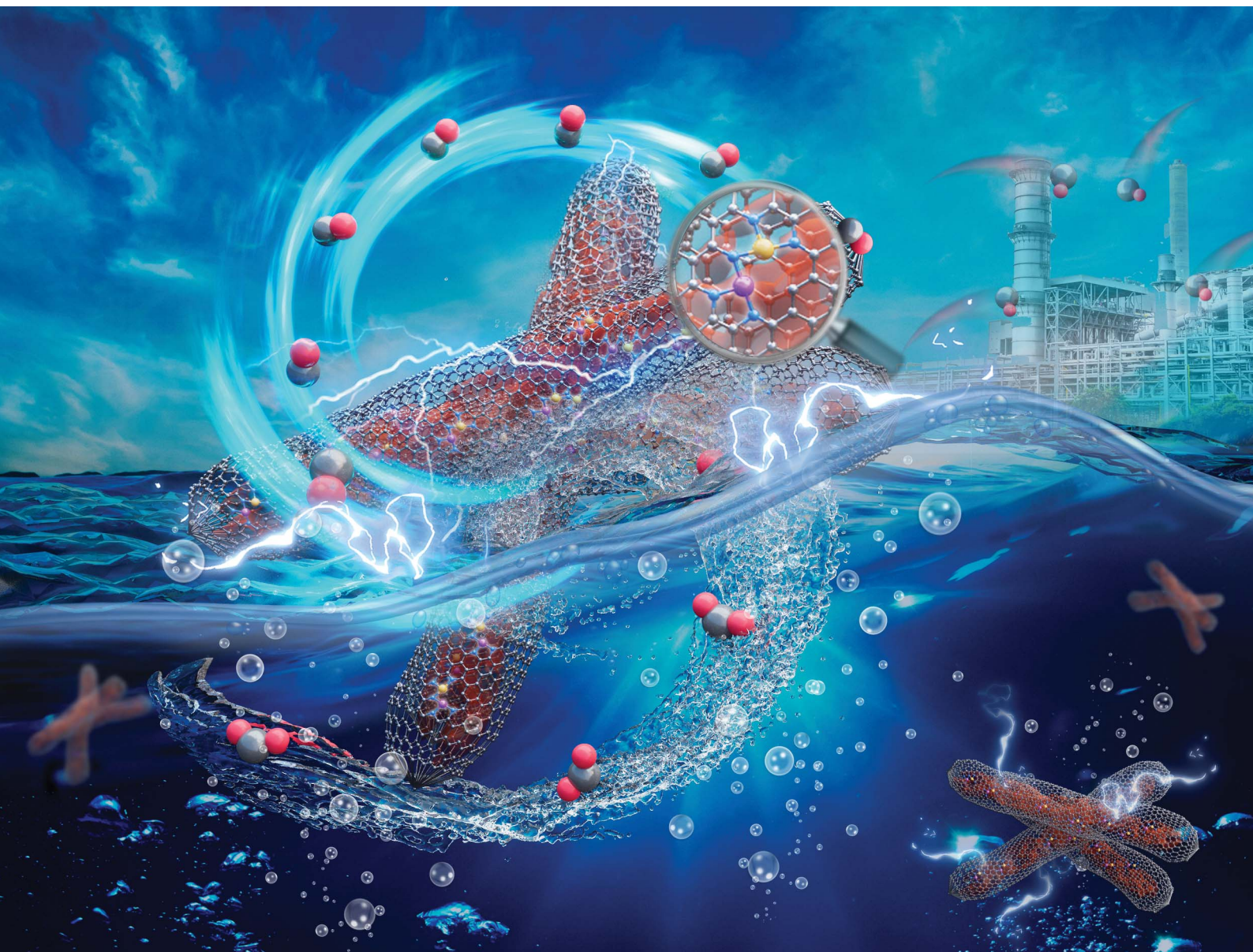


Journal of Materials Chemistry A

Materials for energy and sustainability

rsc.li/materials-a



ISSN 2050-7488

PAPER

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Cite this: *J. Mater. Chem. A*, 2024, 12, 13694

Low-coordinated Co–Mn diatomic sites derived from metal–organic framework nanorods promote electrocatalytic CO₂ reduction†

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The manipulation of the geometric coordination structure of diatomic site (DS) catalysts to promote the CO₂-to-CO conversion process has garnered significant attention. However, the challenge remains in how to rationally design the catalytic microenvironment of DSs to enhance the kinetics of CO product formation. Herein, we present a post-synthetic co-substitution (PSCR) method for producing Co–Mn DS catalysts with low N coordination numbers (referred to as L-Co₁Mn₁-NC) on pre-designed N-doped carbon derived from metal–organic framework nanorods. When utilized in the process of CO₂ electroreduction, the L-Co₁Mn₁-NC catalyst demonstrates a CO faradaic efficiency (FE) of up to 97.6% at –0.47 V, which is significantly higher compared to those of Co₁Mn₁-NC, Co₁-NC, and Mn₁-NC catalysts. *In situ* ATR-SEIRAS and theoretical simulations demonstrate that the creation of Co–Mn DSs with a lower N coordination number can notably facilitate the desorption of CO*, thus expediting the kinetics of the CO₂-to-CO conversion process. This study introduces a novel approach to fine-tune the catalytic microenvironment of multi-atomic sites in order to facilitate the conversion and utilization of CO₂.

Received 3rd April 2024

Accepted 20th May 2024

DOI: 10.1039/d4ta02261k

rsc.li/materials-a

Introduction

The electrochemical reduction of CO₂ to produce high-value fuels and fine chemicals (CO₂RR), powered by electricity generated from renewable clean energy sources, holds significant importance for mitigating environmental issues and

achieving global carbon neutrality.^{1,2} However, challenges persist in the CO₂RR process, such as the intrinsic CO₂ inertness, complex reaction environments, difficulties in controlling the surface interface, and competitive hydrogen evolution reaction (HER).^{3,4} The key to addressing these challenges lies in the development of highly efficient electrocatalysts with

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† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4ta02261k>

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superior catalytic activity, stability, and targeted control over product selectivity.^{5,6} In recent years, single-atom catalysts (SACs), typically atomically dispersed metal–nitrogen–carbon (M–N–C) materials, have sparked a research boom in the CO₂RR process.^{7–10} This is due to their unique chemical environment properties, and they are anticipated to facilitate the large-scale development of electrochemical CO₂-to-CO conversion.^{11–13} Generally, in M–N–C materials, the coordination-saturated M–N₄ moiety is a common structure that determines the activity of CO₂ reduction.¹⁴ Although considerable progress has been made in the CO₂RR application of SACs with the M–N₄ moiety,^{15,16} the highly symmetrical M–N₄ configuration actually restrains the modulation of the electronic structure of the metal center, thereby limiting its potential applications.

Inspired by the fact that single-atom metal centers are stabilized on the carrier by coordination interactions with anchoring nodes based on coordination chemistry, the catalytic activity of SACs is closely related to their catalytic coordination microenvironment.^{17,18} The coordination number of the first shell is a significant indicator that influences the local electronic properties of the catalytic microenvironment in SACs, and can be considered a key factor determining catalytic activity and selectivity.^{19,20} Specifically, altering the coordination number of neighboring N atoms around the metal center can lead to more defects and accumulation of more negative charges in the support.^{21,22} This, in turn, induces differentiation in the local electron density of active sites, enhancing the adsorption mode and capacity of intermediates, ultimately influencing the CO₂RR performance. For instance, a series of Co SACs featuring Co–N_x (where $x = 4, 3, 2$) active sites synthesized at temperatures of 800, 900, and 1000 °C, respectively, were employed to explore the correlation between the Co–N_x sites and the catalytic performance in the CO₂-to-CO conversion process.²³ The findings indicated that a modulation of the Co–N coordination number results in an augmentation of the unoccupied 3d orbitals of the Co center, and such an alteration is conducive to the adsorption of the CO₂[–] intermediate, thus enhancing the CO₂RR performance. Likewise, Ni SACs known as Ni_{SA}–N_x–C derived from metal–organic frameworks with alterable N coordination numbers were developed through precise regulation of the pyrolysis temperature.²⁴ Interestingly, it had been observed that Ni_{SA}–N₂–C catalysts with the lowest N coordination number exhibited the highest selectivity and activity for CO generation. Consequently, there is growing interest in modifying the coordination environment to enable the directed synthesis of SACs with varying coordination numbers, in order to enhance the conversion of CO₂ into CO.^{25,26} In addition, the approach of incorporating a homo/hetero-metal atom to form a diatomic site (DS) has been demonstrated to adjust the electronic configuration of individual metal catalytic sites dispersed within the carbon matrix.^{27–29} The DS catalyst not only retains the same characteristics as SACs, but also exhibits synergistic interactions, altering the chemical surroundings to match the complex reaction mechanism in the CO₂ conversion process, thereby enhancing catalytic efficiency.^{30,31} An atom-pair catalyst composed of stable Cu₁⁰–Cu₁^{x+} pair structures was created, and

the Cu₁⁰ site bound one CO₂ molecule, while the Cu₁^{x+} site absorbed one H₂O molecule to boost the activation of CO₂.³² Despite some research progress in the construction of DSs, to the best of our knowledge, there has been limited attention given to the precise regulation of their low-coordination microenvironment at the atomic level. Furthermore, manipulating the catalytic microenvironment through the low-coordination effect has the potential to enhance the intrinsic activity of DSs, thereby improving the kinetics of CO production, but it remains challenging.

In this study, an approach of post-synthetic co-substitution (PSCR) has been proposed to achieve precise control of Co–Mn DS catalysts in a low-coordination microenvironment on a N-doped carbon host (L-Co₁Mn₁-NC) derived from nanorod-shaped zeolitic imidazolate framework-8 (ZIF-8). In the first stage of pyrolysis, the Zn nodes and ligand within ZIF-8 undergo gradual degradation, resulting in the formation of vacancy defects. Subsequently, the pre-evaporated Zn species are rapidly removed at elevated temperatures, leading to the creation of numerous Zn vacancies surrounded by varying N atoms for the co-infiltration of the Co/Mn DS. At a temperature of 1000 °C, the Co/Mn DS exhibits simultaneous occupation of the Zn vacancy with a 3 N atom coordination, resulting in the creation of a low-coordinated Co/Mn DS. Conversely, at 900 °C, the Co–Mn DS is filled with Zn vacancies with a 4 N atom coordination. The L-Co₁Mn₁-NC DS catalyst exhibits the highest CO faradaic efficiency (FE_{CO}) reaching 97.6%, along with a turnover frequency (TOF) of 1067 h^{–1} at –0.47 V during the CO₂RR, and its CO selectivity and activity exceed the performance of Co₁Mn₁-NC, Co₁-NC, and Mn₁-NC catalysts. *In situ* ATR-SEIRAS reveals that the low coordination microenvironment endows the Co/Mn DS with the ability to produce CO at a low potential. Moreover, theoretical calculations demonstrate that the presence of low-coordinated Co–Mn active sites in the L-Co₁Mn₁-NC catalyst significantly decreases the adsorption energy of the CO* intermediate, leading to an expedited CO₂ reduction process.

Results and discussion

The low-coordinated structure of the L-Co₁Mn₁-NC DS catalyst was rationally modulated by a PSCR strategy, as illustrated in Fig. 1a. Typically, the ZIF-8 substance was chosen as the supporting structure for the metallic active site.³³ The as-prepared ZIF-8 has a uniform nanorod-like morphology, which is different from the conventional dodecahedron morphology, as demonstrated by scanning electron microscopy (SEM) and transmission electron microscopy (TEM) images (Fig. S1†). By encapsulating with a SiO₂ layer, the initial calcination step not only promotes the distortion and disruption of the Zn–N_x structure to form pre-evaporated Zn species, but also effectively preserves the nanorod-like structure of ZIF-8 for subsequent stages. During secondary pyrolysis conducted at 1000 °C, the pre-evaporated Zn species are rapidly eliminated, while simultaneously, a partial loss of N element occurs within the imidazole ligands, resulting in the formation of Zn vacancies and N defects for the co-substitution and insertion of Co–Mn metal sites. When the pyrolysis is performed at 900 °C, both saturated

coordination Co–Mn bimetallic ($\text{Co}_1\text{Mn}_1\text{-NC}$) and individual Co and Mn monometallic ($\text{Co}_1\text{-NC}$ and $\text{Mn}_1\text{-NC}$) catalysts can be synthesized. The L- $\text{Co}_1\text{Mn}_1\text{-NC}$ DS catalyst featuring a low-coordinated N structure can be synthesized. Both TEM and SEM as well as high-angle annular dark-field scanning TEM (HAADF-STEM) images show that the L- $\text{Co}_1\text{Mn}_1\text{-NC}$ DS catalyst inherits the nanorod-like morphology of ZIF-8, but the surface is rougher and the spatial structure distribution is more open (Fig. 1b and c and S2†). This facilitates the desorption

separation of CO intermediates and gas products. Significantly, the absence of metallic Co, Mn, or CoMn is evident in these findings of L- $\text{Co}_1\text{Mn}_1\text{-NC}$, indicating the potential atomic dispersion of Co and Mn components. Fig. 1d displays the energy-dispersive X-ray spectroscopy (EDS) mapping of the L- $\text{Co}_1\text{Mn}_1\text{-NC}$ DS catalyst, illustrating a uniform dispersion of Co, Mn, N, and C atoms within the nanorod-shaped carbon host. To further confirm the distribution of metal Co/Mn species, aberration-corrected HAADF-STEM (AC HAADF-STEM) was

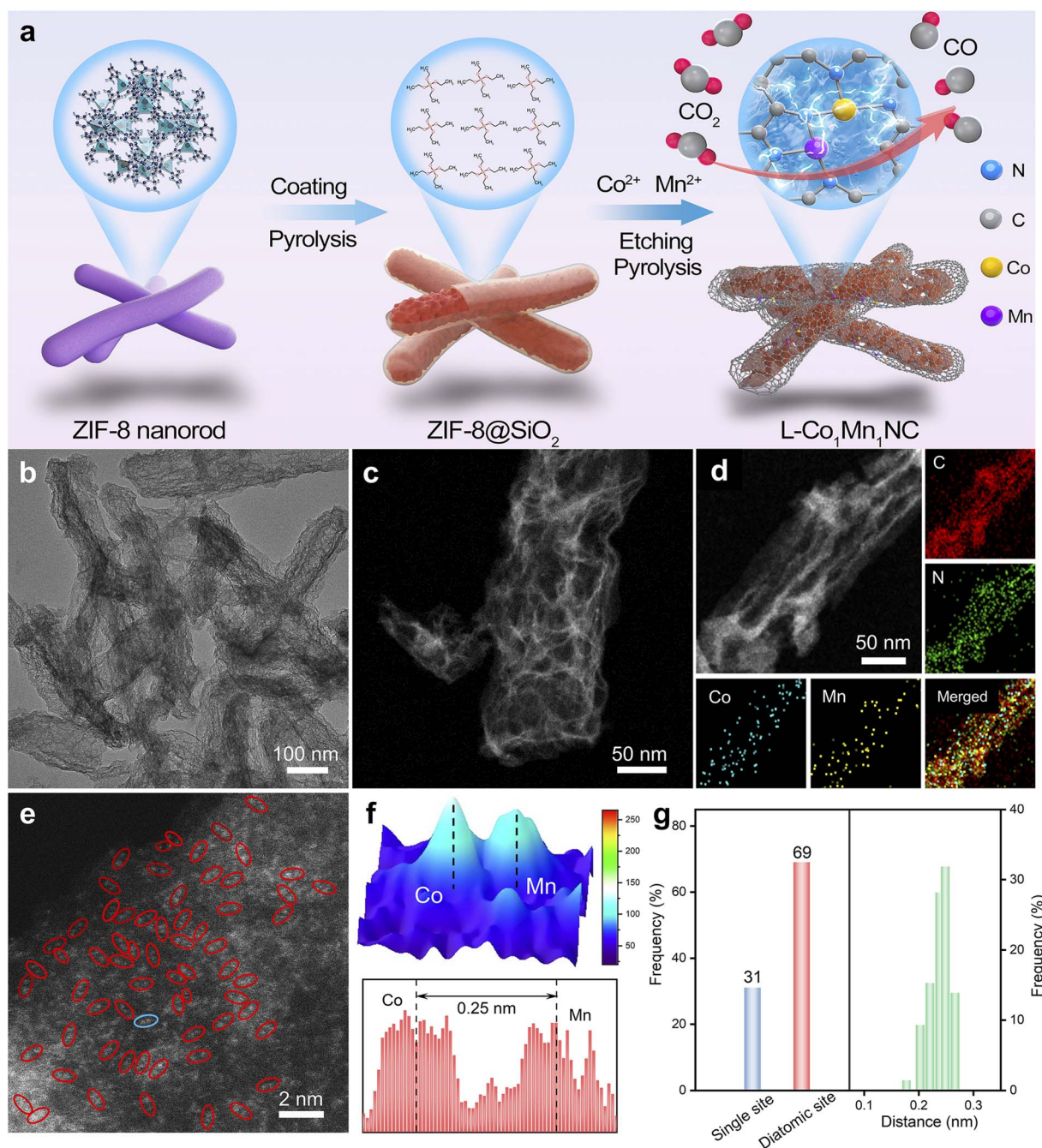


Fig. 1 The morphology and structure characterizations of the L- $\text{Co}_1\text{Mn}_1\text{-NC}$ catalyst. (a) Illustration depicting the PSCR approach utilized in the production of L- $\text{Co}_1\text{Mn}_1\text{-NC}$ catalysts for the CO_2RR . (b) TEM image. (c) HAADF-STEM image. (d) EDS mapping (C in red, N in green, Co in blue, Mn in yellow). (e) AC HAADF-STEM image. (f) 3D map of Co/Mn DSs and intensity spacing diagram. (g) The statistical results of atomic site density and distance distribution.

adopted. As exhibited in Fig. 1e, the area observed in the AC HAADF-STEM image chosen at random indicate the presence of either Co or Mn species, confirming their dispersion at the atomic level. And the selected area electron diffraction (SAED) pattern analysis of the L-Co₁Mn₁-NC DS catalyst, as presented in Fig. S3,[†] reveals the presence of amorphous carbon structure rings exclusively, with no observable diffraction spots corresponding to metal nanoparticles. Following this, a 3D topographic atom imaging analysis was utilized to examine the current distribution of the Co/Mn DS pair based on the blue ellipse in Fig. 1f. The observed hump pair, observed by a minor strength variation, is likely assigned to the Co/Mn DS pair, as indicated by the discrepancy in atomic number. The spacing between the Co/Mn DS pair is measured at 0.25 nm, slightly exceeding the theoretical spacing for a direct Co–Mn connection. In order to minimize the uncertainty attributed to human activities, more than 100 distinct locations (specifically 150 groups of metal sites) were chosen for statistical analyses based on different AC HAADF-STEM images. These statistical analyses include measurements of atomic site density and distance distribution (Fig. 1g and S4[†]). The findings indicate that the L-Co₁Mn₁-NC catalyst primarily exists in the form of Co/Mn DS pairs, with a frequency above 67%, consistent with previous literature on dual-atom structures,^{34,35} and the atomic spacing between Co/Mn pairs is concentrated in the range of 0.24 to 0.26 nm. The metal quantification analysis of the L-Co₁Mn₁-NC is specifically presented in Table S1[†] using the inductively coupled plasma optical emission spectroscopy (ICP-OES) method. In short, the L-Co₁Mn₁-NC catalyst featuring rich Co/Mn pairs undergoes oriented synthesis by the PSCR method.

The X-ray diffraction (XRD) patterns of L-Co₁Mn₁-NC, Co₁Mn₁-NC, Co₁-NC, and Mn₁-NC catalysts exhibit two broad peaks attributed to the (002) and (101) planes of the carbon matrix, with no discernible crystalline phases containing any metal observed (Fig. S5[†]). This observation aligns with the findings from the aforementioned electron microscopy.

The Raman spectra analysis of the four atomic catalysts reveals that the L-Co₁Mn₁-NC DS catalyst exhibits a greater intensity ratio between the D band and G band, suggesting a higher propensity for vacancy defect formation (Fig. S6[†]). To further explain the influence of the low coordination environment, an examination of the chemical composition and electronic state of L-Co₁Mn₁-NC and Co₁Mn₁-NC was conducted through X-ray photoelectron spectroscopy (XPS). In the N 1s XPS spectra (Fig. S7[†]), there is a noticeable shift in the distribution of metallic N (399.5 eV) and graphitic N (401.2 eV) in L-Co₁Mn₁-NC compared to Co₁Mn₁-NC. Specifically, there is a decrease in metallic N and an increase in graphitic N in L-Co₁Mn₁-NC. This suggests a lower N coordination number of Co/Mn DSs in the former, along with a higher level of graphitization overall, likely induced by elevated temperatures. Furthermore, the high-resolution Co 2p and Mn 2p XPS analyses reveal that Co in L-Co₁Mn₁-NC shifts to higher binding energy relative to Co₁Mn₁-NC (Fig. 2a), while Mn migrates toward lower binding energy (Fig. 2b), which may be due to the fact that Mn has fewer d electrons and easily receives external electrons.

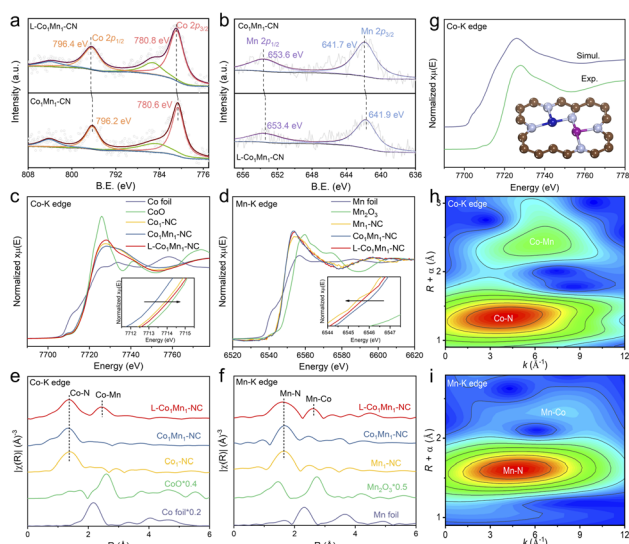


Fig. 2 Geometric and electronic structure characterization of atomic catalysts. XPS spectra of (a) Co 2p and (b) Mn 2p. XANES spectra of (c) the Co K-edge and (d) Mn K-edge. Fourier-transform EXAFS fitting curves of (e) the Co K-edge and (f) Mn K-edge. (g) Comparison between the experimental XANES spectrum of L-Co₁Mn₁-NC at the Co K-edge and the theoretical spectrum based on the structure in the inset. WTs for the EXAFS signals of (h) the Co K-edge and (i) Mn K-edge for L-Co₁Mn₁-NC.

To delve deeply into the variances in the chemical environments of L-Co₁Mn₁-NC and Co₁Mn₁-NC, as well as individual SACs, X-ray absorption near-edge structure (XANES) and extended X-ray absorption fine structure (EXAFS) measurements were carried out. XANES spectra at the Co K-edge for Co₁-NC, Co₁Mn₁-NC and L-Co₁Mn₁-NC exhibit a comparable absorption edge to that of the CoO reference, suggesting a valence state close to +2 (Fig. 2c). In contrast to Co₁Mn₁-NC, the valence state of the low-coordination L-Co₁Mn₁-NC exhibits a slight increase, while the valence state of the Mn atom in L-Co₁Mn₁-NC experiences a decrease (Fig. 2d), thus indicating the electron transfer from Co to Mn. The coordination environments surrounding Co and Mn were additionally investigated using EXAFS analysis. The coordination peaks at 1.4 Å and 2.5 Å in L-Co₁Mn₁-NC can be assigned to the scattering path of Co–N and Co–Mn, respectively (Fig. 2e). Likewise, the presence of Mn–N and Mn–Co coordination can be detected in the vicinity of the Mn atom (Fig. 2f). Furthermore, the theoretical XANES spectrum is calculated based on the structural model in Fig. 2g, and the oscillation characteristics of the experimental spectrum are reproduced well.

The wavelet transform (WT) analysis, along with the quantitative fitting outcomes of EXAFS, provides additional confirmation regarding the presence of Co/Mn–N coordination in the compound. In the WT plots (Fig. 2h and i), the Co–Mn coordination observed at the Co K-edge is marginally shifted to the left relative to the Mn–Co coordination observed at the Mn K-edge. This discrepancy can be rationalized by considering that the scattering ability of Mn, functioning as a coordinating atom, is slightly lower than that of Co. Subsequently, the WT analyses were conducted on the EXAFS spectra obtained at the Co K-edge

for samples including Co foil, Co₁-NC, and L-Co₁Mn₁-NC, as well as at the Mn K-edge for samples comprising Mn foil, Mn₁-NC, and L-Co₁Mn₁-NC (Fig. S8†). These findings illustrate that in L-Co₁Mn₁-NC, the Co–Mn coordination at the Co K-edge lies between the Co–Co coordination in Co foil and Co–N coordination in Co₁-NC. This tendency is also observed at the Mn K-edge. Consequently, it can be inferred that the coordination arrangement of the nearest neighboring metal atom in L-Co₁Mn₁-NC predominantly features Co–Mn pairs rather than self-pairing of identical metals. Moreover, the extracted EXAFS fitting parameters indicate a relatively low N coordination in the L-Co₁Mn₁-NC sample (Fig. S9 and S10, and Tables S2 and S3†). The findings presented suggest that L-Co₁Mn₁-NC possesses a low-coordination structure of Co/Mn dual sites, with evidence of charge transfer occurring between Co and Mn, resulting in a more pronounced interaction.

Given the variances in the coordination microenvironments, electrocatalytic CO₂ reduction over Co–Mn DSs and their corresponding single sites has been investigated. The CO₂RR assessments were conducted on L-Co₁Mn₁-NC, Co₁Mn₁-NC, Co₁-NC, and Mn₁-NC catalysts in an H-type cell utilizing a CO₂-saturated 0.5 M KHCO₃ solution (Fig. S11†). The L-Co₁Mn₁-NC sample was initially examined using linear sweep voltammetry (LSV) profiles in KHCO₃ electrolyte saturated with Ar and CO₂, respectively. As revealed in Fig. S12,† the L-Co₁Mn₁-NC sample exhibits elevated total current densities and a decreased onset potential in the electrolyte saturated with CO₂ compared to that of the Ar saturated environment, indicating that the structure of low-coordinated DSs may facilitate the CO₂RR activity. All electrochemical data were corrected by *i*R compensation

through the test impedance spectra (Fig. S13†), from which it can be found that the L-Co₁Mn₁-NC shows a lower value of solution impedance than those of Co₁Mn₁-NC, Co₁-NC and Mn₁-NC catalysts. Significantly, the L-Co₁Mn₁-NC catalyst also exhibits a notably higher current response and smaller onset potential for CO₂ conversion compared to the Co₁Mn₁-NC, Co₁-NC, and Mn₁-NC catalysts (Fig. 3a). This underscores the enhanced activity associated with the low coordination effect of Co–Mn dual sites. Furthermore, based on the results of online gas chromatography, the products on the four atomic catalysts consist of CO and H₂, with no liquid-phase products generated, as confirmed by ¹H nuclear magnetic resonance spectroscopy (Fig. S14†). The enhancement of CO faradaic efficiency (FE_{CO}) on the four atomic catalysts can be divided into two stages: firstly, a significant increase can be achieved by utilizing DSs; secondly, further improvement can be attained by reducing their coordination number. Specifically, the L-Co₁Mn₁-NC catalyst shows a maximum FE_{CO} of 97.6% obtained at –0.47 V, which can be sustained at levels exceeding 90% across a broad 400 mV potential window (Fig. 3b), which exceeds the performance of Co₁Mn₁-NC (84.7% at –0.57 V), Co₁-NC (64.6% at –0.57 V), and Mn₁-NC (52.8% at –0.67 V) catalysts. Following the creation of a low-coordinated DS microenvironment, there is a notable enhancement in CO selectivity across all operational potential windows, particularly at low overpotential conditions. The carbon substrates produced at varying pyrolysis temperatures may also play a role in influencing the activity of the CO₂RR (Fig. S15†). The polarization curves demonstrate a notable disparity in current density between the carbon carrier and the loading Co–Mn diatomic site, with the CO product

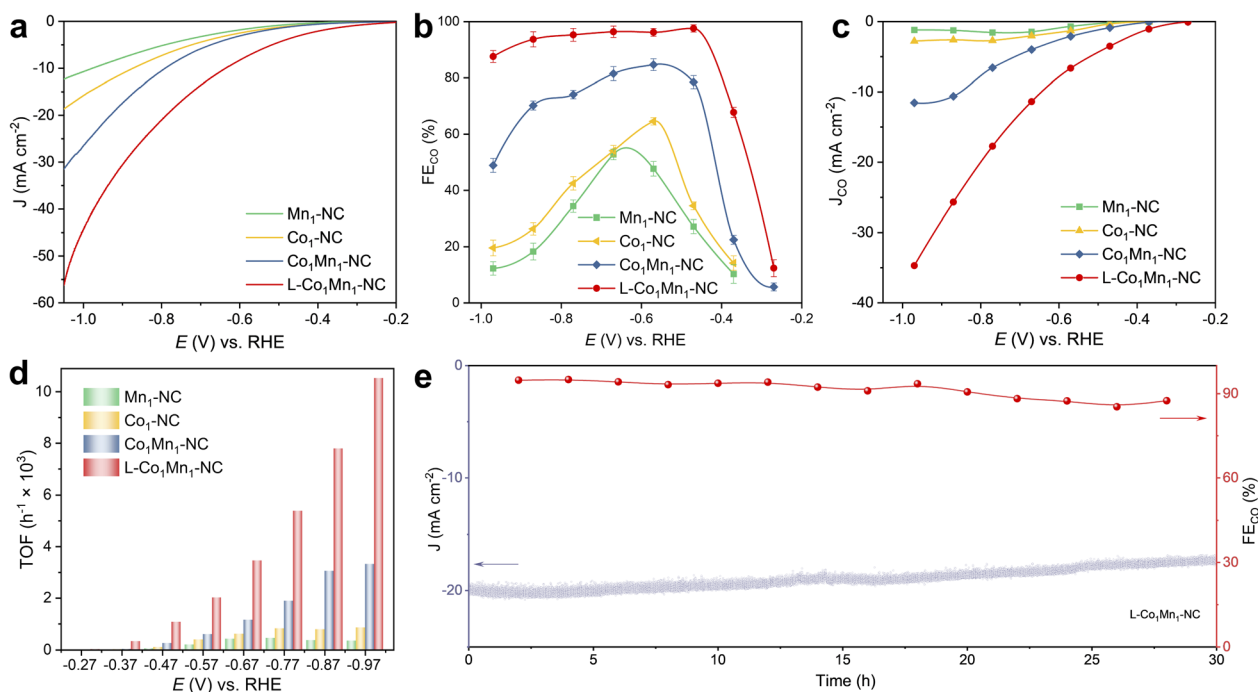


Fig. 3 The CO₂RR performance of L-Co₁Mn₁-NC, Co₁Mn₁-NC, Co₁-NC, and Mn₁-NC catalysts. (a) LSV profiles. (b) FE_{CO} curves. The error bars correspond to the standard deviation of three independent measurements. (c) *J*_{CO} curves. (d) TOF comparison image. (e) The stability test curve of the L-Co₁Mn₁-NC catalyst at –0.8 V.

exhibiting significantly lower efficiency. These findings suggest that the enhanced catalytic performance of the CO₂RR on Co₁Mn₁-NC and L-Co₁Mn₁-NC catalysts primarily stems from the specific microenvironment surrounding the Co–Mn diatomic sites. Besides, a ¹³C isotope labeling experiment was conducted to investigate the precise source of the CO product. By utilizing ¹³CO₂ as the feedstock for CO₂ conversion, it is confirmed that the formation of CO originates solely from the consumption of the CO₂ molecule, rather than from alternative carbon sources within the catalyst (Fig. S16†).³⁶ The CO partial current density (*J*_{CO}) is significantly enhanced after the construction of low-coordinated DSs, as shown in Fig. 3c, achieving a value of 34.7 mA cm⁻² at -0.97 V on the L-Co₁Mn₁-NC catalyst, which demonstrates that the low-coordinated microenvironment promotes the intrinsic activity of the reaction Co/Mn site.

Afterwards, we proceed to quantitatively assess and compare the intrinsic activity of each metal site through the turnover frequency (TOF). As shown in Fig. 3d, the TOF of L-Co₁Mn₁-NC reaches 1067 h⁻¹ at -0.47 V, significantly surpassing the TOFs of Co₁Mn₁-NC (255 h⁻¹), Co₁-NC (104 h⁻¹), and Mn₁-NC (51 h⁻¹) catalysts when subjected to identical operating conditions. In comparison with the reported atomic catalysts,^{37–39} the performance of L-Co₁Mn₁-NC also ranks at the forefront among similar systems reported in the literature (Table S4†). Among these, L-Co₁Mn₁-NC nearly fills the radar chart under equivalent conditions. The construction of Co/Mn DSs and modulation of the coordination environment significantly enhance the activity and selectivity of the CO₂RR. The low-coordinated M–N moieties play a crucial role in promoting this process. In addition, the L-Co₁Mn₁-NC catalyst exhibits acceptable stability, maintaining the current density at approximately 86% at -0.8 V for a duration of 30 hours, while the FE_{CO} level remains consistently above 85% throughout the stability assessment, as illustrated in Fig. 3e. Following the stability assessment, several examinations were conducted to analyze the morphology and structural integrity of the L-Co₁Mn₁-NC DS catalyst. The SEM and TEM images still show that the L-Co₁Mn₁-NC sample maintains a rough and uneven nanorod-like structure after CO₂RR operations (Fig. S17†). The EDS mappings also demonstrate the even dispersion of Co, Mn, N, and C on the carbon host (Fig. S18†). The XRD (Fig. S19†) analysis of L-Co₁Mn₁-NC reveals the presence of only two peaks corresponding to the C (002) and C (101) phases, indicating that the Co/Mn metallic species remain stable during the CO₂RR process without aggregating to form bulk crystalline structures. Furthermore, the micro-distribution state of the Co/Mn active sites was confirmed by AC HAADF-STEM results, still remaining atomically dispersed (Fig. S20†). In short, these findings unequivocally indicate that the Co/Mn DS, characterized by a lower N coordination number, displays superior catalytic activity, selectivity, and stability for the CO₂RR, underscoring the crucial significance of regulating the microenvironment coordination of atoms in catalysis.

To assess the practical application of the L-Co₁Mn₁-NC catalyst under large current density, a custom-designed CO₂ flow cell was utilized, incorporating gas diffusion electrodes (Fig. S21†).

Fig. S22a† displays the polarization curve obtained from flow cell measurements, showing significantly higher current densities due to improved mass transfer compared to those obtained using an H-cell at a specific applied potential. At -0.84 V, the flow cell achieved a remarkable current density of -330 mA cm⁻². Additionally, it exhibited excellent CO selectivity, with FE_{CO} exceeding 90% across a wide current density range from -25 to -275 mA cm⁻² (Fig. R22b†). The endurance of the L-Co₁Mn₁-NC catalyst was evaluated by chronoamperometry at -0.55 V *versus* RHE in the flow cell setup (Fig. R22c†). The L-Co₁Mn₁-NC catalyst demonstrates good durability, with a gradual decrease in current over time from an initial density of -120 mA cm⁻² to -107.7 mA cm⁻² after 80 hours of electrolysis, retaining 89% of its initial value post stability assessment. FE_{CO} levels were monitored every 10 hours, consistently remaining high at 87% throughout the 80 hour stability evaluation. These findings highlight the potential application of the L-Co₁Mn₁-NC catalyst at high current density conditions.

In order to track the catalytic behavior of the reaction interface, we utilized *in situ* attenuated total reflectance surface-enhanced infrared absorption spectroscopy (ATR-SEIRAS) to observe the key intermediates.⁴⁰ The ATR-SEIRAS spectra were acquired by incrementally reducing the potential from 0 to -1.0 V, as illustrated in Fig. S23.† In an electrolyte saturated with CO₂, the IR spectrum recorded at 0 V is regarded as the baseline reference (R0), with no discernible reactive group signal initially observed. Two significant peaks corresponding to the adsorbed CO₂ (2343 cm⁻¹) and COOH* (1425 cm⁻¹) signals are observed starting from a potential of -0.4 V. With a progressive rise in potentials, the CO₂ present at the active interface is continuously depleted within the potential range of -0.4 to -1.0 V. The formation of the COOH* intermediate primarily occurs through the rapid protonation of the adsorbed *CO₂⁻ species, followed by further protonation to generate CO*. Importantly, the emergence of CO* is also identified on the low-coordinated Co/Mn DS at -0.3 V, which is consistent with the experimental results of CO production at a low potential. As the potential increases, the accumulation of adsorbed CO* gradually occurs. Combined with the application potential of CO*, it can be inferred that the CO* desorption is the step that limits the potential of the CO₂RR. Furthermore, across all potential ranges, the strength of H₂O adsorption at the low-coordinated Co/Mn DS remains relatively constant, serving as a potential stable proton donor.⁴¹ Additionally, due to the consumption of H⁺ during the reaction process, a small amount of CO₃²⁻ will form in the localized buffer equilibrium saturated with CO₂.⁴² Furthermore, we also recorded the *in situ* ATR-SEIRAS spectra for the Co₁Mn₁-NC catalyst (Fig. S24†). The adsorption peaks of CO* and COOH* species begin to appear at a higher applied potential compared to the L-Co₁Mn₁-NC catalyst, indicating the effect of low coordination structure has significantly improved the catalytic performance. Therefore, we have provided experimental evidence through *in situ* ATR-SEIRAS to unveil the reduction process of CO₂ on the low-coordination Co/Mn DS.

In view of the excellent catalytic performance of the low-coordinated Co/Mn DS microenvironment toward the CO₂RR, density functional theory (DFT) calculations have been further

performed to explore its essential influence mechanism. Based on structural characterization analysis, we establish the structural models of L-Co₁Mn₁-NC and Co₁Mn₁-NC catalysts and optimize the adsorption models of intermediates at different sites (Fig. S25–S28[†]). The differential charge density diagrams (Fig. 4a–d) illustrate the charge redistribution within Co₁Mn₁-NC and L-Co₁Mn₁-NC. Such redistribution has the potential to alter the electronic configuration of the metal active sites *via* charge interactions, thereby modulating their reactivity.^{43,44} Subsequently, as depicted in Fig. 4e and f, the free energy diagrams for the CO₂RR and HER at $U = 0$ V elucidate the impact of the low coordination effect of the Co/Mn DS structure on performance. Except for Co site in Co₁Mn₁-NC, the potential-limiting step of other sites within Co₁Mn₁-NC and L-Co₁Mn₁-NC is CO desorption (*CO → *+CO), and the Co site at L-Co₁Mn₁-NC exhibits a relatively small energy barrier (0.92 eV). Previous studies have indicated that the slow desorption of CO is a significant factor contributing to the poor selectivity and activity of CO₂-to-CO conversion.^{45,46} The calculation results demonstrate that the adsorption energy of CO can be decreased and the desorption of CO can be enhanced by utilizing a low coordination bimetallic center (CoN₃–MnN₃) in comparison to Co₁Mn₁-NC. The competitive response of the CO₂RR is primarily represented by the HER, thereby necessitating the calculation of the limiting potential difference between the CO₂RR and HER ($U_L(\text{CO}_2) - U_L(\text{H}_2)$) for both catalysts to assess reaction selectivity. The more positive difference value signifies enhanced selectivity, and thus L-Co₁Mn₁-NC shows higher CO selectivity (Fig. 4g).⁴⁷ Therefore, the low-coordinated microenvironment of L-Co₁Mn₁-NC facilitates the attainment of enhanced performance in the CO₂RR, aligning with the outcomes observed in experimental results.

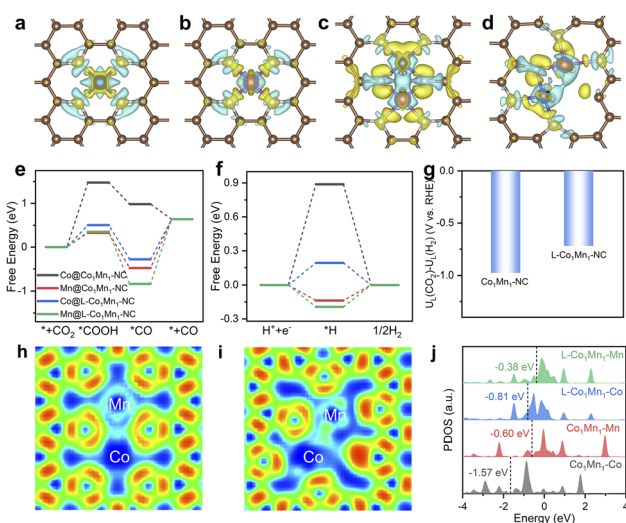


Fig. 4 Charge density difference of (a) Co₁-NC, (b) Mn₁-NC, (c) Co₁Mn₁-NC and (d) L-Co₁Mn₁-NC, where electron depletion and electron accumulation are shown in yellow and cyan, respectively. (e) Gibbs free energy diagrams for the CO₂-to-CO conversion process. (f) Gibbs free energy diagrams for the HER. (g) Calculated limiting potential difference of the CO₂RR and HER. ELF analysis for (h) Co₁Mn₁-NC and (i) L-Co₁Mn₁-NC. (j) PDOS of the 3d-orbital.

Besides, we examined the L-Co₂-NC and L-Mn₂-NC models through DFT calculations, possessing a structure akin to that of the L-Co₁Mn₁-NC model (Fig. S29 and S30[†]). Subsequently, we evaluated their performance in the CO₂RR and HER. As exhibited in Fig. S31,[†] the results show that the overpotentials of the CO₂RR of L-Co₂-NC and L-Mn₂-NC are 1.08 V and 1.54 V, respectively, both of which are significantly higher than that of L-Co₁Mn₁-NC. In addition, their HER overpotential is lower, resulting in a lower limiting potential difference. Consequently, the CO selectivity on L-Co₂-NC and L-Mn₂-NC models is compromised, marking a notable deviation from the observed performance in sample testing.

Moreover, an analysis of the electronic structure was conducted to elucidate the inherent origin of low-coordination L-Co₁Mn₁-NC. The electron localization function (ELF) analysis indicates a higher concentration of electrons shared between the Co and Mn sites within the L-Co₁Mn₁-NC (Fig. 4h and i). This phenomenon suggests an enhanced charge interaction and a direct electron transfer between the Co and Mn sites. Quantitatively, the Bader charge analysis (Fig. S32[†]) reveals a rise in the valence state of Co and a decline in Mn when compared to Co₁Mn₁-NC, which substantiates the charge transfer occurring between Co and Mn sites, aligning with the findings from XPS and XANES analyses. Subsequently, the electronic configuration of Co and Mn 3d orbitals was probed by the projected density of states (PDOS). It is observed that there is a rise in the d-band center of Co and Mn in the L-Co₁Mn₁-NC, with a more pronounced alteration noted in the case of Co (Fig. 4j). This leads to a transfer of electrons from the Co 3d orbital to lower energy states, specifically the Mn 3d orbital. In addition, a strong d–d orbital coupling and delocalization of electrons promote the desorption of CO, thus reducing the reaction energy barrier of the CO₂RR.⁴⁸ Additionally, the total DOS near the Fermi energy within L-Co₁Mn₁-NC surpasses that of Co₁Mn₁-NC, as illustrated in Fig. S33.[†] This indicates that the enhanced conductivity and the facilitation of charge transfer at the reaction interface are attributed to the lower coordination structure that emerged in the L-Co₁Mn₁-NC catalyst.

Conclusions

In brief, a direct and rational PSCR approach was employed to fabricate Co–Mn DS catalysts dominated by a low-coordination microenvironment *via* simultaneously substituting Co and Mn atoms for the atomically dispersed Zn sites in the Zn-N_x-C material derived from ZIF-8 nanorods. When altering the pyrolysis temperature and substituting metal in the PSCR strategy, a relatively high N coordination of Co–Mn DSs and the common 4-coordination of Co/Mn–N₄ structures can be obtained, respectively. The development of this PSCR strategy has simplified the cumbersome steps involved in synthesizing DS catalysts derived from MOFs, and has achieved the regulation of DS coordination environments through simple temperature control. Accordingly, the optimal L-Co₁Mn₁-NC catalyst exhibits superior efficiency in CO₂ electroreduction, achieving a high FE_{CO} of 97.6% and a TOF value of 1067 h^{−1} at −0.47 V, which significantly surpass those of Co₁Mn₁-NC and individual Co/Mn

single sites. A combination of *in situ* investigation and theoretical analysis unveils that the lower Co/Mn coordination number in the L-Co₁Mn₁-NC catalyst efficiently boosts the desorption of the critical CO* species, thereby enhancing the CO₂ reduction kinetics. This study presents a convenient and rational control approach for manipulating the catalytic microenvironment of atomic catalysts, laying a solid experimental foundation for improving the reaction kinetics of energy conversion processes.

Author contributions

J. D. conceived the idea and designed the research. J. P., S. J., P. W. and H. H. performed the synthesis, structural characterisations, and electrochemical measurements. P. A. and J. L. performed the XAFS measurements and analysed the data. G. Z. and S. C. performed DFT simulations. J. P. and G. Z. analysed and discussed the experimental results and wrote the manuscript. All the authors discussed the results and commented on the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

J. D. acknowledges financial support from the Youth Innovation Promotion Association of the Chinese Academy of Sciences (Y2022006) and the Innovation project of Institute of High Energy Physics, Chinese Academy of Sciences. J. P. acknowledges financial support from the China Postdoctoral Science Foundation (grant no. 2023M742047) and the Beijing Natural Science Foundation (no. 1244064).

Notes and references

- C. Chen, J. F. Khosrowabadi Kotyk and S. W. Sheehan, *Chem*, 2018, **4**, 2571–2586.
- B. S. Crandall, S. Overa, H. Shin and F. Jiao, *Acc. Chem. Res.*, 2023, **56**, 1505–1516.
- A. D. Handoko, F. Wei, Jenndy, B. S. Yeo and Z. W. Seh, *Nat. Catal.*, 2018, **1**, 922–934.
- L. Fan, C. Xia, F. Yang, J. Wang, H. Wang and Y. Lu, *Sci. Adv.*, 2020, **6**, eaay3111.
- S. Nitopi, E. Bertheussen, S. B. Scott, X. Liu, A. K. Engstfeld, S. Horch, B. Seger, I. E. L. Stephens, K. Chan, C. Hahn, J. K. Nørskov, T. F. Jaramillo and I. Chorkendorff, *Chem. Rev.*, 2019, **119**, 7610–7672.
- M. B. Ross, P. De Luna, Y. Li, C.-T. Dinh, D. Kim, P. Yang and E. H. Sargent, *Nat. Catal.*, 2019, **2**, 648–658.
- T. N. Nguyen, M. Salehi, Q. V. Le, A. Seifitokaldani and C. T. Dinh, *ACS Catal.*, 2020, **10**, 10068–10095.
- D. Yang, B. Ni and X. Wang, *Adv. Energy Mater.*, 2020, **10**, 2001142.
- C. Zhu, J. Yang, J. Zhang, X. Wang, Y. Gao, D. Wang and H. Pan, *Interdiscip. Mater.*, 2024, **3**, 74–86.
- Y. Wang, Y. Zhang, W. Yu, F. Chen, T. Ma and H. Huang, *J. Mater. Chem. A*, 2023, **11**, 2568–2594.
- S. Zhu, E. P. Delmo, T. Li, X. Qin, J. Tian, L. Zhang and M. Shao, *Adv. Mater.*, 2021, **33**, 2005484.
- J. Zhuang and D. Wang, *Mater. Today Catal.*, 2023, **2**, 100009.
- J. Dong, Y. Liu, J. Pei, H. Li, S. Ji, L. Shi, Y. Zhang, C. Li, C. Tang, J. Liao, S. Xu, H. Zhang, Q. Li and S. Zhao, *Nat. Commun.*, 2023, **14**, 6849.
- T. Tang, Z. Wang and J. Guan, *Adv. Funct. Mater.*, 2022, **32**, 2111504.
- M. Li, H. Wang, W. Luo, P. C. Sherrell, J. Chen and J. Yang, *Adv. Mater.*, 2020, **32**, 2001848.
- Y. Wang, H. Su, Y. He, L. Li, S. Zhu, H. Shen, P. Xie, X. Fu, G. Zhou, C. Feng, D. Zhao, F. Xiao, X. Zhu, Y. Zeng, M. Shao, S. Chen, G. Wu, J. Zeng and C. Wang, *Chem. Rev.*, 2020, **120**, 12217–12314.
- R. Qin, K. Liu, Q. Wu and N. Zheng, *Chem. Rev.*, 2020, **120**, 11810–11899.
- X. Li, L. Liu, X. Ren, J. Gao, Y. Huang and B. Liu, *Sci. Adv.*, 2020, **6**, eabb6833.
- C. Jia, X. Tan, Y. Zhao, W. Ren, Y. Li, Z. Su, S. C. Smith and C. Zhao, *Angew. Chem., Int. Ed.*, 2021, **60**, 23342–23348.
- S. Li, X. Lu, S. Zhao, M. Ceccato, X.-M. Hu, A. Roldan, M. Liu and K. Daasbjerg, *ACS Catal.*, 2022, **12**, 7386–7395.
- L. Yuan, S. Zeng, G. Li, Y. Wang, K. Peng, J. Feng, X. Zhang and S. Zhang, *Adv. Funct. Mater.*, 2023, **33**, 2306994.
- D. Xi, J. Li, J. Low, K. Mao, R. Long, J. Li, Z. Dai, T. Shao, Y. Zhong, Y. Li, Z. Li, X. J. Loh, L. Song, E. Ye and Y. Xiong, *Adv. Mater.*, 2021, **34**, 2104090.
- X. Wang, Z. Chen, X. Zhao, T. Yao, W. Chen, R. You, C. Zhao, G. Wu, J. Wang, W. Huang, J. Yang, X. Hong, S. Wei, Y. Wu and Y. Li, *Angew. Chem., Int. Ed.*, 2018, **57**, 1944–1948.
- Y. N. Gong, L. Jiao, Y. Qian, C. Y. Pan, L. Zheng, X. Cai, B. Liu, S. H. Yu and H. L. Jiang, *Angew. Chem., Int. Ed.*, 2020, **59**, 2705–2709.
- M. Wang, Z. Zhang, S. Zhang, W. Liu, W. Shang, X. Su, Y. Liang, F. Wang, X. Ma, Y. Li and Y. Liu, *Angew. Chem., Int. Ed.*, 2023, **62**, e202300826.
- Y. Deng, J. Zhao, S. Wang, R. Chen, J. Ding, H.-J. Tsai, W.-J. Zeng, S.-F. Hung, W. Xu, J. Wang, F. Jaouen, X. Li, Y. Huang and B. Liu, *J. Am. Chem. Soc.*, 2023, **145**, 7242–7251.
- D.-C. Zhong, Y.-N. Gong, C. Zhang and T.-B. Lu, *Chem. Soc. Rev.*, 2023, **52**, 3170–3214.
- Y.-X. Zhang, S. Zhang, H. Huang, X. Liu, B. Li, Y. Lee, X. Wang, Y. Bai, M. Sun, Y. Wu, S. Gong, X. Liu, Z. Zhuang, T. Tan and Z. Niu, *J. Am. Chem. Soc.*, 2023, **145**, 4819–4827.
- Z.-H. Zhao, J.-R. Huang, P.-Q. Liao and X.-M. Chen, *J. Am. Chem. Soc.*, 2023, **145**, 26783–26790.
- Y. Chen, J. Lin, Q. Pan, X. Liu, T. Ma and X. Wang, *Angew. Chem., Int. Ed.*, 2023, **62**, e202306469.
- Y. N. Gong, C. Y. Cao, W. J. Shi, J. H. Zhang, J. H. Deng, T. B. Lu and D. C. Zhong, *Angew. Chem., Int. Ed.*, 2022, **61**, e202215187.
- J. Jiao, R. Lin, S. Liu, W.-C. Cheong, C. Zhang, Z. Chen, Y. Pan, J. Tang, K. Wu, S.-F. Hung, H. M. Chen, L. Zheng,

- Q. Lu, X. Yang, B. Xu, H. Xiao, J. Li, D. Wang, Q. Peng, C. Chen and Y. Li, *Nat. Chem.*, 2019, **11**, 222–228.
- 33 S. Ma, W. Han, W. Han, F. Dong and Z. Tang, *J. Mater. Chem. A*, 2023, **11**, 3315–3363.
- 34 L. Zhang, R. Si, H. Liu, N. Chen, Q. Wang, K. Adair, Z. Wang, J. Chen, Z. Song, J. Li, M. N. Banis, R. Li, T.-K. Sham, M. Gu, L.-M. Liu, G. A. Botton and X. Sun, *Nat. Commun.*, 2019, **10**, 4936.
- 35 Q. Hao, H.-X. Zhong, J.-Z. Wang, K.-H. Liu, J.-M. Yan, Z.-H. Ren, N. Zhou, X. Zhao, H. Zhang, D.-X. Liu, X. Liu, L.-W. Chen, J. Luo and X.-B. Zhang, *Nat. Synth.*, 2022, **1**, 719–728.
- 36 J. Pei, H. Shang, J. Mao, Z. Chen, R. Sui, X. Zhang, D. Zhou, Y. Wang, F. Zhang, W. Zhu, T. Wang, W. Chen and Z. Zhuang, *Nat. Commun.*, 2024, **15**, 416.
- 37 T. Wang, Q. Zhang, K. Lian, G. Qi, Q. Liu, L. Feng, G. Hu, J. Luo and X. Liu, *J. Colloid Interface Sci.*, 2023, **655**, 176–186.
- 38 S. Liu, M. Jin, J. Sun, Y. Qin, S. Gao, Y. Chen, S. Zhang, J. Luo and X. Liu, *Chem. Eng. J.*, 2022, **437**, 135294.
- 39 S. Gao, S. Chen, Q. Liu, S. Zhang, G. Qi, J. Luo and X. Liu, *ACS Appl. Nano Mater.*, 2022, **5**, 12387–12394.
- 40 W. Cheng, H. Su and Q. Liu, *Acc. Chem. Res.*, 2022, **55**, 1949–1959.
- 41 H. Xiao, W. A. Goddard, T. Cheng and Y. Liu, *Proc. Natl. Acad. Sci. U. S. A.*, 2017, **114**, 6685–6688.
- 42 C. Liu, Y. Wu, K. Sun, J. Fang, A. Huang, Y. Pan, W.-C. Cheong, Z. Zhuang, Z. Zhuang, Q. Yuan, H. L. Xin, C. Zhang, J. Zhang, H. Xiao, C. Chen and Y. Li, *Chem*, 2021, **7**, 1297–1307.
- 43 Z. Chen, X. Zhang, W. Liu, M. Jiao, K. Mou, X. Zhang and L. Liu, *Energy Environ. Sci.*, 2021, **14**, 2349–2356.
- 44 J. Zhu, M. Xiao, D. Ren, R. Gao, X. Liu, Z. Zhang, D. Luo, W. Xing, D. Su, A. Yu and Z. Chen, *J. Am. Chem. Soc.*, 2022, **144**, 9661–9671.
- 45 Y. Li, W. Shan, M. J. Zachman, M. Wang, S. Hwang, H. Tabassum, J. Yang, X. Yang, S. Karakalos, Z. Feng, G. Wang and G. Wu, *Angew. Chem., Int. Ed.*, 2022, **61**, e202205632.
- 46 X. Qin, T. Vegge and H. A. Hansen, *J. Am. Chem. Soc.*, 2023, **145**, 1897–1905.
- 47 W. Ren, X. Tan, W. Yang, C. Jia, S. Xu, K. Wang, S. C. Smith and C. Zhao, *Angew. Chem., Int. Ed.*, 2019, **58**, 6972–6976.
- 48 Z. Zeng, L. Y. Gan, H. Bin Yang, X. Su, J. Gao, W. Liu, H. Matsumoto, J. Gong, J. Zhang, W. Cai, Z. Zhang, Y. Yan, B. Liu and P. Chen, *Nat. Commun.*, 2021, **12**, 4088.