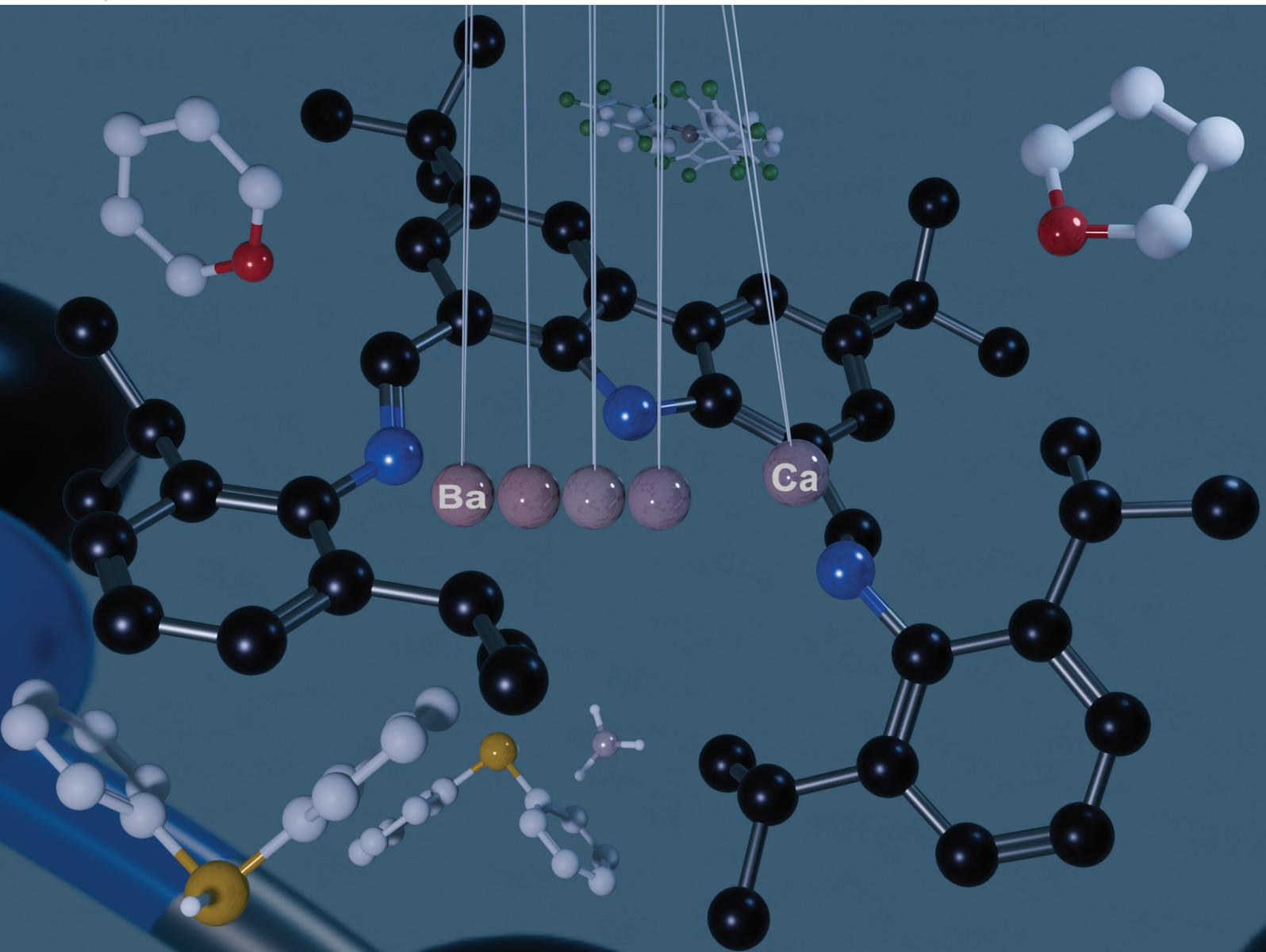


Dalton Transactions

An international journal of inorganic chemistry

rsc.li/dalton



ISSN 1477-9226

Cite this: *Dalton Trans.*, 2024, **53**, 6892

Barium phosphidoboranes and related calcium complexes†

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The attempted synthesis of $[(\text{Carb})\text{BaPPh}_2]$ (**1**) showed this barium-phosphide and its thf adducts, **1**·thf and **1**·(thf)₂, to be unstable in solution. Our strategy to circumvent the fragility of these compounds involved the use of phosphinoboranes HPPH₂·BH₃ and HPPH₂·B(C₆F₅)₃ instead of HPPH₂. This allowed for the synthesis of $[(\text{Carb})\text{Ae}(\text{PPh}_2\text{·BH}_3)]$ (Ae = Ba, **2**; Ca, **3**), $[(\text{Carb})\text{Ca}(\text{H}_3\text{B}_2\text{PPh}_2)\text{·thf}]$ (**4**), $[(\text{Carb})\text{Ba}(\text{PPh}_2\text{·B}(\text{C}_6\text{F}_5)_3)]$ (**5**), $[(\text{Carb})\text{Ba}(\text{O}(\text{B}(\text{C}_6\text{F}_5)_3)\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{PPh}_2)\text{·thf}]$ (**6**), $[\text{Ba}\{\text{O}(\text{B}(\text{C}_6\text{F}_5)_3)\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{PPh}_2\}_2\text{·(thf)}_{1.5}]$ (**7**) and $[\text{Ba}(\text{PPh}_2\text{·B}(\text{C}_6\text{F}_5)_3)_2\text{·(thp)}_2]$ (**8**) that were characterised by multinuclear NMR spectroscopy (thp = tetrahydropyran). The molecular structures of **4**, **6** and **8** were validated by X-ray diffraction crystallography, which revealed the presence of Ba...F stabilizing interactions (ca. 9 kcal mol⁻¹) in the fluorine-containing compounds. Compounds **6** and **7** were obtained upon ring-opening of thf by their respective precursors, **5** and the *in situ* prepared $[\text{Ba}(\text{PPh}_2\text{·B}(\text{C}_6\text{F}_5)_3)_2]_n$. By contrast, thp does not undergo ring-opening under the same conditions but affords clean formation of **8**. DFT analysis did not highlight any specific weakness of the Ba–P bond in **1**·(thf)₂. The instability of this compound is instead thought to stem from the high energy of its HOMO, which contains the non-conjugated P lone pair and features significant nucleophilic reactivity.

Received 19th February 2024,
Accepted 19th March 2024

DOI: 10.1039/d4dt00487f

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Introduction

Soluble complexes of the large alkaline earth metals (= Ae) calcium, strontium and barium are increasingly useful in organic synthesis.^{1,2} They make excellent molecular catalysts for hydrophosphination reactions,^{3,4} e.g. for the *anti*-Markovnikov addition of phosphines on activated alkenes^{5–11} where barium systems perform particularly well.¹² The heteroleptic complex $[(\text{Carb})\text{BaN}(\text{SiMe}_3)_2]$ supported by the bulky bis(imino)carbazolato monoanionic ligand $\{\text{Carb}\}^-$ (Fig. 1) is a typical case of an efficient hydrophosphination precatalyst.¹¹

In vinylarene hydrophosphination promoted by Ae precatalysts of the general composition $[(\text{Lig})\text{AeX}(\text{solvent})_n]$ (where X⁻ is chosen among N(SiMe₃)₂⁻, CH(SiMe₃)₂⁻, H⁻ etc., and {Lig}⁻ is a monoanionic ancillary ligand), it is accepted that the active species responsible for catalytic turnovers is the heteroleptic phosphide $[(\text{Lig})\text{Ae}(\text{PR}_2)\text{·(D)}_n]$, where D is a molecule of the

(donor) solvent or phosphine. This phosphido species undergoes insertion of the C=C unsaturated bond into the Ae–P bond, an elementary step followed by protonolysis of the resulting Ae–alkyl bond with an incoming phosphine to regenerate the active species.¹ In hydrocarbons, the use of an ancillary ligand is required to suppress Schlenk redistribution to the insoluble homoleptic barium diphosphide $[\text{Ba}(\text{PR}_2)_2]_\infty$.^{13,14} A handful of soluble, homoleptic complexes $[\text{Ba}(\text{PAr}_2)_2(\text{D})_n]$ are known for D = thf, dme or 18-c-6.^{13–17} The rare examples of crystallographically characterised heteroleptic Ba-phosphides include the dimeric $[\text{Ba}\{\text{P}(\text{H})\text{Si}^t\text{Bu}_3\}\{\text{N}(\text{SiMe}_3)_2\}_2\text{·(thf)}_2]$ ¹⁵ and $[(\text{IA}^{\text{crown}})\text{Ba}(\text{PPh}_2)]$, where $\{\text{IA}^{\text{crown}}\}^-$ is a multidentate, {amino-tetraether}-anilide (see Fig. 1 for the proligands used to produce heteroleptic Ba-phosphides). We found the synthesis of the latter to be erratic.¹⁰ In this context, it seemed desirable to assess whether the $\{\text{Carb}\}^-$ ligand framework could reliably stabilise the heteroleptic phosphido complex $[(\text{Carb})\text{Ba}(\text{PPh}_2)]$. As the following will show, it soon emerged that this ambition cannot be fulfilled.

A suite of heteroleptic phosphidoborane magnesium (**D–F**, Fig. 2) and calcium (**G–I**) complexes supported by the ubiquitous $\{\text{BDI}^{\text{DIPP}}\}^-$ β-diketiminato ligand have been described by the Hill group.^{18,19} The aim was to assess Ae catalysts for HPPH₂·BH₃ dehydrocoupling and to produce polyphosphinoboranes. In a related topic, the chemistry of the amidoborane derivatives of the Ae metals had provided a mechanistic under-

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† Electronic supplementary information (ESI) available: Experimental details, spectroscopic characterisation, and crystallographic and computational data. CCDC 2314377–2314380. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d4dt00487f>

‡ These two co-authors have contributed equally to the work and must both be considered as first co-authors.



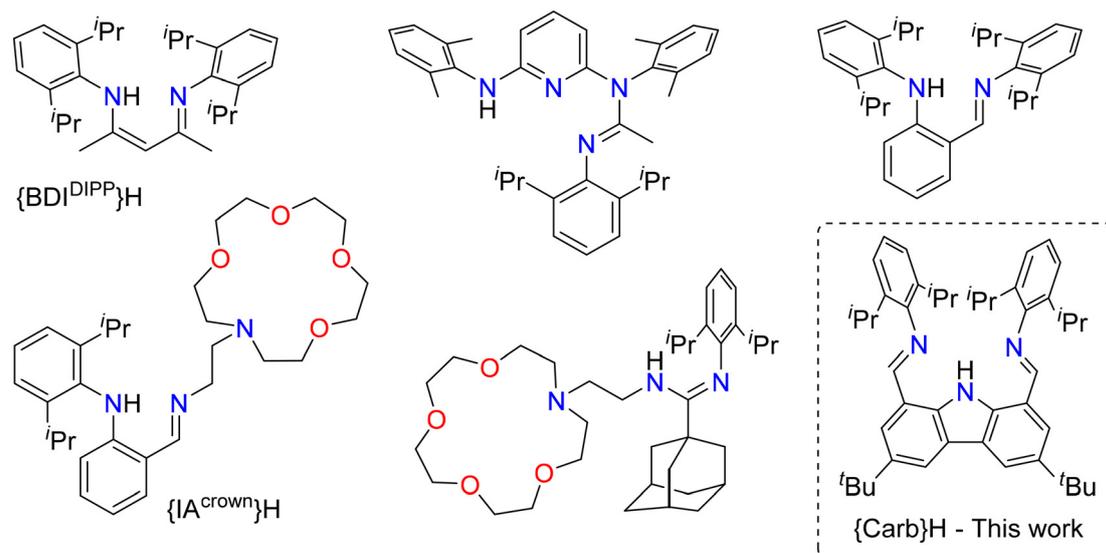


Fig. 1 Proligands used to produce heteroleptic Ba–phosphides. Among these, only $\{\text{IA}^{\text{crown}}\}\text{H}$ has led to a structurally authenticated complex, $\{\{\text{IA}^{\text{crown}}\}\text{Ba}(\text{PPh}_2)\}$.^{5–11}

standing of Ae-catalysed amine–borane dehydrogenation.^{20–22} Of specific interest is that the reaction of $\{[\text{BDI}^{\text{Dipp}}]\text{Ca}\{\text{PPh}_2\cdot\text{BH}_3\}\}$ (**H**) with $\text{B}(\text{C}_6\text{F}_5)_3$ gave $\{[\text{BDI}^{\text{Dipp}}]\text{Ca}\{\text{H}_3\text{B}\cdot\text{PPh}_2\cdot\text{B}(\text{C}_6\text{F}_5)_3\}\cdot(\eta^6\text{-toluene})\}$ (**I**).¹⁹ Beyond Hill's complexes, known examples of phosphidoborane Ae compounds are limited to Izod's homoleptic Ca and Sr complexes and their Mg congeners, synthesised by the reaction of Li- and K-phosphides with AeI_2 , e.g. $\{[(\text{Me}_3\text{Si})_2\text{CH}(\text{Ph})\text{P}(\text{BH}_3)_2]_2\text{Ae}\cdot(\text{thf})_4\}$ (Ae = Ca, K; Sr, L).^{23,24} We are not aware of a previous barium-phosphidoborane complex.

We report here our attempt at producing heteroleptic Ba-phosphides. Confronted by the instability of the sought compounds, we have used phosphinoboranes as starting materials in order to generate more stable Ba–P–B motifs, and the corresponding barium phosphidoboranes, along with related calcium complexes, are described. We also show that $\{[\text{Carb}]\text{Ba}\{\text{PPh}_2\cdot\text{B}(\text{C}_6\text{F}_5)_3\}\}$ ring-opens thf to provide $\{[\text{Carb}]\text{Ba}\{\text{O}(\text{B}(\text{C}_6\text{F}_5)_3)(\text{CH}_2)_4\text{PPh}_2\}\cdot\text{thf}\}$, an unusual Ba-alkoxide. By contrast, the six-member tetrahydropyran (= thp) is inert under similar conditions. The metal-assisted ring-opening of thf with phosphines or phosphides has been previously documented with oxophilic metals including Li,²⁵ Mg,²⁶ lanthanides/actinides^{27–29} or Zr,³⁰ and for a number of phosphine-based frustrated Lewis pairs.^{31–34}

Results and discussion

In an attempt to produce a stable heteroleptic barium–phosphido complex, HPPH_2 was reacted with the solvent-free $\{[\text{Carb}]\text{BaN}(\text{SiMe}_3)_2\}$ (Scheme 1). The bulky bis(imino)carbazolato ligand $\{\text{Carb}\}^-$ has been shown to meet the demanding coordination requirements of the large barium ion ($r_{\text{ionic}}(\text{Ba}^{2+}) = 1.35 \text{ \AA}$).^{11,35} The addition of diphenylphosphine to the

solution of the barium precursor in petroleum ether resulted in an instant colour change from yellow to dark orange, and the precipitation of a vibrant orange solid was observed within 5–10 minutes. Variable temperature NMR analysis of this orange powder revealed two distinct phosphido species, each with a carbazole fragment; no equilibrium between the two species was observed by NMR. One of these species was hypothesised to be the sought Ba–phosphide, $\{[\text{Carb}]\text{Ba}(\text{PPh}_2)\}$ (**1**). However, despite repeated attempts, we were unable to isolate or identify a clean material from this reaction, and neither product could be crystallised from the mixture. Instead of isolating the precipitate, direct addition of thf to the orange suspension in petroleum ether allowed for the isolation of small microcrystals. This material diffracted poorly, and the quality of the dataset is too low to warrant precise discussion of the metric parameters ($R_{\text{int}} = 0.1219$, $R_1 = 0.1235$). Yet, the connectivity in the structure was confirmed to be the twice solvated, heteroleptic Ba–phosphide $\{[\text{Carb}]\text{Ba}(\text{PPh}_2)\cdot(\text{thf})_2\}$ (**1**·(thf)₂), a compound with a six-coordinate distorted octahedral geometry around Ba (Fig. 3). We were not able to record NMR data for this compound due to the insufficient amount of material. ¹H and ³¹P NMR data collected in toluene-*d*₈ for the *in situ* reaction of HPPH_2 and the solvated $\{[\text{Carb}]\text{BaN}(\text{SiMe}_3)_2\cdot(\text{thf})\}$ (see ESI, Fig. S1 and S2†) were consistent with the clean formation of the mono-thf adduct $\{[\text{Carb}]\text{BaPPh}_2\cdot(\text{thf})\}$ (**1**·thf). However, this product proved unstable and decomposed during collection of ¹³C NMR data to a number of other species, including $\text{Ph}_2\text{P}\text{--}\text{PPh}_2$ ³⁶ (the mechanism of formation of this compound, e.g. homolytic cleavage or dehydrocoupling of HPPH_2 released *in situ*, could not be

§ The barium and calcium complexes described herein were too air- and moisture sensitive to allow for elemental and mass spectrometry analyses.



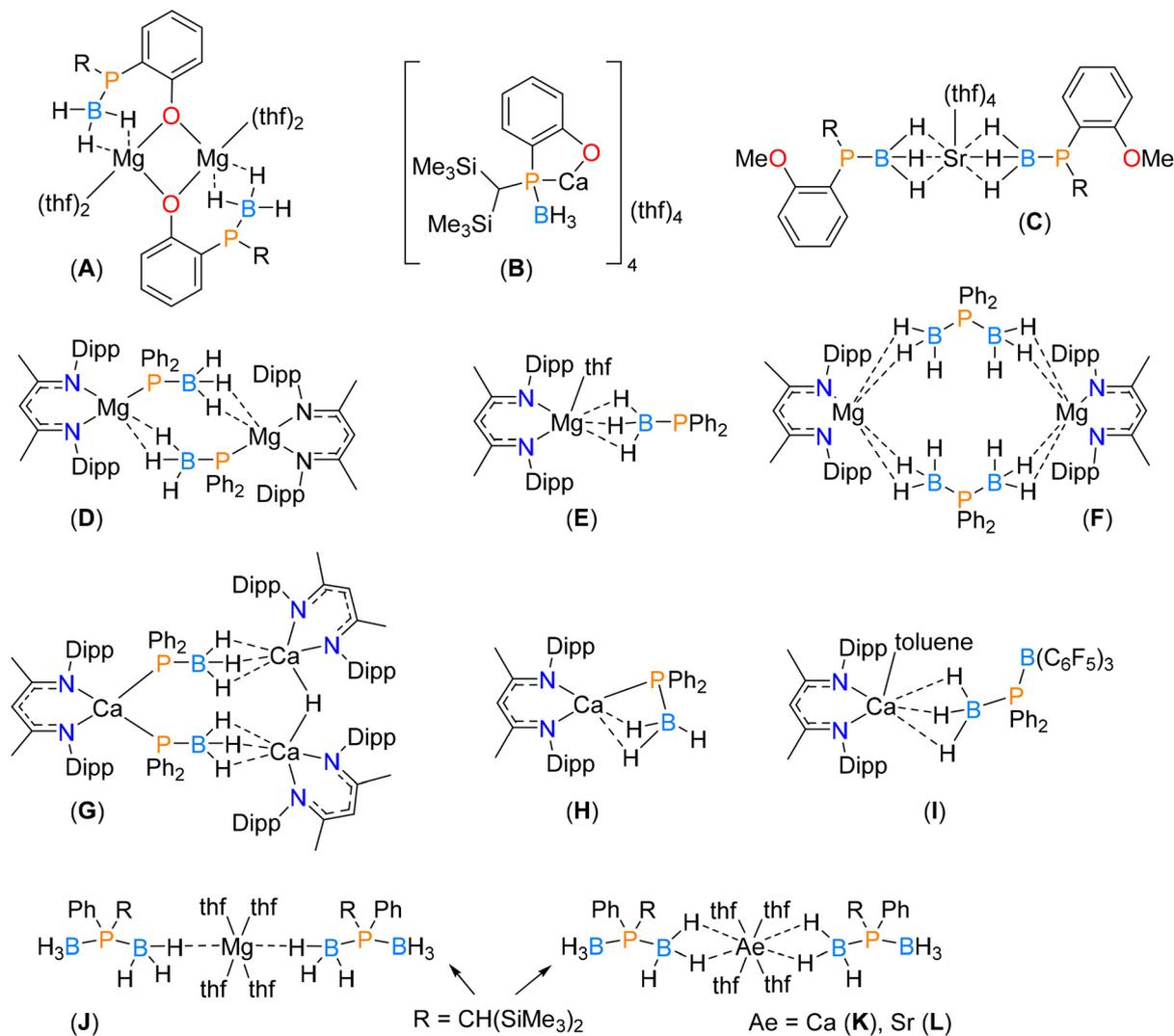


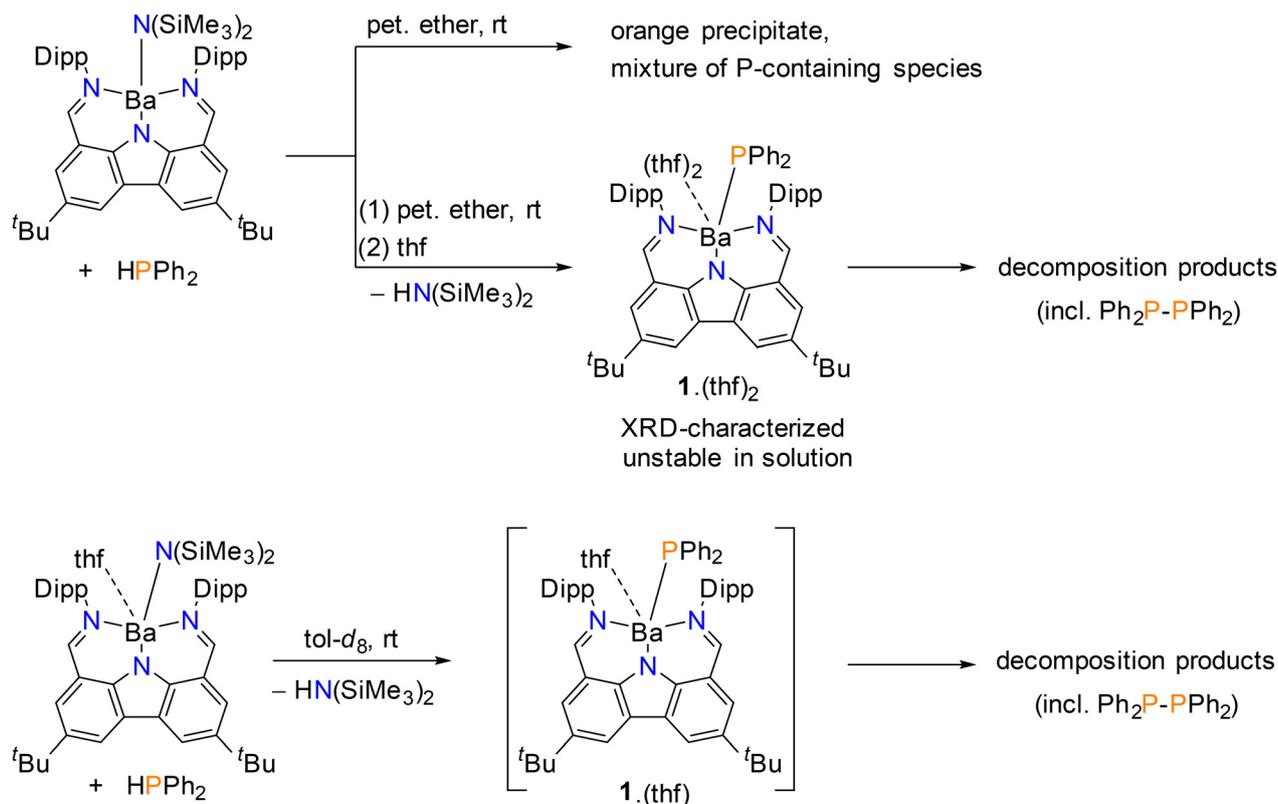
Fig. 2 Known Mg-, Ca- and Sr-phosphidoboranes (A)–(L) characterised by single crystal X-ray diffraction.^{18,19,23,24} R = CH(SiMe₃)₂ and Dipp = 2,6-*i*-Pr₂-C₆H₃.

determined). Some of the unknown resonances were similar to those observed in the NMR spectra for the abovementioned compounds. The instability of the Ba–PPh₂ bond is perhaps unsurprising. For instance, bond dissociation energies in a suite of terminal zirconium–phosphides have been calculated to be low (*ca.* 55.4–65.3 kcal mol^{−1}),³⁷ whereas the homolytic cleavage of Bi(III)–PPh₂ in heteroleptic bismuth–phosphides has been reported to generate Bi(II) species.³⁸ Despite repeated attempts, a sample of **1**·thf could not be isolated to allow full characterisation. Overall, we concluded that the isolation of a stable Ba–phosphido species supported by the {Carb}[−] ligand could not be achieved.

In order to better understand the stability and bonding within Ba–phosphido complexes, density functional theory (DFT) calculations were carried out on the isolated solvated complex **1**·(thf)₂ at the PBE0-ZORA-D3(BJ)/TZ2P level (see the ESI† for computational details). Selected computed data are provided in Table 1. The optimised geometry of **1**·(thf)₂ is con-

sistent with its X-ray structure, in spite of the limitations mentioned above concerning the quality of the XRD data. Except perhaps for the somewhat large Ba–P interatomic distance (optimised value 3.206 Å *vs.* experimental one of *ca.* 3.296(3) Å, although it falls in the typical range for Ba–P bonds^{13–17}), there is no indication that hints at a particularly weak Ba–P bond. To get a better insight, the hypothetical amido homologue [{Carb}Ba(NPh₂)·(thf)₂] was also calculated (selected computed data in Table 1). Its structural features are quite similar to those of **1**·(thf)₂, except that the NPh₂ nitrogen atom is significantly less pyramidalised than its PPh₂ counterpart in **1**·(thf)₂ (the sum of covalent angles 356° and 320° for N and P, respectively). This difference results from the substantial conjugation of the nitrogen lone pair with the phenyl rings. Although comparing the Wiberg bond indices (WBIs, see Table 1) between bonds involving different elements should be made with caution, the corresponding Ba–PPh₂ and Ba–NPh₂ values suggest that the covalent component of the Ba–P bond





Scheme 1 Attempted synthesis of a stable barium–phosphido complex. Dipp = 2,6-diisopropylphenyl.

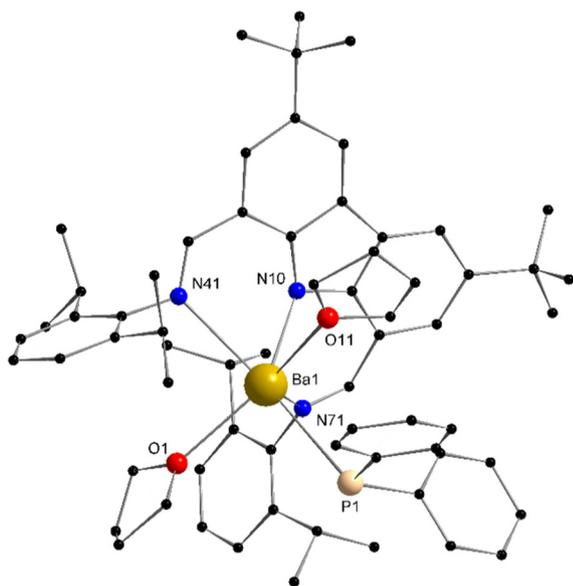


Fig. 3 Representation of the molecular solid-state structure of [(Carb)Ba(PPh₂).(thf)₂] (1.(thf)₂) showing connectivity. H atoms and two non-coordinating thf molecules are omitted for clarity. $R_{\text{int}} = 0.1219$, $R_1 = 0.1224$.

is weaker than its Ba–N homologue. The natural atomic charge distribution (Table 1) indicates also the weaker ionic bonding in the case of the Ba–P bond.

In addition to these qualitative indicators, a more quantitative comparative analysis of the Ba–EPh₂ (E = P, N) bonds can be made by performing an energy decomposition analysis (EDA) of the interaction between two frozen molecular fragments, according to the Morokuma–Ziegler procedure.^{39–41} The decomposition of the total bonding energy (TBE) between the [(Carb)Ba.(thf)₂]⁺ fragment and the [EPh₂][−] (E = P, N) ligand in the two complexes is provided in Table 2. TBE is expressed as the sum of four components: the Pauli repulsion (E_{Pauli}), the electrostatic interaction energy (E_{elstat}), the orbital interaction energy (E_{orb}) and the component associated with the dispersion forces (E_{disp}). It is of note that the TBE components are comparable between both complexes, those of the amido species indicating slightly stronger (positive or negative) interactions, except for the minor dispersion component. The dominating component is E_{elstat} , being about three times larger than E_{orb} . This is in line with a bonding possessing a largely ionic characteristic, but with non-negligible covalency. The TBE values confirm that the Ba–NPh₂ bond is stronger than its Ba–PPh₂ analogue, with both stronger covalent (E_{orb}) and electrostatic (E_{elstat}) interactions. The overall TBE difference of ca. 7 kcal mol^{−1} remains however modest, as in effect are also the differences across the two compounds for each of the four contributions.

The Quantum Theory of Atoms in Molecules (QTAIM) approach^{42,43} provides a different angle to look at these Ba–E (E = P, N) bonds. Selected QTAIM data are collated in Table 3.



Table 1 Relevant DFT-computed data for **1**·(thf)₂, its amido analogue [(Carb)Ba(NPh₂)·(thf)₂], and **5**

		[(Carb)Ba(PPh ₂)·(thf) ₂] (1 ·(thf) ₂)	[(Carb)Ba(NPh ₂)·(thf) ₂] (putative)	[(Carb)Ba{PPh ₂ ·B(C ₆ F ₅) ₃ }] (5)
HOMO–LUMO gap/eV		2.16	3.70	3.76
Computed interatomic distances (Å) and [WBI] ^a (experimental values in italics for comparison)	Ba–P/N	3.206 [0.103] 3.296(3)	2.683 [0.250]	3.303 [0.062]
	Ba–N _{carb.}	2.647 [0.050] 2.629(9)	2.660 [0.047]	2.590 [0.068]
	Ba–N _{imin.} ^b	2.861 [0.027] 2.824(9)	2.883 [0.025]	2.740 [0.037]
	Ba–O/F ^b	2.727 [0.021] 2.737(9)	2.772 [0.020]	2.863 [0.010]
Natural atomic charges	Ba	+1.78	+1.81	+1.82
	P/N	–0.15	–0.79	+0.32
	N _{carb.}	–0.66	–0.66	–0.70
	N _{imin.} ^b	–0.56	–0.56	–0.60
	O/F ^b	–0.60	–0.60	–0.37

^a Wiberg bond indices. ^b Averaged values.

Table 2 Morokuma–Ziegler energy decomposition analysis in **1**·(thf)₂, putative [(Carb)Ba(NPh₂)·(thf)₂], and **5**^a

	[(Carb)Ba(PPh ₂)·(thf) ₂] (1 ·(thf) ₂)	[(Carb)Ba(NPh ₂)·(thf) ₂] (putative complex)	[(Carb)Ba{PPh ₂ ·B(C ₆ F ₅) ₃ }] (5)
Fragmentation	[(Carb)Ba·(thf) ₂] ⁺ + [PPh ₂] [–]	[(Carb)Ba·(thf) ₂] ⁺ + [NPh ₂] [–]	[(Carb)Ba] ⁺ + [PPh ₂ ·B(C ₆ F ₅) ₃] [–]
<i>E</i> _{Pauli}	+2.43	+2.57	+1.84
<i>E</i> _{elstat}	–4.73	–4.95	–3.62
<i>E</i> _{orb}	–1.46	–1.72	–1.50
<i>E</i> _{disp}	–0.85	–0.82	–1.24
TBE ^b	–4.61 (–106 kcal mol ^{–1})	–4.92 (–113 kcal mol ^{–1})	–4.52 (–104 kcal mol ^{–1})

^a Values in eV, unless specified. ^b Total bonding energy (TBE) = *E*_{Pauli} + *E*_{elstat} + *E*_{orb} + *E*_{disp}.

Table 3 QTAIM descriptors (in a.u.) of the Ba–EPh₂ (E = P, N) bonds in **1**·(thf)₂, [(Carb)Ba(NPh₂)·(thf)₂] and **5**^{a,b}

		[(Carb)Ba(PPh ₂)·(thf) ₂] (1 ·(thf) ₂)	[(Carb)Ba(NPh ₂)·(thf) ₂] (putative complex)	[(Carb)Ba{PPh ₂ ·B(C ₆ F ₅) ₃ }] (5)
Atom charge	Ba	+1.61	+1.64	+1.60
	P/N	+0.54	–1.23	+0.71
	N _{carb.}	–1.14	–1.17	–1.17
	N _{imin.} ^a	–1.19	–1.17	–1.21
	O ^a	–1.09	–1.09	
Delocalisation index	δ	0.27	0.27	0.19
bcp indicators ^a	ρ	0.027	0.039	0.022
	∇ ² ρ	0.046	0.111	0.041
	H	–0.003	–0.004	–0.001
	V	–0.018	–0.035	–0.013
	V /G	1.22	1.12	1.13

^a Averaged values. ^b ρ, ∇²ρ, H, V and G are the electron density, Laplacian of ρ density, energy density, potential energy density and kinetic energy density values at the bcp, respectively. All values in a.u.

Surprisingly, the AIM charge on P in **1**·(thf)₂ is now significantly positive, whereas that of N in its amido analogue remains strongly negative. It is consistent with a more ionic (or very polar⁴⁴) characteristic of the Ba–N bond. All Ba–EPh₂ (E = P, N) bond critical point (bcp) indicators have small absolute values. The positive sign of the Laplacian density (larger in the case of N), the negative sign of the (very small) energy density and the |V|/G ratio somewhat larger for **1**·(thf)₂ are compatible with a weakly covalent, strongly ionic bonding.

In any case, the above analyses do not underscore any particular weakness of the Ba–phosphide bond, even if it is not as

strong as its Ba–amide homologue. We suggest that the relative instability of **1**·thf or **1**·(thf)₂ is not of thermodynamic origin, but instead it is related to the reactivity of the lone pair on the P atom. In contrast to its Ba–NPh₂ amido relative, where the nitrogen lone pair is stabilised through conjugation, the phosphido lone pair in **1**·(thf)₂ does not delocalise especially on the phenyl rings and remains rather unperturbed, hence constituting the HOMO in the complex. This HOMO sits isolated 1.60 eV above the HOMO–1, and confers to **1**·(thf)₂ a significant nucleophilic ability, especially as the lone pair is not protected. Very similar differences between the P and N derivatives were



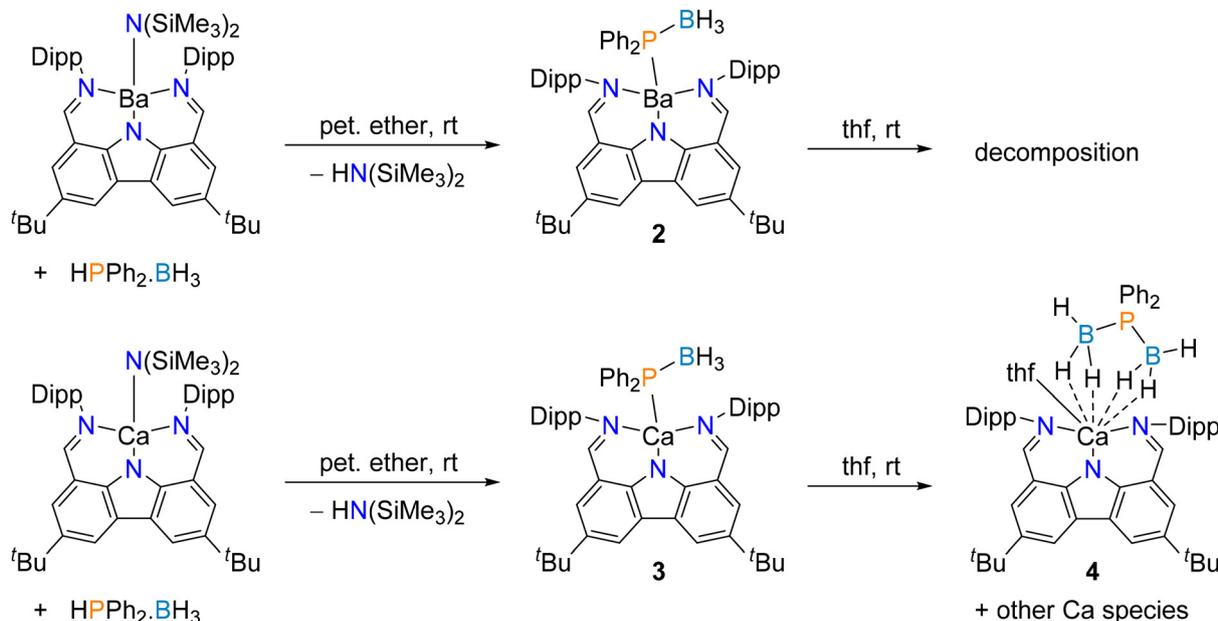
computed for the unsolvated species $[\{\text{Carb}\}\text{Ba}(\text{EPh}_2)]$ ($\text{E} = \text{P}, \text{N}$), as well as for the $[\{\text{Carb}\}\text{BaE}(\text{SiMe}_3)_2]$ ($\text{E} = \text{P}, \text{N}$) pair.

Magnesium, calcium and even strontium phosphidoborane complexes that contain Ae–P bonds are known.^{18,19,23,24} We hence turned our attention to the use of phosphine–borane in order to achieve the stabilisation of a Ba–P bond in a heteroleptic complex. In an NMR-scale reaction, the data recorded in benzene-*d*₆ after addition of one equivalent of HPPh₂·BH₃ to $[\{\text{Carb}\}\text{BaN}(\text{SiMe}_3)_2]$ were consistent with formation of $[\{\text{Carb}\}\text{Ba}\{\text{PPh}_2\cdot\text{BH}_3\}]$ (**2**) (Scheme 2). The ¹H NMR spectrum (Fig. S3†) showed the release of HN(SiMe₃)₂, while only one symmetrical {Carb}[−] ligand environment could be detected. As reported previously for $[\{\text{BDI}^{\text{Dipp}}\}\text{Ca}\{\text{PPh}_2\cdot\text{BH}_3\}]$ and $[\{\text{BDI}^{\text{Dipp}}\}\text{Mg}\{\text{PPh}_2\cdot\text{BH}_3\}]_2$ (**H** and **D**, respectively, in Fig. 2),¹⁸ the resonances for boron-bound H atoms in **2** could not be seen. The ³¹P NMR spectrum of the complex displayed a single broad singlet resonance at $\delta_{\text{P}} -32.3$ ppm, shifted significantly upfield from the broad doublet of HPPh₂·BH₃ (1.6 ppm) and confirming the deprotonation of the phosphine. The ¹¹B NMR spectrum also exhibited a broad singlet ($\delta_{\text{B}} -24.5$ ppm), downfield shifted from HPPh₂·BH₃ (−40.3 ppm). Both chemical shifts are in line with those for $[\{\text{BDI}^{\text{Dipp}}\}\text{Ca}\{\text{PPh}_2\cdot\text{BH}_3\}]$ (toluene-*d*₈, 48 °C: $\delta_{\text{P}} -45.0$ ppm and $\delta_{\text{B}} -30.2$ ppm) and $[\{\text{BDI}^{\text{Dipp}}\}\text{Mg}\{\text{PPh}_2\cdot\text{BH}_3\}]_2$ (toluene-*d*₈, 76 °C: $\delta_{\text{P}} -57.1$ and $\delta_{\text{B}} -34.0$ ppm),¹⁸ though meaningful comparison is difficult due to the differences in metals, ligand environments, solvents and temperature. When the reaction was scaled up in petroleum ether, complex **2** could be isolated in near-quantitative yield as a bright yellow powder. However, in spite of multiple attempts, single-crystals suitable for analysis by XRD could not be obtained. § A solution of **2** in thf rapidly lost its bright yellow colour, with concomitant formation of a yellow precipi-

tate. The ¹H NMR spectrum of the isolated powder revealed a mixture of unassignable {Carb}[−] environments, while no signal could be observed in the ³¹P and ¹¹B NMR spectra. Again, despite several attempts, crystals could not be obtained from this solid to allow identification of any of its components by X-ray diffraction analysis.

In order to improve our understanding of the behaviour of barium complexes, the same experiments were carried out using calcium precursors, as Ca complexes, being less ionic in nature, are generally more robust in solution than their Sr and Ba congeners. Metalation of HPPh₂·BH₃ with $[\{\text{Carb}\}\text{CaN}(\text{SiMe}_3)_2]$ in toluene-*d*₈ resulted in the clean formation of $[\{\text{Carb}\}\text{Ca}\{\text{PPh}_2\cdot\text{BH}_3\}]$ (**3**). The ¹H NMR spectrum displays a single {Carb}[−] ligand environment, though again no identifiable BH₃ resonance could be detected. The ³¹P NMR spectrum displays a broad resonance at $\delta_{\text{P}} -41.3$ ppm, while the ¹¹B NMR spectrum features a broad signal at $\delta_{\text{B}} -26.9$ ppm. Both match those already mentioned for $[\{\text{BDI}^{\text{Dipp}}\}\text{Ca}\{\text{PPh}_2\cdot\text{BH}_3\}]$.¹⁸ Again, despite repeated attempts, satisfactory crystals of **3** for XRD analysis could not be isolated. § A small amount of thf was added to a sample of **3** in petroleum ether, in an attempt to produce a thf adduct that would perhaps allow for solid-state characterisation. Unexpectedly, the calcium–phosphidodiborate $[\{\text{Carb}\}\text{Ca}\{\text{(H}_3\text{B)}_2\text{PPh}_2\}\cdot(\text{thf})]$ (**4**) was isolated in 70% yield as pale yellow crystals. Its identity was established by X-ray diffraction (see Fig. 4) and NMR spectroscopy. In contrast, the addition of thf to a solution of $[\{\text{BDI}^{\text{Dipp}}\}\text{Ca}\{\text{PPh}_2\cdot\text{BH}_3\}]$ (**H** in Fig. 2) in benzene-*d*₆ was reported to give the adduct $[\{\text{BDI}^{\text{Dipp}}\}\text{Ca}\{\text{PPh}_2\cdot\text{BH}_3\}\cdot(\text{thf})]$ (**H**·(thf)), a compound that was also obtained by reaction of $[\{\text{BDI}^{\text{Dipp}}\}\text{CaN}(\text{SiMe}_3)_2\cdot(\text{thf})]$ with HPPh₂·BH₃.¹⁸

The formulation for complex **4** is reminiscent of that reported for $[\{\text{BDI}^{\text{Dipp}}\}\text{Mg}\{\mu\text{-(H}_3\text{B}\cdot\text{PPh}_2\cdot\text{BH}_3)\}]_2$ (**F** in Fig. 2).¹⁸



Scheme 2 Synthesis of Ca and Ba phosphidoborane and phosphidodiborate complexes **2–4**.



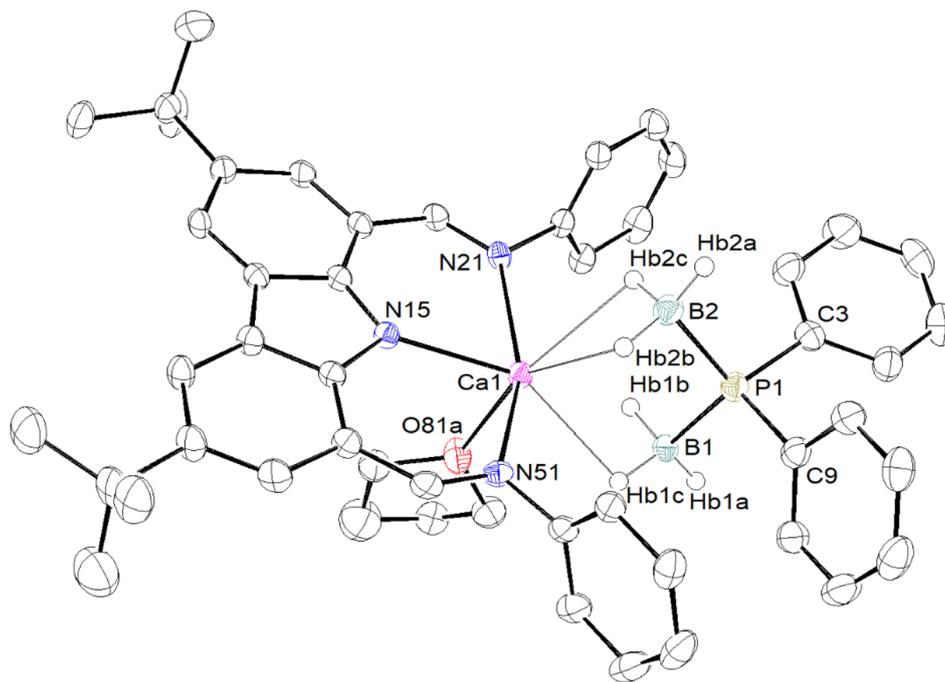


Fig. 4 ORTEP representation of the molecular solid-state structure of $[(\text{Carb})\text{Ca}(\text{H}_3\text{B})_2\text{PPh}_2]\cdot(\text{thf})$ (**4**). Only the major components of disordered ^tBu group and thf molecule are shown. Hydrogen atoms except those bound to boron and ^iPr groups are omitted for clarity. Selected interatomic distances (\AA) and angles ($^\circ$): $\text{Ca1-HB1b} = 2.55(3)$, $\text{Ca1-HB1c} = 2.45(3)$, $\text{Ca1-HB2b} = 2.43(3)$, $\text{Ca1-HB2c} = 2.47(3)$, $\text{Ca1-N15} = 2.385(2)$, $\text{Ca1-N21} = 2.559(3)$, $\text{Ca1-N51} = 2.582(2)$, $\text{Ca1-O81a} = 2.402(2)$, $\text{Ca1-P1} = 3.5483(10)$, $\text{P1-B1} = 1.949(3)$, $\text{P1-B2} = 1.937(3)$, $\text{B1-HB1a} = 1.010(12)$, $\text{B1-HB1b} = 1.066(12)$, $\text{B1-HB1c} = 1.227(12)$, $\text{B2-HB2a} = 1.133(9)$, $\text{B2-HB2b} = 1.139(12)$, $\text{B2-HB2c} = 1.10(3)$; $\text{N15-Ca1-N21} = 79.18(8)$, $\text{N15-Ca1-N51} = 79.58(8)$, $\text{N21-Ca1-N51} = 157.01(7)$, $\text{P1-Ca1-N15} = 155.93(6)$, $\text{P1-Ca1-N21} = 102.59(6)$, $\text{P1-Ca1-N51} = 93.31(6)$, $\text{P1-Ca1-O81a} = 117.07(6)$, $\text{O81a-Ca1-N15} = 86.71(7)$, $\text{O81a-Ca1-N21} = 91.15(8)$, $\text{O81a-Ca1-N51} = 96.32(8)$, $\text{B1-P1-B2} = 109.22(16)$, $\text{B1-P1-C3} = 110.21(14)$, $\text{B1-P1-C9} = 111.82(15)$, $\text{B2-P1-C3} = 110.91(15)$, $\text{B2-P1-C9} = 110.93(14)$, $\text{C3-P1-C9} = 103.67(14)$. Distance from Ca1 to N15N21N51 plane: $0.243(2)$ \AA .

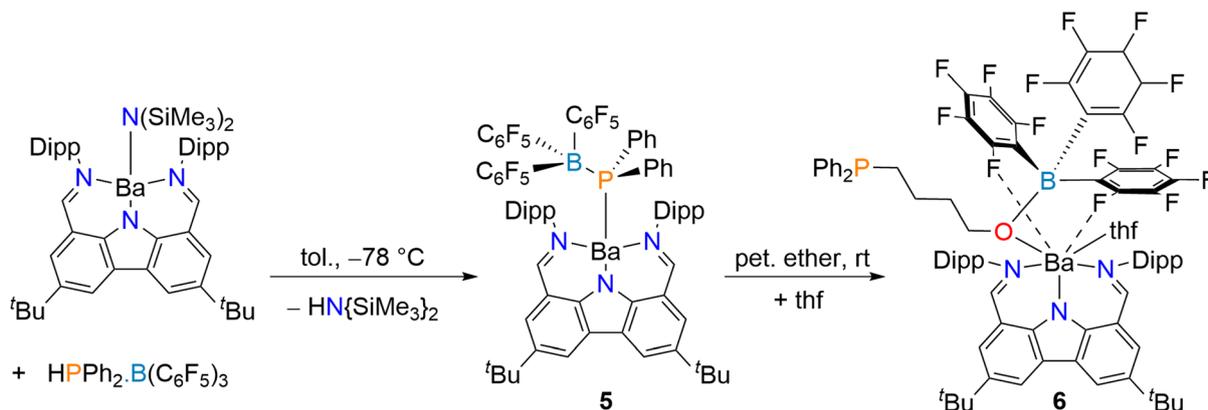
However, the structures of these two complexes differ: whereas the magnesium compound is a dimer with a μ -bridging phosphidodiborate, complex **4** forms a monomeric, thf-solvated complex with a formally six-coordinated, pseudo-octahedral geometry around Ca (Fig. 4). The six boron-bound hydrogens were localised from their electron density in the Fourier difference map. The molecule has four clear $\text{Ca}\cdots\text{H}\cdots\text{B}$ interactions ($2.43(3)$ – $2.55(3)$ \AA) and one coordinated thf molecule. The Ca1-P1 interatomic distance is very long, $3.5483(10)$ \AA , and clearly militates against the existence of a bond between the metal and the phosphorus atom. This is consistent with the near-perfect tetrahedral geometry about P1. By contrast, the Ca-P bond length is much shorter in the four-coordinated heteroleptic $[\{\text{BDI}^{\text{Dipp}}\}\text{CaPPh}_2\cdot(\text{thf})]$ ($2.872(4)$ \AA),⁵ in the six-coordinated $[\text{Ca}(\text{PPh}_2)_2\cdot(\text{thf})_4]$ ($2.9865(6)$ \AA),¹⁴ and in $[\{\text{BDI}^{\text{Dipp}}\}\text{Ca}\{\text{PPh}_2\cdot\text{BH}_3\}\cdot(\text{thf})]$ ($2.8962(4)$ \AA).¹⁸ At a mere $0.243(2)$ \AA above the plane defined by N15, N21 and N51, the calcium centre in **4** fits well inside the sterically shielded cavity created by the bis(imino)carbazolato ligand.

The Mg-phosphidodiborate $[\{\text{BDI}^{\text{Dipp}}\}\text{Mg}\{\mu\text{-}(\text{H}_3\text{B}\cdot\text{PPh}_2\cdot\text{BH}_3)\}_2]$ (**F**) was obtained by addition of two equivalents of $\text{HPPH}_2\cdot\text{BH}_3$ to the dimeric $[\{\text{BDI}^{\text{Dipp}}\}\text{Mg}\{\text{PPh}_2\cdot\text{BH}_3\}_2]$ (**D**) via BH_3 -transfer from $\text{HPPH}_2\cdot\text{BH}_3$.¹⁹ On the other hand, the formation of **4** most likely results from the transfer of a BH_3 moiety from an already coordinated $\{\text{PPh}_2\cdot\text{BH}_3\}^-$ ligand

and concomitant formation of one equivalent of $[\{\text{Carb}\}\text{CaPPh}_2\cdot(\text{thf})]$. Borane transfer of this type has also been documented for lithium compounds, e.g. in the formation of $[\{2\text{-SMe-C}_6\text{H}_4\text{-P}(\text{BH}_3)_2\text{CH}(\text{SiMe}_3)_2\}\text{Li}]$.⁴⁵ The ^{31}P NMR spectrum of **4** displays a singlet at $\delta_{\text{P}} -22.6$ ppm, a significant downfield chemical shift from complex **3** ($\delta_{\text{P}} -41.3$ ppm). The ^{11}B NMR resonance of $\delta_{\text{B}} -32.5$ ppm is slightly upfield from that of **3** ($\delta_{\text{B}} -26.9$ ppm), though similar to Hill's $[\{\text{BDI}^{\text{Dipp}}\}\text{Mg}\{\mu\text{-}(\text{H}_3\text{B}\cdot\text{PPh}_2\cdot\text{BH}_3)\}_2]$ (**F**, $\delta_{\text{B}} -34.3$ ppm).[§]

To circumvent the relative ease of BH_3 migration, $\text{HPPH}_2\cdot\text{B}(\text{C}_6\text{F}_5)_3$ was used due to its potential ability to set up stabilizing $\text{Ba}\cdots\text{F}$ secondary interactions. It is easily available by reaction of HPPH_2 with $\text{B}(\text{C}_6\text{F}_5)_3\cdot(\text{Et}_2\text{O})$ ³⁶ and provides a convenient ^{19}F NMR spectroscopy handle. Addition of one equivalent of the colourless $\text{HPPH}_2\cdot\text{B}(\text{C}_6\text{F}_5)_3$ to a stirred solution of $[\{\text{Carb}\}\text{BaN}(\text{SiMe}_3)_2]$ in toluene at -78 $^\circ\text{C}$ followed by the removal of the volatiles afforded $[\{\text{Carb}\}\text{Ba}\{\text{PPh}_2\cdot\text{B}(\text{C}_6\text{F}_5)_3\}]$ (**5**) in 74% yield, as an extremely moisture-sensitive yellow powder (Scheme 3). The reaction can also be carried out in petroleum ether; however in this case it requires prolonged reaction times. The ^1H NMR spectrum of **5** in toluene- d_8 displays characteristic resonances for the $\{\text{Carb}\}^-$ ligand, a singlet at $\delta_{\text{H}} 8.23$ ($\text{CH}=\text{N}$), a heptet at 2.92 ($\text{CH}(\text{CH}_3)_2$), and two doublets at 1.02 and 0.94 ppm ($^3J_{\text{H-H}} = 6.5$ and 6.7 Hz, $\text{CH}(\text{CH}_3)_2$). The complex is characterised by a single broad resonance in the ^{31}P NMR spectrum at δ_{P}





Scheme 3 Synthesis of [(Carb)Ba(PPh₂·B(C₆F₅)₃)] (5) and [(Carb)Ba(O(B(C₆F₅)₃)(CH₂)₄PPh₂)·thf] (6).

−9.6 ppm, and another broad resonance in the ¹¹B NMR spectrum (δ_B −12.6 ppm). Both values are substantially deshielded compared to those for 3 (δ_P −41.3 and δ_B −26.9 ppm), in agreement with the strongly electron-withdrawing nature of C₆F₅ groups. Three main resonances are visible in the ¹⁹F NMR spectrum (δ_F −128.0, −160.3 and −164.7 ppm for *o*-, *p*- and *m*-F atoms, respectively). The data for 5 are also different from those of the starting material, HPPh₂·B(C₆F₅)₃ (δ_P −0.4 ppm; δ_B −12.4 ppm; δ_F −128.1, −155.2 and −162.9 ppm). Taken collectively, these NMR data all point at the formation of a single product with a unique coordination sphere. Despite repeated recrystallisation attempts from a range of solvents, we could not obtain crystals of 5 suitable for XRD characterisation. Surprisingly, at the time of writing, there was no case of a metal complex containing the “[Met]–PPh₂·B(C₆F₅)₃” fragment in the CSD database,⁴⁶ although the Ti(IV) phosphide (Cp)(η^5 -C₅H₄B(C₆F₅)₂)(μ -PPh₂)TiCl containing a Cp–B–P chelate was reported.⁴⁷ The structure of 5 was then assessed by DFT calculations (Fig. 5). In the optimised geometry, the central metal is unsymmetrically five-coordinated by phosphorus, the three nitrogen atoms from the bis(imino)carbazolato ligand and one fluorine atom (Ba...F = 2.863 Å). The computed Ba–N interatomic distances are somewhat shorter than in 1·(thf)₂ (see Table 1), whereas the Ba–P distance is larger (3.303 Å). Nonetheless, the EDA analysis (Table 2) does not suggest weaker bonding. This apparent contradiction finds its source in the fact that the considered metal–ligand interaction in 5 includes also one Ba...F contact. The Ba–P WBIs (Table 1) and bep indicators (Table 3) are consistent with a somewhat weaker bond in 5 as compared to 1·(thf)₂. The nature of Ba...F bonding is discussed in more detail below in the case of complexes 6 and 8.

The addition of a small amount of thf to a suspension of 5 in petroleum ether did not produce the expected [(Carb)Ba{PPh₂·B(C₆F₅)₃}(thf)_{*n*}], nor did it form a phosphidodiborate akin to 4, e.g. [(Carb)Ca{PPh₂(B(C₆F₅)₃)₂}(thf)_{*n*}]. Instead, [(Carb)Ba{O(B(C₆F₅)₃)CH₂CH₂CH₂CH₂PPh₂}(thf)] (6), a Ba–alkoxide resulting from the ring-opening of one thf molecule, was isolated in 61% yield as yellow crystals (Scheme 3). Complex 6

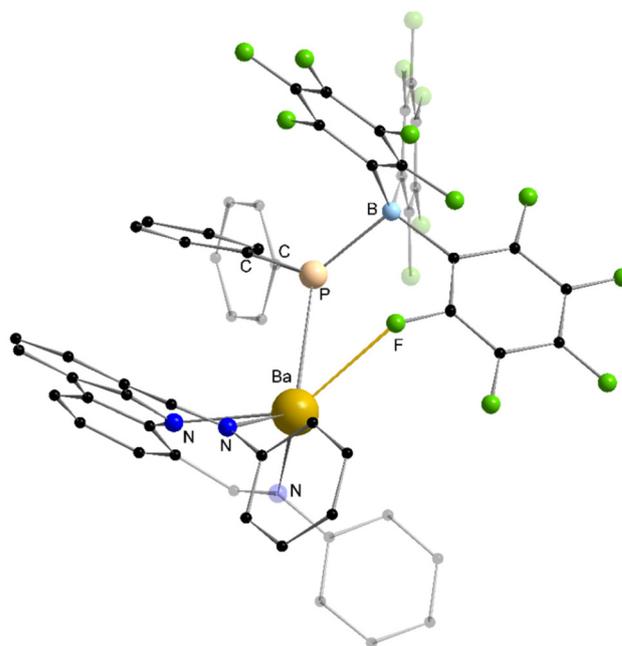


Fig. 5 The DFT-optimised geometry of [(Carb)Ba{PPh₂·B(C₆F₅)₃}] (5). See Table 1 for selected interatomic distances. Additional metrical data: P–B = 2.118 Å, B–C (av.) = 1.641 Å; P–Ba–F = 56°.

is a rare example of a heteroleptic, monometallic barium alkoxide. The ring-opening of thf has been previously reported for a number of systems built around oxophilic metals,^{25–30} frustrated Lewis pairs based on phosphines^{31–34} or NHCs,⁴⁸ even middle-to-late transition metals.^{49,50} The formation of 6 presumably occurs *via* nucleophilic attack by the phosphorus atom on the thf α -carbon atom, resulting in ring opening, while the increased negative charge on the now Ba-coordinated oxygen atom also results in borane migration from the phosphine to generate a more stable Lewis adduct.

The ¹H NMR spectrum of complex 6 in benzene-*d*₆ displays characteristic ligand resonances at δ_H 8.47 (CH=N), 3.00 (CH(CH₃)₂), and four isopropyl resonances at δ_H 1.06, 1.00 and



0.92 (two superimposed) ppm ($\text{CH}(\text{CH}_3)_2$), as well as four methylene resonances at δ_{H} 2.85 (OCH_2), 1.22, 0.56 and -0.28 ppm. It also shows the presence of the protonated ligand {Carb}H which we could not remove. Yet, the ^{31}P NMR spectrum displays a single resonance at δ_{P} -18.8 ppm, an upfield chemical shift from 5, while the $^{11}\text{B}\{^1\text{H}\}$ NMR spectrum has a single broad resonance at δ_{B} -2.6 ppm. This is consistent with the existence of single $[\text{Ba}]\text{-PPh}_2/\text{B}(\text{C}_6\text{F}_5)_3$ species. Both resonances are similar to those for the lithium alkoxide-bound borane $[\text{Li}\{\text{O}(\text{B}(\text{C}_6\text{F}_5)_3)(\text{CH}_2)_4\text{PMe}_2\}]$ (Mes = mesityl; δ_{B} -3.1 and δ_{P} -21.4 ppm).²⁵ Interestingly, the ^{19}F NMR spectrum displays eight resonances in the δ_{F} range -132 to -165 ppm instead of the usually observed three signals. For instance, $[\text{Li}\{\text{O}(\text{B}(\text{C}_6\text{F}_5)_3)(\text{CH}_2)_4\text{PMe}_2\}]$ features the common doublet at δ_{F} -137.6 ppm and triplets at -161.0 and -165.6 ppm. The unusual ^{19}F NMR pattern for **6** suggests that at room temperature, one or more *ortho*-fluorine $\text{Ba}\cdots\text{F}\text{-C}$ interactions inhibit the free rotation of one or two of the C_6F_5 rings in the complex.

The crystal structure of **6** (Fig. 6) reveals the metal to be in a seven-coordinated environment, with coordination of the five N- and O-atoms, along with two clearly identifiable $\text{Ba}\cdots\text{F}\text{-C}$ interactions. At $7.696(2)$ Å, the phosphorus atom P1 is

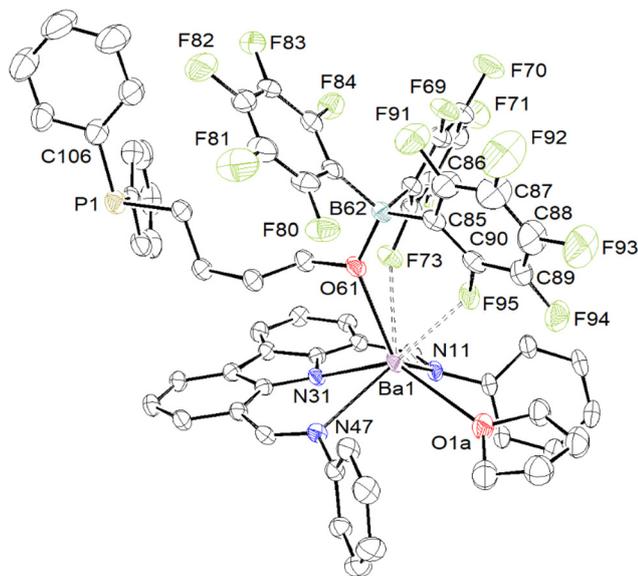


Fig. 6 ORTEP representation of the molecular structure of $[(\text{Carb})\text{Ba}(\text{O}(\text{B}(\text{C}_6\text{F}_5)_3)\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{PPh}_2)\cdot\text{thf}]$ (**6**). H atoms, ^tBu are ^iPr groups are omitted for clarity. Only the major component of the disordered thf molecule is shown. Selected interatomic distances (Å) and angles ($^\circ$): $\text{Ba1-F73} = 2.9889(17)$, $\text{Ba1-F95} = 2.7497(17)$, $\text{Ba1-N11} = 2.798(2)$, $\text{Ba1-N31} = 2.658(2)$, $\text{Ba1-N47} = 2.772(3)$, $\text{Ba1-O1a} = 2.788(2)$, $\text{Ba1-O61} = 2.789(2)$, $\text{C64-F69} = 1.350(4)$, $\text{C65-F70} = 1.340(4)$, $\text{C68-F73} = 1.363(3)$, $\text{C86-F91} = 1.345(4)$, $\text{C88-F93} = 1.338(4)$, $\text{C90-F95} = 1.374(3)$; $\text{N11-Ba1-N31} = 70.07(7)$, $\text{N11-Ba1-N47} = 124.13(7)$, $\text{N11-Ba1-O1a} = 89.19(8)$, $\text{N11-Ba1-O61} = 127.13(7)$, $\text{N31-Ba1-N47} = 70.12(7)$, $\text{N31-Ba1-O1a} = 134.80(7)$, $\text{N31-Ba1-O61} = 98.75(7)$, $\text{N47-Ba1-O1a} = 92.14(7)$, $\text{N47-Ba1-O61} = 96.33(7)$, $\text{O1a-Ba1-O61} = 124.97(7)$. Distance from Ba1 to N11N31N47 plane: $1.2349(4)$ Å.

expulsed far away from the metal Ba1. The Ba1-F95 interatomic distance of $2.7497(17)$ Å testifies to a strong interaction with the fluorine atom, likely one that persists in solution, while the interaction with F73 is slightly weaker ($\text{Ba1-F73} = 2.9889(17)$ Å). The $\text{Ba}\cdots\text{F95}$ distance is at the very low end of values for $\text{Ba}\cdots\text{F}$ secondary interactions, more commonly in the range $2.80\text{--}3.10$ Å. It is for instance significantly shorter than in other previously reported complexes with strong $\text{Ba}\cdots\text{F}\text{-C}$ contacts, such as the bis(hydridoborate) $[\text{Ba}\{\text{HB}(\text{C}_6\text{F}_5)_3\}_2(\text{thf})_4]$ ($2.8228(19)\text{--}2.8386(17)$ Å),⁵¹ the dimeric barium complex $[\text{Ba}\{\mu\text{-N}(\text{C}_6\text{F}_5)_2\}\{\text{N}(\text{C}_6\text{F}_5)_2\}(\text{toluene})_2]$ ($2.825(2)\text{--}3.014(3)$ Å),⁵² or the polymeric $[\text{Ba}\{\text{N}(\text{H})\text{-}2,6\text{-F}_2\text{C}_6\text{H}_3\}_2(\text{thf})_2]_\infty$ ($2.871(2)\text{--}2.901(3)$ Å).⁵³ This short interatomic distance is consistent with the aforementioned ^{19}F NMR data showing inequivalence at room temperature. As relative points of comparison, the sum of van der Waals radii for barium and fluorine (2.68 and 1.47 Å, respectively)⁵⁴ is much greater than the Ba1-F73 and Ba1-F95 distances, whereas the added covalent radii for Ba and F amount to 2.72 Å. Note that it is now established that the nature of such $\text{Ba}\cdots\text{F}$ (and, more generally, $\text{Ae}\cdots\text{F}$) interactions is essentially (albeit not exclusively) electrostatic.^{52,55} Yet, the C68-F73 ($1.363(3)$ Å) and especially the C90-F95 ($1.374(3)$ Å) bonds are stretched compared to the other C-F bonds with fluorine atoms that do not interact with the metal, typically between $1.338(4)\text{--}1.350(4)$ Å. Another notable feature in the structure of **6** is the long Ba1-O61 interatomic distance ($2.788(2)$ Å). Remarkably, it is even longer than the shortest $\text{Ba}\cdots\text{F}$ contact in the complex ($2.7497(17)$ Å). It is significantly greater than for other terminal alkoxides, such as the perfluorinated alkoxides $[\text{Ba}\{\text{OC}(\text{CF}_3)_2(\text{donor})_n\}]$ (donor = thf, diglyme or 1,2-dimethoxyethane; $\text{Ba-O} = 2.5335(9)\text{--}2.544(3)$ Å),⁵⁶ $[\text{Ba}_2(\text{OCPh}_3)(\mu\text{-OCPh}_3)_3(\text{thf})_3]$ ($2.409(4)$ Å),⁵⁷ and even the eight-coordinated $[\text{Ba}\{\text{OC}(\text{CF}_3)_2\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2\text{OME})_2\}_2]$ ($2.558(2)\text{--}2.562(2)$ Å).⁵⁸ The $\text{Ba-O}_{\text{alkoxide}}$ distance in **6** is also much longer than in the related O-ligated barium-boryloxide $[\text{Ba}\{\text{OB}(\text{CH}(\text{SiMe}_3)_2)_2\}_2]$ ($2.3832(16)\text{--}2.4098(16)$ Å)⁵⁹ and barium-siloxide $[\text{Ba}_2\{\text{OSi}(\text{SiMe}_3)_3\}\{\mu\text{-OSi}(\text{SiMe}_3)_3\}_3]$ ($2.414(7)$ Å)⁶⁰ and $[\text{Ba}_2\{\text{OSi}^i\text{Bu}_3\}\{\mu\text{-OSi}^i\text{Bu}_3\}_3]$ ($2.413(16)$ Å)⁵⁷ complexes, despite the reduced donating ability of the O-atoms in these ligands.⁶¹ The $\text{Ba-O}_{\text{alkoxide}}$ distance is still greater in **6** than in the eight-coordinated $[\text{Ba}\{\text{OCH}_2\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2\text{OH})_2\}_2]\cdot\text{EtOH}$ ($2.73(1)$ Å),⁶² the first (and a very rare) structurally verified $\text{Ba}\text{-alkoxide}$ featuring an unexpectedly long Ba-O length due to $\text{EtOH}\cdots\text{OBa}$ hydrogen bonding with lattice ethanol molecules. These comparisons are not ideal due to the structural differences between these compounds and **6**. However, owing to the lack of structurally characterised monometallic barium alkoxides, there is no directly relevant comparison to be made at the current time.⁴⁶ The long Ba1-O61 interatomic distance in **6** likely results from the transfer of electron density from the oxygen atom into the empty p-orbital of the borane. There is a distorted trigonal pyramidal geometry around the O61 atom ($\Sigma_{\theta}(\text{O61}) = 346.8(3)^\circ$). Such relative weakness of the Ba-O bond does raise the question of whether the alkoxide could be substituted for another ligand *via* protonolysis, and, if so, whether



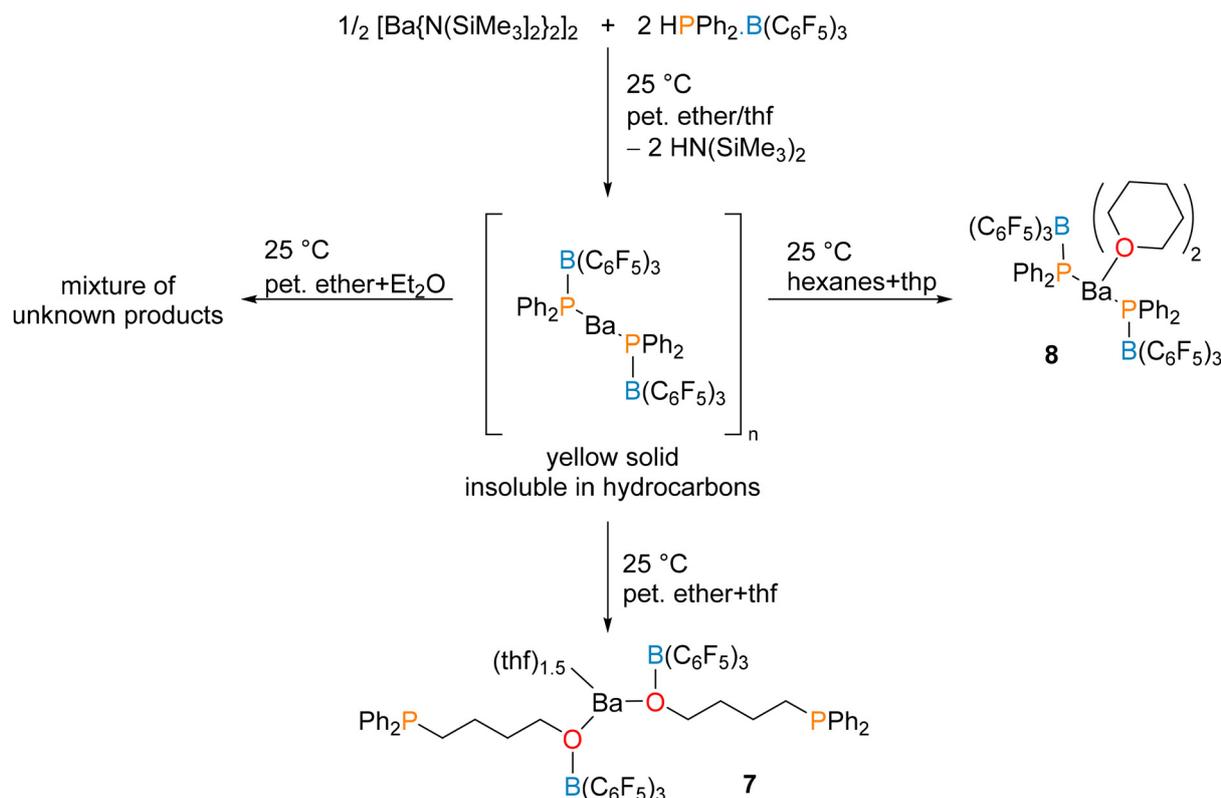
the ring-opening of thf could be turned advantageously into a catalytic manifold.

DFT calculations on complex **6** reproduce well its solid-state structure (ESI, Table S1[†]), with the shortest optimised Ba–F distance (2.813 Å) being slightly longer than its X-ray counterpart. In any case, both Ba...F contacts show similar characteristics, hence they are averaged in Table S1.[†] Their QTAIM descriptors are diagnostic of weak ionic bonds. Such Ae...F–C interactions have been investigated in low-coordinate Ae fluoroarylamides.⁵² Their very small covalent characteristic is also evidenced by the fact that the corresponding F-atoms are more negatively charged (–0.36 (natural), –0.69 (QTAIM), see Table S1[†]) than the other fluorines (range: –0.29 to –0.31 (natural), –0.65 to –0.67 (QTAIM)), indicating charge attraction from Ba onto the fluorine atoms, but very little electron transfer between them.

To further investigate the reactivity of the [Ba]–PPh₂–B(C₆F₅)₃ moiety and the nature of the Ba-to-alkoxide bonding in **6**, the analogous homoleptic barium phosphidoborane starting material was synthesised. The protonolysis reaction between [Ba{N(SiMe₃)₂}₂]₂ and HPPH₂–B(C₆F₅)₃ (two equivalents vs. Ba) yields a pale yellow powder, presumed to be [Ba{PPh₂–B(C₆F₅)₃}₂]_n (Scheme 4). This solid is insoluble in aromatic or aliphatic hydrocarbons. Dissolution by addition of thf to a suspension in petroleum ether gives the product of composition [Ba{O(B(C₆F₅)₃)CH₂CH₂CH₂CH₂PPh₂}₂·(thf)_{1.5}] (7, probably an aggregate), where ring-opening of thf has

occurred. This compound could not be crystallised to allow XRD analysis.[§] Although it is a white solid, crystallisation attempts generated yellow clathrates that separated from the solvent, especially at low temperatures. By contrast, the six-member cyclic tetrahydropyran (thp) does not ring-open under the same conditions, and affords instead the thp-adduct [Ba{PPh₂–B(C₆F₅)₃}₂·(thp)₂] (**8**), which was characterised by NMR (Fig. S30–S35[†]) and XRD (Fig. 7) analyses.[§]

The ¹H NMR spectrum of **7** in thf-*d*₈ displays the resonances for four distinct methylene groups, along with the expected aromatic resonances for the phenyl moieties as two sets of multiplets at δ_H 7.34 (8H) and 7.23 (12H) ppm. The spectrum is consistent with the presence of 1.5 equivalents of non-deuterated thf per metal. The ³¹P NMR spectrum shows a broad singlet at δ_P –16.1 ppm, similar to the chemical shift in **6**, while the ¹¹B NMR spectrum presents a broad singlet at δ_B –2.9 ppm. It is probable that in the solid state, compound **7** creates a larger multimetallic species with bridging alkoxide ligands.^{12,57} The ¹H NMR spectrum of **8** in thf-*d*₈ displays resonances near-identical to those of **7** in this solvent, indicating that deuterated thf molecules displace the initially metal-bound thp ones in solution. Accordingly, three resonances for free thp are also observed. The NMR data confirm that complex **8** is the sole observable species. The ³¹P NMR spectrum displays a single resonance at δ_P –16.6 ppm, while the ¹¹B NMR spectrum displays a sharp resonance at δ_B –2.9 ppm and the ¹⁹F NMR spectrum has three resonances, a doublet at



Scheme 4 Synthesis of [Ba{O(B(C₆F₅)₃)CH₂CH₂CH₂CH₂PPh₂}₂·(thf)_{1.5}] (**7**) and [Ba{PPh₂–B(C₆F₅)₃}₂·(thp)₂] (**8**).



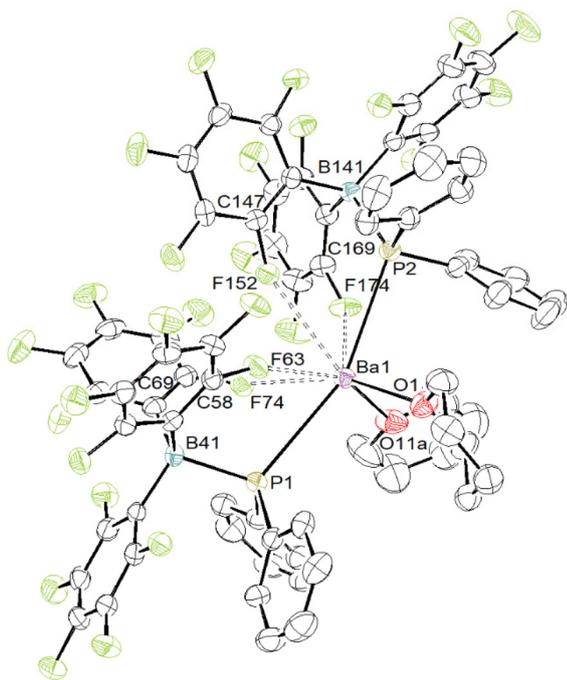


Fig. 7 Representation of the molecular solid-state structure of $[\text{Ba}(\text{PPH}_2)_2\text{B}(\text{C}_6\text{F}_5)_3]_2(\text{thp})_2$ (**8**). Non-interaction lattice toluene molecule and H atoms are omitted for clarity. Only the main component of the disordered thp molecule is shown. Non-interacting F-atoms are shaded in green. Selected interatomic distances (Å) and angles ($^\circ$): Ba1–F63 = 2.7557(17), Ba1–F74 = 2.7634(19), Ba1–F152 = 3.1120(18), Ba1–F174 = 2.7912(18), Ba1–O1 = 2.623(2), Ba1–O11a = 2.658(2), Ba1–P1 = 3.3080(8), Ba1–P2 = 3.2495(8), P1–B41 = 2.071(3), P2–B141 = 2.050(3), C58–F63 = 1.375(3), C69–F74 = 1.379(4), C147–F152 = 1.370(3), C169–F174 = 1.378(4); O1–Ba1–O11a = 97.31(8), O1–Ba1–P1 = 98.22(6), O1–Ba1–P2 = 87.00(6), O11a–Ba1–P1 = 96.21(6), O11a–Ba1–P2 = 106.93(6), P1–Ba1–P2 = 155.47(2).

$\delta_{\text{F}} -133.4$ ($^3J_{\text{F-F}} = 23.7$ Hz) ppm, a sharp triplet at -165.6 ($^3J_{\text{F-F}} = 20.1$ Hz) and a broad triplet at -168.4 ($^3J_{\text{F-F}} = 19.8$ Hz) ppm. All these data are almost identical to those of **7**. Finally, when diethyl ether is added instead of a cyclic ether to a suspension of $[\text{Ba}\{\text{PPH}_2\text{-B}(\text{C}_6\text{F}_5)_3\}_2]_n$ in petroleum ether, the metal species decomposes over the course of a few hours and the precipitation of a black solid is visible. The NMR analysis of the resulting mixture did not provide any insight into the decomposition process.

The solid-state structure of complex **8** depicted in Fig. 7 shows the metal to sit in an eight-coordinate geometry from coordination of the two P- and two O-atoms, along with four additional Ba...F contacts. The complex shows intriguing structural parameters, but there is no [Met]– $\text{PR}_2\text{B}(\text{C}_6\text{F}_5)_3$ metal fragment in the CSD database to allow for a direct comparison at the time of writing.⁴⁶ The P1–Ba1–P2 angle is very obtuse (155.47(2) $^\circ$). Approximately one half of the first coordination sphere around the metal is occupied by Ba...F interactions, whereas the other half is filled by the two coordinated thp molecules and phenyl rings. The Ba1–P1 and Ba1–P2 interatomic distances (3.2495(8)–3.3080(8) Å) are commensurate with those in $[\text{Ba}(\text{PPH}_2)_2\cdot(\text{thf})_5]$ (3.328(2) and 3.345(2) Å)¹⁴ and

in the polymeric $[\text{Ba}_3(\text{PPH}_2)_6\cdot(\text{thf})_4]_\infty$ (3.2958(7)–3.3627(7) Å).¹³ This bonding situation suggests that coordination of the borane onto phosphorus does not affect significantly its electron-sharing properties with the metal. The geometry about the P- and B-atoms forms slightly distorted tetrahedra. The P1–Ba1 and P2–B141 bond lengths (2.071(3) and 2.050(3) Å) are greater than in the calcium complexes $[\{\{\text{BDI}^{\text{Dipp}}\}\text{Ca}\}_3(\mu_2\text{-H})(\text{H}_3\text{BPPH}_2)_2]$ (**G** in Fig. 2, 1.945(3) and 1.946(3) Å) and $[\{\text{BDI}^{\text{Dipp}}\}\text{Ca}\{\text{PPH}_2\text{-BH}_3\}\cdot(\text{thf})]$ (**H**·(thf), 1.9466(18) Å).¹⁸ They compare somewhat better with the P–B(C_6F_5)₃ distance in $[\{\text{BDI}^{\text{Dipp}}\}\text{Ca}\{\text{H}_3\text{B-PPH}_2\text{-B}(\text{C}_6\text{F}_5)_3\}\cdot(\eta^6\text{-toluene})]$ (**I**, 2.1368(18) Å), while the H₃B–P distance to the κ^3 -bound BH₃ group is shorter (1.9469(18) Å). However, the phosphorus atom is not metal-bound in this compound.¹⁹ The P–B interatomic distance is 2.054(4) Å in the chelate complex $(\text{Cp})(\eta^5\text{-C}_5\text{H}_4\text{B}(\text{C}_6\text{F}_5)_2)(\mu\text{-PPH}_2)\text{TiCl}$ where bonding involves a Ti–P σ -bond and a P \rightarrow B donor interaction (Fig. 8),⁴⁷ and where the closest Ti...F distances in this complexes are above 3.6 Å. The Ba1–O1 and Ba1–O11a bond lengths in **8** (2.623(2) and 2.658(2) Å, respectively) are substantially shorter than in the five-coordinated $[\text{Ba}\{\text{N}(\text{SiMe}_3)_2\}_2\cdot(\text{thp})_3]$ (2.829(2)–2.869(2) Å)⁶³ and in the seven-coordinated $[\text{BaI}_2\cdot(\text{thp})_5]$ (2.723(6)–2.789(6) Å).⁶⁴ Finally, three of the four Ba...F interactions in **8**, in the range 2.7557(17)–2.7912(18) Å, compare well with those in complex **6** and can be considered as particularly strong. The last one, with a Ba1–F152 interatomic distance of 3.1120(18) Å, is weaker. As in **6**, the C–F bond lengths for these four metal-interacting fluorine atoms, between 1.370(4) and 1.379(4) Å, are elongated compared to the other C–F bonds (1.342(4)–1.357(4) Å) in the compound. Besides, shortening of the Ba–F interatomic distance is accompanied by a lengthening of the corresponding C–F bond. It is not clear how heavily the multiple Ba...F contacts in **8** weigh on the other metric parameters and bonding properties in the complex, notably concerning the barium-to-phosphorus bonds.

The X-ray structure of complex **8** is well reproduced by DFT calculations (ESI, Table S2[†]). In particular, the four short Ba...F contacts (2.854 Å in average) are also present in the gas phase-optimised geometry. The Ba–P and Ba...F bond indicators (Table S2[†]) are similar to those found in **1**·(thf)₂ and **6**, respectively. In order to compare the strength of the Ba–P bonds in **8** with that in **1**·(thf)₂, an energy decomposition analysis was performed on **8**, based on the $[\text{Ba}(\text{thp})_2]^{2+} + \{\{\text{PPH}_2\text{-B}(\text{C}_6\text{F}_5)_3\}^-\}_2$ fragmentation (see Table 4). It should be pointed out that the reported bonding energies are not only associated

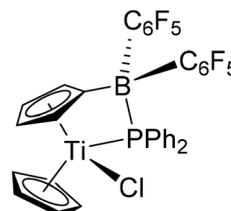


Fig. 8 Structure of $(\text{Cp})(\eta^5\text{-C}_5\text{H}_4\text{B}(\text{C}_6\text{F}_5)_2)(\mu\text{-PPH}_2)\text{TiCl}$.⁴⁷



Table 4 Morokuma–Ziegler energy decomposition analysis in [Ba{PPh₂B(C₆F₅)₃}₂(thp)₂] (**8**) and its H-substituted derivative [Ba{PPh₂B[(C₆H₂F₃)(C₆F₅)₂]}₂(thp)₂] (see text)^a

	[Ba{PPh ₂ B(C ₆ F ₅) ₃ } ₂ (thp) ₂] (8)	[Ba{PPh ₂ B[(C ₆ H ₂ F ₃)(C ₆ F ₅) ₂]} ₂ (thp) ₂] (single point, see text)
Fragmentation	[Ba·(thp) ₂] ²⁺ + {PPh ₂ B[(C ₆ F ₅) ₃]} ₂ ⁻	[Ba·(thp) ₂] ²⁺ + {PPh ₂ B[(C ₆ H ₂ F ₃)(C ₆ F ₅) ₂]} ₂ ⁻
<i>E</i> _{Pauli}	3.22	3.10
<i>E</i> _{elstat}	-11.74	-10.48
<i>E</i> _{orb}	-4.64	-4.24
<i>E</i> _{disp}	-1.21	-1.19
TBE ^b	-14.38 (332 kcal mol ⁻¹)	-12.82 (296 kcal mol ⁻¹)

^a Values in eV, unless specified. ^b Total bonding energy (TBE) = *E*_{Pauli} + *E*_{elstat} + *E*_{orb} + *E*_{disp}.

with two Ba–P bonds, but also with four Ba...F contacts. In order to separate these two contributions, we also ran a single-point calculation on the frozen structure of a molecule of **8** in which the four F-atoms in contact with Ba were replaced by H-atoms (C–H = 1.09 Å). The corresponding EDA decomposition based on the [Ba·(thp)₂]²⁺ + {PPh₂B[(C₆H₂F₃)(C₆F₅)₂]}₂⁻ fragmentation is also given in Table 4. Its TBE values should approximate twice the Ba–P bonding energy and the difference with the TBE value in **8** should approximate four times the Ba...F bonding energy. From there, it ensues that the resulting Ba–P and Ba...F bond energies in **8** are ~148 and ~9 kcal mol⁻¹, respectively. It thus appears that the Ba–P bond in **8** is substantially stronger than in **5** (104–9 = 95 kcal mol⁻¹) and in 1-(thf)₂ (106 kcal mol⁻¹, see Table 2). It should be noted that replacing all F atoms in **8** by hydrogens results in an optimised geometry somehow different from that of **8**, exhibiting a weak Ba-(η²-C₆H₅) interaction in addition to several Ba...H contacts. Finally, replacing the two thp ligands in **8** by thf molecules produces an optimised geometry with very similar metric data and electronic structure, indicating that the unsuccessful isolation of the **8**(thf)₂ relative is only the result of kinetic factors.

Conclusion

Our initial goal, the preparation of soluble and stable Ba-phosphides, in particular supported by the {Carb}⁻ framework, has proved elusive. Compounds such as [{Carb}BaPPh₂(thf)] (**1**) and its thf adducts do not display sufficient stability in solution and can therefore not be used as catalysts for C–P bond forming reactions, such as the hydrophosphination of alkenes where other Ba precatalysts have, on the whole, exhibited very good performances.

On the other hand, we have been able to prepare and, in some cases, crystallise, several barium and calcium phosphidoboranes using the “disguised” phosphines HPPH₂·BH₃ and HPP₂B(C₆F₅)₃. All these complexes are very air-sensitive, and their stability in solution can be limited, which has in some cases hampered NMR analyses. Their structural features are unusual, in particular for complexes such as [{Carb}Ba{PPh₂B(C₆F₅)₃}] (**5**), [{Carb}Ba{O(B(C₆F₅)₃)CH₂CH₂CH₂CH₂PPh₂}]·thf (**6**) and the remarkable [Ba{PPh₂B(C₆F₅)₃}₂(thp)₂] (**8**) that display strong intramolecular Ba...F secondary interactions.

Although BH₃ was involved in unwanted mobility eventually leading to the formation of [{Carb}Ca{(H₃B)₂PPh₂}]·(thf) (**4**), we found that B(C₆F₅)₃ produced stable Ba-phosphidoboranes where the transfer of the electron-depleted borane did not take place. Besides, in a way reminiscent of the behaviour exhibited by frustrated Lewis pairs, the latter compounds featured the ability to ring-open thf, and, in the presence of this solvent, to yield alkoxide derivatives such as **6** and [Ba{O(B(C₆F₅)₃)CH₂CH₂CH₂CH₂PPh₂}₂(thf)_{1.5}] (**7**). No such reactivity was found when the more stable thp was used instead of thf, and we were hence able to isolate and characterise the bis(thp) adduct **8**. We are now trying to exploit this reactivity and assess whether it is possible to turn it into a catalytic cycle for the production of P-functionalised alcohols.

DFT calculations performed to analyse the bonding and stability of the barium complexes do not reveal any particular weakness of the Ba–P bond in 1-(thf)₂. The instability of this compound is better understood from the high energy of its HOMO, which contains the non-conjugated P lone pair, thus conferring it substantial nucleophilic reactivity. The structure of **5**, assessed from DFT calculations, shows non-negligible Ba...F bonding. Calculations indicate that these ionic interactions, also present in **6** and **8**, contribute significantly to the stabilisation of such architectures (Ba...F bond energy ~9 kcal mol⁻¹).

Author contributions

G. Duneş and P. M. Chapple performed the experimental and analytical work and participated in the planning of experiments, analysis of results and writing of the manuscript. S. Kahlal and J.-Y. Saillard performed the DFT calculations and participated in the preparation of the manuscript. T. Roisnel solved the crystallographic structures and prepared the XRD data. J.-F. Carpentier participated in the planning of experiments and analysis of the data. Yann Sarazin was the lead investigator, secured the funding, participated in the planning of experiments, data analysis and writing of the manuscript.

Conflicts of interest

There are no conflicts to declare.



Acknowledgements

The French Agence Nationale de la Recherche is acknowledged for the provision of research grants to P. M. C. (ANR-17-CE07-0017-01) and to G. D. (ANR-21-CE07-0045-02).

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