Synthesis and Thermal Study of Hexacoordinated Aluminum(III) Triazenides for Use in Atomic Layer Deposition

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ABSTRACT: Amidinate and guanidinate ligands have been used extensively to produce volatile and thermally stable precursors for atomic layer deposition. The triazenide ligand is relatively unexplored as an alternative ligand system. Herein, we present six new Al(III) complexes bearing three sets of a 1,3-dialkyltriazenide ligand. These complexes volatilize quantitatively in a single step with onset volatilization temperatures of ~150 °C and 1 Torr vapor pressures of ~134 °C. Differential scanning calorimetry revealed that these Al(III) complexes exhibited exothermic events that overlapped with the temperatures of their mass loss events in thermogravimetric analysis. Using quantum chemical density functional theory computations, we found a decomposition pathway that transforms the relatively large hexacoordinated Al(III) precursor into a smaller dicoordinated complex. The pathway relies on previously unexplored interligand proton migrations. These new Al(III) triazenides provide a series of alternative precursors with unique thermal properties that could be highly advantageous for vapor deposition processes of Al containing materials.

1. INTRODUCTION

Aluminum nitride (AlN) is a semiconductor material widely used in current day electronic devices.1 This is due to its desirable chemical, optical, and electronic properties, such as high thermal stability, a wide direct band gap, and piezoelectricity.2 As electronic devices rapidly miniaturize with increasingly complex surface structures, atomic layer deposition (ALD) becomes a vital technique for depositing uniform thin films of high-performance materials for future microelectronics.3 In ALD, the metal and nonmetal precursors are introduced into the reaction chamber separately, which allows the film mechanism to be governed by two independent and self-limiting half reactions. These are complex surface reactions that can incorporate impurities into the film if the metal precursor does not possess suitable physical and chemical properties. A desirable ALD metal precursor must be thermally stable until reaching the film surface.4 Here, it should undergo a clean and fast reaction to form a single stable monolayer without trapping unwanted byproducts.5 This monolayer should then react with the second precursor (e.g., NH3 or H2O) in the same way. To maximize the growth rate of a thin film, a precursor must be sufficiently volatile and have ligands of low steric bulk for fast surface saturation and maximum density of the deposited precursor. Due to its high volatility and reactivity, trimethylaluminum (AlMe3) has been used to deposit AlN by ALD.5–13 These films contain high levels of carbon impurities due to the strong Al–C bonds, making it difficult to remove all of the methyl ligands of the deposited precursor at low temperatures.5,6 Replacing the Al–C of AlMe3 with more reactive Al–N bonds has led to homoleptic tricoordinated amide precursors (Al(NMe2)3)14 and (Al(NEt2)3)15 which have been used to deposit AlN by ALD.16–19 Although these precursors are highly volatile and reactive, the low thermal stability of the deposited surface species renders films with carbon impurities.20

Amidinate and guanidinate bidentate ligands have been employed to improve thermal stability of group 13 metal precursors. Although these ligands improve thermal stability in comparison to monodentate ligated precursors, their drawback is compounds that lack volatility or surface reactivity, or both, the latter due to crowding of the metal center.21–27 In particular, homoleptic hexacoordinated M–N bonded Al(III) amidinate (Al(amid)3) and guanidinate (Al(guan)3) compounds possess increased thermal stability compared to tricoordinated Al(III) amides,26,27 but have not been used in an ALD process due to insufficient volatility. A ligand closely related to the amidinate and guanidinate is the triazenide, differing by the nitrogen atom in the endocyclic position of the ligand backbone. Homoleptic hexacoordinated Al(III) triazenide complexes have previously been reported;28,29 however, they are not volatile due to their 1,3-diphenyltriazenide ligands.

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Recently, we reported the first examples of highly volatile homoleptic 1,3-dialkyltriazenide complexes, tris(1,3-dipropyltriazenide)In(III) (In(triaz)₃) and Ga(III) (Ga(triaz)₃), and their use as ALD precursors. These new triazened precursors underwent gas-phase decomposition at higher temperatures inside the ALD reactor, giving a smaller and more reactive M(III) species. This in situ thermolysis was highly advantageous for film growth, giving higher growth rates and films with near stoichiometric M/N ratios without unwanted carbon impurities. To further explore the unique properties of the 1,3-dialkyltriazenide ligand, we envisaged its ability to stabilize the Al(III) center to develop a new series of precursors that can be used for future ALD processes. Herein, we describe the synthesis, structure, and thermal properties of six homoleptic Al(III) 1,3-dialkyltriazenide complexes. These compounds were easily synthesized in good yields and are the first example of volatile hexacoordinated M−N bonded aluminum compounds. Furthermore, the compounds exhibit unique thermal properties, similar to Ga(triaz)₃ and In(triaz)₃. Using quantum-chemical density functional theory (DFT) calculations, we mapped out a previously unexplored decomposition pathway utilizing interligand interactions. The pathway is supported by electron impact mass spectrometry (EI-MS) data. The unique thermal properties of these compounds make them potentially advantageous as precursors for vapor deposition processes.

### 2. RESULTS AND DISCUSSION

#### 2.1. Synthesis and Characterization of Aluminum Complexes

Tris(1,3-dialkyltriazenide)aluminum(III) compounds 1−6 were prepared in good yields by reacting the (1,3-dialkyltriazenide)lithium(I) intermediate, generated from an alkylazide and alkyllithium, with AlCl₃ (Scheme 1).

**Scheme 1. Synthesis of Tris(1,3-dialkyltriazenide)aluminum(III) Compounds 1−6**

| R¹ | R² | R³ |
|---|---|---|
| N≡N | N≡N | N≡N |
| R¹ = Pr, tert-C₆H₁₃, Bu | R² = Pr, tert-C₆H₁₃, Bu | R³ = Pr, tert-C₆H₁₃, Bu |

The crystal structure of 6 showed the aluminum in an octahedral coordination geometry bearing three sets of the 1,3-di-tert-butytriazenide ligand (Figure 1). A large degree of disorder was observed in the diffraction data. The ligands are distorted over a multitude of positions (at least 8 individual sets of possible arrangements). The Al−N bond length for 6 (av 1.96(5) Å) is similar to that for tris(1,3-diphenyltriazenide)aluminum(III) (av 1.972(5) Å) but slightly shorter than those of Al(guan)₃ (av 2.024 Å) and Al(amd)₃ (av 2.0195 and 2.0261 Å). Thus, the triazene ligand only has a small effect on the Al−N bond length in comparison to Al(guan)₃ and Al(amd)₃. Compound 1 has an analogous structure to that of 6, but suffers from even more severe disorder and therefore no successful refinement could be completed (see the Supporting Information).

The DFT calculated geometry of 6 is consistent with its crystal structure. The highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO) are both localized on the ligands (Figure 2). While the HOMO is spread over all three ligands, the LUMO only covers the N₃ backbone of two ligands. For the HOMO, one node is centered on the endocyclic nitrogen, while the LUMO has nodes between the endocyclic and exocyclic nitrogens. The experimental EA was not obtained from the sublimed and filtered solid. To obtain satisfactory EA, the compounds were therefore purified by recrystallization from Et₂O/MeCN.

Figure 1. ORTEP drawing for one of two independent molecules in the unit cell of 6. Thermal ellipsoids are displayed at the 50% probability level, and hydrogen atoms are omitted for clarity.

Figure 2. (a) HOMO (−5.74 eV) and (b) LUMO (−0.78 eV) for 6 from DFT calculations.
natural charges of the exocyclic nitrogens (−0.48) and Al metal center (+1.66) indicate a highly polarized Al−N bond character. The primary carbons (−0.60) show negative charges, while the endocyclic nitrogens (+0.044) and tertiary carbons (+0.11) have slightly positive charges. Comparing the natural bond orbital charge of the Al center of 1 (1.61) and its formamidinate analogue (1.86) shows the greater electron donating ability of the triazenide ligand over the formamidinate (see the Supporting Information).

The $^{27}$Al NMR spectra of compounds 1–6 each gave a broad peak at $\delta_{\text{Al}} = 25.1$–27.5 ppm (see the Supporting Information). These chemical shifts are consistent with previously reported hexacoordinated Al(III) triazenide complexes ($\delta_{\text{Al}} = 25$–28 ppm). Dynamic effects were observed by $^1$H NMR for the unsymmetrical ligated compounds 2, 3, and 5 at 25 °C (see the Supporting Information). These effects are most likely caused by isomerization hindered by the bulky ligands surrounding the small Al(III) center. Therefore, the complexes isomerize slowly, resulting in significant lifetimes for the signals in relation to the difference in resonance frequencies. Heating resolves the lifetime broadening by increasing the rate of isomerization. Mild line broadening was observed for compound 2 at 25 °C and was resolved at 35 °C. Line splitting was observed for all but the CH signals for compounds 3 and 5. These more severe effects are caused by the presence of the bulky tert-butyl groups, which further inhibit isomerization. The line splitting was resolved at 45 °C with only line broadening remaining. Heating to 50 °C resolved the line broadening for compound 3. However, mild line broadening was still observed for 5.

2.2. Thermal Analysis of Aluminum Complexes.

Compounds 1, 3, 5, and 6 volatilize quantitatively with exponential mass loss in thermogravimetric analysis (TGA) (Figure 3). Compound 2 has 4% residual mass by TGA, undergoing slight decomposition, as observed by an inflection in the derivative at approximately 200 °C (see the Supporting Information). TGA of 4 showed two distinct events of mass loss, giving 5–7% residual mass. We speculate that 4 decomposes into volatile fragments during the mass loss events. However, we were unable to obtain a satisfactory elemental analysis for 4; therefore, the residual mass may be due to impurities. Overall, TGA shows that compounds 1–6 are sufficiently volatile for use in ALD. In fact, compounds 1–6 are far more volatile than Al(amide)$_3$ and Al(guan)$_3$ which can be explained by more electron density residing on the triazenide ligand. This leads to weaker intermolecular interactions in the crystal structure of the compound, which is conditional for faster volatilization (see the Supporting Information for a comparison of charges between 1 and its amidinate analogue).

Differential scanning calorimetry (DSC) was employed to study exothermic events of 1–6. All compounds have exothermic events, most likely due to decomposition, overlapping with their onset of volatilization displayed in TGA (see the Supporting Information). The TGA and DSC results for 1–6 are summarized in Table 1. Compounds 1 and 5 have exothermic events starting at 150 and 160 °C, respectively. Both compounds give peaks with irregular shape, indicating overlapping exothermic events occurring. Two distinct exotherms are observed for 2, 3, and 6: the first event initiates at 130, 160, and 230 °C, respectively, and the second between 234 and 300 °C. Compound 4 has a small exotherm starting at ~100 °C, followed by a larger event at ~200 °C. The unsymmetrical compounds 2, 3, and 5 undergo exothermic events at similar temperatures to 1. That is, greater ligand bulk does not increase thermal stability of these compounds. However, the exothermic event of 6 is at significantly higher temperatures compared to 1. Interestingly, 1 and 6 have similar calculated 1 Torr vapor pressure temperatures (Table 1).

To study the long-term thermal stability of compounds 1–6, their solids were each flame sealed in an NMR tube and heated to 5 °C above their 1 Torr vapor pressure temperature (Table 1) for 7 days. After heat exposure, compounds 1, 2, and 5 fully dissolved in C$_6$D$_6$ while 3 and 6 had a small amount of insoluble solid. None of the compounds showed signs of decomposition by $^1$H NMR.

A solution of 1 in C$_6$D$_6$ was heated in a flame-sealed NMR tube to various temperatures for 1 h, followed by acquiring $^1$H NMR spectra (Figure 4). After flame sealing but before heating the sample, the $^1$H NMR spectrum showed newly formed, low-intensity impurity signals (red asterisk, Figure 4). Only small changes to 1 occurred when heating up to 180 °C, and the solution remained colorless. Above 210 °C, the solution turned yellow, and $^1$H NMR signals of the doublet and septet, at 1.25 and 3.88 ppm, respectively, decreased in intensity. A white precipitate formed at 225 °C, and the overall $^1$H NMR signal of 1 had decreased significantly. Traces of various decom-

![Figure 3. Thermogravimetric analysis of 1–6.](https://dx.doi.org/10.1021/acs.inorgchem.0c03496)
position products appear in the ranges 3.0−2.7, 2.4−2.5, and 2.0−0.7 ppm (see the Supporting Information).

2.3. Gas-Phase Decomposition by DFT Computations. In previous work, we demonstrated high-quality thin films of indium nitride and gallium nitride by ALD, using the Ga(triaz)₃ and In(triaz)₃ as precursors, respectively.¹³,³⁰ From the thermal properties of the compounds, we speculated that the depositions are activated by gas-phase decomposition of the precursor in the ALD reactor. Compounds 1−6 have similar thermal properties as Ga(triaz)₃ and In(triaz)₃ and are therefore expected to undergo a similar decomposition. DFT was used to study gas-phase thermal decomposition pathways of 1. We found a decomposition pathway relying on Brönsted−Lowry acid−base reactions between neighboring ligands. Overall, the first ligand leaves as triazene while the second decomposes into an imido ligand. Figure 5 shows the free energy profile for the release and decomposition of the first and second ligand, respectively, at 250 °C and 10 hPa. The third ligand decomposes in the same manner as the second, only with minor differences (Figure 6). The largest free energy barriers are 211 and 214 kJ mol⁻¹, for TS-3 (Figure 5) and TS-8 (Figure 6), respectively. These barriers are slightly smaller.

Figure 4. The ¹H NMR (500 MHz, C₆D₆) spectra from a decomposition study of 1 between 0.8−1.3 and 3.7−4.0 ppm separated by an axis break. For visibility, the y-axis is scaled up ~18 times on the left of the axis break compared to the right side. Prior to flame sealing, the compound showed no traces of impurities by ¹H NMR analysis. The peaks marked with an asterisk appeared after flame sealing the tube. Compound 1 was heated in C₆D₆ and all spectra were acquired at 50 °C to suppress line broadening. The decomposition of 1 accelerates after 210 °C, which is shown by the diminished quartet and doublet peaks.

Figure 5. Free energy profile (at 250 °C and 10 hPa) for the first half of the decomposition pathway. Here, 1 loses a triazene ligand (after TS-2), and one ligand decomposes into an imido ligand (TS-5). TS-3 has the largest free energy (211 kJ mol⁻¹) for the displayed part of the decomposition pathway. The overall largest free energy barrier is found at TS-8 (214 kJ mol⁻¹): the analogous step to TS-3 but for the last ligand. At 250 °C and 10 hPa, the adduct structures I-2A separate spontaneously (i.e., the process is barrierless and has a negative free energy difference) and is therefore not included.

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for 6 (192 and 197 \text{kJ mol}^{-1}). The Supporting Information contains pictures and Cartesian coordinates of all optimized geometries and their respective enthalpies and free energies.

Starting from 1, a ligand dechelates from the metal center by a 180° rotation of a N–N bond (TS-1), resulting in I-1. This ligand dechelation enables the isopropyl moiety on the coordinated nitrogen to move closer to neighboring ligands and the metal center. Next, a methyl proton of the isopropyl group migrates to an exocyclic nitrogen on a neighboring ligand (TS-2). Simultaneously, a bond is formed between the deprotonated isopropyl group and the metal center. In I-2A, the deprotonated ligand regains a bidentate binding mode, now with a C,N-coordination to the metal center. Meanwhile, the protonated ligand dechelates and only has a coordination bond to the metal center. Due to the proton transfer in TS-2, the deprotonated and protonated ligand become dianionic and neutral, respectively. I-2A is an adduct structure consisting of a triazene (LH) coordinated to the I-2 structure.

The two steps that transform 1 into I-2A are reversible. In contrast, separating I-2A into LH and I-2 may be reversible or irreversible depending on the reaction conditions (Scheme 2). I-2A separates spontaneously under reduced pressure and elevated temperature, conditions commonly employed in ALD. Therefore, we assume that when the adduct structure separates, LH becomes inaccessible to I-2 and cannot facilitate the backward reaction, i.e., making the forward reaction irreversible.

I-2 has two triazenide ligands: the dianionic C,N-coordinated ligand and an unaltered N,N-coordinated ligand. The dianionic ligand decomposes in three irreversible steps, transforming the C,N-coordinated triazenide into an imido ligand (Scheme S2). First, a methyl proton migrates from the isopropyl group on the \( \beta \)-nitrogen to the \( \alpha \)-nitrogen, with respect to the coordinated nitrogen (TS-3). This proton transfer results in a molecule of propene leaving the structure, giving I-3. Second, the proton on the \( \alpha \)-nitrogen migrates to the coordinated nitrogen (TS-4), releasing dinitrogen to give I-4. Third, passing through TS-5, the former isopropyl group, which coordinated to the metal center in TS-2, leaves as propene to give I-5. Other than the newly formed imido ligand, I-5 has one monoanionic N,N-coordinated and one dianionic C,N-coordinated triazenide ligand.

The reverse step through TS-7 has a significantly larger free energy barrier compared to the analogous TS-2 (290 vs 85 \text{kJ mol}^{-1}, respectively).
Table 2. Summarized EI-MS Signals (Given in m/z) And Their Matching Intermediate Fragments from the Presented Decomposition Pathway for Compounds 1 and 6

|                  | [I-4]⁺ | [I-4 – Me]⁺ | [I-5]⁺ | [I-9 – NH₂]⁺ | [I-10]⁺ |
|------------------|--------|-------------|--------|--------------|---------|
| **1               |        |             |        |              |         |
| Calcd            | 211.15 | 197.13      | 170.11 | 84.04        | 58.01   |
| Found            | 211.21 | 197.20      | 170.18 | 84.07        | 58.09   |
| **2               |        |             |        |              |         |
| Calcd            | 239.18 |             | 198.14 | 98.06        | 58.01   |
| Found            | 239.28 |             | 198.21 | 98.10        | 58.10   |

acts as the Brensted–Lowry base instead of a neighboring triazenide ligand. Furthermore, this step has a significantly lower free energy barrier compared to TS-2 due to the imido being a much stronger base compared to the triazenide ligand. For the same reason, the free energy barrier for the reverse reaction via TS-7 is very large (290 kJ mol⁻¹), essentially blocking the backward reaction. After passing TS-7, the deprotonated and protonated ligands transform into a dianionic C,N-coordinated triazenide and an amido ligand, respectively, to give I-7.

The second dianionic triazenide decompose into an imido ligand via TS-8 to TS-10, which are analogous to TS-3 to TS-5, giving the adduct structure I-10A. The adduct separates spontaneously into the final structure, I-10, and propane. A lack of viable options for I-10 to further decompose makes it thermally stable in the gas phase. However, based on the structure, I-10 is expected to be highly reactive toward surfaces. EI-MS data for I show four signals consistent with fragments for intermediates of the presented decomposition pathway (see the Supporting Information). Three potential fragments are identified for 6 and these fragments are structurally analogous to fragments found for 1. Furthermore, both compounds give a low intensity signal (~1%) at m/z 58, matching I-10. The calculated and found m/z for all fragments are given in Table 2.

3. CONCLUSION

In conclusion, six tris(1,3-dialkyltriazenide)aluminum(III) compounds have been made in good yields. The crystal structure of 6 revealed a homoleptic complex with three sets of the 1,3-di-tert-butyltriazenide ligand chelating to the Al(III) center. The complexes are highly volatile and, with the exception of 4, volatilize in a single step with exponential mass loss. Exponential mass loss indicated that the compounds volatilize without decomposing. However, DSC revealed exothermic events, most likely due to decomposition, overlapping with the temperature range for the mass loss event in TGA. Therefore, the compounds may undergo decomposition upon or after volatilization. Using DFT, a gas-phase decomposition pathway was found that relies on protons migrating between ligands. One ligand leaves the complex as a molecule of triazene, and the two remaining ligands transform into C,N-coordinated dianionic triazenide ligands that decompose into imido ligands. The largest free energy barrier for the pathway is 214 kJ mol⁻¹ (at 10 hPa and 250 °C) during decomposition of the second dianionic ligand (Figure 6, TS-8). A slightly lower free energy barrier of 211 kJ mol⁻¹ (at 10 hPa and 250 °C) is found for TS-3 (decomposition of the first ligand, analogous to TS-8). The final intermediate of the pathway is predicted to be highly reactive. Further studies are required to form a better understanding of how the compounds behave during the mass loss events observed in TGA. To our knowledge, compounds 1–6 are the first hexacoordinated Al–N bonded compounds that are sufficiently volatile for use as Al precursors in vapor deposition. Based on thermal analysis and DFT calculations, we postulate that the compounds decompose into smaller and more reactive species in the gas phase, which would be highly beneficial for ALD.

4. EXPERIMENTAL SECTION

4.1. General Experimental Procedures. Caution! As catenated nitrogen compounds are known to be associated with explosive hazards, alkylazides and compounds 1–6 are possible explosive energetic materials. Although we have not experienced any problems in the synthesis, characterization, sublimation, heating, and handling of compounds 1–6, their energetic properties have not been fully investigated and are therefore unknown. We therefore highly recommend that all appropriate standard safety precautions for handling explosive materials (safety glasses, face shield, blast shield, leather gloves, polymer apron, and ear protection) be used at all times when working with isopropyl-, sec-butyl-, and tert-butylazide and compounds 1–6.

All reactions and manipulations were carried out under a N₂ atmosphere on a Schlenk line using Schlenk air-free techniques, or in a N₂-filled drybox from Glovebox-Systemtechnik. All anhydrous solvents were purchased from Sigma-Aldrich and further dried with molecular sieves 4 Å. Isopropylithium (0.7 M in pentane), sec-butyllithium (1.4 M in cyclohexane), and tert-butyllithium (1.7 M in pentane) were purchased from Sigma-Aldrich, and AlCl₃ (99.985%) was purchased from Alfa Aesar; all were used without further purification. tert-Butyl-, sec-butyl-, and isopropylazide were synthesized according to previously reported literature procedures.32,41 All
NMR spectra were measured with Oxford Varian 300 and AS500 spectrometers at room temperature unless otherwise stated. Solvents' peaks were used as an internal standard for the $^{1}H$ NMR (300 and 500 MHz) and $^{13}$C NMR (75 and 125 MHz) spectra. For temperature stability measurements using NMR, a 40 g L$^{-1}$ solution of 1 in CD$_2$Cl$_2$ was added to a heavy-walled NMR tube, and the tube was flame-sealed. EI-MS data for 1-6 were acquired using a Varian MAT spectrometer operated at 70 eV in the electron ionization mode. Samples were filled into steel cartridges, sealed with lids, and individually fed to the spectrometer via a load-lock chamber which was pumped to ultrahigh vacuum prior to sample transfer to the main chamber. Melting points were determined for samples under N$_2$ atmosphere, flame-sealed in capillaries, using a Stuart SMP10 melting point apparatus and are uncorrected. Elementar analysis was performed by Mikroanalytisches Laboratorium Kolbe, Germany. Purification of compounds 1–6 by sublimation gave unsatisfactory EA results. Satisfactory EA results were obtained by recrystallizing the sublimed compounds.

4.2. General Synthesis Procedure for Al(III) Triazenide Complexes. Alkylithium (3 equiv) was added to a solution of alkyl azide (3 equiv) in Et$_2$O at $-78^\circ$C, and the reaction mixture was stirred at this temperature for 30 min and then at room temperature for 1 h. This solution was then added to a $-78^\circ$C solution of AlCl$_3$ (1 equiv) in Et$_2$O via cannula. The reaction mixture was stirred at this temperature for 30 min and then slowly warmed to room temperature and stirred for 16 h. The reaction mixture was then concentrated under reduced pressure, and the resulting residue was suspended in n-hexane. Solids were filtered off through a pad of Celite and concentrated under reduced pressure to give the crude product. Purifying the crude product by sublimation resulted in unsatisfactory purity by EA. The crude product was therefore purified by recrystallization from Et$_2$O/MecCN at $-35^\circ$C to give the desired trig-[1,3-dialkytriazenide]aluminum(III) complexes, 1–6.

Tris(1,3-disopropyltriazenide)aluminum(III) ([4]). Compound 1 was synthesized according to the general procedure using isopropyl azide (0.42 g, 4.93 mmol) in Et$_2$O (2.5 mL), isopropylithium (7.05 mL, 4.93 mmol), and AlCl$_3$ (0.22 g, 1.65 mmol) in Et$_2$O (25 mL). The solid was purified by recrystallization to give 1 as a solid (0.39 g, 58%).

1: Colorless solid, mp 255–257°C. Sublimation: 90°C (at 0.5 mbar). $^{1}H$ NMR (300 MHz, CD$_2$Cl$_2$): δ 1.25 (d, $J = 6.7$ Hz, 36H, CH$_3$), 3.88 (sept, $J = 6.7$ Hz, 6H, CH$_3$)$_3$. $^{13}$C($^{1}H$) NMR (75 MHz, CD$_2$Cl$_2$): δ 23.4 (s, CH$_3$), 52.6 (s, CH). $^{25}$Al NMR (78 MHz, CD$_2$Cl$_2$): δ 455.3 (s, AlCl$_3$). Anal. Calcld for C$_{21}$H$_{39}$AlN$_9$: C, 55.12%; H, 10.32%; N, 28.97%. Found: C, 55.07%; H, 10.71%; N, 27.53%. 

Tris(1,3-disopropyltriazenide)aluminum(III) (5). Compound 5 was synthesized according to the general procedure using tert-butyl azide (0.45 g, 4.54 mmol) in Et$_2$O (25 mL), tert-butyllithium (2.45 mL, 3.43 mmol), and AlCl$_3$ (0.15 g, 1.14 mmol) in Et$_2$O (20 mL). The solid was purified by recrystallization to give 5 as a solid (0.59 g, 78%).

2: Colorless crystals, mp >300°C. Sublimation: 120°C (at 0.5 mbar). $^{1}H$ NMR (300 MHz, CD$_2$Cl$_2$): δ 0.92 (t, $J = 7.4$ Hz, 9H, CH$_3$), 1.27 (d, $J = 6.6$ Hz, 9H, CH$_3$), 1.35 (s, 27H, CH$_3$), 1.40–1.62 (m, 3H, CH$_3$), 1.75–1.94 (m, 3H, CH$_3$), 3.51–3.66 (m, 3H, CH$_3$). $^{13}$C($^{1}H$) NMR (75 MHz, CD$_2$Cl$_2$): δ 10.4–12.1 (m, 18H, CH$_3$), 17.9–19.7 (m, 18H, CH$_3$), 30.5–30.9 (m, 18H, CH$_3$), 56.4 (C), 57.5–58.6 (m, 6H). $^{25}$Al NMR (78 MHz, CD$_2$Cl$_2$): δ 26.0 (br s). Anal. Calcld for C$_{39}$H$_{78}$AlN$_9$: C, 58.15%; H, 10.98%; N, 25.43%. Found: C, 55.92%; H, 10.66%; N, 24.42%. 

Tris(1,3-disopropyltriazenide)aluminum(III) (6). Compound 6 was synthesized according to the general procedure using tert-butyl azide (0.45 g, 4.54 mmol) in Et$_2$O (25 mL), tert-butyllithium (2.67 mL, 4.54 mmol), and AlCl$_3$ (0.20 g, 1.51 mmol) in Et$_2$O (25 mL). The solid was purified by recrystallization to give 6 as a solid (0.59 g, 78%).

3: Colorless crystals, mp >300°C. Sublimation: 125°C (at 0.5 mbar). $^{1}H$ NMR (300 MHz, CD$_2$Cl$_2$): δ 1.38 (s, 54H, CH$_3$). $^{13}$C($^{1}H$) NMR (75 MHz, CD$_2$Cl$_2$): δ 31.3 (s, C), 57.3 (C). $^{25}$Al NMR (78 MHz, CD$_2$Cl$_2$): δ 23.7 (br s). EI-MS (LR): 41.1 (3.7%), 57.1 (19%), 98.1 (1.1%), 198.2 (9.9%), 239.3 (2.2%), 339.4 (100%), 495.5 (0.51%). Anal. Calcld for C$_{39}$H$_{78}$AlN$_9$: C, 58.15%; H, 10.98%; N, 25.43%. Found: C, 58.16%; H, 10.96%; N, 25.73%. 

4.3. X-ray Crystallographic Analysis. Colorless single crystals were obtained by recrystallization from n-hexanes at $-35^\circ$C. The single crystals were used for X-ray diffraction data collection on a Bruker D8 SMART Apex-II diffractometer, using graphite-monochromated Mo K$\alpha$ radiation ($\lambda = 0.71073$ Å) at 153 K. All data were collected in hemisphere with over 95% completeness to 2θ < 50.05°. The structure is monoclinic, centrosymmetric, space group C2/m, $a = 28.940$(5), $b = 16.958$(3), $c = 9.8864$(17), $\beta = 94.118(2)$. In spite of data collection at low temperature, the data produce an electron density map that is rather flat, which is a result of very heavy disorder, rendering an almost amorphous structure. The structure was solved by direct methods. Coordinates of metal atoms were determined from the initial solutions, and from the N and C methods, located in subsequent differential Fourier syntheses. The solution does not contain much residual electron density, but it stays for a multitude of additional possible positions of light atoms. All nonhydrogen atoms were refined, first in isotropic and then in anisotropic approximation, using Bruker SHELXTL software. Additional crystal data treatment details are available from the Cambridge Crystallographic Data Centre, deposition no. CCDC 2046088.
15.817(15), c = 9.184(9) Å, β = 95.069(14)°. A model analogous to that for 6 was obtained but could not be refined successfully because of poor data quality. The details for the model obtained for the structure of 1 are available in the Supporting Information.

4.4. Thermogravimetric Analysis. Volatilization and vapor pressure curves were collected using a TA Instruments thermogravimetric analysis Q500 tool operating inside a N$_2$-filled glovebox. The ramp experiment of compounds 1–6 was undertaken in tared platinum pans loaded with ∼5–10 mg for low mass volatilization experiments. The furnace was heated at a rate of 10 °C min$^{-1}$ to 500 °C with a maintained N$_2$ flow rate of 60 sccm. The Langmuir vapor pressure equations for compounds 1–4 and 6 were derived from TGA mass-loss derivative data of the ramp experiments according to a previously reported method$^{38}$ employing bis(2,2,6,6-tetramethyl-3,5-heptanedionato)copper(II) as a calibrant.$^{39}$

4.5. Differential Scanning Calorimetry Analysis. DSC measurements were performed using a TA Instruments DSC Q10 tool. For each compound, 1–6, 0.2–0.5 mg of the compound was sealed in a platinum pan in a N$_2$-filled glovebox. All experiments were performed at a heating rate of 10 °C min$^{-1}$ between 25 and 400 °C. Exothermic and endothermic events are indicated by positive and negative heat flow, respectively.

4.6. Quantum-Chemical Computations. All quantum-chemical computations were performed using Gaussian 16 software.$^{40}$ Structural optimization and harmonic normal mode vibrational calculations were performed using the hybrid DFT method B3LYP$^{15,42}$ together with Grimme’s version 3 dispersion correction$^{43}$ and def2TZVP$^{44,45}$ basis set. The decomposition pathway was investigated by searching for possible stable structures as well as finding transition states connecting these structures. Minima were confirmed to have no imaginary frequencies, while transition states were verified to have one imaginary frequency, lying along the reaction path.

ASSOCIATED CONTENT

+ Supporting Information
  The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.inorgchem.0c03496.
  Characterization of the compounds and computational calculation details (PDF)
  Compound 1 (CIF)
  Compound 6 (CIF)

Accession Codes
  CCDC 2046808 contains the supplementary crystallographic data for this paper. These data can be obtained free of charge via www.ccdc.cam.ac.uk/data_request/cif, or by emailing data_request@ccdc.cam.ac.uk, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44 1223 336033.

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Notes
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REFERENCES

(1) Taniyasu, Y.; Kasu, M.; Makimoto, T. An Aluminium Nitride Light-Emitting Diode with a Wavelength of 210 Nanometres. Nature 2006, 441, 325–328.
(2) Li, J.; Nam, K. B.; Nakarmi, M. L.; Lin, J. Y.; Jiang, H. X.; Carrier, P.; Wei, S.-H. Band Structure and Fundamental Optical Transitions in Wurtzite AlN. Appl. Phys. Lett. 2003, 83, 5163–5165.
(3) George, S. M. Atomic Layer Deposition: An Overview. Chem. Rev. 2010, 110, 111–131.
(4) Koponen, S. E.; Gordon, P. G.; Barry, S. T. Principles of Precursor Design for Vapour Deposition Methods. Polyhedron 2016, 108, 59–66.
(5) Van Bui, H.; Nguyen, M. D.; Wiggers, F. B.; Aarnink, A. A. I.; De Jong, M. P.; Kovalgin, A. Y. Self-Limiting Growth and Thickness-And-Deposition Temperature-Dependence of Optical Constants of ALD AlN Thin Films. ECS J. Solid State Sci. Technol. 2014, 3, P101–P106.
(6) Rihelä, D.; Ritala, M.; Matero, R.; Leskelä, M.; Jokinen, J.; Haussalo, P. Low Temperature Deposition of AlN Films by an Alternate Supply of Trimethyl Aluminum and Ammonia. Chem. Vap. Deposition 1996, 2, 277–283.
(7) Liu, X.; Ramanathan, S.; Lee, E.; Seidel, T. E. Atomic Layer Deposition of Aluminum Nitride Thin Films from Trimethyl Aluminum (TMA) and Ammonia. MRS Online Proceedings Library 2003, 811, 158–153.
(8) Jung, Y. C.; Hwang, S. M.; Le, D. N.; Kondusamy, A. L. N.; Mohan, J.; Kim, S. W.; Kim, J. H.; Lucero, A. T.; Ravichandran, A.; Kim, H. S.; Kim, S. J.; Choi, R.; Ahn, J.; Alvarez, D.; Spiegelman, J.; Kim, J. Low Temperature Thermal Atomic Layer Deposition of Aluminum Nitride Using Hydrazine as the Nitrogen Source. Materials 2020, 13, 3387.
(9) Dendooven, J.; Deduytsche, D.; Muschot, J.; Vanmeirhaeghe, R. L.; Detavernier, C. Conformality of Al2O3 and AlN Deposited by Plasma-Enhanced Atomic Layer Deposition. J. Electrochem. Soc. 2010, 157, G111.
(10) Ogist, C.; Donmez, I.; Alevi, M.; Biyikli, N. Self-Limiting Low-Temperature Growth of Crystalline AlN Thin Films by Plasma-Enhanced Atomic Layer Deposition. Thin Solid Films 2012, 520, 2750–2755.

(11) Ogist-Akgun, C.; Goldenberg, E.; Okyay, A. K.; Biyikli, N. Hollow Cathode Plasma-Assisted Atomic Layer Deposition of Crystalline AlN, GaN and AlGaN-XN Thin Films at Low Temperatures. J. Mater. Chem. C 2014, 2, 2123–2126.

(12) Bosund, M.; Sajavaara, T.; Laitinen, M.; Huhtio, T.; Putkonen, M.; Airaksinen, V. M.; Lipsanen, H. Properties of AlN Grown by Plasma Enhanced Atomic Layer Deposition. Appl. Surf. Sci. 2011, 257, 7827–7830.

(13) Nepal, N.; Qadri, S. B.; Hite, J. K.; Mahadik, N. A.; Mastro, M. A.; Eddy, C. R. Epitaxial Growth of AlN Films via Plasma-Assisted Atomic Layer Epitaxy. Appl. Phys. Lett. 2013, 103, 82110.

(14) Waggner, K. M.; Ölmeštad, M. M.; Power, P. P. Structural and Spectroscopic Characterization of the Compounds [Al(NMe2)3][Ga(NMe2)3], [Me2N]2[Al{μ-N(H)-1-Ad}"2 (1-Ad: 1-Adaman-

(15) Wade, C. R.; Silvernail, C.; Banerjee, C.; Soulet, A.; McAndrew, J.; Belot, J. A. Tris(Dialkylamino)Aluminums: Syntheses, Characterization, Volatilization Comparison and Atomic Layer Deposition of Alumina Thin Films. Mater. Lett. 2007, 61, 5079–5082.

(16) Liu, G.; Deguns, E.; Lecordier, L.; Sundaram, G.; Becker, J. Atomic Layer Deposition of AlN with Tris(Dimethylamido) Aluminum and NH. ECS Trans. 2011, 41, 219–225.

(17) Kim, K. H.; Gordon, R. G.; Denton, A.; Antoniadis, D. A. Atomic Layer Deposition of Insulating Nitride Interfacial Layers for Germanium Metal Oxide Semiconductor Field Effect Transistors with High-κ Oxide/Tungsten Nitride Gate Stacks. Appl. Phys. Lett. 2007, 90, 212104.

(18) Abdulagatov, A. I.; Ramazanov, S. M.; Dallaev, R. S.; Murliev, E. K.; Palchaev, D. K.; Rabadanov, M. K.; Abdulagatov, I. M. Atomic Layer Deposition of Aluminum Nitride Using Tris(Diethylamido) Aluminum and Hydrazine or Ammonia. Russ. Microelectron. 2018, 47, 118–130.

(19) Abdulagatov, A. I.; Amashaev, R. R.; Ashurbekova, K. N.; Ashurbekova, K. N.; Rabadanov, M. K.; Abdulagatov, I. M. Atomic Layer Deposition of Aluminum Nitride Using Tris(Diethylamido) Aluminum and Hydrazine or Ammonia. Russ. Microelectron. 2018, 47, 118–130.

(20) Abdulagatov, A. I.; Amashaev, R. R.; Ashurbekova, K. N.; Ashurbekova, K. N.; Rabadanov, M. K.; Abdulagatov, I. M. Atomic Layer Deposition of Aluminum Nitride and Oxynitride on Silicon Using Tris(Dimethylamido)Aluminum, Ammonia, and Water. Russ. J. Gen. Chem. 2018, 88, 1690–1706.

(21) Gordon, R. G. Atomic Layer Deposition for Semiconductors. In Atomic Layer Deposition for Semiconductors; Hwang, C. S., Yoo, C. Y., Eds.; Springer US: New York, 2014; pp 15–46.

(22) Kim, S. B.; Jayaraman, A.; Chua, D.; Davis, L. M.; Zheng, S. L.; Han, J.; Jones, B. G.; Gomperts, R.; Mannucci, B.; Hratchian, H. P.; Ortiz, J. V.; Izmaylov, A. F.; Sonnenberg, J. L.; Williams-Young, D.; Ding, F.; Lipparini, F.; Egidi, F.; Goings, J.; Peng, B.; Petrone, A.; Henderson, T.; Ranasinghe, D.; Zakrzewski, V. G.; Gao, J.; Rega, N.; Zheng, G.; Liang, W.; Hada, M.; Ehrhard, M.; Toyoda, K.; Fukuda, R.; Hasegawa, J.; Ishida, M.; Nakajima, T.; Honda, Y.; Kitao, O.; Nakai, H.; Vreven, T.; Throssell, K.; Montgomery, J. A., Jr.; Petrala, J. E.; Ogliaro, F.; Beard, S.; Heyd, J. J.; Brodsky, E. N.; Kudin, K. N.; Staroverov, V. N.; Kobayashi, R.; Normand, J.; Raghavachari, K.; Rendell, A.; Burant, J. C.; Iyengar, S. S.; Tomasi, J.; Cossi, M.; Millam, J. M.; Klene, M.; Li, X.; Caricato, M.; Marenich, A. V.; Bloino, J.; Zheng, S. L.; Farkas, O.; Foresman, J. B.; Fox, D. J. Gaussian 16, revision B.01; Gaussian, Inc.: Wallingford CT, 2016.

(23) Frisch, M. J.; Trucks, G. W.; Schlegel, H. B.; Scuseria, G. E.; Robb, M. A.; Cheeseman, J. R.; Scalmani, G.; Barone, V.; Petersson, G. A.; Nakatsuji, H.; Li, X.; Caricato, M.; Marenich, A. V.; Bloino, J.; Janssen, B. G.; Gomperts, R.; Mennucci, B.; Hratchian, H. P.; Ortiz, J. V.; Izmaylov, A. F.; Sonnenberg, J. L.; Williams-Young, D.; Ding, F.; Lipparini, F.; Egidi, F.; Goings, J.; Peng, B.; Petrone, A.; Henderson, T.; Ranasinghe, D.; Zakrzewski, V. G.; Gao, J.; Rega, N.; Zheng, G.; Liang, W.; Hada, M.; Ehrhard, M.; Toyoda, K.; Fukuda, R.; Hasegawa, J.; Ishida, M.; Nakajima, T.; Honda, Y.; Kitao, O.; Nakai, H.; Vreven, T.; Throssell, K.; Montgomery, J. A., Jr.; Petrala, J. E.; Ogliaro, F.; Beard, S.; Heyd, J. J.; Brodsky, E. N.; Kudin, K. N.; Staroverov, V. N.; Kobayashi, R.; Normand, J.; Raghavachari, K.; Rendell, A.; Burant, J. C.; Iyengar, S. S.; Tomasi, J.; Cossi, M.; Millan, J. M.; Klene, M.; Adamo, C.; Cammi, R.; Ochterski, J. W.; Martin, R. L.; Morokuma, K.; Farkas, O.; Foresman, J. B.; Fox, D. J. Gaussian 16, revision B.01; Gaussian, Inc.: Wallingford CT, 2016.

(24) Hanson-Heine, M. W. D.; George, M. W.; Besley, N. A. Calculating Excited State Properties Using Kohn-Sham Density Functional Theory. J. Chem. Phys. 2013, 138, 064101.

(25) Lecklinder, T. Evaluation Engineering—Maintaining a Healthy Rhythm, 2011. https://www.evaluationengineering.com/home/article/13005576/maintaining-a-healthy-rhythm.
(44) Weigend, F.; Ahlrichs, R. Balanced Basis Sets of Split Valence, Triple Zeta Valence and Quadruple Zeta Valence Quality for H to Rn: Design and Assessment of Accuracy. *Phys. Chem. Chem. Phys.* **2005**, *7*, 3297−3305.

(45) Metz, B.; Stoll, H.; Dolg, M. Small-Core Multiconfiguration-Dirac-Hartree-Fock-Adjusted Pseudopotentials for Post-d Main Group Elements: Application to PbB and PbO. *J. Chem. Phys.* **2000**, *113*, 2563−2569.