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Interplay of donor–acceptor interactions in stabilizing boron nitride compounds: insights from theory†

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The stability of a variety of linear and cyclic $(\text{BN})_n$ ($n = 1-3$) adducts with N-heterocyclic carbene (ImMe_2 ; $\text{ImMe}_2 = [(\text{HCNMe})_2\text{C}:]$), N-heterocyclic olefin (ImMe_2CH_2) and Wittig (Me_3PCH_2) donors has been examined using M05-2X/cc-pVTZ computations. The strength and nature of the bonds have been investigated using natural bond orbital (NBO) and atoms-in-molecules (AIM) analyses. Complementary energy decomposition analysis (EDA-NOCV) has been carried out based on BP86/TZ2P computations. In agreement with NBO and AIM analyses, the orbital interaction energy obtained from EDA contributes at least 50% to the total attractive interactions for the carbon–boron bonds indicating their largely covalent nature. The feasibility of isolating monomeric $(\text{BN})_n$ units using a donor/acceptor protocol was also investigated in a series of adducts of the general form: $\text{LB}\cdot(\text{BN})_n\cdot\text{BH}_3$ and $\text{LB}\cdot(\text{BN})_n\cdot\text{W}(\text{CO})_5$ ($n = 1-3$; LB = Lewis bases). Moreover, EDA-NOCV analysis of $\text{ImMe}_2\cdot\text{BN}\cdot\text{W}(\text{CO})_5$ and $\text{ImMe}_2\cdot\text{B}_3\text{N}_3\cdot\text{W}(\text{CO})_5$ shows that the carbene–boron bonds are stronger in the presence of $\text{W}(\text{CO})_5$ as a Lewis acid mainly because of a dramatic decrease in the amount of Pauli repulsion rather than an increase in the electrostatic/orbital attraction terms.

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1 Introduction

Boron nitride materials $(\text{BN})_n$ are isoelectronic with various carbon allotropes (e.g., diamond, graphene, nanotubes),^{1–3} however, they possess vastly different properties due to the presence of polarized B–N linkages. Two specific properties, wide electronic band gaps (5.9 eV in hexagonal BN) and chemical inertness,⁴ make boron nitride of significant interest for the electronics industry, with the construction of devices based on BN/graphene heterostructures being a promising recent development.^{5,6} One hindrance to the widespread examination and application of BN materials is the harsh conditions required for their syntheses, i.e., heating above 900 °C and/or the use of plasma conditions.^{1–3} With these challenges in mind, we have embarked on a program wherein complexes of the

general form $[\text{LB}\cdot\text{BN}]_n$ ($n \geq 1$; LB = Lewis base) might be formed with suitable carbon-based donors. Upon heating in solution, the target $[\text{LB}\cdot\text{BN}]_n$ complexes could afford bulk boron nitride and free Lewis base. To provide a solid fundamental basis for future experimental explorations, we present quantum mechanical computations on the Lewis base-stabilized linear and cyclic boron nitride species $(\text{BN})_n$ ($n = 1-3$) including the donor–acceptor adducts $\text{LB}\cdot(\text{BN})_n\cdot\text{LA}$ ($n = 1-3$, LA = Lewis acid). Recent examples of stabilizing main group element units (e.g., Si_2) with the aid of strong carbon-based donors are numerous in the literature.^{7–15} Moreover, donor–acceptor stabilization has been used to great success to isolate heavier group 13–15 element species,^{16–21} while related computational studies have been reported.^{22,23} More specifically, DFT predicted a significant thermodynamical stabilization of group 13–15 cubane systems (e.g., B_4N_4) upon addition of NH_3 and BH_3 as donor and acceptor molecules, respectively.²²

In this work we present our analyses of the bonding within Lewis base-substituted boron nitride compounds in the presence and/or absence of Lewis acid. Specifically we examined the binding of the carbon-based donors, ImMe_2 ($\text{ImMe}_2 = [(\text{HCNMe})_2\text{C}:]$), ImMe_2CH_2 and Me_3PCH_2 to $(\text{BN})_n$ units, given our use of their sterically hindered analogues to bind/stabilize inorganic methylene and ethylene units (EH_2 and $\text{H}_2\text{EE}'\text{H}_2$; E and E' = Si, Ge and/or Sn).^{14,24–31} We also provide computations on $\text{LB}\cdot(\text{BN})_n\cdot\text{BH}_3$ and $\text{LB}\cdot(\text{BN})_n\cdot\text{W}(\text{CO})_5$ adducts ($n = 1-3$) featuring coordinated $(\text{BN})_n$ units

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† Electronic supplementary information (ESI) available: NICS values for all the B_2N_2 and B_3N_3 species (Table S1), NBO results for all the species (Tables S2–S9), M05-2X/cc-pVTZ computed gas phase XYZ coordinates for all the structures (Table S10), BP86/TZ2P computed gas phase XYZ coordinates for all the mono-substituted adducts (Table S11), M05-2X/cc-pVTZ computed gas phase complexation energies for the Lewis acid bound adducts (Table S12), M05-2X/cc-pVTZ optimized structures of the isolated species in the gas phase (Fig. S1), AIM results for all the species (Fig. S2), and frontier molecular orbitals of all the studied systems (Fig. S3). See DOI: 10.1039/c5cp01993a



and show that this overall donor–acceptor approach is a viable means of intercepting a complex of molecular boron nitride. Finally, based on detailed EDA-NOCV computations, we will comment on the strength and nature of both the carbene–boron and nitrogen–tungsten donor acceptor bonds in the ImMe_2 substituted $\text{BN}\cdot\text{W}(\text{CO})_5$ and $\text{B}_3\text{N}_3\cdot\text{W}(\text{CO})_5$ adducts.

2 Computational methods

Geometry optimizations were performed using density functional theory (DFT) with the M05-2X³² functional. The computations employed the following basis sets: cc-pVTZ^{33,34} for all period 1, 2 and 3 atoms and cc-pVTZ-PP,^{35,36} combined with the corresponding small core (60 electrons) effective core potential (ECP) for tungsten (W). The basis set and ECP for tungsten were obtained from the basis set exchange.^{37,38} For convenience, these computations are simply labeled as M05-2X/cc-pVTZ throughout the text. Triplet states for BN, linear BNBN, and cyclic B_2N_2 were computed using an UHF reference. For geometry optimizations, “Tight” convergence criteria were applied: maximum force = 1.5×10^{-5} a.u., RMS force = 1.0×10^{-5} a.u., maximum displacement = 6.0×10^{-5} , and RMS displacement = 4.0×10^{-5} . The grid used for numerical integration in DFT was set to “Ultrafine” with a pruned grid of 99 radial shells and 590 angular points per shell. Harmonic vibrational frequencies were computed analytically at the same level of theory in order to characterize the stationary points as minima, representing equilibrium structures on the potential energy surfaces.

Energy decomposition analyses (EDA) were performed for all the mono-substituted complexes using the GGA BP86 density functional^{39,40} and the TZ2P basis set;⁴¹ relativistic effects were considered for the tungsten atom using the ZORA approximation. As originally developed by Morokuma,⁴² Ziegler and Rauk,⁴³ EDA analysis can provide valuable insight into the nature and strength of a bond. It decomposes the bond dissociation energy (D_e) between two fragments (A and B) into the interaction energy (ΔE_{int}) and the preparation energy (ΔE_{prep}):

$$-D_e = \Delta E_{\text{int}} + \Delta E_{\text{prep}}. \quad (1)$$

The preparation energy, which pertains to the amount of energy required to distort and/or electronically excite the two fragments to their states in the complex, is defined as:

$$\Delta E_{\text{prep}} = E_A - E_A^0 + E_B - E_B^0. \quad (2)$$

$E_{A/B}$ and $E_{A/B}^0$ are the energies of the fragments for their geometries in the complex and as free ligands, respectively. To obtain these energies, all of the boron nitride, Lewis basic (LB) and Lewis acidic (LA) molecules as well as their complexes were re-optimized at the BP86/TZ2P level of theory. When the optimized geometries for the M05-2X/cc-pVTZ and BP86/TZ2P methods were compared, no significant differences were observed, see discussion in (ESI†).

The interaction energy (ΔE_{int}) can be decomposed into three terms: (1) the Pauli exchange repulsion term (ΔE_{Pauli}), (2) the electrostatic interaction energy (ΔE_{elstat}) between charge densities

of the fragments, and (3) the orbital interaction energy (ΔE_{orb}) which results from orbital mixing of the A and B fragments:

$$\Delta E_{\text{int}} = \Delta E_{\text{Pauli}} + \Delta E_{\text{elstat}} + \Delta E_{\text{orb}}. \quad (3)$$

The first term (ΔE_{Pauli}) is always positive while in most cases ΔE_{elstat} and ΔE_{orb} are negative. For more information regarding this method and its application in studying chemical bonds including donor–acceptor complexes the reader is referred to the literature.^{44–48}

The natural orbitals for chemical valence (NOCV) approach can be utilized to obtain both a qualitative and quantitative picture of the chemical bond (eqn (4)).⁴⁹ In this approach, the deformation density $\Delta\rho(r)$ is decomposed into pairwise ψ_k and ψ_{-k} complementary eigenfunctions (NOCVs) with eigenvalues of ν_k and ν_{-k} that have the same magnitude but opposite sign:

$$\Delta\rho(r) = \sum_{k=1}^{N/2} \nu_k [-\psi_{-k}^2(r) + \psi_k^2(r)]. \quad (4)$$

Positive and negative values describe, respectively, density accumulation and density depletion; the bond forms through flowing electron density from the negative part of the molecule (shown later in red color) to the positive part (shown in blue). For quantitative results, one can represent the orbital interaction energy in terms of the NOCV eigenvectors:

$$\Delta E_{\text{orb}} = \sum_{k=1}^{N/2} \nu_k [-F_{-k,-k}^{\text{TS}} + F_{k,k}^{\text{TS}}]. \quad (5)$$

where $F_{-k,-k}^{\text{TS}}$ and $F_{k,k}^{\text{TS}}$ are diagonal transition state Kohn–Sham matrix elements over the corresponding NOCVs. Therefore, eqn (4) and (5) provide the qualitative and quantitative pictures of a chemical bond even for asymmetric complexes. For further details on this approach please see the original paper.⁴⁹

The nature of the bonding in the Lewis base (LB) substituted adducts was also assessed using both natural bonding orbital (NBO)⁵⁰ and atoms-in-molecules (AIM)⁵¹ analyses. NBO population analyses were done at the M05-2X/cc-pVTZ level of theory by using the NBO suite available in Gaussian 09.⁵² AIM analyses were carried out at the same level of theory using the AIMALL software package.⁵³ Nucleus independent chemical shift (NICS)⁵⁴ computations were also performed using the gauge-independent atomic orbital (GIAO) method at the center of and 1 Å above the ring at the M05-2X/cc-pVTZ level of theory. All the electronic structure calculations were performed using Gaussian 09⁵² and ADF 2013⁵⁵ packages.

3 Results and discussion

As mentioned in the Introduction, the use of N-heterocyclic carbene (NHC),^{56–58} N-heterocyclic olefin (NHO)^{59–66} and Wittig ($\text{R}_3\text{PCR}'_2$)^{31,67,68} donors is becoming prevalent in main group element chemistry. Accordingly ImMe_2 , ImMe_2CH_2 and Me_3PCH_2 were chosen as donors within the boron nitride adduct series shown in Fig. 1.

The isolated $(\text{BN})_2$ and $(\text{BN})_3$ species, as well as higher oligomers, have been the subjects of numerous computational



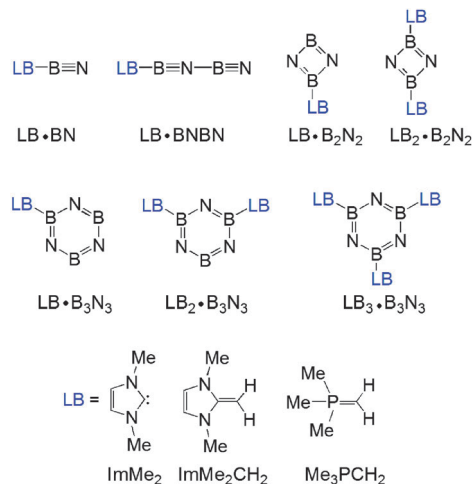


Fig. 1 Lewis base (LB) bound $(\text{BN})_n$ ($n = 1-3$) complexes considered in this study.

investigations;⁶⁹⁻⁷⁴ monomeric BN has also been studied extensively.⁷⁵⁻⁷⁷ The isolated systems are not the focus of the present study, however, results for the isolated species are reported for the corresponding isomers considered in the complexes for completeness. The isolated $(\text{BN})_n$ ($n = 1-3$) molecules are discussed in the ESI.†

3.1 Geometries of the Lewis base (LB) adducts $\text{LB} \cdot (\text{BN})_n$ ($n = 1-3$)

The M05-2X/cc-pVTZ optimized geometries of the mono-ligated boron nitride oligomers, $\text{LB} \cdot (\text{BN})_n$ ($n = 1-3$), are depicted in Fig. 2 ($\text{LB} = \text{ImMe}_2$, ImMe_2CH_2 , and Me_3PCH_2); see Fig. S1, ESI† for the optimized structures of the isolated species.

In each case, the carbon-based donors were bound to electron deficient boron sites, in line with prior adduct formation with amino-boranes ($\text{R}_2\text{N}-\text{BH}_2$).⁷⁸⁻⁸¹ The NHC-bound adducts of the BN chains, $\text{ImMe}_2 \cdot \text{BN}$ and $\text{ImMe}_2 \cdot \text{BNBN}$, each adopt linear $C_{\text{ImMe}_2}(\text{BN})_n$ configurations; geometry optimizations initiated with non-linear configurations return to linearity. On the other hand, appreciably bent C-B-N and intrachain N-B-N angles are found within the related ImMe_2CH_2 and Me_3PCH_2 -capped boron nitride adducts. For example, the C-B-N angles of the ImMe_2CH_2 substituted BN and BNBN adducts are 162.57° and 152.26° , respectively, and the C-B-N bond angles are 156.50° and 156.32° for the corresponding Me_3PCH_2 substituted analogues. In general, within the monoadducts, the B_2N_2 and B_3N_3 rings adopt planar geometries; the B_3N_3 rings in ImMe_2 and ImMe_2CH_2 are slightly puckered. The B-N bond lengths involving the donor-bound boron atoms in these B_2N_2 and B_3N_3 rings are each *ca.* 0.12 Å longer than the remaining B-N bonds involving donor-free, two coordinate boron centers. This bond lengthening implies an increase in B-N π -interactions and/or enhanced ionic contribution to the B-N σ -bonds. Within the ImMe_2 monoadduct series, the formally dative C-B linkages range from 1.510 Å in the terminal adduct $\text{ImMe}_2 \cdot \text{BNBN}$ to elongated values of 1.560 and 1.612 Å in the heterocyclic B_2N_2 and B_3N_3 adducts. For comparison, the C-B bond length in the coordinatively saturated amino-borane adduct $\text{IPr} \cdot \text{BH}_2\text{-NHDipp}$

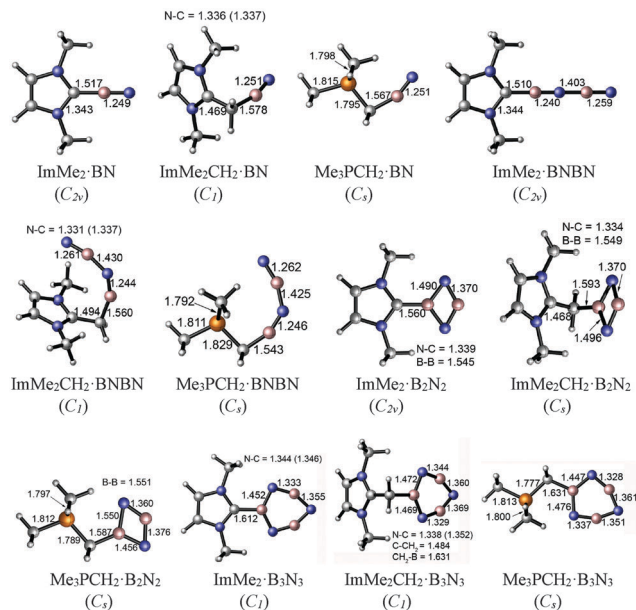


Fig. 2 M05-2X/cc-pVTZ optimized geometries including important bond lengths and corresponding symmetries of the mono-substituted adducts in the gas phase.

($\text{IPr} = [(\text{HCNDipp})_2\text{C}]$; $\text{Dipp} = 2,6\text{-}^i\text{Pr}_2\text{C}_6\text{H}_3$) was determined to be 1.627(4) Å,⁸¹ while in the diboryne adduct $\text{IPr} \cdot \text{B} \equiv \text{B} \cdot \text{IPr}$ this bond length is 1.491(4) Å (avg.).⁷ In general the computed C-B bonds in the ImMe_2 adducts were shorter by *ca.* 0.02 to 0.05 Å compared to the corresponding ImMe_2CH_2 and Me_3PCH_2 complexes; of note, it has been found that N-heterocyclic carbenes are stronger σ -donors than their N-heterocyclic olefin counterparts (such as IPrCH_2).⁶² The coordination of the Me_3PCH_2 units to boron leads to a large increase in the ylidic P-C bond length from 1.672 Å in the free ligand to bond length values as long as 1.829 Å in $\text{Me}_3\text{PCH}_2 \cdot \text{BNBN}$. This observation could be traced to a reduction of $\text{H}_2\text{C} \rightarrow \text{P}-\text{C}(\sigma^*)$ hyperconjugative interactions^{31,82} in Me_3PCH_2 as the terminal CH_2 unit participates in coordination to boron. The same phenomenon can be observed in the case of the ImMe_2CH_2 adducts: the terminal C-C bond length increases from 1.353 Å in the free ligand to 1.494 Å in the BNBN substituted adduct $\text{ImMe}_2\text{CH}_2 \cdot \text{BNBN}$. For the isolated ImMe_2CH_2 and Me_3PCH_2 ligands, the P-C and C-C bonds have Wiberg bond indices (WBIs) of 1.328 and 1.613, respectively, based on NBO analyses, see section 3.3. These are reduced to 0.906–1.046 and 1.026–1.284, respectively, upon binding of the $(\text{BN})_x$ units reflecting the loss of double bond character, and, hence, the corresponding bond elongation.

The M05-2X/cc-pVTZ optimized geometries of the di- and tri-substituted adducts $\text{LB} \cdot (\text{BN})_n$, $n = 2$ and 3) are depicted in Fig. 3. In general, addition of a second equivalent of Lewis base to the B_2N_2 and B_3N_3 units leads to elongation of the average C-B bond length. The C-B bonds increase in length by *ca.* 0.06–0.09 Å for the B_2N_2 rings, while a more modest increase of *ca.* 0.01 to 0.04 Å was noted upon binding two donors to a B_3N_3 unit. Despite the planar nature of the B_2N_2 rings, the intraring B-N distances within the bis-adducts $(\text{ImMe}_2)_2 \cdot \text{B}_2\text{N}_2$, $(\text{ImMe}_2\text{CH}_2)_2 \cdot \text{B}_2\text{N}_2$,



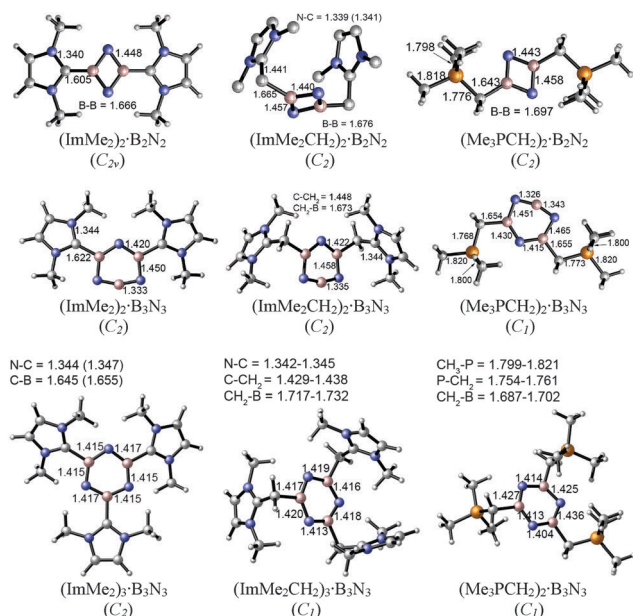


Fig. 3 M05-2X/cc-pVTZ optimized geometries and symmetries of the di- and tri-substituted adducts in the gas phase (C–H bonds are omitted in the case of $(\text{ImMe}_2\text{CH}_2)_2\cdot\text{B}_2\text{N}_2$ for clarity).

and $(\text{Me}_3\text{PCH}_2)_2\cdot\text{B}_2\text{N}_2$ all lie within a narrow range of 1.440 to 1.448 Å and suggest the absence of strong B–N π -bonding. To compare, iminoboranes ($\text{RB}\equiv\text{NR}'$) have B–N triple bond lengths in the range of 1.23 to 1.26 Å,^{83–86} while the diborylamide anion $[\text{Mes}_2\text{B}=\text{N}=\text{BMes}_2]^-$ ($\text{Mes} = 2,4,6\text{-Me}_3\text{C}_6\text{H}_2$), which has significant B=N double bond character, has B–N lengths of 1.343(5) and 1.348(5) Å.⁸⁷ The central B_3N_3 units in each complex adopt nearly planar arrangements with intraring B–N bond lengths that are typical for short B–N single bonds (1.404 to 1.436 Å). Each of the coordinative C–B distances are slightly longer in $(\text{ImMe}_2)_3\cdot\text{B}_3\text{N}_3$ (1.645 to 1.655 Å) in relation to the values found in the bisadduct $(\text{ImMe}_2)_2\cdot\text{B}_3\text{N}_3$ (1.622 Å). The ylide-bound tris adducts $(\text{Me}_3\text{PCH}_2)_3\cdot\text{B}_3\text{N}_3$ and $(\text{ImMe}_2\text{CH}_2)_3\cdot\text{B}_3\text{N}_3$ feature very long C–B bonds of 1.717–1.732 and 1.687–1.702 Å, respectively, suggesting that these species would have reduced stability.

Very recently, Tai and Nguyen have studied the stability of $(\text{ImMe}_2\cdot\text{B})_n$ ($n = 1\text{--}6$) adducts using quantum mechanical computations with the B3LYP method.⁸⁸ They attributed the stability of these systems to the degree of π conjugation and aromatic character within the core B_n ($n = 3\text{--}6$) rings. In order to probe the aromaticity in the $(\text{BN})_x$ rings, NICS analyses of the free (singlet) B_2N_2 and B_3N_3 molecules as well as their adducts were performed using the GIAO method at the M05-2X/cc-pVTZ level of theory. The NICS results were compared to the corresponding values determined at the same level of theory for well known aromatic benzene and anti-aromatic cyclobutadiene molecules (Table S1, ESI[†]) to examine changes in aromaticity upon binding of Lewis bases. NICS data are sensitive to the position at which they are evaluated and to interference from other parts of the molecule, especially for non-planar compounds.²³ The changes in aromaticity/anti-aromaticity are discussed in terms of NICS (1.00)_{zz} values, see Table S1, ESI[†] for complete NICS data.

Interestingly, NICS (1.00)_{zz} values show aromatic character for the free B_2N_2 and B_3N_3 molecules (−15.12 and −6.51 ppm, respectively, compared to −31.06 ppm for benzene). However, the free B_2N_2 loses aromatic character upon binding of one LB ligand (*i.e.*, NICS (1.00)_{zz} values of +2.55, +8.46, and +8.60 ppm for B_2N_2 complexed with Me_3PCH_2 , ImMe_2 , and ImMe_2CH_2 , respectively). For the doubly-bound adducts, $(\text{LB})_2\cdot\text{B}_2\text{N}_2$, the B_2N_2 unit becomes significantly anti-aromatic: +12.62, +13.94, and +9.04 ppm for the Me_3PCH_2 , ImMe_2 , and ImMe_2CH_2 adducts, respectively (see Table S1, ESI[†]). On the other hand, B_3N_3 remains moderately aromatic upon attachment of 1, 2, or 3 equivalents of Lewis base. The NICS (1.00)_{zz} decrease upon attaching two Lewis bases to the B_3N_3 ring with values of −8.55 and −3.19 ppm for ImMe_2 and ImMe_2CH_2 ligands, respectively, but slightly increases from −6.95 ppm to −6.39 ppm upon attaching the second Me_3PCH_2 ligand. For the case of the three LB bound adducts, the NICS (1.00)_{zz} values all increase (−2 to −3 ppm) upon attachment of the third ligand (Table S1, ESI[†]).

3.2 Energies of the Lewis base (LB) bound $\text{LB}\cdot(\text{BN})_n$ ($n = 1\text{--}3$) adducts

The total stabilization energies and Gibbs free energies of the $(\text{BN})_n$ ($n = 1\text{--}3$) molecules upon complexation with the three Lewis bases were computed using the M05-2X/cc-pVTZ level of theory and the results are summarized in Table 1. The sequential stabilization energies, $\Delta E_{\text{seq.}}$, $(\Delta E + \text{ZPE})_{\text{seq.}}$, and $\Delta G_{\text{seq.}}^{\circ}$, which take into account the impact of adding one additional Lewis base to the existing $(\text{LB})_x\cdot\text{B}_2\text{N}_2$ and $(\text{LB})_x\cdot\text{B}_3\text{N}_3$ ($x = 0\text{--}2$) complexes were also evaluated. Notably, in two separate articles, Jones, Frenking and co-workers have studied the ImMe_2 - and phosphine-bound Group 13 element complexes along with their possible applications for hydrogen storage.^{89,90} More specifically, they found that the Gibbs free energies of −29.8 and −45.8 kcal mol^{−1} for the Me_3P and ImMe_2 bound BH_3 adducts, respectively, at the RI-BP86/def2-TZVPP level of theory; which is very close to the −46.9 kcal mol^{−1} computed for the latter complex, $\text{ImMe}_2\cdot\text{BH}_3$, at the M06-2X/cc-pVDZ level of theory.⁹¹ In another recent study, Sarmah *et al.* examined complexes of normal and abnormal N-heterocyclic carbenes with Group 13 element based Lewis acids (EX_3 ; E = B, Al, Ga; X = H, F, Cl, OH, NH₂, CH₃, CF₃) and performed corresponding NBO and AIM analyses of the adducts.⁹² They computed a complexation energy of −49.2 kcal mol^{−1} at the B3LYP/6-31 + G* level of theory for the $\text{ImMe}_2\cdot\text{BH}_3$ adduct which is close to the values found previously by Frenking, Jones and co-workers^{89,90} as well as Brown and coworkers.⁹¹ The complexation (stabilization) energy associated with the formation of our mono-substituted $(\text{BN})_n$ ($n = 1\text{--}3$) adducts was computed to be greater than −100 kcal mol^{−1} for all species except the B_3N_3 adducts, where zero-point corrected energies ($\Delta E + \text{ZPE}$) are in the range of −57.4 to −69.7 kcal mol^{−1}. The ZPE correction to the electronic energies changes the value of ΔE by ~4–10 kcal mol^{−1}. The Gibbs free energy differences are also lower than the ZPE corrected values by ~10–40 kcal mol^{−1}. For the sake of brevity and consistency, the Gibbs free energy differences will be discussed throughout the text. We will comment on the nature of the formed



Table 1 Computed total^a and sequential^b stabilization energies (in kcal mol⁻¹), with ZPE ($\Delta E + ZPE$) and without ZPE (ΔE), and free energies (ΔG°) at the M05-2X/cc-pVTZ level of theory

Species	ΔE_{tot}^a	$(\Delta E + ZPE)_{\text{tot}}^a$	$\Delta G_{\text{tot}}^\circ{}^a$	$\Delta E_{\text{seq.}}^b$	$(\Delta E + ZPE)_{\text{seq.}}^b$	$\Delta G_{\text{seq.}}^\circ{}^b$
ImMe ₂ ·BN	-144.1	-140.5	-131.1	—	—	—
ImMe ₂ CH ₂ ·BN	-128.6	-123.8	-113.5	—	—	—
Me ₃ PCH ₂ ·BN	-145.7	-141.0	-130.7	—	—	—
ImMe ₂ ·BNBN	-129.5	-124.9	-114.5	—	—	—
ImMe ₂ CH ₂ ·BNBN	-126.6	-120.6	-107.7	—	—	—
Me ₃ PCH ₂ ·BNBN	-145.3	-139.7	-127.0	—	—	—
ImMe ₂ ·B ₂ N ₂	-118.2	-114.8	-103.5	-118.2	-114.8	-103.5
(ImMe ₂) ₂ ·B ₂ N ₂	-171.8	-165.9	-142.4	-53.6	-51.2	-38.9
ImMe ₂ CH ₂ ·B ₂ N ₂	-108.6	-104.0	-91.9	-108.6	-104.0	-91.9
(ImMe ₂ CH ₂) ₂ ·B ₂ N ₂	-143.4	-136.1	-109.6	-34.8	-32.0	-17.7
Me ₃ PCH ₂ ·B ₂ N ₂	-121.8	-117.7	-106.6	-121.8	-117.7	-106.6
(Me ₃ PCH ₂) ₂ ·B ₂ N ₂	-172.5	-166.2	-142.7	-50.7	-48.4	-36.1
ImMe ₂ ·B ₃ N ₃	-71.5	-68.7	-57.6	-71.5	-68.7	-57.6
(ImMe ₂) ₂ ·B ₃ N ₃	-122.9	-117.5	-92.6	-51.4	-48.9	-35.0
(ImMe ₂) ₃ ·B ₃ N ₃	-153.6	-147.4	-109.9	-30.7	-29.9	-17.3
ImMe ₂ CH ₂ ·B ₃ N ₃	-61.0	-57.4	-45.4	-61.0	-57.4	-45.4
(ImMe ₂ CH ₂) ₂ ·B ₃ N ₃	-97.9	-91.2	-65.3	-36.9	-33.8	-19.8
(ImMe ₂ CH ₂) ₃ ·B ₃ N ₃	-110.8	-102.0	-61.9	-12.9	-10.8	+3.4
Me ₃ PCH ₂ ·B ₃ N ₃	-72.8	-69.7	-58.5	-72.8	-69.7	-58.5
(Me ₃ PCH ₂) ₂ ·B ₃ N ₃	-121.1	-116.3	-94.1	-48.4	-46.6	-35.6
(Me ₃ PCH ₂) ₃ ·B ₃ N ₃	-141.5	-133.1	-96.3	-21.1	-18.4	-4.6

^a For the reaction: $(\text{BN})_n + x \cdot \text{LB} \rightarrow (\text{LB})_x \cdot (\text{BN})_n$ ($n = 1-3$, $x = 1-3$). ^b For the reaction: $\text{LB}_x \cdot (\text{BN})_n + \text{LB} \rightarrow (\text{LB})_{x+1} \cdot (\text{BN})_n$ ($n = 1-3$, $x = 0-2$).

carbene–boron bonds including their degrees of ionic/covalent character based on EDA as well as NBO/AIM analyses. The gap between the highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO) increases for all of the boron nitride species upon binding of the carbon-based ligands; the exceptions to this trend are the B₃N₃ adducts (Fig. S3, ESI[†]). For B₃N₃, the ImMe₂ and ImMe₂CH₂ bound adducts exhibit a modest decrease in the HOMO–LUMO gap upon binding the first ligand (for Me₃PCH₂ there is a small increase of ≈ 0.3 eV) and then for all ligands, there is a larger (≈ 1 eV) decrease upon binding the second and third ligands.

The free energies associated with the sequential addition of Lewis base equivalents to molecular B₂N₂ and B₃N₃ molecules $\Delta G_{\text{seq.}}^\circ$ follow the general trend that it becomes increasingly less favorable to bind multiple donors to these rings (Fig. 4).

This effect can be explained by a decrease in Lewis acidity of the (BN)_n rings as electron density is being donated from the carbon-based ligands; this phenomenon can be easily observed from the gradual destabilization of the LUMO energy levels of the B₂N₂ and B₃N₃ rings after addition of the Lewis bases (Fig. S3, ESI[†]). Overall, the binding of subsequent equivalents of Lewis base to the BN rings is exergonic, however, a slightly disfavoured binding event was computed for the formation of the tris adduct (ImMe₂CH₂)₃·B₃N₃ (+3.4 kcal mol⁻¹), placing this species on the cusp of stability.

3.3 Bonding properties through NBO and AIM analyses

NBO and AIM computations were performed on all (BN)_n species including their free and as well as their ligand bound

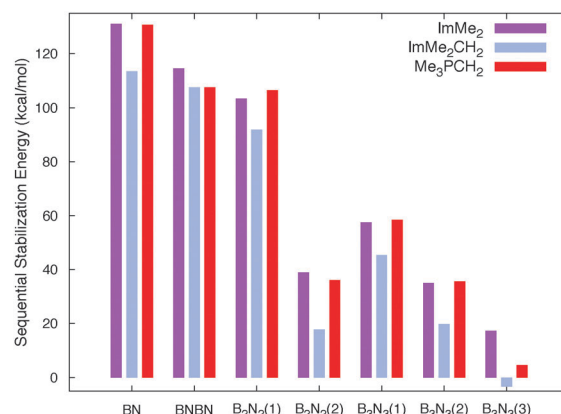


Fig. 4 The absolute values of the computed M05-2X/cc-pVTZ sequential Gibbs free stabilization energies (in kcal mol⁻¹) of all the adducts studied in this work. Numbers in parenthesis denote the number of Lewis base molecules attached to the B₂N₂ and B₃N₃ molecules. See Table 1 for the values.

forms at the M05-2X/cc-pVTZ level of theory (see Table 2; for the NBO and AIM analyses of the di- and tri-substituted compounds see Tables S2–S9 and Fig. S2, ESI[†] respectively). The computed NBO atomic charges of the boron atoms show a significant decrease (0.5 to 0.6 e⁻) upon attachment of the Lewis bases. However, the change in charge of the bonding carbon atom upon complexation is much more modest *ca.* 0.1 to 0.2 (Table 2 and Table S2, ESI[†]). The charge transfer to the boron center is highest for the ImMe₂ and lowest for the Me₃PCH₂ substituted BN and BNBN adducts. For both B₂N₂



Table 2 Selected NBO atomic charges of the carbene carbon and boron atoms (q_C and q_B) along with the total charge of the acceptor molecules (q_{LA}), Wiberg bond indices (WBI) of the C–B and B–N bonds, and the electron density (ρ) and energy density at the bond critical points ($H_{(C-B)}$ and $H_{(B-N)}$) for all the mono-substituted species at the M05-2X/cc-pVTZ level of theory. B–N values refer to bonds adjacent to the carbene carbon atom. Values in parentheses correspond to two different B–N bonds connected to the carbene carbon atom

Species	q_C	q_B	q_{LA}	WBI _{C-B}	WBI _{B-N}	$\rho(r_{C-B})$	$H(r_{C-B})$	$\rho(r_{B-N})$	$H(r_{B-N})$
ImMe ₂ ·BN	+0.203	+0.230	−0.524	0.944	2.622	0.179	−0.175	0.300	−0.324
ImMe ₂ CH ₂ ·BN	−0.821	+0.367	−0.551	0.772	2.656	0.163	−0.158	0.298	−0.324
Me ₃ PCH ₂ ·BN	−1.046	+0.382	−0.565	0.872	2.604	0.166	−0.162	0.299	−0.325
ImMe ₂ ·BNBN	+0.140	+0.608	−0.553	0.966	2.062	0.189	−0.193	0.290	−0.295
ImMe ₂ CH ₂ ·BNBN	−0.794	+0.838	−0.628	0.898	2.018	0.180	−0.184	0.298	−0.320
Me ₃ PCH ₂ ·BNBN	−1.075	+0.858	−0.603	0.931	1.997	0.185	−0.191	0.298	−0.322
ImMe ₂ ·B ₂ N ₂	+0.235	+0.654	−0.651	0.861	0.998	0.171	−0.170	0.192	−0.199
ImMe ₂ CH ₂ ·B ₂ N ₂	−0.738	+0.783	−0.622	0.801	0.969	0.166	−0.168	0.189	−0.194
Me ₃ PCH ₂ ·B ₂ N ₂	−1.037	+0.769	−0.622	0.840	1.096 (0.846)	0.167	−0.168	0.204 (0.172)	−0.214 (−0.171)
ImMe ₂ ·B ₃ N ₃	+0.251	+0.749	−0.592	0.837	1.082	0.154	−0.149	0.192 (0.192)	−0.192
ImMe ₂ CH ₂ ·B ₃ N ₃	−0.763	+0.863	−0.578	0.749	1.022 (1.096)	0.157	−0.157	0.184 (0.185)	−0.183 (−0.182)
Me ₃ PCH ₂ ·B ₃ N ₃	−1.062	+0.690	−0.572	0.689	1.145 (1.143)	0.154	−0.152	0.194 (0.255)	−0.195 (−0.179)

and B₃N₃ adducts, the highest charge transfer to the boron atom belongs to the ImMe₂ and Me₃PCH₂ ligands, respectively, but the lowest to the ImMe₂CH₂ ligand (Table 2). Interestingly, the NBO analysis does not show a significant charge difference for the nitrogen atom attached to the boron center in the LB-BN adducts compared to the isolated species (Tables S2 and S3, ESI†). On the other hand, in LB-BNBN adducts, the terminal nitrogen gains about 0.4–0.8 electrons and the central N atom loses about 0.2–0.4 electrons relative to the free species. For B₂N₂, the boron loses 0.1–0.2 electrons, while the nitrogen shows a 0.4 electron gain upon complexation; the changes in partial charges of B and N in B₃N₃ are very modest (0.1–0.2 e[−]). Therefore, there is stronger electronic communication in the BNBN complexes compared to the other adducts. Also, the Wiberg bond index (WBI) shows the following trend for all the C–B bonds in the mono-substituted adducts: LB-BNBN > LB-BN > LB-B₂N₂ > LB-B₃N₃; a rather similar trend can be observed for the AIM electron densities (ρ) of these bonds

(Table 2). On the other hand, the trend of the WBI for the B–N bonds is LB-BN > LB-BNBN > LB-B₂N₂ ≈ LB-B₃N₃.

Analysis of the energy density ($H(r)$) at the C–B and B–N bond critical points shows that all of these bonds are predominantly covalent in character (*i.e.*, negative values for $H_{(C-B)}$ and $H_{(B-N)}$). This data agrees well with the EDA-NOCV results which will be discussed later and points to the existence of covalent bonding between the carbon donors and boron acceptors, which is accompanied by π -backbonding in these systems as previously noted in LB-BX₃ (X = H, F, Cl) adducts.^{93,94} The computed value of the electron density (ρ) for the C–B bonds also shows that its strength decreases in the order of ImMe₂ > Me₃PCH₂ > ImMe₂CH₂ in the case of the BN, BNBN, and the B₂N₂ adducts. For the mono-substituted B₃N₃ adducts, the trend in ρ is ImMe₂CH₂ > ImMe₂ ≈ Me₃PCH₂ (Table 2) although the differences in ρ are very small (<0.003 e[−]). The ρ values for the B–N bonds in each of the adducts increase from 0.185 e[−] to 0.300 e[−] on going from the B₃N₃ adducts to the BN

Table 3 Computed EDA–NOCV components (in kcal mol^{−1}) for the C–B bonds of the BN and B₃N₃ substituted systems at the BP86/TZ2P level of theory. The C–B bond lengths (R , in Å) are also provided for all the complexes^a

	ImMe ₂ ·BN	ImMe ₂ CH ₂ ·BN	Me ₃ PCH ₂ ·BN	ImMe ₂ ·B ₃ N ₃	ImMe ₂ CH ₂ ·B ₃ N ₃	Me ₃ PCH ₂ ·B ₃ N ₃
ΔE_{int}	−136.1	−127.9	−135.8	−93.8	−84.5	−94.4
ΔE_{Pauli}	211.6	187.7	215.1	203.9	173.5	191.3
ΔE_{elstat}	−168.3	−139.3	−168.2	−151.5	−116.2	−139.0
	(48.4%)	(44.1%)	(47.9%)	(50.9%)	(45.0%)	(48.7%)
ΔE_{orb}	−179.3	−176.3	−182.7	−146.2	−141.8	−146.7
	(51.6%)	(55.9%)	(52.1%)	(49.1%)	(55.0%)	(51.3%)
$\Delta E_{\text{orb},\sigma}$	−115.8	−131.0	−130.6	−112.9	−118.9	−121.5
	(64.6%)	(74.3%)	(71.5%)	(77.2%)	(83.9%)	(82.8%)
$\Delta E_{\text{orb},\pi}$	−40.5	−33.8	−25.2	−16.7	−4.5	−7.6
	(22.6%)	(19.2%)	(13.8%)	(11.4%)	(3.2%)	(5.2%)
$\Delta E_{\text{orb},\text{rest}}$	−19.5	−11.5	−26.9	−16.6	−18.4	−17.6
	(12.8%)	(6.5%)	(14.7%)	(11.4%)	(12.9%)	(12.0%)
ΔE_{prep}	1.7	15.2	8.6	26.8	31.8	31.7
$-D_e$	−134.4	−112.7	−127.2	−67.0	−52.7	−62.7
R	1.508	1.582	1.569	1.605	1.658	1.647

^a For deformation densities see Fig. 5. ^b Percentage contributions to the total attractive interactions ($\Delta E_{\text{elstat}} + \Delta E_{\text{orb}}$) provided in parenthesis.

^c Percentage contributions to the total orbital interactions ΔE_{orb} provided in parenthesis.



adducts, in line with the corresponding increase in WBI values for these species. From the optimized geometries, we found that the C–B bond length increases upon substituting more Lewis bases. AIM data are in agreement with the geometries as, for example, the ρ value for this bond decreases from $0.154 e^-$ in the $\text{ImMe}_2\cdot\text{B}_3\text{N}_3$ complex to $0.136\text{--}0.139 e^-$ in $(\text{ImMe}_2)_3\cdot\text{B}_3\text{N}_3$ while the C–B bond length increases from 1.612 \AA to $\sim 1.650 \text{ \AA}$ (Fig. S2, ESI† and Fig. 2 and 3). This trend mirrors the variation in stabilization energies, and indicate that the interaction between the Lewis base and the boron atoms becomes weaker in the presence of added equivalents of donor. The ρ values of the C–B bonds for $(\text{ImMe}_2\text{CH}_2)_n\cdot\text{B}_3\text{N}_3$ and $(\text{Me}_3\text{PCH}_2)_n\cdot\text{B}_3\text{N}_3$ also decrease by going from mono- to tri-substituted adducts: from $0.157 e^-$ to $0.120 e^-$ and from $0.154 e^-$ to $0.127 e^-$, respectively (Fig. S2, ESI†); a trend reflected in the corresponding C–B bond length.

3.4 Energy decomposition analysis (EDA-NOCV)

To understand the nature of the bonding between different Lewis bases and the cyclic and acyclic boron nitride oligomers ($\text{LB}\cdot(\text{BN})_n$, $n = 1\text{--}3$), EDA-NOCV computations were performed using the GGA BP86 functional and the TZ2P basis set (Table 3). For brevity, we only focus on the most stabilized and least stabilized boron nitride species, *i.e.*, $\text{LB}\cdot\text{BN}$ and $\text{LB}\cdot\text{B}_3\text{N}_3$, respectively. The order of bond dissociation energies (D_e) for the different Lewis bases follows the series $\text{ImMe}_2 > \text{Me}_3\text{PCH}_2 > \text{ImMe}_2\text{CH}_2$. More specifically, the C–B bonds in the ImMe_2 substituted adducts are $4.3\text{--}21.7 \text{ kcal mol}^{-1}$ stronger than their Me_3PCH_2 and ImMe_2CH_2 analogues. For a given boron nitride adduct, there is a clear correlation between C–B bond length on one hand and bond dissociation energy and Pauli repulsion values on the other hand (Table 3).

The percentage contribution of the electrostatic attraction (ΔE_{elstat}) and orbital interaction (ΔE_{orb}) terms to the total attractive energies are also provided in Table 3. Overall, the orbital interaction makes a significant contribution to the total attractive energy (more than 50%) in all complexes except $\text{ImMe}_2\cdot\text{B}_3\text{N}_3$ where it is 49.1%. This high contribution indicates that C–B bonds retain substantial covalent character which is in agreement with our NBO/AIM results discussed above. The percentage contributions of the σ and π orbitals to the total orbital interaction are shown in Table 3 while the relevant deformation densities ($\Delta\rho$) are depicted in Fig. 5. Notably, the $\text{ImMe}_2\text{CH}_2\cdot\text{B}_3\text{N}_3$ adduct shows the lowest π -contribution to the C–B orbital interaction (3.2%) amongst the compounds investigated, while in contrast, the $\text{ImMe}_2\cdot\text{BN}$ and $\text{ImMe}_2\text{CH}_2\cdot\text{BN}$ adducts show the highest degree of π -character with 22.6% and 19.2% contributions, respectively. Thus from both Table 3 and Fig. 5 it is evident that π -backbonding between the boron nitride oligomers and the carbon-based ligands can be quite significant in some cases. The preparation energy (ΔE_{prep}), the difference between the fragment energies in their complexed and free geometries, is the lowest for the $\text{ImMe}_2\cdot\text{BN}$ while it is the highest for the $\text{ImMe}_2\text{CH}_2\cdot\text{B}_3\text{N}_3$ adduct (Table 3).

Our $-D_e$ values for the carbene–boron bonds are significantly more negative than the reported $-D_e$ values for the $\text{H}_3\text{B}\cdot\text{NH}_3$

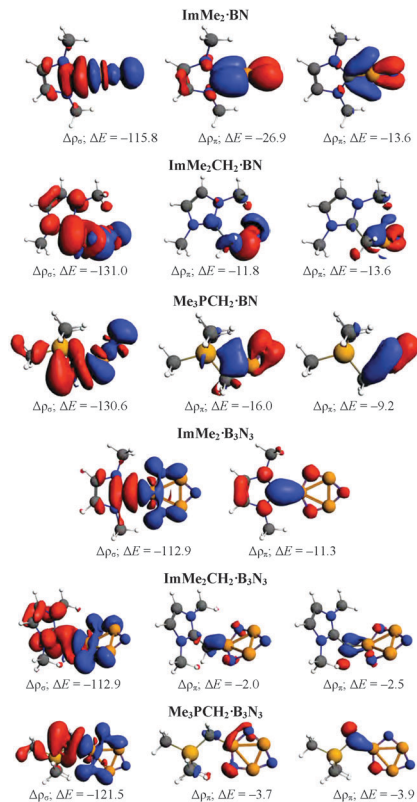


Fig. 5 Deformation densities ($\Delta\rho$) associated with the most important pairwise orbital interactions for C–B bond formation of different Lewis base substituted BN and B_3N_3 adducts. The charge flow is from red \rightarrow blue. Energies in kcal mol^{-1} are also provided.

($-31.9 \text{ kcal mol}^{-1}$) and $\text{H}_3\text{B}\cdot\text{NMe}_3$ ($-36.2 \text{ kcal mol}^{-1}$) bonds computed at the BP86/TZ2P level of theory.^{93–95} Tonner and Frenking have shown that replacing ammonia with the ImMe_2 ligand to form the $\text{ImMe}_2\cdot\text{BH}_3$ adduct changes the $-D_e$ to $-57.9 \text{ kcal mol}^{-1}$ computed at the same level.⁹⁶ Also, the amount

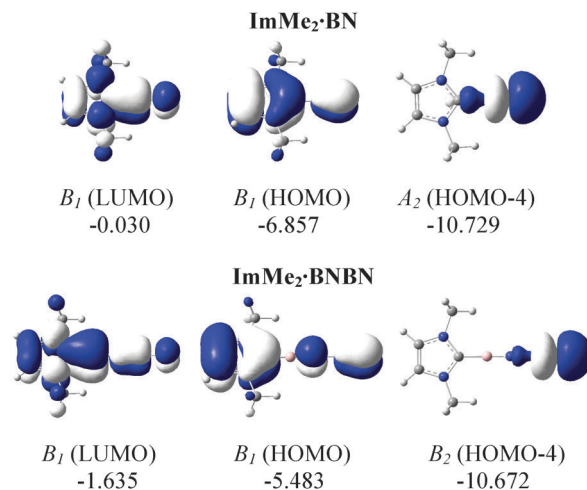


Fig. 6 M05-2X/cc-pVTZ computed relevant MOs for the $\text{ImMe}_2\cdot\text{BN}$ and $\text{ImMe}_2\cdot\text{BNBN}$ substituted adducts in the gas phase. Symmetries as well as energies (in eV) are also provided.



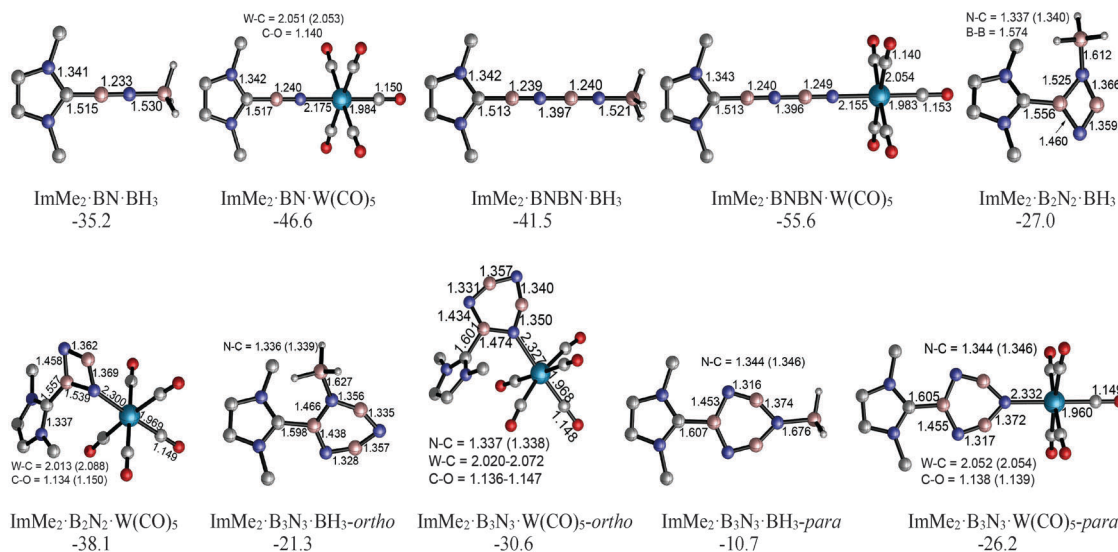


Fig. 7 M05-2X/cc-pVTZ optimized geometries and Gibbs free energies of the BH_3 and $\text{W}(\text{CO})_5$ substituted boron nitride adducts in the gas phase (C–H bonds are omitted for clarity).

of π -backbonding in $\text{H}_3\text{B}\cdot\text{NH}_3$, $\text{Me}_3\text{N}\cdot\text{BH}_3$, and $\text{ImMe}_2\cdot\text{BH}_3$ are 10.1%, 13.0%, and 9.4%, respectively. The values are comparable to each other for all these three systems and are close to the corresponding value for the $\text{ImMe}_2\cdot\text{B}_3\text{N}_3$ complex (Table 3).

3.5 Stabilization through donor–acceptor interactions

The HOMOs of both the $\text{ImMe}_2\cdot\text{BN}$ and $\text{ImMe}_2\cdot\text{BNBN}$ adducts have π character localized on the (terminal) BN unit as well as on the ImMe_2 ring (Fig. 6). On the other hand, the HOMO–4 of both complexes shows a directional lone pair on the terminal nitrogen atom (with some mixing with a B–N σ bond) ready to be captured by a Lewis acid (LA). Herein we consider the previously employed donor–acceptor approach for stabilizing highly reactive heavier Group 14 element dihydrides,¹⁴ by using BH_3 and $\text{W}(\text{CO})_5$ as Lewis acids (LA), and ImMe_2 as a Lewis base. The M05-2X optimized Lewis acid/base bound $(\text{BN})_n$ complexes as well as their complexation Gibbs free energies are shown in Fig. 7 (for a comparison between their electronic energies and Gibbs free energies see Table S12, ESI†). The C–B bond lengths in the Lewis acid bound adducts $\text{ImMe}_2\cdot\text{BN}\cdot\text{LA}$ and $\text{ImMe}_2\cdot\text{BNBN}\cdot\text{LA}$ (LA = BH_3 and $\text{W}(\text{CO})_5$) are in the narrow range of 1.513 to 1.517 Å and are nearly identical to the values found in the Lewis acid free BN and BNBN adducts (1.517 and 1.510 Å, respectively). However, in the donor–acceptor complexes, the B–N bond lengths in BN and BNBN slightly increase by 0.001–0.035 Å showing (modest) π -electron transfer from the B–N bond to the LA molecule. The computed N–B and N–W bond lengths for the BN and BNBN adducts (1.521–1.627 Å and 2.155–2.332 Å, respectively) are shorter than the BP86/TZ2P values for the $\text{H}_3\text{N}\cdot\text{BH}_3$ and $\text{H}_3\text{N}\cdot\text{W}(\text{CO})_5$ complexes (1.657 and 2.350 Å, respectively) due to a change in hybridization at nitrogen from sp^3 to formally sp^5 .

Overall, $\text{W}(\text{CO})_5$ appears to be a stronger Lewis acid compared to BH_3 as the ΔG° values for the former adduct series are more favorable (negative) by 9.3–15.5 kcal mol⁻¹ (Fig. 7); a

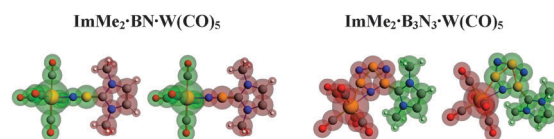


Fig. 8 Different fragments (shown in green and red colors) utilized for the EDA-NOCV computations of the C–B and N–W bonds in the $\text{ImMe}_2\cdot\text{BN}\cdot\text{W}(\text{CO})_5$ and $\text{ImMe}_2\cdot\text{B}_3\text{N}_3\cdot\text{W}(\text{CO})_5$ adducts.

similar conclusion regarding the relative Lewis acidity of $\text{W}(\text{CO})_5$ versus BH_3 has been made previously.⁴⁶ These results support our experimental results within the $\text{IPr}\cdot\text{GeH}_2\cdot\text{LA}$ complexes (LA = BH_3 and $\text{W}(\text{CO})_5$) where the $\text{W}(\text{CO})_5$ adduct is more stable.²⁵

The impact of complexing ImMe_2 and BH_3 molecules concurrently to the B_2N_2 and B_3N_3 units was studied. More specifically, the Gibbs free energies for the addition of the ImMe_2 ligand to the $\text{ImMe}_2\cdot\text{B}_2\text{N}_2\cdot(\text{BH}_3)_2$ and $(\text{ImMe}_2)_2\cdot\text{B}_3\text{N}_3\cdot(\text{BH}_3)_3$ adducts to form the fully saturated $(\text{ImMe}_2)_2\cdot\text{B}_2\text{N}_2\cdot(\text{BH}_3)_2$ and $(\text{ImMe}_2)_3\cdot\text{B}_3\text{N}_3\cdot(\text{BH}_3)_3$ complexes were found to be -74.2 and -60.9 kcal mol⁻¹, respectively (Table S10, ESI†).

Table 4 Computed EDA-NOCV components (in kcal mol⁻¹) for the C–B and N–W bonds of the $\text{ImMe}_2\cdot\text{BN}\cdot\text{W}(\text{CO})_5$ and $\text{ImMe}_2\cdot\text{B}_3\text{N}_3\cdot\text{W}(\text{CO})_5$ adducts at the BP86/TZ2P level of theory. The analogous values for the C–B bonds without Lewis acid are also provided in parenthesis. See Fig. 8 for the corresponding fragments

	$\text{ImMe}_2\cdot\text{BN}\cdot\text{W}(\text{CO})_5$		$\text{ImMe}_2\cdot\text{B}_3\text{N}_3\cdot\text{W}(\text{CO})_5$	
	C–B	N–W	C–B	N–W
ΔE_{int}	-172.6 (-136.1)	-57.6	-92.0 (-93.8)	-45.1
ΔE_{Pauli}	151.1 (211.6)	106.5	203.9 (220.7)	113.0
ΔE_{elstat}	-154.4 (-168.3)	-105.7	-151.5 (-158.1)	-89.8
ΔE_{orb}	-169.4 (-179.3)	-58.4	-146.2 (-154.6)	-68.3
$\Delta E_{\text{orb},\sigma}$	-113.5 (-115.8)	-31.6	-112.9 (-118.8)	-26.7
$\Delta E_{\text{orb},\pi}$	-30.9 (-40.5)	-10.7	-16.7 (-10.9)	-26.9
$\Delta E_{\text{orb},\text{rest}}$	-25.0 (-19.5)	-16.1	-16.6 (-24.9)	-14.7



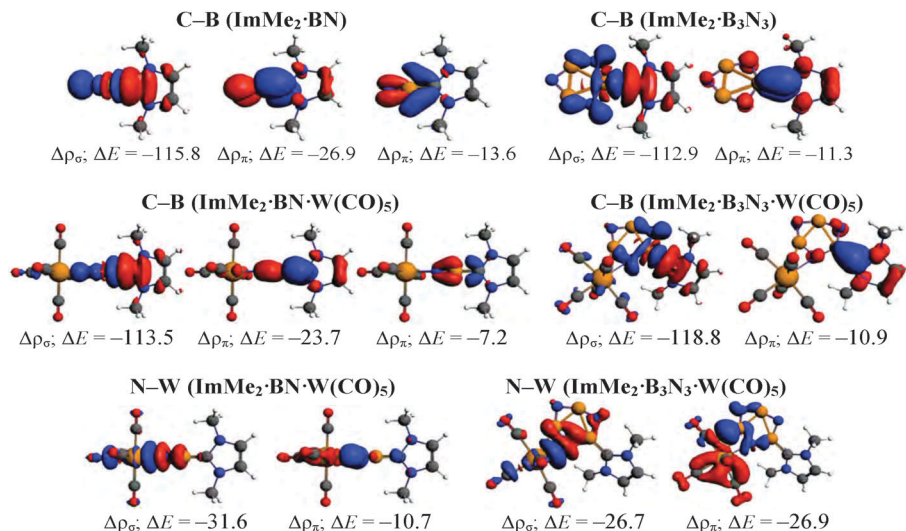


Fig. 9 Deformation densities ($\Delta\rho$) associated with the most important pairwise orbital interactions for the C–B and N–W bond formations in the ImMe₂·BN·W(CO)₅ and ImMe₂·B₃N₃·W(CO)₅ adducts. The charge flow is from red → blue. Energies in kcal mol⁻¹ are also provided.

3.6 EDA-NOCV for BN and B₃N₃ LB/LA substituted adducts

To further study the impact of adding a Lewis acid on stabilizing BN and B₃N₃ molecules, C–B and N–W bonds in the ImMe₂·BN·W(CO)₅ and ImMe₂·B₃N₃·W(CO)₅ adducts were examined using the EDA-NOCV approach (Fig. 8 and Table 4); their corresponding deformation densities are presented in Fig. 9.

Comparing the interaction energies in the BN and the B₃N₃ adducts reveals that the C–B bond becomes 36.5 kcal mol⁻¹ stronger upon W(CO)₅ Lewis acid attachment in the former adduct, but surprisingly it becomes 1.8 kcal mol⁻¹ weaker in the latter (Fig. 7). More specifically, addition of tungsten pentacarbonyl as a Lewis acid significantly decreases the Pauli repulsion portion of the C–B bond; from 211.6 kcal mol⁻¹ in ImMe₂·BN to 151.1 kcal mol⁻¹ in ImMe₂·BN·W(CO)₅ but increases it from 203.9 kcal mol⁻¹ in ImMe₂·B₃N₃ to 220.7 kcal mol⁻¹ in ImMe₂·B₃N₃·W(CO)₅ (Fig. 7). Lewis acid attachment also decreases the contribution of electrostatic and orbital interactions by 13.9 and 9.9 kcal mol⁻¹ for ImMe₂·BN adduct but it increases them to 6.6 and 8.4 kcal mol⁻¹ for the ImMe₂·B₃N₃ adduct. An inspection of the σ and π orbital interaction components for the C–B bonds in the BN adduct proves that the decrease in ΔE_{orb} upon bonding to the Lewis acid comes mainly from the decrease of π -backbonding rather than σ -donation. Moreover, comparing the percent contribution of the ΔE_{orb} component to the total interaction energy confirms the ionic nature of the N–W bonds (35.6% and 43.2% for the BN and B₃N₃ adducts, respectively). It is also worthwhile mentioning that no stationary point was found for the BN·W(CO)₅ or B₃N₃·W(CO)₅ adducts which points towards the instability of the N–W bond in these species in the absence of the Lewis base.

Given that the free energies of complexation associated with coordinating ImMe₂·BN and ImMe₂·BNBN units by BH₃ and W(CO)₅ are quite favorable, Lewis acid coordination can provide even more stability for these highly elusive boron nitride species,^{83,97,98} and research towards preparing these compounds in the laboratory is ongoing.

4 Conclusions

A variety of acyclic and cyclic (BN)_{*n*} (*n* = 1–3) adducts with different Lewis bases including an N-heterocyclic carbene, an N-heterocyclic olefin and a Wittig donor were examined using M05-2X/cc-pVTZ computations. Considering the Gibbs free energies, values greater than –50 kcal mol⁻¹ were found for the complexation energies. From the NBO, AIM and EDA-NOCV approaches, the existence of a polar covalent bond between carbene and boron atom was confirmed in each adduct studied. On the other hand, computed NPA charges illustrated rather significant amounts of charge transfer from the carbene center towards the boron atom upon C–B bond formation. A donor–acceptor strategy, in analogy with our synthesis of heavier group 14 element dihydride adducts,¹⁴ show that LB·(BN)_{*n*}·W(CO)₅ (*n* = 1–3) complexes could be experimentally achievable. Finally, both the C–B donor and N–W acceptor bonds were decomposed into their σ and π bonding components in the ImMe₂ substituted BN and B₃N₃ adducts with and without W(CO)₅ as a Lewis acid. Analysis of the EDA-NOCV results in these adducts showed that the carbene–boron bonds are stronger in the presence of W(CO)₅ as a Lewis acid mainly because of a dramatic decrease in Pauli repulsion rather than an increase in the electrostatic/orbital attraction.

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