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Catalytic Reduction of Dinitrogen to Ammonia at a Single Molybdenum Center

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ABSTRACT

This Account explores the catalytic reduction of dinitrogen by molybdenum complexes that contain the [HIPTN₃N]³⁻ ligand ([HIPTN₃N]³⁻ = [(HIPTNCH₂CH₂)₃N]³⁻, where HIPT = 3,5-(2,4,6-i-Pr₃C₆H₂)₂C₆H₃) at room temperature and pressure with protons and electrons. A total of 7–8 equiv of ammonia is formed out of ~12 possible (depending upon the Mo derivative employed). No hydrazine is formed. Numerous X-ray studies of proposed intermediates in the catalytic cycle suggest that N₂ is being reduced at a sterically protected, single Mo center operating in oxidation states between Mo^{III} and Mo^{VI}. Subtle variations of the [HIPTN₃N]³⁻ ligand are not as successful as a consequence of an unknown shunt in the catalytic cycle that consumes reduction equivalents to yield (it is proposed) dihydrogen.

Introduction

Approximately 10⁸ tons per year of ammonia are formed in the environment from dinitrogen, an amount equivalent to that formed in the Haber–Bosch process (at 350–550 °C and 150–350 atm).¹ It was first recognized in the 1960s that fixation of dinitrogen in the environment is carried out in certain bacteria by a metalloenzyme, an FeMo nitrogenase.^{2–5} “Alternative” nitrogenases are now known, one that contains vanadium instead of molybdenum (which functions when Mo levels are low and V is available) and another that contains only iron (which functions when both Mo and V levels are low).^{6–8} The FeMo nitrogenase has been purified and studied for decades. It also has been crystallized and subjected to X-ray studies that have elicited a great deal of discussion concerning the mechanism of dinitrogen reduction.^{9–12} However, it is fair to say that, despite a huge effort to

understand how dinitrogen is reduced by various nitrogenases over a period of more than 40 years, no definitive conclusions concerning the site and mechanism of dinitrogen reduction in nitrogenase(s) have been reached.¹³

Transition-metal dinitrogen chemistry began in 1965 with the discovery of [Ru(NH₃)₅(N₂)]²⁺, which was prepared by treating Ru salts with hydrazine.¹⁴ As more dinitrogen complexes were uncovered, it seemed possible that a transition-metal complex might be able to catalyze the reduction of dinitrogen to ammonia under mild conditions in solution with protons and electrons or the combination of dinitrogen with other elements in a selective manner under mild conditions.^{15–21} Hundreds of man years have been invested in these goals. Although dinitrogen at 1 atm combines readily with lithium at room temperature, the development of *catalytic* reactions under mild conditions that involve dinitrogen has been a formidable problem. Reduction of dinitrogen to ammonia in solution with protons and electrons carries with it the fact that protons are reduced readily to dihydrogen. Therefore, reduction of one of the most stable molecules known in a true homogeneous nonenzymatic catalytic reaction with protons and electrons has been extraordinarily challenging.

The principles of reduction of dinitrogen to ammonia at a single metal center by Mo(0) and W(0) dinitrogen complexes were established in studies that began in the late 1960's, primarily by the groups directed by Chatt and Hidai. Over a period of years, it was demonstrated that dinitrogen could be bound and reduced to ammonia through a series of approximately a dozen intermediates, the most important of them being M(N₂), M–N=NH, M=N–NH₂, M≡N, M=NH, M–NH₂, and M(NH₃) species. Although examples of virtually all of the proposed intermediates for reduction of dinitrogen at a single Mo or W center in a “Chatt cycle” were isolated, no catalytic reduction of dinitrogen to ammonia was ever achieved.

One reaction is known in which dinitrogen is reduced to a mixture of hydrazine and ammonia (~10:1).²² Molybdenum is required, and dinitrogen reduction is catalytic with respect to molybdenum. The reaction is run in methanol in the presence of magnesium hydroxide and a strong reducing agent such as sodium amalgam. Few details concerning the mechanism of this reaction have been established.

In the last 10 years, we have been exploring early transition-metal complexes that contain a triamidoamine ligand ([RNCH₂CH₂)₃N]³⁻.²³ Recently, we were attracted to the chemistry of Mo complexes that contain the [HIPTN₃N]³⁻ ligand, where HIPT = 3,5-(2,4,6-i-Pr₃C₆H₂)₂-C₆H₃ (hexaisopropylterphenyl; see Figure 1).^{24,25} This ligand was designed to prevent formation of relatively stable and unreactive [ArN₃N]Mo–N=N–Mo[ArN₃N] complexes, maximize steric protection of a metal coordination site in a monometallic species, and provide increased

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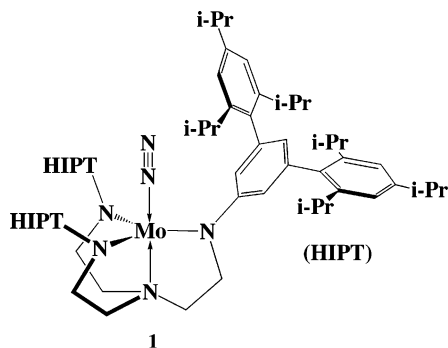


FIGURE 1. Drawing of $[\text{HIPTN}_3\text{N}]\text{Mo}(\text{N}_2) = \text{Mo}(\text{N}_2)$.

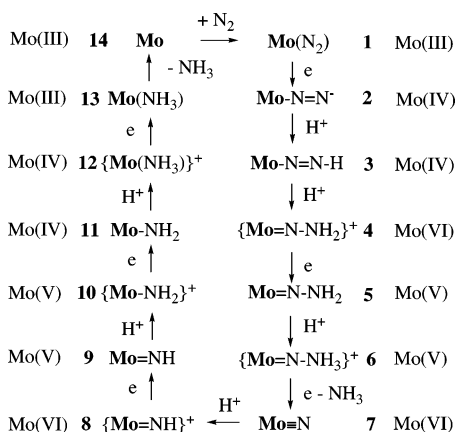


FIGURE 2. Proposed intermediates in the reduction of dinitrogen at a $[\text{HIPTN}_3\text{N}]\text{Mo}$ (**Mo**) center through the stepwise addition of protons and electrons.

solubility in nonpolar solvents. Eight of the proposed intermediates in a hypothetical “Chatt-like” reduction of dinitrogen (Figure 2) were prepared and characterized.^{24–27} These include paramagnetic $\text{Mo}(\text{N}_2)$ (**1**, Figure 1), diamagnetic $[\text{Mo}(\text{N}_2)]^-$ (**2**), diamagnetic $\text{Mo}-\text{N}=\text{N}-\text{H}$ (**3**), diamagnetic $[\text{Mo}=\text{N}-\text{NH}_2]\text{BAR}'_4$ (**4**; $\text{Ar}' = 3,5-(\text{CF}_3)_2\text{C}_6\text{H}_3$), diamagnetic $\text{Mo}\equiv\text{N}$ (**7**), diamagnetic $[\text{Mo}=\text{NH}]\text{BAR}'_4$ (**8**), paramagnetic $[\text{Mo}(\text{NH}_3)]\text{BAR}'_4$ (**12**), and paramagnetic $\text{Mo}(\text{NH}_3)$ (**13**). With the exception of **7**, all are extremely sensitive to oxygen. Several of these species then were employed successfully to reduce dinitrogen catalytically to ammonia with protons and electrons,²⁶ and we have been able to study several steps in the catalytic cycle. Although much remains to be done, we now know something about how this remarkable reaction proceeds. The purpose of this Account is to summarize the major findings in this area since the $[\text{HIPTN}_3\text{N}]\text{Mo}$ species were first reported.²⁴

Some Structural Characteristics of $[\text{HIPTN}_3\text{N}]\text{Mo}$ Complexes

Trigonal bipyramidal molybdenum complexes that contain the $[\text{HIPTN}_3\text{N}]^{3-}$ ligand all contain an approximately trigonal pocket in which N_2 or its partially or wholly reduced products are protected to a dramatic degree by three HIPT groups clustered around it. An example is the structure of $\text{Mo}(\text{NH}_3)$ shown in Figure 3. The basic characteristics of binding of the three amides and the amine in $[\text{HIPTN}_3\text{N}]\text{Mo}$ complexes do not vary dramati-

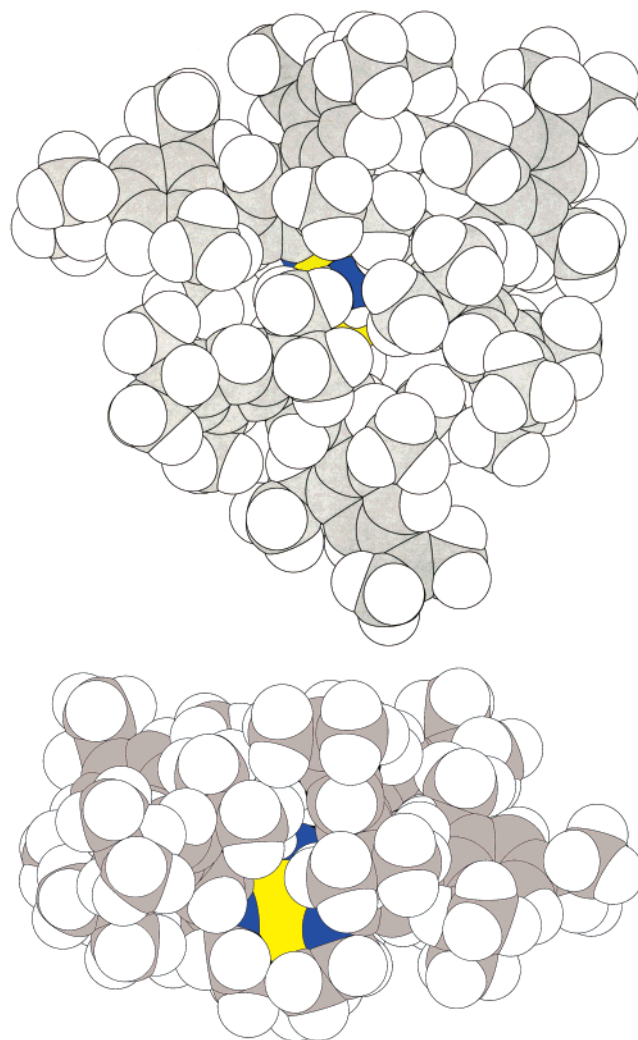
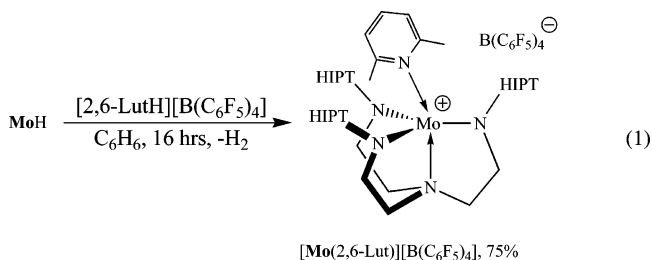


FIGURE 3. Two space-filling views of the structure of $[\text{HIPTN}_3\text{N}]\text{Mo}(\text{NH}_3)$ from the top (above) and from the side (below; Mo is yellow, and N is dark blue).

cally from complex to complex. As a consequence of relatively free rotation of the HIPT substituents about the $\text{N}-\text{C}_{\text{ipso}}$ bond, they adopt a wide variety of orientations in response to what are usually subtle, but sometimes not so subtle, steric demands of the ligand or ligands bound in the trigonal pocket. A dramatic example of the degree to which the HIPT substituents can respond to the steric demands imposed by a ligand bound in the trigonal pocket is $[\text{Mo}(\text{2,6-dimethylpyridine})]^+$ (eq 1).²⁷



The 2,6-lutidine is bound “off-axis” to a significant degree ($\text{N}_{\text{amine}}-\text{Mo}-\text{N}_{\text{lut}} = 157^\circ$) in a “slot” created by two of the HIPT groups. Because no asymmetry is present on

the NMR time scale at room temperature in solution, the HIPT groups, as well as the ligand or ligands in the trigonal pocket, must be rotating and rearranging at a rate near or faster than the NMR time scale.

The orbital to which the lutidine is bound in $[\text{Mo}(2,6\text{-dimethylpyridine})]^+$ can be described as a σ -type hybrid formed from the “ d_z ”, d_{xz} , and d_{yz} orbitals, with the 3-fold axis being the z axis. In general, these three orbitals can be employed to form 3σ , 1π and 2σ , or 2π and 1σ bonds to a ligand or ligands bound in the trigonal pocket.²³ Six-coordinate species are known in TMS or C_6F_5 triamidoamine Mo complexes when strongly binding ligands are present (CO or isonitriles),²⁸ although six-coordinate species have not yet been observed in $[\text{HIPTN}_3\text{N}]\text{MoL}_x$ systems that do not contain CO or RNC ligands.

Catalytic Reduction of Dinitrogen

Dinitrogen is reduced catalytically in heptane with [2,6-lutidinium] BAR'_4 [$\text{Ar}' = 3,5\text{-}(\text{CF}_3)_2\text{C}_6\text{H}_3$] as the proton source and decamethylchromocene as the electron source in the presence of several of the isolated complexes shown in Figure 2.²⁶ We chose 2,6-lutidinium because it is a relatively weak acid ($\text{p}K_a$ in water = 6.75) and because we felt that the 2,6-lutidine that is formed after delivery of the proton is unlikely to bind strongly to a neutral (Mo^{III}) center. The BAR'_4^- anion was chosen because it is large, unlikely to interact with the metal in $\{[\text{HIPTN}_3\text{N}]\text{Mo}\}^+$ complexes, and likely to lead to salts that are soluble in relatively nonpolar solvents. Decamethylchromocene was chosen as the electron source because it was found to reduce $[\text{Mo}(\text{NH}_3)]\text{BAR}'_4$ to $\text{Mo}(\text{NH}_3)$ in benzene. (E° for $\text{Mo}(\text{NH}_3)^{+/0} = -1.51$ V relative to $\text{FeCp}_2^{+/0}$, while E° for $\text{CrCp}^*_{2,0/+} = -1.47$ V in THF.²⁷) However, $\text{CrCp}^*_{2,0/+}$ cannot fully reduce $\text{Mo}(\text{N}_2)$ to $[\text{Mo}(\text{N}_2)]^-$ (E° for $[\text{Mo}(\text{N}_2)]^{0/-}$ is -1.81 V in THF versus $\text{FeCp}_2^{+/0}$).²⁷ Heptane was chosen as the solvent to minimize the solubility of [2,6-lutidinium] BAR'_4 and thereby minimize direct reduction of protons by $\text{CrCp}^*_{2,0/+}$ in solution. Slow addition of the reducing agent in heptane to an Mo complex and [2,6-lutidinium] BAR'_4 in heptane (over a period of 6 h with a syringe pump) was chosen to minimize exposure of protons to $\text{CrCp}^*_{2,0/+}$ at a high concentration.

A growing number of catalytic runs by several researchers with several different Mo derivatives (usually **1**, **3**, **7**, or **12**) reveal that a total of 7–8 equivalents of ammonia are formed out of ~12 possible (depending upon the Mo derivative employed), which suggests an efficiency of ~65% based on the reducing equivalents available. (The efficiency of formation of ammonia from the gaseous dinitrogen present is 55–60%.) A run employing $\text{Mo}-^{15}\text{N}=\text{NH}$ under $^{15}\text{N}_2$ yielded entirely ^{15}N -labeled ammonia. We now know that no hydrazine is formed²⁹ but still have not established that any reducing agent that is not consumed in making ammonia is consumed to form dihydrogen. It is clear, however, that [2,6-lutidinium] BAR'_4 does react rapidly with $\text{CrCp}^*_{2,0/+}$ in benzene to yield $[\text{CrCp}^*_{2,0/+}]\text{BAR}'_4$ and dihydrogen. CoCp_2 , which is a weaker reducing agent than $\text{CrCp}^*_{2,0/+}$ (the $\text{CoCp}_2^{0/+}$ couple is -1.33 V versus

$\text{FeCp}_2^{+/0}$ in either THF or PhF), also can be employed as the reducing agent for catalytic dinitrogen reduction, although it is approximately half as efficient as $\text{CrCp}^*_{2,0/+}$.²⁷

Parent Diazenido Complex, $\text{Mo}-\text{N}=\text{NH}$

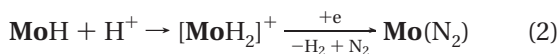
$\text{Mo}-\text{N}=\text{NH}$ (**3** in Figure 2) has been prepared by protonation of $\text{Mo}-\text{N}=\text{N}^-$ with $[\text{Et}_3\text{NH}]\text{BAR}'_4$ and has been characterized structurally.^{25,27} Although the proton on the β nitrogen atom could not be located in the X-ray study, ^{15}N and proton NMR studies have established conclusively that a proton is present on N_β . Although there are many well-characterized examples in the literature of $\text{M}-\text{N}=\text{NR}$ complexes where $\text{R} \neq \text{H}$, *parent* diazenido ($\text{M}-\text{N}=\text{NH}$) complexes are extremely rare.^{30,31} To our knowledge, no other terminal $\text{M}-\text{N}=\text{NH}$ species has been characterized structurally or via NMR studies of ^{15}N -labeled samples, with the exception of the W analogue of $\text{Mo}-\text{N}=\text{NH}$ (see later). Upon heating in benzene, $\text{Mo}-\text{N}=\text{NH}$ slowly decomposes to MoH and dinitrogen in a first-order process ($k = 2.2 \times 10^{-6} \text{ s}^{-1}$, $t_{1/2} = 90 \text{ h}$ at 61°C).²⁵ Decomposition of $\text{Mo}-\text{N}=\text{NH}$ is accelerated an order of magnitude in the presence of 1% $[\text{Et}_3\text{NH}][\text{OTf}]$ or $[\text{Et}_3\text{NH}]\text{BAR}'_4$; mechanistic details and whether the rate of decomposition is further accelerated in the presence of additional $[\text{Et}_3\text{NH}]\text{BAR}'_4$ or $[2,6\text{-LutH}]\text{BAR}'_4$ are not yet available. Decomposition of $\text{Mo}-\text{N}=\text{NH}$ is a problem even at room temperature over the long term, a circumstance that complicated obtaining crystals suitable for an X-ray study.²⁷ In the presence of $[\text{H}(\text{Et}_2\text{O})_2]\text{BAR}'_4$, $\text{Mo}-\text{N}=\text{NH}$ is protonated at N_β to yield $[\text{Mo}=\text{NNH}_2]^+$, but $[2,6\text{-LutH}]\text{BAR}'_4$ only partially and reversibly protonates $\text{Mo}-\text{N}=\text{NH}$.

It has been established that $\text{Mo}-\text{N}=\text{NH}$ is formed rapidly upon treatment of $\text{Mo}(\text{N}_2)$ with CoCp_2 and [2,6-lutidinium] BAR'_4 in benzene, even though neither reacts alone with $\text{Mo}(\text{N}_2)$. CoCp_2 is too weak of a reducing agent to reduce $\text{Mo}(\text{N}_2)$ fully to $[\text{Mo}-\text{N}=\text{N}]^-$, but it is difficult to exclude the possibility that $\text{Mo}-\text{N}=\text{NH}$ is formed through a rapid and reversible reduction of $\text{Mo}(\text{N}_2)$ to give a minute amount of $[\text{Mo}-\text{N}=\text{N}]^-$ in equilibrium with $\text{Mo}(\text{N}_2)$, followed by a rapid protonation of $[\text{Mo}-\text{N}=\text{N}]^-$. Interestingly, the addition of [2,6-lutidinium] BAR'_4 to $\text{Mo}(\text{N}_2)$ has been shown to yield (reversibly) ~30% of a species in fluorobenzene in which ν_{NN} has increased from 1990 to 2056 cm^{-1} , consistent with protonation of $\text{Mo}(\text{N}_2)$ to yield a cationic species. However, *dinitrogen* cannot be the site of protonation. Two possibilities are the metal center or an amido nitrogen. In any cationic species, the potential for addition of an electron should be shifted considerably in the positive direction by as much as 500 or 600 mV.³² In short, the reductive protonation of $\text{Mo}(\text{N}_2)$ to $\text{Mo}-\text{N}=\text{NH}$ could be *proton-catalyzed*.

The Role of MoH

The hydride MoH is most readily prepared by treating $[\text{Mo}(\text{NH}_3)]\text{BPh}_4$ with LiBHET_3 .²⁷ As mentioned above, it is also the product of decomposition of $\text{Mo}-\text{N}=\text{NH}$. MoH may be formed under catalytic conditions, although that has not yet been established.

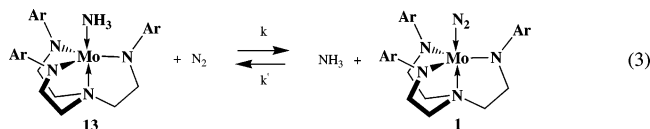
It turns out that **MoH** is as efficient of a catalyst precursor in dinitrogen reduction as any species that we have employed. One can readily imagine how this might happen, as shown in eq 2. In short, **MoH** is a hydrogenase. However, the rate of formation of dihydrogen in the manner shown in eq 2



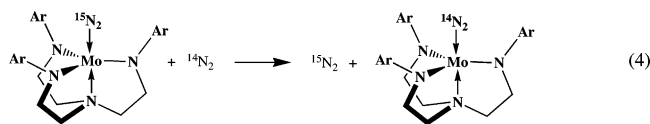
versus the rates of all other reactions in which dihydrogen may be formed under catalytic conditions, including “direct” reduction by electron transfer, is not yet known. It should be noted that formation of 1 equiv of dihydrogen when the FeMo nitrogenase reduces dinitrogen has been proposed to be an integral aspect of the reduction process itself, i.e., not a competitive product that may or may not be formed at the same site at which dinitrogen is reduced. More than one dihydrogen is formed per dinitrogen reduced in alternative nitrogenase systems.

Formation of **Mo(NH₃)** and Its Conversion to **Mo(N₂)**

It has been established that **[Mo(NH₃)]BAR'₄** is formed upon treatment of **Mo(N₂)** in benzene with sufficient **CoCp₂** and **[2,6-lutidinium]BAR'₄** and that a stronger reducing agent such as **CrCp*₂** will reduce **[Mo(NH₃)]BAR'₄** to **Mo(NH₃)** in benzene. A catalytic cycle is then “completed” when **Mo(NH₃)** (**13**) is converted into **Mo(N₂)** (**1**, eq 3).



The equilibrium constant for conversion of **Mo(NH₃)** in the presence of dinitrogen into a mixture of **Mo(N₂)**, **Mo(NH₃)**, and ammonia has been measured in **C₆D₆** at 50 °C and found to be 1.2.²⁷ It also has been shown that an equilibrium ($K_{\text{eq}} \approx 0.1$) between **Mo(NH₃)**, dinitrogen, **Mo(N₂)**, and ammonia is established in 1–2 h under an atmosphere of ammonia (0.28 atm, ~21 equiv versus Mo) and dinitrogen.²⁹ A reaction in which **Mo(¹⁵N₂)** is converted into **Mo(¹⁴N₂)** under 1 atm of ¹⁴N₂ (a large excess versus Mo; eq 4)



has a half-life of ~35 h at 15 psi (1 atm),²⁵ 32 h at 30 psi,³³ and 30 h at 55 psi,³³ which suggests that the exchange of ¹⁵N₂ for ¹⁴N₂ is independent of dinitrogen pressure. Therefore, *dinitrogen* exchange (eq 4) is *dissociative*, with the “naked” species (**14**, Figure 2) the likely intermediate. Because the equilibrium in eq 3 is established in only 1–2 h, conversion of **1** to **13** (and by definition, **13** to **1**) does *not* involve **14** as an intermediate. All data in hand thus far (see below) suggest that this is the case.

The rate of conversion of **Mo(NH₃)** into **Mo(N₂)** under dinitrogen in the absence of **[2,6-lutidinium]BAR'₄** has been studied in a preliminary and semiquantitative fashion by IR spectroscopy and differential pulse voltammetry.^{29,33} The conversion of **Mo(NH₃)** into **Mo(N₂)** in the absence of added ammonia is not a simple reaction, because the ammonia that is formed in solution not only back reacts with **Mo(N₂)** to give **Mo(NH₃)** but also begins to enter the headspace above solvents such as benzene or heptane. (Preliminary measurements suggest that in **C₆D₆** at 22 °C approximately the same amount of ammonia is present in a given volume of **C₆D₆** as in 3 times that volume of headspace at equilibrium.) In heptane, the conversion of **Mo(NH₃)** into **Mo(N₂)** follows what is close to first-order kinetics in Mo with a half-life for the conversion at 22 °C and 1 atm of ~115 min. (The plot of $\ln(1 - \text{Mo}/\text{Mo}_\infty)$ versus time is slightly curved, as a consequence of the back reaction between ammonia in solution and **Mo(N₂)** to reform **Mo(NH₃)**; therefore, these measured rates of conversion are semiquantitative.) At 2 atm, the half-life is ~45 min, which suggests that the conversion depends upon the concentration of dinitrogen to the first power.^{29,33} Conversion of **Mo(NH₃)** into **Mo(N₂)** is dramatically accelerated at 1 atm in the presence of 4 equiv of **BPh₃** ($t_{1/2}$ is ~35 min), and the $\ln(1 - \text{Mo}/\text{Mo}_\infty)$ plot is strictly linear, most likely because **BPh₃** removes much or all of the ammonia formed in solution relatively efficiently.^{29,33} Much remains to be done to confirm that the interconversions of **Mo(NH₃)** and **Mo(N₂)** (eq 3), unlike interconversions of **Mo(¹⁵N₂)** and **Mo(¹⁴N₂)** (eq 4), are *S_N2* reactions. Therefore, an inescapable consequence is that ammonia formed through reduction of dinitrogen will be an inhibitor of further dinitrogen reduction through displacement of the equilibrium shown in eq 3 to the left.

Variations of Standard Conditions

Preliminary results suggest that the amount of ammonia formed from dinitrogen at 2 atm under standard conditions is ~25% greater than at 1 atm, as one would expect on the basis of the nitrogen pressure dependence of the reaction shown in eq 3. If the reducing agent is added over a period of <1 min, then the amount of dinitrogen that is converted into ammonia is <10%; presumably electrons are consumed primarily to form dihydrogen. Because conversion of **Mo(NH₃)** into **Mo(N₂)** depends upon pressure, proton reduction in theory could be minimized at high dinitrogen pressures. An interesting question is whether the efficiency of conversion of dinitrogen into ammonia can be pushed beyond 75% (1 equiv of dihydrogen per dinitrogen reduced, as found in FeMo nitrogenase) under some conditions, e.g., at several atmospheres *N₂* pressure.

Dinitrogen reduction is inhibited by 2,6-lutidine, which is formed after lutidinium delivers a proton. For example, the amount of ammonia formed from dinitrogen decreases from 3.4 in a run employing 18 equiv of reducing agent to 0.5 in a run in which 145 equiv of 2,6-lutidine had been added.²⁹ There are several possible explanations

for this effect, among them is the possibility that excess 2,6-lutidine simply makes 2,6-lutidinium a poorer acid.

Not all [pyridinium]BAR'₄ acids work well. No ammonia is formed from dinitrogen if 2,6-diphenylpyridinium or 3,5-dimethylpyridinium BAR'₄ sources are employed, while 4.1 equiv are formed from dinitrogen when [2,4-dimethylpyridinium]BAR'₄ is employed under standard conditions (36 equiv of reducing agent) and 2.7 equiv are formed when [2,6-diethylpyridinium]BAR'₄ is employed.²⁹ Not only must the solubility of the acid source, its p*K*_a (in water), and the ability of the conjugate base to bind to **Mo** be considered in explaining these results, but the bulk of the acid source as an ion pair must also be considered. Two or more of these factors change when the acid source is changed; therefore, it has been challenging to pinpoint the origin of any [pyridinium]⁺ effect.

Alternatives to the [HIPTN₃N]³⁻ Ligand

Three “symmetric” variations of the [HIPTN₃N]³⁻ ligand have been explored.³⁴ One is a hexa-*tert*-butylterphenyl-substituted ([HTBTN₃N]³⁻) ligand; a second is a hexamethylterphenyl-substituted ([HMTN₃N]³⁻) ligand; and a third is a [*p*BrHIPTN₃N]³⁻ ligand, which is a [HIPTN₃N]³⁻ ligand in which the para position of the central phenyl ring is substituted with a bromide. IR and electrochemical studies suggest that complexes that contain the [HTBTN₃N]³⁻ ligand are slightly more electron-rich than those that contain the parent [HIPTN₃N]³⁻ ligand, while those that contain the [*p*BrHIPTN₃N]³⁻ ligand are slightly more electron-poor. One would expect [HTBTN₃N]Mo complexes to be significantly more crowded sterically than [HIPTN₃N]Mo complexes (an X-ray study of [HTBTN₃N]MoCl shows this to be the case), [HMTN₃N]Mo complexes to be significantly less crowded than [HIPTN₃N]Mo complexes, and [*p*BrHIPTN₃N]Mo complexes to have approximately the same steric crowding as [HIPTN₃N]Mo complexes. In practice, transformations of [HTBTN₃N]³⁻ derivatives that involve electron and proton transfer (e.g., conversion of [HTBTN₃N]Mo(N₂) into [HTBTN₃N]Mo–N≡NH) are slower by perhaps an order of magnitude compared to analogous conversions of [HIPTN₃N]³⁻ derivatives, consistent with a high degree of steric crowding in [HTBTN₃N]³⁻ derivatives.

[*p*BrHIPTN₃N]Mo≡N was found to be a catalyst for the formation of ammonia in yields only slightly less than those observed employing [HIPTN₃N]³⁻ derivatives (~65%). Generation of [*p*BrHIPTN₃N]Mo(NH₃) and observation of its conversion to [*p*BrHIPTN₃N]Mo(N₂) under 1 atm of dinitrogen in heptane at 22 °C showed that the half-life for formation of [*p*BrHIPTN₃N]Mo(N₂) is ~2 h, the same as for the conversion of [HIPTN₃N]Mo(NH₃) into [HIPTN₃N]Mo(N₂).

[HTBTN₃N]Mo≡N was found to be a poor catalyst for the reduction of dinitrogen, with only 1.06 equiv of ammonia being observed. Therefore, only 0.06 equiv are formed from gaseous dinitrogen. One measurement of the rate of conversion of [HTBTN₃N]Mo(NH₃) into [HTBTN₃N]Mo(N₂) under conditions analogous to those employed

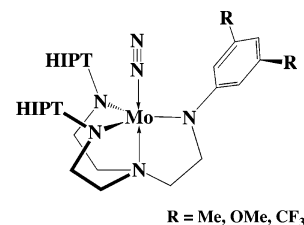


FIGURE 4. Drawing of a [hybrid]Mo(N₂) complex.

for **Mo**(NH₃) reveals that the “half-life” for conversion is approximately 30 h instead of 2 h, as it is for conversion of **Mo**(NH₃) to **Mo**(N₂).²⁹ The observable (slower) coupled reduction of {[HTBTN₃N]Mo(NH₃)}⁺ to [HTBTN₃N]Mo(NH₃) and (slower) conversion of [HTBTN₃N]Mo(NH₃) to [HTBTN₃N]Mo(N₂) apparently cannot compete with the formation of dihydrogen via “direct” reduction of protons.

Use of [HMTN₃N]Mo≡N as a catalyst under standard conditions was also relatively unsuccessful; only 0.47 equiv of ammonia were formed from dinitrogen. Low solubility of [HMTN₃N]Mo≡N and probably other intermediates (in heptane) is one possible problem. However, studies of other “hybrid” alternatives to the [HTBTN₃N]³⁻ ligand (*vide infra*) suggest that there may also be other problems.

Changing three substituents on the ligand at once is a relatively drastic variation. Therefore, we were pleased to find that we could change the substituent on only *one* of the arms, i.e., we could prepare “hybrid” ligands of the type [(HIPTNCH₂CH₂)₂NCH₂CH₂NAr]³⁻, where Ar is 3,5-dimethylphenyl, 3,5-dimethoxyphenyl, or 3,5-bistrifluoromethylphenyl (Figure 4).²⁹ When [(HIPTNCH₂CH₂)₂NCH₂CH₂NAr]Mo≡N species are employed in a standard attempted catalytic reaction, no ammonia is produced from dinitrogen using any of the three Ar variations. The half-life for conversion of [(HIPTNCH₂CH₂)₂NCH₂CH₂NAr]Mo(NH₃) into [(HIPTNCH₂CH₂)₂NCH₂CH₂NAr]Mo(N₂) was shown to be ~170 min when Ar = 3,5-(CF₃)₂C₆H₃ and 42 min when Ar = 3,5-(OMe)₂C₆H₃.²⁹ Because the half-life for conversion of **Mo**(NH₃) into **Mo**(N₂) is ~120 min, reduction must fail in the “hybrid” cases for some reason other than a slow conversion of [hybrid]Mo(NH₃) into [hybrid]Mo(N₂).

The reason that [hybrid]Mo catalysts will not reduce dinitrogen is (we propose) a “shunt” in the catalytic cycle that consumes protons and electrons to yield hydrogen relatively rapidly. Because the three Ar groups are similar sterically, we suspect that the reaction that produces hydrogen is simply accelerated in the less crowded circumstance. Of the several possibilities at this stage, perhaps the simplest is hydrogen formation at the metal center, i.e., hydrogenase activity. In any case, it is extraordinary that dinitrogen reduction fails when only one of the three amido nitrogen substituents is a slightly smaller 3,5-disubstituted phenyl than a HIPT group.

Other Metals

Many tungsten analogues of the species shown in Figure 2 have been synthesized, and several have been structurally characterized, among them are WN≡NK, W(N₂),

$[\mathbf{W}(\text{N}_2)]^-$, $\mathbf{W}\text{N}=\text{NH}$, $[\mathbf{W}=\text{NNH}_2]\text{BAR}'_4$, $\mathbf{W}\equiv\text{N}$, and $[\mathbf{W}(\text{NH}_3)]\text{-BAR}'_4$.³⁵ The structural, spectroscopic, and electrochemical trends are those one might expect. For example, ν_{NN} for $\mathbf{W}(\text{N}_2)$ is found at 1888 cm^{-1} compared to 1990 cm^{-1} for $\mathbf{Mo}(\text{N}_2)$, consistent with \mathbf{W}^{III} being a more powerful reductant of dinitrogen through π back-donation. However, attempted catalytic reduction of dinitrogen using $\mathbf{W}(\text{N}_2)$ as the catalyst under conditions identical or similar to those employed for catalytic reduction of dinitrogen by $\mathbf{Mo}(\text{N}_2)$ and related Mo complexes yielded less than 2 equiv of ammonia; i.e., only the dinitrogen initially bound to tungsten is reduced (partially) to ammonia, and no atmospheric dinitrogen is reduced. At most, 1.51 equiv of ammonia was produced with CoCp^*_2 as the reductant and $[2,6\text{-LutH}]\text{BAR}'_4$ as the acid.

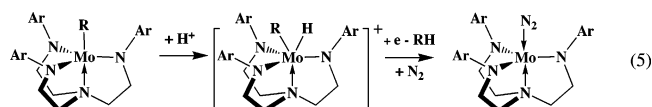
One compound that is conspicuously absent in the list of isolated or observed complexes above is $\mathbf{W}(\text{NH}_3)$. All efforts thus far to observe this compound or to observe $\mathbf{W}(\text{N}_2)$ as a product of reduction of $[\mathbf{W}(\text{NH}_3)]\text{BAR}'_4$ under dinitrogen have failed. Therefore, we suspect that one of the reasons that turnover is not achieved is the failure to generate $\mathbf{W}(\text{NH}_3)$ and exchange ammonia for dinitrogen to yield $\mathbf{W}(\text{N}_2)$. At present, we could only speculate why this is the case. Another problem is that it is more difficult to reduce tungsten analogues of molybdenum compounds. For example, the reduction of $\mathbf{W}(\text{N}_2)$ to $[\mathbf{W}(\text{N}_2)]^-$ is reversible at -2.27 V versus $\text{FeCp}_2^{+/0}$ in 0.1 M $[\text{Bu}_4\text{N}]\text{-BAR}'_4/\text{PhF}$ electrolyte, which is 0.26 V more negative than the $\mathbf{Mo}(\text{N}_2)^{0/-}$ potential under the same conditions. When taken together, and given the already extremely finely balanced nature of the Mo-catalyzed reduction of dinitrogen, it is perhaps not surprising that dinitrogen reduction fails using $\mathbf{W}(\text{N}_2)$ under conditions where reduction using $\mathbf{Mo}(\text{N}_2)$ is successful. It is interesting to note that a tungsten-substituted nitrogenase can be isolated from *Rhodobacter capsulatus* and that it is inactive for reduction of acetylene and dinitrogen, although it is relatively active for reduction of protons.³⁶

Because TMS-substituted or C_6F_5 -substituted triamidoamine complexes that contain vanadium are known,^{37,38} we were not surprised to find that $[\text{HIPTN}_3\text{N}]\text{V}(\text{THF})$ can be prepared readily.³⁹ An attempt to reduce dinitrogen under the standard set of conditions that have been successful for Mo compounds produced only traces of ammonia when $[\text{HIPTN}_3\text{N}]\text{V}(\text{THF})$ was employed. It should be noted that the charges on any complex would be off by one; e.g., if the V(V) nitride were formed, it would have to be an anion. Ideally, therefore, a dianionic ligand might be more desirable, e.g., one in which one of the anionic amido “arms” is instead a neutral donor. However, the “donor arm” in ligands of that type that we have prepared thus far appears to dissociate and allow bimetallic species to form.³⁹ Therefore, this problem is unsolved.

Organometallic Chemistry

As one might expect on the basis of work with other types of triamidoamine Mo complexes,^{23,28,40,41} it is possible to prepare organometallic derivatives such as $\mathbf{Mo}(\eta^2\text{-C}_2\text{H}_4)$,

$\mathbf{Mo}(\eta^2\text{-C}_2\text{H}_2)$, $\mathbf{Mo}(\text{CO})$, $\mathbf{Mo}(\text{CN})$, $\mathbf{Mo}(\text{CH}_2\text{R})$ (R = methyl, pentyl, or heptyl), $\mathbf{Mo}(\eta^2\text{-CHCH}_2)$, $\mathbf{Mo}\equiv\text{CR}$, $[\mathbf{Mo}(\text{C}_2\text{H}_4)]^+$, and $[\mathbf{Mo}(\text{C}_2\text{H}_2)]^+$.⁴² Thus far, we know that $\mathbf{Mo}(\eta^2\text{-C}_2\text{H}_4)$, $\mathbf{Mo}(\eta^2\text{-C}_2\text{H}_2)$, $\mathbf{Mo}(\text{hexyl})$, and $\mathbf{Mo}\equiv\text{CH}$ all yield ammonia when employed under standard conditions as “precatalysts” for the reduction of dinitrogen.⁴³ The most successful is $\mathbf{Mo}(\text{hexyl})$, which yields 6.3 equiv of ammonia from dinitrogen (53%; cf. 7.7 equiv from dinitrogen, 64%, with \mathbf{MoH} as the precursor). One can imagine that $\mathbf{Mo}(\text{hexyl})$ is protonated to yield hexane and $\mathbf{Mo}(\text{N}_2)$ in the presence of the reducing agent and dinitrogen (eq 5).



We have not yet attempted to reduce acetylene, a classic nitrogenase substrate, under the conditions that we have employed to reduce dinitrogen.

Mo^{III} versus Mo^{IV} + Ligand Preferences, and the Nature of $[\text{HIPTN}_3\text{N}]\text{Mo}$

Thus far, we have isolated $[\mathbf{Mo}(\text{L})]^+$ species in which L is ammonia, 2,6-lutidine, pyridine, diethyl ether, and THF. We know that chemical reduction of $[\mathbf{Mo}(2,6\text{-Lut})]^+$ with CrCp^*_2 under dinitrogen in C_6D_6 yields $\mathbf{Mo}(\text{N}_2)$ immediately, while CV studies suggest that reduction of $[\mathbf{Mo}(\text{THF})]^+$ yields $\mathbf{Mo}(\text{N}_2)$ in less than 2 s.³³ However, reduction of $[\mathbf{Mo}(\text{NH}_3)]^+$ yields $\mathbf{Mo}(\text{NH}_3)$, probably because ammonia is the strongest Brønsted base and the smallest of the σ donors in the list above. Therefore, one might expect the rate of displacement of ammonia by dinitrogen to be the slowest.

The “naked” species, \mathbf{Mo} , appears to be an intermediate in the $\text{S}_{\text{N}}1$ conversion of $^{14}\text{N}_2$ into $\mathbf{Mo}(^{15}\text{N}_2)$, but thus far, \mathbf{Mo} has not been implicated in any other reaction and all efforts to observe it have failed thus far. In contrast, high-spin $\text{Mo}[\text{N}(\text{t-Bu})\text{Ar}]_3$ (Ar = 3,5- $\text{Me}_2\text{C}_6\text{H}_3$) has been prepared and studied by Cummins et al.⁴⁴ $\text{Mo}[\text{N}(\text{t-Bu})\text{Ar}]_3$ reacts with dinitrogen to yield a $[\text{N}(\text{t-Bu})\text{Ar}]_3\text{Mo}(\mu\text{-N}_2)\text{Mo}[\text{N}(\text{t-Bu})\text{Ar}]_3$ species that then dissociates to yield 2 equiv of $[\text{N}(\text{t-Bu})\text{Ar}]_3\text{Mo}\equiv\text{N}$. The addition of a donor to a trisamido species, which is analogous to the circumstance in four-coordinate \mathbf{Mo} , would likely raise the energy of the d_{z^2} orbital above d_{xz} and d_{yz} and might be expected to lead to a relatively reactive ground-state low-spin species that is poised to react with dinitrogen efficiently.

Recent calculations have established a detailed energy profile for the proposed steps in the \mathbf{Mo} -catalyzed reduction of dinitrogen.⁴⁵ Interestingly, a free-energy input of $\sim 190\text{ kcal mol}^{-1}$ is calculated to be involved in the conversion of dinitrogen to 2 equiv of ammonia, which is the same as that obtained empirically for biological nitrogen fixation.

Summary

As far as we are aware, the $[\text{HIPTN}_3\text{N}]\text{Mo}$ system and some of its variations are the only nonenzymatic systems

that will reduce dinitrogen catalytically and selectively to ammonia in the presence of protons and electrons at room temperature and pressure. We now can be relatively certain that reduction is accomplished at a single Mo center in a manner that thus far appears to be analogous (approximately) to that proposed by Chatt and his group, with an important difference being that the highest possible oxidation states are involved (from Mo^{III} to Mo^{VI}) in the cycle. Conversion of Mo(NH₃) to Mo(N₂) appears to be one of the slow steps in the successful reductions, in part because ammonia is an inhibitor. Although the catalytic system that we have developed does not involve biologically relevant ligands, reduction of dinitrogen to ammonia in such a simple “artificial” Mo system suggests that Mo is exceptional, as Chatt et al. suspected, in its ability to reduce dinitrogen, although it is probably not unique. Whether dinitrogen is reduced at molybdenum in FeMo nitrogenase is still speculative, although it is attractive to propose that it is. It is possible that dinitrogen is reduced at V or Fe when they substitute for Mo. It may prove to be considerably more difficult to design nonenzymatic V or Fe systems that reduce dinitrogen catalytically under conditions similar to those used in the [HIPTN₃N]Mo reactions, primarily because V and Fe are less efficient than Mo. One also cannot state that dinitrogen must be reduced at low temperature and pressure only in the manner found in [HIPTN₃N]Mo systems. However, at the same time, one should ponder whether the sheer complexity of catalytic dinitrogen reduction to ammonia with protons and electrons automatically severely limits mechanistic alternatives under mild conditions. Finally, it is also interesting to speculate what other “high oxidation state” chemistry and what other catalytic reactions can be developed with complexes that contain the [HIPTN₃N]³⁻ ligand system.

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