This is an electronic reprint of the original article. This reprint may differ from the original in pagination and typographic detail.

Author(s): Brown, Rudy M.; Garcia, Javier Borau; Valjus, Juuso; Roberts, Christopher; Tuononen, Heikki; Parvez, Masood; Roesler, Roland

Title: Ammonia Activation by a Nickel NCN-Pincer Complex featuring a Non-Innocent N-Heterocyclic Carbene: Ammine and Amido Complexes in Equilibrium

Year: 2015

Version:

Please cite the original version:

All material supplied via JYX is protected by copyright and other intellectual property rights, and duplication or sale of all or part of any of the repository collections is not permitted, except that material may be duplicated by you for your research use or educational purposes in electronic or print form. You must obtain permission for any other use. Electronic or print copies may not be offered, whether for sale or otherwise to anyone who is not an authorised user.
Ammonia Activation by a Nickel NCN-Pincer Complex featuring a Non-Innocent N-Heterocyclic Carbene: Ammine and Amido Complexes in Equilibrium**

Rudy M. Brown, Javier Borau Garcia, Juuso Valjus, Christopher J. Roberts, Heikki M. Tuononen, Masood Parvez, and Roland Roesler*

Abstract: An NCN-pincer complex of nickel(0), 3, featuring a six-membered N-heterocyclic carbene (NHC) central platform and ammine pendant arms, was synthesized by deprotonation of its nickel(II) precursor, 2. It retained Cl⁻ in the square planar coordination sphere of nickel and was expected to be highly susceptible to oxidative addition reactions. Complex 3 rapidly activated ammonia at room temperature, in a ligand-assisted process where the carbene carbon played the unprecedented role of proton acceptor. For the first time, the coordinated (ammine, 4) and activated (amido, 5) species were observed together in solution, in a solvent-dependent equilibrium. A structural analysis of 2 and 3 provided insight into the highly unusual, non-innocent behaviour of the NHC ligand.

The activation of ammonia at metal centers is believed to play a crucial step in the catalytic functionalization of organic substrates, most notably olefins. Unfortunately, conducting this reaction with a Lewis base as a substrate poses substantial difficulties related to the activation-coordination equilibrium, which in the vast majority of cases is completely shifted in favor of the latter. Early work on ammonia activation involved TiIII by Armor and ZrIV by Bercaw,2 as well as OsCl₃(CO)₁₂ by Lewis and Süss-Fink.3 Iridium took center stage in the activation of ammonia starting with Milstein’s reports of oxidative addition of the N-H bonds at IrIV to yield dinuclear, bridging amido complexes L₆Ir₂(H)₂(NH₂)₂Ir₂H₆L₆.4 Similar chemistry was also investigated by Braun.5 Another significant development was the introduction of pincer complexes by Hartwig and later Turcuilet, who described the activation of ammonia on IrIII PCP and PSIP pincer complexes, respectively, to yield mononuclear, terminal LIr(NH₂)₂ derivatives.6 Also notable is the homolytic ammonia activation by a dinuclear PdP PNP pincer complex, reported by Ozerov, where a terminal LPd(NH₂)₂ species was obtained along with LPdH.7

Ammonia activation via metathesis, without formally changing the oxidation state of the metal, is well documented on several metals including Ru and Ir8,9 but can pose problems in terms of catalysis because the hydrogen is removed from the catalytic site. Milstein showed evidence for the heterolytic, deprotonative N-H bond activation of ammonia using a RuII complex incorporating non-innocent PNP pincer ligands, without observing the corresponding LRU₂⁻NH₂ species.9a The similar, ligand-assisted cleavage of the N-H bond in ammonia by an IrIV complex featuring a pyridyl-based PNP pincer ligand, to give a terminal LIr⁻NH₂ species, was described by Yoshizawa and Ozawa.9b A related ligand-assisted ammonia activation reaction was reported by Kuwata and Ikariya on a dinuclear LIr₂(H)(μ-H)(μ-NH₂)Ir₂L₆⁻ compound, resulting in the formation of LIr₂(μ-H)(μ-NH₂)Ir₂L₆⁻.9c Piers recently disclosed the use of a carbene-based PCP-pincer complex of Ni0 for the ligand-assisted activation of ammonia to give a terminal LNi₂⁻NH₂ product.10 Noteworthy are also the remarkable progresses made in the activation of ammonia by main group systems.11 The activation of the N-H bond in ammonia proved to be highly sensitive to the nature of the catalyst,12 with the ammine-amide equilibrium being completely shifted upon changing the metal from rhodium (amine) to iridium (amide) for the same ligand system.13 Moreover, the nature of the phosphine ligand was shown to have similar effects in iridium complexes.14 Homogeneous catalytic hydroamination with ammonia remains a major challenge, and notable progress towards this goal was the stoichiometric, intramolecular hydroamination of the coordinated ligand in an iridium amide complex.15

Scheme 1. Synthesis of derivatives 1 - 5.

Continuing our investigations of ammonia activation on pincer complexes incorporating NHC scaffolds, we report herein the reversible, ligand-assisted activation of ammonia on a Ni²⁻ - NCN pincer complex.

Pro-ligand 1 is similar to those reported by Byers and Lavoie and was prepared by the stepwise functionalization of 1,4,5,6-tetrahydropropyrimidine with N-Dipp-benzimidoyl chloride (Dipp =
2,6-diisopropylphenyl), using a modification of a reported procedure. Its room temperature reaction with Ni(cod)₂ (COD = cycloocta-1,5-diene) produced a deep red solution, from which complex 2 was isolated as an orange solid in 67% yield (Scheme 1). The ¹H NMR spectrum of 2 revealed a C₂ symmetry, instead of C₃₂ expected for a Ni-HNC complex. A resonance corresponding to the N₂CH proton appeared at 5.82 ppm, substantially high-field shifted from 8.84 ppm in 1. The crystal structure confirmed the distorted tetrahedral geometry of the N₂CH carbon on the NCN pincer complex, with the sum of the bond angles formed by the heavier elements at C1 measuring 332.5° (Figure 1). With 1.848(5) Å, the C1-Ni1 bond is short when compared to similar bonds in nickel(II) pincer complexes featuring dialkylmethyl or diarylmethyl central platforms (av. Ni-C 1.98(2) and 1.986(3) Å, respectively). Although the preference for alkyl vs. carbene coordination is common for complexes of dialkyl and diaryl carbones, it is very rare for complexes of the more stable NHCS. Only few such examples have been structurally characterized to date, and we propose that the competitive delocalization of the nitrogen lone pairs onto the exocyclic amide moieties is responsible for a reduced stabilization of the carbene structure. In support of this theory, the two C-N bonds of each pendant amide group in 2 are almost equal in length. Treatment with potassium hexamethyldisilazide in toluene yielded nevertheless the NCN complex 3, with nickel retaining the chlorine in its coordination sphere. The ¹H NMR spectrum of this Ni²⁰ complex confirmed the disappearance of the resonance corresponding to the N₂CH proton. A crystal structure of 3 revealed a dimeric arrangement with two anionic LNiCl moieties bridged by the potassium counterions (Figure 2). In turn, the π-coordination of the Dipπ groups completes the distorted square planar coordination geometry of the potassium ions. The carbene carbon C1 is close to planarity, with a sum of bond angles of 350.8°. At 1.795(7) Å, the Ni1-C1 bond in 3 is shorter than in 2, and in fact shorter than all reported Ni²⁰-CNC bonds (1.827(6) to 2.021(4) Å, av. 1.91(4) Å). The conversion to an NHC is accompanied by the expected shortening of the C-N bonds to the carbene carbon, from 1.44 Å in 2 to 1.39 Å in 3. These bonds remain nevertheless a bit long for in comparison to those observed in other NHC complexes (av. 1.36(2) Å), supporting a substantial contribution of resonance structure 3b (Scheme 2).

Complex 3 was expected to be prone to oxidative addition reactions upon elimination of KCl. Exposure to ammonia gas at room temperature in either toluene or THF yielded immediately a dark solution. ¹H NMR spectroscopy indicated the presence of two compounds, one with C₃ and the other with C₄ symmetry, which were assigned to 4 and 5, respectively. For the former compound, the ¹H NMR spectrum of the mixture displayed the two expected doublet resonances corresponding to the diastereotopic methyl groups of the ligand, and a resonance corresponding to the coordinated ammonia at -0.11 ppm in toluene-d₈ and 0.45 in THF-d₈. For the activation product 5, the spectrum displayed four doublet resonances corresponding to the methyl groups of the ligand, and a resonance for the amide protons at -2.97 ppm in toluene-d₈ and -3.63 ppm in THF-d₈. Additionally, the signal corresponding to the N₂CH proton was observed at 5.13 ppm in toluene-d₈ and 5.22 ppm in THF-d₈. The two structural isomers were present in 4 : 5 ratios of ca. 4 : 1 in toluene-d₈ and 2 : 3 in THF-d₈, with intermediate ratios in mixed-solvent systems, indicating that an equilibrium was present in solution (Figure 3). This was further confirmed by
EXSY NMR spectroscopy, which showed cross-peaks between the signals corresponding to the NH₃ protons in 4 and the NH₂ and N₅CH protons in 5 (see Supporting Information). Exposure of a solution of 4 and 5 in toluene-d₅ to ND₃ resulted, as expected, in the rapid decrease of the intensity of the ¹H NMR signals corresponding to the N₅CH, NH₂ and NH₂ protons following the exchange with deuterium. Solvent removal from solutions containing 4 and 5 produced a highly sensitive microcrystalline dark green solid that could be stored for weeks at -38 °C in the absence of air and moisture. Its identity could not be confirmed due to the insufficient quality of the single crystals. No distinctive IR bands corresponding to N-H vibrations were observed. The solid redissolved in THF and toluene giving solutions that displayed properties identical to those observed for the solution of in-situ prepared 4/5. Density functional theory calculations were performed on model compounds 4th and 5th in which the Dipp substituents were replaced by phenyl groups. The gas phase free energies of the optimized structures show that 4th is favored over 5th by 6 kJ·mol⁻¹ in low dielectric media, in good agreement with the value of 3.4 kJ·mol⁻¹ derived from the relative concentrations in toluene-d₅ solutions. When a continuum solvent model for THF was used, the calculations predicted 5th to be lower in energy than 4th by 21 kJ·mol⁻¹. This matches less well the experimentally derived value of 1.0 kJ·mol⁻¹, but it shows the same preference for 5th.

An equilibrium between coordinated (ammine) and activated (amido) ammonia complexes has been proposed before, mainly based on exchange experiments using ND₃. However, in some systems the incorporation of deuterium was likely to be a result of mechanisms other than the reversible oxidative addition of ammonia. The 4 - 5 system represents the first instance where the thermodynamic stability of the two complexes is comparable, allowing for the observation of both species in solution.

In conclusion, we report here a highly active Ni₉ pincer complex that is able to reversibly activate ammonia at room temperature in a ligand-assisted process. Both coordinated (ammine, 4) and activated (amido, 5) coexist in solution in comparable, solvent-dependent proportions. The unprecedented ability of the six-membered NHC in 3 to assist N-H bond activation, accepting the proton at the carbene carbon, opens up a new page in the chemistry of this versatile family of compounds that have found widespread applications as ancillary ligands.


