

Homoleptic Perchlorophenyl “Ate” Complexes of Thorium(IV) and Uranium(IV)

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Abstract

Reaction of $\text{AnCl}_4(\text{DME})_n$ ($\text{An} = \text{Th}, n = 2; \text{U}, n = 0$) with 5 equiv of LiC_6Cl_5 in Et_2O resulted in the formation of homoleptic actinide-aryl “ate” complexes $[\text{Li}(\text{DME})_2(\text{Et}_2\text{O})]_2[\text{Li}(\text{DME})_2][\text{Th}(\text{C}_6\text{Cl}_5)_5]_3$ ([Li][**1**]) and $[\text{Li}(\text{Et}_2\text{O})_4][\text{U}(\text{C}_6\text{Cl}_5)_5]$ ([Li][**2**]). Similarly, reaction of $\text{AnCl}_4(\text{DME})_n$ ($\text{An} = \text{Th}, n = 2; \text{U}, n = 0$) with 3 equiv of LiC_6Cl_5 in Et_2O resulted in formation of heteroleptic actinide-aryl “ate” complexes $[\text{Li}(\text{DME})_2(\text{Et}_2\text{O})][\text{Li}(\text{Et}_2\text{O})_2][\text{ThCl}_3(\text{C}_6\text{Cl}_5)_3]$ ([Li][**3**]) and $[\text{Li}(\text{Et}_2\text{O})_3][\text{UCl}_2(\text{C}_6\text{Cl}_5)_3]$ ([Li][**4**]). Density functional calculations show that the $\text{An-C}_{\text{ipso}}$ σ -bonds are considerably more covalent for the uranium complexes vs. the thorium analogues, in line with past results. Additionally, good agreement between experiment and calculations is obtained for the $^{13}\text{C}_{\text{ipso}}$ NMR chemical shifts in [Li][**1**] and [Li][**3**]. The calculations demonstrate a deshielding by ca. 29 ppm from spin-orbit coupling effects originating at Th, which is a direct consequence of 5f orbital participation in the Th-C bonds.

Introduction

Homoleptic actinide alkyl complexes have proven to be excellent testbeds for the study of electronic structure and bonding within these enigmatic elements.¹⁻³ As a result, a number of homoleptic actinide alkyl complexes have been synthesized in the last 10 years.⁴⁻¹⁸ These complexes include $[\text{Li}(\text{DME})_n][\text{An}(\text{CH}_2\text{SiMe}_3)_5]$ ($\text{An} = \text{Th}, n = 2; \text{An} = \text{U}, n = 3$),^{8,10} reported by us, $[\text{Li}(\text{THF})_4][\text{Li}(\text{THF})_2][\text{UMe}_6]$,⁴ reported by Neidig and co-workers, and $[\text{U}(\text{CH}_2\text{Ph})_4]$,⁷ reported by Bart and co-workers. In contrast, homoleptic actinide aryl complexes remain exceedingly rare, in part because of their tendency to undergo facile *ortho* C-H bond activation.¹⁹⁻²¹ For example, the first structurally characterized homoleptic uranium aryl, $[\text{U}(\text{C}_6\text{H}_3\text{-}2,6\text{-(C}_6\text{H}_4\text{-}4\text{-}^t\text{Bu})_2)_3]$, was only isolated in 2016 by Arnold and co-workers. Its isolation required the use of $4\text{-}^t\text{BuC}_6\text{H}_4$ *ortho* substituents on the aryl ligand to prevent *ortho* C-H bond activation.²⁰ Even with the $4\text{-}^t\text{BuC}_6\text{H}_4$ substituents, however, $[\text{U}(\text{C}_6\text{H}_3\text{-}2,6\text{-(C}_6\text{H}_4\text{-}4\text{-}^t\text{Bu})_2)_3]$ still decomposes on standing at room temperature, specifically by activation of an *ortho* C-H bond located on the $4\text{-}^t\text{BuC}_6\text{H}_4$ substituent. Additionally, we and others have used “ate” complex formation to generate homoleptic An(IV) aryl complexes, including $[\text{Li}(\text{DME})_3]_2[\text{Th}(\text{C}_6\text{H}_5)_6]$ ¹⁹ and $[\text{Li}(\text{THF})_4][(\text{THF})\text{LiU}(\text{C}_6\text{H}_5)_6]$.²² These complexes are stabilized by saturation of the actinide coordination sphere, which presumably disfavors the *ortho* C-H bond activation reaction.¹⁹ Nonetheless, $[\text{Li}(\text{THF})_4][(\text{THF})\text{LiU}(\text{C}_6\text{H}_5)_6]$ is still exceptionally thermally sensitive, and could only be isolated at $-80\text{ }^\circ\text{C}$.²² The use of pendant donor groups can also stabilize An-C_{aryl} bonds, as was observed for $[\text{Th}(\eta^2\text{-}N,C\text{-C}_6\text{H}_4\text{-}o\text{-CH}_2\text{NMe}_2)_4]$,²¹ $[\text{Cp}^*\text{U}(\eta^2\text{-}N,C\text{-}(o\text{-C}_6\text{H}_4)\text{NPh})]$,²³ and $[\text{Cp}^*\text{U}(\eta^2\text{-}Te,C\text{-}(o\text{-C}_6\text{H}_4)\text{Te})]$.²⁴

Many of these actinide aryl complexes are attractive targets for studying covalency via ¹³C NMR spectroscopy. NMR spectroscopy has been previously used as a tool to evaluate covalency

in An-L and Ln-L bonding.^{19,25–33} In the case of actinide aryl complexes, the ¹³C NMR chemical shifts for the C_{ipso} environments are strongly affected by spin-orbit (SO) coupling, a consequence of 5f (and to a lesser extent, 6d) orbital participation in the An-C σ-bond.²⁵ In particular, in 5f⁰ systems with low-lying, empty 5f or 6d orbitals, SO coupling tends to cause ¹³C deshielding as long as the An-C σ-bond has sizable C 2s character.³⁴ For example, the ¹³C NMR spectrum of [Li(DME)₃]₂[Th(C₆H₅)₆] features a C_{ipso} chemical shift of 220.5 ppm, with a calculated 44 ppm downfield contribution from SO coupling.¹⁹ This SO deshielding correlates to the 5f (and 6d) participation in the Th-C bond. In the case of [Th(C₆H₅)₆]²⁻, specifically, the Th-C bonds were calculated to have 15% weight each from Th atomic orbitals (AOs), of which 71/15% was from 6d/5f participation, respectively.

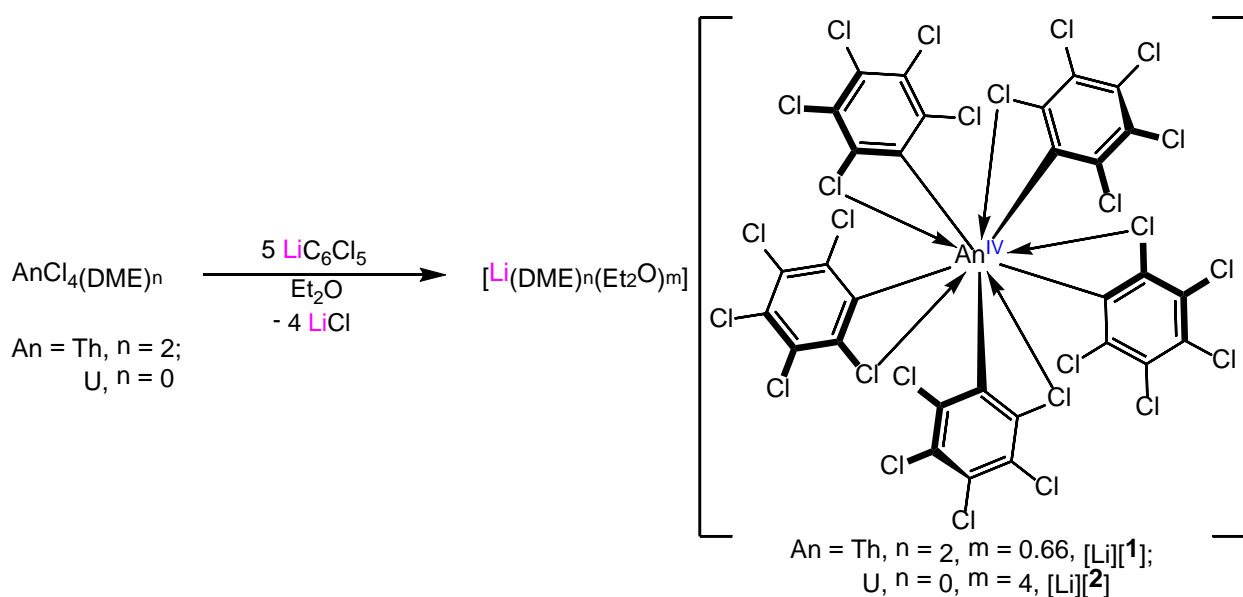
We recently reported the syntheses of the first structurally characterized uranyl(VI) aryl complexes, [Li(Et₂O)₂(THF)][UO₂(C₆Cl₅)₃] and [Li(THF)₄][UO₂(C₆Cl₅)₃(THF)], via reaction of [UO₂Cl₂(THF)₂] with 3 equiv of LiC₆Cl₅. These complexes feature surprisingly good thermal stability,²⁵ which we ascribed, in part, to the *ortho*-Cl substitution of the [C₆Cl₅]⁻ ligand. Given this precedent, we rationalized that the [C₆Cl₅]⁻ ligand could also stabilize homoleptic U(IV) and Th(IV) complexes. Additional support for this hypothesis comes from large number of homoleptic transition metal perhalophenyl complexes that have been reported over the past 25 years.^{35–44} Despite these past synthetic achievements, however, no homoleptic perhalophenyl complexes are known for actinides. Additionally, the only heteroleptic perhalophenyl actinide complexes are the aforementioned uranyl(VI) species,²⁵ making this a potentially fruitful avenue of investigation. Herein, we describe the synthesis and characterization of two homoleptic actinide-aryl “ate” complexes [Li(DME)₂(Et₂O)]₂[Li(DME)₂][Th(C₆Cl₅)₅] ([Li][**1**]) and [Li(Et₂O)₄][U(C₆Cl₅)₅] ([Li][**2**]). During the course of these investigations, we also isolated the closely related heteroleptic

actinide-aryl “ate” complexes, $[\text{Li}(\text{DME})_2(\text{Et}_2\text{O})][\text{Li}(\text{Et}_2\text{O})_2][\text{ThCl}_3(\text{C}_6\text{Cl}_5)_3]$ ($[\text{Li}][\mathbf{3}]$) and $[\text{Li}(\text{Et}_2\text{O})_3][\text{UCl}_2(\text{C}_6\text{Cl}_5)_3]$ ($[\text{Li}][\mathbf{4}]$). The electronic structures of all four complexes were analyzed using relativistic density functional theory (DFT) calculations, with the aim of quantifying the 5f subshell participation in the An-C_{ipso} bonds.

Results and Discussion

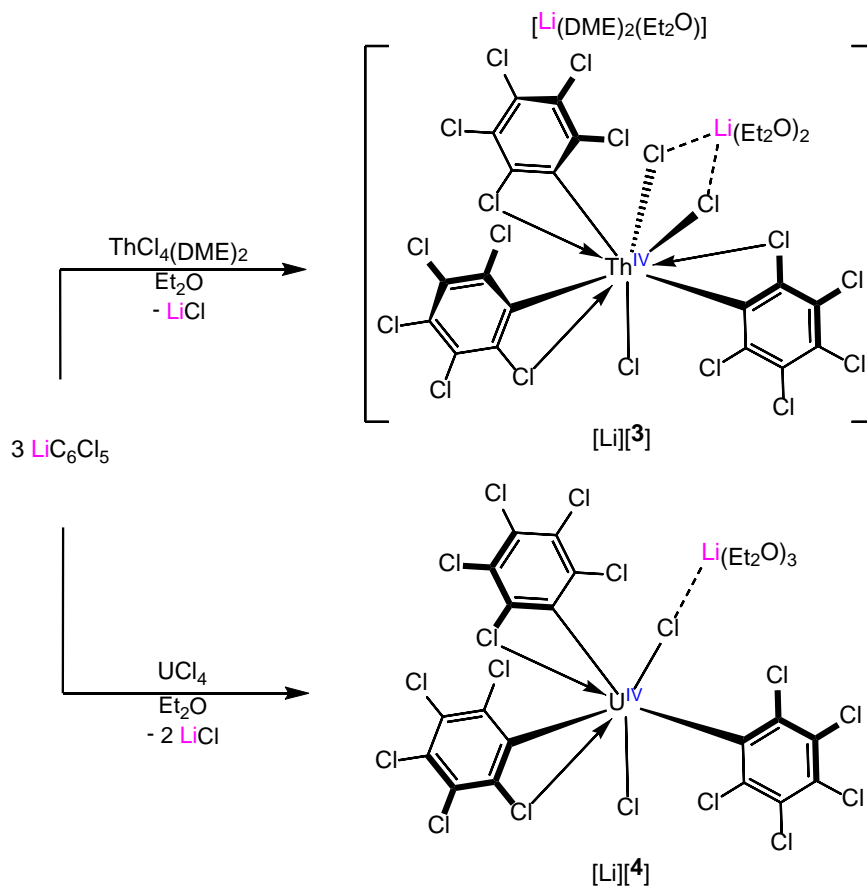
Addition of cold (-25 °C) solutions of 5 equiv of LiC_6Cl_5 ⁴⁵ to cold (-25 °C) suspensions of $\text{AnCl}_4(\text{DME})_n$ (An = Th, $n = 2$; U, $n = 0$) in Et_2O results in immediate formation of orange solutions, concomitant with the deposition of flocculent brown-orange precipitates. Work-up of the thorium reaction mixture, followed by crystallization from dichloromethane, affords $[\text{Li}(\text{DME})_2(\text{Et}_2\text{O})]_2[\text{Li}(\text{DME})_2][\text{Th}(\text{C}_6\text{Cl}_5)_3]$ ($[\text{Li}][\mathbf{1}]$), which can be isolated as colorless plates in 20% yield (Scheme 1). Work-up of the uranium reaction mixture, followed by crystallization from Et_2O , affords $[\text{Li}(\text{Et}_2\text{O})_4][\text{U}(\text{C}_6\text{Cl}_5)_5]$ ($[\text{Li}][\mathbf{2}]$), which can be isolated as yellow plates in 57% yield (Scheme 1). Interestingly, we see no evidence for the formation of octahedral $[\text{An}(\text{C}_6\text{Cl}_5)_6]^{2-}$ -type complexes, even when 6 equiv of $\text{Li}(\text{C}_6\text{Cl}_5)$ are used in the reaction.

Scheme 1.



Curiously, the reaction of $\text{ThCl}_4(\text{DME})_2$ with 4 equiv of LiC_6Cl_5 , performed in an attempt to make the neutral perchlorophenyl complex, $[\text{Th}(\text{C}_6\text{Cl}_5)_4]$, resulted in isolation of heteroleptic Th aryl complex, $[\text{Li}(\text{DME})_2(\text{Et}_2\text{O})][\text{Li}(\text{Et}_2\text{O})_2][\text{ThCl}_3(\text{C}_6\text{Cl}_5)_3]$ ($[\text{Li}][\mathbf{3}]$), in low yield, according to an X-ray crystallographic analysis of the crystals isolated upon work-up. Complex $[\text{Li}][\mathbf{3}]$ can be made rationally by addition of 3 equiv of LiC_6Cl_5 to a cold ($-25\text{ }^\circ\text{C}$) suspension $\text{ThCl}_4(\text{DME})_2$ in Et_2O . When synthesized using this stoichiometry, $[\text{Li}][\mathbf{3}]$ can be isolated as colorless plates in 54% yield (Scheme 2). Similarly, addition of 3 equiv of LiC_6Cl_5 to a cold ($-25\text{ }^\circ\text{C}$) suspension UCl_4 in Et_2O results in formation of $[\text{Li}(\text{Et}_2\text{O})_3][\text{UCl}_2(\text{C}_6\text{Cl}_5)_3]$ ($[\text{Li}][\mathbf{4}]$), which can be isolated as amber plates in 40% yield after work-up (Scheme 2). The reactions of UCl_4 with 4 equiv of LiC_6Cl_5 also results in formation of $[\text{Li}][\mathbf{4}]$, in similar yields to the 3 equiv reaction.

Scheme 2.



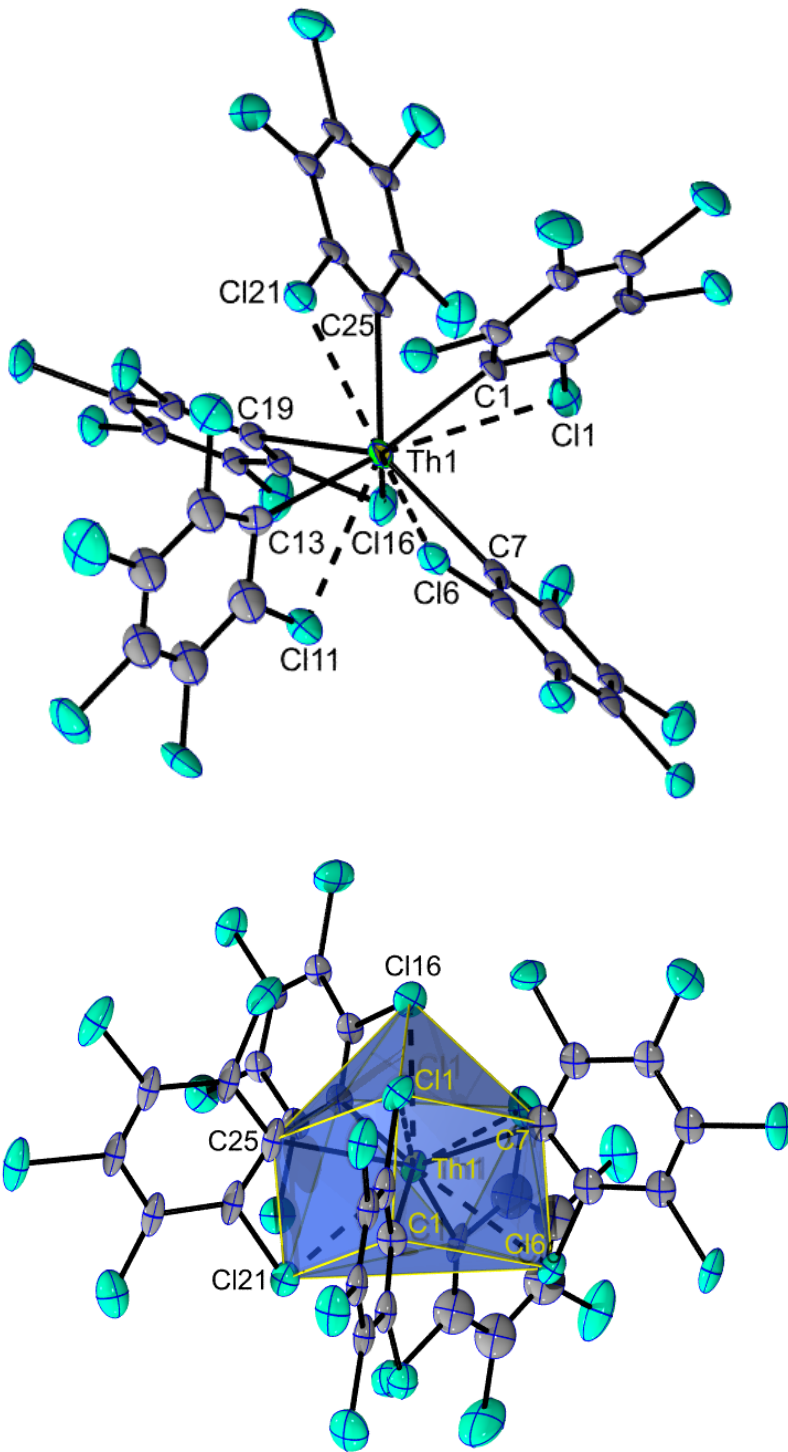


Figure 1. Solid-state molecular structure of $[\text{Li}][\mathbf{1}] \cdot 2.5\text{Et}_2\text{O} \cdot 2\text{CH}_2\text{Cl}_2$ shown with 50% probability ellipsoids (top). Solid-state molecular structure of $[\text{Li}][\mathbf{1}] \cdot 2.5\text{Et}_2\text{O} \cdot 2\text{CH}_2\text{Cl}_2$ with the sphenocorona

polygon shown in blue (bottom). All hydrogen atoms, solvate molecules, two $[\text{Th}(\text{C}_6\text{Cl}_5)_5]^-$ moieties, and Li^+ counterions have been omitted for clarity.

Both $[\text{Li}][\mathbf{1}]$ and $[\text{Li}][\mathbf{2}]$ are air- and moisture-sensitive crystalline solids that are soluble in ethereal solvents, methylene chloride, and benzene, and slightly soluble in hexanes. Additionally, $[\text{Li}][\mathbf{1}]$ is soluble in pyridine and acetonitrile, however $[\text{Li}][\mathbf{2}]$ reacts immediately upon dissolution in pyridine, resulting in formation of an intractable dark brown solution. Similarly, dissolution of $[\text{Li}][\mathbf{2}]$ in acetonitrile results in immediate formation of an intractable orange-brown solution concomitant with precipitation of a dark brown solid. Both $[\text{Li}][\mathbf{1}]$ and $[\text{Li}][\mathbf{2}]$ exhibit moderate thermal stability in solution. For example, solutions of $[\text{Li}][\mathbf{1}]$ or $[\text{Li}][\mathbf{2}]$ in CH_2Cl_2 exhibit minimal evidence of decomposition after standing at room temperature for 1 h. However, on standing at room temperature for 24 h, solutions of $[\text{Li}][\mathbf{1}]$ in methylene chloride- d_2 exhibit partial conversion to $[\text{Li}][\mathbf{3}]$, according to $^{13}\text{C}\{^1\text{H}\}$ NMR spectroscopy (Figure S4), presumably via C-Cl bond activation of the solvent. In contrast, solutions of $[\text{Li}][\mathbf{2}]$ completely convert into intractable mixtures in methylene chloride- d_2 over this time frame (Figure S6). It is not immediately apparent why $[\text{Li}][\mathbf{2}]$ is more reactive than $[\text{Li}][\mathbf{1}]$, but we note that a similar reactivity pattern is observed for $[\text{An}(\text{C}_6\text{H}_5)_6]^{2-}$ and $[\text{Cp}^*_2\text{An}(\text{C}_6\text{H}_5)_2]$ ($\text{An} = \text{Th}, \text{U}$),^{19,22,46} where the uranium analogues exhibit greater thermal sensitivity than the thorium analogues.

Complex $[\text{Li}][\mathbf{1}]$ crystallizes in the monoclinic space group $\text{P}2_1/\text{n}$ with three independent molecules in the asymmetric unit (Figure 1 and Table 1). It crystallizes as the Et_2O and CH_2Cl_2 solvate, $[\text{Li}][\mathbf{1}]\cdot 2.5\text{Et}_2\text{O}\cdot 2\text{CH}_2\text{Cl}_2$. Its asymmetric unit reveals one ten-coordinate thorium center and two nine-coordinate thorium centers. According to the continuous shape measure,⁴⁷ the 10-coordinate Th center, Th1, is best described as a C_{2v} -symmetric sphenocorona, wherein the two square faces are defined by C1, C7, C11, and C116, and C1, C25, C11, and C121, respectively. This

geometry is common for 10-coordinate complexes featuring bidentate ligands.⁴⁸ The two 9-coordinate metal centers are best described as distorted tricapped trigonal prisms.^{49,50} For Th2, the three capping atoms are C31, C43, and Cl46, whereas for Th3, the three capping atoms are C73, C85, and Cl51. The ten coordinate center in [Li][**1**] features five Th-C σ -bonds and five Cl \rightarrow Th dative interactions involving the *ortho*-Cl atoms of the C₆Cl₅ ligands, although one of these Cl \rightarrow Th dative interactions is quite long (Th1-Cl1 = 3.257(5) Å; see below for more discussion). The nine coordinate centers in [Li][**1**] are formed by five An-C σ -bonds and four Cl \rightarrow An dative interactions. The κ^2 coordination mode generated by the Cl \rightarrow An dative interaction has been previously observed for a handful of [C₆Cl₅]⁻ complexes, including Pt(C₆Cl₅)₄.^{38,42,43} The average Th-C bond length in [Li][**1**] is 2.65 Å (range = 2.628(18) – 2.693(18) Å), which is similar those found in other σ -bonded thorium aryl complexes.^{19,21} For example, the Th-C bond length in [Li(DME)₃]₂[Th(C₆H₅)₆]¹⁹ is 2.589(3) Å, whereas the average Th-C bond length in [Th(2-C₆H₄CH₂NMe₂)₄]²¹ is 2.549(2) Å (range = 2.497(3) – 2.544(3) Å). Additionally, both the 10-coordinate and 9-coordinate Th centers in [Li][**1**] exhibit identical average Th-C bond lengths. The aryl rings in [Li][**1**] feature disparate Th-C_{ipso}-C_{ortho} angles, a consequence of the Cl \rightarrow An dative interactions. For example, the Th-C1-C2 angle is 110(1)°, whereas the Th1-C1-C6 angle is 134(1)°. The average Cl \rightarrow Th bond length in [Li][**1**] is 3.09 Å (range = 3.018(5) – 3.257(5) Å). No other complexes with Cl \rightarrow An interactions are available for comparison, but several complexes with F \rightarrow An interactions are known.⁵¹ For example, the F \rightarrow U distances in [Cp*₂Co][U(OB(C₆F₅)₃)₂(^{Ar}acnac)(OEt₂)]⁵² are 2.762(6) and 2.789(5) Å, whereas the F \rightarrow U distances in [U(N(C₆F₅)₂)₄] are 2.6480(11) and 2.5989(11) Å.⁵³ Not surprisingly, these values are much shorter than the Cl \rightarrow An interactions observed in [Li][**1**]. One ten-coordinate and one nine-coordinate center in [Li][**1**] each exhibit an outer sphere cation [Li(DME)₂(Et₂O)]⁺, while the other

nine-coordinate thorium center in [Li][1] features a [Li(DME)₂]⁺ cation that also interacts with the *ortho*-Cl and *meta*-Cl atoms of one C₆Cl₅ ligand. The observation of two different Th coordination geometries in the solid-state can be explained by small differences in local crystal packing, and indicates that the long Cl→Th dative interaction (e.g., Th1-Cl1) is not particularly strong.

Complex [Li][2] crystallizes in the monoclinic space group P2₁/c (Figure S15). The solid-state molecular structure of [Li][2] reveals a ten-coordinate uranium center formed by five An-C σ -bonds and five Cl→An dative interactions involving the *ortho*-Cl atoms of the C₆Cl₅ ligands.⁴⁸ According to the continuous shape measure,⁴⁷ the coordination geometry about the U center is best described as a C_{2v}-symmetric sphenocorona, wherein the two square faces are defined by Cl16, C13, Cl6, and Cl1, and C13, C25, Cl11, Cl16, respectively. The average U-C bond length in [Li][2] is 2.55 Å (range = 2.52(2) – 2.59(2) Å), which is shorter than the Th-C distances in [Li][1], consistent with smaller ionic radius of the U(IV) ion.⁵⁴ The U-C bond lengths in [Li][2] are also similar to those observed in other uranium aryl complexes, such as [Li(Et₂O)₂(THF)][UO₂(C₆Cl₅)₃],²⁵ which features U-C distances from 2.55(1) to 2.63(1) Å, and [Li(THF)₄][Li(THF)][U(C₆H₅)₆],²² which features an average U-C bond length of 2.52 Å. The latter structure also features H→U agostic interactions, which are similar to the Cl→An interactions found in [Li][1] and [Li][2]. The aryl rings in [Li][2] feature disparate U-C_{ipso}-C_{ortho} angles (e.g., U1-C1-C2 = 113(1)°, U1-C1-C6 = 135(1)°), which is a consequence of the Cl→An dative interactions. Finally, the average Cl→U distance in [Li][2] is 3.13 Å (range = 3.006(4) – 3.250(4) Å), which is similar to that found in [Li][1].

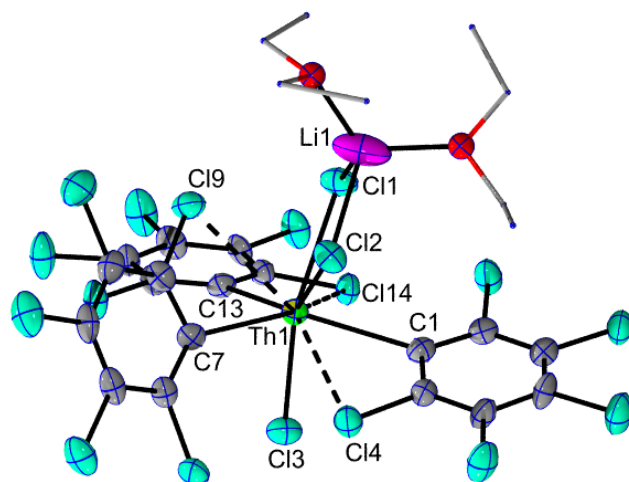


Figure 2. Solid-state molecular structure of [Li][**3**] shown with 50% probability ellipsoids. All hydrogen atoms and a [Li(DME)₂(Et₂O)]⁺ counterion have been omitted for clarity.

Complex [Li][**3**] crystallizes in the triclinic space group *P1* (Figure 2). Its solid-state molecular structure reveals a 9-coordinate thorium center bound by three chloride ligands, three carbon atoms of the pentachlorophenyl ligands, and three Cl→Th dative interactions involving the *ortho*-Cl atoms of the pentachlorophenyl ligands. The coordination geometry about its Th center can be described as distorted tricapped trigonal prism,⁵⁵ wherein the three capping atoms are Cl14, Cl19, and Cl14. The average Th-C bond length in [Li][**3**] is 2.65 Å (range = 2.631(6) – 2.654(6) Å), which is similar to Th-C bond lengths in [Li][**1**]. The average Th-Cl distance in [Li][**3**] is 2.77 Å (range = 2.671(2) – 2.839(2) Å), which is slightly longer than the average Th-Cl bond length in ThCl₄(DME)₂ (2.690 Å),⁵⁶ whereas the average Cl→An dative interactions in [Li][**3**] is 3.19 Å (range = 3.148(2) – 3.248(2) Å). Finally, the [Li(Et₂O)₂]⁺ cation in [Li][**3**] features bridging interaction with two chloride ligands. The resulting Li-Cl distances are 2.37(2) Å and 2.39(2) Å.

Complex [Li][4] also crystallizes in the triclinic space group $P1$ (Figure 3). Its solid-state molecular structure reveals a 7-coordinate uranium center bound by two chloride ligands, three carbon atoms of the pentachlorophenyl ligand, and two long Cl→U dative interactions involving the *ortho*-Cl atoms of the pentachlorophenyl ligands (e.g., U1-Cl3 = 3.226(4), U1-Cl8 = 3.227(3) Å). Ignoring the long Cl→U dative interactions, the coordination geometry about the U center is best described as trigonal bipyramid. The average U-C bond length in [Li][4] is 2.50 Å (range = 2.50(1) – 2.51(1) Å) and is similar to that seen in [Li][2] but shorter than that seen in [Li][3], consistent with smaller ionic radius of U(IV) ion.⁵⁴ The average U-Cl bond length in [Li][4] is 2.59 Å (range = 2.591(4) – 2.594(4) Å), consistent with the average U-Cl bond length of 2.62 Å in $\text{UCl}_4(\text{DME})_2$.⁵⁷ Finally, the $[\text{Li}(\text{Et}_2\text{O})_3]^+$ cation in [Li][4] features a bridging interaction with a chloride ligand. The resulting Li-Cl distance is 2.69(5) Å.

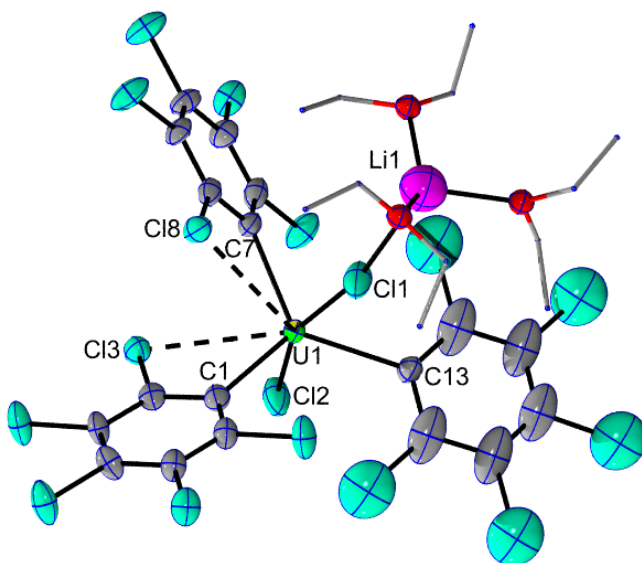


Figure 3. Solid-state molecular structure of [Li][4] shown with 50% probability ellipsoids. All hydrogen atoms have been omitted for clarity.

Table 1. Selected metrical parameters for [Li][1]·2.5Et₂O·2CH₂Cl₂, [Li][2], [Li][3], and [Li][4].

Complex	[Li][1]·2.5Et ₂ O·2CH ₂ Cl ₂			[Li][2]	[Li][3]	[Li][4]
An-C	Th1	Th2	Th3	2.518(17)	2.631(6)	2.497(13)
				2.537(15)	2.653(6)	2.504(13)
	2.633(16)	2.628(18)	2.631(18)	2.541(12)	2.654(6)	2.505(14)
	2.646(17)	2.637(17)	2.641(18)	2.555(17)		
	2.649(18)	2.639(16)	2.645(17)	2.592(16)		
	2.670(19)	2.656(19)	2.66(2)			
	2.673(17)	2.678(19)	2.693(18)			
Cl→An	3.018(5)	3.033(5)	3.073(5)	3.006(4)	3.1489(19)	3.226(4)
	3.068(5)	3.064(5)	3.092(6)	3.071(5)	3.1776(17)	3.227(3)
	3.073(5)	3.069(4)	3.115(5)	3.121(4)	3.2477(17)	
	3.101(4)	3.107(5)	3.122(5)	3.197(4)		
	3.257(5)			3.255(4)		
An-Cl					2.6702(17)	2.591(4)
					2.7887(18)	2.594(4)
					2.8390(18)	

The ¹³C{¹H} NMR spectrum of [Li][1], recorded at room temperature in dichloromethane-*d*₂, features a resonance at 198.78 ppm (Figure S2), attributable to the *ipso* carbon of the pentachlorophenyl ligand, as well as resonances at 138.66, 131.03, and 130.13 ppm, assignable to the *ortho*, *meta*, and *para* resonances of the pentachlorophenyl ligand, respectively. The ¹³C{¹H} NMR spectrum of [Li][3], recorded at room temperature in dichloromethane-*d*₂, features a resonance at 201.01 ppm (Figure S9), attributable to the *ipso* carbon of the pentachlorophenyl ligand, as well as resonances at 137.20, 131.96, and 129.53 ppm, assignable to the *ortho*, *meta*, and *para* resonances of the pentachlorophenyl ligand, respectively. The observation of only one aryl environment for both [Li][1] and [Li][3] in their ¹³C{¹H} NMR spectra is evidence for exchange of the Cl→Th dative interactions at a rate faster than the NMR time scale, which

renders the aryl ligands magnetically equivalent. The $^{13}\text{C}\{^1\text{H}\}$ NMR spectra of both $[\text{Li}][\mathbf{1}]$ and $[\text{Li}][\mathbf{3}]$ feature smaller downfield C_{ipso} resonances compared to $[\text{Li}(\text{Et}_2\text{O})_2(\text{THF})][\text{UO}_2(\text{C}_6\text{Cl}_5)_3]$, which featured a C_{ipso} resonance at 236.7 ppm.²⁵

The $^7\text{Li}\{^1\text{H}\}$ NMR spectra for $[\text{Li}][\mathbf{1}]$ and $[\text{Li}][\mathbf{3}]$ in dichloromethane- d_2 feature sharp resonances at -0.79 ppm and -0.83 ppm, respectively (Figures S1 and S8). In contrast, the $^7\text{Li}\{^1\text{H}\}$ NMR spectrum of $[\text{Li}][\mathbf{2}]$ in dichloromethane- d_2 features a broad resonance at 0.52 ppm (Figure S5), consistent with the paramagnetism of this material. The $^7\text{Li}\{^1\text{H}\}$ NMR spectrum of $[\text{Li}][\mathbf{4}]$ in benzene- d_6 features a broad resonance at 29.59 ppm (Figure S11). The large downfield shift is consistent with formation of a contact ion pair in solution. Similar behavior has been observed previously for uranium aryl and uranium benzyne complexes.⁵⁸

To gain a detailed understanding of the electronic structures and bonding interactions within $[\text{Li}][\mathbf{1}]$, $[\text{Li}][\mathbf{2}]$, $[\text{Li}][\mathbf{3}]$, and $[\text{Li}][\mathbf{4}]$, relativistic DFT calculations were performed on their anionic components, namely, $[\mathbf{1}]^-$, $[\mathbf{2}]^-$, $[\mathbf{3}]^-$, and $[\mathbf{4}]^-$.⁵⁹⁻⁶¹ Full computational details are described in the Supporting Information. The ground states of $[\mathbf{1}]^-$ and $[\mathbf{3}]^-$ are closed-shell spin singlet configurations, whereas the ground states of $[\mathbf{2}]^-$ and $[\mathbf{4}]^-$ are spin triplets because of the two unpaired electrons of the U^{4+} ion. As shown previously by wavefunction calculations,^{62,63} the ground state for Th(IV) is not strongly multi-configurational, and can therefore be described by (single determinant) Kohn-Sham DFT calculations with approximate functionals. Further evidence for the lack of multi-configurational character in the present systems are the large HOMO and LUMO gaps calculated for $[\mathbf{1}]^-$ and $[\mathbf{3}]^-$, at 3.0 and 3.4 eV, respectively. In other words, there are no complications due to the presence of nearly degenerate frontier orbitals in the DFT calculations for these molecules. Natural localized molecular orbital (NLMO) analysis (Figure 4 and Table S1) confirms the $\text{Cl}\rightarrow\text{An}$ dative bonding from the close-contact chlorine atoms of the

C₆Cl₅ ligands that is evident from the crystal structures. The lone pair donation from these chlorines have on average 7/8/6/5% weight at the actinide center in [1/2/3/4]⁻, respectively. Not surprisingly, however, the dative bonding from the chloride ligands in [3]⁻ and [4]⁻ is stronger, with an average of 12 and 14% weight, respectively, at the actinide.

The NLMO analysis further shows that the Th-C_{ipso} interactions in [1]⁻ and [3]⁻ are quite similar (Figure 4). This similarity is also indicated by averaged Th-C_{ipso} Wiberg bond orders of 0.47 for [1]⁻ and 0.49 for [3]⁻, respectively. The interactions between the *ipso*-carbon atoms and thorium can be viewed as two-center two-electron (2c-2e) σ -donation bonding from the negatively charged ligand, with the Th weight in the corresponding σ -bonding NLMOs ranging from 14 to 15%. Of these weights, the Th 5f contributions are 24% in [1]⁻ and 22% in [3]⁻. In comparison, the two uranium complexes, [2]⁻ and [4]⁻, feature stronger covalency in their U-C bonds, along with more pronounced participation of the 5f AOs. The U weights in the σ -bonding NLMOs range from 18% in [2]⁻ to 21% in [4]⁻, of which the 5f contributions are 36% and 30%, respectively. The increased covalency, compared to the Th complexes, is reflected in increased averaged U-C_{ipso} Wiberg bond orders of 0.56 in [2]⁻ and 0.64 in [4]⁻, respectively. Similar bonding trends are observed for both [An(C₆H₅)₆]²⁻ (An = Th, U) and [AnMe₆]²⁻ (An = Th, U).^{4,10,22} For example, the An-C bonds in [U(C₆H₅)₆]²⁻ feature both a greater degree of covalency (23% total metal character for U vs. 16% for Th) and 5f character (38% for U vs. 20% for Th) than its Th analogue.²²

Various approximate density functionals (BP86, PBE, and PBE0) were used for calculating the ¹³C NMR chemical shifts for the *ipso*-carbon in [1]⁻ and [3]⁻. As seen in Table 2, the averaged calculated chemical shifts for both complexes are only weakly dependent on the choice of the functional. The calculated chemical shifts are in acceptable agreement with the measurements. We focus on the PBE/SO-PBE results in the following discussion, unless stated otherwise. For [1]⁻,

the calculated C_{ipso} chemical shift is 203 ppm (expt. = 199 ppm), including a 27 ppm deshielding due to SO coupling. The C_{ipso} chemical shift in $[3]^-$ is calculated to be a bit larger than that in $[1]^-$, in agreement with the experiments, although the shift difference is slightly exaggerated in the calculations. The deshielding caused by SO coupling effects (ca. 29 ppm) is comparable that found in $[1]^-$, which reflects the similarity in their chemical bonding. The *ipso*-carbon SO deshielding in both $[1]^-$ and $[3]^-$ is correlated to the Th 5f and 6d character of the $\sigma(\text{Th-C})$ NLMO, and the overall weights of actinide AOs in these bonds (i.e., the degree of covalency). Our previous calculation²⁵ for $[\text{Li}(\text{Et}_2\text{O})_2(\text{THF})][\text{UO}_2(\text{C}_6\text{Cl}_5)_3]$ found a significantly larger SO deshielding, up to 68 ppm, due to a combination of large U weight (20% overall) and 5f contribution (43% of the U weight). This difference reflects both the change in element and the higher oxidation state in the uranyl example. Similarly, we would expect stronger SO effects for the *ipso*-carbon chemical shifts in our U(IV) complexes, compared to their thorium counterparts. However, the paramagnetism of $[2]^-$ and $[4]^-$ causes too much broadening of the relevant NMR signals for them to be observed. It also renders calculations of the NMR chemical shifts much more difficult,⁶⁴ and the role of SO coupling would not be as clear-cut as it is in the diamagnetic counterparts. Thus, we decided to forego calculations of the shifts for $[2]^-$ and $[4]^-$, because the bond analyses already paint a clear picture. Interestingly, the SO-induced chemical shift in $[\text{Th}(\text{C}_6\text{H}_5)_6]^{2-}$ (40 ppm, present work) is calculated to be 13 ppm larger than that of $[1]^-$ (27 ppm), despite the former complex having a smaller 5f weight in the Th contributions to the Th-C bonds. This decrease, however, is more than compensated for by the slightly greater overall Th weight and carbon 2s character in its Th-C bonds (Table 2 and Table S1).

Table 2. Calculated carbon shielding (σ) and chemical shift (δ) for the C_{ipso} atom center of $[1]^-$, $[3]^-$, and $[\text{Th}(\text{C}_6\text{H}_5)_6]^{2-}$, using various functionals.^a

Complex	Functional	σ_{calc} (ppm)	δ_{calc} (ppm)	δ_{expt} (ppm)
TMS	BP86/SO-BP86	186.9 / 187.8	-	-
	PBE/SO-PBE	187.5 / 188.4	-	
	PBE0/SO-PBE0 (25%) ^b	192.2 / 193.0	-	
	PBE0/SO-PBE0 (40%) ^b	194.7 / 195.6	-	
[1] ⁻	BP86/SO-BP86	10.7 / -16.8	176.2 / 204.6	198.78
	PBE/SO-PBE	11.3 / -14.8	176.2 / 203.2	
	PBE0/SO-PBE0 (25%)	13.9 / -10.8	178.3 / 203.8	
	PBE0/SO-PBE0 (40%)	15.8 / -8.3	178.9 / 203.9	
[3] ⁻	BP86/SO-BP86	3.9 / -25.3	183.0 / 213.1	201.01
	PBE/SO-PBE	4.5 / -23.3	183.0 / 211.7	
	PBE0/SO-PBE0 (25%)	5.5 / -20.8	186.7 / 213.8	
	PBE0/SO-PBE0 (40%)	6.8 / -19.0	187.9 / 214.6	
[Th(C ₆ H ₅) ₆] ²⁻ ^c	PBE/SO-PBE	-21.2 / -60.1	208.7 / 248.5	220.5
	PBE0/SO-PBE0 (40%)	-15.5 / -53.4	210.2 / 249.0	

^a NMR shifts calculated at the SO ZORA- level using the TZ2P basis set, with 'FXC' option. The calculated chemical shifts were averaged over equivalent C_{ipso} nuclei. Dichloromethane (solvent) was considered in the COSMO model for all the computations.

^b The percentages in parentheses of functional column indicate the portion of exact exchange in the functional.

^c For comparison, the carbon shielding and chemical shift for the C_{ipso} atom in C₃-symmetric [Th(C₆H₅)₆]²⁻ was re-calculated using the same procedure as other complexes, such as using the 'FXC' option and same solvent (see computational details). The experimental chemical shift for this complex was taken from Ref. ¹⁹.

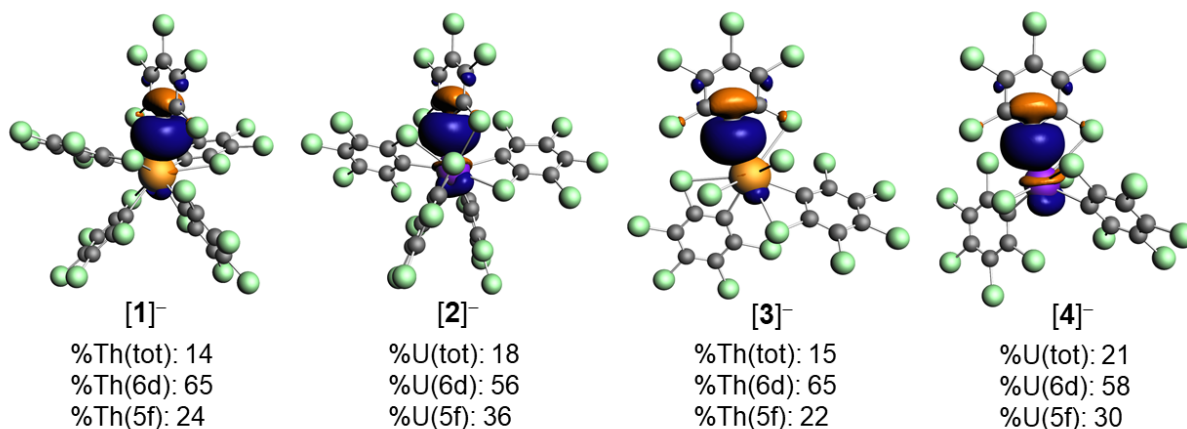


Figure 4. Representative An-C bonding NLMOs (An=Th or U) in [1]⁻, [2]⁻, [3]⁻, and [4]⁻. Weight-% metal character and 6d vs. 5f contributions at the metal averaged over equivalent NLMOs. (Isosurface values ±0.03 a.u. Color code for atoms: Th orange, U purple, Cl green, C gray.)

Conclusion

In summary, we have prepared and characterized two homoleptic actinide-aryl “ate” complexes, [Li(DME)₂(Et₂O)]₂[Li(DME)₂][Th(C₆Cl₅)₅]₃ and [Li(Et₂O)₄][U(C₆Cl₅)₅], and have confirmed their formulation by X-ray crystallography. These two complexes represent the first isolated homoleptic perhaloaryl complexes of the actinides. They exhibit remarkable thermal stability – much greater than past homoleptic actinide aryl complexes – likely on account of the *o*-chloro substitution of the [C₆Cl₅]⁻ ligand, combined with the many Cl→An dative interactions. Additionally, we prepared and characterized two heteroleptic actinide-aryl “ate” complexes, [Li(DME)₂(Et₂O)][Li(Et₂O)₂][ThCl₃(C₆Cl₅)₃] and [Li(Et₂O)₃][UCl₂(C₆Cl₅)₃]. Analysis of the An-C bonding with these complexes by DFT reveals both greater covalency and greater 5f orbital participation in the U(IV) derivatives, consistent with past periodic trends. In addition, a DFT analysis of the C_{ipso} chemical shifts in the two Th complexes reveals modest levels of spin-orbit induced shielding due to 5f participation in the Th-C bonds, providing us with an opportunity to further refine the ¹³C NMR spectroscopic analysis of An-L bonding.

ASSOCIATED CONTENT

Supporting Information. Experimental procedures, crystallographic details (as CIF files), computational results, and spectral data for complexes [Li][1]-[Li][4]. See DOI:

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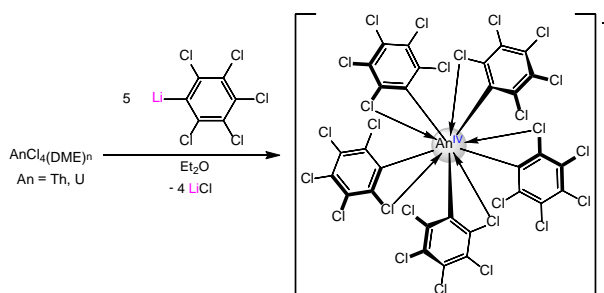
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The homoleptic actinide-aryl “ate” complexes, $[\text{Li}(\text{DME})_2(\text{Et}_2\text{O})]_2[\text{Li}(\text{DME})_2][\text{Th}(\text{C}_6\text{Cl}_5)_5]_3$ and $[\text{Li}(\text{Et}_2\text{O})_4][\text{U}(\text{C}_6\text{Cl}_5)_5]$, exhibit appreciable thermal stability, on account of the *o*-chloro substitution within the $[\text{C}_6\text{Cl}_5]^-$ ligand, combined with the many $\text{Cl} \rightarrow \text{An}$ dative interactions.