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# Anodic SnO<sub>2</sub> porous nanostructures with rich grain boundaries for efficient CO<sub>2</sub> electroreduction to formate†

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Formic acid (HCOOH), the acidic form of formate, is an important hydrogen carrier which can be directly used in fuel cells. Development of earth-abundant element-based catalysts to convert carbon dioxide (CO<sub>2</sub>) into HCOOH or formate with high selectivity and high efficiency has been a vigorous research activity in recent years but remains an unsolved challenge. In this contribution, using one-step anodization, we prepare nanotubular SnO<sub>2</sub> porous nanostructures with high surface area (90.1 m<sup>2</sup> g<sup>-1</sup>), large porosity (0.74 cm<sup>3</sup> g<sup>-1</sup>), and rich grain boundaries for electrochemical CO<sub>2</sub> reduction (CO<sub>2</sub>RR). They exhibit stable 95% faradaic efficiency (FE) towards CO<sub>2</sub>RR and 73% FE for formate at -0.8 V<sub>RHE</sub>. The notable performance of such SnO<sub>2</sub> nanostructures can be attributed to their unique structural and chemical properties, which provide active sites for CO<sub>2</sub> adsorption and conversion, and easy access for CO<sub>2</sub> to the active sites. The insights gained from the structure/property relationships might be beneficial for designing superior electrocatalysts for CO<sub>2</sub> electroreduction into formate.

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## Introduction

Electrochemical CO<sub>2</sub> reduction (CO<sub>2</sub>RR) to carbon-based fuels is a promising strategy for renewable energy storage and net carbon emission.<sup>1–5</sup> Liquid fuels, such as formic acid (HCOOH),<sup>6–8</sup> methanol<sup>9</sup> and ethanol,<sup>10</sup> produced *via* CO<sub>2</sub> electroreduction are actively sought because they can be directly used as energy-intensive carriers for fuel cells. Particularly, formic acid has been identified as an ideal hydrogen carrier owing to its high volumetric density (53.4 g H<sub>2</sub> L<sup>-1</sup>), and low toxicity and volatility.<sup>11</sup> However, achieving highly efficient and selective electrocatalysts for converting

CO<sub>2</sub> into HCOOH is challenging due to the myriad possible reaction pathways and sluggish reaction kinetics in the process of CO<sub>2</sub>RR.

The formation of the intermediate CO<sub>2</sub>\*<sup>-</sup> is a rate-determining step for CO<sub>2</sub>RR in most pathways.<sup>12</sup> Metals like Pb, In, Zn, Sn, Pd and Bi, can hardly bind the intermediate CO<sub>2</sub>\*<sup>-</sup> and hence are able to produce formic acid (or formate when pH > 3.8) at low overpotentials typically *via* the outer-sphere mechanism.<sup>13</sup> Among these metal-based electrocatalysts, Sn and its derivatives (*e.g.*, SnO<sub>x</sub>, Sn alloy), as earth-abundant, low-cost and nontoxic materials, have attracted particular research interest, due to their capability of catalyzing CO<sub>2</sub> into formate with decent faradaic efficiencies (FEs) at moderate potentials.<sup>14–21</sup> For example, Meyer *et al.* found a maximum FE of 86% for formate at -1.8 V<sub>SCE</sub> when loading nanoscale SnO<sub>2</sub> particles on carbon black.<sup>14</sup> Xie *et al.* reported that Sn quantum sheets confined in graphene exhibited ~89% FE towards formate at the same potential.<sup>15</sup> These results point out that the catalyst size and composition, which are correlated with the number and intrinsic properties of active sites exposed during CO<sub>2</sub> reduction, play important roles in determining the formation of formate. In addition, the electrocatalysis of CO<sub>2</sub> in aqueous solution is an electrochemical reaction at the three-phase interface of gas (CO<sub>2</sub>), liquid (electrolyte) and solid (catalyst). The concentration of reactant CO<sub>2</sub> on the surface of the catalyst affects the electrocatalytic performance and is governed by at least two factors: (1) the CO<sub>2</sub> transport within the catalysts; (2) the CO<sub>2</sub> adsorption on

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the catalysts. Therefore, the structure of the catalyst also matters critically in the formate formation activity. For example, hierarchical mesoporous SnO<sub>2</sub> nanosheets were reported to facilitate the mass transfer of CO<sub>2</sub>, leading to a value of 87% FE for formate at a moderate overpotential (0.88 V).<sup>16</sup> SnO<sub>2</sub> nanowires with a high density of grain boundaries were proposed to enhance the CO<sub>2</sub> reduction into formate (*ca.* 80% FE at  $-0.8 V_{\text{RHE}}$ ) *via* reforming the binding energy of the reactant or intermediates.<sup>17</sup> Taken together, these works conclude that formate production by CO<sub>2</sub>RR is sensitive to a variety of parameters, including size, structure and composition *etc.* However, to date, it remains a challenge to incorporate these parameters to achieve an efficient electrocatalyst for the conversion of CO<sub>2</sub> to formate.

In this work, we report the results of an electrochemical study on the CO<sub>2</sub> reduction at SnO<sub>2</sub> porous nanostructures with integrated characteristics of high surface area, large porosity, and rich grain boundaries that favor the CO<sub>2</sub>-to-formate conversion. The SnO<sub>2</sub> porous nanostructures were prepared by an anodization approach which is a facile yet effective technique for fabricating various porous metal oxides with ordered tubular nanostructure.<sup>22</sup> The as-prepared anodic SnO<sub>2</sub> porous nanostructures exhibit a tubular structure (pore-size:  $\sim 45$  nm) with abundant nanograins within the size range of 3–5 nm. The porous structure provides a large specific surface area ( $90.1 \text{ m}^2 \text{ g}^{-1}$ ) and high porosity ( $0.74 \text{ cm}^3 \text{ g}^{-1}$ ), while the grain boundaries render a strong CO<sub>2</sub> affinity with the heat of adsorption up to  $50 \text{ kJ mol}^{-1}$ , the upper limit of the physical adsorption. When tested as the electrocatalyst for the CO<sub>2</sub>RR, the SnO<sub>2</sub> aerogels maintain 95% faradaic efficiency (FE) towards CO<sub>2</sub>RR and 73% FE for HCOOH at  $-0.8 \text{ V vs. RHE}$  for 12 h. Miscellaneous characterizations, including transmission electron microscope (TEM), N<sub>2</sub> (77 K) and CO<sub>2</sub> (273 and 298 K) adsorption, X-ray photoelectron spectroscopy (XPS) and electrochemistry, combined with theoretical calculations were employed to elucidate the correlations between the structure and composition of the aerogels and their CO<sub>2</sub>RR performance.

## Experimental section

### Chemicals and instruments

Sn foil (thickness of 200  $\mu\text{m}$ , purity > 99.99%, Renxin Metal), Nafion solution (5 wt%, Shanghai Hesen), Nafion® 117 cation exchange membrane (H<sup>+</sup> form, Shanghai Hesen), carbon black (VULCAN XC72, Cabot), and other chemicals (Analytic Reagent) are used as received. The X-ray diffraction (XRD) was taken from Rigaku Ultima Iv. A scanning electron microscope (SEM, FEI, Verios 460L) and a high-resolution transmission electron microscope (HRTEM, FEI, Talos F200X) were utilized to characterize the morphology and high-resolution structure. X-ray photoelectron spectrometer (XPS) was determined on a Thermo Scientifica (ESCALAB250xi). Electrochemical measurements were carried on an electrochemical workstation (CHI760e, Shanghai CHI). Products in the gas phase were determined by a gas chromatograph (GC, 9790II, Zhejiang Fuli) equipped with thermal conductivity and a flame ionization

detector. Products in solution was detected by a nuclear magnetic resonance (NMR, Bruker, 400M).

### Synthesis of SnO<sub>2</sub> tubular array by anodic oxidation

The surface oxide layer was firstly removed by soaking an Sn foil in a KOH solution (0.1 M) for 5 min, and then the Sn foil was ultrasonically cleaned in deionized (DI) water for later use. The anodization process was carried in a two-electrode cell, where the cleaned Sn foil was used as a working electrode and a graphite plate was used as the counter electrode with the distance of 1 cm. A constant voltage (8.0–12.0 V) was applied to the working electrode using a potentiostat instrument in a 0.5 M oxalic acid (H<sub>2</sub>C<sub>2</sub>O<sub>4</sub>·2H<sub>2</sub>O, 99.5%) electrolyte for 5 min at room condition. After anodization, the Sn foil was repeatedly washed with DI water before ultrasonically stripped.

### Fabrication of catalytic electrode

The electrode was fabricated through a dropping-coating method. Typically, SnO<sub>2</sub> powders (4 mg), carbon black (2 mg) and Nafion® solution (60  $\mu\text{L}$ ) were mixed with 1 mL mixture of DI water and ethanol (7 : 3 in v/v) and followed with ultrasonic treatment for 5 min to form a homogeneous ink. The suspension (80  $\mu\text{L}$ ) was dropped onto a glassy carbon electrode with an area of  $1 \times 1 \text{ cm}^2$  to form the catalyst layer with a loading amount of  $300 \mu\text{g cm}^{-2}$  and the electrode were naturally dried for further electrochemical measurements.

### Electrochemical measurement

All electrochemical CO<sub>2</sub> reductions were performed by using a well-sealed H-type cell with standard three-electrodes, in which two chambers were separated with a piece of Nafion® 117 membrane. One side of the H-cell was equipped with inlet and outlet, where CO<sub>2</sub> (99.99%) could be bubbled into the reaction cell through an inlet and the product in the gas phase would be introduced into a GC by the outlet. The area of the working electrode was  $1 \text{ cm} \times 1 \text{ cm}$ . An Ag/AgCl electrode (sat. KCl) was used as a reference electrode, and a Pt foil (99.999%,  $1 \text{ cm} \times 1 \text{ cm}$ ) was used as a counter electrode, which was infiltrated into the electrolyte of 0.5 M KHCO<sub>3</sub> aqueous solution for all electrochemical testing. The cyclic voltammetry (CV) under potentials ranging from 0.4 to  $-1.0 V_{\text{RHE}}$  by varying the scan rate of  $50 \text{ mV s}^{-1}$ . The time-current curve was obtained by carrying out constant working potential at various potentials  $-0.7$  to  $-1.2 V_{\text{RHE}}$ . Meanwhile, the average current density was obtained by dividing the total current by the geometric area of the working electrode. The potential values involved in this work were all calibrated to the potential of the reversible hydrogen electrode (RHE) by using eqn (1):<sup>23</sup>

$$E(\text{vs. RHE}) = E(\text{vs. Ag/AgCl}) + 0.197 + 0.0591 \times \text{pH} \quad (1)$$

The high-purity CO<sub>2</sub> was continuously introduced into the electrolyte (0.5 M KHCO<sub>3</sub>) for 30 min to make the solution reach a saturated state of CO<sub>2</sub> (pH = 7.2) before the electrochemical test. The flow rate of CO<sub>2</sub> gas was controlled to be  $5 \text{ mL min}^{-1}$  and the cathode chamber electrolyte was continuously stirred to accelerate



the bubble diffusion on the electrode surface at a rotation speed of 600 rpm during the electrolysis process. All tests for this work were controlled at room temperature and ambient pressure.

### Product determination

The gas-phase products were quantified using a GC during 3 h long bulk electrolysis. Under the same test environment, the gas peak area detected by gas chromatography has a certain linear relationship with its actual concentration, so that the gas with known concentration can be used to calibrate and analyze the gas with unknown concentration, which combined with current density to calculate the faradaic efficiency (FE) by using eqn (2)–(4):<sup>23</sup>

$$j_{\text{H}_2} = \frac{(\text{peak area})/\alpha \times (\text{flow rate}) \times (2FP_0/RT)}{(\text{electrode area})^{-1}} \quad (2)$$

$$j_{\text{CO}} = \frac{(\text{peak area})/\beta \times (\text{flow rate}) \times (2FP_0/RT)}{(\text{electrode area})^{-1}} \quad (3)$$

$$\begin{aligned} \text{FE}_{\text{H}_2/\text{CO}} (\%) &= (i_{\text{H}_2/\text{CO}}/i_{\text{total}}) \times 100 \\ &= (V_{\text{H}_2/\text{CO}} \times Q \times (2FP_0/RT)) \times 100 \end{aligned} \quad (4)$$

where,  $j_{\text{H}_2}$  and  $j_{\text{CO}}$  represent partial current densities of  $\text{H}_2$  and  $\text{CO}$ ,  $\alpha$  and  $\beta$  represent peak area conversion coefficients of  $\text{H}_2$  and  $\text{CO}$  calculated from standard GC mixtures,  $F$  is Faraday constant ( $96485 \text{ C mol}^{-1}$ ),  $R$  is ideal gas constant ( $8.314 \text{ J mol K}^{-1}$ ),  $P_0 = 1.013 \text{ bar}$ ,  $T = 273.15 \text{ K}$ .

The product in solution was detected and quantified by NMR. The 0.4 mL of post-cathodic reaction mixed with 0.1 mL deuterioxide ( $\text{D}_2\text{O}$ ) and 0.1 mL dimethyl sulfoxide (DMSO, 100 ppm) as internal standard. The  $^1\text{H}$  spectrum was measured with water suppression by a presaturation method. The area ratio of the formate peak to the DMSO peak was compared to the standard curve made by using sodium formate and the internal standard DMSO to calculate the amounts of formate products.

### Computational details

All Spin-polarized calculations in this work are carried out using the Quickstep<sup>24</sup> module of the CP2K program package,<sup>24,25</sup>

a program developed for electronic structure calculations and molecular dynamics based on the Gaussian and plane waves formalism.<sup>26</sup> The generalized-gradient approximation (GGA) with spin-polarized revised PBE functional (revPBE) is selected to describe the exchange-correlation energy. All the structures and energies of the stationary points included in our work are evaluated at the DFT-D3 level with the approximation suggested by Grimme<sup>27</sup> added to the revPBE calculated energy. The wavefunctions were expanded in an optimized double- $\zeta$  Gaussian basis set<sup>28,29</sup> with a cutoff energy of 500 Rydberg.<sup>30</sup> Core electrons have been modeled by scalar relativistic norm-conserving pseudo potentials<sup>30</sup> with 4, 6, 1 and 4 valence electrons for Sn, O, H and C, respectively. Brillouin zone integration is performed with a reciprocal space mesh consisting of only the gamma point. The convergence criterion for the maximum force is set as  $2 \times 10^{-3}$ . Thermochemistry calculation is implemented by TAMkin,<sup>31</sup> a toolkit for normal mode analysis A ( $19.11 \text{ \AA} \times 20.10 \text{ \AA}$ ) and 5 O–Sn–O layers thick supercell has been selected to represent the  $\text{SnO}_2\{110\}$  surface. A vacuum region between repeated slabs is set to be 30  $\text{\AA}$ , which ensures the negligible interaction between periodic replicas.

## Results and discussion

Fig. 1 illustrates the typical fabrication of the porous  $\text{SnO}_2$  nanostructures by the anodization approach wherein an Sn foil and a graphite plate were used as the anode and cathode, respectively, in an oxalic acid solution (0.5 M). After being applied with a specific voltage (*i.e.*, 8, 10 and 12 V) for 5 min, the porous  $\text{SnO}_2$  structures were formed on the surface of the Sn foil. They are hereafter denoted as  $\text{SnO}_2\text{-AO}_x$  ( $x$ : the anodization voltage). The  $\text{SnO}_2\text{-AO}_x$  layers can be easily exfoliated from the anode, and then used to construct the catalytic electrode for  $\text{CO}_2\text{RR}$ .

Fig. 2 and S1† report the structural characterization of the as-prepared  $\text{SnO}_2\text{-AO}_x$  aerogels by scanning electron microscopy (SEM) and transmission electron microscope (TEM). From the SEM of the  $\text{SnO}_2\text{-AO}_{10}$  (Fig. 2a and b), highly ordered porous structures are observed with the average pore size within the range of 40–50 nm (Fig. 2c). Varying the anodization voltage (*i.e.*, 8 and 12 V) leads to similarly porous  $\text{SnO}_2\text{-AO}_8$  and  $\text{SnO}_2\text{-}$

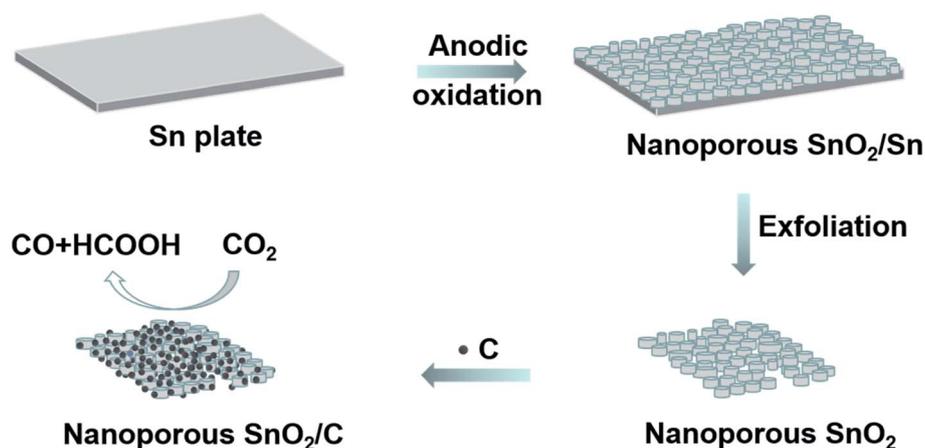


Fig. 1 Schematic representation of the fabrication of anodic  $\text{SnO}_2\text{-AO}_x$  porous nanostructures that are used as the electrocatalysts for  $\text{CO}_2\text{RR}$ .



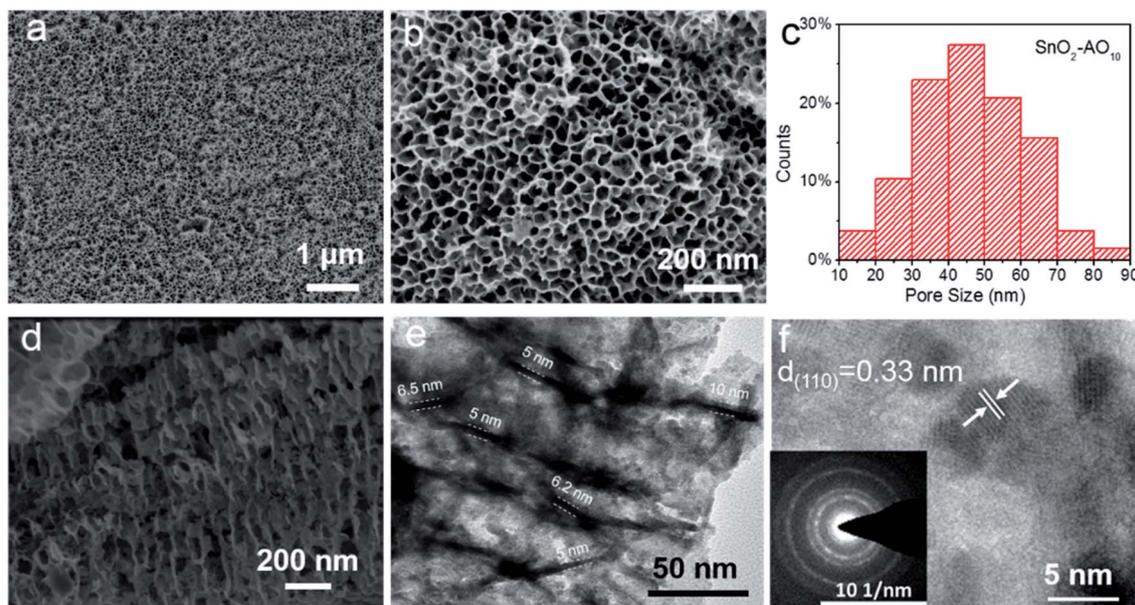


Fig. 2 (a and b) SEM images and (c) the corresponding pore size statistical analysis of the surface of  $\text{SnO}_2\text{-AO}_{10}$ . (d) SEM and (e) TEM images of the cross-section of  $\text{SnO}_2\text{-AO}_{10}$ . (f) High-resolution TEM image and the corresponding electron diffraction pattern (inset) of  $\text{SnO}_2\text{-AO}_{10}$ .

$\text{AO}_{12}$  nanostructures with only slightly different pore sizes (Fig. S1†). The cross-section SEM image further illustrates that the porous  $\text{SnO}_2$  layer is comprised of arrays of tubes that are nearly perpendicular to the Sn substrate (Fig. 2d and S2†). Here, the tubes are expected to promote the mass transfer and reaction during  $\text{CO}_2\text{RR}$ .<sup>14</sup> TEM was additionally employed to study the tubular structure, which shows that the wall thickness of the tubes is 5–10 nm (Fig. 2e). Interestingly, a closer view enabled by the high-resolution TEM (HRTEM) reveals that the tube walls are constituted by many small interconnected nanoparticles (3–

5 nm) through grain boundaries (Fig. 2f and S3†). The individual particles have good crystallinity, as indicated by the interplanar distance of 0.33 nm, which corresponds to the  $d$ -spacing of the (110) plane of  $\text{SnO}_2$ , and further confirmed by some discrete spots in the corresponding electron diffraction pattern of the HRTEM image (inset of Fig. 2f). In addition, the diffraction rings resulting from many spots very close together demonstrate the polycrystalline nature of the tube wall as a whole. Since ultra-small  $\text{SnO}_x$  nanoparticles and grain boundaries have been found to be active sites for  $\text{CO}_2\text{RR}$ ,<sup>16,17,32</sup>

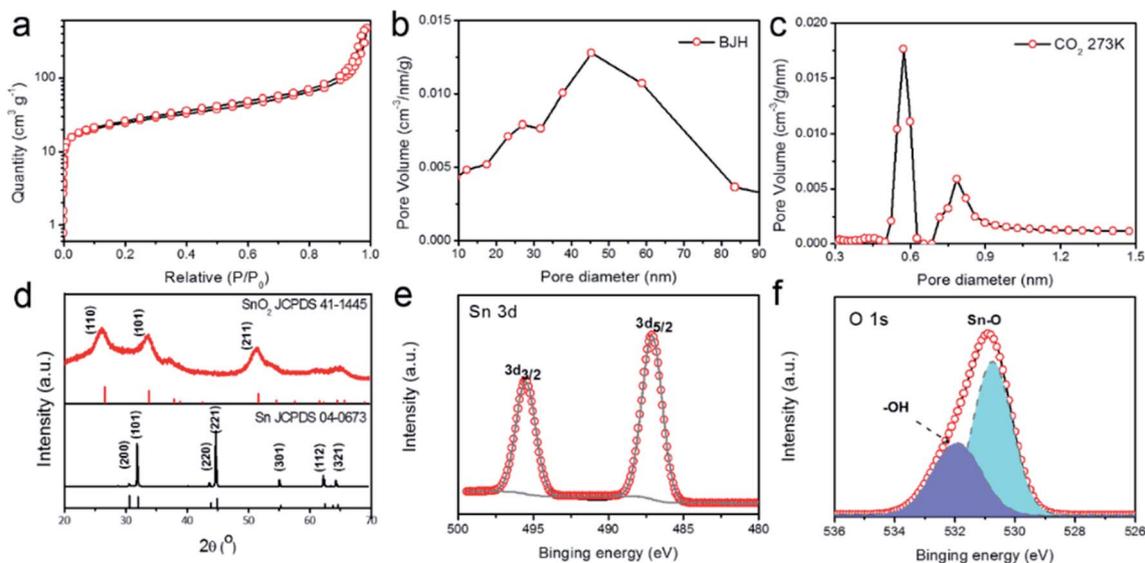


Fig. 3 (a)  $\text{N}_2$  adsorption–desorption isotherm (at 77 K), (b) the corresponding pore size distribution calculated based on the desorption data of the isotherm, and (c) pore size distribution derived from  $\text{CO}_2$  adsorption at 273 K, (d) XRD patterns of  $\text{SnO}_2\text{-AO}_{10}$  and Sn foil, (e) XPS Sn 3d and (f) XPS O 1s spectrum of  $\text{SnO}_2\text{-AO}_{10}$ .



the hierarchical SnO<sub>2</sub>-AO<sub>10</sub> nanostructures that feature tubular structure, ultra-small NPs and abundant grain boundaries show great promise for CO<sub>2</sub>RR.

To gain more insights into the porous structure, N<sub>2</sub> adsorption-desorption isotherms at 77 K were measured to characterize the surface area and porosity of the SnO<sub>2</sub>-AO<sub>10</sub> nanostructures. As shown in Fig. 3a, the N<sub>2</sub> uptake increases slowly with the pressure in the low-pressure range then shoots up when  $P/P_0$  exceeds 0.9. The isotherm thus belongs to the Type-II, as defined by IUPAC,<sup>33</sup> which indicates the presence of both micropores and mesopores in hierarchical porous structures. By applying the Barrett-Joyner-Halenda (BJH) method to the desorption data, the mesopores were found to be within an average size range of 40–50 nm (Fig. 3b), in good agreement with the SEM results (Fig. 2b and c) and reflecting the actual pores created by the anodic oxidation. On the basis of additional data of CO<sub>2</sub> adsorption at 273 K, the micropores were found to display a bimodal size distribution at 0.6 and 0.8 nm (Fig. 3c).

These micropores may arise from the gaps between nanoparticles in the tube walls, as revealed by the HRTEM image of the structure (Fig. 2f). Thanks to these pores of micro- and meso-sizes, the SnO<sub>2</sub>-AO<sub>10</sub> nanostructures exhibit an extremely high BET surface area (90.1 m<sup>2</sup> g<sup>-1</sup>) and porosity (0.74 cm<sup>3</sup> g<sup>-1</sup>). As a result, fast mass transport and easy access of reactants to the active sites during CO<sub>2</sub>RR can be expected.

Finally, we investigated the crystal phase and chemical structure of the SnO<sub>2</sub>-AO<sub>10</sub> aerogels by X-ray diffraction (XRD, Fig. 3d) and X-ray photoelectron spectroscopy (XPS, Fig. 3e and f). From the XRD pattern, the diffraction peaks that can be

unambiguously indexed to rutile tetragonal SnO<sub>2</sub> (JCPDS card no. 41-1445) are observed, confirming the pure SnO<sub>2</sub> phase and crystallinity of SnO<sub>2</sub>-AO<sub>10</sub> nanostructures and demonstrating the successful conversion of pure Sn to SnO<sub>2</sub>.<sup>34</sup> As a comparison, SnO<sub>2</sub>-AO<sub>8</sub> and SnO<sub>2</sub>-AO<sub>12</sub> nanostructures exhibit similar pure SnO<sub>2</sub> phase but much poorer crystallinity (Fig. S4†). However, the mechanism as to how the anodization voltage affects the crystallinity is unclear at this moment, which will be examined in the future. Concerning the compositional information, the two peaks at 495.5 and 487.1 eV in the XPS spectrum are the characteristic peaks that can be assigned to Sn 3d<sub>3/2</sub> and 3d<sub>5/2</sub> ionizations, respectively, confirming the Sn<sup>4+</sup> valence state of Sn element in the tin oxide aerogels (Fig. 3e).<sup>34</sup> Additionally, the XPS spectrum of O 1 s was collected (Fig. 3f). The peak fitting analysis verifies the oxygen bonding either with Sn as Sn-O bond (530.8 eV) or with H as -OH (531.9 eV) adsorbed on the aerogels surface.<sup>35</sup>

To evaluate the CO<sub>2</sub>RR performance, we first examined the catalytic activity of SnO<sub>2</sub>-AO<sub>x</sub> nanostructures and an Sn plate by the Cyclic Voltammetry (CV) (Fig. 4a and S5†). In the N<sub>2</sub>-saturated 0.5 M KHCO<sub>3</sub> solution, the marked increase in the cathodic current at  $\sim -0.80$  V vs. RHE is associated with the hydrogen evolution reaction (HER), a parasite and competing reaction in aqueous solution against CO<sub>2</sub>RR. By contrast, a more drastic current increase occurring at a much lower potential ( $-0.6$  V<sub>RHE</sub>) was observed in the CO<sub>2</sub>-saturated 0.5 M KHCO<sub>3</sub> solution, meaning that the CO<sub>2</sub>RR is catalytically more favored over HER on the SnO<sub>2</sub>-AO<sub>10</sub> catalysts. It is worth noting that the two cathodic peaks correspond to the reduction of SnO<sub>2</sub> to SnO (at  $-0.26$  V<sub>RHE</sub>) and SnO to Sn (at  $-0.34$  V<sub>RHE</sub>), while the anodic peaks arise from the oxidation of Sn to SnO (at  $-0.1$  V

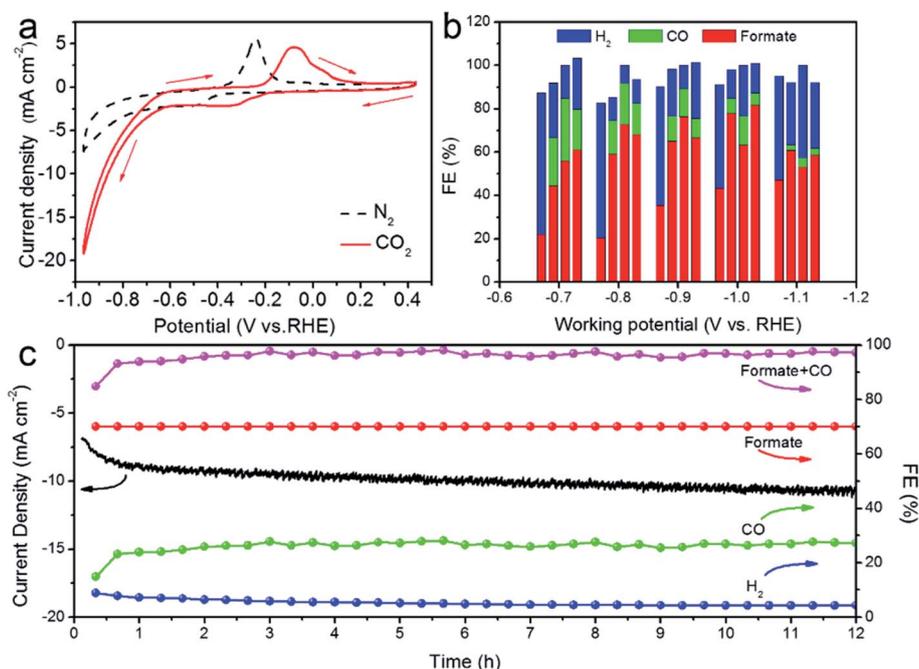


Fig. 4 (a) CV curves of SnO<sub>2</sub>-AO<sub>10</sub> in N<sub>2</sub>- and CO<sub>2</sub>-saturated 0.5 M KHCO<sub>3</sub> solution, scan rate: 50 mV s<sup>-1</sup>. (b) FE of Sn plate, SnO<sub>2</sub>-AO<sub>8</sub>, SnO<sub>2</sub>-AO<sub>10</sub>, and SnO<sub>2</sub>-AO<sub>12</sub> (from left to right at each potential) for H<sub>2</sub>, CO and formate. (c) Temporal evolution of the current density and FE toward H<sub>2</sub>, CO, formate and CO + formate of the SnO<sub>2</sub>-AO<sub>10</sub> catalyst at  $-0.8$  V<sub>RHE</sub> for 12 h.



and SnO to SnO<sub>2</sub> (at 0.05 V).<sup>34</sup> As control samples, SnO<sub>2</sub>-AO<sub>8</sub>, SnO<sub>2</sub>-AO<sub>12</sub> and Sn electrode exhibit slightly more negative onset potentials ( $\sim -0.7$  V<sub>RHE</sub>) for the CO<sub>2</sub>RR and much lower current density at each specific potential (Fig. S6,† for example, SnO<sub>2</sub>-AO<sub>8</sub>:  $\sim 6$  mA cm<sup>-2</sup>, SnO<sub>2</sub>-AO<sub>12</sub>:  $\sim 3$  mA cm<sup>-2</sup>, Sn:  $\sim 2$  mA cm<sup>-2</sup>, SnO<sub>2</sub>-AO<sub>10</sub>:  $\sim 10$  mA cm<sup>-2</sup> at  $-0.8$  V<sub>RHE</sub>). The lower activity of the other two SnO<sub>2</sub> nanostructures (*i.e.*, SnO<sub>2</sub>-AO<sub>8</sub> and SnO<sub>2</sub>-AO<sub>12</sub>) might be due to their poorer crystallinity (Fig. S4†) which limits the electrical conductivity, as corroborated by the Electrochemical Impedance Spectroscopy (EIS) analysis of the electrodes made of the three SnO<sub>2</sub>-AO<sub>x</sub> nanostructures (Fig. S7†). Overall, the CV results demonstrate the superior CO<sub>2</sub>RR activity of SnO<sub>2</sub>-AO<sub>10</sub> aerogels to those of the Sn plate and the SnO<sub>2</sub>-AO<sub>8</sub> and SnO<sub>2</sub>-AO<sub>12</sub> nanostructures. We further assessed the selectivity (FE) of the SnO<sub>2</sub>-AO<sub>x</sub> nanostructures and Sn plate toward each product (*i.e.*, H<sub>2</sub>, CO, formate) at potentials ranging from  $-0.7$  to  $-1.1$  V<sub>RHE</sub>, the results of which are summarized in Fig. 4b and separately presented in Fig. S8–S12.† Consistent with previous reports,<sup>12</sup> the main products detected by NMR and GC during the electrolysis on the Sn plate are H<sub>2</sub> and formate (Fig. 4b, S8 and S9†). The FE for formate progressively increases with the potential from  $\sim 20\%$  at  $-0.7$  V<sub>RHE</sub> until reaching a maximum of  $\sim 48\%$  at  $-1.1$  V<sub>RHE</sub>. Conversely, H<sub>2</sub> dominates in the lower potential range (FE:  $\sim 65\%$  at  $-0.7$  V<sub>RHE</sub>).

Although the FE for H<sub>2</sub> drops as the potential becomes more negative, it sustains at a high level of  $\sim 50\%$  at  $-1.1$  V<sub>RHE</sub>, indicative of strong competition of HER against CO<sub>2</sub>RR on the Sn plate. Interestingly, the anodization of the Sn plate resulted in a dramatic improvement in the CO<sub>2</sub>RR. As illustrated in Fig. 4b and S10–S12,† utilizing SnO<sub>2</sub>-AO<sub>x</sub> nanostructures as the electrocatalyst generates an additional product, *i.e.* CO. The FE toward CO, which decreases with the increasingly negative potential, can be as high as  $\sim 25\%$  at low potentials of  $-0.7$  and  $-0.8$  V<sub>RHE</sub>. Moreover, regardless of the potential, the formation of formate has been significantly boosted. For example, the highest FE achieved on SnO<sub>2</sub>-AO<sub>12</sub> ( $\sim 82\%$  at  $-1.0$  V<sub>RHE</sub>) corresponds to a  $\sim 1.8$ -fold enhancement over that on the Sn plate ( $\sim 46\%$  at  $-1.0$  V<sub>RHE</sub>) and is even comparable with those state-of-the-art results ever reported (Table S1†). Concomitantly, HER is substantially suppressed. In particular, the FE for H<sub>2</sub> obtained over SnO<sub>2</sub>-AO<sub>10</sub> is critically below  $\sim 5\%$  at  $-0.8$  V<sub>RHE</sub>, in stark contrast with the  $\sim 60\%$  over the Sn plate. If we consider the total carbon conversion, the best performance is achieved over the SnO<sub>2</sub>-AO<sub>10</sub> aerogels which display  $\sim 95\%$  total FE (CO:  $\sim 22\%$ ; formate:  $\sim 73\%$ ) towards the CO<sub>2</sub>RR at  $-0.8$  V<sub>RHE</sub>. Essentially, such a superior electrocatalytic performance can be well maintained, as evidenced by the stable current density and the C1 product (CO + formate) conversion, as well as the constantly low FE toward H<sub>2</sub> ( $< \sim 5\%$ ) over a long-term electrolysis of 12 h (Fig. 4c). A plausible explanation for the high

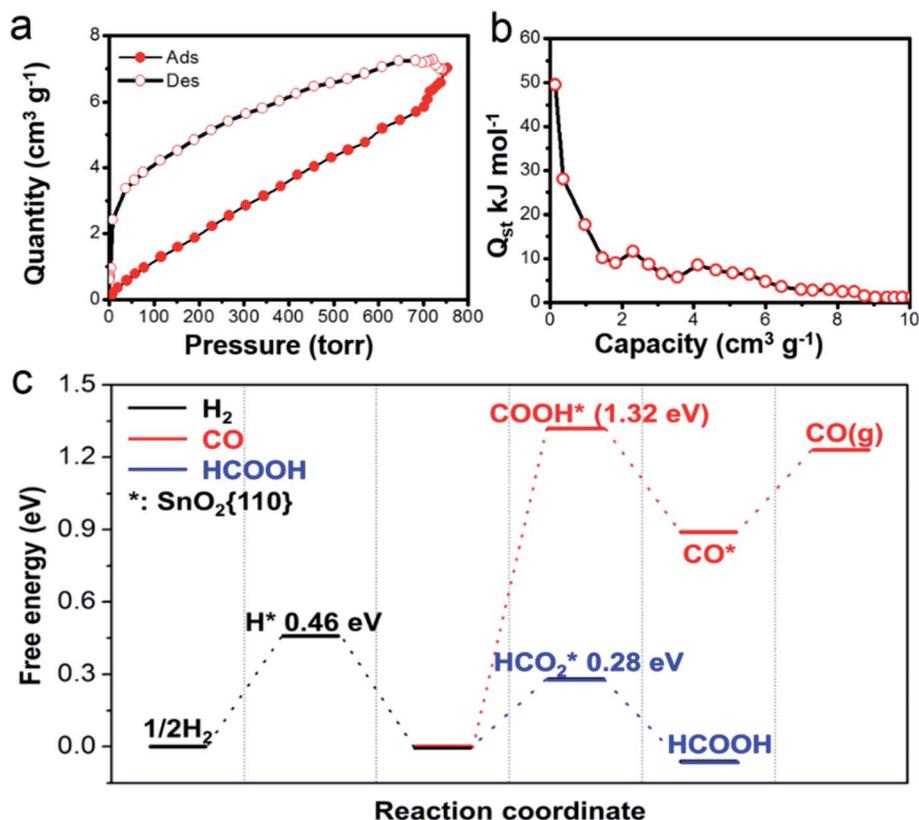


Fig. 5 (a) CO<sub>2</sub> adsorption–desorption isotherm at 298 K and (b) the corresponding heat of CO<sub>2</sub> adsorption derived from the isotherm by employing the Clausius–Clapeyron equation. (c) Free energy diagrams for the reaction pathways to H<sub>2</sub>, CO and HCOOH on SnO<sub>2</sub> (110).



stability is good preservation of the porous structure, grain boundaries and the chemical state of the Sn species (*i.e.*, Sn<sup>4+</sup>), as unravelled by the SEM, XRD and depth-dependent XPS characterizations on the post-mortem SnO<sub>2</sub>-AO<sub>10</sub> catalyst (Fig. S13–S15†). These structural and compositional parameters achieved by simply anodizing the Sn plate are thus the key to understanding the mechanism by which the anodic SnO<sub>2</sub> nanostructures promote pronouncedly the CO<sub>2</sub>RR.

As suggested by earlier studies, porous structures are highly beneficial for CO<sub>2</sub>RR because they provide a large surface area and facilitate the mass transfer.<sup>36</sup> Our SnO<sub>2</sub> aerogels should be particularly advantageous in this aspect, because of their unique porous structure. As a matter of fact, the BET surface area (90.1 m<sup>2</sup> g<sup>-1</sup>) for the SnO<sub>2</sub>-AO<sub>10</sub> nanostructures is considerably higher than that attained by most porous SnO<sub>2</sub> nanostructures (*e.g.*, SnO<sub>2</sub> porous nanowires: 35 m<sup>2</sup> g<sup>-1</sup>,<sup>17</sup> mesoporous tin oxide: 69.2 m<sup>2</sup> g<sup>-1</sup>,<sup>36</sup>) and reaches a comparable level to that of hierarchical mesoporous SnO<sub>2</sub> nanosheets (93.6 m<sup>2</sup> g<sup>-1</sup>).<sup>16</sup> Also, the EIS conducted on the SnO<sub>2</sub>-AO<sub>10</sub> electrode indeed shows a much-improved electrolyte (containing CO<sub>2</sub> in the form of KHCO<sub>3</sub>) transfer over that on the Sn plate (Fig. S7†). In addition to the accessibility to the reactant, another important factor in determining the catalytic behavior is the intrinsic active sites the catalyst owns. As mentioned above, the SnO<sub>2</sub> aerogels present a high density of grain boundaries. Grain boundaries have been proposed to be enhanced active sites for CO<sub>2</sub>RR due to their favorable electronic and chemical properties which tune the binding energy of the reaction intermediates.<sup>37,38</sup> Here, we further propose that the grain boundary might also benefit the CO<sub>2</sub> adsorption on the catalyst, the first step of CO<sub>2</sub>RR. In an experiment of CO<sub>2</sub> adsorption-desorption on the SnO<sub>2</sub>-AO<sub>10</sub> nanostructures at 298 K (Fig. 5a), the isosteric heats of adsorption ( $Q_{st}$ ) at zero coverage is found to be as high as 50 kJ mol<sup>-1</sup> (Fig. 5b), which is the upper boundary of the CO<sub>2</sub> physical adsorption heat (25–50 kJ mol<sup>-1</sup>) and approaches the chemical adsorption heat (60–90 kJ mol<sup>-1</sup>).<sup>39,40</sup> Lastly, the chemical state of the Sn species should be also relevant, especially with the production of formate. Tin oxide has displayed remarkable activity and selectivity towards formate.<sup>41,42</sup> The Sn<sup>4+</sup> species that can mostly survive (~85%, Fig. S14†) during CO<sub>2</sub>RR are thus believed to play a crucial role in the formation of formate on SnO<sub>2</sub>-AO<sub>10</sub> aerogels. Advancing one step, we performed first-principles calculations to assess the key energy pathways of the formation of H<sub>2</sub>, CO and formate on the SnO<sub>2</sub> (110) surface (see details in the ESI†). The free energy diagram in Fig. 5c clearly shows that the production of formic acid is energetically more favorable than either the evolution of H<sub>2</sub> or the formation of CO, thus confirming the importance of the tin oxide in driving the FE toward formate.

## Conclusions

In summary, we have fabricated tubular SnO<sub>2</sub> nanostructures *via* facile anodization of an Sn plate. When tested as electrocatalysts for CO<sub>2</sub>RR, the SnO<sub>2</sub> nanostructures can achieve a relatively high current density at moderate potentials for selective and stable production of formate. The outstanding performance can be attributed to a combination of factors

related to the structural and compositional properties: (1) the high specific surface area (90.1 m<sup>2</sup> g<sup>-1</sup>) and large porosity (0.74 cm<sup>3</sup> g<sup>-1</sup>) enable easy access for CO<sub>2</sub> to the catalytic sites by amply exposing the active sites and advancing the electrolyte transfer; (2) the grain boundaries offer active sites for enhancing the CO<sub>2</sub> adsorption and tuning the binding energies of intermediates; (3) the Sn<sup>4+</sup> species together with its good preservation under CO<sub>2</sub> conditions intrinsically dictates the selectivity for formate and inhibits the H<sub>2</sub> evolution. We expect this study to provide useful guidelines for developing porous electrocatalysts for efficient CO<sub>2</sub> electroreduction.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- 1 D. D. Zhu, J. L. Liu and S. Z. Qiao, *Adv. Mater.*, 2016, **28**, 3423–3452.
- 2 J. F. Huang and R. Buonsanti, *Chem. Mater.*, 2019, **31**, 13–25.
- 3 F. J. Yu, P. H. Wei, Y. Yang, Y. H. Chen, L. M. Guo and Z. Q. Peng, *Nano Mater. Sci.*, 2019, **1**, 60–69.
- 4 J. F. Huang, N. Hörmann, E. Oveisi, A. Loiudice, G. L. DeGregorio, O. Andreussi, N. Marzari and R. Buonsanti, *Nat. Commun.*, 2019, **9**, 3117.
- 5 Y. K. Chen, K. J. Chen, J. W. Fu, A. Yamaguchi, H. M. Li, H. Pan, J. H. Hu, M. Miyauchi and M. Liu, *Nano Mater. Sci.*, 2020, DOI: 10.1016/j.nanoms.2019.10.006.
- 6 R. Kortlever, I. Peters, S. Koper and M. T. M. Koper, *ACS Catal.*, 2015, **5**, 3916–3923.
- 7 T. N. Huan, P. Simon, G. Rousse, I. Genois, V. Artero and M. Fontecave, *Chem. Sci.*, 2017, **8**, 742–747.
- 8 S. B. Liu, J. Xiao, X. F. Lu, J. Wang, X. Wang and X. W. Lou, *Angew. Chem., Int. Ed.*, 2019, **58**, 8499–8503.
- 9 W.-H. Wang, Y. Himeda, J. T. Muckerman, G. F. Manbeck and E. Fujita, *Chem. Rev.*, 2015, **115**, 12936–12973.
- 10 Y. C. Li, Z. Wang, T. Yuan, D.-H. Nam, M. Luo, J. Wicks, B. Chen, J. Li, F. Li, F. P. G. de Arguer, Y. Wang, C.-T. Dinh, O. Voznyy, D. Sinton and E. H. Sargent, *J. Am. Chem. Soc.*, 2019, **141**, 8584–8591.
- 11 T. He, P. Pachfule, H. Wu, Q. Xu and P. Chen, *Nature Rev. Mater.*, 2016, **1**, 16059.
- 12 J.-P. Jones, G. K. S. Prakash and G. A. Olah, *Israel J. Chem.*, 2014, **54**, 1451–1466.
- 13 Y. Zheng, A. Vasileff, X. Zhou, Y. Jiao, M. Jaroniec and S.-Z. Qiao, *J. Am. Chem. Soc.*, 2019, **141**, 7646–7659.



- 14 S. Zhang, P. Kang and T. J. Meyer, *J. Am. Chem. Soc.*, 2014, **136**(5), 1734–1737.
- 15 F. C. Lei, W. Liu, Y. F. Sun, J. Q. Xu, K. T. Liu, L. Liang, T. Yao, B. C. Pan, S. Q. Wei and Y. Xie, *Nat. Commun.*, 2016, **7**, 12697.
- 16 F. Li, L. Chen, G. P. Knowles, D. R. MacFarlane and J. Zhang, *Angew. Chem., Int. Ed.*, 2017, **56**, 505–509.
- 17 B. Kumar, V. Atla, J. P. Brian, S. Kumari, T. Q. Nguyen, M. Sunkara and J. M. Spurgeon, *Angew. Chem., Int. Ed.*, 2017, **56**, 3645–3649.
- 18 S. Liu, F. Pang, Q. Zhang, R. Guo, Z. Wang, Y. Wang, W. Zhang and J. Ou, *Appl. Mater. Today*, 2018, **13**, 135–143.
- 19 X. Wang, J. Lv, J. X. Zhang, X. L. Wang, C. Z. Xue, G. Q. Bian, D. S. Li, Y. Wang and T. Wu, *Nanoscale*, 2020, **12**, 772–784.
- 20 Y.-W. Choi, F. Scholten, I. Sinev and B. R. Cuenya, *J. Am. Chem. Soc.*, 2019, **141**, 5261–5266.
- 21 J. Wang, Y. Ji, Q. Shao, R. Yin, J. Guo, Y. Li and X. Huang, *Nano Energy*, 2019, **59**, 138–145.
- 22 H. C. Shin, J. Dong and M. Liu, *Adv. Mater.*, 2004, **16**, 237–240.
- 23 K. Jiang, Y. F. Huang, G. S. Zeng, F. M. Toma, W. A. Goddard III and A. T. Bell, *ACS Energy Lett*, 2020, **5**, 1206–1214.
- 24 J. Vandevondele, M. Krack, F. Mohamed, M. Parrinello, T. Chassaing and J. Hutter, *Comput. Phys. Commun.*, 2005, **167**, 103–128.
- 25 CP2K version 4.1, CP2K is freely available from, <http://www.cp2k.org>.
- 26 B. Gerdlippert and J. Huttermicheleparrinello, *Mol. Phys.*, 1997, **92**, 477–488.
- 27 S. Grimme, J. Antony, S. Ehrlich and H. Krieg, *J. Chem. Phys.*, 2010, **132**, 154104–154119.
- 28 M. Fronzi, S. Piccinin, B. Delley, E. Traversa and C. Stampfl, *Phys. Chem. Chem. Phys.*, 2009, **11**, 9188–9199.
- 29 V. Shapovalov and H. Metiu, *J. Catal.*, 2007, **245**, 205–214.
- 30 S. Goedecker, M. Teter and J. Hutter, *Phys. Rev. B*, 1995, **54**, 1703–1710.
- 31 A. Ghysels, T. Verstraelen, K. Hemelsoet, M. Waroquier and S. V. Van, *J. Chem. Inf. Model.*, 2010, **50**, 1736–1750.
- 32 J. Gu, F. Heroguel, J. Luterbacher and X. Hu, *Angew. Chem., Int. Ed.*, 2018, **57**, 2943–2947.
- 33 M. Thommes, K. Kaneko, A. V. Neimark, J. P. Olivier, F. Rodriguez-Reinoso, J. Rouquerol and K. S. W. Sing, *Pure Appl. Chem.*, 2015, **87**, 1051–1069.
- 34 R. Daiyan, X. Lu, W. H. Saputera, Y. H. Ng and R. Amal, *ACS Sustainable Chem. Eng.*, 2018, **6**, 1670–1679.
- 35 L. Fan, X. Li, B. Yan, J. Feng, D. Xiong, D. Li, L. Gu, Y. Wen, S. Lawes and X. Sun, *Adv. Energy Mater.*, 2016, **6**, 1502057.
- 36 H. Ge, Z. Gu, P. Han, H. Shen, A. M. Al-Enizi, L. Zhang and G. Zheng, *J. Colloid Interface Sci.*, 2018, **531**, 564–569.
- 37 C. W. Li, J. Ciston and M. W. Kanan, *Nature*, 2014, **508**, 504.
- 38 K.-S. Kim, W. J. Kim, H.-K. Lim, E. K. Lee and H. Kim, *ACS Catal.*, 2016, **6**, 4443–4448.
- 39 Y.-S. Bae and R. Q. Snurr, *Angew. Chem., Int. Ed.*, 2011, **50**, 11586–11596.
- 40 T. Watabe and K. Yogo, *Sep. Purif. Technol.*, 2013, **120**, 20–23.
- 41 Y. Chen and M. W. Kanan, *J. Am. Chem. Soc.*, 2012, **134**, 1986–1989.
- 42 Q. Li, J. Fu, W. Zhu, Z. Chen, B. Shen, L. Wu, Z. Xi, T. Wang, G. Lu, J.-j. Zhu and S. Sun, *J. Am. Chem. Soc.*, 2017, **139**, 4290–4293.

