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Reactivity of halfsandwich rare-earth metal methylaluminates toward potassium (2,4,6-*tert*-butylphenyl)amide and 1-adamantylamine†

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The equimolar reaction of potassium (2,4,6-*tert*-butylphenyl)amide with Cp^{*}Ln(AlMe₄)₂ (Cp^{*} = 1,2,3,4,5-pentamethyl cyclopentadienyl) yielded {Cp^{*}Ln(AlMe₄)[NH(mes^{*})]}_x (Ln = Y, La; mes^{*} = C₆H₂tBu₃-2,4,6). The treatment of Cp^{*}Ln(AlMe₄)[NH(mes^{*})] with tetrahydrofuran led to intramolecular C–H bond activation of a *t*Bu group with the formation of Cp^{*}YMe{NH[C₆H₂tBu₂-2,4-(CMe₂CH₂)-6]}(AlMe₂)(thf). A similar methyl-anilide species Cp^oLuMe{NH[C₆H₂tBu₂-2,4-(CMe₂CH₂)-6]}(AlMe₂) (Cp^o = 2,3,4,5-tetramethyl-1-(8-quinolyl)cyclopentadienyl) with a C–H bond activated ligand backbone formed by the reaction of Cp^oLu(AlMe₄)₂ and K[NH(mes^{*})]. The reactivity of Cp^oY(AlMe₄)₂ toward H₂NAd (Ad = adamantyl) ultimately led to the methyl–amide complex Cp^oYMe{NH(Ad)}(AlMe₃), corroborating the presence of competing deprotonation and donor-induced methylaluminate cleavage reactions. The halfsandwich complexes Cp^oLu(AlMe₄)₂, Cp^{*}Y(AlMe₄)[NH(mes^{*})], Cp^{*}YMe{NH[C₆H₂tBu₂-2,4-(CMe₂CH₂)-6]}(AlMe₂)(thf), Cp^oLuMe{NH[C₆H₂tBu₂-2,4-(CMe₂CH₂)-6]}(AlMe₂), and Cp^oYMe{NH(Ad)}(AlMe₃) as well as the side-product AlMe₃(H₂NAd) were fully characterized by NMR/FTIR spectroscopy, elemental analysis, and X-ray crystallography.

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Introduction

Halfsandwich rare-earth metal bis(hydrocarbyl) complexes¹ have emerged as versatile synthesis precursors (*e.g.*, for hydrido clusters)² and eminent (pre)catalysts for polymerization reactions (*e.g.*, fabrication of syndiotactic polystyrene).³ In 2003, we introduced halfsandwich rare-earth metal bis(methylaluminate) complexes as thermally quite stable variants of the aforementioned bis(alkyl) derivatives.⁴ Upon cationization with fluorinated phenylborane/borate reagents complexes Cp^RLn(AlMe₄)₂ promote the living 1,4-*trans*-selective polymerization of isoprene (synthetic gutta-percha)⁵ and butadiene.⁶ Such catalyst activation proceeds *via* well-established protonolysis and alkyl abstraction pathways.⁷ The feasibility of protonolysis reactions was also demonstrated for the syntheses of metallocenes⁸ as well as silica-grafted

hybrid materials and respective alkoxide/siloxide model complexes.⁹ Furthermore, complexes of the type Cp^RLn(AlMe₄)₂ were successfully applied in C–H bond activation protocols giving access to the first structurally characterized rare-earth metal methylidyne¹⁰ and methylidene complexes.^{11,12} In the following, we adopted the concept of alkylaluminate-based organoaluminium-assisted deprotonation for the synthesis of rare-earth metal imide complexes.¹³ One major finding was that the superbuly monanionic Tp^{*t*Bu,Me} (= hydrotris(3-*tert*-butyl-5-methyl-pyrazolyl)borate) ligand can afford monolanthanide derivatives of the type [(Tp^{*t*Bu,Me})Ln(NAr)(AlHMe₂)] (Ar = C₆H₃Me₂-2,6)¹⁴ and [(Tp^{*t*Bu,Me})Ln(NR)(AlMe₃)] (Ln = Y, Ho; R = *t*Bu, adamantyl).¹⁵ The isolation of rare-earth metal complexes [(Tp^{*t*Bu,Me})Ln=NR(dmap)] (Ln = Y, Ar = C₆H₃Me₂-2,6; Ln = Lu, Ar = C₆H₃(CF₃)₂-3,5) with a terminally bonded imido ligand was achieved by an aluminium-free synthesis procedure.¹⁶ Prior to our studies it has been shown that the deprotonation of rare-earth metal anilide complexes with organoaluminium or alkyl lithium reagents (*e.g.* Scheme 1, V)¹⁷ is a viable strategy toward Ln^{III} imide complexes.^{17,18}

Given the plethora of group 4 imide complexes,¹⁹ rare-earth metal imide chemistry is still in its infancy.²⁰ Mountford and coworkers examined the efficiency of deprotonation reactions utilizing, *e.g.* TiCl₄ or Ti(NMe₂)₄ in amine elimination and transamination/transimination reactions, respectively.²¹ It is noteworthy that such synthesis protocols are not applicable for the rare-earth metals (*vide infra*).

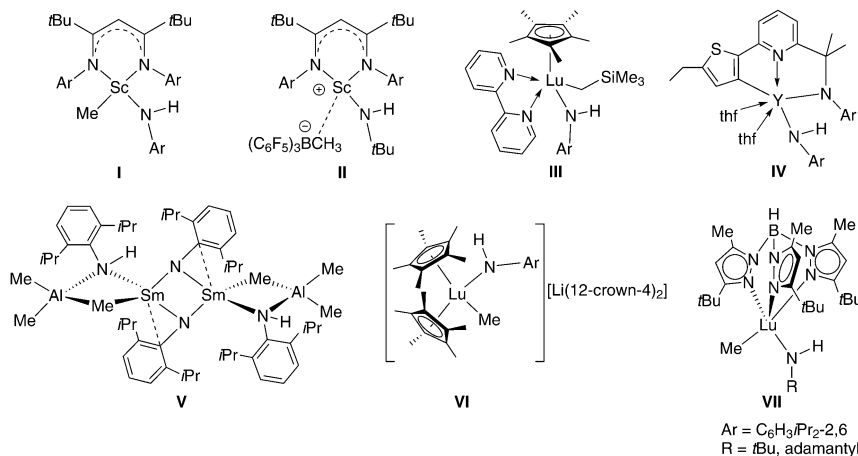
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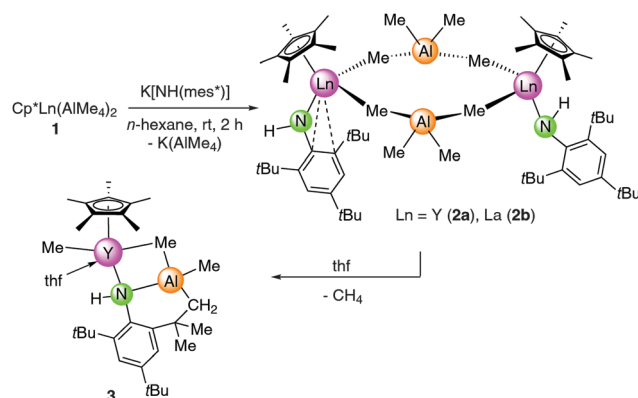
Scheme 1 Structurally characterized rare-earth metal complexes with the amido NH functionality adjacent to Ln-C(alkyl).^{17,22–24}

Based on the successful synthesis of rare-earth metal imide complexes exploiting the synergistic effect associated with organoaluminium moieties and the superbulky monanionic Tp^{tBu,Me}, we have now set out to explore cyclopentadienyl-supported rare-earth metal bis(tetramethylaluminate)s as potential precursors for imide complexes. Herein, we describe the reactivity of Cp*Ln(AlMe₄)₂ and Cp^QLn(AlMe₄)₂ toward potassium (2,4,6-tri-*tert*-butylphenyl)amide and 1-adamantylamine, respectively (Cp* = 1,2,3,4,5-pentamethylcyclopentadienyl, Cp^Q = 2,3,4,5-tetramethyl-1-(8-quinolyl)cyclopentadienyl). Overall, this study discloses a better understanding of why steric/electronic mismatches in isolable mixed amide-alkyl complexes hamper their conversion into the respective imide derivatives.^{22–24} As compiled in Scheme 1, various mixed amide-alkyl complexes (including cationic (II)^{22b} and anionic rare-earth metal entities (VI)^{22c} have been accessed in the presence of a stabilizing N-donor (I,^{22a} II,^{22b} IV,²³ VII,¹⁵) and cyclopentadienyl ancillary ligands (III,²⁴ VI^{22c}).

Results and discussion

Reactivity of Cp*Ln(AlMe₄)₂ (1)

The salt-metathesis reaction of half-sandwich complexes Cp*Ln(AlMe₄)₂ (Ln = Y (1a), La (1b))⁸ with 1 equivalent potassium (2,4,6-tri-*tert*-butylphenyl)amide, K[NH(mes*)], in *n*-hexane at ambient temperature yielded {Cp*Ln(AlMe₄)[NH(mes*)]}_x (Ln = Y (2a), x = 2; Ln = La (2b)) in 83% and 87% yield, respectively (Scheme 2). Rather surprisingly, the reaction came to a halt at the heteroleptic amide-methylaluminate complex and the subsequent deprotonation of the NH anilido functionality did not occur. The mixed amide-imide complex [Sm(NHAr)(AlMe₃)(NAr)]₂¹⁷ is the only other example featuring an NH(Ar) ligand adjacent to an organoaluminium moiety (V, Scheme 1). The latter complex V was obtained from Sm(NHAr)₃ and AlMe₃. For comparison, excess addition of trimethylaluminium to Nd(NHPh)₃ (Ph = phenyl) led to a completely deprotonated imide complex [Nd(AlMe₄)₂(NPh)(AlMe₂)₂] and co-product [(NHPh)AlMe₂]₃ formed by anilido-methyl exchange.^{18a} Aryloxo derivatives [Cp*Ln(AlMe₄)(OAr^{tBu,R})] (Ln = Lu, Y; R = H, Me) reminiscent of complexes 2 were



Scheme 2 Synthesis of anilide complexes 2 and 3.

obtained by the treatment of [Cp*Ln(OAr^{tBu,R})₂] with an excess of trimethylaluminium.^{9b}

Colourless crystals of 2 were sparingly soluble in aliphatic hydrocarbons but soluble in aromatic solvents. The ¹H NMR spectra of complexes 2 show one set of signals for the Cp*, NH(mes*), and AlMe₄ ligands. The five methyl groups of the Cp* ring appear as one singlet at 1.89 and 2.04 ppm in complexes 2a and 2b, respectively. The signals at 5.05 ppm (2a) and 5.83 ppm (2b) can be assigned to the NH functionalities. The metal-bonded methyl groups show one sharp signal at ambient temperature (2a, −0.33 ppm, ²J_{YH} = 2.5 Hz; 2b, −0.43 ppm), which implies a rapid exchange between bridging and terminal methyl groups. For yttrium complex 2a signal splitting of the bridging and terminal methyl groups into two broad singlets at −0.37 ppm and −0.11 ppm occurred at approximately −90 °C. Variable-temperature ¹H NMR studies on lanthanum complex 2b did not reveal any signal splitting at temperatures from +25 to −90 °C indicative of a higher mobility of the La-(AlMe₄) moiety.

Single crystals of {Cp*Y(AlMe₄)[NH(mes*)]}₂ (2a) were grown from saturated *n*-hexane solutions at −35 °C. The X-ray crystallographic analysis revealed a structural motif as found in the solid-state structure of [Cp*₂Ln(AlMe₄)]^{8,25} (Ln = Y, La, Sm) with the yttrium metal centers being bridged by two μ₂-η¹:η¹ coordinated AlMe₄ ligands (Y1-C1-Al1 = 174.8(2)°; Y2-C3-Al1 = 169.3(2)°),



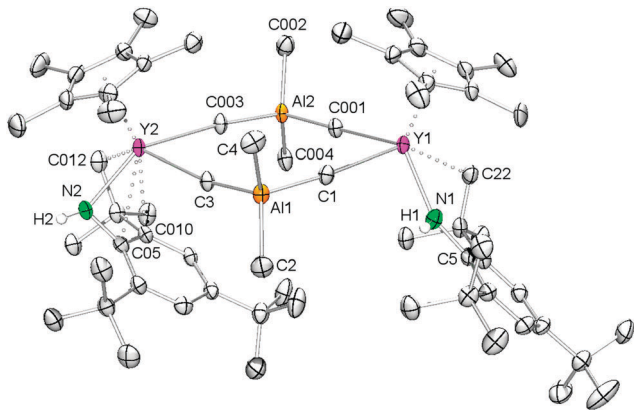


Fig. 1 Molecular structure of **2a**. Atoms are represented by atomic displacement ellipsoids set at the 50% level. Solvent molecules (*n*-hexane) and hydrogen atoms except for the NH protons are omitted for clarity. Selected bond distances [Å] and angles [deg]: Y1–N1 2.186(4), Y2–N2 2.226(3), Y1–C1 2.680(7), Y1–C001 2.653(4), Y2–C3 2.685(4), Y2–C003 2.769(7), Y1···Ct1 2.325, Y2···Ct2 2.328, Y1···C22 2.980(6), Y2···C012 2.819(4), Y2···C05 2.683(4), Y2···C010 2.805(4), Y1–N1–C5 150.7(4), Y2–N2–C05 93.1(3), Y1–C1–Al1 174.8(2), Y2–C3–Al1 169.3(2).

while the Cp* and amido ligands are coordinated terminally (Fig. 1). The NH(mes*) ligands exhibit two distinct coordination modes with the most striking difference being the Y–N(amido)–C(aryl) bond angles of Y1–N1–C5 = 150.7(4)° and Y1–N2–C05 = 93.1(3)°. The Y–N(amido) bond lengths of 2.186(4) Å and 2.226(3) Å are in the expected range.²⁶ The bent amido ligand features additional Ln···arene interactions involving two short contacts to the *ipso* and *ortho* carbon atoms of the aryl ring. Both amido ligands show secondary interactions with the Y(III) center through one of the methyl groups of the *tert*-butyl substituents.

Attempts to synthesize putative Cp*Ln=Nmes* from **2a** *via* Lewis base-induced methane elimination, as described earlier for other alkyl–amide complexes, were not successful.²⁷ Upon addition of thf to **2a** no visual changes were observed, meaning that the mixture remained as a yellow solution. However, evaporation of the solvent, drying *in vacuo*, and recrystallization of the residue from *n*-hexane produced single crystals of complex **3**, revealing AlMe₄ cleavage and C–H bond activation of one of the *tert*-butyl groups (Scheme 2, Fig. 2). The generated monomeric complex is 7-coordinate by the C–H activated primary amido ligand, (μ₂-Me)AlMe[NH[C₆H₂tBu₂-2,4-(CMe₂CH₂)-6]], a η⁵-Cp* ligand, a thf molecule, and a terminal methyl group. The Y–N(amido) bond length of 2.489(2) Å is significantly elongated compared to **2a** (Y–N(amido) = 2.186(4)/2.226(3) Å), which is due to the formation of a four-membered Y–(μ₂-Me)–Al–N metallacycle. This four-membered metallacycle is fused *via* the Al–N bond to a six-membered metallacycle, which itself is annulated to the (anilido)phenyl ring. The Y–C bond lengths differ markedly with the terminal Y–C(CH₃) distance being quite short (2.406(3) Å) similar to the diketiminate complex (L)YMe₂ (L = MeC(N(C₆H₃iPr₂-2,6))CHC(Me)NCH₂CH₂N(Me)CH₂CH₂NMe₂) (av. 2.433 Å)²⁸ and hydrotris(pyrazolyl)borate compound (Tp^{Bu₃Me})YMe(AlMe₄) (Y–C(CH₃) = 2.382(3) Å; Y–C(μ₂-CH₃) = 2.715(3) Å),²⁹ while the Y–C(μ₂-CH₃) (2.827(3) Å) distance is relatively long in

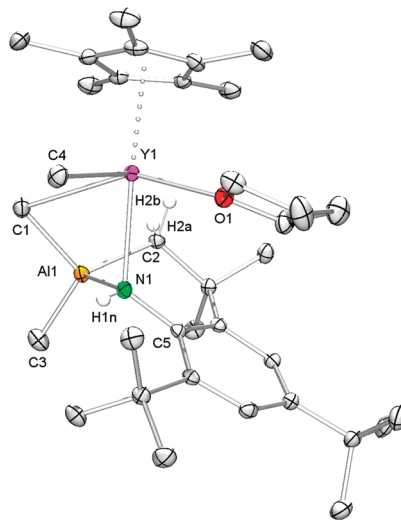


Fig. 2 Molecular structure of **3**. Atoms are represented by atomic displacement ellipsoids set at the 30% level. Hydrogen atoms except for the NH proton and the methylene protons are omitted for clarity. Selected bond distances [Å] and angles [deg]: Y1–C1 2.827(3), Y1–C4 2.406(3), Y1–N1 2.489(2), Y1–O1 2.375(2), Y1···Ct1 2.384, Y1···Al1 3.0705(9), Al1–N1 1.902(2), Al1–C1 2.014(3), Al1–C2 1.985(3), Al1–C3 1.962(3), Y1–N1–C5 123.6(2), C1–Y1–C4 80.4(1), C1–Y1–O1 151.64(8), C4–Y1–O1 92.38(9), N1–Y1–Ct1 162.03, Y1–C1–Al1–N1 39.6(1).

comparison to **2a** (av. 2.697 Å) and [Cp*₂Y(AlMe₄)₂]^{25a} (2.66 Å). Unfortunately, ¹H and ¹³C NMR spectroscopic investigations were hampered by fast decomposition of the product.

Reactivity of Cp^QLn(AlMe₄)₂ (**4** and **6**)

In order to enhance the steric pressure at the rare-earth metal center during the tetramethylaluminate displacement we decided to use the N-donor substituted bis(tetramethylaluminate) precursor Cp^QLn(AlMe₄)₂ (Ln = Y, Lu). The presence of the rigid quinoyl functionality was anticipated to minimize any undesired C–H bond activation at the N-donor substituent.³² Cp^QLu(AlMe₄)₂ was prepared by a slightly modified procedure than described earlier for the yttrium and lanthanum congeners.³³

Accordingly, quinoyl-substituted half-sandwich complex Cp^QLu(AlMe₄)₂ (**4**) was obtained *via* protonolysis of homoleptic Lu(AlMe₄)₃ with HCp^Q. Brown powdery **4** is readily soluble in toluene, but only sparingly soluble in *n*-hexane. The ¹H NMR spectrum of **4** in C₆D₆ at ambient temperature shows one signal for the AlMe₄ ligands at –0.13 ppm, two singlets for the cyclopentadienyl methyl groups at 1.78 and 2.02 ppm and one set of signals for the quinoyl moiety.

Single crystals of **4** suitable for X-ray structure analysis were grown from a toluene–*n*-hexane mixture at –35 °C. As for the yttrium and lanthanum congeners³³ the Cp^Q ligand binds to the metal center in an η⁵ fashion through all five carbon atoms and *via* the quinoyl nitrogen atom (Fig. 3). Not surprisingly, in the sterically more congested lutetium complex one of the AlMe₄ ligands coordinates in an η¹ fashion while the other exhibits the usually observed planar η² coordination mode (Lu1–C24–Al2–C23 = 5.16(8)°), accounting for an overall hepta-coordinate Lu(III) metal center. Such η¹ coordination mode of



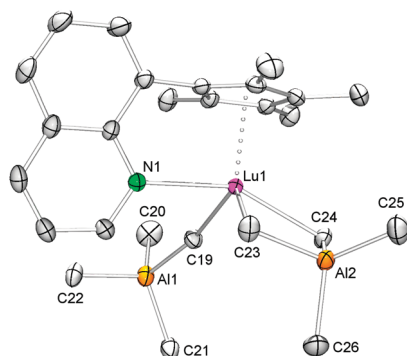


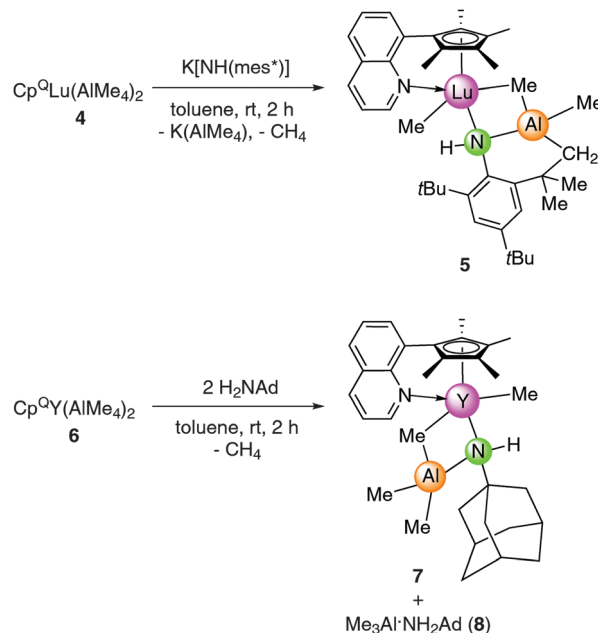
Fig. 3 Molecular structure of **4**. Atoms are represented by atomic displacement ellipsoids set at the 50% level. Hydrogen atoms are omitted for clarity. Selected bond distances [Å] and angles [deg]: Lu1–N1 2.417(1), Lu1–C19 2.539(2), Lu1–C23 2.485(2), Lu1–C24 2.524(2), Lu1···Ct1 2.247, Lu1···Al1 4.5877(6), Lu1···Al2 3.0296(5), Al1–C19 2.081(2), Al1–C20/21/22 1.991(2)/2.001(2)/1.997(2), Al2–C23 2.081(2), Al2–C24 2.069(2), Al2–C25/26 1.964(2)/1.968(2), N1–Lu1–Ct1 96.57, C19–Lu1–Ct1 116.25, C23–Lu1–Ct1 120.30, C24–Lu1–Ct1 110.86, Lu1–C19–Al1 166.51(9), Lu1–C23–Al2 82.63(6), Lu1–C24–Al2 81.87(6), Lu1–C24–Al2–C23 5.16(8).

the AlMe_4 ligand with an almost linear Lu–C(μ -CH₃)–Al bond angle of 166.51(9)° has been observed before in $(\text{Tp}^{\text{tBu,Me}})\text{YMe}(\text{AlMe}_4)$ (Y1–C(μ -CH₃)–Al = 161.0(1)°)²⁹ and $[(\text{ArNMe}_2)_2\text{NC}_5\text{H}_3]\text{La}(\text{AlMe}_4)(\text{thf})$ (Ar = C₆H₃iPr₂-2,6, La1–C(μ -CH₃)–Al = 165.0(4)°),³⁰ supported by sterically demanding tridentate (NNN)ⁿ⁻ ligands ($n = 1, 2$).

Interestingly, the salt metathesis reaction of $\text{Cp}^{\text{Q}}\text{Lu}(\text{AlMe}_4)_2$ (**4**) and $\text{K}[\text{NH}(\text{mes}^*)]$ led to complex $\text{Cp}^{\text{Q}}\text{LuMe}\{\text{NH}[\text{C}_6\text{H}_2\text{tBu}_2\text{-2,4}(\text{CMe}_2\text{CH}_2)\text{-6}]\}(\text{AlMe}_2)$ (**5**) (Scheme 3) revealing a structural motif reminiscent of that found in **3**, with the hard quinolyl donor now occupying the position of the thf coordination site. In the ¹H NMR the proton of the NH functionality was clearly evidenced by a singlet at 4.76 ppm, while the presence of three singlets at 0.21, –0.25, and –0.35 ppm for the metal bonded methyl groups and two doublets at 0.85 and 0.67 ppm with a geminal coupling constant of $^2J_{\text{HH}} = 14.5$ Hz for the methylene group are consistent with the solid-state structure.

An X-ray diffraction study of **5** revealed that the lutetium metal center is coordinated by the Cp^Q ligand *via* the η^5 -C₅Me₄ moiety and the quinolyl nitrogen donor, one terminal and one bridging methyl group as well as the aluminium-linked amido ligand resulting in an overall coordination number of seven (see Fig. 4). The Lu–C(CH₃) and Lu–C(μ -CH₃) bond lengths of 2.334(2) Å and 2.703(2) Å, respectively, and the Lu–N(amido) distance of 2.479(2) Å compare well to those detected in complex **3** considering the difference of the Ln(III) ionic radii. For comparison, complexes $\text{Cp}^*_2\text{LuMe}_2\text{Li}(\text{thf})_3$ ^{22c} and $[\text{Cp}^*_2\text{LuMe}_2]_3$ ³¹ show similar terminal Lu–C(CH₃) bond lengths of 2.361(9) Å and 2.423(3) Å, respectively, whereas the bridging carbon bonds Lu–C(μ -CH₃) can differ markedly ranging from 2.385(9)^{22c} to 2.737(3) Å.³¹ The bending of the nonlinear Lu1–N1–C32(*ipso*) fragment of 129.0(1)° in **5** is much more pronounced than in the related complex $[\text{Cp}^*_2\text{LuMe}(\text{NHAr})][\text{Li}(12\text{-crown-4})_2]$ (Ar = C₆H₃iPr₂-2,6) (Lu–N = 2.245(4) Å, Lu–N–C = 155(3)°).^{22c}

Remarkably, the reaction of $\text{Cp}^{\text{Q}}\text{Y}(\text{AlMe}_4)_2$ with H₂Nmes* or $\text{K}[\text{NH}(\text{mes}^*)]$ led to the formation of several metal-containing



Scheme 3 Synthesis of anilide complexes **5** and **7**.

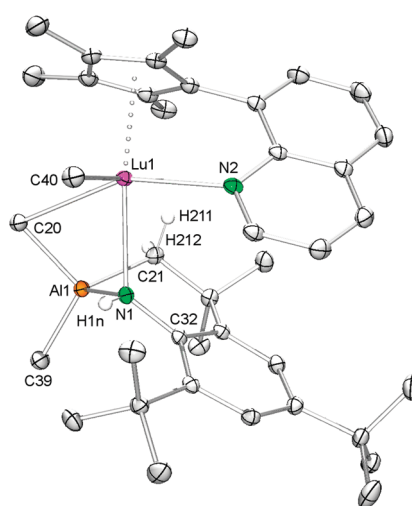


Fig. 4 Molecular structure of **5**. Atoms are represented by atomic displacement ellipsoids set at the 30% level. Hydrogen atoms except for the NH proton and the methylene protons are omitted for clarity. Selected bond distances [Å] and angles [deg]: Lu1–C20 2.703(2), Lu1–C40 2.334(2), Lu1–N1 2.479(2), Lu1–N2 2.401(2), Lu1···Ct1 2.339, Lu1···Al1 2.9694(6), Al1–N1 1.905(2), Al1–C20 2.023(2), Al1–C21 1.991(2), Al1–C39 1.964(2), Lu1–N1–C32 129.0(1), C20–Lu1–C40 89.45(8), C20–Lu1–N2 154.45(6), C20–Lu1–N1 71.84(5), C40–Lu1–N2 97.17(6), C40–Lu1–N1 86.91(6), N1–Lu1–Ct1 165.11, Lu1–C20–Al1–N1 41.27(7).

species. However, we have recently shown that protonolysis of complexes $(\text{Tp}^{\text{tBu,Me}})\text{YMe}(\text{MMe}_4)$ (M = Al, Ga) with primary amines/anilines, is also a viable strategy to access imide species $(\text{Tp}^{\text{tBu,Me}})\text{Ln}=\text{NR}$ or $(\text{Tp}^{\text{tBu,Me}})\text{Ln}(\text{NR})(\text{AlMe}_3)$.^{15,16} For assessing the feasibility of the protonolysis protocol for Cp-based systems, we reacted $\text{Cp}^{\text{Q}}\text{Y}(\text{AlMe}_4)_2$ with 1-adamantylamine (H₂NAd). Primary amine H₂NAd was selected in order to minimize any C–H bond activation at the ligand backbone. Accordingly, the reaction of



6 with H₂NAd afforded amide complex Cp^QYMe[NH(Ad)](AlMe₃) (**7**) together with adduct Me₃Al-NH₂(Ad) (**8**) (Scheme 3). As anticipated C–H bond activation of the amide ligand did not occur. The observed reactivity comes not as a total surprise since the reaction of Ln(AlMe₃)₂ (Ln = Y, Lu) with 1-adamantylamine was previously shown to produce [LnMe₃]_n and Me₃Al-NH₂(Ad). Such distinct reactivity clearly shows that the type of ancillary ligand (Cp *versus* Tp = tris(pyrazolyl)borate) affects organoaluminium-assisted deprotonation^{10,13} and Lewis base-induced aluminate cleavage³⁴ as competing reaction pathways and the action of primary amines as Brønsted acids or Lewis bases.

Anilide species **7** is sparingly soluble in aliphatic hydrocarbons but soluble in aromatic solvents. The ¹H NMR spectrum of **7** shows the expected set of signals for the coordinated Cp^Q ligand. The four methyl groups of the Cp ring appear as two singlets at 1.78 and 2.20 ppm and the signals of the quinoyl protons were found in the range from 6.57 to 8.88 ppm. All signals are shifted 0.1–0.3 ppm upfield compared to those of **6**, indicating the coordination of the adamantylamide ligand to the rare-earth metal center. The resonances assignable to the adamantyl methylene and methine groups were detected as a doublet at 2.28 ppm (6H), a multiplet ranging from 1.79 to 1.61 ppm (6H) and a broad singlet at 2.14 ppm (3H), respectively. Only one doublet was observed for the metal-bonded methyl groups in complex **7** at ambient temperature (−0.06 ppm, 12H), which implies a high mobility of the Y/Al-CH₃ moieties. Such a facile exchange of a terminal Y-CH₃ and a bridging methyl group Y-CH₃-M has been observed before in (Tp^{tBu,Me})YMe[(μ-Me)(GaMe₃)].¹⁶ The open coordination sphere at the yttrium metal center seems to enable enhanced methyl group mobility.

Single crystals of **7** suitable for X-ray structure analysis were grown from toluene-*n*-hexane mixtures at −35 °C. In the solid state, complex **7** is heptacoordinate by one terminal and one bridging methyl group, the amido nitrogen atom as well as the Cp^Q ligand binding *via* the Cp ring (η⁵) and the nitrogen atom of the quinoyl moiety (see Fig. 5). The amido ligand and the

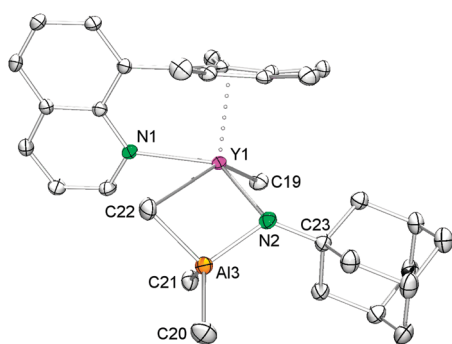
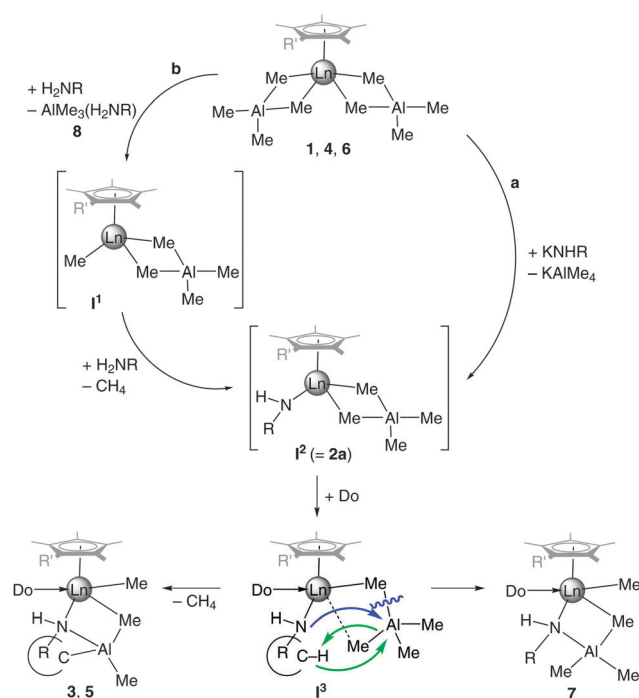


Fig. 5 Molecular structure of **7**. Atoms are represented by atomic displacement ellipsoids set at the 50% level. Selected bond distances [Å] and angles [deg]: Y1–C19 2.392(2), Y1···C22 3.006(2), Y1–N1 2.527(2), Y1–N2 2.428(2), Y1···Ct1 2.366, Y1···Al3 3.0417(6), Al3–C20 1.964(2), Al3–C21 2.007(2), Al3–C22 2.022(2), Al3–N2 1.920(2), Y1–N2–C23 126.9(1), Y1–N2–Al3 87.98(7), C19–Y1–C22 144.61(7), N1–Y1–N2 140.16(6), C19–Y1–N2 103.26(7), C19–Y1–N1 95.24(7), C22–Y1–N1 78.49(6), Y1–N2–Al3–C22 −55.70(9).

(μ-Me)AlMe₂ moiety form a heterobimetallic four-membered ring, which is considerably bent (torsion angle Y1–N2–Al3–C22 −55.70(9)°), resulting in a smaller Y···Al separation of 3.0417(6) Å than in precursor **6** (av. 3.1784(7) Å).³³ The mean metal–ring–carbon distances Ln–C(Cp^Q) (av. 2.656(2) Å) and the Y–N(quinoyl) bond length (2.527(2) Å) are similar to those found in **6**.³³ The bond length of the terminal methyl group Y–C(CH₃) (2.392(2) Å) compares well to that found in **3** while the distance of the bridging Y–C(μ-CH₃) moiety (3.006(2) Å) is significantly longer than the one found in **3** (Y1–C1 = 2.827(3) Å) and markedly longer than the average distances Y–C(CH₃) (2.645(2) Å) in homoaluminate precursor **6**.³³ The Y–N(amido) bond length of 2.428(2) Å is slightly shorter than in **3**, reflecting a less sterically crowded coordination environment around the metal center.

In Scheme 4, we propose a plausible mechanistic scenario for the formation of half-sandwich amide complexes **3**, **5**, and **7**: both salt metathesis involving Cp^RLn(AlMe₃)₂ and KNHR (Scheme 4a) and protonolysis involving Cp^RLn(AlMe₃)₂ and H₂NR (Scheme 4b) afford intermediate **I**², which in the absence of donor functionalities can be isolated as shown for **2a** and **2b**. The protonolysis reaction is preceded by an amine-induced tetramethylaluminate cleavage and the formation of a transient terminal Ln–CH₃ moiety (**I**¹), which reacts instantly with another amine molecule *via* methane elimination to generate **I**². The occurrence of donor-induced tetramethylaluminate cleavage as the rate-determining step along reaction path **b** is supported by the formation of considerable amounts of **7** along with AlMe₃(H₂NR) (**8**) in equimolar reactions of **6** and H₂NAd. Intermediate **I**² is not stable in the presence of donor functionalities (quinoyl or thf), which can exert enhanced steric pressure on the amido as well as tetramethylaluminate ligands. As a consequence the amido



Scheme 4 Proposed reaction mechanism for the formation of **3**, **5**, and **7**.



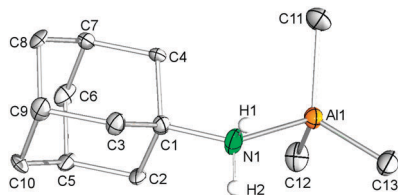


Fig. 6 Molecular structure of **8**. Atoms are represented by atomic displacement ellipsoids set at the 50% level. The disorders in the adamantyl moiety and hydrogen atoms except for the NH₂ protons are omitted for clarity. Selected bond distances [Å] and angles [deg]: Al1–C11 1.988(2), Al1–C12 1.976(2), Al1–C13 1.986(2), Al1–N1 2.032(1), Al1–N1–C1 126.41(9), C12–Al1–C13 113.32(9), C12–Al1–C11 116.32(8), C13–Al1–C11 113.38(9), C12–Al1–N1 107.25(8), C13–Al1–N1 99.39(6), C11–Al1–N1 105.13(8).

nitrogen can approach the aluminate ligand (blue arrow), which might experience a η^2 – η^1 coordination switch, ultimately leading to an aluminate cleavage and concomitantly to the formation of a stable Al–N bond as well as a (second) Ln–CH₃ moiety (isolation of complex **7**). In the case of proximal alkyl moieties, the organo-aluminium fragment can trigger C–H bond activation (green arrows), as evidenced for the formation of complexes **3** and **5**.

Synthesis and characterization of Me₃Al–NH₂(Ad) (**8**)

Byproduct **8**, which could be separated from **7** by extraction with *n*-hexane, was independently synthesized in quantitative yield *via* the addition of AlMe₃ to a toluene solution of 1-adamantylamine. Recrystallization from toluene–*n*-hexane at –35 °C gave colourless crystals. In the solid-state, the aluminium center adopts a distorted tetrahedral coordination geometry (Fig. 6). The average Al–C(CH₃) bond length of 1.983 Å and the Al–N(amine) distance of 2.032(1) Å are in accord with the corresponding metrical parameters observed for the related complex Me₃Al–NH₂(*t*Bu)³⁵ (av. Al–C = 1.96 Å, Al–N = 2.027(3) Å) and **7** (av. Al–C = 1.998 Å), but are considerably elongated compared to the Al–N(amido) bond length in **7** (Al–N = 1.920(2) Å). As expected, the ¹H NMR spectrum of aluminium complex **8** in C₆D₆ shows one high-field signal for the methyl groups at –0.36 ppm (9H), a singlet for the NH₂ functionality at 1.59 ppm (2H), a broad singlet for the methine groups (3H), a doublet at 1.20 ppm and a multiplet at 1.27 ppm for the methylene groups integrating six protons each, respectively.

Conclusion

The reactions of monocyclopentadienyl rare-earth metal bis-(tetramethylaluminate) complexes with 1-adamantylamine or potassium (2,4,6-tri-*tert*-butylphenyl)amide yield half-sandwich mixed alkyl–amide complexes. The reactivity of the metal-bonded alkyl groups toward adjacent NH functionalities appears strongly reduced and further deprotonation was not observed. This is in sharp contrast to previously examined complexes (Tp^{*t*Bu,Me}–)YMe(AlMe₄), which do react with primary amines/anilines to afford Lewis-acid stabilized imido derivatives (Tp^{*t*Bu,Me}–)Ln(NR)–(AlMe₃).¹⁵ Such changed reactivity of complexes Cp*Ln(AlMe₄)₂ and Cp^QLn(AlMe₄)₂ might be caused by a steric misfit of the metal–alkyl and N–H groups, since the former still exhibit high

reactivity as shown for the C–H activation of the NH(mes*) amido ligand backbone. Though substitution of mes* by adamantyl decreases the reactivity of the NH moiety (pK_a criterion), the occurrence of undesired side-reactions is impeded. Previous studies from the Hessen and Hou groups revealed that imides could only be obtained from reactive Cp ligated rare-earth metal hydride and diene complexes *via* the insertion of benzonitrile.^{2b,36} Overall, the present study reinforces that primary alkylamines not only act as protic substrates but also as Lewis bases and that the steric and electronic properties of the ancillary ligands are crucial factors directing the formation of either heteroleptic alkyl–amide complexes or imide species.¹⁵

Experimental section

General considerations

All operations were performed with rigorous exclusion of air and water, using standard Schlenk, high-vacuum, and glovebox techniques (MBraun 200B; <1 ppm O₂, <1 ppm H₂O). THF, toluene and *n*-hexane were purified by using Grubbs columns (MBraun SPS, solvent purification system) and stored inside a glovebox. C₆D₆ and toluene-*d*₈ were obtained from Aldrich, degassed, dried over Na for 24 h, and filtered. 1-Adamantylamine was received from abcr and used as received. Pro-ligand HCp^Q,³⁷ heteroleptic complexes Cp*Ln(AlMe₄)₂ (Ln = Y, La) (**1**)⁸ and Cp^QY(AlMe₄)₂ (**6**),³³ and potassium (2,4,6-Tri-*tert*-butylphenyl)amide¹³ were synthesized according to literature procedures. The NMR spectra of air and moisture sensitive compounds were recorded by using J. Young valve NMR tubes at variable temperature on a Bruker AVII+400 (¹H: 400 MHz; ¹³C: 101 MHz) and a Bruker AVII+500 (¹H: 500 MHz; ¹³C: 126 MHz). ¹H and ¹³C shifts are referenced to internal solvent resonances and reported in parts per million relative to TMS. Coupling constants are given in Hertz. DRIFT spectra were recorded on a NICOLET 6700 FTIR spectrometer using dried KBr and KBr windows. Elemental analyses were performed on an Elementar Vario Micro Cube.

General procedures for the synthesis of

{Cp*Ln(AlMe₄)[NH(mes*)]}_x

A solution of (Cp*)Ln(AlMe₄)₂ in *n*-hexane (2 mL) was added to a stirred suspension of potassium (2,4,6-tri-*tert*-butylphenyl)amide K[NH(mes*)] in *n*-hexane (2 mL). The reaction mixture was stirred for 2 h at ambient temperature and the *n*-hexane solution then separated by centrifugation, decanted, and filtrated. The solid residue product and K(AlMe₄) were extracted with additional *n*-hexane (2 × 2 mL). Compounds **2** were obtained as powders or by crystallization from *n*-hexane solutions at –35 °C.

{Cp*Y(AlMe₄)[NH(mes*)]}₂ (**2a**). Following the procedure described above, (Cp*)Y(AlMe₄)₂ (80 mg, 0.20 mmol) and K[NH(mes*)] (60 mg, 0.20 mmol) yielded **2a** as colourless crystals (121 mg, 0.20 mmol, quant., 83% crystalline yield). ¹H NMR (500 MHz, C₆D₆, 26 °C, TMS): δ = 7.38 (s, 4H, H_{aryl}), 5.05 (s, 2H, NH), 1.89 (s, 30H, CH₃, Cp*), 1.44 (s, 18H, (C(CH₃)₃)_{para}), 1.41 (s, 36H, (C(CH₃)₃)_{ortho}), –0.33 (d, 24H, ²J_{YH} = 2.5 Hz, Al(CH₃)₄). ¹³C NMR (126 MHz, C₆D₆, 26 °C) δ = 151.3 (Ar, C_{ipso}),



138.0 (Ar, C_{para}), 133.0 (Ar, C_{ortho}), 123.4 (Ar, C_{meta}), 121.2 ($C_5(CH_3)_5$), 35.8 ($(C(CH_3)_3)_{ortho}$), 34.9 ($(C(CH_3)_3)_{para}$), 32.4 ($(C(CH_3)_3)_{para}$), 31.3 ($(C(CH_3)_3)_{ortho}$), 12.0 ($C_5(CH_3)_5$), 1.1 ($Al(CH_3)_4$) ppm. DRIFT (KBr): 3416vw (NH), 2959vs, 2912s, 2871s, 1594vw, 1476w, 1456m, 1419s, 1393m, 1361m, 1280w, 1240s, 1195m, 1119w, 1024w, 931w, 912w, 879m, 835m, 778m, 745m, 719m, 677m, 624m, 565m, 545m, 480w, 449w cm^{-1} . Elemental analysis calcd (%) for $C_{32}H_{57}AlNY$ (571.69 $g\ mol^{-1}$): C 67.23, H 10.05, N 2.45; found: C 67.26, H 10.13, N 2.32.

{Cp*La(AlMe₄)[NH(mes*)]}_x (2b). Following the procedure described above, (Cp*)La(AlMe₄)₂ (90 mg, 0.20 mmol) and K[NH(mes*)] (60 mg, 0.20 mmol) yielded **2b** as a light yellow powder (108 mg, 0.17 mmol, 87%). ¹H NMR (500 MHz, C₆D₆, 26 °C, TMS): δ = 7.38 (s, 2H, H_{aryl}), 5.83 (s, 1H, NH), 2.04 (s, 15H, CH₃, Cp*), 1.42 (s, 9H, (C(CH₃)₃)_{para}), 1.33 (s, 18H, (C(CH₃)₃)_{ortho}), -0.43 (s br, 12H, Al(CH₃)₄). ¹³C NMR (126 MHz, C₆D₆, 26 °C) δ = 150.2 (Ar, C_{ipso}), 139.9 (Ar, C_{para}), 134.6 (Ar, C_{ortho}), 125.3 (Ar, C_{meta}), 123.4 ($C_5(CH_3)_5$), 36.5 ($(C(CH_3)_3)_{ortho}$), 35.0 ($(C(CH_3)_3)_{para}$), 32.3 ($(C(CH_3)_3)_{para}$), 31.4 ($(C(CH_3)_3)_{ortho}$), 11.8 ($C_5(CH_3)_5$), 2.6 ($Al(CH_3)_4$) ppm. DRIFT (KBr): 3405vw (NH), 2963vs, 2909vs, 2870s, 2835m, 1597vw, 1498vw, 1477w, 1464w, 1416s, 1393m, 1383m, 1362m, 1279m, 1251m, 1241m, 1220w, 1202w, 1175m, 1114w, 1023vw, 986m, 960s, 883vw, 825w, 809w, 783w, 768w, 727s, 674w, 641vw, 610s, 555s, 539s, 501vw, 478vw, 419w cm^{-1} . Elemental analysis calcd (%) for $C_{32}H_{57}AlLaN$ (621.69 $g\ mol^{-1}$): C 61.82, H 9.24, N 2.25; found: C 61.58, H 9.60, N 1.98.

Cp*YMe{NH[C₆H₂tBu₂-2,4-(CMe₂CH₂)-6]}(AlMe₂)(thf) (3). Tetrahydrofuran (1 ml) was added to solid **2a** (57 mg, 0.10 mmol) at ambient temperature. The obtained yellow solution was dried *in vacuo* and recrystallized in *n*-hexane at -35 °C (estimated yield 29 mg, 46%). Single crystals of **3** suitable for X-ray diffraction were harvested from saturated *n*-hexane solutions at -35 °C. ¹H NMR measurements were hampered by rapid decomposition of **3**. DRIFT (KBr): 3402vw (NH), 2958vs, 2927s, 2907s, 2887s, 2741w, 1457m, 1423s, 1394w, 1361m, 1335w, 1286w, 1228s, 1189w, 1157w, 1120m, 1103w, 1014m, 921vw, 878w, 863w, 830w, 814w, 778w, 752s, 736m, 695s, 627w, 559w, 520vw, 473w cm^{-1} . Elemental analysis calcd (%) for $C_{35}H_{61}AlNOY$ (627.75 $g\ mol^{-1}$): C 66.97, H 9.79, N 2.23; found: C 67.37, H 10.24, N 2.01.

Cp^QLu(AlMe₄)₂ (4). A solution of Lu(AlMe₄)₃ (87 mg, 0.20 mmol) in *n*-hexane (3 mL) was added to a stirred suspension of HCp^Q (50 mg, 0.20 mmol) in *n*-hexane (1 mL). Instant gas formation was observed and the solution turned yellow with a thick, brown slurry at the bottom. The reaction mixture was stirred another 3 h at ambient temperature and then dried under vacuum. To the residue toluene (2 mL) was added and the brown solution was stirred for another 3 h at ambient temperature. The product was dried *in vacuo* and washed with *n*-hexane (5 × 2 mL). The obtained brown solid was dried under reduced pressure (113 mg, 95%). Crystals of **4** were obtained from a saturated, yellow toluene-*n*-hexane solution at -35 °C. ¹H NMR (500 MHz, C₆D₆, 26 °C, TMS): δ = 8.31 (d, 1H, ³J_{HH} = 4.0 Hz, quin-*H*), 7.32 (d, 1H, ³J_{HH} = 8.0 Hz, quin-*H*), 7.29 (dd, 1H, ³J_{HH} = 3.9 Hz and 2.3 Hz), 7.12–7.08 (m, 2H, quin-*H*), 6.53 (m, 1H, quin-*H*), 2.02 (s, 6H, Cp, CH₃), 1.78 (s, 6H, Cp, CH₃), -0.13 (s, 24H, Al-CH₃) ppm. ¹³C{¹H} NMR (126 MHz, C₆D₆, 26 °C) δ = 151.3 (quin-*C*),

150.7 (quin-*C*), 140.2 (quin-*C*), 135.4 (quin-*C*), 134.0 (quin-*C*), 129.8 (quin-*C*), 127.4 (quin-*C*), 123.0 (quin-*C*), 121.9 (quin-*C* and Cp), 121.3 (Cp), 120.0 (Cp), 12.3 (Cp CH₃), 11.5 (Cp CH₃), 2.2 (Al-CH₃) ppm. DRIFT (KBr): 2915s, 2886m, 2816w, 1593w, 1510m, 1435w, 1371w, 1302w, 1220w, 1190m, 1025vw, 970vw, 844w, 824w, 790m, 764w, 706vs, 578m, 519w, 478w cm^{-1} . Elemental analysis calcd (%) for $C_{26}H_{42}Al_2LuN$ (597.55 $g\ mol^{-1}$): C 52.26, H 7.08, N 2.34; found: C 53.43, H 7.36, N 2.54. Multiple attempts to obtain a better microanalysis failed.

Cp^QLuMe{NH[C₆H₂tBu₂-2,4-(CMe₂CH₂)-6]}(AlMe₂) (5)

To a stirred suspension of K[NH(mes*)] in toluene (1 mL) a solution of **4** (87 mg, 0.14 mmol) in toluene (3 mL) was added. The brown suspension was stirred 3 h at ambient temperature and the product then separated by centrifugation, decanted, and filtrated. The solid residue product and K(AlMe₄) was extracted with additional toluene (2 × 1 ml). The combined extracts were dried and washed with *n*-hexane (3 × 2 ml), followed by drying under reduced pressure (96 mg, 0.12 mmol, 91%, 30% crystalline yield). Crystallization from a toluene-*n*-hexane solution at -35 °C afforded yellow crystals of **5** suitable for X-ray diffraction analysis. ¹H NMR (500 MHz, C₆D₆, 26 °C, TMS): δ = 8.03 (d, 1H, ³J_{HH} = 4.0 Hz, quin-*H*), 7.67 (d, 1H, ³J_{HH} = 2.3 Hz, quin-*H*), 7.45–7.43 (m, 3H, quin-*H*), 7.30 (d, 1H, ³J_{HH} = 8.5 Hz, quin-*H*), 7.20 (s, 1H, Ar), 7.13–7.00 (m, 2H, quin-*H* and Ar), 6.41 (dd, 1H, ³J_{HH} = 8.1 Hz and 3.2 Hz), 4.76 (s, 1H, NH), 2.28 (s, 3H, Cp CH₃), 2.06 (s, 3H, Cp CH₃), 2.04 (s, 3H, Cp CH₃), 1.84 (s, 3H, Cp CH₃), 1.53 (s, 9H, C(CH₃)₃), 1.49 (s, 6H, C(CH₃)₂), 1.36 (s, 9H, C(CH₃)₃), 0.85 (d, 1H, ²J_{HH} = 14.5 Hz, Al-CH₂), 0.67 (d, 1H, ²J_{HH} = 14.5 Hz, Al-CH₂), 0.21 (s, 3H, Lu-CH₃), -0.25 (s, 3H, Al-CH₃), -0.35 (s, 3H, Al-CH₃) ppm. ¹³C{¹H} NMR from ¹H¹³C HSQC and HMBC (126 MHz, C₆D₆, 26 °C) δ = 152.3 (quin-*C*), 150.1 (quin-*C*), 145.1 (quin-*C*), 142.2 (Ar), 140.2 (Ar), 138.4 (Ar), 137.3 (Ar), 139.5 (quin-*C*), 128.6 (Ar, quin-*C*, overlapping with solvent signal), 126.4 (Ar, quin-*C*), 124.6 (Ar), 124.0 (quin-*C*), 122.5 (quin-*C*), 121.0 (quin-*C*), 119.2 (Cp), 117.4 (Cp), 38.8 (CH₃), 37.9 (C(CH₃)₃), 34.9 (C(CH₃)₂), 34.2 (C(CH₃)₃), 32.4 (CH₃), 31.9 (C(CH₃)₃), 31.5 (C(CH₃)₃), 29.4 (CH₂), 28.8 (Al-CH₃), 0.9 (Lu-CH₃), -9.2 (Al-CH₃) ppm. DRIFT (KBr): 3394vw (NH), 3348vw, 3045vw, 2951vs, 2902s, 2865s, 2754w, 1603vw, 1586vw, 1506m, 1475w, 1467w, 1423s, 1396w, 1362w, 1334vw, 1303vw, 1281vw, 1261w, 1237w, 1223m, 1183w, 1159w, 1119m, 841w, 829w, 821w, 806m, 788m, 776m, 757m, 738w, 699m, 687m, 656w, 648w, 626vw cm^{-1} . Elemental analysis calcd (%) for $C_{39}H_{56}AlLuN_2$ (754.82 $g\ mol^{-1}$): C 62.06, H 7.48, N 3.71; found: C 60.53, H 5.92, N 3.72. Multiple attempts to obtain a better microanalysis failed.

Cp^QYMe{NH(Ad)}(AlMe₃) (7). A solution of 1-adamantylamine (30 mg, 0.20 mmol) in toluene (2 mL) was added to a solution of Cp^QY(AlMe₄)₂ (51 mg, 0.10 mmol) in toluene (1 mL). The resulting yellow-brown solution was stirred for 1 h at ambient temperature, then dried under reduced pressure, washed with *n*-hexane (2 × 2 ml), and dried *in vacuo* (57 mg, 0.10 mmol, quant.). Crystallization from a toluene-*n*-hexane solution at -35 °C afforded yellow crystals of **7** suitable for X-ray diffraction analysis. ¹H NMR (400 MHz, C₆D₆, 26 °C, TMS): δ = 8.88 (dd, 1H, ³J_{HH} = 4.9 and 1.5 Hz, quin-*H*), 7.36–7.33 (m, 1H, quin-*H*), 7.29 (dd, 1H, ³J_{HH} = 6.0 and 2.7 Hz, quin-*H*),



7.11–7.10 (m, 1H, quin-*H*), 6.96 (m, 1H, quin-*H*), 6.57 (dd, 1H, $^3J_{\text{HH}} = 8.3$ and 5.0 Hz, quin-*H*), 2.28 (d, 6H, $^3J_{\text{HH}} = 2.2$ Hz, CH_2 Ad), 2.20 (s, 6H, CH_3 Cp), 2.14 (m, 3H, CH Ad), 2.11 (s, 1H, *NH*), 1.79–1.61 (m, 6H, CH_2 Ad), 1.78 (s, 6H, CH_3 Cp), -0.06 (d, 12H, $^2J_{\text{YH}} = 1.5$ Hz, Y- CH_3). ^{13}C NMR (126 MHz, C_6D_6 , 26 °C) $\delta = 153.0$ (quin-*C*), 150.1 (quin-*C*), 139.9 (quin-*C*), 135.5 (quin-*C*), 133.0 (quin-*C*), 129.4 (quin-*C*), 127.0 (*C* Cp), 121.9 (CCH_3 Cp), 121.4 (CCH_3 Cp), 121.3 (quin-*C*), 58.3 ($\text{NC}(\text{CH}_2)_3(\text{CH})_3(\text{CH}_2)_3$), 49.5 ($\text{NC}(\text{CH}_2)_3(\text{CH})_3(\text{CH}_2)_3$), 37.0 ($\text{NC}(\text{CH}_2)_3(\text{CH})_3(\text{CH}_2)_3$), 31.8 ($\text{NC}(\text{CH}_2)_3(\text{CH})_3(\text{CH}_2)_3$), 13.2 (CH_3 Cp), 12.7 (CH_3 Cp), 3.8 (CH_3 , $\text{Al}(\text{CH}_3)_3$) ppm. DRIFT (KBr): 3048vw, 2907vs, 2847m, 1584vw, 1558vw, 1540vw, 1507m, 1490w, 1472w, 1456w, 1436w, 1419w, 1366w, 1301w, 1235w, 1184w, 1130vw, 1090vw, 1067w, 1023vw, 985vw, 968vw, 929w, 841vw, 822w, 790m, 760w, 732m, 691vs, 577w, 471w, 419vw cm^{-1} . Elemental analysis calcd (%) for $\text{C}_{32}\text{H}_{46}\text{AlN}_2\text{Y}$ (574.61 g mol^{-1}): C 66.89, H 8.07, N 4.88; found: C 65.33, H 7.48, N 4.12. Multiple attempts to obtain a better microanalysis failed.

AlMe₃(H₂Nad) (8). To a solution of 1-adamantylamine (47 mg, 0.31 mmol) in toluene (2 mL) trimethylaluminium (23 mg, 0.31 mmol) was added. The colourless solution was stirred at ambient temperature for 1 h and then dried under reduced pressure (68 mg, 0.31 mmol, quant., 77% crystalline yield). Crystallization from a toluene-*n*-hexane solution at -35 °C afforded colourless crystals. ^1H NMR (500 MHz, C_6D_6 , 26 °C, TMS): $\delta = 1.68$ (s br, 3H, $\text{NC}(\text{CH}_2)_3(\text{CH})_3(\text{CH}_2)_3$), 1.59 (s br, NH_2), 1.32–1.22 (m, 6H, $\text{NC}(\text{CH}_2)_3(\text{CH})_3(\text{CH}_2)_3$), 1.20 (d, 6H, $^3J_{\text{HH}} = 2.5$ Hz, $\text{NC}(\text{CH}_2)_3(\text{CH})_3(\text{CH}_2)_3$), -0.36 (s, 9H, $\text{Al}(\text{CH}_3)_3$) ppm. ^{13}C NMR (126 MHz, C_6D_6 , 26 °C) $\delta = 52.1$ ($\text{NC}(\text{CH}_2)_3(\text{CH})_3(\text{CH}_2)_3$), 43.6 ($\text{NC}(\text{CH}_2)_3(\text{CH})_3(\text{CH}_2)_3$), 35.9 ($\text{NC}(\text{CH}_2)_3(\text{CH})_3(\text{CH}_2)_3$), 29.9 ($\text{NC}(\text{CH}_2)_3(\text{CH})_3(\text{CH}_2)_3$), -5.7 ($\text{Al}(\text{CH}_3)_3$). DRIFT IR (KBr): 3262m, 3214m, 3117vw, 2910vs, 2851s, 1570m, 1455m, 1363m, 1352w, 1319w, 1298w, 1284w, 1206m, 1182m, 1169s, 1151s, 1102m, 1087s, 1042w, 982w, 936w, 922w, 815w, 711vs, 646w, 620m, 596m, 567w, 526w cm^{-1} . Elemental analysis calcd (%) for $\text{C}_{13}\text{H}_{26}\text{AlN}$ (223.33 g mol^{-1}): C 69.91, H 11.73, N 6.27; found: C 69.84, H 10.97, N 6.41. Multiple attempts to obtain a better microanalysis failed.

X-ray crystallography and crystal structure determination of 2a, 3, 4, 5, 7, and 8

Crystals of 2a and 3 were grown using standard techniques from saturated solutions using *n*-hexane or *n*-hexane-toluene (4, 5, 7 and 8) at -40 °C. Suitable crystals for X-ray structure analyses were selected in a glovebox and coated with Parabar 10312 and fixed on a nylon loop/glass fiber.

X-ray data for 3 were collected on a Bruker AXS TXS rotating anode instrument using a Pt 135 CCD detector and for 4, 7 and 8 on a Bruker APEX DUO instrument equipped with an $\text{I}\mu\text{S}$ microfocus sealed tube and QUAZAR optics for MoK_α radiation ($\lambda = 0.71073$ Å). Data for compounds 2a and 5 were collected on a Bruker SMART APEX II instrument equipped with a fine focus sealed tube and a graphite monochromator using MoK_α radiation ($\lambda = 0.71073$ Å). The Data collection strategy was determined using COSMO³⁸ employing ω - and ϕ scans. Raw data were processed using APEX³⁸ and SAINT,³⁸ corrections for absorption

effects were applied using SADABS.³⁸ The structure was solved by direct methods and refined against all data by full-matrix least-squares methods on F^2 using SHELXTL³⁸ and ShelXle.³⁹ All Graphics were produced employing ORTEP-3⁴⁰ and POV-Ray.⁴¹ Further details of the refinement and crystallographic data are listed in Table S1 (ESI†) and in the CIF files. CCDC 1055068–1055073.

Acknowledgements

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Notes and references

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