

## COMMUNICATION

## Phosphinoborinium cation: a synthon for cationic B–P bond systems

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Accepted 00th January 20xxKinga Kaniewska-Laskowska,<sup>a</sup> Katarzyna Klimsiak,<sup>a</sup> Natalia Szykiewicz,<sup>a</sup> Jarosław Chojnacki<sup>a</sup> and Rafał Grubba<sup>\*a</sup>

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Herein, we report access to phosphinoborinium cations via heterolytic cleavage of the boron-bromide bond in bromophosphinoborane. The product of the reaction was isolated as a dimeric dication possessing a planar P<sub>2</sub>B<sub>2</sub> core. Activation of C–H and C–P bonds in the dication led to formation of the borinium-phosphaborene adduct. Reactivity studies revealed that title cation exhibits ambiphilic properties and intramolecular frustrated Lewis pair features.

Due to Lewis acidic properties resulting from the empty p-orbital and coordinative unsaturation of the B atom, tri-coordinate, neutral boron compounds play key roles in nonmetallic catalysis.<sup>[1]</sup> The discovery of frustrated Lewis pairs (FLPs)<sup>[2]</sup> has stimulated rapid development of the chemistry of triorganoboranes,<sup>[3]</sup> which are most commonly the Lewis acidic components of FLPs capable of activating strong chemical bonds in simple inorganic and organic molecules.<sup>[4–6]</sup>

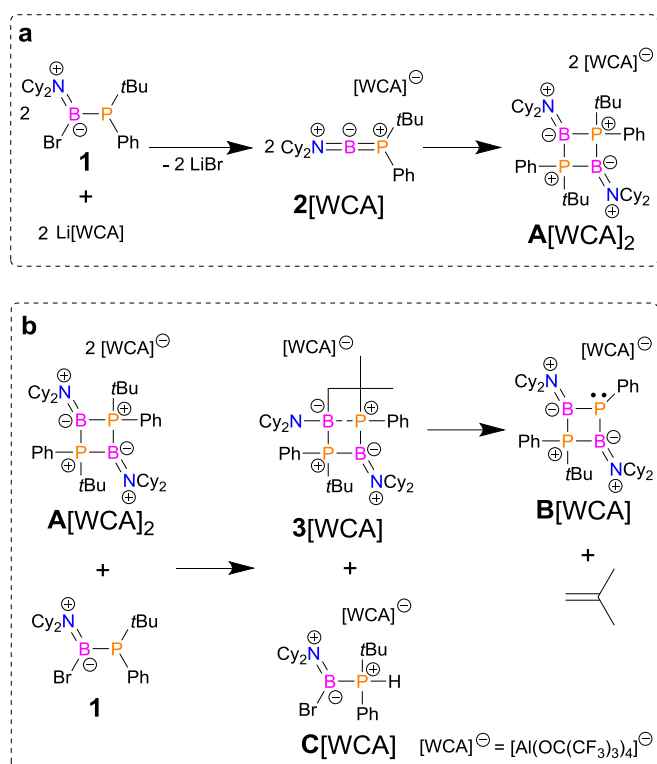
Low-coordinate boronic cations are even more electron deficient than neutral boron Lewis acids. Generally, three main types of boronic cations can be distinguished by taking into account the coordination number of boron: borinium [R<sub>2</sub>B]<sup>+</sup>, borenium [R<sub>2</sub>BL]<sup>+</sup>, and boronium [R<sub>2</sub>BL<sub>2</sub>]<sup>+</sup> ions.<sup>[7,8]</sup> The first two are attractive synthetic targets owing to their strong Lewis acidic properties. In the past, borinium and borenium cations were only considered chemical curiosities; however, the development of modern synthetic methods allowed isolation and characterization of these species in the condensed phase. Due to the high reactivities of low-coordinate boronic cations, their syntheses are not trivial and require the application of electronic and/or steric stabilization. The first borinium cation was obtained by Nöth and coworkers,<sup>[9,10]</sup> and electron-donating amino groups were introduced at the B-centre. Another milestone in this area was the synthesis and isolation of [Mes<sub>2</sub>B]<sup>+</sup> by the Shoji group.<sup>[11],[12],[13]</sup> These results showed

that the presence of π-donating ligands was not necessary for stabilization of the borinium cations. Stephan and coworkers used two phosphoraminate ligands for stabilization of boron centres with a low coordination numbers<sup>[14]</sup>; moreover, they showed that Nöth's and Shoji's cations exhibited high reactivity towards small inorganic and organic molecules.<sup>[15,16]</sup> The recent development of boronic cation chemistry confirms the applicability of these reactive species in stoichiometric reactions and as catalysts for hydrogenation, hydrosilylation, hydroboration, C–H bond borylation, and C–C bond hydrogenolysis reactions.<sup>[17–23]</sup>

Our research interests are focused on the chemistry of low-coordinate B–P bond systems such as phosphinoboranes,<sup>[24]</sup> diphosphinoboranes<sup>[25]</sup>, and triphosphinoboranes.<sup>[26]</sup> Despite the presence of one to three B–P bonds, these systems retain Lewis acidic and Lewis basic character of the reactive boron and phosphorus centres and behave as intramolecular FLPs to activate H<sub>2</sub>, CO<sub>2</sub>, other heterocumulenes, SO<sub>2</sub>, N<sub>2</sub>O, and BH<sub>3</sub>.<sup>[24,26–28]</sup> As a continuation of our previous studies, we decided to introduce this research concept to cationic boron compounds and synthesize low-coordinate boronic cations possessing B–P functionality.

As a precursor to the phosphinoborinium cation, we utilized Cy<sub>2</sub>N(Br)BPtBuPh (**1**), which was obtained via a simple two-step reaction of BBr<sub>3</sub> with Cy<sub>2</sub>NH followed by addition of tBuPhPLi. In a recent study, we used analogous amino(bromo)phosphinoboranes for syntheses of diphosphinoboranes with mixed phosphanyl substituents.<sup>[25]</sup> Spectroscopic and X-ray diffraction data for **1** are presented in the ESI. As a source of weakly coordinating anions (WCAs), we chose Krossing's salt Li[Al(OC(CF<sub>3</sub>)<sub>3</sub>)<sub>4</sub>] due to its synthetic accessibility and chemical inertness.<sup>[29]</sup> According to <sup>31</sup>P and <sup>11</sup>B spectroscopic analyses, an equimolar reaction of **1** with the aforementioned salt in dichloromethane led to formation of three ionic species, A[WCA]<sub>2</sub>, B[WCA], and C[WCA], in a molar ratio ≈ 1:1:1; this was accompanied by precipitation of LiBr (Scheme 1).

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Scheme 1. Syntheses of cations  $\mathbf{A}^{2+}$ ,  $\mathbf{B}^+$  and  $\mathbf{C}^+$  (Synthetic details: solvent  $\text{CH}_2\text{Cl}_2$ ; temperature from  $-30^\circ\text{C}$  to  $25^\circ\text{C}$ ; yields:  $\mathbf{A}[\text{WCA}]_2 = 32\%$ ,  $\mathbf{B}[\text{WCA}] = 61\%$ . The bulk purity of  $\mathbf{A}[\text{WCA}]_2$  and  $\mathbf{B}[\text{WCA}]$  was confirmed by elemental analysis).

Additionally, evolution of gas bubbles was observed during the reaction, especially in the case of pressure equalization between the reaction vessel and the argon manifold of the Schlenk line. This gaseous product together with solvent was collected in a liquid nitrogen trap and unambiguously identified as isobutene by  $^1\text{H}$  and  $^{13}\text{C}$  NMR spectroscopy (Figure S41 and S42).

Due to differences in the solubilities of reaction products in hydrocarbons, they can be separated by fractional crystallization. Colourless crystals of  $\mathbf{A}[\text{WCA}]_2$  were deposited from a dichloromethane solution at  $4^\circ\text{C}$ . Collectively, NMR spectroscopic and X-ray diffraction studies revealed that the obtained salt consisted of dication  $\mathbf{A}^{2+}$  (diphosphoniodiborane), which can be seen as the dimer of the monomeric phosphinoborinium cation  $\mathbf{2}^+$  (Scheme 1a). Indeed, a calculation of the Gibbs free energy for dimerization of  $\mathbf{2}^+$  showed that this reaction is strongly thermodynamically favoured ( $\Delta G_{298\text{K}}^0 = -29.7 \text{ kcal}\cdot\text{mol}^{-1}$ ).<sup>[30]</sup> The  $^{11}\text{B}$  and  $^{31}\text{P}\{^1\text{H}\}$  NMR spectra of  $\mathbf{A}^{2+}$  displayed very broad signals at 36.0 ppm and 11.7 ppm, respectively. Due to the broadness of the signals, boron-phosphorus coupling was not observed. The  $^{11}\text{B}$  chemical shift indicated a low coordination number for the boron atom. Moreover, the  $^{19}\text{F}$  and  $^{27}\text{Al}$  NMR spectra showed characteristic signals for  $[\text{Al}(\text{OC}(\text{CF}_3)_3)_4]$ , confirming that this counterion remained untouched. The molecular structure of  $\mathbf{A}^{2+}$  is presented in Figure 1. The most characteristic structural feature of  $\mathbf{A}^{2+}$  is the planar  $\text{B}_2\text{P}_2$  core. The geometries around the nitrogen and boron atoms are trigonal planar with the sums of angles very close to  $360^\circ$ . The P atoms exhibit distorted tetrahedral geometries.

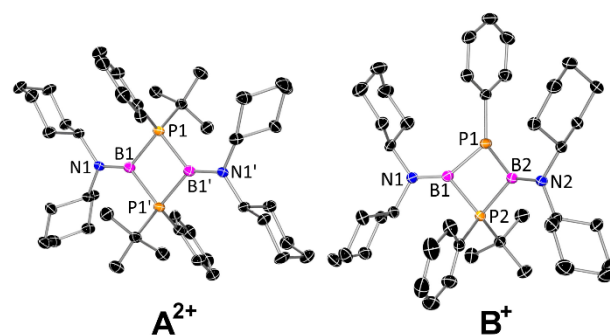


Figure 1. X-ray structures of cations  $\mathbf{A}^{2+}$  and  $\mathbf{B}^+$ . The counterions and H-atoms are omitted for clarity. The thermal ellipsoids are shown at the 50% probability level.

The relatively short N1-B1 distance of  $1.377(6) \text{ \AA}$  together with the planar geometry around the N1 and B1 atoms indicates significant  $\pi$ -interactions occurring between these atoms. The B1-P1 ( $1.979(4) \text{ \AA}$ ) and B1'-P1' ( $1.981(5) \text{ \AA}$ ) distances are in the typical range for single B-P bonds ( $1.96 \text{ \AA}$ ).<sup>[31]</sup> The tBu and Ph substituents are located in *trans* positions with respect to the planar  $\text{B}_2\text{P}_2$  ring. Interestingly, signals attributable to the *cis* isomer were found in the NMR spectra of isolated crystals of  $\mathbf{A}[\text{WCA}]_2$ . The molar ratio of the *trans* and *cis* isomers was calculated as 84:16 based on integration of the well-separated CH signals for Cy groups in the  $^1\text{H}$  NMR spectrum. Furthermore, DFT calculations showed that the *trans* isomer has a lower energy than the *cis* isomer, and the free energy difference is  $-7.4 \text{ kcal}\cdot\text{mol}^{-1}$ .<sup>[30]</sup>

The electronic structure of  $\mathbf{A}^{2+}$  was investigated by theoretical methods.<sup>[32]</sup> Natural population analysis (NPA) revealed high positive charges located at phosphorus and boron atoms of the central planar core with values of  $+0.866$  and  $+0.512$ , respectively. The B-N bonds are significantly polarized with a high negative charge located at the nitrogen atom ( $-0.741$ ). The Wiberg bond index (WBI) calculated for B1-P1 and B1'-P1' ( $0.904$ ,  $0.914$ ) confirmed the single bond character of these bonds. The WBI for B1-N1/B1'-N1' was  $1.196$ , which is consistent with the short boron-nitrogen distance and partial multiple bond character.<sup>[32]</sup> The LUMO of  $\mathbf{A}^{2+}$  is delocalized over boron and phosphorus atoms and is composed of two lobes separated by a nodal plane corresponding to the  $\text{B}_2\text{P}_2$  plane. The HOMO of  $\mathbf{A}^{2+}$  exhibits electron density localized between the B and P atoms ( $\sigma$ -bonding) with a contribution from electron density localized on phenyl substituents (Figure S54).<sup>[32]</sup>

To our knowledge, there are no reports of planar dicationic  $\text{B}_2\text{P}_2$  cyclic systems containing tri-coordinate boron and four-coordinate phosphorus atoms. Analogous planar  $\text{B}_2\text{P}_2$  cores can be found only in very few neutral species, such as the stable diradical  $(\text{tBuBP}(\text{Pr}_2))_2$  reported by the Bertrand group.<sup>[33]</sup> Very recently, Braunschweig and coworkers described the synthesis of a neutral, planar NHC-stabilized  $\text{B}_2\text{P}_2$  ring via a reaction of diboryne  $\text{B}_2(\text{SIDep})_2$  (SIDep=1,3-bis(2,6-diethylphenyl)imidazolin-2-ylidene) with a diphosphane  $(\text{Et}_2\text{P})_2$ .<sup>[34]</sup> Oxidation of the resulting compound led to formation of a dication possessing a butterfly structure and a B-B bond,<sup>[34]</sup> which is in strong contrast with the structural features of  $\mathbf{A}^{2+}$  described above.

After separation of the  $\mathbf{A}[\text{WCA}]_2$  crystals, the dichloromethane solution was layered with pentane and stored at +4 °C. Diffusion crystallization gave large, colourless crystals of  $\mathbf{B}[\text{WCA}]$ . According to NMR spectroscopic and X-ray diffraction studies,  $\mathbf{B}^+$  can be described formally as an adduct of borinium cation  $\mathbf{2}^+$  and the phosphaborene  $\text{PhPBNiPr}_2$ . Cooling the mother liquor to -20 °C gave the mixture of colourless crystals of  $\mathbf{B}[\text{WCA}]$  and  $\mathbf{C}[\text{WCA}]$ . The X-ray study revealed that the latter compound contains a  $\mathbf{C}^+$  borenium cation resulting from protonation of starting  $\mathbf{1}$  (Scheme S3). The presence of the P-H functional group in the structure of  $\mathbf{C}^+$  was confirmed both by analysis of the Fourier density map and  $^1J_{\text{PH}}$  coupling with the expected large absolute value of 346.9 Hz (Figure S40). The  $\mathbf{C}[\text{WCA}]$  decomposes in solution to  $t\text{BuPhPH}$  and  $\text{C}_2\text{NBBR}_2$  which prevented its separation and full characterization (Scheme S4). These experimental observations suggested that  $\mathbf{B}^+$  was formed via the reaction of  $\mathbf{A}^{2+}$  with phosphinoborane  $\mathbf{1}$ . This assumption was additionally confirmed by two independent experiments. First, the reaction of  $\text{Li}[\text{WCA}]$  with an excess of  $\mathbf{1}$  (molar ratio 1:1.5) did not lead to the formation of  $\mathbf{A}^{2+}$  but gave two boronic cations,  $\mathbf{B}^+$  and  $\mathbf{C}^+$ . Similar to the initial, equimolar reaction, precipitation of  $\text{LiBr}$  and evolution of isobutene gas were observed. Second, we added  $\mathbf{1}$  to a solution of the previously isolated crystals  $\mathbf{A}[\text{WCA}]_2$  (molar ratio 1:1), which yielded only  $\mathbf{B}[\text{WCA}]$ ,  $\mathbf{C}[\text{WCA}]$ , and isobutene. Considering all of these experimental observations, we propose a plausible reaction path leading to cation  $\mathbf{B}^+$  (Scheme 1b). In the first step,  $\mathbf{A}^{2+}$  is deprotonated by  $\mathbf{1}$ , which leads to formation of short-lived intermediate  $\mathbf{3}^+$  and borenium cation  $\mathbf{C}^+$ . The proposed structure for  $\mathbf{3}^+$  derived from theoretical studies shows that deprotonation of one methyl group of  $\mathbf{A}^{2+}$  leads to formation of a new bond between the deprotonated C atom and the electrophilic B atom and a significant weakening of one B-P bond (Figure S58). In the next step,  $\mathbf{3}^+$  eliminates isobutene as a result of C-B and C-P bond activation, yielding  $\mathbf{B}^+$ . DFT calculations revealed that the first step of this reaction, which involved the formation of unstable  $\mathbf{3}^+$ , is only slightly endergonic, whereas the second step is strongly exergonic; the  $\Delta G_{298\text{K}}^0$  values were 2.4 kcal·mol<sup>-1</sup> and -24.6 kcal·mol<sup>-1</sup>, respectively.<sup>[30]</sup> Very recently, Stephan and coworkers found that the borinium cation  $[\text{B}(\text{N}(\text{SiMe}_3)_2)_2]^+$  can act as a source of a proton,<sup>[35]</sup> which is in line with our results.

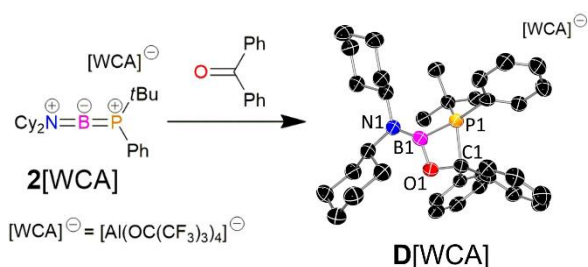
The unprecedented structure of  $\mathbf{B}^+$  was investigated by spectroscopic, diffraction, and theoretical methods. The  $^{31}\text{P}\{\text{H}\}$  NMR spectrum of  $\mathbf{B}^+$  consisted of two broad doublets at 6.5 ppm and -17.3 ppm with large  $^2J_{\text{PP}}$  coupling (391 Hz), which corroborated the presence of two inequivalent P-centres. Similar to  $\mathbf{A}^{2+}$ , only one broad signal was visible in the  $^{11}\text{B}$  spectrum of  $\mathbf{B}^+$ , and it exhibited a chemical shift characteristic of a tri-coordinate boron centre (35.1 ppm). The solid-state structure of  $\mathbf{B}^+$  is depicted in Figure 1. In contrast to the structure of  $\mathbf{A}^{2+}$ , the P1 atom is tri-coordinate and bonded to the B1, B2, and C1 atoms of the phenyl group. Otherwise, the P2 atom retained the tetrahedral geometry, as observed for the phosphorus atoms in  $\mathbf{A}^{2+}$ . Interestingly, the geometry around P1 exhibited a high degree of planarity, and the angles summed to 349.38°. Furthermore, the structure of  $\mathbf{B}^+$  exhibited relatively

short B1-P1 and B2-P1 distances with values of 1.881(2) Å and 1.882(2) Å, respectively. The flattened geometry at the P1 atom together with the significant shortening of the B1-P1 and B2-P1 bonds suggested the presence of  $\pi$ -interactions resulting from overlap of the lone pair electrons on the P1 atom with the formally empty  $p$ -orbitals of the B1 and B2 atoms. The N1-B1 (1.376(2) Å), N2-B2 (1.380(2) Å), B1-P2 (1.970(2) Å) and B2-P2 (1.963(2) Å) bond lengths were very similar to the B-N and B-P distances observed for  $\mathbf{A}^{2+}$ . Moreover, a comparison of the  $\mathbf{A}^{2+}$  and  $\mathbf{B}^+$  structures revealed common planar geometries around the boron and nitrogen atoms. The  $\text{B}_2\text{P}_2$  ring in  $\mathbf{B}^+$  was not planar, as observed for  $\mathbf{A}^{2+}$ , but folded along the P1-P2 vector, and the angle between the B1P1B2 and B1P2B2 planes was 155.65°.

DFT calculations of the frontier molecular orbitals of  $\mathbf{B}^+$  revealed that the HOMO and LUMO represented mainly B1-P1-B2  $\pi$ -bonding and  $\pi^*$ -antibonding orbitals, respectively (Figure S55).<sup>[32]</sup> The WBI indexes calculated for B1-P1 (1.167) and B2-P1 (1.176) were in accordance with the partial multiple bond character of these bonds. On the other hand, the single bond character of B1-P2 and B2-P2 was confirmed by WBIs with values of 0.915 and 0.926, respectively. The WBIs for the boron-nitrogen bonds in  $\mathbf{B}^+$  were slightly smaller than that for  $\mathbf{A}^{2+}$  (1.118-1.132 vs. 1.196).<sup>[32]</sup> A NPA analysis showed that all atoms of the  $\text{B}_2\text{P}_2$  ring were positively charged; the highest positive charge was found at the P2 atom (0.977), and significantly lower positive charges were found at P1 (0.379) and the boron atoms (0.307). As with  $\mathbf{A}^{2+}$ , the highest negative charges were located at the nitrogen atoms (-0.733, -0.744).<sup>[32]</sup> These results showed that despite the intrinsic electron deficiency of  $\mathbf{B}^+$ , this cation also has the feature of a Lewis base (the electron pair at the P1 atom).

We tested the applicability of phosphinoborinium cation in the activation of small molecules. The generated *in situ* cation  $\mathbf{2}^+$  reacted with benzophenone giving adduct  $\mathbf{D}[\text{WCA}]$  as only one product (Scheme 2). The  $^{11}\text{B}$  and  $^{31}\text{P}\{\text{H}\}$  NMR spectra of this compound display very broad signals at 29.3 ppm and 68.4 ppm, respectively, which corroborate with the B-P bond and tricoordinate B atom occurring in the structure. The colourless crystals of  $\mathbf{D}[\text{WCA}]$  were isolated from  $\text{CH}_2\text{Cl}_2$  solution at -20 °C. The X-ray diffraction studies shed more light on the structural features of  $\mathbf{D}^+$  (Scheme 2, Figure S50, Table S1). The adduct  $\mathbf{D}^+$  possesses a planar P1-B1-O1-C1 core resulting from electron donor-acceptor interactions between the phosphinoborinium cation and benzophenone. The Lewis basic P1 atom acts as an electron pair donor to the C1 atom of the carbonyl group, whereas the low-coordinate Lewis acidic B1 atom accepts electron pair from the O1 atom of benzophenone. These observations confirmed the ambiphilic properties of phosphinoborinium cation, which exhibits characteristic reactivity for intramolecular frustrated Lewis pairs. Interestingly, in the case of ambiphilic species containing tri-coordinate B atoms, such as phosphinoboranes and diphosphinoboranes, the small molecules are inserted into the B-P bond (except  $\text{H}_2$ ).<sup>[24,27,28,36]</sup> Similarly to the product of the reaction of phosphinoborinium cation with benzophenone, the B-P bond is retained in adducts of neutral phosphaborene

(containing di-coordinate B atom) with ketones, which were further utilized in phospho-bora-Wittig reaction.<sup>[37]</sup>



Scheme 2. Synthesis and X-ray structure of cation **D**<sup>+</sup> (Synthetic details: solvent CH<sub>2</sub>Cl<sub>2</sub>; temperature from -50°C to 25°C; yield 40%. The bulk purity of **D**[WCA] was confirmed by elemental analysis. The counterion and H-atoms are omitted for clarity. The thermal ellipsoids are shown at the 50% probability level).

In conclusion, we have provided a simple method for generation of phosphinoborinium cations utilizing synthetically accessible bromophosphinoboranes. The obtained borinium cations can act as building blocks for cationic frameworks possessing B-P functionality. The presented results open a new chapter in the chemistry of low-coordinate boronic cations by giving access to species that combine the features of both Lewis acids and Lewis bases. Having a simple method for syntheses of phosphinoborinium cations available, we are currently working on applications of these species in stoichiometric and catalytic transformations that allow activation of strong chemical bonds. Financial support of these studies from Gdańsk University of Technology by the DEC-3/2020/IDUB/III.4.1/Tc grant under the TECHNITIUM and by the DEC-2/2021/IDUB/V.6/Si grant under the SILICIUM - 'Excellence Initiative - Research University' program is gratefully acknowledged. N. S. thanks the TASK Computational Centre and PLGrid Infrastructure for access to computational resources.

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