

Cite this: *Chem. Sci.*, 2021, **12**, 14635

All publication charges for this article have been paid for by the Royal Society of Chemistry

Received 15th September 2021
Accepted 18th October 2021

DOI: 10.1039/d1sc05125c
rsc.li/chemical-science

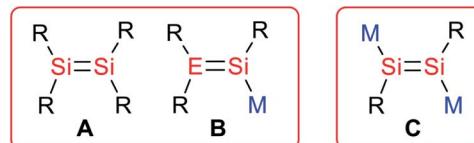
Introduction

There is great interest in the synthesis and reactivity of heavier main group multiply bonded species due to their unusual electronic structures and reaction patterns.¹ A number of alkene analogues of silicon kinetically stabilized by bulky alkyl, aryl and silyl substituents have been isolated (A, Scheme 1).² In light of the diverse bonding modes in disilenes, the corresponding anionic systems are expected to be useful as synthetic reagents for novel disilenes that otherwise could not be obtained.³ In addition, the anionic multiple bonding systems may exhibit cooperative effects in the activation of small molecules. However, anionic multiply bonded silicon species are much less common because of their synthetic challenges and extremely high reactivity. Nevertheless, several silicon analogues of vinyl anions (B, Scheme 1) that have been reported and their synthetic applications for new types of silicon species and unique reaction patterns to small molecules have been demonstrated.^{2a,b,3,4} It is envisioned that 1,2-alkali-metal-substituted disilenes (C, Scheme 1) could be versatile reagents for access to diverse silicon multiply bonded species and activation of inert molecules.

State Key Laboratory of Elemento-organic Chemistry and College of Chemistry, Nankai University, Tianjin 300071, China. E-mail: cmcui@nankai.edu.cn

† Electronic supplementary information (ESI) available. CCDC 2081430, 2081431, 2088661, 2088665 and 2104803. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/d1sc05125c

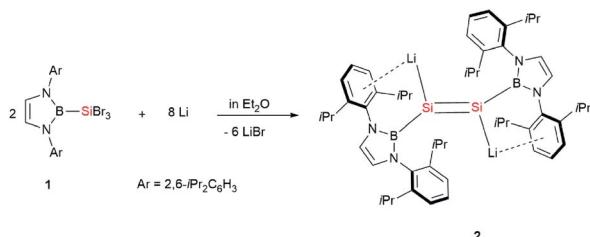
1,2-Dialkali-metal substituted disilenes are also inherently related to disilynes, the silicon analogues of alkynes, because they could be viewed as doubly reduced disilynes. Disilynes have been of considerable interest because of their fundamental importance. By the employment of very bulky terphenyl, alkyl and silyl substituents, a few examples of disilynes have been isolated.⁵ However, their doubly reduced species have not been reported so far. It has been shown that the reduction of a silyl-substituted disilyne yielded the corresponding radical anion, and no further reduction products were disclosed.⁶ It is noted that the doubly reduced alkyne analogues have been isolated for germanium and tin by Power and Aldridge *et al.*⁷ However, to the best of our knowledge 1,2-alkali metal substituted disilenes (disilyne dianions) are still elusive. Very recently, our group reported the synthesis of a disilene and 1-magnesium-2,3-disilacyclop propane stabilized by a bulky boryl substituent.⁸ The strong σ electronic effects of the boryl ligand



R = bulky silyl, aryl, alkyl; E = C, Si, Ge;
M = alkali metal, Zr, Cu, Zn, Mg; boryl = (HCArN)₂B

Scheme 1 Structural motifs of the reported disilenes (A), silicon analogues of vinyl anions (B) and this work (C).





Scheme 2 Synthesis of 2.

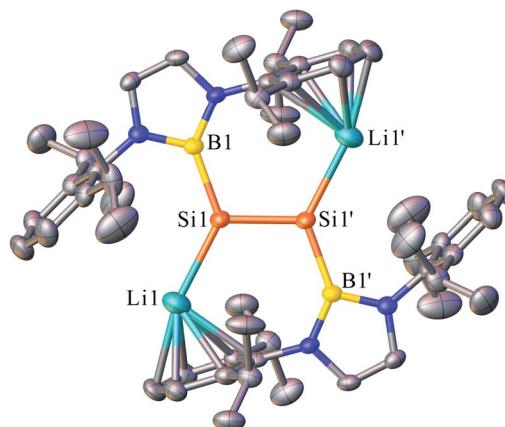
prompted us to investigate the synthetic probability of 1,2-dialkali-metal substituted disilenes.

Herein, we report the first synthesis of a dialkali metal-substituted disilene, the dilithio-diboryl-disilene **2** (Scheme 2), and its reactivity to haloboranes. Remarkably, in contrast to the *trans*-bent geometry observed for most of the disilenes, compound **2** adopts an almost planar geometry. Furthermore, a novel 2π aromatic B-Si-Si three-membered ring system with a 2e-3c bond and the first tetraboryldisilene was obtained from the reaction of **2** with haloboranes.

Results and discussion

The dilithiodisilene $[(\text{boryl})\text{SiLi}]_2$ (**2**) was prepared by the reduction of **1** (ref. 7) with four equivalents of lithium powder in diethyl ether and was isolated as a red powder in 73% yield (Scheme 2). The structure of **2** has been characterized by multinuclear NMR spectroscopy, UV-vis spectroscopy and X-ray single-crystal analysis. The ^{29}Si NMR spectrum of **2** in C_6D_6 shows a resonance at 297.0 ppm, which is down-field shifted in comparison to that reported for 1-magnesium-2,3-disilacyclop propane (204.1 ppm).^{8b} The UV-vis spectrum of **2** in toluene shows the longest wavelength absorption maximum at 450.0 nm, which is slightly red-shifted compared with the maximum reported for 1-magnesium-2,3-disilacyclop propane (431.0 nm) (Fig. S2†).^{8b} This value is very close to the value calculated for the $\pi-\pi^*$ transition of **2** (508.9 nm; HOMO-1 → LUMO+4, $f = 0.2785$) (Fig. S3†). The dilithiodisilene **2** is slightly soluble in *n*-hexane but is well dissolved in benzene and toluene. It is noted that compound **2** is not stable in THF and quickly decomposes into a complicated mixture.

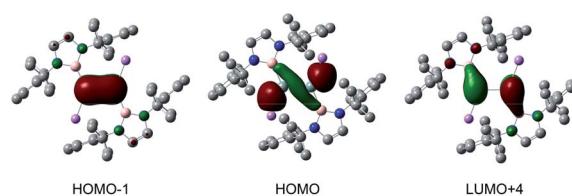
Dark red crystals of **2** suitable for X-ray diffraction studies were obtained from *n*-hexane at 0 °C. There are two independent molecules in the unit cell, and each of them is crystallographically C_2 symmetric. The geometries of the two molecules are similar and thus only one is shown in Fig. 1. The structural analysis of **2** indicated an almost planar Si-Si geometry as indicated by the sum of the bond angles of 359.96° around the Si1 and Si1' atoms and the six almost coplanar atoms B1, Li1, Si1, Si1', Li1', B1' (the largest out-of-plane deviation = 0.06 Å). This is in contrast to the nonplanar, *trans*-bent geometry of the majority of known disilenes. The planarity of **2** could be attributed to the electronic effects of the electropositive boryl and lithium substituents.⁹ The Si-Si bond length of 2.223(4) Å is same as that reported for 1-magnesium-2,3-disilacyclop propane (2.223(17) Å), while the Si-Li bond length of 2.482(12) Å is

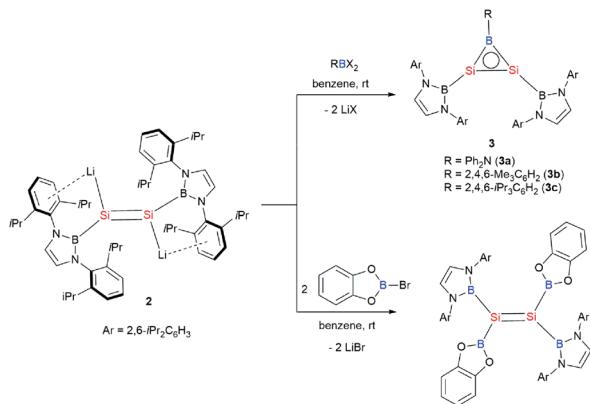
Fig. 1 OLEX2 drawing of **2** with 50% ellipsoid probability. Hydrogen atoms have been omitted for clarity.

remarkably shorter than those observed in contact ion pairs of silenyl lithium species.^{4,10} The two five-membered diazaborolyl $\text{C}_2\text{N}_2\text{B}$ planes are essentially coplanar with the BSiSi plane (the dihedral angle = 1.83°). These structural features indicate that the dilithio-diboryldisilene **2** features an ethylene-like structure.

To get insight into the electronic structure and bonding situations, density functional theory (DFT) calculations were performed for **2** at the B3LYP/6-31G(d,p) level (Fig. S4†). The calculated geometry is very close to that experimentally determined for **2**. Natural bond orbital (NBO) analysis revealed highly filled σ (1.97e) and π (1.81e) bonds for the Si=Si double bonds. The Wiberg bond index (WBI, Table S2†) of 1.77 for the Si-Si bond is consistent with that for the double bond. Natural population analysis (NPA) charges (Si: -0.58; Li: +0.79) of **2** indicate the pronounced anionic character of the silicon atoms. The Si=Si π bond orbital, which is perpendicular to the B-Si-Si-B plane, can be seen in the HOMO-1 (Fig. 2). The HOMO of **2** is mainly composed of the silicon lone pair orbitals. The LUMO+4 consists predominantly of the Si-Si π^* bond with some contribution of the boron-centered p-orbitals.

Alkali metal-substituted disilenes may facilitate extensive studies on their reactions with halogenated reagents for the development of new reaction modes and access to novel multiply bonded silicon compounds.^{2a,b,3,4} Reduction of appropriately disubstituted dihalosilanes ($\text{R}_1\text{R}_2\text{SiX}_2$) is the most general route to disilenes.² However, in contrast to easy availability of alkyl, aryl and silyldihalosilanes, boryl-substituted dihalosilanes could not be easily prepared because of the lack of nucleophilic boron compounds. In view of the interesting

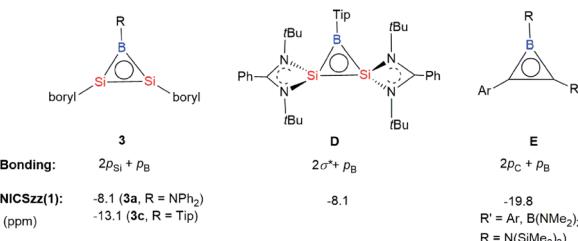
Fig. 2 The calculated frontier orbitals of **2**.



Scheme 3 Reactivity of 2.

electronic effects of boryl substituents, we explored the reaction of dilitihiodesilene 2 with several types of haloboranes.

First, the reactions of 2 with dihaloboranes were explored with the expectation to generate a Si–Si–B three-membered ring system as it is isoelectronic with aromatic cyclopropenylum ions ($[\text{C}_3\text{R}_3]^+$) and borirenes ($\text{R}_2\text{C}_2\text{BR}$). Thus, reactions of dilitihiodesilene 2 with dihaloboranes RBX_2 ($\text{R} = \text{Ph}_2\text{N}$, Mes, Tip; $\text{X} = \text{Cl}, \text{Br}$; Mes = $2,4,6\text{-Me}_3\text{C}_6\text{H}_2$; Tip = $2,4,6\text{-iPr}_3\text{C}_6\text{H}_2$) have been conducted. Treatment of 2 with the haloboranes in benzene at room temperature afforded an orange yellow solution, from which disilaborirenes 3a–c could be obtained as light orange solids in 53%, 86% and 68% yields, respectively (Scheme 3). X-ray single-crystal analysis revealed that compounds 3a–c feature a planar B–Si–Si three-membered ring with both silicon and boron atoms being three-coordinate. The ^{29}Si NMR spectra of compounds 3a–c exhibit a singlet signal at 112.2, 142.0 and 141.1 ppm, respectively. In the ^{11}B NMR spectra of 3a–c, two broad signals are observed (δ 24.2 and 50.1 ppm for 3a, δ 23.5 and 72.9 ppm for 3b, and δ 23.4 and 73.3 ppm for 3c). The UV-vis spectra of compounds 3a–c in toluene show the longest wavelength absorption maximum at 380.0, 364.5 and 370.0 nm, respectively (Fig. S5†). The NMR and UV-vis data indicate that the amino substituent Ph_2N on the boron atom in 3a has pronounced effects on the electronic structure of the B–Si–Si



Scheme 4 Comparison of the bonding and NICS values of disilaborirenes (3 and D) and borirenes (E).

three-membered ring because of the π -interactions of the amino group with the boron atom.

Orange yellow crystals of 3a–c suitable for X-ray diffraction studies were obtained from *n*-pentane at $-20\text{ }^\circ\text{C}$. The molecular structural analysis of 3a and 3c shown in Fig. 3 (for 3a–c, see Fig. S6–S8†) confirmed the B–Si–Si three-membered ring system. The Si–Si bond lengths of 2.1469(11), 2.133(2) and 2.1452(6) \AA in 3a–c are in the range of that of a Si–Si double bond (2.12–2.25 \AA)^{1e} and slightly shorter than that determined for a cyclotrisilene cation (av. 2.217(3) \AA).¹¹ The B–Si bond lengths in the 3a–c BSi_2 cycles ranging from 1.911(7) \AA to 1.952(3) \AA are close to those of 1.9190 and 1.9186(13) \AA reported for $\text{L}_2\text{Si}_2\text{BR}$ (D, Scheme 4) featuring the four-coordinate silicon atoms.¹² The sums of the bond angles around Si1 and Si2 in 3a–c are close to 360° , indicating that the geometries of each silicon atom are essentially planar. A comparison of the B–Si bonds and internal angles of the Si_2B ring in 3a–c reveals that the amino substituent Ph_2N in 3a engages in a relatively weak π -interaction with the empty p orbital of the boron atom.

As borirenes are aromatic systems, we are interested in the aromaticity of the silicon analogues 3a–c for comparison.¹³ Thus, DFT calculations were carried out for amino-substituted 3a and the phenyl-substituted 3c at the B3LYP/6-31G(d,p) level. The optimized structures (Fig. S9 and S10†) are well correlated with the experimentally determined ones. The WBI values (1.25 and 1.28 for 3a and 3c) of the ring B–Si bonds indicate the partial double-bonds caused by 3c–2e bonding of the BSi_2 unit in 3a (1.96e) and 3c (1.95e), as shown by NBO analysis (Tables S3–S6†). The calculated frontier orbitals for 3a

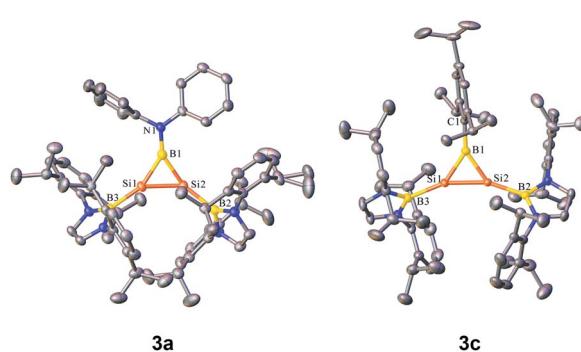


Fig. 3 OLEX2 drawing of 3a and 3c with 50% ellipsoid probability. Hydrogen atoms have been omitted for clarity.

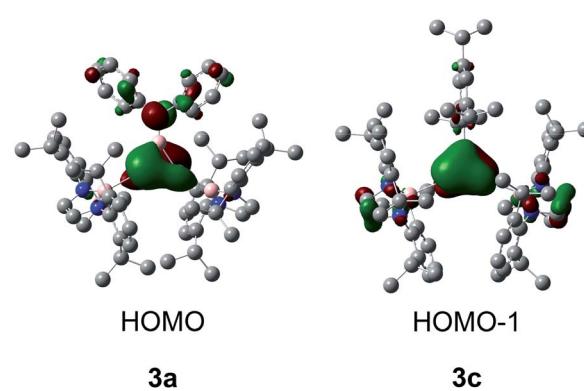


Fig. 4 Frontier orbitals of 3a (left) and 3c (right).



and **3c** are shown in Fig. 4. The HOMO of **3a** consists of the Si=Si π bonding and antiphase nitrogen lone pair, while the HOMO – 1 of **3c** mainly corresponds to the delocalized Si₂B π orbital (Fig. 4).

To determine the π aromaticity of disilaborirenes **3a** and **3c**, the nucleus independent chemical shift (NICS_{zz}(1)) values were calculated. The NICS values of –8.1 and –13.1 ppm calculated for **3a** and **3c** indicate the aromaticity, but it is weaker than that (–19.8 ppm) reported for borirene **E** (Scheme 4).¹³ Although the aromaticity of **3a** with R = Ph₂N is similar to that reported for **D** (Scheme 4, Tip = 2,4,6-iPr₃C₆H₂),¹² that of **3c** is much stronger than that of **D**. It is apparent that the degree of the π aromaticity of **3c** with R = Tip is much greater than that of **3a** with R = Ph₂N because of the electron delocalization in the B–N bond in **3a**. The aromaticity of **3a** and **3c** could be interpreted as the extensive 2 π electron delocalization over the three-center bonding molecular orbitals from the p orbitals of B and Si atoms (Scheme 4). The bonding situation in the Si₂B cycle is analogous to that of borirenes but distinct from that of **D** with four-coordinate Si atoms, in which the delocalized structure was formed by the σ^* MOs of π -symmetry with the boron p orbital.¹²

Reaction of dilithiodisilene **2** with two equivalents of *B*-bromocatecholborane (cat)BBr resulted in the formation of the first tetraboryldisilene **4** (Scheme 3). Although a great number of disilenes have been synthesized, disilenes substituted by group 13 and 15 elements are still limited because of the lack of appropriate synthetic precursors.^{2b,3c,g,m,n,14} As the structure of a disilene is largely dependent on the substituents, the synthesis of boryl-substituents is of current interest. Tetraboryldisilene **4** was fully characterized by NMR spectroscopy, UV-vis spectroscopy and X-ray single-crystal analysis. The ²⁹Si NMR spectrum shows a singlet at 105.6 ppm, which resides at a higher field in comparison to that of diboryldisilene (200.0 ppm). Consistent with the molecular structure of **4**, the ¹¹B NMR spectrum exhibits two broad signals at δ 26.2 and 37.3 ppm. The UV-vis spectra of **4** in toluene show the longest wavelength absorption maximum at 445.0 nm (Fig. S11†), which is same as the value calculated for the π – π^* transition of **4** (445.0 nm; HOMO → LUMO, f = 0.4433) (Fig. S12†).

Yellow crystals of **4** suitable for X-ray diffraction studies were obtained from *n*-hexane at –20 °C. Analysis of the structure of **4**

(Fig. 5 and S13†) revealed that the Si=Si bond adopts an essentially planar geometry with a twist angle of 0° as well as a *trans*-bent angle of 1.86°, indicating the pronounced electronic effects (σ -donor) of the boryl substituents on the Si=Si bond. The diazaborolyl C₂N₂B planes are almost orthogonal to the B–Si–Si planes with a dihedral angle of 86.58°. The structural features of **4** were also examined by DFT calculations at the B3LYP/6-31G(d,p) level. The HOMO and LUMO are mainly Si=Si π and π^* orbitals (Fig. S15†). On the basis of the results, compound **4** represents a planar disilene featuring the smallest *trans*-bent geometry and zero twist angle.¹⁵

Conclusions

In summary, dilithiodisilene **2** was prepared by the reduction of (boryl)SiBr₃ with lithium in Et₂O. Dilithiodisilene **2** represents one of the rare examples of ethylene-like disilenes. It turned out to be a valuable reagent for the construction of novel silicon multiply-bonded species as demonstrated by the first synthesis of a 3c-2 π aromatic system **3a–c** and a fully boryl-substituted disilene **4**. The unique electronic effects of the boryl group allowed facile access to the dianionic species **2**. DFT calculations predicted that the aromaticity of disilaborirenes **3a–c** stems from the delocalization of the 2 π electrons over the three-center bonding molecular orbital from the B and Si p atomic orbitals. Exploration of more challenging synthetic targets from dilithiodisilene **2** is currently underway in our laboratory.

Data availability

All experimental procedures, spectroscopic data, results of the DFT calculations and crystallographic data can be found in the ESI.†

Author contributions

M. Tian performed the major experimental work and wrote the original draft. J. Zhang performed the DFT calculations. L. Guo did the X-ray crystal analysis. C. Cui is the principal investigator and supervised for the project and finalized the draft of the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We are grateful to the National Natural Science Foundation of China (Grant No. 21632006) for generous support.

Notes and references

- (a) For selected reviews, see: R. Borthakur and V. Chandrasekhar, *Coord. Chem. Rev.*, 2021, **429**, 213647–213678; (b) P. P. Power, *Organometallics*, 2020, **39**, 4127–4138; (c) A. Agarwal and S. K. Bose, *Chem.-Asian J.*, 2020,

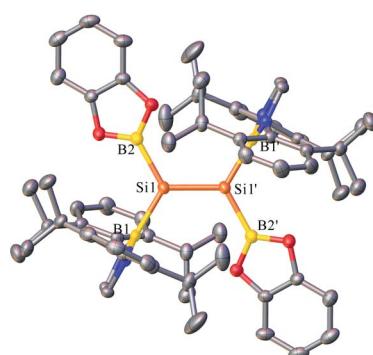


Fig. 5 OLEX2 drawing of **4** with 50% ellipsoid probability. Hydrogen atoms have been omitted for clarity.



15, 3784–3806; (d) L. Zhao, S. Pan, N. Hozlmann, P. Schwerdtfeger and G. Frenking, *Chem. Rev.*, 2019, **119**, 8781–8845; (e) R. C. Fischer and P. P. Power, *Chem. Rev.*, 2010, **110**, 3877–3923.

2 For selected reviews, see: (a) C. Präsang and D. Scheschkewitz, *Chem. Soc. Rev.*, 2016, **45**, 900–921; (b) A. Rammo and D. Scheschkewitz, *Chem.-Eur. J.*, 2018, **24**, 6866–6885; (c) A. Sekiguchi and V. Y. Lee, *Chem. Rev.*, 2003, **103**, 1429–1447.

3 (a) I. Bejan and D. Scheschkewitz, *Angew. Chem., Int. Ed.*, 2007, **46**, 5783–5786; (b) K. Abersfelder and D. Scheschkewitz, *J. Am. Chem. Soc.*, 2008, **130**, 4114–4121; (c) S. Inoue, M. Ichinohe and A. Sekiguchi, *Chem. Lett.*, 2008, **37**, 1044–1045; (d) T. Iwamoto, M. Kobayashi, K. Uchiyama, S. Sasaki, S. Nagendran, H. Isobe and M. Kira, *J. Am. Chem. Soc.*, 2009, **131**, 3156–3157; (e) J. Jeck, I. Bejan, A. J. P. White, D. Nied, F. Breher and D. Scheschkewitz, *J. Am. Chem. Soc.*, 2010, **132**, 17306–17315; (f) M. Hartmann, A. Haji-Abdi, K. Abersfelder, P. R. Haycock, A. J. P. White and D. Scheschkewitz, *Dalton Trans.*, 2010, **39**, 9288–9295; (g) P. Willmes, M. J. Cowley, M. Hartmann, M. Zimmer, V. Huch and D. Scheschkewitz, *Angew. Chem., Int. Ed.*, 2014, **53**, 2216–2220; (h) T. Iwamoto, N. Akasaka and S. Ishida, *Nat. Commun.*, 2014, **5**, 5353–5359; (i) P. Willmes, L. Junk, V. Huch, C. B. Yıldız and D. Scheschkewitz, *Angew. Chem., Int. Ed.*, 2016, **55**, 10913–10917; (j) N. M. Obeid, L. Klemmer, D. Maus, M. Zimmer, J. Jeck, I. Bejan, A. J. P. White, V. Huch, G. Jung and D. Scheschkewitz, *Dalton Trans.*, 2017, **46**, 8839–8848; (k) T. Kosai, S. Ishida and T. Iwamoto, *J. Am. Chem. Soc.*, 2017, **139**, 99–102; (l) T. Kosai, S. Ishida and T. Iwamoto, *Dalton Trans.*, 2017, **46**, 11271–11281; (m) T. Kosai and T. Iwamoto, *J. Am. Chem. Soc.*, 2017, **139**, 18146–18149; (n) T. Kosai and T. Iwamoto, *Chem.-Eur. J.*, 2018, **24**, 7774–7780; (o) Y. Heider, P. Willmes, D. Mühlhausen, L. Klemmer, M. Zimmer, V. Huch and D. Scheschkewitz, *Angew. Chem., Int. Ed.*, 2019, **58**, 1939–1944.

4 (a) D. Scheschkewitz, *Angew. Chem., Int. Ed.*, 2004, **43**, 2965–2967; (b) M. Ichinohe, K. Sanuki, S. Inoue and A. Sekiguchi, *Organometallics*, 2004, **23**, 3088–3090; (c) S. Inoue, M. Ichinohe and A. Sekiguchi, *Chem. Lett.*, 2005, **34**, 1564–1565; (d) K. Abersfelder, D. Güclü and D. Scheschkewitz, *Angew. Chem., Int. Ed.*, 2006, **45**, 1643–1645; (e) M. Ichinohe, K. Sanuki, S. Inoue and A. Sekiguchi, *Silicon Chem.*, 2006, **3**, 111–116; (f) L. Zborovsky, R. Dobrovetsky, M. Botoshansky, D. Bravo-Zhivotovskii and Y. Apeloig, *J. Am. Chem. Soc.*, 2012, **134**, 18229–18232; (g) D. Pinchuk, J. Mathew, A. Kaushansky, D. Bravo-Zhivotovskii and Y. Apeloig, *Angew. Chem., Int. Ed.*, 2016, **55**, 10258–10262; (h) L. Zhu, J. Zhang, H. Yang and C. Cui, *J. Am. Chem. Soc.*, 2019, **141**, 19600–19604; (i) I. Bejan, S. Inoue, M. Ichinohe, A. Sekiguchi and D. Scheschkewitz, *Chem.-Eur. J.*, 2008, **14**, 7119–7122; (j) S. Inoue, M. Ichinohe and A. Sekiguchi, *Angew. Chem., Int. Ed.*, 2007, **46**, 3346–3348; (k) I. Bejan, D. Güclü, S. Inoue, M. Ichinohe, A. Sekiguchi and D. Scheschkewitz, *Angew. Chem., Int. Ed.*, 2007, **46**, 3349–3352; (l) A. Jana, M. Majumder, V. Huch, M. Zimmer and D. Scheschkewitz, *Dalton Trans.*, 2014, **43**, 5175–5181; (m) H. Tanaka, S. Inoue, M. Ichinohe and A. Sekiguchi, *Organometallics*, 2009, **28**, 6625–6628; (n) T. Kosai, S. Nishimura, N. Hayakawa, T. Matsuo and T. Iwamoto, *Chem. Lett.*, 2019, **48**, 1168–1170; (o) A. Jana, V. Huch, M. Repisky, R. J. F. Berger and D. Scheschkewitz, *Angew. Chem., Int. Ed.*, 2014, **53**, 3514–3518; (p) K. Abersfelder, A. J. P. White, H. S. Rzepa and D. Scheschkewitz, *Science*, 2010, **327**, 564–566; (q) M. J. Cowley, K. Abersfelder, A. J. P. White, M. Majumdar and D. Scheschkewitz, *Chem. Commun.*, 2012, **48**, 6595–6597; (r) T. Nguyen and D. Scheschkewitz, *J. Am. Chem. Soc.*, 2005, **127**, 10174–10175; (s) N. Akasaka, K. Fujieda, E. Garoni, K. Kamada, H. Matsui, M. Nakano and T. Iwamoto, *Organometallics*, 2018, **37**, 172–175; (t) P. K. Majhi, V. Huch and D. Scheschkewitz, *Angew. Chem., Int. Ed.*, 2021, **60**, 242–246.

5 (a) A. Sekiguchi, R. Kinjo and M. Ichinohe, *Science*, 2004, **305**, 1755–1757; (b) T. Sasamori, K. Hironaka, Y. Sugiyama, N. Takagi, S. Nagase, Y. Hosoi, Y. Furukawa and N. Tokitoh, *J. Am. Chem. Soc.*, 2008, **130**, 13856–13857; (c) S. Ishida, R. Sugawara, Y. Misawa and T. Iwamoto, *Angew. Chem., Int. Ed.*, 2013, **52**, 12869–12873; (d) M. Asay and A. Sekiguchi, *Bull. Chem. Soc. Jpn.*, 2012, **85**, 1245–1261.

6 R. Kinjo, M. Ichinohe and A. Sekiguchi, *J. Am. Chem. Soc.*, 2007, **129**, 26–27.

7 (a) L. Pu, M. O. Senge, M. M. Olmstead and P. P. Power, *J. Am. Chem. Soc.*, 1998, **120**, 12682–12683; (b) L. Pu, A. D. Philips, A. F. Richards, M. Stender, R. S. Simons, M. M. Olmstead and P. P. Power, *J. Am. Chem. Soc.*, 2003, **125**, 11626–11636; (c) A. F. Richards, M. Brynda and P. P. Power, *Chem. Commun.*, 2004, 1592–1593; (d) A. Rit, J. Campos, H. Niu and S. Aldridge, *Nat. Chem.*, 2016, **8**, 1022–1026.

8 (a) Z. Liu, J. Zhang, H. Yang and C. Cui, *Organometallics*, 2020, **39**, 4164–4168; (b) M. Tian, J. Zhang, H. Yang and C. Cui, *J. Am. Chem. Soc.*, 2020, **142**, 4131–4135.

9 M. Karni and Y. Apeloig, *J. Am. Chem. Soc.*, 1990, **112**, 8590–8592.

10 (a) L. Zhu, J. Zhang and C. Cui, *Inorg. Chem.*, 2019, **58**, 12007–12010; (b) Y. Heider, P. Willmes, V. Huch, M. Zimmer and D. Scheschkewitz, *J. Am. Chem. Soc.*, 2019, **141**, 19498–19504.

11 M. Ichinohe, M. Igarashi, K. Sanuki and A. Sekiguchi, *J. Am. Chem. Soc.*, 2005, **127**, 9978–9979.

12 S. K. Sarkar, R. Chaliha, M. M. Siddiqui, S. Banerjee, A. Münch, R. Herbst-Irmer, D. Stalke, E. D. Jemmis and H. W. Roesky, *Angew. Chem., Int. Ed.*, 2020, **59**, 23015–23019.

13 (a) J. J. Eisch, B. Shafii, J. D. Odom and A. L. Rheingold, *J. Am. Chem. Soc.*, 1990, **112**, 1847–1853; (b) H. Braunschweig, H. Thomas and D. R. F. Seeler, *Angew. Chem.*, 2005, **117**, 7627–7629; (c) H. Braunschweig, A. Damme, R. D. Dewhurst, S. Ghosh, T. Kramer, B. Pfaffinger, K. Radacki and A. Vargas, *J. Am. Chem. Soc.*, 2013, **135**, 1903–1911; (d) H. Braunschweig, R. D. Dewhurst, K. Radacki, C. W. Tate and A. Vargas, *Angew. Chem., Int. Ed.*, 2014, **53**, 6263–6266.

14 (a) K. Takeuchi, M. Ikoshi, M. Ichinohe and A. Sekiguchi, *J. Am. Chem. Soc.*, 2010, **132**, 930–931; (b) K. Takeuchi,



M. Ikoshi, M. Ichinohe and A. Sekiguchi, *J. Organomet. Chem.*, 2011, **696**, 1156–1162; (c) P. Jutzi, A. Mix, B. Neumann, B. Rummel, W. W. Schoeller, H.-G. Stammle and A. B. Rozhenko, *J. Am. Chem. Soc.*, 2009, **131**, 12137–12143; (d) S. Khan, S. S. Sen, H. W. Roesky, D. Kratzert, R. Michel and D. Stalke, *Inorg. Chem.*, 2010, **49**, 9689–9693; (e) K. Izod, P. Evans and P. G. Waddell, *Angew. Chem., Int. Ed.*, 2017, **56**, 5593–5597.

15 (a) H. Watanabe, K. Takeuchi, N. Fukawa, M. Kato, M. Goto and Y. Nagai, *Chem. Lett.*, 1987, **16**, 1341–1344; (b) A. Fukazawa, Y. Li, S. Yamaguchi, H. Tsuji and K. Tamao, *J. Am. Chem. Soc.*, 2007, **129**, 14164–14165; (c) M. Kobayashi, T. Matsuo, T. Fukunaga, D. Hashizume, H. Fueno, K. Tanaka and K. Tamao, *J. Am. Chem. Soc.*, 2010, **132**, 15162–15163.

