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# Facile preparation of organosilanes from benzylboronates and gem-diborylalkanes mediated by $\text{KO}^t\text{Bu}^\dagger$

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Methods to efficiently synthesize organosilanes are valuable in the fields of synthetic chemistry and materials science. During the past decades, boron conversion has become a generic and powerful approach for constructing carbon–carbon and other carbon–heteroatom bonds, but its potential application in forming carbon–silicon remains unexplored. Herein, we describe an alkoxide base-promoted deborylative silylation of benzylic organoboronates, geminal bis(boronates) or alkyltriboronates, allowing for straightforward access to synthetically valuable organosilanes. This selective deborylative methodology exhibits operational simplicity, broad substrate scope, excellent functional group compatibility and convenient scalability, providing an effective and complementary platform for the generation of diversified benzyl silanes and silylboronates. Detailed experimental results and calculated studies revealed an unusual mechanistic feature of this C–Si bond formation.

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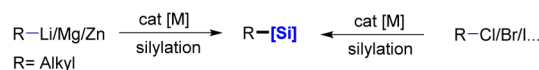
## Introduction

Organosilanes not only play pivotal roles in medicinal chemistry<sup>1</sup> and material science,<sup>2</sup> but they are also highly functionalized and versatile building blocks for further transformation in synthetic chemistry.<sup>3</sup> In the past few decades, considerable attention has been placed on developing synthetic strategies to construct C(sp<sup>3</sup>)–Si bonds by cross-coupling reactions in the presence of transition metals, such as palladium, nickel, and copper salts, as catalysts (Fig. 1A).<sup>4</sup> Highly reactive organometallic species, including Grignard reagents, organolithiums, organozincs or ganoaluminiums, are emerging as the most attractive substrates for the construction of organosilanes.<sup>5,6</sup> Although these transformations are synthetically useful, these organometallic reagents exhibit a high basicity and strong nucleophilicity, making them incompatible with many functional groups. Organohalides are often used as precursors to organometallic reagents; therefore, substantial progress has been recently achieved to access silicon-containing molecules by cross-coupling from these abundant and structurally diverse feedstocks to replace preformed organometallic reagents.<sup>7,8</sup> Despite these advances, a method to easily prepare

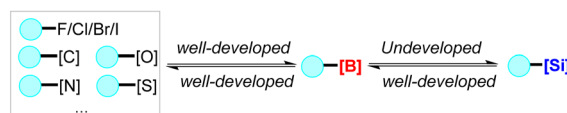
organosilanes without reactive coupling partners and metal catalysts is more attractive.

Interconversion of the main group elements has been a long-standing goal in organic chemistry.<sup>9</sup> In this area, organoboron compounds have become valuable building blocks in organic synthesis because of their flexibility, as they can be synthesized from multiple different functional groups.<sup>10</sup> Significant developments have been achieved for the interconversion of carbon–boron bonds to carbon–carbon and carbon–heteroatom (F, Cl, Br, I, O, N, S ...) bonds (Fig. 1B).<sup>11,12</sup> In this context, the conversion of C–Si bonds into C–B bonds has been developed through the reaction with boron trihalides (BCl<sub>3</sub>, BBr<sub>3</sub>).<sup>13</sup> At the

### (A) Reported routes to organosilanes



### (B) Interconversion of the main group elements



### (C) Opening the route from organoborons to silanes (this work)

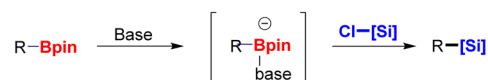


Fig. 1 Background and discovery. (A) Classic synthesis methods of organosilanes; (B) boron conversion; (C) deborylative silylation.

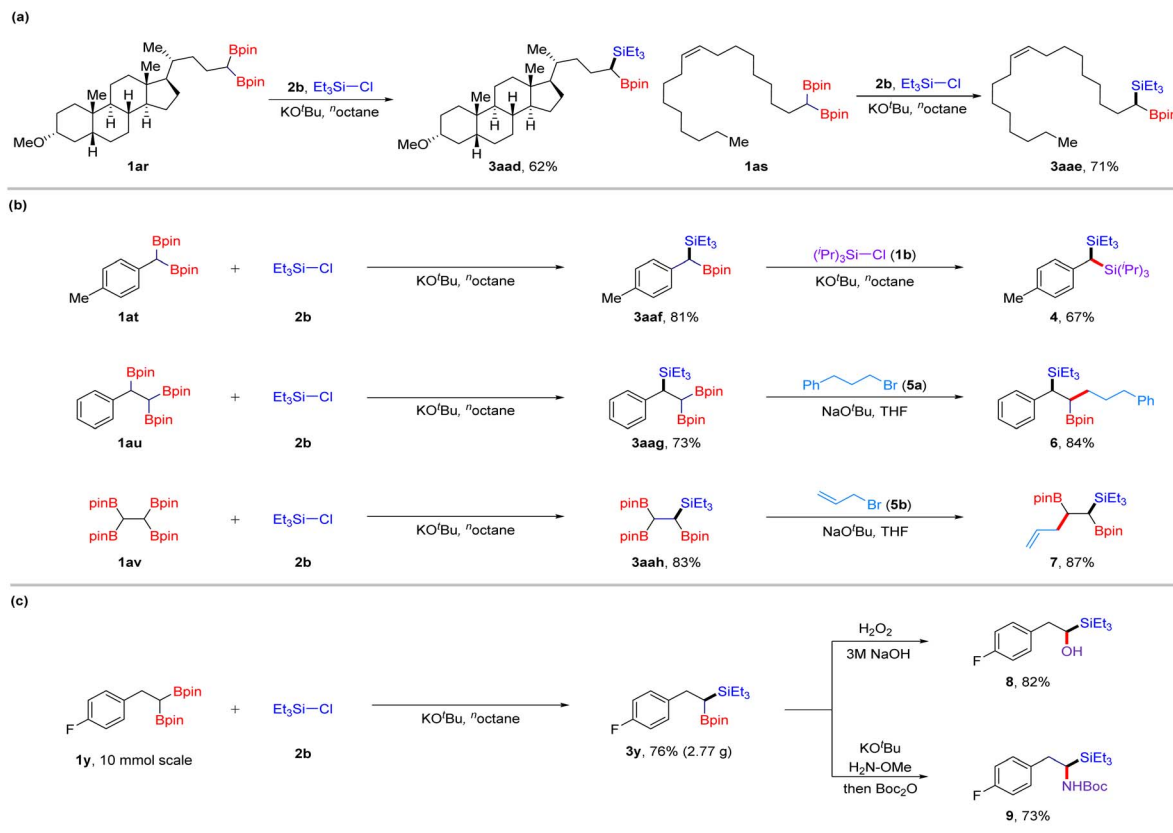
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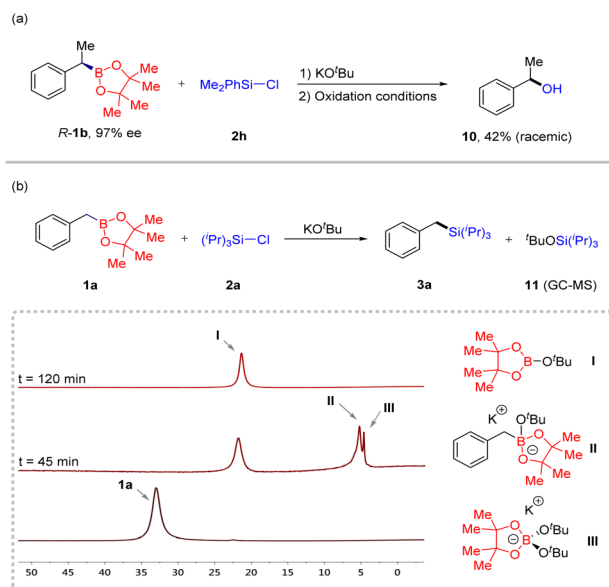
† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d3sc02461j>







Scheme 1 Synthetic applications of the deboronative silylation.



Scheme 2 Mechanistic experiments.

synthesis of geminal silylboronates,<sup>16</sup> which exhibit unique reactivity characteristics that allow them to participate in a variety of complexity-generating procedures. A range of carbon chain 1,1-bis(pinacolboronate) esters were tolerated under modified conditions, affording valuable silicated boronates in

70–79% yields (**3n–3t**). In addition, boronates **1u** underwent cross-coupling with chlorotriethylsilane to form highly congested  $\alpha$ -quaternary carbon centres **3u** in 77% yield. Although oxygen-containing substrates participated in this reaction, their yields were low (42–46%; **3v–3x**). Substrates possessing fluorine, chlorine, bromine, alkenyl and naphthyl groups were well tolerated in this system (**3y–3ad**). Delightedly, the common heterocyclic cores could favourably carry out these deboronative cross-coupling transformations (**3ae–3ah**). Selective functionalization of 1,2-bis(boronate esters) is a challenging topic in organic synthesis.<sup>17</sup> Therefore, we tried the selective cross-coupling reaction with these bis(boronate esters) and chlorosilanes under standard conditions. We found that this cross-coupling exclusively occurs at the benzyl boron, providing 1,2-silylboronates in moderate to good yields (**3ai–3ak**). Subsequently, 1,1,1-alkyltriboronates and 1,1,2-alkyltriboronates were selectively coupled with chlorosilanes to afford the corresponding products **3al–3ap** in 67–89% yield. Most of these silylboronate compounds are difficult to synthesize by known methods. Primary alkylboronates were also examined, but only trace amounts of silanes **3aq** were detected by GC-MS.

Next, we investigated the scope of chlorosilanes for cross-coupling with geminal boronates **1aa**. Substrates with different carbon chains were amenable to the cross-coupling reaction as well (**3ar–3au**). Notably, chlorosilane with a bulky substituent, such as a *tert*-butyl, showed lower yields (**3av**). The reaction of aryl chlorosilanes could afford target products **3aw**,



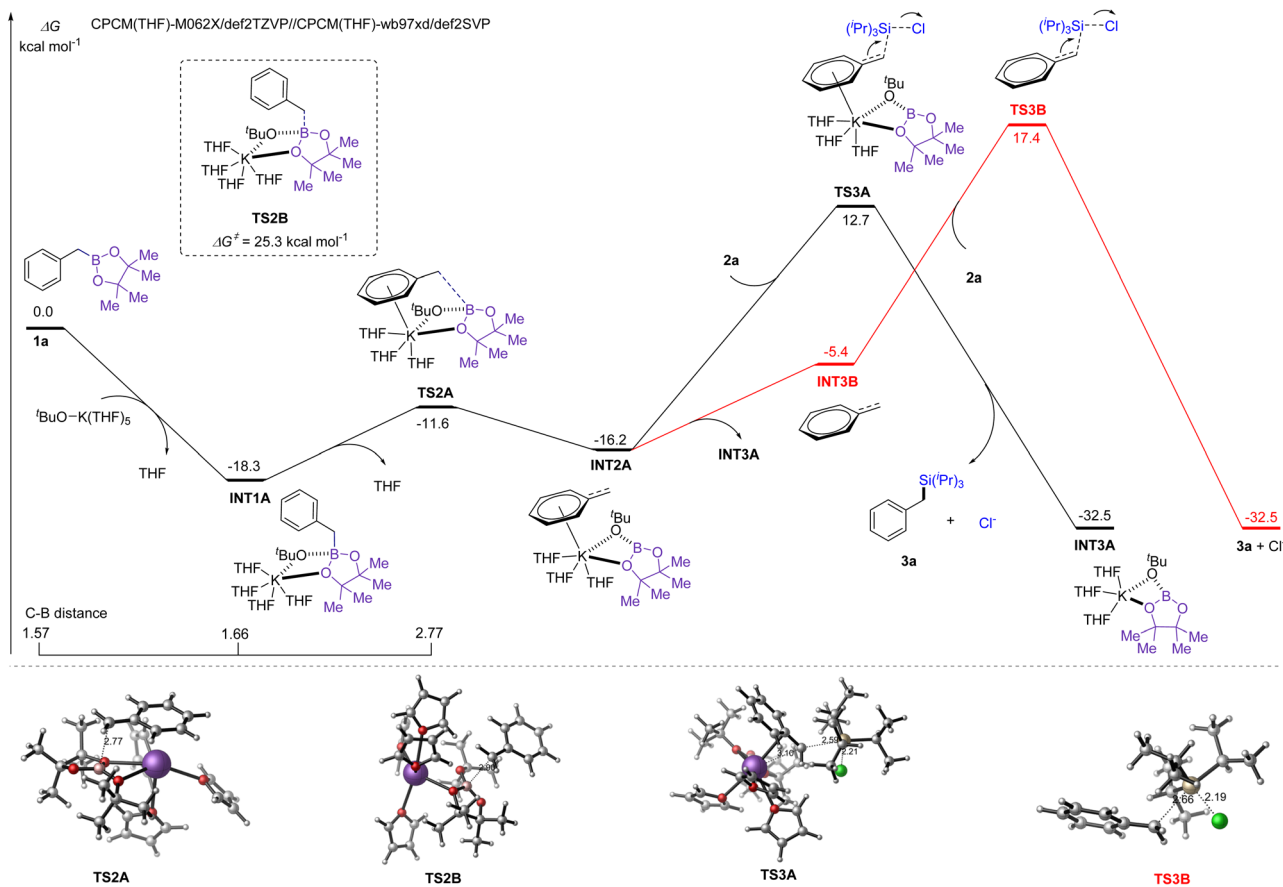


Fig. 2 The calculated energy profiles for the alkoxide-promoted cross-coupling of **1a** and **2a**.

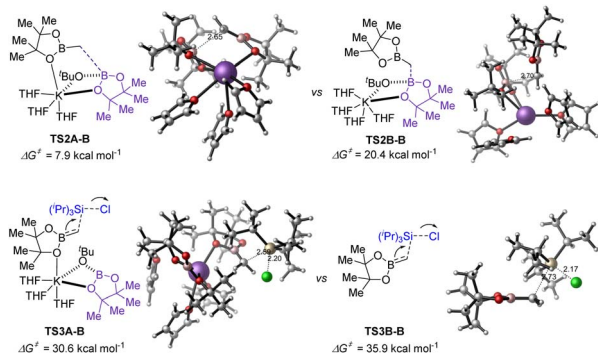


Fig. 3 The additional stabilization for substrate geminal boronates.

**3ax** and **3ay** in satisfactory yields as well. Chlorodisilanes were also feasible substrates with good yields (**3az**). Additionally, the use of a chloro-containing example afforded product **3aaa** in 53% yield. Finally, the presence of alkenyl substituents was tolerated (**3aab** and **3aac**).

To showcase the synthetic applications of this robust deboronative platform, we performed late-stage modification of complex molecules and a series of sequential transformations (Scheme 1). Lithocholic acid derivative **1ar** reacted with  $\text{Et}_3\text{SiCl}$  (**2b**) under optimized conditions, affording product **3aad** in 62% yield. Oleic acid derivative **1as** bearing a *trans*-alkene group

provided silylboronate **3aae** in 71% yield. Interestingly, by continuously treating **1at** with chlorosilanes and  $\text{KO}^t\text{Bu}$ , two deborylative silylations occurred, affording geminal disilanes **4** in 67% yield. When using 1,1,2-alkyltriboronate **1au** with two reactive sites as a substrate, the selective process of silylation was observed to give triethyl(1-phenyl-2,2-bis(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)ethyl) silane **3aag**, which can further undergo cross-coupling with alkyl bromide **5a** under the condition of sodium alkoxide ( $\text{NaO}^t\text{Bu}$ ),<sup>18</sup> providing 1,2-silylboronate **6** in 84% yield. Moreover, successive silylation and allylation led to the desired (1,2-bis(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)pent-4-en-1-yl)triethylsilane **7** in 87% yield. These results demonstrate the excellent chemoselectivity of this method. A gram-scale reaction proceeded with **1y**, affording a 76% isolated yield of silylated product **3y**, which can then undergo oxidation or amination processes to afford hydroxyl- or amino-products (**8**, **9**) in good yields, respectively.

To determine the reaction mechanism, we performed some experiments (Scheme 2). First, when *R*-**1b** (97% ee) was reacted with  $\text{Me}_2\text{PhSi-Cl}$  **2h** under our conditions, racemic compound **10** was observed in 42% yield after oxidation treatment. GC-MS was used to detect the course of the reaction, and we found that byproduct **11** formed. Subsequently, *in situ*  $^{11}\text{B}$  NMR of the reaction process was performed using benzyl boron ester **1a** and chlorosilane **2a** as substrates, and three new resonances (**I**, **II**, and **III**) appeared at 45 minutes. The resonance at 5.1 ppm (**II**) is



assigned to an “ate” complex.<sup>19</sup> Over time, this characteristic peak gradually disappeared. Based on these observations, we speculate that this transformation involves two processes. The first step was the generation of a carbanion through a deborylative pathway of an “ate” complex **II**. The second step was the nucleophilic reaction of the formed alkyl anion with chlorosilane to generate the final product.

The proposed mechanism was further supported by a detailed density functional theory (DFT) study, as shown in Fig. 2. Initially, the boron atom of **1a** is coordinated with the oxygen of KO<sup>t</sup>Bu solvated by five tetrahydrofuran molecules, and then the oxygen of **1a** replaces one molecule of THF at potassium to generate **INT1A** with an exothermic energy of 18.3 kcal mol<sup>-1</sup>. In **INT1A**, the boron centre is more electron-rich than that in **1a**, and the C–B distance is elongated to 1.66 Å from 1.57 Å, decreasing the bond dissociation energy of the C–B bond. Subsequent C–B bond cleavage proceeds smoothly through transition state **TS2A** with an energy barrier of 6.7 kcal mol<sup>-1</sup>, in which the potassium cation binds to the aromatic ring of **1a** to stabilize the developing negative charge centre and simultaneously release one molecule of THF. The direct cleavage of the C–B bond *via* transition state **TS2B** without the assistance of potassium requires a higher activation free energy (25.3 kcal mol<sup>-1</sup>). The negative charge centre in intermediate **INT2A** undergoes nucleophilic attack to chlorotriisopropylsilane **2a** *via* transition state **TS3A** with an activation free energy of 28.9 kcal mol<sup>-1</sup>, producing the desired alkyl silane **3a** and byproduct **INT3A**. The calculational results indicate that the nucleophilic attack step is the rate-determining step, and the overall activation free energy for this transformation is 31.0 kcal mol<sup>-1</sup>. The nucleophilic attack step for the alkyl anion released from **INT2A** must overcome an overall free energy of 35.7 kcal mol<sup>-1</sup> through transition state **TS3B**, which is 4.7 kcal mol<sup>-1</sup> higher than that of **TS3A**. This explains why the deborylative silylation can be promoted by KOMe but failed with sodium alkoxide or lithium alkoxide (Table 1, entries 3–5). Therefore, both the alkoxide anion and potassium cation are indispensable in this transformation, as these entities participate in the formation of a more stable intermediate species, lowering the energy barrier of C–B bond cleavage and C–Si bond formation.

For substrate germinal boronates, the pathways for C–B bond cleavage through transition **TS2A–B** and the subsequent C–Si bond formation *via* **TS3A–B** have the lower energies (Fig. 3). Comparing the energy barrier of transition state **TS3A–B** in the rate-determining step, the stepwise S<sub>N</sub>2 pathway through **TS3B–B** was found to be disfavoured kinetically and ruled out. Overall, DFT analysis further reveals that the transition states **TS2A–B** and **TS3A–B** are substantially stabilized by the coordination between the potassium cation and oxygen of the reserved boronate (see ESI† for details).

## Conclusions

In summary, we have established a general and practical method for synthesizing alkyl silane derivatives from readily available benzylic boronates, geminal bis(boronates) or

alkyltriboronates. This transition-metal-free transformation features excellent chemoselectivity, broad substrate scope, versatility, and scalability. These borylated silanes can readily be further functionalized by well-established organoboron chemistry to enhance the molecular complexity. In view of these features, this transformation should have high synthetic value in the field of materials and pharmaceuticals.

## Data availability

The synthetic procedures, characterization, and spectral data supporting this article have been uploaded as part of the ESI.†

## Author contributions

J. H. conceived the project and directed the research. S. J. and Z. S. supervised the mechanistic study. J. H. and Z. S. wrote the paper. M. T., W. Z., J. W. and H. S. performed the experiments. M. W. performed the DFT calculations.

## Conflicts of interest

The authors declare no competing financial interest.

## Acknowledgements

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## Notes and references

- (a) A. K. Franz and S. O. Wilson, *J. Med. Chem.*, 2013, **56**, 388–405; (b) R. Ramesh and D. S. Reddy, *J. Med. Chem.*, 2018, **61**, 3779–3798.
- (a) R. Pietschnig and S. Spirk, *Coord. Chem. Rev.*, 2016, **323**, 87–106; (b) E. A. Marro and R. S. Klausen, *Chem. Mater.*, 2019, **31**, 2202–2211.
- (a) F. Zhao, S. Zhang and Z. Xi, *Chem. Commun.*, 2011, **47**, 4348–4357; (b) S. E. Denmark and A. Ambrosi, *Org. Process Res. Dev.*, 2015, **19**, 982–994; (c) T. Komiyama, Y. Minami and T. Hiyama, *ACS Catal.*, 2017, **7**, 631–651.
- S. Bähr, W. Xue and M. Oestreich, *ACS Catal.*, 2019, **9**, 16–24.
- (a) M. Lacout-Loustalet, J. Dupin, F. Metras and J. Valade, *J. Organomet. Chem.*, 1971, **31**, 187–204; (b) P. J. Lennon, D. P. Mack and Q. E. Thompson, *Organometallics*, 1989, **8**, 1121–1122; (c) A. P. Cinderella, B. Vulovic and D. A. Watson, *J. Am. Chem. Soc.*, 2017, **139**, 7741–7744; (d) B. Vulovic, A. P. Cinderella and D. A. Watson, *ACS Catal.*, 2017, **7**, 8113–8117; (e) S. Mallick, E.-U. Würthwein and A. Studer, *Org. Lett.*, 2020, **22**, 6568–6572.
- (a) K. Tamao, A. Kawachi and K. Ito, *J. Am. Chem. Soc.*, 1992, **114**, 3989–3990; (b) K. Tamao and A. Kawachi, *Adv. Organomet. Chem.*, 1995, **38**, 1–58; (c) C. K. Chu, Y. Liang



- and G. C. Fu, *J. Am. Chem. Soc.*, 2016, **138**, 6404–6407; (d) W. Xue, R. Shishido and M. Oestreich, *Angew. Chem., Int. Ed.*, 2018, **57**, 12141–12145; (e) R. Chandrasekaran, F. T. Pulikkottil and K. S. Elama, *Chem. Sci.*, 2021, **12**, 15719–15726.
- 7 (a) W. Xue, Z.-W. Qu, S. Grimme and M. Oestreich, *J. Am. Chem. Soc.*, 2016, **138**, 14222–14225; (b) W. Xue and M. Oestreich, *Angew. Chem., Int. Ed.*, 2017, **56**, 11649–11652; (c) Y. Takeda, K. Shibuta, S. Aoki, N. Tohnai and S. Minakata, *Chem. Sci.*, 2019, **10**, 8642–8647; (d) J. Scharfbier, B. M. Gross and M. Oestreich, *Angew. Chem., Int. Ed.*, 2020, **59**, 1577–1580.
- 8 (a) J. Duan, K. Wang, G.-L. Xu, S. Kang, L. Qi, X.-Y. Liu and X.-Z. Shu, *Angew. Chem., Int. Ed.*, 2020, **59**, 23083–23088; (b) L. Zhang and M. Oestreich, *Angew. Chem., Int. Ed.*, 2021, **60**, 18587–18590.
- 9 (a) D. A. Petrone, J. Ye and M. Lautens, *Chem. Rev.*, 2016, **116**, 8003–8104; (b) K. Ouyang, W. Hao, W.-X. Zhang and Z. Xi, *Chem. Rev.*, 2015, **115**, 12045–12090; (c) L.-J. Cheng and N. P. Mankad, *Chem. Soc. Rev.*, 2020, **49**, 8036–8064; (d) R. Sharma and M. R. Yadav, *Org. Biomol. Chem.*, 2021, **19**, 5476–5500; (e) F. Mo, D. Qiu, L. Zhang and J. Wang, *Chem. Rev.*, 2021, **121**, 5741–5829; (f) A. Varenikov, E. Shapiro and M. Gandelman, *Chem. Rev.*, 2021, **121**, 412–484; (g) S. Huang, M. Wang and X. Jiang, *Chem. Soc. Rev.*, 2022, **51**, 8351–8377.
- 10 (a) A. J. L. Lennox and G. C. Lloyd-Jones, *Chem. Soc. Rev.*, 2014, **43**, 412–443; (b) A. Fawcett, J. Pradeilles, Y. Wang, T. Mutsuga, E. L. Myers and V. K. Aggarwal, *Science*, 2017, **357**, 283–286; (c) J. Hu, Y. Zhao and Z. Shi, *Nat. Catal.*, 2018, **1**, 860–869; (d) X. Liu, W. Ming, Y. Zhang, A. Friedrich and T. B. Marder, *Angew. Chem., Int. Ed.*, 2019, **58**, 18923–18927; (e) J. Li, H. Wang, Z. Qiu, C. Y. Huang and C.-J. Li, *J. Am. Chem. Soc.*, 2020, **142**, 13011–13020.
- 11 From boron to other main group elements: (a) C. M. Crudden, B. W. Glasspoole and C. J. Lata, *Chem. Commun.*, 2009, 6704–6716; (b) D. Leonori and V. K. Aggarwal, *Angew. Chem., Int. Ed.*, 2015, **54**, 1082–1096; (c) T. C. Wilson, T. Cailly and V. Gouverneur, *Chem. Soc. Rev.*, 2018, **47**, 6990–7005.
- 12 From other main group elements to boron: (a) E. C. Neeve, S. J. Geier, I. A. I. Mkhaliid, S. A. Westcott and T. B. Marder, *Chem. Rev.*, 2016, **116**, 9091–9161; (b) M. Wang and Z. Shi, *Chem. Rev.*, 2020, **120**, 7348–7398; (c) J. Hu, M. Ferger, Z. Shi and T. B. Marder, *Chem. Soc. Rev.*, 2021, **50**, 13129–13188.
- 13 (a) W. Haubold, J. Herdtle, W. Gollinger and W. Einholz, *J. Organomet. Chem.*, 1986, **315**, 1–8; (b) A. A. Kolomeitsev, A. A. Kadyrov, J. Szczepkowska-Sztolcman, M. Milewska, H. Koroniak, G. Bissky, J. A. Bartene and G. Rösenthaller, *Tetrahedron Lett.*, 2003, **44**, 8273–8277; (c) D. L. Crossley, J. Cid, L. D. Curless, M. L. Turner and M. J. Ingleson, *Organometallics*, 2015, **34**, 5767–5774; (d) A. Lik, L. Fritze, L. Müller and H. Helten, *J. Am. Chem. Soc.*, 2017, **139**, 5692–5695; (e) C. You, M. Sakai, C. G. Daniliuc, K. Bergander, S. Yamaguchi and A. Studer, *Angew. Chem., Int. Ed.*, 2021, **60**, 21697–21701; (f) J. Yao, L. Yu, W. Duan and C.-J. Li, *Org. Chem. Front.*, 2023, **10**, 524–530.
- 14 C. Cheng and J. F. Hartwig, *Chem. Rev.*, 2015, **115**, 8946–8975.
- 15 A. M. Muzafarov, *Silicon Polymers*, Springer, Heidelberg, 2011.
- 16 (a) A. A. Szymaniak, C. Zhang, J. R. Coombs and J. P. Morken, *ACS Catal.*, 2018, **8**, 2897–2901; (b) Z. Cheng, J. Guo, Y. Sun, Y. Zheng, Z. Zhou and Z. Lu, *Angew. Chem., Int. Ed.*, 2021, **60**, 22454–22460; (c) Y. You and S. Ge, *Angew. Chem., Int. Ed.*, 2021, **60**, 20684–20688; (d) S. Jin, K. Liu, S. Wang and Q. Song, *J. Am. Chem. Soc.*, 2021, **143**, 13124–13134; (e) W. Sun, L. Xu, Y. Qin and C. Liu, *Nat. Synth.*, 2023, **2**, 413–422.
- 17 (a) S. N. Mlynarski, C. H. Schuster and J. P. Morken, *Nature*, 2014, **505**, 386–390; (b) R. D. Dewhurst and T. B. Marder, *Nat. Chem.*, 2014, **6**, 279–280; (c) L. Xu, S. Zhang and P. Li, *Chem. Soc. Rev.*, 2015, **44**, 8848–8858.
- 18 K. Hong, X. Liu and J. P. Morken, *J. Am. Chem. Soc.*, 2014, **136**, 10581–10584.
- 19 B. Lee and P. J. Chirik, *J. Am. Chem. Soc.*, 2020, **142**, 2429–2437.

