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A neutral, acyclic, borataalkene-like ligand for group 11 metals: L- and Z-type ligands side by side†‡

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The overall neutral α -borylated phosphorus ylide $\text{Ph}_3\text{PC}(\text{Me})\text{BET}_2$ behaves like a polar borataalkene and can act as acyclic, ambiphilic π -type ligand with L- and Z-type functionalities side by side. In the complexes $[\text{MX}\{\eta^2\text{-Ph}_3\text{PC}(\text{Me})\text{BET}_2\}]$ ($\text{M} = \text{Cu}, (\text{Ag}), \text{Au}; \text{X} = \text{Cl}, \text{NTf}_2$), the bonding is dominated by the highly nucleophilic ylidic carbon atom (L-type ligand). The Lewis acidic boron atom furnishes nonetheless a small but significant bonding contribution (Z-type ligand).

Boron-based ligands for transition metal complexes have received considerable attention for many years.¹ Among others, one focus of interest refers to ambiphilic chelate ligands featuring an L-type donor entity and a Z-type acceptor unit suitable to stabilize metal \rightarrow boron interactions.² Boron-based π -ligands have also been studied, most often cyclic structures.³ Acyclic boron-based π -ligands,⁴ however, are relatively scarce and only a few transition metal complexes have been described.⁵ Interests in these types of ligands are rooted in their isoelectronic relationship to olefin ligands (Scheme 1),⁶ which have been studied in great detail during the past decades and are important intermediates in catalytic processes or serve as leaving groups in starting materials and as steering ligands in homogeneous catalysis, for instance.⁷ An inherent feature of anionic borataalkenes⁸ is their flexibility to potentially act as a “classical” π -ligand according to the well-known Dewar–Chatt–Duncanson (DCD) model^{6,7} or as ambiphilic L-/Z-type ligand (Scheme 1).⁹

Piers and co-workers presented in 1999 the first example of an electron-poor, “early” Ta(0) complex featuring an anionic,

C_6F_5 -substituted borataalkene ligand (**I** in Scheme 1), which was synthesized by converting a Ta-bound alkylidene into the η^2 -coordinated $[\text{H}_2\text{C}=\text{B}(\text{C}_6\text{F}_5)_2]^-$ ligand.¹⁰ Quantum chemical calculations predicted that the latter species readily converts to boryl-substituted methyl ligand, most likely due to steric congestion. In 2020, Erker succeeded in isolating complexes of “late”, *i.e.* electron-rich metals bearing an anionic P/C=B chelate ligand (**II**).¹¹ Neutral heteroelement analogs of η^2 -bound olefins are, however, very rare. To the best of our knowledge, only one example has been described in the literature. Amgoune and Bourissou described the synthesis and full characterization of the platinum complexes $[(\text{Ph}_3\text{P})_2\text{Pt}(\eta^2\text{-}\{\text{R}_2\text{P}=\text{B}(\text{C}_6\text{F}_5)_2\})]$ (**III**; $\text{R} = \text{tBu}, \text{Cy}$) and performed an in-depth quantum-chemical analysis.¹² Although the BP-based π -ligand is bound symmetrically, the polar electronic profile of the ligand is also reflected in its partially asymmetric bonding mode featuring characteristics of both a DCD-type π -ligand and an ambiphilic L-/Z-type ligand.

In this context, we recently reported on a highly polarized π -bond¹³ consisting of a ylide carbon atom and a boron atom within the α -borylated phosphorus ylide $\text{Ph}_3\text{PC}(\text{Me})\text{BET}_2$



Scheme 1 Isoelectronic relationship between olefins and heteroalkenes and selected complexes consisting of anionic and neutral, acyclic boron-based π -ligands from the literature. Scope of this work.

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Scheme 2 Treatment of **L1** with various group 11 precursors to furnish the complexes **1–5**.

(**L1**, Scheme 2).¹⁴ As exemplified in small molecules activation, the competition for the lone-pair of electrons between the boryl (BR₂) and phosphonyl (PR₃) substituents results in electronic frustration within this overall neutral, highly polarized borataalkene-like molecule. We now hypothesized that species **L1** would be a promising candidate to act as an acyclic, ambiphilic π -type ligand with L- and Z-type functionalities side by side.¹⁵ Note that a cyclic analog possessing the 9-borataphenanthrene anion has been recently published by Martin and co-workers.¹⁶ Reported herein is the successful realization of our hypothesis by the synthesis and (full) characterization of electron-rich, “late” transition metal complexes of **L1** and a detailed analysis of their bonding situation.

We were able to synthesize the anticipated group 11 complexes [MCl{ η^2 -Ph₃PC(Me)BET₂}] (M = Cu: **1**; M = Au: **2**) either by reacting CuCl with **L1** in CH₂Cl₂ (74% isolated yield) or *via* the substitution of the CO in [AuCl(CO)] with **L1**, but only in poor yields (no reaction with [Ph₃PAuCl]). To our delight, when the gold(i) complex [AuCl(dms)] (dms = dimethyl sulfide) is used as a precursor, however, an isolated yield of 91% for **2** could be achieved (Scheme 2). The mixture of **L1** with AgCl in 1,2-Cl₂C₂H₄ at 60 °C showed no reaction due to insolubility of AgCl.

Suitable single crystals of **2** for X-ray diffraction were obtained by slow solvent evaporation of a saturated CH₂Cl₂ solution. The title compound crystallizes in the space group *P2*₁/*c* (Fig. 1). Most notably, the P1–C1 (178.2 pm) and the C1–B1 bond (155.2 pm) in **2** are considerably elongated as compared to the free pro-ligand **L1** (171.7 pm and 149.2 pm, respectively). This clearly shows that the electron density from the ylidic carbon atom is involved in the C1–Au bond and, thus,



Fig. 1 Molecular structure of **2**, thermal ellipsoids drawn at the 30% probability level. (structural parameters of the second molecule in the asymmetric unit are given in square brackets). Selected bond lengths (pm) and angles (°): Au1–C1 211.3(6), Au1–B1 239.4(8) [241.8(8)], Au1–Cl1 227.74(18), P1–C1 178.2(7), C1–B1 155.2(11), C1–C2 155.2(10); P1–C1–Au1 109.0(3), B1–C1–Au1 80.0(4), C1–Au–Cl1 178.2(2) [176.5(2)].

is no longer available for both π -interactions with $\sigma^*(\text{PC})$ or $p(\text{B})$ orbitals. The gold atom is coordinated in a linear fashion by the ylidic carbon atom and the chloride ligands ($\angle \text{C1–Au1–Cl1} = 178.2^\circ$). The C1–Au1–Cl1 axis is slightly tilted towards the boron atom ($\angle \text{B1–C1–Au1} = 80.0^\circ$) indicating bonding interaction between the boron center and the gold atom. The short B–Au distance of 239.4 pm found in the solid-state is a strong indication for a Au \rightarrow B interaction. The distance is in the same region found for gold complexes of polydentate B/P ligands.¹⁷

The ¹H NMR spectra of **1** and **2** in d₈-toluene display line broadening at room temperature due to dynamic processes, in particular for the ethyl groups bound to boron. Variable temperature NMR investigations revealed energy barriers of ~ 16 (for **1**) and ~ 15 (for **2**) kcal mol^{−1} (for further details, see section S5, ESI[†]). We assumed that this dynamic process is associated with the rotation around the C_{ylide}–B bond. Indeed, density functional theory (DFT) calculations at the dispersion-corrected BP86-D3/def2-TZVPP level (ESI[†]) indicate that the activation barrier associated with the C_{ylide}–B rotation in **1** and **2** is 17.5 and 17.8 kcal mol^{−1}, respectively. Both values agree reasonably well with the experimental ones (Fig. S1, ESI[†]). Interestingly, the rotation along the C_{ylide}–B bond has a dramatic impact on the bonding situation in the corresponding transition state **TS**. As depicted in Fig. S1 (ESI[†]), the relatively short B–Au distance of 241.4 pm in **2** is significantly longer in **TS** (314.3 pm), thus indicating that the Au \cdots B interaction in the latter saddle-point can be considered as negligible.

Furthermore, the ¹¹B NMR signal of **2** at a chemical shift of $\delta(^{11}\text{B}) = 53.6$ ppm shifted upfield compared to the free pro-ligand **L1** (*cf.* $\delta(^{11}\text{B}) = 56.5$ ppm).¹⁸ This is reminiscent of the ¹¹B highfield shift of the boron–gold complexes featuring Au \rightarrow B interactions (Chart S1, ESI[†]).¹⁷

To check the accessibility of other group 11 complexes, we also prepared the whole series of {M(NTf₂)} complexes **3–5** (**3**: Cu, **4**: Ag, **5**: Au; Scheme 2). It turned out, however, that only highly sensitive, oily products are formed. Due to this, a full characterisation was not possible. We have nevertheless performed comprehensive NMR studies and the results are convincing (*e.g.* ¹¹B NMR chemical shifts of $\delta(^{11}\text{B}) = 53.3$ ppm for **3**, 58.5 ppm for **4**, 53.8 ppm for **5**, respectively, and $\delta(^{31}\text{P}) = 31.3$ ppm (d, ²J_{AgP} = 27,8 Hz) for **4** in d₈-toluene).

Again, line broadening at room temperature was observed for all complexes due to the dynamic process associated with the rotation around the C_{ylide}–B bond. Variable temperature NMR investigations on **3–5** revealed slightly lower energy barriers of ~ 15 (for **3** and **4**) and ~ 14 (for **5**) kcal mol^{−1}.

To further investigate the bonding situation in the newly prepared complexes, we first explored the electronic structure of the pro-ligand **L1** with the help of DFT calculations. From the computational studies it becomes clear that pro-ligand **L1** is best described by the zwitterionic resonance structure (Fig. S2, ESI[†]) with a negative charge on the carbon atom (*i.e.* ylide) and only to a lesser extent by the borataalkene structure. This is confirmed by comparing the charges and Wiberg Bond Indices of related species (see Table S1, ESI[†]). Indeed, the computed HOMO can be viewed as a lone-pair at the carbon atom, which



Table 1 Bonding between MX and Ph₃PC(Me)BR₂ in [MX{η²-Ph₃PC(Me)BR₂}] complexes (M = Cu, Ag, Au; X = Cl, NTf₂) (EDA-NOCV, values in kcal mol⁻¹), computed at the ZORA-BP86-D3/TZ2P//BP86-D3/def2-TZVPP level

	[MCl{η ² -Ph ₃ PC(Me)BR ₂ }]			[(MNTf ₂){η ² -Ph ₃ PC(Me)BR ₂ }]			[AuCl{η ² -Ph ₃ PC(Me)BR ₂ }]	
	M = Cu	M = Ag	M = Au	M = Cu	M = Ag	M = Au	R = CF ₃	R = OMe
ΔE _{int}	-72.6	-57.2	-79.4	-91.4	-74.4	-101.8	-73.5	-80.6
ΔE _{Pauli}	124.1	120.6	182.0	130.0	131.5	204.0	192.8	175.4
ΔV _{elstat}	-127.6	-120.5	-173.8	-134.0	-129.2	-194.0	-170.7	-171.7
ΔE _{orb}	-56.0	-46.5	-75.0	-62.6	-54.4	-88.4	-83.0	-71.4
ΔE _{orb} (ρ ₁)	-24.6	-22.4	-40.0	-27.6	-25.6	-47.6	-37.7	-41.4
ΔE _{orb} (ρ ₂)	-13.6	-8.4	-14.9	-12.3	-8.7	-14.5	-25.3	-9.9
ΔE _{orb} (rest)	-17.8	-15.7	-20.1	-22.7	-20.1	-26.3	-20.0	-20.1
ΔE _{disp}	-13.1	-10.8	-12.5	-24.9	-22.3	-23.4	-12.5	-12.8
r(M-C)/Å	1.979	2.178	2.132	1.971	2.175	2.100	2.144	2.126
r(M-B)/Å	2.318	2.556	2.414	2.322	2.501	2.447	2.258	2.589

is slightly delocalized into the adjacent vacant p_z atomic orbital of boron (see below). The latter finding is reflected in the computed C-B Wiberg bond Index of 1.24, which suggests a partial, highly polarized C=B bond.

More quantitative insight into the bonding situation of the {MCl} complexes (M = Cu, Ag, Au) can be gained by means of the energy decomposition analysis (EDA) method computed at the relativistic ZORA-BP86-D3/TZ2P//BP86-D3/def2-TZVPP level. From the data in Table 1, columns 2–4, it becomes evident that the main contribution to the bonding between **L1** and {MCl} comes from the electrostatic interactions (measured by the ΔV_{elstat} term), which contribute *ca.* 65% to the total interaction (ΔE_{int}). This is consistent with the zwitterionic nature of pro-ligand **L1** commented above. Despite that, the orbital interactions (ΔE_{orb}) are also significant, albeit to a lesser extent (*ca.* 30%). In contrast, interactions coming from dispersion forces (ΔE_{disp}) are much less important in the **L1**...{MCl} interaction (*ca.* 5%).

The natural orbital for chemical valence (NOCV) extension of the EDA method was applied next to further partitioning the orbital term into its main pair-wise interactions. According to the EDA-NOCV, two main orbital interactions dominate the total ΔE_{orb} term, namely the donation from the lone-pair at the carbon atom to the σ*(M-Cl) and the π-backdonation from a doubly-occupied d atomic orbital at the transition metal to the vacant p_z atomic orbital of the boron atom, denoted as ρ₁ and ρ₂, respectively (see Fig. 2). The σ-donation LP(C)→σ*(M-Cl) (ρ₁) is significantly stronger than the π-backdonation (ρ₂). When inspecting the influence of the coordinated group 11 metal, it becomes clear that M = Au leads to stronger interactions in all contributions and that the (hypothetical) silver complexes exhibit the lowest values when compared to Au or Cu complexes. This, however, is consistent with recent computations of Bayat *et al.*^{19a} on phosphorus ylide complexes of group 11 metals and recent bonding analyses of related Au(I) complexes.^{19b} This also correlates with the highest promotion energy for Ag(I) in the series Cu(I) (8.25 eV) < Ag(I) (9.94 eV) > Au(I) (7.83 eV) and the lowest electron affinity in the series Cu(I) (7.72 eV) > Ag(I) (7.59 eV) < Au(I) (9.22 eV). Moreover, the Au→p_z(B) backdonation is comparable to that found in those complexes having a single anchor boron ligand (Chart S1, ESI[†]), which were confirmed to exhibit a significant Au→B



Fig. 2 Deformation densities and associated molecular orbitals of the most important orbital interactions of **2** using the fragments AuCl and **L1** (Table 1) together with the corresponding stabilizing energies ΔE_{orb}(ρ₁) and ΔE_{orb}(ρ₂). The colour code used to represent the flow of charge is red→blue.

interaction.²⁰ This backdonation is also validated by the NBO method, which provides a noticeable Au→B WBI of 0.25 and locates a significant stabilizing delocalization from the doubly-occupied d-atomic orbital of gold to the p_z-atomic orbital of boron (associated ΔE⁽²⁾ = -14.2 kcal mol⁻¹). Therefore, our EDA-NOCV analysis indicates that the orbital interactions in **2** can be described as two donor-acceptor interactions involving the highly nucleophilic ylidic carbon atom as the dominant ligating atom (L-type ligand) and the Lewis acidic boron atom, which furnishes a weaker yet noticeable bonding contribution as a Z-type ligand. Results above suggest that the Au→p_z(B) backdonation can be efficiently modulated by modifying either the geometry or the Lewis acidity of the BR₂ moiety. For instance, in the rotated **TS** (Fig. S1, ESI[†]), the Au-B bond distance is significantly longer than in that in **2** as a consequence of the position of the p_z atomic orbital of boron, which is much less available to accept electron density from the gold atom. As a result, this saddle-point becomes destabilized with respect to **2** and exhibits an almost negligible Au→p_z(B) π-backdonation (ΔE_{orb}(ρ₂) < -1 kcal mol⁻¹).



To check the influence of the Lewis acidity of the BR₂ fragment, we attached both a strong electron-withdrawing (CF₃) and electron-donor (OMe) groups directly to the boron atom. From the data in Table 1, columns 7 and 8, the replacement of the ethyl groups by CF₃ significantly enhances the Au→p_z(B) π-backdonation, which becomes almost twice as strong as that in the parent **2** ($\Delta E_{\text{orb}}(\rho_2) = -25.3 \text{ kcal mol}^{-1}$). This is reflected in a markedly shorter Au–B bond (225.8 pm). At variance, the presence of the donor methoxy groups provokes the opposite effect and the Au→p_z(B) π-backdonation becomes weaker ($\Delta E_{\text{orb}}(\rho_2) = -9.9 \text{ kcal mol}^{-1}$) whereas the corresponding Au–B is much longer (258.9 ppm). We also studied the effect of the M-bound anion and compared {MCl} with {M(NTf₂)} (cf. Table 1, columns 5–7). When changing X[−] = Cl[−] to (NTf₂)[−], the interaction between the {MX} and the ligand **L1** becomes clearly stronger. This is mainly due to a significant increase in the orbital interactions (the electrostatic term also increases albeit to a lesser extent). In particular, the LP(C)→σ*(M–X) (ρ₁) interaction becomes stronger because the M(NTf₂) fragment is a much better acceptor as compared to {MCl}.²⁰ For this same reason, the Au→p_z(B) π-backdonation is slightly weaker.

In summary, this case study clearly depicts that the polar π-bond of the α-borylated phosphorous ylide differs significantly from a C=C π-bond and is even more polar than the phosphinoborane published by Amgoune and Bourissou.¹² In the {MCl} complexes **1** and **2**, the highly nucleophilic ylidic carbon atom is the dominant ligating atom (L-type ligand). The Lewis acidic boron atom furnishes a small but significant bonding contribution as a Z-type ligand. Thus, the bonding is asymmetric from both the geometric and the electronic perspectives. The overall neutral, acyclic pro-ligand **L1** indeed behaves like a polar borataalkene ligand, which is induced by the competition for the lone-pair of electrons. Quantum chemical calculations predict that the strength of the Au→B interaction can be increased by introducing electron-withdrawing groups and decreased by π-donating substituents on boron. We also provided experimental and computational evidence that the L-type ligand contribution can be increased upon moving from X[−] = Cl[−] to (NTf₂)[−]. Thus, it appears that the whole continuum between ambiphilic L-/Z-type coordination and DCD-type π-coordination can be covered by fully exploring the coordination chemistry of variously substituted α-borylated phosphorous ylides. Investigations in this direction are currently performed in our laboratories.

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Conflicts of interest

There are no conflicts to declare.

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