



Cite this: DOI: 10.1039/d5sc06524k

All publication charges for this article have been paid for by the Royal Society of Chemistry

Accelerating $\text{N}_2\text{H}_{4(\text{ads})}$ formation by frustrated Lewis pairs in an oxyhydroxide for electrocatalytic ammonia oxidation into N_2

Meng-Ying Yin,^{ab} Xing-Yuan Xia,^{ab} Ting Dai,^{ab} Xia Chen,^{ab} Qiu-Ju Xing,^{ab} Lei Tian^{*abc} and Jian-Ping Zou^{ID}^{*ab}

Conventional catalysts based on the individual Oswin and Salomon (O–S) or Gerischer and Mauzer (G–M) mechanism cannot achieve direct electrocatalytic ammonia (NH_3) oxidation into nitrogen (N_2) with high activity and selectivity. Herein, a bimetallic nickel–cobalt oxyhydroxide ($\text{Ni}_{0.5}\text{--Co}_{0.5}\text{--OOH}$) with frustrated Lewis pairs was developed through an elaborate analysis of the binding types of NH_3 with the metal-oxide anode, efficiently integrating O–S and G–M mechanisms for converting NH_3 into N_2 with high activity (94%) and selectivity (63%), which is much superior to the anodes in the previous reports. The evidence of batch experiments, *in situ* characterization, and theoretical calculations confirms that two NH_3 molecules bind to Co^{3+} sites (Lewis acid) in CoOOH and hydroxy sites (Lewis base) in NiOOH , respectively. Then, the $\text{NH}_{2(\text{ads})}$ generated on the Lewis acid sites can quickly recombine with the $\text{NH}_{2(\text{ads})}$ desorbed from the Lewis base sites, accelerating the formation of $\text{N}_2\text{H}_{4(\text{ads})}$ and preventing the peroxidation of NH_3 . The electrocatalytic system assembled with the $\text{Ni}_{0.5}\text{--Co}_{0.5}\text{--OOH}$ anode shows excellent performance for NH_3 elimination in the secondary aerobic process effluent. Our work provides precious guidance for the design of novel anodes and sheds light on further promoting the performance of ammonia conversion.

Received 26th August 2025
Accepted 4th October 2025

DOI: 10.1039/d5sc06524k
rsc.li/chemical-science

Introduction

The electrocatalytic ammonia oxidation reaction (EAOR) is viewed as a promising conversion measure due to its ability to address the growing problem of ammonia-containing wastewater pollution.^{1–4} Nitrate and nitrite are the common products during the EAOR process, which could exacerbate secondary pollution.^{5–7} As a comparison, nitrogen (N_2) is the most desirable product of the EAOR.^{8–11} Therefore, it is urgent to exploit effective strategies to efficiently convert ammonia into N_2 .

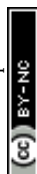
Direct electrocatalytic oxidation, based on anodic electron transfer, has been widely applied to ammonia pollution purification.^{12–14} It is more environmentally friendly than indirect oxidation, which is related to the electrochlorination process because there are no toxic secondary products (chlorine gas and chlorine-containing intermediates) produced.^{15–17} Until now, two possible conversion paths of the EAOR by direct

electrocatalytic oxidation have been proposed. One of them is the Oswin and Salomon (O–S) mechanism,¹⁸ in which the adsorbed NH_3 dehydrogenates to $\text{N}_{(\text{ads})}$ step by step and then converts into N_2 after the recombination of $\text{N}_{(\text{ads})}$ (eqn (1)–(5)). During this process, the active sites in the anode have high Lewis acidity and mainly bind to the N atom in NH_3 , ensuring strong binding energy for subsequent dehydrogenation. Unfortunately, excessive binding prevents the generated $\text{N}_{(\text{ads})}$ from recombining, making it highly susceptible to peroxidation to generate NO_2^- and NO_3^- .^{19–21} The O–S mechanism demonstrates strong activity for the EAOR but extremely low selectivity for N_2 generation. The other reaction path for the EAOR is the Gerischer and Mauzer (G–M) mechanism,^{22,23} in which the formation of N_2 is derived from the recombination of $\text{N}_x\text{H}_y(\text{ads})$ intermediates (eqn (1)–(3) and (6)). The G–M mechanism shows a higher selectivity for N_2 formation than the O–S mechanism because the anodic active sites have high Lewis basicity and mainly bind to the H atom in NH_3 , which is easily desorbed from the anodic surface after dehydrogenation, further efficiently converting into N_2 . However, the binding energy is relatively weak when the active site binds to the H atom in NH_3 , resulting in low efficiency of NH_3 dehydrogenation to $\text{NH}_{2(\text{ads})}$ and sluggish EAOR kinetics.^{24,25} The anode materials reported so far do not exhibit simultaneously high selectivity and high activity of the EAOR. Therefore, it is necessary to develop novel

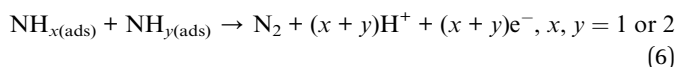
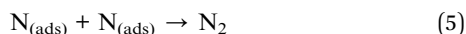
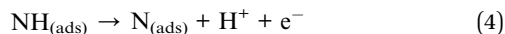
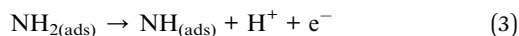
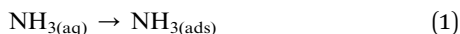
^aKey Laboratory of Jiangxi Province for Persistent Pollutants Prevention Control and Resource Reuse, Nanchang Hangkong University, Nanchang 330063, P. R. China

^bNational-Local Joint Engineering Research Center of Heavy Metals Pollutants Control and Resource Utilization, Nanchang Hangkong University, Nanchang 330063, P. R. China. E-mail: zjp_112@126.com

^cSchool of Chemistry and Chemical Engineering, Institute of Clean Energy and Materials, Guangzhou Key Laboratory for Clean Energy and Materials, Key Laboratory for Water Quality and Conservation of the Pearl River Delta, Ministry of Education, Guangzhou University, Guangzhou 510006, P. R. China



anode materials to realize the highly efficient and selective conversion of NH_3 into N_2 through direct oxidation.



Inspired by the above mechanisms, it is found that the dehydrogenation of NH_3 into $\text{NH}_{2(\text{ads})}$ is the rate-limiting step of the EAOR, while the recombination of $\text{NH}_{x(\text{ads})}$ and $\text{NH}_{y(\text{ads})}$ is the determining step for the selectivity of N_2 generation.²⁶ Theoretically, if the dehydrogenation in the O-S mechanism and the recombination of $\text{NH}_{x(\text{ads})}$ and $\text{NH}_{y(\text{ads})}$ in the G-M mechanism can be integrated together, the oxidation of NH_3 into N_2 can be effectively achieved with high activity and selectivity. Among the numerous anode materials,^{27–30} metal oxyhydroxides (MOOH) are the desired anode materials for the efficient and selective conversion of NH_3 into N_2 due to the presence of frustrated Lewis pairs.^{31,32} In detail, the metal atoms as Lewis acid sites can preferentially bind to the N atom due to the strong interaction between empty d-orbitals and lone pair electrons in the N atom, while the oxygen-containing functional groups as Lewis base sites can bind to the H atom (O–H–N) through hydrogen bonding. $\text{NH}_{x(\text{ads})}$ generated on the Lewis acid sites will quickly recombine with $\text{NH}_{y(\text{ads})}$ desorbed from the Lewis base sites, which can prevent the peroxidation of NH_3 , thereby promoting EAOR efficiency and N_2 generation selectivity. More importantly, the presence of surface hydroxy groups enables oxyhydroxides to effectively resist the erosion of Cl^- , thereby demonstrating high reliability in actual wastewater treatment. Although a small number of hydroxyl oxide electrodes (NiOOH or FeOOH) have been reported for use in the EAOR process, the activity of the EAOR and the selectivity for N_2 generation are poor (below 30%).^{33,34} Therefore, it is urgent to clarify the conversion law of NH_3 on different oxyhydroxides and develop highly active and selective oxyhydroxide anodes to achieve the EAOR.

Herein, a series of mono-metallic oxyhydroxide electrodes (MOOH, M: Mn, Co, Fe, Cu, Ni, and Zn) were synthesized, and their activity and selectivity of N_2 formation in the EAOR were investigated. Combined with the electronic structure of different transition metal ions, the transformation law between NH_3 and the active sites with different Lewis acidity/basicity was summarized. Based on this, a bimetallic oxyhydroxide was designed and synthesized, which can bind to the N and H atoms in NH_3 , respectively, integrating the O-S and the G-M mechanisms together, further enhancing the EAOR efficiency and the selectivity of N_2 formation. The synergistic effect of the bimetallic oxyhydroxide was explored by batch performance

evaluation and structural characterization. Moreover, through XPS, *in situ* diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS), and theoretical calculations, the conversion mechanism of NH_3 mediated by the bimetallic oxyhydroxide was further demonstrated. This work provides a new avenue for the design of anodic materials for the EAOR, which is in favor of the technological development of ammonia pollution purification.

Materials and methods

Materials and characterization

All chemicals were commercially obtained and used without further purification and the detailed information of materials is shown in Text S1 in the SI. The results of electrochemical analysis and characterization for catalyst structure and morphology are displayed in Text S2 in the SI. The evolution of conversion performance is displayed in Text S3 in the SI. The analytical methods for electrochemical characterization are displayed in Text S4 in the SI.

Preparation of the anodes

Nickel foam (NF) was first cleaned through sonicating consecutively in acetone, 3 M HCl, ethanol, and deionized water for 15 min to remove the oxide layer and degrease the surface of the NF. Then, the treated NF was directly used as the substrate to deposit Ni–Co precursors. In a typical procedure of Ni–Co precursor fabrication, 2.0 mmol (582.06 mg) of $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, 2.0 mmol (497.68 mg) of $\text{Ni}(\text{CH}_3\text{COO})_2 \cdot 4\text{H}_2\text{O}$, 3.75 mmol (138.8 mg) of NH_4F and 15 mmol (900.7 mg) of $\text{CO}(\text{NH}_2)_2$ were well dissolved in 50 mL of deionized water to form a mixture solution. The obtained solution was transferred into a 100 mL Teflon-lined stainless-steel autoclave. After immersing the washed NF ($1.5 \times 4 \text{ cm}^2$) substrate in the homogeneous solution, the autoclave was sealed and heated at 90 °C for 7 h. After the autoclave cooled down to room temperature, the Ni–Co precursor loaded NF was taken out from the autoclave and rinsed with ultrapure water several times. The black electrode of Ni–Co oxyhydroxide was obtained by immersing the precursors in 5% NaClO solution as an oxidant for 30 min. The chemical oxidation of the precursors was carried out at pH 3–4 adjusted with H_2SO_4 or NaOH .

Catalytic experiments

The initial concentration of ammonia was 20 mg N L^{-1} and 50 mM Na_2SO_4 electrolyte was added into a 100 mL undivided cylindrical glass electrolytic cell. The different anodes and the Pt cathode were fixed at a gap of 1 cm in the reactor and connected to a potentiostat (CHI 660E electrochemical workstation (Shanghai, Chenhua, China)). Ag/AgCl was used as the reference electrode. The immersed area of the electrode was 4.5 cm^2 . During the reaction process, the samples were taken with syringes at designated time intervals and immediately filtered through 0.45 μm polytetrafluoroethylene (PTFE) membranes. All experiments were conducted three times.



Analysis methods

The concentration of ammonia was measured by Nessler's reagent colorimetric method. The samples were distilled first and the absorbance was obtained from a UV-vis spectrophotometer (UV-9000S, Metash Instruments Ltd, China) at the wavelength of 420 nm.³⁵ The concentrations of nitrate (NO_3^-) and nitrite (NO_2^-) were determined by the ultraviolet spectrophotometry and *N*-(1-naphthyl)-ethylenediamine dihydrochloride spectrophotometric method, respectively.³⁶ The concentration of total nitrogen (TN) was measured by the traditional alkaline persulfate oxidation method.³⁷ The pH value was measured with a pH meter (Mettler Toledo-FE28, Swit). The ammonia removal (*R*) and selectivity of nitrogen conversion (S_N) were calculated using equations below:

$$R = \frac{[\text{NH}_3]_0 - [\text{NH}_3]_t}{[\text{NH}_3]_0} \times 100\% \quad (7)$$

$$S_N = \frac{[\text{N}]_t}{[\text{NH}_3]_0 - [\text{NH}_3]_t} \times 100\% \quad (8)$$

where $[\text{N}]$ denotes the concentration of nitrogen products, including $[\text{NO}_2^- - \text{N}]$, $[\text{NO}_3^- - \text{N}]$ and $[\text{N}_2 - \text{N}]$.

Current efficiency is expressed as eqn (9):

$$\text{CE (\%)} = \frac{n_i \times F \times (C_0 - C_t) \times V}{M \times \int_0^t I \text{d}t} \times 100\% \quad (9)$$

Here, C_0 is the initial concentration of $\text{NH}_3\text{-N}$, C_t is the concentration of $\text{NH}_3\text{-N}$ at degradation time t (s), V is the volume of the electrolyte (0.1 L), M is the molar mass of $\text{NH}_3\text{-N}$ (14 g mol^{-1}), I is the current density (A m^{-2}), A is the effective area of the electrodes (4.5 m^2), F is the Faraday constant ($96485.3 \text{ C mol}^{-1}$) and n is the number of electrons needed for the oxidation of one mole of ammonia ($n = 3$).

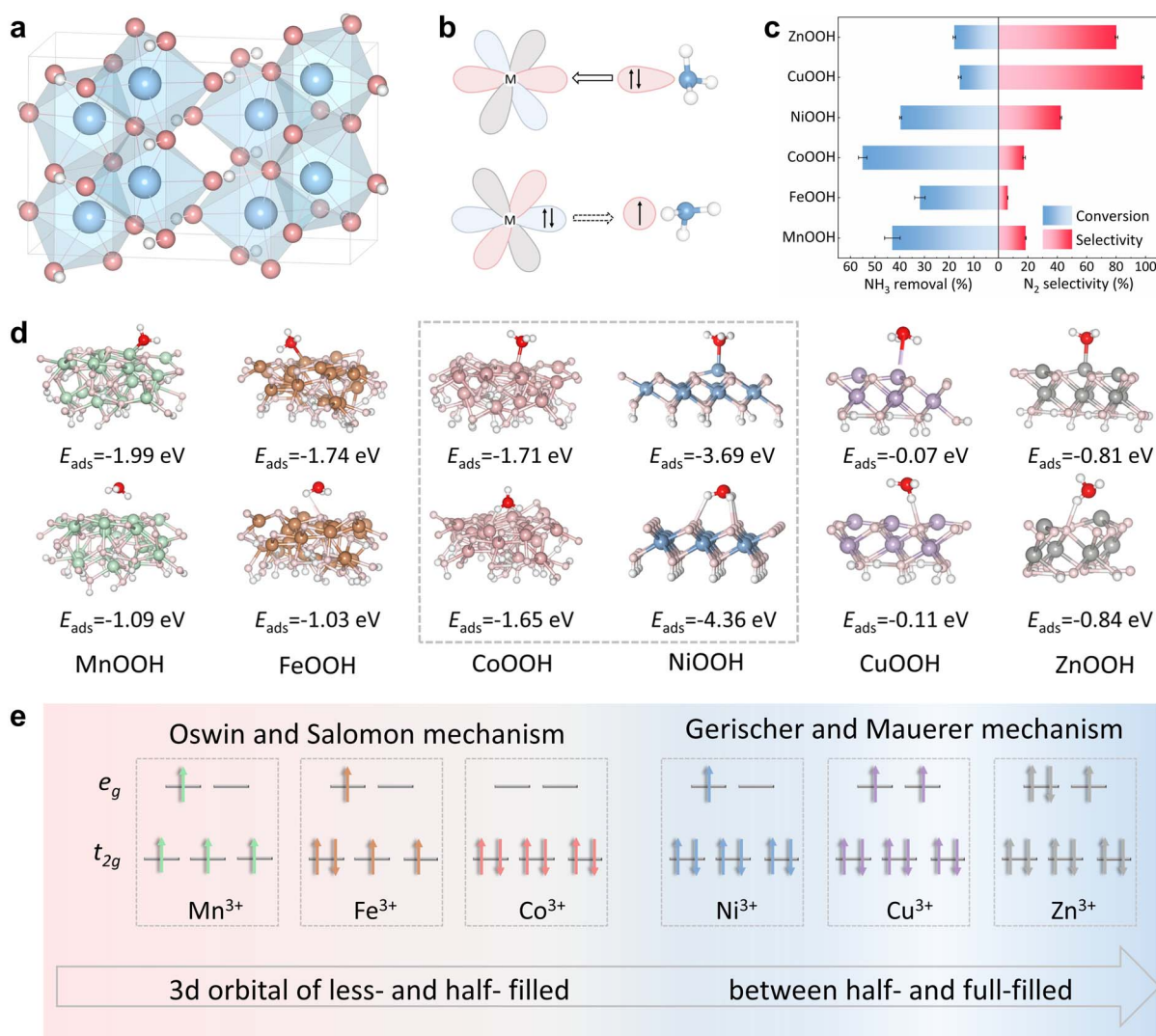


Fig. 1 (a) Crystal structure of MOOH; (b) different adsorption types with NH_3 ; (c) EAOR activity and selectivity of MOOH with different transition metal ions ($E = 1.8 \text{ V}$, $\text{pH} = 11$, $0.05 \text{ M Na}_2\text{SO}_4$); (d) calculated adsorption energies of ammonia molecules at metal and hydroxyl sites in mono-metallic oxyhydroxides; (e) diagram of d orbital electron configuration of different transition metal ions.

The energy consumption (EC, kWh m³) was calculated through eqn (10).

$$EC = \frac{U \times I \times t \times 1000}{\log(C_0/C_t) \times V} \quad (10)$$

where U is the applied voltage (V), I is the average current (A), t is the time (h), V is the solution volume (L), and $\log(C_0/C_t)$ is the logarithm of the initial and instant concentrations of the ammonia.

Theoretical calculation

The present first principle DFT calculations are performed with the Vienna *Ab initio* Simulation Package (VASP) with the projector augmented wave (PAW) method. Detailed information about the calculation procedure is provided in Text S5 in the SI.

Results and discussion

Lewis acidity/basicity on NH₃ conversion

The structure characterization studies of different MOOH are shown in Fig. S1 and S2. The prototypical MOOH oxyhydroxide has a typical cubic crystal structure. Metal atoms are located at the centre of MO₆ octahedra and the layers are linked by

hydrogen bonds (Fig. 1a). When there are many unoccupied d orbitals, metal atoms in MOOH can bind with N atoms in NH₃ during the adsorption process.³⁸ In contrast, the O atoms in MOOH may bind with H in NH₃ through a hydrogen bond (Fig. 1b).³⁹ With the outer 3d orbitals being gradually filled by electrons, the Lewis acidity of metal sites decreases and the Lewis basicity of hydroxy sites increases (Fig. 1c, e and S3). The EAOR performance of MOOH (from Mn³⁺ to Zn³⁺) increases first and then decreases (CoOOH > MnOOH > NiOOH > FeOOH > ZnOOH > CuOOH). When the d orbitals of the metal atom in MOOH are before half-filled, the adsorption energies of NH₃ at the metal sites are higher than those at the hydroxy sites (Fig. 1d), which can strongly bind with the N atom in NH₃, thus giving it strong EAOR performance. However, the overly strong force leads to the peroxidation of NH_{3(ads)} intermediates into NO₂[−]/NO₃[−], which follows the typical O-S mechanism and exhibits low N₂ formation selectivity. As a comparison, after the d orbital is half-filled, NH₃ adsorbs more strongly at the hydroxy sites than at the metal sites. Though the EAOR performance of MOOH is relatively sluggish, the N₂ formation selectivity is dramatically increased, which can be attributed to the change of the catalytic mechanism from O-S to G-M. In the two stages before and after a half-filled electronic orbital, CoOOH and

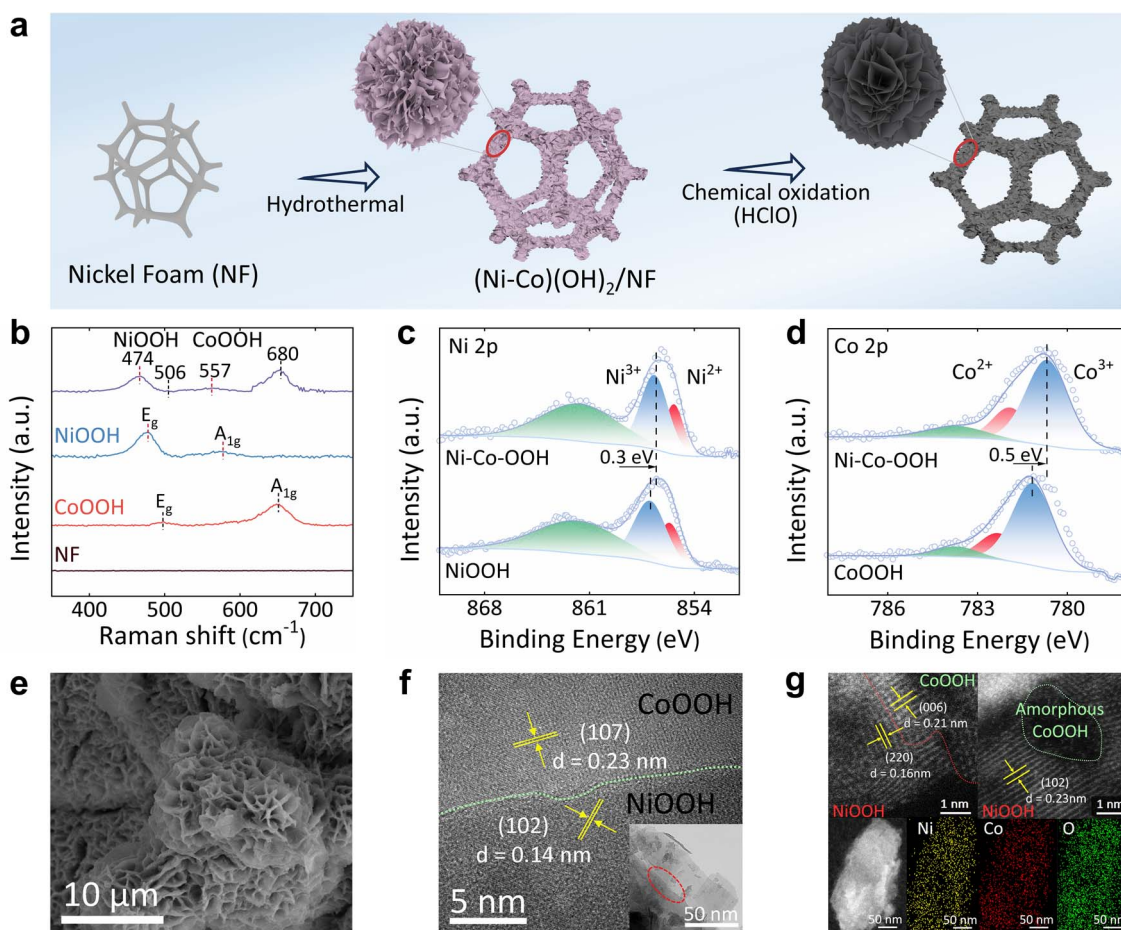


Fig. 2 (a) Synthesis diagram of Ni-Co-OOH; (b) Raman spectra of Ni-Co-OOH, CoOOH and NiOOH and nickel foam (NF); (c) Ni 2p XPS of Ni-Co-OOH and NiOOH; (d) Co 2p XPS of Ni-Co-OOH and CoOOH; (e) SEM images of Ni-Co-OOH; (f) TEM images of Ni-Co-OOH; (g) HAADF-STEM image of Ni-Co-OOH.



NiOOH exhibited optimal Lewis acidity/basicity and superior activity and selectivity for the EAOR compared to other MOOH or bimetallic systems (Fig. S4), respectively. Hence, Co and Ni elements were used to assemble the subsequent bimetallic oxyhydroxide catalysts to integrate the O-S and G-M mechanisms on a catalyst.

Synthesis and characterization of Ni-Co-OOH

The Ni-Co oxyhydroxide (Ni-Co-OOH) was synthesized following the schematic diagram in Fig. 2a. The XRD pattern of Ni-Co-OOH shows the presence of characteristic peaks attributed to NiOOH and CoOOH, respectively (Fig. S5). As shown in Fig. 2b, Raman spectra further demonstrate that the peaks located at 474 and 557 cm^{-1} are attributed to the E_g bending and A_{1g} stretching vibrations of Ni-O of NiOOH, respectively, whereas the peaks observed at 506 and 680 cm^{-1} are assigned to the CoOOH in Ni-Co-OOH.^{40,41} The Raman spectra show a slight shift in the vibrational frequencies of Ni-O and Co-O bonds compared to NiOOH and CoOOH, indicating there is electronic interaction between NiOOH and CoOOH. Besides, X-ray photoelectron spectroscopy (XPS) shows that two peaks in Co 2p_{3/2} at 780.8 and 781.9 eV are assigned to Co³⁺ and Co²⁺,^{42–44} respectively. For Ni 2p_{3/2} spectra, the peaks at 855.36 and 856.66 eV are attributable to the Ni²⁺ and Ni³⁺,^{45–47} respectively (Fig. 2c and d). Ni/Co-O bonds are also identified by the characteristic peak at 529.7 eV from O 1s spectra (Fig. S6). The peaks of Ni³⁺ and Co³⁺ are observed to shift by 0.3 eV and 0.5 eV, respectively, towards a lower binding energy compared to their mono-metallic oxyhydroxide, proving the electron transfer from Co³⁺ to Ni³⁺. The coupling of Ni³⁺ and Co³⁺ *via* the bridging O²⁻ facilitates π -donation from O²⁻ to Ni³⁺, resulting in the overall electron transfer from Co³⁺ to Ni³⁺, which is in agreement with the Raman spectra results. In addition, scanning electron microscopy (SEM) images show that the morphology of Ni-Co-OOH is a nanoplate structure covered with nano-flower spheres, where NiOOH and CoOOH are nanoplates and nano-flower spheres, respectively, confirming that Ni-Co-OOH is a composite and CoOOH is loaded on the surface of NiOOH (Fig. 2e, S7 and S8). A further inspection of these nanosheet structures with high-resolution TEM and spherical aberration-corrected high angle annular dark field scanning transmission electron microscopy (HAADF-STEM) was conducted. As shown in Fig. 2f and g, the images demonstrate the heterojunction region of Ni-Co-OOH, indicating that there are two different phases on both sides of the interface. The lattice distances are 1.6, 2.1 and 2.3 Å, which are consistent with the (2 2 0) facet of NiOOH, (0 0 6) facet of CoOOH and (1 0 2) facet of NiOOH, respectively. Therefore, the aforementioned characterization evidently confirms the successful preparation of the Ni-Co bimetallic oxyhydroxide catalyst by a two-stage method of hydrothermal and chemical oxidation and it is a biphasic composite with a heterojunction structure.

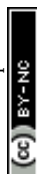
EAOR activity, N₂ selectivity and practical application

The EAOR activity of CoOOH and NiOOH is 93.8% and 24.1%, whereas the selectivity of N₂ is 1.3% and 35.7% and the

selectivity of NO₃⁻ is 74% and 24%, respectively, which typically follows the O-S and G-M mechanisms, respectively (Fig. 3a). Ni_x-Co_{1-x}-OOH catalysts with different Ni/Co ratios were obtained by varying the precursor dosages based on the similar solubility products of Ni(OH)₂ and Co(OH)₂ ($k_{sp}(\text{Ni}(\text{OH})_2) = 2 \times 10^{-15}$, $k_{sp}(\text{Co}(\text{OH})_2) = 1.6 \times 10^{-15}$) (Table S1). Ni_{0.5}-Co_{0.5}-OOH (Ni : Co = 0.5 : 0.5) exhibits higher EAOR activity (94.0%) and N₂ formation selectivity (63.0%) than Ni_{0.75}-Co_{0.25}-OOH, Ni_{0.25}-Co_{0.75}-OOH and the physically mixed CoOOH/NiOOH, which is superior to the electrocatalysts reported in the previous literature (Fig. S9–S11 and Table S2).^{48–54} Since the strong interaction between the Co-based Lewis acid sites and the N atom can cause difficulty in the desorption of N-containing intermediates from the anodic surface, the introduction of excessive CoOOH into NiOOH inhibits the formation of N₂, while generating more NO₃⁻ during the EAOR process. In addition, as shown in Fig. 3b, the NiOOH anode shows low faradaic efficiency (7.5%) and high energy consumption (150 kWh kg N⁻¹) for NH₃ conversion. Conversely, the current efficiency of Co-based oxyhydroxides significantly improves to 20–40%, while the low N₂ selectivity makes the NO₂⁻ and NO₃⁻ predominate the energy consumption. The Ni_{0.5}-Co_{0.5}-OOH can reach the highest faradaic efficiency (42%) and the energy consumption reaches a minimum value of 26.3 kWh kg N⁻¹, which further verifies the superior activity of Ni_{0.5}-Co_{0.5}-OOH for the EAOR.

With the increase of applied voltage from 1.6 V to 2.1 V *vs.* RHE, the EAOR activity of Ni_{0.5}-Co_{0.5}-OOH increases, while the N₂ formation selectivity is optimal (63.0%) at 1.8 V (Fig. 3c and S12). When excessive voltage is applied, the anodic oxygen evolution reaction (OER) may be severe, and NH₃ can be easily peroxidized into NO₃⁻ and NO₂⁻, resulting in a low N₂ formation selectivity. Besides, the removal of NH₃ and the formation selectivity of N₂ is significantly inhibited with increasing NH₃ concentrations (Fig. 3c and S13), which may be due to the saturation of the active sites on the electrode surface. In addition, the oxidation efficiency of NH₃ increases from 3.9% to 99.3%, whereas the N₂ formation selectivity declines from 100% to 38.4% with the increase in pH from 10 to 13 (Fig. 3c and S14). At a relatively low pH, NH₃ can be protonated into NH₄⁺ ($pK_a(\text{NH}_4^+/\text{NH}_3) = 9.25$), leading to the sluggish anodic oxidation of NH₄⁺ caused by electrostatic repulsion. In contrast, at a high pH, N₂ production selectivity is significantly inhibited due to the serious OER and NH₃ peroxidation (eqn (S4)–(S10)) at the anode, leading to the generation of NO₂⁻ and NO₃⁻. These results indicate that the best performance of the Ni_{0.5}-Co_{0.5}-OOH anode can be obtained when the voltage is 1.8 V, the NH₃ concentration is 20 mg L⁻¹ and the pH is 12.

To evaluate the practical application of the Ni_{0.5}-Co_{0.5}-OOH anode, ten cycles of experiments were carried out. As shown in Fig. 3d, the NH₃ conversion is around 90%, while the N₂ formation selectivity is still over 60% within 10 cycles, indicating that Ni_{0.5}-Co_{0.5}-OOH shows remarkable stability. Besides, there are no leaching concentrations of cobalt/nickel ions detected even in chloride-containing solutions and no change is observed in the metal value state and structure during ten cycles (Fig. S15–S18), which further confirms the stability of Ni_{0.5}-Co_{0.5}-OOH. Unlike the lattice oxygen mechanism involved



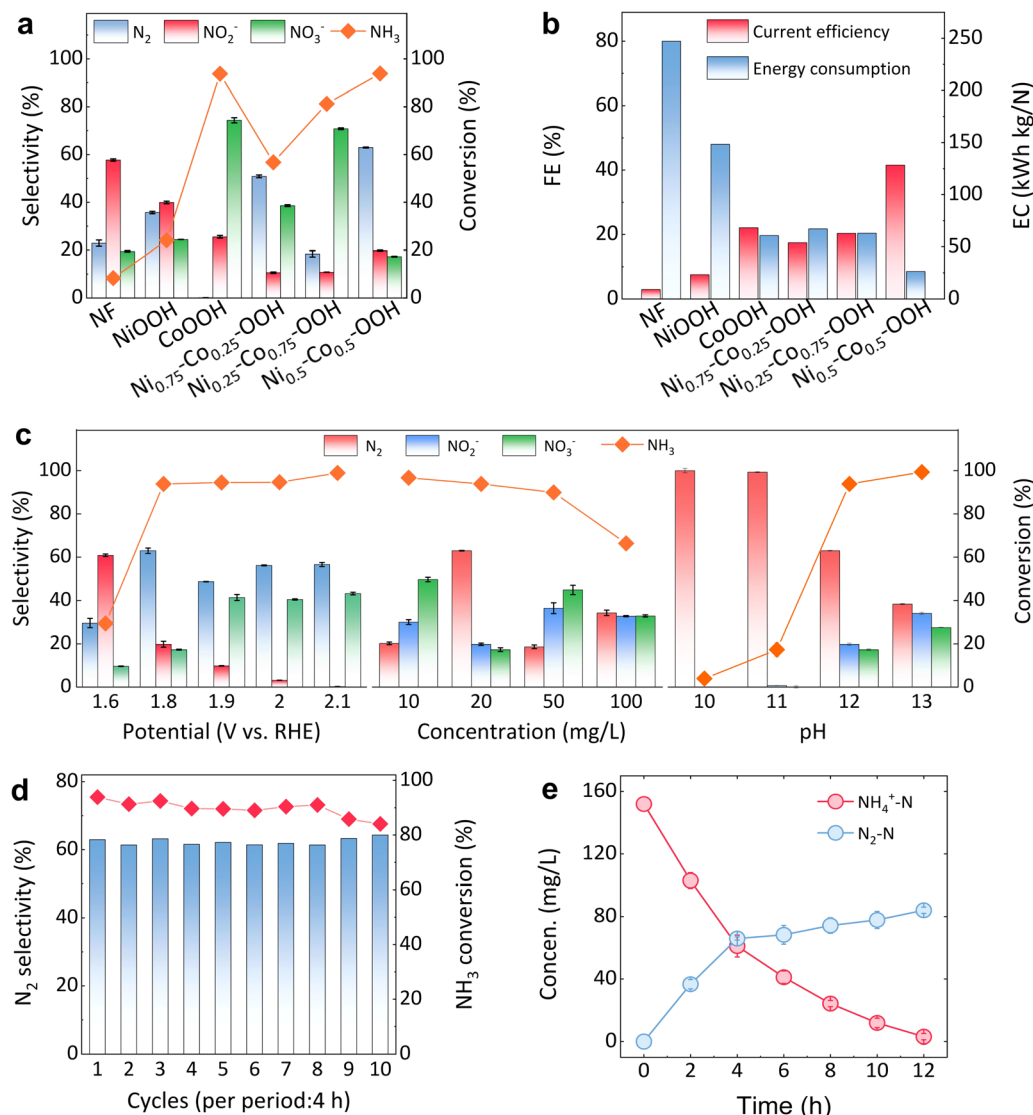


Fig. 3 (a) EAOR activity and product selectivity with different anodes ($E = 1.8$ V, $\text{pH} = 12$, 0.05 M Na_2SO_4); (b) Faradaic efficiency and energy consumption of the EAOR with different anodes ($E = 1.8$ V, $\text{pH} = 12$, 0.05 M Na_2SO_4); (c) EAOR activity and product selectivity of $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ under different conditions; (d) ten cycles of $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ ($E = 1.8$ V, $\text{pH} = 12$, 0.05 M Na_2SO_4); (e) EAOR activity of $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ for practical wastewater ($E = 2.0$ V, 0.05 M Na_2SO_4).

in the OER processes (Fig. S19), surface reconstruction does not occur during the EAOR process. In addition, the electrocatalytic system assembled with the $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ anode was applied to treat the practical ammonia-containing wastewater, which was the secondary aerobic process effluent. As shown in Fig. 3e and S20, the NH_3 conversion is around 99%, while the N_2 formation selectivity (55%) is much greater than that of nitrate (31%) and nitrite (11%). In conclusion, the $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ electrode has excellent stability and shows good application potential in the treatment of practical ammonia-containing wastewater.

Electrochemical characterization

Diverse electrochemical characterization studies were performed to explore the EAOR activity of $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$. As

shown in the linear sweep cyclic voltammetry (LSV) and Tafel curves (Fig. 4a and S21–S23), a lower onset potential (1.28 V) and a lower Tafel slope (128 mV dec^{-1}) are observed in the presence of NH_3 , demonstrating that $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ is more susceptible to the EAOR than the OER (1.45 V, 244 mV dec^{-1}). Besides, the polarization current of $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ in the presence of NH_3 is beyond that of CoOOH and the corresponding Tafel slope is the lowest derived from these catalysts, which further indicates exceptional activity of $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ for the EAOR (Fig. 4b, S24 and S25). Moreover, the electrochemical surface area is calculated from the values of double layer capacitance (C_{dl}) to be 35.5 , 31.75 , 50.25 , 38 and 69.5 cm^2 for NiOOH , CoOOH , $\text{Ni}_{0.75}\text{-Co}_{0.25}\text{-OOH}$, $\text{Ni}_{0.25}\text{-Co}_{0.75}\text{-OOH}$ and $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$, respectively (Fig. 4c and S26). This result confirms that there are more catalytically active sites on the $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$

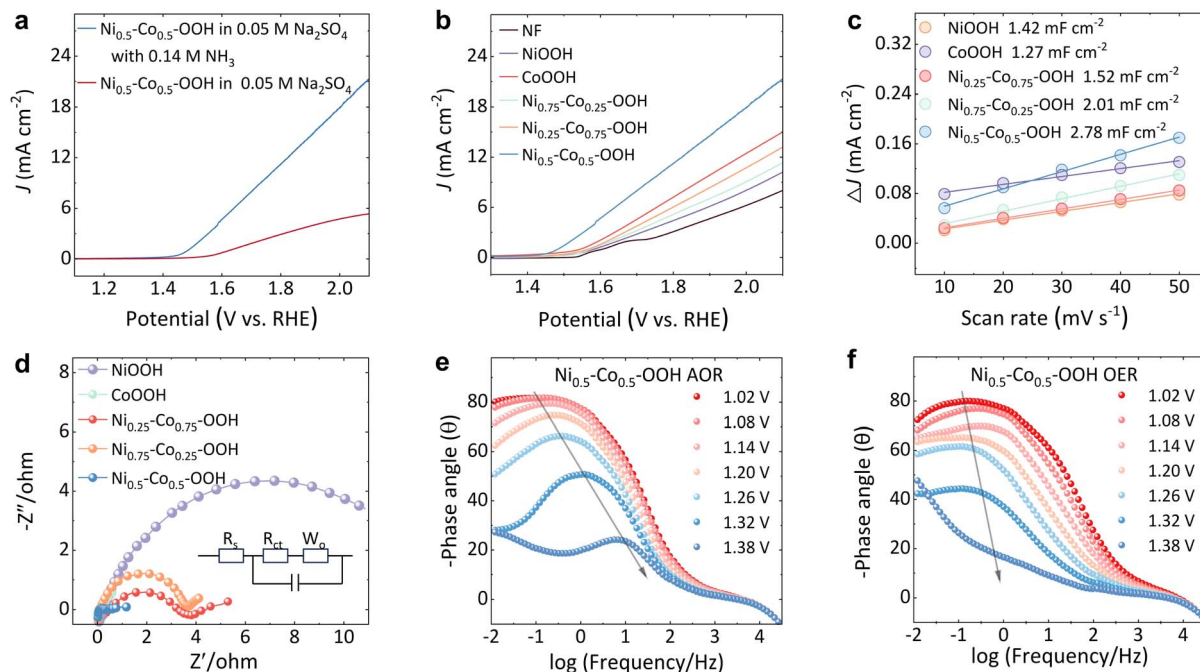


Fig. 4 (a) LSV curves of Ni_{0.5}-Co_{0.5}-OOH with or without NH₃; (b) LSV curves of different oxyhydroxides in the presence of NH₃; (c) electrochemical double-layer capacitance (C_{dl}) of different oxyhydroxides; (d) electrochemical impedance spectroscopy of different oxyhydroxides, the inset is the equivalent circuit for EIS fitting; (e and f) Bode phase plots of Ni_{0.5}-Co_{0.5}-OOH for the EAOR and OER.

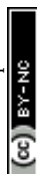
than other oxyhydroxides because of the synergistic effect of CoOOH and NiOOH. Furthermore, Ni_{0.5}-Co_{0.5}-OOH displays the smallest Nyquist semicircle diameter, suggesting the lowest charge transfer resistances (1.2 Ω) during the reaction process (Fig. 4d). Meanwhile, the *in situ* electrochemical impedance spectroscopy (EIS) tests were conducted to uncover the dynamic interfacial changes of the catalyst (Fig. 4e and f). It is observed that the phase angle of the HF region in the NH₃ solution sharply decreased from 1.26 V, indicating the fast charge-transfer kinetics of the EAOR. Thus, the Ni_{0.5}-Co_{0.5}-OOH/NF electrocatalyst exhibits extraordinary activity for the EAOR.

Direct NH₃ oxidation mechanism

XPS and *in situ* diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS) were used to detect the variation of Ni_{0.5}-Co_{0.5}-OOH/NF and NH₃ molecules during the EAOR process. As shown in Fig. 5a and b, the proportions of Ni³⁺/Ni²⁺ and Co³⁺/Co²⁺ first decrease and then increase close to the initial value during the EAOR process (Tables S3 and S4). Regardless of O-S and G-M mechanisms, trivalent metal ions with high oxidative potential in oxyhydroxides act as the electron acceptors to induce the dehydrogenation of NH_{3(ads)} into NH_{2(ads)}. As a result, the decrease of Ni³⁺ and Co³⁺ in Ni_{0.5}-Co_{0.5}-OOH occurs at the initial stage of the EAOR. Subsequently, the formed Ni²⁺ and Co²⁺ in Ni_{0.5}-Co_{0.5}-OOH can be further oxidized into Ni³⁺ and Co³⁺ through anodic electron transfer. Besides, it can be seen from the potential-dependent *in situ* DRIFTS that the peak around 1102 cm⁻¹ is ascribable to the symmetric N-H bending vibration, which stems from the NH_{3(ads)} or NH_{x(ads)} intermediates (Fig. 5c and S27). Notably, there is a new peak at

1244 cm⁻¹ with increasing potential, which is assigned to the stretching of H₂N-NH_{2(ads)}.^{55,56} This phenomenon manifests that the formation of N₂ during the EAOR process is derived from the further conversion of N₂H_{4(ads)}. A negligible peak around 1605 cm⁻¹ in DRIFTS, corresponding to the N-O vibration, is also observed, resulting from a few NO₂⁻ or NO₃⁻ generated from the peroxidation of NH₃. In addition, as shown in Fig. 5d, *in situ* Raman spectra of CoOOH and NiOOH show an increasing intensity of Ni-OH and Co-N peaks, respectively, with the potential increasing from 1.2 V to 1.9 V vs. RHE. It is noted that the stretching vibrations of Co-N and Ni-OH bonds are observed in Ni-Co-OOH throughout the reaction, which serves to further verify the synergistic effect of the frustrated Lewis pairs. The intensity of the Ni-OH peak, which arises from strong hydrogen bonding between NiOOH and NH₃,⁵⁷⁻⁵⁹ initially increases and then decreases as the potential is raised from 1.2 V to 1.9 V. This trend suggests that NiOOH undergoes reversible transformation to Ni(OH)₂ and eventually returns to its initial state, accompanied by the dehydrogenation of NH₃, formation of N₂H_{4(ads)}, and generation of N₂ on frustrated Lewis pairs. A similar reversible behavior is observed for CoOOH in the Ni-Co-OOH system. The change in Co-N strength also confirms that Co³⁺ is the site for NH₃ oxidation. The frustrated Lewis acid-base pair composed of Co³⁺ and hydroxy facilitates N₂H_{4(ads)} formation, thus promoting the selective ammonia oxidation into N₂.

DFT calculations were performed to explore the conversion path of NH₃ mediated by Ni_{0.5}-Co_{0.5}-OOH. According to the structural characterization, the Ni_{0.5}-Co_{0.5}-OOH model with the hetero-junction is simulated (Fig. S28). It is known that NiOOH following



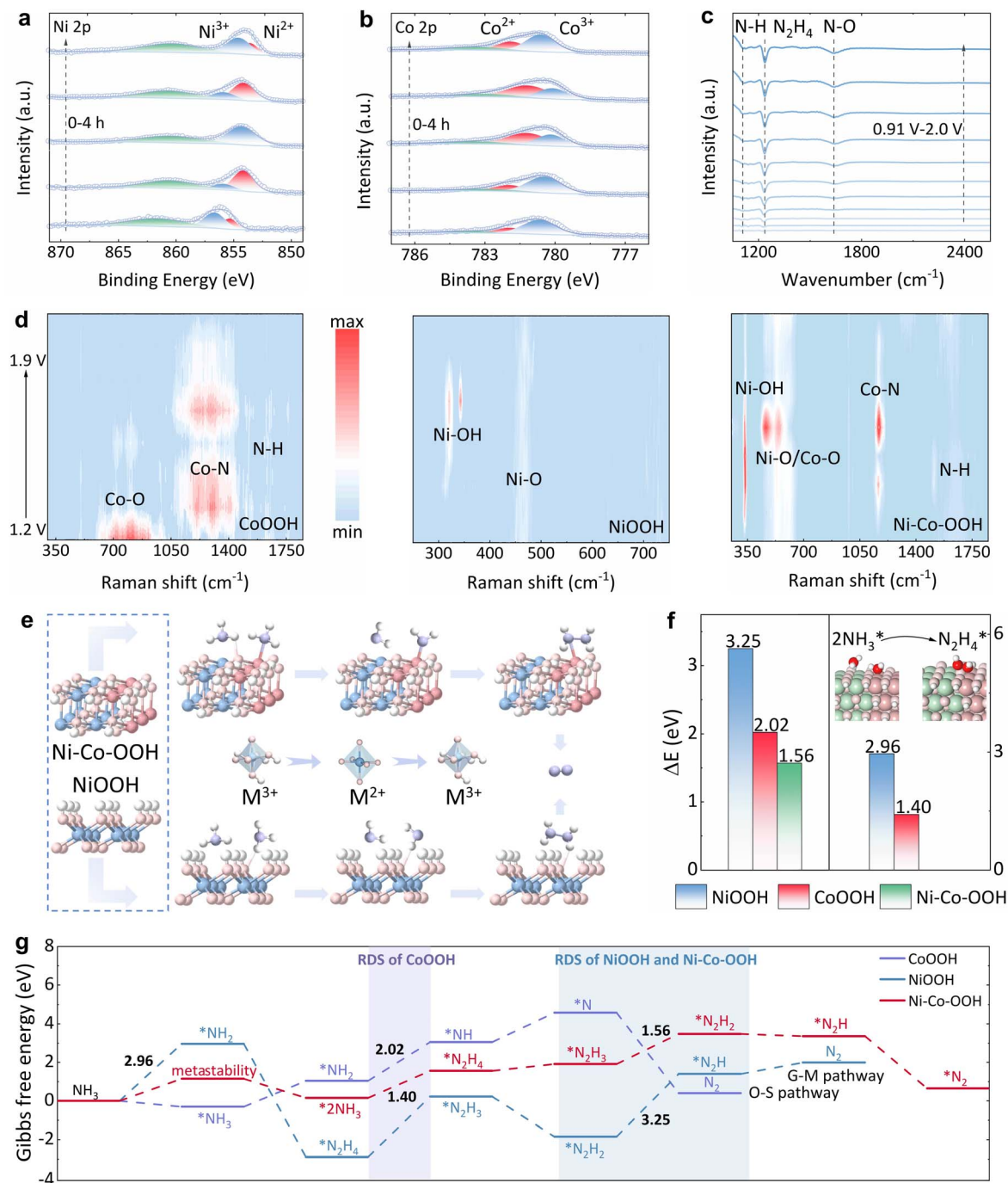
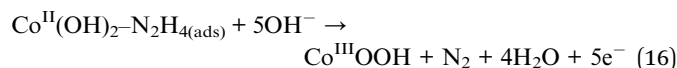
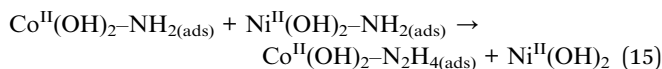
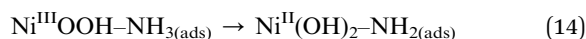
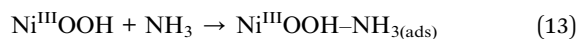
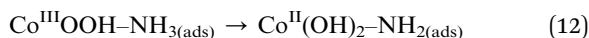
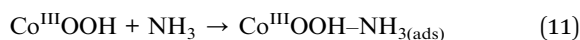


Fig. 5 (a) Ni 2p XPS of the $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ system within 4 h; (b) Co 2p XPS of the $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ system; (c) *in situ* DRIFTS spectra of the $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ system with applied potential positive scanning from 0.91 to 2.0 V vs. RHE; (d) *in situ* Raman test of CoOOH (left), NiOOH (middle) and $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ (right) systems; (e) the modeled $\text{N}_2\text{H}_{4(\text{ads})}$ formation pathway of the $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ and NiOOH system, respectively; (f) the pivotal potential gaps of RDS and the formation step of $\text{N}_2\text{H}_{4(\text{ads})}$ in Ni-Co-OOH , CoOOH and NiOOH systems; (g) Gibbs free energy diagrams of the EAOR by G-M and O-S mechanisms of the $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$ and CoOOH system, respectively.

the G-M mechanism exhibits high selectivity for N_2 generation, which is primarily derived from the conversion of $\text{N}_2\text{H}_{4(\text{ads})}$ intermediates. The $\text{N}_2\text{H}_{4(\text{ads})}$ generation energy barrier is compared to determine the N_2 generation selectivity of Ni-Co-OOH . As shown in Fig. 5e and f, for $\text{Ni}_{0.5}\text{-Co}_{0.5}\text{-OOH}$, two NH_3 molecules bind to Co^{3+} sites and hydroxy sites (the adsorption model as displayed in

Fig. S29), which is the integration of O-S and G-M mechanisms. Then, the $\text{NH}_{2(\text{ads})}$ generated on the Co^{3+} sites can quickly recombine with the $\text{NH}_{2(\text{ads})}$ desorbed from the hydroxy sites into $\text{N}_2\text{H}_{4(\text{ads})}$ (eqn (11)–(16)). As a comparison, for NiOOH , the formation of $\text{N}_2\text{H}_{4(\text{ads})}$ is derived from the recombination of two $\text{NH}_{2(\text{ads})}$ at hydroxy sites. The lower free energy of the $\text{N}_2\text{H}_{4(\text{ads})}$ formation

confirms that the Ni_{0.5}-Co_{0.5}-OOH anode exhibits a higher N₂ selectivity for the EAOR than NiOOH. In addition, the NH₃ conversion path is further analysed to explore the EAOR activity of the Ni_{0.5}-Co_{0.5}-OOH anode. As shown in Fig. 5f and g, after the formation of the N₂H_{4(ads)} intermediate, its continuous dehydrogenation into N₂ occurs at Co³⁺ sites. The rate-limiting step of the EAOR with the Ni_{0.5}-Co_{0.5}-OOH anode is the formation of N₂H_{2(ads)} (1.56 eV), which is the lowest in comparison with the O-S mechanism of CoOOH (2.02 eV) and G-M mechanism of NiOOH (3.25 eV). This result further verifies that the EAOR activity of the Ni_{0.5}-Co_{0.5}-OOH anode is superior to that of CoOOH. In summary, due to the synergetic effect of the frustrated Lewis pairs in Ni_{0.5}-Co_{0.5}-OOH, the O-S and G-M mechanisms are integrated, and the recombination of NH_{2(ads)} intermediates into N₂H_{4(ads)} intermediates is facilitated, thereby immensely promoting the EAOR activity and N₂ formation selectivity.



Conclusions

In conclusion, this study developed a bimetallic nickel-cobalt oxyhydroxide (Ni-Co-OOH) to integrate O-S and G-M mechanisms together. The synergy of the frustrated Lewis acid-base pairs is utilized for the selective conversion of NH₃ into N₂. Noteworthily, two NH₃ molecules bind to Co³⁺ sites (Lewis acid) in CoOOH and hydroxy sites (Lewis base) in NiOOH. Then, the NH_{2(ads)} desorbed from the hydroxy sites can rapidly recombine with the NH_{2(ads)} generated in Co³⁺ into N₂H_{4(ads)}, leading to the formation of N₂. The Ni-Co-OOH anode exhibits excellent activity (94%) and selectivity (63%) for the EAOR beyond other catalysts. This work provides a new idea for designing and preparing novel catalysts to improve the performance of ammonia conversion technology.

Author contributions

Meng-Ying Yin: writing – original draft, formal analysis, data curation, conceptualization. Xing-Yuan Xia: formal analysis, data curation. Ting Dai: formal analysis, data curation, software. Xia Chen: formal analysis, conceptualization, software. Qiu-Ju Xing: resources, supervision, visualization. Lei Tian: project administration, writing – review & editing, methodology. Jian-Ping Zou: writing – review & editing, methodology, resources, project administration, funding acquisition.

Conflicts of interest

There are no conflicts to declare.

Data availability

Data will be made available on request.

The general information, methods and supplementary data associated with this article are provided in the supplementary information (SI). See DOI: <https://doi.org/10.1039/d5sc06524k>.

Acknowledgements

We gratefully acknowledge the financial support of the National Natural Science Foundation of China (Grant No. 52470079, 52170082, 52300081, 51938007, and 52100186), Natural Science Foundation of Jiangxi Province (Grant No. 20212ACB203008), Science and Technology Department Project of Jiangxi Province (No. 20223AEI91001), and Key Laboratory of Jiangxi Province for Persistent Pollutants Prevention Control and Resource Reuse (No. 2023SSY02061).

Notes and references

- M. E. Ahmed, M. R. Boroujeni, P. Ghosh, C. Greene, S. Kundu, J. A. Bertke and T. H. Warren, Electrocatalytic Ammonia Oxidation by a Low-Coordinate Copper Complex, *J. Am. Chem. Soc.*, 2022, **144**, 21136–21145.
- T. Gupta, Sanyam, S. Sarawat, A. Mondal and B. Mondal, Electrochemical Ammonia Oxidation with a Homogeneous Molecular Redox Mediator, *Chem. Sci.*, 2025, **16**, 14377–14383.
- Y. Chen, G. Zhang, Q. Ji, H. Lan, H. Liu and J. Qu, Visualization of Electrochemically Accessible Sites in Flow-through Mode for Maximizing Available Active Area toward Superior Electrocatalytic Ammonia Oxidation, *Environ. Sci. Technol.*, 2022, **56**, 9722–9731.
- M. E. Ahmed, R. J. Staples, T. R. Cundari and T. H. Warren, Electrocatalytic Ammonia Oxidation by Pyridyl-Substituted Ferrocenes, *J. Am. Chem. Soc.*, 2025, **144**, 6514–6522.
- S. Bose, Y. Xia and R. N. Zare, Understanding the formation of nitrate from nitrogen at the interface of gas-water microbubbles, *Chem. Sci.*, 2024, **15**, 19764–19769.
- H. Y. Liu, H. M. C. Lant, J. L. Troiano, G. Hu, B. Q. Mercado, R. H. Crabtree and G. W. Brudvig, Electrocatalytic, Homogeneous Ammonia Oxidation in Water to Nitrate and Nitrite with a Copper Complex, *J. Am. Chem. Soc.*, 2022, **144**, 8449–8453.
- S. He, Y. Chen, M. Wang, H. Nuomin, P. Novello, X. Li, S. Zhu and J. Liu, Metal nitride nanosheets enable highly efficient electrochemical oxidation of ammonia, *Nano Energy*, 2021, **80**, 105528.
- L. Tian, P. Chen, X. H. Jiang, L. S. Chen, L. L. Tong, H. Y. Yang, J. P. Fan, D. S. Wu, J. P. Zou and S. L. Luo, Mineralization of cyanides via a novel Electro-Fenton system generating •OH and •O₂⁻, *Water Res.*, 2022, **209**, 117890.



- 9 Q. Jiang, D. M. Xia, X. F. Li, H. Zhang, R. J. Yin, H. J. Xie, H. B. Xie, J. Jiang, J. W. Chen and J. S. Francisco, Rapid N_2O formation from N_2 on water droplet surfaces, *Angew. Chem., Int. Ed.*, 2025, **64**, e202421002.
- 10 F. Habibzadeh, S. L. Miller, T. W. Hamann and M. R. Smith, Homogeneous electrocatalytic oxidation of ammonia to N_2 under mild conditions, *Proc. Natl. Acad. Sci. U. S. A.*, 2019, **116**, 2849–2853.
- 11 L. Tian, M. Y. Yin, L. L. Zheng, Y. Chen, W. Liu, J. P. Fan, D. S. Wu, J. P. Zou and S. L. Luo, Extremely efficient mineralizing CN^- into N_2 via a newly developed system of generating sufficient $\text{ClO}\bullet/\text{Cl}_2\bullet$ and self-decreasing pH, *Sep. Purif. Technol.*, 2023, **309**, 123021.
- 12 S. I. Venturini, D. R. Martins De Godoi and J. Perez, Challenges in Electrocatalysis of Ammonia Oxidation on Platinum Surfaces: Discovering Reaction Pathways, *ACS Catal.*, 2023, **13**, 10835–10845.
- 13 Y. Li, H. S. Pillai, T. Wang, S. Hwang, Y. Zhao, Z. Qiao, Q. Mu, S. Karakalos, M. Chen, J. Yang, D. Su, H. Xin, Y. Yan and G. Wu, High-performance ammonia oxidation catalysts for anion-exchange membrane direct ammonia fuel cells, *Energy Environ. Sci.*, 2021, **14**, 1449–1460.
- 14 F. Fang, Q. Y. Cheng, M. F. Wang, Y. Z. He, Y. F. Huan, S. S. Liu, T. Qian, C. L. Yan and J. M. Lu, Identifying Upper d-Band Edge as Activity Descriptor for Ammonia Oxidation on PtCo Alloys in Low-Temperature Direct Ammonia Fuel Cells, *ACS Nano*, 2025, **19**, 1260–1270.
- 15 L. Tian, X. Y. Xia, L. J. Zhou, L. L. Zheng, Q. J. Xing, L. S. Zhang, J. P. Zou and Z. Q. Liu, Enhanced chlorine enrichment via electron-deficient centers of Co(III) for efficient electrochlorination and ammonia removal, *Appl. Catal., B*, 2024, **340**, 123260.
- 16 Z. P. Yu, G. J. Xia, V. M. Diaconescu, L. Simonelli, A. P. LaGrow, Z. X. Tai, X. Y. Xiang, D. H. Xiong and L. F. Liu, Atomically dispersed dinuclear iridium active sites for efficient and stable electrocatalytic chlorine evolution reaction, *Chem. Sci.*, 2024, **15**, 9216–9223.
- 17 J. Z. He, C. Zhang, Y. Yang, J. W. Kang, C. Y. Zhang, D. He and J. X. Ma, Chlorine-Mediated Ammonia and Organics Transformation during Electrochemical Ammonia Recovery from Human Urine, *Environ. Sci. Technol.*, 2025, **59**, 13096–13107.
- 18 H. G. Oswin and M. Salomon, The anodic oxidation of ammonia at platinum black electrodes in aqueous KOH electrolyte, *Can. J. Chem.*, 1963, **41**, 1686–1694.
- 19 H. Y. Liu, H. M. C. Lant, C. Decavoli, R. H. Crabtree and G. W. Brudvig, pH-Dependent Electrocatalytic Aqueous Ammonia Oxidation to Nitrite and Nitrate by a Copper(II) Complex with an Oxidation-Resistant Ligand, *J. Am. Chem. Soc.*, 2025, **147**, 1624–1630.
- 20 G. Chen, X. L. Ding, P. He, T. Cheng, Y. Chen, J. Lin, X. Zhang, S. Zhao, N. Qiao and X. Y. Yi, Understanding the factors governing the ammonia oxidation reaction by a mononuclear ruthenium complex, *Chem. Sci.*, 2025, **16**, 7573–7578.
- 21 M. Anand, C. S. Abraham and J. K. Nørskov, Electrochemical oxidation of molecular nitrogen to nitric acid -towards a molecular level understanding of the challenges, *Chem. Sci.*, 2021, **12**, 6442–6448.
- 22 H. Gerischer and A. Mauzer, Untersuchungen Zur anodischen Oxidation von Ammoniak an Platin-Elektroden, *J. Electroanal. Chem. Interfacial Electrochem.*, 1970, **25**, 421–433.
- 23 K. Yang, J. Liu and B. Yang, Mechanism and Active Species in NH_3 Dehydrogenation under an Electrochemical Environment: An *Ab initio* Molecular Dynamics Study, *ACS Catal.*, 2021, **11**, 4310–4318.
- 24 X. Chen, Z. Y. Ke, X. Wang, H. Q. Jin, Y. W. Cheng, Y. K. Xiao, R. Jiang, Y. M. Da, L. Fan, H. X. Li, D. M. Liu, S. F. Yang and W. Chen, Fullerene Network-Buffered Platinum Nanoparticles Toward Efficient and Stable Electrochemical Ammonia Oxidation Reaction for Hydrogen Production, *Angew. Chem., Int. Ed.*, 2025, **137**, e202505180.
- 25 G. L. Li, Y. H. Shi, R. Y. Huang, T. G. Ma, Q. Zhuo, L. Xu, F. Jiang, Y. K. Zhai, Y. H. Xiao, Y. Yan and Y. Zhao, An ingenious structural design of NiCu-MOF bar-like nanosheet array: harnessing synergistic effects to attain exceptional ammonia oxidation performance, *Chem. Eng. J.*, 2025, **521**, 166548.
- 26 X. Yang, L. Sun, X. Liu, Z. Yang, H. Sun, W. Liu and H. Chen, Vacancy-driven ammonia electrooxidation reaction on the nanosized CeO_x electrode in nonaqueous electrolyte, *ACS Catal.*, 2024, **14**, 6236–6246.
- 27 H. Y. Fu, F. C. Gu, Y. Z. Niu, S. X. Liao, Z. Y. Bu, H. N. Wang, D. Yang, X. S. Wang and Q. Li, Spatially confined transition metals boost high initial coulombic efficiency in alloy anodes, *Chem. Sci.*, 2025, **16**, 418–424.
- 28 D. Sun, X. Hong, K. Wu, K. S. Hui, Y. Du and K. N. Hui, Simultaneous removal of ammonia and phosphate by electro-oxidation and electrocoagulation using $\text{RuO}_2\text{-IrO}_2/\text{Ti}$ and microscale zero-valent iron composite electrode, *Water Res.*, 2020, **169**, 115239.
- 29 C. X. Liu, Z. F. Teng, X. Liu, R. Zhang, J. Q. Chi, J. W. Zhu, J. F. Qin, X. B. Liu, Z. X. Wu and L. Wang, Stabilizing NiFe active sites using a high-valent Lewis acid for selective seawater oxidation, *Chem. Sci.*, 2025, **16**, 16321–16330.
- 30 F. Chang, W. Gao, J. Guo and P. Chen, Emerging Materials and Methods toward Ammonia-Based Energy Storage and Conversion, *Adv. Mater.*, 2021, **33**, 2005721.
- 31 Z. Hu, S. Lu, F. Tang, D. Yang, C. Zhang, Q. Xiao and P. Ming, High-performance precious metal-free direct ammonia fuel cells endowed by Co-doped Ni_4Cu_1 anode catalysts, *Appl. Catal., B*, 2023, **334**, 122856.
- 32 H. Zhong, Q. Zhang, J. Yu, X. Zhang, C. Wu, Y. Ma, H. An, H. Wang, J. Zhang, X. Wang and J. Xue, Fundamental Understanding of Structural Reconstruction Behaviors in Oxygen Evolution Reaction Electrocatalysts, *Adv. Energy Mater.*, 2023, **13**, 2301391.
- 33 X. Jiang, D. Ying, X. Liu, M. Liu, S. Zhou, C. Guo, G. Zhao, Y. Wang and J. Jia, Identification of the role of Cu site in Ni-Cu hydroxide for robust and high selective electrochemical ammonia oxidation to nitrite, *Electrochim. Acta*, 2020, **345**, 136157.



- Chem. Sci.