A germaaluminocene†

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The reactions of dipotassium germacyclopentadienediide with two Group 13 dichlorides, Cp*BCl2 and Cp*AlCl2, yield two structurally different products. In the case of boron a borole complex of germanium(II) is obtained. The aluminium halide gives an unprecedented neutral germaaluminocene. Both compounds were fully characterised by multinuclear NMR spectroscopy supported by DFT computations. The molecular structure of the germaaluminocene was determined by XRD.

Introduction

The aim of utilizing readily available and environmentally benign main group element compounds for activation of unreactive materials and strong bonds instead of transition metal-based complexes became increasingly popular during the last decade.1 We attempted to follow this lead by establishing polarised heteroalkenes I as they mimic the electronic situation in transition metal complexes (Fig. 1). The polarised frontier orbitals of these alkenes I provide the needed amphiphilic reactivity. Examples for small molecule activation by polarised heteroalkenes have already been reported.2

Our initial idea was to obtain access to polarised heterofulvenes III in which the Group 14 element is partnered with Group 13–15 elements (Scheme 1). For this reason, we studied the reaction of dipotassium germa- or silacyclopentadienediides II with a series of main group element dihalides. In several cases, we isolated rearranged products with rather unusual structures in high yields.4 For example, the reaction of aminoboron dichlorides with germacyclopentadienediide K2[1] yielded unprecedented borole complexes of germanium(II) 2c. The recent publication by Ruth and Sindlinger who reported the synthesis of a borole-based aluminocene 4 prompted us to communicate our own results on the reactivity of pentamethylcyclopentadiene-substituted boron and aluminium dichlorides versus dipotassium germacyclopentadienediide K2[1] (Scheme 2).

Results and discussion

The reaction of K2[1] with Cp*BCl2 gave the expected borole complex 2c. The NMR spectra suggested quantitative conversion and after work-up complex 2c was isolated as a brown oil in 35% yield (Scheme 2). NMR spectroscopy evidenced the presence of the expected borole ring with a η1-bound cyclopentadienyl substituent. Interestingly, the NMR data indicated frozen rotation around the B-Cσ single bond, giving rise to ten 13C NMR signals for the cyclopentadienyl substituent (Table S3, ESI †). In addition, all four carbon atoms of the borole ring are magnetically non-equivalent. Even at T = 70 °C the NMR signals show no detectable line broadening. Diagnostic for the structure of the product 2c are the high-field shifted 13C NMR resonances of carbon atoms C1/C4 compared to germole dianion (d13C(C1/C4) = 156.2 † (1); 99.8, 99.3 † (2c)). In addition, the 13C NMR signals of C1/C4 in borole 2c are markedly broadened (full width at half height, o1 = 60 Hz), due to the quadrupole moment of the neighbouring boron atom. The position of the 11B NMR resonance (d11B = 37, o2 = 283 Hz) indicates tri-coordination for the boron atom.

Fig. 1 Polarised heteroalkenes I and a sketch of their frontier orbitals.

Scheme 1 Dipotassium sila- or germacyclopentadienediides II, polarised heterofulvenes III and III(Ge).
resonance is significantly deshielded compared to that of borocene tetrachloroaluminate $\text{[AlCl}_4\text{]}$ ($\delta^{11\text{B}} = -42$) which discards the possibility of $\eta^1$-coordination. These NMR chemical shifts are all close to those reported previously for amino-substituted borole complexes $2a$ and $2b$$^{14}$ (Table S3$^\ddagger$). In addition, NMR chemical shift calculations using a DFT-optimised molecular structure of complex $2c$ provide $^{13}\text{C}$ and $^{11}\text{B}$ chemical shifts that are close to the experiment (i.e. $\delta^{11\text{B}}_{\text{calc}} = 30$) and strongly support our structural proposal for $2c$ (Table S3$^\ddagger$).

Applying the same conditions to the reaction of $\text{K}_2[1]$ with $\text{Cp}^*\text{AlCl}_2$$^8$, we noticed complete conversion of both reactants to a single product according to NMR spectroscopy. Due to its high solubility, we isolated compound 3 by crystallization in only moderate yields of 28% as yellow solid. The NMR data indicated a molecular structure very different from that of the germanium borole complex $2c$. Compound 3 is characterised by a relative sharp, high-field shifted $^{27}\text{Al}$ NMR signal at $\delta^{27}\text{Al} = -77$ ($\omega_2 = 703$ Hz). The number of signals in the $^1\text{H}$ and $^{13}\text{C}$ NMR spectra suggests a highly symmetric structure for compound 3 and the two $^{13}\text{C}$ NMR resonances of the $\text{Cp}^*$ ligand are in the typical range of $\eta^3$-coordination ($\delta^{13}\text{C} = 114.9, 10.8$) (Table S3$^\ddagger$). Even at temperatures as low as $T = -90$ $^\circ\text{C}$, all $\text{Cp}^*$-carbon atoms are magnetically equivalent, which supports the presence of a $\eta^3$-bound $\text{Cp}^*$-substituent (see ESI$^\ddagger$). The $^{13}\text{C}$ NMR chemical shifts of the heterole ring are decisively different from that of the borole ring in $2c$. In particular, the carbon atoms $\text{C1}/\text{C4}$ are significantly deshielded compared to the borole complex $2c$ or compared to the germale dianion $1^2^−$ ($\delta^{13}\text{C}(\text{C1/4}) = 167.0 \ (3)$; $99.8$, $99.3 \ (2c)$ and $156.2 \ (1^2^−)$). These $^{13}\text{C}$ and in particular $^{27}\text{Al}$ NMR data are also different from those reported for the dilithium aluminacyclopentadieneide $\text{Li}_2[6]$, ($\delta^{27}\text{Al} = 198$ ($\omega_2 = 7000$ Hz), Table S3$^\ddagger$) or for the tris-$\eta^3$-cyclopentadienyl-substituted alane 7 ($\delta^{27}\text{Al} = 64$).$^{10}$ They are however close to that of $\eta^3$-coordinated cyclopentadienyl aluminium compounds $8^−$$^{11}$ ($\delta^{27}\text{Al} = -59$ to $-150)^{11}$ which suggests a $\eta^3$-coordination of the cyclopentadienyl substituent to the aluminium atom for complex 3. The NMR data is consistent with a sandwich structure of compound 3 as depicted in Scheme 2. Further support comes from NMR chemical shift calculations for a DFT-optimised sandwich structure 3 which predict aluminium and carbon NMR chemical shifts that are very close to the experimental values ($\delta^{27}\text{Al}_{\text{calc}} = -71$, Table S3$^\ddagger$).

Yellow crystals of 3, suitable for X-ray diffraction (XRD) analysis were obtained from hexanes. In the solid state compound 3 adopts a sandwich structure with a $\eta^3$-coordinated $\text{Cp}^*$ ligand and with close contacts to all five atoms of the germale ring (Fig. 3). All germale-C-Al distances (213–228 pm) and the Ge/Al separation (248.8 pm) are significantly shorter than the respective sum of the van der Waals radii $\Sigma\text{vdW}$ ($\Sigma\text{vdW(C/Al)} = 354$ pm; $\Sigma\text{vdW(Ge/Al)} = 395$ pm).$^{12}$ The Ge–Al distance matches even that of Ge–Al single bonds in gemyl–alanes$^{13}$ and -alanates$^{14}$ (244.9–254.5 pm, theoretically predicted: 247 pm$^{15}$). The distance

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**Fig. 3** Left: Molecular structure of 3 in the crystal (Hydrogen atoms are omitted; thermal ellipsoids at 50% probability). Right: Coordination environment of the aluminium atom. Selected atom distances (pm) and angles (°): Ge–C1 199.9(10), Ge–C4 200.0(11), Al–C1 212.8(7), Al–C2 227.18(10), Al–C3 227.58(10), Al–C4 214.16(11), Ge–Al 248.78(4), C1–C2 144.70(13), C2–C3 142.12(14), C3–C4 144.56(13), Al–Ctr. 186.64(3), Al–C($\text{Cp}^*$) 217–228, $\alpha$(Ge) 165.638(9), $\alpha$(Al) 91.628(10).
between the centre of the Cp* ring and the aluminium atom (186.6 pm) is only slightly larger than that found for the Cp*Al(\(\text{I}\)) complex \(^9\) \(^\text{11}\) but significantly smaller than found for monomeric Cp*Al(\(\text{I}\)) \(^9\) \(^\text{11}\) (see Table S2†).

The planes defined by the Cp* ring and by the four carbon atoms of the germole ligand are aligned almost parallel (inter planar angle 3.3°). The C–C bond lengths in the germacycle indicate delocalization (142.1–144.7 pm). Compared to the germole dianion \(^1\text{2–}\) in the triple ion pair \(\text{K}_2[\text{I}]\) (139.5–142.0 pm),\(^3\) or in the germole dianion hafnium complexes \(^1\text{2–}\) (134.7–142.8 pm)\(^\text{17}\) or \(^3\) (140.7–144.5 pm) (Fig. 4),\(^4\) all bonds in the germole ring of \(^3\) are slightly elongated, indicating electron transfer from the germole ring to the aluminium atom (Fig. 3).

Quantum mechanical calculations at the M06-2X/6-311+G(d,p) level of theory were performed to investigate the bonding situation in neutral germaaluminocenes \(^3\) and \(^3\) \(^\text{M}\) \(^3\) \(^\text{M}\) is a close model to \(^3\), in which all substituents at the germole ring and at the cyclopentadiene ring have been replaced by hydrogen atoms).\(^\text{18}\) The analysis of the frontier molecular orbitals suggests a substantial germylene character of the sandwich complex \(^3\) \(^\text{M}\). This is shown by surface diagrams of the frontier orbitals which indicate a significant contribution of an in-plane lone pair at germanium to HOMO–1 (Fig. 5). In addition, the 4p(Ge) atomic orbital dominates the LUMO and the LUMO+1. The HOMO is essentially a combination of an \(\pi\) orbital of the germole cycle that is polarised towards the germanium atom and atomic orbitals of the aluminium. The results of a natural bond orbital (NBO) analysis for complex \(^3\) indicate significant charge transfer from the germole ring to the aluminium atom as shown by the calculated NBO group charges (Fig. 6). The covalent character of the aluminium/germole interaction is supported by the calculated Wiberg bond indices (WBI).\(^1\text{9}\) These exceed those calculated for the ionic interaction between the potassium ions and the germole ring for \(\text{K}_2[\text{I}]\) (Fig. 6). In particular, the WBIs for the C1/C4–Al linkages are larger than those of C(Cp*)–Al bonds in \(^8\) \(^+\) and close to those predicted for the Al–C bond in Me–Al(\(\text{I}\)) (Fig. 6).

This covalent bonding is visualised by a natural localised molecular orbital (NLMO, Fig. 7a), which indicates the electron delocalisation between atomic orbitals of the aluminium and \(\pi\) type orbitals of the germole ligand. The complementary analysis of the calculated electron density of germaaluminocene \(^3\) in the framework of the quantum theory of atoms in molecules (QTAIM) finally reveals a consistent picture with bond paths between the aluminium atom and the carbon atoms C1/4 and a valence shell charge concentration (VSCC) between the aluminium atom and the germanium atom (Fig. 7b).

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Fig. 4 Selected germole dianion metal complexes.

Fig. 5 Calculated surface diagrams of frontier molecular orbitals of germaaluminocene \(^3\) \(^\text{M}\) (M06-2X/6-311+G(d,p), isodensity value 0.04).

Fig. 6 Calculated WBIs of pertinent bonds and NBO group charges \(q\) of complexes \(^3\), \(^\text{K}_2[\text{I}]\), \(^8\) \(^+\) and aluminium methyl compounds (M06-2X/6-311+G(d,p)).

Fig. 7 (a) Surface diagram of the NLMO of \(^3\) \(^\text{M}\) showing the interaction between the germole ring and the CpAl fragment (surface value 0.04, hydrogen atoms omitted). (b) Molecular graph of \(^3\) \(^\text{M}\) projected on a contour plot of the Laplacian of the electron density in the molecular mirror plane (M06-2X/6-311+G(d,p)).
On the basis of our theoretical results we suggest a delocalised bonding scheme for model germaaluminocene compound 3M and likewise for the experimentally investigated compound 3. Scheme 3 shows the three extreme Lewis forms of the delocalisation of the bonding situation in the germaaluminocene.

In the course of our computational investigation of possible reaction channels for the formation of germanium borole or alumole complexes (2c and 14) or germaborocene or -alumina-nocene sandwich compounds (15 and 3), we found that structures with an Y-shaped arrangement\(^\text{26}\) around the two heteroatoms, as it is shown by boraalkene 16\(^1\) or boraasilene 17,\(^2\) are only high lying transition states (see ESI\(^\dagger\) for a more extensive description of the corresponding potential energy surfaces). This is surprising, in view of the recent isolation of an almost linear boragermene 18\(^*\) by Rao and Kinjo, who used an amino substituent at the boron atom (Fig. 9).

As reported previously we synthesised borole complexes of germanium(0) 2a and 2b from the reaction of a germacyclopentadienediide 1\(^{2-}\) with amino boron dichlorides.\(^3\) Now, we found the same structural motif when using the Cp\(^*\) substituent at the boron atom. This suggests that the germacyclopentadienyl fragment, used in the chemistry presented here, has a significant effect on the reaction outcome.

In agreement with the exclusive formation of the borole germanium(0) complex 2c, we found that 2c is significant more stable than the isomeric germaborocene 15 (\(\Delta E = 63\) kJ mol\(^{-1}\)), Fig. 8). The situation is different for the isomeric aluminium compounds 3 and 14. In qualitative agreement with the isolation of the germaaluminocene 3 (Fig. 8), it is more stable than the alumole germanium(0) complex 14 by \(\Delta E = 24\) kJ mol\(^{-1}\). We notice however that their energy difference is significantly smaller. Moreover, the barrier for the interconversion 3 \(\rightarrow\) 14 (78 kJ mol\(^{-1}\)) is relatively small (Fig. 8). These computational results for the aluminium compounds indicate the possibility to influence the product formation significantly by substituent effects. Decisive for the higher stability of the germametallocene 3 vs. the metallocene Ge(0) complex 14 is the larger size of the aluminium atom compared to the boron atom in the isomer pair 2c/15. Its inclusion into the delocalised metallole cycle is disfavoured due to the misfit of 3p(Al) and 2p(C) orbitals but its larger size facilitates the \(\eta^3\)-coordination to the Cp* ligand.\(^7\)\(^9\)\(^23\)

**Conclusions**

In agreement with previous results, the reaction of \(K_2[\text{1}]\) Cp\(^*\)BCl\(_2\) gives a borole complex of Ge(0) 2c. The respective reaction of \(K_2[\text{1}]\) with Cp\(^*\)AlCl\(_2\) provides access to the neutral germaaluminocene 3, which represents a new class of aluminium \(\pi\)-complexes. Similar to aluminocenium cation 8\(^8\)\(^\text{11a,14}\) and boraaluminocene 4,\(^3\) germaaluminocene 3 shows a \(\eta^3,\eta^1\)-coordination of both five-membered rings. In contrast to these previous examples, germaaluminocene 3 exhibits a free electron pair and a low-lying acceptor orbital at the germanium atom, which suggests additional germylene-like reactivity.

The small energy difference between the germole 3 and isomeric alumole 14 structures and the small energetic barriers for their interconversion imply that also the synthesis of alumole complexes of Ge(0) is in reach, when substituent effects are employed advantageously.

**Conflicts of interest**

There are no conflicts to declare.

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