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Neptunium and plutonium complexes with a sterically encumbered triamidoamine (TREN) scaffold

Jessie L. Brown, Andrew J. Gaunt, David M. King, Stephen T. Liddle, Sean D. Reilly, Brian L. Scott, Ashley J. Wooles

The syntheses and characterisation of isostructural neptunium(IV) and plutonium(IV) complexes [An\textsuperscript{IV}(TREN\textsubscript{TIPS})(Cl)] [An = Np, Pu; TREN\textsubscript{TIPS} = (N(CH\textsubscript{3}CH\textsubscript{2}CH\textsubscript{2}NSiPr\textsubscript{3})\textsubscript{3}] are reported, along with the demonstration that they are likely reduced to the corresponding neptunium(III) and plutonium(III) products [An\textsuperscript{III}(TREN\textsubscript{TIPS})]; this chemistry provides new platforms from which to target a plethora of unprecedented molecular functionalities in transuranic chemistry and the neptunium(IV) molecule is the first structurally characterised neptunium(IV)-amide complex.

Non-aqueous molecular uranium chemistry has flourished in recent years, with the isolation of landmark complexes containing unprecedented actinide-ligand functionalities,\textsuperscript{1,7} with examples including terminal nitride, bis(imido), tris(imido), phosphinidene, arsini dine/arsenido, and terminal chalcogenide moieties.\textsuperscript{8-19} The insight afforded by these studies has profoundly enhanced our understanding of the ability of the valence 5f and 6d orbitals to engage in covalent interactions, which is most readily studied in metal-ligand multiple-bonding interactions. However, uranium is not the only 5f element of practical relevance and it is, perhaps, of more telling impact to decipher covalent bonding trends and redox differences across all of the relevant and synthetically available actinide elements, such as neptunium and plutonium.\textsuperscript{20-23} Due to the specialist radiological facilities required to safely handle the transuranium elements, progress pales compared to uranium.\textsuperscript{1,24} Nonetheless, very recently a neptunium(V)-bis(imido) complex was reported;\textsuperscript{25} this is the first example of a non-dioxo transuranic metal-ligand multiple-bond, suggesting that transuranic elements may also offer a rich reservoir of accessible multiple-bond chemistry.

In order to approach general accessibility of multiply-bonded functionalities systematically, a supporting scaffold to control and direct reactivity at a single coordination site is required. In this regard, various substituted triamidoamine (TREN) frameworks have facilitated a wide range of unprecedented uranium chemistry.\textsuperscript{2} For example, the terminal U=N triple bond was prepared by reduction of [U\textsuperscript{IV}(TREN\textsubscript{TIPS})(Cl)] to [U\textsuperscript{II}(TREN\textsubscript{TIPS})] (TREN\textsubscript{TIPS} = (N(CH\textsubscript{3}CH\textsubscript{2}CH\textsubscript{2}NSiPr\textsubscript{3})\textsubscript{3}), followed by oxidation of an azide at a vacant U\textsuperscript{II} coordination site to form a uranium(V)-nitride;\textsuperscript{8} the latter could be oxidised to give the corresponding uranium(V)-nitride.\textsuperscript{9} In a similar manner, a terminal uranium(V)-mono(oxo) species was prepared by oxidation of [U\textsuperscript{IV}(TREN\textsubscript{TIPS})] with Me\textsubscript{3}NO.\textsuperscript{20} Non-reductive reactivity studies with [U\textsuperscript{IV}(TREN\textsubscript{TIPS})(Cl)] has given a range of parent terminal U=EH (E = N, P, As) groups, a U\textsuperscript{V}AsAs arsenide and trapping of HASAs and cyclo-P\textsubscript{5}, and more broadly low-valent [U(TREN)] complexes have facilitated isolation of the first uranium-dinitrogen complex and reductive homologation of carbon monoxide to ethyne diolate.\textsuperscript{13,14,27-31} If the metal-ligand multiple-bond chemistry described above could be developed for transuranic elements it would greatly improve our knowledge and understanding of the electronic structure and reactivity of these elements.

**Scheme 1 Synthesis of complexes 1 and 2.**

Herein, we report coordination of TRE\textsubscript{TIPS} to neptunium(IV) and plutonium(IV) to afford [An\textsuperscript{IV}(TREN\textsubscript{TIPS})(Cl)] (An = Np, 1; Pu, 2). Moreover, NMR spectroscopy suggests that 1 and 2 can be reduced to putative [Np\textsuperscript{IV}(TREN\textsubscript{TIPS})] and [Pu\textsuperscript{IV}(TREN\textsubscript{TIPS})]; complexes that provide a suitable platform for oxidative pathways to new metal-ligand multiple-bonds. Dropwise treatment of [AnCl\textsubscript{2}(DME)]\textsubscript{2} (An = Np, Pu), dissolved in cold THF, with one equivalent of [(L\textsubscript{3})(TREN\textsubscript{TIPS})], also dissolved in cold THF, results in salt metathesis and formation of 1 and 2, respectively (Scheme 1). Isolation of 1 and 2 from hexanes solution

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\textsuperscript{d} Footnotes relating to the title and/or authors should appear here. Electronic Supplementary Information (ESI) available: [details of any supplementary information available should be included here]. See DOI: 10.1039/x0xx00000x

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afforded crystalline yields of 51% and 19%, respectively. The moderate, and low (but repeatable), rather than high isolated yields reflect the high product solubility on the small transuranic reaction scales. Single-crystals suitable for X-ray diffraction were grown from toluene/hexanes for 1 and hexanes for 2. The solid-state structures, Figure 1, reveal that 1 and 2 are isostructural to each other and also to the uranium(IV) congener.8 The actinide(IV) metal centres are five-coordinate, bound to three amide N atoms, one amine N atom, and a chloride anion completes the coordination sphere. The Np–Namide distances in 1 range from 2.206(2) to 2.232(2) Å, and are indistinguishable from the complexes (An=Th, U, Np, Pu) corresponding values in 1. The only other PuIV–amide bond that has been structurally determined is in [Pu(N(SiMe4)3)2(Cl)], with a value of 2.207(6) Å that is comparable to the range observed in 2.33 The Pu–Namide distance of 2.577(2) Å and Pu–Cl distance of 2.600(5) Å are both slightly shorter than the values in 1, again reflecting the actinide contraction. The Pu–Cl distance is also in the range of 2.5745(8) to 2.6033(8) Å observed for PuIV–Cl distances in the [PuCl4(DME)2] precursor.33 The ThIV analogue, [ThIV(TREN)2(Cl)], has been previously reported,34 meaning that isostructural complexes across the AnIV series of Th, U, Np, and Pu have been isolated and characterised – this is rare as few thorium and uranium studies are extended into the transuranic realm due to the specialist facilities required and also changes in redox stability from thorium to plutonium often impart difficulty in being able to isolate Th, U, Np and Pu in the same oxidation state in isostructural complexes. Table 1 (and ESI) provides a comparison of selected bond lengths demonstrating the general trend of shorter metal–ligand bond distances from left to right across the 5f series that is consistent with predominately ionic bonds and the higher charge density of the AnIV ions as the 5f series is traversed.

![Figure 1](image1.png)

**Figure 1** Thermal ellipsoid plot (at the 50% probability level) of the solid-state structure of [PuIV(TREN)2(Cl)] (2). Hydrogen atoms are omitted for clarity. The neptunium analogue, 1, is isostructural (see ESI for details).

**Table 1** Comparison of selected bond distances (Å) in [AnIV(TREN)2(Cl)] complexes (An = Th, U, Np, Pu).

<table>
<thead>
<tr>
<th></th>
<th>Th</th>
<th>U</th>
<th>Np</th>
<th>Pu</th>
</tr>
</thead>
<tbody>
<tr>
<td>An–Cl</td>
<td>2.693(2)</td>
<td>2.6383(6)</td>
<td>2.6200(6)</td>
<td>2.6001(5)</td>
</tr>
<tr>
<td>An–Namide</td>
<td>2.278(3)</td>
<td>2.224(2)</td>
<td>2.2305(19)</td>
<td>2.2221(17)</td>
</tr>
<tr>
<td>An–Namide</td>
<td>2.305(3)</td>
<td>2.215(2)</td>
<td>2.2311(18)</td>
<td>2.2257(17)</td>
</tr>
</tbody>
</table>

Turning to the PuIV complex, 2, the Pu–Namide distances range from 2.212(2) to 2.232(2) Å, and are indistinguishable from the values in 1. The optical absorption spectra reveal a mixture of relatively narrow 5f–5f, 5f–6d, and broader charge transfer bands. For 1, there is excellent correlation between the electronic transitions of the solution phase vis-NIR spectrum in hexanes and those in the solid-state, Figure 2, suggesting that the structure of 1 is maintained in solution (at least with a non-coordinating solvent). For 2, a very dark compound, the charge-transfer band extends past 800 nm and the solid-state spectrum is uninformative in the lower wavelength region; however, a correlation between the electronic transitions in the post-900 nm region is observed (see ESI) in terms of the profile, although the exact values do not match quite as closely as in 1.
Initial reactivity studies of 1 with benzyl potassium suggest that the reaction is slower than the Th\textsuperscript{IV} and U\textsuperscript{VI} congeners,\textsuperscript{38} possibly due to the increased steric encumbrance imparted by the TREN\textsuperscript{NPS} ligand around the smaller Np\textsuperscript{III} ion. Therefore, consideration was given to attempting the same reaction with the more labile iodide anion in place of chloride. NMR spectroscopic data suggests that halide-exchange is achieved upon treatment of 1 with Me\textsubscript{3}SiI in toluene, although under these initial reaction conditions benzyl potassium appears to be essentially unreactive (see ESI). Resonances for 1 disappeared with the emergence of new peaks at 15.97, 10.10, 9.17, and -37.75 ppm, tentatively assigned to [Np\textsuperscript{III}(TREN\textsuperscript{NPS})\textsubscript{3}]\textsuperscript{3+}, with integrals that correlate to the CH, CH(CH\textsubscript{3})\textsubscript{2}, CH(CH\textsubscript{2})\textsubscript{2} and CH protons, respectively. Although single-crystals of the iodide product were obtained, weak diffraction except at high angles prevented conclusive verification of the structure, and the possibility of a different reaction product cannot be entirely ruled out.

Stirring a hexanes solution of 1 or 2 over a potassium mirror for several days results in a colour change, formation of a colourless precipitate (presumed to be KCl), suggestive of reduction of Np\textsuperscript{V} to An\textsuperscript{III}, and formation of [An\textsuperscript{III}(TREN\textsuperscript{NPS})\textsubscript{3}] (Scheme 2), a product that is consistent with H\textsuperscript{1} NMR spectroscopic data. In the case of 1, new resonances appear at 18.75, 18.11, 4.43 and 0.05 ppm, with integral values that match well for the CH, CH, CH(CH\textsubscript{3})\textsubscript{2} and CH(CH\textsubscript{2})\textsubscript{2} protons, respectively, in the presumed reduced Np\textsuperscript{III} complex. Only very minor resonances remain from 1. In the case of 2, the resonances corresponding to 2 disappear and new resonances emerge, which are attributed to a reduced Pu(III) complex, but the resonances are broad making specific assignments less clear cut than in the Np\textsuperscript{III} case (see ESI).

Scheme 2 Suggested reduction process for 1 and 2, yielding An(III) species.

In both cases, the very high solubility of the An\textsuperscript{III} derivatives even in non-coordinating solvents such as hexanes, n-pentane, and HMDSO (hexamethyldisiloxane), coupled with the relatively small-scale of transuranic reactions compared to uranium, has prevented isolation of pure crystalline products. Nonetheless, following filtration to remove the KCl by-product of the reduction process, it should be entirely feasible to conduct a wide range of in situ reactivity studies. For example, initial reactions of Me\textsubscript{3}NO and TEMPO (2,2,6,6-Tetramethylpiperidine 1-oxyl) with [Np\textsuperscript{III}(TREN\textsuperscript{NPS})\textsubscript{3}] exhibit a distinct colour change, suggesting that oxidation occurs. Whether or not the product contains a terminal Np-O or bridging Np-O-Np moiety is the subject of ongoing longer-term studies that are beyond a communication scope, given the non-trivial nature of small-scale transuranic synthetic chemistry.

In conclusion, we have demonstrated that even on the small scales required for safe and practicable handling of transurane the TREN\textsuperscript{NPS} ligand can be installed on elements such as neptunium and plutonium to give actinide(V)-chloride complexes. It appears that the chloride anion can be exchanged for the softer iodide or can be removed by reduction to afford trivalent derivatives. As well as providing rare non-aqueous transuranic molecules that present a rich seam of data through examination of their structural, spectroscopic, and reactivity properties, this work provides potentially important flexibility in subsequent syntheses and invaluable precursors for our currently ongoing pursuit of transuranic metal-ligand multiple-bond species.

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Notes and references

† Caution: Np\textsuperscript{239} and Pu\textsuperscript{239} are high specific activity α-particle emitting radionuclides. This research was conducted in a radiological facility with appropriate analyses of hazards and implementation of controls to allow the safe handling and manipulation of radioactive materials. Experimental details concerning the reduction chemistry 1 and 2 along with other reactivity can be found in the ESI.

Synthesis of 1: THF (1 mL) was added to solid [NpCl\textsubscript{2}(DME\textsubscript{2})\textsubscript{2}] (0.0430 g, 0.0777 mmol) and cooled to -35 °C in the drybox freezer. Separately, [[Li][TREN\textsuperscript{NPS}]] (0.0487 g, 0.0777 mmol) was dissolved in THF (2 mL) and cooled to -35 °C in the drybox freezer. The cold ligand solution was added dropwise to the cold [NpCl\textsubscript{2}(DME\textsubscript{2})\textsubscript{2}]/THF mixture over the course of 45 min (with a few seconds of stirring after each addition followed by placing back in the freezer for a few minutes to keep cold). During the addition process an orange solution was initially formed that became orange-red. The resultant solution was then stirred at ambient temperature for 16 h affording a deep orange-red solution. Volatiles were removed in vacuo and the oily residue dried in vacuo for 15 min. The resulting solid residue was extracted into toluene (3 mL) at 50 °C, allowed to cool to ambient temperature and filtered through a Celite pad supported by a glass fibre filter circle. The volume of the red filtrate was reduced in vacuo to 1 mL and stored at -35 °C. After 2 d the red-orange microcrystalline solid that had deposited was collected and dried in vacuo. The supernatant was layered with HMDSO (1 mL) and a second crop of microcrystalline material was collected after 6 d, and dried in vacuo and combined with the first crop to give a final orange-red powder of 1 (0.0348 g, 51% yield). Single-crystals suitable for X-ray diffraction were obtained from a toluene/hexanes solution of 1 stored at -35 °C. Vis/NIR of 1 dissolved in toluene (in nm): 562, 569, 578, 587, 647, 661, 679, 690(sh), 736, 746, 767, 783, 805, 829, 852(sh), 865, 902, 939, 957(sh), 989(sh), 1003, 1011, 1029, 1058(sh), 1241, 1315. Diffuse reflectance vis/NIR of solid 1 (in nm): 562, 569, 579, 589, 647, 661, 679, 691, 736, 747, 767, 784, 805, 830, 844.
852(sh), 865, 902, 941, 957(sh), 987, 1002, 1014, 1030, 1060(sh), 1243, 1314. $^1$H NMR (in CD$_3$OD, 400 MHz): $\delta$ 7.71 (s, 54H, CH(CH$_3$)$_3$), 6.27 (s, 9H, CH(CH$_3$)$_2$), 16.44 (s, 6H, CH$_3$), -32.15 (s, 6H, CH$_2$).

**Synthesis of 2:** Solid [PuCl$_2$(DMF)$_3$] (0.0413 g, 0.074 mmol) was cooled to -35 °C in the drybox freezer. Separately, [(Li$_2$)[TREN]]$^+$ (0.0466 g, 0.074 mmol) was dissolved in THF (3 mL) and cooled to -35 °C in the drybox freezer. The cold ligand solution was added to the Pu-containing solid resulting in the immediate formation of an intensely coloured solution, which was placed back in the freezer to cool back to -35 °C, and then stirred for 5 min then stored in the freezer for 16 h before being allowed to warm to ambient temperature with stirring over 2 h. Volatiles were removed in vacuo, the residue triturated with hexanes (3 mL), and volatiles removed in vacuo. The residue was dissolved in hexanes (3 mL), filtered through Celite supported on a glass fibre filter circle, and the volume of the filtrate reduced in vacuo to 0.75 mL. Following layering of the solution with HMDSO (0.5 mL) and storage at -35 °C in the drybox freezer for 4 d the red crystalline material was collected and dried in vacuo (0.0122 g, 19% yield). Single-crystals suitable for X-ray diffraction were obtained from a hexane solution of 2 stored at -35 °C. Vis/NIR of 2 dissolved in toluene (in nm): 681, 690, 731, 741(sh), 850, 942, 1030, 1065(sh), 1080, 1120, 1147, 1241. Diffuse reflectance vis/NIR of solid 2 (in nm): 945, 1032, 1063(sh), 1079, 1116(sh), 1147, 1230. $^1$H NMR (in CD$_3$OD, 400 MHz): 6.01 (d, 54H, CH(CH$_3$)$_2$), 1.14 (s, br, 9H, CH(CH$_3$)$_3$), 0.58 (m, 6H, CH$_3$), -1.18 (m, 6H, CH$_2$).

$\Delta$ **Crystal data for 1:** C$_3$H$_5$Sb$_2$Cl$_6$NiSi$_2$, M = 884.69, monoclinic, a = 12.9845(10), b = 15.6008(12), c = 20.4035(15), $\alpha$ = 90.00, $\beta$ = 90.648(1), $\gamma$ = 90.00, $\nu_1$ = 4132.85(5) $\AA$ $^2$, $T = 140(1)$ K, space group P2$_1$/n, Z = 4, $\mu$ = 2.692 mm$^{-1}$, reflections collected/independent = 47211/59786 ($R_{int} = 0.0391$), $R_1$ ($> 2\sigma(I)$) = 0.0241, and $wR_2$ ($> 2\sigma(I)$) = 0.0448.

$\Delta$ **Crystal data for 2:** C$_3$H$_5$Sb$_2$Cl$_6$NiSi$_2$, M = 884.69, monoclinic, a = 13.0136(10), b = 15.5458(12), c = 20.4330(15), $\alpha$ = 90.00, $\beta$ = 90.546(1), $\gamma$ = 90.00, $\nu_1$ = 4133.5(5) $\AA$ $^2$, $T = 100(1)$ K, space group P2$_1$/n, Z = 4, $\mu$ = 1.772 mm$^{-1}$, reflections collected/independent = 94248/17439 ($R_{int} = 0.0250$), $R_1$ ($> 2\sigma(I)$) = 0.0291, and $wR_2$ ($> 2\sigma(I)$) = 0.0764.