

Cite this: *Chem. Sci.*, 2023, 14, 8956

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

Received 28th June 2023

Accepted 28th July 2023

DOI: 10.1039/d3sc03301e

rsc.li/chemical-science

# Selective synthesis of germasila-adamantanes through germanium–silicon shift processes†

Steffen Kühn,<sup>ab</sup> Benedikt Köstler,<sup>b,c</sup> Celine True,<sup>b</sup> Lena Albers,<sup>b</sup> Matthias Wagner,<sup>b,\*c</sup> Thomas Müller<sup>b,\*d</sup> and Christoph Marschner<sup>b,\*a</sup>

The regioselective synthesis of germasila-adamantanes with the germanium atoms in the bridgehead positions is described starting from cyclic precursors by a cationic sila-Wagner–Meerwein (SWM) rearrangement reaction. The SWM rearrangement allows also a deliberate shift of germanium atoms from the periphery and within the cage structures into the bridgehead positions. This opens the possibility for a synthesis of germasila-adamantanes of defined germanium content and controlled regiochemistry. In the same way that sila-adamantane can be regarded as a molecular building block of elemental silicon, the germasila-adamantane molecules represent cutouts of silicon/germanium alloys.

## Introduction

The chemistry of polysilanes has now been studied for more than 100 years.<sup>1–5</sup> While their structural chemistry resembles that of alkanes, their property of electron conjugation suggests some similarity to unsaturated hydrocarbons. Structurally, cyclic and polycyclic polysilanes are the higher congeners of cyclo- and polycycloalkanes.<sup>6,7</sup>

Adamantane cage structures of polysilanes are of particular interest as they represent the building blocks of elemental silicon. The synthesis of sila-adamantane cages nicely reflects the development of preparative approaches towards this type of compounds (Fig. 1). The first synthetic attempt, which was reported by West and Indriksons as early as in 1972, employed a Wurtz type condensation of Me<sub>2</sub>SiCl<sub>2</sub> and MeSiCl<sub>3</sub> with sodium/potassium alloy.<sup>8</sup> As the Wurtz-type approach is not very selective, the formation of the expected permethylsilaadamantane cage **I** could not unambiguously be proven. Salt elimination reactions between oligosilanides and silyl halides represented a major advancement.<sup>9</sup> This method allowed access to a cyclohexasilanyl substituted bicyclo[2.2.1]heptasilane, which served as the structural isomer required for sila-Wagner–Meerwein (SWM) rearrangement to the first structurally verified example of a sila-adamantane molecule **II**.<sup>10</sup>

Recently, Su and co-workers demonstrated site selective functionalization of both the bridging and the bridgehead positions as in adamantane **III**.<sup>11</sup> Wagner and co-workers presented a surprising simple but high yielding synthesis of germasila-adamantanes. Reactions of Si<sub>2</sub>Cl<sub>6</sub> with Me<sub>2</sub>GeCl<sub>2</sub> in the presence of (<sup>t</sup>Bu<sub>4</sub>N)Cl form adamantane cages with silicon atoms in the bridgehead and either SiCl<sub>2</sub>- or GeMe<sub>2</sub>-units in the bridging positions (compounds **IVa–IVc**, Fig. 1).<sup>12</sup>

Our present contribution deals with the selective synthesis of germasila-adamantanes from cyclic germa-oligosilanes, the deliberate formal exchange of a sila-adamantane's bridgehead silicon for an attached germanium atom, and the shift of four

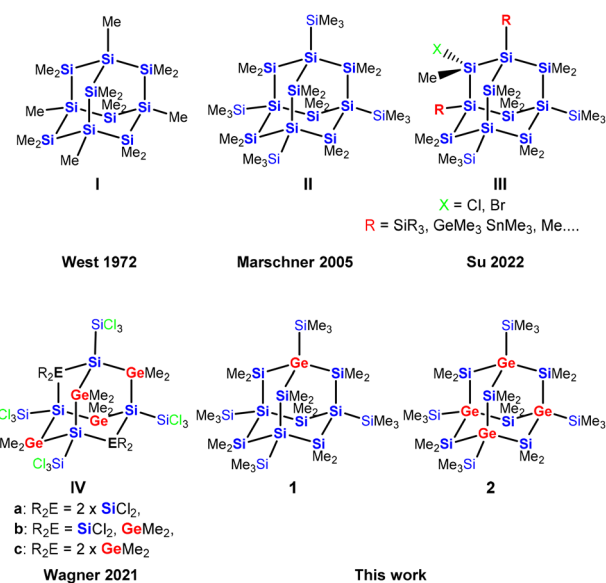


Fig. 1 Progress of the synthesis of oligosilane-adamantane type cage structures.

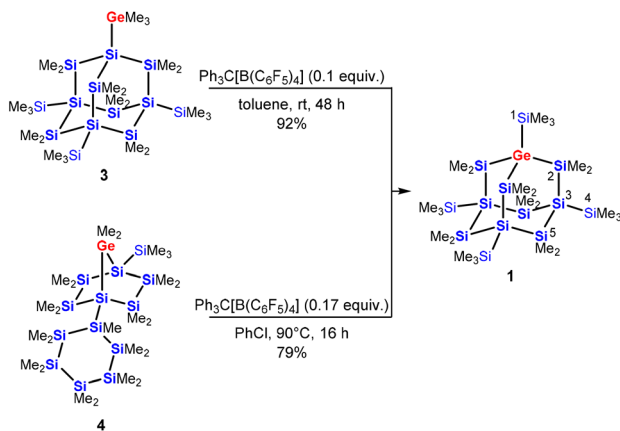
<sup>a</sup>Institut für Anorganische Chemie, Technische Universität Graz, Stremayrgasse 9, 8010 Graz, Austria

<sup>b</sup>Institut für Chemie, Carl Ossietzky Universität Oldenburg, Carl von Ossietzky-Str. 9-11, 26129 Oldenburg, Germany

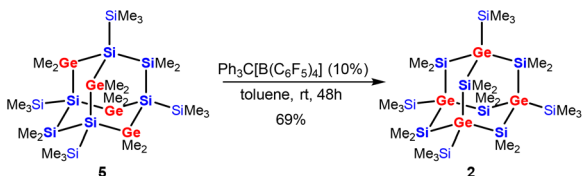
<sup>c</sup>Institut für Anorganische und Analytische Chemie, Goethe Universität Frankfurt am Main, Max-von-Laue-Str. 7, 60438 Frankfurt am Main, Germany

† Electronic supplementary information (ESI) available: Experimental procedures analytical data, NMR spectra, computational details, crystallographic information. CCDC 2257841. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d3sc03301e>

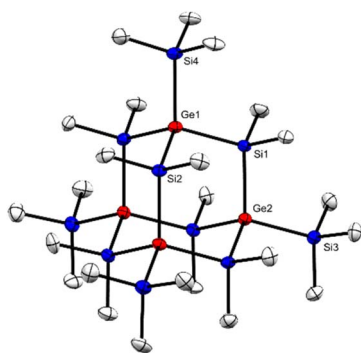




**Scheme 1** Cationic rearrangements of the trimethylgermyl-substituted sila-adamantane **3** and of the polycyclic precursor **4** to adamantane **1** with the germanium atom shifted into a bridgehead position of the adamantane cage. The descriptive numbers at compound **1** refer to NMR spectroscopic assignments.



**Scheme 2** Cationic rearrangement of the tetragermasila-adamantane **5** to the isomeric product **2** with the four germanium atoms of the adamantane cage moving from bridging into bridgehead positions.



**Fig. 2** Molecular structure of **2** (thermal ellipsoid plot drawn at the 30% probability level). All hydrogen atoms are omitted for clarity (bond lengths in [pm], angles in [deg]): Ge(1)–Si(1) 238.59(12), Ge(1)–Si(2) 238.62(11), Ge(1)–Si(4) 239.22(12), Ge(2)–Si(1) 238.72(11), Ge(2)–Si(3) 239.3(2); Si(2)–Ge(1)–Si(1) 108.53(4), Si(4)–Ge(1)–Si(1) 109.76(4), Si(4)–Ge(1)–Si(2) 110.05(4), Si(4)–Ge(1)–Si(2)#1 109.82(4), Si(1)–Ge(2)–Si(1) #2 109.33(3), Si(1)#1–Ge(2)–Si 032 109.33(3), Si(1)#1–Ge(2)–Si(1) 109.33(3), Si(3)–Ge(2)–Si(1)#2 109.61(3), Si(3)–Ge(2)–Si(1) 109.61(3), Ge(2)#1–Si(1)–Ge(1) 109.97(4).

germanium atoms from bridging into bridgehead positions, both without affecting the integrity of the adamantane cage. These transformations showcase the high synthetic capacity of SWM reactions for the tailored construction of complex

germasila-adamantane structures, which might serve as molecular models for nano-sized silicon/germanium materials. Silicon/germanium nanostructures, Si<sub>x</sub>Ge<sub>1–x</sub>, are extensively used in applications such as advanced transistors, quantum devices and photodetectors, electro-optical modulators, photo-voltaics, microelectromechanical systems and thermoelectric generators.<sup>13–19</sup> All silicon and germanium atoms of the here investigated Si/Ge cluster molecules are tetracoordinated<sup>6</sup> and therefore clearly distinguished from substituent-free silicon and germanium atoms in mixed Si/Ge Zintl ions<sup>20–22</sup> and germa-siliconoids.<sup>23–26</sup>

Oligosilanyl cations are capable of undergoing Wagner–Meerwein rearrangement chemistry in a similar way as alkanes.<sup>27</sup> This was discovered by Kumada and Ishikawa, who reported AlCl<sub>3</sub>-mediated rearrangement of permethylated linear oligosilanes to branched structures and related chemistry of cyclosilanes.<sup>28–30</sup> The strictly thermodynamic driving force of this reaction bears some synthetic potential.<sup>10,31,32</sup> In addition to structural rearrangements of oligosilanes, the Wagner–Meerwein chemistry of germa-oligosilanes allows also a selective intramolecular shift of germanium atoms into more silylated positions.<sup>33</sup> For instance, under the influence of a strong Lewis acid tris(trimethylsilyl)trimethylgermylsilane is quantitatively converted to tetrakis(trimethylsilyl)germane.<sup>34,35</sup>

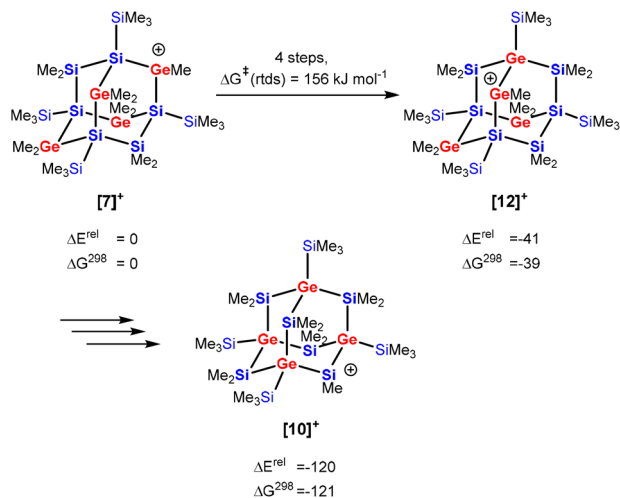
## Results and discussion

Utilizing this chemistry on a more complex substrate, we treated the trimethylgermylated sila-adamantane **3**<sup>11</sup> with a catalytic amount of Ph<sub>3</sub>C[B(C<sub>6</sub>F<sub>5</sub>)<sub>4</sub>] in toluene and observed clean conversion to compound **1** with a germa-nonasila-adamantane cage structure in 92% isolated yield (Scheme 1). Adamantane **1** was also obtained from polycyclic starting materials *via* the established route to all-sila-adamantanes: the tricyclic adamantane precursor **4** with a dimethylgermylene unit rearranges quantitatively and selectively to the germasila-adamantane **1** (79% isolated yield). In this case, slightly enforced reaction conditions had to be applied (Scheme 1).

The NMR spectrum of **1** exhibits the expected pattern. Five signals in the <sup>1</sup>H NMR spectrum at δ = 0.65 (18H), 0.62 (9H), 0.61 (9H), 0.38 (9H), and 0.35 (27H) ppm correspond to the methyl groups at Si<sup>2</sup>, Si<sup>5</sup> (2 × for Me<sub>ax</sub> and Me<sub>eq</sub>), Si<sup>1</sup>, and Si<sup>4</sup> (Si numbering in accordance with the structural formula of **1** in Scheme 1), respectively. The same number of five signals was found for the methyl groups in the <sup>13</sup>C NMR spectrum at 5.5 (Si<sup>1</sup>), 5.0 (Si<sup>4</sup>), 4.8 (Si<sup>2</sup>), 4.2, and 4.1 (Si<sup>5</sup>) ppm. Also the <sup>29</sup>Si NMR spectrum features five signals at 0.8 (Si<sup>1</sup>), –5.6 (Si<sup>4</sup>), –20.9 (Si<sup>2</sup>), –26.3 (Si<sup>5</sup>), and –118.5 (Si<sup>3</sup>) ppm (assignment of Si<sup>2</sup> and Si<sup>5</sup> by <sup>1</sup>H–<sup>29</sup>Si HMB). Comparison to the <sup>1</sup>H NMR spectrum of the all-sila-adamantane **II** (<sup>1</sup>H: δ = 0.60 (s, 36H, SiMe<sub>2</sub>), 0.35 (s, 36H, SiMe<sub>3</sub>)<sup>10</sup> shows the chemical shifts of **1** to be very similar, with all SiMe<sub>2</sub> signals around 0.60 ppm and the SiMe<sub>3</sub> signals around 0.35 ppm. The <sup>29</sup>Si NMR chemical shifts of the cyclohexasilanyl unit of the bottom part of compound **1** (*i.e.* Si<sup>3</sup>, Si<sup>4</sup>, and Si<sup>5</sup>) are very close to those of the all-sila adamantane **II** (<sup>29</sup>Si: δ = –4.8, –26.0, and –118.6), indicating no major influence of the germanium atom.







Scheme 4 Recapitulation of the isomerization reaction of tetra-germa-adamantyl cations  $[7]^+$  into  $[10]^+$  via  $[12]^+$  (energies and free enthalpies are given in  $\text{kJ mol}^{-1}$  at M06-2X/6-311+G(d,p), for details see Fig. S32†).

bonds for two stronger Si-C bonds ( $-94 \text{ kJ mol}^{-1}$ ) and of two Si-Si bonds for two weaker Ge-Si bonds ( $+48 \text{ kJ mol}^{-1}$ ). The estimation of the energy difference **6** vs. **1** based on these increments results in  $-46 \text{ kJ mol}^{-1}$  in preference of the germasila-adamantane **1**, which is close to the DFT-computed energy difference of  $-39 \text{ kJ mol}^{-1}$  (Fig. 3). While the transformation of the germasila-adamantane cages **1**, **3**, **6**, and **2**, **5** can be understood on this basis, their formation from polycyclic precursors such as **4** is less straightforward. In addition, other effects such as release of ring strain and attractive London dispersion forces between silyl groups add to the thermodynamic preference of the adamantane cluster.<sup>43,44</sup> For example, the ring strain of the bicycloheptane part of the tricyclic germasilane **4** is according to an isodesmic reaction  $47 \text{ kJ mol}^{-1}$  and the consideration of dispersion energy favors the adamantane **1** over **4** by  $37 \text{ kJ mol}^{-1}$  (for details, see ESI, Fig. S33 and S34†).

We investigated the intriguing skeletal rearrangement of the germasila-adamantane cage during the transformation  $5 \rightarrow 2$  in detail. Previous experimental and computational studies on related systems demonstrated that the initial step in this type of rearrangement is the formation of silyl- or germyl cations by transfer of a methyl group from the germa-oligosilane to the trityl cation.<sup>33</sup> After rearrangement to the more stable silyl or germyl cation, the reaction terminates with a final methyl group transfer. Methyl-anion abstraction from tetra-germasila-adamantane **5** allows the formation of three different cations  $[7]^+ - [9]^+$ . The secondary germylum cation  $[7]^+$  is the most stable one, followed by the secondary silylium cation  $[8]^+$  and the primary silylium cation  $[9]^+$  with an exo-cage dimethylsilylium substituent is the least stable cation in this series (Fig. 4) (Note: In the context of this manuscript the terms primary or secondary cation are used for cations with either one silyl/germyl substituent or with two such substituents). These cations are significantly higher in energy than the two silyl

cations ( $[10]^+$  and  $[11]^+$ ) that are possible immediate precursors for tetra-germasila-adamantane **2**. The reason for the higher stability of the cations  $[10]^+$  and  $[11]^+$  is the bridgehead position of all four germanium atoms. Therefore, the isomerization of the germylum cation  $[7]^+$  to the secondary silyl cation  $[10]^+$  is exothermic by  $120 \text{ kJ mol}^{-1}$  (Fig. 4). Association of the cations  $[7]^+ - [11]^+$  to the solvent benzene is not of importance as the energy differences between the benzene complexes  $[7(\text{C}_6\text{H}_6)]^+ - [11(\text{C}_6\text{H}_6)]^+$  are similar to those of the isolated cations and inclusion of entropy effects shows that these complexes do not exist under ambient conditions (free association enthalpy  $\Delta G^{\text{A},298} > 0 \text{ kJ mol}^{-1}$  ( $[7]^+$ ,  $[8]^+$ ,  $[10]^+$ )) or are only weakly bonded ( $\Delta G^{\text{A},298} > -15 \text{ kJ mol}^{-1}$  ( $[9]^+$ ,  $[11]^+$ ), see Table S4†).<sup>45</sup>

Based on these relative cation stabilities, we suggest the following mechanistic scenario for the trityl-induced rearrangement  $5 \rightarrow 2$ . The initial step, the exergonic methyl-anion transfer from a germylene unit of adamantane **5** to the trityl cation and formation of the germylum cation  $[7]^+$  ( $\Delta G^{\text{R}} = -68 \text{ kJ mol}^{-1}$ ), is followed by its transformation to the more stable adamantyl cation  $[10]^+$  ( $\Delta G^{\text{R}} = -121 \text{ kJ mol}^{-1}$ ). Also the terminating methyl-anion transfer from adamantane **5** to silyl cation  $[10]^+$  is thermodynamically favored ( $\Delta G^{\text{R}} = -34 \text{ kJ mol}^{-1}$ ) and closes the catalytic cycle (Scheme 3). The endergonic methyl-anion transfer from  $\text{Ph}_3\text{CMe}$  to cation  $[10]^+$  ( $\Delta G^{\text{R}} = +35 \text{ kJ mol}^{-1}$ ) suggests that the trityl borate is only the initiating reagent and not the actual catalyst.

The isomerization of secondary germylum cation  $[7]^+$  with four germanium atoms in the bridging position of the adamantane cage into the silylium cation  $[10]^+$  with all four germanium atoms in the bridgehead position is a multistep process involving several cationic intermediates. To understand this process we investigated the migration of one germanium center of cation  $[7]^+$  into a neighboring bridgehead position, *i.e.* the transformation of cation  $[7]^+$  into the germylum cation  $[12]^+$  (Scheme 4). This isomerization is thermodynamically favored (by  $39 \text{ kJ mol}^{-1}$ ) and involves three cationic intermediates and the rate determining step (rtds) is a [1,2] methyl-group shift,  $159 \text{ kJ mol}^{-1}$  above the initially formed cation  $[7]^+$  (see Fig. S32 and Table S2† for details). Finally, three similar consecutive processes transform the germylum cation  $[12]^+$  into the more stable cation  $[10]^+$ .

## Conclusions

Our results indicate that sila-Wagner-Meerwein rearrangement reactions initiated by trityl cation paired with a weakly coordinating anion can be used to prepare germasila-adamantane from a germa-oligosilane in high yields. The rearrangement reaction is highly selective, only the thermodynamic most stable isomer with the germanium atom in the bridgehead position of the adamantanes is formed. Additionally, we have shown that germanium atoms, installed at the periphery (as in **3**) or in the bridging positions of an all-sila-adamantane cage (as in **5**), can be shuttled into the bridgehead positions of the cage. The major driving force for these rearrangements is the formation of a maximum of strong Si-C bonds. These results suggest that a stepwise doping of sila-adamantanes with germanium is



possible and that adamantanes obtained by different synthetic routes with variable numbers of germanium atoms can be transformed into the most stable isomers without interfering with the cage structure. Therefore, this method allows a rational design of molecular models for mixed silicon germanium alloys. The extension of this method to Sn/Si hetero-adamantanes<sup>36</sup> and related main group element hetero-adamantanes is currently investigated in our laboratories and will further broaden the scope of this method.<sup>46</sup>

## Data availability

All data we wish to disclose are in the ESI.†

## Author contributions

Investigation and validation: S. K., C. T., L. A., B. K., T. M. Conceptualisation, methodology, supervision: M. W., C. M., T. M. Writing – original draft: C. M., T. M. Writing – review and editing: M. W., C. M. T. M.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The authors thank Dr Judith Baumgartner for crystallographic advice. Computations were done at the HPC Cluster, CARL, University of Oldenburg, funded by the DFG (INST 184/108-1 FUGG) and the Ministry of Science and Culture (MWK) of the Lower Saxony State.

## References

- 1 R. D. Miller and J. Michl, *Chem. Rev.*, 1989, **89**, 1359–1410.
- 2 J. Koe and M. Fujiki, in *Organosilicon Compounds*, ed. V. Y. Lee, Academic Press, 2017, pp. 219–300.
- 3 J. Beckmann, in *Comprehensive Organometallic Chemistry III*, ed. D. M. P. Mingos and R. H. Crabtree, Elsevier, Oxford, 2007, pp. 409–512.
- 4 C. Marschner, in *Functional Molecular Silicon Compounds I: Regular Oxidation States*, ed. D. Scheschkewitz, Springer International Publishing, Cham, 2014, pp. 163–228.
- 5 J. Baumgartner and C. Grogger, in *Comprehensive Inorganic Chemistry II*, ed. J. Reedijk and K. Poeppelmeier, Elsevier, Amsterdam, Second Edition, 2013, pp. 51–82.
- 6 Y. Heider and D. Scheschkewitz, *Chem. Rev.*, 2021, **121**, 9674–9718.
- 7 E. Hengge and R. Janoschek, *Chem. Rev.*, 1995, **95**, 1495–1526.
- 8 R. West and A. Indriksons, *J. Am. Chem. Soc.*, 1972, **94**, 6110–6115.
- 9 C. Marschner, *Organometallics*, 2006, **25**, 2110–2125.
- 10 J. Fischer, J. Baumgartner and C. Marschner, *Science*, 2005, **310**, 825.
- 11 T. C. Siu, M. Imex Aguirre Cardenas, J. Seo, K. Boctor, M. G. Shimono, I. T. Tran, V. Carta and T. A. Su, *Angew. Chem., Int. Ed.*, 2022, **61**, e202206877.
- 12 B. Köstler, M. Bolte, H.-W. Lerner and M. Wagner, *Chem.–Eur. J.*, 2021, **27**, 14401–14404.
- 13 J. Aberl, M. Brehm, T. Fromherz, J. Schuster, J. Frigerio and P. Rauter, *Opt. Express*, 2019, **27**, 32009–32018.
- 14 G. L. Wang, M. Moeen, A. Abedin, M. Kolahdouz, J. Luo, C. L. Qin, H. L. Zhu, J. Yan, H. Z. Yin, J. F. Li, C. Zhao and H. H. Radamson, *J. Appl. Phys.*, 2013, **114**, 123511.
- 15 D. Marris-Morini, V. Vakarin, J. M. Ramirez, Q. Liu, A. Ballabio, J. Frigerio, M. Montesinos, C. Alonso-Ramos, X. Le Roux, S. Serna, D. Benedikovic, D. Chrastina, L. Vivien and G. Isella, *Nano*, 2018, **7**, 1781–1793.
- 16 V. Vakarin, W. N. Ye, J. M. Ramirez, Q. Liu, J. Frigerio, A. Ballabio, G. Isella, L. Vivien, C. Alonso-Ramos, P. Cheben and D. Marris-Morini, *Opt. Express*, 2019, **27**, 9838–9847.
- 17 B. Mheen, Y.-J. Song, J.-Y. Kang and S. Hong, *ETRI J.*, 2005, **27**, 439–445.
- 18 S. Sedky, A. Witvrouw and K. Baert, *Sens. Actuators, A*, 2002, **97–98**, 503–511.
- 19 G. Scappucci, C. Kloeffel, F. A. Zwanenburg, D. Loss, M. Myronov, J.-J. Zhang, S. De Franceschi, G. Katsaros and M. Veldhorst, *Nat. Rev. Mater.*, 2021, **6**, 926–943.
- 20 T. Henneberger, W. Klein and T. F. Fässler, *Z. Anorg. Allg. Chem.*, 2018, **644**, 1018–1027.
- 21 M. Waibel, G. Raudaschl-Sieber and T. F. Fässler, *Chem.–Eur. J.*, 2011, **17**, 13391–13394.
- 22 M. Waibel and T. F. Fässler, *Inorg. Chem.*, 2013, **52**, 5861–5866.
- 23 A. Jana, V. Huch, M. Repisky, R. J. F. Berger and D. Scheschkewitz, *Angew. Chem., Int. Ed.*, 2014, **53**, 3514–3518.
- 24 L. Klemmer, V. Huch, A. Jana and D. Scheschkewitz, *Chem. Commun.*, 2019, **55**, 10100–10103.
- 25 N. E. Poitiers, V. Huch, B. Morgenstern, M. Zimmer and D. Scheschkewitz, *Angew. Chem., Int. Ed.*, 2022, **61**, e202205399.
- 26 J. Helmer, J. Droste, M. R. Hansen, A. Hepp and F. Lips, *Dalton Trans.*, 2022, **51**, 10535–10542.
- 27 H. F. T. Klare, L. Albers, L. Süssse, S. Keess, T. Müller and M. Oestreich, *Chem. Rev.*, 2021, **121**, 5889–5985.
- 28 M. Ishikawa and M. Kumada, *Chem. Commun.*, 1969, 567b–568b.
- 29 M. Kumada, *Chem. Commun.*, 1970, 157a.
- 30 M. Ishikawa, M. Watanabe, J. Iyoda, H. Ikeda and M. Kumada, *Organometallics*, 1982, **1**, 317–322.
- 31 H. Wagner, A. Wallner, J. Fischer, M. Flock, J. Baumgartner and C. Marschner, *Organometallics*, 2007, **26**, 6704–6717.
- 32 H. Wagner, J. Baumgartner, C. Marschner and P. Poelt, *Organometallics*, 2011, **30**, 3939–3954.
- 33 T. Müller, in *Organogermanium compounds: theory, experiment and applications*, ed. Lee V. Y., Wiley, Hoboken N.J., 2023, pp. 299–338.
- 34 H. Wagner, J. Baumgartner, T. Müller and C. Marschner, *J. Am. Chem. Soc.*, 2009, **131**, 5022–5023.



- 35 L. Albers, M. Aghazadeh Meshgi, J. Baumgartner, C. Marschner and T. Müller, *Organometallics*, 2015, **34**, 3756–3763.
- 36 B. Köstler, J. Gilmer, M. Bolte, A. Virovets, H.-W. Lerner, P. Albert, F. Fantuzzi and M. Wagner, *Chem. Commun.*, 2023, **59**, 2295–2298.
- 37 J. Hlina, J. Baumgartner and C. Marschner, *Organometallics*, 2010, **29**, 5289–5295.
- 38 J. Fischer, J. Baumgartner and C. Marschner, *Organometallics*, 2005, **24**, 1263–1268.
- 39 Deposition number CCDC 2257841 for **2**, contains the supplementary crystallographic data for this paper. These data are provided free of charge by the joint Cambridge Crystallographic Data Centre and Fachinformationszentrum Karlsruhe Access Structures service.
- 40 P. Pyykkö and M. Atsumi, *Chem.–Eur. J.*, 2009, **15**, 12770–12779.
- 41 C. Hemmert and H. Gornitzka, in *Organogermanium compounds: theory, experiment and applications*, ed. V. Ya. Lee, Wiley, Hoboken, N.J., 2023, pp. 667–743.
- 42 The Gaussian-16 program was used. For details, see ESI† material.
- 43 L. Albers, J. Baumgartner, C. Marschner and T. Müller, *Chem.–Eur. J.*, 2016, **22**, 7970–7977.
- 44 L. Albers, S. Rathjen, J. Baumgartner, C. Marschner and T. Müller, *J. Am. Chem. Soc.*, 2016, **138**, 6886–6892.
- 45 For an arene complex of a siladamantyl cation, see: R. J. Wehmschulte, K. K. Laali, G. L. Borosky and D. R. Powell, *Organometallics*, 2014, **33**, 2146–2149.
- 46 During the refereeing process of this manuscript, we became aware of strongly related work by Timothy Su and coworkers. M. I. Aguirre Cardenas, T. C. Siu, M. G. Shimono, S. Thai, V. Carta and T. Su, *ChemRxiv*, 2023, preprint, DOI: [10.26434/chemrxiv-2023-dk558](https://doi.org/10.26434/chemrxiv-2023-dk558).

