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Highly selective electrocatalytic reduction of nitrate to ammonia over a copper–cobalt bimetallic catalyst[†]

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The electrocatalytic nitrate reduction reaction (NitRR) is a promising alternative to the traditional Haber– Bosch process. However, the competitive hydrogen evolution reaction results in poor NH₃ selectivity (S_{NH3}). Here, a Cu–Co bimetallic catalyst supported on biomass-derived porous carbon (Cu–Co/BPC) is designed and synthesized. Interestingly, the catalyst presents a high NH₃ yield rate of 9114.1 ± 244.8 µg h⁻¹ cm⁻² at -1.4 V (vs. RHE) and a high faradaic efficiency (FE) of 84.5 ± 1.6% at -1.0 V (vs. RHE). Notably, the S_{NH_3} of Cu–Co/BPC catalyst is kept above 94.2% under a broad range from -1.0 to -1.4 V (vs. RHE), indicating the high NitRR-to-NH₃ selectivity of Cu–Co/BPC. The combination of *in situ* characterization and experimental results indicates that the electron transfer occurs between Cu and Co, and many active sites are generated for adsorption and activation of N=O double bonds, and hydrogenation reactions occur with adjacent H protons to improve the selectivity of NH₃.

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Ammonia (NH₃) is a chemical product widely used in industries such as fuel, pharmaceuticals, and fertilizer manufacturing.1-3 Currently, the Haber-Bosch method is the main method used in industry to produce NH3.4,5 Different from the Haber-Bosch technology which requires high temperatures (400-500 °C) and pressures (200-300 atm),6-8 the mild approaches for nitrogen (N_2) fixation such as homogeneous catalysis with organic or reductive agents,9 electrocatalysis,10 nitrogenase catalysis,11 and photocatalysis¹² are promising alternatives. Amongst these, the electrochemical nitrogen reduction reaction (NRR) has gained much attention, since it takes water and air as the synthetic source and can be powered by renewable electricity.13 However, NRR is a representative gas-liquid-solid reaction system in aqueous electrolyte systems, in which the system exhibits poor N_2 solubility, high N \equiv N bonding energy, and sluggish kinetic process resulting in low NH₃ yields and Faraday efficiencies (FE), leading to the limited activity and selectivity in NRR.13-15 To overcome the bottleneck, the electrochemical nitrate reduction reaction (NitRR) is widely recognized for its advantages of mild and environmentally friendly reaction conditions.16 The dissociation energy of the N=O bond (204 kJ mol⁻¹) is lower than that of the N≡N bond (941 kJ mol⁻¹).¹⁷ The solubility of nitrate is also much greater than that of N2.18,19 These two advantages emphasize the research value of NitRR. In addition,

electrochemical NitRR from various industrial wastewater and municipal sewage discharges can efficiently remove polluted nitrate and obtain high-value-added chemical products, affording a popular approach for nitrate-involving wastewater treatment and clean energy regeneration.^{20–22}

Transition metals (TM) have been widely considered in the design of various catalysts owing to their wide distribution, lower cost, and unsaturated d-orbitals, which are capable of inducing a rearrangement between electrons occurring between the adsorbed molecules and the catalysts, leading to orbitals hybridization with the reactants and following activation.23-25 Until now, most of the catalysts that were reported for NitRR are popular by using transition metal as the dominant catalytic center.26,27 In particular, Cu or Co based catalysts are promising and widely implemented in NitRR.26 For example, Zhu et al. reported a nanoporous Cu/CoOOH catalyst that exhibited enhanced *NO2 affinity and reduces the energy barrier in the rate-determining *NO2H formation step, effectively facilitating NO₃⁻ reduction to NH₃.²⁶ Song and co-workers demonstrated that the bimetallic cMOF (Cu_xCo_yHHTP) is proposed for efficient NitRR and the synergy effect between different types of single-metal sites is also revealed. Both theoretical and experimental results reveal that the Co sites can affect the electron structure of Cu sites in Cu₁Co₁HHTP slab and decrease the ΔG

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of potential determining step in NitRR process. Moreover, the Co sites bring a higher selectivity to Cu active sites for reducing $*NO_2$ to *NO, rather than the desorption of $NO_2^{-2.28}$ Additionally, theoretical calculation reveals that the surface oxygen on cobalt sites can stabilize the adsorbed hydrogen on cobalt oxide, which hampers the evolution of hydrogen and leads to an enhanced NitRR activity.²⁷ These findings hold significant implications for electrocatalysis research focused on studying TMs.

Herein, regarding the advantage of Co species in the adsorption of NO₃⁻ and the formation of active H intermediate, we construct the Cu-Co bimetallic nanoparticles supported on biomass-derived porous carbon (Cu-Co/BPC) as an effective electrocatalyst for a high-efficient NH₃ production via NO₃⁻ electro-reduction at ambient conditions. As a result, the assynthesized Cu–Co/BPC affords an excellent $R_{\rm NH_2}$ of 9114.1 \pm 244.8 μ g h⁻¹ cm⁻² at -1.4 V (vs. RHE) and a maximum FE of $84.5 \pm 1.6\%$ at -1.0 V (vs. RHE), with persistent cycle stability of 16 consecutive recycling tests in 0.1 M K₂SO₄ + 0.1 M KNO₃ electrolyte at -1.25 V (vs. RHE). Furthermore, the differential electrochemical mass spectrometry (DEMS) was adopted to monitor the electrochemical separation of the NitRR products on the surface of the Cu-Co/BPC. Accordingly, the possible reaction sites and electrocatalytic NitRR mechanism of Cu-Co/ BPC have been proposed.

Fig. 1a shows the synthetic illustration of the Cu–Co/BPC catalyst supported on coconut powder-derived biomass carbon. X-ray diffraction (XRD) was carried out to confirm the crystal structure of as-synthesized Cu–Co/BPC. As evidenced in Fig. 1b, the broad diffraction peak observed at 26.4° correspond to the (002) plane of graphitic carbon (PDF no. 41-1487),

indicating the presence of graphitic carbon substrate in the catalyst. Additionally, the main diffraction peaks at 32.5°, 35.5°, 38.9° and 48.7° correlate well with the (110), (11-1), (200) and (20-2), planes of copper oxide (PDF no. 48-1548). Furthermore, peaks at 36.5° , 42.4° and 61.5° , correspond to the (111), (200) and (220) planes of CoO (PDF no. 48-1719). The above results indicate that Cu and Co elements in the Cu-Co/BPC catalyst mainly exist in the form of oxidation state. The microstructure and elemental distribution of the as-prepared Cu-Co/BPC catalyst were also investigated. The transmission electron microscopy (TEM) image suggests that the Cu-Co/BPC is composed of nanoparticles with a size of \sim 20 nm (Fig. 1c). From the high-resolution transmission electron microscopy (HR-TEM) image of Cu-Co/BPC in Fig. 1c, the crystal lattice distance of d = 0.21 nm corresponds to the (111) crystal plane of CuO as recorded. Also, the high-angle annular dark-field scanning transmission electron microscopy (HAADF-STEM) along with elemental mapping images confirmed the uniform distribution of Cu, Co, O and C elements in the Cu-Co/BPC catalyst, as illustrated in Fig. 1d, e.

The N₂ physisorption isotherm and pore size distribution in Fig. S1 (ESI[†]) demonstrate its mesopore structure, which are beneficial to exposure of the active sites and the mass transport of electrolytes during electrolysis process.29 The survey XPS spectrum of the Cu-Co/BPC catalyst is illustrated in Fig. S2 (ESI[†]). From the spectrum, it is evident that four elements-C, O, Cu and Co-are distributed on the surface of the Cu-Co/BPC catalyst (Fig. S2, ESI[†]). Fig. 2a presents the high-resolution XPS spectrum of C 1s, which displays a peak at 284.8 eV, indicating the presence of C-C/C=O chemical bonds in the catalyst. Additionally, a peak at 286.6 eV correlates to the existence of C-O bonds.³⁰⁻³² In addition, the high-resolution XPS spectrum of O 1s exhibits two characteristic peaks located at 532.9 eV and 538.2 eV (Fig. 2b). These two peaks are respectively representative of lattice oxygen and oxygen vacancies.33,34 The highresolution XPS spectrum of Cu 2p (Fig. 2c) shows that the peaks for Cu 2p_{3/2} appear at 934.1 eV and 932.3 eV. These two



Fig. 2 High-resolution (a) C 1s, (b) O 1s, (c) Cu 2p and (d) Co 2p spectra of Cu–Co/BPC.



Fig. 1 (a) Schematic for diagram illustrating synthetic procedure of Cu–Co/BPC. (b) XRD pattern and (c) TEM image (inset is the HR-TEM image) of Cu–Co/BPC. (d) HAADF-STEM and (e) corresponding elemental mapping images of Cu–Co/BPC.

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characteristic peaks are respectively caused by Cu²⁺ and Cu⁰/ Cu^{1+} . The peaks for Cu $2p_{1/2}$ appear at 954.2 eV and 952.1 eV. The Cu satellite peaks are located at 941.9 eV, 944.3 eV, and 962.8 eV. These two characteristic peaks are respectively caused by Cu²⁺ and Cu⁰/Cu¹⁺ (Fig. 2c). The high-resolution XPS spectrum of Cu 2p cannot distinguish between Cu⁰ and Cu¹⁺, so the Auger Cu LMM spectrum is used to determine whether both Cu⁰ and Cu^{1+} are present in the sample. As shown in Fig. S3 (ESI^{\dagger}), the Auger kinetic energy peaks in the range of 906-924 eV are broad and asymmetric. Two asymmetric peaks exist at 912.8 eV and 917.3 eV, where 912.8 eV is attributed to Cu¹⁺ and the peak at 917.3 eV is attributed to Cu⁰. Moreover, the Co element in the Cu-Co/BPC catalyst exhibits two main peaks in the highresolution XPS spectrum: Co $2p_{3/2}$ and Co $2p_{1/2}$. The Co $2p_{3/2}$ peak is observed at binding energies of 780.6 eV (Co^{3+}) and 783 eV (Co^{2+}). Similarly, the Co $2p_{1/2}$ peak is found at 795.2 eV (Co^{3+}) and 797 eV (Co^{2+}) (Fig. 2d). In Cu–Co/BPC, the Cu LMM (Fig. S3, ESI^{\dagger}) indicates that copper primarily exists as Cu⁰ within the catalyst at Fig. S6.† Additionally, small amounts of CuO and Cu₂O are also present in Cu-Co/BPC. From the Co 2p XPS spectrum (Fig. 2d), it can be observed that cobalt mainly exists in the Co³⁺ oxidation state. We analyzed the electronic properties of Cu/BPC, Co/BPC, and Cu-Co/BPC using highresolution Cu 2p and Co 2p XPS spectra (Fig. 2c, d and S4 ESI[†]). We were surprised to find that after the electrocatalytic reaction, the binding energy of Cu 2p significantly shifted toward higher energy, while the characteristic peaks of Co 2p shifted toward lower energy. This phenomenon indicates electron transfer between copper and cobalt, leading to the generation of active sites.

The electronic and coordinative structure of Cu-Co/BPC is further investigated by the X-ray absorption spectroscopy (XAS). The Cu K edge XANES spectra reveal that the near-edge absorption of Cu-Co/BPC is close to that of CuO, suggesting that the valence state of Cu in Cu-Co/BPC is close to +2 (Fig. S5a, ESI^{\dagger}). Notably, the characteristic peak at ~1.50 Å is shown in the Fourier-transformation k^3 -weighted Cu K edge EXAFS spectrum of Cu-Co/BPC (Fig. S5b, ESI[†]), confirming the absence of Cu–O bonds, while the presence of the peak at \sim 2.65 Å suggests the existence of the Cu–Co/Cu–Cu bonds. According to the Co K edge XANES spectra (Fig. S5c, ESI[†]), the valence states of Co species in Cu-Co/BPC is close to the valence states of Co species in CoO. As unveiled by Fig. S5d (ESI⁺), the presented peak at ~1.41 Å in the EXAFS spectrum of Cu-Co/BPC confirm the existence of Co-O bonds. Additionally, the peaks at ~ 2.54 Å obtained from the spectra of Cu-Co/BPC are assignable to the Co-Cu/Co-Co bonds, which is significantly inconsistent with that of Co foil, indicating the increased Co valence state of Cu-Co/BPC, ascribed to the electron transfer from Co to Cu due to lower electronegativity of Co (1.88) than Cu (1.90). This observation underscores the strong Cu-Co electronic interaction in Cu-Co/BPC.

To investigate the electrochemical activity of Cu–Co/BPC catalyst for NO_3^- electrochemical reduction, NitRR measurements were performed using a two-chamber cell (h cell) separated by a proton exchange membrane (Nafion 117). The cell consists of a Cu–Co/BPC catalyst coated on a carbon paper as



Fig. 3 (a) LSV curves of Cu–Co/BPC catalyst in 0.1 M K₂SO₄ and 0.1 M K₂SO₄ + 0.1 M KNO₃ solution. (b) R_{NH_3} and FE of Cu–Co/BPC catalyst obtained at different potentials for 2 h NitRR measurement. (c) S_{NH_3} of Cu–Co/BPC catalyst obtained at different potentials. (d) ¹H NMR spectra of Cu–Co/BPC catalyst using $^{15}NO_3^-$ and $^{14}NO_3^-$ as nitrogen source for NitRR and standards ($^{15}NH_4$)₂SO₄. (e) Recycling tests for Cu–Co/BPC catalyst during NitRR at –1.25 V (vs. RHE). (f) The FE and NH₃ yield were monitored over a 14 h NitRR conducted at –1.25 V (vs. RHE).

a working electrode, Ag/AgCl (saturated KCl solution) as a reference electrode, and a Pt mesh as a counter electrode.³⁵ Fig. 3a shows linear sweep voltammetry (LSV) curves of a mixed solution containing 0.1 M K₂SO₄ + 0.1 M KNO₃ and 0.1 M K₂SO₄ solution. Without KNO₃, the onset potential of the hydrogen evolution reaction (HER) was approximately -0.5 V (vs. RHE). With the addition of KNO₃, the potential in the mixed solution shifted to -357 mV (vs. RHE), compared to -721 mV (vs. RHE) in the solution without KNO₃, at a current density of 10 mA cm^{-2} . This indicates an increase in potential of 360 mV (vs. RHE) with the addition of KNO₃ at a current density of 10 mA cm⁻². Furthermore, the solution with nitrate added always had a higher current density when the same potential was applied, which diaplays that the catalyst has a high NitRR activity. The LSV results showed two peaks of reduction. The reduction peak at S1 near 0.08 V is attributed to the reduction of NO_3^- to NO_2^- , while the main reaction at the peak S2 involves the reduction of NO_2^- to NH_3 . Since the chemical reaction at peak S2 is predominant, the onset potential is determined near the vicinity of the S2 peak. The as-synthesized Cu-Co/BPC electrocatalyst was operated at six different potentials, which were selected in the range of -1 V to -1.4 V (vs. RHE). Each potential was tested for two hours of it in a mixture of 0.1 M KNO3 and 0.1 M K_2SO_4 (Fig. S6a, ESI[†]). The ultraviolet-visible (UV-Vis) absorption spectra of electrolytes obtained for each potential colored with indophenol blue reagent were measured (Fig. S6b, ESI[†]). The electrolysis product of NH₃ at a given potential was quantitatively determined by the indophenol blue method

(Fig. S7, ESI[†]). Fig. 3b show the calculated NH₃ yield rate and corresponding FE of Cu-Co/BPC at diverse potentials. It shows that the maximum NH_3 yield rate of 9114.1 \pm 244.8 $\mu g~h^{-1}\,cm^{-2}$ is achieved at -1.4 V (vs. RHE) while the highest FE of 84.5 \pm 1.6% is obtained at -1.0 V (vs. RHE). Obviously, the FE decreases with the applied potentials shifts to more negative values, mainly due to the competition of the HER on catalyst surface.³⁶ After the TEM characterization (Fig. S8, ESI[†]), we found that the structure of the catalyst remained largely unchanged during the reaction. That indicate the morphology and structural stability of the catalyst post-reaction are relatively well. We also tested the performance of Cu-Co/BPC, Cu/BPC and Co/BPC at -1.25 V (vs. RHE), which demonstrated the superiority of Cu-Co/BPC (Fig. S9, ESI†). In addition, the few byproduct NO_2^- is detected in the electrolytes (Fig. S10, ESI^{\dagger}), and the selectivity of NH_3 (S_{NH3}) of Cu–Co/BPC at different applied potentials were calculated as displayed in Fig. 3c. The $S_{\rm NH_2}$ of Cu–Co/BPC can reach the maximum FE of 97.8 \pm 0.65% at -1.0 V (vs. RHE) and excellent FE of NH₃ over 94.0% under a broad range from -1.0 to -1.4 V (vs. RHE) (Fig. 3c), demonstrating the high selective reduction of nitrate to NH₃.

To eliminate potential experimental errors caused by the self-disintegration of catalyst and environmental interference, series control experiments were carried out. As shown in Fig. S11 (ESI[†]), the negligible enhancing of signals associated with ammonia is observed in electrolysis at -1.25 V (vs. RHE) for 2 h in 0.1 M K₂SO₄ electrolyte without NO₃⁻ or carbon paper (CP) without Cu-Co/BPC catalyst, and Cu-Co/BPC electrode under open circuit potential (OCP) for 2 h in 0.1 M K₂SO₄ + 0.1 M KNO3 electrolyte. Furthermore, ¹⁵N isotope labelling experiments are also conducted. The ¹H nuclear magnetic resonance (¹H NMR) spectra were measured to verify the origin of produced NH₃. As exhibited in Fig. 3d, doublet coupled peaks of ¹⁵NH₄⁺ is observed in the ¹H NMR spectrum when using ¹⁵NO₃⁻ as the nitrogen source for NitRR, confirming the generated NH3 is produced from NitRR over Cu-Co/BPC catalyst.

The stability of electrochemical NH₃ production is an important indicator of the electrochemical performance of electrocatalysts. Therefore, the as-synthesized Cu-Co/BPC electrocatalyst was conducted out for 16 consecutive recycling tests in 0.1 M K₂SO₄ + 0.1 M KNO₃ solution at -1.25 V (vs..RHE). After recycling tests, it can be clearly seen in Fig. 3e that no remarkable decay in NH₃ yield rate and FE during the NitRR process. During the two-hour electrochemical reaction, the current gradually increased to 180 mA cm⁻² (Fig. S12, ESI[†]). Besides superior stability, the durability of the catalyst is another crucial parameter for practical applications. Then a 50 ml electrolysis cell was used for the 10 h NitRR experiments in our study. The Cu-Co/BPC catalyst was electrolyzed in a mixed solution of 0.1 M K₂SO₄ + 0.1 M KNO₃ at -1.25 V (vs..RHE) for 10 h with samples of the solution taken every 2 h. As shown in Fig. 3f, NH₃ production can be steadily increased with a slight decrease in FE, further demonstrating the superior stability of the Cu-Co/BPC catalyst. Notably, After electrolysis, the high-resolution Cu 2p and Co 2p XPS spectra were carried out to analyze the electronic properties of Cu-Co/BPC before and after NitRR process (Fig. S13, ESI[†]). Interestingly, after electrocatalytic reaction, the Cu 2p significantly shifted towards higher binding energies (Fig. S13a, ESI[†]), while the characteristic peaks of Co2p shifted to the lower binding energy (Fig. S13b, ESI[†]), indicating the existence of charge transfer from Cu atoms to Co atoms during electrocatalytic NitRR process, which results in high electrocatalytic selectivity and FE of Cu–Co/BPC for NH₃ synthesis. The Nyquist plot (Fig. S14, ESI[†]) shows that the catalyst exhibits the strongest corrosion resistance at this potential of -1.25 V vs. RHE. The improvement in corrosion resistance promotes the stability of the conversion of nitrate to ammonia. However, when the potential is -1.25 V vs. RHE, the reaction activity is relatively low. The reaction activity initially decreases with the rise in potential, reaches a maximum, and then increases.

Moreover, to further investigate the process of the NitRR reaction of the Cu-Co/BPC catalyst, the in situ Differential Electrochemical Mass Spectrometer (DEMS) was performed. The in situ DEMS is an instrument capable of analyzing the intermediates of electrochemical reactions as well as the final products in milliseconds. With the applied voltage is -1.25 V (vs..RHE), the m/z signals of 17, 30, 31, 32, 33 and 46, corresponding to NH₃, NO, HNO, NH₂O, NH₂OH and NO₂, respectively, appeared during ten continuous cycles. As shown in the Fig. 4, the amount of NH₃ gradually increased with the increase of reaction time, and the data were finally stable, indicating the catalytic reaction catalyzed by the catalyst has a relatively good stability. The other intermediate products show a decreasing trend with increasing reaction time, a phenomenon that supports the high FE of the reaction. The other intermediate products were significantly less than NH₃, indicating that NH₃ was the main product (Fig. 4). This above result suggests that the main reaction pathway of NitRR on Cu-Co/BPC catalyst may *NH₂OH \rightarrow *NH₃ \rightarrow NH₃(g). Furthermore, we also utilized *in* situ attenuated total reflection surface-enhanced infrared adsorption spectroscopy (ATR-SEIRAS) to monitor the evolution of NitRR intermediates. Fig. S15 (ESI[†]) shows the infrared signals when the in situ electrocatalytic NitRR on Cu-Co/BPC during the negative scan from OCP V to -1.4 V (vs. RHE). At



Fig. 4 In situ DEMS measurements of NitRR over Cu-Co/BPC.

the applied potential, the significantly enhanced infrared peaks at 1390 cm⁻¹ were assigned to N–O symmetric stretching vibration of NO₃⁻, indicates the activation and consumption of NO₃⁻ catalyzed by Cu–Co/BPC. Meanwhile, the vibration bands of adsorbed NO in bridge adsorption mode were detected at 1525 cm⁻¹. The infrared bands, centred at 1647 cm⁻¹, are assigned to the bending vibrations H–O–H loosely bound water molecules. In addition, the gradually enhanced infrared peaks at 1458 cm⁻¹, which can be ascribed to N–H bending vibration of NH₄⁺. Based on the *in situ* ATR-SEIRAS analysis, we proposed the following pathway for the NitRR on Cu–Co/BPC surface: NO₃⁻ \rightarrow *NO₃ \rightarrow *NO \rightarrow *NH₃. This observation agrees with the *in situ* DEMS results.

In summary, the Cu-Co bimetallic nanoparticles supported on biomass-derived porous carbon (Cu-Co/BPC) have been successfully synthesized as an effective electrocatalyst for electrocatalytic NitRR in 0.1 M K₂SO₄ + 0.1 M KNO₃ electrolyte. As a result, the as-synthesized Cu–Co/BPC affords an excellent $R_{\rm NH_{2}}$ of 9114.1 \pm 244.8 µg h⁻¹ cm⁻² at -1.4 V (vs. RHE) and a maximum FE of 84.5 \pm 1.6% at -1.0 V (vs. RHE), with outstanding stability during 10 h NitRR. We compared the performance of Cu-Co/BPC with other catalysts, demonstrating the superior performance of Cu-Co/BPC (Table S1, ESI†). The ¹⁵N isotope labelling experiments confirmed that the produced NH₃ indeed originated from the nitrate reduction reaction catalyzed by Cu-Co/BPC. Combining electrochemical experiments with in situ characterization, the possible reaction sites and electrocatalytic NitRR mechanism of Cu-Co/BPC have been proposed.

Data availability

The data supporting this article have been included as part of the ESI.†

Author contributions

Daopeng Li: conceptualization, investigation, visualization, writing – original draft. Shengbo Zhang: data curation, resources, supervision, writing – review & editing. Zhixian Mao: investigation, resources, formal analysis. KuiHu: investigation, resources, formal analysis. Dongnan Zhao: investigation, resources, formal analysis. Min Liu: software. Zhengguo Qu: software. Li Zhou: software. Tongfei Shi: funding acquisition, supervision, resources, writing – review & editing.

Conflicts of interest

There are no conflicts to declare.

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