



Cite this: *Phys. Chem. Chem. Phys.*,
2018, 20, 18729

On the possibility of an Eley–Rideal mechanism for ammonia synthesis on $\text{Mn}_6\text{N}_{5+x}$ ($x = 1$)-(111) surfaces†

Constantinos D. Zeinalipour-Yazdi ^{ab}

Recently we reported an Eley–Rideal/Mars–van Krevelen mechanism for ammonia synthesis on cobalt molybdenum nitride ($\text{Co}_3\text{Mo}_3\text{N}$). In this mechanism hydrogenation of activated dinitrogen occurs directly from the gas phase in a low barrier step forming a hydrazinylidene intermediate $=\text{NNH}_2$. In this paper we study whether such a mechanism of ammonia synthesis could occur on the (111) surface of another metal nitride, $\text{Mn}_6\text{N}_{5+x}$ ($x = 1$), as this would explain the low-T ammonia synthesis activity of $\text{Co}_3\text{Mo}_3\text{N}$. We find that although N_2 adsorbs more strongly than H_2 on the (111) surface, having also examined the (110) and the (100) surface, N_2 is not significantly activated when adsorbed in an end-on configuration. The hydrogenation reactions via an Eley–Rideal mechanism are all high barrier processes ($>182 \text{ kJ mol}^{-1}$) and therefore an Eley–Rideal mechanism for ammonia synthesis is predicted to not occur on this material unless there are high temperatures. Our study indicates that the fact that an Eley–Rideal/Mars–van Krevelen mechanism occurs on $\text{Co}_3\text{Mo}_3\text{N}$ is a result of the stronger activation of dinitrogen at nitrogen vacancies when dinitrogen is adsorbed in an end-on configuration.

Received 13th April 2018,
Accepted 21st June 2018

DOI: 10.1039/c8cp02381f

rsc.li/pccp

1. Introduction

Metal nitrides, which are also called interstitial nitrides, have varying amounts of lattice nitrogen that occupies octahedral coordination sites, with the general formula MN_y , with $M = 3d$ metal and $y < 1$.¹ The structure that has received most attention is $\eta\text{-Mn}_3\text{N}_2$,² which was found to be anti-ferromagnetic.³ $\varepsilon\text{-Mn}_4\text{N}^4$ has the lowest percent nitrogen content, in which the atoms occupy a face-centered cubic (fcc) arrangement with interstitial nitrogen. The following in nitrogen content is $\zeta\text{-Mn}_2\text{N}_{0.86}$ ⁵ (occasionally labelled as Mn_2N^6 and Mn_3N_2 ⁷) which has a hexagonal close packed (hcp) structure. The manganese nitride with the largest nitrogen content is $\theta\text{-MnN}^8$ that has a slightly tetragonally distorted rocksalt phase (occasionally labelled as $\theta\text{-Mn}_6\text{N}_{5+x}$ ²). The manganese nitride phase studied here is $\theta\text{-Mn}_6\text{N}_{5+x}$ (for $x = 1$) which however did not contain any intrinsic nitrogen vacancies in the $3 \times 1 \times 1$ unit cell and had therefore a 3 times smaller unit cell as shown in Fig. 1. This manganese nitride is referred to in this paper as $\theta\text{-Mn}_6\text{N}_5$. This manganese nitride has the advantage of having well defined low Miller index surfaces, which are depicted in Fig. 2.

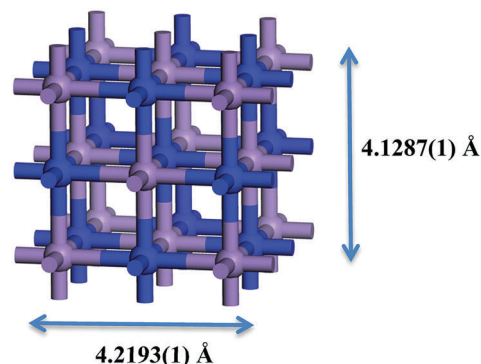


Fig. 1 Bulk unit cell of $\theta\text{-Mn}_6\text{N}_{5+x}$ ($x = 1$). Blue atoms correspond to nitrogen atoms and light blue to manganese atoms. Structure taken from ref. 9.

Manganese nitrides have also been studied as a looping catalyst in ammonia synthesis^{10,11} and for the reverse reaction, ammonia decomposition, by thermogravimetric analysis and neutron diffraction.¹² $\text{Mn}_6\text{N}_{5+x}$ and lithium imide (Li_2NH) when combined result in a larger rate of ammonia decomposition than for either of the separate parts.¹³ Additionally, CaNH can exert a strong synergistic effect on $\text{Mn}_6\text{N}_{5+x}$ leading to greatly enhanced catalytic activity for ammonia decomposition.¹⁴ Manganese nitrides were also found to be an active and durable electrocatalyst in the oxygen evolution reaction (OER) from water under alkaline conditions.¹⁵

^a Department of Chemistry, University College London, 20 Gordon Street, London, WC1H 0AJ, UK. E-mail: c.zeinalipour-yazdi@ucl.ac.uk

^b UK Catalysis Hub, Research Complex at Harwell, Rutherford Appleton Labs, Harwell Campus, OX11 0FA, UK

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8cp02381f



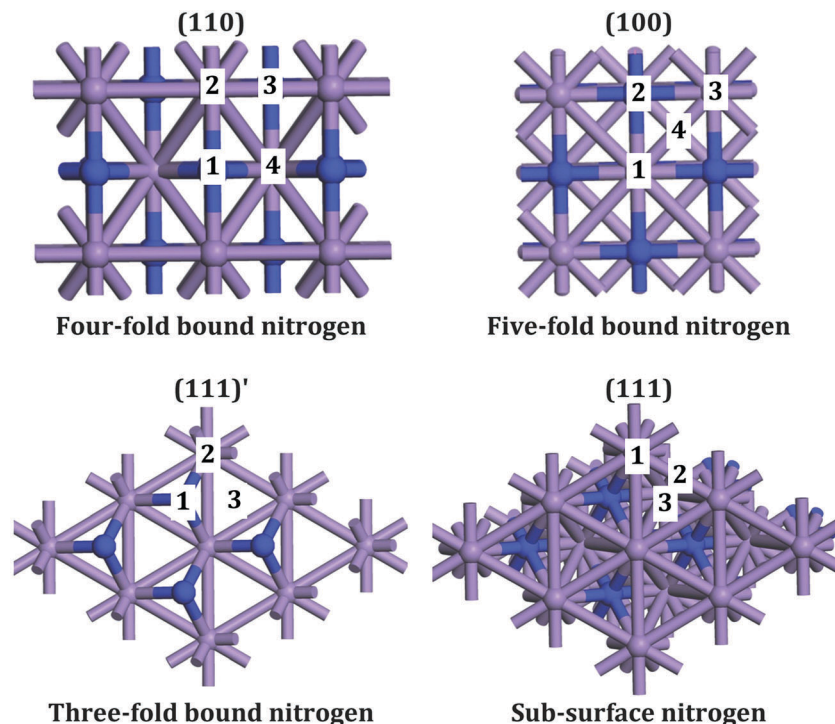


Fig. 2 Low Miller index surfaces of θ - Mn_6N_5 . (111)' indicates that the surface is parallel to the (111) surface. Blue and light-blue atoms are nitrogens and manganese, respectively.

Manganese nitrides can be synthesized by nitriding the metal with nitrogen or ammonia.¹⁶ The rate of manganese nitride formation is higher at lower temperatures when ammonia instead of nitrogen is used as the source of lattice nitrogen,¹⁷ which is a result of ammonia having a lower dissociation barrier than dinitrogen. Most of the manganese nitrides have been studied for their magnetic properties (*e.g.* ε - Mn_4N is ferromagnetic) however little is known about their activity as ammonia synthesis catalysts or nitrogen transfer reagents. There are only a few previous reports that mention a manganese nitride with the chemical formula θ - Mn_6N_5 . There is an electron diffraction study of the phase transition of this material from tetragonal to cubic at temperatures that exceed 440 ± 20 °C. The phase transition is attributed to magnetic reordering of the material rather than changes of the position of the atoms.¹⁸ Another report which addresses the magnetic transition in more detail is a neutron diffraction study of both η - Mn_3N_2 and θ - Mn_6N_5 .⁹

The potential use of metal nitrides in ammonia synthesis as catalysts or as reagents has attracted considerable recent interest.¹⁹ Directly relevant to this is the use of Mg_3N_2 as a nitrogen transfer reagent in the Paal-Knorr pyrrole reaction.²⁰ We have recently reported this material to be an efficient nitrogen transfer reagent especially when doped with lithium.¹¹ The discovery of catalysts or reagents that can synthesize ammonia at lower temperature for small-scale production is highly desirable for *in situ* and “on demand” production of ammonia.¹¹ However to use a catalyst in a decentralised plant, exact knowledge of the reaction mechanism and kinetic models is necessary. Currently the mechanism of ammonia synthesis is not known on manganese nitrides, whether

it follows a conventional Langmuir–Hinshelwood mechanism such as the one observed on the iron or ruthenium²¹ ammonia synthesis catalysts or an Eley–Rideal/Mars–van Krevelen mechanism such as the one we recently found on cobalt molybdenum nitride ($\text{Co}_3\text{Mo}_3\text{N}$).^{22,23} Furthermore, the details of the adsorption and activation of molecular nitrogen and hydrogen are not known and whether hydrogen poisons the surface of the catalyst as the adsorption energy of H_2 usually exceeds that of N_2 , as we have recently calculated on other metal nitrides.^{24,25}

In this DFT study we have calculated the adsorption and activation of N_2 and H_2 on various low Miller index surfaces of θ - Mn_6N_5 . We found the surface of this material that will adsorb N_2 favourably instead of H_2 . We then calculated the complete pathway for ammonia synthesis *via* an Eley–Rideal mechanism. The results are then discussed in reference to another metal nitride, $\text{Co}_3\text{Mo}_3\text{N}$, that has previously exhibited the possibility of an Eley–Rideal/Mars–van Krevelen mechanism for ammonia synthesis.

2. Computational methods

We have generated a slab model of the various surfaces of θ - Mn_6N_5 based on the structure reported in a neutron diffraction study.⁹ The surfaces were modelled *via* slabs about 1 nm in width that had a 20 Å vacuum gap, for which the surface structure is shown in Fig. 2. All DFT calculations were periodic Γ -point²⁶ spin-polarised performed with the VASP 5.4.1 code.^{27,28} Exchange and correlation (XC) effects were evaluated



within the generalized gradient approximation (GGA) using the revised Perdew–Burke–Ernzerhof (revPBE) XC-functional.²⁹ We used the projector augmented-wave (PAW) method^{30,31} to represent core states, 1s for N and H and 1s to 3p for Mn. The cut-off energy for the plane wave expansion was set to 600 eV for periodic bulk and slab calculations. Geometry optimizations were performed with a residual force threshold of 0.01 eV Å⁻¹ using the conjugate-gradient algorithm whereas the convergence criterion for electronic relaxation was set to 10⁻⁴ eV.

In a previous computational study we have tested the bond dissociation enthalpy (BDE) of N₂ with the use of various XC functionals, the % error for revPBE was found to be the lowest among various GGA and hybrid GGA functionals. In particular, we found that the % error was for revPBE = 0.2 < B3LYP = 1.7 < PBE0 = 3.5 < HSE06 = 4.1 < PBE = 4.9 < PW91 = 7.0.²³ Therefore, we have combined this XC functional with pairwise additive correction for van der Waals (VdW) interactions *via* the D3 method,³² denoted as revPBE-D3, with which all subsequent calculations have been performed to yield more accurate barrier heights according to the calculated BDE of the N–N bond and the inclusion of VdW interactions.

For the bulk unit cell, convergence of the energetic and magnetic properties was observed for a 5 × 5 × 5 Monkhorst–Pack grid (MP-grid). The convergence plots are given as supporting information Fig. S1 and S2 (ESI†). A similar or slightly larger grid density (so that the grid obtains numbers to the nearest integer) was used for the slab calculations, for consistency. A Γ -point centred MP *k*-point grid was used with a grid density of 3 × 3 × 1, 2 × 2 × 1, 2 × 2 × 1 and 3 × 2 × 1 for the (100), (111), (111)' and (110) surface of θ -Mn₆N₅, respectively, shown in Fig. 2.

The initial charge density was obtained by superposition of atomic charges. Initial adsorption configurations were such that the distance between the adsorbate and the nearest surface site was set to 2 Å in an end-on configuration for N₂ and H₂. This adsorption configuration was found to also result in end-on and tilt adsorbed adsorbates. The various adsorption sites were every symmetry unique surface site shown in Fig. 2. This resulted in 4, 4, 3 and 3 symmetry unique adsorption sites for the (110), (100), (111)' and (111) surface, respectively. The adsorption energy was taken as the total difference between the energy of the fully relaxed bound state of the surface-adsorbate complex and that of the fully relaxed surface slab and the isolated molecules, given by:

$$\Delta E_{\text{ads,D3}} = E_{\text{slab-X2}} - E_{\text{slab}} - E_{\text{X2}}, \quad (1)$$

where X = N, H. Dispersion corrections were included *via* the DFT-D3 method as implemented in VASP.³² Activation barriers were calculated with the nudged elastic band (NEB) method having five images for each barrier as implemented in VASP.³³

The effective magnetic moment per Mn atom in the optimised bulk structure of θ -Mn₆N₅ was found to be 3.95 μ_{B} , which was in good agreement with the experimentally measured value which ranges between 3.3 and 3.8 μ_{B} .⁹ Furthermore, the optimised structure of the bulk unit cell was found to converge to the lattice parameters determined by neutron diffraction ($a = b = 4.2193(1)$ Å, $c = 4.1287(1)$ Å, and $\alpha = \beta = \gamma = 90^\circ$),⁹ when the initial structure was the neutron diffraction crystal structure, for cutoff energies of the

plane wave expansion greater than 500 eV. When the plane wave expansion had a cutoff smaller than 500 eV, these lattice parameters were severely underestimated.

We have studied an Eley–Rideal mechanism for ammonia synthesis on θ -Mn₆N₅ but without the missing N in the 3 × 1 × 1 unit cell. This reduces the size of the unit cell considerably to the 1 × 1 × 1 unit cell shown in Fig. 1, which is a face-centred cubic arrangement of Mn atoms with N atoms octahedrally coordinated. This unit cell has a chemical formula Mn₄N₄ (or MnN) in which lower Miller index surfaces are shown in Fig. 2. Although our calculations for simplicity were done on Mn₄N₄, we refer to it as θ -Mn₆N₅.

3. Results and discussion

3.1. Adsorption and activation of N₂ and H₂

The adsorption of N₂ was investigated on every atop, bridge and hollow adsorption site of the (111), (111)', (110) and (100) surfaces of θ -Mn₆N₅ as depicted in Fig. 2. The dispersion-corrected adsorption energy and the percent activation of the N–N bond are shown in Fig. 3a. The adsorption energy was found to be between –5 kJ mol⁻¹ and –43 kJ mol⁻¹, which is weak, indicating physisorption except for the (111) surface where a value of –120 kJ mol⁻¹ was calculated. This adsorption energy is unusually large for dinitrogen and it clearly suggests chemisorption. Although this adsorption energy is very large, the percent activation of the N–N bond is less than 4%. This is much lower than the activation of end-on adsorbed N₂ on Co₃Mo₃N at nitrogen vacancies where 11% activation was found. However, it is comparable to the activation of N₂ found on another metal nitride, Ta₃N₅, which was less than 3%.²⁴ Therefore, the adsorption and activation of N₂ on this metal nitride is molecular in the absence of nitrogen vacancies. It is intriguing that among the various metal nitrides studied computationally thus far, the (111) surface of θ -Mn₆N₅ is the only surface where the adsorption energy of N₂ exceeds that of H₂; we will return to this point later.

We have studied the adsorption of H₂ on all low Miller index surfaces of θ -Mn₆N₅. The adsorption of H₂ was found to be molecular with an adsorption energy that ranged between –79 kJ mol⁻¹ and –87 kJ mol⁻¹. This adsorption energy is rather large for molecular hydrogen and identical in strength with the value calculated on other metal nitrides such as Ta₃N₅ where it was found to be between –81 kJ mol⁻¹ and –87 kJ mol⁻¹. In contrast to what was observed on Co₃Mo₃N surfaces, we do not observe dissociative adsorption of hydrogen when nitrogen vacancies are not present and the adsorption of H₂ is molecular on every symmetry unique adsorption site of the (111), (111)', (110) and (100) surfaces of θ -Mn₆N₅. This indicates that there is a barrier for the dissociative chemisorption of H₂ on θ -Mn₆N₅, however the dissociative chemisorption of H₂ at N-vacancies was not considered. In total 15 adsorption sites were investigated and these corresponded to every atop, bridge and hollow site on the three low Miller index surfaces of θ -Mn₆N₅. The adsorption energy of H₂ and the percent activation of the H–H bond are



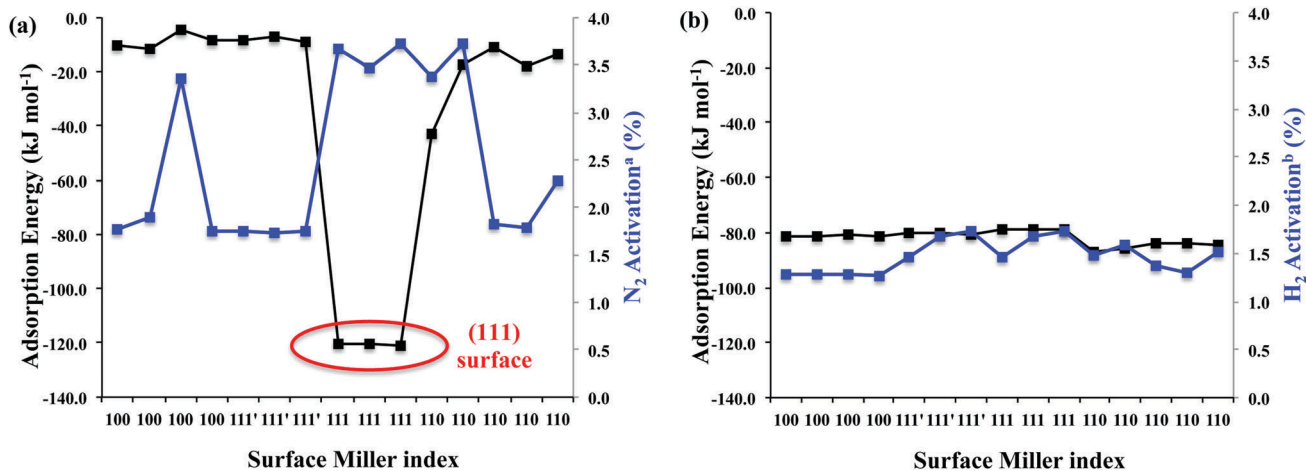


Fig. 3 (a) Nitrogen and (b) hydrogen adsorption energy and percent activation as a function of the Miller index of the surface of θ -Mn₆N₅. Percent N₂ activation is defined as $[r(\text{N}_{2,\text{ads}}) - 1.098] \times 100/1.09$. Percent H₂ activation is defined as $[r(\text{H}_{2,\text{ads}}) - 0.74] \times 100/0.74$.

plotted in Fig. 3b. The activation of the H–H bond is also very constant for these adsorption configurations, which was found to be between 1.3% and 1.7%. The most prevalent adsorption configuration of H₂ was the end-on adsorption although a tilt was also frequent and one side-on configuration was also found.

If we address the question of competitive adsorption of N₂ and H₂ on the various low Miller index surfaces of θ -Mn₆N₅, certain conclusions can be drawn based on these adsorption data. These conclusions are relevant to surfaces of this material that completely lack intrinsic nitrogen vacancies. (i) All surfaces apart from the (111) surface would have mostly hydrogen adsorbed to them, (ii) only the (111) surface would activate the N–N bond significantly, (iii) no activation of the H–H bond is observed when H₂ is adsorbed molecularly, and (iv) there is a barrier for the dissociative adsorption of H₂. For the above four reasons we have modelled the ammonia synthesis mechanism on the (111) surface and only considered hydrogenation reactions *via* an Eley–Rideal type mechanism. We were primarily interested if an Eley–Rideal mechanism is possible on this manganese nitride, as we have seen on another metal nitride, Co₃Mo₃N.²² We note that a potential Langmuir–Hinshelwood mechanism and the participation of intrinsic nitrogen vacancies in the mechanism are not considered in this study and will be the topic of a subsequent study.

3.2. Mechanism of ammonia synthesis on θ -Mn₆N₅

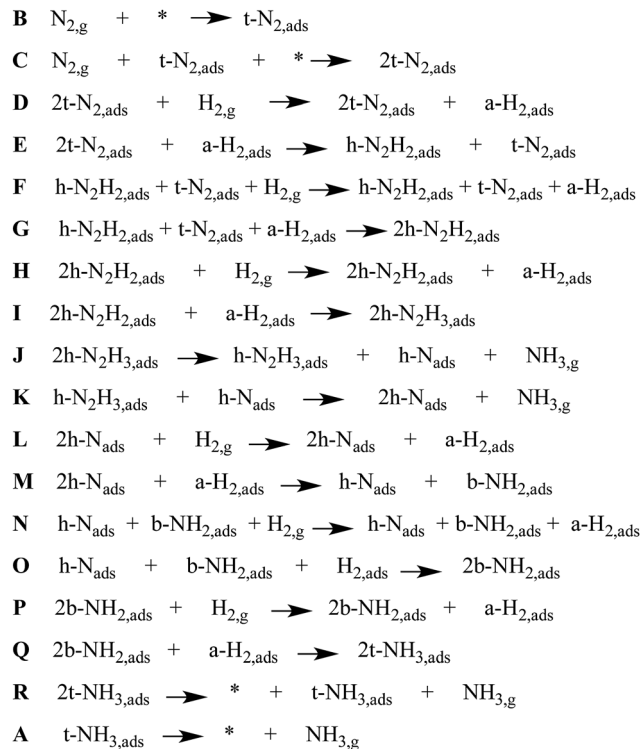
There are only a few studies of the ammonia synthesis mechanism *via* DFT on metal surfaces, nanoclusters and in organometallic complexes. Most of the early computational studies are focused on understanding the adsorption, activation and dissociation of N₂ on the Ru(0001) surface where it was found to occur primarily at steps rather than planar terraces with an activation barrier of 0.5 eV.³⁴ These findings were in agreement with earlier experimental studies by Ertl and co-workers who found through variable-temperature programmed desorption (TPD) measurements the dissociation activation barriers of 0.3 eV³⁵ and 0.6 eV³⁶ on Ru/MgO. The effect of promoters such as Na and Cs

was attributed to an electrostatic interaction that lowers the dissociation activation barrier on Ru(0001).³⁷ A partial study of the ammonia synthesis mechanism by Hu *et al.*³⁸ revealed that the hydrogenation steps of nitrogen can be as high in barrier as the N–N dissociation steps. Logadóttir and Nørskov²¹ studied the ammonia synthesis mechanism on Ru(0001) surfaces at steps and at terraces. The influence of an electric field on ΔE for the dissociation of N₂ and the formation of NNH was studied on Ru surfaces (*i.e.* (0001) and (10 $\bar{1}$ 1)) and found to increase and decrease, respectively.³⁹ Moreover, the barrier for NNH formation was found to decrease with the application of an electric field. There are few theoretical studies of electrochemical ammonia synthesis on metal nanoclusters⁴⁰ and surfaces^{10,41,42} at ambient temperature and pressure. Abghoui and Skúlason⁴³ studied a Mars–van Krevelen mechanism for electrochemical ammonia synthesis on the (100) facet of VN and the (110) facet of RuN. The same authors showed that H-assisted N₂ activation is significantly more facile than direct N₂ dissociation on transition metal nitrides (TMNs).⁴⁴ Michalsky and Steinfeld⁴⁵ explored the possibility of a solar-driven ammonia synthesis cycle where the Fe doping of manganese nitrides was found to increase the number of nitrogen vacancies. Michalsky and Peterson¹⁰ also studied the possibility of low-pressure ammonia synthesis for energy storage through chemical looping of metal nitrides. There are also a few DFT studies of the enzymatic ammonia synthesis mechanism. Rod and Nørskov⁴⁶ studied the ammonia synthesis mechanism on the FeMo cofactor in nitrogenases and found that the reaction is an associative mechanism in which end-on nitrogen adsorbed to the Fe of the cluster undergoes stepwise hydrogenation, forming first hydrazine, which after further hydrogenation forms ammonia. An associative mechanism for ammonia synthesis was studied on mono- and dinuclear molybdenum complexes and cubane-type metal–sulfido clusters.⁴⁷ More recently an associative ammonia synthesis mechanism has been studied on singly dispersed bimetallic catalyst Rh₁Co₃/CoO(011)⁴⁸ similar to how the biological fixation of N₂ occurs by nitrogenases in plants.⁴⁹ The basic characteristic of



these ammonia synthesis mechanisms is that N_2 is hydrogenated by H^+ which causes gradual weakening and dissociation of the $N \equiv N$ bond.^{50,51}

There is only our previous study of an Eley–Rideal/Mars–van Krevelen mechanism on Co_3Mo_3N which we have shown to proceed efficiently by nitrogen activation in an end-on configuration at nitrogen vacancies.²² The question that arises is whether an Eley–Rideal mechanism also exist on other metal nitrides? We have therefore modelled a complete pathway for ammonia synthesis *via* an Eley–Rideal mechanism. The reaction mechanism was modelled on the (111) surface of $\theta\text{-Mn}_6N_5$, which we found adsorbs nitrogen in an end-on configuration. The elementary reaction steps of the mechanism are shown in Scheme 1. The mechanism is an associative mechanism where the hydrogenation of N_2 causes gradual weakening of the $N \equiv N$ bond and the formation of ammonia. The reaction mechanism is depicted in Fig. 4 for which a brief description is given next. Two N_2 adsorb end-on on two manganese atoms that are separated by a nearest neighbour manganese atom (steps B and C). H_2 was then fixed by one of its atoms at a 3 Å separation from the terminal nitrogen of adsorbed N_2 (step E). The reason for this is that H_2 has an exothermic interaction with adsorbed N_2 , and therefore this increases the reaction barrier for its reaction forming diazanylidene, $=NNH_2$. Here a clear distinction has to be made with respect to an Eley–Rideal mechanism occurring at high pressure and at ambient pressure. For the first the adsorption step of H_2 could be ignored as the H_2 coming from the gas phase has high enough velocity that it would not allow any structural perturbation of the adsorbate at the adsorption site, before the bimolecular collision and reaction. So at high pressure this phenomenon would decrease the barrier for hydrogenation by an amount equal to the exothermicity of the adsorption step. Based on this rational higher pressures would act favourably on the barriers of the hydrogenation steps *via* an Eley–Rideal mechanism. In step F H_2 adsorbs again at a 3 Å distance from the terminal nitrogen of the second adsorbed N_2 . A second diazanylidene is formed upon hydrogenation (step G). Two diazanylidene adsorbates then react with an incoming molecular hydrogen (step H), forming two diazanes, $-NNH_3$ (step I). This elementary reaction step although it appears to be trimolecular is in fact bimolecular, as the two diazanylidene molecules are bound to the same substrate. In steps J and K the diazanes decompose forming two ammonia molecules and leaving nitride groups on the surface. H_2 adsorbs to the first nitride group (step L) forming an azanylium, $>NH_2$ (step M). H_2 then adsorbs to the second nitride group (step N) and a second azanylium group forms (step O). The two azanylium groups are aligned in such a way that they can stereochemically interact favourably with a H_2 coming from the gas phase (step P). There are other alignments of the azanylium group that were higher in energy and therefore not considered in the mechanism. This reaction forms two surface adsorbed ammonia groups at positions that are separated by a Mn atom (step Q). The surface adsorbed ammonia desorbs from the manganese nitride surface (step R) and an adsorbate free surface is recovered so that a new catalytic cycle can be initiated.



Scheme 1 Elementary reaction mechanism steps of ammonia synthesis *via* an Eley–Rideal mechanism on $\theta\text{-Mn}_6N_5$ -(111). The letters *t*-, *b*-, *h*- and *a*- are to indicate the adsorption position which is top, bridge, 3-fold hollow and adsorbate, respectively.

3.3. Potential energy surface of reaction

In Fig. 5 we present the potential energy curve for the Eley–Rideal mechanism on $\theta\text{-Mn}_6N_5$ -(111). The values of the activation energy barriers (E_{act}) and the reaction energies (ΔE_{react}) are tabulated in Table 1. We observe that the adsorption of N_2 to the (111) surface is exothermic by -122 kJ mol^{-1} (step B and C). This is followed by an exothermic adsorption of molecular H_2 of -76 kJ mol^{-1} (step D). The previous adsorption energies indicate that the surface will be entirely covered with end-on N_2 before the reaction of H_2 starts to occur. H_2 then reacts with the adsorbed N_2 in a high barrier process with an activation energy of 327 kJ mol^{-1} (step D to E). The adsorption of a second H_2 is only slightly exothermic by -17 kJ mol^{-1} . The next hydrogenation barrier is 478 kJ mol^{-1} high which is considerably higher than the first hydrogenation barrier which could be due to stereochemical repulsion between the diazanylidene and the incoming H_2 . Here the adsorption of H_2 to the two diazanylidenes is slightly exothermic by -20 kJ mol^{-1} . The third hydrogenation barrier of this H_2 that forms two diazanes is much smaller (182 kJ mol^{-1}) compared to the barrier of N_2 hydrogenation to NNH_2 . The decomposition of the diazane into ammonia and a nitride is slightly endothermic by 11 kJ mol^{-1} for the first ammonia (step J) and 67 kJ mol^{-1} for the second ammonia (step K). The adsorption of the fourth H_2 to the nitride was found to be again exothermic by -78 kJ mol^{-1} (step L) with a high hydrogenation barrier of 265 kJ mol^{-1} (step L to M)



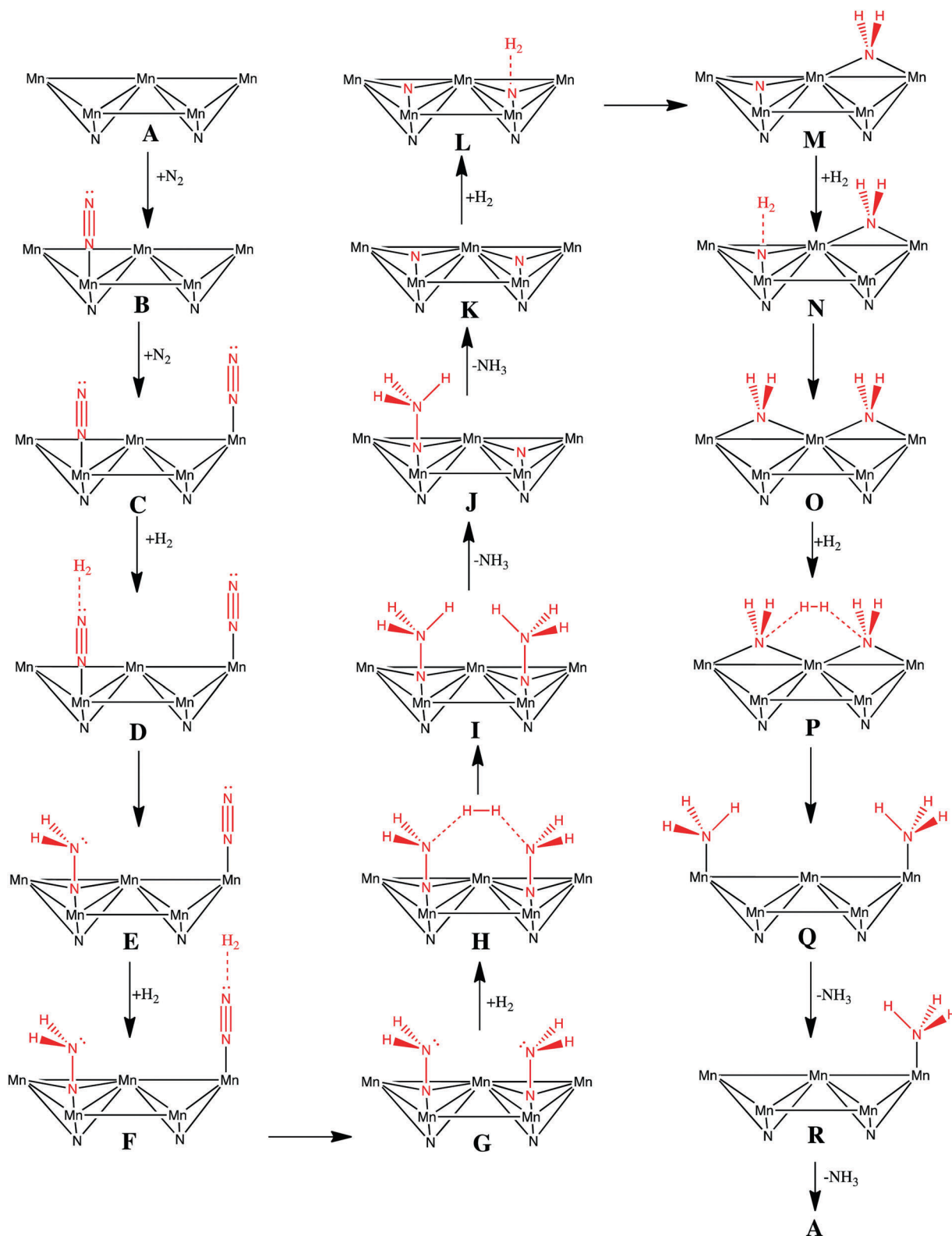


Fig. 4 Reaction mechanism for ammonia synthesis via an Eley-Rideal mechanism on θ - Mn_6N_5 -(111).

in order to form azanylium. This was followed by the adsorption of the fifth H_2 , which was again found to be exothermic by -81 kJ mol^{-1} (step N).

Interestingly, the hydrogenation of the nitride into an azanylium that had an adjacent azanylium (step N to O) had a barrier of only 2 kJ mol^{-1} . We note that this elementary



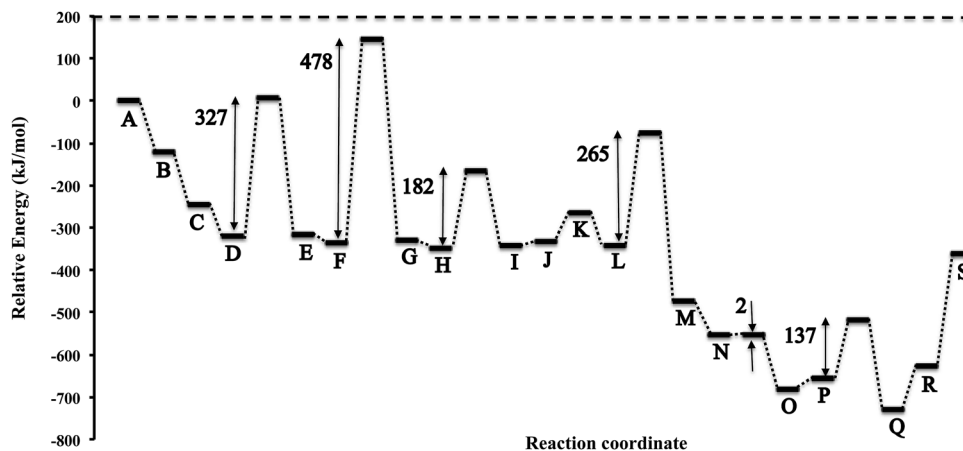


Fig. 5 Potential energy diagram of the ammonia synthesis reaction *via* an Eley–Rideal mechanism on the θ - Mn_6N_5 -(111) slab without surface nitrogen vacancies.

Table 1 Activation energy barriers (E_{act}) and the reaction energies (ΔE_{react}) for an Eley–Rideal mechanism for ammonia synthesis

Label	ΔE_{react} (kJ mol ⁻¹)	Label	E_{act} (kJ mol ⁻¹)
ΔE (AB)	-122		
ΔE (BC)	-122		
ΔE (CD)	-76		
ΔE (DE)	3	E_{act} (DE)	327
ΔE (EF)	-17	E_{act} (FG)	478
ΔE (FG)	6		
ΔE (GH)	-20		
ΔE (HI)	5	E_{act} (HI)	182
ΔE (IJ)	11		
ΔE (JK)	67		
ΔE (KL)	-78		
ΔE (LM)	-131	E_{act} (LM)	265
ΔE (MN)	-81		
ΔE (NO)	-126	E_{act} (NO)	2
ΔE (OP)	26		
ΔE (PQ)	-76	E_{act} (PQ)	137
ΔE (QR)	103		
ΔE (RS)	267		

reaction step may be a low energy pathway for the hydrogenation of surface nitrides in metal nitrides that have 3-fold bound nitrogen which is relevant to the formation of nitrogen vacancies on manganese nitrides on surfaces such as (111)' shown in Fig. 2.

The adsorption of the sixth H_2 becomes endothermic by 26 kJ mol⁻¹ (step P). The subsequent barrier for hydrogenation is 137 kJ mol⁻¹ which forms two ammonia molecules adsorbed to the surface. The desorption of these ammonia from the surface is highly endothermic, with ΔE of 103 kJ mol⁻¹ (step R) and 267 kJ mol⁻¹, respectively, for the desorption of the first and second ammonia. From the above potential energy surface of the reaction it is apparent that the mechanism does not proceed at low temperatures *via* Eley–Rideal chemistry although it is possible at high T . The hydrogenation barriers, apart from the 5th hydrogenation barrier, are high and therefore other mechanisms should be examined, such as Langmuir–Hinshelwood mechanisms or other pathways that occur at nitrogen vacancies. It is intriguing though that an Eley–Rideal

mechanism for the hydrogenation of surface activated nitrogen exists on $\text{Co}_3\text{Mo}_3\text{N}$, which could be the underlying reason for the known enhanced activity of this tertiary metal nitride.

Conclusions

The adsorption and activation of N_2 and H_2 was studied on every low Miller index surface of $\text{Mn}_6\text{N}_{5+x}$ ($x = 1$). The adsorption of N_2 was molecular in the absence of nitrogen vacancies with adsorption energies of -5 kJ mol⁻¹ to -43 kJ mol⁻¹. Only the (111) surface was found to activate N_2 , 4%, with a large exothermic adsorption energy (-120 kJ mol⁻¹). This surface would inhibit the poisoning of the surface by H_2 which was found to bind strongly (-79 kJ mol⁻¹ to -87 kJ mol⁻¹) in a molecular form in the absence of nitrogen vacancies to all surfaces of $\text{Mn}_6\text{N}_{5+x}$ ($x = 1$). The complete reaction Eley–Rideal reaction pathway was modelled on the (111) surface of $\text{Mn}_6\text{N}_{5+x}$ ($x = 1$). It shows that an Eley–Rideal mechanism, in contrast with what was found on $\text{Co}_3\text{Mo}_3\text{N}$, would only proceed at high temperatures. This suggests that other mechanisms such as Langmuir–Hinshelwood or mechanisms that invoke nitrogen vacancies should be the focus of future studies of mechanistic pathways.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This study was supported by EPSRC funding (EP/L026317/1, EP/K014714/1). *Via* our membership of the UK's HEC Materials Chemistry Consortium, which is funded by EPSRC (EP/L000202/1), this work used the ARCHER UK National Supercomputing Service (<http://www.archer.ac.uk>). The authors acknowledge the use of the Grace High Performance Computing Facility



(Grace@UCL), and associated support services, in the completion of this work.

References

- W. R. L. Lambrecht, M. Prokhorodko and M. S. Miao, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2003, **68**, 174411.
- A. Leineweber, R. Niewa, H. Jacobs and W. Kockelmann, *J. Mater. Chem.*, 2000, **10**, 2827–2834.
- G. Kreiner and H. Jacobs, *J. Alloys Compd.*, 1992, **183**, 345–362.
- G. W. Wiener and J. A. Berger, *Trans. Am. Inst. Min., Metall. Pet. Eng.*, 1955, **203**, 360–368.
- M. N. Eddine and E. F. Bertaut, *Solid State Commun.*, 1977, **23**, 147–150.
- M. Mekata, J. Haruna and H. Takaki, *J. Phys. Soc. Jpn.*, 1968, **25**, 234–238.
- F. Lihl, P. Ettmayer and A. Kutzelnigg, *Z. Metallkd.*, 1962, **53**, 715–719.
- K. Suzuki, T. Kaneko, H. Yoshida, Y. Obi, H. Fujimori and H. Morita, *J. Alloys Compd.*, 2000, **306**, 66–71.
- A. Leineweber, R. Niewa, H. Jacobs and W. Kockelmann, *J. Mater. Chem.*, 2000, **10**, 2827–2834.
- R. Michalsky, A. M. Avram, B. A. Peterson, P. H. Pfromm and A. A. Peterson, *Chem. Sci.*, 2015, **6**, 3965–3974.
- S. Laassiri, C. D. Zeinalipour-Yazdi, C. R. A. Catlow and J. S. J. Hargreaves, *Appl. Catal., B*, 2018, **223**, 60–66.
- T. J. Wood, J. W. Makepeace and W. I. F. David, *Phys. Chem. Chem. Phys.*, 2018, **20**, 8547–8553.
- J. Guo, P. Wang, G. Wu, A. Wu, D. Hu, Z. Xiong, J. Wang, P. Yu, F. Chang, Z. Chen and P. Chen, *Angew. Chem., Int. Ed.*, 2015, **54**, 2950–2954.
- P. Yu, J. Guo, L. Liu, P. Wang, G. Wu, F. Chang and P. Chen, *ChemSusChem*, 2016, **9**, 364–369.
- C. Walter, W. Menezes Prashanth, S. Orthmann, J. Schuch, P. Connor, B. Kaiser, M. Lerch and M. Driess, *Angew. Chem., Int. Ed.*, 2017, **57**, 698–702.
- U. Zwicker, *Z. Metallkd.*, 1951, **42**, 274–276.
- M. D. Lyutaya and A. B. Goncharuk, *Powder Metall. Met. Ceram.*, 1977, **16**, 208–212.
- N. Otsuka, Y. Hanawa and S. Nagakura, *Phys. Status Solidi*, 1977, **43**, K127–K129.
- J. S. J. Hargreaves, *Appl. Petrochem. Res.*, 2014, **4**, 3–10.
- G. E. Veitch, K. L. Bridgwood and S. V. Ley, *Org. Lett.*, 2008, **10**, 3623–3625.
- Á. Logadóttir and J. K. Nørskov, *J. Catal.*, 2003, **220**, 273–279.
- C. D. Zeinalipour-Yazdi, J. S. J. Hargreaves and C. R. A. Catlow, *J. Phys. Chem. C*, 2018, **122**, 6078–6082.
- C. D. Zeinalipour-Yazdi, J. S. J. Hargreaves and C. R. A. Catlow, *J. Phys. Chem. C*, 2015, **119**, 28368–28376.
- C. D. Zeinalipour-Yazdi, J. S. J. Hargreaves, S. Laassiri and C. R. A. Catlow, *Phys. Chem. Chem. Phys.*, 2017, **19**, 11968–11974.
- C. D. Zeinalipour-Yazdi, J. S. J. Hargreaves and C. R. A. Catlow, *J. Phys. Chem. C*, 2016, **120**, 21390–21398.
- H. J. Monkhorst and J. D. Pack, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1976, **13**, 5188.
- G. Kresse and J. Furthmüller, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1996, **54**, 11169–11186.
- G. Kresse and J. Hafner, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1993, **47**, 558.
- J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **77**, 3865.
- G. Kresse and D. Joubert, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1999, **59**, 1758–1775.
- P. E. Blöchl, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **50**, 17953.
- S. Grimme, J. Antony, S. Ehrlich and H. Krieg, *J. Chem. Phys.*, 2010, **132**, 154104.
- G. Mills, H. Jónsson and G. K. Schenter, *Surf. Sci.*, 1995, **324**, 305–337.
- S. Dahl, A. Logadóttir, R. C. Egeberg, J. H. Larsen, I. Chorkendorff, E. Törnqvist and J. K. Nørskov, *Phys. Rev. Lett.*, 1999, **83**, 1814–1817.
- F. Rosowski, O. Hinrichsen, M. Muhler and G. Ertl, *Catal. Lett.*, 1996, **36**, 229–235.
- O. Hinrichsen, F. Rosowski, A. Hornung, M. Muhler and G. Ertl, *J. Catal.*, 1997, **165**, 33–44.
- J. J. Mortensen, B. Hammer and J. K. Nørskov, *Phys. Rev. Lett.*, 1998, **80**, 4333–4336.
- C. J. Zhang, M. Lynch and P. Hu, *Surf. Sci.*, 2002, **496**, 221–230.
- R. Manabe, H. Nakatsubo, A. Gondo, K. Murakami, S. Ogo, H. Tsuneki, M. Ikeda, A. Ishikawa, H. Nakai and Y. Sekine, *Chem. Sci.*, 2017, **8**, 5434–5439.
- J. G. Howalt, T. Bligaard, J. Rossmeisl and T. Vegge, *Phys. Chem. Chem. Phys.*, 2013, **15**, 7785–7795.
- E. Skulason, T. Bligaard, S. Gudmundsdóttir, F. Studt, J. Rossmeisl, F. Abild-Pedersen, T. Vegge, H. Jonsson and J. K. Nørskov, *Phys. Chem. Chem. Phys.*, 2012, **14**, 1235–1245.
- Y. Abghoui, A. L. Garden, V. F. Hlynsson, S. Björgvinsdóttir, H. Olafsdóttir and E. Skulason, *Phys. Chem. Chem. Phys.*, 2015, **17**, 4909–4918.
- Y. Abghoui and E. Skúlasson, *Proc. Comp. Sci.*, 2015, **51**, 1897–1906.
- Y. Abghoui and E. Skúlasson, *Catal. Today*, 2017, **286**, 69–77.
- R. Michalsky, P. H. Pfromm and A. Steinfeld, *Interface Focus*, 2015, **5**, 1–9.
- T. H. Rod and J. K. Nørskov, *J. Am. Chem. Soc.*, 2000, **122**, 12751–12763.
- H. Tanaka, Y. Nishibayashi and K. Yoshizawa, *Acc. Chem. Res.*, 2016, **49**, 987–995.
- X.-L. Ma, J.-C. Liu, H. Xiao and J. Li, *J. Am. Chem. Soc.*, 2018, **140**, 46–49.
- S. Kuriyama, K. Arashiba, K. Nakajima, Y. Matsuo, H. Tanaka, K. Ishii, K. Yoshizawa and Y. Nishibayashi, *Nat. Commun.*, 2016, **7**, 12181.
- I. Coric, B. Q. Mercado, E. Bill, D. J. Vinyard and P. L. Holland, *Nature*, 2015, **526**, 96.
- B. M. Hoffman, D. Lukoyanov, Z. Y. Yang, D. R. Dean and L. C. Seefeldt, *Chem. Rev.*, 2014, **114**, 4041.

