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# Synthesis of perhalogenated silylboranes (X = Cl, I) and their application in regiodivergent alkene silaboration

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Silaboration of olefins is a synthetically valuable and atom-economic mode of functionalization; however, it typically requires transition-metal catalysis. We have overcome this requirement by using highly reactive perhalogenated silylboranes,  $X_2B-SiX_3$  (X = Cl, I), for which we herein report a straightforward synthesis, a full characterization, and their key properties. Access to this compound class was enabled by substantial improvement in the synthesis protocol for our previously published compound  $[Et_4N][I_3B-SiI_3]$ , now available on a 40 g scale *via* only two steps. Cation exchange with  $Li[Al(OC(CF_3)_3)_4]$  affords the mixture  $Li[I_3B-SiI_3]/I_2B-SiI_3/LiI$ , serving as a synthetic equivalent of the elusive pure  $I_2B-SiI_3$ . Its chlorine analogue  $Cl_2B-SiCl_3$  is accessible as a distillable liquid *via* treatment of  $[Et_4N][I_3B-SiI_3]$  with  $GaCl_3$ . For both perhalogenated silylboranes, various Lewis base adducts  $Do \cdot X_2B-SiX_3$  were obtained in excellent yields and structurally characterized by X-ray diffraction (Do =  $SMe_2$ , Py, PPh<sub>3</sub>, IDipp; IDipp = 1,3-bis(2,6-diisopropylphenyl)-1,3-dihydro-2H-imidazol-2-ylidene). We demonstrated that  $Me_2S \cdot I_2B-SiI_3$  undergoes efficient 1,2-silaboration of the challenging, non-activated substrate ethylene at rt with 0.1 eq.  $BI_3$  as promoter. In contrast,  $Li[I_3B-SiI_3]/I_2B-SiI_3/LiI$  effects a quantitative, unprecedented 1,1-silaboration of cyclohexene at rt. This remarkable reactivity switch was elucidated by experimental and quantum-chemical studies of the underlying steric and electronic factors.

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## Introduction

Once considered exotic and of limited utility, perhalogenated diborane(4) and disilane compounds (**I**, **III**; Fig. 1a) have recently emerged as valuable building blocks for purposes ranging from organic synthesis to materials development.<sup>1–6</sup> The direct bond between two Lewis-acidic sites in **I** and **III**, each bearing good leaving groups, presents both challenges and opportunities: on the one hand, this unique combination promotes spontaneous disproportionation and vigorous decomposition upon exposure to air and moisture.<sup>7–10</sup> On the other hand, it enables uncatalyzed diboration reactions using **I**,<sup>11–17</sup> the *in situ* generation of versatile  $[SiX_3]^-$  nucleophiles from **III** upon simple halide addition,<sup>18,19</sup> and extensive late-stage derivatization at the B–X and Si–X bonds of the primary products. Thus, in contrast to the abundant bis(pinacolato)diboron (pinB–Bpin), whose B atoms are electronically tamed by  $O=B$   $\pi$ -donation and serve primarily as transmetallation

partners in Suzuki–Miyaura cross-couplings,<sup>20–22</sup> type-I halogenoboranes are tailored for applications where the B atoms are to remain as property-determining functional units in the final molecule.<sup>23–29</sup> Likewise, the  $Si_2X_6/X^-$  trichlorosilylation system and the controlled disproportionation of  $Si_2X_6$  with  $NR_3$  (ref. 30–32) have proven valuable for the synthesis of extensively derivatizable organosilanes,<sup>33–36</sup> oligotetrelanes,<sup>37–42</sup> and silicon clusters.<sup>43–50</sup> In contrast to **I**, **III** undergoes no spontaneous 1,2-additions to unsaturated organic substrates, and theoretical studies predict a prohibitively high activation barrier without a catalyst.<sup>51,52</sup>

Given the indispensable role of borylated<sup>53,54</sup> and silylated<sup>55</sup> building blocks in synthesis, it is desirable to combine both types of functional groups within a single building block, for which silylboranes of the type  $R_2B-SiR_3$  are the most obvious candidates.<sup>56–60</sup> Electronically stabilized representatives such as the prominent pinB– $SiMe_2Ph$  typically require activation by (precious) metal complexes prior to addition across  $C=C$  double<sup>61–63</sup> or  $C\equiv C$  triple bonds.<sup>64–72</sup> In only a few cases, the addition of a (Lewis) base ( $KOtBu$ ,<sup>73,74</sup>  $KN(SiMe_3)_2$ ,<sup>75</sup>  $PR_3$ ,<sup>76</sup> pyridines<sup>77,78</sup>) has been sufficient to replace the transition metal catalyst in silaboration reactions. Yet, a significant proportion of base-catalyzed silylborane transformations results in incorporation of either the boryl<sup>74,79–84</sup> or the silyl<sup>85–91</sup> group,<sup>92</sup> while the respective counterpart is discarded. So far, a single

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Fig. 1 (a) Perhalogenated diboranes(4) (I), disilanes (III), and the perhalogenated silylboranes (II) studied in this work. (b) Previously studied silylboranes (IV) capable of undergoing uncatalyzed silaboration reactions (C atoms marked with asterisks bear *t*Bu substituents); computed transition state (TS) for the silaboration of ethylene with IV<sup>H</sup> (*t*Bu groups have been omitted in the calculations). (c) Silylborates H[V] and VI bearing some structural similarity with II.

uncatalyzed silaboration reaction has been reported, employing compounds IV<sup>H</sup> and IV<sup>Cl</sup> in THF (Fig. 1b).<sup>93,94</sup> Key to this transformation is the incorporation of both the B and Si atoms of IV into planar heterofluorene scaffolds, which—compared to pinB–SiMe<sub>2</sub>Ph—enhances their exposure to the unsaturated substrate while reducing  $\pi$ -electron donation into the vacant B(*p*<sub>z</sub>) orbital (quantum-chemical calculations exclude a promoting effect of the THF ligand on B–Si-bond cleavage; cf. the transition state TS of olefin silaboration shown in Fig. 1b). Based on this background and the high reactivity of I and III, we reasoned that the perhalogenated silylborane II (Fig. 1a) as a silaboration reagent should uniquely combine a strong tendency towards B–Si heterolysis and diverse opportunities for subsequent derivatization. Herein, we demonstrate that type-II compounds with X = Cl, I can indeed be readily synthesized on a multigram scale. We provide a full characterization of their B-adducts with various Lewis bases and show that the Cl derivative Cl<sub>2</sub>B–SiCl<sub>3</sub> can even be isolated in its free form as a distillable liquid. Notably, we disclose that both uncatalyzed 1,2- and rare 1,1-addition reactions to alkenes have been achieved. Only a few previously reported compounds share structural or electronic features with II. Among them are the borate H[V] and

the *nido* cluster VI (Fig. 1c).<sup>95,96</sup> Furthermore, the molecular structure of the anion [Cl<sub>3</sub>B–SiCl<sub>3</sub>]<sup>−</sup> has been determined through single crystal X-ray structure analysis of the salt [(TMS<sub>2</sub>N)SiCl<sub>2</sub>–B( $\eta^5$ -C<sub>5</sub>Me<sub>5</sub>)] [Cl<sub>3</sub>B–SiCl<sub>3</sub>] (TMS = Me<sub>3</sub>Si).<sup>97</sup>

## Results and discussion

The synthesis of B<sub>2</sub>X<sub>4</sub> (I) dates back to 1925, but for decades remained the domain of specialists capable of mastering the technically challenging gas-phase protocols of the time.<sup>98–100</sup> A major breakthrough came in 1981, when Nöth *et al.* obtained B<sub>2</sub>Br<sub>4</sub> in good yields by converting B<sub>2</sub>(OMe)<sub>4</sub> with BBr<sub>3</sub> through a convenient solution-phase synthesis.<sup>101</sup> In 2017, Braunschweig *et al.* extended this approach to the other perhalogenated diboranes(4) *via* solution-phase reactions of B<sub>2</sub>Br<sub>4</sub> with SbF<sub>3</sub>, GaCl<sub>3</sub>, and BI<sub>3</sub>.<sup>5</sup>

Si<sub>2</sub>Cl<sub>6</sub>, a side product of several large-scale processes in the silicon industry,<sup>102</sup> is commercially available; quantitative Cl/F exchange with SbF<sub>3</sub> affords Si<sub>2</sub>F<sub>6</sub>.<sup>103</sup> The perbromo- and periododisilanes are accessible from Si<sub>2</sub>Ph<sub>6</sub> by Ph/X exchange with MeC(O)X/AlX<sub>3</sub> (X = Br, I).<sup>104</sup>

Analogous to how B<sub>2</sub>Br<sub>4</sub> and Si<sub>2</sub>Cl<sub>6</sub> grant access to their respective compound classes, the salt [Et<sub>4</sub>N][I<sub>3</sub>B–SiI<sub>3</sub>] ([Et<sub>4</sub>N][1]; Scheme 1) serves as a key starting material for developing perhalogenated silylboranes. Several years ago, we first reported [Et<sub>4</sub>N][1], primarily to demonstrate the *in situ* formation of [SiCl<sub>3</sub>]<sup>−</sup> as the reactive intermediate in the Si<sub>2</sub>Cl<sub>6</sub>/Cl<sup>−</sup> trichlorosilylation system *via* Lewis-adduct formation with BX<sub>3</sub>.<sup>19</sup> Our study revealed that (excess) BI<sub>3</sub> is a more effective trapping reagent than BCl<sub>3</sub>, because it is the stronger Lewis acid and outcompetes coexisting Si<sub>2</sub>Cl<sub>6</sub> for coordination with [SiCl<sub>3</sub>]<sup>−</sup>, thus suppressing the formation of unwanted oligosilane side products.<sup>18</sup> By thoroughly optimizing the original protocol, the yield of [Et<sub>4</sub>N][1] was increased from  $\approx$  45% to  $\approx$  70%, and the synthesis was scaled to  $\approx$  40 g (Scheme 1). A key improvement is the addition of a second portion of BI<sub>3</sub> (0.1 eq.) toward the end of the reaction, following the initial addition of 2 eq. BI<sub>3</sub>. This prevents contamination of [Et<sub>4</sub>N][1] with [Et<sub>4</sub>N][I<sub>2</sub>ClB–SiI<sub>3</sub>], previously described as an ‘unknown side product’; its identity has now been unequivocally confirmed by X-ray crystallography (Fig. S102). This finding laid the foundation for a systematic exploration of perhalogenated silylboranes.

### Syntheses of new compounds

The quantitative Si–Cl/Si–I exchange during the formation of [Et<sub>4</sub>N][1] is advantageous, as iodinated products crystallize more readily in pure form than their chlorinated congeners. However, in combination with the salt-like nature of [Et<sub>4</sub>N][1], this results in extremely low solubility, posing challenges for subsequent transformations. As initial derivatizations, we consequently replaced one I<sup>−</sup> ligand in [Et<sub>4</sub>N][1] with neutral donor ligands (Do) bearing solubilizing substituents. To this end, suspensions of [Et<sub>4</sub>N][1] and Krossing's salt (Li [Al(OC(CF<sub>3</sub>)<sub>3</sub>)<sub>4</sub>])<sup>105</sup> in CH<sub>2</sub>Cl<sub>2</sub> were treated with the respective ligand at rt [Do: SME<sub>2</sub>, pyridine (Py), PPh<sub>3</sub>, 1,3-bis(2,6-diisopropylphenyl)-1,3-dihydro-2*H*-imidazol-2-ylidene (IDipp);





shielding when going from the  $\text{BI}_2(\text{Do})$  to the  $\text{BCl}_2(\text{Do})$  fragments. The  $^{29}\text{Si}$  NMR resonances of  $2\cdot\text{Do}$  and  $3\cdot\text{Do}$  were not detectable in the solution spectra, owing to unresolved  $^1\text{J}(\text{B},\text{Si})$  coupling and broadening induced by the quadrupolar  $^{11}\text{B}$  nucleus ( $S = 3/2$ ).<sup>108</sup> The  $^{31}\text{P}$  NMR spectra of  $2\cdot\text{PPh}_3$  and  $3\cdot\text{PPh}_3$  are characterized by multiplet resonances at  $-7.2$  and  $2.1$  ppm, respectively.

Electron ionization (EI) mass spectra were recorded for the full series of donor adducts  $2\cdot\text{Do}$  and  $3\cdot\text{Do}$  (introduced as solids). In most cases, we observed the molecular ion peak  $[(\text{Do})\text{X}_2\text{B}-\text{SiX}_3]^+$  and/or the peak corresponding to the donor-free silylborane  $[\text{X}_2\text{B}-\text{SiX}_3]^+$ , typically with low intensity ( $\text{X} = \text{Cl}, \text{I}$ ; see the SI for details). Most adducts appear to release their neutral Do ligand under the measurement conditions—except for IDipp, which resists elimination. Prominent fragmentation products included  $[(\text{Do})\text{BSiX}_4]^+ / [\text{BSiX}_4]^+$  and  $[(\text{Do})\text{BX}_2]^+$ . The latter may arise either by  $[\text{SiX}_3]^-$  loss from the parent ion or *via* a concerted pathway:  $[\text{SiX}_2]$  extrusion from  $[(\text{Do})\text{X}_2\text{B}-\text{SiX}_3]^+$ , followed by  $\text{X}^-$  elimination from the resulting  $[(\text{Do})\text{BX}_3]^+$  intermediate. This, in turn, raises the question—relevant for later reactivity studies—of whether neutral  $2\cdot\text{Do}$  and  $3\cdot\text{Do}$  might undergo thermal  $[\text{SiX}_2]$  extrusion. To probe this,  $2\cdot\text{IDipp}$  was heated with the silylene-trapping reagent 2,3-dimethyl-1,3-butadiene (DMB; 10 eq.) in *o*DFB at  $100^\circ\text{C}$  for 10 days in a flame-sealed NMR tube.  $[\text{SiI}_2]$  was subsequently identified by GC-MS as its cycloadduct, 1,1-diiodo-3,4-dimethyl-1-silacyclopent-3-ene (Fig. S2).<sup>109</sup> Consistently, the reaction mixture showed an  $^{11}\text{B}$  NMR signal corresponding to the byproduct  $\text{BI}_3\cdot\text{IDipp}$  formed through reductive elimination at the  $\text{Si}(\text{IV})$  center of  $2\cdot\text{IDipp}$  ( $-77.3$  ppm; confirmed by comparison with an authentic sample and X-ray crystal structure analysis of a single crystal grown in the NMR tube).

All eight adducts  $2\cdot\text{Do}$  and  $3\cdot\text{Do}$  were structurally characterized by X-ray crystallography (Fig. S103–S106 and S109–S112).<sup>110</sup> Given the particular relevance of  $2\cdot\text{SMe}_2$  to silaboration reactions (see below), the molecular structures of this compound and its perchlorinated congener  $3\cdot\text{SMe}_2$  are shown as representative examples in Fig. 2a. All B–Si-bond lengths in  $2\cdot\text{Do}/3\cdot\text{Do}$  fall within a narrow range of 2.005(3) to 2.043(4) Å, indicating that this parameter is not significantly influenced by either the nature of Do or the halogen ligand. In contrast, the B–Do bond lengths and the degree of pyramidalization at the B atom in the  $\text{X}_2\text{BSi}$  fragments support the *a priori* expectations that (i)  $\text{SMe}_2$  is the weakest among the four donors Do, and (ii) the iodinated compound **2** is more Lewis acidic than its chlorinated analogue **3**.<sup>111</sup>

Each molecule of **4** contains two chiral B atoms, giving rise to four possible diastereomers (Fig. 2b, left). In the lattice of the examined single crystal, two diastereomers [(S,S)/(S,R)] occupy the same crystallographic site in a 73 : 27 ratio, which leads to partial disorder involving the B(2)–Si(3) unit. This disorder, together with the comparatively weak scattering contribution of the light B atoms relative to the multiple heavy I atoms, limits the precision with which the B-atom positions and associated structural parameters can be determined. The five-membered ring in **4** features bridging I atoms (B– $\mu$ (I)–B and B– $\mu$ (I)–Si), resulting in tetracoordinate rather than tricoordinate, electron-

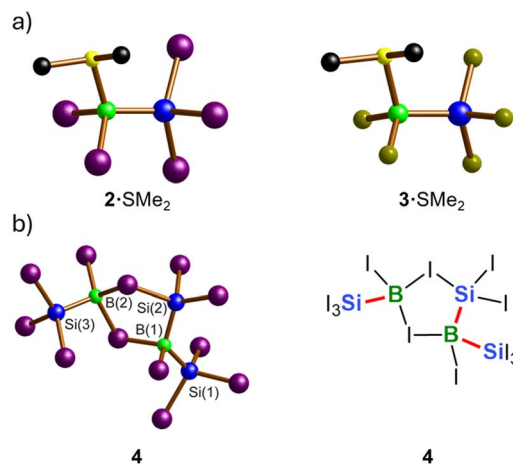


Fig. 2 (a) Solid-state structures of  $2\cdot\text{SMe}_2$  ( $\beta$ -polymorph; left) and  $3\cdot\text{SMe}_2$  (right); (b) solid-state structure of the (S,S)-diastereomer of **4** (left) and its corresponding structural formula, with one BSi and one BSi<sub>2</sub> moiety highlighted in red (right). H atoms are omitted for clarity. Color code: B: green, C: black, Si: blue, S: yellow, Cl: yellow-green, I: violet.

deficient B sites (Fig. 2b, left).<sup>112</sup> This feature prompts speculation that the extremely insoluble species **2** may adopt a polymeric or dimeric structure in the solid state, possibly featuring  $\text{B}_2\text{I}_2\text{Si}$  five-membered rings, with a single I atom replacing the  $\text{Si}(\text{I})_3$  substituent. Moreover, the presence of both a BSi and a BSi<sub>2</sub> moiety in **4** (indicated by red-colored bonds in Fig. 2b, right) suggests that our methodology may provide access not only to perhalogenated silylboranes but also to disilylboranes.

### Silaboration reactions with $2\cdot\text{Do}$ and $3\cdot\text{Do}$

One of the primary motivations for developing  $2\cdot\text{Do}$  and  $3\cdot\text{Do}$  was to create highly reactive silaboration reagents that allow for the simultaneous introduction of both derivatizable functional groups, ideally under catalyst-free conditions. Ethylene was selected as the initial model substrate for two main reasons: (i) its gaseous nature and lack of activating substituents make its silaboration particularly challenging, and (ii) the expected products are highly symmetric molecules with low molecular

Table 1 Conditions and product distributions for the reactions of the adducts  $2\cdot\text{Do}$  or  $3\cdot\text{Do}$  with an excess of ethylene in sealed NMR tubes

Adduct	Conditions	Product(s)
$2\cdot\text{SMe}_2$	$\text{CD}_2\text{Cl}_2$ , 6 d, $80^\circ\text{C}$	$5\cdot\text{SMe}_2$ , 98%
$2\cdot\text{SMe}_2/0.1 \text{ BI}_3$	$\text{CD}_2\text{Cl}_2$ , 12 h, rt	$5\cdot\text{SMe}_2$ , 85%
$2\cdot\text{Py}$	<i>o</i> DFB, 20 d, $120^\circ\text{C}$	$5\cdot\text{Py}$ , 97%
$2\cdot\text{PPh}_3$	<i>o</i> DFB, 20 d, $120^\circ\text{C}$	$5\cdot\text{PPh}_3$ , $\text{BI}_3\cdot\text{PPh}_3^b$
$2\cdot\text{IDipp}$	<i>o</i> DFB, 6 d, $100^\circ\text{C}$	$\text{BI}_3\cdot\text{IDipp}^b$
$3\cdot\text{SMe}_2$	$\text{CD}_2\text{Cl}_2$ , 31 d, $80^\circ\text{C}$	$6\cdot\text{SMe}_2$ , $\text{BCl}_3\cdot\text{SMe}_2^b$
$3\cdot\text{Py}$	<i>o</i> DFB, 7 d <sup>a</sup> , $120^\circ\text{C}^a$	$\text{BCl}_3\cdot\text{Py}^b$
$3\cdot\text{PPh}_3$	<i>o</i> DFB, 17 d, $120^\circ\text{C}$	$\text{BCl}_3\cdot\text{PPh}_3^b$
$3\cdot\text{IDipp}$	<i>o</i> DFB, 7 d <sup>a</sup> , $120^\circ\text{C}^a$	$\text{BCl}_3\cdot\text{IDipp}^b$

<sup>a</sup> After the initial heating period, heating was continued at  $140^\circ\text{C}$  for 1 day and at  $160^\circ\text{C}$  for 1 day. <sup>b</sup> The reaction mixture contained unconsumed starting material.



weight, which facilitates analysis by NMR spectroscopy and mass spectrometry.<sup>61,94</sup>

The reactions were carried out by heating an excess of ethylene with 2·Do or 3·Do in CD<sub>2</sub>Cl<sub>2</sub> or *o*DFB in sealed NMR tubes (Table 1 and Scheme 2a). *o*DFB was used when high temperatures and/or prolonged reaction times posed a risk of I/Cl exchange with CD<sub>2</sub>Cl<sub>2</sub>; for NMR measurements, *o*DFB was replaced with CD<sub>2</sub>Cl<sub>2</sub> prior to measurement.

To evaluate general reactivity trends, we employed the pure silylborane adducts without added promoters. Under these conditions, 2·SMe<sub>2</sub> underwent quantitative conversion with



Scheme 2 (a) 1,2-Silaborations of ethylene with 2·SMe<sub>2</sub> or 2·Py afford 5·SMe<sub>2</sub> or 5·Py. (b) 1,1-Silaboration of cyclohexene with [Et<sub>4</sub>N]1/Li[Al(OC(CF<sub>3</sub>)<sub>3</sub>)<sub>4</sub>] furnishes 7. (c) Reaction of phenylacetylene with 3·SMe<sub>2</sub> leads to the addition of the 3/SMe<sub>2</sub> Lewis pair across the C≡C bond to give 8. (d) Solid-state structures of 5·Py, 7, and 8; H atoms are omitted for clarity. Color code: B: green, C: black, N: pale blue, Si: blue, S: yellow, Cl: yellow-green; I: violet. Reagents and conditions: (i) exc. ethylene, CD<sub>2</sub>Cl<sub>2</sub>, 80 °C, 6 d, 98% yield; (ii) exc. ethylene, 0.1 eq. BI<sub>3</sub>, CD<sub>2</sub>Cl<sub>2</sub>, rt, 12 h, 85% yield; (iii) exc. ethylene, *o*DFB, 120 °C, 20 d, 97% yield; (iv) 10 eq. cyclohexene, 1 eq. Li[Al(OC(CF<sub>3</sub>)<sub>3</sub>)<sub>4</sub>], *o*DFB, rt, 15 min, 99% yield; (v) 5 eq. phenylacetylene, CH<sub>2</sub>Cl<sub>2</sub>, rt, 1 d, 37% yield.

ethylene to afford the 1,2-silaboration product Me<sub>2</sub>S·I<sub>2</sub>B-C<sub>2</sub>H<sub>4</sub>-SiI<sub>3</sub> (5·SMe<sub>2</sub>; Table 1 and Scheme 2a). This transformation proceeded to completion at 80 °C within 6 days. Likewise, 2·Py gave similarly high yields, albeit under even harsher conditions (120 °C, 20 d). At similar temperatures and reaction times, 2·PPh<sub>3</sub> was only partially consumed; the fraction that reacted generated both the silaboration product 5·PPh<sub>3</sub> and the thermolysis product BI<sub>3</sub>·PPh<sub>3</sub>. Among the perchlorinated analogues, only 3·SMe<sub>2</sub> produced a notable amount of the corresponding 1,2-silaboration product. However, this transformation took five times longer than the reaction of 2·SMe<sub>2</sub> and furnished Me<sub>2</sub>S·Cl<sub>2</sub>B-C<sub>2</sub>H<sub>4</sub>-SiCl<sub>3</sub> (6·SMe<sub>2</sub>; Table 1) contaminated with residual starting material and the side product BCl<sub>3</sub>·SMe<sub>2</sub>. No silaboration was observed for the other adducts 2·Do and 3·Do; instead, they formed varying amounts of BX<sub>3</sub>·Do, likely due to thermally induced [SiX<sub>2</sub>] extrusion, as discussed above and corroborated by our previous trapping experiments with DMB. To promote the reaction between 2·SMe<sub>2</sub> and ethylene, BI<sub>3</sub> (0.1 eq.) was added to the mixture. Now, silaboration proceeded smoothly at rt within 12 h, affording 5·SMe<sub>2</sub> in high yields (85%; entry 2 in Table 1). This final result of our systematic screening thus offers a practical and efficient access route to this promising functionalized building block. Notably, neither the Li[1]/2/LiI mixture nor free 3 provided further improvement, as both led to pronounced side reactions, presumably including ethylene polymerization. We further emphasize that haloboration did not compete with silaboration under any of the tested conditions.

Based on these experimental findings, two key principles emerge to guide further synthetic applications: (i) the highly reactive free species 2 and 3 must be tamed by adduct formation with a suitable donor to prevent non-selective transformations. In this regard, the soft ligand SMe<sub>2</sub> performs best in terms of product selectivity while still allowing for reasonable temperatures and reaction times—especially when 0.1 eq. of BI<sub>3</sub> is added as a promoter, which likely generates small concentrations of the free Lewis acid 2 *in situ*. (ii) The iodinated adducts are more effective in silaborations than their chlorinated congeners. From that, we offer the following mechanistic interpretations: (i) silaborations with 2·Do and 3·Do are apparently not initiated by donor-induced B-Si-bond cleavage. Instead, displacement of Do by π-bonded ethylene must precede the 1,2-addition step (comparable B···olefin complexes have been structurally characterized by Yamaguchi *et al.*<sup>113</sup>). (ii) To maximize the interaction between the vacant B(p<sub>z</sub>) orbital and the π-electron cloud of ethylene, competing π-backbonding from X to B must be minimized, which accounts for the superior suitability of X = I (2-type compounds) over Cl (3-type compounds). A comprehensive quantum-chemical assessment of the overall reaction mechanism is provided below.

In a second reactivity test, a mixture of 2·SMe<sub>2</sub> and the internal olefin cyclohexene in *o*DFB was heated to 120 °C for 24 days. Subsequent <sup>11</sup>B NMR analysis of the sample showed essentially one signal at -11.6 ppm, indicating quantitative and selective conversion. As such forcing reaction conditions lack practical relevance, efforts were directed toward significantly accelerating the reaction prior to detailed product analysis.



In this instance, the addition of  $\text{BI}_3$  as a promoter did not prove beneficial. However, a successful outcome was ultimately achieved using an equimolar mixture of  $[\text{Et}_4\text{N}][\mathbf{1}]$  and  $\text{Li}[\text{Al}(\text{OC}(\text{CF}_3)_3)_4]$  in *o*DFB, which effected complete conversion within only 15 min at rt. It is evident that the increased kinetic protection of the  $\text{C}=\text{C}$  double bond in this case suppresses unwanted side reactions, even in the absence of any donor ligand apart from the residual  $\text{I}^-$  ions. More remarkably, olefin internalization exerts a decisive effect on regioselectivity: the reaction with cyclohexene selectively afforded the 1,1-silaboration product **7** rather than the 1,2-isomer (Scheme 2).<sup>114</sup> Such a transformation is unprecedented—not only in silaboration but also in the related diboration or disilylation of olefins.<sup>115</sup>

In a final test experiment, phenylacetylene was chosen as the third representative substrate. Since the iodinated  $2 \cdot \text{SMe}_2$  led to complex product mixtures, we turned to the chlorinated analogue  $3 \cdot \text{SMe}_2$ , which underwent complete conversion at rt after 1 day. From the reaction mixture, the zwitterionic species **8** crystallized in 37% yield (Scheme 2). Unlike  $5 \cdot \text{Do}$  and **7** ( $\text{Do} = \text{SMe}_2, \text{Py}$ ), **8** is not generated *via* silaboration but instead represents the typical outcome of a concerted reaction between a free thioether Lewis base and a free borane Lewis acid acting on the same  $\text{C}\equiv\text{C}$  triple bond.<sup>116–122</sup> This finding thus supports our earlier assumption that replacement of the B-bonded donor  $\text{Do}$  with the unsaturated substrate constitutes the initial step in the reactions of  $2 \cdot \text{Do}$  and  $3 \cdot \text{Do}$ . In the case of olefin substrates, both a boryl and a silyl group are introduced into the molecule. Yet, with phenylacetylene, the B–Si bond remains intact, and  $\text{SMe}_2$  is instead transferred to the substrate. In **8**, the B atom is attached to the terminal position of the resulting olefin, while the  $\text{SMe}_2$  substituent resides near the phenyl ring. This can be explained by the fact that the positive charge accumulated on the carbon framework during electrophilic borylation is better stabilized by resonance at the  $\alpha$ -position relative to the phenyl ring.

### NMR-spectroscopic and X-ray crystallographic characterization of $5 \cdot \text{Do}$ , **7**, and **8** (ref. 107)

The  $^{11}\text{B}$  NMR spectra of the 1,2-silaboration products,  $5 \cdot \text{SMe}_2$  and  $5 \cdot \text{Py}$ , exhibit resonances at  $-18.9$  and  $-14.0$  ppm, respectively, consistent with the presence of tetracoordinate B nuclei.<sup>108</sup> In contrast, the  $^{11}\text{B}$  NMR signal of the 1,1-silaboration product **7** appears at  $53.5$  ppm, indicative of a tricoordinate B center.<sup>108</sup> The  $^{29}\text{Si}$  NMR shifts of  $5 \cdot \text{SMe}_2$ ,  $5 \cdot \text{Py}$ , and **7** are very similar with values of  $-115.1$ ,  $-114.8$ , and  $-122.8$  ppm, respectively. Furthermore, all three compounds give rise to signals exclusively in the aliphatic region of their  $^1\text{H}$  NMR spectra, confirming complete consumption of the  $\text{C}=\text{C}$  double bonds present in the starting materials. The resonances of the axial and equatorial H atoms within the cyclohexyl moiety of compound **7** are distinctly resolved, indicating that bulky substituents on the saturated ring act as effective conformational locks on the NMR timescale.<sup>123,124</sup> The  $^{11}\text{B}$  NMR spectrum of **8** is characterized by a resonance at  $-2.6$  ppm. As in the cases of  $2 \cdot \text{Do}$  and  $3 \cdot \text{Do}$ , the  $^{29}\text{Si}$  NMR signal of the B-bonded Si atom is broadened beyond detection. A singlet at  $\delta(^1\text{H}) = 7.35$ ,

together with a corresponding broad resonance at  $\delta(^{13}\text{C}) = 163.1$ , is consistent with the presence of an olefinic fragment in **8**.

The molecular structures of  $5 \cdot \text{SMe}_2$  (Fig. S115),  $5 \cdot \text{Py}$ , and **7** (Scheme 2d), confirm their proposed identities as 1,2- and 1,1-silaboration products, respectively. The C–C-bond length in  $5 \cdot \text{Py}$  falls within the typical single-bond range ( $1.533(4)$  Å), as do all C–C bonds in **7**. As expected,<sup>123,124</sup> the bulkier  $\text{SiI}_3$  substituent occupies an equatorial position, whereas the less bulky  $\text{BI}_2$  group adopts an axial orientation in the cyclohexane ring of **7**. In contrast to compound **4** (Fig. 2), there is no B– $\mu(\text{I})$ –Si bridge in **7**; rather, the boryl group remains trigonal-planar coordinated. Nonetheless, the vacant B( $p_z$ ) orbital may acquire some electron density from the occupied Si–C  $\sigma$  orbital, reminiscent of the well-known stabilization of carbenium ions bearing  $\beta$ -positioned silyl groups (see below).<sup>125</sup> The C(1)–C(2) distance in compound **8** is  $1.335(5)$  Å, characteristic of a double bond (Scheme 2d). The S and B atoms adopt a mutual *E* configuration, with the sterically demanding (silyl)boryl substituent located at the terminal position of the styrene core.

### Quantum-chemical calculations rationalizing the 1,2- vs. 1,1-silaboration of ethylene vs. cyclohexene to give $5 \cdot \text{SMe}_2$ vs. **7**

For the reactions of  $2 \cdot \text{SMe}_2$  and  $[\text{Et}_4\text{N}][\mathbf{1}]/\text{Li}[\text{Al}(\text{OC}(\text{CF}_3)_3)_4]$  with the olefins, two scenarios were examined: 1,2-silaboration and 1,1-silaboration. Potentially competing haloboration pathways<sup>126</sup> as well as, for  $2 \cdot \text{SMe}_2$ , the hypothetical addition of the  $\text{Me}_2\text{S}/\text{I}_2\text{B}-\text{SiI}_3$  Lewis pair to ethylene, were also considered but found to be irrelevant (see the SI for corresponding reaction pathways, activation barriers, and reaction energies). Fig. 3 shows the plausible silaboration sequences for ethylene (a) and cyclohexene (b). As a first important result, the experimentally observed products correspond to pathways that are both kinetically and thermodynamically favored (highlighted in red).

In the reaction of  $2 \cdot \text{SMe}_2$  with ethylene, the  $\text{SMe}_2$  donor must first dissociate to generate a vacant coordination site at the B atom for olefin binding. The dissociation requires an energy input of  $14.6$  kcal mol $^{-1}$  (*cf.* Fig. S134). However, the presence of  $\text{BI}_3$  renders the *in situ* release of the active silaboration reagent **2** significantly less endergonic ( $2 \cdot \text{SMe}_2 + \text{BI}_3 \rightarrow 2 + \text{BI}_3 \cdot \text{SMe}_2$ ;  $\Delta G = 4.8$  kcal mol $^{-1}$ ). Subsequent ethylene binding to free **2** is endergonic by an additional  $5.4$  kcal mol $^{-1}$ . The resulting intermediate,  $2 \cdot (\text{C}_2\text{H}_4)$ , features a strongly pyramidalized B atom [ $\sum(\text{I}-\text{B}-\text{I}/\text{Si}) = 320.2^\circ$ ]; the ethylene ligand remains essentially planar.<sup>127</sup> The reaction proceeds *via* transition state **TS1**, characterized by B–Si-bond cleavage and the concerted formation of a C–Si bond. The 1,2-silaboration product **5** lies  $-21.7$  kcal mol $^{-1}$  below the starting materials, with an overall activation barrier of only  $15.7$  kcal mol $^{-1}$ . In the final step, **5** acquires an  $\text{SMe}_2$  ligand from  $2 \cdot \text{SMe}_2$  to afford  $5 \cdot \text{SMe}_2$  and free **2** with a similar endoergicity as observed in the case of  $2 \cdot \text{SMe}_2/\text{BI}_3$ , explaining why only minor amounts of  $\text{BI}_3$  are necessary to promote the silaboration at rt. The alternative 1,1-silaboration of ethylene to furnish **12** would have to proceed *via* the much higher-energy transition state **TS2** ( $\Delta G^\ddagger =$





Fig. 3 Computed reaction mechanisms for (a) the observed 1,2-silaboration of ethylene with  $2 \cdot \text{SMe}_2/0.1 \text{ BI}_3$  (red) vs. the not observed 1,1-silaboration pathway (black) and (b) the observed 1,1-silaboration of cyclohexene with  $2$  (red) vs. the not observed 1,2-silaboration pathway (black). Color code: H: white, B: green, C: grey, Si: blue, I: violet. The Gibbs free energy changes ( $\Delta G$ ) were computed at the SMD(DCM)/PBE0-D3(BJ)/def2-QZVPPD level of theory, using geometries optimized at the SMD(DCM)/PBE0-D3(BJ)/def2-SVPD level. Note: compounds 9–11 appear in the SI as part of theoretically examined but energetically unfavorable alternative mechanisms.

25.2 kcal mol<sup>-1</sup> relative to the starting materials) and is thus not observed.

Due to the modified protocol used for the silaboration of cyclohexene, dissociation of  $\text{SMe}_2$  is not an issue here. Instead, free  $2$  can directly interact with the added olefin. Formation of the primary olefin complex  $2 \cdot (\text{C}_6\text{H}_{10})$  is somewhat more endergonic than in the case of ethylene ( $\Delta G = 11.3$  vs. 5.4 kcal mol<sup>-1</sup>), which can be attributed partly to steric factors and partly to a more pronounced reorganization energy: while the B atom in  $2 \cdot (\text{C}_6\text{H}_{10})$  is comparably pyramidalized as in  $2 \cdot (\text{C}_2\text{H}_4)$ , one B-bonded C atom now also deviates significantly from planarity [ $\sum(\text{C}-\text{C}-\text{C}/\text{H}) = 348.2^\circ$ ].<sup>127</sup> Starting from  $2 \cdot (\text{C}_6\text{H}_{10})$ , two subsequent transition states are most relevant:  $\text{TS1}'$  leads, *via* an overall activation barrier of 27.6 kcal mol<sup>-1</sup>, to the (experimentally unobserved) *syn*-1,2-silaboration product  $13$  ( $\Delta G = -7.5$  kcal mol<sup>-1</sup>). In contrast,  $\text{TS2}'$ , which lies 12.4 kcal mol<sup>-1</sup> lower in energy than  $\text{TS1}'$ , corresponds to a 1,2-

hydride shift leading to intermediate  $\text{Int1}$ . Subsequent 1,2-silyl migration *via* the low-lying  $\text{TS3}'$  furnishes the experimentally obtained 1,1-silaboration product  $7$ , with an overall reaction energy of  $\Delta G = -10.9$  kcal mol<sup>-1</sup>.

The differing regioselectivities observed in the silaborations of ethylene and cyclohexene arise as early as in intermediates  $2 \cdot (\text{C}_2\text{H}_4)$  and  $2 \cdot (\text{C}_6\text{H}_{10})$ : In  $2 \cdot (\text{C}_2\text{H}_4)$ , the ethylene coordination is near symmetric with B–C distances of 1.837 and 1.868 Å; the C(2)–C(1)–B–Si torsion angle is 23.0°, which represents an ideal conformation for an ensuing 1,2-silyl shift (Fig. 4a, left). According to a Natural Bond Orbital (NBO) analysis,<sup>128</sup> all three atoms—B, C(1), and C(2)—carry negative partial charges of -0.52, -0.43, and -0.49 e, respectively. Intermediate  $2 \cdot (\text{C}_2\text{H}_4)$  can thus be described as a  $\sigma$ -type donor–acceptor complex, in which charge is transferred from the occupied  $\pi$ -orbital of the olefin to the vacant orbital at the B atom, resulting in a substantial interaction energy of -313 kcal mol<sup>-1</sup>.<sup>129</sup> Notably,





Fig. 4 NBO and IBO analyses rationalizing the divergent silaboration pathways of ethylene and cyclohexene.<sup>129</sup> Color code: H: white, B: green, C: grey, Si: blue, I: violet. (a) Left: optimized structures of 2·(C<sub>2</sub>H<sub>4</sub>) (top) and 2·(C<sub>6</sub>H<sub>10</sub>) (bottom) with selected NBO charges given in elementary charges (e); right: key orbital interactions with associated stabilization energies; top right: IBO representation of the two-electron–three-center (2e–3c) bond in 2·(C<sub>2</sub>H<sub>4</sub>). (b) Left: Optimized structure of **Int1** with selected NBO charges given in elementary charges (e); right: hyperconjugative interactions stabilizing the carbenium ion, including the β-silicon effect and C–H<sub>ax</sub> σ donation; SMD(DCM)/PBE0-D3(BJ)/def2-SVPD level of theory.

an Intrinsic Bond Orbital (IBO)<sup>130</sup> analysis even suggests the presence of a C–B–C two-electron–three-center (2e–3c) bond, with relative contributions of 29.4% (B), 35.4% (C(1)), and 34.7% (C(2)); Fig. 4a, right). In 2·(C<sub>6</sub>H<sub>10</sub>), olefin binding is markedly unsymmetric, likely due to the higher steric bulk of cyclohexene relative to ethylene (Fig. 4a, left):<sup>131</sup> a short σ bond is found between the B center and the pyramidalized C(1) atom (1.829 Å), while the distance to the still planar C(2) atom is significantly longer (B···C = 2.356 Å). Concomitantly, the torsion angle C(2)–C(1)–B–Si is increased to 52.0°, thereby disfavoring a 1,2-silyl shift due to the longer Si···C(2) distance that would have to be traversed in the corresponding transition state. While the NBO charges on B and C(1) in 2·(C<sub>6</sub>H<sub>10</sub>) remain comparably negative to those in 2·(C<sub>2</sub>H<sub>4</sub>), C(2) now carries a positive charge of +0.09e. Cyclohexene can accommodate the steric constraints, as the carbenium ion at C(2) is stabilized by

both the +I effect of the alkyl substituent and hyperconjugative interactions<sup>132–134</sup> between its vacant p<sub>z</sub> orbital and the neighboring B–C and C–H<sub>ax</sub> σ bonds with contributions worth –92.7 and –19.5 kcal mol<sup>–1</sup> (ref. 129; Fig. 4a, right; see section 6.3.6 in the SI for a comparison with 3·(C<sub>6</sub>H<sub>10</sub>) where an NBO analysis reveals that **3** is coordinated primarily through a conventional, symmetric π → B interaction, most likely reflecting the lower Lewis acidity of **3**). Similar hyperconjugative interactions as described for 2·(C<sub>6</sub>H<sub>10</sub>) are also present in the rearrangement intermediate **Int1**—this time between the carbenium ion's p<sub>z</sub> orbital and the B–Si σ bond or two equivalent C–H<sub>ax</sub> σ bonds (relative energy contributions: –41.4 and 2 × –29.1 kcal mol<sup>–1</sup>, respectively; Fig. 4b, right). The former interaction corresponds to the well-known β-effect of a silyl group.<sup>125</sup> The overall stabilizing influence of steric and electronic factors makes **Int1** thermodynamically more favorable than 2·(C<sub>6</sub>H<sub>10</sub>). In



summary, the distinct regioselectivities in ethylene and cyclohexene silaboration originate from substrate-dependent binding geometries to **2**: symmetric coordination of ethylene facilitates direct 1,2-silaboration, whereas the unsymmetric activation of cyclohexene favors a stepwise 1,1-pathway *via* a stabilized carbenium-ion intermediate. The computed energy profiles and bonding analyses offer a coherent explanation for the experimentally observed selectivities and underscore the critical influence of steric and electronic substrate effects in directing the specific silaboration pathway.

## Conclusions

The addition of a reactant X–Y across a C=C double bond is a perfectly atom-economic transformation. When employing versatile (orthogonal) functional groups for X and Y, the primary addition products can be made valuable platforms for a wide range of applications. This is particularly true for X–Y-type reactants featuring covalently bonded boryl and silyl groups: both substituents are not only among the most versatile handles for downstream functionalization, but also play key roles as property-defining units in organic functional materials. Consequently, there is a growing demand for the development of novel silaboration reactions and tailored silylborane reagents  $R_2B-SiR_3$ . We have now found a way to make perhalogenated derivatives ( $R = Cl, I$ ) readily accessible on a multigram scale—both as free Lewis acids (*e.g.*,  $Cl_2B-SiCl_3$ ) and as Lewis base adducts ( $Do \cdot R_2B-SiR_3$ ;  $Do = SME_2, Py, PPh_3, IDipp$ ). These developments create a versatile platform with the following key features: (i)  $Me_2S \cdot I_2B-SiI_3$  and the *in situ*-generated mixture  $Li[I_3B-SiI_3]/I_2B-SiI_3/LiI$  react directly with olefins in silaboration reactions without the need for a catalyst, which is virtually without precedent.<sup>135</sup> (ii) Cyclohexene undergoes a 1,1-addition reaction—so far unobserved not only for silaborations, but also for diboration and disilylation reactions. Combined experimental and quantum-chemical studies revealed that the steric demand of cyclohexene renders symmetrical coordination of the olefin to the B site unfavorable and instead promotes the formation of a zwitterionic  $B^\ominus(sp^3)-C(sp^3)-C^\oplus(sp^2)$  fragment as a key entry point for the 1,1-silaboration cascade. While such a motif is prohibitively high in energy for ethylene, the carbenium center in the zwitterionic cyclohexene intermediate is efficiently stabilized through a combination of positive inductive (+I) and hyperconjugative effects. (iii) The halide substituents on the introduced boryl and silyl units enable diverse late-stage derivatizations—an aspect of particular importance when these functional groups are not merely used for transmetallation purposes in C–C-coupling reactions, but are instead retained as property-defining elements in the final molecule. (iv) Bulk  $Cl_2B-SiCl_3$  can be distilled without decomposition. Considering that  $Si_2Cl_6$  has been successfully used for the gas-phase deposition of silicon thin films,<sup>136</sup> and  $B_2F_4$  for their boron doping,<sup>137,138</sup>  $Cl_2B-SiCl_3$  emerges as a promising single-source precursor for semiconductor fabrication. Taken together, these findings pave the way for the future utilization of perhalogenated silylboranes in both synthesis (i–iii) and materials science (iv).

## Author contributions

J. H. performed the experimental studies and characterized all new compounds. C. D. B. performed the quantum-chemical calculations. A. V. V. and E. P. performed the X-ray crystal structure analyses of all compounds. H.-W. L., F. F. and M. W. supervised the project. The manuscript was written by J. H., C. D. B. and M. W. and edited by all co-authors.

## Conflicts of interest

There are no conflicts to declare.

## Data availability

CCDC 2470869, 2470870, 2470871, 2470872, 2470873, 2470874, 2470875, 2470876, 2470877, 2470878, 2470879, 2470880, 2470881, 2470882, 2470883, 2470884, 2470885, 2470886, 2470887, 2470888, and 2470889, contain the supplementary crystallographic data for this paper.<sup>139a–u</sup>

The data supporting this article have been included as part of the SI. Supplementary information is available. See DOI: <https://doi.org/10.1039/d5sc06234a>.

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- 110 2-SMe<sub>2</sub> is dimorphic. To confirm the phase purity of a freshly prepared sample of 2-SMe<sub>2</sub>, measured powder diffraction data must be compared with simulated patterns of both polymorphs.
- 111 The B-Do bonds within each 2·Do/3·Do pair featuring the same ligand tend to be slightly shorter in 2·Do, *e.g.*, 2-SMe<sub>2</sub> (B-S =  $1.927[2]$  Å)<sup>a</sup> vs. 3-SMe<sub>2</sub> (B-S =  $1.959(2)$  Å). Pyramidalization at boron was assessed by comparing the sum of the three bond angles in each  $SiBX_2$  fragment to that of three equivalent angles in an ideal tetrahedron ( $\Sigma = 328.5^\circ$ ). Almost all B sites in 2·Do/3·Do show equal or even greater pyramidalization [exceptions:  $\beta$ -2-SMe<sub>2</sub> ( $\Sigma = 330.3(6)^\circ$ ) and 3-SMe<sub>2</sub> ( $\Sigma = 333.1(4)^\circ$ )]. Adducts 2·Do are generally more pyramidalized than 3·Do, with the strongest donor IDipp producing the most pronounced effect: 2-IDipp ( $\Sigma = 308.6(5)^\circ$ ), 3-IDipp ( $\Sigma = 315.1[4]^\circ$ ).<sup>b</sup> (a) The average value with standard deviations in square brackets was calculated from the following individual bond lengths: B-S =  $1.925(6)$ ,  $1.927(6)$  ( $\alpha$ -polymorph; two crystallographically unique molecules);  $1.930(5)$  Å ( $\beta$ -polymorph); (b) The average value with standard deviations in square brackets was calculated from the bond angles of three crystallographically unique molecules:  $\Sigma = 314.75(54)$ ,  $314.95(54)$ ,  $315.65(53)^\circ$ .
- 112 The Si(2)-I bond involving a bridging I atom is clearly longer ( $2.611(4)$  Å) than the terminal Si(2)-I bonds ( $2.428(4)/2.403(4)$  Å).



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