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Donor-free 9-aluminafluorenes: molecular structures and reactivity†

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Aluminum-doped polycyclic aromatic hydrocarbons (PAHs) are underexplored despite the broad applications of boron-containing PAHs in areas such as catalysis and optoelectronics. We disclose the donor-free, sterically unprotected 9-methyl-9-aluminafluorene (Me-AlFlu; **2**), synthesized by heating a 9,9-dimethyl-9-stannafuorene and AlMe₃ in hexanes. The compound is a dimer, (**2**)₂, with *trans*-positioned AlMe substituents in the solid state. In solution, (**2**)₂ shows a dynamic *cis/trans*-interconversion rather than a monomer-dimer equilibrium (Tol-*d*₈, RT). Lewis bases L cleave (**2**)₂ into monomeric adducts **2**·L (L = OEt₂, thf, pyridine). Lewis acidic AlBr₃ transforms (**2**)₂ into a 2,2'-(Br₂Al)₂-1,1'-biphenyl (**3**), crystallographically characterized as dimeric (**3**)₂. (**3**)₂ is a synthetic equivalent for the elusive free Br-AlFlu: Treatment with donor molecules furnishes Br-AlFlu·L adducts (L = OEt₂, pyridine); the three-coordinate, monomeric aluminafluorene Mes*–AlFlu was prepared from (**3**)₂, Mes*Li, and a 2,2'-dilithio-1,1'-biphenyl in quantitative yield (Mes* = 2,4,6-(*t*Bu)₃C₆H₂).

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Introduction

Doping organic π -electron systems with other p-block elements is an effective strategy to impart new chemical and physical properties to these species.¹ Specifically, the combination of a polycyclic aromatic hydrocarbon (PAH) such as fluorene with boron as a dopant to generate 9-borafluorenes (BFlu) can have a particularly pronounced effect,^{2,3} as a conjugation barrier (*i.e.*, the CH₂ fragment in the carbonaceous species) is removed and a vacant B(p_z) orbital is introduced instead, which can now: (i) mediate electron delocalization and bring about novel optoelectronic properties,⁴ (ii) facilitate reduction,^{5,6} and (iii) act as a Lewis acid to promote bond-activation reactions⁷ or the expansion of the five-membered central borole ring.⁸

Compared to the extensive research on BFlu, their heavier homologues, the 9-aluminafluorenes (AlFlu),^{9,10} are far less well explored. This is unfortunate, because AlFlu are expected to exhibit a lower degree of aryl-heteroatom double-bond character than BFlu,¹¹ leading to a greater propensity to form structurally intriguing aggregates through Al $\cdots\pi$ (Ar) complexes

or Al–C–Al' two-electron–three-center (2e3c) bonds. Relative to open-chain arylaluminum compounds, AlFlu should possess a structurally enforced enhanced Lewis acidity due to their small endohedral C–Al–C angle. This angle (108° in a regular five-membered ring) deviates more from the ideal 120° angle of three-coordinate AlR₃ species than from the corresponding angles of perfectly tetrahedral (109.5°) adducts. By the same token, the behavior of Al-based Lewis acids is more diverse than that of their B-based counterparts, as Al sites, unlike B centers, can readily accommodate coordination numbers larger than four.

In 1962, Eisch *et al.* reported the formation of Ph-AlFlu through the metalative cyclization of *o*-biphenyl(diphenyl) aluminum at 200 °C. Their claim was mainly based on the analysis of hydrolysis and iodinolysis products.^{12,13} The topic lay dormant until 2015, when Chujo and Tanaka used salt-metalathesis protocols to synthesize AlFlu carrying Al-bonded phenyl rings with one or two chelating (dimethylamino)methyl substituents at their *ortho* positions (Fig. 1). Their research focused on the emission properties of the obtained four- and five-coordinate AlFlu.^{14,15} More recently, Braunschweig *et al.* disclosed the synthesis of various aluminafluorenes R-AlFlu [R = 1,2,4-(*t*Bu)₃C₅H₂ (92%; Fig. 1), Ph₂(*t*Bu)Si (44%), 2-C₄H₃S (79%), *t*Bu (23%; Fig. 1), Br (53%)]. The compounds were again prepared from 2,2'-dilithio-1,1'-biphenyl by salt-metalathesis reactions and isolated and structurally characterized as their ether adducts – with the exception of the η^5 -cyclopentadienide derivative, which is monomeric in the solid state, and the *t*Bu derivative, which crystallizes as a dimer.¹⁶

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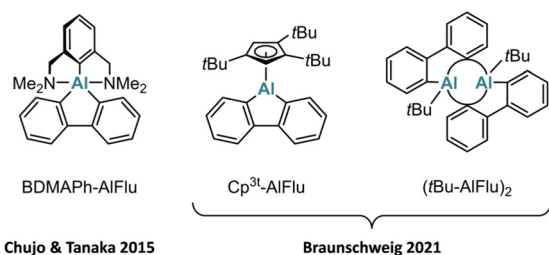


Fig. 1 Known mono- and dimeric 9-aluminafluorenes featuring 2,6-bis[(dimethylamino)methyl]phenyl (BDMAPh), 1,2,4-(*t*Bu)₃C₅H₂ (Cp^{3t}), and *tert*-butyl (*t*Bu) substituents.

One aim of our study outlined herein was to develop straightforward, high-yield synthesis protocols for base-free R-AlFlus featuring (i) the small substituent R = Me to minimize steric shielding of the Al center, and (ii) the reactive substituent R = Br for late-stage derivatization. Particular emphasis was placed on the molecular structure of Me-AlFlu in non-donor solvents and in the solid state, as well as on the synthesis of the first base-free, three-coordinate, monomeric aluminafluorene, Mes^{*}-AlFlu (Mes^{*} = 2,4,6-(*t*Bu)₃C₆H₂). All our AlFlus were equipped with *t*Bu groups in their 2,7-positions to enhance solubility in non-polar solvents and to facilitate NMR-spectroscopic analysis.

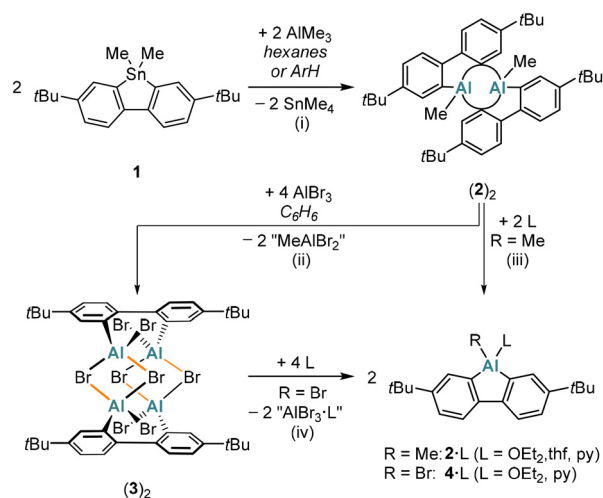
Results and discussion

Syntheses

The base-free Me-AlFlu (**2**) was synthesized by heating the 9,9-dimethyl-9-stannafluorene **1** with 1 equiv. of AlMe₃¹⁷ in either hexanes or C₆H₆/toluene (Scheme 1). The only by-product formed is the volatile and relatively inert SnMe₄.^{18,19} An advantage of using hexanes as the solvent is that the dimer (**2**)₂ precipitates in pure form already upon cooling the reaction mixture to room temperature (yield: 74%); when C₆H₆/toluene is employed, the yield of (**2**)₂ is higher (91%), but some further workup is required. In the presence of the donor molecules Et₂O, THF, or pyridine, (**2**)₂ is cleanly split into its constituting monomers to furnish the monoadducts 2-OEt₂, 2-thf, or 2-py (Scheme 1).

Treatment of (**2**)₂ with 4 equiv. of AlBr₃¹⁷ in C₆H₆ results not only in quantitative AlMe/AlBr exchange but also in the incorporation of two AlBr₃ molecules to afford dimeric 2,2'-(Br₂Al)₂-1,1'-biphenyl [(**3**)₂, 95%; Scheme 1]. Upon addition of Et₂O to (**3**)₂ in C₆H₆, the donor adduct of Br-AlFlu, 4-OEt₂, precipitates quantitatively as a colorless solid. In terms of yield, our overall synthesis cascade to 4-OEt₂ improves upon the published protocol¹⁶ by about 40 percentage points. Although pyridine can also reconstitute the AlFlu scaffold from (**3**)₂, it proved challenging to separate the target product 4-py from by-products such as [AlBr₂(py)₄][X] ([**5**][X]; X = Br[−], AlBr₄[−]; Fig. S44 and S45†).

A particularly notable application of (**3**)₂ as a synthetic equivalent of donor-free Br-AlFlu is the preparation of Mes^{*}-AlFlu

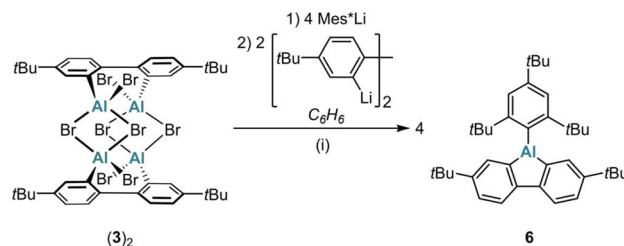


Scheme 1 Synthesis of donor-free (**2**)₂ through Sn/Al exchange between the 9-stannafluorene **1** and AlMe₃ (ArH: C₆H₆/toluene). The addition of AlBr₃ to (**2**)₂ furnishes (**3**)₂. Lewis bases (L: Et₂O, THF, or pyridine), cleave (**2**)₂ or (**3**)₂ into the monomeric adducts 2-L or 4-L. (i) Hexanes, 140 °C, 3 d (74% yield) or C₆H₆/toluene, 120 °C, 3 d (91% yield); sealed glass ampoule. (ii) C₆H₆, room temperature, 1 d (95% yield). (iii) 2-OEt₂: in Et₂O, room temperature; 2-thf: C₆D₆, room temperature; 2-py: C₆H₆, room temperature (quantitative conversions). (iv) 4-OEt₂: C₆H₆, room temperature (quantitative conversion); 4-py: C₆D₆, room temperature (not isolated). Note: in (**3**)₂, four bonds were arbitrarily chosen as formally intermolecular (highlighted in orange) to facilitate the distinction between the monomers M and M'.

(**6**): sequential addition of Mes^{*}Li (4 equiv.) and 2,2'-dilithio-4,4'-di-*tert*-butyl-1,1'-biphenyl (2 equiv.) to (**3**)₂ in C₆H₆ gave **6** in 97% yield (Scheme 2).

Solid-state structures

In the solid state, Me-AlFlu forms centrosymmetric dimers, with the Al-bonded Me substituents adopting a *trans*-configuration (*trans*-(**2**)₂; Fig. 2).²⁰ The individual monomers, M and M', are linked by two Al...C interactions, resulting in two Al(1) ... Al(1') bridging aryl rings (Ar_b) and two terminal rings (Ar_t), with bridging [C(11)] and terminal [C(21)] *ipso*-C atoms. The position of Ar_b is asymmetric between Al(1) and Al(1'), as indicated by the differing angles Al(1)–C(11) ... C(14) = 153.13(17)° and Al(1')–C(11) ... C(14) = 128.38(16)°. The fact that the



Scheme 2 Synthesis of Mes^{*}-AlFlu (**6**) using (**3**)₂ as a synthetic equivalent of the elusive Br-AlFlu. (i) C₆H₆, room temperature, 1 d (97% yield).

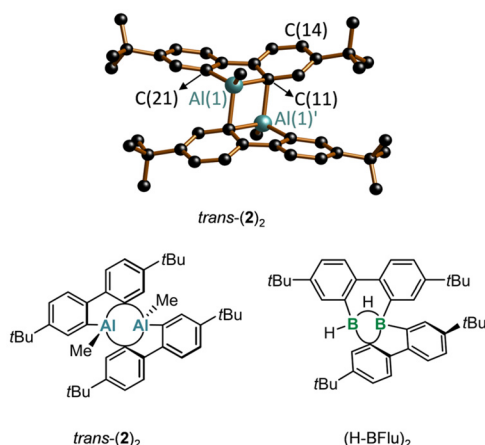


Fig. 2 Top: molecular structure of *trans*-(2)₂ in the solid state; H atoms omitted for clarity (C: black, Al: turquoise). Bottom: structural formulae of *trans*-(2)₂ and of the comparable 9-borafluorene dimer (H-BFlu)₂.

Al(1)′-C(11)-C(14) angle is significantly closer to 90° than the Al(1)-C(11)-C(14) angle can still be viewed as a remnant of the initial intermolecular Al⋯π(Ar) complex when the two heterofluorene units first encountered each other. Correspondingly, the ‘intermonomer’ Al(1)′-C(11) bond (2.148(3) Å) is longer by 0.055 Å than the ‘intramonomer’ Al(1)-C(11) bond (2.093(3) Å; *cf.* Al(1)-C(21) = 1.971(3) Å). The range of C-C bond lengths in Ar_b (1.382(5)–1.427(5) Å) is close to that in Ar_t (1.389(5)–1.409(5) Å), indicating that the bridging mode does not lead to a systematic bond-length alternation. However, the two C-C bonds involving the bridging C(11) atom are slightly longer than the other four (1.409(4) and 1.427(5) Å *vs.* 1.382(5)–1.401(5) Å). Finally, we note that *trans*-(2)₂ has very similar structural parameters to Braunschweig’s (tBu-AlFlu)₂,¹⁶ while the comparable 9-borafluorene dimer (H-BFlu)₂ shows one B-(μ-H)-B two-electron-three-center bond and one B⋯B′-bridging aryl ring (the three other rings remain terminally bonded).²¹

X-ray crystallography reveals that the compound (3)₂ no longer contains the 9-aluminafluorene motif but instead forms a centrosymmetric 2,2′-(Br₂Al)₂-1,1′-biphenyl dimer (Fig. 3).

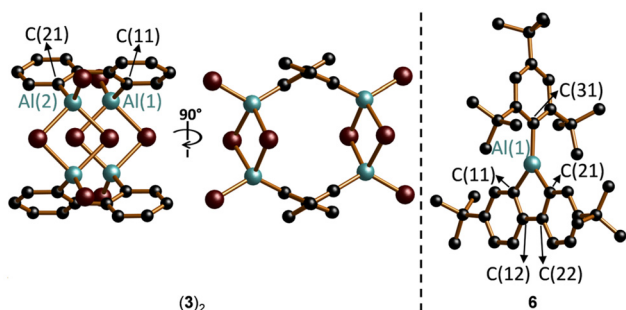


Fig. 3 Left: molecular structure of (3)₂ in the solid state, viewed from two different perspectives; tBu-groups in the 2,7-positions of the biphenyl backbones and H atoms omitted for clarity. Right: molecular structure of 6 in the solid state; H atoms omitted for clarity (C: black, Al: turquoise, Br: brown).

The two Br₂Al substituents in each monomer adopt an approximate *s-trans* configuration with a torsion angle Al(1)-C(11)-C(21)-Al(2) of 129.55(19)° [Al(1)-C(11) = 1.945(5) Å, Al(2)-C(21) = 1.956(4) Å]. Four Br atoms occupy bridging positions between Al centers of different monomers, assembling the cage-like structure of (3)₂. The underlying structural feature, a four-membered R(Br)Al-(μ-Br)₂-Al(Br)R ring, is common not only for aluminum tribromide (R = Br) but also for numerous dibromo(organo)alanes.²²

The 2,4,6-(tBu)₃C₆H₂-substituted Mes*-AlFlu (6) exists as a monomeric species with a three-coordinate Al center in the crystal lattice (Fig. 3). The sum of C-Al-C angles is 360°, confirming a trigonal-planar ligand environment, although the endocyclic C(11)-Al(1)-C(21) bond angle is nearly rectangular (91.79(6)°). All three Al-C bonds are of equal length, regardless of whether they are endo- or exocyclic, or whether the respective *ipso*-C(p_z) orbital is positioned parallel or orthogonal to the vacant Al(p_z) orbital [Al(1)-C(11)/C(21)/C(31) = 1.9611(14)/1.9516(14)/1.9606(13) Å]. Within the five-membered AlC₄ core, the length of the central C(12)-C(22) bond (1.5024(19) Å) approaches that of a typical C-C single bond (1.54 Å),²³ while the benzannulated bonds are significantly shorter [C(11)-C(12)/C(21)-C(22) = 1.4208(18)/1.4134(18) Å]. The other ten C-C bonds within the biphenyl fragment fall within a narrow range of 1.389(2)–1.4041(19) Å, closely matching the corresponding bonds in C₆H₆ (1.39 Å).²⁴ Taken together, this analysis of bond lengths suggests that the AlFlu moiety of 6 preserves two largely unperturbed Clar sextets within its two C₆H₃ fragments, with no indication of a delocalized (antiaromatic) π-system, nor any significant Al(1)=C(11)/C(21) double-bond character in the AlC₄ heterocycle.

The donor adducts 2-OEt₂, 2-py, and 4-py were subjected to X-ray analysis to confirm that (2)₂ and (3)₂ can indeed serve as precursors of Me-AlFlu and Br-AlFlu, respectively (Fig. S40, S41, S43†). Furthermore, compared to donor-free 6, the C-C bond lengths within the C₆H₃-C₆H₃ units of 2-py and 4-py were found to differ by no more than 3σ (and much less for most bonds).²⁵ This observation again suggests that the vacant Al(p_z) orbital exerts no significant electron-withdrawing mesomeric effect on the π-electron system.

NMR analysis

At room temperature, 2 gives severely broadened ¹H NMR signals, providing limited diagnostic value (Tol-*d*₈; Fig. 4 and S7†). At 70 °C, two sharp resonances are detectable in the aliphatic region of the spectrum (integral ratio 3H : 18H); the aromatic region contains one broad feature and two doublets with coupling constants of about 8.2 Hz (Fig. 4 and S6†). At −30 °C, the ¹H NMR spectrum of 2 is characterized by two well-resolved sets of signals attributable to two different but closely similar components (Fig. 4 and S8†); the same is true for the ¹³C{¹H} NMR spectrum (Fig. S9†). The proton-integral values of the two sets indicate a minor-to-major component ratio of approximately 0.12 : 1 (Fig. S8†). Focusing on the major component, the ¹H NMR spectrum reveals one singlet at −0.67 ppm (6H), and two additional singlets at 1.47 and



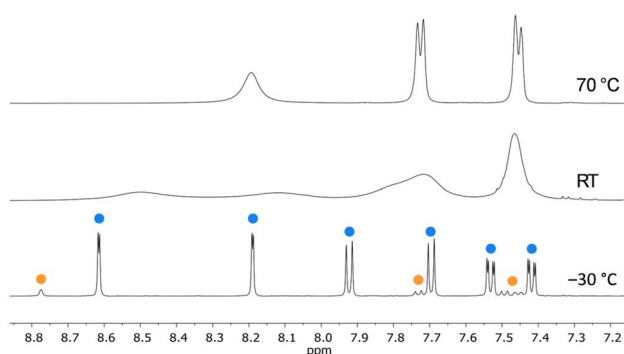


Fig. 4 Aromatic regions of ^1H NMR spectra of $(2)_2$ in Tol- d_8 (500.2 MHz). Top: 70 °C. Middle: room temperature. Bottom: –30 °C. ●: *trans*-(2) $_2$. ●: *cis*-(2) $_2$.

1.28 ppm ($2 \times 18\text{H}$), assignable to two equivalent AlMe substituents and two pairs of non-equivalent *t*Bu groups, respectively. In the aromatic region, four doublets ($4 \times 2\text{H}$; $2 \times {}^3J_{\text{H,H}} = 8.2\text{ Hz}$, $2 \times {}^4J_{\text{H,H}} = 2.2\text{ Hz}$) and two doublets of doublets ($2 \times 2\text{H}$) are observed, indicative of two pairs of non-equivalent C_6H_3 fragments. In principle, these NMR features would align with both the molecular structure of the *cis*- and *trans*-(2) $_2$ dimer (as observed in the solid state). *Vice versa*, the minor signal set likely arises from *trans*- or *cis*-(2) $_2$. At low temperatures, both isomers are present in an (essentially) static mixture, while some dynamic rearrangement equilibrium is established at higher temperatures. This preliminary conclusion raises two questions: (i) Does *cis*- or *trans*-(2) $_2$ dominate at low temperatures? (ii) Is the dynamic equilibrium at high temperatures due to monomer/dimer association/dissociation, or is it the result of a rapidly interconverting *cis/trans* dimeric form of $(2)_2$?

To address question (i), quantum-chemical calculations predict that the crystallographically characterized *trans*-(2) $_2$ is 1.6 kcal mol $^{-1}$ more favorable in energy than *cis*-(2) $_2$ (Scheme S1†; experimental value, determined at –30 °C from the proton-integral values of the minor/major component: $\Delta G^\circ = 0.7\text{ kcal mol}^{-1}$). Furthermore, the relative proportion of the minor component increases with solvent polarity, consistent with the existing dipole moment of *cis*-(2) $_2$ (^1H NMR spectroscopic control; Table S1 and Fig. S1, S2†). Finally, the computed ^{13}C chemical shift values for *cis/trans*-(2) $_2$ align more closely with the assumption that the major component is *trans*-(2) $_2$ rather than *vice versa* (Tables S9–S11†). It is therefore safe to assume that the major component in an equilibrating *cis/trans*-(2) $_2$ mixture is the *trans* isomer.

Regarding question (ii), we note that the computed energy required for cleaving *trans*-(2) $_2$ into its constituting monomers is 19.4 kcal mol $^{-1}$ (in CH_2Cl_2). In contrast, the computed energy barrier of the *cis/trans* interconversion of $(2)_2$ is only $\Delta G^\ddagger = 14.8\text{ kcal mol}^{-1}$, which agrees well with the value of $\approx 14.5\text{ kcal mol}^{-1}$ experimentally determined from the coalescence temperature (T_c) in conjunction with the maximum peak separation ($\Delta\nu$) in the slow-exchange limit (CD_2Cl_2 ; see the

ESI† for full details). The observed NMR features are therefore more convincingly attributed to a dynamic *cis/trans* equilibrium rather than to a monomer/dimer association/dissociation equilibrium.

In the temperature range of –30 to 70 °C, $(3)_2$ exhibited only extremely broadened signals in the ^1H and $^{13}\text{C}\{^1\text{H}\}$ NMR spectra, providing no structural information.

The ^1H and $^{13}\text{C}\{^1\text{H}\}$ NMR spectra of all adducts formed between our R-AlFlus and Lewis bases are in accord with the proposed molecular structures, as is the case for ligand-free **6** (see the ESI† for the fully assigned spectra). In addition to aiding in structure elucidation, $^{13}\text{C}\{^1\text{H}\}$ NMR spectroscopy is also a valuable tool for mapping the π -charge density distribution in conjugated systems, as the shielding of a specific $\text{C}(\text{sp}^2)$ atom depends linearly on the corresponding π -electron density at that position.²⁶ Given this background, we compared the ^{13}C chemical shift values of the C atoms constituting the C_6H_3 – C_6H_3 fragment of **6** with those of the equivalent atoms in the corresponding fragments of the adducts 2-OEt $_2$, 2-thf, 2-py, 4-OEt $_2$, and 4-py. Except for the Al-bonded *ipso*-C atoms, whose shift differences varied from $\delta(\text{6}) - \delta(\text{adduct}) = 3.3$ to –4.7 ppm without a systematic trend, the $\Delta\delta(^{13}\text{C})$ values for all other structurally analogous C atoms were less than $\pm 1.8\text{ ppm}$. In other words, we found no evidence of an overall ^{13}C -deshielding effect or π -electron depletion in **6** that could be attributed to a mesomerically electron-withdrawing $\text{Al}(\text{sp}^2)$ center.

$^{13}\text{C}\{^1\text{H}\}$ NMR spectroscopy on 2-py and 4-py provides a method to evaluate the relative Lewis acidities of free, monomeric Me-AlFlu and Br-AlFlu: in pyridine complexes of main-group elements, stronger acids induce increased shielding of the C-2,6 and deshielding of the C-3,4,5 nuclei of the ligand.²⁷ For 2-py/4-py, our observations consistently indicate that Me-AlFlu is the stronger acid, comparable in this respect to BPh_3 .²⁸ X-ray crystallography, however, offers a contrasting view: 4-py exhibits a shorter Al–N bond and a more pyramidalized Al center, implying higher Lewis acidity for Br-AlFlu.²⁸ Given the small differences in the key NMR and structural parameters between 2-py and 4-py, these conflicting observations highlight the limitation of relying on a single method to determine Lewis acidity, emphasizing the need for complementary approaches.

Conclusions

We synthesized the donor-free 9-aluminafluorene Me-AlFlu (**2**), which was characterized as its dimer $(2)_2$ through X-ray crystallography and VT NMR spectroscopy (Tol- d_8). The key to this success was the highly selective reaction between the 9,9-dimethyl-9-stannafluorene **1** and AlMe_3 ,^{17,18} which proceeds in non-donor solvents and releases volatile SnMe_4 as the sole by-product. Unlike the bulky *tert*-butyl group in *t*Bu-AlFlu,¹⁶ the sterically less demanding methyl substituent in Me-AlFlu allows relatively unhindered access to the electrophilic Al center, as demonstrated by the straightforward formation of



various base adducts 2·L (L = OEt₂, thf, py). In terms of an umpolung of the Al center, exploring the reduction of 2 on a preparative scale could be promising (a non-benzannulated alumole has previously been reduced to its corresponding dianion).¹⁰ The resulting product, [2]^{2−}, could potentially serve as an Al-centered nucleophile – analogous to the [H-BFlu]^{2−} dianion, which is a valuable B-centered nucleophile.^{3,6,29}

Treatment of (2)₂ with AlBr₃ furnishes the 2,2'-(Br₂Al)₂-1,1'-biphenyl (3)₂. Although this ring-opened product no longer retains the AlFlu motif, it rearranges back to afford Br-AlFlu adducts, such as 4·L (L = OEt₂, py), in the presence of Lewis bases. The use of Mes*Li, which provides the extremely bulky, negatively charged Lewis base [Mes*][−], grants unprecedented access to three-coordinate, monomeric aluminafluorenes, specifically Mes*-AlFlu (6), *via* LiBr elimination. This reaction proves the utility of (3)₂ as a synthetic equivalent for the still-elusive free Br-AlFlu. A comparison of characteristic structural and NMR features of 6 with those of 2·L/4·L reveals that the three-coordinate Al center exerts only a negligible π -electron withdrawing effect and does not mediate significant π -electron delocalization.

Author contributions

P. L. L. performed the experimental studies and characterized all new compounds. P. L. L. and J. G. performed the quantum-chemical calculations. A. V. performed the X-ray crystal structure analyses of all compounds. H.-W. L. and M. W. supervised the project. The manuscript was written by P. L. L. and M. W. and edited by all co-authors.

Data availability

The datasets supporting this article have been uploaded as part of the ESI.†

Conflicts of interest

There are no conflicts to declare.

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