**Amide Ligands**

**A Xanthene-Based Mono-Anionic PON Ligand: Exploiting a Bulky, Electronically Unsymmetrical Donor in Main Group Chemistry**

Xiongfei Zheng, Andreas Heilmann, Caitilín McManus, and Simon Aldridge

**Abstract:** The synthesis of a novel mono-anionic phosphino-amide ligand based on a xanthene backbone is reported, together with the corresponding Ga complex, (PON)Ga (PON = 4-(di(2,4,6-trimethylphenyl)phosphino)-5-(2,6-diisopropylanilido)-2,7-di-tert-butyl-9,9-dimethylxanthene). The solid-state structure of (PON)Ga (obtained from X-ray crystallography) reveals very weak O–Ga and P–Ga interactions, consistent with a R3NGa fragment which closely resembles those found in one-coordinate amidogallium systems. Strong N-to-Ga π donation from the amido substituent is reflected in a very short N–Ga distance (1.961(2) Å), while the P–Ga contact (3.076(1) Å) is well outside the sum of the respective covalent radii. While the donor properties of the PON ligand towards Ga are highly unsymmetrical, oxidation to Ga4+ leads to much stronger coordination of the pendant phosphine as shown by P–Ga distances which are up to 20% shorter. From a steric perspective, the PON ligand is shown to be significantly bulkier than related β-diketiminate systems, a finding consistent with reactions of (PON)Ga towards O-atom sources that proceed without oligomerization. Despite this, the enhanced P-donor properties brought about by oxidation at gallium are not sufficient to quench the reactivity of the highly polar Ga–O unit. Instead, intramolecular benzylic C–H activation is observed across the Ga–O bond of a transient gallanone intermediate.

**Introduction**

In recent years, β-diketiminate (‘Nacnac’) ligands have been extensively used in coordination chemistry, to support a wide range of metal complexes from across the Periodic Table. Within main group chemistry, a number of landmark compounds have been reported incorporating these chelating monoanionic LX ligand systems. These include Group 13 metal complexes in the +1 oxidation state—systems which are challenging both in terms of their intrinsic tendency to undergo disproportionation and their lability towards external reagents. Thus in 2000, the groups of Roesky and Power successfully synthesized the monomeric Al and Ga compounds [HCl(Me)(C(Dipp)N)2]E (or (Nacnac)E, I, where E = Al, Ga) using sterically encumbered Dipp-substituted β-diketiminate ligands (Figure 1; Dipp = 2,6-iPr2C6H3).[5–7] These systems show unusual patterns of reactivity: the aluminyl anion can act as a nucleophile in the formation of C–Al and M–Al bonds[8,9] and effect the reversible oxidative addition of the C–C bond in benzene.[10] Given the track record of the Nacnac ligand family in supporting charge neutral E(I) systems, however,[2–4] we targeted related mono-anionic LX donors based on the dimethylxanthene backbone, in which one of the amido groups of the [NON]2+ system is formally replaced by a neutral donor. Moreover, given the successful development by Power, and by Jones, of one-coordinate Ga amides (stabilized to a greater or lesser extent by interactions with the flanking aryl substituents, III),[11] we targeted xanthene systems featur-
ing L substituents which are known to act as relatively weak donors towards Ga. Our aim was to develop amido complexes featuring a weakly interacting (or hemi-labile) L donor that might display the high levels of reactivity associated with genuine one-coordinate systems, while retaining some of the ground state stability of Nacnac compounds. Accordingly, we report in the current manuscript on the development of a mono-anionic [PON]− ligand (Figure 1) and the use of GaⅢ and GaⅣ chemistry to probe its coordination capabilities.

Results and Discussion

(i) Synthesis of H(PON) and (PON)Ga

The H(PON) protio-ligand 2 can be synthesized in good yield (ca. 70%) from the (known) 4-bromo-S-(dimethylphosphino)-xanthene precursor (1)[15] and DippNH₂ via a Buchwald-Hartwig coupling reaction. 2 has been characterized by multinuclear NMR, elemental microanalysis and single crystal X-ray diffraction (see SI). Moreover, it is conveniently deprotonated by benzylpotassium, KCH₂Ph, giving the potassiated ligand K(PON) (3), which can be used for onward reaction chemistry without further purification, or recrystallized from toluene for structural and spectroscopic characterization. 3 is thus obtained free of coordinating solvent as dimeric K₂(PON)₂ in which K(PON) monomer units are linked in head-to-tail fashion through close contacts between the potassium centre of one unit and the Dipp aromatic π system of the other (Figure 2).

With this new ligand system in hand we set out to probe its coordination chemistry, using gallanediyl (GaⅢ) and gallium oxide (GaⅣ) systems as probes of its ability to support highly reactive main group metal centres. Accordingly, the GaⅢ system, (PON)Ga, 4 can be prepared in ca. 60% isolated yield via a salt-metathesis reaction between 3 and “Gal” (Scheme 1);[16] and obtained in crystalline form by recrystallization from benzene. The ¹H NMR spectrum of 4 features a similar pattern of resonances to those of H(PON) and K(PON), with equivalent phosphorus-bound mesityl substituents, Dipp πR groups and xanthene backbone methyl groups implying the presence of a plane of symmetry on the NMR timescale. The ³¹P[¹H] resonance of 4 (at δₚ = −34.3 ppm) is very close to those of protio-ligand 2 (δₚ = −36.9 ppm) and potassiated system 3 (δₚ = −34.4 ppm), implying that the interaction of the phosphine donor with the gallium centre in 4 is relatively weak. Consistently, the (monomeric) structure of 4 in the solid state (Figure 2) features a very long Ga–P distance (3.076(1) Å) which is comfortably outside the sum of the respective covalent radii (1.22(4)+1.07(3) Å)[17] and much longer than that measured for the corresponding GaⅣ system (PON)GaI which was synthesized for comparative purposes (2.612(1) Å; see SI). The Ga–O separation involving the xanthene ether linkage is also relatively long (2.631(2) Å) compared to the sum of the respective covalent radii (1.22(3)+0.66(2) Å);[18] and is broadly comparable to that measured for the potassium gallium system K₃[(NON)Ga]₂ featuring the related [NON]²− diamide ligand (2.542(2) Å).[19] On the other hand, the Ga–N bond is very short (1.961(2) Å) in comparison with those in K₃[(NON)Ga]₂ (2.093(2) and 2.106(2) Å)[19] presumably reflecting the fact that there is only one Ga-X covalent bond in 4. Indeed, the Ga–N separation in 4 is in line with the bond lengths reported by Power and by Jones for GaⅢ amides with geometries which approach mono-coordinate (1.954(2)–1.983(4) Å),[20] consistent with idea that the coordination of the phosphine and ether donors in 4 is very weak.

(ii) Electronic and steric properties of (PON)Ga

The electronic structure of (PON)Ga (4) was probed using Density Functional Theory at the PBE1PBE/TZVP level. These calculations suggest that the LUMO + 2 (−0.68 eV) is comprised predominantly of gallium pₓ character (Figure 3). The HOMO is ligand-based, with the orbital displaying gallium-centred lone pair character being the HOMO-1 (−5.67 eV). The associated energy separation (4.99 eV) is very similar to that calculated for (Nacnac)GaⅢ using the same method (4.86 eV). The essentially identical energies of the formally vacant px orbital in each case (−0.66 eV for (Nacnac)Ga) presumably reflect the fact that in 4, the extent of π donation from the (single) amido substituent is markedly enhanced, consistent with the very short crys-
tallographically determined Ga–N bond length. Both of these systems have wider HOMO–LUMO gaps than that determined for the anionic diamido system [(NON)Ga]−. In that case, the narrow HOMO–LUMO gap (4.21 eV) reflects the fact that the LUMO is not destabilized to the same degree by N-to-Ga π donation due to the constraints of the xanthene backbone (which mean that the amido groups cannot attain coplanarity with the GaN unit).[16] The (PON)Ga system can also be compared with Jones’ one-coordinate amidogallium(I) system,[14b] which is similar in that it features π stabilization from only a single N donor. In that system, the HOMO–LUMO gap (calculated using the same method) is narrower (4.62 eV) and the LUMO somewhat lower than in 4 (−0.91 eV). This presumably reflects the fact that the LUMO is orthogonal to the amido π system, and (compared to 4) has less opportunity for interaction with ancillary neutral donors (featuring as it does, only very weak interaction with the flanking arene π system).

Steric factors are known to be a significant influence on the patterns of reactivity displayed by low-valent main group species. In both (PON)Ga and (Nacnac)Ga, the Ga′ centre sits in a “pocket” between flanking aryl substituents: in the β-diketonate system these are the N-bound Dipp groups, which are aligned such that the distances from the arene centroids to the metal are each ca. 4.0 Å, and the “pocket” defined by the open centroid-Ga-centroid angle occupies 189.1° in angular terms (Figure 4).[20] In 4 the gallium centre is flanked by NDipp and PMes2 substituents (the centroids of which are also ca. 4.0 Å from the metal centre), and the open “pocket” by comparison occupies 156.8°. As such, we hypothesized that 4 might offer the possibility of a more sterically shielded pocket in which to carry out chemistry at the metal centre.

Figure 4. Comparison of the aryl-flanked “pockets” adjacent to the metal centre in (Nacnac)Ga and (PON)Ga (4).

Scheme 2. Oxidation of (PON)Ga by N₂O: formation of products derived from intramolecular C–H or intermolecular B–C bond activation at a putative gallenate intermediate.

To probe the idea of enhanced steric bulk and encouraged by the finding that the weakly-bound phosphine donor becomes more strongly stabilizing on oxidation of the metal centre (cf. (PON)Ga and (PON)GaI), we examined the reactivity of 4 towards oxygen atom transfer agents. The isolation of a (hitherto unknown) gallanone complex containing a terminal GaO moiety, would presumably require both very strongly α-donating and sterically demanding ancillary ligands.[18]

Power et al. have reported the formation of [(Nacnac)Ga(μ-O)], from the reaction of (Nacnac)Ga with N₂O[19] with the oxo-bridged structure presumably reflecting a steric profile which permits ready dimerization. In contrast, oxidation of 4 with N₂O or Me₂NO at room temperature leads to a monomeric product (5, Scheme 2), albeit one which results from intramolecular C–H activation of one of the ortho-methyl groups of the PMes₂ substituent across the transient gallium oxo unit.[20] Compound 5 has been characterized by spectroscopic methods and its structure in the solid state confirmed crystallographically (Figure 5). The formation of 5 is signalled by the appearance of a 13P NMR resonance at δp = −41.0 ppm and by increased complexity in the pattern of 1H NMR signals associated with the ortho methyl substituents. Its structure in the solid state is based around an approximately tetrahedral gallium centre, featuring a significantly shortened Ga–P distance (2.468(1) Å for (PON)Ga), but with a weak Ga–O contact which is essentially unchanged from its precursor (2.578(2) Å vs. 2.631(2) Å). The Ga–C and Ga–OH bond lengths (1.972(4) and 1.828(2) Å) fall within the ranges previously reported for related (Nacnac)Ga-containing compounds.[21]

Nikonov and co-workers have recently reported the trapping of the monomeric [(Nacnac)GaO] fragment by reactions of (Nacnac)Ga with N₂O in the presence of a range of relatively acidic C–H bonds (e.g. those in sulfoxides and ketones, and at the 2-position of pyridine and related heterocycles).[20] In the case of 4/5, the fact that C–H activation occurs in the absence...
of an external trap (and at an unactivated methyl group) suggests that dimerization of the putative [PON]GaO monomer is less facile than in the related Nacnac system—in line with estimates of the comparative sizes of the ligand-enforced “pockets”.

Attempts to identify a labile gallanone intermediate preceding the formation of 5 by VT-NMR methods were unsuccessful, so alternative (chemical) trapping methods were pursued via coordination of either a Lewis acid (at O) or a Lewis base (at Ga). The latter strategy led either to no change in the outcome of the reaction (using iPrMe) or to the precipitation of gallium metal in the presence of the less bulky NHC IMe. On the other hand, addition of N,O in the presence of a boron Lewis acid (either B(C$_6$F$_5$)$_3$, or BPh$_3$) leads to the clean formation of new products which are characterized by $^1$H-NMR signals at $\sim$36.1 and $\sim$35.6 ppm, respectively. Both products could be unambiguously characterized by X-ray crystallography, and are shown to result from cleavage of a B–C bond in BPh$_3$ (6) or B(C$_6$F$_5$)$_3$ (7) across the Ga–O unit formed in the initial reaction of gallylene 4 with N$_2$O (see Scheme 2, Figure 5 and SI). As such, the isolation of compounds 6 and 7 provides further evidence for the formation of a transient, highly reactive galla- nonene species by oxygen atom transfer to the gallium centre of 4. It is worth noting that similar reactivity—leading to the cleavage of one of the B–C bonds in B(C$_6$F$_5$)$_3$ across a terminal Si–O bond—has been reported by Iwamoto and co-workers in studies of a reactive (but isobale) silanone.$^{[22]}$

Conclusion

In summary, we report on the development of a mono-anionic phosphino-amide ligand based on a xanthene backbone, together with the synthesis of the corresponding Ga complex, (PON)Ga (4), as a probe of its coordination properties in low valent main group chemistry. Structural studies of 4 are consistent with very weak O–Ga and P–Ga interactions, and with an R$_4$NGa fragment that closely resembles those found in one-coordinate amidogallium systems. Strong N-to-Ga π donation from the single amido group leads to a very short N-Ga distance (1.961(2) Å), while the P–Ga contact (3.076(1) Å) is comfortably outside the sum of the respective covalent radii. While the electronic properties of the PON donor are shown to be highly unsymmetrical towards Ga, oxidation to Ga$^{IV}$ leads to stronger coordination of the pendant phosphine arm. From a steric perspective, the profile of the PON ligand is shown to be significantly larger than those of related N,N’-chelated β-diketiminato systems, and underpins reactivities of (PON)Ga towards O-atom sources that proceeds without oligomerization. However, the enhanced P-donor properties seen on oxidation are not sufficient to quench the reactivity of the highly polar Ga–O unit. Instead, intramolecular benzyl C–H activation is observed across the Ga–O bond of a transient gallanone intermediate. Further studies of the coordination chemistry of monoanionic [PON]$^-$ (and related) ligands towards main group E(II) centres are in progress and will be reported in due course.

Experimental Section

General procedures

All manipulations were carried out using standard Schlenk line or dry-box techniques under an atmosphere of argon or dinitrogen. Solvents were degassed by sparging with argon and dried by passing through a column of the appropriate drying agent. NMR spectra were measured in [D$_6$]benzene (which was dried over potassium hydroxide), with the solvent then being distilled under reduced pressure after the addition of D$_2$O and stored under argon in Teflon valve ampoules. NMR samples were prepared under argon in 5 mm Wilmad 507-PP tubes fitted with J. Young Teflon valves, $^1$H, $^1$P{${^1}$H}, $^{13}$C{${^1}$H}, $^{19}$B{${^1}$H}, $^{31}$P{${^1}$H} NMR spectra were measured on a Bruker Avance III HD nanobay 400 MHz or Bruker Avance 500 MHz spectrometer at ambient temperature and referenced internally to residual protio-solvent ($^1$H) or solvent ($^{13}$C) resonances and are reported relative to tetramethylsilane ($\delta$ = 0 ppm). Assignments were confirmed using two-dimensional $^{1}$H–$^{1}$H and $^{13}$C–$^{1}$H NMR correlation experiments. Chemical shifts are quoted in ppm and coupling constants in Hz. Elemental analyses were carried out by Elemental Microanalysis Ltd, Okehampton, Devon, UK. Compound 1,$^{[15]}$ B(C$_6$F$_5$)$_3$ and KCH$_2$Ph$_3$ were prepared by literature methods. All other reagents were used as received. The synthetic and characterizing data for H(PON) (2) and (PON)Ga, are included in the Supporting Information.

Crystallographic details

Single-crystal X-ray diffraction data for compounds 2–7 and (PON)Ga, were collected at 150 K on an Oxford Diffraction/Agilent SuperNova diffractometer using Cu–K$_\alpha$ radiation ($\lambda = 1.54184$ Å), and equipped with a nitrogen gas Oxford Cryosystems cooling unit.$^{[25]}$ Raw frame data were reduced using CrysalisPro.$^{[26]}$ The structures were solved using SHELXT$^{[27]}$ and refined to convergence on $\mathbf{R}^2$ by full-matrix least-squares using SHELXL$^{[28]}$ in combination with OLEX2.$^{[29]}$ Distances and angles were calculated using the full covariance matrix. Restraints were used to maintain sensible geometries for the disordered groups and approximate the displacement parameters to typical values. Selected crystallographic data are summarized in the Table S1. Deposition Numbers 2036919, 2036920, 2036921, 2036922, 2036923, 2036924, and 2036925 contain the supplementary crystallographic data for this paper. These data are provided free of charge by the joint Cambridge Crystallographic Centre.
Density functional theory (DFT) calculations

The computational work was performed using DFT within the Gaussian09 (Revision D.01) program package. Geometry optimizations of the monoanionic ligand systems were performed with the PBE1PBE hybrid exchange-correlation functional using a TZVP basis set. Grime’s empirical dispersion correction (DFT-D3) was included in all geometry optimizations. Unless otherwise stated, geometry optimizations were carried out for the full system, and frequency calculations were performed to confirm the nature of the stationary points found (minimum). Graphics were created with the Avogadro program. The natural bond orbital (NBO) analyses were performed using NBO 3.1 as implemented in Gaussian09.

Syntheses of novel compounds

[KP[ON]3]: Toluene (30 mL) was added to a Schlenk flask containing a mixture of 2 (1.01 g, 150 mmol) and benzylpotassium (96 mg, 0.13 mmol) in toluene (5 mL) and the resulting solution subjected to three freeze-pump-thaw cycles before backfilling with N2 (ca. 1 atm) at −78 °C. The solution was stirred at this temperature for 4 h, and slowly warmed to room temperature. The resulting solution was kept at −30 °C for several days to afford colourless crystals of 5 (40 mg, 41 %). 1H NMR (400 MHz, C6D6, 298 K): δ = −0.11 (d, J = 4.1 Hz, 1H, [GaO]H), 0.97 (d, J = 6.6 Hz, 3H, CH3(C)), 1.03 (d, J = 6.6 Hz, 3H, CH3(C)), 1.08 (s, 3H, CH3(C)), 1.28 (d, J = 7.0 Hz, 3H, CH3(C)), 1.42 (s, 3H, CH3(C)), 1.53 (d, J = 7.0 Hz, 3H, CH3(C)), 1.97 (s, 3H, o-CH3 of Mes), 2.01 (m, 6H, o-CH3 of Mes), 2.16 (s, 3H, p-CH3 of Mes), 2.25 (s, 3H, p-CH3 of Mes), 2.82 (m, 2H, [Ga]CH3), 3.48 (sept, J = 6.9 Hz, 1H, CH3(C)), 3.88 (sept, J = 6.9 Hz, 1H, CH3(C)), 6.30 (dd, J = 1.3, 8.2 Hz, 1H, ArH), 6.45 (s, 1H, ArH), 6.50 (dd, J = 1.3, 7.8 Hz, 1H, ArH), 6.56 (m, 1H, ArH), 6.64 (m, 1H, ArH), 6.82 (td, J = 1.2, 7.6 Hz, 1H, ArH), 6.89 (t, J = 8.0 Hz, 1H, ArH), 7.13 (m, 3H, ArH), 7.22 (t, J = 7.6 Hz, 1H, ArH), 7.30 (m, 1H, ArH), 7.45 (td, J = 1.3, 7.8 Hz, 1H, ArH) ppm. 13C NMR (125.8 MHz, C6D6, 298 K): δ = 155.3 (d, J = 13.5 Hz, 151.3, 151.1, 148.9, 147.8, 143.8, 143.7 (d, J = 8.7 Hz, 143.1, 141.8, 141.2, 141.0, 140.9, 139.7, 136.5 (d, J = 3.8 Hz), 133.0, 132.6 (d, J = 6.3 Hz), 131.5 (d, J = 7.2 Hz), 131.0 (d, J = 3.6 Hz), 130.9 (d, J = 7.5 Hz), 128.9 (d, J = 6.0 Hz), 127.3, 126.7 (t, J = 22.0 Hz), 125.2, 124.7 (d, J = 16.8 Hz), 123.4 (d, J = 5.7 Hz), 121.7 (d, J = 31.0 Hz), 119.6, 119.3, 114.1, 109.3, 36.5, 32.0 (Hexane), 31.6, 28.2 (d, J = 8.5 Hz), 25.3, 25.2, 25.1, 25.0, 24.5, 24.0 (d, J = 5.7 Hz), 23.5, 23.2, 23.1 (Hexane), 21.0, 20.8, 14.4 (Hexane) ppm. 31P NMR (162 MHz, C6D6, 298 K): δ = −41.0 ppm. Elemental microanalysis: calc. C 73.18 % H 6.96 % N 1.90 %, meas. C 73.10 % H 6.67 % N 1.86 %.

Reaction of 4 with N2O; synthesis of C–H activation product 5: A solution of 4 (96 mg, 0.13 mmol) in toluene (5 mL) was subjected to three freeze-pump-thaw cycles before backfilling with N2O (ca. 1 atm) at −78 °C. The solution was stirred at this temperature for 4 h, and slowly warmed to room temperature. The resulting solution was kept at −30 °C for several days to afford colourless crystals of 5 (40 mg, 41 %). 1H NMR (400 MHz, C6D6, 298 K): δ = 155.8 (d, J = 12.1 Hz), 149.4, 148.7, 144.4, 143.8, 143.4, 142.9 (d, J = 22.1 Hz), 142.5, 142.2, 141.7, 140.5, 140.2, 139.9, 139.3, 132.5, 132.8, 131.9, 131.8, 131.5, 130.8, 130.4, 128.8, 128.7, 128.6 (d, J = 4.2 Hz), 127.2, 126.4, 125.8, 125.0, 124.5, 119.4, 119.2, 116.1, 109.3, 35.7, 33.0, 32.0 (hexane), 28.7, 28.1, 26.5, 25.8, 25.6 (d, J = 11.6 Hz), 24.1, 23.7, 23.4, 23.1 (hexane), 20.9, 20.5, 14.4 (hexane) ppm. 31P NMR (162 MHz, C6D6, 298 K): δ = −36.0 ppm. 7 Reaction was obtained from 4 (69 mg, 0.096 mmol) and B(C6F5)3, (49 mg, 0.096 mmol) in similar fashion as colourless crystals (41 mg, 34 %). Characterizing data for 7: 1H NMR (400 MHz, C6D6, 298 K): δ = 0.81 (d, J = 6.6 Hz, 3H, CH3(C)), 0.96 (d, J = 6.6 Hz, 3H, CH3(C)), 1.07 (d, J = 6.6 Hz, 3H, CH3(C)), 1.36 (s, 3H, CH3 of Mes), 1.42 (s, 3H, CH3 of Mes), 1.45 (s, 3H, CH3 of Mes), 1.54 (d, J = 6.8 Hz, 3H, CH3(C)), 1.70 (s, 3H, CH3 of Mes), 1.74 (s, 3H, CH3 of Mes), 1.91 (s, 3H, CH3 of Mes), 2.08 (s, 3H, CH3 of Mes), 2.70 (s, 3H, CH3 of Mes),
Conflict of interest

The authors declare no conflict of interest.

Keywords: amide ligands • gallium • low-valent compounds • oxidation • xanthene ligands

[1] For early references, see: a) W. Bradley, I. Wright, J. Chem. Soc. 1956, 640–648; b) J. E. Parks, R. H. Holm, Inorg. Chem. 1968, 7, 1408–1416; c) S. G. McGechin, Can. J. Chem. 1968, 46, 1903–1912; d) C. L. Honeybourne, G. A. Webb, Chem. Commun. (London) 1968, 739–740; e) R. Bonnett, D. C. Bradley, K. J. Fisher, Chem. Commun. (London) 1968, 886–887.

[2] L. Bourget-Merle, M. F. Lappert, J. R. Severn, Chem. Rev. 2002, 102, 3031–3066.

[3] For high profile examples from across the Periodic Table see, for example: a) M. S. Hill, P. B. Hitchcock, R. Pongtavornpinyo, Science 2006, 311, 1904–1907; b) S. P. Green, C. Jones, A. Stasch, Science 2007, 318, 1754–1757; c) M. M. Rodriguez, E. Bill, W. W. Brennessel, P. L. Holland, Science 2011, 334, 780–783.

[4] For selected recent reviews encompassing aspects of Nacnac chemistry, see: a) M. Asay, C. Jones, M. Diess, Rev. Chem. 2011, 111, 354–396; b) Y. Tsai, Coord. Chem. Rev. 2012, 256, 722–758; c) C. Chen, S. M. Bellows, P. L. Holland, Dalton Trans. 2015, 44, 16554–16670; d) C. J. Arnold, Dalton Trans. 2016, 45, 14462–14498.

[5] C. Dohmeier, D. Loos, H. Schnöckel, Angew. Chem. Int. Ed. Engl. 1996, 35, 129–149; Angew. Chem. 1996, 108, 141–161.

[6] N. J. Hardman, B. E. Eichler, P.P. Power, Chem. Commun. 2000, 1991–1992.

[7] C. Cui, H. Roehsy, H.-G. Schmidt, M. Noltemeyer, H. Hao, F. Cimpoesu, Angew. Chem. Int. Ed. 2000, 39, 4274–4276; Angew. Chem. 2000, 112, 4444–4446.

[8] For other examples of n-heterocyclic Ga systems relevant to this study, see ref. [4a] and a) E. S. Schmidt, A. Jockisch, H. Schmidbaur, J. Am. Chem. Soc. 1999, 121, 9758–9759; b) R. J. Baker, R. D. Farley, C. Jones, M. Kluth, D. Murphy, Dalton Trans. 2002, 3848–3850; c) C. Jones, P. C. Junk, J. A. Platts, A. Stasch, J. Am. Chem. Soc. 2006, 128, 2206–2207; d) I. L. Fedushkin, A. N. Lukoyanov, G. K. Fukin, S. Y. Ketkov, M. Hummert, H. Schumann, Chem. Eur. J. 2008, 14, 8465–8468; e) I. L. Fedushkin, A. N. Lukoyanov, A. N. Tishkina, G. K. Fukin, K. A. Lysenkov, M. Hummert, H. Schumann, Chem. Eur. J. 2010, 16, 7563–7571; f) Y. Liu, S. Li, X.-J. Yang, O.-S. Li, Y. Xie, H. F. Schafer, B. Wu, J. Organomet. Chem. 2011, 696, 1450–1455; g) D. L. Choong, W. D. Woodul, D. L. Choong, A. Stasch, C. Schenk, C. Jones, Aust. J. Chem. 2011, 64, 1173–1176; h) D. Dange, S. L. Choong, C. Schenk, A. Stasch, C. Jones, Dalton Trans. 2012, 41, 9304–9315; i) A. A. Hawley, C. A. Ohlin, L. Föhmeister, A. Stasch, Chem. Eur. J. 2017, 23, 447–455.

[9] J. Hicks, P. Vasko, J. M. Goicoechea, S. Aldridge, Nature 2018, 557, 2019–2025.

[10] For other examples of aluminium compounds see: a) R. J. Schwamm, M. D. A. Meier, R. H. M. Coles, Angew. Chem. Int. Ed. 2019, 58, 1489–1493; Angew. Chem. 2019, 131, 1503–1507; b) S. Kurumada, S. Takamori, M. Yamashita, Nat. Chem. 2020, 12, 36–39; c) R. J. Schwamm, M. P. Coles, S. M. Hill, M. F. Ahon, C. L. McMullin, N. A. Rajabi, A. S. Wilson, Angew. Chem. Int. Ed. 2020, 59, 3928–3932; Angew. Chem. 2020, 132, 3956–3960; d) K. Koshina, R. Kinjo, J. Am. Chem. Soc. 2020, 142, 9057–9062; e) S. Grams, E. Eyselein, J. Langer, J. Fäber, S. Harder, Angew. Chem. Int. Ed. 2020, 59, 15982–15986; Angew. Chem. 2020, 132, 16116–16120.

Acknowledgements

ERC (studentship to CM), Leverhulme Trust (studentship to AH; RP-2018-246).

Conflict of interest

The authors declare no conflict of interest.
nesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, Williams, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery, Jr., J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman, D. J. Fox, Wallingford, CT, 2016.

[31] a) C. Adamo, V. Barone, J. Chem. Phys. 1999, 110, 6158–6170; b) M. Ernzerhof, G. E. Scuseria, J. Chem. Phys. 1999, 110, 5029–5036; c) J. P. Perdew, K. Burke, M. Ernzerhof, Phys. Rev. Lett. 1996, 77, 3865–3868; d) J. P. Perdew, M. Ernzerhof, K. Burke, J. Chem. Phys. 1996, 105, 9982–9985; e) J. P. Perdew, K. Burke, M. Ernzerhof, Phys. Rev. Lett. 1997, 78, 1396–1396.
[32] A. Schafer, C. Huber, R. Ahlrichs, J. Chem. Phys. 1994, 100, 5829–5835.
[33] a) S. Grimme, J. Antony, S. Ehrlich, H. Krieg, J. Chem. Phys. 2010, 132, 154104; b) S. Grimme, S. Ehrlich, L. Goerigk, J. Comput. Chem. 2011, 32, 1456–1465.
[34] M. D. Hanwell, D. E. Curtis, D. C. Lonie, T. Vandermeersch, E. Zurek, G. R. Hutchison, J. Cheminf. 2012, 4, 17.
[35] E. D. Glendening, A. E. Reid, J. E. Carpenter, F. Weinhold, NBO Version 3.1.

Manuscript received: October 27, 2020
Revised manuscript received: November 16, 2020
Accepted manuscript online: November 17, 2020
Version of record online: January 14, 2021