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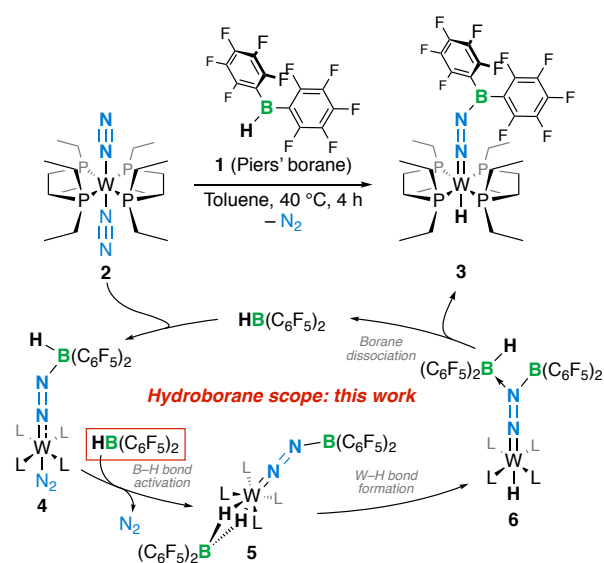
Borane-Catalysed Dinitrogen Borylation by 1,3-B–H Bond Addition

Anaïs Coffinet, Dan Zhang, Laure Vendier, Sébastien Bontemps and Antoine Simonneau*

The borylation of ligated dinitrogen by 1,3-B–H bond addition over a M–N≡N unit using various hydroboranes has been examined. In a previous study, we have shown that Piers' borane (**1**) reacted with the tungsten dinitrogen complex **2** to afford a boryldiazenido-hydrido-tungsten species. The ease and mildness of this reaction has pushed us to extend its scope, with the working hypothesis that **1** could potentially catalyse the 1,3-B–H bond addition of other hydroboranes. Under productive reaction conditions, dicyclohexylborane (HBCy₂) and diisopinocampheylborane (HBIPC₂) underwent retro-hydroboration to give cyclohexylborane (H₂BCy) or isopinocampheylborane (H₂BIPC), respectively, these monoalkylboranes acting as N₂-borylating agents in the presence of a catalytic amount of **1**. Under similar conditions, 9-borabicyclononane (9-BBN) slowly adds over the W–N≡N unit without rearrangement to a monoalkylborane. Catecholborane (HBcat) performs the 1,3-B–H bond addition without the need of a catalyst. We were not able to build more than one covalent B–N bond between the terminal N of the N₂ ligand and the boron reagent with this methodology.

Introduction

The chemistry of dinitrogen complexes has attracted important research efforts over almost 60 years.¹ Indeed, they can serve as platforms to transform the inert but abundant N₂ molecule, and since the early 2000s several catalysts for the reduction of dinitrogen into ammonia² under ambient temperature and pressure have been devised. This contrasts with the harsh conditions found in the Haber-Bosch plant,³ where N₂ is converted into ammonia by hydrogenolysis at the industrial level. N₂ complexes can also serve as models for the active sites of the nitrogenases,⁴ the enzymes responsible for biological nitrogen fixation. Beyond ammonia and hydrazine, the quest for the direct conversion of dinitrogen into more complex nitrogen-containing compounds has fuelled interest towards the construction of N–E bonds (E = C, B, Si, Al, Ga) at ligated dinitrogen.^{1a} In this perspective, our team has reported N₂ silylation and borylation methods inspired by the chemistry of the Frustrated Lewis Pairs.⁵ We have notably shown that in the presence of Piers' borane [HB(C₆F₅)₂, **1**],⁶ compound *trans*-[W(N₂)₂(depe)₂] [**2**, depe = 1,2-bis(diethylphosphino)ethane] underwent a formal 1,3-addition of the B–H bond across the W–N≡N unit (**2** → **3**, Scheme 1),^{5b} a complementary N₂ borylation method to 1,3-B–X bond addition.⁷ The proposed mechanism to account for this transformation, based on experimental observations, involved two molecules of **1**, one being the N₂ borylating agent, the other assuming the role of B–H bond cleavage and hydride shuttling to the metal centre, thus closing a catalytic cycle (Scheme 1). As a continuation of this work, we wanted to check whether this 1,3-B–H addition can be extended to other hydroboranes and if Piers' borane would be a competent catalyst for this reaction.⁸



Scheme 1. Previously reported^{5b} 1,3-B–H bond addition of Piers' borane (**1**) on dinitrogen complex **2** and the proposition of a borane-catalysed mechanism (L = ½ depe).

Results

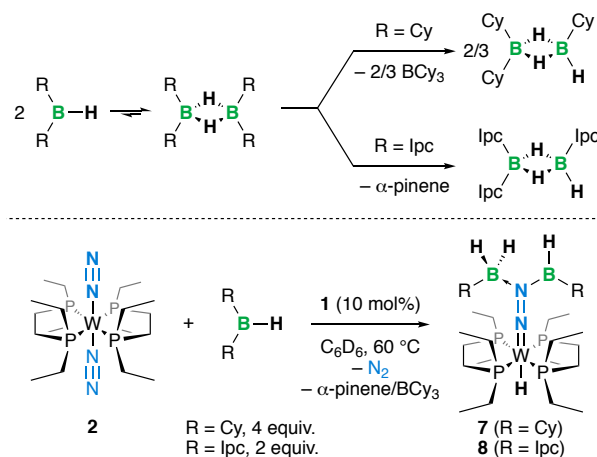
Four different organo-hydroboranes, namely dicyclohexylborane (HBCy₂), diisopinocampheylborane (HBIPC₂), 9-BBN (9-borabicyclononane) and catecholborane (HBcat) were examined. Three types of reactivities have been observed, as detailed below.

Reactions of **2** with dicyclohexylborane (HBCy₂) and diisopinocampheylborane (HBIPC₂)

We reacted *trans*-[W(N₂)₂(depe)₂] **2** with one equivalent of dicyclohexylborane (HBCy₂) at room temperature in C₆D₆. NMR spectroscopy analysis revealed the absence of new tungsten species, and raising the temperature to 80 °C overnight did not change this outcome. This contrasts with the reaction of **2** with the electrophilic hydroborane **1**, occurring under milder conditions and during which the Lewis acid-base adduct thereof (**4**, Scheme 1) could be detected.^{5b} ¹¹B NMR analysis revealed the presence tricyclohexyldiborane and BCy₃ (see Fig. S1). This substituent scrambling presumably arose from a retro-hydroboration, a reaction known for the ethyl- and

isopinocampheyl-substituted boranes, and readily occurs in the absence of **2** (Scheme 2, top and Fig. S1).⁹ To verify whether $\text{HB}(\text{C}_6\text{F}_5)_2$ (**1**) could catalyse 1,3-B–H bond addition of HBCy_2 (or in-situ-formed H_2BCy), we repeated the experiment in the presence of 10 mol% of **1** at 60 °C in C_6D_6 . After a day, NMR analysis revealed again the presence of BCy_3 as well as tricyclohexyldiborane. We also noticed the formation of two new metal complexes according to $^{31}\text{P}\{^1\text{H}\}$ NMR with singlets at δ 45.2 and 41.7 ppm, **2** remaining the major product of the mixture. Hoping to reach higher conversion, we increased the amount of HBCy_2 . ^{31}P NMR analysis of the reaction mixture run with four equivalents thereof during 2 hours at 60 °C revealed conversion to compound **7** (Scheme 2, bottom). The ^1H NMR spectrum of the crude reaction mixture showed a hydride quintet coupling to four magnetically equivalent phosphorus nuclei (δ –3.38, quintet, $^2J_{\text{HP}} = 30$ Hz). Diagnostic signals for a four-coordinated boron species carrying 2 hydrogens were observed in the ^1H and ^{11}B NMR spectra (^1H : δ 3.29, broad singlet; $^1\text{H}\{^{11}\text{B}\}$: δ 3.29, singlet; ^{11}B : δ –30.0, triplet, $J_{\text{BH}} = 91$ Hz). Layering a concentrated toluene solution with pentane afforded crystalline material (76% yield) suitable for an X-ray diffraction study. This allowed us to confirm that the targeted formal 1,3-B–H bond addition had been achieved, with H_2BCy being the actual borylating agent (Fig. 1), thus explaining the need for an excess of HBCy_2 to reach full conversion of **2**.

In the molecular structure, a second, tetracoordinated boron moiety is found. Its presence is explained by the capture of another equivalent of H_2BCy , forming a Lewis pair with the distal, Lewis basic nitrogen of the boryldiazenido moiety [$\text{N}_2\text{--B}2$ 1.618(6) Å]. The latter could not be observed in the ^{11}B NMR spectrum and is characterised by a B–N linkage having double bond character [$\text{N}_2\text{--B}1$ 1.405(6) Å].¹⁰ The distal N atom adopts a trigonal planar geometry ($\sum \text{angles} = 360^\circ$), yet the wide B–N–B angle (126 °), presumably originating from steric repulsions between the boron substituents, deviates it from ideality. The N–N bond length [1.346(3) Å] lies halfway between the ones of double (ca. 1.25 Å) and single (ca. 1.45 Å) N–N bonds. This noticeable elongation ($d_{\text{N--N}}$ 1.1233(17) Å in **2**) results from the formal two-electron reduction of the N_2 ligand upon functionalisation, as well as the effect of H_2BCy complexation pulling electron density from the metal into antibonding orbitals of the boryldiazenido ligand.¹¹ The W–N bond is 1.789(2) Å long, indicative of double bond character,¹² and the W centre lies in a distorted octahedron (all P–W–N angles > 90° and up to 106°). Likewise, the W–N–N unit deviates from linearity [$\text{W}1\text{--N}1\text{--N}2$ 170 °]. Those deviations from ideal geometries are, again, probably of steric origin. Overall, these structural features compare well with those of the previously reported compound **6** (Scheme 1).^{5b}



Scheme 2 Generation of monoalkylboranes from HBR_2 (top) and reactions of $\text{trans-[W}(\text{N}_2)_2(\text{depe})_2]$ (**2**) in the presence of dicyclohexyl- or diisopinocampheylborane catalysed by **1** (bottom).

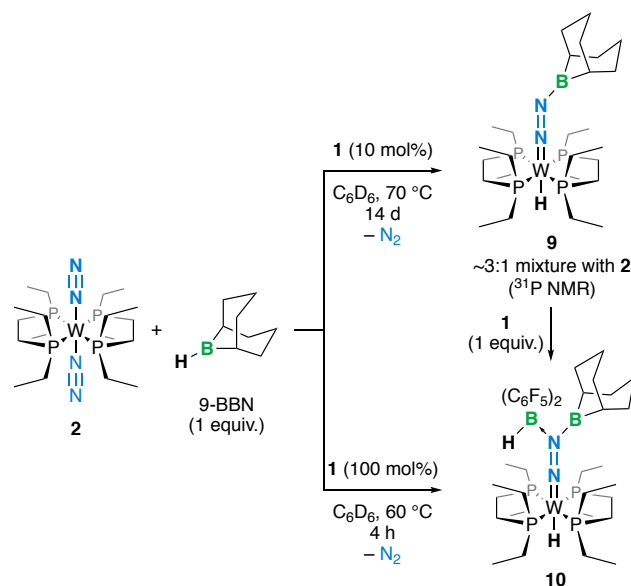
Comparatively to HBCy_2 , diisopinocampheylborane ($\text{HB}(\text{lpc})_2$) did not react with **2** even under forcing conditions. At room temperature and in C_6D_6 , $\text{HB}(\text{lpc})_2$ also underwent a substituent rearrangement (Scheme 2, top). Indeed, its dimer evolves into triisopinocampheylborane and α -pinene (see Fig. S6).^{9a} Having established from the above-discussed reaction involving HBCy_2 that dihydroboranes seem to add more easily to the W– N_2 unit than monohydroboranes, we adjusted the number of equivalents of $\text{HB}(\text{lpc})_2$ in order to generate the right amount of $\text{H}_2\text{B}(\text{lpc})_2$ in the reaction mixture, and added a catalytic amount (10 mol%) of **1**. With two equivalents of $\text{HB}(\text{lpc})_2$, full and selective conversion of **2** to compound **8** was reached after 15 min at 60 °C (Scheme 2, bottom). Although the newly formed compound **8** could not be crystallised, its NMR characterisations are very similar to compound **7**: the ^1H NMR spectrum showed a hydride signal coupling to the four phosphorus (δ –3.33, quintet, $^2J_{\text{HP}} = 30.6$ Hz), and the ^{11}B NMR featured a triplet at –30.0 ppm, leaving little doubt on the identity of **8**.

Reactions with 9-Borabicyclo[3.3.1]nonane (9-BBN)

The reactivity of $\text{trans-[W}(\text{N}_2)_2(\text{depe})_2]$ **2** with one equivalent of 9-borabicyclo[3.3.1]nonane (9-BBN) at room temperature in C_6D_6 was also examined. Likewise, no reaction occurred even under forcing conditions (2 d, 80 °C). Hence, we added 10 mol% of hydroborane **1** to an equimolar solution of **2** and 9-BBN. $^{31}\text{P}\{^1\text{H}\}$ NMR monitoring of the reaction after few days at 60 °C revealed that two new products **9** and **10** started to form (δ 45.0 and 40.7 ppm, respectively), both showing a corresponding hydride quintet in ^1H NMR (δ –3.33 and –4.10 ppm, respectively). Upon prolonged heating, compound **9** formed predominantly, reaching ca. 70% of the mixture according to ^{31}P NMR integration after 14 days at 70 °C and on the basis of NMR spectroscopy, was assigned to the product of 1,3-B–H bond addition (Scheme 3). The other tungsten-containing components of the reaction mixture were **2** and unidentified minor impurities (see Fig. S12), while several unidentified boron species seemed to be present (Fig. S13). The slow rate of this B–H bond addition incited us to perform the stoichiometric

reaction between complex **2**, **1** and 9-BBN. Analysis of the reaction mixture by ^{31}P NMR showed full and selective conversion to compound **10** (Scheme 3) after 4 h at 60 °C. Traces of **10** were also observed in the ^{31}P and ^{19}F NMR spectra acquired at the end of the catalytic reaction (see Fig. S14). Analysis of the ^{19}F and ^{11}B NMR spectra of **10** suggested, in accordance with previous results,^{5b} that it is formed by Lewis acid-base interaction of **1** with the boryldiazenido moiety of **9**. The reaction of **9** with **1** to give **10** brought convincing evidences supporting the latter assumption, as well as the identity of **9**, for which we could not ascertain the structure with X-ray diffraction.

An X-ray diffraction study on single crystals allowed us to confirm the proposed structure for **10** (Fig. 1), the covalent vs. dative bonding between the boron atoms and the terminal N being confirmed by comparison of B–N bond lengths [N2–B1, 1.405(6) Å vs. N2–B2, 1.617(6) Å]. The structure of **10** resembles the one of **7** with a tungsten centre in a distorted octahedron, and the W–N–N unit deviating from linearity (W1–N1–N2, 175°). While W–N bond lengths are similar in **7** and **10**, the N–N bond in **10** is slightly longer (+0.03 Å), probably as a result of the higher electrophilicity of Piers' borane than H_2BCy .



Scheme 3 Reactions of *trans*-[W(N₂)₂(depe)₂] (**2**) with 9-BBN and catalytic or stoichiometric amounts of HB(C₆F₅)₂ (**1**).

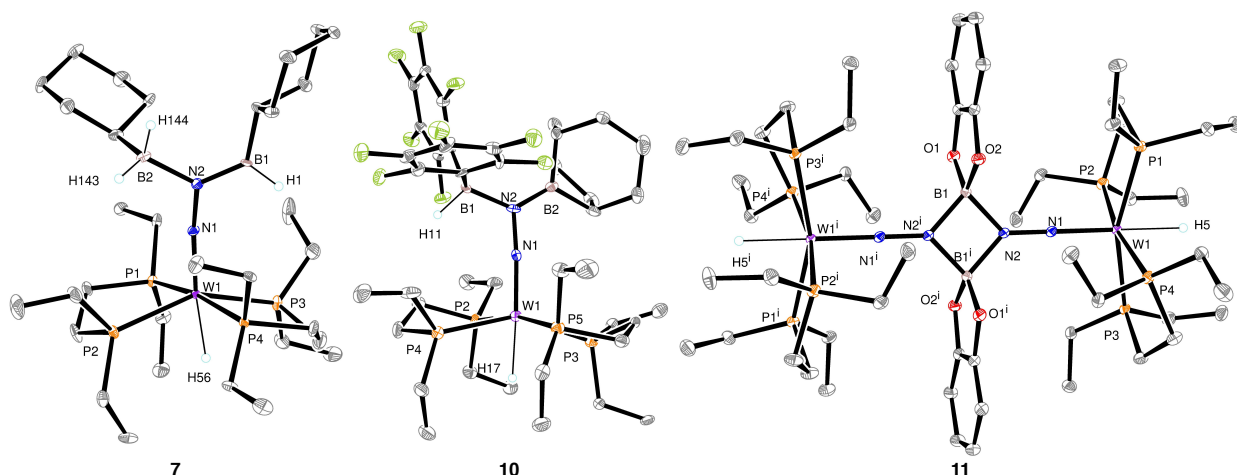
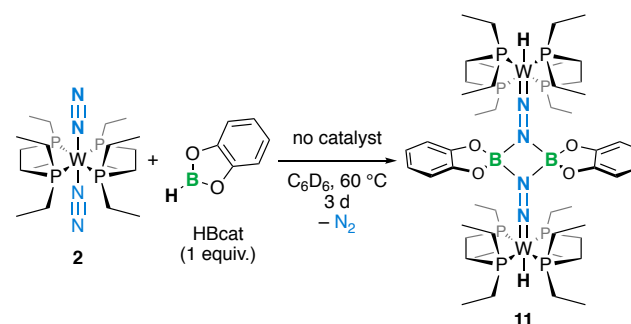


Fig. 1 Solid-state molecular structures of compounds **7**, **10** and **11**. Ellipsoids drawn at the 50% level of probability. All hydrogens have been omitted for clarity except those bound to B or W. Selected distances (Å) and angles (°) for **7**: W1–N1, 1.789(2); N1–N2, 1.346(3); N2–B1, 1.405(6); N2–B2, 1.618(6); W1–H56, 1.77(6); W1–N1–N2 170.3(3); B1–N2–B2 126.4(3). **10**: W1–N1, 1.783(6); N1–N2, 1.383(9); N2–B1, 1.56(1); N2–B2, 1.41(1); W1–H17, 1.82; W1–N1–N2, 175.2(5); B1–N2–B2, 129.3(7). **11**: W1–N1, 1.807(3); N1–N2, 1.295(4); N2–B1, 1.545(6); N2–B1, 1.536(6); W1–H5, 1.87(5); W1–N1–N2, 178.6(3); B1–N2–B1, 94.1(3); N2–B1–N2, 85.9(3).

Reaction with catecholborane (HBcat)

Next, we checked the reactivity of *trans*-[W(N₂)₂(depe)₂] **2** with one equivalent of catecholborane (HBcat) at room temperature in C₆D₆. Quite contrastingly, NMR analysis revealed that a reaction occurred in a selective manner. Indeed, we observed a new diamagnetic compound **11** having a singlet resonance at δ 43.8 ppm in $^{31}\text{P}\{^1\text{H}\}$ NMR with a corresponding hydride quintet at $-\text{3.6}$ ppm in ^1H NMR after few minutes at room temperature. Almost full conversion was obtained after 3 days at 60 °C (>90 % according to ^{31}P NMR). Compared to the previous examples, Piers' borane **1** was not necessary to perform the targeted formal 1,3-B–H bond addition.



Scheme 4 Reaction of *trans*-[W(N₂)₂(depe)₂] **1** with catecholborane (HBcat).

Layering the reaction mixture with pentane allowed us to grow single crystals of **11** (33% yield) suitable for X-ray

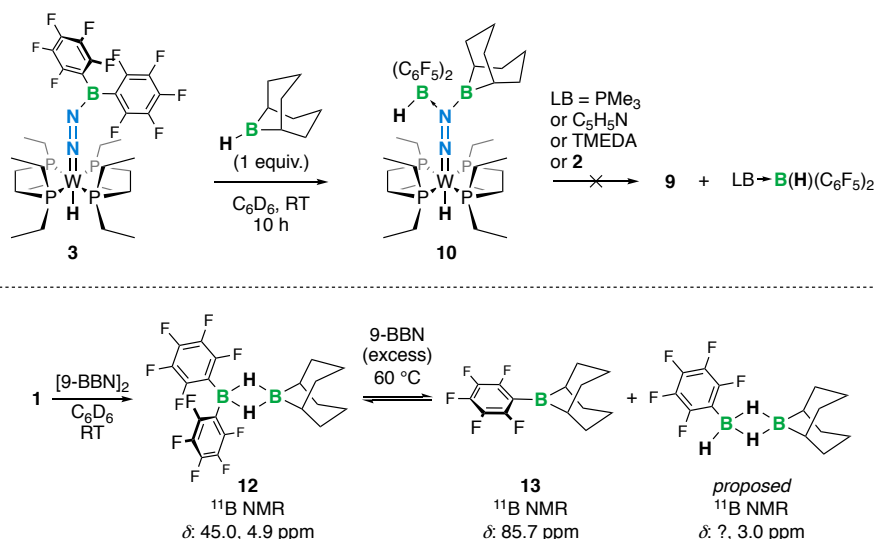
diffraction analysis, that revealed a dimeric solid-state structure of D_{2h} symmetry for **7** (Fig. 1): two boryl groups are bridging two W-N₂ units in an almost square BNB arrangement (N-B bond lengths ca. 1.54 Å; B-N-B angles 94°, N-B-N angles 86°). Compared to the structures of **3** and **6**, the octahedron formed by the W ligands is less distorted, and the W-N-N arrangement is linear (W1-N1-N2, 179°). The N-N bond is also shorter [1.295(4) Å]. Similar dimeric structures have been obtained from the reactions of group 13 chlorides with end-on dinitrogen complexes.¹³ The dimeric nature of **11** in the solid, which contrasts with the one of **7** or **9** (as suggested by diffusion-ordered NMR spectroscopy, see Fig. S15), is likely to result from the diminished steric hindrance of the flat Bcat group.

Discussion

The above-described experiments have confirmed the possibility to use Piers' borane as a catalyst for the borylation of N₂ bound to a group 6 metal centre by 1,3-B-H bond addition when using dialkylhydroboranes (HBCy₂, HBIpc₂ and 9-BBN). Within this family of compounds, two different behaviours have been observed depending on the propensity of the dialkylhydroborane to undergo retro-hydroboration under the reaction conditions to give a monoalkylborane. The latter is the active borylating agent, and in the case of HBCy₂ and HBIpc₂, only the products of 1,3-B-H bond addition of H₂BCy and H₂BIpc over the W-N-N unit were observed. The reason is most probably of kinetic origin, since monoalkylboranes are less sterically hindered than their disubstituted parents, making the interaction with the N₂ complex as well as the B-H bond activation by **1** easier. 9-BBN, which does not undergo retro-hydroboration, reacts much slower and also needs catalytic amounts of **1**. Interestingly, when the reaction of 9-BBN with the dinitrogen complex is conducted in the presence of an equimolar amount of **1**, the B-H bond addition proceeds very rapidly *and* selectively. Indeed, the 1,3-B-H bond addition product of Piers' borane^{5b} (**3**, Scheme 1) is not observed. The following experiment allowed us to shed light on this fact: when product **3** was treated with an equimolar amount of 9-BBN at RT, **10** was obtained quantitatively in a few hours (Scheme 5). This result suggests that in the catalytic 1,3-B-H bond addition mechanism with H₂BCy, H₂BIpc or 9-BBN, addition of Piers' borane to give **3** may precede interaction of the alkylhydroborane with the M-N≡N unit. After complexation of the latter to the thus-formed C₆F₅-substituted boryldiazene moiety in **3**, boron-to-boron hydride transfer then affords a Piers' borane-complexed 1,3-B-H bond addition compound such as **10**. Dissociation of **1** followed by reaction with dinitrogen complex **2** would then close the catalytic sequence. However, further experimental work cast doubts on this conjecture, at least in the case of 9-BBN. Attempts to dissociate

1 in compound **10** in the presence of various Lewis bases such as PMe₃, pyridine, tetramethylethylenediamine (TMEDA) or **2** at elevated temperatures for several hours proved unsuccessful (Scheme 5, top).¹⁴ We have checked the integrity of Piers' borane (**1**) in the presence of a 10-fold excess of 9-BBN: at RT, a new species forms in C₆D₆ within few hours, which we believe is the mixed diborane **12** (Scheme 5, bottom). Upon heating to a temperature relevant to the above-described experiments, the putative dimer seems to get involved in an equilibrated substituent rearrangement process with two other compounds, one being 9-C₆F₅-BBN¹⁵ (**13**) while the second one is presumably a mixed diborane composed of 9-BBN and in-situ-formed H₂BC₆F₅ (see Fig. S22-23 and Scheme 5). The ¹⁹F NMR spectrum recorded at the end of the reaction affording **9** does not feature the spectral signature of **1**, but unambiguously shows the presence of **10** along with another unidentified compound (see Fig. S13-14). These data suggests that **1** is a precatalyst in the reaction involving 9-BBN, but the presence of **10** at the end of the reaction also implies this species might be a catalyst resting state.

HBcat stands out against the 3 other boranes examined. This boron reagent is known to be a much milder hydroboration reagent than HBCy₂ or HBIpc₂, due to its quenched Lewis acidity.¹⁶ As such, HBcat is also a weak hydride donor compared to 9-BBN on the basis of calculated hydride donor abilities [ΔG_{H^-} (298 K) = 159.2 vs. 99.0 kcal/mol, respectively], but this trend reverses upon Lewis base [for example, NHC-HBcat, ΔG_{H^-} (298 K) = 44.8 kcal/mol and NHC-9-BBN, ΔG_{H^-} (298 K) = 47.0 with NHC = 1,3-di(isopropyl)imidazole-2-ylidene] or hydride coordination {[H₂BCat]⁻, ΔG_{H^-} (298 K) = 26.3 kcal/mol and [9-H₂BBN]⁻, ΔG_{H^-} (298 K) = 32.8}.¹⁷ Although the flatness of HBcat and its monomeric nature in non-coordinating solvents also offers a kinetic advantage, we believe the peculiarity of HBcat reactivity lies in its electronic properties. We propose a similar mechanism for 1,3-B-H bond addition of HBcat than with Piers' borane (Scheme 1): Lewis acid-base adduct formation of HBcat with **2** increases the polarisation of the B-H bond to an extent that a second equivalent of HBcat is now able to abstract the hydride, generating the [H₂Bcat]⁻ anion. The latter is a competent hydride donor (*vide supra*), especially when compared to [H₂B(C₆F₅)₂]⁻ [ΔG_{H^-} (298 K) = 61.5 kcal/mol], which also bears a steric penalty due to its C₆F₅ substituents. This may explain why in the reactions with HBcat we were not able to detect species resembling **5** (Scheme 1), the W-H bond formation event being probably too fast under the experimental conditions. In the case of Piers' borane, we surmise that an important factor for W-H bond formation (**5** → **6**, Scheme 1) is the nature of the *trans* ligand, NNB(C₆F₅)₂, which electron-withdrawing properties might counter-balance the weak hydride donor ability of [H₂B(C₆F₅)₂]⁻.¹⁸



Scheme 5 Reaction of **3** with 9-BBN and attempts to dissociate **1** from **10** with various Lewis bases (top) and reaction of **1** with 9-BBN in the absence of a dinitrogen complex (bottom).

Conclusions

In the present article, we expand the scope of 1,3-B–H bond addition over a W–N₂ unit. This mild method for N₂ borylation, which complements 1,3-B–X bond addition,⁷ can be attractive for who seeks to synthesise nitrogen-boron compounds directly from dinitrogen. Having this goal in mind, we have examined the reactivity of 4 different boranes. Only HBCat reacts spontaneously with the dinitrogen complex **2**, affording the 1,3-B–H bond addition product **11**. Hydroboranes HBCy₂, HBIPc₂ and 9-BBN required catalytic amounts of Piers' borane (**1**) to react with **2**. In the case of HBCy₂ and HBIPc₂, monoalkylboranes H₂BCy and H₂BIpc formed in-situ are the actual borylating agents. None of the boranes examined have allowed to build more than one covalent N–B bond. According to experimental observations, we believe the mechanism of these catalytic reactions starts with 1,3-B–H bond addition of **1** to give **3**, followed by complexation of the alkylhydroborane to the terminal N of the resulting W-ligated boryldiazenido ligand and a boron-to-boron hydride transfer ensues. While it is reasonable to think that dissociation of **1** should close the catalytic cycle, such elementary step could not be validated in the case of 9-BBN because **1** is likely to be a precatalyst in this specific reaction. This urges us to explore in detail the reactivity of H₂BC₆F₅ but also L•BH₃ (L = THF, SMe₂, etc...) or diborane (B₂H₆) with zero-valent group 6 dinitrogen complexes. We will share the results of this future work in due time.

Experimental Section

General considerations

All reactions were performed in flame- or oven-dried glassware with rigorous exclusion of air and moisture, using a nitrogen filled *Jacomex* glove box (O₂ < 1 ppm, H₂O < 1 ppm). Solvent used were pre-dried (toluene and *n*-pentane by passing through a Puresolv MD 7 solvent purification machine; *n*-hexane and

hexamethyldisiloxane (HMDSO) by distillation over CaH₂), degassed by freeze-pump-thaw cycles, dried over molecular sieves and stored in the glove box. C₆D₆ (purchased from *Eurisotop*) was degassed by freeze-pump-thaw cycles, dried over molecular sieves and stored in the glove box. HB(C₆F₅)₂ (**1**),^{6a} HBCy₂¹⁹ and HBIPc₂²⁰ were synthesized according to reported procedures and stored in the glove box. 9-BBN and HBCat were purchased from Sigma-Aldrich and used as received in the glove box. ¹H, ¹¹B, ¹⁹F and ³¹P NMR spectra were recorded in C₆D₆ or toluene-*d*₈ using NMR tubes equipped with J. Young valves on a *Bruker Avance III 400* spectrometer. Chemical shifts are in parts per million (ppm) downfield from tetramethylsilane and are referenced to the residual solvent resonance as the internal standard (C₆HD₅: δ reported = 7.16 ppm; C₇HD₇: δ reported = 2.08 ppm for ¹H NMR). ¹¹B, ¹⁹F, ²⁹Si and ³¹P NMR spectra were calibrated according to the IUPAC recommendation using a unified chemical shift scale based on the proton resonance of tetramethylsilane as primary reference.²¹ Data are reported as follows: chemical shift, multiplicity (br = broad, s = singlet, d = doublet, t = triplet, q = quartet, quint = quintet, m = multiplet, m_c = centrosymmetric multiplet), coupling constant (Hz), and integration. Infrared (IR) spectra were recorded in the glove box on an *Agilent Cary 630* FT-IR spectrophotometer equipped with ATR or transmission modules and are reported in wavenumbers (cm⁻¹). Elemental analyses were performed on samples sealed in tin capsules under N₂ by the Analytical Service of the Laboratoire de Chimie de Coordination; results are the average of two independent measurements.

Reaction of **2** with HBCy₂ catalysed by **1**

In a glovebox, *trans*-[W(N₂)₂(depe)₂] (**2**, 52 mg, 80 μ mol, 1 equiv.), HBCy₂ (57 mg, 32 μ mol, 4 equiv.) and HB(C₆F₅)₂ (**1**, 3 mg, 8 μ mol, 0.1 equiv.) were weighed in a 4-mL vial. After addition of C₆D₆ (0.6 mL), the orange-red solution was heated for 2 hours at 60 °C. The resulting orange solution was concentrated to ca. 0.3 mL under reduce pressure and layered with pentane before

storage at $-40\text{ }^{\circ}\text{C}$. After a week, orange crystals of **7** were recovered by decantation and dried under vacuum (50 mg, 61 μmol , 76% yield). Single crystals suitable for an X-ray diffraction study were obtained from the same crop.

7: $^1\text{H NMR}$ (400 MHz, C_6D_6) δ : 3.29 (br s, 2H), 2.21–2.02 (m_c , 8H), 2.06 (t, $J = 11.6\text{ Hz}$, 8H), 1.94 (s, 2H), 1.82–1.63 (m, 5H), 1.63–1.38 (m, 14H), 1.39–1.18 (m, 10H), 1.14 (quint, $J = 7.6\text{ Hz}$, 12H), 0.73 (p, $J = 7.4\text{ Hz}$, 12H), -3.38 (quint, $J_{\text{HW}} = 15.8\text{ Hz}$, $J_{\text{HP}} = 30.4\text{ Hz}$, 1H). $^{31}\text{P}\{^1\text{H}\}$ NMR (162 MHz, C_6D_6) δ : 41.7 ($J_{\text{PW}} = 282\text{ Hz}$). $^{11}\text{B NMR}$ (128 MHz, C_6D_6) δ : -30.0 (t, $J_{\text{BH}} = 91.0\text{ Hz}$).

IR (ATR) $\nu/\text{cm}^{-1} = 2958, 2900, 2783, 1603, 1503, 1456, 1362, 1335, 1274, 1188, 1163, 1124, 1095, 1081, 1029, 982, 867, 802, 755, 732, 710, 693, 686, 662$.

Elem. Anal. calcd. for $\text{C}_{32}\text{H}_{74}\text{B}_2\text{N}_2\text{P}_4\text{W}$: C, 47.08; H, 9.14; N, 3.43. Found: C, 47.10; H, 8.90; N, 3.26.

Reaction of **2** with HBIpc₂ catalysed by **1**

In a glovebox, *trans*-[W(N₂)₂(depe)₂] (**2**, 13 mg, 20 μmol , 1 equiv.), HBIpc₂ (12 mg, 40 μmol , 2 equiv.) and HB(C₆F₅)₂ (**1**, 0.7 mg, 2.0 μmol , 0.1 equiv.) were weighed in a 4-mL vial. After addition of C_6D_6 (0.6 mL), the orange-red solution was heated at $60\text{ }^{\circ}\text{C}$ for 15 minutes, time after which full conversion of **2** into **8** was ascertained by NMR analysis. After evaporation to dryness in the glovebox, the oily residue was triturated in pentane. The supernatant was discarded, and the oily residue was dried under vacuum. Compound **8** could not be isolated otherwise than as a C_6D_6 -containing clathrate.

8: $^1\text{H NMR}$ (400 MHz, C_6D_6) δ : 3.38 (br s, 3H), 2.59–2.50 (m, 2H), 2.43–2.34 (m, 4H), 2.27–2.13 (m, 12H), 2.06–1.97 (m, 3H), 1.82 (m, 2H), 1.74–1.65 (m, 3H), 1.48 (d, $J = 7.2\text{ Hz}$, 8H), 1.45 (s, 6H), 1.37 (s, 6H), 1.20–1.10 (m, 16H), 0.80–0.68 (m, 20H), -3.33 (quint, $J_{\text{HW}} = 13.4\text{ Hz}$, $J_{\text{HP}} = 30.7\text{ Hz}$, 1H). $^{31}\text{P}\{^1\text{H}\}$ NMR (162 MHz, C_6D_6) δ : 41.2 (m_c , $J_{\text{WP}} = 282\text{ Hz}$, second order pattern). $^{31}\text{P NMR}$ (162 MHz, C_6D_6) δ : 41.2 (m_c , second order pattern). $^{11}\text{B NMR}$ (128 MHz, C_6D_6) δ : -3.32 (br s), -30.0 (t, $J_{\text{BH}} = 88.9\text{ Hz}$).

IR (ATR) $\nu/\text{cm}^{-1} = 2962, 2936, 2878, 2357, 2081, 1634, 1501, 1453, 1416, 1293, 1104, 1074, 1027, 961, 869, 806, 751, 735, 683$.

Elem. Anal. calcd. for $\text{C}_{40}\text{H}_{86}\text{B}_2\text{N}_2\text{P}_4\text{W}\cdot 0.55\text{C}_6\text{D}_6$: C, 53.58; H, 8.93; N, 2.89. Found: C, 54.09; H, 9.55; N, 2.56.

Reaction of **2** with 9-BBN catalysed by **1**

In a glovebox, *trans*-[W(N₂)₂(depe)₂] (**2**, 26 mg, 40 μmol , 1 equiv.), 9-BBN (4.9 mg, 40 μmol , 1 equiv.) and HB(C₆F₅)₂ (**1**, 1.4 mg, 4.0 μmol , 0.1 equiv.) were weighed in a 4-mL vial. After addition of C_6D_6 (0.5 mL), the dark orange solution was heated at $70\text{ }^{\circ}\text{C}$ for 14 days, time after which compound **9** become the main component of the reaction mixture (ca. 70% according to ^{31}P NMR) that still contained **2** and other unidentified boron species. Attempts to isolate **9** in an analytically pure form have failed.

9: $^1\text{H NMR}$ (400 MHz, C_6D_6) δ : 2.08 - 1.99 (m, 12H), 1.89 - 1.85 (m, 3H), 1.80 - 1.71 (m, 6H), 1.68 - 1.62 (m, 3H), 1.51 (s, 2H), 1.47 - 1.34 (m, 12H), 1.17 (dt, $J = 14.8, 7.6\text{ Hz}$, 12H), 0.87 (dt, $J = 13.6, 7.5\text{ Hz}$, 12H), -3.33 (quint, $J_{\text{HW}} = 26.1\text{ Hz}$, $J_{\text{HP}} = 26.3\text{ Hz}$, 1H). $^{31}\text{P}\{^1\text{H}\}$ NMR (162 MHz, C_6D_6) δ : 45.03 ($J_{\text{WP}} = 282\text{ Hz}$). $^{11}\text{B NMR}$ (128 MHz, C_6D_6) δ : 45.9.

Reaction of **2** with 9-BBN and **1**

In a glovebox, *trans*-[W(N₂)₂(depe)₂] (**2**, 20 mg, 30 μmol , 1 equiv.), 9-BBN (3.8 mg, 30 μmol , 1 equiv.) and HB(C₆F₅)₂ (**2**, 10 mg, 30 μmol , 1 equiv.) were weighed in a 4-mL vial. After addition of C_6D_6 (0.6 mL), the dark orange solution was heated at $60\text{ }^{\circ}\text{C}$ for 4 hours. The resulting orange solution was concentrated under vacuum and pentane (2 mL) was added before storage at $-40\text{ }^{\circ}\text{C}$. After one day, orange crystals of **10** were recovered by decantation and dried under vacuum (23 mg, 21 μmol , 70% yield). Single crystals suitable for X-ray diffraction crystallography were obtained from the same crop.

10: $^1\text{H NMR}$ (400 MHz, C_6D_6) δ : 3.98 (d, $J = 107.1\text{ Hz}$, 1H), 2.15 - 1.98 (m, 4H), 1.97 - 1.85 (m, 4H), 1.85 - 1.77 (m, 2H), 1.72 (q, $J = 4.7, 4.2\text{ Hz}$, 4H), 1.65 - 1.44 (m, 10H), 1.39 - 1.26 (m, 6H), 1.22 - 1.07 (m, 8H), 1.00 (quint, $J = 7.4\text{ Hz}$, 12H), 0.62 (quint, $J_{\text{HW}} = 9.8\text{ Hz}$, $J_{\text{HP}} = 7.4\text{ Hz}$, 12H), -4.10 (quint, $J_{\text{HP}} = 34.1\text{ Hz}$, 1H). $^{31}\text{P}\{^1\text{H}\}$ NMR (162 MHz, C_6D_6) δ : 40.7 ($J_{\text{PW}} = 287\text{ Hz}$). $^{11}\text{B NMR}$ (128 MHz, C_6D_6) δ : -10.9 (s), the boron shift of the tricoordinated boryl group could not be detected. $^{19}\text{F NMR}$ (377 MHz, C_6D_6) δ : -132.7 (dd, $J_{\text{FF}} = 25.5, 9.3\text{ Hz}$, 4F_{ortho}), -162.5 (t, $J_{\text{FF}} = 20.2\text{ Hz}$, 2F_{para}), -166.0 - -166.5 (m, 4F_{meta}).

IR (ATR) $\nu/\text{cm}^{-1} = 2965, 2937, 2917, 2881, 2354, 1639, 1507, 1452, 1375, 1332, 1264, 1228, 1108, 1091, 1075, 1039, 1028, 964, 936, 887, 869, 844, 803, 736, 712, 680, 662$.

Elem. Anal. calcd. for $\text{C}_{40}\text{H}_{64}\text{B}_2\text{F}_{10}\text{N}_2\text{P}_4\text{W}$: C, 43.98; H, 5.91; N, 2.65. Found: C, 44.00; H, 6.08; N, 2.48.

Reaction of **2** with HBCat

In a glovebox, *trans*-[W(N₂)₂(depe)₂] (**2**, 52 mg, 80 μmol , 1 equiv.) and HBCat (8.6 μL , 80 μmol , 1.0 equiv.) were introduced in a 4-mL vial. After addition of C_6D_6 (0.6 mL), the orange solution was heated at $60\text{ }^{\circ}\text{C}$ for 3 days. The resulting solution was then concentrated under vacuum before adding hexane (2 mL). The resulting turbid solution was centrifuged, giving a yellow liquid that was stored at $-40\text{ }^{\circ}\text{C}$. After a week, a mixture of crystals and powder of **11** were recovered by filtration and dried under vacuum (26.6 mg, 18 μmol , 45% yield). Single crystals suitable for an X-ray diffraction study were obtained by layering pentane onto a saturated C_6D_6 solution of **11** before storage at $-40\text{ }^{\circ}\text{C}$ (33% yield).

11: $^1\text{H NMR}$ (400 MHz, C_6D_6) δ : 6.90–6.86 (m, 4H), 6.83–6.79 (m, 4H), 2.10–1.87 (m, 16H), 1.70 (ddd, $J = 11.1, 9.2, 5.5\text{ Hz}$, 8H), 1.40–1.23 (m, 24H), 1.15 (quint, $J = 7.4\text{ Hz}$, 24H), 0.82–0.70 (m, 24H), -3.60 (quint, $J_{\text{HW}} = 18.1\text{ Hz}$, $J_{\text{HP}} = 29.2\text{ Hz}$, 2H). $^{31}\text{P}\{^1\text{H}\}$ NMR (162 MHz, C_6D_6) δ : 43.8 ($J_{\text{PW}} = 286\text{ Hz}$). $^{11}\text{B NMR}$ (128 MHz, C_6D_6) δ : 5.4 (s).

IR (ATR) $\nu/\text{cm}^{-1} = 2958, 2900, 2839, 2272, 1604, 1503, 1457, 1378, 1362, 1335, 1274, 1188, 1163, 1124, 1095, 1081, 1029, 982, 867, 802, 755, 732, 710, 693, 686, 661$.

Elem. Anal. calcd. for $\text{C}_{52}\text{H}_{106}\text{B}_2\text{N}_4\text{O}_4\text{P}_8\text{W}_2$: C, 41.96; H, 7.18; N, 3.76. Found: C, 42.20; H, 7.40; N, 3.70.

Conflicts of interest

The authors declare no conflict of interest.

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