Reversible Dihydrogen Activation and Hydride Transfer by a Uranium Nitride Complex

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Abstract: Cleavage of dihydrogen is an important step in the industrial and enzymatic transformation of N₂ into ammonia. The reversible cleavage of dihydrogen was achieved under mild conditions (room temperature and 1 atmosphere of H₂) by the molecular uranium nitride complex, \([\text{Cs[U(OSi-(O\text{Bu})_3)_2]_2}\text{(\mu-N)}]_1\) 1, leading to a rare hydride–imide bridged diuranium(IV) complex, \([\text{Cs[U(OSi(O\text{Bu})_3)_2]_2}\text{(\mu-H)}(\mu-NH)]_2\) 2 that slowly releases H₂ under vacuum. This complex is highly reactive and quickly transfers hydride to acetonitrile and carbon dioxide at room temperature, affording the ketimide- and formate-bridged U(IV) species \([\text{Cs[U(OSi-(O\text{Bu})_3)_2]_2}\text{(\mu-H)}(\mu-\text{CH}_3\text{CN})]_3\) 3 and \([\text{Cs[U(OSi-(O\text{Bu})_3)_2]_2}\text{(\mu-HCOO)}(\mu-\text{NHCOO})]_4\) 4.

Heterolytic H₂ cleavage is an important step in many reactions promoted by enzymes or synthetic catalysts.[1] Hydrogen activation is crucial in the industrial Haber–Bosch process that is used in the production of ammonia from N₂ and H₂, and is also important in nitrogen fixation by nitrogenase enzymes, which is enabled by hydride-containing intermediates.[2] Such hydride intermediates are also believed to play a role in the reduction of other substrates such as CO₂ or nitriles effected by these enzymes.[3] Moreover, hydrogenolysis by metal nitride intermediates is believed to be a key step in the Haber–Bosch process.[4] However, only three examples of heterolytic splitting of H₂ by a nitride complex have been reported so far.[5] The reactivity of uranium nitrides is of high current interest because of the relevance of U(IV) nitride (UN) in nuclear industry[6] and in the possibility of obtaining a highly reactive uranium hydride–imide complex.

Notably, despite the relevance of such species in biological and industrial dinitrogen reduction, their formation from H₂ addition to metal nitrides or metal–dinitrogen complexes remains extremely rare.[5a,11] Moreover, despite the fact that the seminal work of Andersen and Marks on organometallic uranium hydrides goes back to the early 80s,[12] uranium hydrides remain rare and most of the reported examples contain carbocyclic ligands.[12c,13] Herein we show that the bifunctional Lewis acid–base character of the U–N bond in the multimetallic U(IV) complex \([\text{Cs[U(OSi(O\text{Bu})_3)_2]_2}\text{(\mu-N)}]_1\) 1,[14] leads to the facile, reversible heterolytic cleavage of dihydrogen under ambient conditions and to the reversible formation of a parent imide–hydride complex that can transfer the hydride to acetonitrile and CO₂.

When a toluene solution of the previously reported[14] complex 1 is exposed to 1 atmosphere of hydrogen, the solution turns from brown to yellow within 15 minutes and the proton NMR spectrum shows the disappearance of the siloxides signal of 1 and the appearance of a new signal at 0.32 ppm, assigned to the siloxide ligands of complex \([\text{Cs[U-(O\text{Si}(O\text{Bu})_3)_2]}_2(\mu-H)(\mu-NH)]_2\) 2 (Scheme 1). The resonance at 571.3 ppm in the ¹H NMR spectrum of 2 is assigned to the U(IV)-bound hydride and its value is consistent with those reported for other U(IV) hydride complexes.[15] This resonance is absent in the ¹H NMR spectrum of the deuterated analogue \([\text{Cs[U(O\text{Si}(O\text{Bu})_3)_2]}_2(\mu-D)(\mu-\text{ND})]_2\), D₂-2. Removal of the gas in the headspace after reaction of 1 with H₂ results in slow H₂ loss and formation of the nitride complex 1. Crystals of complex 2 also lose H₂ once removed from the solution resulting in the isolation of complexes 1 and 2 co-crystallized in 30:70% ratio (see Supporting Information). However, loss of H₂ could be avoided by maintaining at −40°C the crystals of 2, obtained at −40°C under 1 atm H₂, during crystal mounting in oil and by performing X-ray data collection at −133°C. The facile and reversible activation of H₂ by complex 1 is remarkable. Notably, in the only example

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Scheme 1. Reversible cleavage of H₂ by 1 to afford the di-uranium(IV) hydride–imide product 2.
reported so far of cooperative H₂ activation by a bifunctional metal nitride, the H₂ cleavage is not reversible.[8a]

The molecular structure of 2 (Figure 1) could be determined by the X-ray analysis of crystals of 2, and it shows a dinuclear U⁴⁺ complex bridged by an imido and a hydrido ligand. Electron density that could be assigned to the imide and to the hydride was located, and both groups were refined without imposing any constraints. A Cs cation is bound by the siloxide ligands resulting in a disymmetric structure analogous to that found in the nitride complex 1. Notably, two siloxide ligands bridge the U1 and Cs centers binding the Cs in a bidentate fashion while only one siloxide ligand binding the Cs cation in a bidentate fashion bridges U2 and Cs. The Cs cation was found in the apical position of the nitride at a Cs–N distance of 2.73(6) Å and is located at a non-bonding distance of 4.821(3) Å from N1. Moreover, the U–N–U angle changes dramatically from linear in complex 1 (U–N–U angle: 170.2(3)°) to bent in the diamond core geometry of complex 2 (U–N–U angle: 117.5(3)°) and the U–N bond distances are significantly elongated in 2 (2.187(3), 2.256(4) Å) compared to those found in 1 (2.058(5)–2.079(5) Å). These data are in agreement with the presence of a protonated NH group and the U–NH distances are in the range of those found in diuranium(IV) imido bridged complexes (2.156(8)–2.315(8) Å).[36] One terminal parent imido U⁴⁺ complex is known,[27], but a bridging parent imido-U⁴⁺ linkage was not characterized previously. In contrast, the bridging parent imido U⁴⁺U⁴⁺ [K₂[U(OSi(OBu)₃)]₂(μ-NH)(μ-η²-C₃H₅N₂)] and U⁴⁺U⁴⁺ [K₂[U(OSi(OBu)₃)]₂(μ-NH)(μ-Cl)] complexes were previously isolated in our group by protonation of the nitride–hydrazide complex [K₂[U(OSi(OBu)₃)]₂(μ-N)(μ-η²-C₃H₅N₂)].[39] The hydride bridges the two uranium cations (U–U, 3.786 Å) in a non-symmetric fashion (U–H 2.18(6) and 2.36(6) Å) with the shorter distance being comparable to those determined by neutron diffraction studies for the [(C₆Me₆)₂UH₂] complex (2.134(9) Å) where the hydride bridges the two uranium centers in a symmetric fashion. Complex 2 is a rare example of imide–hydride bridged complex.[8a,11,19] Such species are very relevant as probable intermediate in N₂ hydrogenation to yield NH₃.[19b,20]

Complex 2 displays a rich reactivity that both unambiguously confirms the presence of the hydride–imide core in 2 and demonstrates the ability of 2 to effect hydride transfer reactions. Complex 2 reacts with acetonitrile and carbon dioxide to yield insertion products. The addition of MeCN to a toluene solution of 2 (Scheme 2a) led to the formation of the bridging imido–ketimide complex [Cs[U(OSi(OBu)₃)]₂(μ-NH)(μ-CH₂CHN)], 3, that was isolated in high yield (85%) as a yellow crystalline solid. Thus, the hydride addition to the acetonitrile results in the two-electron reduction of the nitrogen–carbon triple bond. The solid-state structure of 3 (Figure 2) shows the presence of two uranium ions bridged by an imido and a ketimido (or azavinylidene) group. A cesium...
cation binds in a dissymmetric fashion the imido nitrogen and the siloxide ligands. The carbon atom C121 has a trigonal planar geometry and the C121–N1 distance of 1.273(7) Å is consistent with a carbon–nitrogen double bond. The pseudo tetrahedral geometry of the N2 atom and the U–N2 distances (U1–N2 2.247(4) and U2–N2 2.209(5) Å) are in agreement with the presence of an imido group NH, indicating that the NH group is not involved in the reactivity with MeCN. The IR spectrum shows a peak at 1629 cm\(^{-1}\) assigned to the N–C double bond stretching. The insertion of nitriles into the U–C bond has been used by Kiplinger and co-workers\(^{[23]}\) as a convenient method for the synthesis of mono- and multi-metallic ketimide complexes. Formation of a ketimide ligand from the insertion of the nitrile triple bond into a uranium hydride complex was also reported previously but the insertion products were not crystallographically characterized\(^{[23]}\). The ability of 2 to perform hydride transfer reactivity and the consequent reducing behavior of the hydride towards substrates were confirmed in the reaction of 2 with carbon dioxide. Addition of two equivalents of CO\(_2\) (Scheme 2b) to 2 led to the formation of complex \([\text{Cs}[\text{U}(\text{OSi}(\text{O}^3\text{Bu}))_3]_2(\mu^-\text{HCOO})(\mu^-\text{NHCOO})]\), 4 in 44 % yield). The addition of less than 2 equivalent of CO\(_2\) shows only the presence of unreacted 2 and 4.

The solid-state structure of 4 (Figure 3) shows the presence of two U\(^{IV}\) cations bridged by a formate and a diaionic carbamate group. A cesium cation is also bound in the pocket formed by three siloxides bound to U1. Complex 4 is only the second example of a crystallographically characterized formate complex obtained from the insertion of CO\(_2\) into a U–H bond and the first one containing a bridging formate.\(^{[13b]}\) The U–O distances for the bridging formate in 4 (2.409(3) and 2.356(3) Å) are slightly shorter than those found in the reported terminal formate (2.451(3) Å). The proton of the formate was located by X-ray diffraction and found in the reported terminal formate (2.409(3) and 2.356(3) Å) of the formate complex obtained from the insertion of CO\(_2\) into an uranium hydride complex.\(^{[24]}\) The conformant cycloaddition of CO\(_2\) to the parent imido linkage affords a bridging κ\(^1\)-carbamate group. This reactivity is a rare example of cycloaddition of CO\(_2\) to a metal imido complex\(^{[25]}\) and differs from that reported for U\(^{IV}\)-bound terminal bulky aryl imido groups (U–N–Ar) that results in N=C bond methathesis and extrusion of the cyanate via a proposed carbamate intermediate.\(^{[25]}\) The low steric pressure of the NH group and its bridging mode result in a high stability of the κ\(^1\)-O,O,N-carbamate group. A few examples of U\(^{IV}\) carbamate have been reported but they all present monoaionic κ\(^2\)-O,O-carbamate groups.\(^{[23,24]}\)

In summary, we have demonstrated that a polymeric U\(^{IV}\) nitride effects the reversible heterolytic activation of dihydrogen in mild conditions (room temperature and 1 atmosphere of H\(_2\)). We have also demonstrated the high reactivity of this species. Notably, the hydride is readily transferred to acetonitrile and CO\(_2\), resulting in the substrate reduction. These results anticipate that metal hydride intermediates may play an important role in the uranium catalyzed transformation of N\(_2\) and H\(_2\) to ammonia and also open new avenues for the development of hydrogenation catalysts.

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Conflict of interest

The authors declare no conflict of interest.

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