIL-88B}(\text{Fe})$  of 44.68%. Fig. 6c demonstrated the effect of different concentrations photocatalytic degradation efficiency of TC-HCl. Among them, as the content of TC-HCl increases, the

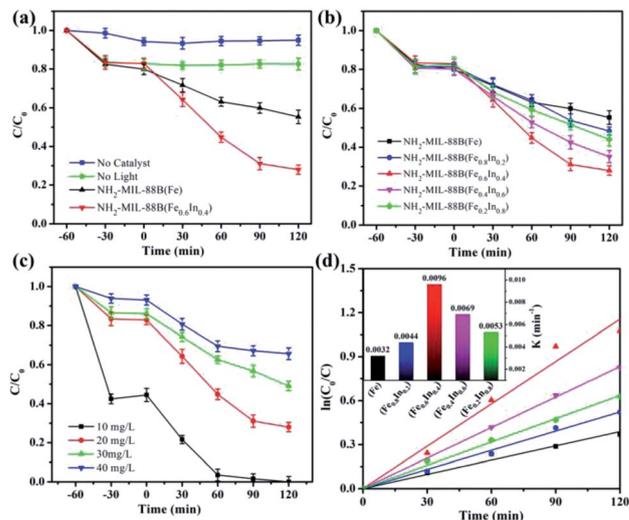


Fig. 6 Photocatalytic degradation of TC-HCl (a) blank group comparison, (b)  $\text{NH}_2\text{-MIL-88B}(\text{Fe}_\alpha\text{In}_{1-\alpha})$ , (c) TC-HCl on different concentrations degradation performance and (d) the fitting of the pseudo-first-order linear line for the photocatalytic reaction. (Reaction conditions:  $0.1 \text{ g L}^{-1}$  photocatalyst, RT).

degradation rate gradually decreases, and when the content is  $10 \text{ mg}$ ,  $\text{NH}_2\text{-MIL-88B}(\text{Fe}_{0.6}\text{In}_{0.4})$  displays the best degradation effect. The relationship between  $\ln(C_0/C)$  and time shows that the photocatalytic degradation efficiency has a good linear relationship with the reaction time, as presented in Fig. 6d, which is in line with the first-order reaction kinetics. The value of  $k$  and  $R^2$  of  $\text{NH}_2\text{-MIL-88B}(\text{Fe})$  and  $\text{NH}_2\text{-MIL-88B}(\text{Fe}_\alpha\text{In}_{1-\alpha})$  was displayed in Tables S6 and S7.†

To detect the reactive oxygen species generation during the photocatalytic degradation of TC-HCl by  $\text{NH}_2\text{-MIL-88B}(\text{Fe}_\alpha\text{In}_{1-\alpha})$ , electron spin resonance (ESR) technique is employed. Fig. 7 depicts the conspicuous characteristic peak signals of  $\text{DMPO}\cdot\text{O}_2^-$  and  $\text{DMPO}\cdot\text{OH}$ , which confirms the generation of  $\cdot\text{O}_2^-$  and  $\cdot\text{OH}$  radicals in the experiment. The peak intensity ratio of the quadruple characteristic peak signal of  $\text{DMPO}\cdot\text{OH}$  is about  $1 : 2 : 2 : 1$ .<sup>59</sup>  $\text{NH}_2\text{-MIL-88B}(\text{Fe}_\alpha\text{In}_{1-\alpha})$  by the photoexcitation generates  $e^-$  under visible light irradiation, which can activate dissolved oxygen in the solution that promotes oxygen molecules to become  $\cdot\text{O}_2^-$  species and can

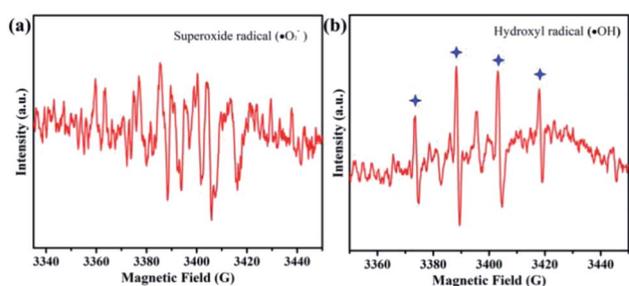


Fig. 7 ESR spectra of  $\text{DMPO}\cdot\text{O}_2^-$  and  $\text{DMPO}\cdot\text{OH}$  adducts generated  $\text{NH}_2\text{-MIL-88B}(\text{Fe}_\alpha\text{In}_{1-\alpha})$  after 60 seconds under visible light irradiation ( $\lambda > 400 \text{ nm}$ ) during the photocatalytic process.



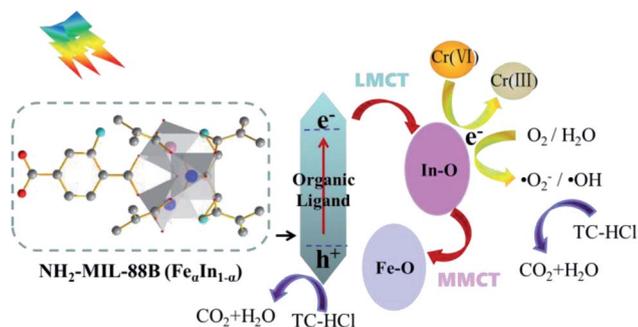
also interact with water molecules to generate  $\cdot\text{OH}$  species, simultaneously.

### Mechanism

Based on the above results, the possible mechanism of  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  photocatalytic treatment of  $\text{Cr(VI)}$  and TC-HCl in water is proposed. First, the electrons transfer process of  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  photocatalytic reduction of  $\text{Cr(VI)}$  is as follows. Under visible light irradiation, the organic ligand of  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  can be excited to generate electrons and holes. A part of the photogenerated electrons transfers to the surface of the catalyst through the In–O clusters through the LMCT to reduce  $\text{Cr(VI)}$  adsorbed directly. Then, the In–O clusters transfer another electron to the Fe–O cluster through MMCT, which promotes the separation of electrons and holes effectively.<sup>49</sup> The excitation and transfer of photogenerated carriers of photocatalytic oxidation of TC-HCl are similar to the process of reduction of  $\text{Cr(VI)}$ . The photogenerated electrons transfer to the surface of the catalyst combine with  $\text{O}_2$  and  $\text{H}_2\text{O}$  in the solution to produce  $\cdot\text{O}_2^-$  and  $\cdot\text{OH}$ . Both the holes generated by  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  excited by visible light and  $\cdot\text{O}_2^-/\cdot\text{OH}$  can interact with TC-HCl directly and eventually generate to  $\text{CO}_2$  and  $\text{H}_2\text{O}$ . Anyway, the MMCT in the  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  expands the active sites of photocatalytic oxidation-reduction. Also, the synergistic process of LMCT and MMCT effectively inhibits the recombination of electrons and holes, thereby enhancing the performance of the  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  (Scheme 2).

### Conclusions

In summary, a series of  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  photocatalysts were designed and synthesized *via* a one-pot method. Proved by characterizations of TEM and XPS, In(III) ions successfully replaced part of Fe(III) ions in the structure, which formed mixed-metal MOFs  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$ . Compared with single  $\text{NH}_2\text{-MIL-88B (Fe)}$ ,  $\text{NH}_2\text{-MIL-88B (Fe}_{0.6}\text{In}_{0.4})$  shows excellent photocatalytic performance: the reduction rate of  $\text{Cr(VI)}$  and the oxidation rate of TC-HCl reach 86.83% and 72.05%, which 1.7 and 1.6 times that of  $\text{NH}_2\text{-MIL-88B (Fe)}$ , respectively. The improvement of photocatalytic performance is



Scheme 2 Possible mechanism of photocatalytic degradation of  $\text{Cr(VI)}$  and TC-HCl by  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$ .

mainly attributed to the more reactive sites when In(III) doped to  $\text{NH}_2\text{-MIL-88B (Fe)}$ . Simultaneously, the synergistic effect of LMCT and MMCT of  $\text{NH}_2\text{-MIL-88B (Fe}_\alpha\text{In}_{1-\alpha})$  effectively expands the separation and transfer of photogenerated carriers and inhibits the recombination of electrons–holes pairs, thereby enhancing the performance of photocatalytic reduction of  $\text{Cr(VI)}$  and oxidative degradation of TC-HCl. This work may provide an insight to design and develop mixed-metal MOFs as photocatalysts for the wastewater treatment.

### Conflicts of interest

There are no conflicts to declare.

### Acknowledgements

Thanks to the 2020 School-level Scientific Research Project of Dalian Vocational & Technical College (Dalian Radio and TV University) and the Key Project of Natural Science Foundation of Liaoning Province (No. 20170540578).

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