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## Introduction

Beginning with the inception of transition metal (TM) carbene complexes by E. O. Fischer,<sup>1</sup> research involving the synthesis and utility of metal carbene complexes has long stood as a pillar of modern organometallic chemistry.<sup>2</sup> This has borne numerous classes of carbene complex, a vast number of which have catalytic implications, from highly reactive alkylidene complexes capable of double-bond metathesis,<sup>3</sup> to those involving N-heterocyclic carbene (NHC) spectator ligands.<sup>4</sup> Whilst considerable efforts have also been given to the study of the heavier tetrylenes in TM complexes, this field has largely focused on synthetic access, and less so on reactivity and utilisation in broader synthetic protocols.<sup>5</sup> The heavier tetrylenes have the capacity to act as ‘single centre ambiphiles’, this effect amplified on descending group 14 due to reduced sp-mixing,<sup>6</sup> with a σ-donating lone electron pair and a Lewis acidic vacant p-orbital. Although heavier tetrylenes have been employed as ligands in catalysis,<sup>6,7</sup> and notably so amidinato silylenes,<sup>7</sup> these, like NHCs, are typically spectator ligands due to common examples being N-heterocyclic in nature. Chelating ligands employing an acyclic, low-coordinate germylene have seen some

attention in the literature, and are typically doubly substituted by a ligand bearing a phosphine arm; a number of examples of such systems have been reported by Cabeza and co-workers,<sup>8,9</sup> which have been combined with first-row TM halides so as to access TM<sup>II</sup> complexes, often with tetrylene insertion into TM–X bonds (X = halide).<sup>10</sup> Related (carbonyl-free) first-row TM<sup>0</sup> complexes bearing acyclic, two-coordinate germylene ligands are rare,<sup>11</sup> and should allow for metal–ligand cooperativity (MLC) due to their amphiphilic nature imparting them the potential to remain Lewis acidic when bound to a metal centre. MLC, whereby a metal-bound ligand plays an active role in bond

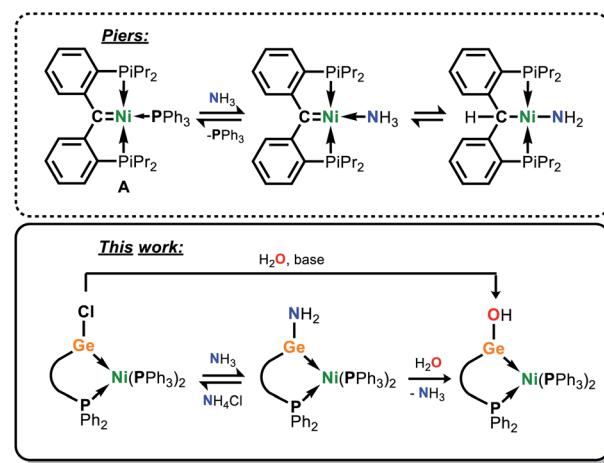


Fig. 1 The nucleophilic role of a carbene ligand in reversible ammonia activation (*Piers*), and the electrophilic role of a germylene ligand in reversible ammonia activation (*this work*).

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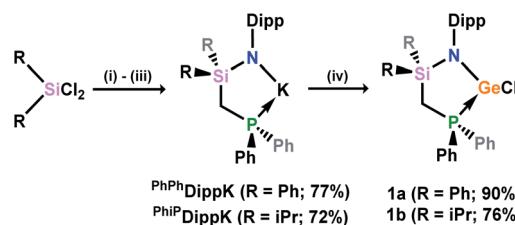


activation,<sup>12</sup> is a powerful concept in catalysis,<sup>13,14</sup> and has allowed for the facile activation of bonds which are otherwise challenging to cleave, a prime example being the N–H bonds in ammonia.<sup>15</sup> In a key demonstration of this process, the phosphino-carbene Ni<sup>0</sup> complex **A** can readily bind ammonia at Ni in the displacement of its Ph<sub>3</sub>P ligand, with the nucleophilic carbene ligand accepting a proton in N–H bond cleavage (Fig. 1).<sup>15c</sup>

Given the potentially Lewis acidic nature of heavier tetrylenes, we envisage that such ligands can reverse the polarity of this MLC process, and act as *electrophilic* binding sites in TM complexes. This would open a new avenue in the dual-centre activation of small molecules, and indeed in catalysis. In order to investigate the capacity of a tetrylene to remain electrophilic in the coordination sphere of a TM, and perhaps more importantly to be more electrophilic than the TM, we sought to develop a chelating ligand incorporating an acyclic, low-coordinate heavier tetrylene. To this end, we have developed novel phosphine-functionalised amine pro-ligands, <sup>PhPh</sup>DippNH and <sup>PhiP</sup>DippNH (<sup>PhPh</sup>DippNH = Ph<sub>2</sub>PCH<sub>2</sub>Si(Ph)<sub>2</sub>N(H)Dipp; <sup>PhiP</sup>DippNH = Ph<sub>2</sub>PCH<sub>2</sub>Si(<sup>i</sup>Pr)<sub>2</sub>N(H)Dipp; Dipp = 2,6-<sup>i</sup>Pr<sub>2</sub>C<sub>6</sub>H<sub>3</sub>), which can be used to generate phosphine-functionalised (amido)(chloro)germylenes which satisfy the targeted acyclic, low-coordinate ligand characteristics when combined with a low-valent TM centre (Fig. 1). Herein we report the synthesis of these compounds, specifically as their Ni<sup>0</sup> complexes, in which the single centre ambiphile ligand centre (*i.e.* Ge<sup>II</sup>) maintains its Lewis acidity, and is capable of binding and activating NH<sub>3</sub> and H<sub>2</sub>O, in the former case reversibly.

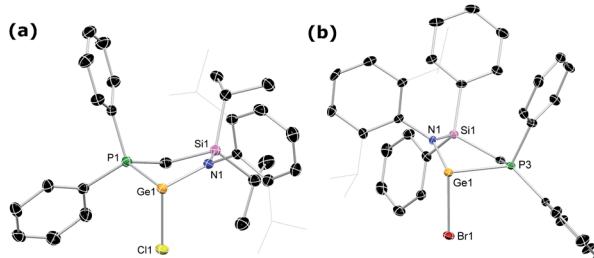
## Results and discussion

The phosphine-functionalised germylene ligands feature a novel ancillary ligand scaffold, namely a phosphine-functionalised amide. The amine pro-ligands are readily accessed through an initial *in situ* synthesis of phosphine-functionalised chlorosilanes, (Ph<sub>2</sub>PCH<sub>2</sub>)(R)<sub>2</sub>SiCl (R = Ph, <sup>i</sup>Pr),<sup>16</sup> which can then be reacted with the lithium anilide, DippN(H)Li, with loss of LiCl. Although the formed pro-ligands, <sup>PhiP</sup>DippNH and <sup>PhPh</sup>DippNH, were not isolated in their pure

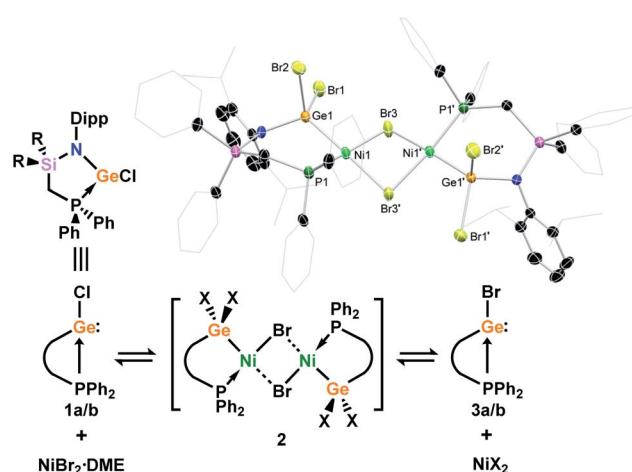


**Scheme 1** The synthesis of phosphine-functionalised amine pro-ligands, their deprotonation, and subsequent synthesis of chloro-germylene complexes. (i) Ph<sub>2</sub>PCH<sub>2</sub>Li·TMEDA, hexane; (ii) DippN(H)Li, THF; (iii) KH, THF; (iv) GeCl<sub>2</sub>·dioxane, THF (yields in parentheses). Dipp = 2,6-<sup>i</sup>Pr<sub>2</sub>C<sub>6</sub>H<sub>3</sub>.

forms, <sup>1</sup>H and <sup>31</sup>P NMR analyses of crude reaction mixtures suggest they are formed near quantitatively.<sup>17</sup> These crude products are readily deprotonated with a suspension of KH in THF, yielding the potassium amides <sup>PhiP</sup>DippNK and <sup>PhPh</sup>DippNK in good isolated yields of ~70 to 75% after work up, based on the Ph<sub>2</sub>PCH<sub>2</sub>Li·TMEDA starting material. Ge<sup>II</sup> chloride complexes <sup>PhPh</sup>DippNGeCl and <sup>PhiP</sup>DippNGeCl (**1a** and **1b**, respectively; (Scheme 1)) are generated through combination of these potassium amides with GeCl<sub>2</sub>·dioxane in either THF or toluene, yielding the desired germylene ligands in high yields. Both **1a** and **1b** show somewhat complex <sup>1</sup>H NMR spectra, when compared with potassium amides <sup>PhiP</sup>DippNK and <sup>PhPh</sup>DippNK, due to the asymmetric coordination at their Ge<sup>II</sup> centres. Nevertheless, the presence of a single peak in the respective <sup>31</sup>P {<sup>1</sup>H} NMR spectra confirms the presence of a single ligand environment in these compounds. A single crystal X-ray diffraction analysis of the two germylenes confirms their monomeric nature (*viz.* Fig. 2), and shows that the Ph<sub>2</sub>P moiety of the amide ligands binds the Ge<sup>II</sup> centre in both species (**1a**: *d*<sub>PGe</sub> = 2.4547(9) Å; **1b**: *d*<sub>PGe</sub> = 2.458(1) Å). Complexes **1a/b** bear some similarity to complexes reported by Wesemann, incorporating bulky aryl ligands at Ge<sup>II</sup>,<sup>18</sup> and Baceiredo, who also developed Ge<sup>II</sup> chloride species bearing phosphine-



**Fig. 2** The molecular structures of (a) **1b** and (b) **3a**, with thermal ellipsoids at 40% probability. Hydrogen atoms omitted for clarity. Selected bond distances (Å) and angles (°) for **1b**: Ge1–Cl1 2.355(1); Ge1–P1 2.472(1); Ge1–N1 1.925(3); N1–Ge1–Cl1 100.69(1); Cl1–Ge1–P1 85.98(4); N1–Ge1–P1 85.85(1). For **3a**: Ge1–Br1 2.512(1); Ge1–P3 2.455(2); Ge1–N1 1.913(4); N1–Ge1–Br1 103.00(1); Br1–Ge1–P3 84.74(4); N1–Ge1–P3 84.98(1).



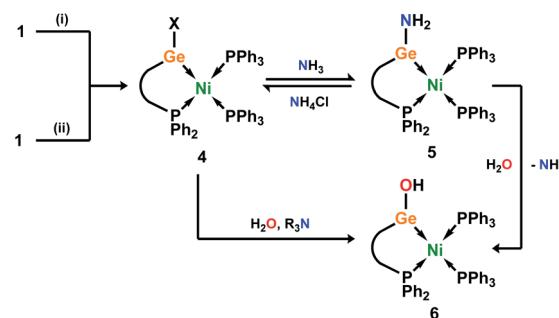
**Scheme 2** The reaction of chloro-germylenes **1a/b** with NiBr<sub>2</sub>·DME, leading to reversible complexation and Cl/Br exchange at Ge<sup>II</sup>. Inset: the molecular structure of dimeric germyl-nickel complex **2**.



functionalised amide ligands.<sup>19</sup> In these cases, Ge–P distances are similar to those in **1a/b**.

With (amido)(chloro)germylene ligands **1a/b** in hand, we began our investigations into the complexation chemistry of these pro-chelating ligands towards nickel dihalides. We found that both complexes do not readily react with  $\text{NiCl}_2$ , even after heating, despite a dark red colouration of reaction solutions. However, reactions with an excess of  $\text{NiBr}_2 \cdot \text{DME}$  in toluene with a small amount of THF led to dark red-brown reaction mixtures, from which dimeric complex **2** could be isolated as large red-brown crystals after filtration (Scheme 2). X-ray structural analysis of these crystals (Scheme 2, inset) revealed that Ni inserts into the Ge–P bond of **1a**, forming a chelating ligand motif incorporating P and Ge as we had hoped. Formal oxidative addition of one Ni–Br bond at  $\text{Ge}^{\text{II}}$  is also observed, akin to previous examples reported by Cabeza and co-workers.<sup>10</sup> Each  $\text{Ni}^{\text{II}}$  centre sits in a square planar geometry, bound by two bridging Br ligands, Ge, and P. Compound **2** represents a surprisingly uncommon example of a germyl-nickel complex, and is unstable in solution in the absence of an excess of  $\text{NiBr}_2$ . This was clear in an attempt to obtain NMR spectroscopic data for **2**, where the only observable species is the bromo germylene  $\text{Ph}_3\text{P}^{\text{Ph}}\text{DippGeBr}$  **3a**, presumably through elimination of the nickel dihalide.<sup>20</sup> Reassessing the reaction of chloro germylene **1a** with  $\text{NiBr}_2 \cdot \text{DME}$ , we found that addition of a single equivalent leads to a  $\sim 50 : 50$  mixture of **1** and **3**, *i.e.* partial Cl/Br exchange (Fig. S24 and S28†). From such a reaction mixture the formation of one or two single crystals of the germyl-nickel mixed halide complex **2'**, the acetonitrile-coordinated monomeric form of **2**, allowed for structural analysis of this ‘intermediate’ (Fig. S80 in ESI†). Addition of 6 equiv. of  $\text{NiBr}_2 \cdot \text{DME}$  allows for full conversion to bromo germylene **3**,<sup>21</sup> whilst a vast excess ( $\sim 10$  equiv.) of  $\text{NiBr}_2 \cdot \text{DME}$  affects the crystallisation of small amounts of germyl-nickel bromide complex **2**. We presume that the presence of an excess of  $\text{NiBr}_2 \cdot \text{DME}$  is required to maintain the stability of **2**, as in all cases NMR analyses showed only the free chloro/bromo-germylene ligands. Taken as a whole, this demonstrates that  $\text{NiBr}_2$  can act as a facile halide exchange reagent for the synthesis of bromo germylenes, which are challenging to access *via* conventional routes due to the lack of readily available  $\text{GeBr}_2$  reagents.<sup>22</sup>

Although the isolation of useful quantities of nickel complex **2** was not possible, the observation that it exists within the reaction mixture seemed a promising start. We found that *in situ* reduction of this reaction mixture, in the presence of  $\text{Ph}_3\text{P}$ , gave facile access to chelating phosphino-germylene complexes of  $\text{Ni}^0$ . Room temperature addition of THF to a mixture of **1a/b**,  $\text{NiBr}_2 \cdot \text{DME}$ ,  $\text{Ph}_3\text{P}$ , and an excess of Zn powder allowed for the clean formation of  $\text{Ni}^0$  complexes **4a/b** after 2 h, albeit as a mixture of the chloro- and bromo-germylene ligated species. Employing  $\text{NiCl}_2 \cdot \text{DME}$  in place of  $\text{NiBr}_2 \cdot \text{DME}$  led to considerably extended reaction times, but gave clean access to the chloro-germylene species (Scheme 3). The pure chloro-germylene analogue can also be accessed through combination of **1**,  $\text{Ni}(\text{COD})_2$ , and  $\text{PPh}_3$  in the stoichiometric ratio 1 : 1 : 2 (Fig. S31†). Satisfyingly,  $^{31}\text{P}$  NMR analysis of crude reaction mixtures in all cases suggest the formation of these novel  $\text{Ni}^0$



Scheme 3 Synthesis of halo-germylene  $\text{Ni}^0$  complexes **4** and subsequent reactions with  $\text{NH}_3$  and  $\text{H}_2\text{O}$ . (i)  $1.1\text{NiX}_2 \cdot \text{DME}$ ,  $2\text{Ph}_3\text{P}$ ,  $6\text{Zn}$ , THF, 2 h ( $\text{X} = \text{Br}$ );  $\sim 24$  h ( $\text{X} = \text{Cl}$ ); (ii)  $1\text{Ni}(\text{cod})_2$ ,  $2\text{Ph}_3\text{P}$ , toluene, 1 h;  $\text{X} = \text{Cl}$ .

complexes as the sole products, with the crystalline compounds isolated in good yields from diethyl ether extracts of the crude reaction mixtures. A single crystal X-ray structural analysis of dark red-brown crystals of **4b** reveal a central  $\text{Ni}^0$ , bound by two  $\text{Ph}_3\text{P}$  ligands, and one chelating phosphino-germylene ligand (Fig. 4(a)). Notably, the chloride moiety in the germylene ligand remains intact. The three-coordinate  $\text{Ge}^{\text{II}}$  centre holds a trigonal planar geometry, and has a relatively short Ge–Ni contact when compared with previously reported examples, indicative of some back-bonding from Ni to Ge (*vide supra*). The  $\text{Ni}^0$  centre holds a tetrahedral geometry, with the three Ni–P distances being as expected when compared with reported phosphine-coordinated  $\text{Ni}^0$  complexes. Complex **4a** is essentially isostructural to **4b**.<sup>23</sup> The UV/vis spectra of these species show two clear absorption bands (**4a**:  $\lambda_{\text{max}} = 484 \text{ nm}$  ( $\epsilon = 1370 \text{ L cm}^{-1} \text{ mol}^{-1}$ ) and  $362 \text{ nm}$  ( $6735 \text{ L cm}^{-1} \text{ mol}^{-1}$ ); **4b**:  $473 \text{ nm}$  ( $1685 \text{ L cm}^{-1} \text{ mol}^{-1}$ ) and  $362 \text{ nm}$  ( $9160 \text{ L cm}^{-1} \text{ mol}^{-1}$ )), in-keeping with related absorptions for Ni-based d–d transitions.<sup>24</sup> Likely due to hindered rotation brought about by the sterics of the chelating ligand system, resonances relating the ligand are broadened in the  $^1\text{H}$  NMR spectrum of both  $\text{Ni}^0$

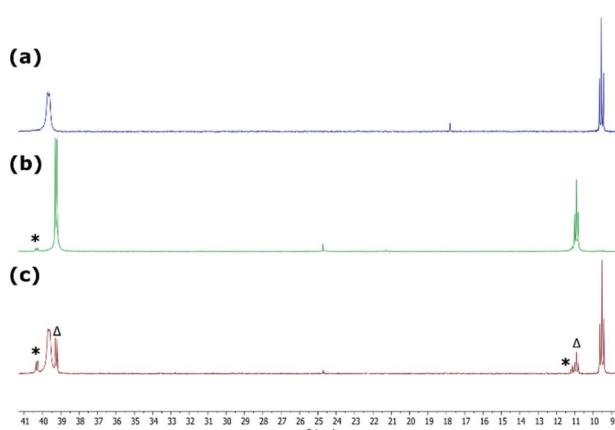


Fig. 3 The  $^{31}\text{P}\{^1\text{H}\}$  NMR spectrum of (a) **4b**, (b) *in situ* addition of  $\text{NH}_3$  to **5b**, and (c) **4b** regenerated after sonication and degassing the sample used for spectrum (b). \* = small amounts of **6b**; Δ = small amounts of residual **5b**.



species. The  $^{31}\text{P}$  NMR spectra show one broadened peak for the  $\text{Ph}_3\text{P}$  ligands, centred at  $\delta = 39.8$  (**4a** and **4b**) ppm, whilst the flanking phosphine arm of the chelating ligand presents as a well resolved triplet centred at  $\delta = 7.1$  (**4a**) and 9.6 (**4b**) ppm (Fig. 3(a)). For the mixed bromide/chloride samples, a second set of peaks relating to the bromide species is observed in the  $^{31}\text{P}$  NMR spectrum, overlapping with the first (Fig. S32 and S40 $\dagger$ ).<sup>25</sup> Separation of the two compounds proved impossible in our hands, although using the mixture in subsequent chemistry discussed herein did not pose any issues.

Given the ambiphilic nature of tetrylenes, particularly in acyclic derivatives, the design of complexes **4a/b** aims to retain a degree of Lewis acidity at  $\text{Ge}^{\text{II}}$ . This forms the central idea of non-innocent single centre ambiphile ligands. This characteristic was probed by Density Functional Theoretical (DFT) analysis of the frontier orbitals in model complex **4'**, employing  $^{\text{PhMe}}\text{XylN}$  in place of  $^{\text{PhiP}}\text{DippN}/^{\text{PhPh}}\text{Dipp}$  ( $^{\text{PhMe}}\text{XylN} = (\text{Ph}_2\text{PCH}_2\text{SiMe}_2)(\text{Xyl})\text{N}$ ). We found that the LUMO of **4'** is located on the Ge centre (Fig. 4(a)), and mainly constitutes a vacant p-orbital which is expected to be Lewis acidic, particularly given the overall NPA charge at Ge of +1.13. Still, Natural Bond Orbital (NBO) analysis of the Ge–Ni bond indicates strong polarisation toward the Ge centre (Table S4 $\dagger$ ), whilst the HOMO in **4'** (Fig. S85 in ESI $\dagger$ ) shows some degree of  $\pi$ -bonding, pointing towards a donor–acceptor description of the Ge–Ni bond. In line with the Lewis acidity of the  $\text{Ge}^{\text{II}}$  centre in **4a/b**, deep red-brown solutions of these compounds readily react with ammonia at 1 atm pressure to form slightly turbid bright orange reaction mixtures containing ammonia activation products **5a/b**. *In situ*  $^{31}\text{P}\{^1\text{H}\}$  NMR spectroscopic analysis indicated the formation of a single new product for both systems (Fig. 3(b)), whilst

complimentary  $^1\text{H}$  NMR spectra indicated new singlet 2H resonances at  $\delta = 3.13$  (**5a**) and 2.96 (**5b**) ppm, which were tentatively assigned to an  $\text{NH}_2$  fragment. The IR spectrum of reaction products is in keeping with this, with two weak N–H stretching bands observed at  $\nu = 3354$  and 3461 (**5a** and **5b**)  $\text{cm}^{-1}$ . Structural analysis of large orange crystals of the product of the reaction of **4a** with ammonia reveal that the Ge–Cl moiety in this species has in fact undergone a  $\sigma$ -metathesis reaction with one N–H bond of ammonia, yielding the bis(amido)germylene  $\text{Ni}^0$  complex **5a** (Scheme 3 and Fig. 4(b)).<sup>26</sup> Presumably, the colourless solid in reaction mixtures is ammonium chloride, through the net loss of HCl in this reaction (*vide supra*). As such, removal of the ammonia atmosphere from reaction mixtures, and sonication of amides **5a/b** over the precipitated  $\text{NH}_4\text{Cl}$  with intermittent degassing leads to regeneration of starting materials **4a/b** (Fig. 3), with the loss of ammonia (Scheme 3).<sup>27</sup> Such a reversible activation of ammonia is, to the best of our knowledge extremely rare, as is the more general redox-innocent metathesis reaction at a tetrylene centre. Beyond the reaction of **4a/b** with ammonia, these species also readily undergo a similar metathesis reaction with water in the presence of nitrogen bases to facilitate HCl abstraction (*i.e.*  $\text{CyNH}_2$ ), to yield (amido)(hydroxyl)germylene  $\text{Ni}^0$  complexes **6a/b** (Scheme 3 and Fig. 4(c)). These species can also be accessed by the reaction of amide complexes **5a/b** with water, with concomitant loss of ammonia. The formation of **6a/b** can be clearly observed by  $^1\text{H}$  NMR and IR spectroscopy, the former containing new 1H resonances pertaining to OH residues ( $\delta = 3.13$  (**6a**) and 2.96 (**6b**) ppm), and the latter clear OH stretching bands ( $\nu = 3547$  (**6a**) and 3541 (**6b**)  $\text{cm}^{-1}$ ). In recent years a flurry of examples of ammonia activation at low-valent group 14

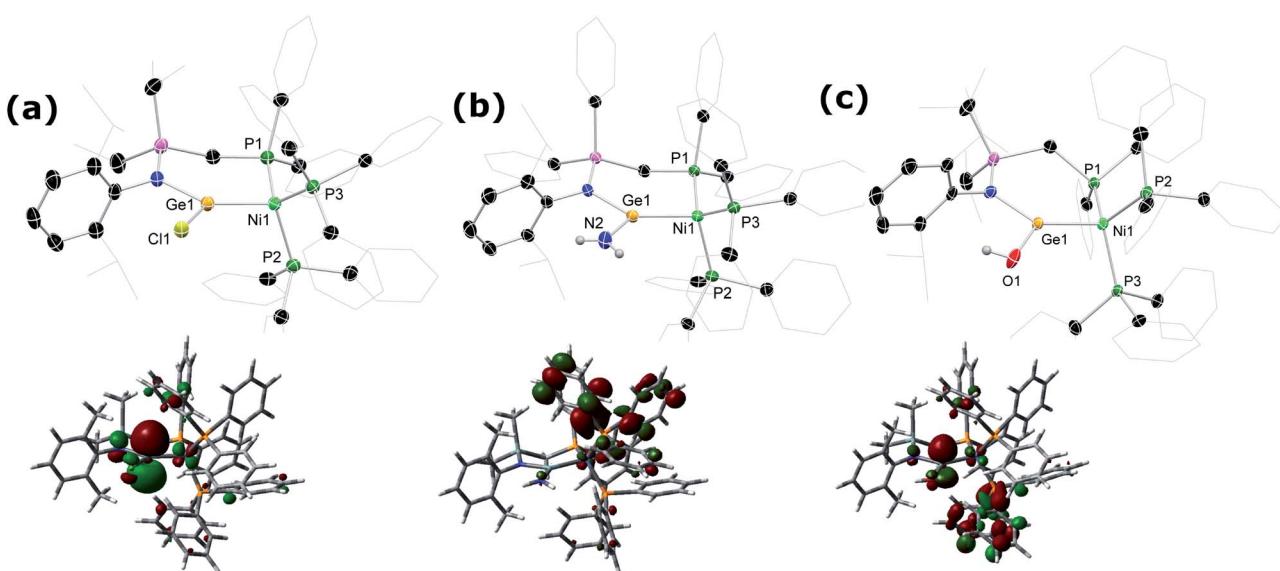


Fig. 4 Molecular structures of compounds (a) **4b**, (b) **5a**, and (c) **6b**, with thermal ellipsoids at 40% probability. Hydrogen atoms omitted for clarity, aside from those at N2 and O1 in **5a** and **6b**, respectively. The LUMO of each compound is inset below the respective structure. Selected bond distances ( $\text{\AA}$ ) and angles ( $^\circ$ ) for **4b**: Ge1–Ni1 2.1877(7); N1–Ge1 1.869(2); P1–Ni1 2.201(1); P2–Ni1 2.2079(8); P3–Ni1 2.2055(8); N1–Ge1–Cl1 99.57(7); Ni1–Ge1–N1 133.09(7); Ni1–Ge1–Cl1 126.89(3). For **5a**: Ge1–Ni1 2.217(1); N1–Ge1 1.890(2); N2–Ge1 1.819(2); P1–Ni1 2.210(1); P2–Ni1 2.201(1); P3–Ni1 2.1892(9); N1–Ge1–N2 99.07(9); Ni1–Ge1–N1 128.63(6); Ni1–Ge1–N2 132.29(7). For **6b**: Ge1–Ni1 2.2077(7); N1–Ge1 1.885(3); O1–Ge1 1.874(2); P1–Ni1 2.210(1); P2–Ni1 2.202(1); P3–Ni1 2.104(1); N1–Ge1–O1 100.98(1); Ni1–Ge1–O1 128.47(7); Ni1–Ge1–N1 130.49(8).



centres have been forthcoming, but lead almost exclusively to oxidative addition reactions, thus yielding  $E^{IV}$  compounds ( $E = C-Sn$ ).<sup>28</sup> That oxidative addition is circumvented in reactions of **4a/b** with both  $NH_3$  and  $H_2O$  likely stems from the binding of the Ge lone electron pair to  $Ni^0$ . Further, it is surprising that the  $Ni^0$  centre in these complexes remains unchanged, given previous examples of MLC in ammonia activation involving  $Ph_3P$ -ligated  $Ni^0$  complexes.<sup>15c,d</sup> Overall, the metathesis reactions of ammonia demonstrated here highlight the targeted high Lewis acidity of the developed acyclic germylene ligands, and are an exciting step towards employing these ligands in complexes which can operate *via* MLC bond activation processes, where a lower coordinate TM centre is utilised.

So as to compare the effects of the differing fragments at Ge<sup>II</sup> in the described  $Ni^0$  complexes, we performed frontier orbital, NBO, Wiberg Bond Index (WBI), Mayer Bond Order (MBO), and Natural Population analyses on **4'**, **5'**, and **6'** (Tables S4–S7†), as model complexes of the real systems, again employing  $^{PhMe}XylN$  in place of  $^{PhP}DippN/^{PhPh}Dipp$ . We found that **4'**, **5'**, and **6'** all show very similar characteristics in general; the main difference is the interaction of the vacant p-orbital on the Ge<sup>II</sup> centre and the lone pair of the Cl/NH<sub>2</sub>/OH moiety. Bond analysis showed the highest bond order for the Ge–NH<sub>2</sub> in **5'** (MBO: 1.10) which is due to the efficient hybridization of the N lone pair and the vacant p-orbital of Ge that are well-described for low-valent germanium compounds.<sup>29</sup> The LUMO of **5'** is the only example in this series which is not represented by a vacant p-orbital at Ge (Fig. 4), most likely due to the described N → Ge donation. Consequently, the Ge–Ni bond order decreases and the Ge–Ni bond length increases (*vide infra*) due to the smaller back donation from the Ni d-orbital to the Ge vacant orbital. All systems show some degree of multiple bond character between the Ge<sup>II</sup> and Ni<sup>0</sup> centres (WBI/MBO for **4'** 1.20/1.13; for **5'**: 1.11/

0.98; for **6'**: 1.17/1.07), with predicted Ge–Ni distances (**4'**: 2.161 Å; **5'**: 2.203 Å; **6'**: 2.177 Å) only slightly contracted relative to those observed experimentally (*e.g.*  $d_{GeNi}$  for **4b**: 2.1877(7) Å; **5a**: 2.217(1) Å; **6a**: 2.1956(7) Å), possibly due to the increased steric encumbrance in the real complexes. An NBO analysis indicates a large degree of s-character in these bonds, indicative of the greatest contributing factor being Ge → Ni lone-pair donation, but, as already described for **4'**, a high degree of polarisation towards Ge is also apparent, making a donor–acceptor interaction the best description for the Ge–Ni interaction in these complexes.

Further, the mechanism for the amination of model complex **4'** was investigated by means of DFT (Fig. 5), given the reversible nature of this interesting process. The most favourable mechanism begins with binding of  $NH_3$  at Ge<sup>II</sup>, highlighting the Lewis acidity of this centre. This process is near thermoneutral (**IM1** in Fig. 5, +1.7 kcal mol<sup>-1</sup>), which corroborates reversibility in this key step. Surprisingly, it was found that two further  $NH_3$  molecules are subsequently required to drive the metathesis, which form a H-bond network between the Ge-bound  $NH_3$  and Cl ligands. The most challenging step in the overall process is the ion pair formation (**TS4** in Fig. 5, +14.5 kcal mol<sup>-1</sup>), which is only possible on the potential energy surface (PES) if the H-bond network is present, which stabilises the formation of the  $NH_4^+$  ion. Then, the  $NH_4^+$  and the additional  $NH_3$  molecule facilitate the removal of  $Cl^-$ . The overall reaction is very close to thermoneutral (**5'** in Fig. 5, -0.9 kcal mol<sup>-1</sup>), making this process reversible simply by changing the reaction conditions, again corroborating the observed experimental results. Finally, we also considered (i) the potential involvement of the Ni centre in  $NH_3$  activation, and (ii) oxidative addition of an N–H bond at the Ge centre. Both of these mechanisms can be excluded from the active PES; the Ni–H complex, formed in (i), is relatively

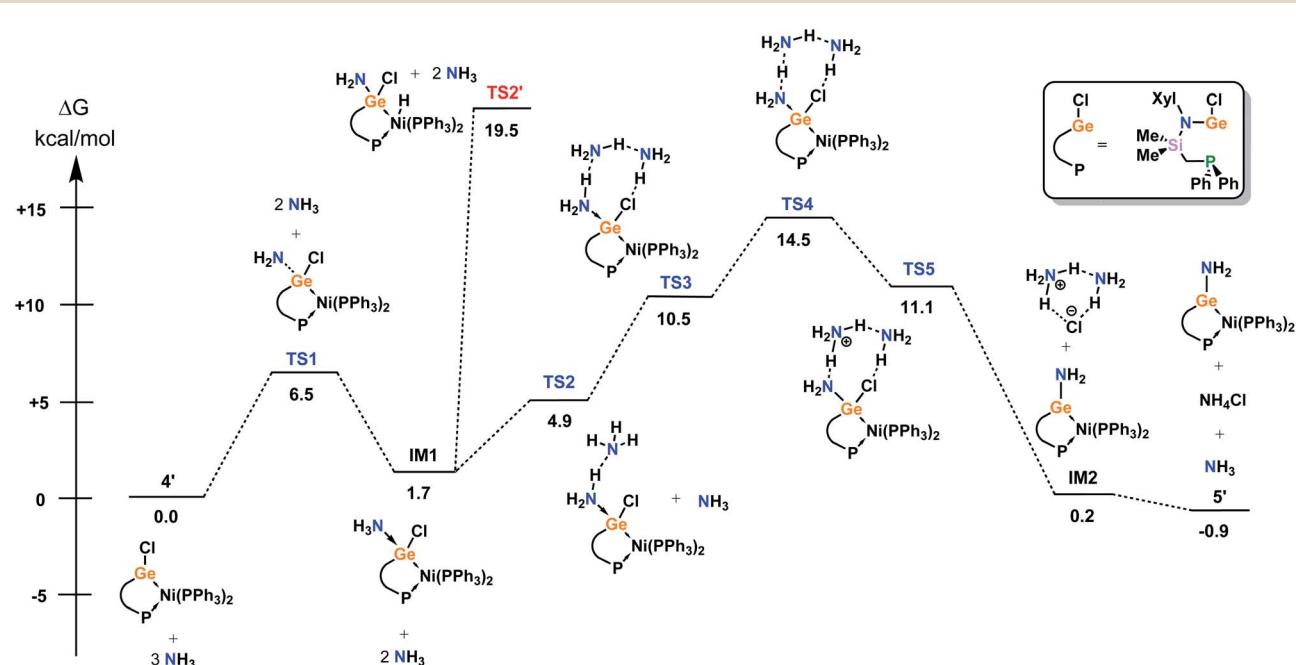


Fig. 5 DFT-derived mechanism for the  $\sigma$ -metathesis of ammonia in **4'** leading to **5'**.



unstable (TS2' in Fig. 5, 19.5 kcal mol<sup>-1</sup>), whilst the Ge oxidative addition product is not a minimum on the PES (route (ii)).

## Conclusions

In conclusion, we have developed a chelating phosphino-germylene ligand scaffold, and a facile route to Ni<sup>0</sup> complexes bearing these ligands in a low-cost, one pot synthetic preparation. These ligands, which are centred around an acyclic (amido)(chloro)germylene, remain Lewis acidic when bound to Ni<sup>0</sup>. This has been demonstrated by the facile and reversible activation of ammonia, as well as the complimentary irreversible reaction with water, both of which lead to Ge<sup>II</sup> products through  $\sigma$ -metathesis of the Ge–Cl bond, thus circumventing oxidation at both germanium and nickel. These results form an initial basis for the single centre ambiphile ligand concept, which we are currently expanding to further low-valent main group species, and further first row transition metals, moving towards cooperative bond activation involving both the ligand and the metal centre.

## Experimental

### General considerations

All experiments and manipulations were carried out under a dry oxygen free argon atmosphere using standard Schlenk techniques, or in a MBraun inert atmosphere glovebox containing an atmosphere of high purity argon. THF and diethyl ether were dried by distillation over a sodium/benzophenone mixture and stored over activated 4 Å mol sieves. C<sub>6</sub>D<sub>6</sub> was dried and stored over a potassium mirror. All other solvents were dried over activated 4 Å mol sieves and degassed prior to use. NiBr<sub>2</sub>·DME,<sup>30</sup> NiCl<sub>2</sub>·DME,<sup>31</sup> DippN(H)Li,<sup>32</sup> and PPh<sub>2</sub>CH<sub>2</sub>Li·TMEDA<sup>33</sup> were synthesized according to known literature procedures. All other reagents were used as received. Commercial CyNH<sub>2</sub>, when used as received, contained enough residual moisture to allow for the synthesis of **6a**. NMR spectra were recorded on a Bruker AV 400 or 500 Spectrometer. The <sup>1</sup>H and <sup>13</sup>C{<sup>1</sup>H} NMR spectra were referenced to the residual solvent signals as internal standards. <sup>29</sup>Si NMR spectra were externally calibrated with SiMe<sub>4</sub>. <sup>31</sup>P NMR spectra were externally calibrated with H<sub>3</sub>PO<sub>4</sub>. LIFDI MS spectra were measured at a Waters Micromass LCT TOF mass spectrometer equipped with an LIFDI ion source (LIFDI 700) from Linden CMS GmbH. The samples were dissolved in dry toluene and filtered using a syringe filter under an inert atmosphere. The TOF setup was externally calibrated using polystyrene. ESI-MS was performed on an exactive plus orbitrap spectrometer from Thermo Fischer Scientific. Infrared spectra were measured with the Alpha FT IR from Bruker containing a platinum diamond ATR device. The compounds were measured as solids under inert conditions in a glovebox. For the ammonia activation experiments water free ammonia 5.0 was used.

Representative procedures for the preparation of compounds **1**, **3**, and **4–6** are given below. Further details for the synthesis and characterisation of all novel compounds are given in the ESI.<sup>†</sup>

### <sup>PhPh</sup>DippNK

A yellow suspension of PPh<sub>2</sub>CH<sub>2</sub>Li·TMEDA (6.0 g, 18.6 mmol) in 100 mL hexane was cooled to -78 °C. The mixture was stirred vigorously and Ph<sub>2</sub>SiCl<sub>2</sub> (3.9 mL, 18.6 mmol) was added. The mixture was allowed to warm to RT overnight. All volatiles were subsequently removed *in vacuo*, leaving a yellow oil. DippN(H)Li (3.4 g, 18.6 mmol) was added to the residue, and the flask cooled to -78 °C, followed by the addition of 50 mL THF. The mixture was stirred until dissolution of all solids was observed. The cold bath was then removed and the reaction allowed to warm to RT, leading to an orange solution. All volatiles were removed *in vacuo* and the oily residue extracted with 50 mL hexane, and filtered. The solvent was removed *in vacuo* and KH (0.9 g, 28.3 mmol) was added. After addition of 50 mL THF, gas started to evolve, and the mixture was vigorously stirred for a further 16 h. The dark brown suspension was filtered, and all volatiles were removed *in vacuo*. To the resulting oil 50 mL hexane was added, and the mixture treated in an ultrasonic bath causing the precipitation of copious pale brown powder, which was filtered and washed multiple times with hexane, and subsequently dried *in vacuo* to yield <sup>PhPh</sup>DippNK as an off-white powder (8.5 g, 14.3 mmol, 77%). Colourless crystals suitable for X-ray diffraction analysis were obtained after two days from a concentrated THF/TMEDA solution layered with hexane stored at -32 °C. <sup>1</sup>H NMR (THF-d<sub>8</sub>, 400 MHz, 298 K):  $\delta$  = 0.91 (d, 12H, <sup>3</sup>J<sub>HH</sub> = 6.9 Hz, Dipp-Pr<sup>i</sup>-CH<sub>3</sub>), 1.97 (d, 2H, <sup>2</sup>J<sub>HP</sub> = 5.4 Hz, Ph<sub>2</sub>P-CH<sub>2</sub>), 3.94 (hept, 2H, <sup>3</sup>J<sub>HH</sub> = 6.9 Hz, Dipp-Pr<sup>i</sup>-CH), 6.24 (t, 1H, <sup>3</sup>J<sub>HH</sub> = 7.4 Hz, Ar-CH), 6.74 (d, 2H, <sup>3</sup>J<sub>HH</sub> = 7.4 Hz, Ar-CH), 7.00 (m, 6H, Ar-CH), 7.10 (m, 6H, Ar-CH), 7.27 (m, 4H, Ar-CH), 7.48 (m, 4H, Ar-CH). <sup>13</sup>C{<sup>1</sup>H} NMR (THF-d<sub>8</sub>, 101 MHz, 298 K):  $\delta$  = 19.7 (d, <sup>1</sup>J<sub>CP</sub> = 29.2 Hz, Ph<sub>2</sub>P-CH<sub>2</sub>), 24.9 (Dipp-Pr<sup>i</sup>-CH<sub>3</sub>), 28.1 (Dipp-Pr<sup>i</sup>-CH), 112.6, 123.00, 127.3, 127.5, 128.3, 128.7, 133.4, 133.6, 135.8, 141.4, 143.9, 144.0, 146.7, 146.7 and 156.2 (Ar-C). <sup>31</sup>P{<sup>1</sup>H} NMR (THF-d<sub>8</sub>, 162 MHz, 298 K):  $\delta$  = -19.1 (s, CH<sub>2</sub>-PPh<sub>2</sub>). <sup>29</sup>Si{<sup>1</sup>H} NMR (THF-d<sub>8</sub>, 99 MHz, 298 K):  $\delta$  = -47.4 (d, <sup>2</sup>J<sub>SiP</sub> = 14.7 Hz, SiPh<sub>2</sub>).

### <sup>PhPh</sup>DippGeCl, **1a**

A pale brown solution of <sup>PhPh</sup>DippNK (7.0 g, 11.7 mmol) in 20 mL THF was added dropwise to a stirring solution of GeCl<sub>2</sub>·dioxane (2.7 g, 11.7 mmol) in 10 mL THF at -78 °C, and subsequently allowed to warm to RT, resulting in the formation of an orange solution. All volatiles were removed *in vacuo*, the residue extracted in 20 mL DCM, and filtered. The solvent was removed *in vacuo* and the residue washed with hexane to yield **1a** as an off-white powder (7.0 g, 10.5 mmol, 90%). Colourless crystals suitable for X-ray diffraction analysis were obtained from a concentrated diethyl ether at RT. <sup>1</sup>H NMR (C<sub>6</sub>D<sub>6</sub>, 400 MHz, 298 K):  $\delta$  = 0.09 (d, 3H, <sup>3</sup>J<sub>HH</sub> = 6.4 Hz, Dipp-Pr<sup>i</sup>-CH<sub>3</sub>), 0.68 (d, 3H, <sup>3</sup>J<sub>HH</sub> = 6.6 Hz, Dipp-Pr<sup>i</sup>-CH<sub>3</sub>), 0.86 (d, 3H, <sup>3</sup>J<sub>HH</sub> = 6.5 Hz, Dipp-Pr<sup>i</sup>-CH<sub>3</sub>), 1.43 (d, 3H, <sup>3</sup>J<sub>HH</sub> = 6.5 Hz, Dipp-Pr<sup>i</sup>-CH<sub>3</sub>), 2.54 (m, 3H, Ph<sub>2</sub>P-CH<sub>2</sub>/Dipp-Pr<sup>i</sup>-CH), 4.28 (m, 1H, Dipp-Pr<sup>i</sup>-CH), 7.02 (m, 14H, Ar-CH), 7.21 (m, 3H, Ar-CH), 7.46 (m, 2H, Ar-CH), 7.65 (m, 2H, Ar-CH), 8.26 (d, 2H, <sup>3</sup>J<sub>HH</sub> = 7.1 Hz, Ar-CH). <sup>13</sup>C{<sup>1</sup>H} NMR (C<sub>6</sub>D<sub>6</sub>, 101 MHz, 298 K):  $\delta$  = 8.1 (Ph<sub>2</sub>P-CH<sub>2</sub>), 21.4, 22.6 and 28.0 (Dipp-Pr<sup>i</sup>-CH<sub>3</sub>), 28.2 and 28.6 (Dipp-Pr<sup>i</sup>-CH), 28.8 (Dipp-Pr<sup>i</sup>-CH<sub>2</sub>), 29.2 (Dipp-Pr<sup>i</sup>-CH), 30.2 (Dipp-Pr<sup>i</sup>-CH<sub>3</sub>), 47.4 (SiPh<sub>2</sub>), 50.2 (SiPh<sub>3</sub>), 51.2 (SiPh<sub>2</sub>), 52.2 (SiPh<sub>2</sub>), 53.2 (SiPh<sub>2</sub>), 54.2 (SiPh<sub>2</sub>), 55.2 (SiPh<sub>2</sub>), 56.2 (SiPh<sub>2</sub>), 57.2 (SiPh<sub>2</sub>), 58.2 (SiPh<sub>2</sub>), 59.2 (SiPh<sub>2</sub>), 60.2 (SiPh<sub>2</sub>), 61.2 (SiPh<sub>2</sub>), 62.2 (SiPh<sub>2</sub>), 63.2 (SiPh<sub>2</sub>), 64.2 (SiPh<sub>2</sub>), 65.2 (SiPh<sub>2</sub>), 66.2 (SiPh<sub>2</sub>), 67.2 (SiPh<sub>2</sub>), 68.2 (SiPh<sub>2</sub>), 69.2 (SiPh<sub>2</sub>), 70.2 (SiPh<sub>2</sub>), 71.2 (SiPh<sub>2</sub>), 72.2 (SiPh<sub>2</sub>), 73.2 (SiPh<sub>2</sub>), 74.2 (SiPh<sub>2</sub>), 75.2 (SiPh<sub>2</sub>), 76.2 (SiPh<sub>2</sub>), 77.2 (SiPh<sub>2</sub>), 78.2 (SiPh<sub>2</sub>), 79.2 (SiPh<sub>2</sub>), 80.2 (SiPh<sub>2</sub>), 81.2 (SiPh<sub>2</sub>), 82.2 (SiPh<sub>2</sub>), 83.2 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(SiPh<sub>2</sub>), 427.2 (SiPh<sub>2</sub>), 428.2 (SiPh<sub>2</sub>), 429.2 (SiPh<sub>2</sub>), 430.2 (SiPh<sub>2</sub>), 431.2 (SiPh<sub>2</sub>), 432.2 (SiPh<sub>2</sub>), 433.2 (SiPh<sub>2</sub>), 434.2 (SiPh<sub>2</sub>), 435.2 (SiPh<sub>2</sub>), 436.2 (SiPh<sub>2</sub>), 437.2 (SiPh<sub>2</sub>), 438.2 (SiPh<sub>2</sub>), 439.2 (SiPh<sub>2</sub>), 440.2 (SiPh<sub>2</sub>), 441.2 (SiPh<sub>2</sub>), 442.2 (SiPh<sub>2</sub>), 443.2 (SiPh<sub>2</sub>), 444.2 (SiPh<sub>2</sub>), 445.2 (SiPh<sub>2</sub>), 446.2 (SiPh<sub>2</sub>), 447.2 (SiPh<sub>2</sub>), 448.2 (SiPh<sub>2</sub>), 449.2 (SiPh<sub>2</sub>), 450.2 (SiPh<sub>2</sub>), 451.2 (SiPh<sub>2</sub>), 452.2 (SiPh<sub>2</sub>), 453.2 (SiPh<sub>2</sub>), 454.2 (SiPh<sub>2</sub>), 455.2 (SiPh<sub>2</sub>), 456.2 (SiPh<sub>2</sub>), 457.2 (SiPh<sub>2</sub>), 458.2 (SiPh<sub>2</sub>), 459.2 (SiPh<sub>2</sub>), 460.2 (SiPh<sub>2</sub>), 461.2 (SiPh<sub>2</sub>), 462.2 (SiPh<sub>2</sub>), 463.2 (SiPh<sub>2</sub>), 464.2 (SiPh<sub>2</sub>), 465.2 (SiPh<sub>2</sub>), 466.2 (SiPh<sub>2</sub>), 467.2 (SiPh<sub>2</sub>), 468.2 (SiPh<sub>2</sub>), 469.2 (SiPh<sub>2</sub>), 470.2 (SiPh<sub>2</sub>), 471.2 (SiPh<sub>2</sub>), 472.2 (SiPh<sub>2</sub>), 473.2 (SiPh<sub>2</sub>), 474.2 (SiPh<sub>2</sub>), 475.2 (SiPh<sub>2</sub>), 476.2 (SiPh<sub>2</sub>), 477.2 (SiPh<sub>2</sub>), 478.2 (SiPh<sub>2</sub>), 479.2 (SiPh<sub>2</sub>), 480.2 (SiPh<sub>2</sub>), 481.2 (SiPh<sub>2</sub>), 482.2 (SiPh<sub>2</sub>), 483.2 (SiPh<sub>2</sub>), 484.2 (SiPh<sub>2</sub>), 485.2 (SiPh<sub>2</sub>), 486.2 (SiPh<sub>2</sub>), 487.2 (SiPh<sub>2</sub>), 488.2 (SiPh<sub>2</sub>), 489.2 (SiPh<sub>2</sub>), 490.2 (SiPh<sub>2</sub>), 491.2 (SiPh<sub>2</sub>), 492.2 (SiPh<sub>2</sub>), 493.2 (SiPh<sub>2</sub>), 494.2 (SiPh<sub>2</sub>), 495.2 (SiPh<sub>2</sub>), 496.2 (SiPh<sub>2</sub>), 497.2 (SiPh<sub>2</sub>), 498.2 (SiPh<sub>2</sub>), 499.2 (SiPh<sub>2</sub>), 500.2 (SiPh<sub>2</sub>), 501.2 (SiPh<sub>2</sub>), 502.2 (SiPh<sub>2</sub>), 503.2 (SiPh<sub>2</sub>), 504.2 (SiPh<sub>2</sub>), 505.2 (SiPh<sub>2</sub>), 506.2 (SiPh<sub>2</sub>), 507.2 (SiPh<sub>2</sub>), 508.2 (SiPh<sub>2</sub>), 509.2 (SiPh<sub>2</sub>), 510.2 (SiPh<sub>2</sub>), 511.2 (SiPh<sub>2</sub>), 512.2 (SiPh<sub>2</sub>), 513.2 (SiPh<sub>2</sub>), 514.2 (SiPh<sub>2</sub>), 515.2 (SiPh<sub>2</sub>), 516.2 (SiPh<sub>2</sub>), 517.2 (SiPh<sub>2</sub>), 518.2 (SiPh<sub>2</sub>), 519.2 (SiPh<sub>2</sub>), 520.2 (SiPh<sub>2</sub>), 521.2 (SiPh<sub>2</sub>), 522.2 (SiPh<sub>2</sub>), 523.2 (SiPh<sub>2</sub>), 524.2 (SiPh<sub>2</sub>), 525.2 (SiPh<sub>2</sub>), 526.2 (SiPh<sub>2</sub>), 527.2 (SiPh<sub>2</sub>), 528.2 (SiPh<sub>2</sub>), 529.2 (SiPh<sub>2</sub>), 530.2 (SiPh<sub>2</sub>), 531.2 (SiPh<sub>2</sub>), 532.2 (SiPh<sub>2</sub</sub>

*CH<sub>3</sub>*), 123.9, 124.6, 125.6, 129.0, 129.1, 129.6, 129.7, 131.1, 131.5, 132.8, 132.9, 133.3, 133.4, 134.8, 135.4, 137.3, 137.8, 140.8, 140.9, 147.8 and 149.7 (Ar-C).  $^{31}\text{P}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 162 MHz, 298 K):  $\delta$  = 4.0 (s,  $\text{CH}_2\text{-PPh}_2$ ).  $^{29}\text{Si}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 79 MHz, 298 K):  $\delta$  = -4.0 (d,  $^2J_{\text{SiP}} = 13.4$  Hz,  $\text{SiPh}_2$ ). MS/LIFDI-HRMS found (calcd) *m/z*: 665.1466 (665.1490) for  $[\text{M}]^+$ .

### $[\text{PhPhDippGeBr}]$ , 3a

To a mixture of **1a** (200 mg, 0.30 mmol) and  $\text{NiBr}_2\cdot\text{DME}$  (540 mg, 1.8 mmol) was added 5 mL toluene and 1 mL THF, and the resulting mixture stirred for 1 h. All volatiles were removed *in vacuo* and the residue extracted with 5 mL toluene. The solution was concentrated and layered with hexane yielding colourless crystals of **3a** suitable for X-ray diffraction analysis (127 mg, 0.18 mmol, 54%).  $^1\text{H}$  NMR ( $\text{C}_6\text{D}_6$ , 400 MHz, 298 K):  $\delta$  = 0.06 (d, 3H,  $^3J_{\text{HH}} = 6.6$  Hz, Dipp- $\text{Pr}^i\text{-CH}_3$ ), 0.67 (d, 3H,  $^3J_{\text{HH}} = 6.7$  Hz, Dipp- $\text{Pr}^i\text{-CH}_3$ ), 0.93 (d, 3H,  $^3J_{\text{HH}} = 6.6$  Hz, Dipp- $\text{Pr}^i\text{-CH}_3$ ), 1.46 (d, 3H,  $^3J_{\text{HH}} = 6.6$  Hz, Dipp- $\text{Pr}^i\text{-CH}_3$ ), 2.55 (m, 3H,  $\text{Ph}_2\text{P-CH}_2$ /Dipp- $\text{Pr}^i\text{-CH}$ ), 4.26 (hept, 1H,  $^3J_{\text{HH}} = 6.9$  Hz, Dipp- $\text{Pr}^i\text{-CH}$ ), 6.99 (m, 14H, Ar-CH), 7.21 (m, 3H, Ar-CH), 7.46 (m, 2H, Ar-CH), 7.60 (m, 2H, Ar-CH), 8.30 (d, 2H,  $^3J_{\text{HH}} = 7.3$  Hz, Ar-CH).  $^{13}\text{C}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 101 MHz, 298 K):  $\delta$  = 8.4 ( $\text{Ph}_2\text{P-CH}_2$ ), 21.4 and 22.6 (Dipp- $\text{Pr}^i\text{-CH}_3$ ), 28.1 and 28.2 (Dipp- $\text{Pr}^i\text{-CH}$ ), 28.5 and 29.0 (Dipp- $\text{Pr}^i\text{-CH}_3$ ), 123.1, 124.0, 124.8, 125.7, 128.9, 129.0, 129.1, 129.6, 129.8, 131.0, 131.6, 132.7, 132.8, 133.3, 133.4, 134.5, 135.4, 137.6, 140.5, 140.6, 147.8 and 149.9 (Ar-C).  $^{31}\text{P}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 162 MHz, 298 K):  $\delta$  = 1.9 (s,  $\text{CH}_2\text{-PPh}_2$ ).  $^{29}\text{Si}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 99 MHz, 298 K):  $\delta$  = -3.9 (d,  $^2J_{\text{SiP}} = 13.9$  Hz,  $\text{SiPh}_2$ ); MS/LIFDI-HRMS found (calcd) *m/z*: 709.1012 (709.0984) for  $[\text{M}]^+$ .

### $[\text{PhPhDippGe(Cl)}\text{Ni}(\text{PPh}_3)_2]$ , 4a

To a mixture of **1a** (1.60 g, 2.4 mmol),  $\text{NiCl}_2\cdot\text{DME}$  (0.53 g, 2.4 mmol),  $\text{PPh}_3$  (1.26 g, 4.8 mmol), and Zn (0.94 g, 14.4 mmol) was added 10 mL THF, and the resulting mixture stirred for 24 h at RT resulting in a deep red reaction mixture. All volatiles were removed *in vacuo* and the residue extracted with 20 mL diethyl ether. Dark red crystals of **4a**, which were suitable for X-ray diffraction analysis, were obtained after storing the solution at RT overnight (1.55 g, 1.8 mmol, 52%).  $^1\text{H}$  NMR ( $\text{C}_6\text{D}_6$ , 400 MHz, 298 K):  $\delta$  = 0.49 (d, 6H,  $^3J_{\text{HH}} = 6.6$  Hz, Dipp- $\text{Pr}^i\text{-CH}_3$ ), 1.28 (bs, 6H, Dipp- $\text{Pr}^i\text{-CH}_3$ ), 3.03 (s, 2H,  $\text{Ph}_2\text{P-CH}_2$ ), 3.72 (hept, 2H,  $^3J_{\text{HH}} = 6.6$  Hz, Dipp- $\text{Pr}^i\text{-CH}$ ), 6.63 (m, 3H, Ar-CH), 6.94 (m, 36H, Ar-CH), 7.32 (d, 5H,  $^3J_{\text{HH}} = 6.2$  Hz, Ar-CH), 7.52 (s, 9H, Ar-CH).  $^{13}\text{C}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 101 MHz, 298 K):  $\delta$  = 20.0 ( $\text{Ph}_2\text{P-CH}_2$ ), 23.8 and 26.1 (Dipp- $\text{Pr}^i\text{-CH}_3$ ), 29.2 (Dipp- $\text{Pr}^i\text{-CH}$ ), 124.0, 125.6, 127.2, 128.8, 128.8, 129.1, 132.3, 134.5, 134.6, 135.3, 136.0, 138.6, 138.9, 142.3 and 145.5 (Ar-C).  $^{31}\text{P}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 81 MHz, 298 K):  $\delta$  = 7.1 (t,  $^2J_{\text{PP}} = 18.2$  Hz,  $\text{Ph}_2\text{P-Ni-(PPh}_3)_2$ ), 39.8 (bs,  $\text{Ph}_2\text{P-Ni-(PPh}_3)_2$ ).  $^{29}\text{Si}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 99 MHz, 298 K):  $\delta$  = -13.5 (d,  $^2J_{\text{SiP}} = 5.5$  Hz,  $\text{SiPh}_2$ ). MS/LIFDI-HRMS found (calcd) *m/z*: 985.1705 (985.1755) for  $[\text{M} - \text{PPh}_3]^+$ .

### $[\text{PhPhDippGe(NH}_2)\text{Ni}(\text{PPh}_3)_2]$ , 5a

Compound **4a** (20 mg, 0.016 mmol) was dissolved in 0.4 mL  $\text{C}_6\text{D}_6$  in an NMR tube. An excess of ammonia was added to the

NMR tube, which was then closed and shaken leading to an immediate colour change from deep red to bright orange, with concomitant formation of a colourless solid ( $\text{NH}_4\text{Cl}$ ). The solution was filtered, and volatiles removed *in vacuo* to yield **5a** as an orange powder (15 mg, 0.013 mmol, 83%). Dark orange crystals suitable for X-ray diffraction analysis were obtained by storage of a concentrated toluene solution layered with hexane at -32 °C.  $^1\text{H}$  NMR ( $\text{C}_6\text{D}_6$ , 400 MHz, 298 K):  $\delta$  = 0.50 (d, 6H,  $^3J_{\text{HH}} = 6.6$  Hz, Dipp- $\text{Pr}^i\text{-CH}_3$ ), 1.14 (m, 6H, Dipp- $\text{Pr}^i\text{-CH}_3$ ), 2.97 (s, 2H,  $\text{Ph}_2\text{P-CH}_2$ ), 3.14 (s, 2H, Ge- $\text{NH}_2$ ), 3.79 (hept, 2H,  $^3J_{\text{HH}} = 7.3$  Hz, Dipp- $\text{Pr}^i\text{-CH}$ ), 6.67 (m, 3H, Ar-CH), 6.81 (m, 6H, Ar-CH), 7.00 (m, 31H, Ar-CH), 7.32 (d, 4H,  $^3J_{\text{HH}} = 6.6$  Hz, Ar-CH), 7.48 (s, 9H, Ar-CH).  $^{13}\text{C}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 101 MHz, 298 K):  $\delta$  = 20.1 ( $\text{Ph}_2\text{P-CH}_2$ ), 23.6 and 26.1 (Dipp- $\text{Pr}^i\text{-CH}_3$ ), 28.8 (Dipp- $\text{Pr}^i\text{-CH}$ ), 124.4, 125.2, 127.1, 131.6, 132.3, 134.3, 134.5, 136.1, 136.3, 140.2, 141.1 and 146.3 (Ar-C).  $^{31}\text{P}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 81 MHz, 298 K):  $\delta$  = 9.3 (t,  $^2J_{\text{PP}} = 15.6$  Hz,  $\text{Ph}_2\text{P-Ni-(PPh}_3)_2$ ), 39.8 (d,  $^2J_{\text{PP}} = 14.2$  Hz,  $\text{Ph}_2\text{P-Ni-(PPh}_3)_2$ ).  $^{29}\text{Si}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 99 MHz, 298 K):  $\delta$  = -14.8 (d,  $^2J_{\text{SiP}} = 3.2$  Hz,  $\text{SiPh}_2$ ); IR,  $\nu/\text{cm}^{-1}$  (ATR): 3354 and 3461 (br, w, Ge- $\text{NH}_2$ ); MS/LIFDI-HRMS found (calcd) *m/z*: 966.2156 (966.2253) for  $[\text{M} - \text{PPh}_3]^+$ .

### $[\text{PhPhDippGe(OH)}\text{Ni}(\text{PPh}_3)_2]$ , 6a

'Wet' cyclohexylaniline (100 mg, 1.00 mmol) was added to a stirring solution of **4a** (200 mg, 0.16 mmol) in 5 mL toluene leading to an immediate colour change from deep red to light red-orange. The mixture was stirred for 30 min. All volatiles were then removed *in vacuo* and the residue extracted with diethyl ether, and filtered. Orange-red crystals of **6a** suitable for X-ray diffraction analysis formed over the course of two hours at ambient temperature (133 mg, 0.11 mmol, 68%).  $^1\text{H}$  NMR ( $\text{C}_6\text{D}_6$ , 400 MHz, 298 K):  $\delta$  = 0.47 (d, 6H,  $^3J_{\text{HH}} = 6.3$  Hz, Dipp- $\text{Pr}^i\text{-CH}_3$ ), 1.10 (m, 6H, Dipp- $\text{Pr}^i\text{-CH}_3$ ), 2.99 (s, 2H,  $\text{Ph}_2\text{P-CH}_2$ ), 3.77 (hept, 2H,  $^3J_{\text{HH}} = 6.4$  Hz, Dipp- $\text{Pr}^i\text{-CH}$ ), 4.15 (s, 1H, Ge- $\text{OH}$ ), 6.67 (m, 3H, Ar-CH), 6.83 (m, 6H, Ar-CH), 7.01 (m, 30H, Ar-CH), 7.34 (d, 4H,  $^3J_{\text{HH}} = 7.0$  Hz, Ar-CH), 7.54 (s, 1H, Ar-CH).  $^{13}\text{C}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 101 MHz, 298 K):  $\delta$  = 19.4 ( $\text{Ph}_2\text{P-CH}_2$ ), 23.3 and 26.2 (Dipp- $\text{Pr}^i\text{-CH}_3$ ), 28.9 (Dipp- $\text{Pr}^i\text{-CH}$ ), 124.5, 125.7, 127.2, 128.9, 132.3, 134.5, 134.5, 134.6, 135.7, 135.8, 136.0, 138.8, 139.7, 139.9, 140.0 and 146.6 (Ar-C).  $^{31}\text{P}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 81 MHz, 298 K):  $\delta$  = 9.0 (t,  $^2J_{\text{PP}} = 15.8$  Hz,  $\text{Ph}_2\text{P-Ni-(PPh}_3)_2$ ), 40.6 (d,  $^2J_{\text{PP}} = 15.8$  Hz,  $\text{Ph}_2\text{P-Ni-(PPh}_3)_2$ ).  $^{29}\text{Si}\{\text{H}\}$  NMR ( $\text{C}_6\text{D}_6$ , 99 MHz, 298 K):  $\delta$  = -14.4 (d,  $^2J_{\text{SiP}} = 4.6$  Hz,  $\text{SiPh}_2$ ); IR,  $\nu/\text{cm}^{-1}$  (ATR): 3547 (m, Ge- $\text{OH}$ ); MS/LIFDI-HRMS found (calcd) *m/z*: 967.2057 (967.2093) for  $[\text{M} - \text{PPh}_3]^+$ .

## Author contributions

PMK carried out the vast majority of the chemical synthesis and characterisation. TS carried out all computational work. TJH designed and planned the project, carried out preliminary experimental work, and wrote the manuscript.

## Conflicts of interest

There are no conflicts to declare.



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## Notes and references

- 1 E. O. Fischer and A. Maasböl, *Angew. Chem., Int. Ed.*, 1964, **8**, 580–581.
- 2 For a broad overview, see the following collections and reviews, and references therein: (a) D. J. Cardin, B. Cetinkaya and M. F. Lappert, *Chem. Rev.*, 1972, **72**, 545–574; (b) A. J. Arduengo and G. Bertrand, *Chem. Rev.*, 2009, **109**, 3209–3210; (c) H. G. Raubenheimer, *Dalton Trans.*, 2014, **43**, 16959–16973; (d) F. Ekkehardt Hahn, *Chem. Rev.*, 2018, **118**, 9455–9456.
- 3 (a) R. R. Schrock and A. H. Hoveyda, *Angew. Chem., Int. Ed.*, 2003, **42**, 4592–4633; (b) D. Astruc, Metal–Carbene and –Carbyne Complexes and Multiple Bonds with Transition Metals, in *Organometallic Chemistry and Catalysis*, Springer, Berlin, Heidelberg, 2007.
- 4 (a) F. Glorius, *N-Heterocyclic Carbenes in Transition Metal Catalysis*, Springer, Berlin, Heidelberg, 2007; (b) S. Diez-González, N. Marion and S. P. Nolan, *Chem. Rev.*, 2009, **109**, 3612–3676; (c) E. Peris, *Chem. Rev.*, 2018, **118**, 9988–10031.
- 5 (a) R. Waterman, P. G. Hayes and T. D. Tilley, *Acc. Chem. Res.*, 2007, **40**, 712–719; (b) J. Baumgartner and C. Marschner, *Rev. Inorg. Chem.*, 2013, **34**, 119–152; (c) L. Álvarez-Rodríguez, J. A. Cabeza, P. García-Álvarez and D. Polo, *Coord. Chem. Rev.*, 2015, **300**, 1–28.
- 6 T. J. Hadlington, M. Driess and C. Jones, *Chem. Soc. Rev.*, 2018, **47**, 4176–4197.
- 7 Y.-P. Zhou and M. Driess, *Angew. Chem., Int. Ed.*, 2019, **58**, 3715–3728.
- 8 (a) L. Álvarez-Rodríguez, J. Brugos, J. A. Cabeza, P. García-Álvarez, E. Pérez-Carreño and D. Polo, *Chem. Commun.*, 2017, **53**, 893–896; (b) L. Álvarez-Rodríguez, J. Brugos, J. A. Cabeza, P. García-Álvarez and E. Pérez-Carreño, *Chem.-Eur. J.*, 2017, **23**, 15107–15115; (c) J. Brugos, J. A. Cabeza, P. García-Álvarez and E. Pérez-Carreño, *Organometallics*, 2018, **37**, 1507–1514; (d) J. A. Cabeza, P. García-Álvarez, C. J. Laglera-Gándara and E. Pérez-Carreño, *Chem. Commun.*, 2020, **56**, 14095–14097.
- 9 A related bis(phosphine) functionalised N-heterocyclic germylene ligand has also been reported by Goicoechea *et al.*, alongside its group 11 halide and Pt<sup>0</sup> complexes: S. Bestgen, N. H. Rees and J. M. Goicoechea, *Organometallics*, 2018, **37**, 4147–4155.
- 10 (a) J. A. Cabeza, I. Fernández, J. M. Fernández-Colinas, P. García-Álvarez and C. J. Laglera-Gándara, *Chem.-Eur. J.*, 2019, **25**, 12423–12430; (b) J. A. Cabeza, I. Fernández, P. García-Álvarez and C. J. Laglera-Gándara, *Dalton Trans.*, 2019, **48**, 13273–13280; (c) A. Arauzo, J. A. Cabeza, I. Fernández, P. García-Álvarez, I. García-Rubio and C. J. Laglera-Gándara, *Chem.-Eur. J.*, 2021, DOI: 10.1002/chem.202005289.
- 11 (a) K. E. Litz, J. E. Bender IV, J. W. Kampf and M. M. Banaszak Holl, *Angew. Chem., Int. Ed. Engl.*, 1997, **36**, 496–498; (b) J. E. Bender IV, A. J. Shusterman, M. M. Banaszak Holl and J. W. Kampf, *Organometallics*, 1999, **18**, 1547–1552; (c) C. Gendy, A. Mansikkamäki, J. Valjus, J. Heidebrecht, P. C.-Y. Hui, G. M. Bernard, H. M. Tuononen, R. E. Wasylishen, V. K. Michaelis and R. Roesler, *Angew. Chem., Int. Ed.*, 2019, **58**, 15–158; (d) T. Watanabe, Y. Kasai and H. Tobita, *Chem.-Eur. J.*, 2019, **25**, 13491–13495; (e) Z. Feng, Y. Jiang, H. Ruan, Y. Zhao, G. Tan, L. Zhang and X. Wang, *Dalton Trans.*, 2019, **48**, 14975–14978; (f) M. Zhong, J. Wei, W.-X. Zhang and Z. Xi, *Organometallics*, 2021, **40**, 310–313.
- 12 (a) H. Grützmacher, *Angew. Chem., Int. Ed.*, 2008, **47**, 1814; (b) J. R. Khusnutdinova and D. Milstein, *Angew. Chem., Int. Ed.*, 2015, **54**, 12236; (c) M. R. Elsby and R. T. Baker, *Chem. Soc. Rev.*, 2020, **49**, 8933–8987.
- 13 (a) V. Lyaskovskyy and B. de Bruin, *ACS Catal.*, 2012, **2**, 270; (b) J. van der Vlugt, *Eur. J. Inorg. Chem.*, 2012, 363; (c) S. Schneider, J. Meiners and B. Askevold, *Eur. J. Inorg. Chem.*, 2012, 412.
- 14 It should be noted that the concept of “Synergistic Catalysis” is perhaps also fitting here, given the generally high reactivity of low-coordinate tetrylenes in their own right. For a review on this concept and its implications, see: U. B. Kim, D. J. Jung, H. J. Jeon, K. Rathwell and S. gi Lee, *Chem. Rev.*, 2020, **120**, 13382–13433.
- 15 (a) A. L. Casalnuovo, J. C. Calabrese and D. Milstein, *Inorg. Chem.*, 1987, **26**, 971–973; (b) E. Khaskin, M. A. Iron, L. J. W. Shimon, J. Zhang and D. Milstein, *J. Am. Chem. Soc.*, 2010, **132**, 8542–8543; (c) D. V. Gutsulyak, W. E. Piers, J. Borau-Garcia and M. Parvez, *J. Am. Chem. Soc.*, 2013, **135**, 11776–11779; (d) R. M. Brown, J. Borau Garcia, J. Valjus, C. J. Roberts, H. M. Tuononen, M. Parvez and R. Roesler, *Angew. Chem., Int. Ed.*, 2015, **54**, 6274–6277.
- 16 Procedures modified from those previously reported for related compounds were used. See: R. D. Holmes-Smith, R. D. Osei and S. R. Stobart, *J. Chem. Soc., Perkin Trans. 1*, 1983, 861–866.
- 17 The free ligands typically form oils at ambient temperature, but a few single crystals sometimes deposit from decomposed reaction mixtures of the metallated ligands. See ESI† for details.
- 18 J. Schneider, K. M. Krebs, S. Freitag, K. Eichele, H. Schubert and L. Wesemann, *Chem.-Eur. J.*, 2016, **22**, 9812–9826.



19 J. Berthe, J. Manuel Garcia, E. Ocando, T. Kato, N. Saffon-Merceron, A. De Cózar, F. P. Cossío and A. Baceiredo, *J. Am. Chem. Soc.*, 2011, **133**, 15930–15933.

20 The bromo-germylene, PhiPDippGeBr (**3b**), can be accessed using an identical method (see ESI† for details).

21 Yields of 50–55% are obtained after recrystallisation.

22 As an example, GeBr<sub>2</sub>·dioxane, used as is GeCl<sub>2</sub>·dioxane, can be synthesised from GeBr<sub>4</sub>, which is a high-cost starting material: N. Hayakawa, T. Sugahara, Y. Numata, H. Kawai, K. Yamatani, S. Nishimura, S. Goda, Y. Suzuki, T. Tanikawa, H. Nakai, D. Hashizume, T. Sasamori, N. Tokitoh and T. Matsuo, *Dalton Trans.*, 2018, **47**, 814–822.

23 Note that single crystals of pure **4a** reproducibly gave poor X-ray diffraction data, not of a publishable standard. However, **4a** co-crystallised with ~19% of **4-Br** gave data of a publishable standard. See ESI† for details.

24 C. A. Tolman, W. C. Seidel and D. H. Gerlach, *J. Am. Chem. Soc.*, 1972, **94**, 2669–2676.

25 Attempts to synthesise pure samples of the bromo-germylene Ni<sup>0</sup> complex, **4-Br**, are outlined in the ESI.†

26 Despite attempts to recrystallise the corresponding amide complex **5b** from a range of solvents, only very fine needles could be isolated which diffracted too poorly to attain meaningful X-ray data.

27 As ascertained by <sup>1</sup>H and <sup>31</sup>P NMR spectroscopic analysis. Due to the extremely high sensitivity of **5a/b** towards moisture, the degassing cycles led to the formation of small amounts of hydroxide complexes **6a/b**.

28 For some recent examples, see: (a) T. J. Hadlington, J. A. B. Abdalla, R. Tirfoin, S. Aldridge and C. Jones, *Chem. Commun.*, 2016, **52**, 1717–1720; (b) A. V. Protchenko, J. I. Bates, L. M. A. Saleh, M. P. Blake, A. D. Schwarz, E. L. Kolychev, A. L. Thompson, C. Jones, P. Mountford and S. Aldridge, *J. Am. Chem. Soc.*, 2016, **138**, 4555–4564; (c) D. C. H. Do, A. V. Protchenko, M. Á. Fuentes, J. Hicks, P. Vasko and S. Aldridge, *Chem. Commun.*, 2020, **56**, 4684–4687.

29 Z. Benedek and T. Szilvási, *Organometallics*, 2017, **36**, 1591–1600.

30 H. Bauer, J. Weismann, D. Saurenz, C. Färber, M. Schär, W. Gidt, I. Schädlich, G. Wolmershäuser, Y. Sun, S. Harder and H. Sitzmann, *J. Organomet. Chem.*, 2016, **809**, 63–73.

31 A. Boudier, P.-A. R. Breuil, L. Magna, H. Olivier-Bourbigou and P. Braunstein, *J. Organomet. Chem.*, 2012, **718**, 31–37.

32 J. T. Patton, M. M. Bokota and K. A. Abboud, *Organometallics*, 2002, **21**, 2145–2148.

33 N. E. Schore, L. S. Benner and B. E. LaBelle, *Inorg. Chem.*, 1981, **20**, 3200–3208.

