


Cite this: *Nanoscale Adv.*, 2019, 1, 1812

# Truncated octahedral bipyramidal $\text{TiO}_2/\text{MXene}$ $\text{Ti}_3\text{C}_2$ hybrids with enhanced photocatalytic $\text{H}_2$ production activity†

Yang Li,<sup>ab</sup> Dainan Zhang,<sup>a</sup> Xionghan Feng,<sup>b</sup> Yulong Liao,<sup>a</sup> Qiye Wen<sup>a</sup> and Quanjun Xiang<sup>id</sup> <sup>\*a</sup>

MXene  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  hybrids were successfully synthesized through a simple calcination of F-terminated  $\text{Ti}_3\text{C}_2$ . The resultant  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  composite photocatalysts retained a 2D multilayer structure like MXene  $\text{Ti}_3\text{C}_2$ , and  $\text{TiO}_2$  exhibited a truncated octahedral bipyramidal structure with exposed (001) facets under the participation of fluorine ions. The residual  $\text{Ti}_3\text{C}_2$  could act as a co-catalyst to enhance the photocatalytic  $\text{H}_2$  production activity by capturing photogenerated electrons from  $\text{TiO}_2$  because of its electron reservoir feature and suitable Fermi level. The (101)–(001) surface heterojunction of the truncated octahedral bipyramidal  $\text{TiO}_2$  further accelerated the separation of photogenerated carriers. As a result, the  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  hybrids with calcining F-terminated  $\text{Ti}_3\text{C}_2$  exhibited photocatalytic hydrogen production that is twofold higher than that of  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  hybrids with calcining OH-terminated  $\text{Ti}_3\text{C}_2$ . This work presented a new strategy to prepare MXene  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  hybrids for photoconversion applications.

Received 14th January 2019

Accepted 3rd March 2019

DOI: 10.1039/c9na00023b

rsc.li/nanoscale-advances

## Introduction

Photocatalytic  $\text{H}_2\text{O}$  decomposition for hydrogen production has great potential in solving environmental pollution and energy crisis because of the high chemical energy and environmental friendliness of hydrogen.<sup>1–5</sup> As the first discovered photocatalyst,<sup>6</sup>  $\text{TiO}_2$  has many advantages, such as low cost, effectiveness, environmental friendliness, and long-term stability.<sup>7–10</sup> However, in practical applications, there are still many problems with  $\text{TiO}_2$  due to the fast recombination of photogenerated carriers and low light utilization capacity.<sup>11,12</sup> Therefore, many strategies, such as modifying co-catalysts,<sup>13–19</sup> doping impurities,<sup>20–23</sup> and changing its shape,<sup>24,25</sup> have been proposed to address these issues. Among these strategies, co-catalysts have been successfully developed for enhancing the activity and stability of  $\text{TiO}_2$ . Unfortunately, the high price, extreme scarcity (such as Pt), and destructive  $\pi$ -conjugated system (such as graphene oxide)<sup>26</sup> of co-catalysts restrict their application. Therefore, exploring a suitable co-catalyst is beneficial to the improvement of photocatalytic hydrogen production.

Recently, 2D MXenes have been proposed as emerging materials because of their hydrophilicity, photostability, large

specific surface area, and high conductivity.<sup>27–29</sup> In general, MXene materials are prepared by etching ternary nitrides/carbides by using the following formula:  $\text{M}_{n+1}\text{AX}_n$ , where M is a transition metal, A is a group III (or IV) element, and X is nitrogen and carbon.<sup>30,31</sup> The resultant MXene samples usually exhibit a multilayer structure with a large layer spacing. Based on these characteristics, 2D MXene materials can be used as a co-catalyst for photocatalytic applications. For instance, Peng *et al.* reported that  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  composite photocatalysts prepared by hydrothermally exhibit an enhanced photocatalytic degradation activity of MO. They found that  $\text{Ti}_3\text{C}_2$  can be used as a reservoir of photogenerated holes by building Schottky batteries because of its lower work function.<sup>32</sup> Li *et al.* synthesized  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  nanoflowers and  $\text{Ti}_3\text{C}_2/\text{MoS}_2/\text{TiO}_2$  hybrids by different methods. They pointed out that  $\text{Ti}_3\text{C}_2$  can enhance the photocatalytic activity of those composite photocatalysts due to the Schottky junction between  $\text{Ti}_3\text{C}_2$  and  $\text{TiO}_2$  and its excellent electronic conductivity.<sup>33,34</sup> Cai *et al.* pointed out that  $\text{Ti}_3\text{C}_2$  in an  $\text{Ag}_3\text{PO}_4/\text{Ti}_3\text{C}_2$  Schottky catalyst can greatly improve the catalytic performance and stability of  $\text{Ag}_3\text{PO}_4$  due to its profuse surface hydrophilic functional groups and the Schottky junction was formed at the interface of  $\text{Ag}_3\text{PO}_4$ – $\text{Ti}_3\text{C}_2$ .<sup>35</sup> In general, MXene  $\text{Ti}_3\text{C}_2$  possesses large surface functionalities (–F, –OH, and –O) because of HF treatment of  $\text{Ti}_3\text{AlC}_2$ . Notably, F-terminated  $\text{Ti}_3\text{C}_2$  has unfavourable Fermi levels (0.15 V);<sup>36</sup> as such, it cannot be an eligible candidate. Nevertheless, fluorine ions play an important role in the preparation of  $\text{TiO}_2$  with exposed high energy (001) facets because it can combine Ti to reduce the surface energy of (001) facets.<sup>37</sup>  $\text{TiO}_2$  with exposed high energy (001) facets can be synthesized with the participation of fluorine.

<sup>a</sup>State Key Laboratory of Electronic Thin Film and Integrated Devices, University of Electronic Science and Technology of China, Chengdu 610054, P. R. China. E-mail: xiangqj@uestc.edu.cn

<sup>b</sup>College of Resources and Environment, Huazhong Agricultural University, Wuhan 430070, P. R. China

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c9na00023b



Meanwhile, fluorine on the surface of  $\text{Ti}_3\text{C}_2$  is sharply decreased during calcination. However, to our knowledge, most studies on  $\text{Ti}_3\text{C}_2$  have directly removed fluorine ions by washing them with a reagent, and few studies have reported about the application of fluorine in photocatalysis.

Herein, a simple calcination synthesis method was first proposed to prepare truncated octahedral bipyramidal  $\text{TiO}_2$  (TOB-T)/MXene  $\text{Ti}_3\text{C}_2$  composite photocatalysts. The resultant  $\text{TiO}_2/\text{Ti}_3\text{C}_2$  hybrids retained the 2D multilayer structure and  $\text{TiO}_2$  exhibited a truncated octahedral bipyramidal structure with exposed (001) and (101) facets. A surface heterojunction between (101) and (001) facets was established, and it could prevent the recombination of photogenerated carriers in  $\text{TiO}_2$ . Moreover, the remaining  $\text{Ti}_3\text{C}_2$  could act as a co-catalyst to accelerate the migration of photoinduced electrons because of its high electronic conductivity. Meanwhile, the concentration of fluorine sharply decreased during calcination, thereby reducing the toxicity and increasing the conductivity of the samples.

## Results and discussion

In Fig. 1, the preparation process of MXene  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  composite photocatalysts was divided into two parts. In Part I, different T-terminated (T = F, O, and OH)  $\text{Ti}_3\text{C}_2$  specimens were prepared. Firstly, the initial  $\text{Ti}_3\text{C}_2$  was prepared by HF-treatment of  $\text{Ti}_3\text{AlC}_2$  power,<sup>38</sup> and the transformation from  $\text{Ti}_3\text{AlC}_2$  into initial  $\text{Ti}_3\text{C}_2$  was confirmed through X-ray diffraction (XRD). In Fig. S1a (see the ESI†), the diffraction curves of the (002) and (004) planes of  $\text{Ti}_3\text{AlC}_2$  at  $9.58^\circ$  and  $19.17^\circ$  were weaker and shifted toward  $8.84^\circ$  and  $18.13^\circ$ , suggesting the removal of the Al layers and the formation of  $\text{Ti}_3\text{C}_2$  after HF treatment.<sup>39</sup> Meanwhile, the obtained precipitates possessed abundant functionalities (–F, –OH, and –O). Then, the first route to the treatment of the initial  $\text{Ti}_3\text{C}_2$  was washing directly with deionized water three times, and the obtained sample was denoted as F- $\text{Ti}_3\text{C}_2$  (the F- $\text{Ti}_3\text{C}_2$  sample contained –F, –OH and –O functionalities). To highlight the effect of fluorine ions, another route to treat the initial  $\text{Ti}_3\text{C}_2$  was chosen to remove fluorine ions. Namely, the obtained initial  $\text{Ti}_3\text{C}_2$  was first disposed with NaOH solution and then washed with deionized water, and the product was referred to as OH- $\text{Ti}_3\text{C}_2$  (the OH- $\text{Ti}_3\text{C}_2$  sample still contained –OH and –O functionalities). The crystal structures

and chemical compositions of F- $\text{Ti}_3\text{C}_2$  and OH- $\text{Ti}_3\text{C}_2$  were determined through XRD and XPS, respectively. In Fig. S1b (see the ESI†), F- $\text{Ti}_3\text{C}_2$  and OH- $\text{Ti}_3\text{C}_2$  exhibited similar XRD patterns, indicating that the treatment of NaOH solution did not change the crystal structure of  $\text{Ti}_3\text{C}_2$ . The survey XPS patterns (Fig. S2a in the ESI†) showed the existence of Ti, O, and C. In comparison with F- $\text{Ti}_3\text{C}_2$ , an obvious weakness of the F peak could be observed in the high-resolution F 1s XPS spectra of OH- $\text{Ti}_3\text{C}_2$  (Fig. 2a), suggesting that the fluorine ions in OH- $\text{Ti}_3\text{C}_2$  were removed by treatment with NaOH solution. The high-resolution XPS spectra of Ti 2p, C 1s, and O 1s are summarized in Tables S1 and S2 (in the ESI†). The existence of  $\text{Ti}_3\text{C}_2$  is demonstrated by various surface compositions (Ti–C, C–Ti–O<sub>x</sub>, and so on). These results confirmed that F- $\text{Ti}_3\text{C}_2$  and OH- $\text{Ti}_3\text{C}_2$  were successfully synthesized, and the difference in F- $\text{Ti}_3\text{C}_2$  and OH- $\text{Ti}_3\text{C}_2$  was only fluorine concentration because the treatment of NaOH solution removed fluorine ions without affecting the crystal structure of  $\text{Ti}_3\text{C}_2$ .

In Part II, different MXene  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  composite photocatalysts were prepared. The thermal oxidation method was selected to synthesize MXene  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  composite photocatalysts because it could remove most of the fluorine ions. Namely, F- $\text{Ti}_3\text{C}_2$  and OH- $\text{Ti}_3\text{C}_2$  were calcined at  $550^\circ\text{C}$  for 4 h. After calcination was performed, the obtained samples of F- $\text{Ti}_3\text{C}_2$  and OH- $\text{Ti}_3\text{C}_2$  were labelled TF and TOH, respectively. The crystal structures of TF and TOH were characterized through XRD and Raman patterns. In Fig. 2b, TF and TOH samples exhibit a typical XRD diffraction line of anatase,<sup>40</sup> confirming that no rutile and brookite, in addition to anatase, were observed after calcination. Furthermore, a diffraction peak assigned to  $\text{Ti}_3\text{C}_2$  could be detected at  $43^\circ$  and  $57.5^\circ$  (see the inset in Fig. 2b), suggesting the generation of  $\text{TiO}_2/\text{Ti}_3\text{C}_2$  hybrids.<sup>28</sup> In the Raman patterns of the TF and TOH samples (Fig. 2c), four characteristic peaks located at  $144$ ,  $400$ ,  $415$ , and  $650\text{ cm}^{-1}$  corresponded to the  $\text{E}_g$ ,  $\text{B}_{1g}$ ,  $\text{A}_{1g}$ , and  $\text{E}_g$  modes of the typical anatase phase.

Notably, the diffraction intensity of  $\text{E}_g$  of TF sample peaks located at  $144$  and  $636\text{ cm}^{-1}$  was weaker than that of the TOH

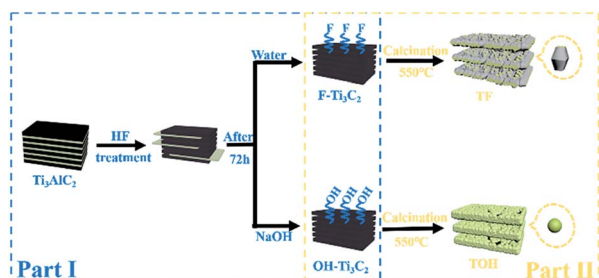


Fig. 1 Schematic of the preparation procedure for the synthesis of MXene  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  composite photocatalysts.

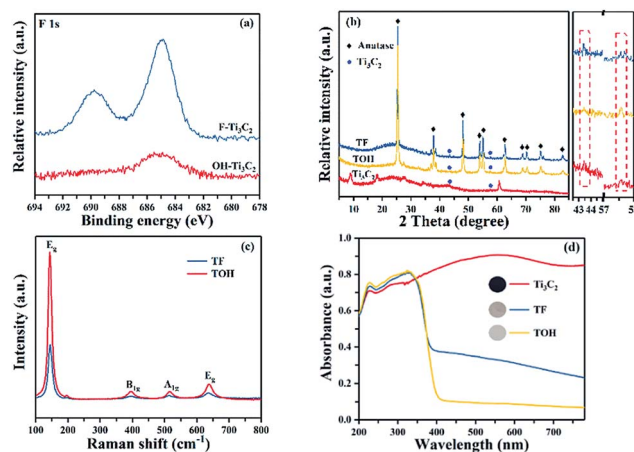


Fig. 2 (a) High-resolution F 1s XPS spectra of F- $\text{Ti}_3\text{C}_2$  and OH- $\text{Ti}_3\text{C}_2$ , (b) XRD patterns of TF, TOH, and  $\text{Ti}_3\text{C}_2$  samples, (c) Raman patterns of TF and TOH samples, and (d) DRS patterns of TF and TOH samples.

sample, suggesting the presence of a large number of (001) facets in the TF samples because the number of the symmetric stretching vibration modes of O–Ti–O decreases when the exposed (001) facets exist.<sup>41,42</sup> This result indicated that  $\text{TiO}_2$  with exposed (001) facets can be found in the TF samples. The UV-Vis diffuse reflectance spectroscopy (UV-Vis DRS) spectra were used to further confirm the existence of  $\text{Ti}_3\text{C}_2$  and the optical properties of the  $\text{Ti}_3\text{C}_2$ , TF, and TOH samples. In Fig. 2d,  $\text{Ti}_3\text{C}_2$  demonstrated the highest absorption intensity in the full spectrum because of the black feature of  $\text{Ti}_3\text{C}_2$ . The curves of the TF and TOH samples exhibited a similar trend, which corresponded to the intrinsic properties of anatase  $\text{TiO}_2$ , suggesting the formation of  $\text{TiO}_2$ .<sup>43,44</sup> By contrast, the background absorption intensity of TF was higher than that of TOH. This phenomenon should be attributed to the concentration of  $\text{Ti}_3\text{C}_2$  in the TF sample, which was higher than that of the TOH sample. This result is easy to understand; generally, the high energy (001) facets is difficult to synthesize. However,  $\text{TiO}_2$  with exposed (001) facets can be found in the TF samples due to the fluorine reduced the surface energy of (001) facets. Therefore, the transformation of  $\text{Ti}_3\text{C}_2$  into  $\text{TiO}_2$  was prevented because the chemical energy in the system is additionally consumed under the action of fluorine. The result indicated that the concentration of  $\text{Ti}_3\text{C}_2$  in the TF sample was higher than that of the TOH sample. These results indicated that the MXene  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  composite photocatalysts were successfully synthesized through a simple calcination method, and the composition of both TF and TOH samples was  $\text{Ti}_3\text{C}_2/\text{TiO}_2$ . Moreover,  $\text{TiO}_2$  exposed (001) facets could be found in the TF samples under the action of fluorine ions.

XPS characterization was conducted to analyse the chemical composition of F- $\text{Ti}_3\text{C}_2$  before and after calcination. In Fig. S2i (see in the ESI†), four elements (Ti, C, O, and F) could be found in the survey curves of TF and F- $\text{Ti}_3\text{C}_2$  samples. Obviously, the diffraction peak of F in the TF samples was considerably weaker than that of the F- $\text{Ti}_3\text{C}_2$  samples, suggesting that fluorine was sharply reduced during calcination. The high resolution Ti 2p curve of the F- $\text{Ti}_3\text{C}_2$  samples is shown in Fig. S2b (see in the ESI†). Six different peaks can be discovered and attributed to the different compositions of Ti.<sup>45–48</sup> However, in the high resolution Ti 2p curve of the TF samples (Fig. S2c in the ESI†), only two obvious peaks were found, and they correspond to  $\text{TiO}_2$  2p<sub>1/2</sub> and  $\text{TiO}_2$  2p<sub>3/2</sub>.<sup>49</sup> This phenomenon indicated the transformation of  $\text{Ti}_3\text{C}_2$  into  $\text{TiO}_2$ . The high resolution spectra of C 1s, O 1s, and F 1s are summarized in Tables S1 and S2 (in the ESI†).

The morphological characteristics of the products before and after calcination treatment can be observed in Fig. 3. In Fig. 3a and b, the  $\text{Ti}_3\text{C}_2$  MXene exhibited a multilayer structure after  $\text{Ti}_3\text{AlC}_2$  was treated with HF. The thickness of the single layer was approximately 5–15 nm. Interestingly, the TF and TOH samples still maintained a 2D multilayer structure on account of the *in situ* transformation of the  $\text{Ti}_3\text{C}_2$  MXene into  $\text{TiO}_2$  (Fig. 3c and e). Actually, the structures of the truncated octahedral bipyramid for the TF sample are  $\text{TiO}_2$  structures, and the layered structures are residual  $\text{Ti}_3\text{C}_2$  structures. So uniformly truncated octahedral bipyramidal  $\text{TiO}_2$  is formed on the surface

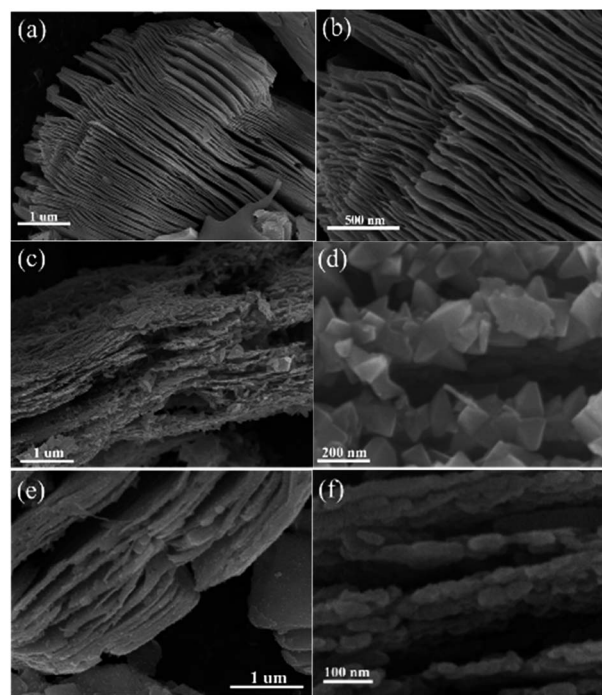


Fig. 3 SEM images of (a and b)  $\text{Ti}_3\text{C}_2$ , (c and d) TF, and (e and f) TOH samples.

of  $\text{Ti}_3\text{C}_2$  for the TF sample. For the TOH sample, the nanoparticles are  $\text{TiO}_2$  nanoparticles and layered structures are  $\text{Ti}_3\text{C}_2$  structures. This result should be attributed to the fact that fluorine on the surface of the TF sample reduced the surface energy of  $\text{TiO}_2$ , which facilitated the formation of  $\text{TiO}_2$  with exposed (001) facets.<sup>28</sup> For the TOH sample, only sphericity with the lowest surface energy was produced during crystallization due to the absence of fluorine to reduce its surface energy. In addition, the surface area of the TOH sample ( $41.3 \text{ m}^2 \text{ g}^{-1}$ ) was larger than that of TF ( $20.5 \text{ m}^2 \text{ g}^{-1}$ ). This results also indicated that smaller particles were formed in the surface of TOH than that for TF.

The transmission electron microscopy (TEM) and high-resolution TEM (HRTEM) images of TOH and TF are shown in Fig. 4. The TEM images further confirmed the 2D multilayer structure of the TOH and TF samples (Fig. 4a and c). Clear lattice fringes were found in the TOH and TF samples. In detail, the lattice fringes of 0.35 and 0.27 nm were measured in the TOH sample, and they corresponded to the (101) facets of anatase  $\text{TiO}_2$  and (0110) facets of  $\text{Ti}_3\text{C}_2$  (Fig. 4b),<sup>50,51</sup> respectively. The truncated octahedral bipyramid of  $\text{TiO}_2$  can be clearly observed in the TF samples, and the lattice fringe of the bevel in the truncated octahedral bipyramid was measured to be 0.35 nm, which was assigned to the (001) plane of  $\text{TiO}_2$ . However, the lattice fringe of the top surface in the truncated octahedral bipyramid could not be detected because TOB-T was too long for the electron beam to penetrate. These results suggested that the  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  hybrids were successfully synthesized in TOH and TF samples. By contrast, TOB-T was formed on the surface of the TF sample under the influence of fluorine ions, and nanoparticles were found on the surface of the TOH







Fig. 4 HRTEM images of (a and b) TOH and (c and d) TF samples.

sample. The percentage of the exposed (001) facets of the TF sample was calculated to be 11% based on the area ratio of the inclined plane to the top surface.

Photocurrent and electrochemical impedance spectra (EIS) were obtained to further investigate the separation and migration of the photogenerated carriers. The photocurrent of pure  $\text{TiO}_2$  without a co-catalyst was hardly detected because of the rapid recombination of photogenerated carriers. In Fig. 5a, upon light irradiation, a remarkable photocurrent was produced in the TF and TOH samples because of the existence of  $\text{Ti}_3\text{C}_2$ . Furthermore, the stronger photocurrent in the TF samples than that in the TOH samples was attributed to its (001)–(101) surface heterojunction. The EIS of TF and TOH is shown in Fig. 5b. The impedance of the TF samples was lower than that of the TOH samples, indicating the low charge

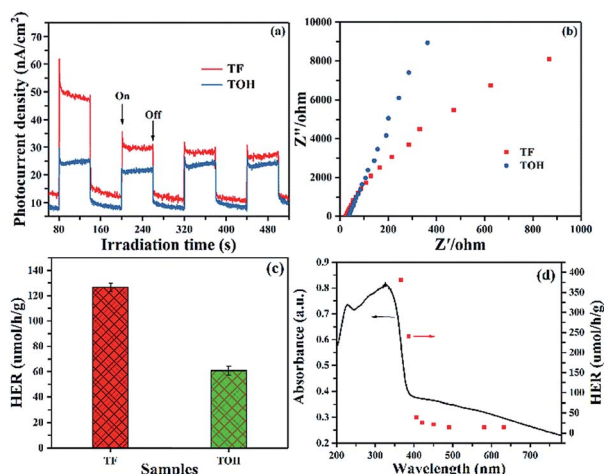


Fig. 5 (a) Photocurrent responses and (b) EIS patterns of TF and TOH samples. (c) Hydrogen evolution rate (HER) of the samples under 1 h light irradiation using glycerinum/water (1 : 10 vol%) solution as a sacrificial agent and (d) wavelength-dependent HER of TF under 1 h monochromatic light irradiation by using glycerinum/water (1 : 10 vol%) solution as a sacrificial agent.

transfer resistance of the TF samples. These results confirmed that the process of charge transfer and the increase in the separation and migration of the photogenerated carriers should be attributed to the surface heterojunction of (001)–(101) facets and the electron reservoir feature of  $\text{Ti}_3\text{C}_2$ .

Photocatalytic hydrogen production was performed in glycerinum/water (1 : 10 vol%) solution under light irradiation to conclude the photocatalytic performance of TF and TOH samples. In Fig. 5c, a remarkable hydrogen evolution rate (HER) of the TF and TOH samples could be obtained because of the existence of the co-catalyst  $\text{Ti}_3\text{C}_2$ . The HER of the TF samples was considerably higher than that of the TOH samples by a factor of 2, suggesting that the exposed (001) facets are key factors for the improvement of the photocatalytic activity of the TF samples. A wavelength-dependent HER of the TF samples was determined using a monochromatic light source. In Fig. 5d, the wavelength-dependent HER increased as the energy of monochromatic light in the excitation region increased. This result indicated that the HER is mainly controlled by light-induced electrons in the TF samples and was in line with the DRS spectrum. Simultaneously, the TF samples exhibited the highest HER with an apparent quantum efficiency of 0.9% under 365 nm monochromatic light irradiation.

Based on the experimental results, Fig. 6 illustrates the tentative mechanism of  $\text{TiO}_2/\text{Ti}_3\text{C}_2$  hybrids for photocatalytic  $\text{H}_2$  evolution over the TF samples. Under light irradiation, the (001) and (101) facets of TOB-T could be excited to produce photogenerated electrons and holes. The surface band gap of (001) facets is misaligned with that of (101) facets because of the different atomic arrangements on the surface of each facet.<sup>52,53</sup> Therefore, the formation of the surface heterojunction between (001) and (101) facets could facilitate the photogenerated electrons to transfer from the (001) to (101) facets because of the staggered band gap. This result led to the accumulation of photogenerated electrons and holes on the (101) and (001) facets, respectively. Moreover, the photogenerated electrons could be transferred from  $\text{TiO}_2$  to  $\text{Ti}_3\text{C}_2$  because of the high conductivity of  $\text{Ti}_3\text{C}_2$ .<sup>54</sup> Namely,  $\text{Ti}_3\text{C}_2$ , as an electron reservoir in  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  hybrids, captured the photogenerated electrons of (101) facets (see the detail of the bandgap information about the hetero-structure in Fig. S6 in the ESI†).

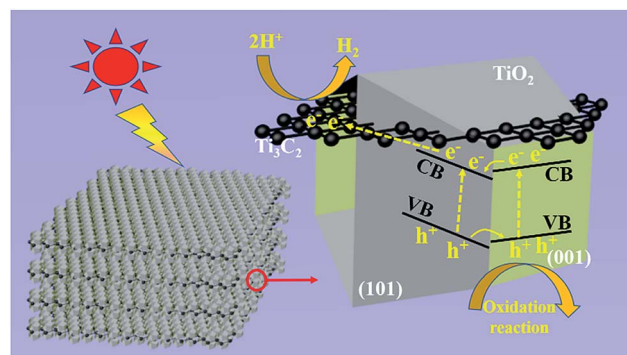


Fig. 6 Schematic of the tentative photocatalytic mechanism of truncated octahedral bipyramidal  $\text{TiO}_2/\text{Ti}_3\text{C}_2$  hybrids under light irradiation.

Therefore, an effective separation of the photogenerated carriers was achieved in the  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  hybrids through the transfer of the photogenerated electrons. Therefore, the synergy of the surface heterojunction of (001)–(101) facets and the electron reservoir feature of  $\text{Ti}_3\text{C}_2$  could effectively increase the HER of  $\text{TiO}_2/\text{Ti}_3\text{C}_2$  composite photocatalysts.

## Conclusions

In summary, binary 2D MXene  $\text{Ti}_3\text{C}_2/\text{TiO}_2$  hybrids were successfully synthesized through a simple calcination of F-terminated  $\text{Ti}_3\text{C}_2$ . TOB-T with exposed (001) facets could be found because of the effect of fluorine ions. The multilayer structure derived from MXene  $\text{Ti}_3\text{C}_2$  is essential for the interaction between  $\text{TiO}_2$  and  $\text{Ti}_3\text{C}_2$ . The enhanced HER should be attributed to the (001)–(101) surface heterojunction in the TF samples, in addition to the electron reservoir feature of  $\text{Ti}_3\text{C}_2$ . A tentative mechanism was proposed based on the experimental results. This work provided a new strategy for the application of 2D MXene materials.

## Experiment

### Materials

Hydrofluoric acid (40 wt%),  $\text{Ti}_3\text{AlC}_2$ , and NaOH were of analytical grade and used without further purification. Deionized water was used in all the experiments.

### Sample preparation

$\text{Ti}_3\text{C}_2$  was synthesized in accordance with previous studies.<sup>29</sup> In brief, 40 mL of hydrofluoric acid (40 wt%) was gradually dropped to disperse 2 g of  $\text{Ti}_3\text{AlC}_2$  powder. The resultant suspension was stirred at normal temperature for 72 h to etch an Al layer. The obtained precipitates were divided into two parts. One part was washed directly with deionized water thrice, and the other part was first disposed with NaOH solution (0.1 mol L<sup>-1</sup>) and then washed with deionized water. Finally, the precipitates were dried at 80 °C in an oven. The product was referred to as F- $\text{Ti}_3\text{C}_2$  for the former part and OH- $\text{Ti}_3\text{C}_2$  for the latter. A  $\text{TiO}_2/\text{Ti}_3\text{C}_2$  hybrid was prepared through primitive calcination. In brief, 0.3 g of F- $\text{Ti}_3\text{C}_2$  (OH- $\text{Ti}_3\text{C}_2$ ) was calcined at 550 °C for 4 h in air. The prepared samples were labeled TF (TOH), milled into powder, and collected.

### Characterization

The XRD patterns were recorded on a D8-Advance diffractometer (Bruker, German) with Ni-filtered Cu K $\alpha$  radiation at a scan rate (2 $\theta$ ) of 10° min<sup>-1</sup>, an accelerating voltage of 40 kV, and a current of 40 mA. The TEM images were obtained using a Hitachi H-7650 (HITACHI, Japan) transmission electron microscope at an accelerating voltage of 200 kV, and HRTEM analysis was performed on a JEM-2100F (JEOL, Japan) microscope. The SEM images were obtained on an S-4800 (HITACHI, Japan). The UV-Vis DRS spectra were obtained using a UV-Vis spectrophotometer (Lambda 650s), and Teflon was used as the reflectance standard. The XPS patterns were

obtained on a Leybold Heraeus-Shenyang SKL-12 X-ray photoelectron spectrometer, and Mg K $\alpha$  served as an excitation source. Raman spectra were obtained on a micro-Raman spectrometer (LabRAM HR Evolution) at room temperature under ambient conditions, and the excitation source was a 532 nm Ar<sup>+</sup> laser.

### Photocatalytic H<sub>2</sub> production activity

The photocatalytic H<sub>2</sub> production of the obtained photocatalysts was measured in a 100 mL three-necked flask at atmospheric pressure and room temperature. A 350 W Xe arc lamp was utilized as a light source and the sacrificial agent was 10% of glycerinum aqueous solution in volume. The detailed steps were as follows: 50 mg of samples was dispersed in 80 mL of sacrificial agent. The suspension solution mixture of photocatalysts was degassed using N<sub>2</sub> for 30 min to evict the remaining oxygen and air in the flask. After 1 h of illumination, 0.4 mL of gas was extracted and detected on a gas chromatograph (GC-2004C, Japan).

The wavelength-dependent HER of TF samples was determined by a similar method to the photocatalytic H<sub>2</sub> production, expect that the light source was replaced by a LED equipped with different monochromatic light. The irradiation intensity of the monochromatic light was about 80 mW cm<sup>-2</sup> by controlling internal power of monochromatic light. The QE was determined by using the nether equation:

$$\begin{aligned}\text{QE}[\%] &= \frac{\text{number of reacted electrons}}{\text{number of incident photons}} \times 100 \\ &= \frac{\text{number of evolved H}_2 \text{ molecules} \times 2}{\text{number of incident photons}} \times 100\end{aligned}$$

### Photoelectrochemical measurements

Photocurrent and EIS were obtained on a CHI660E (Shanghai Chenhua Limited, China) electrochemical analyzer, which was a three-electrode system that used Pt wire and Ag/AgCl as counter and reference electrodes, respectively. The working electrodes were glass electrodes with an effective area of ca. 1.2 cm<sup>2</sup>. A 350 W Xe arc lamp and 0.5 M Na<sub>2</sub>SO<sub>4</sub> aqueous solution were used as a light source and an electrolyte, respectively.

The glass working electrode was synthesized as follows: a mixture of 0.03 g of the sample, 0.03 g of polyethyleneglycol, and 0.5 mL of ethanol were ground into a slurry. A doctor blade method was used to evenly coat the slurry onto a 2 cm × 1.2 cm F-doped SnO<sub>2</sub>-coated glass electrode. Finally, the obtained glass electrodes were dried at 100 °C for 60 min in an oven.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was partially supported by the National Natural Science Foundation of China under Grant No. 51672099,



21403079, 61831012 and 51572042, Fundamental Research Funds for the Central Universities under No. 2017-QR-25, Sichuan Science and Technology Program under No. 2019JDR0027, Science Challenge Project under Grant No. TZ2018003, and International Science & Technology Cooperation Program under Grant No. 2015DFR50870.

## Notes and references

- 1 L. Cheng, Q. J. Xiang, Y. L. Liao and H. W. Zhang, *Energy Environ. Sci.*, 2018, **11**, 1362–1391.
- 2 F. Y. Cheng, H. Yin and Q. J. Xiang, *Appl. Surf. Sci.*, 2017, **391**, 432–439.
- 3 Q. J. Xiang, F. Y. Cheng and D. Lang, *ChemSusChem*, 2016, **9**, 996–1002.
- 4 J. W. Fu, Q. L. Xu, J. X. Low, C. J. Jiang and J. G. Yu, *Appl. Catal., B*, 2019, **243**, 556–565.
- 5 K. He, J. Xie, M. Li and X. Li, *Appl. Surf. Sci.*, 2018, **430**, 208–217.
- 6 A. Fujishima and K. Honda, *Nature*, 1972, **238**, 37–38.
- 7 C. Athanasekou, G. E. Romanos, S. K. Papageorgiou, G. K. Manolis, F. Katsaros and P. Falaras, *Chem. Eng. J.*, 2017, **318**, 171–180.
- 8 X. F. Wang, T. Y. Li, R. Yu, H. G. Yu and J. G. Yu, *J. Mater. Chem. A*, 2016, **4**, 8682–8689.
- 9 Z. Wang and X. J. Lang, *Appl. Catal., B*, 2018, **224**, 404–409.
- 10 T. D. Pham and B. K. Lee, *J. Catal.*, 2017, **345**, 87–95.
- 11 M. S. Akple, J. Low, S. W. Liu, B. Cheng, J. G. Yu and W. K. Ho, *J. CO<sub>2</sub> Util.*, 2016, **16**, 442–449.
- 12 P. N. Paulino, V. M. M. Salim and N. S. Resende, *Appl. Catal., B*, 2016, **185**, 362–370.
- 13 D. Zhang, X. Ma, H. Zhang, Y. Liao and Q. J. Xiang, *Mater. Today Energy*, 2018, **10**, 132–140.
- 14 Y. Xu, Y. G. Li, P. Wang, X. F. Wang and H. G. Yu, *Appl. Surf. Sci.*, 2018, **430**, 176–183.
- 15 X. Zhang, Y. J. Chen, Y. T. Xiao, W. Zhou, G. H. Tian and H. G. Fu, *Nanoscale*, 2018, **10**, 4041–4050.
- 16 D. Lang, F. Y. Cheng and Q. J. Xiang, *Catal. Sci. Technol.*, 2016, **6**, 6207–6216.
- 17 F. Chen, W. Luo, Y. P. Mo, H. G. Yu and B. Cheng, *Appl. Surf. Sci.*, 2018, **430**, 448–456.
- 18 M. Wu, J. Zhang, C. X. Liu, Y. S. Gong, R. Wang and B. B. He, *ChemCatChem*, 2018, **14**, 3069–3077.
- 19 D. Lang, T. Shen and Q. J. Xiang, *ChemCatChem*, 2015, **7**, 943–951.
- 20 P. Wang, Y. G. Lu, X. F. Wang and H. G. Yu, *Appl. Surf. Sci.*, 2017, **391**, 259–266.
- 21 D. N. Zhang, X. Y. Ma, H. W. Zhang, Y. L. Liao and Q. J. Xiang, *Mater. Today Energy*, 2018, **10**, 132–140.
- 22 T. Song, P. Y. Zhang, T. T. Wang, A. Ali and H. P. Zeng, *Nanoscale*, 2018, **10**, 2275–2284.
- 23 W. Chen, Y. H. Wang, S. Liu, L. Gao, L. Q. Mao, Z. Y. Fan, W. F. Shangguan and Z. Jiang, *Appl. Surf. Sci.*, 2018, **445**, 527–534.
- 24 M. Terracciano, V. Galstyan, L. Rea, M. Casalino, L. D. Stefauo and B. Sberveglieri, *Appl. Surf. Sci.*, 2017, **419**, 235–240.
- 25 Y. Y. Duan, L. Liang, K. L. Lv, Q. Li and M. Li, *Appl. Surf. Sci.*, 2018, **456**, 817–826.
- 26 Y. Hernandez, V. Nicolosi, M. Lotya, F. M. Blighe, Z. Y. Sun, S. De, I. T. McGovern, B. Holland, M. Byrne, Y. K. Gun'Ko, J. J. Boland, P. Niraj, G. Duesberg, S. Krishnamurthy, R. Goodhue, J. Hutchison, V. Scardaci, A. C. Ferrari and J. N. Coleman, *Nat. Nanotechnol.*, 2008, **3**, 563–568.
- 27 M. Naguib, M. Kurtoglu, V. Presser, J. Lu, J. J. Niu, M. Heon, L. Hultman, Y. Gogotsi and M. W. Barsoum, *Adv. Mater.*, 2011, **23**, 4248–4253.
- 28 Y. J. Li, X. T. Deng, J. Tian, Z. Q. Liang and H. Z. Cui, *Appl. Mater. Today*, 2018, **13**, 217–227.
- 29 G. R. Berdiyrov and K. A. Mahmoud, *Appl. Surf. Sci.*, 2017, **416**, 725–730.
- 30 M. Naguib, V. N. Mochalin, M. W. Barsoum and Y. Gogotsi, *Adv. Mater.*, 2014, **26**, 992–1005.
- 31 M. Naguib, O. Mashtalir, J. Carle, V. Presser, J. Lu, L. Hultman, Y. Gogotsi and M. W. Barsoum, *ACS Nano*, 2012, **6**, 1322–1331.
- 32 C. Peng, X. F. Yang, Y. H. Li, H. Yu, H. J. Wang and F. Peng, *ACS Appl. Mater. Interfaces*, 2016, **8**, 6051–6060.
- 33 Y. J. Li, Z. H. Yin, G. R. Ji, Z. Q. Liang, Y. J. Xue, Y. C. Guo, J. Tian, X. Z. Wang and H. Z. Cui, *Appl. Catal., B*, 2019, **246**, 12–20.
- 34 Y. J. Li, X. T. Deng, J. Tian, Z. Q. Liang and H. Z. Cui, *Appl. Mater. Today*, 2018, **13**, 217–227.
- 35 T. Cai, L. L. Wang, Y. T. Liu, S. Q. Zhang, W. Y. Dong, H. Chen, X. Y. Yi, J. L. Yuan, X. N. Xia, C. B. Liu and S. L. Luo, *Appl. Catal., B*, 2018, **239**, 545–554.
- 36 J. R. Ran, G. P. Gao, F. T. Li, T. Y. Ma, A. J. Du and S. Z. Qiao, *Nat. Commun.*, 2017, **8**, 13907.
- 37 H. G. Yang, C. H. Sun, S. Z. Qiao, J. Zou, G. Liu, S. C. Smith, H. M. Cheng and G. Q. Lu, *Nature*, 2008, **453**, 638–642.
- 38 C. Y. Xing, S. Y. Chen, X. Liang, Q. Liu, M. M. Qu, Q. S. Zou, J. H. Li, H. Tan, L. P. Liu, D. Y. Fan and H. Zhang, *ACS Appl. Mater. Interfaces*, 2018, **33**, 27631–27643.
- 39 S. W. Cao, B. J. Shen, T. Tong, J. W. Fu and J. G. Yu, *Adv. Funct. Mater.*, 2018, **28**, 1800136.
- 40 Q. J. Xiang, K. L. Lv and J. G. Yu, *Appl. Catal., B*, 2010, **96**, 557–564.
- 41 B. Li, Z. Zhao, Q. Zhou, B. Meng, X. Meng and J. Qiu, *Chem.–Eur. J.*, 2014, **20**, 14763–14770.
- 42 F. Tian, Y. P. Zhang, J. Zhang and C. X. Pan, *J. Phys. Chem. C*, 2012, **116**, 7515–7519.
- 43 X. Y. Ma, Q. J. Xiang, Y. L. Liao, T. L. Wen and H. W. Zhang, *Appl. Surf. Sci.*, 2018, **457**, 846–855.
- 44 Y. Xia, Q. Li, K. L. Lv and M. Li, *Appl. Surf. Sci.*, 2017, **398**, 81–88.
- 45 J. Halim, K. M. Cook, M. Naguib, P. Eklund, Y. Gogotsi, J. Rosen and M. W. Barsoum, *Appl. Surf. Sci.*, 2016, **362**, 406–417.
- 46 V. Schier, H. J. Michel and J. Halbritter, *Fresenius J. Anal. Chem.*, 1993, **346**, 227–232.
- 47 T. Sultana, G. L. Georgiev, G. Auner, G. Newaz, H. J. Herfurth and R. Patwa, *Appl. Surf. Sci.*, 2008, **255**, 2569–2573.



- 48 S. A. Shah, T. Habib, H. Gao, P. Gao, W. Sun, M. J. Green and M. Radovic, *Chem. Commun.*, 2017, **53**, 400–403.
- 49 Y. Li, X. H. Feng, Z. X. Lu, H. Yin, F. Liu and Q. J. Xiang, *J. Colloid Interface Sci.*, 2018, **513**, 866–876.
- 50 J. X. Low, L. Y. Zhang, T. Tong, B. J. Shen and J. G. Yu, *J. Catal.*, 2018, **361**, 255–266.
- 51 H. Ghassemi, W. Harlow, O. Mashtalir, M. Beidaghi, M. R. Lukatskaya, Y. Gogotsi and M. L. Taheri, *J. Mater. Chem. A*, 2014, **2**, 14339–14343.
- 52 N. Liu, Y. Chang, Y. L. Feng, Y. Cheng, X. J. Sun, H. Jian, Y. Q. Feng, X. Li and H. Y. Zhang, *ACS Appl. Mater. Interfaces*, 2017, **9**, 5907–5915.
- 53 A. Y. Meng, J. Zhang, D. F. Xu, B. Cheng and J. G. Yu, *Appl. Catal., B*, 2016, **198**, 286–294.
- 54 C. Peng, P. Wei, X. Li, Y. Liu, Y. Cao, H. Wang, H. Yu, F. Peng, L. Zhang, B. Zhang and K. Lv, *Nano energy*, 2018, **53**, 97–107.

