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## Transition metal oxide clusters: advanced electrocatalysts for a sustainable energy future

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The comprehensive utilization of sustainable green energy is essential to face the global energy and environmental crisis. The oxygen reduction reaction (ORR), oxygen evolution reaction (OER), and electrocatalytic urea synthesis (EUS) are the pivotal electrocatalytic processes, necessitating the development of low-cost electrocatalysts with high efficiency. Small-sized transition metal oxide (TMO) clusters have attracted a lot of attention because of their exceptional qualities, such as exhibiting a dense array of low-coordinated metal active sites (e.g. abundant metal cation defects and oxygen vacancy), amorphous structures with high surface energy, high atom utilization efficiency, and cost-effectiveness. Furthermore, the synergistic actions between metal clusters and TM–N<sub>x</sub> single atom active sites remarkably boost up the electrocatalytic performances, corroborated by density functional theory (DFT). More efforts in this comprehensive feature article are expected to achieve insights into the fundamental understanding of electrocatalytic reaction mechanisms in our lab and serve as a guide for creating cutting-edge electrocatalysts of transition metal oxide clusters.

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### Introduction

Due to expanding global energy use and growing environmental concerns, there is a growing demand for clean, sustainable energy technologies including fuel cells, solar cells, and metal–air batteries.<sup>1–4</sup> The efficiency of catalysts is crucial for the application of these technologies, which rely on catalytic processes, including the oxygen reduction reaction (ORR), the oxygen evolution reaction (OER), and electrocatalytic urea synthesis (EUS) from nitrate ions and CO<sub>2</sub>.<sup>5–7</sup> The high price, lack of stability, and restricted availability of traditional noble metal catalysts like Ir, Pd, and Pt prevent their broad use as energy solutions.<sup>5–7</sup> To overcome these obstacles and make renewable energy systems more practical, lots of efforts have gone into creating metal-free or non-Pt catalysts that are just as efficient for the ORR, OER, EUS, and other processes.<sup>8–10</sup> Multifunctional electrocatalysts as substitutions, which have recently been developed, may provide several methods simultaneously, opening up new possibilities for integrated energy conversion systems.<sup>11–13</sup> There is an alternative for improving electrochemical activity with less expense and resource consumption thanks to new catalysts, such as

ultras-small-sized particles (also named clusters, size of 2–5 nm). Incorporating sophisticated catalysts into comprehensive electrochemical systems may greatly improve energy conversion's life-span and efficiency, propelling us toward a greener, more sustainable energy future.<sup>5,14–17</sup>

Transition metal oxide (TMO) clusters (particle size of 3–6 nm) have a dense array of low-coordinated metal active sites (e.g. abundant metal cation defects and oxygen vacancies), amorphous structures with high surface energy due to the distorted TM–O bonds, high atom utilization efficiency, cost-effectiveness, a few to name.<sup>18–21</sup> These unique geometric characteristics are conducive to electrochemical activity by providing metal–metal contacts and reaction intermediate adsorption sites.<sup>10,22,23</sup> TMO clusters may break scaling relations by lowering reaction barriers, regulating the rate-determining step (RDS) in electrocatalysis and improving catalytic performances (including activity and selectivity) in the ORR, OER, EUS, *etc.* Their ability to create catalysts with co-adsorbates, promoters, ligands, and new alloy structures makes them ideal options for electrocatalysis and energy conversion technology.<sup>24–26</sup> Because of the expensive nature and limited geological resources of precious group metal (PGM) electrocatalysts such as RuO<sub>2</sub> and IrO<sub>2</sub>, there has been growing interest in PGM-free catalysts.<sup>27–29</sup> Therefore, catalysts based on transition metals that are common on Earth (such as Ni, Co, Fe, Mn, *etc.*), particularly transition metal hydroxides (TMHs), have attracted interest due to their electrocatalytic activity, affordability, and strong durability as effective and economical alternatives. In recent decades, several<sup>30–32</sup> TM sulfides,<sup>33–35</sup> nitrides,<sup>36</sup> phosphides,<sup>30,32</sup> oxides,<sup>37,38</sup> and hydroxides<sup>39,40</sup> have been explored as promising

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alternative OER electrocatalysts. And earth-abundant TMHs have attracted attention as promising OER electrocatalysts due to their activity, low cost, excellent stability, and environmentally friendly nature.<sup>41</sup> Fe-based NNMEs have exceptional ORR catalytic activity. These comprise a high concentration of iron-based species such as oxides ( $\text{Fe}_x\text{O}_y$ ), nitrides ( $\text{FeN}_x$ ), carbides ( $\text{FeC}_x$ ), and sulfides ( $\text{FeS}_x$ ).<sup>42–46</sup> Species with high electronegativity can attach stable FeNC electrocatalyst active sites. This might potentially modify the adsorption and desorption of O intermediates on the core Fe atoms, affecting the activities of the oxygen reduction reaction (ORR).<sup>47–49</sup>

To advance the research in TM composites, our group recently explored novel approaches to increase catalytic efficiency in various applications, such as the creation of intrinsically stable  $\text{MnCoO}_x$  solid solutions for long-term water oxidation, the impact of multiple active sites on promotion in Fe-based ORR electrocatalysts, the cooperative role of cobalt clusters– $\text{CoN}_x$  composites in enhancing electrochemical-oxygen-reduction, and the synergistic effect of Fe clusters and  $\text{FeN}_x$  for electrocatalytic urea synthesis (EUS) from nitrate and  $\text{CO}_2$ , Scheme 1. Furthermore, the enhanced electrocatalytic properties of ultrafine metallic hydroxide nanoparticles and FeNi hydroxide nanoclusters modified by ionic liquids for oxygen evolution processes are emphasized. Additionally, the use of atomically precise metal nanoclusters for active-site engineering in heterogeneous catalysis is investigated, offering insights into customizing catalytic processes for increased efficiency. What's more, the mechanism of all these electrocatalysis is well revealed by experiments combined with theoretical calculations. These results highlight the significance of materials design, synthesis techniques, and active-site modification in developing electrocatalysts for sustainable energy systems. This joint effort shows encouraging directions for creating robust and effective electrocatalytic systems necessary to meet urgent global energy issues. Lastly, we provide our thoughts about upcoming initiatives to create efficient TM electrocatalysts.



**Scheme 1** The outline of the articles on metal oxide clusters, including acidic/alkaline OER and alkaline ORR and zinc–air batteries.

## Synthetic methods

### Pechini method

The Pechini process is a sol–gel route invented by Maggio Pechini in which, suitable metal oxides or salts are chelated with hydroxycarboxylic acid (*e.g.* citric acid) in aqueous solutions and then reacted with glycol to form large, cross-linked polymeric networks at 150–250 °C, which are eventually charred and achieves small-sized metal oxide solid solutions at temperatures of 300–350 °C.<sup>50,51</sup> For example, using the potassium permanganate redox technique, the author researched printing  $\text{CoMnO}_x$  compounds on carbon paper (CP).  $\text{Mn}_1\text{Co}_5\text{O}_x$  produced in research differs significantly from doped compounds. Potassium permanganate oxidizes low-valent cobalt to high-valent cobalt, during which, a uniform solid solution of  $\text{Mn}_1\text{Co}_5\text{O}_x$  is formed, Mn-rich doped  $\text{Co}_3\text{O}_4$ . The ability of the solid solution's crystal structure to maintain the solvent's crystal structure, resulting in the same material, structure, and characteristics, is fascinating<sup>51</sup>. This single-phase solid solution has excellent OER catalytic activity. The  $\text{Mn}_1\text{Co}_5\text{O}_x$  catalyst performed well in acid electrolytes, achieving overpotentials of 275 mV@10 mA  $\text{cm}^{-2}$  and 569 mV@100 mA  $\text{cm}^{-2}$ . Additionally,  $\text{MnCoO}_x$  showed exceptional stability during the catalysis for 300 hours at 100 mA  $\text{cm}^{-2}$ , indicating its potential for electrocatalytic applications. As indicated in Fig. 1a,  $\text{MnCoO}_x$  show strong OER catalytic efficiency in an acidic medium. In addition, it highlights this material's unique design and excellent OER performance.<sup>52</sup> DFT results manifested that the Mn dopant can significantly elevate the d-band of  $\text{Co}_3\text{O}_4$  (–2.05 eV) to near Fermi level (–1.78 eV) and sharply reduce the overpotential of the OER from 2.79 eV ( $\text{Co}_3\text{O}_4(311)$ ) to 0.82 eV ( $\text{MnCoO}_x(311)$ ). Thus, the Mn dopant and Ov can stabilize OER intermediates and then reduce the energy barrier of the OER and improve catalytic activity.

Furthermore, Zhang *et al.*<sup>53</sup> developed a straightforward process for mass-producing cobalt-based electrocatalysts by carbonizing at 500 °C as shown in (Fig. 1b). STEM and X-ray absorption showed that pyrolytic treatment merged metallic Co clusters with  $\text{CoN}_x$  sites. In oxygen electroreduction operations, the  $\text{Co/CoN}_x\text{-C}$  composite performed better than the commercial Pt/C catalyst, with an onset potential of 0.926 V vs. RHE and a half-wave potential  $E_{1/2}$  of 0.853 V. The enhanced ORR performance of the  $\text{Co/CoN}_x\text{-C}$  electrocatalyst is supported using first-principles calculations because of the special integration and ideal synergistic effects of electro-catalytically active  $\text{CoN}_x$  and Co cluster species.<sup>55–57</sup> Theoretical results show that the cooperative Co clusters and  $\text{CoN}_x$  have higher electron density compared to bare Co/C and  $\text{CoN}_x\text{-C}$  near the Fermi level, thus,  $\text{Co/CoN}_x\text{-C}$  exhibits better electrical conductivity. And the RDS over  $\text{Co/CoN}_x\text{-C}$  and  $\text{CoN}_x\text{-C}$  occurs at the first step of  $\text{O}_2 \rightarrow \text{OOH}^*$ , and that for Co/C is the last step of  $\text{OH}^* \rightarrow \text{H}_2\text{O}$ . Hence, the superior ORR activity of  $\text{Co/CoN}_x\text{-C}$  catalysts is attributed to the unique integration and synergistic effect of metallic Co clusters and  $\text{CoN}_x$  species.

### Single-step pyrolysis approach

The electrocatalysts were made using a straightforward one-pot method. The N and S precursors, Cy or TCA, were dissolved in





**Fig. 1** (a) Visual representation of the OER process in acidic environments, elucidating the critical points of this investigation. Reproduced with permission from ref. 52, copyright 2023 Royal Society of Chemistry. (b) The synthesis of Co/CoN<sub>x</sub>-C. Reproduced with permission from ref. 53, copyright 2022 Royal Society of Chemistry. (c) An illustrated method for producing electrocatalysts using Hm/Cy@C-900 and Hm/TCA@C-900. Redrawn from ref. 54, copyright 2023 Royal Society of Chemistry. (d) Producing amorphous FeNi hydroxide clusters capped with IL. Redrawn from ref. 55, copyright 2020 Royal Society of Chemistry.

an HNO<sub>3</sub> solution, and hemin was dissolved in a KOH solution. After that, the alkaline solution and the acidic solution were mixed for thirty minutes using ultrasonic shock. There was additional carbon ECP. An overnight drying process at 80 °C was used to eliminate all of the water following a three-hour reaction in a rotary evaporator. Pyrolysis was used to create electrocatalysts at various temperatures in an argon environment. This was accomplished by using an identical synthesis procedure to Hm/Cy@C but replacing Cy with TCA.<sup>54</sup> Notably, producing over 1 g of Fe-based electrocatalysts on a large scale is straightforward (Fig. 1c). M@IL-modified amorphous metallic hydroxide nanoparticles are produced in a single process (Fig. 1d). A IL-NH<sub>2</sub> methanol solution was mixed with an aqueous solution of a metal-containing precursor, such as Fe(NO<sub>3</sub>)<sub>3</sub>, Co(NO<sub>3</sub>)<sub>2</sub>, or Ni(NO<sub>3</sub>)<sub>2</sub>, or their combination. Then, while vigorously shaking the mixture of methanol and water, a NaOH aqueous solution was added. Vulcan XC-72R carbon powder was then added to the M@IL colloidal solution.<sup>55</sup>

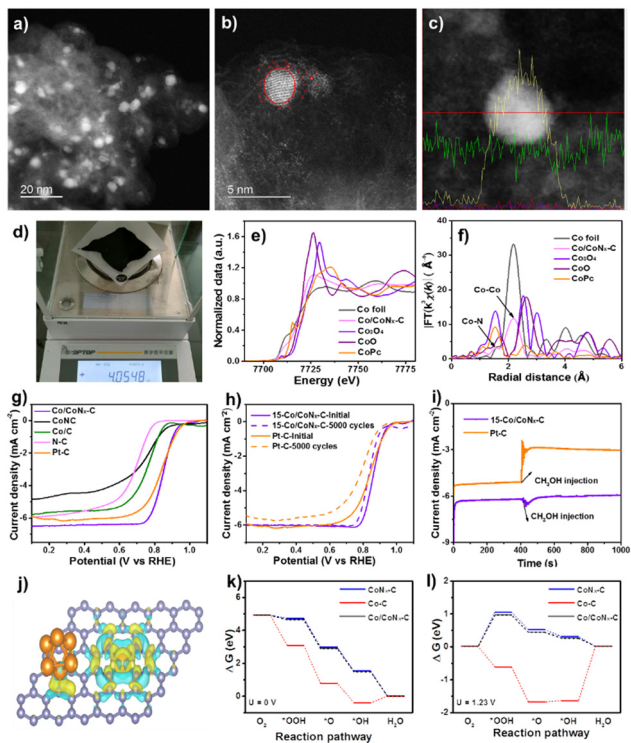
### Mechanistic insights into electrocatalytic activity during the OER/ORR and fuel cell and battery applications

**ORR.** The Co clusters immobilized on N-doped carbon matrixes having a size that is around typical  $4.4 \pm 1.2$  nm is shown in the STEM image of Co/CoN<sub>x</sub>-C (Fig. 2a). As seen in Fig. 2b, red cycles for CoN<sub>x</sub> species and single Co atoms envelop metallic Co clusters. Single-Co-atoms (*e.g.*, CoN<sub>x</sub> species) firmly encircled the metallic Co clusters in N-doped carbon matrixes, according to the elemental mappings of Co/CoN<sub>x</sub>-C on a typical region of a STEM picture (Fig. 2c). The Co K-edge X-ray

absorption near-edge structure (XANES) of Co/CoN<sub>x</sub>-C is between Co foil and CoO<sub>x</sub>, unlike conventional samples like Co foil, CoO, Co<sub>3</sub>O<sub>4</sub>, and CoPc, Fig. 2e. The XANES peak of 7726 eV in Co/CoN<sub>x</sub>-C reduced from 7733 eV in Co foil, perhaps due to strong Co cluster-CoN<sub>x</sub> interaction. Additionally, Fourier-transformed extended X-ray absorption fine structure (FT-EXAFS) analysis revealed a peak at 2.18 Å in the Co/CoN<sub>x</sub>-C electrocatalyst (Fig. 2f), indicating Co<sup>0</sup>-Co<sup>0</sup> bonds and metallic Co clusters. The XAS, XPS, and STEM strongly showed that the Co/CoN<sub>x</sub>-C electrocatalyst's intimate integration of CoN<sub>x</sub> species and Co clusters may synergistically boost ORR catalytic activity.<sup>53</sup>

The Co/CoN<sub>x</sub>-C ORR's catalytically active sites were discovered and identified by analyzing further cobalt-based electrocatalysts of Co/C and CoNC under identical ORR circumstances (Fig. 2g). In terms of  $E_{1/2}$  comparison, ORR activity was as follows: Co/CoN<sub>x</sub>-C > Co/C > CoNC > N-C. Co species increase ORR activity; the N-C deficient Co species exhibited the lowest ORR activity.<sup>58</sup> CoNC had a low restricted current density, suggesting that carbon support improves electrical conductivity. In an O<sub>2</sub>-saturated 0.1 M KOH solution, Co/CoN<sub>x</sub>-C was compared to commercial Pt/C for durability. Electrocatalyst LSV curves (Fig. 2h) were analyzed after 5000 cycles at 100 mV s<sup>-1</sup>.  $E_{1/2}$  losses were ~18 mV for Co/CoN<sub>x</sub>-C and 47 mV for Pt/C, indicating the catalyst's stability and robustness. A possible explanation for the slight reduction in catalytic activity might be the partial formation of Co clusters ( $5.0 \pm 1.5$  nm) in the spent Co/CoN<sub>x</sub>-C catalyst. The *I-t* chronoamperometry in Fig. 2i shows that Co/CoN<sub>x</sub>-C's current density is relatively steady following methanol injection in an O<sub>2</sub>-saturated 0.1 M KOH solution, demonstrating





**Fig. 2** (a)–(c) S/TEM profiles with line scanning of Co/CoN<sub>x</sub>-C. (d) Quality in one-pot synthesis. (e) FT-EXAFS and (f) normalized K-edge XANES spectra. (g) LSV curves. (h) LSV curves before and after 5000 cycles. (i) *I*-*t* responses of Pt/C and Co/CoN<sub>x</sub>-C after 400s of methanol addition. (j) DFT structure. (k) and (l) Diagram of free energy for the ORR. Reproduced from ref. 53, copyright 2023 Royal Society of Chemistry.

a higher resistance to methanol toxicity. The methanol oxidation process (MOR) over the Pt/C catalyst causes the current density to fluctuate following methanol input. CoNC had a low restricted current density, suggesting that carbon support improves electrical conductivity. In the control studies, the Co/CoN<sub>x</sub>-C ORR activity reduced considerably following acid leaching, which preferentially eliminated metallic Co clusters from the electrocatalyst. Because of the optimal Co/CoN<sub>x</sub>-C composition of 15 wt%, which promotes excellent dispersion of Co species and the cooperative effects of CoN<sub>x</sub> sites and Co clusters, the ORR performance is improved.<sup>59–62</sup>

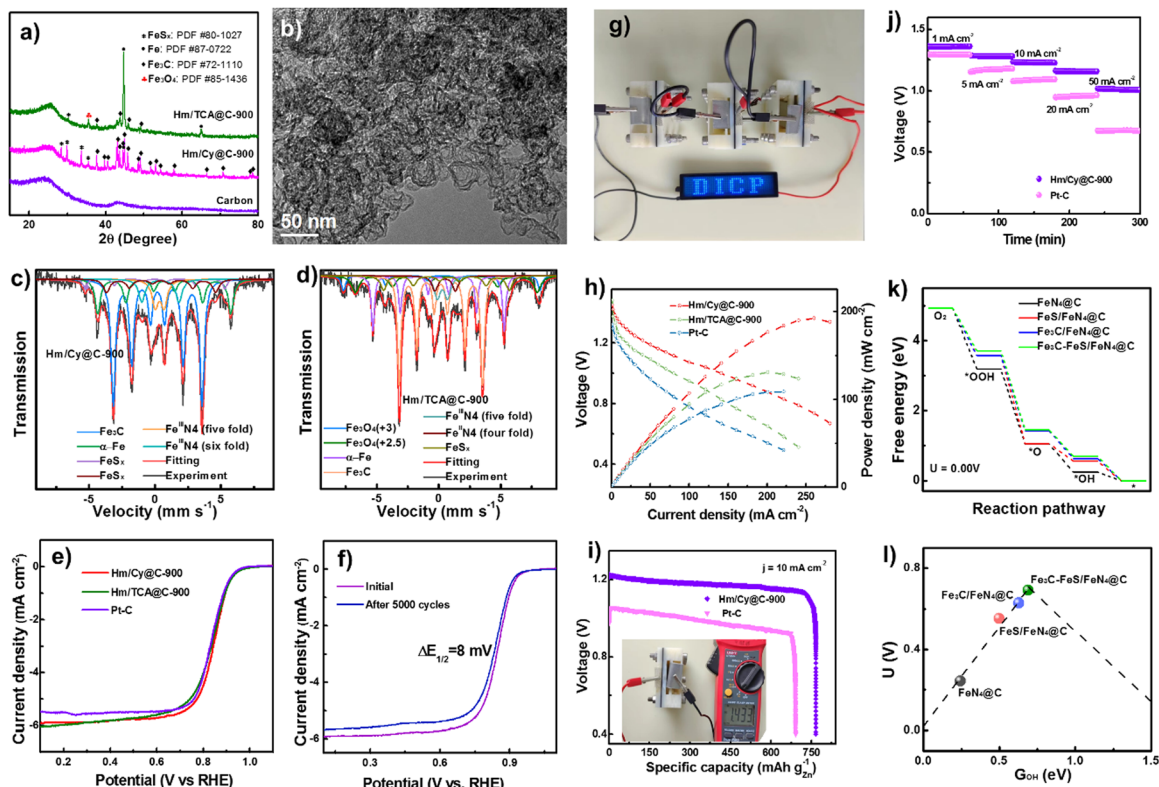
Integration of Co clusters with CoN<sub>x</sub>-C in Co/CoN<sub>x</sub>-C greatly improves stability. The charge density difference indicates several electron clouds surrounding CoN<sub>x</sub> species and Co clusters (Fig. 2j), indicating substantial electron exchange interaction. According to the Bader charge calculations, Co clusters and CoN<sub>x</sub> species in Co/CoN<sub>x</sub>-C transferred 0.35 and 0.96 electrons, respectively; these values are higher than 0.32 and 0.87 electrons transferred in Co/C and CoN<sub>x</sub>-C. The electrocatalytic activity was increased by the interaction and impact of symbiotic Co clusters and CoN<sub>x</sub> species in Co/CoN<sub>x</sub>-C. The rate determining steps for Co/CoN<sub>x</sub>-C and CoN<sub>x</sub>-C occur in the first phase (O<sub>2</sub> → OOH\*), with energy barriers of 0.18 eV and 0.26 eV (*U* = 0 V), according to the ORR process depicted in Fig. 2k and l. Endothermic water generation results from RDS over Co/C in the last step (OH\* → H<sub>2</sub>O). Three Co sites were

assessed for the CoN<sub>4</sub> and Co cluster sites because of the Co/CoN<sub>x</sub>-C's strong ORR performance.<sup>63</sup>

NNME nanomaterials are potential ORR electrocatalysts.<sup>55,64–68</sup> However, the costly and complicated synthetic process will impede NNME expansion. Hemin (Hm) develops ORR-efficient carbon-supported FeN<sub>x</sub> electrocatalysts.<sup>69–71</sup> Iron nitride and sulfide in Hm/Cy@C-900 are confirmed by the overlapping signals of N, S, and Fe species. The different capacities of the N and S-containing functional groups in the Cy and TCA ligands to coordinate could be the cause of the different Fe contents in Hm/Cy@C-900 and Hm/TCA@C-900. The –COOH, –SH, and –NH<sub>2</sub> in N, S-containing ligands bind Fe atoms better than the =S and pyridinic N groups. Next, XRD was used to analyze the electrocatalyst compositions. Carbon ECP exhibited two wide diffraction peaks at ~20–30° and 40–45°, corresponding to graphitic carbon (002) and (101).<sup>72</sup> In contrast, Hm/Cy@C-900 contains Fe<sub>3</sub>C, FeS<sub>x</sub>, and metallic Fe. Hm/TCA@C-900 has a novel Fe<sub>3</sub>O<sub>4</sub> XRD pattern, Fig. 3a, unlike Hm/Cy@C-900. Thus, Hm/TCA@C-900 should include Fe<sub>3</sub>C, FeS<sub>x</sub>, Fe<sub>3</sub>O<sub>4</sub>, and α-Fe nanoparticles. Unlike other iron salts, its metal macrocycle solubility permits iron-based species to self-assemble (Fig. 3b). Solubility-induced scattering of Fe nanoparticles with carbon as active sites was done by Shen *et al.*<sup>73</sup>

Similarly, Hm/Cy@C-900 and Hm/TCA@C-900 composites have 0.39–0.40 at% S. Hm/Cy@C-900 (39.4%) has more C–S–C species than Hm/TCA@C-900 (32.6%), making it more conducive to carbon-S defects. C–S, C–N, and C=N imply that the N and S heteroatoms have been successfully co-doped into the carbon matrixes alongside Fig. 3c's element mapping. The Fe 2p XPS spectrum has low intensity, making it difficult to identify the iron species in the Hm/TCA@C-900 electrocatalyst. So, we used <sup>57</sup>Fe Mössbauer spectra to study the iron components in these Fe-based electrocatalysts. Fig. 3f shows the ORR electrocatalyst performance of Hm/Cy@C-*T* and Hm/TCA@C-900 electrocatalysts with 20 wt% Pt-C. Fig. 3g also shows the Hm/Cy@C-900 electrocatalyst's accelerated durability test. After 5000 cycles, the Hm/Cy@C-900 electrocatalyst showed just an 8 mV change in *E*<sub>1/2</sub>, suggesting it may be durable enough for practical applications. A blue LED screen with DICP writing is illuminated by ZABs that have been built from Hm/Cy@C-900 and coupled in series, as shown in Fig. 3h. In comparison to Hm/TCA@C-900 (131 mW cm<sup>-2</sup>@201.2 mA cm<sup>-2</sup>) and 20 weight percent Pt-C (109 mW cm<sup>-2</sup>@221.3 mA cm<sup>-2</sup>), Fig. 3i demonstrates that Hm/Cy@C-900 had a higher power density of 192 mW cm<sup>-2</sup> at 260.7 mA cm<sup>-2</sup>. These parameters affect the power production and gas diffusion for incorporated electrocatalysts (10 mg cm<sup>-2</sup>) in Zn-air batteries, especially at elevated current densities. With an open-circuit voltage of 1.433 V (Fig. 3j, inset) and a specific capacity of 766.0 mA h g<sup>-1</sup>@10 mA cm<sup>-2</sup>, the Hm/Cy@C-900 electrocatalysts were much more powerful than the Pt-C-assembled battery (691.5 mA h g<sup>-1</sup>@10 mA cm<sup>-2</sup>). In contrast to the ZAB built using Pt-C, the Hm/Cy@C-900 battery displays a consistent discharge voltage at 50 mA cm<sup>-2</sup> (Fig. 3k). Overall, the Hm/Cy@C-900 electrocatalyst performed best in ZAB studies. Computational simulations show that the reduction step of O\* to OH\* is the RDS for FeS/FeN<sub>4</sub>@C in the ORR, but it has a very





**Fig. 3** (a) XRD patterns. (b) High resolution TEM of Hm/Cy@C-900. Mössbauer spectra of (c) Hm/Cy@C-900 and (d) Hm/TCA@C-900. (e) LSV curves. (f) LSV curves of Hm/Cy@C-900 before and after 5000 cycles. (g) Image of ZAB turning on an LED. (h) ZAB polarization and power density. (i) Specific Hm/Cy@C-900 and Pt-C electrocatalyst discharging capabilities. (j) Battery discharge curves at various current densities. (k)  $U = 0.00V$ 's ORR reaction pathway. (l) Free energy volcano diagram with  $\Delta G_{OH}$ . Reproduced from ref. 54, copyright 2023 Royal Society of Chemistry.

strong  $O^*$  chemisorption, leading to a great activation barrier. And the last step of  $OH^*$  to  $OH^-$  is the RDS for  $FeN_4@C$ ,  $Fe_3C/FeN_4@C$ , and  $Fe_3C-FeS/FeN_4@C$ , which have a relatively weak  $O^*$  chemisorption strength and facilitate the ORR process. Furthermore, the offset of d-band center of  $Fe_3C-FeS/FeN_4@C$  was much larger than those of  $FeN_4@C$ ,  $Fe_3C/FeN_4@C$  and  $FeS/FeN_4@C$  systems, indicating that either FeS or  $Fe_3C$  clusters synergize with single sites of  $FeN_4$  to improve the ORR activity.

**OER.** The morphology of ANC- $Fe_xNi_y$  is shown in Fig. 4a using TEM images. The HAADF-STEM picture shows that ANC- $Fe_xNi_y$  has an even size of around 1–2 nm, because of the high nucleation rate and low interfacial tension of ILS.<sup>74,75</sup> According to atomic force microscopy (AFM), ANC- $Fe_1Ni_2$  samples have a thickness of roughly 1.2–1.6 nm. (Fig. 4b), indicating nanosphere shape due to surface IL agent protection.<sup>76,77</sup> The Tyndall effect is well seen in the ANC- $Fe_xNi_y$  solution (Fig. 4c), indicating outstanding dispersibility and subsequent OER electrode construction<sup>78</sup>. The ANC- $Fe_xNi_y$  catalysts' OER performance in 1 M KOH electrolyte was studied to maximize the Fe/Ni ratio. It is revealed in Fig. 3h that the ANC- $Fe_1Ni_2$  cluster has the lowest overpotential (266 mV vs. RHE). LSV curves of the ANC- $Fe_1Ni_2$  catalyst after 1000 and 2000 cycles are shown in Fig. 3(i). The potential losses at  $10 \text{ mA cm}^{-2}$  are 1 and 4 mV. These findings show that ANC- $Fe_1Ni$  has good electrochemical activity and OER endurance.<sup>79,80</sup> The chronopotentiometry measurements (Fig. 3f) reveal that ANC- $Fe_1Ni$  is stable at

$10 \text{ mA cm}^{-2}$  for 6 h. Also, the ANC- $Fe_1Ni$  catalyst's LSV curves are obtained after 1000 and 2000 cycles. The potential losses at  $10 \text{ mA cm}^{-2}$  are 1 mV and 4 mV. These findings show that ANC- $Fe_1Ni$  has good electrochemical activity and OER endurance.

$MnCoO_x$  with a cubic structure was determined by XRD (Fig. 5a).<sup>82,83</sup> XRD patterns of  $MnCoO_x$  samples showed no reflections for  $MnO_x$  species because of the manganese ions' scattering inside the cobaltous matrix, even at high  $MnO_x$  concentrations ( $\sim 40 \text{ mol}\%$ ). The XRD pattern of  $MnCoO_x$  catalysts with different doping levels showed a small shift towards lower angles around  $36.8^\circ$  for (311) as Mn dopants increased, suggesting that Mn cations were well doped into the  $Co_3O_4$  lattice to produce solid solutions. We also examined  $Mn_1Co_5O_x$  annealed at various temperatures using XRD. At  $200^\circ\text{C}$ ,  $Co_3O_4$  developed its crystalline phase. High-temperature calcination enhanced the crystallinity. Despite annealing at  $500^\circ\text{C}$ ,  $Mn_1Co_5O_x$  composites showed no Mn oxide peaks, suggesting thermal stability. TEM was used to study  $MnCoO_x$  solid solutions for elemental composition and morphology. TEM scans showed most oxide particles were 6–8 nm and expose  $Co_3O_4(220)$  and (311) planes (Fig. 5b).<sup>84–86</sup> Interestingly, the  $MnO_x$  species has no lattice fringes, suggesting that Mn was doped in  $Co_3O_4$ .

The linear-scanning profiles in Fig. 5c showed that Mn atoms were scattered inside  $Co_3O_4$ 's lattice. The line sweep region has uniform lattice fringes and a distribution of components that shows that Co and Mn coexist with Co being more





Fig. 4 (a) HAADF-STEM, (b) AFM image with height profiles, and (c) Tyndall effect of the ANC-Fe<sub>1</sub>Ni<sub>2</sub> catalyst. (d) Fe K-edge XANES, (e) Fourier-transformed Fe K-edge, (f) Ni K-edge XANES, and (g) Fourier-transformed Ni K-edge of ANC-Fe<sub>1</sub>Ni<sub>2</sub>. (h) Tafel plots. (i) LSV curves (before and after 1000/2000 CV cycles) of ANC-Fe<sub>1</sub>Ni<sub>2</sub>. (j) Durability tests. (k) Calculated OER free energy diagram. Reproduced from ref. 81, copyright 2022 from Royal Society of Chemistry Royal.

abundant. The elemental mapping (Fig. 5d) supports the homogeneity of Co and Mn species in MnCoO<sub>x</sub> solid solutions. XANES spectra at Co and Mn K-edge were used to study Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub>'s electrical structure and local coordination.<sup>68</sup> Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub> has a peak intensity equal to Co<sub>3</sub>O<sub>4</sub> but distinct from Co-foil (Fig. 5e). Also, Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub>'s pre-line matches Co<sub>3</sub>O<sub>4</sub>. This indicates that Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub>'s primary Co species valence states are Co<sup>2+</sup> and Co<sup>3+</sup>. In contrast, the XANES spectra at Mn K-edge showed that Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub> has a substantially greater peak intensity of the white line than Mn<sub>3</sub>O<sub>4</sub> and Mn<sub>2</sub>O<sub>3</sub>. Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub>'s pre-line moved toward higher binding energies than Mn<sub>2</sub>O<sub>3</sub>, suggesting that its average Mn species valence should be greater than +3. It also shows potassium permanganate oxidation-related electron shortage around the Mn site. The Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub> catalyst's cooperation context was investigated using an *R*-space extended X-ray absorption fine structure (EXAFS) fitting analysis at the Co and Mn K-edges, as depicted in Fig. 5g and h. The Co–O bond distance in Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub> increased from 1.81 Å to 1.83 Å against the control group of CoOOH. In contrast, Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub> reduced the Co–O bond distance from 1.91 Å to 1.83 Å compared to Co<sub>3</sub>O<sub>4</sub>.<sup>81</sup> Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub> reduced the Mn–O bonding distance from 1.80 Å to 1.67 Å compared to Mn<sub>2</sub>O<sub>3</sub>, likely owing to Mn's valence state being greater than +3. The electron density of Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub>'s Co–O bonds was comparable to that of Co<sub>3</sub>O<sub>4</sub>, but somewhat greater than that of CoOOH and shorter. Compared to Mn<sub>2</sub>O<sub>3</sub>, Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub> has a lower electron density in its Mn–O bonds.

This difference in electron density facilitated the creation of a solid-solution structure with a roughly uniform M–O bond length and expedited the flow of electrons between the Mn and Co atoms, hence enhancing OER catalytic activity.

The Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub> composite outperformed Co<sub>3</sub>O<sub>4</sub>, MnO<sub>x</sub>, CP support, and IrO<sub>2</sub> in terms of catalytic efficiency, as indicated by the OER polarization curves (Fig. 5i). This suggests that the OER took place on the surface. Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub> specifically showed overpotentials of 275 and 569 mV at current densities of 10 and 100 mA cm<sup>-2</sup>. (Fig. 5j). Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub>'s increased catalytic OER performance is due to Mn–Co collaboration. The Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub> catalyst had a ~15-fold increase in lifetime compared to Co<sub>3</sub>O<sub>4</sub> at 100 mA cm<sup>-2</sup> (Fig. 5k), suggesting a 20.9% Mn dopant. The CP support is located at 2θ = 26.5°. The Mn<sub>1</sub>Co<sub>5</sub>O<sub>x</sub> catalyst's TEM image exhibited an amorphous layer of ~1 nm thickness (Fig. 5l), confirming Co(OH)<sub>x</sub> production. XPS was also used to analyze the makeup of the pristine surface and its level of oxidation and used MnCoO<sub>x</sub> catalysts. Conversely, the intermediate O\* has the maximum activation energy, which is thermodynamically unfavorable. After Mn doping,<sup>71</sup> the OER overpotential drops from 2.79 eV (Co<sub>3</sub>O<sub>4</sub>(311)) to 0.82 eV (MnCoO<sub>x</sub> (311)). Fig. 5n shows that Mn may raise Co<sub>3</sub>O<sub>4</sub>'s d-band center position from -2.05 eV to near the Fermi level (-1.78 eV). Interestingly, O<sub>v</sub> equilibrates the active site's d-band center (*E*<sub>d</sub> = -1.88 eV). Moderate intermediate species adsorption may reduce the OER reaction energy barrier. Thus, Mn and O<sub>v</sub> stabilize reaction intermediates, lowering the OER reaction





**Fig. 5** (a) XRD patterns. The linear scanning profiles of catalyst's cross-section (c). (b) and (d) HAADF-STEM picture with elemental mapping. XANES spectra of (e) Co and (f) Mn K-edge. (g) OER polarization curves and (h) overpotentials. (i) Chronopotentiometric measurements. (j) STEM picture of used  $\text{Mn}_{1.5}\text{Co}_2\text{O}_x$ . (k) OER routes' reaction-free energies at 0 and 1.23 volts. (l) Co activity sites' d-band center. Reproduced from ref. 52 copyright 2023 from Royal Society of Chemistry.

energy barrier and improving catalytic activity<sup>72</sup> Results of theoretical simulations match experimental phenomena.

We created the hybrid  $\text{MnRuO}_x$  catalyst using  $\text{RuCl}_3$  and  $\text{KMnO}_4$  as metal precursors and controlled crystallization at different annealing temperatures using a simple Pechini approach (Fig. 6a). *Operando* X-ray diffraction (XRD) investigation tracked the catalyst's crystal phase shift in response to temperature to determine how temperature affects crystallization, Fig. 6b.<sup>87</sup> XRD spectra of  $\text{MnRuO}_x$  calcined at 300–350 °C show an amorphous phase without visible precursor peaks.  $\text{MnRuO}_x$ -300 exhibits no distinguishing peaks due to its homogeneous crystal and amorphous dispersion. XRD patterns showed that the  $\text{MnRuO}_x$  hybrid material crystallizes virtually entirely at temperatures over 400 °C, containing Ru oxides. The  $\text{MnRuO}_x$  particle is ~4–5 nm in size (Fig. 6c) and exposes  $\text{RuO}_2$ (110) and (101) planes.<sup>88</sup> An amorphous phase filled the interstitial spaces between the microcrystalline phases, according to the STEM image. The 3D atomic overlap Gaussian fitting findings from the high-angle annular dark-field scanning transmission electron microscopy (HAADF-STEM) pictures showed diverse atomic configurations in different locations.<sup>89</sup> Region 1 had irregular atoms, indicating an amorphous structure (Fig. 6d and e). Region 2 had consistent atom arrangements, suggesting microcrystalline structures (Fig. 6f and g). These findings support the  $\text{MnRuO}_x$ -300 catalyst heterostructures. Thus, the  $\text{MnRuO}_x$  catalyst has both microcrystalline and amorphous structures, with  $\text{RuO}_2$  crystals in

the microcrystalline part. Linear scanning analysis of individual particles revealed a homogenous distribution of Mn and Ru elements, generating a doped phase in the limited size range (Fig. 6h).<sup>68</sup> Local nanoscale elemental line scan investigation shows homogeneous Mn and Ru distribution in solid solution. A large-scale elemental mapping study verifies Mn and Ru's uniform distribution in the catalyst sample.<sup>55</sup>

$\text{MnRuO}_x$  samples were tested for linear sweep voltammetry (LSV) under acidic circumstances and compared to com- $\text{RuO}_2$  (Fig. 6i). The best OER catalyst was  $\text{MnRuO}_x$ -300. At 10  $\text{mA cm}^{-2}$ ,  $\text{RuO}_2$  had an overpotential of 307 mV, consistent with prior results.<sup>91,92</sup> However, a bar chart of the samples' overpotentials at various temperatures showed a significant rise with higher annealing temperatures (Fig. 6j). The  $\text{MnRuO}_x$ -300 catalyst, with its amorphous homogenous structure and microcrystalline particles, had the lowest overpotential (231 mV at 10  $\text{mA cm}^{-2}$  and 282 mV at 50  $\text{mA cm}^{-2}$ ). Additional Tafel slopes were generated to assess catalyst OER kinetics. Comparison of turnover frequencies (TOF) at 1.44 V shows considerable differences between  $\text{RuO}_2$ ,  $\text{MnRuO}_x$ -500, and  $\text{MnRuO}_x$ -300 catalysts (Fig. 6k). The TOF of  $\text{MnRuO}_x$ -300 was 0.071  $\text{s}^{-1}$ , substantially higher than those of  $\text{MnRuO}_x$ -500 (0.0088  $\text{s}^{-1}$ ) and  $\text{RuO}_2$  (0.0030  $\text{s}^{-1}$ ). The catalyst treated at 300 °C has amorphous and crystalline structures, hence it has better OER activity. Stability experiments were performed on  $\text{MnRuO}_x$ -300@TFF samples in acidic circumstances (pH = 0) at varying current





**Fig. 6** (a) Schematic preparation of the  $\text{MnRuO}_x$  catalyst. (b) *Operando* XRD pattern of the  $\text{MnRuO}_x$ -150 precursor treated at different temperatures. (c) HAADF-STEM image of  $\text{MnRuO}_x$ -300 showcases the coexistence of microcrystalline and amorphous phases. 3D atomic overlap Gaussian fitting on selected STEM region 1 (d) and (e) and region 2 (f) and (g). (h) HR-TEM and line scan elemental mapping. (i) Polarization curves, (j) Overpotential bar charts, and comparison of turnover frequency (TOF). (l) Durability tests of  $\text{MnRuO}_x$ -300@TFF in an H-type electrolysis cell. (m) Chronopotentiometry testing at  $100 \text{ mA cm}^{-2}$  in the PEMWE electrolyser at  $80^\circ\text{C}$ . The energy profile of the OER at (n)  $U = 0 \text{ V}$  and (o)  $U = 1.23 \text{ V}$ . (p) Projected DOS. Reproduced with permission from ref. 90, copyright 2024 from Wiley VCH.

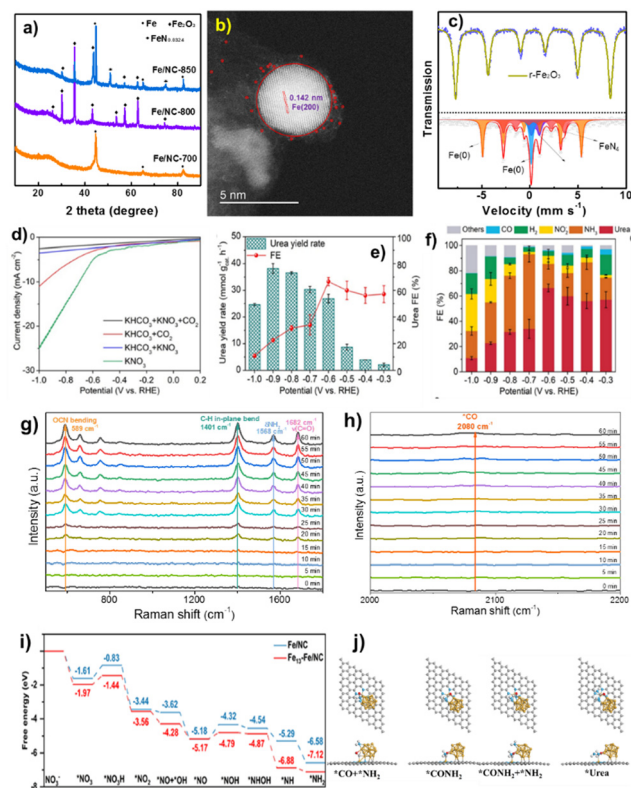
densities using an H-type electrolysis cell. Durability studies of  $\sim 780$  hours at  $100 \text{ mA cm}^{-2}$ ,  $\sim 400$  h at  $200 \text{ mA cm}^{-2}$ ,  $\sim 130$  h at  $500 \text{ mA cm}^{-2}$ , and  $\sim 30$  h at  $1.0 \text{ A cm}^{-2}$  (Fig. 6l) indicate good catalyst stability at high current densities. Under  $100 \text{ mA cm}^{-2}$  for 80 h, the  $\text{MnRuO}_x$ -300 cell maintained PEMWE performance, but a  $\text{RuO}_2$  anode only lasts around 1 hour (Fig. 6m). All reaction steps are endothermic at  $U = 0 \text{ V}$ . At the \*O formation to the \*OOH stage, C/C- $\text{MnRuO}_x$  and C/A- $\text{MnRuO}_x$  heterostructures have OER as the RDS. The OER overpotentials of C/A- $\text{MnRuO}_x$  and C/C- $\text{MnRuO}_x$  are 1.28 eV and 0.96 eV, respectively, at  $U = 1.23 \text{ V}$ , proving that it decreases it. We estimated the predicted density of states (DOS) for C/C- $\text{MnRuO}_x$  and C/A- $\text{MnRuO}_x$  heterostructures (Fig. 6p). The d-band center model describes adsorbate-metal interactions well.<sup>93</sup> After

amorphization, the DOS at the Fermi level rises, raising the d-band center energy level.

### Electrosynthesis of urea with nitrate and carbon dioxide

Simple pyrolysis of  $\text{Fe}^{\text{III}}$ : PVP compounds with carbon ECP at  $700^\circ\text{C}$  in Ar produced the  $\text{FeNC-Fe}_1\text{N}_4/\text{C}$  nanocomposite. Fig. 7a shows that  $\text{FeNC-Fe}_1\text{N}_4$  exhibits metallic Fe particles and the carbon support has high defect and disordered carbon matrix population, promoting electroconductibility and activity during electrocatalysis.<sup>94</sup> It verified that  $\text{FeNC-Fe}_1\text{N}_4/\text{C}$  contained mostly Fe species and the carbon matrix. STEM using EDX elemental mapping technology also examined the distribution of atomically distributed Fe single atoms and metallic Fe clusters. High-resolution TEM images (Fig. 7b) show metallic Fe(200) facets, and the size of Fe clusters is  $\sim 4\text{-}7 \text{ nm}$ .<sup>95</sup>

A typical three-electrode setup in an H-type cell under continuous  $\text{CO}_2$  flow with Nafion 117 membranes between the cathode and anode chambers was used to assess urea electro-synthesis performance. Fig. 7d shows  $\text{FeNC-Fe}_1\text{N}_4/\text{C}$  current densities:  $\text{KNO}_3 > \text{KHCO}_3 + \text{CO}_2 > \text{KHCO}_3 + \text{KNO}_3 > \text{KHCO}_3 + \text{CO}_2$ . Research indicates that the co-electrolysis of  $\text{CO}_2$  and  $\text{NO}_3^-$  for urea production has a lower current density than  $\text{CO}_2\text{RR}$  and  $\text{NO}_3\text{RR}$ . The results show that the  $\text{CO}_2\text{RR}$ ,  $\text{NO}_3\text{RR}$ , and competing HER are effectively regulated, ensuring high urea FE in



**Fig. 7** (a) XRD, (b) STEM image, (c) Mössbauer spectra, (d) LSV curves, (e) potential-dependent urea yield rate, and (f) FEs (at  $-0.6 \text{ V}$ ) of  $\text{FeNC-Fe}_1\text{N}_4/\text{C}$ . (g) and (h) Time-resolved *in situ* Raman spectra in urea electro-synthesis at  $-0.6 \text{ V}$ . (i) Free energy diagrams of  $\text{NO}_3^-$  reduction to  $^*\text{NH}_2$ . (j) Structure diagrams of the C-N coupling process on  $\text{Fe}_{13}\text{-Fe}_1\text{N}_4/\text{C}$ . Reproduced from ref. 91, copyright 2024 from Wiley WCH.



co-reduction.<sup>96</sup> Fig. 7e displays FeNC-Fe<sub>1</sub>N<sub>4</sub>/C sample urea yield rates: 2.1, 3.9, 8.7, 26.9, 30.3, 36.5, 38.2, and 24.6 mmol g<sub>cat.</sub><sup>-1</sup> h<sup>-1</sup> at -0.3, -0.4, -0.5, -0.6, -0.7, -0.8, -0.9, and -1.0 V, respectively. Urea FEs rise and fall when the applied potential is lowered to -1.0 V. Optimal urea production is 38.2 mmol g<sub>cat.</sub><sup>-1</sup> h<sup>-1</sup> at -0.9 V, and an urea FE of 66.5% at -0.6 V surpasses those of existing electrocatalysts. Fig. 7f shows that urea dominates at 60% between -0.3 and -0.6 V, followed by NH<sub>3</sub> and H<sub>2</sub> FEs at 20% and 10%, respectively. When the potential is changed downward, FEs of competing NH<sub>3</sub>, H<sub>2</sub>, and NO<sub>2</sub><sup>-</sup> rise, corresponding to a rapid increase in current density of about -0.6 V in LSV curves.

To understand the complex C- and N-species in co-electrolysis, it is crucial to identify the reaction route and capture important intermediates in C-N coupling.<sup>97</sup> Fig. 7g and h show the time-resolved *in situ* Raman spectra of FeNC-Fe<sub>1</sub>N<sub>4</sub>/C obtained at -0.6 V during urea electrosynthesis. Vibration peaks at 589, 1401, 1568, and 1682 cm<sup>-1</sup> correspond to OCN bending, adsorbed C-H in-plane bending, δ<sub>NH<sub>2</sub></sub>, and C=O stretching modes in formamide. However, <sup>1</sup>H-NMR spectroscopy shows urea production in electrolyte, not formamide. We infer that \*CONH<sub>2</sub> is the intermediary in C-N coupling for urea production. As CO is the reduction product in the CO<sub>2</sub>RR, nucleophilic attack of \*CO and \*NH<sub>2</sub> was used to create C-N coupling. Raman spectra showed the modest signal of \*CO at 2080 cm<sup>-1</sup>, supporting this finding (Fig. 7h). The C-N stretching mode of urea at 1000 cm<sup>-1</sup> is not seen in the Raman spectra owing to its feeble vibration signal.<sup>98</sup> The free energy diagrams in Fig. 7i depict the chemical pathways of NO<sub>3</sub><sup>-</sup> reduction to \*NH<sub>2</sub> on Fe<sub>13</sub>-Fe<sub>1</sub>N<sub>4</sub>/C.

The RDS for Fe<sub>13</sub>-Fe<sub>1</sub>N<sub>4</sub>/C and Fe<sub>1</sub>N<sub>4</sub>/C is \*NO<sub>3</sub> to \*NO<sub>3</sub>H and \*NO to \*NOH, respectively, and the energy barrier of RDS on Fe<sub>13</sub>-Fe<sub>1</sub>N<sub>4</sub>/C is comparatively lower than that on Fe<sub>1</sub>N<sub>4</sub>/C, suggesting that nitrate reduction is more favorable on Fe<sub>13</sub>-Fe<sub>1</sub>N<sub>4</sub>/C. Next, there is increased electron transfer between \*NO<sub>3</sub>, \*NO<sub>3</sub>H and \*COOH intermediates and Fe<sub>13</sub>-Fe<sub>1</sub>N<sub>4</sub>/C. However, the adsorption of the \*COOH intermediate is compromised due to spatial constraints from steric hindrance, as it resides at the interface of the Fe cluster and Fe<sub>1</sub>N<sub>4</sub> site. In all, the Fe cluster and Fe<sub>1</sub>N<sub>4</sub> in Fe<sub>13</sub>-Fe<sub>1</sub>N<sub>4</sub>/C are the primary active sites for nitrate reduction and CO<sub>2</sub> reduction and C-N coupling, respectively.

## Conclusions and future progress

Research on water oxidation and electrocatalysis shows a variety of ways to increase the catalytic stability and activity. They show the synergistic effect of cobalt clusters-CoN<sub>x</sub> composites for electrochemical oxygen reduction, investigating the promotion effect of active sites in iron-based oxygen reduction electrocatalysts for Zn-air batteries, and the high activity for sustainable acidic water oxidation through intrinsically robust cubic MnCoO<sub>x</sub> solid solutions. They also discuss heterogeneous catalysis promoted by ultrafine amorphous metallic hydroxide nanoparticles and oxygen vacancy-rich amorphous FeNi nanoclusters modified by ionic liquids. These findings help develop stable and effective electrocatalysts for energy storage and conversion.

Overall, the highlighted study shows how atomically exact metal nanoclusters and metal oxide clusters may revolutionize electrocatalysis. Research on Fe-based oxygen reduction electrocatalysts has demonstrated their potential to greatly enhance the performance of zinc-air batteries. This Li *et al.* discovery emphasizes the need for several active sites for catalytic effectiveness. Zhang *et al.* showed that cubic MnCoO<sub>x</sub> solid solutions are durable and efficient, demonstrating metal oxide clusters' potential for persistent water oxidation and scalable hydrogen production. Zhang *et al.*'s investigation of the synergistic effect of cobalt clusters-CoN<sub>x</sub> composites on electrochemical oxygen reduction highlights their potential for developing customized catalysts. These examples demonstrate the need to properly manage catalyst's shape and composition to increase catalytic activity and lifespan. In addition, catalyst design and production have been successful, but commercializing and scaling up these advanced electrocatalysts has proven difficult. According to Cao *et al.*, optimizing synthesis processes and studying cheap precursor materials would enable large-scale catalytic material fabrication and deployment. The research found that improving integration and electrochemical systems maximize energy conversion efficiency, lifespan, and cost. These studies demonstrate the revolutionary potential of atomically precise metal nanoclusters and metal oxide clusters in electrocatalysis for sustainable energy technologies, which could lead to cleaner, greener energy sources. Future development may need further electrocatalyst design and manufacturing optimization to increase activity, stability, and selectivity for electrochemical reactions such as oxygen reduction, water oxidation, and energy conversion.

### Essential areas of attention might be:

1. Nanostructuring and active site engineering: researchers are customizing active sites and improving catalytic activity using nanomaterials and atomically precise metal nanoclusters. Identifying catalytic active sites has fascinated academics for decades and is crucial. Active sites may be single atoms or groups of atoms on metal particles. Future research should focus on developing innovative techniques to accurately observe the intricate steps of catalytic reactions at the catalytic site, achieving spatiotemporal accuracy in milliseconds. This endeavor presents a challenging yet highly promising opportunity. Understanding the electronic structure of TMO oxides *via* theoretical studies can help explain catalytic processes. In heterogeneous catalysis, learning the existence and structure of active sites is a huge difficulty, but combining experiments and theory should help. High stability of TMO clusters and novel methods for very stable catalysts throughout catalytic processes may aid mechanistic studies.

2. Synergism in multifunctional materials: composite materials and multifunctional catalysts are being studied to boost catalytic activity and durability. Examining the oxidation state, atom dispersion, and potential behavior of heteroelements is crucial when examining the catalytic properties of the doped element in an alloy catalyst. This information is hard to get. Thus, future research should combine experimentation and theory and build new tools. The link between multifunctional



TMO nanomaterials' geometric and electrical characteristics and catalytic reactivity remains a priority. The fundamental ideas of bimetallic and multimetallic nanocatalysis synergy will influence the composition choices of high-performance nanocatalysts for specific chemical reactions.

3. Understanding reactions using *in situ* and *operando* characterization: a deeper understanding of molecular-level reaction mechanisms might influence selectivity and efficiency-enhancing electrocatalyst design.<sup>99,100</sup> It is important to have a way to analyze TMO catalysts in real time and real space to obtain a more profound comprehension of the catalytic reaction process, its kinetics, and the occurrence of coking and leaching. Studying the high fluxionality of a few TM atom clusters at elevated temperatures and determining whether the nanocluster structure undergoes reversible or irreversible dynamic changes can be achieved through *in situ* and *operando* extended X-ray absorption fine structure (EXAFS) and diffuse reflectance infrared Fourier-transform spectroscopy (DRIFT). Future research should focus on developing advanced *in situ* and *operando* technologies, such as high-resolution imaging and spectroscopy, to better understand and address these challenges. An analysis of this nature would be instrumental in exploring the catalytic factors of TM clusters.

4. Studying new high-entropy oxide clusters: investigation of metal-organic frameworks and ionic liquids as electrolytes and electrocatalysts for devices that convert and store energy. When combined with metal-organic frameworks (MOFs) and ionic liquids (ILs), high-entropy oxide (HEO) clusters—which are defined by several metal cations dispersed randomly inside a single-phase lattice—offer promising advances in energy conversion and storage technologies. Because of their large surface area and adjustable porosity, metal oxide frameworks (MOFs) are a great approach to enhance heteroelectrolytes' ion mobility and structural integrity. Additionally, the metal nodes inside MOFs can function as active catalytic sites. In addition to these hybrids, ILs enhance the electrode-electrolyte interactions by offering a favorable ionic environment due to their high ionic conductivity and stability. The cooperation of HEOs, MOFs, and ILs may be used to create novel materials with enhanced catalytic performance and customized characteristics, greatly enhancing the longevity and efficiency of fuel cells, batteries, supercapacitors, and electrolyzers.

5. Single-atom and single-electron tailoring: atomically precise nanochemistry may govern TMO clusters in new ways, opening up fascinating catalytic research options. Quantum-sized clusters' electronic characteristics are subject to atom count. Customized nanocluster catalysts allow atom-by-atom catalytic reactivity tailoring. Accurate nanocatalysts would not be able to produce novel catalytic phenomena like spin effects in chemical processes, but precise regulation of nanocluster attraction at the single-electron level would.

6. Scale-up and commercialization: this involves improving manufacturing methods and adding advanced electrocatalysts to useful products for affordability, scalability, and sustainability.

The area is well-positioned for sustained innovation and interdisciplinary cooperation to tackle global energy concerns and propel the shift towards a more sustainable energy future.

## Author contributions

This manuscript was collaboratively written by all authors. The final version of the manuscript has been approved by all authors.

## Data availability

Data availability is not applicable to this article as no new data were created or analyzed in this study.

## Conflicts of interest

There are no conflicts to declare.

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