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## Uranium–Nitrogen Multiple Bonding: Isostructural Anionic, Neutral, and Cationic Uranium Nitride Complexes Featuring a Linear U=N=U Core\*\*

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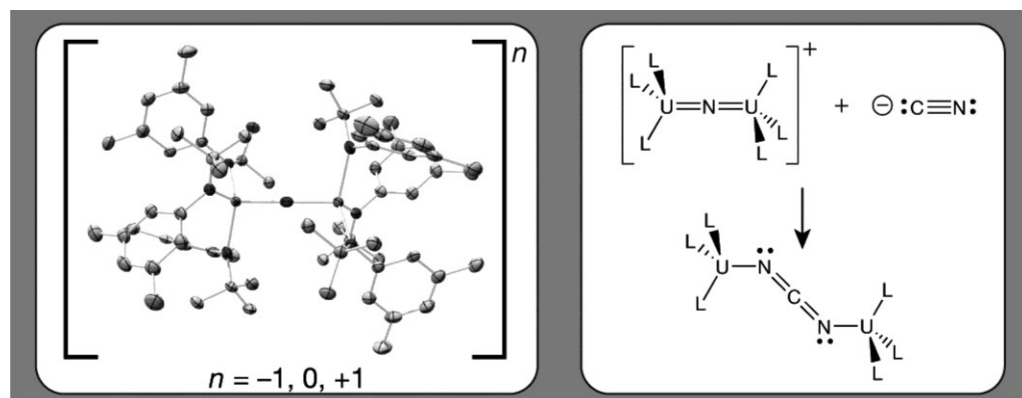
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### Supporting information:

Experimental details, including treatment of  $[3][B(Ar^F)_4]$  with CO and LiCCPh; crystallographic data and files in CIF format; and computational data relevant to the electronic structure of **3**. This material is available free of charge via the Internet at <http://pubs.acs.org>

### Graphical abstract:



## Abstract

Reaction of the uranium(III) tris(anilide) complex (THF)U(N[*t*-Bu]Ar)<sub>3</sub> (**1**, THF = tetrahydrofuran; Ar = 3,5-Me<sub>2</sub>C<sub>6</sub>H<sub>3</sub>) with MN<sub>3</sub> (M = Na, [N(*n*-Bu)<sub>4</sub>]) results in the formation of the bimetallic diuranium(IV/IV) complexes M[(μ-N)(U(N[*t*-Bu]Ar)<sub>3</sub>)<sub>2</sub>] (M[**3**]), which feature a single nitride ligand engaged as a linear, symmetric bridge between two uranium centers. The stability of the U=N=U core across multiple charge states is illustrated by stepwise chemical oxidation of Na[**3**] to the diuranium(IV/V) complex (μ-N)(U(N[*t*-Bu]Ar)<sub>3</sub>)<sub>2</sub> (**3**) and the diuranium(V/V) complex [(μ-N)(U(N[*t*-Bu]Ar)<sub>3</sub>)<sub>2</sub>][B(Ar<sup>F</sup>)<sub>4</sub>] {[**3**][B(Ar<sup>F</sup>)<sub>4</sub>]; Ar<sup>F</sup> = 3,5-(CF<sub>3</sub>)<sub>2</sub>C<sub>6</sub>H<sub>3</sub>}. M[**3**], **3**, and [**3**][B(Ar<sup>F</sup>)<sub>4</sub>] were characterized by NMR spectroscopy, single-crystal X-ray diffraction, and elemental analysis. The cyclic voltammogram of **3** reveals two clean, reversible one-electron electrochemical events at  $E_{1/2} = -1.69$  and  $-0.67$  V, assigned to the [**3**]<sup>-</sup>/**3** and **3**/[**3**]<sup>+</sup> redox couples, respectively. The X-ray crystal structures of [N(*n*-Bu)<sub>4</sub>][**3**], **3**, and [**3**][B(Ar<sup>F</sup>)<sub>4</sub>] reveal a linear U=N=U core that contracts by only ~0.03 Å across the [**3**]<sup>*n*</sup> (*n* = -1, 0, +1) series, an effect that is rationalized as being primarily electrostatic in origin. [**3**][B(Ar<sup>F</sup>)<sub>4</sub>] reacts with NaCN, eliminating Na[B(Ar<sup>F</sup>)<sub>4</sub>] and forming the known diuranium(IV/IV) cyanonitride complex (μ-NCN)(U(N[*t*-Bu]Ar)<sub>3</sub>)<sub>2</sub>, suggesting that the U=N=U core has metallonitrene-like character.

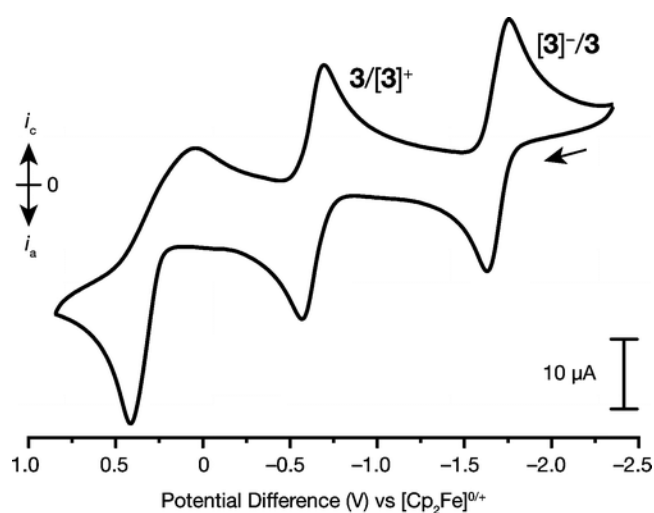
## Introduction

Uranium nitrides have been identified as worthy candidates for N-atom transfer reactions and aziridination of small molecules, and their potential use in catalytic cycles and as advanced nuclear fuels has been noted.<sup>1</sup> To date, only a handful of uranium nitride complexes have been reported, most of which are polymetallic nitride clusters derived from dinitrogen or azide activation<sup>2</sup> and binary and ternary nitrides generated and observed under matrix-isolation conditions.<sup>3</sup> Beyond uranium nitrides, assessing the degree of covalency and mixing of 5f and 6d orbitals in actinide–ligand bonding continues to be an active area of research<sup>4</sup> and has been recently extended to multiply bonded uranium imido complexes shown to accommodate multiple formal oxidation states at uranium.<sup>5</sup>

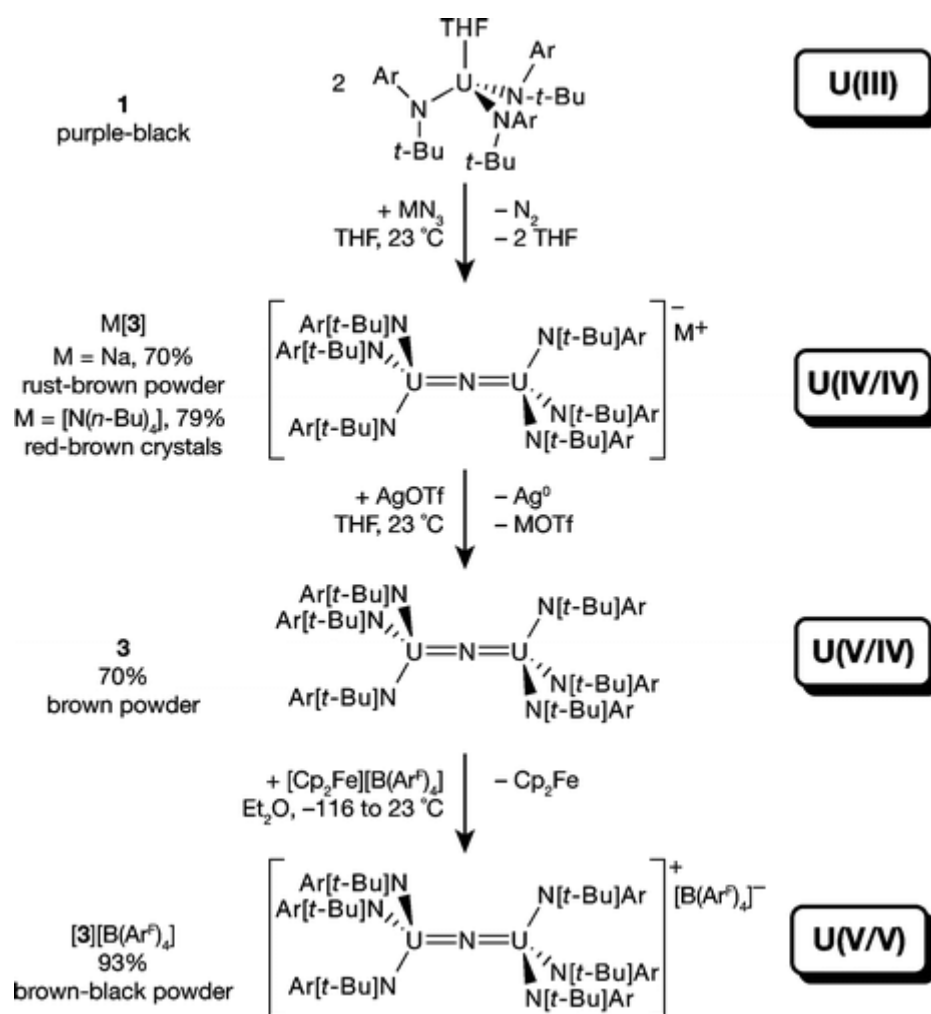
We recently reported on the reaction of (THF)U(N[*t*-Bu]Ar)<sub>3</sub> (**1**, THF = tetrahydrofuran; Ar = 3,5-Me<sub>2</sub>C<sub>6</sub>H<sub>3</sub>) with [N(*n*-Bu)<sub>4</sub>][B(N<sub>3</sub>)(C<sub>6</sub>F<sub>5</sub>)<sub>3</sub>], which directly furnishes the monometallic uranium(V) nitridoborate complex [N(*n*-Bu)<sub>4</sub>][(C<sub>6</sub>F<sub>5</sub>)<sub>3</sub>B–N≡U(N[*t*-Bu]Ar)<sub>3</sub>] ([N(*n*-Bu)<sub>4</sub>][**2**]) and provides access to the neutral uranium(VI) nitridoborate derivative (C<sub>6</sub>F<sub>5</sub>)<sub>3</sub>B–N≡U(N[*t*-Bu]Ar)<sub>3</sub> (**2**) by one-electron oxidation of [N(*n*-Bu)<sub>4</sub>][**2**].<sup>6</sup> In the synthesis of [N(*n*-Bu)<sub>4</sub>][**2**], the tris(pentafluorophenyl)boron fragment serves the role of protecting group by preventing the formation of polymetallic μ-nitrido products, thus facilitating isolation of the first examples of molecular monometallic uranium nitrides. Herein, we report on complexes derived from the action of unprotected azide ion on **1**, a series of bimetallic uranium nitride complexes possessing a linear U=N=U core that is stable across several charge states.

Treatment of a purple-black solution of **1** in THF with  $\text{NaN}_3$  (0.5 equiv) resulted in a gradual color change to red-brown over  $\sim 12$  h. A rust-brown powder was isolated in 70% yield following workup. The paramagnetically broadened  $^1\text{H}$  NMR spectrum, solution magnetic susceptibility ( $\mu_{\text{eff}} = 3.23 \mu_{\text{B}}$ ,  $20^\circ\text{C}$ , THF- $d_8$ ), and combustion analysis of the isolated material were consistent with the formation of the bimetallic diuranium(IV/IV) complex  $\text{Na}[(\mu\text{-N})(\text{U}(\text{N}[\textit{t}\text{-Bu}]\text{Ar})_3)_2]$  (**Na[3]**). Similarly, treatment of **1** with  $[\text{N}(\textit{n}\text{-Bu})_4][\text{N}_3]$  (0.5 equiv) in thawing THF provided  $[\text{N}(\textit{n}\text{-Bu})_4][\mathbf{3}]$  ( $\mu_{\text{eff}} = 3.22 \mu_{\text{B}}$ ,  $20^\circ\text{C}$ , THF- $d_8$ ) in 79% yield.

With a formal  $5f^2$  electron count for each uranium(IV) center in the bimetallic  $[\mathbf{3}]^-$  ion, the possibility for rich redox chemistry was realized (Scheme 1). Indeed, addition of  $\text{AgOTf}$  (1 equiv) to a solution of  $\text{Na}[\mathbf{3}]$  in THF resulted in formation of the neutral diuranium(IV/V) complex  $(\mu\text{-N})[\text{U}(\text{N}[\textit{t}\text{-Bu}]\text{Ar})_3]_2$  (**3**;  $\mu_{\text{eff}} = 3.85 \mu_{\text{B}}$ ,  $20^\circ\text{C}$ , chloroform- $d$ ), which was isolated as a brown powder in 70% yield by precipitating and filtering off the product following removal of precipitated  $\text{Ag}^0$  and separation from  $\text{NaOTf}$ . Furthermore, **3** reacted with  $[\text{Cp}_2\text{Fe}][\text{B}(\text{Ar}^{\text{F}})_4]$  [ $\text{Ar}^{\text{F}} = 3,5\text{-}(\text{CF}_3)_2\text{C}_6\text{H}_3$ ] in  $\text{Et}_2\text{O}$  to form the cationic diuranium(V/V) complex  $[(\mu\text{-N})(\text{U}(\text{N}[\textit{t}\text{-Bu}]\text{Ar})_3)_2][\text{B}(\text{Ar}^{\text{F}})_4]$  ( $[\mathbf{3}][\text{B}(\text{Ar}^{\text{F}})_4]$ ;  $\mu_{\text{eff}} = 2.86 \mu_{\text{B}}$ ,  $20^\circ\text{C}$ , chloroform- $d$ ), which was separated from the ferrocene coproduct by precipitation and isolated in 93% yield by filtration. The electrochemical potentials relating  $[\mathbf{3}]^-$ , **3**, and  $[\mathbf{3}]^+$  were measured via cyclic voltammetry using a solution of **3** in THF. Two reversible electrochemical events at  $E_{1/2} = -1.69$  and  $-0.64$  V (vs  $[\text{Cp}_2\text{Fe}]^{0/+}$ ) were observed and assigned to the  $[\mathbf{3}]^-/\mathbf{3}$  and  $\mathbf{3}/[\mathbf{3}]^+$  couples, respectively (Figure 1). The large separation between the  $[\mathbf{3}]^-/\mathbf{3}$  and  $\mathbf{3}/[\mathbf{3}]^+$  couples ( $|\Delta E_{1/2}| = 1.05$  V) corresponds to an equilibrium constant  $K_c \approx 5.6 \times 10^{17}$  for the comproportionation of  $[\mathbf{3}]^-$  and  $[\mathbf{3}]^+$ , suggesting strong electronic communication between the uranium centers.<sup>7</sup> A third electrochemical event observed at potentials above 0 V was not fully reversible at sweep rates as fast as  $1000 \text{ mV s}^{-1}$  and may be due to formation of an unstable  $[\mathbf{3}]^{2+}$  species.



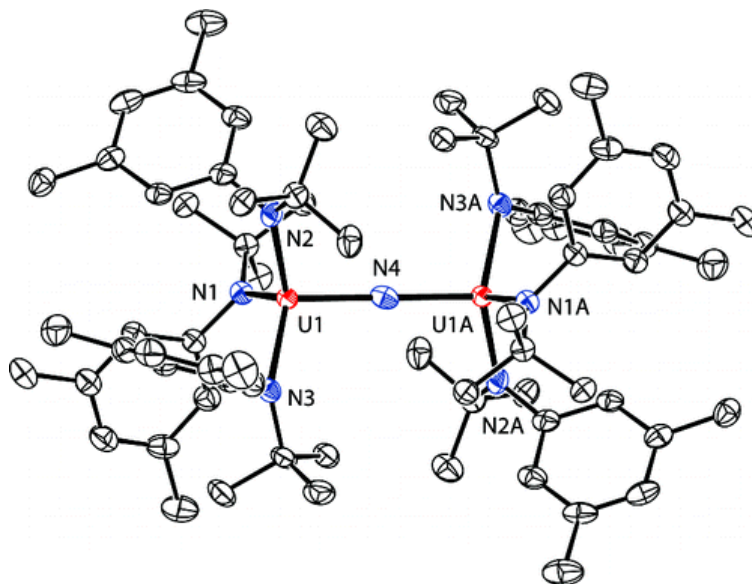
**Figure 1.** Cyclic voltammogram of **3** in THF ( $200 \text{ mV s}^{-1}$  sweep rate,  $\sim 0.1 \text{ M}$   $[\text{N}(\textit{n}\text{-Bu})_4][\text{B}(\text{C}_6\text{F}_5)_4]$  supporting electrolyte) showing the two one-electron redox couples that interconvert  $[\mathbf{3}]^n$  ( $n = -1, 0, +1$ ).



**Scheme 1.**

The solid-state structures of  $[N(n-Bu)_4][3]$ , **3** (Figure 2), and  $[3][B(Ar^F)_4]$  were determined by single-crystal X-ray diffraction, providing a rare opportunity to compare structural parameters of a U–N multiple bond over three charge states. Key metrical parameters describing the structures of  $[3]^n$  ( $n = -1, 0, +1$ ) are summarized in Table 1. In all cases, each uranium center is coordinated in a trigonal pseudo- $C_3$  fashion by three anilide ligands and occupies a terminus of a linear U=N=U bridge. Successive one-electron oxidation of the U=N=U core results in a decrease of the U–N<sub>nitride</sub> distance across a range of  $\sim 0.03$  Å, while the average U–N[t-Bu]Ar distance decreases across a comparably larger range of  $\sim 0.12$  Å. The observed contraction of the U=N=U core upon oxidation is likely primarily electrostatic in origin, where the removal of metal-based nonbonding electrons results in an increasingly electron-deficient U=N=U core. This structural dependence on charge is reminiscent of that displayed by penta- and hexavalent uranyl complexes, where the very covalent multiple bonds along the O=U=O axis are less responsive to the charge state than are the more ionic bonds in the molecular equator.<sup>8</sup> The U–N<sub>nitride</sub> distances of the  $[3]^n$  series fall in a range that is  $\sim 0.1$  Å longer than typical U–N distances observed in uranium imido complexes and are similar to those found in the octanuclear mixed

azide/nitride clusters  $[(C_5Me_4R)_2U(\mu-N)(C_5Me_4R)U(\mu-N_3)]_4$  ( $R = H, Me$ ) reported by Evans and co-workers<sup>2c</sup> (see Table S.2 in the Supporting Information for a more detailed comparison of relevant bond metrics).



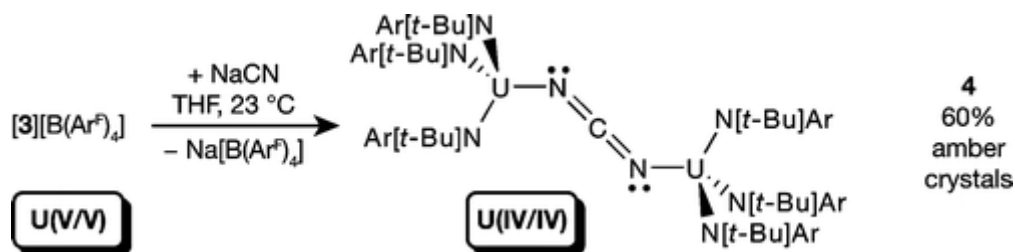
**Figure 2.** ORTEP rendering of **3** with ellipsoids displayed at 50% probability; hydrogen atoms and one molecule of cocrystallized Et<sub>2</sub>O have been omitted for clarity.

**Table 1.** Internuclear Distances (Å) and Angles (deg) for  $[(\mu-N)(U(N[t-Bu]Ar)_3)_2]^n$  ( $n = -1, 0, +1$ )

	<b>[N(<i>n</i>-Bu)<sub>4</sub>][3]</b>	<b>3</b>	<b>[3][B(Ar<sup>F</sup>)<sub>4</sub>]</b>
U–N	2.080(4)	2.0625(2)	2.0470(3)
	2.077(4)		2.0511(3)
U–N[ <i>t</i> -Bu]Ar (avg)	2.323	2.243	2.191
U–N–U	175.1(2)	180	180

Treatment of  $[3][B(Ar^F)_4]$  in THF with NaCN results in elimination of  $Na[B(Ar^F)_4]$  and formation of the known diuranium(IV/IV) cyanoimide complex  $(\mu-NCN)(U(N[t-Bu]Ar)_3)_2$  (**4**), which by this new reaction was isolated in 60% yield (Scheme 2). Prior to its synthesis from NaCN and  $[3][B(Ar^F)_4]$ , **4** had been prepared from **1** and a cyanonitrene ( $N\equiv C-N$ ) group transfer reagent.<sup>9</sup> Here, cyanide serves as a two-electron reducing agent, converting two uranium(V) centers to uranium(IV) while inserting into the nitride bridge. This is similar to a reaction reported by Meyer and co-workers<sup>10</sup> wherein a monometallic uranium(V) imido complex  $[(L)U(NSiMe_3)]$  reacts with methyl isocyanide to form  $Si_2Me_6$  and the uranium(IV) carbodiimide derivative  $[(L)U(NCNMe)]$ . More generally, Scheme 2 relates to the known reactivity of certain electrophilic metal nitride complexes toward reducing Lewis bases such as phosphines or carbon monoxide, yielding

phosphiniminato or isocyanate derivatives, respectively,<sup>11</sup> and shows as well that the U=N=U core of [3]<sup>+</sup> behaves as a masked metallonitrene.<sup>12</sup>



*Scheme 2.*

The complexes reported here provide insight into the structural and electrochemical characteristics of uranium nitrides. Insertion of cyanide into the U=N=U core of  $[3][B(Ar^F)_4]$  shows that the reaction chemistry of uranium nitrides extends beyond simple outer-sphere redox processes. This, along with the high polarity expected of a terminal uranium–nitride bond ( $U^{\delta+}-N^{\delta-}$ ) and the oxidizing nature of uranium in its higher oxidation states, intimates that a high-valent *terminal* uranium nitride would have the capacity to engage in bond-forming reactions at the nitride ligand via multiple mechanistic pathways. Further developments stemming from this system of uranium nitrides may be anticipated, especially regarding reactions that result in heterolytic fragmentation of the U=N=U core.

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