



Cite this: *Green Chem.*, 2024, **26**, 1302

Received 4th November 2023,  
Accepted 4th January 2024

DOI: 10.1039/d3gc04272c

rs.li/greenchem

## Beyond acceptable limits: intrinsic contamination in commercial $^{15}\text{N}_2$ impedes reliable $\text{N}_2$ reduction experiments†

Michiel De Ras,<sup>a</sup> Lander Hollevoet,<sup>b</sup> Johan. A. Martens,<sup>b</sup> Tianxi Liu,<sup>c</sup> Bart M. Nicolai,<sup>d</sup> Maarten L. A. T. M. Hertog,<sup>d</sup> Johan Hofkens<sup>a,e</sup> and Maarten B. J. Roeffaers<sup>id</sup>\*<sup>f</sup>

The validity of  $\text{N}_2$  reduction experiments relies on control experiments with isotopically labeled  $^{15}\text{N}_2$ . Here we discovered significant  $^{15}\text{NH}_3$  contamination (73 ppm) in commercial  $^{15}\text{N}_2$  and also observed the presence of oxidized  $^{15}\text{N}$  in amounts comparable to the typical reported ammonia yields.

### Introduction

The Haber–Bosch (HB) process, with its high carbon intensity (4.21 tCO<sub>2</sub>/tNH<sub>3</sub>), accounts for about 1.3% of the world's CO<sub>2</sub> emissions through the production of ammonia.<sup>1,2</sup> Over the past few years, various alternative synthesis routes for ammonia have been proposed. Amongst them is the direct nitrogen reduction reaction (NRR) in aqueous conditions (eqn (3)). This approach omits the use of a carbon-based hydrogen source and can operate at pressures and temperatures close to ambient. However, irrefutable experimental evidence for actual (photo)electrochemical ammonia formation at the cathode is typically elusive, this is mainly due to the extremely low reported ammonia production rates (10<sup>-12</sup>–10<sup>-9</sup> mol s<sup>-1</sup>). Moreover, the omnipresence of reactive nitrogen species, which are more easily converted to ammonia than the extremely stable dinitrogen, has led to concerns about false posi-

tives in many studies. In response, validation strategies as well as rigorous experimental protocols have been implemented. While most studies stress the use of high-purity reagents, blank tests, controls, *etc.*, the cornerstone of NRR validation lies in the use of isotopically labelled nitrogen gas ( $^{15}\text{N}_2$ ). Detection of  $^{15}\text{NH}_3$  post reaction is typically taken as an indicator of NRR occurrence. However, until now, there has been a lack of comprehensive information regarding the purity of commercial  $^{15}\text{N}_2$  gas, and the impact of activated,  $^{15}\text{N}$  labelled impurities on the measured ammonia yield. Furthermore, there is a notable absence of precise, quantitative comparison between NRR experiments conducted under  $^{14}\text{N}_2$  and  $^{15}\text{N}_2$ .

### False positives and contaminants in the $^{14}\text{N}_2$ feed gas

In a typical electrochemical NRR experiment, nitrogen gas is bubbled through the cathode compartments' aqueous electrolyte solution, where it serves as nitrogen source to generate ammonia. Depending on the pH, either gaseous NH<sub>3</sub> or dissolved NH<sub>4</sub><sup>+</sup> is formed (eqn (1.1) and (1.2)). The required protons originate from the oxidation of water, conducted in a separate anode compartment (eqn (2.1) and (2.2)). The cathode's production rate and selectivity are typically determined by quantifying the ammonia yield over the course of a few hours.

Cathode:

$$\text{pH} = 0$$



$$\text{pH} = 14$$



Anode:

$$\text{pH} = 0$$



<sup>a</sup>Molecular Imaging and Photonics, Department of Chemistry, KU Leuven, Celestijnenlaan 200F, Leuven, 3001, Belgium

<sup>b</sup>Centre of Surface Chemistry and Catalysis: Characterisation and Application team, KU Leuven, Celestijnenlaan 200F—box 2461, 3001 Leuven, Belgium

<sup>c</sup>The Key Laboratory of Synthetic and Biological Colloids, Ministry of Education, School of Chemical and Material Engineering, International Joint Research Laboratory for Nano Energy Composites, Jiangnan University, Wuxi, 214122, China

<sup>d</sup>MeBioS, Department of Biosystems, KU Leuven, Willem de Croylaan 42, 3001 Leuven, Belgium

<sup>e</sup>Max Plank Institute for Polymer Research, Mainz, D-55128, Germany

<sup>f</sup>Centre for Membrane Separations, Adsorption, Catalysis, and Spectroscopy for Sustainable Solutions, Department of Microbial and Molecular Systems, KU Leuven, Celestijnenlaan 200F, 3001 Leuven, Belgium

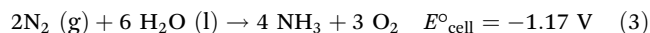
† Electronic supplementary information (ESI) available: Materials and methods, technical drawing of the custom setup and obtained SIFT-MS data. See DOI: <https://doi.org/10.1039/d3gc04272c>



pH = 14



Overall:



Considering that the reported ammonia production rates are typically small ( $10^{-12}$ – $10^{-9}$  mol  $\text{s}^{-1}$   $\text{cm}^{-2}$  of cathode), the challenge lies in verifying that the measured ammonia indeed originates from  $\text{N}_2$  reduction and not from more easily reducible, activated nitrogen species such as  $\text{NO}_3^-$ ,  $\text{NO}_2^-$ ,  $\text{NO}$ ,  $\text{NO}_2$ ,  $\text{NH}_2\text{OH}$ , *etc.* These latter species are often found as contaminants in the reaction system, for instance, being inherently present in the electrolytes, catalyst materials and/or the feed gasses.<sup>3,4</sup> While the concentration of these activated nitrogen species in the reaction system can be reduced, they cannot be entirely eliminated. Notably, even under optimal conditions, their amounts match typical ammonia yields.

A key distinction exists between oxidized nitrogen species like  $\text{NO}_3^-$  or  $\text{NO}_x$  and  $\text{NH}_3/\text{NH}_4^+$ . While background levels of  $\text{NH}_3(\text{g})$  and  $\text{NH}_4^+(\text{aq.})$  in the experimental setup can be factored in *via* blank tests and careful analysis of the reagents and electrodes, identifying contributions from oxidized nitrogen species poses a more significant challenge. Moreover, the ubiquitous and heavily fluctuating presence of ammonia in the atmosphere and on labware, complicate the accurate quantification of a background level.<sup>5–7</sup>

One contamination source of oxidized nitrogen species is the feed gas. Our mass spectrometry results (ESI Fig. 2a†) reveal that a commercial, high-purity (N6)  $^{14}\text{N}_2$  gas cylinder contains approximately 52 ppb of  $^{14}\text{NH}_3$ , 254 ppb of  $^{14}\text{NO}$ , and 22 ppb of  $^{14}\text{NO}_2$  (Table 1). The  $^{14}\text{NO}_x$  and  $^{14}\text{NH}_3$  contamination levels found in a lower purity (N3)  $^{14}\text{N}_2$  gas was found to be comparable and in the same order of magnitude as the 220 ppb of  $^{14}\text{N}_2\text{O}$  reported previously by Choi *et al.* in N5 purity  $^{14}\text{N}_2$ .<sup>5</sup> In these feed gasses, the background level of ammonia is approximately 5–6 times lower than the total amount of oxidized nitrogen species. For typical NRR experiments with  $^{14}\text{N}_2$  feed gas flowing through the cell at 100 ml  $\text{min}^{-1}$ , the total amount of oxidized nitrogen species introduced from high purity  $^{14}\text{N}_2$  gas (N6) can be expected in the order of  $10^{-11}$  mol  $\text{cm}^{-2}$   $\text{s}^{-1}$  (for a cathode of 1  $\text{cm}^2$ ). As this

molar flux of contaminants matches the ammonia production rate of many reported electrocatalysts, it is important to consider that reduction of contaminants in the feed gas alone could potentially account for the entire ammonia output, overshadowing the reduction of  $\text{N}_2$ .

The credibility of a given NRR study is primarily based on the ammonia production rate of the cathode. Consequently, thresholds to determine the plausibility of electrochemical NRR have been suggested by Choi *et al.*<sup>5</sup> Given typical contamination levels, electrodes exhibiting production rates lower than  $10^{-10}$  mol  $\text{cm}^{-2}$   $\text{s}^{-1}$  are not considered promising, while promising NRR experiments should present ammonia production rate exceeding  $10^{-8}$  mol  $\text{cm}^{-2}$   $\text{s}^{-1}$ . Taking this threshold and the above mentioned feed-gas contaminant quantification into account; at modest flow rates, the risk of measuring false positives due to contaminated feed gas is rather limited if high purity (N6)  $^{14}\text{N}_2$  gas is used. Obviously, this needs to be assessed for each individual case and researchers should always establish background contamination levels prior to performing NRR experiments.

## Contamination levels of commercial $^{15}\text{N}_2$ gas

While the contamination level in high-purity (N6)  $^{14}\text{N}_2$  gas is relatively minor compared to the literature-reported ammonia production thresholds, contamination in the form of  $\text{NH}_3/\text{NH}_4^+$  and oxidized nitrogen ( $^{14}\text{N}$ ) species such as  $\text{NO}_3^-$ ,  $\text{NO}_2^-$ ,  $\text{NH}_2\text{OH}$ , *etc.*, present in the used electrolytes, chemicals, and labware is significantly higher – orders of magnitude above the threshold for plausible NRR. As an example, Li *et al.* measured a  $\text{NO}_3^-$  concentration of 11 ppm in an aqueous 0.5 M  $\text{Li}_2\text{SO}_4$  electrolyte solution.<sup>7</sup> Distinctly separating the reduction of these contaminants to ammonia from the NRR proves even more challenging than eliminating the contribution of background  $\text{NH}_3/\text{NH}_4^+$  in the system. As a consequence, adopting isotopically labelled  $\text{N}_2$  gas ( $^{15}\text{N}_2$ ) and subsequent quantification of the produced  $^{15}\text{NH}_3$  has been proposed as a valid strategy to confirm ammonia production truly originates from  $\text{N}_2$ .<sup>5,8</sup> By switching to  $^{15}\text{N}_2$  feed gas, the interference of previously mentioned contaminants decreases sub-

**Table 1** Absolute concentrations of  $\text{NH}_3$ ,  $\text{NO}$  and  $\text{NO}_2$  in commercially obtained high purity (N6) and low purity (N3)  $^{14}\text{N}_2$  gas as well as  $^{15}\text{N}_2$  ( $\geq 98$  at%), determined by SIFT-MS

	Background $\text{NH}_3$ $^{14}\text{NH}_3$ (ppb)	Oxidized nitrogen species		
		$^{14}\text{NO}$ (ppb)	$^{14}\text{NO}_2$ (ppb)	$^{14}\text{N}_2\text{O}$ (ppb)
$^{14}\text{N}_2$ (N6 purity)	52	254	22	220 <sup>a</sup>
$^{14}\text{N}_2$ (N3 purity)	76	220	34	NA
	$^{15}\text{NH}_3$ (ppb)	$^{15}\text{NO}$ (ppb)	$^{15}\text{NO}_2$ (ppb)	$^{15}\text{N}_2\text{O}$ (ppb)
$^{15}\text{N}_2$ (98 at%)	73 293	659	20	230 <sup>a</sup>

<sup>a</sup>  $\text{N}_2\text{O}$  concentration reported by Choi *et al.* for commercially obtained  $^{14}\text{N}_2$  (N5) and  $^{15}\text{N}_2$  (N3).



stantially, as the natural abundance of the  $^{15}\text{N}$  isotope is only 0.33–0.42%.<sup>9</sup>

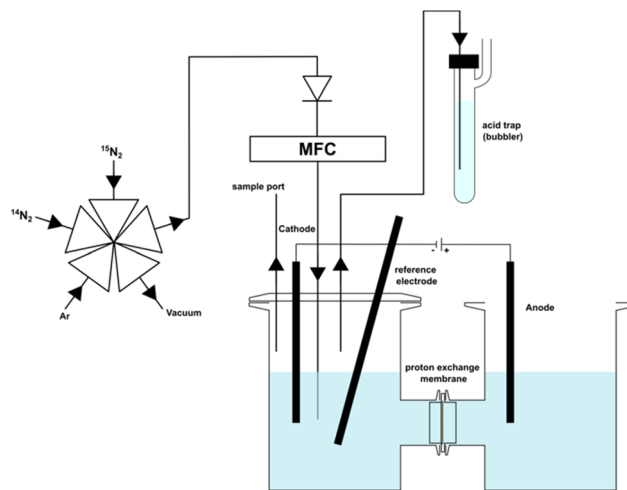
Previous research by Choi and Dabundo pointed out the existence of contaminants such as  $^{15}\text{NO}_x$  in commercial  $^{15}\text{N}_2$  gas. However, these studies were limited in their capability to provide precise quantifications of  $^{15}\text{N}$  labelled contaminants.<sup>10</sup> Dabundo's approach relied on indirect means of measurement, identifying the presence but not the exact amounts of reactive  $^{15}\text{N}$  species in the gas.<sup>10</sup> In contrast, our study employs mass spectrometry to explicitly quantify both oxidized  $^{15}\text{N}$  species ( $^{15}\text{NO}$  and  $^{15}\text{NO}_2$ ) and  $^{15}\text{NH}_3$  in the gas phase. Alarming, our findings show that commercial  $^{15}\text{N}_2$  is considerably contaminated with  $^{15}\text{NH}_3$  (Table 1, ESI Fig. 3†) (approx. 73 ppm), consistent with dissolved  $^{15}\text{NH}_4^+$  concentrations reported by Dabundo (34–1900 ppm). Herein lies a profound risk of measuring false positives: using as-obtained  $^{15}\text{N}_2$  gas bubbled at a  $100\text{ ml min}^{-1}$  flow rate either during the pre-experimental electrolyte saturation step or consistently throughout the entire experiment introduces sufficient  $^{15}\text{NH}_3$  to account for an apparent ammonia yield of approximately  $1 \times 10^{-8}\text{ mol s}^{-1}$ . This exceeds most reported ammonia production rates by factors of 100–1000. It is important to note that despite the evident risks, many recent NRR studies continue to use unpurified  $^{15}\text{N}_2$ , even though pre-experimental gas purification has been suggested.<sup>5,8,11–13</sup>

## NRR experiments using $^{15}\text{N}_2$ and $^{14}\text{N}_2$

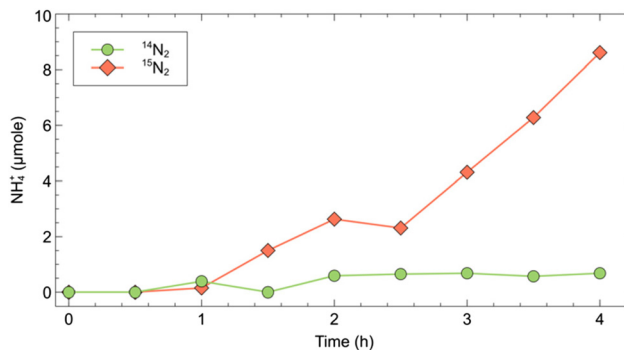
To effectively demonstrate the impact of  $^{15}\text{N}$  labelled impurities on the reported ammonia yield, we conducted NRR experiments at a constant potential ( $-0.91\text{ V vs. RHE}$ ), using bismuth, reportedly a highly active element for  $\text{N}_2$  reduction, as cathode.<sup>14</sup> We performed two sets of experiments under  $^{14}\text{N}_2$  and  $^{15}\text{N}_2$  saturated conditions (Fig. 1). Regularly sampling the catholyte revealed a measurable increase in ammonium content in the  $^{15}\text{N}_2$  experiment. Specifically, after four hours, the  $^{15}\text{N}_2$  experiment showed  $8.6\text{ }\mu\text{mol}$  of  $\text{NH}_4^+$ , compared to only  $0.68\text{ }\mu\text{mol}$   $\text{NH}_4^+$  in the  $^{14}\text{N}_2$  experiment (Fig. 2). This result clearly demonstrates that when using unpurified  $^{15}\text{N}_2$  even at low flow rates ( $20\text{ mL min}^{-1}$ ), a considerable amount of ammonia will be detected, primarily originating from the  $^{15}\text{N}$  labelled impurities rather than from the direct reduction of  $^{15}\text{N}_2$ . If the  $^{15}\text{N}_2$  contamination were overlooked, an apparent ammonia yield of  $6 \times 10^{-10}\text{ mol s}^{-1}$  would be calculated for the  $^{15}\text{N}_2$  experiment. This figure is over ten times higher than the ammonia yield ( $5 \times 10^{-11}\text{ mol s}^{-1}$ ) of the same experiment under  $^{14}\text{N}_2$ .

## Mitigation strategies for the $^{15}\text{N}$ contaminants

Our precise quantification reveals that the level of oxidized  $^{15}\text{N}$  species in this gas is nearly three times higher than in both high (N6) and low purity (N3)  $^{14}\text{N}_2$  gases, exacerbating the risk



**Fig. 1** Schematic representation of the setup used for studying the NRR under  $^{14}\text{N}_2$  or  $^{15}\text{N}_2$  atmospheres. NRR experiments were conducted in an H-cell configuration. Feed gases ( $^{14}\text{N}_2$  and  $^{15}\text{N}_2$ ) were continuously bubbled into the cathode compartment at a predefined flow rate of  $20\text{ mL min}^{-1}$ . An acid trap was employed at the exit to completely isolate the cathode compartment from the surrounding atmosphere and to trap any gaseous ammonia.



**Fig. 2** Cumulative ammonium production ( $^{15}\text{NH}_4^+$  or  $^{14}\text{NH}_4^+$ ) in a four hour NRR experiment with a Bismuth cathode.

of false positives in control experiments. Purifying commercial  $^{15}\text{N}_2$  gas, which comes in extremely limited volumes of 500–1000 cc at 1 to 5 atm, presents significant technical and economic challenges for NRR research. Traditional methods for purification, such as bulky, home-made scrubbers or acid traps, are neither practical nor effective in removing all  $^{15}\text{N}$  contaminants, and even more advanced solutions like small, commercial gas filters require complex, leak-tight setups for recirculating the limited  $^{15}\text{N}_2$  supply. Moreover, to ensure data reliability, continuous monitoring of activated  $^{15}\text{N}_x$  species using sensitive techniques like mass spectrometry is necessary, as is the tracking of  $^{15}\text{NH}_4^+$  levels in solution over time, rather than relying on a single post-reaction measurement, as is often performed nowadays.

Given these constraints, the risk of false positives in isotopic control experiments for NRR studies is significantly larger



when using  $^{15}\text{N}_2$  gas than  $^{14}\text{N}_2$ , regardless of purity level. This casts serious doubt on the reliability and cost-effectiveness of using  $^{15}\text{N}_2$  gas as it currently stands and highlights the urgent need for alternative validation methods in NRR research that can reliably account for these challenges.

## Conclusion

In recent years, the predominant strategy to exclude false positives in NRR experiments has centered on using isotopically labelled  $^{15}\text{N}_2$  gas as a feed. While Choi *et al.* warned researchers about potential interferences from isotopically labelled reactive nitrogen species in these gasses, a comprehensive understanding of the precise contamination levels of these gaseous,  $^{15}\text{N}$  labelled contaminants remained elusive. In this work, we quantify the exact levels of  $^{15}\text{N}_x$  contaminants in commercially obtained  $^{15}\text{N}_2$ , typically used for performing NRR control experiments. Alarming, the contamination levels in these gasses make them unsuitable, increasing the chances of obtaining false positives. Thus, given the current gas purity, validation experiments with  $^{15}\text{N}_2$  do not enhance the credibility of the NRR study when the production rate is low (*e.g.*  $<10^{-8}$  mol  $\text{cm}^{-2}$   $\text{s}^{-1}$ ), neither do they confirm NRR is occurring. Moreover, the  $^{15}\text{N}$  labelling experiments might even mislead researchers into wrongly attributing the measured  $^{15}\text{NH}_3$  to  $^{15}\text{N}_2$  reduction at the cathode; as we clearly demonstrated in our comparative NRR experiments. Indeed, the ammonia production rates originating from  $^{15}\text{N}$  labelled impurities in the feed gas is in the order  $10^{-9}$ – $10^{-10}$  mol  $\text{s}^{-1}$ , which is comparable to most reported NRR studies in aqueous conditions. Therefore, the most convincing proof for successful NRR is consistently measuring high ammonia yields (*e.g.*  $>10^{-8}$  mol  $\text{s}^{-1}$ ), particularly when using purer  $^{14}\text{N}_2$ , which is more realistic to purify, over the scarce  $^{15}\text{N}_2$ .

## Author contributions

M. D. R wrote the paper and assisted in performing the SIFT-MS measurements. M. B. J. R. participated in the writing and editing of the manuscript. L. H., J. A. M. and J. H. participated in the revision of the manuscript. T. L., B. M. N. and M. L. A. T. M. H contributed to scientific discussions and setting up the methodology.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

We thank Tessa Vanempen for her assistance with the SIFT-MS measurements and subsequent quantification. M. D. R. acknowledges the Research Foundation Flanders

(FWO) for his personal grant (1SA3321N). J. H. and M. B. J. R gratefully acknowledge the financial support from the Research Foundation-Flanders (FWO, Grant No. G0983.19N, G0F2322N, VS06523N), Interne Fondsen KU Leuven through project (C3/20/067 and CELSA/21/016). J. H. acknowledges the Flemish Government Through Long-Term Structural Funding Methusalem (CASAS2, Meth/15/04) and the Moonshot csBO project P2C (HBC.2019.0108), and the MPI as MPI fellow.

## References

- 1 R. Nayak-Luke, R. Bañares-Alcántara and I. Wilkinson, *Ind. Eng. Chem. Res.*, 2018, **57**, 14607–14616.
- 2 P. Gilbert and P. Thornley, *Energy and carbon balance of ammonia production from biomass gasification*, ed. A. V. Bridgwater, Proceedings of the Bioten Conference on Biomass, Bioenergy and Biofuels, Host publication, 2010.
- 3 J. Kibsgaard, J. K. Nørskov and I. Chorkendorff, *ACS Energy Lett.*, 2019, **4**, 2986–2988.
- 4 L. F. Greenlee, J. N. Renner and S. L. Foster, *ACS Catal.*, 2018, **8**, 7820–7827.
- 5 J. Choi, B. H. R. Suryanto, D. Wang, H.-L. Du, R. Y. Hodgetts, Federico M. Ferrero Vallana, D. R. MacFarlane and A. N. Simonov, *Nat. Commun.*, 2020, **11**, 5546.
- 6 B. Hu, M. Hu, L. Seefeldt and T. L. Liu, *ACS Energy Lett.*, 2019, **4**, 1053–1054.
- 7 L. Li, C. Tang, D. Yao, Y. Zheng and S.-Z. Qiao, *ACS Energy Lett.*, 2019, **4**, 2111–2116.
- 8 S. Z. Andersen, V. Čolić, S. Yang, J. A. Schwalbe, A. C. Nielander, J. M. McEnaney, K. Enemark-Rasmussen, J. G. Baker, A. R. Singh, B. A. Rohr, M. J. Statt, S. J. Blair, S. Mezzavilla, J. Kibsgaard, P. C. K. Vesborg, M. Cargnello, S. F. Bent, T. F. Jaramillo, I. E. L. Stephens, J. K. Nørskov and I. Chorkendorff, *Nature*, 2019, **570**, 504–508.
- 9 J. R. de Laeter, J. K. Böhlke, P. De Bièvre, H. Hidaka, H. S. Peiser, K. J. R. Rosman and P. D. P. Taylor, *Pure Appl. Chem.*, 2009, **81**, 1535–1536.
- 10 R. Dabundo, M. F. Lehmann, L. Treibergs, C. R. Tobias, M. A. Altabet, P. H. Moisaner and J. Granger, *PLoS One*, 2014, **9**, e110335.
- 11 Y. Ashida, Y. Onozuka, K. Arashiba, A. Konomi, H. Tanaka, S. Kuriyama, Y. Yamazaki, K. Yoshizawa and Y. Nishibayashi, *Nat. Commun.*, 2022, **13**, 7263.
- 12 P. L. Arnold, T. Ochiai, F. Y. T. Lam, R. P. Kelly, M. L. Seymour and L. Maron, *Nat. Chem.*, 2020, **12**, 654–659.
- 13 Q. Wang, J. Pan, J. Guo, H. A. Hansen, H. Xie, L. Jiang, L. Hua, H. Li, Y. Guan, P. Wang, W. Gao, L. Liu, H. Cao, Z. Xiong, T. Vegge and P. Chen, *Nat. Catal.*, 2021, **4**, 959–967.
- 14 Y.-C. Hao, Y. Guo, L.-W. Chen, M. Shu, X.-Y. Wang, T.-A. Bu, W.-Y. Gao, N. Zhang, X. Su, X. Feng, J.-W. Zhou, B. Wang, C.-W. Hu, A.-X. Yin, R. Si, Y.-W. Zhang and C.-H. Yan, *Nat. Catal.*, 2019, **2**, 448–456.

