



Cite this: *Green Chem.*, 2024, **26**, 1443

# Tailoring metal–support interaction over faceted TiO<sub>2</sub> and copper nanoparticles for electrocatalytic nitrate reduction to ammonia†

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The electrocatalytic nitrate reduction reaction (NO<sub>3</sub><sup>−</sup>RR) provides a sustainable route for ammonia production while mitigating nitrate pollutants in the environment. Metal–support interaction has a significant influence on this electrocatalytic process. However, the mechanism of the facet-dependent metal–support interaction in the NO<sub>3</sub><sup>−</sup>RR is still unknown. Herein, we report the modulation of the metal–support interaction by depositing copper nanoparticles on anatase TiO<sub>2</sub> with different facet exposures, *i.e.*, (001) and (101) facets. The result of copper nanoparticles being deposited on TiO<sub>2</sub> with dominant (101) facet exposure is an enhanced ammonia yield rate of 447.5 μg mg<sub>cat</sub><sup>−1</sup> h<sup>−1</sup> at −0.9 V vs. reversible hydrogen electrode (RHE), which is 4.2 times higher than the pristine TiO<sub>2</sub> counterpart. The strong interaction between copper nanoparticles and TiO<sub>2</sub> with dominant (101) facet exposure contributes to a greater increase in catalytic performance than TiO<sub>2</sub> with dominant (001) facet exposure. The strong interaction leads to electron-deficient copper nanoparticles, efficient electron transfer, and stronger binding of the \*NO<sub>2</sub> intermediate, promoting the hydrogenation process in the NO<sub>3</sub><sup>−</sup> reduction reaction for selective NH<sub>3</sub> synthesis.

Received 8th June 2023,  
Accepted 4th December 2023

DOI: 10.1039/d3gc02018e

rsc.li/greenchem

## Introduction

Ammonia (NH<sub>3</sub>) is an essential feedstock chemical for the fertilizer, pharmaceutical, and nitrogen-containing chemical industries, as well as a potential energy carrier.<sup>1–5</sup> Nevertheless, the production of NH<sub>3</sub> still heavily relies on the Haber–Bosch process, which requires high temperature (400–500 °C) and high pressure (100–200 atm).<sup>6,7</sup> Annually, this process needs an energy supply of up to ~2.5 exajoule and is responsible for 1.4% of carbon dioxide emissions.<sup>8,9</sup> Therefore, the development of a more energy-saving and environmentally friendly process is important.

The electrocatalytic nitrate reduction reaction (NO<sub>3</sub><sup>−</sup>RR) to NH<sub>3</sub> has attracted increasing attention as an alternative method for sustainable NH<sub>3</sub> production since it can be performed under ambient conditions. Moreover, NO<sub>3</sub><sup>−</sup> is highly soluble in water and is known as one of the most widespread

pollutants in the environment, especially in water bodies. NO<sub>3</sub><sup>−</sup> mainly comes from industrial wastewater, liquid nuclear waste, livestock excrement, and chemical fertilizers, with a wide range of concentrations up to 2 mol L<sup>−1</sup>.<sup>10,11</sup> The increased concentration of NO<sub>3</sub><sup>−</sup> could lead to eutrophication, which decreases the oxygen level in water and destroys aquatic ecosystems.<sup>12–16</sup> From a thermodynamic point of view, NO<sub>3</sub><sup>−</sup> has a lower dissociation energy of the N=O bond (204 kJ mol<sup>−1</sup>), which could be more readily reduced with a lower thermodynamic barrier compared to bonds in other nitrogen-containing sources, such as N<sub>2</sub>.<sup>17</sup> Therefore, the process of converting NO<sub>3</sub><sup>−</sup> to NH<sub>3</sub> *via* the NO<sub>3</sub><sup>−</sup>RR could potentially provide a sustainable route for green NH<sub>3</sub> synthesis and address current issues with environmental water pollution.

Several electrocatalysts have been developed for the NO<sub>3</sub><sup>−</sup>RR, such as titanium dioxide (TiO<sub>2</sub>), copper(II) oxide (CuO), copper(I) oxide (Cu<sub>2</sub>O), and cobalt oxide (CoO).<sup>18–21</sup> However, the exhibited performance is typically still limited and remains far below the practical performances as mentioned by the US Department of Energy (300 mA cm<sup>−2</sup> with 90% faradaic efficiency for NH<sub>3</sub>, FE<sub>NH<sub>3</sub></sub>).<sup>22</sup> Continuous studies and investigations are still needed to acquire a better understanding of the catalytic system; hence, the catalytic performance can be further improved. In this regard, exposing specific facets of the catalyst is a promising strategy to modulate catalytic activity as different facet exposures reveal different atomic

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† Electronic supplementary information (ESI) available: Experimental procedure, Fig. S1–S19 and Tables S1 and S2. See DOI: <https://doi.org/10.1039/d3gc02018e>

arrangements and electronic properties.<sup>23–28</sup> Typically, the (001) facet of anatase TiO<sub>2</sub> was reported to have higher reactivity due to the dominance of undercoordinated Ti atoms. An ideal (001) surface of TiO<sub>2</sub> is occupied by five-coordinated Ti (Ti<sub>5c</sub>), *i.e.*, undercoordinated Ti, accounting for 100% of coverage. On the other hand, the (101) surface is occupied by 50% Ti<sub>5c</sub> and 50% six-coordinated Ti (Ti<sub>6c</sub>).<sup>29–31</sup> Therefore, the exposure of more (001) facets can, theoretically, lead to higher activity due to the more reactive Ti<sub>5c</sub> species on the surface. Other factors, such as electronic properties, band energy, or electron trapping states, can also contribute to the different catalytic performances induced by the facet effect.<sup>29,32–36</sup> As reported by Pan *et al.*, TiO<sub>2</sub> with (010) facet exposure (TiO<sub>2</sub>-010) showed the highest catalytic activity, followed by TiO<sub>2</sub>-101 and TiO<sub>2</sub>-001.<sup>29</sup> The performances exhibited were attributed to the cooperative effects of the undercoordinated Ti (Ti<sub>5c</sub>) on the (010) surface of TiO<sub>2</sub>-(010), and a more energetic conduction band (CB) position resulting in stronger reducing electrons in the CB. Besides, Mikrut *et al.* reported that the TiO<sub>2</sub>-(100) catalyst showed the highest activity toward the catalytic oxidation reaction compared to its TiO<sub>2</sub>-(001) and the TiO<sub>2</sub>-(101) counterparts.<sup>32</sup> The superior performance was contributed to by the promoted separation of charge carriers in the TiO<sub>2</sub>-(100) catalyst, which facilitated hole transfer to the active sites. Moreover, Sun *et al.* reported a silver (Ag)-loaded TiO<sub>2</sub> for a catalytic NO<sub>3</sub><sup>−</sup>RR.<sup>33</sup> Benefitting from the Ag nanoparticles as trap centres for electron transfer, the removal efficiency of NO<sub>3</sub><sup>−</sup> reached ~95% with high selectivity of 90% towards N<sub>2</sub> production.

In addition to facet control, metal loading is another effective way to improve the catalytic activity of the NO<sub>3</sub><sup>−</sup>RR. For instance, Kim *et al.* reported the strong metal–support interaction between the Pd–Cu cocatalyst and oxygen-deficient TiO<sub>2</sub>, promoting interfacial electron transfer and enhancing the NO<sub>3</sub><sup>−</sup>RR performance.<sup>37</sup> Nevertheless, NO<sub>2</sub><sup>−</sup> is still the dominant product instead of the desired NH<sub>3</sub> using the reported catalysts. Following this study, Li *et al.* reported using Pd–In supported on an FeO<sub>x</sub> substrate as the catalyst.<sup>38</sup> The intrinsic electron transfer from Pd to the support led to the formation of positively charged Pd sites, which facilitates the adsorption of NO<sub>3</sub><sup>−</sup> for better activation. A similar result was also observed in Cu–Pd-loaded nitrogen-doped carbon (Cu–Pd/CN). The interaction between Cu–Pd nanoparticles and the CN support stimulated the interfacial charge polarization and resulted in the electron-deficient Cu as the active site, which is favourable for NO<sub>3</sub><sup>−</sup> adsorption and activation.<sup>39</sup>

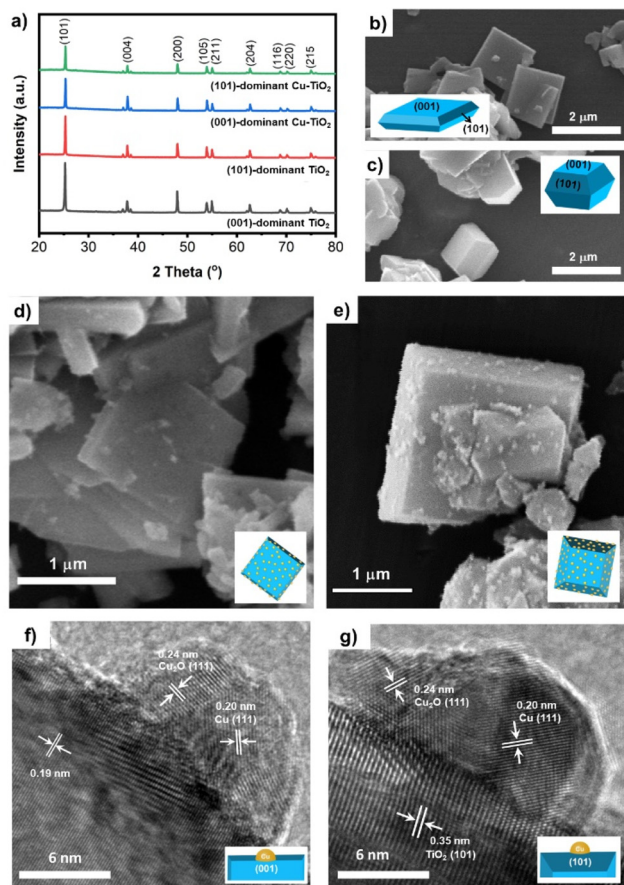
The aforementioned studies indicate the significance of facet exposure and metal–support interaction on the catalytic performance for the NO<sub>3</sub><sup>−</sup>RR. However, the determining factor in the electrocatalytic activity for the faceted TiO<sub>2</sub> is still debatable since many factors could contribute to the apparent catalytic activity.<sup>29,30,33,34</sup> Moreover, the contribution of the exposed TiO<sub>2</sub> facet to the metal–support interaction in metal-loaded faceted TiO<sub>2</sub> in correlation with their electrocatalytic performance for the NO<sub>3</sub><sup>−</sup>RR remains unclear. The deposition of metal on the specific facet of TiO<sub>2</sub> can lead to different

levels of metal–support interaction due to the differing natures of the specific facets of TiO<sub>2</sub>. It can also lead to different reactivities during the catalytic process. In this context, Cu nanoparticles are deposited on the surface of TiO<sub>2</sub> with different facet exposures, *i.e.*, (001) and (101) facets, and catalytic tests toward the NO<sub>3</sub><sup>−</sup>RR as a proof of concept of tailoring the interaction between metal and support are performed. The strong metal–support interaction between the Cu nanoparticles and (101) facets lead to effective electron transfer from the Cu nanoparticles to the TiO<sub>2</sub> support resulting in electron-deficient Cu. In addition to the oxygen vacancies (OVs) in the TiO<sub>2</sub> support, the electron-deficient Cu nanoparticles then serve as active sites for the NO<sub>3</sub><sup>−</sup> adsorption and may also supply the NO<sub>3</sub><sup>−</sup> reactants to the nearby OVs of TiO<sub>2</sub>. As a result, the production of NH<sub>3</sub> by the (101)-dominant Cu–TiO<sub>2</sub> electrocatalyst reaches the highest yield rate of 447.5 μg mg<sub>cat</sub><sup>−1</sup> h<sup>−1</sup> at −0.9 V vs. reversible hydrogen electrode (RHE) among its prepared counterparts, which is 4.2 times higher than that of the pristine (101)-dominant TiO<sub>2</sub> (NH<sub>3</sub> yield: 106.7 μg mg<sub>cat</sub><sup>−1</sup> h<sup>−1</sup>). The increase is more exaggerated compared to the enhancement exhibited by the (001)-dominant Cu–TiO<sub>2</sub> with respect to the pristine (001)-dominant TiO<sub>2</sub>. Moreover, time-dependent experiments and electrochemical analysis disclose that strong metal–support interaction results in a more strongly binding \*NO<sub>2</sub> intermediate, which promotes the electron transfer, facilitating the subsequent hydrogenation process for the selective NH<sub>3</sub> synthesis.

## Results and discussion

### Material characterization

The TiO<sub>2</sub> catalyst with (001) facet or (101) facet as the dominant facet is denoted as (001)-dominant TiO<sub>2</sub> and (101)-dominant TiO<sub>2</sub>, respectively, which was prepared by a hydrothermal process using NaF as a capping agent (details in the Experimental section).<sup>32,40</sup> Subsequently, Cu nanoparticles were evenly deposited on the surface of (001)-dominant TiO<sub>2</sub> and (101)-dominant TiO<sub>2</sub>, resulting in (001)-dominant Cu–TiO<sub>2</sub> and (101)-dominant Cu–TiO<sub>2</sub> (details in the Experimental section). X-ray diffraction (XRD) patterns of the prepared catalysts are presented in Fig. 1a. All XRD peaks can be indexed to the anatase TiO<sub>2</sub> (JCPDS 98-105-4604) without the rutile phase being observed.<sup>31,41</sup> The peak intensities of the (001)-dominant Cu–TiO<sub>2</sub> and (101)-dominant Cu–TiO<sub>2</sub> are slightly lower than those of the pristine (001)-dominant TiO<sub>2</sub> and (101)-dominant TiO<sub>2</sub>, which indicate slight decreases in crystallinity. The decline in the crystallinity can be correlated with the slight reduction of pristine TiO<sub>2</sub> by NaBH<sub>4</sub> during the Cu deposition process.<sup>42</sup> However, no additional peaks from the deposited Cu can be observed in the XRD patterns because of the low amount of deposited Cu and its high dispersion.<sup>43</sup> Inductively coupled plasma optical emission spectroscopy (ICP-OES) analysis shows that the amounts of loaded Cu are 0.86% w/w and 0.82% w/w for the (001)-dominant Cu–TiO<sub>2</sub> and (101)-dominant Cu–TiO<sub>2</sub>, respectively.



**Fig. 1** (a) XRD patterns of the (001)-dominant  $\text{TiO}_2$ , (101)-dominant  $\text{TiO}_2$ , (001)-dominant  $\text{Cu-TiO}_2$ , and (101)-dominant  $\text{Cu-TiO}_2$ . SEM images of (b) the (001)-dominant  $\text{TiO}_2$ , (c) the (101)-dominant  $\text{TiO}_2$ , (d) the (001)-dominant  $\text{Cu-TiO}_2$ , and (e) the (101)-dominant  $\text{Cu-TiO}_2$ . HRTEM images of (f) the (001)-dominant  $\text{Cu-TiO}_2$  with Cu nanoparticles deposited on the (001) facet and (g) the (101)-dominant  $\text{Cu-TiO}_2$  with Cu nanoparticles deposited on the (101) facet.

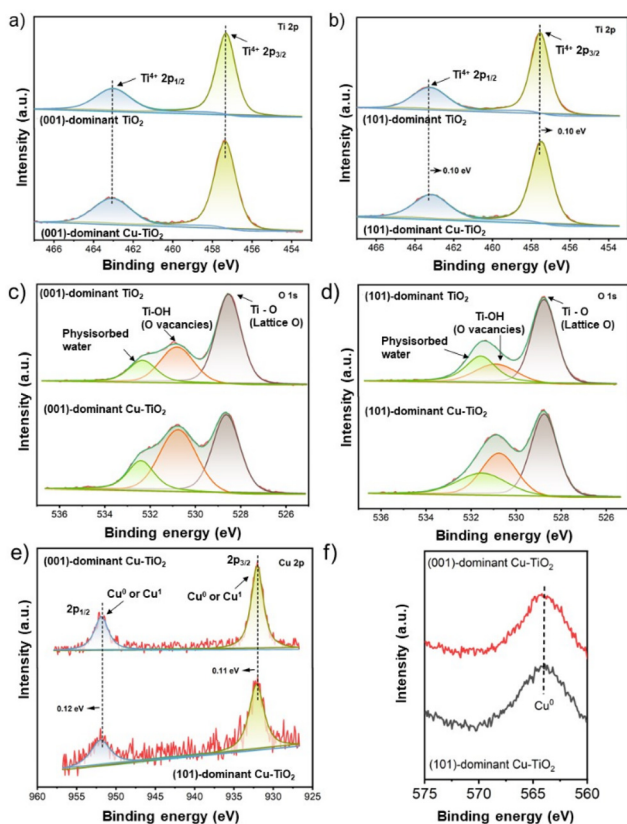
Scanning electron microscopy (SEM) analysis was performed to verify the morphology of the faceted  $\text{TiO}_2$  catalysts. As presented in Fig. 1b and c, the (001)-dominant  $\text{TiO}_2$  catalysts show a flat sheet morphology, while the (101)-dominant  $\text{TiO}_2$  catalysts are truncated octahedral particles. The exposures of (001) facets relative to all facets in the (001)-dominant  $\text{TiO}_2$  and (101)-dominant  $\text{TiO}_2$  were calculated to be  $\sim 70.2\%$  and  $\sim 33.2\%$ , respectively (details in Fig. S1a–c in ESI†). This means that the (001)-dominant  $\text{TiO}_2$  is dominated by (001) facets with an exposure of  $\sim 70.2\%$  from its total facets, while the (101)-dominant  $\text{TiO}_2$  is dominated by (101) facets with an exposure of  $\sim 66.8\%$ . The Cu nanoparticles are deposited on both (001) and (101) facets of the (001)-dominant  $\text{TiO}_2$  and (101)-dominant  $\text{TiO}_2$ , as shown in Fig. 1d and e. The dispersion of the Cu nanoparticles on the surface of the (001)-dominant  $\text{TiO}_2$  and the (101)-dominant  $\text{TiO}_2$  is further verified using transmission electron microscopy (TEM) analysis. The TEM images with the corresponding energy dispersive X-ray (EDX) mapping show the uniform dispersion of the Cu nano-

particles on the (001)-dominant  $\text{Cu-TiO}_2$  and the (101)-dominant  $\text{Cu-TiO}_2$  (Fig. S2a–d and Fig. S3a–d in the ESI†). The average size of the Cu nanoparticle was estimated to be  $6.65 \pm 3.51$  nm and  $6.81 \pm 2.34$  nm for the (001)-dominant  $\text{Cu-TiO}_2$  and the (101)-dominant  $\text{Cu-TiO}_2$ , respectively (Fig. S4a–c and Fig. S5a–d in ESI†). Moreover, the high-resolution TEM (HRTEM) image of the (001)-dominant  $\text{Cu-TiO}_2$  shows a Cu nanoparticle anchored on the surface of the (001) facet of the (001)-dominant  $\text{TiO}_2$  (Fig. 1f). The lattice fringe of 0.19 nm can be attributed to the (200) and (020) crystal planes of anatase  $\text{TiO}_2$ , which are typically correlated with the (001) facet as both (200) and (020) facets have an interfacial angle of  $90^\circ$  between each other on the (001) zone axis diffraction.<sup>31,44</sup> Meanwhile, the attachment of the Cu nanoparticle is identified by the lattice fringe of 0.20 nm coming from the (111) plane of Cu. We note that a lattice fringe of 0.24 nm coming from  $\text{Cu}_2\text{O}$  (111) is also observed,<sup>45,46</sup> which indicates the superficial oxidation of Cu metal under an ambient atmosphere.<sup>47</sup> In addition to the (001) plane, the Cu nanoparticles can also be found on the (101) surface of the (101)-dominant  $\text{TiO}_2$ , as presented in Fig. 1g. The lattice fringe of 0.35 nm refers to the (101) plane of the (101)-dominant  $\text{TiO}_2$ . Similar to the (001)-dominant  $\text{Cu-TiO}_2$ , the crystal planes of both Cu and  $\text{Cu}_2\text{O}$  are also present on the (101)-dominant  $\text{TiO}_2$  catalyst.

The chemical states of all catalysts were investigated using X-ray photoelectron spectroscopy (XPS) analysis. As presented in Fig. 2a, the Ti 2p XPS spectra of the (001)-dominant  $\text{TiO}_2$  and (001)-dominant  $\text{Cu-TiO}_2$  samples show pronounced peaks at 457.3 eV and 462.9 eV, which refer to  $\text{Ti}^{4+}$  2p<sub>3/2</sub> and  $\text{Ti}^{4+}$  2p<sub>1/2</sub>, respectively.<sup>48</sup> The same position of the  $\text{Ti}^{4+}$  2p<sub>3/2</sub> and  $\text{Ti}^{4+}$  2p<sub>1/2</sub> peaks between the pristine (001)-dominant  $\text{TiO}_2$  and (001)-dominant  $\text{Cu-TiO}_2$  indicates that there is no significant change in the  $\text{TiO}_2$  phase of the (001)-dominant  $\text{TiO}_2$  upon depositing Cu nanoparticles. In contrast, the (101)-dominant  $\text{Cu-TiO}_2$  shows a shift of the  $\text{Ti}^{4+}$  2p<sub>3/2</sub> and  $\text{Ti}^{4+}$  2p<sub>1/2</sub> peaks to lower binding energies (457.4 eV and 463.1 eV, respectively) compared to the pristine (101)-dominant  $\text{TiO}_2$  (457.5 eV and 463.2 eV, respectively), as shown in Fig. 2b. The shift of binding energy to lower values suggests the tuned electronic structure of the (101)-dominant  $\text{TiO}_2$  upon Cu deposition.<sup>41,48</sup>

The O 1s XPS spectra are presented in Fig. 2c and d. All catalysts show three deconvoluted peaks at 528.5 eV, 530.8 eV, and 532.3 eV, which can be assigned to the lattice O of  $\text{TiO}_2$ , the O ions associated with bridging Ti–OH (OVs), and physisorbed water, respectively.<sup>41,48</sup> Upon the deposition of the Cu nanoparticles on the surface of the pristine (001)-dominant  $\text{TiO}_2$  and (101)-dominant  $\text{TiO}_2$ , the intensity of OVVs increased, indicating the generation of more OVVs. The formation of more OVVs in the (001)-dominant  $\text{Cu-TiO}_2$  and (101)-dominant  $\text{Cu-TiO}_2$  can be correlated with the Cu deposition process, which involves  $\text{NaBH}_4$  as a reducing agent.<sup>42</sup>

The chemical states analysis of the Cu nanoparticles was performed using argon-etched depth profiling XPS to avoid surface oxidation of Cu in air. As presented in Fig. 2e, two pronounced peaks can be observed at 932.03 eV and 951.85 eV in the (001)-dominant  $\text{Cu-TiO}_2$ , which can be ascribed to the



**Fig. 2** The Ti 2p XPS spectra of (a) the (001)-dominant TiO<sub>2</sub> and (001)-dominant Cu-TiO<sub>2</sub>, and (b) the (101)-dominant TiO<sub>2</sub> and (101)-dominant Cu-TiO<sub>2</sub>. O 1s XPS spectra of (c) the (001)-dominant TiO<sub>2</sub> and (001)-dominant Cu-TiO<sub>2</sub>, and (d) the (101)-dominant TiO<sub>2</sub> and (101)-dominant Cu-TiO<sub>2</sub>. (e) The Cu 2p XPS spectra of the (001)-dominant Cu-TiO<sub>2</sub> and (101)-dominant Cu-TiO<sub>2</sub>. (f) The Cu LMM AES spectra of the (001)-dominant Cu-TiO<sub>2</sub> and (101)-dominant Cu-TiO<sub>2</sub>.

2p<sub>3/2</sub> and 2p<sub>1/2</sub> peaks of Cu<sup>+</sup> or Cu<sup>0</sup>.<sup>41</sup> Further analysis using Cu LMM Auger electron spectroscopy (Fig. 2f) confirmed the presence of Cu<sup>0</sup> species in both the (001)-dominant Cu-TiO<sub>2</sub> and (101)-dominant Cu-TiO<sub>2</sub> as indicated by a single peak at 565.2 eV. The presence of single Cu<sup>0</sup> species was also supported by cyclic voltammetry (CV) measurement over freshly prepared (001)-dominant Cu-TiO<sub>2</sub> and (101)-dominant Cu-TiO<sub>2</sub> samples (Fig. S6 in the ESI†), in which both materials do not show reduction peaks correlated to the reduction of Cu<sup>2+</sup> and Cu<sup>+</sup>, indicating the presence of Cu metal (Cu<sup>0</sup>) on the surface of faceted TiO<sub>2</sub>. However, it is worth noting that in the Cu 2p XPS spectrum of the (101)-dominant Cu-TiO<sub>2</sub> (Fig. 2e), the peaks shifted to the higher binding energies of 932.14 eV and 951.97 eV compared to the (001)-dominant Cu-TiO<sub>2</sub>, suggesting lower electron density in the (101)-dominant Cu-TiO<sub>2</sub> resulting from the electron transfer from the Cu nanoparticles to the (101)-dominant TiO<sub>2</sub> support.<sup>37,41,49</sup>

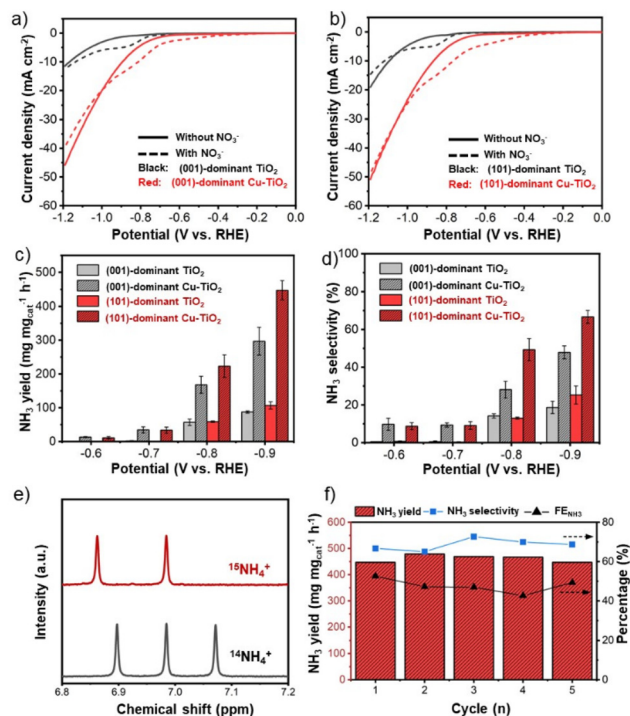
The above XPS results imply that the electron transfer occurred from the Cu nanoparticles to the (101)-dominant TiO<sub>2</sub> support, which decreased the electron density in the Cu nanoparticles and increased the electron density in the OVs

and hence the surrounding Ti atoms. On the other hand, such an electron transfer phenomenon was hardly observed in the (001)-dominant Cu-TiO<sub>2</sub>, which suggests that the Cu nanoparticles have a stronger interaction with (101) facets compared to (001) facets.<sup>50–52</sup> The electron transfer from Cu to TiO<sub>2</sub> correlates with the presence of the OVs on the surface of TiO<sub>2</sub>. In this regard, OVs can serve as electron trapping centres.<sup>53,54</sup> Due to the strong interaction between (101) facets of TiO<sub>2</sub> and the Cu nanoparticles, the electrons are transferred from Cu to TiO<sub>2</sub> and pair with singlet electrons in the OVs near the interface.<sup>55</sup> Additionally, it is also acknowledged that electrons are shallowly trapped in Ti<sub>5c</sub> on the surface of (101) facets, while electrons are also deeply trapped in Ti<sub>6c</sub> on the subsurface of (001) facets.<sup>34</sup> Therefore, the shallow traps on the surface of Ti<sub>5c</sub> in (101) facets can provide a more conducive environment for electron mobility, which result in stronger interactions with the Cu nanoparticles.<sup>34</sup> To sum up, arising from the strong interaction between the Cu nanoparticles and (101)-dominant TiO<sub>2</sub> support, the electrons transfer from Cu nanoparticles to the OVs in (101) facets, leading to electron-deficient Cu nanoparticles.<sup>39,41,49</sup>

### Electrocatalytic NO<sub>3</sub><sup>−</sup>RR evaluation

An electrocatalytic activity test of faceted-TiO<sub>2</sub>-based catalysts was performed in an H-type cell with a typical three-electrode configuration using a Nafion 117 membrane to separate the cathode and the anode chambers. The prepared catalysts were dropcast on carbon paper substrates as the working electrodes (1.0 mg<sub>cat</sub> cm<sup>−2</sup>, details in the Experimental section). Prior to the NO<sub>3</sub><sup>−</sup>RR test, the prepared electrodes were initially examined using linear sweep voltammetry (LSV) from +0.1 V to −1.2 V vs. RHE in 0.5 M Na<sub>2</sub>SO<sub>4</sub> solution with and without the addition of 50 ppm NO<sub>3</sub><sup>−</sup>-N. The LSV curves of both the pristine (001)-dominant TiO<sub>2</sub> and (101)-dominant TiO<sub>2</sub> electrodes showed a slight increase in current density upon the addition of NO<sub>3</sub><sup>−</sup> (Fig. S7 in the ESI†), while the increase in current density was apparently exaggerated with the deposition of Cu nanoparticles (Fig. 3a and b).<sup>19</sup> Besides, the corresponding onset potential of the LSV curves for both faceted-TiO<sub>2</sub> electrodes also shifted to the less negative value (−0.3 V vs. RHE) with the deposition of Cu nanoparticles. Note that the increase of the current density for the (101)-dominant Cu-TiO<sub>2</sub> is greater than that of the (001)-dominant Cu-TiO<sub>2</sub>, suggesting the stronger promoting effect in the NO<sub>3</sub><sup>−</sup>RR upon Cu deposition on the (101) facet of the (101)-dominant TiO<sub>2</sub>.

The electrocatalytic NO<sub>3</sub><sup>−</sup>RR performances were further examined using chronoamperometry (*i*-*t*) tests at different applied potentials in 0.5 M NaSO<sub>4</sub> containing 50 ppm NO<sub>3</sub><sup>−</sup>-N. The concentration levels of NO<sub>3</sub><sup>−</sup>, NH<sub>3</sub>, and NO<sub>2</sub><sup>−</sup> were determined using spectrophotometric methods (Fig. S8a–c in the ESI†).<sup>56</sup> The NO<sub>3</sub><sup>−</sup> conversion rate increased along with the more negative potentials (Fig. S9 in the ESI†). The effect of Cu deposition on the NO<sub>3</sub><sup>−</sup> conversion can be observed clearly at the potentials of −0.8 and −0.9 V vs. RHE, where both the (001)-dominant Cu-TiO<sub>2</sub> and (101)-dominant Cu-TiO<sub>2</sub> show



**Fig. 3** LSV curves of (a) the (001)-dominant TiO<sub>2</sub> and (001)-dominant Cu-TiO<sub>2</sub>, and (b) the (101)-dominant TiO<sub>2</sub> and (101)-dominant Cu-TiO<sub>2</sub> in 0.5 M Na<sub>2</sub>SO<sub>4</sub> without and with the addition of 50 ppm NO<sub>3</sub><sup>-</sup>-N. (c) NH<sub>3</sub> yield and (d) NH<sub>3</sub> selectivity. All catalytic activity experiments were performed in 0.5 M Na<sub>2</sub>SO<sub>4</sub> containing 50 ppm NO<sub>3</sub><sup>-</sup>-N. (e) <sup>1</sup>H NMR spectra of the electrolyte after the electrocatalytic reduction reaction using <sup>15</sup>NO<sub>3</sub><sup>-</sup> and <sup>14</sup>NO<sub>3</sub><sup>-</sup> as the nitrogen sources. The reaction for isotope labelling was performed for the (101)-dominant Cu-TiO<sub>2</sub> at -0.9 V vs. RHE in 0.5 M Na<sub>2</sub>SO<sub>4</sub> containing 50 ppm NO<sub>3</sub><sup>-</sup>-N. (f) Stability test of the (101)-dominant Cu-TiO<sub>2</sub> was performed at -0.9 V vs. RHE in 0.5 M Na<sub>2</sub>SO<sub>4</sub> containing 50 ppm NO<sub>3</sub><sup>-</sup>-N.

higher NO<sub>3</sub><sup>-</sup> conversion rates compared to the pristine TiO<sub>2</sub> counterparts.

Moreover, an obvious trend related to the effect of Cu deposition on the different facets of TiO<sub>2</sub> can be observed in the NH<sub>3</sub> yield rate (Fig. 3c). Within all the applied potentials, both the (001)-dominant Cu-TiO<sub>2</sub> and the (101)-dominant Cu-TiO<sub>2</sub> showed significantly higher NH<sub>3</sub> yield rates compared to the pristine (001)-dominant TiO<sub>2</sub> and (101)-dominant TiO<sub>2</sub>. The activity increased more significantly along with the more negative potentials, reaching the highest value at -0.9 V vs. RHE with NH<sub>3</sub> yields of 296.6 μg mg<sub>cat</sub><sup>-1</sup> h<sup>-1</sup> and 447.5 μg mg<sub>cat</sub><sup>-1</sup> h<sup>-1</sup> for the (001)-dominant Cu-TiO<sub>2</sub> and the (101)-dominant Cu-TiO<sub>2</sub>, respectively. The NH<sub>3</sub> yield rate of the (101)-dominant Cu-TiO<sub>2</sub> is 4.2 times higher than that of the pristine (101)-dominant TiO<sub>2</sub> (106.7 μg mg<sub>cat</sub><sup>-1</sup> h<sup>-1</sup>), while the NH<sub>3</sub> yield of the (001)-dominant Cu-TiO<sub>2</sub> is only 3.3 times higher than the pristine counterpart (87.9 μg mg<sub>cat</sub><sup>-1</sup> h<sup>-1</sup>). Therefore, the (101)-dominant Cu-TiO<sub>2</sub> exhibits an exaggerated NH<sub>3</sub> yield increase compared to the (001)-dominant Cu-TiO<sub>2</sub>. Similar trends can also be observed in the corresponding NH<sub>3</sub> selectivity (Fig. 3d) and FE<sub>NH<sub>3</sub></sub> (Fig. S10 in the ESI<sup>†</sup>). The highest NH<sub>3</sub>

selectivity value was achieved at -0.9 V with values of 66.7% and 47.8% for the (101)-dominant Cu-TiO<sub>2</sub> and the (001)-dominant Cu-TiO<sub>2</sub>, respectively. Additionally, it is worth noting that FE<sub>NH<sub>3</sub></sub> shows a volcano plot with the highest values being reached at -0.8 V vs. RHE with values of 67.1% and 47.0% for the (101)-dominant Cu-TiO<sub>2</sub> and the (001)-dominant Cu-TiO<sub>2</sub>, respectively. Note that the NH<sub>3</sub> selectivity value was calculated based on the concentration change of NO<sub>3</sub><sup>-</sup> only with the corresponding products, while the FE was calculated based on the total current regardless of the reactions (details in the ESI<sup>†</sup>). Therefore, the calculation of the NH<sub>3</sub> selectivity value excluded the contribution of the hydrogen evolution reaction (HER), while the FE<sub>NH<sub>3</sub></sub> took it into consideration.<sup>19,57</sup> The decrease of the FE<sub>NH<sub>3</sub></sub> for all electrodes at -0.9 V vs. RHE is attributed to the intensified HER, as evidenced by the significant increase of the FE<sub>H<sub>2</sub></sub> (Fig. S11 in the ESI<sup>†</sup>).<sup>14,19</sup> Accordingly, both the NO<sub>2</sub><sup>-</sup> selectivity and the FE<sub>NO<sub>2</sub></sub> also decreased at -0.9 V vs. RHE (Fig. S12 and Fig. S13 in the ESI<sup>†</sup>). Another possible byproduct, *i.e.*, hydrazine (N<sub>2</sub>H<sub>4</sub>), was also detected using the Watt and Chrisp method (detailed in the Experimental section in the ESI<sup>†</sup>). In all catalytic reactions, no hydrazine (N<sub>2</sub>H<sub>4</sub>) was detected as presented in Fig. S14a-c in the ESI<sup>†</sup> excluding the possibility of N<sub>2</sub>H<sub>4</sub> as a byproduct. We note that the greatest performance exhibited by the (101)-dominant Cu-TiO<sub>2</sub> catalyst in this work is, in fact, on a par with previous studies reported in the literature, specifically for Ti-based and Cu<sup>0</sup>-based catalysts in neutral media (Table S1 in the ESI<sup>†</sup>). However, it is worth noting that the exhibited performances in this work resulted from lower external potentials (-0.9 V vs. RHE) compared to the other Ti-based catalysts (-1.0 to -1.6 V vs. RHE), which is also one of the important parameters for evaluating the electrocatalytic performance.<sup>58,59</sup>

Control experiments were then performed to verify the origin of the produced NH<sub>3</sub>. As presented in Fig. 3e, the electrolyte taken from the electrocatalytic reduction of Na<sup>15</sup>NO<sub>3</sub> shows typical doublet peaks at δ = 6.86 and δ = 6.98 ppm, while the triplet peak was observed in the spectra when employing Na<sup>14</sup>NO<sub>3</sub>. These <sup>15</sup>N isotope-labelling results strongly confirm that the produced NH<sub>3</sub> originated from the electroreduction of NO<sub>3</sub><sup>-</sup>.<sup>19,56</sup> Moreover, a negligible amount of NH<sub>3</sub> produced in the absence of NO<sub>3</sub><sup>-</sup> and at the open circuit potential (OCP) further supports the results from <sup>15</sup>N isotope-labelling experiments (Fig. S15 in the ESI<sup>†</sup>). No accumulation of NH<sub>3</sub> was observed upon exposing the electrolyte to the air, excluding possible NH<sub>3</sub> contamination from the environment (Fig. S16 in the ESI<sup>†</sup>).

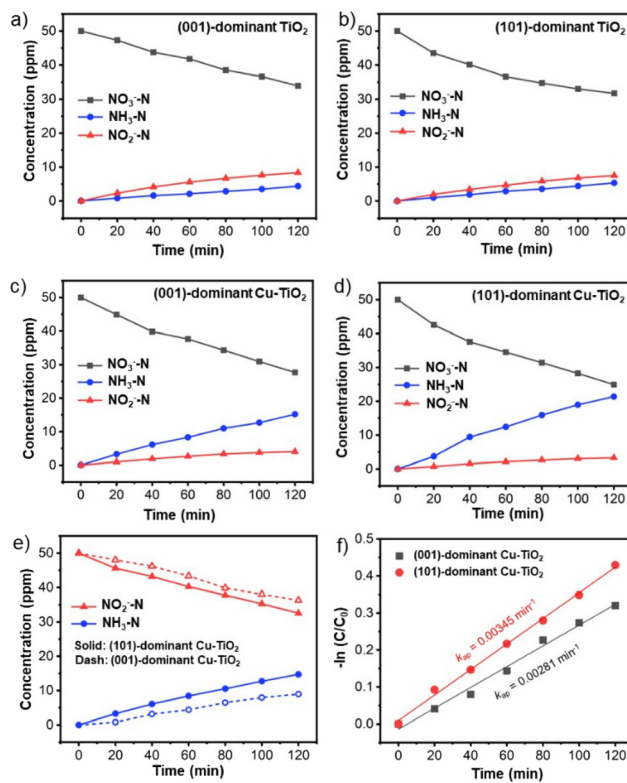
The exaggerated improvement in the NH<sub>3</sub> yield rate and NH<sub>3</sub> selectivity can be correlated with the strong metal-support interaction exhibited by the (101)-dominant Cu-TiO<sub>2</sub> catalyst. On the other hand, we note that the concentration of OV also increases in both the (001)-dominant TiO<sub>2</sub> and (101)-dominant TiO<sub>2</sub> upon Cu deposition as indicated by the O 1s XPS spectra in Fig. 2c and d. One of the oxygen (O) atoms in NO<sub>3</sub><sup>-</sup> can occupy the OV and coordinate with the surrounding Ti<sup>3+</sup> of TiO<sub>2</sub>, which weakens the N-O bonding for better activation.<sup>18</sup> Therefore, to investigate the contribution of OVs, we

also performed the electrocatalytic test on the reduced-(001)-dominant TiO<sub>2</sub> and reduced-(101)-dominant TiO<sub>2</sub> prepared by the NaBH<sub>4</sub> reduction method on the pristine faceted TiO<sub>2</sub> without the addition of Cu salts. As verified by the electron paramagnetic resonance (EPR) spectra of the pristine (101)-dominant TiO<sub>2</sub> and the reduced (101)-dominant TiO<sub>2</sub> (Fig. S17 in the ESI†), the reduced (101)-dominant TiO<sub>2</sub> shows a slightly higher peak intensity at  $g = 2.003$ , indicating a higher concentration of OV's upon implementing the NaBH<sub>4</sub> reduction process.<sup>48</sup> Moreover, as presented in Fig. S18a–d in the ESI,† the reduced-(001)-dominant TiO<sub>2</sub> and the reduced-(101)-dominant TiO<sub>2</sub> also show slight increases in NH<sub>3</sub> yield rate and selectivity, suggesting the benefits of OV's generation in the NO<sub>3</sub><sup>−</sup>RR to a certain degree.<sup>18</sup> However, the improved performances by the reduced-faceted-TiO<sub>2</sub> electrodes are much less significant compared to the enhancements exhibited by the Cu-loaded faceted TiO<sub>2</sub> electrodes.

To further investigate the contribution from the Cu nanoparticles, which serve as the co-catalyst in the Cu-loaded TiO<sub>2</sub> electrode, we also performed electrocatalytic tests over Cu nanoparticles prepared using the NaBH<sub>4</sub> reduction method. For this purpose, we deposited an equal amount of Cu nanoparticles contained in the (001)-dominant Cu-TiO<sub>2</sub> and (101)-dominant Cu-TiO<sub>2</sub> electrodes. As presented in Fig. S19 in the ESI,† Cu nanoparticles show activity for NO<sub>3</sub><sup>−</sup> reduction to NH<sub>3</sub>.<sup>60</sup> NH<sub>3</sub> yield rates increase along with the more negative potentials. However, the magnitude of the NH<sub>3</sub> yield rate is much inferior compared to that of the Cu-loaded faceted TiO<sub>2</sub> catalysts. These control experiments provide evidence that significant enhancement of the NH<sub>3</sub> yield rate, selectivity, and FE<sub>NH<sub>3</sub></sub> over the (001)-dominant Cu-TiO<sub>2</sub> and the (101)-dominant Cu-TiO<sub>2</sub> is mainly determined by the interaction between the Cu nanoparticles and the faceted TiO<sub>2</sub>. Moreover, the stronger metal-support interaction between the Cu nanoparticles and the (101) facets of TiO<sub>2</sub> further enhance the performance. The presumption can be derived from the fact that the increase in NH<sub>3</sub> yield rate and selectivity is exaggerated for the (101)-dominant Cu-TiO<sub>2</sub> compared to the (001)-dominant Cu-TiO<sub>2</sub>. Moreover, the (101)-dominant Cu-TiO<sub>2</sub> shows a stable NH<sub>3</sub> yield rate, NH<sub>3</sub> selectivity, and FE<sub>NH<sub>3</sub></sub> in five consecutive reactions (Fig. 3f), suggesting the appreciable stability of the designed system for the NO<sub>3</sub><sup>−</sup>RR.

### Mechanism study

Time-dependent experiments were performed to understand the kinetics and reaction mechanism of the NO<sub>3</sub><sup>−</sup>RR (Fig. 4a–d). For all prepared catalysts, the NO<sub>3</sub><sup>−</sup> concentration decreases over time accompanied by the increase in NH<sub>3</sub> and NO<sub>2</sub><sup>−</sup> concentrations. The apparent reduction rate constant values of NO<sub>3</sub><sup>−</sup> calculated based on the Langmuir–Hinshelwood model for the (101)-dominant Cu-TiO<sub>2</sub> ( $k_{ap} = 0.00551 \text{ min}^{-1}$ ) and the (001)-dominant Cu-TiO<sub>2</sub> ( $k_{ap} = 0.00476 \text{ min}^{-1}$ ) are higher than those of the pristine (101)-dominant TiO<sub>2</sub> ( $k_{ap} = 0.00368 \text{ min}^{-1}$ ) and (001)-dominant TiO<sub>2</sub> ( $k_{ap} = 0.00322 \text{ min}^{-1}$ ) counterparts, suggesting the positive role of Cu deposition in promoting the NO<sub>3</sub><sup>−</sup>RR (Fig. S20a and b in the ESI†).<sup>38,61,62</sup>



**Fig. 4** NO<sub>3</sub><sup>−</sup>, NH<sub>3</sub>, and NO<sub>2</sub><sup>−</sup> concentration changes over (a) the (001)-dominant TiO<sub>2</sub>, (b) the (101)-dominant TiO<sub>2</sub>, (c) the (001)-dominant Cu-TiO<sub>2</sub>, and (d) the (101)-dominant Cu-TiO<sub>2</sub>. All experiments were performed at  $-0.9 \text{ V}$  vs. RHE in  $0.5 \text{ M Na}_2\text{SO}_4$  containing  $50 \text{ ppm}$  of NO<sub>3</sub><sup>−</sup>-N for 2 h.  $\epsilon$  NO<sub>2</sub><sup>−</sup> and NH<sub>3</sub> concentration changes over the (001)-dominant Cu-TiO<sub>2</sub> and the (101)-dominant Cu-TiO<sub>2</sub>, and (f) the corresponding Langmuir–Hinshelwood plots. All the experiments were performed at  $-0.9 \text{ V}$  vs. RHE in  $0.5 \text{ M Na}_2\text{SO}_4$  containing  $50 \text{ ppm}$  of NO<sub>2</sub><sup>−</sup>-N for 2 h.

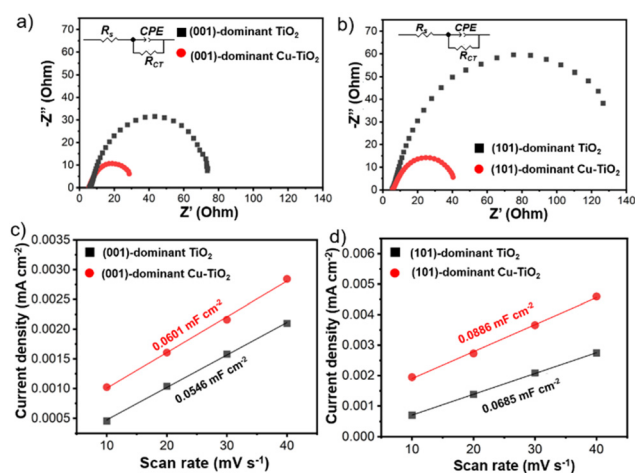
The apparent activation energy ( $E_a$ ) also decreases upon Cu loading on the surface of faceted TiO<sub>2</sub>. The  $E_a$  for the (101)-dominant Cu-TiO<sub>2</sub> is significantly lower ( $E_a = 4.61 \text{ kJ mol}^{-1}$ ) than that of the pristine (101)-dominant TiO<sub>2</sub> ( $E_a = 15.79 \text{ kJ mol}^{-1}$ ). The same trend can be observed in the (001)-dominant Cu-TiO<sub>2</sub> ( $E_a = 13.90 \text{ kJ mol}^{-1}$ ) and the (001)-dominant TiO<sub>2</sub> ( $E_a = 24.03 \text{ kJ mol}^{-1}$ ) (Fig. S21a and b in the ESI†).<sup>63,64</sup> It is noteworthy that among the four samples, the (101)-faceted TiO<sub>2</sub>-based catalysts typically show lower apparent  $E_a$  values than the (001)-faceted TiO<sub>2</sub>-based catalysts, indicating the advantage of (101) facet exposure and Cu loading on these facets for reducing the  $E_a$ . These  $E_a$  values agree well with the Tafel plots (Fig. S22 in the ESI†), in which the (101)-dominant Cu-TiO<sub>2</sub> shows a lower Tafel slope ( $178.7 \text{ mV dec}^{-1}$ ) than the (001)-dominant Cu-TiO<sub>2</sub> ( $203.7 \text{ mV}$ ), indicating the higher kinetics rate of the former for the electrocatalytic NO<sub>3</sub><sup>−</sup>RR.<sup>65,66</sup> These apparent  $E_a$  values and the Tafel slopes are in good agreement with the reduction rate (Fig. 4a–d and Fig. S20a, b in the ESI†), as mentioned before. In addition to the kinetics information, the higher reduction rate of the (101)-dominant Cu-TiO<sub>2</sub> suggests a higher adsorption strength between the NO<sub>3</sub><sup>−</sup> ions with the catalyst surface. This assumption can be rationalized

based on the presence of the electron-deficient Cu nanoparticles in the (101)-dominant Cu-TiO<sub>2</sub>, which are more conducive to attracting the negatively charged NO<sub>3</sub><sup>-</sup> ions.<sup>38,39</sup> This attraction could also lead to the accumulation of NO<sub>3</sub><sup>-</sup> near the catalyst surface, which facilitates the contact between the NO<sub>3</sub><sup>-</sup> ions and the surface OVs in the surrounding Cu nanoparticles.<sup>39</sup>

The influence of the strong metal-support interaction on the catalysis process can be observed by the product distribution, specifically the evolution of NH<sub>3</sub> and NO<sub>2</sub><sup>-</sup> as the two major products. As shown in Fig. 4a and b, the (001)-dominant TiO<sub>2</sub> and the (101)-dominant TiO<sub>2</sub> electrodes show that NO<sub>2</sub><sup>-</sup> is the dominant product at -0.9 V vs. RHE. On the other hand, upon the deposition of Cu on the faceted TiO<sub>2</sub>, NH<sub>3</sub> emerges as the dominant product from the beginning of the reaction (Fig. 4c and d). Notably, the increase in the NH<sub>3</sub> concentration of the (101)-dominant Cu-TiO<sub>2</sub> is much higher than that of the (001)-dominant Cu-TiO<sub>2</sub>, which is in good agreement with the trends of NH<sub>3</sub> yield rate and NH<sub>3</sub> selectivity in the catalytic test. The time-dependent experiments in a prolonged reaction period show similar trends for the concentration changes of the NO<sub>3</sub><sup>-</sup> reactant and the reduction products (Fig. S23a and b in the ESI†). The greater increase in NH<sub>3</sub> production of the (101)-dominant Cu-TiO<sub>2</sub> indicates the benefit of the strong metal-support interaction between the Cu nanoparticles and (101) facets of TiO<sub>2</sub> in promoting the selective NO<sub>3</sub><sup>-</sup>RR to NH<sub>3</sub>.

We note that the detection of NO<sub>2</sub><sup>-</sup> in the electrolyte during the NO<sub>3</sub><sup>-</sup>RR indicates that the production rate of NO<sub>3</sub><sup>-</sup> to \*NO<sub>2</sub> is faster than that of NO<sub>2</sub><sup>-</sup> to NH<sub>3</sub>, and thus leads to the desorption of \*NO<sub>2</sub> from the catalyst surface into the electrolyte.<sup>60</sup> The detection of higher amounts of NO<sub>2</sub><sup>-</sup> in the electrolyte may also indicate that the binding strength of \*NO<sub>2</sub> to the catalyst surface is weaker, hence it can easily be desorbed from the catalyst surface into the electrolyte. Therefore, to further investigate the role of the \*NO<sub>2</sub> intermediate in the NO<sub>3</sub><sup>-</sup>RR, time-dependent NO<sub>2</sub><sup>-</sup> reduction reactions (NO<sub>2</sub><sup>-</sup>RRs) over the (001)-dominant Cu-TiO<sub>2</sub> and the (101)-dominant Cu-TiO<sub>2</sub> catalysts were performed. As presented in Fig. 4e, the decrease in the NO<sub>2</sub><sup>-</sup> concentration using the (001)-dominant Cu-TiO<sub>2</sub> is slower than that using the (101)-dominant Cu-TiO<sub>2</sub>. The corresponding apparent rate constant values based on the Langmuir-Hinshelwood model of the (101)-dominant Cu-TiO<sub>2</sub> and the (001)-dominant Cu-TiO<sub>2</sub> are 0.00345 min<sup>-1</sup> and 0.00281 min<sup>-1</sup> (Fig. 4f), respectively.<sup>61,67</sup> The higher rate constant of NO<sub>2</sub><sup>-</sup> reduction may indicate the higher binding strength between NO<sub>2</sub><sup>-</sup> and the surface of the (101)-dominant Cu-TiO<sub>2</sub> when forming \*NO<sub>2</sub>. These results agree well with the lower amount of NO<sub>2</sub><sup>-</sup> produced using the (101)-dominant Cu-TiO<sub>2</sub> compared with that produced using the (001)-dominant Cu-TiO<sub>2</sub> in the NO<sub>3</sub><sup>-</sup>RR (Fig. S24 in the ESI†). Based on these findings, the exaggerated enhancement in the NH<sub>3</sub> production by the (101)-dominant Cu-TiO<sub>2</sub> can also be attributed to the stronger binding strength of \*NO<sub>2</sub>, which suppresses the desorption of \*NO<sub>2</sub> into the electrolyte and subsequently promotes the subsequent hydrogenation process.

The intrinsic electrochemical properties of the prepared electrodes are important factors in influencing catalytic performance. Therefore, electrochemical impedance spectroscopy (EIS), electrochemical active surface area (ECSA) analysis, and Mott-Schottky analysis were performed for the pristine and Cu-deposited TiO<sub>2</sub> electrodes. The Nyquist plots of the Cu-deposited TiO<sub>2</sub> electrodes show smaller arc radii than the pristine counterparts (Fig. 5a and b). The series resistance (*R<sub>s</sub>*), which is correlated with the resistance from the solution, is comparable among all the electrodes (Table S2 in the ESI†).<sup>61,68,69</sup> On the other hand, the charge transfer resistance (*R<sub>CT</sub>*) decreases significantly upon Cu deposition. The *R<sub>CT</sub>* value of the (001)-dominant Cu-TiO<sub>2</sub> (27.20 Ω) is 2.7 times lower than that of the pristine (001)-dominant TiO<sub>2</sub> (74.38 Ω). Meanwhile, the (101)-dominant Cu-TiO<sub>2</sub> shows a more significant decrease in the *R<sub>CT</sub>* value (40.28 Ω), which is 3.8 times lower than that of the pristine (101)-dominant TiO<sub>2</sub> (152.50 Ω). The more efficient electron transfer as evidenced by the higher decrease in the *R<sub>CT</sub>* value on the (101)-dominant Cu-TiO<sub>2</sub> than on the (001)-dominant Cu-TiO<sub>2</sub> is likely contributed to by the stronger interaction between Cu nanoparticles and (101) facets of TiO<sub>2</sub> in the (101)-dominant Cu-TiO<sub>2</sub>.<sup>41,70</sup> The ECSA was further investigated by measuring the electrochemical double-layer capacitance (*C<sub>dl</sub>*), which is proportional to the ECSA. The linear fitting of the charging current density with various scan rates (Fig. 5c and d and Fig. S25a-d in the ESI†) reveals that the *C<sub>dl</sub>* values of the (001)-dominant Cu-TiO<sub>2</sub> (*C<sub>dl</sub>* = 0.0601 mF cm<sup>-2</sup>) and the (101)-dominant Cu-TiO<sub>2</sub> (*C<sub>dl</sub>* = 0.0886 mF cm<sup>-2</sup>) are higher than those of the pristine (001)-dominant TiO<sub>2</sub> (*C<sub>dl</sub>* = 0.0546 mF cm<sup>-2</sup>) and the (101)-dominant TiO<sub>2</sub> (*C<sub>dl</sub>* = 0.0685 mF cm<sup>-2</sup>), suggesting that more active sites are present in the Cu-deposited TiO<sub>2</sub> catalysts.<sup>41</sup> Moreover, the Mott-Schottky plots show positive slopes for all the electrodes, indi-



**Fig. 5** Nyquist plots of (a) the (001)-dominant TiO<sub>2</sub> and (001)-dominant Cu-TiO<sub>2</sub> and (b) the (101)-dominant TiO<sub>2</sub> and (101)-dominant Cu-TiO<sub>2</sub>. The insets are the equivalent electrical circuits. Charging current density plotted against the scan rate at 0.655 V vs. RHE for (c) the (001)-dominant TiO<sub>2</sub> and (001)-dominant Cu-TiO<sub>2</sub>, and (d) the (101)-dominant TiO<sub>2</sub> and (101)-dominant Cu-TiO<sub>2</sub>.

cating the n-type nature of the faceted-TiO<sub>2</sub> electrodes (Fig. S26a and b in the ESI†).<sup>41,71</sup> The (001)-dominant Cu-TiO<sub>2</sub> and the (101)-dominant Cu-TiO<sub>2</sub> show smaller slopes compared with those of the pristine (001)-dominant TiO<sub>2</sub> and (101)-dominant TiO<sub>2</sub>, also indicating the higher donor density in the Cu-deposited TiO<sub>2</sub>.<sup>72</sup> These results suggest that the significant enhancement in NH<sub>3</sub> production exhibited by the (101)-dominant Cu-TiO<sub>2</sub> is likely contributed to by the promoted electron transfer. In this regard, the nature of (101) facets, which possess shallow traps on the surface of Ti<sub>5c</sub> in (101) facets, can provide a more conducive environment for electron mobility as reported in the literature.<sup>34</sup>

The above electrocatalytic NO<sub>3</sub><sup>-</sup>RR activities, time-dependent experiments, and electrochemical analysis have collectively suggested that (1) the Cu deposition on the faceted-TiO<sub>2</sub> surface leads to higher NH<sub>3</sub> production, NH<sub>3</sub> selectivity, and FE<sub>NH<sub>3</sub></sub> and (2) the deposition of Cu on the (101) facet is more beneficial for promoting NH<sub>3</sub> production compared to Cu deposition on the (001) facet due to the stronger interaction between the Cu nanoparticles and the dominant (101) facet in the (101)-dominant Cu-TiO<sub>2</sub>. The Cu metal was previously reported as an active catalyst for the NO<sub>3</sub><sup>-</sup>RR in the literature.<sup>62,73–75</sup> The activity originates from the d-orbital electronic configuration of Cu, where the NO<sub>3</sub><sup>-</sup> reactant binds with Cu by electron donation from the highest occupied molecular orbital (HOMO) of NO<sub>3</sub><sup>-</sup> to the empty orbital of Cu accompanied by the backdonation from the fully occupied d orbital to the lowest unoccupied molecular orbital (LUMO) of NO<sub>3</sub><sup>-</sup>.<sup>60</sup> Therefore, in addition to OVs as active sites, the deposited Cu nanoparticles can also serve as active sites in the (001)-dominant Cu-TiO<sub>2</sub> and the (101)-dominant Cu-TiO<sub>2</sub> catalysts, resulting in significantly higher NH<sub>3</sub> production than that from the pristine counterparts. Additionally, due to the strong interaction between the Cu nanoparticles and (101) facets of TiO<sub>2</sub>, the electrons are transferred from the Cu nanoparticles to the (101) facets of TiO<sub>2</sub> resulting in electron-deficient Cu, which can strongly attract and accumulate the negatively charged NO<sub>3</sub><sup>-</sup>.<sup>39</sup> The stronger absorption strength is also found in the NO<sub>2</sub><sup>-</sup> (and \*NO<sub>2</sub>), which suppresses the desorption of \*NO<sub>2</sub> to the electrolyte. Finally, the exposure of (101) facets of TiO<sub>2</sub>, possessing shallow traps for electrons on the Ti<sub>5c</sub> surface could provide a more conducive environment for electron mobility, thus promoting the electron transfer in the (101)-dominant Cu-TiO<sub>2</sub> resulting in significant enhancements in NH<sub>3</sub> production and NH<sub>3</sub> selectivity over the (101)-dominant Cu-TiO<sub>2</sub> catalyst.

## Conclusions

In conclusion, the strong interaction between the deposited Cu nanoparticles and the TiO<sub>2</sub> support with the dominant (101) facet exposure promoted selective NH<sub>3</sub> production from the NO<sub>3</sub><sup>-</sup>RR. The (101)-dominant Cu-TiO<sub>2</sub> catalyst demonstrated the highest NH<sub>3</sub> yield rate of 447.5 μg mg<sub>cat</sub><sup>-1</sup> h<sup>-1</sup> at -0.9 V vs. RHE, which is 4.2 times higher than that of the pris-

tine (101)-dominant TiO<sub>2</sub> catalyst with an NH<sub>3</sub> yield of 106.7 μg mg<sub>cat</sub><sup>-1</sup> h<sup>-1</sup>. In comparison, the (001)-dominant Cu-TiO<sub>2</sub> counterpart showed NH<sub>3</sub> yield rate of 296.6 μg mg<sub>cat</sub><sup>-1</sup> h<sup>-1</sup> under identical reaction conditions, which is 3.3 times higher than that of the pristine (001)-dominant TiO<sub>2</sub> with an NH<sub>3</sub> yield of 87.9 μg mg<sub>cat</sub><sup>-1</sup> h<sup>-1</sup>. The greater increase in catalytic performance for the (101)-dominant TiO<sub>2</sub> with Cu deposition was ascribed to the formation of electron-deficient Cu nanoparticles, the stronger binding energy of \*NO<sub>2</sub>, and the promoted electron transfer originating from the strong interaction between the Cu nanoparticles and (101) facets of TiO<sub>2</sub>. Collectively, these factors suppressed the \*NO<sub>2</sub> desorption and promoted the subsequent hydrogenation process, leading to direct NO<sub>3</sub><sup>-</sup> reduction to NH<sub>3</sub>. These findings can stimulate the future surface design over other oxide supports toward the efficient electrocatalytic NO<sub>3</sub><sup>-</sup>RR to produce NH<sub>3</sub>.

## Experimental

The experimental details are provided in the ESI.† This section summarizes the material synthesis, characterization, and electrochemical measurements.

### Preparation of the (001)-dominant TiO<sub>2</sub> (flat sheet) and the (101)-dominant TiO<sub>2</sub> (truncated octahedral)

The TiO<sub>2</sub> samples with different dominant facets were prepared using the hydrothermal method with NaF as a capping agent, as reported by Mikrut *et al.*<sup>40</sup> In a typical procedure for preparing (001)-dominant TiO<sub>2</sub>, 532 mg NaF was mixed with 160 mL of HCl (12.8%). The mixture was then stirred to form a clear solution. Subsequently, 781.6 mg of TiOSO<sub>4</sub> was added to the solution, followed by constant stirring for 1 h to form a clear solution. The resulting solution was then placed into a 200 mL Teflon-lined stainless-steel autoclave and was heated at 180 °C for 12 h. After naturally cooling down to room temperature, the powders were collected and centrifuged in water and ethanol several times. The obtained powders were then dried at 60 °C in a vacuum oven for 12 h and calcined at 600 °C for 2 h at a heating rate of 2 °C min<sup>-1</sup> under an argon atmosphere (20 mL min<sup>-1</sup>). The (101)-dominant TiO<sub>2</sub> (truncated octahedral) was prepared using the same method as that of the (001)-dominant TiO<sub>2</sub> by adjusting the concentration of HCl (6.7%), the mass of NaF (279.6 mg), and the amount of TiOSO<sub>4</sub> (390.8 mg).

### Copper (Cu) deposition on (001)-dominant TiO<sub>2</sub> and (101)-dominant TiO<sub>2</sub>

The Cu deposition was performed using NaBH<sub>4</sub> as a reducing agent.<sup>41</sup> 20 mg of the (001)-dominant TiO<sub>2</sub> or the (101)-dominant TiO<sub>2</sub> was dispersed in 15 mL of DI water, followed by sonication for 30 min. After sonication, 315 μL of 20 mM Cu (NO<sub>3</sub>)<sub>2</sub>·3H<sub>2</sub>O solution was added slowly under vigorous stirring. The stirring was continued for 30 min. 30 mg of NaBH<sub>4</sub> was then added, and the stirring was continued for another 20 min. The powders were subsequently separated by centrifugation.



gation and washed using excess deionized water and ethanol. The powders were finally dried in a vacuum oven at 60 °C for 12 h.

### Characterization

The crystal phase of each prepared sample was analyzed using an X-ray diffraction (XRD) PANalytical X'pert<sup>3</sup> powder X-ray diffractometer operated at 40 kV and 100 mA with Cu K $\alpha$  radiation. The morphology and the composition of the catalyst were observed using a transmission electron microscope (TEM) FEI Tecnai F20 and scanning electron microscope (SEM) Zeiss EVO 10. The oxidation states of Ti, Cu, and O elements were characterized using X-ray photoelectron spectroscopy (XPS, Thermo Scientific NEXSA) conducted with an Al K $\alpha$  X-ray excitation source. All binding energies were referenced to the C 1s peak at 283.67 eV. Electron paramagnetic resonance (EPR) analysis was performed using a Bruker EMXPLUS EPR instrument (Germany). The amount of deposited metal was measured using inductively coupled plasma optical emission spectroscopy (ICP-OES, Optima 8000). The ultraviolet-visible (UV-vis) absorbance data were collected on a Shimadzu UV-3600 UV spectrophotometer.

### Electrochemical measurements

The prepared samples were dropcast on pieces of carbon paper as the working electrodes. Before the drop-casting, the pieces of carbon paper were washed in a mixture of methanol and acetone (1:1; v/v) and ultrasonicated for 10 min. Subsequently, the pieces of carbon paper were washed again in water and ultrasonicated for another 10 min. The pieces of carbon paper were then dried at room temperature. Meanwhile, 10 mg of TiO<sub>2</sub> powders, *e.g.*, the (001)-dominant TiO<sub>2</sub>, were mixed with 0.24 mL of water and 0.72 mL of ethanol. After that, 0.04 mL of Nafion solution (5%) was added. The mixture was then ultrasonicated for 1 h to form a homogeneous suspension. 0.1 mL of the prepared homogeneous suspension was then dropped onto 1.0 × 1.0 cm<sup>2</sup> carbon paper (1.0 mg<sub>cat</sub> cm<sup>-2</sup>). The same procedure was applied to prepare the electrodes loaded with the other catalysts (*i.e.*, the (101)-dominant TiO<sub>2</sub>, (001)-dominant Cu-TiO<sub>2</sub>, and (101)-dominant Cu-TiO<sub>2</sub>).

The electrochemical measurements were carried out using a CHI 660 instrument in an H-type cell using the TiO<sub>2</sub> based-thin film electrode as the working electrode, saturated calomel electrode (SCE) as the reference electrode, and platinum foil as the counter electrode. The surface area of the working electrode was controlled within 1.0 × 1.0 cm<sup>2</sup>. 0.5 M Na<sub>2</sub>SO<sub>4</sub> was used as the electrolyte. The electrolyte volumes in the anodic and cathodic chambers were 30 and 40 mL, respectively. A Nafion 117 membrane was placed between the anodic and cathodic chambers. The catholyte was purged with Ar at a flow rate of 50 mL min<sup>-1</sup> for 15 min under vigorous stirring. Before the NO<sub>3</sub><sup>-</sup>RR measurement, cyclic voltammetry (CV) was performed from +0.1 V to -1.2 V *vs.* RHE to reach a stable curve. After the CV, NaNO<sub>3</sub> (50 ppm NO<sub>3</sub><sup>-</sup>-N) was added to the cathode chamber. The chronoamperometry (*i-t*) test was

carried out at different potentials for 2 h at a stirring rate of 300 rpm. All potentials were referenced to reversible hydrogen electrode (RHE) by the Nernst equation ( $E_{\text{RHE}} = E_{\text{SCE}} + 0.059 \times \text{pH} + 0.241$ ).

To investigate the electrochemically active surface area (ECSA), the working electrode was cycled in the non-faradaic potential region at various scan rates from 10 to 40 mV s<sup>-1</sup> in 0.5 M Na<sub>2</sub>SO<sub>4</sub>. By plotting the charging current density against the scan rate, the double-layer capacitance ( $C_{\text{dl}}$ ) was calculated by determining the slope value. Electrochemical impedance spectroscopy (EIS) was performed at the potential of -0.7 V *vs.* RHE at a frequency of 10<sup>-1</sup>-10<sup>5</sup> Hz and amplitude of 5 mV. Mott-Schottky plots were recorded at a frequency of 0.5 kHz and amplitude of 0.01 V.

### Conflicts of interest

There are no conflicts to declare.

### Acknowledgements

This project was financially supported by the Hong Kong Research Grant Council (RGC) General Research Fund (GRF) CityU 11306920, CityU 11308721, and CityU 11316522. The authors acknowledge the financial support by the Science and Technology Development Fund, Macao SAR (File No. FDCT-0125/2022/A), and the National Natural Science Foundation of China (22305009).

### References

- 1 L. Wang, M. Xia, H. Wang, K. Huang, C. Qian, C. T. Maravelias and G. A. Ozin, *Joule*, 2018, **2**, 1055-1074.
- 2 B. M. Comer, P. Fuentes, C. O. Dimkpa, Y.-H. Liu, C. A. Fernandez, P. Arora, M. Realff, U. Singh, M. C. Hatzell and A. J. Medford, *Joule*, 2019, **3**, 1578-1605.
- 3 L. Hollevoet, F. Jardali, Y. Gorbanev, J. Creel, A. Bogaerts and J. A. Martens, *Angew. Chem., Int. Ed.*, 2020, **59**, 23825-23829.
- 4 D. W. Kim, D. W. Kang, M. Kang, J.-H. Lee, J. H. Choe, Y. S. Chae, D. S. Choi, H. Yun and C. S. Hong, *Angew. Chem., Int. Ed.*, 2020, **59**, 22531-22536.
- 5 W. P. Utomo, M. K. H. Leung, Z. Yin, H. Wu and Y. H. Ng, *Adv. Funct. Mater.*, 2022, **32**, 2106713.
- 6 A. J. Martín, T. Shinagawa and J. Pérez-Ramírez, *Chem*, 2019, **5**, 263-283.
- 7 S. Lin, X. Zhang, L. Chen, Q. Zhang, L. Ma and J. Liu, *Green Chem.*, 2022, **24**, 9003-9026.
- 8 C. J. M. van der Ham, M. T. M. Koper and D. G. H. Hetterscheid, *Chem. Soc. Rev.*, 2014, **43**, 5183-5191.
- 9 O. Elishav, B. M. Lis, E. M. Miller, D. J. Arent, A. Valera-Medina, A. G. Dana, G. E. Shter and G. S. Grader, *Chem. Rev.*, 2020, **120**, 5352-5436.

- 10 L. Barrera and R. Bala Chandran, *ACS Sustainable Chem. Eng.*, 2021, **9**, 3688–3701.
- 11 T. A. Belay, F. M. Lin, C. Y. Lin, H. M. Hsiao, M. F. Chang and J. C. Liu, *Water Sci. Technol.*, 2015, **72**, 960–965.
- 12 X. Zhang, Y. Wang, C. Liu, Y. Yu, S. Lu and B. Zhang, *Chem. Eng. J.*, 2021, **403**, 126269.
- 13 Y. Li, Y. K. Go, H. Ooka, D. He, F. Jin, S. H. Kim and R. Nakamura, *Angew. Chem., Int. Ed.*, 2020, **59**, 9744–9750.
- 14 X. Wan, W. Guo, X. Dong, H. Wu, X. Sun, M. Chu, S. Han, J. Zhai, W. Xia, S. Jia, M. He and B. Han, *Green Chem.*, 2022, **24**, 1090–1095.
- 15 Y. Wang, L. Zhang, Y. Niu, D. Fang, J. Wang, Q. Su and C. Wang, *Green Chem.*, 2021, **23**, 7594–7608.
- 16 Y.-T. Xu, K.-C. Ren, Z.-M. Tao, D. K. Sam, E. Feng, X. Wang, G. Zhang, J. Wu and Y. Cao, *Green Chem.*, 2023, **25**, 589–595.
- 17 A. Stirling, I. Pápai, J. Mink and D. R. Salahub, *J. Chem. Phys.*, 1994, **100**, 2910–2923.
- 18 R. Jia, Y. Wang, C. Wang, Y. Ling, Y. Yu and B. Zhang, *ACS Catal.*, 2020, **10**, 3533–3540.
- 19 Y. Wang, W. Zhou, R. Jia, Y. Yu and B. Zhang, *Angew. Chem., Int. Ed.*, 2020, **59**, 5350–5354.
- 20 R. Daiyan, T. Tran-Phu, P. Kumar, K. Iputera, Z. Tong, J. Leverett, M. H. A. Khan, A. A. Esmailpour, A. Jalili, M. Lim, A. Tricoli, R.-S. Liu, X. Lu, E. Lovell and R. Amal, *Energy Environ. Sci.*, 2021, **14**, 3588–3598.
- 21 J. Fu, F. Yao, T. Xie, Y. Zhong, Z. Tao, S. Chen, L. He, Z. Pi, K. Hou, D. Wang, X. Li and Q. Yang, *Sep. Purif. Technol.*, 2021, **276**, 119329.
- 22 D. R. MacFarlane, P. V. Cherepanov, J. Choi, B. H. R. Suryanto, R. Y. Hodgetts, J. M. Bakker, F. M. F. Vallana and A. N. Simonov, *Joule*, 2020, **4**, 1186–1205.
- 23 G. Liu, H. G. Yang, J. Pan, Y. Q. Yang, G. Q. (Max) Lu and H.-M. Cheng, *Chem. Rev.*, 2014, **114**, 9559–9612.
- 24 Z. Xiong, Z. Lei, Y. Li, L. Dong, Y. Zhao and J. Zhang, *J. Photochem. Photobiol., C*, 2018, **36**, 24–47.
- 25 B. Shao, W. Zhao, S. Miao, J. Huang, L. Wang, G. Li and W. Shen, *Chin. J. Catal.*, 2019, **40**, 1534–1539.
- 26 Z. Xie, H. L. Tan, H. Wu, R. Amal, J. Scott and Y. H. Ng, *Mater. Today Energy*, 2022, **26**, 100986.
- 27 H. L. Tan, X. Wen, R. Amal and Y. H. Ng, *J. Phys. Chem. Lett.*, 2016, **7**, 1400–1405.
- 28 J. Chen, X. Li, B. Lei, L. Zhou and S. Wang, *Green Chem.*, 2019, **21**, 483–490.
- 29 J. Pan, G. Liu, G. Q. (Max) Lu and H.-M. Cheng, *Angew. Chem., Int. Ed.*, 2011, **50**, 2133–2137.
- 30 W. Wang, J. Fang, Y. Zhou, W. Zhang and C. Lu, *RSC Adv.*, 2016, **6**, 67556–67564.
- 31 N. Sutradhar, A. K. Biswas, S. K. Pahari, B. Ganguly and A. B. Panda, *Chem. Commun.*, 2014, **50**, 11529–11532.
- 32 P. Mikrut, D. Mitoraj, R. Beranek and W. Macyk, *Appl. Surf. Sci.*, 2021, **566**, 150662.
- 33 D. Sun, W. Yang, L. Zhou, W. Sun, Q. Li and J. K. Shang, *Appl. Catal., B*, 2016, **182**, 85–93.
- 34 H. Zhou, M. Wang, F. Kong, Z. Chen, Z. Dou and F. Wang, *J. Am. Chem. Soc.*, 2022, **144**, 21224–21231.
- 35 H. Wu, L. Zhang, A. Du, R. Irani, R. van de Krol, F. F. Abdi and Y. H. Ng, *Nat. Commun.*, 2022, **13**, 6231.
- 36 H. Wu, R. Irani, K. Zhang, L. Jing, H. Dai, H. Y. Chung, F. F. Abdi and Y. H. Ng, *ACS Energy Lett.*, 2021, **6**, 3400–3407.
- 37 M.-S. Kim, S.-H. Chung, C.-J. Yoo, M. S. Lee, I.-H. Cho, D.-W. Lee and K.-Y. Lee, *Appl. Catal., B*, 2013, **142–143**, 354–361.
- 38 J. Li, M. Li, X. Yang, S. Wang, Y. Zhang, F. Liu and X. Liu, *ACS Appl. Mater. Interfaces*, 2019, **11**, 33859–33867.
- 39 Y. Xu, K. Shi, T. Ren, H. Yu, K. Deng, X. Wang, Z. Wang, H. Wang and L. Wang, *Small*, 2022, **18**, 2203335.
- 40 P. Mikrut, M. Kobielski and W. Macyk, *Electrochim. Acta*, 2019, **310**, 256–265.
- 41 W. P. Utomo, H. Wu and Y. H. Ng, *Small*, 2022, **18**, 2200996.
- 42 X. Zhang, L. Luo, R. Yun, M. Pu, B. Zhang and X. Xiang, *ACS Sustainable Chem. Eng.*, 2019, **7**, 13856–13864.
- 43 Y. Zhang, J. Hu, C. Zhang, Y. Liu, M. Xu, Y. Xue, L. Liu and M. K. H. Leung, *J. Mater. Chem. A*, 2020, **8**, 9091–9098.
- 44 G. Liu, H. G. Yang, X. Wang, L. Cheng, H. Lu, L. Wang, G. Q. (Max) Lu and H.-M. Cheng, *J. Phys. Chem. C*, 2009, **113**, 21784–21788.
- 45 Y. Zhai, Y. Ji, G. Wang, Y. Zhu, H. Liu, Z. Zhong and F. Su, *RSC Adv.*, 2015, **5**, 73011–73019.
- 46 M. Fernández-Arias, M. Boutinguiza, J. del Val, A. Riveiro, D. Rodríguez, F. Arias-González, J. Gil and J. Pou, *Nanomaterials*, 2020, **10**, 300.
- 47 F. Yao, M. Jia, Q. Yang, F. Chen, Y. Zhong, S. Chen, L. He, Z. Pi, K. Hou, D. Wang and X. Li, *Water Res.*, 2021, **193**, 116881.
- 48 Z. Han, C. Choi, S. Hong, T.-S. Wu, Y.-L. Soo, Y. Jung, J. Qiu and Z. Sun, *Appl. Catal., B*, 2019, **257**, 117896.
- 49 H. Wang, L. Wang, D. Lin, X. Feng, Y. Niu, B. Zhang and F.-S. Xiao, *Nat. Catal.*, 2021, **4**, 418–424.
- 50 Z. Song, M. N. Banis, L. Zhang, B. Wang, L. Yang, D. Banham, Y. Zhao, J. Liang, M. Zheng, R. Li, S. Ye and X. Sun, *Nano Energy*, 2018, **53**, 716–725.
- 51 I. Jiménez-Morales, S. Cavaliere, D. Jones and J. Rozière, *Phys. Chem. Chem. Phys.*, 2018, **20**, 8765–8772.
- 52 L. Lin, S. Yao, R. Gao, X. Liang, Q. Yu, Y. Deng, J. Liu, M. Peng, Z. Jiang, S. Li, Y.-W. Li, X.-D. Wen, W. Zhou and D. Ma, *Nat. Nanotechnol.*, 2019, **14**, 354–361.
- 53 K. Zhu, F. Shi, X. Zhu and W. Yang, *Nano Energy*, 2020, **73**, 104761.
- 54 G. Wu, G. Zhao, J. Sun, X. Cao, Y. He, J. Feng and D. Li, *J. Catal.*, 2019, **377**, 271–282.
- 55 Y. Zhang, F. Du, R. Wang, X. Ling, X. Wang, Q. Shen, Y. Xiong, T. Li, Y. Zhou and Z. Zou, *J. Mater. Chem. A*, 2021, **9**, 17442–17450.
- 56 W. P. Utomo, H. Wu and Y. H. Ng, *Energies*, 2023, **16**, 27.
- 57 C. Wang, Y. Zhang, H. Luo, H. Zhang, W. Li, W. Zhang and J. Yang, *Small Methods*, 2022, **6**, 2200790.
- 58 O. Schmidt, A. Gambhir, I. Staffell, A. Hawkes, J. Nelson and S. Few, *Int. J. Hydrog. Energy*, 2017, **42**, 30470–30492.

- 59 C. A. Fernandez, N. M. Hortance, Y.-H. Liu, J. Lim, K. B. Hatzell and M. C. Hatzell, *J. Mater. Chem. A*, 2020, **8**, 15591–15606.
- 60 G.-F. Chen, Y. Yuan, H. Jiang, S.-Y. Ren, L.-X. Ding, L. Ma, T. Wu, J. Lu and H. Wang, *Nat. Energy*, 2020, **5**, 605–613.
- 61 D. Hartanto, G. Yuhaneka, W. P. Utomo, A. I. Rozafia, Y. Kusumawati, W. Dahani and A. Iryani, *RSC Adv.*, 2022, **12**, 5665–5676.
- 62 X. Wang, M. Zhu, G. Zeng, X. Liu, C. Fang and C. Li, *Nanoscale*, 2020, **12**, 9385–9391.
- 63 P. Wang, L. Yang, Y. Q. Gao and X. S. Zhao, *Nucleic Acids Res.*, 2015, **43**, 7207–7216.
- 64 X. Niu, X. Nie, C. Yang and J. G. Chen, *Catal. Sci. Technol.*, 2020, **10**, 1881–1888.
- 65 Y. Tang, C. Yang, M. Sheng, X. Yin and W. Que, *ACS Sustainable Chem. Eng.*, 2020, **8**, 12990–12998.
- 66 X. Teng, J. Wang, L. Ji, Y. Lv and Z. Chen, *Nanoscale*, 2018, **10**, 9276–9285.
- 67 C. Zhao, Y. Liang, W. Li, Y. Tian, X. Chen, D. Yin and Q. Zhang, *RSC Adv.*, 2017, **7**, 52614–52620.
- 68 K. Ao, Q. Wei and W. A. Daoud, *ACS Appl. Mater. Interfaces*, 2020, **12**, 33595–33602.
- 69 L. L. Zulfa, R. Ediati, A. R. P. Hidayat, R. Subagyo, N. Faaizatunnisa, Y. Kusumawati, D. Hartanto, N. Widiastuti, W. P. Utomo and M. Santoso, *RSC Adv.*, 2023, **13**, 3818–3834.
- 70 H. Wu, S. Qu, Z. Xie and Y. H. Ng, *ACS Appl. Energy Mater.*, 2022, **5**, 8419–8427.
- 71 G. Yuhaneka, A. I. Rozafia, W. P. Utomo, A. Iryani and D. Hartanto, *Malays. J. Fundam. Appl. Sci.*, 2022, **18**, 463–472.
- 72 H. Y. Chung, R. J. Wong, R. Amal and Y. H. Ng, *Energy Fuels*, 2022, **36**, 11550–11558.
- 73 X. Fu, X. Zhao, X. Hu, K. He, Y. Yu, T. Li, Q. Tu, X. Qian, Q. Yue, M. R. Wasielewski and Y. Kang, *Appl. Mater. Today*, 2020, **19**, 100620.
- 74 E. Pérez-Gallent, M. C. Figueiredo, I. Katsounaros and M. T. M. Koper, *Electrochim. Acta*, 2017, **227**, 77–84.
- 75 S.-E. Bae and A. A. Gewirth, *Faraday Discuss.*, 2008, **140**, 113–123.