In contrast to molecular CO and \( \text{CO}_2 \), their heavier mono- and dichalcogenide homologues, \( \text{EX} \) and \( \text{EX}_2 \) (\( \text{E} = \text{Si, Ge, Sn, Pb; } \text{X} = \text{O, S, Se, Te} \)), are important support materials (e.g., \( \text{SiO}_2 \)) and/or semiconductors (e.g., \( \text{SiS}_2 \)) and exist typically as insoluble crystalline or amorphous polymers under normal conditions. Herein, we report the first successful synthesis and characterisation of an extraordinary series of soluble monomeric \( \text{GeX} \) and \( \text{GeX}_2 \) complexes \( (\text{X} = \text{S, Se, Te}) \), representing novel classes of compounds and heavier congeners of \( \text{CO} \) and \( \text{CO}_2 \). This could be achieved by solvent-dependent oxidation reactions of the new zero-valent germanium \( (\text{germynolone})-\text{GaCl}_3 \) precursor adduct \( (\text{bis-NHC})\text{Ge}^{0}\rightarrow\text{GaCl}_3 \) 1 and \( \text{GaCl}_3 (\text{bis-NHC}) \equiv \text{H}_2\text{Cl}[\text{N=CH(NH)}\text{Dipp}]\text{C}(\text{H})(\text{N}(\text{Dipp})\text{C})\text{Cl}_2 \), \( \text{Dipp} = 2,6-\text{Pr}_2\text{C}_6\text{H}_3 \) with elemental chalcogens, affording the donor–acceptor stabilised monomeric germanium(\( \text{iv} \)) dichalcogenide \( (\text{bis-NHC})\text{Ge}^{\text{iv}}(\equiv=\text{X})=\text{X}\rightarrow\text{GaCl}_3 \) \( (\text{X} = \text{S, 2; X = Se, 3}) \) and germanium(\( \text{iv} \)) monochalcogenide complexes \( (\text{bis-NHC})\text{Ge}^{\text{iv}}=\text{X}=\text{GaCl}_3 (\text{X} = \text{Se, 4; X = Te, 5}) \) respectively. Moreover, the reactivity of \( 4 \) and \( 5 \) towards elemental sulphur, selenium, and tellurium has been investigated. In THF, the germanium(\( \text{iv} \)) monoselene complex \( 4 \) reacts with activated elemental selenium to afford the desired germanium(\( \text{iv} \)) diselenide complex \( 3 \). Unexpectedly, both reactions of \( 4 \) and \( 5 \) with elemental sulphur, however, lead to the formation of germanium(\( \text{iv} \)) disulfide complex \( 2 \) under liberation of elemental Se and Te as a result of further oxidation of the germanium centre and replacement of the Se and Te atoms by sulphur atoms. All novel compounds \( 1–5 \) have been fully characterised, including single-crystal X-ray diffraction analyses, and studied by DFT calculations.

Generally, they adopt polymeric structures owing to the relatively weak \( p_n\rightarrow p_n \) bond between germanium and the respective chalcogen atom and the high polarity of the \( \text{E-X} \) bond. It has been shown that the molecular variants of \( \text{EX} \) and \( \text{EX}_2 \) can exist in condensed cryogenic matrices at very low temperature or diluted in the gas phase at high temperature. Thus they have solely been detected spectroscopically\(^2\) under extreme conditions or proposed as reactive intermediates\(^3\) and studied by theoretical calculations.\(^4\)

During the last decades, the concept of kinetic and/or thermodynamic stabilisation has enabled great achievements in synthesising isolable low-coordinate group 14 element complexes as ligands in complexes. Several unusual compounds featuring elusive terminal \( \text{E} = \text{O} (\text{E} = \text{Si, Ge, Sn, Pb})\) and \( \text{E} = \text{X (X = S, Se, Te})\) double bonds could be synthesised and structurally characterised. However, complexes containing \( \text{EX} \) and \( \text{EX}_2 \) are still very rare. The first examples include the Sn=O (I) and Pb=O (II) units stabilised by a benzannulated bis-stannylene reported by Hahn and co-workers (Chart 1).\(^2\) Very recently, Dehnen et al. published the first Pb-Se complex in \( [\text{K}(18\text{-crown-6})][\text{K(en)}_2] \)
Results and discussion

Shortly after communicating the bis-NHC supported germanium(0) species (germylone B, Scheme 1),\textsuperscript{10} we realized that B is sensitive not only towards air and moisture, but also to visible light. For its reactivity investigation, we introduced the Lewis acid GaCl\textsubscript{3} to prepare the more stable germylone–GaCl\textsubscript{3} adduct 1 through a one-pot reaction, starting from the bis-NHC supported chlorogermilumylidene chloride (bis-NHC)GeCl\textsubscript{3} A (Scheme 1).\textsuperscript{10} Accordingly, the germylone B, prepared by reduction of A with two molar equivalents of sodium naphthalenide in THF, reacts \textit{in situ} with one molar equivalent of GaCl\textsubscript{3} to furnish the desired complex 1 as a colourless precipitate in 65% yield after work-up (Scheme 1).

The proton NMR spectrum of 1 reflects the coordination of the germanium centre \textit{via} one of its lone pair of electrons to the GaCl\textsubscript{3} moiety, thus lowering the symmetry of the molecule relative to that of its precursor germylone B. Therefore the spectrum of 1 exhibits four doublets for the methyl protons and two septets for the methine protons in the isopropyl groups. Moreover, the two geminal protons in the CH\textsubscript{2} moiety on the backbone in 1 give two doublets with 2\textsubscript{J}GH = 13.3 Hz (AB-spin system, ESI\textsuperscript{†}). Complex 1 crystallised from acetonitrile solution in triclinic space group P1 with four lattice CH\textsubscript{2}CN molecules in the asymmetric unit (Fig. 1). A single-crystal X-ray diffraction analysis of 1 revealed a three-coordinate germanium centre featuring pyramidal coordination geometry. The sum of the angles around the germanium atom amounts to 266.3°, implying that the vertex of the pyramid is occupied by one pair of electrons. Owing to the coordination of GaCl\textsubscript{3}, the average Ge–C bond length of 2.038(3) Å in 1 is longer than that in germylone B (Ge–C 1.965(2) Å). On the other hand, the Ge1–Ga1 distance of 2.520(1) Å is comparable to the Ge–Ga single bond (2.516(1) Å) supported by bulky substituents in [(Dipp)N–CH=CH–N(Dipp)]Ga–Ge(N(Dipp))\textsubscript{2}CN(iPr)\textsubscript{2}.\textsuperscript{12a}

Complex 1 is much more stable than its precursor B because the zero valent germanium centre is coordinated to the Lewis-acid GeCl\textsubscript{3} by donating one of the two lone pairs of electrons to the electron-deficient gallium atom. It is noteworthy that

\[ \text{K[Rh}_{3}(\text{CN})_{6}(\text{PPh}_{3})_{4}(\text{H}_{2} \text{Se})_{2}(\text{µ-PbSe})_{2}]_{2} \rightarrow \text{en = ethane-1,2-diamine} \]
germylone \( \text{B} \) also reacts readily with other Lewis acids such as \( \text{AlBr}_3, \text{BCl}_3 \), but the desired products could not be isolated in pure form as yet. However, \( \text{B} \) reacts smoothly with \( \text{GaCl}_3 \) to afford \( \text{I} \) as an isolable product. Compound \( \text{I} \) is soluble in \( \text{THF} \) and acetonitrile. Treatment of \( \text{THF} \) solutions of \( \text{I} \) with \( \text{GaCl}_3 \) was also unsuccessful which high-

be recorded in solutions but their low solubility prevents VT-

NMR spectroscopy at low temperature.

Fortunately, single-crystals of \( \text{2} \) and \( \text{3} \) suitable for X-ray

diffraction analyses were obtained in dilute \( \text{THF} \) solutions. Unexpectedly, compounds \( \text{2} \) and \( \text{3} \) exhibit higher symmetry than \( \text{1} \), that is, only two doublets for the methyl protons in the \( \text{CH} (\text{CH}_3)_2 \) groups, instead of four doublets as in \( \text{1} \), could be detected in their \( ^1 \text{H} \) NMR spectra, suggesting the statistically equal position of the \( \text{GaCl}_3 \) moiety between two sulphur or selenium atoms in solution, respectively (see ESI†). The ESI-MS of \( \text{2} \) and \( \text{3} \) show the \( \text{GaCl}_3 \)-free molecular ion peak at \( m/z = 607.1969 \) (calc. 607.19789 for \( [2 - \text{GaCl}_3 + \text{H}^+] \)), corresponding to \( [(\text{bis-NHC})\text{GeS}_2 + \text{H}^+] \) and at \( m/z = 703.08749 \) (calc. 703.08680 for \( [3 - \text{GaCl}_3 + \text{H}^+] \), corresponding to \( [(\text{bis-NHC})\text{GeSe}_2 + \text{H}^+] \), respectively.

Compounds \( \text{2} \) and \( \text{3} \) crystallise isotypic in the monoclinic space group \( \text{Cm} \) with two lattice \( \text{THF} \) molecules in the respective asymmetric unit and the single-crystal X-ray diffraction analyses revealed them to be isostructural, with each of the germanium centres bound to two chalcogen atoms (Fig. 2). Thus the germanium centres in both compounds are four-coordinate and adopt a tetrahedral geometry with almost identical bond angles around the germanium atoms. The \( \text{S1–Ge1–S2} \) angle of \( 115.3(1) \) in \( \text{2} \) and \( \text{Se1–Ge1–Se2} \) angle of \( 115.2(1) \) in \( \text{3} \), respectively, are reminiscent of the corresponding \( \text{S–Si–S} \) value observed in \( [(\text{bis-NHC})\text{Si} (\text{S}) \text{SH} + \text{H}^+] \). The \( \text{C1–Ge1–C'1} \) angles of \( 91.4(2) \) in \( \text{2} \) and \( 91.8(3) \) in \( \text{3} \) are larger than that in \( \text{I} \) (85.7(1)°): accordingly, the \( \text{Ge–C} \) bond distances (1.998(3) \( \AA \) in \( \text{2} \) and 1.987(5) \( \AA \) in \( \text{3} \)) are slightly shorter than that in \( \text{I} \) (2.038 \( \AA \)).

The two \( \text{Ge–S} \) distances in \( \text{2} \) (2.087(1) \( \text{vs.} \) 2.198(1) \( \AA \)) and the two \( \text{Ge–Se} \) bonds in \( \text{3} \) (2.214(1) \( \text{vs.} \) 2.326(1) \( \AA \)) show a significant difference. The \( \text{Ge1–S1} \) bond length of 2.087(1) \( \text{Å} \) in \( \text{2} \) is slightly longer than that in \( \text{Tbt(Tip)Ge} \) (2.049 \( \text{Å} \)), but comparable to the \( \text{Ge=Se} \) double bonds in \( \text{LGe(}\text{S}) \text{SH} \) (2.064(1) \( \text{Å} \)), and
LGe(—S)–SPh 2.071(1) Å (L = monovalent chelating organic groups,

whereas the Ge1–S2 bond of 2.198(1) Å is slightly shorter than the Ge–S single bonds in the latter species (2.242(1) Å,

ref. 12d). Similarly, the Ge1–Se1 distance of 2.214(1) Å in 3 is slightly longer than that in Tbt(Tip)Ge=Se (2.180 Å, Tbt = 2,4,6-tris(3-trimethylsilyl)methylphenyl, Tip = 2,4,6-tris(isopropylphenyl)),

but close to the doubly bonded Ge=Se group in LGe(—Se)–SePh (2.205(1) Å, L = amino-

troponinato ligand),

while the Ge1–Se2 of 3.236(1) Å in 3 is only slightly shorter than the Ge–Se single bond in the latter complex (3.267(1) Å).

The relatively large differences in the two Ge–X lengths (X = S, Se) in 2 and 3 can be understood in light of the calculations presented below.

Although the mechanism of formation of 2 and 3 is still unknown, it is reasonable to propose that the dichalcogenide formation occurs stepwise via the respective monocohalogenides as shown in Scheme 2. The initial step of the reaction implies the oxidation of the zero-valent germanium centre in 1 by one chalcogen atom to yield the divalent GeX complex C. The subsequent GaCl3 migration from the Ge(n) centre to the more Lewis-basic chalcogenide site affords D bearing a three-coordinate Ge(n) centre. Subsequent oxidation of the latter from Ge(n) to Ge(n) by an additional chalcogen atom results in the final product 2 or 3, respectively. These suggestions are supported by results of DFT calculations. 13 The calculated free energies (at room temperature) for the reaction 1 → 2 (and 3) are shown in Fig. S7 in the ESI†. The migration of the GaCl3 fragment, C → D (Scheme 2, Fig. S7†), is exothermic, in the range of 11.8 to 15.8 kcal mol−1, for both X = S and Se, in the gas phase, in acetonitrile and in THF. However, the migration of GaCl3 is less exothermic for X = Se than for X = S by 3 kcal mol−1 in CH3CN. The free energy barriers for the two-step GaCl3 migration (C → D) is 29.9 and 29.5 kcal mol−1 for X = S and X = Se, respectively, in the gas phase (see details in the ESI†). The reaction leading from D (X = S) to 2 is by 5.0 kcal mol−1 more exothermic than for D (X = Se) to 3 (gas phase). The overall reaction from 1 to 2 (X = S) is more exothermic than from 1 to 3 (X = Se) by 10.3 (gas phase) 11.3 (CH3CN), and 10.9 (THF) kcal mol−1, in line with the experimental observation of a faster reaction for X = S than for X = Se. 14

In order to isolate the proposed divalent germanium intermediates C and/or D, 1/8 equivalent of S8 and Se8 were employed for the reaction with 1 at −30 °C in THF solutions. However, even at low temperature, regardless of the ratio of the two reactants, the dichalcogenides 2 and 3 are the exclusive products. In contrast, by employing acetonitrile as solvent the reaction of 1 with 1/8 equivalents of S8 at room temperature became much slower than in THF, and the reaction afforded a mixture containing 2 and, presumably, C and/or D (X = S). Furthermore, the reaction of 1 with 1/8 molar equivalents of activated selenium Se8 at room temperature became so slow that the formation of the germanium(n) monoselsenide complex 4, the selenium version of D (Scheme 2 and 3) was realized. In fact, after 3 days, 4 could be isolated from the resulting solution as a colourless solid in moderate yield (64%). For the reaction of 1 with elemental tellurium, the reaction is even slower than that with selenium and furnishes in THF at room temperature only the germanium(II) monotelluride complex 5 (Scheme 3).

Compound 4 is better soluble in THF and CH3CN than the diselenide complex 3. This allowed us to record its 1H- and 13C-NMR spectra in CD3CN at room temperature. The four doublets for the methyl protons and two septets for the methine protons in the CHMe2 groups indicate similar molecular symmetry to that of 1. Again a similar “roof” effect was observed for the geminal protons of the bridging N–CH2–N group (1JCHH = 13.4 Hz) in the 1H NMR spectrum of 4 (AB-spin system). The ESI-MS spectrum of 4 (positive mode) shows an ion peak of [M – SeGaCl3 + Cl]+ at m/z = 577.21454 (calc. 577.21478) corresponding to the [bis-NHC]GeCl] (A)+ cation. Indeed, the facile formation of similar species with a [LECl]+ cation (L = 1,8-bis(tributylphosphazenylnaphthalene, E = Si, Ge) have been described in our previous reports. 15

Akin to 4, the 1H NMR spectrum of 5 exhibits four doublets for the methyl protons and two septets for the methine protons in the CHMe2 groups. Similarly, the bridging N–CH2–N–N protons are coupled with each other with 1JCHH = 13.1 Hz (AB-spin system). Similar to that of 4, in the ESI-MS spectrum of 5 a signal at m/z = 577.21399 (calc. 577.21478 for [M – Te – GaCl3 + Cl]+ corresponding to A+ was observed.

Compound 4 crystallises in acetonitrile solutions in the monoclinic space group P21/c with one lattice CH3CN molecule in the asymmetric unit (Fig. 3, left). The single-crystal X-ray diffraction analysis confirmed the proposed structure in which the germanium(II) centre is stabilised by the chelating bis-NHC ligand and the Ge=Se moiety is supported by GaCl3 coordination, leading to a Ge–Se–Ga angle of 110.7(1)°. The three-coor-
dinate Ge(n) centre features a pyramidal coordination geometry with a sum of angle of 289.5°. The Ge–C distance of 2.050(3) Å in 4 is very close to that observed in 1 (2.038(3) Å). The Se1–Ga1 length of 2.337(1) Å is comparable to that in 3 (2.374(1) Å) with a similar coordination environment. The Ge1–Se1 distance (2.438(1) Å) is significantly longer than the two Ge–Se bonds in the four-coordinate germanium diselenide complex 3 (2.214(1) and 2.326(1) Å). It is also longer than the Se–Ge length of 2.346 Å in the [CO]W=Ge(SeAr)2 (Ar = 2,4,6-tris(isopropylphenyl)germanium(n) species with a three-coordinate planar Ge centre. Thus a Ge=Se double bond character can not be
concluded from the Ge–Se distance. This is also manifested in the resonance structures 4” calculated by NRT which are discussed below.

Single crystals of 5 suitable for X-ray diffraction analysis were obtained from THF solutions. 5 crystallises in the triclinic space group P1 with three lattice THF molecules in the asymmetric unit. The latter analysis revealed a similar structure to that of 4 (Fig. 3, right). The germanium monoteluride moiety is coordinated by the bis-NHC and GaCl3 ligands, leading to a Ge–Te–Ga angle of 109.0(1)°. The germanium centre also adopts a trigonal–pyramidal coordination geometry with a sum of bond angles of 285.8° around the germanium atom. The average Ge–C bond length of 2.037(4) Å in 5 is close to those in its precursor 1 (2.038(3) Å) and in 4 (2.050(2) Å). The Ge1–Te1 distance of 2.654(1) Å is significantly longer than that in a four-coordinate germanium(v) species LGeR (2.461(7) Å) (L = monovalent chelating organic group, R = monovalent organic group).17 Akin to the situation for the selenide 4, the Ge–Te double bond character in 5 is rather low. It is noteworthy that the 125Te NMR resonance of 5, similar to the 77Se NMR resonances of 3 and 4, could not be observed, presumably owing to the zwiterionic nature of 3 and 4 (Scheme 5 and 6 and Fig. S6 in ESI†) which result in the broadening of the 125Te NMR resonances (for calculated chemical shifts see Table S12 in ESI†). Similar signal broadening occurred in the dimeric SiSe2 species stabilised by CAAC.55

Since the germanium centres in 4 and 5 still feature one lone pair of electrons, their reactivity toward elemental sulphur, selenium, and tellurium has been investigated. Surprisingly, both 4 and 5 react with elemental sulphur to yield 2 immediately either in THF or in CH3CN, as shown by the 1H NMR spectra and by the precipitation of elemental orange Se and black Te, respectively (Scheme 4). Although the intermediates E and F could not be detected, they might play a role in the latter reactions. Accordingly, intermediates E, resulting from oxidation of 4 or 5 by sulphur, might undergo GaCl3 migration from X (X = Se, Te) to sulphur, spontaneously yielding F. Subsequent displacement reaction of the latter should afford 2 as the final product. The latter scenario is supported by DFT calculations. For instance, the calculated reaction free energy (at B3LYP-D3(BJ)/def2-SV) using the PCM model for the solvents’ effect) of 4 + 1/4 $\text{S}_8$ → 2 + 1/8 $\text{Se}_8$ is −21.6 (gas phase), −25.2 (acetonitrile) and −22.6 (THF) kcal mol$^{-1}$. The migration from E to F is nearly thermo-neutral ($\Delta G = −0.74$ (gas phase), −1.60 (acetonitrile), and −0.64 (THF) kcal mol$^{-1}$).14

As expected, the reaction of 4 with activated selenium can afford compound 3 in CH3CN. However, after 24 h only 10% of 4 reacted. In contrast, in THF the reaction is complete after 2 h. This may explain why 4 could be isolated in CH3CN, whereas only 3 was isolable in THF. On the other hand, no reaction of 5 with elemental selenium in both THF and CH3CN could be detected after 24 h, confirming that 5 is less reactive than 4. Furthermore, the reactivity of 4 towards tellurium in THF at room temperature was probed, however, after three days, no reaction occurred.

To obtain better understanding of the electronic properties and reactions of the novel compounds described in this article we performed DFT calculations for the synthesised compounds 1–5 and for the respective model compounds 1′–5′ where the bulky Dipp groups are replaced by Ph groups.41 In general the calculated optimized geometries of 1–5 at B3LYP-D3(BJ)/def2-TZVPP and those of 1′–5′ optimised at B3LYP-D3(BJ)/def2-SV are in good agreement with the corresponding X-ray structures.44 A selection of calculated geometry parameters are presented in Table 1 (more details are provided in Table S11 in ESI†). Unless otherwise stated we discuss the calculated results obtained at the B3LYP-D3(BJ)/def2-0.64 SV level of theory.

The calculated Ge1–X1 distances (2′, X = S, and 3′, X = Se) of 2.091 Å and 2.231 Å are longer than those of Me2Ge= X (X = S: 2.045 Å, X = Se: 2.174 Å), respectively. This trend is in line with the Ge–X Wiberg Bond Indices (WBI)49 which in Me2Ge= X (X = S, 1.82 and X = Se, 1.84) are larger than those of 2′ (1.39) and 3′ (1.36). The calculated Ge= X bond length and WBIs in linear X=Ge=X are 2.016 Å and 1.79, respectively, for X = S, and
The dichalcogenides 2 and 3 as well as the monochalcogenides 4 and 5 are highly delocalised compounds featuring many resonance structures. The most predominant resonance structures according to NRT calculations, for model compounds 2′′ and 3′′ (R = Me), feature two resonance structure types which are responsible for ca. 75% of the total contributions (Scheme 5): (a) structures containing a X1=Ge-X-GaCl3 subunit with a X1=Ge double bond and where the Ge is bound to only one of the carbenes, accounting for 23%, for both X = S and X = Se, of the total. (b) Structures containing a X=Ge-X-GaCl3 subunit with a bond between the Ge and each of the NHC units. These structures account for 53% (X = S) and 55% (X = Se) of the total. Many electron permutations in the NHC rings are possible for both resonance structure types and the values in Scheme 5 present their summation. These resonance structures indicate a partial double bond character of the Ge–X1 and of the endocyclic C–N bonds and a single bond character of the Ge–X2 bond. The corresponding NRT bond orders of Ge–X1 are 1.38, with a covalent contribution of 0.85 and an ionic contribution of 0.52 (X = S) and 1.36 with a covalent contribution of 0.94 and an ionic contribution of 0.41 (X = Se), reflecting the lower polarity of the Ge–Se bonds (see Fig. S6 in the ESI†). The NRT bond orders of Ge–X2 are 0.95 and 0.96 for X = S and X = Se, respectively. This bond orders are in good agreement with the calculated WBIs and explain the trends in the bond lengths (see above and in Table 1).

The relatively long Ge–Se and Ge–Te distances in the monochalcogenides 4 and 5 (see discussion above and Table 1) led to the conclusion that there is no Ge=X double bond in 4 and 5. This is supported by the NRT calculations for 4′′ and 5′′ R = Me which show that the major contribution stems from the resonance structures shown in Scheme 6 and their possible resonance permutations within the NHC fragments with no contribution from a Ge=X (X = Se, Te) double bond.

| Compound | Ge–C | Ge–X1 | Ge–X2 | C–N1 | C–N2 | X2–Ga |
|----------|------|-------|-------|------|------|-------|
| 1, 1′     | 2.043 | —     | —     | 1.349 | 1.354 | —     |
| r (exp.)  | 2.045 | —     | —     | 1.356 | 1.360 | —     |
| WBI       | 0.74  | —     | —     | 1.23  | 1.21  | —     |
| 2, 2′     | 1.998 | 2.087 | 2.198 | 1.329 | 1.358 | 2.249 |
| r (calc.) | 2.022 | 2.091 | 2.213 | 1.344 | 1.354 | 2.280 |
| WBI       | 0.59  | 1.39  | 0.92  | 1.28  | 1.24  | 0.71  |
| 3, 3′     | 1.987 | 2.214 | 2.326 | 1.352 | 1.366 | 2.374 |
| r (exp.)  | 2.020 | 2.231 | 2.348 | 1.344 | 1.354 | 2.404 |
| WBI       | 0.60  | 1.36  | 0.94  | 1.28  | 1.24  | 0.75  |
| 4, 4′     | 2.047 | —     | 2.438 | 1.346 | 1.353 | 2.337 |
| r (exp.)  | 2.071 | —     | 2.435 | 1.352 | 1.358 | 2.381 |
| WBI       | 0.67  | 0.89  | 1.26  | 1.23  | 0.79  |
| 5, 5′     | 2.040 | —     | 2.654 | 1.350 | 1.357 | 2.526 |
| r (calc.) | 2.058 | —     | 2.654 | 1.352 | 1.359 | 2.582 |
| WBI       | 0.69  | 0.92  | 1.25  | 1.23  | 0.85  |

2.145 Å and 1.82, respectively, for X = Se. The significantly smaller WBIs in 2′ and 3′ and the longer bond distances (Table 1) reflect a partial contribution of a double bond character in their Ge–X1 bonds. The Ge–X2 bonds, 2.198 Å (exp.), 2.213 Å (calc.), WBI = 0.92 (X = S); 2.326 Å (exp.), 2.348 Å (calc.), WBI = 0.94 (X = Se), are slightly shorter and have somewhat larger WBIs than those calculated in Me3Ge–XH (2.265 Å, WBI = 0.88, X = S; 2.397 Å WBI = 0.91, X = Se). The NCN bonds are 3-centre 4-electron bonds. This is manifested in the resonance structures which involve these electrons, i.e., N≡C:N ≡ N≡N:≡N, and which are reflected in the C–N WBI of ca. 1.25, suggesting a partial double bond character. The Ge–C bond length is 1.998 Å (2.022 Å, WBI = 0.59, calc.) and 1.987 Å (2.020 Å, WBI = 0.60, calc.) in 2 and 3, respectively. The small WBI may reflect the contribution of resonance structures in which the Ge is bound to only one carbene unit (see below). The Ge–C bonds are longer than that in the germyleone precursor B (average r(Ge–C) = 1.963 Å, exp. 1.981 Å, WBI = 1.0.) and of the acyclic, germyleones supported with two CAAC ligands, synthesised by Roesky et al., of 1.940 Å and 1.954 Å.29
The calculated Natural Population Analysis (NPA) charge distribution in compounds 1-5 is shown in Fig. S6 in the ESI.† In all compounds the total charge on the bis-carbene moiety is positive (0.52 el.), indicating that the carbene units are donating electrons to the GeIVX2GaCl3 and GeIIIXGaCl3 fragments. The positive charge on the Ge atom decreases along the series X = S > Se > Te in 2', 3', 4' and 5', while the charge on the GaCl3 fragment increases along this series. Our results indicate that the chalcogen–GaCl3 interaction is strong. In fact, attempts to remove the coordinated GaCl3 from 2-5 by using strong external Lewis bases such as ‘free’ NHC with methyl ligands at nitrogen or less substituted bis-NHCs or chelating bis-thiols had been performed, but no reaction could be observed. At elevated temperature decomposition occurred, leading to hitherto unidentified structures.

Conclusions

Due to the substantial Lewis acid stabilisation of the Ge(0) atom in 1, the Ge(0) atom can be readily oxidised with elemental chalcogens to form, in solvent-dependent reactions, the first donor–acceptor stabilised isolable monomeric germanium disulphide 2, diselenide 3, monoselenide 4, and monotelluride complex 5, respectively. They represent novel classes of heavier congeners of CO and CO₂ complexes. Apparently, the presence of the GaCl₃ Lewis acid is essential for the stabilisation of all monomeric species bearing highly polar Ge≡X bonds. In THF, the germanium[II] monoselenenide complex 4 can further be oxidised with activated selenium to yield the corresponding germanium diselenide complex 3. Unexpectedly, both selenide and telluride compounds 4 and 5 react with elemental sulphur to produce 2 with liberation of elemental selenium and tellurium, respectively. The unusual structural, spectroscopic, and electronic properties of these novel species could be determined and analysed by combined experimental and computational investigations. One of the important lessons from the calculations is that the bonding framework in all these compounds is complex and cannot be described properly by a single valance bond structure. Currently, we continue to explore the synthesis of Lewis acid-free germanium and silicon chalcogenide complexes and to exploit their reactivity in the context of small molecule activation and ligand ability in metal coordination chemistry. The strategy of donor–acceptor stabilisation is also expected to pave the way to isolable monomeric SiO and SiO₂ complexes. Respective studies are currently in progress.

Acknowledgements

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