

# Synergizing Fe<sub>2</sub>O<sub>3</sub> Nanoparticles on Single Atom Fe-N-C for Nitrate Reduction to Ammonia at Industrial Current Densities

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The electrochemical reduction of nitrates (NO<sub>3</sub><sup>−</sup>) enables a pathway for the carbon neutral synthesis of ammonia (NH<sub>3</sub>), via the nitrate reduction reaction (NO<sub>3</sub>RR), which has been demonstrated at high selectivity. However, to make NH<sub>3</sub> synthesis cost-competitive with current technologies, high NH<sub>3</sub> partial current densities (*j*<sub>NH<sub>3</sub></sub>) must be achieved to reduce the levelized cost of NH<sub>3</sub>. Here, the high NO<sub>3</sub>RR activity of Fe-based materials is leveraged to synthesize a novel active particle-active support system with Fe<sub>2</sub>O<sub>3</sub> nanoparticles supported on atomically dispersed Fe-N-C. The optimized 3×Fe<sub>2</sub>O<sub>3</sub>/Fe-N-C catalyst demonstrates an ultrahigh NO<sub>3</sub>RR activity, reaching a maximum *j*<sub>NH<sub>3</sub></sub> of 1.95 A cm<sup>−2</sup> at a Faradaic efficiency (FE) for NH<sub>3</sub> of 100% and an NH<sub>3</sub> yield rate over 9 mmol hr<sup>−1</sup> cm<sup>−2</sup>. Operando XANES and post-mortem XPS reveal the importance of a pre-reduction activation step, reducing the surface Fe<sub>2</sub>O<sub>3</sub> (Fe<sup>3+</sup>) to highly active Fe<sup>0</sup> sites, which are maintained during electrolysis. Durability studies demonstrate the robustness of both the Fe<sub>2</sub>O<sub>3</sub> particles and Fe-N<sub>x</sub> sites at highly cathodic potentials, maintaining a current of −1.3 A cm<sup>−2</sup> over 24 hours. This work exhibits an effective and durable active particle-active support system enhancing the performance of the NO<sub>3</sub>RR, enabling industrially relevant current densities and near 100% selectivity.

## 1. Introduction

The synthesis of ammonia (NH<sub>3</sub>) based fertilizers is essential to support the growing global food demands. Currently, thermochemical NH<sub>3</sub> synthesis via the Haber Bosch (HB) process, accounts for ≈2% of global energy usage and more than 1.4% of global CO<sub>2</sub> emissions.<sup>[1–3]</sup> The electrochemical reduction of di-nitrogen (N<sub>2</sub>) is a theoretically ideal NH<sub>3</sub> synthesis pathway, however, direct activation of the highly stable and insoluble (in protic electrolytes) N<sub>2</sub> molecule remains challenging and unproven.<sup>[4,5]</sup> As an alternative for the N<sub>2</sub> molecule, recently there has been a revitalized interest in the more oxidized form of nitrogen, nitrate (NO<sub>3</sub><sup>−</sup>). Nitrate is an environmental pollutant present in groundwater runoffs due to heavy overfertilization practices and in industrial waste streams at varying concentrations (0.001–2 M).<sup>[6,7]</sup> The reduction of NO<sub>3</sub><sup>−</sup> to NH<sub>3</sub> is appealing

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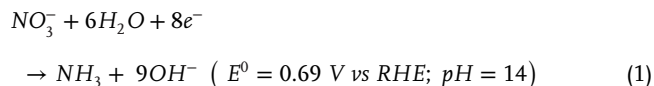
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to reutilize waste nitrogen into value-added  $\text{NH}_3$ , alleviating the demand on the HB process, while also serving as an alternative to traditional denitrification techniques, providing dual benefits for the nitrate reduction reaction ( $\text{NO}_3\text{RR}$ ). It should be noted that the  $\text{NO}_3\text{RR}$  alone is not a replacement for the HB process, because typically, the nitrogen atom in the  $\text{NO}_3^-$  originates from an HB-produced  $\text{NH}_3$ , but the  $\text{NO}_3\text{RR}$  can help to enhance the efficiency of the N-cycle.<sup>[8]</sup> Or the  $\text{NO}_3\text{RR}$  can be coupled with  $\text{N}_2$  plasma oxidation processes to be totally decoupled from the HB process.

Electrochemically, the  $\text{NO}_3\text{RR}$  is a complex  $8e^-$  transfer reaction, consisting of several possible soluble and insoluble intermediates ( $\text{NO}_2$ ,  $\text{NO}_2^-$ ,  $\text{NO}$ ,  $\text{N}_2$ ,  $\text{N}_2\text{O}$ ,  $\text{NH}_2\text{OH}$ ,  $\text{NH}_3$ , and  $\text{N}_2\text{H}_4$ ) and competes directly with the hydrogen evolution reaction (HER).<sup>[9]</sup> Aiming at industrial relevance, it is essential to optimize both catalyst activity and selectivity towards a singular product, in this discussion,  $\text{NH}_3$ . In practice, the use of an alkaline media (pH 13–14) can enhance the  $\text{NO}_3\text{RR}$  current throughput (over most metals) and significantly suppress the formation of the common  $2e^-$  side product, nitrite ( $\text{NO}_2^-$ ), often reporting the highest  $\text{NH}_3$  Faradaic efficiencies ( $FE_{\text{NH}_3}$ ) and yield rates ( $Yield_{\text{NH}_3}$ ) as shown in Table S1, Supporting Information.<sup>[10–14]</sup>

When targeting industrially relevant current densities, there is a competing compromise between the energy efficiency and  $\text{NH}_3$  partial current density ( $j_{\text{NH}_3}$ ), both of which are influenced by the applied cathodic potential and corresponding  $\text{NH}_3$  selectivity. In alkaline media, the standard reduction potential for the  $\text{NO}_3\text{RR}$  to  $\text{NH}_3$  (Equation (1)) is 0.69 V versus RHE.<sup>[12,15]</sup>



Currently, several reports achieve high  $FE_{\text{NH}_3}$  pushing upwards of 90%, some at mildly reductive potentials, resulting in relatively high cathodic energy efficiencies,  $\approx 40\%$ .<sup>[12,13,16–19]</sup> However, most of these reports suffer from limited  $j_{\text{NH}_3}$  ( $0.5 - 100 \text{ mA cm}^{-2}$ ), resulting in the need for largely scaled-up devices or stacks and thus intensive capital costs of these systems. In contrast, the  $\text{NO}_3\text{RR}$  systems with higher  $j_{\text{NH}_3}$  ( $100 < X \text{ mA cm}^{-2}$ ) relied on more cathodic overpotentials, suffering from largely reduced energy efficiencies (e.g., below 30%).<sup>[11,20–22]</sup> Establishing a trade-off between energy efficiency and  $j_{\text{NH}_3}$  remains ambiguous, although, a recent economic analysis quantifying the leveled cost of  $\text{NH}_3$ , suggested that the  $j_{\text{NH}_3}$  (production rate) has a more significant influence than the cell voltage or electricity price in reducing the leveled cost of  $\text{NH}_3$ .<sup>[22]</sup>

Cost-efficient platinum-group-metal-free (PGM-free) metals typically require more cathodic potentials to achieve a desirable  $j_{\text{NH}_3}$ . One way to achieve higher energy efficiency is to utilize PGMs and their alloys as these metals often have earlier (closer to the thermodynamic) reaction onset potentials. However, as more cathodic potentials are applied to achieve higher  $j_{\text{NH}_3}$ , they are typically out-competed by HER.<sup>[13,23]</sup> However, due to their scarcity and price, large-scale systems based on PGMs are not economically feasible. A promising approach was recently demonstrated, utilizing a Cu nanowire decorated with atomically dispersed Ru, able to achieve current densities at over 90%  $FE_{\text{NH}_3}$ .<sup>[10]</sup> Other recently reported rational strate-

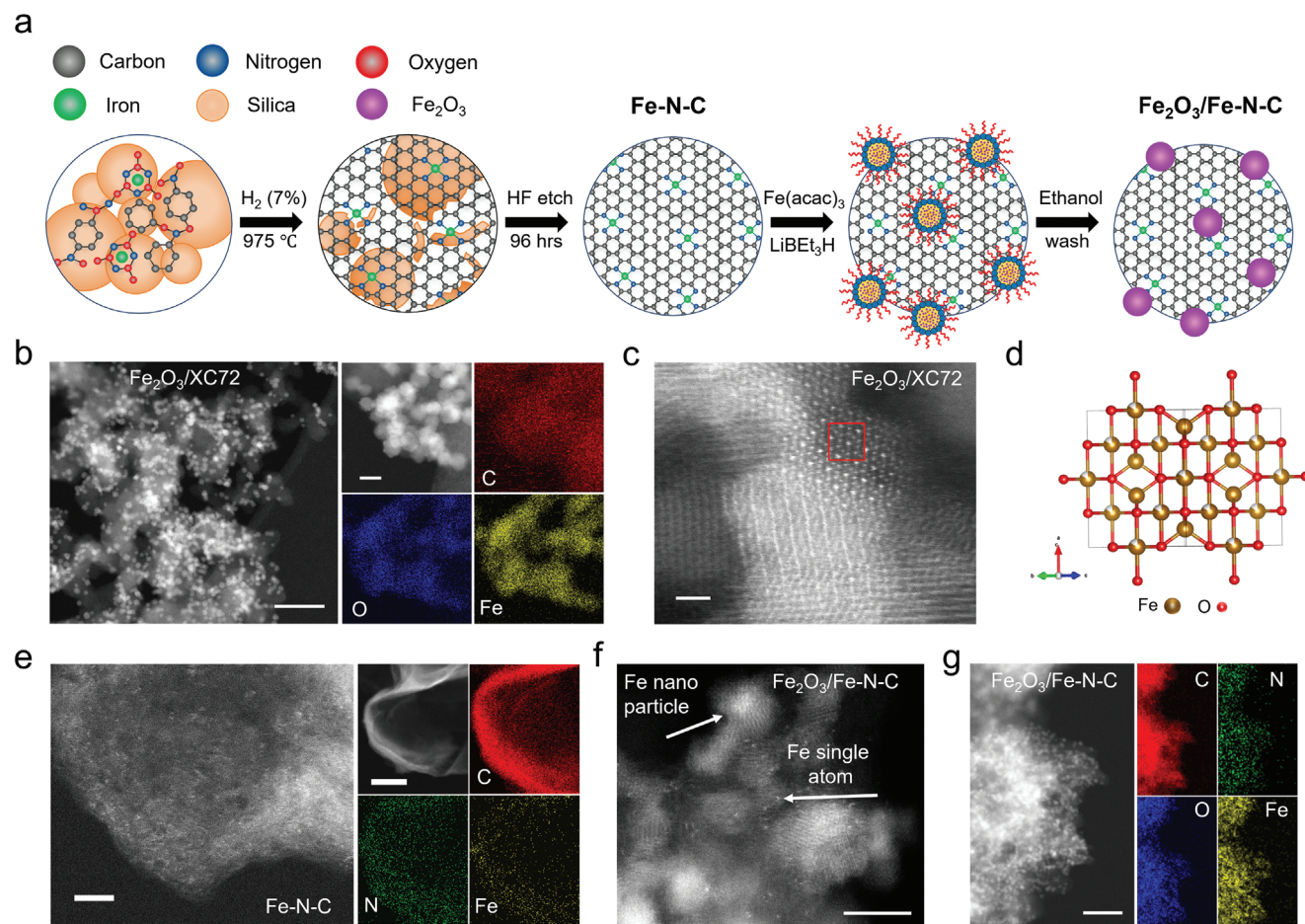
gies involve employing bi-metallic systems to selectively tailor the N-intermediate binding energies or selectively invoke a cascade mechanism for highly selective  $\text{NO}_3^-$  to  $\text{NH}_3/\text{N}_2$  conversion. It was shown that analogous to an approach utilized in the  $\text{CO}_2\text{RR}$ , pulsing cathodic and anodic potentials were shown to enhance  $\text{NO}_3\text{RR}$  by optimizing  $^*\text{NO}$  adsorption, reaching over 97%  $FE_{\text{NH}_3}$ . Another possible strategy currently being investigated for the oxygen reduction reaction (ORR) is the use of active supports to increase the activity and durability of the catalyst systems.<sup>[24–26]</sup> In those cases, a typical inert carbon support (Vulcan / carbon black) is replaced with an ORR active, atomically dispersed metal–nitrogen–carbon (M–N–C) support, often Fe–N–C or Co–N–C. It is speculated that possible electron donation between the platinum nanoparticles and the M–N–C can create more favorable intermediate adsorption energies, increasing the activity. Additionally, it was suggested that the M– $\text{N}_4$  active site can modify the electronic structure of the neighboring carbon, increasing the stability of the nanoparticles.

It has been shown in our previous works and complemented by other studies that atomically dispersed Fe–N–C is highly active for the  $\text{NO}_3\text{RR}$ , achieving a  $FE_{\text{NH}_3}$  greater than 90%.<sup>[12,27–29]</sup> This work builds upon the high  $\text{NO}_3\text{RR}$  activity of Fe–N–C catalysts and utilizes it as an active-support for  $\text{Fe}_2\text{O}_3$  nanoparticles, synthesizing a  $\text{Fe}_2\text{O}_3/\text{Fe–N–C}$  system for ultra-high  $\text{NO}_3\text{RR}$  to  $\text{NH}_3$  performance. Specifically, the  $3\times\text{Fe}_2\text{O}_3/\text{Fe–N–C}$  catalyst exhibits a potential independent selectivity ( $\approx 100\%$   $FE_{\text{NH}_3}$ ) between  $-0.4$  to  $-1.2$  V versus RHE, while increasing the  $j_{\text{NH}_3}$  up to nearly  $2 \text{ A cm}^{-2}$  (at a  $Yield_{\text{NH}_3}$  of more than  $9 \text{ mmol}_{\text{NH}_3} \text{ h}^{-1} \text{ cm}^{-2}$ ). In situ X-ray absorption near edge structure (XANES) supported by post-mortem X-ray photoelectron spectroscopy (XPS) reveals that the pre-reduction activation step is critical in achieving the ultrahigh  $\text{NO}_3\text{RR}$  performance, generating highly active, surface  $\text{Fe}^0$  sites. A durability test showed that the optimized  $3\times\text{Fe}_2\text{O}_3/\text{Fe–N–C}$  catalyst could maintain a  $FE_{\text{NH}_3}$  between 90–100% at a current of  $1.3 \text{ A cm}^{-2}$  for over 24-h, demonstrating the durability of utilizing an active-catalyst/active-support system.

## 2. Results and Discussion

### 2.1. $\text{Fe}_2\text{O}_3/\text{Fe–N–C}$ Synthesis and Characterization

$\text{Fe}_2\text{O}_3$  nanoparticles supported on atomically dispersed Fe–N–C ( $\text{Fe}_2\text{O}_3/\text{Fe–N–C}$ ) were synthesized by utilizing the sacrificial support method (SSM) for the Fe–N–C, followed by an organic solvent synthesis method for the  $\text{Fe}_2\text{O}_3$  nanoparticles. The SSM is a robust technique developed by our group for the synthesis of atomically dispersed M–N–C catalysts.<sup>[28]</sup> Schematically, the SSM is shown in Figure 1a, wherein a catalyst slurry of a carbon-nitrogen precursor is mixed with nanoporous silica and an iron-nitrate salt. The precursor mixture then undergoes a series of ball milling, pyrolyzing, and acid etching steps, yielding an exclusively atomically dispersed Fe–N–C support.<sup>[28]</sup> Afterward, the  $\text{Fe}_2\text{O}_3$  nanoparticles were synthesized on either a Vulcan-XC72 or Fe–N–C support, utilizing an organic solvent method. The high-angle annular dark-field scanning transmission electron microscopy (HAADF-STEM) image (Figure 1b) and TEM images (Figures S1 and S2, Supporting Information) show a homogeneous dispersion of the  $\text{Fe}_2\text{O}_3$  nanoparticles with



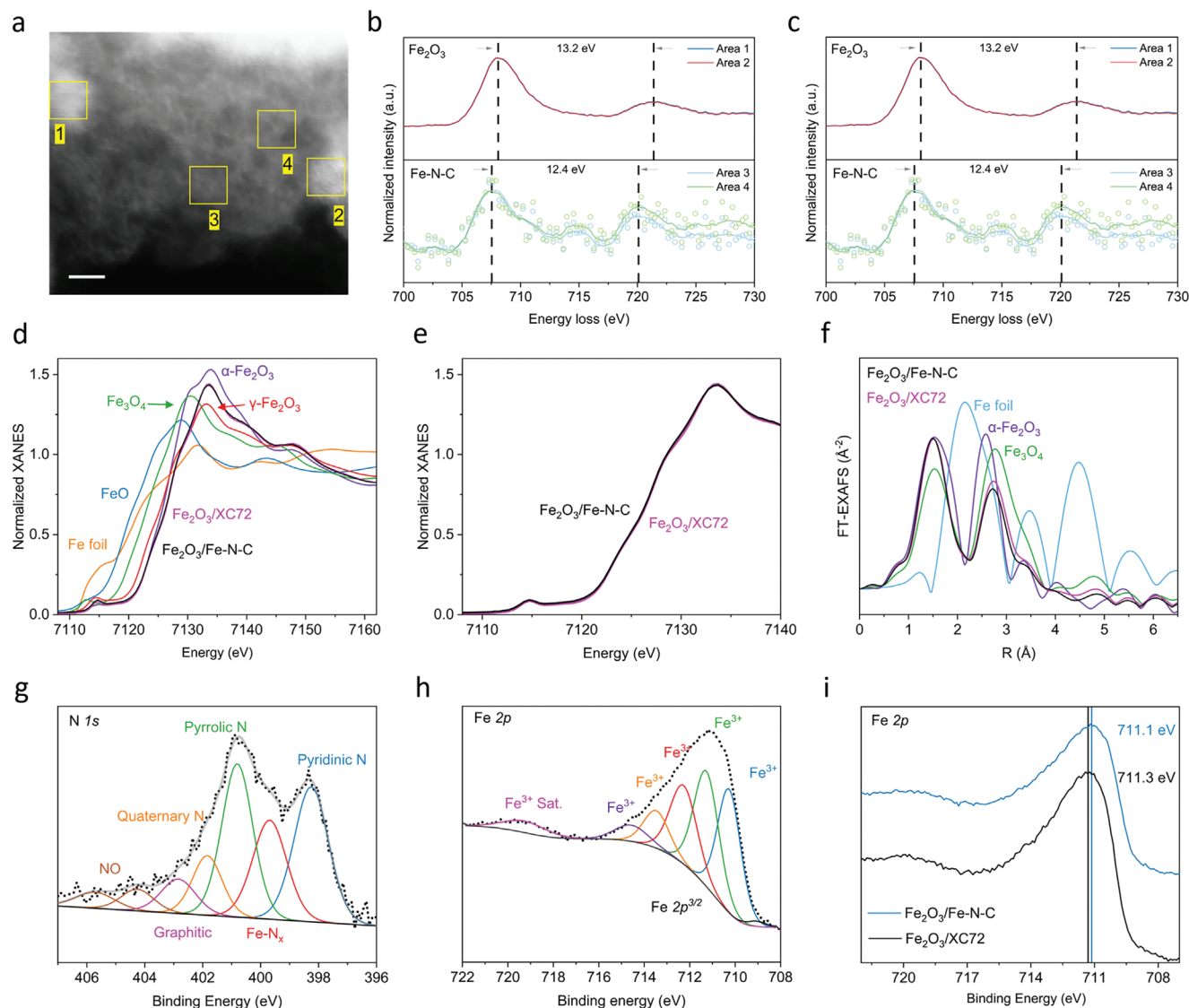
**Figure 1.** Synthesis schematic and AC-HAADF-STEM images of Fe<sub>2</sub>O<sub>3</sub>-based catalysts. a) Synthesis schematic, utilizing the sacrificial support method and an organic solvent synthesis to deposit Fe<sub>2</sub>O<sub>3</sub> nanoparticles on the atomically dispersed Fe-N-C. b) STEM images of the Fe<sub>2</sub>O<sub>3</sub> catalyst supported on XC72 carbon, the scale bar is 50 nm, with its corresponding EDS mapping, scale bar is 5 nm. c) Atomic resolution STEM image showing the Fe<sub>2</sub>O<sub>3</sub> spinel structure, scale bar is 2 nm. d) Schematic representation of the Fe<sub>2</sub>O<sub>3</sub> spinel crystal structure, created using VESTA. e) Atomic resolution STEM image of the atomically dispersed Fe-N-C catalyst support, with its corresponding EDS mapping, scale bar is 2 nm. f) Atomic resolution STEM image showing the Fe<sub>2</sub>O<sub>3</sub> nanoparticles supported on the atomically dispersed Fe-N-C, scale bar is 5 nm. g) EDS mapping of the Fe<sub>2</sub>O<sub>3</sub>/Fe-N-C catalyst, scale bar is 50 nm.

well-controlled sub-5 nm diameter. Additionally, the corresponding elemental mapping is shown in Figure 1b, for the Fe<sub>2</sub>O<sub>3</sub>/XC72 catalyst with a homogeneous distribution of Fe, O, and C. The Fe<sub>2</sub>O<sub>3</sub> nanoparticles have a spinel-like structure, typical of gamma-phase Fe<sub>2</sub>O<sub>3</sub>, as observed in the HAADF-STEM image in Figure 1c. A schematic of the Fe<sub>2</sub>O<sub>3</sub> spinel crystal structure is given in Figure 1d. The high magnification STEM image and energy dispersive X-ray spectroscopy (EDS) mapping in Figure 1e, verify the atomic dispersion of Fe and N-doping in the Fe-N-C support. After reducing the Fe<sub>2</sub>O<sub>3</sub> nanoparticles onto the Fe-N-C support, Figure 1f confirms that the Fe-N-C support retains its atomic dispersion as single atom Fe sites, clearly coexisting with the Fe<sub>2</sub>O<sub>3</sub> nanoparticles. The corresponding EDS mapping of the Fe<sub>2</sub>O<sub>3</sub>/Fe-N-C catalyst in Figure 1g, confirms the presence of nitrogen from the Fe-N-C support. The crystal structure of the catalyst was examined by X-ray diffraction (XRD), confirming the formation of Fe<sub>2</sub>O<sub>3</sub> (Figure S3, Supporting Information). As a comparison to the Fe<sub>2</sub>O<sub>3</sub> catalyst, CoO<sub>x</sub> and RuO<sub>x</sub> were also synthesized analogously and characterized using

TEM and XRD (Figures S4 and S5, Supporting Information). Furthermore, Raman spectroscopy was performed on Fe-N-C and XC72 supports, showing similar graphitic content between the catalyst supports, Figure S6, Supporting Information.

To evaluate the chemical state of the Fe<sub>2</sub>O<sub>3</sub>/Fe-N-C catalyst and the potential electronic structure changes triggered by interactions between Fe-N<sub>x</sub> sites and Fe<sub>2</sub>O<sub>3</sub> nanoparticles, atomic resolution electron energy loss spectroscopy (EELS), X-ray absorption spectroscopy (XAS), and XPS were utilized. EELS was used to probe the valence state of the single-atom Fe and the Fe<sub>2</sub>O<sub>3</sub> sites. Figure 2a shows the locations where EELS spectra were taken for Fe<sub>2</sub>O<sub>3</sub> particles (locations 1 and 2) and single atom Fe sites (locations 3 and 4). Both spectra show the Fe-L<sub>3,2</sub> edges, however, there is an L<sub>3</sub>, L<sub>2</sub> excitation edge shift to lower energy loss and reduced L<sub>3</sub>/L<sub>2</sub> white line ratio for single atom Fe (peak spacing of 12.4 eV), compared to Fe<sub>2</sub>O<sub>3</sub> (13.2 eV) in Figure 2b. This energy shift and quantitative analysis of the Fe-L<sub>3</sub>/L<sub>2</sub> edges suggest the single atom Fe to be in an oxidation state lower than Fe<sup>3+</sup>, in agreement with our previous work





**Figure 2.** Local chemical and coordination environment of the  $\text{Fe}_2\text{O}_3$  nanoparticle catalysts. a) AC-STEM image and EELS spectra locations on the  $\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  catalysts, scale bar is 2 nm. b) EELS spectra of the  $\text{Fe-L}_{3,2}$  edges of the  $\text{Fe}_2\text{O}_3$  nanoparticle (top) and atomically dispersed  $\text{Fe-N}_x$  sites (bottom). c) EELS spectra comparing the energy loss of the  $\text{Fe-L}_{3,2}$  edge of the  $\text{Fe}_2\text{O}_3$  supported on  $\text{Fe-N-C}$  or XC72. Fe K-edge XAS data for the  $\text{Fe}_2\text{O}_3$  catalysts supported on both  $\text{Fe-N-C}$  and XC72. d) XANES spectra with the corresponding references and e) XANES spectra comparing  $\text{Fe}_2\text{O}_3$  supported on XC72 versus  $\text{Fe-N-C}$ . f) Fourier transformed ( $k^2$ -weighted) EXAFS of the  $\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  catalyst and corresponding references. XPS spectra for the  $\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  catalyst, h) N 1s spectra, and i) Fe 2p XPS spectra. j) Comparison of the Fe 2p XPS spectra for the  $\text{Fe}_2\text{O}_3$  catalyst supported on  $\text{Fe-N-C}$  or XC72.

where  $\text{Fe-N-C}$  has an oxidation state  $\text{Fe}^{2.6+}$ .<sup>[28]</sup> At the same time, the analysis for the  $\text{Fe}_2\text{O}_3$  nanoparticles indicated an oxidation state of  $\text{Fe}^{3+}$ , in agreement with the XAS and XPS (Figure S7, Supporting Information) results. Atomic resolution EELS was further applied for a highly localized evaluation of possible  $\text{Fe}_2\text{O}_3$  and  $\text{Fe-N}_x$  interactions modifying the electronic structure of the nanoparticles. Comparing the  $\text{Fe-L}_{3,2}$  edges of the  $\text{Fe}_2\text{O}_3$  nanoparticles supported on  $\text{Fe-N-C}$  or XC72 reveals a 0.1 eV shift in the energy loss, Figure 2c. Such small shifts in energy loss can arise due to experimental conditions and inaccuracies in the selected method for spectra processing (something that requires extreme care and is system-dependent).<sup>[30]</sup> Therefore,

further complementary techniques are employed to evaluate possible nanoparticle–single atom interactions.

XANES of the Fe K-edge (7112 eV) was employed to investigate the chemical state of the  $\text{Fe}_2\text{O}_3$  nanoparticles on both supports, Figure 2d. The Fe K-edge XANES spectra for both  $\text{Fe}_2\text{O}_3$  catalysts again confirm that Fe is in a  $\text{Fe}^{3+}$  oxidation state. The XANES spectra for the catalysts are in between those for the  $\gamma\text{-Fe}_2\text{O}_3$  and  $\alpha\text{-Fe}_2\text{O}_3$  references. We thus further refer to these Fe species simply as  $\text{Fe}_2\text{O}_3$ . Note that our previous work, utilizing the same  $\text{Fe-N-C}$  suggests the  $\text{Fe-N}_x$  sites to be in a  $\text{Fe}^{2.6+}$  oxidation state.<sup>[28]</sup> Again, to evaluate possible  $\text{Fe}_2\text{O}_3$  ( $\text{Fe}^{3+}$ ) and  $\text{Fe-N}_x$  ( $\text{Fe}^{2.6+}$ ) interactions, the rising edge and pre-edge of the  $\text{Fe}_2\text{O}_3$  supported



on Fe–N–C and XC72 were compared, as shown in Figure 2e, however, no meaningful shifts in the energy were observed. The local coordination of Fe species was further analyzed by Fourier-transform extended X-ray adsorption fine structure (FT-EXAFS) in Figure 2f. For the Fe<sub>2</sub>O<sub>3</sub> catalysts, regardless of the support, two dominating peaks are observed. One at a low bond distance ( $\approx 1.4$  Å, phase uncorrected), corresponding to the expected Fe–O coordination and one at a larger bond distance ( $\approx 2.6$  Å, phase uncorrected), which corresponds to the Fe–Fe coordination in Fe<sub>2</sub>O<sub>3</sub>, and is in agreement with the Fe-oxide reference materials. As shown in our previous work, the EXAFS for the Fe–N–C supports exhibits only one peak at low bond distance for Fe–N, confirming its atomically dispersed nature.<sup>[28]</sup> The corresponding EXAFS *k*-space analysis is shown in Figure S8, Supporting Information.

To further investigate the chemical environment and coordination of the Fe species, XPS was performed. Figure 2h shows the N 1s spectra for the Fe<sub>2</sub>O<sub>3</sub>/Fe–N–C catalyst, confirming the Fe–N<sub>x</sub> coordination, along with the (NO<sub>3</sub>RR active) pyridinic and pyrrolic N-moieties, characteristic of the Fe–N–C support (XPS for the Fe–N–C support is given in Figure S9, Supporting Information). From the deconvoluted Fe 2p spectra in Figure 2i, the oxidation state of Fe<sub>2</sub>O<sub>3</sub> is Fe<sup>3+</sup>, regardless of the support used, further corroborating the EELS and XANES analysis. Full XPS deconvolution of the C 1s, O 1s, N 1s, and Fe 2p spectra for the Fe<sub>2</sub>O<sub>3</sub>/Fe–N–C and Fe<sub>2</sub>O<sub>3</sub>/XC72 catalysts are shown in Figures S7 and S10, Supporting Information, respectively. Again, to elucidate possible particle-single atom interactions, the Fe 2p spectra between the Fe<sub>2</sub>O<sub>3</sub> supported on Fe–N–C and XC72 were compared, Figure 2j, a binding energy shift of  $\approx 0.2$  eV is observed, comparable to that of recent nanoparticle-single atom reports in the literature.<sup>[31–33]</sup> However, binding energy shifts during the deconvolution and calibration (commonly to features in C 1s spectrum) of the spectra, can easily induce binding energy shifts on the level of 0.2 eV, especially when the complex nature of the carbon is changing in the compared catalyst supports.<sup>[34]</sup>

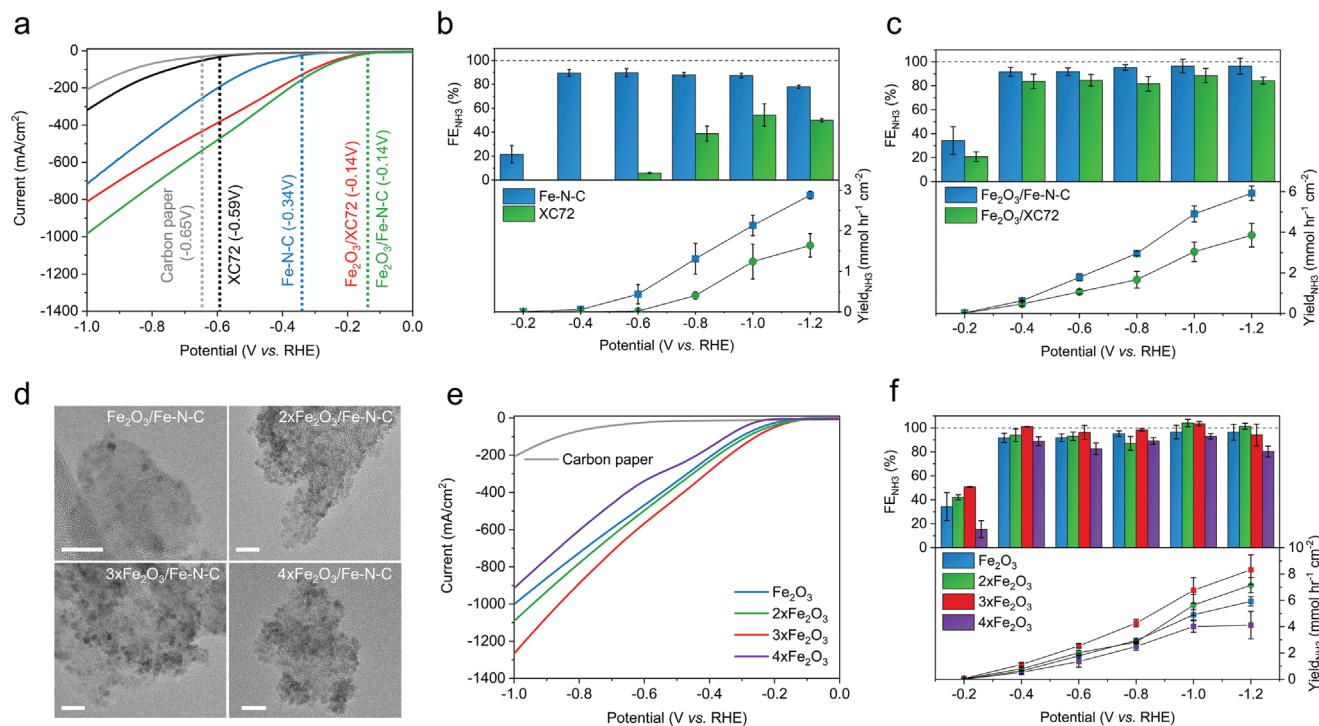
It should be noted that as the popularity of nanoparticle/single atom support systems increases, extreme care must be taken in the interpretation of the chemical state. Often nanoparticle/single atom support electronic interactions are claimed exclusively through ambiguous shifts in the XPS spectra, and subsequently used as the foundation for interesting computational models and reaction mechanisms and attributed to any increased activity and stability. In this work, after rigorously investigating the electronic structure of the Fe<sub>2</sub>O<sub>3</sub> supported on atomically dispersed Fe–N–C and XC72 supports, with highly localized, bulk, and surface sensitive techniques (EELS, XAS, and XPS), no spectroscopically detected interactions were observed. Critically, however, this does not rule out the possibility of electronic interactions between the Fe<sub>2</sub>O<sub>3</sub> nanoparticles and Fe–N<sub>x</sub> sites enhancing NO<sub>3</sub>RR performance. These highly sensitive interactions might require probing through electrochemical processes, in which the nanoparticle is active, while the M–N<sub>x</sub> site is inert, such that CO stripping experiments for Pt/M–N–C, allowing changes in the onset potentials or peak shapes to reflect interactions between the nanoparticle and M–N<sub>x</sub> site. Regardless, a Fe<sub>2</sub>O<sub>3</sub>/Fe–N–C active particle-active support catalyst has been synthesized and robustly charac-

terized. The Fe<sub>2</sub>O<sub>3</sub>/Fe–N–C catalyst contains a multitude of highly active NO<sub>3</sub>RR sites at both the nanoparticle and single atom scales, which are synergized, enhancing the NO<sub>3</sub>RR performance.

## 2.2. Electrochemical NO<sub>3</sub>RR performance

Linear sweep voltammetry (LSV) was performed in an alkaline, 1 M KOH + 0.16 M KNO<sub>3</sub> electrolyte to assess the NO<sub>3</sub>RR activity of the blank carbon paper, XC72, and Fe–N–C catalyst supports, and the Fe<sub>2</sub>O<sub>3</sub> nanoparticles supported on both XC72 and Fe–N–C, as shown in Figure 3a. From the LSV there is a slight positive shift in the reaction onset potential ( $-0.59$  V vs. RHE) in comparison to the blank carbon paper electrode, when using XC72, indicating that even the metal-free carbon support has some, albeit limited NO<sub>3</sub>RR performance. Note in this work all potentials are reported against the reversible hydrogen electrode, RHE. Employing the Fe–N–C catalyst support realizes a significant positive shift in the reaction onset potential ( $-0.34$  V). Interestingly, despite having increased  $j_{\text{NH}_3}$  at higher overpotentials in alkaline media, the Fe–N–C was observed in our previous work to have a higher selectivity to NH<sub>3</sub> at less cathodic potentials in neutral media (0.05 M PBS), which could be due to its hyperactivity toward reducing the NO<sub>2</sub><sup>−</sup> intermediate in the NO<sub>3</sub>RR  $2e^- + 6e^-$  transfer pathway, often formed at lower pH, which is suppressed in alkaline media.<sup>[27,28]</sup> The addition of Fe<sub>2</sub>O<sub>3</sub> nanoparticles further shifts the reaction onset potential even more positively to  $-0.14$  V, regardless of either the XC72 or Fe–N–C support. However, at more cathodic potentials, the current of Fe<sub>2</sub>O<sub>3</sub>/Fe–N–C dominates due to additional activity provided by the active Fe–N<sub>x</sub> sites in the Fe–N–C support. Figure S11, Supporting Information shows the LSV comparison of the Fe, Co, and Ru oxides supported on XC72, where RuO<sub>x</sub> shows the most positive onset potential (approximately  $+0.05$  V, however, is quickly out-competed by HER). Figures S4 and S5, Supporting Information show the LSV performance of CoO<sub>x</sub> and RuO<sub>x</sub> in electrolytes with and without KNO<sub>3</sub>. To evaluate the catalytic performance of the supports towards the NO<sub>3</sub>RR, chronoamperometry measurements were performed at potentials between  $-0.20$  and  $-1.20$  V as shown in Figure 3b. The XC72 support has negligible activity until  $-0.8$  V and reaches a maximum  $FE_{\text{NH}_3}$  of 55%. However, the Fe–N–C support demonstrates superior activity, holding a  $FE_{\text{NH}_3}$  of  $\approx 80\%$  above  $-0.20$  V, reaching a maximum of 90% at  $-0.40$  V and a maximum  $Yield_{\text{NH}_3}$  of  $2.9 \text{ mmol}_{\text{NH}_3} \text{ h}^{-1} \text{ cm}^{-2}$  (at an NH<sub>3</sub> partial current density,  $j_{\text{NH}_3} = 490 \text{ mA cm}^{-2}$  at  $-1.20$  V), surpassing other reported Fe–N–C catalysts for the NO<sub>3</sub>RR (or  $135 \text{ mA cm}^{-2}$  at  $-0.6$  V).<sup>[12,29]</sup> The addition of Fe<sub>2</sub>O<sub>3</sub> nanoparticles enhances the  $FE_{\text{NH}_3}$  (after  $-0.20$  V) and significantly improves the  $Yield_{\text{NH}_3}$  over the potential range reaching a maximum of  $6 \text{ mmol}_{\text{NH}_3} \text{ h}^{-1} \text{ cm}^{-2}$  ( $j_{\text{NH}_3} = 1265 \text{ mA cm}^{-2}$ ), Figure 3c. Furthermore, Fe<sub>2</sub>O<sub>3</sub>/Fe–N–C demonstrates increased  $FE_{\text{NH}_3}$  and  $Yield_{\text{NH}_3}$  over the potential range compared to Fe<sub>2</sub>O<sub>3</sub>/XC72. Interestingly, the Fe<sub>2</sub>O<sub>3</sub>/Fe–N–C can maintain a  $FE_{\text{NH}_3}$  of 90–95% over the potential range, highlighting the catalysts' potential independent NH<sub>3</sub> selectivity, resisting the parasitic HER even at highly cathodic potentials.

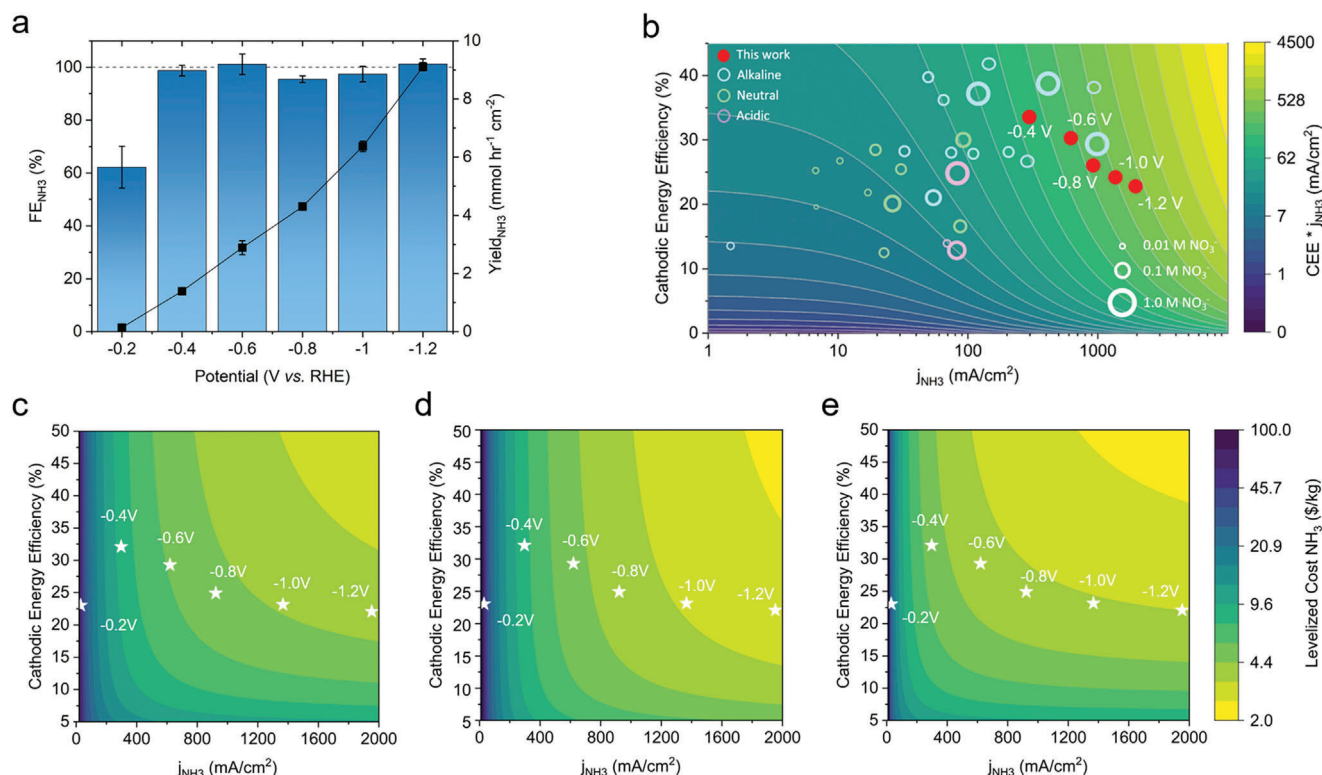
Having established the superior performance of the Fe<sub>2</sub>O<sub>3</sub>/Fe–N–C catalyst, a Fe<sub>2</sub>O<sub>3</sub> loading study was performed by adjusting



**Figure 3.** Electrochemical  $\text{NO}_3\text{RR}$  performance of  $\text{Fe}_2\text{O}_3$  based catalysts and supports in 1 M KOH + 0.16 M  $\text{KNO}_3$  electrolyte. a) Linear sweep voltammetry at a scan rate of  $5 \text{ mV s}^{-1}$ . The red line is  $\text{Fe}_2\text{O}_3/\text{XC72}$  and the green line is  $\text{Fe}_2\text{O}_3/\text{Fe-N-C}$ . Chronoamperometry measurements for 15 min at applied potentials from  $-0.20$  to  $-1.20 \text{ V}$  versus RHE. b) comparing XC72 versus Fe-N-C catalyst supports and c) comparing  $\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  versus  $\text{Fe}_2\text{O}_3/\text{XC72}$ . d) TEM images of increasing  $\text{Fe}_2\text{O}_3$  loadings supported on Fe-N-C, the scale bar is 20 nm. Electrochemical performance of  $\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  with varying  $\text{Fe}_2\text{O}_3$  loadings, e) Linear sweep voltammetry at a scan rate of  $5 \text{ mV s}^{-1}$  and f) Chronoamperometry measurements for 15 min at applied potentials from  $-0.20$  to  $-1.20 \text{ V}$  versus RHE with a catalyst loading on the carbon paper of  $0.2 \text{ mg cm}^{-2}$ . The corresponding chronoamperometry measurements and UV-vis  $\text{NH}_3$  quantification are given in Figures S20–S25, Supporting Information. Note that all samples for the  $\text{NO}_3\text{RR}$  performance in this figure have undergone a pre-reduction activation step, the impact of which is discussed in detail later in the manuscript.

the  $\text{Fe}(\text{acac})_3$  loading, to further enhance the  $\text{NO}_3\text{RR}$  activity. The TEM images in Figure 3d show the  $\text{Fe}(\text{acac})_3$  loading impact on the  $\text{Fe}_2\text{O}_3$  site density and gradual formation of agglomerates. With the standard  $\text{Fe}_2\text{O}_3$  (e.g.,  $1 \times \text{Fe}(\text{acac})_3$ ), a relatively low  $\text{Fe}_2\text{O}_3$  site density is observed with no agglomerates. The optimal loading appears to be at  $3 \times \text{Fe}_2\text{O}_3$ , at this loading, the  $\text{Fe}_2\text{O}_3$  site density significantly increases, with slight agglomerate formation beginning, while at  $4 \times \text{Fe}_2\text{O}_3$ , the agglomerated features dominate, reducing the catalytically active surface area. Figures S12–S15, Supporting Information shows additional TEM images for the  $\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  at loadings between  $1 \times$  –  $4 \times \text{Fe}(\text{acac})_3$ . Thermogravimetric (TGA) analysis, Figure S16, Supporting Information determined a Fe weight loading of 42% for the  $3 \times \text{Fe}_2\text{O}_3$  catalyst and 22% for the  $1 \times \text{Fe}_2\text{O}_3$ , indicating a non-linear increase in the Fe content with precursor loading. The optimal  $3 \times \text{Fe}_2\text{O}_3$  loading is directly observed in the  $\text{NO}_3\text{RR}$  performance and calculated ECSA, Figure 3e and Figure S17, Supporting Information, respectively. Furthermore,  $3 \times \text{Fe}_2\text{O}_3/\text{Fe-N-C}$  demonstrates the highest  $FE_{\text{NH}_3}$  maintaining  $\approx 95$ – $100\%$  from  $-0.40$  to  $-1.20 \text{ V}$ , and the highest  $\text{Yield}_{\text{NH}_3}$  at all potentials, reaching a maximum of  $8.3 \text{ mmol}_{\text{NH}_3} \text{ h}^{-1} \text{ cm}^{-2}$  ( $j_{\text{NH}_3} = 1785 \text{ mA cm}^{-2}$ ). To further optimize the catalyst performance and increase  $j_{\text{NH}_3}$ , the optimal catalyst loading on the carbon paper was investigated. The ink volume of  $3 \times \text{Fe}_2\text{O}_3/\text{Fe-N-C}$  drop cast on the carbon paper electrode was varied to achieve a final

catalyst loading between  $0.2$ – $10.0 \text{ mg cm}^{-2}$ . LSV was performed to screen the activity of the different catalyst loadings, shown in Figure S18, Supporting Information. As the catalyst loading increased from  $0.2$ – $1.0 \text{ mg cm}^{-2}$ , the maximum current increased, and the reaction onset potential shifted positively, with no improvement being observed between  $0.5$ – $1.0 \text{ mg cm}^{-2}$ . However, after  $1.0 \text{ mg cm}^{-2}$  the catalyst layer becomes too thick, inhibiting the optimal use of the porous carbon paper, resulting in reduced activity, with  $10.0 \text{ mg cm}^{-2}$  giving the lowest performance. As an activity comparison, Figure S18c, Supporting Information shows the reaction onset potential and maximum current at  $-0.5 \text{ V}$  (the maximum potential at which a cathodic energy efficiency of 30% is achieved assuming 100%  $FE_{\text{NH}_3}$ ). The optimal catalyst loading is determined to be  $0.5 \text{ mg cm}^{-2}$ , giving an onset potential of approximately  $-0.13 \text{ V}$  and a current density of  $520 \text{ mA cm}^{-2}$  at  $-0.50 \text{ V}$ . Therefore, the optimal catalyst is the  $3 \times \text{Fe}_2\text{O}_3/\text{Fe-N-C}$  with a loading of  $0.5 \text{ mg cm}^{-2}$ , and the electrochemical results discussed further in this manuscript utilize these conditions. Finally, to confirm that the source of  $\text{NH}_3$  originates from the  $\text{NO}_3$  in the electrolyte, rather than contamination or decomposition of the N-doped carbon support, a series of control experiments were performed. Electrolysis from  $-0.2$  to  $-1.2 \text{ V}$  for 15 min each was performed in 1 M KOH electrolyte (without  $\text{NO}_3^-$ ) and tested by UV-vis, showing no detected  $\text{NH}_3$  in the absence of  $\text{NO}_3^-$ , Figure S19a,b, Supporting Information. Next, isotopically



**Figure 4.** Electrochemical  $\text{NO}_3\text{RR}$  performance of the optimized  $3\times\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  catalyst with a  $0.5\text{ mg cm}^{-2}$  catalyst loading on the carbon paper electrode in a  $1\text{ M KOH} + 0.16\text{ M KNO}_3$  electrolyte. a) Optimized  $FE_{\text{NH}_3}$  and  $Yield_{\text{NH}_3}$  as a function of applied potential. b) Comparison of  $\text{NO}_3\text{RR}$  performance in the current literature, evaluating the cathodic energy efficiency versus  $j_{\text{NH}_3}$ . Full details and references for each reported  $\text{NO}_3\text{RR}$  system are provided in Table S1, Supporting Information. Contour plots evaluating the trade-off of cathodic energy efficiency versus  $j_{\text{NH}_3}$  on the levelized cost of  $\text{NH}_3$  for cases with c) grid electricity price, d) idealized cost of renewable energy, and e) reduced electrolyzer stack cost. Where performance improves across contours towards the top right corner.

labeled experiments were performed using a  $1\text{ M KOH} + 0.16\text{ M }^{15}\text{KNO}_3$  electrolyte. Electrolysis was performed at  $-1.0\text{ V}$  for  $15\text{ min}$ , and  $^1\text{H NMR}$  quantified the  $^{15}\text{NH}_3$  produced. The isotopic measurements (Figure S19c,d, Supporting Information) show a strong comparison between the non-isotopically doped experiments, both at a  $FE_{\text{NH}_3} \approx 100\%$  and a yield rate of  $6.5\text{ mmol h}^{-1}\text{ cm}^{-2}$  ( $^{15}\text{KNO}_3$ ) and  $6.2\text{ mmol h}^{-1}\text{ cm}^{-2}$  ( $^{14}\text{KNO}_3$ ), confirming that any  $\text{NH}_3$  detected results from the activation of  $\text{NO}_3^-$ .

Figure 4a, shows the  $\text{NO}_3\text{RR}$  performance of the optimized  $3\times\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  catalyst (with a  $0.5\text{ mg cm}^{-2}$  loading on the working electrode), maintaining a  $FE_{\text{NH}_3}$  above  $95\%$  over the entire  $-0.40$  to  $-1.20\text{ V}$  potential range, again highlighting its potential independent nature towards  $\text{NH}_3$  selectivity. This potential independent behavior on  $\text{NH}_3$  selectivity provides an advantage in practical systems. When coupled with renewable energy, dynamic changes in the supplied energy can lead to cell voltage fluctuations. With the current system, even with cell voltage fluctuations, the product purity would remain unchanged. A maximum  $Yield_{\text{NH}_3}$  of  $9.2\text{ mmol h}^{-1}\text{ cm}^{-2}$  is achieved ( $j_{\text{NH}_3} = 1950\text{ mA cm}^{-2}$ ) at  $\approx 100\% FE_{\text{NH}_3}$ . To compare the performance of the  $3\times\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  catalyst to the current  $\text{NO}_3\text{RR}$  literature, Figure 4b compares the cathodic energy efficiency, CEE (assuming the thermodynamic reduction potential for the anodic oxygen evolution reaction) versus the  $j_{\text{NH}_3}$ . The CEE is a function of the  $FE_{\text{NH}_3}$  and applied potential (taking a penalty for highly ca-

thodic potentials), while the  $j_{\text{NH}_3}$  is a function of the  $FE_{\text{NH}_3}$  and total current, enabling a comparison beyond just the  $FE_{\text{NH}_3}$  or  $Yield_{\text{NH}_3}$ , which can vary significantly based on the applied potential. Larger circles indicate higher concentrations of  $\text{NO}_3^-$ , often resulting in better performance, while colors are used to designate acidic, neutral, or alkaline media. The contours in Figure 4b are the product of  $(CEE \times j_{\text{NH}_3})$ , yielding a performance metric in terms of  $\text{mA cm}^{-2}$ , which is optimized across contours and towards the top right. From Figure 4b, it is apparent that universally, the  $\text{NO}_3\text{RR}$  suffers from relatively low energy efficiencies at meaningful  $j_{\text{NH}_3}$  (above  $100\text{ mA cm}^{-2}$ ), due to the thermodynamic reaction onset potential ( $0.69\text{ V vs RHE}$ ,  $\text{pH} = 14$ ), while more cathodic potentials (approximately  $-0.40$  to  $-0.80\text{ V}$ ) are typically required to achieve large  $j_{\text{NH}_3}$ . The majority of the  $\text{NO}_3\text{RR}$  performances in the literature report limited  $j_{\text{NH}_3}$ , less than  $100\text{ mA cm}^{-2}$ , or utilize expensive PGM metals to realize higher  $j_{\text{NH}_3}$ , hindering industrial relevance. The  $3\times\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  catalyst (red circles) enables high current densities even at mildly reductive potentials,  $297\text{ mA cm}^{-2}$  with a cathodic energy efficiency of  $33\%$ . The potential independent nature of the  $\text{NH}_3$  selectivity, allows the  $3\times\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  system to be operated between  $-0.40$  to  $-1.20\text{ V}$  at nearly  $100\% FE_{\text{NH}_3}$ , giving ultra-high  $j_{\text{NH}_3}$  from  $297$  to  $1950\text{ mA cm}^{-2}$ , outperforming the current  $\text{NO}_3\text{RR}$  literature (see Figure S26, Supporting Information for a linear  $j_{\text{NH}_3}$  scale).



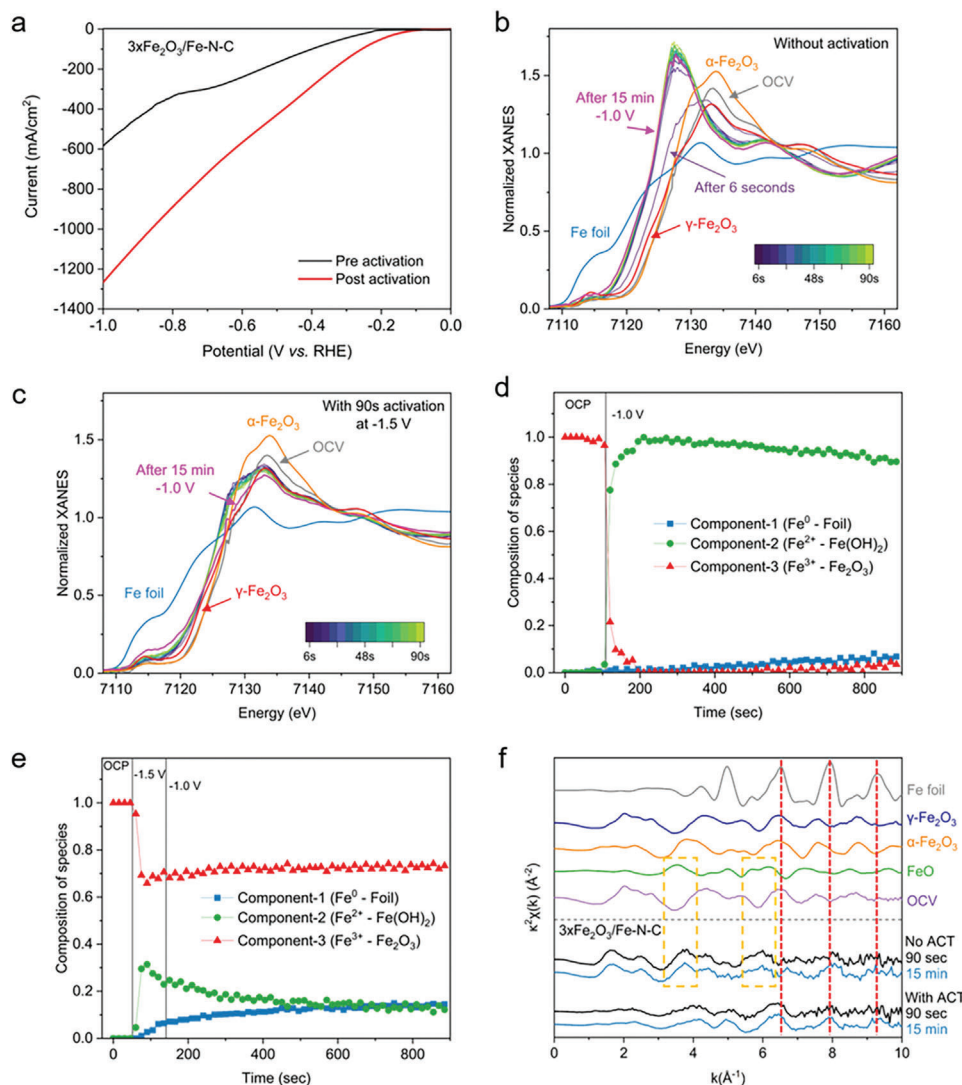
There is significant ambiguity in determining the optimal cathodic potential to yield both, an acceptable CEE and  $j_{\text{NH}_3}$  and depends on many factors including catalyst cost, device costs, CAPEX and OPEX costs, the levelized cost of  $\text{NH}_3$  as  $j_{\text{NH}_3}$  increases, and many others. To offer a semi-quantitative optimal trade-off between energy efficiency and  $j_{\text{NH}_3}$ , Figure 4c–e offers a simplified economic analysis inspired by a recent work from Daiyan et al.<sup>[22]</sup> To construct meaningful contours, the levelized cost of  $\text{NH}_3$  ( $\text{\$ kg}^{-1}$ ) was determined as a function of the CAPEX, OPEX, and yield rates of  $\text{NH}_3$ . These inputs consider the increasing OPEX as the cathodic potential increases, therefore decreasing energy efficiency, while also accounting for increased production rates of  $\text{NH}_3$  (see Experimental Section for details). Three scenarios are considered, where the electricity cost varies from standard grid electricity at  $\text{\$0.07 kWh}^{-1}$  (Figure 4c), idealized renewable energy from solar power at  $\text{\$0.03 kWh}^{-1}$  (Figure 4d), and with a decreased CAPEX resulting from reduced electrolyzer stack costs (Figure 4e). From these contour plots, it is readily observed that the most effective way to cross contours (until  $\approx 1 \text{ A cm}^{-2}$ ) is through increasing the  $j_{\text{NH}_3}$ , rather than achieving low  $j_{\text{NH}_3}$  with increasing energy efficiency. This is observed for both electricity price scenarios, the lowest levelized cost of  $\text{NH}_3$  is achieved at the most cathodic potential of  $-1.2 \text{ V}$ , where the ultrahigh current density ( $j_{\text{NH}_3}$  of  $-1.95 \text{ A cm}^{-2}$ ) is achieved, despite the lower energy efficiency. Interestingly, the impact of energy efficiency has a minor effect at small/moderate  $j_{\text{NH}_3}$  and becomes more effective at higher  $j_{\text{NH}_3}$ .

### 2.3. In Situ Evaluation of Fe Speciation during Pre-Reduction Activation and Electrolysis

Prior to the  $\text{NO}_3\text{RR}$  measurements, a pre-reduction activation step was performed and found to significantly improve the activity of the catalyst, as demonstrated through the LSV in Figure 5a. The pre-reduction activation applies a highly reductive potential of  $-1.5 \text{ V}$  versus RHE for 90 s. To elucidate the chemical state changes of the pristine  $3\times\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  to the now highly active catalyst, in situ quick XAFS was performed and complemented by post-mortem XPS. Note that a milder electrolyte of  $0.1 \text{ M KOH} + 0.016 \text{ M KNO}_3$  was used for the in situ experiments in order to preserve the integrity of the Kapon used in the operando spectroscopy cell. Figure 5b,c show the XANES spectra of the Fe K-edge under  $\text{NO}_3\text{RR}$  conditions for the  $3\times\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  catalyst, without and with the pre-reduction activation step, respectively. Prior to any electrochemical measurements, an initial spectrum was taken at OCV, indicated by the gray curve. Figure S27, Supporting Information demonstrates that there are no significant changes between the air measurements and OCV samples, indicating no changes in the Fe chemical state prior to the applied potential. In Figure 5b, for the sample without the activation step, a potential of  $-1.0 \text{ V}$  was directly applied, and Fe K-edge XAS spectra were recorded every 3 s (with every two spectra being averaged for increased quality), for the first 90 s. Immediately, there is a distinct shift in the absorption edge toward lower energy, with a simultaneous decrease in the pre-edge feature  $\approx 7114 \text{ eV}$  and a significant increase in the intensity of the white line. A final spectrum was acquired after 15 min of a potential hold at  $-1.0 \text{ V}$  (analogous to the  $\text{NO}_3\text{RR}$  experiments dis-

cussed earlier), which demonstrates that changes in the Fe chemical state after the initial 90 s are relatively minor. By comparing these spectra with the Fe-reference materials, it is concluded that in the absence of a pre-reduction activation step, the  $\text{NO}_3\text{RR}$  conditions induce a transformation of  $\text{Fe}^{3+}$  to  $\text{Fe}^{2+}$ . In particular, the final in situ spectrum resembles strongly the  $\text{Fe}(\text{OH})_2$  spectrum.<sup>[35]</sup> In contrast, Figure 5c shows the evolution of Fe K-edge XAS for the catalyst, where the 90 s pre-reduction activation step at  $-1.5 \text{ V}$  has been performed. In this scenario, the changes in Fe K-edge XANES during the first 90 s are remarkably different. The white line intensity decrease is accompanied by a shift in the edge position towards lower energy and an increase in the pre-edge feature  $\approx 7114 \text{ eV}$ . After the 90 s of activation, a  $-1.0 \text{ V}$  potential was applied (replicating the  $\text{NO}_3\text{RR}$  tests, in which an initial 90-s activation at  $-1.5 \text{ V}$  is followed by  $-1.0 \text{ V}$  for 15 min). The XAS spectra collected during this latter stage show a further reduction in the white line intensity and an increase in the pre-edge feature, likely associated with the further reduction of  $\text{Fe}^{3+}$  (or intermediate  $\text{Fe}^{2+}$ ) and the formation of metallic Fe ( $\text{Fe}^0$ ). These results show clearly that the evolution of the chemical state of the Fe is very different in cases with and without the activation step (Figure S28, Supporting Information).

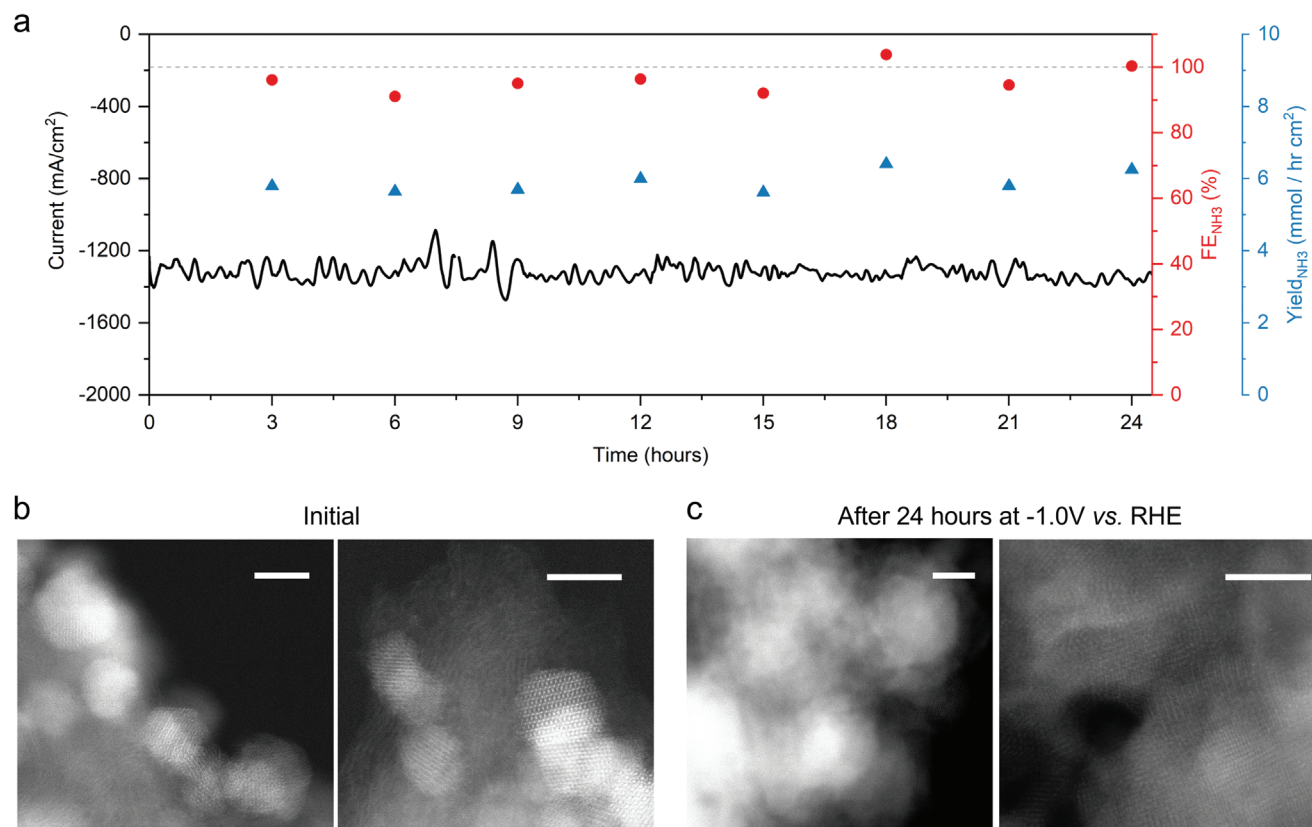
To quantitatively analyze the Fe speciation observed in the in situ quick XAFS experiments, principal component analysis (PCA) and multivariate curve resolution (MCR) are employed and complemented by the EXAFS fitting.<sup>[36–39]</sup> PCA and MCR methods were applied to a combined dataset, consisting of spectra collected in the experiment with and without the activation step. First, PCA identified three spectroscopically distinct species in this combined dataset, as determined by a Scree plot (Figure S29, Supporting Information), showing the relative importance of the principal components. Next, MCR analysis based on the alternating least squares (MCR-ALS) method was used to determine the spectra corresponding to these three species, and the corresponding concentration profiles. For this purpose, a MATLAB code developed by Jaumont et al. was employed.<sup>[40]</sup> A successful convergence was achieved after eight iterations. The details of the fits are shown in Table S2, Supporting Information and discussed in Note S1, Supporting Information. The three spectral components identified by the MCR-ALS method are shown in Figure S30, Supporting Information. Spectral component 1 aligns well with the spectrum of metallic Fe in the  $\text{Fe}^0$  state. Spectral component 2 resembles the spectrum for  $\text{Fe}(\text{OH})_2$ , and thus, can be associated with the  $\text{Fe}^{2+}$  state. Finally, spectral component 3 matches well with the  $3\times\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  sample and the spectrum of metallic Fe in the  $\text{Fe}^0$  state. Figure 5d shows the evolution of the concentration of these identified Fe species over the 15 min  $\text{NO}_3\text{RR}$  electrolysis at  $-1.0 \text{ V}$ , for the  $3\times\text{Fe}_2\text{O}_3/\text{Fe-N-C}$  without the pre-reduction activation. Figure 5d shows that immediately after the  $-1.0 \text{ V}$  is applied, nearly all the  $\text{Fe}^{3+}$  is converted to  $\text{Fe}^{2+}$ , likely in the form of  $\text{Fe}(\text{OH})_2$ . In the following 15 min of the experiment, some  $\text{Fe}^{2+}$  is further reduced to  $\text{Fe}^0$ , but at a very slow rate. In contrast, Figure 5e shows the concentration profiles of Fe species obtained for the case with a pre-reduction activation step. During the first 25 s of the 90 s potential hold at  $-1.5 \text{ V}$ , the conversion from  $\text{Fe}^{3+}$  to  $\text{Fe}^{2+}$  is paralleled by the formation of metallic  $\text{Fe}^0$ . During the 15 min electrolysis at  $-1.0 \text{ V}$ , the remaining  $\text{Fe}^{2+}$  is further converted to  $\text{Fe}^0$  until H550 s, at which point the concentration of  $\text{Fe}^{3+}$ ,  $\text{Fe}^{2+}$ ,



**Figure 5.** In situ investigation of the electronic state of Fe during the pre-reduction activation step and NO<sub>3</sub>RR electrolysis of the 3xFe<sub>2</sub>O<sub>3</sub>/Fe-N-C. a) LSV in 1 M KOH + 0.16 M KNO<sub>3</sub> demonstrating the increased activity from the pre-reduction activation step. b) XANES spectra of the catalyst in the first 90 s of the NO<sub>3</sub>RR at -1.0 V and after 15 min, for the catalyst b) without the pre-reduction activation step and c) with the pre-reduction activation step. Fe speciation over 15 min of NO<sub>3</sub>RR electrolysis at -1.0 V as determined from MCR-ALS analysis of the in situ XANES measurements for the catalyst d) without the pre-reduction activation step and e) with the pre-reduction activation step. f) EXAFS spectra in *k*-space analysis of the in situ measurements, where the EXAFS features related to Fe oxide structures are marked by orange dashed rectangles and features related to metallic Fe are marked by red dashed lines.

and Fe<sup>0</sup> approaches the steady state. Thus, at the highly cathodic potential of -1.5 V, the reduction of Fe<sup>3+</sup> to Fe<sup>0</sup> is triggered, at which point the further reduction of Fe<sup>3+</sup> is hindered. These results agree well with previously reported literature demonstrating that during a cathodic bias of -1.0 V, Fe oxides supported on nitrogen-doped carbon were not fully reduced to Fe<sup>0</sup>, which was attributed to phase contractions and the insertion of H<sub>2</sub>O in the lattice, maintaining a Fe(OH)<sub>2</sub> structure (Fe<sup>2+</sup>), despite being 560 mV lower than thermodynamically expected potential for Fe<sup>0</sup>.<sup>[41]</sup> Additionally, it has been shown that nitrogen dopants can stabilize Fe<sup>2+</sup> species, preventing the complete reduction to Fe<sup>0</sup> under a cathodic bias.<sup>[42]</sup> The in situ EXAFS data in *k*-space shown in Figure 5f agrees with the XANES analysis, showing the

clear formation of Fe<sup>0</sup> during the pre-reduction activation step and further formation during the 15 min electrolysis, resembling the Fe foil reference. For the sample without the activation step, during the electrolysis, the characteristic Fe<sup>0</sup> high-frequency oscillations at larger *k*-values are also present, however, they are much less pronounced than in the case with the pre-reduction activation step. Further analysis of the EXAFS spectra in Figure S31, Supporting Information shows that in the experiments with and without the activation step, a Fe-Fe bond contribution is observed after 15 min of applied potential. However, for the sample with the pre-reduction activation, the Fe-Fe coordination numbers are significantly larger ( $6.2 \pm 0.5$  vs  $1.2 \pm 0.5$ ), and the Fe-O bond distances are shorter (1.97 vs 2.12 Å), suggesting a



**Figure 6.** Durability study of the 3xFe<sub>2</sub>O<sub>3</sub>/Fe–N–C catalyst. a) 24-h electrolysis at –1.0 V in 1 M KOH + 0.16 M KNO<sub>3</sub> electrolyte. HAADF-STEM images investigating the stability of the Fe<sub>2</sub>O<sub>3</sub> nanoparticles b) for the pristine catalyst and c) after the 24-h electrolysis. All scale bars are 5 nm.

larger fraction of Fe<sup>3+</sup> (e.g.,  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> – 1.97 Å) as compared to Fe<sup>2+</sup> (e.g., FeOOH – 2.08 Å). Thus, EXAFS data analysis further confirms the conclusions from the XANES analysis that the activated catalyst contains a significant amount of Fe<sup>3+</sup> species coexisting with Fe<sup>0</sup>, while in the non-activated catalyst mainly Fe<sup>2+</sup> species exist with significantly smaller amounts of Fe<sup>0</sup> (EXAFS fitting parameters given in Tables S3–S5, Supporting Information).

Our XAS analysis shows that even at the highly cathodic potential of –1.5 V, the complete reduction of Fe<sup>3+</sup> to Fe<sup>0</sup> is not observed, with a significant portion of Fe<sup>3+</sup> appearing to be stabilized with the formation of Fe<sup>0</sup>/Fe<sup>2+</sup>. Furthermore, the formation and preservation of the Fe<sup>0</sup> species correspond to a significantly enhanced NO<sub>3</sub>RR performance, which is in agreement with previous works demonstrating the effectiveness of lower oxidation state Fe species towards the NO<sub>3</sub>RR.<sup>[43,44]</sup> The maintained Fe speciation and boosted activity are supported by a 24 h NO<sub>3</sub>RR electrolysis at –1.0 V, following a pre-reduction activation step. To circumvent ultrahigh NH<sub>3</sub> concentrations in the electrolyte and subsequent loss of NH<sub>3</sub> in the gas phase, the electrolysis was performed in eight 3-h segments. Furthermore, the system was modified such that peristaltic pumps were connected to an external reservoir to the working chamber, enabling a working electrolyte volume of 250 mL, with constant circulation. After a 3-h segment, the electrolyte was sampled and refreshed. **Figure 6a** shows a stable current density of approximately –1.3 A cm<sup>–2</sup> at a FE<sub>NH3</sub> between 90–100% and Yield<sub>NH3</sub> of ≈6 mmol h<sup>–1</sup> cm<sup>–2</sup>. The

stable performance over 24 h suggests that no further changes in Fe speciation are occurring, altering catalyst activity or selectivity to NH<sub>3</sub>. Further highlighting the ability of the 3xFe<sub>2</sub>O<sub>3</sub>/Fe–N–C catalyst to preserve this ultrahigh NO<sub>3</sub>RR performance at a near 100% FE<sub>NH3</sub>, resisting the HER even at highly reductive potentials.

To complement the in situ quick XAFS measurements, post-mortem XPS analysis was performed on the working electrode following the pre-reduction activation step and after the 24-h electrolysis at –1.0 V. Following the pre-reduction activation step and 24-h electrolysis, the working electrode was dried under N<sub>2</sub> and stored in a gas-tight vial pre-purged with N<sub>2</sub> for immediate transport to the XPS. To address possible slight re-oxidation of the surface Fe during the transport of the electrode to the XPS, spectra were taken followed by quick (60 s) Ar<sup>+</sup> ion surface etch and re-sampled (see Note S2, Supporting Information for details, considerations, and oxidation consequences of the Ar<sup>+</sup> etching). Although not as rigorous as in situ quick XAFS, the deconvoluted post-mortem XPS spectra in Figures S32–S34, Supporting Information, although more surface sensitive, show a clear qualitative agreement with the XANES measurements, showing the formation of Fe<sup>2+</sup>, Fe(OH)<sub>2</sub> and Fe<sup>0</sup> during the pre-reduction activation step. Throughout the 24-h electrolysis, post-mortem XPS shows only a slight further reduction of Fe<sup>3+</sup>, in agreement with the in situ quick XAFS (first 15 min) and increased formation of Fe<sup>0</sup>. Following the 24-h electrolysis, to investigate changes in the Fe<sub>2</sub>O<sub>3</sub> nanoparticle structure and atomically dispersed Fe



sites, atomic resolution STEM was performed, Figure 6b,c (and Figure S35, Supporting Information). A slight coarsening in the  $\text{Fe}_2\text{O}_3$  nanoparticles is observed ( $\approx 10$  nm), however, the minor physical change in the catalyst structure does not negatively alter the  $\text{NO}_3\text{RR}$  performance as observed in Figure 6c. Additionally, at higher magnification, the coexistence of  $\text{Fe}_2\text{O}_3$  nanoparticles and atomically dispersed Fe sites are maintained (larger images shown in Figure S36, Supporting Information), confirming the durability of both the active  $\text{Fe}_2\text{O}_3$  nanoparticle catalyst and active Fe–N–C support at highly reductive potentials. As a comparison, a 24-h stability test at  $-1.0$  V was performed on the  $\text{Fe}_2\text{O}_3/\text{XC72}$  catalyst, shown in Figure S37, Supporting Information. Without the Fe–N<sub>x</sub> sites, the stability of the  $\text{Fe}_2\text{O}_3$  particles is reduced, leading to the formation of large agglomerated structures (rather than slightly coarsened particles as in the case of  $\text{Fe}_2\text{O}_3/\text{Fe–N–C}$ ). For better visualization of the atomically dispersed sites, a Fe–N–C catalyst without  $\text{Fe}_2\text{O}_3$  nanoparticles after electrolysis (24 h at  $-1.0$  V) was imaged in Figure S38, Supporting Information. These results are supported by other studies in the literature showing high stability of the Fe–N–C sites under reductive potentials.<sup>[12,29,45]</sup> Although it is critical to note that these are ex situ measurements with the sample being exposed to air prior to imaging, which can in some cases enable the re-dispersion of single atoms agglomerated during electrolysis back to their atomically dispersed state.

From the chemical state analysis, it is hypothesized that during the pre-reduction activation step, surface  $\text{Fe}^{3+}$  species are reduced to highly active  $\text{Fe}^0$ , which is preserved throughout the  $\text{NO}_3\text{RR}$  electrolysis maintaining the high  $\text{NO}_3\text{RR}$  activity and selectivity to  $\text{NH}_3$  (Figure 6a). Based on this analysis,  $\text{Fe}^{3+}/\text{Fe}^{2+}$  sites while active for the  $\text{NO}_3\text{RR}$  are less active than  $\text{Fe}^0$  sites. Therefore, to maximize the  $\text{NO}_3\text{RR}$  performance, a pre-reduction step to reduce  $\text{Fe}^{3+}$  sites to  $\text{Fe}^0$  is essential to obtain enhanced  $\text{NO}_3\text{RR}$  performance.

## 2.4. Conclusion

In summary, we have leveraged the high activity of Fe-based catalysts, both at the nanoparticle and single atom level, towards the selective conversion of  $\text{NO}_3^-$  to  $\text{NH}_3$ . We synergized both, the nano and atomic scales to synthesize an active particle-active support catalyst system,  $\text{Fe}_2\text{O}_3/\text{Fe–N–C}$ . The atomically dispersed Fe–N<sub>x</sub> sites of the active Fe–N–C support, mixed  $\gamma/\alpha$  Fe-phase, and spinel structure of the  $\text{Fe}_2\text{O}_3$  nanoparticles were robustly confirmed employing atomic resolution STEM and EELS, XAS, and XPS. The optimized  $3\times\text{Fe}_2\text{O}_3/\text{Fe–N–C}$  catalyst demonstrated potential independent  $\text{NO}_3\text{RR}$  activity, even at highly reductive potentials, reaching a high  $\text{Yield}_{\text{NH}_3}$  of over  $9\text{ mmol h}^{-1}\text{ cm}^{-2}$  at a  $FE_{\text{NH}_3}$  of 100%, and a  $j_{\text{NH}_3}$  up to  $1.95\text{ A cm}^{-2}$  (at  $-1.2$  V vs RHE). In situ XANES and post-mortem XPS revealed the partial reduction of  $\text{Fe}^{3+}$  surface sites to highly active  $\text{Fe}^0$  during the pre-reduction activation, which is maintained throughout the  $\text{NO}_3\text{RR}$  electrolysis and is critical in boosting the  $\text{NO}_3\text{RR}$  performance. A durability study at  $-1.0$  V over 24 h demonstrated the robustness of the  $3\times\text{Fe}_2\text{O}_3/\text{Fe–N–C}$  catalyst, preserving the highly active  $\text{Fe}^0$  sites, maintaining a current of  $1.3\text{ A cm}^{-2}$  and a  $FE_{\text{NH}_3}$  of 91–100%. This work introduces a novel active particle-active support catalyst system for the  $\text{NO}_3\text{RR}$ , utilizing a plurality of active sites

at both the nanoparticle and single atom scale, to significantly enhance  $\text{NO}_3\text{RR}$  activity. While elucidating the importance of a pre-reduction activation step to create highly active surface  $\text{Fe}^0$  species, capable of realizing  $\text{NO}_3^-$  reduction to  $\text{NH}_3$  at industrially relevant current densities ( $1.95\text{ A cm}^{-2}$ ) and near unity  $FE_{\text{NH}_3}$ .

## 3. Experimental Section

**Synthesis of  $\text{Fe}_2\text{O}_3$ ,  $\text{Co}_3\text{O}_4$ , and  $\text{RuO}_x$  Supported on XC72:** The Fe, Co, and Ru nanoparticle catalysts were synthesized using an organic solvent synthesis, employing an oleic acid and oleylamine ligand pair, commonly employed in nanoparticle synthesis to control the size and dispersion of the nanoparticles. Taking  $\text{Fe}_2\text{O}_3$  as an example, first 44.8 mg of XC72 carbon and 0.2 mmol of  $\text{Fe}(\text{acac})_3$  were dispersed by sonication for 30 min in 40 mL of benzyl ether. Next, the mixture was deaerated by purging  $\text{N}_2$  for 30 min. The mixture was then heated to  $100^\circ\text{C}$  and then 400  $\mu\text{L}$  of oleylamine and 200  $\mu\text{L}$  of oleic acid were added and the temperature was held for 10 min. Next, the mixture was heated to  $180^\circ\text{C}$  and 1 mL of lithium triethylborohydride was added and the temperature was held for 10 min. The catalyst mixture was then heated to  $210^\circ\text{C}$  and held for 45 min. The catalyst mixture was then centrifuged with ethanol three times before filtration with ethanol, followed by drying at  $45^\circ\text{C}$  in an oven.

The Co and Ru-based particle syntheses were identical, with the 0.2 mmol of  $\text{Co}(\text{acac})_3$  and  $\text{Ru}(\text{acac})_3$  being added in place of  $\text{Fe}(\text{acac})_3$ .

**Synthesis of Fe–N–C:** The atomically dispersed Fe–N–C active support was synthesized using the sacrificial support method (SSM). First, a catalyst mixture of 6.25 g of nicarbazin, 1.25 g of OX-50 (Evonik), 1.25 g of LM150 (Cabot), 0.5 g of stöber spheres (made in-house), and 0.6 g of iron (III) nitrate was added and dispersed by sonication for 30 min in water. The catalyst slurry was then dried for 24 h at  $45^\circ\text{C}$  under continuous stirring. The partially dried slurry was then transferred to an oven for 24 h for complete drying at  $45^\circ\text{C}$ . The catalyst mixture was then ball-milled at 45 Hz for 60 min. Next, the milled catalyst powder underwent pyrolysis in a 5%  $\text{H}_2$  / 95% Ar atmosphere for 45 min at  $975^\circ\text{C}$ . The pyrolyzed catalyst was then ball-milled a second time at 45 Hz for 1 h before being etched in a concentrated HF (18 M) solution for 96 h to remove the silica support and any nanoparticles. The etched catalyst was then washed with DI water and filtered until neutral pH before drying. A second pyrolysis under a 10%  $\text{NH}_3$  / 90%  $\text{N}_2$  atmosphere was performed at  $950^\circ\text{C}$  for 30 min. The catalyst was then ball-milled a third time at 45 Hz for 1 h.

**Synthesis of  $\text{Fe}_2\text{O}_3/\text{Fe–N–C}$ :** The  $\text{Fe}_2\text{O}_3$  nanoparticles supported on atomically dispersed Fe–N–C ( $\text{Fe}_2\text{O}_3/\text{Fe–N–C}$ ) were synthesized analogously to the nanoparticle catalyst supported on XC72, with the carbon support being switched for the active Fe–N–C support.

**Physical Characterization:** Transmission electron microscopy (TEM) was performed on a JEOL JEM-2100F. To obtain atomic resolution images, aberration-corrected scanning transmission electron microscopy (AC-STEM) and energy dispersive X-ray spectroscopy (EDX) were performed on a JEOL ARM300CF (at 300 keV accelerating voltage). The valence state of the  $\gamma\text{-Fe}_2\text{O}_3$  nanoparticles and atomically dispersed Fe sites were examined through atomic resolution electron energy loss spectroscopy (EELS) on a Nion UltraSTEM200 microscope equipped with a cold FEG, a C3/C5 aberration correction, and a high-energy resolution monochromated EELS system (HERMES). To suppress beam damage, a lower accelerating voltage of 60 keV was used to collect the EELS spectra. For the spectra acquisition, the energy dispersion was set as 0.29 eV per channel at an exposure time of 500 ms per pixel. Background subtraction in the spectrum was achieved by a power-law function and the de-noising of the spectra was performed by the multivariate weighted principal component analysis (PCA) routine in the Digital Micrograph software. The smoothing of the spectra was achieved by a Savitzky-Golay method with points of window of 15 using the Origin software. For energy loss near edge structure (ELNES) analysis on Fe valence state, the spectra collected from different Fe SA locations were summed up and then averaged to improve the signal-to-noise ratio.

The surface valence and chemical bonds of the catalysts were analyzed by XPS performed using a Kratos AXIS Supra spectrometer with a monochromatic Al K $\alpha$  source (1486.6 eV). A pass energy of 160 eV from 1400 to 5 eV at a step size of 1 eV was used to obtain the survey spectra. No charge neutralization was employed. CasaXPS software was used to analyze the XPS data with the spectrum being calibrated by C 1s sp<sup>2</sup> peak at (284.8 eV). For analyzing the data, two backgrounds were used, with a linear background being employed for the C 1s and N 1s spectrum, while a Shirley background was used for the N 1s and Fe 2p spectrum. For analysis of the sp<sup>2</sup> carbon, an asymmetric 50% Gaussian / 50% Lorentzian was applied. While for all other data, a 70% Gaussian / 30% Lorentzian was applied. For the Ar<sup>+</sup> ion etching experiments, a survey was first taken, followed by an etch and another survey, and continued in fashion. Ar<sup>+</sup> ions with an energy of 5 keV were used to etch a 2 × 1 mm area for 60 s per etch cycle.

To examine the crystal phase of the catalysts, X-ray diffraction (XRD) patterns were obtained using a Rigaku Ultima-III powder X-ray diffractometer. The iron metal content of the catalysts was quantified by thermogravimetric analysis (TGA) performed on a Netzsch TG 209 F1 Libra. To quantify the graphitic and amorphous content in the two catalyst supports (XC72 and Fe–N–C), Raman spectra were taken on an InVia, Renishaw Corp., UK system.

The ex situ XAS measurements for the Fe<sub>2</sub>O<sub>3</sub>/Fe–N–C and Fe<sub>2</sub>O<sub>3</sub>/XC72 catalysts were performed on the SAMBA beamline at the SOLEIL Synchrotron Radiation Facility, Paris, France. The sample was measured in fluorescence mode and references in transmission mode using a Si (220) monochromator for the energy selection. Ionization chambers to measure the X-ray intensity before and after the sample were filled with a mixture of Ar/N<sub>2</sub> (10) or pure Ar (11/12).

**In Situ Quick XAFS Measurements:** The in situ quick XAFS measurements were carried out at the P64 beamline of the DESY Synchrotron Radiation Facility, Hamburg, Germany. A tapered undulator was used as an X-ray source. A Si (111) channel-cut single-crystal monochromator was used with an oscillation frequency of 0.17 Hz. The intensity of incident X-ray radiation was measured by an ionization chamber filled with pure N<sub>2</sub>. The beam size was 0.5 × 0.5 mm<sup>2</sup>. The reference and sample were measured in fluorescence mode using a PIPS detector. For the energy calibration, a  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> reference pellet was measured before each sample measurement for 20 s and then moved out of the beam while the sample was moved into the beam. The in situ measurements were performed in a home-built electrochemical single-compartment cell. The electrolyte used was 0.1 M KOH with 0.016 M KNO<sub>3</sub>, which is 10× diluted from the 1 M KOH + 0.16 M KNO<sub>3</sub> used for standard electrochemical experiments. The electrolyte was diluted to remain compatible with the Kapton window of the in situ electrochemical cell. A Biologic SP300 potentiostat was used to control the potential. Two sets of measurements were performed. In the first experiment, the sample was reduced at –1.5 V versus RHE (–2.45 V vs Ag/AgCl) for 90 s as an activation step and then a potential of –1.0 V versus RHE (–1.95 V vs Ag/AgCl) was applied and held for 15 min. In the second experiment, the potential of –1.0 V versus RHE was applied directly without the activation step.

**Preparation of the Working Electrode:** A carbon paper electrode (AvCarb MGL 370, Fuel Cell Store) was used as the working electrode and was cut to a geometric surface area of 0.25 cm<sup>2</sup> (0.5 × 0.5 cm). An oxygen plasma and acid treatment (0.5 M H<sub>2</sub>SO<sub>4</sub>) were employed to remove the PTFE layer on the electrode and increase the hydrophilicity. A catalyst ink comprised of 5 mg of catalyst, 680  $\mu$ L of isopropanol, 300  $\mu$ L of MilliQ water, and 20  $\mu$ L of a 5 wt% Nafion (probe sonicated for 1 min, followed by 30 min in a sonication bath) was drop cast on the electrode. Catalyst loading on the electrode was optimized during the study, by varying the amount of catalyst ink drop cast.

**Electrochemical Nitrate Reduction:** Electrochemical tests were performed in a customized glass H-cell (Adams & Chittenden), separated by a Celgard 3401 membrane (used as received). A carbon paper with catalyst, a reversible hydrogen electrode (Gaskatel), and a graphite rod comprising the three-electrode system were used as the working, reference, and counter electrodes, respectively. An alkaline electrolyte, 1 M potassium hydroxide (KOH) with 0.16 M potassium nitrate (KNO<sub>3</sub>) was used

for the NO<sub>3</sub>RR tests. The electrochemical cell was deaerated prior to electrochemical experiments by purging N<sub>2</sub> gas (research grade 99.9995% – PraxAir) for 30 min at 80 sscm. During the NO<sub>3</sub>RR, N<sub>2</sub> gas was continuously purged at 30 sscm. Control experiments with only 1 M KOH + N<sub>2</sub> gas demonstrate that the catalyst is not active for N<sub>2</sub> reduction to NH<sub>3</sub>, allowing N<sub>2</sub> to be an inert gas in this system (Figure S14a,b, Supporting Information). For the standard NO<sub>3</sub>RR experiments, the working and counter electrolyte volumes were 30 and 25 mL, respectively. Chronoamperometric (CA) tests were performed for 15 min under vigorous stirring. Prior to CA measurements, the electrode was activated by a pre-reduction step at –1.5 V versus RHE for 90 s. Linear sweep voltammetry was performed by cathodically sweeping from 0.5 to –1.0 V versus RHE at a scan rate of 5 mV s<sup>–1</sup>. Electrochemically active surface area (ECSA) was determined by varying the scan rate between 20–100 mV s<sup>–1</sup> between 0.60–0.75 V versus RHE. For the 24-h durability test, which was segmented into eight, 3-h sections, an electrolyte reservoir of 250 mL was connected to the cathodic chamber of the H-cell using peristaltic pumps and was continually circulated throughout the electrolysis. The large circulating reservoir prevented the buildup of produced NH<sub>3</sub> from becoming too high in the cell. After a 3-h segment, all electrolyte was pumped back into the external reservoir, sampled, and then refreshed for the next 3-h segment. The potential reported for all electrochemical tests was not iR-corrected.

**Isotopic (K<sup>15</sup>NO<sub>3</sub>) Nitrate Reduction:** To confirm the N in the detected NH<sub>3</sub> originated from the KNO<sub>3</sub> feed and not from the N-doped catalyst support, the N<sub>2</sub> gas or other sources of contamination, NO<sub>3</sub>RR with isotopically doped K<sup>15</sup>NO<sub>3</sub> (99% – Cambridge isotopes) was performed. A 1 M KOH + 0.16 M K<sup>15</sup>NO<sub>3</sub> electrolyte was used. Isotopically labeled <sup>15</sup>NO<sub>3</sub>RR was performed at –1.0 V versus RHE for 15 min, after which the electrolyte was sampled and quantified by <sup>1</sup>H NMR. Isotopically doped <sup>15</sup>NH<sub>3</sub> yielded a doublet and standard <sup>14</sup>NH<sub>3</sub> resulted in a triplet.

**Calculation of the Yield and FE:** In this study, all error bars were reported based on a 90% confidence interval from a series of three independent measurements.

The yield rate of ammonia (Yield<sub>NH<sub>3</sub></sub>) from the NO<sub>3</sub>RR was calculated from Equation (2).

$$\text{Yield}_{\text{NH}_3} = \frac{c_{\text{NH}_3} \times V}{Mw_{\text{NH}_3} \times t \times A_{\text{electrode}}} \quad (2)$$

The FE for NH<sub>3</sub>, FE<sub>NH<sub>3</sub></sub>, was calculated from Equation (3).

$$\text{FE}_{\text{NH}_3} = \frac{n \times F \times c_{\text{NH}_3} \times V}{Mw_{\text{NH}_3} \times Q} \quad (3)$$

where  $c_{\text{NH}_3}$  is the concentration of NH<sub>3</sub> in the working chamber (mg mL<sup>–1</sup>), V is the volume of the working chamber (30 mL), the molar mass of ammonia, Mw<sub>NH<sub>3</sub></sub> is 17.031 g mol<sup>–1</sup>, t is the electrolysis time (0.25 h), A<sub>electrode</sub> is the area of the working electrode (0.25 cm<sup>2</sup>), n is the number of electrons transferred (8e<sup>–</sup> for NO<sub>3</sub><sup>–</sup> to NH<sub>3</sub>), F is Faradays constant (96485 C), and Q is the charge passed during the electrolysis (C).

**Product Detection:** For typical NO<sub>3</sub>RR tests, the detection and quantification of NH<sub>3</sub> was achieved using a UV–vis spectrophotometer (Shimadzu, UV-2600). NH<sub>3</sub> was detected using the indophenol blue method in which 2 mL of electrolyte (or diluted electrolyte) was mixed with 2 mL of solution A (1 M NaOH, 5 wt% salicylic acid, and 5 wt% sodium citrate), 1 mL of solution B (0.05 M NaClO), and solution C (1 wt% sodium nitro-ferricyanide). After incubating the dark at room temperature for 1 h, the maximum absorbance was taken at  $\approx$ 655 nm and quantified with respective calibration curves. Calibration curves for the detection and quantification of <sup>14</sup>NH<sub>3</sub> were given in Figure S39, Supporting Information. It was noted that in certain instances, the FE<sub>NH<sub>3</sub></sub> was over 100%. This was due to errors propagated during the dilution of the electrolyte. The electrolyte might be diluted such that the quantified NH<sub>3</sub> would fall within the calibration range. Given the ultrahigh yield rates in this work, dilutions up to 400× were employed.

For the detection of isotopic ammonia ( $^{15}\text{NH}_3$ ),  $^1\text{H}$  NMR was used. 3-(Trimethylsilyl)-1-propanesulfonic acid sodium salt (DSS) was selected as an internal standard and dimethylsulfoxide- $d_6$  (DMSO) was used as the locking solvent. The NMR spectra of a solution of 580  $\mu\text{L}$  of electrolyte, 25  $\mu\text{L}$  of DMSO, 20  $\mu\text{L}$  of 3 M  $\text{H}_2\text{SO}_4$ , and 75  $\mu\text{L}$  of 6 mM DSS were obtained on a Bruker CRYO 500 MHz spectrometer. A solvent suppression method was applied to reduce the signal of  $\text{H}_2\text{O}$ , allowing for better resolution. The spectrum was processed using the Topspin 4.0.8 software. The linear calibration for the detection and quantification of  $^{15}\text{NH}_3$  are given in Figure S40, Supporting Information.

**Techno-Economic Analysis—Levelized Cost of Ammonia:** To evaluate the trade-off between energy efficiency and ammonia partial current density, the levelized cost of ammonia ( $\text{LC}_{\text{NH}_3}$ ) was employed as a metric. The  $\text{LC}_{\text{NH}_3}$  was calculated analogously to Daiyan et al. and was determined by Equation (4).<sup>[22]</sup>

$$\text{LC}_{\text{NH}_3} = \frac{R_f \times \text{CAPEX} + \text{OPEX}}{\text{Yield}_{\text{NH}_3}} \quad (4)$$

where  $R_f$  is the capital recovery factor and is set at 0.08%. CAPEX was the capital cost and solely attributed to the cost of the electrolyzer stack (in \$). OPEX was the operational cost and constituted costs associated with electricity, nitrate feed, and water consumption (all in units of \$).  $\text{Yield}_{\text{NH}_3}$  was the yield of ammonia (kg), giving a  $\text{LC}_{\text{NH}_3}$  in  $\text{\$/kg}_{\text{NH}_3}$ . The  $\text{Yield}_{\text{NH}_3}$  was calculated based as a function of the current density and electrolyzer area (with  $8e^-$  transferred per  $\text{NH}_3$ ).

The electrolyzer stack cost was based on parameters in analogous economic calculations and analysis provided in a National Renewable Energy Laboratory report with a cost of  $\$342 \text{ kW}^{-1}$  operating at a cell voltage of 1.9 V and current density of  $2 \text{ A cm}^{-2}$ , which yielded an assumed  $\text{NO}_3\text{RR}$  electrolyzer stack cost of  $12\,996 \text{ \$ m}^{-2}$ .<sup>[22,46]</sup> In the idealized case where the stack cost could be significantly reduced, a cost of  $\$143 \text{ kWh}^{-1}$  was assumed, resulting in an electrolyzer cost of  $5434 \text{ \$ m}^{-2}$ . For the OPEX costs, the cost of the  $\text{NO}_3^-$  ( $\text{NO}_x$ ) input was assumed to be  $\$315$  per metric ton, as estimated in work by Jiang et al.<sup>[22,47]</sup> The cost of water was assumed to be  $\$0.02 \text{ L}$  and the cost of electricity from the grid was assumed to be  $\$70 \text{ MWh}^{-1}$  and electricity generated from renewable sources was assumed to be  $\$30 \text{ MWh}^{-1}$ . The required electrolyzer area required was calculated from Equation (5).

$$A_{\text{electrolyzer}} = \frac{I_{\text{total}}}{j} \quad (5)$$

where  $I_{\text{total}}$  is the current passing through the electrode (A) and  $j$  is the current density of the  $\text{NO}_3\text{RR}$  system.  $I_{\text{total}}$  was determined by dividing the applied power to the stack (assumed here as 1 MW) by the cell voltage (assumed here as 1.9 V). Therefore, assuming a system current density of  $1.3 \text{ A cm}^{-2}$  (the performance demonstrated in the 24-h electrolysis), the total required electrolyzer area was  $37.6 \text{ m}^2$ .

## Supporting Information

Supporting Information is available from the Wiley Online Library or from the author.

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## Conflict of Interest

The authors declare no conflict of interest.

## Author Contributions

E.M. and B.S. contributed equally to this work. E.M.: Conceived the idea, designed and conducted experiments, processed and analyzed the data, and composed the original draft. B.S.: Designed and conducted experiments, and processed the data. M.R.: Performed ex situ and in situ XAS measurements, data analysis, and manuscript editing. Y.L.: Designed experiments and contributed to discussion and manuscript editing. W.Z.: Performed atomic resolution EELS measurements and analysis. S.G.: Assisted in experimental work, discussion, and manuscript editing. Y.C.: Performed durability experiments and STEM imaging for the revision process. U.H.: Performed ex situ and in situ XAS measurements data analysis. Y.H. and A.L.: Assisted in catalyst characterization and manuscript editing. I.V.Z. and X.P. supervised the study and contributed to discussions. J.T.: Supervised and performed ex situ and in situ XAS measurements data analysis and manuscript editing. B.R.C.: Supervised the study, funding acquisition, discussions, and manuscript editing. E.D.S.: Supervised the study, funding acquisition, discussions, and manuscript editing. P.A.: Conceived the idea, supervised the study, funding acquisition, and manuscript editing.

## Data Availability Statement

The data that support the findings of this study are available from the corresponding author upon reasonable request.

## Keywords

active support, alkaline, ammonia, in situ XAS, iron oxide, nitrate reduction, nitrite reduction, single atom catalysts

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