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## The role of oxygen defects in metal oxides for CO<sub>2</sub> reduction

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The abuse of fossil fuels release large amount of CO<sub>2</sub>, causing intense global warming. Using photoreduction and electroreduction to convert CO<sub>2</sub> into highly valuable fuels such as CO and CH<sub>4</sub> can effectively solve this problem. However, due to the limited activity and selectivity, pristine catalyst materials cannot meet the requirements of practical applications, which means that some modifications to these catalysts are necessary. In this review, a series of research reports on oxygen defect engineering have been introduced. First, the methods of preparing oxygen defects by heat treatment, doping, and photoinduction combined with influencing factors in the preparation are introduced. Subsequently, common characterization methods of oxygen defects including EPR, Raman, XPS, EXAFS, and HRTEM are summarized. Finally, the mechanisms of introducing oxygen defects to improve CO<sub>2</sub> reduction are discussed, and include enhancing light absorption, improving CO<sub>2</sub> adsorption and activation, as well as promoting stability of the reaction intermediates. The summary of research on oxygen defects provides guidance for researchers who focus on CO<sub>2</sub> reduction and accelerate the realization of its industrial applications in the future.

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## 1. Introduction

### 1.1 Background

For decades, the consumption of fossil fuels has caused a sharp rise in the level of CO<sub>2</sub> emission, and has resulted in some serious problems related to global warming and climate change.<sup>1,2</sup> Therefore, it is necessary to take some measures to

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limit and reduce CO<sub>2</sub> emission. The capture and conversion of CO<sub>2</sub> into reusable value-added chemicals is considered an effective method to remove excess CO<sub>2</sub> from air.<sup>3,4</sup> Therefore, the photoreduction and electroreduction of CO<sub>2</sub> are considered to be environment-friendly methods for converting CO<sub>2</sub> into combustible chemicals (such as CO and CH<sub>4</sub>), followed by the development of a large number of photocatalysts and electrocatalysts.<sup>5–8</sup> The defects contained in the catalysts play an important role in capturing CO<sub>2</sub> and lowering the activation energy of adsorbed CO<sub>2</sub>.<sup>9–11</sup> In this review, various materials with oxygen defects have been discussed to gain more insight into the correlation between oxygen defects and the activity and selectivity of CO<sub>2</sub> photoreduction and electroreduction. Finally,



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the role of oxygen defect-rich catalysts in the conversion of  $\text{CO}_2$  to reusable fuel is summarized and their future applications are investigated.

## 1.2 Mechanisms of $\text{CO}_2$ photoreduction and electroreduction

Photoreduction and electroreduction processes have been proved to be effective methods for the removal of excessive  $\text{CO}_2$  in air.<sup>12–14</sup> Moreover, there are some differences in the mechanisms between  $\text{CO}_2$  electroreduction and photoreduction.

Since the pioneering work on photoelectrochemical water splitting that was reported by Fujishima and Honda in 1972,<sup>15</sup>  $\text{CO}_2$  photoreduction has gradually developed and studied extensively. As shown in Fig. 1A, the generally accepted process of  $\text{CO}_2$  photoreduction is as follows: first, the photoexcited electrons are excited from the valence band to the conduction band, leaving holes in the valence band. Then, the electrons and holes are separated, which migrate to the surface of the catalysts to reduce  $\text{CO}_2$  to combustible chemicals (such as CO and  $\text{CH}_4$ ).<sup>16</sup> The valence state of carbon in  $\text{CO}_2$  is the highest oxidation state; thus, there are various reduction products of  $\text{CO}_2$ , ranging from CO and  $\text{CH}_4$  to higher hydrocarbons and oxygenates such as  $\text{CH}_3\text{OH}$  and  $\text{HCOOH}$ . On the other hand, the actual process of  $\text{CO}_2$  photoreduction is slightly complicated. Consider the case of CO as the only photocatalytic product. As eqn (1)–(5) show, first, the photogenerated electrons combine with  $\text{CO}_2$  and  $\text{H}^+$  to form the intermediate  $^*\text{COOH}$ . Then,  $^*\text{COOH}$  combines with the photogenerated electrons and  $\text{H}^+$  to release  $\text{H}_2\text{O}$  and to obtain CO absorbed on the catalyst's surface. Finally, CO is desorbed from the catalyst surface. In the meantime, photogenerated holes combine with water to produce oxygen and  $\text{H}^+$ . Due to the complex process of  $\text{CO}_2$  reduction, there are many factors that influence the activity and

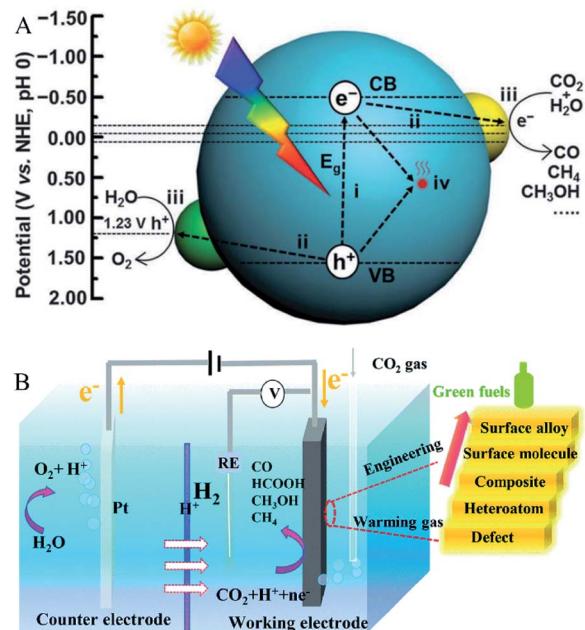


Fig. 1 (A) Schematic illustration of  $\text{CO}_2$  photoreduction. Reproduced with permission: Copyright The Royal Society of Chemistry 2016.<sup>16</sup> (B) schematic illustration of  $\text{CO}_2$  electroreduction. Reproduced with permission: Copyright The Royal Society of Chemistry 2019.<sup>17</sup>

selectivity of  $\text{CO}_2$  photoreduction. Solar light harvesting, charge separation and transportation, as well as surface reactions, are the three main factors, which is recognized as crucial for improving the selectivity and activity of  $\text{CO}_2$  photoreduction. In fact, it is difficult for the pristine catalyst to meet all requirements. While, introducing oxygen defects is an effective method to solve these problems.

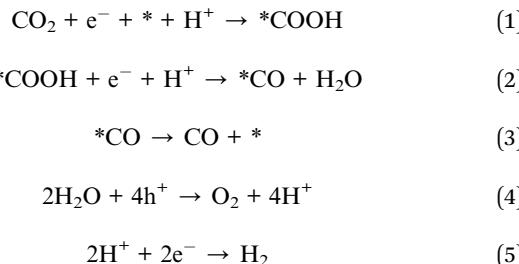


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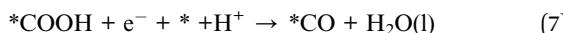
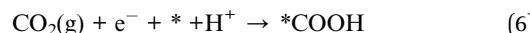


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As for the electric reduction process of  $\text{CO}_2$ , it mainly relies on the electrons generated in the electrode to reduce the  $\text{CO}_2$  adsorbed on the electrocatalyst electrode. As shown in Fig. 1B, electrocatalytic  $\text{CO}_2$  reduction usually occurs in a three-electrode system, including a working electrode loaded with the electrocatalyst for  $\text{CO}_2$  reduction, a reference electrode for determining the potential, and a counter electrode that balances the charge and the current. In the process of  $\text{CO}_2$  electroreduction, electrons transfer to the working electrode and reduce the adsorbed  $\text{CO}_2$  to valuable fuels such as  $\text{HCOOH}$  and  $\text{CO}$ .<sup>17</sup> There are many factors that affect  $\text{CO}_2$  electroreduction, such as the applied potential, the nature of the electrolyte solution, and the electrocatalyst. In this review, the roles of oxygen-defect electrocatalysts applied in  $\text{CO}_2$  electroreduction are mainly discussed. As published in the literature, the process for converting  $\text{CO}_2$  to  $\text{CO}$  involves the steps shown in eqn (6)–(8) and the process of hydrogen evolution involves the steps shown in eqn (9) and (10).<sup>18–20</sup> At the beginning of the reaction,  $\text{CO}_2$  is adsorbed on the electrocatalyst first and then reduced to the intermediate  $* \text{COOH}$ , which is further transformed to  $\text{H}_2\text{O}$  and  $\text{CO}$ , and then desorbed from the electrocatalysts. When the electrocatalyst has the best  $\text{CO}$  binding energy, it can promote further reaction with other reduction products, such as  $\text{CH}_4$  and  $\text{CH}_3\text{OH}$ . As reported in the literature, the two main limitations of  $\text{CO}_2$  electroreduction activity are  $* \text{COOH}$  formation and hydrogen evolution.



### 1.3 The role of oxygen defects

Among the strategies for improving catalytic  $\text{CO}_2$  reduction, introducing oxygen defects into the catalysts has become one of the most likely ways to achieve this goal. The introduction of oxygen defects can enhance the conductivity of the catalysts, and generate a large number of activation sites and adsorption sites on these catalysts, which help to improve their ability for  $\text{CO}_2$  reduction. A large number of studies have reported that

oxygen defect-rich catalysts play a very effective role in  $\text{CO}_2$  reduction.<sup>11,21–24</sup> The formation of oxygen defects can modify the optical absorption by introducing a defective level in order to reduce the bandgap, which further enhances the activity of the catalysts in  $\text{CO}_2$  reduction. Moreover, the formation of the oxygen defects is usually accompanied by the exposure of the metal cation of the metal oxide catalysts and surface atomic structure modification of the catalysts, which may expose more active sites on the catalysts, thus leading to the enhancement in the adsorption and activation of  $\text{CO}_2$  in  $\text{CO}_2$  reduction.<sup>25</sup> According to the studies, the oxygen defects exposed on the surface of the catalysts can offer chemical adsorption sites to reduce the  $\text{CO}_2$  adsorption energy, activate  $\text{CO}_2$  by bending or stretching the  $\text{C}=\text{O}$  bond, and lower the energy barrier by stabilizing the intermediates. It can be anticipated that the introduction of oxygen defects can really improve the  $\text{CO}_2$  reduction efficiency.

Therefore, in order to further intuitively understand the role of oxygen defects in  $\text{CO}_2$  reduction and the corresponding mechanism, we have summarized the different preparation methods of oxygen defect-rich catalysts, the characterization of oxygen defects, and the mechanism involving promotion of  $\text{CO}_2$  reduction by the introduction of oxygen defects.

## 2. The introduction of oxygen defects

Oxygen defect engineering has been widely utilized to adjust the electronic structure of the target catalysts and to improve their performance in  $\text{CO}_2$  reduction. According to the published reports, oxygen defects are usually introduced by methods such as heat treatment, irradiation, or doping modification.

### 2.1 Introducing oxygen defects by heat treatment

Oxygen defects are created by removing oxygen atoms from the catalyst. The common methods for removing oxygen atoms include using a strong reducing agent to reduce the catalyst, reducing the ambient oxygen partial pressure (such as anaerobic condition), and introducing interfacial lattice strain to reduce the Gibbs free energy of oxygen defect formation, so that the reaction balance can move toward the direction of oxygen defects' generation.

There are many methods of introducing oxygen defects through thermal induction operated in vacuum or inert gas environment.<sup>26–28</sup> Wang *et al.*<sup>29</sup> reported a strategy to control the oxygen defect concentration by annealing  $\text{ZnO}$  nanoparticles in different atmospheres with different oxygen partial pressures at 550 °C for 2 h. By comparing the oxygen defect concentration of the  $\text{ZnO}$  samples annealed in oxygen, air, and nitrogen atmospheres, it was found that the order of oxygen defect concentration on the surface of the  $\text{ZnO}$  samples was nitrogen > air > oxygen annealing environments. This result was consistent with the assumption that a lower oxygen defect concentration was obtained in a higher oxygen partial pressure during the heat treatment, demonstrating that the oxygen partial pressure during the heat treatment did have a great effect on the oxygen defect concentration.



In addition to decreasing the partial pressure of oxygen, studies have shown that interfacial lattice strain is another important factor that controls the generation of oxygen defects. Wang *et al.*<sup>30</sup> reported a method for introducing oxygen defects under mild heating conditions by introducing interfacial lattice strain. In this case, as shown in Fig. 2A, interfacial lattice strain was introduced in  $\text{TiO}_2$ (B) by coating transition metal oxides. The results showed that  $\text{TiO}_2$ (B) with coated transition metal oxides had a much higher ratio of oxygen defects than that with uncoated transition metal oxides. As shown in Fig. 2B, it could be calculated that the potential barrier gradually decreased as the interface lattice strain increased. The Gibbs free energy ( $\Delta G$ ) of oxygen defect formation could be defined as a function of epitaxial strain ( $\eta$ ) and temperature ( $T$ ), according to the following relationship, which confirmed the mechanism that the interfacial strain affected the energy barrier of oxygen defect generation (eqn (11)):

$$\Delta G = \Delta E(\eta) + \Delta F(\eta, T) + 0.5 \times \mu_0(T) \quad (11)$$

where  $\mu_0$  is the oxygen chemical potential,  $\Delta E$  represents the free energy at zero temperature, and  $\Delta F$  represents the thermal contribution to free energy. According to the equilibrium, the increase in the interfacial lattice strain led to the decrease in  $\Delta G$ . Therefore, it could be concluded that the interfacial lattice strain introduced by coating made the oxygen defect generation thermodynamically favorable.

The temperature is another significant influencing factor of oxygen defects. According to eqn (11), temperature has a great influence on the Gibbs free energy of oxygen defect formation. The temperature determines whether the oxygen defects can be generated or not. Liu *et al.*<sup>31</sup> reported a method of controlling the concentration of oxygen vacancies (OVs) by annealing  $\text{TiO}_2$  powder in hydrogen gas at different temperatures ranging from 120 °C to 700 °C. In this case, the result of the thermal treatment

demonstrated that the content of oxygen defect increased at first and then decreased with the continuous increase in the annealing temperature. The whole process could be divided into three steps: firstly, hydrogen was physically adsorbed on the surface of  $\text{TiO}_2$  below 300 °C; secondly, when the temperature was higher than 300 °C, the electrons were transferred from the hydrogen atoms to the oxygen atoms, and then the oxygen atoms were removed by the hydrogen atoms in the form of water, leaving oxygen defects on  $\text{TiO}_2$ . As a result, the oxygen defect content increased with the temperature increasing from 300 °C to 450 °C. Thirdly, when the temperature rose up to 450 °C, the electrons were transferred from the oxygen defects to  $\text{Ti}^{4+}$ , and  $\text{Ti}^{3+}$  was formed at the same time, leading to a decrease in the oxygen defects content as the temperature increased over 450 °C. The result of temperature control on the oxygen defect content showed that the OVs formation was determined by thermodynamic control. This means that the oxygen defect content increase for thermodynamically favorable processes and decrease for thermodynamically unfavorable processes.

In addition to introducing oxygen defects under the gas-solid condition by means of heating at high temperature, using a strong reducing agent in the liquid-solid condition can also effectively introduce oxygen defects at a mild temperature. Kim *et al.*<sup>32</sup> reported a strategy for the preparation of  $\text{BaSnO}_{3-x}$  with abundant oxygen defects under the solid-liquid condition at room temperature with  $\text{NaBH}_4$  as the reducing agent. In this case, they could effectively adjust the concentration of oxygen defects by controlling the amount of  $\text{NaBH}_4$ . Compared with the other methods for introducing oxygen defects at high temperature, this mild liquid-solid method could produce oxygen defects without high-temperature processing, which prevented the high temperature from destroying the material structure of the catalysts.

## 2.2 Introducing oxygen defects by doping

The traditional method of generating oxygen defects is heat treatment at a high temperature, which has some problems in keeping the structure and morphology unchanged. The method of introducing oxygen defects by doping heteroatoms is feasible for triggering oxygen defects in order to maintain the stability even under oxygen-rich circumstances. Liu *et al.*<sup>33</sup> reported a facile strategy for introducing oxygen defects by annealing commercial  $\text{MO}_3$  at 300 °C in a  $\text{NH}_3$  atmosphere. The N-doped  $\text{MO}_3$  obtained from the heat treatment had advantages in attaining stable oxygen defects by decreasing the formation energy of the oxygen defects. Zhang *et al.*<sup>34</sup> reported another method of doping boron atom into transition metal oxides to form stable oxygen defects by the pyrolysis approach. The result indicated that boron doping reduced the generation energy of oxygen defect formation in transition metal oxides, which made the generation of oxygen defects thermodynamically favorable. Moreover, substituting the transition metal with the doping cation is also an effective way to introduce oxygen defects. Zhao *et al.*<sup>35</sup> reported a method for precisely controlling the concentration of the oxygen defects by Cu doping under the

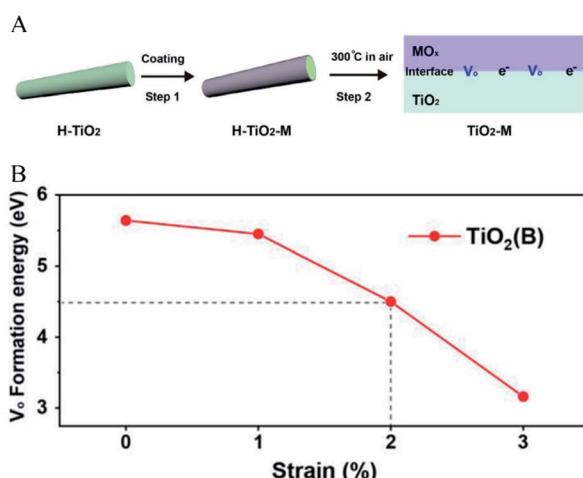


Fig. 2 (A) Illustration of the introduction of interfacial lattice strain by coating transition metal oxides, (B) the relationship between interfacial strain and oxygen defect formation energy. Reproduced with permission: Copyright 2019 Wiley-VCH Verlag GmbH & Co. KGaA, Weinheim.<sup>30</sup>



hydrothermal condition.  $\text{Cu}/\text{TiO}_2$  was synthesized with different amounts of Cu ions, which determined the concentration of the oxygen defects. It was proved that the Ti–O coordination number was decreased as the amount of the doping Cu ions increased, which made it easy for to release oxygen and thereby increase the oxygen defect content. These evidences demonstrate that the concentration of oxygen defects could be controlled by cation substitution. Tong *et al.*<sup>25</sup> also reported an Fe-doping method for the introduction of oxygen defects in  $\text{WO}_{3-x}$ . Fe-doped  $\text{WO}_{3-x}$  was prepared by the hydrothermal method in ethanol solvent, with  $\text{FeCl}_3$  and  $\text{WCl}_6$  used as the precursors. In this case, with the increase in the amount of doped iron, the oxygen defect concentration increased gradually. The low valence state cation substituted the high valence state cation, which would reduce M–O (M is the metal cation) bond and release oxygen; thus, with the increase in cation-doping, the oxygen defect concentration increased gradually. Usually, the activity of the catalysts has an optimal point; thus, that the cation-doping method could control the oxygen defect concentration for obtaining the optimal activity.

### 2.3 Introducing oxygen defects by illumination

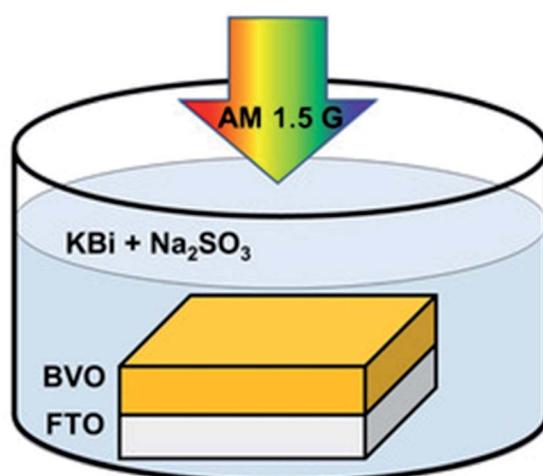
The creation of oxygen defects by photoinduction is a facile method. As for the metal oxide semiconductors, the photoelectrons can be excited by illumination and transferred from oxygen to metal cation, then the oxygen atoms are released and simultaneously form the oxygen defects. Feng *et al.*<sup>36</sup> reported a method for introducing oxygen defects by illumination. The whole process of generating oxygen defects was operated at room temperature by the photoetching method. As shown in Fig. 3,  $\text{BiVO}_4$  with abundant oxygen defects was obtained by immersing the  $\text{BiVO}_4$  photoanode into the electrolyte containing alkali solution and sodium sulfate with illumination. This photoetching method increased the donor density without

introducing a heteroatom and maintained the structure of the  $\text{BiVO}_4$  photoanode. Therefore, the method offered a facile modification strategy to attain oxygen defects. Ye and co-workers<sup>37</sup> have reported a photoinduced oxygen defect engineering method to fabricate black  $\text{In}_{2}\text{O}_{3-x}$  nanosheets. They adopted a 300 W Xe lamp as the irradiation source and introduced oxygen defects in an  $\text{H}_2/\text{CO}$  atmosphere. During the process of introducing oxygen defects,  $\text{h}^+$  was reduced by  $\text{CO}$  or  $\text{H}_2$  and the photogenerated electron reduced  $\text{In}_2\text{O}_3$  to  $\text{In}_2\text{O}_{3-x}$  so as to introduce oxygen defects. Zhang *et al.*<sup>38</sup> have reported a method of introducing oxygen defects in  $\text{BiOCl}$  by illumination.  $\text{BiOCl}$  was irradiated under a 500 W Xe lamp for 5 h, then the white  $\text{BiOCl}$  turned to gray defective  $\text{BiOCl}$ . The introduction of oxygen defects can be effectively controlled by the illumination method. These obtained oxygen defects can effectively widen the photoabsorption range of the catalysts, and improve the  $\text{CO}_2$  adsorption and the catalytic activity. The photoinduction method is an ideal strategy to regenerate oxygen defects in catalysts without introducing a heteroatom and breaking the structure of the catalysts.

## 3. Characterization of the oxygen defects

In the process of studying the role of defective catalysts in  $\text{CO}_2$  reduction, it is usually required to characterize the type and quantity of defects introduced into the catalysts, including the oxygen defects. Common defects characterization methods include high resolution transmission electron microscopy (HRTEM), X-ray photoelectron spectroscopy (XPS), Raman spectroscopy, electron paramagnetic resonance (EPR), X-ray absorption fine structure (XAFS), and photoluminescence (PL); it is usually necessary to combine some of these methods for defect characterization.

Liu *et al.*<sup>39</sup> synthesized OVs-enriched blue  $\text{WO}_{3-x}$  porous nanorods (OBWPN) without N-doping by a simple method. Fig. 4A is the HRTEM image of OBWPN, in which the lattice fringe of 0.375 nm is clearly observed, indexed to the (020) plane of  $\text{WO}_3$ . The amorphous areas of OBWPN marked with a yellow circle proved that its crystallinity had decreased. Comparing the XRD spectra of yellow  $\text{WO}_3$  porous nanorods (YWPN) and OBWPN (Fig. 4B), the two XRD patterns are similar but the peak intensity of OBWPN is relatively weak, further proving that its crystallinity was reduced, which was attributed to the formation of OVs. XPS was employed to confirm the formation of OVs as well. As shown in the O 1s XPS spectra (Fig. 4C), both YWPN and OBWPN contain two typical peaks at 530.3 and 532.6 eV, which represent the W–O bonds and the hydroxyl species of water molecules adsorbed on the surface, respectively. Moreover, the unique peak of OBWPN at 531.2 eV indicates the existence of OVs, proving that the ammonia-assisted reduction strategy introduced OVs in OBWPN. In order to search for more evidences for the existence of OVs, XAFS at W L3-edge and PL spectroscopy were carried out. Fig. 4D is the X-ray absorption near-edge structure (XANES) of YWPN and OBWPN. Compared with YWPN, the peak of OBWPN decreased significantly,



**Photoetching without external bias**

Fig. 3 Illustration of the introduction of oxygen defects by photoetching. Reproduced with permission: Copyright 2020 Wiley-VCH Verlag GmbH & Co. KGaA, Weinheim.<sup>36</sup>



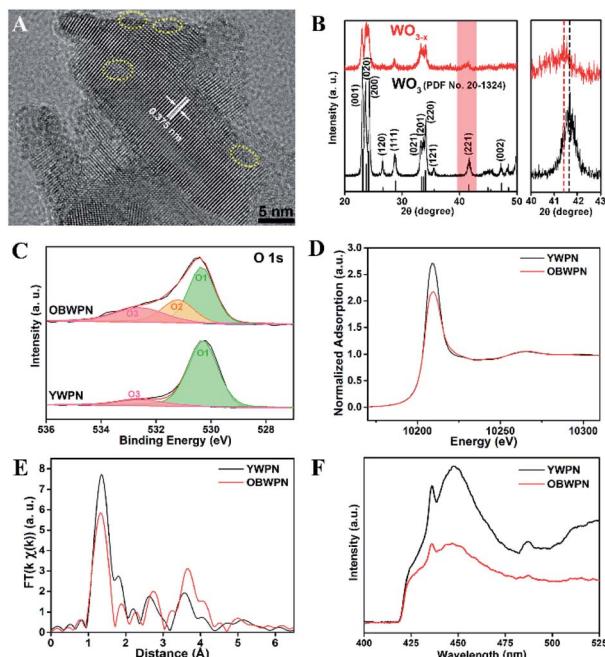


Fig. 4 (A) HRTEM image of OBWPN, (B) XRD patterns of YWPN (black) and OBWPN (red), (C) O 1s XPS spectra of YWPN and OBWPN, (D) normalized W L<sub>3</sub>-edge XANES spectra of YWPN and OBWPN, (E) fourier transform (FT) of k-weighted W EXAFS spectra for YWPN and OBWPN, (F) photoluminescence spectra of YWPN and OBWPN. Reproduced with permission: Copyright 2018 Elsevier Inc.<sup>39</sup>

indicating that OVs caused the decrease in the W valence of OBWPN. Fig. 4E exhibits the fourier transform (FT) of k-weighted W EXAFS spectra for YWPN and OBWPN. The peak at  $\sim 1.33$  Å represents the W–O bond. The W–O peak magnitude of OBWPN was lower than that of YWPN, indicating that the formation of OVs reduced the W–O coordination number in OBWPN. Comparing the fluorescence peaks at 440 nm in the PL spectra of YWPN and OBWPN (Fig. 4F), it can be observed that the peak of OBWPN is weaker, which means that its recombination rate of photogenerated electrons and holes is lower, caused by OVs.

Ahmed *et al.*<sup>40</sup> has synthesized TG (TiO<sub>2</sub> modified by reduced graphene oxide) to reduce the ratio of the relative concentration of the bulk defects to the surface defects so as to improve CO<sub>2</sub> photoreduction. PALS (positron annihilation lifetime spectroscopy) was employed to quantify the relative concentrations of these defects. As observed from Table 1, three positron lifetime components  $\tau_1$ ,  $\tau_2$ ,  $\tau_3$  and their relative intensities  $I_1$ ,  $I_2$ ,  $I_3$  were recorded. The positron lifetimes  $\tau_1$ ,  $\tau_2$ , and  $\tau_3$  are attributed to the bulk defects, surface defects, and micropores as well as the

Table 1 Positron lifetime and relative intensities of TiO<sub>2</sub> and TG.<sup>40</sup> Reproduced with permission: Copyright 2019, Springer Nature

Sample	$\tau_1$ (ns)	$\tau_2$ (ns)	$\tau_3$ (ns)	$I_1$ (%)	$I_2$ (%)	$I_3$ (%)	$I_1/I_2$
TiO <sub>2</sub>	0.1985	0.3937	2.2900	54.4	43.9	1.76	1.23
TG	0.2069	0.3864	2.3130	50.1	48.3	0.67	1.03

annihilation of orthopositronium atoms, respectively. In addition,  $I_1$ ,  $I_2$ ,  $I_3$  are the relative intensity of  $\tau_1$ ,  $\tau_2$ ,  $\tau_3$ , and the ratio of  $I_1/I_2$  is assigned to the relative defect concentration of the bulk to the surface. In this case, the TG had a lower ratio of  $I_1/I_2$ , which indicated that the coupling of graphene reduced the relative concentration of the bulk defects to the surface defects. Considering that TG has a much higher CO<sub>2</sub> reduction activity, this result effectively proved that the surface oxygen defects can improve the photocatalytic activity more effectively.

Karmakar *et al.*<sup>41</sup> explored the role of OVs in enhancing the photoelectrochemical properties of alkali metal (Li, Na, and K) doped ZnO nanorods. The doped alkali metal ions replaced the zinc ions, leaving extra holes, which required electrons to neutralize, thus forming the OVs simultaneously. Through XPS analysis, it could be found that the O 1s spectra of the alkali metal doped ZnO NRs had broad peaks and could be divided into two peaks located at 531.6 eV and 529.9 eV, which represented OVs and O<sup>2-</sup> ions, respectively. The peak intensity of the OVs was related to their concentration. As shown in Fig. 5A, the peak intensities of OVs of Li and Na-ZnO NRs were higher than that of K-ZnO NRs. In this order, the corresponding peak area ratios of their OVs were 0.85, 0.83, and 0.67, respectively, which demonstrated that Li and Na-ZnO NRs contained almost equal OVs concentrations, more than that of the K-ZnO NRs (Fig. 5B). The room temperature PL emission spectra display that the doped ZnO NRs showed significant sub-band gap defect emission near 490 nm, as shown in Fig. 5C, and this deep level emission (DLE) is usually attributed to oxygen-vacancy defects.

In order to enhance the efficiency of photocatalytic CO<sub>2</sub> reduction, Wu *et al.*<sup>42</sup> used high-energy ultraviolet irradiation to produce a large number of OVs on the (001) surface of BiOBr, and characterized the existence of OVs by means of XPS, XAFS, EPR, etc. As shown in Fig. 6A, UV-vis diffuse reflectance spectroscopy was used to confirm the existence of OVs in the BiOBr atomic layer. With the increase in the ultraviolet radiation to the BiOBr atomic layer, a gradually increasing new absorption peak appeared in the visible region, which was due to the formation of OVs. The absorption of visible light by the BiOBr atomic layer reached its maximum value when irradiated with ultraviolet

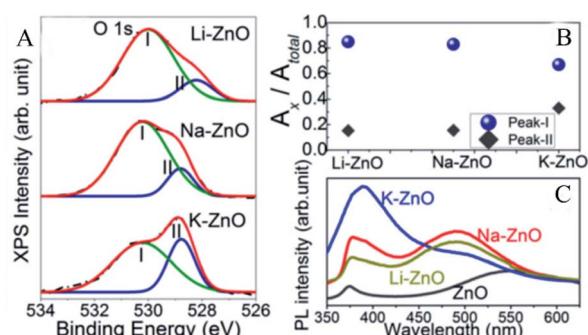


Fig. 5 High-resolution XPS spectra of (A) O 1s for the doped ZnO NRs, (B) the peak area ratios of peak-I and peak-II to the total area, (C) room-temperature PL emission spectra of the alkali metal doped ZnO NRs. Reproduced with permission: Copyright 2018 Wiley-VCH Verlag GmbH & Co. KGaA, Weinheim.<sup>41</sup>



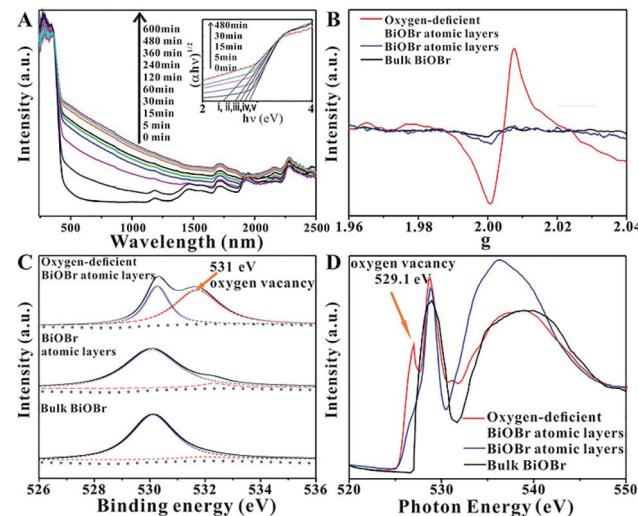


Fig. 6 (A) UV-vis diffuse reflectance spectra for the BiOBr atomic layers after UV irradiation for different periods of time, (B) EPR spectra, (C) high-resolution O 1s XPS spectra, (D) O K-edge XANES spectra for oxygen-deficient BiOBr atomic layers, BiOBr atomic layers, and bulk BiOBr (oxygen-deficient BiOBr atomic layers represent BiOBr atomic layers after UV irradiation for 480 min). Reproduced with permission: Copyright 2018 Wiley-VCH Verlag GmbH & Co. KGaA, Weinheim.<sup>42</sup>

light for 480 minutes, indicating that the oxygen defect concentration reached its maximum value at this time. In response to this situation, EPR technology was used to further prove the existence of the OVs. Fig. 6B shows that the bulk BiOBr and BiOBr atomic layers have no obvious EPR signal, indicating that there are almost no OVs. In contrast, the BiOBr atomic layer, after ultraviolet irradiation for 480 min, showed the strongest EPR signal at  $g = 2.004$ , indicating that ultraviolet irradiation did indeed cause OVs in the BiOBr atomic layer. As observed from Fig. 6C, the high-resolution O 1s XPS spectrum shows that after 480 min of ultraviolet irradiation, a new peak at 531.4 eV appears, which is a peak formed by the oxygen atoms near the OVs. In addition, as shown in Fig. 6D, the same conclusion can be obtained from the O K-edge XANES spectra. The extra peak centered at 529.1 eV can be attributed to the OVs caused by ultraviolet light irradiation.

In order to understand the active site-dependent reaction mechanism, Wang *et al.*<sup>43</sup> used operando spectroscopy methodology to study the Ru/CeO<sub>2</sub> catalytic CO<sub>2</sub> methanation reaction. OVs usually play an important role in the dissociation of oxidative bonds. Therefore, operando Raman was used to detect the changes in the OVs concentration in CeO<sub>2</sub> during the reaction. As shown in Fig. 7A and B, the relative peak intensity ratio between the defect-induced ( $D$ ) mode peak ( $\sim 570$  cm<sup>-1</sup>) and the first-order  $F_{2g}$  peak ( $\sim 460$  cm<sup>-1</sup>) is related to the OV concentration in CeO<sub>2</sub>, and Fig. 7C records the value of  $I_D/I_{F_{2g}}$ . During the reduction, as the temperature increased from 25 °C to 400 °C, the value of  $I_D/I_{F_{2g}}$  increased from 0.02 to 0.52, proving that OVs were gradually generated during the catalyst reduction process. However, when the reaction gases (CO<sub>2</sub> and H<sub>2</sub>) participated, the intensity ratio dropped sharply from 0.43 (at 25 °C) to 0.15 (at 100 °C), and remained low in the temperature range of 200–400 °C.

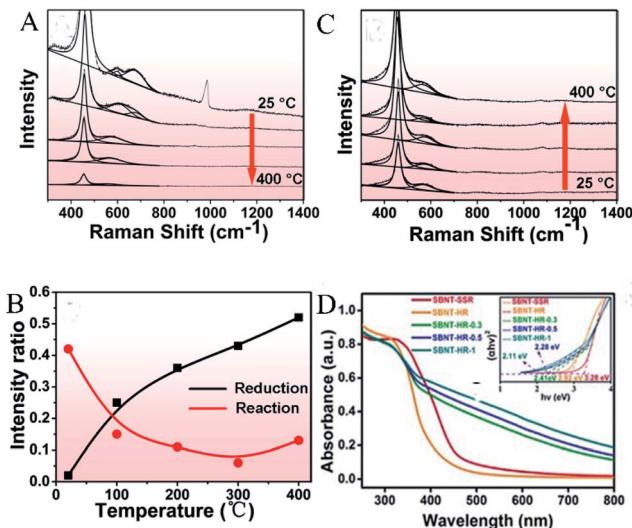


Fig. 7 Operando visible Raman spectra of the Ru/CeO<sub>2</sub> catalyst in (A) the reduction process, (B) the reaction process. The temperature points selected in each process: 25, 100, 200, 300, and 400 °C. (C) The corresponding  $I_D/I_{F_{2g}}$  value as a function of temperature in the reduction and reaction process. Reproduced with permission: Copyright 2020 American Chemical Society.<sup>43</sup> (D) UV-vis diffuse reflectance spectra (DRS), band gap (inset). Reproduced with permission: Copyright 2019 Wiley-VCH Verlag GmbH & Co. KGaA, Weinheim.<sup>44</sup>

## 4. Mechanism of oxygen-defects improving the CO<sub>2</sub> reduction

Due to poor light absorption, slow charge separation, and inefficient surface-active sites, developing effective catalysts for achieving the conversion of CO<sub>2</sub> into useful fuels faces enormous challenges. In recent years, the introduction of oxygen defects has become one of the most effective ways to improve the catalytic efficiency of catalysts.

### 4.1 Increasing the optical absorption

The light absorption capacity of a photocatalyst determines its efficiency of using solar energy. Yu *et al.*<sup>44</sup> reported a simple and controllable *in situ* reduction method for the introduction of oxygen defects on the surface of Aurivillius-phase Sr<sub>2</sub>Bi<sub>2</sub>Nb<sub>2</sub>TiO<sub>12</sub> (SBNT-HR). By the *in situ* treatment of SBNT-HR with 0.3, 0.5, and 1 mL glyoxal, various OVs-rich Sr<sub>2</sub>Bi<sub>2</sub>Nb<sub>2</sub>TiO<sub>12</sub> (SBNT-HR-X,  $X = 0.3, 0.5, 1$ ) were obtained. As shown in Fig. 7D, the light absorption range of SBNT-HR-X extended to 800 nm and as the amount of glyoxal used increased, its light absorption capacity also gradually increased, which might be caused by the increase in the oxygen defect concentration. SBNT-HR (absorption edge of 420 nm) and bulk WO<sub>3</sub> obtained by the high-temperature solid-state reaction (SBNT-SSR, absorption edge of 460 nm) are relatively weak in the visible region. The inset in Fig. 7D shows that the band gaps of SBNT-SSR, SBNT-HR, and SBNT-HR-X ( $X = 0.3, 0.5, 1$ ) are 3.28, 2.92, 2.41, 2.28, and 2.11 eV, respectively. This is because the introduced OVs brings a new defect level to the band gap, thus extending the excitation light range to the entire visible light region.



Similarly, in order to enhance the absorption of visible light, Hou *et al.*<sup>45</sup> prepared sodium tantalate nanocubes with nitrogen and oxygen-vacancy confined (OVs-NaTaON) for efficient photocatalytic CO<sub>2</sub> reduction. Fig. 8A shows the optical absorption of NaTaO<sub>3</sub> and OVs-NaTaON crystals through UV-vis diffuse reflection spectroscopy. The absorption edge of original NaTaO<sub>3</sub> is about 315 nm and its band gap is 3.78 eV (Fig. 8B), while the visible light absorption wavelength range of the OVs-NaTaON nanocrystals is up to 600 nm, corresponding to the band gap of 2.18 eV, as shown in the plots of the transformed Kubelka-Munk function *vs.* the light energy. The greatly reduced band gap proved that the introduction of OVs significantly improved the ability of the catalyst to absorb visible light. The illustration in Fig. 8B shows that the macro color of the catalyst changed from white to orange, which more clearly showed its enhanced visible light absorption capacity.

## 4.2 Increasing the active sites and enhancing CO<sub>2</sub> adsorption

The CO<sub>2</sub> reduction processes includes four steps: adsorption, activation, reaction, and desorption. The introduction of oxygen defects can not only contribute to the enrichment of CO<sub>2</sub> on the catalyst surface but can also serve as an electron capture center to reduce the recombination of electrons and holes as well as to provide more CO<sub>2</sub> activation sites.

Gao *et al.*<sup>46</sup> used the synthesized single-unit-cell layers of anoxic cobalt oxide as an example to construct a model for confining OVs in the atomic layer. As shown in the results of linear sweep voltammetry (LSV) in Fig. 9A, the large cathodic peaks at *ca.* −0.87 V *versus* saturated calomel electrode (SCE) could be ascribed to CO<sub>2</sub> reduction, where the current density of the OVs-rich Co<sub>3</sub>O<sub>4</sub> single-unit-cell layers was approximately twice that of the OVs-poor Co<sub>3</sub>O<sub>4</sub> single-unit-cell layers, indicating that the introduction of OVs was able to improve the ability of catalytic CO<sub>2</sub> reduction. Fig. 9B shows the result that the maximum faradaic efficiency of the OVs-rich and OVs-poor Co<sub>3</sub>O<sub>4</sub> single-unit-cell layers was 87.6% and 67.3% for the production of formate at the potential of −0.87 V *versus* SCE, respectively, further proving the superior selectivity of the former in formate production. The improvement in CO<sub>2</sub> reduction activity and selectivity could be attributed to the involvement of OVs. The increased OVs could provide more active sites for stable reduction intermediates, thereby reducing

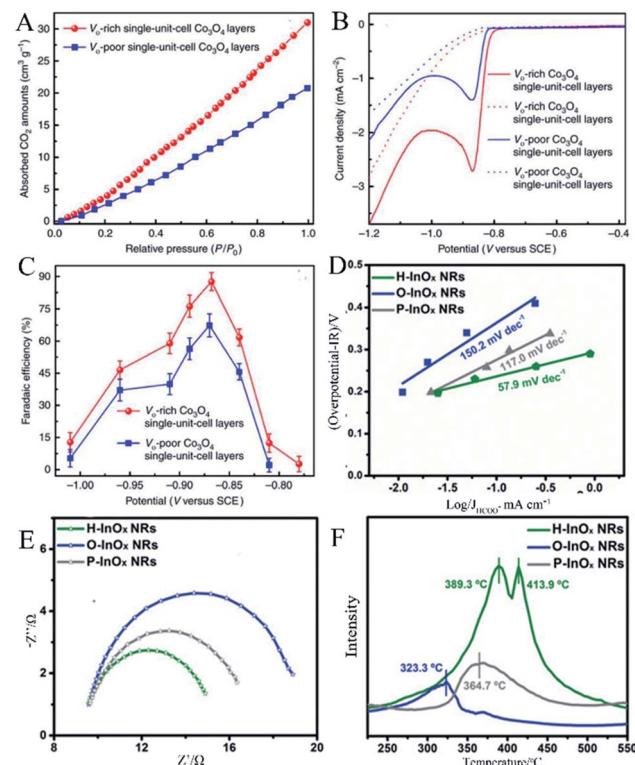


Fig. 9 (A) Linear sweep voltammetric curves in a CO<sub>2</sub>-saturated (solid line) and N<sub>2</sub>-saturated (dashed line) 0.1 M KHCO<sub>3</sub> aqueous solution, (B) faradaic efficiencies of formate at different applied potentials, (C) CO<sub>2</sub> adsorption isotherms. Reproduced with permission: Copyright 2020 Springer Nature Limited.<sup>46</sup> (D) Tafel plots of HCOO<sup>-</sup>, (E) Nyquist plots, (F) CO<sub>2</sub>-TPD profiles of the P-InO<sub>x</sub> NRs, O-InO<sub>x</sub> NRs, and H-InO<sub>x</sub> NRs. Reproduced with permission: Copyright 2019 Wiley-VCH Verlag GmbH & Co. KGaA, Weinheim.<sup>47</sup>

the activation energy barrier. In the measurement of CO<sub>2</sub> adsorption, the result in Fig. 9C exhibits that the OVs-rich Co<sub>3</sub>O<sub>4</sub> single-unit-cell layers had a higher CO<sub>2</sub> adsorption capacity than that of the OVs-poor Co<sub>3</sub>O<sub>4</sub> single-unit-cell layers, indicating that a higher OVs concentration contributed to an increase in CO<sub>2</sub> adsorption.

Zhang *et al.*<sup>47</sup> prepared a series of OV-engineered InO<sub>x</sub> nanoribbons to enhance the absorption and activation of CO<sub>2</sub>, thereby improving the CO<sub>2</sub> electrical reduction ability. Fig. 9D shows the current density of HCOO<sup>-</sup> under various overpotentials and the corresponding fitted Tafel slopes. The Tafel slopes of O-InO<sub>x</sub> NR, P-InO<sub>x</sub> NR, and H-InO<sub>x</sub> NR are 150.2, 117.0, and 57.9 mV dec<sup>-1</sup>, respectively, in which the Tafel slope of H-InO<sub>x</sub> NR was the lowest, demonstrating that the adsorption capacity was enhanced due to the presence of OVs. Nyquist plots were used to evaluate the interface charge transfer resistance, as shown in Fig. 9E. Obviously, the charge transfer resistance of the H-InO<sub>x</sub> NRs was the smallest, which means fastest CO<sub>2</sub> adsorption, charge transfer, and intermediate formation processes. This is because the introduction of OVs enhanced the selectivity of CO<sub>2</sub> reduction. To further investigate the effect of the presence of OVs on the adsorption capacity of CO<sub>2</sub> molecules, temperature programmed CO<sub>2</sub> desorption (CO<sub>2</sub>-

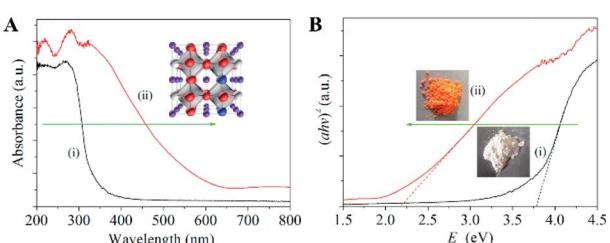


Fig. 8 (A) UV-vis diffuse reflectance spectra of (i) NaTaO<sub>3</sub> and (ii) V<sub>o</sub>-NaTaON crystals. (B) Band-gap energy of (i) NaTaO<sub>3</sub> and (ii) V<sub>o</sub>-NaTaON crystals. Reproduced with permission: Copyright 2016 Elsevier Ltd. All rights reserved.<sup>45</sup>



TPD) was carried out and is shown in Fig. 9F. The CO chemical desorption temperature is arranged in the order O-InO<sub>x</sub> NRs (323.3 °C) < P-InO<sub>x</sub> NRs (364.7 °C) < H-InO<sub>x</sub> NRs (389.3 °C and 413.9 °C), indicating the most capable adsorption of CO<sub>2</sub> by H-InO<sub>x</sub> NRs. The enhanced peaks of the H-InO<sub>x</sub> NRs at higher temperature further showed that the introduction of the OVs could significantly promote the chemisorption capacity of CO<sub>2</sub>, leading to high activity for CO<sub>2</sub> reduction.

## 5. Conclusions

Climate change caused by excessive CO<sub>2</sub> emission has received global attention. Using photoreduction and electric reduction methods, CO<sub>2</sub> can be converted into valuable fuel substances such as CO and CH<sub>4</sub>. However, limited by the lack of activity and selectivity of the original catalytic material, CO<sub>2</sub> reduction still cannot be put into practical use. Therefore, a variety of methods for introducing oxygen defects are listed, including efficient techniques such as heat treatment, doping, and illumination. Various characterization methods of oxygen defects are also summarized. The key point is to perform a qualitative analysis of the oxygen defects contained in the material by combining these characterization methods and to compare the content of oxygen defects generated. Oxygen defects mainly play a role in increasing the light absorption of the catalyst, providing the adsorption and activation sites, effectively improving the stability of the reaction intermediate during the CO<sub>2</sub> reduction process, and maintaining great influence on the catalyst activity and selectivity. A deeper understanding of the structural modification produced by these materials themselves will further promote the industrialization of CO<sub>2</sub> emission reduction, enabling it to be put into practical use on a large scale, and effectively improving actual energy problems and excessive CO<sub>2</sub> emissions.

## Conflicts of interest

There are no conflicts to declare.

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## References

- Y. Fu, D. Sun, Y. Chen, R. Huang, Z. Ding, X. Fu and Z. Li, *Angew. Chem., Int. Ed.*, 2012, **51**, 3364–3367.
- K. Caldeira and M. E. Wickett, *Nature*, 2003, **425**, 365.
- J. Yu, L.-H. Xie, J.-R. Li, Y. Ma, J. M. Seminario and P. B. Balbuena, *Chem. Rev.*, 2017, **117**, 9674–9754.
- D. W. Keith, G. Holmes, D. S. Angelo and K. Heidel, *Joule*, 2018, **2**, 1573–1594.
- Z. W. Seh, J. Kibsgaard, C. F. Dickens, I. Chorkendorff, J. K. Nørskov and T. F. Jaramillo, *Science*, 2017, **355**, eaad4998.
- Y. Pu, Y. Luo, X. Wei, J. Sun, L. Li, W. Zou and L. Dong, *Appl. Catal., B*, 2019, **254**, 580–586.
- H.-P. Liang, A. Acharjya, D. A. Anito, S. Vogl, T.-X. Wang, A. Thomas and B.-H. Han, *ACS Catal.*, 2019, **9**, 3959–3968.
- S. Gao, B. Gu, X. Jiao, Y. Sun, X. Zu, F. Yang, W. Zhu, C. Wang, Z. Feng and B. J. Ye, *J. Am. Chem. Soc.*, 2017, **139**, 3438–3445.
- X. Jiao, Z. Chen, X. Li, Y. Sun, S. Gao, W. Yan, C. Wang, Q. Zhang, Y. Lin and Y. J. Luo, *J. Am. Chem. Soc.*, 2017, **139**, 7586–7594.
- Y. Ji and Y. Luo, *J. Am. Chem. Soc.*, 2016, **138**, 15896–15902.
- N. Guo, H. Xue, A. Bao, Z. Wang, J. Sun, T. Song, X. Ge, W. Zhang, K. Huang and F. He, *Angew. Chem., Int. Ed.*, 2020, **59**, 13778–13784.
- M. Lu, J. Liu, Q. Li, M. Zhang, M. Liu, J. L. Wang, D. Q. Yuan and Y. Q. Lan, *Angew. Chem., Int. Ed.*, 2019, **131**, 12522–12527.
- P. Wu, Y. Li, J.-J. Zheng, N. Hosono, K.-I. Otake, J. Wang, Y. Liu, L. Xia, M. Jiang and S. Sakaki, *Nat. Commun.*, 2019, **10**, 1–8.
- T. Fogeron, P. Retailleau, L. M. Chamoreau, Y. Li and M. Fontecave, *Angew. Chem., Int. Ed.*, 2018, **57**, 17033–17037.
- A. Fujishima and K. Honda, *Nature*, 1972, **238**, 37–38.
- X. Chang, T. Wang and J. Gong, *Energy Environ. Sci.*, 2016, **9**, 2177–2196.
- L. Wang, W. Chen, D. Zhang, Y. Du, R. Amal, S. Qiao, J. Wu and Z. Yin, *Chem. Soc. Rev.*, 2019, **48**, 5310–5349.
- X. Sun, L. Lu, Q. Zhu, C. Wu, D. Yang, C. Chen and B. Han, *Angew. Chem., Int. Ed.*, 2018, **57**, 2427–2431.
- W. Zhu, L. Zhang, S. Liu, A. Li, X. Yuan, C. Hu, G. Zhang, W. Deng, K. Zang and J. Luo, *Angew. Chem., Int. Ed.*, 2020, **59**, 12664–12668.
- F. Y. Gao, S. J. Hu, X. L. Zhang, Y. R. Zheng, H. J. Wang, Z. Z. Niu, P. P. Yang, R. C. Bao, T. Ma and Z. Dang, *Angew. Chem., Int. Ed.*, 2020, **132**, 8784–8790.
- J. Di, C. Zhu, M. Ji, M. Duan, R. Long, C. Yan, K. Gu, J. Xiong, Y. She and J. Xia, *Angew. Chem., Int. Ed.*, 2018, **57**, 14847–14851.
- Z. Geng, X. Kong, W. Chen, H. Su, Y. Liu, F. Cai, G. Wang and J. Zeng, *Angew. Chem., Int. Ed.*, 2018, **57**, 6054–6059.
- X. Jin, C. Lv, X. Zhou, L. Ye, H. Xie, Y. Liu, H. Su, B. Zhang and G. Chen, *ChemSusChem*, 2019, **12**, 2740–2747.
- L. Lu, B. Wang, S. Wang, Z. Shi, S. Yan and Z. Zou, *Adv. Funct. Mater.*, 2017, **27**, 1702447.
- Y. Tong, H. Guo, D. Liu, X. Yan, P. Su, J. Liang, S. Zhou, J. Liu, G. Q. Lu and S. X. Dou, *Angew. Chem., Int. Ed.*, 2020, **59**, 7356–7361.
- M. Vasilopoulou, A. M. Douvas, D. G. Georgiadou, L. C. Palilis, S. Kennou, L. Sygellou, A. Soultati, I. Kostis, G. Papadimitropoulos and D. Davazoglou, *J. Am. Chem. Soc.*, 2012, **134**, 16178–16187.



27 F. C. Calaza, Y. Xu, D. R. Mullins and S. H. Overbury, *J. Am. Chem. Soc.*, 2012, **134**, 18034–18045.

28 T. W. Kim, Y. Ping, G. A. Galli and K.-S. Choi, *Nat. Commun.*, 2015, **6**, 1–10.

29 Y. Wang, B. Wang, Y. Xu, M. Fang, Z. Wu, W. Zhu, J. Hong and C. Li, *J. Chin. Chem. Soc.*, 2017, **64**, 188–194.

30 W. Zhang, L. Cai, S. Cao, L. Qiao, Y. Zeng, Z. Zhu, Z. Lv, H. Xia, L. Zhong and H. Zhang, *Adv. Mater.*, 2019, **31**, 1906156.

31 H. Liu, H. Ma, X. Li, W. Li, M. Wu and X. Bao, *Chemosphere*, 2003, **50**, 39–46.

32 M. Kim, B. Lee, H. Ju, J. Y. Kim, J. Kim and S. W. Lee, *Adv. Mater.*, 2019, **31**, 1903316.

33 K. Liu, W. Zhang, F. Lei, L. Liang, B. Gu, Y. Sun, B. Ye, W. Ni and Y. Xie, *Nano Energy*, 2016, **30**, 810–817.

34 K. Zhang, G. Zhang, J. Qu and H. Liu, *Small*, 2018, **14**, 1802760.

35 Y. Zhao, Y. Zhao, R. Shi, B. Wang, G. I. Waterhouse, L. Z. Wu, C. H. Tung and T. Zhang, *Adv. Mater.*, 2019, **31**, 1806482.

36 S. Feng, T. Wang, B. Liu, C. Hu, L. Li, Z. J. Zhao and J. Gong, *Angew. Chem., Int. Ed.*, 2020, **59**, 2044–2048.

37 Y. Qi, L. Song, S. Ouyang, X. Liang, S. Ning, Q. Zhang and J. Ye, *Adv. Mater.*, 2020, **32**, 1903915.

38 L. Zhang, W. Wang, D. Jiang, E. Gao and S. Sun, *Nano Res.*, 2015, **8**, 821–831.

39 D. Liu, C. Wang, Y. Yu, B.-H. Zhao, W. Wang, Y. Du and B. Zhang, *Chem.*, 2019, **5**, 376–389.

40 G. Ahmed, F. Raziq, M. Hanif, J. Khan, K. S. Munawar, M. Wu, X. Cao and Z. Liu, *Sci. Rep.*, 2019, **9**, 1–8.

41 K. Karmakar, A. Sarkar, K. Mandal and G. G. Khan, *ChemElectroChem*, 2018, **5**, 1147–1152.

42 J. Wu, X. Li, W. Shi, P. Ling, Y. Sun, X. Jiao, S. Gao, L. Liang, J. Xu and W. Yan, *Angew. Chem., Int. Ed.*, 2018, **130**, 8855–8859.

43 F. Wang, S. He, H. Chen, B. Wang, L. Zheng, M. Wei, D. G. Evans and X. Duan, *J. Am. Chem. Soc.*, 2016, **138**, 6298–6305.

44 H. Yu, J. Li, Y. Zhang, S. Yang, K. Han, F. Dong, T. Ma and H. Huang, *Angew. Chem., Int. Ed.*, 2019, **58**, 3880–3884.

45 J. Hou, S. Cao, Y. Wu, F. Liang, L. Ye, Z. Lin and L. Sun, *Nano Energy*, 2016, **30**, 59–68.

46 S. Gao, Z. Sun, W. Liu, X. Jiao, X. Zu, Q. Hu, Y. Sun, T. Yao, W. Zhang and S. Wei, *Nat. Commun.*, 2017, **8**, 1–9.

47 J. Zhang, R. Yin, Q. Shao, T. Zhu and X. Huang, *Angew. Chem., Int. Ed.*, 2019, **58**, 5609–5613.

