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# [(Cp<sub>2</sub>M)<sub>2</sub>B<sub>9</sub>H<sub>11</sub>] (M = Zr or Hf): early transition metal 'guarded' heptaborane with strong covalent and electrostatic bonding†‡

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Among the series of stable *closo*-borate dianions, [B<sub>n</sub>H<sub>n</sub>]<sup>2-</sup>, the X-ray crystallographic structure of [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup> was determined only in 2011. To explore its chemistry and stability, we have isolated and structurally characterized two new transition metal complexes of the heptaborane, [(Cp<sub>2</sub>M)<sub>2</sub>B<sub>9</sub>H<sub>11</sub>] (Cp = η<sup>5</sup>-C<sub>5</sub>H<sub>5</sub>; M = Zr or Hf). The structures of [(Cp<sub>2</sub>M)<sub>2</sub>B<sub>9</sub>H<sub>11</sub>] contain a pentagonal bipyramidal B<sub>7</sub> core, coordinated by two {Cp<sub>2</sub>M} and two {BH<sub>2</sub>} units equatorially. Structural and spectroscopic characterizations and DFT calculations show that [(Cp<sub>2</sub>M)<sub>2</sub>B<sub>9</sub>H<sub>11</sub>] complexes are substantially more stable than the parent dianion, in either [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup> or (nBu<sub>4</sub>N)<sub>2</sub>[B<sub>7</sub>H<sub>7</sub>]. Our theoretical study and chemical bonding analyses reveal that the surprising stability of the two new heptaborane metal complexes is due to multi-center covalent bonds related to the two *exo*-{Cp<sub>2</sub>M} units, as well as electrostatic interactions between the {Cp<sub>2</sub>M} units and the B<sub>7</sub> core. The facile syntheses of the heptaborane metal-complexes will allow further exploration of their chemistry.

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## 1. Introduction

Since the early work of Alfred Stock on borane chemistry,<sup>1</sup> syntheses and characterizations of polyhedral borane compounds, especially with higher vertex *closo* geometries, have attracted significant interest.<sup>2,3</sup> As a result, the syntheses and structural characterizations of *closo*-borate compounds with the general formula [B<sub>n</sub>H<sub>n</sub>]<sup>2-</sup> (*n* = 6–12) have been documented in detail.<sup>4,5</sup> After the first isolation of the *closo*-borate dianion [B<sub>10</sub>H<sub>10</sub>]<sup>2-</sup> by Hawthorne in 1959, other members in the series [B<sub>n</sub>H<sub>n</sub>]<sup>2-</sup> (*n* = 6–12) were discovered successively.<sup>6</sup> Interest in the *closo*-polyhedral borane compounds was stimulated by the first crystal structure report of icosahedral [K<sub>2</sub>(B<sub>12</sub>H<sub>12</sub>)] by Lipscomb in 1960.<sup>7</sup> Interestingly, although extensive research had been carried out, X-ray crystallographic study of the borate

dianion [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup> was still missing at that time, making the [B<sub>n</sub>H<sub>n</sub>]<sup>2-</sup> series incomplete. The only *closo*-heptaborate compound, [B<sub>7</sub>Br<sub>7</sub>]<sup>2-</sup>, was not known until the report of the crystal structure of [(C<sub>5</sub>H<sub>5</sub>N)<sub>2</sub>CH<sub>2</sub>][B<sub>7</sub>Br<sub>7</sub>].<sup>8</sup> It should be noted that the parent [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup> was partially characterized in Cs<sub>2</sub>[B<sub>7</sub>H<sub>7</sub>] using <sup>11</sup>B NMR and it was found to be the least stable polyhedral ion.<sup>9</sup> The reported yield of [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup> synthesized from NaBH<sub>4</sub> after several steps was only 0.008%. As a result, an improved synthesis and complete structural characterization of [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup> were of general interest. Although a number of theoretical studies on the structure and bonding of [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup> have been reported,<sup>10</sup> its convenient synthesis and full structural characterization were established only recently by Bernhardt in 2011.<sup>11</sup> In an effort to synthesize the neutral *hypercloso*-heptaborane analogue, Siebert reported [B<sub>7</sub>Cl<sub>5</sub>(PMe<sub>3</sub>)<sub>2</sub>], which was mainly characterized using <sup>11</sup>B NMR.<sup>12</sup>

The continuous efforts into the development of group 4 metallocenes, both by the authors and others,<sup>14</sup> allowed us to isolate and structurally characterize [(Cp<sub>2</sub>Zr)<sub>2</sub>B<sub>5</sub>H<sub>11</sub>] (**I**) from the reaction of [Cp<sub>2</sub>Zr(BH<sub>4</sub>)<sub>2</sub>] with [BH<sub>3</sub>·THF] recently. The single crystal X-ray structural analysis of **I** revealed an open geometry with a planar B<sub>3</sub> ring. Structural and bonding analyses demonstrated that **I** indeed is a neutral triborane analogue. In addition, theoretical results revealed the intriguing role of early transition metal fragments {Cp<sub>2</sub>Zr} in the stabilization of the borane core. The synthesis of **I** paved the way for accessing group 4 metallocenes readily utilizing the well-known borohydride complexes, [Cp<sub>2</sub>M(BH<sub>4</sub>)<sub>2</sub>] (M = Ti, Zr or Hf).<sup>15</sup> These

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‡ Electronic supplementary information (ESI) available: Detailed synthesis and characterizations; mass spectrometry details; NMR and IR spectral details; X-ray analysis details; bonding analyses and computational details. CCDC 1568096–1568098. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c7sc05014c

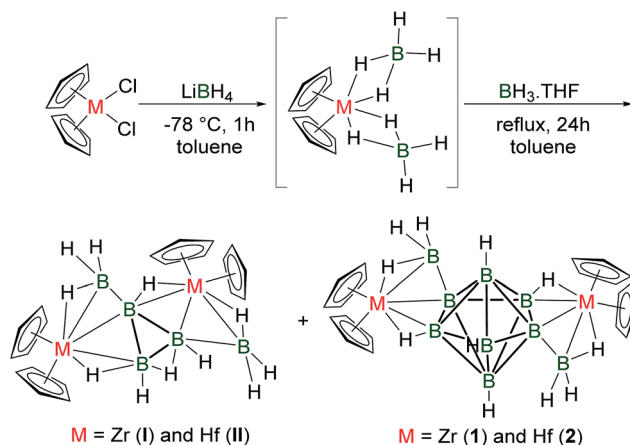
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experimental results suggested that other boranes might also be stabilized synthetically by the  $\{\text{Cp}_2\text{M}\}$  fragments of group 4 metals. This hypothesis motivated us to explore the group 4 metalboranes extensively in a quest to synthesize higher boranes stabilized by the coordination sphere of early transition metals. In the current article, we present an efficient approach for the synthesis of the first neutral transition metal protected heptaboranes **1** and **2** by a simple borane condensation method utilizing group 4 metal borohydrides. In addition, theoretical calculations have been carried out to elucidate the stability and bonding of the two new metallo-borane clusters.

## 2. Experimental

The reaction of  $[\text{Cp}_2\text{Zr}(\text{BH}_4)_2]^{15,16}$  with a large excess of  $[\text{BH}_3, \text{THF}]$  was explored under vigorous conditions, yielding  $[(\text{Cp}_2\text{Zr})_2\text{B}_9\text{H}_{11}]$  (**1**) with a novel  $\text{B}_7$  core (Fig. 1a and Scheme 1). Compound **I** was also isolated as a side product. The isolation of **1** tempted us to explore the Hf analogue, since structurally characterized hafnium–boron compounds are mostly unknown, except for one borohydride compound,  $[\text{Hf}(\eta^3\text{-BH}_4)_4]^{17}$ . Following this idea, we have investigated the reaction of  $[\text{Cp}_2\text{Hf}(\text{BH}_4)_2]^{15}$  with  $[\text{BH}_3, \text{THF}]$  under similar conditions. This indeed led to the formation of  $[(\text{Cp}_2\text{Hf})_2\text{B}_9\text{H}_{11}]$  (**2**) (Fig. 1b and Scheme 1) and  $[(\text{Cp}_2\text{Hf})_2\text{B}_5\text{H}_{11}]$  (**II**) as stable crystalline



Scheme 1 Synthesis of compounds **1** and **2**.

solids. Once compounds **1** and **2** were isolated as colourless solids, they were characterized using mass spectrometry, NMR, IR and single-crystal X-ray crystallography. Detailed syntheses and characterizations can be found in the ESI.†

## 3. Results and discussion

### 3.1. Characterizations

The ESI mass spectrum of compound **1** in 100% MeCN (Fig. S1†) shows the ion signal corresponding to the  $[\text{M} - (\text{Cp}_2\text{Zr})\text{-BH}_2]^-$  fragment (experimental  $m/z = 316.1341$ ; calculated  $m/z = 316.1566$ ). The isotopic distribution of the experimental data matches well with the calculated results. Furthermore, collision-induced dissociation (MS/MS) experiments show that the voltage threshold for the total fragmentation of  $[\text{M} - (\text{Cp}_2\text{Zr})\text{-BH}_2]^-$  is 25–30 V (Fig. S2†). When the mobile phase in the ESI was changed to 95% MeCN + 5%  $\text{H}_2\text{O}$ , the  $[\text{M} + \text{Cl}]^-$  adduct was observed (experimental  $m/z = 585.1156$ ; calculated  $m/z = 585.1369$ ) with a good match in the isotopic distribution between the experiment and simulation (Fig. S3†). The MS/MS experiments indicate that the major fragment is  $[\text{M} - (\text{Cp}_2\text{Zr})]^-$ , with the voltage threshold of total fragmentation observed at 5–8 V (Fig. S4†). Similarly, the mass spectrum of compound **2** in 100% MeCN (Fig. S5†) shows the ion peak of  $[\text{M} - (\text{Cp}_2\text{Hf})\text{BH}_2]^-$  (experimental  $m/z = 404.1802$ ; calculated  $m/z = 404.1919$ ) with the voltage threshold of total fragmentation observed at 30–35 V (Fig. S6†). When 95% MeCN + 5%  $\text{H}_2\text{O}$  was used as the mobile phase,  $[\text{M} + \text{Cl}]^-$  of **2** was observed again (experimental  $m/z = 761.1978$ ; calculated  $m/z = 761.2076$ ) (Fig. S7†), which totally fragments to  $[\text{M} - (\text{Cp}_2\text{Hf})]^-$  at 8–12 V (Fig. S8†). All of the experimental isotopic distributions for compound **2** also match very well with the calculated results. The mass spectrometric analyses clearly show the structural similarity between **1** and **2**. Both of them contain building units of  $\{\text{Cp}_2\text{M}\}$  and  $\{\text{BH}_2\}$  that can fragment easily. The only difference is the slightly higher voltage thresholds for **2**.

In order to confirm the spectroscopic assignments and determine the structures of **1** and **2**, we did X-ray structure

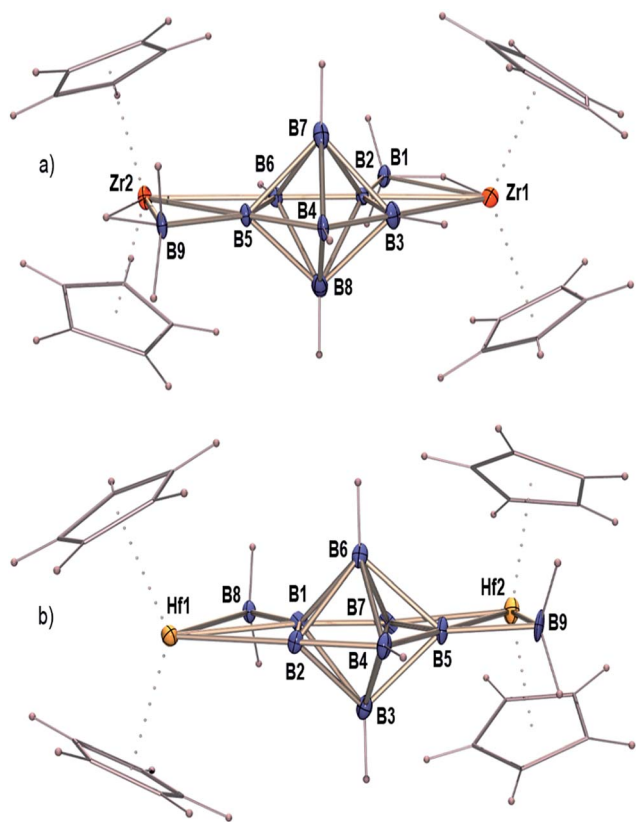


Fig. 1 The crystal structures of  $[(\text{Cp}_2\text{M})_2\text{B}_9\text{H}_{11}]$  [(a)  $\text{M} = \text{Zr}$  (**1**) and (b)  $\text{M} = \text{Hf}$  (**2**)] (some of the hydrogen atoms in **2** could not be located; details can be found in the ESI.†)



analyses (Fig. 1). The crystal structures of **1** and **2** both correspond to a 7-vertex pentagonal bipyramidal core consisting of a planar pentagonal B<sub>5</sub> belt at the equatorial position (the average non-planar deviation of the B<sub>5</sub> ring is 0.0006 Å for **1** and 0.019 Å for **2**). The core geometries of **1** and **2** are virtually identical to the B<sub>7</sub> structure in X<sub>2</sub>[B<sub>7</sub>H<sub>7</sub>] (X = PPh<sub>4</sub> or <sup>n</sup>Bu<sub>4</sub>N),<sup>11</sup> Na<sub>3</sub>B<sub>20</sub> and [(C<sub>5</sub>H<sub>5</sub>N)<sub>2</sub>CH<sub>2</sub>][B<sub>7</sub>Br<sub>7</sub>].<sup>8,18</sup> The sum of the internal angles corroborates well with the ideal pentagon (540° for a pentagon, whereas it is 540.01° for **1** and 539.79° for **2**). As expected, slight distortions are observed for individual internal angles, due to the asymmetry of the *exo*-{Cp<sub>2</sub>M} and *exo*-{BH<sub>2</sub>} moieties. The presence of the *exo*-{Cp<sub>2</sub>M} and *exo*-{BH<sub>2</sub>} moieties also supports the fragmentation channels observed in mass spectrometry.

Another interesting structural feature of compounds **1** and **2** is the presence of the {Cp<sub>2</sub>M} “wings”, which are anchored to the B<sub>7</sub> core through M–B–B and M–H–B interactions. The crystallographic data show that the average B–B bond lengths in the B<sub>7</sub> core of **1** and **2** are slightly shorter compared to [<sup>n</sup>Bu<sub>4</sub>N]<sub>2</sub>[B<sub>7</sub>H<sub>7</sub>] and [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup>, indicating a more stable B<sub>7</sub> core. Counting of the electrons becomes obvious when one realizes that the *exo*-{BH<sub>2</sub>} unit adds only one electron, like hydrogen.<sup>19</sup> The core geometry turns out to be [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup>, which is protected by two 14 electron [(Cp<sub>2</sub>M)H]<sup>+</sup> units. Alternatively, both compounds **1** and **2** can be considered as a fused cluster. As shown in Scheme S1, ‡ cluster **1** or **2** may be described as a fusion<sup>20</sup> of two *arachno*-MB<sub>3</sub> {2[(14 + 12) + 6] = 64} moieties and a *closo*-B<sub>7</sub> {(4 × 7 + 2) = 30} moiety through two B–B edges {2(6 × 2 + 2) = 28}. This arrangement of **1** or **2** generates an electron count of 66e which is equal to the total cluster valence electron (cve) number of 66 (the cve number of [(Cp<sub>2</sub>M)<sub>2</sub>B<sub>9</sub>H<sub>11</sub>] = 2 × 14 + 9 × 3 + 11 × 1 = 66). Although the isolation and structural characterization of **1** and **2** are closely connected to those of [(Cp\*Re)<sub>2</sub>(B<sub>7</sub>H<sub>7</sub>)],<sup>21</sup> the latter is highly electron deficient and shows an *oblate-closo* geometry that is inconsistent with prior theoretical findings.<sup>22</sup> Furthermore, the metal atoms in both **1** and **2** are positioned in the same plane of B<sub>5</sub>. Interestingly, both the Zr and Hf atoms are in pseudo-tetrahedral geometries, contradictory to the theoretical finding which suggests that metal atoms with relatively diffuse frontier orbitals tend to occupy the higher-degree sites.<sup>22</sup>

The <sup>1</sup>H and <sup>11</sup>B{<sup>1</sup>H} NMR spectra of both **1** and **2** apparently suggest a lower symmetry than that in the solid-state structures. The hydrogen atoms in the compounds have been assigned using <sup>1</sup>H–<sup>11</sup>B{<sup>1</sup>H} HSQC spectroscopy. The <sup>11</sup>B{<sup>1</sup>H} chemical shift of **1** at δ = –3.6 ppm can be assigned to the axial boron atoms (B<sub>7</sub> and B<sub>8</sub> in Fig. 1), because the integration is twice that of the other boron signals and the HSQC result clearly indicates the attachment of hydrogen atom(s). To examine the fluxionality of **2**, temperature-dependent <sup>1</sup>H NMR spectroscopy was undertaken (Fig. S21 ‡). The <sup>1</sup>H chemical shifts appearing at δ = –0.64 and –1.14 ppm can be assigned to two Hf–H–B protons and the broad peaks at δ = –3.26 and –3.42 ppm are assigned to the *exo*-{BH<sub>2</sub>} units. In addition, the experimental <sup>11</sup>B{<sup>1</sup>H} NMR chemical shifts of **1** and **2** show reasonable agreement with the theoretical values (error range 3–7 ppm) (Table S1 ‡). The unusual negative <sup>11</sup>B{<sup>1</sup>H} chemical shift for the axial boron

atoms, both in **1** and **2**, prompted us to apply the nucleus-independent chemical shift (NICS)<sup>23</sup> criterion to investigate the magnetic shielding in **1** and **2**. The value of the NICS(0.5) (–21.5 ppm) at the pentagonal plane of **1** is slightly more than that of the NICS(0) (–18.5 ppm). Interestingly, the NICS(3) of –6.2 ppm still shows a ring current indicating the presence of induced diatropic ring currents which is typical for a three-dimensional aromatic system (Fig. S35 ‡). Qualitatively, the NICS values observed for **2** (–22.2, –19.1 and –6.3 ppm; Fig. S36 ‡) are similar to those of **1** with slightly upfield chemical shifts. Therefore, the existence of the aromatic system plays a key role in the stabilization of **1** and **2**.

### 3.2. Bonding analyses

An alternative description of compounds **1** and **2** is that they are metal-stabilized *closo*-compounds without the necessity of having any counter ions. In order to reveal the role of the metals in stabilizing the intriguing B<sub>7</sub> core, bonding analyses of **1** and **2**, as well as the naked dianion [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup>, were performed. However, due to the presence of highly delocalized electrons in such molecules, the bonding pictures established from the molecular orbital (MO) analysis (Fig. S30 and S31 ‡) cannot provide direct insight into the chemical bonding. Thus, adaptive natural density partitioning (AdNDP)<sup>24</sup> analysis was employed to provide the intuitive bonding patterns. The bonding pattern obtained from the AdNDP analysis indicates the existence of both localized and delocalized bonds in compounds **1** and **2** (Fig. S32 and S33 ‡). Since the AdNDP results are similar for both compounds **1** and **2**, only the case of Zr is discussed here for simplicity (Fig. 2). Seven 2c–2e B–H σ bonds (Fig. 2f) can be found in **1**. Four of them are localized at the two *exo*-{BH<sub>2</sub>} units as expected. The other three, which are similar to the B–H σ bonds in [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup> (Fig. 2a), can be localized at the only equatorial non-bridged hydrogen and the two axial hydrogens. The remaining two equatorial bridged hydrogens have different bonding patterns, which will be discussed below. Although five 4c–2e σ bonds (delocalized between two adjacent equatorial boron atoms and two axial boron atoms) can be found in [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup> (Fig. 2b), only three of them are maintained in **1** (Fig. 2g). The other two are replaced by two 5c–2e σ bonds between the *exo*-{Cp<sub>2</sub>M}, the two axial and two equatorial borons (Fig. 2k). Such interactions provide strong stabilizing effects for the B<sub>7</sub> core, which is also the main reason for the slightly shortened equatorial B–B bond lengths (Fig. S34 ‡) in the B<sub>7</sub> core upon coordination by {Cp<sub>2</sub>Zr}. By forming 5c–2e σ bonds with the metal atoms, the high charge density in the naked [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup> can be lowered significantly. Therefore, the strong coulomb repulsion between the equatorial boron atoms is reduced, leading to shorter B–B bond lengths and the more stable B<sub>7</sub> core.

Three delocalized π bonds (Fig. 2h–j) can be found in the B<sub>7</sub> core of **1**. Despite the similarity between them and the three π bonds in the naked [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup> (Fig. 2c–e), the π bonds in **1** have less symmetry due to the presence of *exo*-{Cp<sub>2</sub>M} and *exo*-{BH<sub>2</sub>} moieties. Besides the two 5c–2e σ bonds (Fig. 2k) discussed above, there are six more 3c–2e σ bonds (Fig. 2l–n) that





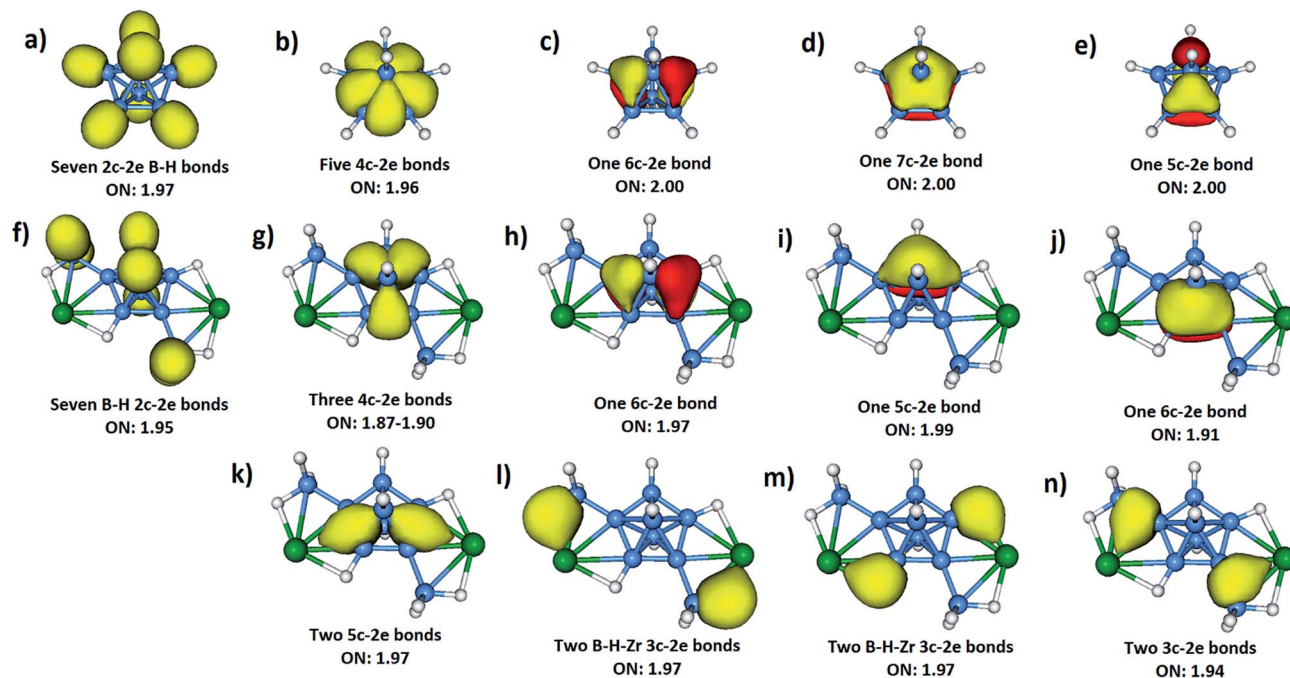


Fig. 2 AdNDP analyses of  $[\text{B}_7\text{H}_7]^{2-}$  (a–e) and **1** (f–n). The four Cp rings of **1** are omitted for clarity.

contribute to the extra stability of **1**. These bonds are absent from  $[\text{B}_7\text{H}_7]^{2-}$ : (a) two 3c-2e  $\sigma$  bonds (Fig. 2l) between the *exo*- $\{\text{Cp}_2\text{M}\}$ , the *exo* boron and an *exo* bridged hydrogen; (b) two 3c-2e  $\sigma$  bonds (Fig. 2m) between the *exo*- $\{\text{Cp}_2\text{M}\}$ , the equatorial boron and hydrogen in the  $\text{B}_7$  unit; (c) two 3c-2e  $\sigma$  bonds (Fig. 2n) between the *exo*- $\{\text{Cp}_2\text{M}\}$ , the *exo* boron and the equatorial boron in  $\text{B}_7$ . All of these multi-center bonds are crucial for keeping the integrity of compound **1**, as also revealed by the large magnitudes of the NICS(0) values for these delocalized bonds (Fig. S35 and S36<sup>†</sup>). The enhanced stability of the  $\text{B}_7$  core in **1** and **2** is also reflected in the large magnitudes of the NICS(0) values (−18.5 to −39.5 for **1**; −19.1 to −39.8 for **2**) in comparison to that of the bare  $[\text{B}_7\text{H}_7]^{2-}$  (−21.3 in Fig. S37<sup>†</sup>).

To further illustrate the strong stabilizing effect of the *exo*- $\{\text{Cp}_2\text{M}\}$  ( $\text{M} = \text{Zr}$  or  $\text{Hf}$ ) units, we studied the binding energies of the system using DFT calculations. The binding energies were calculated as the energy difference between the parent  $[(\text{Cp}_2\text{M})_2\text{B}_9\text{H}_{11}]$  molecule and the sum of two separate fragments of this compound at the PBE1PBE/Def2-TZVP level of theory. The single point energies of two schemes were calculated: (a) one of the *exo*- $\{\text{Cp}_2\text{M}\}$  units (charge state +2) was removed from the molecule, leaving the other part,  $[(\text{Cp}_2\text{M})\text{B}_9\text{H}_{11}]$ , to have a −2 charge; and (b) similar to (a), but both fragments were kept neutral. The energies of these two fragments in each scheme were calculated separately with fixed geometries. The binding energies for these two schemes were 21.2 eV and 9.3 eV for compound **1**, respectively. The difference between them, 11.9 eV, can be attributed to the stabilizing effect from electrostatic interaction or ionic bonding. The 9.3 eV value can be attributed to the stabilizing effect from the four multiple center  $\sigma$  bonds (three 3c-2e  $\sigma$  bonds and one 5c-2e  $\sigma$  bond) related to the *exo*- $\{\text{Cp}_2\text{Zr}\}$  unit or covalent bonding. As for **2**, the

electrostatic interaction was calculated to be 11.9 eV, while the effect of covalent bonding was 9.6 eV. The slightly higher binding energies of **2** also agree well with the slightly higher voltage thresholds observed in mass spectrometry. Therefore, the AdNDP analysis and binding energy calculations have revealed the crucial role of the *exo*- $\{\text{Cp}_2\text{M}\}$  and *exo*- $\text{BH}_2$  moieties for stabilizing the whole  $(\text{Cp}_2\text{M})_2\text{B}_9\text{H}_{11}$  framework.

### 3.3. Characterization of the by-product

Compound **II** (Scheme 1) which was isolated along with the formation of **2** was separated using preparative thin-layer chromatography (TLC), allowing the characterization of the pure material. It was isolated as colourless crystals and characterized using  $^1\text{H}$  and  $^{11}\text{B}\{^1\text{H}\}$  NMR and IR spectroscopy and single-crystal X-ray crystallography.<sup>25</sup> In order to confirm the spectroscopic assignments and to determine the solid-state structure of **II**, X-ray structure analysis was also undertaken (Fig. S29<sup>†</sup>). The crystal structure of **II** resembles that of  $[(\text{Cp}_2\text{Zr})_2\text{B}_5\text{H}_{11}]$  (**I**).<sup>13</sup> The core geometry can be described as *arachno*-triborane.<sup>26</sup> It is worth noting that all of the B–B bond distances of the  $\text{B}_3$  core in **II** are significantly shorter as compared to the other  $[\text{B}_3\text{H}_8]^-$  compounds, such as  $[\text{Na}(\text{B}_3\text{H}_8)]$ ,  $[\text{Cs}(\text{B}_3\text{H}_8)]$ ,  $[\text{Cr}(\text{B}_3\text{H}_8)_2]$  and **I**.<sup>13,27</sup>

## 4. Conclusions

In summary, we have presented a facile synthetic method for two early-transition-metal-protected heptaborane compounds **1** and **2**. They were synthesized using the borane condensation method utilizing group 4 metal borohydride compounds as the starting materials. Single crystal X-ray crystallography revealed



the ubiquitous presence of a pentagonal bipyramidal B<sub>7</sub> core. Our synthetic route successfully overcomes challenges faced previously in the continuing effort of isolating [B<sub>7</sub>H<sub>7</sub>]<sup>2-</sup> and its derivatives. The theoretical study revealed the intriguing role of the early transition metals in stabilizing the borane fragments *via* multicenter coordination bonding. The results described here lead to a new path towards the search for higher borane frameworks that go beyond the icosahedral topology in the coordination sphere of early transition metals. The current work may provide strategies for eventually synthesizing borospherenes and other novel boron clusters,<sup>28</sup> for which CpM-type ligands have been suggested as ideal protecting ligands.<sup>29</sup>

## Conflicts of interest

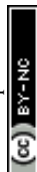
There are no conflicts to declare.

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## Notes and references

- 1 A. Stock, in *Hydrides of Boron and Silicon*, Cornell University Press, Ithaca, NY, 1933.
- 2 (a) J. D. Kennedy, *Prog. Inorg. Chem.*, 1986, **36**, 211; (b) P. v. R. Schleyer, G. Subramanian, H. Jiao, K. Najafian and M. Hofmann, in *Advances in Boron Chemistry*, ed. W. Siebert, The Royal Society of Chemistry, Cambridge, U.K., 1997; (c) E. D. Jemmis, M. M. Balakrishnarajan and P. D. Pancharatna, *Chem. Rev.*, 2002, **102**, 93–144; (d) H. Nöth, in *Homonuclear Boron Clusters in Molecular Clusters of the Main Group Elements*, ed. M. Driess and H. Nöth, Wiley-VCH, Germany, 2005.
- 3 (a) G. B. Dunks, M. M. McKown and M. F. Hawthorne, *J. Am. Chem. Soc.*, 1971, **93**, 2541–2543; (b) A. S. F. Boyd, A. Burke, D. Ellis, D. Ferrer, B. T. Giles, M. A. Laguna, R. McIntosh, S. A. Macgregor, D. L. Ormsby, G. M. Rosair, F. Schmidt, N. M. M. Wilson and A. Welch, *Pure Appl. Chem.*, 2003, **75**, 1325–1333; (c) G.-X. Jin, *Coord. Chem. Rev.*, 2004, **248**, 587–602; (d) J. Zhang and Z. Xie, *Chem.–Asian J.*, 2010, **5**, 1742–1757; (e) D. K. Roy, S. K. Bose, R. S. Anju, B. Mondal, V. Ramkumar and S. Ghosh, *Angew. Chem., Int. Ed.*, 2013, **52**, 3222–3226.
- 4 (a) W. L. Lipscomb, in *Boron Hydrides*, Benjamin, New York, 1963; (b) J. L. Boone, *J. Am. Chem. Soc.*, 1964, **86**, 5036; (c) *Boron Hydride Chemistry*, ed. E. L. Muetterties, Academic Press, New York, 1975.
- 5 (a) A. R. Pitochelli and F. M. Hawthorne, *J. Am. Chem. Soc.*, 1960, **82**, 3228–3229; (b) F. Klanberg and E. L. Muetterties, *Inorg. Chem.*, 1966, **5**, 1955–1960; (c) O. Volkov, N. P. Rath and L. Barton, *J. Organomet. Chem.*, 2003, **680**, 212–217.
- 6 W. N. Lipscomb, A. R. Pitochelli and M. F. Hawthorne, *J. Am. Chem. Soc.*, 1959, **81**, 5833–5834.
- 7 J. A. Wunderlich and W. N. Lipscomb, *J. Am. Chem. Soc.*, 1960, **82**, 4427–4428.
- 8 A. Franken, H. Thomsen and W. Preetz, *Z. Naturforsch., B*, 1996, **51**, 744–746.
- 9 F. Klanberg, D. R. Eaton, L. J. Guggenberger and E. L. Muetterties, *Inorg. Chem.*, 1967, **6**, 1271–1281.
- 10 (a) C. E. Housecroft, R. Snaith, K. Moss, R. E. Mulvey, M. E. O'Neill and K. Wade, *Polyhedron*, 1985, **4**, 1875–1881; (b) Y.-F. Shen, C. Xu and L.-J. Cheng, *RSC Adv.*, 2017, **7**, 36755–36764.
- 11 F. Schlüter and E. Bernhardt, *Inorg. Chem.*, 2011, **50**, 2580–2589.
- 12 W. Einholz, in *Advances in Boron Chemistry*, ed. W. Siebert, The Royal Society of Chemistry, Cambridge, 1997, p. 44.
- 13 (a) R. S. Anju, D. K. Roy, B. Mondal, V. Ramkumar and S. Ghosh, *Organometallics*, 2013, **32**, 4618–4623; (b) D. K. Roy, B. Mondal, A. De, S. Panda and S. Ghosh, *Organometallics*, 2015, **34**, 908–912.
- 14 (a) R. L. Thomas, N. P. Rath and L. Barton, *J. Am. Chem. Soc.*, 1997, **119**, 12358–12375; (b) R. L. Thomas, N. P. Rath and L. Barton, *Inorg. Chem.*, 2002, **41**, 67–75.
- 15 T. J. Marks and J. R. Kolb, *Chem. Rev.*, 1977, **77**, 263–293.
- 16 R. K. Nanda and M. G. H. Wallbridge, *Inorg. Chem.*, 1964, **3**, 1798.
- 17 R. W. Broach, I.-S. Chuang, T. J. Marks and J. M. Williams, *Inorg. Chem.*, 1983, **22**, 1081–1084.
- 18 B. Albert and K. Hofmann, *Z. Anorg. Allg. Chem.*, 1999, **625**, 709–713.
- 19 S. Ghosh, B. C. Noll and T. P. Fehlner, *Angew. Chem., Int. Ed.*, 2005, **44**, 2916–2918.
- 20 (a) D. M. P. Mingos, *Nature (London), Phys. Sci.*, 1972, **236**, 99–102; (b) D. M. P. Mingos, *Acc. Chem. Res.*, 1984, **17**, 311–319.
- 21 B. L. Guennic, H. Jiao, S. Kahlal, J.-Y. Saillard, J.-F. Halet, S. Ghosh, M. Shang, A. M. Beatty, A. L. Rheingold and T. P. Fehlner, *J. Am. Chem. Soc.*, 2004, **126**, 3203–3217.
- 22 E. D. Jemmis, *J. Am. Chem. Soc.*, 1982, **104**, 7017–7020.
- 23 (a) P. v. R. Schleyer, C. Maerker, A. Dransfeld, H. Jiao and N. J. R. v. E. Hommes, *J. Am. Chem. Soc.*, 1996, **118**, 6317–6318; (b) Z. Chen, C. S. Wannere, C. Cominboeuf, R. Puchta and P. v. R. Schleyer, *Chem. Rev.*, 2005, **105**, 3842–3888.
- 24 D. Y. Zubarev and A. I. Boldyrev, *Phys. Chem. Chem. Phys.*, 2008, **10**, 5207–5217.
- 25 The mass spectrum of **II** showed a molecular ion peak at *m/z* 686.4 (Fig. S9†). The <sup>1</sup>H NMR spectrum displayed three signals at δ = 5.92, 5.90 and 5.89 ppm in a 2:1:1 ratio which rationalizes the presence of two types of Cp ligand. The presence of Zr–H–B (δ = –4.16 ppm) and B–H–B (δ =



- 2.74 ppm) was confirmed from  $^1\text{H}$  NMR. The  $^{11}\text{B}\{^1\text{H}\}$  NMR spectrum showed three resonances at  $\delta = 4.6$ , 2.0 and –3.8 ppm in a 2:1:2 ratio.
- 26 Although the positions of the hydrogen atoms were not able to be fully located in the crystal structure, they were assigned based on the spectroscopic data, as shown in Scheme 1. The different B–B distances of the B3 unit in **II** are due to the presence of bridged hydrogen along the B2–B3 bond.
- 27 (a) H. J. Deiseroth, O. Sommer, H. Binder, K. Wolfer and B. Frei, *Z. Anorg. Allg. Chem.*, 1989, **571**, 21–28; (b) D. Y. Kim and G. S. Girolami, *J. Am. Chem. Soc.*, 2006, **128**, 10969–10977; (c) A. C. Dunbar, J. A. Macor and G. S. Girolami, *Inorg. Chem.*, 2014, **53**, 822–826.
- 28 (a) H. J. Zhai, Y. F. Zhao, W. L. Li, Q. Chen, H. Bai, H. S. Hu, Z. A. Piazza, W. J. Tian, H. G. Lu, Y. B. Wu, Y. W. Mu, G. F. Wei, Z. P. Liu, J. Li, S. D. Li and L. S. Wang, *Nat. Chem.*, 2014, **6**, 727–731; (b) Z. A. Piazza, H. S. Hu, W. L. Li, Y. F. Zhao, J. Li and L. S. Wang, *Nat. Commun.*, 2014, **5**, 3113; (c) L. S. Wang, *Int. Rev. Phys. Chem.*, 2016, **35**, 69–142; (d) W. L. Li, X. Chen, T. Jian, T. T. Chen, J. Li and L. S. Wang, *Nat. Rev. Chem.*, 2017, **1**, 0071.
- 29 N. Karmodak and E. D. Jemmis, *Chem.–Asian J.*, 2016, **11**, 3350–3354.

