



Cite as
Nano-Micro Lett.
(2023) 15:110

Received: 11 January 2023
Accepted: 29 March 2023
Published online: 30 April 2023
© The Author(s) 2023

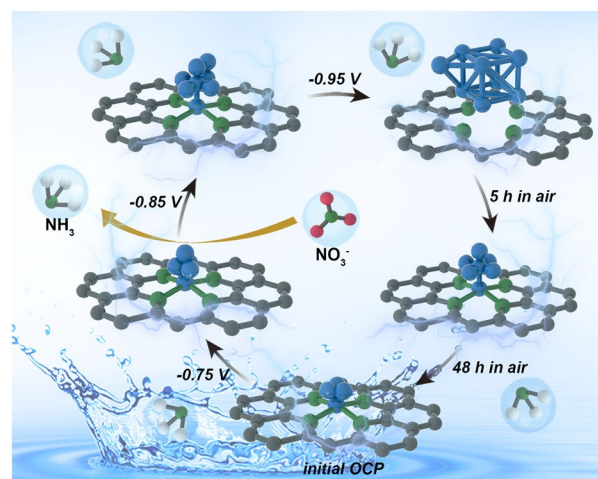
Identification of Dynamic Active Sites Among Cu Species Derived from MOFs@CuPc for Electrocatalytic Nitrate Reduction Reaction to Ammonia

Xue-Yang Ji¹, Ke Sun¹, Zhi-Kun Liu¹, Xinghui Liu^{2,3} ✉, Weikang Dong⁴, Xintao Zuo⁵, Ruiwen Shao⁴, Jun Tao¹ ✉

HIGHLIGHTS

- Cu species with tunable loading supported on N-doped TiO₂/C were successfully fabricated utilizing MOFs@CuPc precursors via the pre-anchor and post-pyrolysis strategy.
- Cu species with tunable loading supported on N-doped TiO₂/C were successfully fabricated utilizing MOFs@CuPc precursors via the pre-anchor and post-pyrolysis strategy.
- Restructured CuN₄&Cu₄ performed the highest NH₃ yield (88.2 mmol h⁻¹ g_{cata}⁻¹) and FE (~94.3%) at -0.75 V due to optimal adsorption of NO₃⁻ and rapid conversion of the key intermediates.

ABSTRACT Direct electrochemical nitrate reduction reaction (NITRR) is a promising strategy to alleviate the unbalanced nitrogen cycle while achieving the electrosynthesis of ammonia. However, the restructuring of the high-activity Cu-based electrocatalysts in the NITRR process has hindered the identification of dynamical active sites and in-depth investigation of the catalytic mechanism. Herein, Cu species (single-atom, clusters, and nanoparticles) with tunable loading supported on N-doped TiO₂/C are successfully manufactured with MOFs@CuPc precursors via the pre-anchor and post-pyrolysis strategy. Restructuration behavior among Cu species is co-dependent on the Cu loading and reaction potential, as evidenced by the advanced *operando* X-ray absorption spectroscopy, and there exists an incompletely reversible transformation of the restructured structure to the initial state. Notably, restructured CuN₄&Cu₄ deliver the high NH₃ yield of 88.2 mmol h⁻¹ g_{cata}⁻¹ and FE (~94.3%) at



✉ Xinghui Liu, liuxinghui119@gmail.com; Jun Tao, taojun@bit.edu.cn

¹ Key Laboratory of Cluster Science of Ministry of Education, School of Chemistry and Chemical Engineering, Liangxiang Campus, Beijing Institute of Technology, Beijing 102488, People's Republic of China

² Department of Chemistry, Sungkyunkwan University (SKKU), Suwon 16419, Republic of Korea

³ Department of Materials Physics, Saveetha School of Engineering, Saveetha Institute of Medical and Technical Sciences (SIMTS), Thandalam, Chennai, Tamilnadu 602105, India

⁴ Beijing Advanced Innovation Center for Intelligent Robots and Systems and Institute of Engineering Medicine, Beijing Institute of Technology, Beijing 100081, People's Republic of China

⁵ Key Laboratory of Bio-inspired Smart Interfacial Science and Technology of Ministry of Education, School of Chemistry, Beihang University, Beijing 100191, People's Republic of China



– 0.75 V, resulting from the optimal adsorption of NO_3^- as well as the rapid conversion of $^*\text{NH}_2\text{OH}$ to $^*\text{NH}_2$ intermediates originated from the modulation of charge distribution and d -band center for Cu site. This work not only uncovers CuN_4 & Cu_4 have the promising NITRR but also identifies the dynamic Cu species active sites that play a critical role in the efficient electrocatalytic reduction in nitrate to ammonia.

KEYWORDS Metal–organic frameworks; Copper phthalocyanine; Electrocatalytic nitrate reduction reaction

1 Introduction

Ammonia (NH_3) is an indispensable chemical used mainly in the manufacture of fertilizer, the synthesis of nitric acid and nitrate, and a refrigerant [1, 2]. The Haber–Bosch process, which utilizes N_2 and H_2 as input gases in a demanding environment (high temperature and pressure), is widely recognized as one of the greatest breakthroughs of the twentieth century and accounts for the overwhelming majority of yearly NH_3 production [3, 4]. Compared to the Haber–Bosch process (high CO_2 emissions and energy consumption), an environment-friendly NH_3 synthesis by an electrocatalytic N_2 reduction reaction (NRR) has garnered conspicuous attention, which enables the electrosynthesis of NH_3 [5, 6]. Furthermore, nitrate (NO_3^-) is the usual contaminant in surface and ground water, exacerbating soil and water eutrophication and posing a threat to the environment and human health [7]. It could be an intriguing tactic for electrocatalytic NO_3^- reduction reaction (NITRR) to NH_3 under ambient circumstances via an eight-electron transfer reaction [8–10]. Hence, seeking low-cost, excellent selective, and efficient electrocatalysts capable of enabling both NH_3 production and NO_3^- conversion rate is critical.

Single-atom catalysts have been extensively recruited in various catalytic reactions due to their maximized atomic utilization and coordination diversity [11–13]. To date, metal–organic frameworks (MOFs) have been selected as the platform for the direct or indirect fabrication of single-atom catalysts on account of the multifarious anchoring mechanisms (e.g., spatial confinement, structural defects, and coordination strategies) in comparison with carbon nanotubes and graphene-based supported materials [14]. Among amine-functionalized MOFs, e.g., $\text{NH}_2\text{-UiO66}$ Zr, $\text{NH}_2\text{-MIL125}$ Ti (here denoted as aMIL), and $\text{NH}_2\text{-MIL101}$ Al, the uncoordinated amine groups ($-\text{NH}_2$) as the electron donors can stabilize and anchor single metal to synthesize single-atom electrocatalysts with well-defined metal- N_4 structure [15–18].

In general, transition-metal atoms are susceptible to migrate and aggregate during pyrolysis with the high temperature-induced effect, forming metal nanoclusters and/or nanoparticles, which is always a thorny problem that seriously affects the targeted synthesis of highly loaded metal single-atom catalysts [19]. In addition, among the single-atom catalysts, Cu electrocatalysts can be restructured and induced by the special potential to form Cu nanoclusters and nanoparticles during electrocatalytic reduction reactions. In situ-formed metal nanoclusters or nanoparticles modulate the d -band center of the metal- N_4 active site, optimize its electronic structure and improve catalytic performance [20, 21]. For instance, Fu and co-workers reported the transformation from Cu single-atom catalysts (CuN_4 configuration) to the compound of Cu single-atom and nanoclusters during the alkaline oxygen reduction reaction (ORR). The restructured Cu-nanoclusters can regulate the d -band center of CuN_4 and balance the free energy of intermediates, benefiting ORR activity [22]. Liu and co-workers found the reversible transformation of atomically dispersed Cu atoms and Cu_3/Cu_4 clusters during the alkaline carbon dioxide reduction reaction (CO_2RR) [23]. In applying NITRR, Wang and co-workers observed the dynamic reversible transformation between Cu single-atom catalysts (Cu loading: 1.0 wt%) and Cu_9 nanoclusters with aggregation driven by the potential in an alkaline electrolyte and redispersion driven by oxidation at environmental conditions. Cao and co-workers verified in situ clustering of single-atom Cu in MOFs (Cu loading: 1.9 wt%) in Na_2SO_4 and NaNO_3 electrolytes [24, 25]. In light of the aforementioned development, exploring the dual-driven behaviors of Cu species loading and reaction potential for electrocatalytic NH_3 production is essential, especially for identifying the dynamical active sites.

Herein, Cu species (single-atom, clusters, and nanoparticles) anchored on N-doped TiO_2/C (NTC) were synthesized via the pre-anchor and post-pyrolysis strategy. As a result, the restructuration behavior of Cu species was jointly related to Cu loading and reaction potential during electrocatalytic NITRR. Specifically, the higher copper loading

and negative potential will more easily transform copper single atoms into copper clusters and particles. Cu1.5/NTC exhibited the highest NH_3 yield with $88.2 \text{ mmol h}^{-1} \text{ g}_{\text{cata}}^{-1}$ ($0.044 \text{ mmol h}^{-1} \text{ cm}^{-2}$) and FE ($\sim 94.3\%$) at the potential of -0.75 V versus RHE, which was superior to NTC, Cu0.7/NTC, and Cu3.2/NTC electrocatalysts. *Operando* XAS and density functional theory (DFT) calculation results confirmed that the restructured $\text{CuN}_4\&\text{Cu}_4$ can enhance the adsorption ability of NO_3^- and promote the rapid potential-determining step conversion of $^*\text{NH}_2\text{OH}$ to $^*\text{NH}_2$ intermediates.

2 Experimental Section

2.1 Preparation of aMIL@CuPc-x Precursors

The synthesis of NH_2 -MIL125 (Ti) refers to our previous work with minor modifications [26]. In the synthesis of NH_2 -MIL125 (Ti)@CuPc precursor (aMIL@CuPc-x), NH_2 -BDC (840 mg, 4.6 mmol), $\text{C}_{12}\text{H}_{28}\text{O}_4\text{Ti}$ (0.90 mL, 3.3 mmol), and a series of amounts of $\text{C}_{32}\text{H}_{16}\text{CuN}_8$ (CuPc) were successively added into the mixed solution containing anhydrous DMF and MeOH (60 mL, v/v = 9:1), and then stirred magnetically at room temperature for 1 h. Subsequently, the resulting solution was transferred to a Teflon-lined stainless steel autoclave and heated at $150 \text{ }^\circ\text{C}$ for 72 h until cooling to the room temperature. DMF and MeOH were alternately washed three times and centrifuged (8,000 rpm, 6 min), and then dried at $120 \text{ }^\circ\text{C}$ under dynamic vacuum for 24 h to obtain the final precursor powder. The additive amounts of CuPc were aMIL (0 mg), aMIL@CuPc-1 (46 mg, $\sim 0.08 \text{ mmol}$), aMIL@CuPc-2 (92 mg, $\sim 0.16 \text{ mmol}$) and aMIL@CuPc-3 (184 mg, $\sim 0.32 \text{ mmol}$), respectively.

2.2 Preparation of Derivatives

The powder precursors were calcined at $800 \text{ }^\circ\text{C}$ with the rate of $2 \text{ }^\circ\text{C min}^{-1}$ for 3 h in Ar. N-doped TiO_2/C (NTC) with different Cu loading could be prepared by the precursors of pyrolysis aMIL@CuPc-x. ICP-OES showed that the Cu content was 0.673 (~ 0.7), 1.540 (~ 1.5) and 3.242 (~ 3.2) wt%, respectively. The samples were named in turn

as NTC, Cu0.7/NTC, Cu1.5/NTC, and Cu3.2/NTC. The entire synthesis process is shown in Scheme S1.

2.3 Electrochemical Measurements

All NITRR electrochemical measurements were performed on an electrochemical workstation (Ivium Vertex, Netherlands) in a three-electrode system using the H-type electrolytic cell separated by a Nafion 117 membrane (DuPont) at the room temperature (Fig. S1). Nafion 117 membrane was pretreated in the 5 wt% H_2O_2 solution at $80 \text{ }^\circ\text{C}$ for 1 h, rinsed by ultrapure water and then in ultrapure water at $80 \text{ }^\circ\text{C}$ for another 1 h, rinsed with ultrapure water several times. Typically, as-prepared sample (5 mg) was dispersed in 900 μL of isopropanol (IPA) aqueous solution (v/v, 1:1) and 100 μL Nafion solution (5%), and ultrasonicated for 30 min. The homogeneous ink was dropped onto the commercial carbon paper (1 cm \times 1 cm, mass loading: ca. 0.5 mg cm^{-2}). Pt plate (1 cm \times 1 cm), saturated calomel electrode (SCE) and above carbon paper were used as the counter, reference and working electrode in 0.5 M Na_2SO_4 and 50 ppm NaNO_3 mixed solutions. The potentiostatic test was conducted at constant potentials for 2 h with a flow rate of Ar for 10 standard cubic centimeters per minute (sccm) and a stirring rate of 400 rpm. Cyclic voltammetry activation was not performed before the experiment, and the pH changes before and after the experiment were ignored. The potential reference equation (Eq. 1) was converted into the reversible hydrogen electrode (RHE).

$$E_{\text{RHE}} = E_{\text{SCE}} + 0.241 \text{ V} + 0.0591 \times \text{pH} \quad (1)$$

2.4 Operando XAS Experiment

The NITRR operando XAS experiment was tested in fluorescence mode at a BSRF 4B9A line station and used the home-built H-type cell referring to the NITRR experimental condition. Where the electrolyte was a mixed aqueous solution with Ar-saturated 0.5 M Na_2SO_4 and 50 ppm NaNO_3 , and the ink is uniformly dropped on a carbon paper of 1 cm \times 2 cm (with a mass of about 2.5 mg cm^{-2}), and the operando XAS experiment is started at a specific voltage without the cyclic voltammetry (CV) activation.

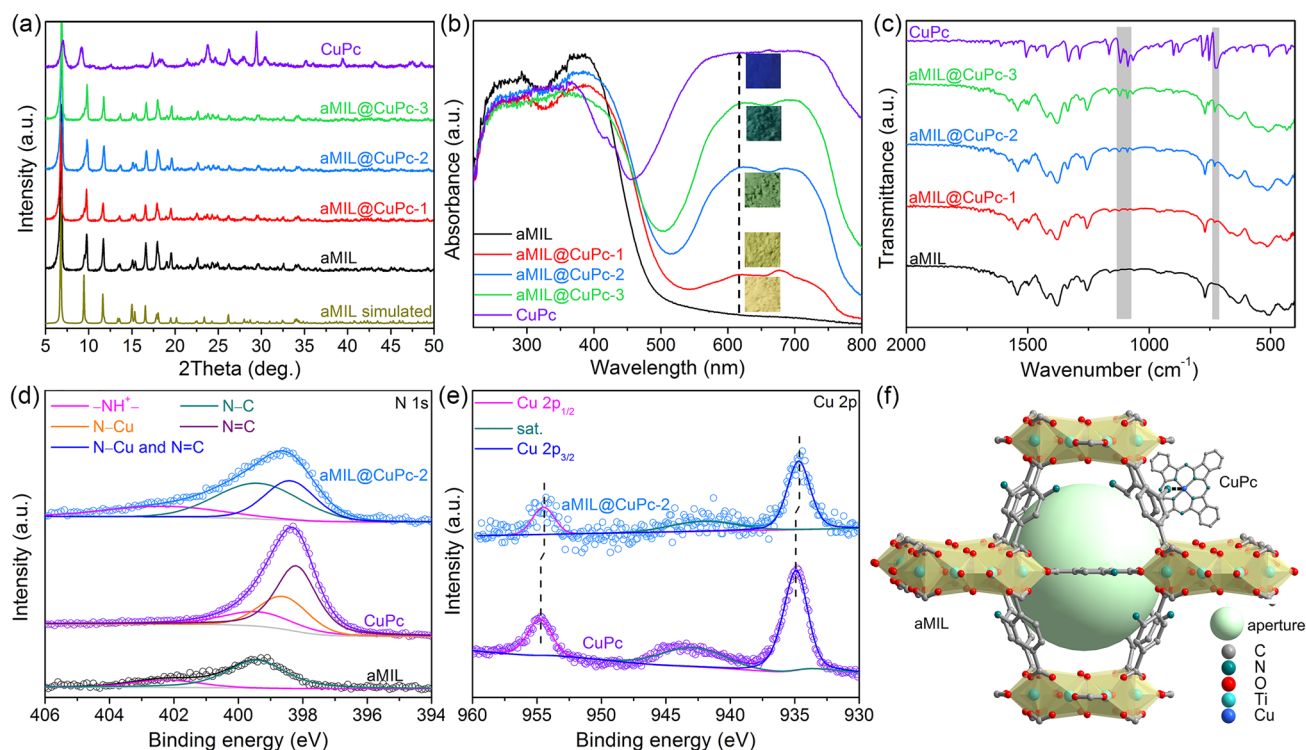


Fig. 1 a–c PXRD patterns, UV–vis DRS and optical photograph, FT-IR spectra of aMIL, aMIL@CuPc-x, CuPc samples. d, e Typical high-resolution XPS spectra of N 1s and Cu 2p of aMIL, CuPc and aMIL@CuPc-2 samples. f Assembly mechanism between aMIL and CuPc

3 Results and Discussion

3.1 Characterization of MOFs@CuPc-x Precursors

All of the experimental powder X-ray diffraction (PXRD) patterns of the as-synthesized aMIL@CuPc-based precursors (Fig. 1a) agree well with the simulated one of pure aMIL due to the similarity of the strong diffraction peaks between aMIL and CuPc or the lower mass loading of CuPc. Increased visible-region absorption and a gradual darkening of color with increasing CuPc loading are observable in the UV–vis diffraction spectra (DRS) and corresponding optical photographs (Fig. 1b) of aMIL, aMIL@CuPc-1, aMIL@CuPc-2, aMIL@CuPc-3, and CuPc samples, proving the presence of aMIL and CuPc composite. The vibrational peaks of aMIL@CuPc-based precursors in Fourier transform infrared (FT-IR) spectra (Fig. 1c) at approximately 1116, 1086, and 725 cm^{-1} correspond to C–H in-plane deformation ($\beta_{(\text{C-H})}$), C–N stretching in pyrrole ($\nu_{(\text{C-N})}$), C–N out-of-plane deformation ($\gamma_{(\text{C-N})}$) except for the characteristic peaks of aMIL [27]. To uncover the link mode between aMIL and

CuPc molecules, X-ray photoelectron spectroscopy (XPS) was characterized for aMIL, CuPc, and aMIL@CuPc-2 precursors. The presence of $-\text{NH}^+$ (402.3 eV) and N–C bonds (399.4 eV) among the $-\text{NH}_2$ group in aMIL is confirmed by typical high-resolution XPS spectra of N 1s (Fig. 1d) [26]. In addition, there exist N–C bond (399.4 eV), pyrrolic N–Cu coordination form (398.6 eV), and N=C bond (398.2 eV) for the CuPc sample [28]. Due to the proximate binding energy of N–Cu and N=C bonds and higher half-peak breadth of N–C bonds given by both aMIL and CuPc, pyrrolic N–Cu and N=C bonds were joint fitted through deconvolution locating at 398.4 eV. Significantly, typical high-resolution XPS spectra of Cu 2p of aMIL@CuPc-2 slightly shift to the lower binding energy compared with that of CuPc in view of the electron-donation effect of $-\text{NH}_2$ group [29], the axial N of $-\text{NH}_2$ group over aMIL could be coordinated with CuPc to form $[\text{Ti}_8(\text{OH})_4\text{O}_8(\text{BDC-NH}_2)_{6-x}][-(\text{BDC-NH})_x-(\text{CuPc})_x]$ (Fig. 1f). Compared to pure CuPc, the average coordination number (CN) of Cu–N bond over the aMIL@CuPc-2 increases from 3.72 to 4.20 (Fig. S4, Table S1). Meanwhile, aMIL@CuPc-based precursors retain the tetragonal plate

shape of the aMIL matrix (Fig. S5), while CuPc molecules can form nanorod structures due to self-nucleation with higher CuPc loading [30].

3.2 Structural Characterization of Cu_x/NTC Derivatives

The rational design of precursors with variable loading of CuPc affects the dispersity of Cu species among aMIL@CuPc-based derivatives to a certain extent. The PXRD patterns (Fig. S6, Table S2) substantiate the existence of

the mixed crystal phase with Anatase and Rutile TiO_2 , and the mass ratio ($m_{Anatase}: m_{Rutile}$) is 2.34. Besides, a distinct diffraction peak exists at 43.3° of metallic Cu phase (JCPDS No: 04–0836) just for Cu3.2/NTC sample. Brunauer–Emmett–Teller (BET) specific surface area, pore volume, and Raman spectra (Fig. S7, Table S3) of NTC, Cu0.7/NTC, Cu1.5/NTC, and Cu3.2/NTC samples show that Cu1.5/NTC possess the biggest specific surface area, larger pore volume and the degree of graphitization, which can provide more active sites and enhance the conductivity of supported materials to accelerate the mass transfer of NITRR. The characteristic peaks of the high-resolution XPS

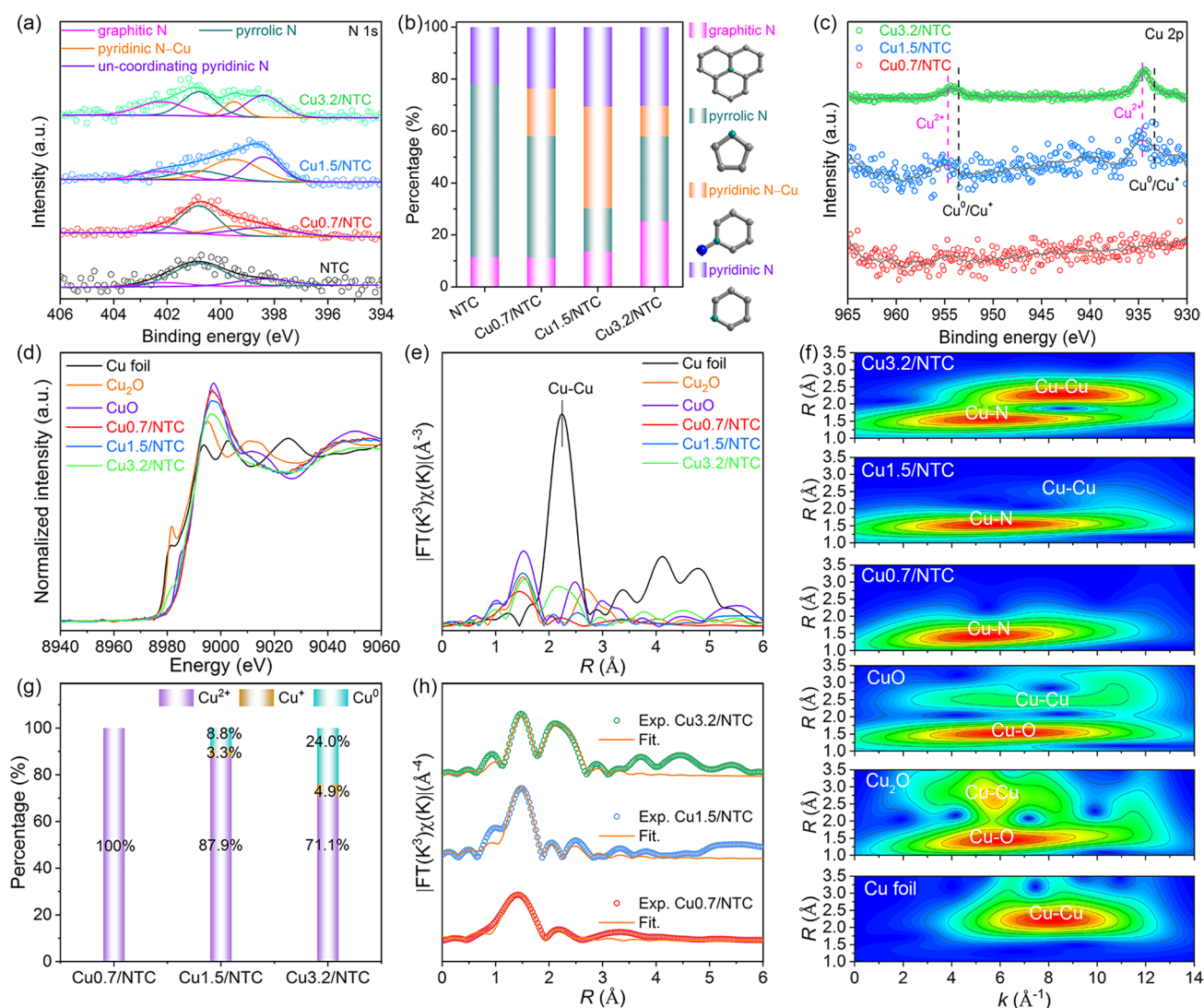


Fig. 2 **a** High-resolution XPS of N 1s; **b** N contents and configurations; **c** Cu 2p for Cu0.7/NTC, Cu1.5/NTC, and Cu3.2/NTC. **d–f** Cu K-edge XANES, k^3 -weighted FT-EXAFS, and WT-EXAFS spectra of Cu foil, Cu₂O, CuO, Cu0.7/NTC, Cu1.5/NTC, Cu3.2/NTC. **g, h** XANES LCF and FT-EXAFS spectra fitting curves of Cu0.7/NTC, Cu1.5/NTC, Cu3.2/NTC



N 1s spectra (Fig. 2a) at 402.2, 400.8, 399.5, and 398.4 eV correspond in turn to graphitized N, pyrrole N, pyridine N–Cu, and un-coordinating pyridine N, revealing the presence of N-doped C and Cu–N species [28]. Additionally, the content of pyrrole N–Cu of Cu3.2/NTC sample is lower than Cu0.7/NTC along with the progressive introduction of CuPc, indicating that Cu3.2/NTC contains a small amount of Cu–N species. High-resolution XPS of Cu 2p spectrum of Cu0.7/NTC (Fig. 2c) has a low signal-to-noise ratio, which can be attributed to the low Cu content. The peaks of Cu1.5/NTC at 955.0 and 934.7 eV can be described as Cu 2p_{1/2} and Cu 2p_{3/2} of Cu²⁺ [31], and the peaks of Cu3.2/NTC samples at 953.5 and 933.3 eV are Cu 2p_{1/2} and Cu 2p_{3/2} of Cu⁰ or Cu⁺ [32]. The high-resolution XPS of Cu 2p spectra for both Cu1.5/NTC and Cu3.2/NTC display a

large half-peak width, indicating mixed valence states of Cu with 0, +1, and +2. Further pre-edge and white line peaks of the X-ray absorption near-edge structure (XANES) spectra at Cu K-edge (Fig. 2d) suggest that the average valence states of Cu successively are +2 ≥ Cu0.7/NTC > Cu1.5/NTC > Cu3.2/NTC > +1. As shown in the *k*³-weighted Fourier-transformed extended X-ray absorption fine structure (FT-EXAFS) and wavelet transform extended X-ray absorption fine-structure (WT-EXAFS) spectra (Fig. 2d, f), Cu0.7/NTC, Cu1.5/NTC, and Cu3.2/NTC have the Cu–N scattering path in the first shell, while Cu1.5/NTC and Cu3.2/NTC sample show the Cu–Cu scattering path in the second shell compared with Cu foil, Cu₂O, and CuO reference samples. The proportions of Cu²⁺, Cu⁺, and Cu⁰ for Cu1.5/NTC are 87.9%, 3.3% and 8.8%, the corresponding proportions for

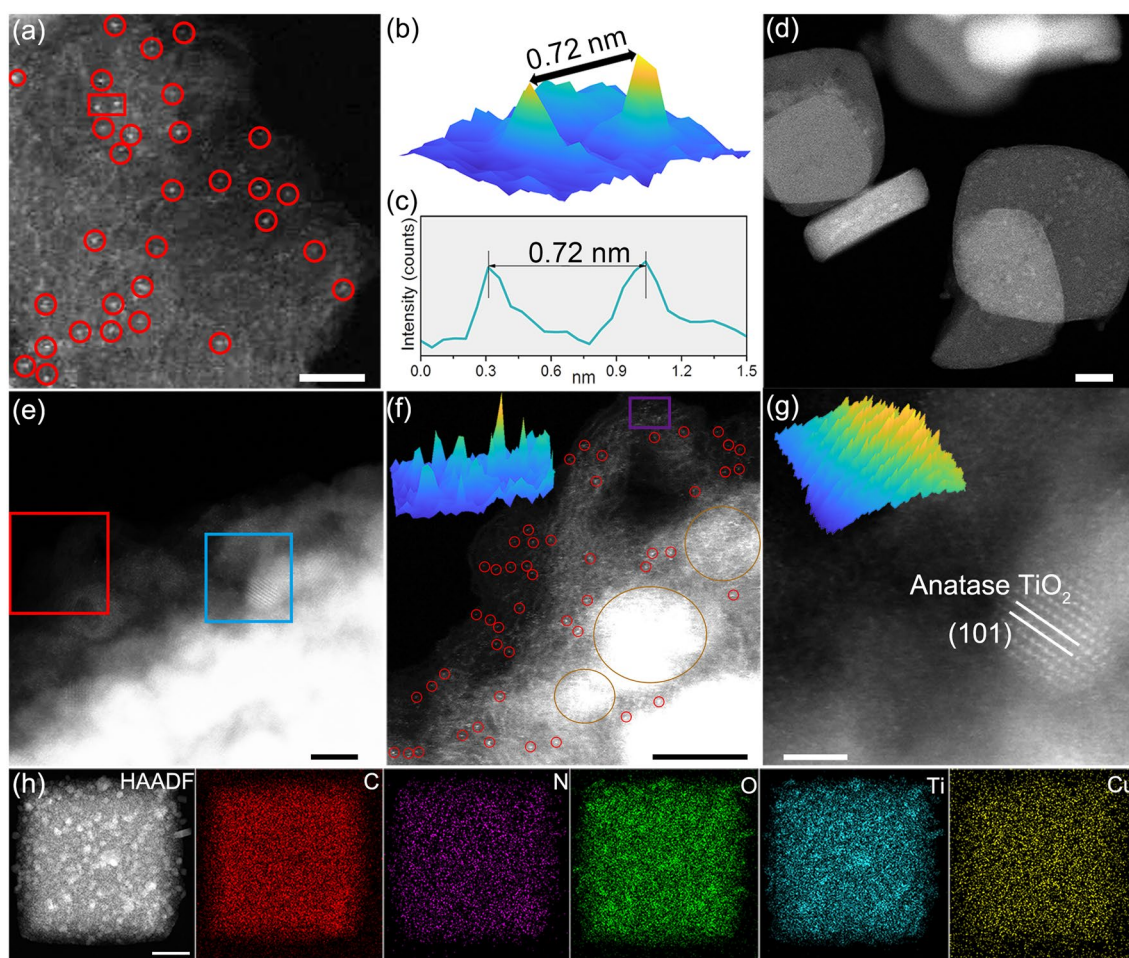


Fig. 3 **a** AC HAADF-STEM image of Cu0.7/NTC. **b, c** 3D and line profile images of the red rectangular area in **a**. **d–g** HAADF-STEM images of Cu1.5/NTC, and **f, g** are the enlarged images of the red and blue rectangular areas in **e**. **h** EDX mapping images of Cu1.5/NTC. Scale bar: **a** 2 nm, **e–g** 5 nm, **d, h** 200 nm

Cu_{3.2}/NTC are 71.1%, 4.9% and 24.0% in the linear combination fitting (LCF) results of XANES spectra (Figs. 2g and S10), manifesting the generation of metallic Cu with the high temperature-induced effect during pyrolysis. The average CN of Cu–N bond (Figs. 2h and S11, Table S4) for Cu_{0.7}/NTC, Cu_{1.5}/NTC, and Cu_{3.2}/NTC are 3.99 (~4), 4.02 (~4), and 3.61 (<4), and the average CN of Cu–Cu bonds for Cu_{1.5}/NTC and Cu_{3.2}/NTC samples are 2.84 (~3) and 7.41 (~7) based on the proportion of Cu⁰ in XANES LCF.

3.3 Morphology of Cu_x/NTC Derivatives

The dispersion state of Cu species and supported materials among Cu_{0.7}/NTC, Cu_{1.5}/NTC, and Cu_{3.2}/NTC were observed by aberration-correction

high-angle-annular-dark-field scanning transmission electron microscopy (AC HAADF-STEM) and high-resolution transmission electron microscopy (HRTEM). AC HAADF-STEM image (Fig. 3a–c) of Cu_{0.7}/NTC reveals that Cu is monodispersed on the NTC and the distance of neighbored Cu atoms is relatively large (~0.72 nm). AC HAADF-STEM images of Cu_{1.5}/NTC (Fig. 3d–f) provide vital evidence for the coexistence form of Cu species with both monodispersity and clusters (small red cycles and big dark yellow circles in Fig. 3f). Further, Fig. 3g (the enlarged image of the blue rectangular area of Fig. 3e) and HRTEM images of Cu_{1.5}/NTC (Fig. S13) uncover the existence of both anatase and rutile TiO₂ on the surface of N-doped C supported materials, which further confirmed with the well-defined elements distribution of C, N, O, Ti, and Cu (Fig. 3h). Furthermore, Cu nanoparticles with exposed (111) crystal plane load on the

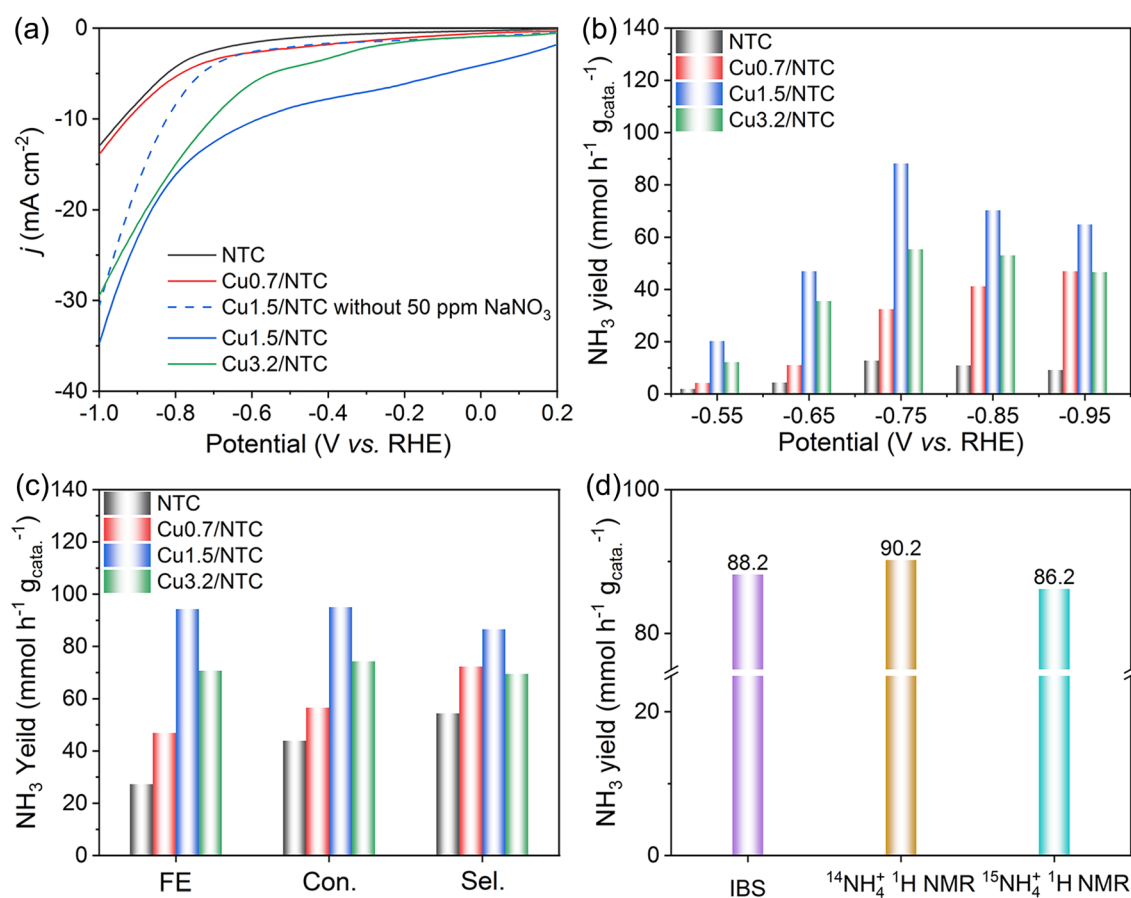


Fig. 4 **a** LSV polarization curves of NTC, Cu_{0.7}/NTC, Cu_{1.5}/NTC and Cu_{3.2}/NTC in 0.5 M Na₂SO₄ electrolytes with 50 ppm NaNO₃, and without 50 ppm NaNO₃ for Cu_{1.5}/NTC. **b** NH₃ yield of NTC, Cu_{0.7}/NTC, Cu_{1.5}/NTC and Cu_{3.2}/NTC. **c** The comparison of FE, conversion rate (Con.) and selectivity (Sel.) of NTC, Cu_{0.7}/NTC, Cu_{1.5}/NTC and Cu_{3.2}/NTC samples at -0.75 V. **d** Comparison of NH₃ yield via IBS and ¹H NMR methods

surface of NTC material for Cu_{3.2}/NTC sample, integrating with the EDX mapping images (Fig. S14).

3.4 Electrocatalytic NITRR Performances

The NITRR performances of NTC, Cu_{0.7}/NTC, Cu_{1.5}/NTC, and Cu_{3.2}/NTC electrocatalysts were evaluated in 0.5 M Na₂SO₄ and 50 ppm NaNO₃ electrolytes. As exhibited in the linear sweep voltammetry (LSV) polarization curves (Fig. 4a), the current density (*j*) of Cu_{0.7}/NTC, Cu_{1.5}/NTC, and Cu_{3.2}/NTC are higher than that of NTC, and Cu_{1.5}/NTC has the highest current density in the range of 0.2 to −1.0 V, indicating that the introduction of appropriate Cu species can promote the NITRR process. Compared to the LSV curves of Cu_{1.5}/NTC before and after adding 50 ppm NaNO₃, the potential range from −0.55 to −0.95 V versus RHE was selected to quantify the NH₃ production performance. After the potentiostatic experiment at the potentials of −0.55, −0.65, −0.75, −0.85, and −0.95 V for 2 h, the products of NH₃, NO₂[−] and catalysis substrate (NO₃[−]) were detected via colorimetric methods (Figs. S15–S17). Cu_{1.5}/NTC delivers best NH₃ yield (88.2 mmol h^{−1} g_{catal}^{−1}) and lower NO₂[−] yield at −0.75 V, which is superior to NTC, Cu_{0.7}/NTC and Cu_{3.2}/NTC at this potential (Figs. 4b and S18). Meanwhile, Cu_{1.5}/NTC presents the optimal FE (94.3%), 95.0% conversion rate, and 86.6% selectivity of NO₃[−] to NH₃ (Fig. 4c), which confirms that Cu is the main active site of NITRR, agreeing well with that the introduction of appropriate Cu species significantly increased the *A*_{ECSA} to enhance the performance of electrocatalytic NH₃ production (Fig. S19). Meanwhile, NH₃ yield and FE for Cu_{1.5}/NTC show no significant decay during the continuous long-term tests at −0.75 V (Fig. S20). To ensure the N source and accurately estimate NH₃ yield, multi-position in one mode of ¹⁵N isotope labeling experiment, indophenol blue spectrophotometric (IBS), and ¹H nuclear magnetic resonance (NMR) methods were engaged by conducting the product from the potentiostatic experiment for Cu_{1.5}/NTC at −0.75 V using blank Na₂SO₄ solution, Na¹⁴NO₃ and Na¹⁵NO₃ as N sources (Figs. S21–S23). Based on the prominent characteristic triple and double peaks of ¹⁴NH₄⁺ and ¹⁵NH₄⁺ (Figs. S22 and S23) [33], NH₄⁺ yield quantified with ¹H NMR method is very close to the results of IBS method (Fig. 4d). The reliable NH₃ yield and FE of Cu_{1.5}/NTC executed in neutral electrolyte are on par with the majority of published Cu-based electrocatalysts (Table S5).

3.5 Restructuration Process Detected with *Operando* XAS

Operando XAS experiment is frequently employed to accurately record the dynamic structural changes of metal activity sites during electrocatalysis reactions of single-atom catalysts [34, 35]. *Operando* XAS experiments of Cu_{0.7}/NTC, Cu_{1.5}/NTC, and Cu_{3.2}/NTC were conducted at open circuit potential (OCP), −0.55, −0.65, −0.75, −0.85, and −0.95 V versus RHE for Cu K-edge using the home-built H-type cell referring to the NITRR experimental condition, and Cu foil, Cu₂O, CuO, Cu(acac)₂, CuPc are reference samples. There are no obvious differences in the structure and valence state at different potentials in the XANES and 1st derivative of XANES spectra for Cu_{0.7}/NTC electrocatalyst (Fig. 5a, d). The characteristic peaks in the XANES spectra of Cu_{1.5}/NTC (Fig. 5b) at 8988.7 eV are attributed to the electron transition from Cu²⁺ 1s to Cu²⁺ 4p, while the peaks of −0.85 and −0.95 V located at ~8982 eV originate the electron transition from Cu⁺ 1s to Cu⁺ 4p_y [36], which indicate that Cu_{1.5}/NTC electrocatalyst begins undergoing the obvious transition from Cu²⁺ to Cu⁺ at −0.85 V. The 1st derivative of XANES spectra (Fig. 5e) at the potentials of −0.85 and −0.95 V have a big half-peak width at ~8979 and 8980 eV [37], revealing that Cu_{1.5}/NTC has the mixed valence states of Cu⁰, Cu⁺, and Cu²⁺ at these potentials. The characteristic peaks of Cu⁺ in the XANES spectra and Cu⁰ in the 1st derivative of XANES spectra for Cu_{3.2}/NTC at all the potentials (Fig. 5c, f) become more robust with the negative shift of reaction potential, suggesting that the Cu species among Cu_{3.2}/NTC experience the significant reduction process of Cu²⁺ → Cu⁺ → Cu⁰.

To understand the valence state and coordination environment of Cu species thoroughly for Cu_{0.7}/NTC, Cu_{1.5}/NTC, and Cu_{3.2}/NTC electrocatalysts, we pay more attention to resolving the FT-EXAFS, XANES LCF, and WT-EXAFS spectra at the representative potentials of OCP, −0.75, −0.85, and −0.95 V. The *k*³-weighted FT-EXAFS fitted spectra together with the fitting results of XANES LCF spectra (Figs. S24 and S25, Table S6) show that Cu_{0.7}/NTC still keeps the dominant configuration of Cu–N₄ at OCP, −0.75, −0.85, −0.95 V. However, the partial Cu⁺–N_x generated at the −0.75 V potential could detach from the anchorage of N-doped C among Cu–N₄ and undergo the significant restructuring driven by the potential at −0.85 V

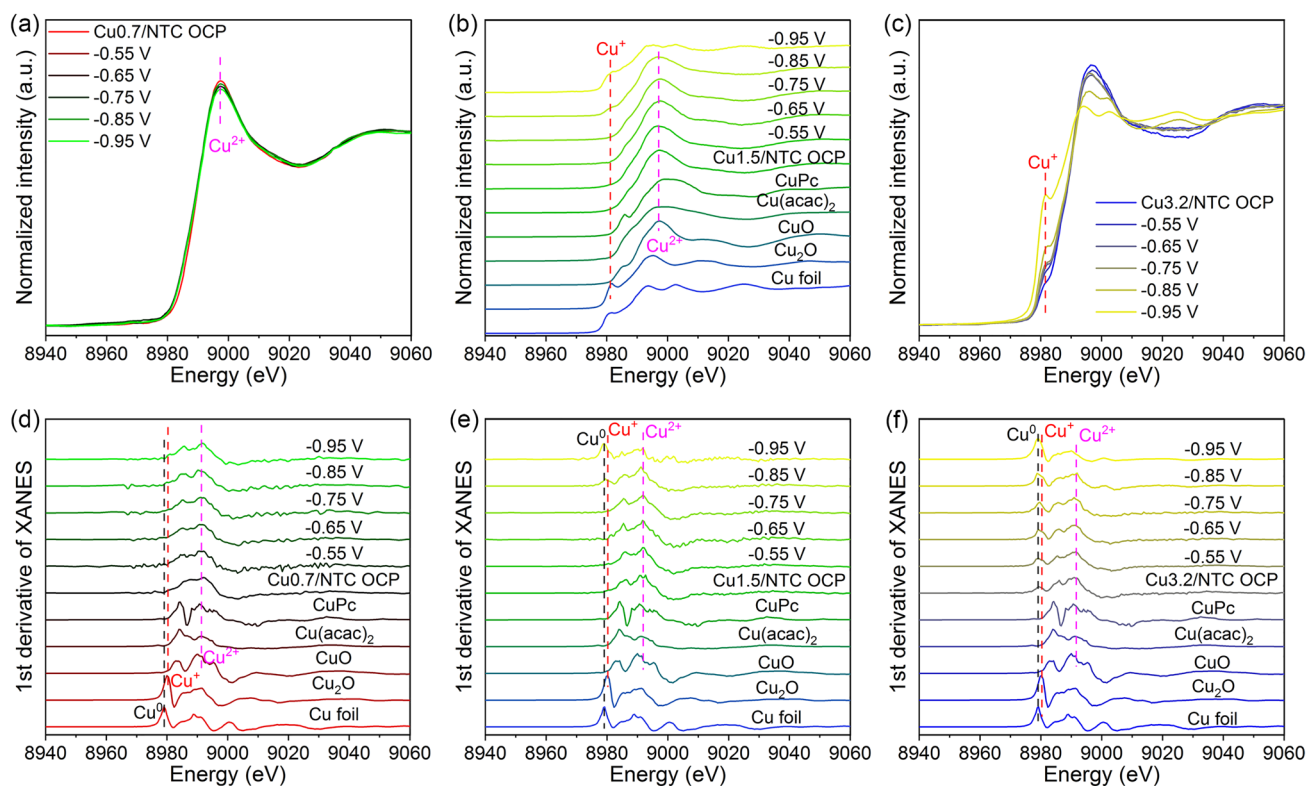


Fig. 5 a–c Operando XANES spectra of Cu_{0.7}/NTC, Cu_{1.5}/NTC, and Cu_{3.2}/NTC. d–f 1st derivative of XANES spectra of Cu_{0.7}/NTC, Cu_{1.5}/NTC and Cu_{3.2}/NTC

to form Cu₂ clusters with Cu–N₄ (CuN₄&Cu₂), and then further restructuring to form CuN₄&Cu₆ at –0.95 V. The weak Cu–Cu bond (k : 8.9 Å⁻¹) and apparent Cu–Cu bonds (k : 9.9 Å⁻¹) could be detected in the WT-EXAFS spectra of –0.85 and –0.95 V for Cu_{0.7}/NTC (Fig. S26), except the findable Cu–N bonds at the potentials from OCP to –0.75 V. Discriminatively, the CN of Cu–N bonds at OCP, –0.75, –0.85, –0.95 V for Cu_{3.2}/NTC are 3.58, 2.89, 1.55, 0.41 in succession, showing that the primary configurations of Cu species at OCP and –0.75 V are Cu²⁺–N₄, while Cu⁰ nanoparticles (NPs) exist at –0.85 and –0.95 V (Figs. S27–S29, Table S7). The restructuring behavior in Cu_{3.2}/NTC system can be jointly dependent on the Cu loading amount and reaction potential since Cu⁰ content of Cu_{3.2}/NTC grows substantially more than that of Cu_{0.7}/NTC at the same potentials.

Given the k^3 -weighted FT-EXAFS fitting spectra integrated with the XANES LCF results of Cu_{1.5}/NTC (Figs. 6a–d, S30–S32, and Table S8), the CN of Cu–N bonds at OCP, –0.75, –0.85, –0.95 V are 3.90, 3.87, 3.58, 0.59, and the CN of Cu–Cu bonds are 2.90, 4.12, 7.29, and

8.85, so it is reasonable to assume that Cu_{1.5}/NTC had CuN₄&Cu₃, CuN₄&Cu₄, CuN₄&Cu₇, and Cu₉ structures at the aforementioned potentials (inset of Fig. 6a–d). Intrigued by the stability of restructuring Cu clusters, *ex-situ* XANES, 1st derivative of XANES, R space, FT-EXAFS fitting spectra, WT-EXAFS, and XANES LCF spectra detected 5 and 48 h after –0.95 V in the air (Figs. 6e–g, S30–S33) clarify the transformation of valence states from metallic Cu to the oxidation state of Cu, accompanied by the incompletely reversible transformation from Cu₉ clusters to CuN₄&Cu₃ (initial OCP). Moreover, the bond length in the first shell of exposure in the air for 5 h after –0.95 V (1.96 Å) is marginally bigger than that of exposure in the air for 48 h (1.94 Å), which can result from the bond between Cu and light atoms (N/O) during the oxidation process of metallic Cu, following by the formation of [Cu(OH)₄]²⁻ intermediate on the wet carbon paper of an alkaline solution. Ultimately, intermediates could be anchored again on the –N₄C_x sites among the supported materials due to the chelating capacity of Cu²⁺ [24, 35].

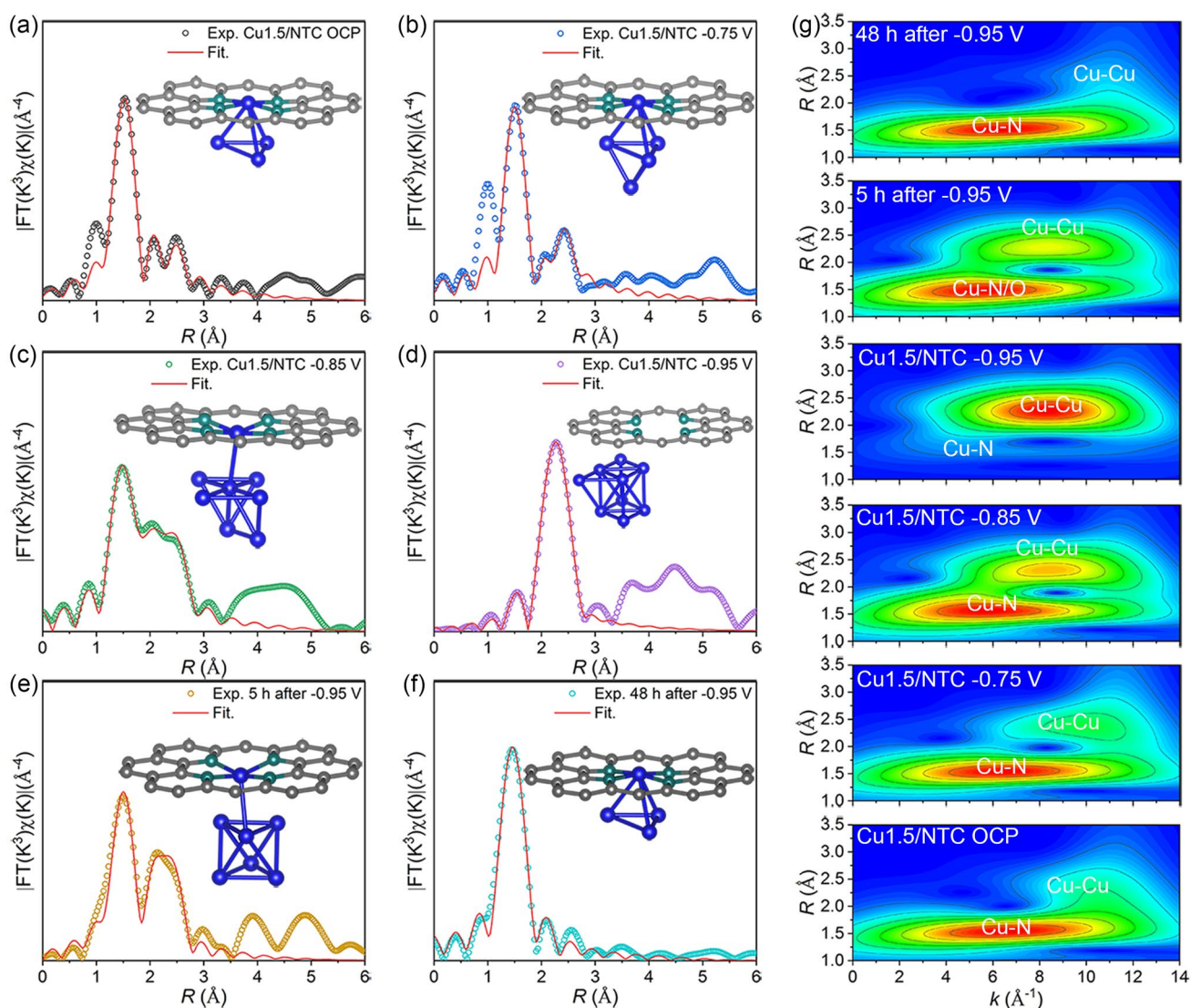


Fig. 6 **a–f** The fitting spectra of R space for Cu_{1.5}/NTC at OCP, –0.75, –0.85, –0.95 V, 5 and 48 h after –0.75 V in air, and the corresponding structures in the inset. **g** The WT-EXAFS spectra. Gray, green and blue balls, respectively, represent C, N, and Cu atoms

The relationships among the CN, Cu loading, and reaction potential were further investigated using the nonlinear fitting curves of the typical reaction potential versus the CN (Cu–N and Cu–Cu bonds) toward Cu_{0.7}/NTC, Cu_{1.5}/NTC, and Cu_{3.2}/NTC electrocatalysts (Fig. 7a, b). Potential intervals of the dominant Cu–N₄ configuration in Cu_{0.7}/NTC, Cu_{1.5}/NTC, and Cu_{3.2}/NTC electrocatalysts are OCP to –0.95 V, OCP to –0.85 V, OCP to –0.75 V, while the CN maximum increments of Cu–Cu bonds occur at –0.95, –0.85, –0.75 V,

indicating that the potentials of electrochemical restructuring are shifted positively with the increased loading of Cu. Consequently, the restructuring behavior among Cu species in this system co-depends on the Cu loading and reaction potential. Furthermore, the possible mechanisms of aggregation driven by potential and redispersion driven by oxidation integrated with chelation for Cu_{1.5}/NTC are described in Fig. 7c and further verified with the EDX mapping of Cu_{1.5}/NTC exposed in the air after –0.95 V (Fig. S34).

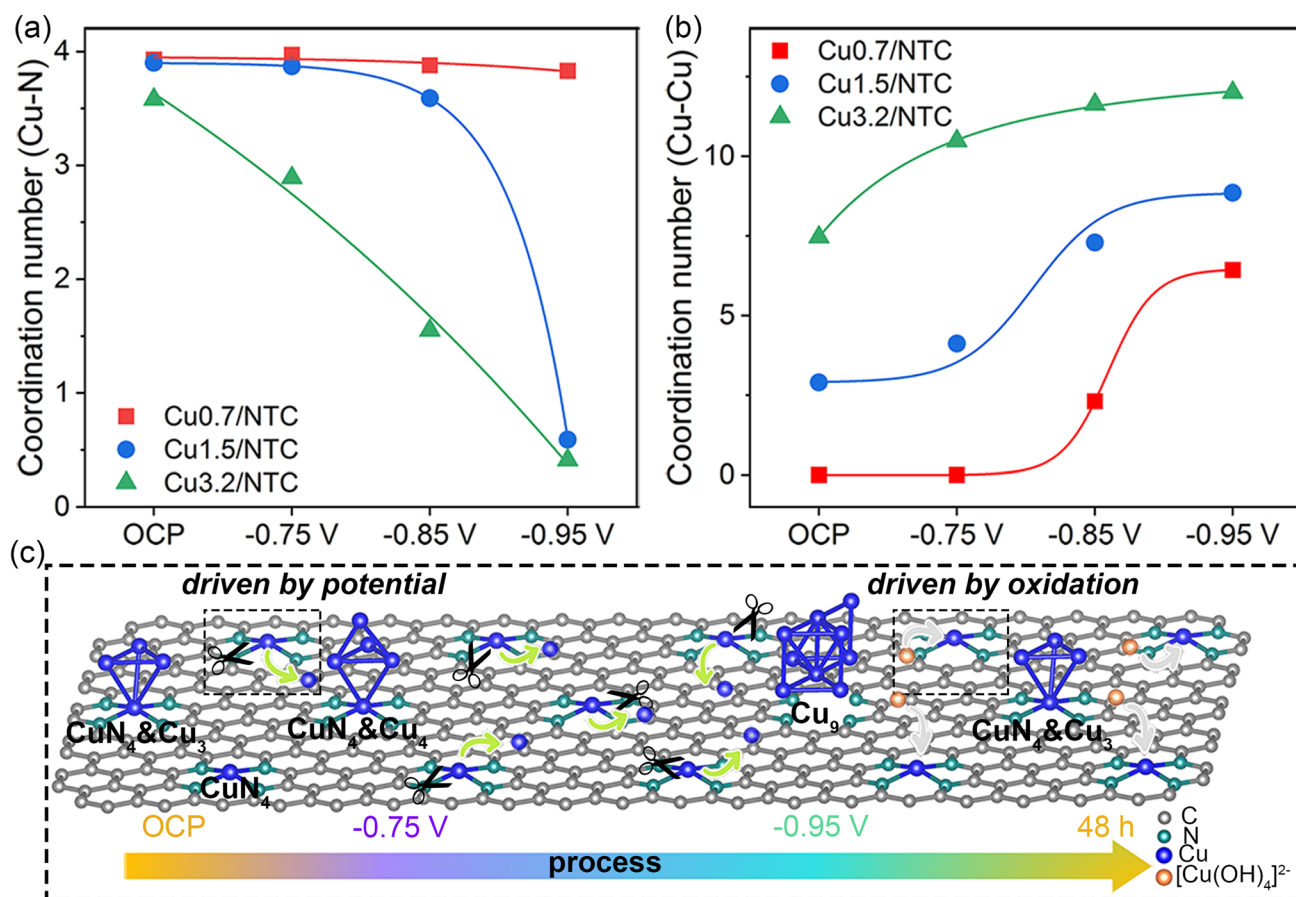


Fig. 7 a, b Nonlinear fitting curves of the typical reaction potential versus the CN (Cu–N and Cu–Cu bonds) among Cu0.7/NTC, Cu1.5/NTC, and Cu3.2/NTC electrocatalysts. c The possible electrochemically driven remodeling and oxidation-driven redispersion mechanisms for Cu1.5/NTC

3.6 Electrocatalytic NITRR Mechanisms

To further evaluate the origin of the NITRR catalytic activity, density functional theory (DFT) calculations were executed with the multi-component models (Fig. S35) of Cu species, including Cu single-atom (CuN₄), Cu single-atom and clusters (CuN₄&Cu₄), Cu single-atom and nanoparticles with (111) facet (CuN₄&Cu NPs), based on the fitting structures of FT-EXAFS and the variation trend of mass activity (Fig. S36) [38, 39]. The *d*-band center of projected density of states (PDOS) for CuN₄&Cu₄ model ($d_{\text{center}} = -2.56$ eV) is closest to the Fermi level than that of CuN₄ and CuN₄&Cu NPs (Figs. 8a and S37), signifying that the strong interaction between CuN₄ and Cu clusters upshifted the *d*-band center to improve the adsorption ability of reactants and intermediates [40, 41]. Charge density difference (EDD) distributions of CuN₄, CuN₄&Cu₄, and CuN₄&Cu NPs were calculated

(Figs. 8b and S38). As a result, introducing the Cu cluster to CuN₄ (CuN₄&Cu₄) can ameliorate the charge distribution of the Cu single-site and modulate the electric structure with high activity for NITRR compared with CuN₄ and CuN₄&Cu NPs, crucial for invigorating the sluggish NITRR kinetics [42]. Online differential electrochemical mass spectrometry (DEMS) offers robust technical support for capturing volatile intermediates and products [43]. Additionally, mass signals were acquired during the chronoamperometry test at -0.75 V for five cycles (Fig. 8c), indicating mass signals of $m/z = 17, 31, 33,$ and 46 are assigned to the intermediate products or the molecular fragments of NH₃, NOH, NH₂OH, and NO₂. According to the results of online DEMS, and integrated with the deoxygenation process of NO₃⁻ ($*\text{NO}_3 \rightarrow *\text{NO}_2 \rightarrow *\text{NO}$) and hydrogenation process of $*\text{NO}$ intermediate [44], we calculated the free energy pathway of NH₃ formation for each step in detail upon the constructed models with the most stable

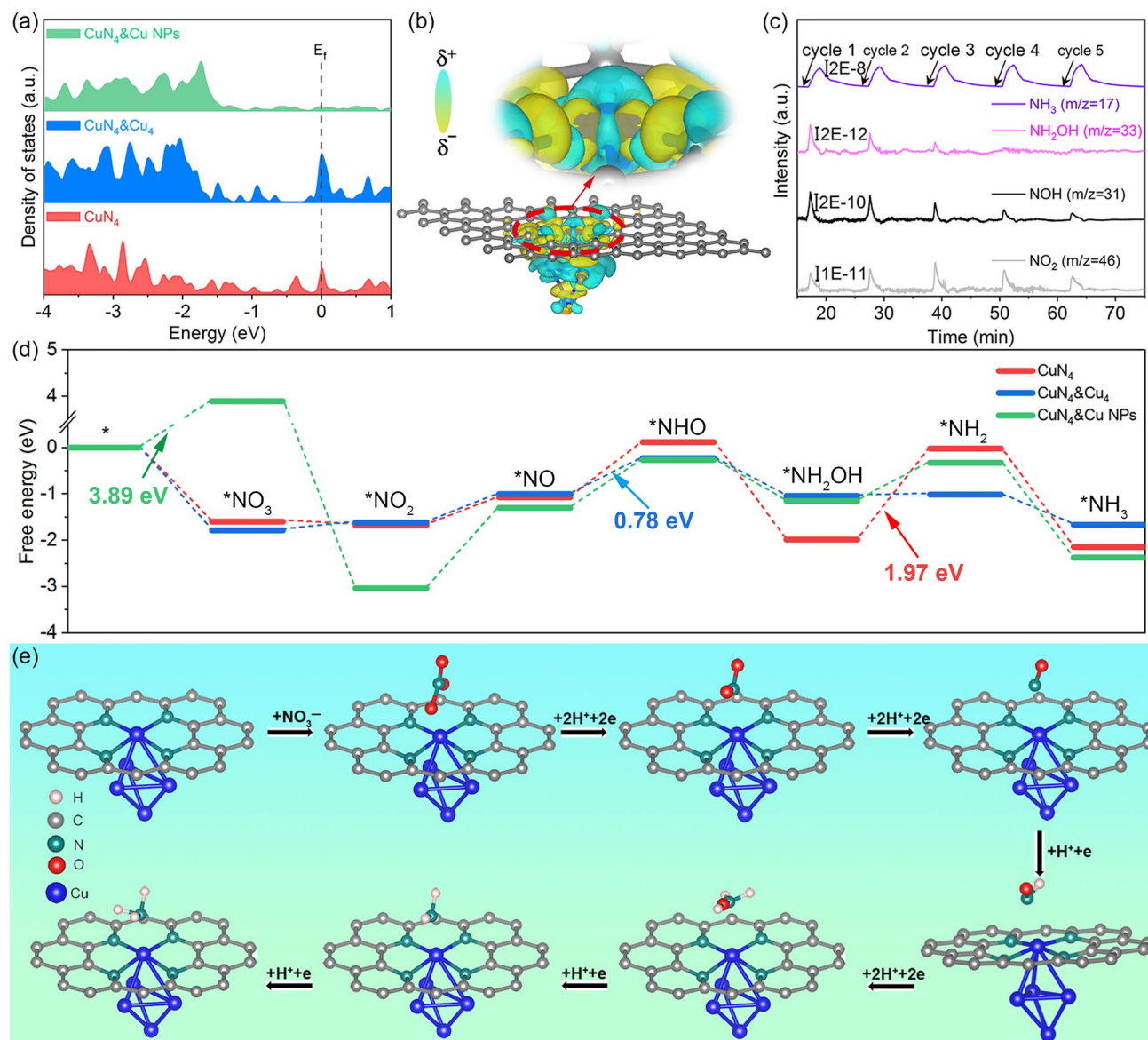


Fig. 8 **a** PDOS CuN₄, CuN₄&Cu₄, and CuN₄&Cu NPs. **b** EDD distribution of CuN₄&Cu₄ model. The yellow and cyan regions represent electron accumulation and depletion. **c** Online DEMS measured at -0.75 V for NITRR over Cu_{1.5}/NTC. **d** The free energy paths of NH₃ production for CuN₄, CuN₄&Cu₄ and CuN₄&Cu NPs. **e** The minimum energy pathway of NH₃ production for CuN₄&Cu₄ configuration. Color code: H (light pink), C (gray), N (green), O (red), Cu (blue). (Color figure online)

adsorption configurations, especially for the first step of hydrogenation site of *NO intermediate. Notably, the first hydrogenation site of the *NO intermediate for CuN₄, CuN₄&Cu₄, and CuN₄&Cu NPs preferentially forms *NHO rather than *ONH (Fig. S39), which coincides with the DEMS results with the by-products (NOH and NH₂OH). At the initiation step of the possible NITRR pathway on CuN₄, CuN₄&Cu₄, and CuN₄&Cu NPs utilized Perdew–Burke–Ernzerhof (PBE)

functional, the adsorption of reactant NO₃⁻ is a spontaneous behavior toward CuN₄ and CuN₄&Cu₄ (more favorable), while the NO₃⁻ adsorption on CuN₄&Cu NPs with the free energy changes (ΔG) of 3.89 eV is thermodynamically unfavorable (Figs. 8d, e, and S40–S41. Tables S9 and S10) [41, 45]. Besides, the potential-determining steps (PDS) of CuN₄ model is *NH₂OH to *NH₂ ($\Delta G = 1.97$ eV), which is apparently higher than CuN₄&Cu₄ model ($\Delta G = 0.03$ eV) despite the

PDS of CuN₄&Cu₄ model is *NH to *NOH ($\Delta G=0.78$ eV), indicating Cu cluster can optimize reaction energy barrier of *NH₂OH to *NH₂. Meanwhile, the results of solvent model and the local density approximation (LDA) functional were adopted in NITRR calculations, which gave similar tendency in CuN₄&Cu₄ configuration to PBE data (Figs. S43-S44). In a nutshell, the synergistic coupling between the CuN₄ and Cu clusters modulates the charge distribution and the electric structure with the suitable *d*-band center, enhancing the adsorption ability of NO₃⁻ and accelerating the rapid conversion from *NH₂OH to *NH₂ simultaneously.

4 Conclusions

In conclusion, we have exhaustively investigated the restructuring behavior in the electrocatalytic NITRR process by employing Cu species with tunable loading supported on N-doped TiO₂/C at potentials ranging from OCP to -0.95 V versus RHE based on the *operando* XAS. Electrochemical restructuring is the reduction process of Cu with a positive valence state to metallic Cu, and then Cu⁰ can migrate and aggregate to form Cu clusters or nanoparticles. The restructuring behavior of Cu species is jointly dependent on Cu loading and reaction potential, i.e., the higher Cu loading and the more negative potential, the easier restructuring of Cu species to form Cu clusters or nanoparticles with higher nucleation numbers. *Operando* XAS and electrocatalytic NITRR performances perform that Cu1.5/NTC with CuN₄&Cu₄ configuration could deliver the highest NH₃ yield with 88.2 mmol h⁻¹ g_{cata}⁻¹ (0.044 mmol h⁻¹ cm⁻²) and FE (~94.3%) at -0.75 V, which is superior to NTC, Cu0.7/NTC (CuN₄ configuration), and Cu3.2/NTC (CuN₄&Cu₁₀ configuration). DFT calculation results reveal that Cu clusters with appropriate nuclear numbers modulate the charge distribution and the *d*-band center of Cu single-site, thus promoting the adsorption ability of NO₃⁻ and accelerating the conversion of the key intermediates (*NH₂OH → *NH₂). This work sheds light on the restructuring behavior trigger, allowing for more informed decisions in the rational design of highly active NITRR electrocatalysts.

Acknowledgments The authors acknowledge the 4B9A station of the Beijing Synchrotron Radiation Facility (BSRF) and the Analytical and Testing Center of Beijing Institute of Technology for technical supports. This work was supported by the National Natural Science Foundation of China (Grant numbers 92061106 and 21971016).

Funding Open access funding provided by Shanghai Jiao Tong University.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit <http://creativecommons.org/licenses/by/4.0/>.

Supplementary Information The online version contains supplementary material available at <https://doi.org/10.1007/s40820-023-01091-9>.

References

1. C.X. Guo, J.R. Ran, A. Vasileff, S.Z. Qiao, Rational design of electrocatalysts and photo(electro)catalysts for nitrogen reduction to ammonia (NH₃) under ambient conditions. *Energy Environ. Sci.* **11**(1), 45–56 (2018). <https://doi.org/10.1039/C7EE022-20D>
2. W.H. Guo, K.X. Zhang, Z.B. Liang, R.Q. Zou, Q. Xu, Electrochemical nitrogen fixation and utilization: theories, advanced catalyst materials and system design. *Chem. Soc. Rev.* **48**(24), 5658–5716 (2019). <https://doi.org/10.1039/C9CS00159J>
3. W.C. Zhang, B.W. Zhang, Bi-atom electrocatalyst for electrochemical nitrogen reduction reactions. *Nano Micro Lett.* **13**, 106 (2021). <https://doi.org/10.1007/s40820-021-00638-y>
4. Y.P. Pang, C. Su, G.H. Jia, L.Q. Xu, Z.P. Shao, Emerging two-dimensional nanomaterials for electrochemical nitrogen reduction. *Chem. Soc. Rev.* **50**(22), 12744–12787 (2021). <https://doi.org/10.1039/D1CS00120E>
5. A. Biswas, S. Kapse, R. Thapa, R.S. Dey, Oxygen functionalization-induced charging effect on boron active sites for high-yield electrocatalytic NH₃ production. *Nano Micro Lett.* **14**, 214 (2022). <https://doi.org/10.1007/s40820-022-00966-7>
6. X.W. Wang, D. Wu, S.Y. Liu, J.J. Zhang, X.Z. Fu et al., Folic acid self-assembly enabling manganese single-atom electrocatalyst for selective nitrogen reduction to ammonia. *Nano Micro Lett.* **13**, 125 (2021). <https://doi.org/10.1007/s40820-021-00651-1>
7. Y.T. Wang, C.H. Wang, M.Y. Li, Y.F. Yu, B. Zhang, Nitrate electroreduction: mechanism insight, in situ characterization, performance evaluation, and challenges. *Chem. Soc. Rev.* **50**(12), 6720–6733 (2021). <https://doi.org/10.1039/D1CS00116G>



8. H. Xu, Y.Y. Ma, J. Chen, W.X. Zhang, J.P. Yang, Electrocatalytic reduction of nitrate—a step towards a sustainable nitrogen cycle. *Chem. Soc. Rev.* **51**(7), 2710–2758 (2022). <https://doi.org/10.1039/D1CS00857A>
9. R. Zhang, Y. Guo, S.C. Zhang, D. Chen, Y.W. Zhao et al., Efficient ammonia electrosynthesis and energy conversion through a Zn-nitrate battery by iron doping engineered nickel phosphide catalyst. *Adv. Energy Mater.* **12**(13), 2103872 (2022). <https://doi.org/10.1002/aenm.202103872>
10. G.F. Chen, Y.F. Yuan, H.F. Jiang, S.Y. Ren, L.X. Ding et al., Electrochemical reduction of nitrate to ammonia via direct eight-electron transfer using a copper–molecular solid catalyst. *Nat. Energy* **5**(8), 605–613 (2020). <https://doi.org/10.1038/s41560-020-0654-1>
11. D.M. Zhao, Y.Q. Wang, C.L. Dong, F.Q. Meng, Y.C. Huang et al., Electron-deficient Zn-N₆ configuration enabling polymeric carbon nitride for visible-light photocatalytic overall water splitting. *Nano Micro Lett.* **14**, 223 (2022). <https://doi.org/10.1007/s40820-022-00962-x>
12. L.L. Han, H. Cheng, W. Liu, H.Q. Li, P.F. Ou et al., A single-atom library for guided monometallic and concentration-complex multimetallic designs. *Nat. Mater.* **21**(6), 681–688 (2022). <https://doi.org/10.1038/s41563-022-01252-y>
13. X. Wu, H.B. Zhang, S.W. Zuo, J.C. Dong, Y. Li et al., Engineering the coordination sphere of isolated active sites to explore the intrinsic activity in single-atom catalysts. *Nano Micro Lett.* **13**, 136 (2021). <https://doi.org/10.1007/s40820-021-00668-6>
14. X.B. Zheng, P. Li, S.X. Dou, W.P. Sun, H.G. Pan et al., Non-carbon-supported single-atom site catalysts for electrocatalysis. *Energy Environ. Sci.* **14**(5), 2809–2858 (2021). <https://doi.org/10.1039/D1EE00248A>
15. W.X. Chen, J.J. Pei, C.T. He, J.W. Wan, H.L. Ren et al., Single tungsten atoms supported on MOF-derived N-doped carbon for robust electrochemical hydrogen evolution. *Adv. Mater.* **30**(30), 180039 (2018). <https://doi.org/10.1002/adma.201800396>
16. B.L. Yan, D.P. Liu, X.L. Feng, M.Z. Shao, Y. Zhang, Ru species supported on MOF-derived N-doped TiO₂/C hybrids as efficient electrocatalytic/photocatalytic hydrogen evolution reaction catalysts. *Adv. Funct. Mater.* **30**(31), 2003 (2020). <https://doi.org/10.1002/adfm.202003007>
17. X.Y. Xie, L.S. Peng, H.Z. Yang, G.I.N. Waterhouse, L. Shang et al., MIL-101-derived mesoporous carbon supporting highly exposed Fe single-atom sites as efficient oxygen reduction reaction catalysts. *Adv. Mater.* **33**(23), 21010 (2021). <https://doi.org/10.1002/adma.202101038>
18. J. Choi, P. Wagner, S. Gambhir, R. Jalili, D.R. MacFarlane et al., Steric modification of a cobalt phthalocyanine/graphene catalyst to give enhanced and stable electrochemical CO₂ reduction to CO. *ACS Energy Lett.* **4**(3), 666–672 (2019). <https://doi.org/10.1021/acseenergylett.8b02355>
19. C. Xia, Y.R. Qiu, Y. Xia, P. Zhu, G. King et al., General synthesis of single-atom catalysts with high metal loading using graphene quantum dots. *Nat. Chem.* **13**(9), 887–894 (2021). <https://doi.org/10.1038/s41557-021-00734-x>
20. W.J. Zhai, S.H. Huang, C.B. Lu, X.N. Tang, L.B. Li et al., Simultaneously integrate iron single atom and nanocluster triggered tandem effect for boosting oxygen electroreduction. *Small* **18**(15), 2107225 (2022). <https://doi.org/10.1002/sml.202107225>
21. Z.M. Ma, S.Q. Liu, N.F. Tang, T. Song, K. Motokura et al., Coexistence of Fe nanoclusters boosting Fe single atoms to generate singlet oxygen for efficient aerobic oxidation of primary amines to imines. *ACS Catal.* **12**(9), 5595–5604 (2022). <https://doi.org/10.1021/acscatal.1c04467>
22. G.Y. Xing, M.M. Tong, P. Yu, L. Wang, G.Y. Zhang et al., Restructuration of highly dense Cu–N₄ active sites in electrocatalytic oxygen reduction characterized by operando synchrotron radiation. *Angew. Chem. Int. Ed.* **61**(40), e202211 (2022). <https://doi.org/10.1002/anie.202211098>
23. H.P. Xu, D. Rebolgar, H.Y. He, L.N. Chong, Y.Z. Liu et al., Highly selective electrocatalytic CO₂ reduction to ethanol by metallic clusters dynamically formed from atomically dispersed copper. *Nat. Energy* **5**(8), 623–632 (2020). <https://doi.org/10.1038/s41560-020-0666-x>
24. J. Yang, H.F. Qi, A.Q. Li, X.Y. Liu, X.F. Yang et al., Potential-driven restructuring of Cu single atoms to nanoparticles for boosting the electrochemical reduction of nitrate to ammonia. *J. Am. Chem. Soc.* **144**(27), 12062–12071 (2022). <https://doi.org/10.1021/jacs.2c02262>
25. Y.T. Xu, M.Y. Xie, H.Q. Zhong, Y. Cao, In situ clustering of single-atom copper precatalysts in a metal-organic framework for efficient electrocatalytic nitrate-to-ammonia reduction. *ACS Catal.* **12**(14), 8698–8706 (2022). <https://doi.org/10.1021/acscatal.2c02033>
26. X.Y. Ji, Y.Y. Wang, Y. Li, K. Sun, M. Yu et al., Enhancing photocatalytic hydrogen peroxide production of Ti-based metal–organic frameworks: the leading role of facet engineering. *Nano Res.* **15**(7), 6045–6053 (2022). <https://doi.org/10.1007/s12274-022-4301-y>
27. J. Janczak, Y.M. Idemori, Synthesis, crystal structure and characterisation of aquamagnesium phthalocyanine—MgPc(H₂O). The origin of an intense near-IR absorption of magnesium phthalocyanine known as ‘X-phase.’ *Polyhedron* **22**(9), 1167–1181 (2003). [https://doi.org/10.1016/S0277-5387\(03\)00110-4](https://doi.org/10.1016/S0277-5387(03)00110-4)
28. J. Gu, C.S. Hsu, L.C. Bai, H.M. Chen, X.L. Hu, Atomically dispersed Fe³⁺ sites catalyze efficient CO₂ electroreduction to CO. *Science* **364**(6445), 1091–1094 (2019). <https://doi.org/10.1126/science.aaw7515>
29. H.D. Li, Y. Pan, Z.C. Wang, Y.D. Yu, J. Xiong et al., Coordination engineering of cobalt phthalocyanine by functionalized carbon nanotube for efficient and highly stable carbon dioxide reduction at high current density. *Nano Res.* **15**(4), 3056–3064 (2022). <https://doi.org/10.1007/s12274-021-3962-2>
30. Q.W. Chang, Y.M. Liu, J.H. Lee, D. Ologunagba, S. Hwang et al., Metal-coordinated phthalocyanines as platform molecules for understanding isolated metal sites in the electrochemical reduction of CO₂. *J. Am. Chem. Soc.* **144**(35), 16131–16138 (2022). <https://doi.org/10.1021/jacs.2c06953>
31. M.Y. Li, T.H. Wang, W.X. Zhao, S.Y. Wang, Y.Q. Zou, A pair-electrosynthesis for formate at ultra-low voltage via

- coupling of CO₂ reduction and formaldehyde oxidation. *Nano Micro Lett.* **14**, 211 (2022). <https://doi.org/10.1007/s40820-022-00953-y>
32. W. Zhou, L. Fu, L. Zhao, X.J. Xu, W.Y. Li et al., Novel core–sheath Cu/Cu₂O–ZnO–Fe₃O₄ nanocomposites with high-efficiency chlorine-resistant bacteria sterilization and trichloroacetic acid degradation performance. *ACS Appl. Mater. Interfaces* **13**(9), 10878–10890 (2021). <https://doi.org/10.1021/acsami.0c21336>
33. X. Zhang, C.H. Wang, Y.M. Guo, B. Zhang, Y.T. Wang et al., Cu clusters/TiO_{2-x} with abundant oxygen vacancies for enhanced electrocatalytic nitrate reduction to ammonia. *J. Mater. Chem. A* **10**(12), 6448–6453 (2022). <https://doi.org/10.1039/D2TA00661H>
34. H. Su, W.L. Zhou, H. Zhang, W. Zhou, X. Zhao et al., Dynamic evolution of solid–liquid electrochemical interfaces over single-atom active sites. *J. Am. Chem. Soc.* **142**(28), 12306–12313 (2020). <https://doi.org/10.1021/jacs.0c04231>
35. D. Karapinar, N.T. Huan, N.R. Sahraie, J.K. Li, D. Wakerley et al., Electroreduction of CO₂ on single-site copper-nitrogen-doped carbon material: selective formation of ethanol and reversible restructuring of the metal sites. *Angew. Chem. Int. Ed.* **58**(42), 15098 (2019). <https://doi.org/10.1002/anie.201907994>
36. X.Z. Su, Z.L. Jiang, J. Zhou, H.J. Liu, D.N. Zhou et al., Complementary *Operando* Spectroscopy identification of in-situ generated metastable charge-asymmetry Cu₂–CuN₃ clusters for CO₂ reduction to ethanol. *Nat. Commun.* **13**, 1322 (2022). <https://doi.org/10.1038/s41467-022-29035-8>
37. J. Yang, W.G. Liu, M.Q. Xu, X.Y. Liu, H.F. Qi et al., Dynamic behavior of single-atom catalysts in electrocatalysis: identification of Cu–N₃ as an active site for the oxygen reduction reaction. *J. Am. Chem. Soc.* **143**(36), 14530–14539 (2021). <https://doi.org/10.1021/jacs.1c03788>
38. X. Ao, W. Zhang, Z.S. Li, J.G. Li, L. Soule et al., Markedly enhanced oxygen reduction activity of single-atom Fe catalysts via integration with Fe nanoclusters. *ACS Nano* **13**(10), 11853–11862 (2019). <https://doi.org/10.1021/acs.nano.9b05913>
39. X. Wan, Q.T. Liu, J.Y. Liu, S.Y. Liu, X.F. Liu et al., Iron atom–cluster interactions increase activity and improve durability in Fe–N–C fuel cells. *Nat. Commun.* **13**, 2963 (2022). <https://doi.org/10.1038/s41467-022-30702-z>
40. X.H. Liu, L.R. Zheng, C.X. Han, H.X. Zong, G. Yang et al., Identifying the activity origin of a cobalt single-atom catalyst for hydrogen evolution using supervised learning. *Adv. Funct. Mater.* **31**(18), 2100547 (2021). <https://doi.org/10.1002/adfm.202100547>
41. Q. Hu, Y.J. Qin, X.D. Wang, Z.Y. Wang, X.W. Huang et al., Reaction intermediate-mediated electrocatalyst synthesis favors specified facet and defect exposure for efficient nitrate–ammonia conversion. *Energy Environ. Sci.* **14**(9), 4989–4997 (2021). <https://doi.org/10.1039/D1EE01731D>
42. A. Kumar, J. Lee, M.G. Kim, B. Debnath, X.H. Liu et al., Efficient nitrate conversion to ammonia on f-block single-atom/metal oxide heterostructure via local electron-deficiency modulation. *ACS Nano* **16**(9), 15297–15309 (2022). <https://doi.org/10.1021/acs.nano.2c06747>
43. Y.T. Wang, H.J. Li, W. Zhou, X. Zhang, B. Zhang et al., Structurally disordered RuO₂ nanosheets with rich oxygen vacancies for enhanced nitrate electroreduction to ammonia. *Angew. Chem. Int. Ed.* **61**(19), e202202604 (2022). <https://doi.org/10.1002/anie.202202604>
44. J. Li, G.M. Zhan, J.H. Yang, F.J. Quan, C.L. Mao et al., Efficient ammonia electrosynthesis from nitrate on strained ruthenium nanoclusters. *J. Am. Chem. Soc.* **142**(15), 7036–7046 (2020). <https://doi.org/10.1021/jacs.0c00418>
45. H.B. Yin, Z. Chen, S.C. Xiong, J.J. Chen, C.Z. Wang et al., Alloying effect-induced electron polarization drives nitrate electroreduction to ammonia. *Chem. Catal.* **1**(5), 1088–1103 (2021). <https://doi.org/10.1016/j.checat.2021.08.014>