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Recent developments in Ti-based nanocatalysts for electrochemical nitrate-to-ammonia conversion

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Recently, electrochemical NO₃⁻-to-NH₃ conversion *via* the nitrate reduction reaction (NO₃⁻-RR) has received much attention because it is regarded as an available option for sewage treatment and ammonia synthesis under mild conditions. Exploring promising electrocatalysts with low cost, reduced overpotential, high yield rate and faradaic efficiency toward NH₃, and sufficient stability is the most crucial factor for NO₃⁻-to-NH₃ conversion and energy efficiencies. Considering the advantages of being nontoxic and having wide availability, outstanding stability, hydrogen evolution inertness, and mature fabrication techniques, titanium-based (mainly metallic Ti- and TiO₂-based) nanomaterials have emerged as potential candidates for NO₃⁻-to-NH₃ conversion over a wide pH range. This review summarizes the overview of the NO₃⁻-RR and fundamental insights into metallic Ti and TiO₂, and clarifies the relationship among the design strategy, material structure, and performance enhancement. Furthermore, the recent progress in next-generation Ti-based nanomaterials is discussed, including Ti-based MXene and single atomic catalysts. Finally, the challenges and future directions of the NO₃⁻-RR and Ti-based nanocatalysts are elucidated. This review aims to provide some inspiration for developing effective electrocatalysts for electrochemical NO₃⁻-to-NH₃ conversion.

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1. Introduction

Ammonia (NH₃) plays an important role in the manufacturing industries such as fertilizers, plastics, pharmaceuticals, *etc.*^{1–3} Recently, the emerging “hydrogen economy” and “carbon neutralization” have caused NH₃ to be regarded as a carbon-free fuel and a portable energy carrier owing to its high energy density (4.32 kW h L⁻¹); in addition, NH₃ can be used as a “green fuel” to produce electricity *via* direct ammonia fuel cells.^{4,5} Importantly, NH₃ is a natural hydrogen carrier with a high H₂ storage capacity of 17.75%,⁶ making liquid ammonia a safer hydrogen storage medium. Thus, the primary innovation directions of “ammonia = hydrogen 2.0” are derived,^{7–10} and a new market for NH₃ is estimated to be created soon.

However, NH₃ is still produced by an energy-intensive Haber–Bosch process which requires harsh conditions of high temperature (400–600 °C) and pressure (200–350 atm), resulting in large amounts of global energy consumption and carbon dioxide emission (450 million metric tons).^{11–15} Exploring alternative NH₃ synthetic techniques driven by sus-

tainable energies at room temperature and under an air atmosphere is significant for developing “green ammonia”.^{16,17} On the other hand, N₂, NH₃, and NO₃⁻ are the key inorganic species for the global nitrogen cycle. Humans utilize the Haber–Bosch process for nitrogen fixation, and as-produced NH₃ is used for chemical fertilizer manufacture or provides nitrates *via* the Ostwald process for the chemical and ammunition industries. Fertilizer-intensive agriculture generates NO₃⁻-containing sewages,¹⁸ and NO₃⁻ is also a major waste byproduct stream of industrial production (for instance, the effluent produced from the ammunition industries contains 65% ammonium nitrate and 20% amine nitrates¹⁹). Extensive nitrogen fixation eventually results in the accumulation of NO₃⁻, excessive NO₃⁻ cannot be converted naturally in time, interferes with the global nitrogen cycle and pollutes the surface and ground waters.²⁰ Therefore, it's necessary to develop a denitrification technique to establish a closed nitrogen cycle. The electrochemical NO₃⁻ reduction reaction (NO₃⁻-RR) for NO₃⁻-to-NH₃ conversion (NO₃⁻ + 6H₂O + 8e⁻ → NH₃ + 9OH⁻) driven by sustainable energies is regarded as an alternative for “green ammonia” production and an efficient artificial denitrification technique to repair the disturbed global nitrogen cycle, which has received great attention in recent years.²¹

Electrochemical NO₃⁻-to-NH₃ conversion is an aqueous-based electrochemical strategy for ammonia synthesis using

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water as a hydrogen source, which gets rid of H_2 that is needed for NH_3 synthesis. Moreover, the high theoretical potential (0.69 V vs. RHE) of NO_3^- -to- NH_3 conversion and the high solubility of NO_3^- make the efficiency of NH_3 synthesis superior to that of the electrochemical nitrogen reduction reaction (NRR) – a traditional electrochemical nitrogen fixation technique.^{4,8,22} Nevertheless, electrochemical NO_3^- -to- NH_3 conversion still suffers from the following issues: although electrochemical NO_3^- -to- NH_3 conversion presents an ideal theoretical potential, it requires a large overpotential (>400 mV) to trigger the NO_3^- -RR in reality,^{23,24} due to which the energy conversion efficiency is seriously retarded; the NO_3^- -RR is a complicated multi-electron coupled proton transfer process, in which a series of nitrogen-containing products such as NO_2 , NO_2^- , NO , N_2O , N_2 , NH_2OH , and NH_3 could be generated, and toxic NO_2^- and low added-value N_2 are the major competitive products relative to NH_3 .^{25,26} Studying the reactive mechanisms of the NO_3^- -RR and exploring efficient electrocatalysts are the keys to solving the above bottlenecks.

Recently, Cu- and Co-based electrocatalysts for electrochemical NO_3^- -to- NH_3 conversion have been intensively developed.^{27–35} However, most of them only work in neutral or alkaline electrolytes, as NO_3^- exhibits strong oxidability in acid solutions. Although noble metal-based electrocatalysts exhibit a strong resistance to acid corrosion, their NO_3^- -RR activity is severely compromised by hydrogen evolution. It is desirable to develop efficient electrocatalysts for NO_3^- -to- NH_3 conversion to treat sewage containing NO_3^- over a wide pH range and potential range. Ti-based materials have the inherent advantages of being non-toxic and cost-effective, and having superior stability over a wide pH range and HER inertness, making them ideal candidates. Currently, increasing studies reveal that Ti-based electrocatalysts, typically metal Ti^{36–38} and TiO_2 composite materials,^{39–42} exhibit comparable electrochemical NO_3^- -to- NH_3 conversion activity. Furthermore, there are a series of low-cost fabrication techniques that enable large-scale fabrication of Ti-based materials with various nanostructures under mild conditions. The above superiorities indicate that exploring Ti-based nanomaterials is a crucial development direction for electrocatalysts for NO_3^- -to- NH_3 conversion. Although some exciting results of Ti-based electrocatalysts have been reported in the literature, there is still a lack of comprehensive evaluation of Ti-based electrocatalysts for NO_3^- -to- NH_3 conversion, and the fundamentals of compositional and structural features associated with electrochemical NO_3^- -to- NH_3 conversion performances have not been thoroughly summarized.

Herein, brief summaries about the fundamental insights of the NO_3^- -RR and inorganic Ti-based materials are proposed. Ti-based electrocatalysts with various structures and their catalytic performances for NO_3^- -to- NH_3 conversion have been reviewed in detail. Finally, the challenges and opportunities for further research are also discussed. This review aims to give some new inspiration to develop highly efficient Ti-based nanocatalysts for electrochemical NO_3^- -to- NH_3 conversion.

2. Brief overview of the NO_3^- -RR

2.1. Reactive pathway

The NO_3^- -RR is a complicated process involving multi-electron coupled multi-proton transfer (Fig. 1a).⁴³ Different NO_3^- concentrations, potential ranges and the pH of electrolytes may trigger diverse possible reactive pathways and result in various nitrogen-containing products, including NO_2^- , NO_x , N_2 , NH_3 , etc. Moreover, the exact reactive pathway of the NO_3^- -RR is still under debate. Niu *et al.* proposed five possible reactive pathways of the NO_3^- -RR. They can be classified into O-end, O-side, N-end, N-side and NO-dimer pathways based on the adsorption configuration of $*NO$ (* denotes the state of being adsorbed), as depicted in Fig. 1b.⁴⁴ However, there is still a lack of sufficient experimental evidence to verify the exact reactive pathway. Some of the studies used the N-end pathway of $NO_3^- \rightarrow *NO_3 \rightarrow *NO_2 \rightarrow *NO \rightarrow *NOH \rightarrow *NHOH \rightarrow *NH \rightarrow *NH_2 \rightarrow *NH_3 \rightarrow NH_3(g)$ to perform the density functional theory (DFT) study, while the O-end pathway of $NO_3^- \rightarrow *NO_3 \rightarrow *NO_2 \rightarrow *HNO_2 \rightarrow *NO \rightarrow *HNO \rightarrow *H_2NO \rightarrow *HNO \rightarrow *O \rightarrow *OH$ was often used for Ti-based electrocatalysts.

2.2. Overpotential

Overpotential is a crucial parameter for evaluating the energy conversion efficiency of the NO_3^- -RR. The theoretical potential for NO_3^- -to- NH_3 conversion is 0.69 V vs. RHE.⁸ In reality, a large overpotential (usually more than 400 mV) is needed for triggering the NO_3^- -to- NH_3 conversion. The ideal electrocatalyst is still to be developed, and the origin of overpotential should be further clarified. The NO_3^- -RR is a cathodic reduction reaction in which NO_3^- is transferred to a series of nitrogen-containing products by accepting electrons on the electrode surface. The rate-limiting step in the entire reaction is usually the reduction of NO_3^- to the intermediate product NO_2^* due to the difficulty of breaking the N–O bond.^{21,45} It is the process of electron transition from electrodes to the lowest occupied molecular orbital (LUMO) of NO_3^- , according to the standpoint of frontier orbital theory. This process can be vividly expressed in Fig. 1c for metal and semiconductor electrodes. The molecular orbital diagram of NO_3^- shown in Fig. 1d suggests that the high-energy LUMO of the NO_3^- ion and the d orbitals of transition metals (except Hg) are hard to match with the LUMO of NO_3^- , thus extra energy is needed for reducing NO_3^- to the intermediate product NO_2^* .⁴⁶ Therefore, exploring electrocatalysts with high energy in the highest occupied state (HOS) is essential for reducing the overpotential.^{47,48}

2.3. Product selectivity

For NO_3^- -to- NH_3 conversion, NH_3 is the target product of the NO_3^- -RR. Thus, the selectivity and faradaic efficiency (FE) of NH_3 are crucial parameters. However, NO_3^- -to- NH_3 conversion involves eight electrons coupled with the transfer of nine protons. Many intermediates with the valence of nitrogen between +5 and –3 may desorb from electrocatalysts to gene-

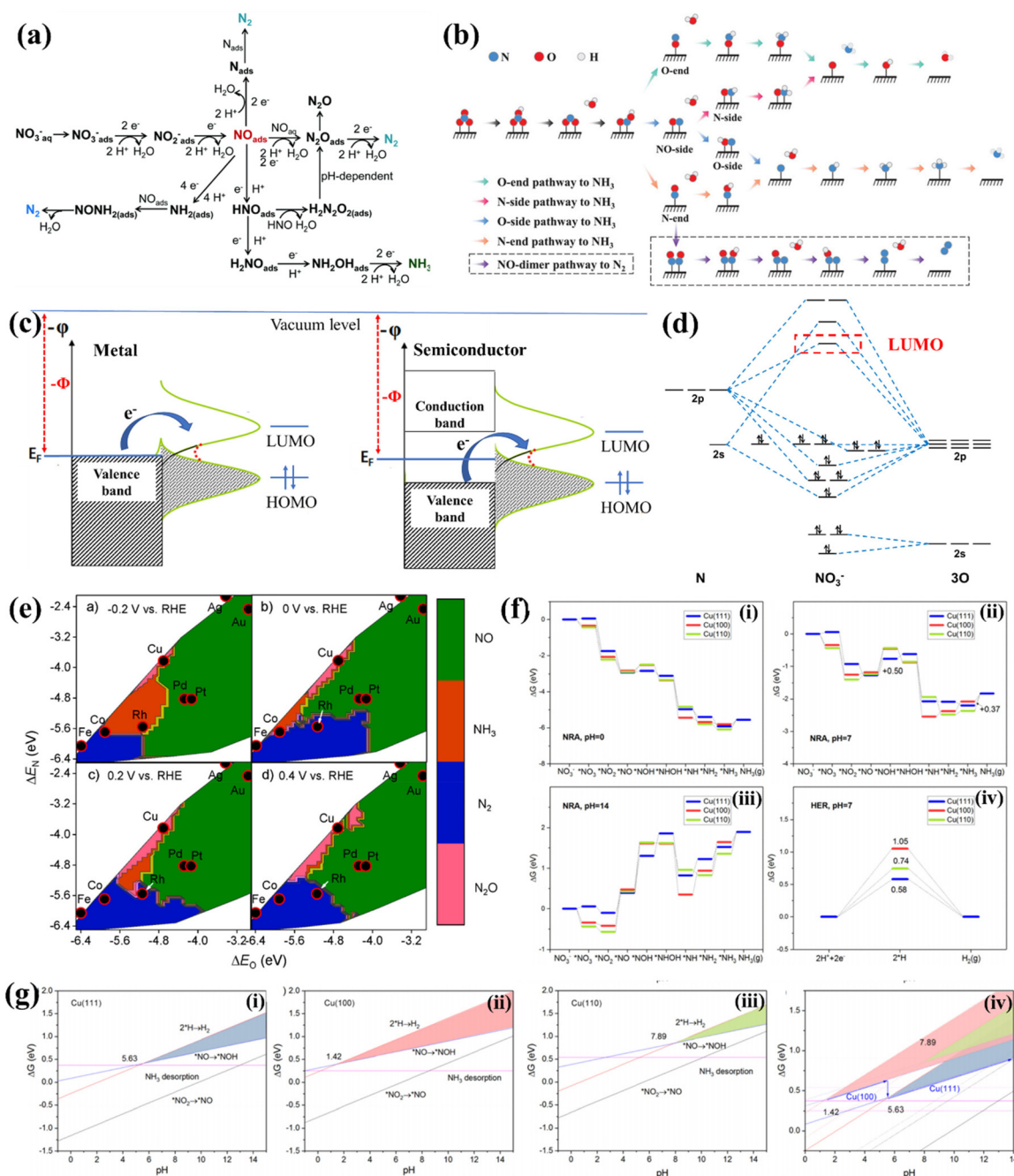


Fig. 1 (a) The possible pathways of the NO₃⁻RR.⁴³ Reproduced from ref. 43 with permission from The Royal Society of Chemistry, copyright 2021. (b) Detailed pathways of the NO₃⁻RR, including O-end, O-side, N-end, and N-side pathways to NH₃, as well as the NO-dimer pathway to N₂.⁴⁴ Reproduced from ref. 44 with permission from Wiley-VCH, copyright 2020. (c) Electronic band structures of metal- and semiconductor-based electrodes, and the electron transfer from the electrode to the frontier orbitals of the adsorbate. (d) Molecular orbital diagram and the LUMO of NO₃⁻. (e) Theoretical selectivity maps to NO, N₂O, N₂ or NH₃ products from electrocatalytic NO₃⁻ reduction as a function of O and N adsorption energy under different applied voltages.⁴⁹ Reproduced from ref. 49 with permission from American Chemical Society, copyright 2019. (f) NO₃⁻RR pathways at Cu(111), Cu(100), and Cu(110) at (i) pH = 0, (ii) pH = 7, (iii) pH = 14, and (iv) the HER pathway at pH = 7. (g) Competing relationship between the NO₃⁻RR and the HER on (i) Cu(111), (ii) Cu(100), and (iii) Cu(110), and (iv) comparison between different facets. The shaded zones are predominated by the NO₃⁻RR.⁵⁶ Reproduced from ref. 56 with permission from American Chemical Society, copyright 2021.

rate various by-products, and the coupling reaction between neighboring *N leads to the generation of N₂, N₂H₄ or N₂O as by-products. Liu *et al.* revealed that the adsorption strengths of O and N atoms (ΔE_O and ΔE_N) could be used as descriptors of

selectivity on transition metals. As depicted in Fig. 1e, moderate adsorption of O and N prefers NH₃ generation, strong adsorption of O and N results in N₂ formation, and the weak adsorption intensity of N and O tends to produce NO. They

proposed that Co and Rh are ideal candidates for electrochemical NO_3^- -to- NH_3 conversion.⁴⁹

Furthermore, the d-band model is commonly used for describing and anticipating the selectivity of the NO_3^- -RR. The d-band center (E_d) is related to the adsorption strengths of intermediates, which increase with the E_d of active sites approaching the Fermi level. NO_2^- is regarded as a major byproduct of NO_3^- -to- NH_3 conversion. A more negative $\Delta(G_{\text{NO}^*} - G_{\text{NO}_2^*})$ is desired for increasing the preference toward nitrite reduction to nitric oxide. As E_d approaches the Fermi level, $\Delta(G_{\text{NO}^*} - G_{\text{NO}_2^*})$ becomes increasingly negative, and $^*\text{NO}_2$ tends to be reduced to $^*\text{NO}$.^{50,51} Subsequently, the adsorbed nitric oxide ($^*\text{NO}$) serves as a critical intermediate in determining the selectivity between nitrogen/oxides and ammonium or hydroxylamine.⁵² NH_3 could be generated from $^*\text{NO}$ via two pathways: one is the so-called Eley-Rideal-like proton-coupled electron transfer, in which $^*\text{NO}$ was reduced to hydroxylamine ($^*\text{NH}_2\text{OH}$),⁵³ and the other involves the dissociation of $^*\text{NO}$ into $^*\text{N}$ and $^*\text{O}$ and the Langmuir-Hinshelwood-like hydrogenation of $^*\text{N}$ by $^*\text{H}$ to NH_3 .^{54,55} The latter has been proposed to favor NH_3 production, namely, dissociative adsorption of nitric oxide is desired for NO_3^- -to- NH_3 conversion. Theoretical studies revealed that the dissociation activation barriers of $^*\text{NO}$ decrease with increasing its adsorption strength, suggesting that the E_d of active sites should be close to the Fermi level to activate the dissociation of nitric oxide for NH_3 formation.⁵⁰

On the other hand, the HER is the major competitive reaction of the NO_3^- -RR, which also retards the selectivity and FE of NO_3^- -to- NH_3 conversion. Hu *et al.* found that the pH influences the competition between the HER and the NO_3^- -RR on the (100), (111), and (110) facets of Cu. As depicted in Fig. 1f, the Gibbs free energies (ΔG) of intermediates along pathways on Cu(111), Cu(100), and Cu(110) are dependent on the pH of the electrolyte, and the ΔG -pH plots of the potential rate-determining step of the NO_3^- -RR and HER are shown in Fig. 1g (i-iii). The pH range corresponding to the triangle zone is suitable for the NO_3^- -RR, and the critical pH values of the NO_3^- -RR are 5.63, 1.42 and 7.89 for Cu(111), Cu(100) and Cu(110), respectively. Fig. 1g(iv) suggests that Cu(100) works at pH ranging from 1.42 to 5.63 while Cu(111) works at pH ranging from 5.63 to 14, indicating that Cu(111) works more effectively. Moreover, the product selectivity is highly dependent on the applied potential, and a negative potential is needed for NH_3 generation while the HER tends to retard the FE and selectivity of NH_3 .⁵⁶ The above results verify that the NO_3^- -RR is potential-dependent and pH-sensitive; thus, seeking HER inert materials for constructing NO_3^- -RR electrocatalysts which can work in a wide pH and potential range is highly desired.

3. Fundamental insights of metallic titanium and titanium oxides

Titanium (Ti) is not only a HER inert element but also one of the most abundant elements in the Earth's crust. Metallic

Ti or its alloys and oxides have been widely used in aerospace, photocatalysis, energy storage, and other fields due to their advantages of low cost, superior mechanical strength, unique electronic structure, excellent corrosion resistance and stability, *etc.*⁵⁷ Some studies revealed that Ti-based inorganic nanomaterials, majorly, metallic Ti- and TiO_2 -based nanomaterials exhibited impressive performances of electrochemical NO_3^- -to- NH_3 conversion. The exploration of low-cost and highly-efficient Ti-based electrocatalysts for the conversion of NO_3^- -to- NH_3 has attracted much attention. In this part, we systematically review the structure of metallic Ti and TiO_2 to help rationalize the design of Ti-based nanomaterials for electrochemical NO_3^- -to- NH_3 conversion.

3.1. Metallic Ti

Metallic Ti is a silver-white transition metal with a high melting point of 1660 ± 10 °C, a low density of 4.506 g cm^{-3} , and excellent corrosion resistance. The valence electronic configuration of Ti is $[\text{Ar}]3d^24s^2$. There are two isomers of titanium: α -Ti and β -Ti; their crystallographic structures are depicted in Fig. 2a and b, respectively. α -Ti belongs to the hexagonal crystal system with an atomic space utilization rate of 74%, while β -Ti presents body-centered cubic dense packing with a utilization rate of 68%. The transition temperature of the above isomers is 882.5 °C, α -Ti is stable below 882 °C, and β -Ti is stable between 882 °C and the melting point; thus α -Ti is the common form in reality.^{58,59} The work function (Φ shown in Fig. 1c) of metallic Ti is 4.33 eV, lower than those of Co (5 eV), Cu (4.65 eV), Fe (4.5 eV), and Ru (4.71 eV) usually regarded as electrocatalysts for the NO_3^- -RR,⁶⁰ suggesting that Ti presents a high-energy Fermi level and may be an ideal candidate with a lower overpotential of the NO_3^- -RR. However, titanium usually forms a stable oxide layer on its surface when exposed to air, and the active sites should be carefully identified while using metallic Ti-based materials as catalysts.

3.2. Titanium oxides

TiO_2 belongs to the category of typical semiconductors.⁶¹ There are many kinds of crystallographic structures, including rutile, anatase, brookite and $\text{TiO}_2(\text{B})$, and several metastable polymorphs, such as $\text{TiO}_2(\text{H})$, $\text{TiO}_2(\text{II})$, and perovskites, have been artificially synthesized.⁶¹ Metastable phases are rarely observed as stable electrocatalysts, and the $\text{TiO}_2(\text{B})$ phase is less common. Thus, they will not be discussed in this review. Anatase and rutile have more comprehensive applications because they are more stable than brookite,⁵⁷ hence they are emphatically discussed. Generally, rutile is the most stable phase and is usually synthesized by high-temperature deposition or annealing. Anatase-phased nanomaterials are commonly obtained by solution-based or low-temperature vapor deposition systems.⁶¹ Anatase and rutile phases present tetragonal structures (as depicted in Fig. 2c and d) with slightly distorted TiO_6 octahedra as a fundamental building block,^{57,62,63} and have bandgaps of 3.2 and 2.96 eV, respectively.^{57,64} A large bandgap results in poor

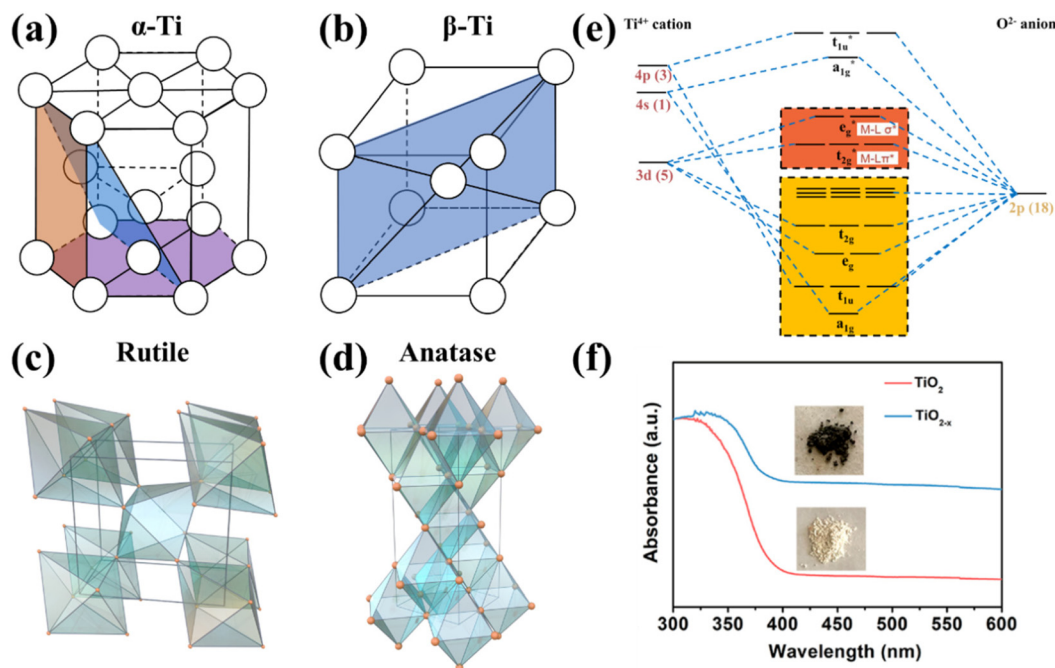


Fig. 2 Unit cells of (a) α -Ti, (b) β -Ti, (c) rutile TiO_2 and (d) anatase TiO_2 .⁶¹ Reproduced from ref. 61 with permission from American Chemical Society, copyright 2014. (e) Molecular orbitals of the octahedral TiO_6 unit. (f) UV-vis absorption spectra (the insets are optical photographs) of TiO_2 and TiO_{2-x} .⁶⁵ Reproduced from ref. 65 with permission from American Chemical Society, copyright 2020.

electron conductivity of TiO_2 . Based on the ligand-field theory, the orbital diagram of the TiO_6 octahedral unit is shown in Fig. 2e. Since the valence electronic configuration of Ti^{4+} is $[\text{Ar}]3d^04s^0$, the bonding orbitals (a_{1g} , t_{1u} , e_g and t_{2g}) are mainly contributed by the 2p orbitals of O^{2-} , and the unoccupied anti-bonding orbitals (a_{1g}^* , t_{1u}^* , e_g^* and t_{2g}^*) are mainly contributed by the 3d, 4s and 4p orbitals of Ti^{4+} . Therefore, the valence band maximum and the conduction band minimum originate from O 2p and Ti 3d, respectively.

Oxygen vacancies (OVs) are easily created on the surface of TiO_2 , thereupon leading to the formation of unsaturated Ti^{3+} sites and nonstoichiometric TiO_{2-x} ($0 < x < 1$). According to the stoichiometry theory for semiconductors, TiO_{2-x} belongs to an n-type semiconductor with a narrow bandgap compared to TiO_2 ,^{57,66} and the electrons in TiO_{2-x} are easily excited from the valence band to the conduction band to form holes and carriers. Therefore, TiO_{2-x} can absorb visible light to show color (Fig. 2f), and the electron conductivity of TiO_{2-x} is strengthened compared to that of intrinsic TiO_2 .⁶⁵ Furthermore, the Fermi level could be upshifted by transforming the intrinsic semiconductor to an n-type semiconductor, which is desired for reducing the overpotential of the NO_3^- RR, as discussed in section 2.2. On the other hand, the electron conductivity of TiO_2 can also be modified by heteroatom doping. By controlling the doping pattern, intrinsic TiO_2 can be transformed to an n- or p-type semiconductor.^{57,67} The bandgap, electron conductivity, and the Fermi level could be regulated subsequently.

4. Nitrate-to-ammonia conversion performance of Ti-based nanocatalysts

4.1. Metallic Ti-based electrocatalysts

Considering that titanium exhibits a higher Fermi level, poor HER activity and high corrosion resistance, as discussed in section 3, some researchers tried to explore metallic Ti-based electrocatalysts for NO_3^- -to- NH_3 conversion.

Fajardo *et al.* studied the NO_3^- RR on a series of transition metals Ti, Fe, Co, Ni, Cu, Zn, and Sn by electroreduction of 100 mg L^{-1} NO_3^- -N in $50 \text{ mM Na}_2\text{SO}_4$ at 20 mA cm^{-2} and 360 min of treatment time. They found that the NO_3^- degradation on the above materials conformed to the pseudo-first-order characteristic. The selectivity of N_2 for Ti is almost negligible, the FE_{NH_3} of Ti is close to those of Co and Fe and higher than those of Ni, Cu, Zn and Sn, and the reaction kinetics of Ti is even comparable to that of Pt. The above results suggest that NH_3 generation on Ti is electrocatalytically preferential over N_2 evolution.³⁶ To further reveal the rule of the NO_3^- RR on Ti, McEnaney *et al.* systematically studied the effect of pH, nitrate concentration, and applied potential on the FE_{NH_3} of the NO_3^- RR on the Ti electrode. The heatmaps constructed by an entire grid of electrolyte conditions and FE_{NH_3} shown in Fig. 3a suggest that (1) more extreme pH values give significantly higher total current densities than those of moderate pH, (2) lower pH generally corresponds to higher FE_{NH_3} , and (3) in moderate base and moderate acid electrolytes, moderate

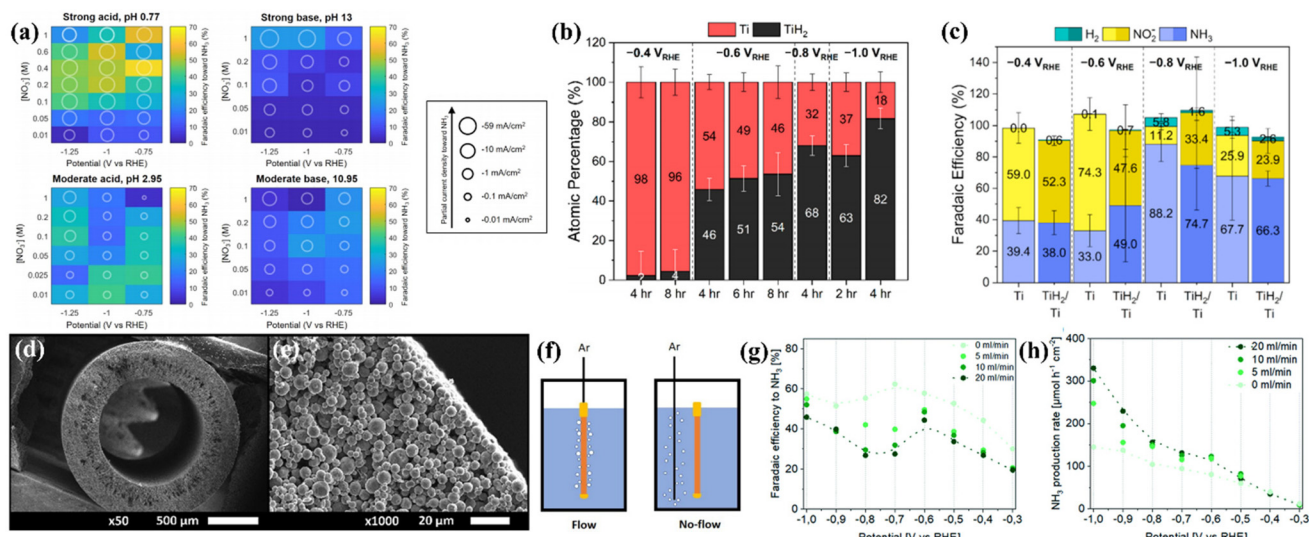


Fig. 3 (a) Heatmap plots at four distinct pH values showing faradaic efficiency to NH_3 by varying applied potentials and nitrate concentrations. Each grid block displays the data from a 30 min potentiostatic experiment performed at the indicated pH, nitrate concentration, and applied potential.¹³ Reproduced from ref. 13 with permission from American Chemical Society, copyright 2020. (b) Ti atomic percentages of the Ti foil cathode after the NO_3^- RR obtained from the EXAFS modeling. (c) Faradaic efficiencies toward the production of ammonia, nitrite, and hydrogen gas on unamended Ti and TiH_2/Ti electrodes.³⁸ Reproduced from ref. 38 with permission from American Chemical Society, copyright 2022. (d) SEM images the cross section of a Ti hollow fiber electrode at different magnifications; (e) gas flow configurations of the working electrode compartment; (f) performance of Ti hollow fiber electrodes for the electrochemical reduction of NO_3^- to NH_3 in 0.1 M HClO_4 using 50 mM KNO_3 ; potential- and Ar flow rate-dependent (g) faradaic efficiency and (h) production rate toward ammonia.³⁷ Reproduced from ref. 37 with permission from the Royal Society of Chemistry, copyright 2022.

nitrate concentrations (between 0.025 and 0.2 M NO_3^-) generate higher FE_{NH_3} . According to the above results, the authors used an electrolytic cell constructed by using a Nafion membrane divider with 0.1 M HNO_3 /0.3 M KNO_3 electrolyte for the NO_3^- RR, and achieved the highest FE_{NH_3} of 82%. Importantly, the authors first revealed that titanium hydride (TiH_x) is generated on the Ti electrode after the reaction.¹³ Subsequently, Liu *et al.* focused on this issue and utilized highly surface-sensitive techniques such as *ex situ* grazing-incidence X-ray diffraction (GIXRD) and total electron yield X-ray absorption spectroscopy (TEY XAS) to deeply elucidate the self-reconstruction of metallic Ti foil. The GIXRD result suggests that α -Ti was the primary phase of unamended Ti foil; the diffraction patterns for TiH_x appeared after the NO_3^- RR was performed at -0.6 V *vs.* RHE in 0.1 M HClO_4 + 0.8 mM KNO_3 . The quantitative analysis of Ti K-edge TEY XAS measurements further suggested that more negative applied potential and longer applied durations promoted near-surface TiH_2 enrichment, as depicted in Fig. 3b. The electrochemical NO_3^- RR performance of unamended Ti and preformed TiH_2/Ti electrodes was assessed with 30 min chronoamperometry experiments at -0.4 , -0.6 , -0.8 , and -1.0 V *vs.* RHE in 1 M NaClO_4 + 10 mM HNO_3 , GIXRD of unamended Ti electrodes after such a short-time chronoamperometry showed that self-reconstruction could not be observed, and the NO_3^- RR performance of unamended Ti could be attributed to the intrinsic Ti foil. Surprisingly, the FE_{NH_3} , $\text{FE}_{\text{NO}_2^-}$ and FE_{H_2} of unamended Ti and preformed TiH_2/Ti electrodes depicted in Fig. 3c followed a similar trend with almost

identical partial current densities of NH_3 .³⁸ Thus, the mystery of whether the active species of the metallic Ti electrode for the NO_3^- RR is Ti, Ti hydride, or a combination of species remains under debate, and the above two studies remind us that the self-reconstruction during the NO_3^- RR cannot be neglected, even if Ti-based materials are usually regarded as stable components. It should be highlighted that the near-surface structure is important for the NO_3^- RR.

A representative study about electrochemical NO_3^- -to- NH_3 conversion on metallic Ti-based nanomaterials reported by Krzywda *et al.* suggested that a tubular porous Ti electrode prepared by dry-wet spinning exhibited interesting catalytic behaviors. The as-prepared Ti electrode exhibited uniform pore distribution over the entire length of the fiber (Fig. 3d and e). They introduced a flow of inert gas exiting the wall of the hollow fiber electrode (Ar flow through the hollow fiber electrode from the inside to the outside, denoted as flow-through conditions), as illustrated in Fig. 3f. The current density of tubular porous Ti electrodes with flow-through conditions is higher than that with no-flow (Ar was introduced through an external sparging line next to a hollow fiber electrode) and vigorous magnetic stirring conditions. This result excluded the possibility that enhanced current density originated from the bubbling induced convective flow. Although the flow-through conditions decreased the FE_{NH_3} (Fig. 3g), the yield rates and partial current densities of NH_3 eventually improved (Fig. 3h). The authors suggested that the flow-through conditions promote the transport of protons and nitrate towards the elec-

trode, thus increasing the catalytic current density. However, the flow-through conditions take NO and N₂O away from the solution, promote the homogeneous reaction of NO₂⁻ and NH₂OH to NO and N₂O, hinder the consecutive electrochemical reduction of NO₂⁻ and NH₂OH towards NH₃, and eventually result in a decrease of FE_{NH₃}.³⁷ Furthermore, Tarpeh *et al.* constructed a representative flow-cell configuration with a polycrystalline titanium electrode for the NO₃⁻RR and revealed that NO₂⁻ and NH₃ accounted for almost all NO₃⁻RR products, the selectivity was flow rate dependent, and NH₃ was favored at the lowest flow rate. The above results suggested that the NO₃⁻RR was subject to mass transport limitations.⁶⁸

4.2. TiO₂-based electrocatalysts

TiO₂ is suitable as a substrate for loading active materials due to its low cost, non-toxic nature and corrosion resistance. The electrons in TiO₂ can be excited from the valence band to the conduction band, and the electrons and holes are able to diffuse to the semiconductor surface for participating in the reaction; thus, modulating the valence electronic structure is an alternative for narrowing the bandgap and improving the electron conductivity of intrinsic TiO₂. In this section, three common strategies for modulating the NO₃⁻-to-NH₃ conversion performances of TiO₂-based nanomaterials including heteroatom doping, creating oxygen vacancies and heterostructures will be introduced.

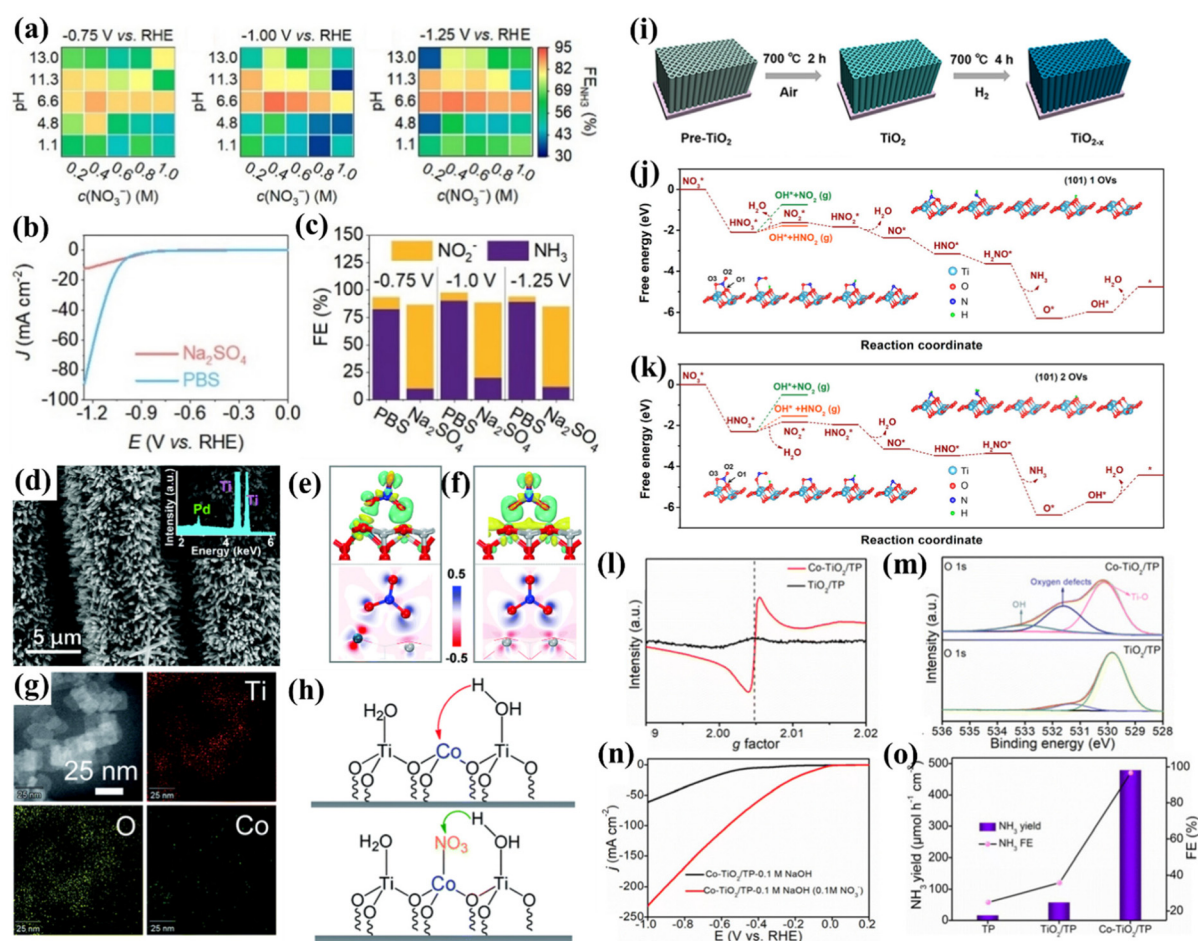


Fig. 4 (a) Heatmaps of FE_{NH₃} in electrolyte solution with different NO₃⁻ concentrations and pH values under three applied potentials; comparison of NO₃⁻RR activities in 1.0 M PBS and 1.0 M Na₂SO₄ with 0.4 M KNO₃, (b) LSV curves, (c) FE_{NH₃},³⁹ reproduced from ref. 39 with permission from Wiley-VCH, copyright 2022; (d) SEM images and the EDX spectrum (inset) of Pd/TiO₂ nanoarrays; (e and f) 3D and 2D electron density difference mappings for the optimized (e) Pd/TiO₂-NO₃⁻ and (f) TiO₂-NO₃⁻ structures;⁴⁰ reproduced from ref. 40 with permission from the Royal Society of Chemistry, copyright 2021; (g) EDS elemental mapping images of Co/TiO₂ NSs; (h) proposed hydrogen-activated routes and the corresponding hydrogen acceptors in the HER and NO₃⁻RR processes;⁴¹ reproduced from ref. 41 with permission from the Royal Society of Chemistry, copyright 2022; (i) schematic illustration for TiO_{2-x} synthesis; calculated free energy changes of the nitrate reduction reaction on the TiO₂ (101) surface with (j) one, and (k) two oxygen vacancies in one 1 × 3 slab at 0 V vs. RHE,⁶⁵ reproduced from ref. 65 with permission from American Chemical Society, copyright 2020; (l) EPR spectra and (m) O 1s regions of Co-TiO₂/TP and TiO₂/TP; (n) LSV curves of Co-TiO₂/TP in 0.1 M NaOH with and without 0.1 M NO₃⁻; (o) NH₃ yields and FEs of Co-TiO₂/TP, TiO₂/TP, and bare TP at -0.5 V vs. RHE.⁴² Reproduced from ref. 42 with permission from The Royal Society of Chemistry, copyright 2022.

Firstly, the electrochemical NO_3^- -to- NH_3 conversion behavior of intrinsic TiO_2 must be clarified. Xu *et al.* synthesized anatase TiO_2 and systematically studied its NO_3^- -RR behavior affected by electrolytic conditions, including pH values, nitrate concentration, and the type of electrolyte. The heatmap in Fig. 4a shows that the relationship between FE_{NH_3} and the NO_3^- concentration exhibits an approximate volcano shape, and the relationship between FE_{NH_3} and the pH of the electrolyte also obeys a similar tendency. According to the heatmap, higher FE_{NH_3} could be achieved in neutral media of 0.4 M KNO_3 + 1.0 M PBS (pH = 6.6), unlike metallic Ti, which is more inclined to produce NH_3 in acidic electrolytes (Fig. 3e). This phenomenon demonstrates that NO_3^- -RR behaviors on both metallic Ti and TiO_2 are pH-sensitive. To further understand the insightful effect of the electrolyte, the electrolyte was switched into another neutral solution of 0.4 M KNO_3 + 1.0 M Na_2SO_4 (pH = 6.7). Fig. 4b and c suggest that TiO_2 displayed a much lower catalytic current density, FE_{NH_3} , and partial current density of NH_3 in an electrolyte of 0.4 M KNO_3 + 1.0 M Na_2SO_4 . To reveal the essence of this phenomenon, the authors tested the pH values of two electrolytes after 30 min electrolysis. Consequently, PBS only increased 0.3 pH units while in Na_2SO_4 solution the pH drastically increased from 7 to 13. Therefore, the pH of the electrolyte could be maintained in a suitable range for NH_3 formation in the buffer. However, the pH of Na_2SO_4 quickly ascended to an inferior area shown in Fig. 4a, which inhibited the NH_3 formation. Moreover, the authors proposed that faster proton transport in PBS is responsible for the larger catalytic current density. This effect could increase the proton concentration on the electrode surface and facilitate the hydrogenation of NO_3^- to NH_3 .³⁹ This study preliminarily revealed the NO_3^- -RR behavior on TiO_2 and demonstrated optimal electrochemical conditions for electrochemical NO_3^- -to- NH_3 conversion using TiO_2 as an electrocatalyst.

4.2.1. Heteroatom doping. Heteroatom doping modifies the valence electronic structure of TiO_2 by elemental interaction, which is a straightforward strategy for modulating the electrochemical performances.

A representative study of heteroatom doping is presented by Guo *et al.*, who successfully doped Pd atoms into TiO_2 (Fig. 4d) for NO_3^- -to- NH_3 conversion. They proved that the lattice stress caused by doping Pd atoms into the TiO_2 phase creates dislocations and distortions forming the active sites of the catalytic reactions. DFT studies revealed that the HOMO of NO_3^- is located at the O atoms. Fig. 4(e and f) further illustrate that the introduction of Pd atoms makes the electrons of TiO_2 transfer to Pd atoms, forming an electron-rich accumulation on Pd, promoting electron transfer to the oxygen atoms of nitrate ions, and eventually accelerating the dissociation of the N–O bond. Meanwhile, a novel Zn-nitrate cell system was assembled for the first time based on the Pd/ TiO_2 catalyst as the cathode and metal Zn as the anode. The battery delivers remarkable dual functions, *i.e.*, it generates electricity and produces NH_3 at the same time with a peak power density of 0.87 mW cm^{-2} and a high FE_{NH_3} of 81.3%. This study demonstrated the feasibility of the galvanic nitrate-based cell, which

broadens the field of Zn-based batteries.⁴⁰ Xu *et al.* synthesized Co(II)-decorated TiO_2 nanosheets (Fig. 4g) exhibiting excellent nitrate performance with an FE and NH_3 yield of 97.4% and 0.348 $\text{mmol cm}^{-2} \text{h}^{-1}$ respectively in 1.0 M phosphate buffer solution (PBS, pH = 6.53). The inhibited HER mechanism was also investigated. They demonstrated a proton pumping mechanism for thermodynamically facilitating proton transfer in the presence of nitrate as follows: in the HER, the hydrogen/proton acceptor should be the Co atoms on Co/ TiO_2 NSs, while the hydrogen/proton acceptor switches into the N/O atoms of nitrate adsorbed on the Co sites in the NO_3^- -RR, and Ti serves as active sites for water dissociation, as depicted in Fig. 4h. Due to the negative charge and delocalized conjugated π electron cloud on NO_3^- , the barrier of proton transportation from H_2O to NO_3^- (0.508 eV) is much lower than that to Co^{2+} (0.805 eV), thus the HER could spontaneously switch to the NO_3^- -RR and the energy input for the NO_3^- -RR could be lowered simultaneously.⁴¹

4.2.2. Oxygen vacancy. The OVs, as an anionic defect, provide electrons to the catalyst surface and endow the lattice oxygen with a higher electron density, making it easier to participate in the catalytic reaction. OVs are easy to form on TiO_2 and they can vary the coordinative structure and the valence state of Ti on the surface (as discussed in section 3.2). Creating OVs is also a common strategy for optimizing the electronic structure of TiO_2 . Moreover, the increase of OVs in TiO_2 is closely related to the NO_3^- -RR activity.

Jia *et al.* synthesized oxygen vacancy-enriched TiO_{2-x} nanotubes by hydrogen reduction (Fig. 4i) and the as-prepared TiO_{2-x} nanotubes were regarded as an efficient electrocatalyst for NO_3^- -to- NH_3 conversion. DFT calculations suggested that introducing OVs on the surface of TiO_2 resulted in the occupation of excess 3d electrons of Ti, and the Fermi level could be moved into the conduction band minimum, eventually giving rise to the metallic behavior of TiO_{2-x} and then improving the electron conductivity. This study revealed that OVs are the adsorption sites filled by the oxygen atoms in nitrate, which weaken the N–O bonding. By comparing the Gibbs free energy diagram of TiO_2 (101) with one and two vacancies, the latter needed a higher reaction barrier for the formation of HNO_2 (Fig. 4j and k), thus suppressing the generation of byproducts.⁶⁵

The enhancement of oxygen vacancies by metal doping has also received much attention. Song *et al.* utilized electrochemical anodic oxidation and electrodeposition to synthesize a highly dispersed Cu-doped TiO_2 nanotube array (Cu/TNTA) cathode with a high electrocatalytic NO_3^- -RR efficiency and long-term stability. They discovered that just doping trace amounts of Cu into TiO_2 could activate more OVs during the NO_3^- -RR process. The OVs highly promoted charge transfer between the NO_3^- and the electrocatalyst, thus reducing the energy barrier of the NO_3^- -RR, eventually reducing the overpotential required for triggering the NO_3^- -RR and improving the yield rate of NH_3 and FE_{NH_3} .⁶⁹ Similarly, Yu's group synthesized a unique structure of Cu clusters homogeneously supported on TiO_2 nanosheets with abundant OVs ($10\text{Cu}/\text{TiO}_{2-x}$).

DFT calculations suggested that OVVs should be the strong adsorption sites of NO_3^- . The N–O bond of NO_3^- breaks automatically during the adsorption process, and the by-products of NO_2 or HNO_2 are suppressed by OVVs with higher reaction barriers, suggesting that the positive effect of interfacial OVVs optimized the NO_3^- RR on $10\text{Cu}/\text{TiO}_2-x$.⁷⁰ Zhao *et al.* skillfully utilized $\text{Na}_2\text{Ti}_2\text{O}_5$ grown on a Ti plate as a precursor to prepare CoTi_2O_5 by an ion exchange strategy, and then the Co-doped TiO_2 nanoribbon array grown on the Ti plate ($\text{Co-TiO}_2/\text{TP}$) was achieved by Ar-annealing of CoTi_2O_5 . XRD patterns suggested that Co in the lattice of TiO_2 reduced the crystal quality. Moreover, the EPR and XPS results verified that OVVs were introduced into TiO_2 after Co doping (Fig. 4l and m). DFT calculations revealed that Co-TiO_2 shows a higher impurity level near the Fermi level compared to TiO_2 , which facilitates charge transfer at the interface. Co doping and OVVs also reduced the Gibbs free energy barrier from $^*\text{NO}$ to $^*\text{N}$, which was regarded as a potential determining step in these systems. These advantages enabled $\text{Co-TiO}_2/\text{TP}$ to attain an improved onset potential compared to TiO_2/TP , a large NH_3 yield of

$1127 \mu\text{mol h}^{-1} \text{cm}^{-2}$ at -0.9 V and a high FE_{NH_3} of 98.2% at -0.5 V vs. RHE (Fig. 4n and o).⁴²

The above Cu/Co induced OVVs generated in TiO_2 could be attributed to the charge difference between $\text{Cu}^{2+}/\text{Co}^{2+}$ and Ti^{4+} . Doping $\text{Cu}^{2+}/\text{Co}^{2+}$ into the TiO_2 lattice must create OVVs to compensate for the missing positive charge while remaining charge neutral.⁷¹

4.2.3. Heterostructure. Stacking any two dissimilar materials with different Fermi levels (work functions, Φ shown in Fig. 1c) to form a heterostructure could form a built-in electric field (BEF). As shown in Fig. 5a, the Fermi level (or work function) represents the electrochemical potential of electrons in the materials, and the difference in Fermi levels results in electrons crossing the heterostructure from the material with a higher Fermi level to the other to form a BEF until their Fermi levels reach equilibrium. Therefore, the electronic interaction that originated from the rectifying effect can also effectively modify the valence electronic structure of TiO_2 -based materials. Based on the above discussions, a series of studies achieved impressive results.



Fig. 5 (a) The formation of a built-in electric field at the heterostructure composed of two components with different Fermi levels. (b) Representative HRTEM image of CuCl_BEF . (c) Molecular dynamics simulation of CuCl and CuCl_BEF in KNO_3 (100 mg L^{-1}) solution. Scale bar, 0.5 nm . (d) Distribution of NO_3^- along the z -axis electrode distance based on the molecular dynamics simulation.⁷² Reproduced from ref. 72 with permission from Wiley-VCH, copyright 2022. (e) TEM and (f) HRTEM images of $\text{FeS}_2@/\text{TiO}_2/\text{TP}$. (g) Charge density difference of $\text{FeS}_2@/\text{TiO}_2$ with adsorbed NO_3^- , where yellow and cyan color indicate electron accumulation and depletion, respectively, and the isosurface values are set to $0.000213 \text{ \AA}^{-3}$.⁷³ Reproduced from ref. 73 with permission from The Royal Society of Chemistry, copyright 2022. (h) SEM images and (i) HRTEM images of CoP/TiO_2 . (j) Calculated PDOS of CoP/TiO_2 .⁷⁶ Reproduced from ref. 76 with permission from Elsevier, copyright 2022.

Sun *et al.* stacked CuCl (111) and rutile TiO₂ (110) layers together (Fig. 5b), a BEF could be induced by electron transfer from TiO₂ to CuCl due to the difference in their Fermi levels, and the BEF strength was roughly estimated to be 8×10^8 V m⁻¹. Molecular dynamics simulation and finite element analysis suggested that the BEF accumulated NO₃⁻ in the diffusion layer near the surface of the electrocatalyst (Fig. 5c and d). This study demonstrated that the BEF increased the free energy of *ON, resulting in a great decrease of Δ*G* of the potential determining step. This phenomenon could be elucidated as the BEF resulted in electron richer Cu(i), which suppressed the electron donation from *NO but facilitated the back donation to the π* anti-bonding orbital of *NO, thus, destabilizing the N–O bond of *NO. The above advantages endowed the heterostructure with an NH₃ selectivity of 98.6% and a yield rate of 1.82 mg h⁻¹ cm⁻² at -1.0 V vs. RHE in 100 mg L⁻¹ NO₃⁻ + 0.5 M Na₂SO₄.⁷²

Wang *et al.* reported a heterostructure of an FeS₂ nanoparticle-decorated TiO₂ nanobelt array supported on a titanium plate (FeS₂@TiO₂/TP), as depicted in Fig. 5e and f. They proved that the abundant heterostructures endowed FeS₂@TiO₂/TP with excellent electrocatalytic performance with an NH₃ yield of 860.3 mmol h⁻¹ cm⁻² and a FE_{NH₃} of 97% at -0.4 V vs. RHE in 0.1 M NaOH + 0.1 M NO₃⁻. Furthermore, they elucidated the positive effect of the FeS₂@TiO₂ heterostructure by DFT studies. The charge density difference and partial density of states suggested that the electrons were spontaneously transferred from TiO₂ to FeS₂ (Fig. 5g). The corresponding charge accumulation and depletion endowed the FeS₂@TiO₂ interface with local nucleophilic and electrophilic regions, thus favoring the adsorption of targeted species. Moreover, such a heterostructure achieves the targeted adsorption of molecules by electrostatic interaction, promoting the charge transfer between the active site and the O atom of NO₃⁻, resulting in the activation of the N–O bond.⁷³ Similar results are obtained for the CoS₂@TiO₂/TP heterostructure synthesized by Zhao *et al.* They utilized XPS to prove the electron transfer from TiO₂ to CoS₂, and the heterostructure eventually resulted in an improved electrocatalytic performance of CoS₂@TiO₂/TP with a high FE_{NH₃} and NH₃ yield rate of 92.80% and 538.21 mmol h⁻¹ cm⁻², respectively.⁷⁴

Based on the conclusion that CoP is proven to be an effective and stable electrocatalyst for NO₃⁻-to-NH₃ conversion,⁷⁵ Deng *et al.* designed a heterostructure of CoP/TiO₂@TP, as depicted in Fig. 5h and i. Interestingly, XPS proved that the BEF of the CoP/TiO₂ heterostructure was established by the electrons transferring from CoP to TiO₂, which is reversed for the heterostructures mentioned above. This phenomenon should be attributed to the fact that CoP exhibits the metallic character of a continuous electron occupation state at the Fermi level, and the work function of CoP should be lower than that of TiO₂ with semiconductor properties. Moreover, the heterostructure displayed an impurity level near the Fermi level compared with CoP and TiO₂, indicating that more charge carriers were created (Fig. 5j), thereby enhancing the electron conductivity of CoP/TiO₂. The Gibbs free energy

diagram suggested that the potential-determining step of conversion of NO₃⁻ to *NO₃ presented a lower energy barrier in the CoP/TiO₂ heterostructure. Therefore, such CoP/TiO₂@TP attained an excellent FE_{NH₃} of 95.0% with a large NH₃ of 499.8 μmol h⁻¹ cm⁻².⁷⁶

4.3. New generation Ti-based MXene and single atomic catalysts

In addition to the traditional metallic Ti- and TiO₂-based nanomaterials, MXene attracts much attention in electrocatalysis. Moreover, single atomic catalysts (SACs) are emerging as fantastic materials due to their ultra-high active site utilization and outstanding catalytic activity. Therefore, developing Ti-based MXene and SACs for electrochemical NO₃⁻-to-NH₃ conversion could be a prevailing trend, although related studies are still rarely reported. In this section, we summarized some preliminary experimental and theoretical achievements of Ti-based MXene and SACs for the NO₃⁻RR and hope to provide inspiration for designing novel Ti-based electrocatalysts.

4.3.1. Ti-based MXene for the NO₃⁻RR. Transition metal carbides, nitrides, and carbonitrides with 2D structures, known as MXenes, were first reported in 2011 and received much attention due to their advantages of large surface area, structural flexibility, and surface tunable chemistry.⁷⁷ MXenes are commonly synthesized by top-down etching of layered ternary MAX phases with the formula M_{n+1}AX_n (*n* = 1–4) as precursors. Here, “M” represents early transition metals, “A” is the elements of group 13–14, and “X” is C or/and N. In the MAX precursors, the edge-sharing octahedron of [MX₆] extends laterally to form an “M–X” layer, while “A” layers are located on both sides of the “M–X” structures to form a sandwich-like structure. By removing the “A” layers from the MAX phase, 2D MXenes with alternately arranged structures “M” and “X” were formed. Moreover, -F, -OH, -Cl could also be coordinated to the surface of the M layer to form abundant surface terminations (denoted as T_x). Therefore, MXene is always written as M₃C₂T_x.⁷⁸ Ti₃C₂T_x is the most widely studied MXene.

Li's group investigated the possible mechanism and catalytic activity of electrochemical NO₃⁻-to-NH₃ conversion on Ti₃C₂ MXene by theoretical calculations. They proposed that the reactive pathway of NO₃⁻ → *NO₃ → *NO₂ → *NO → *N → *NH → *NH₂ → *NH₃ → NH₃(g) is thermodynamically preferable to that of NO₃⁻ → *NO₃ → *NO₂ → *NO → *NOH → *NHOH → *NH → *NH₂ → *NH₃ → NH₃(g). *NO₃ and *NO₂ prefer to adsorb on Ti₃C₂ MXene in parallel adsorption modes due to the O atoms of *NO₃ and *NO₂ being inclined to bond with Ti, resulting in drastic distortion of *NO₃ and *NO₂ (Fig. 6a). Moreover, the Gibbs free energy suggests that the NO₃⁻RR prefers to occur on the basal plane rather than the edge plane of Ti₃C₂ MXene because basal plane sites present lower energy barriers for the potential determining step (Fig. 6b). The termination effects of Ti₃C₂T₂ (T = O, OH, H, Cl, F, representing the terminal atoms located at the surface of Ti₃C₂ MXene) for NO₃⁻-to-NH₃ conversion were also elucidated in detail. As illustrated in Fig. 6c, the NO₃⁻RR is more competitive than the HER on the O-terminated Ti₃C₂O₂ with OVs



Fig. 6 (a) Gibbs free energy diagram of NO_3^- -to- NH_3 on Ti_3C_2 in vertical and parallel $^*\text{NO}_3$ adsorption modes. The grey zone means that $^*\text{NH}_2 \rightarrow ^*\text{NH}_3$ is the RDS. (b) Gibbs free energy diagram of NO_3^- -to- NH_3 on the basal plane and edge plane of Ti_3C_2 . (c) Gibbs free energy diagram of the reaction pathway of $\text{Ti}_3\text{C}_2\text{T}_x$, and the grey zone highlights the step $^*\text{NH}_2 \rightarrow ^*\text{NH}_3$ as the RDS.⁷⁹ Reproduced from ref. 79 with permission from The Royal Society of Chemistry, copyright 2022. (d) NO_3^- RR volcano plot of TM/g-CN with a descriptor of $\Delta G_{+\text{NO}_3}$. (e) Contour plot of limiting potential as a function of two potential-determining steps ($^*\text{NO} + \text{H}^+ + \text{e}^- \rightarrow ^*\text{NOH}$ and $^*\text{NH}_2 + \text{H}^+ + \text{e}^- \rightarrow ^*\text{NH}_3$). The promising zone is highlighted.⁴⁴ Reproduced from ref. 44 with permission from Wiley-VCH, copyright 2020. (f) Gibbs free energy change of NO_3^- adsorption on $\text{Ti}_3\text{C}_2\text{O}_2\text{-TM}_{5\text{A}}$. (g) Summary of limiting potentials on $\text{Ti}_3\text{C}_2\text{O}_2\text{-TM}_{5\text{A}}$.⁸¹ Reproduced from ref. 81 with permission from Springer Nature, copyright 2023. (h) Schematic illustration of preparing the $\text{Ru}_1\text{-TiO}_x/\text{Ti}$ electrode via an inherent surface oxide anchoring strategy. (i) (i and ii) Low-magnified HAADF-STEM images of $\text{Ru}_1\text{-TiO}_x/\text{Ti}$. (iii) Atomic-resolution HAADF-STEM image of $\text{Ru}_1\text{-TiO}_x/\text{Ti}$; Ru single atoms are indicated by white circles. (iv) Line intensity profile (inset) taken along the yellow dashed rectangle in HAADF-STEM and the corresponding 3D surface intensity profile. (j) Schematic illustration of the work functions of $\text{Ru}_1\text{-TiO}_x/\text{Ti}$ and TiO_x/Ti .⁸² Reproduced from ref. 82 with permission from Wiley-VCH, copyright 2022.

(denoted as $\text{Ti}_3\text{C}_2\text{O}_2\text{-O}_v$), and all free energy changes of NO_3^- -to- NH_3 on the $\text{Ti}_3\text{C}_2\text{O}_2\text{-O}_v$ are negative, signifying that the termination effect of oxygen promotes the hydrogenation of nitrogen-containing intermediates.⁷⁹ Another study utilized $\text{Ti}_3\text{C}_2\text{T}_x$ nanosheets as a promising molecular catalyst substrate. CuPc can be spontaneously dispersed on delaminated $\text{Ti}_3\text{C}_2\text{T}_x$ to overcome their large π -conjugated stacking. The prepared CuPc@MXene exhibited a high yield rate and selectivity for NH_3 , which is superior to other counterparts of FePc@MXene, NiPc@MXene, and CoPc@MXene. DFT calculations suggested that NO_3^- -to- NH_3 conversion tends to obey the ONH pathway

because the Gibbs free energy required only 0.27 eV to form $^*\text{ONH}$.⁸⁰

4.3.2. Ti-based SACs for the NO_3^- RR. Recently, SACs have become a frontier of research due to their outstanding catalytic performance and relatively simple coordinative structure. A typical structure of SACs denoted as M- N_x/C is a transition metal atomic site singly separated on nitrogen (N)-doped carbon (C)-based substrates.⁸³⁻⁸⁶ Moreover, metallic oxides,⁸⁷ phosphides,⁸⁸ and sulfides⁸⁹ could also be used as substrates for anchoring the atomic metal sites. Recently, significant advances in SACs were achieved in water splitting and CO_2

reduction. However, Ti-based SACs designed for NO_3^- -to- NH_3 conversion are currently rare. In this section, we summarized a theoretical prediction of the catalytic activity of Ti SAs and Ti-based substrates for anchoring the noble metal SAs, to provide some inspiration for the design of Ti-based SACs for the NO_3^- -RR.

Niu *et al.* established several transition metal atoms from Ti to Au supported on graphitic carbon nitrides (g-CN) as representatives (TM/g-CN) to study the NO_3^- -RR rule of SACs. They found that the adsorption energies of NO_3^- ($\Delta G_{\text{NO}_3^-}$), protons (ΔG_{H^+}), and the N_2 molecule (ΔG_{N_2}) on TM/g-CN exhibited periodic regulations. Furthermore, they proposed that the N-end pathway is the most reasonable on TM/g-CN because the N-end adsorption of NO presents the lowest energy. Based on the above reactive pathway, the performance criteria of limiting potential on TM/g-CN were evaluated. A volcano plot of limiting potential vs. $\Delta G_{\text{NO}_3^-}$ on TM/g-CN was proposed. As shown in Fig. 6d, too strong adsorption of NO_3^- results in a larger energy barrier in the step from $^*\text{NH}_2$ to $^*\text{NH}_3$. In contrast, too weak adsorption leads to a sluggish step from $^*\text{NO}$ to $^*\text{NOH}$, and Ti/g-CN and Zr/g-CN stand exactly near the top of the volcano. Regarding selectivity, the large energy barriers prevented the formation of NO_2 , NO, N_2O , and N_2 on Ti/g-CN, guaranteeing a high FE_{NH_3} of Ti/g-CN. In addition, the linear scaling relationship between ($\Delta G_{\text{NH}_3} - \Delta G_{\text{NH}_2}$) and ($\Delta G_{\text{NOH}} - \Delta G_{\text{NO}}$) hinders the exploration of more effective electrocatalysts (Fig. 6e), hence, a critical strategy to break such a scaling relationship is desired.⁴⁴ Another study that predicts TM/g- C_3N_4 (TM = Sc to Au, except Tc, Cd, and Hg) for the NO_3^- -RR also proposes a similar volcano plot and suggests that Ti/g- C_3N_4 should be an ideal candidate with outstanding limiting potential and FE_{NH_3} .⁹⁰

Recently, Ti-based MXene and oxides are also predicted to be suitable substrates for anchoring SAs. Wang *et al.* screened out 30 single transition metal atoms (3d: Sc–Zn, 4d: Y–Cd, and 5d: La–Hg) anchored on $\text{Ti}_3\text{C}_2\text{O}_2$ by means of first principles calculations. They found that SAs with half-filled d orbitals (around d^5) are favorable for NO_3^- activation due to high electronic states at the Fermi level (Fig. 6f). Further screening suggested that Cr, Re and Os SAs anchored on $\text{Ti}_3\text{C}_2\text{O}_2$ present negative free energy changes in the whole NO_3^- -to- NH_3 conversion process and thus exhibit positive limiting potentials (Fig. 6g); hence, they are recognized as the most efficient candidates.⁶⁶

An experimental study utilized Ti foil as a substrate to prepare binder-free monolithic single-atom electrodes (MSAEs). Metallic Ti inevitably undergoes oxidation in air to form a thin metal oxide layer (TiO_x) with abundant dangling bonds and defects, thereby providing unique on-site hosts for the dispersion of SACs. A two-step annealing approach depicted in Fig. 6h was employed to successfully anchor Ru atoms into the TiO_x layer (Fig. 6i). The as-prepared MSAEs were named $\text{Ru}_1\text{-TiO}_x/\text{Ti}$. Anchoring of Ru atoms upshifted the Fermi level, as proven by the lower work function of $\text{Ru}_1\text{-TiO}_x/\text{Ti}$ compared to TiO_x/Ti , as shown in Fig. 6j. This characteristic endowed $\text{Ru}_1\text{-TiO}_x/\text{Ti}$ with much improved catalytic current

density and FE_{NH_3} at various applied potentials. The highest FE_{NH_3} and NH_3 yield rate of 87.6% and 22.2 mol g^{-1} were respectively achieved at -0.3 V vs. RHE for $\text{Ru}_1\text{-TiO}_x/\text{Ti}$.⁸²

Although DFT studies predicted Ti-based SACs to be outstanding candidates as electrocatalysts for NO_3^- -to- NH_3 conversion, there is still a lack of experimental achievement to fulfill the above results. Further synthesis strategies and performance studies should be conducted.

5. Conclusion and perspective

Emerging “hydrogen economy” and “carbon neutralization” have caused NH_3 to be regarded as a carbon-free fuel and a portable energy carrier, endowing the electrochemical nitrate reduction reaction for NO_3^- -to- NH_3 conversion with a new opportunity for “green ammonia” synthesis. Compared to noble metal-based electrocatalysts, Ti-based electrocatalysts mainly composed of metallic Ti- and TiO_2 -based nanomaterials are attractive due to their inherent advantages of being nontoxic and cost-effective and having outstanding stability and HER inertness. Fundamental studies revealed that metallic Ti and TiO_2 are more suitable for NO_3^- -to- NH_3 conversion over a wide pH range. Many strategies including morphological design, heteroatom doping, creation of OVs, and the construction of heterostructures have been developed for reducing the overpotential while improving the FE_{NH_3} . To sum up, regulating the work function (for metallic Ti) or ionizing energy (for TiO_2) to reduce the gap between the HOS and the LUMO of nitrogen-containing intermediates is essential for reducing the overpotential, modulating the d-band structure to optimize the binding energy of O- and N-containing intermediates is crucial for regulating the reactive route and faradaic efficiency of NH_3 , and designing secondary active sites for splitting H_2O on Ti-based nanomaterials is also crucial for the hydrogenation of NO_3^- and consequent intermediates in neutral and alkaline electrolytes. In particular, since potential induced self-reconstruction exists for Ti-based electrocatalysts, the use of *in situ* characterization techniques is encouraged to reveal the real active sites for the NO_3^- -RR.

Finally, the difficulties and opportunities of Ti-based nanocatalysts for electrochemical NO_3^- -to- NH_3 conversion are discussed.

The large bandgap of TiO_2 (~ 3 eV) results in poor electronic conductivity which is the major reason for low electrochemical activity. Narrowing the bandgap of TiO_2 is an essential strategy to solve the problem of electronic conductivity. Therefore, characterizing the bandgap by UV-Vis spectroscopy, UPS, or DFT calculations is suggested to be performed for TiO_2 -based nanomaterials. Moreover, constructing metal/ TiO_2 heterostructures (such as Ti/TiO_2) and *in situ* compositing the TiO_2 -based nanomaterials with electronic conductive additives such as graphene, carbon nanotubes or acetylene black are effective strategies for solving the electronic conductivity issue. Recently, novel Ti-based MXenes with a large specific area and outstanding electron conductivity have been proposed as ideal

candidates for nitrate-to-ammonia conversion; thus developing Ti-based MXene electrocatalysts is an important development direction in the future.

The effect of lattice facets is rarely considered for Ti-based nanomaterials for the NO_3^- RR. Different lattice facets present unique atomic arrangements, signifying that the nanostructure with different exposed lattice facets exhibits disparate catalytic performance. Typically, high-index facets represent better catalytic activity. Therefore, exploring the effect of the lattice facets of Ti-based nanomaterials on the catalytic performance of the NO_3^- RR is important for designing Ti-based nanomaterials with desired exposed lattice facets, reducing the overpotential and improving the FE_{NH_3} .

Although heteroatom doping is a common modification strategy for inorganic materials, OV s in TiO_2 could also be simultaneously formed by doping metallic atoms with lower valence than Ti^{4+} . Therefore, doping-induced OV s should be considered but unfortunately, they are often neglected.

In addition to NH_3 , producing high value-added organic molecules with deeply reduced nitrogen functional groups (such as urea,^{91,92} methylamine,⁹³ formamide,⁹⁴ glycine,⁹⁵ etc.) by electrochemical reduction of NO_3^- is also an emerging direction. Since TiO_2 -based nanomaterials exhibit profound reduction ability of NO_3^- , some preliminary studies successfully coupled the CO_2 reduction reaction and the NO_3^- RR on TiO_2 -based nanomaterials to produce urea.⁹¹ Zhang *et al.* revealed that constructing secondary active sites on TiO_2 for adsorbing carbon-containing intermediates is a feasible strategy for the gC–N coupled reaction.⁹²

We hope that this review will shed light on the exploration of Ti-based nanocatalysts and pave the way for low-cost and efficient Ti-based nanocatalysts to realize large-scale industrial applications.

Conflicts of interest

The authors declare no competing financial interest.

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