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Aqueous electrocatalytic N₂ reduction for ambient NH₃ synthesis: recent advances in catalyst development and performance improvement

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Electrochemical N₂ reduction has emerged as an environmentally benign alternative to the Haber–Bosch process for sustainable NH₃ synthesis under ambient reaction conditions, and considerable recent attention has focused on electrocatalytic NH₃ synthesis from N₂ and H₂O in aqueous media. In this Minireview, we summarize the recent advances in the development of electrocatalysts for the N₂ reduction reaction (NRR). Strategies to boost the NRR performances are also discussed. Perspectives for further research directions are provided finally.

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1. Introduction

NH₃ is not only widely utilized as an essential activated nitrogen source to manufacture agricultural fertilizers, dyes, polymers, explosives, *etc.*, but also provides a carbon-free chemical energy carrier solution for the transportation sector.^{1–3} As the dominant route for industrial-scale NH₃ production using N₂ and H₂ as the feed gases, the century-old Haber–Bosch process suffers from harsh reaction conditions, complicated factory infrastructure, high energy consumption, and serious CO₂ emissions.⁴ One alternative approach to solve the NH₃ synthesis problem is to use electricity to drive the NH₃ production reaction,⁵ and this electrically driven process is compatible with intermittent operation and enables utilization of renewable electricity without needing transmission capacity expansion. Electrochemical N₂ reduction has emerged as an attractive method for artificial N₂-to-NH₃ conversion under ambient conditions; however it is severely challenged by N₂ activation and needs efficient electrocatalysts to break the rather inert molecular structure of N₂ with an extremely high bond energy of about 941 kJ mol⁻¹.^{6–11} One big issue for the NRR in aqueous electrolytes is that the competitive hydrogen evolution reaction (HER) limits its current efficiency and leads to a low overall reaction rate (Fig. 1).¹² Although they have better NH₃ selectivity, molecular catalysts are fragile, which can be circumvented by using heterogeneous NRR catalysts.

Two basic mechanisms (Scheme 1) are involved in the NRR over heterogeneous catalysts: the dissociative and associative mechanisms.¹³ In the dissociative mechanism, the N≡N triple bond is broken before its hydrogenation, leaving individual N-adatoms on the catalyst surface which are transformed into NH₃ independently (Scheme 1a). The NRR based on the associative mechanism proceeds *via* an alternating (Scheme 1b) and a distal (Scheme 1c) pathway. For the alternating pathway, each

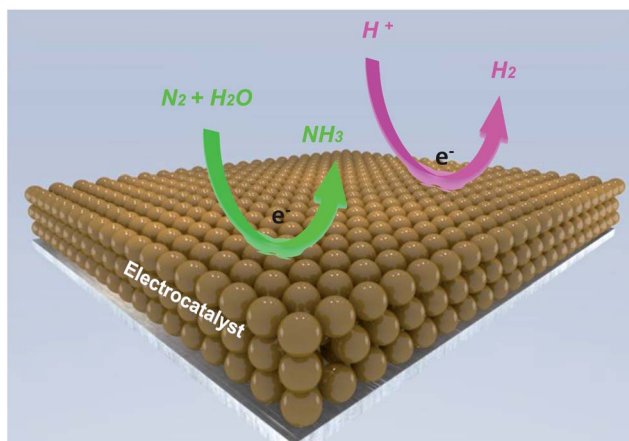
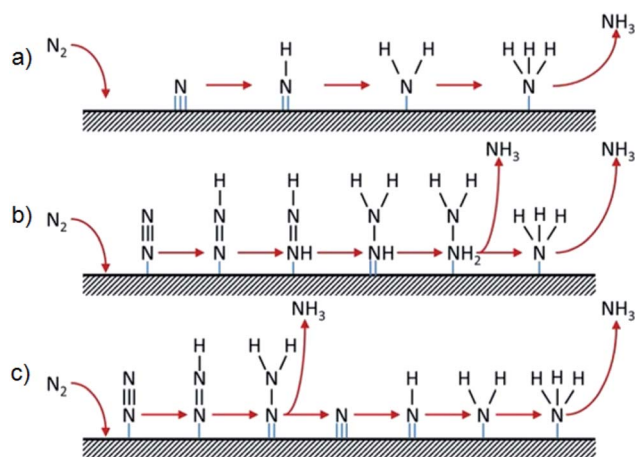


Fig. 1 Competition between the electrocatalytic NRR and HER processes on the catalyst. The NRR process involves the transfer of six electrons for one N_2 molecule: $N_2(g) + 6H_2O(l) + 6e^- \rightleftharpoons 2NH_3(aq.) + 6OH^-(aq.)$ ($E^\circ = 0.092$ V vs. the reversible hydrogen electrode, RHE). In this process, N_2 reacts with protons from the electrolyte on the surface of the catalyst to form NH_3 using electrons from the electrode. Because of the involvement of only two electrons per H_2 molecule, $2H^+(aq.) + 2e^- \rightleftharpoons H_2(g)$ ($E^\circ = 0.00$ V vs. RHE), the HER is kinetically preferred over the multi-step six-electron NRR process.



Scheme 1 Generic NRR mechanisms on heterogeneous catalysts: (a) dissociative pathway; (b) associative alternating pathway; and (c) associative distal pathway. Reproduced from ref. 13 with permission from Elsevier, copyright 2018.

of the two N atoms of adsorbed N_2 is hydrogenated in turn until one N atom is converted into NH_3 and the triple bond is broken. For the distal one, preferential hydrogenation of the N atom furthest away from the surface releases one equivalent of NH_3 , followed by the hydrogenation of the other N-atom to release a second equivalent of NH_3 .

In this Minireview, we summarize the recent advances in developing NRR electrocatalysts including noble-metal, non-noble-metal and non-metal catalysts. Following this, we discuss the strategies to boost the NRR performances including electronic structure tailoring, active site enrichment and HER

suppression. Finally, future research directions are also proposed in the concluding remarks.

2. Advances in NRR electrocatalysts

2.1. Noble-metal catalyst

Based on previous density functional theory (DFT) calculations predicting a stronger binding of intermediates to the stepped facets than to the flat terraces on a N_2 -fixing metal catalyst,¹⁴ Yan and co-workers demonstrated a proof-of-concept that a Au nanorod with a tetrahedral structure is capable of catalyzing the NRR with an NH_3 yield of $1.648 \mu g h^{-1} cm^{-2}$ and a faradaic efficiency (FE) of 4.02%.¹⁵ Wang *et al.* reported that Pd/C is superior in NRR activity to Au and Pt in phosphate buffer solution (PBS), attaining a FE of 8.2%.¹⁶ Theoretical calculations further reveal that the *in situ* formed α -PdH enables N_2 activation through a thermodynamically more favorable Grothuss-like hydride transfer pathway compared to direct surface hydrogenation or proton-coupled electron transfer steps. Ru is industrially used as an alternative to conventional Fe catalysts for more efficient NH_3 synthesis at lower temperatures.¹⁷ Recent work also shows that Ru nanoparticles perform efficiently in ambient electrocatalytic N_2 reduction to NH_3 ($5.5 mg h^{-1} m^{-2}$; 5.4%).¹⁸ Of note, Ag nanosheets also provides quite positive results (4.8%).¹⁹

2.2. Non-noble-metal catalysts

Compared with noble-metal materials, non-noble-metal ones have much higher earth abundance and hence hold greater promise for application as attractive NRR catalysts. Biological nitrogenases containing Mo, Fe and V are involved in natural N_2 fixation,^{20,21} and a variety of NRR electrocatalysts made of these metals have been developed for artificial electrochemical NH_3 synthesis under ambient conditions.

Inspired by the fact that Mo and S elements play significant roles in nitrogenases, we performed theoretical calculations to study the electronic structures of MoS_2 and mapped out the energy profile of the NRR on MoS_2 , which suggests that the positively charged Mo-edge is the key to polarizing and activating the N_2 molecules (Fig. 2).²² To prove this, we hydrothermally grew a MoS_2 nanosheet array on carbon cloth (MoS_2/CC) to catalyze the NRR with a FE of 1.17%. Interestingly, other Mo compounds are also active for the NRR. For instance, an MoN catalyst can suppress electron transfer from Mo to the adsorbed N_2 , facilitating the release of produced NH_3 and thus enhancing the NRR performance, and MoN nanosheets achieve an NH_3 yield of $3.01 \times 10^{-10} mol s^{-1} cm^{-2}$ with a FE of 1.15%.²³ Superior NRR performances have been reported for Mo_2C nanorods²⁴ and Mo_2C nanodots embedded in ultrathin carbon nanosheets²⁵ with FEs of 8.13% and 7.8%, respectively. The exposed Mo atoms of the $Mo_2C(121)$ surface have large adsorption energies to dissociate N_2 and the subsequent exothermic hydrogenation reactions are energetically preferable *via* the alternating pathway.²⁵ Compared with the above Mo-based catalysts, its oxide counterpart can be more easily prepared on a large scale, and our study also verifies that MoO_3

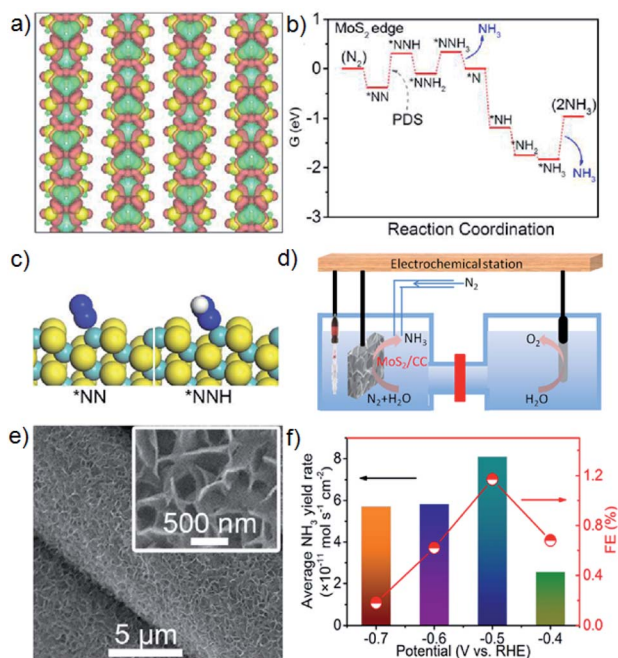


Fig. 2 (a) Isosurface of deformation charge density as viewed from the top. Red and green represent charge accumulation and loss, respectively. The isosurface is 0.0025 a.u. (b) Free energy profiles for the NRR at the MoS₂ edge site. The asterisk (*) denotes the adsorption site. (c) Structures of key intermediates of the potential-determining step (PDS). (d) A schematic diagram to illustrate the electrochemical setup for the NRR tests. (e) Scanning electron microscopy (SEM) images for MoS₂/CC. (f) Average NH₃ yields and FEs of MoS₂/CC at different potentials in 0.1 M Na₂SO₄. Reproduced from ref. 22 with permission from Wiley-VCH, copyright 2018.

nanosheets are active for electrocatalytic N₂-to-NH₃ fixation ($4.80 \times 10^{-10} \text{ mol s}^{-1} \text{ cm}^{-2}$; 1.9%).²⁶

As the cheapest and one of the most abundant transition metals, Fe-based NRR catalysts have also been widely explored. Licht *et al.* reported efficient N₂ reduction using Fe₂O₃ in a molten hydroxide electrolyte cell at temperatures ≥ 200 °C.²⁷ At room temperature and pressure, however, only a low FE of 0.15% was obtained for Fe₂O₃ nanoparticles supported on carbon nanotubes in a flow electrochemical cell operating in the gas phase.²⁸ Since then, much effort has been made to explore Fe oxides with higher NRR activities in aqueous media.^{29–31} An Fe₃S₄ nanocatalyst was also shown by Zhao *et al.* to be active with a FE of 6.45% for NH₃ formation.³² For the first time, we demonstrate that FeOOH nanorods achieve a FE of 6.7%, and DFT calculations evidence that the Fe-edge atom of the FeOOH(110) surface plays a key role in polarizing and activating the N₂ molecules for both charge exchange and transfer.³³ Quite surprisingly, a more recent study by us suggests that a P-rich FeP₂ nanoparticle-reduced graphene oxide hybrid exhibits superhigh performances with a large NH₃ yield of $35.26 \mu\text{g h}^{-1} \text{ mg}_{\text{cat.}}^{-1}$ and a high FE of 21.99%.³⁴ DFT calculations reveal decreased HER activity, higher N₂ adsorption energy, and a larger number of NRR active sites for FeP₂ compared to FeP.

VN has previously been identified by DFT analysis as a promising candidate for the NRR under ambient conditions,³⁵ but only recently has this prediction been experimentally verified.^{36–38} For transition metal nitrides, a Mars-van Krevelen mechanism is considered for NH₃ generation: a surface N atom is reduced to form NH₃ after which the resulting vacancy is replenished by a N₂ molecule from the electrolyte. The N-vacancy must be stable at the surface, thus avoiding migration into the bulk of the catalyst. If not so, the reacted N on the surface is only replaced with more N atoms from the catalyst itself until all the N atoms of the metal nitrides are completely depleted, leaving only pure metal. V oxides are also able to catalyze the NRR.^{39,40} It should be noted that many other metals without biological implications, including Ti,^{41–43} Nb,^{44,45} Cr,⁴⁶ Mn,⁴⁷ Co,⁴⁸ Cu,⁴⁹ Bi,^{50,51} Sn,⁵² La,⁵³ *etc.*, have also been identified to actively electrocatalyze the reduction.



Fig. 3 Transmission electron microscopy (TEM) images of non-metal catalysts: (a) B nanosheet. Reproduced from ref. 56 with permission from The Royal Society of Chemistry, copyright 2019; (b) black P nanosheet. Reproduced from ref. 57 with permission from Wiley-VCH, copyright 2019; (c) B₄C nanosheet. Reproduced from ref. 58 with permission from the Nature Publishing Group, copyright 2018; (d) BP nanoparticles. Reproduced from ref. 59 with permission from The Royal Society of Chemistry, copyright 2019; (f) PCN. Reproduced from ref. 62 with permission from Wiley-VCH, copyright 2018; (e) atomic force microscopy image and corresponding height profile for BN nanosheets. Reproduced from ref. 61 with permission from Tsinghua University Press and Springer-Verlag, copyright 2019.

Table 1 Performance metrics of aqueous-based NRR electrocatalysts under ambient conditions

Catalyst development	Catalyst	Electrolyte	Potential (V) vs. RHE	NH ₃ yield	FE (%)	Ref.
Noble-metal catalysts	Au nanorods	0.10 M KOH	-0.2	1.648 $\mu\text{g h}^{-1} \text{cm}^{-2}$	4.02	<i>Adv. Mater.</i> , 2017, 29, 1604799
	Pd/C	0.01 M HCl		4.5 $\mu\text{g h}^{-1} \text{mg}_{\text{Pd}}^{-1}$ (-0.05 V)	8.2 (0.1 V)	<i>Nat. Commun.</i> , 2018, 9, 1795
	Ru nanoparticles	0.01 M HCl		5.5 $\mu\text{g h}^{-1} \text{cm}^{-2}$ (-0.1 V)	5.4 (0.01 V)	<i>ChemSusChem</i> , 2018, 11, 3416–3422
	Ag nanosheets	0.1 M HCl	-0.6	4.62 $\times 10^{-11} \text{ mol s}^{-1} \text{cm}^{-2}$	4.8	<i>Chem. Commun.</i> , 2018, 54, 11427–11430
	MoS ₂ nanosheet array	0.1 M Na ₂ SO ₄	-0.5	8.08 $\times 10^{-11} \text{ mol s}^{-1} \text{cm}^{-2}$	1.17	<i>Adv. Mater.</i> , 2018, 30, 1800191
	MoN nanosheet array	0.1 M HCl	-0.3	3.01 $\times 10^{-10} \text{ mol s}^{-1} \text{cm}^{-2}$	1.15	<i>ACS Sustainable Chem. Eng.</i> , 2018, 6, 9550–9554
	Mo ₂ C/C	0.5 M Li ₂ SO ₄	-0.3	11.3 $\mu\text{g h}^{-1} \text{mg}_{\text{Mo}_2\text{C}}^{-1}$	7.8	<i>ACS Mater.</i> , 2018, 30, 1803694
	Mo ₂ C nanorods	0.1 M HCl	-0.3	91.5 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	8.31	<i>ACS Central Sci.</i> , 2019, 5, 116–121
	MoO ₃ nanosheets	0.1 M HCl	-0.3	4.8 $\times 10^{-10} \text{ mol s}^{-1} \text{cm}^{-2}$ (-0.5 V)	1.9 (-0.3 V)	<i>J. Mater. Chem. A</i> , 2018, 6, 12974–12977
	Fe ₂ O ₃ -CNT	KHCO ₃	-2.0	0.22 $\mu\text{g h}^{-1} \text{cm}^{-2}$	0.15	<i>Angew. Chem., Int. Ed.</i> , 2017, 56, 2699–2703
Non-noble-metal catalysts	Fe/Fe ₃ O ₄	0.1 M PBS	-0.3	0.19 $\mu\text{g h}^{-1} \text{cm}^{-2}$	8.29	<i>ACS Catal.</i> , 2018, 8, 9312–9319
	Fe ₂ O ₃ nanorods	0.1 M Na ₂ SO ₄	-0.8	15.9 $\mu\text{g h}^{-1} \text{mg}^{-1}$	0.94	<i>ChemCatChem</i> , 2018, 10, 4530–4535
	Fe ₃ O ₄ nanorods	0.1 M Na ₂ SO ₄	-0.4	5.6 $\times 10^{-11} \text{ mol s}^{-1} \text{cm}^{-2}$	2.6	<i>Nanoscale</i> , 2018, 10, 14386–14389
	β -FeOOH nanorods	0.5 M LiClO ₄	-0.4	23.32 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$ (-0.75 V)	6.7 (-0.7 V)	<i>Chem. Commun.</i> , 2018, 54, 11332–11335
	Fe ₃ S ₄ nanosheets	0.1 M HCl	-0.4	75.4 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	6.45	<i>Chem. Commun.</i> , 2018, 54, 13010–13013
	VN nanoparticles	0.05 M H ₂ SO ₄	-0.1	3.3 $\times 10^{-10} \text{ mol s}^{-1} \text{cm}^{-2}$	6.0	<i>J. Am. Chem. Soc.</i> , 2018, 140, 13387–13391
	VN nanosheet array	0.1 M HCl	-0.5	8.4 $\times 10^{-11} \text{ mol s}^{-1} \text{cm}^{-2}$	2.25	<i>ACS Sustainable Chem. Eng.</i> , 2018, 6, 9545–9549
	TiO ₂ nanosheet array	0.1 M Na ₂ SO ₄	-0.7	9.16 $\times 10^{-11} \text{ mol s}^{-1} \text{cm}^{-2}$	2.5	<i>ACS Appl. Mater. Interfaces</i> , 2018, 10, 28251–28255
	Ti ₃ C ₂ T _x nanosheets	0.1 M HCl	-0.4	20.4 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	9.3	<i>J. Mater. Chem. A</i> , 2018, 6, 24031–24035
	Ti ₃ C ₂ T _x /FeOOH	0.05 M H ₂ SO ₄	-0.4	0.53 $\mu\text{g h}^{-1} \text{cm}^{-2}$ (-0.5 V)	5.78 (-0.2 V)	<i>Joule</i> , 2019, 3, 279–289
Non-metal catalysts	NbO ₂ nanoparticles	0.05 M H ₂ SO ₄	-0.65	11.6 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	32	<i>Small Methods</i> , 2019, 3, 1800386
	Nb ₂ O ₅ nanofibers	0.1 M HCl	-0.55	43.6 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	9.26	<i>Nano Energy</i> , 2018, 52, 264–270
	Hollow Cr ₂ O ₃ microspheres	0.1 M Na ₂ SO ₄	-0.9	25.3 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	6.78	<i>ACS Catal.</i> , 2018, 8, 8540–8544
	MnO particles	0.1 M Na ₂ SO ₄	-0.39	1.11 $\times 10^{-10} \text{ mol s}^{-1} \text{cm}^{-2}$	8.02	<i>Adv. Sci.</i> , 2019, 6, 1801182
	LaF ₃ nanoplates	0.5 M LiClO ₄	-0.45	55.9 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	16.0	<i>J. Mater. Chem. A</i> , 2019, 7, 17761–17765
	Mosaic Bi nanosheets	0.1 M Na ₂ SO ₄	-0.8	2.54 $\pm 0.16 \mu\text{g}_{\text{NH}_3} \text{cm}^{-2} \text{h}^{-1}$	10.46 ± 1.45	<i>ACS Catal.</i> , 2019, 9, 2902–2908
	Bi nanosheet array	0.1 M HCl	-0.5	6.89 $\times 10^{-11} \text{ mol s}^{-1} \text{cm}^{-2}$	10.26	<i>Chem. Commun.</i> , 2019, 55, 5263–526
	CuO/rGO	0.1 M Na ₂ SO ₄	-0.75	1.8 $\times 10^{-10} \text{ mol s}^{-1} \text{cm}^{-2}$	3.9%	<i>ChemCatChem</i> , 2019, 11, 1441–1447
	B nanosheets	0.1 M HCl	-0.14	31.37 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	4.84	<i>Chem. Commun.</i> , 2019, 55, 4246–4249
	Black P nanosheets	0.1 M Na ₂ SO ₄	-0.8	13.22 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	4.04	<i>ACS Catal.</i> , 2019, 9, 4609–4615
Performance Tailoring electronic improvement structures	B ₃ C nanosheets	0.01 M HCl	-0.75	31.37 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$ (-0.7 V)	5.07 (-0.6 V)	<i>Angew. Chem., Int. Ed.</i> , 2019, 58, 2612–2616
	BP nanoparticles	0.1 M HCl	-0.6	26.42 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	15.95	<i>Nat. Commun.</i> , 2018, 9, 3485
	BN nanosheets	0.1 M HCl	-0.75	22.4 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	12.7	<i>J. Mater. Chem. A</i> , 2019, 7, 16117–16121
	Mesoporous BN	0.1 M Na ₂ SO ₄	-0.7	18.2 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	4.7	<i>Nano Res.</i> , 2019, 12, 919–924
	PCN	0.1 M HCl	-0.2	8.09 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	5.5	<i>Nanoscale</i> , 2019, 11, 4231–4235
	Defective TiO ₂	0.1 M HCl	-0.15	1.24 $\times 10^{-10} \text{ mol s}^{-1} \text{cm}^{-2}$	11.59	<i>Angew. Chem., Int. Ed.</i> , 2018, 57, 10246–10250
	MnO _x nanowire array	0.1 M Na ₂ SO ₄	-0.5	1.63 $\times 10^{-10} \text{ mol s}^{-1} \text{cm}^{-2}$	9.17	<i>Nanoscale</i> , 2019, 11, 1555–1562
	R-WO ₃ nanosheets	0.1 M HCl	-0.3	17.28 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	11.40	<i>Chem. Commun.</i> , 2019, 55, 4627–4630
	r-CeO ₂ nanorods	0.1 M Na ₂ SO ₄	-0.2	16.4 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$ (-0.5 V)	7.0	<i>Nanoscale</i> , 2019, 11, 19274–19277
	W ₂ N ₃ nanosheets	0.1 M KOH	-0.2	11.66 $\pm 0.98 \mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	3.7 (-0.4 V)	<i>ACS Sustainable Chem. Eng.</i> , 2019, 7, 2889–2893
Defect-rich MoS ₂ nanoflowers	0.1 M Na ₂ SO ₄	-0.4	29.28 $\mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$	11.67 ± 0.93	<i>Adv. Mater.</i> , 2019, 31, 1902709	
Defect-Bi nanoplates	0.2 M Na ₂ SO ₄	-0.6	5.453 $\mu\text{g h}^{-1} \text{mg}_{\text{Bi}}^{-1}$	8.34	<i>Adv. Energy Mater.</i> , 2018, 8, 1801357	
				11.6	<i>Angew. Chem., Int. Ed.</i> , 2019, 58, 9464–9469	

Table 1 (Contd.)

Catalyst	Electrolyte	Potential (V) vs. RHE	NH ₃ yield	FE (%)	Ref.
NPC	0.05 M H ₂ SO ₄	-0.9	1.40 mmol g ⁻¹ h ⁻¹	1.42	ACS Catal., 2018, 8, 1186-1191
NCM-Au NPs	0.1 M KOH	-0.5	0.36 g m ⁻² h ⁻¹ (-0.2 V)	22 (-0.1 V)	Angew. Chem., Int. Ed., 2018, 57, 12360-12364
B-Doped graphene	0.05 M H ₂ SO ₄	-0.5	9.8 μg h ⁻¹ cm ⁻²	10.8	Joule, 2018, 2, 1610-1622
O-Doped graphene	0.1 M HCl	-0.5	21.3 μg h ⁻¹ mg _{cat.} ⁻¹	12.6 (-0.45 V)	Chem. Commun., 2019, 55, 7502-7505
S-Doped graphene	0.1 M HCl	-0.5	27.3 μg h ⁻¹ mg _{cat.} ⁻¹ (-0.6 V)	11.5 (-0.5 V)	Chem. Commun., 2019, 55, 3371-3374
Defect-rich fluorographene	0.1 M Na ₂ SO ₄	-0.7	9.3 μg h ⁻¹ mg _{cat.} ⁻¹	4.2	Chem. Commun., 2019, 55, 4266-4269
FeO(OH,F) nanorods	0.5 M LiClO ₄	-0.6	42.38 μg h ⁻¹ mg _{cat.} ⁻¹	9.02	Chem. Commun., 2019, 55, 3987-3990
F-SnO ₂ nanosheets	0.1 M Na ₂ SO ₄	-0.45	19.3 μg h ⁻¹ mg _{cat.} ⁻¹	8.6	Inorg. Chem., 2019, 58, 10424-10431
Cu-CeO _{2-x} nanorods	0.1 M Na ₂ SO ₄	-0.45	5.3 × 10 ⁻¹⁰ mol s ⁻¹ cm ⁻²	19.1	Chem. Commun., 2019, 55, 2952-2955
C-Ti _x O _y /C	0.1 M LiClO ₄	-0.4	14.8 μg h ⁻¹ mg _{cat.} ⁻¹	17.8	Angew. Chem., Int. Ed., 2019, 58, 13101-13106
V-TiO ₂ nanorods	0.5 M LiClO ₄	-0.45	17.73 μg h ⁻¹ mg _{cat.} ⁻¹ (-0.5 V)	15.3 (-0.4 V)	Small Methods, 2019, 3, 1900355
Zr-TiO ₂ nanotubes	0.1 M KOH	-0.45	8.90 μg h ⁻¹ cm ⁻²	17.3	Nat. Commun., 2019, 10, 2877
Fe-TiO ₂ nanoparticles	0.5 M LiClO ₄	-0.4	25.47 μg h ⁻¹ mg _{cat.} ⁻¹	25.6	Angew. Chem., Int. Ed., 2019, 131, 18620-18624
O-CNT	0.1 M LiClO ₄	-0.4	32.33 μg h ⁻¹ mg _{cat.} ⁻¹	12.50	Chem. Commun., 2019, 55, 4997-5000
TA-rGO	0.5 M LiClO ₄	-0.75	17.02 μg h ⁻¹ mg _{cat.} ⁻¹	4.83	ACS Sustainable Chem. Eng., 2019, 7, 14368-14372
Pd-TA	0.1 M Na ₂ SO ₄	-0.45	24.12 μg h ⁻¹ mg _{cat.} ⁻¹	9.4	J. Mater. Chem. A, 2019, 7, 21674-21677
a-Au/CeO _x -rGO	0.1 M HCl	-0.2	8.31 μg h ⁻¹ mg _{cat.} ⁻¹	10.10	Adv. Mater., 2017, 29, 1700001
Bi ₄ V ₂ O ₁₁ /CeO ₂	0.1 M HCl	-0.2	23.21 μg h ⁻¹ mg _{cat.} ⁻¹	10.16	Angew. Chem., Int. Ed., 2018, 57, 6073-6073
Au ₆ /Ni	0.05 M H ₂ SO ₄	-0.14	7.4 μg h ⁻¹ mg _{cat.} ⁻¹	67.8	J. Am. Chem. Soc., 2019, 141, 14976-14980
Enriching active sites TA-reduced Au/TiO ₂	0.1 M HCl	-0.2	21.4 μg h ⁻¹ mg _{cat.} ⁻¹	8.11	Adv. Mater., 2017, 29, 1606550
Au flowers	0.1 M HCl	-0.2	25.57 μg h ⁻¹ mg _{cat.} ⁻¹	6.05	ChemSusChem, 2018, 11, 3480-3485
AuHNCs	0.5 M LiClO ₄	-0.9	3.9 μg h ⁻¹ cm ⁻² (-0.5 V)	30.2 (-0.4 V)	Nano Energy, 2018, 49, 316-323
TiO ₂ -rGO	0.1 M Na ₂ SO ₄	-0.85	15.13 μg h ⁻¹ mg _{cat.} ⁻¹	3.3	J. Mater. Chem. A, 2018, 6, 17303-17306
Mn ₃ O ₄ -rGO	0.1 M Na ₂ SO ₄	-0.85	17.4 μg h ⁻¹ mg _{cat.} ⁻¹	3.52	Nano Res., 2019, 12, 1093-1098
Cr ₂ O ₃ -rGO	0.1 M HCl	-0.5	33.3 μg h ⁻¹ mg _{cat.} ⁻¹ (-0.7 V)	7.33 (-0.6 V)	Inorg. Chem., 2019, 58, 2257-2260
SnO ₂ /rGO	0.1 M Na ₂ SO ₄	-0.7	25.6 μg h ⁻¹ mg _{cat.} ⁻¹	7.1	ACS Appl. Mater. Interfaces, 2019, 11, 31806-31815
Pd _{0.2} Cu _{0.8} -rGO	0.1 M KOH	-0.85	2.8 μg h ⁻¹ mg _{cat.} ⁻¹	0.6	Adv. Energy Mater., 2018, 8, 1800124
S dots-rGO	0.5 M LiClO ₄	-0.85	28.56 μg h ⁻¹ mg _{cat.} ⁻¹	7.07	Chem. Commun., 2019, 55, 3152-3155
PTCA-rGO	0.1 M HCl	-0.5	24.7 μg h ⁻¹ mg _{cat.} ⁻¹	6.9	J. Mater. Chem. A, 2019, 7, 12446-12450
MnO ₂ -Ti ₃ C ₂ T _x	0.1 M HCl	-0.55	34.12 μg h ⁻¹ mg _{cat.} ⁻¹	11.39	J. Mater. Chem. A, 2019, 7, 18 823-18 827
TiO ₂ /Ti ₃ C ₂ T _x	0.1 M HCl	-0.55	32.17 μg h ⁻¹ mg _{cat.} ⁻¹ (-0.55 V)	16.07 (-0.45 V)	Adv. Energy Mater., 2019, 9, 1803406
BP/SnO _{2-x} nanotubes	0.1 M Na ₂ SO ₄	-0.4	48.87 μg h ⁻¹ mg _{cat.} ⁻¹	14.6	Angew. Chem., Int. Ed., 2019, 131, 1-7
Au ₁ /C ₃ N ₄	0.05 M H ₂ SO ₄	-0.1	1.305 μg h ⁻¹ mg _{Au} ⁻¹	11.1	Sci. Bull., 2018, 63, 1246-1253
AuSAs-NDPCs	0.1 M HCl	-0.2	2.32 μg h ⁻¹ cm ⁻²	12.3	Small Methods, 2018, 2, 1800202
Ru SAs/N-C	0.05 M H ₂ SO ₄	-0.2	120.9 μg h ⁻¹ mg _{cat.} ⁻¹	29.6	Adv. Mater., 2018, 30, 1803498
SA-Mo/NPC	0.1 M KOH	-0.3	34.0 ± 3.6 μg h ⁻¹ mg _{cat.} ⁻¹	14.6 ± 1.6	Angew. Chem., Int. Ed., 2019, 58, 2321-2325
Enriching active sites FeS _N -C	0.1 M KOH	0.0	7.48 μg h ⁻¹ mg _{cat.} ⁻¹	56.55	Nat. Commun., 2019, 10, 341
PEBCD	0.5 M Li ₂ SO ₄	-0.2	2.01 μg h ⁻¹ cm ⁻² (-0.7 V)	2.91 (-0.4 V)	J. Am. Chem. Soc., 2017, 139, 9771-9774
MoS ₂ /BCCF	0.1 M Li ₂ SO ₄	-0.6	43.4 μg h ⁻¹ mg _{cat.} ⁻¹	9.81	Adv. Energy Mater., 2019, 9, 1803935
BiNCs	Acidic 0.5 M K ₂ SO ₄ (pH = 3.5)	-0.6	200 mmol g ⁻¹ h ⁻¹	66	Nat. Catal., 2019, 2, 448-456
Ag-Au@ZIF	0.1 M HCl	-2.5	10 pmol cm ⁻² s ⁻¹	18 ± 4	Sci. Adv., 2018, 4, eaar3208
NPG@ZIF-8	0.1 M Na ₂ SO ₄	-2.5	28.7 ± 0.9 μg h ⁻¹ cm ⁻² (-0.8 V)	44 (-0.6 V)	Angew. Chem., Int. Ed., 2019, 131, 15506-15510

2.3. Non-metal catalysts

From economic and environmental viewpoints, using non-metal catalysts could lower the cost and avoid the residues of metal ions. Nanocarbons, which are mainly composed of carbon and can even be made directly out of biomass, feature wide potential windows and structural diversity, which make them promising for electrochemical application as sustainable materials.⁵⁴ Unfortunately, pristine nanocarbons have poor NRR activities and effective strategies must be implemented to improve their catalytic performances (see the following section for details).

Interestingly, nanomaterials of elemental B^{55,56} and black P⁵⁷ perform efficiently in electrochemical NH₃ synthesis. Other non-metal materials including B₄C,⁵⁸ BP,⁵⁹ BN,^{60,61} and polymeric carbon nitride (PCN)⁶² are highly active NRR catalysts. Fig. 3 presents the morphologies and the performance metrics are detailed in Table 1.

3. Strategies for boosting the NRR performances

The catalytic performances of current catalysts are still far from meeting the needs of practical applications. From the point of view of thermodynamics, the NRR should proceed at negative potentials that are dominated by the competing HER, leading to a tough selectivity issue and thus unsatisfactory current efficiency. It is clear that strategies should be developed to enable more efficient N₂ reduction electrocatalysis. In the following section, we will summarize the most recent progress in strategies to boost the NRR performances, which include: (1) tailoring the electronic structures to promote intrinsic NRR activity; (2) enriching active sites to increase the apparent NRR activity; (3) suppressing the HER to improve the selectivity.

3.1. Tailoring the electronic structures of catalysts

Tailoring the electronic structure of a catalyst is the most common method to change the adsorption behaviour and tune the intrinsic activity of each active site.⁶³ Defect engineering and heteroatom doping have been proven to be the two most effective methods to boost the intrinsic activity of NRR electrocatalysts. Surface functionalization and interface engineering provide us with two other good choices for this purpose.

3.1.1 Defect engineering. O vacancies (V_O) can manipulate the electronic structure of metal oxides to achieve enhanced conductivity and could also act as the active sites to adsorb the reactants and reaction intermediates leading to a lowered activation energy barrier.⁶⁴ Li *et al.* reported that V_O in amorphous TiO₂ promote the adsorption and activation of N₂ facilitating N₂ photoreduction to NH₃.⁶⁵ Recently, we found that a TiO₂ nanosheet array can electrocatalyze the NRR with a FE of 2.50% in Na₂SO₄, and the *in situ* generated V_O during the NRR are supposed to be responsible for the enhanced adsorption and activation of N₂.⁴¹ To gain further experimental and theoretical support for this, V_O were intentionally introduced into TiO₂ *via* cathodic polarization, and the resulting defective TiO₂ achieves a much higher FE (9.17%) than the pristine one (0.95%) in

acid.⁶⁶ Such an enhancement is also observed for other metal oxides like MnO₂,⁶⁷ WO₃,⁶⁸ Fe₂O₃ (ref. 69) and CeO₂.⁷⁰

The Yu group reported experimentally and theoretically that electrochemical N₂ reduction can be achieved by using PCN with N vacancies (V_N) which chemisorbs and significantly increases the bond length of N₂ resulting in strong N₂ activation.⁶² Because of the lack of ¹⁵N₂ isotopic experiments, one cannot confirm that the N of NH₃ only comes from the N₂ feed gas. A more recent study by the Qiao group shows that V_N confined on 2D W₂N₃ nanosheets provide an electron-deficient environment to facilitate N₂ adsorption and lower the thermodynamic limiting potential of the NRR.⁷¹ A nuclear magnetic resonance (NMR) test using ¹⁵N₂ as the feed gas verifies that the NH₃ was indeed generated *via* the NRR instead of decomposition of the catalyst or other contaminants.

The intrinsic high HER activity of MoS₂ limits its current efficiency for NH₃ formation.^{22,72} We have addressed this issue by designing S-rich defective MoS₂ nanoflowers.⁷³ This catalyst attains a much higher FE of 8.34% than its defect-free counterpart (2.18%). The DFT results suggest that the defects lead to the d-band center of Mo atoms moving close to the Fermi level (−0.26 eV → −0.14 eV) related to a stronger interaction between the catalyst surface and N₂ molecule (−0.65 eV). Moreover, the defective MoS₂ has a lower energy barrier of the PDS (0.60 eV) than the pristine one (0.65 eV).

For pure metal catalysts, the defect effect is rarely studied. A recent study by the Yan group demonstrates the modulation of the electronic properties of Bi *via* defect engineering (Fig. 4).⁷⁴ In their study, Bi(110) nanoplates with a high fraction of isolated Bi vacancies were fabricated from Bi₂O₃ nanoplates using a low-temperature plasma bombardment approach. This defect-Bi achieves an NH₃ yield of 5.453 μg h^{−1} mg_{cat.}^{−1} and a FE of 11.68%. Theoretical calculations indicate that defect-Bi(110) has a much lower activation energy (0.56 eV) in comparison with perfect-Bi(110) (1.03 eV) for the rate-determining step.

3.1.2 Heteroatom doping. Chemical doping with heteroatoms has also been proven to be an effective strategy to improve the electrocatalytic NRR activity. In Liu's study, N-doped porous carbon with a high N content and tunable N species was derived from zeolite imidazole framework-8 (ZIF-8) pyrolysis and tested for the NRR leading to an NH₃ yield of 23.8 μg h^{−1} mg_{cat.}^{−1} and a FE of 1.42%.⁷⁵ Although the authors established the correlation between the level of N doping and the NRR performance based on their experimental and theoretical data, a persuasive conclusion is still pending because of the unavailability of ¹⁵N₂ isotopic experiment results. Wang *et al.* directly utilized a hierarchically structured N-doped nanoporous carbon membrane (NCM) as the working electrode for electrochemical N₂-to-NH₃ conversion with a FE of 5.2% and an NH₃ yield of 0.08 g m^{−2} h^{−1}.⁷⁶ Decorating this NCM electrode with Au nanoparticles further improves the performance metrics to 22% and 0.36 g m^{−2} h^{−1}, respectively.

As a 2D nanocarbon with high conductivity and chemical stability, graphene with B doping attains an NH₃ yield of 9.8 μg h^{−1} cm^{−2} and a FE of 10.8%.⁷⁷ B incorporation causes redistribution of electron density and the as-formed electron-deficient B sites have higher N₂-binding capability. The much

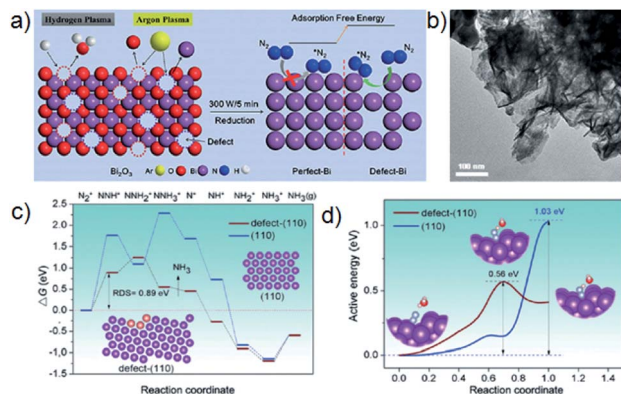


Fig. 4 (a) Illustration of the synthesis of a defect-Bi nanoplate and its application for the NRR; (b) TEM image of defect-Bi nanoplates; (c) free-energy diagrams for the NRR on the perfect-Bi(110) and defect-Bi(110) facets via a reaction pathway; (d) activation energy of the rate-determining step. Reproduced from ref. 74 with permission from Wiley-VCH, copyright 2019.

larger electronegativity of O with respect to C endows it with a stronger ability to manipulate the electronic properties of carbon catalysts. O-doped graphene derived from sodium gluconate was also identified by us as a superior NRR catalyst with a larger NH_3 yield of $21.3 \mu\text{g h}^{-1} \text{mg}_{\text{cat.}}^{-1}$ and a higher FE of 12.6%.⁷⁸ DFT data show that both C=O and O-C=O groups make a greater contribution to the NRR than the C-O group. Other heteroatoms like S⁷⁹ and F⁸⁰ also work effectively as dopants to boost the NRR performances of graphene.

Like carbon materials, metal-based catalysts can also be doped with non-metal and metal heteroatoms to boost the NRR performances. F was utilized by us to dope FeOOH to decrease the reaction energy barrier, and the performance metrics are greatly boosted from $10.01 \mu\text{g h}^{-1} \text{mg}_{\text{cat.}}^{-1}$ and 2.16% to $42.38 \mu\text{g h}^{-1} \text{mg}_{\text{cat.}}^{-1}$ and 9.02%, respectively.⁸¹ The same idea was followed by Liu *et al.* for SnO₂ mesoporous nanosheets,⁸² and it could also be applicable to other metal oxides. Cu doping in CeO₂ was reported to form multiple V_O for dramatically enhanced catalytic activities.⁸³ For such a strategy, TiO₂ is the most extensively studied metal oxide and several dopants like C,^{84,85} B,⁸⁶ V,⁸⁷ and Zr⁸⁸ work effectively. The C-doped TiO₂/C material derived from MIL-125(Ti) affords a high FE of 17.8%, which is attributed to the doping of C atoms into V_O and the formation of Ti-C bonds, which enable energetically more favourable N₂ activation.⁸⁵ In Zheng's work, given that Zr⁴⁺ has a similar d-electron configuration and oxide structure but relatively larger ionic size, it was doped into TiO₂ to induce the formation of adjacent bi-Ti³⁺ pairs as the most active centers for efficient N₂ lying-down chemisorption and activation ($8.90 \mu\text{g h}^{-1} \text{mg}_{\text{cat.}}^{-1}$; 17.3%) (Fig. 5).⁸⁸ Impressively, high-performance NRR catalysis can also be enabled over TiO₂ using Fe as a dopant due to the synergistic effect of bi-Ti³⁺ and V_O ($25.47 \mu\text{g h}^{-1} \text{mg}_{\text{cat.}}^{-1}$; 25.6%).⁸⁹

3.1.3 Surface functionalization and interface engineering.

To avoid using high-temperature thermal annealing to prepare carbon-based NRR catalysts, carbon nanotubes were acidically

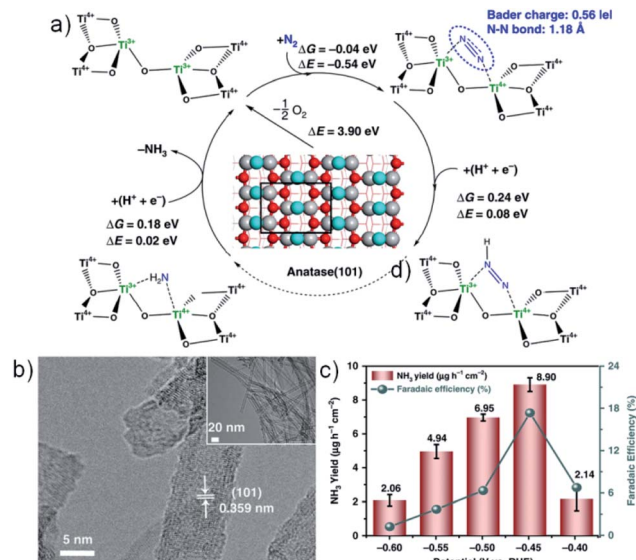


Fig. 5 (a) DFT prediction of the NRR activity for adjacent bi-Ti³⁺ on anatase (101) with V_O (light-blue spheres: lattice oxygen at the bridge sites where the surface V_O are formed most easily; red spheres: other lattice oxygens on the surfaces; gray spheres: titanium cations; ΔG : free energy; ΔE : electronic energy. (b) TEM image for Zr-TiO₂ nanotubes. (c) NH_3 yields and FEs of Zr-TiO₂ at each given potential. Reproduced from ref. 88 with permission from the Nature Publishing Group, copyright 2019.

oxidized to introduce functional groups (C-O, OH, COOH, C=O) on their surface, and as a NRR catalyst, they exhibit superior performances ($32.33 \mu\text{g h}^{-1} \text{mg}_{\text{cat.}}^{-1}$; 12.50%).⁹⁰ Our recent work further shows an enhancement in the NRR activity of reduced graphene oxide (rGO) after surface modification with oxygen-rich tannic acid (TA).⁹¹ The strong π - π stacking interactions between π -rich TA and rGO brings TA into very close proximity with rGO, which leads to intimate contact between the oxygen functional groups of TA and rGO favouring more effective manipulation of its electronic properties. The same strategy has also been used to improve the NRR performances of Pd catalysts ($24.12 \mu\text{g h}^{-1} \text{mg}_{\text{cat.}}^{-1}$; 9.49%).⁹² Compared to Pd, the much larger electronegativity of the O of TA could drive the formation of electron-deficient Pd with enhanced N₂-adsorption ability,⁶⁹ but the exact enhancement mechanism is not completely understood at the present time.

Because amorphous catalysts are in a metastable state with abundant unsaturated coordination sites, they exhibit higher catalytic activity than the crystalline ones. To this end, Shi *et al.* achieved the amorphization of Au nanoparticles anchored on rGO using CeO_x as a trigger, and the resulting a-Au/CeO_x-rGO hybrid shows superior NRR performances ($8.31 \mu\text{g h}^{-1} \text{mg}_{\text{cat.}}^{-1}$; 10.10%) to other control catalysts.⁹³ The Yu group also reported highly active NRR electrocatalysis enabled by a CeO₂/amorphous Bi₄V₂O₁₁ hybrid ($23.21 \mu\text{g h}^{-1} \text{mg}_{\text{cat.}}^{-1}$; 10.16%).⁹⁴ The amorphous phase has more abundant defective sites and lowers the energy barrier, and CeO₂ triggers its amorphization during calcination. More recently, metal/metal interface engineering has also been applied to boost NRR catalysis, and the donor-

acceptor couple of Ni and Au nanoparticles with a strong electronic connection attains a superhigh FE of 67.8%.⁹⁵ In this study, Au nanoparticles were directly deposited on Ni nanoparticles *via* Galvanic replacement, and the electron-rich Au nanoparticles accepting electrons from Ni nanoparticles facilitate the adsorption and activation of N₂ molecules.

3.2. Enriching the active sites of catalysts

Enriching the active sites to increase the apparent activity is regarded as the most straightforward way to boost the catalytic performances of electrocatalysts. To obtain a more active catalyst, one promising approach is nanostructuring of the catalyst to significantly increase the number of active sites. Previous studies point out that the size and shape of catalysts have a large influence on their NRR activities. The Yan group reported using Au sub-nanoclusters on TiO₂ as a more efficient catalyst (21.4 μg h⁻¹ cm⁻²; 8.11%) due to the fact that the small size of the sub-nanoclusters with high surface energy facilitates the formation of strong Au–O–Ti bonds and the resulting positively charged Au centers favour N₂ adsorption.⁹⁶ Au flowers were also reported to show higher performances than Au spheres, which is attributed to the exposure of more active sites by the dendritic structures of Au flowers.⁹⁷ Using hollow gold nanocages as a NRR catalyst, Nazemi *et al.* attained a FE of 30.2% with an NH₃ yield of 3.9 μg h⁻¹ cm⁻².⁹⁸ Such a unique hollow nanostructure not only exposes abundant active sites, but also provides a confined reaction environment that increases the residence time of N₂ molecules on its inner surface to enhance the conversion of N₂ to NH₃. In another study, we analyzed the performances of Cr₂O₃ catalysts, including hollow microspheres, solid microspheres, and nanoparticles with the following performance metrics: 25.3 μg h⁻¹ mg_{cat.}⁻¹ & 6.78%, 11.4 μg h⁻¹ mg_{cat.}⁻¹ & 2.94%, and 13.8 μg h⁻¹ mg_{cat.}⁻¹ & 4.73%.⁴⁶

Another attractive approach is to disperse electrocatalysts on supports with a high surface area and conductivity. Using such a support can better disperse and reduce aggregation of the nanocatalysts on one hand and enhance the charge transport on the other hand. Thus far, graphene has been largely used to support CoO,⁴⁸ CuO,⁴⁹ Au/CeO_x,⁹³ TiO₂,⁹⁹ Mn₃O₄,¹⁰⁰ Cr₂O₃,¹⁰¹ SnO₂,¹⁰² Fe₂O₃,¹⁰³ FeOOH,¹⁰⁴ PdCu,¹⁰⁵ PdP₂,¹⁰⁶ Ru₂P,¹⁰⁷ S dots,¹⁰⁸ and perylene-3,4,9,10-tetracarboxylic acid nanorods.¹⁰⁹ Graphene is electrocatalytically inert for the NRR, and its use as an inactive support would thus have an adverse effect on NH₃ yield per unit mass. The high conductivity and large surface area of 2D Ti₃C₂T_x MXene,¹¹⁰ together with its intrinsic NRR activity,^{43,111} are promising for its use as an ideal support for NRR catalysts. We grew whisker-like MnO₂ on Ti₃C₂T_x flakes and the MnO₂–Ti₃C₂T_x hybrid attains a large NH₃ yield of 34.12 μg h⁻¹ mg_{cat.}⁻¹ and a high FE of 11.39%, superior to those of each component due to the synergistic catalytic effect.¹¹² In another study, Ti₃C₂T_x MXene nanosheets act as both the support and Ti source for *in situ* solvothermal development of V_O-rich TiO₂ nanoparticles, and this hybrid efficiently catalyzes the NRR (32.17 μg h⁻¹ mg_{cat.}⁻¹; 16.07%).¹¹³ More recently, black P quantum dots were stably confined on electrochemically

active, electrically conductive black SnO_{2-x} nanotubes *via* Sn–P coordination, and both components in the hybrid synergistically catalyze the NRR to afford amazing performance metrics (48.87 μg h⁻¹ mg_{cat.}⁻¹; 14.6%).¹¹⁴

Single-atom catalysts (SACs) with maximum atom-utilization efficiency and unique properties enable reasonable use of metal resources and achieve atom economy, and they have recently emerged as a new Frontier in materials and catalysis sciences. Encouragingly, the establishment of reliable synthetic strategies to guarantee the stabilization of single metal atoms against migration and agglomeration has aroused huge recent interest in exploring their electrocatalytic applications.¹¹⁵ Several recent studies have described highly active N₂ reduction electrocatalysis enabled by SACs. Wang *et al.* synthesized atomically dispersed Au catalysts on carbon nitride for the NRR with a FE of 11.1% and an NH₃ yield of 1.305 μg h⁻¹ mg_{Au}⁻¹, which is ~22.5 times as high as that for supported Au nanoparticles.¹¹⁶ Au single sites stabilized on N-doped porous and highly oxidized carbon also enable efficient N₂ electroreduction (2.32 μg h⁻¹ cm⁻²; 12.3%).¹¹⁷ An NH₃ yield of 120.9 μg h⁻¹ mg_{cat.}⁻¹ was reported by Geng *et al.* over single Ru atoms anchored on N-doped carbon (Ru SAs/N–C).¹¹⁸ Han *et al.* prepared single Mo atoms anchored to N-doped porous carbon, and the high-density active sites and hierarchically porous carbon frameworks led to an NH₃ yield of 34.0 ± 3.6 μg h⁻¹ mg_{cat.}⁻¹ and a FE of 14.6 ± 1.6%.¹¹⁹ Another recent study by Wang *et al.* confirms that single-atom Fe on N-doped carbon achieves a much lower NH₃ yield of 7.48 μg h⁻¹ mg_{cat.}⁻¹ than Ru and Mo-based SACs.¹²⁰ These studies would open up an exciting new avenue to explore using SACs for electrocatalytic N₂ reduction, but care should be taken regarding these and other N-containing catalysts because they may decompose and leach N leading to inaccurate results.

3.3. Suppressing the HER

Suppressing the HER at the catalyst/electrolyte interface by manipulating the availability of H⁺ is an attractive route to improving NRR selectivity. DFT calculations predict that polyimide has intrinsic sluggish HER activity with a large activation barrier and Li⁺ association can further reduce the active sites to passivate the HER.¹²¹ Inspired by this, Chen *et al.* reported a Li⁺-incorporated poly(*N*-ethyl-benzene-1,2,4,5-tetracarboxylicdiimide) (PEBCD) to boost NRR selectivity, and the NRR and HER processes are effectively promoted and retarded, respectively, by associating Li⁺ ions with the O atoms in the PEBCD matrix.¹²² Although it achieves significant HER passivation, this catalyst attains a FE of only 2.85%, which could be due to its intrinsic low NRR activity. In another study by the Zhao group, *in-operando* created strong Li–S interactions endow S-rich MoS₂ with superior NRR activity (43.4 μg h⁻¹ mg_{cat.}⁻¹; 9.81%).¹²³ Such interactions suppress the HER, facilitate N₂ adsorption and increase NRR activity.

Hao *et al.* recently reported a strategy to simultaneously promote NRR selectivity and activity using Bi nanocrystals and K⁺ cations (Fig. 6).¹²⁴ The K⁺ cations not only stabilize the key nitrogen-reduction intermediates but also regulate proton transfer to increase the selectivity. A previous study suggests

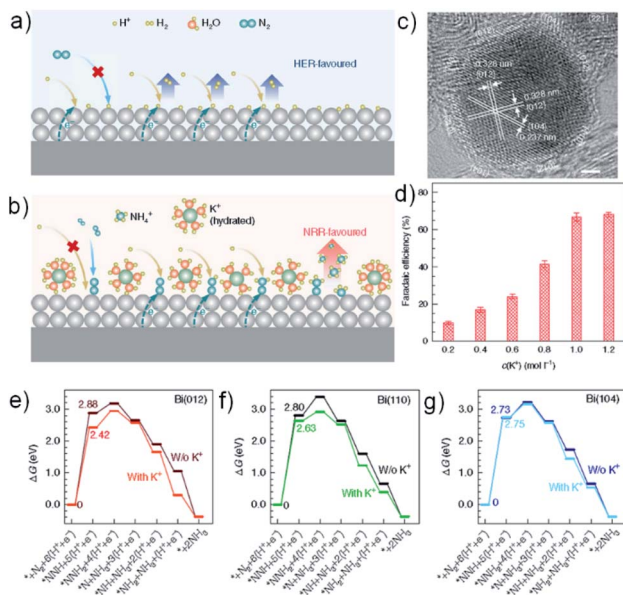


Fig. 6 Mass transfer of H⁺ and N₂ to the catalyst surface in electrolytes (a) without and (b) with K⁺. (a) In acidic solutions without K⁺, H⁺ can be transferred to the surface readily, and the HER will dominate. (b), K⁺ hinders H⁺ transfer to the catalyst surface. N₂ will be adsorbed preferentially, and the NRR is promoted. (c) High-resolution TEM image of BiNCs, showing the exposed surfaces of {012}, {104} and {110}. (d) FE vs. c(K⁺). Gibbs free-energy diagrams for the NRR on (e) Bi(012), (f) Bi(110), and (g) Bi(104) facets in the presence or absence of K⁺ (pH 0 and U = 0 V vs. SHE). Reproduced from ref. 124 with permission from the Nature Publishing Group, copyright 2019.

that supporting cations with high concentration can retard H⁺ migration from the bulk solution to the electrode surface leading to significant HER suppression.¹²⁵ In Hao's work, optimized NRR performances (200 mmol g⁻¹ h⁻¹; 66%) were obtained in acidic 0.5 M K₂SO₄ electrolytes (pH = 3.5). Of note, such NRR promotion and HER suppression by K⁺ can also apply to Au and Pt catalysts.

Coating a porous hydrophobic nanolayer over a NRR catalyst to repel water molecules could be another attractive alternative. This concept was first demonstrated in Ling's work.¹² The authors deposited Ag nanocubes onto an Au electrode, followed by coating with a bifunctional ZIF-71 thin film as a sorption layer for confining N₂ molecules to improve reactant-catalyst interactions and as a superhydrophobic barrier to inhibit water access. This structure affords a FE of 18 ± 4%, which is boosted by 10% compared to that of the uncoated catalyst. However, this system is neither cost-effective nor environmentally friendly because of the use of tetrahydrofuran and ethanol as the medium and proton source, respectively. An improved catalyst was recently designed by Yang *et al.*¹²⁶ In their work, nanoporous gold (NPG) as the core was coated with a ZIF-8 shell which weakens hydrogen evolution and retards reactant diffusion, leading to a superb aqueous-based NPG@ZIF-8 catalyst (28.7 ± 0.9 μg h⁻¹ cm⁻²; 44%). However, the authors have not commented on why the hydrophobic ZIF-8 can operate efficiently in an aqueous electrolyte.

The higher barrier for mass- and charge-transfer of neutral electrolytes leads to less favourable HER kinetics.¹²⁷ Wang *et al.* tested a Pd/C catalyst in PBS with a FE of 8.2%, but this catalyst only affords a FE lower than 0.1% in both H₂SO₄ and NaOH.¹⁶ It is argued that PBS is a promising electrolyte for the electrochemical NRR due to its effective suppression of HER activity. Given the wide variety of experimental parameters tested in each study, this conclusion should be taken with caution because the large amount of research experience of our group suggests that the choice of electrolyte for optimized NRR performances also relies strongly on the catalyst itself.

4. Conclusions and outlook

The electrocatalysis community has made some recent progress in the design and development of catalytic systems to enable efficient electrochemical N₂ reduction to NH₃ under ambient conditions. An ideal catalyst should effectively adsorb and activate N₂ molecules to overcome the slow kinetics. By manipulating the electronic structures of NRR catalysts, we can promote their intrinsic activities. Increasing the number of exposed active sites provides us with the most straightforward way to enhance the apparent catalytic activity. As the HER has lower overpotentials, the NRR in aqueous electrolytes always suffers from strong HER competition, leading to limited selectivity for NH₃ formation. Despite some successful examples demonstrating the effective suppression of the HER to boost NRR selectivity, the versatility of this strategy has not been fully exploited. New strategies for catalyst design and HER suppression are still needed. Obviously, the selectivity issue will be better addressed by more general HER-retarding strategies like adding organic additives in the electrolytes or modifying the catalyst surface with water solvable H⁺-repelling molecules. We should point out that it is not appropriate to only emphasize current efficiency, and NH₃ yield should also be treated seriously as an important metric for evaluating the overall performances of a NRR catalyst.

Recently, the reliability of the NRR results has provoked discussions of experimental protocols,¹²⁸ and strict standards have been set to obtain correct scientific reports.¹²⁹ Control measurements with Ar under the exact same conditions and with N₂ at open circuit potential are required to probe NH₃ contamination within the cell, catalyst, and feed gas.¹³⁰ Possible contaminants like NO_x and other labile nitrogen compounds can be effectively removed through adsorption on a reduced Cu catalyst,¹²⁹ and possible NH₃ and NO_x impurities can also be removed through a saturator filled with 0.05 M H₂SO₄ (aq).¹³⁰ Provided that all these protocols are carefully taken into consideration,^{129–131} both isotope-specific nuclear magnetic resonance and colorimetric measurements should give similar quantified results.¹²⁹ As a well-established technique for ion analysis, ion chromatography (IC) also provides quantitative results which are consistent with those of colorimetry.^{47,66,80} Both colorimetry and IC measurements should also be considered as reliable alternatives to quantitatively gauge the NH₃ product. The future development of electrochemical NH₃ synthesis will heavily rely on establishing more efficient

catalytic systems to achieve high selectivity and large current density for massive NH₃ production, which however will still remain a big challenge for quite a long time.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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