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Magnesium–halobenzene bonding: mapping the halogen sigma-hole with a Lewis-acidic complex†

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Complexes of the Lewis base-free cations (^{Me}BDI)Mg⁺ and (^{tBu}BDI)Mg⁺ with Ph–X ligands (X = F, Cl, Br, I) have been studied (^{Me}BDI = HC[C(Me)N-DIPP]₂ and ^{tBu}BDI = HC[C(tBu)N-DIPP]₂; DIPP = 2,6-diisopropylphenyl). For the smaller β-diketiminato ligand (^{Me}BDI) only complexes with PhF could be isolated. Heavier Ph–X ligands could not compete with bonding of Mg to the weakly coordinating anion B(C₆F₅)₄[−]. For the cations with the bulkier ^{tBu}BDI ligand, the full series of halobenzene complexes was structurally characterized. Crystal structures show that the Mg⋯X–Ph angle strongly decreases with the size of X: F 139.1°, Cl 101.4°, Br 97.7°, I 95.1°. This trend, which is supported by DFT calculations, can be explained with the σ-hole which increases from F to I. Charge calculation and Atoms-In-Molecules analyses show that Mg⋯F–Ph bonding originates from electrostatic attraction between Mg²⁺ and the very polar C^{δ+}–F^{δ−} bond. For the heavier halobenzenes, polarization of the halogen atom becomes increasingly important (Cl < Br < I). Complexation with Mg leads in all cases to significant Ph–X bond activation and elongation. This unusual coordination of halogenated species to early main group metals is therefore relevant to C–X bond breaking.

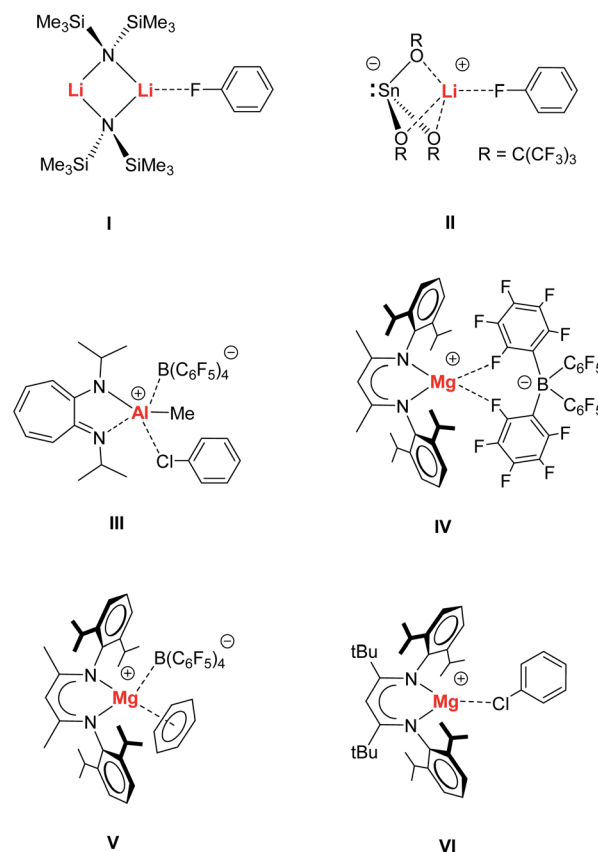
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Introduction

Halogenated benzenes (Ph–X, X = F, Cl, Br, I) are key building blocks in organic syntheses and also often used as polar but weakly coordinating solvents. This last property explains the scarcity of metal⋯X–Ph complexes. There is only a handful of well-characterized metal⋯X–Ph complexes.^{1,2} Despite the availability of three lone-pairs of electrons, Ph–X is a very weak Lewis base and electron donor. This is illustrated by the fact that its oxidized form, Ph–X=O, is only known for iodosobenzene (X = I) and even this compound is highly explosive.³ In the series Ph–X (X = F, Cl, Br, I), iodobenzene is likely the best electron pair donor especially for the softer, larger transition metals. It is, however, also most susceptible to C–X bond cleavage and indeed the formation of transition metal⋯X–Ph complexes is proposedly the first step towards oxidative addition. Metal–halobenzene complexes are crucial to C–X bond activation and catalysis^{4,5} and especially of great importance in the challenging activation of strong C–F bonds.^{2,6}



Main group metal⋯X–Ph complexes are even rarer and most examples deal with metal bonding of the most polar halobenzene Ph–F, e.g. I⁷ or II.⁸ Metal bonding of the heavier

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halobenzenes is much more uncommon. While metal...Cl-Ph complexes have been identified for Re, Zr and Rh,^{9a-d} the cationic Al complex **III** represents a rare example for binding of a main group metal to a heavier halobenzene.^{9e} Complexation of Ph-Br and Ph-I is limited to the softer metals Pd, Pt and Ag¹⁰ and to the best of our knowledge, main group complexes with such ligands are unknown.

We recently introduced syntheses and structures of cationic, Lewis base-free, alkaline earth metal complexes.¹¹⁻¹⁶ The cationic β -diketiminate Mg complex $[(^{\text{Me}}\text{BDI})\text{Mg}^+][\text{B}(\text{C}_6\text{F}_5)_4^-]$ (**IV**) shows a chelating $\text{B}(\text{C}_6\text{F}_5)_4^-$ anion but cation-anion association is weak and $(^{\text{Me}}\text{BDI})\text{Mg}^+$ is more Lewis acidic than $(^{\text{Me}}\text{BDI})\text{AlMe}^+$ or the benchmark Lewis acid $\text{B}(\text{C}_6\text{F}_5)_3$.¹¹ Highly Lewis acidic **IV** was shown to be able to form π -arene complexes (e.g. **V**),^{11,13-15} alkynes complexes^{11,16} and even a complex of the silyl ether $\text{O}(\text{SiMe}_3)_2$, that is normally fully inert to metal bonding, has been isolated.¹²

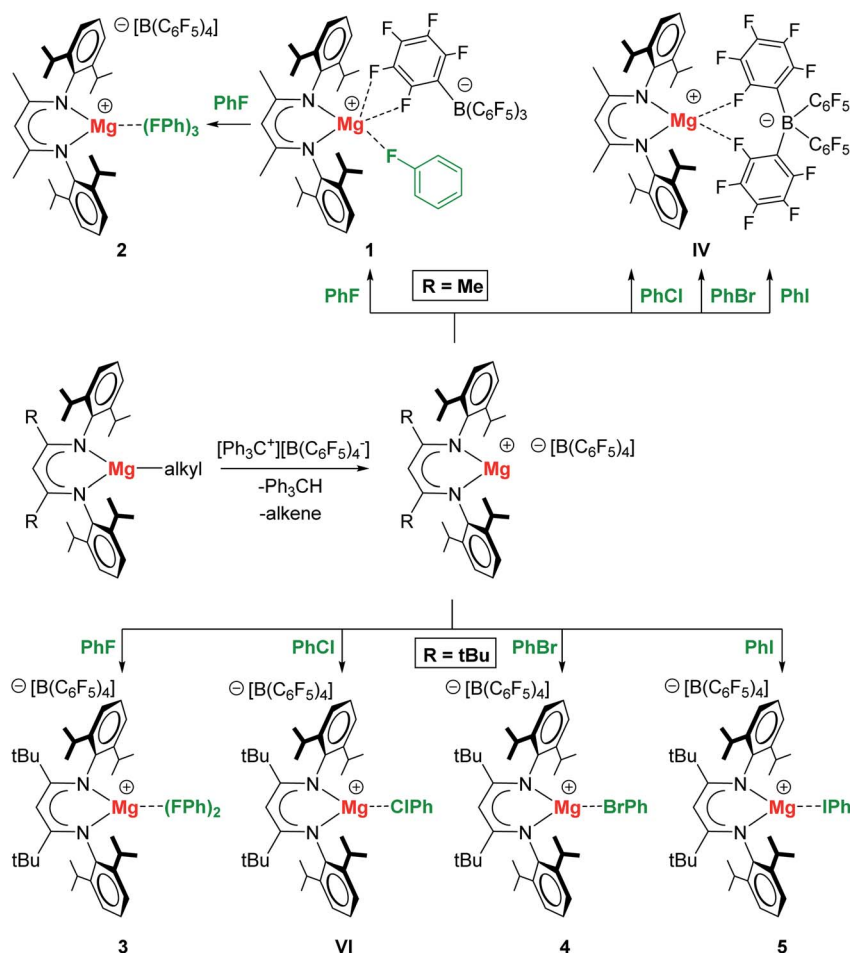
While Hill and co-workers reported a similar set of cationic complexes with the somewhat weaker coordinating anion $\text{Al}(\text{OC}_4\text{F}_9)_4^-$,¹⁷ we introduced a cationic β -diketiminate Mg complex with a much bulkier β -diketiminate ligand carrying *t*Bu groups in the ligand's backbone. Restriction of the Mg coordination sphere weakens the $\text{Mg}\cdots\text{B}(\text{C}_6\text{F}_5)_4^-$ contacts and this

allowed for isolation of a unique $(^{\text{tBu}}\text{BDI})\text{Mg}^+\cdots\text{Cl-Ph}$ species (**VI**).¹⁵ Herein, we present extensive experimental and theoretical investigations on Mg-halobenzene complexes and discuss bonding and bond activation.

Results and discussion

Synthesis and solid state structures of cationic $\text{Mg}\cdots$ halobenzene complexes

In order to prepare complexes of the type $[(\text{BDI})\text{Mg}^+\cdot\text{XPh}][\text{B}(\text{C}_6\text{F}_5)_4^-]$ ($\text{X} = \text{F}, \text{Cl}, \text{Br}, \text{I}$) any coordinating ethereal solvent and aromatic solvents should be strictly avoided. Reaction of $[(^{\text{Me}}\text{BDI})\text{Mg}\eta\text{Pr}]_2$ or $(^{\text{tBu}}\text{BDI})\text{Mg}\eta\text{Bu}$ with excess $[\text{Ph}_3\text{C}^+][\text{B}(\text{C}_6\text{F}_5)_4^-]$ in neat halobenzene generally led to a color change from dark orange to pale yellow, which indicated the reaction to be completed. For the cation with the smaller BDI ligand, $(^{\text{Me}}\text{BDI})\text{Mg}^+$, only the PhF complex could be isolated (**1**); Scheme 1. The $\text{B}(\text{C}_6\text{F}_5)_4^-$ anion is still weakly bound to Mg with two $\text{Mg}\cdots\text{F}$ contacts. Recrystallization from PhF led to $[(^{\text{Me}}\text{BDI})\text{Mg}^+(\text{FPh})_3][\text{B}(\text{C}_6\text{F}_5)_4^-]$ (**2**) in which cation-anion contacts are fully cleaved. All attempts to isolate PhCl, PhBr or PhI complexes failed. These halobenzenes are clearly less polar than



Scheme 1 Syntheses of $\text{Mg}\cdots\text{XPh}$ complexes ($\text{X} = \text{F}, \text{Cl}, \text{Br}, \text{I}$).



PhF and cannot compete with the $\text{Mg} \cdots \text{B}(\text{C}_6\text{F}_5)_4^-$ interaction in IV.

Switching to the cation with the bulkier ${}^t\text{BuBDI}$ ligand, however, weakens the $\text{Mg} \cdots \text{B}(\text{C}_6\text{F}_5)_4^-$ interaction and in this case we were able to isolate and structurally characterize the complete series of halobenzene complexes. The increased bulkiness of the BDI ligand led in all cases to cation–anion separation. For the most polar and least sterically encumbered PhF, a complex with two PhF ligands was isolated (3); Fig. 1 and Table 1. The geometry of the $({}^t\text{BuBDI})\text{Mg}^+ \cdot (\text{FPh})_2$ cation in 3 closely matches that of the cation in a similar complex with a different counter anion: $[(\text{CF}_3)_3\text{O}]_3\text{Al-F-Al}[\text{O}(\text{CF}_3)_3]_3^-$, see Fig. S26.† This shows that the nature of the non-coordinating anion has no effect on $\text{Mg} \cdots \text{FPh}$ coordination. For the heavier, less polar, halobenzenes only complexes with one PhX

ligand could be isolated: $[(\text{BDI})\text{Mg}^+ \cdot \text{XPh}][\text{B}(\text{C}_6\text{F}_5)_4^-]$ ($\text{X} = \text{Cl}, \text{Br}, \text{I}$) (4–5, VI); Fig. 1 and Table 1. Complexes 4 and 5 are the first examples for the coordination of PhBr and PhI to a main group metal.

While the Mg metal center in 1 and 2 is five-coordinate, that in the complexes with the bulkier ${}^t\text{BuBDI}$ ligand is four- (3) or three-coordinate (4–5, VI). The $\text{Mg} \cdots \text{F}(\text{Ph})$ contacts range from 1.946(1) to 2.0533(14) Å and, although PhF is a neutral ligand that is known to be a weak donor, the shortest contact is just as short as the $\text{Mg}^{2+} \cdots \text{F}^-$ bonds in the F-bridged dimer $[({}^{\text{Me}}\text{BDI})\text{Mg}(\mu\text{-F})(\text{THF})_2]_2$ (Mg–F av. 1.95 Å).¹⁸ Likewise, the distances $\text{Mg} \cdots \text{Cl}(\text{Ph})$ and $\text{Mg} \cdots \text{Br}(\text{Ph})$ of 2.414(1) Å and 2.551(1) Å, respectively, can be compared to the corresponding compounds containing X^- anions: $[({}^{\text{Me}}\text{BDI})\text{Mg}(\mu\text{-Cl})_2]_2$ (Mg–Cl av. 2.3884 Å) and $[({}^{\text{Me}}\text{BDI})\text{Mg}(\mu\text{-Br})_2]_2$ (Mg–Br av. 2.539 Å).¹⁹ The $\text{Mg} \cdots \text{I}(\text{Ph})$

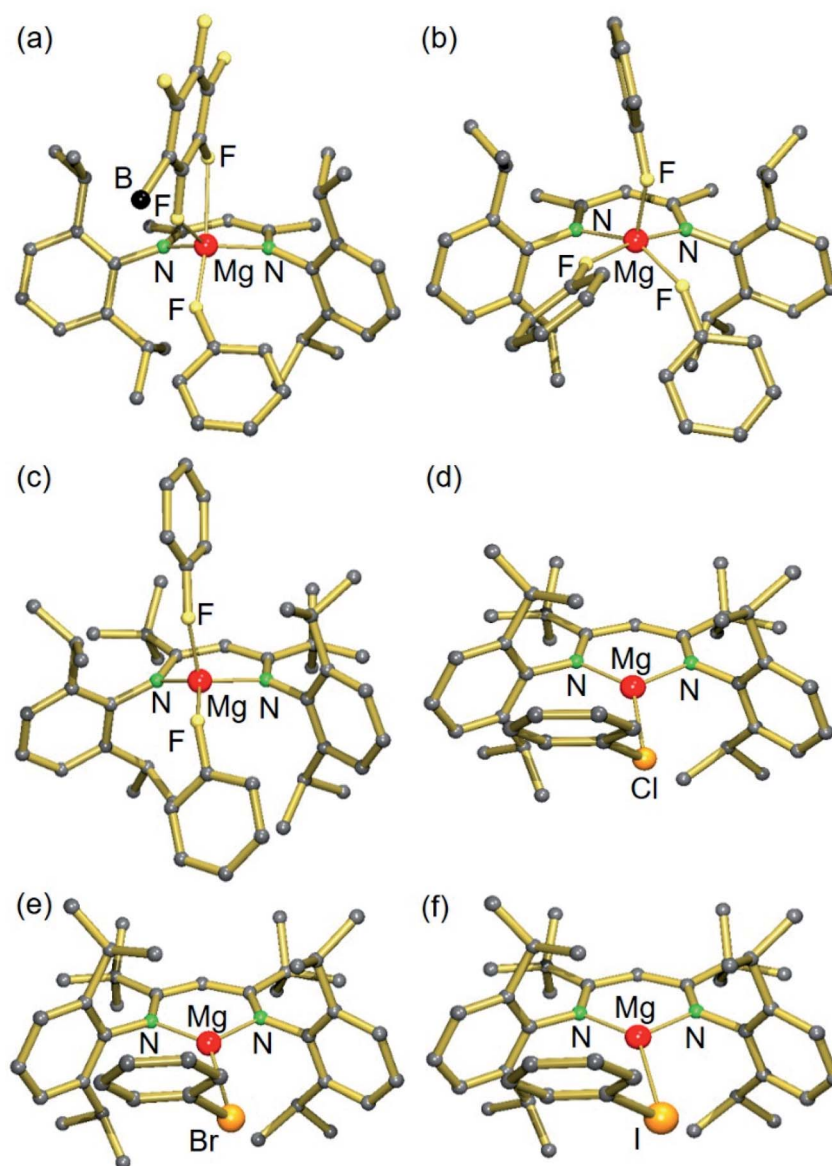


Fig. 1 Crystal structures of (a) $[({}^{\text{Me}}\text{BDI})\text{Mg}^+ \cdot \text{FPh}][\text{B}(\text{C}_6\text{F}_5)_4^-]$; borate anion partially shown (1), (b) $[({}^{\text{Me}}\text{BDI})\text{Mg}^+ \cdot (\text{FPh})_3]$ (2), (c) $[({}^t\text{BuBDI})\text{Mg}^+ \cdot (\text{FPh})_2]$ (3), (d) $[({}^t\text{BuBDI})\text{Mg}^+ \cdot \text{ClPh}]$ (VI),¹⁵ (e) $[({}^t\text{BuBDI})\text{Mg}^+ \cdot \text{BrPh}]$ (4) and (f) $[({}^t\text{BuBDI})\text{Mg}^+ \cdot \text{IPh}]$ (5). Anions and H atoms are not shown for clarity.



Table 1 Selected distances and angles in [(BDI)Mg⁺·halobenzene] complexes; numbers without standard deviations are average values. Calculated values are shown between squared brackets

| | Mg···X(Ph) [Å] | Angle Mg–X–C [°] | (Ph)C–X [Å] | Free (Ph)C–X [Å] ²¹ |
|--|-------------------|------------------|------------------|--------------------------------|
| [(^{Me} BDI)Mg ⁺ ·FPh][B(C ₆ F ₅) ₄ [−]] (1) | 2.007 (2) | 151.9 (1) | 1.417 (3) | 1.364 (15) |
| [(^{Me} BDI)Mg ⁺ ·(FPh) ₃] (2) | 2.044 | 147.1 | 1.408 | |
| [(^{tBu} BDI)Mg ⁺ ·(FPh) ₂] (3) | 1.977 [1.961] | 139.1 [131.4] | 1.417 [1.431] | |
| [(^{tBu} BDI)Mg ⁺ ·ClPh] (VI) | 2.414 (1) [2.422] | 101.4(1) [101.2] | 1.788(2) [1.792] | 1.737 (5) |
| [(^{tBu} BDI)Mg ⁺ ·BrPh] (4) | 2.551 (1) [2.563] | 97.7(1) [97.9] | 1.938(1) [1.949] | 1.899 (12) |
| [(^{tBu} BDI)Mg ⁺ ·IPh] (5) | 2.748(1) [2.759] | 95.1(1) [93.8] | 2.118(2) [2.136] | 2.095 (15) |

contact of 2.748(1) Å in 5 is considerably longer than in monomeric (^{tBu}BDI)MgI (Mg–I 2.597(1) Å)²⁰ which is due to the terminal character of the Mg–I bond in the latter.

Interaction of the highly Lewis acidic (BDI)Mg⁺ cations with Ph–X causes substantial activation of the C–X bond, indicated by significant C–X bond lengthening. Compared to other metal···F(Ph) complexes the average C–F bond of 1.417 Å in 3 is elongated by 0.053 Å compared to that in free PhF (Table 1). Also the Mg complexes of PhCl, PhBr and PhI show a significant activation of the C–X bond. The extent of bond elongation decreases along the row C–F (0.053 Å) > C–Cl (0.051 Å) > C–Br (0.039 Å) > C–I (0.023 Å), *i.e.* with decreasing C–X bond polarity. There are only very few examples in literature in which a larger extent of C–X bond activation was observed. The elongation of the C–X bonds by cationic Mg complexes is of similar magnitude as C–X bond activation by transition metals like Zr, Re, Pd, Pt, Rh or Ag (see Table S2[†] for a detailed compilation of literature Ph–X···metal complexes).

Another interesting aspect of these Mg–halobenzene complexes is the Mg···X–Ph angle which steadily decreases with halogen size: F 139.1–151.9(1)° > Cl 101.4(1)° > Br 97.7(1)° > I 95.1(1)°. While the Ph–F···Mg angle is closest to linear, the Ph–I···Mg in 5 is nearly perpendicular. This trend is also found for the DFT calculated complexes and can be explained by an increasing σ -hole along the row F < Cl < Br < I (*vide infra*).

Cationic Mg···halobenzene complexes in solution

It is difficult to identify the Mg···halobenzene complexes in solution. They are insoluble in inert solvents like pentane or hexane and use of aromatic solvents would lead to Mg··· π -arene complexes. Complexes were therefore characterized by NMR techniques in bromobenzene-*d*₅. The existence of Mg···FPh interactions has been proven with ¹⁹F NMR. Complexes [(^{Me}BDI)Mg⁺·FPh][B(C₆F₅)₄[−]] (1) and [(^{tBu}BDI)Mg⁺·(FPh)₂][B(C₆F₅)₄[−]] (3) show signals at −120.9 ppm and −116.3 ppm, respectively. These are downfield shifted in respect to uncoordinated PhF in this solvent (−111.7 ppm). Concentration dependent NMR analysis gave upon dilution a shift of the ¹⁹F NMR resonance towards the value for uncoordinated PhF. The ¹H NMR signals for the PhCl complex (VI) dissolved in bromobenzene-*d*₅ overlap with the residual solvent signals and are not informative. For the PhBr complex (4) and PhI complex (5) exchange of the halobenzene with bromobenzene-*d*₅ is expected. Indeed, the ¹H NMR spectrum of 5 in bromobenzene-*d*₅ only shows signals for free PhI.

DFT calculations

DFT calculations were performed at the B3LYP/def2TZVP level of theory with the Grimme D3 dispersion correction using Becke–Johnson damping, unless indicated otherwise (see ESI[†] for further details). Preliminary studies on the interaction of very simple model systems (HMg⁺ and HC[C(H)NH]₂Mg⁺) with Ph–X (X = F, Cl, Br, I) showed that, apart from Mg···X, also Mg··· π -arene interactions contribute to their stability (see Fig. S29 and S30[†]). Although the latter interactions were not found in the Mg···F–Ph complexes, they become more prevalent along the row Mg··· π -PhCl < Mg··· π -PhBr < Mg··· π -PhI, *i.e.* with increasing electron density in the ring π -system. This influence of the inductive effect of the X-substituent on metal··· π -PhX interactions has been previously recognized experimentally and

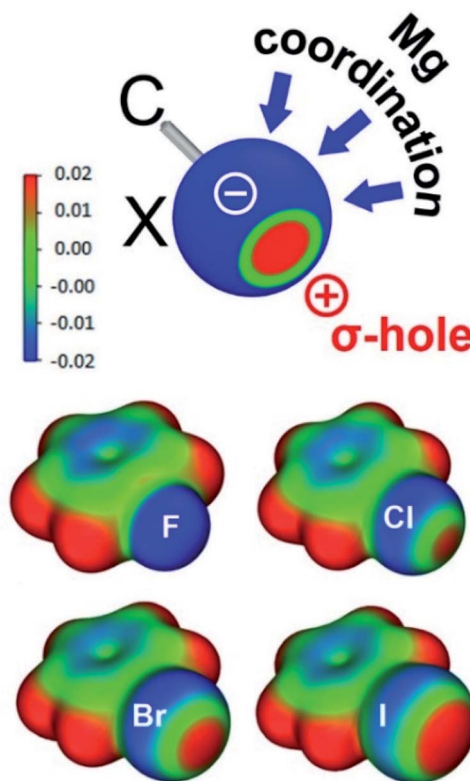


Fig. 2 (Top) Schematic representation of the σ -hole and possible coordination sites for Mg. (Bottom) Electrostatic maps of PhF, PhCl, PhBr and PhI at an isosurface of 0.003 a.u. of the total electron density.



theoretically.^{22,23} Such metal $\cdots\pi$ -PhX interactions could be avoided by calculations on the full ligand system. Calculated structures for $({}^t\text{BuBDI})\text{Mg}^+\cdots\text{XPh}$ (X = F, Cl, Br, I) closely match the crystal structures (Fig. S31† and Table 1). Not only the $\text{Mg}^+\cdots\text{XPh}$ distances but especially the $\text{Mg}^+\cdots\text{X-Ph}$ angles are in excellent agreement with those found in the crystal structures (Table 1). Given the fact that angular distortions are generally very facile, this close match between experiment and theory is surprising.

The gradually decreasing $\text{Mg}\cdots\text{X-Ph}$ angle along the row X = F > Cl > Br > I finds its origin in the concept of halogen bonding. Dating back to the first observation of $\text{I}_2\cdots\text{NH}_3$,²⁴ halogen bonding has been reviewed as early as 1968.²⁵ Halogen bonding, generally defined as a noncovalent interaction between a halogen atom and a Lewis base,²⁶ consists of different contributions including electrostatic interaction, dispersion, charge transfer and halogen atom polarization.²⁷ The angular geometries of large numbers of halogen-bonded complexes have been statistically evaluated and were rationalized by the halogen's σ -hole.²⁸ Previously dubbed as "polar flattening",²⁶ the σ -hole is a region of positive electrostatic potential on the halogen's surface (Fig. 2). Although halogen atoms are rich in free electron pairs, the surface of the atom opposite of the C-X bond is positively charged. This region increases with halogen size and is generally discussed to explain Lewis base-halogen interactions,²⁹ however, less frequently is also taken into

account when discussing Lewis acid-halogen interactions.³⁰ Decreasing $\text{Mg}\cdots\text{X-Ph}$ angles can be explained by the gradually decreasing belt of electron density on the halogen with increasing halogen size (Fig. 2). An alternative way to look at increased bending of the $\text{Mg}\cdots\text{X-Ph}$ geometries along the series X = F < Cl < Br < I is the change in the halogen atom hybridization³¹ which could be seen as the origin of the σ -hole. Two of the free electron pairs on the halogen are in p-orbitals perpendicular to the C-X axis. The third unshared pair along the C-X axis is in a hybrid orbital of s-character and some degree of p-character which is strongly reduced with increasing halogen size (for F circa 25% and for I circa 8%) explaining that the σ -hole is particularly large for the heavier halogen atoms.²⁸

In order to quantify the driving force for the bent $\text{Mg}\cdots\text{X-Ph}$ structures, the energies for linearization have been calculated for the full cation $({}^t\text{BuBDI})\text{Mg}^+\cdots\text{XPh}$ (X = F, Cl, Br, I) (Fig. S33†). While linearization of the $\text{Mg}\cdots\text{F-Ph}$ unit hardly costs any energy ($\Delta H = 2.0 \text{ kcal mol}^{-1}$), linearization free energies strongly increase for PhCl ($\Delta H = 15.1 \text{ kcal mol}^{-1}$), PhBr ($\Delta H = 18.8 \text{ kcal mol}^{-1}$) and PhI ($\Delta H = 22.1 \text{ kcal mol}^{-1}$).

Formation of the $({}^t\text{BuBDI})\text{Mg}^+\cdots\text{XPh}$ complexes is in all cases exothermic. The following ΔH values have been calculated for the gas phase interaction of $({}^t\text{BuBDI})\text{Mg}^+$ with PhX (kcal mol^{-1}): PhF -21.6 , PhCl -28.0 , PhBr -29.3 and PhI -30.6 . However, using a solvent correction (PCM = benzene) turns this order and gives lower, more realistic, values (kcal mol^{-1}): PhF -12.8 ,

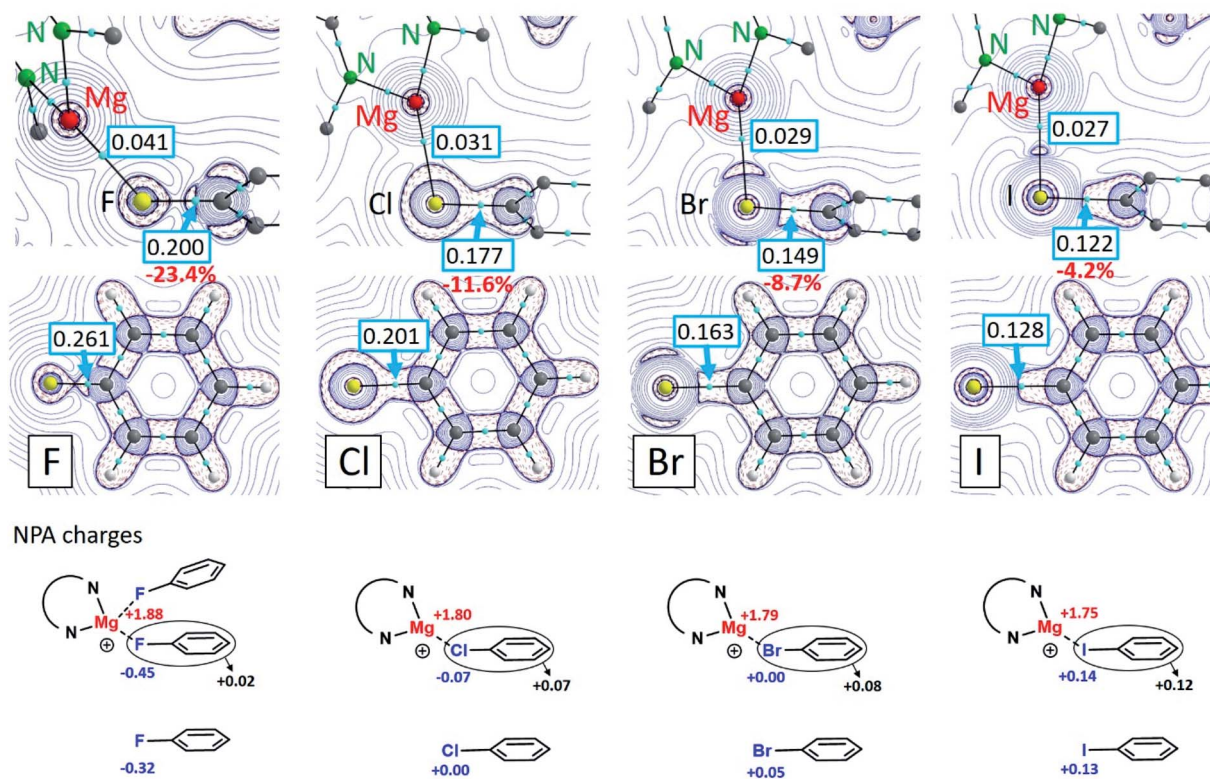


Fig. 3 Contour plots of the negative Laplacian, $-\nabla^2\rho(r)$, for the cations $({}^t\text{BuBDI})\text{Mg}^+\cdots\text{XPh}$ (X = F, Cl, Br, I) in the Mg \cdots X-C plane (top row) and for the free halobenzenes PhX in the molecule plane (middle row). Light-blue dots are bond-critical-points (BCP) and boxed numbers show the electron densities in these points. NPA charges are shown in the bottom part.



PhCl -12.3 , PhBr -11.8 and PhI -10.9 . Although the complexation energies decrease from PhF to PhI, the difference within the series is very small. The strong $\text{Mg}\cdots\text{FPh}$ interaction can be attributed to the highly polar $\text{C}^{\delta+}\text{-F}^{\delta-}$ bond which upon bonding with Mg^{2+} is even further polarized (see NPA charges in Fig. 3). The charge distribution also shows that there is hardly electron transfer from PhF to Mg. However, electron transfer from PhX to Mg increases along the row $\text{X} = \text{F} < \text{Cl} < \text{Br} < \text{I}$. This is likely related to decreasing electronegativity differences in the $\text{Mg}\cdots\text{X}$ bond along the same row which makes the $\text{Mg}\cdots\text{I}$ interaction partially covalent. The $\text{Mg}\cdots\text{IPh}$ bond is strong on account of polarization of the core electron density of the iodine atom. The latter has been visualized by an Atoms-In-Molecules (AIM) analysis.

Contour plots of the negative Laplacian, $-\nabla^2\rho(\mathbf{r})$, for the cations ($^{\text{tBu}}\text{BDI}\text{Mg}^+\cdots\text{XPh}$ ($\text{X} = \text{F}, \text{Cl}, \text{Br}, \text{I}$) are shown in Fig. 3 which for comparison also shows the electron densities for the free halobenzene ligands. In all cases, distinct $\text{Mg}\cdots\text{X}$ bond paths have been found. The electron density in the bond-critical-point (BCP) decreases gradually from $\text{X} = \text{F}$ to I . The electron density in the BCP on the C-X axis of the coordinated PhX ligands is always smaller when compared to the free PhX ligands. This is in line with the observed C-X bond activation by Mg coordination. Weakening of the C-X bond decreases along the row: C-F (-23.4%) > C-Cl (-11.6%) > C-Br (-8.7%) > C-I (-4.2%). This is in agreement with the experimentally observed elongations of the C-X bond (Table 1): C-F ($+3.7\%$) > C-Cl ($+2.9\%$) > C-Br ($+2.0\%$) > C-I ($+1.1\%$). Analysis of the Wiberg bond indices for the C-X bonds by the NBO method also shows that Mg coordination leads in all cases to C-X bond weakening (Fig. S32[†]). While the bonding interaction between Mg^{2+} and PhF can be explained by electrostatic interaction with the strongly polar $\text{C}^{\delta+}\text{-F}^{\delta-}$ bond, for the heavier halogens mainly polarization of the halogen atom is important. This is clearly visible by accumulation of electron density on the $\text{Mg}\cdots\text{X}$ axis (Fig. 3).

Conclusion

Highly Lewis acidic cationic Mg complexes with “naked” Lewis base-free Mg centers have been the key to the isolation of the full series of halobenzene complexes: $\text{Mg}\cdots\text{XPh}$ ($\text{X} = \text{F}, \text{Cl}, \text{Br}, \text{I}$). Although bonds between PhF and PhCl and main group metals have been reported, complexation of the softer ligands, PhBr and PhI, by main group metals is unprecedented.

Tuning the bulk of the β -diketiminato ligand (BDI) is crucial to the isolation of $(\text{BDI})\text{Mg}^+\cdots\text{XPh}$ complexes. For the smaller $^{\text{Me}}\text{BDI}$ ligands a competition between $(^{\text{Me}}\text{BDI})\text{Mg}^+\cdots\text{B}(\text{C}_6\text{F}_5)_4^-$ and $(^{\text{Me}}\text{BDI})\text{Mg}^+\cdots\text{XPh}$ interaction is observed and only complexes with the most polar halobenzene, PhF, could be isolated. Using the bulkier $^{\text{tBu}}\text{BDI}$ ligand shuts off cation-anion contacts and the full series $(^{\text{tBu}}\text{BDI})\text{Mg}^+\cdots\text{XPh}$ ($\text{X} = \text{F}, \text{Cl}, \text{Br}, \text{I}$) could be structurally characterized.

Complexation of PhX with Mg leads to C-X bond activation. Elongation of the C-X bond is most extreme for PhF and decreases for the heavier halogens. Most remarkable is the $\text{Mg}\cdots\text{XPh}$ coordination mode: whereas the $\text{Mg}\cdots\text{F-C}$ angle

tends to be large and linear, the $\text{Mg}\cdots\text{I-C}$ angle is close to perpendicular. The steadily decreasing $\text{Mg}\cdots\text{X-C}$ angle ($\text{X} = \text{F} > \text{Cl} > \text{Br} > \text{I}$) can be explained with the halogen σ -hole. This region of positive electrostatic potential on the halogen's surface, opposite of the C-X bond, increases with halogen size. Previous reports on the statistical examination of non-bonded interactions of arylhalides with neighboring molecules in crystal structures have shown the preferred bonding sites for electrophiles and nucleophiles.³⁰ For the heavier arylhalides (Br, I), however, only scatter plots for interactions with nucleophiles have been identified. This is due to the lack of metal interactions with arylbromide and iodide species. Here we show mapping of the σ -hole down the full halogen group with a single Lewis acidic Mg cation.

The experimentally observed structures are surprisingly well modelled by DFT calculations. Since bending angles are generally subject to variation, there is clearly a distinct preference for the bent $\text{Mg}\cdots\text{X-Ph}$ coordination modes observed for the heavier halogens. Indeed, the halogen-metal bond is highly directional²⁷ and the linearization energies strongly increase along the row $\text{X} = \text{F} < \text{Cl} < \text{Br} < \text{I}$.

Although weaker than a classical O-H \cdots O hydrogen bond, the complexation enthalpies between $11\text{--}13\text{ kcal mol}^{-1}$ (ΔH) for $(^{\text{tBu}}\text{BDI})\text{Mg}^+\cdots\text{XPh}$ formation are still considerable and relevant for physical interaction. Bonding between Mg and PhF is mainly dictated by the very polar $\text{C}^{\delta+}\text{-F}^{\delta-}$ bond. Bonding between Mg and the heavier PhX ligands relies increasingly on halogen polarization. Evidence for electrostatic $\text{Mg}\cdots\text{X-Ph}$ interactions is mainly based on solid state structures. In solution there is competition with $\text{Mg}\cdots\text{solvent}$ bonding and dissociation equilibria lie largely on the side of free uncoordinated PhX. Despite their weakness, we embrace the idea that metal $\cdots\text{X-Ph}$ interactions can play an important role in halogen exchange reactions. Being stronger than typical van der Waals interactions, the preferred approach of early main group metal reagents and halobenzenes is defined by metal $\cdots\text{X-Ph}$ bonding. Since the interaction between a positively charged s-block metal and organic halides activates the C-X bond, they could be especially important in the C-X bond breaking process. Although the here presented $\text{Mg}\cdots\text{XPh}$ complexes are in solution short-living intermediates, their challenging isolation and structural characterization contribute to mechanistic insights in the emerging field of s-block metal catalysis.³²

Experimental section

General considerations

All experiments were conducted under an inert nitrogen atmosphere using standard Schlenk and glovebox techniques (MBraun, Labmaster SP). All solvents were degassed with nitrogen, dried over activated aluminium oxide (Solvent Purification System: Pure Solv 400-4-MD, Innovative Technology) and stored over 3 Å molecular sieves unless noted otherwise. Fluorobenzene, chlorobenzene, bromobenzene and iodobenzene were dried over calcium hydride, distilled under N_2 atmosphere and stored over molecular sieves 3 Å. C_6D_6 and $\text{C}_6\text{D}_5\text{Br}$ (99.6% D, Sigma Aldrich) were dried over 3 Å molecular sieves. $[\text{CPh}_3]^+$



$[B(C_6F_5)_4]^-$ (Boulder Scientific) was used as received. Complexes $[(^{Me}BDI)MgnPr]_2$,¹¹ $[(^{tBu}BDI)MgnBu]$,³³ and $[(^{tBu}BDI)Mg^+ \cdot (PhCl)] [B(C_6F_5)_4]^-$ ¹⁵ were synthesized according to literature procedures. NMR spectra were recorded with a Bruker Avance III HD 400 MHz or a Bruker Avance III HD 600 MHz spectrometer. The spectra were referenced to the respective residual signals of the deuterated solvents. Elemental analysis was performed with an Euro EA 3000 (Euro Vector) analyzer. All crystal structures have been measured on a SuperNova (Agilent) diffractometer with dual Cu and Mo microfocus sources and an Atlas S2 detector. Crystallographic data have been deposited with the Cambridge Crystallographic Data Centre as supplementary publication no. 2042836 (1), 2042837 (2), 2042838 (3), 2042839 (4), 2042840 (5) and 2042841 for $[(^{tBu}BDI)Mg^+ \cdot (FPh)_2][\{(CF_3)_3O\}_3Al-F-Al\{O(CF_3)_3\}_3]^-$.

Synthesis of $[(^{Me}BDI)Mg^+ \cdot (PhF)][B(C_6F_5)_4]^-$ (1) and $[(^{Me}BDI)Mg^+ \cdot (PhF)_3][B(C_6F_5)_4]^-$ (2). $[(^{Me}BDI)MgnPr]_2$ (0.0858 g, 0.0884 mmol) and $[Ph_3C^+][B(C_6F_5)_4]^-$ (0.1513 g, 0.1640 mmol) were dissolved in fluorobenzene (2 mL). The brownish solution was stirred until it became almost colorless (1 min) and was allowed to stand for 18 h. The solution was filtered and all volatiles were removed under reduced pressure resulting in a slightly yellow foam. The material was extracted with hexane (4 × 1 mL) and directly crystallized from the solution. The colorless crystals were dried *in vacuo* to yield 0.0412 g (21%) of the desired product. Complex 2 was crystallized by slow solvent evaporation from a $[(^{Me}BDI)Mg^+ \cdot (PhF)][B(C_6F_5)_4]^-$ solution in neat PhF. ¹H NMR (C_6D_5Br , 600 MHz, 298 K) δ 7.19 (m, 2H, aryl-*H*), 7.06 (m, 6H, aryl-*H*, PhF), 6.95 (m, 1H, PhF), 6.68 (m, 2H, PhF), 4.99 (s, 1H, CCHC), 2.78 (hept, ³*J*_{HH} = 6.9 Hz, 4H, CH(CH₃)₂), 1.60 (s, 6H, CCH₃), 1.05 (d, ³*J*_{HH} = 6.9 Hz, 12H, CH(CH₃)₂), 0.89 (d, ³*J*_{HH} = 6.9 Hz, 12H, CH(CH₃)₂). ¹³C NMR (C_6D_5Br , 151 MHz, 298 K) δ 173.5 (s, CCHC), 164.2 (d, ¹*J*_{CF} = 233 Hz, PhF), 149.1 (br d, ¹*J*_{CF} = 238 Hz, B(C₆F₅)₄⁻), 142.4 (s, aryl-*C*), 141.8 (s, aryl-*C*), 138.1 (br t, ¹*J*_{CF} = 238 Hz, B(C₆F₅)₄⁻), 131.0 (d, ³*J*_{CF} = 9.8 Hz, PhF), 127.7 (s, aryl-*C*), 126.4 (s, aryl-*C*), 125.2 (s, PhF), 114.8 (d, ²*J*_{CF} = 21.4 Hz, PhF), 96.8 (s, CCHC), 29.1 (s, CH(CH₃)₂), 24.6 (s, CH(CH₃)₂), 24.5 (s, CCH₃), 24.4 (s, CH(CH₃)₂). ¹⁹F NMR (C_6D_5Br , 565 MHz, 298 K) δ -165.1 (t, ³*J*_{FF} = 21 Hz, 8F, meta aryl-F), -160.6 (t, ³*J*_{FF} = 22 Hz, 4F, para aryl-F), -131.3 (d, ³*J*_{FF} = 19 Hz, 8F, ortho aryl-F), -120.9 (s, 1F, PhF). ¹¹B NMR (C_6D_5Br , 193 MHz, 298 K) δ -15.7 (s, B(C₆F₅)₄⁻). Elemental analysis found: C, 58.04; H, 3.96; N, 2.25. Calc. for C₅₉H₄₆BF₂₁MgN₂: C, 58.22; H, 3.81; N, 2.30%.

Synthesis of $[(^{tBu}BDI)Mg^+ \cdot (PhF)_2][B(C_6F_5)_4]^-$ (3). $(^{tBu}BDI)MgnBu$ (0.121 g, 0.207 mmol, 1.1 eq.) and $[Ph_3C^+][B(C_6F_5)_4]^-$ (0.173 g, 0.188 mmol, 1 eq.) were dissolved in fluorobenzene (3 mL). The red-brown solution was stirred for 2 h and a change to a clear yellow solution was observed. Subsequently all volatiles were removed under reduced pressure. The resulting foam was washed with hexane (8 × 3 mL) giving the product as a white powder. $[(^{tBu}BDI)Mg^+ \cdot (PhF)_2][B(C_6F_5)_4]^-$ (85 mg) was obtained in 30% yield. Crystals suitable for X-ray diffraction were grown from a concentrated fluorobenzene solution layered with hexane. ¹H NMR (C_6D_5Br , 600 MHz, 298 K): δ 7.09 (m, 4H, aryl-*H*), 6.95 (m, 5H, aryl-*H*), 6.83 (m, 2H, aryl-*H*), 5.54 (s, 1H, CCHC), 2.89 (sept, ³*J*_{HH} = 6.8 Hz, 4H, CH(CH₃)₂), 1.16 (d, ³*J*_{HH} = 6.8 Hz,

12H, CH(CH₃)₂), 1.04 (s, 18H, C(CH₃)₃), 0.84 (d, ³*J*_{HH} = 6.8 Hz, 12H, CH(CH₃)₂) ppm. ¹³C NMR (C_6D_5Br , 151 MHz, 298 K): δ 182.0 (s, CCHC), 163.7 (d, ¹*J*_{CF} = 240 Hz, PhF), 148.9 (d, ¹*J*_{CF} = 244 Hz, B(C₆F₅)₄⁻), 142.9 (s, aryl-*C*), 141.6 (s, aryl-*C*), 138.7 (d, ¹*J*_{CF} = 244 Hz, B(C₆F₅)₄⁻), 137.2 (d, ¹*J*_{CF} = 244 Hz, B(C₆F₅)₄⁻), 130.8 (s, aryl-*C*), 130.7 (s, aryl-*C*), 127.2 (s, aryl-*C*), 125.5 (s, aryl-*C*), 125.1 (s, aryl-*C*), 115.5 (s, aryl-*C*), 115.4 (s, aryl-*C*), 97.5 (s, CCHC), 45.3 (s, C(CH₃)₃), 32.8 (s, C(CH₃)₃), 28.7 (s, CH(CH₃)₂), 26.2 (s, CH(CH₃)₂), 23.9 (s, CH(CH₃)₂) ppm. ¹⁹F NMR (C_6D_5Br , 565 MHz, 298 K): δ -165.7 (br s, 8F, meta aryl-F), -161.0 (br s, 4F, para aryl-F), -130.5 (br s, 8F, ortho aryl-F), -116.3 (br s, 1F, PhF) ppm. ¹¹B NMR (C_6D_5Br , 193 MHz, 298 K): δ -15.6 (s, 1B, B(C₆F₅)₄⁻) ppm. Elemental analysis found: C, 60.79; H, 4.60; N, 2.04. Calc. for C₇₁H₆₃BF₂₂MgN₂: C, 61.03; H, 4.54; N, 2.00%.

Synthesis of $[(^{tBu}BDI)Mg^+ \cdot (PhBr)][B(C_6F_5)_4]^-$ (4). $(^{tBu}BDI)MgnBu$ (63.7 mg, 0.109 mmol, 1.1 eq.) was dissolved in bromobenzene (2 mL). Addition of $[Ph_3C^+][B(C_6F_5)_4]^-$ (91.6 mg, 0.0993 mmol, 1 eq.) gave a red-brown solution which immediately changed color to pale yellow upon shaking for 1 min. After layering the reaction mixture with hexane (0.4 mL) colorless crystals were obtained after 3 days. The crystals were washed with hexane (3 × 1 mL) giving the product $[(^{tBu}BDI)Mg^+ \cdot (PhBr)][B(C_6F_5)_4]^-$ in 58% yield (78 mg). Crystals suitable for X-ray diffraction were grown from the reaction mixture. ¹H NMR (C_6D_5Br , 600 MHz, 298 K): δ 7.09 (t, ³*J*_{HH} = 7.7 Hz, 2H, aryl-*H*), 6.93 (m, 4H, aryl-*H*), 5.51 (s, 1H, CCHC), 2.89 (sept, ³*J*_{HH} = 6.8 Hz, 4H, CH(CH₃)₂), 1.17 (d, ³*J*_{HH} = 6.8 Hz, 12H, CH(CH₃)₂), 1.03 (s, 18H, C(CH₃)₃), 0.83 (d, ³*J*_{HH} = 6.8 Hz, 12H, CH(CH₃)₂) ppm. ¹³C NMR (C_6D_5Br , 151 MHz, 298 K): δ 182.0 (s, CCHC), 149.1 (d, ¹*J*_{CF} = 244 Hz, B(C₆F₅)₄⁻), 143.0 (s, aryl-*C*), 141.6 (s, aryl-*C*), 138.8 (d, ¹*J*_{CF} = 244 Hz, B(C₆F₅)₄⁻), 137.4 (d, ¹*J*_{CF} = 244 Hz, B(C₆F₅)₄⁻), 133.1 (s, PhBr), 132.0 (s, PhBr), 130.5 (s, PhBr), 127.2 (s, PhBr), 125.1 (s, aryl-*C*), 121.4 (s, aryl-*C*), 97.7 (s, CCHC), 45.3 (s, C(CH₃)₃), 32.8 (s, C(CH₃)₃), 28.7 (s, CH(CH₃)₂), 26.2 (s, CH(CH₃)₂), 23.9 (s, CH(CH₃)₂) ppm. ¹⁹F NMR (C_6D_5Br , 565 MHz, 298 K): δ -165.8 (br s, 8F, meta aryl-F), -160.8 (br s, 4F, para aryl-F), -130.4 (br s, 8F, ortho aryl-F) ppm. ¹¹B NMR (C_6D_5Br , 193 MHz, 298 K): δ -15.6 (s, 1B, B(C₆F₅)₄⁻) ppm. Elemental analysis found: C, 58.37; H, 4.35; N, 2.09. Calc. for C₆₅H₅₈BBF₂₀MgN₂: C, 57.31; H, 4.29; N, 2.06%.

Synthesis of $[(^{tBu}BDI)Mg^+ \cdot (PhI)][B(C_6F_5)_4]^-$ (5). $(^{tBu}BDI)MgnBu$ (46.1 mg, 0.079 mmol, 1.05 eq.) was dissolved in iodobenzene (0.6 mL). Addition of $[Ph_3C^+][B(C_6F_5)_4]^-$ (69.4 mg, 0.075 mmol, 1 eq.) gave a red-brown solution that changed color to pale yellow upon shaking for 1 min. The reaction mixture was allowed to stand at room temperature and colorless crystals were obtained overnight. The crystals were washed with hexane (4 × 1 mL) giving the product $[(^{tBu}BDI)Mg^+ \cdot (PhI)][B(C_6F_5)_4]^-$ in 80% yield (85 mg). Crystals suitable for X-ray diffraction were grown from the reaction mixture. ¹H NMR (C_6D_5Br , 600 MHz, 298 K): δ 7.50 (m, 2H, PhI), 7.09 (t, ³*J*_{HH} = 7.7 Hz, 2H, aryl-*H*), 7.06 (m, 1H, PhI), 6.94 (d, ³*J*_{HH} = 7.7 Hz, 4H, aryl-*H*), 6.81 (m, 2H, PhI), 5.51 (s, 1H, CCHC), 2.90 (sept, ³*J*_{HH} = 6.8 Hz, 4H, CH(CH₃)₂), 1.17 (d, ³*J*_{HH} = 6.8 Hz, 12H, CH(CH₃)₂), 1.03 (s, 18H, C(CH₃)₃), 0.83 (d, ³*J*_{HH} = 6.8 Hz, 12H, CH(CH₃)₂) ppm. ¹³C NMR (C_6D_5Br , 151 MHz, 298 K): δ 182.0 (s, CCHC), 149.0 (d, ¹*J*_{CF} = 244 Hz, B(C₆F₅)₄⁻), 143.0 (s, aryl-*C*), 141.6 (s, aryl-*C*), 138.7 (d,



$^1J_{CF} = 244$ Hz, $B(C_6F_5)_4^-$, 137.2 (d, $^1J_{CF} = 244$ Hz, $B(C_6F_5)_4^-$), 133.0 (s, PhI), 130.7 (s, PhI), 127.9 (s, PhI), 126.8 (s, aryl-C), 125.1 (s, aryl-C), 97.7 (s, CCHC), 95.3 (s, *ipso* C-PhI), 45.3 (s, $C(CH_3)_3$), 32.8 (s, $C(CH_3)_3$), 28.7 (s, $CH(CH_3)_2$), 26.2 (s, $CH(CH_3)_2$), 23.9 (s, $CH(CH_3)_2$) ppm. ^{19}F NMR (C_6D_5Br , 565 MHz, 298 K): δ -165.8 (br s, 8F, *meta* aryl-F), -160.8 (br s, 4F, *para* aryl-F), -130.3 (br s, 8F, *ortho* aryl-F) ppm. ^{11}B NMR (C_6D_5Br , 193 MHz, 298 K): δ -15.6 (s, 1B, $B(C_6F_5)_4^-$) ppm. Elemental analysis found: C, 54.91; H, 4.31; N, 1.75. Calc. for $C_{65}H_{58}BF_{20}IMgN_2$: C, 55.40; H, 4.15; N, 1.99%.

Synthesis of $[(^{tBu}BDI)Mg^+ \cdot (PhF)_2][\{(CF_3)_3O\}_3Al-F-Al\{O(CF_3)_3\}_3]^-$. ($^{tBu}BDI)Mg^+nBu$ (37.5 mg, 64.4 μ mol, 1.1 eq.) was dissolved in fluorobenzene (3 mL). A solution of $[Ph_3C^+][\{(CF_3)_3O\}_3Al-F-Al\{O(CF_3)_3\}_3]^-$ (101.0 mg, 58.5 μ mol, 1 eq.) in PhF (0.5 mL) was added and stirred for 5 min. The pale yellow solution was layered with hexane (0.5 mL) and after 2 d colorless crystals were obtained. The crystals were collected and washed with hexane (3×1 mL) giving the product as a white crystalline solid. $[(^{tBu}BDI)Mg^+ \cdot (PhF)_2][\{(CF_3)_3O\}_3Al-F-Al\{O(CF_3)_3\}_3]^-$ (75 mg) was obtained in 58% yield. Crystals suitable for X-ray diffraction were grown from the reaction mixture. 1H NMR (C_6D_5F , 600 MHz, 298 K): δ 7.21 (t, 2H, $^3J_{HH} = 7.7$ Hz, aryl-H), 7.08 (d, 4H, $^3J_{HH} = 7.7$ Hz, aryl-H), 5.88 (s, 1H, CCHC), 3.06 (sept, $^3J_{HH} = 6.8$ Hz, 4H, $CH(CH_3)_2$), 1.21 (d, $^3J_{HH} = 6.8$ Hz, 12H, $CH(CH_3)_2$), 1.18 (s, 18H, $C(CH_3)_3$), 0.93 (d, $^3J_{HH} = 6.8$ Hz, 12H, $CH(CH_3)_2$) ppm. ^{13}C NMR (C_6D_5F , 151 MHz, 298 K): δ 182.0 (s, CCHC), 143.4 (s, aryl-C), 142.0 (s, aryl-C), 126.7 (s, *para* aryl-C), 124.9 (s, *meta* aryl-C), 122.5 (s, [Al-F-Al]), 120.6 (s, [Al-F-Al]), 96.9 (s, CCHC), 45.1 (s, $C(CH_3)_3$), 32.4 (s, $C(CH_3)_3$), 28.4 (s, $CH(CH_3)_2$), 25.4 (s, $CH(CH_3)_2$), 23.3 (s, $CH(CH_3)_2$) ppm. ^{19}F NMR (C_6D_5F , 565 MHz, 298 K): δ -74.7 (s, 54F, [Al-F-Al]), -118.2 (s, 1F, PhF), -183.8 (s, 1F, [Al-F-Al]) ppm. ^{27}Al NMR (C_6D_5F , 565 MHz, 298 K): no signal from δ -200 to +200 ppm. Elemental analysis found: C, 38.12; H, 2.52; N, 1.43. Calc. for $C_{71}H_{63}Al_2F_{57}MgN_2O_6$: C, 38.74; H, 2.88; N, 1.27%.

Author contributions

A. Friedrich: conceptualization, investigation, validation, formal analysis, writing – original draft, visualization. J. Pahl: conceptualization, investigation, validation, formal analysis. J. Eysel: formal analysis, validation. J. Langer: formal analysis, validation. N. van Eikema Hommes: formal analysis, validation, supervision. A. Görling: validation, supervision. Sjoerd Harder: conceptualization, writing – original draft – review and editing, visualization, validation, supervision, project administration.

Conflicts of interest

The authors declare no competing financial interest.

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References

- R. J. Kulawiec and R. H. Crabtree, *Coord. Chem. Rev.*, 1990, **99**, 89–115.
- H. Plenio, *Chem. Rev.*, 1997, **97**, 3363–3384.
- V. V. Zhdankin, *Hypervalent Iodine Chemistry*, John Wiley & Sons Ltd, Chichester, UK, 2013.
- T. G. Richmond, *Topics in Organometallic Chemistry – Activation of Unreactive Bonds and Organic Synthesis*, Springer-Verlag, Berlin Heidelberg, 1999.
- F. J. Fernandez-Alvarez, M. Iglesias, L. A. Oro and V. Passarelli, *Comprehensive Inorganic Chemistry II – Bond Activation and Catalysis*, Elsevier, Amsterdam, 2013.
- U. Rosenthal, V. V. Burlakov, P. Arndt, A. Spannenberg, U. Jaeger-Fiedler, M. Klahn and M. Hapke, *Activating Unreactive Substrates*, Wiley-VCH, Weinheim, 2009.
- P. G. Williard and Q. Y. Liu, *J. Org. Chem.*, 1994, **59**, 1596–1597.
- A. V. Korolev, F. Delpech, S. Dagorne, I. A. Guzei and R. F. Jordan, *Organometallics*, 2001, **20**, 3367–3369.
- (a) X.-Y. Liu, K. Venkatesan, H. W. Schmalte and H. Berke, *Organometallics*, 2004, **23**, 3153–3163; (b) A. M. Chapman, M. F. Haddow and D. F. Wass, *J. Am. Chem. Soc.*, 2011, **133**, 18463–18478; (c) O. J. Metters, S. J. K. Forrest, H. A. Sparkes, I. Manners and D. F. Wass, *J. Am. Chem. Soc.*, 2016, **138**, 1994–2003; (d) P. Ren, S. D. Pike, I. Pernik, A. S. Weller and M. C. Willis, *Organometallics*, 2015, **34**, 711–723; (e) A. V. Korolev, E. Ihara, I. A. Guzei, V. G. Young and R. F. Jordan, *J. Am. Chem. Soc.*, 2001, **123**, 8291–8309; (f) S. Balasubramaniam, S. Kumar, A. P. Andrews, B. Varghese, E. D. Jemmis and A. Venugopal, *Eur. J. Inorg. Chem.*, 2019, 3265–3269.
- (a) M. D. Butts, B. L. Scott and G. J. Kubas, *J. Am. Chem. Soc.*, 1996, **118**, 11831–11843; (b) L. P. Press, B. J. McCulloch, W. Gu, C.-H. Chen, B. M. Foxman and O. V. Ozerov, *Chem. Commun.*, 2015, **51**, 14034–14037; (c) J. Powell, M. Horvath and A. Lough, *J. Organomet. Chem.*, 1993, **456**, 27–28; (d) J. Powell, M. J. Horvath, A. Lough, A. Phillips and J. Brunet, *J. Chem. Soc., Dalton Trans.*, 1998, 637–646.
- J. Pahl, S. Brand, H. Elsen and S. Harder, *Chem. Commun.*, 2018, **54**, 8685–8688.
- J. Pahl, H. Elsen, A. Friedrich and S. Harder, *Chem. Commun.*, 2018, **54**, 7846–7849.
- J. Pahl, A. Friedrich, H. Elsen and S. Harder, *Organometallics*, 2018, **37**, 2901–2909.
- S. Brand, H. Elsen, J. Langer, S. Grams and S. Harder, *Angew. Chem., Int. Ed.*, 2019, **58**, 15496–15503.
- A. Friedrich, J. Pahl, H. Elsen and S. Harder, *Dalton Trans.*, 2019, **48**, 5560–5568.
- J. Pahl, T. E. Stennett, M. Volland, D. M. Guldi and S. Harder, *Chem.–Eur. J.*, 2019, **25**, 2025–2034.
- L. Garcia, M. D. Anker, M. F. Mahon, L. Maron and M. S. Hill, *Dalton Trans.*, 2018, **47**, 12684–12693.
- (a) H. Hao, H. W. Roesky, Y. Ding, C. Cui, M. Schormann, H. G. Schmidt, M. Noltemeyer and B. Žemva, *J. Fluorine Chem.*, 2002, **115**, 143–147; (b) C. Bakewell, A. J. P. White



- and M. R. Crimmin, *J. Am. Chem. Soc.*, 2016, **138**, 12763–12766.
- 19 A. P. Dove, V. C. Gibson, P. Hormnirun, E. L. Marshall, J. A. Segal, A. J. P. White and D. J. Williams, *Dalton Trans.*, 2003, 3088–3097.
- 20 S. J. Bonyhady, C. Jones, S. Nembenna, A. Stasch, A. J. Edwards and G. J. McIntyre, *Chem.–Eur. J.*, 2010, **16**, 938–955.
- 21 Crystal structure of PhF: V. R. Thalladi, H. C. Weiss, D. Bläser, R. Boese, A. Nangia and G. R. Desiraju, *J. Am. Chem. Soc.*, 1998, **120**, 8702–8710; crystal structure of PhCl: N. P. Penionzhkevich, N. I. Sadova and L. V. Vilkov, *Zh. Strukt. Khim.*, 1979, **20**, 527–529; crystal structures of PhBr and PhI: F. H. Allen, O. Kennard, D. G. Watson, L. Brammer, A. G. Orpen and R. Taylor, *J. Chem. Soc., Perkin Trans. 2*, 1987, 1–19.
- 22 J.-F. Gal, P.-C. Maria, M. Decouzon, O. Mó and M. Yáñez, *Int. J. Mass Spectrom.*, 2002, **219**, 445–456.
- 23 N. Hallowita, E. Udonkang, C. Ruan, C. E. Frieler and M. T. Rodgers, *Int. J. Mass Spectrom.*, 2009, **283**, 35–47.
- 24 F. Guthrie, *J. Chem. Soc.*, 1863, **16**, 239–244.
- 25 H. A. Bent, *Chem. Rev.*, 1968, **68**, 587–648.
- 26 R. Sedlak, M. H. Kolár and P. Hobza, *J. Chem. Theory Comput.*, 2015, **11**, 4727–4732.
- 27 P. Politzer, J. S. Murray and T. Clark, *Phys. Chem. Chem. Phys.*, 2010, **12**, 7748–7757.
- 28 T. Clark, M. Henneman, J. S. Murray and P. Politzer, *J. Mol. Model.*, 2007, **13**, 291–296.
- 29 (a) G. Cavallo, P. Metrangolo, R. Milani, T. Pilati, A. Priimagi, G. Resnati and G. Terraneo, *Chem. Rev.*, 2016, **116**, 2478–2601; (b) J. Y. C. Lim and P. D. Beer, *Chem*, 2018, **4**, 731–783.
- 30 N. Ramasubbu, R. Parthasarathy and P. Murray-Rust, *J. Am. Chem. Soc.*, 1986, **108**, 4308–4314.
- 31 W. Kutzelnigg, *Angew. Chem., Int. Ed. Engl.*, 1984, **23**, 272–295.
- 32 (a) S. Harder, *Chem. Rev.*, 2010, **110**, 3852–3876; (b) M. S. Hill, D. J. Liptrot and C. Weetman, *Chem. Soc. Rev.*, 2016, **45**, 972–988; (c) *Early Main Group Metal Catalysis – Concepts and Reactions*, ed. S. Harder, Wiley-VCH, Weinheim, Germany, 2020.
- 33 V. Balasanthiran, M. H. Chisholm, K. Choojun, C. B. Durr and P. M. Wambua, *Polyhedron*, 2016, **103**, 235–240.

