

## PAPER

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[View Journal](#) | [View Issue](#)Cite this: *J. Mater. Chem. A*, 2022, 10, 20018A hexacoordinated Bi<sup>3+</sup>-based ellagate MOF with acid/base resistance boosting carbon dioxide electroreduction to formate†Junjun Li,<sup>‡a</sup> Congyong Wang,<sup>‡bd</sup> Dingjia Wang,<sup>‡c</sup> Chenhuai Yang,<sup>‡a</sup> Xiaoya Cui,<sup>ef</sup> Xuejiao J. Gao<sup>ib\*ac</sup> and Zhicheng Zhang<sup>ib\*aa</sup>

Electrochemical conversion of CO<sub>2</sub> into energy-dense liquids, such as formic acid, as hydrogen carriers and chemical feedstocks is desirable. However, the development of highly efficient and stable electrocatalysts toward formate production in a wide potential window still remains a great challenge. Herein, rod-shaped bismuth ellagate metal–organic framework (referred to as SU-101) catalysts were demonstrated to exhibit excellent activity, selectivity, and stability for the conversion of CO<sub>2</sub> into HCOO<sup>−</sup>. In particular, the SU-101 nanorods (NRs) achieved a high HCOO<sup>−</sup> Faraday efficiency (FE) up to 93.66% at −1.10 V vs. RHE and a partial current density of −14.57 mA cm<sup>−2</sup>. Significantly, the FE of HCOO<sup>−</sup> could be maintained over 87.51% during 10 hours of continuous electrolysis without significant deterioration. The results indicated that the excellent performance could be attributed to the special structure of Bi<sub>2</sub>O(H<sub>2</sub>O)<sub>2</sub>(C<sub>14</sub>H<sub>2</sub>O<sub>8</sub>)·*n*H<sub>2</sub>O (SU-101). DFT calculations show that the unique hexacoordinated Bi<sup>3+</sup> site of SU-101 is highly efficient for formate production with very low overpotentials. This work paves a way to develop a Bi-MOF as a highly stable, molecularly tunable catalytic material for selective CO<sub>2</sub> reduction to specific valuable chemicals.

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## Introduction

The electrochemical CO<sub>2</sub> reduction reaction (CO<sub>2</sub>RR) has been considered one of the most attractive pathways to convert CO<sub>2</sub> into useful feedstocks and fuels, which can mitigate environmental issues and alleviate the energy crisis.<sup>1–6</sup> Formic acid or formate, as a common liquid CO<sub>2</sub> reduction product, has been widely applied in livestock feed as a preserving and antibacterial agent, as liquid fuel in direct formate fuel cells, and in textile and leather production.<sup>7–9</sup> Significant efforts have been devoted to exploring and designing novel electrocatalysts for producing value-added products, such as formate.<sup>1,5,10,11</sup>

In recent years, various metal-based electrocatalysts, such as Bi, In, Sn, Pb, Pd, Cu, Hg, *etc.*,<sup>12–21</sup> have been proven to be active in reducing CO<sub>2</sub> into formate. Notably, Bi-based catalysts have been considered suitable candidates owing to their captivating characteristics: (i) compared to the toxic metals of Pb and Hg, Bi-based electrocatalysts display lower toxicity and an

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environmentally benign nature; (ii) their high hydrogen evolution overpotential is beneficial for promoting CO<sub>2</sub> reduction; (iii) Bi-based electrocatalysts exhibit strong affinity toward OCHO intermediates, giving rise to high selectivity for formate production; (iv) their low cost and good durability can facilitate their practical application. Recently, Bi-based metal–organic frameworks (MOFs) have attracted considerable interest for electrocatalysis due to their readily accessible active sites, tailored porosity, convenient functionalization, and large specific surface area.<sup>22–27</sup> To date, there have been very few reports on the application of MOFs in the CO<sub>2</sub>RR, thus deserving an in-depth exploration and understanding of Bi-based MOF electrocatalysts.<sup>28,29</sup> However, the reported Bi-based MOFs usually suffer from structural degradation and rearrangement at reducing potentials during the CO<sub>2</sub>RR process,<sup>6,12,30</sup> which would lead to the complexity of the study of the catalytic mechanism. Therefore, the rational design and synthesis of Bi-based MOFs with robust structural stability is of

great significance to the development of highly efficient electrocatalysts for the conversion of CO<sub>2</sub> into HCOO<sup>−</sup>.

To effectively tackle this issue, we fabricated highly stable Bi<sub>2</sub>O(H<sub>2</sub>O)<sub>2</sub>(C<sub>14</sub>H<sub>2</sub>O<sub>8</sub>)·*n*H<sub>2</sub>O·*n*H<sub>2</sub>O (SU-101) nanorod (NR) catalysts *via* a facile wet chemical method at room temperature. Impressively, the crystalline structure of the SU-101 NRs could be well maintained over a large pH range (from 2 to 14). The as-prepared SU-101 NRs exhibited high activity and selectivity toward HCOO<sup>−</sup> for electrochemical CO<sub>2</sub> reduction under ambient conditions, with an excellent faradaic efficiency (FE) up to 93.66% and a partial current density as high as 14.57 mA cm<sup>−2</sup> at −1.06 V *versus* the reversible hydrogen electrode (*vs.* RHE). Moreover, it was also found that the SU-101 NRs are capable of retaining their good structure and chemical stability after a long-term electrocatalytic process. DFT calculation results indicated that the outstanding performance could be attributed to the unique hexacoordinated Bi<sup>3+</sup> site of SU-101. The adsorption of CO<sub>2</sub> on the central Bi site is mainly

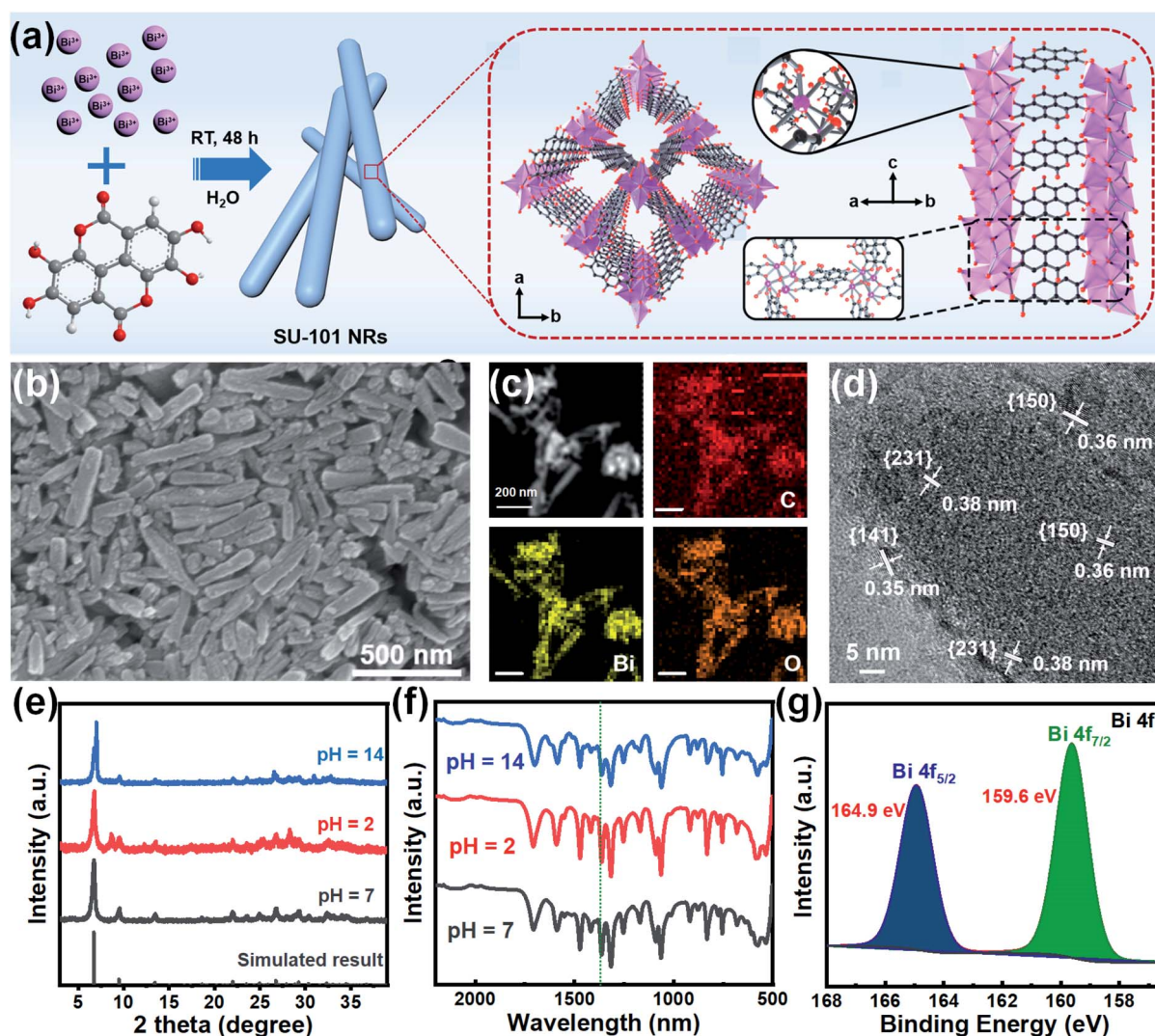


Fig. 1 (a) Schematic illustration of the preparation procedure of the SU-101 NRs. (b) SEM image of the SU-101 NRs. (c) HAADF-STEM and the corresponding EDS elemental mapping images of Bi, O, and C. (d) HRTEM image of the SU-101 NRs. (e) PXRD patterns and (f) FT-IR spectra of the SU-101 NRs treated with different pH solutions. (g) Bi XPS spectrum of the SU-101 NRs.

achieved by coordination interaction. The active Bi sites exhibit moderate adsorption strength, which is neither so weak as to allow the hydrogen evolution reaction (HER) to occur preferentially nor so strong as to cause CO formation, resulting in high selectivity toward  $\text{HCOO}^-$  production.

## Results and discussion

The SU-101 NRs were synthesized in the presence of ellagic acid, bismuth acetate, and glacial acetic acid *via* a facile wet chemical method, as schematically illustrated in Fig. 1a and S1.† Scanning electron microscope (SEM) and transmission electron microscopy (TEM) images show the rod-like morphology with an average length and width of 334.9 nm and 69.2 nm, respectively (Fig. 1b, S2 and S3†). High angle annular dark-field scanning TEM (HAADF-STEM) and corresponding energy dispersive spectroscopy (EDS) element mapping images of the as-synthesized SU-101 NRs confirm the homogeneous distribution of Bi, C, and O elements (Fig. 1c). The HRTEM image of the SU-101 NRs shows obvious lattice fringes with lattice spacings of 0.35 nm, 0.36 nm and 0.38 nm, which can be assigned to the {141}, {150}, and {231} crystal planes of SU-101 (Fig. 1d). Their powder X-ray diffraction (PXRD) patterns (Fig. 1e) show the typical diffraction peaks of SU-101, which are reasonably consistent with the simulated results.<sup>31</sup> Besides, the samples display strong diffraction peaks, indicating good crystallinity of the SU-101 NRs. More importantly, the crystalline structure of the SU-101 NRs could be well maintained over a wide pH region ranging from 2 to 14. The Fourier transform infrared (FT-IR) spectra indicated that there is coordination interaction between the metallic ions and hydroxyl group of ellagic acid, as shown by an obvious red shift in the C–OH stretching to 1363 nm from 1322 nm for the uncoordinated building blocks of ellagic acid (Fig. 1f).<sup>32</sup> It is worth mentioning that similar FT-IR spectra of the SU-101 NRs treated with different pH solutions could be observed, further indicating that the SU-101 NRs possess excellent chemical stability. X-ray photoelectron spectroscopy (XPS) characterization was also utilized to study the chemical composition and surface chemical state of the as-fabricated SU-101 NRs. The survey spectrum of the SU-101 NRs exhibited typical spectrum characteristics of Bi, C, O, and N elements (Fig. S4†). As exhibited in Fig. 1g, the binding energies of Bi 4f were located at 159.6 eV and 164.9 eV, which could be assigned to Bi 4f<sub>5/2</sub> and Bi 4f<sub>7/2</sub>, respectively, suggesting that the oxidation state of Bi is +3. Thermogravimetric analysis (TGA) demonstrates that the SU-101 NRs show high thermal stability up to 300 °C (Fig. S5†). In addition, it was calculated that the Brunauer–Emmett–Teller surface area and average pore diameter of the SU-101 NRs are 25.96 m<sup>2</sup> g<sup>−1</sup> and 20.43 nm, respectively (Fig. S6†).

The electrocatalytic CO<sub>2</sub>RR performances of the SU-101 NRs were investigated in a typical three-electrode H-type cell. To confirm the reduction products, the liquid and the gas products were analyzed by nuclear magnetic resonance (NMR) spectroscopy (Fig. S7†) and online gas chromatography (GC) (Fig. S8†), respectively. The linear sweep voltammetry (LSV) curves of the SU-101 NRs in CO<sub>2</sub>/N<sub>2</sub>-saturated 0.5 M KHCO<sub>3</sub> aqueous solution

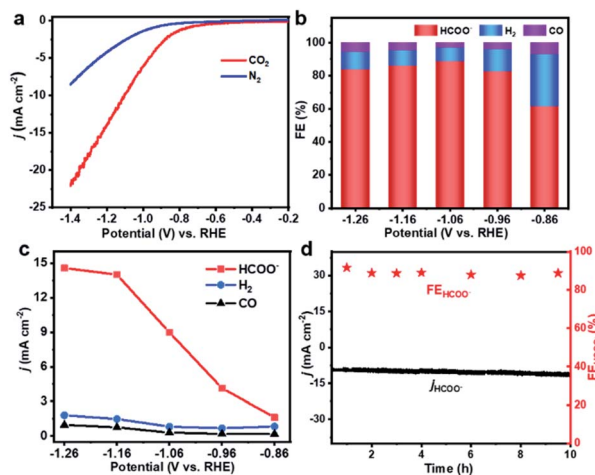


Fig. 2 Electrocatalytic CO<sub>2</sub>RR performance of the SU-101 NRs. (a) LSV curves in N<sub>2</sub>- (blue line) or CO<sub>2</sub>- (red line) saturated 0.5 M KHCO<sub>3</sub> aqueous solution at a scan rate of 5 mV s<sup>−1</sup>. (b) FEs of H<sub>2</sub>, CO and HCOO<sup>−</sup> at different applied potentials. (c) Partial current densities of H<sub>2</sub>, CO and HCOO<sup>−</sup>. (d) Total current density and faradaic efficiency toward HCOO<sup>−</sup> over 10 h electrolysis at −1.06 V vs. RHE.

at a scan rate of 5 mV s<sup>−1</sup> are presented in Fig. 2a. In N<sub>2</sub>-saturated 0.5 M KHCO<sub>3</sub> electrolyte, the increase in the current below  $\sim -0.70$  V vs. RHE could be ascribed to the HER which is the main competing reaction during the CO<sub>2</sub>RR. With the CO<sub>2</sub>-purged 0.5 M KHCO<sub>3</sub> electrolyte, a dramatic increase of reductive current is observed at a potential of  $-0.56$  V vs. RHE, which indicates the predominant occurrence of CO<sub>2</sub> reduction over the SU-101 catalyst.<sup>33</sup> Obviously, the SU-101 NRs displayed a higher cathode current density in CO<sub>2</sub>-saturated solution than in N<sub>2</sub>-saturated solution, manifesting their remarkable catalytic performances towards the CO<sub>2</sub>RR. The conversion of CO<sub>2</sub> into HCOO<sup>−</sup> over the SU-101 catalyst occurred in the range from  $-0.86$  to  $-1.26$  V vs. RHE (Fig. 2b). Impressively, the highest FE of the SU-101 NRs could reach up to 93.66% at  $\sim -1.06$  V vs. RHE and a maximum formate partial current density ( $j_{\text{HCOO}^-}$ ) of 14.57 mA cm<sup>−2</sup> could be achieved at  $-1.26$  V vs. RHE (Fig. 2c), which are comparable or even superior to those of recently reported Bi-based and other advanced catalysts (Fig. 3 and Table

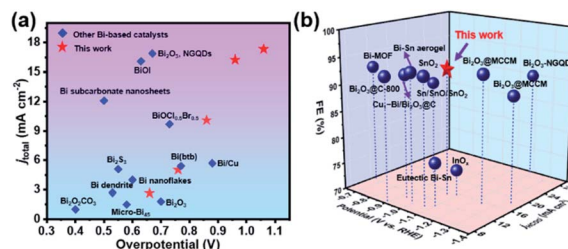


Fig. 3 (a) Literature overview of Bi-based electrocatalysts for the CO<sub>2</sub>-to-formate conversion. Overpotentials ( $\eta$ ) calculated using the equilibrium potential for CO<sub>2</sub>/HCOO<sup>−</sup>,  $E = -0.2$  V vs. RHE.<sup>34</sup> (b) Comparison of partial current density and FE of formate for the SU-101 NRs with those of previously reported Bi-based and other advanced catalysts at the applied potentials.

S1†). Moreover, substantial quantities of  $\text{HCOO}^-$  are produced with FE over 80.0% over a wide applied cathodic potential region ranging from  $\sim -0.96$  to  $\sim -1.26$  V vs. RHE. The long-term stability characterization of the SU-101 NRs was carried out at a constant potential of  $-1.06$  V vs. RHE for 10 h. As presented in Fig. 2d, the FE of  $\text{HCOO}^-$  could be maintained over 87.51% during 10 hours of continuous electrolysis and there was almost no evident deterioration in the current density and the NRs were well maintained after  $\text{CO}_2\text{RR}$  measurement (Fig. S9†), and the PXRD spectrum shows that the crystalline structure of the SU-101 NRs remains almost unchanged after the long-term stability test (Fig. S10†). The above-mentioned results demonstrate the superior structural robustness of the SU-101 NRs for the  $\text{CO}_2\text{RR}$ . In addition, the electrocatalytic  $\text{CO}_2\text{RR}$  performances of the SU-101 NRs in a flow cell were also investigated (Fig. S11–S13†). Note that the highest FE of the SU-101 NRs could reach up to 88.25% at  $\sim -1.06$  V vs. RHE and the partial current density towards formate is  $40.46 \text{ mA cm}^{-2}$  (Fig. S12†).

DFT calculations were performed to explore the possible molecular mechanisms for the  $\text{CO}_2\text{RR}$  and HER catalytic activities of SU-101 and gain insights into its high selectivity for

formic acid production. As shown in Fig. 4, the crystal structure of SU-101 can be regarded as  $\text{Bi}_2\text{O}$  nanowires connected by  $(\text{C}_{14}\text{H}_2\text{O}_8)$  linkers in four directions, and its chemical composition is  $\text{Bi}_2\text{O}(\text{H}_2\text{O})_2(\text{C}_{14}\text{H}_2\text{O}_8)$ . To save computational cost, we constructed a nanowire model, simplifying the two linkers on one side of the  $\text{Bi}_2\text{O}$  wire to  $(\text{C}_2\text{H}_2\text{O}_2)$ , while leaving the two  $(\text{C}_{14}\text{H}_2\text{O}_8)$  linkers connected to the reaction site (Fig. 4b). The linker terminal oxygen is saturated with H and thus the chemical composition of the constructed nanowire is  $\text{Bi}_2\text{O}(\text{H}_2\text{O})_2(-\text{C}_{14}\text{H}_2\text{O}_8)(\text{C}_2\text{H}_2\text{O}_2)$ , which preserves the valence state of  $\text{Bi}^{3+}$  and its chemical environment. Three possible pathways were proposed for the  $\text{CO}_2\text{RR}$  (paths 1–3 in Fig. 4c). Path 1 leads to the production of CO, paths 2 and 3 lead to the yield of  $\text{HCOOH}$  and path 4 represents the HER pathway. The  $\text{Bi}^{3+}$  in the active sites is hexacoordinated and one of the ligands is the water molecule (Fig. 4b), and thus the active site is denoted as  $\text{H}_2\text{O}^*$  (\* represents the adsorption state). The energy profiles in Fig. 4b indicate that the first addition of H is the rate-determining step (RDS) and path 3 is the most favourable way leading to the generation of  $\text{HCOOH}$  since it has an energy barrier of only 0.87 eV for the RDS which is significantly lower than that of paths 1 and 2 (1.64 eV). The comparison of path 1 with path 2

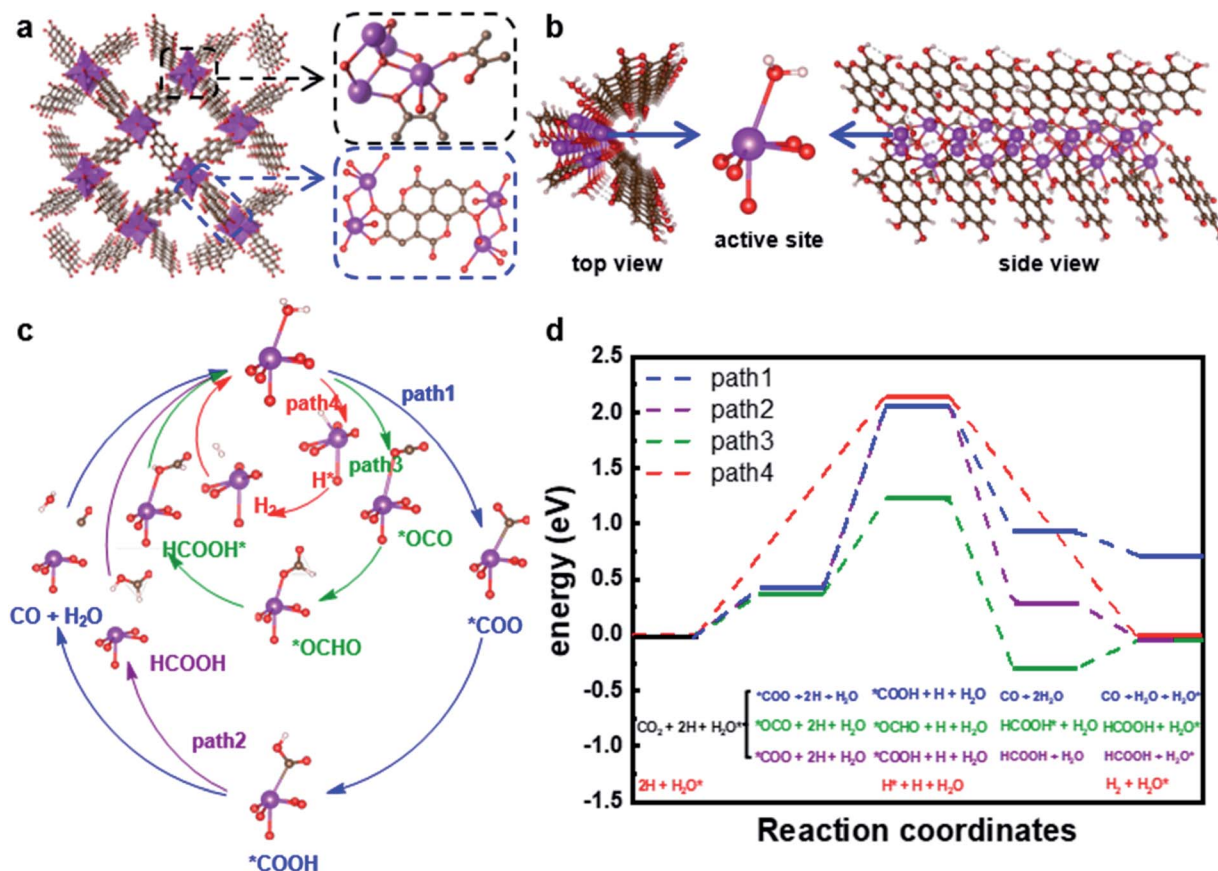


Fig. 4 (a) The crystal structure of the SU-101 and enlarged local structural details for the coordination microenvironment of Bi metal and the connection between Bi and the  $(\text{C}_{14}\text{H}_2\text{O}_8)$  linker. (b) The top and side views for the constructed nanowire periodic structure model mimicking SU-101 and the active site for the  $\text{CO}_2\text{RR}$  and HER. (c) Four proposed reaction pathways on the active site leading to different products which correspond to the  $\text{CO}_2\text{RR}$  and HER. (d) The energy profiles in eV for the four pathways proposed in (c) with material compositions provided.

also indicates the better selectivity for HCOOH production, which is rooted in that the adsorption of  $^*COOH$  or  $(^*CO_2)$  on the  $Bi^{3+}$  site is not strong enough to weaken the carbon–oxygen double bond, thus leading to the generation of hardly any CO. Path 4 responsible for the HER has the highest energy barrier of 2.13 eV, which indicates that the generation of  $H_2$  is unfavourable. Thus, the easier adsorption of  $CO_2$  than H makes the  $CO_2RR$  a more advantageous reaction than the HER, from which it is understandable that the O atom in  $CO_2$  has stronger coordination capability than H. Moreover, as illustrated in Fig. S6,† the SU-101 catalyst exhibited a contact angle of  $145.5^\circ$  (Fig. S14†), demonstrating its inherent hydrophilic character which could contribute to inhibition of the competitive HER. The above results consistently indicate the high selectivity for HCOOH formation on SU-101, which corresponds well with the experimental results. This can be explained by two aspects: on one hand, the easier adsorption of  $CO_2$  than H makes the  $CO_2RR$  a more advantageous reaction than the HER;<sup>35,36</sup> on the other hand, the adsorption of  $CO_2$  on SU-101 is not strong enough to weaken the C=O bond and thus the CO generation is not favourable.

## Conclusions

In summary, rod-shaped SU-101 catalysts were synthesized *via* a wet chemical method. Importantly, the crystalline structure of the SU-101 NRs could be well maintained over a wide pH region ranging from 2 to 14. The as-prepared SU-101 NRs exhibited high activity and selectivity toward  $HCOO^-$  for electrochemical  $CO_2$  reduction under ambient conditions. An exceptional FE of 93.66% for  $HCOO^-$  production was achieved at a potential of  $-1.06$  V vs. RHE with an outstanding stability over 10 h. Theoretical calculations confirm that the suitable adsorption strength of SU-101 to  $CO_2$  is the reason for its high selectivity for  $HCOO^-$  production: neither so weak as to allow the HER to occur preferentially, nor so strong as to cause CO formation. This work offers new opportunities for developing and designing highly stable framework structures for the electrochemical conversion of  $CO_2$  into value-added products.

## Author contributions

Junjun Li, Congyong Wang, Dingjia Wang and Chenhuai Yang contributed equally to this work.

## Conflicts of interest

There are no conflicts to declare.

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## References

- 1 K. Sordakis, C. Tang, L. K. Vogt, H. Junge, P. J. Dyson, M. Beller and G. Laurenczy, *Chem. Rev.*, 2018, **118**, 372–433.
- 2 D. H. Nam, P. De Luna, A. Rosas-Hernandez, A. Thevenon, F. Li, T. Agapie, J. C. Peters, O. Shekhah, M. Eddaoudi and E. H. Sargent, *Nat. Mater.*, 2020, **19**, 266–276.
- 3 P. Deng, F. Yang, Z. Wang, S. Chen, Y. Zhou, S. Zaman and B. Y. Xia, *Angew. Chem., Int. Ed.*, 2020, **59**, 10807–10813.
- 4 N. Wang, K. Yao, A. Vomiero, Y. Wang and H. Liang, *SmartMat*, 2021, **2**, 423–425.
- 5 Y. Wang, X. Zheng and D. Wang, *Nano Res.*, 2022, **15**, 1730–1752.
- 6 S. Chen, W.-H. Li, W. Jiang, J. Yang, J. Zhu, L. Wang, H. Ou, Z. Zhuang, M. Chen, X. Sun, D. Wang and Y. Li, *Angew. Chem., Int. Ed.*, 2022, **61**, e202114450.
- 7 J. Li, S. U. Abbas, H. Wang, Z. Zhang and W. Hu, *Nano-Micro Lett.*, 2021, **13**, 216.
- 8 Z. Gao, J. Li, Z. Zhang and W. Hu, *Chin. Chem. Lett.*, 2022, **33**, 2270–2280.
- 9 Y. Zhu, X. Cui, H. Liu, Z. Guo, Y. Dang, Z. Fan, Z. Zhang and W. Hu, *Nano Res.*, 2021, **14**, 4471–4486.
- 10 Z. Gao, C. Wang, J. Li, Y. Zhu, Z. Zhang and W. Hu, *Acta Phys.-Chim. Sin.*, 2021, **37**, 2010025.
- 11 P. Lamagni, M. Miola, J. Catalano, M. S. Hvid, M. A. H. Mamakhel, M. Christensen, M. R. Madsen, H. S. Jeppesen, X. M. Hu, K. Daasbjerg, T. Skrydstrup and N. Lock, *Adv. Funct. Mater.*, 2020, **30**, 1910408.
- 12 J. Yang, X. Wang, Y. Qu, X. Wang, H. Huo, Q. Fan, J. Wang, L. M. Yang and Y. Wu, *Adv. Energy Mater.*, 2020, **10**, 2001709.
- 13 M. Zhang, W. Wei, S. Zhou, D.-D. Ma, A. Cao, X.-T. Wu and Q.-L. Zhu, *Energy Environ. Sci.*, 2021, **14**, 4998–5008.
- 14 Z. H. Zhu, B. H. Zhao, S. L. Hou, X. L. Jiang, Z. L. Liang, B. Zhang and B. Zhao, *Angew. Chem., Int. Ed.*, 2021, **60**, 23394–23402.
- 15 Z. Wu, H. Wu, W. Cai, Z. Wen, B. Jia, L. Wang, W. Jin and T. Ma, *Angew. Chem., Int. Ed.*, 2021, **60**, 12554–12559.
- 16 J. Li, Z. Zhang and W. Hu, *Green Energy Environ.*, 2022, DOI: [10.1016/j.gee.2021.11.004](https://doi.org/10.1016/j.gee.2021.11.004).
- 17 T. Zheng, C. Liu, C. Guo, M. Zhang, X. Li, Q. Jiang, W. Xue, H. Li, A. Li, C. W. Pao, J. Xiao, C. Xia and J. Zeng, *Nat. Nanotechnol.*, 2021, **16**, 1386–1393.
- 18 H. Yang, N. Han, J. Deng, J. Wu, Y. Wang, Y. Hu, P. Ding, Y. Li, Y. Li and J. Lu, *Adv. Energy Mater.*, 2018, **8**, 1801536.
- 19 N. Han, P. Ding, L. He, Y. Li and Y. Li, *Adv. Energy Mater.*, 2019, **10**, 1902338.
- 20 E. Zhang, T. Wang, K. Yu, J. Liu, W. Chen, A. Li, H. Rong, R. Lin, S. Ji, X. Zheng, Y. Wang, L. Zheng, C. Chen, D. Wang, J. Zhang and Y. Li, *J. Am. Chem. Soc.*, 2019, **141**, 16569–16573.
- 21 K. Ye, Z. Zhou, J. Shao, L. Lin, D. Gao, N. Ta, R. Si, G. Wang and X. Bao, *Angew. Chem., Int. Ed.*, 2020, **59**, 4814–4821.
- 22 H. Wu, M. Zeng, X. Zhu, C. Tian, B. Mei, Y. Song, X.-L. Du, Z. Jiang, L. He, C. Xia and S. Dai, *ChemElectroChem*, 2018, **5**, 2717–2721.

- 23 M. K. Birhanu, M.-C. Tsai, A. W. Kahsay, C.-T. Chen, T. S. Zeleke, K. B. Ibrahim, C.-J. Huang, W.-N. Su and B.-J. Hwang, *Adv. Mater. Interfaces*, 2018, **5**, 1800919.
- 24 Y. Wang, P. Hou, Z. Wang and P. Kang, *ChemPhysChem*, 2017, **18**, 3142–3147.
- 25 N. Devarajan and P. Suresh, *Org. Chem. Front.*, 2018, **5**, 2322–2331.
- 26 W. Guo, X. Sun, C. Chen, D. Yang, L. Lu, Y. Yang and B. Han, *Green Chem.*, 2019, **21**, 503–508.
- 27 Q. Mu, W. Zhu, G. Yan, Y. Lian, Y. Yao, Q. Li, Y. Tian, P. Zhang, Z. Deng and Y. Peng, *J. Mater. Chem. A*, 2018, **6**, 21110–21119.
- 28 F. Li, G. H. Gu, C. Choi, P. Kolla, S. Hong, T.-S. Wu, Y.-L. Soo, J. Masa, S. Mukerjee, Y. Jung, J. Qiu and Z. Sun, *Appl. Catal., B*, 2020, **277**, 119241.
- 29 K. Ye, A. Cao, J. Shao, G. Wang, R. Si, N. Ta, J. Xiao and G. Wang, *Sci. Bull.*, 2020, **65**, 711–719.
- 30 C. Cao, D. D. Ma, J. F. Gu, X. Xie, G. Zeng, X. Li, S. G. Han, Q. L. Zhu, X. T. Wu and Q. Xu, *Angew. Chem., Int. Ed.*, 2020, **59**, 15014–15020.
- 31 E. S. Grape, J. G. Flores, T. Hidalgo, E. Martinez-Ahumada, A. Gutierrez-Alejandre, A. Hautier, D. R. Williams, M. O'Keeffe, L. Ohrstrom, T. Willhammar, P. Horcajada, I. A. Ibarra and A. K. Inge, *J. Am. Chem. Soc.*, 2020, **142**, 16795–16804.
- 32 H. Wang, X. Xu, C. Lee, C. Johnson, K. Sohlberg and H.-F. Ji, *J. Phys. Chem. C*, 2012, **116**, 4442–4448.
- 33 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.
- 34 C. W. Li and M. W. Kanan, *J. Am. Chem. Soc.*, 2012, **134**, 7231–7234.
- 35 L. Liu, K. Yao, J. Fu, Y. Huang, N. Li and H. Liang, *Colloids Surf., A*, 2022, **633**, 127840.
- 36 Y. Xue, C. Li, X. Zhou, Z. Kuang, W. Zhao, Q. Zhang and H. Chen, *ChemElectroChem*, 2022, **9**, e202101648.