

Low-T Mechanisms of Ammonia Synthesis on $\text{Co}_3\text{Mo}_3\text{N}$

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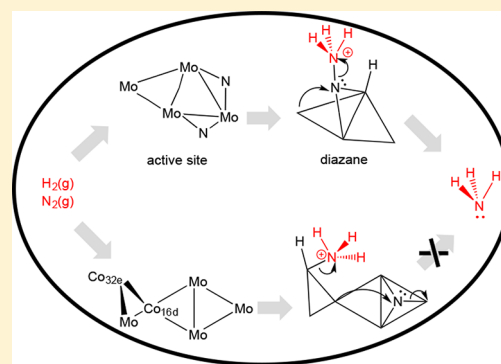
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Supporting Information

ABSTRACT: Dispersion-corrected periodic DFT calculations have been applied to elucidate the Langmuir–Hinshelwood (dissociative) and an Eley–Rideal/Mars–van Krevelen (associative) mechanism for ammonia synthesis over $\text{Co}_3\text{Mo}_3\text{N}$ surfaces, in the presence of surface defects. Comparison of the two distinct mechanisms clearly suggests that apart from the conventional dissociative mechanism, there is another mechanism that proceeds via hydrazine and diazane intermediates that are formed by Eley–Rideal type chemistry, where hydrogen reacts directly with surface activated nitrogen, in order to form ammonia at considerably milder conditions. This result clearly suggests that via surface defects ammonia synthesis activity can be enhanced at milder conditions on one of the most active catalysts for ammonia synthesis.



INTRODUCTION

The development of alternative nitrogen-fixation processes could have a profound economic and environmental impact, as more than 50% of ammonia for soil fertilizers is produced by man. The other 50% is produced naturally by nitrogenase¹ in nitrogen fixation plants where the reduction of N_2 occurs by the FeMo-cofactor.^{2–4} Currently ammonia synthesis industrially is achieved mostly via the classical Haber–Bosch (H-B) process,^{5–7} with a Fe– K_2O – Al_2O_3 catalyst, which operates under high temperatures (>400 °C) and pressures (150–200 atm). Some industrial plants have changed to the Kellogg advanced ammonia process that uses a graphite-supported alkali/alkaline-earth promoted Ru catalyst, which operates at milder conditions but is expensive.⁸ Researchers are now seeking catalytic materials that could potentially produce ammonia at low temperatures ($T = 200$ – 300 °C) in order to save energy. The generally accepted mechanism for ammonia synthesis on the iron K-promoted Fe catalyst is a Langmuir–Hinshelwood (L-H) mechanism which is dissociative.^{9,10} An L–H mechanism has also been found on the Ru(0001) surfaces, via density functional theory (DFT) calculations in order to model the Ru–graphite catalyst.^{11,12} The rate-determining step (RDS) was found to be the activation step for N_2 ,¹³ due to the high bond dissociation enthalpy of N_2 (945 kJ mol^{-1}) compared to H_2 (436 kJ mol^{-1}).¹⁴ An associative mechanism has recently been found on Ru that proceeds through a $-\text{N}_2\text{H}$ intermediate.¹⁵

$\text{Co}_3\text{Mo}_3\text{N}$ is known to be a more active catalyst than the one currently used by the industry, especially when doped with cesium.^{16–20} Although the high activity of this and related materials has been established, there is a need to understand its origin and to elucidate the mechanism of ammonia synthesis.

Only a few previous DFT studies have attempted to model the kinetics of ammonia synthesis on this catalyst, based on linear-scaling relationships, that correlated the dissociation barrier to the adsorption energy of N_2 ,^{16,21} where an L–H mechanism was assumed.¹³ However, a detailed study of the elementary mechanistic steps and the role of surface defects in particular is currently lacking. In two recent DFT studies we have identified possible sites for the adsorption and activation of the reactants of the ammonia synthesis reaction on a model $\text{Co}_3\text{Mo}_3\text{N}$ surface with heterogeneity due to surface nitrogen vacancies.²² Such vacancies are present in large concentrations even at ambient temperatures (i.e., 10^{13} cm^{-2}) and can efficiently activate N_2 .²³ Our earlier work found that there are two activation sites for N_2 , the first is a surface cavity, where N_2 is bound *side-on* a Co_8 cluster at the 16d Wyckoff site, where N_2 is activated 21% (as measured by the percentage change in the N–N bond length); the second, a somewhat weaker activation site (11%) that was found to be located at surface 3f-nitrogen vacancies for which we have suggested that nitrogen vacancies can participate in the elementary reaction steps toward the formation of ammonia.²² Indeed, recent DFT studies show that N-vacancies can participate in the mechanism for the electrochemical reduction of ammonia on Zr, Nb, Cr, and V mononitrides^{24,25} and in the two-step solar-energy driven ammonia synthesis on metal-nitrides.^{26,27} Additionally, in a recent DFT study, the electrochemical synthesis of ammonia was studied for the Mars–van Krevelen mechanism on group III–VII transition metal nitrides and a N–N was found to be

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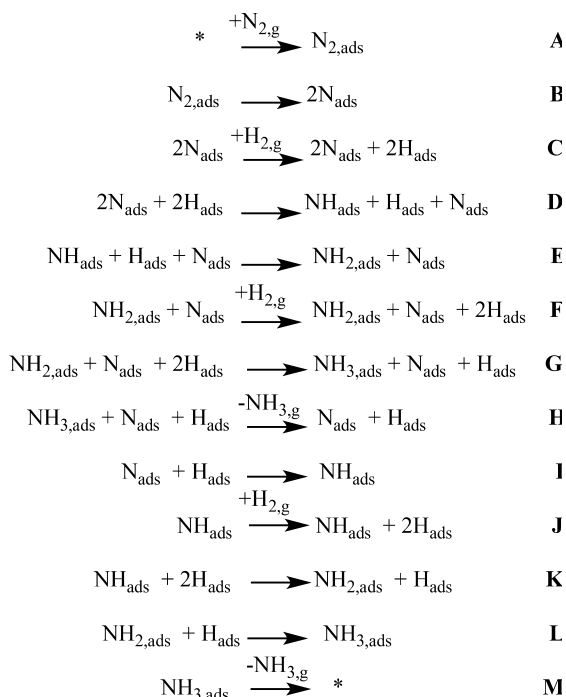
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suitable, whereas the other metal nitrides decompose to their pure metals.²⁸

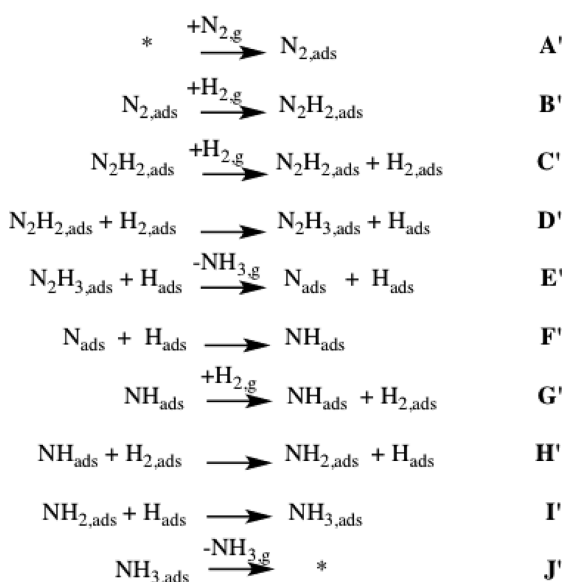
RESULTS AND DISCUSSION

In this communication we have considered a Langmuir–Hinshelwood (L-H; see Schemes 1 and 3) and an Eley–Rideal/Mars–van Krevelen (E-R/MvK; see Schemes 2 and 4) mechanism for ammonia synthesis on $\text{Co}_3\text{Mo}_3\text{N}$ -(111) surfaces.

Scheme 1. Elementary Reaction Steps for the L-H (Dissociative) Mechanism



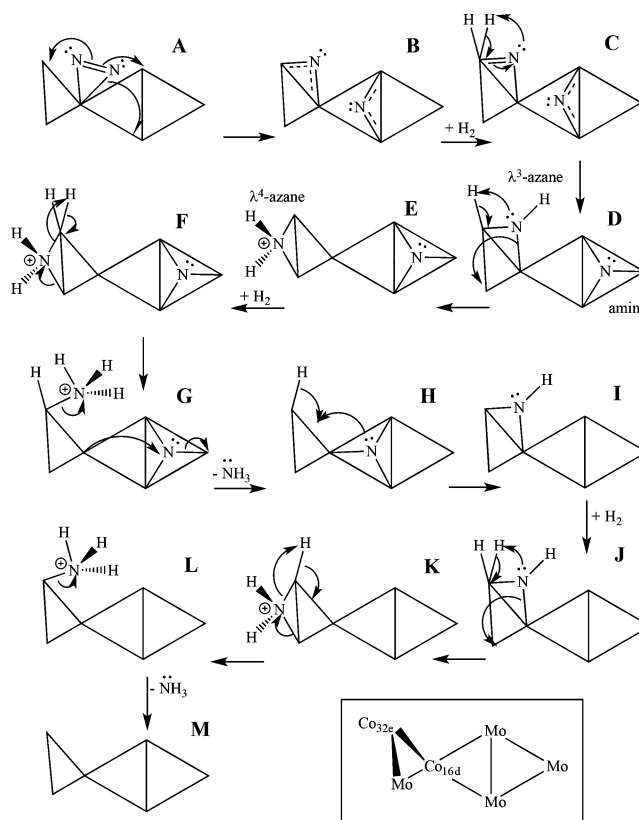
Scheme 2. Elementary Reaction Steps for the E-R/MvK (Associative) Mechanism



In particular, we have applied dispersion-corrected (D3) DFT in order to calculate the potential energy diagram for

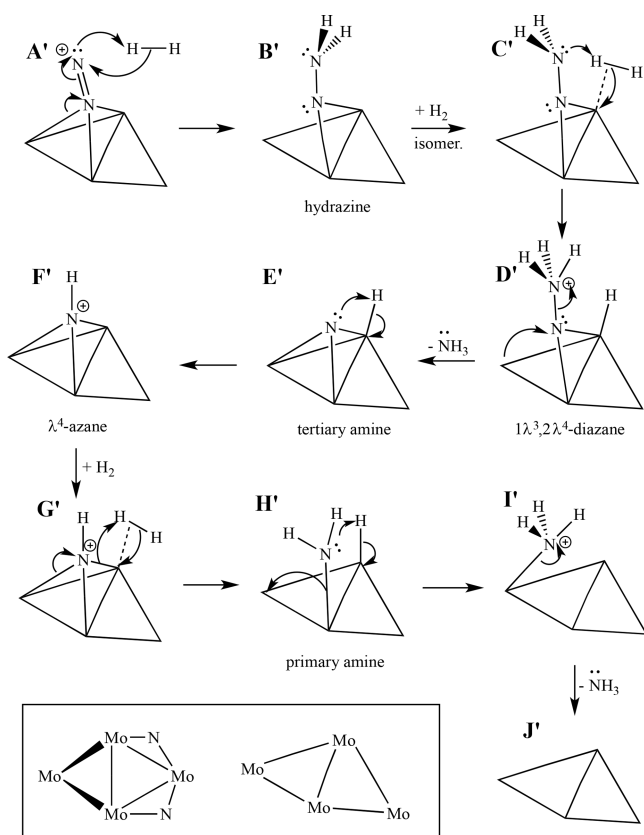
ammonia synthesis for two mechanistic pathways for ammonia synthesis on the (111)-surfaces of $\text{Co}_3\text{Mo}_3\text{N}$ in the presence of surface defects, such as nitrogen vacancies and intrinsic surface cavities. The computational methodology has been described in detail elsewhere.^{22,23} Activation barriers were obtained by the nudged elastic band (NEB) method in which the barrier was modeled first by 10 images to acquire the details of the potential energy surface. Once the various intermediates were found and fully optimized, the transition states were located by a separate NEB run of 3 images. Remarkably, we find that there are two possible mechanisms for ammonia synthesis; the first is the commonly known L-H mechanism⁸ shown in Scheme 3;

Scheme 3. L-H (Dissociative) Mechanism for Ammonia Synthesis on $\text{Co}_3\text{Mo}_3\text{N}$



the second resembles an Eley–Rideal/Mars–van Krevelen (associative) mechanism in which surface lattice nitrogen participates as shown in Scheme 4. The relative energy diagram of the two reactions is shown in Figure 1.

L-H (Dissociative) Ammonia Synthesis Mechanism. N_2 adsorbs generally less favorably than hydrogen on $\text{Co}_3\text{Mo}_3\text{N}$ -(111) surfaces. However, at the surface cavity, the adsorption of both N_2 and H_2 is slightly endothermic: 40 and 21 kJ/mol, respectively (other less active adsorption sites have exothermic adsorption energies), indicating that some pressure maybe required. A detailed study of the various adsorption sites is given in ref 23, which showed that H_2 and N_2 can essentially coadsorb at the activation sites; the activation barriers of the L-H mechanism has hydrogenation barriers higher than the mechanism modeled on $\text{Ru}(0001)$, which indicates that at higher-T there maybe additional reaction mechanisms occurring on nitrogen free $\text{Co}_3\text{Mo}_3\text{N}$ surfaces. Therefore, in the L-H modeled here the mechanism for NH_3 synthesis follows the

Scheme 4. E-R/MvK (Associative) Mechanism for Ammonia Synthesis on $\text{Co}_3\text{Mo}_3\text{N}$ 

route A: gas phase nitrogen adsorbs side-on to the catalyst and is activated by 21% by stretching of the N–N bond; B: activated N_2^* undergoes dissociation through a relatively low activation barrier that results in two bridged-N intermediates; C: H_2 adsorbs dissociatively on a Co_8 cluster, forming two atomic H species; D: $-\text{H}$ chemisorbed on the Co_8 cluster reacts with a

bridged-N forming an λ^3 -azane intermediate ($>\text{NH}$), which, due to steric congestion, causes the adjacent bridge-N to move to a hollow position at Mo_3 sites; E: H reacts with $>\text{NH}$ forming $>\text{NH}_2$ at a bridge position formed between $\text{Co}_{32e}-\text{Mo}$; F: another hydrogen chemisorbs dissociatively on a Co_8 cluster; G: the 3f-bound-N to the Mo_3 hollow moves due to surface diffusion to the adjacent $\text{Co}-\text{Mo}_2$ hollow, while displacing NH_3 chemisorbed on the Co_8 cluster; H, I: 3f-bound-N at $\text{Co}_{16d}-\text{Mo}_2$ hollow moves to adjacent bridge position at Co_8 reacting with H and forming an λ^3 -azane intermediate; J: another hydrogen dissociates on the Co_8 cluster forming two $-\text{H}$ species; K: $-\text{H}$ reacts with λ^3 -azane intermediate ($>\text{NH}$) forming a λ^4 -azane intermediate ($>\text{NH}_2$) at the adjacent $\text{Co}_{32e}-\text{Mo}$ site; L: $-\text{H}$ reacts with $>\text{NH}_2$ forming surface-adsorbed ammonia ($-\text{NH}_3$) on the Co_8 clusters; M: ammonia desorbs from the Co_8 clusters in a relatively high barrier process; the surface cavity is free and the catalytic cycle resumes, starting from step A.

E-R/MvK (Associative) Ammonia Synthesis Mechanism. In the E-R/MvK mechanism, ammonia is synthesized at the 3f-nitrogen vacancy sites on the Mo_3N framework. The active site is shown in the box of Scheme 4 and has some similarities with FeMo-cofactor in nitrogenases: A': nitrogen adsorbs in an end-on configuration at a 3-fold N-vacancy on the Mo_3N framework with 11% activation; B': gas phase hydrogen reacts directly in an Eley–Rideal mechanism with surface activated nitrogen forming a *trans* hydrazine intermediate ($>\text{NNH}_2$, similar to $\text{Mo}=\text{NNH}_2^{29}$ and $\text{Fe}-\text{NNH}_2^{30}$), in a low barrier process; C': hydrogen adsorbs molecularly onto the Mo_3N framework adjacent to the nitrogen vacancy site, while *trans* hydrazine isomerizes into a conformation that can react with H_2 ; D': H_2 reacts with the hydrazine-like intermediate forming diazane and $-\text{H}$; E': Diazane readily decomposes due to its positive charge into a tertiary amine and $-\text{H}$; F': $-\text{H}$ reacts with the tertiary amine forming an azane bound to a Mo_3 hollow; G': hydrogen adsorbs molecularly at Mo_3N framework adjacent to where previously the nitrogen vacancy site was located; H': the precursor-mediated adsorbed state of H_2 reacts with azane forming primary amine, through a high barrier step; J': ammonia desorbs from the Co_8 clusters in a relatively high barrier process; the surface cavity is free and the catalytic cycle resumes, starting from step A.

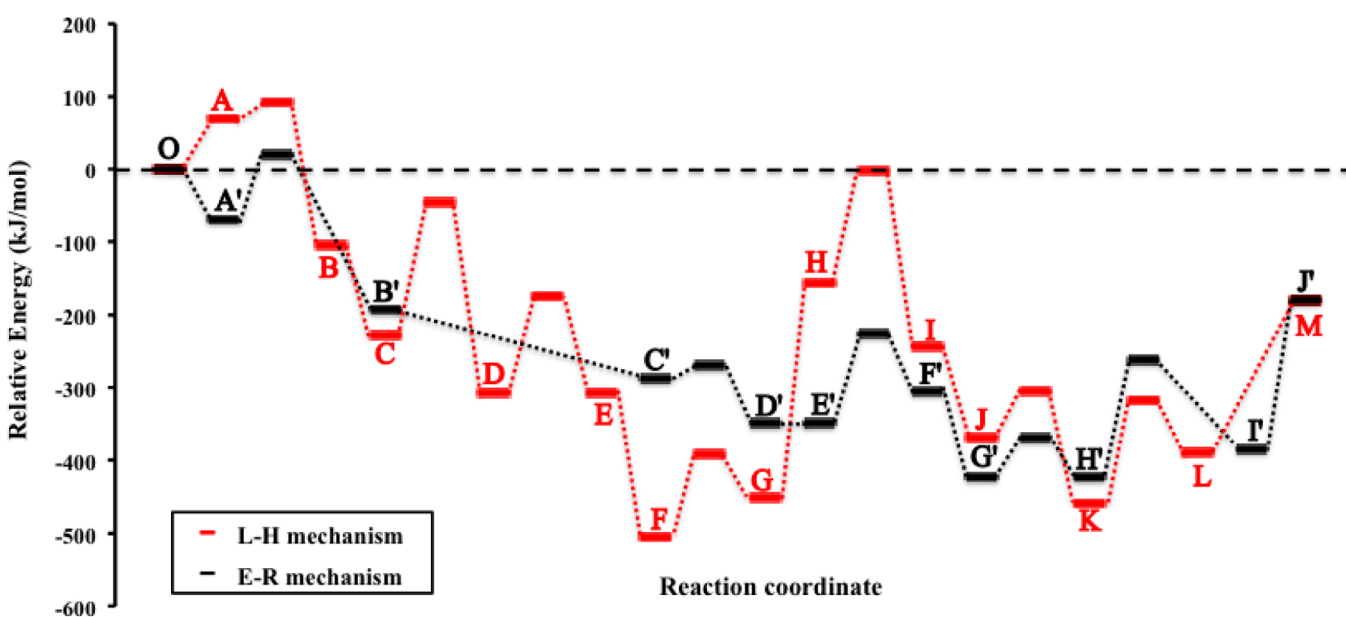


Figure 1. Relative energy diagram of L-H (red) and E-R/MvK (black) mechanism for ammonia synthesis on $\text{Co}_3\text{Mo}_3\text{N}$ surfaces. Letter labels given in Schemes 1–4

I': the second $-H$ reacts with the primary amine forming surface chemisorbed ammonia that is bound to a Mo_3 hollow site; J': finally, the second NH_3 desorbs through a high barrier step >200 kJ/mol.

Comparison of the two distinct mechanisms shows that the barriers for the corresponding hydrogenation steps are lower for the associative E-R/MvK (associative) mechanism and the barrier of the rate-determining step (RDS) is 90 kJ mol⁻¹ smaller in the E-R/MvK mechanism. Additionally, the E-R/MvK mechanism lacks the second and third activation barrier in the L-H mechanism that are high. These results clearly suggest that apart from the conventional L-H (dissociative) mechanism, there is also a mechanism which proceeds via diazane and hydrazine intermediates that are formed by a direct associative Eley–Rideal/Mars–van Krevelen mechanism, where molecular hydrogen reacts directly with surface activated nitrogen, in order to form ammonia at considerably milder conditions. Intriguingly, if the mechanism is cycled in two phases, phase I: steps A' through D' as depicted in Scheme 4, and phase II: the catalyst is activated to form nitrogen vacancies, then barriers are <70 kJ/mol for phase I which produces half the stoichiometric amount of ammonia. The second half would be generated by annealing the catalyst to cause desorption of NH_3 ($\Delta H_{des} = 204$ kJ mol⁻¹) and regenerate the amount of nitrogen vacancies on the catalyst surface.

CONCLUSIONS

We have studied with dispersion-corrected DFT two complete mechanisms for ammonia synthesis on cobalt molybdenum nitride. The first that occurs at surface cavities is a Langmuir–Hinshelwood (dissociative) mechanism, and the second that occurs at surface nitrogen vacancies is an Eley–Rideal/Mars–van Krevelen (associative) mechanism. We show that, in the Eley–Rideal/Mars van Krevelen mechanism, hydrogen reacts directly with surface activated nitrogen, in order to form ammonia, at considerably milder conditions. The mechanism proceeds via hydrazine and diazane intermediates. This result clearly suggests that, via surface defects, ammonia synthesis activity can be enhanced at milder conditions on one of the most active catalysts for ammonia synthesis.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.jpcc.7b12364.

The coordinates of the structure of the various intermediates and transition states (Figure S2) and the detailed barrier calculations (Figure S1) (PDF)

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Notes

The authors declare no competing financial interest.

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REFERENCES

- (1) Owens, C. P.; Katz, F. E. H.; Carter, C. H.; Oswald, V. F.; Tezcan, F. A. Tyrosine-Coordinated P-Cluster in G. diazotrophicus Nitrogenase: Evidence for the Importance of O-Based Ligands in Conformationally Gated Electron Transfer. *J. Am. Chem. Soc.* **2016**, *138*, 10124–10127.
- (2) Lukoyanov, D.; Khadka, N.; Yang, Z.-Y.; Dean, D. R.; Seefeldt, L. C.; Hoffman, B. M. Reductive Elimination of H₂ Activates Nitrogenase to Reduce the N≡N Triple Bond: Characterization of the E4(4H) Janus Intermediate in Wild-Type Enzyme. *J. Am. Chem. Soc.* **2016**, *138*, 10674–10683.
- (3) Čorić, I.; Holland, P. L. Insight into the Iron–Molybdenum Cofactor of Nitrogenase from Synthetic Iron Complexes with Sulfur, Carbon, and Hydride Ligands. *J. Am. Chem. Soc.* **2016**, *138*, 7200–7211.
- (4) Siegbahn, P. E. M. Model Calculations Suggest that the Central Carbon in the FeMo-Cofactor of Nitrogenase Becomes Protonated in the Process of Nitrogen Fixation. *J. Am. Chem. Soc.* **2016**, *138*, 10485–10495.
- (5) Haber, F. Über die Darstellung des Ammoniaks aus Stickstoff und Wasserstoff. *Naturwissenschaften* **1922**, *10*, 1041–1049.
- (6) Haber, F. Bemerkung zu vorstehender Notiz. *Naturwissenschaften* **1923**, *11*, 339–340.
- (7) Smil, V. *Enriching the Earth: Fritz Haber, Carl Bosch, and the Transformation of World Food Production*; MIT Press: Cambridge, MA, 2004.
- (8) Saadatjou, N.; Jafari, A.; Sahebdehfar, S. Ruthenium Nanocatalysts for Ammonia Synthesis: A Review. *Chem. Eng. Commun.* **2015**, *202*, 420–448.
- (9) Ertl, G. Surface Science and Catalysis—Studies on the Mechanism of Ammonia Synthesis: The P. H. Emmett Award Address. *Catal. Rev.: Sci. Eng.* **1980**, *21*, 201–223.
- (10) Bowker, M. Modelling of Ammonia Synthesis Kinetics. *Catal. Today* **1992**, *12*, 153–163.
- (11) Logadóttir, Á.; Nørskov, J. K. Ammonia Synthesis over a Ru(0001) Surface Studied by Density Functional Calculations. *J. Catal.* **2003**, *220*, 273–279.
- (12) Zhang, C. J.; Lynch, M.; Hu, P. A Density Functional Theory Study of Stepwise Addition Reactions in Ammonia Synthesis on Ru(0001). *Surf. Sci.* **2002**, *496*, 221–230.
- (13) Jacobsen, C. J. H.; Dahl, S.; Clausen, B. S.; Bahn, S.; Logadóttir, Á.; Nørskov, J. Catalyst Design by Interpolation in the Periodic Table: Bimetallic Ammonia Synthesis Catalysts. *J. Am. Chem. Soc.* **2001**, *123*, 8404–8405.
- (14) Blanksby, S. J.; Ellison, G. B. Bond Dissociation Energies of Organic Molecules. *Acc. Chem. Res.* **2003**, *36*, 255–263.
- (15) Garden, A. L.; Skúlason, E. The Mechanism of Industrial Ammonia Synthesis Revisited: Calculations of the Role of the Associative Mechanism. *J. Phys. Chem. C* **2015**, *119*, 26554–26559.
- (16) Boisen, A.; Dahl, S.; Jacobsen, C. J. H. Promotion of Binary Nitride Catalysts: Isothermal N₂ Adsorption, Microkinetic Model, and Catalytic Ammonia Synthesis Activity. *J. Catal.* **2002**, *208*, 180–186.
- (17) Jacobsen, C. J. H. Novel Class of Ammonia Synthesis Catalysts. *Chem. Commun.* **2000**, 1057–1058.
- (18) Jacobsen, C. J. H.; Dahl, S.; Clausen, B. S.; Bahn, S.; Logadóttir, Á.; Nørskov, J. Catalyst Design by Interpolation in the Periodic Table: Bimetallic Ammonia Synthesis Catalysts. *J. Am. Chem. Soc.* **2001**, *123*, 8404–8405.
- (19) Kojima, R.; Aika, K.-I. Cobalt Molybdenum Bimetallic Nitride Catalysts for Ammonia Synthesis. *Chem. Lett.* **2000**, *29*, 514–515.

(20) Kojima, R.; Aika, K.-I. Cobalt Molybdenum Bimetallic Nitride Catalysts for Ammonia Synthesis: Part 1. Preparation and Characterization. *Appl. Catal., A* **2001**, *215*, 149–160.

(21) Nørskov, J. K.; Bligaard, T.; Logadóttir, A.; Bahn, S.; Hansen, L. B.; Bollinger, M.; Bengard, H.; Hammer, B.; Sljivančanin, Z.; Mavrikakis, M.; et al. Universality in Heterogeneous Catalysis. *J. Catal.* **2002**, *209*, 275–278.

(22) Zeinalipour-Yazdi, C. D.; Hargreaves, J. S. J.; Catlow, C. R. A. Nitrogen Activation in a Mars–van Krevelen Mechanism for Ammonia Synthesis on Co₃Mo₃N. *J. Phys. Chem. C* **2015**, *119*, 28368–28376.

(23) Zeinalipour-Yazdi, C. D.; Hargreaves, J. S. J.; Catlow, C. R. A. DFT-D3 study of molecular N₂ and H₂ activation on Co₃Mo₃N surfaces. *J. Phys. Chem. C* **2016**, *120*, 21390–21398.

(24) Abghoui, Y.; Garden, A. L.; Hlynsson, V. F.; Björgvinsdóttir, S.; Olafsdóttir, H.; Skúlason, E. Enabling Electrochemical Reduction of Nitrogen to Ammonia at Ambient Conditions Through Rational Catalyst Design. *Phys. Chem. Chem. Phys.* **2015**, *17*, 4909–4918.

(25) Abghoui, Y.; Garden, A. L.; Howalt, J. G.; Vegge, T.; Skúlason, E. Electroreduction of N₂ to Ammonia at Ambient Conditions on Mononitrides of Zr, Nb, Cr, and V: A DFT Guide for Experiments. *ACS Catal.* **2016**, *6*, 635–646.

(26) Michalsky, R.; Pfromm, P. H.; Steinfeld, A. Rational Design of Metal Nitride Redox Materials for Solar-Driven Ammonia Synthesis. *Interface Focus* **2015**, *5*, 20140084–10.

(27) Michalsky, R.; Avram, A. M.; Peterson, B. A.; Pfromm, P. H.; Peterson, A. A. Chemical Looping of Metal Nitride Catalysts: Low-Pressure Ammonia Synthesis for Energy Storage. *Chem. Sci.* **2015**, *6*, 3965–3974.

(28) Abghoui, Y.; Skúlason, E. Electrochemical Synthesis of Ammonia via Mars–van Krevelen Mechanism on the (111) Facets of Group III–VII Transition Metal Mononitrides. *Catal. Today* **2017**, *286*, 78–84.

(29) Rittle, J.; Peters, J. C. An Fe–N₂ Complex That Generates Hydrazine and Ammonia via Fe=NNH₂: Demonstrating a Hybrid Distal-to-Alternating Pathway for N₂ Reduction. *J. Am. Chem. Soc.* **2016**, *138*, 4243–4248.

(30) Del Castillo, T. J.; Thompson, N. B.; Peters, J. C. A Synthetic Single-Site Fe Nitrogenase: High Turnover, Freeze-Quench ⁵⁷Fe Mössbauer Data, and a Hydride Resting State. *J. Am. Chem. Soc.* **2016**, *138*, 5341–5350.